Greenhouse gas fluxes in mangrove forest soil in an Amazon estuary

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- 12 | Abstract: Tropical mangrove forests are important carbon sinks, the soil being the
- main carbon reservoir. Understanding the variability and the key factors that control
- 14 fluxes is critical to accounting for greenhouse gas (GHG) emissions, particularly in the
- 15 current scenario of global climate change. This study is the first to quantify carbon
- dioxide (CO₂) and methane (CH₄) emissions using a dynamic chamber in a natural
- 17 mangrove soil of the Amazon. The plots for the trace gases study were allocated at
- 18 contrasting topographic heights. The results showed that the mangrove soil of the
- Amazon estuary is a source of CO_2 (6.66 g CO_2 m⁻² d⁻¹) and CH_4 (0.13 g CH_4 m⁻² d⁻¹) to
- 20 the atmosphere. The CO₂ flux was higher in the high topography (7.86 g CO₂ m⁻² d⁻¹)
- 21 than in the low topography (4.73 g CO_2 m⁻² d⁻¹) in the rainy season, and CH_4 was higher
- 22 in the low topography (0.13 g CH₄ m⁻² d⁻¹) than in the high topography (0.01 g CH₄ m⁻²
- 23 d⁻¹) in the dry season. However, in the dry period, the low topography soil produced
- 24 more CH₄. Soil organic matter, carbon and nitrogen ratio (C/N), and redox potential
- 25 influenced the annual and seasonal variation of CO₂ emissions; however, they did not
- 26 affect CH₄ fluxes. The mangrove soil of the Amazon estuary produced 35.40 Mg CO_{2-eq}
- 27 ha⁻¹ y⁻¹. A total of 2.16 kg CO₂ m⁻² y⁻¹ needs to be sequestered by the mangrove
- 28 ecosystem to counterbalance CH₄ emissions.

1 Introduction

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- 30 Mangrove areas are estimated to be the main contributors to greenhouse gas emissions
- 31 in marine ecosystems (Allen et al., 2011; Chen et al., 2012). However, mangrove forests
- 32 are highly productive due to a high nutrient turnover rate (Robertson et al., 1992) and
- 33 have mechanisms that maximize carbon gain and minimize water loss through plant
- 34 transpiration (Alongi and Mukhopadhyay, 2015). A study conducted in 25 mangrove
- 35 forests (between 30° latitude and 73° longitude) revealed that these forests are the

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- 36 richest in carbon (C) storage in the tropics, containing on average 1,023 Mg C ha⁻¹ of
- which 49 to 98% is present in the soil (Donato et al., 2011).
- 38 The estimated soil CO₂ flux in tropical estuarine areas is 16.2 Tg C y⁻¹ (Alongi, 2009).
- 39 However, soil efflux measurements from tropical mangroves revealed emissions
- 40 ranging from 2.9 to 11.0 g CO₂ m⁻² d⁻¹ (Castillo et al., 2017; Chen et al., 2014; Shiau
- 41 and Chiu, 2020). In situ CO₂ production is related to the water input of terrestrial,
- 42 riparian, and groundwater brought by rainfall (Rosentreter et al., 2018b). Due to the
- 43 periodic tidal movement, the mangrove ecosystem is daily flooded, leaving the soil
- anoxic and consequently reduced, favoring methanogenesis (Dutta et al., 2013). Thus,
- estuaries are considered hotspots for CH₄ production and emission (Bastviken et al.,
- 46 2011; Borges et al., 2015). Organic material decomposition by methanogenic bacteria in
- 47 anoxic environments, such as sediments, inner suspended particles, zooplankton gut
- 48 (Reeburgh, 2007; Valentine, 2011), and the impact of freshwater should change the
- 49 electron flow from sulfate-reducing bacteria to methanogenesis (Purvaja et al., 2004),
- 50 which also results in CH₄ formation. On the other hand, high salinity levels, above 18
- 51 ppt, may result in an absence of CH₄ emissions (Poffenbarger et al., 2011), since CH₄
- 52 dissolved in pores is typically oxidized anaerobically by sulfate (Chuang et al., 2016).
- 53 Currently the uncertainty in emitted CH₄ values in vegetated coastal wetlands is
- 54 approximately 30% (EPA, 2017). Soil flux measurements from tropical mangroves
- revealed emissions range from 0.3 to 4.4 mg CH₄ m⁻² d⁻¹ (Castillo et al., 2017; Chen et
- 56 al., 2014; Kreuzwieser et al., 2003).
- 57 The production of greenhouse gases from soils is mainly driven by biogeochemical
- 58 processes. Microbial activities and gas production are related to soil properties,
- 59 including total carbon and nitrogen concentrations, moisture, porosity, salinity, and
- 60 redox potential (Bouillon et al., 2008; Chen et al., 2012). Due to the dynamics of tidal
- 61 movements, mangrove soils may become saturated and present reduced oxygen
- 62 availability, or suffer total aeration caused by the ebb tide. Studies attribute soil carbon
- 63 flux responses to moisture perturbations because of seasonality and flooding events
- 64 (Banerjee et al., 2016), with fluxes being dependent on tidal extremes (high tide and low
- 65 tide), and flood duration (Chowdhury et al., 2018). In addition, phenolic compounds
- 66 inhibit microbial activity and help keep organic carbon intact, thus leading to the
- 67 accumulation of organic matter in mangrove forest soils (Friesen et al., 2018).

- 68 The Amazonian coastal areas in the State of Pará (Brazil) cover 2,176.8 km² where
- 69 mangroves develop under the macro-tide regime (Souza Filho, 2005), representing
- 70 approximately 85% of the entire area of Brazilian mangroves (Herz, 1991). The
- objective of this study is to investigate the monthly flux of CO₂ and CH₄ from the soil,
- at two topographic heights, in a pristine mangrove area in the Mojuim River Estuary,
- belonging to the Amazon biome. The gas fluxes were studied together with the analysis
- of the vegetation structure and soil physical-chemical parameters.

