



Organic soil carbon balance in drained and undrained hemiboreal forests

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Abstract. Drainage of organic soils is associated with increasing soil carbon (C) efflux, which is typically linked to losses in soil C stock. In previous studies, drained organic forest soils have been reported as both C sinks and sources depending on, e.g., soil nutrient and moisture regime. However, most of the earlier research was done in boreal region, and both the magnitude of C efflux and the impact of soil moisture regime on soil C stock are likely to vary across different climatic conditions and ecosystems, depending further on vegetation. A two-year study was conducted in hemiboreal forest stands with nutrient-rich organic soil (including current and former peatlands) and a range of dominant tree species (black alder, birch, Norway spruce, Scots pine) in the Baltic states (Estonia, Latvia, Lithuania). In this study, we analysed the C balance of organic soil in drained (19) and undrained (7) sites. To assess the C balance, soil respiration was measured along with evaluation of C influx into the soil through aboveground and belowground litter. To characterize the sites and factors influencing the C fluxes, we analysed soil temperature, water table level, physical and chemical parameters of soil and soil water. On average, no changes in soil C stocks ($0.45 \pm 0.50 \text{ t C ha}^{-1} \text{ year}^{-1}$) were observed in drained sites dominated by black alder, birch, or Norway spruce, while drained Scots pine sites showed soil C removals with a mean rate of $2.77 \pm 0.36 \text{ t C ha}^{-1} \text{ year}^{-1}$. In undrained birch- and spruce-dominated sites, soil functioned as mean C sink at $1.33 \pm 0.72 \text{ t C ha}^{-1} \text{ year}^{-1}$, while the undrained black alder stands showed an uncertain C balance of $1.12 \pm 2.47 \text{ t C ha}^{-1} \text{ year}^{-1}$. The variability in C balances were influenced by the nutrient-rich soil exhibiting a wide range of nutrient conditions and organic matter quality. Thus, indicating that soil macronutrient concentrations and pH can determine whether the soil functions as a C source or sink.

1 Introduction

Soil in peatlands, characterised by its high content of partially decomposed plant matter, is a major terrestrial organic carbon (C) stock, estimated to range from 504 to 3000 Gt C (Scharlemann et al., 2014). Although northern peatlands make up only 2–4% of the global land area, they contain a substantial amount of soil C, estimates ranging from around 500 to 1,055 Gt C (Yu, 2012; Nichols and Peteet, 2019), highlighting the significance of these lands in the global C budget. About 28% of the pristine (undrained) peatlands globally are inherently covered by forest (Zoltai and Martikainen, 1996), and those peatland forests in the boreal biome can accumulate C into the soil at similar rates to non-forested peatlands; the higher decomposition rates observed in peatland forests (Beaulne et al., 2021) can be compensated by higher litter inputs (Straková et al., 2010). To enhance tree growth, peatland drainage for forestry has been common in the past. Drainage facilitates oxygen access to deeper peat layers, thereby promoting tree root survival and function, but also the mineralization of organic matter and the release of C into the atmosphere in the form of CO₂. Therefore, the conservation of organic soil C stocks in managed current and former peatlands has attracted attention in the context of climate change.



41 The approximately 13 million ha of forestry-drained organic soils in Europe have been estimated to emit 17 million tons of
42 CO₂ per year (European Environment Agency, 2023). Despite the temperate region being characterized by wide climatic
43 gradients, currently, only a single default emission factor (EF) by the Intergovernmental Panel on Climate Change (IPCC) is
44 available for the entire temperate climate region, to which the Baltic states correspond according to the IPCC (Hiraishi et al.,
45 2013). The EF was developed using study results from 8 drained sites (Hiraishi et al., 2013), which were published in 5 articles
46 (Glenn et al., 1993; Minkinen et al., 2007; Yamulki et al., 2013; Von Arnold et al., 2005). These studies employed different
47 CO₂ estimation methods, complicating the comparability of the aggregated results (Jauhiainen et al., 2023; Jauhiainen, 2019).
48 None of the sites are in the Baltic states. Given that the soil emissions in the boreal zone are smaller than in the temperate zone
49 (Jauhiainen et al., 2023), and Baltic states are situated in the hemiboreal vegetation zone (Ahti et al., 1968) – in transition
50 between the temperate and boreal zones – the IPCC's default temperate EF may not be suitable for application in this region.
51 The same issue arises on a broader geographic scale, where the use of unharmonized country-specific and default EFs creates
52 challenges in comparing estimated emissions both within and across different countries and climate regions. Discrepancies in
53 CO₂ emissions between regions can be expected due not only to climate gradient (Ojanen et al., 2010) but also to site
54 productivity (Janssens et al., 2001). While higher ecosystem productivity is associated with increased soil respiration rates
55 (Janssens et al., 2001) it also facilitates higher C influx through litter (Krasnova et al., 2019). Therefore, EFs are most
56 appropriately used when applied to areas with similar environmental conditions, rather than being limited by national
57 boundaries or applied too broadly across diverse geographic regions.

58 In the Baltic states (Estonia, Latvia, and Lithuania) organic soils (Eggleston et al., 2006) are current or former peatlands where
59 a peat layer is still identifiable, or, due to high decomposition, no longer meet the typical characteristics of peat. However,
60 these soils by definition contain at least a 20 cm thick layer rich in organic matter (organic layer). In the region, the total area
61 of drained organic forest soils is reported to be 0.8 million ha, with estimated emissions of 1.8 million tons of CO₂ per year
62 (Ministry of the Environment of Republic of Estonia, 2021; Konstantinavičiūtė et al., 2023; Skrebele et al., 2023). Thus,
63 countries with a relatively small total land area yet a substantial proportion of organic soil can have a considerable role in
64 organic soil management. This underscores the importance of acquiring precise estimates of the CO₂ emissions from organic
65 forest soils in this region. However, despite the Baltic States being located next to each other and thus expectedly showing
66 similar soil CO₂ emissions from comparable sites and land uses, the emission estimation approach is currently not harmonized
67 as the countries use different EF to estimate and report emissions (Ministry of the Environment of Republic of Estonia, 2021;
68 Konstantinavičiūtė et al., 2023; Skrebele et al., 2023). According to National Greenhouse Gas Inventories submissions of
69 2023, the CO₂ emissions of drained organic forest soil in Estonia and Lithuania were estimated using the default EF provided
70 by IPCC for the temperate region (Calvo Buendia et al., 2019), while Latvia applied a country-specific EF. Due to similarities
71 in biogeography, climate, and land-use practices, common EFs based on regionally representative, congregated data could be
72 a better option.

73 A recent synthesis evaluated whether default IPCC EF can be improved by compiling results from the most recent studies
74 (Jauhiainen et al., 2023). Still, only modest, and insignificant changes judging by confidence intervals (CI) of IPCC EFs could
75 be introduced for the temperate climate region as a whole, due to limited data (Jauhiainen et al., 2023). Although the general
76 driving factors of CO₂ emissions are known, the number and geographical representation of studies on drained soils,
77 particularly in the temperate zone, remain too limited for stratification of EFs based on local conditions. It is widely known
78 that soil temperature is the primary factor influencing gross CO₂ emissions. However, while temperature can explain variations
79 in emissions, it does not fully account for their magnitude. The extent of CO₂ emissions is affected by soil properties (Basiliko
80 et al., 2007), incorporating complex interactions between a soil's physical, chemical, and biological characteristics, including
81 the impact of the litter quality (Berger et al., 2010). The recent synthesis confirmed that key factors influencing the magnitude
82 of CO₂ emissions include soil C concentration, carbon-to-nitrogen (C:N) ratio, and bulk density, as well as stand type



(Jauhiainen et al., 2023). Yet, the list likely remains quite incomplete, as individual studies in different site types using various methods can provide unharmonized results making it difficult to identify the relationships influencing CO₂ emissions. Varying and often insufficient reporting of study site conditions in previous studies (Jauhiainen, 2019) further hampers the ability to compile the results for effective synthesis and meta-analysis (Jauhiainen et al., 2023). This limitation may have hindered the identification of emission-impacting factors and the ability to quantify their relationships, underscoring the need for more localized studies to address these gaps, particularly in the Cool Temperate Moist climate region (Calvo Buendia et al., 2019), which overlaps with the hemiboreal vegetation zone.

In the few studies on drained and undrained soil C balance conducted in the Baltic states, using both chamber and soil inventory methods, findings have been inconsistent (Butlers et al., 2022; Lazdiņš et al., 2024; Bārdule et al., 2022). Drained organic soils have been identified as both C sinks and sources, with no decisive conclusions reached regarding the factors driving such variation. Soil C loss of nutrient-rich organic soil has been identified using the soil inventory method (Lazdiņš et al., 2024) but not confirmed by the chamber method (Butlers et al., 2022; Bārdule et al., 2022). Given that nutrient-rich organic forest soils can make up to 72 % of total organic forest soils (Līcīte et al., 2019) and are associated with a higher risk of soil C loss, there is a need to enhance our understanding of the underlying drivers to improve the accuracy of C balance estimates in Baltic states. Therefore, studies using harmonized data collection methods are necessary covering variety of nutrient-rich forest soils, recording or monitoring essential environmental conditions, and accounting for both soil C efflux and influx from various litter sources.

