Greenhouse gas fluxes in mangrove forest soil in the a Amazon estuary

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Abstract: Tropical mangrove forests are important carbon sinks, the soil being the main reservoir of this chemical element. Understanding the variability and the key factors that control fluxes is critical to account for greenhouse gas (GHG) emissions, particularly in the current scenario of global climate change especially in a scenario of global climate change. The current study is the first to quantify methane (CH₄) and carbon dioxide (CO₂) emissions using a dynamic chamber in Amazon natural mangrove soils. The plots for the trace gases study were allocated at contrasting topographic heights. Sampling points were selected in a contrasting topographic gradient, the highest point being where flooding occurs only at high tides during the solstice and on the high tides of the rainy season of the new and full moons. The results showed that the mangrove soil of the Amazon estuary is a source of CO2 (6.66 g CO2 m-2 d-1) and CH4 (0.13 g CH4 m-2 d-1) to the atmosphere. In the spatial variation, the CO₂ flux was higher in the high topography (7.858 g CO₂ m⁻² d⁻¹) in comparison to the low topography (4.734 g CO₂ m⁻¹ ² d⁻¹) in the rainy season, and the CH₄ was higher in the low topography (0.128 g CH₄ m⁻² d⁻¹) than in the high topography (0.014 g CH₄ m⁻² d⁻¹) in the dry season. The results showed that mangrove soils are sources of greenhouse gases, and CO2 fluxes were not different between seasons, and only in the dry period were they greater in the high topography. Only in the low topography, the CH₄ fluxes were higher in the rainy season. However, in the dry period, the low topography soil produced more CH₄. Soil organic matter, carbon and nitrogen ratio (C/N), and redox potential influenced the annual and seasonal variation of CO2 emissions; however, they did not influence CH4 flux. The mangrove soil of the Amazon estuary produced 35.4 Mg CO2-eq ha-1 year-1, and to counterbalance CH4 emissions is necessary to sequester 2.16 kg CO2 m-2 y-1 by the mangrove ecosystem. To account for global GHG emissions, in the Amazonian estuary mangrove soil produced 35.4 Mg CO_{2-eq} ha⁻¹ yr⁻¹.

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1 Introduction

The Amazon coastal areas in the State of Pará (Brazil) cover 2,176.8 km² where mangroves develop under the macro tide regime in the (Souza Filho, 2005), representing approximately 85% of the entire area of Brazilian mangroves (Herz, 1991). These_mangrove areas are estimated to be the main contributors to greenhouse gas emissions in marine ecosystems (Allen et al., 2011; Chen et al., 2012). However, mangrove forests are highly productive due to a high nutrient turnover rate (Robertson et al., 1992) and have mechanisms that maximize carbon gain and minimize water loss through plant transpiration (Alongi and Mukhopadhyay, 2015). A study conducted in 25 mangrove forests (between 30° latitude and 73° longitude) revealed that these forests are the richest in carbon 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). In addition, phenolic compounds inhibit microbial activity and help keep organic earbon intact, thus accumulating organic matter in mangrove forest soils (Friesen et al., 2018).

The production of greenhouse gases from soils is mainly attributable to biogeochemical processes. Microbial activities and gas production are related to soil properties, including total carbon and total nitrogen concentrations, moisture, porosity, salinity, and redox potential (Bouillon et al., 2008; Chen et al., 2012). Due to the dynamics of tidal movements, mangrove soils may become saturated and present a reduced oxygen availability or total acration caused by the ebb tide. Studies attribute soil carbon flux responses to moisture perturbations because of seasonality and flooding events (Banerjee et al., 2016), with fluxes being dependent on tidal extremes (high tide and low tide), and flood duration (Chowdhury et al., 2018).

The estimated <u>soil CO₂ outgassingCO₂ production to the atmosphere</u>, in tropical estuarine areas, is 16.2 Tg_C_y⁻¹ (Alongi, 2009). <u>However, soil efflux measurements from tropical mangroves revealed emissions ranging from 2.9 to 11.0 g CO2 m-2 d-1 However, the most recent estimate between latitude 0° to 23.5° S reveals an emission of 2.3 g CO₂ m⁻² d⁻¹ (Castillo et al., 2017; Chen et al., 2014; Shiau and Chiu, 2020). In situ CO₂ production is related to the water input of terrestrial, riparian, and groundwater brought by rainfall (Rosentreter et al., 2018b).</u>

Due to thise periodic tidal influencemovement, the mangrove ecosystem is regularly daily flooded, leaving the soil anoxic and consequently reduced, and favoring methanogenesis (Dutta et al., 2013). Thus, estuaries are considered hot spots for CH₄

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production and emission (Bastviken et al., 2011; Borges et al., 2015). The organic material decomposition by methanogenic bacteria in anoxic environments, such as sediments, inner suspended particles, zooplankton gut (Reeburgh, 2007; Valentine, 2011), and the impact of freshwater should change the electron flow from sulfatereducing bacteria to methanogenesis reduction of sulfate in anoxic marine sediments (Purvaja et al., 2004), also results in CH₄ formation. On the other hand, an ecosystem with salinity levels greater than 18 ppt may show an absence of CH₄ emissions (Poffenbarger et al., 2011), since methane dissolved in pores is typically oxidized anaerobically by sulfate (Chuang et al., 2016). Currently the uncertainties in emitted CH₄ values in vegetated coastal wetlands are approximately 30% (EPA, 2017). Soil flux measurements from tropical mangroves revealed emissions ranging from 0.3 to 4.4 mg CH4 m-2 d-1. The total emission of 0.010 Tg CH₄ y⁴ or 0.64 g CH4 m² d⁴ was estimated between 0 and 5° latitude (Castillo et al., 2017; Chen et al., 2014; Kreuzwieser et al., 2003). The production of greenhouse gases from soils is mainly driven by attributable to biogeochemical processes. Microbial activities and gas production are related to soil properties, including total carbon and total nitrogen concentrations, moisture, porosity, salinity, and redox potential (Bouillon et al., 2008; Chen et al., 2012). Due to the dynamics of tidal movements, mangrove soils may become saturated and present a reduced oxygen availability or total aeration caused by the ebb tide. Studies attribute soil carbon flux responses to moisture perturbations because of seasonality and flooding events (Banerjee et al., 2016), with fluxes being dependent on tidal extremes (high tide and low tide), and flood duration (Chowdhury et al., 2018). In addition, phenolic compounds inhibit microbial activity and help keep organic carbon intact, thus accumulating organic matter in mangrove forest soils (Friesen et al., 2018). The Amazon coastal areas in the State of Pará (Brazil) cover 2,176.8 km² where mangroves develop under the macro-tide regime in the (Souza Filho, 2005), representing 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 gases fluxes were studied together with the analysis of the vegetation structure and soil physical-chemical parameters. The

objective of this study is to investigate the spatial and seasonal variation in the monthly

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fluxes of CO₂-and CH₄ from the soil in a non-anthropized mangrove area in the Mojuim
River Estuary, belonging to the Amazon biome. The environmental factors and
physicochemical analysis of the soil were investigated from 2017 to 2018 to understand
the gas fluxes.

