



Wintertime Production and Storage of Methane in Thermokarst Ponds of Subarctic Norway

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Abstract. The ongoing climate change in permafrost areas can trigger abrupt thaw processes, leading to the formation of thermokarst lakes and ponds. These water bodies, especially in organic-rich areas, are recognized as strong methane emitters during the ice-free periods and have the potential to accumulate high amounts of methane in and under the ice, which can be released during the ice melt. We estimated wintertime CH₄ storage and daily bottom flux in nine shallow ponds within two permafrost peatlands in Northern Norway, Iškoras and Áidejávri, during the 2023–2024 ice cover season. The wintertime CH₄ storage ranged from 0.6 to 24 g CH₄-C m⁻² and contributed up to 40 % of the annual CH₄ budget at the Iškoras site. The heterogeneity of the CH₄ wintertime accumulation is related to pond depth, differences in vegetation, and the thermokarst pond formation age. The latter has been investigated using a space-for-time substitution approach along chronosequences of thermokarst formation spanning more than 70 years. The winter CH₄ bottom flux increased from 3 mg CH₄-C m⁻² d⁻¹ in two-year-old pond to 107 mg CH₄-C m⁻² d⁻¹ in a pond formed between 30 and 60 years ago. Ponds that formed more than 70 years ago and are currently experiencing sedge regrowth exhibited a high CH₄ bottom flux of 60 mg CH₄-C m⁻² d⁻¹, while older ponds dominated by Sphagnum mosses showed 4 to 10 times lower CH₄ bottom fluxes.

1 Introduction

Lakes and ponds are key sources of methane (CH₄) at high northern latitudes (Wik et al., 2016), particularly in ice-rich permafrost regions where abrupt thaw processes lead to the formation of thermokarst waterbodies (Turetsky et al., 2020). Thermokarst lakes and ponds are recognized as strong methane sources (Heslop et al., 2015; Vonk et al., 2015; Kuhn et al., 2021, Knutson et al., 2025). Recent synthesis studies have summarized existing data on CH₄ fluxes from waterbodies in boreal and Arctic ecosystems (Wik et al., 2016; Denfeld et al., 2018; Kuhn et al., 2021). However, most of this data represents ice-free periods, while studies addressing seasonal variations remain limited. Northern lakes are ice-covered for roughly 60 % of the year underscoring the importance of estimating wintertime CH₄ storage and bottom flux during frozen periods (Walter Anthony et al., 2010, Wik et al., 2011; Boereboom et al., 2012; Greene et al., 2014, Langer et al., 2015). Ice-



covered lakes and ponds often experience anoxic conditions which lead to significant CH₄ accumulation both under and in the ice. This methane is released upon ice-off from the oversaturated water column and from ice bubble storage (Greene et al., 2014; Sepulveda-Jauregui et al., 2015; Vonk et al., 2015). It remains unclear to what extent the emission of CH₄ accumulated during winter contributes to the annual CH₄ budget of arctic landscapes. Several studies have reported high emissions during ice melt (Phelps et al., 1998, Karlsson et al., 2012; Jammet et al., 2015), but quantification of wintertime CH₄ production in northern lakes and ponds is scarce (Boereboom et al., 2012; Langer et al., 2015) and may result in underestimation of annual CH₄ emissions, especially for peatland thermokarst lakes (Matveev et al., 2019).

Northern peatlands cover approximately 3.7 ± 0.5 million km² and store around 415 ± 150 Pg of carbon (Hugelius et al., 2020). Given this large carbon pool, decomposition of peat may trigger significant additional greenhouse gas (GHG) emissions, potentially amplifying global warming. The sporadic permafrost zone outside the mountain regions of Fennoscandia is largely represented by peat plateaus, which are currently undergoing significant degradation due to thermokarst processes and may completely disappear from most of these areas in the coming decades (Borge et al., 2017). The intensification of abrupt thaw is leading to the expansion of small (< 1000 m² in surface area) and shallow (< 2 m in depth) thermokarst lakes and ponds. Despite their significance as active CH₄ sources, these small water bodies are often overlooked in the global assessments, although high CH₄ emissions have been reported in Canada, Northern Sweden, and Russia (Matveev et al., 2016; Kuhn et al., 2018; Burke et al., 2019; Serikova et al., 2019).

In this study, we assessed wintertime CH₄ production and storage in nine ponds across two permafrost peatland sites within the sporadic permafrost zone of Finnmark, Northern Norway. We estimated CH₄ storage under and in the ice during the 2023-2024 winter season. Our specific objectives were to (1) quantify CH₄ winter bottom fluxes in various ponds across two permafrost peatland sites in Finnmark, (2) assess the contribution of wintertime CH₄ storage to the annual CH₄ budget at one of the sites, (3) identify the main factors causing differences in CH₄ winter bottom fluxes, and (4) explore the relationship between the age of thermokarst pond formation and CH₄ winter bottom fluxes.

2 Study area

We investigated nine ponds located within two peat plateau complexes in the sporadic permafrost zone of Finnmark, Northern Norway (Fig. 1a): Iškoras (69°20'26.4"N; 25°17'42.2"E; 380 m a.s.l.) and Áidejávri (68°45'14"N; 23°19'01"E, 398 m a.s.l.). These peatlands have experienced thawing for decades, and the thermokarst pond formation has accelerated significantly in the last 20 years (Borge et al., 2017). Using a chronosequence approach, the formation age of each thermokarst pond was determined using historical aerial images from the Norwegian Mapping Authority (Norgebilder.no, 2025) dating back to 1955 for Iškoras and 1958 for Áidejávri, as well as drone survey images obtained between 2015 and 2023 for both sites (method as described in Martin et al., 2021). The thermokarst pond formation age was identified as the time of permafrost collapse, which typically coincided with water accumulation; however, there can be exceptions with the accumulation of water starting several years after collapse, as observed in the case of pond A4 (Fig. S1). The year 2023



served as the baseline for age estimations. While some thermokarst ponds are currently undergoing a transition to terrestrial wetland ecosystems through sedge and/or peat succession, recently formed ponds continue to expand due to thawing and collapse of peat plateau edges.

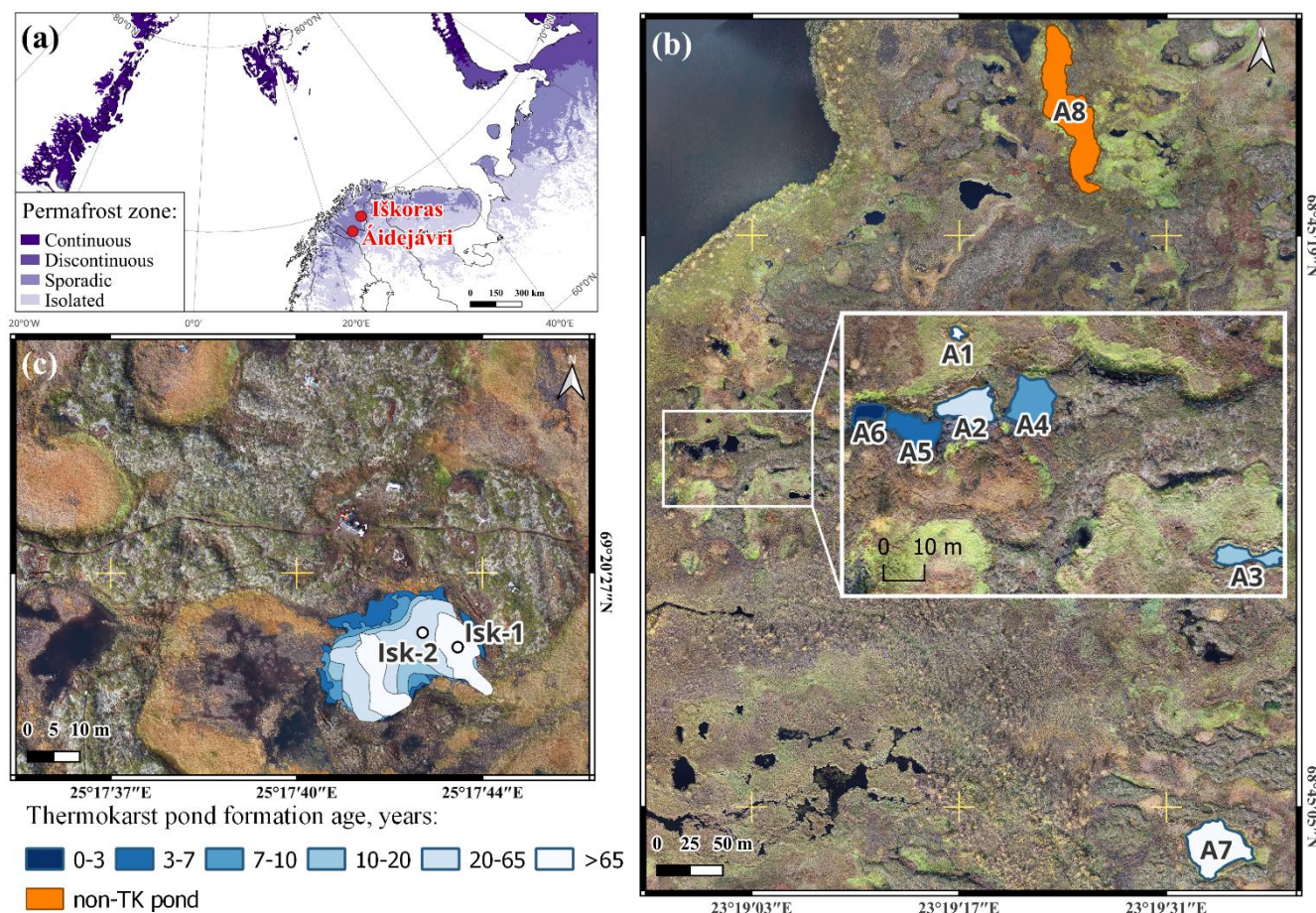


Figure 1. Study area: (a) map showing permafrost distribution in Northern Scandinavia and surrounded territories with the location of two studied peat plateaus (adapted from Obu et al., 2019); (b) studied ponds in Áidejávri (A1-A8); (c) studied pond Isk in Iškoras with two sampling locations (Isk-1 and Isk-2). The colour of the ponds represents the thermokarst pond formation age (Sect. 2). The orthophotos used in (b) and (c) were obtained from a drone survey conducted in 2023 and processed as described in Martin et al., 2021.

