



- 1 Water chemistry and greenhouse gas concentrations in waterbodies of a thawing permafrost
- 2 peatland complex in northern Norway
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12 13	Abstract Thermokarst ponds in thawing permafrost landscapes play a considerable role in greenhouse
14	gas (GHG) emissions despite their small size, yet they remain underrepresented in Earth
15	system models. At the Iškoras site in northern Norway, a peat plateau with decaying
16	permafrost and thermokarst ponds adjacent to a wetland, we studied water chemistry,
17	dissolved organic matter (DOM) processing, and GHG fluxes over two years. Thermokarst
18	ponds exhibited low pH, high organic acidity, and high oversaturation of dissolved carbon
19	dioxide (CO <sub>2</sub> ) and especially high dissolved methane (CH <sub>4</sub> ). Adjacent wetland streams,
20	however, with near-neutral pH, showed lower CH4 and organic acidity but significantly
21	higher CO <sub>2</sub> emissions despite moderate saturations driven by turbulence and bicarbonate
22	replenishment. By contrast, CO <sub>2</sub> emissions in ponds were primarily linked to DOM
23	mineralization.
24	DOM mineralization rates were similar between ponds and streams, suggesting that
25	environmental factors like pH and microbial community differences counteract DOM lability
26	variations. As permafrost decays and transitions from peat plateaus to wetlands, ponds as
27	hotspots of CH4 emissions will disappear. However, total GHG fluxes across the peatland-
28	wetland continuum will depend on wetland emissions, where CH4 emissions usually are
29	considerable, and the fate of organic matter within the plateau. Lateral DOC fluxes may
30	represent a significant loss of soil organic carbon, highlighting the importance of
31	hydrological connectivity in linking terrestrial and aquatic systems. This study emphasizes
32	the need to account for the relationship between hydrological and chemical processes when
33	assessing C and GHG fluxes in permafrost-impacted regions.





35	Northern latitude regions, which store approximately 1,300 Pg of organic carbon (OC)
36	(Hugelius et al., 2014), represent one of the largest terrestrial carbon reservoirs on Earth
37	(Schuur et al., 2008; Schuur et al., 2015; Walter et al., 2006). Sequestered under cold and
38	oxygen-limited conditions, this carbon (C) is increasingly vulnerable to release as permafrost
39	thaws due to climate warming, generating significant feedbacks that complicate predictions
40	of future climate trajectories (Schuur et al., 2008; Schuur et al., 2015; Walter et al., 2006). As
41	permafrost degrades, the release of greenhouse gases, particularly methane (CH4) and carbon
42	dioxide (CO <sub>2</sub> ), through the microbial decomposition of previously frozen organic matter
43	(OM) can rapidly escalate the impact of this feedback (Schuur et al., 2008; Walter et al.,
44	2008; Wik et al., 2016; Zimov et al., 2006). While the large-scale thaw of permafrost is
45	widely recognized (Leppiniemi et al., 2023), the timing, magnitude, and pathways of carbon
46	release remain uncertain, influenced by processes such as burial, mobilization, lateral export,
47	and mineralization (Verdonen et al., 2023; Vonk et al., 2015).
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59	mirrors processes observed across the northern hemisphere, including in the Canadian Arctic,
59	
60	European Russia, and the Kola Peninsula, highlighting the vulnerability of sporadic
61	permafrost regions to warming climates (Krutskikh et al., 2023; Payette et al., 2004; Sannel
62	and Kuhry, 2011). While the processes driving these transformations, such as thermal
63	disturbances, vegetation shifts, and subsidence, are relatively well-studied, their
64	consequences for GHG fluxes and C cycling remain uncertain, limiting our ability to project
65	future climate feedbacks (Holmes et al., 2022; Olefeldt et al., 2021; Turetsky et al., 2020).
66	Among the new landscape forms that emerge from degrading peat plateaus, thermokarst
67	ponds and wetlands play a critical role in greenhouse gas dynamics. These small aquatic
68	systems, formed by the thaw and collapse of permafrost, are characterized by high
69	concentrations of dissolved organic carbon (DOC) and inorganic carbon (DIC) (Abnizova et
70	al., 2012; Martin et al., 2021; Matveev et al., 2018). Thermokarst ponds, in particular, act as
71	hotspots for CH4 and CO2 emissions due to their unique biogeochemical conditions,
72	including hydrological isolation, anoxic sediments, and high organic matter availability (In 't
73	Zandt et al., 2020; Polishchuk et al., 2018; Vonk et al., 2015; Ward and Cory, 2015). Despite
74	their small size, these water bodies can contribute significantly to regional C fluxes, with CH4
75	and CO2 supersaturation levels often surpassing those of larger lakes or surrounding tundra
76	ecosystems (Abnizova et al., 2012; Kuhn et al., 2018; Shirokova et al., 2012). However, their
77	contributions are often overlooked in large-scale C assessments, as their small size makes
78	them difficult to detect using satellite-based methods (Holgerson and Raymond, 2016; Muster
79	et al., 2017).
80	As permafrost that progresses the transition of isolated thermokarst ponds to interconnected

As permafrost thaw progresses, the transition of isolated thermokarst ponds to interconnected
wetland systems further alters GHG dynamics. While northern permafrost wetlands currently
act as a C sink, the inclusion of thaw pond emissions into broader wetland carbon budgets
reveals their potential to offset the sink capacity by 39% (Kuhn et al., 2018). Compared to





84	thermokarst ponds, wetlands have sustained CH4 fluxes over larger areas due to persistent
85	waterlogging and OM decomposition (Pirk et al., 2024; Swindles et al., 2015; Turetsky et al.,
86	2020), thus constituting important long term CH <sub>4</sub> sources (Bansal et al., 2023). The
87	transformation from stable permafrost to thermokarst landscapes is accompanied by shifts in
88	hydrology, OM lability, and microbial activity, which collectively shape CO2 and CH4
89	production pathways (Holmes et al., 2022; Laurion et al., 2020). Understanding the dynamics
90	of these evolving systems is critical for assessing the broader impacts of permafrost thaw on
91	regional C uptake and emissions as well as global C cycles.
92	Northern Norway's sporadic permafrost zone, with its abundant small thermokarst ponds and
93	emerging wetlands, provides a valuable opportunity to study these processes. The region's
94	rapidly degrading peat plateaus host significant C stocks, yet small aquatic systems,
95	especially those in Fennoscandia, remain underrepresented in Earth system models
96	(Abnizova et al., 2012; Muster et al., 2019; Muster et al., 2017). Existing studies emphasize
97	the importance of quantifying CH <sub>4</sub> and CO <sub>2</sub> fluxes in these environments and their
98	implications for C budgets (Abnizova et al., 2012; Matveev et al., 2018). However, critical
99	questions remain regarding how transitions between permafrost, thermokarst, and wetland
100	systems influence C dynamics, and whether these landscapes function as net C sources or
101	sinks under changing climatic conditions (Sim et al., 2021).
102	This study aims to address these gaps by examining the GHG dynamics and C
103	biogeochemistry of thermokarst ponds and wetland streams in the sporadic permafrost zone
104	of northern Norway. Over two years, we measured dissolved CO2 and CH4 concentrations,
105	water chemistry, and OM lability to evaluate the processes driving C fluxes in these systems.
106	We hypothesize that (1) thermokarst ponds serve as hotspots of $CH_4$ and $CO_2$ production
107	relative to wetland streams, (2) the transition from isolated ponds to wetlands significantly
108	alters GHG emission pathways, driven by shifts in hydrology and OC availability, and (3)





- 109 recently mobilized OM presents a labile source of C promoting CO<sub>2</sub> production in
- 110 thermokarst water bodies compared to wetland streams. By exploring these dynamics, this
- study provides insights into the role of small water bodies in permafrost C feedbacks,
- advancing our understanding of sub-Arctic and boreal C cycling.

