Future inhibition of ecosystem productivity by increasing wildfire pollution over boreal North America

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Abstract

Biomass burning is an important source of tropospheric ozone (O$_3$) and aerosols. These air 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 significantly in boreal North America by the midcentury, while little is known about the impacts of enhanced emissions on the terrestrial carbon budget. Here, combining site-level and satellite observations and a carbon-chemistry-climate model, we estimate the impacts of fire emitted O$_3$ and aerosols on net primary productivity (NPP) over boreal North America. Fire emissions are calculated based on an ensemble projection from 13 climate models. In the present day, wildfire enhances surface O$_3$ by 2 ppbv (7%) and aerosol optical depth (AOD) at 550 nm by 0.03 (26%) in the summer. By midcentury, area burned is predicted to increase by 66% in boreal North America, contributing more O$_3$ (13%) and aerosols (37%). Fire O$_3$ causes negligible impacts on NPP because ambient O$_3$ concentration (with fire contributions) 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 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 aerosols enhance NPP by 72 Tg C yr$^{-1}$ in the present day but decrease NPP by 118 Tg C yr$^{-1}$ in the future, mainly because of the soil moisture perturbations. Our results suggest that future wildfire may accelerate boreal carbon loss, not only through direct emissions increasing from 68 Tg C yr$^{-1}$ at present day to 130 Tg C yr$^{-1}$ by midcentury, but also through the biophysical impacts of fire aerosols.
1 Introduction

Wildfire area burned is increasing in recent decades over North America boreal regions (Stocks et al., 2002; Kasischke and Turetsky, 2006). Fire activity is closely related to weather conditions and large-scale atmospheric oscillations (Gillett et al., 2004; Duffy et al., 2005), and is projected to increase significantly in the future due to climatic changes (Flannigan et 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-Lamberty et al., 2007; Turetsky et al., 2011). Meanwhile, fire-induced air pollution, including ozone (O\textsubscript{3}) and aerosols, is predicted to increase in boreal and downwind regions by midcentury (Yue et al., 2013; Yue et al., 2015). Wildfire emissions have large impacts on air 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., 2017). However, little is known about how these pollutants affect ecosystem carbon assimilation, and how this impact will change with the increased wildfire activity in the future.

Surface O\textsubscript{3} causes damages to photosynthesis through stomatal uptake (Sitch et al., 2007). In the present climate state, fire-induced O\textsubscript{3} enhancements are predicted to reduce net primary productivity (NPP) in the Amazon forest by 230 Tg C yr\textsuperscript{-1} (1 Tg = 10\textsuperscript{12} g), a magnitude comparable to the direct release of CO\textsubscript{2} from fires in South America (Pacifico et al., 2015). 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 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) estimated that biomass burning aerosols increase Amazon NPP by 78–156 Tg C yr\textsuperscript{-1}, which offsets about half of the damage caused by fire O\textsubscript{3} (Pacifico et al., 2015). In contrast, strong light attenuation associated with high aerosol loading may decrease canopy photosynthesis (Cohan et al., 2002; Oliveira et al., 2007; Cirino et al., 2014). Furthermore, the aerosol radiative effects indirectly influence ecosystem productivity through concomitant meteorological perturbations that are only beginning to be examined (Yue et al., 2017).
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; Nitschke and Innes, 2008; Amiro et al., 2009; Balshi et al., 2009; Bergeron et al., 2010; Wotton et al., 2010; de Groot et al., 2013; Wang et al., 2016). For example, Amiro et al. (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 (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 2045-2050 relative to 1991-2000, using the Multivariate Adaptive Regression Splines (MARS) 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 projections originating from both fire models and simulated future climate. However, even 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 models (Moritz et al., 2012; Yue et al., 2013). The multi-model ensemble approach has shown superior predictability over single models in historical climate simulations (Flato et al., 2013) and near-term climate predictions (Kirtman et al., 2014), and has been used as a standard technique to assess changes of climate variables in the long-term projections (Collins et al., 2013). Following this strategy, Yue et al. (2015) used output from 13 climate 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 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 Canada. The study found regional maximum O₃ enhancements as high as 15 ppbv, suggesting the potential for possible vegetation damage and land carbon loss due to the enhanced boreal fire-related air pollution. Wildfire aerosols are also expected to increase significantly but not predicted in Yue et al. (2015).

In this study, we quantify the impacts of O₃ 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 relationships between gross primary production (GPP) and aerosol optical depth (AOD) at 550 nm over the boreal regions based on observations. We then perform a suite of Earth system model simulations using NASA GISS ModelE2 that embeds the Yale Interactive Terrestrial Biosphere model (YIBs), a framework known as ModelE2-YIBs (Yue and Unger, 2015). Future projections of wildfire emissions from Yue et al. (2015) are applied as input to ModelE2-YIBs model to project fire-induced O\textsubscript{3} and aerosol concentrations in the 2010s and 2050s. The impacts of the boreal fire O\textsubscript{3} on forest photosynthesis are predicted using the flux-based damage algorithm proposed by Sitch et al. (2007), which has been fully evaluated against available O\textsubscript{3} 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 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 isolate the contributions of changes in the individual meteorological drivers.

2 Materials and methods

2.1 Observed GPP-AOD relationships

Following the approach by Strada et al. (2015), we investigate the GPP sensitivity to diffuse 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 boreal sites than those in temperate regions. We select AMF sites providing hourly (or half-hourly) simultaneous measurements of GPP (non gap-filled) and photosynthetically active 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 Quebec Mature Boreal Forest Site (CA-Qfo, 73.4°W, 49.7°N), an evergreen needleleaf forest (ENF). At the two selected sites, we calculate the Pearson’s correlation coefficients between half-hourly GPP and different components of PAR. In total, we select 2432 and 3201 pairs of GPP and PAR measurements at CA-Gro and CA-Qfo, respectively. We then apply instantaneous Level 2 Collection 6 of AOD pixels at 3-km resolution retrieved by the Moderate Resolution Imaging Spectroradiometer (MODIS, https://ladsweb.nascom.nasa.gov/)
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., 2013) and urban/suburban (Munchak et al., 2013) scales. Strada et al. (2015) used ground-based AOD observations from the Aerosol Robotic Network (AERONET) near AMF sites to validate the sampling technique of MODIS 3-km AOD product. They found high correlations of 0.89-0.98 and regression slopes from 0.89 to 1.03 for daily AOD between AERONET and 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 sites.

