Citation

URL
https://oro.open.ac.uk/38753/

License
None Specified

Policy
This document has been downloaded from Open Research Online, The Open University's repository of research publications. This version is being made available in accordance with Open Research Online policies available from Open Research Online (ORO) Policies

Versions
If this document is identified as the Author Accepted Manuscript it is the version after peer review but before type setting, copy editing or publisher branding
Methane fluxes during the initiation of a large-scale water table manipulation experiment in the Alaskan Arctic tundra


Received 5 February 2009; accepted 24 February 2009; published 14 May 2009.

1 Much of the 191.8 Pg C in the upper 1 m of Arctic soil of Arctic soil organic mater is, or is at risk of, being released to the atmosphere as CO\textsubscript{2} and/or CH\textsubscript{4}. Global warming will further alter the rate of emission of these gases to the atmosphere. Here we quantify the effect of major environmental variables affected by global climate change on CH\textsubscript{4} fluxes in the Alaskan Arctic. Soil temperature best predicts CH\textsubscript{4} fluxes and explained 89\% of the variability in CH\textsubscript{4} emissions. Water table depth has a nonlinear impact on CH\textsubscript{4} efflux. Increasing water table height above the surface retards CH\textsubscript{4} efflux. Decreasing water table depth below the surface has a minor effect on CH\textsubscript{4} release once an aerobic layer is formed at the surface. In contrast with several other studies, we found that CH\textsubscript{4} emissions are not driven by net ecosystem exchange (NEE) and are not limited by labile carbon supply.


1. Introduction

Peatland regions of the world are sites of past accumulation of organic materials due to generally anaerobic conditions [Harriss et al., 1985]. These regions could be significant current and future global sources of CH\textsubscript{4} and CO\textsubscript{2} release [Ehhalt, 1974; Harriss et al., 1985; Oechel et al., 1993; Oechel et al., 1994; Zimov et al., 1997; Harazono et al., 2003]. The amount of carbon stored in the upper 1 m layer of Arctic tundra soil is approximately 191.8 Pg [Post et al., 1982; Schlesinger, 1991; Hobbie et al., 2000], 14\% of the global soil organic carbon [Post et al., 1982; Billings, 1987]. A more recent estimate reported the carbon stock in the first 1 m of cryosols in the northern circumpolar regions to about 268 Pg of organic carbon [Tarnocai et al., 2003]. However, large amounts of soil organic carbon lie below 100 cm in both mineral and organic soils [Sombroek et al., 1993; Tarnocai, 1994] and reserves of carbon in the upper 2 m of the world’s peat soils is 679 Pg of C [Batjes, 1996]. The large carbon storage is attributed to low decomposition rates in saturated soils and to low temperatures [Clymo, 1984]. As the global climate warms, carbon stored in these regions could be released to the atmosphere [Grulke et al., 1990; Oechel et al., 1993; Oechel et al., 1995], providing a strong positive feedback to global warming. The current release of CH\textsubscript{4} from northern peatlands is estimated to range from 14.1 Tg C-CH\textsubscript{4} [Fung et al., 1991; Bartlett and Harriss, 1993] to 46 Tg C-CH\textsubscript{4} each year [Gorham, 1991].

Recent evidence indicates a heterogeneous response of soil moisture in the Arctic to global warming. Soil drying in continuous permafrost has been reported and appears to be due to an increasing gap between potential summer evapotranspiration and summer precipitation [Oechel et al., 1993; Oechel et al., 2000; Barber et al., 2000, Dickson, 2000; Klein et al., 2005]. Drying in discontinuous permafrost is often due to increased drainage following permafrost degradation [Yoshikawa and Hinzman, 2003; Smith et al., 2005; Riordan et al., 2006]. On the other hand, some Arctic areas on continuous permafrost are showing increased lake numbers and/or extent [Smith et al., 2005]. In addition to lake expansion, slumping and thermokarst erosion can cause new wet areas. These latter situations could result in extensive new areas of anaerobic soils. Therefore, even under scenarios of warming and drying of the Arctic, many regions underlain by continuous permafrost are likely to show increased water availability and anoxic condition in the soil in coming decades.

The drying of currently wet and anaerobic areas of the Arctic will likely affect the net rates of CH\textsubscript{4} emission from soils [Moore and Knowles, 1989; Freeman et al., 1993; Funk et al., 1994; Bubier, 1995; Nykänen et al., 1998]. In particular, the increase in surface soil aeration resulting from a drop in the water table can increase methanotrophy near the soil surface, and lead to the almost complete oxidation...
of CH₄ produced in deeper, anaerobic layers [Conrad and Rothfuss, 1991; Updegraff et al., 1995; Cao et al., 1996; Bridgham et al., 1998]. Field and laboratory manipulations generally show a decrease in CH₄ emissions with the increase in drainage (or a lowering of the water table) [Moore and Roulet, 1993; Sihvola et al., 1996; Oechel et al., 1998a; Updegraff et al., 2001; Whittington and Price, 2006]. However, some other experiments showed contrasting results where sometimes CH₄ emissions were not affected by lowered water table [Updegraff et al., 2001; Chimner and Cooper, 2003; Strack and Waddington, 2007]. These contradictory results make generalization and prediction of CH₄ release with changes in water table difficult.