113





- 115 2. Methods
- 116 2.1 Study area
- 117 The Iškoras field site (69.34°N, 25.29°E; 381 m a.s.l.) is a permafrost peatland plateau
- 118 located in the interior of the Finnmark province, northern Norway, on the Finnmarksvidda
- 119 plateau (Fig. 1). The region of Finnmarksvidda lies between 300 and 500 m a.s.l. and is
- 120 characterized by a subarctic continental climate. The topography was shaped by Pleistocene
- 121 glaciations, which deposited ground moraines, glaciofluvial, and glaciolacustrine sediments

122 (Sollid et al., 1973). The depressions in the landscape are commonly filled with peatlands

123 (Borge et al., 2017), and peat plateaus underlain by permafrost are common.

124 The Iškoras peat plateau covers an area of approximately 4 ha and is part of a 3.3 km<sup>2</sup>

- subarctic headwater catchment that drains into the Báhkiljohka river (91 km<sup>2</sup>). Mean annual
- 126 air temperature and precipitation for the 30-year normal (1991-2020) period was  $-1.9^{\circ}C \pm$
- 127  $1.0^{\circ}$ C, and  $513 \pm 90$  mm, respectively (Table 1). For our study period 2021 to 2022, MAAT

and MAP were -1.1°C  $\pm$  0.4°C, and 589.5  $\pm$  62.5 (SeNorge, 2023). Iškoras lies within the

20 zone of sporadic permafrost and the peat soils extend down to about 1.5 m in the plateau

areas (Kjellman et al., 2018) and active layers depths up to 90 cm. The plateau exhibits a

131 complex surface of intact and degrading palsas, along with thermokarst ponds, and is

surrounded by wetlands and a stream to the northwest (Martin et al., 2019). Between 2019

and 2022, up to 0.8 m of subsidence of palsas was measured at localized sites (Pirk et al.,

134 2024). The site is located about 90 km south of the nearest coastal fjord and is dominated by

135 mountain birch forest (Betula pubescens) and tundra vegetation, including dwarf birch (B.

136 *nana*). The plateau consists primarily of low heath shrubs, Ericaceae (*Empetrum nigrum*,

137 Rhododendron tomentosum), lichen crusts, mosses, and cloudberry (Rubus chamaemorus) or

- 138 bare ground, while the surrounding wetlands are dominated by Sphagnum mosses, sedges
- 139 (*Carex* spp.), and cotton grass (*Eriophorum* spp.) (Kjellman et al., 2018; Martin et al., 2019).





	Unit	Mean ± std for 1991-2020	Mean + std for 2021-2022
annual temperature	°C	$-1.9 \pm 1.0$	$-1.1 \pm 0.4$
summer temperature	°C	$10.4 \pm 2.2$	$11.8 \pm 0.2$
annual precipitation	mm	513 ± 90	589.5 ± 62.5
summer precipitation	mm	196 ± 53	207 ± 48

140 Table 1 Mean and standard deviation (std) of climate parameters for the Iškoras catchment for the

142 September.

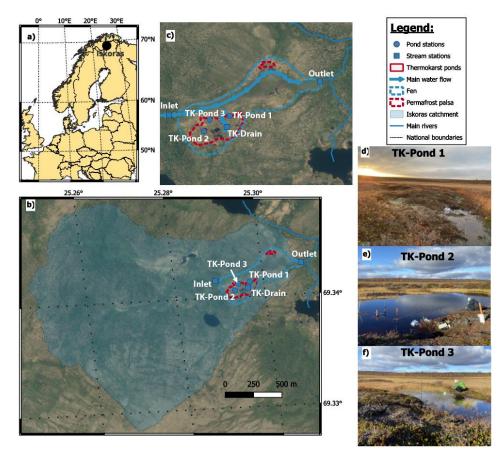
144	The study area included water bodies within a peat plateau and the adjacent wetland, selected
145	for sampling and monitoring. Measurements and samples were taken approximately monthly
146	in the ice-free season from May until October in 2021 and 2022. The waterbodies consisted
147	of three thermokarst ponds (TK-Pond 1, 2, and 3), a seasonal drainage channel (TK-Drain)
148	connecting the peat plateau to the wetland, and the wetland's inlet and outlet streams (Inlet,
149	Outlet), with the outlet also marking the terminus of the Iškoras catchment. From mid-May to
150	early November, monitoring showed that the thermokarst ponds and Inlet were ice-free for
151	$170 \pm 5$ days, while the Outlet remained ice-free for 184 days (Table SI 1).
152	The thermokarst ponds varied in hydrological connectivity and permafrost influence,
153	reflecting differences in age and physical characteristics. TK-Pond 1 (0.4 m depth) is small
154	and located at the peat plateau-wetland transition, experiencing periodic hydrological
155	isolation. TK-Pond 2, the largest and deepest (1.5 m depth), lies centrally on the peat plateau,
156	surrounded by degrading palsas. TK-Pond 3 (0.6 m depth) is situated at the plateau's edge and
157	was initially isolated by a surrounding permafrost mound.
158	The TK-Drain is a shallow, ephemeral drainage channel that provides the primary
159	hydrological connection between the peat plateau and wetland. The wetland's inlet stream
160	(0.6 m wide, 15–40 cm deep) begins approximately 200 m upstream of the plateau, flowing
161	through birch forest and mires without permafrost before entering the wetland where the

<sup>141</sup> normal period (1991–2020) and the study period 2021-2022. Summer is defined as May to





- 162 channel becomes less defined. The outlet stream (0.8 m wide, 30–60 cm deep) re-emerges
- approximately 700 m downstream at the wetland's far end, serving as the catchment outlet.
- 164 Between September 2020 and October 2022, a total of nine field campaigns were conducted
- 165 for regular sampling of water chemistry, dissolved gases, CO<sub>2</sub> emissions, dark incubations
- and high-frequency monitoring of water height and temperature.



- 168 Figure 1 Location map of the study area in Europe (a) and the Iškoras catchment (b; determined
- 169 from the outlet station Outlet). Close-up of the wetland (c) with regular sampling sites and main
- 170 water flow direction. Pictures of the three main pond sites are also shown (d–f).

