

1 Methane exchange in a poorly-drained black spruce
2 forest over permafrost observed using the eddy
3 covariance technique

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15 **Abstract**

16 Ecosystem-scale methane (CH₄) exchange was observed in a poorly-drained
17 black spruce forest over permafrost in interior Alaska during the snow-free
18 seasons of 2011–2013, using the eddy covariance technique. The magnitude
19 of average CH₄ exchange differed depending on wind direction, reflecting spa-
20 tial variation in soil moisture condition around the observation tower, due
21 to elevation change within the small catchment. In the drier upper posi-
22 tion, the seasonal variation in CH₄ emission was explained by the variation
23 in soil water content only. In the wetter bottom, however, in addition to
24 soil temperature and soil water content, seasonal thaw depth of frozen soil
25 was also an important variable explaining the seasonal variation in CH₄ ex-
26 change for this ecosystem. Total snow-free season (day of year 134–280) CH₄
27 exchanges were 12.0 ± 1.0 , 19.6 ± 3.0 , and 36.6 ± 4.4 mmol m⁻² season⁻¹ for

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28 the drier upper position, moderately wet area, and wetter bottom of the
29 catchment, respectively. Observed total season CH₄ emission was nearly one
30 order smaller than those reported in other northern wetlands, due proba-
31 bly to the relatively low ground water level and low soil temperature. The
32 interannual variation of total snow-free season CH₄ emission in the wetter
33 bottom of the catchment was influenced by the amount of rainfall and thaw
34 depth. On the other hand, in the drier upper position the amount of rainfall
35 did not strongly affect the total season CH₄ emission. Different responses of
36 CH₄ exchange to seasonal change in environmental conditions, depending on
37 the position of a small catchment, should be considered when estimating the
38 spatial variation in CH₄ exchange accurately in ecosystems over permafrost.
39 *Keywords:* Boreal forest, CH₄ flux, Path analysis, Spatial variability,
40 Thaw depth

41 **1. Introduction**

42 Methane (CH₄) is an important greenhouse gas, contributing about 20 %
43 to the total direct radiative forcing from long-lived greenhouse gases since
44 pre-industrial times (Forster et al., 2007). Clarifying the spatial and tempo-
45 ral variations of CH₄ exchange is thus of urgent importance for understanding
46 variations in atmospheric CH₄ concentration and its influence toward climate
47 changes.

48 Wetlands are identified as a major natural source of CH₄ (Matthews and
49 Fung, 1987; Bousquet et al., 2006; Schlesinger and Bernhardt, 2013). Many
50 studies were conducted to clarify the characteristics of CH₄ emission from
51 wetlands (e.g., Sebacher et al., 1986; Moore and Knowles, 1989; Whalen and

52 Reeburgh, 1990; Morrissey and Livingston, 1992; Bartlett and Harriss, 1993;
53 Harazono et al., 2006; Mastepanov et al., 2008; Olefeldt et al., 2013; Turet-
54 sky et al., 2014). Efforts have been made to clarify ecosystem-scale CH₄
55 emission from wetlands using the eddy covariance technique (e.g., Fan et al.,
56 1992; Verma et al., 1992; Friborg et al., 1997; Hargreaves et al., 2001; Sachs
57 et al., 2008; Zona et al., 2009; McDermitt et al., 2011; Pypker et al., 2013; Eu-
58 skirchen et al., 2014), and to develop CH₄ exchange components in ecosystem
59 models (e.g., Zhuang et al., 2004; Riley et al., 2011; Ringeval et al., 2011; Ito
60 and Inatomi, 2012). Despite these efforts, a recent review by Kirschke et al.
61 (2013) has suggested that CH₄ emissions from natural wetlands estimated
62 using ecosystem models were overestimated compared to results from inver-
63 sion models. A recent wetland model inter-comparison (Melton et al., 2013)
64 also showed large variations in CH₄ emissions between ecosystem models,
65 suggesting the need for improving the parameters and structures of ecosys-
66 tem models. Discrepancy between estimates clearly suggests the need for
67 more efforts to clarify the spatial and temporal variation in ecosystem-scale
68 CH₄ exchange for various wetland types, and for model validations based on
69 ecosystem-scale CH₄ exchange data.

70 In a boreal forest region, especially with lowlands and north-facing slopes,
71 permafrost is a characteristic soil condition. Ice-rich permafrost impedes
72 infiltration, and soils tend to be wet or saturated (Hinzman et al., 2006).
73 Thus, quite a large portion of boreal forest can be classified as wetland forest.
74 For example, in boreal Alaska, roughly 40–60% of the landscape is poorly
75 drained due to the presence of permafrost and characterized by shallow water
76 table conditions (Harden et al., 2003; Myers-Smith et al., 2007). A number

77 of studies (e.g., Crill et al., 1988; Whalen and Reeburgh, 1988; Moore et al.,
78 1990; Bartlett et al., 1992; Bubier et al., 1993; Dise, 1993; Moosavi et al.,
79 1996; Bellisario et al., 1999; Wickland et al., 2006; Turetsky et al., 2008;
80 Ullah et al., 2009; Matson et al., 2009) have conducted chamber observations
81 of CH₄ exchange in the boreal region in Alaska and Canada, and the seasonal
82 variations in CH₄ exchange and its dependence on environmental variables
83 such as soil temperature, soil moisture, and ground water table depth were
84 examined. These studies indicated that different environmental variables
85 appeared to affect CH₄ exchange at different temporal and spatial scales,
86 and general and quantitative relationships between environmental variables
87 and CH₄ exchange have not yet been found (Olefeldt et al., 2013).

88 One of the difficulties in studying CH₄ exchange is its heterogeneous
89 source/sink distributions with respect to both space and time, making it dif-
90 ficult to cover using chamber observations (Turetsky et al., 2014). Although
91 the chamber technique is useful for examining the influence of environmental
92 conditions on CH₄ exchange at the local scale, heterogeneous source/sink
93 distributions have hindered the accurate quantification of CH₄ exchange at
94 the ecosystem scale. Poorly-drained boreal forests typically show a large
95 spatial variability in soil moisture from the meter scale of tussock-hollow
96 microtopography to the few hundred-meter scale, due to elevation changes
97 within small catchments. Eddy covariance observation can provide such
98 ecosystem-scale CH₄ exchange data, covering a spatial area on a tens- to
99 hundreds-square-kilometer order (Baldocchi et al., 2001). The method is
100 thus useful for quantifications of CH₄ exchange and validations of ecosystem
101 models, especially in a boreal region. In addition, eddy covariance observa-

102 tion can obtain almost continuous data without disturbing the measurement
103 environment, which may provide detailed insight into a temporal variation in
104 CH₄ exchange. This insight will help to obtain more general and quantita-
105 tive relationship between environmental variables and CH₄ exchange. Eddy
106 covariance observations of CH₄ exchange in poorly-drained boreal forests,
107 however, has been seldom reported in the literature.

108 We applied the eddy covariance technique here to observe ecosystem-scale
109 CH₄ exchange in a poorly-drained black spruce forest over permafrost in inte-
110 rior Alaska for three snow-free seasons. Black spruce is the dominant species
111 in the Interior, and tends to grow in poorly-drained lowland over permafrost.
112 Our objectives here are 1) to clarify the variations in ecosystem-scale CH₄ ex-
113 change from diurnal to interannual time scale, 2) to identify the influence of
114 environmental conditions on ecosystem-scale CH₄ exchange, and 3) to quan-
115 tify the total CH₄ exchange during snow-free period in this poorly-drained
116 black spruce forest using eddy covariance flux data. To our knowledge, very
117 few studies have reported on an interannual variation of ecosystem-scale CH₄
118 exchange in boreal and arctic region. This study presents new information
119 regarding how ecosystem-scale CH₄ exchange during snow-free period re-
120 sponds to seasonal and interannual variations in environmental conditions in
121 a boreal forest using observations of three snow-free seasons.

122 **2. Observations and Data Analyses**

123 *2.1. Study Site*

124 Data were obtained in a poorly-drained black spruce (*Picea mariana*)
125 forest (64°52'N, 147°51'W, 159 m a.s.l.), standing on ice-rich permafrost, in

126 Fairbanks, Alaska, USA (Ueyama et al., 2006, 2009, 2014; Iwata et al., 2010).
127 Mean tree age is approximately 90 years (Ueyama et al., 2015), and tree
128 height typically ranges from 1 to 5 m, though there are sparsely distributed
129 taller trees of more than 6 m. Tree density is 4500 trees ha⁻¹; however, due
130 to the narrow canopy architecture of black spruce, the forest canopy is rel-
131 atively open. The forest floor has a pronounced tussock-hollow microtopog-
132 raphy, and standing water is seen in the hollows when ground water level is
133 high. The understory is dominated by low evergreen shrubs (*Ledum groen-*
134 *landicum*, *Vaccinium vitis-idaea*), deciduous shrubs (*Vaccinium uliginosum*,
135 *Rubus chamaemorus*, *Betula glandulosa*), and sedges (*Carex* species). The
136 ground is almost completely covered with mosses (*Sphagnum* and feather
137 mosses). Leaf area index (LAI) of black spruce and understory vegetation,
138 measured with a plant canopy analyzer (LAI-2000, Li-Cor, USA), varied
139 from 0.2 m² m⁻² during snow season to 1.9 m² m⁻² during mid-summer. Soil
140 is silt-loam overlain by an organic layer of 25–45 cm (Heijmans et al., 2004),
141 and is poorly drained due to the presence of ice-rich permafrost. The pH
142 of ground water above the frozen soil layer was 5–6. Active layer depth was
143 40–50 cm (Iwata et al., 2012).

