The impacts of modelling prescribed vs. dynamic land cover in a high CO₂ future scenario – greening of the Arctic and Amazonian dieback

Sian Kou-Giesbrecht[1], Vivek K. Arora[2], Christian Seiler[3], and Libo Wang[4]

5 [1]Department of Earth and Environmental Sciences, Dalhousie University, Halifax, NS, Canada
[2]Canadian Centre for Climate Modelling and Analysis, Climate Research Division, Environment Canada, Victoria, BC, Canada
[3]School of Environmental Studies, Queen's University, Kingston, ON, Canada
[4]Climate Processes Section, Climate Research Division, Environment and Climate Change Canada, Toronto, ON, Canada

10 *Correspondence to*: Sian Kou-Giesbrecht (sian.kougiesbrecht@dal.ca)

Abstract. Terrestrial biosphere models are a key tool in investigating the role played by the land surface in the global climate system. However, few models simulate the geographic distribution of biomes dynamically, opting to prescribe them instead using remote sensing products. While prescribing land cover still allows for the simulation of the impacts of climate change on vegetation growth as well as the impacts of land use change, it prevents the simulation of climate change-driven

15 biome shifts, with implications for projecting the future terrestrial carbon sink. Here, we isolate the impacts of prescribed vs. dynamic land cover implementations in a terrestrial biosphere model. We first introduce a framework for evaluating dynamic land cover (i.e., the spatial distribution of plant functional types across the land surface), which can be applied across terrestrial biosphere models alongside standard benchmarking of energy, water, and carbon cycle variables. After establishing confidence in simulated land cover, we then show that the simulated terrestrial carbon sink differs significantly

20 between simulations with dynamic vs. prescribed land cover for a high CO₂ future scenario. This is because of important range shifts that are only simulated when dynamic land cover is implemented: tree expansion into the Arctic and Amazonian transition from forest to grassland. In particular, the projected net land-atmosphere CO₂ flux at the end of the 21st century is twice as large in simulations with dynamic land cover than in simulations with prescribed land cover. Our results illustrate the importance of climate change-driven biome shifts for projecting the future terrestrial carbon sink.

25 ## 1 Introduction

Terrestrial biosphere models simulate the exchange of CO₂, water, and energy between the atmosphere and the land surface under both current and future climatic and socioeconomic conditions. The results from these models have led to the critical understanding that the terrestrial biosphere currently sequesters approximately a third of anthropogenic CO₂ emissions (Friedlingstein et al., 2022). Vegetation currently covers ~75% of global land area and plays a major role in regulating the

30 land-atmosphere exchange of CO₂ (Fisher and Koven, 2020). Vegetation is a dynamic component of the global climate

system that responds and feeds back to changes in environmental conditions at timescales ranging from seconds to centuries. Stomata, pores through which plants exchange CO_2 and water with the outside air, respond to changes in environmental conditions within seconds (Vialet-Chabrand et al., 2017). Over the course of a day, vegetation responds to diurnal cycles of energy. Seasonality determines annual cycles of plant phenology and interannual climate variability regulates the land-

35 atmosphere exchange of CO_2 over decades. Over decades to centuries, the geographical distribution of natural vegetation can also change as the geographical distribution of different biomes shift with long-term climate variations (Pecl et al., 2017). These changes to vegetation have critical implications for the terrestrial carbon (C) sink as well as feedbacks to the climate system. Because changes to the geographical distribution of natural vegetation occur over such long timescales, terrestrial biosphere models generally opt to prescribe the geographical distribution of natural vegetation rather than simulate it

40 dynamically. This avoids the challenges associated with accurately reproducing the geographical distribution of natural vegetation which introduces another degree of uncertainty. However, when modelling the terrestrial biosphere over longer timescales, changes to the geographical distribution of natural vegetation are magnified and become critically important (Renwick and Rocca, 2015).

Terrestrial biosphere models generally represent vegetation using a set of plant functional types (PFTs) due to the

45 intractability associated with modelling ecosystem processes for each plant species individually. This classification scheme allows terrestrial biosphere models to simplify the diversity in physiology and abiotic interactions across plant species by clustering plant species by their fundamental structure and function (Box, 1996; Reich et al., 2003). Models may choose to represent a set of PFTs that are distinguished based on leaf form (needleleaf or broadleaf), leaf phenology (evergreen or deciduous), stature (trees, grasses, or shrubs), photosynthetic pathway (C_3 or C_4), or geographical location (tropical,

50 temperate, or boreal). When a terrestrial biosphere model prescribes land cover rather than represents it dynamically, the spatial distribution of PFTs across the land surface is specified using current observations of land cover based on remote sensing products (National Research Council, 2008). Remote sensing products are first reclassified into the PFTs that a given model represents (Hartley et al., 2017). Then historical information on land use change is incorporated: crop and pasture areas have increased while natural vegetation area has correspondingly decreased over the past century (Chini et al., 2021).

55 The resulting land cover forcing allows for the simulation of how deforestation and the conversion of natural vegetation to crop and pasture, i.e., land use change, impacts net terrestrial C sequestration (Houghton et al., 2012). Because the expansion of crop and pasture areas has been greater than climate-driven changes to the geographical distribution of natural vegetation over the historical period, prescribing land cover using a forcing that only accounts for land use change is a reasonable assumption for simulations over the historical period (although historical climate-driven changes to the geographical

60 distribution of natural vegetation are not negligible (Pecl et al., 2017)). However, prescribing land cover does not capture the shifting ranges of different biomes driven by climate change and its subsequent impacts on the net land-atmosphere CO_2 flux. Changes to the geographical distribution of natural vegetation should be especially important for high CO_2 future scenarios with correspondingly strong climate warming because these scenarios exhibit the largest changes to vegetation productivity (Arora et al., 2020; Koven et al., 2022) and thus the largest potential changes to the ranges of different biomes.

65   In its simplest form, the dynamic behaviour of vegetation can be classified into two aspects: vertical and horizontal. Vertical changes in vegetation structure include changes to leaf area index, vegetation height, rooting depth, etc. given a certain spatial extent. Horizontal changes in vegetation structure describe changes in this spatial extent. Global change influences both aspects of the dynamic behaviour of vegetation: the response of plant growth to global change drivers such as CO_2 fertilisation and climate variability (Dusenge et al., 2019; Huang et al., 2018; Wu et al., 2011), as well as range shifts of

70   different PFTs. The range of a given PFT is modulated by competitive interactions for space and resources with other PFTs and by the bioclimatic limits within which a PFT can exist. Both competitive interactions and bioclimatic limits are affected by climate change (Lenoir and Svenning, 2015; Thomas, 2010; Walther, 2010). While a terrestrial biosphere model with prescribed land cover can capture how global change modulates plant growth of a given PFT within its specified range (vertical changes), it cannot capture accompanying climate change-driven range shifts of its PFTs (horizontal changes) aside

75   from specified land use change. Conversely, a terrestrial biosphere model with dynamic land cover can capture both variation in plant growth (vertical changes) as well as range shifts (horizontal changes) driven by climate change alongside specified land use change.

The Global Carbon Project produces an annual quantification of the global C budget, which includes an estimate of the current terrestrial C sink (i.e., the positive global atmosphere-land CO_2 flux) from an ensemble of terrestrial biosphere

80   models (Friedlingstein et al., 2022). In the most recent budget, only 3 out of the 16 contributing terrestrial biosphere models implemented dynamic land cover. Furthermore, only 3 out of the 11 contributing Earth System Models to the Coupled Climate–Carbon Cycle Model Intercomparison Project (C^4MIP) within the 6th phase of the Coupled Model Intercomparison Project (CMIP), which coordinates the analysis of C cycle interactions, included dynamic land cover (Arora et al., 2020). Here, we isolate the impacts of prescribed vs. dynamic land cover by comparing two versions of the Canadian Land Surface

85   Scheme Including Biogeochemical Cycles (CLASSIC): a version with prescribed land cover and a version with dynamic land cover. All other aspects of the model are identical between the two versions, including land use change, thereby isolating the effects of land cover implementation (prescribed vs. dynamic). CLASSIC has contributed to the Global Carbon Project since 2016 and CLASSIC is the land component of the Canadian Earth System Model (CanESM), which contributes to CMIP and the assessment reports of the Intergovernmental Panel on Climate Change (IPCC).

90   We first demonstrate that CLASSIC with dynamic land cover successfully reproduces the current geographical distribution of natural vegetation when compared to remote sensing products following a statistical framework, which we adapt for the first time to include an evaluation of the geographical distribution of natural vegetation. This framework can be applied across terrestrial biosphere models alongside standard benchmarking of energy, water, and carbon cycle variables. We then analyse CLASSIC simulations of Shared Socioeconomic Pathway 585 (SSP5-8.5), which describes a "fossil-fueled

95   development scenario" from 2015 to 2100, and examine how climate change could influence the future geographical distribution of natural vegetation. We identify differences between projections with prescribed land cover vs. dynamic land cover and examine how different land cover implementations impact the future terrestrial C sink.

2 Methods

2.1 CLASSIC overview

100 The Canadian Land Surface Scheme Including Biogeochemical Cycles (CLASSIC) (Melton et al., 2020; Seiler et al., 2021) is the successor to and is based on the coupled Canadian Land Surface Scheme (CLASS (Verseghy, 1991; Verseghy et al., 1993)) and the Canadian Terrestrial Ecosystem MODEL (CTEM (Arora and Boer, 2005a; Melton and Arora, 2016)). Older versions of CLASSIC (under the name CLASS-CTEM) have served as the land component in the family of Canadian Earth System Models (CanESM), including CanESM5 which contributes to CMIP (Swart et al., 2019).

105 The physical component of CLASSIC simulates fluxes of energy, momentum, and water (Verseghy, 1991; Verseghy et al., 1993). The structural attributes of vegetation are characterized by leaf area index (LAI), canopy mass, vegetation height, and rooting depth, all of which are dynamically simulated by the biogeochemical component of CLASSIC (described below). The soil profile is represented by 20 soil layers, starting with 10 soil layers of 0.1 m thickness followed by soil layers of increasing thickness up to a soil layer of 30 m thickness for a total depth of 61.4 m. The depth of permeable soil layers and

110 thus the depth to bedrock soil layers varies geographically and is specified based on the SoilGrids250m data set (Hengl et al., 2017). Soil temperature and soil moisture content (liquid and frozen) are simulated for each permeable soil layer. Where the climate permits snow to exist, the temperature, mass, density, and albedo of a single snowpack layer are simulated. The physical calculations yield net radiation, soil heat flux, latent and sensible heat fluxes, evapotranspiration, and runoff at the land-atmosphere boundary. Each grid cell is simulated independently and there are no lateral transfers between grid cells.

115 The biogeochemical component of CLASSIC simulates the land-atmosphere exchange of CO_2 via photosynthesis, autotrophic respiration, heterotrophic respiration, land use change, and fire (Arora and Boer, 2005a). CLASSIC prognostically simulates the amount of C in vegetation, litter, and soil organic matter pools for each PFT and over the bare soil fraction in each grid cell. Vegetation C is represented by leaf, stem, and root components, each of which consists of structural and non-structural carbohydrate pools. Photosynthesis generates non-structural carbohydrates which are allocated

120 between the non-structural leaf, stem, and root C pools. Autotrophic respiration occurs from the non-structural leaf, stem, and root C pools (Arora and Boer, 2005a). Non-structural C is converted to structural C within each vegetation component (Asaadi et al., 2018). Leaf, stem, and root turnover transfer C from the vegetation C pool to the litter C pool. In addition to normal leaf turnover, leaf turnover also occurs due to drought stress, cold stress, and shorter day lengths, affecting leaf phenology (Arora and Boer, 2005a). Land use change transfers C from the vegetation C pool to the litter C pool as well as

125 the land use change product C pool, whereas fire emits C to the atmosphere and also transfers C from the vegetation C pool to the litter C pool (Arora and Boer, 2005b; Arora and Melton, 2018). Decomposition transfers C from the litter C pool to the soil C pool. Finally, heterotrophic respiration occurs from both the litter and soil C pools (Melton et al., 2015). While CLASSIC does include a representation of nitrogen cycling (Asaadi and Arora, 2021; Kou-Giesbrecht and Arora, 2022), the interactions between C and nitrogen cycling are not considered in this study.

2.2 CLASSIC land cover implementation

Biogeochemical processes in CLASSIC are modelled for nine plant functional types (PFTs): needleleaf evergreen trees (NE), needleleaf deciduous trees (ND), broadleaf evergreen trees (BE), deciduous broadleaf cold trees (DBC), deciduous broadleaf dry trees (DBD), C_3 crops, C_4 crops, C_3 grasses (C3G), and C_4 grasses (C4G). These nine PFTs map directly to four PFTs used for simulating physical processes (needleleaf trees, broadleaf trees, crops, and grasses). When prescribed land cover is implanted, the time-varying fractional coverage of each PFT in each grid cell is specified by a land use forcing (described below). Land use change is the only driver of variation in the fractional coverage of PFTs over time. When crop area increases, natural vegetation area proportionally decreases. When crop area decreases, natural vegetation are proportionally increases. When dynamic land cover is implemented, the time-varying fractional coverages of C_3 crops and C_4 crops in each grid cell are specified by a land use forcing and the fractional coverages of natural PFTs in each grid cell evolve due to competition and mortality (within the non-crop area of each grid cell). For both prescribed and dynamic land cover implementations, the fractional coverages of all PFTs and bare ground in a grid cell sum to 1:

$$\sum_{n=1}^{N+1} f_n = 1 \tag{1}$$

f_n is the fractional coverage of a given natural PFT $n \in \{1, \ldots, N\}$ and f_{N+1} is the fractional coverage of bare ground in a given grid cell.

When dynamic land cover is implemented, the fractional coverage of a natural PFT is the result of colonization and mortality and is described in detail in Melton & Arora (2016) and in the Supplementary Information. Competition between natural PFTs is based on modified Lotka-Volterra equations which describe the interactions between two populations using first-order nonlinear differential equations. Here, competition is for area in a grid cell rather than population size. The colonization rate of natural PFT n (c_n; day^{-1}) is determined by its net primary productivity and leaf area index. The mortality rate of natural PFT n (m_n; day^{-1}) is the sum of intrinsic or age-related mortality, mortality due to reduced growth, mortality due to fire, and mortality when a PFT exists outside its bioclimatic limits. Intrinsic or age-related mortality is determined by its maximum age. Mortality due to reduced growth is determined by its growth rate over the previous year. Mortality due to fire is described in detail in Arora & Melton (2018). Area burned, which generates bare ground, depends on wind speed and soil moisture as well as the probability of fire occurrence (which depends on the availability of vegetation biomass as a fuel source, the combustibility of the fuel source based on soil moisture, and the presence of an ignition source based on lightning and population density) and fire suppression (which depends on population density).

Mortality when a PFT exists outside its bioclimatic limits is determined by six bioclimatic indices: air temperature of the coldest month, air temperature of the warmest month, aridity index (ratio of potential evaporation to precipitation), growing degree days (cumulative number of days with air temperature above 5°C), dry season length (number of consecutive months with precipitation less than potential evaporation), and precipitation surplus (difference between precipitation and potential evaporation). Each bioclimatic index for each grid cell is updated annually on a 25-year timescale using exponential smoothing:

$$X(t + 1) = X(t)e^{-1/25} + x(t)\left(1 - e^{-1/25}\right) \tag{2}$$

$x(t)$ represents a bioclimatic index at year t and $X(t)$ represents the smoothed bioclimatic index at year t. This accounts for time lags in the response of vegetation to climate change drivers (Wu et al., 2015). $m_{bioclim,n} = 0.25$ when any bioclimatic index is outside its corresponding bioclimatic limit for a given PFT and $m_{bioclim,n} = 0$ when all bioclimatic indices are inside their corresponding bioclimatic limits for a given PFT. Bioclimatic limits for each natural PFT are given in Table B1. These were derived by first calculating the bioclimatic indices for each grid cell using the CRUJRA climate dataset averaged over 1900 to 1920 (Harris et al., 2020). Then, the 1% and 99% quantiles of these bioclimatic indices over all grid cells for each land cover type in the Collection 5 MODIS Global Land Cover Type International Geosphere-Biosphere Programme (IGBP) product (Friedl et al., 2010) were calculated and associated with a CLASSIC PFT. Note that this assumes that the current biome ranges are in equilibrium with the 1900-1920 climate.

Each natural PFT is given a dominance rank (i) where PFTs with higher dominance ranks invade PFTs with lower dominance ranks and bare ground. Tree PFTs are given higher dominance ranks than grass PFTs. Within tree PFTs and within grass PFTs, a PFT with a higher c_n is dominant over all PFTs with lower c_n. Each PFT is thus ranked from $1, 2, \ldots, i - 1, i, i + 1, \ldots N$ where the PFT with dominance rank 1 is the most dominant. Overall, the change in fractional coverage of a PFT with dominance rank i is due to (1) its colonization into the areas of PFTs with lower dominance ranks ($i + 1, \ldots, N$) and bare ground ($N + 1$), (2) colonization by PFTs with higher dominance ranks ($i, \ldots, i - 1$) into its area, and (3) its mortality:

$$\frac{df_i}{dt} = (c_i f_{i+1} + \cdots + c_i f_N + c_i f_{N+1}) - (c_1 f_i + \cdots + c_{i-1} f_i) - m_i f_i \tag{3}$$

The change in fractional coverage of bare ground is due to mortality of the PFTs and colonization of bare ground by the PFTs:

$$\frac{df_{N+1}}{dt} = \sum_{n=1}^{N} m_n f_n - \sum_{n=1}^{N} c_n f_{N+1} \tag{4}$$

The conceptual form of Equations (3) and (4) is different from the standard Lotka-Volterra equation and allows coexistence of PFTs (Arora and Boer, 2006).

2.3 Simulation descriptions

We use CLASSIC (offline) to simulate the principal aspects of the land surface energy, water, and C cycles at the global scale over the historical period (1851 – 2020) and over the future period (2015 – 2100) for Shared Socioeconomic Pathways 585 (SSP5-8.5) at a spatial resolution of 1°. Simulations are described in Table 1.

