Interacting effects of vegetation components and water level on methane dynamics in a boreal fen

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Abstract. Vegetation and hydrology are important controlling factors in peatland methane dynamics. This study aimed at investigating the role of vegetation components, sedges, dwarf shrubs, and Sphagnum mosses, in methane fluxes of a boreal fen under natural and experimental water level drawdown conditions. We measured the fluxes during growing seasons 2001–2004 using the static chamber technique in a field experiment where the role of the ecosystem components was assessed via plant removal treatments. The first year was a calibration year after which the water level drawdown and vegetation removal treatments were applied. Under natural water level conditions, plant-mediated fluxes comprised 68 %–78 % of the mean growing season flux (1.73 ± 0.17 g CH₄ m⁻² month⁻¹ from June to September), of which Sphagnum mosses and sedges accounted for one-fourth and three-fourths, respectively. The presence of dwarf shrubs, on the other hand, had a slightly attenuating effect on the fluxes. In water level drawdown conditions, the mean flux was close to zero (0.03 ± 0.03 g CH₄ m⁻² month⁻¹) and the presence and absence of the plant groups had a negligible effect. In conclusion, water level acted as a switch; only in natural water level and vegetation removal treatments were applied. Under natural water level conditions, plant-mediated fluxes comprised 68 %–78 % of the mean growing season flux (1.73 ± 0.17 g CH₄ m⁻² month⁻¹ from June to September), of which Sphagnum mosses and sedges accounted for one-fourth and three-fourths, respectively. The presence of dwarf shrubs, on the other hand, had a slightly attenuating effect on the fluxes. In water level drawdown conditions, the mean flux was close to zero (0.03 ± 0.03 g CH₄ m⁻² month⁻¹) and the presence and absence of the plant groups had a negligible effect. In conclusion, water level acted as a switch; only in natural water level conditions did vegetation regulate the net fluxes. The results are relevant for assessing the response of fen peatland fluxes to changing climatic conditions, as water level drawdown and the consequent vegetation succession are the major projected impacts of climate change on northern peatlands.

1 Introduction

Approximately one-third of all terrestrial carbon is stored in boreal and subarctic peatlands (e.g., Yu, 2012) that generally act as CO₂ sinks in current climatic conditions. However, pristine wetlands, including peatlands, marshes, and floodplains, are also the largest natural source of methane (CH₄) in the atmosphere (Ciais et al., 2014; Kirschke et al., 2013; Saunois et al., 2016). The carbon sink function of peatlands is mostly due to the slow decomposition rate resulting from waterlogged, anaerobic conditions sustained by a high water level, which simultaneously favor CH₄ production. CH₄ is the end product of anaerobic decomposition by strictly anaerobic methanogenic archaea. It is released from the peat into the atmosphere via diffusion through the peat column, ebullition or plant-mediated transport (Lai, 2009). A considerable part, from 20 % to up to 90 % (Le Mer and Roger, 2001; Pearce and Clymo, 2001; Whalen, 2005) of the CH₄ diffusing through the upper, aerobic part of the peat layer is oxidized to CO₂ by methanotrophic bacteria (MOB) before reaching the atmosphere.

Vegetation is a major factor controlling peatland CH₄ fluxes (Koelbener et al., 2010; Ström et al., 2005, 2012). Fresh root litter and exudates are important substrates for the methanogenic microbes, and a significant proportion of the CH₄ is formed from this easily available organic matter instead of from old, recalcitrant peat (Koelbener et al., 2010; Ström et al., 2012). Therefore, CH₄ fluxes have a strong, positive correlation with the CO₂ uptake (Bellisario et al., 1999; Christensen et al., 2000; Rinne et al., 2018), since
higher primary productivity leads to a higher input of substrates. Of the vegetation components, deep-rooting aerenchymatous species such as sedges (Cyperaceae) and aerenchymatous herbs are especially important (Leppälä et al., 2011; Ward et al., 2013). In sedge-dominated wetlands, most of the CH4 is released through vascular plants (Whalen and Reeburgh, 2000; Fritz et al., 2011) and stimulate CH4 oxidation (King, 1994; Popp et al., 2000). The net effect of the presence of aerenchymatous species on CH4 fluxes is positive in most cases (Bellisario et al., 1999; Greenup et al., 2000; Rinnan et al., 2003; Couwenberg and Fritz, 2012; Ward et al., 2013), although opposite results have also been reported (Roura-Carol and Freeman, 1999; Strack et al., 2006). Although the influence of the non-aerenchymatous species on the fluxes has been studied relatively little, Gray et al. (2013) showed that plant functional groups based on more complex traits than those related to aerenchyma were good proxies of CH4 flux. In open boreal peatlands, the most abundant non-aerenchymatous vascular plant functional group is dwarf shrubs, which are generally shallow rooted (Korrensalo et al., 2018a) and have a negligible CH4 transport capacity (Shannon et al., 1996; Garnet et al., 2005) compared to deep-rooting aerenchymatous species. In plant removal experiments, the presence of shrubs has been shown to decrease CH4 fluxes (Ward et al., 2013; Robroek et al., 2015). Recently, trees have been shown to transport significant amounts of CH4 from soil in certain ecosystems, but so far not in forested boreal peatlands (Covey and Megonigal, 2019). Sphagnum mosses, in turn, have an impact on CH4 oxidation as they host partly endophytic methanotrophs in the water-filled, hyaline cells of their leaves and stem (Raghoebarsing et al., 2005; Larmola et al., 2010; Putkinen et al., 2012).

