

**Authors' responses are in red in this file, and the revisions in the manuscript are done using Track Changes.**

**#Reviewer 1 Tim Moore**

This manuscript examines the emission of methane from a small wetland during the summer over a three year period (though the upper parts of the wetland were sampled in only the third year) and provides environmental measurements to help explain the patterns that were observed. The methane emission rates are consistent with other peatland systems in this type of environment. The originality of the study is the focus on emissions in the central part of the wetland, a mesotrophic sedge fen, which show that methane emission rates are smallest at the upper, drier part of the wetland, increase in the middle section which is wetter and then decrease in the central part which is essentially a very wet riparian strip with slowly flowing water in a stream. Methane production, consumption and emission are complex processes in terms of a multitude of influencing factors and the authors attempted to identify properties which may be important in the creation of the emission patterns. These include dissolved oxygen, temperature, plant distribution and estimates of plant production (GPP), LAI and presence of arenchymous species. Through essentially correlative analysis, the authors demonstrate the importance of flowing water and dissolved oxygen is reducing methane emissions in the central part of the fen, the drier conditions producing small emissions at the margin, resulting in maximum emissions in the middle part of the upland-stream transect. The manuscript concludes with speculation on the effect of climate change in these types of wetlands, though no strong predictions were made, rather an identification of environmental changes' linkages to methane emission.

The authors have assembled a good data base on a complicated northern wetland (compared to bogs for example). It is unfortunate that the upper parts of the wetland were not sampled until the third year, which provides evidence of the peak methane emissions part way to the fluvial section, but it seems that the third year results could probably occur in the earlier two years. As with such a complex system, with counter-acting properties influencing methane emission rates, it is difficult to tell how generally applicable these results will be. I think the paper would be strengthened by a greater consideration of the setting of the wetland within the overall watershed about which there is little mention. Climate change effects will be moderated by the other parts of the watershed, which seems to be forested from photographs and thus it would be useful to know the size of the wetland (which is about 1 km<sup>2</sup>) and the size, topography and soils/vegetation of the overall watershed. It would also be useful to know how common such mesotrophic sedge wetlands are in these landscapes and whether they are 'unique' so that application of the broad principles from this study may be inapplicable. Fens are more difficult to understand and model, than 'boring bogs'.

**-We thank the reviewer for suggesting modifying the paper from a larger view of the topic. In general, the studied fen type is rather common in many regions and therefore our results have global significance. Please find some texts in the third paragraph of Introduction. Also, we now describe the site better from the catchment point of view in section 2.1.**

Specific comments:

Line 26: lower peat temperatures→ cooler peat temperatures

-Changed as suggested.

Lines 27-28: move this sentence to after the next one, makes it a bit more 'fluent'.

-Changed as suggested.

Line 50: litter degradation rates→faster litter degradation rates

-Changed as suggested.

Line 112: a scale added to 1c would be useful.

-A scale was added to Fig. 1c. However, some parts of the illustration might not be scaled into the actual size, such as the size of the chambers and the distance between each chamber. Therefore, we also added some text in the caption.

Line 114: 1f does not seem to have 50, 60 and 90 plots visible, thus taken in 2017 or 2018?

-The drone image was taken in year 2018, this information is now added to the figure caption.

Line 118: peat temperatures were greater→ peat temperatures were warmer

-Changed as suggested.

Line 119: there were also larger temperature variations between the different depths in the drier parts?

-Yes, we meant in the drier parts. We now added the information there.

Line 121: deepest→maximum. What is the thickness of the peat along the transect from the central area to the margin.

-We changed deepest to maximum. Mathijssen et al. (2014) has demonstrated the peatland development history of this site based on multiple basal cores collected from different locations of the site. According to their study, the peat thickness of our sampled area ranges from *c.* 1 to 2.5 m.

Line 123: dry is “relative” so “relatively” is redundant.

-We deleted “relatively”.

Line 124: seem 57% is pretty precise to quality for a “c”, which means “about”.

-We deleted “c”.

Line 128: comment on overall catchment characteristics, relevant to the present study.

-We added the information on the size of the site, the surrounding vegetation and some text related to the water flowing feature. Please see details in the text in section 2.1.

Line 180: dissolved oxygen concentration, the unites appear to be %: % of what? Perhaps % of saturation? Please clarify.

-Yes, it is percent of air saturation. The information has been added.

Lines 252: S. warnsdorfii → S.warnstorffii

-Corrected.

Line 329: I assume the data for the three plot sets further from the stream are from only 2019. In (a) I cannot see the 60 m set, though it does appear in Fig. 4d, at the same median as at 50 m. What are the red circles?

-Yes, the three plot sets (50, 60 and 90) further from the stream are only from 2019. According to our definition, distances from the stream between 40 to 50 m are set as 50 m set, 50 to 60 are 60 m set, and 80 to 90 are 90 m set. In Figure 5, the actual distances of the plots to the stream in m were used in the analysis. The actual distance of 60 m set is 51 m, so the data points almost in the middle of x-axis 40 to 60 are from 60 m set. To avoid confusion, we now used “Distance to stream sampling set” when mention plot sets 10, 20, ... 60 (Fig.2, Fig.4 and Figs. S4-6) and used “Distance to stream (m)” for the actual distance (Fig.5). The red circles are model predicted CH<sub>4</sub> fluxes for different distances to stream.

Line 403: lower peat temperature → cooler peat temperature

-Changed as suggested.

Line 498: lower panel → lowest panel

-Changed as suggested.

Line 639: italicize journal title

-Changed as suggested.

Lines 775-777: misplaced reference

-Corrected.

## #Reviewer 2

This manuscript investigates the spatial patterning and environmental controls (namely water table depth, dissolved oxygen, soil temperature, GPP and various vegetation parameters) of methane emissions from a small sedge valley fen over a three-year period. They found methane emissions were lowest in the driest part of the fen, increased as moved towards the stream in the middle where the water table was seemingly in the optimal position and then decreased again closest to the stream due to the high amount of dissolved oxygen in the peat. Given the complexity in understanding CH<sub>4</sub> emissions from northern peatlands, the authors have done a good job of identifying the properties that are likely to influence them.

The methane emissions recorded from this site are within the range of other similar peatland systems.

I thank the authors for putting together a neat study which is well written and reads well. It will be a useful addition to the literature. The results section could do with some clarity, as the wording used is a little confusing and I found myself having to go back and remind myself a few times. I shuffle of some figures to and from the supplementary information will provide the reader with better clarity. I think the discussion overall is well written and the authors clearly have a strong understanding on the controls on CH<sub>4</sub> emissions in this site. However, I do think there needs to be some more information included for this system in the context of the wider landscape/hydrogeomorphic setting? The link to ‘climate change’ is a little tangible – with much of the work being focused on the environmental characteristics measured. I think giving the reader a greater understanding of the context of these ecosystems in wider landscape will help improve the ‘impacts of climate change’.

-Thank you for suggesting modifying the paper from a larger view of the topic, this was a very good point. In general, the studied fen type is rather common in many regions and therefore our results have global significance. Please find some texts in Introduction. Also, we now describe the site better from the catchment point of view in section 2.1.

Specific comments:

Line 19: I find the phrase ‘climate changed caused variations’ rather awkward and would recommend tweaking it.

-We modified the expression.

Line 55: Given the novelty of this study is the focus on valley fens which according to the authors are understudied, it would be useful to note approximately how much land they cover in Finland? It’s not clear to me if these are a common peatland type in this region or whether this study (although scientifically sound and interesting) may not be applicable across larger areas?

-Valley fens are commonly found in many regions; unfortunately, we are not able to estimate the coverage of them either in Finland or globally. Please see detailed information in the Introduction.

Section 2.1 Give more context to the wider area here.

-We added the information on the size of the site, the surrounding vegetation and some text related to the water flowing feature. Please see details in the text in section 2.1.

Line 116: What do you mean by different habitats? Distance from stream or different vegetation communities – this is not clear here.

-Here we meant generally different locations of the site. We rephrased the text.

Line 123: What do you mean by relatively dry areas? Are they just a ‘bit drier’ than the completely saturated area or is the water table consistently well below the surface?

-With *relatively dry areas* we meant basically having a water table below the surface unlike the surrounding inundated areas. Considering together with the comment from referee one, we deleted “relatively”.

Figure 1: Would be useful to include a scale bar to 1c and change the colour of the floating chambers so the difference between the static collars is clearer to the reader. Or use the drone image as the base image for the experimental design.

-As suggested, a scale bar was added to Figure 1c, and we changed the symbol of floating chambers.

Line 199: I find it confusing that use the word ‘clusters’ for location of sampling plots but also for the different plant communities. I think you should change one for clarity. When it comes to the results, it becomes difficult to follow.

-As suggested, we now use “clusters” only for plant communities. For flux sampling plots, we used “sets”.

Figure 3: I would recommend moving this to the supplementary information. I would actually replace it with Figure A2 as I think it is interesting for the reader to see the spatial variation in the landscape of the different vegetation communities rather than an ordination plot.

-We carefully considered the reviewer’s suggestion, but we prefer to keep them as they are – Figures 3 demonstrates the vegetation of the flux sampled plots, and the vegetation clustering information that we used in the data analyses. It also underpins the information that Figure 2A indicates, i.e. the vegetation structure is dominantly controlled by the stream.

Line 252: Typo: should be *S. warnstorffii*

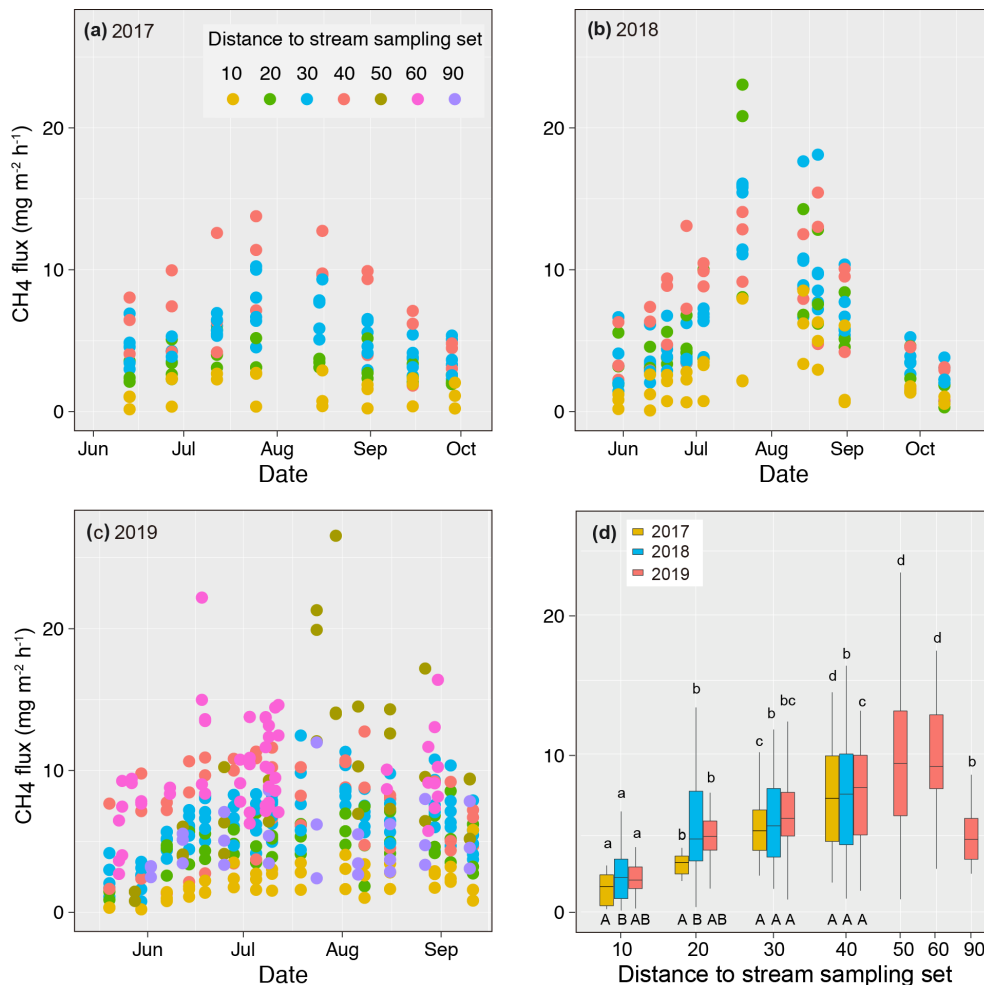
-Corrected.

Line 262: This is where the use of cluster becomes confusing. Could you change it to Plots?

-As suggested, we now use “clusters” only for plant communities, and “sets” for sampling plots.

Figure 4: Given you have 7 symbols; I think these figures would benefit from colour rather than being black. It’s hard to tell the difference between some of them. Again, the use of cluster here is confusing – I think plots would make much more sense. I would maybe use capitalised letters for the significant difference between studied year for each plot or cluster. I’m not sure I understand this as it seems you are looking at (for example) CH<sub>4</sub> in 2017 at 10, 20, 30, 40 m from stream and see there is no significant difference? How does that differ from the letters above the plot? Are the letters below just looking at differences WITHIN plot/cluster? And not across years?

-We tried new plots with sampling sets indicated using colors (shown below). However, we feel the quality is quite the same as that using symbols, especially for data in 2019 that have many overlapping data points. We therefore prefer to keep the use of symbols to make the figures readable in both online and printed versions. However, to improve the quality of the symbol figures, we now enlarged the symbols. We changed “cluster” to “set”.



We now used capitalized letters for the differences between studied years for each sampling set. The letters below the plots are used to indicate the difference between studied years for each sampling set, for example, if we look at set 10, the letters (A, B, AB) below the plots indicate the difference of CH<sub>4</sub> flux at set 10 between years 2017, 2018, and 2019.

Line 306: Do you mean there was variation within each of the three plots at each distance from the stream but also there is overall difference in CH<sub>4</sub> with distance from stream. This is a little confusing.

-Yes, this is what we meant. We now rephrased the text.

I have to admit, I find it rather confusing you create vegetation community classes but then don't use them once you start focusing on distance from stream?

-When we built mixed models, we tested if the plant community structure was important in driving CH<sub>4</sub> by adding the vegetation community clusters as potential predictors in the model, but they were not significant. Instead, the distance to stream is a significant predictor. In our case, it seems the stream has an override direct control on CH<sub>4</sub> which leaves the importance of plant species composition negligible. The relevant information was presented in section 3.3 and also in the first paragraph of section 4.2.

