

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

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Abstract

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

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52

53 **1 Introduction**

54

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
60 consequent fire emissions are accelerating carbon loss in boreal North America (Bond-
61 Lamberty et al., 2007; Turetsky et al., 2011). Meanwhile, fire-induced air pollution, including
62 ozone (O₃) and aerosols, is predicted to increase in boreal and downwind regions by
63 midcentury (Yue et al., 2013; Yue et al., 2015). Wildfire emissions have large impacts on air
64 quality (Wotawa and Trainer, 2000; Morris et al., 2006), weather/climate conditions
65 (Randerson et al., 2006; Zhao et al., 2014), and public health (Zu et al., 2016; Liu et al.,
66 2017). However, little is known about how these pollutants affect ecosystem carbon
67 assimilation, and how this impact will change with the increased wildfire activity in the
68 future.

69

70 Surface O₃ causes damages to photosynthesis through stomatal uptake (Sitch et al., 2007). In
71 the present climate state, fire-induced O₃ enhancements are predicted to reduce net primary
72 productivity (NPP) in the Amazon forest by 230 Tg C yr⁻¹ (1 Tg = 10¹² g), a magnitude
73 comparable to the direct release of CO₂ from fires in South America (Pacífico et al., 2015).
74 The aerosol effects are more uncertain because both positive and negative feedbacks occur.
75 Appearance of aerosols increases diffuse light, which is beneficial for shaded leaves in the
76 lower canopy. Consequently, photosynthesis of the whole ecosystem will increase as long as
77 the total light availability is not compromised (Kanniah et al., 2012). Rap et al. (2015)
78 estimated that biomass burning aerosols increase Amazon NPP by 78–156 Tg C yr⁻¹, which
79 offsets about half of the damage caused by fire O₃ (Pacífico et al., 2015). In contrast, strong
80 light attenuation associated with high aerosol loading may decrease canopy photosynthesis
81 (Cohan et al., 2002; Oliveira et al., 2007; Cirino et al., 2014). Furthermore, the aerosol
82 radiative effects indirectly influence ecosystem productivity through concomitant
83 meteorological perturbations that are only beginning to be examined (Yue et al., 2017).

84

85 Future wildfire activity is projected to increase over boreal North America but with large
86 uncertainties (Flannigan et al., 2005; Tymstra et al., 2007; Girardin and Mudelsee, 2008;
87 Nitschke and Innes, 2008; Amiro et al., 2009; Balshi et al., 2009; Bergeron et al., 2010;
88 Wotton et al., 2010; de Groot et al., 2013; Wang et al., 2016). For example, Amiro et al.
89 (2009) predicted an increase of 34% in Canadian area burned for a 2×CO₂ scenario (2040-
90 2060) relative to a 1×CO₂ condition (1975-1995), using the Canadian Fire Weather Index
91 (CFWI) and output from Canadian global climate model (CGCM) version 1. Balshi et al.
92 (2009) projected that area burned in boreal North America would double by the year 2045-
93 2050 relative to 1991-2000, using the Multivariate Adaptive Regression Splines (MARS)
94 approach and meteorological output from CGCM version 2. The increasing rate in Balshi et
95 al. (2009) is higher than that in Amiro et al. (2009), indicating substantial uncertainties in fire
96 projections originating from both fire models and simulated future climate. However, even
97 with the same fire models and climate change scenario, large uncertainties (in both
98 magnitude and signs) are found in the projection of area burned among individual climate
99 models (Moritz et al., 2012; Yue et al., 2013). The multi-model ensemble approach has
100 shown superior predictability over single models in historical climate simulations (Flato et al.,
101 2013) and near-term climate predictions (Kirtman et al., 2014), and has been used as a
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
105 area burned at 2046-2065 relative to 1981-2000 under the IPCC A1B scenario (Solomon et
106 al., 2007). Yue et al. (2015) further calculated that the wildfire emission increase by the
107 2050s would increase mean summertime surface O₃ by 5 ppbv in Alaska and 3 ppbv in
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
111 predicted in Yue et al. (2015).

112

113 In this study, we quantify the impacts of O₃ and aerosols emitted from boreal wildfires on the
114 land carbon uptake in North America in the present climate state and in the future world at
115 2050, taking advantage of the ensemble projection of future wildfire emissions by Yue et al.
116 (2015). The major chain we investigate includes i) generation of aerosols and surface ozone
117 from wildfire emissions and ii) impact of fire-emitted aerosols and ozone on plant

118 photosynthesis through physical and biogeochemical processes (Fig. 1). We first analyze
119 relationships between gross primary production (GPP) and aerosol optical depth (AOD) at
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
125 2050s. The impacts of the boreal fire O₃ on forest photosynthesis are predicted using the flux-
126 based damage algorithm proposed by Sitch et al. (2007), which has been fully evaluated
127 against available O₃ damage sensitivity measurements globally and over North America (Yue
128 and Unger, 2014; Yue et al., 2016; Yue et al., 2017). Fire aerosols induce perturbations to
129 radiation, meteorology, and hydrology, leading to multiple influences on the land carbon
130 uptake. Sensitivity experiments are performed using the YIBs model in offline mode to
131 isolate the contributions of changes in the individual meteorological drivers.

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133

134 **2 Materials and methods**

135

136 **2.1 Observed GPP-AOD relationships**

137

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

151 onboard the Aqua and Terra satellites (Levy et al., 2013). The MODIS 3-km AOD product
152 has been fully validated against ground-based sun photometers at both global (Remer et al.,
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
158 observations was not possible because no AERONET stations exist near to the selected AMF
159 sites.

160

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

181

182 **2.2 Wildfire emissions**

183

184 Wildfire emissions used in climate modeling are calculated as the product of area burned,
185 fuel consumption, and emission factors. To predict area burned, we build stepwise
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
191 whole year), as well as the mean and maximum values of 7 fire indexes from the CFWI
192 system during fire season. We consider the impacts of antecedent factors on current fire
193 activity by including all above variables at the same year and those in the previous two years,
194 making a total of 132 (44×3) factors. The final formats of regression are different among
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
198 Project Phase 3 (CMIP3) climate models under A1B scenario (Meehl et al., 2007) to predict
199 area burned at present day (1981-2000) and midcentury (2046-2065). In the A1B scenario,
200 CO₂ concentration is projected to 532 ppm by the year 2050, similar to the value of 541 ppm
201 in IPCC RCP8.5 scenario (van Vuuren et al., 2011) archived for the Coupled Model
202 Intercomparison Project Phase 5 (CMIP5).

203

204 We derive $1^\circ \times 1^\circ$ gridded area burned based on the prediction for each ecoregion following
205 the approach by Yue et al. (2015). Temporally, the annual area burned estimated with
206 regressions is first converted to monthly area burned using the mean seasonality for each
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 $1^\circ \times 1^\circ$
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
211 attribute 85% of area burned within each ecoregion to a number of fires with fixed size of
212 1000 ha. We then allocate these large fires among the $1^\circ \times 1^\circ$ grid cells based on the observed
213 spatial probability of large fires. For example, if one grid box (named grid 'A') bears 1% of
214 large fires (>1000 ha) within an ecoregion at present day, the same grid will bear the same
215 possibility for large fires in the future. On the other hand, fuel availability limits reburning
216 and fire spread during the forest return interval, suggesting that current burning will decrease

217 the possibility of future fires in the same location. To consider such impact, we scale the
218 observed probabilities by the fraction remaining unburned in each grid box, and then use this
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% ($1\% \times 0.8$, 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
224 gridded area burned for boreal North America by eliminating reburning issues. Sensitivity
225 tests show that specifying different area burned to the large fires (100 or 10 000 ha rather
226 than 1000 ha) yields $< 1\%$ changes in predicted biomass burned, suggesting that this
227 approach is not sensitive to the presumed fire size in the allocation procedure.

228

229 Fuel consumption, the dry mass burned per fire area, is the product of fuel load and burning
230 severity. For fuel load in Alaska, we use 1-km inventory from the US Forest Service (USFS)
231 Fuel Characteristic Classification System (FCCS, McKenzie et al., 2007). For fuel load in
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
237 boreal North America with the predicted area burned. As in Amiro et al. (2009) and Yue et al.
238 (2015), we apply constant fuel load for both present day and midcentury because opposite
239 and uncertain factors influence future projections (Kurz et al., 2008; Heyder et al., 2011;
240 Friend et al., 2014; Knorr et al., 2016; Kim et al., 2017). Instead, we consider changes in
241 burning severity due to perturbations in fuel moisture as indicated by CFWI indexes (Yue et
242 al., 2015). On average, we estimate a 9% increase in fuel consumption over boreal North
243 America by the midcentury, because higher temperature and lower precipitation result in a
244 future with drier fuel load (Flannigan et al., 2016).

245

246 Fire emissions for a specific species are then estimated as the product between biomass
247 burned and the corresponding emission factor, which is adopted from measurements by
248 Andreae and Merlet (2001) except for NO_x . We use the average value of 1.6 g NO per Kg dry
249 mass burned (DM) from six studies as NO_x emission factor, because the number of 3.0 g NO

250 per Kg DM reported in Andreae and Merlet (2001) is much higher than that of 1.1 g NO per
251 Kg DM from field observations (Alvarado et al., 2010). Based on projected area burned and
252 observation-based fuel consumption and emission factors, we derive fire emissions of NO_x,
253 carbon monoxide (CO), non-methane volatile organic compounds (NMVOCs, Alkenes and
254 Alkanes), NH₃, SO₂, black (BC) and organic carbon (OC) in the present day and midcentury.

255

256 **2.3 NASA ModelE2-YIBs model**

257

258 The NASA ModelE2-YIBs is an interactive climate-carbon-chemistry model, which couples
259 the chemistry-climate model NASA ModelE2 (Schmidt et al., 2014) and the YIBs vegetation
260 model (Yue and Unger, 2015). NASA ModelE2 is a general circulation model with
261 horizontal resolution of 2°×2.5° latitude by longitude and 40 vertical layers up to 0.1 hPa. It
262 dynamically simulates both the physical (emissions, transport, and deposition) and chemical
263 (production, conversion, and loss) processes of gas-phase chemistry (NO_x, HO_x, O_x, CO, CH₄,
264 and NMVOCs), aerosols (sulfate, nitrate, ammonium, BC, OC, dust, and sea salt), and their
265 interactions. In the model, oxidants influence the photochemical formation of secondary
266 aerosol species (e.g., sulfate, nitrate, and biogenic secondary organic aerosol), in turn,
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
270 al., 2014). The model also considers interactions between climate and atmospheric
271 components. Simulated climate affects formation, transport, and deposition of atmospheric
272 components, in turn, both O₃ and aerosols influence climate by altering radiation, temperature,
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).

277

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

283 1986). Plant respiration considers thermal dependence as well as acclimation to temperature
284 (Atkin and Tjoelker, 2003). Soil respiration is calculated based on the carbon flows among 12
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
290 responses to ambient O₃ (Yue and Unger, 2014). For this scheme, O₃ damaging level is
291 dependent on excess O₃ stomatal flux within leaves, which is a function of ambient O₃
292 concentration, boundary layer resistance, and stomatal resistance. Reduction of
293 photosynthesis is calculated on the basis of plant functional types (PFTs), each of which
294 bears a range of low-to-high sensitivities to O₃ uptake.

295

296

297 **2.4 Simulations**

298

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
301 explore the impacts of fire emissions on NPP in boreal North America (Table 1). Simulations
302 F10CTRL and F50CTRL turn off all fire emissions as well as O₃ vegetation damage for the
303 2010s and 2050s, respectively. However, climatic feedbacks of aerosols from other sources
304 (both natural and anthropogenic) and related photosynthetic responses are included.
305 Simulations F10AERO and F50AERO consider the responses of plant productivity to
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
310 twice with either low or high O₃ sensitivity. However, both of these GPP changes are not fed
311 back into the model to influence carbon allocation and tree growth. Plant respiration is
312 changing in response to meteorological perturbations, either due to climate change or aerosol
313 radiative effects. We assume no impact of O₃ damage to plant respiration and examine
314 vegetation NPP, the net carbon uptake by biosphere, for the current study. The difference
315 between AERO and CTRL runs isolates the impacts of fire aerosols on NPP, and the

316 difference between O3 and CTRL runs isolates O₃ vegetation damage caused by fire and non-
317 fire emission sources.

318
319 All simulations are conducted for 20 years and outputs for the last 15 years are used for
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
323 conditions for present-day (2010s) runs while that of 2046-2055 are used for future (2050s)
324 runs. In the RCP8.5 scenario, global average SST increases by 0.62 °C while sea ice area
325 decreases by 13.8% at the midcentury compared to the present-day level. Decadal average
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
332 vegetation model cannot simulate changes in PFT fractions. The RCP8.5 land cover change
333 dataset shows limited changes in land cover fractions between 2010s and 2050s (Oleson et al.,
334 2010). For example, relative to the 2010s, a maximum gain of 5% is predicted for grassland
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
338 2050s.

