

1 Spatiotemporal Variability and Environmental Controls on Aquatic Methane Emissions in an Arctic  
2 Permafrost Catchment

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## 8 Abstract

9 Understanding spatiotemporal dynamics and drivers of methane ( $\text{CH}_4$ ) fluxes from rapidly changing  
10 permafrost regions is critical for improving our understanding of such changes. Between May and  
11 August 2023 and 2024, we measured  $\text{CH}_4$  using floating chambers in a small Arctic permafrost  
12 catchment on Disko Island, Greenland. Fluxes were derived from 707 chamber measurements using  
13 a semi-automated algorithm incorporating boosted regression trees and generalized additive  
14 models. Highest fluxes occurred in streams and along lakeshores associated with inlets. Diffusive  
15 fluxes dominated (~98% of observations), while only ~1% of chamber deployments exhibited non-  
16 linear concentration increases indicative of ebullition, while the other ~1% were attributed to  
17 uptake. Median diffusive fluxes were  $5.0 \text{ nmol m}^{-2}\text{s}^{-1}$ , (-0.1 to 271.8), peaking at ice-break. Ebullition  
18 had a median of  $939 \text{ nmol m}^{-2}\text{s}^{-1}$  (5.2 - 14,893), but did not impact overall fluxes. Model results  
19 suggest that thaw-season  $\text{CH}_4$  fluxes were initially driven by meteorological conditions and  
20 catchment soil conditions, but shifted rapidly—within approximately one week after ice-off—to  
21 biogeochemical controls, including dissolved organic matter, oxygen saturation, and pH.

## 22 1 Introduction

23 Permafrost regions across the Arctic store substantial amounts of carbon. Climate warming  
24 is rapidly changing permafrost regions and consequently their carbon storage dynamics, creating a  
25 critical climate feedback mechanism (Schuur et al., 2015; Miner et al., 2022). At current warming  
26 rates, models project approximately 77% of shallow permafrost will be lost by 2100 (Fox-Kemper et  
27 al. 2021), suggesting large implications for the global carbon budget and how carbon emissions are  
28 distributed across permafrost landscapes. The underlying issue is that thawing permafrost can  
29 release previously frozen organic matter, delivering labile nutrients to soil microbes which accelerate  
30 the decomposition of soil organic carbon as a result of their metabolic processes (Schuur et al., 2015;  
31 Keskitalo et al., 2021; Olefeldt et al., 2021). Subtle changes in microbial processes in soils can  
32 enhance positive feedback mechanisms which compounds atmospheric warming. Lateral movement  
33 of water through active layer soils is a critical pathway for  $\text{CH}_4$  emissions from surface waters (Street  
34 et al., 2016; Olid et al., 2021, 2022; Fazi et al., 2021). Hydrological and catchment system dynamics in  
35 particular play a critical role in distributing dissolved carbon throughout permafrost environments.  
36 Catchment systems, such as thermokarst lakes and wetlands, have been shown to be “hotspots” for  
37  $\text{CH}_4$  release, where daily emission rates between  $10$  and  $200 \text{ mg m}^{-2}\text{d}^{-1}$  have been reported (Walter  
38 Anthony et al., 2018; Elder et al., 2020). However, while localized high emissions have been  
39 reported, the overall contribution of Arctic and permafrost freshwater bodies to global methane  
40 budgets is fairly low at 2-6% when compared to other ecoregions such as the tropics at 64%  
41 (Bastviken et al., 2004; Saunois et al., 2025; Virkkala et al., 2024). Nonetheless, with such drastic  
42 change expected, well designed field studies exploring which processes are the most important for

43 governing CH<sub>4</sub> emissions from permafrost regions, are critical data sources for validating climate  
44 models and simulations (Bartsch et al. 2025).

45 Carbon dynamics in permafrost regions have been shown to be governed by interactions  
46 between soil, vegetation, hydrology, and atmospheric processes (Walter Anthony et al., 2012;  
47 Virkkala et al., 2024; Yuan et al., 2024; Kleber et al. 2025). Located on the central-west coast of  
48 Greenland, Qeqertarsuaq, also known as Disko Island, has become an important data point for  
49 understanding environmental interactions which govern Arctic tundra carbon dynamics. The island  
50 provides a natural laboratory for observing interactions between permafrost, vegetation, microbial  
51 activity and aquatic ecosystems (Humlum, 1998; Humlum et al., 1999; Callaghan et al., 2011;  
52 Christiansen et al., 2015; D'Imperio et al., 2017). Previous work from the study area has suggested  
53 shifting hydrology, historic permafrost thaw, nutrient cycling, and microbial activity in the active and  
54 permafrost layers as possible drivers of CH<sub>4</sub> fluxes from surface water bodies (Zastruzny et al., 2017;  
55 Kluge et al., 2021; Stevenson et al., 2021; Juncher Jørgensen et al., 2024). These studies highlight the  
56 interconnectedness of terrestrial and aquatic ecosystems, and the effect they may have on CH<sub>4</sub>  
57 fluxes from lakes and streams on Disko Island. There is yet to be an extensive study on CH<sub>4</sub> fluxes  
58 from the island's lakes and streams. However, it has been suggested that permafrost thaw and  
59 warming air temperatures may have an effect on greenhouse gas fluxes (Kluge et al., 2021; Juncher  
60 Jørgensen et al., 2024). Soil warming experiments and studies of increased snow cover in winter  
61 were shown to regulate carbon fluxes through accelerated carbon turnover (Ravn et al., 2020; Xu et  
62 al., 2021). Carbon fluxes are further controlled by plant uptake and through microbial activity  
63 regulating the availability of nutrients and subsequent CH<sub>4</sub> production (Laanbroek, 2010; Liebner et  
64 al., 2011; D'Imperio et al., 2017). Sedimentary processes in lakes promote carbon storage, whereas  
65 methanotrophic and methanogenic microbial assemblages along an upland–wetland environmental  
66 gradient regulate CH<sub>4</sub> consumption and emission, respectively. Therefore, freshwater ecosystems  
67 play a critical role storing, producing, and emitting CH<sub>4</sub> (Christiansen et al., 2015; Žárský et al., 2018;  
68 Stevenson et al., 2021).

69 The hydrology of Disko Island is strongly influenced by past volcanic activity during the  
70 Paleocene epoch. With extensive basaltic lava flows characterizing the landscape, the island's terrain  
71 is formed by the Maligåt and Vaigat Formations, which are comprised of highly permeable layers of  
72 basalt interbedded with fluvial and lacustrine sediments (Westergaard-Nielsen et al., 2020; Larsen &  
73 Larsen, 2022). The high permeability of these geologic formations enables substantial subsurface  
74 flow, subsequently forming perennial water features such as warm springs. In spring, the soils which  
75 make up the active layer allow for rapid infiltration of meltwater, which laterally distributes  
76 nutrients and organic matter throughout the island's aquatic ecosystems (Westergaard-Nielsen et  
77 al., 2020). For example, during spring runoff meltwater and hillside topography was found to largely  
78 drive the distribution of nitrates from terrestrial to aquatic ecosystems (Zastruzny et al., 2017;  
79 Stevenson et al., 2021). Thus, pools of nutrients available during the growing season may vary  
80 dramatically from one year to the next. Lateral flow of snowmelt and permafrost thaw may influence  
81 CH<sub>4</sub> fluxes due to changes in physio- and biogeochemical properties of the lakes, streams and rivers  
82 on the island (Liebner et al., 2011; Rautio et al., 2011; Walvoord & Kurylyk, 2016; Stevenson et al.,  
83 2021). Although Disko Island has discontinuous permafrost (Christiansen et al., 2015; Kluge et al.,  
84 2021), thawing can release trapped organic matter and nutrients into aquatic ecosystems,  
85 potentially affecting CH<sub>4</sub> fluxes by providing new substrates for microbial activity (Ravn et al., 2020;  
86 Stevenson et al., 2021; Westergaard-Nielsen et al., 2020; Xu et al., 2021).

87 The distribution and drivers of aquatic CH<sub>4</sub> emissions in permafrost regions remain poorly  
88 constrained, particularly across small lakes and streams which may arise as emission hotspots.

89 Previous studies on Disko Island have highlighted the potential importance of hydrology, permafrost  
90 thaw, and microbial processes for greenhouse gas fluxes, but comprehensive spatial and seasonal  
91 assessments of CH<sub>4</sub> are lacking. In this study we address this gap by quantifying CH<sub>4</sub> fluxes from 707  
92 floating chamber measurements across a permafrost-affected catchment (Sanningasup Tasia). Using  
93 boosted regression trees, we evaluate the partial effects of physiochemical water conditions,  
94 catchment soil conditions, and meteorology in regulating emissions from ice-break through the  
95 growing season. Our objective was to determine how spatial heterogeneity and seasonal dynamics  
96 shape CH<sub>4</sub> emissions from Arctic freshwater ecosystems and to identify the key processes that  
97 control flux variability in permafrost catchments.

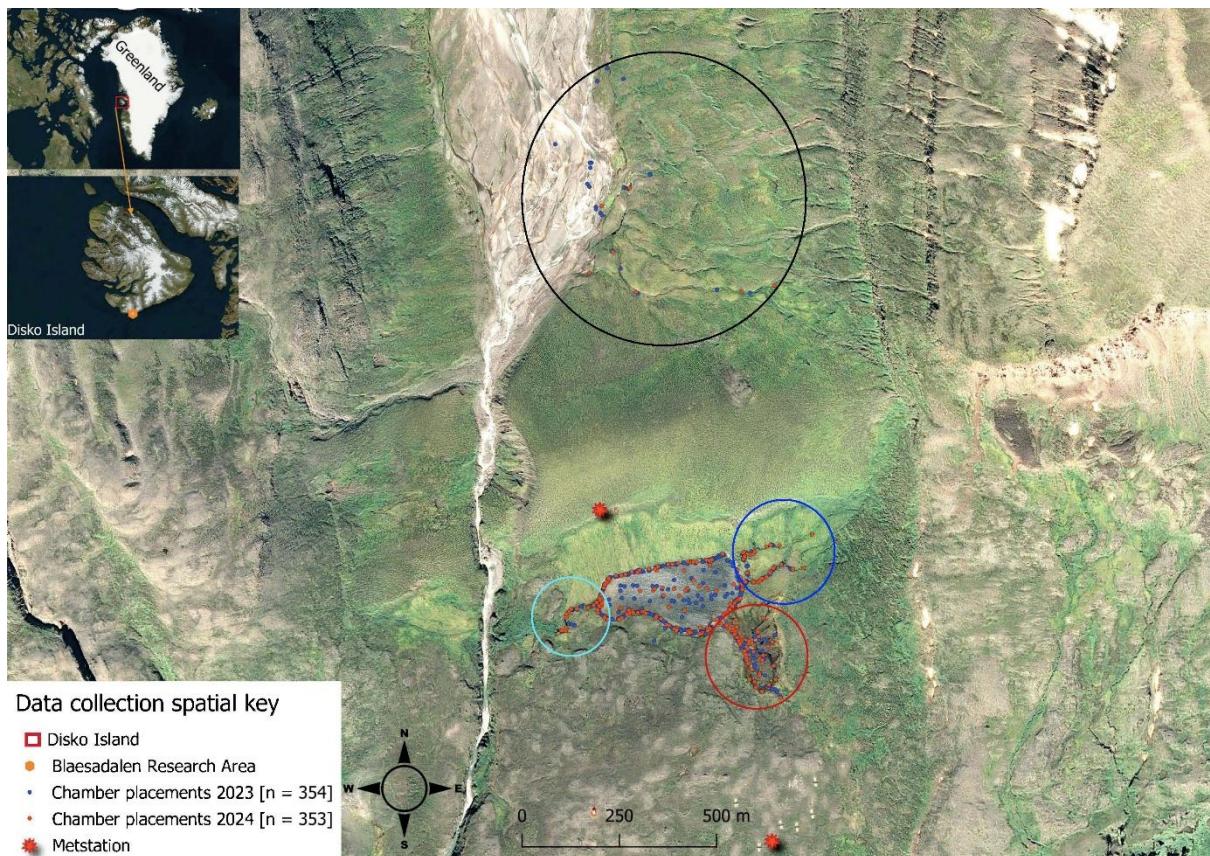
## 98 2 Methods

### 99 2.1 Study Site

100 Lake Sanningasup Tasia in Greenlandic, or Moræne sø in Danish, is situated between  
101 moraines in the north and east and an outlet which drains into the Red River to the west (Figure 1).  
102 The lake is primarily fed by a large warm spring which enters the lake from the southeast, forming a  
103 wetland type ecosystem. The other inlets of the lake are primarily fed by seasonal snowmelt. The  
104 heterogeneity of the catchment provides an exemplary study site, allowing us to understand the  
105 mechanisms regulating CH<sub>4</sub> emissions from a lake, streams, and wetland. According to a 2018 report  
106 from the University of Copenhagen, the lake has a maximum depth of 4.5 m and is generally  
107 phosphorus limited with nitrogen concentrations being seasonally variable, where concentrations  
108 during ice cover are higher than during periods of no ice cover (Westergaard-Nielsen et al., 2020).  
109 We found water temperature of the lake to range between 1.1 and 13.9 °C with a mean of 7.9 °C. To  
110 our knowledge there has never been an extensive study on the greenhouse gas fluxes from the lake  
111 and surrounding water bodies.

112

113



114

115 Figure 1. Map showing the 707 chamber measurements (blue and brown dots). Points in the south  
116 are concentrated around Lake Sanningasup Tasia and its connected streams. The blue circle indicates  
117 streams fed by snowmelt and the turquoise circle indicates the outlet of the lake. The red circle  
118 indicates the warm spring area which forms a wetland type ecosystem. Points inside the black circle  
119 north of the lake indicate measurements taken from the Red River and its stream tributaries.  
120 Orthomosaic background image © CNES (2024), Distribution Airbus DS, produced from Pléiades 1B  
121 satellite imagery.

