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RESEARCH ARTICLE

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Lowering water table reduces carbon sink strength and carbon stocks in northern peatlands

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Abstract

Peatlands at high latitudes have accumulated >400 Pg carbon (C) because saturated soil and cold temperatures suppress C decomposition. This substantial amount of C in Arctic and Boreal peatlands is potentially subject to increased decomposition if the water table (WT) decreases due to climate change, including permafrost thawrelated drying. Here, we optimize a version of the Organizing Carbon and Hydrology In Dynamic Ecosystems model (ORCHIDEE-PCH4) using site-specific observations to investigate changes in $CO₂$ and $CH₄$ fluxes as well as C stock responses to an experimentally manipulated decrease of WT at six northern peatlands. The unmanipulated control peatlands, with the WT <20 cm on average (seasonal max up to 45 cm) below the surface, currently act as C sinks in most years $(58 \pm 34 \text{ g C m}^{-2} \text{ year}^{-1})$; including $6±7g$ C-CH₄ m⁻² year⁻¹ emission). We found, however, that lowering the WT by 10cm reduced the CO₂ sink by $13±15$ g Cm^{-2} year⁻¹ and decreased CH₄ emission by 4 \pm 4 $\rm g$ CH₄ m^{−2} year^{−1}, thus accumulating less C over 100 years (0.2 \pm 0.2 $\rm kg$ Cm^{−2}).

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Yet, the reduced emission of $CH₄$, which has a larger greenhouse warming potential, resulted in a net decrease in greenhouse gas balance by 310 ± 360 gCO_{2-eq} m⁻² year⁻¹. Peatlands with the initial WT close to the soil surface were more vulnerable to C loss: Non-permafrost peatlands lost >2 kg C m−2 over 100 years when WT is lowered by 50 cm, while permafrost peatlands temporally switched from C sinks to sources. These results highlight that reductions in C storage capacity in response to drying of northern peatlands are offset in part by reduced CH_A emissions, thus slightly reducing the positive carbon climate feedbacks of peatlands under a warmer and drier future climate scenario.

KEYWORDS

carbon flux, carbon stock, drainage, high latitude, land surface model, manipulation experiment, permafrost thaw

1 | **INTRODUCTION**

The carbon (C) stock of northern peatlands is estimated to be 265– 621 Pg (Gorham, [1991](#page-14-0); Hugelius et al., [2020](#page-14-1); Treat et al., [2019](#page-17-0); Yu et al., [2010](#page-18-0)) with some estimates as high as 1055 PgC (Nichols & Peteet, [2019](#page-16-0)) despite controversy (Nichols & Peteet, [2021](#page-16-1); Ratcliffe et al., [2021](#page-17-1); Yu et al., [2021](#page-18-1)). Approximately half of northern peatlands are underlain by permafrost (Hugelius et al., [2020](#page-14-1)), contributing to a large portion of permafrost C stock of 1035 ± 150 Pg in the first 3 m depth (Hugelius et al., [2014;](#page-14-2) Schuur et al., [2015](#page-17-2)). Most of the northern peatlands C have accumulated after the last glacial maximum (Kleinen et al., [2012](#page-15-0); MacDonald et al., [2006](#page-15-1); Treat et al., [2019](#page-17-0); Yu et al., [2010](#page-18-0)), and undisturbed peatlands continue to accumulate C at present (Bridgham et al., [2006](#page-13-0); Hugelius et al., [2020](#page-14-1); Tarnocai et al., [2009](#page-17-3)). Peatlands act as C sinks because plant productivity sustains litter C input but anaerobic conditions, low pH, and poor litter quality inhibit peat decomposition. However, there is an increasing concern that this substantial C stock is becoming vulnerable to decomposition in response to drying. Drying, as a result of a lower water table (WT), exposes the upper peat horizons to aerobic conditions, which dramatically increases C decomposition rates.

In low to mid-latitudes, substantial areas of peatlands have become drier due to anthropogenic drainage by ditches for forestry, agriculture, and urbanization (Byrne et al., [2004](#page-13-1); Evans et al., [2021](#page-14-3); Wijedasa et al., [2018](#page-18-2)), in addition to climate warming and drying over natural peatlands (Swindles et al., [2019](#page-17-4)). In northern high latitudes, however, peatland drying is mostly associated with climate change through several processes. First, if evapotranspiration increases faster than precipitation, the WT is likely to decrease. Generally, increasing precipitation is expected due to an enhanced hydrologic cycle (Bintanja & Andry, [2017](#page-13-2); Bintanja & Selten, [2014](#page-13-3)) but the patterns vary spatially and temporally (Greve et al., [2014](#page-14-4)) together with more frequent extreme precipitation events (Palmer & Räisänen, [2002](#page-16-2); Shiogama et al., [2016](#page-17-5)). Furthermore, in the high latitudes, a large fraction of precipitation can be lost during

the spring melt (Douville et al., [2021](#page-14-5); Kirtman et al., [2013](#page-15-2)) and the growing season can have a negative water balance. Coupled Model Intercomparison Project (CMIP6) climate models predict shorter but stronger drought events at high latitudes by 2100 (Ukkola et al., [2020](#page-17-6)). Warmer temperatures also enhance evapotranspiration (Helbig et al., [2020](#page-14-6)), which decreases soil moisture, and possibly the WT in peatlands. Second, higher air temperatures thaw permafrost and melt extensive ground ice complexes, which subsequently increases active layer thickness and changes surface morphology. This change in surface morphology alters the spatial and vertical distribution of water in permafrost peatlands and exposes surface peat layers to aerobic decomposition in some areas (Lewis et al., [2012;](#page-15-3) Olefeldt et al., [2016](#page-16-3)). This can occur at small spatial scales, with a variation of local wet versus dry patches in polygonal tundra, or at large scales, such as thermokarst formation or thaw lake drainage, which are highly variable in space and time (Fewster et al., [2022;](#page-14-7) Jones et al., [2022](#page-14-8); Liljedahl et al., [2016](#page-15-4)).

Currently, Arctic and boreal peatlands are C sinks (Virkkala et al., [2021](#page-17-7)), and peatland drying has the potential to alter C fluxes and soil organic C (SOC) stocks (Avis et al., [2011](#page-13-4); Lara et al., [2015;](#page-15-5) Lawrence et al., [2015](#page-15-6); Vaughn et al., [2016](#page-17-8); Wainwright et al., [2015](#page-18-3)). Generally, the exposure of peat soils to oxygen in the undersaturated part of the soil profile decreases CH_A production and emissions but increases $CO₂$ emissions due to SOC decomposition (Huang et al., [2021](#page-14-9); Leifeld et al., [2019](#page-15-7); Leifeld & Menichetti, [2018](#page-15-8)), potentially contributing to a positive carbon-climate feedback (Günther et al., 2020). Huang et al. (2021) estimated $CO₂$ emissions to be increased by 1.12 mg Cm⁻² h⁻¹ cm⁻¹ and CH₄ emissions to be decreased by 0.09 mg Cm⁻² h⁻¹ cm⁻¹ due to drying in global peatlands, but the emission changes of northern peatlands were highly sensitive and variable compared to lower latitudes. In addition, temperature sensitivity of $CH₄$ emissions varies with WT and can add complexity to the CH_4 :CO₂ ratio under temperature and WT variations (Chen et al., [2021](#page-13-5)).

