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Reducing uncertainties of future global soil carbon responses to climate and land

use change with emergent constraints

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Key Points:

- The uncertainty in Soil organic carbon (SOC) change is dominated by differences between model structure rather than by climate forcing.
- Soil input changes explain most variations in projected SOC change for natural vegetation across models at global and region.
- The effective reduction in constrained SOC change depends on climate forcing and region considered.

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1 Abstract

2 Soil organic carbon changes (\triangle SOC) are regulated by climate and land use change. Here, we 3 analyze regional and global ΔSOC from 1861 to 2099 based on five Terrestrial Biosphere Models (TBMs) simulations of the Inter-Sectorial Impact Model Inter-comparison Project 4 Phase 2b. The TBMs were driven by harmonized gridded land use change and bias-adjusted 5 6 climate forcing data from different General Circulation Models (GCMs) for climate scenarios 7 RCP2.6 and RCP6.0. Between 2005 and the end of this century, we estimated an increase of SOC for two scenarios with large uncertainty, which is dominated by differences between 8 9 TBMs. We present a new emergent constraint approach to constrain future modeled \triangle SOC over natural vegetation from RCP6.0 simulations using recent observed trends of net primary 10 11 productivity as a proxy of litter inputs to soil pools. Our results showed that the uncertainties in constrained \triangle SOC can be reduced in comparison with the original model ensemble, but 12 13 constrained values of \triangle SOC depend on the choice of a GCM and climate regions. For the reduction of the SOC density in areas where cropland expanded ($\Delta soc_{cropland\ expansion}$) over 14 natural vegetation as a result of land use change, the constrained $\Delta soc_{cropland\ expansion}$ still 15 features large uncertainties due to uncertain observed data. Our proposed emergent constraint 16 17 approach appears to be valuable to reduce uncertainty on SOC projections, but it is limited here by the small number of models (five) and by the uncertainty in the observational data. 18 Applications to larger ensembles from Earth System Models should be tested for the future. 19

1. Introduction

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Soil organic carbon (SOC) is the largest carbon pool in the terrestrial biosphere, containing 2.3~5.3 times more carbon than the vegetation and the atmosphere (Ciais et al. 2014). Due to its large pool size and gross exchange fluxes representing annually more than 10% of the mass of carbon in the atmosphere, soil carbon plays a very important role in regulating the global carbon cycle. Soil organic carbon change (Δ SOC) is controlled by input from plants (litter, exudates), by lateral fluxes (e.g. erosion of particulate organic matter, dissolved organic matter runoff) and by the rate of soil organic matter decomposition (Todd-Brown et al., 2013; Carvalhais et al., 2014; Yan et al., 2014; Wu et al., 2018). These processes are affected by climate and land-use change. Land-use change over the last century was dominated by the conversion of forests and natural grasslands to cropland and pasture. During this process, SOC inputs are reduced because most of agricultural net primary production (NPP) is lower than that of natural systems (Kolby Smith et al., 2014; Neumann and Smith, 2018) and because only the non-harvested fraction of agricultural NPP is returned to SOC (Haberl et al., 2007; Krausmann et al., 2008). Climate change through temperature and precipitation changes, directly modifies both carbon input rates to SOC and decomposition rates. Quantifying and separating the effects of climate change and land use change on SOC change at regional and global scales to improve future Δ SOC projections is a key research challenge, which has implications for mitigation solutions based on increasing soil organic matter stocks in soils. Observational data are valuable to quantify the global spatial distribution of SOC (Harmonized World Soil Database (HWSD; FAO/IIASA/ISRIC/ISSCAS/JRC, 2012), the Northern Circumpolar Soil Carbon Database (NCSCD; Tarnocai et al., 2009), the World Inventory of Soil Emission Potentials (WISE30sec; Batjes, 2016), the Unified North American Soil Map (UNASM; Liu et al., 2013), and the SoilsGrids250 database (Hengl et al., 2017)). For evaluating, historical changes of SOC, there are only few sites where long-term measurements are available, especially for natural ecosystems. Meta-analysis of SOC changes after land use change were reported by previous studies (Post and Kwon, 2000; Guo and Gifford 2002; Li et al., 2017; 2018). Observed regional changes of SOC from inventories

are reported by some studies (Bellamy et al., 2005; Hamdi et al., 2013; Doetterl et al., 2015).

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Modeled estimates of global and regional SOC changes during the the historical period have been reported by terrestrial biosphere models (TBMs; e.g. Tian et al., 2015b) and in some cased data-driven models, e.g. Sanderman et al., (2017) for grasslands SOC losses due to past land use. For future projections, estimations of SOC changes are based on TBMs coupled with General Circulation Models (GCMs) or run offline with climate forcing from GCM simulations. Projections of SOC from coupled models are particularly uncertain (Nishina et al., 2014; Todd-Brown et al., 2014; Koven et al., 2015; Luo et al., 2016), partly because models are not well calibrated and evaluated against observed data (Xiao et al., 2014; Luo et al., 2016), and partly because carbon cycle coupled models have climate biases. Global SOC stocks were found to vary from 510 to 3040 Pg C in the period 1995-2005 among 11 models of the Coupled Model Inter-comparison Project 5 (CMIP5) (Todd-Brown et al., 2013). This large range was attributed to differences in model structure, parameter values, and climate input fields. To better understand the different sources of model uncertainties, model-to-model variation in Δ SOC was decomposed into uncertainties due to initial SOC stocks (the SOC stocks during 1997-2006), relative changes in soil inputs and decomposition rates / turnover times following ideas proposed by Todd-Brown et al., (2014) and Koven et al., (2015). Compared to ensembles of SOC simulations from fully coupled GCMs that differ in their climate, ensembles of SOC simulations from offline TBMs forced by bias-adjusted climate forcing data allow us to focus on structural errors of TBMs. Over the historical period (i.e., 2010) during which climate forcing can be obtained from observations to drive TBMs, Tian et al., (2015b) analyzed SOC from 10 models of the Multi-scale Synthesis and Terrestrial Model Inter-comparison Project (MsTMIP). They found that i) the magnitude of SOC stocks ranged from 425 to 2111 Pg C across models, slightly narrower the range (510-3040 Pg C during 1995-2005) of TBMs reported by Todd-Brown et al., (2013) and ii) cumulative SOC changes during the historical period differed from -70 to 86 PgC. This large spread suggests that model structural errors are dominant in both initial SOC stock and SOC changes simulations. Up to now, no study has linked systematic errors of modeled SOC

change errors between the historical period and future projections.

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The emergent constraint approach allows using historical simulations and observed data to reduce uncertainty in future projections of earth system variables (Hall et al., 2019). This approach relies on the assumption that historical or present-day differences between models and observed data are preserved in future projections and reflect stationary differences explained by models' structure. Thus, if we can estimate an effective emergent constraint using contemporary observations, it helps to downweigh less realistic models and reduce the spread of the ensemble. This approach was applied to constrain for instance snow albedo temperature sensitivities (Hall et al., 2019), tropical carbon cycle sensitivity to warming (Cox et al., 2013), global ratio of plant transpiration to total terrestrial evapotranspiration (Lian et al., 2018), future yield changes (Zhao et al., 2016), and CO₂ fertilization of land photosynthesis (Wenzel et al., 2016). Such an approach was also applied to reduce the uncertainty in projections of permissible emissions for climate stabilization (Jones et al., 2006). In this study, we attempt to apply a new emergent constraint approach to reduce uncertainties related to future SOC changes (Δ SOC) by an ensemble of offline terrestrial carbon cycle models, hereafter called terrestrial biosphere models or TBMs. Specifically, we aim to:

- (1) Compare Δ SOC in past and future from five different ISIMIP2b terrestrial biosphere models (LPJ-GUESS, LPJmL, VISIT, ORCHIDEE-MICT and DLEM) forced by the same set of bias-adjusted climate forcing from different climate models under two different greenhouse gas concentration pathways (RCP 2.6 and RCP 6.0) and corresponding land use scenarios.
- (2) Quantify the contributions of initial soil carbon, changes in decomposition rate, and changes in soil inputs to the model spread of Δ SOC in natural ecosystems, that is ecosystems where the climate and CO₂ perturbation dominates SOC changes.
- (3) Reduce the model spread of ΔSOC in natural ecosystems caused by climate and CO_2 driven soil carbon inputs changes, using observed input changes approximated by NPP trends, with an emergent constraint approach.

(4) Reduce the model spread of Δ SOC in ecosystems where land use conversion of natural ecosystems to croplands has been driving SOC, using observations of SOC densities changes before and after land use, with an emergent constraint approach.

2. Methods

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2.1 ISIMIP2b biome models and simulation set-up

The Inter-Sectoral Impact Model Intercomparison Project Phase 2b (ISIMIP2b) provides simulations of TBMs driven with several bias-adjusted climate fields and land use change scenarios for the period from 1861 to 2099 (Frieler et al., 2017). The ISIMIP2b models were driven by gridded, daily bias-adjusted climate from different CMIP5 GCMs (Lange 2016; Frieler et al., 2017), global annual atmospheric CO₂ concentration, and harmonized annual land use maps (Klein Goldewijk et al., 2017). Models performed a spin up to simulate land carbon pools in 1860 as described in the protocol (https://www.isimip.org/protocol/#isimip2b). The use of bias-adjusted climate data ensures that TBMs are forced by climate that match observations in the last 40 years of the historical period, and that there is no discontinuity of climate forcing between the past and the future. Note however that decadal and inter-annual variations of the ISIMIP2b climate forcing do not match observed climate variability since variability follows the one of each GCM. Decadal and inter-annual climate variability as well as historical climate trends thus differ between bias-adjusted GCMs. The key point is that the use of common bias-adjusted climate forcing for the historical period and the future in this study reduces the spread in SOC projections from TBMs compared to using TBMs fully coupled with climate models that have considerable climate differences. This makes it possible for us to focus on structural uncertainties from TBMs, and yet to examine the impact of different GCMs and scenarios for the future.

Five TBMs from the ISIMIP2b biome sector were used (Table S1): LPJ-GUESS (Smith et al., 2014), LPJmL (Bondeau et al., 2007), VISIT (Ito and Inatomi 2012), DLEM (Tian et al., 2015a), and ORCHIDEE-MICT (Guimberteau et al., 2018). These models differ in their biogeochemical parameterizations and thus in their simulated response of SOC to climate and land use change (Table S1; Text S1) but they nevertheless share the same

philosophy for their soil carbon modules using first order kinetics equations applied to one to three pools adjusted by soil temperature and moisture. SOC stock at soil depth of 0-1 m was calculated from carbon mass in soil pool (litter was not included) based on ISIMIP2b simulations. All models simulated carbon cycling in terrestrial ecosystem with different discretization of vegetation into plant functional types (PFTs). Three models (ORCHIDEE-MICT, LPJmL and DLEM) include permafrost. None of the models includes wetlands (Table S1). We selected TBM output from simulations driven by bias-adjusted daily climate forcing of four different GCMs: GFDL-ESM2M, HadGEM2-ES, IPSL-CM5A-LR and MIROC5 (Frieler et al., 2017) at a spatial resolution of 0.5°×0.5° for the RCP 2.6 and RCP 6.0. An exception is ORCHIDEE-MICT, which used climate forcing at a resolution of 1.0°×1.0° and its results were downscaled to 0.5°×0.5°.

Historical land use change (LUC) forcing for ISIMIP2b was derived from the LUH2 gridded reconstruction based on HYDE3.2 data (Klein Goldewijk et al., 2017). The LUH2 data was further disaggregated into annual land use maps with major crop types, rainfed and irrigated (Monfreda et al., 2008) for ISIMIP2b. Future land use change forcing was based on projections from the MAgPIE land use model (Popp et al., 2014; Stevanović et al., 2016) assuming population growth and economic development following the SSP2 storyline (Popp et al., 2017) and including climate change impacts on crop yields estimated by the LPJmL crop model (Müller and Robertson, 2014) for each RCP scenario (Frieler et al., 2017). To ensure continuity of spatially explicit land use change forcing from historical to future period, the LUH2 harmonization method was applied (Frieler et al., 2017).

Variable agricultural area (cropland and pasture) from LUH2 was used as input to TBMs. Each model started from the same 1860 agricultural area from LUH2 and non-harmonized pre-industrial natural vegetation distributions, and used different transition rules for converting a fraction of natural vegetation to LUH2 agriculture land (or vice-versa) in each grid cell, each year. The models did not report SOC for each PFT in each grid cell, which would have allowed a precise evaluation of SOC changes in agricultural land use type vs. natural PFTs, separately. To overcome this limitation, we calculated 'ΔSOC from cropland dominated areas' by selecting only grid cells where the cropland fraction is larger than 30%

in 2005 (Figure S1).

Three groups of simulations defined by the ISIMIP2b protocol were analyzed (Frieler et al., 2017; Table 1). Group 1 contains simulations driven by historical climate and land use change during 1861-2005. Group 2 contains simulations driven by future climate change with a fixed future land use map equal to that of year 2005. Group 2 simulations are thus driven only by climate change and named after CC. Group 3 contains simulations driven by both future climate and land use change (hereafter, CC+LUC). The difference of SOC between Group 2 and Group 3 simulations gives the effect of future land use change (LUC) assuming drivers are additive.

<<Table 1>>

We separated the analysis of ΔSOC between grid-cells dominated by cropland (cropland fraction more than 30% in 2005) and grid-cells with no or little cropland (cropland fraction less than 30% in 2005), defined as 'natural vegetation'. 54.3% of these natural vegetation grids cells still include small cropland fractions (83.7% out of the 54.3% have a cropland fraction lower than 15.0%; Figure S1f). Grid cells dominated by historical cropland summed up to 10.0% of the global land grid cells (Figure S1f). For each of these two categories, a separate approach is used to constrain ΔSOC with different types of observations.

2.2 Constraining ASOC in areas dominated by cropland

Bookkeeping land use models, data-driven models and TBMs indicate that agricultural expansion caused a net soil carbon loss in the past (Hansis et al., 2015; Houghton and Nassikas 2017; Li et al., 2017; Sanderman et al., 2017). Generally, after conversion to cropland there is a SOC loss during the first years because cultivated land has a lower NPP than natural ecosystems (Kolby Smith et al., 2014; Neumann and Smith 2018), and because agricultural NPP is harvested and tillage accelerates SOC decomposition.

