

1 **A full year of continuous net soil and ditch CO<sub>2</sub>, CH<sub>4</sub>, N<sub>2</sub>O  
2 fluxes, soil hydrology and meteorology for a drained fen in  
3 Denmark**

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11 **Abstract.** We here present a detailed dataset of automated greenhouse gas (GHG) net soil and ditch fluxes of  
12 carbon dioxide (CO<sub>2</sub>), methane (CH<sub>4</sub>), and nitrous oxide (N<sub>2</sub>O) from a drained fen in Denmark covering a full  
13 year. The dataset resolves small scale spatial and hourly-daily-seasonal dynamics of GHG soil fluxes. The GHG  
14 flux dataset is accompanied by simultaneous time series of soil temperature and moisture, as well as  
15 groundwater table depth and covers spatiotemporal gradients in soil hydrological and climatic variability. The  
16 GHG fluxes of CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O were measured simultaneously by a high-precision cavity ring down laser  
17 spectrometer connected with a novel automated GHG system platform called SkyLine2D (Earthbound Scientific  
18 Ltd., UK) that allowed up to 27 individual chamber measurement points along a 24 meter transect. In total  
19 47.483 chamber measurements were completed and after quality control 44.631 CO<sub>2</sub> fluxes, 44.099 N<sub>2</sub>O and  
20 42.515 CH<sub>4</sub> fluxes remained.

21 The average ( $\pm$ SE) net soil CO<sub>2</sub> efflux observed at the site ( $2.6 \pm 0.02 \text{ } \mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$  or  $35 \pm 0.3 \text{ tCO}_2 \text{ ha}^{-1} \text{ y}^{-1}$ )  
22 aligns with findings from similar drained fens in northern Europe. However, this transect average masks  
23 substantial spatial variability and highlights the role of episodic emission bursts related to hydrological  
24 variability. The organic soil at the site was a larger net source of N<sub>2</sub>O ( $8.9 \pm 0.1 \text{ nmol N}_2\text{O m}^{-2} \text{ s}^{-1}$  or  $123 \pm 1.4 \text{ kg N}_2\text{O m}^{-2} \text{ ha}^{-1} \text{ y}^{-1}$ ) to the atmosphere compared to other temperate drained organic grassland soils in northern  
25 Europe. The soil N<sub>2</sub>O emissions were similarly variable in space as soil CO<sub>2</sub> effluxes, but were more dynamic in  
26 time, where increasing groundwater table depth in response to precipitation during warmer seasons led to  
27 emission bursts of soil N<sub>2</sub>O emissions that dominated the annual net budget of soil N<sub>2</sub>O and decreased to near-  
28 zero fluxes in drier warmer periods. Net soil CH<sub>4</sub> fluxes were near-zero and the site overall acted as a smaller  
29 net source ( $0.18 \pm 0.06 \text{ nmol CH}_4 \text{ m}^{-2} \text{ s}^{-1}$  or  $0.91 \pm 0.3 \text{ kg CH}_4 \text{ ha}^{-1} \text{ y}^{-1}$ ) compared to other drained organic  
30 grassland soils, although net uptake of atmospheric CH<sub>4</sub> was observed as well especially in drier conditions.

32 Diurnal and seasonal patterns of net soil CO<sub>2</sub> and N<sub>2</sub>O emissions align with variations of soil temperature, but  
33 no clear patterns were observed for net soil CH<sub>4</sub> uptake or emission. Compared to soil GHG fluxes, the ditch  
34 was a smaller net source of CO<sub>2</sub> ( $0.94 \pm 0.05 \text{ } \mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$  or  $1.3 \pm 0.7 \text{ tCO}_2 \text{ ha}^{-1} \text{ y}^{-1}$ ) and N<sub>2</sub>O ( $0.35 \pm 0.03 \text{ nmol N}_2\text{O m}^{-2} \text{ s}^{-1}$  or  $4.9 \pm 0.4 \text{ kg N}_2\text{O ha}^{-1} \text{ y}^{-1}$ ) to the atmosphere. The ditch was also a net source of CH<sub>4</sub> ( $161 \pm 13 \text{ nmol CH}_4 \text{ m}^{-2} \text{ s}^{-1}$  or  $812 \pm 66 \text{ kg CH}_4 \text{ ha}^{-1} \text{ y}^{-1}$ ) average of diffusive and ebullition fluxes) to the atmosphere and  
35 annual cumulative emissions were more than two orders of magnitude larger than net the soil CH<sub>4</sub> emissions,  
36  
37

38 confirming earlier findings that ditches can be CH<sub>4</sub> emission hotspots, where the ditch CH<sub>4</sub> is emitted in bursts  
39 with little seasonal variability, including emissions as ebullitions.

40 The data set (<https://doi.org/10.60612/DATADK/BZQ8JE>) is well suited for testing and developing  
41 biogeochemical models, with emphasis on the soil thermal-hydrology interactions with the peat C and N cycles.

42 **1 Introduction**

43 Understanding the climate feedbacks of temperate drained and rewetted wetlands require robust observational  
44 datasets of net fluxes, e.g. whether the rewetted peatlands act as net sources or sinks of greenhouse gases  
45 (GHG). This necessitates being able to capture spatial and temporal variability from these systems. Flux data  
46 covering all three major GHGs are rare for temperate peatlands, and despite growing efforts to quantify GHG  
47 fluxes from drained peatlands, existing datasets often suffer from limited temporal resolution, short monitoring  
48 periods, or a lack of concurrent hydrological and meteorological data. Many studies rely on chamber-based  
49 measurements or short-term campaigns that fail to capture seasonal dynamics and extreme events. Moreover,  
50 current datasets typically offer either high temporal resolution (e.g., eddy covariance or automatic chambers)  
51 with poor spatial coverage, or manual measurements with good spatial resolution but very low temporal  
52 frequency. In turn this hampers the ability to model and forecast GHG fluxes, and hence climatic feedbacks, in  
53 these systems under land use and climatic changes.

54 However, automated GHG closed chamber flux measurements from ecosystems are becoming increasingly  
55 common, also in peatland research (Anthony and Silver 2023; Boonman et al. 2024) as equipment costs  
56 decrease and awareness grows about the importance of resolving temporal variability of GHG fluxes to better  
57 understand soil biogeochemical processes and soil-climate feedback. But high-frequency data of GHG fluxes  
58 are still scarce for peatlands and spatial variability of fluxes is rarely represented as well due to limited number  
59 of spatial replicates. Thus, most automated chamber systems are setup around a multiplexer control unit linking  
60 multiple chambers with one or more GHG analysers. State-of-the-art automatic chamber systems, like the LI-  
61 8250 Automated Gas Flux System (LiCOR, USA) or the eosAC-LT/LO (Eosense Inc. Canada), i.e. allow for a  
62 standard number of 8 or 16 chambers, respectively, that can be upgraded to 36 chambers with additional  
63 manifolds. Such large replicate chambers allow for improved characterization of spatial variation or treatment  
64 effects coupled with temporal variations, but are costly to establish.

65 Additionally, the introduction of automated chamber systems raises the need for improved data handling and  
66 flux calculation tailored to handle a wide range of flux magnitudes and chamber behaviour or design (Kroon et  
67 al. 2008; Pihlatie et al. 2013). Recent examples of novel flux calculation software are based on publicly  
68 available R codes and include goFlux (Rheault et al. 2024), HMR (Pedersen et al. 2010; Pullens et al. 2023) and  
69 fluxfinder (Wilson et al. 2024). Furthermore, unsupervised automated chamber flux measurements increases the  
70 likelihood of misinterpretation of fluxes, such as overestimated night-time fluxes due to atmospheric  
71 stratification that disturbs the steady-state diffusion gradient between soil and the atmosphere (Brændholt et al.  
72 2017) or leaky chambers that disturb chamber headspace concentrations. This is a significant challenge of  
73 automated chamber systems producing thousands of data points, where manual control of each data point may  
74 not be practical or feasible calling for automated and objective quality control such as used with the eddy  
75 covariance methodology.

76 We here present a dataset that addresses the abovementioned limitations by uniquely combining high-frequency,  
77 continuous measurements of net soil fluxes of carbon dioxide (CO<sub>2</sub>), methane (CH<sub>4</sub>) and nitrous oxide (N<sub>2</sub>O)  
78 with detailed hydrological and meteorological variables. The GHG fluxes were measured with an automated  
79 GHG chamber system over 12 months resolving spatiotemporal patterns of GHG fluxes including 27 individual  
80 collars (26 on organic soil and 1 in a ditch) over a 24 m transect on a temperate drained fen peatland. Integrated

81 quality control, flagging of erroneous or uncertain flux measurements enabled objective filtering of poor quality  
82 data on the entire dataset. This comprehensive spatiotemporal coverage enables robust calibration and validation  
83 of biogeochemical and hydrological models, particularly those aiming to simulate the complex interactions  
84 between water table dynamics, soil processes, and GHG emissions in managed peatland systems.

85 Considering the critical need for obtaining high-quality data on soil GHG fluxes from natural and restored  
86 peatlands in Europe and globally, our dataset marks an important contribution to this endeavour as it addresses  
87 current data shortcomings for Danish and European peatlands by providing detailed data on temporal and spatial  
88 patterns of GHG fluxes from organic soils and drainage ditches together with environmental drivers of soil  
89 hydrology and temperature, organic soil properties and groundwater geochemistry. We publish this data with the  
90 aim of it being used by the scientific community for both experimentalists to test hypothesis of how GHG  
91 dynamics are related to hydrology, soil, geochemistry and climate, as well as for the modelers to test and  
92 develop biogeochemical models for peat lands.

## 93 **2 Materials and Methods**

### 94 **2.1 Site description and the SkyLine2D system**

95 The field site, Vejrumbro (N 56.43819 E 9.54527 (WGS 84)), is located in Central Jutland, in Denmark near the  
96 city of Viborg (Fig. 1) with a mean annual temperature of 8.3°C and annual precipitation of 675 mm for the  
97 period 1991–2020 (measured 6 km away at Aarhus University Viborg Meteorological Station in Foulum  
98 (Jørgensen et al. 2023)). It is situated in the Nørre Å valley and is characterized as a riparian fen peat soil (Reza  
99 Mashhadi et al. 2024). The riparian fen developed in a former glacial river valley with flat topography gently  
100 sloping (<2.5 meters over 300 meters) towards the Nørre Å that forms the central river in this area (Fig. S1).  
101 The site was drained in 1950 with ditches and tile drains for cultivation but has primarily served as grassland in  
102 recent decades due to the wet conditions (Nielsen et al. 2024). Since 2018, Vejrumbro has been a living lab for  
103 agroecological research managed by the Department of Agroecology at Aarhus University. From 2018, the site  
104 had a passive rewetting strategy by terminating maintenance of the open ditches. During 2022, the main ditches  
105 were gradually blocked.

#### 106 2.1.1 Site preparation and disturbance

107 Initially, we chose to perform the flux measurements without aboveground plants as the small chamber  
108 dimensions (height of 20 cm) prohibited inclusion of these in the chamber as the plants typically reach over 100  
109 cm in height at this site. The strategy was therefore to focus on measuring net soil GHG fluxes, where we  
110 assume the contribution of gases are derived from heterotrophic respiration of older peat C/N, root exudated  
111 C/N from adjacent plants, dissolved N in groundwater and belowground autotrophic respiration (CO<sub>2</sub>) from  
112 roots inhabiting the peat below the collars. We are aware that omitting plants prohibit a full evaluation of the net  
113 ecosystem exchange of GHG and hence its net climate impact, as the aboveground plants represent a net sink of  
114 atmospheric CO<sub>2</sub> and also can increase the emission of CH<sub>4</sub> and N<sub>2</sub>O (Jørgensen et al. 2012; Vroom et al. 2022).  
115 However, by avoiding plants we also isolate the soil processes leading to net soil emission/uptake of the GHG  
116 and resolve spatiotemporal patterns to a higher degree than previous studies at this site have achieved and what  
117 other commercial platforms are capable of. Collectively, this can provide a mechanistic insight into the

118 regulation of fluxes by hydrology and temperature. We acknowledge that future studies of GHG fluxes in  
119 peatlands should seek to include the aboveground plant component to the net GHG flux from the ecosystem.

120 The disturbance to the transect related to initial harvesting and removal of aboveground plants and continuous  
121 removal of aboveground live plant inside the collars and in a small perimeter outside the collar. In this way we  
122 kept an approximate area of 40 x 40 cm clear of vegetation at each collar. Two months prior to installation of  
123 collars in summer of 2021, the transect (Fig. 2) was harvested and remaining living aboveground vegetation was  
124 killed by applying one recommended dose of glyphosate (~100 mg m<sup>-2</sup>) to the plants only across the transect and  
125 avoiding spraying on the soil surface. The half-life of labile glyphosate in mineral soils range between 6-87 days  
126 (average 21 days) (Padilla and Selim 2020) with clay contents increasing half-life. The absence of clay and low  
127 dosage indicate that there were no, or only little traces of glyphosate left once the flux measurements began and  
128 hence the glyphosate treatment likely did not have a direct impact on the measured fluxes. Continued glyphosate  
129 application would potentially have reduced microbial activity in the soil and thus lower microbial respiration  
130 (Nguyen et al. 2016). Considering that we sprayed the vegetation only one time with glyphosate months prior to  
131 flux measurements, we assume the direct impact on soil microbial processes to be small. However, we cannot  
132 fully rule out that glyphosate may have led to a transient response. Because we did not have an undisturbed  
133 control we cannot quantify the effects of glyphosate. Subsequently, regrowth inside the collars was restricted by  
134 manual harvesting of emerging plants at a minimum of once every 7 days and throughout the period. Plant  
135 removal from collars is considered a common practice to isolate net soil GHG fluxes as the aboveground  
136 autotrophic respiration is removed. Since the individual collars were not trenched it is unavoidable to include  
137 belowground autotrophic respiration from plants growing adjacent to the collars. To avoid excessive disturbance  
138 of the site we did not remove these roots. Since we did not have a control, untreated/unharvested plot it is not  
139 possible to assess the direct impact of the disturbance on the GHG fluxes.



