

# 1 **High-resolution spatial patterns and drivers of terrestrial** 2 **ecosystem carbon dioxide, methane, and nitrous oxide fluxes** 3 **in the tundra**

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24

25 **Abstract.** Arctic terrestrial greenhouse gas (GHG) fluxes of carbon dioxide (CO<sub>2</sub>), methane (CH<sub>4</sub>) and nitrous  
26 oxide (N<sub>2</sub>O) play an important role in the global GHG budget. However, these GHG fluxes are rarely studied  
27 simultaneously, and our understanding of the conditions controlling them across spatial gradients is limited.  
28 Here, we explore the magnitudes and drivers of GHG fluxes across fine-scale terrestrial gradients during the  
29 peak growing season (July) in sub-Arctic Finland. We measured chamber-derived GHG fluxes and soil  
30 temperature, soil moisture, soil organic carbon and nitrogen stocks, soil pH, soil carbon-to-nitrogen (C/N) ratio,  
31 soil dissolved organic carbon content, vascular plant biomass, and vegetation type from 101 plots scattered  
32 across a heterogeneous tundra landscape (5 km<sup>2</sup>). We used these field data together with high-resolution remote  
33 sensing data to develop machine learning models for predicting (i.e., upscaling) daytime GHG fluxes across the  
34 landscape at 2-m resolution. Our results show that this region was on average a daytime net GHG sink during  
35 the growing season. Although our results suggest that this sink was driven by CO<sub>2</sub> uptake, it also revealed small  
36 but widespread CH<sub>4</sub> uptake in upland vegetation types, almost surpassing the high wetland CH<sub>4</sub> emissions at the  
37 landscape scale. Average N<sub>2</sub>O fluxes were negligible. CO<sub>2</sub> fluxes were controlled primarily by annual average  
38 soil temperature and biomass (both increase net sink) and vegetation type, CH<sub>4</sub> fluxes by soil moisture  
39 (increases net emissions) and vegetation type, and N<sub>2</sub>O fluxes by soil C/N (lower C/N increases net source).  
40 These results demonstrate the potential of high spatial resolution modeling of GHG fluxes in the Arctic. They  
41 also reveal the dominant role of CO<sub>2</sub> fluxes across the tundra landscape, but suggest that CH<sub>4</sub> uptake in dry  
42 upland soils might play a significant role in the regional GHG budget.

## 43 **1 Introduction**

44 Over the past millennia, Arctic soils in the treeless tundra biome have played an important role in the global  
45 climate system by accumulating large amounts of carbon (C) and nitrogen (N), thus cooling the climate  
46 (Hugelius et al., 2014, 2020; Strauss et al., 2017). However, the ongoing climate warming is changing the C and  
47 N cycles, leading to potentially increased net greenhouse gas (GHG) emissions from Arctic ecosystems to the  
48 atmosphere (Belshe et al., 2013; McGuire et al., 2012; Masyagina and Menyailo, 2020). Yet, even the current  
49 GHG balance of Arctic ecosystems is insufficiently understood due to severe gaps in flux measurement  
50 networks and poorly performing coarse-resolution models (Virkkala et al., 2021; Treat et al., 2018c). Thus, the  
51 contribution of Arctic regions to the global climate feedback remains uncertain.

52 One of the main uncertainties in understanding the Arctic GHG balance is related to the inadequately quantified  
53 magnitudes of all three main GHG fluxes - carbon dioxide (CO<sub>2</sub>), methane (CH<sub>4</sub>) and nitrous oxide (N<sub>2</sub>O) -  
54 which show pronounced spatial variability across the diverse terrestrial environmental gradients in tundra  
55 (Virkkala et al., 2018; Pallandt et al., 2021; Voigt et al., 2020). In most tundra ecosystems, CO<sub>2</sub> fluxes are the  
56 largest flux driving the GHG balance due to the strong growing season photosynthetic activity and relatively  
57 high non-growing season respiratory CO<sub>2</sub> losses (Natali et al., 2019; Euskirchen et al., 2012; Heiskanen et al.,  
58 2021). However, growing evidence points to the importance of CH<sub>4</sub> and N<sub>2</sub>O fluxes, which are more potent  
59 GHGs than CO<sub>2</sub> (Voigt et al., 2017b). All three gasses have distinct spatiotemporal dynamics (Emmerton et al.,  
60 2014; Bruhwiler et al., 2021). However, only a few studies have simultaneously considered the contributions of  
61 all three main GHG fluxes to the tundra GHG balance (Voigt et al., 2017b; Kelsey et al., 2016; Brummell et al.,  
62 2012; Wagner et al., 2019).

63 The largest fine-scale differences in Arctic GHG fluxes occur in ecosystems with spatially varying soil moisture  
64 conditions (McGuire et al., 2012). Broadly speaking, the Arctic can be divided into wetlands and drier uplands  
65 (i.e., shrublands, grasslands, and barren lands; see e.g. (Treat et al., 2018a; Virkkala et al., 2021). Wetlands  
66 cover between 5 and 25 % of the Arctic (Olefeldt et al., 2021; Kåresdotter et al., 2021; Reynolds et al., 2019).  
67 They are hotspots for soil C and N stocks and have the potential for high CH<sub>4</sub> emissions (Euskirchen et al.,  
68 2014; Hugelius et al., 2020); therefore they have been intensively studied (Rinne et al., 2018; Peltola et al.,  
69 2019; Turetsky et al., 2014). However, uplands cover the largest part of the Arctic (75 to 95 %) and can have  
70 significant variability in environmental conditions and GHG fluxes. These uplands have been relatively well  
71 studied for CO<sub>2</sub> fluxes (Williams et al., 2006; Cahoon et al., 2012a). Upland CH<sub>4</sub> and N<sub>2</sub>O fluxes, on the other  
72 hand, remain less well understood in terms of their magnitudes and drivers (Virkkala et al., 2018; Voigt et al.,  
73 2020). There are still likely some GHG flux hotspots to be discovered and coldspots to be verified, particularly  
74 in the upland tundra ecosystems.

75 The Arctic tundra is characterised by fine-scale environmental heterogeneity even within upland and wetland  
76 tundra environments. Thus, local-scale study settings that cover the main spatial environmental gradients are  
77 highly important (Treat et al., 2018c; Davidson et al., 2017). Such fine-scale variabilities are often measured  
78 with chambers, but most chamber-based study designs are limited to relatively small environmental gradients  
79 focusing on only a handful of different land cover types and environmental variables, leaving large gaps in our  
80 understanding of GHG flux hotspots (Virkkala et al. 2018). In this study, using an extensive spatial study design

81 with chamber GHG flux measurements from 101 plots, we aim to understand the magnitudes and environmental  
82 drivers of Arctic terrestrial CO<sub>2</sub>, CH<sub>4</sub>, and N<sub>2</sub>O fluxes in a heterogeneous tundra landscape dominated by upland  
83 heaths. By combining in-situ measurements and remote sensing data, we investigate the fine-scale (2 m) spatial  
84 heterogeneity of GHG fluxes across the landscape, and estimate the contribution of the three gases to the total  
85 landscape-scale GHG flux.

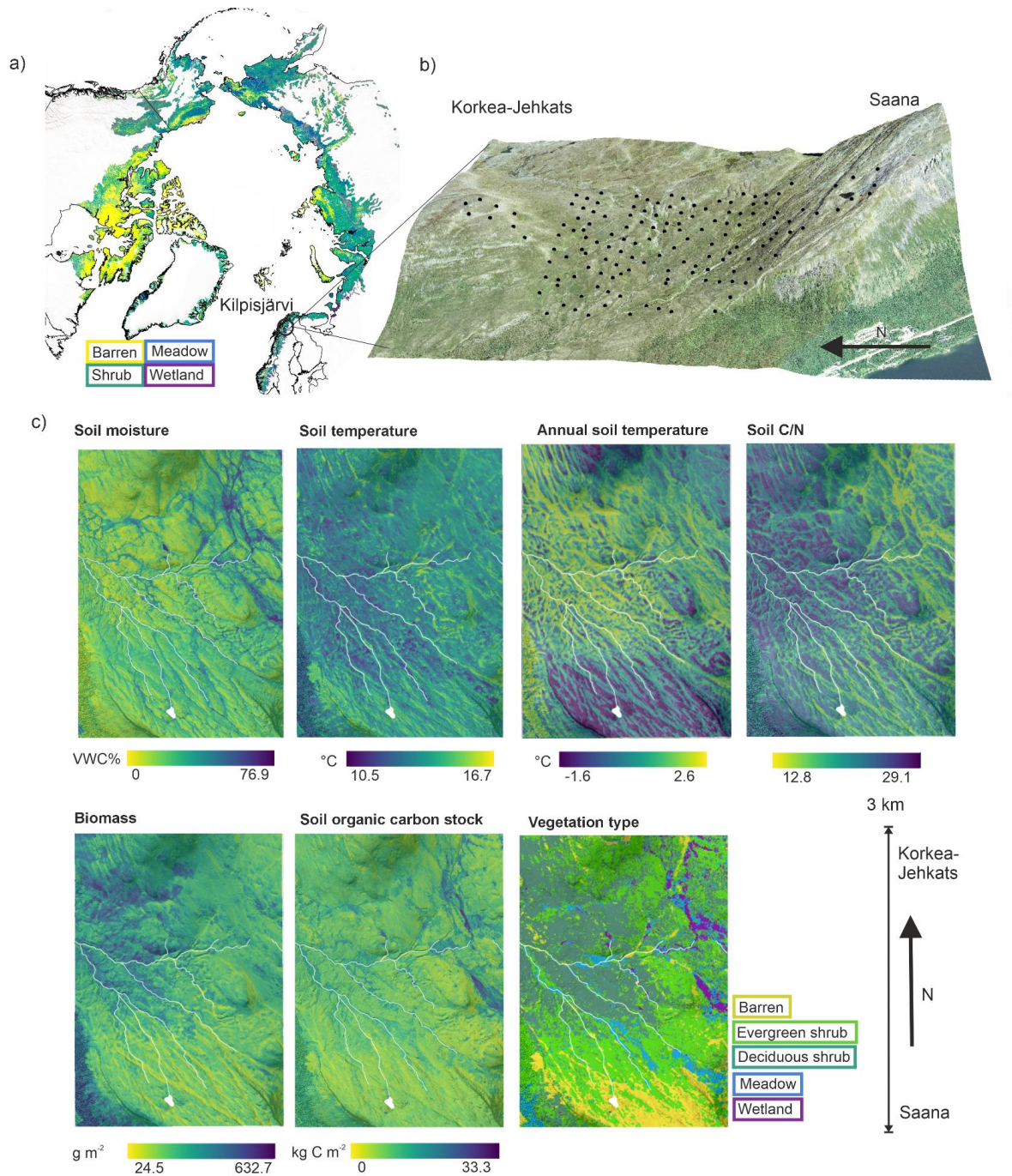
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## 87 **2 Materials and Methods**