Table 1. Description of simulations. Simulations analysed (S1-S5) are indicated for the corresponding time period. Corresponding pre-industrial spin-ups and/or historical simulations (to initialise simulations of the future period) are also indicated.

Land cover	Protocol	Pre-industrial	Historical period	Future period / SSP5-8.5

implementation		period	(1851 – 2020)	(2015 – 2100)
GLC2000 (prescribed)	TRENDY	S1 spin-up	S1	na
ESACCI (prescribed)	TRENDY	S2 spin-up	S2	na
Dynamic	TRENDY	S3 spin-up	S3	na
ESACCI (prescribed)	ISIMIP	S4 spin-up	S4 historical simulation	S4
Dynamic	ISIMIP	S5 spin-up	S5 historical simulation	S5

To evaluate dynamic land cover over the historical period (1851 – 2020), we conducted three simulations: two simulations
with prescribed land cover (S1 and S2) and a simulation with dynamic land cover (S3). These three historical simulations
were initialised from corresponding pre-industrial spin-ups (described below). Simulations with prescribed land cover use
land cover forcings derived from two remote sensing products: the Global Land Cover 2000 (GLC2000) product (S1) and
the European Space Agency Climate Change Initiative (ESACCI) product (S2). The process of generating a land cover
forcing has three steps. First, the present-day fractional coverages of the model's PFTs are obtained from a remote sensing
product, which involves reclassifying the land cover classes in the remote sensing product to the model's PFTs. A. Wang et
al. (2006) describes how the 22 land cover categories in the GLC2000 product are reclassified to the nine CLASSIC PFTs.
L. Wang et al. (2023) show how the 37 land cover categories in the ESACCI product are reclassified to the nine CLASSIC
PFTs. Second, the fractional coverages of crop PFTs are replaced with values from a land use change forcing for a given
year in the present day and the fractional coverages of natural PFTs are adjusted accordingly such that the total fractional
coverage of vegetation is unchanged in that year. Third, the timeseries of fractional coverages of crop PFTs from the land
use change forcing is incorporated by adjusting the fractional coverages of natural PFTs over a given time period relative to
the single-year land cover forcing generated in the previous step. These three steps are used to generate a land cover forcing
derived from a remote sensing product that specifies the fractional coverages of the nine CLASSIC PFTs over a given time
period. Simulations with dynamic land cover also use land cover forcings but only use the fractional coverages of crop PFTs.
S1-S3 follow the TRENDY protocol (used for contributions to the Global Carbon Project) (Friedlingstein et al., 2022). The
TRENDY protocol uses the merged monthly Climate Research Unit (CRU) and 6-hourly Japanese 55-year Reanalysis (JRA-
55) dataset from Harris et al. (2020), atmospheric CO_2 forcing from Dlugokencky and Tans (2022), population density
forcing from HYDE3.3 (Klein Goldewijk et al., 2017), and land use change forcing from the LUH2-GCB2022 (Land-Use
Harmonization 2) dataset (Chini et al., 2021; Hurtt et al., 2020; Klein Goldewijk et al., 2017). S1-S3 were initialised from
corresponding pre-industrial spin ups (Table 1). We conducted different pre-industrial spin ups for each simulation. Pre-
industrial simulations used atmospheric CO_2, land use change, and population density forcings corresponding to the year
1851. Pre-industrial simulations used meteorological forcings from 1901 to 1920 repeatedly. We ran the pre-industrial spin
up until the C pools came into equilibrium. A threshold of 0.05 Pg C yr^{-1} for the global net atmosphere-land CO_2 flux was
used to assess if equilibrium was achieved.

To examine prescribed vs. dynamic land cover over the future period (2015 – 2100) for SSP5-8.5 (Riahi et al., 2017), we conducted two simulations: one simulation with prescribed land cover from the ESACCI product (S4) and one simulation with dynamic land cover (S5). These future simulations for SSP5-8.5 were initialised from the end of corresponding historical simulations, which were in turn initialised from corresponding pre-industrial spin-ups (described below). S4 and

225 S5 use bias-corrected meteorological forcings from CanESM5. They use atmospheric CO_2, land use change, and population density forcings from CMIP and the Inter-Sectoral Impact Model Intercomparison Project (ISIMIP) (Buchner and Reyer, 2021; Warszawski et al., 2014). Our study uses a single meteorological forcing (from the corresponding model CanESM5). The effects of different meteorological forcings have been explored elsewhere (Arora et al., 2023) as have the effects of bias correction (Seiler et al., in prep). S4 and S5 were initialised from the end of corresponding historical simulations, which were

230 in turn initialised from corresponding pre-industrial spin-ups (Table 1). We conducted different pre-industrial spin ups for each simulation. Pre-industrial simulations used atmospheric CO_2, land use change, and population density forcings corresponding to the year 1851. Pre-industrial simulations used pre-industrial meteorological forcings provided by ISIMIP. We ran the pre-industrial spin up until the C pools came into equilibrium. A threshold of 0.05 Pg C yr^{-1} for the global net atmosphere-land CO_2 flux was used to assess if equilibrium was achieved.

235 **2.4 Model evaluation**

We evaluated historical simulations using the Automated Model Benchmarking R (AMBER) package developed by Seiler et al. (2021), which quantifies model performance in reproducing observation-based datasets using a skill score system that is based on ILAMB (Collier et al., 2018). Five scores compare model output to a reference dataset for a given variable, assessing the simulated time-mean bias (S_{bias}), monthly centralised root-mean-square-error (S_{rmse}), seasonality (S_{phase}), inter-

240 annual variability (S_{iav}), and spatial distribution (S_{dist}) in comparison to the observation-based dataset. Scores are dimensionless and range from 0 to 1, where higher values indicate better model performance. The overall score for each variable ($S_{overall}$) is calculated as:

$$S_{overall} = \text{mean}(S_{bias}, S_{rmse}, S_{phase}, S_{iav}, S_{dist}) \tag{5}$$

For variables with more than one observation-based dataset, we calculated a benchmark score for the observation-based

245 datasets, which quantifies how well the independently derived observation-based datasets agree with each other, thereby providing an estimate of the uncertainty of the observation-based datasets themselves. The benchmark score was calculated by iteratively comparing pairs of independently derived observation-based datasets, where one observation-based dataset is treated as model output and the other observation-dataset is treated as the reference dataset. If the two observation-based datasets were exactly the same, the benchmark score would be 1. The calculation of each score and benchmark scores is

250 described in detail in Seiler et al. (2022).

We adapted this statistical framework, which operates at a grid cell scale, to evaluate land cover fraction for each natural PFT within a grid cell. For simulations with dynamic land cover, we evaluated the land cover fraction for each natural PFT against both the GLC2000-derived land cover product and the ESACCI-derived land cover product. We compared the

GLC2000-derived land cover product and the ESACCI-derived land cover product to obtain a benchmark score that
255 quantifies the uncertainty of these land cover products. Note that a qualitative assessment of dynamic land cover at a spatial
resolution of 2.81˚ was presented in Melton & Arora (2016).

Additionally, we evaluated the standard set of energy, water, and C cycle variables following Seiler et al. (2022).
Specifically, we evaluated aboveground biomass C (AGB), surface albedo (ALBS), area burned (BURNTAREA), soil C
(CSOIL), vegetation C (CVEG), fire emissions (FIRE), gross primary productivity (GPP), soil heat flux (HFG), latent heat
260 flux (HFLS), sensible heat flux (HFSS), leaf area index (LAI), runoff (MRRO), soil moisture (MRSLL), net biome
productivity (NBP), net ecosystem exchange (NEE), ecosystem respiration (RECO), net surface long wave radiation (RLS),
net surface radiation (RNS), net surface shortwave radiation (RSS), and snow water equivalent (SNW). Observation-based
datasets used for evaluation are summarized in Table B2.

3 Results

265 ### 3.1 Historical simulations and evaluation

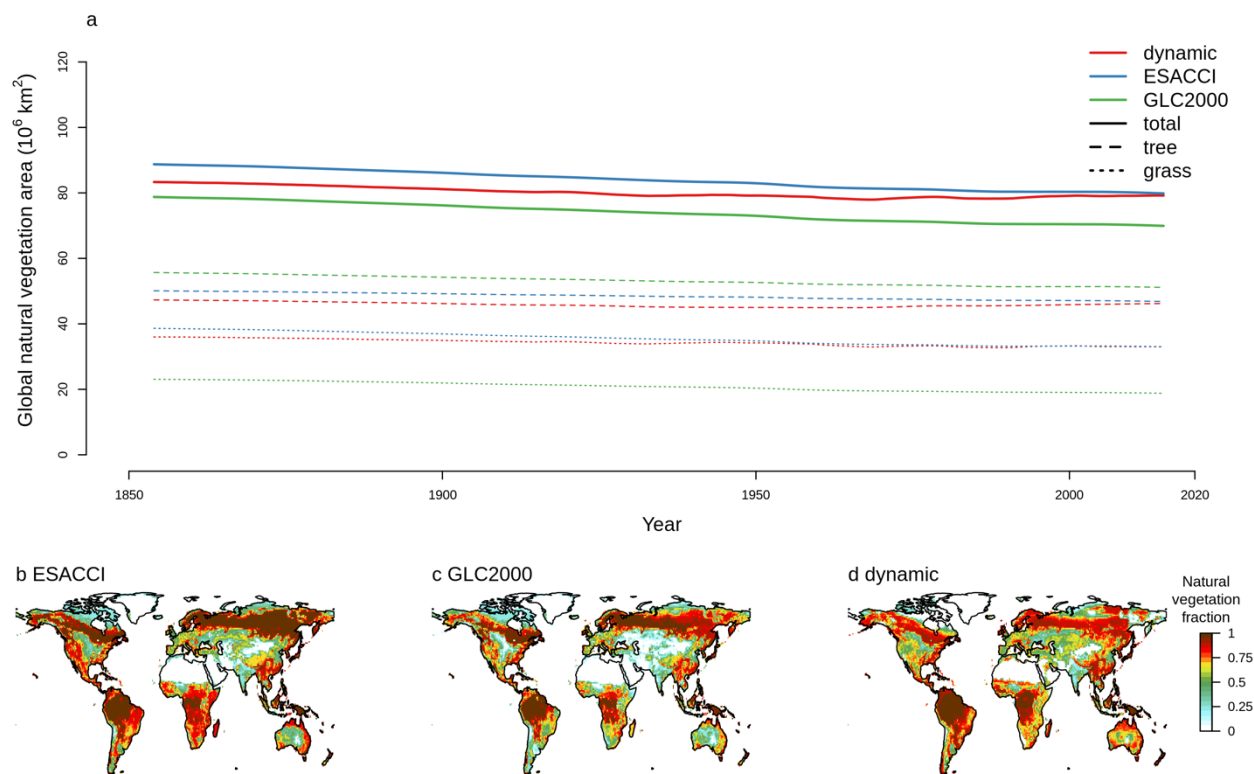
CLASSIC with dynamic land cover successfully reproduces the global area of trees, grasses, and total natural vegetation
over the historical period, falling within the range of ESACCI-derived and GLC2000-derived land cover products as shown
in Figure 1. Simulated global tree, grass, and natural vegetation areas are closer to the ESACCI-derived land cover product
than the GLC2000-derived land cover product for the present day (2000-2020). Over the historical period, simulated global
270 natural vegetation area decreases due to increasing crop area, i.e., land use change (Figure B1). Land use change is the only
driver of variation in natural vegetation area in the ESACCI-derived and GLC2000-derived land cover products. CLASSIC
with dynamic land cover simulates a smaller decrease in natural vegetation area than that attributable to land use change.
This is because decreasing natural vegetation area due to land use change is partially offset by increasing natural vegetation
area driven by stimulated plant growth from CO_2 fertilisation and climate change.

275

Figure 1. Figure 1: Natural vegetation area over the historical period. Global area of a. natural vegetation, trees, and grasses simulated by CLASSIC with dynamic land cover in comparison to ESACCI-derived and GLC2000-derived land cover products. Vegetated fraction per grid cell in the b. ESACCI land cover product, c. GLC2000 land cover product, and d. simulated by CLASSIC with dynamic land cover. Simulations and land cover products are averaged over 2000-2020 in b-d.

280 CLASSIC with dynamic land cover also broadly reproduces the latitudinal distribution of the natural PFTs in comparison to ESACCI-derived and GLC2000-derived land cover products as shown in Figure 2. However, there remain some differences between simulations and observations. CLASSIC does not simulate needleleaf evergreen trees between 20°N and 40°N whereas observations suggest that this PFT exists at these latitudes (e.g., high-elevation tropical and subtropical coniferous forests). Rather, CLASSIC simulates broadleaf deciduous cold trees at these latitudes whereas observations suggest that this

285 PFT declines between 40°N and 20°N. Furthermore, CLASSIC slightly overestimates C_4 grass area and slightly underestimates C_3 grass area in the Northern Hemisphere. Reasons for these disagreements are discussed below.

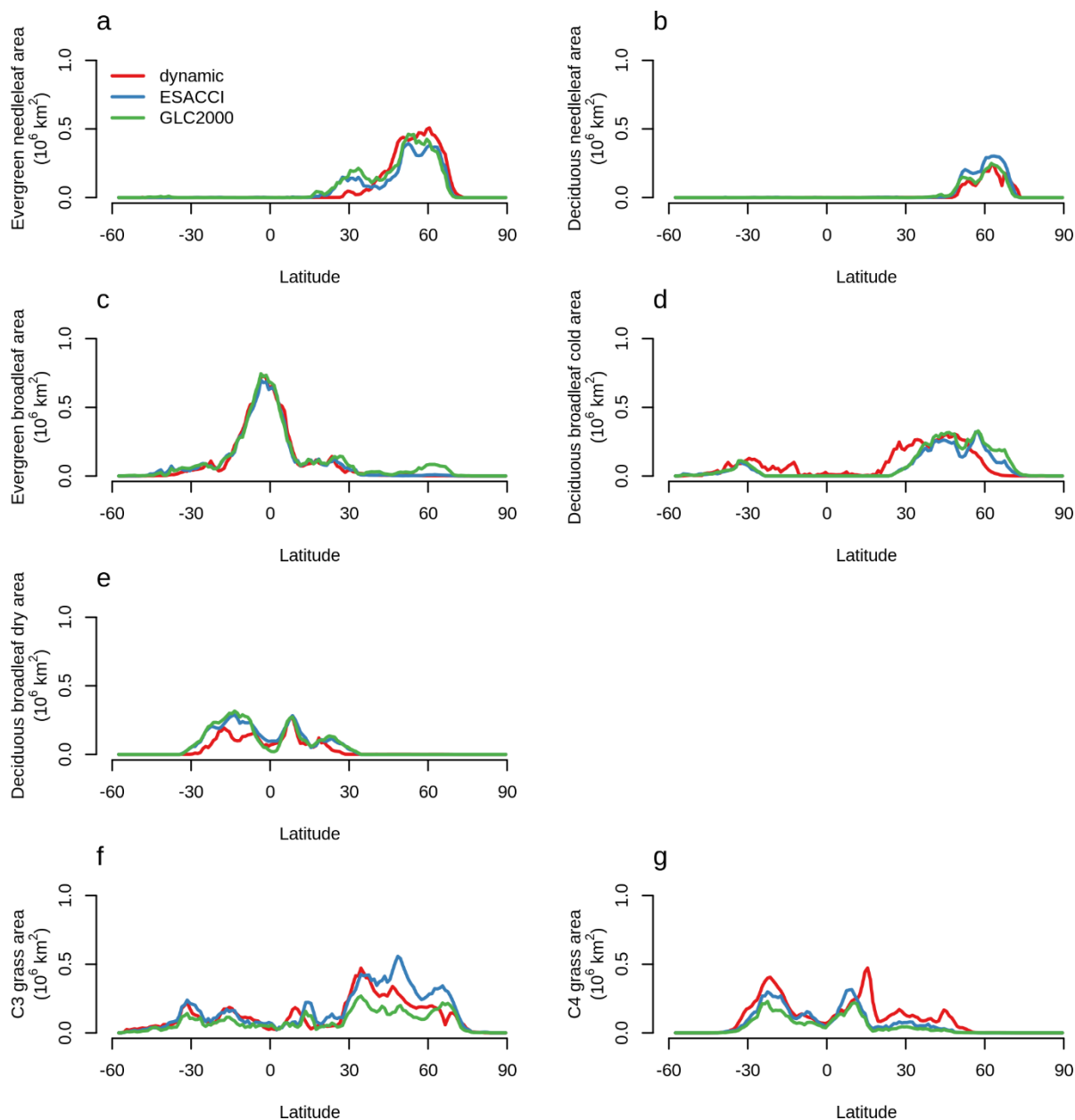



Figure 2. Latitudinal distributions of each natural plant functional type (PFT) simulated by CLASSIC with dynamic land cover in comparison to ESACCI-derived and GLC2000-derived land cover products. a. Evergreen needleleaf, b. deciduous needleleaf, c. evergreen broadleaf, d. deciduous broadleaf cold, e. deciduous broadleaf dry, f. C_3 grass, and g. C_4 grass. Simulations and land cover products are averaged over 2000-2020.

Both CLASSIC with prescribed land cover (simulated with an ESACCI-derived and GLC2000-derived land cover products) and CLASSIC with dynamic land cover reasonably capture the primary C, water, and energy cycle variables (Figure B2). The agreement between CLASSIC simulations and observations is quantified by scores of model performance in

reproducing observation-based datasets for each biogeochemical and biophysical variable. These scores indicate that CLASSIC with dynamic land cover performed similarly to CLASSIC with prescribed land cover (Figure B3). Figures 3a and 3b show model scores (averaged across all CLASSIC simulations, with both prescribed and dynamic land cover) plotted against benchmark scores for biogeochemical and biophysical variables, respectively. Figure 3 shows the correspondence between model performance in reproducing observation-based datasets (i.e., model scores) and the uncertainty of the observation-based datasets themselves (i.e., benchmark scores). Model scores for both versions of CLASSIC and benchmark scores are similar, falling close to the 1:1 line, which indicates that model performance is comparable to the uncertainty of the observation-based datasets themselves. Both versions of CLASSIC perform slightly better for biophysical variables (Figure 3b) than for biogeochemical variables (Figure 3a). Finally, both CLASSIC with prescribed and dynamic land cover simulate the terrestrial C sink over the historical period in comparison to other models in the Global Carbon Project (Friedlingstein et al., 2022) (Figure B4).

