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1 **Carbon and nutrients accumulation in tropical mangrove creeks, Amazon region**

2

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21

22 **ABSTRACT**

23 The Marapanim River estuary (MRE) is part of the Amazon estuarine system located in  
24 northern Brazil, which is characterized as having extensive mangrove forests. Given that previous  
25 studies reported CO<sub>2</sub> and CH<sub>4</sub> fluxes from mangrove creeks in this region, here we investigate the

26 potential organic carbon sequestration of the creek mudflats to get a better understanding of the  
27 carbon cycling through these systems. Sediment accumulation rates derived from  $^{210}\text{Pb}$  dating  
28 indicated that sampled cores represent the previous 24 ( $\pm 4$ ) yr. The approximately 24-year total  
29 organic carbon (TOC), total nitrogen (TN) and total phosphorus (TP) burial rates were estimated to  
30 be 192.5 ( $\pm 43.5$ ), 15.3 ( $\pm 4.1$ ) and 3.2 ( $\pm 0.8$ )  $\text{g m}^{-2} \text{yr}^{-1}$ , respectively. A binary source mixing model  
31 based on carbon stable isotopes ( $\delta^{13}\text{C}$ ) revealed that the sedimentary organic matter (OM) is mainly  
32 influenced by marine phytoplankton input (49% to 95%). Furthermore, the TOC accumulation rates  
33 found here were slightly higher than the global averages estimated for within mangrove forests,  
34 suggesting that these unaccounted carbon sinks along creek mudflat environments are relevant for  
35 carbon budgets in mangrove-colonized coastal zones. The highest contents, stocks and accumulation  
36 rates were found in the tidal creek sediments that are most influenced by nearby mangroves and are  
37 more protected than sediments from major river margin. Our results indicate that the creek mudflats  
38 play a major role in carbon and nutrients sequestration, directly related to grain size and OM sources.

39

40 Keywords: Blue carbon, Organic matter source, Macrotidal estuary, creek mudflat, Brazilian  
41 Amazon coast.

42

### 43 **1. Introduction**

44

45 Mangrove forests, along with other vegetated coastal ecosystems such as seagrasses and  
46 saltmarshes, are recognized as blue carbon ecosystems due to their capacity to sequester carbon at a  
47 far higher rate than terrestrial forests (Nellemann et al., 2009; McLeod et al., 2011). Although  
48 mangrove forests occupy <1% of the global coastal area, these ecosystems sequester and store high  
49 amounts of organic carbon in plant biomass and sediment, contributing 10 - 15% to coastal sediment  
50 carbon storage and exporting 10 - 11% of the particulate terrestrial carbon to the ocean (Jennerjahn

51 and Ittekkot, 2002; Dittmar et al., 2006; Alongi, 2014). For instance, the carbon stocks in the Amazon  
52 mangrove are over twice those of upland evergreen forests and almost 10-fold those of tropical dry  
53 forests (Kauffman et al., 2018), which underscores their potential value to mitigating greenhouse gas  
54 emissions.

55 Two important measurements to determining the rate of organic carbon sequestration are the  
56 sediment total organic carbon (TOC) content and the sediment accumulation rate (SAR). The first  
57 term provides information about the TOC stock sequestered in sediments (Howard et al., 2014). The  
58 SAR is used to measure the TOC accumulation rates, which address the question of how much TOC  
59 is sequestered in a specified period and quantifies the ongoing sink capacity (Arias-Ortiz et al., 2018).  
60 From the  $^{238}\text{U}$  decay series, the  $^{210}\text{Pb}$  dating method has been an ideal tracer for dating aquatic  
61 sediments deposited during the previous 100 years, enabling the determination of TOC accumulation  
62 rates in vegetated coastal ecosystems (Smoak et al., 2013; Marchio et al., 2016; Sanders et al., 2016,  
63 Sasmito et al., 2020).

64 Total organic carbon to total nitrogen ratios TOC/TN ratios, and carbon and nitrogen stable  
65 isotope signatures ( $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$ ) have been widely used as effective geochemical proxies to estimate  
66 the relative proportions of terrigenous and marine OM in estuarine and coastal sediments (Lamb et  
67 al., 2006; Ranjan et al., 2011; Liu et al., 2015; Vilhena et al., 2018; Kusumaningtyas et al., 2019).  
68 Previous studies have shown that the terrestrial OM is preserved, stored and accumulated more  
69 efficiently than marine-derived OM in sediments (Hedges et al., 1997; Ranjan et al. 2011; Watanabe  
70 and Kuwae, 2015; Kusumaningtyas et al., 2019), due to the selective preservation of refractory OM  
71 at the expense of labile components, which is more susceptible to degradation by microorganisms  
72 (Hedges et al., 1997; Zonneveld et al., 2010). Therefore, identifying the source of OM is important  
73 in evaluating the effectiveness of the mangrove ecosystems as blue carbon sinks, since the variability  
74 of the origin of the OM stored in sediments contributes differently to longer-term carbon burial.

75 Data on the contribution of intertidal creek environments are more limited than mangrove  
76 forests as carbon sinks. However, intertidal mudflats situated along mangrove creeks can play an  
77 important role in the carbon cycling of tropical land-sea interfaces, such as acting as a conduit for  
78 exchanges between mangrove forests and coastal waters. For instance, Call et al. (2019) demonstrated  
79 that a mangrove tidal creek in the Amazon region presented large CO<sub>2</sub> and CH<sub>4</sub> fluxes, suggesting  
80 that recent global estimates of these fluxes based mostly on data from higher latitudes (exceeding 5°)  
81 without considering macrotidal mangrove system, may be underestimated. Moreover, tidal creeks  
82 may be more sensitive to land-use changes than open water systems due to a broader connectivity  
83 with watersheds, implying that OC accumulation rate changes by up to one order of magnitude in  
84 response to urban effluents input (Darrow et al., 2017). Therefore, the balance between carbon export,  
85 and sediment burial may present an important conduit to the mangrove and coastal ocean carbon  
86 cycle.