144 The observation tower was located near the bottom of a gentle northwest-facing
145 slope (of approximately one degree; Fig. 1) within a small catchment. North-
146 ward is the bottom of the small catchment, which is flat and extends approxi-
147 mately 200 m from the tower. The terrain gains elevation again to the further
148 north, at approximately one degree. Snowmelt and rain water flows within
149 the surface soil, following topography and converging to the west of the tower,
150 and then flowing into a lake located 270 m west of the tower. As a result,

151 the western portion tends to be wetter than others. *Sphagnum* moss is the
152 typical surface cover there. Ground water level was generally below ground,
153 except for a short period just after snowmelt. To the south, on the other
154 hand, ground water was not observed due to higher elevation. The ground
155 in that area is typically covered with feather moss and lichen.

156 Mean monthly air temperature in Fairbanks between 1971 and 2000 ranged
157 from -23.2°C in January to 16.9°C in July, and the mean annual precipita-
158 tion was 263 mm yr^{-1} (Shulski and Wendler, 2007). The observation site was
159 typically free of snow from late April through early October.

160 2.2. Observation

161 CH_4 flux was observed using the closed-path eddy covariance technique
162 during three snow-free seasons (early May through early October), in 2011–
163 2013. An ultrasonic anemo-thermometer (CSAT3, Campbell Scientific, USA)
164 was attached to a 10-m aluminum tower (UT930, Campbell Scientific, USA)
165 at a height of 6 m above ground. Sample air was drawn from the same height
166 as the anemo-thermometer, and fed to a closed-path CH_4 analyzer (RMT-200
167 Fast Methane Analyzer or Greenhouse Gas Analyzer, Los Gatos Research
168 Inc., USA) placed in a box on the forest floor. The air inlet was placed at
169 a distance of 0.4 m from the measurement path of the anemo-thermometer,
170 and polyethylene tubing with 9.5 mm inner diameter was used for sampling.
171 An external pump was placed at the end of the flow line to draw sample
172 air. Two buffer tanks with $1.3 \times 10^{-3}\text{ m}^3$ volume were inserted between the
173 CH_4 analyzer and the external pump to reduce pressure fluctuation in the
174 sample air. Fluctuation in water vapor concentration for the sampled air was
175 suppressed using a Nafion dryer (PD-200T-48, Perma Pure, Inc., USA). All

176 data were recorded at 10 Hz using a datalogger (CR3000, Campbell Scientific,
177 USA). During the observation period of three years, the observation system
178 was gradually modified, aiming to improve the flux accuracy. The detailed
179 observation system and its modifications are shown in Table 1. Sampling
180 air flow rate was increased by changing the external pump and modifying
181 the flow line. The resultant refreshing time of measurement cell and the
182 Reynolds number of air flow in the sampling tube was, respectively, 2.4 s and
183 1490 for 2011, 1.9 s and 1940 for 2012, and 0.8 s and 4620 for 2013.

184 Relevant micrometeorological observations were also conducted at the
185 tower, including air temperature and relative humidity (HMP45AC or HMP155,
186 Väisälä, Finland) at 2 m, solar radiation (CMP3 or CNR4, Kipp & Zonen,
187 The Netherlands) at 6 m, and atmospheric pressure (PTB101B, Väisälä, Fin-
188 land) at 1 m. Within 20-meter distance from the tower, rainfall (TR-525M-R3,
189 Texas Electronics, USA) at 1 m, volumetric soil water content (CS616, Camp-
190 bell Scientific, USA) for mean values at 0–0.1, 0.1–0.2, and 0.2–0.3 m depth,
191 soil temperature (thermocouple) at 0.1 and 0.2 m depths at one location,
192 ground water level (CS445 and CS450, Campbell Scientific, USA) at two
193 locations were all measured. All data were scanned every ten seconds, with
194 half-hourly mean values stored in dataloggers (CR10X and CR23X, Camp-
195 bell Scientific, USA). Thaw depth was measured manually by inserting a
196 metal rod into the soil approximately once a week. This measurement was
197 conducted with five measurements at each of ten plots.

198 *2.3. Data Processing*

199 *2.3.1. Flux Calculation and Corrections*

200 Covariances of vertical wind velocity (m s^{-1}), w , and CH_4 density ($\mu\text{mol m}^{-3}$),
201 m , were calculated for half-hourly intervals from the raw 10 Hz data. Prior
202 to covariance calculation, removal of spike noises (Vickers and Mahrt, 1997),
203 coordinate rotation of wind velocities (double rotation), and synchronization
204 of CH_4 density data to wind velocity (Moncrieff et al., 1997) were performed.
205 The median delay time derived from a certain time period (typically, one
206 month) was applied for the synchronization of the whole period. Depen-
207 dence of delay time on relative humidity (Ibrom et al., 2007) was not ob-
208 served. In addition, a low-pass filter with two-sec running mean was applied
209 to CH_4 density data, to suppress high-frequency instrumental noise, which
210 facilitated the determination of transfer functions described below.

211 The high-frequency loss of CH_4 flux was corrected with an empirical trans-
212 fer function approach. Transfer functions were determined against cospec-
213 tra of w and T_{sv} . Figure 2 indicates the cospectral ratios of CH_4 flux to
214 sensible heat flux. Although the scatter was large, median values follow a
215 smooth decline with increasing frequency. The transfer function was deter-
216 mined by fitting an equation, $y = 1/(1 + ax^b)$ where a and b are parameters,
217 to median values. Parameters were changed every time the eddy covari-
218 ance system was modified. These transfer functions were combined with an
219 empirical cospectral model, which depends on wind speed and atmospheric
220 stability, to estimate the magnitude of high-frequency loss. More details can
221 be found in Iwata et al. (2014). The cut-off frequency of transfer function
222 was 0.22, 0.22, and 0.08 Hz for May/2011–June/17/2012, June/17/2012–

223 September/2012, and 2013, respectively. The mean correction coefficients
224 and its standard deviations for each period above were 1.5 ± 0.3 , 1.3 ± 0.2 , and
225 1.9 ± 0.6 , respectively. Unfortunately, the faster flow rate in 2013 decreased
226 the signal-to-noise ratio, probably resulted in the lower cut-off frequency and
227 the larger correction coefficients.

228 CH_4 exchange was evaluated as the sum of turbulent flux and storage
229 within the atmospheric column below observation height. Storage was esti-
230 mated from changes of CH_4 density over a half-hourly period at the height
231 of the eddy covariance observation.

232 *2.3.2. Data Selection Criteria and Gap-Filling*

233 Data used in the analysis were selected from visual inspection of raw
234 10-Hz data, data quality criteria, footprint analysis, and u_* thresholding.
235 First, all 10-Hz data were checked visually for malfunction of instruments.
236 Next, spikes in the raw data were removed using a method of Vickers and
237 Mahrt (1997). Data with abundant spikes, large discontinuities, and strong
238 non-stationarity were discarded (Vickers and Mahrt, 1997; Mahrt, 1998).
239 The fetch contributing 80 % of observed flux was calculated using a footprint
240 model by Kormann and Meixner (2001), and data were selected so that
241 the 80 % fetch did not overlap the non-black spruce area. Then, data with
242 $u_* < 0.10 \text{ m s}^{-1}$ were rejected, for insufficient turbulence conditions (e.g.,
243 Rinne et al., 2007; Zona et al., 2009; Long et al., 2010). A sensitivity test for
244 threshold value showed that total CH_4 exchange over the observation period,
245 calculated from gap-filled data (described below), was clearly underestimated
246 when data with $u_* < 0.10 \text{ m s}^{-1}$ were included in the analysis, and that even
247 increasing the threshold value above 0.10 m s^{-1} did not change the total CH_4

248 exchange.

249 To obtain daily and seasonal total CH₄ exchange, data gaps were filled
250 using the multiple imputation technique (Rubin, 1987; Hui et al., 2004; En-
251 ders, 2010). This technique generates several data sets by filling gaps with a
252 different estimation for each data set, based on multiple regression and un-
253 certainty of the regression. Twenty data sets were generated and combined
254 to obtain daily total CH₄ exchange and its uncertainty due to gap-filling
255 (Enders, 2010). As independent variables, solar radiation, thaw depth, soil
256 temperature at 0.2 m depth, and volumetric soil water content between 0.1
257 and 0.2 m below ground were used. Thaw depth data was linearly interpo-
258 lated, and a constant value was assigned for a day. Gap-filling was applied
259 to data using a one-month moving window to fill the gaps from a local rela-
260 tionship between CH₄ exchange and environmental variables. The multiple
261 imputation was performed with an *Amelia II* package (Honaker et al., 2013)
262 in the *R* statistical software. Comparisons of artificially-removed half-hourly
263 data and imputed half-hourly data for certain one-month intervals suggested
264 that the imputed data can, at least, reconstruct the average CH₄ exchange
265 over one-month time scale, but individual imputed half-hourly data might
266 have large uncertainty. This was, in part, because clear relationships between
267 CH₄ exchange and environmental variables were not found using half-hourly
268 data due to large uncertainties of observed flux for its small flux magnitude
269 (Appendix A). Thus, we focused on averaged CH₄ exchange over half months
270 to examine its temporal variations and the influence of environmental con-
271 ditions on the exchange.