In addition to the effects of water table, and soil aeration, CH₄ fluxes may also be influenced by availability of labile C. Ecosystem productivity has been positively correlated with CH₄ release in a number of studies, including some in Arctic tundra [Whiting and Chanton, 1993; Reeburgh, 1996; Nylänen et al., 2003]. This link could be explained by an increase in energy available to methanogens associated with increased plant productivity. Prediction of future CH₄ release requires a solid understanding of the controls on CH₄ production, including the role that substrate availability plays in limiting methanogenesis.

Because trace gas fluxes are so tightly linked to soil moisture and water table in the Arctic, we initiated a large-scale manipulation in the Alaskan Arctic at the Barrow Environmental Observatory (BEO) in 2005 as part of the NSF Biocomplexity Program. The target of this manipulation includes increasing and decreasing the water table over large areas of tundra and observing the effects over the diverse microtopography of the region. Here, we present results from the initial year of the manipulation. The differences in water table reported are caused by the installation and initiation of the dikes, as well as by seasonal progression in water table, but do not reflect the full long-term effect of the planned manipulation. However, these results do show the impact of the observed variation and progression in water table on CH₄ fluxes. The Biocomplexity experiment described here is the first large-scale (1.2 km x 0.3 km) water table manipulation in the Arctic. The experiment was designed to investigate the link between CH₄ and CO₂ fluxes as affected by changes in soil water, over complex terrain.

This paper focuses on the patterns and controls on CH₄ flux and the relationship between NEE and CH₄ fluxes during summer (June and July) 2007, the first year of manipulation. We also present the comparison of the two locations within the experimental area with different water table depth and soil moisture. We hypothesize that (1) water table and soil moisture are major predictors for CH₄ fluxes; an alternative hypothesis is that (2) NEE is a major predictor of CH₄ fluxes.

2. Site Description

The investigation site was located in the wet sedge tundra in the northern part of the Arctic Coastal Plain which generally occurs across the northernmost region in Alaska and is distinguished by low elevation, gentle slopes, and proximity to the Arctic Ocean [Brown, 1967]. The area is characterized by continuous permafrost and by an average seasonal thaw (active layer) depth of about 37 cm [Hinkel et al., 2001]. More than half of the surface is covered by polygonal ground [Brown, 1967]. Low center and high center polygons are common features of this area. The soils of the site are characterized by the presence of the main horizons: an organic-rich surface layer, a horizon of silty clay to silt loam textured mineral material, and an underlying perennially frozen organic-rich mineral layer [Brown et al., 1980]. The soils are classified as Gelisols in the suborders Histels (organic soils with permafrost within 100 cm of the surface) [Bockheim et al., 1999].

The study site is located about 10 km east of the town of Barrow, Alaska. This 3,021 ha BEO reserve was set aside by the Ukpeagvik Iñupiat Corporation (UIC, the Barrow village native corporation), in perpetuity, for scientific research. The vegetation shows very high heterogeneity reflecting variation in microtopographic and other environmental factors and the presence of thaw lakes, frost boils and other patterned ground features [Walker et al., 1998]. In particular lakes and revegetated drained lake basins represent about 20% and 50% respectively of the northern part of the Arctic Coastal Plain [Hussey and Michelson, 1966; Hinkel et al., 2003; Hinkel et al., 2005]. After the (usually) natural drainage of the lakes, as part of the thaw lake cycle [Billings and Peterson, 1980; Hinkel et al., 2003], vegetation establishes (or reestablishes) on the exposed lacustrine sediment. The vegetation mainly consists of wet sedge meadow tundra [Webber, 1978; Brown et al., 1980; Billings et al., 1982]. In our site, a vegetated drained lake basin that drained 50–300 years ago [Hinkel et al., 2003], mosses represent the major component of the living biomass (about 80% of the total living biomass in the second week of August 2006); the main vascular plant is represented by Carex aquatilis (average of 0.64 leaf area index), followed by Eriophorum vaginatum (0.04 leaf area index) and Dupontia fisheri (0.02 leaf area index) (P. C. Olivas and S. F. Oberbauer, unpublished data, 2006). The vegetated drained lake basin chosen for this study (1.6 km long and about 0.4 km wide at the widest point) was divided into three sections separated by two dikes (Figure 1). Dikes were built from water impermeable, interlocking rigid plastic barriers inserted in trenches dug in the frozen soil during the spring of 2007. Sand bags were placed at strategic locations to provide support for the barrier and impede any seepage. The dikes effectively slowed or prevented the water from moving from the north site to the south area toward the natural outlet for the basin and the associated drainage channel (Figure 1). The imposed treatments resulted in different water table heights between the various treatment areas, with the north area having the highest water table and the south the lowest.