87 The Brazilian coastal region holds, over 960,000 ha, the third-largest mangrove area  
88 worldwide (Giri et al., 2011) including the North coast of Brazil (known as the Brazilian Amazon  
89 coast) which is considered the largest continuous and best-preserved mangrove forest in the world  
90 (Nascimento et al., 2013; Kauffman et al., 2018). The mangroves of the Marapanim River estuary  
91 (MRE) are part of this extensive range of mangroves on the Brazilian north coast, which present  
92 minimal anthropogenic impact. Mangroves are enormously important in this region as they sustain  
93 traditional communities found on the Amazon coast (Fernandes et al., 2018). Information on the  
94 biogeochemical cycling of carbon and nutrients in a non-impacted environment is necessary to better  
95 understand how potential changes to the mangrove dynamics may impact the carbon and nutrient  
96 fluxes between mangrove forests and coastal waters.

97 The main goal of this study is to obtain a better understanding of the cycling of sedimentary  
98 OM in the Marapanim mangrove estuarine system, located in the Brazilian Amazon region. To this  
99 aim, the hypothesis that creeks mudflats play an important role as carbon and nutrient sinks, rather

100 than only act as a conduit for carbon and nutrients cycling in coastal regions, was tested. We analyzed  
101 TOC, TN and TP stocks (sediment) and accumulation rates, as well as estimated the sources of OM  
102 in unvegetated mudflats situated along mangrove creeks to: 1) characterize the different OM sources,  
103 and 2) investigate potential relationships between OM sources and TOC, TN, and TP burial based on  
104 accumulation rates. In addition, we characterized the mangrove forest, according to topographical  
105 gradients of the tidal flats and vegetation heights, since biogeochemical cycles and carbon budget are  
106 also affected by the mangrove structure and environmental conditions (Chambers et al., 2013; Perez  
107 et al., 2018; Steinmuller et al., 2020). As such, we suggest that mangrove characteristics should be  
108 considered when comparing the biogeochemical data obtained here with other studies.

109

## 110 **2. Materials and methods**

111

### 112 *2.1. Study area*

113

114 The Marapanim River estuary is part of the Amazon estuarine system in Para, Northern Brazil,  
115 between 00°30' to 01°00'S and 47°32' to 47°00'W (Fig. 1a). The main River channel has a funnel  
116 form with a length of more than 70 km and a width of 8 km at the mouth (Atlantic Ocean) (Silva et  
117 al., 2009). The water of the estuary was extremely mixed as a result of tidal pumping and wave action,  
118 with the ocean water penetrating approximately 62 km up the estuary mouth during the dry season  
119 and 42 km during the wet season (Berrêdo et al., 2008). This system is dominated by macrotidal  
120 regime with semidiurnal tides; their amplitude ranges from 3.5 m during neap tide to over 6 m during  
121 spring tides.

122 The coastal region of Para is characterized by a tropical climate, with a wet season from  
123 January to June, and a dry season from July to December, high annual precipitation (2500–3000 mm)  
124 and average annual temperature of 27.7 °C (Martorano et al., 1993). The water temperature of the

125 estuary varies from 27 to 30 °C. pH values indicate alkaline conditions during the dry season (7.9 to  
126 8.0) and slight acidic during wet season (5.7 to 6.7). The salinity at low and high fluvial discharge  
127 varies from 24 to 3, respectively, in the estuarine channel (Berrêdo et al. 2008).

128 The estuary is part of the Master Lucindo Marine Extractivist Reserve, a protected  
129 Conservation Unit region. The estuary contains 130 km<sup>2</sup> of continuous and pristine mangrove forest  
130 (Vilhena et al., 2013). The dominant mangrove species are *Rhizophora* spp., *Avicennia germinans*,  
131 and *Laguncularia racemosa*. The trees are tall (up to approximately 35 m, Fig. 1d), generally  
132 distributed in mixed forests of *Rhizophora* spp. and *Avicennia germinans*. *Laguncularia racemosa* is  
133 found in the mangrove fringe. Saltmarsh vegetation is represented by *Spartina brasiliensis*, which  
134 occupies pioneering position in mudflat accretion areas.

135 The Marapanim catchment extends over an area of 2500 km<sup>2</sup> (Silva et al., 2009), with no  
136 industrial development, where about 28,000 people live. Like other Brazilian Amazon coastal cities,  
137 the Marapanim city's economy is based mainly on the sustainable use of natural resources,  
138 particularly fishing (crabs, shrimps, mollusks and fish), as well as commerce and tourism (Kjerfve  
139 and Lacerda, 1993; ICMBio, 2018; Fernandes et al., 2019).