Every day, MODIS satellite sensors pass a specific region between 10:00 and 14:00 Local 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 the GPP-AOD relationships to the noontime (10:00-14:00 LT) from June to August. The chosen noontime window limits the contributions that confounding factors such as low solar angles and high diffuse fraction may have on the amount of diffuse PAR and plant productivity (Niyogi et al., 2004). For each summer day, we select instantaneous MODIS 3-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. 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 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 between “quasi-coincident” GPP and AOD at the selected sites. Negative GPP is considered as a missing value. To further reduce the influence of cloud cover, we discard instants (both AMF and MODIS data) when precipitation is non-zero. In total, we select 65 pairs of GPP 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 and GPP data are available. In addition, AOD is screened for clear instants to exclude the impacts of clouds.

2.2 Wildfire emissions
Wildfire emissions used in climate modeling are calculated as the product of area burned, fuel consumption, and emission factors. To predict area burned, we build stepwise regressions for area burned in 12 boreal ecoregions (Yue et al., 2015). Observed area burned aggregated from inter-agency fire reports is used as the predictand. Predictors are selected from 44 (5×6+7×2) variables including five meteorological parameters (mean and maximum temperature, relative humidity, precipitation, and geopotential height at 500 hPa) of six 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 system during fire season. We consider the impacts of antecedent factors on current fire 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 ecoregions, depending on the selection of the factors that contribute the maximum observed variance in predictand but remain the minimum collinearity among predictors. These 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 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 in IPCC RCP8.5 scenario (van Vuuren et al., 2011) archived for the Coupled Model Intercomparison Project Phase 5 (CMIP5).

We derive 1°×1° gridded area burned based on the prediction for each ecoregion following 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 boreal ecoregion during 1980-2009. Spatially, large fires tend to burn in ecosystems where historical fires are frequent because of favorable conditions (Keane et al., 2008). In each 1°×1° grid square, we calculate the frequency of large fires (>1000 ha) during 1980-2009; these 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 1000 ha. We then allocate these large fires among the 1°×1° grid cells based on the observed spatial probability of large fires. For example, if one grid box (named grid ‘A’) bears 1% of large fires (>1000 ha) within an ecoregion at present day, the same grid will bear the same 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...
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 modified probability distribution to allocate large fires for the remaining months. For example, if present-day fires have consumed 20% of the total area within the grid ‘A’, then the possibility of large fire will be 0.8% (1%×0.8, instead of 1%) for this grid. Finally, we disaggregate the remaining 15% of area burned into fires 10 ha in size, and randomly 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 tests show that specifying different area burned to the large fires (100 or 10 000 ha rather than 1000 ha) yields < 1 % changes in predicted biomass burned, suggesting that this approach is not sensitive to the presumed fire size in the allocation procedure.

Fuel consumption, the dry mass burned per fire area, is the product of fuel load and burning severity. For fuel load in Alaska, we use 1-km inventory from the US Forest Service (USFS) Fuel Characteristic Classification System (FCCS, McKenzie et al., 2007). For fuel load in Canada, we use a 1-km fuel type map from the Canadian Fire Behavior Prediction (FBP) system (Nadeau et al., 2005), combined with fuel-bed definition from the FCCS. Burning severity, the fraction of fuel load burned by fires, is calculated with the USFS CONSUME model 3.0 following the approach described in Val Martin et al. (2012). With both fuel load 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. (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; Friend et al., 2014; Kim et al., 2017). Instead, we consider changes in burning severity due to perturbations in fuel moisture as indicated by CFWI indexes (Yue et al., 2015). On average, 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 (Flannigan et al., 2016).

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 NOx. 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 NOx, 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.

2.3 NASA ModelE2-YIBs model

The NASA ModelE2-YIBs is an interactive climate-carbon-chemistry model, which couples 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 horizontal resolution of 2°x2.5° latitude by longitude and 40 vertical layers up to 0.1 hPa. It dynamically simulates both the physical (emissions, transport, and deposition) and chemical (production, conversion, and loss) processes of gas-phase chemistry (NOx, HOx, Ox, CO, CH4, and NMVOCs), aerosols (sulfate, nitrate, ammonium, BC, OC, dust, and sea salt), and their interactions. In the model, oxidants influence the photochemical formation of secondary aerosol species (e.g., sulfate, nitrate, and biogenic secondary organic aerosol), in turn, aerosols alter photolysis rates and influence the online gas-phase chemistry. Size-dependent optical parameters computed from Mie scattering, including extinction coefficient, single scattering albedo, and asymmetry parameters, are applied for each aerosol type (Schmidt et al., 2014). The model also considers interactions between climate and atmospheric components. Simulated climate affects formation, transport, and deposition of atmospheric components, in turn, both O3 and aerosols influence climate by altering radiation, temperature, precipitation, and other climatic variables. Both observation-based evaluations and multi-model inter-comparisons indicate that ModelE2 demonstrates skill in simulating climatology (Schmidt et al., 2014), soil moisture (Fig. S1), radiation (Wild et al., 2013), atmospheric composition (Shindell et al., 2013b), and radiative effects (Shindell et al., 2013a).