339
340 To evaluate the simulated GPP responses to changes in diffuse radiation, we perform site-
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
343 moisture) and both diffuse and direct PAR at sites CA-Gro and CA-Qfo. To isolate the
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
347 driven with variables from the online simulation of F10CTRL (Table 1). The run Y10_TAS
348 adopts the same forcing as Y10_CTRL except for temperature, which is simulated by the
349 climate simulation of F10AERO. In this case, we quantify the NPP responses to individual

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

353

354 **2.5 Observation datasets**

355

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

366

367

368 **3 Results**

369

370 **3.1 Observed GPP-AOD relationships**

371

372 Positive correlations between GPP and diffuse PAR are found at the two boreal sites (Figs
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.
376 Consequently, the slope of regression between GPP and PAR_{dif} is higher at CA-Gro than that
377 at CA-Qfo, suggesting that GPP of DBF (or MF) is more sensitive to changes in diffuse PAR
378 than that of ENF. We find almost zero correlation between GPP and PAR_{dir} at the two sites
379 (Table 4), indicating that photosynthesis is in general light-saturated for sunlit leaves at these
380 sites during boreal summer noontime. As a result, modest reductions in direct light by
381 aerosols will not decrease GPP of the whole canopy.

382

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

390

391 **3.2 Model evaluations**

392

393 Simulated summer GPP shows high values in mid-western Canada (Alberta and
394 Saskatchewan) and the Southeast (Ontario) (Fig. 3a). Forest GPP at high latitudes is low
395 because of the cool weather and light limitation there. Simulated GPP reasonably captures the
396 spatial distribution with a high correlation coefficient of 0.77 ($p \ll 0.01$) and relatively small
397 biases within 20% of the data product. Simulated AOD reproduces the observed spatial
398 pattern including the high values in boreal forests (Fig. 3b). In contrast to the MODIS
399 observations, predicted AOD is relatively uniform over the West with a background value of
400 ~ 0.1 . This discrepancy explains the low correlation coefficient ($R = 0.25$, $p < 0.01$) between
401 the model and MODIS data. The simulation fails to capture the high values in the west,
402 possibly due to a climate model underestimation of biogenic secondary organic aerosol,
403 which may be an important contribution over the western boreal forest. Simulated maximum
404 daily 8-hour average (MDA8) $[\text{O}_3]$ shows low values in boreal North America and high
405 values in the western and eastern U.S. (Fig. 4a). This pattern is consistent with surface
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.

409

410 With the Sitch et al. (2007) scheme, the YIBs model simulates reasonable GPP responses to
411 $[\text{O}_3]$ in North America (Yue and Unger, 2014; Yue et al., 2016). Generally, damage to GPP
412 increases with the enhancement of ambient $[\text{O}_3]$, but with varied sensitivities for different
413 plant species (see Fig. 6 of Yue and Unger (2014)). In responses to the same level of $[\text{O}_3]$,
414 predicted O_3 damages are higher for deciduous trees than that for needleleaf trees, consistent
415 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 half-
418 hourly) time step during summer. For 1342 pairs of GPP and PAR_{dif} at the site CA-Gro, the
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
422 PAR_{dif} . The simulated correlation is 0.61 and the regression is 0.011 at the same site. The
423 GPP sensitivity to PAR_{dif} in the model is slightly higher than that of the available
424 observations, likely because the latter are affected by additional non-meteorological abiotic
425 factors. To remove the influences of compound factors other than radiation, we follow the
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
428 GPP-to-PAR sensitivities. Increase in PAR boosts GPP, but the efficiency is much higher for
429 diffuse light than that for direct light, suggesting that increase of diffuse radiation is a benefit
430 for plant growth.

431

432 **3.3 Simulation of wildfire O_3 and aerosols**

433

434 During 1980-2009, wildfire is observed to burn 2.76×10^6 ha and 156.3 Tg DM every year
435 over boreal North America. Similarly, the ensemble prediction with fire regression models
436 estimates present-day area burned of 2.88×10^6 ha yr^{-1} and biomass burned of 160.2 Tg DM
437 yr^{-1} (Yue et al., 2015). By the midcentury, area burned is projected to increase by 77% (to
438 5.10×10^6 ha yr^{-1}) in boreal North America, mainly because of the higher temperature in
439 future fire seasons. Consequently, biomass burned increases by 93% (to 308.6 Tg DM yr^{-1})
440 because fuel consumption also increases by 9% on average in a drier climate (Yue et al.,
441 2015). Enhanced fire emissions increase concentrations of surface O_3 and column AOD,
442 especially over Alaska and central Canada (Fig. 6). The maximum centers of air pollutants
443 are collocated for O_3 and AOD but with unproportional magnitudes, suggesting non-linear
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 $7.1 \pm$
446 3.1% (2.1 ± 0.9 ppbv) to surface O_3 and $25.7 \pm 2.4\%$ (0.03 ± 0.003) to AOD in the summer
447 over boreal North America in the present day. By midcentury, these ratios increase
448 significantly to $12.8 \pm 2.8\%$ (4.2 ± 0.9 ppbv) for O_3 and $36.7 \pm 2.0\%$ (0.05 ± 0.003) for AOD.

449

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

451

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

468

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

480

481 Atmospheric circulation patterns respond to the aerosol-induced radiative perturbations (Figs
482 10c-10d). Surface radiative cooling and atmospheric heating together increase air stability
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
488 range of subsidence expands southward to 42°N (Fig. 10d) due to strengthened atmospheric
489 heating (Fig. 10b). The downward convection of warm air offsets surface radiative cooling
490 (Fig. 9b), leading to a significant warming in the Southwest (Fig. 11b). The expanded
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
499 larger maintenance and growth respiration. In the 2010s, forest NPP increases by 5-15% in
500 Alaska and southern Canada, but decreases by 5-10% in northern and eastern Canada. This
501 pattern of NPP changes (Δ NPP) is connected to the climatic effects of aerosols, especially
502 changes in soil moisture (Fig. 11). The correlation between Δ NPP (Fig. 12a) and changes in
503 soil moisture (Fig. 11e) reaches $R = 0.56$ ($n = 356$), much higher than the values of $R = -0.11$
504 for temperature change (Fig. 11a) and $R = 0.22$ for precipitation change (Fig. 11c). At the
505 continental scale, the patchy responses of NPP offset each other. Since the dominant fraction
506 of carbon uptake occurs in southern Canada (Fig. 3a), where positive NPP change is
507 predicted (Fig. 12a), wildfire aerosols enhance the total NPP by 72 Tg C yr⁻¹ in the present
508 day (Table 5). In contrast, increased wildfire emissions in the 2050s inhibit precipitation (Fig.
509 11d) and decrease soil moisture in boreal North America (Fig. 11f), leading to widespread
510 NPP reductions and a total NPP loss of 118 Tg C yr⁻¹ (Fig. 12b, Table 5).

511

512

513 **4 Discussion**

514

515 **4.1 Roles of aerosol climatic feedback**

516

517 The contrasting sign of NPP responses in the present day and midcentury are closely related
518 to the aerosol-induced surface climatic feedback. Sensitivity experiments using offline YIBs
519 model (Table 3) allowed assessment of the impacts of individual changes in the major
520 meteorological drivers, including temperature, radiation (diffuse and direct), and soil
521 moisture (Table 5). The offline simulations driven with changes in all three variables yield
522 ΔNPP of 126 Tg C yr^{-1} for the 2010s and -97 Tg C yr^{-1} for the 2050s. These values are
523 different from the online simulations, which predict ΔNPP of 72 Tg C yr^{-1} for the 2010s and -
524 118 Tg C yr^{-1} for the 2050s. Missing of other aerosol climatic feedbacks in the offline model,
525 for example, changes in relative humidity, surface pressure, soil temperature, and turbulence
526 momentum, may cause such discrepancy between the online and offline simulations.
527 Seasonal analyses show that summertime ΔNPP is 99 Tg C at present day and -95 Tg C at
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
533 ground data also show positive correlations between GPP and PAR_{dif} (Fig. 2 and Table 4),
534 and the model reproduces observed GPP responses to perturbations in direct and diffuse PAR
535 (Fig. 5). Wildfire aerosols enhance diffuse radiation by 2.6 W m^{-2} (1.7%) at present day and
536 4.0 W m^{-2} (2.3%) at midcentury in boreal North America (Fig. 9). With these changes,
537 simulated NPP increases by 8 Tg C yr^{-1} at the 2010s and 14 Tg C yr^{-1} at the 2050s (Table 5).
538 Near the two AmeriFlux sites (Fig. 2a), wildfires increase local AOD by 0.03 (Fig. 6c).
539 Meanwhile, we estimate that summer average (00:00-24:00) GPP increases by $0.04 \mu\text{mol m}^{-2}$
540 s^{-1} in the same region due to aerosol diffuse fertilization effects (DFE) based on the results of
541 $(\text{Y10_PAR} - \text{Y10_CTRL})$. This change suggests a simulated GPP sensitivity of $1.2 \mu\text{mol m}^{-2}$
542 s^{-1} (22%) per unit AOD. Observed GPP sensitivity to AOD at the two sites are 2.3 (19%) and
543 $4.5 \mu\text{mol m}^{-2} \text{ s}^{-1}$ (58%) per unit AOD, respectively (Figs 2d-2e). The absolute value of GPP
544 sensitivity from simulations is much smaller than that of observations, because the former is
545 for 24-h average while the latter is only for noontime (10:00-14:00). The relative change of
546 22% in YIBs model falls within the observed range of 19-58%.

547

548 The estimated NPP changes of 8 Tg C yr^{-1} by the radiative effects of boreal fire aerosols are
549 much weaker than the enhancement of $78\text{-}156 \text{ Tg C yr}^{-1}$ by fires in Amazon basin (Rap et al.,
550 2015). There are at least two reasons for such a difference in the DFE between boreal and
551 Amazon fire aerosols. First, wildfire emissions and associated impacts on radiation are much
552 smaller in boreal regions. Wildfires in Alaska and Canada directly emit 68 Tg C yr^{-1} at the
553 2010s, resulting in enhancement of summer AOD by 35% and diffuse radiation by 1.7%.
554 These boreal emissions are much smaller than the $\sim 240 \text{ Tg C yr}^{-1}$ in Amazon basin (van der
555 Werf et al., 2010), where fires enhances regional PM_{2.5} concentrations by 85% and diffuse
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,
558 most of NPP changes occur at high latitudes of boreal regions (Fig. 12), where total
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
561 light-saturation to light-limitation, offsetting the benefit from enhanced diffuse radiation for
562 shaded leaves. For this study, we do not find GPP reduction by the decline of direct light at
563 the two Ameriflux sites (Table 4), possibly because these sites are located at middle latitudes
564 ($<50^\circ\text{N}$). In the future, more observations at higher latitudes ($> 55^\circ\text{N}$) are required to explore
565 the sensitivity of GPP to AOD at the light-limited conditions.

566

567 Simulations have shown that absorbing aerosols can cause regional drought by increasing air
568 stability (Liu, 2005; Cook et al., 2009; Tosca et al., 2010). Our results confirm such tendency
569 but with varied range of hydrological responses depending on the magnitude of wildfire
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 ΔNPP at present day and 70.5% of
574 ΔNPP at midcentury (Table 5). These results suggest that aerosol-induced changes in soil
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;
577 Girardin et al., 2016; Chen et al., 2017).

578

579 **4.2 Limitations and uncertainties**

580

581 In this study, we examine the interactions among climate change, fire activity, air pollution,
582 and ecosystem productivity. To reduce the complexity of the interactions, we focus on the
583 most likely dominant feedback and thus main chain of events: “climate → fire → pollution →
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
586 vegetation cover/types can influence fire activity by alteration of fuel load, and air pollution
587 by BVOC emissions (climate → biosphere → 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 → pollution →
590 climate → 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
597 is not a complete assessment of all known processes and feedbacks, within these limitations,
598 this study for the first time quantifies the indirect impacts of wildfire on long-range
599 ecosystem productivity under climate change.

600

601 We use the ensemble projected fire emissions from Yue et al. (2015). Area burned is
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
604 et al., 2013; Kirtman et al., 2014), but cannot remove the possible biases in the selection of
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
612 the 2×CO₂ scenario increases only by 34% relative to the 1×CO₂ scenario. This ratio is only

613 half of the estimate in Yue et al. (2015), which compared results between periods with
614 $1.44\times\text{CO}_2$ and $1\times\text{CO}_2$. The discrepancies among these studies are more likely attributed to
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,
620 which helps retain extreme values at both the temporal and spatial domain. Compared to
621 previous estimates, Yue et al. (2015) predicted median increases in future fire emissions over
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,
627 using a dynamic global vegetation model (DGVM) and an ensemble of climate change
628 projections, Heyder et al. (2011) predicted a large-scale dieback in boreal-temperate forests
629 due to increased heat and drought stress in the coming decades. On the contrary, projections
630 using DGVMs show a widespread increase in vegetation carbon under the global warming
631 scenario with CO_2 fertilization of photosynthesis (Friend et al., 2014; Knorr et al., 2016). In
632 addition, compound factors such as greenhouse gas mitigation (Kim et al., 2017), population
633 change (Knorr et al., 2016), pine beetle outbreak (Kurz et al., 2008), and fire management
634 (Doerr and Santin, 2016) may exert varied impacts on future vegetation and fuel load.
635 Although we apply constant fuel load, we consider changes of fuel moisture because warmer
636 climate states tend to dry fuel and increase fuel consumption (Flannigan et al., 2016). With
637 constant fuel load but climate-driven fuel moisture, we calculate a 9% increase in boreal fuel
638 consumption by the midcentury (Yue et al., 2015). Although such increment is higher than
639 the prediction of 2-5% by Amiro et al. (2009) for a doubled- CO_2 climate, the consumption-
640 induced uncertainty for fire emission is likely limited because changes in area burned are
641 much more profound.