2 Material and Methods

2.1 Study site

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This study was conducted in the Amazonian coastal zone, Macaca Island (-0.746491 latitude, -47.997219 longitude), located in the Mojuim River estuary, at the Mocapajuba Marine Extractive Reserve, municipality of São Caetano de Odivelas (Figure 1), state of Pará (Brazil). Macaca island has an area of 1,322 ha with pristine mangroves exclusively untouched mangrove forests, which belongs to a coastal stripmangrove area of 2177 km² in the state of Pará (Souza Filho, 2005). The climate is Am type (tropical monsoon) according to the Köppen classification (Peel et al., 2007). The climatological data were obtained from the Meteorological Database for Teaching and Research of the National Institute of Meteorology (INMET). The area has a rainy season from January to June (2,296 mm of precipitation) and a dry season from July to December (687 mm). March and April are the rainiest months with 505 and 453 mm of precipitation, while October and November are the driest (53 and 61 mm, respectively). The minimum temperatures occur in the rainy period (26 °C) and the maximum in the dry period (29 °C). The Mojuim estuary has a macrotidal regime, with an average amplitude height of 4.9 m during spring tide and 3.2 m during low tide (Rollnic et al., 2018). During the wet season the Mojuim River has a flow velocity of 1.8 m s⁻¹ at the ebb tide and 1.3 m s⁻¹ at the flood tide. During the dry season, the maximum currents are 1.9 m s⁻¹ at the flood and 1.67 m s⁻¹ at the ebb tide (Rocha, 2015). The annual mean salinity is 26.95 ± 0.98 PSU (Valentim et al., 2018).

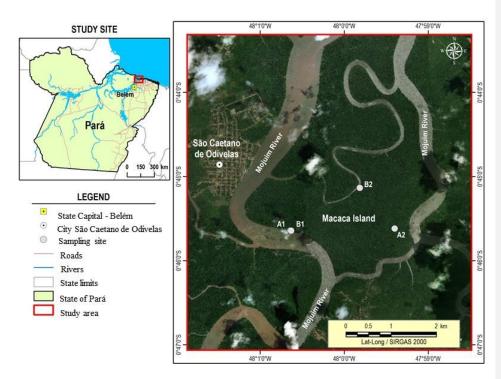


Figure 1. Macaca Island located in the mangrove coast of Northern Brazil, Municipality of São Caetano de Odivelas (state of Pará), with the sampling points at low (B1 and B2) and high topographies (A1 and A2). Image Source: © Google Earth

The Mojuim River region is geomorphologically formed by partially submerged river basins consequent of an 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). This river forms the entire watershed of the municipality of São Caetano de Odivelas and borders the municipality of São João da Ponta (Figure 1). 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) with various ages (Fernandes and Pimentel, 2019). The population economically exploited the estuary, primarily by artisanal fishing, crab (*Ucides cordatus* L.) extraction, and oyster farms.

Four sampling sites were selected in the Macaca Island: two where flooding occurs every day (B1 and B2; Figure 1), called low topography, and two where flooding occurs only at high tides during the solstice and on the high tides of the rainy season of the new and full moons (A1 and A2; Figure 1), called high topography. Once a month, the gas

flux for each chamber was measured during periods of waning or crescent moon, these are the times when the soil in the low topography is more exposed. The flora of the mangrove area on the Macaca Island is little anthropized and comprises the genera Rhizophora, Avicenia, Laguncularia, and Acrostichum (Ferreira, 2017; França et al., 2016). The estuarine plains are influenced by a-macro-tide dynamics and can be physiographically divided into four sectors (França et al., 2016). The Macaca Island is classified as being from the fourth sector, which consists of woods of adult trees of the genus Ryzophora with an average height of 10 to 25 m, located at an elevation of 0 to 5 m, with silt-clay soil (França et al., 2016).

Four sampling sitesplots were selected in the Macaca Island (Figure 1) on 19/05/2017, when the moon was in the waning quarter phase: two plots where flooding occurs every day (plots B1 and B2; Figure 1), called low topography, and two plots where flooding occurs only at high tides during the solstice and on the high tides of the rainy season of the new and full moons (plots A1 and A2; Figure 1), called high topography. Once a month, the gas flux for each chamber was measured during periods of waning or erescent moon, as these are the times when the soil in the low topography is more exposed.

2.2 Greenhouse gas flux measurements

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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 an area of 0.028 m⁻² (volume of 3.47 L) and were fixed 0.05 m into the ground, and remained in place until the study was completed. Once a month, the Greenhouse gas flux was measured during periods of waning or crescent moon, as these are the times when the soil in the low topography is more exposed. To avoid the influence of mangrove roots on the gas fluxes, the rings were placed in locations without any seedlings or aboveground mangrove roots. CO₂ and CH₄ concentration (ppm) were measured using the dynamic chamber methodology (Norman et al., 1997; Verchot et al., 2000), sequentially connected to a Los Gatos Research portable gas analyzer (Mahesh et al., 2015). The device was calibrated monthly with high quality standard gas (500 ppm CO₂; 5 pmm CH₄). The rings were sequentially closed for three minutes with a PVC cap, which enabled the connection to the analyzer via two 12.0 m polyethylene hoses. The gas concentration was measured every two seconds and automatically stored by the analyzer. CO2 and CH4 fluxes were calculated from the linear regression of Formatado: Título 2

increasing/decreasing CO_2 and CH_4 concentrations within the chamber, usually between one and three minutes after the ring cover was placed (Frankignoulle, 1988; McEwing et al., 2015). Analyzing the literature, we found that the flux is considered zero when the linear regression reaches an $R^2 < 0.30$ (Sundqvist et al., 2014). However, in our analyses, the vast majority of regressions reached an $R^2 > 0.70$, and the regressions were weak in only 6% of the data, which were considered to be zero. At the end of each flux measurement, the height of the ring above ground was measured at four equidistant points with a ruler at each gas flux measurement. For the seasonal data is it average monthly fluxes in the wet compared with average monthly fluxes in the dry season.

2.22.3 Vegetation structure and biomass

The floristic survey was conducted in October 2017, at the same topographies as the gas flux analysis at the same sites as the gas flow study, using circular plots of 1,256.6 m² (Kauffman et al., 2013), divided into four subplots of 314.15 m², which is the equivalent to 0.38 ha (Figure 1). All trees with DBH (diameter at breast height) greater than 0.05m had their diameter above the aerial roots, the diameter of the stem, and total height recorded. The allometric equations (Howard et al., 2014) to calculate tree biomass (aboveground biomass; AGB) were: *R. mangle* AGB = 0.1282 * DBH^{2.6} (R² = 0.92); *A. germinans* AGB = 0.140 * DBH^{2.4} (R² = 0.97); Total AGB = 0.168 * ρ * DBH^{2.47} (R² = 0.99), where $\rho_{R. mangle} = 0.87$; $\rho_{A. germinans} = 0.72$ (ρ = wood density) The allometric equation to calculate tree biomass (AGB) was: *AGB* = 0.168 × ρ × 2.471, where ρ represents wood density, using 0.87 g cm³ for *R. mangle* and 0.72 g cm³ for *A. germinans* (Howard et al., 2014).