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,
Plant respiration considers thermal dependence as well as acclimation to temperature (Atkin and Tjoelker, 2003). Soil respiration is calculated based on the carbon flows among 12 biogeochemical pools (Schaefer et al., 2008). Net carbon uptake is allocated among leaves, stems, and roots to support leaf development and plant growth (Cox, 2001). The YIBs model has been benchmarked against in situ GPP from 145 eddy covariance flux tower sites and satellite retrievals of LAI and phenology (Yue and Unger, 2015). An interactive flux-based 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 dependent on excess O₃ stomatal flux within leaves, which is a function of ambient O₃ concentration, boundary layer resistance, and stomatal resistance. Reduction of photosynthesis is calculated on the basis of plant functional types (PFTs), each of which bears a range of low-to-high sensitivities to O₃ uptake.

### 2.4 Simulations

Using the NASA ModelE2-YIBs model, we perform 6 time-slice simulations, three for 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 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 (both natural and anthropogenic) and related photosynthetic responses are included. Simulations F10AERO and F50AERO consider the responses of plant productivity to perturbations in radiation and meteorology caused by aerosols, including emissions from wildfires and other sources, but do not include any O₃ vegetation damage. In contrast, simulations F10O3 and F50O3 calculate offline O₃ damage based on the simulated O₃ from 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 back into the model to influence carbon allocation and tree growth. Plant respiration is changing in response to meteorological perturbations, either due to climate change or aerosol radiative effects. We assume no impact of O₃ damage to plant respiration and examine vegetation NPP, the net carbon uptake by biosphere, for the current study. The difference between AERO and CTRL runs isolates the impacts of fire aerosols on NPP, and the
difference between O3 and CTRL runs isolates O₃ vegetation damage caused by fire and non-fire emission sources.

All simulations are conducted for 20 years and outputs for the last 15 years are used for analyses. The simulations apply sea surface temperatures (SSTs) and sea ice distributions from previous NASA GISS experiments under the IPCC RCP8.5 scenario (van Vuuren et al., 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) 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 well-mixed greenhouse gas concentrations and anthropogenic emissions of short-lived species, both at present day and midcentury, are adopted from the RCP8.5 scenario (Table 2). The enhancement of CO₂ will affect climate (through longwave absorption) and ecosystem productivity (through CO₂ fertilization), but not the fire activity and related emissions directly. Natural emissions of soil and lightning NOₓ, biogenic volatile organic compounds (BVOC), dust, and sea salt are climate-sensitive and simulated interactively. The YIBs vegetation model cannot simulate changes in PFT fractions. The RCP8.5 land cover change 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 in the 2050s, resulting from a 1% loss in deciduous forest and another 1% loss in needleleaf forest over boreal North America. As a result, a land cover dataset derived from satellite retrievals (Hansen et al., 2003) is applied as boundary conditions for both the 2010s and 2050s.

To evaluate the simulated GPP responses to changes in diffuse radiation, we perform site-level simulations using standalone YIBs model, which is driven with observed hourly 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 impact of individual aerosol-induced climatic perturbations on NPP, we perform 10 sensitivity experiments using the offline YIBs model driven with offline meteorology 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 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
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.

2.5 Observation datasets

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 covariance data using an ensemble of regression trees (Jung et al., 2009). For AOD observations, we use satellite retrieval at 550 nm from Terra MODIS Level 3 data product. 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 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 period to represent present-day climatological conditions. Gridded datasets are interpolated to the same 2°×2.5° resolution as ModelE2-YIBs model.

3 Results

3.1 Observed GPP-AOD relationships

Positive correlations between GPP and diffuse PAR are found at the two boreal sites (Figs 2b-2c). The magnitude of diffuse PAR is similar for these sites, possibly because they are located at similar latitudes (Fig. 2a). GPP values at CA-Gro are generally higher than that at CA-Qfo, likely because deciduous broadleaf forest (DBF) has higher photosynthetic rates. Consequently, the slope of regression between GPP and PAR₃₉ is higher at CA-Gro than that at CA-Qfo, suggesting that GPP of DBF (or MF) is more sensitive to changes in diffuse PAR than that of ENF. We find almost zero correlation between GPP and PAR₃₉ at the two sites (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 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 ± range) is estimated as 3.5 ± 1.1 µmol m$^{-2}$ s$^{-1}$ per unit AOD at lower latitudes of boreal regions in the summer.

### 3.2 Model evaluations

Simulated summer GPP shows high values in mid-western Canada (Alberta and 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 spatial distribution with a high correlation coefficient of 0.77 ($p << 0.01$) and relatively small biases within 20% of the data product. Simulated AOD reproduces the observed spatial pattern including the high values in boreal forests (Fig. 3b). In contrast to the MODIS observations, predicted AOD is relatively uniform over the West with a background value of ~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, 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 daily 8-hour average (MDA8) [O$_3$] 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 observations (Fig. 4b), but the model overestimates the measured surface O$_3$ by 22%. The Canadian measurement sites are located near the southern boundary, and as a result do not represent the average state over the vast boreal region at higher latitudes.

With the Sitch et al. (2007) scheme, the YIBs model simulates reasonable GPP responses to [O$_3$] in North America (Yue and Unger, 2014; Yue et al., 2016). Generally, damage to GPP increases with the enhancement of ambient [O$_3$], but with varied sensitivities for different plant species (see Fig. 6 of Yue and Unger (2014)). In responses to the same level of [O$_3$], predicted O$_3$ 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
observed light responses of GPP to diffuse radiation in boreal regions. With the site-level simulations, we evaluate the modeled GPP-PAR\textsubscript{dif} relationships at the hourly (instead of half-hourly) time step during summer. For 1342 pairs of GPP and PAR\textsubscript{dif} at the site CA-Gro, the observed correlation coefficient is 0.42 and regression slope is 0.011, while the results for the simulation are 0.60 and 0.014, respectively. At the site CA-Qfo, the observations yield a correlation coefficient of 0.46 and regression slope of 0.007 for 1777 pairs of GPP and PAR\textsubscript{dif}. The simulated correlation is 0.61 and the regression is 0.011 at the same site. The GPP sensitivity to PAR\textsubscript{dif} in the model is slightly higher than that of the available observations, likely because the latter are affected by additional non-meteorological abiotic factors. To remove the influences of compound factors other than radiation, we follow the approach of Mercado et al. (2009) to discriminate GPP responses to ‘diffuse’ and ‘direct’ 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 diffuse light than that for direct light, suggesting that increase of diffuse radiation is a benefit for plant growth.