To constrain historical Δ SOC from cropland dominated grid cells (Figure 1a) by observations, we hypothesized that there is a strong relationship between i) Δ SOC per unit area over the grid cells dominated by historical cropland expansion (Δ soc_{cropland expansion},

calculated as ΔSOC divided by the increased area of cropland) and ii) the difference of SOC 192 per unit area between cropland and initial natural vegetation 193 $(\Delta soc_{cropland\ minus\ initial\ natveg})$. For example, in the case of a grid cell that was 100% 194 covered by natural vegetation in 1861 and is now 100% covered by cropland, 195 196 $\Delta soc_{cropland\ expansion}$ is strictly equal to $\Delta soc_{cropland\ minus\ initial\ natveg}$. The idea is that $\Delta soc_{cropland\ minus\ initial\ nativeg}$ can be obtained from observations of SOC density across 197 different land use types in the same region, and then used to constrained modeled output of 198 $\Delta soc_{cropland\ expansion}$. The principle of this emergent constraint is illustrated in Figure 1a. It 199 should be noted that the $\Delta soc_{cropland\ minus\ initial\ natveg}$ will be affected by transient 200 climate, especially by transient CO₂, which would lead to a bias. 201 202 To verify the above hypothesis, we established regional regressions between

 $\Delta soc_{cropland\ expansion}$ and $\Delta soc_{cropland\ minus\ initial\ natveg}$ for each model between 1861 and 2005. Eight regions were considered (Eurasia, North America, South America, West Eurasia, Australia, South Asia, and East Asia; Figure S1c-d). In each region, $\Delta soc_{cropland\ minus\ initial\ natveg}$ was calculated as the difference of soil carbon stocks densities of grid cells dominated by cropland grid cells with a cropland fraction larger than 50% in 2005 and grid cells with a natural vegetation fraction higher than 50% in 1861 in each region of Figure 1c-d, based on historical simulations (Group 1). The choice of a 50% fraction threshold was made by considering the tradeoff between a strong relationship between $\Delta soc_{cropland\ expansion}$ and $\Delta soc_{cropland\ minus\ initial\ natveg}$ (Figure S2) and a sufficient number of dominated cropland grids number (Figure S1e). The results of these regressions confirmed that $\Delta soc_{cropland\ expansion}$ and $\Delta soc_{cropland\ minus\ initial\ natveg}$ are indeed strongly positively correlated across different TBMs (see in Sect. 3.3.3). Thus, it is justified to constrain modeled $\Delta soc_{cropland\ expansion}$ by observations of $\Delta soc_{cropland\ minus\ initial\ nature\ q}$ using the emergent relationship illustrated in Figure 1a. We compiled field observations of the SOC density (Deng et al., 2016; Li et al., 2018; Nyawira et al., 2016), hereafter referred to as soc, for natural vegetation and cropland in each region, and calculated $\Delta soc_{cropland\ minus\ initial\ natveg}$ as their difference; Figure S3).

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During the selection of field soc data, two criteria were considered: (1) studies must report cropland soc measurements; (2) soc measurements must come from paired adjacent sites, one with natural vegetation type and the other with cropland. Overall, 274 paired data were selected from 147 study sites (Figure S1c) to assess $\Delta soc_{cropland\ minus\ initial\ natureg}$. Those data were further classified into four transitions types involving cropland in different climate regions, including forest to cropland transitions in tropical region (F-C, Trop; n=78), forest to cropland transitions in temperate region (F-C, Temp; n=49), grassland to cropland in tropical region (G-C, Trop; n=15) and grassland to cropland in temperate region (G-C, Temp; n=132). In order to constrain modeled $\Delta soc_{cropland\ expansion}$ by observed $\Delta soc_{cropland\ minus\ initial\ natureg}$, we selected the dominant type of transition to croplands in each region from the models (Figure S1b-d; Table 5) and corresponded it with observed $\Delta soc_{cropland\ minus\ initial\ natureg}$ in the climate zone of each region (Figure S1b-d).

2.3 Constraining ΔSOC in areas of natural vegetation where SOC change is dominated by climate change

In grid cells dominantly covered by natural vegetation with a cropland fraction less than 30% in 2005, we assumed that Δ SOC can be mainly explained by climate- and CO₂-induced shifts in the balance between litter input, and decomposition rates (Todd-Brown et al., 2013). Todd-Brown et al., (2014) showed that model-to-model variation in Δ SOC across the CMIP5 models could be explained (R²=0.89, p<0.01) by differences in initial soil carbon stocks combined with relative changes in soil inputs and decomposition rates. We used the same attribution method to quantify the impact of the three key variables on Δ SOC from ISIMIP2b models, based on Group 2 simulations (CC). Todd-Brown et al., (2014) assumed that Δ SOC from transient ESM model runs is equal to the difference of their equilibrium SOC pools between the end and the start of each run, so that Δ SOC can be written as:

$$\Delta SOC = C_{end} - C_{start} = \frac{I_{end}}{k_{end}} - \frac{I_{start}}{k_{start}}$$
 (1)

Where I is the soil carbon input approximated by net primary productivity (NPP), k the decomposition rate and calculated from global heterotrophic respiration divided by soil carbon stocks; and subscripts *end* and *start* are for the initial and final state of a simulation.

Here we consider the period of 2040-2049 for RCP 2.6 and 2090-2099 for RCP 6.0 as the final state, and the period of 1995-2005 as the initial state. The choice of 2040-2049 as the final state for RCP 2.6 is because in this scenario, atmospheric CO₂ concentration that drives the positive trend of NPP and soil C inputs through the CO₂ fertilization effect present in all TBMs, peaks by 2050s and decreases thereafter (Meinshausen et al., 2011). After that date, decreasing CO₂ may cause a decrease of NPP and soil C input, inducing a decrease of SOC with a time delay, which complicates the use of equation (1). Equation (1) can be rearranged into:

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$$\Delta SOC = C_{end} - C_{start} = \left(\frac{1 + \frac{\Delta I}{I_{start}}}{1 + \frac{\Delta k}{k_{start}}} - 1\right) \times C_{start}$$
 (2)

Using regression analysis of modeled ΔSOC with the terms on the right-hand side of Eq. (2), we assessed the relative contributions of changes in soil inputs $(1 + \frac{\Delta I}{I_{start}})$, changes in decomposition rate $(1 + \frac{\Delta k}{k_{start}})$, and initial soil carbon stocks (C_{start}) to the modeled ΔSOC . The regression is executed across different TBMs for each GCM (i.e., one regression for each GCM) averaging all variables over grid cells with natural vegetation.

We used a two-steps emergent constraint (Figure 1b). The first step is to constrain future $\frac{\Delta I}{I_{start}}$ from past NPP trends, if there is a strong enough linear relationship between these two variables across TBMs. To test for such a relationship, we established linear regressions between future $\frac{\Delta I}{I_{start}}$ and past NPP trends during 2001-2015 from the different models, and then, we used the observed NPP trends to constrain $\frac{\Delta I}{I_{start}}$. In the second step, we established linear regressions between future Δ SOC and $1 + \frac{\Delta I}{I_{start}}$ from models from Eq. 2, and then used the constrained future $\frac{\Delta I}{I_{start}}$ from the first step to constrain future Δ SOC. This strategy is summarized in Figure 1b. We constrain Δ SOC for global natural vegetation and each climate region using the same emergent constraint than above.

The hypothesis behind this two-step emergent constraint is that future carbon input changes can be constrained from observation-based trends of past NPP. The trends of

observed NPP were derived from trends of observation-based photosynthesis (GPP) from 274 gridded datasets assuming a constant ratio of NPP to GPP equal to 0.45 (He et al., 2018). GPP 275 trends were estimated from two data-driven models that include both the effect of rising CO₂ 276 277 on photosynthesis and satellite observed trends of leaf area (Jiang and Ryu 2016; Wang et al., 278 2017). Note that we did not to use trends of satellite based NPP models based on AVHRR greenness data and light-use efficiency (LUE) models (Kolby Smith et al., 2015) and on 279 MODIS (Zhao and Running 2010) because their LUE formulation ignores the fertilization 280 281 effect of increasing CO₂ and thus likely underestimates NPP trends in this approach (De 282 Kauwe et al., 2016). 283 The two GPP data-driven models are the P-model (Wang et al., 2017; Stocker et al., 2019) and the breathing earth system simulator (BESS) model simulations (Ryu et al., 2011; 284 Jiang and Ryu 2016) during the period of 2001-2015. The P-model is a LUE model in which 285 286 LUE is depends on environmental condition (air temperature, vapor pressure deficit, 287 elevation) and CO₂ concentrations, with an optimality principle that predicts stomatal conductance and foliar photosynthetic traits based on a standard model for C3 leaf 288 photosynthesis. The bias of the P-model for global GPP is 3.81% (Wang et al., 2017). BESS 289 290 is a process-based GPP model that use remotely sensed data of land surface and air 291 temperature, leaf area index (LAI), CO₂ concentrations and canopy information. The bias of 292 BESS for global GPP is 1.92% (Jiang and Ryu 2016). Significant increase in NPP are produced by those two data-driven approaches. Models in natural ecosystem during the 293 period of 2001-2015, with a trend of 0.11 Pg C yr⁻² in BESS and a trend of 0.21 Pg C yr⁻² in 294 295 P-Model. Larger increase in NPP was found in P-model for tropical, temperate and boreal regions (Table S2). 296 297 In the two-step emergent constraint approach illustrated in Figure 1b, uncertainties in constrained Δ SOC are a function of uncertainties in litter carbon input trend constrained by 298 the observed trend of NPP, and in the linear regression slopes of regressions between $\frac{\Delta I}{I_{start}}$ 299 and past input changes, and between ΔSOC and $(1 + \frac{\Delta I}{I_{start}})$ are considered. The uncertainty in 300 constrained \triangle SOC is calculated as in Stegehuis et al., (2013) by: 301

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$$\sigma_{\Delta SOC} = \sqrt{\beta^2 \sigma_{\frac{\Delta I}{I_{start}}}^2 + \sigma_{res_\Delta SOC}^2}$$
 (3)

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$$\sigma_{\frac{\Delta I}{I_{start}}} = \sqrt{\alpha^2 \sigma_{obs}^2 + \sigma_{res_obs}^2}$$
 (4)

- where $\sigma_{\Delta SOC}$, $\sigma_{\frac{\Delta I}{I_{start}}}$ and σ_{obs} are the uncertainties in constrained ΔSOC , the
- uncertainties in $\frac{\Delta I}{I_{start}}$ and uncertainties in the past NPP trend based on two datasets. β and
- 306 $\sigma_{res_\Delta SOC}$ indicate the slope and standard deviation of the residuals from linear regression
- between \triangle SOC and $1 + \frac{\Delta I}{I_{start}}$. Similarly, α and σ_{res_obs} present the linear regression between
- 308 $\frac{\Delta I}{I_{start}}$ and past NPP trend.
- Last, in the attribution of \triangle SOC differences for natural vegetation between models
- 310 given by equation (2), the term related to initial SOC stocks differences across models can
- also be constrained from observations. Three global SOC datasets were used for this purpose,
- the HWSD (FAO/IIASA/ISRIC/ISSCAS/JRC, 2012), the NCSCD (Tarnocai et al., 2009) and
- 313 the WISE30sec (Batjes 2016).

3. Results

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3.1 Changes in the modeled global soil carbon

- We found large differences in simulated global \triangle SOC for the historical period (1861-
- 317 2005; Group 1 simulations), ranging from -81.3 Pg C (LPJmL driven by IPSL-CM5-LR
- 318 climate) to 88.8 Pg C (VISIT driven by HadGEM2-ES climate; Figure 2; Table 2). For the
- future period of 2006-2099 in Group 2 simulations with CC+LUC effects, model differences
- of global ΔSOC are also large, going from -9.4 Pg C (LPJmL driven by HadGEM2-ES
- 321 climate forcing) to 114.7 Pg C (VISIT driven by GFDL-ESM2M climate) for RCP 2.6 and
- 322 from -30.1 Pg C (LPJmL driven by IPSL-CM5-LR climate) to 176.5 Pg C (VISIT driven by
- 323 MIROC5 climate) for RCP 6.0 (Figure 2; Table 2).
- 324 <<Figure 2>>
- 325 <<Table 2>>
- The interquartile range (IQR, the difference between 75th and 25th percentile of the

data) of future ΔSOC across all GCMs forcing data and TBMs is larger for RCP 6.0 (81.9 Pg C) than that for RCP 2.6 (73.9 Pg C; Figure 3a, Table S3). The larger IQR of ΔSOC for RCP 6.0 is partly explained by diverging model responses to climate change alone, with an IQR of 69.6 Pg C from the effects of climate change alone in RCP 6.0 compared to a climate change induced IQR of 19.9 Pg C in RCP 2.6 (Figure 3b; Table S3). The difference in IQR of ΔSOC between RCP 6.0 (IQR=14.2) and RCP 2.6 (IQR=25.5) was reduced when considering the effects of LUC alone, even though a larger IQR was found in RCP 2.6. The IQR of ΔSOC caused by different GCMs forcing, obtained by averaging all TBMs outputs for the same GCM, is smaller than the IQR across TBMs, with an IQR across GCMs of 15.2 Pg C in RCP 2.6 and 30.6 Pg C in RCP 6.0 (Figure 3a). The spread of ΔSOC is thus mainly due to structural differences in TBMs, with an IQR of 56.8 Pg C across TBMs for RCP 2.6 and 73.7 Pg C for RCP 6.0 (fourth column of Figure 3a). The relative shares of both GCM versus TBM-related uncertainties are similar for both scenarios (Figure 3a-c).

<< Figure 3>>

Under effects of CC+LUC, the change of SOC during 2006-2099 is a net increase of 41.8 ± 43.9 Pg C ($3.2 \pm 3.4\%$) for RCP 2.6, and of 48.5 ± 63.3 Pg C ($3.8 \pm 4.8\%$) for RCP 6.0 across all TBMs and GCMs (Figure 2; Table 2). Climate change alone caused larger SOC changes of 46.8 ± 58.2 Pg C ($3.4 \pm 4.3\%$) under RCP 6.0 than 15.3 ± 20.8 Pg C ($1.2 \pm 1.3\%$) under RCP 2.6 (Table 2). In addition, LUC alone after 2005 caused a SOC increase of 28.2 ± 31.4 Pg C under RCP 2.6 and 2.7 ± 5.9 Pg C under RCP 6.0. The LUC forcing alone impacted future Δ SOC in LPJmL, ORCHIDEE-MICT, DLEM and VISIT, but had no obvious effect in LPJ-GUESS (Figure S4a-b).