Despite growing research on the sensitivity of peatlands to drying, both $CO₂$ and $CH₄$ gas responses and the change of C **Blobal Change Biology**

stocks in northern circumarctic peatlands are still uncertain, es pecially for peatlands underlain by permafrost. Individual-site studies in Arctic and Boreal regions consistently showed in creased CO ² emissions from ecosystem respiration (*R*eco, [Kittler et al., [2016](#page-15-9); Martikainen et al., [1995](#page-15-10); Natali et al., [2015](#page-16-4)]) and decreased CH ⁴ emissions (Kittler et al., [2017](#page-15-11); Kwon et al., [2017](#page-15-12); Natali et al., [2015](#page-16-4); Nykänen et al., [1998](#page-16-5); Olefeldt et al., [2017](#page-16-6); Turetsky et al., [2008](#page-17-9), [2014](#page-17-10); Zona et al., [2009](#page-18-4)) in response to lower WT. However, gross primary production (GPP) showed contrasting responses, that is, reduced (Olefeldt et al., [2017](#page-16-6)) or increased GPP (Kittler et al., [2016](#page-15-9); Natali et al., [2015](#page-16-4)), depending on the plant community composition and the occurrence of shift in species due to drying. Some studies synthesized the drying effects in northern circumarctic peatlands, but they were limited to CH ⁴ (Nykänen et al., [1998](#page-16-5); Olefeldt et al., [2013](#page-16-7); Turetsky et al., [2014\)](#page-17-10) or to a small region (Laine et al., [1996](#page-15-13)). To quantify the response of both CO $_{\rm 2}$ and CH_4 fluxes and SOC stock to drying in northern peatlands, we compiled CO $_2$ and CH $_4$ fluxes from six field WT manipulation experiments from Arctic and Boreal sites, optimized key parame ters of the ORCHIDEE-PCH4 land surface model using data assim ilation to reproduce varying WT conditions, and quantified the C flux and stock change in response to drying. Specifically, using a site specifically optimized model, we first quantify the sensitivity of C fluxes of each site when WT is sequentially lowered by 5, 10, 20, and 50 cm relative to the control WT, and then evaluate the changes in C stocks when the WT is lowered for 100 years. Lastly, we compare what drives the variations in C flux sensitivity among sites.

2 | **METHOD**

2.1 | **Field drying manipulation experiment sites**

We used data from six field WT manipulation (drainage) experi ments, which are located between 62- and 71-degree North: (1) Särkkä, Finland (FI-SAR), (2) Lakkasuo, Finland (FI-LAK), (3) Healy, Alaska (US-HEA), (4) Bonanza, Alaska (US-BZF), (5) Chersky, Russia (RU-CHE), and (6) Utqiagvik (a.k.a. Barrow), Akaska (US-BES; Table [1](#page-2-0)). The specific peatland types include boreal bog, fen, and moist/wet tundra. Three sites are underlain by permafrost, while the others are not (Table [1](#page-2-0)). Peat depths range from 0.2 to 2.7 m (Table [2](#page-3-0); note that RU-CHE has peat depths of 0.2–0.4 m consid ering spatial heterogeneity, and to be exact it is not peatland by definition—peat depth should be >0.4 m). The average control WT varies from 20 cm (belowground) to −5 cm (aboveground), with the lowest seasonal WT ranging from 45 cm to −1 cm (Figure [S1\)](#page-18-5). The drainage experiments were carried out in parallel with the con trol experiments, and the drainage intensity varies by site, ranging from −17 (17 cm lower WT compared to the control) to 0 cm (no av erage difference). Flux rates were measured 0–30 years after the drying experiments started. CO_2 and CH_4 fluxes were measured using eddy covariance method and/or chambers. More details on

Site characteristics of northern peatlands with field drying manipulation experiments used for the model calibration **TABLE 1** Site characteristics of northern peatlands with field drying manipulation experiments used for the model calibration \overline{a} 쁰 $\overline{\mathbf{r}}$ $\overline{}$

TABLE 2 Initialization parameters for each site for fitting soil temperature and soil organic carbon (C) content

Note: The number of dry peat layers (and corresponding thickness in cm) was included to increase goodness of fit between simulated and observed soil temperature, which does not influence the actual WT or soil moisture. Simulated peat depth and SOC are from the corresponding observation years for comparison to the observations, and may differ over the simulation years. The amount of SOC was simulated by adjusting the number of spin-up years, that is, SOC accumulation years.

site characteristics and experimental setups are found in the references in Table [1](#page-2-0).

2.2 | **Model description**

We used a modified version of the land surface model ORCHIDEE-PEAT that was developed to simulate northern peatlands (Qiu et al., [2018](#page-16-8)). This version is called ORCHIDEE-PCH4, and simulates hydrology, surface energy, and C cycle processes resulting in $CO₂$ and CH₄ fluxes in northern peatlands using the parameteriza-tions from Qiu et al. ([2018](#page-16-8)) and a $CH₄$ module (Salmon et al., [2021](#page-17-11)). Photosynthesis and plant respiration are simulated for peatland vegetation type (represented as C3 graminoids) as described in section 2.2.1 in Qiu et al. ([2018](#page-16-8)). After senescence, litterfall goes into two litter pools (metabolic and structural) and three soil C pools (active, slow, and passive) after a series of decomposition processes following the CENTURY scheme (Krinner et al., [2005](#page-15-14); Parton et al., [1987](#page-16-9); Paustian et al., [1992](#page-16-10)). The decomposition rates of active, slow, and passive soil C pools, including the transfer among soil pools are 1.0, 0.027, and 0.0006 year−1 at 30°C, and the actual rates are simulated considering soil temperature, moisture, and depth (Qiu et al., [2019](#page-16-11)). Peat C from three pools is decomposed to $CO₂$ and CH_A , with the decomposition rates affected by soil temperature and moisture, peat depth, and additionally oxygen concentration in soil pores in the case of $CH₄$ (Qiu et al., [2019](#page-16-11); Salmon et al., [2021](#page-17-11)). Oxygen is diffused between the atmosphere and the top soil layer, or through snow layers when existing, as well as between soil layers, and is also provided up to the rooting depths by plant roots (Salmon et al., [2021](#page-17-11)). Oxygen in soils oxidizes CH_A , and the oxidation rate is determined by the turnover time of CH₄ (Salmon et al., [2021](#page-17-11)). The residual CH₄ after oxidation in each layer is then emitted to the atmosphere through diffusion, ebullition, and plant-mediated transport. Diffusion occurs between soil layers as well as between top soil layer and the atmosphere based on the concentration gradients, soil moisture, and soil pore size. Methane bubbles form due to hydrostatic pressure in soil, and the probability of these bubbles to reach the atmosphere is simulated as ebullition. The amount of plant-mediated transport is

influenced by the gas transport efficiency (representing aerenchyma density) and seasonal plant productivity. More details on these processes are described in section 2.1 in Salmon et al. ([2021](#page-17-11)). Peat growth (accumulation of peat C; C input into soil minus $CO₂$ & CH₄ production/decomposition) is simulated by transferring excessive peat C of one layer to the one below within the 32 discretized soil layers (section 2.1 in Qiu et al. ([2019](#page-16-11))).