2.2 Water chemistry





1/1	2.2 Water chemistry
172	Water samples for chemical analyses were collected using standardized procedures. From
173	each site, 500 mL unfiltered water was collected in HDPE rectangular bottles (Emballator
174	Melledrud AB, Stockholm, Sweden) after rinsing with sample waters three times, kept dark
175	after sampling, carried out of the field, and stored within hours after sampling at 4°C. The
176	samples were then transported by car and plane back to the laboratory and delivered for
177	chemical analysis where they were kept at 4°C until analysis. Chemical analysis of pH,
178	electrical conductivity (EC), alkalinity, sulfate (SO <sub>4</sub> ), silica (SiO <sub>2</sub> ), ammonium (NH <sub>4</sub> ), nitrate
179	(NO <sub>3</sub> ), total phosphorous (totP), total organic carbon (TOC), DOC (filtered by 0.45 $\mu m)$ and
180	particulate organic carbon (POC) (filtered and then combusted at $1800$ °C) in the water
181	samples was performed at accredited laboratories at the Norwegian Institute for Water
182	Research (NIVA); methods for analysis and quality control are described in Vogt and
183	Skancke (2022). The samples were not fully digested according to standard procedures
184	required for the determination of total nitrogen (totN), hence totN values are expected to be
185	underestimated and are therefore not shown in the manuscript (Thrane et al., 2020).
186	Absorption spectra of DOM were measured at NIVA for wavelengths between 200 and 900
187	nm, using 1 nm intervals, with a 5 cm cuvette length and Milli-Q water as a reference, using a
188	Lambda 40 UV/Vis spectrophotometer (Perkin Elmer, USA) and expressed in absorbance pr
189	cm. In two samples, incomplete filtration caused excess scattering, and these spectra were
190	removed. The absorbencies were used to calculate specific UV absorbency (sUVa =
191	$A_{\lambda 254nm}/mg \ C \ L^{-1}$ ) and the specific UV absorption ratio (SAR = $A_{\lambda 254nm}/A_{\lambda 400nm}$ ).
192	
193	2.3 Dissolved gas analysis
194	Dissolved gases (CO <sub>2</sub> CH <sub>4</sub> ) were sampled in the field using the acidified headspace

- 194 Dissolved gases ( $CO_2$ ,  $CH_4$ ) were sampled in the field using the acidified headspace
- 195 technique (Åberg and Wallin, 2014). Duplicate gas samples were collected according to





196	Valiente et al. (2022). Two 50 mL syringes were filled and sealed underwater without air
197	bubbles to prevent gas loss. Excess water was expelled to retain 30 mL, and 20 mL of
198	ambient air was drawn in to create a headspace. Samples were acidified with 0.6 mL of 3%
199	HCl to achieve pH <2, ensuring DIC was present as CO <sub>2</sub> . Equilibrium was reached by
200	shaking for one minute, followed by a 30-second rest, repeated thrice. Fifteen mL of
201	headspace gas was transferred to 12 mL evacuated vials, and water temperature was
202	measured immediately after. Samples were stored at room temperature and flown to southern
203	Norway for analysis. A 15 mL ambient air sample was taken daily for background correction.
204	Analysis was performed via automated gas chromatography (GC) at the Norwegian
205	University of Life Sciences (NMBU), as described by Yang et al. (2015). A GC autosampler
206	(GC-Pal, CTC, Switzerland) injected 2 mL headspace samples into an Agilent 7890A GC
207	(Santa Clara, CA, USA) with a 20-m wide-bore Poraplot Q column at 38°C, using He as the
208	carrier gas to separate CH <sub>4</sub> and CO <sub>2</sub> from Ar, N <sub>2</sub> , and O <sub>2</sub> . For calibration, certified standards
209	of $CO_2$ and $CH_4$ in He were used (AGA, Germany) and $N_2$ , $O_2$ , and Ar were calibrated using
210	laboratory air. CH4 was measured with a flame ionization detector (FID). A thermal
211	conductivity detector (TCD) was used to measure all other gases.
212	Dissolved gas concentrations were calculated from headspace concentrations corrected for
213	background air, applying temperature-adjusted Henry's law constants (Wilhelm et al., 1977)
214	based on the recorded water temperature. At pH >4, a non-negligible amount of DIC is in the
215	form of (bi)carbonates (HCO <sub>3</sub> <sup>-</sup> , CO <sub>3</sub> <sup>2-</sup> ). The bicarbonate concentrations were calculated based
216	on pH, dissolved $CO_2$ and the temperature-adjusted first dissociation constant (pK <sub>1</sub> = 6.41 at
217	25°C; Stumm and Morgan (2013)) of the carbonic acid equilibrium. Dissolved CO <sub>2</sub> was
218	calculated as DIC minus bicarbonate. To facilitate comparisons with existing studies that
219	report dissolved gases in $\mu$ atm, we converted dissolved gas concentrations to CO <sub>2</sub> or CH <sub>4</sub>





- saturation indexes  $(GHG_{SI})$  assuming atmospheric partial pressures of CO<sub>2</sub> and CH<sub>4</sub> as 400
- 221 μatm and 1.9 μatm, respectively:

222 
$$GHG_{SI} = \frac{[GHG]}{[GHG]_{saturation}}$$

- 223 Where [GHG] is the measured dissolved CO<sub>2</sub> or CH<sub>4</sub> concentration, and [GHG]<sub>saturation</sub> is
- the concentration of dissolved CO<sub>2</sub> or CH<sub>4</sub> at equilibrium with their respective atmospheric
- 225 partial pressure.

226

- 227 2.4 Diffusive CO<sub>2</sub> fluxes from water to atmosphere
- 228 Measurements of CO<sub>2</sub> fluxes from water to atmosphere (diffusive CO<sub>2</sub> fluxes) were measured
- at each site for 30-60 minutes using self-made, opaque flux chambers as described by
- Bastviken et al. (2015) at the water-air interface. The chamber consists of a Senseair K30
- sensor (Senseair AB, Delsbo, Sweden) housed within a plastic bucket that records pCO<sub>2</sub>,
- temperature, and relative humidity every 30 seconds. Fluxes are calculated from the linear
- increase in pCO<sub>2</sub> corrected for ambient temperature and humidity in the chamber (Bastviken
- et al., 2015) considering the internal air volume and the water surface area covered by the
- chamber. Single measurements with a linear increase in pCO<sub>2</sub> with time associated with a
- coefficient of determination  $(R^2)$  lower than 0.9 were discarded.
- 237

#### 238 2.5 Dark incubations

- 239 Water samples were collected for short term dark incubations started directly in the field
- lasting between 18 and 30 hours to estimate DOM mineralization and GHG processing rates.
- 241 Serum flasks (120 mL) were filled with 80 mL of water with a 50 mL syringe equipped with
- 242 a long tube. The syringe was filled and closed under water, and the water was gently pushed
- 243 at the bottom of the serum flask to prevent gas loss. The remaining 40 mL were left with
- ambient air as headspace. The flasks were crimp-sealed with gas-tight, butyl-rubber septa,





sealed, covered with aluminium foil and kept at field temperature (for maximum 6 hours), transported back from the field to be stored at room temperature (18-20°C). The day following the sampling (18 to 30 hours after sampling), the incubations were stopped by adding 1.6 mL 3% HCl to reach a final pH below 2, after which gas samples were taken following the protocols described above. Results from the dark incubation were expressed as rates of DIC production over the course of the incubation period by comparison with initial DIC concentrations and reported as  $\mu$ Mh<sup>-1</sup>:

252 
$$DIC_{rate} = \frac{[DIC]_f - [DIC]_0}{h}$$
 (Eq. 2)