2.32.4 Soil sampling and environmental characterization

In July 2017 and January 2018, four soil samples were collected with an auger at a depth of 0.10 m in all the studied sites plots for gas flux measurements (Figure 1). Before the soil samples were 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. In the laboratory, the soil analyzes began on the following day of collection, and the samples were kept in a freezer. The soil samples were properly stored and taken to the Chemical Analysis

210	Laboratory of the <i>Museu Paraense Emílio Goeldi</i> . Salinity (Sal; ppt) was measured with			
211	PCE-0100, and soil moisture (Sm; %) by the residual gravimetric method (EMBRAPA,			
212	1997).			
213	Organic Matter (OM; g kg ⁻¹), Total Carbon (T _C ; g kg ⁻¹) and Total Nitrogen (T _N ; g kg ⁻¹)	F	ormatado: Subscrito	
214	were calculated by volumetry (oxidoreduction) using the Walkley-Black method	F	ormatado: Subscrito	
215	(Kalembasa and Jenkinson, 1973). Microbial carbon (C _{mic} ; mg kg ⁻¹) and microbial	F	ormatado: Subscrito	
216	nitrogen (N _{mic} ; mg kg ⁻¹) were determined through the 2,0 min of Irradiation-extraction	F	ormatado: Subscrito	
217	method of soil by microwave technique (Islam and Weil, 1998). Microwave heated soil			
218	extraction proved to be a simple, fast, accurate, reliable and safe method to measure soil			
219	microbial biomass (Araujo, 2010; Ferreira et al., 1999; Monz et al., 1991). The C _{mic} was	F	ormatado: Subscrito	
220	determined by dichromate oxidation (Kalembasa and Jenkinson, 1973; Vance et al.,			
221	1987). The N _{mic} was analyzed following the method described by Brookes et al. (1985),	F	ormatado: Subscrito	
222	changing fumigation to irradiation, which uses the difference between the amount of TN			
223	in irradiated and non-irradiated soil. We used the flux conversion factor of 0.33			
224	(Sparling and West, 1988) and 0.54 (Almeida et al., 2019; Brookes et al., 1985), for			
225	carbon and nitrogen, respectively. Particle size analysis was performed separately on			
226	four soil samples collected at each flux siteplot, in the two seasons (October 2017 and			
227	March 2018), according to EMBRAPA (1997).			
228	At each gas flux flow-measurement, environmental variables such as air temperature			
229	(Tair, °C), relative humidity (RH, %), wind speed (Ws, m s-1) were quantified with a	F	ormatado: Subscrito	
230	portable thermo-hygrometer (model AK821) at the height of 2.0 m above the soil	F	ormatado: Subscrito)
231	surface. Soil temperature (Ts, °C) was measured with a portable digital thermometer	F	ormatado: Subscrito	
232	(model TP101) sequentially after each gas fluxflow measurement. Daily precipitation			
233	was obtained from an automatic precipitation station installed at a pier on the banks of			
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234	the Mojuim River in São Caetano das Odivelas (coordinates: 0°44'18.48 "S;			
234	the Mojuim River in São Caetano das Odivelas (coordinates: 0°44'18.48 "S; 48°00'47.94 "W).			
	·	F	ormatado: Título 3	
235	48°00'47.94 "W).	F	ormatado: Título 3	
235236	48°00'47.94 "W). 2.4 Fluxes Measurements	F	ormatado: Título 3	
235236237	48°00'47.94 "W). 2.4 Fluxes Measurements In each plot, eight Polyvinyl Chloride rings with 0.20 m diameter and 0.12 m height	F	ormatado: Título 3	
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235236237238239	48°00'47.94 "W). 2.4 Fluxes Measurements 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 an area of 0.028 m ⁻² (volume of 3.47 L) and were fixed 0.05 m into the ground. The	Fi	ormatado: Título 3	
235 236 237 238 239 240	48°00'47.94 "W). 2.4 Fluxes Measurements 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 an area of 0.028 m ⁻² (volume of 3.47 L) and were fixed 0.05 m into the ground. The height of the ring above ground was measured at four equidistant points with a ruler at	Fi	ormatado: Título 3	

CO₂ and CH₄ fluxes (g CO₂ or CH₄ m⁻² d⁻¹) were measured using the dynamic chamber methodology (Norman et al., 1997; Verchot et al., 2000), sequentially connected to a Los Gatos Research portable gas analyzer (Mahesh et al., 2015). The device was calibrated monthly with high quality standard gas. The rings were sequentially closed for three minutes with a PVC cap, which enabled the connection to the analyzer via two 12.0 m polyethylene hoses. The gas concentration was measured (ppm) every two seconds and automatically stored by the analyzer. CO₂ and CH₄ fluxes were calculated from the linear regression of increasing/decreasing CO₂ and CH₄ concentrations within the chamber, usually between one and three minutes after the ring cover was placed (Frankignoulle, 1988; McEwing et al., 2015). Analyzing the literature, we found that the flux is considered zero when the linear regression reaches an R² < 0.30 (Sundqvist et al., 2014). However, in our analyses, the vast majority of regressions reached an R² > 0.70, and the regressions were weak in only 6% of the data.

2.5 Statistical analyses

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On Macaca Island, two treatments were allocated (low and high topography), with two plots in either. In each plot, eight chambers were randomly distributed, which were considered sample repetitions. The normality of the data of FCH₄ and FCO₂ and soil physicochemical parameters was determined by the Shapiro-Wilks method. The student's t test was used to test the differences (p < 0.05) in the emissions between the different sites and seasonal periods. An ANOVA and Tukey's test (p < 0.05) were used when the distributions were normal. The soil CO2 and CH4 flux showed a non-normal distribution, and for this reason, we used the non-parametric ANOVA (Kruskal-Wallis, p < 0.05) for testing the differences between the two treatments at months and season. For non-parametric data the Kruskal-Wallis test was used (p < 0.05). The physicochemical parameters showed a normal distribution, and therefore parametric ANOVA was used to test statistical differences (p < 0.05) between two treatments at months and season. Pearson correlation coefficients were calculated to determine the relationships between soil properties and gas fluxes in the months (dry and wet season) in which soil chemical properties analyzes were performed at the same time as gas fluxes were measuredPearson correlation coefficients were calculated to determine the relationships between soil properties and gas fluxes. Statistical analyses were performed with and free statistical software Infostat 2015®.

3 Results

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276 **3.1 Precipitation**

277 3.23.1 Carbon Dioxide dioxide and methane effluxes

278 The CO₂ and CH₄ fluxes in the mangrove soil were not normally distributed, so the 279 statistical analysis was performed using a non-parametric method. CO2 fluxes only 280 differed significantly among between topographies in January (H = 3.915; p = 0.048), 281 July (H = 9.091; p = 0.003), and November (H = 11.294; p < 0.000) (Figure 2; 282 Supplementary Information Table 1), with generally higher fluxes at the high 283 topography than at the low topography. CH4 fluxes were statistically different between 284 topographies only in November (H = 9.276; p = 0.002) and December (H = 4.945; p = 285 0.005), with higher fluxes at the low topography (Table 1) At the high topography, CO2 286 fluxes were significantly higher in July compared to August and December, March, 287 October, and May, not differing from the other months of the year (H = 24.510; p = 288 0.011). In the same way, aAt the low topography, CO₂ fluxes were statistically (H = 19.912; p = 0.046) higher in September and February compared to January and 289 November, not differing from the other months. We found a mean monthly flux of 290 $327.9 \pm 78.0 \text{ mg CO}_2 \text{ m}^{-2} \text{ h}^{-1}$ (mean \pm standard error) and $217.2 \pm 51.0 \text{ mg CO}_2 \text{ m}^{-2} \text{ h}^{-1}$, 291 292 at the high and low topography, respectively.

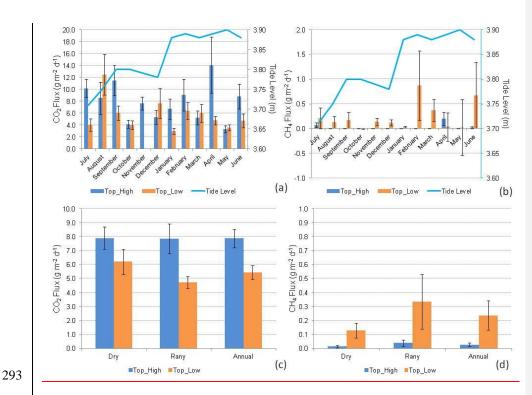


Figure 2. CO_2 (a) and CH_4 (b) fluxes (g CO_2 or CH_4 m⁻² d⁻¹) monthly (July 2018 to June 2019) (n = 16). Seasonal (Dry and Rany), and annual fluxes of CO_2 (c) and CH_4 (d), at high (Top_High) and low (Top_Low) topographies (n = 96), in mangrove forest soil compared to tide level (Tide Level). The bars represent the standard error of the mean.