2.3 | **Model setup and parameter optimization**

The ORCHIDEE model allows multiple vegetation types in one grid cell, with distinct soil tiles to compute the hydrology of peat, herbaceous, and woody vegetation types. The fraction of the grid cell occupied by peatland receives runoff from the other nonpeat vegetation fractions to maintain a high WT (Qiu et al., [2018](#page-16-8)), and the prognostic WT in peatland depends on the fraction size of non-peatland vegetation. Here, we do not use the prognostic WT from the model but set the entire grid cell to be covered by peatlands. Then, we prescribed soil moisture according to the observed daily WT (the resolution can be 30 min), and set soil moisture to 0.80 (80% of soil porosity filled with water and 20% with air) for the soil layers below WT and to 0.50 for the soil layers above WT. These values are the averages soil moisture observed in US-HEA and RU-CHE sites, where volumetric soil moisture was measured.

To more accurately simulate soil temperature, which is one of the critical drivers for CO_{2} - and CH_{4} -related processes and their rates, we used the thermal properties (heat capacity of 2.5 $\cdot10^6$ JK $^{-1}$ m $^{-3}$ and thermal conductivity of 0.05 $Wm^{-1}K^{-1}$ [dry] and 0.25 $Wm^{-1}K^{-1}$ [solid]) of 100% organic soil in the model. The apparent heat capacity and thermal conductivity were calculated considering the water and ice content in soil (Guimberteau et al., [2018](#page-14-13)). Plus, when simulated soil temperature was higher than observations, we overwrote the thermal properties of dry peat soils at the top soil layers within the soil thermic sub-module to mimic the insulating function of an overlying moss or organic soil layer to accurately simulate observed soil temperature profiles (Gornall et al., [2007](#page-14-14); Soudzilovskaia et al., [2013](#page-17-12)) as in (Ekici et al., [2014](#page-14-15)). These dry peat layers were decoupled from the soil hydrology module and only used for simulating soil temperature profiles. The number of dry peat layers was optimized for each site by minimizing the root mean squared error (RMSE) and Nash-Sutcliffe model efficiency coefficient (MEF) from the observed soil temperatures (shown in Table [2](#page-3-0) and Figure [S2](#page-18-5)).

We calibrated the key parameters that are associated with $CO₂$ and $CH₄$ fluxes to simulate site-specific fluxes of the control treatment (Table [3](#page-4-0)). Thus, the differences in C fluxes between control and dry treatments are driven by the prescribed WT of each treatment and model parameters that change by soil moisture. For GPP, the Vc_{max} at 25°C (maximum carboxylation rate in photosynthesis at 25°C; default 40 μmol m^{−2} s^{−1} [Qiu et al., [2018](#page-16-8)]) was calibrated. In default setting, GPP decreases with decreasing soil moisture availability. However, some plants can photosynthesize more actively under dry condition (Sulman et al., [2010](#page-17-13)). In addition, some sites (e.g., RU-CHE) showed significant changes in vegetation communities due to drying, with greater shrub abundance (Kwon et al., [2016](#page-15-15)) and GPP (Kittler et al., [2016](#page-15-9)). Increasing shrub abundance is often observed in drying northern peatlands unless taller trees outcompete them for light (Harris et al., [2020](#page-14-16); Laiho et al., [2003](#page-15-16); Murphy et al., [2009](#page-16-12)). To test the effects of increased photosynthetic rates on flux and stock change with or without composition changes, we additionally ran simulations with the increased Vc_{max} at 25°C by 10% to mimic increased plant productivity. The phenology was selected between C_3 grasses and C_3 grasses + mosses by minimizing errors (RMSE and MEF) between simulations and observations. Phenology determines the onset time of photosynthesis, with the earlier onset when mosses are present, as they can photosynthesize at lower temperature and lower light level than C_3 grasses (Atanasiu, [1971](#page-13-6)). The growing degree days (GDD) threshold for the onset of C_3 grass photosynthesis is calculated using

GDD threshold =
$$
320 + 6.25 \cdot \text{temp} + 3.125 \cdot 10^{-2} \cdot \text{temp}^2
$$
 (1)

where temp is the average air temperature of the past 3 years. The coefficients of Equation ([1](#page-4-1)) are calibrated globally for different vegetation types (Botta et al., [2000](#page-13-7); Krinner et al., [2005](#page-15-14)). Earlier onset of photosynthesis for C_3 grasses + mosses is achieved by reducing the threshold of GDD using

GDD threshold =
$$
\frac{1.93 \cdot 10^5}{\left(1 + e^{\left(-8,13 \cdot 10^{-2} \cdot (\text{temp}-87.87)\right)}\right)}
$$
(2)

which is calibrated by GPP observations in 19 northern peatlands (Qiu et al., [2018](#page-16-8)). Different from the default setting of respiration (CO₂ production) cutoff below −1°C, we allowed a continuous CO₂ respiration at sub-zero temperatures, with the temperature control on respiration using.

Temperature control =
$$
e^{(0.69 * \frac{\text{soil temperature}-30}{10})}
$$
, max = 1 (3)

as non-growing season respiration can be considerable (Natali et al., [2019](#page-16-13)).

The amount of plant-mediated CH_A transport was calculated after subtracting the fixed fraction of methanotrophy at roots (mrox in the equation 9 in Salmon et al. ([2021](#page-17-11))) in the original model. Instead, we excluded the methanotrophy (mrox) term in the plant-mediated $CH₄$ transport equation but used mrox as the amount of oxygen provided into soil through roots, and let it oxidize $CH₄$ during the methanotrophy process in soils, similar to Morel et al. ([2019](#page-16-14)). This allowed the amount of plant-mediated CH_A transport decoupled from methanotrophy in soils. Then, six parameters were optimized targeting the best fit to the observed $CH₄$ emissions of control treatment of each site using the ORCHIDEE Data Assimilation Systems (ORCHIDAS; Bastrikov et al. ([2018](#page-13-8)); <https://orchidas.lsce.ipsl.fr/>). Daily CH₄ emissions of growing season were linearly interpolated for days without observations, and the earliest (latest) observations of each year were used for CH_4 emissions before (after) the first (last) observation of that year. Within the ORCHIDAS framework, we used the genetic algorithm (Goldberg & Holland, [1988](#page-14-17); Haupt & Haupt, [2004\)](#page-14-18) to find the best set of parameters within the defined boundary of each parameter. This stochastic algorithm is a global random search method based on the principles of genetics and natural selection, and was found in Bastrikov et al. ([2018](#page-13-8)) to outperform traditional

TABLE 3 Parameter sets for each site for fitting CO₂ and CH₄ fluxes: *V*_{Cmax} at 25°C (maximum carboxylation rate in photosynthesis at 25°C), *k* (methanogenesis rate relative to the oxic decomposition), k_{MT} (turnover time of methanotrophy), tveg (the amount of CH₄ transported through aerenchymatous plants), mrox (the amount of oxygen provided into soil through roots), mxr (the mixing ratio of CH₄ in bubbles in soils), and wsize (the extent of the connected network of water-filled pores)