Figure 5: Can you use a different colour for the circles, the red and pink colour used are quite similar. Also do the pink circles in 5a represent 26m? Why 26 m? It's unclear to me here why you have 1m and 26 m?

-In Fig.5a, the color circles are predicted CH<sub>4</sub> flux for different distances to stream under a constant temperature. We used the color to separate them from the observed data circles. The idea in the original Figure 5b was to create a transect of distance to stream to demonstrate how the distance modified the responses to temperature. The use of 1, 26, and 60 was rather random, but we always used numbers within the range of the collected dataset, i.e., from 0 to 89 m.

This applied also to the original Figure 6 when setting different levels of the interactive variables. However, during the revision process, we realized that we hadn't considered the distribution of residuals of the final three-year data model in the original submission, and learned that a variance function is needed for our dataset as the residuals clearly show a cone shape indicating violation of the homogeneity assumption. In the revised version, we corrected this. While doing so, the interaction between distance to stream and temperature became nonsignificant ( $p = 0.113$ ), and therefore we ended excluding the interaction and correcting the related text, figures and tables. We sincerely apologise for this.

Figure 6: I find these figures quite hard to interpret given you have a combination of each interaction between variables. Again, the colours are very similar and should be changed to more contrasting colours.

-In Figure 6, the relationship between CH<sub>4</sub> and each significant predictor is presented in one

plot. If the predictor has interactions with another predictor, the interactions are presented by setting a transect of values for that interactive predictor and are presented using different colors. To simplify the figures a bit and also being inspired by the reviewer's comments on Figure 5, we made some modifications to Figure 6. Instead of setting four different levels, we now use only three levels for the interactive variables, i.e. min., average and max. values of the variables in the collected dataset. We also tried to clarify the figure text to help the readers.



# Water Flow Controls the Spatial Variability of Methane Emissions in a Northern Valley Fen Ecosystem

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## 15 **Abstract**

Northern peatlands are projected to be crucial in future atmospheric methane (CH<sub>4</sub>) budgets and have a positive feedback on global warming. Fens receive nutrients from catchments via inflowing water and are more sensitive than bogs to ~~climate change caused~~ variations in their ecohydrology. Yet, due to a lack of data detailing the impacts of moving water on microhabitats and CH<sub>4</sub> fluxes in fens, there remains large uncertainties in predicting CH<sub>4</sub> emissions from these sites under ~~climate changes~~. We measured CH<sub>4</sub> fluxes with manual chambers over three growing seasons (2017–2019) at a northern boreal fen. To address the spatial variation at the site where a stream flows through the long and narrow valley fen, we established sample plots at varying distances from the stream. To link the variations in CH<sub>4</sub> emissions to environmental controls, we quantified water levels, peat temperature, dissolved oxygen concentration, vegetation composition and leaf area index in combination with flux measurements during the growing season in 2019. We found that due to the flowing water, there was a higher water level, ~~lower-cooler~~ peat temperatures, and more oxygen in the peat close to the stream, which also had the highest total leaf area and gross primary production (GPP) values but the lowest CH<sub>4</sub> emissions. ~~Further from the stream, the conditions were drier and produced low CH<sub>4</sub> emissions. In contrast,~~ CH<sub>4</sub> emissions were highest at an intermediate distance from the stream where the oxygen concentration in the surface peat was low but GPP was still high. Further from the stream, the conditions were drier and produced low CH<sub>4</sub> emissions. Our results emphasise the key role of ecohydrology in CH<sub>4</sub> dynamics in fens, and for the first time show how a stream controls CH<sub>4</sub> emissions in a flow-through fen. As valley fens are common peatland ecosystems from the arctic to the temperate zones, future projections of global CH<sub>4</sub> budgets need to take flowing water features into account.

## 1 Introduction

Northern peatlands, which cover approximately 15 % of the boreal and arctic regions, are long-term sources of the greenhouse gas methane (CH<sub>4</sub>) (Korhola et al., 2010; MacDonald et al., 2006), partly counteracting the cooling impact of related long-term carbon dioxide (CO<sub>2</sub>) uptake. The response of northern peatlands to global warming has partly contributed to the recent increase in atmospheric CH<sub>4</sub> concentrations (Bousquet et al., 2011; Ciais et al., 2014; Kirschke et al., 2013), and modelling projections have suggested that, globally, wetland CH<sub>4</sub> emissions will continue to increase during the 21<sup>st</sup> century and have a positive feedback on global warming (Zhang et al., 2017). However, large uncertainties remain in the global CH<sub>4</sub> budget models due to limited knowledge of the relative contribution of the various environmental drivers that control CH<sub>4</sub> fluxes (Riley et al., 2011). To upscale observed CH<sub>4</sub> fluxes and produce realistic scenarios for future projections of atmospheric CH<sub>4</sub> concentrations, it is crucial to understand and quantify the correlations between peatland CH<sub>4</sub> emissions and their environmental drivers.

In peatlands, CH<sub>4</sub> is produced in wet and anoxic conditions below the water level by methanogens, and then released from the peat to the atmosphere. During the transport process, part of the produced CH<sub>4</sub> is consumed/oxidised by methanotrophs. The processes of CH<sub>4</sub> production, consumption, transport and final release to the atmosphere are affected by several environmental factors, such as water level, organic substrates, and temperature (Abdalla et al., 2016; Bellisario et al., 1999; Larmola et al., 2010). There is also evidence that peatland vascular plant functional types can affect CH<sub>4</sub> emissions by altering microbial community structure (Robroek et al., 2015). Sedge-dominated fens are CH<sub>4</sub> emission hotspots due to greater methanogenic activity (Juottonen et al., 2005) and [faster](#) litter degradation rates (Aerts et al., 1999). Also, the greater abundance of sedges (*Carex* spp.) in fens provides both a direct route for CH<sub>4</sub> movement to the atmosphere through aerenchyma tissue, thereby avoiding the oxidation of CH<sub>4</sub>, and also provides high-quality litter into the soil, which promotes CH<sub>4</sub> production (Noyce et al., 2014).

Fens, unlike bogs, are fed by mineral-rich water as seepage from the mineral soil below (soligenous fens) or from surface water flow from the catchment (topogenous fens) (Wheeler and Proctor, 2000). Valley fens that are located in water collecting depressions can receive water from both sources. [Valley fens are widespread in shallow water bodies in river or stream valleys with a slow flow of mineral-rich water \(e.g., Everglades, USA; Biebrza, Poland\), or in pools, lakes or other landscape depressions receiving a slow flow of discharging groundwater and/or surface water \(e.g., rich fens in Norfolk Broads, UK; Weerribben-Wieden, The Netherlands\) \(Lamers et al., 2015\). In addition, in boreal permafrost peatlands in Siberia and north America, the running water-controlled systems probably are common due to the difficulty of water penetration into the soil. However, it is difficult to provide a number for the percentage of peatlands globally that may be classified as valley fens, because of the complex spatial structure and gradients between different peatland types, and differences in terminology.](#) The spatial variation in the quantity and quality of incoming water creates spatial patterns in vegetation and microbial communities (e.g., methanogens and methanotrophs), and thus CH<sub>4</sub> production and oxidation, transportation and ultimately emissions to the atmosphere (Andersen et al., 2011; Juottonen et al., 2015; Kokkonen et al.,

2019; Robroek et al., 2015). Several studies have focused on the interactions of CH<sub>4</sub> with vertical water level fluctuations. For example, long-term lowering of the water level has been associated with a decreased abundance of *Sphagnum* mosses and aerenchymous plants, decreased CH<sub>4</sub> emissions and CH<sub>4</sub> production potential (Yrjälä et al., 2011). However, due to the heterogeneity of peatlands, inconsistent patterns can also be found. For instance, several studies have indicated that greater  
70 CH<sub>4</sub> emissions occur when the water level is close to the surface of the peatland (Bubier et al., 2005; Pelletier et al., 2007), while other studies have found maximum fluxes occurred at intermediate water levels (Turetsky et al., 2014), or found no connection between CH<sub>4</sub> emissions and water level (Euskirchen et al., 2019, Korrensalo et al., 2018). Nevertheless, water level has been suggested as a more important forcing factor on CH<sub>4</sub> cycling in fens than either temperature or vegetation composition alone (Laine et al., 2019; Mäkiranta et al., 2018; Riutta et al., 2020). In addition to vertical water level changes,  
75 the lateral flow of water in fens can be even more important in driving the processes that underpin CH<sub>4</sub> emissions, because flowing water not only ensures a water supply for the vegetation, but also transports nutrients, which benefits vegetation and microbial communities (Laitinen et al., 2007). At the same time, flowing water is likely to transport more oxygen (Ingram, 1983), thus enhancing CH<sub>4</sub> oxidation and suppressing production. While fens are typically the highest CH<sub>4</sub> emitters of all peatlands (Turetsky et al., 2014), the influence of lateral water flow on fen CH<sub>4</sub> emissions has not been studied to date.

80 At a global scale, climate warming is projected to continue in the decades ahead, while changes in precipitation patterns are projected to be more regional (Collins et al., 2013). The Coupled Model Intercomparison Project (CMIP5), under a RCP8.5 scenario, predicts a warmer and wetter climate for Fennoscandia (Collins et al., 2013). As peatland hydrology is driven by several processes, such as precipitation, lateral water fluxes, transpiration and evaporation, climate model predictions cannot be directly applied to infer peatland hydrological conditions (Helbig et al., 2020; Tuittila et al., 2007; Wu et al., 2010; Zhang  
85 et al., 2018), especially in minerotrophic fens. Nevertheless, peatland habitats can be impacted under both warming-dry and warming-wet scenarios (Bjorkman et al., 2018; Strack et al., 2006). In addition, fens may be more sensitive to water level changes than bogs; in particular, their plant communities have been shown to experience clear species turnover under drier conditions (Kokkonen et al., 2019). Aside from the vertical fluctuations in the water level, climate change is also likely to affect the water that enters fens as it will control the hydrological conditions within the catchments, e.g., the temperature sum  
90 in spring strongly controls the timing and amount of snowmelt water that enters the fen. This type of change in catchment conditions is likely to impact, for example, plant phenology and biomass production (Mäkiranta et al., 2018). This will, in turn, impact on C cycling between the peatland and the atmosphere due to different photosynthesis, decomposition and gas transportation rates, and on other factors at the plant functional type and even at the species levels (Hajek et al., 2009; Laine et al., 2011; Turetsky et al., 2008). Hence, a full insight into the complex climate-peatland-ecohydrology-CH<sub>4</sub> relationship is  
95 needed to predict the impact of changing catchment hydrology on fen CH<sub>4</sub> emissions under climate change scenarios. Prior to importing peatland-scale CH<sub>4</sub> emissions into global circulation models, we first need to bridge the gap of understanding as to how water flows control fen microhabitats and CH<sub>4</sub> emissions.

In this study, we aimed to assess the role of flowing water in regulating spatial variations in valley fen vegetation and CH<sub>4</sub> emissions. More specifically, we asked the following research questions: 1) How does a flowing stream within a valley fen impact microhabitat conditions, vegetation composition and biomass production? 2) Does the distance to a stream modify CH<sub>4</sub> fluxes? 3) How does vegetation composition and stream-related variables control CH<sub>4</sub> emissions? We hypothesised that: (H1) water table, temperature, oxygen concentration, vegetation structure and biomass are related to the proximity of the stream; (H2) spatial variation in CH<sub>4</sub> fluxes is related to the distance from the stream; (H3) regulation of CH<sub>4</sub> fluxes by the stream is mediated by the vegetation and by environmental variables, such as oxygen concentration.

## 105 2 Material and methods

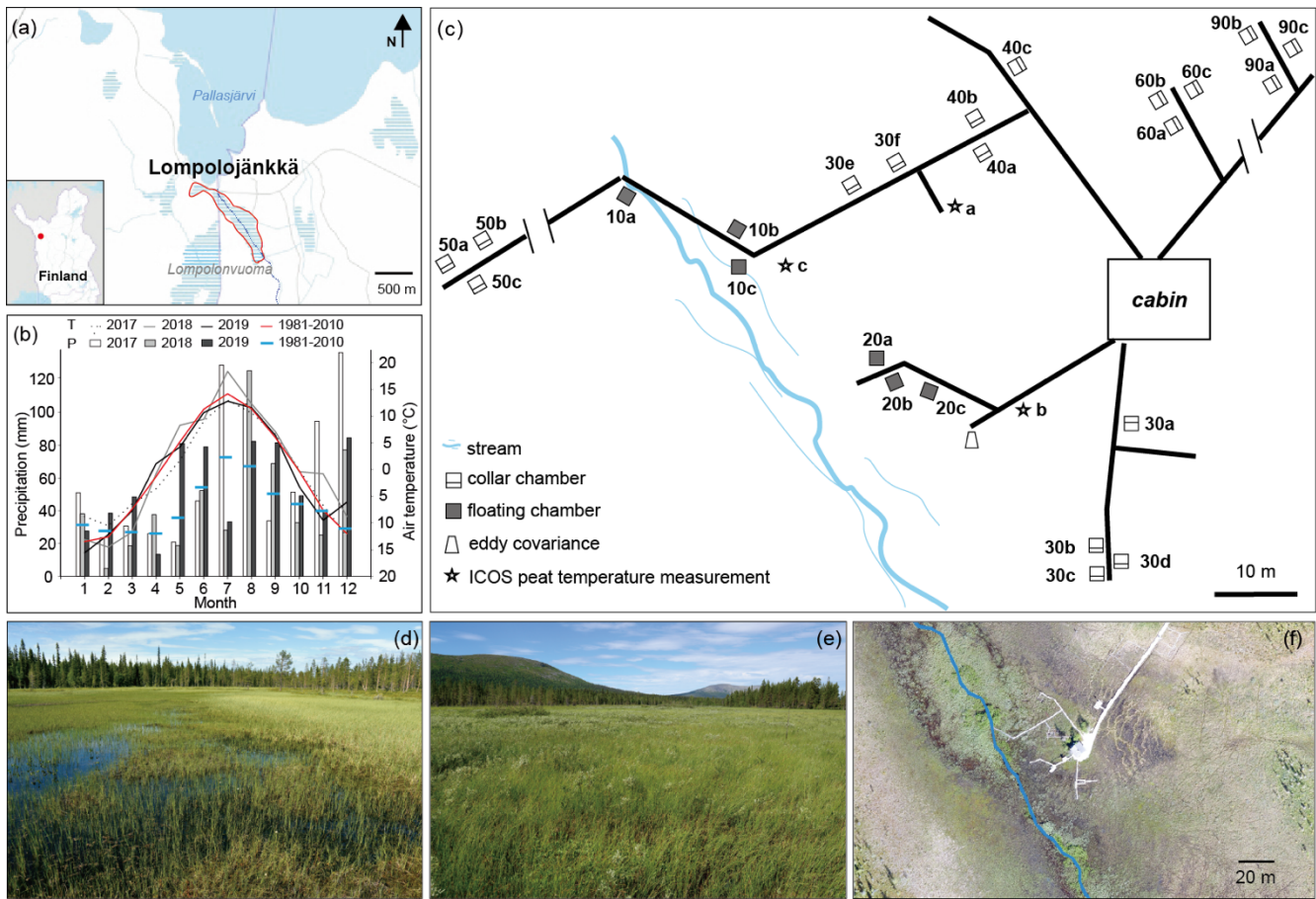
### 2.1 Study site

Lompolojänkkä (67.997 °N, 24.210 °E, 269 m a.s.l.) is one of the Finnish Integrated Carbon Observation System (ICOS) sites. It is an open mesotrophic sedge fen that is located in a valley in the hilly Pallas region of northern Finland (Figure 1). Based on the 30-year average (1981–2010; Kittilä Pokka meteorological station), the annual average temperature and total precipitation are -1.3 °C and 547 mm, respectively (Pirinen et al., 2012).