3.3 Simulation of wildfire O\textsubscript{3} and aerosols

During 1980-2009, wildfire is observed to burn 2.76×10\textsuperscript{6} ha and 156.3 Tg DM every year over boreal North America. Similarly, the ensemble prediction with fire regression models estimates present-day area burned of 2.88 ×10\textsuperscript{6} ha yr\textsuperscript{-1} and biomass burned of 160.2 Tg DM yr\textsuperscript{-1} (Yue et al., 2015). By the midcentury, area burned is projected to increase by 77% (to 5.10 ×10\textsuperscript{6} ha yr\textsuperscript{-1}) in boreal North America, mainly because of the higher temperature in future fire seasons. Consequently, biomass burned increases by 93% (to 308.6 Tg DM yr\textsuperscript{-1}) because fuel consumption also increases by 9% on average in a drier climate (Yue et al., 2015). Enhanced fire emissions increase concentrations of surface O\textsubscript{3} and column AOD, especially over Alaska and central Canada (Fig. 6). The maximum centers of air pollutants are collocated for O\textsubscript{3} and AOD but with unproportional magnitudes, suggesting non-linear conversion among fire emission species as well as the interactions with natural emission sources (e.g., lightning/soil NO\textsubscript{x} and BVOC). On average, wildfire emissions contribute 7.1 ± 3.1\% (2.1 ± 0.9 ppbv) to surface O\textsubscript{3} and 25.7 ± 2.4\% (0.03 ± 0.003) to AOD in the summer over boreal North America in the present day. By midcentury, these ratios increase significantly to 12.8 ± 2.8\% (4.2 ± 0.9 ppbv) for O\textsubscript{3} and 36.7 ± 2.0\% (0.05 ± 0.003) for AOD.
3.4 Simulation of fire pollution impacts on NPP

Surface O$_3$, including both fire and non-fire emissions (Table 2), causes limited (1-2%) damages to summer GPP in boreal North America (Fig. 7). The most significant damage is predicted over eastern U.S., where observed [O$_3$] is high over vast forest ecosystems (Fig. 4). In the western U.S., [O$_3$] is also high but the O$_3$-induced GPP reduction is trivial because low stomatal conductance in the semi-arid ecosystems limits O$_3$ uptake there (Yue and Unger, 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 cover). Both species have shown relatively high O$_3$ tolerance with a damaging threshold of 40 ppbv as calculated with Sitch’s scheme (Yue and Unger, 2014). For boreal regions, the mean [O$_3$] of 28 ppbv (Fig. 4a) is much lower than this damaging threshold, explaining why the excess O$_3$ stomatal flux (the flux causing damages) is low there (Fig. 8). Statistics in Yue et al. (2015) show that maximum daily 8-hour average (MDA8) [O$_3$] with fire contributions can be higher than 40 ppbv in Alaska and Canada. However, such episodes appear at 95 percentile for present day and 90 percentile for midcentury, suggesting that O$_3$ vegetation damage is rare in boreal North America and fire-induced O$_3$ enhancement does not exacerbate such damages. Therefore, we do not consider O$_3$ damage effects further.

Fire aerosols cause significant perturbations in shortwave radiation at surface (Fig. 9). The 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 scattering. In the present day, the average reduction of 5.6 W m$^{-2}$ in the direct light component is in part offset by the enhancement of 2.6 W m$^{-2}$ in the diffuse light component, leading to a net reduction of 3.0 W m$^{-2}$ in solar radiation over boreal North America. By the midcentury, a stronger reduction of 9.5 W m$^{-2}$ in direct light is accompanied by an increase of 4.0 W m$^{-2}$ in diffuse light, resulting in a net reduction of 5.5 W m$^{-2}$ in solar radiation. Fire-induced BC aerosols strongly absorb solar radiation in the atmospheric column (Figs 10a-10b). On average, fire aerosols absorb 1.5 W m$^{-2}$ in the present day and 2.6 W m$^{-2}$ by the midcentury.
Atmospheric circulation patterns respond to the aerosol-induced radiative perturbations (Figs 10c-10d). Surface radiative cooling and atmospheric heating together increase air stability and induce anomalous subsidence. In the present day, such descending motion is confined to 55-68°N, accompanied by a rising motion at 52-55°N (Fig. 10c). As a result, fire aerosols induce surface warming at higher latitudes but cooling at lower latitudes in boreal regions (Fig. 11a). Meanwhile, precipitation is inhibited by the subsidence in northwestern Canada 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 heating (Fig. 10b). The downward convection of warm air offsets surface radiative cooling (Fig. 9b), leading to a significant warming in the Southwest (Fig. 11b). The expanded subsidence further inhibits precipitation in vast domain of Canada (Fig. 11d). Soil moisture is closely related to rainfall and as a result exhibits dipole changes (drier north and wetter south) in the present day (Fig. 11e) but widespread reductions (Fig. 11f) by the midcentury.