140

## 141 2.2. Acquisition of drone images and processing

142

143 Very high resolution (3 cm) images of the study area were obtained using a Drone Phantom 4  
144 DJI (FC 330 digital 4K/12MP camera). The planialtimetric data were processed by the Agisoft  
145 Photoscan version 1.6.1 (AgisoftPhotoScan, 2018), and Global Mapper version 19 (GlobalMapper,  
146 2017). Planialtimetric data of ten ground control points were acquired by a smartphone connected to  
147 an Antenna Trimble Catalyst with a differential Global Navigation Satellite System (GNSS). A sub-  
148 metric correction ( $\pm 30$  cm), provided by the Trimble website upon payment of a subscription  
149 (<https://geospatial.trimble.com/catalyst-subscriptions>), was applied to the GNSS data. The vegetation

150 was manually classified by photointerpretation in the Global Mapper Software. *Rhizophora*,  
151 *Avicennia*, and *Laguncularia* trees were identified according to color, geometry, and texture of the  
152 canopy. This work followed procedures described by the software developer (Agisoft, 2018; “Global  
153 Mapper User’s Manual,” 2020) and adapted for mangrove areas (Cohen et al. 2018, 2019). A detailed  
154 description of the data processing can be obtained in the supplementary information section.

155

### 156 2.3. Field sampling

157

158 The sediment core collection was conducted during the wet (May 2017) and dry (Sept 2017)  
159 seasons. The sampling sites were situated near the mouth (Atlantic Ocean) of the MRE, along a  
160 lobular structure, where mangrove vegetation has developed. We compared tidal mangrove creeks  
161 (P2 and P3) with the Marapanim River margin (P1), which is located in an area under influence of a  
162 nearby sand bar (Fig. 1b).

163

#### 164 2.3.1. Sediment

165 One sediment core was collected from the unvegetated mudflat (devoid of macrophytes) at  
166 each site by inserting an acrylic tube (50 cm length) vertically into the substrate during low tide.  
167 Immediately after extraction, the sediment core was sectioned at 1-cm depth intervals from the core  
168 top to 6 cm depth, then 2-cm intervals until the 20 cm depth, finally at 5-cm intervals until the 35 cm  
169 depth. The sub-samples were bagged, preserved on ice and then transported to the laboratory. In  
170 addition, pore waters were retrieved using Rhyzon® collectors (Seeberg-Elverfeldt et al., 2005) at  
171 the same intervals to the solid phase for salinity analysis. Pore water salinity was measured in situ  
172 with a portable refractometer (Atago).

173

#### 174 2.3.2. Vegetation and phytoplankton



175 The phytoplankton and vegetation collection was conducted just during the wet (May 2017)  
176 season. Approximately 15 fresh leaves of six different adult trees of each dominant mangrove species  
177 (*Rhizophora* spp., *Avicennia germinans* and *Laguncularia racemose*) were collected. The leaves were  
178 washed in deionized water to remove the adhered sedimentary particles and salt, then the samples  
179 were frozen and lyophilized. The phytoplankton samples were collected along the mangrove tidal  
180 creek during the flood tides (n=3); horizontal hauls were performed on the water surface (maximal  
181 depth 50 cm), using a standard-type plankton mesh, with mesh opening 64  $\mu\text{m}$ . This mesh opening is  
182 the most used one in the Amazon estuaries due to the strong local hydrodynamics (Paiva et al., 2006).  
183 Each sample was observed in a binocular microscope after it was washed in deionized water and  
184 subjected to wet sieving (mesh opening 20  $\mu\text{m}$ ) to remove the contaminant particles (leaves and  
185 shells) or possible zooplankton predators. Finally, the samples were frozen and lyophilized.

186

## 187 2.4. Analyses and data treatment

188

### 189 2.4.1. DBD and Grain size

190 Dry bulk density (DBD,  $\text{g cm}^{-3}$ ) was determined as the dry sediment weight (g) divided by the  
191 initial volume ( $\text{cm}^3$ ). From the original wet section, a portion was taken for grain size analysis. The  
192 sediment grain size was determined using a Fritsch particle size meter model Analysette 22, after  
193 calcium carbonate and OM removal, and dispersion in sodium hexametaphosphate 4% (Loring and  
194 Rantala, 1992). The GRADISTAT 9.1 software (Blott and Pye, 2001) was used for treating results,  
195 where the grain size scale was modified from Udden (1914) and Wentworth (1922), with the  
196 classification of clay ( $< 2 \mu\text{m}$ ), silt (2-63  $\mu\text{m}$ ) and sand ( $> 63 \mu\text{m}$ ) are established for these fractions.

197

### 198 2.4.2. Elemental and stable isotope analysis

199 Carbon and nitrogen stable isotope ratios of phytoplankton, leaves and sediments were  
200 measured to identify the sources of OM contributing to the sediments column at each site. A  
201 subsample of each core fraction was acidified to remove carbonate material; then it was washed in  
202 deionized water, dried (60 °C) and ground to powder before TOC and  $\delta^{13}\text{C}$  analyses. TN and  $\delta^{15}\text{N}$   
203 were analyzed in non-acidified subsamples. Isotopic signatures ( $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$ ) and TOC and TN  
204 contents were analyzed using a Leco Flash Elemental Analyser coupled to a Thermo Fisher Delta V  
205 isotope ratio mass spectrometer (Thermo Flash EA 1112). Analytical precision was as follows: TOC  
206 = 0.1%, TN = 0.1%,  $\delta^{13}\text{C}$  = 0.1‰, and  $\delta^{15}\text{N}$  = 0.15‰. Working standards were used (glucose, 10.7  
207 ppt and urea, 41.3 ppt) to calibrate for  $\delta^{13}\text{C}$ . A pair of standards were measured with every 20 samples.  
208 These standards were calibrated initially against international absolute standards LSVEC and  
209 NIST8542. TP contents in the sediments were determined by colorimetry, according to Grasshoff et  
210 al. (1999), following the extraction procedure from Aspila et al. (1976). The results of TOC, TN and  
211 TP, expressed in % in this work, were converted to  $\mu\text{mol g}^{-1}$  dry weight sediment to calculate  
212 individual TOC/TN/TP molar ratios.

