



## Nitrous oxide (N<sub>2</sub>O) and methane (CH<sub>4</sub>) in rivers and estuaries of northwestern Borneo

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30 Abstract

Nitrous oxide ( $\text{N}_2\text{O}$ ) and methane ( $\text{CH}_4$ ) are atmospheric trace gases which play important roles of the climate and atmospheric chemistry of the Earth. However, little is known about their emissions from rivers and estuaries which seem to contribute significantly to the atmospheric budget of both gases. To  
35 this end concentrations of  $\text{N}_2\text{O}$  and  $\text{CH}_4$  were measured in the Rajang, Maludam, Sebuyau and Simunjan Rivers draining peatland in northwestern (NW) Borneo during two campaigns in March and September 2017. The Rajang River was additionally sampled in August 2016 and the Samusam and Sematan Rivers were additionally sampled in March 2017. The Maludam, Sebuyau, and Simunjan Rivers are typical ‘blackwater’ rivers with very low pH, very high dissolved organic carbon (DOC)  
40 concentrations and very low  $\text{O}_2$  concentrations. The spatial and temporal variability of  $\text{N}_2\text{O}$  and  $\text{CH}_4$  concentrations (saturation) in the six rivers/estuaries was large and ranged from  $2.0 \text{ nmol L}^{-1}$  (28 %) to  $41.4 \text{ nmol L}^{-1}$  (570 %) and from  $2.5 \text{ nmol L}^{-1}$  (106 %) to  $1372 \text{ nmol L}^{-1}$  (57,459 %), respectively. We found no overall trends of  $\text{N}_2\text{O}$  with  $\text{O}_2$  or  $\text{NO}_3^-$ ,  $\text{NO}_2^-$ ,  $\text{NH}_4^+$  and there were no trends of  $\text{CH}_4$  with  $\text{O}_2$  or dissolved nutrients or DOC.  $\text{N}_2\text{O}$  concentrations showed a positive linear correlation with  
45 rainfall. We conclude, therefore, that rainfall is the main factor determining the riverine  $\text{N}_2\text{O}$  concentrations since  $\text{N}_2\text{O}$  production/consumption in the ‘blackwater’ rivers themselves seems to be unlikely because of the low pH. In contrast  $\text{CH}_4$  concentrations showed an inverse relationship with rainfall.  $\text{CH}_4$  concentrations were highest at salinity = 0 and most probably result from methanogenesis as part of the decomposition of organic matter under anoxic conditions. We speculate  
50 that  $\text{CH}_4$  oxidation, which can be high when the water discharge is high (e.g. after rainfall events), is responsible for the decrease of the  $\text{CH}_4$  concentrations along the salinity gradients. The rivers and estuaries studied here were an overall net source of  $\text{N}_2\text{O}$  and  $\text{CH}_4$  to the atmosphere. The total annual  $\text{N}_2\text{O}$  and  $\text{CH}_4$  emissions were  $1.09 \text{ Gg N}_2\text{O yr}^{-1}$  ( $0.7 \text{ Gg N yr}^{-1}$ ) and  $23.8 \text{ Gg CH}_4 \text{ yr}^{-1}$ , respectively. This represents about 0.3 – 0.7 % of the global annual riverine and estuarine  $\text{N}_2\text{O}$  emissions and about  
55 0.1 – 1 % of the global riverine and estuarine  $\text{CH}_4$  emissions. Therefore, we conclude that rivers and estuaries in NW Borneo –despite the fact their water area covers only 0.05 % of the global river/estuarine area– contribute significantly to global riverine and estuarine emissions of  $\text{N}_2\text{O}$  and  $\text{CH}_4$ .

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## 1. Introduction

Nitrous oxide ( $\text{N}_2\text{O}$ ) and methane ( $\text{CH}_4$ ) are atmospheric trace gases which influence the climate and atmospheric chemistry of the Earth (IPCC, 2013; WMO, 2014). They act as greenhouse gases in the troposphere and are indirectly involved in stratospheric ozone depletion. Emission estimates indicate that rivers and estuaries contribute significantly to the atmospheric budget of both  $\text{N}_2\text{O}$  and  $\text{CH}_4$ .  $\text{N}_2\text{O}$  emissions estimate for rivers and estuaries range from 0.05 to 3.3 Tg  $\text{N}_2\text{O yr}^{-1}$  and from 0.09 to 5.7 Tg  $\text{N}_2\text{O yr}^{-1}$ , respectively (see overview in (Maavara et al., 2019)). Thus, the combined riverine and estuarine emissions may contribute up to 32 % to the global natural and anthropogenic emissions of  $\text{N}_2\text{O}$  (28.1 Tg  $\text{N}_2\text{O yr}^{-1}$ ; IPCC, 2013).  $\text{CH}_4$  emission estimates for rivers and estuaries are in the range of 1.5 – 26.8 Tg  $\text{CH}_4 \text{ yr}^{-1}$  (Bastviken et al., 2011; Stanley et al., 2016) and 0.8 – 6.6 Tg  $\text{CH}_4 \text{ yr}^{-1}$  (see overview in (Borges and Abril, 2011)), respectively. The combined emissions from rivers and estuaries can contribute up to 6% of the global natural and anthropogenic atmospheric emissions of  $\text{CH}_4$  (556 Tg  $\text{CH}_4 \text{ yr}^{-1}$ ; IPCC, 2013). As indicated by the wide range of the estimates cited above, the emission estimates of both gases are associated with a high degree of uncertainty, which is mainly caused by an inadequate coverage of the temporal and spatial distributions of  $\text{N}_2\text{O}$  and  $\text{CH}_4$  in rivers and estuaries and the inherent errors of the model approaches to estimate their release across the water/atmosphere interface (see e.g. (Alin et al., 2011; Borges and Abril, 2011)).

$\text{N}_2\text{O}$  is produced by microbial processes such as nitrification (i.e. oxidation of ammonia,  $\text{NH}_3$ , to nitrite,  $\text{NO}_2^-$ ) in estuarine waters (see e.g. (Barnes and Upstill-Goddard, 2011)) and heterotrophic denitrification (i.e. reduction of nitrate,  $\text{NO}_3^-$ , to dinitrogen,  $\text{N}_2$ ) in river sediments (Beaulieu et al., 2011). The yields of  $\text{N}_2\text{O}$  from these processes are enhanced under low oxygen (i.e. suboxic) conditions (see e.g. (Brase et al., 2017; Zhang et al., 2010)), whereas  $\text{N}_2\text{O}$  can be reduced to  $\text{N}_2$  under anoxic conditions via sedimentary denitrification in rivers (see e.g. (Upstill-Goddard et al., 2017)). Apart from ambient oxygen ( $\text{O}_2$ ) concentrations, riverine and estuarine  $\text{N}_2\text{O}$  production is also dependent on the concentrations of dissolved inorganic nitrogen, DIN ( $= \text{NH}_4^+ + \text{NO}_2^- + \text{NO}_3^-$ ). There seems to be a general trend towards high estuarine  $\text{N}_2\text{O}$  concentrations when DIN concentrations are high as well (Barnes and Upstill-Goddard, 2011; Zhang et al., 2010). However, this trend masks the fact that in many cases the spatial and temporal variability of riverine and estuarine  $\text{N}_2\text{O}$  is often not related to DIN (see e.g. (Borges et al., 2015; Brase et al., 2017; Müller et al., 2016a)).

$\text{CH}_4$  is produced during microbial respiration of organic matter by anaerobic methanogenesis in riverine and estuarine sediments (see e.g. (Borges and Abril, 2011; Romeijn et al., 2019; Stanley et al., 2016)). A significant fraction of the  $\text{CH}_4$  produced in sediments can be oxidized to carbon dioxide ( $\text{CO}_2$ ) via anaerobic  $\text{CH}_4$  oxidation in sulphate-reducing zones of estuarine sediments (see e.g. (Maltby et al., 2018)). When released to the overlying riverine/estuarine water  $\text{CH}_4$  can be oxidized by aerobic



CH<sub>4</sub> oxidation before reaching the atmosphere (see e.g. (Borges and Abril, 2011; Sawakuchi et al., 2016; Steinle et al., 2017)).

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In general, the temporal and spatial distributions of N<sub>2</sub>O and CH<sub>4</sub> in rivers and estuaries are driven by the complex interplay of microbial production and consumption pathways (see above) as well as physical processes such as input via shallow groundwater, river discharge, tidal pumping, release to the atmosphere and export to coastal waters (Barnes and Upstill-Goddard, 2011; Borges and Abril, 2011; Stanley et al., 2016).

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Peatlands, which are found in the tropics and at high latitudes, constitute one of the largest reservoirs of organic-bound carbon worldwide (Page et al., 2011; Treat et al., 2019; Yu et al., 2010). Rivers and streams draining peatlands have exceptionally high concentrations of dissolved organic carbon (DOC) and low pH and, thus, belong to the ‘blackwater’ river type which is also found in southeast (SE) Asia (Alkhatib et al., 2007; Baum et al., 2007; Martin et al., 2018; Moore et al., 2011; Rixen et al., 2008; Wit et al., 2015)..

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Despite the fact that a number of studies about N<sub>2</sub>O and CH<sub>4</sub> emissions from peatlands in southeast (SE) Asia have been published (see e.g. (Couwenberg et al., 2010; Hatano et al., 2016; Jauhiainen et al., 2012), only a few studies about their emissions from peatland draining rivers in SE Asia have been published so far (Jauhiainen and Silvennoinen, 2012; Müller et al., 2016a). Therefore, our knowledge about the biogeochemistry and emissions of N<sub>2</sub>O and CH<sub>4</sub> from peatland draining rivers is still rudimentary at best.