## 122 2.2 Data Collection

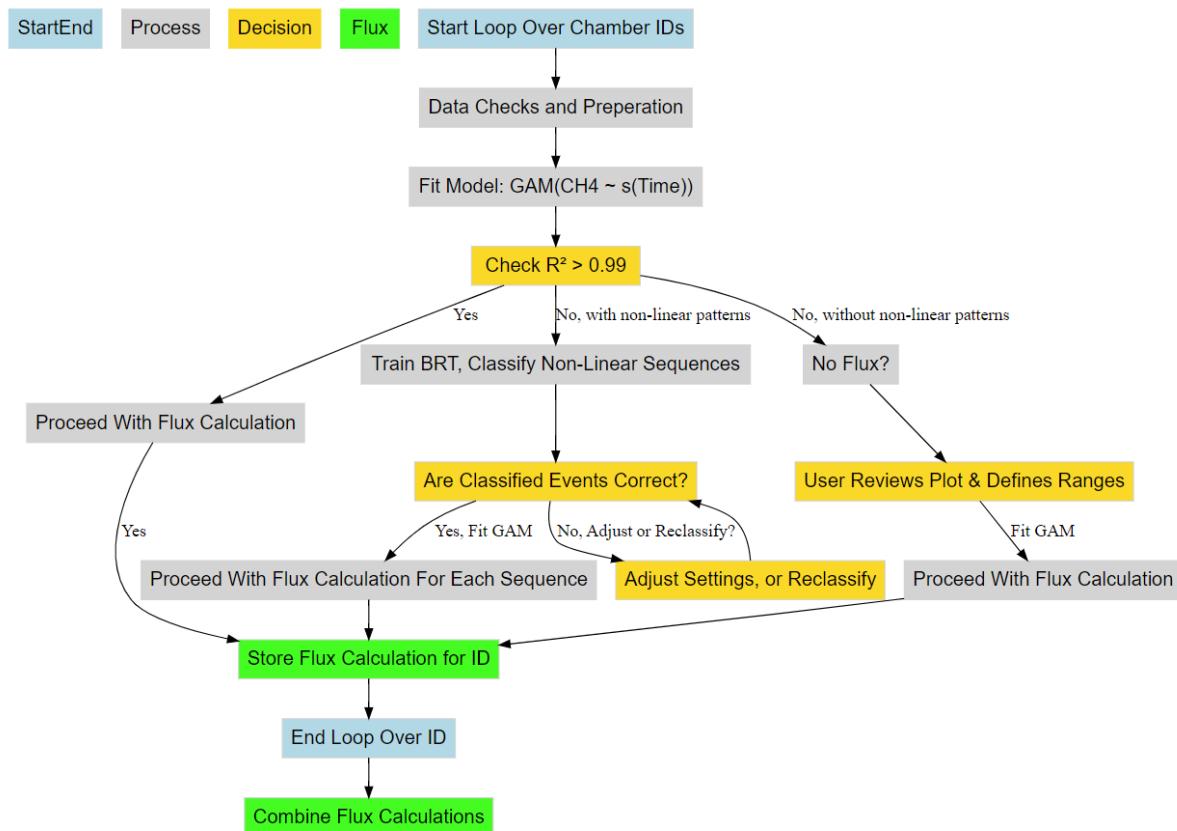
123 We used closed floating chamber systems connected to an ABB/Los Gatos Research GLA131  
124 Series Micro portable Gas Analyzer in 2023, and to a LI-COR® LI-7810 Trace Gas Analyzer in 2024,  
125 with the goal of capturing the spatial and temporal variability of CH<sub>4</sub> fluxes in the catchment area. In  
126 2023, we used a self-built cylindrical chamber made of semi-transparent plastic material with  
127 volumetric capacity of 0.016 m<sup>3</sup> and a basal area of 0.096 m<sup>2</sup>. In 2024, we used a West Systems type  
128 C flux chamber made of stainless steel with a volumetric capacity of 0.013 m<sup>3</sup> and a basal area of  
129 0.07 m<sup>2</sup>. Both chambers included a fan for mixing air and a temperature sensor. The semi-  
130 transparent plastic chamber used a circular foam floater that was wrapped around the outside of  
131 the chamber, allowing 2cm of the chamber to be below the water surface, forming a 100% airtight  
132 seal. The West Systems chamber was inserted into a closed-cell foam floater, where the seal was  
133 created once the chamber was inserted into the floater. Despite differing chamber materials and  
134 flotation devices, median fluxes between 2023 and 2024 were identical at 5.0 nmol m<sup>-2</sup>s<sup>-1</sup>. However,  
135 to evaluate whether changes in chamber construction between years introduced systematic bias in  
136 CH<sub>4</sub> flux estimates, we fit a linear model using log-transformed flux as the response and chamber  
137 type, latitude, longitude, and Julian day as predictors. Chamber type was not a significant predictor

138 ( $p = 0.13$ ), and QQ plots of log-transformed fluxes across years showed no consistent deviation  
139 across the flux distribution, except at lower emission rates (Figure S1). These findings suggest that  
140 differences in chamber construction did not substantively influence the calculated fluxes. Chamber  
141 measurements were conducted on the surface of the lake and surrounding water bodies at a spatial  
142 distance of 10 to 20 m with a closure time of 10 minutes. The chamber and gas analyzer were  
143 connected in a closed loop, and sample air was continuously pumped through the gas analyzer. CH<sub>4</sub>  
144 concentrations were measured with a frequency of 1 Hz. Each flux measurement was given a unique  
145 id based on its spatial location or water body type. For the lake, ids were given based on what  
146 shoreline we were measuring on. For example, if on the east shore, ids would be e1, e2, e3, ... etc.  
147 The majority of flux measurements on the lake were conducted within 0.2 m from the shore. Open  
148 water measurements in the lake were made using a small boat and anchor system. CH<sub>4</sub>  
149 concentrations in streams were measured by starting at, or near the headwaters and then taking  
150 measurements progressively downstream with consideration to the changing terrestrial vegetation  
151 and stream dynamics (i.e., fast, slow, or eddie pool). In 2023, we measured isolated meltwater pools  
152 during the thaw to represent control conditions (water not yet connected to the lake or streams). In  
153 2024, to capture an even earlier baseline, we conducted chamber measurements on top of snow  
154 and lake ice prior to thaw onset, providing a true pre-thaw control period. Overall, we made 707  
155 chamber measurements, representing ~10-15 chamber placements taking place daily, around the  
156 lake and surrounding streams giving us an extensive view of the spatial and temporal variability of  
157 CH<sub>4</sub> fluxes in the study area. While floating chambers isolate the headspace from light wind  
158 disturbance, increased surface turbulence may influence gas exchange in the open space of the  
159 chamber (Vachon and Prairie 2013). Our approach captures diffusive exchange under mostly  
160 calm-water conditions (i.e., wind speed up to ~4 ms<sup>-1</sup>), but we acknowledge that regional wind-  
161 driven mixing may contribute to flux variability beyond individual chamber footprints. We  
162 simultaneously measured water temperature using Truebner EC-100 RS-485 EC/Temperature  
163 sensors in 2023 and a suite of water parameters were collected in 2024 using an AquaTroll 600  
164 water sonde (see section: Decoding Methane Drivers). Meteorological data and soil characteristics  
165 were collected from nearby meteorological stations maintained by Aarhus University which are part  
166 of the Greenland Ecosystem Monitoring Database (Greenland Ecosystem Monitoring, 2025a-d) (see  
167 section “Decoding Methane Drivers” for list of variables used).

### 168 2.3 Flux Algorithm & Ebullition Detection

169 In collaboration with ChatGPT 4.0, we wrote an interactive algorithm in R which leverages  
170 General Additive Models (GAM) and Boosted Regression Trees (BRT) to robustly and flexibly  
171 calculate CH<sub>4</sub> fluxes from individual floating chamber measurements (Figure 6). The flux calculation  
172 procedure was applied identically to individual chamber time series for both 2023 and 2024, while  
173 controlling for different the chamber constructions.

174



175

176 Figure 2. CH<sub>4</sub> flux calculation workflow from concentration data using predictions from GAM and  
 177 BRT. Blue boxes represent the start and end of a single chamber measurement working through the  
 178 algorithms processes (grey boxes) and decision logic (gold boxes). Green boxes represent the storage  
 179 and combination of the results for further analysis.

180 Rather than fitting concentration data with linear, exponential, and/or polynomial models  
 181 (Kutzbach et al., 2007; Pedersen et al., 2010; Hoffmann et al., 2017), the algorithm fits GAM, which  
 182 are capable of modelling non-linear patterns without a-priori specification of the functional form of  
 183 the relationship between time and concentration. However, before fitting a GAM, the concentration  
 184 and accompanying data is checked and processed (Figure 2; “Data Checks and Preparation”) as  
 185 follows: The algorithm conducts a preliminary check for the required chamber parameters which  
 186 are; id, ordered times of measurement, air temperature (°C), volume, area, and air pressure. In  
 187 addition, air temperature is expected to be initially in Celsius, which is automatically converted to  
 188 Kelvin during the processing of concentration data in preceding steps. The data is then ordered  
 189 based on id and time to maintain correct chronological order of chamber measurements. CH<sub>4</sub> is then  
 190 converted from ppm to moles using Ideal Gas Law:

191

$$CH_4_{moles} = \frac{(CH_4_{ppm} \times P \times V)}{(R \times T)} \quad (1)$$

193

194 where P is air pressure (Pa), V is chamber volume (m<sup>3</sup>), R (8.314 J / (mol × K) is the universal gas  
 195 constant, and T is air temperature inside the chamber (K). After preliminary data checks and initial

196 processing of the concentration data, the concentration time series is then fit to a GAM (i.e.  
197  $\text{gam}(\text{CH4}_{\text{moles}} \sim \text{s}(\text{time}, \text{k} = \text{gam\_knots}))$ ), where the smoothing parameter 'k' is a user defined  
198 parameter named 'gam\_knots' which has a default value of 5. The value of 'k' cannot exceed 3 times  
199 the degrees of freedom for a given concentration time series, or the algorithm defaults to fitting a  
200 linear model. The algorithm then checks the  $R^2$  value of the fitted GAM to see if it meets the default  
201 conditional value of  $\geq 0.99$ , if so, it proceeds to calculate fluxes following these steps:

202

203 
$$\Delta \text{CH4}_{\text{moles}} = \left[ \frac{\text{CH4}(t_2) - \text{CH4}(t_1)}{t_2 - t_1}, \frac{\text{CH4}(t_3) - \text{CH4}(t_2)}{t_3 - t_2}, \dots, \frac{\text{CH4}(t_n) - \text{CH4}(t_{n-1})}{t_n - t_{n-1}} \right] \quad (2)$$

204

205 Where  $\Delta \text{CH4}_{\text{moles}}$  is the rate of change, calculated as the quotient of the predicted differences in  
206  $\text{CH}_4$  concentration between successive time points. Flux is then calculated between successive time  
207 points by:

208

209 
$$\bar{F}_{\text{Flux}} = \frac{1}{n-1} \sum_{i=1}^{n-1} \frac{\Delta \text{CH4}}{A} \quad (3)$$

210

211 Where mean flux of the chamber measurement is estimated by calculating flux at each  
212 successive time step, where flux is determined by dividing  $\Delta \text{CH4}_{\text{moles}}$  by the basal area (A) of the  
213 chamber, expressed in ( $\text{m}^2$ ). A plot of the time series and model fit is generated and saved in the file  
214 directory defined by the user by setting the parameter 'save\_directory' (Figures S1-5). Because  
215 fluxes were derived from high-frequency (1 Hz) concentration data fitted using a GAM with a strict  
216 acceptance threshold of  $R^2 \geq 0.99$ , analytical uncertainty in the rate-of-change estimation is  
217 negligible. Conventional uncertainty propagation (e.g., based on regression slope error or replicate  
218 chambers) is not meaningful in this context because the GAM approach fits a smooth curve through  
219 hundreds of data points per deployment, effectively minimizing noise and preventing poor-quality  
220 fits from contributing to the final flux values. This ensures that the dominant source of variability in  
221 the dataset reflects true environmental heterogeneity rather than analytical error. Furthermore,  
222 because the chamber headspace was fully sealed and isolated from external turbulence, wind-  
223 induced variability—which often motivates uncertainty corrections—is mechanically removed from  
224 the flux calculation process. For these reasons, we report spatial variability (e.g., medians, ranges,  
225 and interquartile spread) rather than analytical uncertainty, as it provides a more ecologically  
226 relevant representation of flux variability across the catchment.

227 In the cases where the initial GAM fit does not meet the  $R^2 \geq 0.99$  condition, the algorithm  
228 can follow two pathways (Figure 2). Pathway (1) is a result of the algorithm having detected non-  
229 linear concentration increases using BRT, while pathway (2) the algorithm has found the chamber  
230 measurement has not met any of the conditional requirements for flux calculations, or more  
231 generally stated, there was no measurable concentration increase detected automatically. Both  
232 pathways are interactive as the user is prompted to confirm the classification of "ebullition" versus.  
233 diffusive data sequences in pathway (1), while in pathway (2) the user confirms there is indeed no  
234 concentration increase by reviewing the diagnostic plots (see Supplemental text and Figures S2-S6).  
235 Once confirmed, the user initiates flux calculations by manually entering the time range of the  
236 measurement that should be fitted (Figure 2). "Ebullition" in the context of the algorithm refers to a

237 sudden, non-linear  $\text{CH}_4$  increases identified by the algorithm, which likely includes ebullitive events,  
238 but does not strictly infer all fluxes calculated this way were from bubbles entering the chamber.