2 Material and Methods

76 2.1 Study site

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- 77 This study was conducted in the Amazonian coastal zone, Macaca Island (-0.746491
- 78 latitude and -47.997219 longitude), located in the Mojuim River estuary, at the
- 79 Mocapajuba Marine Extractive Reserve, municipality of São Caetano de Odivelas
- 80 (Figure Fig. 1), state of Pará (Brazil). The Macaca island has an area of 1,322 ha of
- 81 pristine mangroves, and belongs to a mangrove area of 2,177 km² in the state of Pará
- 82 (Souza Filho, 2005). The climate is type Am (tropical monsoon) according to the
- 83 Köppen classification (Peel et al., 2007). The climatological data were obtained from
- 84 the Meteorological Database for Teaching and Research of the National Institute of
- 85 Meteorology (INMET). The area has a rainy season from January to June (2,296 mm of
- 86 precipitation) and a dry season from July to December (687 mm). March and April were
- 87 the rainiest months with 505 and 453 mm of precipitation, while October and November
- 88 were the driest (53 and 61 mm, respectively). The minimum temperatures occur in the
- 89 rainy period (26 °C) and the maximum in the dry period (29 °C). The Mojuim estuary
- 90 has a macrotidal regime, with an average amplitude of 4.9 m during spring tide and 3.2
- 91 m during low tide (Rollnic et al., 2018). During the wet season the Mojuim River has a
- 92 flow velocity of 1.8 m s⁻¹ at the ebb tide and 1.3 m s⁻¹ at the flood tide, whereas in the
- 93 dry season, the maximum currents reach 1.9 m s⁻¹ at the flood and 1.67 m s⁻¹ at the ebb
- 94 tide (Rocha, 2015). The annual mean salinity of the river water is 26.95 PSU (Valentim
- 95 et al., 2018).

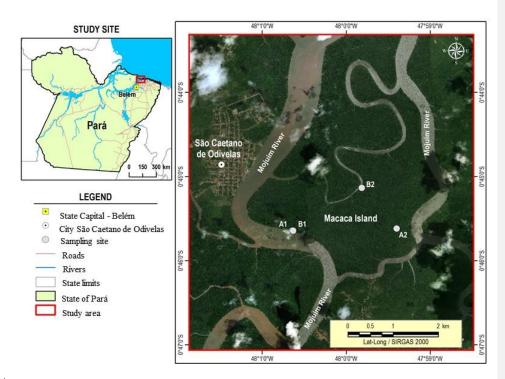


Figure 1. The Macaca Island located in the mangrove coast of Northern Brazil, Municipality of São Caetano de Odivelas (state of Pará), with sampling points at low (plot B1 and plot B2) and high (plot A1 and plot A2) topographies. Image Source: © Google Earth

The Mojuim River region is geomorphologically formed by partially submerged river basins consequent of the increase in the relative sea level during the Holocene (Prost et al., 2001) associated with the formation of mangroves, dunes, and beaches (El-Robrini et al., 2006). Before reaching the estuary, the Mojuim River crosses an area of a dryland forest highly fragmented by family farming, forming remnants of secondary forest (< 5.0 ha) of various ages (Fernandes and Pimentel, 2019). The population economically exploited the estuary, primarily by artisanal fishing, crab (*Ucides cordatus* L.) extraction, and oyster farms.

The flora of the mangrove area of Macaca Island is little anthropized and comprises the plant genera *Rhizophora*, *Avicenia*, *Laguncularia*, and *Acrostichum* (Ferreira, 2017; França et al., 2016). The estuarine plains are influenced by macrotide dynamics and can be physiographically divided into four sectors according to the different vegetation covers, associated with the landforms distribution, topographic gradient, tidal

- inundation, and levels of anthropic transformation(França et al., 2016). The Macaca
- 115 Island is ranked as being from the fourth sector, which implies having woods of adult
- trees of the genus Ryzophora with an average height of 10 to 25 m, is located at an
- elevation of 0 to 5 m, and having silt-clay soil (França et al., 2016).
- 118 Four sampling plots were selected in the Macaca Island (Figure Fig. 1) on 19/05/2017,
- 119 when the moon was in the waning quarter phase: two plots where flooding occurs every
- 120 day (plots B1 and B2; Figure-Fig. 1), called low topography (Top_Low), and two plots
- where flooding occurs only at high tides during the solstice and on the high tides of the
- 122 | rainy season of the new and full moons (plots A1 and A2; Figure Fig. 1), called high
- topography (Top_High).

2.2 Greenhouse gas flux measurements

125 In each plot, eight Polyvinyl Chloride rings with 0.20 m diameter and 0.12 m height were randomly installed within a circumference with a diameter of 20 m. The rings had 126 127 an area of 0.028 m⁻² (volume of 3.47 L), were fixed 0.05 m into the ground, and 128 remained in place until the study was completed. Once a month, gas fluxes were 129 measured during periods of waning or crescent moon, as these are the times when the 130 soil in the low topography is more exposed. To avoid the influence of mangrove roots 131 on the gas fluxes, the rings were placed in locations without any seedlings or 132 aboveground mangrove roots. The CO₂ and CH₄ concentrations (ppm) were measured 133 using the dynamic chamber methodology (Norman et al., 1997; Verchot et al., 2000), 134 sequentially connected to a Los Gatos Research portable gas analyzer (Mahesh et al., 135 2015). The device was calibrated monthly with a high quality standard gas (500 ppm 136 CO₂; 5 ppm CH₄). The rings were sequentially closed for three minutes with a PVC cap, 137 being connected to the analyzer through two 12.0 m polyethylene hoses. The gas 138 concentration was measured every two seconds and automatically stored by the 139 analyzer. CO2 and CH4 fluxes were calculated from the linear regression of 140 increasing/decreasing CO₂ and CH₄ concentrations within the chamber, usually between 141 one and three minutes after the ring cover was placed (Frankignoulle, 1988; McEwing et al., 2015). The flux is considered zero when the linear regression reaches an R² < 142 143 0.30 (Sundqvist et al., 2014). However, in our analyses, most regressions reached R^2 > 144 0.70, and the regressions were weak and considered zero in only 6% of the samples. At 145 the end of each flux measurement, the height of the ring above ground was measured at four equidistant points with a ruler. The seasonal data were analyzed by comparing the

average monthly fluxes in the wet season and dry season separately.

2.3 Vegetation structure and biomass

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- The floristic survey was conducted in October 2017 using circular 1,256.6 m² plots
- 150 (Kauffman et al., 2013) divided into four 314.15 m² subplots, which is the equivalent to
- 151 0.38 ha, at the same topographies as the gas flux analysis (Figure Fig. 1). We recorded
- the diameter above the aerial roots, the diameter of the stem, and total height of all trees
- with DBH (diameter at breast height; m) greater than 0.05m. The allometric equations
- 154 (Howard et al., 2014) to calculate tree biomass (aboveground biomass; AGB) were:
- AGB = $0.1282 * DBH^{2.6}$ (R² = 0.92) for *R. mangle*; AGB = $0.140 * DBH^{2.4}$ (R² = 0.97)
- 156 for A. germinans; and Total AGB = $0.168 * \rho * DBH^{2.47}$ (R² = 0.99), where $\rho_{R. mangle}$ =
- 157 0.87; $\rho_{A. \; germinans} = 0.72 \; (\rho = wood \; density).$