In response to the climatic effects of fire aerosols, boreal NPP shows distinct changes between the present day and midcentury (Fig. 12). Such changes in NPP are a consequence of changes in GPP and autotrophic respiration (Fig. S2). Variations in plant respiration resemble those of GPP, because higher photosynthesis leads to faster leaf/tissue development, resulting in larger maintenance and growth respiration. In the 2010s, forest NPP increases by 5-15% in Alaska and southern Canada, but decreases by 5-10% in northern and eastern Canada. This pattern of NPP changes (ΔNPP) is connected to the climatic effects of aerosols, especially changes in soil moisture (Fig. 11). The correlation between ΔNPP (Fig. 12a) and changes in soil moisture (Fig. 11e) reaches $R = 0.56$ ($n = 356$), much higher than the values of $R = -0.11$ 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 of carbon uptake occurs in southern Canada (Fig. 3a), where positive NPP change is predicted (Fig. 12a), wildfire aerosols enhance the total NPP by 72 Tg C yr$^{-1}$ in the present day (Table 5). In contrast, increased wildfire emissions in the 2050s inhibit precipitation (Fig. 11d) and decrease soil moisture in boreal North America (Fig. 11f), leading to widespread NPP reductions and a total NPP loss of 118 Tg C yr$^{-1}$ (Fig. 12b, Table 5).

4 Discussion
4.1 Roles of aerosol climatic feedback

The contrasting sign of NPP responses in the present day and midcentury are closely related to the aerosol-induced surface climatic feedback. Sensitivity experiments using offline YIBs model (Table 3) allowed assessment of the impacts of individual changes in the major meteorological drivers, including temperature, radiation (diffuse and direct), and soil moisture (Table 5). The offline simulations driven with changes in all three variables yield $\Delta$NPP of 126 Tg C yr$^{-1}$ for the 2010s and -97 Tg C yr$^{-1}$ for the 2050s. These values are different from the online simulations, which predict $\Delta$NPP of 72 Tg C yr$^{-1}$ for the 2010s and -118 Tg C yr$^{-1}$ for the 2050s. Missing of other aerosol climatic feedbacks in the offline model, for example, changes in relative humidity, surface pressure, soil temperature, and turbulence momentum, may cause such discrepancy between the online and offline simulations.

Seasonal analyses show that summertime $\Delta$NPP is 99 Tg C at present day and -95 Tg C at midcentury, dominating the NPP changes all through the year, because both wildfire emissions and ecosystem photosynthesis maximize in boreal summer.

Observations show that aerosols can promote plant photosynthesis through increasing diffuse 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), and the model reproduces observed GPP responses to perturbations in direct and diffuse PAR (Fig. 5). Wildfire aerosols enhance diffuse radiation by 2.6 W m$^{-2}$ (1.7%) at present day and 4.0 W m$^{-2}$ (2.3%) at midcentury in boreal North America (Fig. 9). With these changes, simulated NPP increases by 8 Tg C yr$^{-1}$ at the 2010s and 14 Tg C yr$^{-1}$ at the 2050s (Table 5). Near the two AmeriFlux sites (Fig. 2a), wildfires increase local AOD by 0.03 (Fig. 6c).

Meanwhile, we estimate that summer average (00:00-24:00) GPP increases by 0.04 µmol m$^{-2}$ s$^{-1}$ in the same region due to aerosol diffuse fertilization effects (DFE) based on the results of (Y10_PAR – Y10_CTRL). This change suggests a simulated GPP sensitivity of 1.2 µmol m$^{-2}$ s$^{-1}$ (22%) per unit AOD. Observed GPP sensitivity to AOD at the two sites are 2.3 (19%) and 4.5 µmol m$^{-2}$ s$^{-1}$ (58%) per unit AOD, respectively (Figs 2d-2e). The absolute value of GPP sensitivity from simulations is much smaller than that of observations, because the former is 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%.
The estimated NPP changes of 8 Tg C yr\(^{-1}\) by the radiative effects of boreal fire aerosols are much weaker than the enhancement of 78-156 Tg C yr\(^{-1}\) by fires in Amazon basin (Rap et al., 2015). There are at least two reasons for such a difference in the DFE between boreal and 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 2010s, resulting in enhancement of summer AOD by 35% and diffuse radiation by 1.7%. These boreal emissions are much smaller than the ~240 Tg C yr\(^{-1}\) in Amazon basin (van der Werf et al., 2010), where fires enhances regional PM2.5 concentrations by 85% and diffuse radiation by 6.2% in dry seasons (Rap et al., 2015). Second, larger solar insolation in lower 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 insolation is not so abundant as that at the tropical areas. Consequently, decline of direct 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 shaded leaves. For this study, we do not find GPP reduction by the decline of direct light at the two Ameriflux sites (Table 4), possibly because these sites are located at middle latitudes (<50°N). In the future, more observations at higher latitudes (> 55°N) are required to explore the sensitivity of GPP to AOD at the light-limited conditions.

Simulations have shown that absorbing aerosols can cause regional drought by increasing air 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 emissions (Figs 11c-11f). Observations suggest that precipitation (and the associated soil moisture) is the dominant driver of the changes in GPP over North America, especially for the domain of cropland (Beer et al., 2010). Sensitivity experiments with offline YIBs model show that changes in soil moisture account for 82.5% of ΔNPP at present day and 70.5% of ΔNPP at midcentury (Table 5). These results suggest that aerosol-induced changes in soil water availability, instead of temperature and radiation, dominantly contribute to the changes of boreal NPP, consistent with observational and experimental results (Ma et al., 2012; Girardin et al., 2016; Chen et al., 2017).

4.2 Limitations and uncertainties
In this study, we examine the interactions among climate change, fire activity, air pollution, 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 → fire → pollution → biosphere” (Fig. 1). However, our choice of feedback analysis does not mean that the 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 by BVOC emissions (climate → biosphere → fire/pollution). In addition, other feedbacks may amplify ecosystem responses but are not considered. For example, the drought caused by fire aerosols in the midcentury (Fig. 11) may help increase fire activity (fire → pollution → climate → fire). Furthermore, we apply fixed SSTs in the climate simulations because reliable ocean heat fluxes for the future world were not available. Many previous studies have investigated regional aerosol-climate feedbacks without ocean responses. For example, Cook et al. (2009) found that dust-climate-vegetation feedback promotes drought in U.S., with a climate model driven by prescribed SSTs. Similarly, Liu (2005) found fire aerosols enhance regional drought using a regional climate model, which even ignores the feedback between 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, this study for the first time quantifies the indirect impacts of wildfire on long-range ecosystem productivity under climate change.