This study aimed to quantify the soil C balance in hemiboreal forests with nutrient-rich organic soil and different dominant tree species. The research was carried out in 26 forest stands with organic soil in Estonia (EE), Latvia (LV), and Lithuania (LT), including both undrained and drained sites, over two years. We analysed soil CO₂ emissions as well as C inputs by tree fine roots, ground vegetation (below- and aboveground) and fine foliar litter. We examined factors contributing to soil C balance by indirectly influencing soil processes, such as soil and soil water properties and water-table level (WTL). We also evaluated the impact of dominant tree species and whether soil emissions differ between countries. To facilitate the use of the results in future syntheses or meta-analyses, the data used for C balance estimations have been openly published.

2 Materials and methods

2.1 Study sites

In total, 26 study sites (Figure 1) were established in stands dominated by black alder (*Alnus glutinosa* (L.) Gärtner), birch (*Betula pendula* Roth, *Betula pubescens* Ehrh.), Scots pine (*Pinus sylvestris* L.), or Norway spruce (*Picea abies* (L.) Karst.) of different ages (Table S1). The study sites included both drained (n=19) and undrained (n=7) organic soil (Eggleston et al., 2006), with the organic layer thickness ranging from 27 cm to over 2 meters, measured by a rod insertion. According to forest type classification, all sites were characterised as nutrient-rich based on ground vegetation and stand productivity (Bušs, 1981). Drained sites were represented by two site types: *Oxalidosa turf. mel.* (*Oxalidosa*), which has relatively higher nutrient availability (pH, macronutrients), and *Myrtillosa turf. mel.* (*Myrtillosa*). Undrained sites were represented by the *Dryopterioso-caricosa* site type. Soil drainage status was determined based on the presence of drainage ditches within or along the respective forest compartment (a rectangular forest area of 50 ± 25 ha used as a management unit).

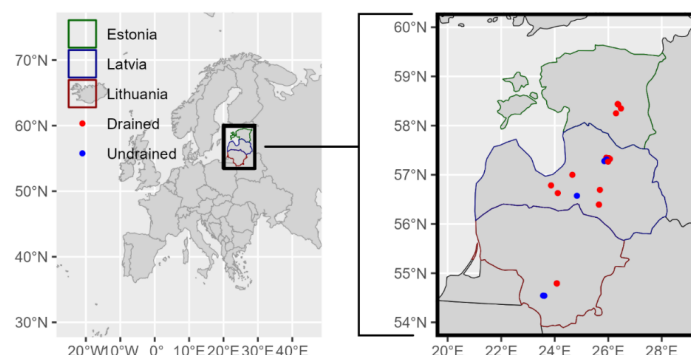


Figure 1: Locations of the study sites. Points indicate the locations of study site clusters.

Despite the more numerous drained sites showing greater variation in the stand characteristics than the undrained sites, overall, the two groups of sites were comparable (Table 1). The mean stand age of both groups was 74 years, with a range of 26-162 years for drained and 44-96 years for undrained sites. Average basal areas in turn were 27 for drained and 30 m² ha⁻¹ for undrained sites, respectively. More information on stand characteristics, including mean WTL and coordinates, is provided in Table S 1. The projective cover of the most common ground vegetation species in the study sites is presented in Table S 2.

Both groups represent historically naturally forested lands that have undergone active forest management. While the exact time since drainage is unknown, active drainage of the forests was initiated in the mid-19th century and resumed in the mid-20th century. Drainage ditches were dug to depths of 0.8 to 1.2 m, with distances between them ranging from 100 to 400 m, and they are maintained to remain functional (Zālītis, 2012).

Table 1: Range (minimum; maximum) of tree stand characteristics in the study sites.

Parameter	Dominant tree species							
	Black alder		Birch		Pine		Spruce	
	Drained	Undrained	Drained	Undrained	Drained	Undrained	Drained	Undrained
Site count	n = 2	n = 2	n = 5	n = 2	n = 5	-	n = 7	n = 3
Age, year	30; 80	44; 74	24; 45	44; 61	60; 141	-	40; 162	81; 96
Mean height, m	13; 20	16; 28	13; 18	9; 20	12; 21	-	10; 23	15; 20
Mean diameter, cm	12; 21	16; 28	12; 22	8; 21	12; 22	-	10; 25	17; 21
Basal area, m ² ha ⁻¹	26; 36	30; 36	15; 23	22; 23	17; 48	-	18; 36	25; 42

At each study site, three subplots (Figure S 1) for data collection were selected, with a minimum distance of 30 m, in an area representing typical ecosystem characteristics of the floristically defined site type, as classified under the local forest site type system (Bušs, 1981). The subplots were arranged along a transect positioned perpendicular to the closest ditch in drained areas and perpendicular to stand borders in undrained areas. In the drained sites, the first subplot was located about 20 m from the nearest drainage ditch. In the undrained sites, the first subplot was located at least 20 m from the forest border.

Empirical data was gathered from January 2021 to December 2022 in Estonia and Latvia, and from July 2021 to June 2023 in Lithuania. The sites were visited monthly in Latvia and Lithuania, and biweekly in Estonia. The meteorological conditions during the study were typical for the region (Table S 3).



139 2.2 Respiration

140 Measurements of forest floor respiration, which we will hereafter refer to as total respiration (R_{tot}), included both soil
141 heterotrophic respiration and autotrophic dark respiration of ground vegetation (above and below-ground) as well as tree roots
142 extending to the measurement locations. Gas samples were collected from manual closed static dark (opaque) chambers (PVC,
143 volume 0.0655 m^3) as described in the literature (Hutchinson and Livingston, 1993) for subsequent laboratory analysis. R_{tot}
144 monitoring included five to six monitoring locations, divided over 3 subplots, at each site. Ring-shaped chamber collars (area
145 0.196 m^2) were permanently installed in the soil at a depth of five cm at least one month before the first sampling to avoid the
146 installation effect on fluxes. Collar locations reflected local diversity in vegetation and potential WTL gradient at each subplot
147 and the encircled soil surface and vegetation were kept intact. Thus, the soil heterotrophic respiration component in R_{tot}
148 includes CO_2 emissions caused by the decomposition of both litter and soil organic matter.

149 Gas samples for analyses were collected by obtaining four air samples from a closed chamber into pre-evacuated (0.3 mbar)
150 glass vials (100 cm^3). Air samples were taken from the chamber outlet, equipped with a valve attached in the sampling tube
151 reaching approximately the centre of the airspace. The air within the chamber was not artificially mixed during sampling. Air
152 sampling was done by first removing the residual air left from the sampling tube (to avoid potential impact on the concentration
153 readout) by a syringe, and thereafter pre-evacuated glass vial was attached into the outlet. The first sample was taken
154 immediately after attaching the chamber on the collar, and subsequent samples were taken at either 10 (Latvia) or 20 (Estonia
155 and Lithuania) minute intervals over 30- or 60-minute monitoring periods, respectively (Vigricas et al., 2024; Butlers et al.,
156 2022).

157 The gas samples were analysed using a Shimadzu GC-2015 gas chromatograph (Shimadzu USA Manufacturing, Inc., Canby,
158 OR, USA) equipped with an electron capture detector (ECD). The uncertainty of the method used was estimated to be 20 ppm
159 of CO_2 (Magnusson et al., 2017). Linear regression was applied to relate the CO_2 concentrations with the time elapsed since
160 chamber closure for each measurement. Subsequently, the measurement data was screened to identify deviations from the
161 recognized trend, considering the removal of measurements with identified errors. All measurements were discarded if the
162 regression coefficient of determination (R^2) was less than 0.9 ($p < 0.01$), except for cases where the difference between the
163 highest and lowest measured CO_2 concentration in the chamber was less than the uncertainty of the method (specifically
164 applicable during non-vegetation periods). Consequently, an insignificant amount of data was discarded.

165 The data that met the quality criteria were used to determine the slope coefficient of the linear regression, which was then used
166 to calculate the instantaneous R_{tot} according to the ideal gas law equation (Fuss and Hueppi, 2024):

$$R_{tot} = \frac{M \times P \times V \times \text{slope}}{R \times T \times A \times 1000} \quad (1)$$

167 where R_{tot} is the instantaneous total respiration, $\text{mg CO}_2\text{-C m}^2 \text{ h}^{-1}$; M is the molar mass of $\text{CO}_2\text{-C}$, 12.01 g mol^{-1} ; R is the
168 universal gas constant, $8.314 \text{ m}^3 \text{ Pa K}^{-1} \text{ mol}^{-1}$; P is the assumption of air pressure inside the chamber, 101.300 Pa ; T is the air
169 temperature in the chamber, K ; V is the chamber volume, 0.0655 m^3 ; the slope is the CO_2 concentration change over time,
170 ppm h^{-1} ; and A is the collar area, 0.19625 m^2 .

171 We also conducted measurements of heterotrophic respiration (R_{het}) for comparison, as described in the Supplementary text.

172 2.3 Environmental variables

173 Manual WTL measurements were carried out using nylon-mesh-coated, perforated piezometer tubes (5 cm in diameter)
174 installed down to a 140 cm depth in all subplots. Manual soil temperature measurements were done at depths of 5, 10, 20, and
175 40 cm in all subplots by Comet data logger (COMET SYSTEM, s.r.o., 756 61 Roznov pod Radhostem, Czech Republic)
176 equipped with Pt1000 temperature probes. All manual measurements were carried out at the same time as CO_2 flux



177 measurements. Continuous soil temperature measurements at depths of 10 and 40 cm were carried out at 30-minute interval in
178 the centermost subplot (Maxim Integrated DS1922L2F, iButtonLink Technology, Whitewater, WI 53190 USA).

