

Impact of forest plantation on methane emissions from tropical peatland

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Abstract

Tropical peatlands are a known source of methane (CH₄) to the atmosphere, but their contribution to atmospheric CH₄ is poorly constrained. Since the 1980s, extensive areas of the peatlands in Southeast Asia have experienced land-cover change to smallholder agriculture and forest plantations. This land-cover change generally involves lowering of groundwater level (GWL), as well as modification of vegetation type, both of which potentially influence CH₄ emissions. We measured CH₄ exchanges at the landscape scale using eddy covariance towers over two land-cover types in tropical peatland in Sumatra, Indonesia: (a) a natural forest and (b) an *Acacia crassicaarpa* plantation. Annual CH₄ exchanges over the natural forest (9.1 ± 0.9 g CH₄ m⁻² year⁻¹) were around twice as high as those of the *Acacia* plantation (4.7 ± 1.5 g CH₄ m⁻² year⁻¹). Results highlight that tropical peatlands are significant CH₄ sources, and probably have a greater impact on global atmospheric CH₄ concentrations than previously thought. Observations showed a clear diurnal variation in CH₄ exchange over the natural forest where the GWL was higher than 40 cm below the ground surface. The diurnal variation in CH₄ exchanges was strongly correlated with associated changes in the canopy conductance to water vapor, photosynthetic photon flux density, vapor pressure deficit, and air temperature. The absence of a comparable diurnal pattern in CH₄ exchange over the *Acacia* plantation may be the result of the GWL being consistently below the root zone. Our results, which are among the first eddy covariance CH₄ exchange data reported for any tropical peatland, should help to reduce the uncertainty in the estimation of CH₄ emissions from a globally important ecosystem, provide a more complete estimate of the impact of land-cover change on tropical peat, and develop science-based peatland management practices that help to minimize greenhouse gas emissions.

KEYWORDS

Acacia crassicaarpa, eddy covariance measurements, forest plantation, Indonesia, land-use change, methane emissions, peatland management, tropical peatlands

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1 | INTRODUCTION

Methane (CH_4) is the second most important anthropogenic greenhouse gas after carbon dioxide (CO_2) and its concentration is continuing to increase (Dlugokencky, 2019; Nisbet et al., 2019). The global warming potential (GWP) of CH_4 is 34 times that of CO_2 on a 100 year basis when including climate-carbon feedbacks (Myhre et al., 2013). Due to its short atmospheric life span of about 10 years and relatively high GWP, there is increasing interest in reducing CH_4 emissions in order to meet global temperature targets (Collins et al., 2018). Current and future regional and global CH_4 budgets and mitigation strategies require better quantitative and process-based understanding of CH_4 sources, pathways, and removals under climate and land-use change (Saunois et al., 2016).

Natural wetlands are the single largest source of atmospheric CH_4 (Kirschke et al., 2013; Poulter et al., 2017; Saunois et al., 2016). The coastal lowlands of Southeast Asia host one-third of tropical peatlands, with the majority located in Indonesia (Gumbricht et al., 2017; Page, Rieley, & Banks, 2011), and they represent unrecognized and poorly understood components of the CH_4 cycle (Pangala, Moore, Hornibrook, & Gauci, 2013; Wong et al., 2018). Since the 1980s, extensive areas of Southeast Asian peatlands have experienced land-cover changes (Miettinen, Shi, & Liew, 2016; Wijedasa et al., 2018), driven by transmigration, local population growth, and ongoing economic development. The 2015 land-cover distribution for the insular Southeast Asian peatlands reveals that half of all former peatland forest is managed as either small-holder agriculture or industrial plantation, while around 29% is characterized as intact or degraded natural peat swamp forest (Miettinen et al., 2016). The remaining 21% of the peatlands are covered by open undeveloped areas, fern, low/tall shrub, and secondary regrowth forest (Miettinen et al., 2016). Agriculture and forest plantation on peatlands require the maintenance of groundwater level (GWL) below the root zone to support the required level of productive growth. Maintaining the GWL below the surface alters the CH_4 dynamic by weakening the potential for CH_4 production and increasing the potential for CH_4 oxidation in the upper peat layers (Furukawa, Inubushi, Ali, Itang, & Tsuruta, 2005; Melling, Hatano, & Goh, 2005). Given the potential importance of tropical peatlands in global CH_4 budgets, it is important to understand any effects of land-cover changes on CH_4 emissions from tropical peatlands.

When the balance between CH_4 production and consumption is positive, CH_4 can be emitted to the atmosphere via: (a) diffusion from soil and water surfaces, (b) ebullition from water surfaces, or (c) vegetation-mediated transport through aerenchymatous and air-filled tissues in herbaceous plants and trees (Jauhiainen & Silvennoinen, 2012; Pangala et al., 2013). In addition, CH_4 can be emitted from terrestrial arthropods such as termites (Jeeva, Bignell, Eggleton, & Maryati, 1999) and plants producing CH_4 in aerobic conditions (Keppler, Hamilton, Brass, & Röckmann, 2006). The contribution of each pathway to total ecosystem CH_4 exchange varies within and among peatland ecosystems depending on surface microtopography

(hummock vs. hollow), GWL, peat temperature, vegetation composition and structure, and land-use practices (Melling et al., 2005; Pangala et al., 2013). Variation in plant physiological processes driven by solar radiation might substantially influence vegetation-mediated transports as observed in northern peatlands (Kim, Verma, Billesbach, & Clement, 1998; Long, Flanagan, & Cai, 2010; Nisbet et al., 2009; van der Nat, Middelburg, van Meteren, & Wielemaers, 1998). Thus, significant spatial and temporal variability in CH_4 emissions from tropical peatlands can be anticipated, yet available data rarely allow analysis of how such variability influences annual emissions.

Measurements of CH_4 emissions from tropical peatlands are sparse and have focused mainly on soil surfaces based on small-scale chamber measurements (Furukawa et al., 2005; Hadi et al., 2005; Inubushi, Hadi, Okazaki, & Yonebayashi, 1998; Ishikura et al., 2019; Jauhiainen, Limin, Silvennoinen, & Vasander, 2008; Melling et al., 2005). The large and heterogeneous nature of forest vegetation, together with dynamic biotic and abiotic processes, makes it difficult to measure vegetation-mediated transport accurately using chamber systems (Barba et al., 2018). Notably, vegetation-mediated transport, principally through trees, could be 62%–87% of total ecosystem CH_4 exchange (Pangala et al., 2013) and represents a “science frontier” in our understanding of biosphere-atmosphere exchange in forests (Barba et al., 2018).

Knowledge of the magnitude of CH_4 exchanges including all existing sources and removals in tropical peatland ecosystems is limited (Pangala et al., 2013; Sakabe, Itoh, Hirano, & Kusin, 2018; Tang et al., 2018; Wong et al., 2018). Many process-based CH_4 models lack sufficient details in their treatment and parameterization of transport pathways to derive reliable emissions estimates (Gedney, Huntingford, Comyn-Platt, & Wiltshire, 2019; Parker et al., 2018). This leads to uncertainty in estimates of the current and future contribution of tropical peatlands to regional and global CH_4 budgets (Saunois et al., 2016).

Given these uncertainties, we need to improve our understanding of the spatiotemporal and environmental variability that drive exchange strength and direction in order to better understand the potential CH_4 exchanges that may result from any future climate or land-use change scenarios. Micrometeorological methods (such as eddy covariance) provide half-hourly measurements of turbulent CH_4 exchanges between an entire ecosystem and the atmosphere above the vegetation canopy (Aubinet et al., 2000). Hence, eddy covariance measurements incorporate all existing CH_4 sources and removals that can vary significantly within an ecosystem in both space and time arising from variation in environmental conditions.

In the above context, we used the eddy covariance technique to measure net ecosystem CH_4 exchange over two land-covers in a single peatland hydrological unit on the Kampar Peninsula in Sumatra, Indonesia: (a) a natural forest, and (b) a forest plantation (*Acacia crassicaarpa*). Measurements were conducted for more than four site-years (October 2016–May 2019 over the *Acacia* plantation and June 2017–May 2019 over the natural forest). The main

objectives of this study were to: (a) determine the magnitudes of CH_4 exchanges from tropical peatlands while incorporating all existing sources and removals, and (b) understand the link between temporally varying CH_4 exchanges and associated changes in the environmental controls.

We hypothesized that a lower GWL would reduce vegetation-mediated CH_4 transport to the atmosphere in the managed peatland. We evaluated this hypothesis over timescales ranging from diurnal to annual. These results were then used to quantify the impact of *Acacia* plantation, considering the change in CH_4 exchanges due to the associated altered landscape, as one component of the ecosystem greenhouse gas balance. Finally, we considered the relevance of these results for tropical peatland greenhouse gas emissions reporting, climate change mitigation policies and land-use management.

2 | MATERIALS AND METHODS

2.1 | Study area

The Kampar Peninsula is a coastal tropical peatland of around 700,000 ha (Figure 1a). This ombrotrophic (acidic and nutrient-poor) peatland is largely formed within the past 8,000 years (Dommain, Couwenberg, & Joosten, 2011). The study area has a humid tropical climate (warm year-round) with average monthly air temperature ranging from 29 to 32°C (Badan Meteorologi, Klimatologi dan Geofisika, 1994–2017 data). Average annual rainfall for the last 5 years (2014–2018) is ~1,800 mm with two wet seasons (March–April and October–December) and two dry seasons (January–March and May–August). The peninsula is characterized by a large, relatively intact central forest area surrounded by a mosaic of smallholder

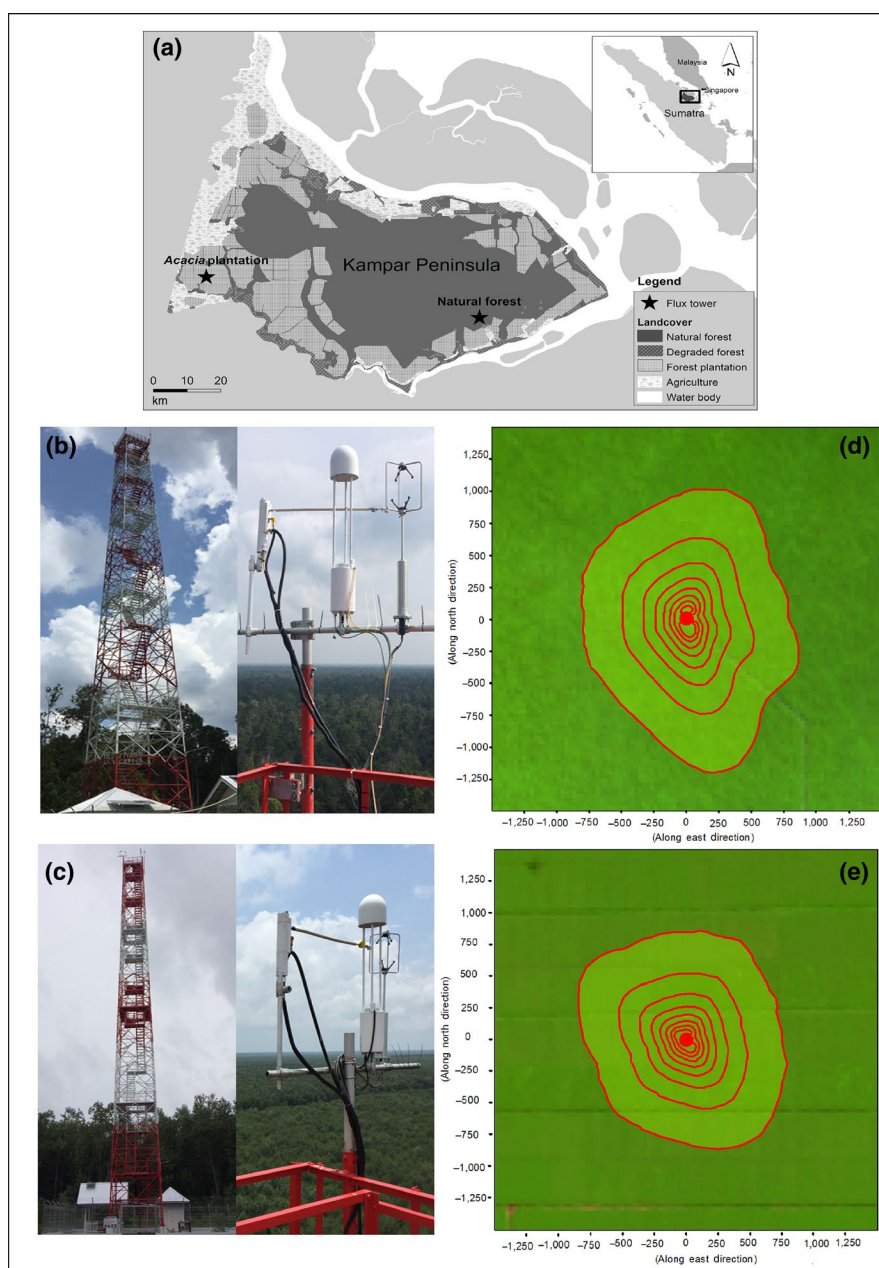


FIGURE 1 Land-cover map of the Kampar Peninsula, Sumatra, Indonesia and the location of research flux tower sites (a), photos of the eddy covariance instruments installed at the top of the tower at the natural forest (b), and the *Acacia* plantation (c), and integrated eddy covariance footprint contour lines from 10% to 80% in 10% intervals over the natural forest for June 2017–May 2019 (d), and the *Acacia* plantation for October 2016–May 2019 (e)

agricultural land (largely oil palm, *Elaeis guineensis*), and industrial fiber wood plantation (largely *A. crassicaarpa*), smaller secondary forest areas, and undeveloped open and degraded land (Figure 1a; Miettinen et al., 2016). Natural forest and *Acacia* plantation together occupy around 80% of the peninsula (Figure 1a).

Above-canopy eddy covariance flux towers were established at the *Acacia* plantation and the natural forest in 2016 and 2017, respectively, for the purpose of measuring net ecosystem CO₂ and CH₄ exchange (Figure 1b,c; note that CO₂ flux measurements will be reported separately). The terrain around the towers is flat (slope <0.05%) and land-cover and topography are homogenous for at least 3 km in all directions at both sites, ensuring a good fetch and a consistent land-cover-related signal regardless of wind direction. The relatively close proximity of the natural forest and the *Acacia* plantation sites (~80 km apart) within the same peatland hydrological unit avoids potentially confounding variables such as climatic differences, past natural succession, and to some extent geomorphological formation (Figure 1a). Thus, although it is inherently difficult and expensive to replicate flux measurements using the eddy covariance technique, our sites should provide a robust and unbiased basis for evaluating the impact of land-cover change (from peat swamp forest to *Acacia* plantation) on CH₄ exchanges.