3. Materials and Methods

3.1. CO₂, H₂O, and CH₄ Eddy Covariance Measurements

Three eddy covariance towers for CO₂, H₂O vapor, and energy flux were installed in the vegetated drained lake in the BEO (Figure 1) in July 2005, one in each of the three
manipulation sites (north 71°17’11.80”N 156°36’12.23”W, central 71°17’1.71”N 156°35’54.77”W and south 71°16’51.17”N 156°35’47.28”W). The eddy covariance towers operated continuously from July 2005 to the present.

CO₂ and H₂O fluxes were measured through the use of an open path infrared analyzer (Li-7500, Li-COR, Lincoln, Nebraska, United States) with a sampling rate of 10Hz [Oechel et al., 1998b; Vourlitis and Oechel, 1997, 1999; Harazono and Miyata, 1997]. The Li-7500 was positioned 10 cm from the center of the sonic anemometer. CO₂ and H₂O vapor were calibrated every 2 to 4 weeks using ultra-high-purity nitrogen as the H₂O and CO₂ zero and 729 ppm CO₂ in air standard gas (certified grade ± 1 ppm) (Matheson Gas Product, Montgomeryville, Pennsylvania, United States) for the CO₂ span. A dew point generator (Li-610, Li-COR, Lincoln, Nebraska, United States) was used to produce an air stream with a known water vapor dew point (typically 7°C lower than the ambient air temperature). A sonic anemometer (WindMasterPro, Gill Instruments, Limited, Lymington, Hampshire, United Kingdom) was used to measure the three wind velocity components and to determine the high-speed sonic temperature fluctuations.

In addition, two closed-path DLT-100 fast response CH₄ analyzers (Los Gatos Research, Mountain View, California, United States) were added to the system of the two sites studied in detail in 2007. The DLT-100 methane analyzers were not calibrated because the instruments do not require external calibration (see T. Owano and D. Baer, DLT-100 Fast Methane Analyzer Manual, Los Gatos Research, Mountain View, California, United States). The CH₄, CO₂ and H₂O raw signals were synchronized with the sonic anemometer data and sent through a wireless connection to a computer located in a protected control shelter 170 to 800 m from the eddy covariance tower sites.

3.2. Eddy Covariance Flux Computation and Gap Filling

Figure 1. Site of the Manipulation experiment in the Barrow Environmental Observatory (BEO), Barrow, Alaska. Indicated are the orientation of the main axis of the vegetated drained lake (north) and the dominant wind direction (east). Two dikes (highlighted in gray) separate the three sites and prevent water from draining through the drainage channel in the southern part of the basin. Boardwalks run along and across the basin and provide access to the site avoiding disturbance.
components to CO₂ and H₂O fluxes according to Webb et al. [1980]. We did not correct the CH₄ fluxes either for high-speed variation in sensible heat since the temperature fluctuations are dampened by passage of air through the tube [Leuning and Moncrieff, 1990; Leuning and King, 1992], nor for latent heat since it was a very small correction. Obvious data outliers (values more than 6 standard deviations from the 30 min mean for CO₂ and H₂O vapor and more than 10 deviations from the 30 min mean for the wind velocity components, u, v, and w) were removed. Data quality was assessed through the analysis of energy budget closure and by comparing cospectra of wiTi, wiCO₂, wiH₂O, wiCH₄ [Kaimal et al., 1972]. The data were filtered by wind direction, and only wind directions from 350 to 180 degrees were used, excluding all results when the winds were coming from the back of the tower and outside of the experimental area and the footprint of interest. A friction velocity (u*) threshold 0.25 m/s was used as a cut off, and data below this value were removed. The percentage of data loss for the closed path Los Gatos CH₄ analyzer was 15%, while for the open path LI-COR 7500 it was 32%. The reason for the loss was mainly due to rain, fog, or ice blocking the sonic transducers or the LI-COR 7500 mirrors, low turbulence condition or wind coming from the back of the tower. The gap filing procedures used were: linear interpolation for short gaps (from 1/2 h to 2 h) with approximately uniform environmental conditions [Falge et al., 2001]. The mean diurnal of four to seven adjacent days (averaged for the same time period) was used to fill larger gaps in the data, up to 5 days long [Falge et al., 2001]. For more details about the eddy covariance techniques see Baldocchi [2003].

