

Methane fluxes from three ecosystems in tropical peatland of Sarawak, Malaysia

Lulie Melling^{a,b,*}, Ryusuke Hatano^a, Kah Joo Goh^c

^aSoil Science Laboratory, Graduate School of Agriculture, Hokkaido University, Kita 9, Nishi 9, Kitaku, Sapporo 060-8589, Japan

^bDepartment of Agriculture Sarawak, Jalan Badruddin, 93400 Kuching, Sarawak, Malaysia

^cApplied Agricultural Research Sdn Bhd, Locked Bag 212, Sg. Buloh, P.O. Box 47000, Sg. Buloh, Selangor, Malaysia

Received 9 April 2004; received in revised form 26 December 2004; accepted 3 January 2005

Abstract

Methane fluxes were measured monthly over a year from tropical peatland of Sarawak, Malaysia using a closed-chamber technique. The CH₄ fluxes in forest ecosystem ranged from -4.53 to $8.40 \mu\text{g C m}^{-2} \text{h}^{-1}$, in the oil palm ecosystem from -32.78 to $4.17 \mu\text{g C m}^{-2} \text{h}^{-1}$ and in the sago ecosystem from -7.44 to $102.06 \mu\text{g C m}^{-2} \text{h}^{-1}$. A regression tree approach showed that CH₄ fluxes in each ecosystem were related to different underlying environmental factors. They were relative humidity for forest and water table for both sago and oil palm ecosystems. On an annual basis, both forest and sago were CH₄ source with an emission of $18.34 \text{ mg C m}^{-2} \text{yr}^{-1}$ for forest and $180 \text{ mg C m}^{-2} \text{yr}^{-1}$ for sago. Only oil palm ecosystem was a CH₄ sink with an uptake rate of $-15.14 \text{ mg C m}^{-2} \text{yr}^{-1}$. These results suggest that different dominant underlying environmental factors among the studied ecosystems affected the exchange of CH₄ between tropical peatland and the atmosphere.

© 2005 Elsevier Ltd. All rights reserved.

Keywords: CH₄; Oil palm; Peatswamp forest; Sago; Tree regression

1. Introduction

Tropical peatland covers an area ranging from 33 to 49 Mha, which is over 8% of the world's total (386–409 Mha) peatland. However, the relatively greater depth of tropical peat allows the storage of more than 70 Gt carbon, which constitutes almost 20% of the global peatland carbon (Maltby and Immirzi, 1993). This carbon store makes it an important source of atmospheric CH₄ (Bartlett and Harriss, 1993) since, the amount of CH₄ in the atmosphere is a balance between CH₄ emission, both natural and anthropogenic, and CH₄ removal (Watson et al., 1990). Tropical peatland is, therefore, commonly considered to play an essential role in the global cycling of carbon and in climate change (Sorenson, 1993).

Methane (CH₄) is a potent greenhouse gas with a relative global warming potential 23 times that of carbon dioxide over a time horizon of 100 years (IPCC, 2001a). The atmospheric CH₄ concentration has doubled, since, the start of the industrial revolution to 1750 ppb in 2000 (IPCC, 2001b).

CH₄ emissions from peat soils to the atmosphere are dependent on the rates of methane production and consumption and the ability of the soil and plants to transport the gas to the surface. Three major environmental factors that control emission rates from peatland were water table position (Crill et al., 1988; Moore and Dalva, 1993; Bartlett and Harriss, 1993), temperature (Crill et al., 1988; Nyakanen et al., 1995), and substrate properties such as pH and mineral nitrogen concentration (Mosier et al., 1991). It has also been suggested that the CH₄ consumption rate depend on management factors such as drainage, compaction and nitrogen (N) fertilization (Hansen et al., 1993; Ball et al., 1997).

Large areas of tropical peatland have been developed in recent years for agricultural plantations in Southeast Asia especially for oil palm and sago whereby drainage is a prerequisite. Drainage may decrease CH₄ flux

* Corresponding author. Address: Soil Science Laboratory, Graduate School of Agriculture, Hokkaido University, Kita 9, Nishi 9, Kitaku, Sapporo 060-8589, Japan. Tel.: +60 82 429621; fax: +60 82 429624.

E-mail address: lulie_melling@yahoo.com (L. Melling).

(Martikainen et al., 1992; Flessa et al., 1998) because the lower ground water table may result in a thicker aerobic top layer in the soil, thereby possibly decreasing CH₄ production and increasing CH₄ consumption. Our knowledge of CH₄ fluxes from different ecosystems on tropical peatland is still fragmentary (Bartlett and Harriss, 1993; Smith et al., 2000; Inubushi et al., 2003). Thus, information on the different ecosystems on tropical peatland and their major controlling factors affecting CH₄ fluxes are needed to determine their role as a source or sink of CH₄.

Therefore, this study was conducted with the following objectives: (1) to quantify the amount of CH₄ flux under the forest, sago and oil palm ecosystems and (2) to determine the underlying environmental factors that explain the differences in CH₄ flux in these three ecosystems, particularly the effects of drainage and compaction.

2. Materials and methods

2.1. Sites description

The three experimental sites were all located in the Mukah Division of Sarawak, Malaysia representing three ecosystems namely mixed peat swamp forest, sago (*Metroxylon sagu*) plantation and oil palm (*Elaeis guineensis*) plantation. The main characteristics of these ecosystems are shown in Table 1. The sites were part of a contiguous peat swamp whereby part of it was converted to oil palm plantation and sago plantation. Distance between forest site and oil palm plantation site was about 3.7 km and these sites were about 8.3–11.7 km away from sago plantation site. The climate at the study sites was equatorial characterized by high, even temperatures and heavy rainfall without a distinct dry season.