The natural forest is characterized as pristine peat swamp forest (Miettinen et al., 2016). The forest structure is mixed, and the canopy is uneven with the tallest canopy in a range of 28–35 m. Tree density with diameter at breast height >5 cm was 1,343 trees per hectare. The dominant tree species of the overstory are *Shorea uliginosa*, *Calophyllum ferrugineum*, and *Syzygium* spp.; together they represent around 75% of the overstory vegetation (Table 1). The understory is dominated by *Pandanus* spp., *Cyrtostachys renda*, and *Nepenthes* spp. The forest floor is uneven with a hummock-hollow microtopography, and covered with tree debris, root mat, and leaf litter. Hollow surfaces are often 20–40 cm lower than hummock tops. The average area ratio of hollow to hummock was 3:1 around

the tower. The surface peat type is fibric and the average peat thickness is $\sim 9 \pm 1$ m in the area surrounding the tower. The surface peat pH is 3.6 ± 0.1 and the GWL fluctuates seasonally with the rainfall variation (see Section 3). An integrated climatologic footprint analysis (Kljun, Calanca, Rotach, & Schmid, 2015) indicated that approximately 80% of fluxes were derived within 1,200 m in the upwind direction (Figure 1d), and thus originated within the pristine peat swamp forest as characterized by Miettinen et al. (2016).

At the forest plantation, *A. crassicaarpa* trees are grown for fiber production on a 5 year rotation from planting to harvesting. When measurements began in October 2016, the trees were already at the end of the plantation cycle. In March–April 2017, the mature trees, which had achieved an average height of 20 m, were harvested. Replanting at a density of 1,667 trees per hectare (3 m × 2 m spacing) took place within 2 weeks after harvesting. One kg boiler wood ash per tree was applied around the seedlings during planting as per the standard operational procedure, without additional fertilizers. In May 2019, 2 years after replanting, the canopy height was ~17 m. The ground surface within the plantation area is relatively even, without a hummock-hollow microtopography, and with very little understory vegetation. The surface peat type is hemic and the average peat thickness is $\sim 7 \pm 0.8$ m in the surrounding area of the tower. The surface peat pH is 3.4 ± 0.1 . GWLs in plantation are actively managed to support the required level of productive growth via an extensive network of topographically defined water management zones, controlled by outlet sluices, and supported by large-scale and continuous rainfall and water level monitoring (Evans et al., 2019). Water management zones comprise of ditches and canals (also used for transportation). The integrated climatologic footprint analysis (Kljun et al., 2015) indicated that (a) approximately 80% of fluxes were estimated to occur within 1,000 m in the upwind direction, and thus originated within the *Acacia* plantation; and (b) the water surface of ditches and canals represented 2.1% of the flux footprint (Figure 1e).

Parameter	Natural forest	<i>Acacia</i> plantation
Tower location	Latitude: 0°23'42.735"N Longitude: 102°45'52.382"E	Latitude: 0°30'57.221"N Longitude: 102°2'11.090"E
Tower height (m)	48	40
Average canopy height (m)	32 ± 6	17 ± 6
Dominant understory species	<i>Nepenthes</i> , <i>Pandanus</i> , <i>Cyrtostachys renda</i>	Not available
Dominant trees species	<i>Shorea uliginosa</i> , <i>Calophyllum</i> <i>ferrugineum</i> , <i>Syzygium</i> spp., <i>Camnosperma macrophylla</i> , <i>Tetramerista glabra</i> , <i>Palaquium burckii</i>	<i>Acacia crassicaarpa</i>
Peat depth (m)	9 ± 1.0	7 ± 0.8
Surface (0–50 cm) peat type	Fibric	Hemic
Surface (0–50 cm) peat bulk density (g/cm ³)	0.08 ± 0.03	0.14 ± 0.02
Surface (0–50 cm) peat pH	3.6 ± 0.10	3.4 ± 0.03

TABLE 1 Site characteristics. Value represents average ± standard deviation

2.2 | Net ecosystem CH₄ exchange and environmental variables measurements

The eddy covariance system consisted of an open path gas analyzer (LI-7700, LI-COR Inc.) to measure the atmospheric CH₄ concentration, and a three-dimensional sonic anemometer (WindMaster Pro 3-Axis Anemometer, Gill Instruments Limited) to measure orthogonal components of wind speed fluctuations. The mirrors of the CH₄ analyzer were self-cleaned either at 5:00 (local time) every day or if the received signal strength indicator (RSSI) dropped below 20%. Furthermore, the upper and lower mirrors of the CH₄ analyzer were manually cleaned on a biweekly basis. Dew condensation, rain, and dirty window events were excluded using an RSSI value of 20% because CH₄ data become noisy below this threshold (Chu et al., 2014; McDermitt et al., 2011). Water vapor densities were measured using an enclosed path CO₂/H₂O analyzer (LI-7210, LI-COR Inc.). The sensors were mounted at the top of the tower to ensure complete exposure in all directions (Figure 1b,c). The raw turbulence eddy covariance data were recorded at 10 Hz using an analyzer interface unit (LI-7550, LI-COR Inc.) and stored on a removable flash disk (APRO, Industrial Grade USB Flash Disk).

Quantum sensors (LI-190SL-50, LI-COR Inc.) were mounted at the top of the towers to measure the incoming photosynthetic photon flux density (PPFD, $\mu\text{mol m}^{-2} \text{day}^{-1}$). Relative humidity (RH, %), and thereby the vapor pressure deficit (VPD, hPa) and air temperature (T_{air} , °C) were measured using the air temperature and humidity probes (Vaisala HMP155 Humidity Temperature Probe, Vaisala, Inc), which were installed inside a ventilated radiation shield at the top of the towers. Estimates of canopy conductance to water vapor, an indicator of transpiration, were made from measurements of latent heat flux following the approach described in Long et al. (2010).

Daily rainfall (mm/day) rates were measured using three and six manual bucket systems within 10 km distance from the tower location in the natural forest and the *Acacia* plantation, respectively. Manual bucket systems were installed 1.5 m above the ground, in an open area so that rainfall was not intercepted by the tree canopy. Soil temperature (T_{soil} , °C) was measured at 15 cm below the peat hollow surface using temperature probe (Stevens Hydra Probe II, Stevens Water Monitoring Systems, Inc.) with three replicates at each tower site. GWLs (m) were monitored as the water elevation relative to the ground surface (taking the base of the hollows as a datum) every 30 min using a GWL logger (Solinst Levellogger Model 3001). The GWL logger was placed in a perforated polyvinyl chloride (PVC) tube that was inserted vertically into the peat at a distance approximately 30 m away from the towers. The GWL logger also recorded temperature at 150 cm below the peat surface that is below GWL. Additionally, PVC poles were randomly distributed within a 3 km radius around the tower locations to monitor GWL fortnightly. All meteorological sensors took measurements every second and were recorded as one minute averages with a datalogger (Sutron Model 9210 XLITE, Sutron Corporation).

All measuring systems were powered using solar panels along with a rechargeable battery system (65 Watt Solar Package, SunWize Power & Battery). Owing to the large power requirement

and cost of a separate CH₄ analyzer, we could not conduct CH₄ profile measurements to calculate CH₄ storage below the flux measurement height (Finnigan, 2006). In theory, accumulated CH₄ below the canopy during nighttime is likely to be released and measured by the EC system following the onset of turbulence after sunrise and the bias on annual sums should be negligible (Xu et al., 2019).

2.3 | Eddy covariance data processing

Net ecosystem CH₄ exchange (NEE-CH₄) was computed from the 10 Hz vertical wind velocity and CH₄ concentration data using EddyPro software (version 6.2.0, LI-COR Inc.) at a standard averaging interval of half hour period (Aubinet et al., 2000). A de-spiking procedure was applied to detect and eliminate individual out-of-range values for vertical wind velocity and CH₄ concentrations (Vickers & Mahrt, 1997). De-trending was carried out using the block averaging method. A coordinate correction was applied to force the average vertical wind velocity to zero by the planar fit method (Wilczak, Oncley, & Stage, 2001). Frequency response loss corrections were applied to compensate the flux losses at different frequencies (Massman, 2000, 2001; Moncrieff, Clement, Finnigan, & Meyers, 2004). Fluctuations in CH₄ density due to temperature (thermal expansion) and water vapor (dilution) were corrected using the Webb-Pearman-Leuning correction (Webb, Pearman, & Leuning, 1980) and spectroscopic effects taken into account by EddyPro (Burba, Anderson, & Komissarov, 2019). Differences between deployment-specific variables, that is, sensor separation distance and instrument placement, were considered while processing the data. We adopted the standard meteorological notation whereby a positive value of NEE-CH₄ represents a net CH₄ flux to the atmosphere, and a negative value indicates net CH₄ uptake from the atmosphere (Aubinet et al., 2000). All NEE-CH₄ values in the paper are reported in mass of CH₄ per unit area per time.

After a set of quality controls, the numbers of high-quality measurements during the course of the study were 38% and 29% for the natural forest and the *Acacia* plantation, respectively, including measuring system malfunctions due to lightning strikes and power supply failure (Table 2). In other words, we obtained a total

TABLE 2 Summary of the percentage of half-hourly net ecosystem CH₄ exchange data that were removed using various quality control criteria and accepted high quality data

	Natural forest	Acacia plantation
Stationarity criteria (Mauder et al., 2013)	19%	24%
Extreme outlier (Papale et al., 2006)	6%	10%
u^* threshold (Wutzler et al., 2018)	12%	7%
Instrument malfunction	25%	30%
Accepted high quality data	38%	29%

of 13,637 and 13,548 half-hourly measurements that met all quality criteria for the natural forest and the *Acacia* plantation, respectively. We gap-filled both low-quality and missing data due to instrument malfunction, as is commonly done in eddy covariance studies.

We applied two gap-filling approaches (a) mean diurnal course (MDC; Dengel, Levy, Grace, Jones, & Skiba, 2011; Sakabe et al., 2018; Wong et al., 2018) and (b) marginal distribution sampling (MDS; Alberto et al., 2014; Dalmagro et al., 2019; Tang et al., 2018) using the REddyProc package (Wutzler et al., 2018). The MDC is a simple interpolation technique where the missing value is replaced with the averaged value of the adjacent days at exactly that time of day (Falge et al., 2001). The MDS considers the covariation of the fluxes with the environmental variables and the temporal autocorrelation of fluxes. We performed MDS gap filling separately for the daytime (06:00–16:00 hr) and the nighttime (18:00–06:00 hr) data. GWL and PPFD were used during the daytime, whereas GWL and T_{soil} above the GWL were used during the nighttime gap-filling. The emissions were similar from both methods at the natural forest (Mann–Whitney test; $p = .34$, Table 4), whereas the emissions were different from both methods at the *Acacia* plantation (Mann–Whitney test; $p < .05$, Table 4). To provide a conservative estimate, we used the average of the MDC and MDS approaches. Flux random uncertainty (σ_1) was calculated following Finkelstein and Sims (2001). The standard deviation of three different flux values derived from friction velocity (u^*) thresholds of 5th, 50th, and 95th percentiles were applied as an uncertainty due to u^* threshold (σ_2) using the REddyProc package (Wutzler et al., 2018). The flux uncertainty due to gap-filling (σ_3) was calculated as the standard deviation of the binned records used to fill the missing value (Wutzler et al., 2018). The total uncertainty in NEE-CH₄ was calculated with the law of propagation of errors (Deventer et al., 2019). Only high quality measurements were used in the qualitative analysis (Figures 4a,b, 5, and 6) and gap-filled data were used in the quantitative analysis (Figures 3g,h, 4c,d, and 7 and Table 4).

2.4 | Statistical analyses

Differences between groups of data were examined using *t* test in GraphPad Prism (GraphPad Software, Inc., v5.04). The choice of the non-parametric test (Mann–Whitney test compare median values) was dependent on non-normal behavior of the datasets. All statistical tests used a significance level of 5%.

3 | RESULTS

3.1 | Environmental conditions

During the course of the study, the PPFD, T_{air} , VPD, and canopy conductance to water vapor above the canopy showed typical diurnal patterns reaching their maximum around noon (Figure 2a–h). No significant diurnal variation in T_{soil} below the GWL was observed at either site (Figure 2i,j). The diurnal variation in T_{soil}

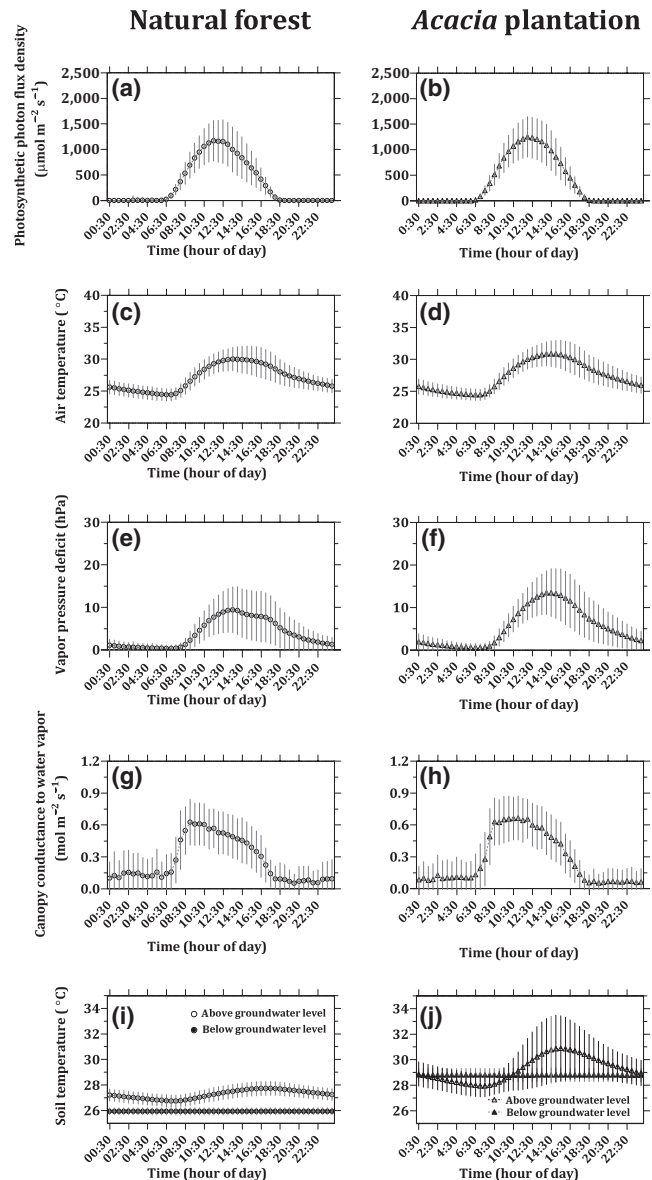


FIGURE 2 Average diurnal variation in the photosynthetic photon flux density (a, b), air temperature (c, d), vapor pressure deficit (e, f), canopy conductance to water vapor (g, h), and soil temperature (i, j) at the natural forest (left panels) and the *Acacia* plantation (right panels). Data were binned by time of day and then presented for all days during the measurement periods. The error bars show the standard deviation

above the GWL was small ($<1^{\circ}\text{C}$) at the natural forest, due to the closed canopy and high GWL (Figure 2i). Before canopy closure, the *Acacia* plantation showed a clear diurnal variation (amplitude of 3°C) in the T_{soil} above the GWL, but after canopy closure, the observed diurnal T_{soil} above the GWL amplitude was similar to the natural forest.