We use the ensemble projected fire emissions from Yue et al. (2015). Area burned is predicted based on the simulated meteorology from multiple climate models. Such an 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 climate scenarios and fire models. All predictions in Yue et al. (2015) are performed under the IPCC A1B scenario. With two different scenarios, A2 of high emissions and B2 of low emissions, Balshi et al. (2009) showed that future area burned in boreal North America increases at a similar rate until the 2050s, after which area burned in A2 scenario increases much faster than that in B2 scenario. On average, boreal area burned in Balshi et al. (2009) increases by ~160% at 2051-2060 compared with 2001-2010, much higher than the change of 66% in Yue et al. (2015). In contrast, Amiro et al. (2009) predicted that boreal area burned at the 2×CO₂ scenario increases only by 34% relative to the 1×CO₂ scenario. This ratio is only
half of the estimate in Yue et al. (2015), which compared results between periods with 1.44×CO₂ and 1×CO₂. The discrepancies among these studies are more likely attributed to the differences in fire models. Although both Amiro et al. (2009) and Yue et al. (2015) developed fire-weather regressions in boreal ecoregions, the former study did not include geopotential height at 500 hPa and surface relative humidity as predictors, which make dominant contributions to area burned changes in the latter study. On the other hand, Balshi 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 previous estimates, Yue et al. (2015) predicted median increases in future fire emissions over boreal North America.

We apply constant land cover and fuel load for both present day and midcentury, but we estimate an increase in fuel consumption due to changes in fuel moisture. Future projection of 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 projections, Heyder et al. (2011) predicted a large-scale dieback in boreal-temperate forests due to increased heat and drought stress in the coming decades. On the contrary, projections 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 addition, compound factors such as greenhouse gas mitigation (Kim et al., 2017), pine beetle outbreak (Kurz et al., 2008), and fire management (Doerr and Santin, 2016) may exert varied 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 fuel consumption (Flannigan et al., 2016). With constant fuel load but climate-driven fuel moisture, we calculate a 9% increase in boreal fuel consumption by the midcentury (Yue et al., 2015). Although such increment is higher than the prediction of 2-5% by Amiro et al. (2009) for a doubled-CO₂ climate, the consumption-induced uncertainty for fire emission is likely limited because changes in area burned are much more profound.

Predicted surface [O₃] is much higher than observations over boreal North America (Fig. 4). This bias does not affect main conclusions of this study, because predicted O₃ causes limited damages to boreal GPP even with the overestimated [O₃] (Fig. 7). The result confirms that fire-induced O₃ vegetation damage is negligible in boreal North America. For aerosols, the
model captures reasonable spatial pattern of AOD but with a background value of ~0.1 outside fire-prone regions, where the observed AOD is usually 0.1-0.2 (Fig. 3). This discrepancy may be related to the insufficient representations of physical and chemical processes in the model, but may also result from the retrieval biases in MODIS data due to the poor surface conditions (Liu et al., 2005) and small AOD variations (Vachon et al., 2004) at high latitudes.

Simulated aerosol climatic effects depend on radiative and physical processes implemented in the climate model. We find that present-day boreal fire aerosols on average absorb 1.5 W m\(^{-2}\) in the atmosphere (Fig. 10), which is much smaller than the value of 20.5 ± 9.3 W m\(^{-2}\) for fires in equatorial Asia (Tosca et al., 2010). This is because boreal fires enhance AOD only 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 (Feingold et al., 2005; Tosca et al., 2010; Liu et al., 2014). Most of these results were based on the direct and/or semi-direct radiative effects of fire aerosols. Inclusion of the indirect aerosol effect may further inhibit precipitation and amplify drought, but may also introduce additional uncertainties for the simulations. The fire-drought interaction may promote fire activity, especially in a warmer climate. Ignoring this interaction may underestimate future area burned and the consequent emissions.

4.3 Implications

Inverse modeling studies have shown that the land ecosystems of boreal North America are carbon neutral in the present day, with the estimated land-to-air carbon flux from -270 ± 130 Tg C yr\(^{-1}\) to 300 ± 500 Tg C yr\(^{-1}\) (Gurney et al., 2002; Rodenbeck et al., 2003; Baker et al., 2006; Jacobson et al., 2007; Deng et al., 2014). Here, we reveal a missing land carbon source 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 Tg C yr\(^{-1}\) in the present day, comparable to the direct carbon loss of 68 Tg C yr\(^{-1}\) from wildfire CO\(_2\) emissions (product of biomass burned and CO\(_2\) emission factors). By midcentury, increasing fire emissions instead cause a NPP reduction of 118 Tg C yr\(^{-1}\) due to 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)
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 wildfire-related air pollution risk to food production.

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 variability in plant species, topography, and climatology across different boreal regions. Such differences indicate distinct GPP sensitivities as well as fire characteristics. At lower latitudes, where anthropogenic pollution emissions are more abundant, ambient ozone concentrations may have exceeded damaging thresholds for most plant species. In those regions, additional 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 NPP by 230 Tg C yr\(^{-1}\) through the generation of surface ozone (Pacifico et al., 2015). Meanwhile, solar radiation is more abundant at lower latitudes, indicating more efficient increases in photosynthesis through aerosol DFE because the sunlit leaves receive saturated direct light in those regions. As shown in Beer et al. (2010), partial correlations between GPP and solar radiation are positive in boreal regions but negative over the subtropics/tropics, 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 to understand the net impacts of ozone and aerosols from biomass burning on the global carbon cycle.