272 We observed infrequent spike-like CH₄ emission, which met the data selec-

273 tion criteria above. These spike-like CH₄ emission data were excluded when
274 constructing the relationship between CH₄ exchange and environmental vari-
275 ables in the gap-filling procedure, but were retained in calculating the final
276 total CH₄ exchange as spike-like CH₄ emission is possible when ebullition oc-
277 curs. Data were identified as the spike-like CH₄ emissions when half-hourly
278 CH₄ exchange exceeded three times the typical uncertainty due to random
279 error from the local median value of CH₄ exchange within a two-week moving
280 window.

281 *2.4. Analysis*

282 We first examined the spatial variability of CH₄ exchange by analyzing
283 its relationship with wind direction and footprint area. This revealed that
284 the magnitude of CH₄ exchange was different across the direction sectors,
285 and we thereafter separated CH₄ exchange data into groups according to
286 wind direction. However, we also used identical environmental variables in
287 analyzing each group of data, as information regarding spatial variations in
288 environmental variables between areas was not available. We assumed that
289 seasonal variation in environmental variables such as soil temperature and
290 moisture are correlated between areas, and thus variables obtained at a single
291 place are adequate for explaining seasonal variation in CH₄ exchange in all
292 areas.

293 We applied path analysis (Schemske and Horvitz, 1988; Bassow and Baz-
294 zaz, 1998) to CH₄ exchange data, in order to examine the relative importance
295 of environmental variables controlling CH₄ exchange. Path analysis is an ex-
296 tension of multiple regression and is useful when independent variables have
297 a causal or correlated relationship. In path analysis, a hypothesized model

298 (i.e., causal connections between variables) is constructed, and path coeffi-
299 cients are calculated by fitting the model to data. A path coefficient is a
300 standardized partial regression coefficient, and represents the magnitude of
301 the direct effect of the independent variable on the dependent variable, with
302 all other independent variables held constant (Schemske and Horvitz, 1988).
303 In this study, the model was constructed to evaluate relative importance
304 of soil temperature, soil water content, and seasonal thaw depth on CH₄
305 exchange. Rather than finding the best model by including other environ-
306 mental variables, we intended to identify the change in importance of these
307 fundamental variables under different conditions. The adequacy of the model
308 was determined using a goodness-of-fit index. When the goodness-of-fit in-
309 dex was greater than 0.8, the model was considered adequate. Path analysis
310 was performed with a *sem* package (Fox, 2006) in the *R* statistical software.

311 **3. Results**

312 *3.1. Environmental Conditions*

313 Among environmental variables, rainfall and soil water conditions var-
314 ied distinctly across the three seasons. The 2011 season, defined as day of
315 year (DOY) 120–270, had total rainfall of 195.1 mm (Table 2), which was
316 17% larger than the 1971–2000 mean (Shulski and Wendler, 2007), with rain
317 events occurring more frequently compared to other years. As a result, vol-
318 umetric soil water content was higher than other years, and ground water
319 level did not decline below 0.35 m during mid-summer (Fig. 3). Contrarily,
320 the 2013 season had less rainfall, at 146.1 mm—13% less than the 1971–2000
321 mean. Rainfall was especially low in the early half of the season: no rainfall

322 was observed during DOY 200–231. As a result, mean volumetric soil water
323 content at 0.1–0.2 m depth during DOY 200–231 was 0.54, while in the other
324 years, this value was close to 0.90. Subsequently, volumetric soil water con-
325 tent in 2013 increased due to increased rainfall. The 2012 season had total
326 rainfall comparable to 2011, though there was a long period with little rain-
327 fall during DOY 207–236. Volumetric soil water content and ground water
328 level for 2012 declined during this period.

329 As for temperature, spring 2013 was unusually cold, and snow melt was
330 delayed by about a month compared to typical years. As a result, soil thaw
331 also started later, compared to the other two years (Fig. 3). After snow melt,
332 air temperature rose rapidly, reaching 25.5 °C on DOY 177, and resulting in
333 a larger range for air temperature in 2013 (Table 2). Soil temperature in
334 summer of 2013 was also higher than the other two years.

335 Maximum thaw depth was largest in 2011 (Table 2). Higher soil water
336 content enhanced thermal conductivity in the soil (Brown, 1963) in 2011,
337 resulting in the largest thaw depth at the end of the season (0.40 m). In
338 2013, contrarily, maximum thaw depth was the lowest (0.33 m). This is due
339 to drier soil and delayed soil thawing.

340 *3.2. Spatial Variability of Methane Exchange*

341 Spatial variability in CH₄ exchange reflected the expected spatial varia-
342 tion of soil water conditions due to topography around the observation tower
343 (Fig. 4 and Table 3). CH₄ emissions tended to be higher in the 240–300 di-
344 rection (Table 3), where soil was considered to be wetter. To the northwest
345 (300–360), CH₄ emission tended to be comparable to western emissions. In
346 comparison, the southern area emitted less CH₄, because the area (120–240

347 direction) shows higher elevation and relatively dry soil. Ground water above
348 the frozen soil layer was not seen in this area. The northeast (0–120 direc-
349 tion) showed intermediate CH₄ emission levels. Footprint analysis showed
350 no relationship between CH₄ exchange and footprint area within the wind
351 direction sectors, suggesting the surface heterogeneity within the direction
352 sectors did not influence the variability in CH₄ exchange. Hence, we sep-
353 arated data according to wind direction into 0–120, 120–240, and 240–360
354 directional sectors. Hereafter, sectors of 0–120, 120–240, and 240–360 degree
355 from the tower are referred to as moderately wet, drier, and wetter areas,
356 respectively.

357 *3.3. Methane Exchange and Environmental Conditions*

358 CH₄ exchange in this ecosystem had indiscernible diurnal variations (Fig. 5).
359 Most diurnal variation was within a 95 % confidence interval for all areas. For
360 May–June, the difference in median values between areas was not obvious,
361 though median values for CH₄ exchange tended to be higher for the wetter
362 area. For July–August, CH₄ emissions tended to increase in all areas. The
363 increases were larger in the wetter area, and CH₄ emission from the wetter
364 area was clearly higher than in the drier area. Although Fig. 5 showed 2012
365 data only, other years showed similar patterns.

366 CH₄ emission generally increased from May to August, and at the end
367 of the season, CH₄ emission showed a decreasing tendency compared to
368 mid-summer (Fig. 6). The magnitude of seasonal variation was largest for
369 the wetter area. In the wetter area, CH₄ emission clearly increased from
370 May/June to July/August in 2011 and 2012: average CH₄ emission was
371 1.9 ± 0.1 and 1.2 ± 0.4 nmol m⁻² s⁻¹ in May/June, and 3.6 ± 0.8 and $3.3 \pm$

372 0.1 nmol m⁻² s⁻¹ for July/August of 2011 and 2012, respectively. The grad-
373 ual increase in CH₄ emission until August was observed in 2011 and 2012
374 with increases in thaw depth, soil temperature, and soil moisture. At the
375 end of season, CH₄ emission decreased, suggesting that CH₄ emission was
376 likely suppressed by low soil temperature. In 2013, CH₄ emission reduced
377 significantly in the latter half of August (1.4 nmol m⁻² s⁻¹), corresponding
378 with the end of the drought period of 2013. Similar but somewhat smaller
379 seasonal variations were also observed in the moderately wet area. In the
380 drier area, seasonal variation was the smallest of all areas.

381 Linear regression analysis was applied in order to examine the relationship
382 between CH₄ exchange and environmental variables, based on half-monthly
383 average data (Table 4). Analysis showed that CH₄ exchange was positively
384 correlated with thaw depth and soil water content at 0.1–0.2 m and 0.2–0.3 m
385 depths for all areas. These correlations were less strong for drier area data
386 than in other areas. In the wetter and moderately wet areas, soil temperature
387 at 0.2 m depth and soil water content at 0–0.1 m depth were also positively
388 correlated with CH₄ exchange. No significant correlation was found between
389 soil temperature at 0.1 m depth and CH₄ exchange.

390 Soil temperature dependence of CH₄ emission showed a complicated pat-
391 tern (Fig. 7). For example, in 2011 the relationship between CH₄ emission
392 and soil temperature displayed a hysteresis pattern, with an emission peak
393 observed in the early half of September, when soil temperature had already
394 begun to decline. In addition, CH₄ emission in the late season was slightly
395 larger than early-season emission, although soil temperature was similar. In
396 2012, an emission peak was not obvious, though the pattern was similar to

397 2011. The 2013 season showed a rather different pattern: CH₄ emission re-
398 duced from July to August, due probably to decreased soil moisture content
399 (Fig. 3). As a result, emission was largest when soil temperature was highest,
400 in the first half of July, 2013.

401 Path analysis effectively revealed that the environmental variables con-
402 trolling CH₄ exchange varied between different areas (Fig. 8). In the drier
403 area, soil water affected CH₄ exchange (path coefficient of 0.68, $p = 0.13$),
404 and effects from thaw depth and soil temperature were only marginal. In
405 the wetter area, thaw depth was most important (path coefficient of 0.44,
406 $p = 0.29$). The path coefficient for soil temperature on CH₄ exchange for the
407 wetter area was 0.17 ($p = 0.20$), which was greater than in the drier area
408 (-0.06 , $p = 0.69$). On the other hand, the path coefficient for soil water
409 content for the wetter area (0.20, $p = 0.62$) was lower than in the drier area.
410 Thus, the relative importance of soil water content was higher in the drier
411 area, while that of thaw depth was higher in the wetter area. In the moder-
412 ately wet area, the path coefficients of three variables for CH₄ exchange took
413 values between those of the drier and wetter areas.