Table 1. Monthly and seasonal (dry and rainy seasons) fluxes of CO_2 and CH_4 (g CO_2 or CH_4 -m⁻²-d⁻¹) at the high and low topographies. Numbers represent the mean (standard error). Lower case letters compare topographies in the same month. Upper case letters compare stations at each topography. Different boldface letters have statistically significant variation (Kruskal Wallis, p < 0.05).

	CO ₂ flux (g m⁻²-d⁻¹)	CH ₄ flux (g m ⁻² -d ⁻¹)			
	High topography	Low topography	High topography	Low topography		
July/2017	10.166(1.555) ^a	4.036(1.027) ^b	0.0724(0.0518) ^a	0.2129(0.2087) ^a		
August	8.513(2.672) ^a	12.462(3.400) ^a	0.0033(0.0016) ^a	0.1270(0.1185) ^a		
September	11.506(2.515) ^a	6.020(1.207) ^a	0.0014(0.0008) ^a	0.1738(0.1608) ^a		
October	4.147(0.653) ^a	3.993(0.731) ^a	0.0000(0.0000) ^a	-0.0004(0.0056) ^a		

November	7.648(1.064) ^a	$\frac{0.007(0.002)^{b}}{}$	-0.0004(0.0001) ^b	0.1395(0.0708) ^a
December	5.302(1.176) ^a	7.622(2.505) ^a	$0.0009(0.0009)^{b}$	0.1210(0.0575) ^a
Dry period	7.902(0.803) ^{aA}	6.202(0.895) ^{bA}	$0.0141(0.010)^{bB}$	0.1280(0.053)* Formatado: Fonte: Não Negrito
January/2018	6.697(1.717) ^a	$2.995(0.493)^{b}$	0.0007(0.0004) ^a	0.0294(0.0183) ^a
February	9.053(2.650) ^a	6.384(1.428) ^a	0.0049(0.0022) ^a	0.8743(0.7024) *
March	5.225(1.135) ^a	5.970(1.534) ^a	0.0077(0.0056) ^a	0.3736(0.2197) *
April	14.077(4.695) ^a	4.785(0.711) ^a	0.1968(0.1304) ^a	0.0372(0.2841)*
May	3.299(0.587) ^a	3.565(0.472) ^a	0.0014(0.0019) ^a	0.0218(0.5648) ^a
June	8.796(2.053) ^a	4.704(1.183) ^a	0.0226(0.0191) ^a	0.6739(0.6665) *
Rainy period	7.858(1.058) ^{aA}	4.734(0.440) ^{aA}	0.0390(0.023) ^{aA}	$0.3350(0.194)^{aA}$
At the high top	ography, CO ₂ flux	es were significant	tly higher in July co	ompared to

August and December, March, October, and May, not differing from the other months of the year (H = 24.510; p = 0.011). CH₄ fluxes at the high topography were significantly (H = 40.073; p < 0.001) higher in April and July compared to the other months studied, and in November there was consumption of CH₄ from the atmosphere (Table 1). At the low topography, CO₂ fluxes were statistically (H = 19.912; p = 0.046) higher in September and February compared to January and November, not differing from the other months. CH4 fluxes at the low topography did not show a significant variation between months (H = 10.114; p = 0.407). 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 2; Supplementary Information). 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, and in November there was consumption of CH₄ from the atmosphere (Figure 2; Supplementary Information). In the same way, CH₄ fluxes at the low topography did not show a significant variation between months (H = 10.114; p = 0.407). Greenhouse gas fluxes (Figure 2) were only significativey different between topographies in the dry season (Figure 3) where CO_2 fluxes were higher (H = 7.378; p = 0.006) at the high topography and CH_4 fluxes were higher (H = 8.229; p < 0.001) at the low topography Although seasonal CO2 fluxes were higher at the high topography than

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topographies in the dry season (Figure 3) where CO_2 fluxes were higher (H = 7.378; p = 0.006) at the high topography and CH_4 fluxes were higher (H = 8.229; p < 0.001) at the low topography Although seasonal CO_2 fluxes were higher at the high topography than at the low topography (Table 1), they were only statistically different in the dry season (H = 7.378; p = 0.006). In contrast, seasonal CH_4 fluxes were higher at the low topography (Table 1) but were only statistically different in the dry season (H = 8.229; p

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<0.001). In the Macaca Island, With this the mean annual fluxes of CO₂ and CH₄ were $6.659 \pm 0.419 \text{ g CO}_2 \text{ m}^{-2} \text{ d}^{-1} \frac{\text{(mean} \pm \text{standard error)}}{\text{and } 0.132 \pm 0.053 \text{ g CH}_4 \text{ m}^{-2} \text{ d}^{-1},$ respectively.

3.33.2 Weather data

There was a marked seasonality during the study period (Figure 2), with 2,155.0 mm of precipitation during the rainy period and 1,016.5 mm during the dry period. As a consequence of the rains, the highest tides occurred in the period when the precipitation was greater (Figure 3). However, the rainfall distribution was different from the climatological average (Figure 23). The rainy season had 553.2 mm less precipitation, and the dry season had 589.1 mm more than the climatological normal. Thus, in the period studied, the dry season was rainier, and the rainy season was drier than the climatological normal, may already be a consequence of global climate change.



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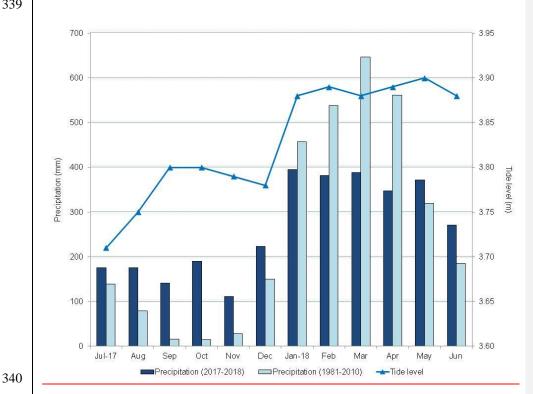
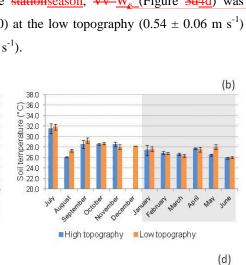
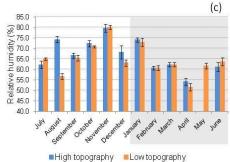


Figure 23. Monthly climatological normal in the municipality of Soure (1981-2010, mm), monthly precipitation (mm), and maximum tide height (m) for the from 2017 to 2018, in the municipality of São Caetano de Odivelas (PA).

 T_{air} was significantly higher (LSD = 0.72, p = 0.01) at the high topography (31.24 \pm 0.26 °C) than at the low topography (30.30 \pm 0.25 °C) only in the rainy season (Figure $\frac{3a4a}{a}$). No significant variation in T_s was found between the topographies in both seasons (Figure $\frac{3b4b}{}$). The 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 $\frac{3c4c}{}$). At this same $\frac{stationseason}{}$, $\frac{VV-W_s}{}$ (Figure $\frac{3d4d}{}$) was significantly higher (LSD = 0.15, p < 0.00) at the low topography (0.54 \pm 0.06 m s⁻¹) than at the high topography $(0.24 \pm 0.04 \text{ m s}^{-1})$.

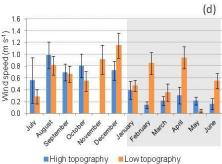
(a)





Low topography

■High topography



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Figure 34. 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.