## 239 2.4 Lake and Stream Metabolism

240 In 2024, we calculated the metabolic parameters net ecosystem production (NEP), gross  
241 primary production (GPP), and ecosystem respiration (ER) of oxygen for the lake and streams using  
242 dissolved oxygen (DO) collected at a one-minute frequency during chamber measurements (DO  
243 sensor accuracy:  $\pm 0.1 \text{ mg L}^{-1}$ ; resolution  $0.01 \text{ mg L}^{-1}$ ). Including metabolic parameters as predictors  
244 in the BRT models gave us an understanding of the role microbial oxygen production plays in  
245 regulating or not  $\text{CH}_4$  emissions from water. DO saturation was adjusted using temperature-  
246 dependent solubility constants (Garcia & Gordon, 1992). Oxygen flux at the air-water interface was  
247 determined using wind derived gas exchange coefficients and adjusted for water temperature (Cole  
248 & Caraco, 1998). NEP was calculated as the rate of change in DO concentration over each chamber  
249 deployment, adjusted for air-water exchange (Hall & Madinger, 2018; Noss et al., 2018). GPP and ER  
250 were partitioned from NEP by applying a threshold of  $200 \text{ } \mu\text{mol m}^{-2} \text{ s}^{-1}$  photosynthetically active  
251 radiation (PAR), which distinguishes intervals with effective photosynthesis from those with  
252 negligible light-driven production, despite continuous daylight during Arctic summer. Aggregating  
253 each chamber DO measurement into one-minute intervals, we calculated mean NEP, GPP, and ER for  
254 each chamber placement (Winslow et al., 2016). While this methodology is sound for the lake, there  
255 is some caveats in relation to stream metabolism. Because we are using a model which assumes  
256 wind driven gas exchange for small streams, we likely underestimate gas exchange in parts of the  
257 streams where turbulence from streambed roughness dominates. Nonetheless, the approach  
258 captures broadly the metabolic trends in lake and stream metabolism observed in other Arctic and  
259 Boreal waterbodies (Mulholland et al., 2001; Rocher-Ros et al., 2021; Ayala-Borda et al., 2024; Klaus  
260 et al., 2022; Myrstener et al., 2021) and is useful for comparing fluxes across aquatic biomes.

## 261 2.5 Spatial Flux Evaluation

262 We uploaded as a spatial layer in QGIS version 3.40.1 (QGIS Development Team, 2025) an  
263 orthomosaic image produced by Airbus satellite Pléiades 1B and the chamber placements as points  
264 with their associated flux estimates. We spatially analyzed the  $\text{CH}_4$  fluxes by creating bi-weekly  
265 emission heatmaps using the Kernel Density Estimation (KDE) algorithm in QGIS. The use of KDE  
266 allowed us to smooth across discrete chamber measurements, yielding an intuitive continuous  
267 surface representation of  $\text{CH}_4$  flux hotspots and their evolution through time (Figure 2). We set the  
268 radius to between 30-35 meters to allow some connectivity between points which allows for a  
269 smooth representation of any environmental gradients that might be captured. We used the default  
270 method using a quartic kernel shape weighted by the flux calculated for each chamber placement.  
271 While “ebullitive” fluxes were not considered in further statistical analysis (i.e., in the BRT), those  
272 fluxes are depicted in the resulting heatmaps. Overall, using KDE allowed for an intuitive  
273 interpretation of the seasonal and spatial development of flux hotspots in the research area.

274 We additionally compared daily  $\text{CH}_4$  emissions from Sanningasup Tasia catchment relative to  
275 other Arctic-Boreal Lake classes compiled in the Boreal-Arctic Wetland and Lake Dataset (Kuhn et al.  
276 2021; Olefeldt et al., 2021). A Kruskal-Wallis test was performed to determine significant differences  
277 ( $p < 0.05$ ) in the log transformed daily fluxes from Sanningasup Tasia ( $n = 48$ ) and its streams ( $n = 35$ )  
278 relative to broader biome-scale fluxes from Small Peat Lakes ( $n = 50$ ), Medium Peat Lakes ( $n = 36$ ),  
279 Large Lakes ( $n = 10$ ), and Small Yedoma Lakes ( $n = 7$ ). We then performed pairwise Wilcoxon rank-  
280 sum tests with Benjamini-Hochberg adjustment for multiple comparisons to highlight specific  
281 contrasts between Sanningasup Tasia catchment and the other lake classes. Conducting this

282 provides an understanding of how Sanningasup Tasia catchment emissions compares to other Arctic  
283 waterbody types.

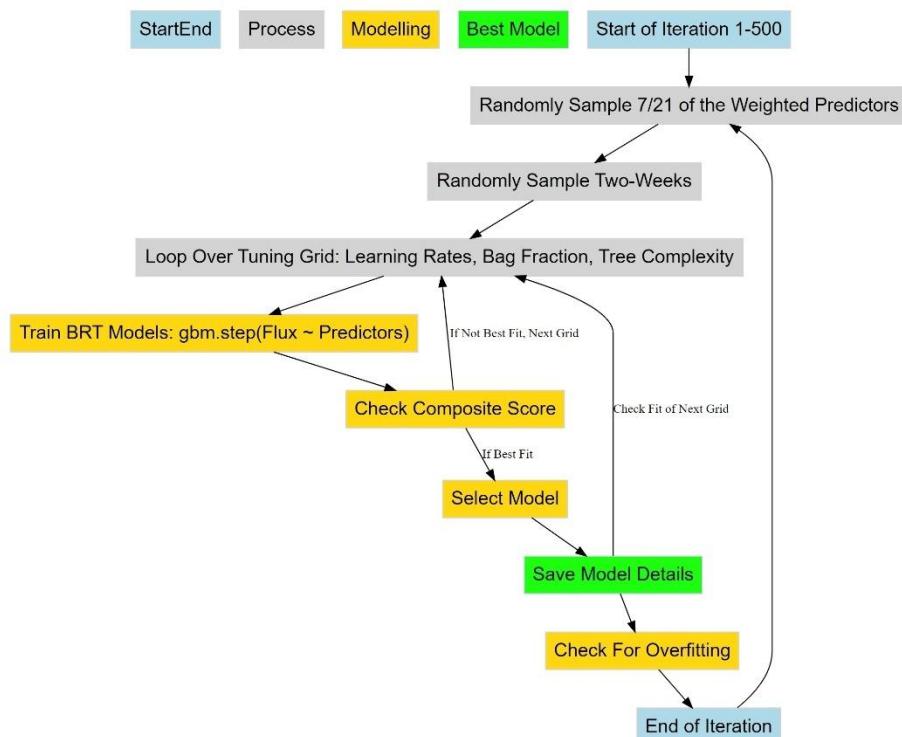
## 284 2.6 Decoding Methane Drivers

285 To determine important drivers and their partial effects on diffusive CH<sub>4</sub> fluxes, we trained  
286 BRT with physicochemical water parameters, catchment soil temperatures, catchment soil moisture,  
287 surface air temperatures, local meteorology, and Julian day (Figure 3). Considering we only collected  
288 water temperature in 2023, we used flux data from 2024 for training the BRT. We have focused on  
289 diffusive fluxes due to the unpredictability of fluxes when “ebullitive” processes were considered in  
290 the models. The diffusive fluxes give us a detailed view of environmental controls shaping CH<sub>4</sub> fluxes  
291 coming from the catchment. To characterize fluxes we assembled a comprehensive set of predictors  
292 including; (a) aquatic variables measured *in-situ* with a water sonde at each chamber deployment  
293 (e.g., Conductivity (µS/cm), pH, redox potential (mV), dissolved oxygen (mg/L), oxygen saturation  
294 (%), water temperature (°C), and fluorescent dissolved organic matter (FDOM; RFU)) (Figure S7), (b)  
295 catchment soil characteristics collected at nearby climate stations (e.g., soil volumetric water  
296 content at 10 cm and soil temperature at 40 cm), and (c) meteorological variables collected from a  
297 nearby climate station (e.g., Surface air temperature at 2 cm (°C), Air temperature (°C), relative  
298 humidity (%), air pressure (mbar), precipitation (mm), PAR (µmol m<sup>-2</sup>s<sup>-1</sup>), and mean wind speed (ms<sup>-1</sup>)  
299 and direction (°)). Lake water levels (mm) were included to characterize the effect of changing  
300 hydrologic conditions and its influence on lake CH<sub>4</sub> fluxes. Although water depth was recorded  
301 during chamber deployments using the AquaTroll 600, it was not included as a predictor in the BRT  
302 models. In shallow Arctic lakes like Sanningasup Tasia (<4.5 m), CH<sub>4</sub> is primarily sediment-derived,  
303 with deeper zones more likely to promote oxidation or dilution due to greater oxygen exposure  
304 (Bogard et al., 2014; Bulínová et al., 2025; Emerson et al., 2021; Li et al., 2020). Shoreline fluxes  
305 often dominate due to anoxic, vegetated sediments, while interior zones tend to suppress emissions  
306 (Thompson et al., 2016; Kyzivat et al., 2022; Rasilo et al., 2015). We therefore prioritized  
307 biogeochemical water column predictors—FDOM, dissolved oxygen saturation, and GPP—over  
308 depth (Christiansen et al., 2015; Singleton et al., 2018), and explicitly captured depth gradients via  
309 boat-based chamber deployments across the lake interior. Catchment soil characteristics were  
310 included to capture the hydrogeological conditions surrounding the catchment. We used catchment  
311 soil temperature at 40 cm to represent subsurface active-layer conditions that influence deeper  
312 thermal dynamics, groundwater inflow, and delayed soil heat retention through the thaw  
313 season. Soil volumetric water content (VWC) at 10 cm was included to gain an understanding if  
314 dryer, or wetter catchment conditions effect surface water CH<sub>4</sub> fluxes, and to act as a substitute for  
315 water level in the lake early in the season as these two share a Pearson’s correlation of  $r = 0.93$ .  
316 Additionally, we used VWC at 10 cm depth because it was the most complete and continuous  
317 dataset across the measurement depths, and highly correlated with VWC at 20 cm, 30 cm and 40  
318 cm. To reduce multicollinearity amongst the predictors, we set up weighting for random feature  
319 selection by calculating the average absolute Pearson correlations between predictors and assigning  
320 weights inversely proportional to the correlations, resulting in higher weights given to predictors  
321 with decreased collinearity and thus more likely to be included as a predictor.

322 Using the “gbm.step” algorithm from the R package “dismo” version 1.3.14 (Elith et al.,  
323 2008; Hijmans et al. 2023), we iteratively attempted to fit 500 BRT with a subset of randomly  
324 sampled two-week time series of flux calculations and 7 of the 21 weighted predictors. Each two  
325 weeks must have at least 90 observations, or the date range is buffered on either end of the time  
326 series to meet the minimum observation requirement. The algorithm uses 10-fold cross-validation to  
327 minimize overfitting the models (Elith et al., 2008). If a randomly sampled two weeks did not meet

328 the minimum required observation of 90, the time series was buffered on both ends of the date  
329 range to meet the minimum required observations. BRT were optimized using a grid search where  
330 hyperparameters such as learning rate (0.001, 0.002, 0.003, 0.004, 0.005), tree complexity (1, 3, 5,  
331 7), and bag fraction (0.30, 0.40, 0.50), were tuned for each model fit. While bag fraction values in the  
332 range of 0.5–0.8 are more commonly used, a lower bag fraction increases stochasticity in tree  
333 construction, which helps reduce overfitting—especially important for modeling noisy and highly  
334 non-linear CH<sub>4</sub> flux data. This conservative approach favors identifying robust general patterns  
335 rather than fitting noise or outliers. Variable monotonicity was handled dynamically for each subset  
336 of predictors using Spearman's rank correlation. Monotonicity for categorical variables was set to  
337 zero, while positive correlations were given a +1 and negative correlations were given a -1. The  
338 model with the best composite score was selected for each iteration. The composite score was  
339 calculated by adding together the standardized cross-validation error, standardized correlation  
340 error, and the cross-validation correlation. The model was finally saved after checking for over fitting  
341 by taking the difference between cross validated mean deviance and training mean deviance and  
342 dividing the difference by training mean deviance. Because the inherently noisy nature of ecological  
343 data, we allowed for 40% difference between cross validated predictions and training data. We  
344 further calculated the percent deviance explained for each BRT model using the formula: % deviance  
345 explained = 100 × ((null deviance – residual deviance) / null deviance), where the null deviance  
346 represents the deviance of a model using only the mean response, and the residual deviance is from  
347 the fitted BRT model. Each fitted model and its metadata were saved for further analysis. This  
348 modelling structure ensures robustness against outliers and ensures data integrity through  
349 dynamically handling monotonicity and applying overfitting constraints. Furthermore, the structure  
350 ensures robust predictions of fluxes by accounting for multi-collinearity amongst predictors and flux  
351 heterogeneity throughout the season.

352 To visualize the results of the models, we plotted partial dependence two ways. First, we  
353 extracted partial dependence information for understanding model structure, i.e., those predictors  
354 and interactions which were used to split trees and decrease cross validated prediction error. In  
355 addition, we made isolated predictions for each environmental feature in the model by holding all  
356 other predictors at their median to gain a more mechanistic understanding of which conditions  
357 and/or processes are directly affecting CH<sub>4</sub> fluxes. The two ways of visualizing partial dependence  
358 give us an ecological understanding of how integrated direct and indirect effects regulate fluxes from  
359 the catchment, but also how individual variables and/or processes regulate fluxes from the water  
360 surfaces, respectively. All visualizations were generated using the R package “ggplot2” version 3.5.0  
361 (Wickham, 2016), and the package “DiagrammeR” version 1.0.11 for flowcharts (Iannone, 2024).