414 *3.4. Total Snow-Free Season Methane Exchange*

415 Total snow-free season CH₄ emission (Table 5) was also, on average, great-
416 est in the wetter area, followed by the moderately wet area. For the wet-
417 ter area, total CH₄ emission was greatest in 2011 (45.0 mmol m⁻² season⁻¹),
418 when deepest thaw depth and highest soil water content were observed. The
419 2013 season had lowest total CH₄ emission (30.3 mmol m⁻² season⁻¹), while
420 total CH₄ emission in the 2012 season (34.6 mmol m⁻² season⁻¹) was slightly
421 greater than in 2013. In the drier area, total season CH₄ emission did not

422 vary largely across the three years ($12.0 \pm 1.7 \text{ mmol m}^{-2} \text{ season}^{-1}$).

423 **4. Discussion**

424 In general, CH_4 exchange responds to environmental conditions such as
425 soil temperature and soil moisture (e.g., Sebacher et al., 1986; Crill et al.,
426 1988; Christensen et al., 1995; von Fischer et al., 2010), as CH_4 is produced
427 by methanogens and consumed by methanotrophs, and the activity of these
428 bacteria is influenced by temperature and oxygen availability. In the diurnal
429 cycle, soil temperature can be the main controlling variable, as moisture con-
430 dition does not change over single days. CH_4 exchange in the poorly-drained
431 black spruce forest, however, did not show any obvious diurnal variations
432 (Fig. 5). Similar results have been reported in a boreal fen (Rinne et al.,
433 2007), an Arctic wet tundra (Harazono et al., 2006; Tagesson et al., 2012),
434 and a sub-boreal peatland (Pypker et al., 2013). At our site, the lack of
435 obvious diurnal variation in CH_4 exchange may suggest that most CH_4 was
436 likely produced in a deeper active layer soil, where diurnal variation in soil
437 temperature was not significant (Moosavi and Crill, 1997). The ground cov-
438 ered by moss showed low thermal conductivity and the typical magnitude
439 of daily soil temperature variation was about 1°C at 0.2 m depth (data not
440 shown), with ground water table below this depth during most of the ob-
441 servation period (Fig. 3). In contrast, CH_4 oxidation was also expected to
442 occur in aerobic surface soil, where temperature variation was larger than in
443 deeper soil, and to increase during daytime. A part of this expected increase
444 in CH_4 oxidation could be canceled out by an increase of CH_4 production
445 during daytime, depending on the strength of both CH_4 production and ox-

446 idation, and the magnitude of their temperature dependence (Segers, 1998;
447 van Winden et al., 2012).

448 At the bottom of the small catchment, soil tended to be wetter, with
449 ground water below the soil surface. Here, thaw depth was the most impor-
450 tant variable controlling CH₄ emission (Fig. 8). Thaw depth is a variable,
451 integrating conditions favorable for CH₄ production (Whalen and Reeburgh,
452 1992). Sturtevant et al. (2012) and Kim (2015) also showed that thaw depth
453 is a key environmental variable in regulating CH₄ exchange in Arctic tundra,
454 although their studies showed far stronger sensitivity of CH₄ emission to in-
455 crease in thaw depth when thaw depth was more than 30 cm. Soil thaw likely
456 regulated the vertical extent to which methanogens can be active, thus influ-
457 encing the base CH₄ emission rate. Microbial population could also increase
458 later in the season compared to spring season (Funk et al., 1994; Moosavi
459 et al., 1996; van Hulzen et al., 1999). These factors can explain the higher
460 CH₄ emission rate later in the season, though soil temperature was similar
461 (Fig. 7). The relative importance of soil temperature was also higher in the
462 wetter area than the drier area. This may be because soil was wet for most
463 of the observation period in the wetter area, with soil temperature more ef-
464 fective in enhancing methanogen activity than in the drier area (Morrissey
465 and Livingston, 1992; Moosavi et al., 1996; Olefeldt et al., 2013). Jackowicz-
466 Korczyński et al. (2010) and Parmentier et al. (2011) similarly reported that
467 responses of CH₄ emission to environmental variables varied depending on
468 vegetation and surface conditions in a single eddy covariance site.

469 Permafrost condition may also affect the spatial variability of CH₄ ex-
470 change. Degradation of permafrost in the northwestern portion has occurred

471 under inundated standing water there; on the other hand, permafrost in the
472 southern area is still stable (V. Romanovsky, 2014, personal communication).
473 In this poorly-drained black spruce forest, higher CH₄ emission was observed
474 in the northwestern area, where degradation of permafrost had occurred. Fur-
475 ther degradation of permafrost may enhance CH₄ emission (Olefeldt et al.,
476 2013) in this ecosystem.

477 Even in the drier area, where no ground water was present above the
478 frozen soil layer, small net CH₄ emission was observed in our black spruce
479 forest (Fig. 6 and Table 5). This suggests that CH₄ was also produced within
480 an anaerobic microsite in the unsaturated soil (von Fischer and Hedin, 2002;
481 Blankinship et al., 2010). The increase in soil water content likely extended
482 the anaerobic microsite area, resulting in enhanced CH₄ emission later in the
483 season (Fig. 6 and 8).

484 Total snow-free season CH₄ emission in this poorly-drained black spruce
485 forest (Table 5) was nearly one order smaller than emissions reported for
486 other northern wetland ecosystems: e.g., 788 mmol m⁻² over one year in a
487 boreal fen in southern Finland (Rinne et al., 2007), 633 mmol m⁻² over
488 four months in an Arctic tundra in Greenland (Tagesson et al., 2012), 258–
489 515 mmol m⁻² over four months in an Arctic wet tundra (Harazono et al.,
490 2006), 311 mmol m⁻² over six months in a collapsed scar bog in Alaska (Eu-
491 skirchen et al., 2014), 200 mmol m⁻² over four months in a peatland in south-
492 ern Canada (Long et al., 2010), and 121 mmol m⁻² over three months in an
493 Arctic tundra in northern Siberia (Sachs et al., 2008). The relatively low
494 ground water level (approximately 0.2–0.4 m below ground; Fig. 3) was at-
495 tributable to low CH₄ emission in this forest. The tundra sites in colder

496 climate listed above had both higher ground water levels and CH₄ emissions.
497 At landscape scale, spatial variation of CH₄ emission was reported to be
498 related to that of ground water level (Sebacher et al., 1986; Olefeldt et al.,
499 2013). The low CH₄ emission in this forest also indicates that quite a large
500 fraction of CH₄ produced in deeper soil could be consumed in aerobic sur-
501 face soil and *Sphagnum* moss layer while transported to the atmosphere by
502 diffusion (Conrad and Rothfuss, 1991; Whalen et al., 1996; Kip et al., 2010).
503 In addition, soil temperature at this site was relatively low underlain by per-
504 mafrost, thus constraining CH₄ emission, compared to wetland sites without
505 permafrost in warmer climates (Turetsky et al., 2014).

506 So far, few studies have reported the interannual variation in CH₄ ex-
507 change in a boreal and arctic wetland. Our study showed that the CH₄
508 emission of the wetter area within the forest was the largest in 2011 (Ta-
509 ble 5), a year with the largest amount of rainfall and the greatest thaw depth
510 among the three years (Table 2). The combination of large amount of rainfall
511 and deep thaw led to greater vertical extent of anaerobic soil layer, and thus
512 potentially enhanced the CH₄ production. Similarly, a few studies (Parmen-
513 tier et al., 2011; Tagesson et al., 2012; Brown et al., 2014) in the literature
514 reported a higher CH₄ emission in a wetter year from a two-year observation.
515 On the other hand, our study showed that the interannual variation in CH₄
516 emission in the drier area was insignificant regardless of the amount of rain-
517 fall. The lower amount of rainfall in 2013 resulted in lower soil water content
518 in the soil surface layer (Fig. 3). However, the deeper soil layer where CH₄
519 was presumably produced was relatively unaffected due to water input from
520 soil thawing, especially later in the season. Thus, the interannual variation

521 in CH₄ emission in the drier area was not strongly affected by the amount of
522 rainfall.

523 Finally, we discuss the contribution of CH₄ emission to the greenhouse
524 gas budget in this black spruce forest. In 2011, the same forest emitted CO₂
525 of 5.5 mol m⁻² y⁻¹ (Ueyama et al., 2014). Whereas, the moderately wet area
526 of the black spruce forest emitted CH₄ of 23.2 mmol m⁻² in 2011 snow-free
527 season (Table 5). Kim et al. (2007) reported from an observation conducted
528 in 2005–06 winter in the same forest that the CH₄ emission was 9.4 mmol m⁻²
529 during winter time. Assuming the similar winter CH₄ emission in 2011, we
530 estimated that the annual CH₄ emission in 2011 could be 32.6 mmol m⁻²,
531 which was equivalent to CO₂ emission of 0.3 mol m⁻² with the global warming
532 potential of CH₄ as 9.1 for a molar basis (Forster et al., 2007). Thus, the
533 contribution of CH₄ emission to the greenhouse gas budget was 5% in this
534 forest in 2011. The CH₄ emission in this black spruce forest could not be
535 overlooked when considering the greenhouse gas budget due to its stronger
536 global warming potential.