3.43.3 Soil characteristics Environmental characterization

Silt concentration was higher at the low topography (LSD: 14.763; p= 0.007) and clay concentration was higher at the high topography plotsites (LSD: 12.463; p= 0.005), in both stations seasons studied (Table 21). Soil particle size analysis did not vary

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statistically (p > 0.05) between the two stations seasons (Table $\frac{21}{2}$). Soil moisture did not vary significantly (p > 0.05) between topographies at each stationseason, or between seasonal periods at the same topography (Table $\frac{21}{2}$). The variable pH varied statistically only at the low topography when the two stations seasons were compared (LSD: 5.950; p= 0.006), being more acidic in the dry period (Table $\frac{21}{2}$). On average pH was significantly (LSD: 0.559; p= 0.008) higher in the dry season (Table $\frac{21}{2}$). No variation in Eh was identified between topographies and seasons (Table $\frac{21}{2}$), 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 $\frac{21}{2}$). In addition, when comparing the two seasons, Sal was significantly higher in the dry season, both in the high (LSD: 2.916; p < 0.001) and in the low (LSD: $\frac{3.003}{2}$; p < 0.001) topography (Table 1). In addition, Sal was significantly higher in the dry season in both the high (LSD: 2.916; p < 0.001) and low (LSD: 3.003; p < 0.001) topographies (Table $\frac{21}{2}$).

Table $\frac{21}{2}$. Concentration analysis of Sand, Silt, Clay, Moisture, pH, Redox Potential (Eh) and salinity (Sal; ppt) in mangrove soil in the high and low topographies, and in the rainy and dry seasons, at Macaca island, São Caetano das Odivelas. Numbers represent the mean \pm (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 variation (LSD, p < 0.05).

C	Т1	Sand	Silt	Clay	Moisture		Eh	Sal
Season	Topography	(%)	(%)	(%)	pH (%)		(mV)	(ppt)
	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(<u>±</u> 45.53) ^a	35.25(±1.11) ^a A
Dry	Low	9.7 (± 2.5) ^{aA}	63.6 (± 6.1) ^{aA}	26.6(±5.2) ^{bA}	86.9 <u>(±</u> 3.4) ^{aA}	5.3 <mark>(±</mark> 0.3) ^{aA}	106.38 <mark>(±</mark> 53.76) ^a A	30.13 <u>(±</u> 1.16) ^b A
	Mean	10.9(<u>±</u> 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 <u>(±</u> 35.71)	32.69 <u>(±</u> 1.02)
	High	12.1 (± 1.4) ^{aA}	41.8 (± 3.3) ^{bA}	46.1(±2.6) ^{aA}	88.9 (± 3.5) ^{aA}	4.9 (± 0.4) ^{aA}	92.50 <u>(±</u> 56.20) ^{aA}	7.50(±0.78) ^{aB}
Rainy	Low	9.7 (± 2.5) ^{aA}	63.6 (± 6.1) ^{aA}	26.6 (± 5.2) bA	88.6 (± 3.7) ^{aA}	4.4(±0.1) ^{aB}	36.25 (± 49.97) ^{aA}	8.13 (± 0.79) ^{aB}
	Mean	10.9(<u>±</u> 1.4) ^A	52.7 (± 4.4) ^A	36.4 (± 3.8) ^A	88.7 (± 2.5) ^A	4.6 (± 0.2) ^B	64.38 <u>(±</u> 37.04) ^A	$7.81(\pm 0.54)^{B}$

380 The C_{mic} did not differ between topographies in the two seasons (Table $\frac{32}{2}$); however 381 CT-T_C was significantly higher in the low topography in the dry season (LSD: 5.589; p 382 < 0.000) and in the rainy season (LSD: 5.777; p = 0.024). In addition, C_{mic} was higher in 383 the dry season in both the high (LSD: 11.325; p < 0.010) and low (LSD: 9.345; p < 0.000) topographies (Table $\frac{32}{2}$). N_{mic} did not vary between topographies seasonally. 384 385 However, N_{mic} in the high (LSD: 9.059; p = 0.013) and low topographies (LSD: 4.447; 386 p = 0.001) was higher during the dry season (Table $\frac{32}{2}$). The C/N ratio (Table $\frac{32}{2}$) was 387 higher in the low topography in both the dry (LSD: 3.142; p < 0.000) and rainy seasons 388 (LSD: 3.675; p = 0.033), when compared to the high topography. However, only in the 389 low topography was the C/N ratio higher (LSD: 1.863; p < 0.000) in the dry season 390 compared to the rainy season (Table 32). Soil MO OM was higher at the low 391 topography in the rainy (LSD: 9.950; p = 0.024) and in the dry seasons (LSD: 9.630; p 392 < 0.000). However, only in the lowland topography was the MO-OM concentration 393 higher in the dry season than in the rainy season (Table $\frac{32}{2}$).

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Table 32. 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 (standard error). Lower case letters compare topography at each stationseason, and upper-case letters compare topography among between stationseasons.

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

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3.53.4 Vegetation structure and biomass

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 43). 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 variables did not show significant variation (Table 43). A total aboveground biomass of 322.1 ± 49.6 Mg ha⁻¹ was estimated.

G .	T 1	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(<u>±</u> 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 <u>(±</u> 29.3) ^{aB}	11.8 <u>(±</u> 8.8) ^{aA}	136.0 <u>(±</u> 108.3) ^{aA}
Total	High	350.2 (± 18.4)	325.6(±33.6) ^a	31.1 (± 7.5) ^a	304.5(<u>±</u> 99.8) ^a
10141	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: $R. mangle = 0.1282*DBH^{2.6}$; $A. germinans = 0.14*DBH^{2.4}$; Total = $0.168*\rho*DBH^{2.47}$, where $\rho_{R. mangle} = 0.87$; $\rho_{A. germinans} = 0.72$ (Howard et al., 2014).

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3.63.5 Drivers of greenhouse gas fluxes

In the rainy season, CO_2 efflux was correlated with T_{air} (Pearson = 0.23, p = 0.03), RH (Pearson = -0.32, p < 0.00) and T_s (Pearson = 0.21, p = 0.04) only at the low topography. In the dry season CO₂ flux was correlated with T_s (Pearson = 0.39, p < 0.00) at low topography. The dry season was the period in which we found the greatest amount of significant correlations between CO2 efflux and soil chemical parameters, while the C:N ratio, OM, and Eh were correlated with CO2 efflux in both seasons (Table 54). The negative correlation between T_C, N_T, C/N, and OM, along with the positive correlation of N_{mic} with soil CO₂ flux-(Table 5), in the dry period, indicates that microbial activity is a decisive factor for CO₂ efflux (Table 4). (Poungparn et al., 2009). Soil moisture in the Mojuim River mangrove forest negatively influenced CO2 flux in both seasons (Table 54). However a correlation between soil moisture with the flux of CH₄ was not identified. No significant correlations were found between CH₄ efflux and the chemical properties of the soil in the mangrove of the Mojuim River estuary (Table 54). However, with an average flux of 4.70 mg C m⁻² h⁻¹ and with extreme monthly and seasonal variation, more detailed studies are needed on CH₄ efflux and on the relationship with methanotrophic bacteria and interactions with abiotic factors (mainly ammonia and sulfate).