213

#### 214 2.4.3 Calculation of marine and terrestrial organic matter

215 The relative proportions of mangrove/terrestrial OM ( $\text{OM}_{\text{terr}}$ ) and marine OM ( $\text{OM}_{\text{mar}}$ ) in  
216 sediment cores were estimated using the two end-member mixing model described in Schultz and  
217 Calder (1976):

218

$$219 F_{\text{terr}} = (\delta^{13}\text{C}_{\text{mar}} - \delta^{13}\text{C}_{\text{sed}})/(\delta^{13}\text{C}_{\text{mar}} - \delta^{13}\text{C}_{\text{terr}}) \times 100 \text{ and } F_{\text{mar}} = 100 - F_{\text{terr}} \quad (1)$$

220

221 where  $F_{\text{terr}}$  is the contribution from the mangrove/terrestrial fraction,  $F_{\text{mar}}$  is the contribution from the  
222 marine fraction,  $\delta^{13}\text{C}_{\text{sed}}$  value of the sediment interval, and  $\delta^{13}\text{C}_{\text{terr}}$  and  $\delta^{13}\text{C}_{\text{mar}}$  are the terrestrial and  
223 marine end-members value, respectively. We estimated the terrestrial (an average of the plants that

224 dominate in the Marapanim mangrove) and marine (phytoplankton-derived OM) end-member  $\delta^{13}\text{C}$   
225 values to be  $-29.9 \pm 1.0 \text{ ‰}$  ( $n = 18$ ) and  $-23.9 \pm 0.7 \text{ ‰}$  ( $n = 3$ ), respectively, from our results.

226

#### 227 2.4.4. $^{210}\text{Pb}$ dating and rates of organic carbon, nitrogen and phosphorus accumulation

228 To date sediments, we measured radionuclide activities from the  $^{238}\text{U}$  decay series in a high-  
229 purity germanium (HPGe) well gamma detector with 40% efficiency coupled to a multichannel  
230 analyzer. Sediments at each interval were sealed in gamma tubes for at least three weeks to establish  
231 secular equilibrium between  $^{226}\text{Ra}$  and its daughter products  $^{214}\text{Pb}$  and  $^{214}\text{Bi}$ . The  $^{210}\text{Pb}$  activities were  
232 determined by the direct measurement of 46.5 KeV gamma peaks, while  $^{226}\text{Ra}$  activity was calculated  
233 averaging its daughters' peaks  $^{214}\text{Pb}$  and  $^{214}\text{Bi}$  (295.2 KeV) (351.9 KeV) (609.3 KeV) (Sanders et al.,  
234 2016). The excess  $^{210}\text{Pb}$  ( $^{210}\text{Pb}_{\text{ex}}$ ) activity was estimated by subtracting the  $^{226}\text{Ra}$  from the total  $^{210}\text{Pb}$   
235 activity. The sediment accumulation rate (SAR) was calculated according to the Constant Initial  
236 Concentration (CIC) method as a net downcore decrease in  $^{210}\text{Pb}_{\text{(ex)}}$  activities was noted in all three  
237 cores, implying a consistent rate of sedimentation (Appleby and Oldfield, 1992). Accumulation rates  
238 and stocks (for 0-35 cm) of TOC, TN and TP were estimated for each depth interval (cm), using  
239 values of SAR ( $\text{cm yr}^{-1}$ ), DBD ( $\text{g cm}^{-3}$ ), and TOC, TN as well as TP contents ( $\text{g g}^{-1}$ ), respectively:

$$240 \text{ Accumulation rates (AR) (g m}^{-2} \text{ yr}^{-1}) = [\text{SAR}] \times [\text{DBD}] \times [\text{TOC, TN or TP content}] \quad (2)$$

$$241 \text{ Stocks (g m}^{-2}) = [\text{DBD}] \times [\text{depth}] \times [\text{TOC, TN or TP content}] \quad (3)$$

242

#### 243 2.4.5. Statistical analysis

244 Statistical analyses were performed using statistical package PAST version 3.26 (Hammer et  
245 al., 2001). One-way analysis of variance (ANOVA) was used to assess the significant difference of a  
246 single variable (clay, silt, sand, TOC, TN, TP,  $\delta^{13}\text{C}$ ,  $\delta^{15}\text{N}$ ) between the sampling sites (P1, P2 and P3)  
247 and seasons (wet and dry), with a Tukey HSD post hoc test for distinguishing pairwise relationships  
248 among sites. The normality of data distribution was tested using Shapiro-Wilk prior to statistical

249 analysis. When the variables were not normally distributed (clay, silt, sand, TOC, TN and TP), they  
250 were log-transformed to fit a normal distribution. Statistical significance at  $\alpha < 0.05$  was used for all  
251 tests. Principal Components Analysis (PCA) was applied to identify the multivariate relationships  
252 between the geochemical variables. Significant factors were selected based on eigenvalues  $> 1$ . The  
253 relation between each pair of variables was measured by Pearson's correlation coefficient. Correlation  
254 coefficients greater than 0.5 were considered significant.