253 where  $[DIC]_f$  is the final solute concentrations in the dark incubation and  $[DIC]_0$  is the initial solute concentration taken in the field (see Sect 2.3) in  $\mu$ M, and h is the incubation duration 254 in hours. In addition, we normalized the DIC production rate with DOC concentration to 255 estimate DOM mineralization rates (per time unit). Also, we calculated the first-order DOM 256 decay rate (yr<sup>-1</sup>) using the exponential decay rate model (Mostovaya et al., 2017). The 257 258 exponential decay model, based on early studies on sediment diagenesis (Boudreau and 259 Ruddick, 1991; Westrich and Berner, 1984), is often the best model to describe decay rates 260 from bioassays in closed systems (Vähätalo et al., 2010) and has been widely used to describe 261 DOM degradation reactions. Under the exponential decay model, the decay constant ( $k_{DOM}$ ; yr<sup>-1</sup>) can be expressed as: 262

263 
$$k_{DOM} = ln\left(\frac{DOC}{DOC - ([DIC]_f - [DIC]_0)M_c}\right) \times \frac{8766}{h}$$
(Eq. 3)

where *DOC* is the DOC concentration in  $\mu$ g L<sup>-1</sup>,  $M_C$  is the molecular mass of C in g mol<sup>-1</sup> and 8766 is the number of hours in a year. Where  $[DIC]_f$  was equal to or below  $[DIC]_0$ , we removed the values from the dataset assuming that the temperature correction of  $[DIC]_0$  was not precise enough (three of 39 samples) to allow quantification of CO<sub>2</sub> processing rates.





- 268 These occurred in September 2020 and October 2021, under cold field conditions, when
- $[DIC]_0$  was overestimated because of unknown sample temperature in the field.

- 271 2.6 Statistical methods
- 272 Statistical analyses were conducted to evaluate differences between sites for various
- 273 measured parameters. One-way analysis of variance (ANOVA) was employed to test for
- 274 differences among groups. Pairwise comparisons of group means were performed using
- 275 Student's t-test using JMP 18.0.11 (2024 JMP Statistical Discovery LLC). For data that did
- 276 not conform to normal distribution assumptions, non-parametric methods were applied,
- 277 specifically the Wilcoxon rank-sum test, to ensure robust comparisons across sites. Results
- are displayed in the form of connecting letters reports within the tables. Sites with the same
- 279 letter (e.g., "A" or "B") indicate no statistically significant differences in the measured
- parameter between those groups at the p < 0.05 significance level. Groups with different
- letters (e.g., "A" vs. "B") are significantly different. When overlapping letters (e.g., "AB") are
- reported, those groups are statistically similar to others with at least one shared letter but may
- 283 differ from groups with entirely distinct letters. Figures were created using the ggplot2
- package (Wickham, 2016) using R software (R Core Team, 2021).

3. Results



285



3.1 Water chemistry 286 287 The thermokarst water bodies were more acidic, richer in DOC and total P, and lower in SO4 and SiO<sub>2</sub> compared with the wetland streams (all differences statistically significant; Table 2; 288 289 Fig. 2). The low pH of the ponds is consistent with their high DOC, and thus high organic 290 acidity. The water bodies aligned along the inverse DOC-pH relationship with TK-Pond 3 at the top, followed by TK-Pond 2 and TK-Pond 1. The TK-Drain usually held an intermediate 291 292 position between the thermokarst ponds and the wetland streams, which were found at the high pH – low DOC end of the curve. Similar patterns were found for DOC-SO4 and DOC-293  $SiO_2$  relationships (Fig. 2). Particulate OC was on average <5% of TOC in the wetland 294 streams, while POC showed considerably more variation in the thermokarst water bodies, 295 possibly related to inputs from destabilized organic matter from the thawing permafrost. 296 All water bodies had NO3 concentrations at, or close to, the detection limit, while the 297 298 thermokarst water bodies had considerable levels of NH<sub>4</sub> contrary to the wetland streams 299 (Table 2). Despite incomplete digestion prior to totN determination, totN values were enough 300 to confirm that the dominant form of N was organic. Total P was highest, and most variable, 301 in the ponds which to some extent mirrored the pattern in DOC, understandably given that in 302 these nutrient-poor sites most P would be in an organic form just like N. 303 The DOM quality indicator SAR was highest in the thermokarst ponds (p<0.03). SAR was positively strongly correlated with DOC concentration (positive, R<sup>2</sup> 0.57, p<0.0001), 304 305 implying that lowest SAR was found in the wetland streams. Other DOM quality indicators 306 (sUVa, associated with aromaticity) tended to be somewhat higher in the wetland streams but did not show significant differences between wetlands and thermokarst water bodies. 307 308





## 309 Table 2. Water chemistry parameters for thermokarst ponds and wetland sites during nine

- 310 sampling campaign. Median values with standard deviations are shown for all water chemistry
- 311 variables, except for pH, which is shown as the median with minimum and maximum values. EC:
- electrical conductivity; SO4: Sulfate; SiO2: silica; DOC: dissolved organic carbon; sUVa: specific UV
- 313 absorbency, SAR: specific UV absorption ratio; TOC: total organic carbon; NH<sub>4</sub>: ammonium; NO<sub>3</sub>:
- 314 nitrate, totP: total organic phosphorous; POC: particulate organic carbon (POC, % of TOC). Letters
- 315 indicate significant differences between sites for each variable (Tukey's t-test, pairwise comparisons,
- 316 *p*<0.05; see Sect. 2.6).

	рН		EC		SO <sub>4</sub>		SiO <sub>2</sub>	
	-		mS m <sup>-1</sup>		mg SO₄ L <sup>-1</sup>		mg SiO <sub>2</sub> L <sup>-1</sup>	
TK-Pond 1	4.49 (4.16-4.79)	В	2.1 (0.7)	В	0.12 (0.07)	С	2.4 (1.8)	BC
TK-Pond 2	4.23 (4.03-4.37)	С	3.2 (0.7)	А	0.18 (0.20)	С	1.0 (1.0)	С
TK-Pond 3	4.06 (3.79-4.32)	С	4.0 (1.6)	А	0.11 (0.02)	С	4.3 (1.7)	В
TK-Drain	4.79 (4.63-4.88)	В	1.6 (0.1)	В	0.16 (0.08)	С	2.3 (1.9)	BC
Inlet	6.69 (6.11-7.36)	А	2.0 (0.5)	В	0.85 (0.18)	А	8.7 (2.4)	А
Outlet	6.56 (6.04-6.97)	А	1.9 (0.3)	В	0.60 (0.25)	В	8.6 (3.0)	Α
	DOC		sUVa		SAR		тос	
	mg C L <sup>-1</sup>		$A_{\lambda 254nm}/mg \ C \ L^{-1}$		$A_{\lambda 254nm}/A_{\lambda 400nm}$		mg C L <sup>-1</sup>	
TK-Pond 1	19.2 (4.4)	С	3.9 (0.4)	ABC	8.4 (0.3)	BC	20.8 (5.1)	С
TK-Pond 2	25.7 (6.4)	В	4.2 (0.3)	А	8.8 (0.5)	AB	27.4 (7.7)	В
TK-Pond 3	34.1 (6.8)	А	3.6 (0.6)	BC	9.2 (0.7)	А	34.8 (9.4)	А
TK-Drain	17.3 (6.6)	С	3.8 (0.4)	С	8.1 (0.2)	AB	19.5 (4.5)	С
Inlet	8.5 (2.1)	D	4.1 (0.4)	AB	7.6 (0.2)	С	8.4 (1.6)	D
Outlet	9.0 (1.5)	D	4.3 (0.3)	А	7.8 (0.2)	С	9.4 (1.6)	D
	NH <sub>4</sub>		NO <sub>4</sub>		TotP		POC	
	μg N L <sup>-1</sup>		μg N L <sup>-1</sup>		μg Ρ L <sup>-1</sup>		% of TOC	
TK-Pond 1	23 (26)	В	2.0 (0.0)	А	27 (17)	В	5.6 (3.0)	В
TK-Pond 2	94 (105)	А	2.1 (0.3)	А	18 (9)	BC	6.1 (4.1)	В
TK-Pond 3	38 (77)	AB	1.9 (0.3)	А	54 (32)	А	8.2 (5.9)	В
TK-Drain	33 (20)	В	2.0 (0.0)	А	20 (18)	BC	17.1 (11.6)	А
Inlet	2 (2)	В	1.9 (0.3)	А	9 (6)	С	4.4 (1.8)	В
Outlet	4 (8)	В	1.9 (0.3)	А	7 (4)	С	4.1 (1.9)	В