Given the fact that differences in the projected Δ SOC are partly driven by different trends in NPP (see equation 2), we further examined the simulated evolution of NPP with time (Figure 3; Figure S4-5 for each model). All models simulated increased NPP in the future, of 7.3 \pm 3.2 Pg C yr⁻¹ (11.9 \pm 5.2%) under RCP 2.6 and of 18.3 \pm 4.9 Pg C yr⁻¹ (29.7 \pm 7.9%) under RCP 6.0 driven by CC+LUC (Figure 3d; Table S4). Similar to the uncertainty of Δ SOC (here, expressed as IQR), the uncertainty of Δ NPP mainly come from differences of TBMs and from the two RCP scenarios rather than from differences of GCM forcing, a result

consistent with the dominant attribution of uncertainties on ΔSOC to differences of TBMs (Figure 3d).

An accelerated decomposition rate (increase of k; positive Δk) of global SOC was simulated by all models, with a mean increase of $2.8 \pm 1.0 \ 10^{-3} \ yr^{-1}$ ($-8.0 \pm 3.9\%$) under RCP $2.6 \ and \ 7.7 \pm 1.8 \ 10^{-3} \ yr^{-1}$ ($21.0 \pm 4.5\%$) under RCP 6.0, respectively (Figure 3g; Table S5). Similar to the variations in the simulated global Δ SOC and global Δ NPP, the spread of global decomposition rate (Δk) among simulations were mostly attributed to differences in TBMs and RCPs rather than to differences of GCMs (Figure 3g).

3.2 Contribution of initial soil carbon, decomposition rate, soil inputs to soil carbon changes of natural ecosystems

For the RCP 6.0 scenario, we decomposed model differences of Δ SOC into differences explained by soil inputs, decomposition rates, initial soil carbon stocks using Equation 2. We found that these three variables altogether explain 84% - 91% of the variation in global \triangle SOC across TBMs, this range being from the different GCMs (all p < 0.1; Figure 4). The initial soil carbon stocks and change in decomposition rate do not show significant correlation with global ΔSOC among TBMs, for any GCM. Instead, most of the ΔSOC differences between TBMs can be explained by their different changes in soil inputs (Figure 4d, i, n, s). Different changes in soil inputs explain 52% - 89% of the global ΔSOC across the five TBMs, depending on the GCM considered (Figure 4). Regression slopes (between original modeled ΔSOC and predicted values from Eq. 2) are similar between GCMs (ranging from 0.63 to 0.76; Figure 4a, f, k, p). For the RCP 2.6 scenario, no significant relationship between simulated \triangle SOC and predicted values from Eq. 2 was found across the different TBMs (Figure S6). This is because in RCP 2.6, climate change is small and CO₂ concentration increases much less (63.9 ppm from 2005-2050) compared to RCP6.0 (287.6 ppm from 2005-2099) and does not produce a change of SOC large enough to be attributed to the factors considered in equation 2.

<< Figure 4>>

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Using Eqn. 2, we further separated ΔSOC for tropical, temperate and boreal regions (Figure 5-7). The relative changes in soil input, decomposition rates, and initial soil carbon

stocks altogether explain 81% - 96% of tropical Δ SOC (slopes from 0.58 to 0.83), 81% - 96% of temperate Δ SOC (slopes from 0.47 to 0.71), and 75% - 95% of boreal Δ SOC (slopes range from 0.26 to 0.45) (Figure 5-7). In the tropical region, most of Δ SOC differences between TBMs can be attributed to differences in changes of soil inputs (R²=0.43-0.92) and initial soil carbon (R²=0.44-0.90). For simulations driven by HadGEM2-ES GCM in the tropical region, initial soil carbon can explain more of the Δ SOC differences between TBMs than change in soil inputs (Figure 5). In both temperate and boreal region, differences in changes in soil inputs explain most of the differences in Δ SOC across TBMs (Figure 6-7).

393 <<Figure 5>>

394 <<Figure 6>>

395 <<Figure 7>>

3.3 Constraining future \triangle SOC by observations

3.3.1 Constrained global Δ SOC for natural vegetation

As explained in Section 2.3, we used observed NPP trends to constrain Δ SOC over grid cells dominated by natural vegetation (Figure 1b) for simulations driven by RCP 6.0. We recall here that the results of RCP 2.6 are not shown because they produced small Δ NPP and Δ SOC except for MIROC5 climate forcing in the tropical region (Figure S6-S9), and thus are not suitable for applying our emergent constraint approach (see Methods). We found significant linear relationships between modeled $\frac{\Delta I}{I_{start}}$ and modeled NPP trend during 2001-2015 across TBMs (R² ranging from 0.85 to 0.95) for three out of four GCM forcing (GFDL-ESM2M, HadGEM2-ES, and IPSL-CM5A-LR). This is shown in Figure 4, the last column of plots. For MIROC5, the relationship between modeled $\frac{\Delta I}{I_{start}}$ and modeled NPP trend was not significant (p=0.20; R² = 0.47; Figure 4t). Here, we use observed NPP trends (grey areas) and the linear relationships above to constrain future input changes $\frac{\Delta I}{I_{start}}$ (purple areas in Figure 4e, j, o, t). Then, the constrained $\frac{\Delta I}{I_{start}}$ (purple areas in the fourth column of Figure 4) was used to constrain Δ SOC (green areas in Figure 4d, i, n, s), according to the principle

illustrated in Figure 1b.

Using the two-step emergent constraint with all uncertainties propagated (Eqs. 3 and 4), we constrained global Δ SOC values of 25.2 \pm 42.4 Pg C with GFDL-ESM2M, -36.9 \pm 67.3 Pg C with HadGEM2-ES, 1.4 \pm 42.6 Pg C with IPSL-CM5A-LR, and 38.8 \pm 54.1 Pg C with MIROC5 (Table 3; Figure 4). These constrained global Δ SOC values were all lower than the original ensemble means of Δ SOC (Figure 4; Table 3). For HadGEM2-ES, the constrained global Δ SOC was even constrained to be a net loss whereas it was simulated as a gain in the original ensemble mean of the TBMs (19.0 \pm 53.9 Pg C). We acknowledged that uncertainties of observed NPP trends combined with uncertainties in the regressions led to only a marginal uncertainty reduction of constrained vs. original Δ SOC, given the small set of TBMs examined in this study. The linear relationships shown in Figure 4 were based only on five TBMs, and a larger ensemble of models should make the emergent constraint more effective, with more expected model outliers. Here, we found that 15 out of 19 of the original Δ SOC simulations were within 1-sigma uncertainty of constrained Δ SOC, one outlier being the VISIT model (Figure 4; Table 3).

<<Table 3>>

3.3.2 Constrained regional ΔSOC for natural vegetation

Significant linear relationships between modeled $\frac{\Delta I}{I_{start}}$ and modeled NPP trends during 2001-2015 were found in all GCMs in the temperate region for natural vegetation (Figure 6e, j, o, t) and in one GCM (i.e., HadGEM2-ES) forcing in the boreal region (Figure 7j), but not in the tropical region (Figure 5e, j, o, t). Constrained temperate Δ SOC showed large differences between different GCMs (Table 3; Figure 6), ranging from -96.5 \pm 61.7 Pg C for HadGEM2-ES to 65.5 \pm 44.2 Pg C for GFDL-ESM2M. Constrained Δ SOC in the temperate region can be either higher than (driven by GFDL-ESM2M and MIROC5 climate) or lower than (driven by HadGEM2-ES and IPSL-CM5A-LR climate) the original model ensemble mean (Table 3). The uncertainty range of constrained temperate Δ SOC is slightly reduced with GFDL-ESM2M, IPSL-CM5A-LR, and MIROC5 climate (by 3.6 – 13.2 Pg C). With HadGEM2-ES climate, the uncertainty range of constrained Δ SOC was not reduced,

because observed NPP trends are much smaller than in the model ensemble forced by this GCM, and constrained Δ SOC is extrapolated outside the range of TBMs in Table 3. For temperate region only 10 out of 19 constrained Δ SOC values were within the 1-sigma uncertainty of the original Δ SOC ensemble, which indicates that observed NPP trends imply a strong change of constrained vs. original SOC changes, and thus that the quality of observational data is critical.

The lower constrained Δ SOC values obtained for HadGEM2-ES climate compared to other GCMs, globally and for the temperate and boreal regions is due to the much higher regression slope and lower intercept between $\frac{\Delta I}{I_{start}}$ and observed NPP trend for this GCM, which caused lower constrained $1 + \frac{\Delta I}{I_{start}}$ and thus lower constrained Δ SOC. In the tropical region, however, the constrained Δ SOC for HadGEM2-ES is significantly higher than under the other GCMs because less TBMs used this GCM; DLEM did not provide HadGEM2-ES GCM output and Figure S10 shows the results after excluding DLEM. In addition, the uncertainty of the constrained Δ SOC from HadGEM2-ES is larger than those from other GCMs, due to the high regression slope between Δ SOC and $1 + \frac{\Delta I}{I_{start}}$ (simulations driven by HadGEM2-ES have the highest slopes globally, and in tropical and temperate regions) and the uncertainty of constrained $\frac{\Delta I}{I_{start}}$ is determined by the regression slope between $\frac{\Delta I}{I_{start}}$ and modeled NPP trend.

We found no significant relationships between ΔSOC and initial soil carbon across TBMs at global scale (Figure 4), which prevent us using observed global initial SOC stocks to constrain future ΔSOC . However, there are positive relationships in the tropics, but significant only for the HadGEM2-ES GCM (R²=0.90, p = 0.05; Figure 5g). A lower constrained ΔSOC was found from HadGEM2-ES (-4.8 \pm 2.3 Pg C) than in the original ensembles (4.7 \pm 21.1 Pg C) using initial SOC stocks as a constraint. Such constrained ΔSOC was smaller than that constrained by NPP trend (14.4 \pm 21.6 Pg C; Figure 5; Table 3). It should be noted that none of the constrained ΔSOC values were within a 1-sigma uncertainty of the original ΔSOC ensemble in such constrained ΔSOC .

3.3.3 Constrained future ΔSOC from cropland expansion

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We found a significant positive relationship between the simulated SOC density changes across grid-cells with cropland expansion $\Delta soc_{cropland\ expansion}$ and the soil carbon density difference between cropland in 2005 and initial natural vegetation in 1861, Δsoc_{cropland minus initial native} in all LUC regions except for East Asia (Figure 8). The coefficient of determination (R²) of those relationships are high in all regions (ranging from 0.79 to 0.97 across regions) except for South Asia (R^2 =0.49). That means the Δ SOC due to LUC outweighs \triangle SOC due to climate change over grid cells with cropland expansion. The 473 slopes of the relationships between $\Delta soc_{cronland\ expansion}$ and $\Delta soc_{cropland\;minus\;initial\;natveg}\;$ are different in each region, ranging from 0.90 kg C m^-2 (kg C m⁻²)⁻¹ in South Asia to 3.15 kg C m⁻² (kg C m⁻²)⁻¹ in West Eurasia. Following the emergent constraint principle described in the method section, we constrain $\Delta soc_{cropland\ expansion}$ from observed values of $\Delta soc_{cropland\ minus\ initial\ natveg}$. << Figure 8>>

In LUC dominated areas, the constrained global $\Delta soc_{cropland\ expansion}$ was -0.29 \pm 5.56 kg C m^{-2} which is a smaller loss than the original modeled range of $-0.61 \pm 2.15 \text{ kg C m}^{-1}$ ² but this constrained value has a large uncertainty (Figure 8; Figure S3; Table 4). However, larger constrained carbon loss ($\Delta soc_{cropland\ expansion}$) than from the unconstrained simulations were found in Eurasia, North America, South America, Africa, West Eurasia and Australia (Table 4). For example, constrained $\Delta soc_{cronland\ expansion}$ is of -2.78 \pm 9.92 kg C m^{-2} in North America compared to the original mean value of -1.43 \pm 2.85 kg C m^{-2} for that region. Constrained $\Delta soc_{cropland\ expansion}$ is a loss of -3.45 \pm 12.84 kg C m⁻² in Australia compared to the original ensemble mean of 0.45 ± 2.53 kg C m⁻². In addition, large differences in constrained $\Delta soc_{cropland\ expansion}$ also can be found in regions characterized by carbon losses, ranging from -5.94 \pm 17.68 kg C m⁻² (West Eurasia) to -1.66 \pm 2.27 kg C m⁻² ² (Africa) .We found that all simulated $\Delta soc_{cropland\ expansion}$ is within 1-sigma uncertainty of the constrained values in LUC region (Figure 8; Table 4) but the spread of constrained $\triangle soc$ is always larger than that in the original model ensemble.

By multiplying these constrained changes of SOC densities by the area of historical

cropland expansion, we constrained a carbon loss induced by cropland expansion of -1.03 \pm 19.94 Pg C during 1861-2005 (Table 5). There are differences between regions of historical cropland expansion, with a maximum loss in North America (-2.79 \pm 9.96 Pg C), a minimum loss in South America (-0.36 \pm 1.92 Pg C), and a neutral carbon change in East Asia (0.05 \pm 0.01 Pg C) and South Asia (0.42 \pm 1.53 Pg C).

Future regional SOC changes associated with cropland expansion were constrained by multiplying future cropland expansion areas (Figure S11) by constrained estimates of $\Delta soc_{cropland\ expansion}$ (Table 5). We inferred small future carbon losses from future cropland expansion for RCP 2.6 and RCP 6.0 in Eurasia, North America, South America, Africa, West Eurasia, and Australia, and small carbon gains in South Asia and East Asia (Table 5). In North America, no cropland expansion occurs under RCP 6.0. Overall, there is no large cropland expansion. Global constrained carbon losses from future cropland expansion are -0.19 \pm 3.72 Pg C for RCP 2.6 and -0.18 \pm 3.52 Pg C for RCP 6.0, respectively.