140

141 **Figure 1: The Vejrumbro location in Jutland (N 56.43819 E 9.54527 (WGS 84)) in the Nørre Å valley near the village**  
 142 **of Vejrumbro. The grey circle marks the placement of the SkyLine2D system. Satellite images: © Google Earth.**

143 2.1.2 Peat and organic soil characteristics

144 In November 2023 the peat across the SkyLine2D transect was sampled to 1 meter depth using a Russian auger  
 145 and cores split into five layers of 20 cm thickness. Collars 1, 2, 5, 6, 8, 13 – 27 were sampled. For the remaining  
 146 collars it was not possible to retrieve a sample due to excessive wetness of the peat. The decomposition of the  
 147 peat samples were assessed by a 10-point Von Post scale of humification (1 = completely undecomposed and 10  
 148 = completely decomposed) together with quantification of the pH<sub>H<sub>2</sub>O</sub> (1:5 peat:water mix), dry bulk density (g  
 149 cm<sup>-3</sup>) and total C and N by dry combustion (g C/N 100 g peat<sup>-1</sup> or %) (Table 1).

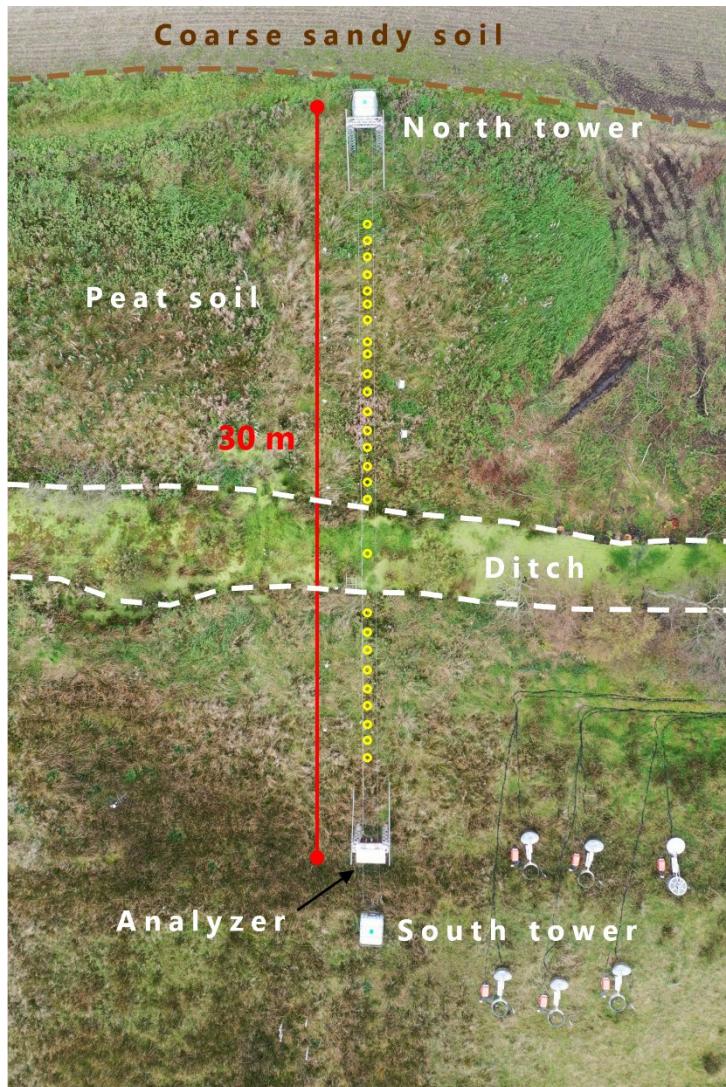
150 **Table 1 Mean (±standard error of the mean (SE)) peat/organic soil characteristics of humification degree (Von Post),**  
 151 **pH (H<sub>2</sub>O), dry bulk density (ρ<sub>dry</sub>), total C (TC) concentration, total N concentration (TN) and the C/N ratio for collars**  
 152 **1, 2, 5, 6, 8 and 13 - 27 at the Vejrumbro transect.**

Depth (cm)	N	Von post		pH (H <sub>2</sub> O)		ρ <sub>dry</sub> (g cm <sup>-3</sup> )		TC (%)		TN (%)		C/N	
		Min	Max	Mean	±SE	Mean	±SE	Mean	±SE	Mean	±SE	Mean	±SE
0-20	20	7	10	4.2	0.08	0.31	0.02	26	1.1	1.6	0.06	16	0.4
20-40	20	5	10	4.6	0.06	0.20	0.01	43	1.3	1.8	0.04	24	0.7
40-60	11	3	8	4.9	0.10	0.15	0.01	48	1.8	1.9	0.05	25	1.1
60-80	11	3	6	5.3	0.09	0.11	0.01	47	1.8	1.9	0.05	24	0.6
80-100	10	1	8	5.4	0.09	0.10	0.02	44	2.1	1.9	0.05	24	0.6

153 Generally, there was peat/organic soil to one meter depth except for one collar (25) where gyttja was found in a  
154 depth of 80 cm (Table 1). The organic soil was more decomposed in the top 40 cm indicated by higher Von Post  
155 values between 5 and 10. Below 40 cm peat still displayed high levels of decomposition along the transect, but  
156 was more often found to be less decomposed, values ranging from 1-8 (Table 1). This corresponds well to the  
157 previous land use with drainage of the topsoil leading to higher degree of humification. Also, the organic soil  
158 was most dense in the top 20 cm (on average  $0.31\pm0.02\text{ g cm}^{-3}$ ) and bulk density decreased to  $0.10 - 0.12\text{ g cm}^{-3}$   
159 from 40 – 100 cm depth. Total C and N was lowest in the 0-20 cm layer, but still classified as organic soil.  
160 Below 20 cm total C and N concentrations, respectively were similar. C/N ratio was lowest in the top 20 cm  
161 ( $16\pm0.4$ ) and increased to 22-25 in 20 – 100 cm depth (Table 1).

162 2.1.3 Groundwater water sampling and chemical analysis

163 Groundwater was sampled monthly in the piezometers placed at collars 1, 5, 13, 18, 22 and 27 (Fig. 3) by  
164 retrieving a 200 mL sample 20-30 cm below the groundwater level at the sampling time. The water sample was  
165 retrieved using a syringe and transferred to a plastic bottle that was capped to avoid air bubbles. Water samples  
166 were frozen immediately after sampling and subsequently after thawing analyzed for pH, EC and alkalinity on a  
167 855 Robotic Titrosampler (Metrohm, Germany). Total N and DOC were measured on a TOC-V CPH Analyzer  
168 with Total Nitrogen Unit TNM-1 & ASI-V Autosampler (Shimadzu, Japan). Ion chromatograph (IC) analyses  
169 of  $\text{Cl}^-$ ,  $\text{NO}_3^-$ , and  $\text{SO}_4^{2-}$  were performed on a 930Compact IC Flex (Metrohm, Germany) and  $\text{NH}_4^+$   
170 concentrations were measured with continuous flow analysis using a Seal AA500 Autoanalyzer (SEAL  
171 Analytic, USA). Total dissolved Fe and P were analyzed with coupled plasma–mass spectrometry (ICP-MS) on  
172 an iCAP-Q ICP-MS (Thermo Fisher Scientific, USA) in KED mode using He as the collision gas. Prior to  
173 analysis the 10 mL subsamples were acidified with 200  $\mu\text{L}$  concentrated nitric acid to a 10 mL sample.  
174 Elemental ICP-MS analyses also included dissolved base cations of  $\text{Ca}^{2+}$ ,  $\text{Mg}^{2+}$ ,  $\text{K}^+$ ,  $\text{Na}^+$  as well as total  
175 dissolved Al and Mn cations (not shown, but included in the data set).



176

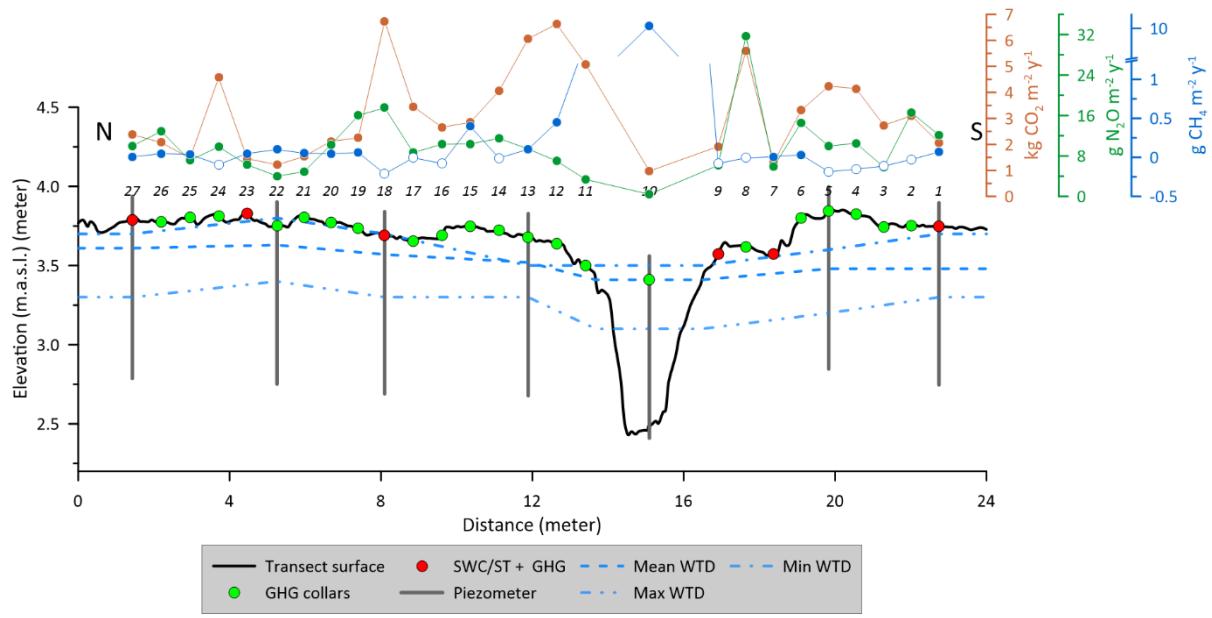
177 **Figure 2: Drone image of the measurement transect (September 27<sup>th</sup>, 2023) after flux measurements had stopped.**  
 178 **Dashed brown line marks the approximate boundary between the agricultural field, coarse sandy soil (north) and the**  
 179 **peat/organic soil (south). The red line marks the end points of the SkyLine2D system (30 meters). The open yellow**  
 180 **circles (n=27) mark the approximate position of individual collars across the transect of the field (24 meters in length)**  
 181 **where greenhouse gas fluxes were measured. The ditch is placed between the dashed white lines. The analyser was**  
 182 **placed at the south tower. Elevation above sea level along the 24 meter collar transect varied little from 3.77 m in the**  
 183 **south to 4.06 m in the north.**

184 2.1.4 SkyLine2D system configuration at Vejrumbro

185 The SkyLine2D system is an automated chamber based system for measuring GHG fluxes. The system is  
 186 designed and built by Earthbound Scientific Ltd. (United Kingdom). We used the SkyLine2D system to measure  
 187 the net soil fluxes of carbon dioxide (CO<sub>2</sub>), methane (CH<sub>4</sub>) and nitrous oxide (N<sub>2</sub>O) measured with an automated  
 188 GHG chamber system over 12 months resolving spatiotemporal patterns of GHG fluxes including 27 individual  
 189 collars (26 on organic soil and 1 in a ditch) over a 24 m transect on a temperate drained fen peatland.

190 The SkyLine2D system transect was oriented in an north-south direction (Fig. 2). Two 2.5 meter-tall scaffold  
 191 towers marked the end of the 30 m SkyLine2D system (Fig. 2 and Fig. S2D). The towers were fixed by ropes

192 attached to 1000L pallet tanks filled with water (Fig. S2D) that maintained a stable position of the towers and  
 193 ropes and hence placement of the chamber over the collars. The measurement transect was in total 24 m with 27  
 194 individual measurement collars for GHG fluxes on the ground, 26 on organic soil and 1 in a drainage ditch (Fig.  
 195 2 and 3). The GHG analyser (model G2508, Picarro Inc., USA) was installed in a waterproof and temperature-  
 196 controlled shelter at the south end of the transect (Fig. 2 and Fig. S2C). The transect was situated on the edge of  
 197 the riparian fen in close proximity to the mineral upland soils, where active agriculture was practiced (Fig. 2).  
 198 Along the transect volumetric soil water content (SWC) and soil temperature (ST) as well as water table depth  
 199 (WTD) were measured at seven locations (Fig. 4). The agricultural field north of the SkyLine2D was sown with  
 200 annual crops in rotation according to normal practice.



201 **Figure 3: Schematic representation of the measurement transect at Vejrumbro and associated measurement**  
 202 **variables. The annual cumulative fluxes of CO<sub>2</sub> (red) (kg CO<sub>2</sub> m<sup>-2</sup> y<sup>-1</sup>), N<sub>2</sub>O (green) (g N<sub>2</sub>O m<sup>-2</sup> y<sup>-1</sup>) and CH<sub>4</sub> (blue) (g**  
 203 **CH<sub>4</sub> m<sup>-2</sup> y<sup>-1</sup>) are shown for each collar across the measurement transect at Vejrumbro. Closed and open symbols for**  
 204 **CH<sub>4</sub> represent net cumulative emission and uptake, respectively. Mean WTD is the mean water table depth measured**  
 205 **in piezometers (blue dashed line). GHG collars (green symbols) mark the positions of greenhouse gas flux**  
 206 **measurements of CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O. SWC/ST + GHG mark the positions where volumetric soil water content**  
 207 **(SWC) and soil temperature (ST) at 5 cm depth were measured alongside greenhouse gas fluxes. Numbers on top of**  
 208 **plot show the collar numbers (from 1 – 27). N and S mark the north and south ends of the transect (see Fig. 3). The**  
 209 **peat depth was at least one meter in all points.**

211 2.2 Overview of time series of GHG fluxes, soil temperature/moisture, air temperature, wind direction and  
 212 groundwater level

213 The dataset is comprised of a 12-month time series of net soil fluxes of CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O, accompanied by a  
 214 longer timeseries of soil temperature and moisture at 5 cm depth, meteorological variables (air temperature,  
 215 wind speed and direction measured at 2 meter height) and a shorter time series groundwater table level, depth  
 216 and temperature (Fig. 3, Table 2). Due to equipment failure of the SkyLine2D the GHG flux measurements  
 217 started on February 2<sup>nd</sup>, 2022 (Table 2). Groundwater level measurements started between March 9<sup>th</sup> to 31<sup>st</sup>,  
 218 2022 (Table 1). All other variables were measured continuously from July 1<sup>st</sup>, 2021, until January 31<sup>st</sup>, 2023

219 (Table 1). In the period between December 7<sup>th</sup> and 19<sup>th</sup>, 2022 intermittent periods of snow cover (depth was not  
220 measured) on the ground occurred. This snow cover did not impede flux measurements.