642

643 Predicted surface $[\text{O}_3]$ is much higher than observations over boreal North America (Fig. 4).
644 This bias does not affect main conclusions of this study, because predicted O_3 causes limited
645 damages to boreal GPP even with the overestimated $[\text{O}_3]$ (Fig. 7). The result confirms that

646 fire-induced O₃ vegetation damage is negligible in boreal North America. For aerosols, the
647 model captures reasonable spatial pattern of AOD but with a background value of ~0.1
648 outside fire-prone regions, where the observed AOD is usually 0.1-0.2 (Fig. 3). This
649 discrepancy may be related to the insufficient representations of physical and chemical
650 processes in the model, but may also result from the retrieval biases in MODIS data due to
651 the poor surface conditions (Liu et al., 2005) and small AOD variations (Vachon et al., 2004)
652 at high latitudes.

653

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

666

667 **4.3 Implications**

668

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

679 reduction of NPP is always accompanied with the decline of net ecosystem exchange (NEE)
680 and the enhanced carbon loss. In combination with the enhanced carbon emission of 130 Tg
681 C yr⁻¹, future boreal wildfire presents an increasing threat to the regional carbon balance and
682 global warming mitigation. Furthermore, the NPP reductions are mostly located in southern
683 Canada, where cropland is the dominant ecosystem, newly exposing the future wildfire-
684 related air pollution risk to food production.

685

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

703

704

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Table 1. Online simulations with ModelE2-YIBs climate model ^a

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

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^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

Species	Fire emissions (Tg yr ⁻¹)		Non-fire emissions (Tg yr ⁻¹)	
	2010s	2050s	2010s	2050s
NO _x ^a	0.39	0.74	2.43	2.08
CO	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

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^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).

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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

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

Site	Period ^b	Pearson’s <i>R</i>					
		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)

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^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

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^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.

777 **References**

- 778 Alvarado, M. J., Logan, J. A., Mao, J., Apel, E., Riemer, D., Blake, D., Cohen, R. C., Min, K.
779 E., Perring, A. E., Browne, E. C., Wooldridge, P. J., Diskin, G. S., Sachse, G. W.,
780 Fuelberg, H., Sessions, W. R., Harrigan, D. L., Huey, G., Liao, J., Case-Hanks, A.,
781 Jimenez, J. L., Cubison, M. J., Vay, S. A., Weinheimer, A. J., Knapp, D. J., Montzka, D.
782 D., Flocke, F. M., Pollack, I. B., Wennberg, P. O., Kurten, A., Crouse, J., St Clair, J. M.,
783 Wisthaler, A., Mikoviny, T., Yantosca, R. M., Carouge, C. C., and Le Sager, P.:
784 Nitrogen oxides and PAN in plumes from boreal fires during ARCTAS-B and their
785 impact on ozone: an integrated analysis of aircraft and satellite observations, *Atmos.*
786 *Chem. Phys.*, 10, 9739-9760, doi:10.5194/Acp-10-9739-2010, 2010.
- 787 Amiro, B. D., Cantin, A., Flannigan, M. D., and de Groot, W. J.: Future emissions from
788 Canadian boreal forest fires, *Can. J. For. Res.*, 39, 383-395, doi:10.1139/X08-154, 2009.
- 789 Andreae, M. O., and Merlet, P.: Emission of trace gases and aerosols from biomass burning,
790 *Global Biogeochemical Cycles*, 15, 955-966, 2001.
- 791 Atkin, O. K., and Tjoelker, M. G.: Thermal acclimation and the dynamic response of plant
792 respiration to temperature, *Trends in Plant Science*, 8, 343-351, doi:10.1016/S1360-
793 1385(03)00136-5, 2003.
- 794 Baker, D. F., Law, R. M., Gurney, K. R., Rayner, P., Peylin, P., Denning, A. S., Bousquet, P.,
795 Bruhwiler, L., Chen, Y. H., Ciais, P., Fung, I. Y., Heimann, M., John, J., Maki, T.,
796 Maksyutov, S., Masarie, K., Prather, M., Pak, B., Taguchi, S., and Zhu, Z.: TransCom 3
797 inversion intercomparison: Impact of transport model errors on the interannual
798 variability of regional CO₂ fluxes, 1988-2003, *Global Biogeochemical Cycles*, 20,
799 Gb1002, doi:10.1029/2004gb002439, 2006.
- 800 Ball, J. T., Woodrow, I. E., and Berry, J. A.: A model predicting stomatal conductance and its
801 contribution to the control of photosynthesis under different environmental conditions,
802 in: *Progress in Photosynthesis Research*, edited by: Biggins, J., Nijhoff, Dordrecht,
803 Netherlands, 110-112, 1987.
- 804 Balshi, M. S., McGuirez, A. D., Duffy, P., Flannigan, M., Walsh, J., and Melillo, J.:
805 Assessing the response of area burned to changing climate in western boreal North
806 America using a Multivariate Adaptive Regression Splines (MARS) approach, *Global*
807 *Change Biology*, 15, 578-600, doi:10.1111/J.1365-2486.2008.01679.X, 2009.
- 808 Beer, C., Reichstein, M., Tomelleri, E., Ciais, P., Jung, M., Carvalhais, N., Rodenbeck, C.,
809 Arain, M. A., Baldocchi, D., Bonan, G. B., Bondeau, A., Cescatti, A., Lasslop, G.,
810 Lindroth, A., Lomas, M., Luysaert, S., Margolis, H., Oleson, K. W., Rouspard, O.,
811 Veenendaal, E., Viovy, N., Williams, C., Woodward, F. I., and Papale, D.: Terrestrial
812 Gross Carbon Dioxide Uptake: Global Distribution and Covariation with Climate,
813 *Science*, 329, 834-838, doi:10.1126/Science.1184984, 2010.
- 814 Bergeron, Y., Cyr, D., Girardin, M. P., and Carcaillet, C.: Will climate change drive 21st
815 century burn rates in Canadian boreal forest outside of its natural variability: collating
816 global climate model experiments with sedimentary charcoal data, *International Journal*
817 *of Wildland Fire*, 19, 1127-1139, doi:10.1071/Wf09092, 2010.

818 Bond-Lamberty, B., Peckham, S. D., Ahl, D. E., and Gower, S. T.: Fire as the dominant
819 driver of central Canadian boreal forest carbon balance, *Nature*, 450, 89-92,
820 doi:10.1038/Nature06272, 2007.

821 Chen, L., Huang, J.-G., Alam, S. A., Zhai, L., Dawson, A., Stadt, K. J., and Comeau, P. G.:
822 Drought causes reduced growth of trembling aspen in western Canada, *Global Change*
823 *Biology*, in press, doi:10.1111/gcb.13595, 2017.

824 Cirino, G. G., Souza, R. A. F., Adams, D. K., and Artaxo, P.: The effect of atmospheric
825 aerosol particles and clouds on net ecosystem exchange in the Amazon, *Atmospheric*
826 *Chemistry and Physics*, 14, 6523-6543, doi:10.5194/acp-14-6523-2014, 2014.

827 Cohan, D. S., Xu, J., Greenwald, R., Bergin, M. H., and Chameides, W. L.: Impact of
828 atmospheric aerosol light scattering and absorption on terrestrial net primary productivity,
829 *Global Biogeochemical Cycles*, 16, 1090, doi:10.1029/2001gb001441, 2002.

830 Collins, M., Knutti, R., Arblaster, J., Dufresne, J.-L., Fichet, T., Friedlingstein, P., Gao, X.,
831 Jr., W. J. G., Johns, T., Krinner, G., Shongwe, M., Tebaldi, C., Weaver, A. J., and
832 Wehner, M.: Long-term Climate Change: Projections, Commitments and Irreversibility,
833 in: *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I*
834 *to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*,
835 edited by: Stocker, T. F., Qin, D., Plattner, G.-K., Tignor, M., Allen, S. K., Boschung, J.,
836 Nauels, A., Xia, Y., Bex, V., and Midgley, P. M., Cambridge University Press,
837 Cambridge, United Kingdom and New York, NY, USA., 1029-1136, 2013.

838 Cook, B. I., Miller, R. L., and Seager, R.: Amplification of the North American "Dust Bowl"
839 drought through human-induced land degradation, *Proceedings of the National Academy*
840 *of Sciences of the United States of America*, 106, 4997-5001,
841 doi:10.1073/pnas.0810200106, 2009.

842 Cox, P. M.: Description of the "TRIFFID" Dynamic Global Vegetation Model, Hadley
843 Centre technical note 24, 2001.

844 de Groot, W. J., Flannigan, M. D., and Cantin, A. S.: Climate change impacts on future
845 boreal fire regimes, *Forest Ecology and Management*, 294, 35-44,
846 doi:10.1016/j.foreco.2012.09.027, 2013.

847 Deng, F., Jones, D. B. A., Henze, D. K., Bousserez, N., Bowman, K. W., Fisher, J. B., Nassar,
848 R., O'Dell, C., Wunch, D., Wennberg, P. O., Kort, E. A., Wofsy, S. C., Blumenstock, T.,
849 Deutscher, N. M., Griffith, D. W. T., Hase, F., Heikkinen, P., Sherlock, V., Strong, K.,
850 Sussmann, R., and Warneke, T.: Inferring regional sources and sinks of atmospheric
851 CO₂ from GOSAT XCO₂ data, *Atmospheric Chemistry and Physics*, 14, 3703-3727,
852 doi:10.5194/acp-14-3703-2014, 2014.

853 Doerr, S. H., and Santin, C.: Global trends in wildfire and its impacts: perceptions versus
854 realities in a changing world, *Phil. Trans. R. Soc. B*, 371, doi:10.1098/rstb.2015.0345,
855 2016.

856 Duffy, P. A., Walsh, J. E., Graham, J. M., Mann, D. H., and Rupp, T. S.: Impacts of large-
857 scale atmospheric-ocean variability on Alaskan fire season severity, *Ecological*
858 *Applications*, 15, 1317-1330, doi:Doi 10.1890/04-0739, 2005.

859 Farquhar, G. D., Caemmerer, S. V., and Berry, J. A.: A Biochemical-Model of
860 Photosynthetic Co₂ Assimilation in Leaves of C-3 Species, *Planta*, 149, 78-90,
861 doi:10.1007/Bf00386231, 1980.

862 Feingold, G., Jiang, H., and Harrington, J. Y.: On smoke suppression of clouds in Amazonia,
863 *Geophysical Research Letters*, 32, L02804, doi:10.1029/2004GL021369, 2005.

864 Flannigan, M. D., Logan, K. A., Amiro, B. D., Skinner, W. R., and Stocks, B. J.: Future area
865 burned in Canada, *Clim. Change*, 72, 1-16, doi:10.1007/S10584-005-5935-Y, 2005.

866 Flannigan, M. D., Wotton, B. M., Marshall, G. A., Groot, W. J. d., Johnston, J., Jurko, N.,
867 and Cantin, A. S.: Fuel moisture sensitivity to temperature and precipitation: climate
868 change implications, *Climatic Change*, 134, 59, doi:10.1007/s10584-015-1521-0, 2016.

869 Flato, G., Marotzke, J., Abiodun, B., Braconnot, P., Chou, S. C., Collins, W., Cox, P.,
870 Driouech, F., Emori, S., Eyring, V., Forest, C., Gleckler, P., Guilyardi, E., Jakob, C.,
871 Kattsov, V., Reason, C., and Rummukainen, M.: Evaluation of Climate Models, in:
872 *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to*
873 *the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, edited
874 by: Stocker, T. F., Qin, D., Plattner, G.-K., Tignor, M., Allen, S. K., Boschung, J.,
875 Nauels, A., Xia, Y., Bex, V., and Midgley, P. M., Cambridge University Press,
876 Cambridge, United Kingdom and New York, NY, USA, 465-570, 2013.

877 Friend, A. D., Lucht, W., Rademacher, T. T., Keribin, R., Betts, R., Cadule, P., Ciais, P.,
878 Clark, D. B., Dankers, R., Falloon, P. D., Ito, A., Kahana, R., Kleidon, A., Lomas, M. R.,
879 Nishina, K., Ostberg, S., Pavlick, R., Peylin, P., Schaphoff, S., Vuichard, N.,
880 Warszawski, L., Wiltshire, A., and Woodward, F. I.: Carbon residence time dominates
881 uncertainty in terrestrial vegetation responses to future climate and atmospheric CO₂,
882 *Proc. Natl. Acad. Sci. u. s. a.*, 111, 3280-3285, doi:10.1073/pnas.1222477110, 2014.

883 Gillett, N. P., Weaver, A. J., Zwiers, F. W., and Flannigan, M. D.: Detecting the effect of
884 climate change on Canadian forest fires, *Geophys. Res. Lett.*, 31, L18211,
885 doi:10.1029/2004gl020876, 2004.

886 Girardin, M. P., and Mudelsee, M.: Past and future changes in Canadian boreal wildfire
887 activity, *Ecological Applications*, 18, 391-406, doi:Doi 10.1890/07-0747.1, 2008.