509 <<Table 4>>

510 <<Table 5>>

3.4 Comparison of initial SOC stocks for natural vegetation

We compared initial soil carbon stocks between modeled and observed datasets for grid-cells with natural vegetation. The mean global soil carbon stock over the period of 1995-2005 across all GCMs and TBMs was 1421.6 Pg C with a range of 702.6-2008.3 Pg C (Table 6). The mean value was higher than that from three datasets of 1202.4 Pg C, but the observed range of 1094.8-1283.9 Pg C was much smaller than the spread of models, indicating that even with the same climate forcing, models are inconsistent with the observed SOC stocks (Table 6). Yet, there was a slightly smaller spread between ISIMIP2b TBMs than the CMIP5 ESMs, that gave a mean value of 1520 Pg C and a range of 510~3040 Pg C (Todd-Brown et al., 2013). DLEM forced by MIROC5 has the lowest SOC (702.6 Pg C) and ORCHIDEE-MICT forced by IPSL-CM5A-LR has the highest value (2008.3 Pg C). In temperate region, SOC from all GCMs and TBMs had a median value of 762.8 Pg C with an IQR of 708.5 Pg

C, slightly higher but comparable to the observed value (mean 717.3 Pg C with a range of 645.9~758.0 Pg C across datasets). Modeled SOC in the tropical region had a median value of 381.6 Pg C with an IQR of 231.6 Pg C, consistent with observed datasets (mean of 371.8 Pg C with a range of 362.4~376.5 Pg C). Modeled SOC in the boreal region took a median value of 110.8 Pg C with an IQR of 238.6 Pg C, consistent with observed datasets (mean of 113.3 Pg C with a range of 72.6-159.3 Pg C). Largest IQR of SOC in temperate region rather than in tropical and boreal region was found, which indicate that large uncertainties in SOC of TBMs comes from the temperate region.

<<Table 6>>

4. Discussion

4.1 Large modeled differences of projected future ΔSOC

Previous studies reported that a large range of initial or present-day SOC stocks simulated by TBMs when they are coupled to climate model or run offline with the same climate forcing, going from 510 Pg C to 3040 Pg C for 11 TBMs part of the CMIP5 ESMs (Todd-Brown et al., 2014) and from 425 Pg C to 2111 Pg C among 10 offline TBMs in MsTMIP (Tian et al., 2015b). The CMIP5 models have biases in climate causing a bias of SOC, whereas the MsTMIP models only covered the historical period. In this study, historical and future projections of SOC with bias-adjusted climate and harmonized land use change forcing make it possible to examine ΔSOC driven by climate and land use change continuously for the historical period and the future.

We found that simulated global ΔSOC during the historical period (1861-2005) is a small increase, with a median value of 16.51 Pg C and a large range going from -81.3 to 88.8 Pg C (Table 2). This result from ISIMIP2b models is higher but comparable to MsTMIP that gave a median change of 3.39 Pg C from 1901 to 2010, with a range of -70.2 – 85.9 Pg C (Tian et al., (2015b). Besides, our results show that most models project a future global SOC increase under the RCP 2.6 scenario (2005-2099; median value of 31.91 Pg C) and under RCP 6.0 scenario (2005-2099; median value of 31.40 Pg C). The global ΔSOC reported here is slightly higher but narrower than estimates from 11 ESMs presented in Todd-Brown et al.,

(2014) for the RCP 8.5 scenario, with a median increase of 15 Pg C and a range of -72 – 253 Pg C.

Our results showed that uncertainties in Δ SOC were more attributed to structural differences between TBMs rather than to differences in GCMs (Figure 3). The ISMIP2b TBMs used the same protocol for spin-up, the same input data, land use change data, and climate forcing (Tian et al., 2015b; Frieler et al., 2017). Yet, even if GCM bias adjustment was applied to match observed mean climate period 1960-1990 (Hempel et al., 2013) the different GCM forcing data have distinct climate variability for the historical period, and differences in future climate as well (Table S6) due to different GCMs climate sensitivities. Furthermore, the vegetation distribution was not harmonized between models which introduced inconsistencies in the simulation of SOC between models.

4.2 Uncertainties in constrained Δ SOC for dominated area by natural vegetation

It is known that future ΔSOC is sensitive to climate via both changes in soil input and decomposition (Jones et al., 2005; Todd-Brown et al., 2013; Carvalhais et al., 2014; Yan et al., 2014). Our results showed that soil input changes explain most of the simulated ΔSOC across the ISIMIP2b TBMs at global scale and also in different regions, the rest of variation being explained by the interaction between initial soil carbon stocks, decomposition rate and changes in soil inputs for RCP 6.0 (Figure 4-7). Such results are consistent with previous study with 11 ESMs from CMIP5 (Todd-Brown et al., 2014).

We showed that the success of using observed recent NPP trend as a constraint for future ΔSOC over natural vegetation depends on the choice of the GCM. Our proposed emergent constrained worked well with the HadGEM2-ES in the boreal region and with all GCMs in the temperate region. It also works for constraining global ΔSOC , with a given GCM but provides diverging results between different GCMs. This is because GCMs differ in their regional patterns of climate change, with possible compensating effects of climate change on ΔSOC between different regions, for instance decreased rainfall reducing input and increased temperature increasing them. The failure to reliably constrain ΔSOC at the regional scale for the tropical and boreal regions with most GCMs may be attributed to the uncertainties in observed NPP trends and the weak relationships established for the emergent

constraint when it is applied at regional scale. The NPP trend in the P-model is stronger than in the BESS model (Table S2), possibly because nutrient limitations are not included in the P-model (Jiang and Ryu 2016; Wang et al., 2017). Previous studies indicated that NPP is strongly limited by N availability in many ecosystems, especially in boreal forests (Hickler et al., 2015). In addition, observed NPP trend based on satellite observation includes implicitly the management of forest and pasture, which is ignored in ISIMIP2b models. Due to the small number of TBMs used to establish emergent constraint relationships, with only five TBMs in this study, the R^2 of linear regressions between NPP trend and future input change (0.27-0.51) were low in the tropical region (Figure 5). It should be noted that an effective constrained Δ SOC is based on a significant relationship between Δ SOC and changes in soil input. Applying the same approach for CMIP5 and CMIP6 models should offer larger ensembles with likely a larger spread and a better-defined relationship to constrain SOC changes.

4.3 Uncertainties in constrained $\triangle soc$ for dominated area by cropland

We showed that soil carbon losses in LUC dominated regions are mostly caused by conversion of natural vegetation to cropland, consistent with previous studies (Guo and Gifford 2002; Don et al., 2011; Poeplau et al., 2011; Wei et al., 2014; Deng et al., 2016; Li et al., 2018). For example, the meta-analysis by Guo and Gifford (2002) found a SOC decline of 42% and 59% after LUC from forest to cropland and pasture to cropland. Models that do not reduce soil C input from harvested cropland NPP tend to underestimate SOC reductions from cropland expansion (-2.64 \pm 8.43 Pg C vs. -1.03 \pm 19.94 Pg C for modeled vs. estimated $\Delta soc_{cropland\ expansion}$; Table 5). In addition, we found that the simulated $\Delta soc_{cropland\ expansion}$ is generally within 1-sigma uncertainty of the constrained $\Delta soc_{cropland\ expansion}$. Such large uncertainties in constrained $\Delta soc_{cropland\ expansion}$ reflect uncertainties of $\Delta soc_{cropland\ minus\ initial\ natureg}$ from meta-analysis data (Table 4).

Previous studies indicated that anthropogenic land use changes have resulted into about 50 million km² being used for cropland (about 12% of the total ice-free land area) and pasture (about 26% of the total ice-free land area) (Foley et al., 2007; 2011). In ISIMIP2b simulations, the area of cropland and pasture increases in both the past the future (Figure

S11-S12; Table S7). For example, the cropland area expands from 5.9×10^6 km² in 1861 to 14.6×10^6 km² in 2005 (Table S7).

Based on constrained Δ*soc_{cropland expansion*} and future cropland area, we tentatively suggest that future cropland expansion will result in carbon losses in most of regions (Table 5; Figure S11-S12). These results are consistent with previous studies reporting that land use will be an important driver of SOC in the future (Lozano-Garcia et al., 2017; Molotoks et al., 2018). The degree of cropland expansion results in SOC change partially dependent upon the land management practices (i.e., harvest), soil condition (e.g., soil properties and soil type) and climate condition. Crop harvest for bioenergy production can reduce inputs to the soil and diminish soil fertility (Powlson et al., 2011). In general, SOC decreases when land use conversion is from forest to cropland, but varies with forest type and cultivation stage (Wei et al., 2014). The conversion from natural vegetation to cropland breaks down the aggregate structure that physically protects SOC from microbial decomposition (Wei et al., 2013), leading to more available SOC for microbial attacks. Future increased temperature will result in greater SOC losses by increasing decomposition rate but also cause a positive feedback between SOC mineralization and global warming.

Previous studies have highlighted the importance of accounting for agricultural land and management (e.g., harvest, grazing, tillage, residue management) in model simulations (Pugh et al., 2015). In this study, modeled SOC from cropland expansion showed large difference between models, which are related to different cropland management schemes and implies large uncertainty in future projections. Theoretically, crop harvest reduces carbon input into soil. In this study, four models (i.e., VISIT, LPJmL, ORCHIDEE-MICT and DLEM) include cropland harvest in their parameterization. LPJmL, ORCHIDEE-MICT and DLEM actually consider that harvested carbon is released as CO_2 to the atmosphere. As a result, negative $\Delta soc_{cropland\ minus\ initial\ natveg}$ were observed from those three models that do not allocate crop harvest to soil pools (Figure S3). However, the residual part of the harvest NPP of cropland from VISIT return to field as litter, which explains positive $\Delta soc_{cropland\ minus\ initial\ natveg}$ values in most regions. The LPJ-GUESS version used in ISIMIP2b did not consider crop harvest. Therefore, how to treat crop harvest and the

management of crop residue is a key source of uncertainty for modeling SOC changes from cropland expansion in the ISIMIP2b models. We recommend a better treatment of harvested crop carbon in TBMs, accounting for the harvest flux to leave the system rather than allocating harvests to SOC pools.

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In addition, it is important to note that not all models have taken into account tillage process except for ORCHIDEE-MICT. ORCHIDEE-MICT considers the effect of tillage by increasing decomposition rate in cropland to mimic increased soil oxygenation and accelerated decomposition of SOC after tillage (Gervois et al., 2008). In general, tillage can improve the decomposition of crop residues by facilitating contact between plant tissue and soil aggregate surface (Bronick and Lal, 2005) and increases the availability of nutrients for plant growth through distributing organic matter. In conjunction with this, the effectiveness of tillage on SOC in comparison to no-tillage is controversial (Angers and Eriksen-Hamel, 2008; Virto et al., 2012). Several studies observed an increase of soil organic matter and carbon with no-tillage or conservation tillage (minimum tillage) in the top soil layer (Vogeler et al., 2009; Powlson et al., 2012; Pinheiro et al., 2015). However, such effect is partly or completely offset by greater SOC content in the deeper soil layers under conventional tillage (complete inversion of soil through ploughing; Álvaro-Fuentes et al., 2013). These discrepancies are not surprising since tillage effects integrate a complex set of biological and environmental factors, such as the management practices (e.g., fertilization; Gregorich et al., 2005), crop performance (e.g., cropping intensity and crop types; VandenBygaart et al., 2003), and climate conditions (e.g., soil temperature and soil moisture; Snyder et al., 2009). In the TBMs used here, the effect of tillage is either represented as a scaling factor increasing the SOC decomposition rate in ORCHIDEE-MICT (Gervois et al., 2008) or ignored in other models. A newer version of LPJmL now incorporates two processes directly affected by tillage, including surface litter reduction from tillage management and decreased bulk soil density affecting soil hydrology (Lutz et al., 2019).

4.4 How to improve emergent constraints on soil carbon changes

The use of an emergent constraint to constrain an ensemble of model results requires

i) a strong regression relationship between the target variable to be constrained and the

variable used to predict this target (e.g., NPP trends and $\Delta soc_{cropland\ minus\ initial\ nativeg})$ and ii) available observed data. If a strong enough emergent relationship can be established, then it can be confidently combined with observation to produce a constrained target variable. In our study, the hypothesis behind this two-step emergent constraint for SOC changes of natural vegetation is that future carbon input changes can be constrained from observation-based trends of past NPP. This hypothesis was verified but observed trends of NPP were different between the two products considered. The hypothesis to constrain $\Delta soc_{cropland\ expansion}$ is that this target variable is related to mean soc differences between cropland and historical natural vegetation. This hypothesis was verified but observed $\Delta soc_{cropland\ minus\ initial\ natureg}$ shows a large spread from current meta-analysis, limiting the success of this approach. We recommend to pursue this approach but select meta-analysis data and use PFT specific SOC output from models, instead of grid cell averages with cropland fraction above a threshold as it was done here because ISIMIP models did not report PFT specific carbon variables.

4.5 Limitations

Despite much work to incorporate agricultural process in current models (Ciais et al., 2011; Ito and Inatomi, 2012; Lutz et al., 2019; Pugh et al., 2015; Tian et al., 2010; Wu et al., 2016), some limitations still remain for projection of SOC in our modeling approach. Carbon capture and storage in cropland are dependent on management practices. Practices such as tillage, crop rotation, crop residue management, etc. are very important to soil carbon decomposition but are not included in all models. Moreover, the inclusion of nitrogenous fertilizers was found to be a limiting factor in the amount of carbon stored (Drewniak et al., 2015). Likewise, nitrogen limitation is not considered in all models (Table S1). In addition, the application of genetically engineered crop with enhanced root exudates may affect soil functions and microbial diversity (Motavalli et al., 2004), which are important factors affecting soil carbon decomposition. Further research is needed to develop approaches able to represent such important management practices in models to fully evaluate the importance of agricultural practices for soil carbon.