Both peat plateau complexes are located within the continental subarctic climate zone, with a mean annual air temperature of -2°C at Áidejávri (meteorological station Sihccajavri) and -1.5°C at Iškoras (meteorological station Cuovddatmohkki) for the period 1980-2024 (<https://seklima.met.no>). The mean air temperature measured at 2 m in Iškoras over the past three years, recorded by the eddy covariance tower at the Iškoras site (Pirk et al., 2024) was 0.15°C . The ice growth season at Iškoras in 2023 began between October 1 and 2, coinciding with the first substantial drop in air temperature (Fig. S2). In Áidejávri, the onset of ice growth occurred between October 6th and 13th as indicated by temperature data from the meteorological station Sihccajavri and Sentinel-2 satellite imagery (Fig. S3). During the sampling in March 2024, the snow thickness varied from



80 0.2 to 0.8 m (Table 1). Ponds less than 0.5 m deep (Table 1) were frozen to the bottom, including the bottom peat layers (ponds A5 and A6), which correspond to the former surface layers of peat plateaus that became submerged due to thermokarst formation. The ice cover remained until mid-May 2024 and disappeared completely by the end of May (Fig. S4). During the fieldwork in October 2024, the ponds at both sites were ice-covered since the 4th of October.

At Áidejávri, we focused on eight shallow ponds (ranging from 0.3 to 1.5 m in depth) located in the northern and eastern parts of the large peat plateau complex (Fig. 1b, Table 1). In the main study area, we studied six thermokarst ponds (A1–A6) 85 formed within the last 80 years. The ponds were selected and named in chronological order, the lowest number representing the oldest pond. All ponds covered an area < 500 m². Pond A1, the oldest in the study area, and pond A3, formed over the last two decades, are rapidly overgrowing with sedges. Ponds A2 and A4–6 are currently expanding and are to some degree hydrologically connected, but initially formed as separate water bodies. The ponds A2, A4–6 share similar vegetation and 90 location conditions but differ in their thermokarst pond formation ages (Table 1). For comparison, we included two reference ponds (A7 and A8) located to the north and east of the main study area at Áidejávri (Fig. 1b). Pond A7, situated in the eastern part and being the oldest studied, has retained its boundaries since 1958 and is currently undergoing slow peat formation (Table 1). Pond A8, which has also been stable since 1958, differs in shape from typical thermokarst ponds in permafrost peatland environments and was therefore classified as non-thermokarst.

95 At Iškoras, the majority of thermokarst ponds is located along the southern edge of the plateau (Fig. 1c). Here, we sampled two locations in the oldest Isk pond (Fig. 1c). The sampling locations represent different pond formation ages, with the oldest part (Isk-1) formed before 1955 and the younger part collapsed between 1955 and 2003 (Isk-2). The pond depth was estimated to be 0.6 m at both locations (Table 1), but the depth determination was challenging due to plant growth and new peat formation within the pond. Despite the ongoing retreat of the surrounding peat plateau edges, the pond borders appear 100 relatively stable. The pond is undergoing Sphagnum colonization, especially on its eastern shore, indicating that it is approaching the final stage of thermokarst pond development. More detailed information about the ponds is given in Table 1. The water temperature in Isk pond at Iškoras was measured at two depths (0.4 m and 0.6 m) using automatic data loggers (Fig. S5), indicating that the pond is well mixed during the ice-free period. However, temperature differences of up to 5°C between the two depths were observed during the initial ice formation and ice melt periods. In winter, the temperature varied 105 due to top-down ice formation which reached the upper logger, but temperature differences did not exceed 2°C.



Table 1. Characteristics of the studied ponds. The thermokarst pond formation age was determined as the period of peat plateau collapse based on historical aerial images. Numbers in parentheses represent the thermokarst pond formation age used for comparisons and figures in this study (the year 2023 was the baseline for age estimations). Snow and ice thicknesses were measured during the field campaign in March 2024. Total ice thickness included the frozen bottom peat layer in ponds A5 and A6, which corresponds to the former surface of the peat plateau that has subsided due to thermokarst.

Pond name	Area, m ²	Depth, m	Stage	Thermokarst pond formation age	Total ice thickness, m	Snow thickness, m
Áidejávri, main study area						
A1	8	1.3	overgrowing with sedges	> 65 (70)	0.49	0.2
A2	88	1.5	expanding	20-65 (40)	0.46	0.5
A3	53	1.1	overgrowing with sedges	10-20 (15)	0.53	0.4
A4	112	0.8	expanding	3-10 (7)	0.58	0.3
A5	84	0.4	expanding	3-7 (5)	0.56	0.8
A6	27	0.4	expanding	1-3 (2)	0.53	0.8
Áidejávri, reference ponds						
A7	1386	0.5	stable, new peat formation	> 65 (100)	0.58	0.3
A8	2577	0.8	non-thermokarst pond		0.45	0.2
Iškoras, Isk pond						
Isk-1	538	0.6	stable, new peat formation	> 68 (70)	0.50	0.6
Isk-2	538	0.6	stable, new peat formation	20-68 (44)	0.55	0.5

3 Methods

3.1. Field Sampling

Fieldwork was carried out in 2024 during March 11–17, September 10–13, and October 4–5. During the winter sampling in March 2024, ice cores and below-ice pond water samples were collected in the center area of each pond. At each sampling location, after measuring the snow thickness, we cleaned the ice surface and drilled through the ice using an ice auger. If the pond was sufficiently deep, water was sampled from both near the bottom and directly below the ice table using a custom-made sampler consisting of a 120 mL serum bottle on a rotating arm. The sampler was lowered to a specific depth with the bottle neck pointing downwards, before turning the bottle upwards to fill it with water. Immediately after bringing the water samples to the surface, dissolved gases were extracted on-site using the acidified headspace method (Åberg and Wallin, 2014). For this, we collected 30 mL of water with a disposable syringe equipped with a 3-way valve and created a 20 mL headspace with ambient air before adding 0.6 mL of 3 % HCl. After shaking for five minutes, the headspace gas was transferred to Helium (He) washed and evacuated 12 mL septum vials (Chromacol). Ambient air was collected at each sampling point to correct for background concentrations for gas extraction. After taking dissolved gas samples, another 50



130 mL of water from the serum bottles was transferred into 50 mL Falcon tubes for dissolved organic carbon (DOC) analysis. These samples were filtered through a 0.45 μm sterile syringe filter with an RC membrane (VWR International) on the same day and subsequently stored at 4°C before being analyzed. Water temperature and pH were measured in the remaining water sample.

After processing the water samples, we extracted ice columns close to the sampling hole using an ice auger. The ice core was
135 subsampled horizontally according to visible differences in texture, shipped frozen to a storage in a cold container (-5°C) at the University of Oslo. In September 2024, we repeated the sampling of dissolved gases and water at the same locations using the procedure described above, except that water was collected from a depth of 0.1 m directly from the pond for dissolved gas measurements. Water for DOC measurements was sampled from the same location and depth, pH and temperature were recorded. In October 2024, additional water samples were extracted for dissolved gases from selected
140 ponds (A1, A2, A4, A5, Isk) both from below the first ice formed and the deeper parts of the pond, again using the water sampler and the acidified headspace method described above.

3.2 Ice sample preparation

Ice samples were cleaned and cut in the cold container at -5°C at the University of Oslo. The bubble structure was visually described, partly following the classification of Boereboom et al. (2012). Samples were characterized as "Superimposed ice",
145 "Clear ice", "Spherical and nut-shaped bubbles", "Elongated bubbles", "Mixed bubbles", "Methane ebullition bubbles" and "Frozen peat". Superimposed ice was found on a top of the ice cores, identified by its texture, brownish color, higher impurities content and high DOC content compared to the ice layers below (Manispurov et al., 2015). Methane ebullition bubbles were identified as relatively large (1-2 cm in diameter), flat bubbles near the surface layers. In the deeper layers, ebullition bubbles were categorized as mixed bubbles.

150 When enough sample material was available, the ice monoliths were divided into three subsamples. Each subsample was placed into a 1050 mL glass jar and sealed with an airtight lid equipped with a sampling septum. The jars were flushed with He using an automated manifold and a vacuum pump, and after releasing He-overpressure, the ice was left to melt at room temperature (+23°C) overnight. The bottles were shaken vertically at 120 rpm for 1 hour to equilibrate gases between the sample and the headspace. Immediately following shaking, the jar headspace was analyzed for CO₂ and CH₄ using a gas
155 chromatograph (GC).

Thereafter, the meltwater was collected to measure the ice water equivalent, pH and DOC content, using the same filtration procedure as for the pond water samples (Sect. 3.1). If the bottom samples contained plant material or peat, they were dried in an oven at +40°C to estimate dry weight and liquid volume. A bulk density of the peat of 0.2 g cm⁻³ was calculated from the dry weight of ca. 100 g of field-wet sample of known volume after freeze-drying for at least 72 h.