537 **5. Conclusions**

538 Here we examined seasonal and interannual variations in ecosystem-scale
539 CH₄ exchange at a poorly-drained black spruce forest over permafrost, rep-
540 resenting one of the typical ecosystems of interior Alaska and boreal Canada.
541 The magnitude of CH₄ emission and its dependence on environmental vari-
542 ables varied, depending on the position within a small catchment. CH₄ emis-
543 sion was greater at the wetter bottom of the small catchment than at a drier
544 upper position. At the drier upper position, soil water content affected the

545 seasonal variation in CH₄ emission. At the wetter bottom, in addition to soil
546 temperature and soil water content, seasonal thaw depth of frozen soil was
547 also an important variable explaining the seasonal variation in CH₄ exchange.
548 These different responses to changes in environmental conditions within the
549 ecosystem should be considered when estimating the spatial variation in CH₄
550 exchange in ecosystems over permafrost. The interannual variation of total
551 snow-free season CH₄ emission in the wetter bottom of the catchment (30.3–
552 45.0 mmol m⁻² season⁻¹) was influenced by the amount of rainfall and thaw
553 depth. On the other hand, in the drier upper position the amount of rainfall
554 did not strongly affect the total season CH₄ emission, because the deeper
555 soil layer where CH₄ was presumably produced was kept wet from soil thaw-
556 ing even in a year with low rainfall. Total season CH₄ emission was nearly
557 one order smaller than those reported in other northern wetland ecosystems,
558 likely due to the relatively low ground water level and soil temperature.
559 However, degradation of the ice-rich permafrost, expected in future warmer
560 environment, may enhance CH₄ emission in boreal forests with permafrost.
561 CH₄ exchange components in ecosystem models have not sufficiently been
562 validated for various wetland types, and it has not been assured whether
563 the models can reproduce both spatial and temporal variations in CH₄ ex-
564 change. Further efforts are needed to improve the ecosystem models using
565 eddy covariance observations for accurate estimates of regional and global
566 CH₄ exchange.

567 **Acknowledgments**

568 This research was funded through grants to the University of Alaska Fair-
569 banks, International Arctic Research Center from NSF under the Carbon Cy-
570 cle Program of IARC/NSF, from the Japan Aerospace Exploration Agency
571 (JAXA) under the “Arctic Research Plan Utilizing the IARC-JAXA Infor-
572 mation System (IJIS) and Satellite Imagery”, and from the Japan Agency for
573 Marine-Earth Science and Technology (JAMSTEC) under the JAMSTEC-
574 IARC Collaboration Study, and by JSPS KAKENHI Grant Number 23310009
575 (FY2011–2013) and 23248023 (FY2011-2013). We would like to thank Mr.
576 N. Bauer for proofreading the manuscript, and the editor and two anonymous
577 reviewers for their constructive comments.

578 **Appendix A. Uncertainty Evaluation**

579 Random errors in CH_4 flux due to limited averaging time and the in-
580 strumental noise were evaluated using the method from Meyers et al. (1998)
581 and Finkelstein and Sims (2001). To account for the effect of high-frequency
582 loss, calculated random errors were multiplied with the same correction coef-
583 ficients as fluxes. Random error tended to increase with increasing absolute
584 magnitude of flux (Fig. 9), with typically 30 % of flux for positive values and
585 80 % of flux for negative values.

586 The total uncertainty of observed CH_4 exchange was estimated by com-
587 bining uncertainties due to random error and gap-filling, assuming uncertain-
588 ties are independent and random (Taylor, 1997). Half-hourly uncertainties
589 due to random error were added in quadrature, to obtain uncertainties of

590 total CH₄ exchange due to random error—i.e.,

$$U_{\text{total,RE}} = \sqrt{\sum_{i=1}^N U_{\text{RE},i}^2} \quad (1)$$

591 where $U_{\text{total,RE}}$ is the uncertainty of total exchange due to random error,
592 $U_{\text{RE},i}$ is half-hourly uncertainty due to random error, and N is the number
593 of data to be summed. For gap-filled data, the obtained relationships, as
594 shown in Fig. 9, were used to estimate random errors of fluxes. For gap-filled
595 fluxes with an absolute magnitude smaller than $1.25 \text{ nmol m}^{-2} \text{ s}^{-1}$, median
596 value was obtained for this range, and this constant value was assigned as the
597 random error of gap-filled fluxes. Similarly, uncertainties in daily total CH₄
598 exchange due to gap-filling described in the previous section were also added
599 in quadrature, to obtain the uncertainty in total CH₄ exchange. Finally, the
600 uncertainty in total CH₄ exchange due to both random error and gap-filling
601 was combined by adding them in quadrature.

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Table 1:

System modification	Year		
	2011	2012	2013
CH ₄ analyzer	Fast Methane Analyzer [#]	→ Aug/23	Greenhouse Gas Analyzer ^{##}
Pump	Piston vacuum pump [§]	→ Jun/18	Dry scroll vacuum pump ^{§§}
Nafion dryer	Reflex method [*]	→	Circulation of untreated air ^{**}
Length of sampling tube	8 m	→ Jun/18	22 m ^{&}
Flow rate	Approx. 10 L min ⁻¹	→ Jun/18	Approx. 13 L min ⁻¹ → Approx. 31 L min ⁻¹
Output rate	1 Hz ^{&&}	→	10 Hz

[#]RMT-200 Fast Methane Analyzer (Los Gatos Research, Inc., USA).

^{##}Greenhouse Gas Analyzer with the enhanced cell temperature control (Los Gatos Research Inc., USA) customized to conduct CH₄ and water vapor concentration measurements only to improve the signal-to-noise ratio (D. Baer, personal communication, 2011).

[§]Model 4VCF-10-M450X, Gast, USA.

^{§§}Model ISP-500C, Anest Iwata, USA.

^{*}The exhaust air from the CH₄ analyzer was used as the dry purge air.

^{**}To reduce the resistance of flow line, the external pump was placed directly after the CH₄ analyzer, and untreated air was circulated for the dry purge air using another small pump.

[&]The gas analyzer was moved away from the tower to prevent damage of vegetation.

^{&&}To improve the signal-to-noise ratio under the limited flow rate condition.

Table 2:

Environmental conditions	Year		
	2011	2012	2013
Mean air temperature (°C)	12.1	11.5	11.2
Range of air temperature (°C)	0.7 to 22.0	−4.0 to 20.8	−3.8 to 25.5
Mean soil temperature (°C)	3.2	3.5	3.3
Range of soil temperature (°C)	−0.7 to 7.6	−0.5 to 8.0	−7.9 to 9.4
Total solar radiation (GJ m ^{−2})	2.38	2.33	2.54
Total rainfall (mm)	195.1	185.3	146.1
Maximum thaw depth (m)	0.40	0.36	0.33

Table 3:

Wind direction (degree)	Year		
	2011	2012	2013
0–60	2.0 (1.1–2.7)	1.5 (1.1–1.8)	1.3 (0.8–1.7)
60–120	1.5 (1.1–1.9)	1.6 (1.4–1.8)	0.9 (0.7–1.1)
120–180	0.4 (0.0–1.0)	0.6 (0.5–0.8)	0.5 (0.0–0.8)
180–240	0.8 (0.3–1.5)	0.9 (0.7–1.1)	0.7 (0.5–1.0)
240–300	3.0 (2.5–3.3)	2.6 (2.4–2.9)	2.6 (2.4–3.0)
300–360	2.0 (1.3–2.7)	2.7 (2.4–3.2)	1.8 (1.5–2.1)

Table 4:

Area (wind direction: degree)	Variable						
	$T_{s,0.1\text{ m}}$	$T_{s,0.2\text{ m}}$	Thaw depth	$SWC_{0-0.1\text{ m}}$	$SWC_{0.1-0.2\text{ m}}$	$SWC_{0.2-0.3\text{ m}}$	
Drier (120-240)	-0.06	0.16	0.53***	0.20	0.57***	0.61***	
Moderately wet (0-120)	0.17	0.39**	0.70***	0.46**	0.71***	0.61***	
Wetter (240-360)	0.18	0.36**	0.66***	0.43**	0.65***	0.62***	

Table 5:

Area (wind direction: degree)	Year		
	2011	2012	2013
Drier (120–240)	10.6 ± 1.6	13.9 ± 0.8	11.3 ± 1.2
Moderately wet (0–120)	23.2 ± 1.7	21.9 ± 1.1	13.7 ± 1.4
Wetter (240–360)	45.0 ± 1.6	34.6 ± 1.0	30.3 ± 1.1