255

### 256 **3. Results**

257

#### 258 *3.1. Mangrove structure*

259

260 The A-B profile (Fig. 1d-e) revealed a young mangrove fringe (1-10 m) mainly represented  
261 by *Laguncularia racemosa* in the topographically lowest sector (~1 m above mean sea-level, amsl)  
262 of the tidal flat. In an intermediate topography (1 – 2 m amsl) a dense mangrove forest characterized  
263 by *Laguncularia racemosa* and *Rhizophora* spp. (10 - 20 m tall) was found. Following the  
264 topographical gradient (~2 m amsl) a mixed forest of *Avicennia germinans* and *Rhizophora* spp. is  
265 noted (20 - 30 m tall). These trees become taller (25-33 m) in the inner parts of this mangrove forest.  
266 The sampling sites P2 and P3 (Fig. 1b) were located in topographically highest areas, inundated only  
267 during spring tides, of which sediments are strongly oxidized during the dry season, coexisting with  
268 more mature mangroves (15-25 m).

269

#### 270 *3.2. Sediment chronology*

271

272 The  $\log^{210}\text{Pb}_{\text{ex}}$  of the sedimentary profiles from the three examined sampling sites were  
273 depicted by an almost linear decline with depth (Fig. 2), implying a consistent rate of sedimentation.

274 Based on the  $^{210}\text{Pb}_{(\text{ex})}$  profiles, the top 6 cm of the P3 and bottom of the P2 sediment cores were  
275 determined to be mixed and therefore, were excluded when calculating SAR. The SAR were  
276 calculated as 1.8 (core P1), 1.5 (core P3) and 1.3 (core P2)  $\text{cm yr}^{-1}$ , respectively (Fig. 2), with a  
277 regional mean of  $1.5 \pm 0.3 \text{ cm yr}^{-1}$ . According to the extrapolated  $^{210}\text{Pb}_{\text{ex}}$  - derived age, sediment at  
278 the core basis corresponds to an age of about 1997 year in P1 (35 cm), 1989 year in P2 (35 cm) and  
279 1994 in P3 (35 cm).

280

### 281 *3.3. Grain size and dry bulk density*

282

283 The textural composition of the sediments was mostly sand and silt (34–80% of silt, 12–64%  
284 of sand) with a low percentage of clay (<10%) (Fig. 3). This textural distribution differs among the  
285 three sites ( $p < 0.05$ , Table 2), with the distribution of fine-grained higher at sites P2 and P3 compared  
286 with P1 (Fig. 3). The contents of the grain size fractions also showed significant seasonal variations  
287 (Table 2), except to silt at P3 and sand at P1 and P2. In general, values of DBD ( $\text{g cm}^{-3}$ ) in the sediment  
288 sites increased substantially with the depth (Fig. 3). The highest DBD was found in the P1, exceeding  
289  $1.0 \text{ g cm}^{-3}$ , where the sedimentary profile graded to predominately sand.

290

### 291 *3.4. Elemental and isotopic composition*

292

293 The values and vertical profiles of elemental and isotopic composition for the sediments,  
294 plants and phytoplankton are shown in Table 1 and Fig. 3. Differences in TOC, TN, TP contents of  
295 sediments between sampling locations and seasons (except to TN at site P3) were statistically  
296 significant (Table 2). The range of TOC was highest at sites P2 (2.28 to 3.84 %) and P3 (2.21 to 3.04  
297 %) compared to P1 (0.53 to 1.91 %). The range of TN was similar to P2 (0.19 to 0.27 %) and P3 (0.19  
298 to 0.26%) compared to P1 (0.04 to 0.17 %). Like TOC and TN, TP contents was much larger in the

299 P2 (0.028 to 0.052 %) and P3 (0.033 to 0.059 %) than to P1 (0.009 to 0.044%). In all three sites,  
300 TOC, TN and TP contents decreased slightly with depth.

301 The  $\delta^{13}\text{C}$  values of sediments varied from -26.9 to -24.2‰, with significant difference between  
302 sites ( $p < 0.05$ ). The downcore  $\delta^{13}\text{C}$  profile was relatively homogeneous at site P1, without significant  
303 difference between seasons ( $p > 0.05$ ), with an average of  $-25.2 \pm 0.4\text{‰}$ . However, P2 and P3 showed  
304 significant differences between seasons ( $p < 0.05$ ), due to results from upper 18 cm, with highest values  
305 during dry season, with average of  $-25.7 \pm 0.6\text{‰}$  and  $-25.2 \pm 0.5\text{‰}$ , compared to wet season, with  
306 average of  $-26.6 \pm 0.2\text{‰}$  and  $-26.3 \pm 0.2\text{‰}$ , respectively.

307 The  $\delta^{15}\text{N}$  values of sediments did not differ significantly between seasons ( $p > 0.05$ ), but  
308 differed significantly between sites ( $p < 0.05$ ). The  $\delta^{15}\text{N}$  values ranged from 5.1 to 8.4‰, with an  
309 average of  $6.1 \pm 0.8\text{‰}$  to P1,  $5.9 \pm 0.6\text{‰}$  to P2 and  $6.5 \pm 0.3\text{‰}$  to P3. Downcore profiles of  $\delta^{15}\text{N}$   
310 showed a small increase with depth, excepted for site P1 that showed oscillations and no tendency  
311 with depth, with a peak at 14 cm.