During the three flux measurement years (2017–2019), the summer of 2018 was exceptionally warm, up to 5 °C warmer than the long-term average (Figure 1b). Based on the ICOS continuous peat profile temperature measurements (at 5, 10, 20, 30, 50 and 100 cm; Figure 1c) in 2018, peat temperature at Lompolojänkkä varied along depth and also for different [locations of the site habitats](#) (Figure A1). During summer, peat temperature decreased from the surface to the deeper layers and the pattern was reversed for the other seasons. Peat temperatures were ~~greater~~ warmer in the drier parts of the study site compared to the wetter parts (closer to the stream) at all measured depths, and there were also larger temperature variations between the different depths [in the drier parts](#).

Peat accumulation at Lompolojänkkä initiated around 10,000 cal. yr BP (calibrated years before the present 1950 AD) and the ~~deepest maximum~~ peat depth is approximately 2.5 m (Mathijssen et al., 2014). [The peat thickness of the sampled area ranges from c. 1 to 2.5 m \(Mathijssen et al., 2014\). The site currently spans an area of c. 14 ha and is surrounded by boreal forests.](#) Almost the whole peatland is water saturated throughout the year. The relatively dense vegetation layer is dominated by different sedges (e.g., *C. rostrata*, *C. chordorrhiza*) in the wet areas and various deciduous shrubs (e.g., *Betula nana*, *Salix phylicifolia*) in the ~~relatively~~ dry areas. Moss cover (e.g., *Sphagnum* spp.) is patchy with ~~e-~~57 %-cover (Aurela et al., 2009). A small stream flows through the long and narrow valley fen (outlined in Figure 1a) and empties into the nearby Pallasjärvi lake. [The special catchment feature therefore creates both vertical and lateral water movement in the fen. However, the whole site clearly has different water transfer mechanisms operating in different areas of the site, and in different period of the year. For example, the lateral water movement mainly occurs in the center of the site that close to the stream.](#) The flow and size of the stream varies seasonally; being largest in spring after snow melt in the catchment. During

130 summer, the stream water level in many locations is below the vegetation surface and may not be visible (Figure 1d). For more detailed descriptions of Lompolojänkki, see Aurela et al. (2009) and Lohila et al. (2010).



135 **Figure 1:** (a) Location of the Lompolojänkki study site outlined in red, with the stream marked in blue. The base map was downloaded from the National Land Survey of Finland dataset under a CC 4.0 open source license. (b) Monthly air temperature (T) and precipitation (P) during 2017–2019, and long-term mean T and P values (1981–2010; Kittilä Pokka meteorological station <https://en.ilmatieteenlaitos.fi/statistics-from-1961-onwards>). The data for 2017–2019 were obtained from the nearest meteorological stations; Lompolonvuoma (for temperature) and Kenttäröva (for precipitation) (<https://en.ilmatieteenlaitos.fi/download-observations#!/>). (c) Schematic illustration of the field measurement setup. Note some parts may not be scaled accurately. (d, e) Photos of the study site. (f) Drone image of the field measurement area [on 20 August 2018](#) with the stream ~~indicated~~ marked in blue.

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[The catchment area of the stream has a size of 5.1 km<sup>2</sup> and is draining to Pallasjärvi lake a few hundred meters after leaving the fen. The lowest and highest points of the catchment area range from 268 to 375 m a.s.l. The soils consist mainly of glacial till, while the land cover at the catchment consists of coniferous and mixed coniferous-deciduous forests \(c. 80%\) and](#)

145 forested and open peatlands (c. 20%). Dominating tree species include Norway spruce (*Picea abies*), Scots pine (*Pinus sylvestris*) and Downy birch (*Betula pubescens*). The coniferous forests dominate at the catchment. Furthermore, some of the peatlands in the eastern part have been drained for forestry purposes during the same period. In such valley mires with streams, the watercourse is small compared to those with, e.g., large rivers and does not provide significant amounts of water through overbank flooding. However, they can form a complex mosaic of habitats around streams with small catchments, for example, at our site the central stream with a limited floodplain has developed a riparian strip characterised by e.g.,  
 150 *Equisetum fluviatile*, *Carex limosa* and *Salix lapponum*. The impact of flowing water on a particular site also depends on the shape of the site, compared to other sites that with streams presented, the long narrow shape of Lompolojänkkä therefore undergoes stronger effects by the stream than many other sites.

## 2.2 Sampling / sample plot set up

To quantify the spatial variability in CH<sub>4</sub> fluxes in the valley fen, we installed 15 permanent sample plots 60 cm x 60 cm (W x L) at varying distances from the stream in 2017 (Table 1). The sample plots were set up as ~~set~~ clusters of three to six plots that were typically located within a metre of each other. Initially, the closest ~~cluster~~ set from the stream was located within a 10-m distance, and the furthestmost at a 40-m distance. In 2019, we sought a better mechanistic understanding of the controls on CH<sub>4</sub> fluxes and so we added nine more sample plots, located in three ~~clusters~~ sets at 50, 60 and 90 m distance from the stream (Table 1).

160

**Table 1:** Methane (CH<sub>4</sub>) flux sample plot setup and measured variables at Lompolojänkkä. DTS: distance to stream; PCT: plant community type; T<sub>air</sub>: air temperature; T<sub>5</sub>: peat temperature at depth 5 cm below moss surface; WT: water table; DO20 and DO40: dissolved oxygen concentration at 20 and 40 cm below the peat surface; LAI: leaf area index.

Year	No. of plots	Plot codes	Measured environmental variables
2017	15	10a-c, 20a-c, 30a-f, 40a-c	DTS, PCT, T <sub>air</sub>
2018	15	10a-c, 20a-c, 30a-f, 40a-c	DTS, PCT, T <sub>air</sub>
2019	24	10a-c, 20a-c, 30a-f, 40a-c, 50a-c, 60a-c, 90a-c	DTS, PCT, T <sub>air</sub> , T <sub>5</sub> , WT, DO20, DO40, LAI

165 In total, 24 permanent gas flux measurement plots were established (Table 1, Figure 1c). The sample plots are coded according to their distance to the stream/visible flowing water as 10a-c (a ~~cluster~~ set within 10 m to the stream with three replicates a-c), 20a-c, 30a-f, 40a-c, 50a-c, 60a-c and 90a-c. The location of each plot was measured with a Trimble R8 GPS device with ±5 cm accuracy and the distance to the stream from each sample plot was calculated based on the National Land Survey of Finland topographic database.

## 170 2.3 CH<sub>4</sub> and CO<sub>2</sub> flux measurements

Seasonal CH<sub>4</sub> and CO<sub>2</sub> (dark respiration) fluxes were measured for three years (2017–2019) in sample plot [elusters-sets\\_10–40](#), and for one year (2019) in [series-sets\\_50–90](#). Measurements at [elusters-sets\\_10–40](#) were conducted eight times in 2017 (between 13 June–29 September), 11 times in 2018 (between 30 May–11 October) and 15 times in 2019 (between 20 May–11 September). At [elusters-sets\\_50](#) and 90, measurements were conducted 11 times in 2019, and 21 measurements were  
175 performed at [eluster-set\\_60](#). In total, these measurements yielded 126, 163 and 330 CH<sub>4</sub> flux records for 2017, 2018 and 2019, respectively.

For determining fluxes, the closed chamber method with fixed collars was used for [elusters-sets\\_30–90](#), and a floating chamber method without collars was employed for [elusters-sets\\_10](#) and 20 (Alm et al., 2007). The size of the opaque aluminium chamber was 60 cm x 60 cm x 40 cm (W x L x H) and each chamber was equipped with a fan. The sample gas  
180 was sucked from the chamber at a flow rate of 200–200 ml min<sup>-1</sup> using 50-m long tubing (d=6 mm) into a LGR gas analyser (LGR GCA-24p-EP, model 911-0011-0004, Los Gatos Research Inc., Ca, USA) located in a temperature-controlled cabin. The duration of one measurement was approximately 5 mins. The floating chamber (60 cm x 60 cm x 30 cm) was used at  
plots with permanently high, flowing water. In addition, gross primary production (GPP) was measured at [elusters-sets\\_10–40](#) using a transparent chamber on 24–25 July 2019 at the time of peak growing season. Same gas analyser as described  
185 above was used. Photosynthetically active radiation in the chamber was measured using a Kipp&Zonen PQS1 PAR Quantum Sensor (Kipp & Zonen B.V., Delft, the Netherlands). In order to fit a light-response curve to the net CO<sub>2</sub> exchange (NEE) data, NEE was first measured in full light, after which the chamber was covered with fabrics to create four different light levels (white shade, black shade, double black shade, and double black with green shade). In addition, one measurement with full shading to capture dark respiration was performed.

The CH<sub>4</sub> and CO<sub>2</sub> fluxes from each measurement were calculated from the linear slope ( $R^2 > 99\%$  for over 90 % measurements and  $R^2 > 90\%$  for other measurements) in gas concentration over time, taking into account chamber volume, chamber air temperature and air pressure at the measuring point. The volume in the chamber during each measurement was specified according to the instant ambient water level. The air temperature and air pressure data were derived from the nearest meteorological station, and air pressure was calibrated for each chamber, taking into account the altitude of the plot.  
190 We determined the GPP-light response curve for each sample plot (based on the NEE measurements with the transparent chamber), and derived sample plot specific GPP<sub>max</sub> values at a photosynthetic photon flux density level of 800  $\mu\text{mol m}^{-2} \text{s}^{-1}$ .  
195

## 2.4 Environmental data collection

To reach a mechanistic understanding of the spatial pattern of CH<sub>4</sub> fluxes, we collected data on the potential environmental factors that control emissions in combination with each flux measurement conducted in 2019. These factors were air and  
200 peat temperature, water table, dissolved oxygen concentration, leaf area index, and plant community cluster (Table 1).

Air temperature was either measured using a temperature sensor fixed inside the chamber or measured at 2-m height at the site (Lompolonvuoma meteorological station of Finnish Meteorological Institute (FMI)). Peat temperature was measured at 5 cm below the moss surface (T5) using a Pt100 thermometer (Omega HH376, Omega Engineering Inc., CT, USA). Water table relative to the moss surface (WT) was measured from a plastic tube installed in the peat next to each sample plot. Dissolved oxygen concentrations ([percent of air saturation](#)) at 20 (DO20) and 40 cm (DO40) below the surface (except [eluster-serieset](#) 60) were measured using a YSI Professional Series Digital handheld meter.

The leaf area index (LAI) of four vascular plant functional types (PFTs; deciduous shrub, evergreen shrub, forb and graminoid), and moss cover were estimated. The estimation of LAI followed Juutinen et al. (2017). First, we selected 31 square plots (50 cm × 50 cm) located within the fen and surrounding areas in July–August 2019, and estimated green projection cover (%) and measured mean height for each PFT in the plots. Second, to measure LAI of the samples, we harvested the aboveground parts of the vascular plant species, scanned them with an A4 scanner and calculated the proportion of green pixels in GIMP 2.8 (The GIMP Team, [www.gimp.org](http://www.gimp.org)). Third, we constructed empirical relationships between cover or plant volume (cover × height) and LAI with ordinary least squares (OLS) regressions for four PFTs found in the site. We chose the optimal predictor (cover or volume) by minimising the root mean square error value, and in the final models, the adjusted coefficient of determination (Adj.R<sup>2</sup>) varied between 0.73 and 0.89 (Table A1). Fourth, we used the equations from the OLS regressions to model seasonal LAI development curves for each CH<sub>4</sub> sample plot in which we had measured green projection cover and height for the four PFTs throughout the summer of 2019. Finally, we derived LAI values for each flux measurement time from the seasonal LAI development curves. We also calculated LAI values for the aerenchymous plants in each plot, which included *C. aquatilis*, *C. canescens*, *C. chordorrhiza*, *C. lasiocarpa*, *C. limosa*, *C. rostrata*, *Comarum palustre*, *Equisetum fluviatile*, *Eriophorum vaginatum*, and *Menyanthes trifoliata*. The calculation of aerenchymous LAI was carried out by applying the same OLS regression equations used for forb and graminoid PFTs to datasets that included only aerenchymous plant species.

In addition, we delineated four plant community types/clusters for the CH<sub>4</sub> sample plots as follows. First, we calculated the Bray-Curtis distance matrix of the plant species projection cover data from the sample plots and, in addition, 200 systematically sampled vegetation plots that were inventoried in the fen in 2018. Second, we derived four non-metric multidimensional scaling ordination (NMDS) axes from the distance matrix. Third, we delineated four plant community clusters from the NMDS axes with the Partitioning Around Medoids (PAM) method. The clustering was conducted in R with packages ‘vegan’ (Oksanen et al., 2019) and ‘cluster’ (Maechler et al., 2019). A map showing the location of the vegetation community clusters in the study site can be found in Figure A2.

## 2.5 Data analysis

NMDS was used to explore the linkages between peak season vegetation composition, distance to the stream, biomass production and flowing water. Peak season total LAI was used as a proxy for biomass production, and early summer DO20



and DO40 were used as proxies for flowing water/nutrient availability. For a robust analysis, plant species with occurrence lower than 3 % were excluded from the analysis.