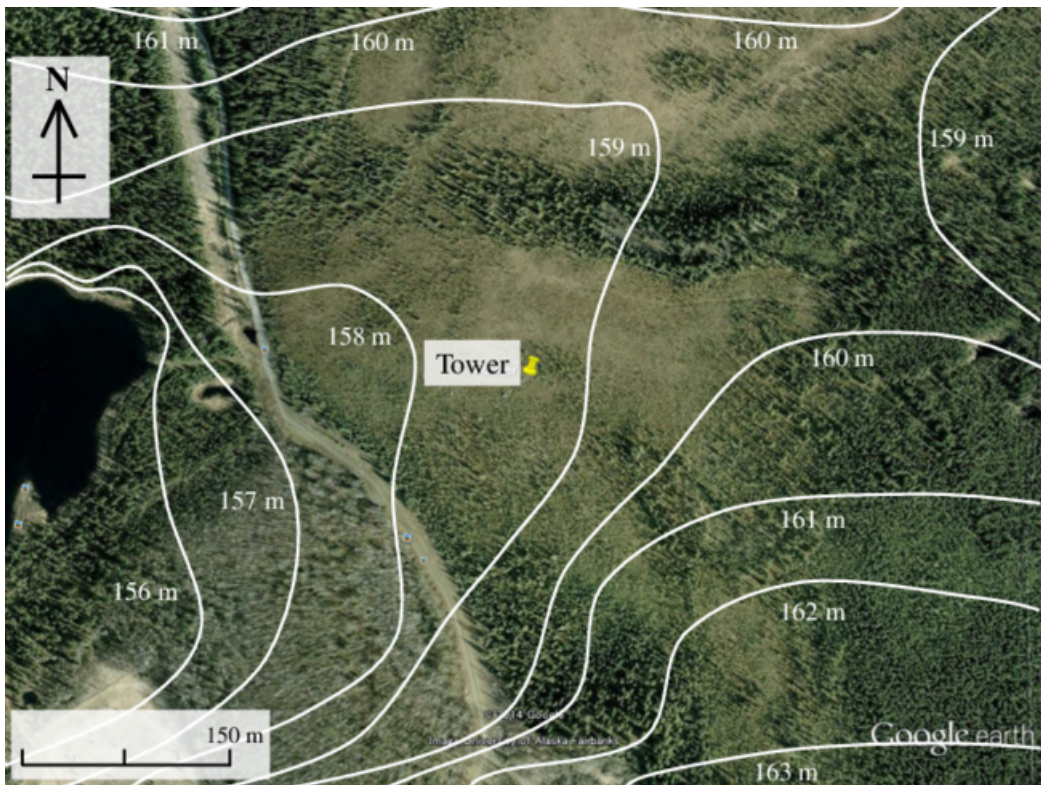


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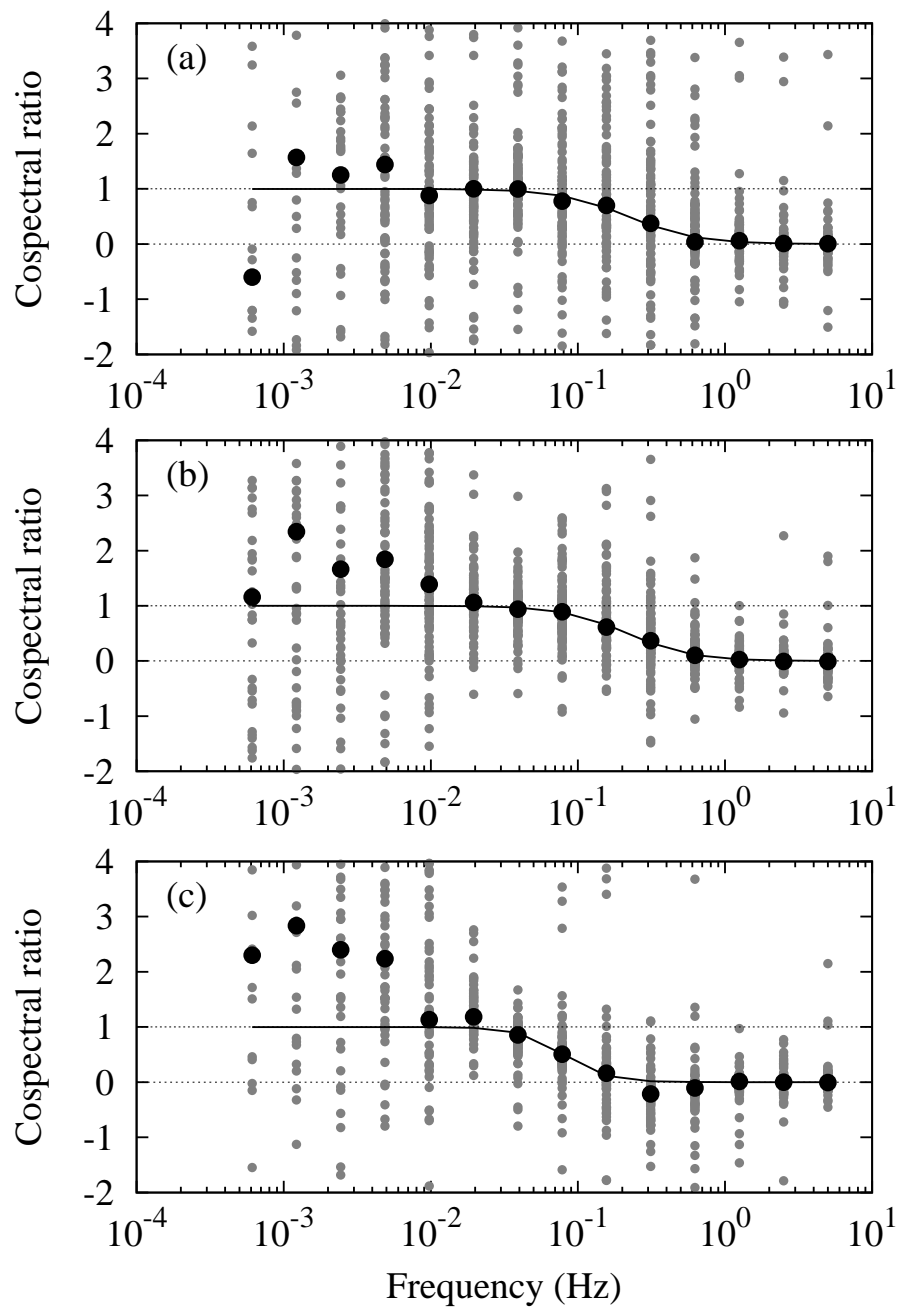


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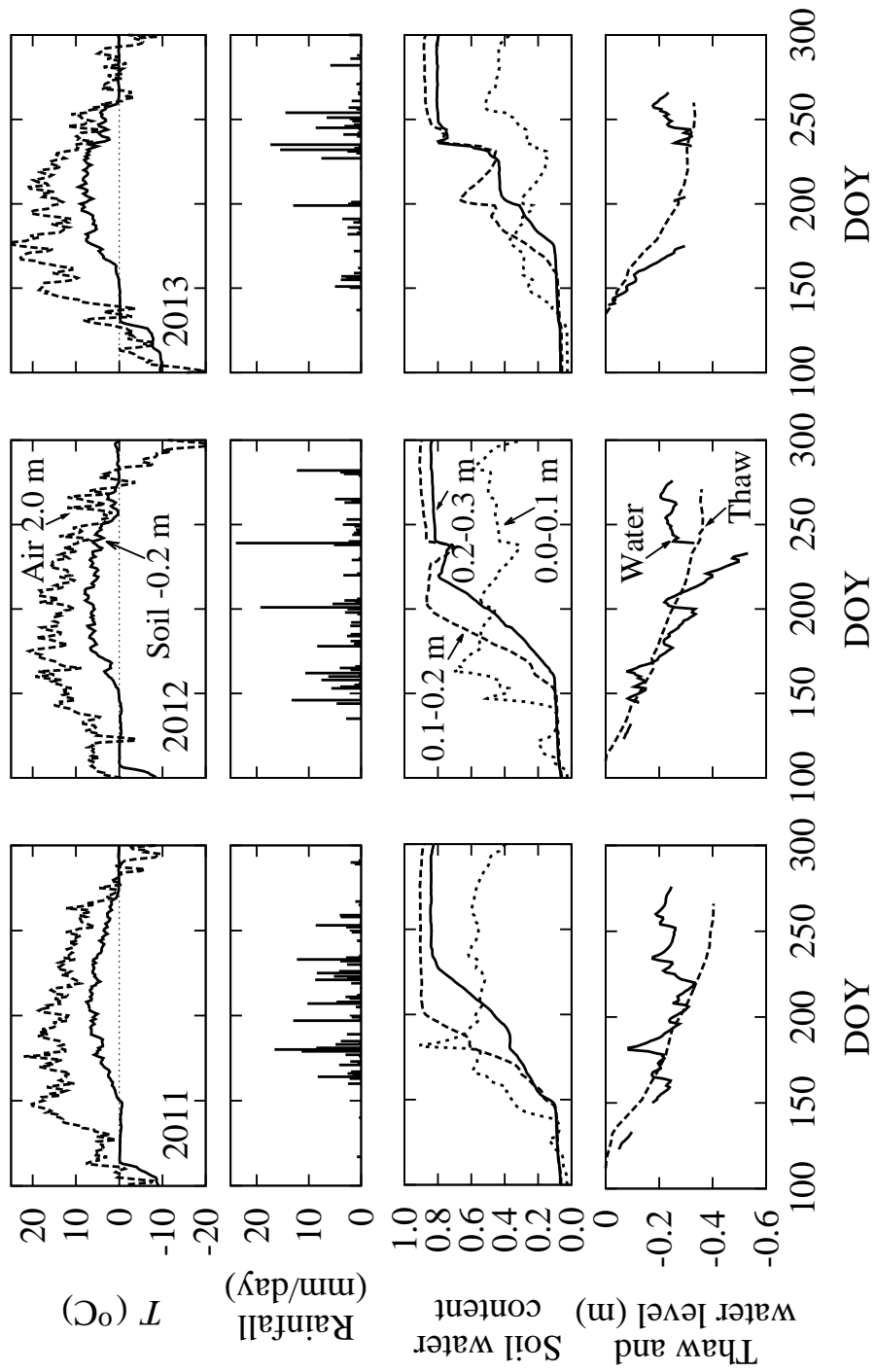