312 Differences in TOC/TN molar ratios between sampling locations and seasons were  
313 statistically significant (Table 2), varying from 12 to 18.5, with an average of  $15.3 \pm 1.3$  to P1,  $15.6$   
314  $\pm 1.3$  to P2 and  $13.7 \pm 0.8$  to P3. Downcore profiles of TOC/TN showed a small increase with the  
315 depth at all sites. The TN/TP molar ratios in sediments were not significantly different in terms of  
316 spatial variability ( $p > 0.05$ ), but were significantly different seasonally, with highest ratios in the wet  
317 season, varying from 8.3 to 13.3, compared to dry season, that varied from 2.9 to 11.8.

318 The  $\delta^{13}\text{C}$  values of *Rhizophora* spp., *Avicennia germinans* and *Laguncularia racemosa* leaves  
319 were similar and ranged from -31.8 to -28.2‰ ( $29.8 \pm 1.0\text{‰}$ ), which is within the range of  $\text{C}_3$   
320 terrestrial plants. The  $\delta^{15}\text{N}$  and TOC/TN values varied slightly from species to species from 1.7 to  
321 6.4‰ ( $4.4 \pm 1.3\text{‰}$ ) and from 30.0 to 40.1 ( $34.0 \pm 3.2$ ), respectively. The phytoplankton presented  
322  $\delta^{13}\text{C}$  values from -24.4 to -23.4‰ ( $23.9 \pm 0.7\text{‰}$ ),  $\delta^{15}\text{N}$  from 3.2 to 3.7‰ ( $3.4 \pm 0.4\text{‰}$ ), and TOC/TN  
323 from 6.9 to 7.2 ( $7.1 \pm 0.2$ ).

324

### 325 *3.5 Principal Component Analysis (PCA)*

326

327 The significant components (ie., eigenvalue >1) loading matrix of PCA are listed in Table 3.

328 The PCA (Fig. 4) showed that the first two components together explained 74.5% of the data

329 variation. The first component (PC1) accounted for the largest proportion with 61.7% of the total

330 variance. It showed significant positive loading (> 0.5) for TN, TOC, TP, silt and clay, and negative

331 loading of sand and  $\delta^{13}\text{C}$ . The first component explained the variations between the sites, separating

332 the site P1 from P2 and P3 due to its high sand content (Fig. 4a). This component also gives strong

333 evidence of TOC, TN and TP increase with increase of silt and clay, with the highest TOC, TN and

334 TP contents for P2 and P3 (Fig 4b). The second component (PC2) explained 12.9 % of the total

335 variance and showed significant loading only for  $\delta^{15}\text{N}$ .

336

### 337 *3.6. Proportions of organic matter sources*

338

339 The proportions of terrestrial ( $\text{OM}_{\text{terr}}$ ) and marine ( $\text{OM}_{\text{mar}}$ ) OM sources were derived from the

340  $\delta^{13}\text{C}$  data and Eq. (1). Downcore profiles of  $\text{OM}_{\text{mar}}$  and  $\text{OM}_{\text{terr}}$  remained relatively constant

341 throughout the sediment columns during the wet season, but during the dry season an increase in

342 marine input in the upper layers (< 18 cm) was observed at sites P2 and P3 (Fig. 3). Relatively high

343 proportions of  $\text{OM}_{\text{mar}}$  were observed at all sites in both wet and dry seasons. Overall,  $\text{OM}_{\text{mar}}$  varied

344 from 49.1 to 95.2% and  $\text{OM}_{\text{terr}}$  from 4.8 to 50.9% (Table 1). In the wet season,  $\text{OM}_{\text{mar}}$  contribution

345 was highest in the site P1 (76.4 %) compared to sites P2 (54.5 %) and P3 (60 %). During dry season,

346  $\text{OM}_{\text{mar}}$  increased to 79.6, 69 and 78 % at sites P1, P2 and P3, respectively.

347

### 348 *3.7. Stocks and accumulation rates*

349

350 The TOC, TN and TP accumulation rates and stocks are shown in Fig. 5 and Table 4. The  
351 TOC, TN, and TP accumulation rates were estimated from SAR, DBD and TOC, TN and TP content  
352 results (Eq. (2)). The downcore sediment densities along with the TOC, TN and TP contents were  
353 used to determine stocks (Eq. (3)), the stocks were calculated to 30 cm sediment depth for the three  
354 sampling sites. The TOC, TN and TP accumulation rates showed significant site-difference ( $p < 0.05$ ),  
355 with a regional mean of  $192.5 \pm 50.6 \text{ g m}^{-2} \text{ y}^{-1}$  for TOC,  $15.3 \pm 4.3 \text{ g m}^{-2} \text{ y}^{-1}$  for TP and  $3.2 \pm 1.0$  for  
356 TP. The highest TOC, TN, and TP accumulation rates were measured at sites P2 (C: 218.7, N: 16.4,  
357 and P:  $3.1 \text{ g m}^{-2} \text{ yr}^{-1}$ ) and P3 (C: 215.8, N: 18.4, and P:  $3.9 \text{ g m}^{-2} \text{ yr}^{-1}$ ) and the lowest were measured  
358 at site P1 (C: 143.2, N: 11.0, and P:  $2.7 \text{ g m}^{-2} \text{ yr}^{-1}$ ). The TOC, TN and TP stocks also differed  
359 significantly among the three sites ( $p < 0.05$ ), with a regional mean of  $3811 \pm 1389 \text{ g m}^{-2}$  for TOC,  $296$   
360  $\pm 109 \text{ g m}^{-2}$  for TN and  $58 \pm 15 \text{ g m}^{-2}$  for TP. Similarly, the TOC, TN and TP stocks were higher at  
361 sites P2 (C: 5217.8, N: 386.3, P:  $64.9 \text{ g m}^{-2}$ ) and P3 (C: 3878.6, N: 325.8, P:  $55.4 \text{ g m}^{-2}$ ) than P1 (C:  
362  $2337.5$ , N: 177.1, P:  $17.1 \text{ g m}^{-2}$ ).