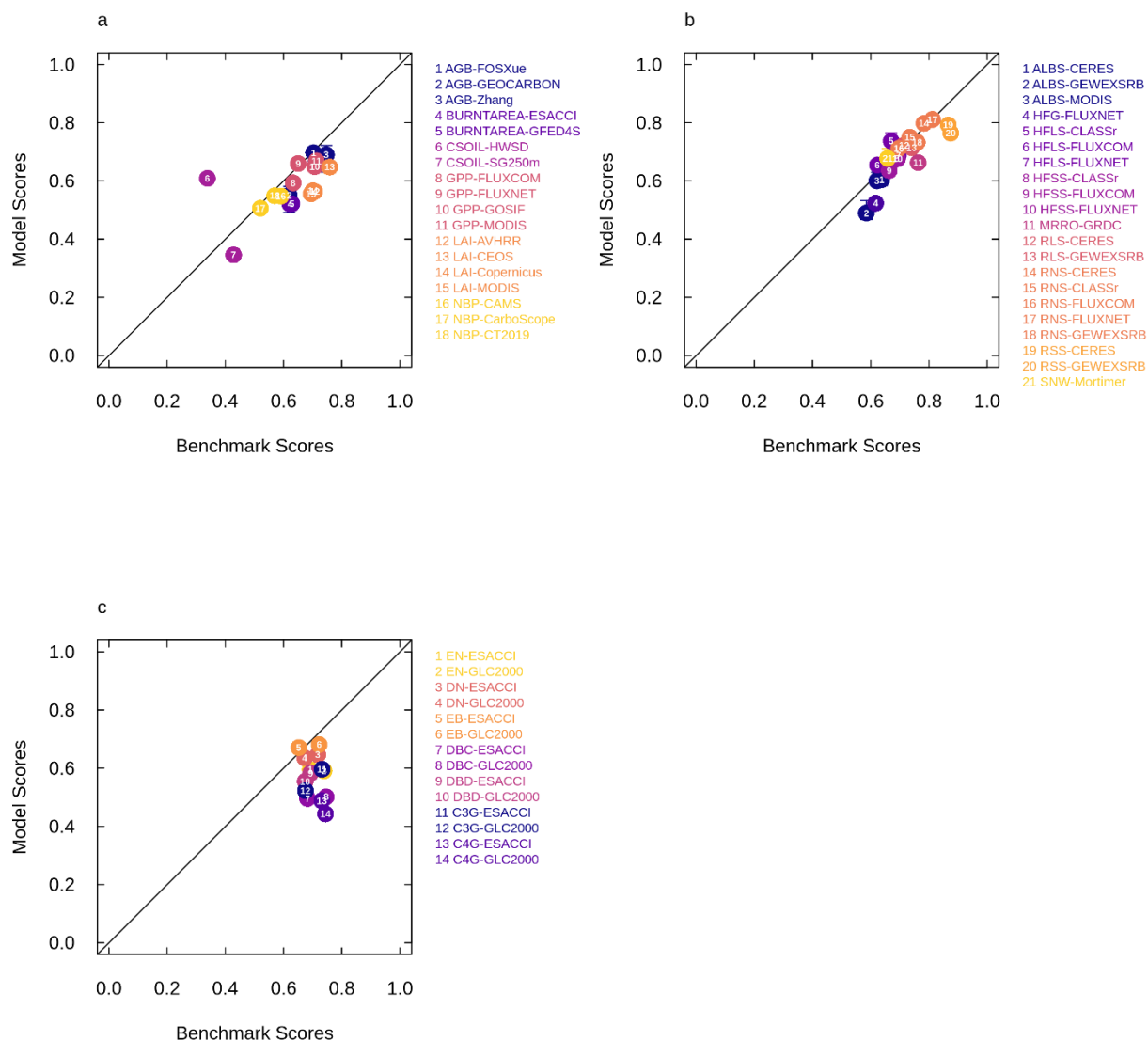



Figure 3. Model performance in reproducing observation-based datasets (model scores) relative to the uncertainty of the observation-based datasets themselves (benchmark scores) for a. biogeochemical variables, b. biophysical variables, and c. land cover fractions of each natural plant functional type (PFT). Model scores are averaged across simulations of CLASSIC with prescribed land cover (with an ESACCI-derived land cover forcing and a GLC2000-derived land cover forcing) and dynamic land cover for biogeochemical and biophysical variables. Whiskers indicate the maximum and minimum model scores. Abbreviations for C, water, and energy variables and observation-based datasets are described in the Methods. Abbreviations for PFTs are evergreen needleleaf (EN), deciduous needleleaf (DN), evergreen broadleaf (EB), deciduous broadleaf cold (DBC), deciduous broadleaf dry (DBD), C$_3$ grass (C3G), and C$_4$ grass (C4G).

310

315   When dynamic land cover is implemented in CLASSIC, the ability of CLASSIC to reproduce land cover fractions of each natural PFT can be assessed alongside C, water, and energy variables as shown in Figure 3c. Additionally, benchmark scores can be calculated for the land cover fractions of each natural PFT by comparing the ESACCI-derived land cover product to the GLC2000-derived land cover product. Model scores tend to be lower than benchmark scores, especially for tropical PFTs (broadleaf deciduous dry trees and C_4 grasses). For these PFTs uncertainty is introduced because remote sensing products do

320   not distinguish between cold and dry broadleaf deciduous trees and do not distinguish between C_3 and C_4 grasses. This distinction is made during the reclassification of the land cover categories to the nine CLASSIC PFTs and introduces some subjectivity (Wang et al., 2023). Specifically, deciduous trees above 30° in both hemispheres are assumed to be cold deciduous, deciduous trees below 20° in both hemispheres are assumed to be dry deciduous, and the fraction of trees that are cold vs. dry deciduous varies linearly between 0 and 1 with latitude for deciduous trees between 20° and 30°. The separation

325   of grasses into C_3 and C_4 is based on the time-invariant global product from (Still et al., 2003).

3.2 Future simulations

For SSP5-8.5, CLASSIC with dynamic land cover simulates increasing global area of trees, grasses, and total natural vegetation (Figure 4a). Global natural vegetation area decreases in the ESACCI-derived land cover product due to land use change, which is again the only driver of variation in natural vegetation area in the ESACCI-derived land cover product

330   (Figure 4a, Figure B1). Decreasing natural vegetation area due to land use change occurs primarily in South America, Africa, and Asia (Figure 4b). CLASSIC with dynamic land cover simulates increasing natural vegetation area because the loss of natural vegetation area due to land use change is offset by increasing natural vegetation area especially at high latitudes (Figure 4c).

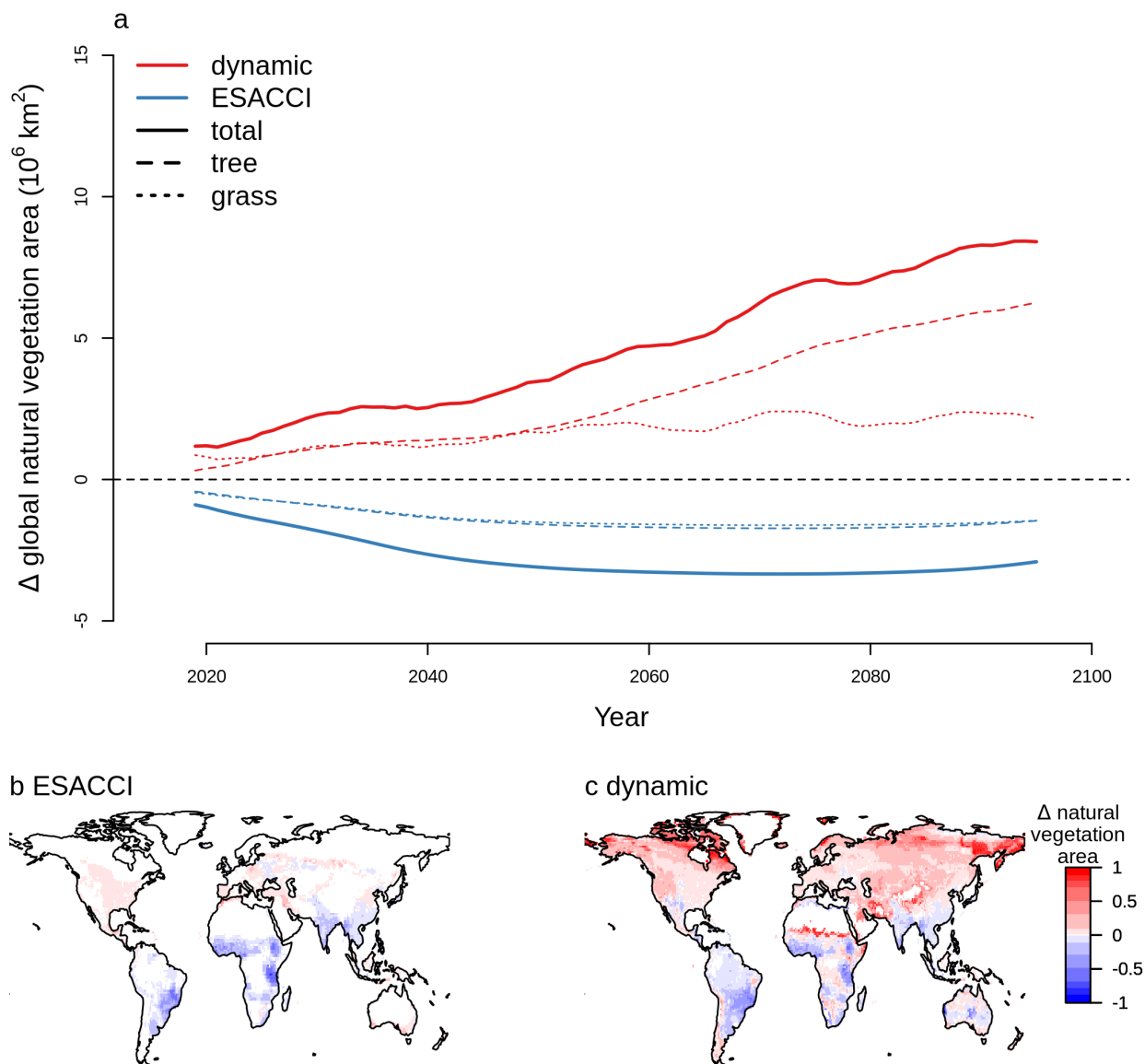



Figure 4. Change in natural vegetation area for SSP5-8.5 (2015 – 2100). a. Change in global area of natural vegetation, trees, and grasses simulated by CLASSIC with dynamic land cover in comparison to the ESACCI-derived land cover product. Change in vegetated fraction per grid cell b. in the ESACCI-derived land cover product and c. simulated by CLASSIC with dynamic land cover. Differences reflect the change between the average over 1995-2015 and 2080-2100. Figure B4 shows the change in fraction per grid cell for each PFT simulated by CLASSIC with dynamic land cover. Figure B7 shows the vegetated fraction per grid cell averaged over 1995-2015 and 2080-2100.

At high latitudes, CLASSIC with dynamic land cover simulates increasing areas of needleleaf evergreen trees, broadleaf deciduous cold trees, and C_3 and C_4 grasses, expanding into what was previously bare ground (Figure B5). This corresponds

to increasing temperatures at high latitudes (Figure B6). At low latitudes, CLASSIC with dynamic land cover simulates decreasing natural vegetation area in South America but variably decreasing and increasing natural vegetation area in Africa

345 and Asia. In the Amazon, the area of broadleaf evergreen trees decreases while the areas of C_3 and C_4 grasses increase (Figure B5). This corresponds to decreasing precipitation in this region (Figure B6). Conversely, in Africa and Asia, the area of broadleaf evergreen trees increases while the areas of C_3 and C_4 grasses decrease (Figure B5). This corresponds to increasing precipitation in this region (Figure B6). As an exception, in the Sahel, the areas of C_3 and C_4 grasses increase, expanding into what was previously bare ground (Figure B5). This also corresponds to increasing precipitation in this region

350 (Figure B6).

CLASSIC with prescribed land cover (simulated with an ESACCI-derived land cover product) and CLASSIC with dynamic land cover both simulate increasing NPP at higher latitudes for SSP5-8.5 (Figure 5a-b). NPP increases to a greater extent at higher latitudes in the simulation with dynamic land cover because of increasing natural vegetation area and NPP, especially that of trees (Figures 5c-f and 6a-c). NPP changes are similar at lower latitudes in both versions of CLASSIC (Figure 5a-b).

355 Both versions of CLASSIC project decreasing NPP in the Amazon (Figure 5a-b). NPP decreases in simulations with prescribed land cover simulations due to decreasing precipitation in the Amazon (Figure B6). NPP decreases slightly less in the simulation with dynamic land cover because decreasing tree area and NPP are offset by increasing grass area and NPP in the Amazon (Figures 5c-f and 6d-f). In the Sahel, NPP increases slightly in the simulation with dynamic land cover but decreases slightly in the simulation with prescribed land cover (Figure 5a-b). NPP decreases in simulations with prescribed

360 land cover due to land use change in the Sahel (Figure 6h). NPP increases slightly in the simulation with dynamic land cover because of increasing grass area and NPP, outweighing the effects of NPP in the Sahel(Figures 5c-f and 6g-i).

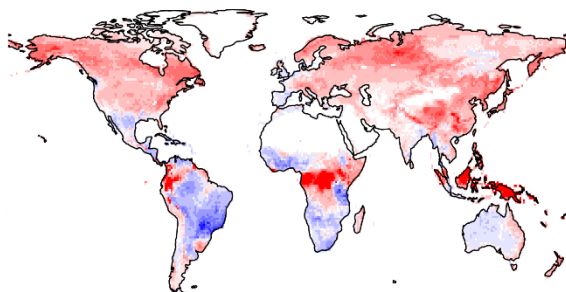
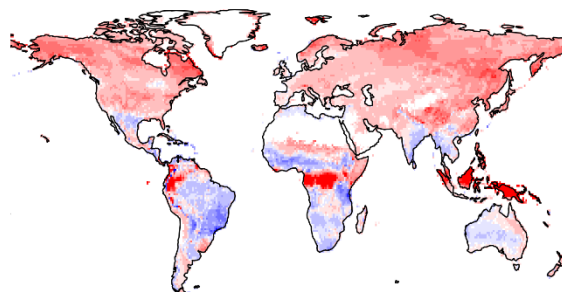
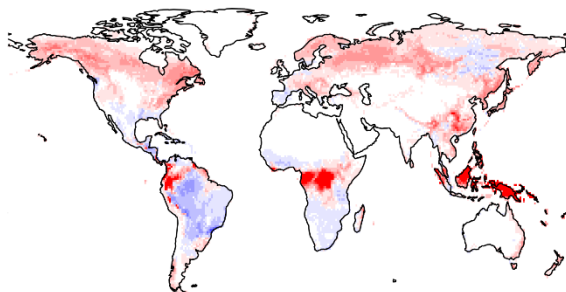
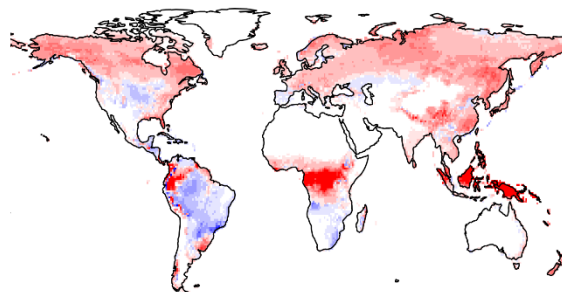
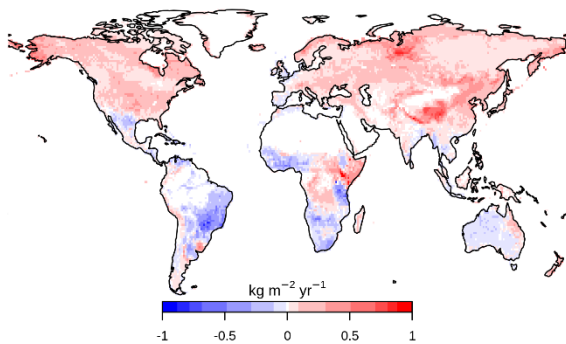
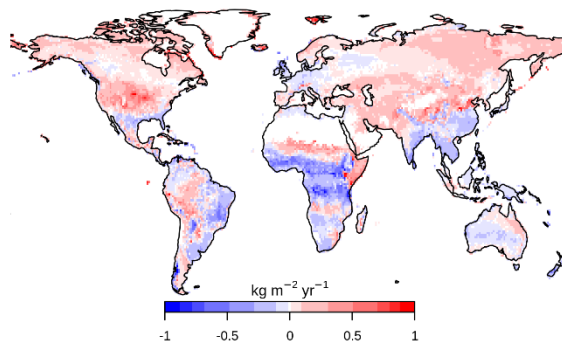



Figure 5. Change in net primary productivity (NPP) simulated by CLASSIC with prescribed land cover (with an ESACCI-derived land cover forcing) and CLASSIC with dynamic land cover for SSP5-8.5 (2015 – 2100). ab. Total NPP, cd. tree NPP, and ef. grass NPP. Differences reflect the change between the average over 1995-2015 and 2080-2100.

