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Empirical estimates of regional carbon budgets imply reduced global soil heterotrophic respiration

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ABSTRACT

Resolving regional carbon budgets is critical for informing land-based mitigation policy. For nine regions covering nearly the whole globe, we collected inventory estimates of carbon-stock changes complemented by satellite estimates of biomass changes where inventory data are missing. The net land–atmospheric carbon exchange (NEE) was calculated by taking the sum of the carbon-stock change and lateral carbon fluxes from crop and wood trade, and riverine-carbon export to the ocean. Summing up NEE from all regions, we obtained a global ‘bottom-up’ NEE for net land anthropogenic CO₂ uptake of -2.2 ± 0.6 PgC yr⁻¹ consistent with the independent top-down NEE from the global atmospheric carbon budget during 2000–2009. This estimate is so far the most comprehensive global bottom-up carbon budget accounting, which set up an important milestone for global carbon-cycle studies. By decomposing NEE into component fluxes, we found that global soil heterotrophic respiration amounts to a source of CO₂ of 39 PgC yr⁻¹ with an interquartile of 33–46 PgC yr⁻¹—a much smaller portion of net primary productivity than previously reported.

Keywords: carbon budget, human appropriation of ecosystems, soil carbon

REGIONAL NET LAND–ATMOSPHERE CARBON-EXCHANGE ESTIMATES FROM BOTTOM-UP INVENTORIES AND LATERAL CARBON FLUXES

Net ecosystem exchange (NEE) is defined as the land–atmosphere flux of carbon excluding fossil-fuel emissions [1,2]. Accurate and consistent observation-based estimates of NEE at regional scales are needed to understand the global land-carbon sink, to evaluate land-carbon models used

for carbon-budget assessments and future climate projections and to define baselines for land-based mitigation efforts. Currently, land-carbon models show significant disagreement in their quantification of regional carbon fluxes [3,4]. A comprehensive global assessment of regional NEE from bottom-up approaches using land observations is still missing, which hampers an independent evaluation of the top-down global NEE deduced from the atmospheric CO₂ growth rate [5]. In addition, understanding the components of regional NEE allows

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the turnover time of carbon lost to the atmosphere by soil heterotrophic respiration (SHR) and other processes and the response of land-carbon storage to increasing atmospheric CO₂ (β) and warming (γ) to be better quantified [6].

We quantified NEE and its component fluxes for nine large land regions that cover nearly the entire land surface, using land-carbon-stock-change estimates reported from different inventories and observation-based data [7–16]. Special attention was brought to include uncertainties reported in each original publication and to use different data sources wherever available to account for uncertainty between data sets (Supplementary Table 1). NEE, the net carbon flux exchanged between each region and the atmosphere, excluding fossil-fuel emissions, is diagnosed from:

$$NEE = \Delta C + F_{rivers} + F_{trade}, \quad (1)$$

where ΔC is the net carbon-stock change in living and dead biomass, soil carbon, wood and crop products; F_{rivers} is the flux of particulate and dissolved organic and inorganic carbon lost by each region to the ocean; and F_{trade} is the net carbon flux from crop and wood products. F_{trade} is a gain of carbon by a region if it imports more than it exports or a loss if otherwise (Supplementary Fig. 1). In Equation (1), the sign convention is that NEE is negative when a region removes CO₂ from the atmosphere, ΔC is negative when carbon storage increases in that region, F_{rivers} is always positive and F_{trade} can be positive (net export) or negative (net import).

We estimated ΔC in each region based on publications from the REgional Carbon Cycle Assessment and Processes (RECCAP) project [7–14] completed by other data sources (Supplementary Table 1). The nine RECCAP regions cover the entire land surface except the Middle East, Ukraine, Belarus, Kazakhstan and New Zealand. These latter regions, given their surface area and/or their sparse vegetation, represent only a small fraction of global ΔC (Supplementary Table 1). For humid forests in tropical South America and Africa, ΔC of biomass was provided from long-term forest-plots data [8,12], but there were no such data for Southeast Asia and South Asia. To fill this gap, we used biomass ΔC from remote-sensing data combining Lidar and optical measurements [17]. This remote-sensing-based estimate of biomass ΔC is consistent with data from forest plots in South America and Africa, giving support to using it for Southeast Asia and South Asia (Supplementary Fig. 2). Further, unlike for other regions where regional data on soil-carbon-stock changes, albeit uncertain, were available, soil-

carbon ΔC was ignored in the tropical regions where no data exist with adequate coverage. We estimated from global carbon-model output that ignoring soil carbon in our estimate of tropical ΔC could lead to an average underestimate of 26%, which is still within the range of uncertainties (Supplementary Text 1 and Table 2). To assess F_{trade} , we used statistics of the wood and crop-products trade converted into carbon units [18]. To assess F_{rivers} , we calculated the total export of fluvial carbon to the oceans using data from references [19] and [20] and subtracted the fraction of fluvial carbon that originates from weathered carbonate minerals, which represents a transfer from the lithosphere to the ocean that does not involve a contemporary atmospheric C flux (see ‘Methods’ section). Error propagation was applied to each term of Equation (1) to quantify regional NEE uncertainties.

A NEW GLOBAL ESTIMATION OF LAND–ATMOSPHERE NET CARBON EXCHANGE

We obtained a bottom-up anthropogenic NEE of -2.2 ± 0.6 PgC yr⁻¹, totally independently of the value derived from the global CO₂ budget [5] of -2.4 ± 0.7 PgC yr⁻¹ but remarkably consistent with it. Our bottom-up estimate is calculated from the sum of all the regional NEE values (Supplementary Table 1) plus NEE from regions not covered by the RECCAP publications being a sink of -0.4 PgC yr⁻¹ (Supplementary Text 2 and Table 3). This gives a net uptake of atmospheric CO₂ equal to -2.8 ± 0.7 PgC yr⁻¹ for the period 2000–2009 (Fig. 1a). This estimate includes the sum of the anthropogenic sink linked to the human-induced perturbation of the global carbon cycle and the background natural CO₂ sink associated with the fact that there is always a fraction of atmospheric CO₂ absorbed on land by the vegetation, which is leached to rivers and transferred to the ocean [21]. In contrast, the NEE calculated from the global CO₂ budget [5] only quantifies the anthropogenic land-CO₂ uptake. We thus subtracted from our bottom-up NEE value the background sink from reference [21] to derive our bottom-up anthropogenic NEE estimate.

The contributions of each region to the global bottom-up NEE differ in magnitude. Tropical regions are small sinks of atmospheric CO₂ and have NEE values of -0.25 ± 0.11 PgC yr⁻¹ in South Asia, -0.17 ± 0.19 PgC yr⁻¹ in Southeast Asia, -0.07 ± 0.29 PgC yr⁻¹ in South America and -0.06 ± 0.29 PgC yr⁻¹ in Africa, summing up to a small tropical land-CO₂ sink of -0.55 ± 0.46 PgC yr⁻¹.