5. Conclusion

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SOC changes have been significantly influenced by climate and land use change over the past century. Even though an increasing number of data sets are available to constrain future simulations, large differences between climate scenarios and TBMs remain, which implies large uncertainty of future projections. We show that uncertainties in future ΔSOC can be primarily attributed to the structural differences between TBMs rather than difference in GCMs. For RCP 2.6 land use change is the dominant driver of future Δ SOC, while for RCP 6.0 the climate change effect dominates. Soil input changes explain most of variations in projected ΔSOC across the TBMs globally and in different climate regions. Applying an emergent constraint for ΔSOC to climate change under RCP 6.0, our results showed a reduction in constrained Δ SOC compared to original modelled ensembles for all GCMs in the temperate region, and one GCM (i.e., HadGEM2-ES) globally and in the boreal region. In cropland dominated areas, SOC will continue to diminish under RCP 2.6 (-0.19 \pm 3.72 Pg C) and RCP 6.0 (-0.18 \pm 3.52 Pg C) due to cropland expansion, but with gains and losses compensating between regions. In cropland dominated areas, the large spread in constrained $\Delta soc_{cropland\ expansion}$ comes from uncertainties of observations. The idea of an emergent constraint approach purposefully reduced uncertainties in future projections of SOC. Although the uncertainties in constrain Δ SOC are still relatively high, as more accurate observation data and more model simulations become available, applying an emergent constraint approach to improve the accuracy of future Δ SOC projections is a promising research avenue. More importantly, understanding how SOC could be impacted by future climate change and land use changes can effectively help land managers and policymakers to develop appropriate land planning strategies.

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Data available statement

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Model data from ISIMIP2b are publicly available at https://esg.pik-729 730 potsdam.de/projects/isimip/. GPP products from P-Model are available from Stocker et al. (2019) [https://zenodo.org/record/1423484#.XKNO8pj7Q2x]. GPP products from BESS are 731 publicly available from Jiang and Ryu, (2016) [http://environment.snu.ac.kr/bess_flux/]. The 732 soil organic carbon density data from meta-analysis are available from Deng et al. (2016), Li 733 734 et al. (2018) and Nyawira et al. (2016). HWSD data are available from FAO/IIASA/ISRIC/ISSCAS/JRC (2012) at http://www.fao.org/soils-portal/soil-survey/soil-735 maps-and-databases/harmonized-world-soil-database-v12/zh/, the NCSCD are avaiable from 736 737 Tarnocai et al. (2009) at data link:https://bolin.su.se/data/ncscd/, and the WISE30sec are 738 available from Batjes, (2016) at https://www.isric.org/explore/wise-databases.

References

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- 740 Álvaro-Fuentes, J., F. J. Morell, E. Madejón, J. Lampurlanés, J. L. Arrúe, and C. Cantero-
- Martínez (2013), Soil biochemical properties in a semiarid Mediterranean agroecosystem as
- affected by long-term tillage and N fertilization, *Soil and Tillage Research*, 129, 69-74.
- 743 doi:10.1016/j.still.2013.01.005.
- Angers, D. A., and N. S. Eriksen-Hamel (2008), Full-Inversion Tillage and Organic Carbon
- 745 Distribution in Soil Profiles: A Meta-Analysis, Soil Science Society of America Journal,
- 746 72(5), 1370-1374. doi:10.2136/sssaj2007.0342.
- 747 Batjes, N. H. (2016), Harmonized soil property values for broad-scale modelling

- 748 (WISE30sec) with estimates of global soil carbon stocks, *Geoderma*, 269, 61-68.
- 749 doi:10.1111/j.1365-2389.1996.tb01386.x.
- 750 Bellamy, P. H., P. J. Loveland, R. I. Bradley, R. M. Lark, and G. J. Kirk (2005), Carbon losses
- 751 from all soils across England and Wales 1978-2003, *Nature*, 437(7056), 245-248.
- 752 doi:10.2136/sssaj2009.0036.
- 753 Bondeau, A., et al. (2007), Modelling the role of agriculture for the 20th century global
- terrestrial carbon balance, Global Change Biology, 13(3), 679-706. doi:10.1111/j.1365-
- 755 2486.2006.01305.x.
- 756 Bronick, C. J., and Lal, R. (2005). Manuring and rotation effects on soil organic carbon
- 757 concentration for different aggregate size fractions on two soils in northeastern Ohio,
- 758 USA. Soil and Tillage Research, 81(2), 239-252. doi:10.1016/j.still.2004.09.011.
- 759 Carvalhais, N., et al. (2014), Global covariation of carbon turnover times with climate in
- 760 terrestrial ecosystems, *Nature*, *514*(7521), 213-217. doi:10.1038/nature13731.
- 761 Ciais, P., S. Gervois, N. Vuichard, S. L. Piao, and N. Viovy (2011), Effects of land use change
- and management on the European cropland carbon balance, Global Change Biology, 17(1),
- 763 320-338.
- Ciais, P., C. Sabine, G. Bala, L. Bopp, V. Brovkin, J. Canadell, A. Chhabra, R. DeFries, J.
- Galloway, and M. Heimann (2014), Carbon and other biogeochemical cycles, in *Climate*
- 766 change 2013: the physical science basis. Contribution of Working Group I to the Fifth
- 767 Assessment Report of the Intergovernmental Panel on Climate Change, edited, pp. 465-570,
- 768 Cambridge University Press.
- 769 Cox, P. M., D. Pearson, B. B. Booth, P. Friedlingstein, C. Huntingford, C. D. Jones, and C.
- 770 M. Luke (2013), Sensitivity of tropical carbon to climate change constrained by carbon
- dioxide variability, *Nature*, 494(7437), 341-344. doi:10.1038/nature11882.
- 772 DeAngelis, A. M., X. Qu, M. D. Zelinka, and A. Hall (2015), An observational radiative
- constraint on hydrologic cycle intensification, *Nature*, 528(7581), 249-253.
- 774 doi:10.1038/nature15770.

- De Kauwe, M. G., T. F. Keenan, B. E. Medlyn, I. C. Prentice, and C. Terrer (2016), Satellite
- based estimates underestimate the effect of CO₂ fertilization on net primary productivity,
- 777 *Nature Climate Change*, 6(10), 892-893. doi:10.1038/nclimate3105.
- 778 Deng, L., G.-y. Zhu, Z.-s. Tang, and Z.-p. Shangguan (2016), Global patterns of the effects of
- land-use changes on soil carbon stocks, *Global Ecology and Conservation*, 5, 127-138.
- 780 doi:10.1016/j.gecco.2015.12.004.
- Doetterl, S., et al. (2015), Soil carbon storage controlled by interactions between
- 782 geochemistry and climate, *Nature Geoscience*, 8(10), 780-783. doi:10.1038/ngeo2516.
- 783 Don, A., J. Schumacher, and A. Freibauer (2011), Impact of tropical land-use change on soil
- organic carbon stocks a meta-analysis, Global Change Biology, 17(4), 1658-1670.
- 785 doi:10.1111/j.1365-2486.2010.02336.x.
- 786 Drewniak, B. A., U. Mishra, J. Song, J. Prell, and V. R. Kotamarthi (2015), Modeling the
- 787 impact of agricultural land use and management on US carbon budgets, *Biogeosciences*,
- 788 12(7), 2119-2129. doi:10.5194/bg-12-2119-2015.
- 789 FAO/IIASA/ISRIC/ISSCAS/JRC: Harmonized World Soil Database (version 1.10), FAO,
- 790 Rome, Italy and IIASA, Laxenburg, Austria, 2012.
- 791 Foley, J. A., et al. (2005), Global consequences of land use., *Science*, 309(5734), 570–574,
- 792 doi:10.1126/science.1111772.
- Foley, J. A., et al. (2011), Solutions for a cultivated planet, *Nature*, 478(7369), 337–342,
- 794 doi:10.1038/nature10452.
- 795 Frieler, K., et al. (2017), Assessing the impacts of 1.5 °C global warming simulation
- 796 protocol of the Inter-Sectoral Impact Model Intercomparison Project (ISIMIP2b),
- 797 Geoscientific Model Development, 10(12), 4321-4345. doi:10.5194/gmd-10-4321-2017.
- 798 Gervois, S., Ciais, P., de Noblet-Ducoudré, N., Brisson, N., Vuichard, N., & Viovy, N. (2008).
- 799 Carbon and water balance of European croplands throughout the 20th century. *Global*
- 800 *Biogeochemical Cycles*, 22(2). doi:10.1029/2007gb003018.
- Gregorich, E., P. Rochette, A. VandenBygaart, D. J. S. Angers, and T. Research (2005),

- 802 Greenhouse gas contributions of agricultural soils and potential mitigation practices in
- 803 Eastern Canada. *Soil and Tillage Research*, 83(1), 53-72. doi:10.1016/j.still.2005.02.009.
- Guimberteau, M., et al. (2018), ORCHIDEE-MICT (v8.4.1), a land surface model for the
- 805 high latitudes: model description and validation, Geoscientific Model Development, 11(1),
- 806 121-163. doi:10.5194/gmd-11-121-2018.
- 807 Guo, L. B., and R. M. Gifford (2002), Soil carbon stocks and land use change: a meta
- analysis, Global Change Biology, 8(4), 345-360. doi:10.1046/j.1354-1013.2002.00486.x.
- Haberl, H., K. H. Erb, F. Krausmann, V. Gaube, A. Bondeau, C. Plutzar, S. Gingrich, W.
- Lucht, and M. Fischer-Kowalski (2007), Quantifying and mapping the human appropriation
- of net primary production in earth's terrestrial ecosystems, *Proceedings of the National*
- 812 *Academy of Sciences*, 104(31), 12942-12947. doi:10.1073/pnas.0704243104.
- Hall, A., P. Cox, C. Huntingford, and S. Klein (2019), Progressing emergent constraints on
- 814 future climate change, *Nature Climate Change*, 9(4), 269-278. doi:10.1038/s41558-019-
- 815 0436-6.
- Hamdi, S., F. Moyano, S. Sall, M. Bernoux, and T. Chevallier (2013), Synthesis analysis of
- 817 the temperature sensitivity of soil respiration from laboratory studies in relation to incubation
- methods and soil conditions, *Soil Biology and Biochemistry*, 58, 115-126.
- 819 doi:10.1016/j.soilbio.2012.11.012.
- Hansis, E., S. J. Davis, and J. Pongratz (2015), Relevance of methodological choices for
- accounting of land use change carbon fluxes, Global Biogeochemical Cycles, 29(8), 1230-
- 822 1246. doi:10.1002/2014gb004997.
- He, Y., S. Piao, X. Li, A. Chen, and D. Qin (2018), Global patterns of vegetation carbon use
- 824 efficiency and their climate drivers deduced from MODIS satellite data and process-based
- models, *Agricultural and Forest Meteorology*, 256-257, 150-158.
- 826 doi:10.1016/j.agrformet.2018.03.009.
- Hempel, S., Frieler, K., Warszawski, L., Schewe, J., and Piontek, F. (2013). A trend-
- preserving bias correction—the ISI-MIP approach, Earth Syst. Dynam., 4, 219-236.

- 829 doi:10.5194/esd-4-219-2013.
- Hengl, T., et al. (2017), SoilGrids250m: Global gridded soil information based on machine
- learning, *PLoS One*, 12(2), e0169748. doi:10.1371/journal.pone.0169748.
- Hickler, T., A. Rammig, and C. Werner (2015), Modelling CO₂ Impacts on Forest
- 833 Productivity, *Current Forestry Reports*, *1*(2), 69-80. doi:10.1007/s40725-015-0014-8.
- Houghton, R. A., and A. A. Nassikas (2017), Global and regional fluxes of carbon from land
- use and land cover change 1850–2015, Global Biogeochemical Cycles, 31(3), 456-472.
- 836 doi:10.1002/2016GB005546.
- 837 Ito, A., and M. Inatomi (2012), Water-Use Efficiency of the Terrestrial Biosphere: A Model
- Analysis Focusing on Interactions between the Global Carbon and Water Cycles, *Journal of*
- 839 *Hydrometeorology*, *13*(2), 681-694. doi:10.1175/jhm-d-10-05034.1.
- Jiang, C., and Y. Ryu (2016), Multi-scale evaluation of global gross primary productivity and
- evapotranspiration products derived from Breathing Earth System Simulator (BESS), *Remote*
- *Sensing of Environment*, 186, 528-547. doi:10.1016/j.rse.2016.08.030.
- Jones, C., C. McConnell, K. Coleman, P. Cox, P. Falloon, D. Jenkinson, and D. Powlson
- 844 (2005), Global climate change and soil carbon stocks; predictions from two contrasting
- models for the turnover of organic carbon in soil, *Global Change Biology*, 11(1), 154-166.
- 846 doi:10.1111/j.1365-2486.2004.00885.x.
- Klein Goldewijk, K., A. Beusen, J. Doelman, and E. J. E. S. S. D. Stehfest (2017), New
- anthropogenic land use estimates for the Holocene: HYDE 3.2, 9(2), 927-953.
- Köchy, M., A. Don, M. K. van der Molen, and A. Freibauer (2015), Global distribution of soil
- organic carbon Part 2: Certainty of changes related to land use and climate, Soil, 1(1), 367-
- 851 380. doi:10.5194/soil-1-367-2015.
- 852 Kolby Smith, W., C. C. Cleveland, S. C. Reed, and S. W. Running (2014), Agricultural
- 853 conversion without external water and nutrient inputs reduces terrestrial vegetation
- productivity, Geophysical Research Letters, 39, 363-391. doi:10.1002/2013GL058857.
- Kolby Smith, W., S. C. Reed, C. C. Cleveland, A. P. Ballantyne, W. R. L. Anderegg, W. R.