4. Environmental Variables

[13] Micrometeorological variables were recorded continuously at each site. Soil moisture was measured at three depths (0–30 cm, 0–10 cm, 20–30 cm) in five different locations in proximity to the eddy covariance towers using Time Domain Reflectometry (TDR) (CS616 Campbell Scientific, Logan, Utah, United States) moisture probes. Soil temperature was recorded by type T thermocouples, (Omega Engineering, Stamford, Connecticut, United States) in nine different locations at 6 different depths (at surface, 1 cm, 5 cm, 10 cm, 20 cm and 30 cm depth). Surface temperature was recorded using an Apogee infrared sensor (Apogee Instruments, Incorporated, Logan, Utah, United States) pointing into the footprint of the tower at an angle of about 45° with the ground (with field of view 22°, a height above surface of 2.31 m, corresponding to a footprint of about 1.82 m²). Air temperature and relative humidity was recorded at three heights on the eddy tower structure (0.46, 1.6 and 2.95 m from the ground) using Vaisala HMP45C probes (Vaisala, Helsinki, Finland). Air pressure was measured with an electronic barometer (model PTB 101B, Vaisala, Helsinki, Finland). Incoming, reflected and surface photosynthetically active radiation (PAR) (400–700 nm) was recorded using quantum sensors (Li-190, Li-COR, Nebraska, United States). Net radiation was recorded using a net radiometer (REBS Q7, 0.25–60 μm), shortwave global solar radiation measurements in the spectral range from 310 to 2800 nm incoming and reflected from the ground was collected using two pyranometers (model CMP3, Kipp & Zonen, Delft, The Netherlands). The PAR sensors (except for a surface sensor placed in the vegetation layer), the net radiometer and the pyranometers were mounted on a tripod at about 1.5 m above the ground and at about 5 m to the side of the eddy covariance towers. To obtain a representative ground heat flux measurement, five heat flux plates (HFT3, REBS Incorporated, Seattle, Washington, United States) were installed near each tower positioned at 2 cm depth. Wind speed and wind direction were measured using a wind vane (RM Young Wind Sentry, R.M. Young Company, Traverse, Michigan, United States). Precipitation was recorded using tipping bucket rain gauges (TR-525M, Texas Electronics, Dallas, Texas, United States). All the instruments were connected to a data logger (model 23X, Campbell Scientific, Logan, Utah, United States) and each environmental variable was read once every 10 s and the 30 min averages were recorded.

[14] Water table and thaw depth measurements were made every 3–4 days about every 13 m along first 200m downwind from the towers during summer 2007. Thaw depth was measured using a graduated, pointed metal rod approximately 6 mm in diameter. Water table was measured in 2.5 cm diameter PVC pipe water wells installed at 12 locations randomly selected within the first 200 m of an upwind transect at each site. Three boardwalks were installed across the vegetated drained lake basin to provide access for sampling while avoiding disturbance of the vegetation (Figure 1).

5. Statistical Analysis

[15] General linear modeling was used to identify the most important predictors of CH₄ fluxes in the north site (Systat version 10, Systat Software Incorporated, 2002). Both a single variable model and a forward stepwise multiple regression approach were used to discriminate between and rank the most important environmental variables correlated with CH₄ fluxes. The models were applied to CH₄ fluxes averaged both in half-hour and daily blocks. The data from the south site were not included in these models because we lacked a continuous seasonal data set for this site.

[16] Thaw depth was divided into two groups after examination of the position of the water table (Figure 2a). The variable “Logit thaw depth” was a binary variable with 0 for thaw depth of <15 cm and 1 for thaw depths of >15 cm. We considered this value as a threshold because after 7 July, when thaw depths exceeded 15 cm, the CH₄ fluxes are consistently higher. The water table in the north site (Figure 2a) likely never dropped below 15 cm, and as a consequence the soil layer below this depth never became aerobic during the whole summer season and it is probably the soil layer where the majority of the methanogens occur. This interpretation is in agreement with past measurement of CH₄ concentration profiles at different saturated soil.
the differences in CH₄ fluxes between the two sites, averaged both in half-hour and daily blocks.

6. Results and Discussion

6.1. CH₄ Fluxes and Environmental Variables

[19] In the summer 2007 from visual inspection during daily visit to the sites, and from albedo measurements general snowmelt occurred on 10 June, however the north site showed a higher amount of snow and snowmelt at the tower occurred at 16 June at the north and 11 June at the south tower. The thaw depth was higher in the south site than in the north one until 2 July when the two sites showed the same thaw depth. After that date the north site had a larger thaw depth (Figure 2b). The water table was consistently higher in the north site than in the south during the entire summer (Figure 2a), and the average difference in water table between the two site was 6 cm. Summer 2007 was exceptionally dry and warm; the rainfall from mid-June to the end of July was 3.9 mm and the average air temperature for the same period was 4.7°C.