Daily average T_{air} fluctuated between 23.3 and 29.9°C as a function of rainfall and cloud cover, without showing any clear seasonality (Figure 3a,b). Daily average T_{soil} above the GWL at the natural forest ranged from 25.6 to 28.3°C depending on the GWL and cloudiness, again without clear seasonality. At the *Acacia*

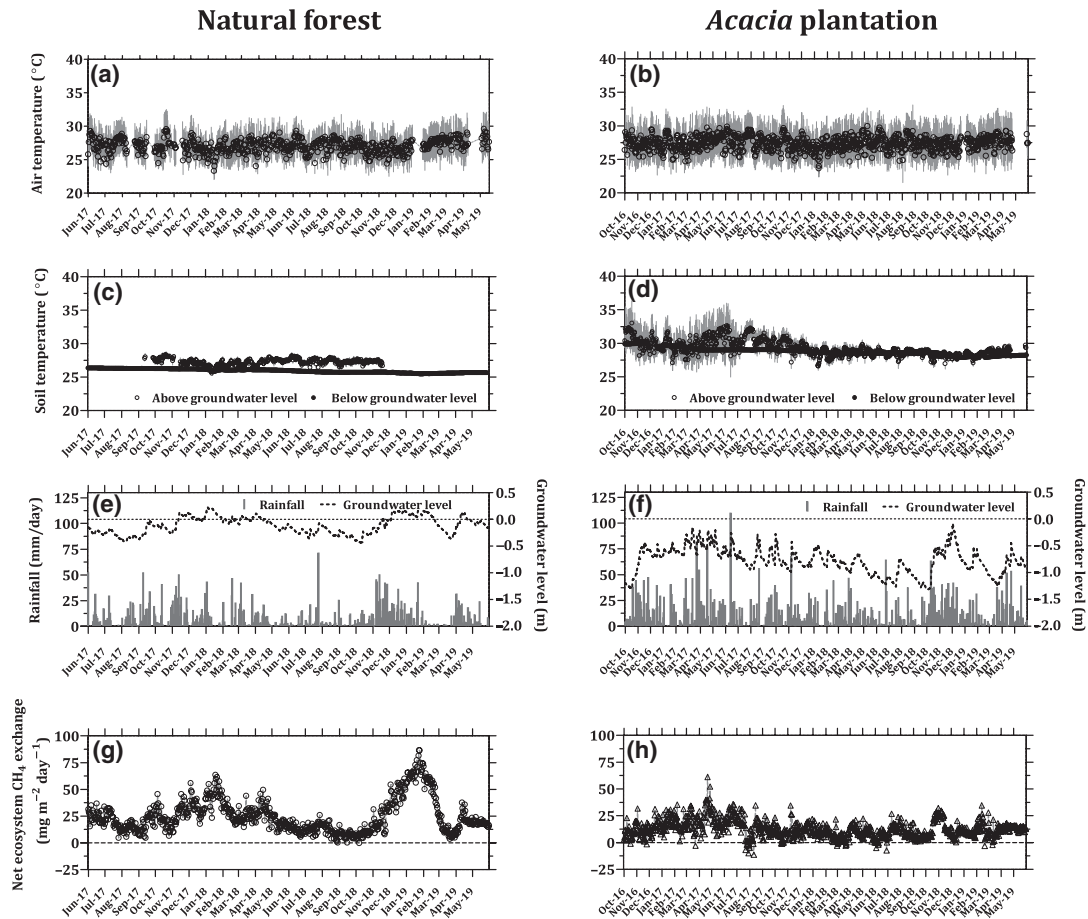


FIGURE 3 Variations in daily air temperature (a, b), soil temperature above and below groundwater level (c, d), cumulative rainfall and groundwater level (e, f), and net ecosystem CH₄ exchanges (g, h) at the natural forest (left panels) and the Acacia plantation (right panels). The vertical bar in panels (a, b, c, d) represents standard deviation. Positive value of groundwater level indicates water level above the peat surface, and negative values indicate water level below the soil surface

TABLE 3 The average and standard deviation of environmental variables

Site	Natural forest			Acacia plantation		
	Year 1 (June 2017– May 2018)	Year 2 (June 2018– May 2019)	Study period (June 2017– May 2019)	Year 1 (June 2017– May 2018)	Year 2 (June 2018– May 2019)	Study period (October 2016– May 2019)
Air temperature (°C)	26.9 ± 1.05	27.1 ± 0.9	27.0 ± 1.0	27.4 ± 1.1	27.4 ± 0.9	27.4 ± 1.0
Photosynthetic photon flux density (μmol m ⁻² s ⁻¹)	342 ± 90	328 ± 83	335 ± 87	343 ± 78	365 ± 83	358 ± 86
Vapor pressure deficit (hPa)	3.79 ± 2.23	3.72 ± 1.7	3.75 ± 1.98	5.85 ± 2.15	4.97 ± 2.31	5.52 ± 2.33
Canopy conductance to water vapor (mol m ⁻² s ⁻¹)	0.32 ± 0.11	0.34 ± 0.11	0.33 ± 0.11	0.29 ± 0.09	0.38 ± 0.1	0.32 ± 0.11
Cumulative rainfall (mm)	2,019	1,756	3,775	1,907	2,034	5,705
Groundwater level (m)	0.20 ± 0.13	0.28 ± 0.15	0.24 ± 0.14	0.69 ± 0.09	0.75 ± 0.17	0.73 ± 0.14
Soil temperature above groundwater level (at 15 cm below ground surface; °C)	27.2 ± 0.5	27.4 ± 0.3	27.3 ± 0.4	29.6 ± 1.3	28.5 ± 0.4	29.3 ± 1.2
Soil temperature below groundwater level (at 150 cm below ground surface; °C)	26.2 ± 0.1	25.7 ± 0.1	25.9 ± 0.3	28.8 ± 0.2	28.3 ± 0.3	28.8 ± 0.5
Wind speed (m/s)	2.12 ± 0.45	2.11 ± 0.43	2.11 ± 0.44	2.32 ± 0.56	1.98 ± 0.38	2.16 ± 0.53
Friction velocity (m/s)	0.2 ± 0.04	0.2 ± 0.04	0.2 ± 0.04	0.19 ± 0.04	0.17 ± 0.06	0.18 ± 0.05

plantation, daily average T_{soil} above the GWL ranged from 26.6 to 33.0°C as a function of canopy development, GWL, and cloudiness, without any clear seasonality (Figure 3c,d). The daily average T_{soil} above and below the GWL at the natural forest was statistically different (t test; $p < .05$; Table 3) and around $\sim 2^\circ\text{C}$ lower than at the *Acacia* plantation (Figure 3c,d). The average VPD at the natural forest of 3.7 ± 1.9 hPa was significantly lower (40%) than the average of 5.6 ± 2.2 hPa at the *Acacia* plantation (t test; $p < .05$; Table 3).

At both sites, the daily cumulative rainfall was highly variable, ranging from 0 to 137 mm (Figure 3e,f), but did not significantly differ between sites (Mann–Whitney test; $p > .05$). Annual average rainfall was 1,887 and 1,970 mm for the natural forest and the *Acacia* plantation, respectively, similar to the previously reported average annual rainfall of the study area as a whole. GWL at the natural forest showed periodic sharp rises and steady decreases corresponding to rain events (Figure 3e). At the natural forest, the GWL rose up to 23 cm above the peat surface in the wet season, then in the late dry season reached -44 cm. The annual GWL pattern at the natural forest was almost the same as reported for other undrained peatland in Southeast Asia (Cobb et al., 2017). At the *Acacia* plantation, GWL rose up during rain events, but remained always below the ground surface (Figure 3f). During the study period, the average GWL from six sampling points around the natural forest tower of -0.24 ± 0.14 m was significantly shallower than that of -0.73 ± 0.14 m from 10–20 sampling points around the *Acacia* plantation tower (Mann–Whitney test; $p < .05$).

3.2 | Net ecosystem CH_4 exchanges

At both sites, the NEE-CH_4 showed a marked peak at around 07:00–10:30 hr (Figure 4a,b), consistent with flushing of CH_4 accumulated

in the vegetation canopy at night following the onset of turbulent mixing in the morning (Wong et al., 2018). NEE-CH_4 over the natural forest remained much higher than the nighttime during the remaining day hours and began to decline late in the afternoon (Figure 4a). NEE-CH_4 over the *Acacia* plantation began to decline and reached levels similar to the nighttime values after around 10:30 hr (Figure 4b). Thus, the diurnal variation in NEE-CH_4 was more pronounced over the natural forest (Figure 4a,b).

In order to avoid bias due to flushing of accumulated CH_4 , we considered nighttime NEE-CH_4 from 18:30 to 10:30 hr and daytime from 10:30 to 18:30 hr. This threshold might principally be site specific, but offered an opportunity to examine the diurnal variation in the NEE-CH_4 over our sites. Over the natural forest, daytime median NEE-CH_4 was more than three times higher (29 $\text{mg m}^{-2} \text{ day}^{-1}$) than at nighttime (8.4 $\text{mg m}^{-2} \text{ day}^{-1}$; Mann–Whitney test; $p < .05$; Figure 4c). Furthermore, daytime median NEE-CH_4 was almost three times higher over the natural forest than over the *Acacia* plantation (Mann–Whitney test; $p < .05$; Figure 4c,d). In contrast, the nighttime medians NEE-CH_4 were 8.3 and 7.9 $\text{mg m}^{-2} \text{ day}^{-1}$, respectively, over the natural forest and the *Acacia* plantation (Figure 4c,d). The diurnal variation in NEE-CH_4 over the natural forest was positively correlated with corresponding changes in canopy conductance to water vapor, PPFD, VPD, and T_{air} (Figure 5a–d). However, we did not observe a corresponding dependency of diurnal NEE-CH_4 on any environmental variables at the *Acacia* plantation (Figure 5e–h).

During the study period, daily NEE-CH_4 ranged from -0.15 to 86.6 $\text{mg m}^{-2} \text{ day}^{-1}$ and -11.3 to 61.2 $\text{mg m}^{-2} \text{ day}^{-1}$ at the natural forest and the *Acacia* plantation, respectively (Figure 3g,h). Daily NEE-CH_4 was almost two times higher (median = 20.7 $\text{mg m}^{-2} \text{ day}^{-1}$ and mean = 25.0 $\text{mg m}^{-2} \text{ day}^{-1}$) over the natural forest than over the *Acacia* plantation (median = 11.6 $\text{mg m}^{-2} \text{ day}^{-1}$ and

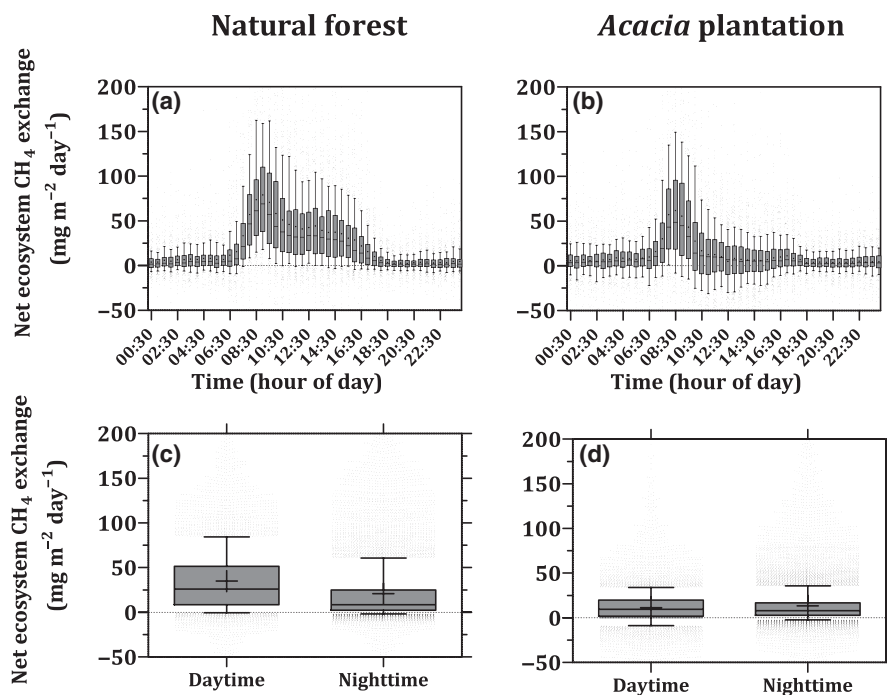


FIGURE 4 Diurnal variation in the net ecosystem CH_4 exchanges (a, b), and daytime (10:30–18:30 hr) and nighttime (18:30–10:30 hr) ranges for net ecosystem CH_4 exchanges (c, d) at the natural forest (left panels), and the *Acacia* plantation (right panels). The boxes show the median and the interquartile range, and whiskers denote the 10–90 range of all values

FIGURE 5 Response of the half-hourly net ecosystem CH_4 exchanges to canopy conductance to water vapor (a, e), photosynthetic photon flux density (b, f), vapor pressure deficit (c, g), and air temperature (d, h) at the natural forest (left panels), and the *Acacia* plantation (right panels). Data were binned by subgroups of 50 values of independent variable and corresponding net ecosystem CH_4 exchange rates and then averaged for the subgroup. The vertical and horizontal bars represent the standard deviation for the subgroup. Note: we excluded measurements from 7:00 to 10:30 hr to avoid the possible bias due to flushing of nighttime accumulated CH_4 . The exclusion of data may have created biases in actual response curves of both ecosystems, but this bias would not change the interpretation

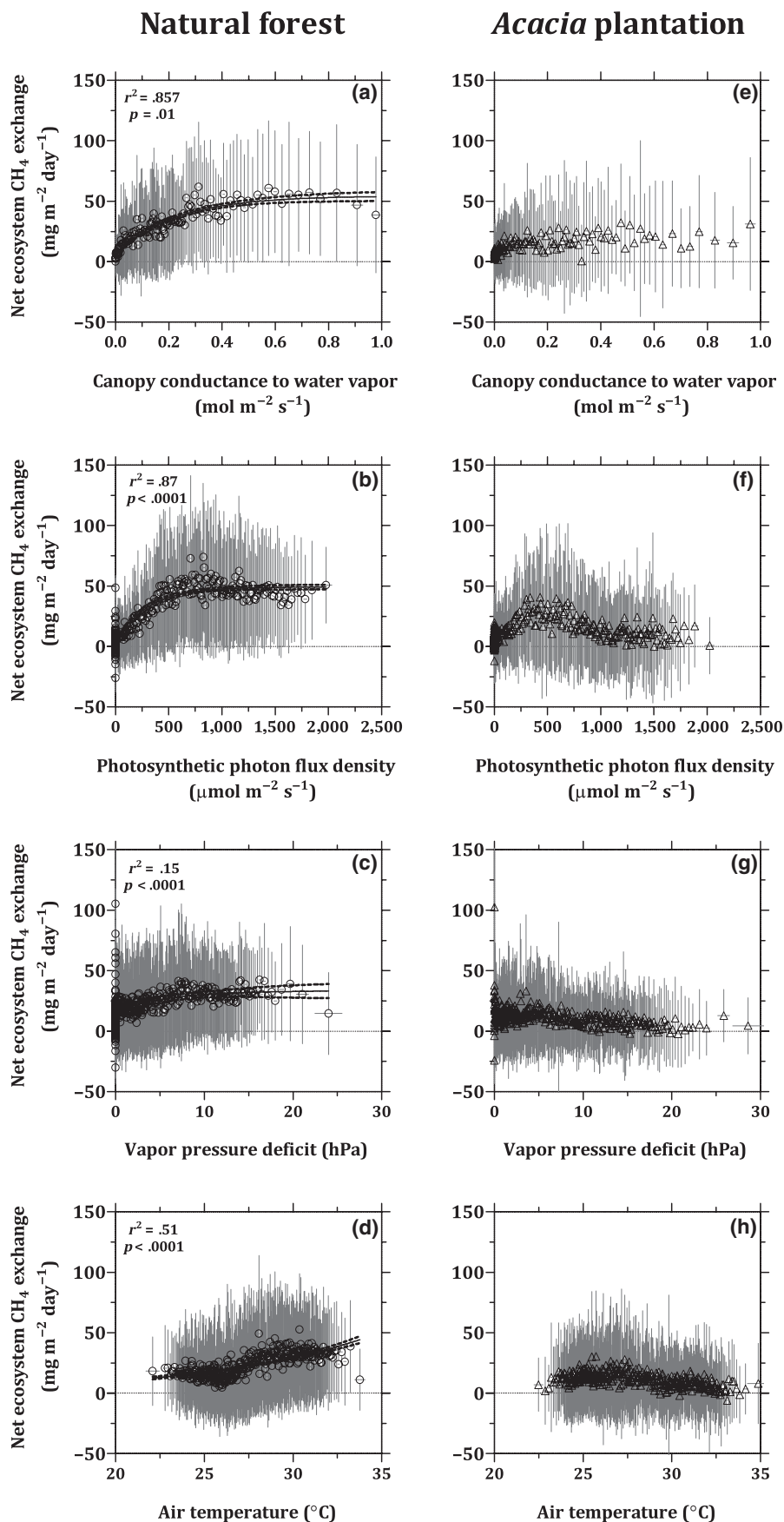


TABLE 4 Net ecosystem CH_4 exchanges over the natural forest and the *Acacia* plantation using the eddy covariance technique. Daily average values are with standard error, whereas cumulative annual values are with cumulative flux uncertainty

	Natural forest				Acacia plantation				Study period including matured <i>Acacia</i> plantation and harvesting activity (October 2016–May 2019)
	Gap-filling approach		Year 1 (June 2017– May 2018)	Year 2 (June 2018– May 2019)	Average (June 2017– May 2019)	Year 1 (June 2017– May 2018)	Year 2 (June 2018– May 2019)	Average (June 2017– May 2019)	
Daily average ($\text{mg m}^{-2} \text{ day}^{-1}$)	Mean diurnal course		26.0 ± 0.3	24.7 ± 0.3	25.3 ± 0.2	12.3 ± 0.2	12.0 ± 0.2	12.2 ± 0.1	13.6 ± 0.1
	Marginal distribution sampling		25.4 ± 0.2	23.7 ± 0.3	24.6 ± 0.2	10.4 ± 0.2	10.8 ± 0.1	10.6 ± 0.1	12 ± 0.1
	Average		25.8 ± 0.2	24.2 ± 0.3	25.0 ± 0.3	11.4 ± 0.2	11.4 ± 0.2	11.4 ± 0.1	12.8 ± 0.1
Cumulative annual ($\text{mg m}^{-2} \text{ year}^{-1}$)	Mean diurnal course		9,510 ± 637	8,997 ± 635	9,255 ± 899	4,476 ± 654	4,332 ± 917	4,404 ± 1,126	4,928 ± 1,444
	Marginal distribution sampling		9,280 ± 716	8,655 ± 615	8,967 ± 944	3,798 ± 693	3,938 ± 870	3,868 ± 1,112	4,547 ± 1,495
	Average		9,395 ± 677	8,826 ± 625	9,111 ± 922	4,137 ± 674	4,135 ± 894	4,136 ± 1,120	4,738 ± 1,470
Impact of <i>Acacia</i> plantation on CH_4 emissions ($\text{mg m}^{-2} \text{ year}^{-1}$)	Average							–4,975 ± 1,451	–4,373 ± 1,735

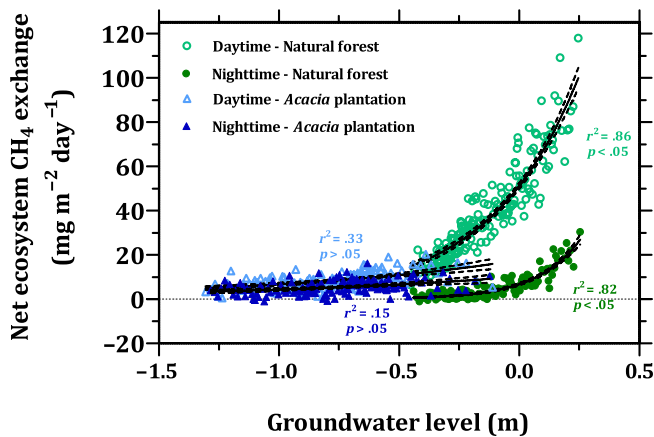


FIGURE 6 The relationship between the half-hourly net ecosystem CH_4 exchange and the groundwater level. Data were binned by subgroups of 50 values of groundwater level and corresponding net ecosystem CH_4 exchange rates and then averaged for the subgroup. Note: we excluded measurements from 7:00 to 10:30 hr to avoid the possible bias due to flushing of nighttime accumulated CH_4 . The exclusion of data may have created biases in actual response curves of both ecosystems, but this bias would not change the interpretation

mean = $12.8 \text{ mg m}^{-2} \text{ day}^{-1}$; Mann-Whitney test; $p < .05$; Table 4). Variation in daytime and nighttime NEE- CH_4 was positively correlated with associated changes in GWL at both sites (Figure 6). Notably, the relationships between NEE- CH_4 and GWL for daytime and nighttime over the natural forest were significantly different, whereas the relationships were quite similar over the *Acacia* plantation (Figure 6). There was no clear relationship between NEE- CH_4 and the T_{soil} either above or below the GWL (data not shown).

Our measurements showed that the natural forest emitted $9.1 \pm 0.9 \text{ g m}^{-2} \text{ year}^{-1}$ to the atmosphere (Table 4). Annual NEE- CH_4 over the *Acacia* plantation were approximately 50% lower than the natural forest, at $4.7 \pm 1.5 \text{ g m}^{-2} \text{ year}^{-1}$, suggesting a net reduction of CH_4 exchanges from natural forest to *Acacia* plantation of $-4.4 \pm 1.7 \text{ g m}^{-2} \text{ year}^{-1}$ (Table 4).

4 | DISCUSSION

4.1 | High GWL supports diurnal variability in NEE- CH_4

Our results show substantial and apparent diurnal variation in the NEE- CH_4 over the natural forest where GWL are high, highlighting the importance of high-frequency ecosystem-scale flux measurements. There is increasing evidence that woody vegetation can emit a significant CH_4 amount to the atmosphere (Barba et al., 2018; Covey & Megonigal, 2019; Pangala et al., 2013, 2017; Pitz & Megonigal, 2017; Rice et al., 2010; Wang et al., 2016; Welch, Gauci, & Sayer, 2019). In tropical peatlands, the majority of root biomass occurs within the upper 50 cm of the peat column (Brady, 1997; Sulistiyanto, 2004), and dissolved CH_4 in the root zone can

be significant (100–1,500 $\mu\text{mol/L}$; Hoyt, 2017; Pangala et al., 2013). The magnitude of vegetation-mediated transport seems to be directly regulated by a well-connected root-stem pathway for the CH_4 transport, although it is strongly (if not primarily) controlled by the availability of dissolved CH_4 in the root zone (Covey & Megonigal, 2019; Pangala et al., 2013, 2017; Waddington, Roulet, & Swanson, 1996). At the natural forest site, *S. uliginosa*, *C. ferrugineum*, and *Syzygium* spp. are the dominant species; together they represent around 75% of the tall-canopy vegetation. *Shorea uliginosa*, *Mesua* sp. 1, and *Xylopia fusca* emit significant CH_4 in tropical peatlands (Pangala et al., 2013). *Mesua* sp. and *C. ferrugineum* belong to the same family. Thus, *S. uliginosa* and *C. ferrugineum* may have contributed significantly to the vegetation-mediated transport.

When the root zone is inundated, changes in biological processes in vegetation driven by solar energy input might be the most important factors controlling diurnal variation in measured NEE- CH_4 (Figure 5a–d), as reported in northern peatlands (Chanton, Whiting, Happell, & Gerard, 1993; Garnet, Megonigal, Litchfield, & Taylor, 2005; Kim et al., 1998; Long et al., 2010; Whiting & Chanton, 1996) and recently reported over tropical peatland (Tang et al., 2018) and flooded forest (Dalmagro et al., 2019). At the natural forest, the observed positive correlation between NEE- CH_4 and canopy conductance to water vapor suggests that CH_4 could be dissolved in the water, absorbed by the roots, transported with sap flow, and emitted through the stem by effervescence (Garnet et al., 2005; Nisbet et al., 2009). In addition, the positive correlation between NEE- CH_4 and PPFD, VPD, and temperature (Figure 5b–d) may suggest vegetation-mediated transport through either diffusion or convective throughflow (Brix, Sorrell, & Orr, 1992; Chanton, Martens, Kelley, Crill, & Showers, 1992; Dacey, 1981). Our results are in line with a study in a temperate forested wetland which showed a sudden decrease in CH_4 emissions from *Betula pubescens* after leaf loss, suggesting physiological control on gas transport (Pangala, Hornibrook, Gowing, & Gauci, 2015). Furthermore, labile organic compounds released from root tissues during photosynthesis and respiration can then be used as substrates by methanogenic archaea, contributing to the diurnal variation in NEE- CH_4 (Chanton et al., 1995; Christensen et al., 2003). Our study did not aim to conduct direct measurements to establish the relative importance of these different processes. Quantifying the pathway-specific emissions and improving our understanding on the impact of root distribution by depth and dissolved CH_4 concentration profile are important future study (Barba et al., 2018; Megonigal, Brewer, & Knee, 2019).

Nighttime and daytime NEE- CH_4 was positively correlated with associated changes in GWL at both sites (Figure 6). Nighttime NEE- CH_4 can be considered as the emissions from soil and water surfaces since there would be negligible vegetation-mediated transport. A higher GWL may support larger CH_4 concentration gradients between the peat surface and the atmosphere. Thus, GWL seems to be the key indirect control on CH_4 emissions via diffusion from soil surfaces (Winton, Flanagan, & Richardson, 2017). Overall, the lack of a difference between nighttime NEE- CH_4 over the natural forest and

the *Acacia* plantation can be attributed to the high GWL at the natural forest and to the potential presence of emissions from the water surfaces of ditches and canals in the *Acacia* plantation (Jauhiainen & Silvennoinen, 2012; Manning, Kho, Hill, Cornulier, & Teh, 2019). In addition, the higher soil temperature at the *Acacia* plantation might have increased CH_4 production (Sjögersten et al., 2018), while the higher peat bulk density and the absence of a hollow-hummock microtopography at the *Acacia* plantation might have lowered CH_4 oxidation by increasing soil moisture content and lowering oxygen diffusion in the peat (Estop-Aragones, Knorr, & Blodau, 2012). For these reasons, the *Acacia* plantation seems to produce higher nighttime CH_4 emissions compared to the natural forest if the same range of GWL (−0.4 to −0.1) at both sites is considered. The difference between daytime and nighttime NEE- CH_4 can be attributed to vegetation-mediated transport. Thus, our estimated vegetation-mediated transport over the natural forest is 71% of the total daytime emissions, which is in line with the published range for tropical peatlands (Pangala et al., 2013). Overall, at the same GWL range, the natural forest emits higher CH_4 as compared to the *Acacia* plantation during the daytime, most likely due to the presence of CH_4 emitting trees (i.e., *S. uliginosa* and *C. ferrugineum*). However, it should be noted that only a few measurements are available for the *Acacia* plantation for the same range of GWL (Figure 6).

The influence of vegetation on CH_4 emissions is strongly dependent on the GWL, and therefore, the interaction among hydrology, vegetation, and CH_4 emissions must be carefully taken into account for process-based modeling (Figure 6). Predicted changes in rainfall amount, intensity, duration, and frequency and water management practices could affect the dynamics of hydrology in tropical peatlands (Ge et al., 2019), and thereby CH_4 emissions (Saunois et al., 2016).

4.2 | GWL controls seasonal variability in NEE- CH_4

The seasonal variation is controlled by the GWL driven by rainfall. Our results show higher NEE- CH_4 during the wet season as compared to the dry season. Other eddy covariance studies in tropical peatlands have reported a similar seasonal pattern in CH_4 emissions (Sakabe et al., 2018; Wong et al., 2018). A study in Amazonian peatland reported lower soil- CH_4 emissions in the wet season as compared to the dry season, where the GWL was 54 cm above the peat surface during the wet season (Teh, Murphy, Berrio, Boom, & Page, 2017). If GWL rises above a limit, soil CH_4 emissions can decrease with flooding depth as gas diffusion may be restricted more as hydrostatic pressure increases along with increasing flooding depth (Ishikura et al., 2019). Furthermore, the standing water can enhance CH_4 oxidation because it would increase dissolved oxygen and prolong traveling time of CH_4 to the atmosphere (Strack, Waddington, & Tuittila, 2004). Notably, Teh et al. (2017) only reported emissions from soil and water surfaces and did not measure vegetation-mediated transport which can be significant in Amazonian wetlands (Pangala et al., 2017). This highlights that seasonality differs from one pathway to another; thus, caution should

be taken when modeling seasonality in CH_4 emissions from tropical peatlands.

In northern peatlands, temperature exerts a strong effect on seasonal variation in CH_4 emissions with an exponential dependence via its influence over enzyme kinetics of CH_4 production and plant growth and development (Desai et al., 2015; Rinne et al., 2007; Tagesson et al., 2012). Observed fluctuations in both T_{air} and T_{soil} in this study are much smaller than those of northern peatlands. During the study period, the T_{soil} below the GWL varied within a very narrow range ($\sim 2^\circ\text{C}$) at both sites. This suggests that variation in T_{soil} would have only a minor effect (if any) on variation in NEE- CH_4 . Furthermore, T_{soil} tended to be higher when GWLs were lower; thus, it is difficult to determine the independent effect (if any) that a change in temperature had on CH_4 production and oxidation (Sjögersten et al., 2018). For example, if CH_4 oxidation above the GWL increased more rapidly (due to the combination of a deeper aerobic zone and higher rates of microbial activity at a higher temperature) than rates of CH_4 production below the GWL, the net effect of warmer and drier conditions would be a lower NEE- CH_4 . Our results suggest that the effects of changing rainfall and land management on peat hydrology will be more important than rising temperature as a driver of changes in tropical peatland CH_4 balance in the future.

4.3 | Low GWL reduces NEE- CH_4 over the *Acacia* plantation

At the *Acacia* plantation, the lower GWL leads to an aerobic root zone (indeed, this is the specific aim of water management in the plantation, to support *Acacia* growth) which is likely to reduce (but not eliminate) CH_4 production and transport. Firstly, aerobic conditions are unfavorable to methanogens and promote methanotrophy (Furukawa et al., 2005; Moore & Roulet, 1993; Strack et al., 2004). Secondly, as most of *Acacia* roots are mainly restricted above GWL in the aerated peat layer, this may result in inadequate CH_4 in the root zone to be taken and transported to the atmosphere. But given the GWL fluctuation, it is possible that when GWL rises after a heavy rain event, some portion of the root system will be below GWL, at least for a few days. However, our measurements over the *Acacia* plantation do not show a diurnal variation in NEE- CH_4 , and this may confirm that the root system remained above the GWL. Finally, it is likely that a substantial fraction of CH_4 emission from the *Acacia* plantation area could be occurring from the open water surface of the ditch and canal network (Evans, Renou-Wilson, & Strack, 2016; Jauhiainen & Silvennoinen, 2012; Manning et al., 2019), and therefore subject to different environmental controls (Deshmukh et al., 2014). The CH_4 uptake rates in the *Acacia* plantation are similar to those previously reported over tropical peatlands during the dry season (Sakabe et al., 2018). The CH_4 uptakes might be due to methanotrophy in the aerobic upper peat layer (Arai et al., 2014). However, CH_4 uptake by tree may also be possible (Sundqvist, Crill, Mölder, Vestin, & Lindroth, 2012).

4.4 | Potential effects of GWL on CH₄ production and oxidation

Variation in soil redox conditions driven by GWL fluctuation plays an essential role in influencing not only the quantity but also the quality of organic substrate used by the methanogenic archaea for CH₄ production (Girkin et al., 2018; Hoyos-Santillan et al., 2016; Reiche, Gleixner, & Küsel, 2010; Winton et al., 2017). Higher GWLs promote CH₄ production in a relatively large portion of the peat column and restrict the zone in which aerobic CH₄ oxidation can occur (Moore & Roulet, 1993; Moore et al., 2011; Strack et al., 2004). In contrast, lower GWL would narrow the zone of CH₄ production in the peat column and further supporting aerobic CH₄ oxidation above the GWL. In tropical peatlands, the availability of labile organic matter is largely limited to near-surface peat, for example, via root exudation and leaching from fresh litter (Brady, 1997; Könönen et al., 2016). When GWLs are low, most of this labile organic matter will be aerobically decomposed to CO₂ (Itoh, Okimoto, Hirano, & Kusin, 2017) and unavailable for CH₄ production. Therefore, when GWLs are low only organic matter with a greater aromatic content derived from the deeper peat would be available for anaerobic decomposition, restricting CH₄ production (Sakabe et al., 2018). In the *Acacia* plantation, most of the labile organic matter supplied from harvested vegetation residues (leaf litter, small branches, and roots) and boiler wood ash might be restricted above GWL in the surface peat layer and expected to be aerobically decomposed to CO₂ (Jauhiainen, Hooijer, & Page, 2012). Therefore, the effect (if any) of harvested vegetation residues and boiler wood ash on CH₄ production would be minor.

4.5 | Comparison of NEE-CH₄ with other studies

Our annual NEE-CH₄ over the natural forest are in the same range as those measured using the eddy covariance technique above a tropical peatland in the presence of CH₄-transporting trees (10.0–14.4 g m⁻² year⁻¹; Tang et al., 2018; Wong et al., 2018). In the absence of CH₄-transporting trees, a study in a tropical peatland reported no significant diurnal pattern in NEE-CH₄ (Sakabe et al., 2018) and far lower annual CH₄ emissions (0.12–0.23 g m⁻² year⁻¹), despite similar GWLs to our forest site. The chamber-based total ecosystem flux including tree CH₄ emissions to an average height of 15 m based on the power function relationship from a tropical peatland is lower than our results over the natural forest (Pangala et al., 2013). Despite the higher GWL as compared to our study, the lower emissions in Pangala et al. (2013) can be attributed to (a) a lower hollow to hummock area ratio (1:1), as CH₄ emissions from hollows can be up to 50 times higher as compared to hummocks (Pangala et al., 2013); and (b) possible underestimation of vegetation-mediated transport. Emissions from young trees exceed those of mature trees by orders of magnitudes (Pangala, Gowing, Hornibrook, & Gauci, 2014), but Pangala et al. (2013) reported on emissions from mature trees. Also shoots can emit up to 10 times more than stems

in a boreal forest (Machacova et al., 2016), but were not included in Pangala et al. (2013). Furthermore, entire trees may release CH₄, albeit at the lower rates from their higher portions.

Our annual NEE-CH₄ over the natural forest are around two times higher than the IPCC CH₄ emissions factor for rewetted tropical peatland, derived from undrained sites (Blain et al., 2014). This difference could be attributable to vegetation-mediated transport, which was not captured by most of the studies used to derive the IPCC CH₄ emission factor (Blain et al., 2014). Our annual NEE-CH₄ over the natural forest are nevertheless lower than those reported from Amazonian peatlands (Teh et al., 2017) and floodplain wetlands (Dalmagro et al., 2019; Pangala et al., 2017). In Amazonian peatlands, CH₄ production is greater owing to high nutrient status and soil pH, and low recalcitrant carbon (Wassmann et al., 1992). In addition, methanotrophy is generally less effective because of increased anoxic and stratified, water-submerged sediments (Bartlett et al., 1988; Devol & Rickey, 1990).

Our annual NEE-CH₄ over the natural forest is similar to emissions from northern bogs (average = 9.5 g m⁻² year⁻¹) and around two times lower than CH₄ emissions from northern fens (average = 20.5 g m⁻² year⁻¹; Abdalla et al., 2016). Higher temperatures in tropical peatlands favor greater humification, selective removal of reactive labile carbohydrates, and accumulation of aromatic content leading to a highly recalcitrant residual peat (Brady, 1997; Hodgkins et al., 2018). This results in low substrate availability for CH₄ production in the woody peat where there is a high aromatic lignin content (Miyajima, Wada, Hanba, & Vijarnsorn, 1997; Sakabe et al., 2018). In northern peatlands, peat is mainly derived from mosses, sedges, and herbs which contain a high carbohydrate and lower aromatic content (Hodgkins et al., 2018). This supports higher CH₄ production in northern peatlands, despite lower temperatures (Sundh, Nilsson, Granberg, & Svensson, 1994; Updegraff, Pastor, Bridgman, & Johnston, 1995). Tropical peatlands also typically have higher vertical and lateral recharge rates, driven by higher hydraulic conductivity than northern peatlands (Evans et al., 2014), making them susceptible to rapid flushing of the dissolved CH₄ after rainfall. This could limit CH₄ accumulation in near-surface porewaters, reducing the potential for diffusion, ebullition, and vegetation-mediated transport, but could increase emissions via drainage waters. In contrast, lower vertical and lateral recharge rates in northern peatlands support the buildup of dissolved CH₄ concentrations, and result in substantial ebullition and a high CH₄ concentration near the surface soil causing high diffusive and vegetation-mediated transport (Hoyt, 2017).

Our annual NEE-CH₄ at the *Acacia* plantation is around 18 times higher than the IPCC CH₄ soil-derived emission factor for this category, which is mostly based on soil CH₄ flux measurements (Dröslér et al., 2014). The IPCC methodology does, however, provide a separate emission factor for CH₄ emissions from drainage canals in tropical peatlands, of 225 g/m² ditch surface area year⁻¹. Based on 2.1% of the flux footprint area occupied by canals and ditches at the *Acacia* plantation, the water surfaces are contributed to generate an area-weighted emission of 4.74 g m⁻² year⁻¹, which is very similar to our ecosystem-scale

estimate over the *Acacia* plantation. The *Acacia* plantation in this study is situated within one of the largest tracts of *Acacia* plantation in Southeast Asia, managed by the same company in a similar way. The fraction of water surface in the majority of land uses involving drainage in tropical peatlands is approximately 2% (Drösler et al., 2014). Hence, our *Acacia* plantation with 2.1% of water surface can be considered as both representative and conservative in Southeast Asia. The results indicate that despite their small fractional area, higher emissions from water management ditches could be large enough to partly compensate for the reduced CH_4 emissions by lower GWL on the remainder of the managed peatland area. Notably, this suggests some potential for targeted mitigation measures to reduce CH_4 emissions, for example, by keeping the ditches clear and vegetation free (Jauhiainen & Silvennoinen, 2012; Waldron et al., 2019).

Regardless of land-cover on tropical peatland, if GWL is lower than 20 cm below the ground surface, most studies indicate that the peat acts as a net sink for atmospheric CH_4 (Couwenberg, Dommain, & Joosten, 2010; Hergoualc'h & Verchot, 2012; Ishikura et al., 2019). Our results show that even when GWL is lower than 20 cm below the peat surface, both ecosystems acted as a CH_4 source. Hence, it is possible for the ecosystem as a whole to act as a net CH_4 source to the atmosphere due to emissions from vegetation and water surfaces, despite the soil surface likely acting as a net CH_4 sink (Melling et al., 2005). Lower frequency and/or below-canopy measurements, for example, daytime chamber measurements of soil surface exchange, are likely to give highly erroneous estimates of CH_4 emissions in both ecosystems. In the case of the *Acacia* plantation, it is likely that such measurements would suggest that the system is a net sink for CH_4 , when in fact it remains a source (albeit smaller than the natural forest).

Overall, our results highlight that tropical peatlands, including natural forest and areas managed for forest plantation, are significant sources of CH_4 , and probably have a greater impact on global atmospheric CH_4 concentrations than previously thought. The associated radiative forcing effect of CH_4 emissions has the potential to partly offset net CO_2 uptake.

If we follow IPCC GWP accounting methodology and apply a 100 year GWP of 34 for CH_4 (Myhre et al., 2013), this implies a CH_4 emission of $3.1 \text{ t CO}_2\text{eq ha}^{-1} \text{ year}^{-1}$ from natural forest. Applying a long-term peat accumulated CO_2 rate of around $2.6 \text{ t CO}_2 \text{ ha}^{-1} \text{ year}^{-1}$ since their formation (Dommain et al., 2011), the 100 year net warming impact for tropical peatland would be $0.5 \text{ t CO}_2\text{eq ha}^{-1} \text{ year}^{-1}$. Over longer time-horizons, the shorter atmospheric lifetime of CH_4 compared to CO_2 means that an ecosystem that is in approximate greenhouse gas balance based on 100 year net warming impact will have a net cooling impact if it acts as a sustained CO_2 sink and a steady CH_4 source (Allen et al., 2018; Froking, Roulet, & Fuglestedt, 2006). However, according to the current IPCC assessment, tropical peatlands are in approximate CO_2 balance (Drösler et al., 2014); therefore, the net warming impact value would be $3.1 \text{ t CO}_2 \text{ ha}^{-1} \text{ year}^{-1}$. Nevertheless, our data confirm that CH_4 emissions from tropical peatlands should be included in landscape level

greenhouse gas budgets (Miettinen, Hooijer, Vernimmen, Liew, & Page, 2017; Wijedasa et al., 2018).

4.6 | Impact of *Acacia* plantation on CH_4 emissions

We present here an assessment of the impact of forest plantation on CH_4 emissions associated with the altered landscape (i.e., *Acacia* plantation). By definition, the impact represents the actual CH_4 exchange with the atmosphere in addition to the exchange that existed in the pre-existing natural landscape, and thus represents the exchange that can be directly attributed to the creation and existence of the *Acacia* plantation (Prairie et al., 2018; Teodoru et al., 2012).

Our measurements indicate that both studied ecosystems in the tropical peatland functioned as net CH_4 sources to the atmosphere (Table 4, Figure 7). Therefore, our results indicated that the impact of the *Acacia* plantation was to reduce CH_4 emissions by $4.4 \pm 1.7 \text{ g m}^{-2} \text{ year}^{-1}$ (Table 4, Figure 7). If we apply a 100 year GWP of 34 (Myhre et al., 2013), this implies an emission reduction of $1.5 \text{ t CO}_2\text{eq ha}^{-1} \text{ year}^{-1}$. For comparison, the IPCC's Tier 1 default emission factor for CO_2 from *Acacia* plantation on tropical peat is $73 \text{ t CO}_2 \text{ ha}^{-1} \text{ year}^{-1}$ (Drösler et al., 2014), which is larger than the natural forest. Measurements of net ecosystem CO_2 exchanges over the natural forest and the *Acacia* plantation are being conducted (C. Deshmukh, unpublished data); results of this ongoing study will be published in due course, following the completion of one 5 year *Acacia* plantation cycle, and will also take into consideration the biomass harvested from the plantation. These measurements will lead to a better understanding of the climate footprint of *Acacia* plantation (Dommain et al., 2018; Petrescu et al., 2015).

The estimated impact of *Acacia* plantation on CH_4 exchange related to land-cover change that we present here is by no means invariant in time and space. In addition to variations related to natural hydrology, the impact is also likely to vary with actual water

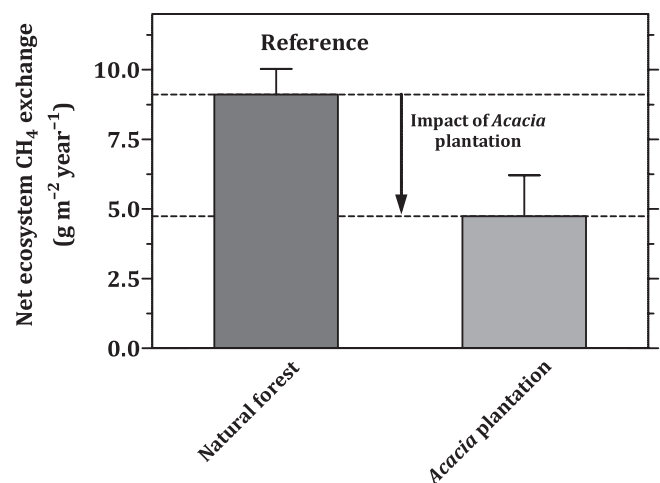


FIGURE 7 Impact of the *Acacia* plantation on net ecosystem CH_4 exchange from tropical peatland

management practices in plantation landscapes. Furthermore, results presented here are specific for *Acacia* plantation; thus, caution should be taken when extrapolating to other agriculture in the region (e.g., sago, oil palm, rubber plantations, etc.) with different water management practices and fertilizer applications (Hergoualc'h & Verchot, 2012). To evaluate the impact of land-cover change on global peatland CH₄ emissions, more ecosystem-scale flux measurement studies are needed.

In conclusion, our half-hourly multi-year NEE-CH₄ measurements directly captured and integrated “hot spot and hot moment” dynamics of all known and unknown sources and removals in the studied ecosystems. The observed high variability in NEE-CH₄ suggests complex nonlinear process-level controls on CH₄ exchange between tropical peatlands and the atmosphere. Our results provide some of the first reliable information on the magnitudes of CH₄ exchange at a tropical peatland ecosystem scale, demonstrating that traditional manual soil chamber techniques provide an incomplete picture of the total CH₄ flux, and improving mechanistic understanding based on high temporal resolution measurements of NEE-CH₄ and key environmental variables such as the sensitivity of emissions to GWL. Our data indicate that the *Acacia* plantation on tropical peatland results in significant reductions in CH₄ emissions compared to the natural system, although the associated cooling impact is likely to be smaller than the accompanying warming impact of higher CO₂ and nitrous oxide emissions. More ecosystem-scale measurements are needed to fully evaluate the effect of land-cover change on the greenhouse gas balance, at a larger number of sites and over long time periods, in order to develop science-based, climate-smart management practices for tropical peatlands.

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DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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REFERENCES

- Abdalla, M., Hastings, A., Truu, J., Espenberg, M., Mander, Ü., & Smith, P. (2016). Emissions of methane from northern peatlands: A review of management impacts and implications for future management options. *Ecology and Evolution*, 6(19), 7080–7102. <https://doi.org/10.1002/ece3.2469>
- Alberto, M. C. R., Wassmann, R., Buresh, R. J., Quilty, J. R., Correa, T. Q., Sandro, J. M., & Centeno, C. A. R. (2014). Measuring methane flux from irrigated rice fields by eddy covariance method using open-path gas analyzer. *Field Crops Research*, 160, 12–21. <https://doi.org/10.1016/j.fcr.2014.02.008>
- Allen, M. R., Shine, K. P., Fuglestedt, J. S., Millar, R. J., Cain, M., Frame, D. J., & Macey, A. H. (2018). A solution to the misrepresentations of CO₂-equivalent emissions of short-lived climate pollutants under ambitious mitigation. *npj Climate and Atmospheric Science*, 1, 16. <https://doi.org/10.1038/s41612-018-0026-8>
- Arai, H., Hadi, A., Darung, U., Limin, S. H., Hatano, R., & Inubushi, K. (2014). A methanotrophic community in a tropical peatland is unaffected by drainage and forest fires in a tropical peat soil. *Soil Science and Plant Nutrition*, 60(4), 577–585. <https://doi.org/10.1080/00380768.2014.922034>
- Aubinet, M., Grelle, A., Ibrom, A., Rannik, U., Moncrieff, J., Foken, T., ... Vesala, T. (2000). Estimates of the annual net carbon and water exchange of European forests: The EUROFLUX methodology. *Advances in Ecological Research*, 30, 113–175. [https://doi.org/10.1016/S0065-2504\(08\)60018-5](https://doi.org/10.1016/S0065-2504(08)60018-5)
- Barba, J., Bradford, M. A., Brewer, P. E., Bruhn, D., Covey, K., Haren, J., ... Vargas, R. (2018). Methane emissions from tree stems: A new frontier in the global carbon cycle. *New Phytologist*, 222(1), 18–28. <https://doi.org/10.1111/nph.15582>
- Bartlett, K. B., Crill, P. M., Sebach, D. I., Harriss, R. C., Wilson, J. O., & Melack, J. M. (1988). Methane flux from the central Amazonian floodplain. *Journal of Geophysical Research*, 93(D2), 1571–1582. <https://doi.org/10.1029/JD093iD02p01571>
- Blain, D., Murdiyarso, D., Couwenberg, J., Nagata, O., Renou-Wilson, F., Sirin, A., ... Wilson, D. (2014). Rewetted organic soils. In T. Hiraishi, T. Krug, K. Tanabe, N. Srivastava, J. Baasansuren, M. Fukuda, & T. G. Troxler (Eds.), *2013 Supplement to the 2006 IPCC guidelines for national greenhouse gas inventories: Wetlands* (pp. 3.1–3.42). Geneva, Switzerland: IPCC.
- Brady, M. A. (1997). Effect of vegetation changes on organic matter dynamics in three coastal peat deposits in Sumatra, Indonesia. In J. O. Rieley & S. E. Page (Eds.), *Biodiversity and sustainability of tropical peatlands* (pp. 113–134). Cardigan, UK: Samara Publishing.
- Brix, H., Sorrell, B. K., & Orr, P. T. (1992). Internal pressurization and convective gas flow in some emergent freshwater macrophytes. *Limnology and Oceanography*, 37(7), 1420–1433. <https://doi.org/10.4319/lo.1992.37.7.1420>
- Burba, G., Anderson, T., & Komissarov, A. (2019). Accounting for spectroscopic effects in laser-based open-path eddy covariance flux measurements. *Global Change Biology*, 25(6), 2189–2202. <https://doi.org/10.1111/gcb.14614>
- Chanton, J. P., Bauer, J. E., Glaser, P. A., Siegel, D. I., Kelley, C. A., Tyler, S. C., ... Lazrus, A. (1995). Radiocarbon evidence for the substrates supporting methane formation within northern Minnesota peatlands. *Geochimica et Cosmochimica Acta*, 59(17), 3663–3668. [https://doi.org/10.1016/0016-7037\(95\)00240-Z](https://doi.org/10.1016/0016-7037(95)00240-Z)
- Chanton, J. P., Martens, C. S., Kelley, C. A., Crill, P. M., & Showers, W. J. (1992). Methane transport mechanisms and isotopic fractionation in emergent macrophytes of an Alaskan tundra lake. *Journal of Geophysical Research-Atmospheres*, 97(D15), 16681–16688. <https://doi.org/10.1029/90JD01542>
- Chanton, J. P., Whiting, G. J., Happell, J. D., & Gerard, G. (1993). Contrasting rates and diurnal patterns of methane emission from emergent aquatic macrophytes. *Aquatic Botany*, 46(2), 111–128. [https://doi.org/10.1016/0304-3770\(93\)90040-4](https://doi.org/10.1016/0304-3770(93)90040-4)
- Christensen, T. R., Ekberg, A., Ström, L., Mastepanov, M., Panikov, N., Öquist, M., ... Oskarsson, H. (2003). Factors controlling large scale variations in methane emissions from wetlands. *Geophysical Research Letters*, 30(7), 1414–1419. <https://doi.org/10.1029/2002GL016848>

- Chu, H., Chen, J., Gottgens, J. F., Ouyang, Z., John, R., Czajkowski, K., & Becker, R. (2014). Net ecosystem methane and carbon dioxide exchanges in a Lake Erie coastal marsh and a nearby cropland. *Journal of Geophysical Research: Biogeosciences*, 119(5), 722–740. <https://doi.org/10.1002/2013JG002520>
- Cobb, A. R., Hoyt, A. M., Gandois, L., Eri, J., Dommain, R., Abu Salim, K., ... Harvey, C. F. (2017). How temporal patterns in rainfall determine the geomorphology and carbon fluxes of tropical peatlands. *Proceedings of the National Academy of Sciences of the United States of America*, 114(26), E5187–E5196. <https://doi.org/10.1073/pnas.1701090114>
- Collins, W. J., Webber, C. P., Cox, P. M., Huntingford, C., Lowe, J., Sitch, S., ... Powell, T. (2018). Increased importance of methane reduction for a 1.5 degree target. *Environmental Research Letters*, 13, 54003. <https://doi.org/10.1088/1748-9326/aab89c>
- Couwenberg, J., Dommain, R., & Joosten, H. (2010). Greenhouse gas fluxes from tropical peatlands in south-east Asia. *Global Change Biology*, 16(6), 1715–1732. <https://doi.org/10.1111/j.1365-2486.2009.02016.x>
- Covey, K. R., & Megonigal, J. P. (2019). Methane production and emissions in trees and forests. *New Phytologist*, 222(1), 35–51. <https://doi.org/10.1111/nph.15624>
- Dacey, J. W. H. (1981). Pressurized ventilation in the yellow waterlily. *Ecology*, 62(5), 1137–1147. <https://doi.org/10.2307/1937277>
- Dalmagro, H. J., Arruda, P. H., Vourlitis, G. L., Lathuillière, M. J., Nogueira, J. D., Couto, E. G., & Johnson, M. S. (2019). Radiative forcing of methane fluxes offsets net carbon dioxide uptake for a tropical flooded forest. *Global Change Biology*, 25(6), 1967–1981. <https://doi.org/10.1111/gcb.14615>
- Dengel, S., Levy, P. E., Grace, J., Jones, S. K., & Skiba, U. M. (2011). Methane emissions from sheep pasture measured with an open-path eddy covariance system. *Global Change Biology*, 17(12), 3524–3533. <https://doi.org/10.1111/j.1365-2486.2011.02466.x>
- Desai, A. R., Xu, K., Tian, H., Weishampel, P., Thom, J., Baumann, D., & Kolka, R. (2015). Landscape-level terrestrial methane flux observed from a very tall tower. *Agricultural and Forest Meteorology*, 201, 61–75. <https://doi.org/10.1016/j.agrformet.2014.10.017>
- Deshmukh, C., Serça, D., Delon, C., Tardif, R., Demarty, M., Jarnot, C., ... Guérin, F. (2014). Physical controls on CH₄ emissions from a newly flooded subtropical freshwater hydroelectric reservoir: Nam Theun 2. *Biogeosciences*, 11, 4251–4269. <https://doi.org/10.5194/bg-11-4251-2014>
- Deventer, M. J., Griffis, T. J., Roman, D. T., Kolka, R. K., Wood, J. D., Erickson, M., ... Millet, D. B. (2019). Error characterization of methane fluxes and budgets derived from a long-term comparison of open- and closed-path eddy covariance systems. *Agricultural and Forest Meteorology*, 278, 107638. <https://doi.org/10.1016/j.agrformet.2019.107638>
- Devol, A. H., Richey, J. E., Forsberg, B. R., & Martinelli, L. A. (1990). Seasonal dynamics in methane emissions from the Amazon River floodplain to the troposphere. *Journal of Geophysical Research*, 95(D10), 16417–16426. <https://doi.org/10.1029/JD095iD10p16417>
- Dlugokencky, E. (2019). NOAA/ESRL. Retrieved from www.esrl.noaa.gov/gmd/ccgg/trends_ch4/
- Dommain, R., Couwenberg, J., & Joosten, H. (2011). Development and carbon sequestration of tropical peat domes in south-east Asia: Links to post-glacial sea-level changes and Holocene climate variability. *Quaternary Science Reviews*, 30(7–8), 999–1010. <https://doi.org/10.1016/j.quascirev.2011.01.018>
- Dommain, R., Frolking, S., Jeltsch-Thömmes, A., Joos, F., Couwenberg, J., & Glaser, P. H. (2018). A radiative forcing analysis of tropical peatlands before and after their conversion to agricultural plantations. *Global Change Biology*, 24(11), 5518–5533. <https://doi.org/10.1111/gcb.14400>
- Dröslér, M., Verchot, L. V., Freibauer, A., Pan, F., Evans, C. D., Bourbonniere, R. A., ... Wang, C. (2014). Drained inland organic soils. In T. Hiraishi, T. Krug, K. Tanabe, N. Srivastava, J. Baasansuren, M. Fukuda, & T. G. Troxler (Eds.), *2013 Supplement to the 2006 IPCC guidelines for national greenhouse gas inventories: Wetlands* (pp. 2.1–2.79). Geneva, Switzerland: IPCC.
- Estop-Aragones, C., Knorr, K. H., & Blodau, C. (2012). Controls on in situ oxygen and dissolved inorganic carbon dynamics in peats of a temperate fen. *Journal of Geophysical Research-Biogeosciences*, 117, G02002. <https://doi.org/10.1029/2011JG001888>
- Evans, C. D., Page, S. E., Jones, T., Moore, S., Gauci, V., Laiho, R., ... Garnett, M. H. (2014). Contrasting vulnerability of drained tropical and high-latitude peatlands to fluvial loss of stored carbon. *Global Biogeochemical Cycles*, 28(11), 1215–1234. <https://doi.org/10.1002/2013GB004782>
- Evans, C. D., Renou-Wilson, F., & Strack, M. (2016). The role of water-borne carbon in the greenhouse gas balance of drained and re-wetted peatlands. *Aquatic Sciences*, 78, 573–590. <https://doi.org/10.1007/s00027-015-0447-y>
- Evans, C. D., Williamson, J. M., Kacaribu, F., Irawan, D., Suardiwerianto, Y., Hidayat, M. F., ... Page, S. E. (2019). Rates and spatial variability of peat subsidence in Acacia plantation and forest landscapes in Sumatra, Indonesia. *Geoderma*, 338, 410–421. <https://doi.org/10.1016/j.geoderma.2018.12.028>
- Falge, E., Baldocchi, D., Olson, R., Anthoni, P., Aubinet, M., Bernhofer, C., ... Wofsy, S. (2001). Gap filling strategies for defensible annual sums of net ecosystem exchange. *Agricultural and Forest Meteorology*, 107(1), 43–69. [https://doi.org/10.1016/S0168-1923\(00\)00225-2](https://doi.org/10.1016/S0168-1923(00)00225-2)
- Finkelstein, P. L., & Sims, P. F. (2001). Sampling error in eddy correlation flux measurements. *Journal of Geophysical Research*, 106(D4), 3503–3509. <https://doi.org/10.1029/2000JD900731>
- Finnigan, J. (2006). The storage term in eddy flux calculation. *Agricultural and Forest Meteorology*, 136(3–4), 108–113. <https://doi.org/10.1016/j.agrformet.2004.12.010>
- Frolking, S., Roulet, N., & Fuglestad, J. (2006). How northern peatlands influence the Earth's radiative budget: Sustained methane emission versus sustained carbon sequestration. *Journal of Geophysical Research*, 111, G01008. <https://doi.org/10.1029/2005JG000091>
- Furukawa, Y., Inubushi, K., Ali, M., Itang, A. M., & Tsuruta, H. (2005). Effect of changing water table depths caused by land-use changes on greenhouse gas fluxes from tropical peat lands. *Nutrient Cycling in Agroecosystems*, 71(1), 81–91. <https://doi.org/10.1007/s10705-004-5286-5>
- Garnett, K. N., Megonigal, J. P., Litchfield, C., & Taylor, G. E. (2005). Physiological control of leaf methane emission from wetland plants. *Aquatic Botany*, 81(2), 141–155. <https://doi.org/10.1016/j.aquabot.2004.10.003>
- Ge, F., Zhu, S., Peng, T., Zhao, Y., Sielmann, F., Zhi, X., ... Ji, L. (2019). Risks of precipitation extremes over Southeast Asia: Does 1.5 °C or 2 °C global warming make a difference? *Environmental Research Letters*, 14, 044015. <https://doi.org/10.1088/1748-9326/aaff7e>
- Gedney, N., Huntingford, C., Comyn-Platt, E., & Wiltshire, A. (2019). Significant feedbacks of wetland methane release on climate change and the causes of their uncertainty. *Environmental Research Letters*, 14, 084027. <https://doi.org/10.1088/1748-9326/ab2726>
- Girkin, N. T., Vane, C. H., Cooper, H. V., Moss-Hayes, V., Craigon, J., Turner, B. L., ... Sjögersten, S. (2018). Spatial variability of organic matter properties determines methane fluxes in a tropical forested peatland. *Biogeochemistry*, 142(2), 231–245. <https://doi.org/10.1007/s10533-018-0531-1>
- Gumbrecht, T., Roman-Cuesta, R. M., Verchot, L., Herold, M., Wittmann, F., Householder, E., ... Murdiyarso, D. (2017). An expert system model for mapping tropical wetlands and peatlands reveals South America as the largest contributor. *Global Change Biology*, 23(9), 3581–3599. <https://doi.org/10.1111/gcb.13689>
- Hadi, A., Inubushi, K., Furukawa, Y., Purnomo, E., Rasmadi, M., & Tsuruta, H. (2005). Greenhouse gas emissions from tropical peatlands of Kalimantan Indonesia. *Nutrient Cycling in Agroecosystems*, 71, 73–80. <https://doi.org/10.1007/s10705-004-0380-2>

- Hergoualc'h, K. A., & Verchot, L. V. (2012). Changes in soil CH₄ fluxes from the conversion of tropical peat swamp forests: A meta-analysis. *Journal of Integrative Environmental Sciences*, 9(2), 93–101. <https://doi.org/10.1080/1943815X.2012.679282>
- Hodgkins, S. B., Richardson, C. J., Dommair, R., Wang, H., Glaser, P. H., Verbeke, B., ... Chanton, J. P. (2018). Tropical peatland carbon storage linked to global latitudinal trends in peat recalcitrance. *Nature Communications*, 9(1), 3640. <https://doi.org/10.1038/s41467-018-06050-2>
- Hoyos-Santillan, J., Lomax, B. H., Large, D., Turner, B. L., Boom, A., Lopez, O. R., & Sjögersten, S. (2016). Quality not quantity: Organic matter composition controls of CO₂ and CH₄ fluxes in neotropical peat profiles. *Soil Biology and Biochemistry*, 103, 86–96. <https://doi.org/10.1016/j.soilbio.2016.08.017>
- Hoyt, A. M. (2017). *Carbon fluxes from tropical peatlands: methane carbon dioxide and peatland subsidence*. Doctoral dissertation, Massachusetts Institute of Technology. Retrieved from <https://dspace.mit.edu/handle/1721.1/113476>
- Inubushi, K., Hadi, A., Okazaki, M., & Yonebayashi, K. (1998). Effect of converting wetland forest to sago palm plantations on methane gas flux and organic carbon dynamics in tropical peat soil. *Hydrological Processes*, 12(13–14), 2073–2080. [https://doi.org/10.1002/\(SICI\)1099-1085\(19981030\)12:13/14<2073:AID-HYP720>3.0.CO;2-K](https://doi.org/10.1002/(SICI)1099-1085(19981030)12:13/14<2073:AID-HYP720>3.0.CO;2-K)
- Ishikura, K., Hirata, R., Hirano, T., Okimoto, Y., Wong, G. X., Melling, L., ... Ishii, Y. (2019). Carbon dioxide and methane emissions from peat soil in an undrained tropical peat swamp forest. *Ecosystems*, 22, 1852–1868. <https://doi.org/10.1007/s10021-019-00376-8>
- Itoh, M., Okimoto, Y., Hirano, T., & Kusin, K. (2017). Factors affecting oxidative peat decomposition due to land-use in tropical peat swamp forests in Indonesia. *Science of the Total Environment*, 609, 906–915. <https://doi.org/10.1016/j.scitotenv.2017.07.132>
- Jauhiainen, J., Hooijer, A., & Page, S. E. (2012). Carbon dioxide emissions from an Acacia plantation on peatland in Sumatra, Indonesia. *Biogeosciences*, 9, 617–630. <https://doi.org/10.5194/bg-9-617-2012>
- Jauhiainen, J., Limin, S., Silvennoinen, H., & Vasander, H. (2008). Carbon dioxide and methane fluxes in drained tropical peat before and after hydrological restoration. *Ecology*, 89(12), 3503–3514. <https://doi.org/10.1890/07-2038.1>
- Jauhiainen, J., & Silvennoinen, H. (2012). Diffusion GHG fluxes at tropical peatland drainage canal water surfaces. *Suo*, 63(3–4), 93–105.
- Jeeva, D., Bignell, D. E., Eggleton, P., & Maryati, M. (1999). Respiratory gas exchanges of termites from the Sabah (Borneo) assemblage. *Physiological Entomology*, 24(1), 11–17. <https://doi.org/10.1046/j.1365-3032.1999.00106.x>
- Keppler, F., Hamilton, J. T. G., Brass, M., & Röckmann, T. (2006). Methane emissions from terrestrial plants under aerobic conditions. *Nature*, 439, 187–191. <https://doi.org/10.1038/nature04420>
- Kim, J., Verma, S. B., Billesbach, D. P., & Clement, R. J. (1998). Diel variation in methane emission from a midlatitude prairie wetland: Significance of convective throughflow in *Phragmites australis*. *Journal of Geophysical Research*, 103(D21), 28029–28039. <https://doi.org/10.1029/98JD02441>
- Kirschke, S., Bousquet, P., Ciais, P., Saunio, M., Canadell, J. G., Dlugokencky, E. J., & Zeng, G. (2013). Three decades of global methane sources and removals. *Nature Geoscience*, 6, 813–823. <https://doi.org/10.1038/ngeo1955>
- Kljun, N., Calanca, P., Rotach, M. W., & Schmid, H. P. (2015). A simple two-dimensional parameterisation for Flux Footprint Prediction (FFP). *Geoscientific Model Development*, 8, 3695–3713. <https://doi.org/10.5194/gmd-8-3695-2015>
- Könönen, M., Jauhiainen, J., Laiho, R., Spetz, P., Kusin, K., Limin, S., & Vasander, H. (2016). Land-use increases the recalcitrance of tropical peat. *Wetlands Ecology Management*, 24(6), 717–731. <https://doi.org/10.1007/s11273-016-9498-7>
- Long, K. D., Flanagan, L. B., & Cai, T. (2010). Diurnal and seasonal variation in methane emissions in a northern Canadian peatland measured by eddy covariance. *Global Change Biology*, 16(9), 2420–2435. <https://doi.org/10.1111/j.1365-2486.2009.02083.x>
- Machacova, K., Bäck, J., Vanhatalo, A., Halmeenmäki, E., Kolari, P., Mammarella, I., ... Pihlatie, M. (2016). *Pinus sylvestris* as a missing source of nitrous oxide and methane in boreal forest. *Scientific Reports*, 6, 23410. <https://doi.org/10.1038/srep23410>
- Manning, F. C., Kho, L. K., Hill, T. C., Cornulier, T., & Teh, Y. A. (2019). Carbon emissions from oil palm plantations on peat soil. *Frontiers in Forests and Global Change*, 2(37). <https://doi.org/10.3389/ffgc.2019.00037>
- Massman, W. J. (2000). A simple method for estimating frequency response corrections for eddy covariance systems. *Agriculture and Forest Meteorology*, 104(3), 185–198. [https://doi.org/10.1016/S0168-1923\(00\)00164-7](https://doi.org/10.1016/S0168-1923(00)00164-7)
- Massman, W. J. (2001). Reply to comment by Rannik on "A simple method for estimating frequency response corrections for eddy covariance systems". *Agricultural and Forest Meteorology*, 107, 247–251. [https://doi.org/10.1016/S0168-1923\(00\)00237-9](https://doi.org/10.1016/S0168-1923(00)00237-9)
- Mauder, M., Cuntz, M., Drüe, C., Graf, A., Rebmann, C., Schmid, H. P., ... Steinbrecher, R. (2013). A strategy for quality and uncertainty assessment of long-term eddy-covariance measurements. *Agricultural and Forest Meteorology*, 169, 122–135. <https://doi.org/10.1016/j.agrformet.2012.09.006>
- McDermitt, D., Burba, G., Xu, L., Anderson, T., Komissarov, A., Riensche, B., ... Hastings, S. (2011). A new low-power, open-path instrument for measuring methane flux by eddy covariance. *Applied Physics B*, 102(2), 391–405. <https://doi.org/10.1007/s00340-010-4307-0>
- Megonigal, J. P., Brewer, P. E., & Knee, K. L. (2019). Radon as a natural tracer of gas transport through trees. *New Phytologist*, 225(4), 1470–1475. <https://doi.org/10.1111/nph.16292>
- Melling, L., Hatano, R., & Goh, K. J. (2005). Methane fluxes from three ecosystems in tropical peatland of Sarawak, Malaysia. *Soil Biology and Biochemistry*, 37(8), 1445–1453. <https://doi.org/10.1016/j.soilbio.2005.01.001>
- Miettinen, J., Hooijer, A., Vernimmen, R., Liew, S. C., & Page, S. E. (2017). From carbon sink to carbon source: Extensive peat oxidation in insular Southeast Asia since 1990. *Environmental Research Letters*, 12, 024014. <https://doi.org/10.1088/1748-9326/aa5b6f>
- Miettinen, J., Shi, C., & Liew, S. C. (2016). Land cover distribution in the peatlands of Peninsular Malaysia Sumatra and Borneo in 2015 with changes since 1990. *Global Ecology and Conservation*, 6, 67–78. <https://doi.org/10.1016/j.gecco.2016.02.004>
- Miyajima, T., Wada, E., Hanba, Y., & Vijarnsorn, P. (1997). Anaerobic mineralization of indigenous organic matters and CH₄ production in tropical wetland soils. *Geochimica et Cosmochimica Acta*, 61(17), 3739–3751. [https://doi.org/10.1016/S0016-7037\(97\)00189-0](https://doi.org/10.1016/S0016-7037(97)00189-0)
- Moncrieff, J., Clement, R., Finnigan, J. J., & Meyers, T. (2004). Averaging, detrending, and filtering of eddy covariance time series. In B. E. Law, X. Lee & W. J. Massman (Eds.), *Handbook of micrometeorology: A guide for surface flux measurement and analysis* (pp. 7–31). Dordrecht, The Netherlands: Kluwer Academic. Retrieved from <http://hdl.handle.net/102.100.100/189360?index=1>
- Moore, T. R., & Roulet, N. T. (1993). Methane flux: Water table relations in northern wetlands. *Geophysical Research Letters*, 20(7), 587–590. <https://doi.org/10.1029/93GL00208>
- Moore, T. R., Young, A., Bubier, J. L., Humphreys, E. R., Lafleur, P. M., & Roulet, N. T. (2011). A multi-year record of methane flux at the Mer Bleue bog, southern Canada. *Ecosystems*, 14(4), 646–657. <https://doi.org/10.1007/s10021-011-9435-9>
- Myhre, G., Shindell, D., Brèon, F. M., Collins, W., Fuglestedt, J., Huang, J., ... Zhang, H. (2013). Anthropogenic and natural radiative forcing. In T. F. Stocker, D. Qin, G. K. Plattner, M. Tignor, S. K. Allen, J. Boschung, ... P. M. Midgley (Eds.), *Climate change 2013: The physical science basis. Contribution of working group I to the fifth assessment report of the*

- Intergovernmental Panel on Climate Change (pp. 659–740). Cambridge, UK and New York, NY: Cambridge University Press. <https://doi.org/10.1017/CBO9781107415324.018>
- Nisbet, E. G., Manning, M. R., Dlugokencky, E. J., Fisher, R. E., Lowry, D., Michel, S. E., ... White, J. W. C. (2019). Very strong atmospheric methane growth in the 4 years 2014–2017: Implications for the Paris Agreement. *Global Biogeochemical Cycles*, 33(3), 318–342. <https://doi.org/10.1029/2018GB006009>
- Nisbet, R., Fisher, R., Nimmo, R. H., Bendall, D. S., Crill, P. M., Gallego-Sala, A. V., ... Nisbet, E. G. (2009). Emission of methane from plants. *Proceedings Biological Sciences*, 276, 1347–1354. <https://doi.org/10.1098/rspb.2008.1731>
- Page, S. E., Rieley, J. O., & Banks, C. J. (2011). Global and regional importance of the tropical peatland carbon pool. *Global Change Biology*, 17, 798–818. <https://doi.org/10.1111/j.1365-2486.2010.02279.x>
- Pangala, S. R., Enrich-Prast, A., Basso, L. S., Peixoto, R. B., Bastviken, D., Hornibrook, E. R. C., ... Gaudi, V. (2017). Large emissions from floodplain trees close the Amazon methane budget. *Nature*, 552, 230–234. <https://doi.org/10.1038/nature24639>
- Pangala, S. R., Gowing, D. J., Hornibrook, E. R. C., & Gaudi, V. (2014). Controls on methane emissions from *Alnus glutinosa* samplings. *New Phytologist*, 201(3), 887–896. <https://doi.org/10.1111/nph.12561>
- Pangala, S. R., Hornibrook, E. R., Gowing, D. J., & Gaudi, V. (2015). The contribution of trees to ecosystem methane emissions in a temperate forested wetland. *Global Change Biology*, 21(7), 2642–2654. <https://doi.org/10.1111/gcb.12891>
- Pangala, S. R., Moore, S., Hornibrook, E. R., & Gaudi, V. (2013). Trees are major conduits for methane egress from tropical forested wetlands. *New Phytologist*, 197(2), 524–531. <https://doi.org/10.1111/nph.12031>
- Papale, D., Reichstein, M., Aubinet, M., Canfora, E., Bernhofer, C., Kutsch, W., ... Yakir, D. (2006). Towards a standardized processing of net ecosystem exchange measured with eddy covariance technique: Algorithms and uncertainty estimation. *Biogeosciences*, 3, 571–583. <https://doi.org/10.5194/bg-3-571-2006>
- Parker, R. J., Boesch, H., McNorton, J., Comyn-Platt, E., Gloor, M., Wilson, C., ... Bloom, A. A. (2018). Evaluating year-to-year anomalies in tropical wetland methane emissions using satellite CH₄ observations. *Remote Sensing of Environment*, 211, 261–275. <https://doi.org/10.1016/j.rse.2018.02.011>
- Petrescu, A. M. R., Lohila, A., Tuovinen, J.-P., Baldocchi, D. D., Desai, A. R., Roulet, N. T., ... Cescatti, A. (2015). The uncertain climate footprint of wetlands under human pressure. *Proceedings of the National Academy of Sciences of the United States of America*, 112(15), 4594–4599. <https://doi.org/10.1073/pnas.1416267112>
- Pitz, S., & Megonigal, J. P. (2017). Temperate forest methane removal diminished by tree emissions. *New Phytologist*, 214(4), 1432–1439. <https://doi.org/10.1111/nph.14559>
- Poulter, B., Bousquet, P., Canadell, J. G., Ciais, P., Peregon, A., Saunio, M., ... Zhu, Q. (2017). Global wetland contribution to 2000–2012 atmospheric methane growth rate dynamics. *Environmental Research Letters*, 12(9), 094013. <https://doi.org/10.1088/1748-9326/aa8391>
- Prairie, Y. T., Alm, J., Beaulieu, J., Barros, N., Battin, T., Cole, J., ... Vachon, D. (2018). Greenhouse gas emissions from freshwater reservoirs: What does the atmosphere see? *Ecosystems*, 21, 1058–1071. <https://doi.org/10.1007/s10021-017-0198-9>
- Reiche, M., Gleixner, G., & Küsel, K. (2010). Effect of peat quality on microbial greenhouse gas formation in an acidic fen. *Biogeosciences*, 7, 187–198. <https://doi.org/10.5194/bg-7-187-2010>
- Rice, A. L., Butenhoff, C. L., Shearer, M. J., Teama, D., Rosenstiel, T. N., & Khalil, M. A. K. (2010). Emissions of anaerobically produced methane by trees. *Geophysical Research Letters*, 37, L03807. <https://doi.org/10.1029/2009GL041565>
- Rinne, J., Riutta, T., Pihlatie, M., Aurela, M., Haapanala, S., Tuovinen, J.-P., ... Vesala, T. (2007). Annual cycle of methane emission from a boreal fen measured by the eddy covariance technique. *Tellus*, 59(3), 449–457. <https://doi.org/10.1111/j.1600-0889.2007.00261.x>
- Sakabe, A., Itoh, M., Hirano, T., & Kusin, K. (2018). Ecosystem-scale methane flux in tropical peat swamp forest in Indonesia. *Global Change Biology*, 24, 5123–5136. <https://doi.org/10.1111/gcb.14410>
- Saunio, M., Bousquet, P., Poulter, B., Peregon, A., Ciais, P., Canadell, J. G., ... Zhu, Q. (2016). The global methane budget 2000–2012. *Earth System Science Data*, 8, 697–751. <https://doi.org/10.5194/essd-8-697-2016>
- Sjögersten, S., Aplin, P., Gaudi, V., Peacock, M., Siegenthaler, A., & Turner, B. L. (2018). Temperature response of ex-situ greenhouse gas emissions from tropical peatlands: Interactions between forest type and peat moisture conditions. *Geoderma*, 324, 47–55. <https://doi.org/10.1016/j.geoderma.2018.02.029>
- Strack, M., Waddington, J. M., & Tuittila, E. (2004). Effect of water table drawdown on northern peatland methane dynamics: Implications for climate change. *Global Biogeochemical Cycles*, 18(4), GB4003. <https://doi.org/10.1029/2003GB002209>
- Sulistiyanto, Y. (2004). *Nutrient dynamics in different sub-types of peat swamp forest in Central Kalimantan, Indonesia*. Doctoral dissertation, University of Nottingham. Retrieved from <http://eprints.nottingham.ac.uk/12597/>
- Sundh, I., Nilsson, M., Granberg, G., & Svensson, B. H. (1994). Depth distribution of microbial production and oxidation of methane in northern peatlands. *Microbial Ecology*, 27, 253–265. <https://doi.org/10.1007/BF00182409>
- Sundqvist, E., Crill, P., Mölder, M., Vestin, P., & Lindroth, A. (2012). Atmospheric methane removal by boreal plants. *Geophysical Research Letters*, 39, <https://doi.org/10.1029/2012GL053592>
- Tagesson, T., Mölder, M., Mastepanov, M., Sigsgaard, C., Tamstorf, M. P., Lund, M., ... Ström, L. (2012). Land-atmosphere exchange of methane from soil thawing to soil freezing in a high-Arctic wet tundra ecosystem. *Global Change Biology*, 18, 1928–1940. <https://doi.org/10.1111/j.1365-2486.2012.02647.x>
- Tang, A. C. I., Stoy, P. C., Hirata, R., Musin, K. K., Aeries, E. B., Wenceslaus, J., & Melling, L. (2018). Eddy covariance measurements of methane flux at a tropical peat forest in Sarawak, Malaysian Borneo. *Geophysical Research Letters*, 45, 4390–4399. <https://doi.org/10.1029/2017GL076457>
- Teh, Y. A., Murphy, W. A., Berrio, J. C., Boom, A., & Page, S. E. (2017). Seasonal variability in methane and nitrous oxide fluxes from tropical peatlands in the western Amazon basin. *Biogeosciences*, 14, 3669–3683. <https://doi.org/10.5194/bg-14-3669-2017>
- Teodoru, C. R., Bastien, J., Bonneville, M.-C., del Giorgio, P. A., Demarty, M., Garneau, M., ... Tremblay, A. (2012). The net carbon footprint of a newly created boreal hydroelectric reservoir. *Global Biogeochemical Cycles*, 26, GB2016. <https://doi.org/10.1029/2011GB004187>
- Updegraff, K., Pastor, J., Bridgman, S. D., & Johnston, C. A. (1995). Environmental and substrate controls over carbon and nitrogen mineralization in northern wetlands. *Ecological Applications*, 5, 151–163. <https://doi.org/10.2307/1942060>
- van der Nat, F. J. W. A., Middelburg, J. J., van Meteren, D., & Wielemakers, A. (1998). Diel methane emission patterns from *Scirpus lacustris* and *Phragmites australis*. *Biogeochemistry*, 41, 1–22. <https://doi.org/10.1023/A:1005933100905>
- Vickers, D., & Mahrt, L. (1997). Quality control and flux sampling problems for tower and aircraft data. *Journal of Atmospheric and Oceanic Technology*, 14, 512–526. [https://doi.org/10.1175/1520-0426\(1997\)014%3c0512:QCAFSP%3e2.0.CO;2](https://doi.org/10.1175/1520-0426(1997)014%3c0512:QCAFSP%3e2.0.CO;2)
- Waddington, J. M., Roulet, N. T., & Swanson, R. V. (1996). Water table control of CH₄ emission enhancement by vascular plants in boreal peatlands. *Journal of Geophysical Research*, 101(D17), 22775–22785. <https://doi.org/10.1029/96jd02014>

- Waldron, S., Vihermaa, L., Evers, S., Garnett, M. H., Newton, J., & Henderson, A. C. G. (2019). C mobilisation in disturbed tropical peat swamps: Old DOC can fuel the fluvial efflux of old carbon dioxide but site recovery can occur. *Scientific Reports*, 9(1), 11429. <https://doi.org/10.1038/s41598-019-46534-9>
- Wang, Z.-P., Gu, Q., Deng, F.-D., Huang, J.-H., Megonigal, J. P., Yu, Q., ... Han, X.-G. (2016). Methane emissions from the trunks of living trees on upland soils. *New Phytologist*, 211, 429–439. <https://doi.org/10.1111/nph.13909>
- Wassmann, R., Thein, U. G., Whiticar, M. J., Rennenberg, H., Seiler, W., & Junk, W. J. (1992). Methane emissions from the Amazon Floodplain: Characterization of production and transport. *Global Biogeochemical Cycles*, 6, 3–13. <https://doi.org/10.1029/91GB01767>
- Webb, E. K., Pearman, G. I., & Leuning, R. (1980). Correction of flux measurements for density effects due to heat and water vapor transfer. *Quarterly Journal of the Royal Meteorological Society*, 106, 85–100. [https://doi.org/10.1002/\(ISSN\)1477-870X](https://doi.org/10.1002/(ISSN)1477-870X)
- Welch, B., Gauci, V., & Sayer, E. J. (2019). Tree stem bases are sources of CH₄ and N₂O in a tropical forest on upland soil during the dry to wet season transition. *Global Change Biology*, 25(1), 361–372. <https://doi.org/10.1111/gcb.14498>
- Whiting, G. J., & Chanton, J. P. (1996). Control of the diurnal pattern of methane emission from emergent aquatic macrophytes by gas transport mechanisms. *Aquatic Botany*, 54, 237–253. [https://doi.org/10.1016/0304-3770\(96\)01048-0](https://doi.org/10.1016/0304-3770(96)01048-0)
- Wijedasa, L. S., Sloan, S., Page, S. E., Clements, G. R., Lupascu, M., & Evans, T. A. (2018). Carbon emissions from South-East Asian peatlands will increase despite emission-reduction schemes. *Global Change Biology*, 24, 4598–4613. <https://doi.org/10.1111/gcb.14340>
- Wilczak, J. M., Oncley, S. P., & Stage, S. A. (2001). Sonic anemometer tilt correction algorithms. *Boundary Layer Meteorology*, 99, 127–150. <https://doi.org/10.1023/A:1018966204465>
- Winton, R. S., Flanagan, N., & Richardson, C. J. (2017). Neotropical peatland methane emissions along a vegetation and biogeochemical gradient. *PLoS ONE*, 12(10), e0187019. <https://doi.org/10.1371/journal.pone.0187019>
- Wong, G. X., Hirata, R., Hirano, T., Kiew, F., Aeries, E. B., Musin, K. K., & Melling, L. (2018). Micrometeorological measurement of methane flux above a tropical peat swamp forest. *Agricultural and Forest Meteorology*, 256–257, 353–361. <https://doi.org/10.1016/j.agrformet.2018.03.025>
- Wutzler, T., Lucas-Moffat, A., Migliavacca, M., Knauer, J., Sickel, K., Šigut, L., ... Reichstein, M. (2018). Basic and extensible post-processing of eddy covariance flux data with REddyProc. *Biogeosciences*, 15, 5015–5030. <https://doi.org/10.5194/bg-15-5015-2018>
- Xu, K. E., Pingintha-Durden, N., Luo, H., Durden, D., Sturtevant, C., Desai, A. R., ... Metzger, S. (2019). The eddy-covariance storage term in air: Consistent community resources improve flux measurement reliability. *Agricultural and Forest Meteorology*, 279, 107734. <https://doi.org/10.1016/j.agrformet.2019.107734>

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