888 Girardin, M. P., Hogg, E. H., Bernier, P. Y., Kurz, W. A., Guo, X. J., and Cyr, G.: Negative
889 impacts of high temperatures on growth of black spruce forests intensify with the
890 anticipated climate warming, *Global Change Biology*, 22, 627-643,
891 doi:10.1111/gcb.13072, 2016.

892 Groot, W. J. d., D.Flannigan, M., and S.Cantin, A.: Climate change impacts on future boreal
893 fire regimes, *Forest Ecology and Management*, 294, doi:10.1016/j.foreco.2012.09.027,
894 2013.

895 Gurney, K. R., Law, R. M., Denning, A. S., Rayner, P. J., Baker, D., Bousquet, P., Bruhwiler,
896 L., Chen, Y. H., Ciais, P., Fan, S., Fung, I. Y., Gloor, M., Heimann, M., Higuchi, K.,
897 John, J., Maki, T., Maksyutov, S., Masarie, K., Peylin, P., Prather, M., Pak, B. C.,
898 Randerson, J., Sarmiento, J., Taguchi, S., Takahashi, T., and Yuen, C. W.: Towards
899 robust regional estimates of CO₂ sources and sinks using atmospheric transport models,
900 *Nature*, 415, 626-630, doi:10.1038/415626a, 2002.

901 Hansen, M. C., DeFries, R. S., Townshend, J. R. G., Carroll, M., Dimiceli, C., and Sohlberg,
902 R. A.: Global Percent Tree Cover at a Spatial Resolution of 500 Meters: First Results of
903 the MODIS Vegetation Continuous Fields Algorithm, *Earth Interactions*, 7, 1-15,
904 doi:10.1175/1087-3562(2003)007<0001:GPTCAA>2.0.CO;2, 2003.

905 Heyder, U., Schaphoff, S., Gerten, D., and Lucht, W.: Risk of severe climate change impact
906 on the terrestrial biosphere, *Environmental Research Letters*, 6, 034036,
907 doi:10.1088/1748-9326/6/3/034036, 2011.

908 Jacobson, A. R., Fletcher, S. E. M., Gruber, N., Sarmiento, J. L., and Gloor, M.: A joint
909 atmosphere-ocean inversion for surface fluxes of carbon dioxide: 2. Regional results,
910 *Global Biogeochemical Cycles*, 21, Gb1020, doi:10.1029/2006gb002703, 2007.

911 Jung, M., Reichstein, M., and Bondeau, A.: Towards global empirical upscaling of
912 FLUXNET eddy covariance observations: validation of a model tree ensemble approach
913 using a biosphere model, *Biogeosciences*, 6, 2001-2013, doi:10.5194/bg-6-2001-2009,
914 2009.

915 Kanniah, K. D., Beringer, J., North, P., and Hutley, L.: Control of atmospheric particles on
916 diffuse radiation and terrestrial plant productivity: A review, *Progress in Physical
917 Geography*, 36, 209-237, doi:10.1177/0309133311434244, 2012.

918 Kasischke, E. S., and Turetsky, M. R.: Recent changes in the fire regime across the North
919 American boreal region - Spatial and temporal patterns of burning across Canada and
920 Alaska, *Geophys. Res. Lett.*, 33, doi:10.1029/2006gl025677, 2006.

921 Keane, R. E., Agee, J. K., Fule, P., Keeley, J. E., Key, C., Kitchen, S. G., Miller, R., and
922 Schulte, L. A.: Ecological effects of large fires on US landscapes: benefit or catastrophe?,
923 *International Journal of Wildland Fire*, 17, 696-712, doi:10.1071/Wf07148, 2008.

924 Kim, J. B., Monier, E., Sohngen, B., Pitts, G. S., Drapek, R., McFarland, J., Ohrel, S., and
925 Cole, J.: Assessing climate change impacts, benefits of mitigation, and uncertainties on
926 major global forest regions under multiple socioeconomic and emissions scenarios,
927 *Environmental Research Letters*, 12, 045001, doi:10.1088/1748-9326/aa63fc, 2017.

928 Kirtman, B. P., Min, D., Infanti, J. M., Kinter, J. L., III, Paolino, D. A., Zhang, Q., van den
929 Dool, H., Saha, S., Mendez, M. P., Becker, E., Peng, P., Tripp, P., Huang, J., DeWitt, D.
930 G., Tippet, M. K., Barnston, A. G., Li, S., Rosati, A., Schubert, S. D., Rienecker, M.,
931 Suarez, M., Li, Z. E., Marshak, J., Lim, Y.-K., Tribbia, J., Pegion, K., Merryfield, W. J.,
932 Denis, B., and Wood, E. F.: The North American Multimodel Ensemble: Phase-1
933 Seasonal-to-Interannual Prediction; Phase-2 toward Developing Intraseasonal Prediction,
934 *Bulletin of the American Meteorological Society*, 95, 585-601, 2014.

935 Knorr, W., Jiang, L., and Arneeth, A.: Climate, CO2 and human population impacts on global
936 wildfire emissions, *Biogeosciences*, 13, 267-282, doi:10.5194/bg-13-267-2016, 2016.

937 Kurz, W. A., Dymond, C. C., Stinson, G., Rampley, G. J., Neilson, E. T., Carroll, A. L.,
938 Ebata, T., and Safranyik, L.: Mountain pine beetle and forest carbon feedback to climate
939 change, *Nature*, 452, 987-990, doi:10.1038/nature06777, 2008.

940 Levy, R. C., Mattoo, S., Munchak, L. A., Remer, L. A., Sayer, A. M., Patadia, F., and Hsu, N.
941 C.: The Collection 6 MODIS aerosol products over land and ocean, *Atmospheric
942 Measurement Techniques*, 6, 2989-3034, doi:10.5194/amt-6-2989-2013, 2013.

943 Liu, H. Q., Pinker, R. T., and Holben, B. N.: A global view of aerosols from merged transport
944 models, satellite, and ground observations, *J. Geophys. Res.*, 110, -, 2005.

945 Liu, J. C., Wilson, A., Mickley, L. J., Ebisu, K., Wang, Y., Sulprizio, M. P., Peng, R. D., Yue,
946 X., Son, J.-Y., Anderson, G. B., Dominici, F., and Bell, M. L.: Wildfire-specific Fine
947 Particulate Matter and Risk of Hospital Admissions in Urban and Rural Counties,
948 *Epidemiology*, 28, 77-85, doi:10.1097/EDE.0000000000000556, 2017.

949 Liu, Y.: Enhancement of the 1988 northern U.S. drought due to wildfires, *Geophysical*
950 *Research Letters*, 32, L10806, doi:10.1029/2005GL022411, 2005.

951 Liu, Y., Goodrick, S., and Heilman, W.: Wildland fire emissions, carbon, and climate:
952 Wildfire–climate interactions, *Forest Ecology and Management*, 317, 80-96,
953 doi:10.1016/j.foreco.2013.02.020, 2014.

954 Ma, Z. H., Peng, C. H., Zhu, Q. A., Chen, H., Yu, G. R., Li, W. Z., Zhou, X. L., Wang, W. F.,
955 and Zhang, W. H.: Regional drought-induced reduction in the biomass carbon sink of
956 Canada's boreal forests, *Proceedings of the National Academy of Sciences of the United*
957 *States of America*, 109, 2423-2427, doi:10.1073/pnas.1111576109, 2012.

958 McKenzie, D., Raymond, C. L., Kellogg, L. K. B., Norheim, R. A., Andreu, A. G., Bayard, A.
959 C., Kopper, K. E., and Elman, E.: Mapping fuels at multiple scales: landscape
960 application of the Fuel Characteristic Classification System, *Can. J. For. Res.*, 37, 2421-
961 2437, doi:10.1139/X07-056, 2007.

962 Meehl, G. A., Covey, C., Delworth, T., Latif, M., McAvaney, B., Mitchell, J. F. B., Stouffer,
963 R. J., and Taylor, K. E.: The WCRP CMIP3 multi-model dataset: A new era in climate
964 change research, *Bull. Am. Meteorol. Soc.*, 88, 1383-1394, doi:10.1175/BAMS-88-9-
965 1383, 2007.

966 Mercado, L. M., Bellouin, N., Sitch, S., Boucher, O., Huntingford, C., Wild, M., and Cox, P.
967 M.: Impact of changes in diffuse radiation on the global land carbon sink, *Nature*, 458,
968 1014-1017, doi:10.1038/Nature07949, 2009.

969 Moritz, M. A., Parisien, M.-A., Batllori, E., Krawchuk, M. A., Dorn, J. V., Ganz, D. J., and
970 Hayhoe, K.: Climate change and disruptions to global fire activity, *Ecosphere*, 3,
971 doi:10.1890/ES11-00345.1, 2012.

972 Morris, G. A., Hersey, S., Thompson, A. M., Pawson, S., Nielsen, J. E., Colarco, P. R.,
973 McMillan, W. W., Stohl, A., Turquety, S., Warner, J., Johnson, B. J., Kucsera, T. L.,
974 Larko, D. E., Oltmans, S. J., and Witte, J. C.: Alaskan and Canadian forest fires
975 exacerbate ozone pollution over Houston, Texas, on 19 and 20 July 2004, *Journal of*
976 *Geophysical Research-Atmospheres*, 111, D24s03, doi:10.1029/2006jd007090, 2006.

977 Munchak, L. A., Levy, R. C., Mattoo, S., Remer, L. A., Holben, B. N., Schafer, J. S.,
978 Hostetler, C. A., and Ferrare, R. A.: MODIS 3 km aerosol product: applications over
979 land in an urban/suburban region, *Atmospheric Measurement Techniques*, 6, 1747-1759,
980 doi:10.5194/amt-6-1747-2013, 2013.

981 Nadeau, L. B., McRae, D. J., and Jin, J. Z.: Development of a national fuel-type map for
982 Canada using fuzzy logic, Natural Resources Canada, Canadian Forest Service, Northern
983 Forestry Centre, Edmonton, Alberta. Information Report NOR-X-406, 2005.

984 Nitschke, C. R., and Innes, J. L.: Climatic change and fire potential in South-Central British
985 Columbia, Canada, *Global Change Biology*, 14, 841-855, doi:10.1111/j.1365-
986 2486.2007.01517.x, 2008.

987 Niyogi, D., Chang, H.-I., Saxena, V. K., Holt, T., Alapaty, K., Booker, F., Chen, F., Davis, K.
988 J., Holben, B., Matsui, T., Meyers, T., Oechel, W. C., Sr., R. A. P., Wells, R., Wilson, K.,
989 and Xue, Y.: Direct observations of the effects of aerosol loading on net ecosystem CO₂
990 exchanges over different landscapes, *Geophysical Research Letters*, 31,
991 doi:10.1029/2004GL020915, 2004.

992 Oleson, K. W., Lawrence, D. M., Bonan, G. B., Flanne, M. G., Kluzek, E., Lawrence, P. J.,
993 Levis, S., Swenson, S. C., and Thornton, P. E.: Technical Description of version 4.0 of
994 the Community Land Model (CLM), National Center for Atmospheric Research,
995 Boulder, CONCAR/TN-478+STR, 2010.

996 Oliveira, P. H. F., Artaxo, P., Pires, C., de Lucca, S., Procópio, A., Holben, B., Schafer, J.,
997 Cardoso, L. F., Wofsy, S. C., and Rocha, H. R.: The effects of biomass burning aerosols
998 and clouds on the CO₂ flux in Amazonia, *Tellus Series B-Chemical and Physical
999 Meteorology*, 59, 338-349, doi:10.1111/j.1600-0889.2007.00270.x, 2007.

1000 Pacifico, F., Folberth, G. A., Sitch, S., Haywood, J. M., Rizzo, L. V., Malavelle, F. F., and
1001 Artaxo, P.: Biomass burning related ozone damage on vegetation over the Amazon forest:
1002 a model sensitivity study, *Atmospheric Chemistry and Physics*, 15, 2791-2804,
1003 doi:10.5194/acp-15-2791-2015, 2015.

1004 Randerson, J. T., Liu, H., Flanner, M. G., Chambers, S. D., Jin, Y., Hess, P. G., Pfister, G.,
1005 Mack, M. C., Treseder, K. K., Welp, L. R., Chapin, F. S., Harden, J. W., Goulden, M. L.,
1006 Lyons, E., Neff, J. C., Schuur, E. A. G., and Zender, C. S.: The impact of boreal forest
1007 fire on climate warming, *Science*, 314, 1130-1132, doi:10.1126/Science.1132075, 2006.

1008 Rap, A., Spracklen, D. V., Mercado, L., Reddington, C. L., Haywood, J. M., Ellis, R. J.,
1009 Phillips, O. L., Artaxo, P., Bonal, D., Coupe, N. R., and Butt, N.: Fires increase Amazon
1010 forest productivity through increases in diffuse radiation, *Geophysical Research Letters*,
1011 42, 4654-4662, doi:10.1002/2015gl063719, 2015.

1012 Remer, L. A., Mattoo, S., Levy, R. C., and Munchak, L. A.: MODIS 3 km aerosol product:
1013 algorithm and global perspective, *Atmospheric Measurement Techniques*, 6, 1829-1844,
1014 doi:10.5194/amt-6-1829-2013, 2013.