[20] Previous studies reported a wide range of estimates for CH₄ emission from the Arctic tundra. Svensson et al. [1975] and Svensson and Rosswall [1984] reported emissions from Swedish subarctic mire ranging from 0.34 to 950 mg CH₄ m⁻² d⁻¹. Sebacher et al. [1986] and Whalen and Reeburgh [1990] measured CH₄ emission along a latitudinal gradient from the Arctic to sub-Arctic between −0.3 to 265 mg CH₄ m⁻² d⁻¹. Vourlitis and Oechel [1997] estimated Arctic and subarctic CH₄ flux from 12 mg CH₄ m⁻² d⁻¹ from moist tundra to 100 mg CH₄ m⁻² d⁻¹ from wet sedge ecosystems. Wet sedge ecosystems appeared to be the dominant Arctic and subarctic sources for CH₄ and are estimated to emit between 30 to 128 mg CH₄ m⁻² d⁻¹ [Vourlitis et al., 1993; Vourlitis and Oechel, 1997]. CH₄ emission was shown to vary along a microtopographic gradient of soil moisture, with the highest fluxes occurring in wet, vegetated areas [Morrisey and Livingston, 1992]. In the Barrow region flooded and wet sites are also characterized by the highest emission, with an average around 48 mg CH₄ m⁻² d⁻¹, and dry sites are a small source (4.8 mg CH₄ m⁻² d⁻¹) or sometimes a small sink for CH₄ [Rhew et al., 2007]. Integration of CH₄ fluxes for different vegetation types in Greenland showed this area to be net source of CH₄ of 45.6 mg CH₄ m⁻² d⁻¹ [Christensen et al., 2000] during the summer growing season. In our study we measured CH₄ emission on average of 24.6 mg CH₄ m⁻² d⁻¹ during June and July, which is consistent with previous estimates and very similar to the 20 mg CH₄ m⁻² d⁻¹ determined by Fan et al. [1992] for the wet meadow tundra using micro-meteorological measurements.

6.2. Environmental Controls on CH₄ Fluxes

[21] In the early season 2007, the south site shows a larger CH₄ release than the north site (Figure 3a). CO₂ uptake during this period is pronounced for the south site, with substantial midday uptake. At the same time, the north site shows a small CH₄ release, with very little evidence of midday CO₂ uptake. The earlier snowmelt at the south site is probably responsible for the earlier activation of the
ecosystem for both \( \text{CH}_4 \) and \( \text{CO}_2 \) fluxes. Because of the earlier snowmelt, the south site shows higher thaw depth (Figure 2b), higher soil temperature, and higher soil moisture (Figures 3c and 3d) in late June when compared to the north site. In fact the ground is still largely frozen in the north site and the water availability to the plants is lower (from 37 to 61% volumetric water content, VWC, in the soil layer 0–10 cm depth) compared to the south site (from 82 to 94% VWC) during the period from 18 to 22 June 2007.

Later in the season (16 to 23 July) the south site shows \( \text{CH}_4 \) emission about 21% lower than the north site (Figure 4). At this time the north site has 20% higher soil moisture (at 0–10 cm depth), 1\(^\circ\)C lower average soil temperature (at –10 cm) than the south site (Figures 4c and 4d); the water table depth is 6 cm lower in the south site than in the north, and thaw depth is about 2 cm lower in the north than in the south (Figure 2). The differences between \( \text{CH}_4 \) fluxes in the north and south sites during both early and late season proved to be highly significant (t test, Bonferroni adjusted probability of 0.009).

The general linear model chosen to explain the differences in \( \text{CH}_4 \) fluxes between north and south sites, for the data averaged in daily blocks, included only the difference in soil moisture in the 0–10 cm layer between the two sites. This model explained 88% of the difference in

![Figure 3](image-url). Half-hour (a) \( \text{CH}_4 \) fluxes and (b) \( \text{CO}_2 \) fluxes in the north and south sites in early season (18 to 22 June 2007); (c) daily averaged volumetric water content in the first 10 cm depth and (d) soil temperature at 10 cm depth.

![Figure 4](image-url). Half-hour (a) \( \text{CH}_4 \) fluxes and (b) \( \text{CO}_2 \) fluxes in the north and south site in late season (16 to 23 July 2007); (c) daily averaged volumetric water content in the first 10 cm depth and (d) soil temperature at 10 cm depth.
CH$_4$ fluxes between the two sites. Lower water table and soil moisture probably allow more oxygen penetration to the upper soil layers in the south site. This, in turn, would result in greater CH$_4$ oxidation and direct inhibition of methanogenesis. However, single variable models that included only the difference in thaw depth or the difference in water table between north and south explained about 86% of the variability in the difference in CH$_4$ fluxes between the two sites, very similar to the one that included difference in soil moisture (0–10 cm) between the two sites. CH$_4$ production most likely occurs in the deepest, most anaerobic layers, i.e., in the thawed zone below the water table. For this reason we created a new variable, the difference between thaw depth and water table. Once we modeled the difference in CH$_4$ emission between sites as a function of this variable, the single variable model was able to explain 90% of the variability in CH$_4$ fluxes. The deeper thaw depth and the higher water table in the north site are linked to a larger amount of anaerobic peat available for methanogenesis. However, the correlation and colinearity among the different environmental variables presents a major challenge in the selection of the best statistical model in explaining the differences in CH$_4$ fluxes. Better knowledge of the processes occurring in the soil could improve the analysis and avoid underestimation of the role of variables with slightly lower explanatory power.