160 3.3 Analysis of gas concentrations

Dissolved gases from both the ponds and the ice monoliths were analyzed at the Norwegian University of Life Sciences (Ås, Norway) using a gas chromatograph (GC; Model 7890A, Agilent, Santa Clara, CA, USA) equipped with an autosampler (GC-Pal, CTC, Switzerland). Approximately 2 mL of headspace gas was sampled by a hypodermic needle connected to a peristaltic pump (Gilson Minipuls 3) and admitted to two heated 250 µL sampling loops loading the analyte on two
165 separation columns: a 20 m wide-bore (0.53 mm) Poraplot Q column for the separation of CH₄, CO₂, and N₂O from bulk gases, and a 60 m wide-bore Molsieve 5Å PLOT column for separating Ar, O₂, and N₂. Calibration and conversion of peak areas to ppm were performed using dry bottles with standard gas mixtures (AGA, Norway). The precision of the GC measurements was within 1 % determined by repeated analyses of certified gas standards.

Dissolved concentrations at in situ water temperature were calculated from measured headspace concentrations using
170 temperature-corrected solubility constants (Wilhelm et al., 1977) considering the temperature and volume of the sample at extraction. Dissolved CO₂ concentrations were back calculated to in situ pH using bicarbonate equilibrium constants (Appelo and Postma, 1993). Dissolved oxygen (O₂) concentrations were analyzed for the samples collected in September 2024. To assess the oxygen conditions in the ponds in September 2024, we determined the O₂ saturation in the water relative to atmospheric equilibrium. We classified the oxygen conditions as oxic when the saturation exceeded 30 %, as hypoxic when
175 the saturation was below 30 %, and as anoxic when saturation values were below 1 %. We estimated a high absolute error of 20 % in oxygen saturation measurements due to significant differences observed between measured values in replicate samples. Dissolved oxygen concentrations below ice were not measured.

3.4 pH and Dissolved Organic Carbon (DOC) analysis

The pH was measured using a HANNA instruments pH meter HI9124 with a HI1230B pH electrode, which was 3-point
180 calibrated using Cetripur® buffer solutions from Supelco® (buffer solutions pH 4.01, 7, 10).

The Dissolved Organic Carbon content (DOC) was analyzed using the Total Organic Carbon Analyser (TOCV, Shimadzu, Japan) coupled to an autosampler (ASI-V) using combustion and near infrared detection of CO₂ after removing carbonates by HCl.

3.5 Winter methane bottom flux

185 In this study, we determine the winter methane storage (CH₄) as the sum of methane stored in the ice and in the unfrozen water below the ice. To derive the “winter methane bottom flux” (i.e. the methane flux at the sediment-water interface), we subtract the CH₄ storage prior to freezing (see below) and divide by the time interval between the start of ice formation and the day of sampling in March 2024. The CH₄ storage in the ice was determined by combining the concentrations of the individual ice layers, excluding superimposed ice (Sect. 3.2) which was in contact with the atmosphere. Frozen peat was
190 considered separately; its volumetric CH₄ content was calculated from the concentration in the water phase, corrected by the



volume of the peat using the bulk density (Sect. 3.2). We estimated the uncertainty for individual ice layers using the standard deviation of all available subsamples. In cases where the number of subsamples was insufficient, we assigned an uncertainty based on the average relative error observed in comparable ice layers (Sect. 3.2) from similar ponds. For the frozen peat samples, we calculated the uncertainty by applying the average relative error derived from the deepest ice layers in other ponds. The uncertainty of each layer thickness was estimated during the sample preparation to be around 0.02 m.

We also estimated the potential contribution of superimposed ice, for which it is unclear whether the contained CH₄ is derived from the air (in case of frozen melt- or rainwater) or from pond water pressed upwards through cracks. In eight of the ice cores, the superimposed ice contributed less than 1% of the total CH₄ storage, while it was 5 % for pond A3. In these cases, the uncertainty due to the superimposed ice was considered negligible compared to the uncertainty of the CH₄ concentrations in the individual ice layers. However, in pond A6, superimposed ice comprised half of the ice column, contributing 50 % to the total CH₄ storage. For this pond, we therefore assumed a 50 % uncertainty in the ice storage.

The CH₄ storage in the water column under the ice was estimated for ponds that had not frozen completely to the bottom. We assumed a 5 % uncertainty for the CH₄ measured by the field headspace method, based on existing uncertainty estimates of the method (Åberg and Wallin, 2014; Koschorreck et al., 2021). The presence of plants or loose peat layers on the pond bottoms created challenges for determining whether the probe reached the true bottom of the pond, and we therefore assumed a depth uncertainty of ± 0.1 m. For the pond at Iškoras, disturbances caused by the drilling made it impossible to determine whether it was frozen to the bottom or whether a 0.05 to 0.1 m thick water layer remained below the ice. Furthermore, it was not possible to collect dissolved CH₄ samples from the stirred-up water. As even a thin water layer can contain large amounts of CH₄, we report two confining CH₄ fluxes for the Isk pond: one assuming that the pond had frozen completely to the bottom, and a second flux estimate assuming a water layer of 0.05 m (Isk-2) and 0.1 m (Isk-1) with CH₄ concentration corresponding to the average CH₄ partial pressures from the Áidejávri ponds.

As we do not have data for the water column CH₄ storage prior to freezing in October 2023, we use two confining estimates. First, we calculate the average of the CH₄ concentrations measured in September 2024, with solubility constants corrected for a temperature of 0°C, at which the actual pond freezing occurs. Secondly, we use the CH₄ concentrations in the uppermost winter ice layers which might reflect the gas concentrations in the water prior to freezing. As both estimates are associated with considerable uncertainty, we use the average of both estimate and assign the range to the maximum and minimum values as uncertainty (Sect. 4.3). For the pond depth, we estimated uncertainties of ± 0.1 m for the Áidejávri ponds, and ± 0.2 m for Iškoras. We emphasize that the CH₄ storage prior to freezing is generally small compared to the combined ice and water storage during winter, so that it does not contribute strongly to the overall uncertainty of the winter methane bottom fluxes.

The accumulation period was estimated using meteorological data and Sentinel-2 satellite imagery available for October 2023 (Sect. 2 and Fig. S2-S3). Initial ice formation on both peat plateaus started between October 1 and 13, 2023, while sampling was conducted from March 11 to 17, 2024. For calculation purposes, we set the accumulation period from October 7, 2023 to March 14, 2024, and considered a temporal error of ± 10 days.



225 To compute the uncertainty of our winter CH₄ bottom flux estimates, we use Gaussian error propagation taking into account
all the uncertainties of the individual terms (CH₄ ice and water storage, ice and water depth/thickness, freezing period, etc.).
Spearman's rank correlation coefficients (r_s) were calculated to assess relationships between biogeochemical pond
parameters, estimated fluxes, and environmental factors. The strength of the correlation was characterized as weak ($r_s=0.20$ -
0.39), moderate ($r_s=0.40$ -0.59), strong ($r_s=0.60$ -0.79) and very strong ($r_s=0.80$ -1). A p-value smaller than 0.05 was
230 considered a statistically significant correlation.

4 Results

4.1 Seasonal variations of pH, DOC, and dissolved CH₄ and CO₂ in water

All studied ponds were acidic, with pH levels ranging from 4 to 6.4 (Table 2). The highest pH (6.4) was measured in the
non-thermocarst pond A8. Apart for the latter, the pH increased from September to March for all ponds not frozen to the
235 bottom.

Table 2. Water column pH and Dissolved Organic Carbon (DOC) concentration in studied ponds sampled in March and September 2024. Ponds A5-A7 were frozen to the bottom in March 2024.

Site	Áidejávri							Iškoras	
	A1	A2	A3	A4	A5	A6	A7	A8	Isk
pH									
September	4	4.4	5.5	4.2	4.1	4	4.7	6.4	4.2
March	4.8	4.9	5.9	5	frozen	frozen	frozen	6.2	5
DOC, mg L⁻¹									
September	95	85	75	90	105	105	36	44	42
March	58	78	70	95	frozen	frozen	frozen	36	50