235 Linear mixed-effect models were applied to the CH<sub>4</sub> flux and environmental data to identify the potential drivers of CH<sub>4</sub> flux using two different approaches. First, we explored the spatial variation in CH<sub>4</sub> fluxes by constructing a model with CH<sub>4</sub> data from all three measured years. Here, potential fixed predictors for CH<sub>4</sub> flux were distance to the stream, air temperature and the factorial plant community cluster. To account for repeated measurements, we included the nested random effects of year, month and measurement plot. Second, to gain [more](#) mechanistic understanding of the controls on CH<sub>4</sub> fluxes, we used a  
240 dataset with additional variables gathered during 2019. Here, potential fixed predictors were DO20, DO40, T5, air temperature, WT, GPP<sub>max</sub>, LAI of all vascular, aerenchymous and ericoid plants, moss cover (% coverage), CO<sub>2</sub> dark respiration, distance to the stream, and the factorial plant community cluster. To account for repeated measurements from the plots over the growing season, we included the crossed random effects of measurement day and plot.

In building the models, we manually added the potential fixed predictors one by one and tested whether the resultant, more  
245 complex model was significantly better than the model without the added predictor, using conditional *F*-test and Akaike information criterion (AIC). To account for the nonlinear relationship between CH<sub>4</sub> flux and some environmental variables (such as temperature), we tested several response shapes for the fixed predictors: i) linear response, ii) quadratic response, iii) linear response above or below a certain threshold value, but constant otherwise, and iv) quadratic response above or below a certain threshold value, but constant otherwise. In cases iii) and iv), the response type and threshold value were determined  
250 visually by plotting the residuals of the previous model against the fixed predictor to be added. The final response shape and threshold value were selected based on the conditional *F*-test and AIC values. Furthermore, we tested the interactions between all fixed predictors in the final models and only included those predictors that led to a significant improvement in model performance. The first explorative model was fitted with function `lme` of the package ‘nlme’, and the second, more  
complex [and](#) mechanistical model was fitted with function `lmer` of the package ‘lme4’ in R.

255

### 3 Results

#### 3.1 Variations in vegetation and environmental factors

The studied valley fen exhibited clear but distinctive patterns in vegetation composition, WT, LAI, and DO concentrations related to distance from the stream (Figures 2 and A2–5). Moreover, the temporal patterns in WT and DO concentration  
260 showed distinct variations at locations further away and closer to the stream, respectively (Figure A4).

In total, four plant community types were identified (Figure 3, Table A2). Community type (1) fluvial, which was found in the wetter parts of the fen, was dominated by *E. fluviatile* and *C. limosa*. Community type (2) riparian represented riparian vegetation that were taller, such as *C. aquatilis*, *S. lapponum*, *S. phlycifolica* and *Comarum palustre*. Community type (3) lawn, and community type (4) hummock contained vegetation typical of drier fen conditions, with the hummock type found

265 in the driest areas. The dominant species in these community types included *S. riparium*, *Vaccinium oxycoccos* and *C. livida*  
(lawn), and *S. russowii*, *V. uliginosum*, *Betula nana* and *Rubus chamaemorus* (hummock). The overriding pattern was related  
to the distance to the stream (Figure 2a and A2), i.e. fluvial and riparian community types were recorded in the locations  
closest to the stream, while lawn and hummock types were located at the plots furthest from the stream. In addition, the plant  
communities in the sample plots were suggestive of a spatially heterogeneous structure in the fen, i.e. different types were  
270 recorded within a short distance (Figures 2a and 3). The NMDS ordination (Figure 3) revealed that the main pattern in  
vegetation structure related to the distance to the stream was correlated strongly with, and was better explained by, peak  
season oxygen concentration. Total LAI increased with peak season oxygen concentration, which was negatively correlated  
with distance. Aquatic species, such as *C. aquatilis* and species that typically benefit from moving water, such as *S.*  
*lapponum*, *C. palustre* and *M. trifoliata*, exhibited relatively high positive values on the first NMDS axis, revealing a strong  
275 relationship between the stream and some specific plant species. Species adapted to drier surfaces, such as ~~mosses~~  
~~*Rhizomnium* sp. and *S. warnstorfii*, and the sedge *E. vaginatum*~~, were located at the other end of the axis. As peak season  
GPP data were only available for ~~clusters~~ 10–40, they were not included in the NMDS analysis, but were analysed  
separately against oxygen concentration and total vascular LAI data (Figure A6). GPP was clearly higher closer to the stream  
( $> 0.45 \text{ mg CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ ) than further from the stream ( $< 0.35 \text{ mg CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ ). In addition, GPP was strongly related to total  
280 vascular LAI, at least when LAI  $< 2$ . In the only sampling point with a LAI value  $> 2$ , GPP did not increase any further.

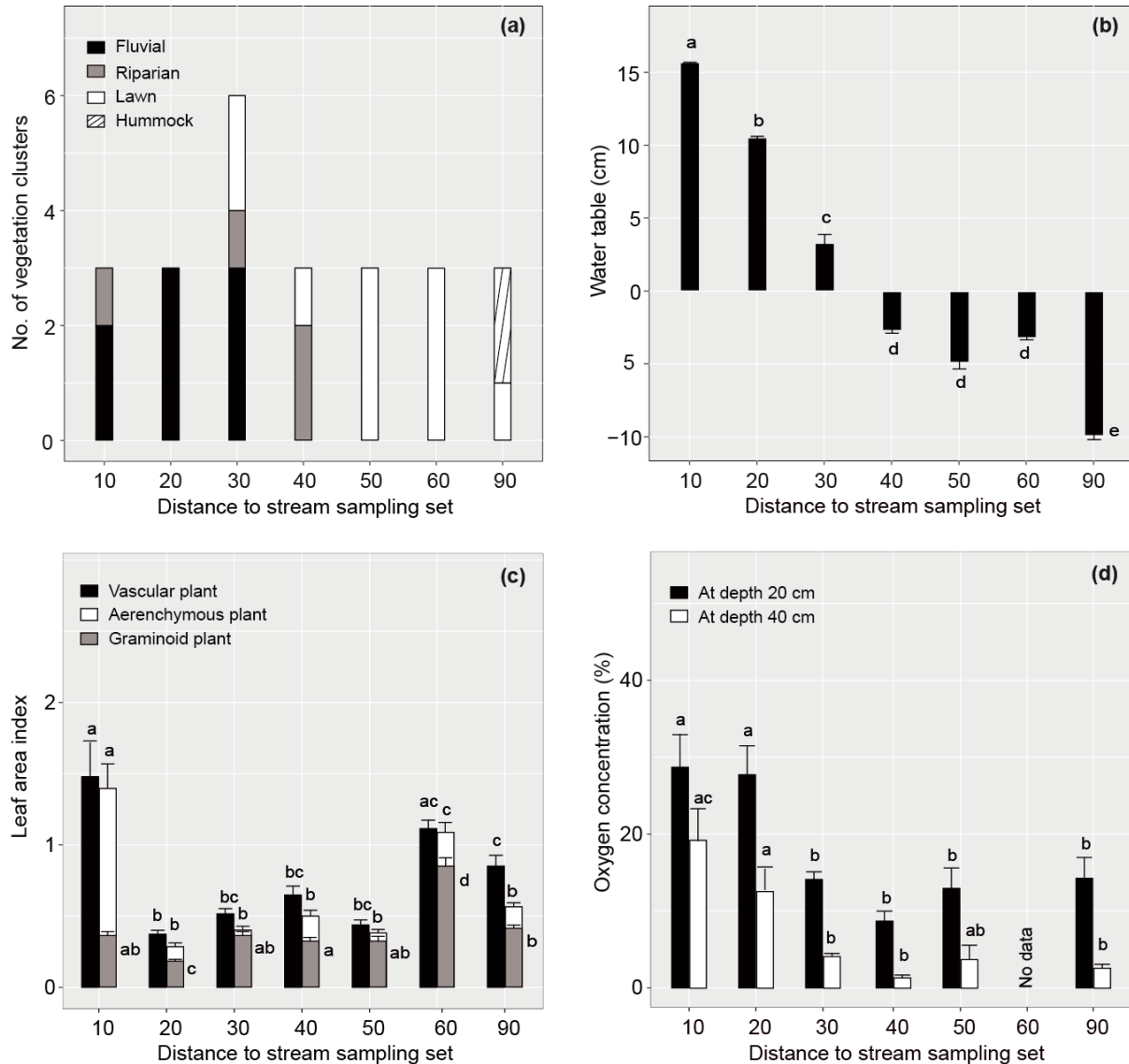
The WT pattern at the sample plots was strongly linked to their distance to the stream, i.e. WT was higher closer to the  
stream (Figure 2b). At ~~sample set~~ 10 and 20, close to the stream, there was approximately 10 cm of water above the  
peat surface, while at ~~cluster set~~ 90, furthest from the stream, the WT was approximately 10 cm below the surface. The other  
~~clusters~~ displayed intermediate WT values. In general, the lowest (deepest) WT levels were measured at all sample plots  
285 during late July, when precipitation was low and air temperature had reached the seasonal peak (Figures A4 and A7).

The sample plots located next to the stream (~~set~~ 10) showed significantly larger mean seasonal vascular LAI values  
(mean 1.5) but were similar to ~~set~~ 60 (with lawn vegetation) (Figure 2ac). ~~Clusters-Sets~~ 10 and 60 both showed  
significantly higher aerenchymous LAI values than the other ~~clusters-sets~~ ( $\sim 0.5$ ), although ~~set~~ 10 (mean 1.4) had a  
significantly higher value than ~~set~~ 60 (mean 1.1) (Figure 2c). Plot 10a appeared to be an outlier with higher total and  
290 aerenchymous LAI values ( $\sim 4$ ) than the other plots ( $< 2$ ), which was attributed to the presence of the abundant forb *C.*  
*palustre* in that plot (Figure A3). Graminoid LAI values (that excluded *C. palustre* and two other occasionally recorded  
species: *M. trifoliata* and *E. fluviatile*) were significantly higher in ~~set~~ 60 (mean 0.8) than in the other ~~set~~ ( $<$   
0.5). The development of LAI showed a clear seasonal pattern (Figure A5), with the peak occurring around late July.

Dissolved oxygen concentrations at both 20 and 40 cm depths showed a similar spatial pattern, with higher concentrations  
295 recorded close to the stream (in ~~set~~ 10 and 20) (Figure 2d). However, large temporal variations existed in DO values  
at both the 20 and 40 cm depths, which generally peaked in early summer during a high flow of water (Figure A4). Also, DO

concentrations showed a similar temporal pattern to precipitation, with higher concentrations recorded during periods with higher precipitation (Figures A4 and A7).

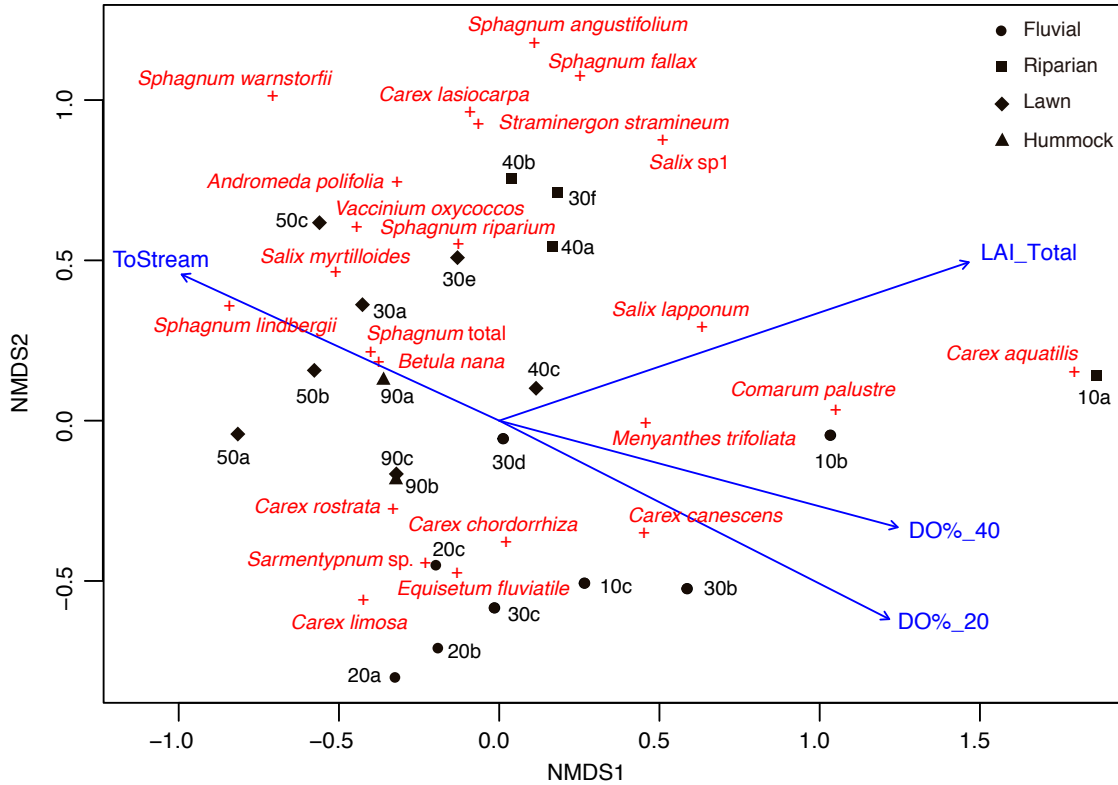
The proximity of the stream reduced the temporal variation in the peat temperature measured at 5 cm depth in 2019 (Figure A4); while the temperature at sample plots further away (clusters 50–90) varied between 3 and 23 °C, and the temperature at sample plots close to the stream (clusters 10–40) stayed between 7 and 15 °C.



**Figure 2:** Spatial variation in vegetation and environmental factors in relation to the distance of the sample plot from the stream in summer 2019. (a) Occurrence of different plant community types, (b) mean ( $\pm$  standard error) water table relative to peat surface (cm), (c) vascular, aerenchymous and graminoid plant leaf area index, and (d) dissolved oxygen concentration at 20 and 40 cm below the peat

surface for each sample plot [clusterset](#). Note that dissolved oxygen concentration was not measured at [cluster-set](#) 60. Different letters above the bars indicate significant differences ( $p < 0.05$ ) between the sample plot [clustersets](#) calculated using Tukey's Honest Significant Difference method. For species composition in the different plant community types, see Figure 3 and Table A2.