179 Once per month, soil water samples were collected from perforated tubes (7.5 cm in diameter) explicitly installed for water
180 sampling for chemical analysis. Water chemical parameters including pH, electrical conductivity (EC), and concentrations of
181 dissolved organic carbon (DOC), total nitrogen (N), nitrate (NO_3^-), ammonium (NH_4^+), and phosphate (PO_4^{3-}) ions were
182 determined. Soil samples were taken up to a depth of 75 cm (0-10; 10-20; 20-30; 30-40; 40-50; 50-75cm) at two locations in
183 each subplot during the establishment of the study sites. Two separate sample sets were collected – for the determination of
184 bulk density, ash content and chemical parameters (pH, concentrations of total carbon (TC), nitrogen (N), and HNO_3
185 extractable phosphorus (P), potassium (K), calcium (Ca), and magnesium (Mg)). The samples were collected with a volumetric
186 100 cm³ cylinder (Cools and De Vos, 2010) at 10 cm intervals to a depth of 50 cm. Two additional samples were taken from
187 soil depths of 50-75 and 75-100 cm with a soil auger. Soil samples collected for determination of bulk density were oven-dried
188 (105 °C) and weighed, while soil samples for chemical analyses were prepared by air drying (≤ 40 °C), sieving and
189 homogenizing (LVS ISO 11464:2006). Organic carbon (Corg) content was calculated by multiplying the ash content
190 measurement result derived soil organic matter content by factor 0.5, thus assuming that organic matter is 50% Corg (Pribyl,
191 2010). All soil and water analyses were done in an ISO 17025-certified laboratory using ISO standard methods (Table S 4).

192 2.4 Litter input

193 Annual litter inputs to be used in C balance estimation were either measured (foliar litter) or estimated based on biomass
194 components measured (ground vegetation, fine roots).

195 Foliar fine litter (fLF) was collected with conical litter traps (area 0.5 m²) set one meter above ground (Latvia, Lithuania), or
196 with square mesh frames (0.5 x 0.5 m) placed on the ground (Estonia). In each study site, five replicate litter traps were placed
197 in the centremost subplot of the transect. It included all fine fractions of litter, such as needles, leaves, and branches with a
198 diameter up to 1 cm and a length up to 10 cm. Branches with larger dimensions were collected from coarse woody litter (cLF)
199 traps (square mesh frames, 0.5 x 0.5 m). The litter samples were collected from the traps once every four weeks. Due to the
200 heterogenous nature and large dimensions of cLF, respective decomposition emissions could not be representatively included
201 in R_{tot} measurements. Therefore, we considered only fLF as the litter input source, and the cLF results are presented only as
202 indicative of the site conditions.

203 Ground vegetation (GV) aboveground (aGV) and belowground (bGV) biomass samples were collected at the end of the
204 growing season (August) in 2021 in five replicates per subplot, from square sampling locations with an area of 0.0625 m².
205 Aboveground biomass was separated into herbaceous (aGV) species and dwarf shrubs. bGV was collected from the top soil
206 layer, extending down to 20 - 30 cm from the same location and area as aGV. Herbaceous ground vegetation roots (bGV) were
207 separated from tree and shrub roots by wet sieving based on root morphological properties. We assumed that the measured
208 herbaceous ground vegetation aboveground (aGV) and belowground (bGV) biomasses were equal to annual ground vegetation
209 litter inputs. The contribution of shrub litter was thus not included but was assumed to be minor (see Table S 5).

210 Moss biomass was measured by collecting samples from 0.01 m² square areas, with four replicates per subplot, and visually
211 removing any dead parts. Collections were conducted concurrently with moss production sampling. Moss production (MP)
212 samples in four replicates per subplot were collected by anchoring a square mesh (0.01 m²) on the moss at the end of
213 the growing season and harvesting the moss biomass grown through the mesh by the end of the next growing season. Including
214 moss litter in the C balance estimation by aligning C inputs and outputs from areas with and without moss cover, without
215 increasing uncertainty, would however have required doubling the number of spatial replicates in chamber measurements.
216 Therefore, we did not consider moss as a litter source, which causes underestimation of litter inputs to a varying extent. The
217 moss results are presented only as indicative of site conditions (see Table S 5).



Tree fine-root production (FRP) was estimated with the ingrowth core method (Laiho et al., 2014; Bhuiyan et al. 2017). Five replicate ingrowth cores (cylindrical mesh bags with diameter 2.5 cm, mesh size 2 mm) per subplot, filled with soil collected from the subplot, were installed in autumn or spring and removed after two growing seasons. In the laboratory, the biomass of the ingrown fine roots was determined after wet sieving, and ground vegetation roots were separated from tree roots by morphological properties. We assumed that tree fine-root biomass was essentially not changing over the study years, and thus we could assume that FRP equalled litter production. Since the ingrowth cores were removed from the soil after two growing seasons, the FRP estimate was calculated by dividing the fine-root biomass in the cores by two (Bhuiyan et al., 2017).

All litter and biomass samples were oven-dried (70 °C), weighed and milled before further analysis. Chemical analyses were performed according to ISO standard methods (Table S 4). To estimate the annual C inputs to be used in the C balance estimation (section 1.5), the estimated annual litter inputs on a unit area were transformed to C input by using C content values measured for each component (Table 2).

Table 2: Mean C and N content (% of dry matter) in foliar litter from trees and biomass components used for estimating litter inputs. The values are means \pm standard deviation including both drained and undrained sites. Abbreviations: aGV and bGV – above- and belowground biomass of herbaceous vegetation, FR – tree fine roots, M – moss, fLF – foliar fine litter, cLF – coarse woody litter.

Element	aGV	bGV	FR	M	fLF	cLF
C	49.34 \pm 2.45	50.95 \pm 2.02	51.21 \pm 5.16	48.38 \pm 2.13	52.50 \pm 0.25	53.88 \pm 0.67
N	2.18 \pm 0.64	1.53 \pm 0.43	1.47 \pm 0.44	1.10 \pm 0.75	1.30 \pm 0.41	1.04 \pm 0.20

2.5 Estimation of annual soil carbon balance

We estimated the annual soil C balance of the sites by combining annualized C input (section 1.4) and output data. In exceptional cases where the C input from specific source was not estimated in certain countries, such as bGV and FRP in EE and FRP in some sites in LT and LV (Table S 10), we assumed the C input to be equivalent to the average results from sites with the same drainage status in other countries. While we directly measured R_{het} , we utilized the estimated R_{het} derived from R_{tot} as output value (marked from here onwards as R_{het}^*). Such an approach was necessary because, at all sites, our R_{het} values were significantly higher (by mean $5.8 \pm 3.1 \text{ t CO}_2\text{-C ha}^{-1} \text{ year}^{-1}$) than R_{tot} , which would be logically impossible if the conditions in the measurement locations were the same. This pattern was probably mostly due to the high and variable CO_2 efflux from roots killed by the trenching, noted also in other studies (e.g., Hermans et al., 2022). This discrepancy could not be remedied with the root data at hand (for more details, please refer to the Supplementary text and Discussion section).

To estimate annual C output, at first, site-specific relationships between instantaneous R_{tot} fluxes and soil temperatures were established. To identify the best approach for to express the relationship, we compared the suitability of exponential regression on untransformed data and linear regression on logarithmically or Box-Cox transformed data. The performance of these models for flux data interpolation based on continuous soil temperature data was evaluated by using the root mean square error (RMSE) of prediction. It was found that the Box-Cox approach achieved the best conformity of flux data to a normal distribution and provided the best fit for the models. Previous studies also indicate that this method effectively addresses the typical underestimation of fluxes caused by their nonlinearity (Box and Cox, 1964; Liaw et al., 2021; Wutzler et al., 2020). Interpolation was performed by evaluating the relationship between R_{tot} and soil temperature (at 10-cm depth) measured at each study site and constructing site-specific linear regression equations (Table S 6) after applying Box-Cox transformation to the flux data for normalization (Box and Cox, 1964). This approach, compared to alternative methods, more successfully (lower RMSE of prediction) accounted for the exponential nature of the relationship between flux and temperature, and prevented underestimation (Wutzler et al., 2020; Liaw et al., 2021). Hourly R_{tot} estimates were then formed by using the hourly recorded soil temperatures (logger data) for each study site. Consecutively, site-specific annual R_{tot} estimate was calculated by summing the interpolated hourly emission estimates of the year.



We derived site-specific annual R_{het} from estimated annual R_{tot} empirically. To avoid potential bias introduced by using a single fixed factor, we instead used Equation (2), accounting for the observed pattern of decreasing R_{het} proportion as soil surface respiration (R_s) increases (Subke et al., 2006; Bond-Lamberty et al., 2004). The equation characterizes the relationship between R_s and R_{het} , and was created using results of previous studies (Jian et al., 2021) in the boreal zone (Figure S 2). We assumed R_{tot} is equal to R_s , i.e., that aboveground autotrophic respiration has a minor role in R_{tot} (Hermans et al., 2022; Munir et al., 2017), and applied the equation to annual R_{tot} directly.

$$R_{het} = -0.70 + 0.78 \times R_{tot} \quad (2)$$

Neglecting aboveground autotrophic respiration leads to a minor overestimation of R_{het} .