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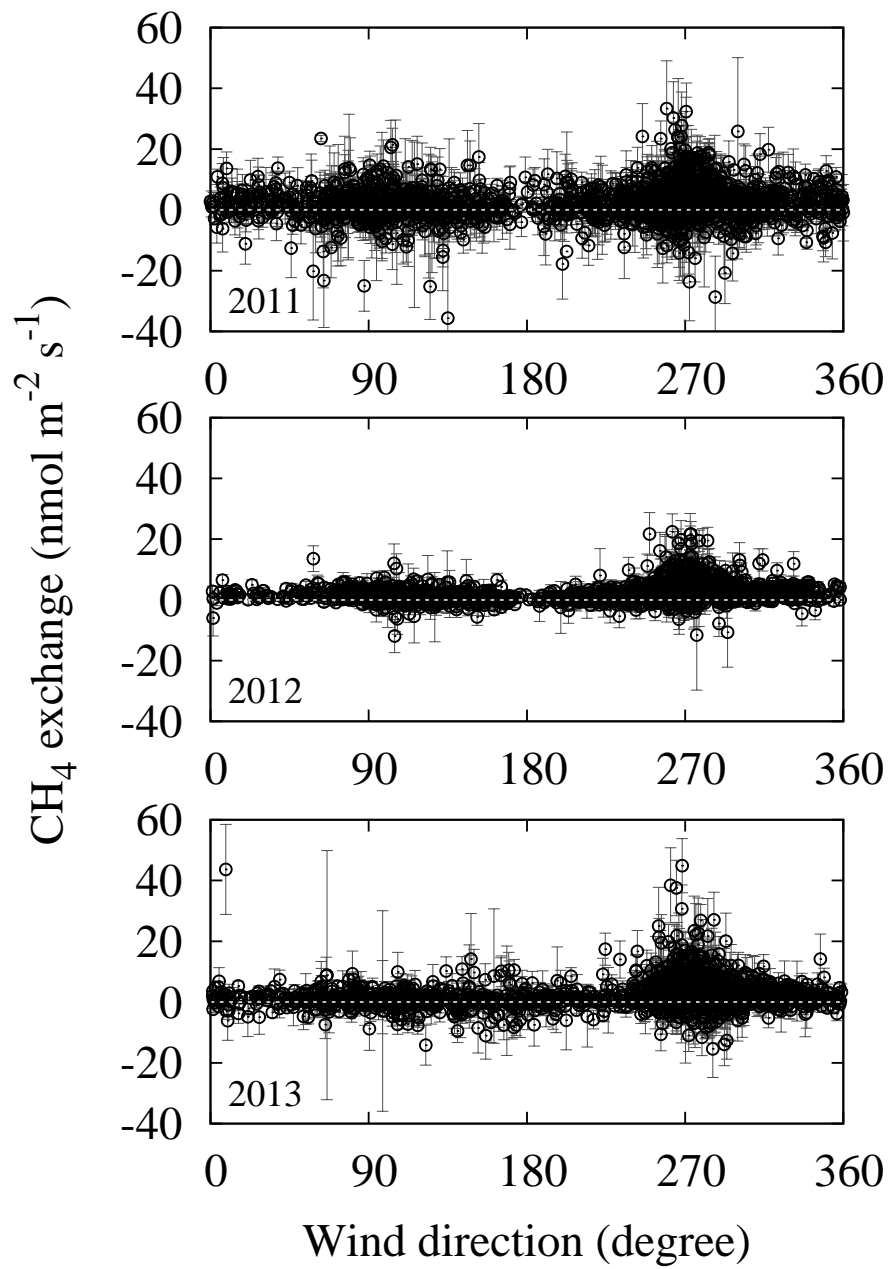


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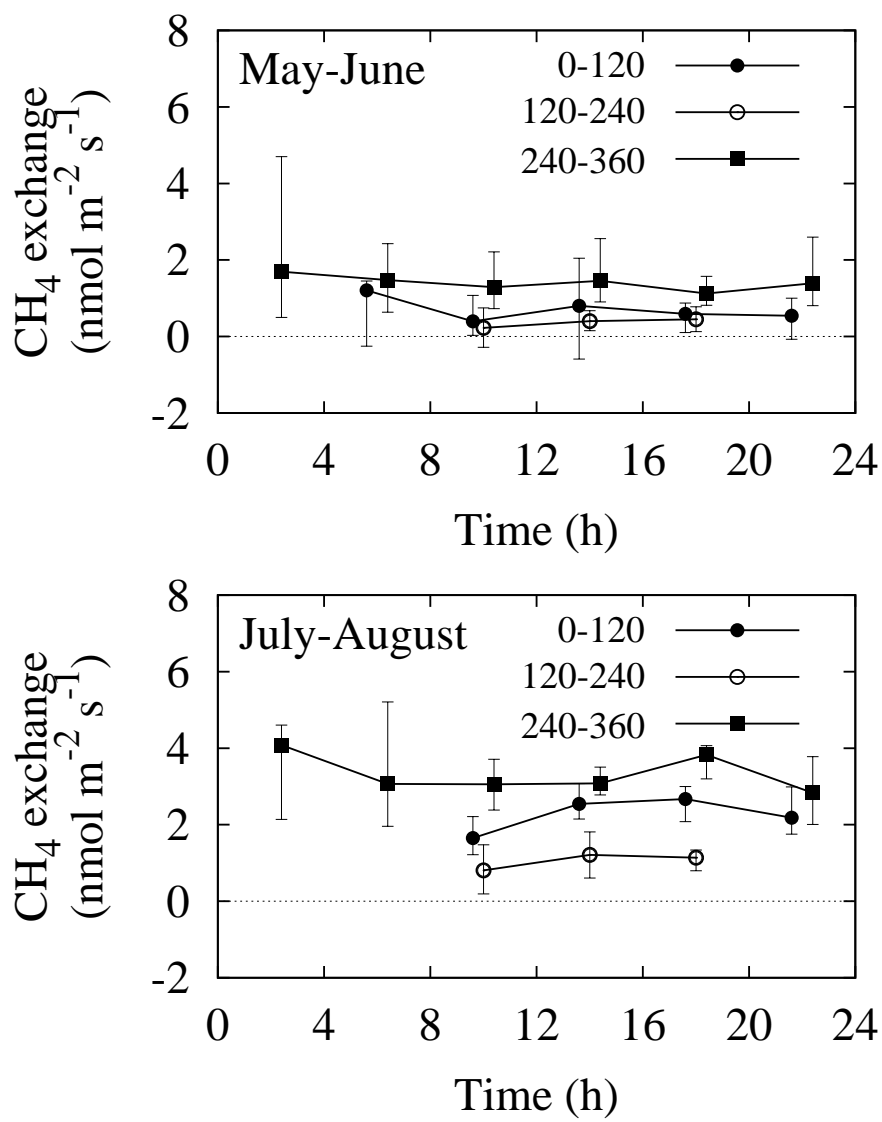


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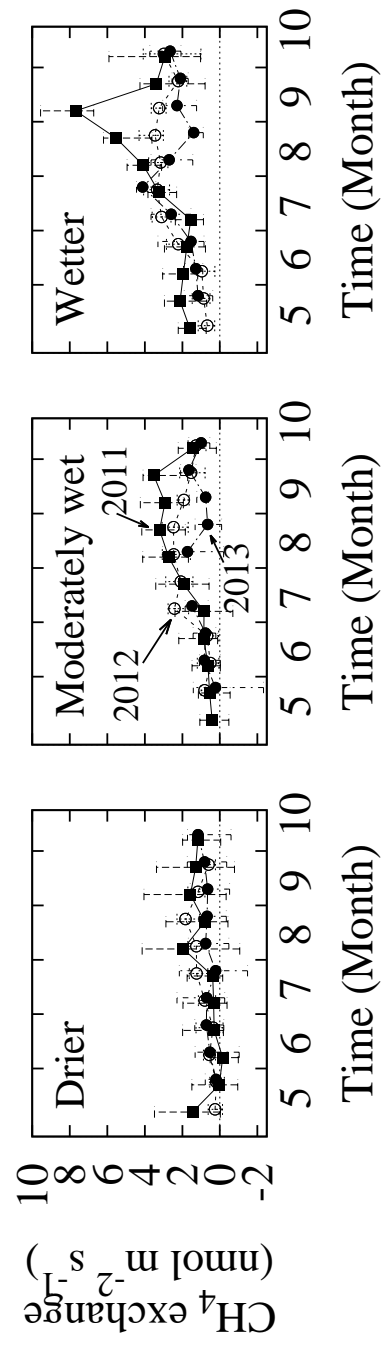