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Table 1. Online simulations with ModelE2-YIBs climate model

<table>
<thead>
<tr>
<th>Simulations</th>
<th>SST</th>
<th>[CO₂]</th>
<th>Emissions</th>
<th>Fires</th>
<th>O₃ effect</th>
<th>Aerosol effect</th>
</tr>
</thead>
<tbody>
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<td>F10O3</td>
<td>2010s</td>
<td>2010s</td>
<td>2010s</td>
<td>2010s</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>F10AERO</td>
<td>2010s</td>
<td>2010s</td>
<td>2010s</td>
<td>2010s</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>F10CTRL</td>
<td>2010s</td>
<td>2010s</td>
<td>2010s</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>F50O3</td>
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<td>2050s</td>
<td>2050s</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>F50AERO</td>
<td>2050s</td>
<td>2050s</td>
<td>2050s</td>
<td>2050s</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>F50CTRL</td>
<td>2050s</td>
<td>2050s</td>
<td>2050s</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
</tr>
</tbody>
</table>

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

<table>
<thead>
<tr>
<th>Species</th>
<th>Fire emissions (Tg yr(^{-1}))</th>
<th>Non-fire emissions (Tg yr(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>2010s</td>
<td>2050s</td>
</tr>
<tr>
<td>NO(_x)^a</td>
<td>0.39</td>
<td>0.74</td>
</tr>
<tr>
<td>CO</td>
<td>15.7</td>
<td>28.8</td>
</tr>
<tr>
<td>SO(_2)^a</td>
<td>0.12</td>
<td>0.22</td>
</tr>
<tr>
<td>NH(_3)</td>
<td>0.22</td>
<td>0.40</td>
</tr>
<tr>
<td>BC</td>
<td>0.08</td>
<td>0.16</td>
</tr>
<tr>
<td>OC</td>
<td>1.10</td>
<td>2.04</td>
</tr>
<tr>
<td>NM VOC</td>
<td>0.39</td>
<td>1.34</td>
</tr>
<tr>
<td>BV OC</td>
<td>N/A</td>
<td>N/A</td>
</tr>
</tbody>
</table>

^a Natural emissions are included for NO\(_x\) (lightning and soil) and SO\(_2\) (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

<table>
<thead>
<tr>
<th>Simulations</th>
<th>Base forcing</th>
<th>Temperature</th>
<th>PAR</th>
<th>Soil moisture</th>
</tr>
</thead>
<tbody>
<tr>
<td>Y10_CTRL</td>
<td>F10CTRL</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Y10_ALL</td>
<td>F10CTRL</td>
<td>F10AERO</td>
<td>F10AERO</td>
<td></td>
</tr>
<tr>
<td>Y10_TAS</td>
<td>F10CTRL</td>
<td>F10AERO</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Y10_PAR</td>
<td>F10CTRL</td>
<td></td>
<td>F10AERO</td>
<td></td>
</tr>
<tr>
<td>Y10_SLM</td>
<td>F10CTRL</td>
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<td>F10AERO</td>
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<tr>
<td>Y50_CTRL</td>
<td>F50CTRL</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Y50_ALL</td>
<td>F50CTRL</td>
<td>F50AERO</td>
<td>F50AERO</td>
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<tr>
<td>Y50_TAS</td>
<td>F50CTRL</td>
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<tr>
<td>Y50_PAR</td>
<td>F50CTRL</td>
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<td>F50AERO</td>
<td></td>
</tr>
<tr>
<td>Y50_SLM</td>
<td>F50CTRL</td>
<td></td>
<td></td>
<td>F50AERO</td>
</tr>
</tbody>
</table>
Table 4. Pearson’s correlation coefficients for GPP-PAR and GPP-AOD relationships at Ameriflux (AMF) sites

<table>
<thead>
<tr>
<th>Site</th>
<th>Period</th>
<th>GPP-PAR</th>
<th>GPP-PAR\textsubscript{dir}</th>
<th>GPP-PAR\textsubscript{dif}</th>
<th>GPP-AOD</th>
<th>AOD-PAR\textsubscript{dir}</th>
<th>AOD-PAR\textsubscript{dif}</th>
</tr>
</thead>
<tbody>
<tr>
<td>CA-Gro</td>
<td>2004-2013</td>
<td>0.19 (2432)</td>
<td>-0.01 (2432)</td>
<td>0.42 (2432)</td>
<td>0.15 (65)</td>
<td>0.60 (65)</td>
<td>-0.52 (65)</td>
</tr>
<tr>
<td>CA-Qfo</td>
<td>2003-2014</td>
<td>0.16 (3201)</td>
<td>-0.04 (3201)</td>
<td>0.45 (3201)</td>
<td>0.36 (59)</td>
<td>0.91 (34)</td>
<td>-0.80 (34)</td>
</tr>
</tbody>
</table>

\textsuperscript{a} Both GPP and PAR (direct PAR\textsubscript{dir} and diffuse PAR\textsubscript{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.