Table 2: Available time series data from the Vejrumbro SkyLine2D system. Coloured time periods in 2021 to 2023 for each variable indicate data availability.

Variable	Unit	Model/sensor type	Data availability																		
			2021			2022			2023												
			Frequency (minutes)	Aug	Sep	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Jan
CO <sub>2</sub> flux*	µmol CO <sub>2</sub> m <sup>-2</sup> s <sup>-1</sup>	G2508 (Picarro Inc., USA)	~10**																		
CH <sub>4</sub> flux*	nmol CH <sub>4</sub> m <sup>-2</sup> s <sup>-1</sup>	G2508 (Picarro Inc., USA)	~10**																		
N <sub>2</sub> O flux*	nmol N <sub>2</sub> O m <sup>-2</sup> s <sup>-1</sup>	G2508 (Picarro Inc., USA)	~10**																		
Soil temperature at 5 cm depth***	°C	RXW-TMB-868 (Onset, USA)	5																		
Soil water content at 5 cm depth***	(cm <sup>3</sup> cm <sup>-3</sup> )	RXW-SMD-868 (5HS) (Onset, USA)	5																		
Air temperature at 2 m height	°C	S-THC-M002 (Onset, USA)	5																		
Wind speed	m s <sup>-1</sup>	S-WSB-M003 (Onset, USA)	5																		
Wind direction	°	S-WDA-M003 (Onset, USA)	5																		
Groundwater level****	m.a.s.l.	DCL532 (BD sensors, Germany)	15																		
Groundwater table ****	cm	DCL532 (BD sensors, Germany)	15																		
Groundwater temperature***	°C	Dallas DS 18B20	15																		

\*Net soil/ditch fluxes for all collars 1 - 27.

\*\*Time in between two consecutive flux measurements. The 10 minutes comprise actual flux measurement of 5 minutes and 5 minutes headspace flushing between flux measurements.

\*\*\*Measured for a subset of collars: 4, 7, 9, 23, 27.

\*\*\*\*Measured for a subset of collars: 1, 5, 10 (ditch), 13, 18, 22, 27.

226 **2.3 Soil moisture and temperature measurements**

227 Soil moisture was measured at collars 1, 7, 9, 18, 23, 27 (Figure 4) and probes (6 cm length) were inserted at an  
228 approximate 30° angle 5 cm outside the collar, while the soil temperature probes were inserted vertically  
229 adjacent to the soil moisture probe.

230 **2.4 Groundwater table level and depth**

231 Piezometers (inner diameter 5 cm) were installed at collars 1, 5, 10 (ditch), 13, 18, 22, 27 (Figure 4) to 1 meter  
232 depth below the surface, which is deeper than the lowest groundwater level in summer (~60 cm below the  
233 surface) with openings from 0.1 – 1.2 meter below terrain. In the ditch the piezometer bottom was deeper than  
234 one meter to secure anchoring in the peat. The piezometers were installed approximately 50-60 cm beside the  
235 collars to avoid interference with the SkyLine2D system. After installation, piezometers were cleaned and  
236 sealed at the surface with bentonite pellets to avoid surface infiltration along the piezometers which can distort  
237 water level measurements.

238 Pressure transducers (Table 2) connected to Arduino-loggers were installed in each piezometer (at collars 1, 5,  
239 10, 13, 18, 22 and 27 – Fig. 3) approximately 1 m below terrain measuring water levels every 15 minutes. The  
240 pressure transducers were vented and thus do not need correction for atmospheric pressure.

241 The groundwater levels were described using two metrics: hydraulic head and groundwater depth (GWD).  
242 Hydraulic head represents the water level relative to mean sea level, based on the Danish Vertical Reference  
243 (DVR90), while GWD indicates the depth of the groundwater below the surface terrain. The elevation of top of  
244 the piezometers were measured using a GPS (model GS07 High Precision GNSS Antenna with a CS20  
245 Controller, Leica, Germany) and used as a local reference for hydraulic head. Manual measurements of  
246 groundwater levels were conducted every 2 months and used to calibrate the logger water levels to hydraulic  
247 head and GWD.

248 **2.5 Wireless data transfer**

249 Wireless sensors for air temperature, wind speed, wind direction, soil temperature and volumetric soil water  
250 content were set up with Wi-Fi data transfer to HOBO RX3000 Weather Station (Onset, USA) equipped with  
251 HOBOnet Manager (RXMOD-RXW-868) module for wireless communication with sensors and logged data  
252 every 5 minutes. Data access was through the HOBOlink cloud software.

253 Groundwater loggers were interfaced with the I<sup>2</sup>C (Inter-integrated Circuit) protocol and data was collected on  
254 Arduino custom-built logger (<https://vandstande.dk/logger.php>) with wireless connection via LoRaWAN or  
255 SigFox.

256 **2.6 Greenhouse gas flux measurements with the SkyLine2D system at Vejrumbro**

257 Along the SkyLine2D transect the 26 individual collars (Ø19 cm) along the 24 meter transect on organic soil  
258 (Fig. 3) were inserted 5 cm into the peat leaving 5 cm above the surface. The collars were distanced app. 70 cm  
259 apart. One collar was installed in the ditch by inserting a tube (Ø19 cm, length 100 cm) to the bottom of the  
260 ditch with holes deeper than the minimum water level in the ditch to allow water flow. Thus, it was avoided that

261 air entered in the collar in the ditch due to low water levels in the ditch. On top of this longer tube a collar (Ø19  
262 cm, length 10 cm) was glued allowing for flux measurements. The chamber was programmed to stop when the  
263 bottom of the chamber sat the water surface if the water level in the ditch extended above the top of the collar.  
264 For most of the time the collar was not submerged and the chamber therefore hit the collar.

265 There was one round transparent chamber (height: 39.5 cm and inner Ø: 19 cm, volume: 11.2 L) on the  
266 SkyLine2D, hanging below a moving trolley, which was suspended on two ropes stretched between the north  
267 and south towers (Fig. S2A and B). At defined positions along the rope, neodymium magnets had been inserted,  
268 and a magnet sensor (Fig. S2B) on the trolley informed the internal computer to stop and lower the chamber  
269 over positions with a collar on the surface. The chamber was lowered and guided down to the collar by  
270 supporting rods shaping a funnel (Fig. S2A). The chamber stopped when it hit the collar, achieved through a  
271 pressure sensor on top of the chamber connected to a hollow rubber gasket (Ø 3 cm) at the bottom, which also  
272 sealed the chamber with the collar. There was no fan installed in the chamber as the mixing was ensured by the  
273 main pump (Fig. S2C). A vent was installed in the top of the chamber to allow for pressure equilibration under  
274 windy conditions and chamber deployment.

275 One entire flux + flushing sequence lasted 10 minutes (Table 1). The chamber closure period was set to 5  
276 minutes with a purging time of 5 minutes in between measurements when chamber was open and hanging  
277 underneath the trolley at approximately 1 meter above the ground (Fig. S2D). This provided on average 10 min  
278 between flux measurements on consecutive collars (Table 1). Due to small variations in mechanical operations,  
279 flux measurements were occasionally farther apart than 10 minutes, but overall, the timing of the SkyLine2D  
280 system was consistent. After each cycle of 27 flux measurements there was a 30-minute delay until the start of  
281 the next cycle. On average this resulted in 4-5 flux measurements per collar per day throughout the period.

282 To determine the concentrations of CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O in the chamber air, a laser spectroscopy GHG analyser  
283 (model G2508, Picarro Inc., USA) was used. The sample output frequency was set to 1 Hz with a manufactured  
284 specified raw precision on 1 Hz data for CO<sub>2</sub>: 240 ppb, CH<sub>4</sub>: 0.3 ppb and N<sub>2</sub>O: 5 ppb at ambient conditions  
285 (Picarro Inc., USA). A main pump (model: N86 KN.18, KNF, Germany) circulated the air to and from the  
286 chamber at 6 L min<sup>-1</sup>. The GHG analyser was installed in parallel to the inflow from the chamber due to the  
287 much lower flow of 250 mL min<sup>-1</sup> of the vacuum pump. There was a 30-meter tube between the chamber and  
288 main pump to allow for the GHG analyser to remain stationary in the hut while the trolley moved.

## 289 2.7 Calculation of diffusive fluxes

290 Fluxes were calculated and quality checked using the goFlux R package (Rheault et al. 2024) and presented as  
291  $\mu\text{mol CO}_2 \text{ m}^{-2} \text{ s}^{-1}$ ,  $\text{nmol N}_2\text{O m}^{-2} \text{ s}^{-1}$  and  $\text{nmol CH}_4 \text{ m}^{-2} \text{ s}^{-1}$ . Prior to flux calculations, the gas concentration data  
292 from the G2508 was matched to the chamber closure time and chamber id in order to determine the start time of  
293 the chamber measurement, so it was possible to separate individual flux measurements from each collar over the  
294 measurement time (see examples of flux detection and calculation in Fig. S3A-D). An automatic deadband  
295 detection method was applied based on maximal R<sup>2</sup> of a linear regression over the first 180 s (in 10 s steps) after  
296 chamber closure. The deadband was allowed to attain values between 0 to 150 seconds thereby also allowing for  
297 compensation for the ~60 s delay between chamber headspace gas concentration change and GHG analyser  
298 detection due to transport time through the 30 m tube connecting the chamber and GHG analyser.

299 Flux calculations were done with both linear (LM) and non-linear (Hutchinson-Mosier – HM) regression models  
300 (Pihlatie et al. 2013) to determine the slope at time zero. The best flux estimates with either the LM or HM  
301 regression model was determined using the *best.flux* function in the goFlux package (Rheault et al. 2024).  
302 Shortly, if the RMSE of the HM model was lower than minimum detectable flux (MDF), HM was chosen.  
303 However, if the ratio (g-factor) between HM and LM was larger than 2, LM was chosen, as this indicates over-  
304 fitting of the HM, which may result in unrealistic large HM flux estimates. If the relative SE of the slope  
305 (SE/slope) at time zero for the HM model was larger than 100% it indicated overfitting of the HM model and  
306 the LM was chosen. This approach is conservative as it will discard non-linear flux behaviour and instead  
307 provide a conservative linear flux estimate. Out of 47.438 detected flux measurements for CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O,  
308 respectively, a total of 2807 CO<sub>2</sub> fluxes (5.9%), 3339 N<sub>2</sub>O fluxes (7%) and 4923 CH<sub>4</sub> fluxes (10.3%) were  
309 discarded either due to chamber mechanical malfunction (imperfect sealing on collar due to erroneous lowering  
310 of chamber on collar indicated by background atmospheric or fluctuating gas concentrations in the headspace).  
311 At low flux levels non-significant fluxes were discarded as it was not possible to visibly detect whether there  
312 was a flux due to high noise-signal ratio of the analyser and/or it was because the chamber had malfunctioned. It  
313 is acknowledged that discarding low fluxes can bias annual means and cumulative values, but the data quality  
314 did not allow us to determine whether the flux measurement was performed correctly and hence a conservative  
315 approach was chosen as including false low fluxes would also bias the data set.

316 For flux measurements the air temperature in 2 meters was used as an estimate of the chamber headspace  
317 temperature along with a 1 atm air pressure.

318 The annual cumulated fluxes from the soil or the ditch were estimated simply by multiplying the daily average  
319 CO<sub>2</sub>, CH<sub>4</sub> or N<sub>2</sub>O flux for the measurement period with 365 days. We believe for the purpose of data  
320 presentation that this simplistic methodology is adequate here, also given the very few data gaps in the  
321 timeseries. However, there are other more sophisticated methods using interpolation and response variable  
322 functions that may refine the annual budget. However, it is not the goal of this manuscript to present these  
323 methodologies but to provide the data so other users can test different temporal upscaling methodologies.

## 324 **2.8 Calculation of ebullition fluxes in the ditch**

325 Methane ebullition fluxes were occasionally observed only in the ditch. The resultant CH<sub>4</sub> time series for the  
326 chamber would have a characteristic appearance (Fig. S4) where the measurement would essentially start out as  
327 diffusive flux measurement, then CH<sub>4</sub> bubbles entered the chamber headspace, and the concentration would  
328 quickly increase to a maximum value and reach a threshold concentration corresponding to the mixed headspace  
329 concentration. In these cases, the LM/HM flux calculation assumptions are violated and instead the ebullition  
330 flux would be calculated as the total increase in CH<sub>4</sub> mass m<sup>-2</sup> per 5 min enclosure. The mass flux of CH<sub>4</sub> per  
331 enclosure (nmol m<sup>-2</sup> per 5 min enclosure) was calculated according to Eq. (1):

$$332 F_{CH_4-ebu} = dCH_4 * \frac{V_{system}*P}{A*R*T} \quad (1)$$

333 Where dCH<sub>4</sub> is the concentration difference in nmol between start of chamber enclosure (CH<sub>4,start</sub>) and end CH<sub>4</sub>  
334 concentration (CH<sub>4,end</sub>) after it reached a plateau (Fig. S4), V<sub>system</sub> is the total volume (11.7 L) of the system  
335 (collar, chamber, tubes and GHG analyser) in L, P is the pressure (1 atm), A is the area of the collar (0.028 m<sup>2</sup>),

336 R is the gas constant (0.082057 L atm K<sup>-1</sup> mol<sup>-1</sup>) and T is the chamber headspace temperature (K). The time step  
337 of dCH<sub>4</sub> was assumed to be 1 second meaning that the flux unit is nmol CH<sub>4</sub> m<sup>-2</sup> s<sup>-1</sup>.

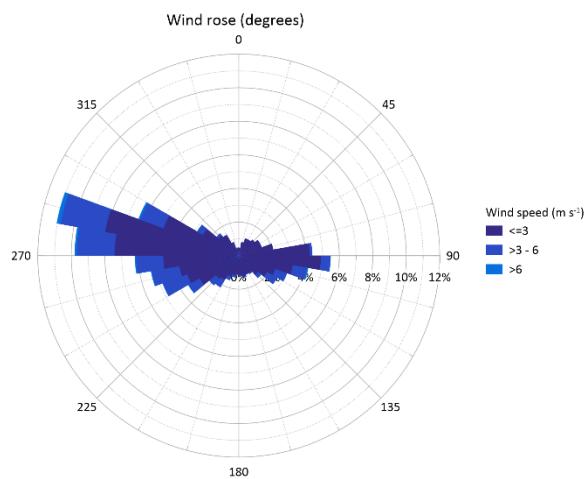
338 Out of a total of 1728 flux measurements from the ditch (collar 10), 334 were classified as ebullitions indicating  
339 that ebullition was erratic which is in line with studies of ebullition of fluxes from ponds (Wik et al. 2016; Sø et  
340 al. 2023). Hence, it can be assumed that ebullition occurred around 19.3% of the time during the measurement  
341 period (360 days). Furthermore, the ebullition flux is calculated as the accumulated CH<sub>4</sub> in the chamber  
342 headspace during the entire flux measurement, e.g. 5 minutes here (Sø et al. 2023), and the calculated ebullition  
343 flux in the data set is therefore representative of 5 minute enclosure and not per second. To extrapolate to an  
344 annual estimate the number of 5 minute enclosures in 19% of 360 days is therefore estimated (N=20049 5-min  
345 360 days<sup>-1</sup>), multiplied with the average ebullition flux (nmol CH<sub>4</sub> m<sup>-2</sup> 5 min<sup>-1</sup>).

346 Ebullitions could also be caused by mechanical disturbance of the chamber landing on the collar. Ebullition  
347 fluxes were discarded if the sudden increase in CH<sub>4</sub> headspace concentration (Fig. S4) occurred 30 seconds after  
348 recorded chamber closure as this indicated bubbles released by chamber deployment on top of the collar.

349 **3 Data presentation**

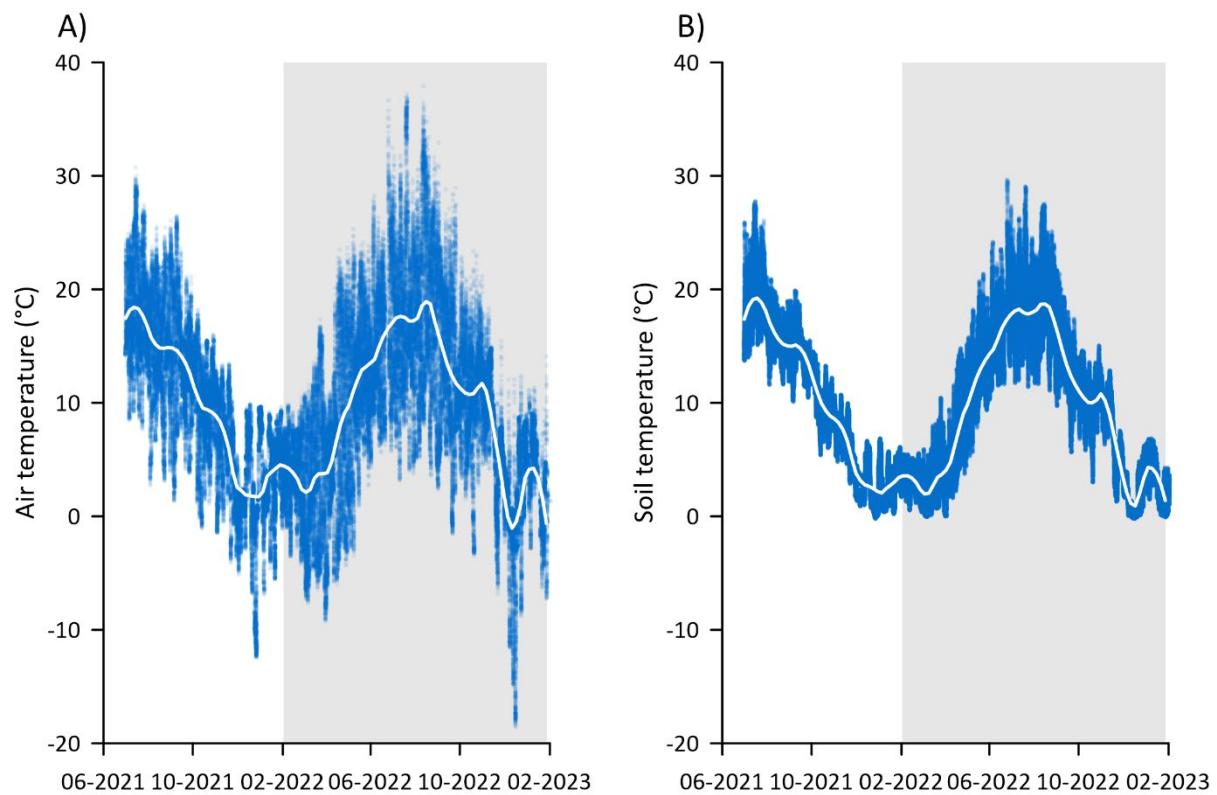
350 **3.1 Wind speed and direction**

351 Generally, the wind regime during the measurement period was rather mild with monthly average wind speeds  
352 ranging between  $1.2$  to  $2.9 \text{ m s}^{-1}$  and maximum gust up to  $20 \text{ m s}^{-1}$ . The wind direction was uniformly from the  
353 west for 52% of the time, with easterly winds constituting 27% and northern and southern winds 8 and 13% of  
354 the time (Fig. 4). Winds from western directions were highest for the longest period, while easterly winds were  
355 of similar magnitude, but less frequent (Fig. 4). Northern and southerly winds were generally below  $3 \text{ m s}^{-1}$  and  
356 represented periods with still conditions. The very uniform western-eastern wind field at Vejrumbro may also  
357 partly be explained by the W-E direction of the valley in which the site is situated, that effectively blocks or  
358 dampens winds from S and N.



359

360 **Figure 4: Wind regime at Vejrumbro for the period July 1<sup>st</sup>, 2021 to January 31<sup>st</sup>, 2023 presented as a wind rose**  
361 **diagram with wind speed and direction for the period.**



363  
 364 **Figure 5:** Time series of A) air temperature in °C measured at 2 meter height above the surface and B) soil  
 365 temperature (°C) at 5 cm depth for collars 4, 7, 9, 23 and 27 along the measurement. The blue dots are the raw 5 min  
 366 measurements of air temperature and the white lines represent are LOESS fit to show overall seasonal trend. The  
 367 periods of GHG measurements with the SkyLine2D system are shown with the shaded area.

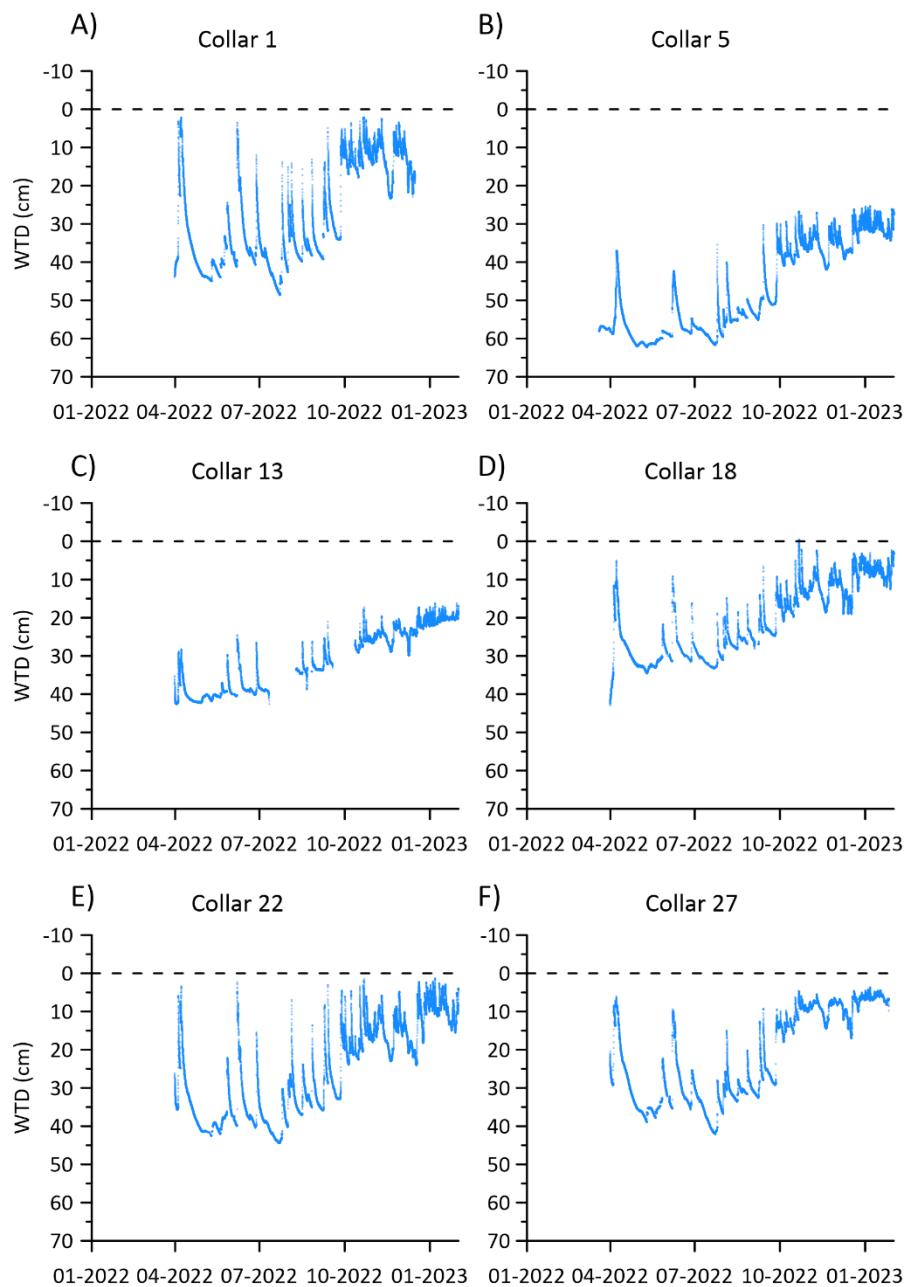
368 Over the study period the average air temperature was 9.6°C ranging between maximum 37.9°C and minimum  
 369 of -18.6°C (Fig. 5A). Monthly ranges of air temperatures (Tab. 2) show >20°C variation between minimum and  
 370 maximum, except for February, pointing towards large diurnal variations. Soil temperature magnitude and  
 371 temporal variation were similar across the transect, varying between 0 to 28°C (Fig. 5B) and followed that of air  
 372 temperature (Fig. 5A) with less variability (Fig. 5B and Table 3). The annual site average soil temperature was  
 373 similar to the air temperature (Table 3).

374 **Table 3: Monthly mean, maximum and minimum air temperature and soil temperature (°C), groundwater table**  
 375 **depth (cm) and volumetric soil water content (cm<sup>3</sup> cm<sup>-3</sup>) at Vejrumbro in the measurement period from February 1<sup>st</sup>,**  
 376 **2022 to January 31<sup>st</sup>, 2023.**

Variable	Month	Year												
		2022						2023						
Air temperature (°C)	Mean	3.8	3.0	6.6	12.0	15.4	17.7	16.6	13.4	10.7	6.9	1.2	3.7	9.6
	Max	10.6	17.4	23.7	25.3	36.7	37.2	37.9	32.9	23.3	18.4	12.4	14.1	-
	Min	-4.3	-9.3	-8.3	-3.4	4.3	3.2	2.7	-1.5	-3.5	-6.9	-18.6	-7.3	-
Soil temperature (°C)	Mean	3.0	3.2	2.9	6.4	12.3	16.1	18.4	17.0	13.8	10.3	7.2	2.1	9.6
	Max	6.5	5.3	9.1	12.5	18.8	25.1	27.0	24.7	19.3	14.3	12.6	6.3	-
	Min	0.3	1.1	0.4	0.8	6.6	10.7	12.4	11.8	7.0	4.0	2.1	0.0	-
Groundwater table depth (cm)	Mean	-	39	35	41	36	41	35	31	20	18	17	13	29
	Max	-	58	39	58	43	52	46	36	30	31	28	28	-
	Min	-	23	5	24	9	28	22	9	5	6	3	2	-
Volumetric soil water content (cm <sup>3</sup> cm <sup>-3</sup> )	Mean	0.53	0.45	0.40	0.37	0.38	0.43	0.43	0.45	0.50	0.53	0.52	0.51	0.46
	Max	0.56	0.51	0.50	0.41	0.47	0.55	0.56	0.56	0.57	0.58	0.56	0.57	-
	Min	0.43	0.39	0.37	0.33	0.32	0.26	0.32	0.35	0.40	0.47	0.42	0.34	-

### 377 3.3 Groundwater table depth

378 Average groundwater table depth (WTD) below terrain during the period was between 47 to 21 cm across the  
 379 transect (Fig. 3, Table 3). During summer, the peat drained between 18 – 31 cm below the annual average and in  
 380 winter the WTD increased to 0 – 22 cm above the annual average across the transect (Fig. 3, Table 3).  
 381 Generally, the WTD was lower in the ditch across the entire study period (Fig. 3). It was only on the northern  
 382 end of the transect that the surface occasionally was flooded during winter periods (Fig. 3).



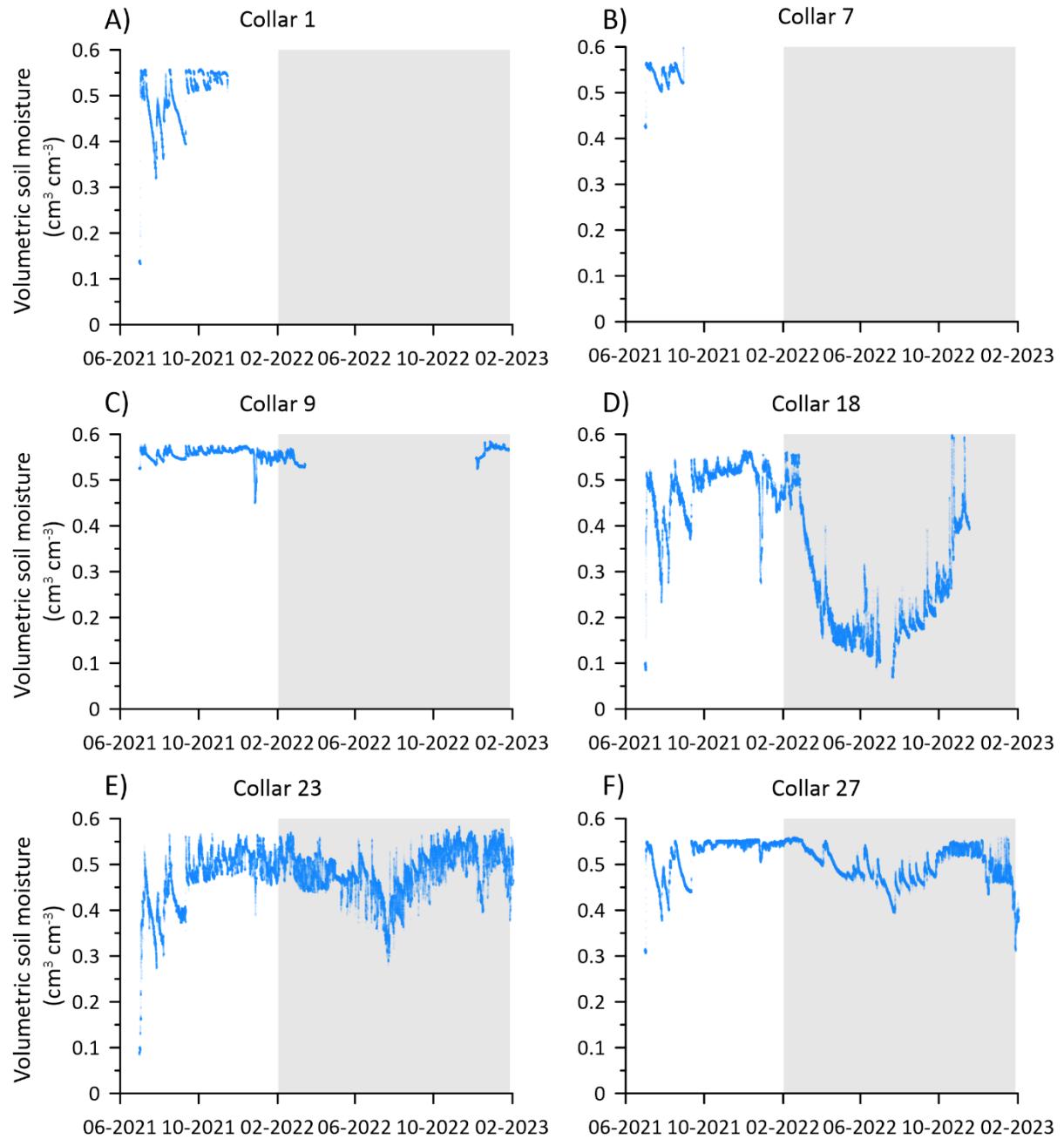
383

384 **Figure 6: Time series of groundwater table depth (WTD) below terrain for the six piezometer locations along the**  
 385 **SkyLine2D transect in the period March 31<sup>st</sup>, 2022 and January 31<sup>st</sup>, 2023 when the flux measurements stopped.**  
 386 **Dashed line show surface.**

387 The temporal variability of WTD was similar across the transect despite different absolute water table depths  
 388 (Fig. 6A-F). In the summer periods, the WTD was most variable decreasing to below -40 for collars 1, 13, 18,  
 389 22 and 27, whereas the WTD for collar 5 showed the deepest groundwater measured at the site. WTD responded  
 390 quickly (within hours) to precipitation events that could increase the WTD by almost 40 cm at some plots,  
 391 indicating that the entire aerated soil volume above the groundwater table was flooded. There was a slight  
 392 tendency to lower response to precipitation events for piezometers at collar 5 and collar 13 that were placed  
 393 closer to the ditch (Fig. 3 and 6B and C). As the ditch water level was lower than in the peat this could be  
 394 explained by more efficient lateral drainage into the ditch from the areas closer to the ditch. In the winter

395 periods, the WTD was less responsive to precipitation and was closer to the surface (Fig. 6A-F) across the  
396 transect.

397 **3.5 Soil water content**



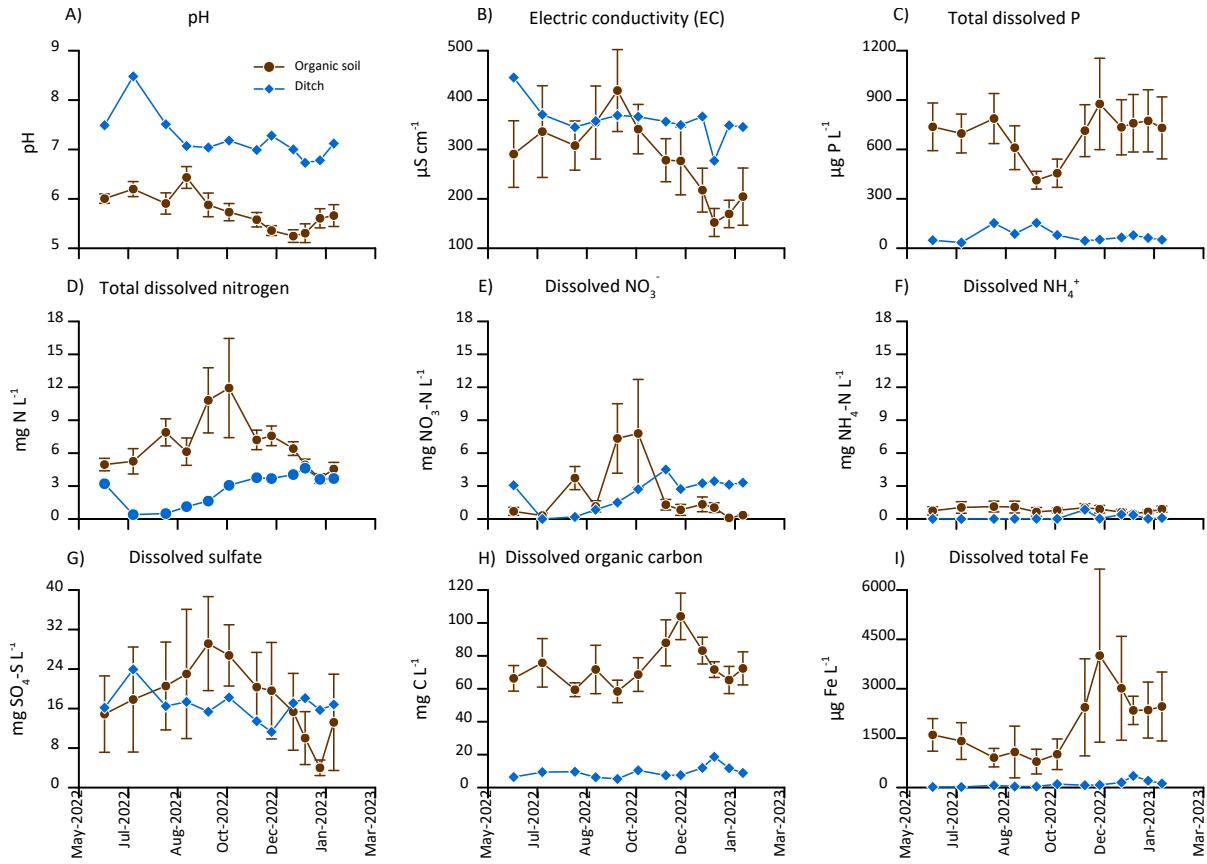
398  
399 **Figure 7: Time series of volumetric soil water content ( $\text{cm}^3 \text{cm}^{-3}$ ) in 0-5 cm for the six collars 1, 7, 9, 18, 23 and 27**  
400 along the SkyLine2D transect in the period July 1<sup>st</sup>, 2021 – January 31<sup>st</sup>, 2023 when the measurements terminated.  
401 The periods of GHG measurements with the SkyLine2D system are shown (green lines) on the x-axis.

402 Due to instrument failure the temporal coverage of soil moisture in the topsoil (5 cm) was not similar across the  
403 transect (Fig. 7A-F). For collars 18, 23 and 27 the entire period of greenhouse gas measurements was covered  
404 by soil moisture measurements (Fig. 7D-F). While SWC for collars 1, 9, 18, 23 and 27 was similar in the winter  
405 periods (around  $0.55 \text{ cm}^3 \text{cm}^{-3}$ ) the SWC for collar 18 decreased to lower minima between  $0.1 – 0.2 \text{ cm}^3 \text{cm}^{-3}$ ,

406 than the minima observed between  $0.3 - 0.4 \text{ cm}^3 \text{ cm}^{-3}$  for collars 23 and 27 in the summer periods (Fig. 7, Table 407 3). Similar for all collars it was observed that SWC was more variable in summer, responding similarly as WTD 408 to precipitation events (Fig. 7, Table 3). Since plants were removed regularly from the collars the decrease of 409 SWC for collar 18 cannot be explained by plant transpiration, and the dynamic behaviour could indicate the 410 impact of soil evaporation, but the different levels of SWC also show that there is spatial variation across the 411 transect in the drying properties of the soil. However, it cannot be ruled out that the SWC sensor at collar 18 412 experienced malfunction or that soil contact was lost in the dry periods of 2022 (Fig. 7D) which could lead to 413 erroneous and too low SWC. Therefore, these data should be considered with care.

#### 414 **3.6 Groundwater and ditch water chemical composition**

415 Site mean pH of the groundwater in the organic soil was  $5.8 \pm 0.1$  and was lower than the pH of the ditch 416 ( $7.3 \pm 0.6$ ). There was a tendency towards lower pH in groundwater and ditch towards the end of the 417 measurement period (Fig. 8A). Electric conductivity was generally higher in the ditch water ( $359 \pm 36 \mu\text{S cm}^{-1}$ ) 418 compared to the groundwater in the organic soil ( $276 \pm 18 \mu\text{S cm}^{-1}$ ) but varied less over the season. The 419 groundwater shows a clear peak in EC around September 2022 (Fig. 8B). Total dissolved P was markedly 420 higher in the groundwater ( $687 \pm 45 \mu\text{g P L}^{-1}$ ) compared to the ditch water ( $76 \pm 10 \mu\text{g P L}^{-1}$ ). Whereas there was 421 little seasonal trend in ditch P concentrations, dissolved P in groundwater dipped to below average 422 concentrations between August to October, likely indicating plant uptake during the growing season (Fig. 8C). 423 Similarly, total dissolved N was higher in groundwater ( $6.7 \pm 0.5 \text{ mg N L}^{-1}$ ) than in ditch ( $2.6 \pm 1.6 \text{ mg N L}^{-1}$ ) with 424 increasing concentrations during the growing season (Fig. 8D). Similar, temporal trend was observed for  $\text{NO}_3^-$  425 (Fig. 8E), but average groundwater ( $2 \pm 0.5 \text{ mg NO}_3\text{-N L}^{-1}$ ) and ditch ( $2.2 \pm 1.5 \text{ mg NO}_3\text{-N L}^{-1}$ ) concentrations 426 were similar. As expected, dissolved  $\text{NH}_4\text{-N}$  was lowest among investigated N-species and there was more 427 dissolved  $\text{NH}_4\text{-N}$  present in groundwater ( $0.8 \pm 0.1 \text{ mg NH}_4\text{-N L}^{-1}$ ) than in the ditch ( $0.14 \pm 0.25 \text{ mg NH}_4\text{-N L}^{-1}$ ). 428 However, there was no discernable temporal trend for  $\text{NH}_4^+$  (Fig. 8F). Collectively, the temporal trend of TN 429 and  $\text{NO}_3^-$  could point to temperature driven mineralization of the peat. Also, the organic N (TN – inorganic N- 430 species) was on average 10 times higher in the groundwater than in the ditch. Average  $\text{SO}_4^{2-}$  concentrations 431 were similar between the groundwater ( $17.5 \pm 2.4 \text{ mg SO}_4\text{-S L}^{-1}$ ) and ditch ( $17 \pm 1.5 \text{ mg SO}_4\text{-S L}^{-1}$ ), but  $\text{SO}_4^{2-}$  432 concentration peaked during September and October in the groundwater whereas it remained more constant in 433 the ditch over the season (Fig. 8G). Similar to dissolved organic N, DOC concentrations were consistently 434 higher in the groundwater ( $73 \pm 3.1 \text{ mg DOC L}^{-1}$ ) than in the ditch ( $9.4 \pm 3.5 \text{ mg DOC L}^{-1}$ ), but peaked later in the 435 season, around December 2022, whereas there was little temporal variability of DOC in the ditch (Fig. 8H). 436 Dissolved total Fe displayed the same temporal trend as DOC (Fig. 8I) but was higher groundwater ( $1916 \pm 163$  437  $\mu\text{g Fe L}^{-1}$ ) compared to the ditch ( $98 \pm 95 \mu\text{g Fe L}^{-1}$ ). The geochemical parameters of groundwater and ditch 438 water point to different mechanisms regulating especially elements related to peat decomposition and possibly 439 plant uptake, where groundwater was more dynamic over time than ditch water. Generally, there were no 440 systematic spatial pattern of groundwater chemistry across the transect.

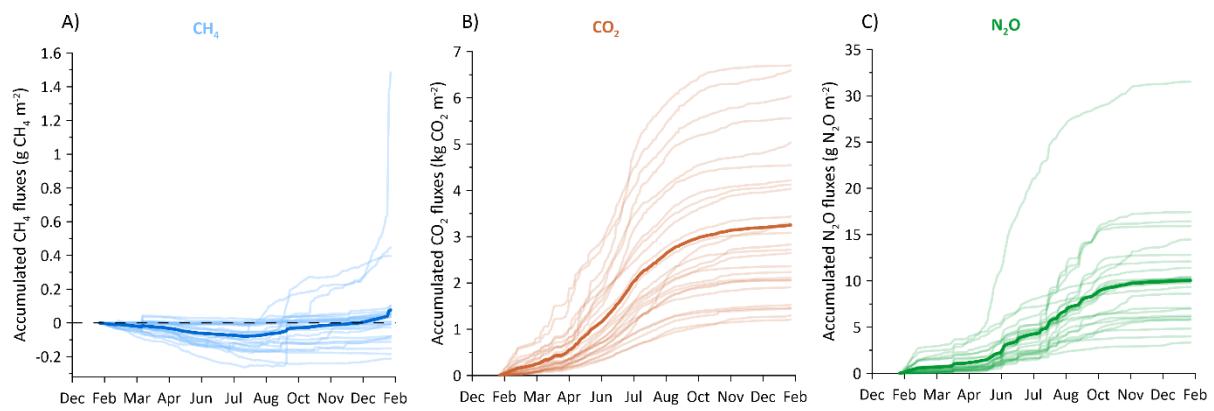


441

442 **Figure 8** Groundwater (brown closed circles) and ditch water (closed blue diamonds) chemistry at Vejrumbro for the  
443 period June 2022 to February 2023 for A) pH, B) Electric conductivity and dissolved C) total phosphor (P), D) total  
444 nitrogen (N), E) nitrate ( $\text{NO}_3^-$ ), F) ammonium ( $\text{NH}_4^+$ ), G) sulfate ( $\text{SO}_4^{2-}$ ), H) organic carbon and I) total iron (Fe).  
445 Values for organic soils are site means with error bars showing the standard error of the mean (N=6 per sampling  
446 date).

447 **3.7 Net soil and ditch  $\text{CO}_2$ ,  $\text{CH}_4$  and  $\text{N}_2\text{O}$  fluxes**

448 **3.7.1 Spatial variation of net soil  $\text{CO}_2$ ,  $\text{CH}_4$  and  $\text{N}_2\text{O}$  fluxes**



449

450 **Figure 9:** Cumulative fluxes of A)  $\text{CH}_4$ , B)  $\text{CO}_2$ , and C)  $\text{N}_2\text{O}$  for 26 individual collars along the SkyLine2D transect.  
451 Units for  $\text{CH}_4$  and  $\text{N}_2\text{O}$  are in  $\text{g CH}_4/\text{N}_2\text{O} \text{ m}^{-2}$  and for  $\text{CO}_2$  in  $\text{kg CO}_2 \text{ m}^{-2}$ . The cumulative fluxes represent the raw  
452 dataset. The ditch data was excluded. Site average is shown as thick lines.

453 Within the transect, cumulative CH<sub>4</sub> fluxes over the study period (360 days) varied between -0.21 to 1.48 g CH<sub>4</sub>  
454 m<sup>-2</sup> over the study period, with a site average ( $\pm$ SE) cumulative flux of 0.07 $\pm$ 0.06 g CH<sub>4</sub> m<sup>-2</sup> (Fig. 3 and 9A).  
455 Out of the 26 collars, excluding the ditch collar, 11 displayed a net uptake over the measurement period and the  
456 remaining were small net emitters (Fig. 3 and 9A). There was generally little spatial variation in the absolute  
457 CH<sub>4</sub> fluxes among the soil collars, but three collars (11, 12 and 15) showed increasing net positive cumulative  
458 fluxes towards the ditch (Fig. 3). The low spatial and similar temporal variation between collars indicate both  
459 hydrological indicators of SWC and WTD are poor predictors of CH<sub>4</sub> fluxes at this site. However, as we  
460 excluded plants from the collars we might have decreased the net emission of CH<sub>4</sub> directly by restricting gas  
461 transport in aerenchyma (Askaer et al. 2011; Vroom et al. 2022) and indirectly by potentially reducing plant  
462 carbon supply to methanogens. However, visible inspection at the site confirmed lateral root growth from  
463 vegetation adjacent to the collar. This could indicate that plant derived C and N was still available for microbes  
464 underneath the collars, but the impact on gas transport is uncertain. However, we did not excavate roots during  
465 the study to avoid excessive disturbance. Furthermore, considering that the WTD in the growing season was  
466 mostly 20-40 cm below terrain the potential for CH<sub>4</sub> production in the topsoil would limited (Koch et al. 2023).  
467 Also, the lack of consistent hot moments of CH<sub>4</sub> emissions and low cumulative emissions from the soil despite  
468 hydrological conditions in the subsoil being conducive for CH<sub>4</sub> production could indicate that redox potential is  
469 elevated due to presence of other electron acceptors. The presence of both free NO<sub>3</sub><sup>-</sup>, SO<sub>4</sub><sup>2-</sup>, Fe (Fig. 8E, G, I) in  
470 the groundwater could indicate that there are alternative electron acceptors that prevent lowering of the redox  
471 status of the soil and hence suppresses CH<sub>4</sub> production.

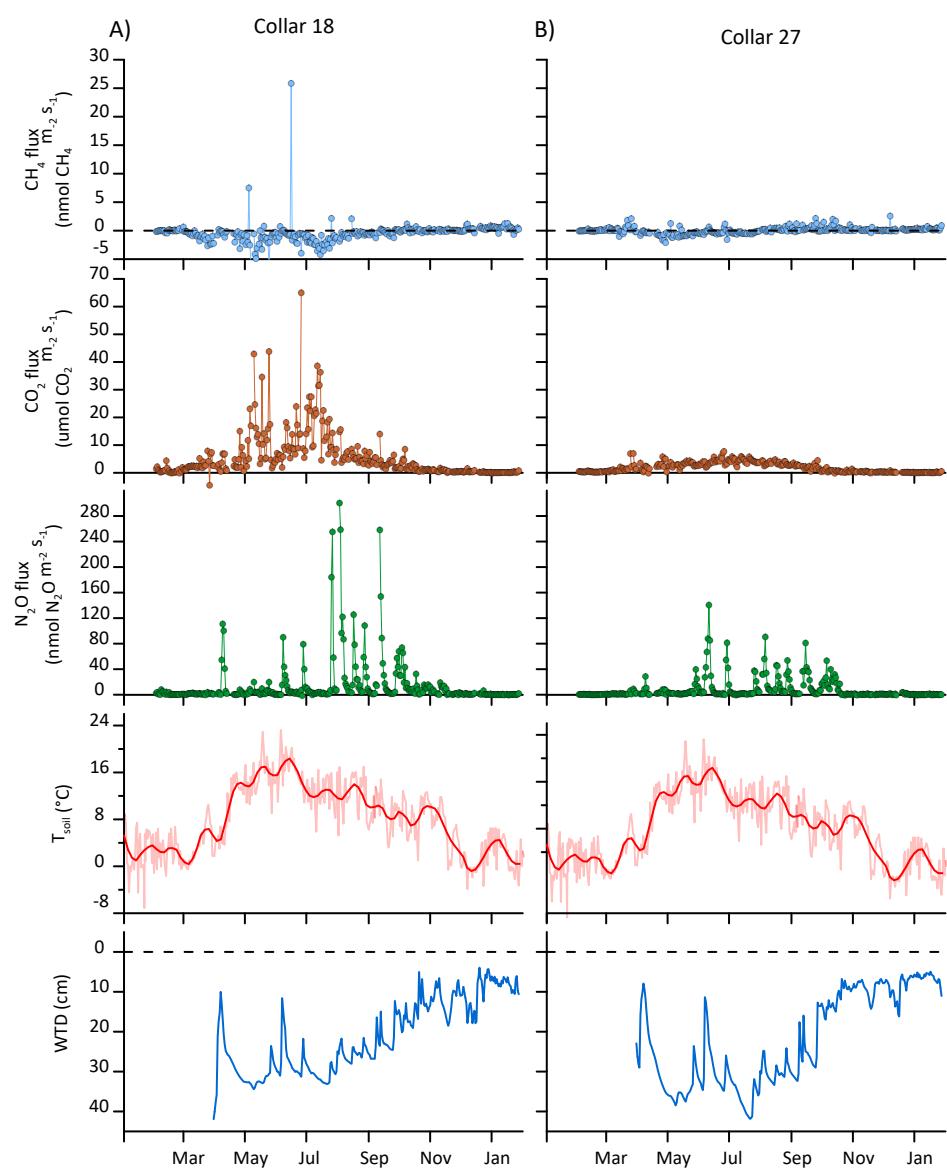
472 The CO<sub>2</sub> effluxes displayed tremendous spatial variation across the 24-meter transect (Fig. 3 and 9B) and  
473 measurements indicated that the drained organic soil was a net source of CO<sub>2</sub>, with cumulative fluxes over the  
474 study period ranging between 1214 – 6740 g CO<sub>2</sub> m<sup>-2</sup>, and a site average ( $\pm$ SE) of 3269 $\pm$ 328 g CO<sub>2</sub> m<sup>-2</sup>, over  
475 the study period of 360 days (Fig. 3 and 9B). There was no apparent relation between the magnitude of  
476 cumulative CO<sub>2</sub> efflux to the position along the transect and average WTD (Fig. 3). The cumulative net soil CO<sub>2</sub>  
477 emission is equal to 8.9 tCO<sub>2</sub>-C ha<sup>-1</sup> y<sup>-1</sup> (range of 3.3 to 18 tCO<sub>2</sub>-C ha<sup>-1</sup> y<sup>-1</sup> across the transect) and compares  
478 well to estimates of annual soil C loss (8.8 tCO<sub>2</sub>-C ha<sup>-1</sup> y<sup>-1</sup>) from a drained unfertilized grassland on organic soil  
479 in Denmark (Kandel et al. 2018) as well as annual carbon budgets of similar Danish, British and German  
480 wetlands (Tiemeyer et al. 2020; Evans et al. 2021; Koch et al. 2023).

481 Similarly, the site was overall a net source of N<sub>2</sub>O, with cumulative fluxes ranging between 3.3 – 32 g N<sub>2</sub>O m<sup>-2</sup>,  
482 with a site average ( $\pm$ SE) of 10.1 $\pm$ 1.1 g N<sub>2</sub>O m<sup>-2</sup> (Fig. 3 and 9C) over the study period (360 days). Thus, there is  
483 a 10-fold difference between minimum and maximum cumulative N<sub>2</sub>O fluxes within the transect, without any  
484 apparent relation to the position along the transect and WTD. The highest cumulative N<sub>2</sub>O fluxes occurred at  
485 collar 8 situated close to the ditch (Fig. 3). The site average cumulative N<sub>2</sub>O emission is equivalent to a net N  
486 loss from N<sub>2</sub>O emission alone of 64 kg N ha<sup>-1</sup> y<sup>-1</sup>, was very high and exceeding previously reported fluxes from  
487 this site (1.5 – 2.1 g N<sub>2</sub>O m<sup>-2</sup> y<sup>-1</sup>) (Nielsen et al., 2024) and German organic soils (0.04 – 6.3 g N<sub>2</sub>O m<sup>-2</sup> y<sup>-1</sup> for  
488 grassland and cropland land uses) (Tiemeyer et al. 2020). The high N<sub>2</sub>O emission from this site during the  
489 measurement period indicate that N<sub>2</sub>O may in fact dominate the GWP budget at this site had gross primary  
490 production been included in the measurements. It is important to reiterate here that the flux measurements of  
491 this study were done on bare soil whereas the studies referenced above included vegetation.

492 The high N<sub>2</sub>O fluxes may be a result of high rates of denitrification in the subsoil from either *in situ* produced  
 493 NO<sub>3</sub><sup>-</sup> from peat decomposition or as NO<sub>3</sub><sup>-</sup>-enriched agricultural runoff from the surrounding intensively  
 494 cultivated areas, which was not affecting groundwater NO<sub>3</sub><sup>-</sup> concentration in the center of the wetland with  
 495 lower N<sub>2</sub>O (Nielsen et al. 2024). The groundwater enters the northern peripheral zone of the wetland at  
 496 Vejrumbro coinciding with the position of the measurement transect. The highest NO<sub>3</sub><sup>-</sup> concentrations in  
 497 groundwater at the SkyLine2D transect corresponded roughly with highest N<sub>2</sub>O emission during summer and  
 498 early autumn (Fig. 8D-F and Fig. 12D), but the frequency of water sampling was too low to fully link  
 499 groundwater NO<sub>3</sub><sup>-</sup> temporal dynamics to N<sub>2</sub>O emissions.

500 **3.7.2 Temporal variability of net soil CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O fluxes**

501 **3.7.2.1 Time series of raw data of net soil CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O fluxes**



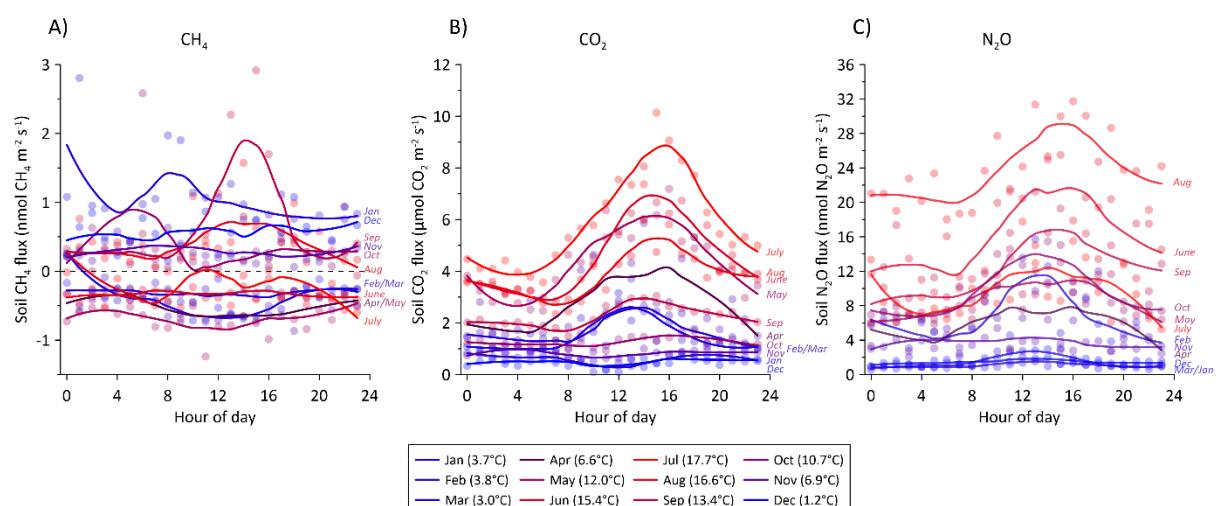
502  
 503 **Figure 10:** Examples of daily average time series of CH<sub>4</sub>, CO<sub>2</sub> and N<sub>2</sub>O fluxes for collars 18 and 27 at the SkyLine2D  
 504 transect in Vejrumbro, soil temperature (ST) in celsius (°C) and groundwater table depth (WTD) in cm below terrain  
 505 is shown in two lower panels.

506 With the high frequency of GHG flux measurements (on average 5 measurements per day per collar) it was  
 507 possible to observe short term flux phenomena that in most studies deploying manual chambers are missed or if  
 508 captured can lead to biased conclusions on flux magnitudes. For example, in most of the measurement points,  
 509 CH<sub>4</sub> fluxes were generally near zero, but occasionally displayed elevated net emission for short periods even in  
 510 periods with deeper WTD (Fig. 10A) for most chambers (see supplementary Fig. S5). This flux dynamic might  
 511 be related to episodic release of accumulated CH<sub>4</sub> from deeper soil layers that are not fully oxidized in the  
 512 aerated root zone and that were not released through plants (Askaer et al. 2011). As plants were not included in  
 513 the collars these bursts cannot be attributed to plant emission pathways.

514 Generally, it was observed that soil CO<sub>2</sub> fluxes increased over the season with increasing temperature. However,  
 515 for some collars displayed rapid bursts of CO<sub>2</sub> emissions (example in Fig. 10A), while other collars at the same  
 516 period did not display this behaviour (Fig. 10B). This dynamic points to different emission pathways from the  
 517 soil not related to plant mediated transport. Thus, while we purposely omitted aboveground autotrophic  
 518 respiration by clipping the vegetation, it cannot be ruled out that living roots inhabited the soil below the  
 519 chambers and hence contributed to the observed CO<sub>2</sub> emission rates.

520 For N<sub>2</sub>O, the spatiotemporal pattern was even more pronounced than for CO<sub>2</sub>, with N<sub>2</sub>O primarily emitted in  
 521 bursts related to rapidly increasing or decreasing WTD that coincided with precipitation events. In drier periods  
 522 with deeper WTD and little fluctuations, N<sub>2</sub>O fluxes quickly dropped to near zero (Fig. 10A and B). Despite  
 523 N<sub>2</sub>O being emitted in similar temporal patterns across the site, the magnitude of the N<sub>2</sub>O peaks were not similar  
 524 across the transect (Fig. 10 and supplementary Fig. S5). Hence, the majority of N<sub>2</sub>O is emitted in hot moments  
 525 driven by fluctuations in WTD mainly (Fig. 10) as it has also been shown in other drained temperate peatland  
 526 soils (Anthony and Silver 2023).

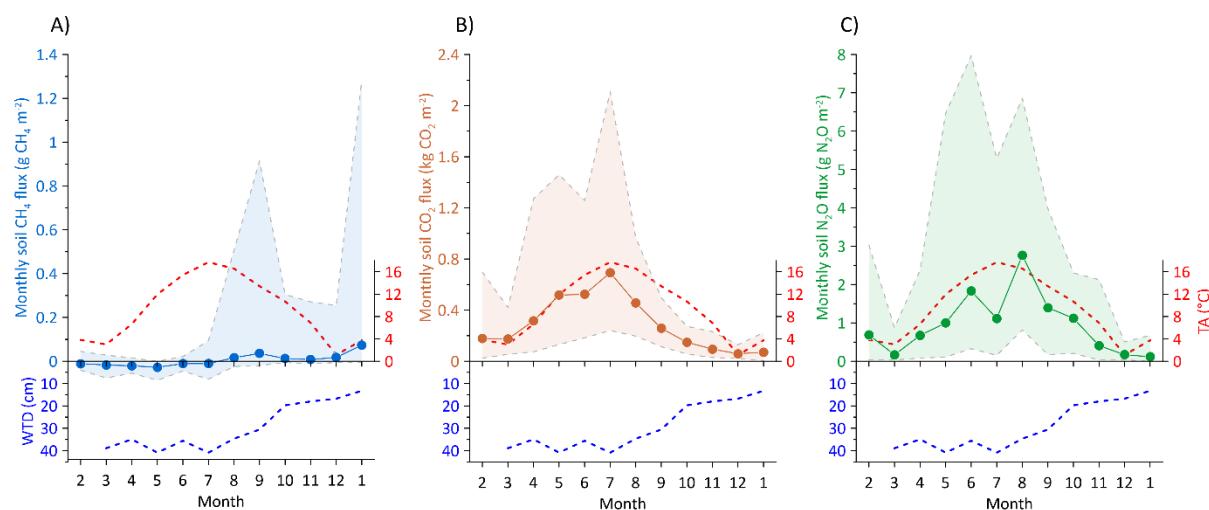
### 527 3.7.2.2 Diurnal variation of net soil CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O fluxes



528  
 529 **Figure 11: Average hourly flux for all soil collars of A) CH<sub>4</sub>, B) CO<sub>2</sub>, and C) N<sub>2</sub>O during a 24 hour period. The**  
 530 **diurnal variation is split between each month during the 2022-2023 measurement period. The fluxes were assigned**  
 531 **the hour of measurement during the day and averaged per month. Color shade between blue and red corresponds to**  
 532 **average air temperature for the specific month shown in parenthesis in the figure legend. Solid lines are loess fits for**  
 533 **visualization of the diurnal variation in each month.**

534 With the SkyLine2D system we observed a clear diurnal cycle for CO<sub>2</sub> and N<sub>2</sub>O fluxes, but not for CH<sub>4</sub> (Fig. 535 11A-C). The lack of diurnal variability of CH<sub>4</sub> fluxes could also be due the removal of plants from the collars 536 that would have facilitated light-driven fluxes (Askaer et al. 2011). The amplitude of diurnal variability 537 increased with higher air temperature for CO<sub>2</sub> (Fig. 11B) and partly for N<sub>2</sub>O (Fig. 11C). The month of July was 538 an exception as it resembled the pattern observed in May although the July soil temperature was about 5°C 539 higher (Table 2). The lower N<sub>2</sub>O fluxes observed in July can be attributed to lower and more constant WTD in 540 July compared to May, June and September across the transect (Fig. 6). Diurnal variability of soil CO<sub>2</sub> fluxes 541 are well known and can be related to both increased heterotrophic respiration during the warmer day and 542 autotrophic respiration in response to photosynthesis. Previously, similar diurnal patterns of N<sub>2</sub>O emissions were 543 observed in a Danish fen (Jørgensen et al. 2012).

544 **3.7.2.3 Monthly variability of net soil GHG fluxes**



545 **Figure 12: Monthly summed soil fluxes of A) CH<sub>4</sub> in g CH<sub>4</sub> m<sup>-2</sup>, B) CO<sub>2</sub> in kg CO<sub>2</sub> m<sup>-2</sup>, and C) N<sub>2</sub>O in g N<sub>2</sub>O m<sup>-2</sup> for 546 all organic soil collars. Shaded areas for CH<sub>4</sub>, CO<sub>2</sub> and N<sub>2</sub>O graphs represent the maximum and minimum monthly 547 average fluxes. Blue dashed line below CH<sub>4</sub>, CO<sub>2</sub> and N<sub>2</sub>O represent the measured monthly average transect 548 groundwater table depth (WTD) in cm below terrain. Red dashed line shows the monthly average air temperature 549 (TA). 550**

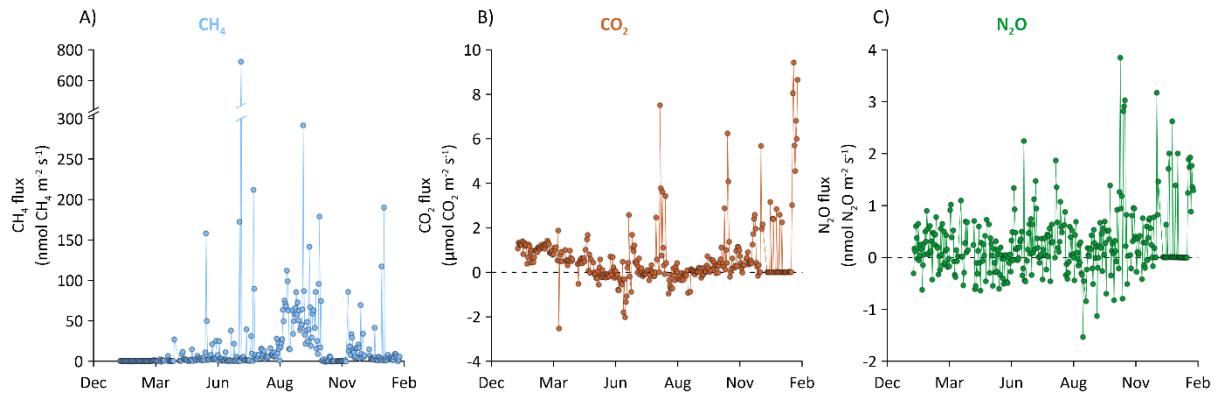
551 The average soil GHG fluxes for all collars were summed to monthly site sums to illustrate long term drivers on 552 the flux magnitude. Overall, monthly sums of CO<sub>2</sub> and N<sub>2</sub>O emissions increase with temperature and fluxes are 553 highest under deeper WTD, but CH<sub>4</sub> net fluxes were less responsive to long term changes in both temperature 554 and hydrology (Fig. 12A-C). Net uptake increased slightly with increasing temperature and lower WTD during 555 the spring and summer. With increasing water table and high temperatures in August the site turned into a small 556 net CH<sub>4</sub> source continuing in fall and winter (Fig. 12A).

557 For CO<sub>2</sub> the seasonal variation was pronounced and closely followed soil temperature until peak values in July 558 for both site average, minimum and maximum fluxes, respectively (Fig. 12B). From July to August, it was 559 observed that WTD at the site began to increase again and CO<sub>2</sub> fluxes departed from the close relation to soil 560 temperature, indicating an inhibitory role of the WTD in this period, but reaching minimum fluxes in December, 561 corresponding to the wettest and coldest month (Fig. 12B).

562 Similarly,  $\text{N}_2\text{O}$  fluxes increased with soil temperature reaching peak monthly values in August, corresponding to  
 563 the period of the year with highest soil temperature and increasing WTD (Fig. 12C). This supports the  
 564 promoting role of soil water saturation on the production of  $\text{N}_2\text{O}$  when temperature is favourable for  
 565 denitrification.  $\text{N}_2\text{O}$  fluxes reached minimum values in December when WTD and ST were lowest (Fig. 12C).

### 566 3.7.3 Ditch $\text{CO}_2$ , $\text{CH}_4$ and $\text{N}_2\text{O}$ fluxes

#### 567 3.7.3.1 Time series of raw data of ditch $\text{CO}_2$ , $\text{CH}_4$ and $\text{N}_2\text{O}$ fluxes



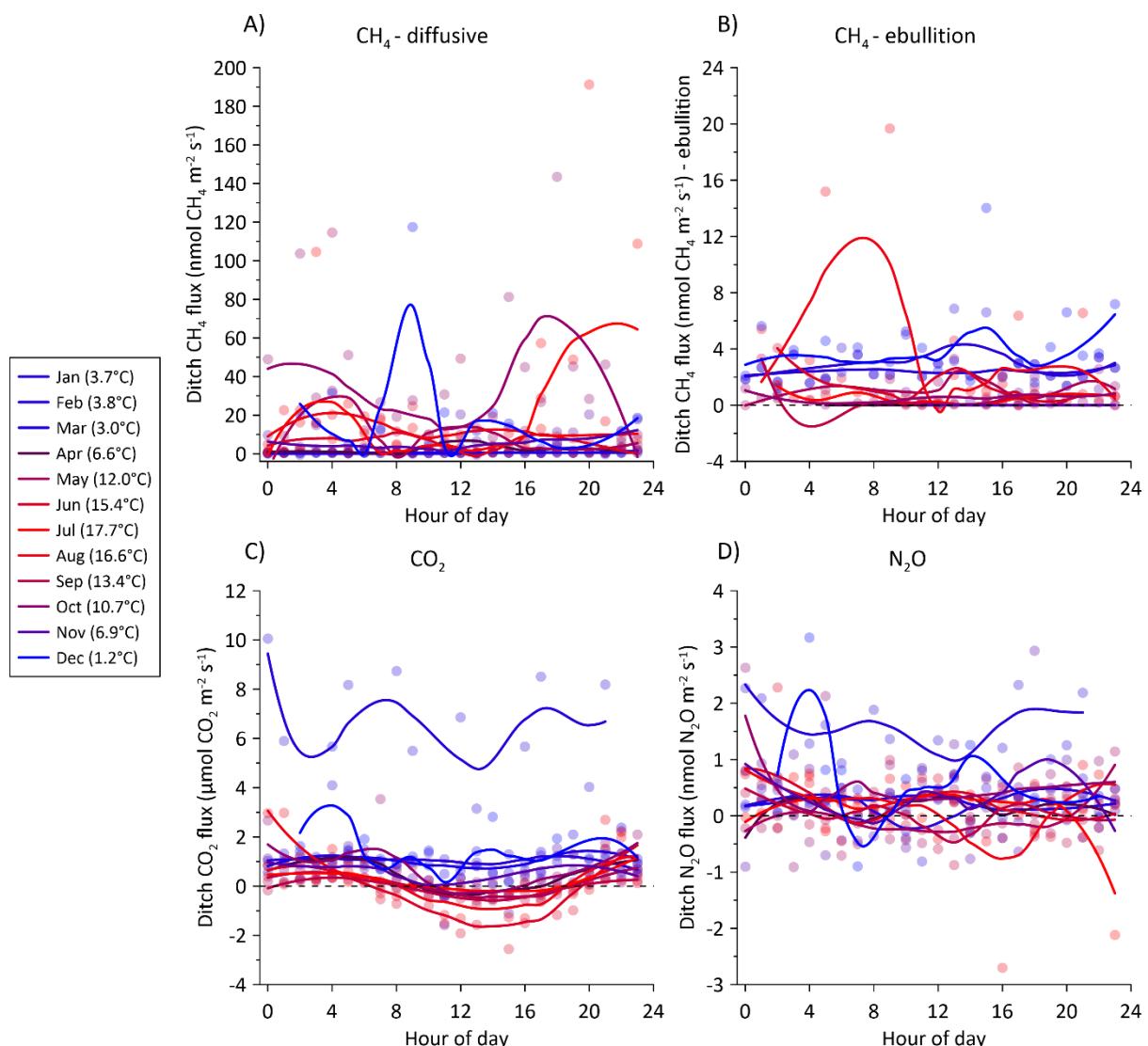
568  
 569 **Figure 13: Daily average time series of net ditch total A)  $\text{CH}_4$  (diffusion and ebullition), B)  $\text{CO}_2$ , and C)  $\text{N}_2\text{O}$  fluxes at**  
 570 **the Vejrumbro site.**

571 Common for all three gases is that ditch emissions are dynamic and net fluxes change from zero to large net  
 572 positive or negative fluxes within hours or days (Fig. 13A-C). Compared to net soil  $\text{CH}_4$  fluxes the ditch can be  
 573 considered an emission hotspot at the site (sum of diffusive and ebullition:  $8.3 \text{ g CH}_4 \text{ m}^{-2} \text{ y}^{-1}$ ), but fluxes are  
 574 lower than earlier reports for ditches in other drained wetlands (between  $0.1 - 44.3 \text{ g CH}_4 \text{ m}^{-2} \text{ y}^{-1}$ ) (Peacock et  
 575 al., 2021). Methane is most dynamic with maximum diffusive flux close to  $700 \text{ nmol CH}_4 \text{ m}^{-2} \text{ s}^{-1}$  and there was  
 576 a tendency toward consistently higher net  $\text{CH}_4$  emission from August to September, becoming close to zero in  
 577 colder seasons (Fig. 13A). Ebullition of  $\text{CH}_4$  did occur occasionally in the ditch, e.g. about 19.3% of flux  
 578 measurements for the ditch was comprised of ebullitions but constituted on average only 2.9% of the total  $\text{CH}_4$   
 579 emission ( $0.24 \text{ g CH}_4 \text{ m}^{-2} \text{ y}^{-1}$ ) from the ditch which is lower, but in the same range as a recent estimate from a  
 580 ditch in a similar drained German peatland (Köhn et al. 2021). According to the flux calculation methodology,  
 581 flux separation and extrapolation to daily sums, diffusive fluxes dominated ( $6.56 \text{ g CH}_4 \text{ m}^{-2} \text{ y}^{-1}$ ). However, it  
 582 cannot be ruled out that the classification as diffusive flux may in fact be ebullition by nature. It has been  
 583 suggested that microbubbles resulting from mass transport can resemble diffusive fluxes in a chamber making it  
 584 difficult, if not impossible, to fully separate the two emission mechanisms in a continuous time series if  
 585 headspace  $\text{CH}_4$  concentrations do not abruptly increase (Prairie and del Giorgio 2013), such as in the example  
 586 shown in Fig. S4.

587 For  $\text{CO}_2$ , there was a general tendency towards lower fluxes during the summer months and fluxes increased in  
 588 magnitude and variability towards the end of the study period (Fig. 13B). For  $\text{N}_2\text{O}$ , the fluxes fluctuated around  
 589 zero for most of the study period, except towards the end (December and January) where net fluxes became  
 590 positive (Fig. 13C).

591 Compared to the net soil  $\text{N}_2\text{O}$  and  $\text{CO}_2$  fluxes the ditch fluxes of these gases are low showing that the ditch is  
 592 not contributing significantly to the  $\text{CO}_2$  and  $\text{N}_2\text{O}$  budget at this site.  
 593 Per square meter, the ditch emitted less  $\text{N}_2\text{O}$  ( $0.41 \text{ g N}_2\text{O m}^{-2}$  or  $2.6 \text{ kg N}_2\text{O-N ha}^{-1} \text{ y}^{-1}$ ) and  $\text{CO}_2$  ( $961 \text{ g CO}_2 \text{ m}^{-2}$   
 594  $\text{y}^{-1}$  or  $2.6 \text{ tCO}_2\text{-C ha}^{-1} \text{ y}^{-1}$ ) than the organic soil, but was a hotspot of  $\text{CH}_4$  emission ( $8.4 \text{ g CH}_4 \text{ m}^{-2} \text{ y}^{-1}$  or  $63 \text{ kg}$   
 595  $\text{CH}_4\text{-C ha}^{-1} \text{ y}^{-1}$ ) during the measurement period. Although these emissions estimates are lower than previously  
 596 reported for ditches in organic soil (up to  $44 \text{ g CH}_4 \text{ m}^{-2} \text{ y}^{-1}$ ) (Peacock et al. 2021). For the ditch  $\text{CH}_4$  budget,  
 597 ebullition only constitutes 2.9% of net  $\text{CH}_4$  emissions during the study period. This proportion may be  
 598 underestimated as the count of ebullition events may have been underestimated (Prairie and del Giorgio 2013).

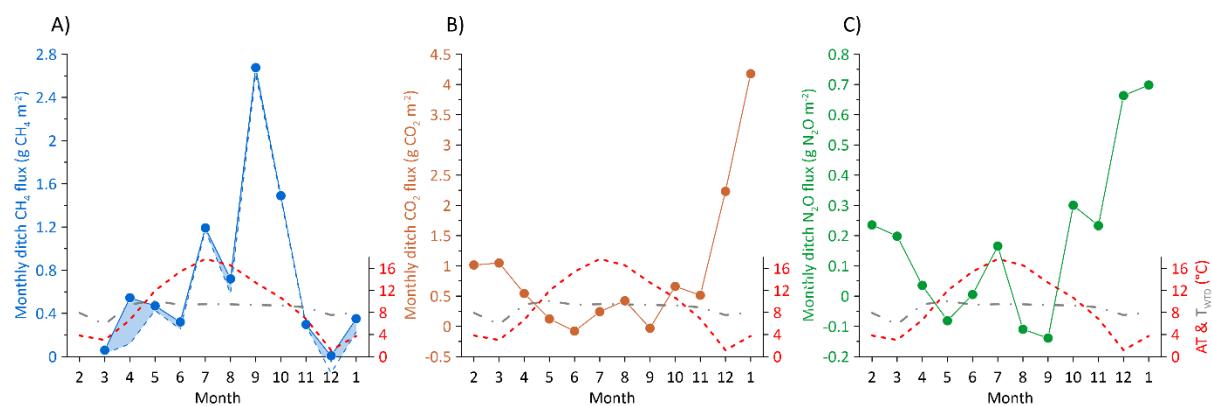
599 **3.7.3.2 Diurnal variability in ditch fluxes**



600  
 601 **Figure 14: Average hourly fluxes for the ditch collar of A) diffusive  $\text{CH}_4$  fluxes, B)  $\text{CH}_4$  ebullition fluxes, C)  $\text{CO}_2$ , and**  
 602 **C)  $\text{N}_2\text{O}$  during a 24 hour period. The fluxes were assigned the hour of measurement during the day and averaged per**  
 603 **month. The diurnal variation is split between each month during the 2022-2023 measurement period. Color shade**  
 604 **between blue and red corresponds to average air temperature for the specific month shown in parenthesis in the**  
 605 **figure legend. Solid lines are loess fits for visualization of the diurnal variation in each month. Note different axes.**

606 For CH<sub>4</sub> fluxes, both diffusive and ebullition, there was no clear diurnal variability in any month (Fig. 14A and  
 607 B). This is expected for ebullition emissions which is known to be erratic without any clear diurnality (Wik et al.  
 608 2016; Sø et al. 2023). For net CO<sub>2</sub> fluxes from the ditch there was no diurnal variability in colder seasons (Jan,  
 609 Feb, Mar, Nov and Dec), but consistent positive net CO<sub>2</sub> efflux (Fig. 14C). Diurnal patterns became clearer with  
 610 higher temperatures from May to October (Fig. 14C) and in this period CO<sub>2</sub> fluxes decreased during the day to  
 611 sometimes reach net negative fluxes (net uptake of CO<sub>2</sub>) during and after midday (Fig. 14C), although the net  
 612 emissions were also observed in the daytime period (Fig. 14C). The net negative fluxes can likely be explained  
 613 by photosynthetic activity of aquatic plants on the surface of the ditch or by algae in the water column which  
 614 was measured due to the transparency of the chamber. Using an opaque chamber instead would likely have  
 615 resulted in different net CO<sub>2</sub> efflux in daytime. For N<sub>2</sub>O, the same pattern as for CH<sub>4</sub> was observed, where flux  
 616 magnitude across the day fluctuated around zero, except for January where N<sub>2</sub>O fluxes were consistently above  
 617 zero (Fig. 14D).

### 618 3.7.3.3 Monthly variability in ditch fluxes



619  
 620 **Figure 15: Monthly summed ditch fluxes of A) CH<sub>4</sub> in g CH<sub>4</sub> m<sup>-2</sup>, B) CO<sub>2</sub> in g CO<sub>2</sub> m<sup>-2</sup> and C) N<sub>2</sub>O in g N<sub>2</sub>O m<sup>-2</sup>. In A)**

621 the blue dashed line is the contribution of diffusive fluxes and the shaded blue area between the full and dashed blue

622 lines represent the monthly contribution of ebullition to the total flux. Red and grey dashed lines show the monthly

623 average air (AT) and groundwater temperature (T<sub>WTD</sub>) in °C, respectively.

624 The monthly sums of CH<sub>4</sub> tend to increase with air temperature, although peak CH<sub>4</sub> emissions (September)  
 625 occurred after air temperature peak (July) (Fig. 15A). Diffusive fluxes comprised the major emission pathway of  
 626 CH<sub>4</sub> in the ditch (between 21% - 99%), with the contribution from ebullition being highest in March (55%) and  
 627 April (78%) (Fig. 15A). Water temperature in the ditch was relatively stable throughout the year, varying  
 628 between 5.8 – 10.1°C being highest from April to November and lowest from December to March. However,  
 629 there is little indication of a direct relation between ditch water temperature and net GHG fluxes (Fig. 15A-C).  
 630 For CO<sub>2</sub> and N<sub>2</sub>O, the seasonal pattern is reversed with lowest fluxes during the warmest periods, approaching  
 631 net zero or even net negative fluxes (Fig. 15B and C).

### 632 3.7.4 Estimate of the annual soil and ditch GHG budgets at the Vejrumbro location

633 The annual GHG budgets for N<sub>2</sub>O, CO<sub>2</sub> and CH<sub>4</sub> were adjusted from the cumulated values by multiplying with a  
634 factor of 365/360. It showed that for the drained organic soil its gaseous carbon loss was mostly as CO<sub>2</sub>, while  
635 CH<sub>4</sub> played a negligible role in the C cycle and consequently also for global warming potential (GWP) budget.

## 636 **4 Data availability**

637 Data for this publication is available for download via  
638 <https://dataVERSE.deic.dk/previewurl.xhtml?token=abda26d4-a430-4830-ad30-fbf5ff1d352e> (Skov Nielsen et al.  
639 2025).

## 640 **5 Conclusion**

641 The dataset presented here is unique for temperate fens and demonstrate the advantage of using automated GHG  
642 measurements systems to resolve temporal and spatial patterns of GHG dynamics in high detail. The dataset also  
643 demonstrate how especially temporal variation of soil hydrology and temperature is linked to the dynamics of  
644 fluxes and highlight that spatial variability in hydrology and temperatures not necessarily is the best predictor of  
645 flux magnitudes within the site. The cause for the spatial variability of GHG fluxes remains unresolved and do  
646 not clearly link directly to either WTD, soil temperature and soil/groundwater chemical parameters.

647 Interestingly it appears that the temporal variability of GHG fluxes across the transect is lower than the spatial  
648 variation.

649 The data only represents one full year in 2022-2023 and hence must be considered specific for this period. It  
650 must therefore be expected that the annual budget of all GHG's in other years will be different due to other  
651 climatic and hydrological conditions.

652 The initial harvest and herbicide application represent ecosystem disturbances that potentially can alter soil  
653 biogeochemistry, but they were done months prior to the start of the flux measurements and hence the direct  
654 effect of herbicide would be minimal. The continued plant removal from inside collars was necessary for the  
655 flux measurements with the consequence that our fluxes may only be regarded as net soil GHG fluxes, and not  
656 as being representative of the net ecosystem exchange. Excluding the influence of vegetation have influenced  
657 the measured fluxes of soil respiration (e.g. excluding root exudates etc.) and reduced plant mediated CH<sub>4</sub> and  
658 N<sub>2</sub>O emissions and lowered most likely also reduced interannual variability. However, the data set represents a  
659 unique ability to continue to develop models that predict the soil GHG fluxes in response to soil temperature and  
660 hydrology (WTD) that can aid in prediction of reliable budgets for sites.

661 The measurements of the soil GHG fluxes show that the magnitude of annual cumulative CO<sub>2</sub> fluxes are in the  
662 same range as in other studies of temperate fens, and that temporal variability are largely governed by the  
663 seasonality of WTD and ST. However, spatial variation of cumulative fluxes for all GHG were not directly  
664 related to WTD levels, contradicting the general assumption that WTD is the primary driver of GHG emissions.  
665 Cumulative soil N<sub>2</sub>O fluxes exceed what has been previously reported for temperate fens, but show similar  
666 seasonal regulation by ST. However, in contrast to soil CO<sub>2</sub> fluxes, soil N<sub>2</sub>O is emitted largely in pulses related  
667 to rapid fluctuations of WTD that increase in size with temperature. These measurements therefore point to an  
668 important, but difficult to capture dynamic of N<sub>2</sub>O in peatlands where hot moments during the warm periods  
669 determine most of the annual emissions. A likely cause for the high soil N<sub>2</sub>O emissions could be a combination  
670 of leaching of inorganic nitrogen from surrounding agricultural fields and release of organic N from the

671 decomposing peat. The site was during the measurement period an insignificant source of soil CH<sub>4</sub>, which is  
672 likely due to the well-drained summer period, a cold wet winter and presence of the major electron acceptors  
673 (NO<sub>3</sub><sup>-</sup>, SO<sub>4</sub><sup>2-</sup> and Fe<sup>3+</sup>), providing suboptimal conditions for CH<sub>4</sub> production. However, it cannot be ruled out  
674 that the vegetation removal impeded CH<sub>4</sub> emissions, as we effectively restricted plant mediated CH<sub>4</sub> emissions.  
675 Therefore, caution should be taken when comparing the CH<sub>4</sub> flux data to other drained peatlands. Soil CO<sub>2</sub> and  
676 N<sub>2</sub>O fluxes both showed diurnal variability with higher fluxes during midday where the amplitude between  
677 night and day was augmented with ST. This was not observed for soil CH<sub>4</sub> fluxes. The ditch at the site was a net  
678 source of both N<sub>2</sub>O and CO<sub>2</sub>, but at rates 27 and 4 times lower than the soil GHG fluxes respectively. However,  
679 the ditch acted as a CH<sub>4</sub> source mostly comprised of diffusive emissions from the water surface, but with  
680 observations of ebullition.

681 We wish to publish this dataset to the research community with the intention that experimentalists and modellers  
682 can use the data to test hypothesis on basic hydrological and thermal regulation of GHG fluxes and develop  
683 models to predict spatiotemporal variability of the GHG fluxes.

#### 684 Competing interests

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

#### 686 Author contributions

687 JRC, PEL and KSL designed the experiment and carried them out. ASN performed flux calculation and quality  
688 checking. RJP and PEL installed the equipment for groundwater measurements. All authors contributed to  
689 writing of this manuscript.

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