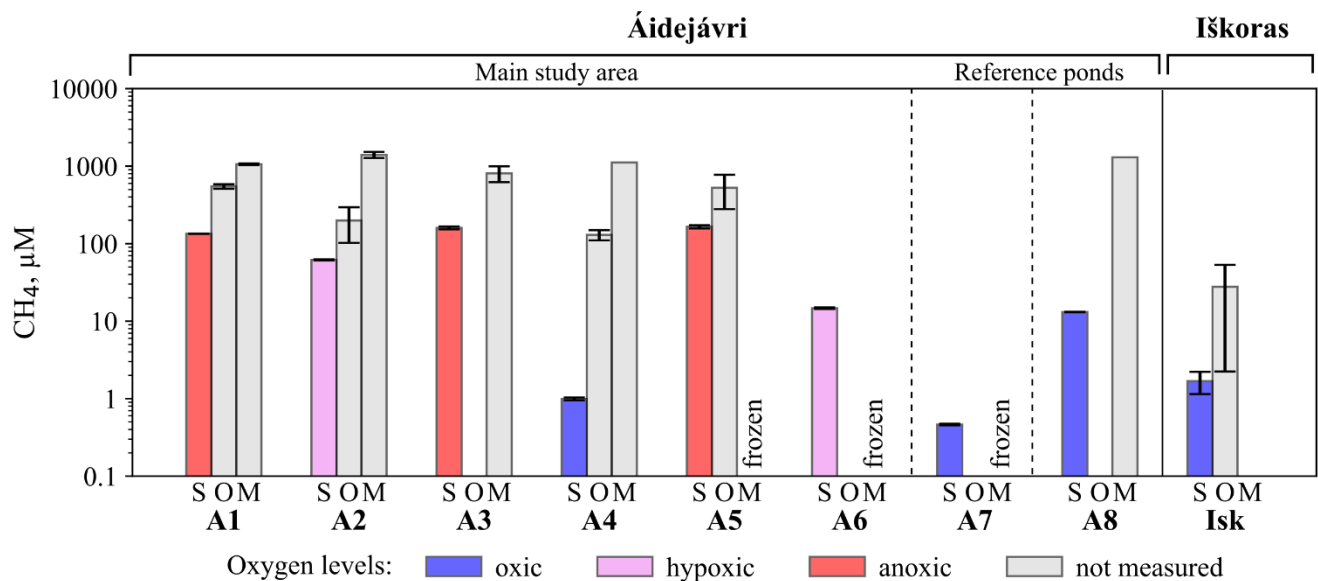
DOC concentrations varied from 36 mg L⁻¹ in the non-thermocarst pond A8, sampled in March, to 105 mg L⁻¹ in the young,
240 shallow ponds A5 and A6, sampled in September (Table 2). In contrast to pH, DOC concentrations showed no clear seasonal
pattern. The largest seasonal difference was observed in pond A1, which is overgrowing with sedges (Table 2). A strongly
negative correlation ($p < 0.05$) was observed between pond size and DOC concentration in September. Concentrations were
low (< 50 mg L⁻¹) in the ponds larger than 500 m² A7 and A8 in Áidejávri, and Isk in Iškoras. Although September DOC
concentrations showed no distinct pattern with pond age, there was a moderate negative correlation ($p < 0.05$) with pond age
245 in the main study area (A1-A6) in Áidejávri. DOC concentrations in the overgrowing ponds A1 and A3 ranged from 58 to 95
mg L⁻¹. DOC concentrations in A2, A4, A5, and A6 averaged at 91 mg L⁻¹ and correlated negatively with ponds age,
increasing from 78 mg L⁻¹ in the oldest pond A2, to a maximum of 105 mg L⁻¹ in the recently formed ponds A5 and A6.



Large ($> 500 \text{ m}^2$) ponds (A7, 8 and Isk) were saturated with oxygen in September (Fig. S6), while the smaller ponds A2 and A6 were hypoxic, and ponds A1, A3 and A5 – anoxic. The only pond with O_2 levels $> 30 \%$ saturation in September was A4 (Fig. S6). Dissolved oxygen concentrations below ice were not measured, but available literature data clearly indicate that ponds experience anoxic conditions (Matveev et al., 2019).

Dissolved CO_2 and CH_4 concentrations generally exceeded atmospheric equilibrium irrespective of season and pond characteristics. Similar to DOC, a strongly negative correlation was observed between pond size and concentrations of both CO_2 ($p < 0.05$) and CH_4 ($p < 0.05$) in September 2024; dissolved CO_2 concentrations were significantly lower in large ponds (Isk and A8 ponds), averaging $30 \mu\text{M}$ (Fig. S7a). The highest dissolved CO_2 concentrations were recorded in the overgrowing ponds A1 and A3, with concentrations of 1740 and $2071 \mu\text{M}$, respectively. Although ponds A2, A4, A5, and A6 were to some extent hydrologically connected, they differed in dissolved concentrations of both CO_2 and CH_4 . CO_2 concentrations in September ranged from $422 \mu\text{M}$ in pond A4 to $1010 \mu\text{M}$ in pond A5 (Fig. S7a). Dissolved CO_2 concentrations increased rapidly with the start of the ice formation in October, in some ponds increasing by an order of magnitude, e.g. in the Isk pond. Winter CO_2 concentrations ranged from $2424 \mu\text{M}$ in pond A3 to $4941 \mu\text{M}$ in pond A4 (Fig. S7a).

In September, the dissolved CH_4 concentrations ranged from 0.4 to $170 \mu\text{M}$, with the lowest values found in the stable ponds Isk, A7, and in expanding pond A4 (Fig. 2). The highest concentrations in September, up to $170 \mu\text{M}$, were measured in the anoxic ponds A1, A3, and pond A5. Notably, the recently formed pond A6, which is connected to A5, had a mean CH_4 concentration that was at least one order of magnitude lower than A5. As with CO_2 , sampling in early October revealed that all ponds experienced a rapid increase in CH_4 concentrations with the start of ice formation with the maximum concentration measured in pond A5 at $701 \mu\text{M}$ (Fig. 2). In pond A4, dissolved CH_4 increased by two orders of magnitude, reaching $144 \mu\text{M}$. The increase at Isk pond was smaller than at Áidejávri ponds, with a maximum increase from $1.7 \mu\text{M}$ to $82 \mu\text{M}$ (Fig. 2). Dissolved CH_4 concentrations in the remaining water column in March ranged from 660 to $1487 \mu\text{M}$ (Fig. 2). Interestingly, the overgrowing ponds A1 and A3, which exhibited high concentrations in September and October, showed relatively smaller CH_4 accumulation over the winter. In contrast, the highest concentrations were measured in the ponds that were hypoxic in September, with the maximum mean value in pond A2 ($1398 \mu\text{M}$). The non-thermokarst pond aligned with the others, exhibiting a value of $1302 \mu\text{M}$ (Fig. 2).



275 **Figure 2. Dissolved water column CH₄ concentrations in March (M), September (S), and October (O), 2024. Error bars represent the standard deviations of multiple samples collected from different depths in March and October, as well as replicate samples taken in September from the same depth.**

In all ponds, the CO₂:CH₄ ratio was higher than 1, with the highest values observed in the oxic ponds during the ice-free season in Áidejávri (Fig. S7b). Ratios were also high at Isk pond, both in the ice-free season and under the first ice formed.

280 In most of the ponds, the ratio decreased from September to March, indicating a shift from oxic to anoxic metabolism.

4.2. pH, DOC, and CH₄ storage in ice

pH values in the ice samples ranged from 4.4 to 8.2, with an average of 5.9 (Table S1). A pH higher than 7 was only observed in the superimposed ice of non-thermocarst pond A8. DOC concentrations ranged from 1.9 mg L⁻¹ to 160 mg L⁻¹ (Table S1). However, for most ice samples without visible organic material, DOC concentrations were confined to a range of

285 1.9 to 7.7 mg L⁻¹. Superimposed ice formed a distinct group, with DOC concentrations ranging from 7.4 mg L⁻¹ in the non-thermocarst pond A8 to 42 mg L⁻¹ in pond A6. Values exceeding 60 mg L⁻¹ were only measured in the bottom ice layers containing peat or plants, or within the frozen peat layers (Table S1).

All ice core layers were classified (Sect. 3.2) as “Superimposed ice”, “Clear ice”, “Methane ebullition bubbles”, “Spherical and nut-shaped bubbles”, “Elongated bubbles”, “Mixed bubbles”, and “Frozen peat” (Fig. 3).

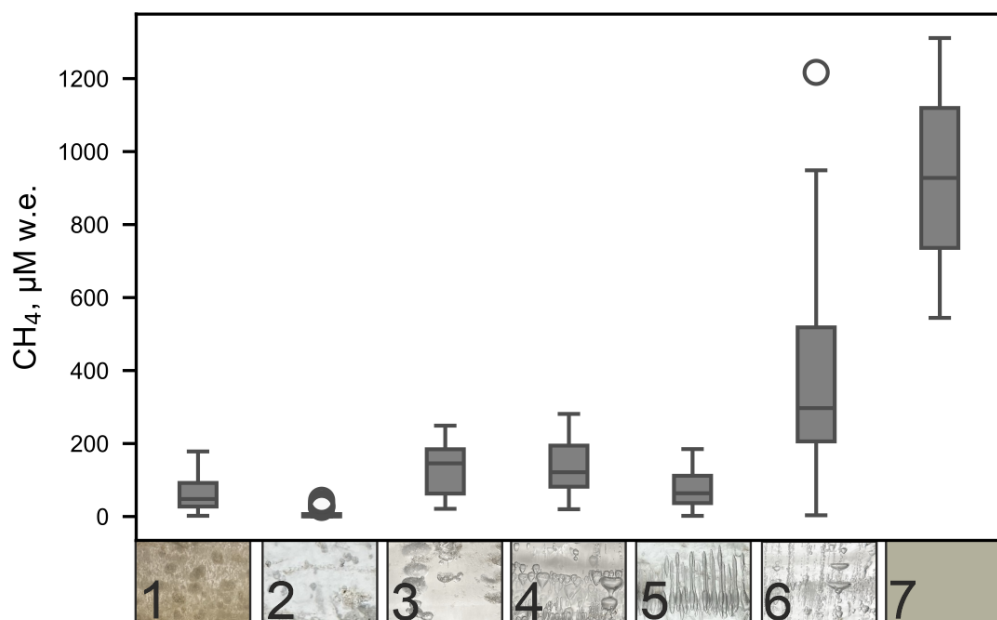


Figure 3. Box plots illustrating methane (CH₄) concentrations in distinct ice types (ice types from Boereboom et al., 2012 with adjustments): 1 – Superimposed ice, 2 – Clear ice, 3 – Methane ebullition bubbles, 4 – Spherical and nut-shaped bubbles, 5 – Elongated bubbles, 6 – Mixed bubbles, 7 – Frozen peat. Error bars are standard deviations across multiple samples within the same ice type category.