363

## 364 **4. Discussion**

365

### 366 *4.1. Sedimentary composition*

367

368 The mudflat sediments were found to be mainly composed of a mixture containing silt and  
369 sand, reflecting a moderately high energetic environment, influenced by tide and fluvial process. The  
370 low hydrodynamics flow in the mangrove creeks (P2 and P3) likely caused an increase in the  
371 deposition of fine sediments. However, the MRE margin, where P1 was collected, appears to be more  
372 affected by tidal currents. In addition, the P1 site is located closer to the sand bar that likely  
373 contributed to the increase of fine sand in the sedimentary profile at this study site.



374 The spatial variations in the grain size distribution of the mudflat sediments play an important  
375 role in controlling the OM content. Silt and clay had positive strong to moderate correlation with  
376 TOC, TN and TP contents ( $r$  varied from 0.52 to 0.81). The first component of the PCA also indicated  
377 strong evidence that TOC, TN and TP increase as the silt and clay increase, with the highest TOC,  
378 TN and TP contents for P2 and P3 (Fig 4a-b). Generally, fine-grained (silt + clay) sediments have  
379 higher %TOC than coarse sediments (Canfield, 1994), such relationships may be attributed to the  
380 fine-grained sediments which have large specific surface areas that provide higher capacity to adsorb  
381 OM (Loring and Rantala, 1992; Mayer, 1994).

382 The TOC and TN contents measured in the sediments during the present study are comparable  
383 to those reported in the adjacent coastal areas (Kauffman et al. 2018; Vilhena et al., 2018) and the TP  
384 contents are within the global average for mangrove (0.01–0.16%; Alongi et al., 1992). The gradual  
385 decrease of TOC, TN and TP content with depth in all cores likely reflect the decomposition of OM  
386 by microorganisms as noted in other systems (Kristensen et al., 2008) or a general increase in the OM  
387 content with time. The significant positive correlation between TOC and TN ( $r= 0.96$ ) and TP ( $r=$   
388  $0.83$ ) in the sediments indicate that the nitrogen and phosphorous in the samples are predominantly  
389 associated with the organic fraction.

390 The TOC/TN/TP molar ratios in sediments were not significantly different in terms of spatial  
391 variability, but were significantly different seasonally: 171:11:1 in the wet season and 71:5:1 in the  
392 dry season. In both cases the TOC/TN/TP was different from the phytoplankton sources (Redfield:  
393 106:16:1), with TN/TP ratios below 16 that suggest nitrogen to be the limiting factor for  
394 phytoplankton growth in these systems. This seasonal variation may be related to the diagenesis of  
395 OM. Higher TOC/TN molar ratios in wet than dry season indicate a greater contribution from OM of  
396 terrestrial origin, which is more refractory, during the wet season.

397

398 *4.2. Sources of sedimentary organic matter*

400 Different sources of OM often exhibit distinct elemental (carbon and nitrogen) and/or isotopic  
401 signatures ( $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$ ), representing useful indicators to quantify their relative contribution to the  
402 sedimentary OM. For example,  $\text{C}_3$  plants are assumed to have  $\text{TOC}/\text{TN} > 12$  (Meyers, 1997),  $\delta^{13}\text{C}$   
403 between  $-32\text{‰}$  and  $-21\text{‰}$  (Deines, 1980) and  $\delta^{15}\text{N}$  around  $0.4 \pm 0.9\text{‰}$  (Peterson and Howard, 1987),  
404 while marine-derived OM is characterized by  $\text{TOC}/\text{TN}$  between 5 and 7 (Redfield et al., 1963),  $\delta^{13}\text{C}$   
405 from  $-16\text{‰}$  to  $-23\text{‰}$  (Meyers, 1994) and  $\delta^{15}\text{N}$  around  $8.6 \pm 1.0\text{‰}$  (Peterson and Howard, 1987).

406 The potential sources to the sedimentary OM pool in mangrove-estuarine ecosystems can be  
407 upland forest and mangrove tissues (fresh leaf, stem, root, and litter), soils from river flow, aquatic  
408 macrophytes, microphytobenthos, and phytoplankton (Bouillon et al. 2008; Sasmito et al., 2020).  
409 Some studies have also identified the presence of microphytobenthos as a potential source of OM in  
410 mudflats (Gontharet et al., 2014; Gorman et al., 2020), when microphytobenthos layers are visually  
411 detectable and thick enough to be sampled and separated from the sediments. However, the presence  
412 of microphytobenthos was not visually detectable in our sampling sites, therefore prohibiting the  
413 separation and analyses of a potential benthic microalgae endmember. Saltmarsh vegetation (*Spartina*  
414 spp.), dominated by  $\text{C}_4$  plants, can be found at the mudflats along MRE. However, given that *Spartina*  
415 spp. and their sediments have enriched  $\delta^{13}\text{C}$  signature, ranging from  $-14$  to  $-12\text{‰}$  and  $-18$  to  $-14\text{‰}$ ,  
416 respectively (Currin et al., 1995; Kemp et al., 2020, and references therein), the depleted  $\delta^{13}\text{C}$  values  
417 found in sediment samples suggests that *Spartina* spp. is not a major contributor of OM to the studied  
418 mudflat sediments. Therefore, we assumed mangrove plants and phytoplankton as the principal OM  
419 sources to the mudflats along MRE.