- Wieder, Y. Y. Liu, and S. W. Running (2015), Large divergence of satellite and Earth system
- model estimates of global terrestrial CO2 fertilization, *Nature Climate Change*, 6(3), 306-
- 858 310. doi:10.1038/nclimate2879.
- 859 Koven, C. D., J. Q. Chambers, K. Georgiou, R. Knox, R. Negron-Juarez, W. J. Riley, V. K.
- Arora, V. Brovkin, P. Friedlingstein, and C. D. Jones (2015), Controls on terrestrial carbon
- feedbacks by productivity versus turnover in the CMIP5 Earth System Models,
- 862 *Biogeosciences*, 12(17), 5211-5228. doi:10.5194/bg-12-5211-2015.
- Krausmann, F., K.-H. Erb, S. Gingrich, C. Lauk, and H. Haberl (2008), Global patterns of
- socioeconomic biomass flows in the year 2000: A comprehensive assessment of supply,
- consumption and constraints, *Ecological Economics*, 65(3), 471-487.
- 866 doi:10.1016/j.ecolecon.2007.07.012.
- Lange, S. (2016), EartH2Observe, WFDEI and ERA-Interim data Merged and Bias-corrected
- for ISIMIP (EWEMBI), GFZ Data Services, https://doi.org/10.5880/pik.2019.004.
- 869 Li, W., et al. (2018), Temporal response of soil organic carbon after grassland-related land-
- 870 use change, *Glob Chang Biol*, 24(10), 4731-4746. doi:10.1111/gcb.14328.
- Li, W., et al. (2017), Land-use and land-cover change carbon emissions between 1901 and
- 872 2012 constrained by biomass observations, *Biogeosciences*, 14(22), 5053-5067.
- 873 doi:10.5194/bg-14-5053-2017.
- Lian, X., et al. (2018), Partitioning global land evapotranspiration using CMIP5 models
- constrained by observations, *Nature Climate Change*, 8(7), 640-646. doi:10.1038/s41558-
- 876 018-0207-9.
- Liu, S., Y. Wei, W. M. Post, R. B. Cook, K. Schaefer, and M. M. Thornton (2013), The
- 878 Unified North American Soil Map and its implication on the soil organic carbon stock in
- 879 North America, *Biogeosciences*, 10(5), 2915-2930. doi:10.5194/bg-10-2915-2013.
- 880 Lozano-García, B., Muñoz-Rojas, M., & Parras-Alcántara, L. (2017). Climate and land use
- changes effects on soil organic carbon stocks in a Mediterranean semi-natural area. Science of
- the Total Environment, 579, 1249-1259. doi:10.1016/j.scitotenv.2016.11.111.

- 883 Luo, Y., et al. (2016), Toward more realistic projections of soil carbon dynamics by Earth
- 884 system models, *Global Biogeochemical Cycles*, *30*(1), 40-56. doi:10.1002/2015gb005239.
- Lutz, F., T. Herzfeld, J. Heinke, S. Rolinski, S. Schaphoff, W. von Bloh, J. J. Stoorvogel, and
- 886 C. Müller (2019), Simulating the effect of tillage practices with the global ecosystem model
- 887 LPJmL (version 5.0-tillage), Geoscientific Model Development, 12(6), 2419-2440.
- 888 doi:10.5194/gmd-12-2419-2019.
- Meinshausen, M., et al. (2011), The RCP greenhouse gas concentrations and their extensions
- 890 from 1765 to 2300, *Climatic Change*, 109(1-2), 213-241. doi:10.1007/s10584-011-0156-z.
- Molotoks, A., E. Stehfest, J. Doelman, F. Albanito, N. Fitton, T. P. Dawson, and P. Smith
- 892 (2018), Global projections of future cropland expansion to 2050 and direct impacts on
- biodiversity and carbon storage, Globle Change Biology, 24(12), 5895-5908.
- 894 doi:10.1111/gcb.14459.
- Monfreda, C., N. Ramankutty, and J. A. Foley (2008), Farming the planet: 2. Geographic
- distribution of crop areas, yields, physiological types, and net primary production in the year
- 897 2000, Global Biogeochemical Cycles, 22(1), GB1022. doi:10.1029/2007gb002947.
- Motavalli, P. P., Kremer, R. J., Fang, M., and Means, N. E. (2004). Impact of genetically
- 899 modified crops and their management on soil microbially mediated plant nutrient
- 900 transformations. *Journal of environmental quality*, 33(3), 816-824. doi:
- 901 10.2134/jeq2004.0816
- 902 Müller, C., Robertson, R.D. (2014) Projecting future crop productivity for global economic
- 903 modeling. *Agricultural Economics*, 45(1), 37-50. doi:10.1111/agec.12088.
- Neumann, M., and P. Smith (2018), Carbon uptake by European agricultural land is variable,
- and in many regions could be increased: Evidence from remote sensing, yield statistics and
- models of potential productivity, *The Science of the total environment*, 643, 902-911.
- 907 doi:10.1016/j.scitotenv.2018.06.268.
- 908 Nishina, K., et al. (2014), Quantifying uncertainties in soil carbon responses to changes in
- global mean temperature and precipitation, *Earth System Dynamics*, 5(1), 197-209.

- 910 doi:10.5194/esd-5-197-2014.
- 911 Nyawira, S. S., J. E. M. S. Nabel, A. Don, V. Brovkin, and J. Pongratz (2016), Soil carbon
- 912 response to land-use change: evaluation of a global vegetation model using observational
- 913 meta-analyses, *Biogeosciences*, 13(19), 5661-5675. doi:10.5194/bg-13-5661-2016.
- 914 Pinheiro, É. F. M., D. V. B. de Campos, F. de Carvalho Balieiro, L. H. C. dos Anjos, and M.
- 915 G. Pereira (2015), Tillage systems effects on soil carbon stock and physical fractions of soil
- 916 organic matter, *Agricultural Systems*, *132*, 35-39. doi:10.1016/j.agsy.2014.08.008.
- Poeplau, C., A. Don, L. Vesterdal, J. Leifeld, B. A. S. Van Wesemael, J. Schumacher, and A.
- 918 Gensior (2011), Temporal dynamics of soil organic carbon after land-use change in the
- 919 temperate zone carbon response functions as a model approach, Global Change Biology,
- 920 *17*(7), 2415-2427. doi:10.1111/j.1365-2486.2011.02408.x.
- Popp, A., et al. (2014), Land-use protection for climate change mitigation, *Nature Climate*
- 922 *Change*, 4, 1095. doi:10.1038/nclimate2444.
- Popp, A., et al. (2017), Land-use futures in the shared socio-economic pathways, Global
- 924 Environmental Change, 42, 331-345. doi:10.1016/j.gloenvcha.2016.10.002.
- Post, W. M., & Kwon, K. C. (2000). Soil carbon sequestration and land-use change: processes
- 926 and potential. *Global change biology*, 6(3), 317-327. doi:10.1046/j.1365-2486.2000.00308.x.
- 927 Powlson, D. S., A. Bhogal, B. J. Chambers, K. Coleman, A. J. Macdonald, K. W. T.
- 928 Goulding, and A. P. Whitmore (2012), The potential to increase soil carbon stocks through
- 929 reduced tillage or organic material additions in England and Wales: A case study, *Agriculture*,
- 930 Ecosystems & Environment, 146(1), 23-33. doi:10.1016/j.agee.2011.10.004.
- 931 Pugh, T. A. M., A. Arneth, S. Olin, A. Ahlström, A. D. Bayer, K. Klein Goldewijk, M.
- Lindeskog, and G. Schurgers (2015), Simulated carbon emissions from land-use change are
- 933 substantially enhanced by accounting for agricultural management, *Environmental Research*
- 934 *Letters*, 10(12), 124008. doi:10.1088/1748-9326/10/12/124008.
- Ryu, Y., et al. (2011), Integration of MODIS land and atmosphere products with a coupled-
- process model to estimate gross primary productivity and evapotranspiration from 1 km to

- 937 global scales, *Global Biogeochemical Cycles*, 25(4), GB4017. doi:10.1029/2011GB004053.
- 938 Sanderman, J., T. Hengl, and G. J. Fiske (2017), Soil carbon debt of 12,000 years of human
- 939 land use, Proceedings of the National Academy of Sciences of the United States of America,
- 940 114(36), 9575-9580. doi:10.1073/pnas.1706103114.
- 941 Smith, B., D. Wårlind, A. Arneth, T. Hickler, P. Leadley, J. Siltberg, and S. Zaehle (2014),
- 942 Implications of incorporating N cycling and N limitations on primary production in an
- 943 individual-based dynamic vegetation model, *Biogeosciences*, 11(7), 2027-2054.
- 944 doi:10.1111/gcb.14677.
- 945 Snyder, C. S., T. W. Bruulsema, T. L. Jensen, and P. E. Fixen (2009), Review of greenhouse
- gas emissions from crop production systems and fertilizer management effects, *Agriculture*,
- 947 Ecosystems & Environment, 133(3), 247-266. doi:10.1029/2018MS001277.
- 948 Stevanović, M., A. Popp, H. Lotze-Campen, J. P. Dietrich, C. Müller, M. Bonsch, C. Schmitz,
- 949 B. L. Bodirsky, F. Humpenöder, and I. Weindl (2016), The impact of high-end climate change
- on agricultural welfare, *Science Advances*, 2(8), e1501452. doi:10.1126/sciadv.1501452.
- 951 Stocker, B. D., J. Zscheischler, T. F. Keenan, I. C. Prentice, S. I. Seneviratne, and J. Peñuelas
- 952 (2019), Drought impacts on terrestrial primary production underestimated by satellite
- 953 monitoring, *Nature Geoscience*, 12(4), 264-270. doi:10.1038/s41561-019-0318-6.
- 954 Tarnocai, C., J. G. Canadell, E. A. G. Schuur, P. Kuhry, G. Mazhitova, and S. Zimov (2009),
- 955 Soil organic carbon pools in the northern circumpolar permafrost region, *Global*
- 956 *Biogeochemical Cycles*, 23(2), GB2023. doi:10.1029/2008gb003327.
- 957 Tian, H., G. Chen, M. Liu, C. Zhang, G. Sun, C. Lu, X. Xu, W. Ren, S. Pan, and A.
- 958 Chappelka (2010), Model estimates of net primary productivity, evapotranspiration, and
- 959 water use efficiency in the terrestrial ecosystems of the southern United States during 1895–
- 960 2007, Forest Ecology and Management, 259(7), 1311-1327.
- Tian, H., G. Chen, C. Lu, X. Xu, D. J. Hayes, W. Ren, S. Pan, D. N. Huntzinger, and S. C.
- 962 Wofsy (2015a), North American terrestrial CO₂ uptake largely offset by CH4 and N₂O
- emissions: toward a full accounting of the greenhouse gas budget, *Climatic Change*, 129(3),

- 964 413-426. doi:10.1007/s10584-014-1072-9.
- 965 Tian, H., et al. (2015b), Global patterns and controls of soil organic carbon dynamics as
- simulated by multiple terrestrial biosphere models: Current status and future directions,
- 967 *Global Biogeochem Cycles*, 29(6), 775-792. doi:10.1002/2014GB005021.
- 968 Todd-Brown, K. E. O., J. T. Randerson, W. M. Post, F. M. Hoffman, C. Tarnocai, E. A. G.
- 969 Schuur, and S. D. Allison (2013), Causes of variation in soil carbon simulations from CMIP5
- 970 Earth system models and comparison with observations, *Biogeosciences*, 10(3), 1717-1736.
- 971 doi:10.5194/bg-10-1717-2013.
- 972 Todd-Brown, K. E. O., et al. (2014), Changes in soil organic carbon storage predicted by
- 973 Earth system models during the 21st century, *Biogeosciences*, 11(8), 2341-2356.
- 974 doi:10.5194/bg-11-2341-2014.
- 975 VandenBygaart, A., E. Gregorich, and D. J. C. J. o. S. S. Angers (2003), Influence of
- agricultural management on soil organic carbon: A compendium and assessment of Canadian
- 977 studies, 83(4), 363-380. doi:10.4141/S03-009.
- 978 Virto, I., P. Barré, A. Burlot, and C. Chenu (2012), Carbon input differences as the main
- 979 factor explaining the variability in soil organic C storage in no-tilled compared to inversion
- 980 tilled agrosystems, *Biogeochemistry*, 108(1-3), 17-26. doi:10.1007/s10533-011-9600-4.
- Vogeler, I., J. Rogasik, U. Funder, K. Panten, and E. Schnug (2009), Effect of tillage systems
- and P-fertilization on soil physical and chemical properties, crop yield and nutrient uptake,
- 983 *Soil and Tillage Research*, *103*(1), 137-143. doi:10.1016/j.still.2008.10.004.
- Wang, H., I. C. Prentice, T. F. Keenan, T. W. Davis, I. J. Wright, W. K. Cornwell, B. J. Evans,
- and C. Peng (2017), Towards a universal model for carbon dioxide uptake by plants, *Nat*
- 986 *Plants*, 3(9), 734-741. doi:10.1038/s41477-017-0006-8. doi: 10.1111/j.1466-
- 987 8238.2011.00678.x.
- 988 Wei, X., M. Shao, W. J. Gale, X. Zhang, and L. Li (2013), Dynamics of aggregate-associated
- organic carbon following conversion of forest to cropland, Soil Biology and Biochemistry, 57,
- 990 876-883. doi:10.1016/j.soilbio.2012.10.020.

- 991 Wei, X., M. Shao, W. Gale, and L. Li (2014), Global pattern of soil carbon losses due to the
- onversion of forests to agricultural land, *Scientific reports*, 4, 4062. doi:10.1038/srep04062.
- 993 Wenzel, S., P. M. Cox, V. Eyring, and P. Friedlingstein (2016), Projected land photosynthesis
- onstrained by changes in the seasonal cycle of atmospheric CO₂, *Nature*, 538(7626), 499-
- 995 501. doi:10.1038/nature19772.
- 996 Wu, D., Piao, S., Liu, Y., Ciais, P., and Yao, Y. (2018). Evaluation of CMIP5 Earth System
- 997 Models for the spatial patterns of biomass and soil carbon turnover times and their linkage
- 998 with climate. Journal of Climate, 31(15), 5947-5960. doi: 10.1175/JCLI-D-17-0380.1.
- 999 Wu, X., et al. (2016), ORCHIDEE-CROP (v0), a new process-based agro-land surface model:
- model description and evaluation over Europe, Geoscientific Model Development, 9(2), 857-
- 1001 873.
- 1002 Xiao, J., K. J. Davis, N. M. Urban, and K. Keller (2014), Uncertainty in model parameters
- and regional carbon fluxes: A model-data fusion approach, Agricultural and Forest
- 1004 *Meteorology*, 189-190, 175-186. doi:10.1016/j.agrformet.2014.01.022.
- Yan, Y., Y. Luo, X. Zhou, and J. Chen (2014), Sources of variation in simulated ecosystem
- carbon storage capacity from the 5th Climate Model Intercomparison Project (CMIP5), *Tellus*
- 1007 *B: Chemical and Physical Meteorology*, *66*(1), 22568. doi:10.3402/tellusb.v66.22568.
- Thao, C., et al. (2016), Plausible rice yield losses under future climate warming, *Nat Plants*,
- 1009 *3*, 16202. doi:10.1038/nplants.2016.202.
- 1010 Zhao, M., and S. W. Running (2010), Drought-Induced Reduction in Global Terrestrial Net
- 1011 Primary Production from 2000 Through 2009, *Science*, *329*(5994), 940.
- 1012 doi:10.1126/science.1192666.