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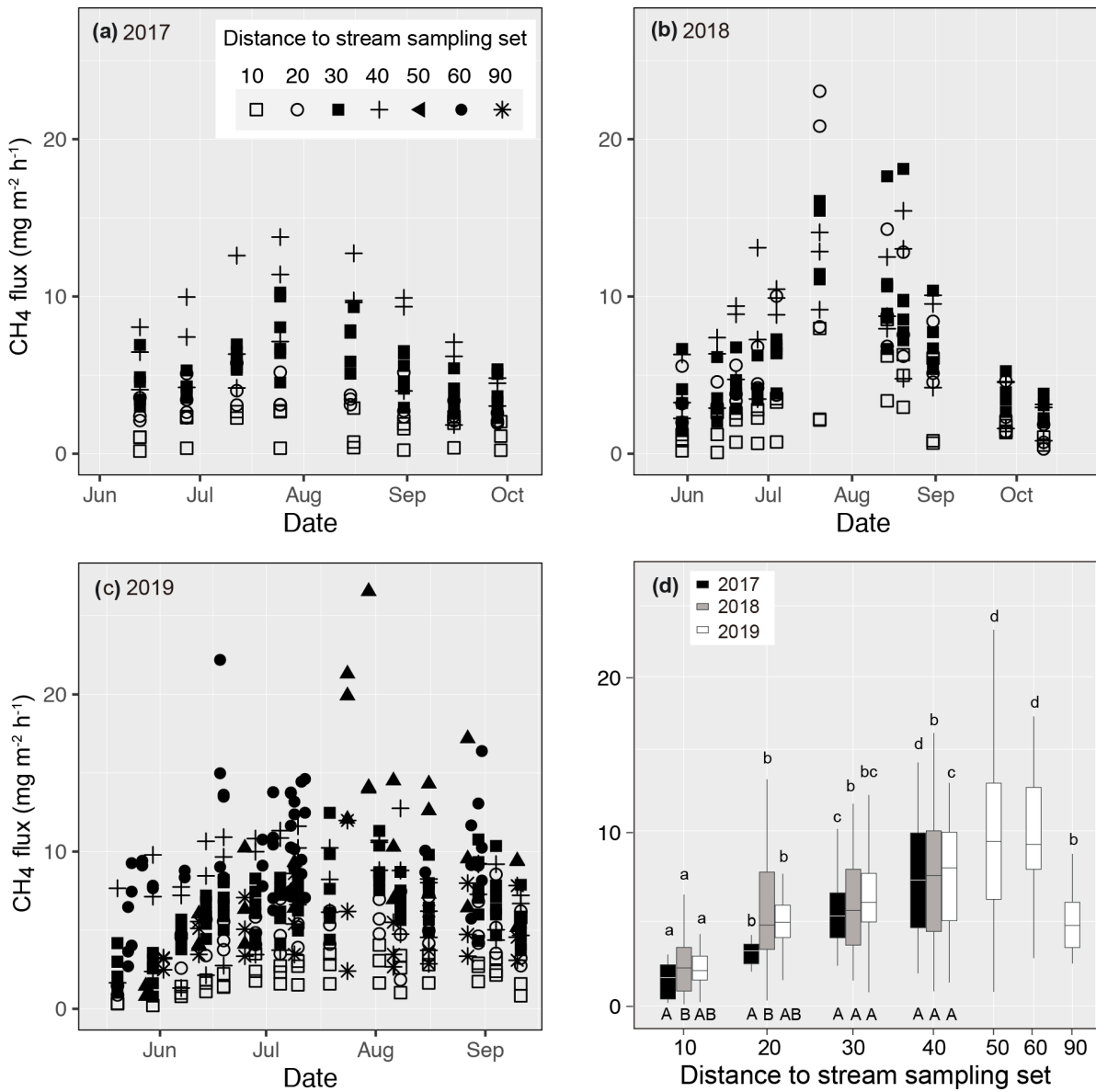
**Figure 3:** Non-metric multidimensional scaling (NMDS; stress 0.15) ordination showing peak season (late July) vegetation structure in the sample plots, including distance to stream (ToStream,  $p = 0.042$ ), peak season (early summer) dissolved oxygen concentration at 20 cm (DO%\_20,  $p = 0.006$ ) and 40 cm depths (DO%\_40,  $p = 0.015$ ), and peak season (late July) total vascular plant leaf area index (LAI\_Total,  $p = 0.003$ ) as fitted environmental variables. Four plant community types that were derived from the regional vegetation data are indicated using different symbols.

315

### 3.2 Variations in CH<sub>4</sub> fluxes

Measured CH<sub>4</sub> fluxes ranged from 0.16 to 13.78, 0.08 to 23.05, and 0.21 to 26.55 mg m<sup>-2</sup> h<sup>-1</sup> in 2017, 2018 and 2019, respectively (Figure 4). In all three years, CH<sub>4</sub> fluxes increased gradually from the early summer, peaking in early August, after which the fluxes decreased. In 2018 and 2019, higher fluxes (> 20 mg m<sup>-2</sup> h<sup>-1</sup>) were observed in the middle of the growing season compared to 2017 (< 15 mg m<sup>-2</sup> h<sup>-1</sup>).

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**Figure 4:** Measured methane (CH<sub>4</sub>) fluxes during the growing season in (a) 2017, (b) 2018, and (c) 2019 at Lompolojänkä. Distance to the stream of the sample plot [clusters-sets](#) are indicated by different symbols. (d) Box plots of CH<sub>4</sub> fluxes for each measurement [cluster-set](#) in 2017, 2018 and 2019. The different letters on top of the box plots indicate significant differences ( $p < 0.05$ ) between the [clusters-sets](#) for each year, calculated using Tukey's Honest Significant Difference method, and the different letters below the box plots indicate significant differences ( $p < 0.05$ ) between the studied years for each [cluster-set](#).

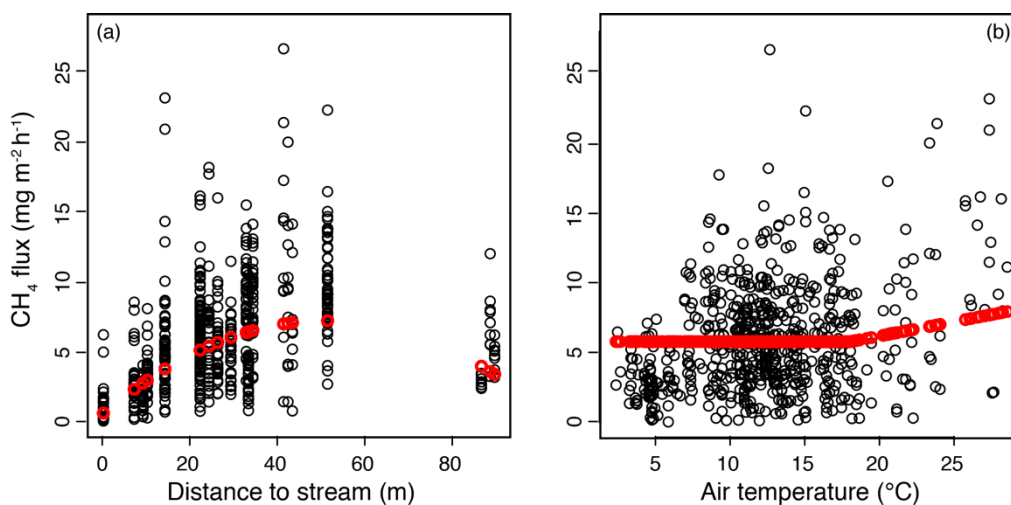
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Even though there were variations in CH<sub>4</sub> fluxes within ~~each of the replicated the~~ sample plots for each ~~elusters~~ sample sets, clear spatial patterns related to their distance to the stream were also evident (Figure 4). In all three years, CH<sub>4</sub> fluxes next to the stream (~~eluster-set~~ 10) were the lowest. In 2019, when additional sample plots were established (see Materials and methods), fluxes peaked at a 50 m distance from the stream. In the previous two years, when sampling only took place to a distance of 40 m from the stream, there was an increasing trend in fluxes with distance. In 2017, CH<sub>4</sub> fluxes measured from the various sample plot ~~elusters-sets~~ were significantly different from each other. In 2018, CH<sub>4</sub> fluxes from ~~elusters-sets~~ 20–40 were similar, but were significantly higher than the ~~elusters-sets~~ located next to the stream (~~eluster-set~~ 10).

Close to the stream, CH<sub>4</sub> emissions differed between years; emissions from ~~elusters-sets~~ 10 and 20 were significantly lower in 2017 than in 2018, while emissions in 2019 were intermediate and did not differ from the previous two years. At the intermediate distance (~~elusters-sets~~ 30 and 40), CH<sub>4</sub> emissions were at the same level in all three years.

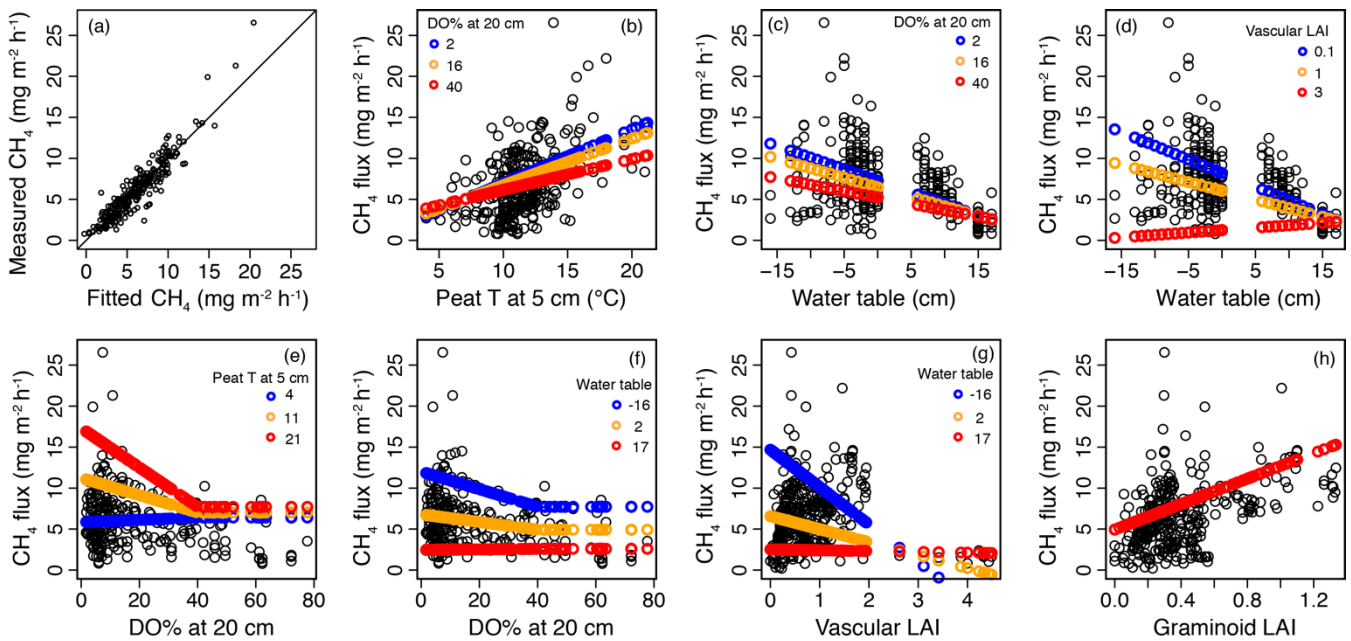
### 3.3 Response of CH<sub>4</sub> fluxes to environmental forcing

In the mixed-effect model (three-year dataset), which was constructed to examine spatial variability, CH<sub>4</sub> fluxes were controlled by the distance to the stream and by air temperature (fixed predictors), while plant community type was not a significant predictor when distance to the stream was included (Table A3a). There was a quadratic relationship between CH<sub>4</sub> fluxes and distance to the stream, with the highest fluxes observed at an intermediate distance (Figure 5a). There was a positive linear correlation between air temperature and CH<sub>4</sub> fluxes only at temperatures above a threshold value of 18 °C. Below that threshold, CH<sub>4</sub> fluxes remained unaffected (Figure 5). ~~There was a significant interaction between distance to the stream and air temperature ( $p = 0.03$ ), with the greater impact of temperature on CH<sub>4</sub> flux observed at plots closer to the stream (Figure 5b).~~



**Figure 5:** Results of mixed-effect model constructed to examine the spatial variation in methane (CH<sub>4</sub>) fluxes in the valley fen using a three-year dataset. Response curves (in red) of CH<sub>4</sub> flux to (a) distance to stream, with air temperature kept constant at 18 °C, (b) air temperature, with distance to stream kept constant at 1 m (blue), 276 m (mean value of the dataset) (pink) and 60 m (red).

In the second model (2019 dataset), which was constructed to provide a robust mechanistic understanding of CH<sub>4</sub> dynamics in the fen, temporal and spatial variation in CH<sub>4</sub> flux were found to be best explained by peat temperature at 5 cm (T5), WT, DO concentration at 20 cm below the surface (DO20), graminoid LAI and vascular LAI as fixed predictors (Figure 6, Table A3b). When DO20 was included, the distance to the stream and plant community type were not significant predictors. Of these predictors, DO20 linearly decreased the flux until a threshold value of 40 % was reached, above which it remained constant, while there was a linear relationship between CH<sub>4</sub> fluxes and the other predictors (Figure 6). Both T5 and graminoid LAI were observed to linearly increase CH<sub>4</sub> fluxes, while fluxes were negatively correlated with WT and vascular LAI (i.e., fluxes were lower at higher water levels and greater vascular LAI values). DO20 interacted with T5 and WT (Figure 6e, f), so that DO20 decreased CH<sub>4</sub> flux more steeply at lower-higher T5 and lower WT values. Also, T5 and WT responses were steeper at low DO20 values. Furthermore, vascular LAI had less impact on CH<sub>4</sub> flux at high WT levels.



**Figure 6:** Results of the mixed-effect model, which was constructed to provide a robust mechanistic understanding on the controls of methane (CH<sub>4</sub>) flux in the valley fen using the 2019 data set. (a) Measured CH<sub>4</sub> flux values plotted against fitted values of the mixed-effect model. (b-h) Response curves of CH<sub>4</sub> flux to (b) peat temperature (T) at 5 cm below the surface, (c, d) water table, (e, f) dissolved oxygen concentration (DO%) at 20 cm below the surface, (g) vascular plant leaf area index (LAI) and (h) graminoid plant LAI with the interactive variable adjusted at three-four different levels (rounded minimum, average and maximum values of the observed dataset; indicated by

375 ~~threefour~~ different colours) and the other variables kept constant. When setting variables as constant, the median values of the dataset were chosen, i.e. 11 °C for peat T at 5 cm, -11 (response to DO % at 20 cm) and -1 cm (other responses) for water table, 7 for DO% at 20 cm (when < 40 %), 0.6 for vascular LAI and 0.3 for graminoid LAI.