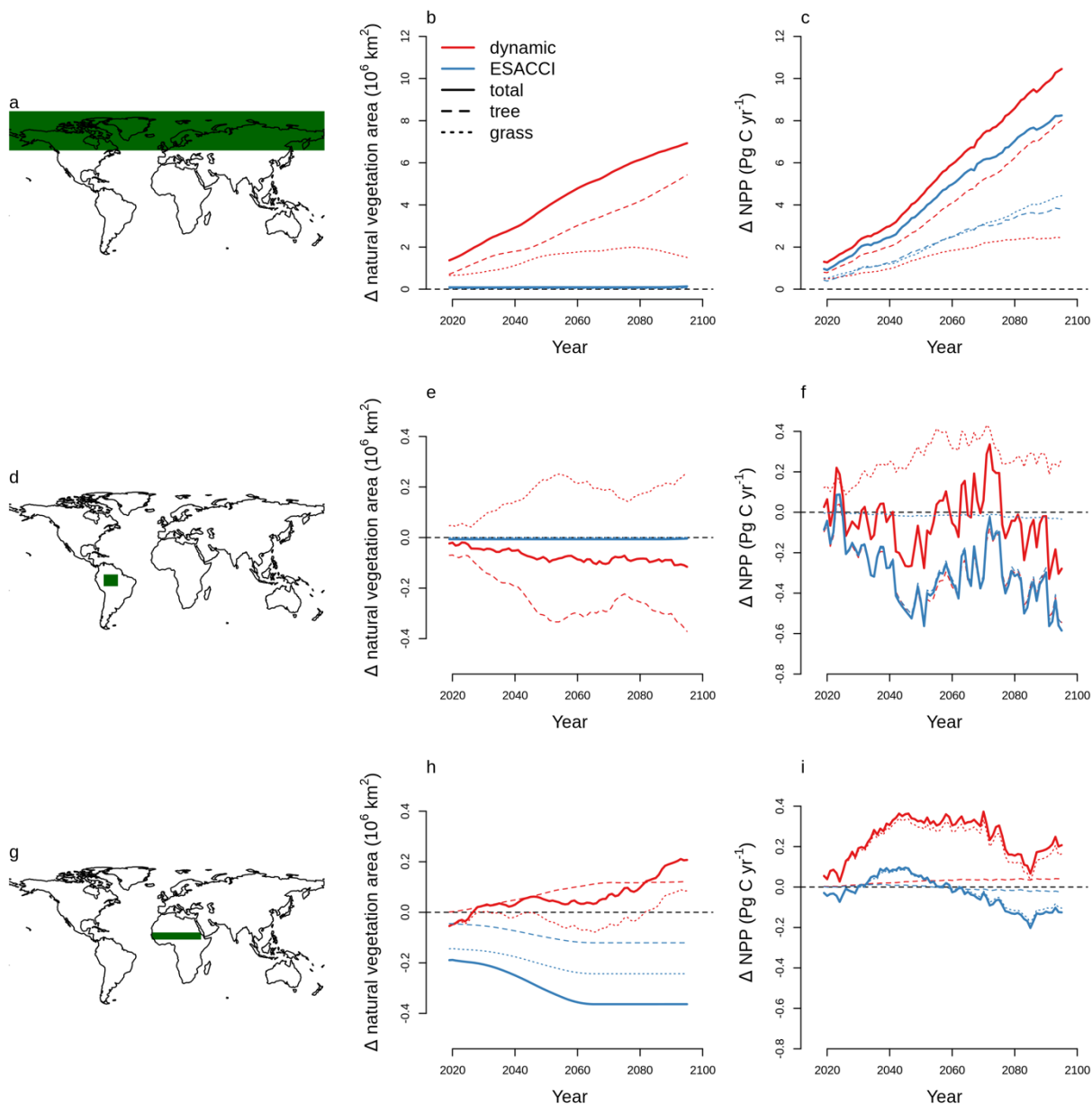



Figure 6. Change in natural vegetation area and net primary productivity (NPP) at high latitudes, the Amazon, and the Sahel by CLASSIC with prescribed land cover (with an ESACCI-derived land cover forcing) and CLASSIC with dynamic land cover for SSP5-8.5 (2015 – 2100). a. Map of high latitudes. b. Change in global area of natural vegetation, trees, and grasses in high latitudes. c. Change in NPP in high latitudes. d. Map of the Amazon. e. Change in global area of natural vegetation, trees, and grasses in the Amazon. f. Change in NPP in the Amazon. d. Map of the Sahel. e. Change in global area of natural vegetation, trees, and grasses in the Sahel. f. Change in NPP in the Sahel.

Net biome productivity (NBP), the net land-atmosphere CO_2 flux, increases in CLASSIC simulations of SSP5-8.5 with both prescribed and dynamic land cover (Figure 7a). The increase in NBP is almost twofold greater in the simulation with

dynamic land cover (5.28 Pg C yr^{-1} averaged over 2080 – 2100) than in the simulation with prescribed land cover (2.29 Pg C yr^{-1} averaged over 2080 – 2100). At the global scale, this increase in NBP is driven by increasing NPP (Figure 7b-c) which is primarily due to CO_2 fertilisation in both versions of CLASSIC. In CLASSIC with dynamic land cover, there is a stronger NPP increase at high Northern latitudes (Figure 7c). NPP increases at low latitudes in both versions of CLASSIC because increasing NPP in Africa and Asia outweighs decreasing NPP in South America. The change in NPP at low latitudes is similar between both versions of CLASSIC despite changing tree vs. grass areas as described above due to offsetting effects. Surface albedo decreases due to snow loss driven by increasing temperature in both versions of CLASSIC (Figure 7d), but this decrease is slightly stronger at high Northern latitudes in CLASSIC with dynamic land cover due to the expansion of natural vegetation into higher latitudes (Figure 7e). Evapotranspiration increases in both versions of CLASSIC (Figure 7f), but this increase is slightly stronger at high Northern latitudes in CLASSIC with dynamic land cover due to the expansion of natural vegetation into higher latitudes (Figure 7g).

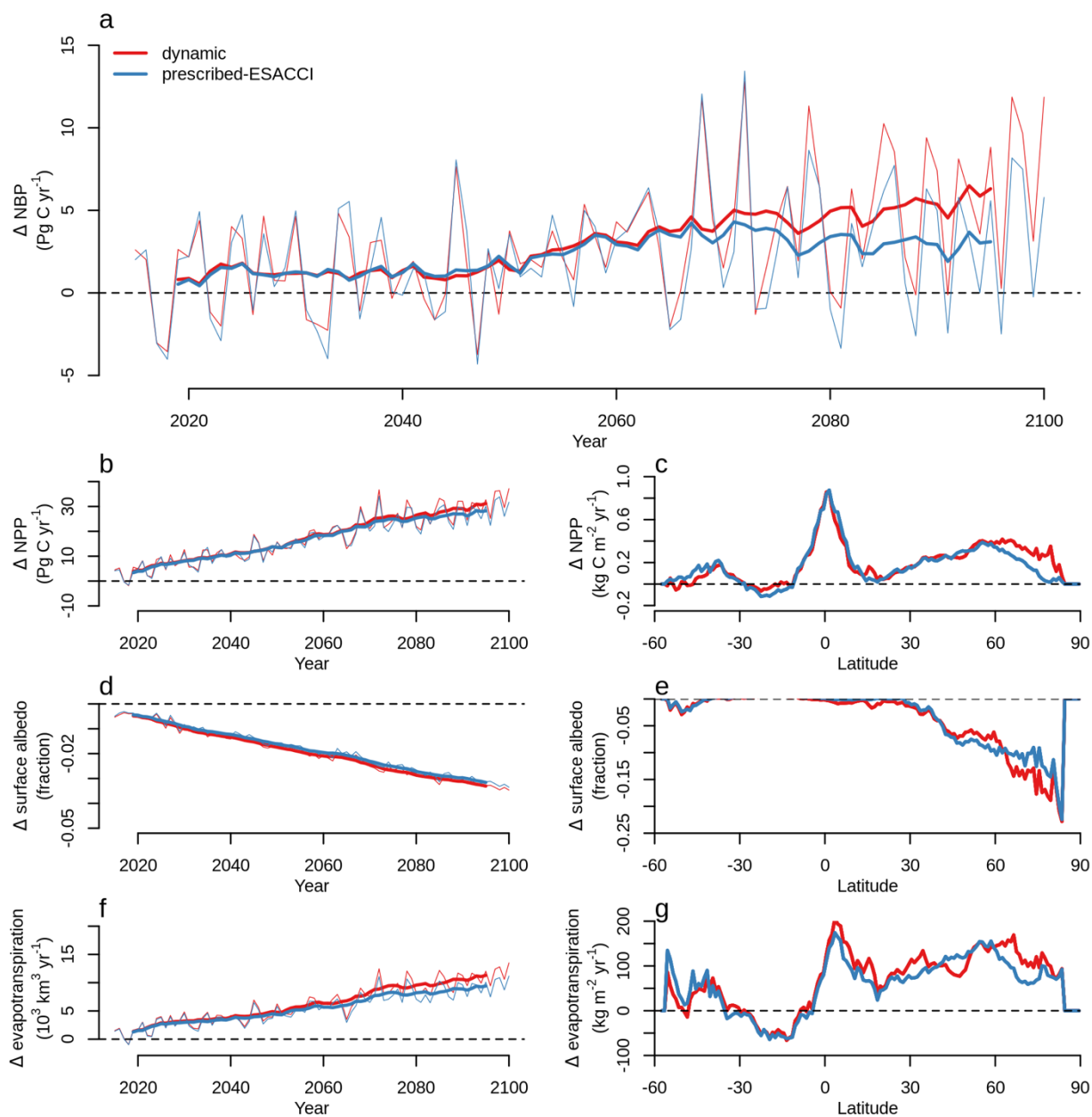



Figure 7. Change in a. net biome productivity (NBP), bc. net primary productivity (NPP), de. surface albedo, and fg. evapotranspiration simulated by CLASSIC with prescribed land cover (with an ESACCI-derived land cover forcing) and CLASSIC with dynamic land cover for SSP5-8.5 (2015 – 2100). For the latitudinal distributions, CLASSIC simulations were averaged over 2080-2100. Thin lines indicate the annual value and thick lines indicate the moving average over 10 years.

390

4. Discussion

Dynamic land cover is rarely implemented in terrestrial biosphere models (e.g., in only 3 out of 16 contributing models to the Global Carbon Project and in only 3 out of 11 contributing models to C[4]MIP within the 6th phase of CMIP) and the

simulated geographical distribution of natural vegetation has not yet been evaluated robustly as it lacked the methodology to
395   do so. The Global Carbon Project uses a statistical framework to evaluate the ability of its models to reproduce observations
of the C, water, and energy cycles – the International Land Model Benchmarking (ILAMB) framework (Collier et al., 2018),
which has since been expanded to include a method to quantify the uncertainty of the observation-based datasets themselves
(Seiler et al., 2022). We build on this to introduce a framework with which the dynamically simulated geographical
distribution of natural vegetation can be quantitatively evaluated against observations. We use this framework to show that
400   CLASSIC with dynamic land cover successfully simulates the geographical distribution of its natural PFTs for the present
day, reproducing observations (Figures 1-3). This framework can be applied across terrestrial biosphere models to evaluate
dynamic land cover alongside evaluations of the principal C, water, and energy cycle processes.

Future projections for Shared Socioeconomic Pathway 5-8.5 ("fossil-fueled development scenario", SSP5-8.5) differ
significantly between simulations with prescribed and dynamic land cover because in the latter the geographical distribution
405   of natural vegetation is able to respond to changing environmental conditions. The net land-atmosphere CO_2 flux (or net
biome productivity, NBP) is nearly twice as large in the simulation with dynamic land cover (5.28 Pg C yr^{-1} averaged over
2080 – 2100) than the simulation with prescribed land cover (2.29 Pg C yr^{-1} averaged over 2080 – 2100) (Figure 7). This
difference is attributed to three important range shifts that are only simulated when dynamic land cover is implemented: (1)
the expansion of trees into high latitudes, (2) the recession of trees accompanied by the expansion of grasses in the Amazon,
410   and (3) the expansion of grasses into the Sahel (Figures 5, 6, and 7). These responses occurred over the historical period as
well but to a lesser extent (Figure 1). We now discuss each of these range shifts in greater detail.

"Greening of the Arctic" describes the expansion of vegetation into high-latitude regions which occurs as previously
uninhabitable regions become habitable due to increasing temperatures (Chen et al., 2011; Elmendorf et al., 2012; Pearson et
al., 2013; Tape et al., 2006). While terrestrial biosphere models with prescribed land cover capture increasing plant growth at
415   high latitudes within their present-day range, these models are not able to simulate range shifts. Our results suggest these
range shifts could make a sizeable contribution to the terrestrial C sink via C sequestration in vegetation at high latitudes due
to the northward expansion of trees and grasses. Although the greening of the Arctic drives increased terrestrial C
sequestration in vegetation at high latitudes, there are other important feedbacks to the climate system, such as reduced
albedo (Chapin et al., 2005) and enhanced evapotranspiration (Swann et al., 2010).

420   "Amazonian dieback" describes the combination of deforestation and the increased mortality of trees in the Amazon
rainforest due to changing climate in the region (Malhi et al., 2008; Parry et al., 2022). Tree mortality occurs as previously
habitable regions become inhabitable due to the sensitivity of trees to drought and heat stress (Aleixo et al., 2019; Phillips et
al., 2009). While the direct effects of reduced precipitation and elevated temperature on plant growth as well as deforestation
are captured by terrestrial biosphere models with prescribed land cover, these models are incapable of simulating transitions
425   from forest to grassland. Additionally, CLASSIC dynamically simulates fire and the loss of natural vegetation area due to
fire (Arora and Melton, 2018), which influences and feeds back to these transitions (Davidson et al., 2012). Finally,
Amazonian dieback self-amplifies via feedbacks to the climate system, as decreased forest cover decreases

evapotranspiration which could amplify regional drought (Zemp et al., 2017). The effects of these feedbacks as well as the feedbacks associated with the greening of the Artic are not evaluated in our offline simulations and warrant further study.

430 Finally, the "greening of the Sahel" describes the expansion of vegetation into the Sahara which occurs as previously uninhabitable regions become habitable due to increasing precipitation (Brandt et al., 2015; Herrmann et al., 2005; Olsson et al., 2005). This range shift is not captured in terrestrial biosphere models with prescribed land cover. However, there is large uncertainty in future projections of precipitation in this region (Monerie et al., 2020). While less explored than the previous two effects, greening of the Sahel is regionally important and could make a non-trivial contribution to the global C, water,

435 and energy cycles, warranting further study.

These responses are consistent with those of other terrestrial biosphere models that implement dynamic land cover. Both Scholze et al. (2006) and Alo & Wang (2008) observed increasing tree area at high latitudes and the replacement of trees with grasses in the Amazon. Sitch et al. (2008), which compared four models with dynamic land cover, found similar results, although the magnitudes of the responses differed between models. Port et al. (2012) conducted and examined a fully

440 coupled simulation with a land model with dynamic land cover and similarly observed both greening of the Arctic and Amazonian dieback. Because there is large uncertainty in climate projections in the Sahel region, greening of the Sahel is often but not always simulated (Sitch et al., 2008). Our results follow the consistent pattern simulated by these other models, but our unique experimental design allows us to isolate the impact of dynamic land cover implementation (vs. prescribed land cover implementation) and compare it against the effects of other global change drivers.

445 **5. Conclusions**

Our results illustrate the potential effects of range shifts alongside variation in plant growth that is driven by climate change. While all terrestrial biosphere models account for the influence of global change drivers such as CO_2 fertilisation, increasing temperatures, and variable precipitation regimes on plant growth, terrestrial biosphere models with prescribed land cover cannot capture range shifts that are driven by climate change. However, few terrestrial biosphere models incorporate

450 dynamic land cover. Importantly, the shifting ranges of biomes have critical feedbacks to climate change as well as critical consequences for biodiversity and human well-being (Pecl et al., 2017) alongside their consequences to the global C cycle.

Appendix A: Detailed description of dynamic land cover in CLASSIC.

When dynamic land cover is implemented, the fractional coverage of a natural PFT is the result of colonization and mortality and is described in detail in Melton & Arora (2016). The colonization rate of PFT n (c_n; day^{-1}) is determined by its net

455 primary productivity (NPP):

$$c_n = \min\left[\max\left[\lambda_{1,n}, \lambda_{2,n}\right], 0.1\right] NPP_n \frac{1}{s_{sap,n} \max[0.25, \min[C_{veg,n}, 5.0]]} \tag{A1}$$

NPP_n is the NPP of PFT n and $C_{veg,n}$ is the vegetation C biomass of PFT n. $s_{sap,n}$ is a factor for converting vegetation C biomass to seedling C biomass of PFT n (unitless) (Table B3). $\lambda_{1,n}$ and $\lambda_{2,n}$ determine the fraction of NPP that is used for spatial expansion within a given grid cell of PFT n (unitless):

$$\lambda_{1,n} = \begin{cases} 0, & LAI_n \leq LAI_{min,n} \\ 0.1 \frac{LAI_n - LAI_{min,n}}{LAI_{max,n} - LAI_{min,n}}, & LAI_{min,n} < LAI_n < LAI_{max,n} \\ 0.1, & LAI_{max,n} \leq LAI_n \end{cases} \tag{A2}$$

$$\lambda_{2,n} = \begin{cases} 0, & LAI_n \leq 0.25 LAI_{min,n} \\ \cosh\left(0.115\left(LAI_n - 0.25 LAI_{min,n}\right)\right) - 1, & 0.25 LAI_{min,n} < LAI_n \end{cases} \tag{A3}$$

$LAI_{min,n}$ and $LAI_{max,n}$ are minimum and maximum leaf area index thresholds, respectively, of PFT n (m² m⁻²) (Table B3). The mortality rate of PFT n (m_n; day⁻¹) is the sum of intrinsic or age-related mortality ($m_{intr,n}$; day⁻¹), mortality due to reduced growth ($m_{ge,n}$; day⁻¹), mortality due to fire ($m_{dist,n}$; day⁻¹), and mortality when a PFT exists outside its bioclimatic limits ($m_{bioclim,n}$; day⁻¹):

$$m_n = m_{intr,n} + m_{ge,n} + m_{dist,n} + m_{bioclim,n} \tag{A4}$$

$$m_{intr,n} = 1 - \exp\left(\frac{-4.605}{A_{max,n}}\right) \tag{A5}$$

$$m_{ge,n} = \frac{m_{ge,max,n}}{1 + \left(300\,^{m^2}/_{kg\,C}\right)(\max[0, ge_n])} \tag{A6}$$

$$m_{dist,n} = \varsigma_{r,n} \frac{f_{burned,n}}{f_n} \tag{A7}$$

$A_{max,n}$ is the maximum age of PFT n (yr) (Table B3). $m_{ge,max,n}$ is the maximum growth-related mortality rate occurring when no growth occurs of PFT n (day⁻¹) (Table B3). ge_n is growth efficiency over the previous year of PFT n (kg C m⁻²):

$$ge_n = \frac{\Delta C_{s,n} + \Delta C_{r,n}}{L_{max,n}} \tag{A8}$$

$\Delta C_{s,n}$ is the stem C increment over the previous year of PFT n (kg C m⁻²), $\Delta C_{r,n}$ is the root C increment over the previous year of PFT n (kg C m⁻²), and $L_{max,n}$ is the maximum LAI over the previous year of PFT n (m² m⁻²). $\varsigma_{r,n}$ is the susceptibility to stand replacing fire of PFT n (fraction) (Table B3). $f_{burned,n}$ is the fractional area burned within a day of PFT n and depends on wind speed and soil moisture as well as the probability of fire occurrence (which depends on the availability of vegetation biomass as a fuel source, the combustibility of the fuel source based on soil moisture, and the presence of an ignition source based on lightning and population density) and suppression (which depends on population density). This is described in detail in Arora & Melton (2018).