2.6 Statistical analysis

Statistical analyses were performed and figures prepared using the software R version 4.3.1 (packages ‘MASS’, ‘stats’, ‘nlme’, ‘Hmisc’, ‘lmerTest’, ‘lme4’, ‘vegan’, ‘pls’, and ‘caret’), using $p=0.05$ as the limit for statistical significance. The compliance of the data with the normal distribution was checked with the Shapiro-Wilk normality test and visually by density and quantile-quantile (Q-Q) plots. To calculate the uncertainty of the study results when combining multiple data sources, we used the root sum of the squares method to aggregate the individual uncertainties (95 % confidence interval). Therefore, for the C influx, for instance, we combined the uncertainties from various influx (fLF, GV, FRP) sources. The uncertainty of Equation (2) used for the calculation of R_{het} was expressed as the root mean square error (RMSE) of the corresponding regression model. The soil C balances were calculated by summing the C influx and efflux for individual sites. The uncertainties of the averaged C balances, categorized by drainage status, country, site type, or dominant tree species, are expressed as standard error. Figures are prepared by using packages ‘ggplot2’, ‘corplot’, ‘ggbiplot’.

Correlations between the sample groups were expressed with the Pearson correlation coefficient (r). We compared differences between two sample groups using pairwise Wilcoxon rank sum tests with continuity correction and adjusted the p -values using Bonferroni correction. The method was used to compare soil parameters and instantaneous or annualized R_{tot} data segregated by site type, drainage status, country or dominant tree species. In the same way, soil characteristics were compared between drained and undrained sites. Multivariate flux and impacting factor relationships testing involved assessing the significance of these factors on the relationship between soil temperature and R_{tot} using mixed-effects linear models. As flux impacting factors, we considered country, dominant tree species, drainage status, WTL, and a 30 cm WTL threshold. In this analysis, Box-Cox transformed flux data was fitted to linear models using the study site as a random effect. In addition, multivariate relationships were observed through Principal Component Analysis (PCA) to visualize covariation and seek observational confluences with the results of other analyses. To assess the contribution of influencing factors on the soil C balance, Redundancy Analysis (RDA) and Partial Least Squares Regression (PLSR) were conducted.

3 Results

3.1 Soil and soil water characteristics

The organic layer depth in the drained sites ranged from 27 to 212 cm (mean 81 ± 47 cm) and in undrained sites from 100 to 230 cm (mean 167 ± 49 cm). Soil bulk density (0-30 cm depth) in the drained sites (mean 314 ± 215 kg m⁻³) was characterized by both higher variation and higher mean density ($p=0.003$) compared to undrained sites (mean 168 ± 32 kg m⁻³) (Table S 7). Soil drainage status had no impact on Corg content in the 0-30 cm soil layer ($p=0.11$, total mean 416 ± 130 g kg⁻¹). However, drained soils had a higher mean C:N ratio (22 ± 7 ; $p=0.01$) than the undrained soils (17 ± 3). A trend could be observed that undrained soils had higher nutrient concentrations and higher pH than the drained soils. Soil analysis confirmed that site types



classified based on vegetation presence served as an indicator of nutrient availability (Figure 2). In drained *Oxalidosa* sites, mean macronutrient concentrations in the 0–30 cm soil layer was higher compared to *Myrtillosa* sites. Notably, N, Mg, Ca, and P showed statistically significant differences ($p < 0.05$). Additionally, when comparing drained sites, pH values in *Oxalidosa turf. mel.* sites were, on average, 1.89 units higher ($p = 0.018$).

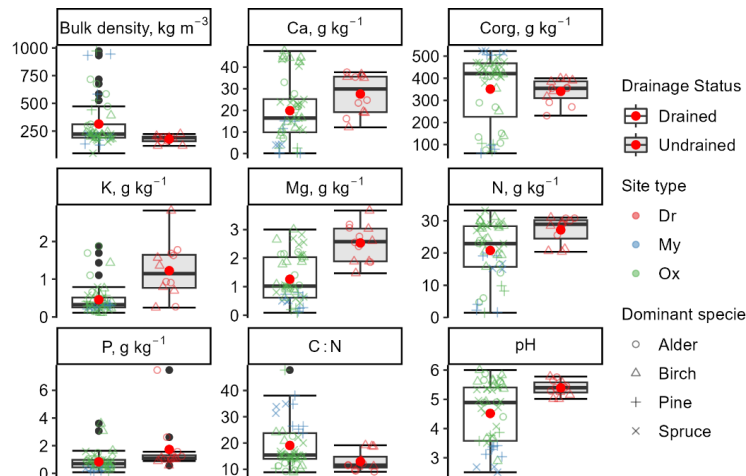


Figure 2: Variation of soil chemical and physical properties at soil depth 0-30 cm. The clear box represents the drained, and the grey-shaded box the undrained sites. The bottom and top edges of the box represent the 25th and 75th percentiles, summarizing the interquartile range (IQR). The whiskers extend to the smallest and largest values within $1.5 \times$ IQR from the 25th and 75th percentiles, respectively. Black dots mark outliers. A red dot and a solid horizontal line in the box indicate mean and median values, respectively. Corg – organic carbon; N – total nitrogen. Site types: Dr - *Dryopteris-caricosa*; Ox - *Oxalidosa turf. mel.*; My - *Myrtillosa turf. mel.*

The range of mean WTL over the study period was from -23 to -112 cm (mean -60 ± 25 cm) in the drained sites and from -7 to -17 cm (mean -13 ± 4 cm) in the undrained sites, respectively. In the undrained sites, the WTL was mainly rather high (see interquartile range in Figure 3) and had comparably smaller variation (mean standard deviation 16 cm) than in the drained sites (mean standard deviation 23 cm); however, in all sites except LTC108, WTLs below 30 cm were also observed (Figure 4). In the undrained sites, the range of min-max WTL was from 3 ± 3 cm to -63 ± 27 cm, while the WTL in drained sites had a greater absolute variation ranging from -14 ± 19 cm to -104 ± 28 cm.

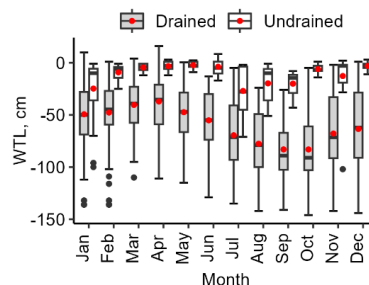


Figure 3: Yearly variation of soil water-table level (WTL) in the study sites. The edges of the box represent the 25th and 75th percentiles, encapsulating the interquartile range (IQR). The whiskers extend to the smallest and largest values within $1.5 \times$ IQR from the 25th and 75th percentiles, respectively. Black dots mark outliers. A red dot and a solid horizontal line indicate the average values of the date represented – mean and median, respectively.



The concentrations of all measured chemical parameters in the soil water, except for NH_4^+ , were, on average, higher in the drained sites (Figure S 3). However, the results were highly variable (Table S 8).

3.2 Instantaneous total respiration

In the drained sites, the mean instantaneous R_{tot} varied from 48 to 125 $\text{mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$, and in the undrained sites from 38 to 80 $\text{mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$ (Figure 4). In all sites combined, during the summer months (June, July, August), the interquartile range of R_{tot} varied from 111 to 198 $\text{mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$ with a mean of $160 \pm 78 \text{ mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$. In contrast, during the winter (December, January, February), it ranged from 8 to 24 $\text{mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$ with a mean of $17 \pm 14 \text{ mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$. The relative standard deviations of the instantaneous R_{tot} in drained ($90 \pm 9\%$) and undrained ($106 \pm 29\%$) sites were comparable. Although the study sites represented a broad soil WTL gradient, no significant impact of the site mean WTL on the mean instantaneous R_{tot} emission was observed ($r=0.16$, $p>0.05$). Furthermore, no significant correlations were found between instantaneous R_{tot} and soil water parameters.

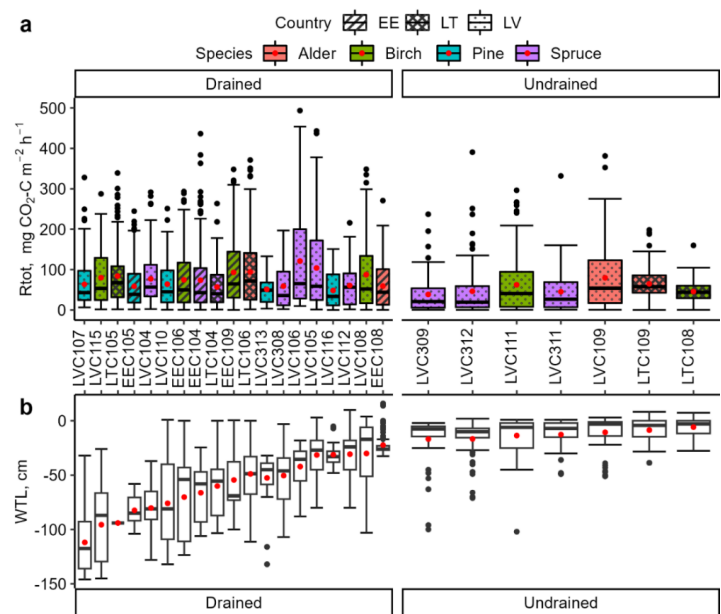
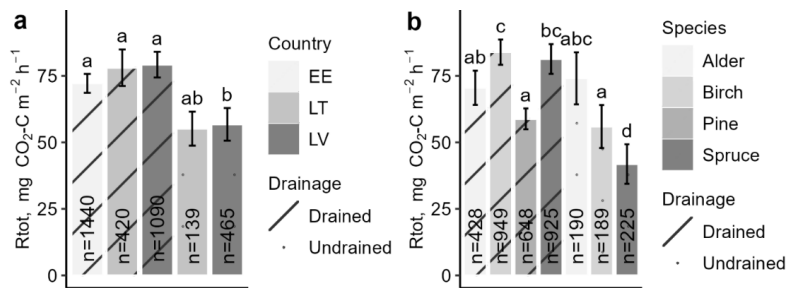


Figure 4: Variation of instantaneous total respiration (R_{tot} , a) and soil water-table level (WTL, b) in the study sites. The bottom and top edges of the box represent the 25th and 75th percentiles, summarizing the interquartile range (IQR). The whiskers extend to the smallest and largest values within $1.5 \times \text{IQR}$ from the 25th and 75th percentiles, respectively. Black dots mark outliers. A red dot and a solid horizontal line in the box indicate mean and median values, respectively.