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Table 54. 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	<u>Season</u>	$\underline{\mathrm{T_{C}}}$	$\underline{\mathrm{T_{N}}}$	<u>C</u> mic	N _{mic}		<u>OM</u>	<u>Sal</u>	<u>Eh</u>		Moisture
(g m ⁻² d ⁻¹)		(g kg ⁻¹)	(g kg ⁻¹)	(mg kg ⁻¹)	(mg kg ⁻¹)	<u>C/N</u>	(g kg ⁻¹)	(ppt)	<u>(mV)</u>	<u>pH</u>	<u>(%)</u>
	Dry	<u>-0.68**</u>	<u>-0.59*</u>	0.18 ^{NS}	0.61**	<u>-0.66**</u>	<u>-0.67**</u>	<u>-0.07^{NS}</u>	<u>0.51*</u>	0.21 ^{NS}	<u>-0.49</u> *
<u>CO</u> 2	Rainy	<u>-0.44^{NS}</u>	<u>-0.20^{NS}</u>	<u>-0.15^{NS}</u>	<u>-0.32^{NS}</u>	<u>-0.50*</u>	<u>-0.63**</u>	<u>-0.54</u> *	0.53*	0.47 ^{NS}	<u>-0.54*</u>
	<u>Annual</u>	<u>-0.50**</u>	<u>-0.35</u> *	<u>-0.18^{NS}</u>	0.00^{NS}	<u>-0.53**</u>	<u>-0.48**</u>	-0.30 ^{NS}	0.39*	0.23 ^{NS}	<u>-0.56**</u>
	<u>Dry</u>	<u>0.30^{NS}</u>	<u>0.07^{NS}</u>	<u>-0.14^{NS}</u>	<u>-0.24^{NS}</u>	0.34 ^{NS}	0.02^{NS}	-0.04 ^{NS}	-0.38 ^{NS}	0.26 ^{NS}	0.26 ^{NS}
<u>CH</u> ₄	Rainy	<u>0.05^{NS}</u>	<u>-0.09^{NS}</u>	<u>0.44^{NS}</u>	<u>-0.27^{NS}</u>	0.09^{NS}	<u>-0.11^{NS}</u>	<u>-0.04^{NS}</u>	<u>-0.13^{NS}</u>	<u>-0.07^{NS}</u>	<u>0.04^{NS}</u>
	Annual	<u>0.04^{NS}</u>	<u>-0.10^{NS}</u>	<u>-0.01^{NS}</u>	<u>-0.18^{NS}</u>	0.08 ^{NS}	<u>-0.01^{NS}</u>	<u>-0.17^{NS}</u>	-0.21 ^{NS}	<u>-0.08^{NS}</u>	<u>0.02^{NS}</u>

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 $p \le 0.01$

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4 Discussion

4.1 Carbon dioxide and methane flux

It is important to consider that when compared with the climatological average (1981-2010), the year under study was rainier in the dry season (2017) and less rainy in the wet season (2018) It is important to consider that when compared with the elimatological average (1981-2010), the year under study was rainier in the dry season (2017) and less rainy in the wet season (2018)(Figure 23). Perhaps this variation is already related to the effects of global climate changes. Under these conditions, negative and positive flows of the two greenhouse gases were found (negative values represent gas consumption) Under these conditions, the CO₂ flux from the mangrove soil ranged from -5.06 to 68.96 g CO₂ m⁻² 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 d⁻¹ (mean 0.13 g CH₄ m⁻² d⁻¹), resulting in a total carbon rate of 1.92 g C m⁻² d⁻¹ or 7.00 Mg C ha⁻¹ y⁻¹ (negative values represent atmospheric consumption of the gas) (Figure 3). The negative CO₂ flux appears to be a consequence of the increased CO2 solubility in tidal waters, or of the increased sulfate reduction, which has already been described in the literature (Borges et al., 2018; Chowdhury et al., 2018; Nóbrega et al., 2016). Fluctuations in redox potential altered the availability of the terminal electron acceptor and donor and the forces of recovery of their concentrations in the soil, such that a disproportionate release of CO2 can result from alternative anaerobic degradation processes such as sulfate and iron reduction (Chowdhury et al., 2018). The soil carbon flux in mangrove area in the Amazon region was within the range of findings for other tropical mangrove areas (2.57 to 11.00 g CO₂ $m^{\text{--}2}\ d^{\text{--}1};$ Shiau and Chiu, 2020). However, the mean flux of 6.2 mmol CO $_2\ m^{\text{--}2}\ h^{\text{--}1}$ 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). We found a mean monthly flux of 327.9 ± 78.0 mg CO₂-m²-h⁻¹ and 217.2 ± 51.0 mg CO₂-m⁻²-h⁻¹, at the high and low topography, respectively. An emission of 0.010 Tg CH₄ y⁻¹, 0.64 g CH₄ m⁻² d⁻¹ (Rosentreter et al., 2018a), or 26.7 mg CH₄ m⁻² h⁻¹ is estimated at tropical latitudes (0 and 5°). In our study, the monthly average in CH₄ flux was higher at the low topography $(7.3 \pm 8.0 \text{ mg CH4 m}^{-2} \text{ h}^{-1})$ than

at the high topography $(0.9 \pm 0.6 \text{ mg C m}^{-2} \text{ h}^{-1})$, resulting in a total of 0.13 g CH4 m-2 d-1 or 0.48 Mg CH4 ha-1 y-1 (Figure 32). Therefore, the CH₄-C fluxes from the mangrove soil in the Mojuim River estuary were much lower than expected. It is known that there is a microbial functional module for CH4 production and consumption (Xu et al., 2015) and diffusibility (Sihi et al., 2018), which considers three key mechanisms: aceticlastic methanogenesis (acetate production), hydrogenotrophic methanogenesis (H2 and CO2 production) and aerobic methanotrophy (CH4 oxidation and O2 reduction), and this will be discussed below. The average emission from the soil of 8.4 mmol CH₄ m⁻² d⁻¹ was well below the fluxes recorded in the Bay of Bengal, with 18.4 mmol CH₄ m⁻² d⁻¹ (Biswas et al., 2007). In the Amazonian mangrove studied the mean annual carbon equivalent efflux was 429.6 mg CO_{2-eq} m⁻² h⁻¹. This value is 0.00004% of the erosion losses of 103.5 Tg CO_{2-eq} ha⁻¹ y⁻¹ projected 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 wettest, and the rainy season was drier when compared to the climatological normal.

4.2 Drivers of greenhouse gas fluxes Topography variation

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The 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 CO₂ efflux (Dai et al., 2012; Davidson et al., 2000; Ehrenfeld, 1995). In the two climatic periods of the year, the high topography produced more CO₂ (7,869 ± 1,873 g CO₂ m⁻² d⁻¹) than the low topography $(5,212 \pm 1,225 \text{ g CO}_2 \text{ m}^{-2} \text{ d}^{-1})$ (Figure $\frac{32}{2}$). No significant influence on CO₂ flux was observed due to the low variation in high tide level throughout the year (0.19 m) (Figure 32), 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 (7.858 \pm 0.039 g CO₂ m⁻² d⁻¹) than at the low topography (4.734 \pm 0.335 g CO₂ m⁻² d⁻¹) (Figure 32; Supplementary Information Table 1). This result is because the root systems of most flood-tolerant plants remain active when flooded (Angelov et al., 1996). Still, the high topography has longer flood-free periods, because this only happens when the tides are in the form of triangle or when the rains are torrential.

 CO_2 efflux was higher in the high topography than in the low topography, i.e., 39.8% lower in the forest soil exposed to the atmosphere for less time, in the rainy season (when soils are more subject to inundation). Measurements performed on 62 mangrove forest soils showed an average flux of 2.87 mmol CO_2 m⁻² h⁻¹ when the soil is exposed to the atmosphere, while 75 results on flooded mangrove forest soils showed an average emission of 2.06 mmol CO_2 m⁻² h⁻¹ (Alongi, 2007, 2009), i.e., 28.2% less than for the dry soil. Reflecting the more significant facility gases have for molecular diffusion than fluids, and the increased surface area for aerobic respiration and chemical oxidation during air exposure (Chen et al., 2010). Some studies attribute this variation to the temperature of the soil when exposed to tropical air (Alongi, 2009), increasing the export of dissolved inorganic carbon (Maher et al., 2018). However, although there was no significant variation in soil temperature between topographies at each time of year (Figure 4b), there was a positive correlation (Pearson = 0.15, p = 0.05) between CO_2 efflux and soil temperature at the low topography.