## 4 Discussion

### 4.1 The role of the stream in driving fen vegetation and biomass production

380 As hypothesised, the spatial heterogeneity in environmental variables in this valley fen site was highly related to the distance from the stream. Peatlands typically have spatially heterogeneous microhabitats due to wide variations in hydrology and nutrient availability (Rydin and Jeglum, 2013), which impact microbial activities and subsequent CH<sub>4</sub> emissions (Juottonen et al., 2005; Juottonen et al., 2015; Noyce et al., 2014; Ström et al., 2003). Water table level is one of the most important influences on plant occurrence and growth in peatlands (Rydin and Jeglum, 2013), and in this study, was highest closer to  
385 the stream. As a result, hydrophilic species, such as *C. aquatilis*, *S. lapponum*, *C. palustre*, and *M. trifoliata*, were abundant in places close to the stream. Even though we did not measure the chemical composition of the water, the abundance of these species implies a minerogenic supply established by water flow (Wassen et al., 1990).

The observed positive link between early summer oxygen concentrations (a proxy for flowing water) and total LAI further confirmed that flowing water likely delivers more nutrients and better supply plant growth and photosynthesis, and therefore  
390 provides more C substrates for microbial activities (Bellisario et al., 1999). In addition, GPP, the key driver of the peatland C cycle (Whiting and Chanton, 1993) and influences peatland vegetation composition and abiotic factors, such as air temperature and water level (Peichl et al., 2018), was consistently higher in plots located nearer the stream. Similarly, dissolved oxygen concentrations that acted as a proxy of the mineral-nutrient rich water were also higher in those plots. It has been shown that increased water supply alone can cause substantial increases in biomass on nutrient-rich soils, while  
395 fertilisation/nutrient addition has little effect (Eskelinen and Harrison, 2015). As such, the forbs and mosses that dominate such wet fens might benefit from higher water tables for biomass production (Mäkiranta et al., 2018). In this study, as the stream can bring both water and nutrients to the site at the same time, we are not able to distinguish whether the impact of the stream on the vegetation at our site was caused by the water or by nutrient supply, or both. Nevertheless, our results suggest that flowing water acted as a decisive factor on peatland vegetation composition and biomass production, through  
400 the addition of either water or nutrients. Therefore, the stream is likely to play a key role in regulating peatland CH<sub>4</sub> emission patterns.

### 4.2 Role of stream-induced microhabitats in driving CH<sub>4</sub> emissions

Consistent with our second hypothesis, the overall pattern of CH<sub>4</sub> fluxes showed clear spatial variations in relation to the distance from the stream. The impact of the stream was greater than the influence of vegetation community types, which  
405 represent general microform conditions and have been commonly reported to regulate CH<sub>4</sub> emissions (e.g., Riutta et al.,



2007). Specifically, as expected in the third hypothesis, factors such as peat temperature at 5 cm depth (T5), WT, DO20 and LAI, which were to some extent shaped by the stream, were all significant factors in driving CH<sub>4</sub> emissions at this site.

Our data suggest that CH<sub>4</sub> emissions increased with higher T5 values, similarly to many previous studies (e.g., Korrensalo et al., 2018; Rinne et al., 2018). Rising temperature is known to increase the activity of CH<sub>4</sub> producing microbes, and also enhance CH<sub>4</sub> transport through aerenchymous plants (Dunfield et al., 1993; Grosse, 1996; Kolton et al., 2019). ~~In support of this finding, higher DO20 values were found to decrease temperature sensitivity. Moreover, the temperature sensitivity in our study site was stronger closer to the stream, which is possibly due to a higher dissolution rate in cold water.~~ The pattern implies that in the fertile conditions next to the stream, higher oxygen concentrations in the cool water limits emissions by suppressing CH<sub>4</sub> production or by enhancing oxidation, and that warming of the water removes this limitation. ~~In support of this finding, higher DO20 values were found to decrease temperature sensitivity.~~ Similar to the CH<sub>4</sub> response to T5, higher DO20 values also reduced the impact of WT position on CH<sub>4</sub> emissions. Both responses highlight the importance of oxidation when considering how CH<sub>4</sub> emissions respond to environmental changes (Song et al., 2020). The patterns might also indicate higher CH<sub>4</sub> production under warmer conditions within the catchment and, consequently, on higher CH<sub>4</sub> concentrations in the flowing water (Juutinen et al., 2013). However, in this study we were not able to determine the origin of the emitted CH<sub>4</sub>.

In our sampling campaign, WT levels were observed both above and below the soil surface, and CH<sub>4</sub> emissions were found to generally decrease with rising WT levels. This decrease is in contradiction with many other studies that mainly cover sites with WT levels below the soil surface (Bubier et al., 2005; Pelletier et al., 2007; Turetsky et al., 2008). However, low emissions were also observed in the drier parts of the fen, which is in agreement with previous studies, in addition to very low emissions observed close to the stream. The lower emissions and a generally unimodal response to WT level were overridden in the whole dataset by the much stronger pattern captured close to the stream. Two plausible explanations for the observed WT–CH<sub>4</sub> emission pattern are, 1) the potential CH<sub>4</sub> production zone is smaller and the potential CH<sub>4</sub> oxidation zone is greater in drier conditions (Lai, 2009), and 2) in wet conditions, where there is moving water, lower CH<sub>4</sub> emissions can be attributed to enhanced CH<sub>4</sub> oxidation in the oxygen-rich water column, and a lower CH<sub>4</sub> production rate due to the presence of oxygen (Bubier, 1995). Also, ~~lower-cooler~~ peat temperatures due to the higher water table and flowing water likely contribute to a lower CH<sub>4</sub> production rate.

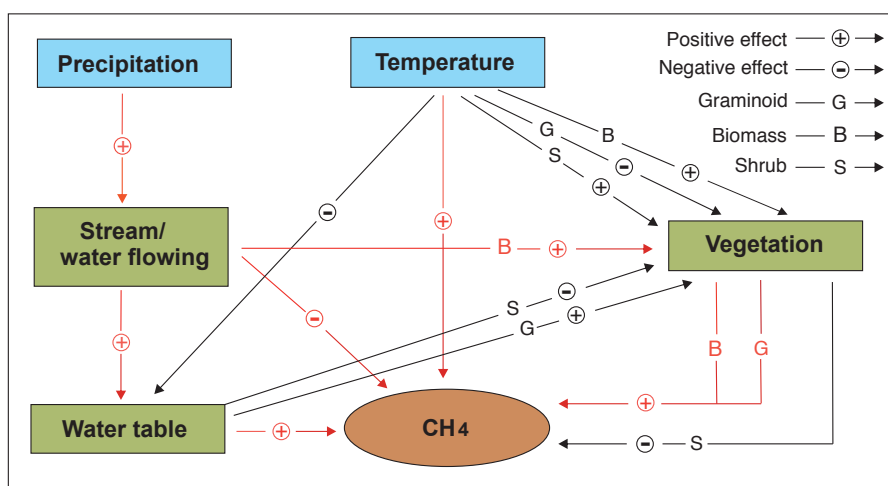
In our study, vascular LAI was found to have a negative linear correlation with CH<sub>4</sub> emissions. Plots nearest the stream had the highest vascular LAI values but the lowest CH<sub>4</sub> fluxes, i.e. the impact of the stream was again predominant over the impact of LAI. Studies have shown that shrubs can hinder CH<sub>4</sub> production because of their poor-quality substrate for methanogenesis (Riutta et al., 2020, Yavitt et al., 2019), although the cover of shrubs at our study site was very small. Therefore, low CH<sub>4</sub> emissions at high vascular LAI values is likely due to *in situ* lower peat temperature and the higher oxygen concentrations caused by the moving water. As aerenchymous LAI showed a very similar pattern to vascular LAI, it was not included in the mechanistic model. Instead, graminoid LAI, which was not impacted by the stream, showed a

440 positive link with CH<sub>4</sub> emissions, which is in line with several previous studies (e.g. Bhullar et al., 2013ab). The exceptionally high CH<sub>4</sub> fluxes measured at [cluster-set\\_50](#) where the graminoid LAI was low is potentially linked to one aerenchymous species growing in the [clusterset](#), i.e. *Eriophorum vaginatum*, which can enhance CH<sub>4</sub> release and increase C substrate input to methanogens (Greenup et al., 2000).

445 In general, CH<sub>4</sub> emissions in stream-dominated fens are likely to show a quadratic response pattern in regard to their distance to the stream, with low emissions occurring at both the closest and farthest locations from the stream, mainly due to high oxygen concentrations and water depletion, respectively. The highest CH<sub>4</sub> emissions were found in places at intermediate distances to the stream, which benefit from both sufficient water and nutrient supply but have inherently low soil oxygen concentrations. However, we acknowledge the challenge of defining the stream at our site due to the seasonal variation in catchment hydrological conditions. Hence, this study only demonstrates the stream-CH<sub>4</sub> emission pattern, rather than providing quantitative information for projections.

### 450 4.3 Future peatland CH<sub>4</sub> emission trajectories under climate change

Projection of global peatland CH<sub>4</sub> emissions under different climate change scenarios is a major challenge due to the reported variabilities in emissions, and also because of the interactions between the various environmental predictors (Strack and Waddington, 2007; Strack et al., 2004; Weltzin et al., 2000; Zhang et al., 2002). Our study further highlights that the impacts of climate change on CH<sub>4</sub> emissions in flow-through peatland systems are even more complicated due to the additional effects of the flowing water, which poses a challenge for accurate predictions of the global CH<sub>4</sub> budget. 455 Nevertheless, despite the complexity, clear patterns emerged that are informative in placing current peatland habitat-based CH<sub>4</sub> emission measurements into a broader context, and supplement the current understanding of peatland CH<sub>4</sub> emissions (Figure 7).



460

**Figure 7:** Schematic illustration showing the potential independent and interactive impacts of precipitation, temperature, vegetation and hydrology on methane (CH<sub>4</sub>) fluxes in northern peatlands. Arrows in red and black are derived from this study and previous studies, respectively (e.g., Mäkiranta et al., 2018; Roulet et al., 1992; Yavitt et al., 2019).

465 The majority of peatlands are located in northern high latitudes where the climate is currently experiencing a greater rate of  
change than in other regions (Collins et al., 2013). Climate warming is expected to promote microbial activity, and therefore  
faster C cycling. However, warming in tandem with drying has been shown to decrease the methanogenic archaea  
populations (Peltoniemi et al., 2015). In our study, vegetation composition, as such, was not a significant controller of CH<sub>4</sub>  
emissions, although biomass production (GPP and LAI), influenced by the stream, was a very important controller as it  
470 likely provides substrates for methanogens. However, this is in contradiction with the suggestion that vegetation mainly  
influences CH<sub>4</sub> emissions at minerotrophic sites by facilitating transportation, while at ombrotrophic sites it is through  
substrate-based interactions regulated by plant photosynthetic activity (Oquist and Svensson, 2002). Climate warming and/or  
peat surface drying can alter vegetation composition and affect the contribution of the biomass that is produced. For example,  
shrubs can benefit from these environmental changes, while forbs and mosses may suffer (Kokkonen et al. 2019, Mäkiranta  
475 et al., 2018, Strack et al. 2006). Even though such hydroclimatic impacts on vegetation might be modified by nitrogen  
availability (Luan et al., 2019), high latitudes generally experience little nitrogen deposition (Du et al., 2020). The abundance  
and functional types of the plants, especially graminoid plants, regulate CH<sub>4</sub> fluxes, but such impacts might be overruled if  
the water table level drops substantially (Riutta et al., 2020). In addition, there is some evidence of microtopographic  
differences in peatland nutrient dynamics in response to drying (Macrae et al., 2013), whereas flowing water will benefit the  
480 nutrient supply at a specific site. Furthermore, the expansion of shrubs, in response to drying, might potentially decrease peat  
temperatures due to increased shading and the evaporative cooling effect (Strakova et al., 2012), and thereby reduce CH<sub>4</sub>  
emissions.

Flowing water also tends to keep the peat temperature lower, even though fens with moving water warm up earlier than other  
peatlands in the spring and early summer (Rydin and Jeglum, 2013). In contrast to temperature predictions, predicting  
485 precipitation remains more uncertain, although in general, it is expected to increase in several regions (Collins et al., 2013).  
Although peatland hydrological processes are not directly impacted by precipitation due to, for example, evapotranspiration  
and runoff, it has been shown that precipitation can decrease CO<sub>2</sub> uptake and GPP due to cloudiness and associated reduced  
light availability (Nijp et al., 2015), thus influencing CH<sub>4</sub> emissions. Precipitation can also cause more dynamics of water  
and decrease CH<sub>4</sub> emissions by providing more oxygen for CH<sub>4</sub> oxidation (Mitchell and Branfireun 2005, Radu and Duval  
490 2018), which can be further accelerated under a warmer and drier peat surface scenario.