The peat soils were classified as Typic Tropofibrin in the USDA soil classification system (Soil Survey Staff, 1992) and Fibric Histosols in the FAO classification (FAO-UNESCO, 1974). They were acidic (pH < 4.0) with a very high loss of ignition of about 99%. A detailed description of the site properties is provided by Melling et al. (2005).

Measurements of climatic variables and CH₄ emissions were made at monthly intervals from August 2002 to July 2003.

Table 1
Main characteristics of the forest, sago and oil palm ecosystem

Ecosystem	Forest	Sago	Oil palm
	2°49'N, 111°51'E	2°45'N, 111°50'E	2°49'N, 111°56'E
Site code	F	S	P
Peat thickness (cm)	480	650	555
Bulk density (g/cm ³)	0.15 ± 0.004	0.16 ± 0.006	0.20 ± 0.007
C:N	27.24 ± 0.96	22.63 ± 0.78	23.43 ± 1.14
Base saturation (%)	22.92 ± 1.68	30.09 ± 2.59	32.32 ± 4.50

The figures show the mean ± SE.

2.2. Methane flux measurements

Methane (CH₄) fluxes were determined using a closed-chamber method (Crill, 1991). Three open-bottom stainless steel chambers, 20 cm in diameter and 25 cm in height was placed directly on the soil surface in each site for about 30 min before sampling (Norman et al., 1997) to establish an equilibrium state. The chamber was pushed into the ground by cutting the soil along the edge of the chamber to a depth of 3 cm from the soil surface to avoid gas leakage through the bottom of the chamber by lateral diffusion (Melling et al., 2005). The chambers were sealed with acrylic cover after attaining equilibrium. It had two ports, one for gas sampling and the other for attaching a sampling bag to equilibrate the chamber pressure with the atmospheric pressure. Headspace samples of 20 ml were extracted from the chamber at 0, 10, 20 and 40-min using a polypropylene syringe with a three-way stopcock. The extracted gas was transferred to a 10 ml vacuum vial bottle.

The sampling was conducted between 11.00 and 13.00 h each day.

2.3. Soil gas

Soil gas was sampled using a stainless steel pipe of 9 mm in diameter equipped with a silicon tube and a three-way stopcock inserted to a depth of 5, 10, 20, 40 and 80 cm. The pipes were pushed into the peat with a steel rod inserted in the pipe to prevent physical damage or obstruction during placement. Triplicate measurements were made for each depth. After the pipes were set up, 50 ml of air was siphoned out from each pipe. The pipes were then close by using the three-way stopcock. The pipes were kept overnight to allow the gas concentrations in the pipes to equilibrate with the soil air. Thus, the pipes were installed a day before sampling. From each pipe, 50 ml of the soil gas were extracted to a Tedlar bag using a polypropylene syringe whereby 20 ml was transferred into a 10 ml vacuum vial bottle using a double-ended hypodermic needle.

If only water samples were extracted from the pipes, dissolved CH₄ concentrations were measured using the methods of McAullife (1971). Using a 60 ml syringe, 30 ml water samples were extracted from the pipe, and then 30 ml of air was immediately drawn into the syringe. The syringe was shaken vigorously for 3 min and a 20 ml headspace sample was then transferred to a 10 ml vacuum vial bottle. Extraction of CH₄ by this method was 98% efficient. Soil pore space air and water samples were collected in each site at the time of flux measurements.

2.4. Methane analysis

CH₄ concentration was determined by a gas chromatograph equipped with a flame ionization detector (Hewlett Packard 6890N) maintained at 250 °C, using a 2 m long Poropak N column (80/100 mesh) maintained at 50 °C with

a N₂ carrier gas flowing at 40 ms⁻¹. CH₄ fluxes were calculated from the linear increase or decrease in gas concentration in the chamber with time, using a linear regression equation (Christensen et al., 1995). The annual CH₄ fluxes were calculated from the monthly averages as follows:

$$\text{Cumulative gas flux} = \sum_{i=1}^n R_i D_i$$

where R_i is the mean gas flux (g m⁻² d⁻¹) of the two sampling times, D_i is the number of days in the sampling interval, and n is the number of sampling times. In this study, negative fluxes indicated the uptake of atmospheric CH₄, while positive fluxes indicated the net production of CH₄ from the peat soil.

2.5. Environmental variables

Air temperature, soil temperatures at 5 and 10 cm below the soil surface, relative humidity (RH) and the gaseous flux were measured at the same time. Water table depth was also recorded when the CH₄ flux was measured. Monthly precipitation in each ecosystem was recorded. Three soil samples were collected monthly from each ecosystem at the same time as gas samplings were bulked for both physical and chemical analysis. Three other undisturbed core samples were also taken to determine their bulk density and moisture content. Details of the measurements have already been described by Melling et al. (2005).

2.6. Statistical analysis

Repeated measure analysis was used to compare methane fluxes with ecosystems as the subject and time as repeated measure using GenStat (GenStat, 2002). Tree regression was used instead of the classical multiple linear regression to avoid multicollinearity problem between the independent variables.

Tree regression is a non-parametric, non-monotonic, and non-linear exploratory data analysis technique for uncovering structure in the data (Clark and Pregibon, 1992), which allows the nesting of variables that predict gas fluxes (Melling et al., 2005). This technique develops a decision tree based on a binary partitioning algorithm that divides data in a recursive manner until each group is either homogenous or contain a user-defined minimum number of observations. At each split, the algorithm considers each explanatory variable and chooses the one that results in the greatest reduction in deviance. This approach usually creates an overly complex decision tree that needs to be pruned in order to convey the most important information i.e. the nodes that explain the largest amount of deviance (Breiman et al., 1993). The advantage of regression trees is the transparency of results, and the relative importance of inputs can be easily assessed. Regression trees are generally

regarded as superior to standard regression with respect to capturing interactions and non-addictive behavior (Mathsoft, 1999). The tree regression analysis was performed using S-PLUS 2000 (Mathsoft, 1999).