The inverse relationship between DOC and pH points towards organic acidity as a strong 318 319 driver of pH. Additionally, the near-to-neutral pH in the wetland streams is consistent with groundwater influences from the catchment, which also would explain the elevated SiO<sub>2</sub> and 320 321 SO<sub>4</sub> concentrations. A limited set of water samples were analysed for base cations (Table SI 322 3), confirming that these were highest in the wetland streams. 323 The water chemical composition of the ponds mirrored the impact of thawing permafrost: the TK-Pond 3 is hydrologically most isolated with the lowest pH, highest conductivity, and 324 highest DOC. TK-Pond 1, located at the transition from peat plateau to wetland, had a higher 325 326 pH and lower EC, DOC and NH<sub>4</sub> than the other ponds, which is consistent with some hydrological influences from the wetland and hence less permafrost impact. TK-Pond 2 is 327 located in the middle of the peat plateau and is by far the largest pond and, under wet 328 conditions, hydrologically connected to neighbouring ponds. The water chemistry of TK-329 Drain was usually most similar to that of TK-Pond 1. An example of pH and EC gradients 330 from the peat plateau into the wetland is consistent with the influence of thermokarst 331 332 waterbodies gradually becoming less dominant in the transition from the peat plateau complex to the wetland (Fig. SI1). 333





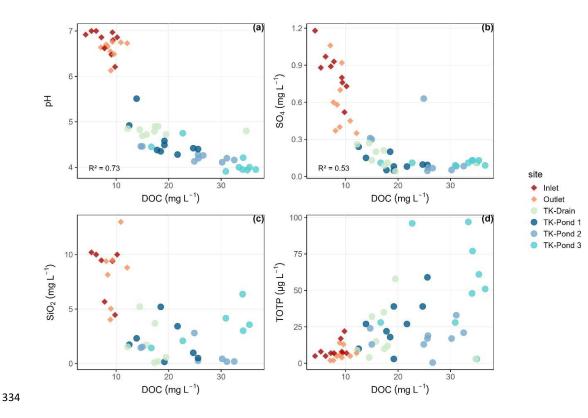


Figure 2. Relationships between Dissolved Organic Carbon (DOC) and various water quality parameters across different sites. The scatter plots demonstrate the relationships between dissolved organic carbon (DOC) and pH (a), sulfate (SO<sub>4</sub>) (b), silica (SiO<sub>2</sub>) (c), and total organic phosphorus (totP) (d).

339

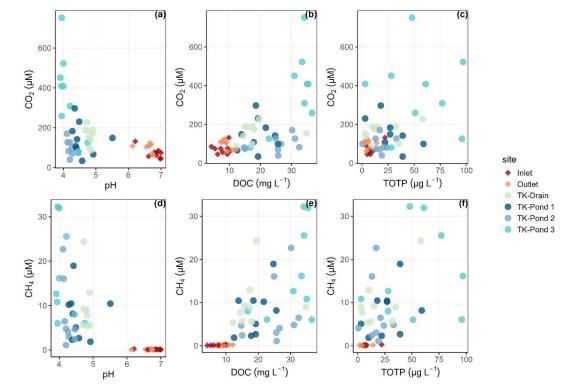
340 3.2 Dissolved gases and gas evasion

All water bodies were oxygenated and dissolved O<sub>2</sub> concentrations were on average 61 to 81% of water O<sub>2</sub> saturation (Table 3). The ponds are shallow which allow for wind mixing and they host sphagnum, suggesting active O<sub>2</sub> production through photosynthesis. All water bodies were oversaturated with CH<sub>4</sub> and CO<sub>2</sub>. Dissolved CH<sub>4</sub> concentrations were 2000–5000 and ~30 times higher than atmospheric equilibrium, in the ponds and in the wetland streams,





- 346 respectively, indicating that all water bodies thermokarst ponds in particular are net
- 347 sources of CH<sub>4</sub> to the atmosphere. The lower CH<sub>4</sub> oversaturation in streams compared with
- 348 ponds is likely related to higher CH<sub>4</sub> losses caused by stream turbulence and/or higher
- 349 production rates of  $CH_4$  in the thermokarst ponds (Fig. 3).





351 Figure 3 Variations in carbon dioxide (CO<sub>2</sub>) and methane (CH<sub>4</sub>) concentrations in relation to pH,

352 DOC, and totP across sampling sites. The upper panels show the relationships between CO<sub>2</sub>

353 concentrations (µM) and pH (a), dissolved organic carbon (DOC, mg L<sup>-1</sup>) (b), and total organic

354 phosphorous (totP,  $\mu$ g L<sup>-1</sup>) (c).

355 CO<sub>2</sub> saturation indexes in the thermokarst waterbodies reached 5 to 20 while in the streams

- they ranged between 3 and 4. By contrast, the  $CO_2$  evasion from the streams (e.g. 1-2 g C m<sup>-2</sup>
- $day^{-1}$ ) was higher than from the ponds (0.3-0.5 C m<sup>-2</sup> day<sup>-1</sup>), consistent with the higher
- turbulence in the streams, and the replenishment of CO<sub>2</sub> from bicarbonates from groundwater





in these streams, which is a geological rather than a recent source of CO<sub>2</sub>. Bicarbonates
contributed about 60-70% to DIC in the wetland streams (with pH between 6.0 and 7.4),
while bicarbonates in thermokarst water bodies were almost negligible (with pH below 4.5),
which is consistent with equilibrium between bicarbonates and CO<sub>2</sub> over these pH ranges.
TK-pond 3 had the lowest O<sub>2</sub> concentrations and the highest CH<sub>4</sub> and CO<sub>2</sub> concentrations of
all thermokarst water bodies. Concentrations of CH<sub>4</sub> and DIC were positively related in the
thermokarst water bodies (r2 0.48, p<0.0001, F-test) but not in the wetland streams (Fig. 4).</li>

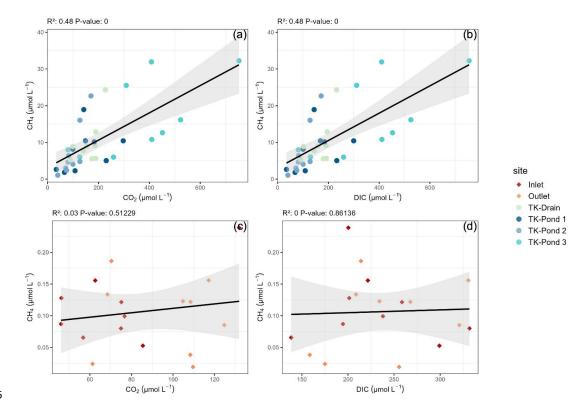




Figure 4 Relationships between CH<sub>4</sub> and (a) CO<sub>2</sub>, (b) DIC for thermokarst waterbodies, and (c) CO<sub>2</sub>,
 (d) DIC for Inlet and Outlet sites including linear regression lines and corresponding R<sup>2</sup> and p-value
 statistics. Note scale differences for CH<sub>4</sub> between thermokarst waterbodies and the wetland
 streams.



371



372	those from the thermokarst waterbodies (Table 3). The mean CO <sub>2</sub> emission at the Inlet site
373	was $1.12\pm0.46$ g C m $^{-2}$ day $^{-1},$ and at the Outlet site $2.20\pm1.15$ g C m $^{-2}$ day $^{-1}.$ These values
374	are 3 to 7 times higher than the fluxes observed from the thermokarst waterbodies, which
375	ranged from 0.30 $\pm$ 0.22 g C m $^{-2}$ day $^{-1}$ (TK-Pond 2) to 0.51 $\pm$ 0.28 g C m $^{-2}$ day $^{-1}$ (TK-Pond 3).
376	Annual CO <sub>2</sub> fluxes for the ice-free period, assuming negligible flux during the ice-covered
377	months, ranged between 51 g C m $^{-2}$ yr $^{-1}$ and 87 g C m $^{-2}$ yr $^{-1}$ for the thermokarst ponds, and
270	the streams ranged between 190 g C m <sup>-2</sup> yr <sup>-1</sup> (Inlet) and 405 g C m <sup>-2</sup> yr <sup>-1</sup> (Outlet).
378	the streams ranged between 196 g e m yr (milet) and 465 g e m yr (Outlet).
378	Table 3. Mean and standard deviations of dissolved gas concentrations and associated metrics for
379	Table 3. Mean and standard deviations of dissolved gas concentrations and associated metrics for
379 380	Table 3. Mean and standard deviations of dissolved gas concentrations and associated metrics for thermokarst ponds and wetland sites across nine sampling campaigns. Mean concentrations of CO <sub>2</sub>
379 380 381	Table 3. Mean and standard deviations of dissolved gas concentrations and associated metrics for thermokarst ponds and wetland sites across nine sampling campaigns. Mean concentrations of $CO_2$ ( $\mu$ M) and $CH_4$ ( $\mu$ M) with their respective saturation ratios, along with $CO_2$ emission flux (g C m <sup>-2</sup> day <sup>-2</sup> )
379 380 381 382	Table 3. Mean and standard deviations of dissolved gas concentrations and associated metrics for thermokarst ponds and wetland sites across nine sampling campaigns. Mean concentrations of $CO_2$ ( $\mu$ M) and $CH_4$ ( $\mu$ M) with their respective saturation ratios, along with $CO_2$ emission flux (g C m <sup>-2</sup> day <sup>-1</sup> ), DIC ( $\mu$ M), and oxygen concentrations ( $\mu$ M) with percent saturation. The saturation ratio is defined

The CO<sub>2</sub> emissions from the stream sites (Inlet and Outlet) were substantially larger than

	CO <sub>2</sub>		DIC	CO <sub>2</sub> emission
		Saturation ratio		
	µmol L⁻¹	CO <sub>2</sub>	µmol L⁻¹	g C m <sup>-2</sup> day <sup>-1</sup>
TK-Pond 1	146 (82) B	7.0 (3.2) BC	149 (83) BC	0.36 (0.28) C
TK-Pond 2	97 (39) B	5.1 (2.2) BC	98 (39) C	0.30 (0.22) C
TK-Pond 3	369 (206) A	18.8 (11.3) A	371 (206) A	0.51 (0.28) C
TK-Drain	161 (45) B	8.1 (2.7) B	165 (46) BC	0.37 (0.15) C
Inlet	73 (26) B	3.1 (1.0) C	232 (59) B	1.12 (0.46) B
Outlet	97 (24) B	4.4 (1.1) BC	241 (59) B	2.20 (1.15) A
	CH <sub>4</sub>		O <sub>2</sub>	
		CH <sub>4</sub> saturation		
	µmol L <sup>-1</sup>	ratio	µmol L⁻¹	% saturation
TK-Pond 1	7.8 (5.5) B	2 330 (1 719) B	273 (59) AB	81 (14) A
TK-Pond 2	7.2 (7.2) B	2150 (2 110) B	266 (85) AB	79 (19) A
TK-Pond 3	16.6 (10.6) A	5 109 (3 470) A	210 (89) B	61 (22) B
TK-Drain	9.8 (5.9) B	2976 (1973) B	260 (65) AB	77 (18) AB
Inlet	0.1 (0.1) C	30 (15) C	297 (59) A	81 (14) A





Outlet 0.1 (0.1) C 28 (18) C 241 (63) AB 67 (14) AB

38	6
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## 387 3.3 DOM processing rates

- 388 Average DIC production rates in the different water bodies were highly variable (5.8 35.7)
- $\mu$ M day<sup>-1</sup>, Table 4), but tended to be highest in the thermokarst ponds compared with the
- 390 wetland streams, while the TK-drain had the lowest rates (Tukey's t-test, p<0.05. The non-
- 391 parametric Wilcoxon tests supported these trends, confirming minimal site-specific effects
- 392 overall, with TK-Drain showing lower activity). These results reflect the in-situ processing of
- 393 DOM in both thermokarst ponds and streams. The DOM mineralization rate did not vary
- significantly between sites and neither did the exponential decay rate kDOM (Table 4).
- kDOM in the thermokarst ponds ranged from 5.5 yr<sup>-1</sup> to 6.7 yr<sup>-1</sup> (Table 4), while the stream
- showed higher kDOM values from 8.3 yr<sup>-1</sup> to 8.8 yr<sup>-1</sup>. The TK-Drain site was substantially
- 397 lower  $(2.0 \text{ yr}^{-1})$ .
- 398 Table 4. Rates of production and decay. DIC rates reflect DIC production. The DIC rate/DOC ratio
- 399 indicates the relative efficiency of converting DOC to DIC, while kDOM indicates the exponential
- 400 decay rate of DOM, showing how quickly DOM is decomposed over time. Letters indicate significant
- 401 differences between sites for each variable (p<0.05).

	DIC rate µmol day⁻¹		DIC rate/DOC µmol g C⁻¹ day⁻¹		kDOM yr <sup>-1</sup>	
TK-Pond 1	26.7 (13.7)	AB	62.9 (34.2)	Α	6.7 (3.7)	Α
TK-Pond 2	34.5 (39.5)	В	62.3 (74.3)	Α	6.7 (8.2)	А
TK-Pond 3	35.7 (33.9)	А	51.2 (36.3)	Α	5.5 (4.0)	А
TK-Drain	5.8 (6.1)	В	19.2 (21.2)	Α	2.0 (2.3)	А
Inlet	18.8 (11.3)	В	77.3 (27.5)	А	8.3 (3.0)	Α
Outlet	20.5 (16.7)	AB	82.0 (64.7)	Α	8.8 (7.0)	Α

## 402 4. Discussion

403 4.1 Thermokarst ponds as hotspots of methane emissions

404 Thermokarst ponds in Iškoras display CH<sub>4</sub> saturation indexes of 2300 to 5000, which is

405 among the highest values reported in the literature for natural waterbodies. These findings





406	align with Shirokova et al. (2012) and Matveev et al. (2018) who documented saturation
407	indexes of 50 to 5000 in Siberian thermokarst depressions, and of 5 to 50 in subarctic lithalsa
408	lakes, respectively. Such high CH4 concentrations in these poorly connected, small, and
409	relatively protected water bodies are consistent with the established inverse relationship
410	between CH4 concentrations and water body size, hydrological connectivity, and turbulence
411	exposure (Abnizova et al., 2012; Kankaala et al., 2013; Polishchuk et al., 2018). Similarly, at
412	Iškoras, the smallest pond exhibited the highest CH4 oversaturation, in combination with the
413	highest DOC, totP, and lowest pH values, likely linked to destabilization of thawing
414	permafrost combined with limited hydrological connectivity. In fact, low pH and elevated
415	totP concentrations are commonly related to increased DOM concentrations (Holmes et al.,
416	2022; Ward and Cory, 2015), which could originate from destabilized permafrost (Turetsky
417	et al., 2020).
418	Thermokarst ponds were highly oversaturated in CH4 despite the presence of dissolved O2.
419	CH4 production is known to occur mainly in anoxic sediments (Bastviken et al., 2004; Clayer
420	et al., 2016; Wik et al., 2016), from where CH4 is subsequently transported to overlying
421	water. The microbial activity responsible for CH4 production may be enhanced by fresh OM
422	input from ongoing thermokarst development (Crevecoeur et al., 2017), as recently observed
423	in laboratory incubations with recently inundated peat material from the Iškoras site (Kjær,
424	2024).
425 426	4.2 Impact of thawing peat plateaus on water chemistry The transition of permafrost-underlain peat plateaus to thermokarst ponds and further to
427	wetlands can markedly shift the landscape-scale GHG dynamics as permafrost continues to
428	thaw (Hugelius et al., 2020; Sannel and Kuhry, 2011). Thermokarst ponds are localized CH4
429	
429	hotspots; however, as these systems transition into wetlands, emissions patterns change due

430 to altered hydrology and biogeochemistry (Holmes et al., 2022; Peura et al., 2019). In





431	wetlands, persistent inundation creates anoxic conditions favourable for methanogenesis,
432	often leading to significant CH4 emissions (Cui et al., 2024). Pirk et al. (2024) demonstrated
433	that fens at Iškoras, as an example of such inundated wetland systems, emit large amounts of
434	CH <sub>4</sub> , particularly where fresh organic carbon is available from peat decomposition from
435	degrading palsa edges. Similarly, Turetsky et al. (2020) noted that newly formed wetlands,
436	arising from permafrost thaw become additional CH4 sources due to labile OC availability.
437	Holmes et al. (2022) observed that inundation in permafrost landscapes increases CH4
438	emissions because of waterlogged conditions that limit oxygen diffusion and enhance
439	anaerobic decomposition. Additionally, Kjær et al. (2024) highlighted that recently thawed
440	permafrost peat in wetland systems, including peat from Iškoras, features a high CH4
441	production potential due to the presence of labile C and methanogenic microbial
442	communities.
443	Unlike thermokarst ponds, which are spatially limited, wetlands have a wider spatial extent.
443 444	Unlike thermokarst ponds, which are spatially limited, wetlands have a wider spatial extent. The carbon balance in wetlands depends not only on $CH_4$ fluxes but also on changes in $CO_2$
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444 445	The carbon balance in wetlands depends not only on CH <sub>4</sub> fluxes but also on changes in CO <sub>2</sub> dynamics as organic matter mineralization occurs under waterlogged conditions (Turetsky et
444 445 446	The carbon balance in wetlands depends not only on $CH_4$ fluxes but also on changes in $CO_2$ dynamics as organic matter mineralization occurs under waterlogged conditions (Turetsky et al., 2020). In this context, the transition from thermokarst ponds to wetlands represents a shift
444 445 446 447	The carbon balance in wetlands depends not only on CH <sub>4</sub> fluxes but also on changes in CO <sub>2</sub> dynamics as organic matter mineralization occurs under waterlogged conditions (Turetsky et al., 2020). In this context, the transition from thermokarst ponds to wetlands represents a shift from localized emissions of both CH <sub>4</sub> and CO <sub>2</sub> emissions to more spatially homogenous CH <sub>4</sub>
444 445 446 447 448	The carbon balance in wetlands depends not only on CH <sub>4</sub> fluxes but also on changes in CO <sub>2</sub> dynamics as organic matter mineralization occurs under waterlogged conditions (Turetsky et al., 2020). In this context, the transition from thermokarst ponds to wetlands represents a shift from localized emissions of both CH <sub>4</sub> and CO <sub>2</sub> emissions to more spatially homogenous CH <sub>4</sub> emissions, while CO <sub>2</sub> is sequestered (Pirk et al., 2024).
444 445 446 447 448 449	The carbon balance in wetlands depends not only on CH <sub>4</sub> fluxes but also on changes in CO <sub>2</sub> dynamics as organic matter mineralization occurs under waterlogged conditions (Turetsky et al., 2020). In this context, the transition from thermokarst ponds to wetlands represents a shift from localized emissions of both CH <sub>4</sub> and CO <sub>2</sub> emissions to more spatially homogenous CH <sub>4</sub> emissions, while CO <sub>2</sub> is sequestered (Pirk et al., 2024). Although a focus on vertical fluxes dominates many studies, lateral DOC fluxes also play a
444 445 446 447 448 449 450	The carbon balance in wetlands depends not only on CH <sub>4</sub> fluxes but also on changes in CO <sub>2</sub> dynamics as organic matter mineralization occurs under waterlogged conditions (Turetsky et al., 2020). In this context, the transition from thermokarst ponds to wetlands represents a shift from localized emissions of both CH <sub>4</sub> and CO <sub>2</sub> emissions to more spatially homogenous CH <sub>4</sub> emissions, while CO <sub>2</sub> is sequestered (Pirk et al., 2024). Although a focus on vertical fluxes dominates many studies, lateral DOC fluxes also play a role in carbon dynamics in permafrost-affected systems. Wetland and peatland ecosystems
444 445 446 447 448 449 450 451	The carbon balance in wetlands depends not only on CH <sub>4</sub> fluxes but also on changes in CO <sub>2</sub> dynamics as organic matter mineralization occurs under waterlogged conditions (Turetsky et al., 2020). In this context, the transition from thermokarst ponds to wetlands represents a shift from localized emissions of both CH <sub>4</sub> and CO <sub>2</sub> emissions to more spatially homogenous CH <sub>4</sub> emissions, while CO <sub>2</sub> is sequestered (Pirk et al., 2024). Although a focus on vertical fluxes dominates many studies, lateral DOC fluxes also play a role in carbon dynamics in permafrost-affected systems. Wetland and peatland ecosystems exhibit high rates of lateral DOC export due to hydrological connectivity (Tank et al., 2018).



455

456



457 contributors and increased hydrological connectivity leading to greater potential lateral C 458 fluxes highlights the need for integrated C budget models that capture the evolving landscape dynamics in permafrost regions. 459 4.3 Carbon dioxide dynamics in thermokarst water bodies 460 Both thermokarst ponds and streams in Iškoras are oversaturated with CO<sub>2</sub>, a common feature 461 462 of Arctic and subarctic aquatic systems (Allesson et al., 2022; Bastviken et al., 2004). However, the mechanisms driving  $CO_2$  fluxes differ between ponds and streams. In ponds, 463 464 despite high CO<sub>2</sub> concentrations, CO<sub>2</sub> release is lower than in streams and likely limited by 465 the lack of turbulence. By contrast, streams exhibit enhanced CO2 fluxes due to high turbulence and carbonate inputs, as described by Zolkos et al. (2019), who emphasized the 466 467 role of groundwater-derived bicarbonates in sustaining CO<sub>2</sub> fluxes in Arctic streams. Quantitatively, CO<sub>2</sub> efflux from streams at Iškoras averages 0.4 g C m<sup>-2</sup> day<sup>-1</sup>, aligning 468 closely with values observed in Siberian permafrost streams (0.3-0.5 g C m<sup>-2</sup> day<sup>-1</sup>; 469 (Shirokova et al., 2012)). This flux reflects the significant contribution of turbulent flow and 470 471 bicarbonate-rich groundwater inputs, processes shown to facilitate CO<sub>2</sub> release (Lundin et al., 2013; Raymond et al., 2013). Streams benefit from continuous replenishment of  $CO_2$  from 472 473 bicarbonates, which account for 60-70% of DIC in these environments (Wallin et al., 2018; Zolkos and Tank, 2020). 474 475 In contrast, CO<sub>2</sub> emissions from ponds at Iškoras are notably lower, similar to findings of

aquatic systems, where it may later contribute to downstream GHG emissions or C

sequestration. The potential for fens and wetlands to transition into significant GHG

- 476 Campeau and Del Giorgio (2014), attributed to the ponds' high DOC-to-bicarbonate ratios,
- which restrict bicarbonate formation and subsequent CO<sub>2</sub> production (Abnizova et al., 2012;
- 478 Bastviken et al., 2004). The limited water mixing in ponds further diminishes CO<sub>2</sub> flux due to
- 479 low gas exchange rates compared to streams. However, DIC production rates in ponds (26.7–





480	35.7 $\mu$ M day <sup>-1</sup> ) remain sufficient to sustain CO <sub>2</sub> effluxes of 5–10 mol C m <sup>-2</sup> year <sup>-1</sup> , consistent
481	with findings by (Shirokova et al., 2012), who linked DIC production directly to DOC
482	concentrations in Arctic aquatic systems. These DIC production rates are relatively high
483	compared to other reported values in the literature (Catalán et al., 2016).
484	High CO <sub>2</sub> evasion from lower-order streams is caused by high flow velocities and associated
485	turbulence causing high gas exchanges (Schelker et al., 2016). These drivers likely explain
486	the relatively higher CO <sub>2</sub> efflux from Iškoras streams compared to ponds. Furthermore, DOM
487	lability, influenced by pH and nutrients, plays a critical role. High molecular-weight DOM, as
488	indicated by low SUVA and SAR values, can inhibit microbial oxidation, thereby slowing
489	DOM decay rates (Shirokova et al., 2019). Ward and Cory (2015) also noted that DOM from
490	thawing permafrost, while less aromatic and more labile compared to active layer DOM, may
491	become limited by environmental factors such as pH and nutrient availability, resulting in
492	lower mineralization rates. The acidic conditions in ponds could shift microbial communities
493	and affect activity (Vigneron et al., 2019), a factor that can further limit CO <sub>2</sub> fluxes from
494	ponds in addition to the low gas exchange rate.
495	In summary, the higher CO <sub>2</sub> fluxes observed in streams at Iškoras are likely driven by the
496	combined effects of turbulent flow, bicarbonate input, and mineral weathering, whereas lower
497	emissions from ponds are shaped by organic acidity, limited hydrological connectivity, and
498	lack of turbulence. These dynamics emphasize the need to account for the interplay of
499	hydrological and chemical factors when assessing C fluxes in permafrost-impacted regions.
500 501	4.4 Climate feedback implications The distinct roles of $\rm CH_4$ and $\rm CO_2$ in the Iškoras landscape underscore their unique climate
502	feedback potentials. The transition from thermokarst ponds to wetlands modifies the overall
503	GHG footprint of the peatland-wetland continuum, balancing the loss of localized CH4
504	emission hotspots with the emergence of sustained, long-term CH4 emissions from wetlands,





505	while the fate of organic matter currently stored in permafrost remains uncertain. At the same
506	time, CO <sub>2</sub> fluxes from the streams and rivers may increase due to enhanced hydrological
507	connectivity and increased organic matter input (Zolkos et al., 2019), in agreement with the
508	results of our study. These findings reflect the complex interplay of ecological and
509	hydrological factors shaping GHG emissions in permafrost landscapes. Turetsky et al. (2020)
510	and Pirk et al. (2024) both emphasized the need for further research on spatiotemporal
511	variability, particularly during thaw cycles, to refine predictions of permafrost C feedbacks.
512	
513	5. Conclusions
514	This study highlights the distinct biogeochemical roles of thermokarst ponds and wetland
515	streams in a landscape of sporadic permafrost in subarctic Norway. Thermokarst ponds at the
516	Iškoras site, characterized by low pH, high organic acidity, and elevated DOC concentrations,
517	are currently hotspots for CH4 emissions, with stable DOM lability driving sustained carbon
518	processing. In contrast, wetland streams exhibit higher CO2 fluxes, largely driven by
519	turbulence and bicarbonate replenishment from groundwater. Despite similarities in DOM
520	mineralization rates between ponds and streams, environmental constraints, such as pH,
521	microbial community composition, and hydrodynamic mixing, are likely controls of the
522	observed differences in GHG fluxes. As thermokarst ponds transition into wetlands, they will
523	no longer function as hotspots for CH4 emissions. Instead, CH4 emissions are likely to
524	increase across the entire landscape, as sustained waterlogging promotes elevated CH4
525	production. These ecological shifts, coupled with lateral DOC losses from peat plateaus,
526	highlight the importance of hydrological connectivity in linking terrestrial and aquatic C
527	dynamics. Such transitions emphasize the need for integrated C budget models that account
528	for the evolving contributions of small aquatic systems to regional and global C cycles.





Future research should prioritize direct measurements of CH4 fluxes, microbial community 529 530 contributions to DOM decomposition under varying environmental constraints, and the temporal variability of gas production and emissions. Additionally, exploring seasonal 531 532 dynamics, lateral carbon transport, and hydrological processes will provide critical insights 533 into C cycling. Investigating catchment-scale signals, such as DOC concentrations across 534 entire river systems and their links to permafrost contributions, can further advance our understanding of landscape-level processes. By addressing these questions, we can better 535 predict the trajectory of permafrost-impacted landscapes and their feedbacks to the global C 536 537 cycle in a warming climate.

## 538 Data availability

All data supporting this study will be made available on a permanent repository uponacceptance.

# 541 Author contributions

- 542 JKK and HDW conceptualized the study. JKK, HDW and FC participated in data collection.
- 543 JKK, FC, HDW and PD conducted the experiments and performed data analysis. JKK, HDW
- and FC created the figures. JKK and HDW drafted the initial manuscript, and HDW, FC, SW
- 545 and PD revised and edited the final version.

# 546 Competing interests

- 547 The authors declare that they have no conflict of interest.
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