Mean R_{tot} in sites with the same drainage status did not differ ($p>0.05$) between countries (Figure S 4). The instantaneous R_{tot} in drained sites (mean $76 \pm 3 \text{ mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$) was overall higher ($p<0.05$) than those from undrained soil (mean $56 \pm 5 \text{ mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$). The measured R_{tot} of undrained soil were smaller in both Latvia (mean $57 \pm 6 \text{ mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$) and Lithuania (mean $55 \pm 6 \text{ mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$) compared to R_{tot} from drained soil in the Baltic states ranging from mean 72 ± 4 to $79 \pm 5 \text{ mg CO}_2\text{-C m}^{-2} \text{ h}^{-1}$ (Figure 5a).



340

341 **Figure 5: Mean instantaneous total respiration (Rtot) throughout the study period, categorized by drainage status and country (a)**
342 **or dominant tree species (b).** Error bars indicate confidence interval. A shared letter indicates that differences are not significant.

343 There were few apparent differences in the mean Rtot between stands of different tree species (Figure 5b). Rtot was the lowest
344 at undrained sites dominated by spruce and the highest at drained sites dominated by birch. Rtot was significantly different
345 ($p < 0.05$) between coniferous forest sites with different dominant tree species and soil moisture regimes, where Rtot ranged
346 from mean 42 ± 7 mg CO₂-C m⁻² h⁻¹ in undrained spruce forests to 59 ± 4 and 81 ± 6 mg CO₂-C m⁻² h⁻¹ in drained pine and spruce
347 forests, respectively. In deciduous stands, the moisture regime and dominant tree species had less impact on the mean flux;
348 Rtot was higher ($p < 0.05$) in drained birch stands (mean 84 ± 5 mg CO₂-C m⁻² h⁻¹) than in undrained sites (56 ± 8 mg CO₂-
349 C m⁻² h⁻¹), while in alder stands the mean Rtot was similar regardless of the soil moisture regime (total average 67 ± 9 mg CO₂-
350 C m⁻² h⁻¹), (Figure 5b). In drained coniferous and deciduous sites, on average, the mean Rtot was similar, but in undrained
351 sites, emissions in deciduous forests were about 40% higher.

352 The impact of WTL is reflected in the mean Rtot, which was 87 ± 3 mg CO₂-C m⁻² h⁻¹ when WTL was below 30 cm and
353 57 ± 3 mg CO₂-C m⁻² h⁻¹ when WTL was closer to soil surface. However, when evaluating the effect of WTL on Rtot variation
354 in mixed-effects models predicting Rtot based on soil temperature, WTL was found to have an insignificant impact on Rtot
355 variation. Similarly, the contribution of country and dominant tree species to Rtot prediction was marginal (Table S9). The
356 inclusion of dominant species provided minimal model improvement ($\Delta AIC = +5$, $\Delta \log Lik = 0$), while country effects captured
357 some additional variability in Rtot. However, the increase in R² due to country variables was minor (from 0.77 to 0.78),
358 indicating limited explanatory power.

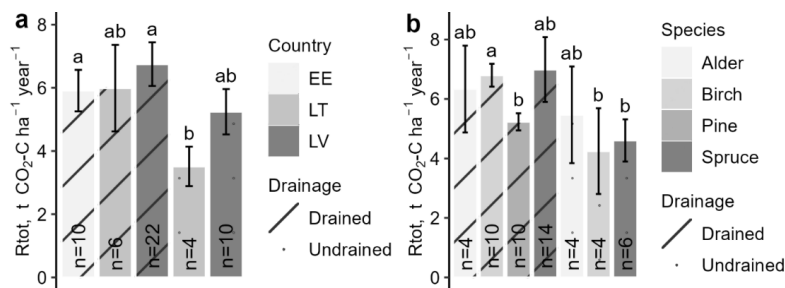
359

360 3.3 Annual total respiration

361 Soil temperature at 10 cm depth (Figure S 5) was used in constructing Rtot prediction models and for emission interpolation
362 needed for annualizing Rtot. The 10 cm depth was chosen because it showed the strongest correlation between instantaneous
363 Rtot and soil temperatures measured at different depths, with a mean Pearson correlation coefficient (r) of 0.86 ± 0.04 across
364 the study sites. For the other soil depths (5, 20, 30, 40 cm), r ranged from 0.71 ± 0.07 to 0.79 ± 0.05 . Linear models developed
365 using Box-Cox transformed data provided the best Rtot prediction power. A lambda value of 0.3411 was used for all data
366 transformations, as individual data transformations for each site resulted in comparatively less successful data normalization.
367 With this approach, the RMSE of instantaneous Rtot predictions for individual sites decreased by an average of $16 \pm 14\%$,
368 compared to linear models with logarithmically transformed data or non-linear models with untransformed data (Table S 6).

369 Annualized Rtot indicated similar mutual relationships among the study site dominant tree species and drainage status
370 categories as the instantaneous Rtot. Consequently, the estimated annual emissions from neither drained sites (overall mean
371 6.21 ± 0.43 t CO₂-C ha⁻¹ year⁻¹) nor undrained sites (overall mean 4.38 ± 1.20 t CO₂-C ha⁻¹ year⁻¹) differed significantly
372 between countries and were generally higher from drained soils (Figure 6a).

373



374

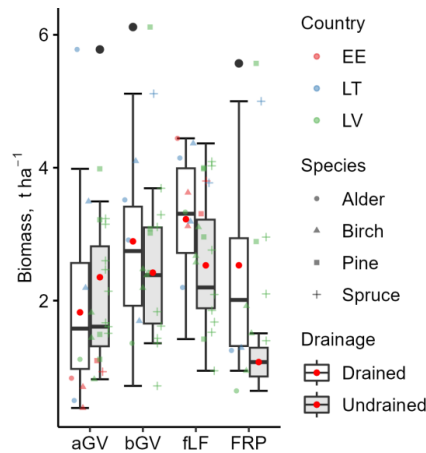
375 **Figure 6: Annualized total respiration (R_{tot}) in study sites stratified by drainage status and country (a) or dominant tree species (b).**
376 Error bars indicate confidence interval. A shared letter indicates that differences are not significant.

377 However, when categorizing data according to drainage status and dominant tree species, fewer differences were found in the
378 annualized R_{tot} than in the instantaneous R_{tot} (Figure 6b). For instance, among the drained sites, the lowest mean annual R_{tot}
379 was estimated for pine forests (5.23 ± 0.29 t CO₂-C ha⁻¹ year⁻¹), while in spruce, birch, and alder forests, the means were similar
380 ($p > 0.05$) (6.71 ± 0.31 t CO₂-C ha⁻¹ year⁻¹). Emissions from undrained soils in alder, birch, and spruce forests were similar to
381 each other and lower than from drained sites, ranging from 4.6 ± 0.71 in spruce forests to 5.47 ± 1.63 t CO₂-C ha⁻¹ year⁻¹ in alder
382 forests (overall mean 4.86 ± 0.71).

383 The correlation between R_{tot} and WTL was low; however, a drainage status (drainage ditch presence) impact on R_{tot} is
384 indicated by the PCA results, where undrained sites tend to have more similar characteristics while drained sites show greater
385 diversity concerning R_{tot}. However, clear covariation of dominant tree species and R_{tot} are not recognized by PCA (Figure S
386 6 and Figure S 7). When comparing the chemical and physical properties of different soil layers with the estimated annual
387 R_{tot}, as well as the measured mean R_{het}, the mean measured R_{het} consistently shows a higher correlation with evaluated soil
388 parameters (Figure S 8). The only exception is Corg, where in all correlation combinations, it was not present (r around -0.1).
389 Excluding Corg, the other soil chemical parameters generally have a low to moderate correlation (mean $r = 0.4$) with respiration.
390 The highest correlation is with pH, K, Mg, and P (mean $r = 0.5 \pm 0.07$, $p < 0.05$), and it is consistent across all evaluated soil
391 layers, while the correlation with BD (mean $r = -0.2$, $p > 0.05$) tends to increase with deeper soil layers reaching the highest
392 correlation ($r = -0.3$) in layer 20-30 cm. In addition, a higher C:N ratio is associated with lower CO₂ emissions (mean $r = -0.4$,
393 $p < 0.05$).

394 3.4 Annual litter inputs

395 The estimated mean litter inputs in both drained and undrained sites were mostly similar (Table S 10 and S11), typically
396 differing by less than 20%. Only fLF and FRP tended to be considerably higher in the drained sites, FRP on average even more
397 than twice as high. Compared to undrained sites, bGV in drained sites was about 20% higher on average, while aGV was about
398 20% lower on average (Figure 7). However, regardless of the soil drainage status, the proportion of aGV in the total GV
399 biomass was $54 \pm 18\%$ (Table 3).