2.4 Soil sampling and environmental characterization

- Four soil samples were collected with an auger at a depth of 0.10 m in all the studied
- plots for gas flux measurements (Figure-Fig. 1) in July 2017 (beginning of the dry
- season) and January 2018 (beginning of the rainy season). Before the soil samples were
- 162 removed, pH and redox potential (Eh; mV) were measured with a Metrohm 744
- equipment by inserting the platinum probe directly into the intact soil at a depth of 0.10
- m (Bauza et al., 2002). The soil samples collected in the field were transported to the
- laboratory (Chemical Analysis Laboratory of the Museu Paraense Emílio Goeldi) in
- thermal boxes containing ice. The soil samples were analyzed on the day after collection
- at the laboratory, and the samples were kept in a freezer. Salinity (Sal; ppt) was
- 168 measured with PCE-0100, and soil moisture (Sm; %) by the residual gravimetric
- 169 method (EMBRAPA, 1997).
- Organic Matter (OM; g kg⁻¹), Total Carbon (T_C; g kg⁻¹) and Total Nitrogen (T_N; g kg⁻¹)
- were calculated by volumetry (oxidoreduction) using the Walkley-Black method
- 172 (Kalembasa and Jenkinson, 1973). Microbial carbon (C_{mic}; mg kg⁻¹) and microbial
- nitrogen (N_{mic}; mg kg⁻¹) were determined through the 2.0 min of Irradiation-extraction
- method of soil by microwave technique (Islam and Weil, 1998). Microwave heated soil
- extraction proved to be a simple, fast, accurate, reliable, and safe method to measure
- soil microbial biomass (Araujo, 2010; Ferreira et al., 1999; Monz et al., 1991). The C_{mic}
- was determined by dichromate oxidation (Kalembasa and Jenkinson, 1973; Vance et al.,

- 178 1987). The N_{mic} was analyzed following the method described by Brookes et al. (1985),
- 179 changing fumigation to irradiation, which uses the difference between the amount of T_N
- 180 in irradiated and non-irradiated soil. We used the flux conversion factor of 0.33
- 181 (Sparling and West, 1988) and 0.54 (Almeida et al., 2019; Brookes et al., 1985), for
- 182 carbon and nitrogen, respectively. Particle size analysis was performed separately on
- 183 four soil samples collected at each flux plot, in the two seasons (October 2017 and
- 184 March 2018), according to EMBRAPA (1997).
- 185 At each gas flux measurement, environmental variables such as air temperature (Tair,
- 186 °C), relative humidity (RH, %), and wind speed (W_s, m s⁻¹) were quantified with a
- portable thermo-hygrometer (model AK821) at the height of 2.0 m above the soil
- surface. Soil temperature (T_s, °C) was measured with a portable digital thermometer
- (model TP101) after each gas flux measurement. Daily precipitation was obtained from
- an automatic precipitation station installed at a pier on the banks of the Mojuim River in
- 191 São Caetano das Odivelas (coordinates: -0.738333 latitude; -48.013056 longitude).

192 2.5 Statistical analyses

- 193 On the Macaca Island, two treatments were allocated (low and high topography), with
- 194 two plots in either treatment. In each plot, eight chambers were randomly distributed,
- which were considered sample repetitions. The normality of the data of CH₄ and FCO₂
- 196 flux, and soil physicochemical parameters was evaluated using the Shapiro-Wilks
- 197 method. The soil CO₂ and CH₄ flux showed a non-normal distribution. Therefore, we
- 198 used the non-parametric ANOVA (Kruskal-Wallis, p < 0.05) to test the differences
- 199 between the two treatments among months and seasons. The physicochemical
- 200 parameters were normally distributed. Therefore, a parametric ANOVA was used to test
- 201 the statistical differences (p < 0.05) between the two treatments among months and
- 202 seasons. Pearson correlation coefficients were calculated to determine the relationships
- between soil properties and gas fluxes in the months (dry and wet season) when the
- 204 chemical properties of the soil were analyzed at the same time as gas fluxes were
- 205 measured. Statistical analyses were performed with the free statistical software Infostat
- 206 2015®.

3 Results

3.1 Carbon dioxide and methane fluxes

 CO_2 fluxes differed significantly between topographies only in January (H = 3.915; p = 0.048), July (H = 9.091; p = 0.003), and November (H = 11.294; p < 0.001) (Figure Fig. 2; Supplementary Information, SI 1), with generally higher fluxes at the high topography than at the low topography. At the high topography, CO_2 fluxes were significantly higher (H = 24.510; p = 0.011) in July compared to August and December, March, October, and May, not differing from the other months of the year. Similarly, at the low topography, CO_2 fluxes were statistically significantly higher (H = 19.912; p = 0.046) in September and February than inwhen compared to January and November, not differing from the other months. We found a mean monthly flux of 7.9 ± 0.7 g CO_2 m⁻² d⁻¹ (mean \pm standard error) and 5.4 ± 0.5 g CO_2 m⁻² d⁻¹ at the high and low topographies, respectively.



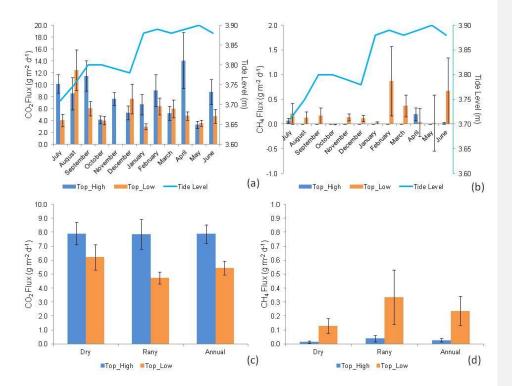


Figure 2. CO₂ (a) and CH₄ (b) fluxes (g CO₂ or CH₄ m⁻² d⁻¹) monthly (July 2018 to June 2019) (n = 16). Seasonal (Dry and Rainy) and annual fluxes of CO₂ (c) and CH₄

- 224 (d), at high (Top_High) and low (Top_Low) topographies (n = 96), in a mangrove forest
- soil compared to tide level (Tide Level). The bars represent the standard error of the
- 226 mean.
- 227 The CH₄ fluxes were statistically different between topographies only in November (H
- = 9.276; p = 0.002) and December (H = 4.945; p = 0.005), with higher fluxes at the low
- topography (Figure Fig. 2; SI 1). At the high topography, CH₄ fluxes were significantly
- (H = 40.073; p < 0.001) higher in April and July compared to the other months studied,
- 231 and in November CH₄ was consumed from the atmosphere (Figure Fig. 2; SI 1).
- Similarly, CH₄ fluxes at the low topography did not vary significantly among months
- 233 (H = 10.114; p = 0.407).
- Greenhouse gas fluxes (Figure Fig. 2) were only significantly different between
- topographies in the dry season (Figure Fig. 3), period when CO₂ fluxes were higher (H
- = 7.378; p = 0.006) at the high topography and CH₄ fluxes at the low topography (H =
- 8.229; p < 0.001). In the Macaca Island, the mean annual fluxes of CO_2 and CH_4 were
- 238 $6.659 \pm 0.419 \text{ g CO}_2 \text{ m}^{-2} \text{ d}^{-1}$ and $0.132 \pm 0.053 \text{ g CH}_4 \text{ m}^{-2} \text{ d}^{-1}$, respectively. During the
- study year, the CO₂ flux from the mangrove soil ranged from -5.06 to 68.96 g CO₂ m⁻²
- 240 d^{-1} (mean 6.66 g CO₂ m⁻² d⁻¹), while the CH₄ flux ranged from -5.07 to 11.08 g CH₄ m⁻²
- 241 | d^{-1} (mean 0.13 g CH₄ m⁻² d^{-1}), resulting in a total carbon <u>efflux</u> rate of 1.92 g C m⁻² d^{-1}
- 242 or 7.00 Mg C ha⁻¹ y⁻¹ (Figure Fig. 2).