1015 Rodenbeck, C., Houweling, S., Gloor, M., and Heimann, M.: CO₂ flux history 1982-2001
1016 inferred from atmospheric data using a global inversion of atmospheric transport,
1017 *Atmospheric Chemistry and Physics*, 3, 1919-1964, 2003.

1018 Schaefer, K., Collatz, G. J., Tans, P., Denning, A. S., Baker, I., Berry, J., Prihodko, L., Suits,
1019 N., and Philpott, A.: Combined Simple Biosphere/Carnegie-Ames-Stanford Approach
1020 terrestrial carbon cycle model, *J. Geophys. Res.*, 113, G03034,
1021 doi:10.1029/2007jg000603, 2008.

1022 Schmidt, G. A., Kelley, M., Nazarenko, L., Ruedy, R., Russell, G. L., Aleinov, I., Bauer, M.,
1023 Bauer, S. E., Bhat, M. K., Bleck, R., Canuto, V., Chen, Y. H., Cheng, Y., Clune, T. L.,
1024 Del Genio, A., de Fainchtein, R., Faluvegi, G., Hansen, J. E., Healy, R. J., Kiang, N. Y.,
1025 Koch, D., Lacis, A. A., LeGrande, A. N., Lerner, J., Lo, K. K., Matthews, E. E., Menon,
1026 S., Miller, R. L., Oinas, V., Oloso, A. O., Perlwitz, J. P., Puma, M. J., Putman, W. M.,

1027 Rind, D., Romanou, A., Sato, M., Shindell, D. T., Sun, S., Syed, R. A., Tausnev, N.,
1028 Tsigaridis, K., Unger, N., Voulgarakis, A., Yao, M. S., and Zhang, J. L.: Configuration
1029 and assessment of the GISS ModelE2 contributions to the CMIP5 archive, *Journal of*
1030 *Advances in Modeling Earth Systems*, 6, 141-184, doi:10.1002/2013ms000265, 2014.

1031 Shindell, D. T., Lamarque, J. F., Schulz, M., Flanner, M., Jiao, C., Chin, M., Young, P. J.,
1032 Lee, Y. H., Rotstayn, L., Mahowald, N., Milly, G., Faluvegi, G., Balkanski, Y., Collins,
1033 W. J., Conley, A. J., Dalsoren, S., Easter, R., Ghan, S., Horowitz, L., Liu, X., Myhre, G.,
1034 Nagashima, T., Naik, V., Rumbold, S. T., Skeie, R., Sudo, K., Szopa, S., Takemura, T.,
1035 Voulgarakis, A., Yoon, J. H., and Lo, F.: Radiative forcing in the ACCMIP historical and
1036 future climate simulations, *Atmospheric Chemistry and Physics*, 13, 2939-2974, doi:Doi
1037 10.5194/Acp-13-2939-2013, 2013a.

1038 Shindell, D. T., Pechony, O., Voulgarakis, A., Faluvegi, G., Nazarenko, L., Lamarque, J. F.,
1039 Bowman, K., Milly, G., Kovari, B., Ruedy, R., and Schmidt, G. A.: Interactive ozone
1040 and methane chemistry in GISS-E2 historical and future climate simulations,
1041 *Atmospheric Chemistry and Physics*, 13, 2653-2689, doi:10.5194/Acp-13-2653-2013,
1042 2013b.

1043 Sitch, S., Cox, P. M., Collins, W. J., and Huntingford, C.: Indirect radiative forcing of climate
1044 change through ozone effects on the land-carbon sink, *Nature*, 448, 791-794,
1045 doi:10.1038/Nature06059, 2007.

1046 Solomon, S., Qin, D., Manning, M., Chen, Z., Marquis, M., Averyt, K. B., Tignor, M., and
1047 Miller, H. L.: *Climate Change 2007: Working Group I: The Physical Science Basis*,
1048 Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA,
1049 2007.

1050 Spitters, C. J. T.: Separating the Diffuse and Direct Component of Global Radiation and Its
1051 Implications for Modeling Canopy Photosynthesis .2. Calculation of Canopy
1052 Photosynthesis, *Agricultural and Forest Meteorology*, 38, 231-242, doi:10.1016/0168-
1053 1923(86)90061-4, 1986.

1054 Stocks, B. J., Mason, J. A., Todd, J. B., Bosch, E. M., Wotton, B. M., Amiro, B. D.,
1055 Flannigan, M. D., Hirsch, K. G., Logan, K. A., Martell, D. L., and Skinner, W. R.: Large
1056 forest fires in Canada, 1959-1997, *Journal of Geophysical Research*, 108, 8149,
1057 doi:10.1029/2001jd000484, 2002.

1058 Strada, S., Unger, N., and Yue, X.: Observed aerosol-induced radiative effect on plant
1059 productivity in the eastern United States, *Atmospheric Environment*, 122, 463-476,
1060 doi:10.1016/j.atmosenv.2015.09.051, 2015.

1061 Tosca, M. G., Randerson, J. T., Zender, C. S., Flanner, M. G., and Rasch, P. J.: Do biomass
1062 burning aerosols intensify drought in equatorial Asia during El Niño?, *Atmospheric*
1063 *Chemistry and Physics*, 10, 3515-3528, doi:10.5194/acp-10-3515-2010, 2010.

1064 Turetsky, M. R., Kane, E. S., Harden, J. W., Ottmar, R. D., Manies, K. L., Hoy, E., and
1065 Kasischke, E. S.: Recent acceleration of biomass burning and carbon losses in Alaskan
1066 forests and peatlands, *Nature Geoscience*, 4, 27-31, doi:10.1038/Ngeo1027, 2011.

1067 Tymstra, C., Flannigan, M. D., Armitage, O. B., and Logan, K.: Impact of climate change on
1068 area burned in Alberta's boreal forest, *Int. J. Wildland Fire*, 16, 153-160,
1069 doi:10.1071/Wf06084, 2007.

1070 Unger, N., Harper, K., Zheng, Y., Kiang, N. Y., Aleinov, I., Arneth, A., Schurgers, G.,
 1071 Amelynck, C., Goldstein, A., Guenther, A., Heinesch, B., Hewitt, C. N., Karl, T.,
 1072 Laffineur, Q., Langford, B., McKinney, K. A., Misztal, P., Potosnak, M., Rinne, J.,
 1073 Pressley, S., Schoon, N., and Serça, D.: Photosynthesis-dependent isoprene emission
 1074 from leaf to planet in a global carbon–chemistry–climate model, *Atmos. Chem. Phys.*, 13,
 1075 17717-17791, doi:10.5194/acp-13-10243-2013, 2013.

1076 Vachon, F., Royer, A., Aube, M., Toubbe, B., O'Neill, N. T., and Teillet, P. M.: Remote
 1077 sensing of aerosols over North American land surfaces from POLDER and MODIS
 1078 measurements, *Atmospheric Environment*, 38, 3501-3515,
 1079 doi:10.1016/j.atmosenv.2004.01.046, 2004.

1080 Val Martin, M., Kahn, R. A., Logan, J. A., Paugam, R., Wooster, M., and Ichoku, C.: Space-
 1081 based observational constraints for 1-D plume rise models, *J. Geophys. Res.*, 117,
 1082 D22204, doi:10.1029/2012JD018370, 2012.

1083 van der Werf, G. R., Randerson, J. T., Giglio, L., Collatz, G. J., Mu, M., Kasibhatla, P. S.,
 1084 Morton, D. C., DeFries, R. S., Jin, Y., and van Leeuwen, T. T.: Global fire emissions and
 1085 the contribution of deforestation, savanna, forest, agricultural, and peat fires (1997-2009),
 1086 *Atmospheric Chemistry and Physics*, 10, 11707-11735, doi:10.5194/Acp-10-11707-2010,
 1087 2010.

1088 van Vuuren, D. P., Edmonds, J., Kainuma, M., Riahi, K., Thomson, A., Hibbard, K., Hurtt, G.
 1089 C., Kram, T., Krey, V., Lamarque, J. F., Masui, T., Meinshausen, M., Nakicenovic, N.,
 1090 Smith, S. J., and Rose, S. K.: The representative concentration pathways: an overview,
 1091 *Climatic Change*, 109, 5-31, doi:10.1007/s10584-011-0148-z, 2011.

1092 Wang, X., Thompson, D. K., Marshall, G. A., Tymstra, C., Carr, R., and Flannigan, M. D.:
 1093 Increasing frequency of extreme fire weather in Canada with climate change, *Climatic*
 1094 *Change*, 130, 573-586, doi:10.1007/s10584-015-1375-5, 2015.

1095 Wang, X. L., Parisien, M. A., Taylor, S. W., Perrakis, D. D. B., Little, J., and Flannigan, M.
 1096 D.: Future burn probability in south-central British Columbia, *International Journal of*
 1097 *Wildland Fire*, 25, 200-212, doi:10.1071/Wf15091, 2016.

1098 Wild, M., Folini, D., Schar, C., Loeb, N., Dutton, E. G., and Konig-Langlo, G.: The global
 1099 energy balance from a surface perspective, *Climate Dynamics*, 40, 3107-3134,
 1100 doi:10.1007/s00382-012-1569-8, 2013.

1101 Wittig, V. E., Ainsworth, E. A., and Long, S. P.: To what extent do current and projected
 1102 increases in surface ozone affect photosynthesis and stomatal conductance of trees? A
 1103 meta-analytic review of the last 3 decades of experiments, *Plant Cell and Environment*,
 1104 30, 1150-1162, doi:10.1111/J.1365-3040.2007.01717.X, 2007.

1105 Wotawa, G., and Trainer, M.: The influence of Canadian forest fires on pollutant
 1106 concentrations in the United States, *Science*, 288, 324-328, 2000.

1107 Wotton, B. M., Nock, C. A., and Flannigan, M. D.: Forest fire occurrence and climate change
 1108 in Canada, *International Journal of Wildland Fire*, 19, 253-271, doi:10.1071/Wf09002,
 1109 2010.

1110 Yue, X., Mickley, L. J., Logan, J. A., and Kaplan, J. O.: Ensemble projections of wildfire
 1111 activity and carbonaceous aerosol concentrations over the western United States in the

1112 mid-21st century, *Atmos. Environ.*, 77, 767-780, doi:10.1016/J.Atmosenv.2013.06.003,
1113 2013.

1114 Yue, X., and Unger, N.: Ozone vegetation damage effects on gross primary productivity in
1115 the United States, *Atmospheric Chemistry and Physics*, 14, 9137-9153, doi:10.5194/acp-
1116 14-9137-2014, 2014.

1117 Yue, X., Mickley, L. J., Logan, J. A., Hudman, R. C., Martin, M. V., and Yantosca, R. M.:
1118 Impact of 2050 climate change on North American wildfire: consequences for ozone air
1119 quality, *Atmospheric Chemistry and Physics*, 15, 10033-10055, doi:10.5194/acp-15-
1120 10033-2015, 2015.

1121 Yue, X., and Unger, N.: The Yale Interactive terrestrial Biosphere model version 1.0:
1122 description, evaluation and implementation into NASA GISS ModelE2, *Geoscientific
1123 Model Development*, 8, 2399-2417, doi:10.5194/gmd-8-2399-2015, 2015.

1124 Yue, X., Keenan, T. F., Munger, W., and Unger, N.: Limited effect of ozone reductions on
1125 the 20-year photosynthesis trend at Harvard forest, *Global Change Biology*, 22, 3750-
1126 3759, doi:10.1111/gcb.13300, 2016.

1127 Yue, X., Unger, N., Harper, K., Xia, X., Liao, H., Zhu, T., Xiao, J., Feng, Z., and Li, J.:
1128 Ozone and haze pollution weakens net primary productivity in China, *Atmospheric
1129 Chemistry and Physics*, 17, 6073-6089, doi:10.5194/acp-17-6073-2017, 2017.

1130 Zhao, Z., Kooperman, G. J., Pritchard, M. S., Russell, L. M., and Somerville, R. C. J.:
1131 Investigating impacts of forest fires in Alaska and western Canada on regional weather
1132 over the northeastern United States using CAM5 global simulations to constrain
1133 transport to a WRF-Chem regional domain, *Journal of Geophysical Research*, 119, 7515-
1134 7536, doi:10.1002/2013jd020973, 2014.

1135 Zu, K., Tao, G., Long, C., Goodman, J., and Valberg, P.: Long-range fine particulate matter
1136 from the 2002 Quebec forest fires and daily mortality in Greater Boston and New York
1137 City, *Air Quality Atmosphere and Health*, 9, 213-221, doi:10.1007/s11869-015-0332-9,
1138 2016.