Site	$V_{\rm C_{max}}$ $(\mu \text{mol m}^{-2} \text{ s}^{-1})$	Phenology	k (ratio to oxic decomposition)	k_{MT} (s)	tveg	mrox	mxr (fraction)	wsize (m)
FI-SAR	40	$C_3 + \text{moss}$	4.92	129.047	0.10	1.30	0.05	0.49
FI-LAK	40	$C_2 + \text{moss}$	4.56	86,400	0.25	1.03	0.07	0.48
US-HEA	40	C_3	9.82	153.127	0.41	0.39	0.52	0.24
US-BZF	45	C_3	6.65	86,400	1.30	0.5	0.20	0.38
RU-CHE	40	C_3	1.02	97,695	2.23	1.07	0.11	0.41
US-BES	40	$C_3 + \text{moss}$	2.00	161,472	3.71	0.70	0.39	0.05

Note: The parameters related to CH₄ processes are optimized using the ORCHIDAS.

gradient-based approaches when optimizing key ORCHIDEE model parameters. Calibrated parameters include *k* (methanogenesis rate relative to the oxic decomposition; for example, *k* of 2 represents the CO₂ and CH₄ maximum production ratio of 2:1; ranging from 1 to 10; Khvorostyanov et al., [2008](#page-15-17); Wania et al., [2010](#page-18-7)), k_{MT} (turnover time of methanotrophy; ranging from 1 to 5 days; Juncher Jørgensen et al., [2014;](#page-14-19) Khvorostyanov et al., [2008](#page-15-17); Morel et al., [2019](#page-16-14)), tveg (the amount of $CH₄$ transported through aerenchymatous plants; ranging from 0 to 30; Walter & Heimann, [2000](#page-18-8)), mrox (the amount of oxygen provided into soil through roots, which can oxidize CH_4 in soils; ranging from 0 to 5, Walter & Heimann, [2000](#page-18-8)), mxr (the mixing ratio of $CH₄$ in bubbles in soils; ranging from 0.05 to 0.53; Baird et al., [2004](#page-13-9); Morel et al., [2019](#page-16-14); Riley et al., [2011](#page-17-14)), and wsize (the extent of the connected network of water-filled pores that affects the probability of $CH₄$ bubbles to reach the soil surface; ranging from 0.001 to 0.5; Khvorostyanov et al., [2008](#page-15-17)) are summarized in Table [3.](#page-4-0) The range of each parameter was defined and modified following Salmon et al. ([2021](#page-17-11)), which evaluated $CH₄$ fluxes across 14 northern peatlands. Optimized parameter sets were used during the experimental simulations (at the start of the drying experiment; described in the next section). The first 20 years of simulations were excluded, which may include transitional status due to changed parameters.

2.4 | **Model simulations of each site**

The simulation was forced by the measured meteorology of random years at each site on a 30-min time-step. When meteorological measurements were not available, they were substituted by CRUNCEP v8 for the corresponding grid cell for the site-specific observation years ([https://vesg.ipsl.upmc.fr/thredds/catalog/work/p529viov/](https://vesg.ipsl.upmc.fr/thredds/catalog/work/p529viov/cruncep/V8_1901_2016/catalog.html) [cruncep/V8_1901_2016/catalog.html\)](https://vesg.ipsl.upmc.fr/thredds/catalog/work/p529viov/cruncep/V8_1901_2016/catalog.html). The model simulations were performed as follows: (1) the full model was run for 50 years to obtain steady state daily values for soil C inputs, (2) a soil accumulating spin-up sub-model was run for a site-specific time period to match simulated SOC content with observed SOC (Table [2](#page-3-0)), and (3) another 50 years of the full model run was conducted to achieve the equilibrium of physical variables. During the spin-up, atmospheric $(CO₂)$ was set to the pre-industrial level of 286 ppm, and (4) transient simulations were carried out with the observed rising atmospheric $[CO₂]$ for 100 years until the starting year of drying experiment. During the simulations, WT was prescribed with the site-specific daily WT of the control treatment.

Afterwards, another 100 years of experimental simulations were conducted at each site, using (a) the WT of the control sites (Figure [S1\)](#page-18-5), (b) the WT of the dried sites (Figure [S1\)](#page-18-5), and (c) the WT sequentially lowered by 5, 10, 20, and 50 cm relative to the WT of the control sites. Sequential drying was applied until the WT did not exceed the maximum thaw depth for permafrost-affected peatlands, which was equivalent to a WT depth 20 cm lower than controls, and until the WT up to 50 cm belowground for non-permafrost peatlands because hydrological self-regulation of peatlands may not sustain low WT in the long term (Dise, [2009](#page-13-10); Waddington et al., [2015](#page-18-9)).

 KWON et al. **[|] 6757**

When WT and fluxes were measured at multiple locations within the study site (chamber-based method), we used the average WT for one representative simulation of that site. In addition, we excluded sub-plots, which experienced significant tree growth following drying (two sub-plots were excluded out of six in the case of FI-LAK).

2.5 | **Model-observation comparison and statistical analysis**

Model simulation errors against observations were estimated: RMSE, and the squared of its two partitioned components, squared difference of standard deviations (SDSD) and the lack of positive correlation weighted by the standard deviation (LCS) following Kobayashi and Salam ([2000](#page-15-18)). They measure the magnitudes, for example, maximum and minimum seasonal flux rates, and patterns of fluctuations, for example, seasonality, respectively. We also estimated MEF for each site and treatment. When the model simulation is fully matching with the observation, RMSE, SDSD, and LCS are 0 and MEF is 1.

To compare the flux sensitivity to drying, we fitted exponential equations for NEE (net ecosystem CO₂ exchange; R_{e^ce} – GPP) and $CH₄$ for each site with the sequential drying.

$$
NEE or CH_4 = a * e^{(b*WT)} \tag{4}
$$

using the nls fit function in R (R Development Core Team, [2013](#page-17-15)).

2.6 **|** Combined effects of CO₂ and CH₄

Using the new global warming potential metrics (GWP*; Allen et al., [2018](#page-13-11); Cain et al., [2019](#page-13-12); Lynch et al., [2020](#page-15-19)), we estimated the combined effects of $CO₂$ and $CH₄$ flux changes. Different from the conventional GWP, GWP* considers the removal of short-lived gases over time, for example, the removal of CH_A from the atmosphere after the residence time of 12 years. Thus, its application can be especially beneficial when the CH_A emission is stable or decreasing over time.