1013 **Table 1** Description of scenario design used from ISIMIP2b.

1015

ISIMIP2b simulations		Driver	Description
Group 1	Historical (1861-2005)	Climate + Land use	The effects of historical climate change with varying land use change
Group 2	only CC (2006-2099)	Climate + Fixed land use	Pure effect of future climate change assuming fixed year 2005 levels of land
			use change under RCP 2.6 and 6.0 scenario
Group 3	CC+LUC (2006-2099)	Climate + Land use	The effects of future climate change and land use change from 2005 onwards
			associated with RCP 2.6 and 6.0 scenario
Group3-Group2	only LUC (2006-2099)	Land use	Pure effect of future land use change under RCP 2.6 and 6.0 scenario

Note: you can find different climate and land use change -impacts simulation data (the simulation round, sectors, scenarios, variables, time period etc.) at https://esg.pik-potsdam.de/projects/isimip/.

Table 2 Modeled global soil carbon changes (Δ SOC) during historical period (1861-2005), and during future period (2006-2099) under both the effects of climate change and land use change (CC+LUC), the effect of only climate change (CC) and the effect of only land use change (LUC) based on RCP 2.6 and 6.0. The bold and bold-italic indicates the largest and smallest value, respectively. The Δ SOC is the difference in SOC compared to the means of 1861-1870.

ΔSOC (Pg C)		TT: -4	Future (2099-2005)							
ΔSOC (Pg C)	History		RCP 2.6			RCP 6.0			
TBMs	GCM	2005-1861	CC+LUC	CC	LUC	CC+LUC	CC	LUC		
	GFDL-ESM2M	16.5	14.9	15.1	-0.2	18.3	18.4	-0.1		
LPJ-GUESS	HadGEM2-ES	26.0	-0.5	-0.6	0.1	-11.3	-11.4	0.1		
LPJ-GUESS	IPSL-CM5A-LR	9.3	1.9	1.8	0.1	-1.9	-1.8	-0.1		
	MIROC5	23.4	5.6	5.8	-0.2	13.2	13.4	-0.2		
	GFDL-ESM2M	-52.1	31.9	11.7	20.2	10.0	12.1	-2.1		
LPJmL	HadGEM2-ES	-38.6	-9.4	-28.2	18.8	-22.9	-19.4	-3.5		
LPJIIIL	IPSL-CM5A-LR	-81.3	-8.0	-25.5	17.4	-30.1	-25.3	-4.8		
	MIROC5	-45.0	28.0	5.9	22.1	31.4	36.3	-4.9		
	GFDL-ESM2M	71.4	114.7	38.5	76.3	154.9	148.9	6.0		
VISIT	HadGEM2-ES	88.8	109.9	22.8	87.1	118.4	109.4	8.9		
V1511	IPSL-CM5A-LR	46.1	112.7	28.4	84.3	141.0	132.5	8.5		
	MIROC5	72.5	103.9	25.5	78.4	176.5	171.5	5.0		
	GFDL-ESM2M	22.6	71.7	55.1	16.6	80.9	80.6	0.3		
ORCHIDEE-MICT	HadGEM2-ES	18.0		11.3			21.4			
ORCHIDEE-MICT	IPSL-CM5A-LR	-17.4	33.3	17.9	15.4	35.0	36.9	-1.9		
	MIROC5	34.2		47.8			89.9			
DLEM	GFDL-ESM2M	7.2	34.5	20.1	14.4	38.6	26.8	11.8		
DLEM	IPSL-CM5A-LR	4.3	30.3	15.2	15.0	30.8	19.7	11.0		

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	MIROC5	9.7	36.0	21.8	14.2	40.9	29.2	11.7
Model range		-81.3~88.8	-9.4~114.7	-28.2~55.1	-0.2~87.1	-30.1~176.5	-25.3~171.5	-4.9-11.8
Model mean		11.4	41.8	15.3	28.2	48.5	46.8	2.7
Model Median		16.51	31.91	15.22	16.61	31.40	26.83	0.14

Table 3 Modeled ΔSOC (unit, Pg C), constrained ΔSOC by NPP trends over the period of 2001-2015 and constrained ΔSOC by initial SOC (1995-2005) under RCP 6.0 scenario in natural vegetation grid-cells (exclude the cropland) and its different climate region across different GCMs. The number indicates the number of TBMs with modeled Δ SOC within 1-sigma uncertainty of the constrained Δ SOC. Bold values indicate the constrained Δ SOC based on a significant relationship between Δ SOC and NPP or initial SOC, respectively. A p value \leq 0.1 was considered significant.

Classificat	tion	Modele	ed ΔSOC	Constrai	ned ΔSO	OC by NPP	Constrained ΔSOC by initial SOC		
		Mean	1σ	Mean	1σ	Within 1σ uncertainty	Mean	1σ	Within 1σ uncertainty
Global	Natural vegetation								
	GFDL-ESM2M	47.9	49.4	25.2	42.2	4.0	49.0	0.6	0
	HadGEM2-ES	19.0	53.9	-36.9	67.3	3.0	45.6	6.8	0
	IPSL-CM5A-LR	26.1	54.6	1.4	42.6	4.0	33.1	3.4	0
	MIROC5	54.9	53.1	38.8	54.1	4.0	54.8	0.1	0
Climate	Tropical region								
Region	GFDL-ESM2M	14.4	10.4	4.9	9.0	3.0	16.1	0.6	0
	HadGEM2-ES	4.7	21.1	14.4	21.6	3.0	-4.8	2.3	0
	IPSL-CM5A-LR	9.3	14.1	6.2	13.3	4.0	10.3	0.8	0
	MIROC5	20.3	15.2	9.2	13.0	3.0	20.5	0.7	0
	Temperate region								
	GFDL-ESM2M	31.3	46.2	65.5	44.2	2.0	36.3	2.0	0
	HadGEM2-ES	12.7	56.8	-96.5	61.7	1.0	48.1	8.0	0
	IPSL-CM5A-LR	14.8	57.0	-11.6	49.5	4.0	27.6	4.6	0
	MIROC5	32.9	48.8	37.2	47.0	3.0	37.0	1.8	0
	Boreal region								
	GFDL-ESM2M	2.2	7.1	2.5	7.1	3.0	3.1	1.1	0
	HadGEM2-ES	1.6	19.3	-20.6	5.2	1.0	9.2	4.7	0

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IPSL-CM5A-LR	2.0	11.8	0.9	11.4	3.0	3.7	2.1	0
MIROC5	1.6	9.7	2.1	7.4	3.0	2.6	1.5	2

Table 4 The global and regional soil carbon changes across cropland expansion area ($\Delta soc_{cropland\ expansion}$, calculated as ΔSOC divided by the increased area of cropland; unit, kg C m⁻²) and the soil carbon density changes ($\Delta soc_{cropland\ minus\ initial\ natureg}$; unit, kg C m⁻²) between initial natural vegetation (in 1861) and current cropland (in 2005) from original ISIMIP2b models and from the estimate constrained by meta-analysis data ($\Delta soc_{cropland\ minus\ initial\ natureg}$; unit, kg C m⁻²) in land use dominated areas. The number indicates the total model number with modeled $\Delta soc_{cropland\ expansion}$ within 1-sigma uncertainty of the constrained $\Delta soc_{cropland\ expansion}$.

Region	LUC type	$\Delta soc_{cropland\ m}$	inus initial natveg	$\Delta soc_{cropland\ expansion}$					
	LOC type	Meta-analysis	ISMIP2b	Constrained	ISMIP2b	number			
Eurasia	G-C (temp)	-1.51 ± 5.61	-0.39 ± 1.32	-2.46 ± 12.22	-0.15 ± 2.95	19			
North America	G-C (temp)	-1.51 ± 5.61	-0.76 ± 1.57	-2.78 ± 9.92	-1.43 ± 2.85	19			
South America	G-C (temp)	-1.51 ± 5.61	-0.50 ± 1.62	-1.89 ± 10.08	-0.22 ± 3.09	19			
Africa	G-C (trop)	-1.62 ± 1.76	-0.42 ± 1.16	-1.66 ± 2.27	-0.18 ± 1.67	14			
West Eurasia	G-C (temp)	-1.51 ± 5.61	-0.27 ± 0.99	-5.94 ± 17.68	-1.82 ± 3.35	19			
Australia	G-C (temp)	-1.51 ± 5.61	0.20 ± 1.09	-3.45 ± 12.84	0.45 ± 2.53	19			
South Asia	F-C (trop)	-2.14 ± 2.72	-2.25 ± 1.45	0.67 ± 2.45	0.12 ± 1.28	18			
East Asia	G-C (temp)	-1.51 ± 5.61	-4.52 ± 3.85	0.27 ± 0.03	0.20 ± 1.81	0			
LUC region		-1.74 ± 4.61	-1.74 ± 1.66	-0.29 ± 5.56	-0.61 ± 2.15	19			

Table 5 The estimated regional soil carbon change (ΔSOC , Pg C) due to land use change in historical (1861-2005) and future period (2005-2099). The estimated ΔSOC was equal to the constrained ΔSOC was equal to the constrained ΔSOC region.

	Modeled	Estimated ΔSOC							
Region	Historical		Historical		RCP 2.6		RCP 6.0		
	ΔSOC	Area	ΔSOC	Area	ΔSOC	Area	ΔSOC		
Eurasia	-0.08 ± 1.32	0.406	-1.00 ± 4.96	0.020	-0.05 ± 0.24	0.011	-0.03 ± 0.14		
North America	-1.74 ± 3.08	1.004	-2.79 ± 9.96	0.039	-0.11 ± 0.38	0.000	0		
South America	-0.05 ± 0.65	0.190	-0.36 ± 1.92	0.179	-0.34 ± 1.80	0.152	-0.29 ± 1.53		
Africa	-0.09 ± 0.75	0.408	-0.68 ± 0.93	0.251	-0.42 ± 0.57	0.120	-0.20 ± 0.27		
West Eurasia	-0.10 ± 1.62	0.452	-2.69 ± 8.00	0.009	-0.05 ± 0.15	0.001	-0.01 ± 0.03		
Australia	0.18 ± 0.93	0.334	-1.15 ± 4.28	0.001	-0.003 ± 0.002	0.001	-0.002 ± 0.007		
South Asia	0.09 ± 0.88	0.623	0.42 ± 1.53	0.163	0.11 ± 0.40	0.348	0.23 ± 0.85		
East Asia	0.04 ± 0.34	0.171	0.05 ± 0.01	0.009	0.002 ± 0.001	< 0.0001	< 0.0001		
LUC region	-2.64 ± 8.43	3.588	-1.03 ± 19.94	0.670	-0.19 ± 3.72	0.633	-0.18 ± 3.52		

Note, there is no crop expansion under RCP 6.0 in North America.

Table 6 Global and natural ecosystem initial soil carbon stocks from different
 database (without permafrost C) and GCMs during the period of 1995-2005 at soil
 depth of 0-1 m. The bold and bold-italic indicates the largest and smallest value,
 respectively.

Detabase/TDMs	Data/GCMs	Global	N	atural ecosyste	em (Pg C))
Database/TBMs	Data/GCMS	(Pg C)	Tropical	Temperate	Boreal	Total
Database	HWSD	1265.8	376.4	645.9	72.6	1094.9
	WISE30sec	1419.8	362.4	758.0	108.1	1228.5
	HWSD+NCSCD	1454.4	376.5	748.1	159.3	1283.9
	Mean	1380.0	371.8	717.3	113.3	1202.4
LPJ-GUESS	GFDL-ESM2M	1340.0	356.4	748.7	61.0	1166.1
	HadGEM2-ES	1380.5	381.6	762.8	65.3	1209.7
	IPSL-CM5A-LR	1357.4	367.0	746.2	52.3	1165.5
	MIROC5	1380.9	401.1	755.1	46.4	1202.6
LPJmL	GFDL-ESM2M	2024.9	414.5	1210.3	271.6	1896.3
	HadGEM2-ES	2055.3	450.6	1195.5	285.3	1931.4
	IPSL-CM5A-LR	2074.2	432.3	1235.5	273.7	1941.5
	MIROC5	2012.5	457.5	1168.7	259.7	1885.9
VISIT	GFDL-ESM2M	1287.6	308.6	745.8	110.8	1165.2
	HadGEM2-ES	1334.8	317.0	782.4	112.7	1212.1
	IPSL-CM5A-LR	1271.5	307.8	729.6	106.9	1144.3
	MIROC5	1302.4	317.6	751.1	109.1	1177.8
ORCHIDEE-	GFDL-ESM2M	2164.6	459.8	1187.9	276.6	1924.4
MICT	HadGEM2-ES	2207.6	472.4	1227.5	269.6	1969.5
	IPSL-CM5A-LR	2269.9	472.8	1250.9	284.7	2008.3
	MIROC5	2099.1	469.4	1136.0	252.9	1858.2
DLEM	GFDL-ESM2M	790.3	211.0	469.9	30.8	711.7
	IPSL-CM5A-LR	817.2	224.4	483.5	<i>29.1</i>	737.0
	MIROC5	780.3	204.0	468.8	29.7	702.6
Model mean		1576.4	369.8	897.7	154.1	1421.6
Model median		1380.5	381.6	762.8	110.8	1209.7
Model IQR		1250.7	231.6	708.5	238.6	1175.7

Notes: the total indicates the total soil carbon in grid cells dominated by natural

1041

vegetation where the cropland fraction is less than 30% in 2005.