### 88 **2.1 Study area**

89 The field measurements were collected during 2016-2018 in a sub-Arctic tundra environment in Kilpisjärvi  
90 (Gilbbesjávri in Northern Sámi language), northwestern Finland (69.06 N, 20.81 E). The study area is located on  
91 an elevational gradient between two fells, Saana (Sána; 1029 m.a.s.l) and Korkea-Jehkats (Jiehkkaš; 960 m.a.s.l),  
92 and the valley in between (~600 m.a.s.l.). The study area is above a mountain birch (*Betula pubescens* ssp.  
93 *czerepanovii*) forest and is dominated by dwarf-shrub evergreen and deciduous heaths. Dominant vascular plant  
94 species are, e.g., *Empetrum nigrum* ssp. *hermaphroditum*, *Betula nana*, *Vaccinium myrtillus*, *Vaccinium vitis-*  
95 *idaea*, and *Phyllodoce caerulea*. Vegetation in the wetlands is dominated by species common to fen wetlands,  
96 such as *Eriophorum* sp. or *Carex* sp. Mesic meadows are rich in forbs and grasses whereas barren heaths  
97 accommodate mostly lichens (e.g. *Cladonia* spp.) and mat-forming cushion plants (e.g. *Diapensia lapponica*) with  
98 scattered patches of *E. nigrum* and *B. nana*. Soils in the area are shallow (mean organic layer depth 6.6 cm, mean  
99 mineral layer depth 13.0 cm), and permafrost is absent from soils but can be found in the bedrock above 800 m  
100 a.s.l. (King and Seppälä, 1987). The environment is relatively undisturbed but experiences reindeer (*Rangifer*  
101 *tarandus tarandus*) grazing. The mean annual temperature in Saana fell (1002 m.a.s.l.) is -3.1 °C and the annual  
102 precipitation in Kilpisjärvi village ca. 5 km from the study site (480 m.a.s.l.) is 518 mm in 1991-2018 (Finnish  
103 Meteorological Institute, 2019a, b).

104 Our study design covered an area of ca. 3 x 1.5 km and consisted of 101 plots with GHG flux measurements and  
105 their supporting environmental data (Fig. 1). To produce continuous maps of soil temperature, moisture,  
106 vegetation type, biomass, soil C/N, and soil organic carbon stock, we utilized an extended dataset where some of  
107 the variables were measured from 50 plots while others from close to 6000 plots (Table S1). We selected the plots  
108 based on a combination of stratified sampling and systematic grid approaches, and the plots contain a variety of  
109 environmental gradients and habitats as well as the transition zones between them (Kemppinen et al., 2021). We  
110 recorded the locations of the plots using a hand-held Global Navigation Satellite  
111 System receiver with an accuracy of up to ≤6 cm under optimal conditions  
112 (GeoExplorer GeoXH 6000 Series; Trimble Inc., Sunnyvale, CA, USA).



113  
 114 **Figure 1:** The distribution of the main vegetation types across the Arctic tundra (Dinerstein et al., 2017; Agency,  
 115 2017) and the location of our study area (a), the distribution of plots (b) and environmental conditions derived  
 116 from statistical upscaling of in-situ measurements (see Sect. 2.4.2 Machine learning models) across the study area  
 117 (c). Soil moisture and temperature represent mean daytime (8 am to 8 pm) conditions from the 1st of July to the  
 118 2nd of August and annual soil temperature is an average within the entire year (July 2017-June 2018). Other  
 119 conditions represent growing season conditions and are considered static in this study. The aerial image is  
 120 produced by the National Land Survey of Finland (accessed in 2016).

## 121 2.2 Data

122 We measured GHG fluxes from 101 plots during the peak growing season (from now on, growing season).  
123 Environmental conditions explaining these GHG fluxes were measured at each plot. Most environmental variables  
124 had near complete spatial coverage; missing data were filled using the environmental predictions (see Sect. 2.4.2  
125 Machine learning models, Table S1). We used additional in-situ environmental data to upscale and visualize  
126 environmental conditions across the entire landscape (see Sect. 2.4.2 Machine learning models and Fig. 2):  
127 continuous soil moisture loggers (50 plots), continuous soil temperature loggers (250), soil samples for carbon  
128 and nitrogen stock and C/N estimation (168), and vegetation classification data (5280). The full set of variables  
129 at a plot consisted of the plot for GHG flux measurements, and of a nearby complementing plot (max. 2 m distance)  
130 where we monitored soil moisture and temperature continuously and did a vegetation classification. The additional  
131 plot was separated from the GHG plot to avoid disturbance of the continuous recordings. The additional plot was  
132 carefully situated to similar vegetation and microtopographic conditions as the GHG plot. Soil samples were  
133 collected as close as possible to the GHG plot.

### 134 2.2.1 Chamber measurements

135 We measured GHG exchange using a static, non-steady state non-flow-through system (Livingston and  
136 Hutchingson, 1995) composed of an acrylic chamber (20 cm diameter, 25 cm height). The chamber was placed  
137 on top of a collar and ventilated before each measurement. Prior to the measurements, we installed steel collars,  
138 which were 21 cm in diameter and 6 - 7 cm in height. Each collar was visited once during the growing-season,  
139 and measurements were conducted between 10 am and 5 pm. Although we did not have any temporal replicates,  
140 the spatial variation in our plots covered most of the temperature variation during the growing season. For more  
141 details, see Sect. S1.

142 For CO<sub>2</sub> flux measurements, transparent and opaque chamber measurements were conducted during 1<sup>st</sup> of July  
143 and 27<sup>th</sup> of July, 2018. The chamber included a small fan, a carbon dioxide probe GMP343 and an air humidity  
144 and temperature probe HMP75 (Vaisala, Finland). In the chamber, CO<sub>2</sub> concentration, air temperature and  
145 relative air humidity were recorded at 5-s intervals for 90 s. Photosynthetically active radiation was logged  
146 manually outside the chamber at 10-s intervals during the same period using a MQ-200 quantum sensor with a  
147 hand-held meter (Apogee Instruments, Inc, USA). MQ-200 measures photosynthetic photon flux density  
148 (PPFD) at a spectral range from 410 to 655 nm in  $\mu\text{mol m}^{-2} \text{s}^{-1}$ . For more details of the equipment, see  
149 Happonen et al. (2022).

150 We progressively decreased the light intensity of net ecosystem exchange (NEE) measurements from ambient  
151 conditions to ca. 80%, 50% and 30% PPFD by shading the chamber with layers of white mosquito net (replicate  
152 measurements per collar = 5 - 9). Ecosystem respiration (ER) was measured in dark conditions (0 PPFD), which  
153 were obtained by covering the chamber with a space blanket (replicates = 2 - 3). Before flux calculations, we  
154 discarded the first 0 - 5 s as well as the last 5 s of the measurements to remove potentially disturbed  
155 observations. Fluxes were calculated from the concentration change within the chamber headspace over time  
156 using linear regression (for performance statistics see Sect. S2).

157 We standardized NEE, GPP, and ER measurements conducted at different light and temperature conditions to  
158 allow across-plot comparison of the fluxes. We fitted light-response curves using a non-linear hierarchical  
159 bayesian model with the plot as a random effect (Sect. S5). We used the Michaelis-Menten equation to model  
160 instantaneous NEE with plot-specific ER, maximum photosynthetic rate ( $GPP_{max}$ ) and the half-saturation  
161 constant (K) as parameters using the same formula as in (Williams et al., 2006; Cahoon et al., 2012b). ER also  
162 had an exponential air temperature (T) response (for more details, see (Happonen et al., 2022)). We used this  
163 model to predict NEE at dark (0 PPFD, i.e. ER) and average light (600 PPFD) conditions, and an air temperature  
164 of 20 °C at each plot. 20 °C was chosen as it represents a typical air temperature inside the chamber during flux  
165 measurements, and 600 PPFD because it is widely used in tundra literature (Dagg and Lafleur, 2011; Shaver et  
166 al., 2007). We then subtracted ER from the NEE normalized to average light conditions to arrive at an estimate  
167 of normalized gross primary productivity (GPP). Negative values in NEE indicate a net sink of CO<sub>2</sub> from the  
168 atmosphere to the ecosystems. GPP and ER are given as positive values.

169 We measured CH<sub>4</sub> and N<sub>2</sub>O fluxes with an opaque chamber (0 PPFD). Measurements were conducted during  
170 the 2<sup>nd</sup> of July and 2<sup>nd</sup> of August, 2018. Five gas samples were taken within a 50-min enclosure time and  
171 transferred into 12-mL vials (Labco Exetainer, Labco Ltd.). The vials were pre-evacuated in the laboratory and  
172 filled with 25 mL of the sample in the field. Gas samples were analyzed at the University of Eastern Finland  
173 with a gas chromatograph (Agilent 7890B; Agilent Technologies, Santa Clara, CA, USA), equipped with an  
174 autosampler (Gilson Inc., Middleton, WI, USA), with thermal conductivity detector (TCD) for CO<sub>2</sub>, flame  
175 ionization detector (FID) for CH<sub>4</sub> and an electron capture detector (ECD) for N<sub>2</sub>O. We calculated gas  
176 concentrations from GC peak areas relative to peak areas derived by analyzing gas standards (CO<sub>2</sub>: 7  
177 concentration levels ranging from 0-10000 ppm; CH<sub>4</sub>: 7 concentration levels ranging from 0-100 ppm; N<sub>2</sub>O: 5  
178 concentration levels ranging from 0-5000 ppb). Fluxes were calculated from the concentration change within the  
179 chamber headspace over time using linear regression. Quality control was based on visual inspection and  
180 RMSE. We also verified that the RMSE was less than 3 \* standard deviation of gas standards in a similar  
181 concentration range. Negative values in these fluxes represent net CH<sub>4</sub> and N<sub>2</sub>O sinks from the atmosphere to  
182 the ecosystems.