Superimposed ice (1) was found on most ponds, forming an upper, up to 0.1 m thick layer (Fig. 4). In ponds A3 and A6, superimposed ice was up to 0.3 m thick and contained bubbles which may indicate that the ice was indeed formed from pond water (and the contained CH₄ thus contributes to the winter CH₄ bottom flux, Sect. 3.2). In addition to our visual inspection, a higher DOC content than in ice layers below was another indicator for the presence of superimposed ice derived from pond water (Table S1). The Isk pond had the lowest CH₄ content in superimposed ice, averaging 2.9 µM w.e. In Áidejávri, the CH₄ content in the superimposed ice ranged from 21 µM w.e. in A3 to 178 µM w.e. in A1.

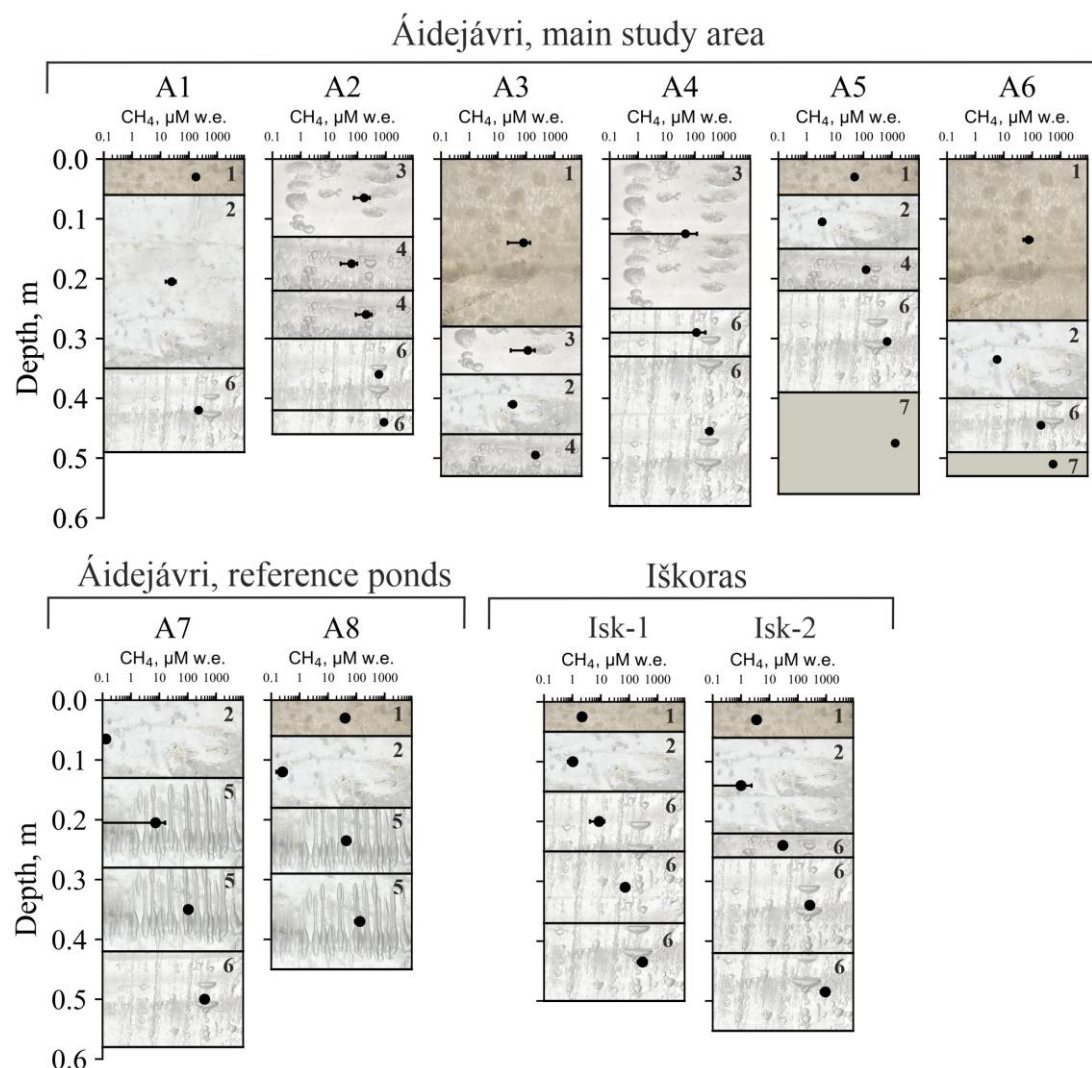


Figure 4. Ice stratigraphy and methane concentration profiles across various ice and frozen peat layers in ponds from Áidejávri (A1–A8) and Iškora (Isk-1 and Isk-2). Error bars are standard deviations across multiple samples. The numbers and the background picture of the ice layers correspond to the ice types indicated in the legend in Fig. 3.

Below the superimposed ice, the CH_4 content in the ice increased with depth. Clear ice (2) layers represented the first winter ice formed and were characterized by small, irregularly distributed bubbles with diameters < 2 mm, typically found as the initial ice layer or beneath the superimposed ice. The CH_4 content in clear ice varied from 0.1 to 45 $\mu\text{M w.e.}$, with the highest value in the anoxic pond A3.

Methane ebullition bubbles (3), relatively flat and up to 2 cm in diameter, were identified in the surface layers (0 – 0.25 m) of ice cores from ponds A2 and A4, as well as in the first ice layer formed under the superimposed ice in pond A3 (depth from 0.3 to 0.4 m). The CH_4 content of ice layers with methane ebullition bubbles varied between 21 and 249 $\mu\text{M w.e.}$. Distinct layers of spherical and nut-shaped bubbles (4) were found in only four ice cores: A2, A3, and as individual thin



layers in A5 and Isk-2 (Fig. 3). CH₄ content of these bubbles ranged from 20 to 281 μM w.e. within the same ice column of pond A2, but at different depths. In pond A3, this type was found the middle ice layer, with an average CH₄ content of 212 μM w.e. Elongated bubbles (5) were present only in two large ponds in Áidejávri: the old thermokarst pond A7 and the non-thermokarst pond A8. The CH₄ content in the ice of these ponds increased with depth, ranging from 1.9 to 122 μM w.e. in A8 and from 43 to 185 μM w.e. in A7.

When spherical, nut-shaped, elongated, and methane ebullition bubbles were found together in an ice layer, it was attributed to the mixed bubbles group (6), typically located in the lower parts of the ice core. This group was identified in all ice cores except for A3 and A8 and exhibited the highest CH₄ content among the ice types (Fig. 3). The minimum (3.4 μM w.e.) and maximum (1217 μM w.e.) CH₄ content for the mixed bubbles group, spanning all sites, were recorded in the parts of Isk pond (Isk-1 and Isk-2, respectively). The concentrations in the deepest layer of the old (Isk-1) and young (Isk-2) parts of the Iskora pond were significantly different, with an average CH₄ content of 933 μM w.e. in ice core Isk-2 (young) and 299 μM w.e. in ice core Isk-1 (old). In Áidejávri, the recently formed pond A6 and the deep sedge-growing pond A1 exhibited the lowest average CH₄ content in the mixed bubble layers around 200 μM w.e., respectively. In contrast, pond A5, despite its connection to A6 and similar depth, had a more than three times higher methane content. The CH₄ content of the mixed bubble layers increased with depth from 567 to 856 μM w.e. in A2, and from 114 to 321 μM w.e. in A4. The shallow ponds A5 and A6 (each with a depth of less than 0.4 m) were completely frozen to the bottom, including the uppermost peat bottom layer. Within the frozen peat (7), the measured CH₄ content were 1311 μM w.e. for pond A5 and 544 μM w.e. for pond A6.

4.3 CH₄ winter storage

The largest contribution of ice storage to total winter CH₄ storage, up to 5010 ± 473 mg CH₄-C m⁻², was found in ponds that were fully or nearly frozen to the bottom, as observed in the Áidejávri pond A5 and the young part of the Isk pond (Isk-2). In contrast, other shallow ponds that were completely frozen to the bottom – such as A6, A7, and Isk-1 – exhibited lower values, from 443 ± 221 to 854 ± 102 mg CH₄-C m⁻², highlighting the significance of thermokarst pond formation age (Sect. 5.4) as a contributing factor. Little CH₄ storage in the ice was observed in the deeper ponds A1, A3, and A8 (Table S2), while the expanding deep thermokarst ponds A2 and A4 showed significantly more CH₄ storage in ice with values of 1580 ± 259 mg CH₄-C m⁻² and 1120 ± 147 mg CH₄-C m⁻², respectively, standing out from the patterns described for the other ponds. The uncertainty of the ice storage estimation did not exceed 30 % in most of the ponds, being highest with 50 % in the ice core A6 with a significant contribution of superimposed ice in the ice column (Sect. 3.5).

Below-ice CH₄ storage was highest in the deep ponds A1 and A2, reaching 16626 ± 1874 mg CH₄-C m⁻² in the pond A2. In the Áidejávri ponds A3, A4, and A8, values ranged from 3347 ± 1349 to 5524 ± 1008 mg CH₄-C m⁻² (Table S2). The uncertainty in these estimations was mostly related to depth error and was less than 40 % for the Áidejávri ponds. The below-ice storage calculated for the Isk pond was 636 mg CH₄-C m⁻² for Isk-2 part and 1273 mg CH₄-C m⁻² for Isk-1 part, with a 100 % uncertainty due to the ambiguity of whether a residual water layer remained below the ice (Sect. 3.5).