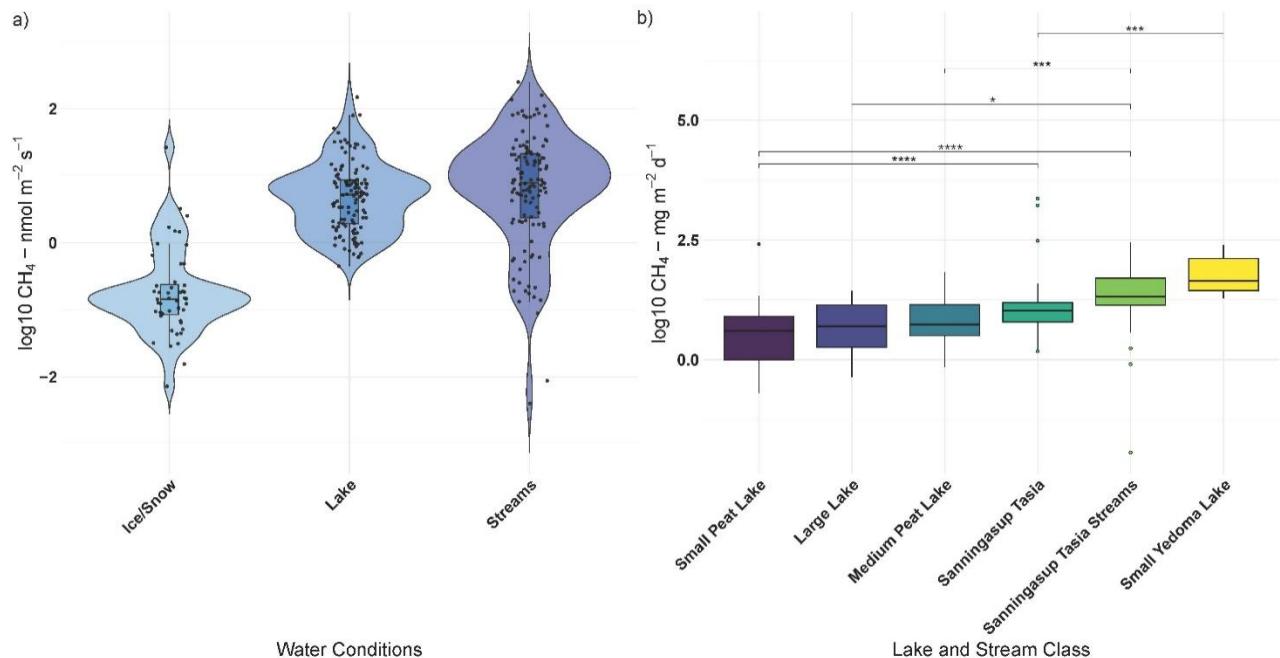
364 Figure 3. Workflow of the iterative randomized process for selecting the best BRT for predicting  
 365 diffusive CH<sub>4</sub> fluxes using various environmental predictors. Blue boxes represent the start and end  
 366 of a single iteration through an index of weighted predictors, time periods, and tuning grid (grey  
 367 boxes). Yellow boxes represent the model selection logic, while the green box represents the storage  
 368 of flux predictions and selected model details for the included predictors and two-week sub-sample  
 369 of chamber measurements.

### 370 3 Results and Discussion

#### 371 3.1 Overview

372 Methane fluxes from permafrost affected catchments are influenced by a complex interplay  
 373 between climatological, hydrogeological, and biogeochemical processes. This study highlights the  
 374 transient nature of CH<sub>4</sub> fluxes from a permafrost catchment in west central Greenland and the  
 375 partial effects of physiochemical water conditions, local meteorology and catchment conditions  
 376 (Figures 4-7). CH<sub>4</sub> emissions from the catchment were variable across water conditions, with streams  
 377 exhibiting the highest emissions (Figure 4). In comparison to the global coverage of the Boreal-Arctic  
 378 Wetland and Lake Dataset (Olefeldt et al., 2021), besides small peat lakes, daily fluxes from  
 379 Sanningasup Tasia at 8.3 mg m<sup>-2</sup> d<sup>-1</sup> were mostly comparable to other permafrost waterbodies across  
 380 the Arctic-Boreal region, which ranges between 3.8-5.4 mg m<sup>-2</sup> d<sup>-1</sup> (Figure 4). Highlighting the  
 381 importance of emissions from small streams, Sanningasup Tasia streams showed significantly higher  
 382 daily fluxes (18.2 mg m<sup>-2</sup> d<sup>-1</sup>) than all inland water body classes, except Yedoma lakes (43.7 mg m<sup>-2</sup> d<sup>-1</sup>). Our results indicate that CH<sub>4</sub> fluxes were seasonally variable and controls on fluxes shifted from  
 383 hydroclimatic factors during colder periods to biogeochemical processes as the catchment warmed  
 384 and increased in productivity (Figure 3-4 and Figure A1). The seasonal thaw of annual snow and ice  
 385 accumulation in the two study years varied in timing and duration due to 2023 staying anomalously  
 386 snowy until the beginning of July, where in 2024 the number of snow free days aligned with  
 387 historical records. In 2023, our initial chamber measurements between July 03-15 captured peak  
 388

389 median fluxes at  $8.9 \text{ nmol m}^{-2} \text{s}^{-1}$  just as the ice began to break on the lake. In an effort to capture  
 390 similar conditions in 2024, we used an index of historical snow free days on the island which led to  
 391 us capturing median fluxes of  $0.18 \text{ nmol m}^{-2} \text{s}^{-1}$  between May 24-June 05 atop ice and snow. Peak  
 392 median fluxes of  $8.1 \text{ nmol m}^{-2} \text{s}^{-1}$  in 2024, were comparable to 2023, but occurred a month earlier  
 393 between June 11-19. As runoff water receded and the catchment warmed, growing season  
 394 commenced in conjunction with steadily decreasing median fluxes between  $3.9$  and  $4.5 \text{ nmol m}^{-2} \text{s}^{-1}$ .  
 395 This study further provides methods to disentangle important drivers and their partial effects on  $\text{CH}_4$   
 396 fluxes using BRT (Figure 3-4). In general,  $\text{CH}_4$  fluxes were strongly dependent on discrete oxic-anoxic  
 397 aquatic environments under the chamber (Figure 4 and Figure A1). This research emphasizes the  
 398 importance of integrating field measurements with GIS-based spatial analysis to monitor  $\text{CH}_4$  fluxes  
 399 in permafrost catchments. It further drives home the importance of seasonal transition periods in  
 400 predicting fluxes from Arctic waterbodies.

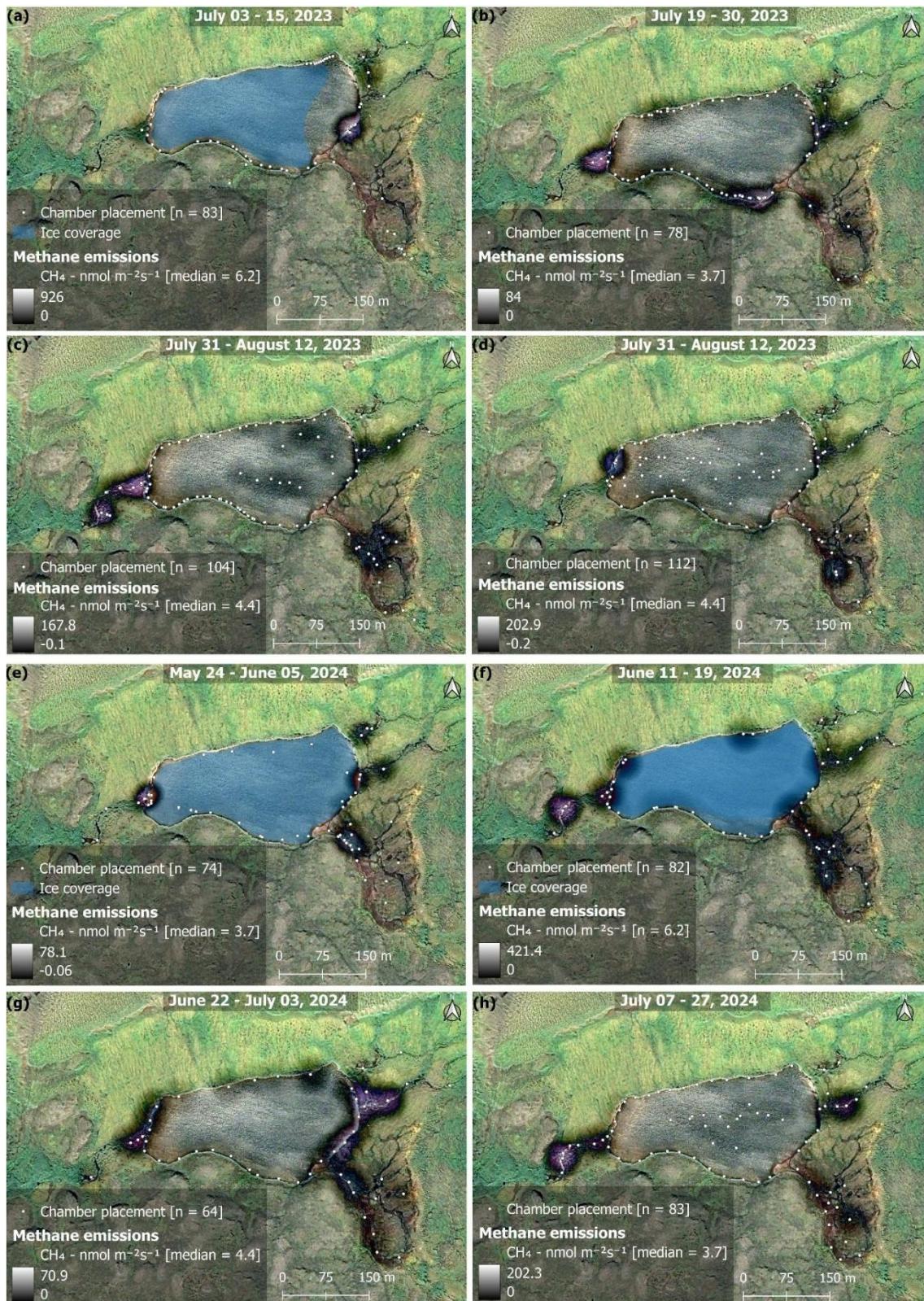


401  
 402 Figure 4. Panel a) shows log transformed  $\text{CH}_4$  fluxes across the different catchment water conditions  
 403 during 2024 field season and b) compares log transformed daily  $\text{CH}_4$  fluxes (y axis) between  
 404 Sanningasup Tasia catchment and other permafrost waterbodies (Kuhn et al., 2021) across the  
 405 Arctic-boreal region. Connecting brackets and stars show, for example, that Sanningasup Tasia had  
 406 significantly ( $p < 0.05$ ) higher daily emissions when compared to Small Peat Lakes, while daily  
 407 emissions from Sanningasup Tasia Streams were significantly higher than all lake classes except  
 408 Yedoma Lakes.

### 409 3.2 Spatial and Temporal Evolution of Methane Fluxes

410 In both 2023 and 2024, spatial and temporal evolution of fluxes occurred somewhat  
 411 heterogeneously in the catchment, but generally “hotspots” occurred in the streams and where they  
 412 enter the lake (Figure 5 a-b, e-f). Despite different time periods of the thaw, fluxes in the catchment  
 413 in both years followed a similar trajectory, with peak fluxes occurring post thaw and decreasing  
 414 through the growing season. In 2023, snow persisted anomalously late into the summer season, and  
 415 soil temperatures were the coldest recorded in a 6-year record (Figure S8). We found that local  
 416 climate and catchment soil characteristics were at times, (i.e., during the thaw season and towards  
 417 the peak of growing season) more important than water temperature in predicting 2023 fluxes,  
 418 suggesting catchment contributions to surface waters plays an indirect role in  $\text{CH}_4$  fluxes (Figure S9).

419 While water temperature was found to be relatively important in both years, catchment CH<sub>4</sub> fluxes in  
 420 2024 suggest the system is more driven by variability in dissolved organic matter and microbial  
 421 production of oxygen (Figure 4-7 and Figure A1).



422

423 Figure 5. Maps showing the bi-weekly spatial evolution of CH<sub>4</sub> emissions during 2023 (a-d) and 2024  
 424 (e-h). The time period covered in each map is given at the top of each map. Whiter colors represent

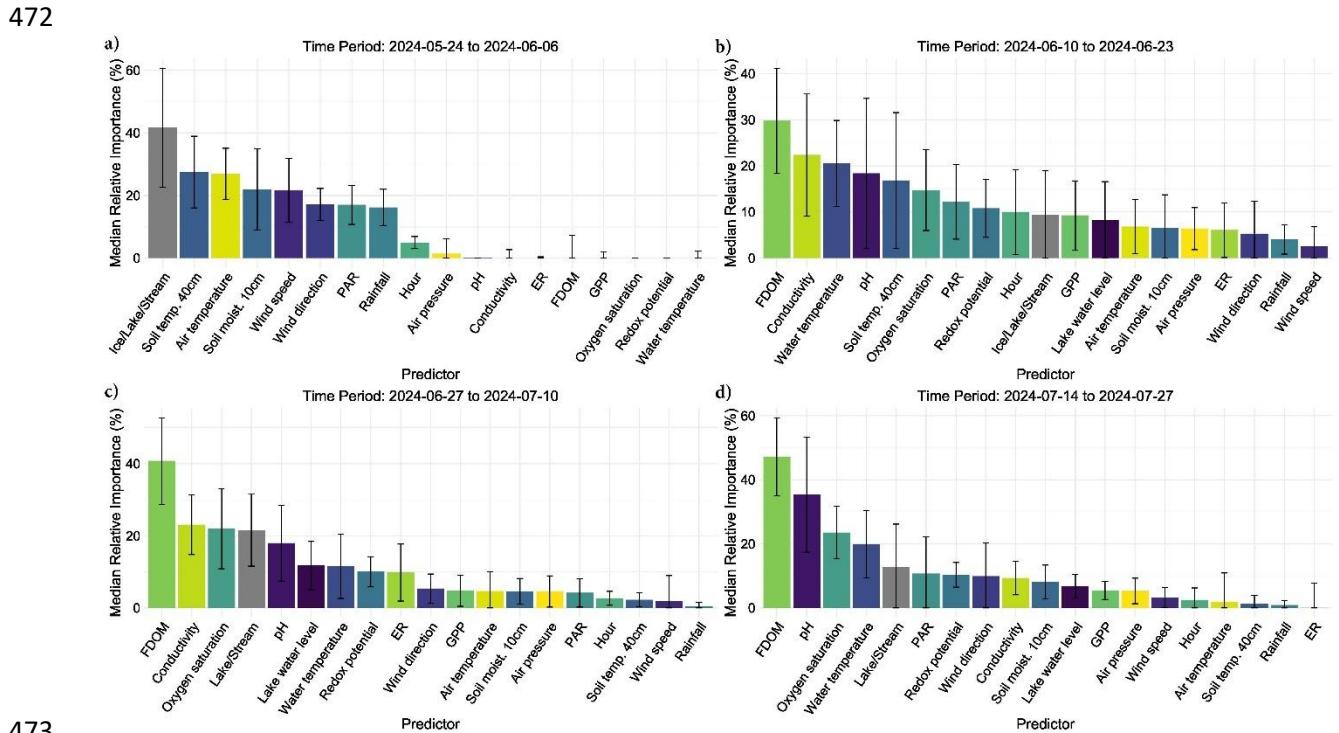
425 areas of high emissions, while darker, or no color represent areas of lower, or no emissions (see  
426 color bars in map legends). Areas outside of the water were not measured and are artifacts from the  
427 KDE mapping algorithm. Each white point on the map represents the placement of the floating  
428 chamber. Orthomosaic background image © CNES (2024), Distribution Airbus DS, produced from  
429 Pléiades 1B satellite imagery.