400

401

402 **Figure 7: Variation in the biomass components used as litter input estimates.** Abbreviations: aGV and bGV – above- and belowground
403 biomass of herbaceous vegetation, fLF – fine foliar litter (needles, leaves, fine woody litter), FRP – tree fine root production. The bottom
404 and top edges of the box represent the 25th and 75th percentiles, summarizing the interquartile range (IQR). The whiskers extend to the
405 smallest and largest values within $1.5 \times \text{IQR}$ from the 25th and 75th percentiles, respectively. Black dots mark outliers. A red dot and a solid
406 horizontal line in the box indicate mean and median values, respectively.

407 **Table 3: Biomass (mean \pm CI, t dm. ha⁻¹) components used as litter input estimates, stratified by drainage status.** Abbreviations: aGV
408 and bGV – above- and belowground biomass of herbaceous vegetation; respectively; FRP – tree fine root production; fLF – fine foliar litter
409 (needles, leaves, fine woody litter).

Category	Drained	Undrained
aGV	1.82 \pm 0.52	2.35 \pm 1.61
bGV	2.89 \pm 0.85	2.42 \pm 0.84
FRP	2.53 \pm 0.77	1.08 \pm 0.57
fLF	3.22 \pm 0.44	2.53 \pm 1.06

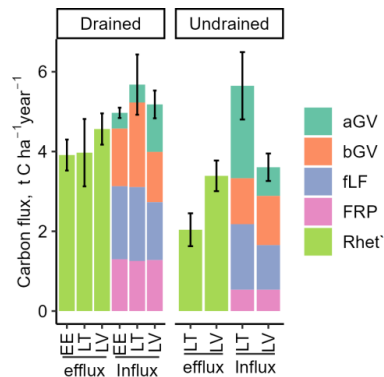
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411 Both bGV ($r=0.6$) and FRP ($r=0.7$) had a significant negative correlation with soil pH but a positive with the C:N ratio in
412 soil layer 0-30 cm. Additionally, FRP had a significant negative correlation ($r=0.7$) with the contents of N, Ca, and Mg in the
413 soil. No explanatory factors for aGV could be identified. Moderate correlation ($r=0.5$, $p<0.05$) was found between stand age
414 and fLF.

415 3.5 Annual soil carbon balance

416 The estimated R_{het} (Table S 10) proportion of R_{tot} varied between 54 and 71% (mean 65%). Consequently, the estimated
417 annual gross C losses from drained soils in the form of R_{het} emissions ranged from 2.36 to 7.49 t CO₂-C ha⁻¹ year⁻¹
418 (mean 4.30 \pm 1.20), while for undrained soils the range was from 1.63 to 4.68 t CO₂-C ha⁻¹ year⁻¹ (mean 3.00 \pm 0.99). According
419 to the RMSE of the model (Equation 2, Figure S 2) prediction, the R_{tot} to R_{het} calculation introduced an uncertainty of
420 approximately 0.32 t CO₂-C ha⁻¹ year⁻¹. In drained and undrained sites, the total estimated C input ranged from 3.81 to
421 7.03 t C ha⁻¹ year⁻¹ (mean 5.20 \pm 0.91) and 2.89 to 5.98 t C ha⁻¹ year⁻¹ (mean 4.19 \pm 1.10), respectively (Figure 8). The
422 uncertainties (relative CI) in C efflux and influx in both drained and undrained soils were relatively uniform, with a mean
423 uncertainty of 36 \pm 14% for the estimated individual annual C fluxes. The largest source of error was C influx in undrained soils
424 (49 \pm 13%), while the uncertainty of individual C fluxes in drained sites averaged 27 \pm 6% (Figure 8).

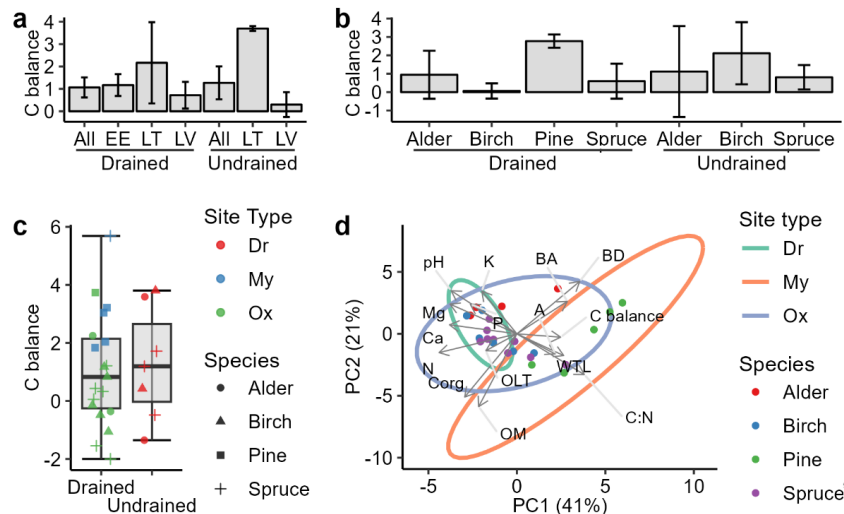
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426

427 **Figure 8: Components of the estimated soil carbon balance (sum±combined CI).** Efflux is soil heterotrophic respiration (Rhet')
428 calculated from Rtot, and influx is the estimated litter input; both are expressed as C. Abbreviations: aGV and bGV – above- and belowground
429 biomass of herbaceous vegetation; respectively; fLF – fine foliar litter; FRP – tree fine root production.

430 The mean soil C balance during the study period was 1.06 ± 0.45 and 1.27 ± 0.73 t C ha⁻¹ year⁻¹ for the drained and undrained
431 sites, respectively (Figure 9a). These results indicate that long-term drainage reduced soil C sequestration capacity by an
432 average of 0.20 t C ha⁻¹ year⁻¹. The higher mean soil C removals estimated for both drained and undrained sites in Lithuania,
433 compared to those in Estonia and Latvia, are primarily attributed to the greater C input from GV litter (Figure 8). However,
434 the sites in Lithuania also stand out due to greater uncertainty in both C input and output, resulting in the soil C balance in
435 drained sites across the countries being equivalent within the margin of error (Figure 9a). The low number of undrained sites
436 in Lithuania (n=2) limited the ability to investigate the patterns behind the observed lower soil CO₂ emissions and higher aGV
437 litter, contributing to significantly higher soil C removals (3.70 ± 0.11 t C ha⁻¹ year⁻¹) compared to sites in Latvia
438 (0.30 ± 0.56 t C ha⁻¹ year⁻¹, n=5).



439

440 **Figure 9: Soil carbon balance (a, b, c: mean t C ha⁻¹ year⁻¹ ± SE) and impacting factors (d: PCA biplot).** Abbreviations: A – stand age;
441 BA – basal area, BD – bulk density; C:N – ratio between organic carbon and nitrogen in soil; OLT – soil organic layer thickness; WTL –
442 water table level; pH – soil pH value; K, Ca, Mg, P, OM, Corg, N represent the content of potassium, calcium, magnesium, phosphorus,
443 organic matter, organic carbon and nitrogen in the top 0-30 cm layer of soil, respectively. Site types: Dr - Dryopterioso-caricosa (undrained
444 sites); Ox - Oxalidosa turf. mel. (drained); My - Myrtilliosa turf. mel. (drained).

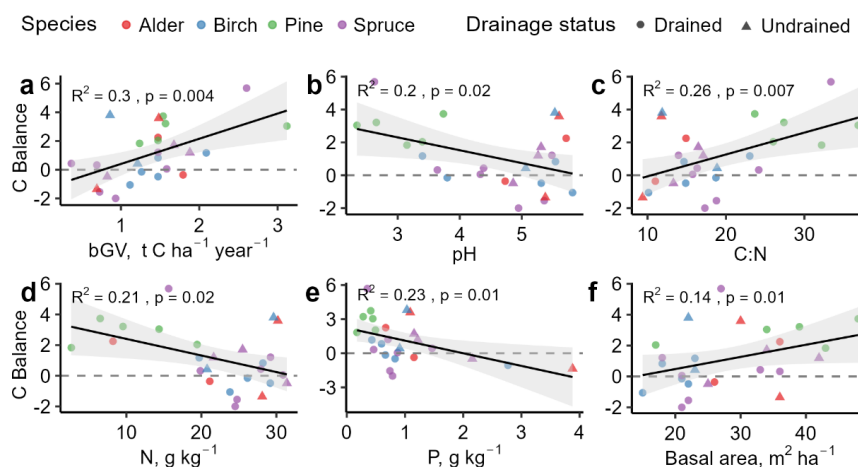
445 The soil C balance stratified by dominant tree species (Figure 9, b) indicates that the drained sites were skewed towards C
446 removals due to significantly higher C removals in pine stands. Pine sites (n=5) showed high C removals with low uncertainty



(mean $2.77 \pm 0.36 \text{ t C ha}^{-1} \text{ year}^{-1}$), in contrast to the other drained sites ($n=19$), where the mean soil C balance was estimated at $0.45 \pm 0.50 \text{ t C ha}^{-1} \text{ year}^{-1}$, suggesting that soils in drained alder, birch and spruce sites were in near C equilibrium during the study period. The soil C removals identified for undrained birch and spruce stands ($1.33 \pm 0.72 \text{ t C ha}^{-1} \text{ year}^{-1}$, $n=5$) were consistent, while in alder stands, the C balance of $1.12 \pm 2.47 \text{ t C ha}^{-1} \text{ year}^{-1}$ was highly uncertain due to the number of sites being just two.