In the rainy season, CO_2 efflux was correlated with $T_{\rm air}$ (Pearson = 0.23, p = 0.03), RH (Pearson = -0.32, p < 0.00) and T_s (Pearson = 0.21, p = 0.04) only at the low topography. In the dry season CO_2 flux was correlated with T_s (Pearson = 0.39, p < 0.00) at low topography. Some studies show that CH_4 efflux is a consequence of the seasonal temperature variation in mangrove forest in temperate/monsoon climate (Chauhan et al., 2015; Purvaja and Ramesh, 2001; Whalen, 2005). However, in this study CH_4 efflux was correlated with Ta (Pearson = -0.33, p < 0.00) and RH (Pearson = 0.28, p = 0.01) only in the dry season and at the low topography. The results show that the physical parameters do not act in the fluxes in a standardized way, and their influence depends on the topography and seasonality. These results show that hardly does only one physical parameter interfere with the fluxes, and that they do not interact similarly in a different topography and seasonality.

A compilation of several studies showed that the total CH₄ emissions from soil in mangrove ecosystem range from 0 to 23.68 mg C m⁻² h⁻¹ (Shiau and Chiu, 2020), and our study showed a range of -0.01 to 31.88 mg C m⁻² h⁻¹, with a mean of 4.70 ± 5.00 mg C m⁻² h⁻¹. The monthly CH₄ fluxes were generally higher at the low (0.232 \pm 0.256) than at the high (0.026 \pm 0.018) topography, especially during the high tide—rainy season when the tides were higher (Figure 32). Compared to the high topography, only in the dry season was there a significantly higher production at the low topography

(Figure 32). The low topography produced 0.0249 g C m⁻² h⁻¹ more to the atmosphere in 535 the rainy season than in the dry season (Figure 32), and the same seasonal variation was 536 537 recorded in other studies (Cameron et al., 2021). 538 The mangrove soil in the Mojuim River estuary is rich in silt and clay (Table 21), which 539 reduces sediment porosity and fosters the formation and retention of anoxic conditions 540 (Dutta et al., 2013). In addition, the lack of oxygen in the flooded mangrove soil favor 541 microbial processes such as denitrification, sulfate reduction, methanogenesis, and 542 redox reactions (Alongi and Christoffersen, 1992). A significant amount of CH4 543 produced in wetlands is dissolved in the pore water due to high pressure resulting in a 544 significant supersaturation, which allows CH4 to be released by diffusion from the sediment to the atmosphere and by boiling through the formation of 545 bubbles Furthermore, plenty of the CH₄ produced in wetlands is dissolved in situ in the 546 pore water caused by the high pressure, which can result in supersaturation in the water, 547 enabling CH₄ to be released from the sediment to the atmosphere by diffusion and by 548 549 boiling in the water (Neue et al., 1997). 550 Only the species R. mangle and A. germinans were found in the floristic survey carried 551 out, which agrees with other studies in the same region (Menezes et al., 2008). Thus, the 552 variations found in the flux between the topographies in the Mojuim River estuary are 553 not related to the mangrove forest structure because there was no significant difference in the aboveground biomass. Since there was no difference in the species composition, 554 it is expected that the belowground biomass would not be different either (Table 4). 555 556 Soil moisture in the Mojuim River mangrove forest negatively influenced CO2 flux in both seasons (Table 5). However a correlation with the flux of CH₄ was not identified. 557 Studies show that CO₂ flux tends to be lower with high soil saturation (Chanda et al., 558 559 2014; Kristensen et al., 2008). A total of 395 Mg C ha⁻¹ was found at the soil surface (0.15 m) in the mangrove of the Mojuim River estuary, which was slightly higher than 560 the 340 Mg C ha⁻¹ found in mangroves in the Amazon (Kauffman et al., 2018), however 561 562 being significantly 1.8 times greater at the low topography (Table 32). The finer soil 563 texture at the low topography (Table 21) reduces groundwater drainage which facilitates 564 the accumulation of C in the soil (Schmidt et al., 2011).

4.3 Mangrove biomass

Only the species *R. mangle* and *A. germinans* were found in the floristic survey carried out, which agrees with other studies in the same region (Menezes et al., 2008). Thus, the variations found in the flux between the topographies in the Mojuim River estuary are not related to the mangrove forest structure because there was no significant difference in the aboveground biomass. Since there was no difference in the species composition, it is expected that the belowground biomass would not be different either (Table 3).

Assuming that the amount of carbon stored is 0.42 of the total biomass (Sahu and Kathiresan, 2019), the mangrove forest biomass of the Mojuim River estuary stores 127.9 and 138.9 Mg C ha⁻¹ at the high and low topography, respectively. This result is well below the 507.8 Mg C ha⁻¹ estimated for Brazilian mangroves (Hamilton and Friess, 2018), but are near the 103.7 Mg C ha⁻¹ estimated for a mangrove at Guará´s island (Salum et al., 2020), 108.4 Mg C ha⁻¹ for the Bragantina region (Gardunho, 2017), and 132.3 Mg Mg C ha⁻¹ in French Guiana (Fromard et al., 1998). The biomass found in the Mojuim estuary does not seem to be different from the biomass found in other Amazonian mangroves, however much lower than that found in other Brazilian mangroves. The estimated primary production for tropical mangrove forests is 218 ± 72 Tg C y (Bouillon et al., 2008).

4.4 Biogeochemical parameters

The dry season was the period in which we found the greatest amount of significant correlations between CO2 efflux and soil chemical parameters, while the C:N ratio, OM, and Eh were correlated with CO2-efflux in both seasons (Table 5). During the seasonal and annual periods, CH4 efflux was not correlated significantly with chemical parameters (Table 5), similar to the observed in another study (Chen et al., 2010). The soil waterlogging Increased soil moisture reduces gas diffusion rates, which directly affects the physiological state and microbial activities, by limiting the supply of the dominant electron acceptors, such as oxygen, and gases such as CH4 (Blagodatsky and Smith, 2012). The importance of soil moisture was evident in the richness and diversity of bacterial communities in a study comparing the different pore spaces filled with water (Banerjee et al., 2016). Furthermore, sulfate reduction in flooded soils (another pathway of organic matter metabolism) is dependent on the redox potential of the soil. However, no sulfate reduction occurs when the redox potential has values above -150

mv (Connell and Patrick, 1968). In our study, Eh was above 36.0 mV, this indicates indicating that sulfate reduction probably did not influence the OM metabolism.