[25] CH$_4$ fluxes in the north site were measured continuously for a large part of the growing season. The daily averaged CH$_4$ fluxes in the north site (Figure 5) showed an increase in CH$_4$ release from 17 June to 7 July 2007, followed by a decrease until 10 July with little reduction from 10 to 30 July. Several univariable and multivariable models were used to explain seasonal changes in CH$_4$ fluxes in the north site as a function of environmental factors (see Table 1 for the univariable models and Table 2 for the multivariate analysis). On the daily timescale, the single variable model with the highest explanatory power of averaged CH$_4$ fluxes identified soil temperature at −10 cm and explained 89% of the variability in CH$_4$ fluxes (Table 1). In contrast, the best single variable model on the data averaged for half-hour periods also used soil temperature (−10 cm) and had an $R$-square of 54%. Increased soil temperature not only increases the CH$_4$ production but also CH$_4$ diffusion through the soil [Stensson and Rosswall, 1984]. Our results suggest that a large part of the half-hour variability in CH$_4$ fluxes is not well explained by the short-term (half-hour) variation in the key environmental variables measured. In fact, since methanogenesis occurs across a range of soil depths which have different soil temperatures and different diurnal temperature lags, determining the relationship between short-term soil temperature patterns and CH$_4$ efflux is not straight forward. The multiple variable model of the daily averaged CH$_4$ flux includes soil temperature at −10 cm, Logit thaw depth (or Logit water table) and soil moisture at 20–30 cm depth, and explained 94% of the variability in CH$_4$ fluxes (Table 2). The other environmental variables increased the explanatory power of the model less than 1%, their $p$ values were only slightly significant or not significant and, consequently they were not included in the model (Table 2). Also, the different variables were correlated with each other and their inclusion in the model could lead to colinearity problems.

Figure 5. (a) Daily averaged CH$_4$ fluxes (mg m$^{-2}$ h$^{-1}$), thaw depth, and water table and (b) soil temperature at 10 cm depth in the north site for the entire measurement period (17 June to 30 July 2007).
the results of other researchers [e.g., Jauhiainen et al., 2005; Pelletier et al., 2007].

[26] Once the water table dropped to a level lower than the surface, the layer of dry soil would be expected to lead to CH₄ consumption [Del Grosso et al., 2000] causing the observed decreases in CH₄ efflux. In fact, according to Kelker and Chanton [1997], CH₄ release from Carex may be affected by the water level if the water level covers the areas from which CH₄ exits from plant tissues (that is, at the location of the plant stem bases). If CH₄ could escape from any part of the leaf blades, changes in water levels would have a minor effect on CH₄ emission as the CH₄ would simply exit from farther up the leaf blades [Kelker and Chanton, 1997].

[27] A decrease in water table from a few centimeters below the soil surface to 11 cm below the surface seems to have little effect on CH₄ emissions (Figure 6a). This result is in opposition to the conclusions of other researchers where a drawdown of the water table was connected to decreased emission and/or to the sink activity for CH₄ in peatlands [Harriss et al., 1982; Whalen and Reeburgh, 1990; Conrad and Rothfuss, 1991; Updegraff et al., 1995; Cao et al., 1996; Bridgham et al., 1998]. Our interpretation is that the water table dropping to a few centimeters below the surface would lead to CH₄ consumption, causing decreases in CH₄ efflux. In fact, according to Kelker and Chanton [1997], CH₄ release from Carex may be affected by the water level if the water level covers the areas from which CH₄ exits from plant tissues (that is, at the location of the plant stem bases). If CH₄ could escape from any part of the leaf blades, changes in water levels would have a minor effect on CH₄ emission as the CH₄ would simply exit from farther up the leaf blades [Kelker and Chanton, 1997].

Table 1. Univariable Model Results for the Daily Average of CH₄ Fluxes as a Function of Several Environmental Variables at the North Site From 17 June to 30 July 2007