A trend of higher soil C removals was observed in site types associated with relatively lower nutrient availability (Figure 9, c). In *Oxalidosa* sites, the mean soil C balance was $0.32 \pm 0.40 \text{ t C ha}^{-1} \text{ year}^{-1}$, while in *Myrtillosa* sites, it was $3.16 \pm 0.69 \text{ t C ha}^{-1} \text{ year}^{-1}$. Thus, drained nutrient-rich soils were approximately at C equilibrium, whereas comparably nutrient-poorer soils acted as a C sink. The observed tendency is supported by PCA (Figure 9d) which indicates that higher soil C removals are associated with lower nutrient concentrations and pH levels. In addition, the PCA reveals that the risk of soil C loss is reduced in stands with higher basal area and age, as well as with a higher soil C:N ratio, which in our study likely reflects the variability in peat quality and decomposability under different vegetation types.

According to correlation analysis, soil parameters such as pH, C:N ratio, N, and P showed the strongest correlations with the soil C balance (Figure 10). Basal area was the stand characteristic with the strongest correlation with C balance, while, among the C flux components, bGV demonstrated the most consistent role in C balance. No meaningful relationship was identified between soil C balance and soil organic matter or C content, nor with the depth of the WTL or organic layer thickness.



463

464 **Figure 10: Relationships between soil C balance ($\text{t C ha}^{-1} \text{ year}^{-1}$) and impacting factors.** The figures include impacting factors
465 showing the highest correlations found in the study dataset.

Soil nutrient availability as C balance affecting factor is confirmed by RDA and PLSRS models. RDA and PLSRS models ($p < 0.05$) explained 78% and 70% of the soil C variance, respectively. After excluding variables introducing multicollinearity, the models included WTL, organic layer thickness, pH, N, K, Ca, Mg, P, organic matter content of the soil, stand age, and basal area. The variables pH, K, and Mg showed significant contributions ($p < 0.05$) in the RDA model explaining the variation of soil C balance. The variable N was not significant ($p = 0.087$) but was close to the threshold of significance. PLSR model indicated that pH, K, Mg and N explained 45% of the C balance variance. Although the VIP values for all potential explanatory variables were below 0.4, suggesting limited predictive power for C balance with the current dataset.



473 4 Discussion

474 4.1 Soil carbon balance

475 The soil C balance of the studied drained and undrained organic forest soils fluctuated around equilibrium, demonstrating both
476 C sink and source dynamics. The reason for the uncertain C balance in undrained alder stands could not be determined, as site
477 characteristics were consistent with the patterns observed in other undrained sites showing soil C removals. However, we
478 identified soil properties as the likely reasons being why the soil in drained pine stands showed C removals, in contrast to the
479 C-neutral soil observed at other drained sites.

480 Although observing a C sink in drained nutrient-rich soils may seem somewhat unexpected, given that these soils have
481 generally been estimated to on average act as a net C sources in both boreal and temperate zones (Jauhiainen et al. 2023), it is
482 not entirely novel. Both soil C sinks and sources have also been observed in earlier studies under a wide range of site conditions
483 (e.g., Ojanen et al. 2013; Minkkinen et al., 2018; Bjarnadottir et al., 2021; Hermans et al., 2022). Many of the soil C sink sites
484 may be classified as nutrient-poor, but not all (e.g., Ojanen et al. 2013). In soil inventory studies carried out in Latvia, the soil
485 C stocks of forestry-drained peatlands were found to be stable in all but the most nutrient-rich soil conditions, under which the
486 C stock was reduced in the long term (Lazdiņš et al., 2024; Dubra et al., 2023). However, interpretation of soil C balances
487 solely based on the nutrient status of a forest site should be approached with caution, as it is typically derived from indirect
488 indicators such as vegetation or stand productivity, rather than a quantitative assessment of soil nutrient concentrations.
489 Consequently, sites with variable conditions may be classified under a given nutrient status category. Similarly, drainage status
490 does not guarantee specific WTL levels (Figure 4). This may be the reason for varying findings on soil C balances across
491 studies that allegedly target the same soil type and drainage status, i.e. the category of drained, nutrient-rich soils may be too
492 broad, encompassing varying nutrient and moisture regimes, which prevents the expectation of similar C balances, especially
493 in different climates. In our study, this aspect was evident, as the soil C balance, regardless of drainage status, appeared to be
494 influenced by the composition and relative proportions of the dominant tree species in combination with variation in soil
495 nutrient conditions and organic matter quality, as indicated by the C:N ratio, across the study sites.

496 The soils at our study sites represented a wide range, from highly mineralized soils close to the threshold of organic soil
497 definition (Hiraishi et al., 2013) to deep peat (Figure 2). Similarly, the WTLs varied widely, with sites ranging from an average
498 WTL close to the soil surface to depths exceeding one meter (Figure 4). However, no meaningful relationship was identified
499 between WTL variation and soil CO₂ emissions, nor between mean WTL and soil C balance. Also, soil C and organic matter
500 contents, or the depth of the organic layer were poor predictors of soil C balance. Probably the importance of these factors
501 decreases over time since initial disturbance by drainage system implementation. Rhet has been found to decrease over time
502 following drainage (Qiu et al., 2021), except for the initially wettest, waterlogged hollow surfaces (Munir et al., 2017). There
503 are few long-term monitoring or chronosequence studies, but those suggest that the most intensive period of soil C loss
504 following drainage is the first decade/decades (Hargreaves et al. 2003; Vanguelova et al., 2019). During that period, also peat
505 subsidence following drainage is highest (Lukkala 1949), and labile substances in peat that due to lowering WTL and peat
506 subsidence becomes exposed to oxic decomposition are largely lost, leaving more decomposition-resistant substrates behind
507 (e.g., Jayasekara et al., 2025). Simultaneously, major changes in litter inputs and their decomposability take place (Straková
508 et al., 2010, 2012). Considering the long period since drainage of our sites, dating back to around a century ago and potentially
509 even to the mid-19th century (Zālītis, 2012), the initially high soil C loss due to drainage-induced increase in gross soil CO₂
510 emissions has likely been offset by enhanced biomass growth and the resulting increased litter inputs (Hommeltenberg et al.,
511 2014). However, a comparison of soil C balances between drained and undrained sites still shows a negative impact of
512 historical drainage (Table S 11). To more accurately assess the impact of drainage on soil C stock changes and its evolution in
513 time, long-term studies would be required.



Soil nutrient conditions explained the observed C balances better than WTL and soil C characteristics did. In our study, drained nutrient-rich soils were represented by Oxalis (*Oxalidosa turf. mel.*) and Myrtillosa (*Myrtillosa turf. mel.*) site types. All our drained birch and alder sites and most of drained spruce sites belonged to Oxalis site type, but most of the pine sites belonged to the Myrtillosa site type. While the soil C balance under the other species in drained sites was practically neutral during the study period, pine stands showed relatively high C removals. In the Myrtillosa site type nutrient concentrations were, on average, 1.5 to 5.4 times lower compared to Oxalis sites. In addition, Myrtillosa sites had significantly lower pH levels and a higher C:N ratio (Figure 2). The distinctive C balance patterns between the site types suggest that, in addition to lower nutrient concentrations, increased soil acidity and differences in soil organic matter and litter input quality (decomposability) had a role in the observed C removals in these sites. Previous studies have also reported a negative correlation between pH and soil C content (Zhou et al., 2019) and have linked soil acidification with increasing soil C stocks (Madsen et al., 2025; Marinos and Bernhardt, 2018). The relevance of soil chemical parameters in determining soil C balance was supported by RDA and PLSR analysis, revealing that soil pH and macronutrient concentrations were key parameters determining soil C balance.

Among the drained sites, soil in pine stands had the lowest mean CO₂ emissions (Figure 6b). Additionally, nutrient availability correlated negatively with belowground biomass (bGV, FRP) confirming previous observations that greater belowground biomass is associated with reduced nutrient availability (Zhang et al., 2024). Higher CO₂ emissions observed in Oxalis sites were consistent with the previous observations of higher organic matter decomposition rates typically observed in sites with high nutrient availability (Shahbaz et al., 2022; Hiraishi et al., 2013). However, an increased total soil C influx from litter was also observed in these sites during the study period, effectively offsetting soil C loss from Rhet. Another potential contributor to the stability of C stocks in drained soils was soil compaction induced by drainage, as the increased BD of drained soil was found to be associated with lower Rhet emissions. The reason may be reduced soil porosity limiting gas exchange between the soil and the atmosphere (Ball, 2013; Novara et al., 2012). Thus, while drained soil may be prone to a higher decomposition rate, flux-driving processes seem to be countered by increased soil compaction.

In any case, the mean C removals we estimated are uncertain and remain indicative, preventing a definitive conclusion that the soil functions as a C sink. The uncertainty in the results arises from the variety of included study sites. The results obtained reflect C balances only for the study period, specific to the respective stands in their specific developmental stages and site conditions. They do not represent average changes in soil C stocks over a longer timeframe, such as an entire forest management cycle. In similar conditions, organic soils have been found to result in C stock loss following regenerative felling (Korkiakoski et al., 2023; Butlers et al., 2022). Therefore, if the impact of management practices were considered, the soil C balance would likely shift towards reduced C sequestration.