3.2 Weather data

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- 244 There was a marked seasonality during the study period (Figure Fig. 2), with 2,155.0
- 245 mm of precipitation during the rainy period and 1,016.5 mm during the dry period. The
- highest tides occurred in the period of greater precipitation (Figure Fig. 3) due to the
- rains. However, the rainfall distribution was different from the climatological normal
- 248 (Figure Fig. 3). The precipitation in the rainy season was 553.2 mm below and in the
- 249 dry season was 589.1 mm above the climatological normal. Thus, in the period studied,
- 250 the dry season was rainier and the rainy season drier than the climatological normal,
- which may be a consequence of the La Niña event (Wang et al., 2019).

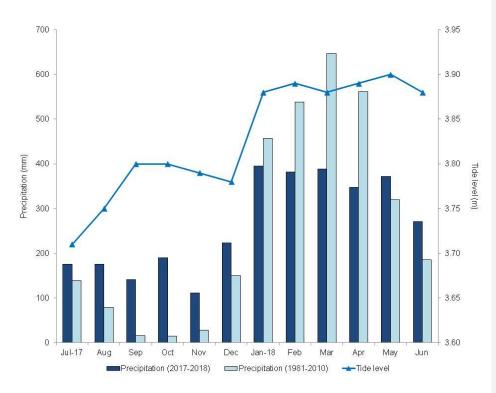


Figure 3. Monthly climatological normal in the municipality of Soure (1981-2010, mm), monthly precipitation (mm), and maximum tide height (m) from 2017 to 2018, in the municipality of São Caetano de Odivelas (PA).

 T_{air} was significantly higher (LSD = 0.72, p = 0.01) at the high (31.24 \pm 0.26 °C) than at the low topography (30.30 \pm 0.25 °C) only in the rainy season (Figure Fig. 4a). No significant variation in T_s was found between topographies in either season (Figure Fig. 4b). RH was significantly higher (LSD = 2.55, p = 0.01) at the high topography (70.54 \pm 0.97%) than at the low topography (66.85 \pm 0.87%) only in the rainy season (Figure Fig. 4c). W_s (Figure Fig. 4d) was significantly higher (LSD = 0.15, p < 0.00) at the low (0.54 \pm 0.06 m s⁻¹) than at the high topography (0.24 \pm 0.04 m s⁻¹) also in the rainy season.

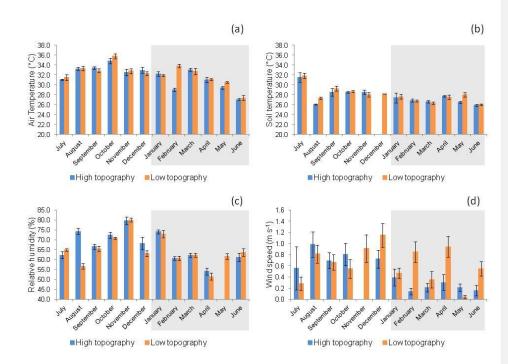


Figure 4. a) Air temperature (°C), b) soil temperature (°C), c) relative humidity (%), and d) wind speed (m s⁻¹) at high and low topographies, from July 2017 to June 2018 in a mangrove area in the Mojuim River estuary. Bars highlighted in grey correspond to the rainy season (n = 16). The bars represent the standard error.

3.3 Soil characteristics

Silt concentration was higher at the low topography (LSD: 14.763; p=0.007) and clay concentration was higher at the high topography plots (LSD: 12.463; p=0.005), in both seasons studied (Table 1). Soil particle size analysis did not differ statistically (p>0.05) between the two seasons (Table 1). Soil moisture did not vary significantly (p>0.05) between topographies at each season, or between seasonal periods at the same topography (Table 1). The pH varied statistically (LSD: 5.950; p=0.006) only at the low topography when the two seasons were compared, being more acidic in the dry period (Table 1). The pH values were significantly (LSD: 0.559; p=0.008) higher in the dry season (Table 1). No variation in Eh was identified between topographies and seasons (Table 1), although it was higher in the dry season than in the rainy season. However, Sal values were higher (LSD: 3.444; p=0.010) at the high topography than at the low topography in the dry season (Table 1). In addition, Sal was significantly higher

- in the dry season than in the rainy season, in both high (LSD: 2.916; p < 0.001) and low
- 284 (LSD: 3.003; p < 0.001) topographies (Table 1).

286 hig 287 err 288 sea

Table 1. Analysis of Sand (%), Silt (%), Clay (%), Moisture (%), pH, Redox Potential (Eh, mV) and salinity (Sal; ppt) in the mangrove soil of high and low topographies, and in the rainy and dry seasons (Macaca island, São Caetano das Odivelas). Numbers represent the mean ± standard error of the mean. Lower case letters compare topographies in each seasonal period and upper-case letters compare the same topography between seasonal periods. Different letters indicate statistical difference (LSD, p < 0.05).

Season	Topography	Sand	Silt	Clay	Moisture	**	Eh	Sal
		(%)	(%)	(%)	(%)	pН	(mV)	(ppt)
Dry	High	12.1±1.4 ^{aA}	41.8±3.3 ^{bA}	46.1±2.6 ^{aA}	73.1±6.6 ^{aA}	5.5±0.2 ^{aA}	190.25±45.53 ^{aA}	35.25±1.11 ^{aA}
	Low	$9.7{\pm}2.5^{aA}$	63.6 ± 6.1^{aA}	26.6 ± 5.2^{bA}	86.9±3.4 ^{aA}	5.3 ± 0.3^{aA}	106.38±53.76 ^{aA}	30.13±1.16 ^{bA}
	Mean	10.9±1.4 ^A	52.7±4.4 ^A	36.4±3.8 ^A	80.0±4.0 ^A	5.4±0.2 ^A	148.31±35.71 ^A	32.69±1.02 ^A
Rainy	High	12.3±1.0 ^{aA}	39.3±2.1 ^{bA}	48.4±1.6 ^{aA}	88.9±3.5 ^{aA}	4.9±0.4 ^{aA}	92.50±56.20 ^{aA}	7.50±0.78 ^{aB}
	Low	7.8 ± 1.4^{bA}	63.4 ± 5.2^{aA}	28.8 ± 4.2^{bA}	88.6±3.7 ^{aA}	$4.4{\pm}0.1^{aB}$	36.25±49.97 ^{aA}	8.13±0.79 ^{aB}
	Mean	10.1±1.1 ^A	51.4±4.1 ^A	38.6±3.4 ^A	88.7±2.5 ^A	4.6±0.2 ^B	64.38±37.04 ^A	7.81±0.54 ^B