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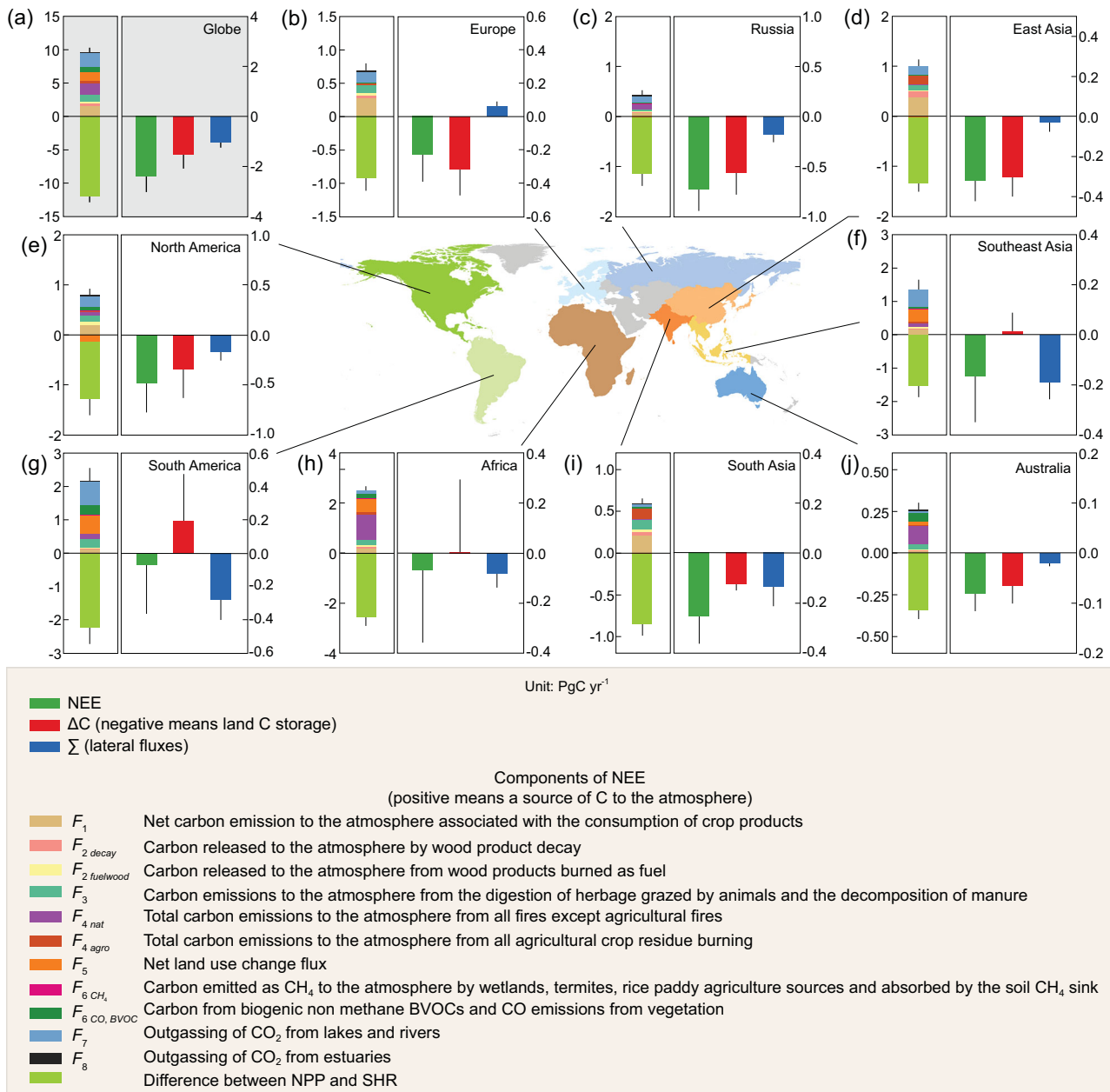


Figure 1. (a–j) Carbon-storage change from inventories (ΔC in red) and lateral fluxes from trade- and riverine-carbon export to the ocean for different regions of the globe (in blue) for the 2000s (Supplementary Fig. 1). The resulting bottom-up NEE from the sum of ΔC and lateral fluxes is given in green. An atmospheric convention is used, so that $\Delta C < 0$ denotes an increase in land-carbon stocks and $NEE < 0$ is also a net uptake of atmospheric CO₂. The upper stacked bars on the left show the NEE subcomponents that are sources of C to the atmosphere, excluding soil heterotrophic respiration, and the green bar is the resulting imbalance between the net primary productivity (NPP) and soil heterotrophic respiration, a negative value indicating that the soil heterotrophic respiration is smaller than NPP.

Northern-hemisphere regions tend to have larger CO₂ sinks: -0.23 ± 0.16 PgC yr⁻¹ in Europe, -0.32 ± 0.10 PgC yr⁻¹ in East Asia, -0.49 ± 0.3 PgC yr⁻¹ in North America and -0.73 ± 0.22 PgC yr⁻¹ in Russia. The NEE from these northern regions amounts to -1.8 ± 0.4 PgC yr⁻¹, which is within the range of independent estimates given by atmospheric inversions [5].

WHY LAND–ATMOSPHERE NET CARBON EXCHANGE DIFFERS FROM CARBON-STOCK CHANGES

Regional differences between NEE and ΔC are displayed in Fig. 1. In Russia, North America, South Asia and Australia, NEE is a larger CO₂ sink than the increase in carbon stocks (ΔC) because a fraction of

CO₂ fixed from the atmosphere is exported by trade and rivers. In Europe, however, NEE is a smaller CO₂ sink than ΔC because trade imports exceed riverine-carbon exports, given the fact that imported products are oxidized by humans and animals, releasing CO₂ into the atmosphere in these two regions. In Africa, Southeast Asia and South America, inventories show a reduction in carbon stocks (positive ΔC values in Fig. 1) because of deforestation, but NEE is still a small net sink of atmospheric CO₂ from the atmosphere, due to strong lateral exports from rivers and trade. These regional differences between NEE and ΔC clearly demonstrate that the results of top-down atmospheric inversions estimating NEE are not comparable to bottom-up carbon-stock change ΔC from inventories at the regional scale. We thus recommend that stock-change-based regional carbon budgets should be corrected for lateral fluxes to be properly compared with inversion results [11]. At the scale of smaller regions that exchange a lot of carbon by trade circuits, such as crop-production basins, plantation areas exporting carbon products or populated areas importing and consuming those products, differences between inversions NEE and ΔC are expected to be even larger in relative value.

DEDUCING SHR CO₂ EMISSIONS FROM NET ECOSYSTEM CARBON EXCHANGE

SHR is one of the largest and arguably the most uncertain flux of the global land-carbon cycle. Here, we combined our new bottom-up NEE estimates with other component fluxes exchanged with the atmosphere to infer SHR ('Methods' section and Supplementary Fig. 3). Regional values of net primary productivity (NPP) were taken from three satellite-based products [22–24] and a global land-carbon-cycle data-assimilation system [25]. These products based on different approaches and different satellite sensors give a global NPP of -50 PgC yr^{-1} with an interquartile range (IQR) of -57 to -44 PgC yr^{-1} over all RECCAP regions (Monte-Carlo standard deviation across the four NPP estimates and the uncertainty of each estimate; see 'Methods' section). Those four different NPP estimates are also consistent with each other for each individual region, within their respective uncertainties (Supplementary Fig. 4). Once fixed by NPP, carbon turns over in ecosystem pools and is returned back to the atmosphere mainly by SHR, but also by land-use-change emissions; fires; livestock grazing and the harvest wood and crop products subsequently oxidized by humans and animals; outgassing

of carbon by lakes, rivers, and estuaries; and biogenic emissions of reduced-carbon compounds including methane and biogenic volatile organic compounds (Supplementary Fig. 3). All these gross fluxes and their uncertainties were estimated for each region using observational data sets, considering wherever possible different independent estimates for consistency checking ('Methods' section and Supplementary Table 1). Then, SHR, which is the largest single flux of CO₂ lost by the land to the atmosphere, was calculated for each region as a residual, namely as the difference between NEE, NPP and all other non-SHR carbon exchanges. Uncertainties of SHR were obtained using a Monte-Carlo approach, sampling different estimates and their individual uncertainties (Supplementary Table 1).

Among individual sources of carbon to the atmosphere that are not from SHR, we found that the largest flux is the outgassing by rivers, lakes and estuaries [26,27] ranging from 0.8 to 2.3 PgC yr⁻¹. This flux is followed by carbon emissions from fires (1.6 PgC yr⁻¹), the consumption of harvested crop products (1.5 PgC yr⁻¹), land-use-change emissions (1.0–1.2 PgC yr⁻¹), emissions from grazing (1.0 PgC yr⁻¹), biogenic reduced-carbon emissions such as methane and volatile biogenic compounds (0.8 PgC yr⁻¹) and the decay and burning of wood products (0.7 PgC yr⁻¹). Altogether, these fluxes represent a globally large source of carbon to the atmosphere of $8.3 \pm 0.9 \text{ PgC yr}^{-1}$. From these data combined with NPP (and our bottom-up NEE estimates), we infer a global median value of SHR equal to 39 PgC yr⁻¹ with an IQR of 33–46 PgC yr⁻¹ and a non-Gaussian uncertainty distribution obtained from a Monte-Carlo analysis across different available data sets and their internal uncertainty (Fig. 2). This new global value of SHR is significantly lower than the estimate of $\sim 50 \text{ PgC yr}^{-1}$ given in previous global assessments [28,29]. To obtain an independent verification of this lower SHR estimation, we upscaled data from 455 site-level measurements of SHR from a global database [30] (see 'Methods' section) into regional SHR budgets using two approaches. The first approach is a machine-learning algorithm (Supplementary Text 3) and the other one is an empirical model based on functions of soil moisture and soil temperature fitted to the site data [31] ('Methods' section and Supplementary Fig. 7). The two estimates of SHR obtained from site-data upscaling agree well with our value deduced from regional carbon budgets, at both regional and global scales (Fig. 2a), which leads to high confidence that SHR amounts to $\sim 40 \text{ PgC yr}^{-1}$.