## 5 Conclusions

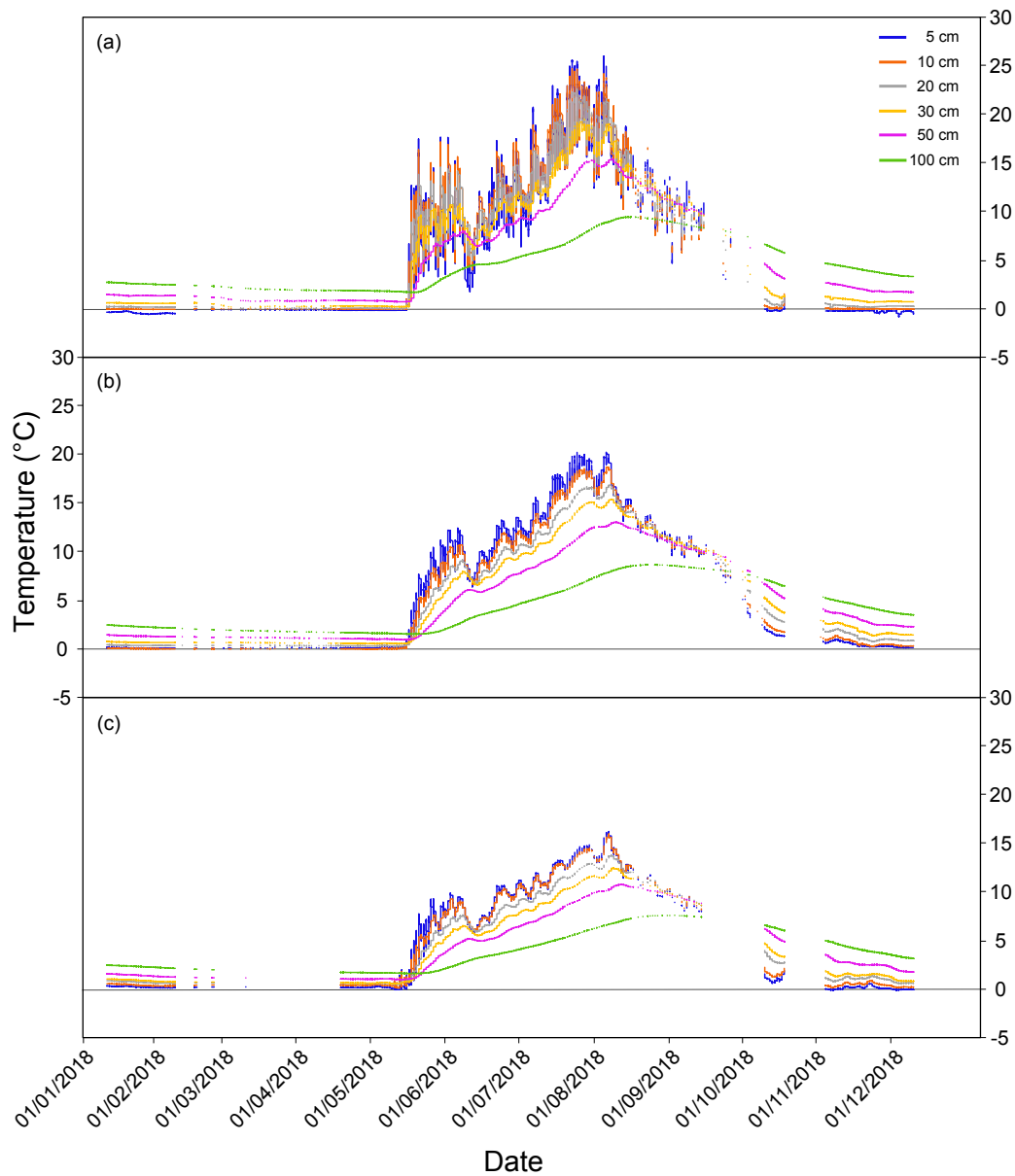
Our data from a flow-through valley fen demonstrates that hydrology in northern fen systems has a dual role in controlling CH<sub>4</sub> emissions, depending on the presence or absence of a stream. Flowing water not only enhances nutrient transportation and oxygen availability, but also decreases peat temperature, all of which are significant direct or indirect controllers of CH<sub>4</sub> emissions. At places close to the stream there were higher water levels, lower peat temperatures, and greater oxygen concentrations; these supported the highest total leaf area and gross primary production rates but resulted in the lowest CH<sub>4</sub> emissions. Further from the stream, the conditions were drier and CH<sub>4</sub> emissions were also low. CH<sub>4</sub> emissions were highest in the intermediate distance from the stream where oxygen concentration in the surface peat was low but gross primary production was still high. Our results show how a stream controls CH<sub>4</sub> emissions in a flow-through fen, which is a common peatland ecosystem type from the arctic to the temperate zones. Therefore, future projections of the global CH<sub>4</sub> budget need to take into account flowing water features in fen systems.

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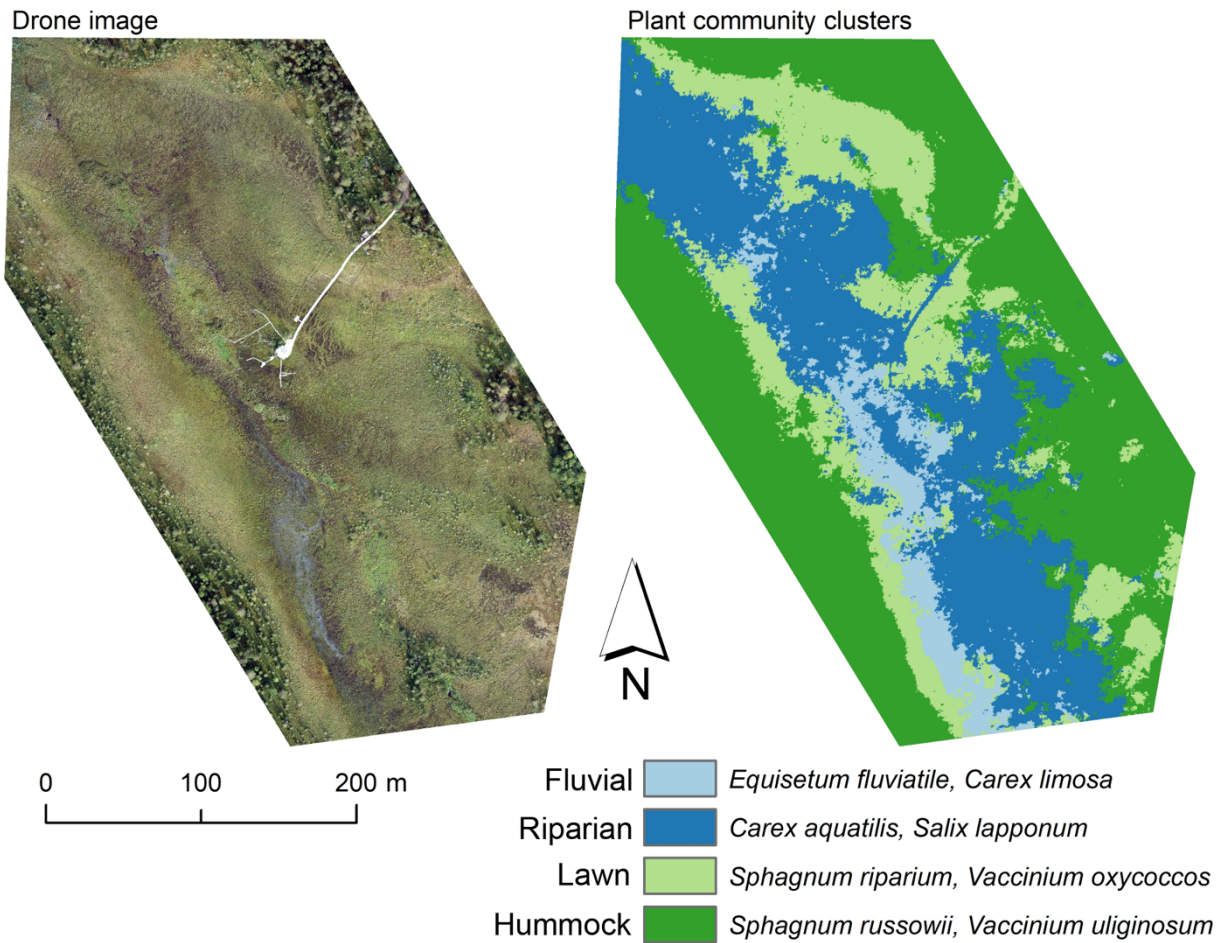
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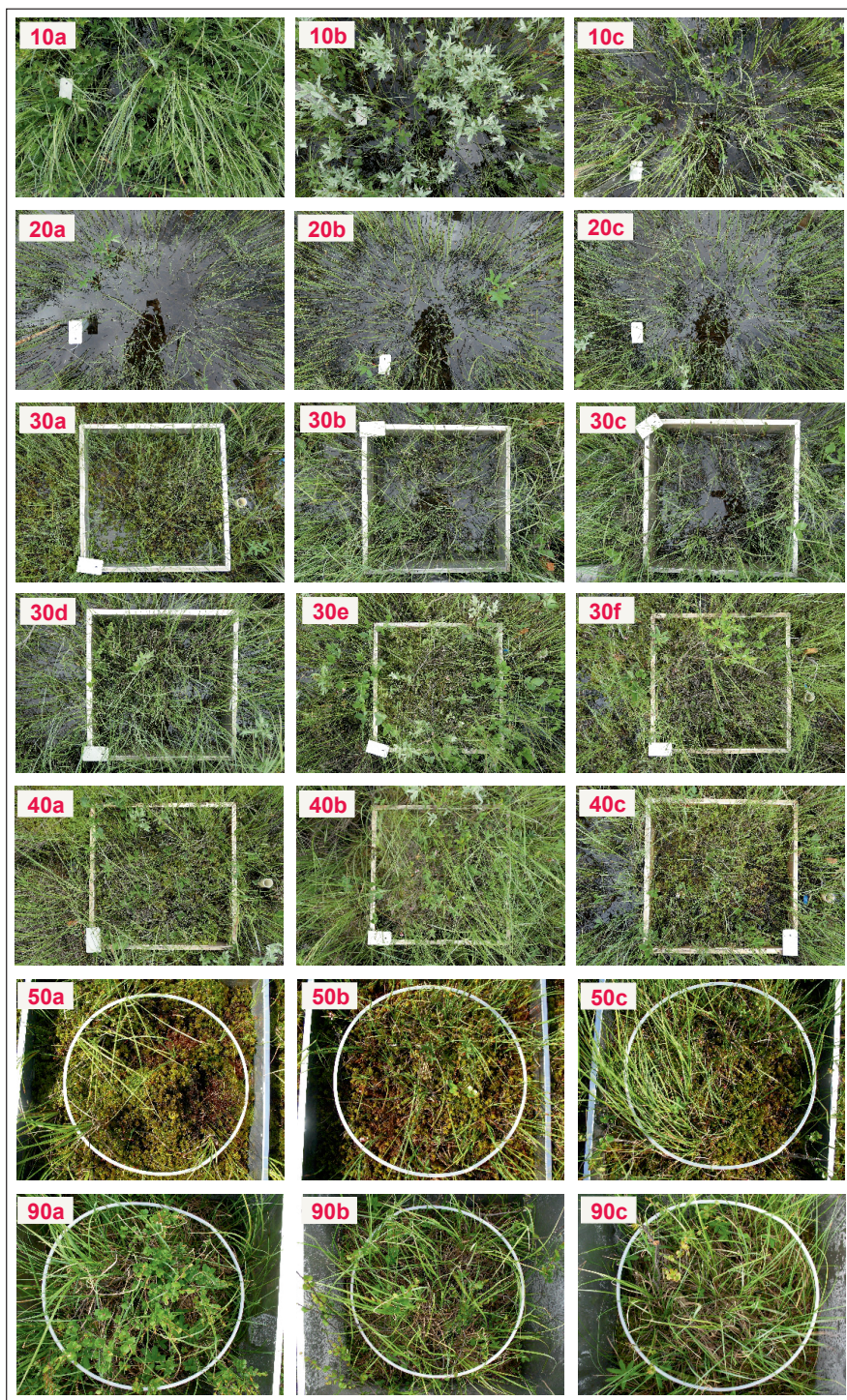
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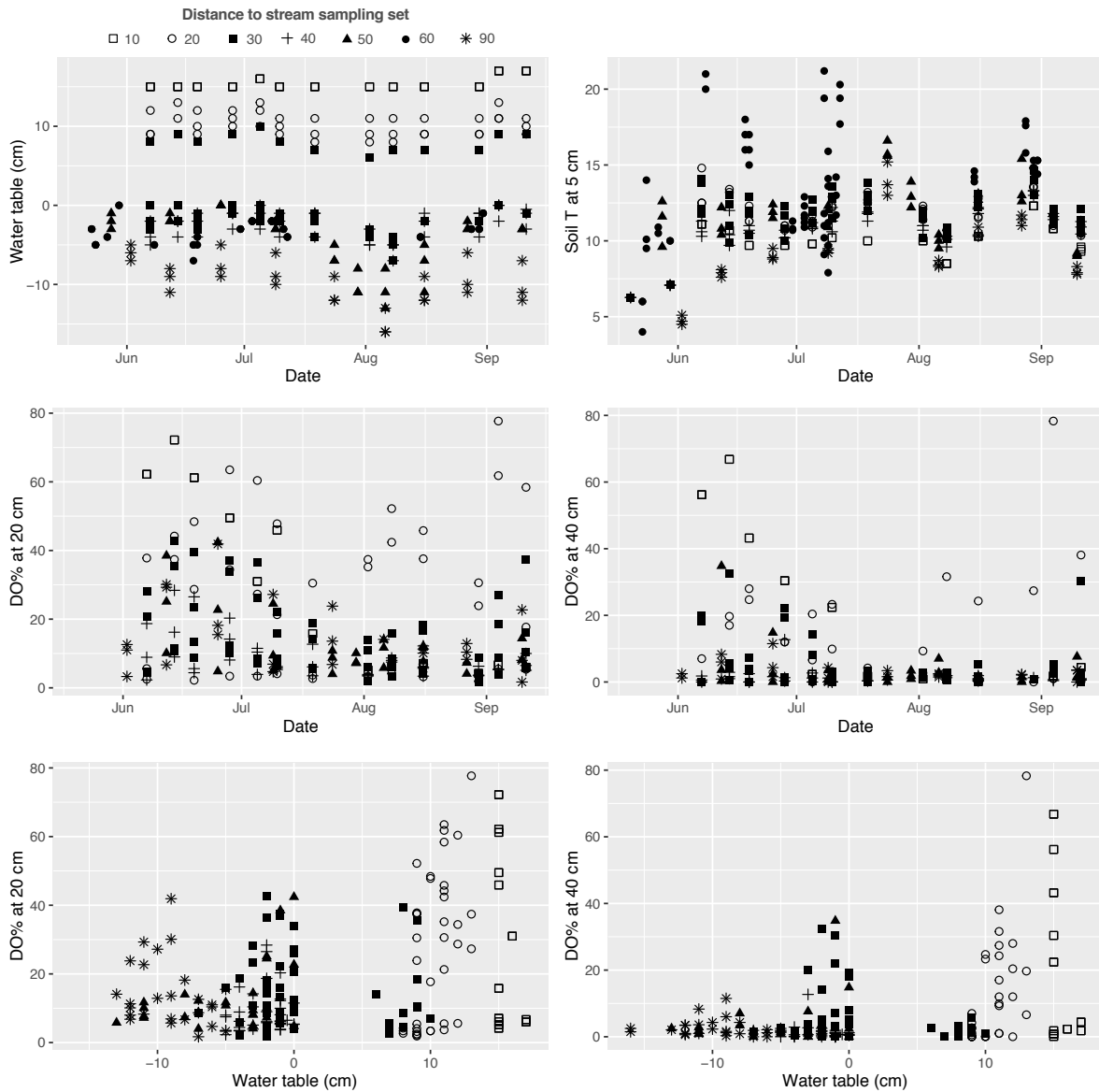
**Figure A1:** Continuous peat temperature (at 5, 10, 20, 30, 50 and 100 cm depth below the surface) in 2018 at Lompolojänkka at measuring points, (a) far from the stream, (b) intermediate distance to the stream, and (c) in the stream. Detailed description of the locations of the sample points can be found in Figure 1.