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Here we present measurements of dissolved N<sub>2</sub>O and CH<sub>4</sub> in six rivers and their estuaries in northwestern (NW) Borneo during August 2016, March 2017 and September 2017. The objectives of our study were (i) to measure the distributions of dissolved N<sub>2</sub>O and CH<sub>4</sub>, (ii) to identify the major factors influencing their distributions and (iii) to estimate the N<sub>2</sub>O and CH<sub>4</sub> emissions to the atmosphere.

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## 2. Study site description

Discrete samples of surface water were taken at several stations along the salinity gradients of the Rajang, Maludam, Sebuyau and Simunjan Rivers in NW Borneo during two campaigns in March and September 2017 (Figure 1, Table 1). The Rajang River was additionally sampled in August 2016 and the Samusam and Sematan Rivers were additionally sampled in March 2017. The environmental settings of the river basins are summarized in Table 2. Based on the areas affected by oil palm plantations and logging in combination with our own observations during several samplings

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135 campaigns, we classified the Rajang and Simunjan river basins as ‘disturbed’, the Maludam, Sebuyau,  
Sematan and Samusam river basins as ‘undisturbed’ (Table 2).

### 3. Methods

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#### 3.1 Measurements of N<sub>2</sub>O and CH<sub>4</sub>

Discrete water samples were taken as duplicates or triplicates in 20 or 37 mL glass vials from a water  
depth of approximately 1 m. The samples were poisoned immediately after sampling with a saturated  
aqueous mercuric chloride (HgCl<sub>2</sub>) solution. The samples were shipped to GEOMAR Helmholtz  
145 Centre for Ocean Research Kiel, Germany, for further analysis within a few weeks after sampling. For  
the determination of the N<sub>2</sub>O and CH<sub>4</sub> concentrations we applied the static-headspace equilibration  
method followed by gas chromatographic separation and detection with an electron capture detector  
(ECD, for N<sub>2</sub>O) and a flame ionization detector (FID, for CH<sub>4</sub>) as described in (Bastian, 2017) and  
(Kallert, 2017). Calibration of the ECD and FID were performed with standard gas mixtures of N<sub>2</sub>O  
150 and CH<sub>4</sub> in synthetic air which have been calibrated against NOAA-certified primary gas standards.

Dissolved N<sub>2</sub>O/CH<sub>4</sub> concentrations ( $C_{obs}$  in nmol L<sup>-1</sup>) were calculated with

$$C_{obs} = x'PV_{hs} / (RTV_{wp}) + x'\beta P \quad (1),$$

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where  $x'$  is the dry mole fraction of N<sub>2</sub>O or CH<sub>4</sub> in the headspace of the sample,  $P$  is the ambient  
pressure (set to 1013.25 hPa),  $V_{hs}$  and  $V_{wp}$  are the volumes of the headspace and the water phase,  
respectively.  $R$  stands for the gas constant (8.31451 m<sup>3</sup> Pa K<sup>-1</sup> mol<sup>-1</sup>),  $T$  is the temperature during  
equilibration and  $\beta$  is the solubility of N<sub>2</sub>O or CH<sub>4</sub> (Weiss and Price, 1980; Wiesenburg and Guinasso  
160 Jr., 1979). The estimated mean relative errors of the measurements were +/- 9 % and +/- 13 % for N<sub>2</sub>O  
and CH<sub>4</sub>, respectively. These comparably high relative errors most probably resulted from the long  
storage time for some of the samples. It was shown that CH<sub>4</sub> samples are more sensitive to storage  
time than N<sub>2</sub>O samples (Wilson et al., 2018).

#### 3.2 Ancillary measurements

Water temperature, dissolved oxygen, and salinity were recorded with an Aquaread® 2000. Nutrient  
measurements are described in detail in (Sia et al., 2019). In short, all samples were collected within  
the upper 1 m (surface) using pre-washed bottles via a pole-sampler to reduce contamination from the  
surface of the boat and engine coolant waters (Zhang et al., 2015). Samples were filtered through a 0.4  
170 µm pore-size polycarbonate membrane filters (Whatman) into pre-rinsed bottles, killed with  
concentrated HgCl<sub>2</sub> solution and kept in a cool, dark room. Nutrients were determined utilizing a



Skalar SANplus auto analyser with an analytical precision <5%. The measurements of dissolved organic carbon (DOC) are described in detail in (Martin et al., 2018). The DOC data are available from the supplementary material in (Martin et al., 2018).

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### 3.3 Computations of saturations and flux densities

The saturations ( $Sat$ , %) for  $N_2O$ ,  $CH_4$  and  $O_2$  were calculated as

$$Sat = 100 C_{obs} / C_{eq} \quad (2)$$

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where  $C_{eq}$  is the equilibrium concentration of  $N_2O/CH_4/O_2$  calculated according to (Weiss and Price, 1980), (Wiesenburg and Guinasso Jr., 1979) or (Weiss, 1970), respectively, with the *in-situ* temperature and salinity as well as the mean dry mole fractions of  $N_2O/CH_4$  at the time of the sampling. Mean monthly  $N_2O/CH_4$  dry mole fractions of  $329/1841 \cdot 10^{-9}$  (ppb),  $331/1880$  ppb and  $330/1852$  ppb for August 2016, March 2017 and September 2017, respectively, were measured at the atmospheric monitoring station Bukit Kototabang, located on the west coast of Sumatra (Indonesia). This station is operated by the NOAA/ESRL Global Monitoring Division program and data are available from <http://www.esrl.noaa.gov/gmd>. A saturation < 100 % indicates a concentration lower than the theoretical equilibrium concentration (i.e. undersaturation) and a saturation > 100 % indicates supersaturation.

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Flux densities ( $F$ ,  $nmol \cdot m^{-2} \cdot s^{-1}$ ) were calculated as

$$F = k_w (C_{obs} - C_{eq}) \quad (3)$$

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$$k_w = k_{600} (Sc/600)^{0.5} \quad (4)$$

$k_w$  is the gas transfer velocity and  $Sc$  is the Schmidt number, which was calculated with the equations for the kinematic viscosity of water (Siedler and Peters, 1986) and the diffusion of  $N_2O$  or  $CH_4$  in water (Jähne et al., 1987; Rhee et al., 2009).  $k_{600}$  was determined in a seasonal study for the Lupur and Saribas Rivers which are located in close vicinity to the Maludam River (Müller et al., 2016a; Müller et al., 2016b). We assume that the  $k_{600}$  values measured by (Müller et al., 2016a) are representative for the rivers in NW Borneo studied here. Mean  $k_{600}$  range from  $13.2 \text{ cm h}^{-1}$  (Lupur River) to  $23.9 \text{ cm h}^{-1}$  (Saribas River tributary). On the basis of the data in (Müller et al., 2016a) we computed a mean  $k_{600}$  of  $19.2 \text{ cm h}^{-1}$  ( $5.33 \cdot 10^{-5} \text{ m s}^{-1}$ ) which we used to estimate the flux densities of  $N_2O$  and  $CH_4$ . This  $k_{600}$  is in good agreement with the mean  $k_{600}$  for rivers and estuaries listed in (Alin et al., 2011) which range from  $4.8$  to  $35.3 \text{ cm h}^{-1}$ .

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### 3.4 Rainfall data



In order to account for the regional variability of the rainfall in NW Borneo, we used mean monthly  
210 rainfall data recorded at the weather stations in Kuching, Bandar Sri Aman and Sibul (all in NW  
Borneo). The rainfall data were provided by World Weather Online (Dubai, UAE, and Manchester,  
UK) and are available via <https://www.worldweatheronline.com/>. Representative weather stations  
were chosen for each river basin studied here and allocated as follows: The rainfall data for the  
Simunjan, Sematan and Samsuman River basins are represented by the data from Kuching, the  
215 Maludam/Sebuyau and the Rajang River basins are represented by the data from the Bandar Sri Aman  
and Sibul weather stations, respectively.

#### 4 Results and Discussion

220 All rivers showed low concentrations of DIN in the range from 1.1 to 29  $\mu\text{mol L}^{-1}$  (Table 1).  $\text{NO}_3^-$   
concentrations ranged from below the detection limit of 0.14  $\mu\text{mol L}^{-1}$  up to 19  $\mu\text{mol L}^{-1}$  and  $\text{NH}_4^+$   
concentrations were in the range of 0.3 to 17  $\mu\text{mol L}^{-1}$ . The Maludam, Sebuyau, and Simunjan Rivers  
can be classified as ‘blackwater’ rivers with low pH (3.7 – 4.8), high DOC concentrations (1960 –  
4387  $\mu\text{mol L}^{-1}$ ) and low  $\text{O}_2$  concentrations (31 – 95  $\mu\text{mol L}^{-1}$ ; 13 – 39 % saturation) at salinity = 0  
225 (Table 1). Comparable settings have been reported from other tropical ‘blackwater’ rivers in SE Asia  
as well (Alkhatib et al., 2007; Baum et al., 2007; Moore et al., 2011; Rixen et al., 2008; Wit et al.,  
2015).