290 The C_{mic} did not differ between topographies in the two seasons (Table 2). However, T_C 291 was significantly higher in the low topography in the dry season (LSD: 5.589; p < 292 0.000) and in the rainy season (LSD: 5.777; p=0.024). In addition, C_{mic} was higher in 293 the dry season in both the high (LSD: 11.325; p < 0.010) and low (LSD: 9.345; p < 294 0.000) topographies (Table 2). N_{mic} did not vary between topographies seasonally. 295 However, N_{mic} in the high (LSD: 9.059; p = 0.013) and low topographies (LSD: 4.447; 296 p = 0.001) was higher during the dry season (Table 2). The C/N ratio (Table 2) was 297 higher in the low than in the high topography in both the dry (LSD: 3.142; p < 0.000) 298 and rainy seasons (LSD: 3.675; p = 0.033). However, only in the low topography was 299 the C/N ratio higher (LSD: 1.863; p < 0.000) in the dry season than in the rainy season 300 (Table 2). Soil OM was higher at the low topography in the rainy (LSD: 9.950; p = 301 0.024) and in the dry seasons (LSD: 9.630; p < 0.000). Only in the lowland topography 302 was the OM concentration higher in the dry season than in the rainy season (Table 2).

Table 2. Seasonal and topographic variation in microbial Carbon (C_{mic} ; mg kg⁻¹), microbial Nitrogen (N_{mic} , mg kg⁻¹), Total Carbon (T_C ; g kg⁻¹), Total Nitrogen (N_T ; g kg⁻¹), Carbon/Nitrogen ratio (C/N) and Soil Organic Matter (OM; g kg⁻¹). Numbers represent the mean (\pm standard error). Lower case letters compare topographies at each season, and upper-case letters compare the topography between seasons.

Season		$C_{ m mic}$	$N_{ m mic}$	$T_{\rm C}$	T_N		OM	
	Topography	mg kg ⁻¹	mg kg ⁻¹	g kg ⁻¹	g kg ⁻¹	C/N	g kg ⁻¹	
	High	22.12±5.22 ^{aA}	12.76±4.20 ^{aA}	14.12±2.23 ^{bA}	1.43±0.06 ^{aA}	9.60±1.20 ^{bA}	24.35±3.84 ^{bA}	
Dry	Low	26.34±4.23 ^{aA} 10.34±2.05 ^{aA}		26.44±1.35 ^{aA} 1.56±0.04 ^a		16.98±0.84 ^{aA}	45.59±2.32 ^{aA}	
	Mean	24.23±3.29 A	11.55±2.28 ^A	20.28 ±2.03 ^A	1.49±0.04 A	13.29±1.19 ^A	34.97±3.50 A	
Rainy	High	7.40±0.79 ^{aB}	0.75±0.41 ^{aB}	11.46±2.48 ^{bA}	1.32±0.04 ^{aA}	8.42±1.70 ^{bA}	19.75±4.27 ^{bA}	
	Low	5.95 ± 1.06^{aB}	1.23 ± 0.28^{aB}	18.27 ± 1.06^{aB}	1.46 ± 0.06^{aA}	12.47 ± 0.22^{aB}	31.51 ± 1.83^{aB}	
	Mean	6.68±0.67 ^B	0.99±0.25 ^B	14.86 ±1.57 ^B	1.39±0.04 A	10.44±0.98 ^A	25.63±2.71 ^B	

3.4 Vegetation structure and biomass

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Only the species R. mangle and A. germinans were found in the floristic survey carried out. The DBH did not vary significantly between the topographies for either species (Table 3). However, R. mangle had a higher DBH than A. germinaris at both high (LSD: 139.304; p = 0.037) and low topographies (LSD: 131.307; p = 0.001). The basal area (BA) and AGB did not show significant variation (Table 3). A total aboveground biomass of 322.1 ± 49.6 Mg ha⁻¹ was estimated.

- ·		N ha ⁻¹	DBH	BA	AGB	
Specie	Topography		(cm)	$(m^2 ha^{-1})$	(Mg ha ⁻¹)	
Rhizophora	High	302.4±20.5	238.8±24.9 ^{aA}	17.3±2.0 ^{aA}	219.3±25.7 ^{aA}	
mangle	Low	310.4±37.6	283.5±45.0 ^{aA}	24.2±4.3 ^{aA}	338.7±62.9 ^{aA}	
Avicennia	High	47.7±20.5	86.8±51.2 ^{aB}	13.8±9.2 ^{aA}	135.3±94.7 ^{aA}	
germinans	Low	15.9±9.2	46.1 ± 29.3^{aB}	11.8±8.8 ^{aA}	136.0±108.3 ^{aA}	
T-4-1	High	350.2±18.4	325.6±33.6 ^a	31.1±7.5 ^a	304.5±99.8 ^a	
Total	Low	346.2±41.0	296.0±23.7 ^a	30.0±4.1 ^a	330.8 ± 60.4^{a}	

The equations for biomass estimates (AGB) were: $R. mangle = 0.1282*DBH^{2.6}$; $A. germinans = 0.14*DBH^{2.4}$; and $Total = 0.168*\rho*DBH^{2.47}$, where $\rho_{R. mangle} = 0.87$; $\rho_{A. germinans} = 0.72$ (Howard et al., 2014).

3.5 Drivers of greenhouse gas fluxes

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321 In the rainy season, CO_2 efflux was correlated with T_{air} (Pearson = 0.23, p = 0.03), RH 322 (Pearson = -0.32, p < 0.00) and T_s (Pearson = 0.21, p = 0.04) only at the low 323 topography. In the dry season CO₂ flux was correlated with T_s (Pearson = 0.39, p < 324 0.00) at the low topography. The dry season was the period in which we found the 325 greatest amount of significant correlations between CO2 efflux and soil chemical 326 parameters, while the C:N ratio, OM, and Eh were correlated with CO2 efflux in both 327 seasons (Table 4). The negative correlation between T_C, N_T, C/N, and OM, along with 328 the positive correlation of N_{mic} with soil CO₂ flux, in the dry period, indicates that 329 microbial activity is a decisive factor for CO2 efflux (Table 4). Soil moisture in the 330 Mojuim River mangrove forest negatively influenced CO2 flux in both seasons (Table 331 4). However, soil moisture was not correlated with CH₄ flux. No significant correlations 332 were found between CH₄ efflux and the chemical properties of the soil in the mangrove 333 of the Mojuim River estuary (Table 4).

