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Methane emissions from fens in Alberta's boreal region: Reference data for functional evaluation of restoration outcomes

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Abstract:	<p>The aim of the study was to document methane (CH₄) dynamics from fen ecosystems in the Athabasca Oil Sands Region (AOSR) in northern Alberta to create a reference database for evaluation of peatland restoration and reclamation projects in the region. The study included three types of fens commonly occurring in this region: poor fen (open and treed), moderately-rich treed fen, and open saline fen. We quantified CH₄ fluxes, pore water concentration (PW[CH₄]), and production potential together with ecohydrological variables that may influence CH₄ dynamics over four growing seasons. Mean (standard deviation) fluxes for open and treed poor fen (99.8 (269.7) and 68.3 (118.8) mg CH₄ m⁻² d⁻¹, respectively) were higher than for treed rich (32.8 (63.7) mg CH₄ m⁻² d⁻¹) and open saline fens (34.6 (91.3) mg CH₄ m⁻² d⁻¹). The total growing season CH₄ emissions from these fens ranged between 3.7 and 11.3 g CH₄ m⁻². Methane production potential varied from 0.1 (0.1) μmol CH₄ g peat⁻¹ d⁻¹ at the saline fen to 4.6 (0.8) μmol CH₄ g peat⁻¹ d⁻¹ at the treed rich fen. The variability of CH₄ fluxes and pore water concentration between study sites and years was mostly controlled water table (WT) and soil temperature indicating that these variables should be used to assess the expected CH₄ flux in peatland reclamation projects. Large inter-annual variability in CH₄ flux illustrates the importance of multi-year records for data used in functional</p>	

	evaluation of restoration outcomes.
<p>Response to Reviewers:</p>	<p>Thank-you for the opportunity to revise our manuscript. The comments from the reviewer have helped to clarify many points in the manuscript. We present each reviewer comment, followed by our response. Line numbers in our response refer to those in the revised version.</p> <p>Reviewer report: This report specifically examines methane dynamics in four different fen peatland ecosystem types in the Athabasca Oil Sands region, over four years. The principal objective was to establish "reference" conditions for methane efflux to be used as benchmarks for restoration; this goal appears to be appropriate for 'Wetland Ecology and Management'. The writing is concise and describes the data quite well. The figures and tables are straightforward, and there is enough detail to replicate the field monitoring protocols. The authors place their findings in the broader context of northern fen research (table 4). I offer minor comments by line number, below, and hope they are helpful with revision.</p> <p>Line 164: Might be good to specify that this work focuses on diffusive efflux, and does not reflect ebullition events. Response: A good suggestion and we have added the following sentence at the end of this paragraph to clarify what is included in the fluxes based on our quality control methods: "Therefore, results presented here largely represent diffusive and plant-mediated fluxes as sporadic ebullition events would have been removed from the data set based upon our quality control criteria; steady ebullition, if it occurred, would be included as it would result in a linear increase in concentration change over the chamber closure period". (lines 168-172)</p> <p>Line 174-175: It would be good to specify the micron size, or brand, of nylon screening. Response: The screening had a 250 µm mesh size and this has now been specified in the text (line 179).</p> <p>Line 188-189: It would be nice to briefly provide a little information on how the incubation headspace N2 was flushed (glovebox? Replacing headspace?). Response: The jars were prepared and sealed in a glovebox flushed with N2. We have clarified this in the text (line 193).</p> <p>Line 202-203: This is noted again on line 343, but it would be good to note right up front that since canopy is not captured in the 30 cm tall chambers, and yet tree roots are, ecosystem respiration is likely higher than what is represented by the plants present, and GEP is therefore likely to be an underestimation. Response: Since we only use GEP in our analysis of potential controls on CH4 flux (and not ecosystem respiration), we have chosen to only highlight here the underestimation of ecosystem GEP and not the effects of tree roots on respiration. Since including tree root respiration in our chamber measurements will not effect the GEP estimate directly, we do not want to confuse the reader by highlighting that the respiration measured is higher than that from only the ground layer plants as it is not relevant to the data used in further analysis. In short, we have modified the last sentence in this paragraph so that it now reads: "As these measurements included only the vegetation within the collar, they represent GEP of the understory vegetation only and do not include trees present at TPF and TRF and therefore underestimate total ecosystem GEP."</p> <p>Line 333: Would it be appropriate to briefly list the statistic used to establish if a value was truly an "outlier" (Cook's D statistic, or similar?)? Response: These values were greater than 2 orders of magnitude different than other samples from the same study site and so appeared as true outliers. We did not use a specific statistical test, but have now clarified how far outside the other replicates they lie in the text (line 340).</p> <p>Line 436: The authors introduce the "multifaceted role" that different plant functional groups play on CH4 efflux/oxidation; perhaps this could be a little more detailed here, coming back to aerenchymatous plants?</p>

Response: We have added the following sentence and associated reference here to provide more detail on plant roles in CH₄ cycling
“For example, while aerenchymatous plants have been shown to increase the transport of CH₄ from the soil to the atmosphere, in some cases, CH₄ emissions can be reduced due to greater oxidation in the rooting zone (Bhullar et al., 2013).” (lines 444-447)

Line 445: I think these references pertaining to trees venting CH₄ to the atmosphere are for tropical trees? I don't think black spruce or tamarack have pneumatophores or lenticels?

Response: More and more recent research is suggesting that even trees without pneumatophores or lenticels may vent methane from wetland soils, although little research specific to black spruce or tamarack has been conducted. While the Pangala reference is related to tropical wetlands, the Gauci reference actually refers to alders in a temperate fen. We also already acknowledge here that more research is needed in boreal peatlands to determine if trees are really playing an important role, so we feel this addresses the reviewer's concerns and have not made any further changes here.

Line 450-465: It seems there is something interesting going on with pore water [CH₄] in 2014, with a large spike that doesn't necessarily correspond with CH₄ efflux. Can the authors speculate as to what may be contributing to the spike in PW[CH₄] in that year?

Response: We did not specifically investigate what might cause this decoupling, but hypothesize that it could reflect the potentially long mean residence time of porewater CH₄ (months to years). So, we posit that the porewater pool lags hydrological conditions slightly and that the continuing increase in porewater concentration in 2014 may reflect increases in CH₄ production in 2013 and continued favourable conditions as most sites in 2014, allowing the pool to continue to grow, while falling water table would enhance oxidation and start to reduce emissions. We have added the following to this section:

“However, there appears to be some decoupling between CH₄ emissions and PW[CH₄] (Figure 3). The continuing increase in PW[CH₄] in 2014 while emissions declined compared to the previous year may reflect the fact that mean residence time of CH₄ in peat can be months to years (Strack and Waddington, 2008). Therefore, the dissolved CH₄ pool in 2014 potentially reflects the favourable CH₄ production conditions in 2013 when all sites had shallow water tables.” (lines 475-479).

Line 478: See comment above about apparent decoupling of PW and CH₄ efflux for some of the sites in 2014.

Response: We agree and have addressed this in detail in the previous comment. Here, we now say “In our study PW[CH₄] was generally linked to fluxes but showed less spatial and temporal variability”, adding the word “generally” to reflect that there is some level of decoupling.

[Click here to view linked References](#)

1 **Methane emissions from fens in Alberta's boreal region: Reference data for**
2 **functional evaluation of restoration outcomes**

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11 **Keywords:** CH₄, dissolved methane, ecological restoration, peatland, reclamation

12 **Abstract**

13 The aim of the study was to document methane (CH₄) dynamics from fen ecosystems in the
14 Athabasca Oil Sands Region (AOSR) in northern Alberta to create a reference database for
15 evaluation of peatland restoration and reclamation projects in the region. The study included three
16 types of fens commonly occurring in this region: poor fen (open and treed), moderately-rich treed
17 fen, and open saline fen. We quantified CH₄ fluxes, pore water concentration (PW[CH₄]), and
18 production potential together with ecohydrological variables that may influence CH₄ dynamics
19 over four growing seasons. Mean (standard deviation) fluxes for open and treed poor fen (99.8
20 (269.7) and 68.3 (118.8) mg CH₄ m⁻² d⁻¹, respectively) were higher than for treed rich (32.8 (63.7)
21 mg CH₄ m⁻² d⁻¹) and open saline fens (34.6 (91.3) mg CH₄ m⁻² d⁻¹). The total growing season CH₄
22 emissions from these fens ranged between 3.7 and 11.3 g CH₄ m⁻². Methane production potential
23 varied from 0.1 (0.1) μmol CH₄ g peat⁻¹ d⁻¹ at the saline fen to 4.6 (0.8) μmol CH₄ g peat⁻¹ d⁻¹ at
24 the treed rich fen. The variability of CH₄ fluxes and pore water concentration between study sites
25 and years was mostly controlled water table (WT) and soil temperature indicating that these
26 variables should be used to assess the expected CH₄ flux in peatland reclamation projects. Large
27 inter-annual variability in CH₄ flux illustrates the importance of multi-year records for data used
28 in functional evaluation of restoration outcomes.

29

30 **Introduction**

31 Northern peatlands play an important role in the global carbon cycle by acting as large soil carbon
32 stocks, contributing significant amounts of dissolved carbon to downstream ecosystems and
33 accounting for 5 – 10% of global CH₄ emissions (Blodau 2002). Boreal and subarctic peatlands
34 release 17 – 61 Tg of CH₄ per year and inter-annual variations in these emissions may contribute
35 to fluctuations in atmospheric CH₄ concentration (Bridgham et al. 2013). In the Athabasca Oil
36 Sands Region (AOSR) of Alberta, Canada, open pit mining for oil sands extraction has disturbed
37 895 km² boreal forest (Government of Alberta 2018), ~50% of which is covered by peatlands (Vitt
38 et al. 1996), with 90% of these peatlands being fens (Vitt et al. 2000). The Alberta government
39 requires land disturbed by oil sands extraction to be returned to equivalent land capability
40 (Province of Alberta 2018), with recently more focus placed on including peatlands in the post-
41 mining landscape (e.g., Daly et al. 2012). Therefore, understanding and quantifying key processes
42 in fen ecosystems in near-pristine condition is essential to develop reference baselines for future
43 evaluation of reclaimed landscapes (Nwaishi et al. 2015). This study focuses specifically on fen
44 CH₄ dynamics in the AOSR.

45 Methane is produced in soils under highly reducing conditions by methanogenic *Archaea*
46 (Rosenberry et al. 2006; Lai 2009). The saturated soil conditions in peatlands allow for CH₄
47 production; however, presence of alternative terminal electron acceptors (TEAs), such as NO₃⁻,
48 SO₄²⁻, Fe³⁺, and anaerobic bacteria that utilize them, can inhibit or reduce rates of methanogenesis
49 by creating conditions where the reaction is not thermodynamically favoured (Bridgham et al.
50 2013; Madigan et al. 2009; Minderlein & Blodau 2010). As fens receive water from ground and
51 surface water sources, concentrations of TEAs can be higher than in bogs, thereby reducing CH₄
52 production (Estop-Aragónés et al. 2013). Methane production is also dependent on organic matter
53 substrate quality, plant community composition and productivity (Bridgham et al. 2013; Tuittila
54 et al. 2000). The presence of highly-productive graminoids in many fens provides large quantities
55 of fresh substrate through root exudates and litter accumulation that should enhance CH₄
56 production (Strack et al. 2017).

57 The atmospheric flux of CH₄ from a peatland is dependent not only on production, but also on CH₄
58 oxidation rate and transport pathways. Methane is oxidized by methanotrophic bacteria, with
59 greatest rates usually measured just above the mean water table position, where there is a source
60 of both CH₄ and oxygen (Andersen et al. 2013; Clymo & Bryant 2008; Sundh et al. 1995). Since

61 the majority of CH₄ is oxidized in the unsaturated zone, water table is often a good predictor of
62 flux (Couwenberg & Fritz 2012). Plants with aerenchymatous tissue, which transports oxygen to
63 roots growing in saturated soils, can contribute to CH₄ oxidation below the water table, as radial
64 oxygen loss from roots supports methanotrophic activity (Popp et al. 2000). Anaerobic oxidation
65 has also been reported from peatlands (Gupta et al. 2013) and is likely linked to reduction of TEAs
66 including NO₃⁻, SO₄²⁻; however, its role in reducing CH₄ flux *in situ* remains unclear.