3 | **RESULTS**

3.1 | **Comparison between observations and simulations**

The model simulated soil temperature profiles well (Figure [S2\)](#page-18-5). Although peat pore $CO₂$ and $CH₄$ concentrations can vary with WT and site-specific characteristics, simulated concentrations (ca. $3-40$ g CH₄ m⁻²) were comparable with those of the previous studies (Saarnio et al., [1997](#page-17-16); Waddington et al., [2009](#page-17-17)). Together with the prescribed WT, reasonable soil temperature and pore gas concentrations, and calibration of flux-related parameters, model simulations and observations of daily $CO₂$ and $CH₄$ fluxes agreed well

FIGURE 1 Comparison of NEE (a, e), CH₄ (b, f), R_{eco} (c, g), and GPP (d, h) between the model simulations (X-axis) and the observations (Yaxis) for control (a–d) and dry (e–h) treatments after the calibration for each site. Lines are drawn for 1:1 comparisons (solid) and regressions (dashed). Asterisks next to the site name indicate non-permafrost peatlands.

for both NEE (average RMSE: 1.04^g C–CO₂ m⁻² day⁻¹ and MEF: −0.36, Table [S1](#page-18-5) and Figures [1, 2](#page-6-0) and Figure [S3](#page-18-5)) and CH₄ (average RMSE: $0.03g$ CH₄ m⁻² day⁻¹ and MEF: -8.27, Table [S1](#page-18-5); Figures [1, 2](#page-6-0) and Figure [S3\)](#page-18-5) for control and drying treatments (for NEE, average RMSE: 1.20 $g \text{ C-CO}_2$ m⁻² day⁻¹ and MEF: -2.05; for CH₄, average

RMSE: 0.03 g CH₄ m⁻² day⁻¹ and MEF: -2.84, Table [S1](#page-18-5); Figures [1, 2](#page-6-0) and Figure [S3\)](#page-18-5). The model was calibrated using observations from control treatments, and similar model performance for the drying treatment implies that the model can capture the response of C flux to WT variations well (Tables [S1](#page-18-5) and [S2\)](#page-18-5). Fluxes of some sites

FIGURE 2 Taylor diagrams of daily NEE (a), CH₄ (b), R_{eco} (c), and GPP (d) using model simulations and observations for control and dry treatments. Asterisks next to the site name indicate non-permafrost peatlands.

FIGURE 3 The response of NEE (a) and CH₄ (b) fluxes to sequential drying by 5, 10, 20, and 50cm (filled circles) starting from the initial WT of control treatment (unfilled circles; highest WT of each site). Points and error bars represent the averages and standard deviations of annual WT and C fluxes for 100-year simulations of each site (color). The exponential lines were fitted for NEE and CH $_A$. Positive values of WT denote WT above the soil surface. Asterisks next to the site name indicate non-permafrost peatlands.

showed opposite differences between the simulated and observed dynamics. For example, in the case of US-BZF, drying reduced $CH₄$ emissions in simulations but the observed data showed increases in

 $CH₄$ emissions. These differences were related to measurements being made at multiple plots with high heterogeneity. Across most sites and fluxes model errors were due to seasonal patterns rather

than magnitude, as inferred from larger LCS than SDSD (Table [S1](#page-18-5); Figures [1, 2](#page-6-0) and Figure [S3](#page-18-5)). The drying treatment (deeper WT) of RU-CHE showed that errors from magnitude were larger than those from the seasonality for $CO₂$ fluxes, where drying decreased GPP in simulation but increased GPP in observations. With a good agreement between simulations and observations, we described the results based on the simulations from here.

3.2 | **Response of C fluxes to drying**

With the initial WT of the control treatment (simulation during the observation years), NEE ranged from −133 g C–CO₂ m⁻² year⁻¹ (US-BZF) to $-21g$ C–CO₂ m⁻² year⁻¹ (US-BES; Figure [5a](#page-10-0)). Across all sites drying reduced the net $CO₂$ uptake (i.e., less negative NEE) by $16g$ C–CO₂ m⁻² year⁻¹ on average, with the smallest uptake decrease in US-HEA by 1g C–CO₂ m⁻² year⁻¹ (with the smallest drying intensity) and the largest uptake decrease in RU-CHE by 50 g C-CO₂ m^{-2} year⁻¹ (with the largest drying intensity). The change in NEE was driven primarily by reduced GPP (11 g C–CO₂ m⁻² year⁻¹ on average) and increased $R_{\text{e}c}$ (5g C–CO₂ m⁻² year⁻¹ on average). The annual CH₄ emissions were largest at US-BZF (19^g C–CH₄ m⁻² year⁻¹) and smallest at US-HEA (0.1 g C–CH₄ m⁻² year⁻¹) for the control treat-ment (Figure [5b](#page-10-0)). CH₄ emissions decreased by 2^g C–CH₄ m⁻² year⁻¹ on average due to the drying treatment, with the smallest decrease in US-HEA by 0.003 g C–CH₄ m⁻² year⁻¹ and the largest decrease in RU-CHE by 5^g C–CH₄ m⁻² year⁻¹.

The drying intensity (change in WT) and the number of years since the start of the drainage differed among sites. To analyze the C flux sensitivity to drying by site, we compared C flux responses to sequential decrease of the WT by 5, 10, 20, and 50 cm compared to the control simulation. As WT draws down, net $CO₂$ uptake decreased (Figure [3a](#page-7-0)). This decreased uptake was driven both by re-duced GPP and increased R_{eco} (Figure [S4\)](#page-18-5). The response of NEE to a change of the WT was large when the initial WT was close to the soil surface, and insignificant and almost linear when the initial WT was below 10cm (Figure [3a](#page-7-0) and Figure [S4\)](#page-18-5). The average decrease in the CO₂ sink was $13±15$ g C–CO₂ m⁻² year⁻¹ with 10 cm decrease in WT, ranging from 0 to 36 g C–CO₂ m⁻² year⁻¹. A similar response was found for CH_A emissions. They decreased sharply when the initial WT was close to zero cm but showed negligible changes when the initial WT was below 10cm (Figure [3b](#page-7-0) and Figure [S4](#page-18-5)). The average decrease in the CH₄ emissions due to 10cm lower WT cm was $4\pm 4g$ CH₄ m⁻² year⁻¹ with the range between 0 and 9^g CH₄ m⁻² year⁻¹. These responses were mostly driven by reduced $CH₄$ production and CH₄ transport by plants, while CH₄ oxidation was decreased (US-HEA, RU-CHE, and US-BES), increased (FI-SAR and FI-LAK), or unchanged (US-BZF) with lowered WT (Figure [S5](#page-18-5)).

Permafrost peatlands showed smaller net $CO₂$ uptake and lower $CH₄$ emissions compared to the non-permafrost peatlands (Figure [3](#page-7-0); note that * indicates non-permafrost sites), which was also represented by larger and smaller coefficient values of 'a' of the exponential fits for NEE and $CH₄$, respectively (Figure [S6](#page-18-5)). Despite smaller

net $CO₂$ uptake due to drying, the average NEE remained negative during the 100 years of simulation, indicating the persistence of peatland C sinks, even under drier conditions. Two permafrost underlain peatlands (RU-CHE and US-BES), which show low net $CO₂$ uptake due to the existence of permafrost and large response to drying due to high initial WT, showed net $CO₂$ emission in some years (Figure [3a](#page-7-0)), indicating reduced C sinks and the potential switch to C sources.