Six bioclimatic indices are used to determine the spatial range of PFTs: air temperature of the coldest month, air temperature of the warmest month, aridity index (ratio of potential evaporation to precipitation), growing degree days (cumulative number of days with air temperature above 5°C), dry season length (number of consecutive months with precipitation less than potential evaporation), and precipitation surplus (difference between precipitation and potential evaporation). Bioclimatic indices for each grid cell are updated annually on a 25-year timescale using exponential smoothing:

485 $\quad X(t+1) = X(t)e^{-1/25} + x(t)\left(1 - e^{-1/25}\right)$ $\hfill$ (A9)

$x(t)$ represents a bioclimatic index at year t and $X(t)$ represents the smoothed bioclimatic index at year t. This accounts for time lags in the response of vegetation to climate change drivers (Wu et al., 2015). $m_{bioclim,n} = 0.25$ when any bioclimatic index is outside its corresponding bioclimatic limit for a given PFT and $m_{bioclim,n} = 0$ when all bioclimatic indices are inside their corresponding bioclimatic limits for a given PFT. Bioclimatic limits for each PFT are given in Table B1. These

490 were derived by first calculating the bioclimatic indices for each grid cell using the CRUJRA climate dataset averaged over 1900 to 1920 (Harris et al., 2020). Then, the 1% and 99% quantiles of these bioclimatic indices over all grid cells for each land cover type in the Collection 5 MODIS Global Land Cover Type International Geosphere-Biosphere Programme (IGBP) product (Friedl et al., 2010) were calculated and associated with a CLASSIC PFT. Note that this assumes that the current biome ranges are in equilibrium with the 1900-1920 climate.

495 Each natural PFT is given a dominance rank (i) where PFTs with higher dominance ranks invade PFTs with lower dominance ranks and bare ground. Natural PFTs cannot invade crop area. Tree PFTs are given higher dominance ranks than grass PFTs. Within tree PFTs and within grass PFTs, a PFT with a higher c_n is dominant over all PFTs with lower c_n. Each PFT is thus ranked from $1, 2, \dots, i-1, i, i+1, \dots N$ where the PFT with dominance rank 1 is the most dominant. Overall, the change in fractional coverage of a PFT with dominance rank i is due to (1) its colonization into the areas of PFTs with lower

500 dominance ranks ($i+1, \dots, N$) and bare ground ($N+1$), (2) colonization by PFTs with higher dominance ranks ($i, \dots, i-1$) into its area, and (3) its mortality:

$$\frac{df_i}{dt} = (c_i f_{i+1} + \cdots + c_i f_N + c_i f_{N+1}) - (c_1 f_i + \cdots + c_{i-1} f_i) - m_i f_i \qquad (A10)$$

505 The change in fractional coverage of bare ground is due to mortality of the PFTs and colonization of bare ground by the PFTs:

$$\frac{df_{N+1}}{dt} = \sum_{n=1}^{N} m_n f_n - \sum_{n=1}^{N} c_n f_{N+1} \qquad (A11)$$

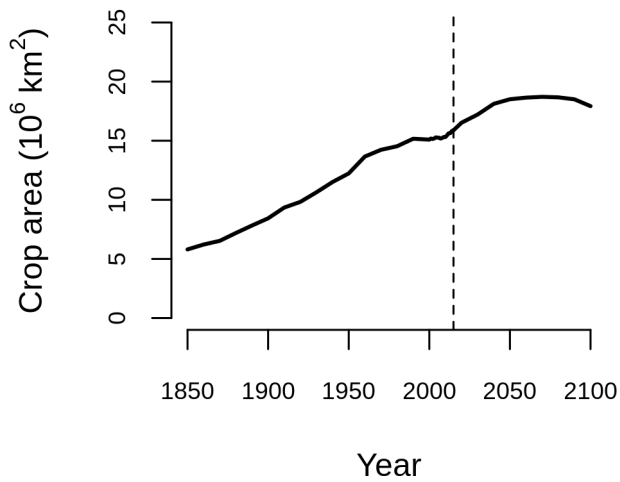
Appendix B: Supplemental figures and tables.

510

Figure B1: Crop area over the historical period (1850 – 2015) and SSP5-8.5 (2015 – 2100) from the LUH2-GCB2022 (Land-Use Harmonization 2) dataset (Chini et al., 2021; Hurtt et al., 2020; Klein Goldewijk et al., 2017).

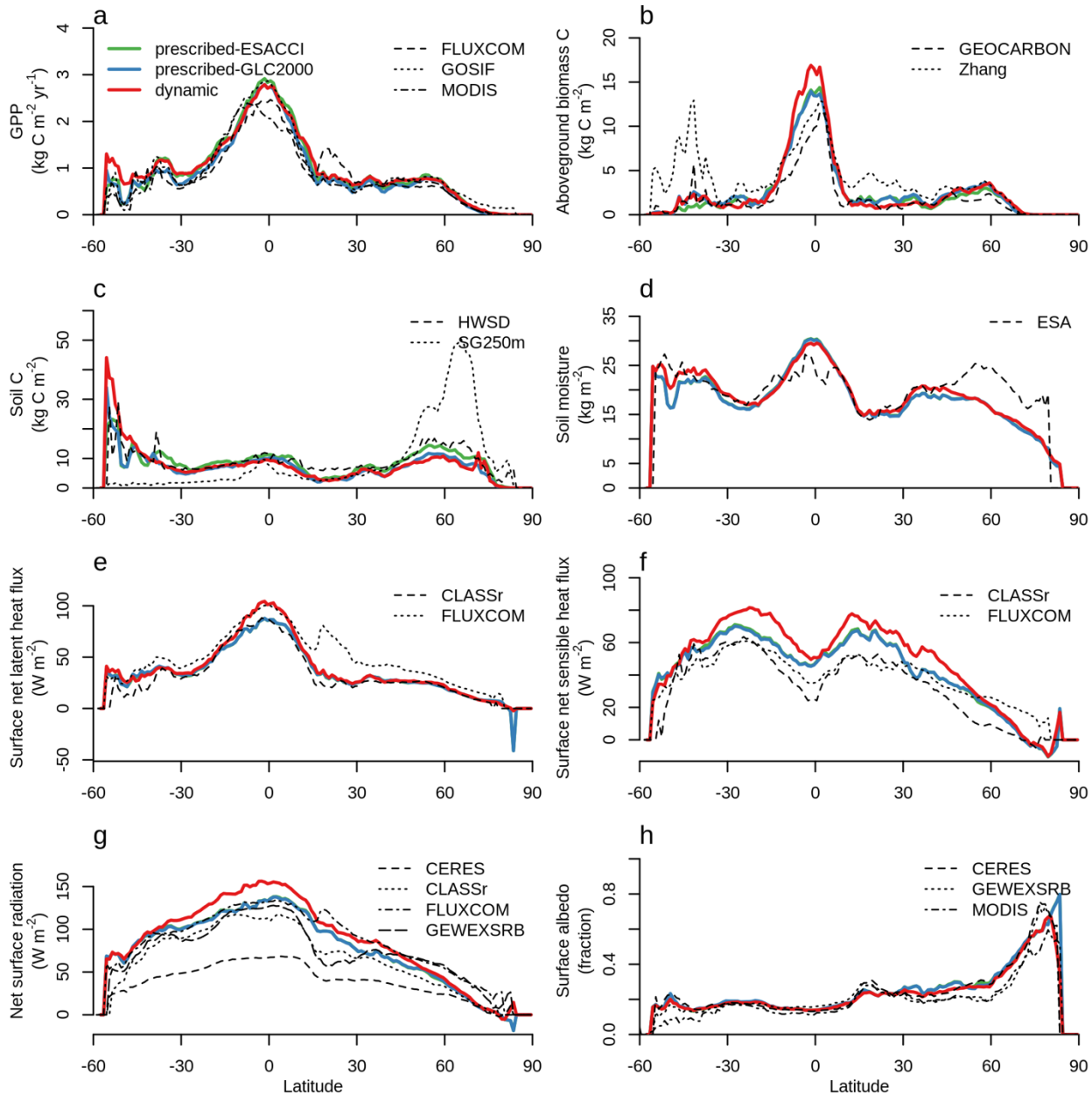



Figure B2: Latitudinal distributions of a. gross primary productivity (GPP), b. aboveground biomass C, c. soil C, d. soil moisture, e. surface net latent heat flux, f. surface net sensible heat flux, g. net surface radiation, and h. surface albedo simulated by CLASSIC with prescribed land cover (with an ESACCI-derived land cover forcing and a GLC2000-derived land cover forcing) and CLASSIC with dynamic land cover in comparison to present-day observations. CLASSIC simulations were averaged over 2000-2020. Observation-based datasets are described in the Methods. Figure B2 shows the scores of model performance in reproducing observation-based datasets for each variable.

520

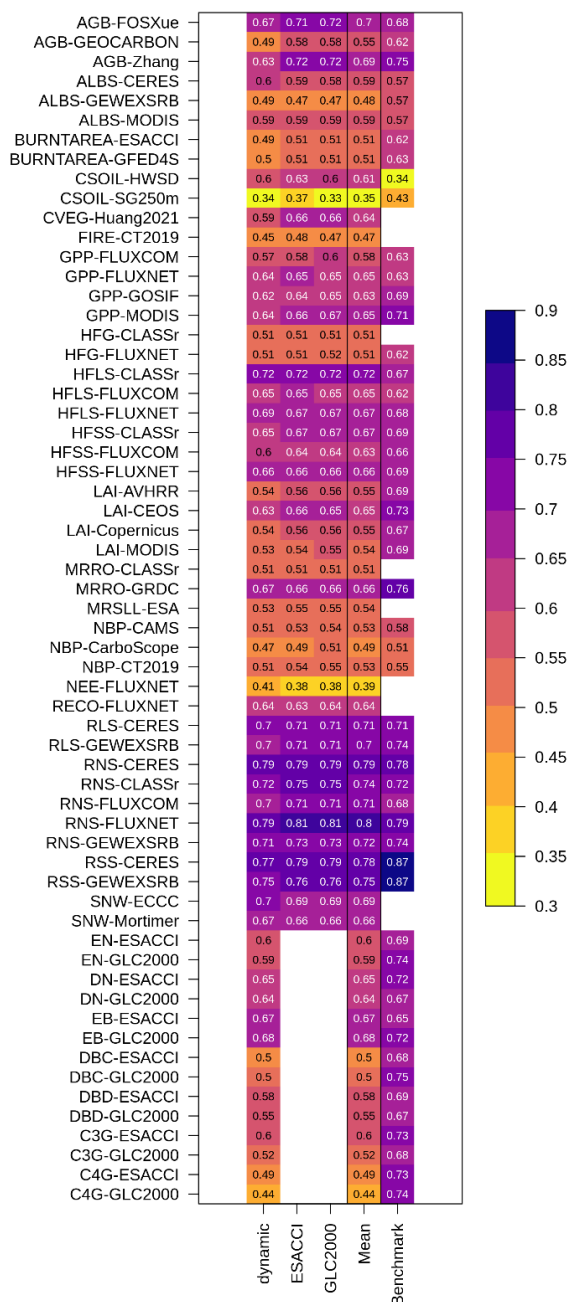



Figure B3: Scores of model performance in reproducing observation-based datasets of principal C, water, and energy variables as well as land cover fractions of each natural plant functional type (PFT) for simulations of CLASSIC with prescribed land cover (with an ESACCI-derived land cover forcing and a GLC2000-derived land cover forcing) and dynamic land cover. Abbreviations for C, water, and energy variables and observation-based datasets are described in the Methods. Abbreviations for PFTs are:
525   **evergreen needleleaf (EN), deciduous needleleaf (DN), evergreen broadleaf (EB), deciduous broadleaf cold (DBC), deciduous broadleaf dry (DBD), C$_3$ grass (C3G), and C$_4$ grass (C4G).**

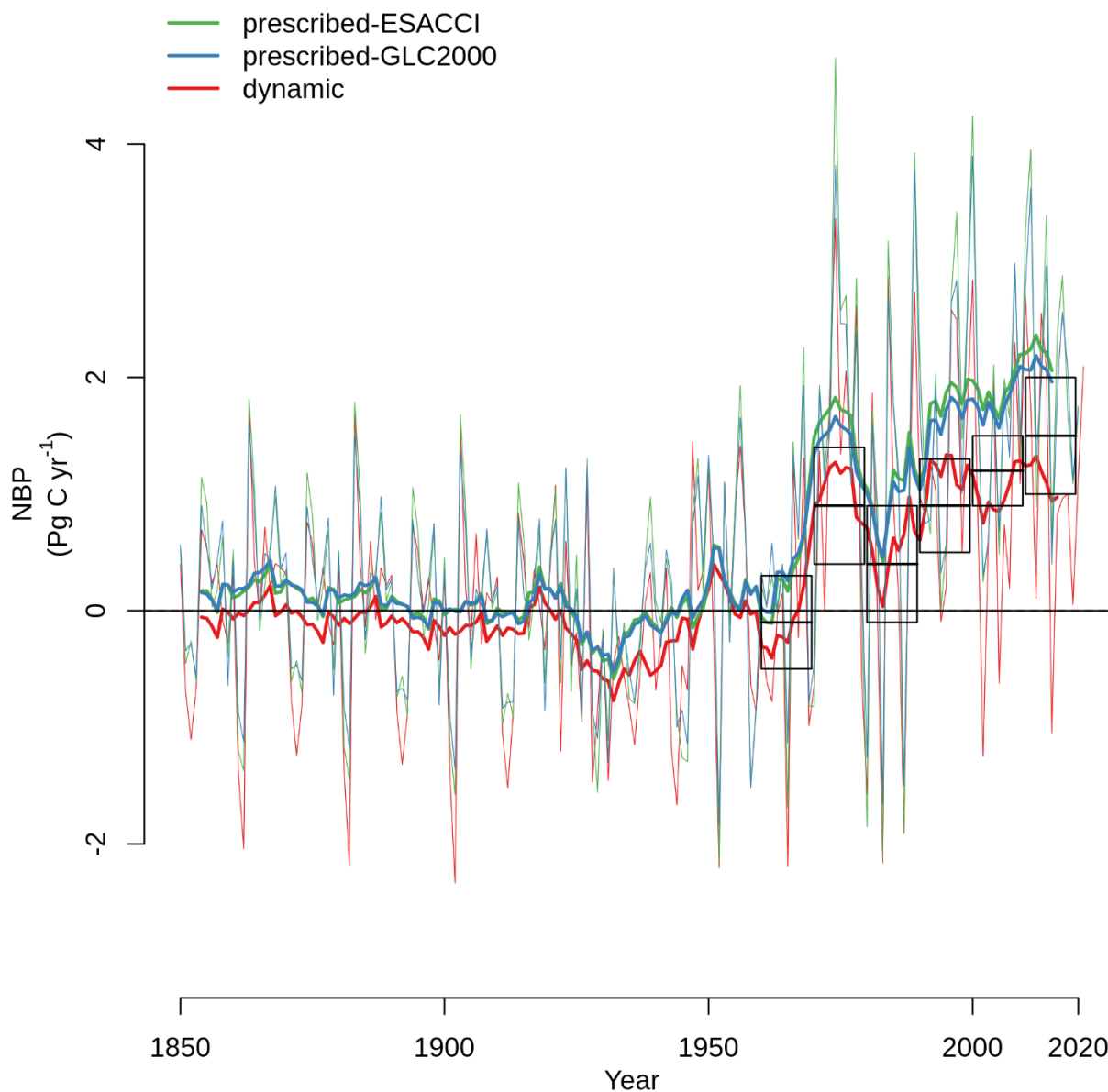



Figure B4: Net biome productivity (NBP) simulated by CLASSIC with prescribed land cover (with an ESACCI-derived land cover forcing and a GLC2000-derived land cover forcing) and CLASSIC with dynamic land cover. Boxes indicate estimates from the suite of models in the Global Carbon Project where dark horizontal lines indicate the mean and the extent indicates one standard deviation (Friedlingstein et al., 2022). Thin lines indicate the annual value and thick lines indicate the moving average over 10 years.