Water level regulates the volume ratio of the aerobic and anaerobic peat and, consequently, the extent of the CH4 production and oxidation zones. Therefore, a positive correlation between the water level and CH4 fluxes has been reported in numerous studies (Moore and Roulet, 1993; Laine et al., 2007a; Pearson et al., 2015; Turetsky et al., 2014; Chimner et al., 2017). However, the relationship between the water level and CH4 fluxes is complex due to the vegetation–water level interaction. Because the plant communities in the wettest habitats are often associated with the sparsest vascular plant cover and lowest productivity (Waddington and Roulet, 2000; Laine et al., 2007b; Riutta et al., 2007b), less substrate for CH4 production is available in those communities. At the dry end of the water level gradient, fewer roots reach the anaerobic layer of the peat (Waddington et al., 1996; Kützbach et al., 2004). Hence, CH4 fluxes may also show a unimodal relationship to water level (Strack et al., 2004; Brown et al., 2014) or no relationship at all (Rask et al., 2002; Korrensalo et al., 2018b).

In this study, we aim to disentangle the intertwined relationships among water level, vegetation, and fen CH4 fluxes. We test the role assumed for different plant functional groups based on earlier literature and quantify how these roles are modulated by changing water level. Our objective is to quantify the contribution of the different components of fen plant community, namely sedges, dwarf shrubs, Sphagnum mosses, and the underlying peat, to the CH4 fluxes under wet and dry conditions. To achieve this, we applied removal treatments of plant functional groups under both natural and experimentally lowered water level in a factorial study design. We hypothesized that aerenchymatous plant species enhance CH4 fluxes and that this effect would be less pronounced under lowered water level as a smaller proportion of the roots would extend to the anaerobic peat layer. Further, we hypothesized Sphagnum mosses and dwarf shrubs to reduce CH4 fluxes.

2 Materials and methods

2.1 Study site

The study was carried out at Lakkasuo peatland complex, an eccentric raised bog with minerotrophic laggs situated in the southern boreal vegetation zone (Ahti et al., 1968) in southern Finland (61°47′N, 24°18′E). Annual precipitation in the region totals 710 mm, of which about a third falls as snow. The average temperatures for January and July are −8.9 and 15.3 °C, respectively (Juupajoki Hyytiälä weather station).

The study site was situated on a nutrient-poor, oligotrophic, treeless fen part of the peatland complex. Surface topography at the site is uniform, mostly lawn. The pH of the surface peat at the site was 4.9 (Juutonen et al., 2005). The field layer is dominated by sedges and dwarf shrubs. The most abundant sedge species is Carex lasiocarpa Ehrh. (percentage of cover in 2001 3.4±3.9, mean ± standard deviation of 40 inventory plots), and other common sedge species are Eriophorum vaginatum L. (0.9±1.8) and Trichophorum cespitosum (L.) Hartm. (0.5±2.4). The most abundant shrubs are the deciduous Betula nana L. (4.0±4.2) and ericaceous Andromeda polifolia L. (6.6±5.7) and Vaccinium oxycoccos L. (4.9±4.2). Note that due to the erect growth form of sedges, their percentage of cover is lower than that of shrubs, although their leaf area is higher; see Table 1 and Fig. 2. The moss layer forms a continuous carpet dominated by Sphagnum papillosum Lindh. (40.1±31.3) and the species of the S. recurvum complex (S. fallax (Klinggr.) Klinggr., S. flexuosum Dozy & Molk, and S. angustifolium (C.E.O.Jensen ex Russow) C.E.O.Jensen) (together 32.7±24.0). The vegetation inventory and variation conducted at the site are described in detail in Kokkonen et al. (2019).
Table 1. Growing season average (standard deviation) water level (WL) relative to moss surface (in centimeters), with negative values indicating water level below the surface, growing season peak LAI of sedges (LAIc) and dwarf shrubs (LAId), and projection cover of Sphagnum mosses (Spha) (unit is square meters per square meter) in different plant removal treatments in wet and WL drawdown subsites. The year 2001 was a calibration year without the WL drawdown and plant removal treatments, which were implemented in 2002. Vegetation treatments: PSCD – plots with intact vegetation, consisting of peat, Sphagnum mosses, sedges, and shrubs; PSC – plots consisting of peat, Sphagnum mosses, and sedges (shrubs removed); PS – plots consisting of peat and Sphagnum mosses (shrubs and sedges removed); P – plots consisting of bare peat (all vegetation removed).

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2.2 Experimental design

The study was carried out during four growing seasons from 2001 to 2004. The first season of the study, 2001, served as a calibration year without the water level drawdown (WLD) treatment, which was implemented in April 2002. The study site was divided into two subsites approximately 20 m apart, namely the wet and the drier WLD subsite, by surrounding the WLD subsite with a shallow ditch that lowered the water level by an average 17 ± 1.6 cm (22 ± 3.0 cm in 2002, 12 ± 3.4 cm in 2003, and 16 ± 1.9 cm in 2004). The shallow ditch was located approximately 10 m from the wet subsite and drained to a larger, old ditch.

We studied the contribution of the ecosystem components to the net CH4 fluxes in wet and dry conditions by means of plant removal treatments. At the site, we established permanent sample plots of 56 cm × 56 cm consisting of

- peat, Sphagnum mosses, sedges, and dwarf shrubs (PSCD, intact vegetation, n = 8 at the wet subsite and n = 8 at the WLD subsite)
- peat, Sphagnum mosses, and sedges (PSC, dwarf shrubs removed, n = 5 + 4)
- peat and Sphagnum mosses (PS, sedges, and shrubs removed, n = 3 + 3)
- peat (P, all vegetation removed, n = 4 + 4).

The plant removal treatment plots (PSC, PS, and P) were established April 2002. In the plant removal treatment plots vascular plants were cut with scissors to the level of the moss (PS plots) or peat (P plots) surface and their aboveground litter was removed. In the P plots the top 1.5 cm of the Sphagnum moss carpet was cut off with scissors. All emerging regrowth was clipped off once a week as necessary. Over the course of the study, progressively less clipping was needed, with hardly any in 2004. Prior to CH4 flux measurements, sedge stubble in P and PS plots was treated with paraffin wax to seal the aerenchymatous pathway of CH4.

2.3 Measurements

CH4 fluxes were measured using the closed-chamber method. A stainless steel collar (56 cm × 56 cm × 30 cm, length × width × height) was permanently inserted into each sample plot prior to the start of the study. The collars had a water groove to allow chamber placement and airtight sealing during the measurement. For the flux measurements, an aluminum chamber of 60 cm × 60 cm × 30 cm was placed on the water groove of the collar. After the chamber placement, a vent on the chamber roof that ensured pressure equilibration was sealed with a septum plug. A battery-operated fan circulated the air inside the chamber. A 40 mL air sample was drawn into a polypropylene syringe at 5, 15, 25, and 35 min after closure. The samples were stored at +4 °C before analysis, which was carried out within 36 h. Samples were analyzed with a HP-5710A gas chromatograph (GC) from 2001 to 2003 and with a HP-5890A GC in 2004. Both GCs were equipped with a 1 mL loop, 6 × 1/8” packed column (HayeSep Q in HP-5710A; Porapak Q in HP-5890A) and flame ionization detector. The carrier gas was helium with a flow rate of 30 mL min⁻¹. Column and detector temperatures were 40 and 300°C, respectively. The precision of the analysis was ±0.16 %, determined as the coefficient of variation of the replicate samples.

To relate the fluxes to prevailing environmental conditions, peat temperatures at 5, 10, 20, and 30 cm below the moss surface and water level in a perforated tube adjacent to each plot were measured during the flux measurements. Air and peat temperatures and precipitation were also continuously recorded at the weather station at the site. Green leaf area index (LAI) of each vascular plant species in each plot was determined with the method of Wilson et al. (2007) from April until November, as a product of the total number of leaves (counted monthly) and the average leaf size of marked indi-
viduals (measured every 2 weeks). Species-specific Gaussian curves (Wilson et al., 2007) were fitted to the observations to describe the continuous development of LAI throughout the season. LAI of different species was summed up to sedge, dwarf-shrub, and total LAI (LAIc, LAId, and LAIg, respectively). Moss cover at each plot was visually estimated annually.

In addition to CH4 exchange, CO2 exchange was measured at the study site. The methods and results are reported elsewhere (Riutta et al., 2007a) in more detail, but some CO2 exchange estimates are used here to study the relationship between the CO2 and CH4 fluxes. In summary, net ecosystem CO2 exchange (NEE) was measured weekly or once every 2 weeks by employing the closed-chamber technique in the same plots and during the same period as the CH4 fluxes. Measurements were carried out in both light and dark, which enabled the partitioning of the fluxes into gross photosynthesis and ecosystem respiration. We constructed nonlinear regression models for photosynthesis and respiration, with water level, temperature, and LAI as explanatory factors, separately for each vegetation treatment, to reconstruct the fluxes for the whole growing season.

### 2.4 Data analyses

CH4 flux was calculated as the linear change in CH4 concentration as a function of time by fitting a least-squares regression line. Of the 1300 measurements, < 0.5% were rejected due to clear errors, such as leakage or problems in the GC analysis, and 2% were classified as episodic fluxes.

To reconstruct seasonal (June–September) estimates for each sample plot, the fluxes measured once every 2 weeks were linearly interpolated between measurement days, and the obtained daily values were integrated. In the interpolation, rejected values and episodic fluxes were replaced with the median flux of the corresponding vegetation and water level treatment on the same measurement day. The impact of the episodic fluxes on the seasonal flux was taken into account by using the episodic values as the CH4 flux estimates of the day they were measured. The reconstructed seasonal fluxes at the wet and WLD subsites were converted to CO2 equivalent according to Myhre et al. (2013).

We used linear mixed-effect models to test the impact of the plant removal treatments and the WLD treatment on WL, LAI, and daily measured CH4 flux. First, we tested the differences in WL, LAIc, LAId, LAIg, and CH4 flux between the wet and WLD subsites before the WLD treatment was applied (year 2001) and over the years after the WLD treatment (2002–2004), with WLD treatment, year, and their interaction as potential fixed predictors. This model included only the plots with intact vegetation (PSCD). The wet subsite in 2001 was the constant against which WLD and other years were compared. Therefore, the difference between the wet and WLD treatment in the model describes the pre-treatment difference among the two subsites in the calibration year 2001, and the interaction between WLD treatment and the years 2002–2004 describes the impact of WLD after the treatment.

Second, we tested the impact of plant removal on CH4 flux over the years and the interaction of the plant removal treatments with the WLD treatment with data from the years 2002 to 2004 (no plant removal treatments in 2001). For each year separately, we fitted a model with plant removal treatments, WLD treatment, and the interaction between them as potential fixed predictors.

Third, we tested the response of CH4 flux to leaf area and environmental variables by extending the model fitted to the data of the year 2004, which had the maximum amount of time for stabilization after the treatments. In addition to plant removal and WLD treatments, potential fixed predictors were LAIc, LAId, cover of Sphagnum mosses, measured WL, temperature in the chamber, and peat temperature at the depths of 5, 10, 20, and 30 cm (T5, T10, T20, and T30) as well as the potential interactions among these parameters. Potential new predictors were sequentially added and after each addition the significance of all predictors was tested. We reported both models separately for the year 2004: one including plant removal and WLD treatments as fixed predictors for CH4 flux and another including the response of CH4 flux to leaf area and cover of plant groups and environmental variables.

In each case, a conditional F test was used to test if the full model with all fixed predictors and their interactions was significantly better (p < 0.05) than a simpler model. Plot and date were included as crossed random effects. Resulting models are reported in Table 2. The models were fitted using the function lmer of the package lme4 (Bates et al., 2015) of RStudio version 1.1.383.

### 3 Results

#### 3.1 Impact of the water level drawdown

The pre-treatment water level did not differ between the wet and WLD subsites (p = 0.174, comparison between wet and WLD treatment during the calibration year 2001) (Fig. 1a, Table 1). Following the drainage in April 2002, the water level was significantly lower in the WLD subsite (p < 0.001, interaction between WLD and the year 2002). The WLD treatment lowered the water level by approximately 17 cm, except in July and August 2003 when a severe drought lowered the water level below the ditch, resulting in similar water levels at wet and WLD subsites. At the wet subsite, the water level during the years 2001 and 2004 was similar to the long-term average of the site, approximately 5 to 10 cm below the moss surface (Table 1) (Laine et al., 2004). During July and August 2002 and 2003, however, the water level was lower than the long-term average. More information on the weather conditions during the study is given in Riutta et al. (2007b).
Table 2. Parameter estimates of the linear mixed-effect model describing (a) the differences, water level (WL, cm), total, sedge, and dwarf-shrub leaf area index (LAI_T, LAI_C, and LAI_D), and CH4 flux between wet (WLD0) and water level drawdown (WLD1) subsites and years before (2001) and after (2002–2004) the WLD treatment in plots without vegetation removal, (b) the differences in CH4 flux between the vegetation removal treatments in the years 2002–2004 and (c) the response of CH4 flux in the year 2004 to leaf area and environmental variables. Vegetation treatments: PSCD – intact vegetation; PSC – plots consisting of peat, Sphagnum mosses, and sedges (shrubs removed); PS – plots consisting of peat and Sphagnum (sedges and shrubs removed); P – plots consisting of bare peat (all vegetation removed).

(a) Fixed part

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Random part

| SD (measurement day) | 7.5 | 0.28 | 0.22 | 0.06 | 1.2 |
| SD (plot code) | 2.0 | 0.25 | 0.21 | 0.09 | 0.5 |
| Residual SD | 2.9 | 0.19 | 0.19 | 0.05 | 1.1 |

(b) Fixed part

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Random part

| SD (measurement day) | 1.6 | 0.7 | 0.9 |
| SD (plot code) | 3.0 | 0.7 | 0.4 |
| Residual SD | 3.8 | 1.3 | 0.9 |
Prior to the drainage, vegetation composition in the plots with intact vegetation (PSCD) was similar at both subsites (Table 1, Fig. 1b). In the mixed-effect model, LAI_C, LAI_D, and LAI_T did not differ between wet and WLD subsites in the year 2001 (p values 0.996, 0.656, and 0.878, respectively). In 2001 the peak season average LAI_T was approximately 1.0 m^2 m^{-2}, of which sedges composed 70%. The mean Sphagnum cover was 80%. By the third year since WLD, 2004 LAI_C had decreased (p < 0.001) and LAI_D increased (p < 0.001) at the WLD subsite, resulting in an overall decrease in LAI_T (p = 0.007) (Table 1, Fig. 1b).

In the PSCD plots, the pre-treatment CH_4 fluxes did not differ between the wet and WLD subsites (p = 0.654) (Fig. 1c). After the treatment, in 2002–2004, fluxes were significantly lower in the WLD than at the wet subsite (p < 0.001 for all years). During the 3-year WLD treatment, the mean flux was approximately 51 and 7.0 mg CH_4 m^{-2} d^{-1} at the wet and WLD subsites, respectively. Converted to CO_2 equivalents, the seasonal reconstructed fluxes at the wet and WLD subsites in 2002–2004 were 236 and 32 g CO_2 eq. m^{-2} per growing season, respectively.

3.2 Impact of the plant removal treatments

Plant removal treatments did not lead to major changes in vegetation composition beyond the clipped target groups. Vascular plant removal did not affect the Sphagnum moss cover, and the removal of dwarf shrubs did not change the LAI of sedges. LAI_C was similar in PSC and PSCD plots (data for 2004 shown in Table 1) during all years at the wet subsite and during 2003 and 2004 at the WLD subsite (all p values > 0.05). LAI_C was higher in the PSC plots than in the PSCD plots at the WLD subsite in 2002 (p = 0.016).

During the first season of the removal treatments (2002) at the wet subsite, CH_4 fluxes were higher in the plant removal plots (P, PS, and PSC) than in the intact plots (PSCD), in some cases almost triple (p < 0.05 for all treatments, Fig. 2a–c). The fluxes in the plant removal treatment plots also showed a stronger seasonal pattern and larger spatial variation. After the first year of removal treatments, the fluxes of the P, PS, and PSC plots decreased, and in 2003 P plots had a similar CH_4 flux to the intact plots (p = 0.908), while PS and PSC plots still had a higher flux than PSCD plots (p = 0.033 and p = 0.005, respectively).

By the third year of the plant removal treatments (2004), the fluxes in all treatments showed a seasonal pattern sim-
ilar to that of the intact plots. Bare peat plots had lower fluxes than the intact PSCD plots ($p < 0.001$). Fluxes of the PSC plots (shrubs removed) were marginally significantly higher ($p = 0.060$) than those of the PSCD plots (shrubs present). In WLD conditions, the fluxes in the plant removal plots (P, PS, and PSC) were mostly lower than the fluxes in the intact PSCD plots during all three vegetation treatment years (Fig. 2d–f), but the differences were not significant (Table 2b). WLD and plant removal treatments had a significant interaction: in 2004 WLD lowered the fluxes more in PSCD plots than in the P plots and more in PSC plots than in the P and PS plots ($p < 0.05$ for the interaction terms). Seasonal fluxes visualize the patterns tested with the nonlinear mixed-effect models: at the WLD subsite fluxes were lower than at the wet subsite in all plant removal treatments (Fig. 3b). In wet conditions, the seasonal flux of the P and PS plots was lower than that of the PSCD and PSC plots in which vascular plants were present (Fig. 3a). Taking the fluxes from bare peat plots as a baseline, the presence of vegetation enhanced the fluxes. Compared with the situation of sedges and Sphagna present (PSC), the presence of shrubs (PSCD) seemed to slightly attenuate the fluxes (Fig. 3b, c). In WLD conditions, the differences between plant removal treatments were negligible. The differences between the plant removal treatments can be used as an estimate of the contribution of each plant group to the total flux, although due to the propagation of the errors, uncertainty in these estimates is large. In normal hydrological conditions, plant-mediated flux accounted for 68% ± 23% (comparison of P and PSCD plots) or 78% ± 17% (comparison of P and PSC plots) of the total growing season flux, of which Sphagnum mosses and sedges accounted for approximately one-fourth and three-fourths, respectively (Fig. 3c).

The seasonal CH$_4$ fluxes displayed a clear positive, exponential relationship with the seasonal net CO$_2$ flux (Fig. 4). The relationship was similar among the plant removal treatments in wet and dry conditions. However, the plots with intact vegetation (PSCD) were an exception; they had lower CH$_4$ fluxes than could have been expected based on their net CO$_2$ flux, pointing towards the potential suppressing effect of shrubs on CH$_4$ emissions.

### 3.3 Response of CH$_4$ flux to environmental variables and interaction with leaf area

The best predictors of the CH$_4$ flux in the extended model for the year 2004 were the categorical WLD treatment (which was a better predictor than the measured WL), $T_{20}$ (best out of the measured temperatures), and LAI$_C$ (which was a better predictor than the categorical vegetation removal treatment). The abundance of the other plant functional groups, LAI$_D$, or Sphagna cover did not have a significant effect on the fluxes. CH$_4$ flux was increased by LAI$_C$ and $T_{20}$ in wet conditions (Table 2c). In the WLD conditions, however, neither LAI$_C$ nor $T_{20}$ had any impact on the fluxes (coefficient estimates for LAI$_C$*WLD1 and $T_{20}^*$WLD1 cancel out the coefficient estimates for LAI$_C$ and $T_{20}$ in wet conditions; Table 2c). The positive coefficient of the WLD treatment seemingly indicated a larger flux at the WLD treatment site compared with the wet site, when LAI$_C$ and $T_{20}$ both equal zero; however, the measured minimum $T_{20}$ during the growing season in 2004 was 6.1°C, and the model was not intended for any extrapolation. The predicted CH$_4$ flux in the WLD treatment was similar to or lower than the flux in the wet treatment in the observed $T_{20}$ and LAI$_C$ range.

### 4 Discussion

#### 4.1 The effect of the plant types and substrate on the CH$_4$ fluxes in natural water level conditions

In line with previous studies, the plant removal treatments of this study indicated that sedges were the most important plant group in regulating CH$_4$ fluxes. In other sedge-dominated sites, plant-mediated flux has accounted for 75% to 97% of the total flux (Schimel, 1995; Kelker and Chanton, 1997; Ström et al., 2005; Sun et al., 2012; Noyce et al., 2014) and plant removal experiments have shown that of different plant functional types, removal of graminoids causes the largest decrease in CH$_4$ production and flux (Ward et al., 2013; Robroek et al., 2015). Compared with the bare peat surfaces, the presence of Sphagnum mosses seemed to have a slight, although not statistically significant, enhancing effect on the CH$_4$ fluxes, similar to the results of Roura-Carol and Freeman (1998), who found the presence of mosses to have a slightly attenuating effect on the fluxes, while Greenup et al. (2000) did not find significant differences in fluxes after Sphagnum removal. Based on this, the CH$_4$ oxidation by the loosely symbiotic methanotrophs within Sphagnum mosses (Raghoebarsing et al., 2005; Larmola et al., 2010; Putkinnen et al., 2012) seems to play a minor role in CH$_4$ dynamics at our site.

Similarly to Ward et al. (2013), we found that the presence of shrubs seemed to have a slightly attenuating effect on the fluxes under natural water level conditions. Robroek et al. (2015) made a similar finding with potential CH$_4$ production. In contrast, an aerenchymatous shrub, Myrica gale, supported similar potential CH$_4$ production to a sedge, Carex aquatilis, and did not suppress CH$_4$ flux (Strack et al., 2017). Furthermore, in line with the attenuating effect of shrubs, the CH$_4$ flux : NEE ratio was lower in the plots with intact vegetation (PSCD, shrubs present) than in the other vegetation treatments. Mechanisms behind that might relate to the impact of shrubs on soil chemistry, microbial community, or the biomass allocation of sedges. Shrub litter has higher lignin and leaf dry matter content than sedges, which are both related to lower methanogenesis (Yavitt et al., 2019). Shrub removal has been observed to result in higher dissolved organic C and N and lower C:N ratio (Ward et al., 2013) as
Figure 2. Difference of the measured CH$_4$ fluxes in plots with plant removal treatments and the mean flux in the plots with intact vegetation on each measurement day at the control subsite (a, b, c) and water level drawdown subsite (d, e, f). Positive values indicate that fluxes in the plant removal treatment plots are higher than in the intact plots. Units on the x axis give the day of year. Note the difference scales of the y axes in the upper and lower panels. Error bars are standard errors of the mean. Vegetation treatments: PSC – plots consisting of peat, Sphagnum mosses, and sedges (shrubs removed); PS – plots consisting of peat and Sphagnum (sedges and shrubs removed); P – plots consisting of bare peat (all vegetation removed). Intact plots consisted of peat, Sphagnum mosses, sedges, and shrubs. Removal treatments were established in 2002.

Figure 3. Seasonal (June–September) CH$_4$ flux (mean ± 1 standard error) at wet and water level (WLD) drawdown subsites (a) in plots with intact vegetation (PSCD) during the 4 study years (2001 was a calibration year before the implementation of the WLD treatment), (b) in different plant removal treatments plots in 2004, and (c) by each plant group, the contribution of which to the total flux in 2004 was estimated from differences between the different plant removal treatments. Letters above bars denote differences among treatments, where bars with no letter in common are significantly different based on a mixed-effect models presented in Table 2 (a, b) and based on two-way ANOVA test with Tukey pairwise comparisons (c). Plant removal treatments in (b): PSCD – plots with intact vegetation, consisting of peat, Sphagnum mosses, sedges, and shrubs; PSC – plots consisting of peat, Sphagnum mosses, and sedges (shrubs removed); PS – plots consisting of peat and Sphagnum mosses (sedges and shrubs removed); P – plots consisting of bare peat (all vegetation removed). Plant groups in (c): P – bare peat; S – Sphagnum mosses; C – sedges; D – dwarf shrubs.
Figure 4. (a) The relationship between the net ecosystem CO$_2$ uptake (NEE) and CH$_4$ flux during the growing season of 2004 described with an exponential model and (b) the residuals of the model, in the different plant removal treatments at wet (solid symbols) and water level drawdown (open symbols) subsites. Vegetation treatments: PSCD – plots with intact vegetation, consisting of peat, Sphagnum mosses, sedges, and shrubs; PSC – plots consisting of peat, Sphagnum mosses, and sedges (shrubs removed); PS – plots consisting of peat and Sphagnum mosses (shrubs and sedges removed); P – plots consisting of bare peat (all vegetation removed). NEE is positive when the fen is a net sink of atmospheric CO$_2$. Methane flux is positive when the fen is a source of CH$_4$ to the atmosphere.

well as higher fungal biomass (Robroek et al., 2015). A study on the competitive ability and biomass allocation of a wetland grass, Molinia caerulea, revealed that M. caerulea allocated more biomass to the roots when it did not face competition by shrubs (Aerts et al., 1991). Similarly, in our study, sedges in the plots where shrubs were removed may have allocated more biomass to the roots than the sedges growing in the sedge and shrub mixture. As a result, methanogenic microbes may have benefited from the higher substrate availability in the shrub removal plots (PSC). CH$_4$ production has a negative relationship and CH$_4$ oxidation has a positive relationship with the concentration of certain woody lignin compounds in peat pore water (Yavitt et al., 2000). In our study, this may be the reason behind the lower fluxes in the presence of the arboreals. The results concerning the attenuating effect of shrubs on CH$_4$ fluxes are, however, only indicative and further process-orientated research is needed.

4.2 Delay in the plant removal treatment effect

We observed a considerable disturbance in the fluxes following the plant removal treatments. In other clipping studies in which the shoots were cut above the water level, clipping increased the CH$_4$ flux during the first growing season after clipping (Schimel, 1995), had no effect (Kelker and Chanton, 1997; Greenup et al., 2000), or decreased the flux (Waddington et al., 1996; Rinnan et al., 2003). Thus, we assumed that the higher fluxes at the clipped plots during the first 2 years after the vegetation removal treatments were mainly caused by treatment artifacts. The removal of the aboveground parts of vascular plants led to the gradual death of the belowground parts, creation of an unnatural amount of new root necromass, and, thereby, a peak in the amount of available substrate. Methanogenesis at the study site may have been substrate limited (Bergman et al., 1998; Rinne et al., 2007), which could explain the initially high fluxes in the plant removal plots. The mass loss of Carex roots and rhizomes is only 10% to 45% during the first 12 months of decomposition, although the litter quality deteriorates (Scheffer and Aerts, 2000). However, after 2 years the mass loss can be as much as 75% of the original mass (Thormann et al., 2001), which gives more confidence in the results of the third year of the plant removal treatments. Thus, we used the third year of the plant removal treatments to quantify the contribution of the vegetation components to the fluxes and the response of fluxes to environmental conditions. King et al. (1998) likewise reported the effects of the plant removal 2 years after the treatment began. Shrub litter, especially belowground litter, decomposes slower than sedge litter (Moore et al., 2007), due to the high lignin content (Yavitt et al., 2019). On the other hand, the majority of dwarf shrub roots grow in the uppermost 20 cm peat layer, while sedge roots extend deeper (Korrensalo et al., 2018a; Mäkiranta et al., 2018), causing a larger proportion of dwarf shrub roots to decompose in oxic conditions, thus counteracting the differences in litter quality. Even 2 years after the start of the vegetation removal treatments, some shrub roots still probably remained. However, they were mostly located above the CH$_4$ production zone.

4.3 Water level regulates the role of the vegetation

Experimental water level drawdown has been used to mimic climate change impact on northern peatland CH$_4$ fluxes in the mesocosm (Freeman et al., 1992; Blodau et al., 2004; Dinsmore et al., 2009) and in the field studies ranging from bogs to rich fens (Laine et al., 2007a; Strack and Waddington, 2007; Turetsky et al., 2008; Ballantyne et al., 2014; Mu-
Our results, all these studies report some level of decrease in CH$_4$ flux due to WLD ranging from 3 to $\sim$20 cm. Together with temperature and vegetation, water level is a major regulator of CH$_4$ flux (Lai, 2009; Turetsky et al., 2014). However, the mechanistic understanding of this process is still limited. While Strack et al. (2004) found only small differences in the CH$_4$ production and consumption potentials between control and WLD sites, and thus attributed the decrease in fluxes mainly to the change in the volume ratio of the anaerobic and aerobic zones, Yrjälä et al. (2011) and Peltoniemi et al. (2016) found that WLD had a stronger impact on emissions through decreasing CH$_4$ production than through increasing oxidation.

In this study, the presence or absence of the plant types or LAI$_4$ had no effect on the CH$_4$ flux in the WLD conditions. This supports the findings by Waddington et al. (1996) as well as Strack et al. (2006) that the impact of the vegetation on the fluxes is strongly dependent on the water level conditions. CH$_4$ flux also responded to peat temperature only in wet conditions. A similar result with water level and temperature response has been previously reported by Moosavi et al. (1996). Our results showed that water level acts as a switch; it turns CH$_4$ flux on and off, after which temperature and vegetation regulate the flux magnitude. This result is further emphasized by the response model, where WLD treatment including change in the ecosystem following a new WT regime rather than seasonally varying WL was a better predictor for CH$_4$ fluxes. In conclusion, vegetation is a major controlling factor of the peatland CH$_4$ dynamics, but only in wet conditions.

5 Conclusions

Vegetation, sedges in particular, regulates the level of fen CH$_4$ fluxes in normal hydrological conditions, but this vegetation control is strongly dependent on the water level regime. In water level drawdown conditions, CH$_4$ fluxes are significantly lowered, practically to zero, and vegetation composition has no influence on the fluxes. The results are relevant for assessing the response of fen peatlands to changing climatic conditions, as water level drawdown and the consequent vegetation changes are the major projected impacts of climate change on northern peatlands.

Data availability. The data associated with the paper are published in the PANGAEA repository (https://doi.pangaea.de/10.1594/PANGAEA.911742, Riutta et al., 2020a, and https://doi.pangaea.de/10.1594/PANGAEA.911740, Riutta et al., 2020b).

Author contributions. The study was designed and the field experiment established by EST, TR, and JL. Fieldwork, flux calculation, and flux reconstruction was conducted by TR. Statistical analysis was done by AK. TR wrote the first version of the manuscript, which was further processed by all other authors.

Competing interests. The authors declare that they have no conflict of interest.

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