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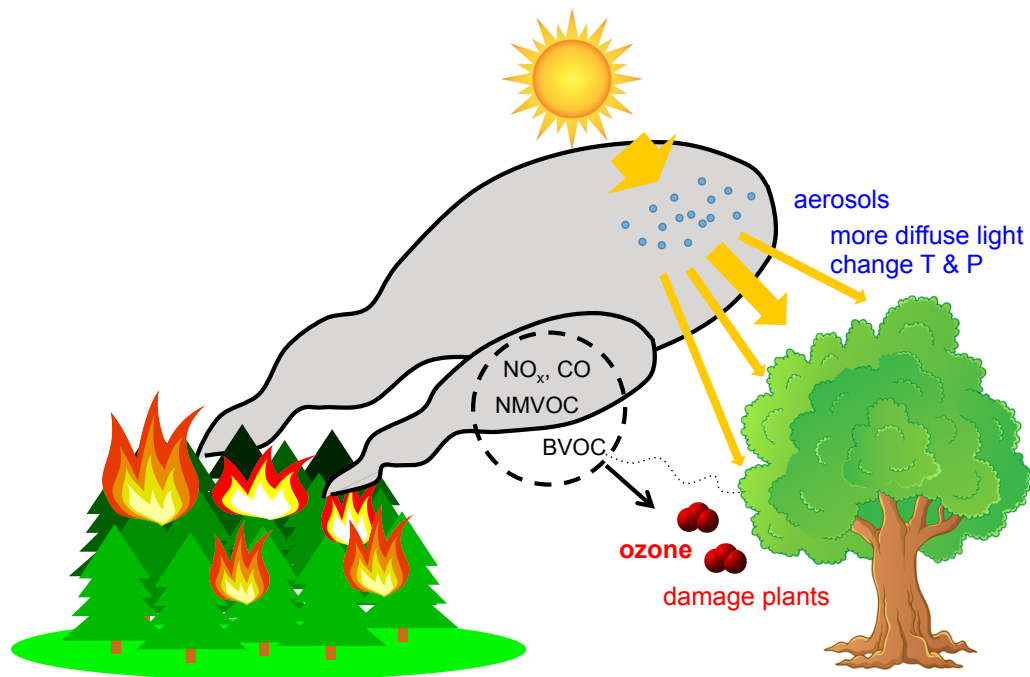
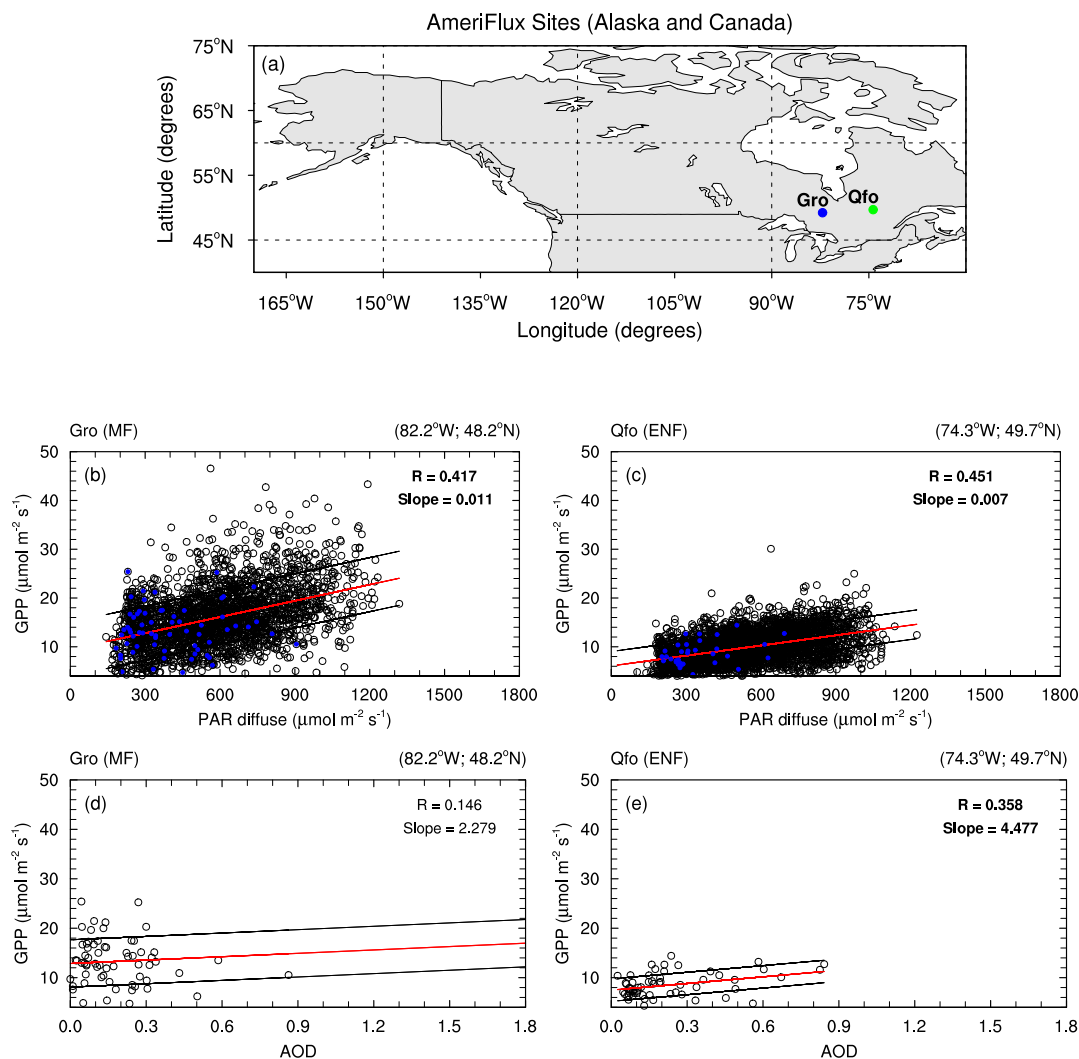


Figure 1. Illustration of atmospheric chemistry and physics, and biospheric processes investigated in the study. Carbonaceous aerosols from fire plumes increase diffuse light and change temperature and precipitation, influencing vegetation photosynthesis. Ozone generated photochemically from fire-emitted precursors (NO_x, CO, and non-methane volatile organic compound (NMVOC)) and associated BVOC changes causes direct damage to plant photosynthesis.

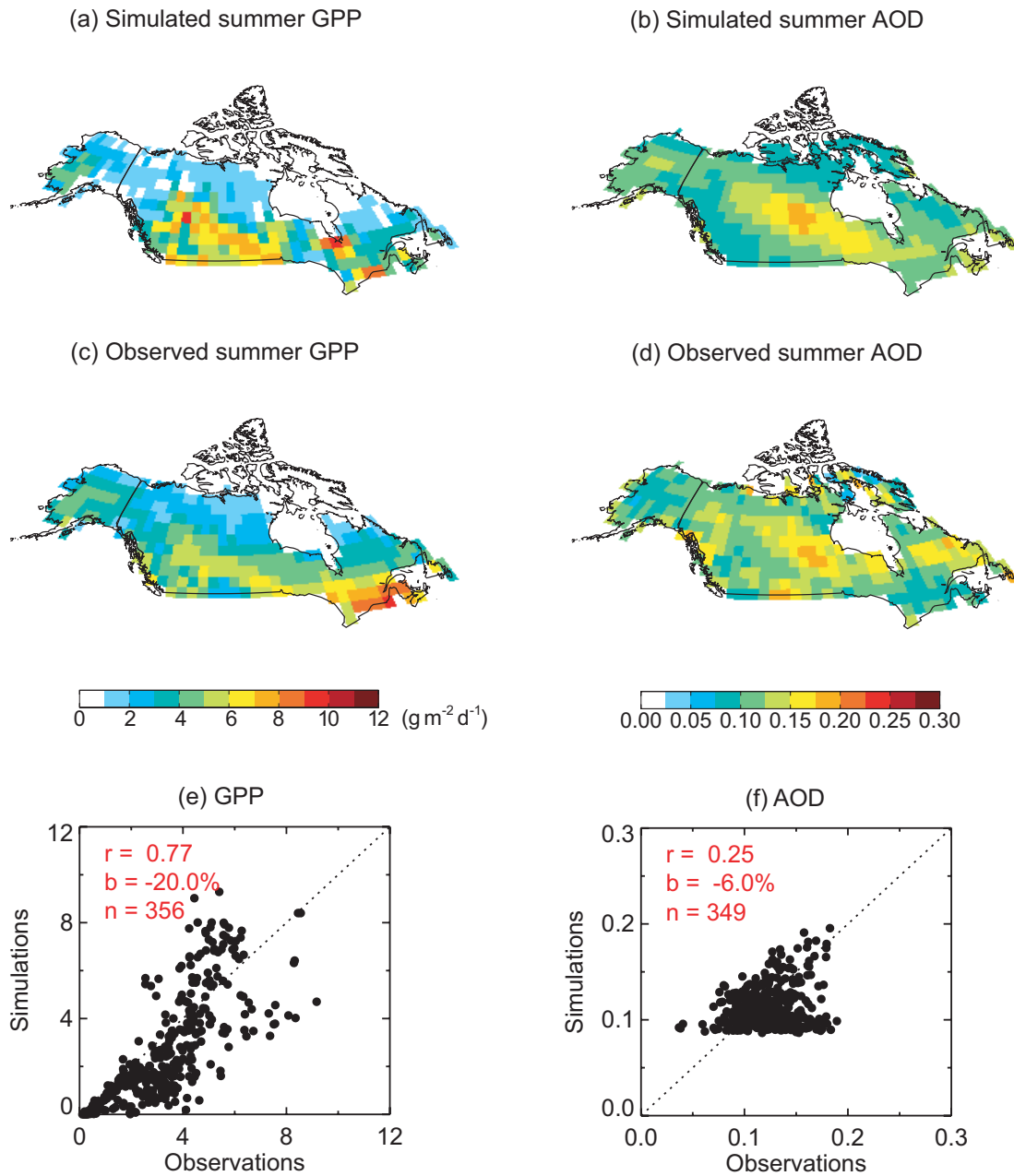


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 1174 **Figure 2.** Relationships between (b, c) GPP and diffuse PAR and (d, e) GPP and MODIS
 1175 AOD at (a) two boreal sites: Groundhog River (Gro) and Quebec Mature Boreal Forest Site
 1176 (Qfo). The two sites are from the AmeriFlux network in Canada and are dominated by mixed
 1177 forest (MF at Gro) and evergreen needleleaf forest (ENF at Qfo) (Table 1). Data cover
 1178 summer days (June-August). AmeriFlux diffuse PAR and GPP (in $\mu\text{mol m}^{-2} \text{s}^{-1}$) are half-
 1179 hourly observations (10:00-14:00 LT). Instantaneous MODIS Aqua and Terra 3-km AOD are
 1180 selected in a time span centered on AmeriFlux record time. For each plot: the red line
 1181 indicates the regression line, black lines depict the 1- σ interval; the regression slope and
 1182 correlation coefficient are both included for each site (in bold if statistically significant at 95%
 1183 confidence level). Blue dots in (b, c) show instants when MODIS Aqua and Terra 3-km
 1184 AODs overlap AmeriFlux data.

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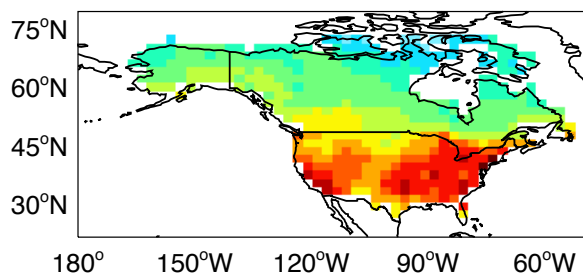


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1191 **Figure 3.** Evaluation of simulated summer (a) GPP and (b) AOD at 550 nm with (c, d)
1192 observations. Simulation results are from F10AERO (Table 1). Each point on the (e, f) scatter
1193 plot represents one grid square in boreal North America. The number of points (n),
1194 correlation coefficient (r), and relative bias (b) for the evaluation are presented on the plot.