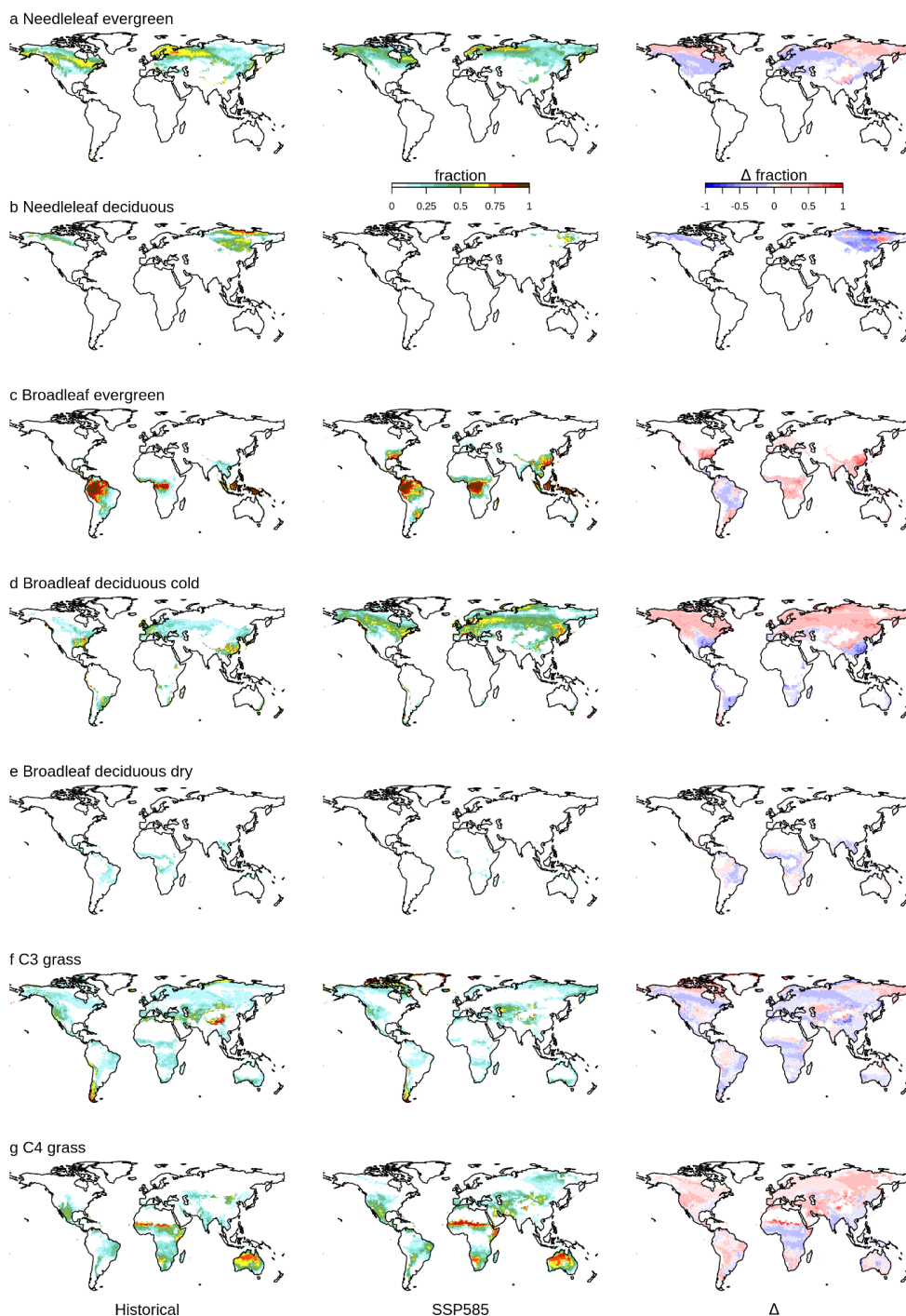



Figure B5: Fraction per grid cell in the historical period (averaged over 1995-2015), fraction per grid cell in SSP5-8.5 (averaged over 2080-2100), and change in fraction per grid cell for each plant functional type (PFT) simulated by CLASSIC with dynamic land cover. a. Evergreen needleleaf, b. deciduous needleleaf, c. evergreen broadleaf, d. deciduous broadleaf cold, e. deciduous broadleaf dry, f. C_3 grass, and g. C_4 grass. Differences reflect the change between the average over 1995-2015 and 2080-2100.

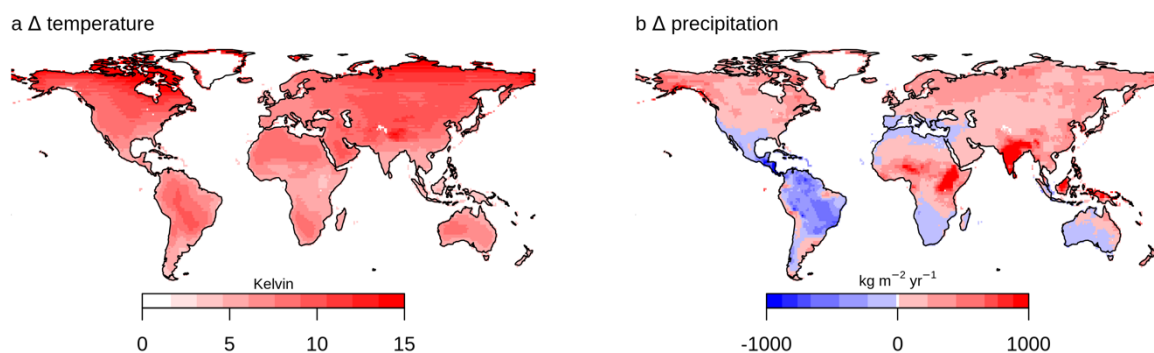



Figure B6: Change in a. temperature and b. precipitation between the historical period (averaged over 1995-2015) and SSP5-8.5 (averaged over 2080-2100).

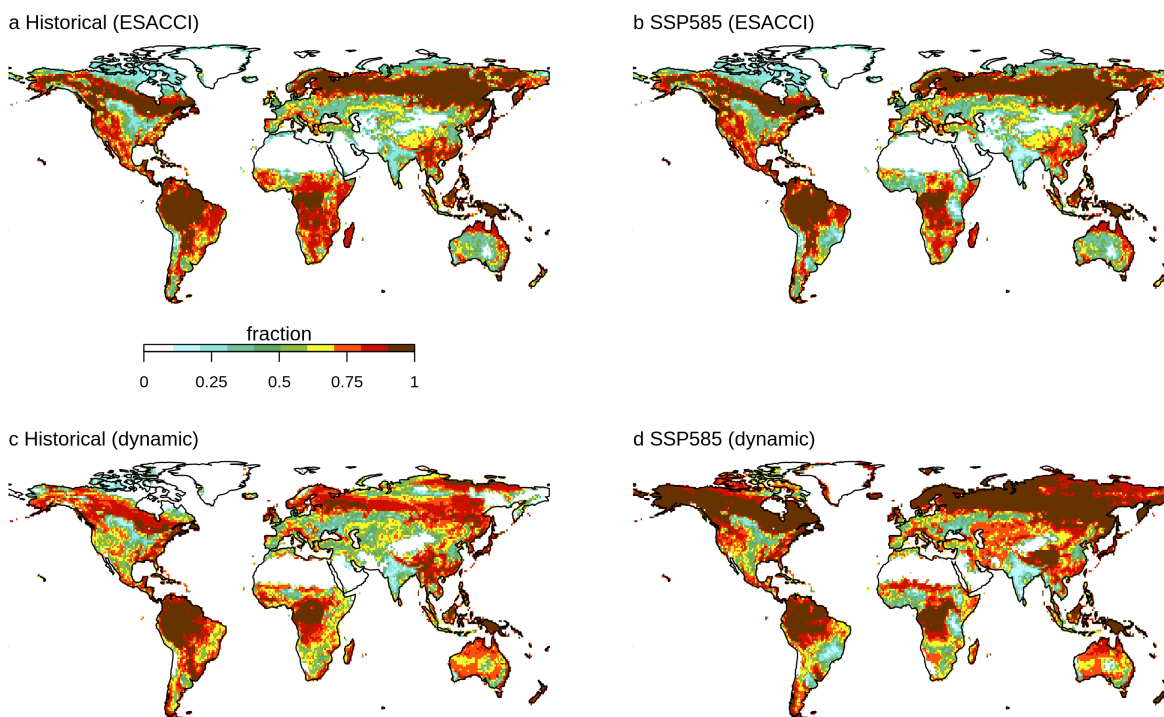



Figure B7: Vegetated fraction per grid cell in the ab. ESACCI land cover product and cd. simulated by CLASSIC with dynamic land cover over ac. the historical period (averaged over 1995-2015) and bd. for SSP5-8.5 (averaged over 2080-2100).

Table B1: Parameters associated with bioclimatic indices.

Parameter	EN	DN	EB	DBC	DBD	C3G	C4G
Maximum air temperature of the coldest month	-2	-27		17			
Minimum air temperature of the coldest month			1	-33	15		
Maximum air temperature of the warmest month	28	19		30			
Minimum air temperature of the warmest month							20
Maximum aridity index	8	8	8	8	8	8	8
Minimum aridity index					0.5		
Minimum growing degree days	370	340	1320	430	4920		
Maximum dry season length			8				
Minimum dry season length					5		
Precipitation surplus	150	150	150	150	150		

Table B2: Observation-based datasets used in AMBER. Described in detail in (Seiler et al., 2022). Datasets are globally grided or in situ (specified).

Variable	Variable abbreviation	Observation-based dataset
Aboveground biomass carbon	AGB	FOSXue (in situ) (Schepaschenko et al., 2019; Xue et al., 2017) GEOCARBON (Avitabile et al., 2016; Santoro et al., 2015) Zhang (Zhang and Liang, 2020)
Albedo	ALBS	CERES (Kato et al., 2013) GEWEXSRB (Stackhouse et al., 2011) MODIS (Strahler and Muller, 1999)
Area burned	BURNTAREA	ESACCI (Chuvieco et al., 2018) GFED4S (Giglio et al., 2010)
Soil carbon	CSOIL	HWSD (Todd-Brown et al., 2013; Wieder, 2014) SG250m (Hengl et al., 2017)
Vegetation carbon	CVEG	Huang2021 (Huang et al., 2021)
Fire emissions	FIRE	CT2019 (Jacobson et al., 2020)
Gross primary productivity	GPP	FLUXCOM (Jung et al., 2020) GOSIF (Li and Xiao, 2019) MODIS (Zhang et al., 2017) FLUXNET (in situ) (Pastorello et al., 2020)
Soil heat flux	HFG	CLASSr (Hobeichi et al., 2020) FLUXNET (in situ) (Pastorello et al., 2020)
Latent heat flux	HFLS	CLASSr (Hobeichi et al., 2020) FLUXCOM (Jung et al., 2019) FLUXNET (in situ) (Pastorello et al., 2020)
Sensible heat flux	HFSS	CLASSr (Hobeichi et al., 2020) FLUXCOM (Jung et al., 2019) FLUXNET (in situ) (Pastorello et al., 2020)
Leaf area index	LAI	AVHRR (Claverie et al., 2016) CEOS (in situ) (Garrigues et al., 2008)

		Copernicus (Verger et al., 2014)
		MODIS (Myneni et al., 2002)
Streamflow	MRRO	CLASSr (Hobeichi et al., 2020)
		GRDC (in situ) (Dai and Trenberth, 2002)
Soil moisture	MRSLL	ESA (Liu et al., 2011)
Net biome productivity	NBP	CAMS (Agustí-Panareda et al., 2019)
		CarboScope (Rödenbeck et al., 2018)
		CT2019 (Jacobson et al., 2020)
Net ecosystem exchange	NEE	FLUXNET (in situ) (Pastorello et al., 2020)
Ecosystem respiration	RECO	FLUXNET (in situ) (Pastorello et al., 2020)
Net surface long wave radiation	RLS	CERES (Kato et al., 2013)
		GEWEXSRB (Stackhouse et al., 2011)
Net surface radiation	RNS	CERES (Kato et al., 2013)
		CLASSr (Hobeichi et al., 2020)
		FLUXCOM (Jung et al., 2019)
		FLUXNET (Pastorello et al., 2020)
		GEWEXSRB (Stackhouse et al., 2011)
Net surface shortwave radiation	RSS	CERES (Kato et al., 2013)
		GEWEXSRB (Stackhouse et al., 2011)
Snow water equivalent	SNW	Mortimer (Mortimer et al., 2020)

550

Table B3: Parameters associated with the calculation of colonization and mortality rates for each PFT.

Parameter	EN	DN	EB	DBC	DBD	C3G	C4G
$s_{sap,n}$	0.30	0.10	0.10	0.14	0.30	0.10	0.10
$LAI_{min,n}$	1.0	0.25	1.5	1.5	1.5	0.5	0.5
$LAI_{max,n}$	4.0	2.0	6.0	6.0	6.0	3.0	3.0
$A_{max,n}$	250	400	600	250	500	0	0
$m_{ge,max,n}$	0.005	0.005	0.005	0.005	0.005	0.1	0.1
$\varsigma_{r,n}$	0.20	0.20	0.50	0.20	0.15	0.25	0.25

Code availability

The source code for CLASSIC is available on the CLASSIC community Zenodo page (https://zenodo.org/communities/classic?q=&l=list&p=1&s=10&sort=newest). AMBER is available
555 at https://gitlab.com/cseiler/AMBER.

Author contribution

S.K.G. designed and conducted the study and prepared the initial manuscript. V.K.A. provided input on the study design and the initial manuscript. C.S. and L.W. provided input on the subsequent revisions.

Competing interests

560   The authors declare that they have no conflict of interest.

Acknowledgements

The authors would like to thank the input from the CLASSIC team (Paul Bartlett, Mike Brady, Ed Chan, Salvatore Curasi, Joe Melton, and Gesa Meyer).

References

565   Agustí-Panareda, A., Diamantakis, M., Massart, S., Chevallier, F., Muñoz-Sabater, J., Barré, J., Curcoll, R., Engelen, R., Langerock, B., Law, R. M., Loh, Z., Morguí, J. A., Parrington, M., Peuch, V.-H., Ramonet, M., Roehl, C., Vermeulen, A. T., Warneke, T., and Wunch, D.: Modelling CO_2 weather – why horizontal resolution matters, Atmospheric Chemistry and Physics, 19, 7347–7376, https://doi.org/10.5194/acp-19-7347-2019, 2019.

Aleixo, I., Norris, D., Hemerik, L., Barbosa, A., Prata, E., Costa, F., and Poorter, L.: Amazonian rainforest tree mortality
570   driven by climate and functional traits, Nature Climate Change, 9, 384–388, https://doi.org/10.1038/s41558-019-0458-0, 2019.

Alo, C. A. and Wang, G.: Potential future changes of the terrestrial ecosystem based on climate projections by eight general circulation models, Journal of Geophysical Research: Biogeosciences, 113, https://doi.org/10.1029/2007JG000528, 2008.

Arora, V. K. and Boer, G. J.: A parameterization of leaf phenology for the terrestrial ecosystem component of climate
575   models, Global Change Biology, 11, 39–59, https://doi.org/10.1111/j.1365-2486.2004.00890.x, 2005a.

Arora, V. K. and Boer, G. J.: Fire as an interactive component of dynamic vegetation models, Journal of Geophysical Research: Biogeosciences, 110, https://doi.org/10.1029/2005jg000042, 2005b.

Arora, V. K. and Boer, G. J.: Simulating Competition and Coexistence between Plant Functional Types in a Dynamic Vegetation Model, Earth Interactions, 10, 1–30, https://doi.org/10.1175/EI170.1, 2006.

580   Arora, V. K. and Melton, J. R.: Reduction in global area burned and wildfire emissions since 1930s enhances carbon uptake by land, Nature Communications, 9, 1–10, https://doi.org/10.1038/s41467-018-03838-0, 2018.

Arora, V. K., Katavouta, A., Williams, R. G., Jones, C. D., Brovkin, V., Friedlingstein, P., Schwinger, J., Bopp, L., Boucher, O., Cadule, P., Chamberlain, M. A., Christian, J. R., Delire, C., Fisher, R. A., Hajima, T., Ilyina, T., Joetzjer, E., Kawamiya, M., Koven, C. D., Krasting, J. P., Law, R. M., Lawrence, D. M., Lenton, A., Lindsay, K., Pongratz, J., Raddatz, T., Séférian,
585   R., Tachiiri, K., Tjiputra, J. F., Wiltshire, A., Wu, T., and Ziehn, T.: Carbon–concentration and carbon–climate feedbacks in CMIP6 models and their comparison to CMIP5 models, Biogeosciences, 17, 4173–4222, https://doi.org/10.5194/bg-17-4173-2020, 2020.

Arora, V. K., Seiler, C., Wang, L., and Kou-Giesbrecht, S.: Towards an ensemble-based evaluation of land surface models in light of uncertain forcings and observations, Biogeosciences, 20, 1313–1355, https://doi.org/10.5194/bg-20-1313-2023,
590   2023.

Asaadi, A. and Arora, V. K.: Implementation of nitrogen cycle in the CLASSIC land model, Biogeosciences, 18, 669–706, https://doi.org/10.5194/bg-18-669-2021, 2021.

Asaadi, A., Arora, V. K., Melton, J. R., and Bartlett, P.: An improved parameterization of leaf area index (LAI) seasonality in the Canadian Land Surface Scheme (CLASS) and Canadian Terrestrial Ecosystem Model (CTEM) modelling framework,
595 Biogeosciences, 15, 6885–6907, https://doi.org/10.5194/bg-15-6885-2018, 2018.

Avitabile, V., Herold, M., Heuvelink, G. B. M., Lewis, S. L., Phillips, O. L., Asner, G. P., Armston, J., Ashton, P. S., Banin, L., Bayol, N., Berry, N. J., Boeckx, P., de Jong, B. H. J., DeVries, B., Girardin, C. A. J., Kearsley, E., Lindsell, J. A., Lopez-Gonzalez, G., Lucas, R., Malhi, Y., Morel, A., Mitchard, E. T. A., Nagy, L., Qie, L., Quinones, M. J., Ryan, C. M., Ferry, S. J. W., Sunderland, T., Laurin, G. V., Gatti, R. C., Valentini, R., Verbeeck, H., Wijaya, A., and Willcock, S.: An integrated
600 pan-tropical biomass map using multiple reference datasets, Global Change Biology, 22, 1406–1420, https://doi.org/10.1111/gcb.13139, 2016.

Box, E. O.: Plant functional types and climate at the global scale, Journal of Vegetation Science, 7, 309–320, 1996.

Brandt, M., Mbow, C., Diouf, A. A., Verger, A., Samimi, C., and Fensholt, R.: Ground- and satellite-based evidence of the biophysical mechanisms behind the greening Sahel, Global Change Biology, 21, 1610–1620,
605 https://doi.org/10.1111/gcb.12807, 2015.