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On the other hand, increasing soil moisture provides the microorganisms with essential substrates such as ammonium, nitrate, and soluble organic carbon, and increases gas diffusion rates in the water (Blagodatsky and Smith, 2012). Biologically available nitrogen often limits marine productivity (Bertics et al., 2010), and thus can affect the fluxes of CO₂ to the atmosphere. However, a mangrove fertilization experiment showed that CH4 emission rates were not affected by N addition (Kreuzwieser et al., 2003). A higher concentration of C_{mic} and N_{mic} in the dry period (Table $\frac{32}{2}$), both in the high and low topographies, indicated that microorganisms are more active when the soil spends more time aerated in the dry period (Table 32), the period in which only the high tides produce anoxia in the mangrove soil mainly in the low topography. the period in which the high tides produce anoxia in the mangrove soil. Additionally, the C/N ratio was well below 40, indicating that soil microorganisms and roots do not compete for nitrogen (Stevenson and Cole, 1999). Under reduced oxygen conditions in laboratory incubated mangrove soil, the addition of nitrogen resulted in a significant increase in the microbial metabolic quotient, but no concomitant change in microbial respiration, which was explained by a decrease in microbial biomass (Craig et al., 2021). Sulfate reducing bacteria (SO₄²) are important diazotrophs in coastal ecosystems and can contribute with significant nitrogen (N2) fixation in mangrove ecosystems (Bertics et al., 2010; Shiau et al., 2017; Welsh et al., 1996). The negative correlation between T_C, N_⊥, C/N, and OM, along with the positive correlation of N_{mic} with soil CO₂ flux (Table 5), in the dry period, indicates that microbial activity is a decisive factor for CO₂

can contribute with significant nitrogen (N₂) fixation in mangrove ecosystems (Berties et al., 2010; Shiau et al., 2017; Welsh et al., 1996). The negative correlation between T_C, N₄, C/N, and OM, along with the positive correlation of N_{mic} with soil CO₂ flux (Table 5), in the dry period, indicates that microbial activity is a decisive factor for CO₂ efflux (Poungparn et al., 2009). The high OM concentration at the two topographic heights (Table 32), at the two seasons studied, and the respective negative correlation with CO₂ flux (Table 5) confirm the importance of microbial activity in mangrove soil (Gao et al., 2020). Also, CH₄ produced in flooded soils can be converted mainly to CO₂ by the anaerobic oxidation of CH₄ (Boetius et al., 2000; Milucka et al., 2015; Xu et al., 2015) which may contribute to the higher CO₂ efflux in the Mojuim River estuary compared to other tropical mangroves (Rosentreter et al., 2018b). The belowground C stock is considered the largest C reservoir in mangrove ecosystem resulting from the low rate of OM decomposition due to flooding (Marchand, 2017).

Table 5. 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	∓ ∈	∓ _N	€ _{mie}	N _{mie}	C/N	OM	Sal	Eh		Moisture
(g m ² d ¹)		(g kg ⁺)	(g kg ⁺)	(mg kg ⁻¹)	(mg kg ⁻¹)	C/N	(g kg ¹)	(ppt)	(mV)	pH	(%)
	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 $p \le 0.01$

The higher water salinity in the dry season (Table 21) seems to result in a lower CH₄ flux at the low topography, more influenced by the tidal movement in this season (Dutta et al., 2013; Lekphet et al., 2005; Shiau and Chiu, 2020). Another essential factor for the reduced CH₄ emissions is when sulfate (SO₄²) in the brine affects the competition between SO₄²- reduction and methanogenic fermentation, because theas sulfate-reducing bacteria are more efficient at utilizing in hydrogen utilization than the methanotrophic bacteria (Abram and Nedwell, 1978; Kristjansson et al., 1982). At high SO₄²concentrations methanotrophic bacteria use CH₄ as an energy source and oxidize it to CO₂ (Coyne, 1999; Segarra et al., 2015), resulting in a consequent increasing increase the in CO2 efflux and reducing reduced the CH4 efflux (Megonigal and Schlesinger, 2002; Roslev and King, 1996). This may explain the high CO₂ efflux found throughout the year at the high and, especially, at the low topography (Figure 3). Only in the rainy season was a significant correlation recorded between salinity and CO2 flux. Still, in all seasonal periods the correlation between salinity and CO2 and CH4 fluxes were negative. Studies in other coastal ecosystems in Taiwan have recorded that methanotrophic bacteria can be sensitive to soil pH, and reported an optimal growth at pH ranging from 6.5 to 7.5 (Shiau et al., 2018). The higher soil acidity in the Mojuim River wetland (Table 21) may be inhibiting the activity of methanogenic bacteria by increasing the population of methanotrophic bacteria, which are efficient in consuming CH₄ consumption (Chen et al., 2010; Hegde et al., 2003; Shiau and Chiu, 2020). In addition, the pneumatophores present in R. mangle increase soil aeration and reduce CH₄ emissions (Allen et al., 2011; He et al., 2019). Spatial differences (topography) in CH₄ emissions in the soil can be attributed to substrate heterogeneity, salinity, and the abundance of methanogenic and methanotrophic bacteria (Gao et al., 2020). The high Eh values found in both topographies, mainly in the dry period (Table 21), are unfavorable for CH₄ emission. Soil Eh above -150 mV was considered limiting for CH₄ production (Yang and Chang, 1998). Increases in CH₄ efflux with reduced salinity were found due to intense oxidation or reduced competition from the more energetically efficient SO₄²⁻ and NO³⁻reducing bacteria than the methanogenic bacteria (Biswas et al., 2007). This fact can be observed in the CH₄ efflux in the mangrove of the Mojuim River, because in the rainy season (Figure 3), when there is a reduced water salinity

(Table 21) due to increased precipitation, there was an increased CH₄ production,

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- especially in the low topography (Figure 3). However, we did not find a correlation between CH₄ efflux and salinity, as already reported (Purvaja and Ramesh, 2001)
- 669 No significant correlations were found between CH₄-efflux and the chemical properties
- of the soil in the mangrove of the Mojuim River estuary (Table 5). However, with an
- 671 average flux of 4.70 mg C m⁻² h⁻⁴ and with extreme monthly and seasonal variation.
- 672 more detailed studies are needed on CH4 efflux and on the relationship with
- 673 methanotrophic bacteria and interactions with abiotic factors (mainly ammonia and
- 674 sulfate).

5 Conclusions

- 676 Between latitude 0° to 23.5° S the most recent estimate shows an emission of 2.3 g CO₂
- 677 m⁻² d⁻¹ (Rosentreter et al., 2018b). However, the efflux in the mangrove of the Mojuim
- 678 River estuary was 6.7 g CO₂ m⁻² d⁻¹. For the same latitudinal range, the Rosentreter et
- 679 <u>al. authors (2018c)</u> estimated an emission of 0.64 g CH₄ m⁻² d⁻¹, and we found an efflux
- 680 of 0.13 g CH₄ m⁻² d⁻¹. Seasonality was important for CH₄ efflux but did not influence
- 681 CO₂ efflux. Due to the rainfall variation compared to the climatology, that is, rainier in
- 682 the dry season and drier in the rainy season, the differences in fluxes may be an effect of
- 683 global climate changes on the terrestrial biogeochemistry at the plant-soil-atmosphere
- 684 interface, making it necessary to extend this study for more years. Using the factor of 23
- to convert the global warming potential of 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, the
- radiative forcing due to the continuous emission of 0.05 kg CH4 m-2 y-1, found in this
- work, would be offset if CO2 sequestration rates were 2.16 kg CO2 m-2 v-1 (Neubauer
- 689 and Megonigal, 2015).
- 690 Microtopography should be considered when determining the efflux of CO₂ and CH₄ in
- 691 mangrove forest in the Amazon estuary. The low topography in the mangrove forest of
- Rio Mojuim contained a higher concentration of organic carbon in the soil. However, it
- did not produce a higher CO₂ efflux because this was negatively influenced by soil
- moisture, which was indifferent to CH₄ efflux. MO, C/N ratio, and Eh were critical in
- soil microbial activity, which resulted in a variation in CO₂ flux during the year and
- 696 seasonal periods. In this sense, physicochemical properties of the soil are important for
- 697 CO₂ flux, especially in the rainy season; however, they did not influence CH₄ fluxes.

- 698 Data availability: The data used in this article belong to the doctoral thesis of Saul
- 699 Castellón, within the Postgraduate Program in Environmental Sciences, at the Federal
- 700 University of Pará. Access to the data can be requested from Dr. Castellón
- 701 (saulmarz22@gmail.com), which holds the set of all data used in this paper.
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