430 The main inlet to the lake is a warm spring with median temperatures of 7.4 °C and tended  
431 to be a persistent location throughout the season for increased CH<sub>4</sub> fluxes. The warm spring area in  
432 the southeast is a complex area where a perennial spring bubbles out of the ground forming a  
433 perennial tributary to the lake. However, the spring seeps out along the base of the hillslope to the  
434 east, subsequently forming a peat fen environment (Figure 5). The eastern most inlets are fed by  
435 meltwater flowing through the vadose zone, but also served as a persistent location for increased  
436 CH<sub>4</sub> fluxes. Fluxes from inlet streams followed along an upland-wetland environmental gradient,  
437 where fluxes at the headwaters of streams were generally close to 0, or slightly taking up CH<sub>4</sub>, but  
438 steadily increased as steeper upland slopes gave way to more gradual water saturated slopes and  
439 pooling sediments. This is consistent with previous work on the island showing CH<sub>4</sub> fluxes association  
440 with soil microbial assemblages shifting from methanotrophic to methanogenic along an upland-  
441 wetland gradient, respectively(Christiansen et al., 2015). Additionally, the catchment serves as  
442 micro-topography previously described as an area of snowmelt retention, and subsequently an area  
443 of nutrient and/or dissolved organic matter pooling (Westergaard-Nielsen et al., 2020), which has an  
444 impact on CH<sub>4</sub> fluxes throughout the season (Figure 7). While high fluxes were recorded along the  
445 shore and in the open water of the lake, fluxes tended to be patchy and decreased moving away  
446 from the inlet streams (Figure 5 and Figure S10). However, as soon as the lake water flowed to the  
447 outlet, fluxes increased substantially. As the summer season progressed, CH<sub>4</sub> fluxes declined across  
448 most of the catchment, becoming largely confined to the warm spring inlet and the eastern inlet  
449 streams (Figure 5). Field observations of late-season fluxes in 2024, found decreased fluxes were  
450 associated with submerged filamentous green algae in stream channels, while assemblages of iron-  
451 oxidizing bacteria on the stream banks were associated with increased fluxes, (Figure A2). The  
452 spatial and temporal evolution of fluxes was driven by seasonally shifting environmental conditions.

### 453 3.3 Boosted Regression Tree Results

454 Out of 500 iterations, 321 BRT were fit, and showed good alignment and consistently  
455 performed well in cross-validation, with a correlation median of 0.40 between observed and  
456 predicted values, and a median deviance standard error of 131. Between 8.4% and 62.4% with a  
457 median of 27.3% of the CH<sub>4</sub> flux variability was explained by the various models and included  
458 environmental conditions, suggesting a substantial proportion of CH<sub>4</sub> fluxes were explained by the  
459 environmental conditions included. The calculated root mean squared error (RMSE), which reflects  
460 the average magnitude in prediction error of the BRT, ranged between 6.5 to 28 nmol m<sup>-2</sup>s<sup>-1</sup>, with a  
461 median of 13.7 nmol m<sup>-2</sup>s<sup>-1</sup>. The summary statistics reflect models that performed reliably and with  
462 fairly good accuracy in predicting diffusive CH<sub>4</sub> fluxes from the catchment in 2024. The models  
463 predicted shifting relative importance (Figure 6) and partial effects of the various environmental  
464 conditions throughout the season (Figure 7). The magnitude of CH<sub>4</sub> fluxes predicted by the BRT  
465 models were strongly influenced by localized biogeochemical conditions within the water column  
466 based on whether the flux was originating from the lake, stream, or if it was influenced by ice or  
467 snow (Figure 4-7, and Figure A1a-d). Visualizing partial dependence of predictors important for  
468 model structure, revealed integrated ecological effects between local meteorology, catchment  
469 conditions and physiochemical water conditions (Figure 7). However, isolated direct marginal effects

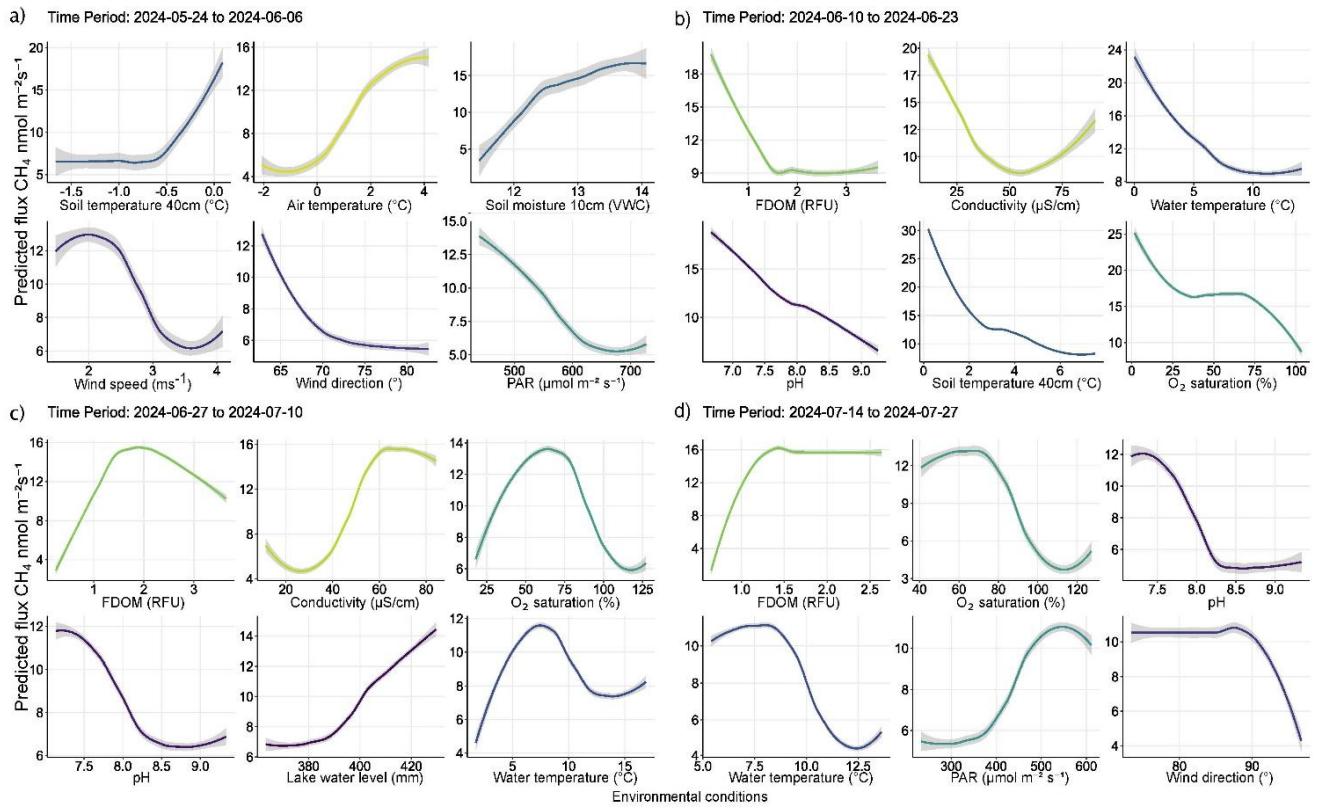
470 of the various environmental conditions suggest fluxes from water surfaces are directly regulated via  
 471 biochemical processes associated with GPP and ER of oxygen (Figure A1b-d).



473  
 474 Figure 6. The figure illustrates the relative importance of environmental conditions predicting  
 475 diffusive  $\text{CH}_4$  fluxes using bar-plots and standard error bars. Each predictor variable is on the x-axis,  
 476 while its percent importance for its inclusion in a fitted BRT is given on the y-axis where the standard  
 477 error bars represent variability in importance based on presence or absence in a given model. Each  
 478 bar color represents a distinct environmental condition.

### 479 3.3.1 Spring Thaw Phase: Peak Fluxes Driven by Hydrological and Climatic Controls

480 The spring thaw phase marks a shift in catchment conditions, where a frozen landscape gives  
 481 way to thaw season and hydro-connectivity between land and water is strong (Figure 5c-d, g-h). In  
 482 the spring thaw phase, initial peak fluxes of  $\text{CH}_4$  were primarily dependent on increasing rainfall,  
 483 changing wind conditions, warming air and soil temperatures, and increased soil moisture content,  
 484 while low dissolved organic matter (i.e., FDOM) indicated increased fluxes (Figure 6a-b and 7a-b).  
 485 Soil moisture was found to have a Pearson's  $r = 0.93$  with lake water levels, suggesting the lake levels  
 486 are strongly connected to snowmelt and groundwater hydrology (Figure S11). The distribution of  
 487 nutrients on the island has been shown to be linked to snowmelt and hill slope topography  
 488 (Westergaard-Nielsen et al., 2020), which is likely playing a role during the early part of the season,  
 489 but especially later in the year as DOM, a proxy for nutrients, becomes the primary limiting factor in  
 490 predicting higher fluxes (Figure 7b-c) (Olid et al., 2021, 2022). The processes driving  $\text{CH}_4$  fluxes from  
 491 water surfaces is likely two-fold.



492

493 Figure 7. Partial dependency plots illustrating the indirect and direct marginal effects of various  
 494 environmental conditions predicting diffusive CH<sub>4</sub> fluxes. Each panel displays the effect of a  
 495 numerical predictor used during model training to predict CH<sub>4</sub> fluxes. Although some predictors may  
 496 not have been directly involved in regulating fluxes from water surfaces, their evaluation reveals the  
 497 relationship between water column conditions and catchment processes regulating CH<sub>4</sub> fluxes. The  
 498 figures are ordered by importance in each time period and the colors correspond to those seen in  
 499 Figure 6. The colored lines represent the result of a fitted general additive model ( $y \sim s(x)$ ) and thus  
 500 a smooth representation across the 321 fitted BRT models. The grey shaded area around the line  
 501 represents  $\pm SE$  (0.02 - 0.2). Each predictor and its numerical range are given on the x-axis, while  
 502 predicted fluxes are given on the y-axis.

503 As snowfall turned to rain, the thawing of soils accelerated and water content in the active  
 504 layer increased, potentially driving peak emissions via the lateral mobilization of dissolved CH<sub>4</sub>  
 505 toward surface waters (Figure 7a-b) (Walter Anthony et al., 2012; Neumann et al., 2019; Olid et al.,  
 506 2022). However, as the thaw progressed, contributions to fluxes from catchment soils decreased as  
 507 the upper layers began to dry and lake water levels reached their maximum (Figure 7b-c). DOM  
 508 serves as a critical substrate for both CH<sub>4</sub> production and oxidation, particularly in permafrost-  
 509 influenced regions where thawing can release large amounts of particulate matter (Keskitalo et al.,  
 510 2021; Bouranis et al. 2025). Anaerobic conditions in water saturated soils and low oxygenated  
 511 waters may have further driven fluxes during the thaw period as methanogenic microbial  
 512 communities rapidly consumed incoming labile organic compounds (Neumann et al., 2019;  
 513 Stevenson et al., 2021). However, as the thaw progressed, the lake briefly shifted to net-autotrophy  
 514 (Figure S6), marking an important shift from hydroclimatic controls on fluxes, to a patchwork of  
 515 biochemical transport pathways and barriers (Figure 7 b-d and Figure A1b-d). During this transition,  
 516 increased GPP and the resulting oxygen saturation in the lake and streams enhance methanotrophic  
 517 activity, thereby promoting methane oxidation and reducing net CH<sub>4</sub> emissions (Figure 7b-c and  
 518 Figure A1b-c). The central role of oxygen availability and active methanotroph communities in

519 regulating methane fluxes during Arctic thaw has been demonstrated in both tundra landscapes and  
520 permafrost mires (Christiansen et al., 2015; Graef et al., 2011; Singleton et al., 2018). As oxygen  
521 production increases, greater amounts of oxygen become available to support methanotrophic  
522 activity in the water column. Methanotrophic activity is likely represented in the observed reduction  
523 of CH<sub>4</sub> fluxes during periods of decreased ER (i.e., when oxygen consumption is elevated, Figure A1c),  
524 as higher oxygen availability enhances aerobic methane oxidation (Figure 7c). However, the streams  
525 remain net heterotrophic despite increasing GPP and ER during this time period, therefore providing  
526 a more suitable environment for CH<sub>4</sub> production and flux (Figure S6). Thus, the lake and streams  
527 reflect distinct ecosystems for the production and flux of CH<sub>4</sub>, where the lake being an overall  
528 methane source, has a greater tendency to oxidize CH<sub>4</sub> following the thaw, while the streams follow  
529 along a low-to-high flux gradient as the slope flattens and soils become saturated (Westergaard-  
530 Nielsen et al., 2020; Stevenson et al., 2021).

### 531 3.3.2 Growing Season: Oxic-Anoxic Transport Pathways and Barriers

532 Early in the growing season the lake and streams enter into a transition phase, where peak  
533 fluxes become increasingly dependent on biochemical pathways related to DOM availability,  
534 conductivity, oxygen saturation and pH (Figure 7c). DOM is the limiting factor late in the season  
535 where higher fluxes are increasingly dependent on water column conditions with greater levels of  
536 DOM (Figure 6c-d and 7c-d). CH<sub>4</sub> fluxes during this time period were strongly associated with  
537 indicators of microbial activity forming oxic-anoxic transport barriers, or pathways, respectively. For  
538 example, microbial activity in anoxic sediments maybe producing CH<sub>4</sub>, but whether it is diffused to  
539 the atmosphere is directly affected by the micro-conditions of the water column (e.g., DOM,  
540 conductivity, pH, and/or GPP/oxygen saturation), either forming an oxidative barrier, or an anoxic  
541 pathway (Figure 7c-d and Figure A1). For example, groundwater transport of CH<sub>4</sub> from anoxic  
542 sediments to oxygen-rich streams may result in water with high concentrations of both. While some  
543 of the CH<sub>4</sub> is likely oxidized during transport, it can be that both are respiration at turbulent sections of  
544 the stream, which were the highest fluxes observed from the streams and during this phase of the  
545 season (Street et al., 2016; Neumann et al., 2019; Olid et al., 2022; Kleber et al. 2025).

546 Fluxes are further affected by water conditions either favoring methanogenic, or  
547 methanotrophic activity (Conrad, 2007; Cunada et al., 2021; Emerson et al., 2021). pH levels near  
548 neutral likely indicate water conditions favorable to methanogenesis at the sediment-water  
549 interface, while increasing alkalinity may reduce methanogenic and/or favor increased  
550 methanotrophic activity as growing season progresses (Figure 7c-d). For example, during growing  
551 season micro-pH and oxygen saturation conditions in the lake and streams are influenced by the  
552 ever-increasing presence of macrophytes, mosses and plankton, which tend to drive pH and oxygen  
553 levels higher (Liebner et al., 2011; Cunada et al., 2021). Here we show that increasing pH and oxygen  
554 saturation, as a result of primary production, create an aerobic environment that favors  
555 methanotrophic activity, thereby driving CH<sub>4</sub> emissions down through the growing season (Figure  
556 7c-d, Figure A1). Declines in oxygen saturation driven by microbial respiration can create anoxic  
557 conditions that enable CH<sub>4</sub> emissions from sediment to surface waters (Conrad, 2007; Michel et al.,  
558 2010; Street et al., 2016; Cheng et al. 2024). Such a mechanism likely explains the formation of flux  
559 hotspots associated with decomposing iron-oxidizing bacterial mats along stream banks (Figure A2)  
560 (Wallenius et al., 2021; Cheng et al. 2024). In the case of the bacterial mats, we observed fluxes  
561 were highest in the streams where bacterial assemblages had become exposed to the atmosphere  
562 and were decomposing in stagnant water (Figure A2), which may suggest that the decomposition of  
563 the bacteria was releasing dissolved organic substrates in a low-oxygen environment already primed  
564 for methanogenic activity (Wallenius et al., 2021; Cheng et al. 2024). This idea is supported late in

565 the season when increased CH<sub>4</sub> emissions become dependent on niche environments where  
566 moderate levels of dissolved organic matter (FDOM) and low oxygenated water predict higher fluxes  
567 (Figure 7d and Figure A1d). However, submerged bacterial mats along with filamentous green algae  
568 the presence of macrophytes and/or mosses in the lake and streams, were associated with lower  
569 fluxes, suggesting they form oxidative barriers for CH<sub>4</sub> fluxes from the sediment when submerged  
570 (Figure 7d and Figure A1d) (Heilman & Carlton, 2001; Laanbroek, 2010; Liebner et al., 2011; Esposito  
571 et al., 2023). The submerged, or not status of bacterial assemblages' points to an interesting  
572 feedback mechanism between CH<sub>4</sub> fluxes and dropping water levels creating variability in emission  
573 pathways. Similar processes have been shown in relation to submerged brown mosses in Arctic  
574 tundra ecosystems have been shown to promote CH<sub>4</sub> oxidation and thus decrease CH<sub>4</sub> emissions  
575 from sediments (Žárský et al., 2018). Overall, these results highlight the broader importance of fine-  
576 scale biogeochemical dynamics shaping CH<sub>4</sub> fluxes from a permafrost catchment and provide an  
577 important data point in an uncertain region of the world.

## 578 4 Conclusion

579 This research provides a temporally resolved catchment scale CH<sub>4</sub> flux analysis across  
580 different waterbody types and conditions—lake, streams, and ice/snow-covered surfaces—  
581 subsequently describing important biogeochemical and climatic controls on emissions. Often lost in  
582 temporally coarse assessments is a detailed understanding of seasonal transitions in processes  
583 related to CH<sub>4</sub> fluxes and environmental control mechanisms. Leveraging BRT to fit hundreds of  
584 randomized models and visualizing the direct, and indirect controls on CH<sub>4</sub> fluxes reveals variability  
585 in how, for example, DOM and/or water temperature affect fluxes differently as the Arctic summer  
586 progresses. We presented an approach which captures ecosystem-scale effects, but furthermore  
587 describes isolated mechanistic effects related to, for example, GPP, revealing that primary  
588 productivity plays a critical role in regulating CH<sub>4</sub> emissions from permafrost affected waterbodies.  
589 This work contributes to understanding carbon feedback mechanisms in a region where process-  
590 level knowledge is needed to scale global models simulating CH<sub>4</sub> emissions from permafrost affected  
591 waterbodies.

## 592 Code Availability

593 R code for calculating methane fluxes can be found here: <https://github.com/mthayne527/fluxCH4>.

## 594 Data Availability

595 Meteorological can be accessed here: <https://doi.org/10.17897/FEGK-0632>, and soil data here:  
596 <https://doi.org/10.17897/6G78-P793>, <https://doi.org/10.17897/9N7Z-GA63>, and can be  
597 accessed via the Greenland Ecosystem Monitoring website: <https://data.g-e-m.dk/datasets?theme=climate>. BAWLD circum-Arctic waterbody dataset can be found here: DOI:  
598 10.5194/essd-13-5151-2021. Water chemistry and chamber data can be requested from Torsten  
599 Sachs at Helmholtz Centre for Geosciences in Potsdam, Germany (GFZ).

## 601 Author contributions

602 MWT collected, compiled and analyzed data, and wrote the manuscript. KK collected,  
603 compiled, and analyzed data, and contributed writing parts of the manuscript. CW provided  
604 methodological guidance and feedback, and contributed to writing parts of the methodology. AK  
605 provided comments, feedback, and guidance on interpreting results, and contributed to writing

606 various parts of the manuscript. TS collected data, provided comments, feedback, and guidance on  
607 interpreting results, and contributed to writing the manuscript.

## 608 Competing interests

609 The authors declare that they have no conflict of interest.

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## 621 References

622 Ayala-Borda, P., Bogard, M. J., Grosbois, G., Preskienis, V., Culp, J. M., Power, M., &  
623 Rautio, M. (2024). Dominance of net autotrophy in arid landscape low relief polar lakes,  
624 Nunavut, Canada. *Global Change Biology*, 30(2), e17193. DOI: 10.1111/gcb.17193

625

626 Bastviken, D., Cole, J., Pace, M., & Tranvik, L. (2004). Methane emissions from lakes:  
627 Dependence of lake characteristics, two regional assessments, and a global estimate. *Global  
628 Biogeochemical Cycles*, 18(4). <https://doi.org/10.1029/2004GB002238>

629

630 Bastviken, D., Cole, J. J., Pace, M. L., & van de-Bogert, M. C. (2008). Fates of methane from  
631 different lake habitats: Connecting whole-lake budgets and CH<sub>4</sub> emissions. *Journal of  
632 Geophysical Research: Biogeosciences*, 113(2). <https://doi.org/10.1029/2007JG000608>

633 Bartsch, A., Gay, B. A., Schüttemeyer, D., et al. (2025). Advancing the Arctic Methane  
634 Permafrost Challenge (AMPAC) with future satellite missions. *IEEE Journal of Selected  
635 Topics in Applied Earth Observations and Remote Sensing*, 15, 1234–1245. DOI:  
636 10.1109/JSTARS.2025.3538897

637 Bogard, M. J., del Giorgio, P. A., Boutet, L., Chaves, M. C. G., Prairie, Y. T., Merante, A., &  
638 Derry, A. M. (2014). Oxic water column methanogenesis as a major component of aquatic  
639 CH<sub>4</sub> fluxes. *Nature Communications*, 5. <https://doi.org/10.1038/ncomms6350>

640 Bouranis, J. A., McGivern, B. B., Makke, G., et al. (2025). Metabolic redox coupling controls  
641 methane production in permafrost-affected peatlands through organic matter  
642 quality-dependent energy allocation. *Global Change Biology*, 31(8), e70390.  
643 <https://doi.org/10.1111/gcb.70390>

644 Bulínová, M., Schomacker, A., Kjellman, S. E., Gudasz, C., Olid, C., Rydberg, J., Panieri, G.,  
645 et al. (2025). Increased ecosystem productivity boosts methane production in Arctic lake

646 sediments. *Journal of Geophysical Research: Biogeosciences*, 130(7), e2024JG008508.  
647 <https://doi.org/10.1029/2024JG008508>

648 Emerson, J. B., Varner, R. K., Wik, M., Parks, D. H., Neumann, R. B., Johnson, J. E.,  
649 Singleton, C. M., Woodcroft, B. J., Tollerson, R., Owusu-Dommey, A., Binder, M., Freitas,  
650 N. L., Crill, P. M., Saleska, S. R., Tyson, G. W., & Rich, V. I. (2021). Diverse sediment  
651 microbiota shape methane emission temperature sensitivity in Arctic lakes. *Nature  
652 Communications*, 12(1). <https://doi.org/10.1038/s41467-021-25983-9>

653

654 Callaghan, T. v., Christensen, T. R., & Jantze, E. J. (2011). Plant and vegetation dynamics on  
655 Disko island, west Greenland: Snapshots separated by over 40 years. In *Ambio* (Vol. 40, Issue  
656 6). <https://doi.org/10.1007/s13280-011-0169-x>

657 Cheng, S., Meng, F., Wang, Y., Zhang, J., & Zhang, L. (2024). The potential linkage between  
658 sediment oxygen demand and microbes and its contribution to the dissolved oxygen depletion  
659 in the Gan River. *Frontiers in Microbiology*, 15, 1413447.  
660 <https://doi.org/10.3389/fmicb.2024.1413447>

661 Christiansen, H. H. (1999). Active layer monitoring in two Greenlandic permafrost areas:  
662 Zackenberg and Disko Island. *Geografisk Tidsskrift*, 99.

663

664 Christiansen, J. R., Romero, A. J. B., Jørgensen, N. O. G., Glaring, M. A., Jørgensen, C. J.,  
665 Berg, L. K., & Elberling, B. (2015). Methane fluxes and the functional groups of  
666 methanotrophs and methanogens in a young Arctic landscape on Disko Island, West  
667 Greenland. *Biogeochemistry*, 122(1). <https://doi.org/10.1007/s10533-014-0026-7>

668

669 Cole, J. J., & Caraco, N. F. (1998). Atmospheric exchange of carbon dioxide in a low-wind  
670 oligotrophic lake measured by the addition of SF6. *Limnology and Oceanography*, 43(4).  
671 <https://doi.org/10.4319/lo.1998.43.4.0647>

672 Conrad, R. (2007). Microbial Ecology of Methanogens and Methanotrophs. In *Advances in  
673 Agronomy* (Vol. 96). [https://doi.org/10.1016/S0065-2113\(07\)96005-8](https://doi.org/10.1016/S0065-2113(07)96005-8)

674 Cunada, C. L., Lesack, L. F. W., Tank, S. E., & Hesslein, R. H. (2021). Methane flux  
675 dynamics among CO<sub>2</sub>-absorbing and thermokarst lakes of a great Arctic delta.  
676 *Biogeochemistry*, 156(1), 25–39. <https://doi.org/10.1007/s10533-021-00853-0>

677

678 D'Imperio, L., Nielsen, C. S., Westergaard-Nielsen, A., Michelsen, A., & Elberling, B.  
679 (2017). Methane oxidation in contrasting soil types: responses to experimental warming with  
680 implication for landscape-integrated CH<sub>4</sub> budget. *Global Change Biology*, 23(2).  
681 <https://doi.org/10.1111/gcb.13400>

682

683 Elder, C. D., Thompson, D. R., Thorpe, A. K., Hanke, P., Walter Anthony, K. M., & Miller,  
684 C. E. (2020). Airborne Mapping Reveals Emergent Power Law of Arctic Methane Emissions.  
685 *Geophysical Research Letters*, 47(3). <https://doi.org/10.1029/2019GL085707>

686

687 Elith, J., Leathwick, J. R., & Hastie, T. (2008). A working guide to boosted regression trees.  
688 In *Journal of Animal Ecology* (Vol. 77, Issue 4). <https://doi.org/10.1111/j.1365-2656.2008.01390.x>

689

690 Emerson, J. B., Varner, R. K., Wik, M., Parks, D. H., Neumann, R. B., Johnson, J. E.,  
691 Singleton, C. M., Woodcroft, B. J., Tollerson, R., Owusu-Domme, A., Binder, M., Freitas,  
692 N. L., Crill, P. M., Saleska, S. R., Tyson, G. W., & Rich, V. I. (2021). Diverse sediment  
693 microbiota shape methane emission temperature sensitivity in Arctic lakes. *Nature  
694 Communications*, 12(1). <https://doi.org/10.1038/s41467-021-25983-9>

695

696 Esposito, C., Nijman, T. P. A., Veraart, A. J., Audet, J., Levi, E. E., Lauridsen, T. L., &  
697 Davidson, T. A. (2023). Activity and abundance of methane-oxidizing bacteria on plants in  
698 experimental lakes subjected to different nutrient and warming treatments. *Aquatic Botany*,  
699 185. <https://doi.org/10.1016/j.aquabot.2022.103610>

700

701 Fazi, S., Amalfitano, S., Venturi, S., Pacini, N., Vazquez, E., Olaka, L. A., Tassi, F.,  
702 Crognale, S., Herzsprung, P., Lechtenfeld, O. J., Cabassi, J., Capecchiacci, F., Rossetti, S.,  
703 Yakimov, M. M., Vaselli, O., Harper, D. M., & Butturini, A. (2021). High concentrations of  
704 dissolved biogenic methane associated with cyanobacterial blooms in East African lake  
705 surface water. *Communications Biology*, 4(1). <https://doi.org/10.1038/s42003-021-02365-x>

706 Fox-Kemper, B., Hewitt, H. T., Xiao, C., et al. (2021). Chapter 9: Ocean, cryosphere and sea  
707 level change. In: Climate Change 2021: The Physical Science Basis. *Contribution of Working  
708 Group I to the Sixth Assessment Report of the Intergovernmental Panel on Climate Change*.  
709 Cambridge University Press, 1211–1362. <https://doi.org/10.1017/9781009157896.011>

710 Garcia, H. E., & Gordon, L. I. (1992). Oxygen solubility in seawater: Better fitting equations.  
711 In *Limnology and Oceanography* (Vol. 37, Issue 6).  
712 <https://doi.org/10.4319/lo.1992.37.6.1307>

713

714 Graef, C., Hestnes, A. G., Svenning, M. M., & Frenzel, P. (2011). The active methanotrophic  
715 community in a wetland from the High Arctic. *Environmental Microbiology Reports*, 3(4).  
716 <https://doi.org/10.1111/j.1758-2229.2010.00237.x>

717

718 Hall, R. O., & Madinger, H. L. (2018). Use of argon to measure gas exchange in turbulent  
719 mountain streams. *Biogeosciences*, 15(10). <https://doi.org/10.5194/bg-15-3085-2018>

720 Heilman, M. A., & Carlton, R. G. (2001). Methane oxidation associated with submersed  
721 vascular macrophytes and its impact on plant diffusive methane flux. *Biogeochemistry*, 52(2),  
722 207-224. DOI: 10.1023/A:1006427712846

723 Hijmans RJ, Phillips S, Leathwick J, Elith J (2023). dismo: Species Distribution Modeling. R  
724 package version 1.3-14. DOI: 10.32614/CRAN.package.dismo

725 Hoffmann, M., Schulz-Hanke, M., Garcia Alba, J., Jurisch, N., Hagemann, U., Sachs, T.,  
726 Sommer, M., & Augustin, J. (2017). A simple calculation algorithm to separate high-  
727 resolution CH<sub>4</sub> flux measurements into ebullition- and diffusion-derived components.  
728 *Atmospheric Measurement Techniques*, 10(1). <https://doi.org/10.5194/amt-10-109-2017>

729

730 Humlum, O. (1998). Active layer thermal regime 1991-1996 at Qeqertarsuaq, Disko Island,  
731 Central West Greenland. *Arctic and Alpine Research*, 30(3). <https://doi.org/10.2307/1551977>

732

733 Humlum, O., Hansen, B. U., & Nielsen, N. (1999). Meteorological observations 1998 at the  
734 arctic station, Qeqertarsuaq (69°15'N), Central West Greenland. *Geografisk Tidsskrift*, 99.  
735 <https://doi.org/10.1080/00167223.1999.10649428>

736 Iannone R, Roy O (2024). *DiagrammeR: Graph/Network Visualization*. R package version  
737 1.0.11. DOI: 10.32614/CRAN.package.DiagrammeR

738 Juncher Jørgensen, C., Schlaikjær Mariager, T., & Riis Christiansen, J. (2024). Spatial  
739 variation of net methane uptake in Arctic and subarctic drylands of Canada and Greenland.  
740 *Geoderma*, 443. <https://doi.org/10.1016/j.geoderma.2024.116815>

741 Klaus, M., Verheijen, H. A., Karlsson, J., & Seekell, D. A. (2022). Depth and basin shape  
742 constrain ecosystem metabolism in lakes dominated by benthic primary producers.  
743 *Limnology and Oceanography*, 67(12), 2763–2778. <https://doi.org/10.1002/lno.12236>

744 Keskitalo, K. H., Bröder, L., Shakil, S., Zolkos, S., Tank, S. E., van Dongen, B. E., Tesi, T.,  
745 Haghipour, N., Eglinton, T. I., Kokelj, S. v., & Vonk, J. E. (2021). Downstream Evolution of  
746 Particulate Organic Matter Composition From Permafrost Thaw Slumps. *Frontiers in Earth  
747 Science*, 9. <https://doi.org/10.3389/feart.2021.642675>

748 Kleber, G. E., Magerl, L., Turchyn, A. V., Schloemer, S., Trimmer, M., Zhu, Y., & Hodson,  
749 A. (2025). Proglacial methane emissions driven by meltwater and groundwater flushing in a  
750 high-Arctic glacial catchment. *Biogeosciences*, 22(3), 659-674. <https://doi.org/10.5194/bg-22-659-2025>

751 Kluge, M., Wauthy, M., Clemmensen, K. E., Wurzbacher, C., Hawkes, J. A., Einarsson, K.,  
752 Rautio, M., Stenlid, J., & Peura, S. (2021). Declining fungal diversity in Arctic freshwaters  
753 along a permafrost thaw gradient. *Global Change Biology*, 27(22).  
754 <https://doi.org/10.1111/gcb.15852>

755 Korrensalo, A., Alekseychik, P., Mammarella, I., et al. (2024). High-resolution spatial  
756 patterns and drivers of terrestrial ecosystem greenhouse gas fluxes in sub-Arctic Finland.  
757 *Biogeosciences*, 21, 335–352. <https://doi.org/10.5194/bg-21-335-2024>

758 Kuhn, M. A., Varner, R. K., Bastviken, D., Crill, P., MacIntyre, S., Turetsky, M., Walter  
759 Anthony, K., McGuire, A. D., and Olefeldt, D. (2021). BAWLD-CH4: a comprehensive  
760 dataset of methane fluxes from boreal and arctic ecosystems, *Earth System Science Data*, 13,  
761 5151–5189. DOI: 10.5194/essd-13-5151-2021

762 Kutzbach, L., Schneider, J., Sachs, T., Giebels, M., Nykänen, H., Shurpali, N. J.,  
763 Martikainen, P. J., Alm, J., & Wilmking, M. (2007). CO<sub>2</sub> flux determination by closed-  
764 chamber methods can be seriously biased by inappropriate application of linear regression.  
765 *Biogeosciences*, 4(6). <https://doi.org/10.5194/bg-4-1005-2007>

766 Kyzivat, E. D., Smith, L. C., Garcia-Tigreros, F., Huang, C., Wang, C., Langhorst, T., et al.  
767 (2022). The importance of lake emergent aquatic vegetation for estimating Arctic–Boreal  
768 methane emissions. *Journal of Geophysical Research: Biogeosciences*, 127(6),  
769 e2021JG006635. <https://doi.org/10.1029/2021JG006635>

770

771

772

773

774

775

776

777

778 Laanbroek, H. J. (2010). Methane emission from natural wetlands: Interplay between  
779 emergent macrophytes and soil microbial processes. A mini-review. In *Annals of Botany*  
780 105(1). <https://doi.org/10.1093/aob/mcp201>

781

782 Larsen, J. G., & Larsen, L. M. (2022). Lithostratigraphy, geology and geochemistry of the  
783 Tertiary volcanic rocks on Svartenhuk Halvø and adjoining areas, West Greenland. *GEUS*  
784 *Bulletin*, 50. <https://doi.org/10.34194/geusb.v50.8295>

785

786 Liebner, S., Zeyer, J., Wagner, D., Schubert, C., Pfeiffer, E. M., & Knoblauch, C. (2011).  
787 Methane oxidation associated with submerged brown mosses reduces methane emissions  
788 from Siberian polygonal tundra. *Journal of Ecology*, 99(4). <https://doi.org/10.1111/j.1365-2745.2011.01823.x>

790 Li, M., Peng, C., Zhu, Q., Zhou, X., Yang, G., Song, X., & Zhang, K. (2020). The significant  
791 contribution of lake depth in regulating global lake diffusive methane emissions. *Water*  
792 *Research*, 172, 115465. <https://doi.org/10.1016/j.watres.2020.115465>

793 Michel, G., Tonon, T., Scornet, D., Cock, J. M., & Kloareg, B. (2010). Central and storage  
794 carbon metabolism of the brown alga *Ectocarpus siliculosus*: Insights into the origin and  
795 evolution of storage carbohydrates in Eukaryotes. *New Phytologist*, 188(1).  
796 <https://doi.org/10.1111/j.1469-8137.2010.03345.x>

797

798 Miner, K. R., Turetsky, M. R., Malina, E., Bartsch, A., Tamminen, J., McGuire, A. D., Fix,  
799 A., Sweeney, C., Elder, C. D., & Miller, C. E. (2022). Permafrost carbon emissions in a  
800 changing Arctic. In *Nature Reviews Earth and Environment* (Vol. 3, Issue 1).  
801 <https://doi.org/10.1038/s43017-021-00230-3>

802

803

804 Mulholland, P. J., Fellows, C. S., Tank, J. L., Grimm, N. B., Webster, J. R., Hamilton, S. K.,  
805 Martí, E., Ashkenas, L., Bowden, W. B., Dodds, W. K., McDowell, W. H., Paul, M. J., &  
806 Peterson, B. J. (2001). Inter-biome comparison of factors controlling stream metabolism.  
807 *Freshwater Biology*, 46(11). <https://doi.org/10.1046/j.1365-2427.2001.00773.x>

808

809 Myrstener, M., Sponseller, R. A., Bergström, A. K., & Giesler, R. (2021). Organic carbon  
810 availability regulates aquatic metabolism in a boreal stream network across land-use settings.  
811 *Freshwater Biology*, 66(10), 1899–1911. <https://doi.org/10.1111/fwb.13793>

812

813 Neumann, R. B., Moorberg, C. J., Lundquist, J. D., Turner, J. C., Waldrop, M. P., McFarland,  
814 J. W., Eus Kirchen, E. S., Edgar, C. W., & Turetsky, M. R. (2019). Warming Effects of Spring  
815 Rainfall Increase Methane Emissions From Thawing Permafrost. *Geophysical Research*  
816 *Letters*, 46(3). <https://doi.org/10.1029/2018GL081274>

817

818 Noss, C., Bodmer, P., Koca, K., & Lorke, A. (2018). Flow and Turbulence driven Water  
819 Surface Roughness and Gas Exchange Velocity in Streams. *E3S Web of Conferences*, 40.  
820 <https://doi.org/10.1051/e3sconf/20184005018>

821

822 Olefeldt, D., Hovemyr, M., Kuhn, M. A., Bastviken, D., Bohn, T. J., Connolly, J., Crill, P.,  
823 Eus Kirchen, E. S., Finkelstein, S. A., Genet, H., Grosse, G., Harris, L. I., Heffernan, L.,  
824 Helbig, M., Hugelius, G., Hutchins, R., Juutinen, S., Lara, M. J., Malhotra, A., ... Watts, J. D.

825 (2021). The boreal-arctic wetland and lake dataset (BAWLD). *Earth System Science Data*,  
826 13(11). <https://doi.org/10.5194/essd-13-5127-2021>

827

828 Olid, C., Zannella, A., & Lau, D. C. P. (2021). The Role of Methane Transport From the  
829 Active Layer in Sustaining Methane Emissions and Food Chains in Subarctic Ponds. *Journal*  
830 *of Geophysical Research: Biogeosciences*, 126(3). <https://doi.org/10.1029/2020JG005810>

831

832 Olid, C., Rodellas, V., Rocher-Ros, G., Garcia-Orellana, J., Diego-Feliu, M., Alorda-  
833 Kleinglass, A., Bastviken, D., & Karlsson, J. (2022). Groundwater discharge as a driver of  
834 methane emissions from Arctic lakes. *Nature Communications*, 13(1).  
835 <https://doi.org/10.1038/s41467-022-31219-1>

836

837 Pedersen, A. R., Petersen, S. O., & Schelde, K. (2010). A comprehensive approach to soil-  
838 atmosphere trace-gas flux estimation with static chambers. *European Journal of Soil Science*,  
839 61(6). <https://doi.org/10.1111/j.1365-2389.2010.01291.x>

840

841 QGIS Development Team: QGIS Geographic Information System, Open Source Geospatial  
842 Foundation Project, available at: <https://qgis.org>

843

844 Rasilo, T., Prairie, Y. T., & del Giorgio, P. A. (2015). Large-scale patterns in summer  
845 diffusive CH<sub>4</sub> fluxes across boreal lakes, and contribution to diffusive carbon emissions.  
846 *Global Change Biology*, 21(3), 1124–1139. <https://doi.org/10.1111/gcb.12741>

847

848 Rautio, M., Dufresne, F., Laurion, I., Bonilla, S., Vincent, W. F., & Christoffersen, K. S.  
849 (2011). Shallow freshwater ecosystems of the circumpolar Arctic. *Ecoscience*, 18(3).  
850 <https://doi.org/10.2980/18-3-3463>

851 Ravn, N. R., Elberling, B., & Michelsen, A. (2020). Arctic soil carbon turnover controlled by  
852 experimental snow addition, summer warming and shrub removal. *Soil Biology and*  
853 *Biochemistry*, 142. <https://doi.org/10.1016/j.soilbio.2019.107698>

854 Rocher-Ros, G., Sponseller, R. A., Bergström, A.-K., Myrstener, M., & Giesler, R. (2020).  
855 Stream metabolism controls diel patterns and evasion of CO<sub>2</sub> in Arctic streams. *Global*  
856 *Change Biology*, 26(3), 1400–1413. DOI: 10.1111/gcb.14895

857 Saunois, M., Martinez, A., Poulter, B., Zhang, Z., Raymond, P. A., Regnier, P., Canadell, J.  
858 G., Jackson, R. B., Patra, P. K., Bousquet, P., Ciais, P., Dlugokencky, E. J., Lan, X., Allen,  
859 G. H., Bastviken, D., Beerling, D. J., Belikov, D. A., Blake, D. R., Castaldi, S., Crippa, M.,  
860 Deemer, B. R., Dennison, F., Etiope, G., Gedney, N., Höglund-Isaksson, L., Holgerson, M.  
861 A., Hopcroft, P. O., Hugelius, G., Ito, A., Jain, A. K., Janardanan, R., Johnson, M. S.,  
862 Kleinen, T., Krummel, P. B., Lauerwald, R., Li, T., Liu, X., McDonald, K. C., Melton, J. R.,  
863 Mühle, J., Müller, J., Murguia-Flores, F., Niwa, Y., Noce, S., Pan, S., Parker, R. J., Peng, C.,  
864 Ramonet, M., Riley, W. J., Rocher-Ros, G., Rosentreter, J. A., Sasakawa, M., Segers, A.,  
865 Smith, S. J., Stanley, E. H., Thanwerdas, J., Tian, H., Tsuruta, A., Tubiello, F. N., Weber, T.  
866 S., van der Werf, G. R., Worthy, D. E. J., Xi, Y., Yoshida, Y., Zhang, W., Zheng, B., Zhu, Q.,  
867 Zhu, Q., and Zhuang, Q.: Global Methane Budget 2000–2020, *Earth Syst. Sci. Data*, 17,  
868 1873–1958, <https://doi.org/10.5194/essd-17-1873-2025, 2025>.

869

870 Schuur, E. A. G., McGuire, A. D., Schädel, C., Grosse, G., Harden, J. W., Hayes, D. J.,  
871 Hugelius, G., Koven, C. D., Kuhry, P., Lawrence, D. M., Natali, S. M., Olefeldt, D.,  
872 Romanovsky, V. E., Schaefer, K., Turetsky, M. R., Treat, C. C., & Vonk, J. E. (2015).

873 Climate change and the permafrost carbon feedback. In *Nature* (Vol. 520, Issue 7546).  
874 <https://doi.org/10.1038/nature14338>

875

876 Singleton, C. M., McCalley, C. K., Woodcroft, B. J., Boyd, J. A., Evans, P. N., Hodgkins, S.  
877 B., Chanton, J. P., Frolking, S., Crill, P. M., Saleska, S. R., Rich, V. I., & Tyson, G. W.  
878 (2018). Methanotrophy across a natural permafrost thaw environment. *ISME Journal*, 12(10).  
879 <https://doi.org/10.1038/s41396-018-0065-5>

880

881 Street, L. E., Dean, J. F., Billett, M. F., Baxter, R., Dinsmore, K. J., Lessels, J. S., Subke, J.  
882 A., Tetzlaff, D., & Wookey, P. A. (2016). Redox dynamics in the active layer of an Arctic  
883 headwater catchment; examining the potential for transfer of dissolved methane from soils to  
884 stream water. *Journal of Geophysical Research: Biogeosciences*, 121(11).  
885 <https://doi.org/10.1002/2016JG003387>

886

887 Stevenson, M. A., McGowan, S., Pearson, E. J., Swann, G. E. A., Leng, M. J., Jones, V. J.,  
888 Bailey, J. J., Huang, X., & Whiteford, E. (2021). Anthropocene climate warming enhances  
889 autochthonous carbon cycling in an upland Arctic lake, Disko Island, West Greenland.  
890 *Biogeosciences*, 18(8). <https://doi.org/10.5194/bg-18-2465-2021>

891

892 Thompson, H. A., White, J. R., Pratt, L. M., & Sauer, P. E. (2016). Spatial variation in flux,  
893  $\delta^{13}\text{C}$  and  $\delta^2\text{H}$  of methane in a small Arctic lake with fringing wetland in western Greenland.  
894 *Biogeochemistry*, 131(1–2), 17–33. <https://doi.org/10.1007/s10533-016-0261-1>

895

896 Vachon, D., & Prairie, Y. T. (2013). The ecosystem size and shape dependence of gas  
897 transfer velocity versus wind speed relationships in lakes. *Canadian Journal of Fisheries and  
898 Aquatic Sciences*, 70(12), 1757–1764. <https://doi.org/10.1139/cjfas-2013-0241>

899

900 Virkkala, A. M., Niittynen, P., Kemppinen, J., Marushchak, M. E., Voigt, C., Hensgens, G.,  
901 Kerttula, J., Happonen, K., Tyystjärvi, V., Biasi, C., Hultman, J., Rinne, J., & Luoto, M.  
902 (2024). High-resolution spatial patterns and drivers of terrestrial ecosystem carbon dioxide,  
903 methane, and nitrous oxide fluxes in the tundra. *Biogeosciences*, 21(2).  
904 <https://doi.org/10.5194/bg-21-335-2024>

905

906 Walter Anthony, K. M., Anthony, P., Grosse, G., & Chanton, J. (2012). Geologic methane  
907 seeps along boundaries of Arctic permafrost thaw and melting glaciers. *Nature Geoscience*,  
908 5(6). <https://doi.org/10.1038/ngeo1480>

909

910 Walter Anthony, K., Schneider von Deimling, T., Nitze, I., Frolking, S., Emond, A., Daanen,  
911 R., Anthony, P., Lindgren, P., Jones, B., & Grosse, G. (2018). 21st-century modeled  
912 permafrost carbon emissions accelerated by abrupt thaw beneath lakes. *Nature  
913 Communications*, 9(1). <https://doi.org/10.1038/s41467-018-05738-9>

914

915 Walvoord, M. A., & Kurylyk, B. L. (2016). Hydrologic Impacts of Thawing Permafrost—A  
916 Review. *Vadose Zone Journal*, 15(6). <https://doi.org/10.2136/vzj2016.01.0010>

917

918 Wallenius, A. J., Dalcin Martins, P., Slomp, C. P., & Jetten, M. S. M. (2021). Anthropogenic  
919 and Environmental Constraints on the Microbial Methane Cycle in Coastal Sediments. In  
920 *Frontiers in Microbiology* (Vol. 12). <https://doi.org/10.3389/fmicb.2021.631621>

921

922 Westergaard-Nielsen, A., Balstr m, T., Treier, U. A., Normand, S., & Elberling, B. (2020).  
923 Estimating meltwater retention and associated nitrate redistribution during snowmelt in an  
924 Arctic tundra landscape. *Environmental Research Letters*, 15(3).  
925 <https://doi.org/10.1088/1748-9326/ab57b1>

926 Winslow, L. A., Zwart, J. A., Batt, R. D., Dugan, H. A., Woolway, R. I., Corman, J. R.,  
927 Hanson, P. C., & Read, J. S. (2016). LakeMetabolizer: an R package for estimating lake  
928 metabolism from free-water oxygen using diverse statistical models. *Inland Waters*, 6(4).  
929 <https://doi.org/10.1080/IW-6.4.883>

930 H. Wickham. *ggplot2: Elegant Graphics for Data Analysis*. Springer-Verlag New York, 2016.  
931 DOI: 10.1007/978-3-319-24277-4

932 Xu, W., Lambæk, A., Holm, S. S., Furbo-Halken, A., Elberling, B., & Ambus, P. L. (2021).  
933 Effects of experimental fire in combination with climate warming on greenhouse gas fluxes  
934 in Arctic tundra soils. *Science of the Total Environment*, 795.  
935 <https://doi.org/10.1016/j.scitotenv.2021.148847>

936

937 Yuan, K., Li, F., McNicol, G., Chen, M., Hoyt, A., Knox, S., Riley, W. J., Jackson, R., &  
938 Zhu, Q. (2024). Boreal–Arctic wetland methane emissions modulated by warming and  
939 vegetation activity. *Nature Climate Change*, 14(3). <https://doi.org/10.1038/s41558-024-01933-3>

940

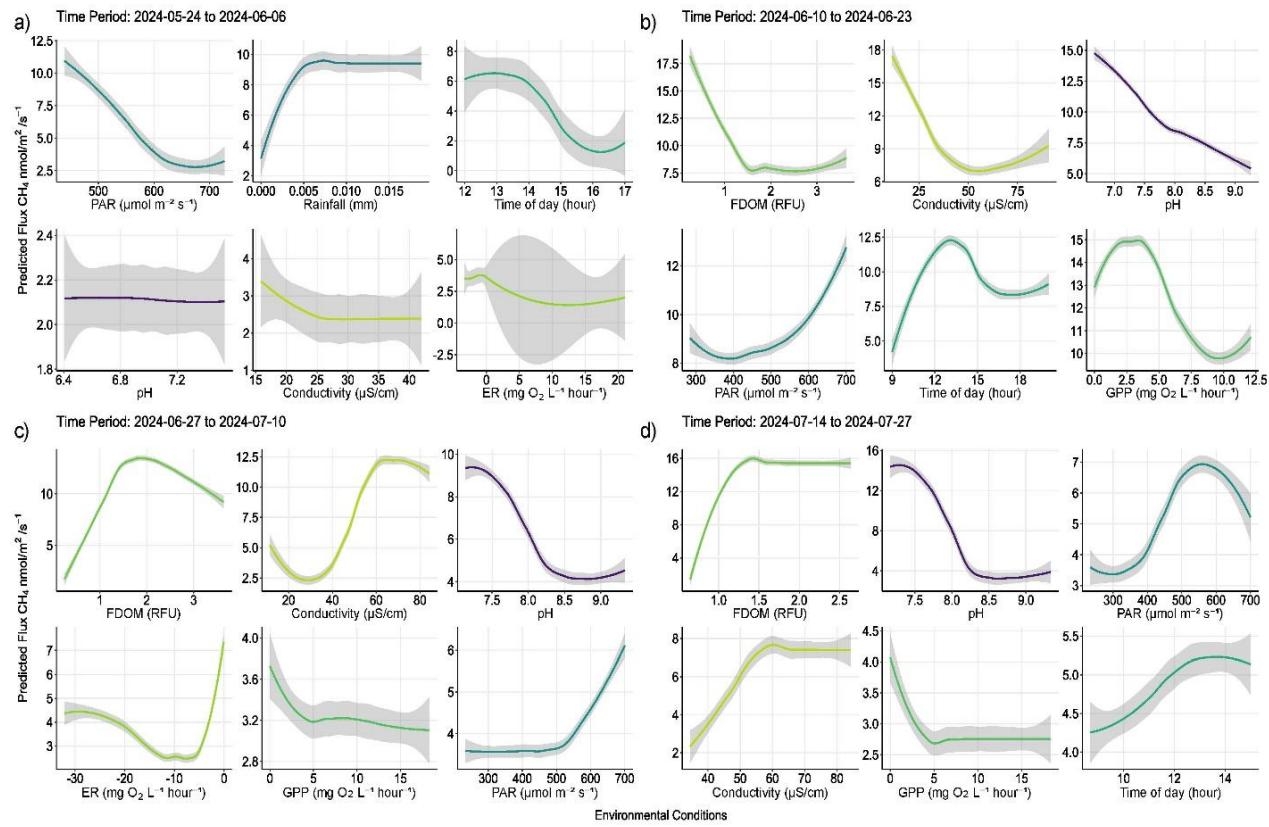
941 Žárský, J. D., Kohler, T. J., Yde, J. C., Falteisek, L., Lamarche-Gagnon, G., Hawkings, J. R.,  
942 Hatton, J. E., & Stibal, M. (2018). Prokaryotic assemblages in suspended and subglacial  
943 sediments within a glacierized catchment on Qeqertarsuaq (Disko Island), west Greenland.  
944 *FEMS Microbiology Ecology*, 94(7). <https://doi.org/10.1093/femsec/fiy100>

945

946 Zastruzny, S. F., Elberling, B., Nielsen, L., & Jensen, K. H. (2017). Water flow in the active  
947 layer along an arctic slope—An investigation based on a field campaign and model  
948 simulations. *The Cryosphere Discussions*, 1-32.

949

950 **Appendix A**



951

952 Figure A1. Partial dependency plots illustrating predicted marginal effects of meteorological, and  
 953 biochemical water conditions predicting diffusive CH<sub>4</sub> fluxes. Each figure represents the direct  
 954 marginal effect on CH<sub>4</sub> fluxes when all other predictors are held at their median, therefore giving a  
 955 more mechanistic understanding of those conditions and processes regulating fluxes from water  
 956 surfaces. The figures are ordered by importance in each time period and colors correspond to those  
 957 seen in Figure 6 of the main text. The colored lines represent the result of a fitted general additive  
 958 model ( $y \sim s(x)$ ) and thus a smooth representation across the 321 fitted BRT models. Each predictor  
 959 and its numerical range are given on the x-axis, while predicted fluxes are given on the y-axis.

960



970 Figure A2. Illustrates the various microbial forms encountered and found to influence CH<sub>4</sub> fluxes  
971 from the catchment area. In the photo on the far left we can see gas film on the water surface which  
972 was associated with submerged iron-oxidizing microbial assemblages, i.e. similar to what is shown in  
973 the middle photo, however exposed to the atmosphere in lower water levels. The photo on the right  
974 shows a brown alga which formed in the warm spring area southeast of the lake. In all cases,  
975 increased fluxes were generally encountered when measuring atop the middle and right microbial  
976 assemblages.

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