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(a) Simulated summer MDA8 O₃



(b) Observed summer MDA8 O₃

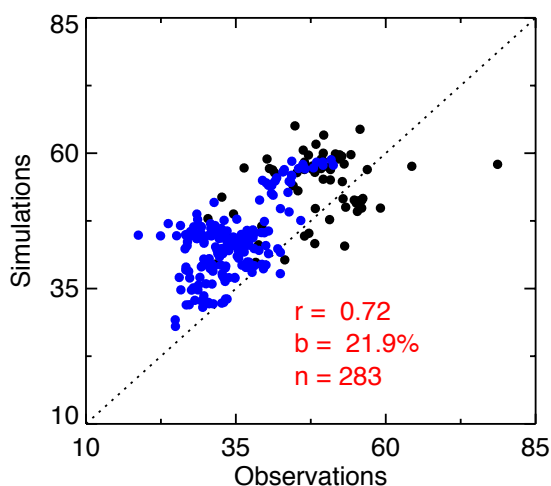
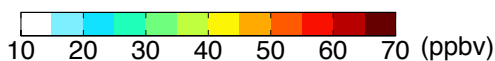
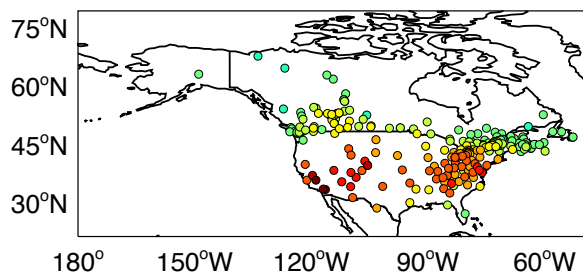
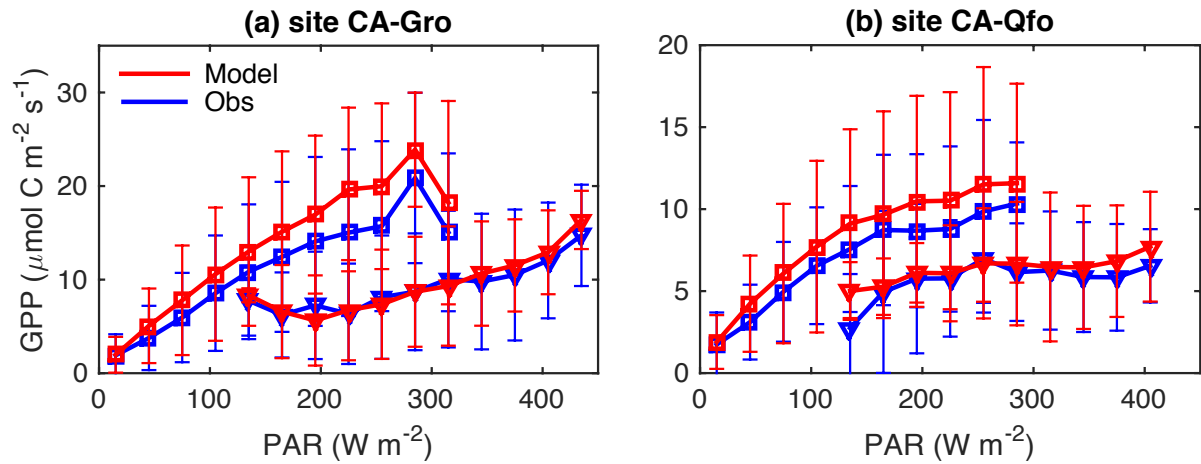


Figure 4. Evaluation of simulated summer surface maximum daily 8-hour average [O₃] with observations for 2008-2012. Observations are collected from 81 U.S. sites at the Clean Air Status and Trends Network (CASTNET) and 202 Canadian sites at the National Air Pollution Surveillance (NAPS) program. The 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.

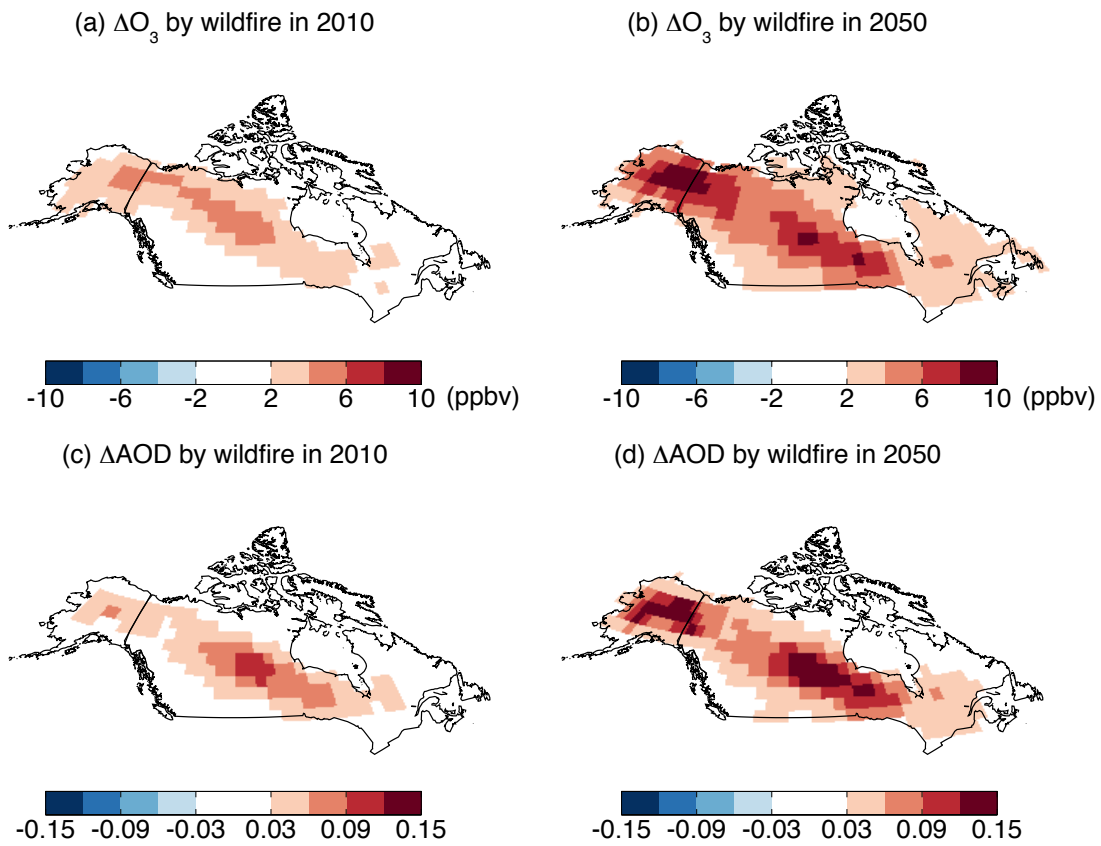
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Figure 5. Observed (blue) and simulated (red) response of GPP to diffuse (square) and direct (triangle) PAR at boreal sites (a) CA-Gro (2004-2013) and (b) CA-Qfo (2004-2010). Observations and simulations are split into ‘diffuse’ and ‘direct’ conditions 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.

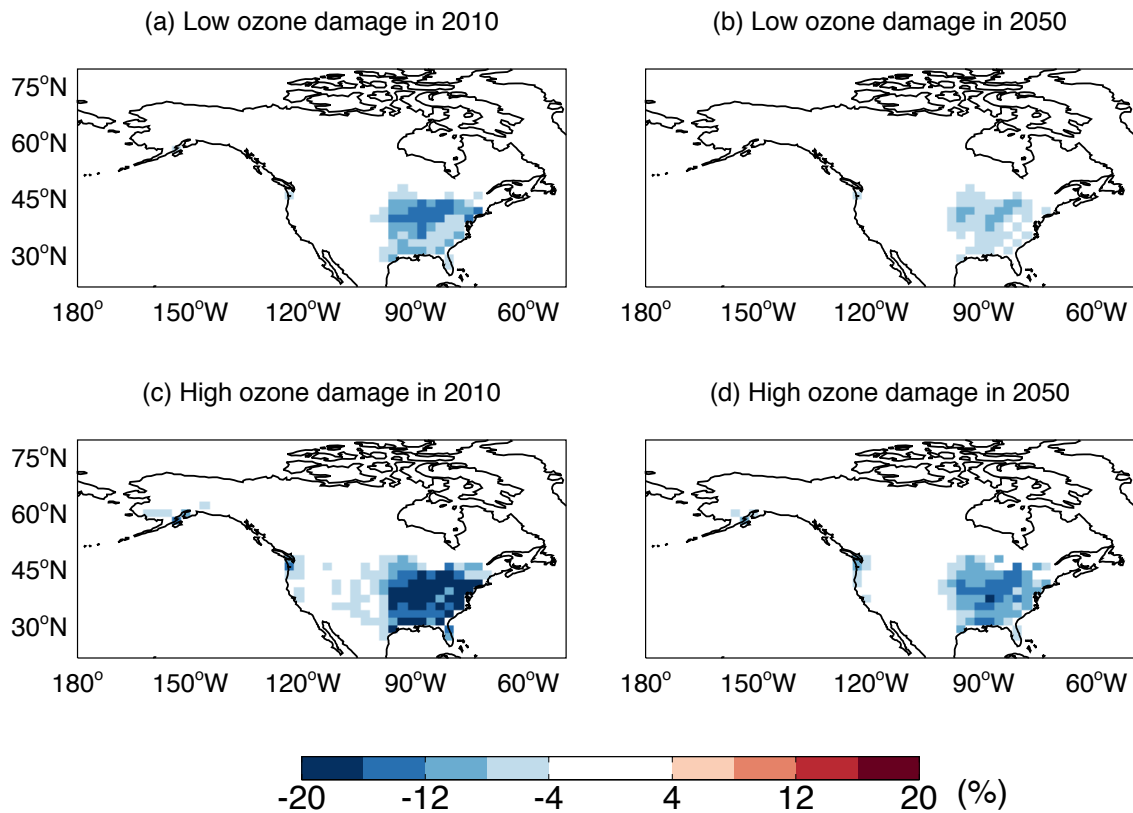
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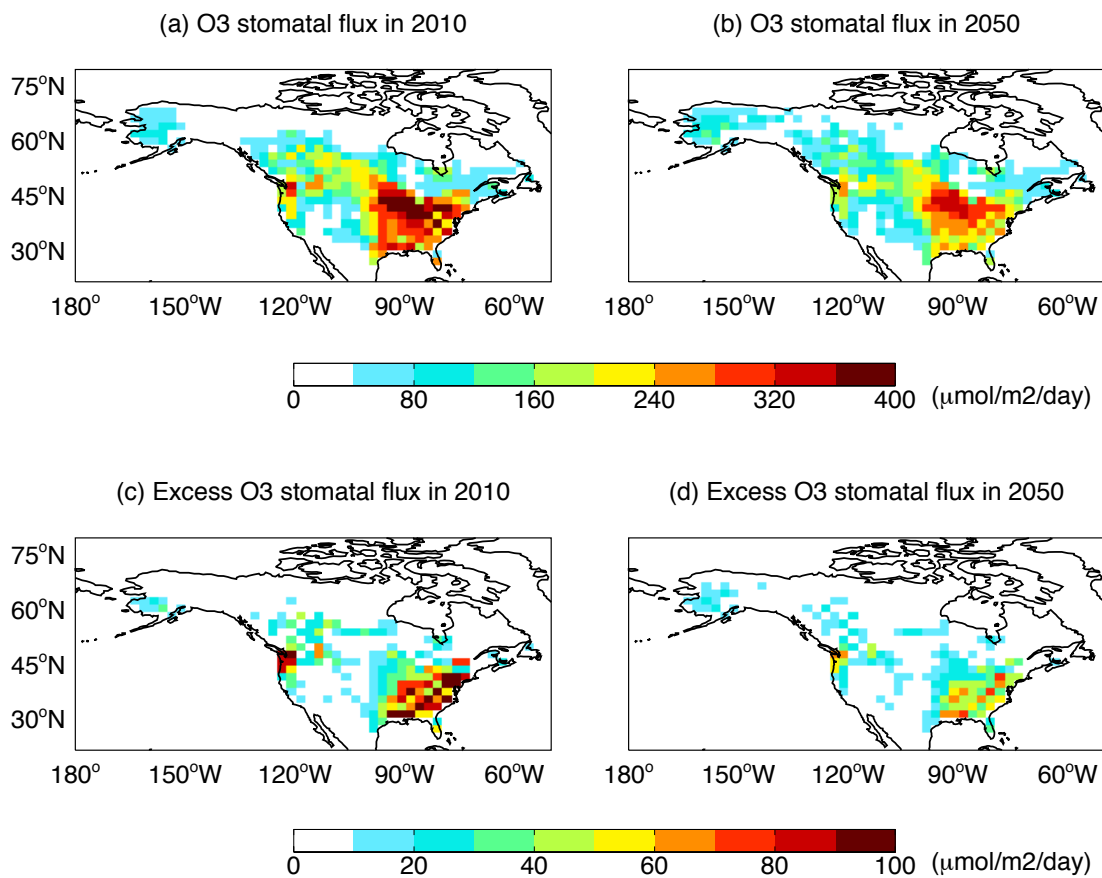
Figure 6. Changes in summer (a, b) $[O_3]$ and (c, d) AOD at 550 nm induced by wildfire emissions in (a, c) the 2010s and (b, d) the 2050s over boreal North America. Only significant changes ($p < 0.05$) are shown.

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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 $(F10O3/F10CTRL-1) \times 100\%$. Results for 2050 are derived as $(F50O3/F50CTRL-1) \times 100\%$.