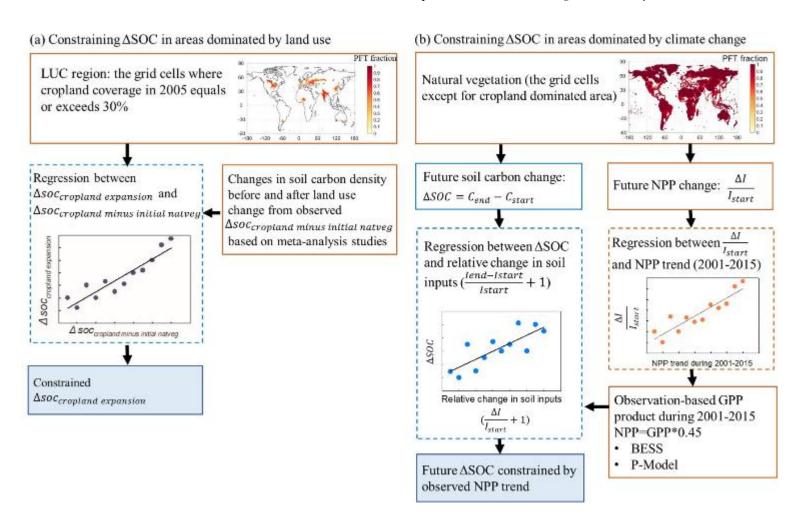


Figure 1 The framework of emergent constraint approach in areas dominated by land use change (a) and areas dominated by climate change (b).

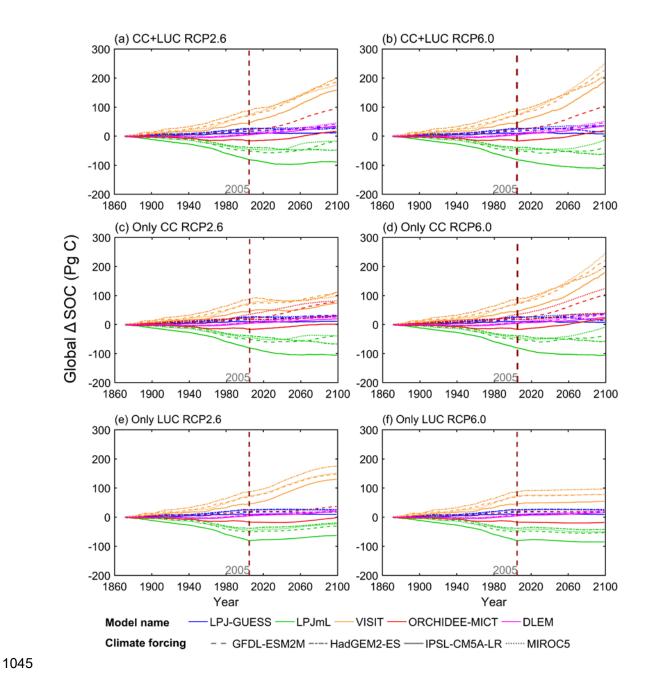


Figure 2 Changes of global soil organic carbon (ΔSOC) compared to the historical (1861-1870) under both the effects of climate change and land use change (CC+LUC), the effect of climate change (Only CC) and the effect of land use change (Only LUC) based on RCP 2.6 and 6.0 during the period of 1871-2099.

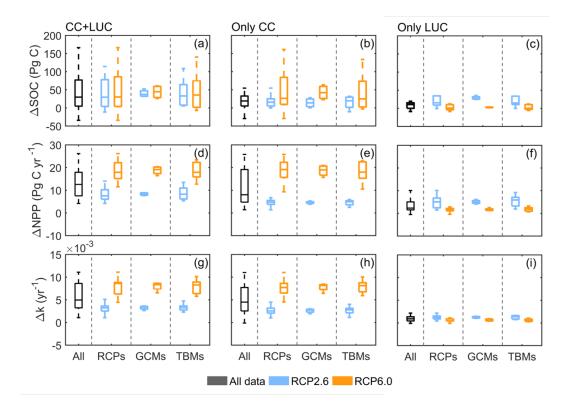


Figure 3 Change in global soil carbon (ΔSOC), net primarily productivity (ΔNPP), and decomposition rate (Δk) according the effects of all data, RCPs (i.e., RCP 2.6 and RCP 6.0), GCMs (i.e., four climate forcing, including GFDL-ESM2M, HadGEM2-ES, IPSL-CM5A-LR and MIROC5), and TBMs (i.e., LPJ-GUESS, LPJmL, VISIT and ORCHIDEE-MICT) under both effects of climate change and land use change (CC+LUC), the effect of climate change (Only CC) and the effect of land use change (Only LUC) over the period of 2090-2099 compared to the means of 1996-2005. 'All' indicates the range obtained by averaging all data; 'RCPs' indicates the range obtained by averaging all TBMs and GCMs outputs; 'GCMs' indicates the range obtained by averaging all TBMs outputs for each GCM; 'TBMs' indicates the range obtained by averaging all GCMs outputs for each TBM.

Global natural ecosystem under RCP 6.0

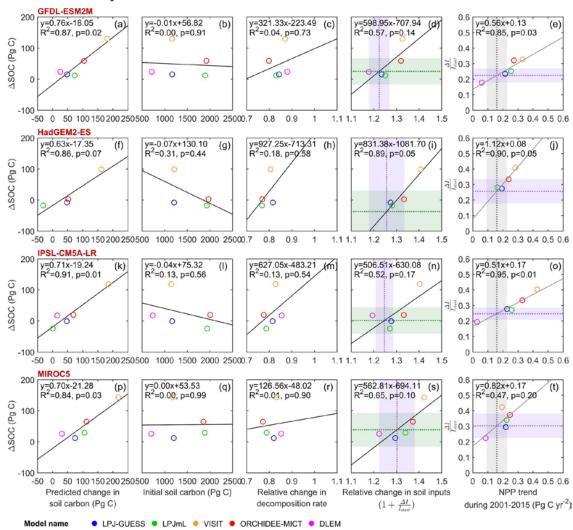


Figure 4 Change in global soil carbon stocks in area dominated by natural ecosystems 1062 between start time (1995-2005) and end time (2090-2099) global means as a function of Eq. 1063 (2) (the first column), the initial soil carbon stocks (C_{start} ; the second column), the relative 1064 change in decomposition rate $(\frac{1}{1+\frac{\Delta k}{k_{ctout}}})$; the third column), and the relative change in soil 1065 inputs $(1 + \frac{\Delta I}{I_{\text{above}}})$; the fourth column), and the relationship between future input change 1066 $(\frac{I_{end}-I_{start}}{I_{etent}})$ and NPP trend during the period of 2001-2015 across the ISIMIP2b TBMs (the 1067 1068 fifth column). All results shown here are from simulations driven by different GCMs' climate forcing under RCP 6.0 scenario. Group 2 simulations with climate change effect only are 1069 1070 used. The different colors indicate different TBMs. The black lines indicate the linear regression across TBMs for each GCMs climate forcing. The dotted grey lines and grey areas 1071 1072 indicate the observation-based NPP trend for the period 2001-2015. The dotted purple lines and purple areas indicate the constrained $\frac{\Delta I}{I_{start}}$ and $1 + \frac{\Delta I}{I_{start}}$. The dotted green lines and 1073 green areas indicate the constrained Δ SOC. 1074

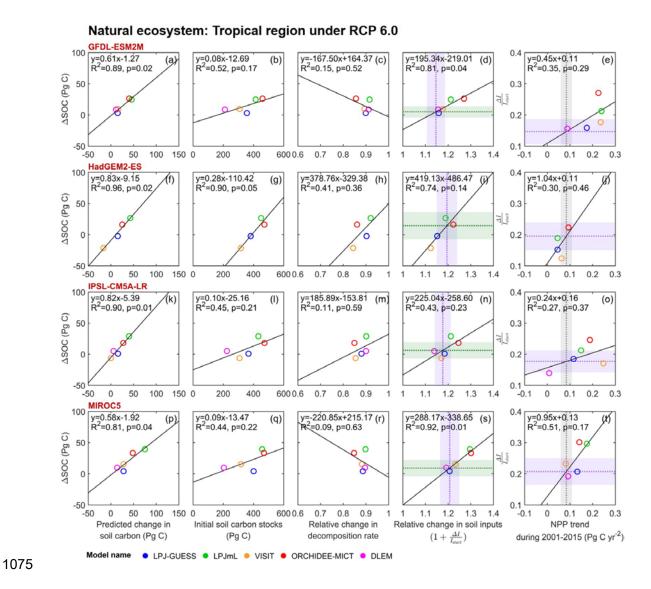


Figure 5 Change in global soil carbon stocks in tropical region dominated by natural ecosystems between start time (1995-2005) and end time (2090-2099) global means as a function of Eq. (2) (the first column), the initial soil carbon stocks (C_{start} ; the second column), the relative change in decomposition rate ($\frac{1}{1+\frac{\Delta I}{k_{start}}}$; the third column), and the relative change in soil inputs ($1+\frac{\Delta I}{I_{start}}$; the fourth column), and the relationship between future input change ($\frac{I_{end}-I_{start}}{I_{start}}$) and NPP trend during the period of 2001-2015 across the ISIMIP2b models (the fifth column). Detailed symbol and line information are in Figure 4.

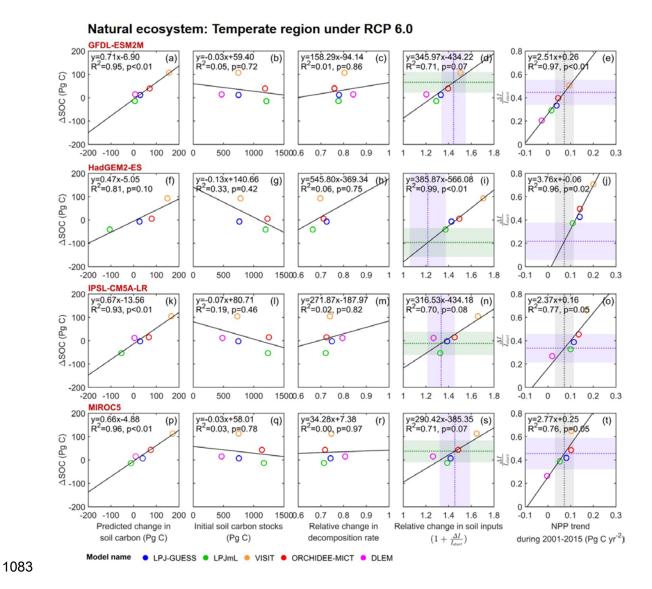


Figure 6 Change in global soil carbon stocks in temperate region dominated by natural ecosystems between start time (1995-2005) and end time (2090-2099) global means as a function of Eq. (2) (the first column), the initial soil carbon stocks (C_{start} ; the second column), the relative change in decomposition rate ($\frac{1}{1+\frac{\Delta k}{k_{start}}}$; the third column), and the relative change in soil inputs ($1+\frac{\Delta I}{I_{start}}$; the fourth column), and the relationship between future input change ($\frac{I_{end}-I_{start}}{I_{start}}$) and NPP trend during the period of 2001-2015 across the ISIMIP2b models (the fifth column). Detailed symbol and line information are in Figure 4.

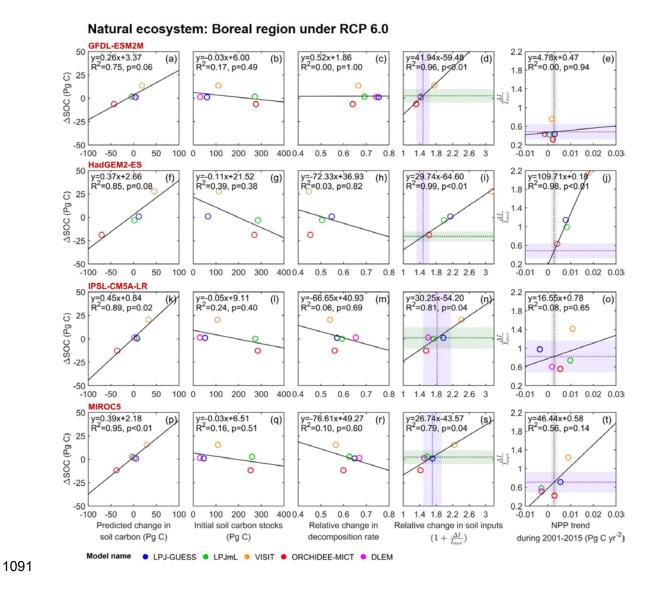


Figure 7 Change in global soil carbon stocks in boreal region dominated by natural ecosystems between start time (1995-2005) and end time (2090-2099) global means as a function of Eq. (2) (the first column), the initial soil carbon stocks (C_{start} ; the second column), the relative change in decomposition rate ($\frac{1}{1+\frac{\Delta k}{k_{start}}}$; the third column), and the relative change in soil inputs ($1+\frac{\Delta I}{I_{start}}$; the fourth column), and the relationship between future input change ($\frac{I_{end}-I_{start}}{I_{start}}$) and NPP trend during the period of 2001-2015 across the ISIMIP2b models (the fifth column). Detailed symbol and line information are in Figure 4.

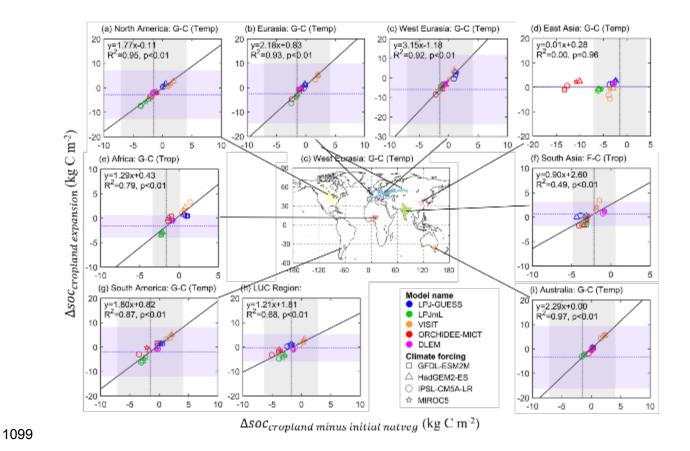


Figure 8 Relationship between modeled SOC density changes across cropland expansion area ($\Delta soc_{cropland\ expansion}$) during the period of 1861-2005 and the soil carbon density difference ($\Delta soc_{cropland\ minus\ initial\ natureg}$) between current cropland (in 2005) and initial natural vegetation (in 1861). Group 1 simulations were used in this analysis. The bottom center panel shows the results at global land use dominated areas. The different colors and symbols indicate different climate forcing and models, respectively. The black lines indicate the linear regression across all TBMs and all GCMs. The dotted grey lines and grey areas indicate the observation-based meta-analysis data. The dotted purple lines and purple areas

indicate the constrained $\Delta soc_{cropland\ expansion}$.