345 To estimate the CH₄ storage prior to freezing, we use both measurements from September 2024 and from the first winter ice layer (Sect. 3.5). In the latter, the CH₄ contents (Sect 4.2) were up to 5 times lower than the dissolved CH₄ concentrations measured in September 2024 for most of the ponds (Fig. S8), with the exceptions of A5 and A8 where it was 50 times lower. The largest CH₄ dissolved storage before ice growth was measured in the deepest ponds A1, A2, and A3, with values ranging from 1174 ± 86 to 1364 ± 881 mg CH₄-C m⁻² (Table S2). At the same time, the shallow pond A5 exhibited a storage of 419 ± 415 mg CH₄-C m⁻², while the remaining ponds showed values not exceeding 70 mg CH₄-C m⁻² (Table S2). The initial storage of CH₄ prior to freezing introduces an uncertainty of 5 % to 100 %. However, considering the comparatively low pre-freezing storage relative to the CH₄ storage in and below the ice, this error does not contribute strongly to the final storage uncertainty.

4.4 Winter CH₄ bottom flux

355 The winter CH₄ bottom fluxes (Sect. 3.5) for thermokarst ponds in Áidejávri varied significantly, ranging from 2.8 ± 1.4 mg CH₄-C m⁻² d⁻¹ in the recently formed shallow pond A6 to 107 ± 14 mg CH₄-C m⁻² d⁻¹ in the pond A2 formed 40 years ago (Fig. 5, Table S2). In comparison, the fluxes at the Isk pond were lower than those observed in Áidejávri with similar formation age (Fig. 5). Even assuming that the Isk pond was not frozen to the bottom and adding a potential below-ice CH₄ storage (Sect. 3.5), the fluxes were estimated to be 11 ± 8 mg CH₄-C m⁻² d⁻¹ in the old part of the pond (Isk-1) and 15 ± 5 mg CH₄-C m⁻² d⁻¹ in the section of the pond formed between 20 and 68 years ago (Isk-2).

A significantly higher CH₄ flux was estimated for the non-thermokarst pond A8 with 36 ± 10 mg CH₄-C m⁻² d⁻¹ in comparison to the thermokarst ponds of the same size. The uncertainty in the calculated fluxes was less than 31 % for all ponds, except for pond A6, where the presence of superimposed ice resulted in a 50 % uncertainty in the estimated flux, and the oldest part of Isk pond Isk-1, where uncertainty in below-ice CH₄ water storage led to a higher value of 72 % (Sect. 3.5).

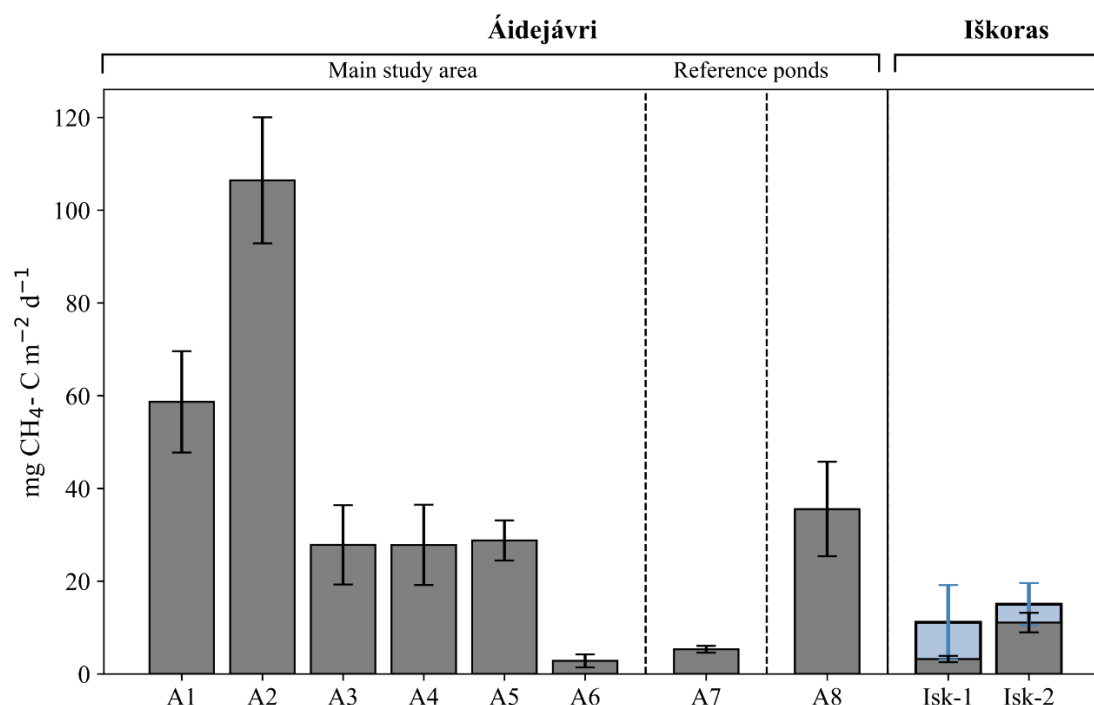
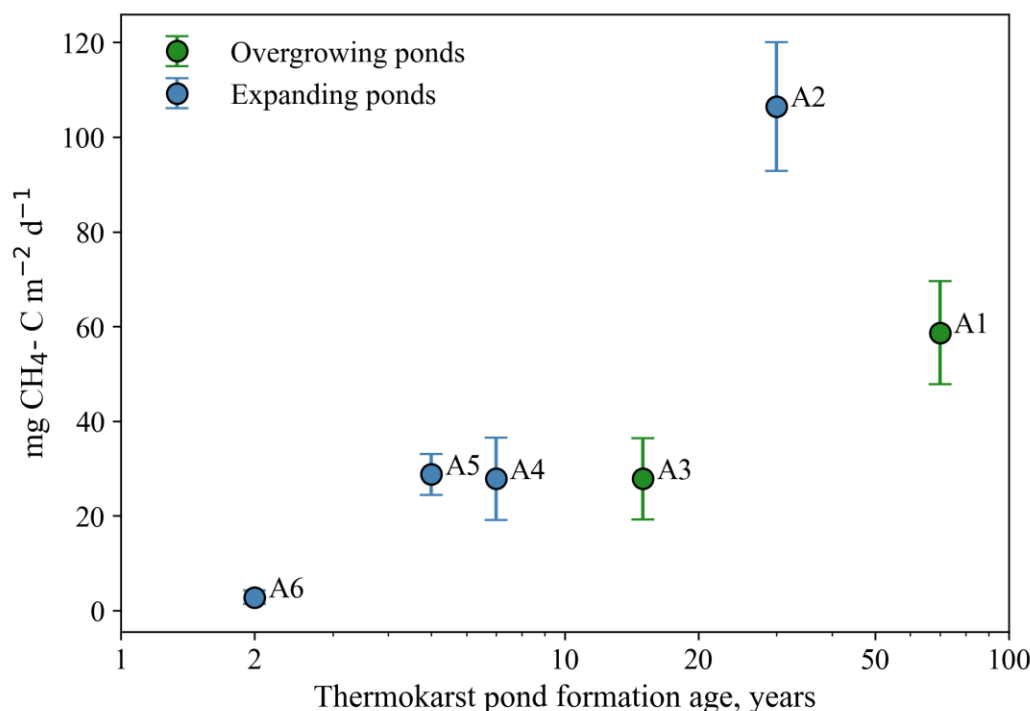


Figure 5. CH₄ winter bottom flux in the studied ponds. Error bars represent the uncertainties calculated using Gaussian error propagation (Sect. 3.5). Blue error bars indicate the values estimated based on the assumption of below-ice dissolved methane storage in the Isk pond (Sect. 3.5) The vertical lines separate the main study area in Áidejávri (A1–A6) from the reference ponds A7 and A8, and the Isk pond.

365 There was a strongly positive correlation ($p < 0.05$) between pond depth and winter bottom CH₄ flux, resulting in the highest fluxes estimated for lakes A1 and A2. However, this pattern does not explain the similar fluxes observed in ponds A3, A4, and A5, which have different depths. For ponds A1-A6, which originated from the same section of the peat plateau (Sect. 2), we compare the relationship between the age of thermokarst pond formation and winter CH₄ bottom fluxes (Fig. 6). We observed an increase in winter CH₄ bottom flux with age, from about 3 mg CH₄-C m⁻² d⁻¹ in the recently formed pond A6 to

375 around 30 mg CH₄-C m⁻² d⁻¹ in ponds A3, A4, and A5, which were formed between 3 and 20 years ago. The highest flux was recorded in the approximately 40-year-old pond A2. The oldest pond A1 had a smaller CH₄ bottom flux than A2, but still higher than all younger ponds. While the statistical evaluation of this age relationship is challenging due to the limited number of ponds, the relationship between pond age and bottom flux is close to being statistically significant ($p = 0.07$).



380 **Figure 6. Winter CH₄ bottom flux vs. thermokarst pond formation age in ponds of the main study area in Áidejávri (A1-A6).**

5 Discussion

5.1 Methodological limitations

We estimated winter bottom CH₄ flux for the 2023–2024 winter season for nine ponds located within two permafrost peatlands in Northern Scandinavia. We based our estimates on methane storage in the ice (ice bubble storage) and the unfrozen water column below, corrected for the storage prior to freezing. Conceptionally, we used the pond ice and the remaining water as a natural chamber to quantify winter methane bottom flux which captures both ebullition and diffusive fluxes of CH₄. Our approach entails the following uncertainties and limitations:

- A significant uncertainty arises from measuring the depth of the water column in the shallow ponds, especially under the winter ice. The loose peat material and presence of plants at the bottom renders accurate measurements of pond depth challenging (Sect. 3.5). We accounted for this by including an estimated uncertainty of 10 to 100 % of the water depth, with 100 % for thin water layers of only 0.05–0.1 m thickness under the ice. Moreover, sampling was conducted at the central part of the ponds, but lateral variations in pond depth could lead to biases in the CH₄ storage accumulated during the winter, particularly in the larger ponds. The presence of plants on the bottom and difficulties in sampling dissolved CH₄ from the residual water layer (Sect 3.5) resulted in high flux uncertainties of about 70 % for the Isk pond locations Isk-1 and Isk-2.