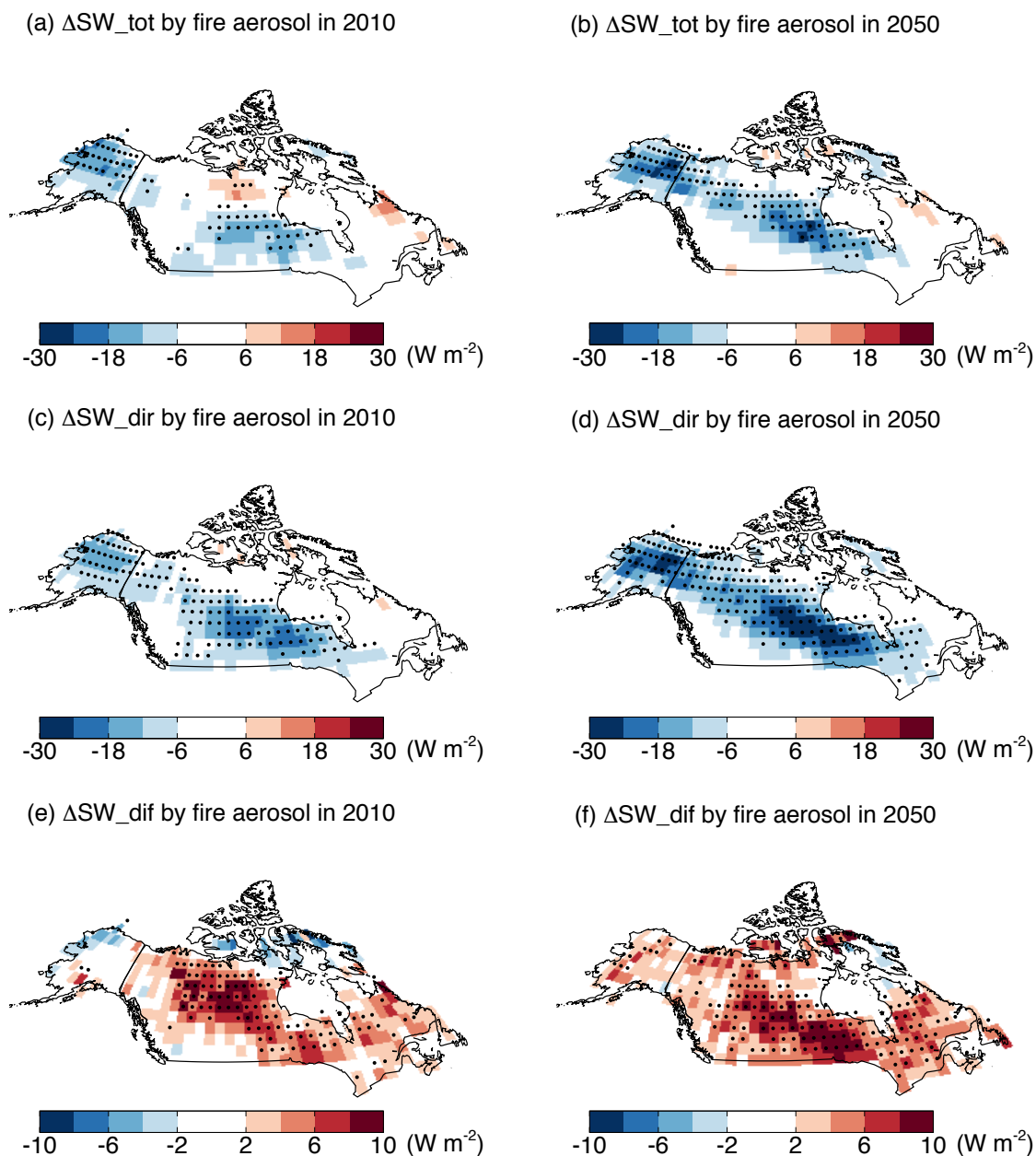
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1273 **Figure 8.** Simulated summertime O₃ stomatal fluxes in boreal North America. Results shown
 1274 are the (a, b) mean and (c, d) excess flux at (a, c) 2010 and (b, d) 2050. Simulated [O₃]
 1275 includes contributions from both wildfire and non-fire emissions. Excess O₃ stomatal flux is
 1276 calculated as the difference between the stomatal flux and a PFT-specific threshold as defined
 1277 in Sitch et al. (2007).

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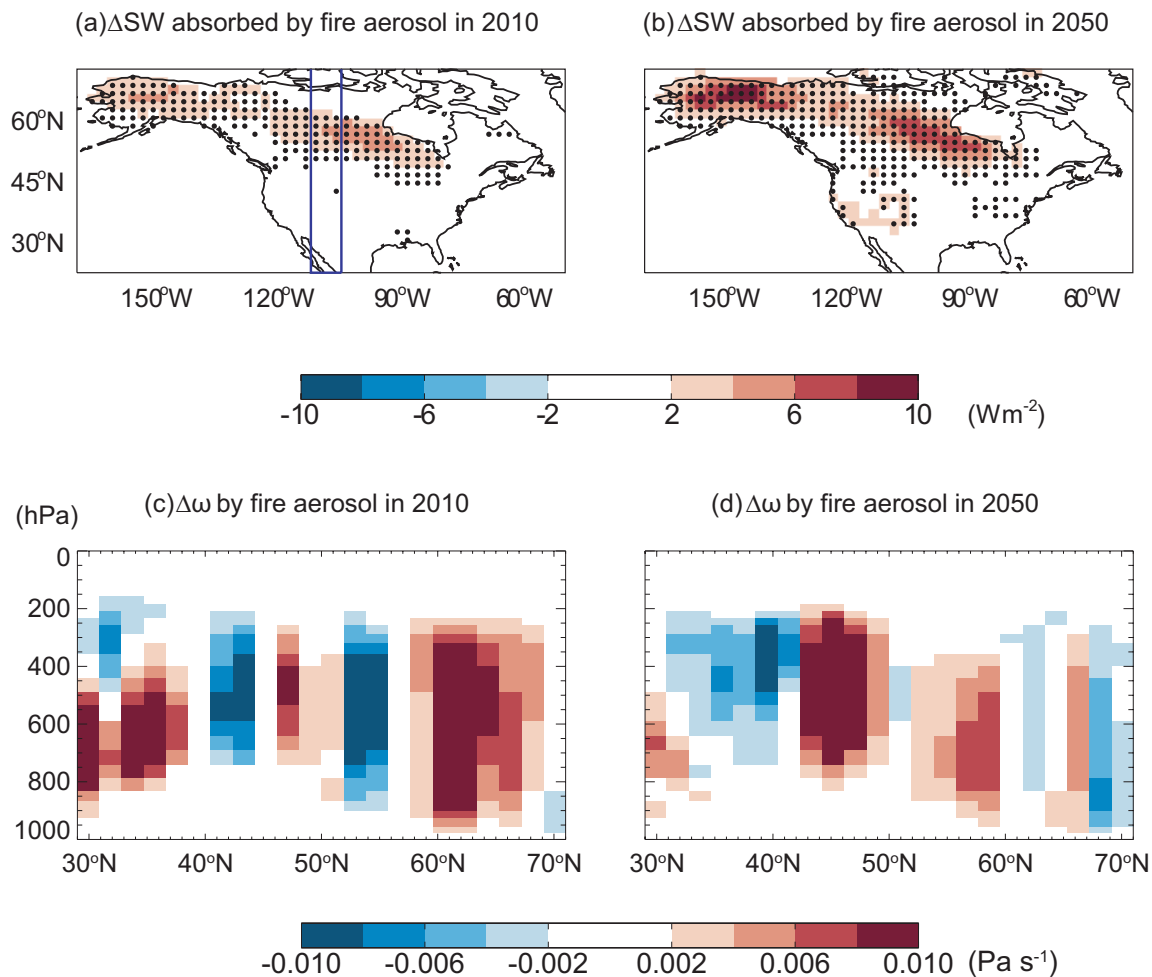
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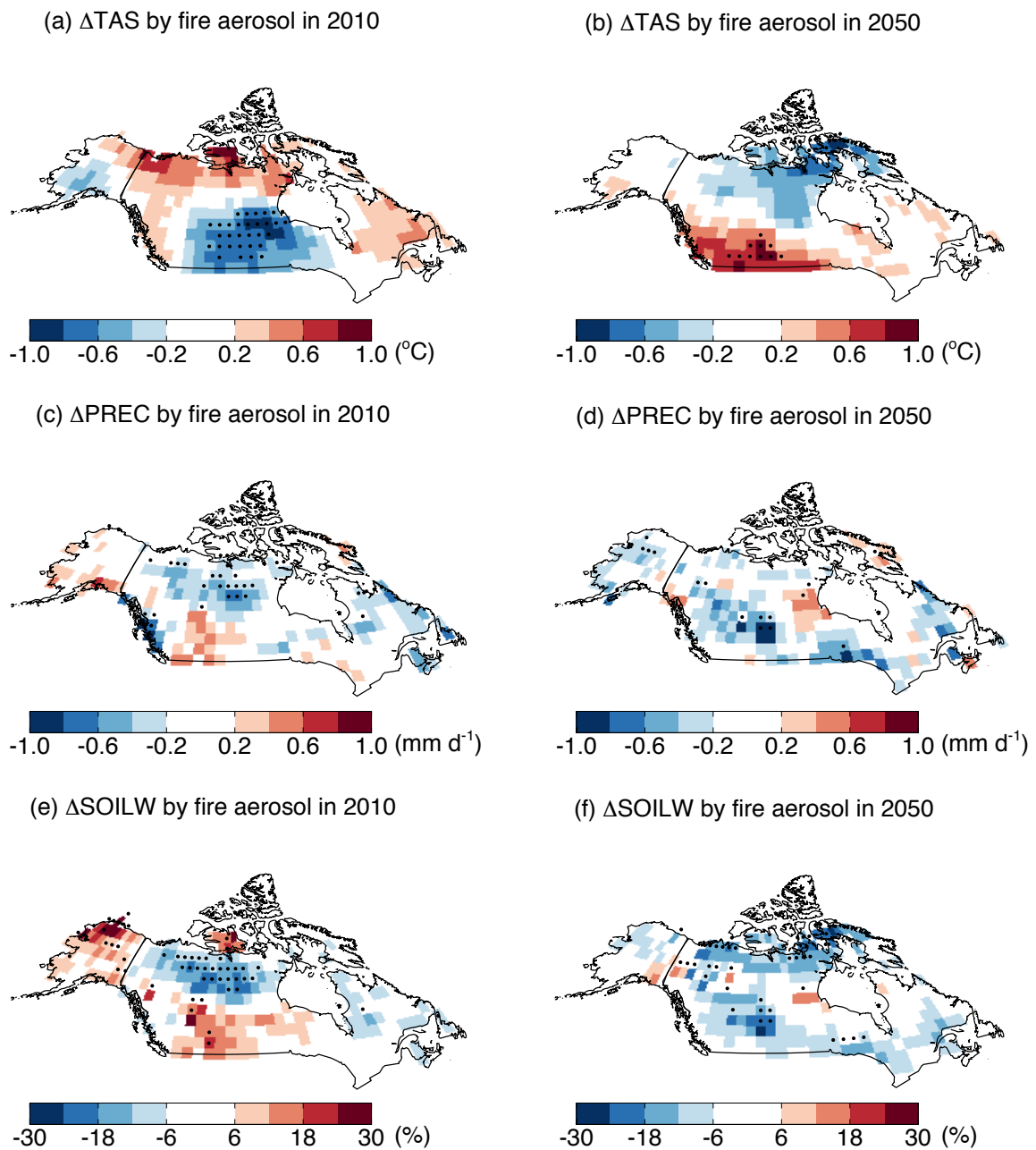
Figure 9. Changes in surface radiative fluxes induced by wildfire aerosols in boreal North America. Results shown are for the changes in summertime (June-August) (a, b) total, (c, d) direct, and (e, f) diffuse solar radiation at surface caused by aerosols from wildfire emissions at (a, c, e) present 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).

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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.

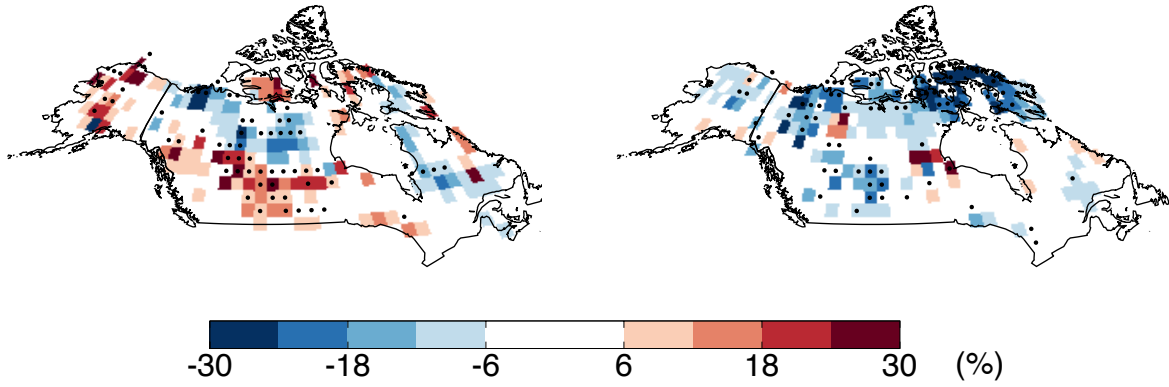


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 1307 **Figure 11.** Predicted changes in summertime (a, b) surface air temperature, (c, d)
 1308 precipitation, and (e, f) soil water content at surface caused by aerosols from wildfire
 1309 emissions at (a, c, e) present day and (b, d, f) midcentury. Results for temperature and
 1310 precipitation are shown as absolute changes. Results for soil water are shown as relative
 1311 changes. Results for the 2010s are calculated as (F10AERO - F10CTRL). Results for the
 1312 2050s are calculated as (F50AERO - F50CTRL). Significant changes ($p < 0.05$) are marked
 1313 with black dots.
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(a) Δ NPP by fire aerosol in 2010

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(b) Δ NPP by fire aerosol in 2050



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1321 **Figure 12.** Predicted percentage changes in summer NPP caused by wildfire aerosols at (a)
1322 present day and (b) midcentury. Results for the 2010s are calculated as $(F10AERO/F10CTRL$
1323 $- 1) \times 100\%$. Results for the 2050s are calculated as $(F50AERO/F50CTRL - 1) \times 100\%$.
1324 Significant changes ($p < 0.05$) are marked with black dots.

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