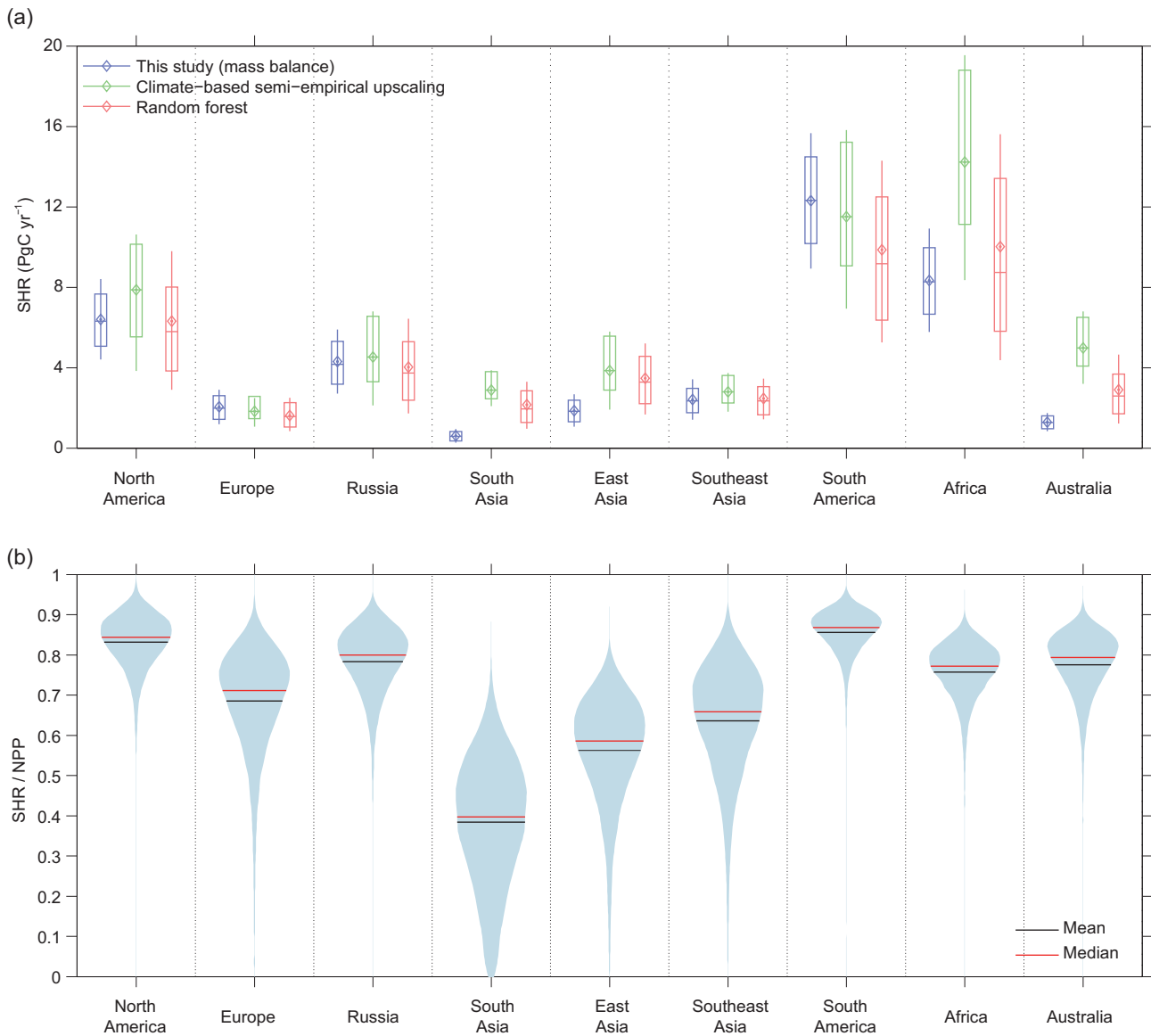


Figure 2. (a) Regional estimates of soil heterotrophic respiration from this study in blue and from two other estimates obtained using an independent approach from upscaling a global data set of site-level field measurements [38] by reference [31]. The whisker bars indicate the interquartile range and the full range. (b) Regional estimates of the ratio between soil heterotrophic respiration and NPP for each region with mean and median values of distributions calculated using a Monte-Carlo analysis (see ‘Methods’ section).

SHR CO₂ EMISSION IS A SMALLER FRACTION OF NPP THAN IN PREVIOUS ASSESSMENTS

The median ratio of SHR to NPP is 0.78 (IQ range from 0.76 to 0.80), which is smaller than the value reported by the third Intergovernmental Panel on Climate Change (IPCC) assessment report [28] of 0.9 and also less than the simulation results from Dynamic Global Vegetation Model (DGVM) carbon-cycle models [5] that give a median of 0.89 and a range of 0.77–0.9. DGVM-based estimates are likely to be overestimated because those models lack a description of the carbon lost to rivers and, in

some models, of the components of harvest, which represent carbon not delivered to the soil and thus not available as a substrate for SHR [32]. The ratios of SHR to NPP displayed in Fig. 2b show the lowest values in South Asia (0.38) and East Asia (0.56) where the NPP appropriation by human activities is large, as are fire losses and riverine export. The ratio of SHR to NPP varies between 0.64 and 0.86 in other regions (Fig. 3b). In regions with a large fraction of natural ecosystems like South America and Russia, the values of this ratio are mainly determined by fire losses, reduced-carbon emissions and inland-water outgassing.

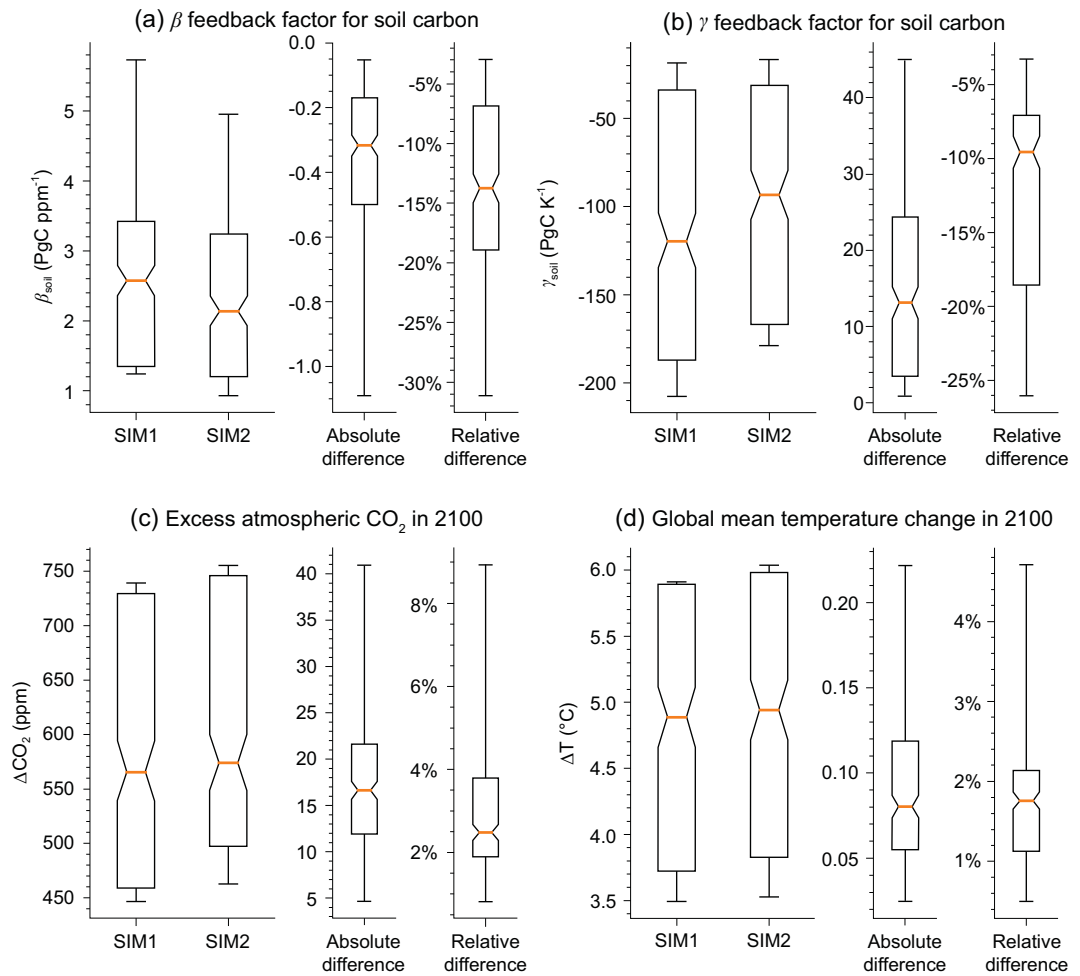


Figure 3. Carbon and climate feedback from soil-carbon simulated with the OSCAR coupled carbon-climate model when emulating seven Earth System Models of the 5th IPCC Assessment Report and following the high fossil-fuel emissions RCP8.5 scenario. In ‘SIM-1’, harvest fluxes, reduced-carbon-compound emissions and riverine export are ignored if not already accounted for in the ESM. In ‘SIM-2’, these lateral fluxes are always included, which altogether have the effect of reducing the flux of litter delivered to the soil and hence soil heterotrophic respiration relative to the net primary production. (a) and (b) β_{soil} is the change in soil-carbon storage per unit change in atmospheric CO_2 and γ_{soil} is the change in soil-carbon storage in response to a global warming of 1 K. (c) and (d) Atmospheric CO_2 and global mean temperature by year 2100. The configuration of SIM-1 that corresponds to that of the current-generation Earth System Models leads to under-prediction of the future increase in CO_2 and temperature. Boxes and whiskers show the interquartile and minimum-to-maximum ranges, from 224 sensitivity simulations of OSCAR (Supplementary Text 4). Both absolute (SIM2 – SIM1) and relative differences (SIM2 \div SIM1 – 1) are shown.

Overestimating the ratio of SHR to NPP in global land-carbon-cycle models [5] implies that too much of their NPP is transferred to soils compared to the reality, which gives rise to simulated carbon stocks that are too large and associated turnover times that are too long. However, a positive bias of SHR to NPP in models is not direct evidence that soil-carbon stocks and turnover times in model simulations are inaccurate because of the highly uncertain available soil-carbon observations and compensating errors of model parameterizations. Nevertheless, this structural bias of models should have two consequences for modeling future carbon storage in response to rising CO_2 and climate change. The

first one is that β_{soil} (the change in soil-carbon stock per ppm increase in CO_2) will be overestimated by models because the overestimated fraction of the future NPP will continue to reach the soil as litterfall, compared to the real world. The second consequence is that models will also overestimate γ_{soil} (the temperature sensitivity of soil carbon, defined by the loss of soil carbon in response to a 1°C global warming) because they over-predict the soil stock exposed to warming. In a coupled carbon-cycle-climate model, overestimating β_{soil} translates into underestimating the amount of CO_2 that will remain in the atmosphere in the future, whereas overestimating γ_{soil} results in overestimating the

amount of CO₂ released by soil warming in the future.

IMPLICATIONS FOR THE LAND-CARBON-CYCLE RESPONSES TO RISING CO₂ AND CLIMATE WARMING

To assess the net effect of these two compensatory effects of overestimating β_{soil} and γ_{soil} on future climate projections, we used the compact coupled carbon-climate model OSCAR [33]. OSCAR was calibrated to reproduce the same pre-industrial regional carbon stocks and NPP, the same sensitivity of NPP to CO₂ and temperature, and the same sensitivity of soil-carbon decomposition to temperature as seven of the Earth System Models (ESMs) from the 5th IPCC Assessment Report. This calibration ensures that, when run for a future scenario with imposed fossil CO₂ emissions, OSCAR gives results similar to each original ESM model for future CO₂ and temperature but at much lower computational costs.

We ran two sensitivity simulations with OSCAR, both forced by fossil CO₂ emissions from the RCP8.5 pathway [34]. Those emissions lead to an intensive warming and high future CO₂ concentration, and can be viewed as an extreme test to examine the effect of overestimating β_{soil} and γ_{soil} . In the first simulation (SIM-1), SHR was simulated according to each original ESM model (Supplementary Text 4). In the second simulation (SIM-2), we prescribed crop and wood harvest, land use, fires and grazing CO₂ emissions as a fraction of future NPP in order to reproduce our estimate of present-day SHR-to-NPP ratios for each region (Supplementary Text 4). For the future evolution of riverine-carbon export and outgassing, we constructed an empirical model of this flux based on relationships established by reference [26]. The set-up for SIM-2 is a compromise that avoids a detailed—and uncertain—mechanistic, complex representation of the river export, harvest and grazing. Yet it provides plausible magnitudes for future changes in those fluxes relative to NPP (Supplementary Table 6).

The values of β_{soil} and γ_{soil} calculated by OSCAR from SIM-1 and SIM-2 are shown in Fig. 3 for a combination of seven different ESMs and 32 estimates to account for uncertainty in the SHR-to-NPP ratios for each region. As expected, β_{soil} is 3%–31% lower in SIM-2 than in SIM-1, confirming that SIM-2 systematically reduces soil-carbon storage. Conversely, γ_{soil} is 3%–26% lower (less negative) in SIM-2 than in SIM-1 because there is less carbon in the soil exposed to warming. Although partially cancelling one another out, the reduction of β_{soil} in SIM-2

versus SIM-1 dominates over the reduction of γ_{soil} , leading to higher atmospheric CO₂ in the range of 5–41 ppm in 2100, as shown in Fig. 3c. Similarly, the increase of temperature in 2100 is larger in SIM-2 than in SIM-1, by 0.03–0.22°C (Fig. 3d).

The SIM-2 simulation should be seen as conservative because increasing future wood demand and livestock production in the RCP8.5 storyline [34] should enhance the human appropriation of NPP more than we assumed in the set-up of this simulation. The river-outgassing flux increases significantly in SIM-2, by 61% in 2100 compared to 2000–2009 in response to increased temperature, population and NPP. The development of reservoirs was ignored, which could also lead to both additional outgassing and carbon burial [35]. Presumably, other scenarios such as the RCP2.6, with its vast areas of bio-energy crops and harvested residues, should also significantly affect β_{soil} and γ_{soil} .

CONCLUSION

We conclude that soil carbon is temporally and spatially decoupled from NPP by lateral carbon fluxes from biomass harvest, grazing and carbon export to rivers, as well as by emissions of reduced biogenic carbon compounds. These fluxes are already important today in regions with high human appropriation of NPP and a strong leaching of soil carbon to rivers. The current generation of land-carbon models lack a representation of these lateral fluxes and the underpinning processes, and of their responses to climate change and human pressure. These fluxes are a first-order effect for estimating carbon-climate feedback and we thus recommend incorporating them in the next generation of ESMs and using spatially explicit data sets of harvest and grazing for model evaluation [36]. Likewise, national, regional and global carbon budgets used for the purpose of reporting progress towards targets in climate mitigation all need to account for lateral carbon flows and their consequences for the vertical exchange of carbon between the land biosphere and the atmosphere [37].

METHODS

Bottom-up NEE

For each region, NEE is the net flux of carbon exchanged with the overlying atmosphere, excluding fossil-carbon emissions given by Equation (1) in which the total carbon-stock change during the period 2000–2009 (ΔC) is the sum of inventory-based estimates of carbon-storage changes in crop products δ_1 ; in wood products, including products

decaying in landfills δ_2 ; in biomass, litter and soil pools δ_3 ; and in inland-water pools corresponding to the carbon burial in sediments δ_4 . The lithogenic carbon-storage decrease in carbonate rocks from weathering δ_5 is not counted in ΔC because it does not contribute to NEE as it does not involve an exchange with the atmosphere, and it was subtracted from the total riverine export of carbon. For δ_1 , we used data reported by the RECCAP publications in Europe [16] and Australia [15] and assumed zero elsewhere, this storage term being nevertheless very small. δ_2 was taken as reported from RECCAP publications [7–14] or calculated by a wood-product-pools bookkeeping model with input being wood products used in each region from reference [39] and decay functions from reference [40]. δ_3 was taken from the RECCAP publications [1–18] based on inventories for forest-biomass and data-driven models for soil-carbon changes, for instance models calibrated using soil-carbon inventory data. Those inventories have generally a large uncertainty for soil, litter and dead-wood carbon stocks [41]. For the four tropical regions, changes in soil carbon were ignored and the magnitude of this bias on ΔC is estimated using global models (Supplementary Text 1). Extensive forest-biomass inventories cover northern-hemisphere regions and there are data from research plots in South America and Africa. The primary biomass inventory data are similar to those synthesized over 2000–2007 by reference [42] with some updates to cover the period 2000–2009. In South Asia and Southeast Asia, we used satellite-based estimates of biomass stock changes from Lidar and optical observations [17] updated for this study to a global coverage and averaged over the period 2003–2009 to maximize overlap with the period covered by RECCAP. One-sigma uncertainties of ΔC were taken as reported by the original RECCAP publications (all data given in Supplementary Table 1). Supplementary Fig. 3 presents a comparison of ΔC between RECCAP inventories and the remote-sensing product [17] for each region. Although the remote-sensing estimates of ΔC were only used to fill the gap of missing inventory data in South and Southeast Asia regions, the two independent estimates are within their respective uncertainty for all the other regions, except for Russia, where inventories indicate a larger carbon-storage increase. δ_4 is the burial of carbon in freshwater sediments in lakes and reservoirs, reported in the original RECCAP publications only for North America (20 TgC yr⁻¹), Europe (41 TgC yr⁻¹) and Russia (20 TgC yr⁻¹) and here completed by data from reference [43] for other regions.

The bottom-up estimate of NEE was calculated using Equation (1), which requires lateral fluxes

from trade and river exports of carbon to oceans, obtained as explained below.

F_{trade} is the net lateral flux of crop and wood products related to trade across the boundaries of each region, calculated as the sum of the export and import fluxes of crop and wood products. By convention, this flux is positive if a region is a net exporter. For crop products, we considered all products, directly and indirectly, thus covering a broader spectrum of processed crop products than appears in the FAOSTAT database. The carbon in crops is estimated based on FAO crop-production statistics with standard conversion factors to adjust for water and then carbon content [18]. The lateral flux of wood products is calculated in a similar manner, based on reference [18] for different forestry products. For roundwood (FAO code 1861), FAO-STAT data [44] were used and, for the products processed from roundwood and potentially entering international trade, the GTAP-MRIO data [45] were used. This approach considers all products containing roundwood, directly and indirectly, and covers a broader spectrum of processed wood products than appears in the FAO database.

F_{rivers} is the net export of biogenic carbon by inland waters to the ocean, including dissolved organic carbon (DOC), particulate organic carbon (POC), and dissolved inorganic carbon (DIC). This biogenic carbon export is in fact from atmospheric origin as it was fixed by NPP (predominately terrestrial). The border between inland waters and the ocean is the end of estuaries. The mask of the RECCAP regions is such that there is no river carbon flowing from one region into another, so that imports/export between regions can be ignored. F_{rivers} was calculated specifically for this study using DOC, POC and DIC at 0.5-degree resolution aggregated into each region, based on the GLOBALNEWS model [19] and data from reference [20]. Following Resplandy *et al.* [46], who recently re-estimated fluvial exports of DOC and POC and showed that these fluxes were underestimated by the GLOBALNEWS model, we used their estimates to rescale spatial explicit estimates from GLOBALNEWS. Only a fraction of DIC transported by rivers is biogenic, the rest being from the lithosphere. The fraction of river DIC from the lithosphere varies in each region because the weathering of carbonate minerals consumes 1/2 mole of atmospheric carbon per mole of DIC transported by rivers, whereas the weathering of silicate minerals consumes 1 mole of atmospheric carbon per mole of DIC transported by rivers. The biogenic and lithogenic proportions of river DIC exports were calculated for this study based on reference [20] and the lithogenic component was subtracted from F_{rivers} .

To determine SHR from bottom-up estimates of NEE, we used a mass balance equation, whose terms are detailed below:

$$\begin{aligned} \text{SHR} = & \text{NEE} - \text{NPP} - F_{\text{crop products}} \\ & - F_{\text{wood products}} - F_{\text{grazing}} - F_{\text{fires}} \\ & - F_{\text{LUC}} - F_{\text{reduced}} - F_{\text{outgas rivers+ lakes}} \\ & - F_{\text{outgas estuaries}} \end{aligned} \quad (2)$$

Uncertainties of SHR were obtained by error propagation from uncertainties of each term from Equation (2), which are independent of each other (see below). A Monte-Carlo approach was used to randomly sample different empirical estimates of component fluxes when available, assuming that each estimate has an equal probability, and then sampling the internal Gaussian uncertainty of each estimate. Different estimates reported in Supplementary Table 1 include four estimates of NPP, two estimates of F_{fires} , two estimates of F_{LUC} and two estimates of $F_{\text{outgas rivers + lakes}}$. Other fluxes have a single estimate and only their internal uncertainty was considered. It is important to note that the error in SHR is positively correlated with the error in NPP and other fluxes, so that the uncertainty of the SHR-to-NPP ratio is lower than that of each flux separately. This is accounted for by the Monte-Carlo approach used to calculate this ratio and shown theoretically (Supplementary Text 5).

NPP was estimated using four different empirical products averaged over each region: the MODIS product [22,47] evaluated against site data [48] reporting an accuracy of $\approx 20\%$, the GIMMS product based on AVHRR and MODIS satellite-vegetation-absorbed radiation fraction [24], the BETHY/DLR product based on SPOT-VEGETATION satellite albedo and LAI data [49] and the CARDAMOM carbon-cycle data-assimilation product [25]. Those four NPP products are based on different methods and different sensors. GIMMS and MODIS-NPP use a calibrated light-use-efficiency model while BETHY assimilates vegetation indices in a Soil-Vegetation-Atmosphere model. CARDAMOM does not use satellite observations of NPP. Supplementary Fig. 4 shows that the four largely independent NPP products give consistent estimates in each region, following the Global Climate Observing System definition of consistency [50]: ‘when the independent measurements agree to within their individual uncertainties.’ The global NPP from the four empirical products is also consistent with empirical estimates compiled by references [51] and [52]. We performed a Monte-Carlo analysis by sampling the four different NPP products and

assuming for each product a Gaussian uncertainty [48] derived from MODIS-NPP because MODIS-NPP is the only product with a formal uncertainty estimation [48] (relative std. dev. $\approx 20\%$). This gives a global median NPP of -50 PgC yr^{-1} with an IQR of -57 to -44 PgC yr^{-1} for the area of all the RECCAP regions (Supplementary Table 1).

$F_{\text{crop products}}$ is the carbon emission to the atmosphere from the consumption of crop products, calculated as the sum of domestically harvested products minus net export minus storage (δ_1) in each region [18] from FAO data. Crop products are consumed by animals and humans and no distinction was made between these two groups. Digestion of crop products by ruminants emits CH_4 -carbon counted in $F_{\text{crop products}}$ and hence not in F_{reduced} to avoid double counting. The fraction of consumed products channeled to sewage systems and lost to rivers instead of being emitted to the atmosphere was ignored, as this flux is globally small [21] (100 TgC yr^{-1}). The 1-sigma uncertainty of $F_{\text{crop products}}$ could not be formally established in the absence of global trade-statistics data independently from FAO and was set to 20%.

$F_{\text{wood products}}$ is the carbon emission from the decay and burning of wood products. The term $F_{\text{wood products decay}}$ was calculated using a bookkeeping model forced by inputs from the domestic harvest of non-fuelwood [39] minus net export plus imports, to simulate pool changes and losses using decay functions from reference [40]. The small fraction of product waste going to sewage waters and rivers was ignored. The term $F_{\text{wood products decay}}$ was calculated in carbon units, including carbon lost to the atmosphere as CH_4 in landfills. The uncertainty of this flux was set to 20%. $F_{\text{wood products burning}}$ was calculated from statistics of fuelwood consumption [44] and carbon-emission factors (including CO_2 , CO and CH_4) compiled in reference [53] and updated to include the fraction of carbon emitted as black carbon [54]. This flux includes commercial fuelwood, fuelwood gathered locally and burned as a fuel in households and industrial fuelwood. The uncertainty of this flux was estimated by accounting for uncertain fuel-consumption and emission factors [53].

F_{grazing} is the carbon emission from the digestion of herbage grazed by animals and the decomposition of manure, here only from grass digestion, because manure from crop-products digestion is already included in $F_{\text{crop products}}$. The grass requirement by animals was derived from a grass-biomass-use data set [55] for the year 2000. The evolution of grazed biomass during the period 2001–2009 was calculated based on changes in the total metabolizable energy (ME) of ruminants [56,57]. Actual grass intake was modeled by the ORCHIDEE-GM global

model of pasture ecosystems [56] constrained by data from reference [55] including grazing and cut-and-carry forage intake. F_{grazing} includes CH_4 -carbon emissions from manure and enteric fermentation from reference [58], CO_2 respiration during grazing and emissions from the use of milk and meat. Animal and manure products are assumed to decay in 1 year. The uncertainty of F_{grazing} mainly comes from the grass-biomass-use data [55], obtained from an IPCC tier 3 digestion-metabolism model [59,60] and livestock-population statistics, diet composition and feed quality from databases and surveys. The evolution of the grass-biomass use was estimated based on livestock-population and IPCC tier 2 methods for ME. Uncertainty associated with livestock populations should be <20% and uncertainty of digestibility, a key parameter describing feed quality, is also <20%. Because the uncertainty of the digestion-metabolism model was not estimated, we used the upper-bound uncertainty of the two above input data sources to assess a relative uncertainty of F_{grazing} of 20%.

F_{fires} is the carbon emission from fires including CO_2 , CO, CH_4 and black carbon, separated into emissions from crop-residues burning and emissions from other fire types. The residues-burning occurs through small-scale fires that are underestimated by global burned-area products. Further, some residues are burned out of the field and their emissions are missed by satellites. $F_{\text{fires crop residues}}$ was calculated from fuel-consumption and carbon-emission factors [53] updated to include black carbon [54]. Its uncertainty was estimated using a Monte-Carlo method accounting for uncertainties of fuel-consumption and emission factors [53]. Emissions from other fires $F_{\text{fires others}}$ were estimated using two different satellite-based data sets: GFED4 (www.globalfiredata.org) based on burned areas [61] and GFAS based on fire radiative power [62]. GFED4 is an update of the GFED3 product [63] with updated burned area [61] completed by a data set of small fires [64]. In tropical regions, deforestation causes fires and those emissions, being already included in F_{LUC} , were subtracted from $F_{\text{fires others}}$ using reference [63] to minimize double accounting. Because the uncertainty of $F_{\text{fires others}}$ was not formally established, we diagnosed it from the interannual standard deviation of annual emissions in each region during 2000–2009. We verified that this estimate of uncertainty is consistent with formal uncertainties reported for GFED3 [63]. Supplementary Fig. 8 compares GFED4 and GFAS fire emissions in each region.

F_{LUC} is the net land-use-change carbon flux including gross deforestation, secondary ecosystem regrowth, soil CO_2 emissions after land-use change

and forest-degradation emissions, with the latter estimated only for Africa (Supplementary Table 1). F_{LUC} can be positive or negative, depending on the region considered. F_{LUC} results from the imbalance between NPP, SHR and deforestation fires over areas historically affected by land-use change. In the absence of NPP and SHR measurements over those areas, F_{LUC} was treated as a separate flux component of NEE, so that SHR in Equation (2) is not including legacy heterotrophic respiration after LUC, but only respiration on natural and managed lands. We used two different estimates of F_{LUC} . The first estimate was based on estimates from regional data provided by the RECCAP publications, except for South Asia and Africa, for which the results from the global bookkeeping model of Houghton *et al.* [65] were used. The uncertainty of F_{LUC} was taken as reported in those publications. The second estimate of F_{LUC} is from the ‘Bookkeeping of Land Use Emissions’ (BLUE) model described by reference [66], which tracks changes in soil- and biomass-carbon stocks following land-use conversion and due to management practices (e.g. shifting cultivation) using biome-specific carbon densities and exponential response curves. The LUC-transition maps applied were those from LUH2v2.1h [67] from 1850 to 2017. In BLUE, key uncertainties relate to land-use-change areas and carbon densities. We performed sensitivity simulations using different versions of the model to estimate the uncertainty of F_{LUC} from the BLUE model. For land-use-change input data, the largest uncertainties for the period 2000–2009 (when FAO data and remote sensing enter the LUH2 land-use forcing) arise from shifting cultivation and the allocation of extensive grazing (‘rangelands’) on natural vegetation types, which are both hard to monitor in satellite imagery and inventories. We added and subtracted the difference of simulations with and without shifting cultivation [66] and used two simulations that respectively clear natural vegetation for rangeland expansion or keep natural vegetation intact under this process, the latter as in reference [68]. For carbon densities, we used the original carbon densities of BLUE and an alternative lower map of carbon densities [66]. The standard deviation of the five sensitivity runs plus the default run for each grid cell was used to quantify uncertainty assessment. BLUE data were extrapolated by the last available year (2012 in reference [66] and 2016 in reference [68]). Supplementary Fig. 9 compares RECCAP and BLUE estimates of F_{LUC} in each region.

F_{reduced} is the emission of reduced carbon, including biogenic CH_4 , non-methane biogenic volatile organic compounds (BVOC) and biogenic CO, excluding fires to minimize double counting. Biogenic

CH₄ emissions from wetlands, termites, rice-paddy agriculture and the small sink in soils were estimated using atmospheric inversions from reference [69]. CH₄ emissions from crop and wood products in landfills are already counted in $F_{crop\ products}$ and $F_{wood\ products}$ and CH₄ from animals and manure in $F_{crop\ products}$. The uncertainty of $F_{reduced}$ was calculated from the standard deviation of 11 inversions [69]. CH₄ emission from rice paddies was estimated by assuming a fraction of 18% of the total CH₄ emission from the agriculture and waste source from inversion results [69]. For biogenic carbon emissions BVOC and CO, we used the CLM-MEGAN2.1 gridded product [70] converted into units of carbon mass. CLM-MEGAN2.1 estimates biogenic emissions of CO and of ~150 BVOC compounds with the main contributions being from terpenes, isoprene, methanol, ethanol, acetaldehyde, acetone, α -pinene, β -pinene, t - β -ocimene, limonene, ethene and propene. The uncertainty of that flux was estimated as a percentage of the total flux for each region following sensitivity tests and multi-model comparisons [71], assuming the same relative uncertainty in percentage for all species. The regional uncertainties are: for North America 21%; Europe 35%; Russia 33%; South Asia, East Asia and Southeast Asia 30%; South America 22%; Africa 35%; and Australia 40%. The global emission relative uncertainty is 22%.

$F_{outgas\ rivers + lakes}$ is the outgassing of carbon from lakes and rivers. We used two recent spatially resolved estimates of this flux [26,27] calculated using the GLORICH river pCO₂ database, but with different data-selection criteria and upscaling techniques. The one of reference [27] was made for the COSCAT groups of watersheds, re-interpolated to the RECCAP regions. $F_{outgas\ rivers + lakes}$ from reference [26] produced on a 0.5° × 0.5° global grid does not include lakes (lakes were added from reference [27]). River outgassing from reference [26] amounts to half the value of reference [27] due to lower estimates of average river pCO₂ for the tropics and Siberia resulting from a more restrictive data-selection process and additional averaging effects from the statistical model applied in reference [26]. For estimating SHR (Equation 2), we assumed that carbon emitted by $F_{outgas\ rivers + lakes}$ originates entirely from terrestrial NPP. In reality, a fraction of this flux may originate from autotrophic carbon fixation in inland waters [72] and autotrophic respiration from flooded plants [73]. Uncertainty of $F_{outgas\ rivers + lakes}$ was taken from original publications [26,27]. It ranges from 20% to 50% of the mean, depending upon the region.

$F_{outgas\ estuaries}$ is the outgassing of CO₂ from estuaries [74] calculated based on a compilation of local

flux measurements and a global segmentation of the coastal zone into MARCATS units, which aggregate COSCAT regions. The estimation also accounts for different estuarine types (small deltas, tidal systems, lagoons, fjords). A fraction of $F_{outgas\ estuaries}$ may originate from autotrophic C fixation in tidal wetlands [75], which was ignored. The uncertainty of $F_{outgas\ estuaries}$ was set to 50%, based on expert judgment.

Other SHR estimates from site-level data

Two independent estimates of SHR were obtained to be compared with the one obtained from regional carbon data (Fig. 2). The first estimate of SHR is a gridded product from the extension of the approach of Hashimoto *et al.* [29], which estimates soil respiration (SR) (including live-root respiration) using non-linear functions of climate variables and using a global relationship between annual soil respiration and SHR [29,76]. Here, we use the version of this approach by Konings *et al.* [31], for which the parameters of both the climate-SR and SR-SHR relationships were recalculated using site-level data of SR and SHR from the most recent version of the Soil Respiration Data Base [77] (SRDB) (v20180216). These parameters were fitted to SR data from 1979 sites instead of the original 1638 sites in reference [38]. Similarly, 362 measurement sites of SHR distributed across all biomes were used—a much larger data set than the original 53 measurements in the previous version of the SRDB database used by Hashimoto *et al.* [29].

The second SHR gridded product was produced for this study using site-level data from the same SRDB data set upscalled but with a random forest (RF) machine-learning algorithm (Supplementary Text 3). We used 455 site-year observations in total after data filtering (Supplementary Fig. 5) from the following criteria: (i) removing records without complete temporal, coordinates information and annual SHR information; (ii) excluding observations from manipulation experiments and soda-lime-application experiments, which would underestimate soil-CO₂ fluxes [78]. The RF algorithm applied here is composed of an ensemble of regression trees from bootstrapped training data [79]. We trained RF models with predictor variables including annual temperature, precipitation, soil moisture and radiation, gross primary productivity (GPP), nitrogen deposition, soil-carbon (depths of 0–30 and 30–100 cm) and soil-nitrogen content (Supplementary Table 4). Missing meteorological measurements at SHR sites are filled by gridded climatological driver data based on their coordinates. The values of the remaining predictor variables are

extracted at each site location from gridded driving data (Supplementary Table 4). We evaluated the RF-model performance using Leave One Out cross-validation (Supplementary Fig. 6) giving a correlation (R^2) of 0.57 and a RMSE of $111 \text{ gC m}^{-2} \text{ yr}^{-1}$ between observed and predicted values. Simulations of SHR were performed on a 0.5° grid over the globe using gridded predictors fields and averaged for each RECCAP region. The uncertainty of regional mean values of SHR was obtained by modeling the quantiles of this flux using the ‘QuantregForest’ package [80] in R.

SUPPLEMENTARY DATA

Supplementary data are available at [NSR](#) online.

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AUTHOR CONTRIBUTIONS

P. Ciais designed the study and wrote the manuscript. Y. Yao computed estimates of heterotrophic respiration and key figures. T. Gasser designed and performed simulations with the OSCAR model. A. Baccini provided new estimates of biomass changes from EO-data. Y. Wang prepared key figures and performed the uncertainty analysis. Other co-authors contributed specific data and analysis and helped to improve the manuscript.

Conflict of interest statement. None declared.

REFERENCES

- Canadell JG, Ciais P and Gurney K *et al.* An international effort to quantify regional carbon fluxes. *EOS (Washington DC)* 2011; **92**: 81–2.
- Ito A, Inatomi M and Huntzinger DN *et al.* Decadal trends in the seasonal-cycle amplitude of terrestrial CO_2 exchange resulting from the ensemble of terrestrial biosphere models. *Tellus Ser B Chem Phys Meteorol* 2016; **68**: 28968.
- Sitch S, Huntingford C and Gedney N *et al.* Evaluation of the terrestrial carbon cycle, future plant geography and climate-carbon cycle feedbacks using five Dynamic Global Vegetation Models (DGVMs). *Glob Chang Biol* 2008; **14**: 2015–39.
- Bastos A, O’Sullivan M and Ciais P *et al.* Sources of uncertainty in regional and global terrestrial CO_2 exchange estimates. *Glob Biogeochem Cycles* 2020; **34**: e2019GB006393.
- Friedlingstein P, Jones MW and O’Sullivan M *et al.* Global Carbon Budget 2019. *Earth Syst Sci Data* 2019; **11**: 1783–838.
- Friedlingstein P, Cox P and Betts R *et al.* Climate-carbon cycle feedback analysis: results from the C4MIP model intercomparison. *J Clim* 2006; **19**: 3337–53.
- King AW, Andres RJ and Davis KJ *et al.* North America’s net terrestrial CO_2 exchange with the atmosphere 1990–2009. *Biogeosciences* 2015; **12**: 399–414.
- Valentini R, Arneeth A and Bombelli A *et al.* A full greenhouse gases budget of Africa: synthesis, uncertainties, and vulnerabilities. *Biogeosciences* 2014; **11**: 381–407.
- Schulze ED, Ciais P and Luysaert S *et al.* The European carbon balance. Part 4: integration of carbon and other trace-gas fluxes. *Glob Chang Biol* 2010; **16**: 1451–69.
- Dolman AJ, Shvidenko A and Schepaschenko D *et al.* An estimate of the terrestrial carbon budget of Russia using inventory-based, eddy covariance and inversion methods. *Biogeosciences* 2012; **9**: 5323–40.
- Patra PK, Canadell JG and Houghton RA *et al.* The carbon budget of South Asia. *Biogeosciences* 2013; **10**: 513–27.
- Gloor M, Gatti L and Brienen R *et al.* The carbon balance of South America: a review of the status, decadal trends and main determinants. *Biogeosciences* 2012; **9**: 5407–30.
- Piao SL, Ito A and Li SG *et al.* The carbon budget of terrestrial ecosystems in East Asia over the last two decades. *Biogeosciences* 2012; **9**: 3571–86.
- Cervarich M, Shu S and Jain AK *et al.* The terrestrial carbon budget of South and Southeast Asia. *Environ Res Lett* 2016; **11**: 105006.
- Haverd V, Raupach MR and Briggs PR *et al.* The Australian terrestrial carbon budget. *Biogeosciences* 2013; **10**: 851–69.
- Luyssaert S, Abril G and Andres R *et al.* The European land and inland water CO_2 , CO, CH_4 and N_2O balance between 2001 and 2005. *Biogeosciences* 2012; **9**: 3357–80.
- Baccini A, Walker W and Carvalho L *et al.* Tropical forests are a net carbon source based on aboveground measurements of gain and loss. *Science* 2017; **358**: 230–4.
- Peters GP, Davis SJ and Andrew R. A synthesis of carbon in international trade. *Biogeosciences* 2012; **9**: 3247–76.

19. Mayorga E, Seitzinger SP and Harrison JA *et al.* Global nutrient export from WaterSheds 2 (NEWS 2): model development and implementation. *Environ Model Softw* 2010; **25**: 837–53.
20. Hartmann J, Jansen N and Dürr HH *et al.* Global CO₂-consumption by chemical weathering: what is the contribution of highly active weathering regions? *Glob Planet Chang* 2009; **69**: 185–94.
21. Regnier P, Friedlingstein P and Ciais P *et al.* Anthropogenic perturbation of the carbon fluxes from land to ocean. *Nat Geosci* 2013; **6**: 597–607.
22. Running SW, Nemani RR and Heinsch FA *et al.* A continuous satellite-derived measure of global terrestrial primary production. *Bioscience* 2004; **54**: 547–60.
23. Wißkirchen K, Tum M and Günther KP *et al.* Quantifying the carbon uptake by vegetation for Europe on a 1 km² resolution using a remote sensing driven vegetation model. *Geosci Model Dev* 2013; **6**: 1623–40.
24. Kolby-Smith W, Reed SC and Cleveland CC *et al.* Large divergence of satellite and Earth system model estimates of global terrestrial CO₂ fertilization. *Nat Clim Chang* 2015; **6**: 306–10.
25. Bloom AA, Exbrayat J-F and van der Velde IR *et al.* The decadal state of the terrestrial carbon cycle: global retrievals of terrestrial carbon allocation, pools, and residence times. *Proc Natl Acad Sci USA* 2016; **113**: 1285–90.
26. Lauerwald R, Laruelle GG and Hartmann J *et al.* Spatial patterns in CO₂ evasion from the global river network. *Glob Biogeochem Cycles* 2015; **29**: 534–54.
27. Raymond PA, Hartmann J and Lauerwald R *et al.* Global carbon dioxide emissions from inland waters. *Nature* 2013; **503**: 355–9.
28. Prentice IC, Farquhar GD and Fasham MJR *et al.* The carbon cycle and atmospheric carbon dioxide. In: Houghton JT, Ding Y and Griggs DJ (eds.). *Climate Change 2001: the Scientific Basis. Contribution of working group I to the third assessment report of the Intergovernmental Panel on Climate Change*. Cambridge: Cambridge University Press, 2001.
29. Hashimoto S, Carvalhais N and Ito A *et al.* Global spatiotemporal distribution of soil respiration modeled using a global database. *Biogeosciences* 2015; **12**: 4121–32.
30. Beer C, Reichstein M and Tomelleri E *et al.* Terrestrial gross carbon dioxide uptake: global distribution and covariation with climate. *Science* 2010; **329**: 834–8.
31. Konings AG, Bloom AA and Liu J *et al.* Global, satellite-driven estimates of heterotrophic respiration. *Biogeosciences* 2019; **16**: 2269–84.
32. Arneith A, Sitch S and Pongratz J *et al.* Historical carbon dioxide emissions caused by land-use changes are possibly larger than assumed. *Nat Geosci* 2017; **10**: 79–84.
33. Gasser T, Ciais P and Boucher O *et al.* The compact Earth system model OSCAR v2.2: description and first results. *Geosci Model Dev* 2017; **10**: 271–319.
34. Riahi K, Rao S and Krey V *et al.* RCP 8.5: a scenario of comparatively high greenhouse gas emissions. *Clim Chang* 2011; **109**: 33–57.
35. Tranvik LJ, Downing JA and Cotner JB *et al.* Lakes and reservoirs as regulators of carbon cycling and climate. *Limnol Oceanogr* 2009; **54**: 2298–314.
36. Krausmann F, Erb K-H and Gingrich S *et al.* Global human appropriation of net primary production doubled in the 20th century. *Proc Natl Acad Sci USA* 2013; **110**: 10324–9.
37. Kirschbaum MUF, Zeng G and Ximenes F *et al.* Towards a more complete quantification of the global carbon cycle. *Biogeosciences* 2019; **16**: 831–46.
38. Bond-Lamberty B. New techniques and data for understanding the global soil respiration flux. *Earth's Future* 2018; **6**: 1176–80.
39. Food and Agriculture Organization of the United Nations. *Forest Products 2016*. Rome: FAO, 2018. Available at <http://www.fao.org/3/I9987M/I9987m.pdf>.
40. Mason Earles J, Yeh S and Skog KE. Timing of carbon emissions from global forest clearance. *Nat Clim Chang* 2012; **2**: 682–5.
41. Pan Y, Birdsey RA and Phillips OL *et al.* The structure, distribution, and biomass of the world's forests. *Annu Rev Ecol Evol Syst* 2013; **44**: 593–622.
42. Pan Y, Birdsey RA and Fang JY *et al.* A large and persistent carbon sink in the world's forests. *Science* 2011; **333**: 988–93.
43. Mendonça R, Müller RA and Clow D *et al.* Organic carbon burial in global lakes and reservoirs. *Nat Commun* 2017; **8**: 1694.
44. Food and Agriculture Organization of the United Nations. FAOSTAT Statistical Database. 1997.
45. Peters GP, Andrew R and Lennox J. Constructing a multi-regional input-output table using the GTAP database. *Econ Syst Res* 2011; **23**: 131–52.
46. Resplandy L, Keeling RF and Rödenbeck C *et al.* Revision of global carbon fluxes based on a reassessment of oceanic and riverine carbon transport. *Nat Geosci* 2018; **11**: 504–9.
47. Zhao M, Heinsch FA and Nemani RR *et al.* Improvements of the MODIS terrestrial gross and net primary production global data set. *Remote Sens Environ* 2005; **95**: 164–76.
48. Turner DP, Ritts WD and Cohen WB *et al.* Evaluation of MODIS NPP and GPP products across multiple biomes. *Remote Sens Environ* 2006; **102**: 282–92.
49. Tum M, Zeidler JN and Günther KP *et al.* Global NPP and straw bioenergy trends for 2000–2014. *Biomass Bioenerg* 2016; **90**: 230–6.
50. Immler FJ, Dykema J and Gardiner T *et al.* Reference quality upper-air measurements: guidance for developing GRUAN data products. *Atmos Meas Tech* 2010; **3**: 1217–31.
51. Carvalhais N, Forkel M and Khomik M *et al.* Global covariation of carbon turnover times with climate in terrestrial ecosystems. *Nature* 2014; **514**: 213–7.
52. Ito A. A historical meta-analysis of global terrestrial net primary productivity: are estimates converging? *Glob Chang Biol* 2011; **17**: 3161–75.
53. Wang R, Tao S and Ciais P *et al.* High-resolution mapping of combustion processes and implications for CO₂ emissions. *Atmos Chem Phys* 2013; **13**: 5189–203.
54. Wang R, Balkanski Y and Boucher O *et al.* Estimation of global black carbon direct radiative forcing and its uncertainty constrained by observations. *J Geophys Res Atmos* 2016; **121**: 5948–71.
55. Herrero M, Havlík P and Valin H *et al.* Biomass use, production, feed efficiencies, and greenhouse gas emissions from global livestock systems. *Proc Natl Acad Sci USA* 2013; **110**: 20888–93.
56. US Climate Change Science Program. *The First State of the Carbon Cycle Report (SOCCR): The North American Carbon Budget and Implications for the Global Carbon Cycle*. A Report by the US Climate Change Science Program and the Subcommittee on Global Change Research. Asheville: National Oceanic and Atmospheric Administration, National Climatic Data Center, 2007.
57. Eggleston S, Buendia L and Miwa K *et al.* *2006 IPCC Guidelines for National Greenhouse Gas Inventories. Volume 4: Agriculture, Forestry and Other Land Use*. Hayama: Institute for Global Environmental Strategies (GES), 2006.
58. Pinares-Patiño CS, D'Hour P and Jouany J-P *et al.* Effects of stocking rate on methane and carbon dioxide emissions from grazing cattle. *Agric Ecosyst Environ* 2007; **121**: 30–46.
59. Herrero M, Thornton PK and Kruska R *et al.* Systems dynamics and the spatial distribution of methane emissions from African domestic ruminants to 2030. *Agric Ecosyst Environ* 2008; **126**: 122–37.
60. Thornton PK and Herrero M. Potential for reduced methane and carbon dioxide emissions from livestock and pasture management in the tropics. *Proc Natl Acad Sci USA* 2010; **107**: 19667–72.
61. Giglio L, Randerson JT and van der Werf GR. Analysis of daily, monthly, and annual burned area using the fourth-generation global fire emissions database (GFED4). *J Geophys Res Biogeosci* 2013; **118**: 317–28.

62. Di Giuseppe F, Rémy S and Pappenberger F *et al.* Using the Fire Weather Index (FWI) to improve the estimation of fire emissions from fire radiative power (FRP) observations. *Atmos Chem Phys* 2018; **18**: 5359–70.
63. van der Werf GR, Randerson JT and Giglio L *et al.* Global fire emissions and the contribution of deforestation, savanna, forest, agricultural, and peat fires (1997–2009). *Atmos Chem Phys* 2010; **10**: 11707–35.
64. Randerson JT, Chen Y and van der Werf GR *et al.* Global burned area and biomass burning emissions from small fires. *J Geophys Res* 2012; **117**: G04012.
65. Houghton RA, House JI and Pongratz J *et al.* Carbon emissions from land use and land-cover change. *Biogeosciences* 2012; **9**: 5125–42.
66. Hansis E, Davis SJ and Pongratz J. Relevance of methodological choices for accounting of land use change carbon fluxes. *Glob Biogeochem Cycles* 2015; **29**: 1230–46.
67. Hurtt GC, Chini LP and Frolking S *et al.* Harmonization of land-use scenarios for the period 1500–2100: 600 years of global gridded annual land-use transitions, wood harvest, and resulting secondary lands. *Clim Chang* 2011; **109**: 117–61.
68. Le Quéré C, Andrew RM and Friedlingstein P *et al.* Global carbon budget 2017. *Earth Syst Sci Data* 2018; **10**: 405–48.
69. Bousquet P, Ringeval B and Pison I *et al.* Source attribution of the changes in atmospheric methane for 2006–2008. *Atmos Chem Phys* 2011; **11**: 3689–700.
70. Guenther AB, Jiang X and Heald CL *et al.* The Model of Emissions of Gases and Aerosols from Nature version 2.1 (MEGAN2.1): an extended and updated framework for modeling biogenic emissions. *Geosci Model Dev* 2012; **5**: 1471–92.
71. Sindelarova K, Granier C and Bouarar I *et al.* Global data set of biogenic VOC emissions calculated by the MEGAN model over the last 30 years. *Atmos Chem Phys* 2014; **14**: 9317–41.
72. Hotchkiss ER, Hall Jr RO and Sponseller RA *et al.* Sources of and processes controlling CO₂ emissions change with the size of streams and rivers. *Nat Geosci* 2015; **8**: 696–9.
73. Abril G, Martinez J-M and Artigas LF *et al.* Amazon River carbon dioxide outgassing fuelled by wetlands. *Nature* 2014; **505**: 395–8.
74. Laruelle GG, Dürr HH and Lauerwald R *et al.* Global multi-scale segmentation of continental and coastal waters from the watersheds to the continental margins. *Hydrol Earth Syst Sci* 2013; **17**: 2029–51.
75. Bauer JE, Cai W-J and Raymond PA *et al.* The changing carbon cycle of the coastal ocean. *Nature* 2013; **504**: 61–70.
76. Bond-Lamberty B, Wang C and Gower ST. A global relationship between the heterotrophic and autotrophic components of soil respiration? *Glob Chang Biol* 2004; **10**: 1756–66.
77. Bond-Lamberty B and Thomson A. A global database of soil respiration data. *Biogeosciences* 2010; **7**: 1915–26.
78. Haynes BE and Gower ST. Belowground carbon allocation in unfertilized and fertilized red pine plantations in northern Wisconsin. *Tree Physiol* 1995; **15**: 317–25.
79. Breiman L. Random forests. *Mach Learn* 2001; **45**: 5–32.
80. Meinshausen N. Quantile regression forests. *J Mach Learn Res* 2006; **7**: 983–99.