- The upper 0.05–0.1 m of the ice column of most of the frozen ponds was composed of superimposed ice (Sect. 4.2), with nearly half of the column composed of this ice for two of the cores. This type of ice forms primarily by freezing of melt- or rainwater within the snowpack on the ice surface. However, it can also be formed when unfrozen water spreads across the ice surface following crack formation and subsequently refreezes, resulting in high DOC contents in the ice (Manasypov et al., 2015). In the first case, any stored CH₄ would originate from the ambient air, thus not contributing to the bottom CH₄ flux. In the latter case, the CH₄ trapped in the ice originates at least partly from the water body below. Since we cannot distinguish between these cases, we excluded the superimposed ice layer from our CH₄ storage calculations to avoid overestimating CH₄ flux. In any case, CH₄ contents in the superimposed ice were relatively low compared to those in the deeper ice layers and the dissolved CH₄ storage in the water column. For almost all cores, the contribution of superimposed ice to the total CH₄ storage was less than 5 % and thus was negligible in the final uncertainty of winter bottom fluxes. However, for pond A6, superimposed ice comprised half of the ice column, resulting in a contribution of 50 % which was included in the final uncertainty. Therefore, the winter CH₄ bottom flux of A6 features a high relative uncertainty, but it is still clear that the absolute flux values are significantly below those of the other ponds A1–A5 (Fig. 6, Sect. 4.4).
- Another source of uncertainty is related to the lack of measured concentrations of dissolved CH₄ prior to freezing in 2023. To estimate initial CH₄ storage, we used the average concentration from the first winter ice layer combined with available September data from 2024. In some cases, this approach led to an uncertainty of up to 100 % for the CH₄ storage prior to freezing. However, this uncertainty has a limited effect on the final CH₄ flux estimates, as the initial CH₄ storage was relatively small compared to the total CH₄ accumulation during winter. For example, CH₄ storage after winter accumulation was 4 to 200 times higher than the initial CH₄ storage. Notably, in ponds that were anoxic before freezing, the differences between initial and accumulated CH₄ storages (Fig. S8) were less pronounced (4 to 14 times).

Despite the challenges in estimating the individual terms of the CH₄ storage, our approach captures both ebullition and diffusive fluxes during the winter, thus making it well-suited to estimate wintertime CH₄ storage in high latitudes. In this study, we have estimated the uncertainties of the individual terms of the CH₄ winter storage, including ice storage, below-ice water storage and the CH₄ storage prior to freezing, and used Gaussian error propagation to determine an overall uncertainty for the winter CH₄ bottom flux. Our ice records suggested that CH₄ production in shallow ponds continued until the pond was frozen to the bottom in mid-March (Fig. S5). This highlights the importance of assessing both the ice storage and the water storage under the ice. In ponds shallower than 0.6 m, the ice storage is the dominant term for methane winter storage, while the ice storage accounted for less than 25 % of the estimated CH₄ storage in deeper ponds. The absolute magnitude of these individual storage terms strongly influences the final uncertainty, with the uncertainty in shallow ponds largely related to ice storage, while the below-ice water storage dominates the uncertainty for deep ponds.



5.2 Annual CH₄ budget at the Iškoras site

In Finnmark, shallow ponds are ice-covered for at least seven months in a year. Anoxic conditions are likely to establish shortly after the formation of the first winter ice. The lack of oxygen and the reduced activity of methanotrophs at low temperatures lead to accumulation of CH₄ in and under the ice, which is released to the atmosphere during ice-off in late spring (Denfeld et al., 2018). This highlights the importance of accurately estimating wintertime CH₄ production and storage to understand and predict the CH₄ budget of northern water bodies.

Assuming a constant production rate and 228 days of ice cover prior to ice melt on May 22, 2024 (Fig. S4), we estimate the wintertime CH₄ storage in the ponds to be 0.6–24 g CH₄-C m⁻² for Áidejávri and 3 g CH₄-C m⁻² for Iškoras. These estimated storage values in our study are generally higher than those reported for small peatland lakes (Kuhn et al., 2021) and thermokarst lakes in Alaska (Sepulveda-Jauregui et al., 2015). However, they align with CH₄ storage values accumulated until ice melt for thermokarst lakes at a Canadian palsa site, which were around 5 g CH₄-C m⁻² (Matveev et al., 2019).

At Iškoras, methane flux is measured by an eddy covariance tower, and a footprint analysis combined with machine learning has delivered average CH₄ fluxes for the thermokarst ponds within the peat plateau complex (Pirk et al., 2024). As the Isk pond investigated in this study comprises a large fraction of the “thermokarst pond” class within the eddy covariance footprint, it is possible to establish a rough annual CH₄ balance by combining the eddy covariance measurements during the ice-free season with winter-time CH₄ bottom fluxes. For the ice-free season, the mean CH₄ flux for ponds measured by the eddy covariance system is 38 mg CH₄-C m⁻² d⁻¹ (Pirk et al., 2024; Table S3), while our mean daily winter-time CH₄ bottom flux for the Isk pond is about three times lower (13 mg CH₄-C m⁻² d⁻¹). This is consistent with findings from Alaskan lakes, where summer CH₄ production rates increase by a factor of two to three due to warmer bottom sediment temperatures (Matheus Carnevali et al., 2015). The apparent Q₁₀ value for the estimated CH₄ bottom flux at Iškoras is 2.7, based on a mean bottom water temperature of 1°C under ice cover and an average temperature of 11°C during the ice-free period (Fig. S5, Table S3). This value aligns well with previously reported Q₁₀ values for northern lakes (Kuhn et al., 2021).

Assuming that Áidejávri ponds follow the same Q₁₀ relationship as inferred at Iškoras, the corresponding ice-free methane flux for Áidejávri would range from 8 to 300 mg CH₄-C m⁻² d⁻¹, with an average value of 103 mg CH₄-C m⁻² d⁻¹. This assumption allows us to estimate the annual CH₄ budget for each pond system, which ranges from 2 to 66 g CH₄-C m⁻² year⁻¹, with averages (over all ponds) of 8.3 and 23 g CH₄-C m⁻² year⁻¹ at Iškoras and Áidejávri, respectively.

Our back-of-the-envelope calculations suggest that winter methane accumulation at Iškoras constitute up to 40 % of the yearly flux in the ponds. Therefore, the resulting CH₄ storage likely results in elevated emissions during the ice-melt period. Multi-year data from the Iškoras eddy covariance tower showed only small methane flux peaks during the ice-melt period. A direct comparison of these pond flux estimates is difficult, as the eddy covariance flux footprint only comprises about 7 % pond surfaces on average, and the wind direction only seldomly moved the footprint to pond surfaces during the ice-melt period. Hence, a dedicated experimental design would be required to reliably capture the CH₄ ice-off signal in such complex landscapes. In some northern lakes, ice-melt emissions have been reported to contribute up to 60 % of the annual CH₄ budget



(Jansen et al., 2019; Denfeld et al., 2018). By contrast, the CH₄ ice-free emissions from thermokarst ponds in the sporadic permafrost zone of the West Siberian Lowland (WSL) did not differ significantly from ice-off emissions (Serikova et al., 2019). Interestingly, in areas of continuous permafrost, diffusive CH₄ flux during the ice-off and ice-free periods showed pronounced variability (Serikova et al., 2019). This highlights the need for additional observational data for small peatland ponds to better constrain the magnitude of CH₄ fluxes during ice melt.

5.3 Key factors controlling the CH₄ budget

The wintertime CH₄ flux is smaller than the summertime flux, partly caused by temperature differences in the sediment between winter and summer (e.g. Fig. S5). Furthermore, the winter microbiome in thermokarst lake water was found to differ significantly from the summer microbiome in both microbial composition and metabolic functions, with taxa supporting multiple reductive pathways dominating the winter microbiome and enhancing the degradation of permafrost-derived organic matter (Vigneron et al., 2019). This taxonomic shift in the C cycling microbiome goes together with a general shift in oxygen availability; while pO₂ may play an important role for CH₄ formation and oxidation in summer, thermokarst ponds are generally anoxic in winter. Many small ponds in Áidejávri area exhibited anoxic conditions as early as September (Sect. 4.1, Fig. S6), suggesting that CH₄ oxidation is marginal during the ice-free period. By contrast, ponds larger than 500 m² were O₂ saturated during this period (Sect. 4.1, Fig. S6). In small ponds, CH₄ oxidation likely does not play a significant role for the overall CH₄ budget, while in larger ponds CH₄ oxidation might occur at the beginning of the freezing season. However, the effect on the overall CH₄ balance is likely small, as oxygen would be rapidly depleted after an ice cover has formed.

In our study, the highest CH₄ flux was estimated for thermokarst ponds smaller than 500 m². This is in agreement with previous studies that found that pond size is a critical factor for CH₄ fluxes during the ice-free period (Bastviken et al., 2004; Shirokova et al., 2013; Zabelina et al., 2021, Manasypov et al., 2024). Among the key factors driving CH₄ production in our dataset, CH₄ winter bottom flux showed a strongly positive correlation with the pond depth, with deeper lakes consistently exhibiting a higher CH₄ bottom flux.