\textsuperscript{b} For CA-Gro site, diffuse PAR observations of 2005-2009 have been discarded because of poor calibration, as documented on the AMF website.
Table 5. Changes in NPP (Tg C yr\(^{-1}\)) caused by composite and individual climatic effects of fire aerosols

<table>
<thead>
<tr>
<th></th>
<th>2010s</th>
<th>2050s</th>
</tr>
</thead>
<tbody>
<tr>
<td>Online (^a)</td>
<td>72</td>
<td>-118</td>
</tr>
<tr>
<td>Offline total (^b)</td>
<td>126</td>
<td>-97</td>
</tr>
<tr>
<td>Temperature</td>
<td>11</td>
<td>-22</td>
</tr>
<tr>
<td>Radiation</td>
<td>8</td>
<td>14</td>
</tr>
<tr>
<td>Soil moisture</td>
<td>104</td>
<td>-86</td>
</tr>
</tbody>
</table>

\(^a\) 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.
References


Shindell, D. T., Lamarque, J. F., Schulz, M., Flanner, M., Jiao, C., Chin, M., Young, P. J.,
Lee, Y. H., Rotstayn, L., Mahowald, N., Milly, G., Faluvegi, G., Balkanski, Y., Collins,
W. J., Conley, A. J., Dalsoren, S., Easter, R., Ghan, S., Horowitz, L., Liu, X., Myhre, G.,
Nagashima, T., Naik, V., Rumbold, S. T., Skeie, R., Sudo, K., Szopa, S., Takemura, T.,
Voulgarakis, A., Yoon, J. H., and Lo, F.: Radiative forcing in the ACCMIP historical and
Shindell, D. T., Pechony, O., Voulgarakis, A., Faluvegi, G., Nazarenko, L., Lamarque, J. F.,
and methane chemistry in GISS-E2 historical and future climate simulations,
Atmospheric Chemistry and Physics, 13, 2653-2689, doi:10.5194/Acp-13-2653-2013,
2013b.
Sitch, S., Cox, P. M., Collins, W. J., and Huntingford, C.: Indirect radiative forcing of climate
change through ozone effects on the land-carbon sink, Nature, 448, 791-794,
Solomon, S., Qin, D., Manning, M., Chen, Z., Marquis, M., Averyt, K. B., Tignor, M., and
Miller, H. L.: Climate Change 2007: Working Group I: The Physical Science Basis,
Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA,
2007.
Spitters, C. J. T.: Separating the Diffuse and Direct Component of Global Radiation and Its
Implications for Modeling Canopy Photosynthesis .2. Calculation of Canopy
Photosynthesis, Agricultural and Forest Meteorology, 38, 231-242, doi:10.1016/0168
1923(86)90061-4, 1986.
Stocks, B. J., Mason, J. A., Todd, J. B., Bosch, E. M., Wotton, B. M., Amiro, B. D.,
Strada, S., Unger, N., and Yue, X.: Observed aerosol-induced radiative effect on plant
productivity in the eastern United States, Atmospheric Environment, 122, 463–476,
Tosca, M. G., Randerson, J. T., Zender, C. S., Flanner, M. G., and Rasch, P. J.: Do biomass
burning aerosols intensify drought in equatorial Asia during El Niño?, Atmospheric
Chemistry and Physics, 10, 3515-3528, doi:10.5194/acp-10-3515-2010, 2010.
Turetsky, M. R., Kane, E. S., Harden, J. W., Ottmar, R. D., Manies, K. L., Hoy, E., and
Kasischke, E. S.: Recent acceleration of biomass burning and carbon losses in Alaskan
Tymstra, C., Flannigan, M. D., Armitage, O. B., and Logan, K.: Impact of climate change on
area burned in Alberta's boreal forest, Int. J. Wildland Fire, 16, 153-160,
Amelynck, C., Goldstein, A., Guenther, A., Heinesch, B., Hewitt, C. N., Karl, T.,


Yue, X., and Unger, N.: Ozone vegetation damage effects on gross primary productivity in the United States, Atmospheric Chemistry and Physics, 14, 9137-9153, doi:10.5194/acp-14-9137-2014, 2014.


Figure 1. Illustration of atmospheric chemistry and physics, and biospheric processes investigated in the study. Carbonaceous aerosols from fire plumes increase diffuse light and change temperature and precipitation, influencing vegetation photosynthesis. Ozone generated photochemically from fire-emitted precursors (NOx, CO, and non-methane volatile organic compound (NMVOC)) and associated BVOC changes causes direct damage to plant photosynthesis.
Figure 2. Relationships between (b, c) GPP and diffuse PAR and (d, e) GPP and MODIS AOD at (a) two boreal sites: Groundhog River (Gro) and Quebec Mature Boreal Forest Site (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 summer days (June-August). AmeriFlux diffuse PAR and GPP (in µmol m$^{-2}$ s$^{-1}$) are half-hourly observations (10:00-14:00 LT). Instantaneous MODIS Aqua and Terra 3-km AOD are selected in a time span centered on AmeriFlux record time. For each plot: the red line indicates the regression line, black lines depict the 1-σ interval; the regression slope and correlation coefficient are both included for each site (in bold if statistically significant at 95% confidence level). Blue dots in (b, c) show instants when MODIS Aqua and Terra 3-km AODs overlap AmeriFlux data.
**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.
Figure 5. Observed (blue) and simulated (red) response of GPP to diffuse (square) and direct (triangle) PAR at boreal sites (a) CA-Gro (2004-2013) and (b) CA-Qfo (2004-2010). Observations and simulations are split into ‘diffuse’ and ‘direct’ conditions if the diffuse fraction is >0.8 and < 0.2, respectively. Data points are then averaged over PAR bins of 30 W m$^{-2}$ 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 O$_3$ damages to summer GPP in North America. Results shown are from simulations with (a, b) low and (c, d) high O$_3$ sensitivities for (a, c) 2010 and (b, d) 2050. Simulated [O$_3$] 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%.
Figure 8. Simulated summertime O₃ 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 [O₃] includes contributions from both wildfire and non-fire emissions. Excess O₃ stomatal flux is calculated as the difference between the stomatal flux and a PFT-specific threshold as defined in Sitch et al. (2007).
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.
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 \((\frac{F_{10\text{AERO}}}{F_{10\text{CTRL}}} - 1) \times 100\%\). Results for the 2050s are calculated as \((\frac{F_{50\text{AERO}}}{F_{50\text{CTRL}}} - 1)\times100\%\). Significant changes \((p<0.05)\) are marked with black dots.