Buchner, M. and Reyer, C.: ISIMIP3b bias-adjusted atmospheric climate input data, , https://doi.org/10.48364/ISIMIP.842396.1, 2021.

Chapin, F. S., Sturm, M., Serreze, M. C., McFadden, J. P., Key, J. R., Lloyd, A. H., McGuire, A. D., Rupp, T. S., Lynch, A. H., Schimel, J. P., Beringer, J., Chapman, W. L., Epstein, H. E., Euskirchen, E. S., Hinzman, L. D., Jia, G., Ping, C.-L.,
610 Tape, K. D., Thompson, C. D. C., Walker, D. A., and Welker, J. M.: Role of Land-Surface Changes in Arctic Summer Warming, Science, 310, 657–660, https://doi.org/10.1126/science.1117368, 2005.

Chen, I.-C., Hill, J. K., Ohlemüller, R., Roy, D. B., and Thomas, C. D.: Rapid Range Shifts of Species Associated with High Levels of Climate Warming, Science, 333, 1024–1026, https://doi.org/10.1126/science.1206432, 2011.

Chini, L., Hurtt, G., Sahajpal, R., Frolking, S., Goldewijk, K. K., Sitch, S., Ganzenmüller, R., Ma, L., Ott, L., Pongratz, J.,
615 and Poulter, B.: Land-use harmonization datasets for annual global carbon budgets, Earth System Science Data, 13, 4175–4189, https://doi.org/10.5194/essd-13-4175-2021, 2021.

Chuvieco, E., Lizundia-Loiola, J., Pettinari, M. L., Ramo, R., Padilla, M., Tansey, K., Mouillot, F., Laurent, P., Storm, T., Heil, A., and Plummer, S.: Generation and analysis of a new global burned area product based on MODIS 250\,m reflectance bands and thermal anomalies, Earth System Science Data, 10, 2015–2031, https://doi.org/10.5194/essd-10-2015-2018, 2018.

620 Claverie, M., Matthews, J. L., Vermote, E. F., and Justice, C. O.: A 30+ Year AVHRR LAI and FAPAR Climate Data Record: Algorithm Description and Validation, Remote Sensing, 8, https://doi.org/10.3390/rs8030263, 2016.

Collier, N., Hoffman, F. M., Lawrence, D. M., Keppel-Aleks, G., Koven, C. D., Riley, W. J., Mu, M., and Randerson, J. T.: The International Land Model Benchmarking (ILAMB) System: Design, Theory, and Implementation, Journal of Advances in Modeling Earth Systems, 10, 2731–2754, https://doi.org/10.1029/2018MS001354, 2018.

625 Dai, A. and Trenberth, K. E.: Estimates of Freshwater Discharge from Continents: Latitudinal and Seasonal Variations, Journal of Hydrometeorology, 3, 660–687, https://doi.org/10.1175/1525-7541(2002)003<0660:EOFDFC>2.0.CO;2, 2002.

Davidson, E. A., de Araújo, A. C., Artaxo, P., Balch, J. K., Brown, I. F., C. Bustamante, M. M., Coe, M. T., DeFries, R. S., Keller, M., Longo, M., Munger, J. W., Schroeder, W., Soares-Filho, B. S., Souza, C. M., and Wofsy, S. C.: The Amazon basin in transition, Nature, 481, 321–328, https://doi.org/10.1038/nature10717, 2012.

630 Dlugokencky, E. and Tans, P.: Trends in atmospheric carbon dioxide, National Oceanic and Atmospheric Administration, Global Monitoring Laboratory (NOAA/GML), 2022.

Dusenge, M. E., Duarte, A. G., and Way, D. A.: Plant carbon metabolism and climate change: elevated CO_2 and temperature impacts on photosynthesis, photorespiration and respiration, New Phytologist, 221, 32–49, 2019.

Elmendorf, S. C., Henry, G. H. R., Hollister, R. D., Björk, R. G., Boulanger-Lapointe, N., Cooper, E. J., Cornelissen, J. H.
635 C., Day, T. A., Dorrepaal, E., Elumeeva, T. G., Gill, M., Gould, W. A., Harte, J., Hik, D. S., Hofgaard, A., Johnson, D. R., Johnstone, J. F., Jónsdóttir, I. S., Jorgenson, J. C., Klanderud, K., Klein, J. A., Koh, S., Kudo, G., Lara, M., Lévesque, E., Magnússon, B., May, J. L., Mercado-Dı´az, J. A., Michelsen, A., Molau, U., Myers-Smith, I. H., Oberbauer, S. F., Onipchenko, V. G., Rixen, C., Martin Schmidt, N., Shaver, G. R., Spasojevic, M. J., Þórhallsdóttir, Þ. E., Tolvanen, A., Troxler, T., Tweedie, C. E., Villareal, S., Wahren, C.-H., Walker, X., Webber, P. J., Welker, J. M., and Wipf, S.: Plot-scale
640 evidence of tundra vegetation change and links to recent summer warming, Nature Climate Change, 2, 453–457, https://doi.org/10.1038/nclimate1465, 2012.

Fisher, R. A. and Koven, C. D.: Perspectives on the Future of Land Surface Models and the Challenges of Representing Complex Terrestrial Systems, Journal of Advances in Modeling Earth Systems, 12, https://doi.org/10.1029/2018MS001453, 2020.

645 Friedl, M. A., Sulla-Menashe, D., Tan, B., Schneider, A., Ramankutty, N., Sibley, A., and Huang, X.: MODIS Collection 5 global land cover: Algorithm refinements and characterization of new datasets, Remote Sensing of Environment, 114, 168–182, https://doi.org/10.1016/j.rse.2009.08.016, 2010.

Friedlingstein, P., O'Sullivan, M., Jones, M. W., Andrew, R. M., Gregor, L., Hauck, J., Le Quéré, C., Luijkx, I. T., Olsen, A., Peters, G. P., Peters, W., Pongratz, J., Schwingshackl, C., Sitch, S., Canadell, J. G., Ciais, P., Jackson, R. B., Alin, S. R.,
650 Alkama, R., Arneth, A., Arora, V. K., Bates, N. R., Becker, M., Bellouin, N., Bittig, H. C., Bopp, L., Chevallier, F., Chini, L. P., Cronin, M., Evans, W., Falk, S., Feely, R. A., Gasser, T., Gehlen, M., Gkritzalis, T., Gloege, L., Grassi, G., Gruber, N., Gürses, Ö., Harris, I., Hefner, M., Houghton, R. A., Hurtt, G. C., Iida, Y., Ilyina, T., Jain, A. K., Jersild, A., Kadono, K., Kato, E., Kennedy, D., Klein Goldewijk, K., Knauer, J., Korsbakken, J. I., Landschützer, P., Lefèvre, N., Lindsay, K., Liu, J., Liu, Z., Marland, G., Mayot, N., McGrath, M. J., Metzl, N., Monacci, N. M., Munro, D. R., Nakaoka, S.-I., Niwa, Y.,
655 O'Brien, K., Ono, T., Palmer, P. I., Pan, N., Pierrot, D., Pocock, K., Poulter, B., Resplandy, L., Robertson, E., Rödenbeck, C., Rodriguez, C., Rosan, T. M., Schwinger, J., Séférian, R., Shutler, J. D., Skjelvan, I., Steinhoff, T., Sun, Q., Sutton, A. J., Sweeney, C., Takao, S., Tanhua, T., Tans, P. P., Tian, X., Tian, H., Tilbrook, B., Tsujino, H., Tubiello, F., van der Werf, G. R., Walker, A. P., Wanninkhof, R., Whitehead, C., Willstrand Wranne, A., et al.: Global Carbon Budget 2022, Earth System Science Data, 14, 4811–4900, https://doi.org/10.5194/essd-14-4811-2022, 2022.

660 Garrigues, S., Lacaze, R., Baret, F., Morisette, J. T., Weiss, M., Nickeson, J. E., Fernandes, R., Plummer, S., Shabanov, N. V., Myneni, R. B., Knyazikhin, Y., and Yang, W.: Validation and intercomparison of global Leaf Area Index products derived from remote sensing data, Journal of Geophysical Research: Biogeosciences, 113, https://doi.org/10.1029/2007JG000635, 2008.

Giglio, L., Randerson, J. T., van der Werf, G. R., Kasibhatla, P. S., Collatz, G. J., Morton, D. C., and DeFries, R. S.:
665 Assessing variability and long-term trends in burned area by merging multiple satellite fire products, Biogeosciences, 7, 1171–1186, https://doi.org/10.5194/bg-7-1171-2010, 2010.

Harris, I., Osborn, T. J., Jones, P., and Lister, D.: Version 4 of the CRU TS monthly high-resolution gridded multivariate climate dataset, Scientific Data, 7, 1–18, https://doi.org/10.1038/s41597-020-0453-3, 2020.

Hartley, A. J., MacBean, N., Georgievski, G., and Bontemps, S.: Uncertainty in plant functional type distributions and its impact on land surface models, Remote Sensing of Environment, 203, 71–89, https://doi.org/10.1016/j.rse.2017.07.037, 2017.

Hengl, T., De Jesus, J. M., Heuvelink, G. B. M., Gonzalez, M. R., Kilibarda, M., Blagotić, A., Shangguan, W., Wright, M. N., Geng, X., Bauer-Marschallinger, B., Guevara, M. A., Vargas, R., MacMillan, R. A., Batjes, N. H., Leenaars, J. G. B., Ribeiro, E., Wheeler, I., Mantel, S., and Kempen, B.: SoilGrids250m: Global gridded soil information based on machine learning, PLoS ONE, 12, e0169748–e0169748, https://doi.org/10.1371/journal.pone.0169748, 2017.

Herrmann, S. M., Anyamba, A., and Tucker, C. J.: Recent trends in vegetation dynamics in the African Sahel and their relationship to climate, Global Environmental Change, 15, 394–404, https://doi.org/10.1016/j.gloenvcha.2005.08.004, 2005.

Hobeichi, S., Abramowitz, G., and Evans, J.: Conserving Land–Atmosphere Synthesis Suite (CLASS), Journal of Climate, 33, 1821–1844, https://doi.org/10.1175/JCLI-D-19-0036.1, 2020.

Houghton, R. A., House, J. I., Pongratz, J., Van Der Werf, G. R., Defries, R. S., Hansen, M. C., Le Quéré, C., and Ramankutty, N.: Carbon emissions from land use and land-cover change, Biogeosciences, 9, 5125–5142, 2012.

Huang, K., Xia, J., Wang, Y., Ahlström, A., Chen, J., Cook, R. B., Cui, E., Fang, Y., Fisher, J. B., Huntzinger, D. N., Li, Z., Michalak, A. M., Qiao, Y., Schaefer, K., Schwalm, C., Wang, J., Wei, Y., Xu, X., Yan, L., Bian, C., and Luo, Y.: Enhanced peak growth of global vegetation and its key mechanisms, Nature Ecology & Evolution, 2, 1897–1905, https://doi.org/10.1038/s41559-018-0714-0, 2018.

Huang, Y., Ciais, P., Santoro, M., Makowski, D., Chave, J., Schepaschenko, D., Abramoff, R. Z., Goll, D. S., Yang, H., Chen, Y., Wei, W., and Piao, S.: A global map of root biomass across the world's forests, Earth System Science Data, 13, 4263–4274, https://doi.org/10.5194/essd-13-4263-2021, 2021.

Hurtt, G. C., Chini, L., Sahajpal, R., Frolking, S., Bodirsky, B. L., Calvin, K., Doelman, J. C., Fisk, J., Fujimori, S., Klein Goldewijk, K., Hasegawa, T., Havlik, P., Heinimann, A., Humpenöder, F., Jungclaus, J., Kaplan, J. O., Kennedy, J., Krisztin, T., Lawrence, D., Lawrence, P., Ma, L., Mertz, O., Pongratz, J., Popp, A., Poulter, B., Riahi, K., Shevliakova, E., Stehfest, E., Thornton, P., Tubiello, F. N., van Vuuren, D. P., and Zhang, X.: Harmonization of global land use change and management for the period 850–2100 (LUH2) for CMIP6, Geoscientific Model Development, 13, 5425–5464, https://doi.org/10.5194/gmd-13-5425-2020, 2020.

Jacobson, A. R., Schuldt, K. N., Miller, J. B., Oda, T., Tans, P., Andrews, A., Mund, J., Ott, L., Collatz, G. J., and Aalto, T.: CarbonTracker CT2019, NOAA Earth System Research Laboratory, Global Monitoring Division, 10, 2020.

Jung, M., Koirala, S., Weber, U., Ichii, K., Gans, F., Camps-Valls, G., Papale, D., Schwalm, C., Tramontana, G., and Reichstein, M.: The FLUXCOM ensemble of global land-atmosphere energy fluxes, Scientific Data, 6, 74, https://doi.org/10.1038/s41597-019-0076-8, 2019.

Jung, M., Schwalm, C., Migliavacca, M., Walther, S., Camps-Valls, G., Koirala, S., Anthoni, P., Besnard, S., Bodesheim, P., Carvalhais, N., Chevallier, F., Gans, F., Goll, D. S., Haverd, V., Köhler, P., Ichii, K., Jain, A. K., Liu, J., Lombardozzi, D., Nabel, J. E. M. S., Nelson, J. A., O'Sullivan, M., Pallandt, M., Papale, D., Peters, W., Pongratz, J., Rödenbeck, C., Sitch, S., Tramontana, G., Walker, A., Weber, U., and Reichstein, M.: Scaling carbon fluxes from eddy covariance sites to globe: synthesis and evaluation of the FLUXCOM approach, Biogeosciences, 17, 1343–1365, https://doi.org/10.5194/bg-17-1343-2020, 2020.

Kato, S., Loeb, N. G., Rose, F. G., Doelling, D. R., Rutan, D. A., Caldwell, T. E., Yu, L., and Weller, R. A.: Surface Irradiances Consistent with CERES-Derived Top-of-Atmosphere Shortwave and Longwave Irradiances, Journal of Climate, 26, 2719–2740, https://doi.org/10.1175/JCLI-D-12-00436.1, 2013.

Klein Goldewijk, K., Beusen, A., Doelman, J., and Stehfest, E.: Anthropogenic land use estimates for the Holocene – HYDE 3.2, Earth System Science Data, 9, 927–953, https://doi.org/10.5194/essd-9-927-2017, 2017.

Kou-Giesbrecht, S. and Arora, V. K.: Representing the Dynamic Response of Vegetation to Nitrogen Limitation via Biological Nitrogen Fixation in the CLASSIC Land Model, Global Biogeochemical Cycles, 36, e2022GB007341, https://doi.org/10.1029/2022GB007341, 2022.

Koven, C. D., Arora, V. K., Cadule, P., Fisher, R. A., Jones, C. D., Lawrence, D. M., Lewis, J., Lindsay, K., Mathesius, S., Meinshausen, M., Mills, M., Nicholls, Z., Sanderson, B. M., Séférian, R., Swart, N. C., Wieder, W. R., and Zickfeld, K.: Multi-century dynamics of the climate and carbon cycle under both high and net negative emissions scenarios, Earth System Dynamics, 13, 885–909, https://doi.org/10.5194/esd-13-885-2022, 2022.

Lenoir, J. and Svenning, J.: Climate-related range shifts–a global multidimensional synthesis and new research directions, Ecography, 38, 15–28, 2015.

Li, X. and Xiao, J.: Mapping Photosynthesis Solely from Solar-Induced Chlorophyll Fluorescence: A Global, Fine-Resolution Dataset of Gross Primary Production Derived from OCO-2, Remote Sensing, 11, https://doi.org/10.3390/rs11212563, 2019.

Liu, Y. Y., Parinussa, R. M., Dorigo, W. A., De Jeu, R. A. M., Wagner, W., van Dijk, A. I. J. M., McCabe, M. F., and Evans, J. P.: Developing an improved soil moisture dataset by blending passive and active microwave satellite-based retrievals, Hydrology and Earth System Sciences, 15, 425–436, https://doi.org/10.5194/hess-15-425-2011, 2011.

Malhi, Y., Roberts, J. T., Betts, R. A., Killeen, T. J., Li, W., and Nobre, C. A.: Climate Change, Deforestation, and the Fate of the Amazon, Science, 319, 169–172, https://doi.org/10.1126/science.1146961, 2008.

Melton, J. R. and Arora, V. K.: Competition between plant functional types in the Canadian Terrestrial Ecosystem Model (CTEM) v. 2.0, Geoscientific Model Development, 9, 323–361, https://doi.org/10.5194/gmd-9-323-2016, 2016.

Melton, J. R., Shrestha, R. K., and Arora, V. K.: The influence of soils on heterotrophic respiration exerts a strong control on net ecosystem productivity in seasonally dry Amazonian forests, Biogeosciences, 12, 1151–1168, https://doi.org/10.5194/bg-12-1151-2015, 2015.

Melton, J. R., Arora, V. K., Wisernig-Cojoc, E., Seiler, C., Fortier, M., Chan, E., and Teckentrup, L.: CLASSIC v1.0: The open-source community successor to the Canadian Land Surface Scheme (CLASS) and the Canadian Terrestrial Ecosystem Model (CTEM)-Part 1: Model framework and site-level performance, Geoscientific Model Development, 13, 2825–2850, https://doi.org/10.5194/gmd-13-2825-2020, 2020.

Monerie, P.-A., Wainwright, C. M., Sidibe, M., and Akinsanola, A. A.: Model uncertainties in climate change impacts on Sahel precipitation in ensembles of CMIP5 and CMIP6 simulations, Climate Dynamics, 55, 1385–1401, https://doi.org/10.1007/s00382-020-05332-0, 2020.

Mortimer, C., Mudryk, L., Derksen, C., Luojus, K., Brown, R., Kelly, R., and Tedesco, M.: Evaluation of long-term Northern Hemisphere snow water equivalent products, The Cryosphere, 14, 1579–1594, https://doi.org/10.5194/tc-14-1579-2020, 2020.