4.2 Total respiration

No significant differences in R_{tot} were observed between the countries, likely because the gradient in the mean air temperature from Estonia to Lithuania, ranging from 6.4 to 8.5 °C, was not substantial enough to introduce distinguishable differences. Nevertheless, while temperature is generally recognized as a strong factor influencing soil respiration and its variation, it should not be regarded as the sole predictor of respiration. Relationships observed in one region may not directly apply to another, as differences in soil moisture (Jovani-Sancho et al., 2018) or nutrient status, as discussed earlier, can significantly alter the respiration dynamics. Similarly, no clear impact of dominant tree species on R_{tot} was found. This points to a minor role of dominant tree species on emissions. However, there is some evidence that emissions in undrained sites tended to be higher in deciduous stands, particularly alder stands, according to the measured instantaneous emissions. The enhanced soil CO₂ efflux observed in the presence of alder can probably be attributed to the symbiotic nitrogen fixation associated with these trees (Warlo et al., 2019), which increases nitrogen availability in the soil. Nitrogen availability, in turn, can stimulate decomposition processes, leading to a higher rate of CO₂ release. However, we did not observe increased nitrogen levels in



neither the soil nor soil water of alder sites. Although statistically unconfirmed, a tendency can be noticed that in drained sites Rtot emissions tend to be higher in birch stands, but lower in pine forests. Also, previous studies indicated that deciduous stands show higher CO₂ emissions (Jauhiainen et al., 2023).

While it was found that both drainage status and WTL threshold above or below 30 cm can be used as a predictor of Rtot, a meaningful correlation between WTL and Rtot was not found. Furthermore, although the absolute variation of the WTL was higher in drained sites, the relative variation in both WTL level and Rtot was indifferent to the drainage status. The observations suggest that higher WTL conditions in undrained sites, while decreasing Rtot emissions, do not guarantee higher resilience to moisture regime disturbances, i.e., more stable emissions. The main reason is that the presence of drainage ditches is not the only factor constraining WTL both spatially and temporally, and in undrained sites too, WTL frequently falls below 30 cm (Butlers et al., 2023) ensuring oxic conditions in soil layers containing labile organic matter. Furthermore, this typically happens in summer (Butlers et al., 2023) when increased temperatures further promote organic matter mineralization. The role of WTL dynamics is reflected also in PCA, showing higher dispersion of drained sites likely due to higher variation in WTL depths. This may be the reason complicating the quantification of relationships between Rtot and the affecting factors, especially in drained sites. The wide range of mean WTL measured at the drained sites also helps explain why Rtot at these sites is not necessarily significantly higher compared to undrained sites.

To achieve accurate Rtot annualization using data from periodic flux measurements, data interpolation through modelling approaches was applied. For Rtot interpolation we compared nonlinear models and linear models after logarithmic or Box-Cox transformation. Both the advantages and shortcomings of these data transformation methods and modelling approaches have been reported in previous studies. (Yueqian, 2020; Wutzler et al., 2020; Liaw et al., 2021; Moulin et al., 2014; Box and Cox, 1964; Khomik et al., 2009). Although the bias in predicted annual Rtot varied among study sites, the overall impact of different flux modelling approaches on estimated mean annual Rtot of drained and undrained sites was minimal. Specifically, the mean bias of results obtained through the implementation of the Box-Cox transformation compared to other approaches was $-2\pm 9\%$. Thus, while such an impact has been observed in previous studies, the skewing of results due to the annualization of respiration was not identified in our study.

4.3 Soil heterotrophic respiration

We used a Rhet value derived from $R_{tot} - R_{het'}$ as the soil C efflux. Rhet' was derived from Rtot empirically using the Rhet/Rs data from a large dataset (Jian et al. 2021), thereby reducing the effect of potential systematic and random errors in individual studies. Such empirical derivation of Rhet has been acknowledged to be an applicable approach and has been utilized in previous studies (Jauhiainen et al., 2019, 2023). To elaborate the recalculation model we used Rhet and Rs values from the database (Jian, J. et al., 2021) on forest soil flux in the boreal zone, as existing experience suggests that organic soil emissions in hemiboreal forests are more likely to align with boreal rather than temperate conditions (Krasnova et al., 2019; Heikkinen et al., 2023; Bårdule et al., 2022; Butlers et al., 2022; Dubra et al., 2023; Lazdiņš et al., 2024; Jauhiainen et al., 2023). The choice of using only boreal data tends towards estimating higher Rhet', compared to the use of temperate data, as illustrated in Figure S 2. This approach aimed to avoid the underestimation of soil C efflux. The mean share of Rhet acquired using boreal data was 0.65 ± 0.04 while using temperate data - 0.60 ± 0.15 , or around 10% difference. Accordingly, the Rhet proportion values we applied were higher than the typically observed range of 0.5 to 0.6 (Bond-Lamberty et al., 2004; Hanson et al., 2000), demonstrating that our approach avoided underestimating Rhet.

The role of ground vegetation autotrophic respiration in Rtot increases with its biomass (Munir et al., 2017). Therefore, the risk of underestimating Rhet by using Rhet' is further reduced because the Rhet/Rs ratio used to recalculate Rtot to Rhet' does not account for the impact of autotrophic respiration from aboveground vegetation, consequently, the approach tends to rather overestimate the Rhet. This aspect should be considered when assessing our results. When estimating the impact of historical



drainage on the soil C balance by comparing sites by drainage status, this bias was likely negligible, because the mean ground vegetation biomass did not significantly differ between drained and undrained sites ($\Delta=0.53 \text{ t dm. ha}^{-1}$).

The applicability of the approach is supported by the comparability of the estimated R_{het} values with R_{het} reported in previous studies (Bond-Lamberty and Thomson, 2010). We estimated R_{het} of drained soil to be mean $4.30 \pm 1.20 \text{ t CO}_2\text{-C ha}^{-1} \text{ year}^{-1}$, which is slightly higher than both mean R_{het} of forest organic soil found in the boreal zone ($4.09 \text{ t C CO}_2\text{-C ha}^{-1} \text{ year}^{-1}$) (Ojanen et al., 2010) and in a broader regional scale ($3.71 \pm 0.53 \text{ t CO}_2\text{-C ha}^{-1} \text{ year}^{-1}$) (Jian, J. et al., 2021). When attempting to correct for the approximated overestimation of R_{het} introduced by trenching (see the Supplementary text), the resulting mean R_{het} would be $2.4 \text{ t CO}_2\text{-C ha}^{-1} \text{ year}^{-1}$, which is considerably lower than the R_{het} values we used in the C balance estimates.

4.4 Carbon influx by litter inputs

In the estimation of C influx, we considered data only for fLF, aGV, bGV, and FRP, excluding cLF, MP and dwarf shrubs. This approach was chosen because the CO_2 emissions produced in decomposition of these litter types was directly included in the measured R_{tot} . For instance, cLF due to its dimensions and scarce coverage could not be objectively included in chamber measurements. Furthermore, while fLF is relatively uniform in forest areas, the coverage of mosses and dwarf shrubs is not always so, therefore it is necessary to know their area of projection to be included in the C balance estimation. One solution for incorporating the “missing” litter inputs would be to use modelling approaches (Alm et al., 2023). However, we did not attempt to include those litter sources in the soil C balance estimation, as doing so would have introduced additional uncertainty. We estimated that annual moss production was $22 \pm 10\%$ of the average total moss biomass of $0.50 \pm 0.09 \text{ kg dry matter (dm.) m}^{-2}$ measured for moss patches in our sites. Thus, mosses could potentially provide annual litter inputs reaching up to $0.98 \pm 0.25 \text{ t dm. ha}^{-1}$ if their cover was 100%. The mean annual cLF was $0.74 \pm 0.23 \text{ t dm. ha}^{-1}$. Therefore, the litter input estimates (mean $4.70 \pm 1.43 \text{ t C ha}^{-1} \text{ year}^{-1}$) used in the calculation of soil C balance likely led to overestimated soil C loss. This highlights the need to consider all litter input components in further studies, even though that may clearly increase the workload.

5. Conclusions

Although all soils in our study sites were classified as nutrient-rich based on forest site type taxonomy, they included a wide variety, ranging from those near the threshold of organic soil definition to soils with deep peat layers. Consequently, the soils exhibited broad variability in pH, macronutrient concentrations, and C:N ratio. That in turn contributed to the observed behaviour of the soils demonstrating both C sink and source dynamics under both drained and undrained conditions. During the study period, drained soils under birch, black alder, and Norway spruce remained C neutral, while in pine stands the soils were C sinks, presumably due to the significantly lower nutrient availability limiting mineralization of the organic matter. The disparity in soil nutrient conditions also explains why some undrained soils, characterized by relatively high nutrient availability, acted as C sources. These findings highlight the potential to improve predictions of soil C balance by complementing the broad “nutrient-rich” soil classification - typically assigned using site vegetation as a proxy - with quantitative measurements of soil nutrient status. The results can be used as input for constructing emission factors or more elaborate forest organic soil CO_2 emission prediction methods. Additional research is necessary to expand the dataset for establishing robust quantitative relationships that can be used to reliably identify and predict whether organic soils function as C sinks, sources, or remain in neutral balance, depending on site-specific conditions and annual weather variations.



634 **Data availability**

635 Data used for carbon balance estimations is available at <https://doi.org/10.5281/zenodo.14968843>

636 **Author contributions**

637 KS, JJ, RL, AL and KA developed a harmonized methodology. ABu, DČ, TS and MKS managed and processed the study
638 data. ABu wrote the original manuscript, with significant reviewing contributions from RL, JJ, TS, ABā, IL, VS, HV, IL, AH
639 and AJ provided critical reviews and edits to the manuscript.

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