67 Once produced, CH₄ can be transported to the atmosphere via diffusion through the peat matrix,
68 plant-mediated transport, and ebullition. Diffusion through peat is slow due to its high water
69 content, and has the potential to result in high rates of CH₄ oxidation, particularly if the water table
70 is deep (Lai 2009). Plant-mediated transport is the movement of CH₄ through plant tissue as
71 diffusion through aerenchyma, pressure-driven flow, or dissolved in water lost through
72 transpiration (Lai 2009). Emission by plants can account for the majority of CH₄ flux from
73 peatlands, particularly when plants with aerenchyma are present (Couwenberg & Fritz 2012).
74 Ebullition may also account for a large proportion of CH₄ emission; Glaser et al. (2004) estimated
75 that several large, episodic ebullition events accounted for over 50% of annual emissions from a
76 bog. The importance of ebullition in fens is less clear, but losses of CH₄ through plant-mediated
77 transport may reduce subsurface CH₄ pools (Strack et al. 2017), potentially limiting bubble
78 accumulation and thus ebullition.

79 Following oil sands extraction in Alberta, government regulations require the return of the
80 landscape to equivalent land capability. While this does not necessarily require the return to
81 conditions identical to those present pre-disturbance, focus on returning peatland ecosystems to
82 the post-mining landscape has increased in recent years (Daly et al. 2012; Environment and Parks
83 2017). Reclamation criteria for particular disturbances related to oil sands extraction, such as well-
84 pads and associated roads, focus largely on returning appropriate vegetation communities
85 (Environment and Parks 2017); however, Nwaishi et al. (2015) argue that outcomes should be
86 evaluated using functional indicators. Methane production and emission indicates decomposition
87 of organic matter under high-reduced, anoxic conditions, those characteristics of peat-forming
88 conditions, indicating that CH₄ accumulation in pore water and atmospheric flux are useful
89 indicators of peatland function. However, to be used as a functional indicator, data from
90 representative reference ecosystems, in this case fens in boreal western Canada, is required. A
91 review of previous measurements of CH₄ flux in northern fens reports mean annual emissions of

92 15.4 g CH₄-C m⁻² yr⁻¹, with large variation between sites being significantly related to water table
93 position (Abdalla et al. 2016). According to Abdalla et al.'s (2016) compiled data set, Canadian
94 fens emitted 0 to 154 g CH₄-C m⁻² yr⁻¹; however, this only incorporated one reported measurement
95 from Alberta, namely 2.8 g CH₄-C m⁻² yr⁻¹ from a treed moderately-rich fen (Long et al. 2010).
96 Given the limited available data on fen CH₄ flux from the AOSR and continued disturbance of
97 natural peatlands, it is critical to quantify CH₄ fluxes from a range of fen types in the region that
98 could be used as reference ecosystems and generate data for evaluation of current and future
99 reclamation projects. Therefore, the objectives of this study were to: 1) quantify CH₄ flux, pore
100 water concentration and potential production rates from representative fens in the AOSR, 2)
101 evaluate spatial and temporal variation in CH₄ dynamics at each fen type, and 3) investigate the
102 relationship of CH₄ dynamics to ecohydrological conditions.

103 **Methods**

104 *Study sites*

105 In May 2011, three main study sites (poor fen, moderate-rich fen and saline fen) were chosen in
106 the AOSR that represented a range of fen types in the region, had large sections where hydrology
107 was unaffected by human disturbance, and were sufficiently accessible to allow for frequent
108 measurements. The poor fen sites included distinct open (OPF) and treed (TPF) areas, whereas the
109 moderately-rich fen was treed (TRF), and the saline fen was open (SF). We acknowledge that
110 resource exploration and extraction is widespread in the region, therefore it is virtually impossible
111 to find truly undisturbed sites; however, in all cases, sampling plots were located more than 50 m
112 from any disturbance (e.g. road, cutlines).

113 The poor fen (Pauciflora fen, see also Wells et al. 2017) is located ~40 km south of Fort McMurray
114 (56° 22.610 N, 111° 14.164 W). This fen receives discharge from peatlands upstream and a forested
115 upland surrounding the fen. The peat is 4 m deep on average; however, thickness varies widely
116 ranging from <1 m to >10 m. The site is situated close to a road at its north end, leading to wetter
117 conditions in this portion of the site (Bocking et al. 2017). Plant species include *Sphagnum* spp.,
118 *Chamaedaphne calyculata*, *Carex* spp., *Picea mariana* and *Betula pumila*. The poor fen basin is
119 dominated by *Sphagnum* moss species (*Sphagnum fuscum* and *Sphagnum angustifolium*);
120 however, distinct plant communities are observed in the north and south of the basin. TPF occupies

121 the central part of the fen, where water table (WT) is deeper than in the northern and southern
122 parts, which are wetter and dominated by sedges (OPF). Mean pH and corrected electrical
123 conductivity (corrEC; Sjors 1950) were 5.6 and 45 $\mu\text{S cm}^{-1}$ at OPF and 4.9 and 25 $\mu\text{S cm}^{-1}$ at TPF.

124 The rich fen (TRF; Poplar fen, Elmes et al. 2018) is located ~20 km north of Fort McMurray
125 ($56^{\circ} 56.330 \text{ N}$, $111^{\circ} 32.934 \text{ W}$). This site was disturbed by cutlines and a pipeline and dirt roads
126 passing through the broader fen boundaries, although the actual study area has not been directly
127 impacted. The site's vegetation is dominated by *Larix laricina*, *Betula pumila*, *Equisetum*
128 *fluviatile*, *Smilacina trifolia*, *Carex* spp. and brown mosses, dominated by *Tomenthypnum nitens*.
129 The peat was about 1 to 1.5 m thick, mean pH was 7.0 and corrEC was 330 $\mu\text{S cm}^{-1}$.

130 The saline fen (SF) is located 10 km south of Fort McMurray ($56^{\circ} 34.398 \text{ N}$, $111^{\circ} 16.518 \text{ W}$) and
131 is dominated by *Juncus balticus*, *Calamagrostis stricta* and *Triglochin maritima*. It is an extremely
132 saline site due to its geological setting that causes discharge of saline groundwater (Wells & Price
133 2015). The site is surrounded by forested peatland but there are no trees in the study area. Peat
134 depth was 0.75 to 1.5 m, pH was 6.1 and corrEC was 12,000 $\mu\text{S cm}^{-1}$. Although saline fens are not
135 widespread in the region, they represent a potentially important reference system for peatland
136 reclamation, as construction materials in the post-mining AOSR landscape will include tailings
137 sand, which represents a source of salinity (Simhayov et al. 2018), and is likely to result in saline
138 wetlands (Trites & Bayley 2009).

139 At each of the four sites, three replicate pairs of sampling plots were established, each pair
140 encompassing a hummock/ridge and a hollow/depression. The plots included collars for
141 greenhouse gas measurement (GHG) where vegetation surveys were also undertaken in 2011 and
142 2014, pore-water samplers, temperature measurements, dipwells, and were also used for peat
143 sampling in 2014.

144 ***Methane flux***

145 All study sites were monitored for CH_4 flux between June 2011 and August 2014. Gas samples
146 were collected weekly to biweekly in 2011 from June 23rd to August 11th, resulting in 6 – 7
147 measurements at each plot. In 2012 samples were also collected weekly to biweekly between May
148 9th and August 25th, with an additional measurement in mid-October, resulting in 9 – 15
149 measurements at each plot. In 2013 and 2014, sampling frequency declined to once every three
150 weeks between May 19th and August 22nd, with 5 – 6 measurements made at each plot.

151 Methane flux was measured using static closed chambers. At each sampling location, stainless
152 steel collars (60 cm x 60 cm) were installed 10 – 15 cm deep in the peat in early June 2011 and
153 left in place for the remainder of the study. During a measurement, an opaque acrylic chamber (60
154 cm x 60 cm x 30 high) was placed on the collar and the collar was filled with water to prevent air
155 leakage. A hole in the top of the chamber prevented over-pressurization during chamber
156 placement. Once the chamber was in place, the hole was blocked with a stopper equipped with
157 tubing sealed with a three-way valve. The headspace was mixed with a battery-operated fan and
158 samples were collected using a syringe at 7, 15, 25, and 35 minutes post-chamber closure, and
159 immediately injected into pre-evacuated Exetainers (Labco Ltd. UK). Samples were analyzed for
160 CH₄ content on a Varian 3800 gas chromatograph (GC, Varian 3800) equipped with a flame
161 ionization detector. The GC was calibrated every eight samples and standards were within +/- 10%
162 of known concentrations. Methane flux was estimated from the linear change in CH₄ concentration
163 in the headspace over time after correcting for actual headspace volume and temperature, except
164 when concentration change was less than the precision of the GC, in which case flux was assigned
165 a value of 0. Patterns of concentration change suggesting disturbance during the measurement
166 period (e.g., 7-minute concentration > 5 ppm with concentration falling over the rest of the closure
167 period or rapid increase in the middle of the closure period followed by decline) were removed
168 from the data set. This resulted in a loss of 20% of the data over the entire study period. Therefore,
169 results presented here largely represent diffusive and plant-mediated fluxes as sporadic ebullition
170 events would have been removed from the data set based upon our quality control criteria; steady
171 ebullition, if it occurred, would be included as it would result in a linear increase in concentration
172 change over the chamber closure period.

173 *Pore water CH₄ concentration*

174 Pore water CH₄ concentration was determined from water samples (Strack et al. 2004) collected
175 from samplers installed ~10 cm deeper than the WT position in early June 2011. As WT fluctuated
176 over the study period, some samplers were occasionally above the WT, and could not be sampled
177 at these times. Samplers consisted of 20 cm long segments of 2.5 cm diameter plastic pipe with
178 holes drilled in the middle 10 cm. Samplers were sealed at both ends and covered in synthetic
179 nylon screening (250 μm mesh size) to prevent clogging. Tubing extended from the bottom end of
180 the sampler to the soil surface where it was sealed with a three-way valve. The entire sampler was

181 filled with water and the valve was closed to prevent oxygen leakage to the sampler. To collect a
182 sample, 40 – 60 ml of water was removed from the sampler using a syringe and discarded. Then,
183 20 ml was collected and equilibrated with 20 ml of ambient air in the syringe by shaking for 5
184 minutes. The headspace in the syringe was then transferred to a pre-evacuated Exetainer, and CH₄
185 concentration determined on the GC. Pore water concentration was calculated according to
186 Kampbell and Vandergrift (1998).

187 ***Methane production potential***

188 Five composite replicate peat samples per site were collected at OPF, TRF, and SF in July 2014.
189 Each sample consisted of five homogenized cores 15 cm long and 10 cm in diameter, three of them
190 being collected adjacent to collars. Fresh peat was stored in plastic zipper bags at 4°C in the dark
191 until analysed. From each sample, triplicate sub-samples were prepared for incubation by placing
192 10 g of peat in Erlenmeyer flasks and adding sufficient Milli-Q water to reach saturation. The
193 flasks were prepared in a glovebox flushed with N₂ to ensure anoxic conditions. Flasks were closed
194 with butyl rubber stoppers in the glovebox prior to incubation in the dark at 24.5°C for 6 weeks.
195 Gas was sampled at 0, 1, 2, 5, 7, 14, 21, 28, 38, and 42 days of incubation using a syringe flushed
196 three times with N₂. During sampling, 20 mL of gas from the headspace was transferred into a pre-
197 evacuated Exetainer and replaced with 20 mL of 99% N₂. Methane concentration was measured
198 using a Shimadzu GC (GC-2014) with flame ionization detector.

199 ***Gross Ecosystem Productivity (GEP)***

200 Productivity of the vegetation within the collars was estimated under full light conditions using
201 dynamic transparent closed chambers (60 cm x 60 cm x 30 cm) to determine net ecosystem
202 exchange of carbon dioxide. Measurements were carried out over 2 minute closure periods with
203 CO₂ concentration measured every 15 seconds using a portable infrared gas analyzer (EGM-4,
204 PPSystems). Ecosystem respiration was determined by placing an opaque shroud over the chamber
205 and GEP calculated as the difference between net exchange and respiration (see Munir et al. 2015
206 for further details). As these measurements included only the vegetation within the collar, they
207 represent GEP of the understory vegetation only and do not include trees present at TPF and TRF
208 and therefore underestimate total ecosystem GEP.

209 ***Environmental conditions***

210 Water table position was measured in wells installed adjacent to each collar during each flux
211 measurement. Water table was also monitored hourly in one additional well at all representative
212 fen types using a pressure transducer (Solinst levellogger), corrected for barometric pressure
213 (Solinst barologger). This continuous water table record was regressed with manual measurements
214 at each plot and was expressed relative to the soil surface of hummocks at each study site. Soil
215 temperature was measured adjacent to each collar during flux measurements using an Omega
216 HH200A temperature probe at depths from -5 cm to -30 cm with 5 cm depth increments.
217 Meteorological data for the region was compiled from Environment and Climate Change Canada's
218 climate data, using the Fort McMurray AWOS station for 2011 – 2012 and its replacement station
219 Fort McMurray A for 2013 – 2014 (Environment and Climate Change Canada 2018)

220 ***Growing season CH₄ emissions***

221 The growing season length was determined according to Robeson (2002) as the number of days
222 between the last freeze in spring and first freeze in autumn. A freeze was defined as a threshold of
223 daily minimum temperatures (Linderholm 2006); in this study, a threshold of 0 °C was used. The
224 average length of the growing season in 2011 – 2014 was calculated to be 113 days, using data
225 recorded at the Fort McMurray CS meteorological station (56°39'04''N, 111°12'48''W, elevation
226 368.80 m, Environment and Climate Change Canada 2018). CH₄ emission per growing season was
227 estimated by multiplying the mean flux for each site by 113 days.

228 ***Data analysis***

229 Statistical analysis, graphs and tables were prepared using R (R Core Team 2017). Temporal and
230 spatial variability in CH₄ flux and dissolved CH₄ concentration in pore water (PW[CH₄]) was
231 analyzed using linear mixed effect (lme) models built with the 'nlme' package (Pinhero et al.
232 2017). In all cases, the mean for each plot for each study season was used. We chose to use seasonal
233 means as several studies have observed stronger correlations between peatland CH₄ flux and
234 environmental conditions on monthly to seasonal time scales compared to shorter time periods
235 (Treat et al. 2007; Turetsky et al. 2015). Moreover, the aim was to develop understanding of fen
236 CH₄ flux as a functional indicator for restoration/reclamation outcomes. In this case it is the overall
237 CH₄ emission expected under a given set of environmental conditions established in the reclaimed

238 site that is of interest, as opposed to the daily variation in that flux. The model included site, year,
239 microform, and paired interactions between them. Separate lme models were created to investigate
240 the significance of environmental factors on CH₄ flux and PW[CH₄]. These models contained fixed
241 effects of GEP, WT, T5, T20, and categorical (site, year) variables and paired interactions between
242 them. All lme models in the study included plot as a random factor to account for multiple sampling
243 at the same locations. F-values for models and type III (marginal) errors were generated using the
244 ‘anova’ function. To determine the variables explaining important amounts of variation in CH₄
245 flux between collars, non-significant controls and their interactions were removed from the model
246 one at a time, in order of the highest p-value (Zuur et al. 2009). The R² of each model was
247 calculated using the ‘r.squaredGLMM’ function in the ‘MuMIn’ package (Barton 2016). All
248 models were built for measured CH₄ flux and PW[CH₄] and for log transformed data (log CH₄ and
249 log PW[CH₄]). Models were validated for normality and distribution of residuals; only those
250 incorporating log transformed values were used for further spatial-temporal and environmental
251 analysis, and only the results from the models using log transformed data are reported here.

252 To evaluate which sites and years within sites showed similar CH₄ and PW[CH₄] patterns
253 and which ones varied significantly from each other, Tukey pairwise comparison was conducted
254 using the ‘lsmeans’ function in the ‘lsmeans’ package (Lenth 2016).

255 The significance of single environmental factors in CH₄ flux and PW[CH₄] was likely
256 obfuscated by the presence of significant interactions between model components. When such
257 interaction occurred between a categorical (site or year) and a continuous variable, regression was
258 calculated for each category using the ‘lstrends’ function in the ‘lsmeans’ package (Lenth, 2016).
259 The significance of regression was assessed based on the 0.95 confidence interval. Further
260 comparison (e.g., how the relationship between CH₄ flux and water table differs between sites
261 given the significance of water table x site interaction) was conducted using Tukey pairwise
262 comparisons.

263 Graphs were prepared using the ‘ggplot2’ (Wickham 2009), ‘cowplot’ (Wilke 2016) and
264 ‘gridExtra’ (Auguie 2016) packages. The ‘tables’ package (Murdoch 2017) was used to calculate
265 descriptive statistics for parameters reported in Table 1. Although the data are not normally
266 distributed, which is common for greenhouse gas fluxes, we have included mean and standard
267 deviation values to enable comparison with other published data on CH₄ pore water concentration
268 and fluxes, which are frequently reported as mean values.

269 Potential rate of CH₄ production was calculated as a slope of CH₄ concentration increase
270 over time. For both sets of data, linear models were built using the ‘lm’ function in the ‘nlme’
271 package (Pinheiro et al. 2017), followed by analysis of variance (‘anova’ function) to investigate
272 if the rates of CH₄ production varied significantly between sites. If the relationship was found to
273 be significant ($P < 0.05$), Tukey pairwise comparison was conducted using the ‘lsmeans’ function
274 in the ‘lsmeans’ package (Lenth 2016) to identify significant differences between sites.

275 **Results**

276 *Environmental conditions*

277 Over the 2011 – 2014 study period, mean monthly growing season (May – October) temperatures
278 were similar to or warmer than the 30-year average (1980-2010; Table 1). It was 1 – 2 °C warmer
279 most months in 2012 and 2014, while 2013 had a particularly warm May and June. With only
280 213.5 mm of precipitation, 2011 was relatively dry compared to the long-term average of 313.5
281 mm over the same period (Table 1). Conditions were wetter than average in 2012, receiving over
282 50 mm more than normal, 2013 and 2014 received just over the average amount of precipitation.
283 However, in both 2012 and 2013, there were several summer months with > 100 mm of
284 precipitation, which led to shallow water table position and frequent inundation at many of the
285 study sites by 2013 (Figure 1).

286 Water table position at OPF remained close to the surface throughout all study years, falling
287 slightly in August and September (Figure 1). At TPF, water table was generally 10 to 20 cm below
288 the surface, also varying comparatively little between years. Water table at TRF was much more
289 variable, declining from just below the surface in 2011 to deeper than -40 cm by September 2012,
290 followed by largely flooded conditions in 2013 and 2014. At SF, water table also dropped below
291 the surface throughout 2011 and early 2012, but rapidly rebounded with the large amount of
292 precipitation in July 2012. This led to standing water in many locations that persisted into 2013.
293 Water table at SF then gradually declined throughout 2014.

294 *Methane flux*

295 Generally, CH₄ fluxes increased from May into June and July, declining by late August (Figure
296 2). Measured fluxes in 2011 did not exceed 125 mg m⁻² d⁻¹ at sites TPF and TRF, were close to 0
297 mg m⁻² d⁻¹ at SF, and were up to ~350 mg m⁻² d⁻¹ at OPF. In 2012, CH₄ fluxes did not increase

298 until the end of June (days 158 – 168). The peak in CH₄ emission overlapped with increased
299 precipitation in July and decreased gradually towards the end of August. In 2013, increased CH₄
300 emission at both OPF and TPF overlapped with high precipitation in June and July (~170 mm and
301 ~80 mm, respectively, Table 1). Mean CH₄ fluxes in 2013 at SF gradually increased during the
302 growing season from ~25 mg m⁻² d⁻¹ to ~200 mg m⁻² d⁻¹. Mean CH₄ fluxes in 2014 were lower
303 than in 2013 at all sites except TRF.

304 While CH₄ flux from hollows was generally higher than from hummocks, the linear mixed effects
305 models built to characterize the spatio-temporal patterns in CH₄ flux and PW[CH₄] indicated that
306 microform type (i.e., hummock vs. hollow) did not explain a significant amount of variation. Thus,
307 microforms were removed from the models. There was a significant interaction between year and
308 site ($F_{9,60} = 5.2$, $p < 0.001$) and CH₄ fluxes varied significantly between sites ($F_{3,20} = 16.4$, $p <$
309 0.001), but not years ($F_{3,60} = 1.5$, $p = 0.22$). The highest fluxes were measured at OPF in 2012 with
310 mean (standard deviation) of 152.7 (446.6) mg CH₄ m⁻² d⁻¹, while other sites showed relatively
311 low fluxes that year (Table 2). The lowest flux was observed at SF in 2011, a mean flux of -0.02
312 (0.7) mg CH₄ m⁻² d⁻¹ (negative values indicate net removal of CH₄ from the atmosphere). The CH₄
313 fluxes were similar across years at OPF, but were significantly lower in 2011 than in the other
314 years at TPF and SF, and significantly higher in 2013-2014 than 2011-2012 at TRF (Figure 3A).
315 Over all study years, mean CH₄ at OPF was significantly higher than at TRF and SF (99.8, 32.8,
316 and 34.6 mg CH₄ m⁻² d⁻¹, respectively), while on average 68.3 mg CH₄ m⁻² d⁻¹ was emitted from
317 TPF, which was not significantly different from mean fluxes at OPF and TRF (Table 2, Figure
318 3A).

319 *Pore water CH₄ concentrations*

320 Concentrations of CH₄ dissolved in pore water generally followed the same pattern as CH₄ fluxes:
321 the highest mean PW[CH₄], over 5 mg CH₄ L⁻¹, were observed at OPF and TPF sites, while TRF
322 and SF had lower values (2.7 (4.0) and 0.9 (1.8) mg CH₄ L⁻¹, respectively, Table 2). Pore water
323 samples were taken less frequently than CH₄ fluxes, thus temporal trends are not as clear as for
324 CH₄ fluxes; however, mean PW[CH₄] increased over the growing season at TPF and TRF in 2011,
325 2012, and 2013 (Figure 4). A few measurements taken in 2014 at all sites showed much higher
326 values than in other years, but no consistent patterns of increase or decrease over the study period
327 in mean PW[CH₄], aside from decreasing values at the TPF.

328 In the linear mixed effects model, year ($F_{3,53} = 10.8$, $p < 0.001$), site ($F_{3,20} = 30.0$, $p < 0.001$) and
329 their interaction ($F_{9,53} = 7.4$, $p < 0.001$) explained a significant fraction of spatio-temporal
330 variability of PW[CH₄]. At OPF and TPF, PW[CH₄] was significantly higher in 2014 compared to
331 the early study years (Figure 3B). A similar pattern was observed at TRF; however, at this site
332 PW[CH₄] was also significantly lower in 2012 compared to 2011 and 2013. At SF, concentration
333 was significantly lower in 2011 than all other study years. Over the whole study period, both poor
334 fen sites had significantly higher PW[CH₄] than TRF and SF, which were also significantly
335 different from each other (Figure 3B).

336 *Potential CH₄ production*

337 The mean potential CH₄ production was 0.8 (0.5), 0.1 (0.1) and 4.6 (0.8) $\mu\text{mol CH}_4 \text{ g peat}^{-1} \text{ d}^{-1}$ at
338 OPF, SF, and TRF, respectively. Outliers (30.9 and 0.023 $\mu\text{mol CH}_4 \text{ g peat}^{-1} \text{ d}^{-1}$ from OPF and
339 TRF, respectively) were rejected before the calculation of the mean values; in both cases, the
340 removed values were two orders of magnitude differ than the other replicates from the same study
341 site. The potential rate of CH₄ production was significantly higher at TRF than OPF and SF ($p =$
342 0.0025 and $p = 0.0009$, respectively).

343

344 *Relationship of CH₄ flux and pore water concentrations to environmental conditions*

345 Generally, all parameters, except temperature at -5 and -20 cm depth, varied widely between years
346 and across sites (Table 2). Ground layer GEP was negative at all sites and in all years, meaning
347 that there was net uptake of CO₂ from the atmosphere, which is expected for natural peatlands. SF
348 had the highest CO₂ uptake (from -21.5 to -38.2 $\text{g CO}_2 \text{ m}^{-2} \text{ d}^{-1}$), and TRF the lowest (from -10.1 to
349 -17.3 $\text{g CO}_2 \text{ m}^{-2} \text{ d}^{-1}$). However, it should be noted that as the overstory production was not included
350 at TRF, total GEP would be underestimated.

351 Methane flux was significantly related to pore water CH₄ concentration and WT position
352 across all plots (Table 3, Figure 5), but there was also a significant WT-site interaction, indicating
353 that the relationship was site specific. Within each site, WT explained a significant amount of the
354 variation in CH₄ flux, but at OPF, shallower WT resulted in lower CH₄ emission, while the
355 opposite was true at all other study sites; therefore, the slope of the regression at OPF was
356 significantly different than all other sites (OPF vs. SF, $p < 0.001$, OPF vs. TPF, $p = 0.002$, OPF vs.

357 TRF, $p = 0.005$). Moreover, the slope of the WT-CH₄ flux relationship was steeper at SF, resulting
358 in a significant difference from the slope at TRF ($p = 0.001$).

359 Variation in pore water CH₄ concentration was significantly explained by WT, soil temperature at
360 20 cm (T20) and interactions between temperature at both 5 cm (T5) and T20 with site. Across the
361 whole data set, PW[CH₄] was higher when WT was shallow (Figure 5) and T20 was cooler. The
362 regression for T5 and PW[CH₄] was only significant at SF, where cooler temperatures resulted in
363 higher concentrations. Considering T20, regressions were significant for all sites except TPF. At
364 OPF and TRF, cooler temperatures resulted in higher concentration, while at SF, warmer
365 temperatures results in higher PW[CH₄].

366 *Discussion*

367 Results from the present study provide a valuable starting point for building a reference database
368 of CH₄ flux and pore water concentrations for fens in the AOSR specifically, and western Canada
369 in general. Following oil sands extraction, the function of restored and constructed peatlands can
370 be assessed in comparison to these reference ecosystems. While measurements of CH₄ flux in
371 western Canada are limited, our results agree well with previous studies in the region (Table 4).
372 OPF had similar carbon emissions to an open poor fen in the Northwest Territories (mean 99 mg
373 CH₄ m⁻² year⁻¹, Liblik et al. 1997), while TRF (mean 32.8 mg CH₄ m⁻² d⁻¹) had slightly greater
374 mean flux compared to a moderate-rich treed fen in boreal Alberta (mean 25.6 mg CH₄ m⁻² d⁻¹;
375 Long et al. 2010). Measured PW[CH₄] concentrations also were similar to literature values at sites
376 across North America (1.3 – 6.4 mg L⁻¹; Chasar et al. 2000; Murray et al. 2017a; Strack et al.
377 2004).

378 Mean flux from data compilation across boreal and temperate fens was approximately 80 mg CH₄
379 m⁻² d⁻¹ (Turetsky et al. 2015), with the mean across study years in the present study of 68.3 to 99.8
380 at poor fen sites and 32.8 mg CH₄ m⁻² d⁻¹ at the treed rich fen. We also compared our fluxes to data
381 estimated on an annual scale by estimating annual emissions of 9.9, 6.8, 3.3 and 3.4 g CH₄-C m⁻²
382 year⁻¹ for OPF, TPF, TRF, and SF, respectively, assuming that the non-growing season flux
383 contributed 15% of the total yearly flux (Saarnio et al. 2007). This assumption is similar to the
384 median non-growing season contribution of 16% determined through data compilation across
385 northern wetlands (Treat et al. 2018), but may also represent an underestimate of non-growing
386 season fluxes given that some studies report up to 47% of CH₄ flux can occur outside the growing

387 season (Treat et al. 2018). When compared to mean fluxes for northern peatlands (7.6 – 15.7 g
388 CH₄-C m⁻² year⁻¹, where lower values represented bogs and higher fens; Abdalla et al. 2016), our
389 values were closer to natural northern bogs than to fens. In the context of peatland reclamation,
390 comparing values with averages from northern fens dispersed at latitudes above 45°N may
391 therefore not be relevant, and highlights the importance of compiling data from local reference
392 ecosystems to assess ecosystem progress following intervention (e.g., Gorham & Rochefort 2003).
393 Furthermore, a wide range of mean annual CH₄ emission has been reported from northern poor
394 fens (e.g., 1.5 g CH₄-C m⁻² year⁻¹ (Godin et al. 2012) and 31 g CH₄-C m⁻² year⁻¹ (Treat et al. 2007))
395 and rich fens (e.g., 4.1 g CH₄-C m⁻² year⁻¹ (Pelletier et al. 2007) and 154 g CH₄-C m⁻² year⁻¹ (Godin
396 et al. 2012)) suggesting that the type of fen alone cannot be a proxy for the level of emitted CH₄.
397 Instead, if CH₄ flux is used as a functional indicator for restoration or reclamation evaluation,
398 specific environmental conditions prevailing at each site should be considered. Indeed, the factors
399 that significantly controlled CH₄ flux in our research, e.g., WT, and soil temperature, have been
400 previously recognized in other peatland studies (Pelletier et al. 2007; Strack et al. 2004; Treat et
401 al. 2007; Whalen 2005; Rinne et al. 2018).

402 Water table is a widely reported control on peatland CH₄ flux, and our data further supports this
403 observation (Table 3; Figure 4). Interestingly, WT affected CH₄ fluxes differently at OPF than the
404 other three sites (Table 4). Generally, CH₄ emission increases at sites with shallower WT (Abdalla
405 et al. 2016), as deep WT position increases the size of the oxic zone and the likelihood that CH₄
406 will be oxidized to CO₂ before it reaches the peat surface (Whalen 2005). This pattern was
407 observed at TPF, TRF and SF. At OPF, CH₄ emission declined with increasing WT. While this
408 site was characterized by more stable (Wells et al. 2017) and shallower WT than the other sites, it
409 also had a complete cover of *Sphagnum*, which are known to host symbiotic CH₄ oxidizing bacteria
410 (Kip et al. 2010). These communities can support high rates of CH₄ oxidation even in submerged
411 conditions (Parmentier et al. 2011) when CH₄-rich pore water comes in contact with the living part
412 of *Sphagnum*. This CH₄ emission pattern could also be caused by reduced vegetation productivity
413 under inundated conditions (Strack et al. 2004) that would reduce substrate availability for
414 methanogens and thus limit CH₄ production. These wet conditions at OPF, and other sites in some
415 study years, may have also contributed to the lack of a statistically significant effect of microform
416 type in spatial variability of CH₄ fluxes. Although fluxes > 50 mg CH₄ m⁻² d⁻¹ were observed in
417 hollows during the dry growing season of 2011, large fluxes were measured at both hummocks

418 and hollows in wetter years, with hummocks emitting large amounts of CH₄ at the OPF. At this
419 site, hummocks were often only a few cm higher than hollows, resulting in limited WT differences
420 between microforms.

421 Differences in the slope of the CH₄ flux-WT and PW[CH₄]-T20 relationship between sites may
422 also be driven by chemical differences between the study fens. The poor fen sites were more acidic
423 than TRF and SF, where pH of 6 – 7 is optimal for methanogens (Blodau 2002). This suggests that
424 under saturated conditions, CH₄ emissions should be higher from TRF and SF. Although fluxes at
425 all sites were similar when WT was near the surface (Figure 4), high CH₄ production potential at
426 TRF does suggest more favourable substrate availability and/or chemical conditions than the other
427 sites. CH₄ production and flux can also be limited by the availability of terminal electron acceptors
428 (TEAs), such as nitrate, iron and sulphate (Lai 2009), that are more likely to be present in higher
429 concentrations at TRF and SF than the poor fen sites. In fact, Murray et al. (2017a) found a strong
430 negative relationship between sulphur availability and CH₄ flux and PW[CH₄] across OPF and SF.
431 The relatively steep slope of the CH₄ flux-WT relationship at SF may reflect the importance of
432 sulphate as a control on fluxes at this site; once WT drops below the surface, reduced sulphur could
433 be oxidized to sulphate, quickly limiting CH₄ production. The presence of sulphate could also
434 account for the low CH₄ production potential measured at SF. Given that studies of fen reclamation
435 projects in the AOSR report high sulphate concentrations (Nwaishi et al. 2015; Murray et al.
436 2017a; Clark et al. 2019), measuring pore water chemistry, specifically for TEAs will be important
437 when using CH₄ as a functional indicator of reclamation outcomes.

438 Substrate availability for methanogens is also a control on CH₄ production and hence PW[CH₄]
439 and CH₄ flux (Lai 2009). Some previous studies have found a significant positive correlation
440 between plant productivity and CH₄ flux (e.g., Whiting & Chanton 1993; Bellisario et al. 1999),
441 but GEP was not a significant predictor of variation in mean flux between measured plots in the
442 present study. This could be due to the multifaceted role that plants play in CH₄ flux, both
443 increasing production and transport (Waddington et al. 1996, Strack et al. 2017), while also
444 enhancing oxidation (Popp et al. 2000, Sutton-Grier & Megonigal 2011). For example, while
445 aerenchymatous plants have been shown to increase the transport of CH₄ from the soil to the
446 atmosphere, in some cases, CH₄ emissions can be reduced due to greater oxidation in the rooting
447 zone (Bhullar et al. 2013). When WT is favourable for CH₄ production, other environmental
448 variables such as soil temperature and GEP can drive spatial and temporal variation in CH₄ flux

449 (Strack et al. 2004). Since WT varied widely between sites and years, it may be that this acted as
450 an overriding control compared to GEP. It should be noted that measured GEP did not include the
451 overstory productivity, which is likely substantial at TPF and TRF (Murray et al. 2017b). Including
452 an estimate of total ecosystem GEP potentially would further explain differences in CH₄ flux
453 between the studied fens. Further, some studies indicate that trees can vent CH₄ in wetlands (Gauci
454 et al. 2010; Pangala et al. 2013), suggesting that the role that trees play in total CH₄ emissions in
455 western Canadian peatlands also requires further investigation.

456 One of the most striking patterns observed was the large inter-annual variability in CH₄ fluxes. In
457 the year following the data collection period presented here, Murray et al. (2017a) measured CH₄
458 emission at some of these same sites and report a flux of 23.9 mg CH₄ m⁻² d⁻¹ at OPF and 4.4 mg
459 CH₄ m⁻² d⁻¹ at SF. Both, if used to recalculate mean CH₄ flux for years 2011 – 2015, would reduce
460 the flux by ~20% at OPF and ~10% at SF. This inter-annual variability can be captured only with
461 long-term studies, illustrating their importance to the development of robust reference datasets.
462 Wells et al. (2017) also underlined the importance of long-term studies on peatland hydrology
463 (main driver of CH₄ fluxes) in relation to long-term climate variability.

464 The studied fens also varied in their CH₄ flux response to inter-annual variation in summer
465 precipitation, a pattern likely driven by local hydrogeologic setting. Hydrological studies at TRF
466 indicate that it has a longer “memory” than the other study fens, with WT position carrying over
467 from available moisture in previous years (Elmes et al. 2018). For example, while most sites
468 exhibited wetter conditions in 2012 than 2011, TRF remained dry and consequently had lower CH₄
469 emissions than the other sites in this year. Furthermore, availability of TEAs at TRF, particularly
470 after long periods of low water table may have also contributed to lower CH₄ emissions.

471 Compared to inter-annual variation in flux, PW[CH₄] was less variable; values remained relatively
472 similar over the first 3 years of study, but increased in 2014 at TPF, OPF, and TRF. This may
473 indicate that PW[CH₄] is a useful functional indicator for reclamation outcomes as it appears to be
474 more stable in response to inter-annual hydrologic variations. However, there appears to be some
475 decoupling between CH₄ emissions and PW[CH₄] (Figure 3). The continuing increase in PW[CH₄]
476 in 2014 while emissions declined compared to the previous year may reflect the fact that mean
477 residence time of CH₄ in peat can be months to years (Liblik et al. 1997; Strack and Waddington
478 2008). Therefore, the dissolved CH₄ pool in 2014 potentially reflects the favourable CH₄
479 production conditions in 2013 when all sites had shallow water tables.

480 **Conclusions**

481 We measured CH₄ production potential and CH₄ flux and pore water CH₄ concentration
482 (PW[CH₄]) over four growing seasons, across a variety of fen types in the Athabasca Oil Sands
483 Region (AOSR) in order to contribute to the development of a reference fen dataset that can be
484 used for functional evaluation of fen restoration and reclamation projects in the region. Mean flux
485 over the study period was 32.8 to 99.8 mg CH₄ m⁻² d⁻¹, while mean PW[CH₄] was 0.9 to 5.5 mg
486 CH₄ L⁻¹. While differences within and between fens and across study years were largely driven by
487 WT position, this relationship varied between sites likely linked to local plant community and
488 chemistry. The high level of variability in CH₄ flux between study years indicates the importance
489 of multi-year studies of CH₄ flux, not only for developing reference datasets, but also for
490 measuring ecosystem function in reclamation projects. In our study, PW[CH₄] was generally
491 linked to fluxes but showed less spatial and temporal variability. Thus, PW[CH₄] may be useful
492 independently as a functional indicator for reclamation. Our study generated a multi-year baseline
493 for fens in the AOSR, which could be used by the energy industry in the context of post-mining
494 management of disturbed or reclaimed fens in this region. Ideally, similar baseline CH₄ emission
495 and PW[CH₄] databases, including high frequency of measurements, would need to be constructed
496 for other regions.

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505 **Data availability**

506 All data is available from the corresponding author by request.

507 **Author contributions**

508 MS, MSM and RA conceived the field study and implemented the design. JP, VD, RA and FN
509 conceived the incubation study. VD, RA, FN, MSM and MS collected data and contributed to data
510 analysis. AB completed final statistical analysis and wrote the first draft of the paper. All authors
511 edited the final manuscript.

512 **References**

- 513 Abdalla M, Hastings A, Truu J, Espenberg M, Mander Ü, Smith P (2016) Emissions of methane
514 from northern peatlands: A review of management impacts and implications for future
515 management options. *Ecology and Evolution* 6: 7080-7102.
- 516 Andersen R, Chapman SJ, Artz R (2013) Microbial communities in natural and disturbed
517 peatlands: A review. *Soil Biology and Biochemistry* 57: 979-94.
- 518 Auguie B (2016) gridExtra: Miscellaneous Functions for "Grid" Graphics. R package version
519 2.2.1. URL: <https://CRAN.R-project.org/package=gridExtra>.
- 520 Barton K (2016) MuMIn: Multi-Model Inference. R package version 1.15.6. URL:
521 <https://CRAN.R-project.org/package=MuMIn>.
- 522 Bellisario LM, Bubier JL, Moore TR, Chanton JP (1999) Controls on CH₄ emissions from a
523 northern peatland. *Global Biogeochemical Cycles* 13: 81-91.
- 524 Bhullar GS, Edwards PJ, Venterink HO (2013) Variation in plant-mediated methane transport and
525 its importance for methane emission from intact wetland peat mesocosms. *Journal of Plant*
526 *Ecology* 6: 298-304.
- 527 Blodau C (2002) Carbon cycling in peatlands A review of processes and controls. *Environmental*
528 *Reviews* 10: 111-134.
- 529 Bocking E, Cooper DJ, Price J (2017) Using tree ring analysis to determine impacts of a road on
530 a boreal peatland. *Forest Ecology and Management* 404: 24-30.
- 531 Bridgham SD, Cadillo-Quiroz H, Keller JK, Zhuang Q (2013) Methane emissions from wetlands:
532 Biogeochemical, microbial, and modeling perspectives from local to global scales. *Global*
533 *Change Biology* 19: 1325-1346.

534 Chasar LS, Chanton JP, Glaser PH, Siegel DI (2000) Methane concentration and stable isotope
535 distribution as evidence of rhizospheric processes: Comparison of a fen and bog in the glacial
536 lake Agassiz peatland complex. *Annals of Botany* 86: 655-663.

537 Clark MG, Humphrey E.R, Carey SK (2019) Low methane emissions from a boreal wetland
538 constructed on oil sand mine tailings, *Biogeosciences Discussions*, [https://doi.org/10.5194/bg-](https://doi.org/10.5194/bg-2019-271)
539 [2019-271](https://doi.org/10.5194/bg-2019-271).

540 Clymo RS, Bryant C (2008) Diffusion and mass flow of dissolved carbon dioxide, methane, and
541 dissolved organic carbon in a 7-m deep raised peat bog. *Geochimica et Cosmochimica Acta* 72:
542 2048-2066.

543 Couwenberg J, Fritz C (2012) Towards developing IPCC methane 'emission factors' for peatlands
544 (organic soils). *Mires and Peat* 10: 03.

545 Daly C, Price J, Rezanezhad F, Pouliot R, Rochefort L, Graf MD (2012) Initiatives in oil sand
546 reclamation: considerations for building a fen peatland in post mined oil sands landscape. In:
547 Vitt DH, Bhatti J (eds) *Restoration and Reclamation of Boreal Ecosystems*, Cambridge
548 University Press, New York.

549 Elmes MC, Thompson DK, Sherwood JH, Price JS (2018) Hydrometeorological conditions
550 preceding wildfire, and the subsequent burning of a fen watershed in Fort McMurray, Alberta,
551 Canada. *Natural Hazards and Earth System Sciences* 18: 157-170.

552 Environment and Climate Change Canada (2018) Historical Climate Data, [cited 2018 Jul 19].
553 Available from: <http://climate.weather.gc.ca/>.

554 Environment and Parks (2017) Reclamation Criteria for Wellsites and Associated Facilities for
555 Peatlands, March, 2017, Edmonton, Alberta, pp 142.

556 Estop-Aragónés C, Knorr K-H, Blodau C (2013) Belowground in situ redox dynamics and
557 methanogenesis recovery in a degraded fen during dry-wet cycles and flooding, *Biogeosciences*
558 10: 421-436.

559 Gauci V, Gowing DJG, Hornibrook ERC, Davis JM, Dise NB (2010) Woody stem methane
560 emission in mature wetland alder trees. *Atmospheric Environment* 44: 2157-2160.

561 Glaser PH, Chanton JP, Morin P, Rosenberry DO, Siegel DI, Ruud O, Chasar LI, Reeve AS (2004)
562 Surface deformations as indicators of deep ebullition fluxes in a large northern peatland, *Global*
563 *Biogeochemical Cycles* 18: GB1003.

564 Godin A, McLaughlin JW, Webster KL, Packalen M, Basiliko N (2012) Methane and methanogen
565 community dynamics across a boreal peatland nutrient gradient. *Soil Biology and Biochemistry*
566 48: 96-105.

567 Gorham E, Rochefort L (2003) Peatland restoration: A brief assessment with special reference to
568 sphagnum bogs. *Wetlands Ecology and Management* 11: 109-119.

569 Government of Alberta (2018) [cited 2018 Jul 19]. Available from:
570 <https://www.energy.alberta.ca/OS/AOS/Pages/FAS.aspx#Environment>.

571 Gupta V, Smemo KA, Yavitt JB, Fowle D, Branfireun B, Basiliko N (2013) Stable isotopes reveal
572 widespread anaerobic methane oxidation across latitude and peatland type. *Environmental*
573 *Science and Technology* 47: 8273-8279.

574 Kampbell DH, Vandegrift SA (1998) Analysis of dissolved methane, ethane, and ethylene in
575 ground water by a standard gas chromatographic technique. *Journal of Chromatographic*
576 *Science* 36: 253-256.

577 Kip N, van Winden JF, Pan Y, Bodrossy L, Reichart G, Smolders AJP, Jetten MSM, Damste JSS,
578 Op den Camp HJM (2010) Global prevalence of methane oxidation by symbiotic bacteria in
579 peat-moss ecosystems. *Nature Geoscience* 3: 617-621.

580 Lai D (2009) Methane dynamics in northern peatlands: A review. *Pedosphere* 19: 409-421.

581 Lenth RV (2016) Least-Squares Means: The R Package lsmeans. *Journal of Statistical Software*
582 69: 1-33.

583 Liblik LK, Moore TR, Bubier JL, Robinson SD (1997) Methane emissions from wetlands in the
584 zone of discontinuous permafrost: Fort Simpson, Northwest Territories, Canada. *Global*
585 *Biogeochemical Cycles* 11: 485-494.

586 Linderholm HW (2006) Growing season changes in the last century. *Agricultural and Forestry*
587 *Meteorology* 137: 1-14.

588 Long KD, Flanagan LB, Tiebo C (2010) Diurnal and seasonal variation in methane emissions in a
589 northern Canadian peatland measured by eddy covariance. *Global Change Biology* 16: 2420-
590 2435.

591 Madigan MT (2009) *Brock biology of microorganisms*. 12th ed. San Francisco (CA),
592 Pearson/Benjamin Cummings, 1061 pp.

593 Minderlein S, Blodau C (2010) Humic-rich peat extracts inhibit sulfate reduction, methanogenesis,
594 and anaerobic respiration but not acetogenesis in peat soils of a temperate bog. *Soil Biology*
595 and *Biochemistry* 42: 2078-2086.

596 Munir TM, Perkins M, Kaing E, Strack M (2015) Carbon dioxide flux and net primary production
597 of a boreal treed bog: Responses to warming and water-table-lowering simulations of climate
598 change. *Biogeosciences* 12, 1091-1111.

599 Murdoch D (2017) tables: Formula-Driven Table Generation. R package version 0.8.3. URL:
600 <https://CRAN.R-project.org/package=tables>.

601 Murray KR, Barlow N, Strack M (2017a) Methane emissions dynamics from a constructed fen
602 and reference sites in the Athabasca oil sands region, Alberta. *Science of the Total Environment*
603 583: 369-381.

604 Murray KR, Borkenhagen AK, Cooper DJ, Strack M (2017b) Growing season carbon gas
605 exchange from peatlands used as a source of vegetation donor material for restoration. *Wetlands*
606 *Ecology and Management* 25: 501-515.

607 Nwaishi F, Petrone R, Price J, Andersen R (2015) Towards developing a functional-based approach
608 for constructed peatlands evaluation in the Alberta oil sands region, Canada. *Wetlands* 35: 211-
609 225.

610 Pangala SR, Moore S, Hornibrook ERC, Gauci V (2013) Trees are major conduits for methane
611 egress from tropical forested wetlands. *New Phytologist* 197: 524-531.

612 Parmentier FJW, van Huissteden J, Kip N, Op den Camp HJM, Jetten MSM, Maximov TC,
613 Dolman AJ (2011) The role of endophytic methane-oxidizing bacteria in submerged *Sphagnum*
614 in determining methane emissions of Northeastern Siberian tundra. *Biogeosciences* 8: 1267-
615 1278.

616 Pelletier L, Moore TR, Roulet NT, Garneau M, Beaulieu-Audy V (2007) Methane fluxes from
617 three peatlands in the la Grande Rivière watershed, James Bay lowland, Canada. *Journal of*
618 *Geophysical Research* 112: 1-12.

619 Pinheiro J, Bates D, DebRoy S, Sarkar D, R Core Team (2017) nlme: Linear and Nonlinear Mixed
620 Effects Models. R package version 3.1-131, URL: <https://CRAN.R-project.org/package=nlme>.

621 Popp TJ, Chanton JP, Whiting GJ, Grant N (2000) Evaluation of methane oxidation in the
622 rhizosphere of a *Carex* dominated fen in north central Alberta, Canada. *Biogeochemistry* 51:
623 259-281.

624 Province of Alberta (2018) Environmental Protection and Enhancement Act, Conservation and
625 Reclamation Regulation. Alberta Queen's Printer, Edmonton, AB.

626 R Core Team (2017) R: A language and environment for statistical computing. R Foundation for
627 Statistical Computing, Vienna, Austria. URL: <https://www.R-project.org/>.

628 Rinne J, Tuittila E-S, Peltola O, Li X, Raivonen M, Alekseychik P, Haapanala S, Pihlatie M,
629 Aurela M, Mammarella I, Vesala T (2018) Temporal variation of ecosystem scale methane
630 emission from a boreal fen in relation to temperature, water table position, and carbon dioxide
631 fluxes. *Global Biogeochemical Cycles* 32: 1087–1106.

632 Robeson SM (2002) Increasing growing-season length in Illinois during the 20th century. *Climatic*
633 *Change* 52: 219-238.

634 Rosenberry DO, Glaser PH, Siegel DI (2006) The hydrology of northern peatlands as affected by
635 biogenic gas: Current developments and research needs. *Hydrological Processes* 20: 3601-
636 3610.

637 Saarnio S, Morero M, Shurpali NJ, Tuittila , Mäkilä M, Alm J (2007) Annual CO₂ and CH₄ fluxes
638 of pristine boreal mires as a background for the lifecycle analyses of peat energy. *Boreal*
639 *Environment Research* 12: 101-113.

640 Simhayov RB, Weber TKD, Price JS (2018) Saturated and unsaturated chemical non-equilibrium
641 salt transport in peat from a constructed fen. *Soil* 4: 63-81.

642 Sjörs H (1950) On the relation between vegetation and electrolytes in north Swedish mire waters.
643 *Oikos* 2: 241-258.

644 Strack M, Mwakanyamale K, Hassanpour Fard G, Bird M, Bérubé V, Rochefort L (2017) Effect
645 of plant functional type on methane dynamics in a restored minerotrophic peatland. *Plant and*
646 *Soil* 410: 1-16.

647 Strack M, Waddington JM (2008) Spatiotemporal variability in peatland subsurface methane
648 dynamics. *Journal of Geophysical Research* 113: G02010, doi: 10.1029/2007JG000472.

649 Strack M, Waddington JM, Tuittila E-S (2004) Effect of water table drawdown on northern
650 peatland methane dynamics: Implications for climate change. *Global Biogeochemical Cycles*
651 18: GB4003, doi: 10.1029/2003GB002209.

652 Sundh I, Mikkilä C, Nilsson M, Svensson BH (1995) Potential aerobic methane oxidation in a
653 sphagnum-dominated peatland--controlling factors and relation to methane emission. *Soil*
654 *Biology and Biochemistry* 27: 829-837.

655 Sutton-Grier AE, Megonigal JP (2011) Plant species traits regulate methane production in
656 freshwater wetland soils. *Soil Biology & Biochemistry* 43: 413-420.

657 Treat CC, Bubier JL, Varner RK, Crill PM (2007) Timescale dependence of environmental and
658 plant-mediated controls on CH₄ flux in a temperate fen. *Journal of Geophysical Research* 112:
659 G01014, doi: 10.1029/2006JG000210.

660 Treat CC, Bloom AA, Marushchak ME (2018) Nongrowing season methane emissions – a
661 significant component of annual emissions across northern ecosystems. *Global Change Biology*
662 24: 3331-3343.

663 Trites M, Bayley SE (2009) Vegetation communities in continental boreal wetlands along a
664 salinity gradient: Implications for oil sands mining reclamation, *Aquatic Botany* 91: 27-39.

665 Tuittila E, Komulainen V, Vasander H, Nykanen H, Martikainen P, Laine J (2000) Methane
666 dynamics of a restored cut-away peatland. *Global Change Biology* 6: 569-581.

667 Vitt DH, Halsey LA, Bauer IE, Campbell C (2000) Spatial and temporal trends in carbon storage
668 of peatlands of continental western Canada through the Holocene. *Canadian Journal of Earth*
669 *Sciences* 37: 683-693.

670 Vitt D, Halsey L, Thormann M, Martin T (1996) Peatland inventory of Alberta. Phase 1: Overview
671 of peatland resources in the natural regions and subregions of the province, University of
672 Alberta, Edmonton, AB, Canada.

673 Waddington JM, Roulet NT, Swanson RV (1996) Water table control of CH₄ emission
674 enhancement by vascular plants in boreal peatlands. *Journal of Geophysical Research* 101:
675 22775-22785.

676 Wells C, Ketcheson S, Price J (2017) Hydrology of a wetland-dominated headwater basin in the
677 boreal plain, Alberta, Canada. *Journal of Hydrology* 547: 168-183.

678 Wells CM, Price JS (2015) A hydrologic assessment of a saline-spring fen in the Athabasca oil
679 sands region, Alberta, Canada – a potential analogue for oil sands reclamation. *Hydrological*
680 *Processes* 29: 4533-4548.

681 Whalen SC (2005) Biochemistry of methane exchange between natural wetlands and the
682 atmosphere. *Environmental Engineering Science* 22: 73-94.

683 Whiting GJ, Chanton JP (1993) Primary production control of methane emission for wetlands.
684 *Nature* 364: 794-795.

685 Wickham H (2009) *ggplot2: Elegant Graphics for Data Analysis*. Springer-Verlag New York.

686 Wilke CO (2016) cowplot: Streamlined Plot Theme and Plot Annotations for 'ggplot2'. R package
687 version 0.7.0. URL: <https://CRAN.R-project.org/package=cowplot>.
688 Zuur AF, Ieno EN, Walker NJ, Saveliev AA, Smith GM (2009) Mixed Effects Models and
689 Extensions with R. Springer, New York.

690 **Tables**

691 Table 1. Mean monthly precipitation (P, mm) and temperature (T, °C)

Month	Year									
	2011		2012		2013		2014		Normal ^b	
	P	T	P	T	P	T	P	T	P	T
May	23.5	11.0	19.6	10.6	4.7	12.7	82.9	7.9	36.5	9.9
June	51.0	14.6	38.2	15.3	165.9	16.1	69.7	15.3	73.3	14.6
July	53.0	17.2	130.8	19.2	87.1	16.9	55.7	19.0	80.7	17.1
August	61.5	16.0	16.4	16.8	3.9	17.6	36.3	17.0	57.1	15.4
September	21.0	12.5	116.9	12.1	35.0	13.4	62.8	9.4	39.7	9.5
October	3.5	6.2	42.8	-0.2	21.1	3.6	30.3	5.2	26.2	2.3
Growing season ^a	213.5	12.9	364.7	12.3	317.7	13.4	337.7	12.3	313.5	11.5

692 a. total P and average T for May to October

693 b. normal average for years 1981 – 2010

Table 2. Descriptive statistics for measured variables

Site	Year	WT (cm)			T5 ^a (°C)			T20 ^b (°C)			GEP ^c (g CO ₂ m ⁻² d ⁻¹)			CH ₄ flux (mg CH ₄ m ⁻² d ⁻¹)			PW[CH ₄] ^d (mg L ⁻¹)		
		n	mean	sd	n	mean	sd	n	mean	sd	n	mean	sd	n	mean	sd	n	mean	sd
OPF	2011	36	-2.3	6.7	38	17.2	3.9	38	15.2	2.8	24	-23.0	10.6	31	50.0	75.6	46	4.5	1.7
	2012	45	-7.5	4.7	38	16.1	3.5	38	12.7	5.1	48	-18.2	16.1	42	152.7	446.6	32	4.5	2.0
	2013	42	-3.9	7.9	42	16.2	3.9	42	13.8	3.5	40	-16.0	10.7	29	104.1	103.7	26	4.4	2.0
	2014	29	-1.5	4.6	29	16.2	2.9	29	13.0	4.8	30	-19.3	17.9	24	66.3	97.9	12	12.8	6.8
TPF	2011	36	-9.9	12.0	36	16.4	3.2	36	13.8	2.1	37	-21.6	22.5	28	13.1	21.9	48	3.9	1.8
	2012	74	-11.6	8.1	52	15.3	3.7	52	12.0	6.1	54	-17.7	9.5	49	38.0	54.4	37	4.8	2.5
	2013	46	-3.0	8.7	46	16.9	3.7	46	14.3	3.1	36	-20.8	10.7	38	140.0	171.6	30	5.0	2.4
	2014	30	-3.4	7.1	30	17.4	2.0	30	13.4	4.2	30	-25.1	18.5	26	79.9	127.1	15	13.5	10.5
TRF	2011	28	-7.9	11.2	28	16.5	3.2	28	12.5	2.4	51	-17.3	20.0	32	20.2	32.8	29	1.9	1.7
	2012	87	-23.6	17.8	63	13.4	5.4	63	10.4	6.5	55	-15.5	16.1	65	7.7	12.3	31	1.0	3.6
	2013	36	2.2	10.8	36	13.6	3.3	36	11.3	4.2	35	-10.1	10.7	29	66.3	105.2	29	2.5	2.6
	2014	24	1.4	7.2	28	13.6	3.1	28	10.4	4.5	29	-13.8	12.0	26	73.8	74.0	14	9.1	4.8
SF	2011	13	-13.5	5.1	42	19.1	3.2	42	17.2	2.2	50	-38.2	16.0	37	0.0	0.7	37	0.2	0.1
	2012	66	-3.6	11.4	55	16.5	4.6	55	15.0	5.4	55	-29.5	20.6	55	11.1	19.9	24	0.2	0.3
	2013	38	1.0	8.3	42	16.7	2.4	42	16.0	2.2	30	-23.6	14.2	34	117.6	159.1	34	2.0	2.2
	2014	30	-6.3	10.1	30	16.9	4.8	30	15.7	4.4	32	-21.7	16.9	19	21.8	41.4	8	2.3	3.6
OPF	2011 - 2014	152	-4.1	6.6	147	16.4	3.6	147	13.7	4.2	142	-18.6	14.4	126	99.8	269.7	116	5.3	3.7
TPF	2011 - 2014	186	-7.8	9.7	164	16.4	3.4	164	13.3	4.4	157	-20.7	15.6	141	68.3	118.9	130	5.5	5.0
TRF	2011 - 2014	175	-12.4	18.5	155	14.1	4.3	155	11.0	5.1	170	-14.6	16.0	152	32.8	63.7	103	2.8	4.0
SF	2011 - 2014	147	-3.8	10.6	169	17.3	4.0	169	15.9	4.0	167	-29.5	18.5	145	34.6	91.3	103	0.9	1.9

a. T5 = soil temperature at 5 cm depth

b. T20 = soil temperature at 20 cm depth

c. GEP = gross ecosystem photosynthesis

d. PW[CH₄] = CH₄ concentration in pore water

Table 3. Results from the linear mixed effect models for CH₄ flux and pore water CH₄ concentration

Factor^a	F^b	p-value^b
$\log_{10}[\text{CH}_4 \text{ flux (mg m}^{-2} \text{ d}^{-1})]; R^2_{\text{GLMM}} = 0.66^c$		
Site	F _{3,20} = 0.4	0.77
WT	F _{1,60} = 7.5	0.008
$\log_{10}(\text{PW}[\text{CH}_4])$	F _{1,60} = 10.4	0.002
Site x WT	F _{3,60} = 10.6	<0.0001
Intercept	F _{1,60} = 24.8	<0.0001
$\log_{10}[\text{pore water CH}_4 \text{ concentration (mg L}^{-1})]; R^2_{\text{GLMM}} = 0.74^c$		
Site	F _{3,20} = 2.1	0.13
WT	F _{1,56} = 24.9	<0.0001
T5	F _{1,56} = 0.10	0.76
T20	F _{1,56} = 6.41	0.014
Site x T5	F _{3,56} = 4.9	0.004
Site x T20	F _{3,56} = 6.08	0.001
Intercept	F _{1,56} = 10.4	0.002

- 5 a. Factors considered included water table (WT), soil temperature at 5 cm (T5), soil temperature at 20 cm (T20), ground layer gross ecosystem photosynthesis (GEP), pore water CH₄ concentration (PW[CH₄])
- b. Results from the linear mixed effect models. All models included year, site, WT, T5, T20, GEP and two-way interaction with site, as fixed factors and plot as a random factor. The flux model also included $\log_{10}(\text{PW}[\text{CH}_4])$ and its interaction with site. Factors were removed sequentially starting with the highest p-value.
- 10 c. Reported R^2_{GLMM} is the marginal value, representing variation described by fixed factors only.

Table 4. Compilation of literature values for western Canadian fen methane flux

Fen type	Location	No. seasons measured, months covered	Mean CH₄ flux (std. error) mg CH₄ m⁻² d⁻¹	Reference
Open poor fen	56.38, 111.24	4, May-Oct	99.8 (269.7)	This study
Open poor fen	61.8, 121.4	1, Jul-Aug	162.5 (94.3) ^c	Liblik et al. 1997
Open poor fen	56.67, 113.53	1, May-Oct	23.5 (18.7)	Malhotra 2010
Open rich fen (graminoid)	61.8, 121.4	1, Jul-Aug	63.5 (36.2) ^c	Liblik et al. 1997
Open rich fen (low shrub)	61.8, 121.4	1, Jul-Aug	19.0 (13.9) ^c	Liblik et al. 1997
Open rich fen	55.85, 107.68	2, Jul-Oct	0.2 (0.1)	Turetsky et al. 2002
Open fen	54, 113	3, May-Oct	56.2 (11.2) ^a	Whiting & Chanton, 2001
Open saline fen	56.57, 111.28	4, May-Oct	91.3 (103)	This study
Patterned rich fen	53.77, 104.60	1, Apr-Oct	121.8 (85.9)	Rask et al. 2002
Patterned rich fen	53.95, 105.95	2, Apr-Oct	194.4 ^d	Sukyter et al. 1996
Treed poor fen	56.38, 111.24	4, May-Oct	68.3 (118.9)	This study
Treed moderate rich fen	56.94, 111.55	4, May-Oct	63.7 (103)	This study
Treed moderate rich fen	56.40, 116.89	2; May-Sep	30.4 (41.6)	Strack et al. 2018
Treed moderate rich fen	54.82, 113.52	1, May-Sep	25.4 ^b	Long et al. 2010
Treed moderate rich fen	56.94, 111.55	2; May-Aug	34.3 (18.4)	Murray et al. 2017b
Treed rich fen	61.8, 121.4	1, Jul-Aug	3.7 (3.8) ^c	Liblik et al. 1997

- a. Mean and standard error calculated from mean flux for each year presented in the publication
- b. Daily mean calculated from seasonal total measured by eddy covariance, divided by number of days in the study period. No uncertainty in the estimate was presented in the original publication
- c. Value in brackets represents standard deviation as reported in the original publication
- d. Measured with eddy covariance, no uncertainty presented in original publication

Figure captions

Figure 1: Daily mean water table position during the growing season over each study year. Water table is presented relative to the surface of hummocks at each site, where negative values indicate depth below the surface. When available, data from May 1 (day of year 121) to October 31 (day 304) is presented.
 25 Sensor failure in early 2013 at TRF resulted in a large data gap; manual measurements are plotted as points during this period.

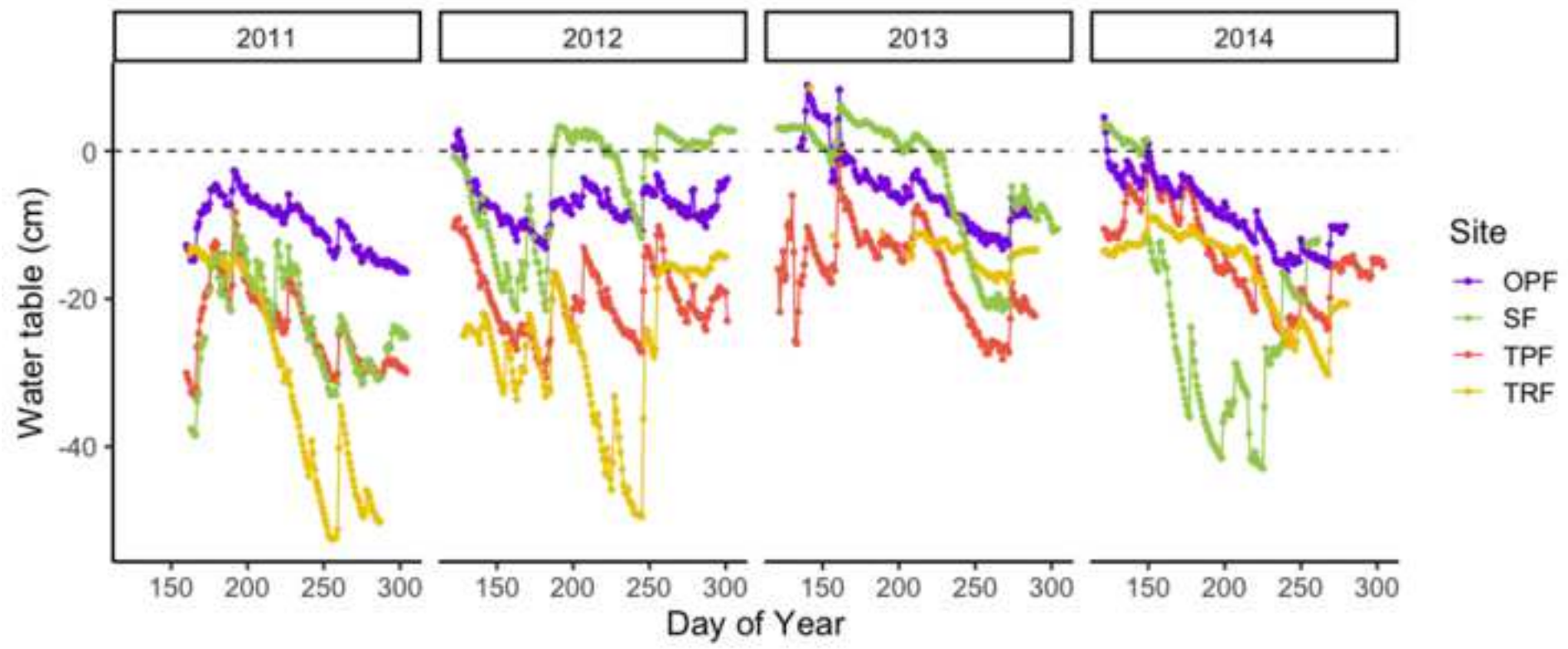
Figure 2: Measured CH₄ fluxes over the growing seasons 2011 – 2014 at each site. Sites are open poor fen (OPF), treed poor fen (TPF), treed rich fen (TRF) and open saline fen (SF). DOY is day of year. Note
 30 the log scale used on the y-axis. In order to apply the log scale to negative fluxes, a value of 10 was added to each flux; therefore, 10 represents a zero flux.

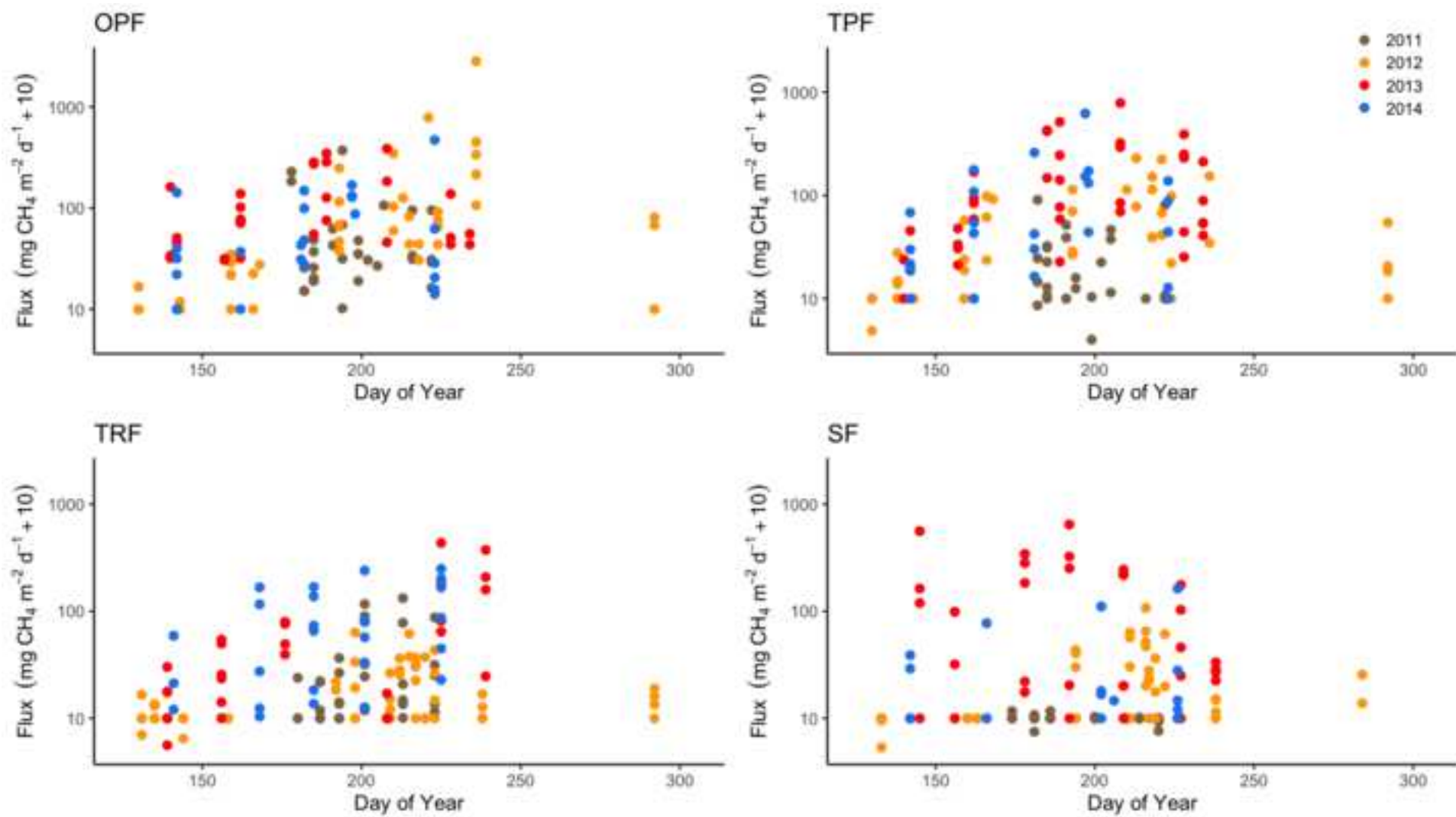
Figure 3. Mean log CH₄ fluxes (A) and mean log PW[CH₄] (B) at each site separated by years. Significantly different sites are labelled with no capital letter in common. Significantly different years are
 35 marked with no lower-case letter in common and should be compared within one site. The upper and lower edges of the boxes show 25th and 75th percentile, respectively, and the median (50th percentile) is located between them. The extent of the upper and lower whiskers away from the box to the most extreme data is no longer than 1.5 times the length of the box, thus outliers are present in the graphs.

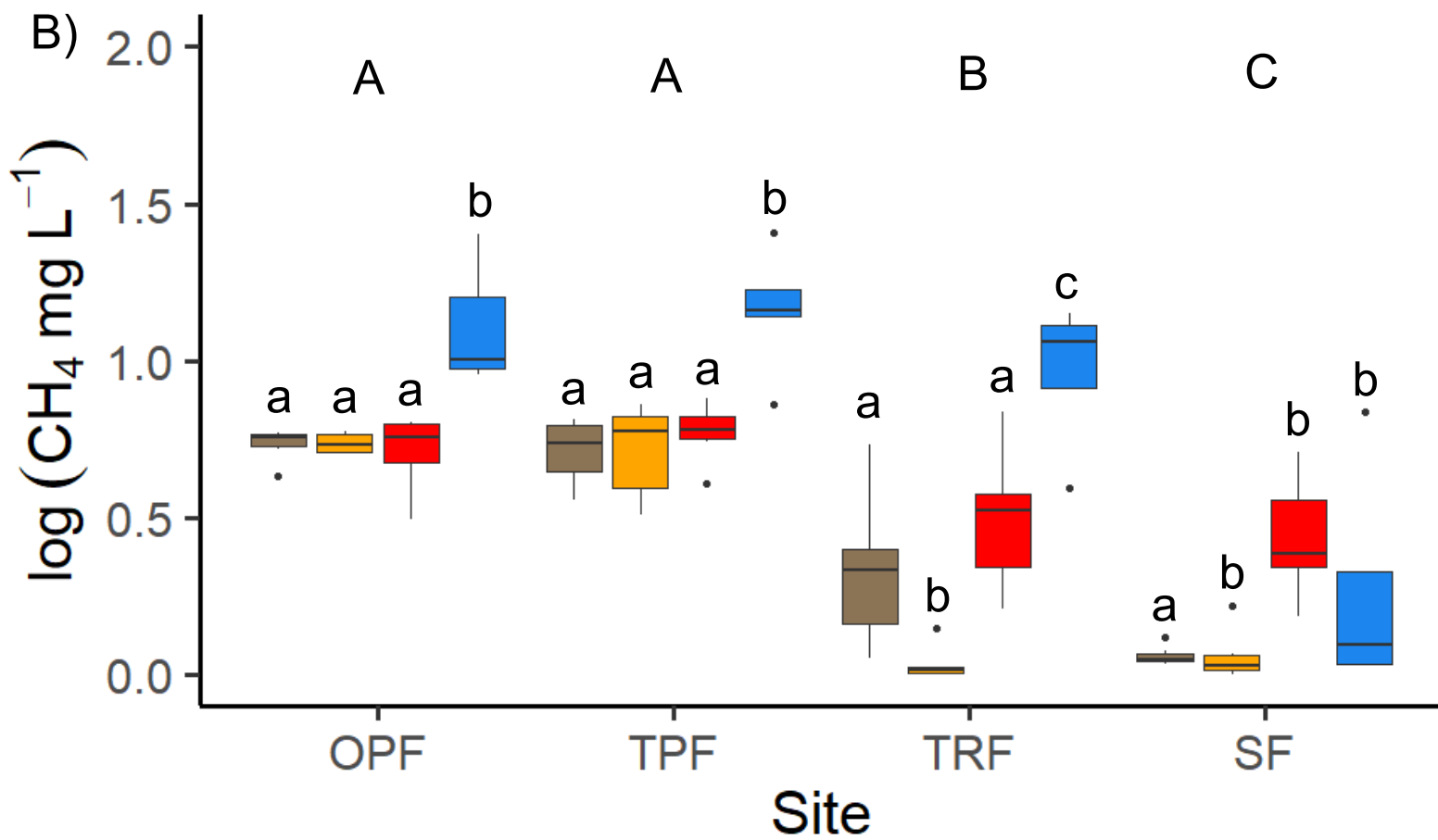
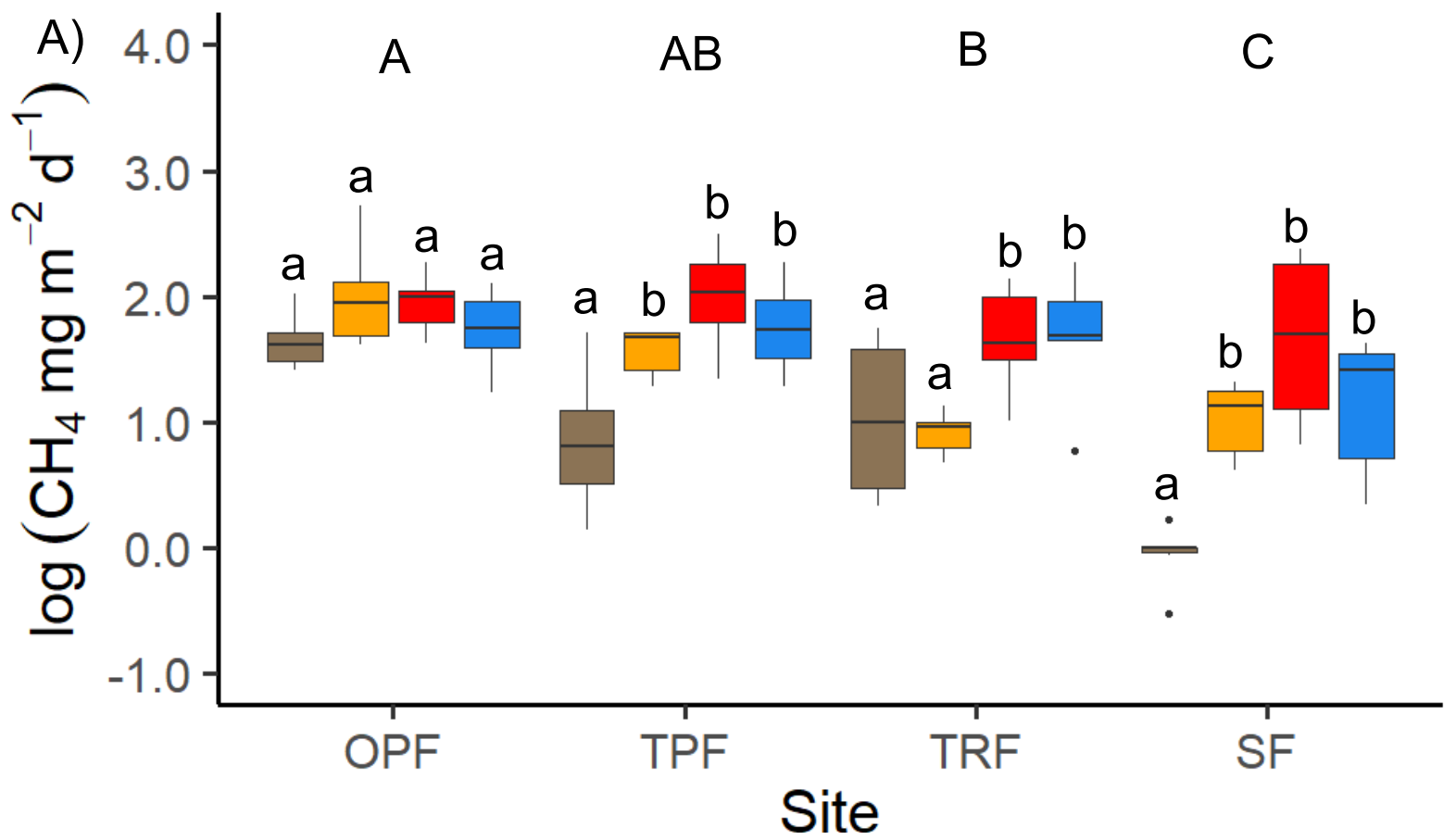
40 Figure 4. Measured pore water CH₄ concentration over the growing seasons of 2011-2014 at each site. Sites are open poor fen (OPF), treed poor fen (TPF), treed rich fen (TRF) and open saline fen (SF). DOY is day of year.

Figure 5. Water table relationship to CH₄ fluxes (A) and PW[CH₄] (B). A: $y=0.04x + 1.6415$, $r^2=0.2775$;
 45 B: $y=0.0184x + 0.6799$, $r^2=0.175$.

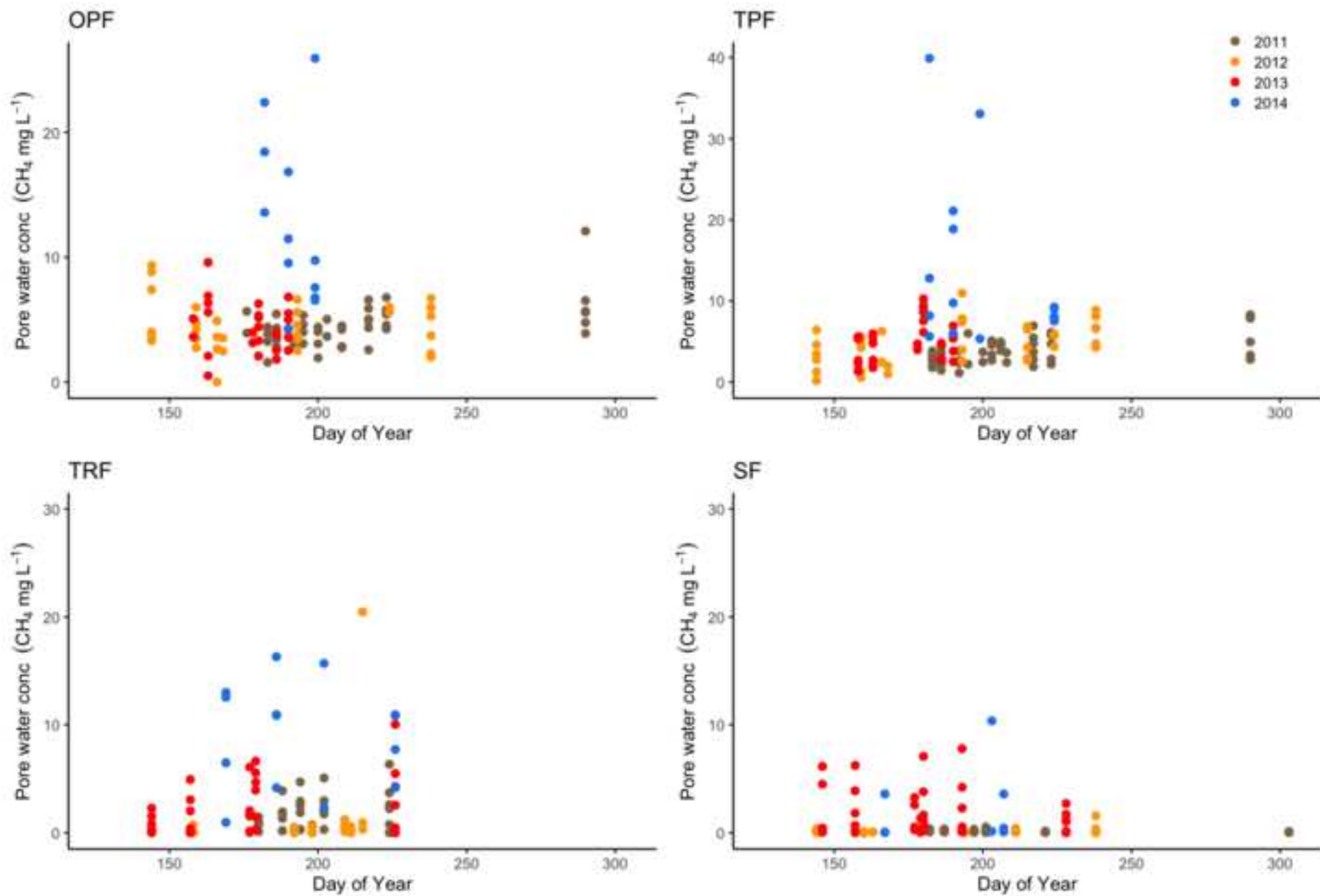
Figure 1

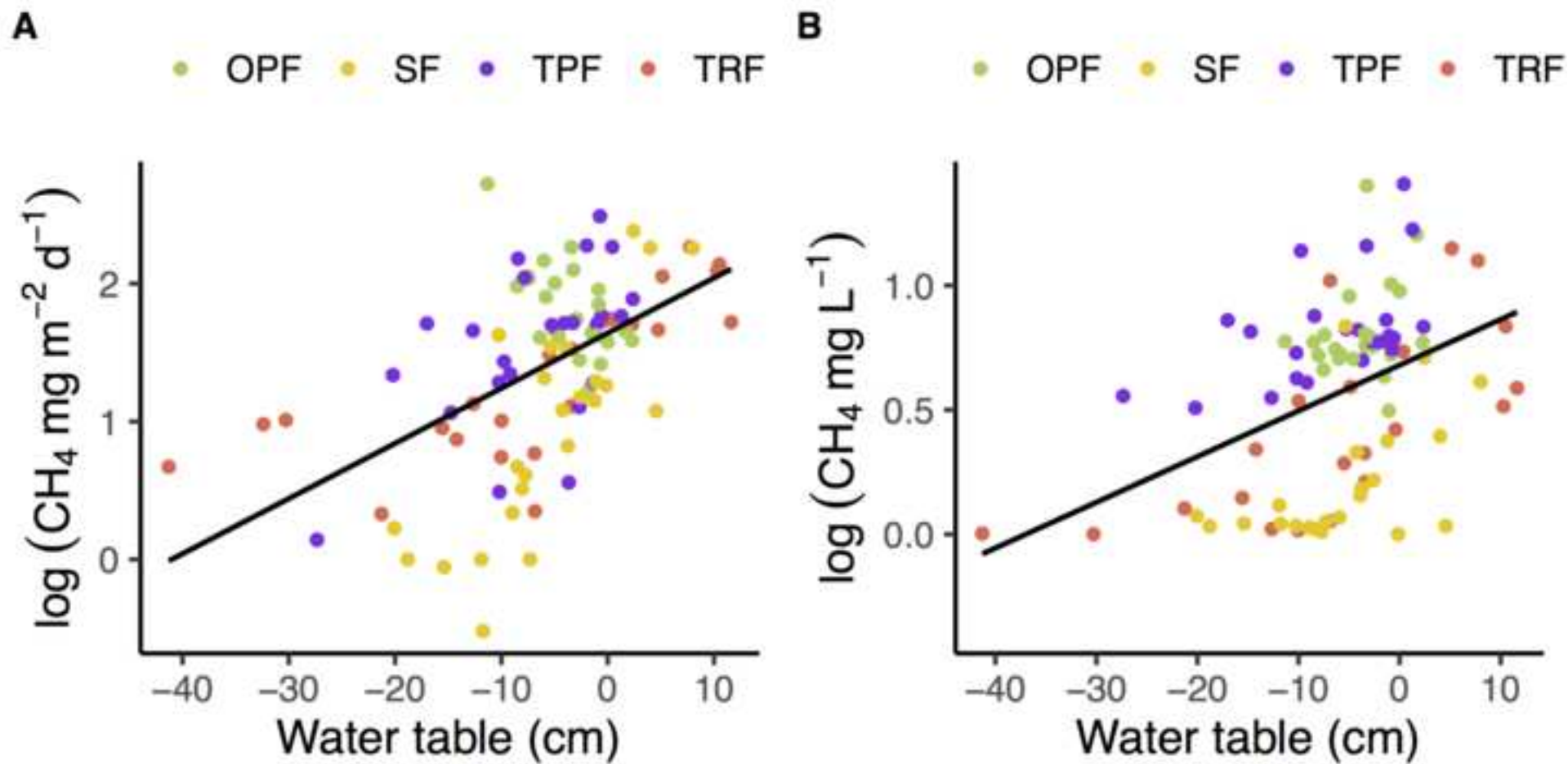






2011 2012 2013 2014





Methane emissions from fens in Alberta's boreal region: Reference data for functional evaluation of restoration outcomes

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