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Table 4. Correlation coefficient (Pearson) of CO₂ and CH₄ fluxes with chemical parameters of the soil in a mangrove area in the Mojuim River estuary.

Gas Flux	Season	T_{C}	T_N	C_{mic}	N_{mic}	CAL	OM	Sal	Eh	11	Moisture
$(g m^{-2} d^{-1})$		$(g kg^{-1})$	$(g kg^{-1})$	(mg kg ⁻¹)	(mg kg ⁻¹)	C/N	$(g kg^{-1})$	(ppt)	(mV)	pН	(%)
	Dry	-0.68**	-0.59*	0.18 ^{NS}	0.61**	-0.66**	-0.67**	-0.07 ^{NS}	0.51*	0.21 ^{NS}	-0.49*
CO_2	Rainy	-0.44 ^{NS}	-0.20 ^{NS}	-0.15 ^{NS}	-0.32 ^{NS}	-0.50*	-0.63**	-0.54*	0.53*	0.47^{NS}	-0.54*
	Annual	-0.50**	-0.35*	-0.18 ^{NS}	0.00^{NS}	-0.53**	-0.48**	-0.30^{NS}	0.39*	0.23^{NS}	-0.56**
	Dry	0.30 ^{NS}	0.07 ^{NS}	-0.14 ^{NS}	-0.24 ^{NS}	0.34 ^{NS}	0.02 ^{NS}	-0.04 ^{NS}	-0.38 ^{NS}	0.26 ^{NS}	0.26 ^{NS}
CH_4	Rainy	0.05^{NS}	-0.09 ^{NS}	0.44 ^{NS}	-0.27^{NS}	0.09^{NS}	-0.11 ^{NS}	-0.04 ^{NS}	-0.13 ^{NS}	-0.07 ^{NS}	0.04 ^{NS}
	Annual	0.04^{NS}	-0.10 ^{NS}	-0.01 ^{NS}	-0.18 ^{NS}	0.08^{NS}	-0.01 ^{NS}	-0.17 ^{NS}	-0.21 ^{NS}	-0.08 ^{NS}	0.02^{NS}

Total Carbon (T_C; g kg⁻¹); Total Nitrogen (T_N; g kg⁻¹); Microbial Carbon (Cmic, g kg⁻¹); Microbial Nitrogen (N_{mic}, g kg⁻¹); Carbon and Nitrogen ratio (C/N); Organic Matter (OM; g kg⁻¹); Salinity (Sal; ppt); Redox Potential (Eh; mV); Soil Moisture (Moisture, %).

NS= not significant; * significant effects at $p \le 0.05$; ** significant effects at $p \le 0.01$

4 Discussion

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4.1 Carbon dioxide and methane flux

343 It is important to consider that the year under study was rainier in the dry season (2017) 344 and less rainy in the wet season (2018) when the climatological average is concerned 345 (1981-2010) (Figure Fig. 3). Perhaps this variation is related to the La Niña effects 346 (extreme event), taking into account that and the intensification and higher frequency of 347 extreme events result from is considered as climate change (Barichivich et al., 2018). 348 Under these conditions, negative and positive fluxes of the two greenhouse gases were found (negative values represented gas consumption). The negative CO2 flux is 349 apparently a consequence of the increased CO2 solubility in tidal waters or of the 350 351 increased sulfate reduction, as described in the literature (Borges et al., 2018; 352 Chowdhury et al., 2018; Nóbrega et al., 2016). Fluctuations in redox potential altered 353 the availability of the terminal electron acceptor and donor, and the forces of recovery 354 of their concentrations in the soil, such that a disproportionate release of CO2 can result 355 from the alternative anaerobic degradation processes such as sulfate and iron reduction 356 (Chowdhury et al., 2018). The soil carbon flux in the mangrove area in the Amazon region was within the range of findings for other tropical mangrove areas (2.6 to 11.0 g 357 CO₂ m⁻² d⁻¹; Shiau and Chiu, 2020). However, the mean flux of 6.2 mmol CO₂ m⁻² h⁻¹ 358 359 recorded in this Amazonian mangrove was much higher than the mean efflux of 2.9 mmol CO₂ m⁻² h⁻¹ recorded in 75 mangroves during low tide periods (Alongi, 2009). 360 An emission of 0.01 Tg CH₄ y⁻¹, 0.6 g CH₄ m⁻² d⁻¹ (Rosentreter et al., 2018a), or 26.7 361 mg CH₄ m⁻² h⁻¹ has been reported for tropical latitudes (0 and 5°). In our study, the 362 monthly average of CH₄ flux was higher at the low $(7.3 \pm 8.0 \text{ mg CH}_4 \text{ m}^{-2} \text{ h}^{-1})$ than at 363 the high topography (0.9 \pm 0.6 mg C m⁻² h⁻¹), resulting in 0.1 g CH₄ m⁻² d⁻¹ or 0.5 Mg 364 CH₄ ha⁻¹ y⁻¹ (Figure Fig. 2). Therefore, the CH₄-C fluxes from the mangrove soil in the 365 366 Mojuim River estuary were much lower than expected. It is known that there is a 367 microbial functional module for CH₄ production and consumption (Xu et al., 2015) and 368 diffusibility of CH₄ (Sihi et al., 2018), and this module considers three key mechanisms: 369 aceticlastic methanogenesis (acetate production), hydrogenotrophic methanogenesis (H₂ and CO₂ production), and aerobic methanotrophy (CH₄ oxidation and O₂ reduction). 370 The average emission from the soil of 8.4 mmol CH₄ m⁻² d⁻¹ was well below the fluxes 371 recorded in the Bay of Bengal, with 18.4 mmol CH₄ m⁻² d⁻¹ (Biswas et al., 2007). In the 372 373 Amazonian mangrove studied the mean annual carbon equivalent efflux was 429.6 mg

CO_{2-eq} m⁻² h⁻¹. This value is—was insignificant—very low compared to the projected erosion losses of 103.5 Tg CO_{2-eq} ha⁻¹ y⁻¹ for the next century in tropical mangrove forests (Adame et al., 2021). These higher CO₂ flux concomitantly with lower CH₄ flux in this Amazonian estuary are probably a consequence of changes in the rainfall pattern already underway, where the dry season was wetter and the rainy season drier when compared to the climatological normal. The most recent estimate between latitude 0° to 23.5° S shows an emission of 2.3 g CO₂ m⁻² d⁻¹ (Rosentreter et al., 2018b). However, the efflux in the mangrove of the Mojuim River estuary was 6.7 g CO₂ m⁻² d⁻¹. For the same latitudinal range, Rosentreter et al. (2018c) estimated an emission of 0.6 g CH₄ m⁻² d⁻¹, and we found an efflux of 0.1 g CH₄ m⁻² d⁻¹.