The CH₄ bottom fluxes observed in Áidejávri ponds were generally higher than those in the Iškoras pond. This aligns with the results of the incubation experiments under controlled temperature and anoxic conditions of permafrost samples from these sites (Kjær et al., 2024). The CH₄ production potential of the Iškoras permafrost samples was lower than for the samples from Áidejávri, which might be related to differences in peat formation history, resulting in different geochemistry of the peat (e.g. a high iron content in Áidejávri) or other local factors (Kjær et al., 2024).

The CH₄ winter bottom fluxes in the studied ponds in Finnmark ranged from 3 to 107 mg CH₄-C m⁻² d⁻¹. While data on wintertime CH₄ bottom flux from thermokarst ponds remain limited, our findings are within the range of winter fluxes reported for other northern regions. In polygonal ponds in the Lena River Delta (Eastern Siberia), winter CH₄ fluxes ranged from 0.01 mg CH₄-C m⁻² d⁻¹ in ponds at the initial development stage to 104 mg CH₄-C m⁻² d⁻¹ in ponds exhibiting signs of thermal erosion (Langer et al., 2015). The mean winter daily flux estimates for glacial lakes in Northern Sweden ranged from



0.001 to 9.5 mg CH₄-C m⁻² d⁻¹ which is at the lower limit of the fluxes measured in Finnmark. This comparison renders thermokarst ponds, particularly those formed in organic-rich regions such as yedoma and permafrost peatlands, to be significant sources of CH₄ compared to other lake types in northern ecosystems (Wik et al., 2016; Kuhn et al., 2021). The relatively higher CH₄ fluxes in thermokarst systems are related to the input of permafrost-derived organic matter, which accelerates CH₄ production rates. In our study, we observed a strongly negative correlation between pond size, dissolved CH₄, and CO₂ concentrations and DOC at the end of the ice-free season in September (Sect. 4.1). Higher DOC concentrations enhance oxygen consumption, leading to oxygen depletion or complete anoxia, which favors methanogenesis and CH₄ release with minimal oxidation during the summer. Together with the predominance of ebullition as a major emission pathway in small thermokarst peatland ponds (Kuhn et al., 2018), this suggests that most of the CH₄ produced during the summer is released to the atmosphere. Factors controlling the production of CH₄ are likely similar in both winter and summer, with seasonal differences largely driven by variations in sediment temperature.

5.4 Winter CH₄ bottom flux vs. thermokarst pond formation age

The CH₄ production in thermokarst water bodies is influenced by a number of factors, including permafrost type (organic-rich vs. non-organic-rich), pond depth and size, as well as vegetation (Burke et al., 2019). Our study design makes it possible to explore the role of the age of thermokarst pond for CH₄ production. For this purpose, we focused on the ponds in the main study area of Áidejávri (A1-A6, Sect. 2) which shared the same peat properties, vegetation, and hydrological regimes, while exhibiting similar DOC concentrations (ranging from 90 to 105 mg L⁻¹). Our results suggest that there may be a functional relationship between the winter CH₄ bottom flux and the thermokarst pond formation age. Thermokarst lakes and ponds undergo several development stages, from initial thawing and expansion to succession into wetland ecosystems. Methane production is generally strongest during the first decades of thermokarst pond evolution when permafrost thawing is most active (Walter et al., 2006; Desyatkin et al., 2009; Shirokova et al., 2013; Vonk et al., 2015). However, in our study, the youngest pond, A6 (2 years old), exhibited the lowest winter CH₄ bottom flux among all the studied lakes. Meanwhile, the pond A5, which is only 2 years older, showed the flux 10 times higher. This delayed onset of CH₄ production could possibly be explained by the lag phase required for methanogenic archaea to adapt and recover after prolonged dormancy in frozen permafrost to new environmental conditions (Rivkina et al., 2007; Knoblauch et al., 2018). This lag is also associated with the shift from previously oxic conditions in the active layer to newly established anoxic conditions conducive to methanogenesis. Such lag phases in CH₄ production are well-documented from incubation studies conducted with Siberian permafrost (Rivkina et al., 2007, Knoblauch et al., 2018) and from the same peatlands in Finnmark studied here (Kjær et al., 2024). Similar to our results, a low CH₄ production during the initial stages of thermokarst pond formation has also been estimated for the polygonal tundra ponds of the Lena River delta (Langer et al., 2015). An alternative explanation for the low CH₄ winter bottom flux in the only two years old pond A6 could be that more CH₄ is indeed produced but is trapped below the still intact root zone of the freshly submerged peat plateau surface, thus preventing its release to the water column. During summer, we observed that a large amount of gas bubbles could be released by disturbing this uppermost root layer



with a probe which suggests that gas is indeed accumulated. While the exact dynamics of this process remain unclear, it is at least possible that not all the produced CH₄ is directly released to the water column and eventually the atmosphere in such young thermokarst water bodies.

530 Pond A2 shows the highest winter CH₄ bottom flux among all studied ponds, while the younger pond A4, which shares similar physical characteristics (i.e., pond depth, location, and evidence of ebullition bubbles in ice column), exhibits fluxes almost four times lower. This difference may indicate that thermokarst pond formation age is a major factor influencing CH₄ production, when other environmental conditions are comparable.

Methane production and emissions from newly formed thermokarst lakes or along thermokarst margins in organic-rich
535 permafrost areas are generally higher than those from older ones (Vonk et al., 2015; Walter Anthony et al., 2016; Heslop et al., 2020). In the Iškora pond, the estimated winter bottom CH₄ flux from the younger part of the pond was at least 30 % higher than from the central part older than 68 years. In the main study area of Áidejávri (A1-A6), our results show that CH₄ production can remain high in ponds older than 60 years. For example, the winter bottom CH₄ flux was about 60 mg CH₄-C m⁻² d⁻¹ for the old pond A1, which is more than double the values found for the relatively young thermokarst ponds (A3-
540 A6).

While the limited number of investigated ponds does not allow a robust statistical evaluation, ponds in the later stages of development during the transition to a permafrost-free mire (A1, A3, A7, Isk) displayed significant differences in CH₄ wintertime production which may be related to vegetation type. Ponds undergoing a transition through sedge regrowth exhibited higher CH₄ winter bottom fluxes than the ones experiencing Sphagnum colonization. Sedges release highly labile
545 organic acids, such as root exudates, including sugars and organic acids, which are rapidly converted into CH₄ by methanogens (Ström et al., 2012; Dorodnikov et al., 2011). Furthermore, vascular plants can reduce oxidation rates and deplete oxygen levels, particularly toward the end of the growing season (Turner et al., 2020). In contrast, the old, stable lakes at both Iškora and Áidejávri (Isk and A7) in which Sphagnum mosses were strongly growing at the edges displayed low winter methane bottom flux (< 15 mg CH₄-C m⁻² d⁻¹). The presence of the Sphagnum mosses may directly reduce CH₄
550 emissions due to methane oxidation processes facilitated by symbiotic relationships between Sphagnum mosses and methane-consuming endophytic bacteria (Raghoebarsing et al., 2005), as well as the presence of aerated surface peat layers (Parmentier, Huissteden, et al., 2011; Magnusson et al., 2020).

Conclusions

This study assesses wintertime methane (CH₄) flux in nine shallow ponds within two permafrost peatlands in Northern
555 Norway (Iškora and Áidejávri) during the 2023–2024 winter season. These ponds form a chronosequences of thermokarst formation spanning more than 70 years. Using pond ice and the below-ice water column as natural chambers, we estimate winter CH₄ bottom flux and storage. Our key findings are:



- At the study sites, ponds remained ice-covered for more than seven months per year. Limited oxygen availability and reduced methanotrophic activity led to CH₄ accumulation. The average wintertime CH₄ bottom flux ranged from 3 to 107 mg CH₄-C m⁻² d⁻¹ across the studied ponds in the two permafrost peatlands.
- The wintertime CH₄ storage ranges from 0.6 to 24 g CH₄-C m⁻² which at the Iškora site constitutes up to 40 % of the annual CH₄ budget.
- Ponds dominated by sedge regrowth have larger CH₄ bottom fluxes, while older ponds experiencing colonization by Sphagnum mosses feature smaller CH₄ bottom fluxes.
- Pond age appears to be a significant factor influencing the wintertime CH₄ bottom flux. A young pond that formed two years prior showed the smallest flux, but fluxes increased considerably with pond age. Our results suggest that the wintertime CH₄ bottom flux can remain at high values (up to 60 mg CH₄-C m⁻² d⁻¹) in thermokarst ponds older than 70 years.

The current atmospheric warming trends may lead to an acceleration of thermokarst processes, resulting in an increase of small shallow thaw ponds in organic-rich permafrost regions. While this could increase CH₄ emissions from these permafrost landscapes, studies on the seasonality and especially the wintertime production of CH₄ remain sparse. This study highlights the need to constrain the wintertime CH₄ production from thermokarst ponds to accurately estimate the present-day and project the future CH₄ budgets at high latitudes.

Data availability. Data supporting this study will be made available in the Zenodo permanent repository upon acceptance (<https://zenodo.org>).

Author contributions. AP, SW and PD conceptualized the research. AP, SW, MRI, CW and SS conducted the field sampling. AP conducted the laboratory analyses, with PD providing expertise and help. AP performed the data analysis, with input from PD, NP and SW. NP contributed with data from eddy covariance measurements in Iškora. AP prepared the manuscript, and all co-authors revised and edited the final version.

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