3.3 | **Carbon stock changes due to drying**

Although CH₄ emissions decreased in response to drying, larger decreases in net $CO₂$ uptake (less negative NEE) resulted in lower soil C stocks compared to the control (Figure [4a,b\)](#page-9-0). When subjected to a lowered WT by 10 cm, peatlands still accumulate C but the rate of accumulation is lower by 0.2 ± 0.2 kg C m−2 over 100 years. We found that most of this reduction in C accumulation came from the active (labile) C pool (Figure [4d\)](#page-9-0). Larger decreases in the active C pool and C stock were observed in US-BZF and RU-CHE, where the initial WT is close to 0 and the NEE response to drying was stronger than at other sites (Figure [4a,b](#page-9-0)). The most significant reduction in C stock (over 2 kg C m−2) was observed at US-BZF with 50 cm lower WT, where permafrost does not exist (Figure [4a](#page-9-0)). Increasing photosynthetic uptake due to drying (increasing Vc_{max25} by 10%) reduced the differences in GPP between simulation and observation at US-BZF and RU-CHE, where drying increased GPP (Table [S2\)](#page-18-5). Because of the larger C input to soil, increased GPP increased C stock compared to the control treatment in most sites (Figure [4c\)](#page-9-0), out-compensating the C loss by drying. However, it was not the case for US-BZF when dried by >10 cm, that additional C input due to increased productivity was not enough to offset enhanced R_h.

3.4 | **Net GHG balance change**

When comparing C fluxes between control and a lower WT by 10 cm over 100 years, drying reduced net CO₂ uptake (Figure [5a,d\)](#page-10-0), but decreased $CH₄$ emissions (Figure [5b,e\)](#page-10-0). These contrasting effects are combined using GWP*, with the following patterns observed across the study sites: control treatments of US-BZF and RU-CHE acted as net CO_{2-eq} sources, FI-SAR, US-HEA, and US-BES as CO_{2-eq} sinks, and FI-LAK switched from a $CO_{2\textrm{-}ea}$ source to sink after 50 years (Figure [5c](#page-10-0)). Lowering the WT by 10cm reduced $CO_{2\text{-eq}}$ emissions by 310 \pm 360g CO_{2-eq} m⁻² year⁻¹ because reduced CH₄ emissions played a larger role than the reduced $CO₂$ uptake (Figure [5f\)](#page-10-0). It is noteworthy that when the initial $CH₄$ emission rates were high, as in the case of US-BZF, the peatland still acted as a net CO_{2-eq} source after drying (Figure [5c](#page-10-0)). Peatlands that had the initial WT close to the soil surface showed larger responses than those that experienced a lower WT (Figure [5](#page-10-0)). Similar trends were shown for drying by 5cm and 20 cm , but we observed smaller reductions in CO_{2-en} emissions with smaller drying intensity (Figure [S7\)](#page-18-5).

FIGURE 4 Carbon stock change due to decreasing WT by 5, 10, 20, and 50 cm compared to the control treatment. Total C change (a), total C change in % (b), total C change when *V*c_{max25} is increased by 10% (c), and C change by pool (d). Negative (positive) values represent C loss (gain) or smaller (larger) C accumulation compared to the control treatment. Positive values of WT denote WT above the soil surface. Asterisks next to the site name indicate non-permafrost peatlands.

4 | **DISCUSSION**

4.1 | Responses of CO₂ and CH₄ fluxes to drying

Carbon sequestration and storage is one of the vital climate regulating services provided by peatlands; however, the C storage capacity of peatlands may be greatly altered under changing climatic conditions. We showed that drying of peatlands by lowering WT resulted in less C sequestration. First, lowered WT decreased C input into soil, due to reduced photosynthetic activity (i.e., GPP) under decreased water availability in shallow soil layers, where most roots exist. This response can be, however, different depending upon plant species. For example, vascular plants show higher productivity with lowered WT, while mosses show lower productivity (Sulman et al., [2010](#page-17-13)). In addition to the instant responses of GPP to WT variations, plant composition can shift (Breeuwer et al., [2009](#page-13-13); Potvin et al., [2015](#page-16-15)) and GPP may decrease (Churchill et al., [2014;](#page-13-14) McPartland et al., [2019](#page-16-16)) or increase (Kittler et al., [2016](#page-15-9)) when WT in peatlands persistently lowers. Increasing productivity due to drying can compensate the C loss, but may not be enough under intense drying as shown in Figure [4c](#page-9-0).

In addition to reduced photosynthesis, peatland drying generally increased peat C loss through heterotrophic respiration (R_h) despite decreased autotrophic respiration (R_a) offsetting this change. Respiration rates are higher under aerobic conditions compared to water-saturated conditions (Moyano et al., [2013](#page-16-17)), and this mechanistic relationship is well represented in the model. Although drying did not significantly affect soil temperatures in the simulation, drying reduced deep soil temperature at US-HEA and RU-CHE in the observations because of insulation effects of drier peat at the surface (Kwon et al., [2019](#page-15-20)). This implies that the respiration response of deep soil layers to drying can be limited as compared to the surface soil layers. The combined effects of drying on GPP and R_{eco} , thus the direction of C gain or loss, are largely dependent on the response of the plant productivity to drying, as shown in the reversed C stock change with the increased productivity in most sites (Figure [4a,c](#page-9-0)). The initial WT also plays a role that higher initial WT with a large proportion of labile C can show a stronger R_h response to drying, driving the net CO_2 response toward a larger loss.

In contrast to the reduced net $CO₂$ uptake following peatland drying, $CH₄$ emissions decreased with lowered WT with similar decreasing rates to those of the previous studies (Evans et al., [2021](#page-14-3); Huang et al., [2021](#page-14-9); Kuhn et al., [2021](#page-15-21); Nykänen et al., [1998](#page-16-5); Olefeldt et al., [2013](#page-16-7)). This exponential decrease with lower WT (Figure [3b](#page-7-0)) can be attributed to thinner anaerobic peat layers at deep soil layers and thicker aerobic peat layers at the surface (Kuhn et al., [2021](#page-15-21); Kwon et al., [2017](#page-15-12); Olefeldt et al., [2013](#page-16-7)). Although atmospheric $CH₄$ can be oxidized in top soils by highaffinity methanotrophs, thereby reducing net CH_A emission to the atmosphere (Oh et al., [2020](#page-16-18)), the response of $CH₄$ emissions

FIGURE 5 Cumulative CO₂ (a), CH₄ (b), and combined (c) flux change over 100 years for control (solid) and dry by 10cm (dashed) treatments. Positive (negative) values represent net emission (uptake) to (from) the atmosphere (a–c). The differences between control and dry by 10 $\rm cm$ treatments are depicted in d–f, with positive values representing increased CO_{2-eq} emissions or decreased CO_{2-eq} uptake compared to the control treatment. Asterisks next to the site name indicate non-permafrost peatlands.

to WT was largely driven by $CH₄$ production because most $CH₄$ is oxidized by low-affinity methanotrophs when abundant $CH₄$ exists in peat soils (Kwon et al., [2017](#page-15-12), [2021](#page-15-22)). The initial WT, thus, largely affected CH_A emissions through the rate of methanogenesis (aerobic respiration to methanogenesis ratio; *k*) and methanotrophy (k_{MT} ; turnover time of methanotrophy; Figure [S8\)](#page-18-5). For example, a low ratio of aerobic respiration to methanogenesis (smaller k; higher CH₄ production potential) in RU-CHE and US-BES compared to other sites can be attributed to higher average WT and smaller temporal variations (Figure [S1](#page-18-5)), which kept the soil more anaerobic than other sites. The variations in these optimized parameters among sites may represent other environmental status that the model does not include, such as substrate status (Chang et al., [2020](#page-13-15); Roy Chowdhury et al., [2021](#page-17-18)).