3. Results

3.1. Environmental variables

The mean air temperatures in the forest, sago and oil palm ecosystems were 27.4±0.19, 33.0±0.42 and 30.9±0.35 °C, respectively. The mean soil temperatures at 5 cm for sago and oil palm ecosystems were about 28.2±0.19 and 28.0±0.17 °C, respectively, whilst the forest showed a lower mean temperature of 25.9±0.11 °C. For soil temperature at 10 cm, the forest also showed the lowest temperature of 26.0±0.10 °C followed by sago at 27.1±0.08 °C and oil palm at 27.6±0.14 °C.

The rainfall and water table patterns in the three ecosystems also followed a similar seasonal variation with the highest recorded in January at all sites (Fig. 1a and b). The monthly rainfall in the forest, sago and oil palm ecosystems ranged from 33–418, 37–610 and 43–537 mm, respectively. The highest mean water table in the sago ecosystem was 27.4 cm compared with 45.3 cm in forest and 60.2 cm in oil palm ecosystem.

Similarly, soil water-filled pore space (WFPS) was the highest in sago ecosystem with a mean of 78.1% compared with 57.6 and 60.4% in forest and oil palm ecosystem, respectively (Fig. 1c). In contrast, RH was highest in the forest followed by oil palm and sago ecosystems. Their monthly RH ranges from 70–99, 43–85 and 48–77%, respectively (Fig. 1d).

3.2. Methane flux

The monthly CH₄ fluxes varied significantly between the three ecosystems (Table 2). The seasonal variation in CH₄ fluxes in forest and oil palm ecosystems did not follow the measured climatic and environmental variables whereas in sago ecosystem, it correlated with the monthly rainfall ($r=0.73$) and depth of water table ($r=0.73$) (Figs. 2 and 3).

Both CH₄ emission and uptake were observed in all the three ecosystems. The CH₄ fluxes in forest ecosystem ranged from -4.53 to 8.40 μg C m⁻² h⁻¹, oil palm ecosystem -32.78 to 4.17 μg C m⁻² h⁻¹ and sago ecosystem -7.44 to 102.06 μg C m⁻² h⁻¹. The highest CH₄ emission rate in forest ecosystem was recorded in December 2002 and a month later in oil palm and sago ecosystems. The highest CH₄ uptake rates were recorded in different months being February 2003, August 2002 and December 2002 for forest, oil palm and sago ecosystem, respectively.

The repeated measure analysis showed that ecosystem, time and ecosystem×time were significantly different (Table 2). Annually, both forest and sago ecosystems were

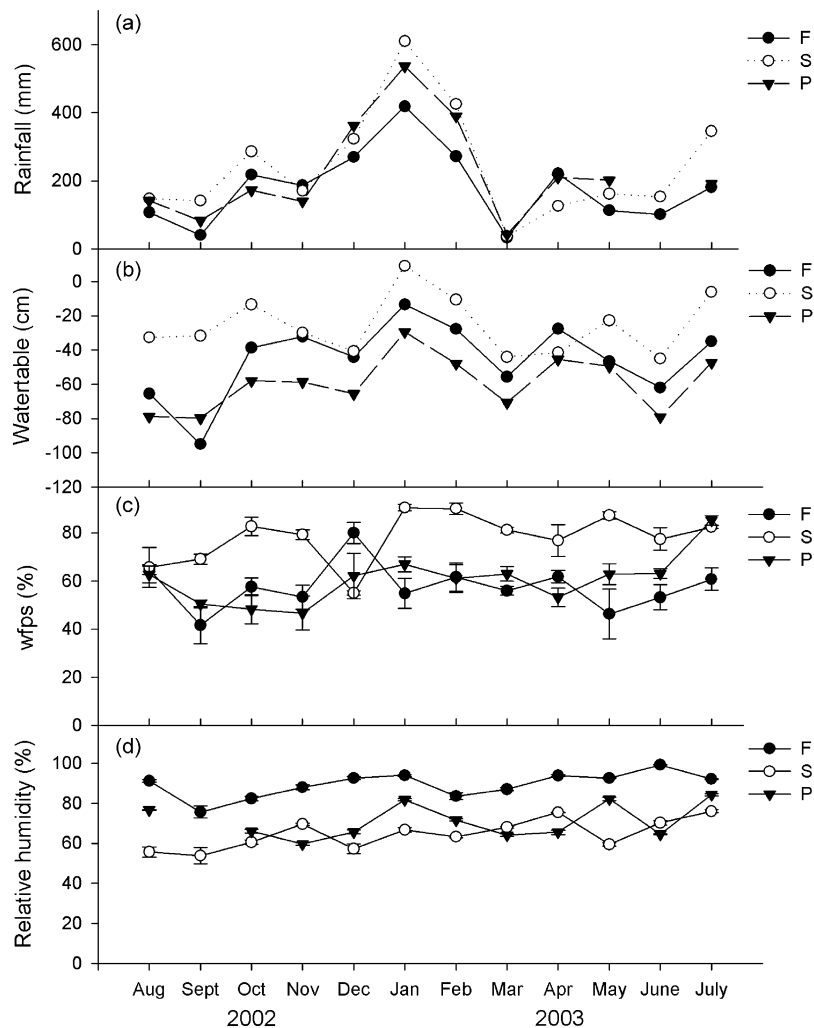


Fig. 1. Rainfall (a), watertable (b), water-filled pore space (c), and relative humidity (d). Data represent means \pm SE ($n=3$). Error bars indicate SE of the mean. Symbols without error bars have errors smaller than the symbols.

CH₄ sources (Table 2) with sago ecosystem producing an order higher than forest. The oil palm ecosystem was a CH₄ sink with an uptake rate of 15 mg C m⁻² yr⁻¹.