<table>
<thead>
<tr>
<th>Variable</th>
<th>F</th>
<th>p Value</th>
<th>R² (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Soil T – 10 cm</td>
<td>332.40</td>
<td>&lt;0.001</td>
<td>88.8</td>
</tr>
<tr>
<td>Soil moist (0–30 cm)</td>
<td>113.30</td>
<td>&lt;0.001</td>
<td>73.0</td>
</tr>
<tr>
<td>Logit thaw depth or Logit water table</td>
<td>103.20</td>
<td>&lt;0.001</td>
<td>71.1</td>
</tr>
<tr>
<td>Soil T – 20 cm</td>
<td>86.10</td>
<td>&lt;0.001</td>
<td>67.2</td>
</tr>
<tr>
<td>Water table</td>
<td>68.04</td>
<td>&lt;0.001</td>
<td>62.0</td>
</tr>
<tr>
<td>Thaw depth</td>
<td>68.50</td>
<td>&lt;0.001</td>
<td>62.0</td>
</tr>
<tr>
<td>Soil T – 1 cm</td>
<td>60.00</td>
<td>&lt;0.001</td>
<td>60.0</td>
</tr>
<tr>
<td>fCO₂</td>
<td>50.40</td>
<td>&lt;0.001</td>
<td>57.8</td>
</tr>
<tr>
<td>Soil T surface</td>
<td>42.00</td>
<td>&lt;0.001</td>
<td>50.0</td>
</tr>
<tr>
<td>Soil moist (0–10 cm)</td>
<td>37.00</td>
<td>&lt;0.001</td>
<td>47.0</td>
</tr>
<tr>
<td>Air T</td>
<td>36.40</td>
<td>&lt;0.001</td>
<td>46.4</td>
</tr>
<tr>
<td>Soil moist (0–10 cm)</td>
<td>22.33</td>
<td>&lt;0.001</td>
<td>34.7</td>
</tr>
<tr>
<td>Soil heat flux</td>
<td>21.30</td>
<td>&lt;0.001</td>
<td>34.3</td>
</tr>
<tr>
<td>Soil moist (20–30 cm)</td>
<td>21.50</td>
<td>&lt;0.001</td>
<td>33.8</td>
</tr>
<tr>
<td>RH</td>
<td>0.80</td>
<td>0.37</td>
<td>1.9</td>
</tr>
<tr>
<td>PAR</td>
<td>0.70</td>
<td>0.39</td>
<td>1.8</td>
</tr>
<tr>
<td>Wind speed</td>
<td>0.14</td>
<td>0.71</td>
<td>0.3</td>
</tr>
</tbody>
</table>

*Reported are F ratio, p value, and R² of each of the variables in the model. The best single variable model included soil temperature at 0 cm, and explained 89% of the variability in CH₄ fluxes. RH, relative humidity; PAR, photosynthetically active radiation. T, temperature; Units are mg m⁻² h⁻¹.

Table 2. Multivariable Model Results for the Daily Average of CH₄ Fluxes as a Function of Several Environmental Variables at the North Site From 17 June to 30 July 2007

<table>
<thead>
<tr>
<th>n</th>
<th>Multivariate</th>
<th>F</th>
<th>p Value</th>
<th>ΔR² (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Soil T – 10 cm</td>
<td>332.43</td>
<td>&lt;0.001</td>
<td>88.8</td>
</tr>
<tr>
<td>2</td>
<td>Logit thaw depth or Logit water table</td>
<td>103.24</td>
<td>&lt;0.001</td>
<td>3.1</td>
</tr>
<tr>
<td>3</td>
<td>Soil moist (20–30 cm)</td>
<td>21.48</td>
<td>0.001</td>
<td>2.1</td>
</tr>
<tr>
<td>4</td>
<td>Soil moist (0–10 cm)</td>
<td>5.02</td>
<td>0.031</td>
<td>0.7</td>
</tr>
<tr>
<td>4</td>
<td>Soil moist (0–30 cm)</td>
<td>4.54</td>
<td>0.04</td>
<td>0.7</td>
</tr>
<tr>
<td>4</td>
<td>Water table</td>
<td>5.16</td>
<td>0.029</td>
<td>0.7</td>
</tr>
<tr>
<td>4</td>
<td>Thaw depth</td>
<td>5.02</td>
<td>0.031</td>
<td>0.7</td>
</tr>
<tr>
<td>4</td>
<td>Soil T – 1 cm</td>
<td>1.96</td>
<td>0.17</td>
<td>0.3</td>
</tr>
<tr>
<td>4</td>
<td>Soil T surface</td>
<td>1.99</td>
<td>0.167</td>
<td>0.3</td>
</tr>
<tr>
<td>4</td>
<td>PAR</td>
<td>0.79</td>
<td>0.381</td>
<td>0.2</td>
</tr>
<tr>
<td>4</td>
<td>Soil T – 5 cm</td>
<td>0.93</td>
<td>0.341</td>
<td>0.2</td>
</tr>
<tr>
<td>4</td>
<td>Air T</td>
<td>1.10</td>
<td>0.301</td>
<td>0.2</td>
</tr>
<tr>
<td>4</td>
<td>fCO₂</td>
<td>0.67</td>
<td>0.418</td>
<td>0.1</td>
</tr>
<tr>
<td>4</td>
<td>RH</td>
<td>0.49</td>
<td>0.487</td>
<td>0.1</td>
</tr>
<tr>
<td>4</td>
<td>Soil T – 20 cm</td>
<td>0.61</td>
<td>0.439</td>
<td>0.1</td>
</tr>
<tr>
<td>4</td>
<td>Wind speed</td>
<td>0.00</td>
<td>0.971</td>
<td>0</td>
</tr>
<tr>
<td>4</td>
<td>Soil heat flux</td>
<td>1.67</td>
<td>0.204</td>
<td>0</td>
</tr>
</tbody>
</table>