##### 4.1 Nitrous oxide

230 The measured ranges of  $\text{N}_2\text{O}$  concentrations and saturations are listed in Table 3 and the distributions  
of  $\text{N}_2\text{O}$  saturations along the salinity gradients are shown in Figure 2.  $\text{N}_2\text{O}$  concentrations (saturations)  
were highly variable and ranged from 2.0  $\text{nmol L}^{-1}$  (28 %) in the Rajang River (at salinity = 0 in  
August 2016) to 41.4  $\text{nmol L}^{-1}$  (570 %) in the Simunjan River (at salinity = 0 in March 2017).  $\text{N}_2\text{O}$   
concentrations in the Rajang, Maludam and Sebuyau Rivers were generally higher in September 2017  
235 compared to March 2017 (Figure 2a-c). A decreasing linear trend of the  $\text{N}_2\text{O}$  saturations with salinity  
was only observed for the Rajang River in March 2017 (Figure 2a) indicating a conservative mixing  
and no  $\text{N}_2\text{O}$  sources or sinks along the salinity gradient. Our results are in general agreement with the  
 $\text{N}_2\text{O}$  measurements in the Lupar and Saribas Rivers (which are located in close vicinity of the  
Maludam River) in June 2013 and March 2014: Müller et al. (2016) measured  $\text{N}_2\text{O}$  concentrations  
240 (saturations) from 6.6 to 117  $\text{nmol L}^{-1}$  (102 to 1679 %) in the Lupar and Saribas Rivers. Salinity and  
 $\text{N}_2\text{O}$  concentrations in the Lupar and Saribas Rivers were negatively correlated in June 2013 but were  
not correlated in March 2014 (Müller et al., 2016a). In contrast to our study, no  $\text{N}_2\text{O}$  undersaturations  
have been observed by (Müller et al., 2016a). Our results are at the lower end of  $\text{N}_2\text{O}$  concentrations  
reported from rivers around the globe which can range from extreme undersaturation (down to about 3  
245 %, i.e. almost devoid of  $\text{N}_2\text{O}$ ) as measured in a tropical river in Africa (Borges et al., 2015; Upstill-



Goddard et al., 2017) to extreme supersaturation (of up to 6500%) as measured in a river in Europe ((Barnes and Upstill-Goddard, 2011).

250 Maximum N<sub>2</sub>O saturations measured in March 2017 were in the range from 106 % to 142 % for the rivers classified as undisturbed (Maludam, Sebuyau, Sematan and Samusam) whereas the maximum saturation for the rivers classified as disturbed (Rajang and Simunjan) were in the range from 329 % to 570 % (Tables 2 and 3) indicating higher emissions from the disturbed rivers. The maximum N<sub>2</sub>O saturations in September 2017 ranged from 329 % to 390 % and no differences were observed between undisturbed and disturbed rivers (Table 3).

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We found no overall trends of N<sub>2</sub>O with O<sub>2</sub> or NO<sub>3</sub><sup>-</sup>, NO<sub>2</sub><sup>-</sup>, NH<sub>4</sub><sup>+</sup> and DIN. Therefore, it is difficult to decipher the major consumption or production processes of N<sub>2</sub>O or to locate the influence of (local) anthropogenic input of nitrogen compounds on riverine N<sub>2</sub>O cycling. This is in line with results from studies of other tropical rivers (Borges et al., 2015; Müller et al., 2016a). N<sub>2</sub>O production via nitrification depends on the prevailing pH because nitrifiers prefer to take up ammonia (NH<sub>3</sub>). The concentration of dissolved NH<sub>3</sub> is dropping significantly at pH < 8-9 (Bange, 2008) because of its easy protonation to ammonium (NH<sub>4</sub><sup>+</sup>). A low pH of about 5-6 can reduce nitrification (NH<sub>4</sub><sup>+</sup> oxidation) significantly as it was recently shown for the Tay Ninh River in Vietnam (Le et al., 2019). Moreover, the optimum for a net N<sub>2</sub>O production by nitrification, nitrifier-denitrification and denitrification lies between a pH of 7 – 7.5 (Blum et al., 2018). Therefore, a net N<sub>2</sub>O production may be unlikely in the ‘blackwater’ rivers studied here with their low pH (see Table 1). The observed N<sub>2</sub>O supersaturations, therefore, might have been the result of external inputs of N<sub>2</sub>O-enriched waters or groundwater. The observed N<sub>2</sub>O undersaturations were most probably resulting from heterotrophic denitrification which could have taken place either in organic matter-enriched anoxic river sediments or in anoxic environments of the surrounding soils. However, the main factor for riverine N<sub>2</sub>O under- or supersaturation might be rainfall, because rainfall events determine the height of the water table in the surrounding soils, which in turn determines the amount of suboxic/anoxic conditions favourable for N<sub>2</sub>O production or consumption. See also discussion in Section 4.3.

#### 275 4.2 Methane

The measured ranges of CH<sub>4</sub> concentrations and saturations are listed in Table 3 and the distributions of CH<sub>4</sub> saturations along the salinity gradients are shown in Figure 3. CH<sub>4</sub> concentrations (saturations) were highly variable and ranged from 2.5 nmol L<sup>-1</sup> (106 %) in the Simunjan River (at salinity = 0 in September 2017) to 1372 nmol L<sup>-1</sup> (57,459 %) in the Simunjan River (at salinity = 0 in March 2017). (Please note that we also measured a CH<sub>4</sub> concentration of 14,999 nmol L<sup>-1</sup> (624,070 %) at one station in the Simunjan River at salinity = 0 in March 2017 which, however, was not included in Figure 3 and which was not used in further computations because of statistical reasons.) CH<sub>4</sub> saturations in the



Rajang, Maludam, Sebuyau and Simunjan Rivers were higher in March 2017 compared to September 2017. Maximum CH<sub>4</sub> concentrations were measured at salinity = 0 and there was a general decrease of  
285 CH<sub>4</sub> with increasing salinity. Exceptions from this trend occurred at individual stations in the Maludam, Sebuyau and Samusam Rivers which point to local sources of CH<sub>4</sub> (Figure 3). The range of CH<sub>4</sub> concentrations (saturations) from our study is larger compared to the concentration range measured in the Lupar and Saribas Rivers (3.7 – 113.9 nmol L<sup>-1</sup>; 168 – 5058 %) ((Müller et al., 2016a). (Borges et al., 2015) reported a maximum CH<sub>4</sub> concentration (saturation) of 62,966 nmol L<sup>-1</sup>  
290 (appr. 954,000 %) in their study of tropical rivers in Africa which is much higher than the maximum concentration measured in our study. (Bouillon et al., 2014)

We found no overall trends of CH<sub>4</sub> with O<sub>2</sub> or dissolved nutrients or DOC along the salinity gradients. High CH<sub>4</sub> concentrations, which were often associated with high DOC and low O<sub>2</sub> concentrations at  
295 salinity = 0, might have been produced by methanogenesis in anoxic riverine sediments rich in organic material or in anoxic parts of the surrounding soils drained by the rivers. The decrease of CH<sub>4</sub> with increasing salinity can be attributed to the gas exchange across the river water/atmosphere interface in combination with CH<sub>4</sub> oxidation (Borges and Abril, 2011; Sawakuchi et al., 2016).

300 We found no differences in the CH<sub>4</sub> saturations between the rivers classified as undisturbed and those classified as disturbed in both March and September 2017.

#### 4.3 N<sub>2</sub>O/CH<sub>4</sub> concentrations and rainfall

Mean N<sub>2</sub>O concentrations showed a linear correlation with rain fall (Figure 4a). Enhanced N<sub>2</sub>O  
305 emissions from (peat) soils are usually associated with rainfall when the water table approaches the soil surface (Couwenberg et al., 2010; Jauhiainen et al., 2016). A high water table, in turn, allows decomposition of previously deposited fresh organic material (Jauhiainen et al., 2016) and, thus, will result in favourable conditions for microbial N<sub>2</sub>O production mainly via denitrification in a suboxic/anoxic soil environment (Pihlatie et al., 2004). N<sub>2</sub>O production via nitrification may be less  
310 important at high water table (Pihlatie et al., 2004). Therefore, the positive linear relationship of the riverine N<sub>2</sub>O concentrations with rainfall might result from enhanced N<sub>2</sub>O production in the adjacent soils drained by the rivers.

In contrast to N<sub>2</sub>O, the mean CH<sub>4</sub> concentrations decrease with increasing rainfall (Figure 4b). Under  
315 the assumption that rainfall is a predictor for river discharge/high water we can argue that our result are in agreement with (i) the often observed inverse relationship between CH<sub>4</sub> concentrations and river discharge (Anthony et al., 2012; Bouillon et al., 2014; Dinsmore et al., 2013; Hope et al., 2001) and (ii) the enhancement of CH<sub>4</sub> oxidation during high waters: (Sawakuchi et al., 2016) showed that CH<sub>4</sub> oxidation in ‘blackwater’ rivers of the Amazon basin was maximal during the high water season



320 resulting in a reduction of up to 96% of the diffusive flux of CH<sub>4</sub> (i.e. its input to the river and its  
release to the atmosphere) (Sawakuchi et al., 2016). This was explained by the higher river water  
levels which, in turn, could enhance CH<sub>4</sub> oxidation because of a longer residence time of CH<sub>4</sub> in the  
sediment and river water (Sawakuchi et al., 2016).

#### 325 4.4 Emission estimates

The N<sub>2</sub>O flux densities from the six rivers studied here are comparable to the N<sub>2</sub>O flux densities from  
other aqueous and soil systems reported from Borneo and other sites in SE Asia, see Table 4. The  
corresponding CH<sub>4</sub> flux densities are higher than the CH<sub>4</sub> flux densities reported for the Lupar and  
Saribas Rivers but much lower than the flux densities from drainage canals in Central Kalimantan and  
330 Sumatra (Jauhainen and Silvennoinen, 2012) (Table 4). Our CH<sub>4</sub> flux densities are, however,  
comparable to recently published CH<sub>4</sub> eddy covariance measurements (Tang et al., 2018) in the  
Maludam National Park, which is drained by the Maludam River, and measurements of the CH<sub>4</sub>  
release from peat soils when the water table is high and CH<sub>4</sub> from rice paddies (Couwenberg et al.,  
2010), see Table 4. The mean annual N<sub>2</sub>O and CH<sub>4</sub> emissions for the individual rivers were calculated  
335 by multiplying the mean flux density, *F*, for each river (Table 4) with the river surface area given in  
Table 2. The results are listed in Table 5. The resulting total annual N<sub>2</sub>O emissions for the rivers in  
NW Borneo -including the emissions from the Lupar and Saribas Rivers (Müller et al., 2016a)- are  
1.09 Gg N<sub>2</sub>O yr<sup>-1</sup> (0.7 Gg N yr<sup>-1</sup>). This represents about 0.3 – 0.7 % of the global annual riverine and  
estuarine N<sub>2</sub>O emissions of 166 – 322 Gg N<sub>2</sub>O (106 – 205 Gg N yr<sup>-1</sup>) recently estimated by (Maavara  
340 et al., 2019). The total annual CH<sub>4</sub> emissions from rivers in NW Borneo are 23.8 Gg CH<sub>4</sub> yr<sup>-1</sup>. This  
represents about 0.1 – 1 % of the global riverine and estuarine CH<sub>4</sub> emissions of 2300 – 33,400 Gg  
CH<sub>4</sub> yr<sup>-1</sup> (the emission range is based on the minimum and maximum estimates given in (Bange et al.,  
1994; Bastviken et al., 2011; Borges and Abril, 2011; Stanley et al., 2016). However, we caution that  
our estimates are associated with a high degree of uncertainty because (i) our data are biased by the  
345 fact that for some rivers it was not possible to cover the entire salinity gradient and (ii) seasonal and  
internannual variabilities are not adequately represented in our data set.

## 5 Summary and Conclusions

350 N<sub>2</sub>O and CH<sub>4</sub> were measured in the Rajang, Maludam, Sebuyau and Simuntan Rivers and Estuaries in  
NW Borneo during two campaigns in March and September 2017. The Rajang River was additionally  
sampled in August 2016 and the Samusam and Sematan Rivers were additionally sampled in March  
2017. The spatial and temporal variability of N<sub>2</sub>O and CH<sub>4</sub> concentrations was large. N<sub>2</sub>O  
concentrations (saturations) ranged from 2.0 nmol L<sup>-1</sup> (28 %) in the Rajang River (at salinity = 0 in  
355 August 2016) to 41.4 nmol L<sup>-1</sup> (570 %) in the Simunjan River (at salinity = 0 in March 2017). CH<sub>4</sub>  
concentrations (saturations) were in the range from 2.5 nmol L<sup>-1</sup> (106 %) in the Simunjan River (at



salinity = 0 in September 2017) to 1372 nmol L<sup>-1</sup> (57,459 %) in the Simunjan River (at salinity = 0 in March 2017). N<sub>2</sub>O concentrations showed a positive linear correlation with rainfall. We conclude, therefore, that rainfall, which determines the N<sub>2</sub>O production/consumption in the surrounding soils, is the main factor determining the riverine N<sub>2</sub>O concentrations. N<sub>2</sub>O production in the ‘blackwater’ rivers themselves seems to be unlikely because of the low pH. In contrast CH<sub>4</sub> concentrations showed an inverse relationship with rainfall. CH<sub>4</sub> concentrations were highest at salinity = 0 and most probably results from methanogenesis as part of the decomposition of organic matter under anoxic conditions. We speculate that CH<sub>4</sub> oxidation, which can be high when the water discharge is high (e.g. after rainfall events), is responsible for the reduction of the CH<sub>4</sub> concentrations along the salinity gradient. The rivers and estuaries studied here were an overall net source of N<sub>2</sub>O and CH<sub>4</sub> to the atmosphere. The total annual N<sub>2</sub>O and CH<sub>4</sub> emissions were 1.09 Gg N<sub>2</sub>O yr<sup>-1</sup> (0.7 Gg N yr<sup>-1</sup>) and 23.8 Gg CH<sub>4</sub> yr<sup>-1</sup>, respectively. This represents about 0.3 – 0.7 % of the global annual riverine and estuarine N<sub>2</sub>O emissions and about 0.1 – 1 % of the global riverine and estuarine CH<sub>4</sub> emissions. Rivers and estuaries in NW Borneo contribute only 0.05 % (= 7.9 10<sup>2</sup> km<sup>2</sup> including the surface areas of the Lupar and Saribas Rivers; (Müller et al., 2016a) to the global water surface area of rivers and estuaries (= 1.7 10<sup>6</sup> km<sup>2</sup>; (Maavara et al., 2019)). Therefore we conclude that rivers and estuaries in NW Borneo contribute significantly to the global riverine and estuarine emissions of both N<sub>2</sub>O and CH<sub>4</sub>.

The environment of Borneo (and SE Asia) is affected by rapid changes due to (i) anthropogenic activities such as conversion of peatland into oil palm plantations etc. (see e.g. (Austin et al., 2018; McAlpine et al., 2018; Schoneveld et al., 2019)) and (ii) climatic changes (see e.g. (Sa’adi et al., 2017a, b; Tang, 2019)) which, in turn, could significantly affect N<sub>2</sub>O and CH<sub>4</sub> emissions from soils (see e.g. (Jauhiainen et al., 2016; Oktarita et al., 2017)). But little is known about how these changes will affect N<sub>2</sub>O and CH<sub>4</sub> emissions from aqueous systems such as rivers and estuaries in the future. The obvious relationship of N<sub>2</sub>O and CH<sub>4</sub> concentrations and rainfall could be used to predict future concentrations and its associated emissions to the atmosphere. However, the trends of rainfall and river discharge in Borneo show a high local variability and no general common trend (Sa’adi et al., 2017a; Tang, 2019). Therefore, predictions of future trends of N<sub>2</sub>O and CH<sub>4</sub> emissions will be associated with high degree of uncertainty. In order to improve our knowledge to predicted future changes of N<sub>2</sub>O and CH<sub>4</sub> riverine/estuarine emissions we suggest establishing regular measurements in the rivers and along the salinity gradients. This will help deciphering the temporal and spatial variability of N<sub>2</sub>O and CH<sub>4</sub> emissions from tropical rivers and estuaries. Moreover, studies of the relevant production/consumption pathways (and their main driving factors) for both gases are required. A suitable framework for this could be the recently published concept of the global N<sub>2</sub>O Ocean Observation Network (N<sub>2</sub>O-ON) (Bange et al., 2019).



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## 7 References

- Alin, S. R., de Fátima F. L. Rasera, M., Salimon, C. I., Richey, J. E., Holtgrieve, G. W., Krusche, A.  
V., and Snidvongs, A.: Physical controls on carbon dioxide transfer velocity and flux in low-  
415 gradient river systems and implications for regional carbon budgets, *Journal of Geophysical  
Research: Biogeosciences*, 116, 2011.
- Alkhatib, M., Jennerjahn, T. C., and Samiaji, J.: Biogeochemistry of the Dumai River estuary,  
Sumatra, Indonesia, a tropical black-water river, *Limnology and Oceanography*, 52, 2410-2417,  
2007.
- 420 Allen, K., Hassler, E., Kurniawan, S., Veldkamp, E., and Correa, M. D.: Canopy soil of oil palm  
plantations emits methane and nitrous oxide, *Soil Biology and Biochemistry*, 122, 1-6, 2018.
- Anthony, S. E., Prahl, F. G., and Peterson, T. D.: Methane dynamics in the Willamette River, Oregon,  
*Limnology and Oceanography*, 57, 1517-1530, 2012.
- Austin, K. G., Harris, N. L., Wijaya, A., Murdiyarso, D., Harvey, T., Stolle, F., and Kasibhatla, P. S.:  
425 A review of land-based greenhouse gas flux estimates in Indonesia, *Environmental Research  
Letters*, 13, 055003, 2018.
- Bange, H. W.: Gaseous nitrogen compounds (NO, N<sub>2</sub>O, N<sub>2</sub>, NH<sub>3</sub>) in the ocean. In: *Nitrogen in the  
Marine Environment*, 2nd Edition, Capone, D. G., Bronk, D. A., Mulholland, M. R., and Carpenter,  
E. J. (Eds.), Elsevier, Amsterdam, 2008.
- 430 Bange, H. W., Arévalo-Martínez, D. L., de la Paz, M., Farías, L., Kaiser, J., Kock, A., Law, C. S.,  
Rees, A. P., Rehder, G., Tortell, P. D., Upstill-Goddard, R. C., and Wilson, S. T.: A Harmonized  
Nitrous Oxide (N<sub>2</sub>O) Ocean Observation Network for the 21st Century, *Frontiers in Marine  
Science*, 6, 2019.
- 435 Bange, H. W., Bartell, U. H., Rapsomanikis, S., and Andreae, M. O.: Methane in the Baltic and North  
Seas and a reassessment of the marine emissions of methane, *Global Biogeochem. Cycles*, 8, 465-  
480, 1994.



- Barnes, J. and Upstill-Goddard, R. C.: N<sub>2</sub>O seasonal distribution and air-sea exchange in UK estuaries: Implications for tropospheric N<sub>2</sub>O source from European coastal waters *Journal of Geophysical Research*, 116, G01006; doi:10.1029/2009JG001156, 2011.
- 440 Bastian, D.: N<sub>2</sub>O und CH<sub>4</sub> Verteilung in Ästuaren und Flüssen im Nordwesten von Borneo, 2017.BSc thesis, Kiel University, Kiel, 50 pp., 2017.
- Bastviken, D., Tranvik, L. J., Downing, J. A., Crill, P. M., and Enrich-Prast, A.: Freshwater methane emissions offset continental carbon sink, *Science*, 331, 50, 2011.
- 445 Baum, A., Rixen, T., and Samiaji, J.: Relevance of peat draining rivers in central Sumatra for the riverine input of dissolved organic carbon into the ocean, *Estuarine, Coastal and Shelf Science* 73 563-570, 2007.
- Beaulieu, J. J., Tank, J. L., Hamilton, S. K., Wollheim, W. M., Hall, R. O., Mulholland, P. J., Peterson, B. J., Ashkenas, L. R., Cooper, L. W., Dahm, C. N., Dodds, W. K., Grimm, N. B., Johnson, S. L., McDowell, W. H., Poole, G. C., Valett, H. M., Arango, C. P., Bernot, M. J., Burgin, A. J., 450 Crenshaw, C. L., Helton, A. M., Johnson, L. T., O'Brien, J. M., Potter, J. D., Sheibley, R. W., Sobota, D. J., and Thomas, S. M.: Nitrous oxide emission from denitrification in stream and river networks, *Proceedings of the National Academy of Sciences*, 108, 214-219, 2011.
- Blum, J.-M., Su, Q., Ma, Y., Valverde-Pérez, B., Domingo-Félez, C., Jensen, M. M., and Smets, B. F.: 455 The pH dependency of N-converting enzymatic processes, pathways and microbes: effect on net N<sub>2</sub>O production, *Environmental Microbiology*, 20, 1623-1640, 2018.
- Borges, A. V. and Abril, G.: Carbon dioxide and methane dynamics in estuaries. In: *Treatise on estuarine and coastal science - vol. 5: Biogeochemistry*, Wolanski, E. and McLusky, D. (Eds.), Academic Press, Waltham, 2011.
- Borges, A. V., Darchambeau, F., Teodoru, C. R., Marwick, T. R., Tamooh, F., Geeraert, N., Omengo, 460 F. O., Guérin, F., Lambert, T., Morana, C., Okuku, E., and Bouillon, S.: Globally significant greenhouse-gas emissions from African inland waters, *Nature Geoscience*, 8, 673-642, 2015.
- Bouillon, S., Yambélé, A., Gillikin, D. P., Teodoru, C., Darchambeau, F., Lambert, T., and Borges, A. V.: Contrasting biogeochemical characteristics of the Oubangui River and tributaries (Congo River basin), *Scientific Reports*, 4, 5402, 2014.
- 465 Brase, L., Bange, H. W., Lendt, R., Sanders, T., and Dähnke, K.: High resolution measurements of nitrous oxide (N<sub>2</sub>O) in the Elbe estuary, *Frontiers in Marine Science*, 4, 2017.
- Couwenberg, J., Dommain, R., and Joosten, H.: Greenhouse gas fluxes from tropical peatlands in south-east Asia, *Global Change Biology*, 16, 1715-1732, 2010.
- Dinsmore, K. J., Billett, M. F., and Dyson, K. E.: Temperature and precipitation drive temporal 470 variability in aquatic carbon and GHG concentrations and fluxes in a peatland catchment, *Global Change Biology*, 19, 2133-2148, 2013.
- Hatano, R., Toma, Y., Hamada, Y., Arai, H., Susilawati, H. L., and Inubushi, K.: Methane and nitrous oxide emissions from tropical peat soil. In: *Tropical Peatland Ecosystems*, Osaki, M. and Tsuji, N. (Eds.), Springer Japan, Tokyo, 2016.
- 475 Hope, D., Palmer, S. M., Billett, M. F., and Dawson, J. J. C.: Carbon dioxide and methane evasion from a temperate peatland stream, *Limnology and Oceanography*, 46, 847-857, 2001.
- IPCC (Ed.): *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, Cambridge University Press, Cambridge UK and New York, NY, USA, 2013.
- 480 Jähne, B., Heinz, G., and Dietrich, W.: Measurements of the diffusion coefficients of sparingly soluble gases in water, *Journal of Geophysical Research*, 92, 10,767-710,776, 1987.
- Jauhiainen, J., Page, S. E., and Vasander, H.: Greenhouse gas dynamics in degraded and restored tropical peatlands, *Mires and Peat*, 17, 1-12, 2016.
- Jauhiainen, J. and Silvennoinen, H.: Diffusion GHG fluxes at tropical peatland drainage canal water 485 surfaces, *Suo* 63, 93-105, 2012.
- Jauhiainen, J., Silvennoinen, H., Hämäläinen, R., Kusin, K., Limin, S., Raison, R. J., and Vasander, H.: Nitrous oxide fluxes from tropical peat with different disturbance history and management, *Biogeosciences*, 9, 1337-1350, 2012.
- Kallert, J.: Verteilung von Lachgas (N<sub>2</sub>O) und Methan (CH<sub>4</sub>) im Fluss Rajang (Malaysia), 2017.BSc, University of Kiel, Kiel, 25 pp., 2017.
- 490 Le, T. T. H., Fettig, J., and Meon, G.: Kinetics and simulation of nitrification at various pH values of a polluted river in the tropics, *Ecology & Hydrobiology*, 19, 54-65, 2019.



- 495 Maavara, T., Lauerwald, R., Laruelle, G. G., Akbarzadeh, Z., Bouskill, N. J., Van Cappellen, P., and  
Regnier, P.: Nitrous oxide emissions from inland waters: Are IPCC estimates too high?, *Global  
Change Biology*, 25, 473-488, 2019.
- Maltby, J., Steinle, L., Löscher, C. R., Bange, H. W., Fischer, M. A., Schmidt, M., and Treude, T.:  
Microbial methanogenesis in the sulfate-reducing zone of sediments in the Eckernförde Bay, SW  
Baltic Sea, *Biogeosciences*, 15, 137-157, 2018.
- 500 Martin, P., Cherukuru, N., Tan, A. S. Y., Sanwani, N., Mujahid, A., and Müller, M.: Distribution and  
cycling of terrigenous dissolved organic carbon in peatland-draining rivers and coastal waters of  
Sarawak, Borneo, *Biogeosciences*, 15, 6847-6865, 2018.
- McAlpine, C. A., Johnson, A., Salazar, A., Syktus, J., Wilson, K., Meijaard, E., Seabrook, L.,  
Dargusch, P., Nordin, H., and Sheil, D.: Forest loss and Borneo's climate, *Environmental Research  
Letters*, 13, 044009, 2018.
- 505 Melling, L., Hatano, R., and Goh, K. J.: Methane fluxes from three ecosystems in tropical peatland of  
Sarawak, Malaysia, *Soil Biology & Biochemistry* 37, 1445-1453, 2005.
- Melling, L., Hatano, R., and Goh, K. J.: Nitrous oxide emissions from three ecosystems in tropical  
peatland of Sarawak, Malaysia, *Soil Science and Plant Nutrition*, 53, 792-805, 2007.
- 510 Moore, S., Gauci, V., Evans, C. D., and Page, S. E.: Fluvial organic carbon losses from a Bornean  
blackwater river, *Biogeosciences*, 8, 901-909, 2011.
- Müller, D., Bange, H. W., Warneke, T., Rixen, T., Müller, M., Mujahid, A., and Notholt, J.: Nitrous  
oxide and methane in two tropical estuaries in a peat-dominated region of northwestern Borneo,  
*Biogeosciences*, 13, 2415-2428, 2016a.
- 515 Müller, D., Warneke, T., Rixen, T., Müller, M., Mujahid, A., Bange, H. W., and Notholt, J.: Fate of  
terrestrial organic carbon and associated CO<sub>2</sub> and CO emissions from two Southeast Asian  
estuaries, *Biogeosciences*, 13, 691-705, 2016b.
- Oktarita, S., Hergoualc'h, K., Anwar, S., and Verchot, L. V.: Substantial N<sub>2</sub>O emissions from peat  
decomposition and N fertilization in an oil palm plantation exacerbated by hotspots, *Environmental  
Research Letters*, 12 2017.
- 520 Page, S. E., Riley, J. O., and Banks, C. J.: Global and regional importance of the tropical peatland  
carbon pool, *Global Change Biology*, 17, 798-818, 2011.
- Pihlatie, M., Syväsalo, E., Simojoki, A., Esala, M., and Regina, K.: Contribution of nitrification and  
denitrification to N<sub>2</sub>O production in peat, clay and loamy sand soils under different soil moisture  
conditions, *Nutrient Cycling in Agroecosystems*, 70, 135-141, 2004.
- 525 Rhee, T. S., Kettle, A. J., and Andreae, M. O.: Methane and nitrous oxide emissions from the ocean: A  
reassessment using basin-wide observations in the Atlantic *Journal of Geophysical Research*, 114,  
D12304, doi: 12310.11029/12008JD011662, 2009.
- Rixen, T., Baum, A., Pohlmann, T., Balzer, W., Samiaji, J., and Jose, C.: The Siak, a tropical black  
water river in central Sumatra on the verge of anoxia, *Biogeochemistry*, 90, 129-140, 2008.
- 530 Romeijn, P., Comer-Warner, S. A., Ullah, S., Hannah, D. M., and Krause, S.: Streambed Organic  
Matter Controls on Carbon Dioxide and Methane Emissions from Streams, *Environmental Science  
& Technology*, 53, 2364-2374, 2019.
- Sa'adi, Z., Shahid, S., Ismail, T., Chung, E.-S., and Wang, X.-J.: Distributional changes in rainfall and  
river flow in Sarawak, Malaysia, *Asia-Pac. J. Atmos. Sci.*, 53, 489-500, 2017a.
- 535 Sa'adi, Z., Shahid, S., Ismail, T., Chung, E.-S., and Wang, X.-J.: Trends analysis of rainfall and  
rainfall extremes in Sarawak, Malaysia, using modified Mann-Kendall test, *Meteorology and  
Atmospheric Physics*, <https://doi.org/10.1007/s00703-017-0564-3>, 2017b.
- Sawakuchi, H. O., Bastviken, D., Sawakuchi, A. O., Ward, N. D., Borges, C. D., Tsai, S. M., Richey,  
J. E., Ballester, M. V. R., and Krusche, A. V.: Oxidative mitigation of aquatic methane emissions  
in large Amazonian rivers, *Global Change Biology*, 22, 1075-1085, 2016.
- 540 Schoneveld, G. C., Ekowati, D., Andrianto, A., and van der Haar, S.: Modeling peat- and forestland  
conversion by oil palm smallholders in Indonesian Borneo, *Environmental Research Letters*, 14,  
014006, 2019.
- 545 Sia, E. S. A., Zhang, J., Shan, J., Zhu, Z., Cheah, W., Carrasco, G., Jang, F. H., Mujahid, A., and  
Müller, M.: Behavior of dissolved phosphorus with the associated nutrients in relation to  
phytoplankton biomass of the Rajang River-South China Sea continuum, *Biogeosci. Discuss.*,  
submitted, 2019.



- Siedler, G. and Peters, H.: Properties of sea water. In: *Oceanography*, Sündermann, J. (Ed.), Landolt-Börnstein, New Series, Springer Verlag, Berlin, 1986.
- 550 Stanley, E. H., Casson, N. J., Christel, S. T., Crawford, J. T., Loken, L. C., and Oliver, S. K.: The ecology of methane in streams and rivers: Patterns, controls, and global significance, *Ecological Monographs*, 86, 146-171, 2016.
- Staub, J. R., Among, H. L., and Gastaldo, R. A.: Seasonal sediment transport and deposition in the Rajang River delta, Sarawak, East Malaysia, *Sedimentary Geology*, 133, 249–264, 2000.
- 555 Steinle, L., Maltby, J., Treude, T., Kock, A., Bange, H. W., Engbersen, N., Zopfi, J., Lehmann, M. F., and Niemann, H.: Effects of low oxygen concentrations on aerobic methane oxidation in seasonally hypoxic coastal waters, *Biogeosciences*, 14, 1631-1645, 2017.
- Tang, A. C. I., Stoy, P. C., Hirata, R., Musin, K. K., Aeries, E. B., Wenceslaus, J., and Melling, L.: Eddy Covariance Measurements of Methane Flux at a Tropical Peat Forest in Sarawak, Malaysian Borneo, *Geophysical Research Letters*, 45, 4390-4399, 2018.
- 560 Tang, K. H. D.: Climate change in Malaysia: Trends, contributors, impacts, mitigation and adaptations, *Science of the Total Environment* 650, 1858–1871, 2019.
- Treat, C. C., Kleinen, T., Broothaerts, N., Dalton, A. S., Dommain, R., Douglas, T. A., Drexler, J. Z., Finkelstein, S. A., Grosse, G., Hope, G., Hutchings, J., Jones, M. C., Kuhry, P., Lacourse, T., 565 Lähteenoja, O., Loisel, J., Notebaert, B., Payne, R. J., Peteet, D. M., Sannel, A. B. K., Stelling, J. M., Strauss, J., Swindles, G. T., Talbot, J., Tarnocai, C., Verstraeten, G., Williams, C. J., Xia, Z., Yu, Z., Väliiranta, M., Hättestrand, M., Alexanderson, H., and Brovkin, V.: Widespread global peatland establishment and persistence over the last 130,000 y, *Proceedings of the National Academy of Sciences*, 116, 4822-4827, 2019.
- 570 Upstill-Goddard, R. C., Salter, M. E., Mann, P. J., Barnes, J., Poulsen, J., Dinga, B., Fiske, G. J., and Holmes, R. M.: The riverine source of CH<sub>4</sub> and N<sub>2</sub>O from the Republic of Congo, western Congo Basin, *Biogeosciences*, 14, 2267-2281, 2017.
- Weiss, R. F.: The solubility of nitrogen, oxygen and argon in water and seawater, *Deep-Sea Res.*, 17, 721-735, 1970.
- 575 Weiss, R. F. and Price, B. A.: Nitrous oxide solubility in water and seawater, *Marine Chemistry*, 8, 347-359, 1980.
- Wiesenburg, D. A. and Guinasso Jr., N. L.: Equilibrium solubilities of methane, carbon monoxide, hydrogen in water and seawater, *J. Chem. Eng. Data*, 24, 356-360, 1979.
- Wilson, S. T., Bange, H. W., Arévalo-Martínez, D. L., Barnes, J., Borges, A. V., Brown, I., Bullister, 580 J. L., Burgos, M., Capelle, D. W., Casso, M., de la Paz, M., Farías, L., Fenwick, L., Ferrón, S., Garcia, G., Glockzin, M., Karl, D. M., Kock, A., Laperriere, S., Law, C. S., Manning, C. C., Marriner, A., Myllykangas, J. P., Pohlman, J. W., Rees, A. P., Santoro, A. E., Tortell, P. D., Upstill-Goddard, R. C., Wisegarver, D. P., Zhang, G. L., and Rehder, G.: An intercomparison of oceanic methane and nitrous oxide measurements, *Biogeosciences*, 15, 5891-5907, 2018.
- 585 Wit, F., Müller, D., Baum, A., Warneke, T., Pranowo, W. S., Müller, M., and Rixen, T.: The impact of disturbed peatlands on river outgassing in Southeast Asia, *Nature Communications*, 6, 10155, 2015.
- WMO: Scientific Assessment of Ozone Depletion: 2014, Geneva, Switzerland Global Ozone Research and Monitoring Project—Report No. 55, 416 pp., 2014.
- 590 Yu, Z., Loisel, J., Brosseau, D. P., Beilman, D. W., and Hunt, S. J.: Global peatland dynamics since the Last Glacial Maximum, *Geophysical Research Letters*, 37, 2010.
- Zhang, G.-L., Zhang, J., Liu, S.-M., Ren, J.-L., and Zhao, Y.-C.: Nitrous oxide in the Changjiang (Yangtze River) estuary and its adjacent marine area: Riverine input, sediment release and atmospheric fluxes, *Biogeosciences*, 7, 3505-3516, 2010.
- 595 Zhang, R., John, S. G., Zhang, J., Ren, J., Wu, Y., Zhu, Z., Liu, S., Zhu, X., Marsay, C. M., and Wenger, F.: Transport and reaction of iron and iron stable isotopes in glacial meltwaters on Svalbard near Kongsfjorden: From rivers to estuary to ocean, *Earth Planet. Sci. Lett.*, 424, 201–211, 2015.

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## 8 Tables

605 Table 1: Overview of sampling and sampled ranges of salinity, pH as well as O<sub>2</sub> concentration and saturation (in %, given in parenthesis) and concentrations of dissolved inorganic nitrogen (DIN = NO<sub>3</sub><sup>-</sup> + NO<sub>2</sub><sup>-</sup> + NH<sub>4</sub><sup>+</sup>), silicate (SiO<sub>2</sub>) and dissolved organic carbon (DOC). All concentrations are given in μmol L<sup>-1</sup>. na stands for not available and Stat. stands for sampling station. DOC data were taken from (Martin et al., 2018).

River	Date	# of Stat.	Range of					
			Salinity	pH	O <sub>2</sub>	DIN	SiO <sub>2</sub>	DOC
Rajang	20 – 27 Aug '16	30	0 – 32	6.5 – 8.1	85 – 153 (42 – 73)	6.7 – 29	4.0 – 179	na
	4 – 7 Mar '17	14	0 – 30	6.0 – 8.2	142 – 237 (58– 109)	8.1 – 18	16 – 158	96 – 201
	5 – 14 Sept '17	8	0 – 18	6.9 – 8.2	164 – 227 (76 – 90)	6.7 – 14	12 – 98	na
Maludam	9 Mar '17	9	0 – 20	3.7 – 7.6	34 – 213 (13 – 100)	3.9 – 10	5.8 – 32	266 – 4387
	14/15 Sept '17	9	0 – 15	4.1 – 6.7	43 – 155 (17 – 74)	2.1 – 3.0	0.1 – 8.0	3072 – 3245
Sebuyau	11 Mar '17	11	0 – 24	4.3 – 7.8	43 – 246 (18 – 116)	2.9 – 13	33 – 78	206 – 1968
	15 Sept '17	5	0 – 10	7.2 – 7.7	65 – 179 (27 – 75)	1.1 – 13	0.9 – 44	235 – 2052
Simunjan	12 Mar '17	6	0 – 0.4	4.7 – 6.3	31 – 81 (13 – 34)	2.2 – 16	73 – 114	2016 – 3039
	17 Sept '17	6	0 – 4.6	4.8 – 6.7	95 – 131 (39 – 53)	2.0 – 13	1.4 – 2.6	925 – 1960
Sematan	9 Mar '17	5	0 – 28	6.8 – 8.3	184 – 208 (81 – 102)	5.9 – 10	6.3 – 141	100 – 240
Samusam	11 Mar '17	5	0 – 27	6.3 – 8.2	174 – 208 (72 – 102)	3.9 – 6.6	9.7 – 98	87 – 1188

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Table 2: Summary of the environmental settings of the river basins. Based on the area percentage of oil palm, logging and our own surveys and observations, we classified the river basins into undisturbed (U) and disturbed (D). All areas are given in km<sup>2</sup>.

River	Areas				River water surface <sup>4</sup>	Remarks	Classification
	Total Basin	Peatland <sup>1</sup>	Oil palm plantations <sup>2</sup>	Logging <sup>3</sup>			
Rajang	50,000 <sup>5</sup>	3844	4514	29,379	455 <sup>5</sup>	The longest river in Malaysia. Major town is Sibul (163,000 population). Smaller townships are Kapit, Kanowit and Sarikei. There is a large number of villages and longhouses (traditional buildings inhabited by local communities) located along the river and its tributaries. Two hydroelectric power plants were built at two tributaries in the upper Rajang basin. The river mouth is surrounded by peat lands, and most of these peat lands have been converted to commercial oil palm plantations.	D
Maludam	197	172	16	0	0.36	The upstream of the river is surrounded by the Maludam National Park. The Maludam Peninsula is bordered by the Lupar and Saribas Rivers and is the biggest undisturbed peat forest in Malaysia. The National Park had been subjected to selective logging before it was gazetted as a totally protected area in 2000. Well preserved peat land. There are oil palm cultivations near the few villages.	U
Sebuyau	538	288	24	0	2.11	Major town is Sebuyau (14,000 population), surrounded by a few villages. Other agricultural activities were observed.	U
Simunjan	788	346	240	0	4.73	Major town is Simunjan (22,000 population), a few villages. Two streams combine to form the main Simunjan River. One of the streams passes an oil palm mill which discharges into the river.	D
Sematan	287	0	0	0	1.47	Major town is Sematan (7,600 population), small villages. We observed agricultural activities by the local people.	U
Samusam	163	0	0	0	0.85	Well preserved tropical forest. Some peat in the upper catchment area.	U

<sup>1</sup> Estimate is based on 'Wetlands International': 'Malaysia peat lands'. Accessed through Global Forest Watch on 22<sup>nd</sup> November 2018 ([www.globalforestwatch.org](http://www.globalforestwatch.org)).

<sup>2</sup> Estimate is based on 'Oil palm concessions'. Accessed through Global Forest Watch on 22<sup>nd</sup> November 2018 ([www.globalforestwatch.org](http://www.globalforestwatch.org)).

<sup>3</sup> Estimate is based on 'Managed forest concessions'. Accessed through Global Forest Watch on 22<sup>nd</sup> November 2018 ([www.globalforestwatch.org](http://www.globalforestwatch.org)).

<sup>4</sup> Area estimates are based on the length and width of the primary course and main tributaries of the rivers. Length and width of the rivers were estimated using Google Earth (multiple readings).

<sup>5</sup> Estimate from (Staub et al., 2000).



Table 3: Overview of N<sub>2</sub>O and CH<sub>4</sub> concentrations, saturations and flux densities in rivers and estuaries of NW Borneo.

River	Date	N <sub>2</sub> O			CH <sub>4</sub>		
		concentration nmol L <sup>-1</sup>	saturation %	flux density nmol m <sup>-2</sup> s <sup>-1</sup>	concentration nmol L <sup>-1</sup>	saturation %	flux density nmol m <sup>-2</sup> s <sup>-1</sup>
Rajang	Aug '16	2.0 – 14.1	28 – 215	-0.33 – 0.48	13.2 – 233	719 – 9988	0.77 – 15
	Mar '17	5.9 – 24.0	100 – 329	0 – 1.08	11.1 – 1008	455 – 40,598	0.34 – 62
	Sept '17	18.6 – 24.6	277 – 390	0.76 – 1.22	7.4 – 150	350 – 6019	0.35 – 9.05
Maludam	Mar '17	4.5 – 6.7	62 – 106	-0.20 – 0.03	312 – 829	12,603 – 32,988	19 – 50
	Sept '17	10.8 – 20.7	150 – 331	0.23 – 1.00	3.3 – 18	163 – 717	0.09 – 0.93
Sebuyau	Mar '17	3.5 – 7.7	55 – 118	-0.18 – 0.08	8.4 – 1228	396 – 50,774	0.41 – 78
	Sept '17	12.8 – 23.0	176 – 335	0.36 – 1.08	6.4 – 29	299 – 1285	0.28 – 1.79
Simunjan	Mar '17	2.5 – 41.4	35 – 570	-0.31 – 2.20	39 – 1372	1642 – 57,459	2.37 – 88
	Sept '17	5.1 – 26.5	73 – 365	-0.13 – 1.24	(14,999) <sup>1</sup>	(624,070) <sup>1</sup>	
Sematan	Mar '17	4.3 – 8.2	71 – 109	-0.11 – 0.04	2.5 – 21	106 – 878	0.01 – 1.18
Samusam	Mar '17	4.0 – 9.5	67 – 142	-0.13 – 0.19	8.6 – 12	433 – 47,055	0.43 – 72
					16.5 – 978	830 – 43,807	0.95 – 63

620 <sup>1</sup> This extreme value was not included in further computations.



Table 4: Overview of N<sub>2</sub>O and CH<sub>4</sub> flux densities from aqueous and soils ecosystems in SE Asia. (na stands for not available/not measured.)

Site	Location	N <sub>2</sub> O flux density, mmol m <sup>-2</sup> s <sup>-1</sup>		CH <sub>4</sub> flux density, mmol m <sup>-2</sup> s <sup>-1</sup>		Measurement or sampling dates	Reference
		Range	Mean <sup>†</sup>	Range	Mean <sup>†</sup>		
Aqueous systems							
Rajang River/Estuary	Sarawak, NW Borneo	-0.33 – 1.22	0.53	0.34 – 62	5.52	Aug. 2016; Mar.; Sept. 2017	This study
Maludam River/Estuary	Sarawak, NW Borneo	-0.20 – 1.00	0.32	0.09 – 50	15.9	March 2017; September 2017	
Sebuyau River/Estuary	Sarawak, NW Borneo	-0.18 – 1.08	0.39	0.28 – 78	15.4	March 2017; September 2017	
Sinunjan River/Estuary	Sarawak, NW Borneo	-0.31 – 2.20	0.50	0.01 – 88	18.7	March 2017; September 2017	
Sematan River/Estuary	Sarawak, NW Borneo	-0.11 – 0.04	-0.05	0.43 – 72	21.1	March 2017	
Samusum River/Estuary	Sarawak, NW Borneo	-0.13 – 0.19	0.05	0.95 – 63	21.7	March 2017	
Lupar River/Estuary	Sarawak, NW Borneo	0.04 – 0.04	0.04	0.59 – 0.84	0.72	June 2013; March 2014	(Müller et al., 2016a)
Saribas River/Estuary	Sarawak, NW Borneo	0.04 – 0.08	0.06	0.45 – 1.01	0.73	June 2013; March 2014	(Jauhainen and Silvennoinen, 2012)
Saribas River tributary	Sarawak, NW Borneo	0.37 – 0.39	0.38	0.81 – 4.84	2.83	June 2013; March 2014	
Drainage canal, Kalimantan, settled	Central Kalimantan, S Borneo	-0.02 – 0.03	0	0 – 943	119	September 2007; April 2008	
Drainage canal, Kampar, settled	Riau, eastern central Sumatra	0.03 – 5.80	0.73	0 – 3672	776	September 2007; April 2008	
Drainage canal, Kampar, disturbed	Riau, eastern central Sumatra	0.02 – 0.84	0.20	2.17 – 281	64.4	September 2007; April 2008	
Soil systems							
Forest	Sarawak, NW Borneo	-0.03 – 0.20	0.08	-0.10 – 0.19	0.04	August 2002 – July 2003	(Melling et al., 2005, 2007)
Sago plantation	Sarawak, NW Borneo	0.01 – 1.75	0.88	-0.17 – 2.36	1.10	August 2002 – July 2003	
Oil palm plantation	Sarawak, NW Borneo	0.01 – 0.58	0.29	-0.76 – 0.11	-0.33	August 2002 – July 2003	(Jauhainen et al., 2012)
Undrained forest	Central Kalimantan, S Borneo	-0.09 – 1.16	0.02	na	na	Dry/wet seasons in 2000/2001	
Drained forest	Central Kalimantan, S Borneo	-0.42 – 22.9	1.11	na	na	monitoring 2004 – 2007	
Drained recovering forest	Central Kalimantan, S Borneo	-0.06 – 0.45	0.02	na	na	Dry/wet seasons in 2001/2002	
Drained burned peat	Central Kalimantan, S Borneo	-0.70 – 0.88	0.11	na	na	Dry/wet seasons in 2001/2002; monitoring 2004 – 2007	
Agricultural peat in Kalampagan	Central Kalimantan, S Borneo	-0.95 – 0.89	0.12	na	na	Dry/wet seasons in 2001/2002	
Agricultural peat in Marang	Central Kalimantan, S Borneo	-0.86 – 0.59	0.07	na	na	Dry/wet seasons in 2001/2002	
Canopy soil of oil palm	Jarabi, eastern central Sumatra	na	0.001	na	0.0004	February 2013 – May 2014	
Drained agricultural land (fertilized)	Various locations in SE Asia	0.81 – 29.3	10.3	0.05 – 6.74	3.39	Various dates	(Allen et al., 2018) (Couvemberg et al., 2010); Review of results from various studies.
Drained open vegetation (abandoned, not fertilized)	Various locations in SE Asia	-0.12 – 0.45	0.08	na	na	Various dates	
Forested (drained and undrained peat swamp, agro-forestry)	Various locations in SE Asia	-0.06 – 1.51	0.39	-0.73 – 11.6	5.45	Various dates	
Rice paddies	Various locations in SE Asia	-0.04 – 0.23	0.07	7.17 – 98.1	52.7	Various dates	
Peat soil	Various locations in SE Asia	na	na	0 – 52.1	26.0	Various dates	
Maludam Natl. Park	Sarawak, NW Borneo	na	na	na	23.1	November – December 2013	(Tang et al., 2018)

<sup>†</sup> Values in italics indicate a mean flux density computed from the range given in the table (when no mean flux density was given in the ref.)



625 Table 5: Mean annual emissions of N<sub>2</sub>O and CH<sub>4</sub> from rivers and estuaries in NW Borneo. The data from Lupar and Saribas Rivers are from (Müller et al., 2016a).

River	Emissions	
	Gg N <sub>2</sub> O yr <sup>-1</sup>	Gg CH <sub>4</sub> yr <sup>-1</sup>
Rajang	0.33	1.27
Maludam	0.20	3.65
Sebuyau	0.24	3.53
Simunjan	0.32	4.30
Sematan	-0.03	5.99
Samusam	0.03	4.99
Lupar	0.01	0.08
Saribas	0.01	0.04
Sum	1.09	23.8



#### Figure Captions

Figure 1: Map of the study area with locations of the sampling stations. Sampling stations from  
630 August 2016 are displayed in red circles, from March 2017 in blue triangles, and from September  
2017 in green diamonds. Major cities are highlighted in bold plus symbols. Inset is adapted from  
(Staub et al., 2000).

Figure 2: N<sub>2</sub>O saturations along the salinity gradients of (a) Rajang, (b) Maludam, (c) Sebuyau, (d)  
Simutan, (d) Sematan and (e) Samusam. The dashed lines indicate the equilibrium (100%) saturation.  
635 The open cycles depict measurements from August 2016, the filled red cycles depict measurements  
from March 2017 and the filled blue cycles depict measurements from September 2017.

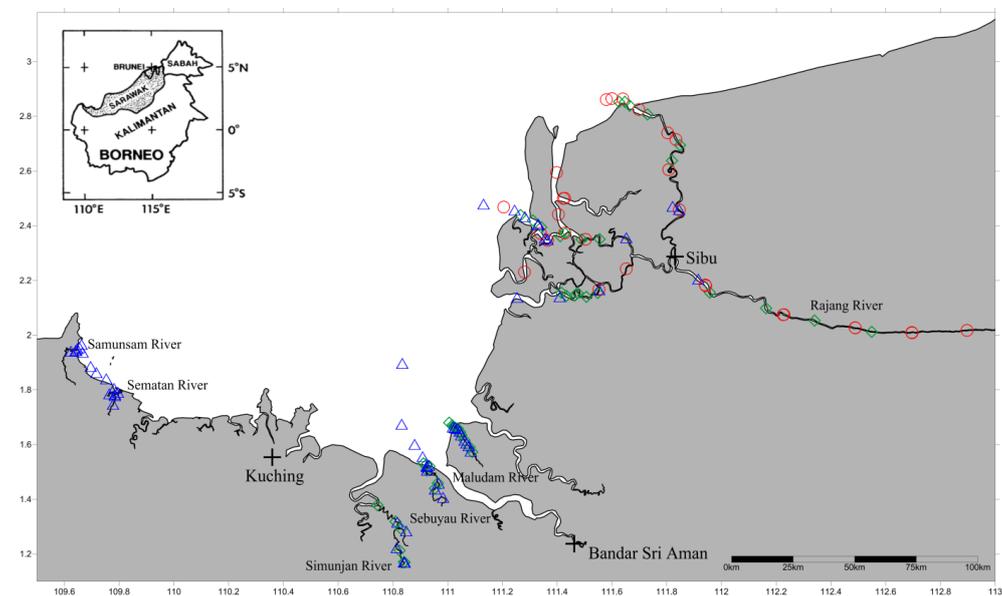
Figure 3: CH<sub>4</sub> saturations along the salinity gradients of (a) Rajang, (b) Maludam, (c) Sebuyau, (d)  
Simutan, (d) Sematan and (e) Samusam. The dashed lines indicate the equilibrium (100%) saturation.  
640 The open cycles depict measurements from August 2016, the filled red cycles depict measurements  
from March 2017 and the filled blue cycles depict measurements from September 2017.

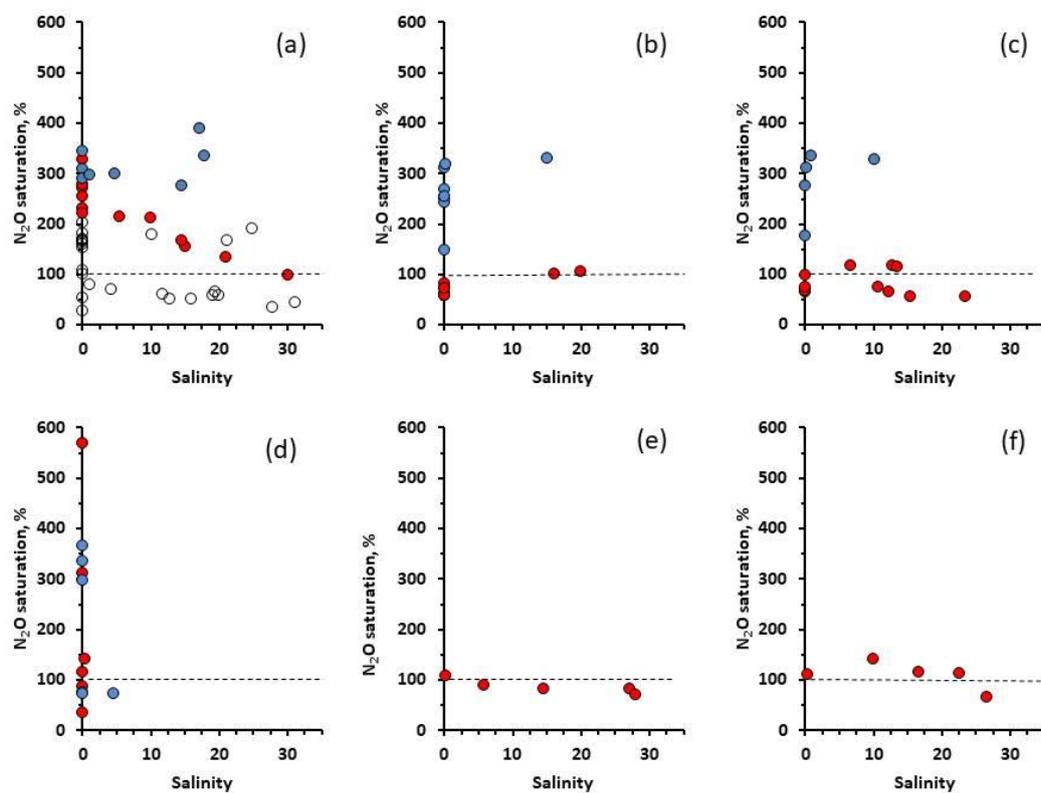
Figure 4: (a) Mean N<sub>2</sub>O and (b) mean CH<sub>4</sub> concentrations for the individual rivers vs. the mean  
monthly rainfall amount during the month of the sampling. We also included the mean N<sub>2</sub>O and CH<sub>4</sub>  
concentration for the Lupar, Saribas Rivers and Saribas tributary from (Müller et al., 2016a). The  
linear correlation in (a) is described by  $y = 0.08x + 5.76$  ( $r = 0.72$ ,  $n = 17$ , significant at the 99% level).  
645 The linear correlation in (b) is described by  $y = -9.57x + 713.15$  ( $r = 0.88$ ,  $n = 13$ , significant at the  
99% level; please note that the encircled data were not included in the correlation).



650 6 Figures

Figure 1.





655 Figure 2.

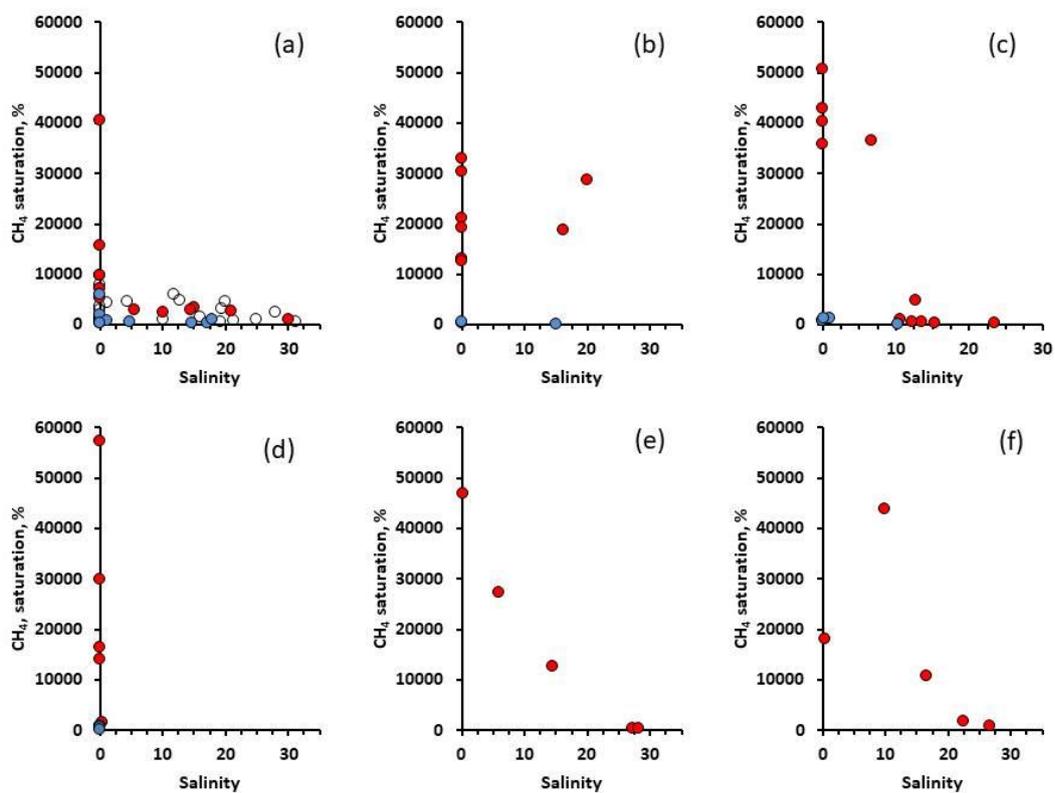


Figure 3.



Figure 4