4.2 Drivers of greenhouse gas fluxes

Mangrove areas are periodically flooded, with a larger flood volume during the syzygy tides, especially in the rainy season. The hydrological condition of the soil is determined by the microtopography and can regulate the respiration of microorganisms (aerobic or anaerobic), being a decisive factor in controlling the CO_2 efflux (Dai et al., 2012; Davidson et al., 2000; Ehrenfeld, 1995). No significant influence on CO_2 flux was observed due to the low variation in high tide level throughout the year (0.19 m) (Figure Fig. 2), although it was numerically higher at the high topography. However, tidal height and the rainy season resulted in a higher CO_2 flux (rate high/low =1.7) at the high topography (7.86 \pm 0.04 g CO_2 m⁻² d⁻¹) than at the low topography (4.73 \pm 0.34 g CO_2 m⁻² d⁻¹) (Figure Fig. 2; SI 1). This result may be due to the root systems of most flood-tolerant plants remaining active when flooded (Angelov et al., 1996). Still, the high topography has longer flood-free periods, which only happens when the tides are syzygy or when the rains are torrential.

CO₂ efflux was higher in the high topography than in the low topography in the rainy season (when soils are more subject to inundation), i.e., 39.8% lower in the forest soil exposed to the atmosphere for less time. Measurements performed on 62-mangrove forest soils showed an average flux of 2.87 mmol CO₂ m⁻² h⁻¹ when the soil was exposed to the atmosphere (dry soil), while 75-results on flooded mangrove forest soils showed an average emission of 2.06 mmol CO₂ m⁻² h⁻¹ (Alongi, 2007, 2009), i.e., 28.2% less than for the dry soil. This reflects the increased facility gases have for molecular diffusion than fluids, and the increased surface area available for aerobic respiration and chemical oxidation during air exposure (Chen et al., 2010). Some studies

- 407 attribute this variation to the temperature of the soil when it is exposed to tropical air
- 408 (Alongi, 2009), which increases the export of dissolved inorganic carbon (Maher et al.,
- 409 2018). However, although despite the lack of significant variation in soil temperature
- 410 between topographies at each time of year (Figure Fig. 4b), there was a positive
- 411 correlation (Pearson = 0.15, p = 0.05) between CO_2 efflux and soil temperature at the
- 412 low topography.
- 413 Some studies show that CH₄ efflux is a consequence of the seasonal temperature
- variation in mangrove forest under temperate/monsoon climates (Chauhan et al., 2015;
- Purvaja and Ramesh, 2001; Whalen, 2005). However, in your study CH₄ efflux was
- 416 correlated with Ta (Pearson = -0.33, p < 0.00) and RH (Pearson = 0.28, p = 0.01) only
- 417 in the dry season and at the low topography. The results show that the physical
- 418 parameters do not affect the fluxes in a standardized way, and their greater or lesser
- influence depends on the topography and seasonality.
- 420 A compilation of several studies showed that the total CH₄ emissions from the soil in a
- 421 mangrove ecosystem range from 0 to 23.68 mg C m⁻² h⁻¹ (Shiau and Chiu, 2020), and
- 422 our study showed a range of -0.01 to 31.88 mg C m⁻² h⁻¹ (mean of 4.70 ± 5.00 mg C m⁻²
- 423 h^{-1}). The monthly CH₄ fluxes were generally higher at the low $(0.232 \pm 0.256 \text{ g CH}_4 \text{ m}^{-2})$
- 424 d^{-1}) than at the high $(0.026 \pm 0.018 \text{ g CH}_4 \text{ m}^{-2} \text{ d}^{-1})$ topography, especially during the
- rainy season when the tides were higher (Figure Fig. 2). Only in the dry season was
- 426 there a significantly higher production at the low than at the high topography (Figure
- 427 Fig. 2; SI 1). The low topography produced 0.0249 g C m⁻² h⁻¹ more to the atmosphere
- 428 in the rainy season than in the dry season (Figure Fig. 2), and a similar seasonal pattern
- was recorded in other studies (Cameron et al., 2021).
- The mangrove soil in the Mojuim River estuary is rich in silt and clay (Table 1), which
- 431 reduces sediment porosity and fosters the formation and maintenance of anoxic
- conditions (Dutta et al., 2013). In addition, the lack of oxygen in the flooded mangrove
- 433 soil favors microbial processes such as denitrification, sulfate reduction,
- 434 methanogenesis, and redox reactions (Alongi and Christoffersen, 1992). A significant
- amount of CH₄ produced in wetlands is dissolved in the pore water due to high pressure,
- 436 causing supersaturation, which allows CH₄ to be released by diffusion from the
- sediment to the atmosphere and by boiling through the formation of bubbles.
- Studies show that the CO₂ flux tends to be lower with high soil saturation (Chanda et
- al., 2014; Kristensen et al., 2008). A total of 395 Mg C ha⁻¹ was found at the soil surface

- 440 (0.15 m) in the mangrove of the Mojuim River estuary, which was slightly higher than
- 441 the 340 Mg C ha⁻¹ found in other mangroves in the Amazon (Kauffman et al., 2018),
- 442 however being significantly 1.8 times greater at the low topography (Table 2). The finer
- 443 soil texture at the low topography (Table 1) reduces groundwater drainage which
- facilitates the accumulation of C in the soil (Schmidt et al., 2011).

445 **4.3 Mangrove biomass**

- Only the species R. mangle and A. germinans were found in the floristic survey carried
- 447 out, which is aligned with the results of other studies in the same region (Menezes et al.,
- 448 2008). Thus, the variations found in the flux between the topographies in the Mojuim
- 449 River estuary are not related to the mangrove forest structure, because there was no
- 450 difference in the aboveground biomass. Since there was no difference in the species
- composition, the belowground biomass is not expected to differ either (Table 3).
- 452 Assuming that the amount of carbon stored is 42.0% of the total biomass (Sahu and
- 453 Kathiresan, 2019), the mangrove forest biomass of the Mojuim River estuary stores
- 454 127.9 and 138.9 Mg C ha⁻¹ at the high and low topographies, respectively. This result is
- lower than the 507.8 Mg C ha⁻¹ estimated for Brazilian mangroves (Hamilton and
- 456 Friess, 2018), but are near the 103.7 Mg C ha⁻¹ estimated for a mangrove at Guará's
- 457 island (Salum et al., 2020), 108.4 Mg C ha⁻¹ for the Bragantina region (Gardunho,
- 458 2017), and 132.3 Mg C ha⁻¹ in French Guiana (Fromard et al., 1998). Thus, the biomass
- 459 found in the Mojuim estuary does not differ from the biomass found in other
- 460 Amazonian mangroves. The estimated primary production for tropical mangrove forests
- 461 is 218 ± 72 Tg C y^{-1} (Bouillon et al., 2008).