Myneni, R. B., Hoffman, S., Knyazikhin, Y., Privette, J. L., Glassy, J., Tian, Y., Wang, Y., Song, X., Zhang, Y., Smith, G. R., Lotsch, A., Friedl, M., Morisette, J. T., Votava, P., Nemani, R. R., and Running, S. W.: Global products of vegetation leaf area and fraction absorbed PAR from year one of MODIS data, Remote Sensing of Environment, 83, 214–231, https://doi.org/10.1016/S0034-4257(02)00074-3, 2002.

National Research Council: Earth observations from space: The first 50 years of scientific achievements, National Academies Press, 2008.

Olsson, L., Eklundh, L., and Ardö, J.: A recent greening of the Sahel—trends, patterns and potential causes, Journal of Arid Environments, 63, 556–566, https://doi.org/10.1016/j.jaridenv.2005.03.008, 2005.

Parry, I. M., Ritchie, P. D. L., and Cox, P. M.: Evidence of localised Amazon rainforest dieback in CMIP6 models, Earth System Dynamics, 13, 1667–1675, https://doi.org/10.5194/esd-13-1667-2022, 2022.

Pastorello, G., Trotta, C., Canfora, E., Chu, H., Christianson, D., Cheah, Y.-W., Poindexter, C., Chen, J., Elbashandy, A., Humphrey, M., Isaac, P., Polidori, D., Reichstein, M., Ribeca, A., van Ingen, C., Vuichard, N., Zhang, L., Amiro, B., Ammann, C., Arain, M. A., Ardö, J., Arkebauer, T., Arndt, S. K., Arriga, N., Aubinet, M., Aurela, M., Baldocchi, D., Barr, A., Beamesderfer, E., Marchesini, L. B., Bergeron, O., Beringer, J., Bernhofer, C., Berveiller, D., Billesbach, D., Black, T. A., Blanken, P. D., Bohrer, G., Boike, J., Bolstad, P. V., Bonal, D., Bonnefond, J.-M., Bowling, D. R., Bracho, R., Brodeur, J., Brümmer, C., Buchmann, N., Burban, B., Burns, S. P., Buysse, P., Cale, P., Cavagna, M., Cellier, P., Chen, S., Chini, I., Christensen, T. R., Cleverly, J., Collalti, A., Consalvo, C., Cook, B. D., Cook, D., Coursolle, C., Cremonese, E., Curtis, P. S., D'Andrea, E., da Rocha, H., Dai, X., Davis, K. J., Cinti, B. D., Grandcourt, A. de, Ligne, A. D., De Oliveira, R. C., Delpierre, N., Desai, A. R., Di Bella, C. M., Tommasi, P. di, Dolman, H., Domingo, F., Dong, G., Dore, S., Duce, P., Dufrêne, E., Dunn, A., Dušek, J., Eamus, D., Eichelmann, U., ElKhidir, H. A. M., Eugster, W., Ewenz, C. M., Ewers, B., Famulari, D., Fares, S., Feigenwinter, I., Feitz, A., Fensholt, R., Filippa, G., Fischer, M., Frank, J., Galvagno, M., et al.: The FLUXNET2015 dataset and the ONEFlux processing pipeline for eddy covariance data, Scientific Data, 7, 225, https://doi.org/10.1038/s41597-020-0534-3, 2020.

Pearson, R. G., Phillips, S. J., Loranty, M. M., Beck, P. S. A., Damoulas, T., Knight, S. J., and Goetz, S. J.: Shifts in Arctic vegetation and associated feedbacks under climate change, Nature Climate Change, 3, 673–677, https://doi.org/10.1038/nclimate1858, 2013.

Pecl, G. T., Araújo, M. B., Bell, J. D., Blanchard, J., Bonebrake, T. C., Chen, I.-C., Clark, T. D., Colwell, R. K., Danielsen, F., and Evengård, B.: Biodiversity redistribution under climate change: Impacts on ecosystems and human well-being, Science, 355, eaai9214, 2017.

Phillips, O. L., Aragão, L. E. O. C., Lewis, S. L., Fisher, J. B., Lloyd, J., López-González, G., Malhi, Y., Monteagudo, A., Peacock, J., Quesada, C. A., van der Heijden, G., Almeida, S., Amaral, I., Arroyo, L., Aymard, G., Baker, T. R., Bánki, O., Blanc, L., Bonal, D., Brando, P., Chave, J., de Oliveira, Á. C. A., Cardozo, N. D., Czimczik, C. I., Feldpausch, T. R., Freitas, M. A., Gloor, E., Higuchi, N., Jiménez, E., Lloyd, G., Meir, P., Mendoza, C., Morel, A., Neill, D. A., Nepstad, D., Patiño, S., Peñuela, M. C., Prieto, A., Ramírez, F., Schwarz, M., Silva, J., Silveira, M., Thomas, A. S., Steege, H. ter, Stropp, J., Vásquez, R., Zelazowski, P., Dávila, E. A., Andelman, S., Andrade, A., Chao, K.-J., Erwin, T., Di Fiore, A., C., E. H., Keeling, H., Killeen, T. J., Laurance, W. F., Cruz, A. P., Pitman, N. C. A., Vargas, P. N., Ramírez-Angulo, H., Rudas, A., Salamão, R., Silva, N., Terborgh, J., and Torres-Lezama, A.: Drought Sensitivity of the Amazon Rainforest, Science, 323, 1344–1347, https://doi.org/10.1126/science.1164033, 2009.

Port, U., Brovkin, V., and Claussen, M.: The influence of vegetation dynamics on anthropogenic climate change, Earth System Dynamics, 3, 233–243, https://doi.org/10.5194/esd-3-233-2012, 2012.

Reich, P. B., Wright, I. J., Cavender-Bares, J., Craine, J. M., Oleksyn, J., Westoby, M., and Walters, M. B.: The Evolution of Plant Functional Variation: Traits, Spectra, and Strategies, International Journal of Plant Sciences, 164, S143–S164, https://doi.org/10.1086/374368, 2003.

Renwick, K. M. and Rocca, M. E.: Temporal context affects the observed rate of climate-driven range shifts in tree species, Global Ecology and Biogeography, 24, 44–51, https://doi.org/10.1111/geb.12240, 2015.

Riahi, K., Vuuren, D. P. van, Kriegler, E., Edmonds, J., O'Neill, B. C., Fujimori, S., Bauer, N., Calvin, K., Dellink, R., Fricko, O., Lutz, W., Popp, A., Cuaresma, J. C., KC, S., Leimbach, M., Jiang, L., Kram, T., Rao, S., Emmerling, J., Ebi, K., Hasegawa, T., Havlik, P., Humpenöder, F., Silva, L. A. D., Smith, S., Stehfest, E., Bosetti, V., Eom, J., Gernaat, D., Masui, T., Rogelj, J., Strefler, J., Drouet, L., Krey, V., Luderer, G., Harmsen, M., Takahashi, K., Baumstark, L., Doelman, J. C., Kainuma, M., Klimont, Z., Marangoni, G., Lotze-Campen, H., Obersteiner, M., Tabeau, A., and Tavoni, M.: The Shared Socioeconomic Pathways and their energy, land use, and greenhouse gas emissions implications: An overview, Global Environmental Change, 42, 153–168, https://doi.org/10.1016/j.gloenvcha.2016.05.009, 2017.

Rödenbeck, C., Zaehle, S., Keeling, R., and Heimann, M.: How does the terrestrial carbon exchange respond to inter-annual climatic variations? A quantification based on atmospheric CO_2 data, Biogeosciences, 15, 2481–2498, https://doi.org/10.5194/bg-15-2481-2018, 2018.

Santoro, M., Beaudoin, A., Beer, C., Cartus, O., Fransson, J. E. S., Hall, R. J., Pathe, C., Schmullius, C., Schepaschenko, D., Shvidenko, A., Thurner, M., and Wegmüller, U.: Forest growing stock volume of the northern hemisphere: Spatially explicit estimates for 2010 derived from Envisat ASAR, Remote Sensing of Environment, 168, 316–334, https://doi.org/10.1016/j.rse.2015.07.005, 2015.

Schepaschenko, D., Chave, J., Phillips, O. L., Lewis, S. L., Davies, S. J., Réjou-Méchain, M., Sist, P., Scipal, K., Perger, C., Herault, B., Labrière, N., Hofhansl, F., Affum-Baffoe, K., Aleinikov, A., Alonso, A., Amani, C., Araujo-Murakami, A., Armston, J., Arroyo, L., Ascarrunz, N., Azevedo, C., Baker, T., Bałazy, R., Bedeau, C., Berry, N., Bilous, A. M., Bilous, S. Yu., Bissiengou, P., Blanc, L., Bobkova, K. S., Braslavskaya, T., Brienen, R., Burslem, D. F. R. P., Condit, R., Cuni-Sanchez, A., Danilina, D., del Castillo Torres, D., Derroire, G., Descroix, L., Sotta, E. D., d'Oliveira, M. V. N., Dresel, C., Erwin, T., Evdokimenko, M. D., Falck, J., Feldpausch, T. R., Foli, E. G., Foster, R., Fritz, S., Garcia-Abril, A. D., Gornov, A., Gornova, M., Gothard-Bassébé, E., Gourlet-Fleury, S., Guedes, M., Hamer, K. C., Susanty, F. H., Higuchi, N., Coronado, E. N. H., Hubau, W., Hubbell, S., Ilstedt, U., Ivanov, V. V., Kanashiro, M., Karlsson, A., Karminov, V. N., Killeen, T., Koffi, J.-C. K., Konovalova, M., Kraxner, F., Krejza, J., Krisnawati, H., Krivobokov, L. V., Kuznetsov, M. A., Lakyda, I., Lakyda, P. I., Licona, J. C., Lucas, R. M., Lukina, N., Lussetti, D., Malhi, Y., Manzanera, J. A., Marimon, B., Junior, B. H. M., Martinez, R. V., Martynenko, O. V., Matsala, M., Matyashuk, R. K., Mazzei, L., Memiaghe, H., Mendoza, C., Mendoza, A. M., Moroziuk, O. V., Mukhortova, L., Musa, S., Nazimova, D. I., Okuda, T., Oliveira, L. C., Ontikov, P. V., et al.: The Forest Observation System, building a global reference dataset for remote sensing of forest biomass, Scientific Data, 6, 198, https://doi.org/10.1038/s41597-019-0196-1, 2019.

Scholze, M., Knorr, W., Arnell, N. W., and Prentice, I. C.: A climate-change risk analysis for world ecosystems, Proceedings of the National Academy of Sciences, 103, 13116–13120, https://doi.org/10.1073/pnas.0601816103, 2006.

Seiler, C., Melton, J., Arora, V., and Wang, L.: CLASSIC v1.0: the open-source community successor to the Canadian Land Surface Scheme (CLASS) and the Canadian Terrestrial Ecosystem Model (CTEM) – Part 2: Global Benchmarking, Geoscientific Model Development, 14, 2371–2417, https://doi.org/10.5194/gmd-2020-294, 2021.

Seiler, C., Melton, J. R., Arora, V. K., Sitch, S., Friedlingstein, P., Anthoni, P., Goll, D., Jain, A. K., Joetzjer, E., Lienert, S., Lombardozzi, D., Luyssaert, S., Nabel, J. E. M. S., Tian, H., Vuichard, N., Walker, A. P., Yuan, W., and Zaehle, S.: Are Terrestrial Biosphere Models Fit for Simulating the Global Land Carbon Sink?, Journal of Advances in Modeling Earth Systems, 14, e2021MS002946, https://doi.org/10.1029/2021MS002946, 2022.

825 Sitch, S., Huntingford, C., Gedney, N., Levy, P., Lomas, M., Piao, S., Betts, R., Ciais, P., Cox, P., and Friedlingstein, P.: Evaluation of the terrestrial carbon cycle, future plant geography and climate-carbon cycle feedbacks using five Dynamic Global Vegetation Models (DGVMs), Global change biology, 14, 2015–2039, 2008.

Stackhouse, P. W. J., Gupta, S. K., Cox, S. J., Zhang, T., Mikovitz, C., and Hinkelman, L. M.: The NASA/GEWEX surface radiation budget release 3.0: 24.5-year dataset, Gewex news, 21, 10–12, 2011.

830 Still, C. J., Berry, J. A., Collatz, G. J., and DeFries, R. S.: Global distribution of C3 and C4 vegetation: Carbon cycle implications, Global Biogeochemical Cycles, 17, 6–1, https://doi.org/10.1029/2001GB001807, 2003.

Strahler, A. H. and Muller, J.-P.: MODIS BRDF/Albedo Product: Algorithm Theoretical Basis Document Version 5.0, 1999.

Swann, A. L., Fung, I. Y., Levis, S., Bonan, G. B., and Doney, S. C.: Changes in Arctic vegetation amplify high-latitude warming through the greenhouse effect, Proceedings of the National Academy of Sciences, 107, 1295–1300,
835 https://doi.org/10.1073/pnas.0913846107, 2010.

Swart, N. C., Cole, J. N. S., Kharin, V. V., Lazare, M., Scinocca, J. F., Gillett, N. P., Anstey, J., Arora, V., Christian, J. R., Hanna, S., Jiao, Y., Lee, W. G., Majaess, F., Saenko, O. A., Seiler, C., Seinen, C., Shao, A., Sigmond, M., Solheim, L., and Von, K.: The Canadian Earth System Model version 5 (CanESM5.0.3), Geoscientific Model Development, 12, 4823–4873, https://doi.org/10.5194/gmd-12-4823-2019, 2019.

840 Tape, K., Sturm, M., and Racine, C.: The evidence for shrub expansion in Northern Alaska and the Pan-Arctic, Global change biology, 12, 686–702, 2006.

Thomas, C. D.: Climate, climate change and range boundaries, Diversity and Distributions, 16, 488–495, 2010.

Todd-Brown, K. E. O., Randerson, J. T., Post, W. M., Hoffman, F. M., Tarnocai, C., Schuur, E. A. G., and Allison, S. D.: Causes of variation in soil carbon simulations from CMIP5 Earth system models and comparison with observations,
845 Biogeosciences, 10, 1717–1736, https://doi.org/10.5194/bg-10-1717-2013, 2013.

Verger, A., Baret, F., and Weiss, M.: Near Real-Time Vegetation Monitoring at Global Scale, IEEE Journal of Selected Topics in Applied Earth Observations and Remote Sensing, 7, 3473–3481, https://doi.org/10.1109/JSTARS.2014.2328632, 2014.

Verseghy, D. L.: CLASS — A Canadian land surface scheme for GCMS. I. Soil model, International Journal of
850 Climatology, 11, 111–133, 1991.

Verseghy, D. L., McFarlane, N. A., and Lazare, M.: CLASS — A Canadian land surface scheme for GCMS, II. Vegetation model and coupled runs, International Journal of Climatology, 13, 347–370, https://doi.org/10.1002/joc.3370130402, 1993.

Vialet-Chabrand, S. R. M., Matthews, J. S. A., McAusland, L., Blatt, M. R., Griffiths, H., and Lawson, T.: Temporal Dynamics of Stomatal Behavior: Modeling and Implications for Photosynthesis and Water Use, Plant Physiology, 174, 603–
855 613, https://doi.org/10.1104/pp.17.00125, 2017.

Walther, G.-R.: Community and ecosystem responses to recent climate change, Philosophical Transactions of the Royal Society B: Biological Sciences, 365, 2019–2024, https://doi.org/10.1098/rstb.2010.0021, 2010.

Wang, A., Price, D. T., and Arora, V.: Estimating changes in global vegetation cover (1850–2100) for use in climate models, Global Biogeochemical Cycles, 20, https://doi.org/10.1029/2005GB002514, 2006.

860 Wang, L., Arora, V. K., Bartlett, P., Chan, E., and Curasi, S. R.: Mapping of ESA's Climate Change Initiative land cover data to plant functional types for use in the CLASSIC land model, Biogeosciences, 20, 2265–2282, https://doi.org/10.5194/bg-20-2265-2023, 2023.

Warszawski, L., Frieler, K., Huber, V., Piontek, F., Serdeczny, O., and Schewe, J.: The inter-sectoral impact model intercomparison project (ISI–MIP): project framework, Proceedings of the National Academy of Sciences, 111, 3228–3232,
865 2014.

Wieder, W.: Regridded Harmonized World Soil Database v1.2, https://doi.org/10.3334/ORNLDAAC/1247, 2014.

Wu, D., Zhao, X., Liang, S., Zhou, T., Huang, K., Tang, B., and Zhao, W.: Time-lag effects of global vegetation responses to climate change, Global change biology, 21, 3520–3531, 2015.

Wu, Z., Dijkstra, P., Koch, G. W., Peñuelas, J., and Hungate, B. A.: Responses of terrestrial ecosystems to temperature and
870 precipitation change: A meta-analysis of experimental manipulation, Global Change Biology, 17, 927–942, 2011.

Xue, B.-L., Guo, Q., Hu, T., Wang, G., Wang, Y., Tao, S., Su, Y., Liu, J., and Zhao, X.: Evaluation of modeled global vegetation carbon dynamics: Analysis based on global carbon flux and above-ground biomass data, Ecological Modelling, 355, 84–96, https://doi.org/10.1016/j.ecolmodel.2017.04.012, 2017.

Zemp, D. C., Schleussner, C.-F., Barbosa, H. M. J., Hirota, M., Montade, V., Sampaio, G., Staal, A., Wang-Erlandsson, L.,
875 and Rammig, A.: Self-amplified Amazon forest loss due to vegetation-atmosphere feedbacks, Nature Communications, 8, 14681, https://doi.org/10.1038/ncomms14681, 2017.

Zhang, Y. and Liang, S.: Fusion of Multiple Gridded Biomass Datasets for Generating a Global Forest Aboveground Biomass Map, Remote Sensing, 12, 2559, https://doi.org/10.3390/rs12162559, 2020.

Zhang, Y., Xiao, X., Wu, X., Zhou, S., Zhang, G., Qin, Y., and Dong, J.: A global moderate resolution dataset of gross
880 primary production of vegetation for 2000-2016, Scientific data, 4, 1–13, https://doi.org/10.1038/sdata.2017.165, 2017.