535 **Figure A2:** Drone image and a map of plant community clusters over the study area. The cluster map was produced with the multi-source remote sensing data listed in Räsänen et al. (2020). We calculated remote sensing features for the vegetation plots and vegetation patches delineated from the drone image with mean-shift segmentation in Orfeo ToolBox (Grizonnet et al. 2017). We trained a random forest classification (Breiman 2001) with the vegetation plot data and predicted the classification for the vegetation patches. Classification accuracy (random forest out-of-bag estimate) was 61 %.



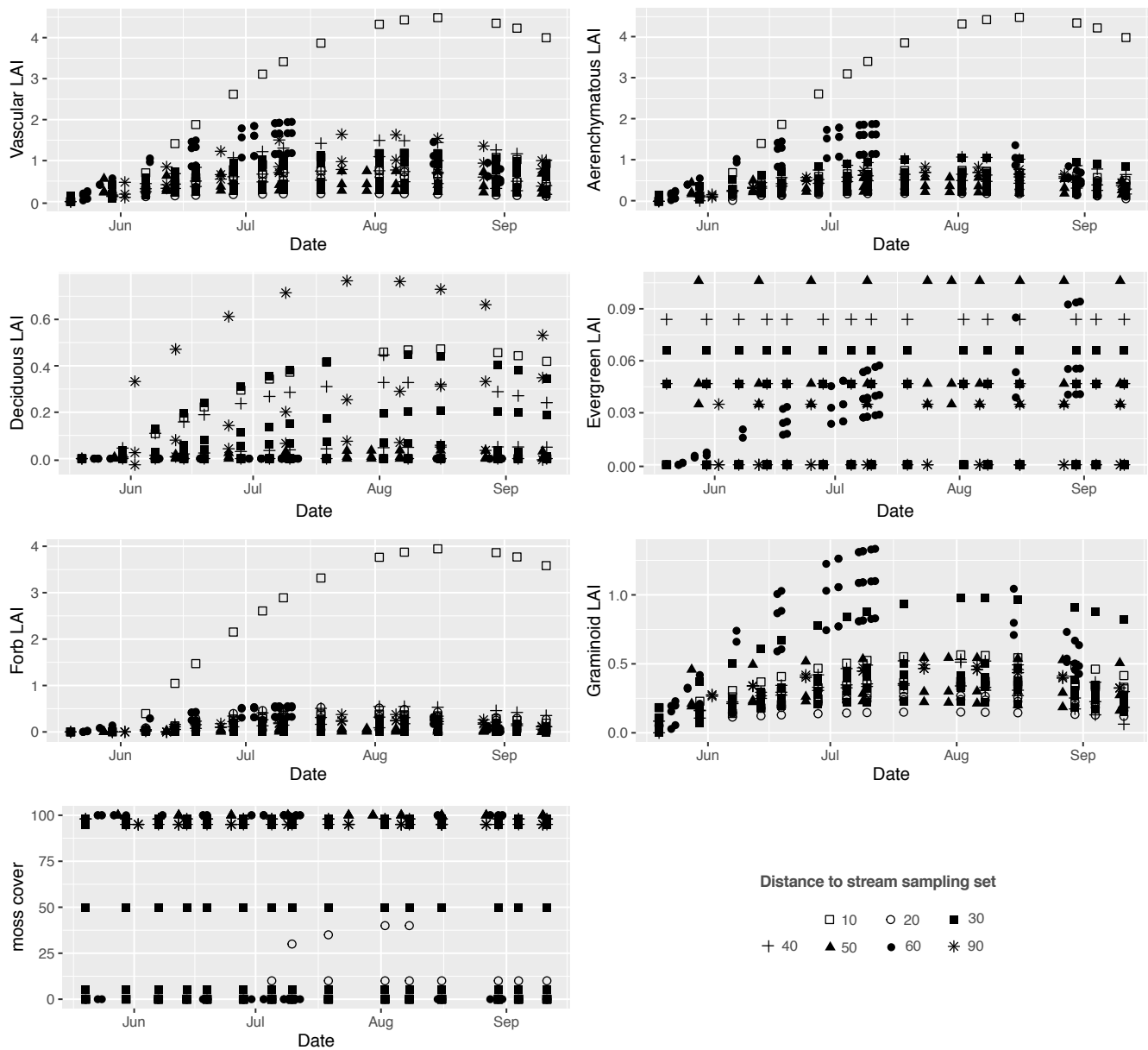
540 **Figure A3:** Photos of the methane ( $\text{CH}_4$ ) flux sample plots at Lompolojärnkä taken on 1 July 2019 ([elusters-sets\\_10](#) to 40) and 28 June 2019 ([elusters-sets\\_50](#) and 90). Photos for [eluster-set\\_60](#) were not taken.



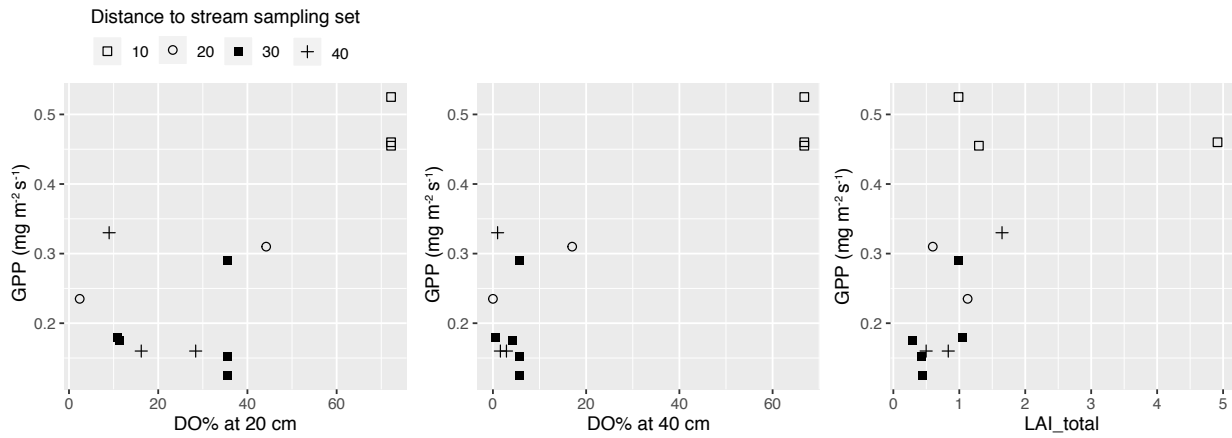
**Figure A4:** Upper two panels: temporal variations in water table, soil temperature (T) at 5 cm below the peat surface, and dissolved oxygen concentration (DO%) at 20 cm and 40 cm below the surface during the 2019 growing season at each methane (CH<sub>4</sub>) flux sample plot. Lower panel: DO% at 20 and 40 cm plotted against water table. Plots in the same cluster set are labeled using the same symbol.

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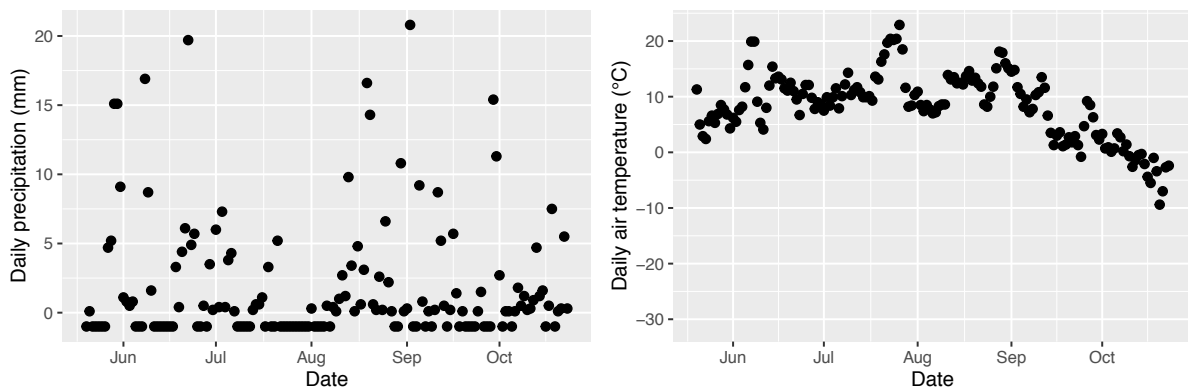


**Figure A5:** Temporal development of leaf area index (LAI) for total vascular, aerenchymatous, deciduous, evergreen, forb, and graminoid plants, and moss cover (%) in each methane (CH<sub>4</sub>) flux sample plot. Plots in the same [cluster-set](#) are labeled using the same symbol.



**Figure A6:** Gross primary production (GPP) at a photosynthetic photon flux density level of  $800 \mu\text{mol m}^{-2} \text{s}^{-1}$  in the methane ( $\text{CH}_4$ ) flux sample plots in [clusters-sets](#) 10–40 plotted against peak season (early summer) dissolved oxygen concentration (DO%) at 20 and 40 cm below the surface, and peak season (late July) total vascular leaf area index (LAI\_total). Plots in the same [cluster-set](#) are labeled using the same symbol.

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**Figure A7:** Daily precipitation and air temperature at Lompolojänkkä during summer 2019. Data were originated from the nearest weather stations; Lompolonvuoma (for temperature) and Kenttäröva (for precipitation).

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**Table A1:** Equations for calculating leaf area index (LAI) for four plant functional types (PFTs) based on %-cover (c) and height (h; cm) data.

PFT	Equation	RMSE	Adj.R <sup>2</sup>
evergreen shrub	$LAI = 0.023066 + 0.011866*c$	0.1413066	0.7488
deciduous shrub	$LAI = -0.034458 + 0.020706*c$	0.3275706	0.7261
forb	$LAI = -2.193e-02 + 1.360e-03*c*h$	0.188271	0.8877
graminoid	$LAI = 0.045542 + 0.024965*c$	0.1697018	0.7346

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**Table A2:** Indicator plant species of plant community clusters of the methane (CH<sub>4</sub>) flux sample plots derived from non-metric multidimensional scaling (NMDS) analysis.

Species	Cluster	Indicator_ value	Species	Cluster	Indicator_ value	Species	Cluster	Indicator_ value
<i>Equisetum fluviatile</i>	1	0.505	<i>Menyanthes trifoliata</i>	2	0.208	<i>Empetrum nigrum</i>	4	0.254
<i>Carex limosa</i>	1	0.355	<i>Sphagnum riparium</i>	3	0.361	<i>Sphagnum fallax</i>	4	0.189
<i>Carex canescens</i>	1	0.116	<i>Vaccinium oxycoccos</i>	3	0.284	<i>Aulacomnium palustre</i>	4	0.188
<i>Maksasammal sp.</i>	1	0.088	<i>Carex livida</i>	3	0.194	<i>Carex pauciflora</i>	4	0.143
<i>Sarmentypnum sarmentosum</i>	1	0.080	<i>Sphagnum lindbergii</i>	3	0.146	<i>Eriophorum vaginatum</i>	4	0.086
<i>Carex aquatilis</i>	2	0.429	<i>Sphagnum russowii</i>	4	0.538	<i>Picea abies</i>	4	0.076
<i>Salix lapponum</i>	2	0.380	<i>Vaccinium uliginosum</i>	4	0.536	<i>Vaccinium vitis- idaea</i>	4	0.076
<i>Salix phylicifolica</i>	2	0.278	<i>Betula nana</i>	4	0.441	<i>Dicranum sp.</i>	4	0.073
<i>Mnium sp</i>	2	0.273	<i>Sphagnum total</i>	4	0.402	<i>Sphagnum compactum</i>	4	0.061
<i>Comarum palustre</i>	2	0.250	<i>Rubus chamaemorus</i>	4	0.394	<i>Equisetum sylvaticum</i>	4	0.061
<i>Sphagnum teres</i>	2	0.234	<i>Sphagnum fuscum</i>	4	0.275			

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**Table A3:** Parameter estimates of the linear mixed-effect models describing the response of methane (CH<sub>4</sub>) fluxes to environmental variables in the (a) 2017-2019, and (b) 2019 datasets. Tostream: the distance from the plot to the stream; Tair.18: air temperature greater than the threshold value 18 °C; T5: peat temperature at 5 cm depth below the surface; DO20.40: dissolved oxygen concentration at 20 cm depth below the surface lower than the threshold value 40%; WT: water level relative to the moss surface; G\_LAI: graminoid LAI; V\_LAI: vascular LAI.

(a) marginal R <sup>2</sup> = 0.37, conditional R <sup>2</sup> = 0.99					
	Coeff.	SE	DF	t-value	P-value
<b>Fixed part</b>					
Intercept	0641	0.538	349	1.191	0.235
Tostream	0.260	0.017	243	15.673	0.000
Tair.18	0.204	0.048	349	4.270	0.000
I(Tostream^2)	-0.003	0.000	243	-13.224	0.000
<b>Random part</b>					
SD (measured year)	0.000				
SD (measured month)	1.814				
SD (plot)	1.507				
Residual SD	0.317				
<b>Variance function</b>					
Power	0.973				

(b) marginal R <sup>2</sup> = 0.42, conditional R <sup>2</sup> = 0.87					
	Coeff.	SE	DF	t-value	P-value
<b>Fixed part</b>					
Intercept	-0.936	1.930	98	-0.485	0.629
T5	0.675	0.159	88	4.252	<0.001
DO20.40	0.111	0.075	220	1.487	0.139
WT	-0.370	0.066	107	-5.607	<0.001
G_LAI	7.756	2.241	136	3.461	0.001
V_LAI	-2.398	0.745	190	-3.219	0.002
T5:DO20.40	-0.015	0.006	218	-2.348	0.020
DO20.40:WT	0.003	0.001	232	2.339	0.020
WT:V_LAI	0.135	0.046	197	2.946	0.004
<b>Random part</b>					
	Variance	SD			
measurement day	1.648	1.284			
plot	5.402	2.324			
Residual	1.967	1.402			

## Data availability

590 The data used in this study will be made available on the Figshare repository after the article is accepted for publication.

## Author contribution

AL designed the experiments with the help from TL. SG, VL and HZ contributed to the field measurements. HZ, EST, AK, AR analysed the data. HZ prepared the manuscript with contributions from all co-authors.

## Competing interests

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

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