*Reported are F ratio, p value, and ΔR² of each of the variables in the model. The multivariable model chosen included soil temperature at 0 cm, and explained 94% of the variability in CH₄ fluxes. Units are mg m⁻² h⁻¹.
the surface leads to the formation of an oxic layer, which results in the oxidation of CH\textsubscript{4} diffusing through the soil to the atmosphere. A further decrease in water table does not affect CH\textsubscript{4} emissions. At this time thaw depth increases simultaneously with the drop in water table, thereby maintaining a constant active soil volume below the water table (the area most suitable for methanogenesis) and methane bypasses the oxidative zone in soil via plant aerenchyma [Joabsson et al., 1999].

[26] The temporal pattern observed in the north site, that was described in section 6.1, probably also occurred in the south site, presumably due to similar processes. In fact in the south site there is a steady increase in methane release (Figure 3) with a concomitant decrease in water table (Figure 2a), until the water table reaches the surface on about 23 June. After that date the water table at the south site continues to drop and on 17 July it is ~9 cm. However, the limited seasonal trends observed in the south site would make any further discussion unwarranted.

[29] Ecosystem carbon supply was highly correlated with CH\textsubscript{4} release in several studies [Whiting and Chanton, 1993; Reeburgh, 1996; Nylänen et al., 2003]. In our experiment we could not find a significant correlation between NEE and CH\textsubscript{4} emission rates in a multiple regression analysis. The correlation between NEE and methane emission in a single variable model (Table 1) is probably due to dependence of these two variables on similar environmental parameters. It appears that CH\textsubscript{4} production in this system is not limited by available carbohydrates, or other labile carbon sources, and is therefore not likely to respond to increases in net CO\textsubscript{2} uptake from the atmosphere and is not expected to be stimulated by increased plant productivity or increased root exudation. It is well established that a significant fraction (~20%) of C fixed by photosynthesis is exuded by roots, in the form of labile compounds like sugars and organic acids [Hinsinger et al., 2005; Jones et al., 2004], supporting a link between NEE and labile carbon in soil. Dissolved organic carbon (DOC) and CO\textsubscript{2} fluxes were reported to be strongly correlated for the northern latitude soils [Neff and Hopper, 2002] that would suggest a correlation between CH\textsubscript{4} release and available carbon pool in the soil. Because of the very high concentration of available carbohydrate for microbes in soils in our site (D. A. Lipson, unpublished data, 2007), CH\textsubscript{4} emission is probably not limited by available carbon. The upper 15 cm or so of these peat soils is entirely organic matter. The high levels of complex organic matter in these soils appear to support high fluxes of labile C as well: the average DOC concentration at our manipulation site is 43 mg C/L, of which 43% was found to be glucose; furthermore, the activities of polysaccharide-degrading enzymes, such as cellulases, appear to be close to saturation (D. A. Lipson, unpublished data, 2007). The interpretation that CH\textsubscript{4} production is not carbon limited in this system is supported by the results of the experiments performed by von Fischer and Hedin [2002], von Fischer and Hedin [2007], Owens and von Fischer [2007] and von Fischer et al. [2007] where acetate addition in a location very close to the site of the experiment described in this paper did not lead to any significant changes in CH\textsubscript{4} emission rates, indicating that the system is not carbon limited.

[30] In conclusion, our study shows that the effect of a decrease in water table is not necessarily a decrease in CH\textsubscript{4} release. An increase in water table above the surface could increase the diffusive resistance to CH\textsubscript{4} release. Additionally, a drop in water table below the surface may not decrease CH\textsubscript{4} emissions, because of the simultaneous increase in thaw depth, and therefore soil volume available for methanogenesis. This study was performed at a landscape level and some of the interpretations presented here should be tested with more mechanistic microscale studies. Moreover, in the long-term, vegetation and species composition may change in response to altered water level, which will mostly likely affect CH\textsubscript{4} fluxes. Better understanding of the complex, holocoenotic [Billings, 1952], and nonlinear controls on CH\textsubscript{4} flux is necessary to confidently predict future CH\textsubscript{4} feedbacks from the Arctic.

[31] Acknowledgments. We thank Robert Clement, University of Edinburgh, for having written the code for the data reduction and for help in the data analysis; Joseph Verfaille and Hiroki Ikawa for field and technical assistance; Rob Rhew, Steven Hastings, and Rommel Zulueta for insight and advice; Douglas Deutschman for the helpful suggestions on the statistics analysis; and the Barrow Arctic Science Consortium (BASC) and Glenn Sheehan for logistic support. This manuscript was improved by the constructive comments of the two reviewers. This work was funded by the Biocomplexity Program of the National Science Foundation (award OPP 0421588).

References