4.2 | **Variations in sensitivity to drying**

The fluxes of $CO₂$ and CH₄ were sensitive to drying when the initial WT was close to 0. Unless there had been an abrupt hydrological change before the observations, the current WT can be represented as the long-term WT. Water-saturated peatlands, which are less exposed to aerobic conditions, have a large fraction of partially decomposed C in the topsoil compared to peatlands with lower WT, the topsoil of which has already been largely decomposed. Thus, a large fraction of remaining labile C can be rapidly decomposed to $CO₂$ instead of $CH₄$ when WT lowers, thus showing a stronger drying response. Non-permafrost peatland with high initial WT in US-BZF showed the strongest response to drying up to 50 cm, with >2 kg C m−2 less C accumulating over 100 years compared to the initial WT condition. In contrast, peatlands with low initial WT may show no or subtle responses to drying due to less available labile C in deeper soil layers (Huang et al., [2021](#page-14-9); Muhr et al., [2011](#page-16-19)). Larger temperature fluctuations in shallow layers compared to deeper layers may contribute to these stronger responses (Kwon et al., [2017](#page-15-12)). Fluctuations in WT can also affect the magnitude of the responses: Although the largest response was observed in US-BZF because of the combination of high WT and a large C stock, temporal lowering of WT resulted in a smaller slope change than in RU-CHE (Figure [3](#page-7-0) and Figure [S6](#page-18-5)), where WT was constantly high.

Permafrost underlain peatlands showed lower $CO₂$ and $CH₄$ fluxes compared to the other sites, as shown in previous studies (Kuhn et al., [2021](#page-15-21); Olefeldt et al., [2013](#page-16-7); Treat et al., [2018](#page-17-19)). This is mainly because permafrost limits the plant rooting depth and productivity, subsequently leading to reduced C input to soils, and reduced decomposition. However, the responses to drying in terms of direction and magnitude were similar between permafrost and nonpermafrost peatlands. Low absolute C fluxes in permafrost peatlands affected the variations, leading to almost no response in C fluxes and stock to drying in US-HEA, where initial WT is low and the existence of permafrost retards deep soil processes. Furthermore, low absolute CO₂ fluxes in RU-CHE and US-BES resulted in a temporal switch from net CO_2 uptake to net CO_2 emission in response to drying.

The simulated exponential decrease in CH_A fluxes as a function of WT was very similar to previous observations in natural (Huang et al., [2021](#page-14-9); Kuhn et al., [2021](#page-15-21); Nykänen et al., [1998](#page-16-5); Olefeldt et al., [2013](#page-16-7)) and managed peatlands (Evans et al., [2021](#page-14-3)). However, our simulations showed exponentially decreasing NEE to near-neutral NEE with lowered WT, in contrast to the linear relationship between WT and NEE in the managed peatlands of Evans et al. ([2021](#page-14-3)). The difference can be attributed to the management (harvest): Fens and bogs without harvest hardly showed the net $CO₂$ emissions with low WT, but grasslands and croplands showed net $CO₂$ emissions with low WT possibly because of different biogeochemical processes from natural peatlands, for example, disturbance and nutrient status (Evans et al., [2021](#page-14-3)). Nevertheless, we can further investigate whether natural peatlands can act as the net $CO₂$ source with extremely low WT by including more sites and more years with climate variability (Fenner & Freeman, [2011](#page-14-20); Qiu et al., [2022](#page-16-20)).

4.3 | **C stock change and warming effects**

Stronger decreases in the net $CO₂$ uptake than decreases in $CH₄$ emission resulted in smaller C accumulation in dry peatlands. Also, water-saturated peatlands were more vulnerable to C loss in response to drying compared to peatlands with initially lower WT. Furthermore, a significant C loss was observed in non-permafrost peatlands and a possible switch from C sink to source in permafrost peatlands. It is challenging to detect the change in SOC in field manipulation experiments since the soil C pool changes slowly, but we can infer from a large accumulation of C in peatlands that wet

 KWON et al. **[|] 6763**

conditions are favorable for slower turnover and higher C accumulation in soils (Hugelius et al., [2020](#page-14-1); MacDonald et al., [2006](#page-15-1)). Whether these C sinks become C sources or not is highly uncertain, but it is likely that C sink capacity decreases or even turns into a C source (Hugelius et al., [2020](#page-14-1)), combined with higher risk of peat fire may make these C stocks more vulnerable to future warming (Turetsky et al., [2011](#page-17-20); Witze, [2020](#page-18-10)). Günther et al. ([2020](#page-14-10)) showed that peatland drying has a net warming effect due to larger increases in longlived CO₂ emission than decreases in short-lived CH₄ emission, which contrasts with this present study. Günther et al. ([2020](#page-14-10)) used emission factors to estimate warming effects, with much larger $CO₂$ emission rates than $CH₄$ in temperate and tropical zones compared to the smaller warming effects reported here for the drying response of boreal peatlands. As both $CO₂$ and $CH₄$ fluxes are responding non-linearly to a lower WT and the relative changes between $CO₂$ and CH_A fluxes differ by site combined with temperature sensitivity (Chen et al., [2021](#page-13-5)), more thorough analyses are needed to quantify the compound effects of warming and drying on peatland C.

We did not include climate change other than WT drawdown in this study, but WT dynamics are tightly connected with climate change. For example, warming itself can increase evapotranspiration and warming-induced increases in plant biomass can accelerate this change (Helbig et al., [2020](#page-14-6)), further lowering WT in peatlands. In addition, drier peatlands with lower water availability can increase sensible heat flux, which can warm lower atmosphere (Göckede et al., [2017](#page-14-21)). Changing precipitation patterns can modify WT of peatlands in the local and regional scales (Qiu et al., [2022](#page-16-20)). Because of high uncertainty and large temporal and spatial variations in the relationship among temperature, precipitation, and peatland WT, it is challenging to forecast the direction and magnitude of C response of northern peatlands and their feedback with climate. However, studies including climate change predict that northern peatlands will reduce their C sink capacity (Chaudhary et al., [2020](#page-13-16)) or lose C (Treat et al., [2021](#page-17-21); Wu et al., [2013](#page-18-11)) especially under strong warming scenarios (Chaudhary et al., [2022](#page-13-17); Qiu et al., [2022](#page-16-20)). Although the magnitude of C loss in response to WT changes in our study is less than the previously studied temperature and precipitation responses, our results indicate that drying may exacerbate C loss due warming of northern peatlands.

4.4 | **Limitations and future directions**

Despite well simulated fluxes after optimizing the model parameters, some processes can be added for further improvement. For example, our model does not differentiate peatland types, such as fens and bogs, which have distinguishable hydrologic connectivity to groundwater and nutrient supply (Charman, [2009](#page-13-18)). Greater parameter value (Vc_{max25}) for plant productivity in US-BZF (fen) than other sites could have represented higher nutrient supply, but different seasonality among peatland types may not be captured depending on the relative importance of hydrologic connectivity on the C cycle. Furthermore, processes in the standing water in fens, such as $CH₄$ oxidation in the water column (Ward et al., [2020](#page-18-12)), contribution of algae (Kane et al., [2021](#page-14-22)) and predation (Wyatt et al., [2021](#page-18-13)), can be incorporated.