### 183 2.2.2 Soil temperature and moisture data

184 Soil moisture and soil temperature were measured simultaneously during the flux measurements. We measured  
185 soil moisture with a time-domain reflectometry sensor (FieldScout TDR 300; Spectrum Technologies Inc.,  
186 Plainfield, IL, USA; 0 to 7.5 cm depth). Soil temperature measurements conducted at the same time as CO<sub>2</sub> flux  
187 measurements were taken with a thermometer (TD 11 thermometer; VWR International bvba; Leuven,  
188 Germany; 6.0 to 7.5 cm depth). Soil temperature measurements (TM-80N measure and ATT-50 sensor)  
189 conducted at the same time as CH<sub>4</sub> and N<sub>2</sub>O flux measurements were taken with a thermometer in the uppermost  
190 10 cm. We refer to these variables as soil moisture and soil temperature throughout the text.

191 Temperature loggers (Thermochron iButton DS1921G and DS1922L, San Jose, CA, USA and TMS-4; TOMST  
192 s.r.o., Prague, Czech Republic) monitored temperatures at 7.5 cm and 6.5 cm (iButton and TMS-4, respectively)  
193 belowground at 0.25–4 h intervals (Sect. S3). We calculated a variable describing soil temperature conditions

194 during the previous 12 months by averaging the iButton measurements from the study design (n=138) from July,  
195 2017 to June 2018. We refer to this variable as annual soil temperature. In addition to temperature, the TMS-4  
196 loggers also monitored soil moisture (raw time-domain transmission data between 1 and 4095) to a depth of c.  
197 14 cm (Wild et al., 2019). The raw time-domain transmission data was transformed into volumetric water  
198 content (VWC%) (Tyystjärvi et al., 2022).

199 These continuous soil moisture and temperature data were used to upscale soil microclimatic conditions at 2-  
200 hour timesteps during daytime (8 am to 8 pm) and from the 1st of July to the 2nd of August (see section Models  
201 used to predict environmental conditions). This period was chosen because the GHG fluxes were measured  
202 during this period and we did not want to extrapolate outside our main measurement campaign. Moreover, this  
203 period represents the peak growing season of this region.

### 204 2.2.3 Vegetation data

205 We took images from CH<sub>4</sub> and N<sub>2</sub>O collars on the measurement day and used them to classify the dominant  
206 vegetation to five distinct classes, following the Circumpolar Arctic Vegetation Map physiognomic  
207 classification system (Walker et al. 2005) with minor modifications (Fig. 1). We used the following classes:  
208 barren (< 10 % vegetation cover), evergreen shrub, deciduous shrub, meadow (graminoids and forbs), and  
209 wetlands. The sample sizes were not even between vegetation types, rather they roughly represent the spatial  
210 coverage of each vegetation type (8 observations of barren, 38 of evergreen shrub, 14 of deciduous shrub, 26 of  
211 meadow, and 15 of wetland). We utilized a larger dataset of 5820 vegetation descriptions estimated in the field  
212 and from aerial imagery from the study design to create the vegetation type map (for more details, see S4.1). We  
213 collected biomass samples from above-ground vascular plants using the clip-harvest method during late peak  
214 season, between 17<sup>th</sup> of July and 10<sup>th</sup> of August. Samples were collected within the chamber collars, and were  
215 oven-dried at 70 °C for 48 h and weighed after drying. We refer to this variable as biomass (g dry-weight m<sup>-2</sup>).

### 216 2.2.4 Soil sampling and analyses

217 We measured the thickness of the organic and mineral soil layers using a metal probe reaching up to 80 cm  
218 depth. We collected soil samples (ca. 1 dl) from the organic and mineral layers using metal soil core cylinders (4  
219 - 6 cm Ø, 5 - 7 cm height) during August in 2016-2018. The organic samples were collected from the top soil,  
220 and mineral samples directly below the organic layer which was on average 6.6 cm deep. Large roots were  
221 excluded from the samples. The soil samples were freeze-dried and analysed in the Laboratory of Geosciences  
222 and Geography and Laboratory of Forest Sciences (University of Helsinki). Bulk density (kg m<sup>-3</sup>) was estimated  
223 by dividing the dry weight by the sample volume. Soil organic layer pH was analyzed following ISO standard  
224 10390. Total carbon and nitrogen content (C%, N%) analyses were done using Vario Elementar Micro cube and  
225 Vario Elementar Max -analyzer (Elementar Analysensysteme GmbH, Germany). Prior to CN% analysis,  
226 mineral samples were sieved through a 2 mm plastic sieve. Organic samples were homogenized by hammering  
227 the material into smaller pieces.

228 Soils in this landscape are acidic and likely have a minimal amount of carbonates; consequently, we assumed  
229 C% to equal organic C%. Soil organic carbon and nitrogen stocks were calculated for the entire soil horizon up

230 to 80 cm (in 95 % of plots soil depth was less than that). Some plots lacked CN% data (30 % of the plots), and  
231 therefore, we used soil organic matter content estimated with the loss-on-ignition method according to SFS 3008  
232 (1990). We utilized a similar stock calculation framework using the bulk density, layer depth, and C% and N%  
233 data as in Kemppinen et al. (2021) except we used average bulk density and mineral C% estimates in each  
234 vegetation type in case that information was missing in stock calculation.

235 Soil samples for dissolved organic carbon concentration analyses in dry soil were collected between the 5th and  
236 14th of July 2018. After the collection, samples were stored at 4 °C and then dried at 60 °C for at least 5 days.  
237 Extraction of dissolved organic carbon was done using pure water extractions with 0.5 to 3 grams of dried soil  
238 added to 40 ml of water following the WEOC protocol from (Hensgens et al., 2021). Extracts were immediately  
239 filtered (0.7µm) using glass fibre filters, diluted, acidified to remove inorganic carbon, and measured on a  
240 Shimadzu TOC V-CPN analyzer set on the Nonpurgeable Organic Carbon mode. We refer to this variable as  
241 dissolved organic carbon.

#### 242 **2.2.5 Remotely sensed data**

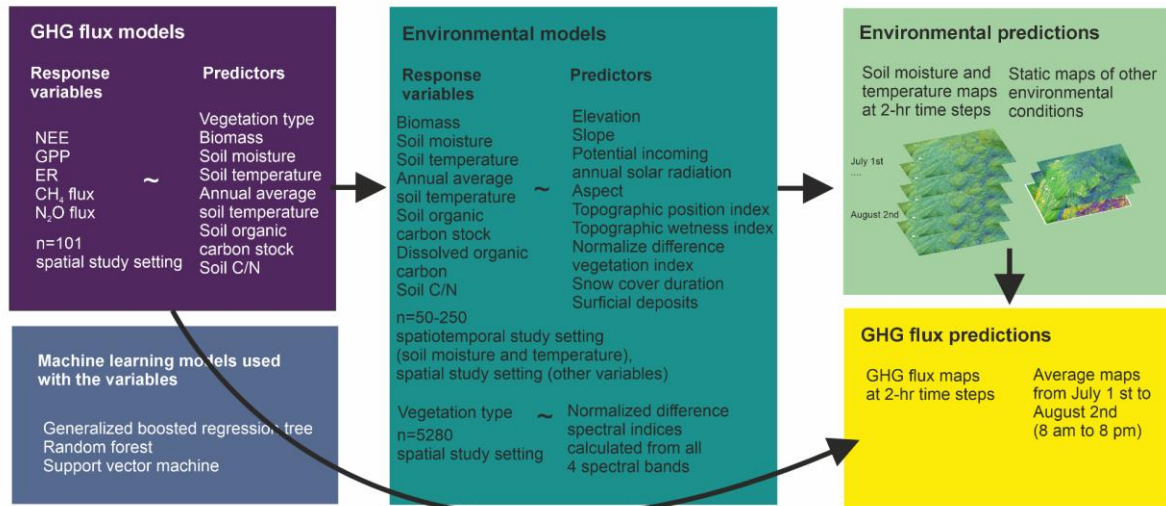
243 Remotely sensed optical and light detection and ranging-based (LiDAR) data describing topographic,  
244 vegetation, snow, and surficial deposit conditions was used for upscaling the in-situ measured environmental  
245 variables (Fig. 2, Sect. S4 and Fig. S1).

#### 246 **2.3 Statistical analyses**

247 We investigated the dependencies of GPP, ER, NEE, CH<sub>4</sub> flux, and N<sub>2</sub>O flux on environmental variables using  
248 statistical analyses which included analysis of variance (ANOVA), and machine learning modeling and  
249 prediction. We developed machine learning models, in which we 1) upscaled environmental data (annual soil  
250 temperature, soil temperature, soil moisture, soil C/N, soil organic carbon stock, dissolved organic carbon,  
251 biomass) using remotely sensed variables as predictors; 2) modeled GHG fluxes using the environmental data as  
252 predictors, and 3) upscaled GHG fluxes using the upscaled environmental data maps at a 2-meter spatial  
253 resolution across the landscape (Fig. 2). This two-step upscaling approach enabled us to focus on the  
254 relationships between GHG fluxes with their physical and ecological, in-situ measured environmental controls  
255 instead of the remotely sensed data that are proxies by nature. We ran all analysis in the R statistical  
256 programming environment (R Core Team 2020; version 4.0.3).

257





258

259 **Figure 2:** The upscaling framework used in this study. We first linked GHG fluxes to the in-situ environmental  
 260 drivers using machine learning models. Then we trained three machine learning models to upscale environmental  
 261 conditions across the landscape using remote sensing data. Then we used the GHG flux models and environmental  
 262 predictions to upscale GHG fluxes across the landscape throughout the entire growing season.

### 263 2.3.1 Analysis of variance (ANOVA)

264 We used one-way ANOVAs to test for vegetation type differences in environmental conditions, GHG fluxes,  
 265 and tested significance using multiple comparisons with a Tukey's honest significant difference test ( $p < 0.05$ ).  
 266 CH<sub>4</sub> flux, soil moisture, soil organic carbon and nitrogen stock, and biomass were not normally distributed, thus  
 267 we used Kruskal-Wallis test instead of ANOVA at first.

### 268 2.3.2 Machine learning models

269 We modeled our response variables using three machine-learning methods (generalized boosted regression  
 270 models, GBM; random forest, RF, and support vector machine regression, SVM), all of which have been widely  
 271 used in flux upscaling studies (see e.g. (Natali et al., 2019; Peltola et al., 2019; Tramontana et al., 2016). These  
 272 three approaches are non-parametric and can handle linear and non-linear relationships and different data  
 273 distributions. We chose RFs and GBMs because they utilize several decision trees in an ensemble model  
 274 framework and thus avoid overfitting, have high accuracy, are highly adaptable, and are not significantly  
 275 impacted by outliers. We chose SVMs because they are good at generalizing the relationships in the data. Based  
 276 on these models, we visualized the partial dependence plots characterizing the relationships between the  
 277 response and predictor variables while accounting for the average effect of the other predictors in the model  
 278 using the pdp package (Greenwell, 2017). Further, we calculated variable importance using the vip package  
 279 (Greenwell et al., 2020). Variable importance scores were estimated by randomly permuting the values of the  
 280 predictor in the training data and exploring how this influenced model performance based on the adjusted R<sup>2</sup>  
 281 values, with the idea that random permutation would decrease model performance (Breiman, 2001). We used  
 282 100 simulations to calculate 100 importance scores which were averaged. A standard deviation across these

283 scores was used as an uncertainty estimate, together with the differences in average importance across models.  
284 For more details, see Sect. S5.

285 We used ten topography, snow, vegetation, and surficial deposits variables to construct landscape-wide  
286 predictors matching the in-situ environmental conditions that we used to model the GHG flux values. These  
287 variables were the following: elevation, topographic wetness index, topographic position index at 5 and 30 m  
288 radii, aspect, slope, potential incoming solar radiation, normalized difference vegetation index, snow cover  
289 duration, and surface deposits. Soil organic carbon stocks, dissolved organic carbon, soil C/N, biomass, and  
290 annual soil temperature models were calibrated only once and a single prediction was made to the landscape.  
291 Soil temperatures and moisture vary throughout the growing season, thus, we calibrated each model at each time  
292 step and created 231 predictions over the study period (every 2 hours between 8 am and 8 pm from July 1st until  
293 August 2nd). For each variable, an ensemble prediction was produced by calculating a median prediction over  
294 the three predictions from the different modeling methods. Soil organic carbon stock was log+1 and biomass  
295 were log-transformed prior to tuning the models, and after making the predictions, values were transformed  
296 back to the original scale.

297 We examined the relationship between the five primary response variables (GPP, ER, NEE, CH<sub>4</sub> flux, N<sub>2</sub>O flux)  
298 and environmental predictors that describe (i) soil resources and conditions (soil moisture, soil C/N, soil pH)  
299 which are relevant to, for example, the growth of organisms (Nobrega and Grogan, 2008; Happonen et al.,  
300 2022); (ii) soil C and N stocks and dissolved organic carbon which are one of the main sources for the GHG  
301 emissions (Bradley-Cook and Virginia, 2018); (iii) soil temperatures which regulate enzymatic processes (St  
302 Pierre et al., 2019; Mauritz et al., 2017); and (iv) biomass and vegetation type which describe resource-use  
303 strategies, carbon inputs to soils and plant photosynthetic capacity, and integrate multiple environmental  
304 properties into one variable (Magnani et al., 2022). We excluded soil pH and soil nitrogen stock from modeling  
305 analyses due to high correlations (Pearsons's  $r > 0.7$ ) with soil moisture and soil organic carbon stock,  
306 respectively. Further, dissolved organic carbon was excluded due to its low importance in all the models. We  
307 did not use air temperature as a predictor as we already controlled for it in CO<sub>2</sub> fluxes in the light-response  
308 model, and we assumed that soil microbes regulating CH<sub>4</sub> and N<sub>2</sub>O cycling are most importantly driven by soil  
309 temperatures (Kuhn et al., 2021). The final predictors for our models were soil moisture, soil temperature,  
310 annual soil temperature, soil organic carbon stock, soil C/N, biomass, and vegetation type. The machine learning  
311 parameters tuned for each model can be found from Sect. S5.

312 We used the machine learning models to predict GHG fluxes across the landscape for each 2-hour time step  
313 from July 1st until August 2nd. Similar to the environmental predictions, an ensemble prediction was produced  
314 by calculating a median prediction over the three predictions from the different modeling methods. As our focus  
315 was on understanding the spatial patterns in the mean growing season fluxes, we averaged GHG flux predictions  
316 over the study period. However, a visualization of the predicted mean daily patterns in soil moisture and  
317 temperatures, and the consequent GHG fluxes is provided in the supplementary material (Fig. S2).

318 To compare the magnitude of all three important GHGs, namely CO<sub>2</sub>, CH<sub>4</sub>, and N<sub>2</sub>O, we calculated the radiative  
319 forcing strength of the three GHGs over a 100-year period from our measurements and ensemble predictions.  
320 We used the Global Warming Potential (GWP; 27 for CH<sub>4</sub> and 273 for N<sub>2</sub>O (IPCC 2021)) and sustained GWP

321 (45 for CH<sub>4</sub> and 270 for N<sub>2</sub>O (Neubauer 2015), which are, to our knowledge, the best and most widely used  
322 approaches that exist to compare flux magnitudes. We acknowledge that these approaches are designed to  
323 quantify an effect of a change in emission to the radiative forcing, and are thus not fully suitable to be used to  
324 quantify the climatic effect of natural continuous fluxes in our study (Mathijssen et al., 2022; Frolking et al.,  
325 2006).

326

327 For all of our models, we used a leave-one-plot-out cross validation scheme in which each plot was iteratively  
328 left out from the data set, and the remaining data were used to predict fluxes for the excluded plot to assess the  
329 predictive performance of the models (Bodesheim et al., 2018). Estimates of bias were calculated as an average  
330 of the absolute error (MAE) between prediction and actual observation. Coefficient of determination (R<sup>2</sup>) was  
331 used to determine the strength of the linear relationship between the observed and predicted fluxes.

332 The root mean squared error (RMSE) was used to estimate the model error. The same evaluation metrics were  
333 also calculated based on the prediction to the full model training data to represent model fit (Virkkala et al.  
334 2021); see table S3 which presents these for the individual models. Uncertainty in GHG flux predictions was  
335 derived by bootstrapping (fractional resampling with replacement based on vegetation type classes). We subset  
336 the model training data into 30 different data sets, all of which had the same number of observations as the  
337 original data itself. These 30 data sets were then used to produce 30 individual predictions for a subset of the  
338 times with all three machine learning models and their ensemble for each response variable (Sect. S5). The  
339 uncertainty estimates represent how different distributions of the input data as well as model parameters  
340 influence the upscaled flux maps.

341

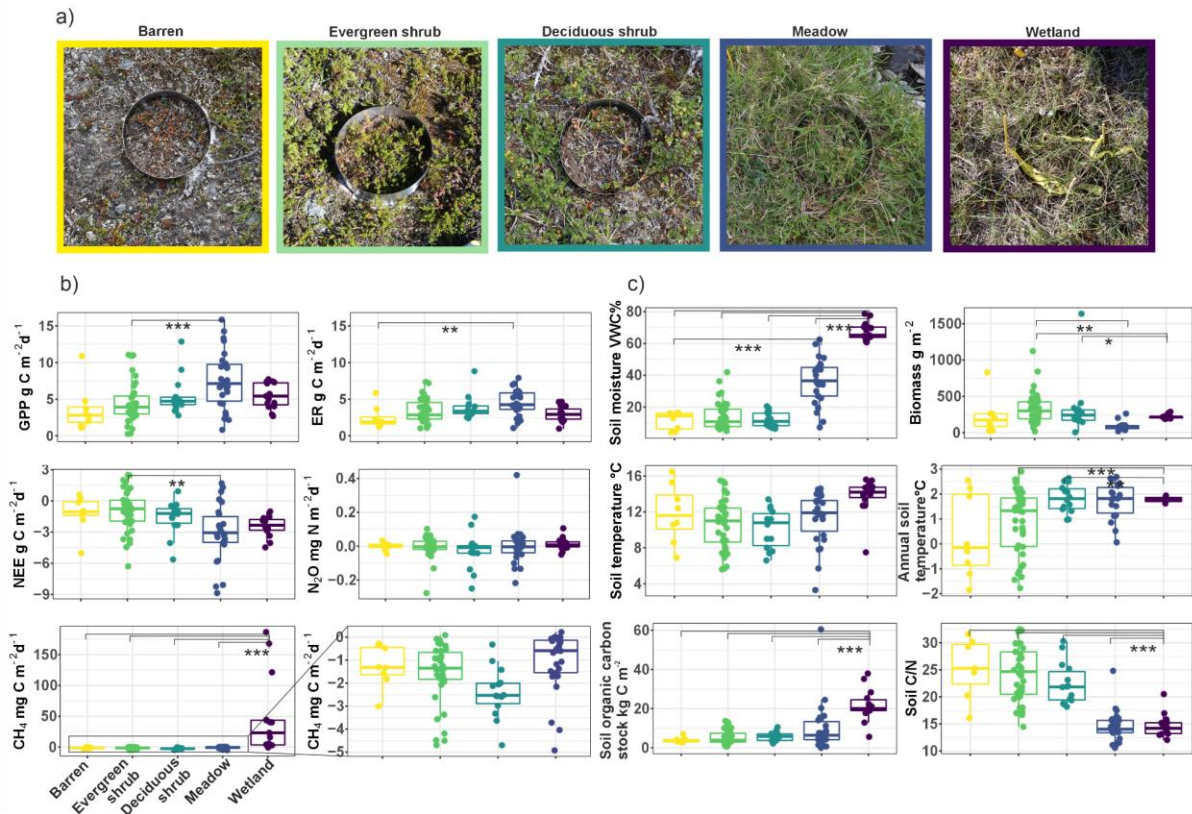
## 342 **3 Results**

### 343 **3.1 Environmental conditions and GHG fluxes across vegetation types**

344 We observed large variability in GHG fluxes and environmental conditions within and across vegetation types  
345 (Fig. 3, Table S2). The variability within the vegetation types differed depending on the flux and environmental  
346 variable considered (e.g., meadow class had large variability in GPP and evergreen shrub class in soil C/N).  
347 Frequently, wetlands differed clearly from the other vegetation types. While wetlands had high CH<sub>4</sub> emissions,  
348 all the other vegetation types with significantly lower soil moisture showed CH<sub>4</sub> uptake. Meadows were a  
349 significantly larger net CO<sub>2</sub> sink than evergreen shrub sites, while other vegetation types had intermediate NEE  
350 values. The N<sub>2</sub>O fluxes were low from all vegetation types, and varied between small sinks and small sources.

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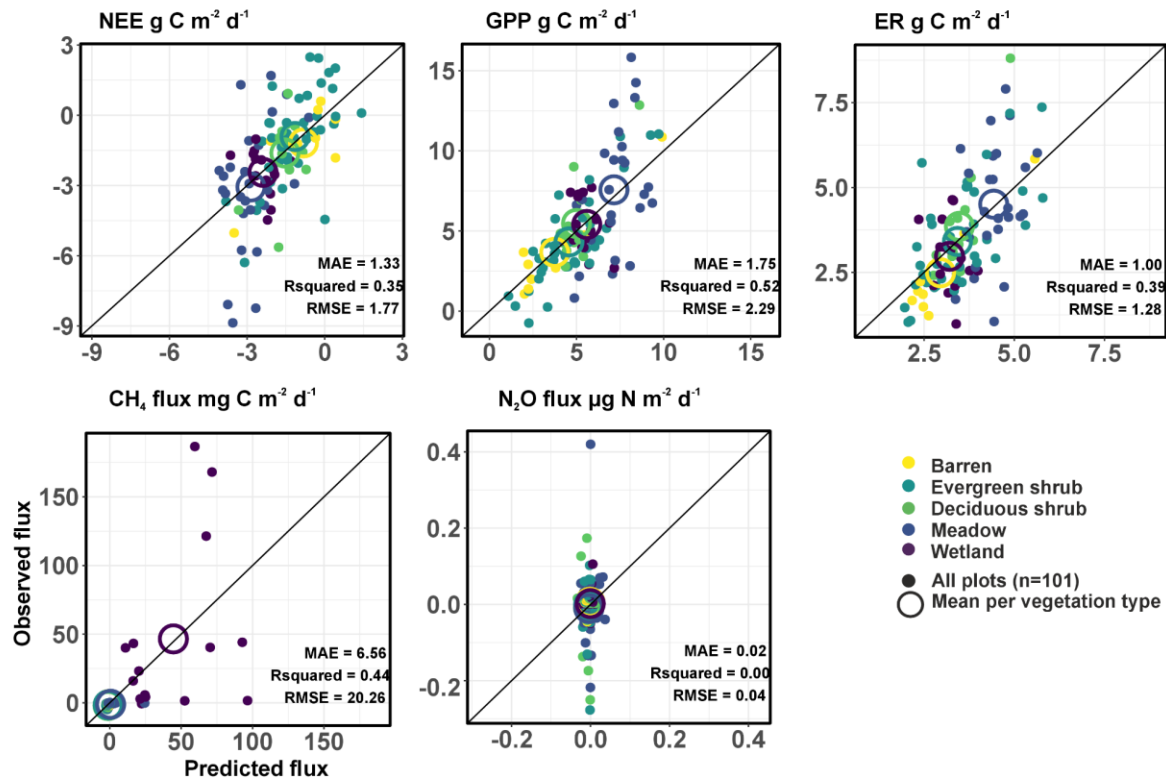
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**Figure 3:** The vegetation types considered in this study (a), the distribution of GHG fluxes (b), and environmental conditions (c) across the vegetation types. Lines represent Tukey's test results (\* =  $p \leq 0.05$ , \*\* =  $p \leq 0.01$ , \*\*\* =  $p \leq 0.001$ ). The box corresponds to the 25th and 75th percentiles, and the line within the box represents the median. The lines denote the 1.5 IQR of the lower and higher quartile, where IQR is the inter-quartile range, or distance between the first and third quartiles.

### 361 3.2 The performance of environmental and greenhouse gas flux models

362 The predictive performance of the ensemble environmental variable models was rather high but varied  
 363 depending on the variable ( $R^2$ : 0.26-0.71; Fig. S3). The predictive performance of the GHG models was for  
 364 most variables lower ( $R^2$ : 0.00-0.52), with  $\text{N}_2\text{O}$  flux models being close to random and GPP models performing  
 365 the best (Fig. 4). Model fit was significantly higher than predictive performance for all the fluxes (Table S3).  
 366 The scatterplots of observed and cross-validation-based predicted GHG fluxes suggest that the highest flux  
 367 estimates are often predicted most poorly, but the mean fluxes in each vegetation type were predicted  
 368 accurately, as expected.

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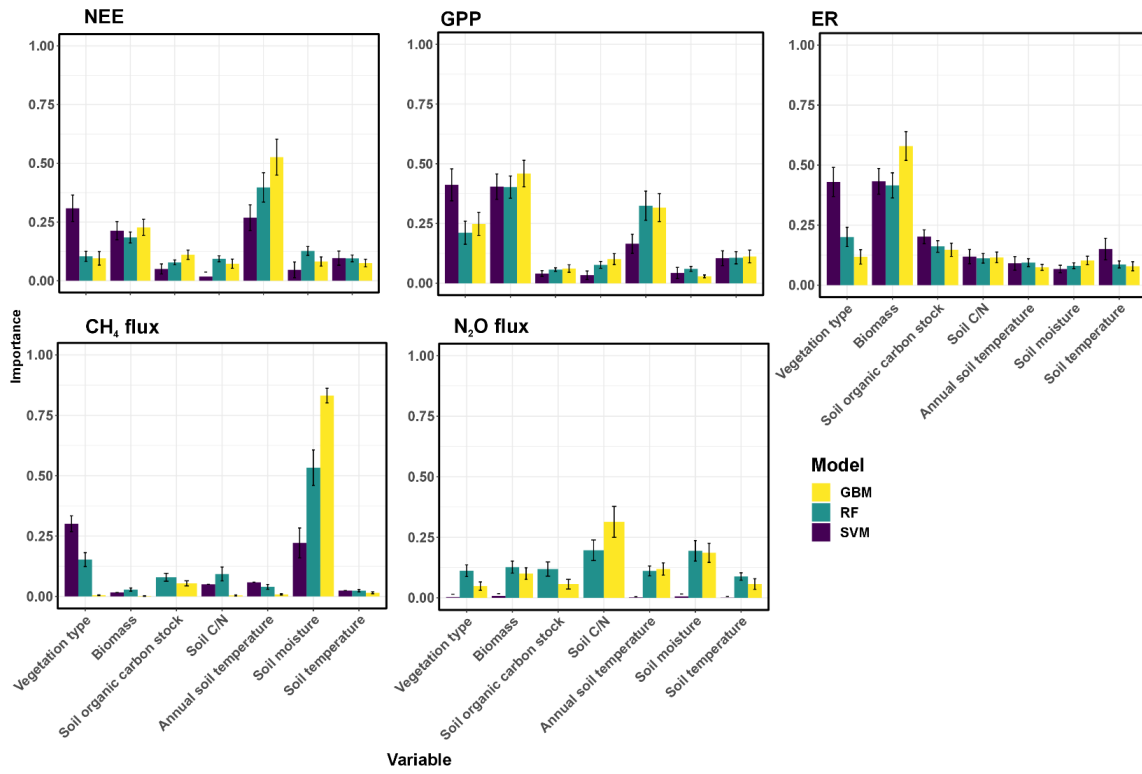
372 **Figure 4:** The correlation between observed and predicted values based on the ensemble model predictions (i.e.,  
 373 median of the three machine learning model outputs). Model predictive performance is described with mean  
 374 absolute error (MAE), R<sup>2</sup> (Rsquared), and RMSE (root mean square error).

### 375 3.3 Drivers of greenhouse gas fluxes

376 The most important controlling variables and the response shapes differed depending on the GHG flux (Fig. 5,  
 377 Fig. 6 and Fig. S4), and sometimes also depending on the machine learning model type applied. CO<sub>2</sub> fluxes  
 378 were driven by annual average soil temperature, biomass, and vegetation type. In addition, soil organic carbon  
 379 stocks were an important predictor for ER. Soil moisture and vegetation type were the most important predictors  
 380 for CH<sub>4</sub> fluxes, and soil C/N and soil moisture for N<sub>2</sub>O fluxes. In general, warmer and wetter conditions  
 381 increased net emissions of CH<sub>4</sub> and N<sub>2</sub>O and net uptake of CO<sub>2</sub>. Some fluxes were further positively correlated  
 382 with soil organic carbon stocks (ER, CH<sub>4</sub> flux) and negatively with soil C/N (GPP, ER, N<sub>2</sub>O). The importance  
 383 for variables explaining the N<sub>2</sub>O flux is low because the model predictive performance is close to random.

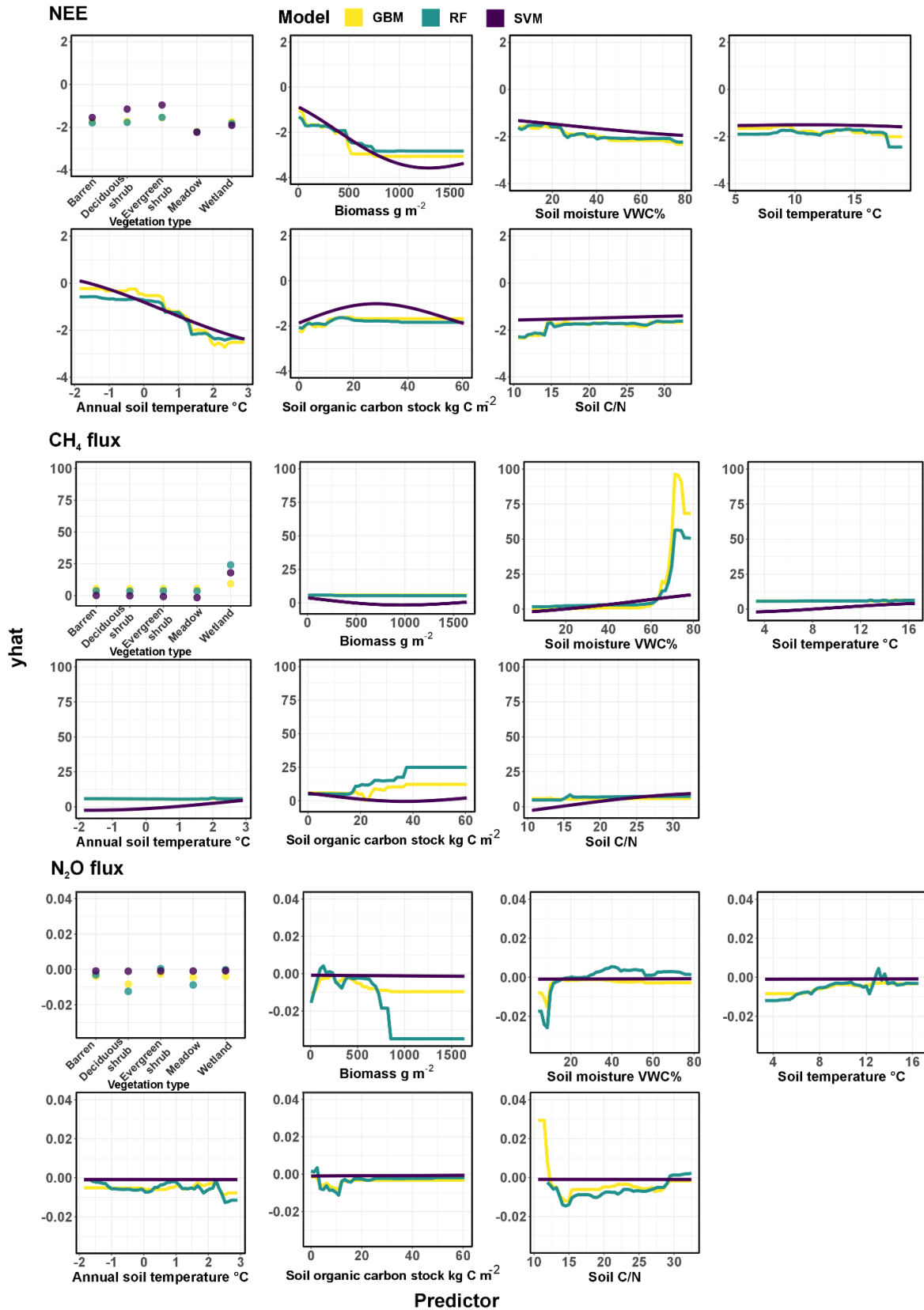
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**Figure 5:** The variable importance of the environmental variables used to predict GHG fluxes. The models were generalized boosted regression models (GBM), random forest (RF), and support vector machine regression (SVM).



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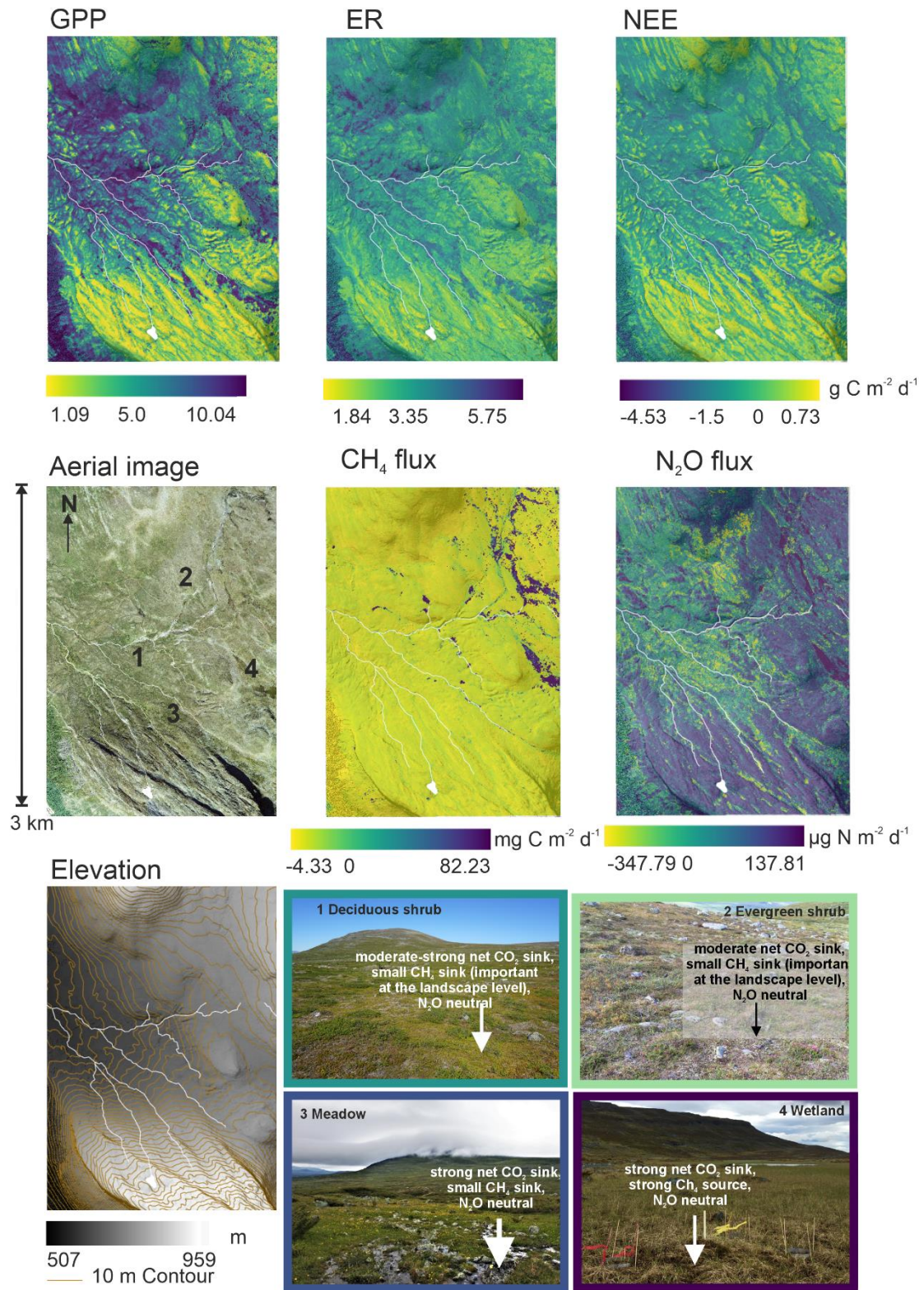
**Figure 6:** Partial dependence plots showing the relationships between GHG fluxes and environmental conditions across the three models (generalized boosted regression models, GBM; random forest, RF; and support vector machine regression, SVM). The y-axis of the plot (y-hat) represents the marginal effect of the

394 predictor on the response and should not be directly compared with observed or predicted values, rather the  
395 shape and direction of the response instead. RFs and GBMs are based on decision trees, where trees are split  
396 based on a certain threshold in the data, which can be seen as thresholds in the partial dependence plots as well.  
397 SVMs map the data into a high-dimensional space where a hyperplane is fit to separate them, creating smoother  
398 response shapes. Partial dependence plots for GPP and ER are found in Fig. S4.

### 399 **3.4 Spatial patterns and contributions in greenhouse gas flux predictions**

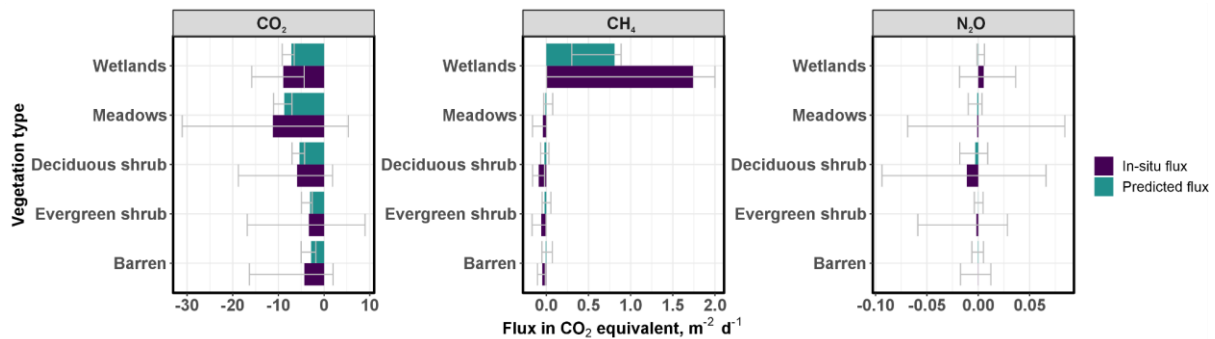
400 The model predictions show large spatial variability in GHG fluxes across the landscape (Fig. 7, Fig. S5). Net  
401 CO<sub>2</sub> uptake as well as GPP and ER were highest in the warm and productive meadow locations of the valley  
402 whereas CH<sub>4</sub> and N<sub>2</sub>O fluxes were highest in the eastern parts of the landscape that is dominated by wetlands.  
403 The prediction suggests small but widespread net CH<sub>4</sub> uptake across the entire upland region. CO<sub>2</sub> was the most  
404 important flux contributing to the net GHG sink (Fig. 8). Mean fluxes calculated based on the upscaled flux  
405 maps differ from the in-situ based ones, particularly for wetland CH<sub>4</sub> emissions (Fig. 8, Fig. S6).





407 **Figure 7:** Ensemble predictions of growing season GHG fluxes, averaged over the 1st of July to the 2nd of  
 408 August (only daytime variability between 8 am and 8 pm considered) and photographs summarizing the main  
 409 sink-source patterns in the landscape. Note that the southwestern corner of the study design has mountain birch  
 410 forest for which we did not have any data; we did not have measurements from the northeastern corner either.

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416 **Figure 8:** Growing season mean and percentile (0.025 and 0.975) GHG fluxes in CO<sub>2</sub> equivalents based on in-  
 417 situ data and upscaled flux predictions, averaged across the entire study period (only daytime variability  
 418 between 8 am and 8 pm considered) and across vegetation types. Note that the scale for the x axis is different for  
 419 each gas species, and that the uncertainties in in-situ versus predicted mean fluxes cannot be directly compared  
 420 with each other. The uncertainty in in-situ wetland CH<sub>4</sub> continues up to 6.7 but was cropped for visualization  
 421 purposes. The same graph using the sustained GWP approach can be found in the Supplement Fig. 6 and  
 422 demonstrates the potentially larger role of CH<sub>4</sub> fluxes over a 100-year horizon in this landscape.

## 423 4 Discussion

### 424 4.1 CO<sub>2</sub> fluxes driven by both biotic and abiotic variables

425 Our results show the importance of several environmental variables for CO<sub>2</sub> fluxes, demonstrating the strong  
 426 dependence of GPP and ER on a wide range of soil microclimatological, hydrological, soil biogeochemical, and  
 427 ecological processes (Sørensen et al., 2019; Dagg and Lafleur, 2011; Nobrega and Grogan, 2008; Cahoon et al.,  
 428 2016). Overall, the relationships with environmental conditions and GPP and ER were rather similar. Biomass  
 429 was a more important predictor than vegetation type for all the CO<sub>2</sub> fluxes, indicating that the quantity of plant  
 430 material producing and emitting carbon was potentially more important than the different types of plants  
 431 associated with CO<sub>2</sub> cycling in this study setting (Happonen et al., 2022). The high importance of plant-related  
 432 variables (e.g., leaf area index) as drivers of spatial variability in CO<sub>2</sub> fluxes has been previously found in other  
 433 tundra landscapes (Marushchak et al. 2013 and references therein).

434

435 Our models also show that annual soil temperatures have a different and stronger relationship with CO<sub>2</sub> fluxes  
 436 than instantaneous growing season soil temperatures, and these two soil temperature variables are indeed  
 437 negatively correlated in this study design. This is because annual soil temperatures are driven by winter soil

438 temperatures which increase with thicker snow cover that is found particularly in the valley and in  
439 microtopographic depressions, which are colder in the summer. Moreover, annual soil temperatures integrate  
440 many other environmental conditions from the entire year: they reflect growing season length and temperature  
441 conditions, regulate C and N availability, and control vegetation and microbial community composition and  
442 functioning over long time scales. These conditions have been shown to be important drivers of CO<sub>2</sub> fluxes  
443 across a range of Arctic sites (Zona et al., 2022; Lund et al., 2010). Similar to these previous studies, we  
444 observed that plots with warmer annual soil conditions have larger growing season GPP and ER fluxes and  
445 stronger net uptake. Our results also show other logical relationships between environmental conditions and CO<sub>2</sub>  
446 fluxes. For example, GPP and ER increased with soil moisture (Nobrega and Grogan, 2008). However, at  
447 around 50-60 % VWC this relationship plateaued and turned negative. This was likely due to the lack of oxygen  
448 for plant roots restricting growth of non-aerenchymous plants and for microbes, allowing only anoxic metabolic  
449 pathways, such as CH<sub>4</sub> productions in methanogenesis, where CO<sub>2</sub> production is low (Bridgham et al., 2013).  
450 Further, soil organic carbon stock was an important predictor for ER, but not so much for GPP. This was likely  
451 related to the higher soil carbon contents boosting decomposition (Schlesinger and Andrews, 2000).

#### 452 **4.2 Small but consistent net CH<sub>4</sub> uptake mostly driven by soil moisture**

453 Net CH<sub>4</sub> flux was strongly controlled by soil moisture due to its effect on regulating the anoxic and oxic soil  
454 conditions, and therefore CH<sub>4</sub> production (methanogenesis) and CH<sub>4</sub> consumption (CH<sub>4</sub> oxidation, or  
455 methanotrophy) (Kelsey et al., 2016; Christensen et al., 1996; Treat et al., 2018b). Our results demonstrate that  
456 the rate of CH<sub>4</sub> emissions increases sharply in water-logged soil conditions, i.e. at soil moisture levels of > 60  
457 VWC% (Vainio et al. 2021). In drier conditions (VWC < 60%), soils contain more oxygen, which prevents CH<sub>4</sub>  
458 production and increases net CH<sub>4</sub> uptake. This result supports findings from recent studies that show that drier  
459 upland tundra areas can be habitats for methane oxidizing bacteria which can use CH<sub>4</sub> from the atmosphere as  
460 their main energy source, transforming these environments to net CH<sub>4</sub> sinks (Christiansen et al. 2015; Juncher  
461 Jørgensen et al. 2015; Lau et al. 2015; Emmerton et al. 2014; Wagner et al. 2019; St Pierre et al. 2019; Voigt et  
462 al. 2023). Given the large area of the Arctic, even minor fluxes such as those observed here for CH<sub>4</sub> uptake can  
463 be of global importance. This CH<sub>4</sub> uptake can strengthen the GHG sink of the Arctic and prevent CH<sub>4</sub> from  
464 entering the atmosphere.

465  
466 Our results show that net CH<sub>4</sub> uptake increases not only in drier conditions but also in soils with low C/N, soil  
467 dissolved organic carbon, and carbon stocks. This is likely due to microbes needing and getting C and energy  
468 from the atmosphere due to limited soil C supply (Lau et al., 2015; Juutinen et al., 2022), and the capability of  
469 methanotrophs to effectively compete against classical heterotrophs dependent on larger organic  
470 macromolecules in these environments. The models did not clearly identify a particular vegetation type  
471 controlling net CH<sub>4</sub> uptake, however some individual models demonstrated deciduous shrubs and meadows to  
472 be more closely related to net CH<sub>4</sub> uptake (Larmola et al., 2010). Overall, our results indicate that net CH<sub>4</sub>  
473 uptake potential is present in any kind of upland tundra vegetation type (Fig. S7) as long as the abiotic  
474 conditions for microbes responsible for atmospheric CH<sub>4</sub> consumption are favourable.

475 Methane fluxes had a rather uniform distribution across the mineral upland regions (i.e., small but consistent net  
476 uptake). High CH<sub>4</sub> emissions were located in wetland regions dominated by high soil organic carbon stocks and  
477 moisture levels. Our observations demonstrated similar, or even higher net CH<sub>4</sub> uptake than previous studies.  
478 For example, dry tundra was CH<sub>4</sub> neutral in a recent Arctic-Boreal CH<sub>4</sub> flux synthesis (mean=3.83, median= -  
479 0.01 mg CH<sub>4</sub> m<sup>-2</sup> d<sup>-1</sup>; primarily based on growing season daytime fluxes; (Kuhn et al., 2021) whereas our study  
480 showed higher uptake rates for the non-wetland plots (mean=-2.05, median=-1.81 mg CH<sub>4</sub> m<sup>-2</sup> d<sup>-1</sup>). However,  
481 studies focusing on individual sites have recorded similar CH<sub>4</sub> flux magnitudes as observed here (Emmertson et  
482 al., 2014; Lau et al., 2015), but to the best of our knowledge, such extensive spatial patterns in CH<sub>4</sub> flux uptake  
483 using fine spatial resolution models as presented here have not been published so far.

#### 484 **4.3 N<sub>2</sub>O fluxes remain neglectable and unpredictable**

485 We observed moderate, and to a large extent unpredictable variability in N<sub>2</sub>O fluxes in this landscape. The  
486 differences in average fluxes between the vegetation types were small. Based on our observations, most  
487 vegetation types were on average N<sub>2</sub>O sinks or neutral but deciduous and evergreen shrubs and meadows had  
488 some variability from moderate N<sub>2</sub>O sinks (up to -300 μg N m<sup>-2</sup> d<sup>-1</sup>) to moderate N<sub>2</sub>O sources (up to 400 μg N  
489 m<sup>-2</sup> d<sup>-1</sup>). Overall, our average N<sub>2</sub>O fluxes were close to zero and thus low in the light of the recent review (Voigt  
490 et al., 2020), which demonstrated that vegetated soils in permafrost regions are often small but evident sources  
491 of N<sub>2</sub>O during the growing season (~30 μg N m<sup>-2</sup> d<sup>-1</sup>), and that barren or sparsely vegetated soils serve as  
492 substantial sources of N<sub>2</sub>O (~455 μg N m<sup>-2</sup> d<sup>-1</sup>). The relatively small N<sub>2</sub>O fluxes observed here can be explained  
493 by the nitrogen-limited nature of the studied soils and the strong competition between plants and microbes for  
494 nutrients: with shallow soils and low stocks of soil organic nitrogen, nitrogen release in labile forms by  
495 mineralization remains low (Voigt et al., 2020). Most of the data in the synthesis came from ecosystems that are  
496 not as much nitrogen-limited as our site (e.g., peatlands, grasslands).

497  
498 We were unable to explain the patterns in N<sub>2</sub>O fluxes with the predictors used here. This was likely related to  
499 the relatively low variability in N<sub>2</sub>O fluxes in most of the plots in general, and the complexity of the soil  
500 microbial N cycle, where N<sub>2</sub>O is produced (nitrification, denitrification, DNRA) and consumed (denitrification)  
501 by multiple co-occurring processes, differently regulated by environmental variables (Butterbach-Bahl et al.,  
502 2013). Nevertheless, the most important driver of N<sub>2</sub>O flux was soil C/N, and the models suggested that lower  
503 C/N ratios were linked to higher net N<sub>2</sub>O emissions. This is expected as the excess soil N in soils with low C:N  
504 ratio allows more rapid N mineralization, nitrification and denitrification as compared to microbial  
505 immobilisation which accelerates N<sub>2</sub>O emissions (Klemetsson et al., 2005; Liimatainen et al., 2018). Further,  
506 N<sub>2</sub>O emissions were highest in the wetlands, similar to (Ma et al., 2007) who explained this by high ammonia or  
507 nitrate levels boosting N<sub>2</sub>O production. The uppermost soil layers were also likely not fully saturated by water at  
508 the time of the wetland measurements, which can induce higher N<sub>2</sub>O emissions in oxic but still moist conditions,  
509 which allow aerobic nitrification and anaerobic denitrification to co-occur (Voigt et al., 2020; Takakai et al.,  
510 2008). In contrast to C fluxes, vegetation type did not play an important role for N<sub>2</sub>O fluxes. This might be  
511 related to our study having no measurements in the previously observed, clear N<sub>2</sub>O flux hot spots located in  
512 barren permafrost peatlands, such as peat plateaus or palsas, with thick organic layers and high inorganic N  
513 content (Repo et al., 2009; Voigt et al., 2017a).

#### 514 **4.4 The sub-Arctic tundra landscape is a strong growing season GHG sink**

515 Our results demonstrate a high level of spatial heterogeneity in the growing season GHG fluxes across the  
516 landscape, with areas acting as both net CO<sub>2</sub>, CH<sub>4</sub>, and N<sub>2</sub>O sinks and sources in some parts of it. Areas acting  
517 as GHG sinks covered most of the landscape (CO<sub>2</sub>: 91 %, CH<sub>4</sub>: 87 %, N<sub>2</sub>O: 73 %; 62 % of the area was a sink  
518 for all the three GHGs). We observed clear differences in flux magnitudes driven by key environmental  
519 conditions. Moist, and carbon and nitrogen-rich meadows and deciduous shrub heaths were strong GHG sinks.  
520 Wet sedge-dominated fens were GHG sinks with CH<sub>4</sub> emissions being compensated by net CO<sub>2</sub> uptake. Barren  
521 lands and evergreen shrubs were more resource-limited and closer to GHG neutral. These results are interesting  
522 in the light of the shrubification patterns observed across the entire Arctic (Myers-Smith et al., 2011; Parker et  
523 al., 2015; Vowles and Björk, 2018), and indicate that deciduous or evergreen shrub expansion may increase or  
524 decrease the growing season GHG sink. If shrubs expand to meadows, the GHG sink may decrease, whereas if  
525 they invade barren areas, the GHG sink may increase. However, our results did not quantify this change over  
526 time, or cover the entire year to confirm the net annual effect.

527

528 Our results indicate that this heterogeneous Arctic landscape was a cumulative net GHG sink during the  
529 measurement period during daytime (8 am to 8 pm) in July 2018. The July budget for CO<sub>2</sub> was -4.7 g C m<sup>-2</sup>  
530 month<sup>-1</sup>, for CH<sub>4</sub> 0.73 mg C m<sup>-2</sup> month<sup>-1</sup> and for N<sub>2</sub>O -10.0 µg N m<sup>-2</sup> month<sup>-1</sup>. The CO<sub>2</sub> sink is relatively small,  
531 likely due to the high cover of patchy and sparsely vegetated areas that were often CO<sub>2</sub> sources. This small sink  
532 value is an overestimation of the sink activity considering the whole course of the day as we did not have  
533 measurements from the night time and did thus not upscale fluxes in night-time conditions when ecosystems are  
534 net CO<sub>2</sub> sources due to the lack of light required for photosynthesis. It also overestimates the importance of CO<sub>2</sub>  
535 as a radiative forcing agent, since ecosystem CO<sub>2</sub> production during autumn and winter contributes substantially  
536 to the annual C balance (Celis et al., 2017; Commane et al., 2017), thereby reducing the CO<sub>2</sub> sink strength on an  
537 annual basis. Further, CH<sub>4</sub> uptake might continue even in rather cold conditions as long as soils remain dry and  
538 unfrozen (Emmerton et al., 2014). Nevertheless, our results demonstrate that net CO<sub>2</sub> uptake plays the most  
539 important role for the net growing season GHG budget. CH<sub>4</sub> emissions from wetlands are almost balanced by  
540 the net CH<sub>4</sub> uptake of other ecosystems. The role of N<sub>2</sub>O fluxes for the net GHG budget across the entire  
541 landscape is negligible for the growing season.

#### 542 **4.5 Methodological considerations in GHG flux modeling**

543 Our study creates new understanding about high-resolution upscaling of GHG fluxes by incorporating more  
544 chamber measurements, predictors, models, and environmental gradients compared to earlier efforts (Fox et al.,  
545 2008; Dinsmore et al., 2017; Räsänen et al., 2021; Juutinen et al., 2022; Vainio et al., 2021). For example, we  
546 included chamber measurements from 101 plots whereas earlier local-scale upscaling studies have usually had  
547 circa 30 plots. Further, we included eight different environmental predictors while other studies have often used  
548 only one or two, focusing on predictors describing vegetation type or soil moisture. Finally, we studied a tundra  
549 landscape that consists of almost all the main vegetation types of the entire Arctic, whereas earlier studies have  
550 investigated a narrower range of vegetation conditions, with a focus on wet ecosystems. However, at the same  
551 time, our models showed some signs of overfitting as demonstrated by the high model fit statistics and the  
552 mismatch between model fit and predictive performance statistics (Supplementary Text S5.3). This is a common

553 issue in upscaling (Kemppinen et al. 2018; Shi et al. 2022), and could indicate that the models have potentially  
554 learned to fit some noise or specific patterns unique to the training set instead of broadly generalizable  
555 relationships. Nevertheless, the relationships we observed were logical and comparable to those observed in  
556 other studies - both based on spatial and time series study designs (e.g., positive soil moisture-CH<sub>4</sub> flux or soil  
557 temperature-ER relationships (Euskirchen et al. 2014; Davidson et al. 2016; Zona et al. 2023)). Moreover, our  
558 study is based on a dataset focusing on spatial variation in GHG fluxes and correlations between variables.  
559 Therefore, the dataset should not directly be used to infer causal relationships or estimates of flux change over  
560 time (Damgaard 2019), and we advise caution when extrapolating these results to areas outside our study  
561 domain or different time periods.

562

563 Our study showed that using means of in-situ GHG fluxes in each vegetation class to derive a landscape-level  
564 GHG budget might produce significantly different results compared to the upscaled budget. This was apparent  
565 particularly for CH<sub>4</sub> fluxes, where the in-situ based average wetland CH<sub>4</sub> emission was more than twotimes  
566 larger CH<sub>4</sub> compared to the upscaled one. This mismatch is likely explained by the heterogeneity of  
567 environmental conditions and CH<sub>4</sub> fluxes within the wetland class that the chamber measurements alone could  
568 not cover (Fig. S7). A multivariate machine learning modeling approach with variables describing not only  
569 vegetation type but also soil moisture and other conditions were likely able to characterize the resulting CH<sub>4</sub>  
570 flux variability in a more representative way. For example, our soil moisture maps showed high variation in soil  
571 moisture between ca. 50 and 70 VWC% within the wetland areas, and high CH<sub>4</sub> emissions were observed only  
572 in areas with 60 VWC%. Overall, this result suggests that simple land cover-based upscaling efforts might lead  
573 to biased budget estimates, especially when spatial variability within land cover types is high, emphasizing the  
574 need for multivariate models in flux upscaling.

575

576 The performance of our models varied from good (GPP, CH<sub>4</sub> flux), moderate (ER and NEE) to low (N<sub>2</sub>O). CH<sub>4</sub>  
577 fluxes - both sources and sinks - were most accurately modeled, providing important support for future studies  
578 predicting not only the large CH<sub>4</sub> emissions but also the previously unquantified CH<sub>4</sub> uptake in Arctic  
579 landscapes. The lower predictive performance of the models for other GHG fluxes might be explained by the  
580 dynamic nature of fluxes not being represented in our spatial study design with no temporal chamber replicates  
581 in the plots, our models lacking important predictors, or our model structure not being ideal. The performance of  
582 the models could potentially be improved by describing plant functional composition using plant traits  
583 (Happonen et al., 2022), and including more detailed information about soil nutrients (e.g., soil nitrate or  
584 ammonium concentrations as soil C/N captures only very roughly how much N is available) or microbial  
585 communities (e.g., communities or genes associated with nitrification or methanogenesis or methanotrophy;  
586 (Pessi et al., 2022)).

587

588 Rainfall events are another source of uncertainty in our upscaling because they might increase soil moisture  
589 levels and activate processes related to methanogenesis, photosynthesis and respiration as well as nitrogen  
590 cycling. While our soil moisture predictions should capture these variations in soil wetness, we only made  
591 measurements once per plot under clear conditions and do not have information about how GHG fluxes might

592 respond to rainfall events. We might thus underestimate some of the instantaneous and longer-term changes in  
593 GHG fluxes during and after rain (see Text S1 and Fig. S10 for details).

594

595 We chose to use in-situ environmental data as predictors of GHG fluxes in our upscaling framework instead of  
596 linking remotely sensed variables with GHG fluxes directly. This was done to increase understanding about the  
597 mechanistic and ecological relationships but required us to first produce spatially continuous maps of  
598 environmental conditions, which might have added an additional layer of uncertainty into our framework.  
599 However, the most important environmental variables (i.e., soil moisture, temperature, biomass) had a high  
600 predictive performance. Nevertheless, future studies could explore the performance and information derived by  
601 upscaling GHG fluxes using high-resolution satellite or drone-derived remotely sensed indices directly (Siewert  
602 and Olofsson, 2020; Vainio et al., 2021; Berner et al., 2018).

603

604 Overall, the performance of our machine learning models predicting spatial variability in GHG fluxes was  
605 weaker than in other studies focusing on temporal variability (e.g., (López-Blanco et al., 2017; Celis et al.,  
606 2017), even though we had a comprehensive set of environmental measurements. Our results thus highlight the  
607 need for more focus on the spatial patterns in GHG fluxes. While the temporal variability is widely  
608 acknowledged as a source of uncertainty in GHG budget estimates (Baldocchi et al., 2018), the spatial  
609 variability may be just as important but remains insufficiently studied (Treat et al., 2018c). Study designs  
610 focusing on spatial variation in GHG fluxes using a combination of intensive measurement campaigns, remotely  
611 sensed datasets, and modeling approaches are informative although they do not produce direct information on  
612 the trends and drivers of GHG flux change following climate change. They provide new knowledge about the  
613 heterogeneity in GHG fluxes and their environmental drivers which is highly important for understanding flux  
614 magnitudes from local to global scales. Further, they can be used as a space-for-time substitution to understand  
615 ecosystem functions in locations that are assumed to be at different stages of development. Moreover, this  
616 knowledge is valuable for designing representative field studies in the future.

617

## 618 **5 Conclusions**

619 This study showed that predicting fluxes in heterogeneous tundra landscapes at high spatial resolutions is  
620 possible for CH<sub>4</sub>, GPP, and to some extent also NEE and ER fluxes but remains a challenge for N<sub>2</sub>O fluxes. This  
621 is a promising result for future high spatial resolution modeling studies that aim to understand the fine-scale  
622 biogeochemistry of the rapidly changing Arctic environments. Our study further demonstrates high spatial  
623 variability of GHG fluxes which is driven by a multitude of vegetation, soil microclimatological, hydrological,  
624 and biogeochemical conditions. The upscaling shows the importance of net CO<sub>2</sub> uptake for the peak growing  
625 season net GHG budget, and suggests that annual soil temperature and vegetation parameters are the most  
626 important drivers. Most importantly, it reveals small but widespread CH<sub>4</sub> uptake across the entire upland tundra  
627 in our domain that almost surpasses the high wetland CH<sub>4</sub> emissions. This provides more evidence to the  
628 relatively unquantified but important CH<sub>4</sub> sink in the Arctic GHG budget.

629

630 **Code/Data availability**

631 The field data, analysis codes and most of the results are available in a repository (Virkkala et al. 2023).  
632 Upscaling results for each individual timestep were not included in the repository due to their large size, but  
633 they can be acquired from the author upon request.

634 **Author contribution**

635 AMV and ML conceptualized the research with input from PN, JK, MEM, and CV. AMV, PN, JK, MEM, JK,  
636 CV, GH, VT, JH and ML contributed to data collection. AMV analyzed the data and wrote the manuscript draft.  
637 All the coauthors reviewed and edited the manuscript.

638 **Competing interests**

639 The authors declare no competing interests.

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