4.4 Biogeochemical parameters

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- 463 During the seasonal and annual periods, CH₄ efflux was not significantly correlated
- with chemical parameters (Table 5), similar as observed in another study (Chen et al.,
- 465 2010). Flooded soils present reduced gas diffusion rates, which directly affects the
- 466 physiological state and activity of microbes, by limiting the supply of the dominant
- 467 electron acceptors (e.g., oxygen), and gases (e.g., CH₄) (Blagodatsky and Smith, 2012).
- 468 The importance of soil can be reflected in bacterial richness and diversity compared to
- 469 pore spaces filled with water (Banerjee et al., 2016). On the other hand, increasing soil
- 470 moisture provides the microorganisms with essential substrates such as ammonium,
- 471 nitrate, and soluble organic carbon, and increases gas diffusion rates in the water

472 (Blagodatsky and Smith, 2012). Biologically available nitrogen often limit marine 473 productivity (Bertics et al., 2010), and thus can affect CO₂ fluxes to the atmosphere. 474 However, a mangrove fertilization experiment showed that CH₄ emission rates were not 475 affected by N addition (Kreuzwieser et al., 2003). A higher concentration of C_{mic} and 476 N_{mic} in the dry period (Table 2), both in the high and low topographies, indicated that 477 microorganisms are more active when the soil spends more time aerated in the dry 478 period (Table 2), time when only the high tides produce anoxia in the mangrove soil 479 mainly in the low topography. Under reduced oxygen conditions, in a laboratory 480 incubated mangrove soil, the addition of nitrogen resulted in a significant increase in the 481 microbial metabolic quotient, showing no concomitant change in microbial respiration, 482 which was explained by a decrease in microbial biomass (Craig et al., 2021). 483 The high OM concentration at the two topographic locations (Table 2), at the two 484 seasons studied, and the respective negative correlation with CO₂ flux (Table 5) confirm 485 the importance of microbial activity in mangrove soils (Gao et al., 2020). Also, CH₄ 486 produced in flooded soils can be converted mainly to CO₂ by the anaerobic oxidation of 487 CH₄ (Boetius et al., 2000; Milucka et al., 2015; Xu et al., 2015) which may contribute to 488 the higher CO₂ efflux in the Mojuim River estuary compared to other tropical 489 mangroves (Rosentreter et al., 2018b). The belowground C stock is considered the 490 largest C reservoir in a mangrove ecosystem, and it results from the low OM 491 decomposition rate due to flooding (Marchand, 2017). 492 The higher water salinity influenced by the tidal movement in the dry season (Table 1) 493 seems to result in a lower CH₄ flux at the low topography (Dutta et al., 2013; Lekphet et al., 2005; Shiau and Chiu, 2020). High SO_4^{2-} concentration in the marine sediments 494 inhibits methane formation due to competition between SO_4^{2-} reduction and 495 496 methanogenic fermentation, as sulfate-reducing bacteria are more efficient at using hydrogen than methanotrophic bacteria (Abram and Nedwell, 1978; Kristjansson et al., 497 1982), a key factor fostering reduced CH₄ emissions. At high SO₄²- concentrations 498 methanotrophic bacteria use CH₄ as an energy source and oxidize it to CO₂ (Coyne, 499 500 1999; Segarra et al., 2015), increasing the efflux of CO₂ and reduced CH₄ (Megonigal 501 and Schlesinger, 2002; Roslev and King, 1996). This may explain the high CO₂ and low 502 CH₄ efflux found throughout the year at the high and, especially, at the low

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topographies (Figure Fig. 3).

504 Studies in coastal ecosystems in Taiwan have reported that methanotrophic bacteria can 505 be sensitive to soil pH, and reported an optimal growth at pH ranging from 6.5 to 7.5 506 (Shiau et al., 2018). The higher soil acidity in the Mojuim River wetland (Table 1) may 507 be inhibiting the activity of methanogenic bacteria by increasing the population of 508 methanotrophic bacteria, which are efficient in CH₄ consumption (Chen et al., 2010; 509 Hegde et al., 2003; Shiau and Chiu, 2020). In addition, the pneumatophores present in 510 R. mangle increase soil aeration and reduce CH₄ emissions (Allen et al., 2011; He et al., 511 2019). Spatial differences (topography) in CH₄ emissions in the soil can be attributed to 512 substrate heterogeneity, salinity, and the abundance of methanogenic and 513 methanotrophic bacteria (Gao et al., 2020). Increases in CH₄ efflux with reduced 514 salinity were found as a consequence of intense oxidation or reduced competition from the more energetically efficient SO₄²⁻ and NO³⁻ reducing bacteria when compared to the 515 methanogenic bacteria (Biswas et al., 2007). This fact can be observed in the CH₄ efflux 516 517 in the mangrove of the Mojuim River, because there was an increased CH₄ production 518 especially in the low topography in the rainy season (Figure Fig. 3), when water salinity 519 is reduced (Table 1) due to the increased precipitation. However, we did not find a 520 correlation between CH₄ efflux and salinity, as previously reported (Purvaja and 521 Ramesh, 2001).

522 **5 Conclusions**

- 523 Seasonality was important for CH₄ efflux but did not influence CO₂ efflux. The
- 524 differences in fluxes may be an effect of global climate changes on the terrestrial
- 525 biogeochemistry at the plant-soil-atmosphere interface, as indicated by the deviation in
- 526 precipitation values from the climatology normal, making it necessary to extend this
- study for more years. Using the factor of 23 to convert the global warming potential of
- 528 CH₄ to CO₂ (IPCC, 2001), the CO₂ equivalent emission was 35.4 Mg CO_{2-eq} ha⁻¹ yr⁻¹.
- Over a 100-year time period, a radiative forcing due to the continuous emission of 0.05
- kg CH₄ m⁻² y⁻¹ found in this study, would be offset if CO₂ sequestration rates were 2.16
- kg CO_2 m⁻² y⁻¹ (Neubauer and Megonigal, 2015).
- 532 Microtopography should be considered when determining the efflux of CO₂ and CH₄ in
- mangrove forests in an Amazon estuary. The low topography in the mangrove forest of
- 534 Mojuim River had a higher concentration of organic carbon in the soil. However, it did
- 535 not produce a higher CO₂ efflux because it was negatively influenced by soil moisture,
- 536 which was indifferent to CH₄ efflux. MOOM, C/N ratio, and Eh were critical in soil

- 537 microbial activity, which resulted in a variation in CO2 flux during the year and
- seasonal periods. Thus, the physicochemical properties of the soil are important for CO₂
- flux, especially in the rainy season. Still, they did not influence CH₄ fluxes.
- 540 Data availability: The data used in this article belong to the doctoral thesis of Saul
- Castellón, within the Postgraduate Program in Environmental Sciences, at the Federal
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- (saulmarz22@gmail.com), which holds the set of all data used in this paper.
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