Our current model has one peatland-representative plant functional type and does not distinguish multiple peatland plant species, such as sedges, mosses, shrubs, and trees, which can respond differently to short-term hydrological changes (Sulman et al., [2010](#page-17-13)), and their potential compositional change following long-term hydrological changes (Breeuwer et al., [2009](#page-13-13)). We did not include peatland forestry in this study, but intensive drainage that results in significant tree growth can be another aspect to consider, which may increase (Krüger et al., [2016](#page-15-23); Minkkinen et al., [1999](#page-16-21); Simola et al., [2012](#page-17-22)) or decrease (Krüger et al., [2016](#page-15-23); Minkkinen et al., [1999](#page-16-21); Nykänen et al., [2020](#page-16-22); Simola et al., [2012](#page-17-22)) soil C but increase plant biomass and total terrestrial C stock (Minkkinen et al., [1999](#page-16-21)). Furthermore, potentially deepening rooting depth (priming effects) may accelerate C loss as well in deeper soils (Keuper et al., [2020](#page-14-23)). In addition, our current model has constant peatland surface elevation, although peatland surfaces are not static and changes in their physical properties can alter soil biogeochemical properties: Persistent dry conditions can change the peat properties, such as bulk density, which can subsequently influence hydraulic and thermal properties, decomposition rates, and plant composition (Kreyling et al., [2021](#page-15-24); Nykänen et al., [1998](#page-16-5); Turetsky et al., [2014\)](#page-17-10). After a substantial portion of the dry peat at the surface is decomposed and becomes compact, the surface subsides and the relative WT becomes higher (hydrological self-regulation of peatlands; Belyea & Baird, [2006](#page-13-19); Dise, [2009](#page-13-10); Waddington et al., [2015](#page-18-9)). We prescribed the WT with the observations (relative to the surface), and short-term responses up to the observation point were well simulated. However, without dynamic surface elevation (e.g., subsidence) represented in the current model, WT drawdown and its effect on the C cycle can be overestimated in the long-term especially in non-permafrost peatlands (Nijp et al., [2017](#page-16-23)). This overestimation can be less of a concern in permafrost peatlands, because permafrost and ground ice block or retard the vertical and lateral water drainage, and permafrost thaw and ground ice melt will remove surface water that connects the hydrological feedback process. The intensity of this change can be also species specific and can be strong at the surface, where abundant macro-pores exist (McCarter et al., [2020](#page-15-25)). Additional model development to take into account these short term and long term as well as aboveground and belowground factors could further reduce the uncertainty of the direction and magnitude of C stock changes.

With a finely calibrated land surface model with a multilayer soil module, we demonstrated decreasing $CO₂$ sink strength and $CH₄$ emissions in response to drying, and these responses were stronger when initial WT conditions were close to the soil surface. There is limitation in upscaling this response to the whole northern peatlands based on the results from only six northern peatlands. For example, US-BZF showed the highest CH_A fluxes among all six sites of this study, but its $CH₄$ flux rate is comparably

lower than other northern peatlands (Kuhn et al., [2021](#page-15-21)) possibly because methanogens are outcompeted by reducers of alternative electron acceptors (Kane et al., [2013](#page-14-24); Rupp et al., [2021](#page-17-23)). Variations in parameters that are associated with methanogenesis (*k*) partially represent redox status, yet such processes can be included. Using averaged parameters for the whole region can over- or under-estimate fluxes (Qiu et al., [2018](#page-16-8); Salmon et al., [2021](#page-17-11); Treat et al., [2018](#page-17-19)), or converge to the average fluxes. In addition, despite several available peatland/wetland maps (Hugelius et al., [2020](#page-14-1); Olefeldt et al., [2021](#page-16-24); Xu et al., [2018](#page-18-14)) and possible usage of precipitation minus evapotranspiration as a proxy of relative WT variations (Gulev et al., [2021](#page-14-25)), the lack of highly heterogeneous local WT data in the northern peatlands will not allow us to accurately assess the current responses of northern peatlands to changes in water balance. In addition to the high uncertainty in the temporal and spatial variations in WT depending on climate and anthropogenic activities, permafrostthaw-related changes add more complexity to evaluation of long-term trends. When excessive ground ice melts, it can induce further hydrological changes (Andresen et al., [2020](#page-13-20); Lewis et al., [2012](#page-15-3); Nitzbon et al., [2020](#page-16-25); Rodenhizer et al., [2020](#page-17-24)). Thus, the inclusion of ground ice dynamics and the associated topographical and hydrological change in the model are essential to constrain C flux and stock change in permafrost peatlands (Cai et al., [2020](#page-13-21); O'Neill et al., [2019](#page-16-26)).

Although peatland drying reduced the net CO_{2-eq} fluxes, it decreased C accumulation, which is one of the crucial functions of northern peatlands. In particular, peatlands with a high WT are more vulnerable to C loss, including a large potential C loss in nonpermafrost peatlands and a possible switch from C sink to source in permafrost peatlands. Human-induced drainage can be avoided to reduce this C loss, but climate-driven peatland drying, including permafrost-thaw related drying, cannot be prevented without strong climate change mitigation. Furthermore, re-wetting does not neces-sarily restore peatlands to pre-drying conditions (Harris et al., [2020;](#page-14-16) Kreyling et al., [2021](#page-15-24)), and the intact wet conditions are critical for maximizing C storing function of peatlands.

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CONFLICT OF INTEREST

The authors declare no conflicts of interest.

DATA AVAILABILITY STATEMENT

All the model simulation data are stored at zenodo.org ([10.5281/](https://doi.org/10.5281/zenodo.6817633) [zenodo.6817633](https://doi.org/10.5281/zenodo.6817633)). Observation data used for the model optimization is available as follows: (1) FI-SAR & FI-LAK: [https://doi.org/10.5281/ze](https://doi.org/10.5281/zenodo.6967198) [nodo.6967198](https://doi.org/10.5281/zenodo.6967198), (2) US-BZF: [https://doi.org/10.17190/AMF/1756433,](https://doi.org/10.17190/AMF/1756433) (3) US-HEA: [https://doi.org/10.6073/pasta/34c20956a48edb45](https://doi.org/10.6073/pasta/34c20956a48edb4507add832b98a7fa6) [07add832b98a7fa6](https://doi.org/10.6073/pasta/34c20956a48edb4507add832b98a7fa6) and [https://doi.org/10.6073/pasta/2959f51f7](https://doi.org/10.6073/pasta/2959f51f79d067ce01d2c74a8d5fbb75) [9d067ce01d2c74a8d5fbb75,](https://doi.org/10.6073/pasta/2959f51f79d067ce01d2c74a8d5fbb75) (4) RU-CHE: [https://doi.org/10.18140/](https://doi.org/10.18140/FLX/1669654) [FLX/1669654](https://doi.org/10.18140/FLX/1669654) and [https://doi.org/10.18140/FLX/1669655,](https://doi.org/10.18140/FLX/1669655) and (5) US-BES:<https://doi.org/10.18739/A20Z70Z1H>.

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 KWON et al. **[|] 6767**

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