3.3. Soil gas concentrations

The mean annual concentrations of CH₄ in the air above soil surface were 2.28 ± 0.08 ppm for forest, 2.21 ± 0.05 ppm for sago and 2.32 ± 0.08 ppm for oil palm ecosystem (Fig. 3a–c). Depth profiles of soil CH₄ concentrations in forest and oil palm ecosystems had similar patterns at all sampling dates (Fig. 3a and b). Both have discontinuities in CH₄ gradients at approximately 20 cm depth. The soil air CH₄ concentrations in the sago ecosystem increased throughout the depth profile (Fig. 3b). The highest soil CH₄ concentration for all the ecosystems occurred at 80 cm depth but in the forest ecosystem, it was only 33.7 ± 7.0 ppm compared with 1086 ± 123.2 and 1465 ± 309.9 ppm in sago and oil palm ecosystems respectively.

3.4. Regression trees

The regression tree method selected RH, water table and WFPS as predictor variables for CH₄ flux in forest ecosystem (Fig. 4). The RH was the most important splitting variable in determining the CH₄ uptake and emission; the critical value being 90.55%. The highest mean CH₄ emission of $9.23 \mu\text{g C m}^{-2} \text{h}^{-1}$ occurred when the RH

Table 2
Soil CH₄ flux and cumulative flux for forest, sago and oil palm ecosystem

Ecosystem	Methane flux, $\mu\text{g C m}^{-2} \text{h}^{-1}$	Cumulative flux, $\text{mg C m}^{-2} \text{yr}^{-1}$
Forest	2.27 ± 1.37	18.34
Sago	22.06 ± 5.68	179.54
Oil palm	-3.58 ± 2.28	-15.14

Note: (a) The p -value for ecosystem was 0.005. The SE of differences of means for ecosystem was 4.91. (b) The p -value for time was 0.05. The SE of differences of means for time was 7.51. (c) The p -value for ecosystem \times time was 0.047. The SE of differences of means for ecosystem \times time was 13.38.

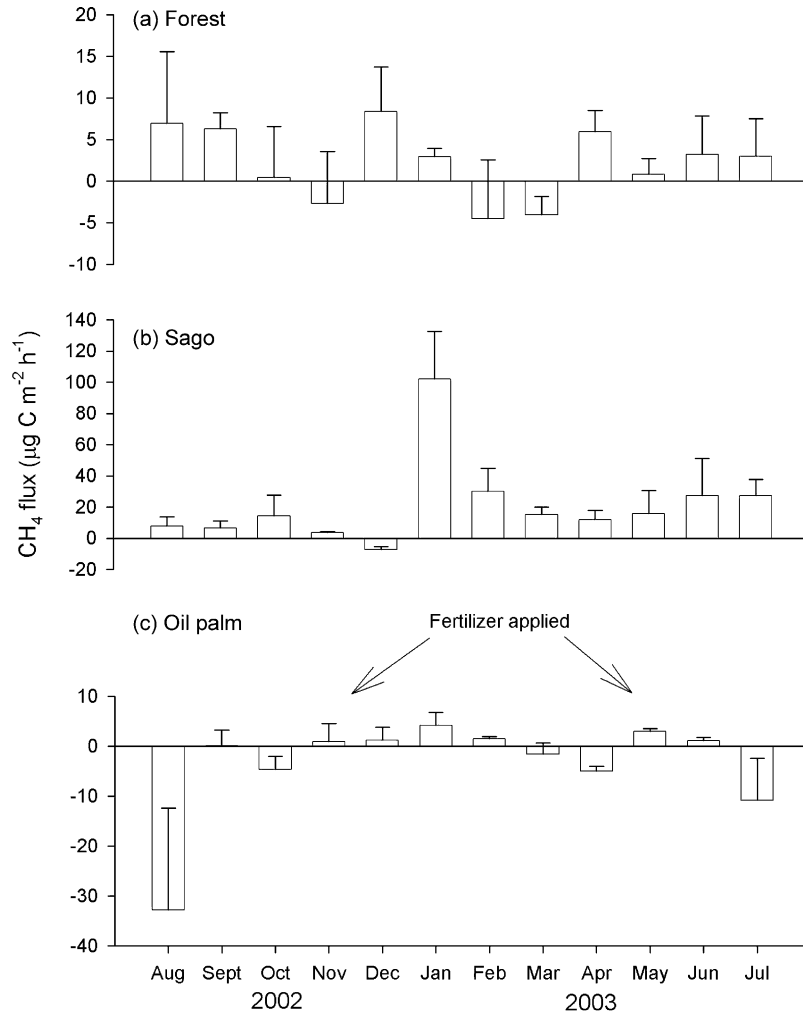


Fig. 2. Monthly methane flux ($\mu\text{g C m}^{-2} \text{h}^{-1}$) at the forest, sago and oil palm ecosystem. Data represent means \pm SE ($n=3$). Error bars indicate SE of the mean. Symbols without error bars have errors smaller than the symbols.

was more than 90.55% and water table more than 49 cm. The highest mean CH_4 uptake of $5.42 \mu\text{g C m}^{-2} \text{h}^{-1}$ occurred when the RH was less than 90.55%, and WFPS more than 54.38%.

Unlike the forest ecosystems, water table, soil temperature at 10 cm and WFPS were chosen as the predictor variables in the sago ecosystem (Fig. 5). The first split for the CH_4 flux in the sago ecosystem was water table with

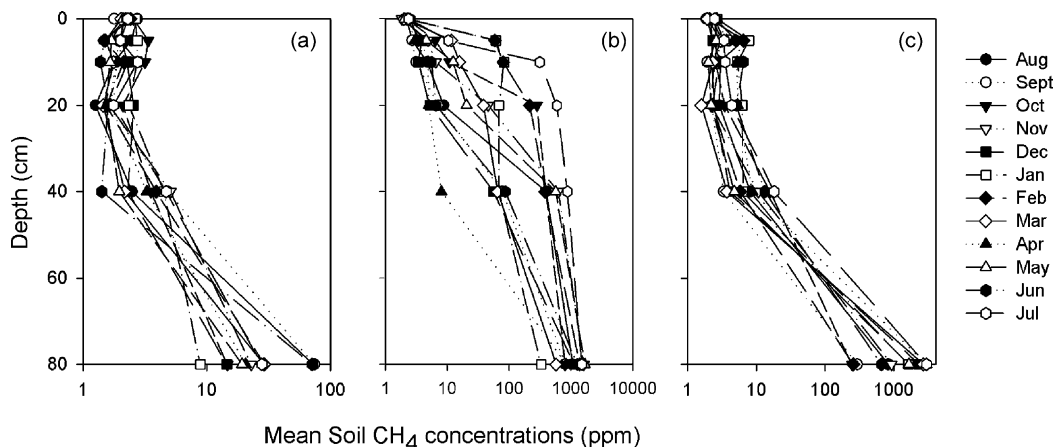


Fig. 3. Mean soil profile CH_4 concentration (ppm) of forest (a), sago (b) and oil palm (c) ecosystem.

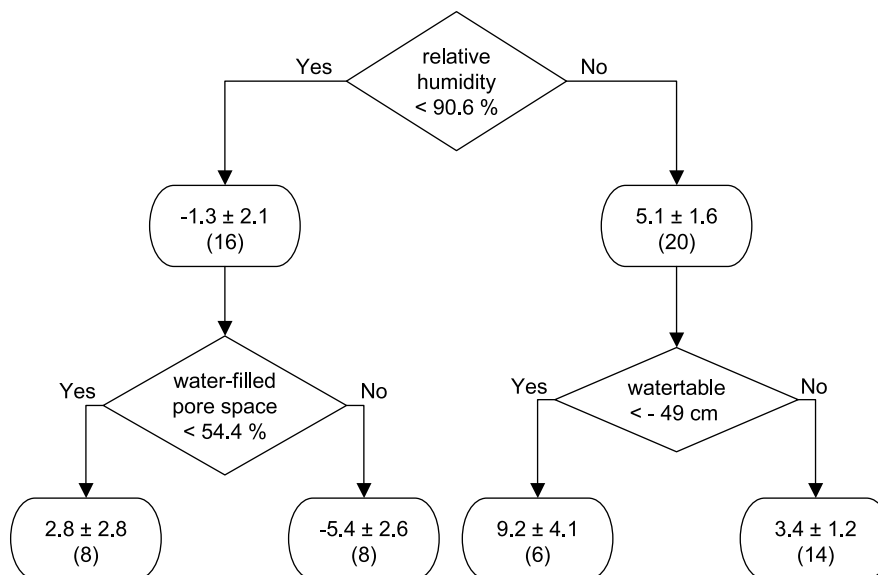


Fig. 4. Regression tree for forest ecosystem ($R^2=0.33$). The CH_4 flux ($\mu\text{g C m}^{-2} \text{h}^{-1}$) in the group is shown in each box. Values are means \pm SE. The number in bracket is the count of samples in the group.

a critical value of 17.2 cm. The highest mean CH_4 emission of $78.7 \mu\text{g C m}^{-2} \text{h}^{-1}$ occurred when the water table was less than 17.2 cm and WFPS greater than 87.6%. On the other hand, water table more than 17.2 cm and soil temperature at 10 cm less than 27.3°C resulted in the lowest CH_4 emission of $1.7 \mu\text{g C m}^{-2} \text{h}^{-1}$.

For the oil palm ecosystem, water table, WFPS and soil temperature at 5 cm were chosen as the predictor variables (Fig. 6). Water table was used to separate between low and high CH_4 uptake with critical value at 77.4 cm. The highest mean CH_4 uptake of $17.44 \mu\text{g C m}^{-2} \text{h}^{-1}$ occurred when the water table was more than 77.4 cm. Highest mean CH_4

emission of $1.13 \mu\text{g C m}^{-2} \text{h}^{-1}$ occurred when the water table was less than 77.4 cm, WFPS less than 72.5% and soil temperature at 5 cm less than 29°C .

4. Discussion

4.1. Methane flux and soil gas concentrations

No distinct seasonal variation was observed in the methane flux in both the forest and oil palm ecosystems even though precipitation and water table depth were

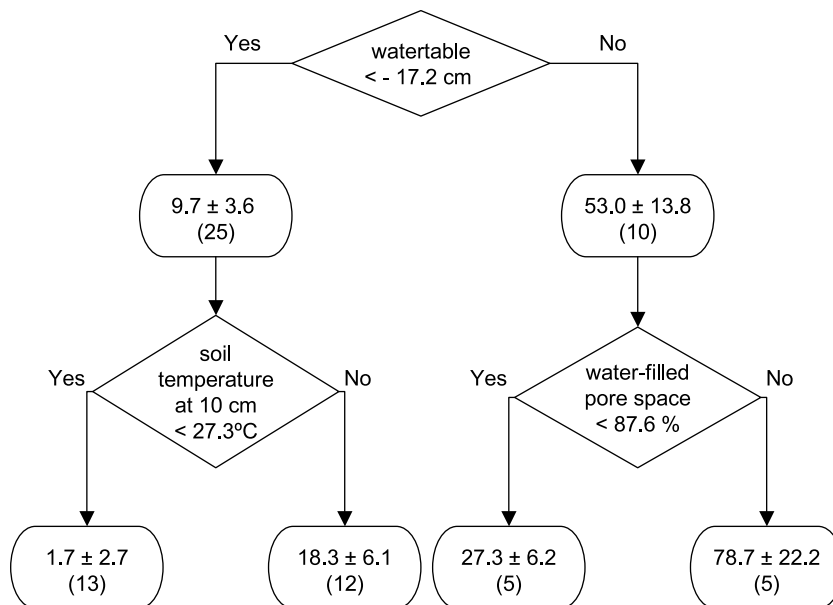


Fig. 5. Regression tree for sago ecosystem ($R^2=0.57$). The CH_4 flux ($\mu\text{g C m}^{-2} \text{h}^{-1}$) in the group is shown in each box. Values are means \pm SE. The number in bracket is the count of samples in the group.

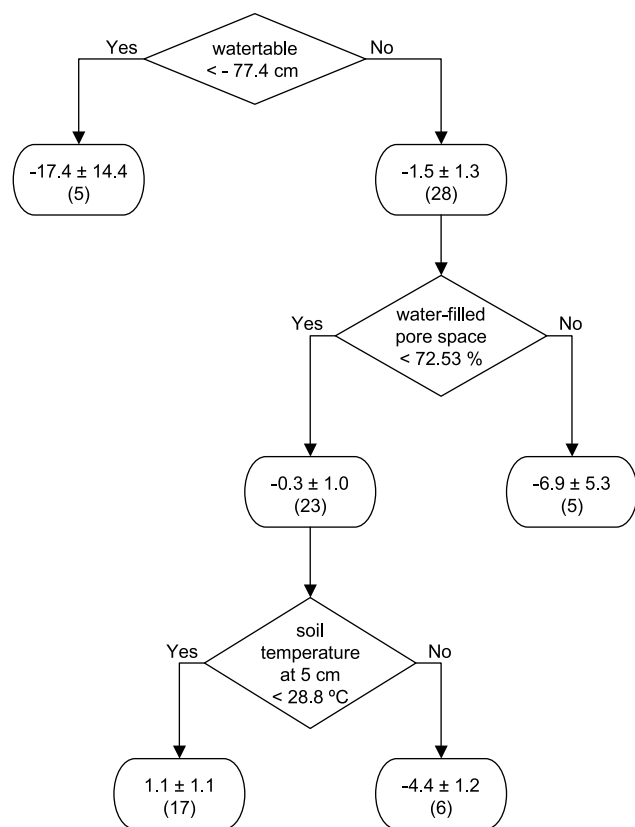


Fig. 6. Regression tree for oil palm ecosystem ($R^2=0.22$). The CH_4 flux ($\mu\text{g C m}^{-2} \text{h}^{-1}$) in the group is shown in each box. Values are means \pm SE. The number in bracket is the count of samples in the group.

observed to have distinct seasonal variation. However, in the sago ecosystem, the seasonal variation in CH_4 flux was positively correlated with rainfall due to the necessary retention of high water table for the crop which increased its susceptibility to flooding (and anaerobic conditions) as shown in January 2003 resulting in high CH_4 emission (Fig. 2).

Using the regression tree approach, we found that different dominant predictor environmental variables had influenced the variations in CH_4 flux in each ecosystem. They were RH for forest (Fig. 4) and water table for both sago (Fig. 5) and oil palm ecosystems (Fig. 6).

Although, RH was well correlated with rainfall and water table, it showed a larger monthly fluctuation indicating its higher sensitivity to environmental changes. It probably had an indirect effect, since, it was associated with high water table and WFPS resulting in larger CH_4 production and emission. RH had no significant effect on CH_4 fluxes in oil palm and sago ecosystems probably due to their more open canopies resulting in lower RH and poorer correlation with water table and WFPS.

Water table was the most important environmental variable that controlled CH_4 fluxes in both the sago and oil palm ecosystems (Figs. 5 and 6), which was agreeable with other researchers (Moore and Dalva, 1993;

Martikainen et al., 1992). In fact, the largest monthly averaged CH_4 emission rates were observed in January 2003 for both sago and oil palm ecosystems when the highest water table was measured at the sites, which probably caused an anoxic condition that was essential for methanogenesis. In the oil palm ecosystem, the water table was also used to separate high and low CH_4 uptake where low water table depth generally resulted in the former (Fig. 6). This might be attributed to the thicker aerobic layer with lower water table, which probably increased the oxidation of CH_4 .

When the WFPS was very high ($> 87.6\%$) as in the sago ecosystem (Fig. 5), high CH_4 emission occurred probably due to high process of methanogenesis. Under the forest and oil palm ecosystem, where the upper horizons were more aerobic, the increase in WFPS may probably have facilitate CH_4 consumption resulting in CH_4 uptake. In the sago ecosystem, an increase in temperature would increase CH_4 emission due to higher gas diffusion rate. However, in the oil palm ecosystem, which has a thick aerobic layer, the higher temperature probably also had increased CH_4 oxidation resulting in higher CH_4 uptake.

Upon cultivation of sago and oil palm on tropical peatland, the CH_4 gas concentration had increased more than 10-fold as shown at 80 cm depth (Fig. 3a–c). This might be ascribed to the increased in soil temperature and decomposition rate as indicated by the lower C:N ratio (Table 1) after land conversion.

The marked increase in CH_4 gas concentration with deeper soil depth (Fig. 3a–c) was probably an interaction of higher WFPS and greater anaerobicity, CH_4 diffusion rate through the peat profile, and CH_4 emission rate. In forest and oil palm ecosystems where CH_4 emissions were not dominant, there were clear declines in CH_4 gas concentrations at 20 cm depth due to diffusion and CH_4 consumption. In the sago ecosystem where CH_4 emissions were dominant, clear CH_4 gradients were observed above 20 cm depth only. This indicated a slow diffusion of CH_4 from the lower depth probably due to high water table.

4.2. Effect of types of ecosystem on CH_4 flux

CH_4 fluxes from the three ecosystems on tropical peatland showed high variation, both temporally and spatially. This has also been reported by Waddington and Roulet (1996), Turetsky et al. (2002) and Inubushi et al. (2003) for different types of peat ecosystem in both temperate and tropical regions. The CH_4 fluxes for forest with a range of -4.53 to $8.40 \mu\text{g C m}^{-2} \text{h}^{-1}$ were similar to grassland on peat in Netherlands (van den Pol-van Dasselaar et al., 1997). The annual soil CH_4 emission of $18.34 \text{ mg C m}^{-2} \text{yr}^{-1}$ for the forest ecosystem, $179.54 \text{ mg C m}^{-2} \text{yr}^{-1}$ for the sago ecosystem and $-15.14 \text{ mg C m}^{-2} \text{yr}^{-1}$ for oil palm ecosystem (Table 2) were similar to those observed by other researchers (Jauhainen et al., 2001; Morishita, Unpublished) but they

were very much lower than those reported by Bartlett and Harriss (1993) and Inubushi et al. (2003).

Forest had a well-developed soil structure and a more permeable surface layer whereby it can easily emit CH₄. This was supported by the slight increases in CH₄ concentration with depth indicating high gas permeability (Fig. 3a). The low water table in the forest probably contributed to the minimal CH₄ flux to the atmosphere because CH₄ produced at depth was oxidized as it diffused upward through the aerobic surface peat layers (Roulet and Moore, 1995; Turetsky et al., 2002).

Upon cultivation to sago, the annual CH₄ emission increased to 179.54 mg C m⁻² yr⁻¹ which was similar to the unfertilized grassland (Glatzel and Stahr, 2001). This shift from a low to a high CH₄ emission might be attributed to the well controlled, high water table, which enhanced the potential for high rates of methanogenesis and CH₄ emissions. Furthermore, higher temperature and low C:N ratio (Table 1) would also contribute to the higher rates of methanogenesis and CH₄ emissions.

The cultivation of tropical peatland primary forest to oil palm promoted CH₄ oxidation due to the lowering of water table by drainage which increased the thickness of aerobic soil layer. This improved CH₄ uptake making the oil palm ecosystem a CH₄ sink. Similar results have also been reported for soils drained for agriculture, which were generally attributed to the reduced CH₄ fluxes and higher CH₄ oxidation (Crill et al., 1988; Roulet and Moore, 1995; Nyakanen et al., 1995; Maljanen et al., 2003).

Drainage and compaction in the oil palm ecosystem had also increased the soil bulk density by 33% (Table 1), thus reducing the diffusion of CH₄ and oxygen (Ball et al., 1997). Hansen et al. (1993) also reported that compaction of agricultural soil can reduce CH₄ uptake by 52% due to above reasons. The higher bulk density would similarly increase the micropores resulting in an increase in moisture retention and probably the amount of anaerobic sites even at lower water content. These anaerobic sites had stimulated the increase in CH₄ production but with the reduced CH₄ diffusion, led to a higher accumulation of soil gas concentration as compared to the forest ecosystem (Fig. 3a and c).

The positive effects of water table and compaction on CH₄ emission in tropical peatland are also found in the boreal and temperate zones.

5. Conclusion

The underlying environmental factors that cause the variations in CH₄ flux between the three peatland ecosystems in Sarawak, Malaysia differed significantly being RH in forest ecosystem and water table in oil palm and sago ecosystems. The conversion of tropical forest on peat to sago increased CH₄ emission by 10-fold due to the necessary retention of high water table. Conversely, its

conversion to oil palm resulted in CH₄ uptake which might be attributed to the lowering of water table and soil compaction. These results have important implications for land-use planning and agricultural management on tropical peatland especially to minimize the adverse effects of human activity on CH₄ emission to the atmosphere.

Acknowledgements

We thank Margaret Abat, Zakri bin Besri, Donny Sudid and Gan bin Haip for assistance in the laboratory or field. We also wish to thank Dr Takuji Sawamoto and Dr Tomoaki Morishita on their assistance in verifying the methods. We acknowledge Gan Huang Huang for her assistance in statistical analyses. This study was supported by IRPA Grant for Scientific Research from the Ministry of Science, Technology and Environment, Malaysia.

References

- Ball, B.C., Dobbie, K.E., Parker, J.P., Smith, K.A., 1997. The influence of gas transport and porosity on methane oxidation in soils. *Journal of Geophysical Research* 102, 23301–23308.
- Bartlett, K.B., Harriss, R.C., 1993. Review and assessment of methane emissions from wetlands. *Chemosphere* 26, 261–320.
- Breiman, L., Friedman, J.H., Olshen, R.A., Stone, C.J., 1993. *Regression Trees*. CRC Press, Boca Raton, FL.
- Christensen, T.R., Jonasson, S., Callaghan, T.V., Hvstrom, M., 1995. Spatial variation in high-latitude CH₄ flux along a transect across Siberian and European tundra environments. *Journal of Geophysical Research* 100, 21035–21045.
- Clark, L.A., Pregibon, D., 1992. Tree-based models, in: Hastie, T.J. (Ed.), *Statistical Models*. Wadsworth, Pacific Grove, CA.
- Crill, P.M., 1991. Seasonal patterns of methane uptake and carbon dioxide release by a temperate woodland soil. *Global Biogeochemical Cycles* 5, 319–334.
- Crill, P.M., Bartlett, K.B., Harris, R.C., Gorham, E., Verry, E.S., Sebacher, D.I., Madzar, I., Sanner, W., 1988. Methane flux from Minnesota peatlands. *Global Biogeochemical Cycles* 2, 371–384.
- FAO-UNESCO, 1974. *Soil Map of the World*. vol. 1. Legend. Unesco, Paris.
- Flessa, H., Wild, U., Klemisch, M., Pfadenhauer, J., 1998. Nitrous oxide and methane fluxes from organic soils under agriculture. *European Journal of Soil Science* 49, 327–335.
- GenStat, 2002. *GenStat® for Windows (Release 6.1): Reference Manual and Guides*. VSN International, Oxford, UK.
- Glatzel, S., Stahr, K., 2001. Methane and nitrous oxide exchange in differently fertilized grassland in Southern Germany. *Plant and Soil* 231, 21–35.
- Hansen, S., Maehlum, J.E., Bakken, L.R., 1993. N₂O and CH₄ fluxes in soil influenced by fertilization and tractor traffic. *Soil Biology & Biochemistry* 25, 621–630.
- Inubushi, K., Furukawa, Y., Hadi, A., Purnomo, E., Tsuruta, H., 2003. Seasonal changes of CO₂, CH₄ and N₂O fluxes in relation to land-use change in tropical peatlands located in coastal area of South Kalimantan. *Chemosphere* 52, 603–608.
- IPCC, 2001a. *Climate change 2001: the scientific basis*, in: Houghton, J.T., Ding, Y., Griggs, D.J., Noguier, M., van der Linden, P.J., Xiaosu, D.

- (Eds.), Contribution of Working Group 1 to the Third Assessment Report of the Intergovernmental Panel on Climate Change. Cambridge University Press, Cambridge, UK.
- IPCC, 2001b. Climate change 2001; synthesis report, in: Watson, R.T., The Core Writing Team (Eds.), A Contribution of Working Groups I, II, and III to the Third Assessment Report of the Intergovernmental Panel on Climate Change. Cambridge University Press, Cambridge, UK, p. 398.
- Jauhiainen, J., Heikkinen, J., Martikainen, P.J., Vasander, H., 2001. CO₂ and CH₄ fluxes in pristine peat swamp forest and peatland converted to agriculture in central Kalimantan, Indonesia. *International Peat Journal* 11, 43–49.
- Maljanen, M., Liikanen, A., Silvola, J., Martikainen, P.J., 2003. Methane fluxes on agricultural and forested boreal organic soils. *Soil Use and Management* 19, 73–79.
- Maltby, E., Immirzi, P., 1993. Carbon dynamics in peatlands and other wetland soils. Regional and global perspectives. *Chemosphere* 27, 999–1023.
- Martikainen, P.J., Nykanen, H., Crill, P., Silvola, J., 1992. The effect of changing water table on methane fluxes at two Finnish mire sites. *Suo* 43, 237–240.
- Mathsoft, 1999. SPLUS 2000 Professional. Release 2. User's Manual. Mathsoft, Cambridge, MA.
- McAuliffe, C., 1971. GC determination of solutes by multiple phase equilibration. *Chemical Technology* 1, 46–51.
- Melling, L., Hatano, R., Goh, K.J., 2005. Soil CO₂ flux from three ecosystems in tropical peatland of Sarawak, Malaysia. *Tellus* 57B, 1–11.
- Moore, T.R., Dalva, M., 1993. The influence of temperature and water table position on carbon dioxide and methane emissions from laboratory columns of peatland soils. *Journal of Soil Science* 44, 651–664.
- Morishita, T., Unpublished PhD Thesis. Effect of various environmental changes on CH₄ dynamics in soil ecosystems. Hokkaido University.
- Mosier, A., Schimel, D., Valentine, D., Bronson, K., Parton, W., 1991. Methane and nitrous oxide fluxes in native, fertilized and cultivated grasslands. *Nature* 350, 330–332.
- Norman, J.M., Kucharik, C.J., Gower, S.T., Baldocchi, D.D., Crill, P.M., Rayment, M., Savage, K., Striegl, R.G., 1997. A comparison of six methods for measuring soil-surface carbon dioxide fluxes. *Journal of Geophysical Research* 102, 28771–28777.
- Nykanen, H., Alm, J., Lang, K., Silvola, T., Martikainen, P.J., 1995. Emissions of CH₄, N₂O and CO₂ from a virgin fen drained for grassland in Finland. *Journal of Biogeography* 22, 351–357.
- Roulet, N.T., Moore, T.R., 1995. The effect of forestry drainage practices on the emission of methane from Northern peatlands. *Canadian Journal of Forest Research* 25, 491–499.
- Smith, K.A., Dobbie, K.E., Ball, B.C., Bakken, L.R., Sitaula, B.K., Hansen, S., Brumme, R., Borken, W., Christensen, S., Prieme, A., Fowler, D., Macdonald, J.A., Skiba, U., Klemetsson, L., Kasimir-Klemetsson, A., Degorska, A., Orlanski, P., 2000. Oxidation of atmospheric methane in Northern European soils, comparison with other ecosystems, and uncertainties in the global terrestrial sink. *Global Change Biology* 6, 791–803.
- Soil Survey Staff, 1992. Keys to Soil Taxonomy, fifth ed. SMSS Technical Monograph No.19. Blacksberg, Virginia, USA.
- Sorenson, K.M., 1993. Indonesian peat swamp forests and their role as a carbon sink. *Chemosphere* 27, 1065–1082.
- Turetsky, M.R., Wieder, R.K., Vitt, D.H., 2002. Boreal peatland C fluxes under varying permafrost regimes. *Soil Biology & Biochemistry* 34, 907–912.
- van den Pol-van Dasselaar, van Beusichem, M.L., Oenema, O., 1997. Effects of grassland management on the emission of methane from intensively managed grasslands on peat soil. *Plant and Soil* 189, 1–9.
- Waddington, J.M., Roulet, N.T., 1996. Atmosphere-wetland carbon exchanges; scale dependency of CO₂ and CH₄ exchange on the developmental topography of a peatland. *Global Biogeochemical Cycles* 10, 233–245.
- Watson, R.T., Rodhe, H., Oeschger, H., Siegenthaler, U., 1990. Greenhouse gases and aerosols, in: Houghton, J.T., Jenkins, G.J., Ephraums, J.J. (Eds.), *Climate Change: the IPCC Scientific Assessment*. Cambridge University Press, Cambridge, New York.