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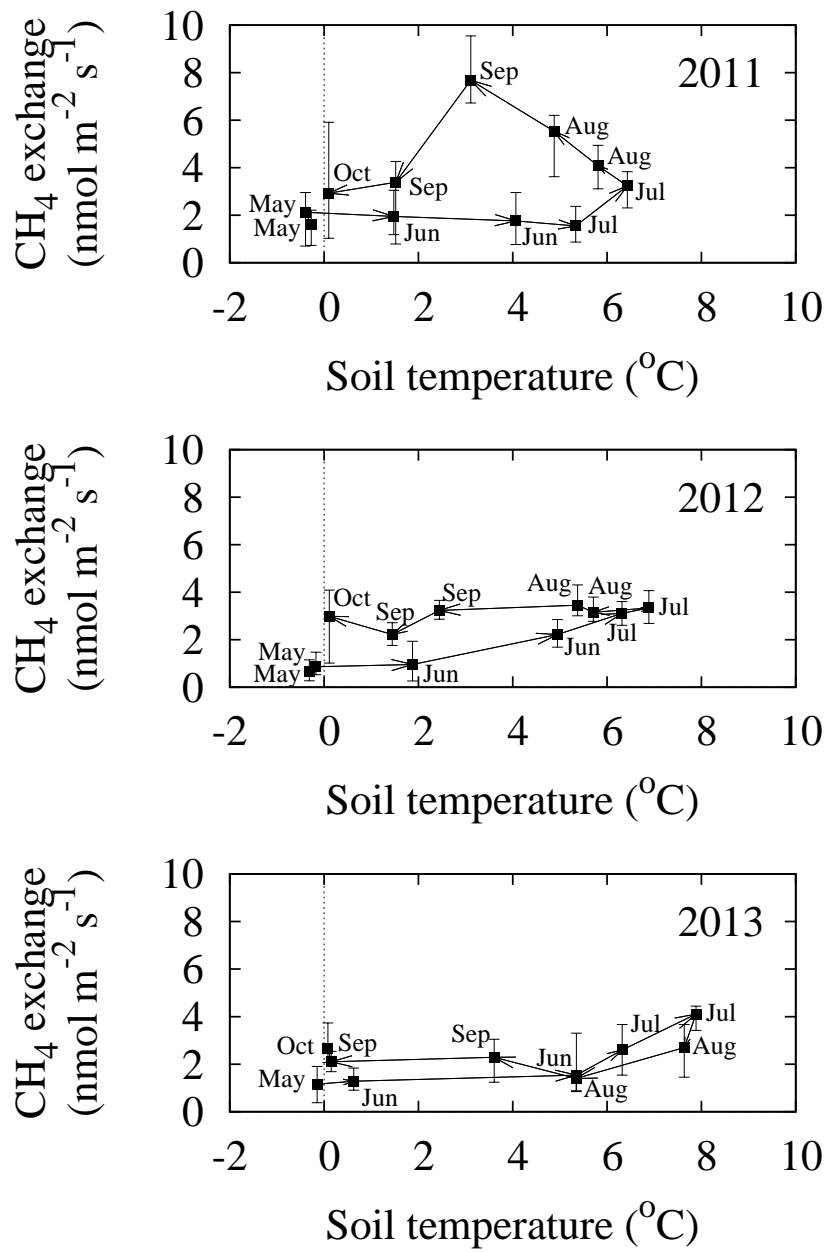
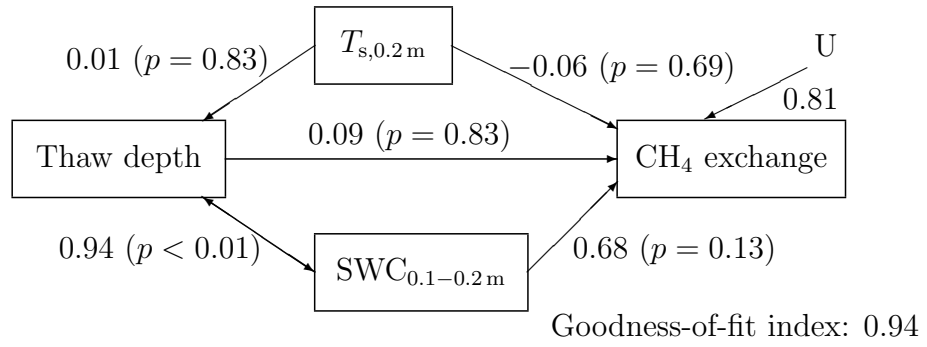
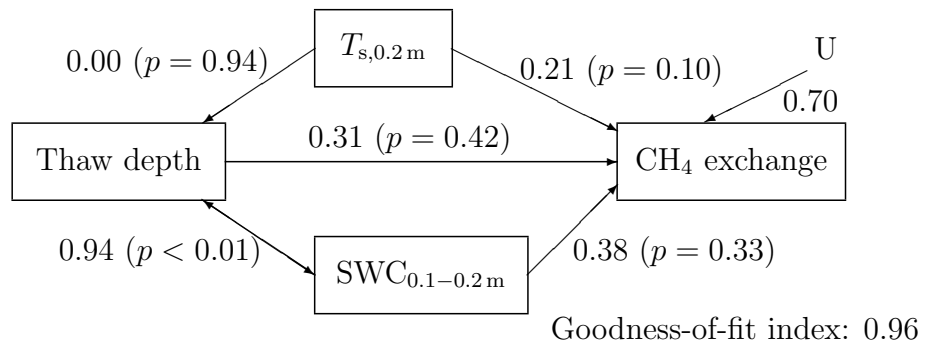


Figure 7:

Drier (120–240 degree)



Moderately wet (0–120 degree)



Wetter (240–360 degree)

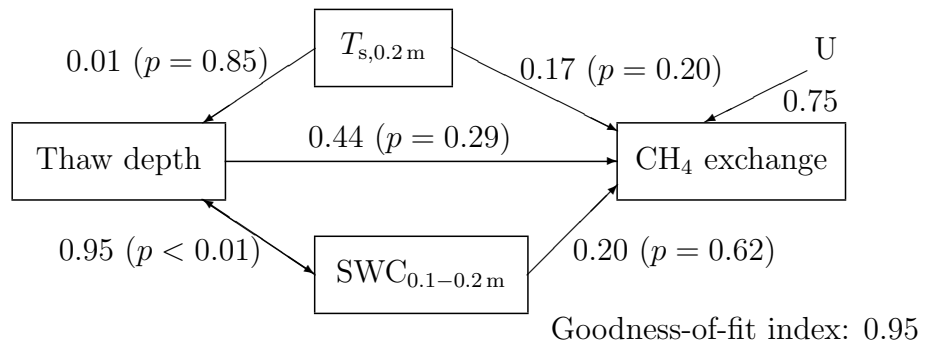


Figure 8:

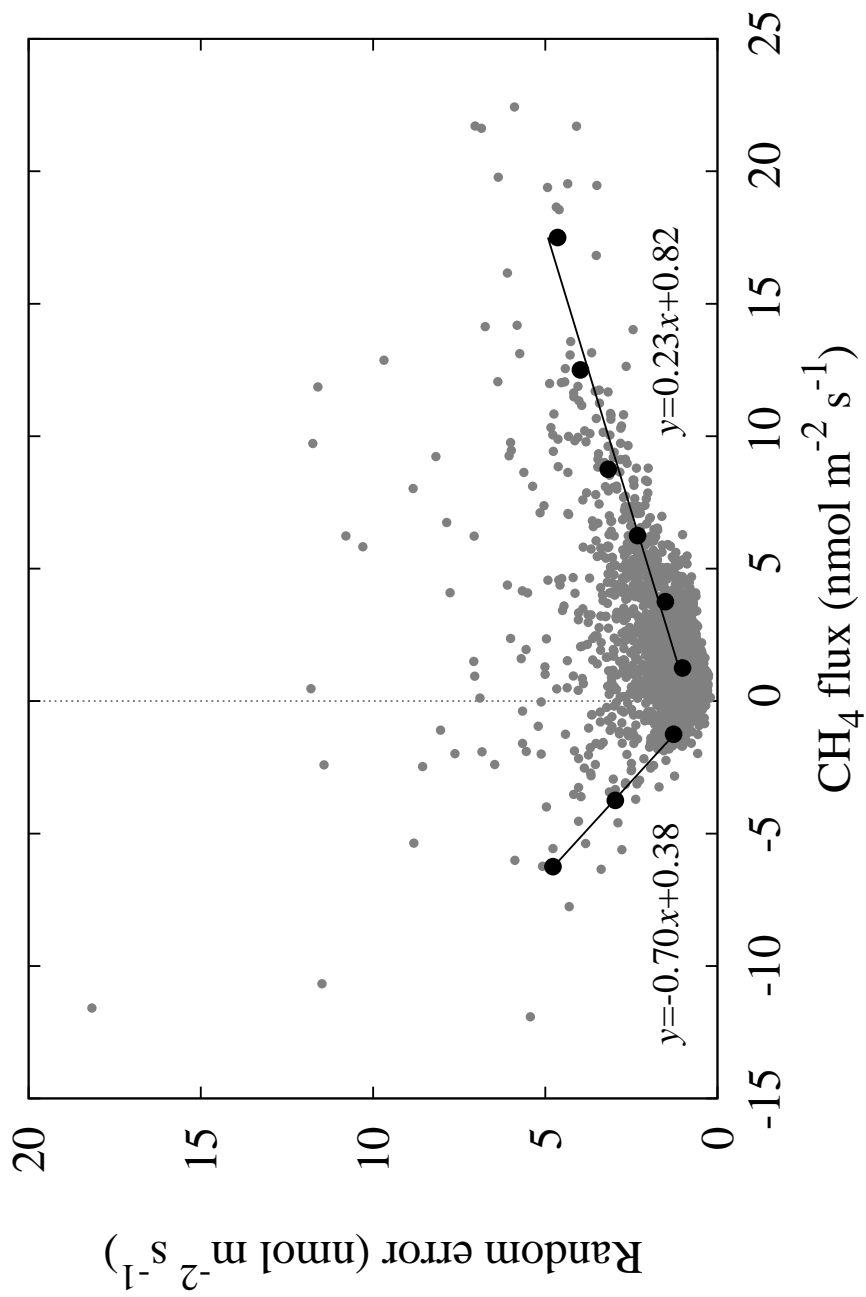


Figure 9: