



No support for carbon storage of >1,000 GtC in northern peatlands

Zicheng Yu ^{1,2}✉, Fortunat Joos ³, Thomas K. Bauska ⁴, Benjamin D. Stocker^{5,6},
Hubertus Fischer ³, Julie Loisel ⁷, Victor Brovkin ⁸, Gustaf Hugelius ⁹,
Christoph Nehrbass-Ahles¹⁰, Thomas Kleinen ⁸ and Jochen Schmitt³

ARISING FROM J. E. Nichols and D. M. Peteet *Nature Geoscience* <https://www.nature.com/articles/s41561-019-0454-z> (2019)

Northern peatlands store large amounts of carbon: 500 ± 100 GtC, according to a consolidated estimate from a diversity of methods^{1–6}. However, Nichols and Peteet⁷ presented an estimate of 1,055 GtC, exceeding previous estimates of carbon stock in global peatlands² and in northern peatlands by a factor of two. Here we argue that this is an overestimate, caused by systematic bias introduced by their inclusion of ¹⁴C dates from mineral deposits and other unsuitable sites, the use of records that lack direct measurements of carbon density, and the methodology issues. Furthermore, their estimate is difficult to reconcile within the top-down constraints imposed by ice-core and marine records, and estimated contributions from other processes that affected the terrestrial carbon storage during the Holocene epoch.

Unsuitable datasets and methodology issues

Nichols and Peteet⁷ used the time-history approach² to estimate peatland carbon stocks and their evolution over time. Their area-specific net carbon accumulation rates (j_c), as shown in their Fig. 2c, have a Holocene mean value of $33.4\text{--}37.6$ gC m⁻² yr⁻¹ (median across three methods), which is 46–102% higher than previous estimates of $18.6\text{--}22.9$ gC m⁻² yr⁻¹ (refs 2,3). Why this difference? Nichols and Peteet calculated j_c from sedimentation rates (cm yr⁻¹) and carbon density (gC cm⁻³). We argue that both of these parameters were overestimated by the authors.

Sedimentation rates are biased by the inclusion of ¹⁴C dates derived from mineral-rich non-peat deposits. Nichols and Peteet claimed to include all the sites in the Neotoma Paleoecology Database labelled as peatlands or synonyms, such as bogs or fens. However, many of these records, despite being called ‘bogs’, are deposits that developed from initial lake stages. For example, Chatsworth Bog in Illinois (Neotoma ID 364) contains >12 m sediments but was a lake for most of its 14,000-year history. Mineral lacustrine sediment had almost completely filled the basin about 3,000 years ago, when it changed from a lake to a marl fen that accumulated peat. The large difference in j_c —up to 30 gC m⁻² yr⁻¹ during the early Holocene—between Nichols and Peteet⁷ and ref. 3 (using the same data compilation) was partly due to Nichols and Peteet’s inclusion of rapidly accumulating mineral deposits. In addition, many sites from

Nichols and Peteet originate from low-latitude locations that are not representative of the areas where the vast majority of northern peatland areas are located (their Fig. 1a and Supplementary Fig. 1); this also compromises their estimates.

As stated by Nichols and Peteet, “rather than individual measurements of carbon density, a median carbon density (g cm⁻³) was used to calculate the j_c from sedimentation rate (cm yr⁻¹)”. Thus, Nichols and Peteet fail to account for the variability in carbon density in different regions and among different types of peatland^{3,8}. For example, there is a more than twofold difference in carbon density between western European islands/continental Europe (0.028 gC cm⁻³; $n=449$) and western Canada (0.076 gC cm⁻³; $n=3,441$)³. Also, peat undergoes different degrees of decomposition and compaction with age, resulting in highly variable carbon density often observed along a single peat profile. Furthermore, using one median carbon density value for all sites that lack direct measurements is prone to introducing bias and greatly inflates j_c calculations, especially for mineral-rich deposits. The propagation analysis of carbon density uncertainties by Nichols and Peteet⁷ does not resolve this problem.

Furthermore, we find an inherent problem in Nichols and Peteet’s algorithm that inflates the sedimentation rates and total carbon storage. Their probabilistic method was initially developed in a case study from an Alaskan peatland⁹. Using their data⁹ and algorithm, we find that a composite stratigraphy of 197 cm in length (in their Table 2) would change to a 246-cm-long core. We arrived at this 24.5% increase in core length by summing the product of the sedimentation rate (as annotated on their Fig. 3d) and time duration (shown in their Fig. 3e) of their 10 core intervals. By the same argument, the observed peat carbon storage of 126.3 kgC m⁻² (as calculated from their Table 2) would change to 155.8 kgC m⁻², an increase of 23.4%. This case study demonstrates that the assumptions behind their probabilistic method artificially create new carbon mass. The same problem exists in Nichols and Peteet⁷, but unfortunately they did not provide their specific and complete data in order to reproduce their results and quantify the effects of this carbon mass inflation, as well as the effects of the inclusion of the erroneous data and the use of median carbon density values for filling data gaps.

¹Key Laboratory of Geographical Processes and Ecological Security in Changbai Mountains, School of Geographical Sciences, Northeast Normal University, Changchun, China. ²Department of Earth and Environmental Sciences, Lehigh University, Bethlehem, PA, USA. ³Climate and Environmental Physics, Physics Institute and Oeschger Centre for Climate Change Research, University of Bern, Bern, Switzerland. ⁴British Antarctic Survey, Cambridge, UK. ⁵Department of Environmental Systems Science, ETH Zürich, Zürich, Switzerland. ⁶Swiss Federal Institute for Forest, Snow and Landscape Research WSL, Birmensdorf, Switzerland. ⁷Department of Geography, Texas A&M University, College Station, TX, USA. ⁸Max-Planck Institute for Meteorology, Hamburg, Germany. ⁹Department of Physical Geography, Stockholm University, Stockholm, Sweden. ¹⁰Department of Earth Sciences, University of Cambridge, Cambridge, UK. ✉e-mail: PeatHunter@gmail.com

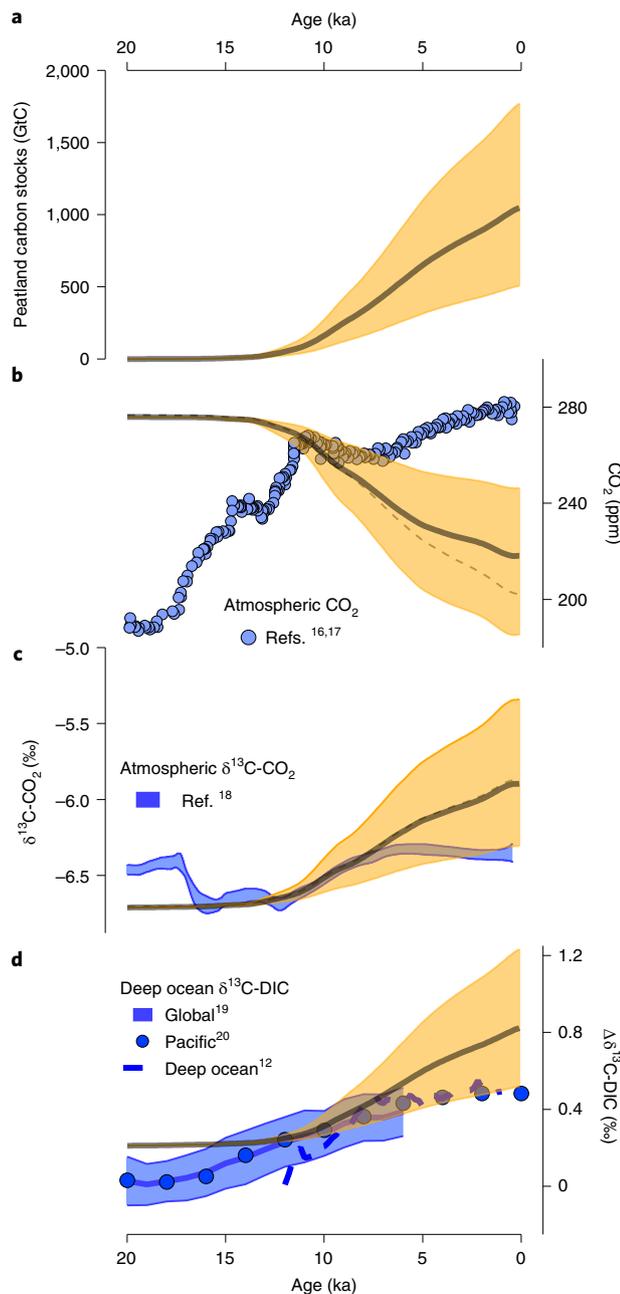


Fig. 1 | Unrealistic consequences of large peat carbon storage. **a**, Peat carbon storage change (line) over time with uncertainties (orange shading)⁷. **b**, Observed atmospheric CO₂ concentration from ice-core (circles) and box-model-calculated CO₂ concentration. The dashed line represents the outcome without ‘carbonate compensation’ mechanism in the model. **c**, Observed atmospheric δ¹³C-CO₂ from ice-core (blue shading) and model-calculated value. **d**, Observed deep ocean δ¹³C-DIC from the global ocean (blue shading), deep Pacific (circles) and a stack of 33 deep-ocean cores (dashed blue line) and model-calculated values. The δ¹³C values are plotted as anomalies relative to model results. Data from refs.^{12,16–20}.

Lack of support from global carbon budget constraints

The exceptionally large peat carbon storage⁷ is not supported by top-down constraints from the global carbon budget reconstructions. Our model simulation results show that an increase in peat carbon storage of >1,000 GtC during the Holocene would induce a decrease in atmospheric CO₂ to below 220 ppm, an increase in

atmospheric δ¹³C-CO₂ to a value more than 0.8‰ higher than the observed and a steady rise in deep ocean δ¹³C of dissolved inorganic carbon (δ¹³C-DIC) throughout the Holocene (Fig. 1).

First, our box-model calculations demonstrate that the simplified conversion of peat carbon uptake into an atmospheric signal of >600 ppm, as shown in their Fig. 2f, was erroneous due to the neglect of the compensating effect by the ocean that acts to reduce any atmospheric perturbation by up to 80% on the millennial time scale relevant here¹⁰. We assume that Nichols and Peteet instead converted their estimated terrestrial carbon stock increase by a division factor of 2.12 GtC per ppm to arrive at the claimed peat carbon uptake-related decrease in atmospheric CO₂ of >300 ppm during the Holocene. Translating the same peat carbon uptake into an atmospheric CO₂ signal with our model yielded a decrease of about 60 ppm (Fig. 1b).

Second, our simulations suggest that exceptionally large peat carbon storage is difficult to reconcile with the atmospheric and oceanic carbon budgets. Previously, the observed changes in atmospheric CO₂ concentration and in δ¹³C from ice cores have been used to partition the contributions from the land biosphere and ocean, providing a global constraint on land carbon budget during the Holocene. The measured increase in CO₂ concentration from 265 ppm 11 kyr ago (ka) to 278 ppm in 1750 CE and the small change in δ¹³C (Fig. 1b,c) were used to reconstruct the preindustrial terrestrial net carbon uptake over the Holocene epoch to be about 250 GtC (ref. 11). This total Holocene land carbon balance reflects a strong uptake in the early Holocene through the growth of boreal forests and early peat buildup—consistent with the observed early Holocene increase in atmospheric and oceanic δ¹³C values¹²—and a carbon release of 50 GtC during the late Holocene¹¹. The small decrease in land carbon storage in the past 5 kyr contrasts with the large estimated increase in peat carbon storage of ~400 GtC during the same time period as in their Fig. 2e. A compensating carbon source of 400–500 GtC with a biogenic δ¹³C signature would have to be invoked to close the budget. A detailed analysis of this budget concluded that CO₂ emissions from land use change by early agriculturalists were not sufficient to close the gap¹³. The twofold higher estimates of peat carbon storage by Nichols and Peteet⁷—compared with the one used¹³—make it even harder to reconcile the budget. This conflict is not discussed in Nichols and Peteet⁷.

Rather than balancing the carbon budget with terrestrial carbon sources, Nichols and Peteet suggest that the “most important mechanism for balancing the peatland sink” is a continued carbon release from the deep ocean by the wind-driven upwelling during the Holocene. This mechanism requires an even greater loss of carbon from the deep ocean than implied by the peatland carbon sink alone, and is not supported by observation and simulation of marine δ¹³C and carbonate ion changes. For example, an increase in Southern Ocean upwelling would further increase δ¹³C-DIC in the deep ocean¹⁴ from the already untenable increase in δ¹³C-DIC from peatland regrowth (Fig. 1d), yet δ¹³C values remained constant after 7 ka, as observed from a stack of benthic δ¹³C data from 33 deep-ocean (>3,000 m) cores around the world oceans¹² (Fig. 1d). Furthermore, the CO₂ release from the deep ocean would lead to an increase in the carbonate ion concentration and enhanced preservation of carbonates in the deep ocean, but deep-ocean cores show the opposite—a reduction in the carbonate ion and an increase in carbonate dissolution during the Holocene¹⁵.

In summary, we conclude that the evidence presented by Nichols and Peteet⁷ is not sufficient to support their claim of doubled carbon storage in northern peatlands compared with earlier estimates.

Online content

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of

author contributions and competing interests; and statements of data and code availability are available at <https://doi.org/10.1038/s41561-021-00769-2>.

Received: 26 November 2019; Accepted: 6 May 2021;
Published online: 28 June 2021

References

- Gorham, E. Northern peatlands: role in the carbon cycle and probable responses to climatic warming. *Ecol. Appl.* **1**, 182–195 (1991).
- Yu, Z., Loisel, J., Brosseau, D. P., Beilman, D. W. & Hunt, S. J. Global peatland dynamics since the Last Glacial Maximum. *Geophys. Res. Lett.* **37**, L13402 (2010).
- Loisel, J. et al. A database and synthesis of northern peatland soil properties and Holocene carbon and nitrogen accumulation. *Holocene* **24**, 1028–1042 (2014).
- Jackson, R. B. et al. The ecology of soil carbon: pools, vulnerabilities, and biotic and abiotic controls. *Annu. Rev. Ecol. Evol. Syst.* **48**, 419–445 (2017).
- Lindgren, A., Hugelius, G. & Kuhry, P. Extensive loss of past permafrost carbon but a net accumulation into present day soils. *Nature* **560**, 219–222 (2019).
- Hugelius, G. et al. Large stocks of peatland carbon and nitrogen are vulnerable to permafrost thaw. *Proc. Natl Acad. Sci. USA* **117**, 20438–20446 (2020).
- Nichols, J. E. & Peteet, D. M. Rapid expansion of northern peatlands and doubled estimate of carbon storage. *Nat. Geosci.* **12**, 917–921 (2019).
- Treat, C. C. et al. Effects of permafrost aggradation on peat properties as determined from a pan-Arctic synthesis of plant macrofossils. *J. Geophys. Res. Biogeosciences* **121**, 78–94 (2016).
- Nichols, J. E. et al. A probabilistic method of assessing carbon accumulation rate at Imnavait Creek Peatland, Arctic Long Term Ecological Research Station, Alaska. *J. Quat. Sci.* **32**, 579–586 (2017).
- Joos, D. et al. Carbon dioxide and climate impulse response functions for the computation of greenhouse gas metrics: a multi-model analysis. *Atmos. Chem. Phys.* **13**, 2793–2825 (2013).
- Elsig, J. et al. Stable isotope constraints on Holocene carbon cycle changes from an Antarctic ice core. *Nature* **461**, 507–510 (2009).
- Menviel, L. & Joos, F. Toward explaining the Holocene carbon dioxide and carbon isotope records: results from transient ocean carbon cycle-climate simulations. *Paleoceanography* **27**, PA1207 (2012).
- Stocker, B. D., Yu, Z., Massa, C. & Joos, F. Holocene peatland and ice-core data constraints on the timing and magnitude of CO₂ emissions from past land use. *Proc. Natl Acad. Sci. USA* **114**, 1492–1497 (2017).
- Tschumi, T., Joos, F., Gehlen, M. & Heinze, C. Deep ocean ventilation, carbon isotopes, marine sedimentation and the deglacial CO₂ rise. *Clim. Past* **7**, 771–800 (2011).
- Yu, J., Anderson, R. F. & Rohling, E. J. Deep ocean carbonate chemistry and glacial-interglacial atmospheric CO₂ changes. *Oceanography* **27**, 16–25 (2014).
- Monnin, E. et al. Atmospheric CO₂ concentrations over the last glacial termination. *Science* **291**, 112–114 (2001).
- Monnin, E. et al. Evidence for substantial accumulation rate variability in Antarctica during the Holocene, through synchronization of CO₂ in the Taylor Dome, Dome C and DML ice cores. *Earth Planet. Sci. Lett.* **224**, 45–54 (2004).
- Schmitt, J. et al. Carbon isotope constraints on the deglacial CO₂ rise from ice cores. *Science* **336**, 711–714 (2012).
- Peterson, C. D. & Lisiecki, L. E. Deglacial carbon cycle changes observed in a compilation of 127 benthic δ¹³C time series (20–6 ka). *Clim. Past* **14**, 1229–1252 (2018).
- Lisiecki, L. E., Raymo, M. E. & Curry, W. B. Atlantic overturning responses to Late Pleistocene climate forcings. *Nature* **456**, 85–88 (2008).

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

© The Author(s), under exclusive licence to Springer Nature Limited 2021

Methods

To illustrate the effect of such large peat carbon perturbations on the global carbon cycle we carried out a sensitivity analysis using a simple carbon-cycle box model²¹. The model considers the carbon exchange among the atmosphere, land biosphere, oceans and marine sediments. We used the ranges (median $\pm 1\sigma$) from all three scenarios (literature, combined, grid box) in Nichols and Peteet⁷ as model inputs. All scenarios essentially yielded the same solutions. Therefore, we show the results from only the 'combined' approach here. We also ran a separate sensitivity experiment by turning off the simple 'carbonate compensation' mechanism using just the median scenario.

References

21. Bauska, T. K. et al. Carbon isotopes characterize rapid changes in atmospheric carbon dioxide during the last deglaciation. *Proc. Natl Acad. Sci. USA* **113**, 3465–3470 (2016).

Author contributions

Z.Y. and F.J. designed the research, T.K.B. carried out the box-model simulation and created the figure, and all authors were involved in writing and revising the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s41561-021-00769-2>.

Correspondence and requests for materials should be addressed to Z.Y.

Peer review information Primary Handling Editor: James Super.

Reprints and permissions information is available at www.nature.com/reprints.