420 The  $\delta^{13}\text{C}$  values of mangrove leaves ( $-31.8$  to  $-28.2\text{‰}$ ) are consistent with previously  
421 published results from other mangrove-estuarine ecosystems (Bouillon et al., 2008; Prasad and  
422 Ramanathan 2009; Ranjan et al., 2011; Vilhena et al., 2018). The  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  values and  $\text{TOC}/\text{TN}$   
423 molar ratio of phytoplankton (average of  $-23.9 \pm 0.7\text{‰}$ ,  $3.4 \pm 0.4\text{‰}$  and  $7.1 \pm 0.2$ , respectively), which

424 we extracted along the studied tidal creek, are close to the marine phytoplankton values. The  $\delta^{13}\text{C}$   
425 values are also similar to the reported value for marine DOC (average  $-23.7\text{‰}$ ) extracted from  
426 Atlantic deep water in the adjacent study area (Dittmar et al., 2006).

427 The  $\delta^{13}\text{C}$  ( $-26.9$  to  $24.2\text{‰}$ ) versus TOC/TN ( $12.0$  to  $18.5$ ) (Fig. 6) of the sedimentary OM  
428 suggests a mixture of sources contributing to the sedimentary OM pool, with a higher contribution of  
429 marine phytoplankton ( $\delta^{13}\text{C}$ :  $-23.9 \pm 0.7 \text{‰}$ , TOC/TN:  $7.1 \pm 0.2$ ) than mangrove-derived OM ( $\delta^{13}\text{C}$ :  
430  $-29.8 \pm 1.0\text{‰}$ , TOC/TN:  $34 \pm 3.1$ ). The  $\delta^{15}\text{N}$  values of sediments ( $5.2 - 8.4\text{‰}$ , average of  $6.2\text{‰}$ ) are  
431 within a typical range of aquatic OM produced from assimilation of the nitrate pool. The slight  
432 increase of  $\delta^{15}\text{N}$  values and TOC/TN molar ratios with depth, with exception to site P1 in the wet  
433 season, suggests anaerobic microbial degradation of OM in the deep reducing sediments (Meyers,  
434 2003; Routh et al., 2009; Jennerjahn, 2012; Prasad et al., 2017). The peak of  $\delta^{15}\text{N}$  at 14 cm in the site  
435 P1 during wet season could reflect an event provoked by input from a phytoplankton bloom or  
436 accumulation of allochthonous material deposited during and after a rainfall.

437 The consistently similar values of the  $\delta^{13}\text{C}$  throughout the sediment column suggest that the  
438 source of the depositional OM has been invariable during the past  $\sim 25$  years at site P1. However, at  
439 sites P2 and P3, the enrichment of  $\delta^{13}\text{C}$ , during the dry season compared to wet season, indicated a  
440 shift in OM source, because the decomposition of OM usually does not cause a significant enrichment  
441 of  $\delta^{13}\text{C}$  (Saintilan et al., 2013), and isotopic fractionation during the decomposition is  $<3\text{‰}$ . This shift  
442 in OM accumulation may be due to an increase in marine input in the upper layer ( $< 18$  cm), likely  
443 associated with a larger sediment mixing provoked by bioturbation in the surface sediments, as the  
444  $\delta^{13}\text{C}$  values were relatively similar along the bottom layers.

445 The difference in OM source for terrestrial/mangrove  $\text{C}_3$  plants ( $\delta^{13}\text{C}$ :  $-29.8 \pm 1.0 \text{‰}$ ) vs.  
446 marine phytoplankton ( $\delta^{13}\text{C}$ :  $-23.9 \pm 0.7 \text{‰}$ ) is distinct and hence, suitable for indicating the OM  
447 sources in the study area. However,  $\delta^{15}\text{N}$  values and TOC/TN molar ratios can be unreliable because  
448 these are influenced by diagenetic alterations (Prah et al., 1997). Additionally,  $\delta^{13}\text{C}$  did not show

449 significant positive correlations with  $\delta^{15}\text{N}$  ( $r < 0.08$ ) and TOC/TN molar ratios ( $r < 0.05$ ) in either the  
450 wet or dry season. Therefore, only  $\delta^{13}\text{C}$  was used for the quantification analysis of OM. Based on the  
451 assumption of differing OM source input to the mudflat sediments, the end-member mixing model of  
452 terrestrial and marine OM showed that the contribution of OM sources varied between sites and  
453 seasons (Fig 3, Table 1).

454 The  $\text{OM}_{\text{mar}}$  predominated in the studied sites likely due to the positioning of the sampling  
455 sites, which are situated near the mouth of the MRE (Atlantic Ocean). During the wet season (pore  
456 water salinity: 4-20), the  $\text{OM}_{\text{mar}}$  contribution was highest in the mudflat sediment from MRE margin  
457 (P1: 76.4 %) compared to the mangrove tidal creeks (P2: 54.5, P3: 60 %). During the dry season (pore  
458 water salinity: 18-25), due to less dilution from Marapanim River discharge, the influence of salt  
459 water is greater, thereby increasing the contribution of  $\text{OM}_{\text{mar}}$  by up to 18 % (P1: 79.6, P2: 69 and  
460 P3: 78 %). Overall, the downcore profiles of  $\text{OM}_{\text{mar}}$  and  $\text{OM}_{\text{terr}}$  remained relatively constant  
461 throughout the sediment columns during the wet season, but during the dry season an increase in  
462 marine input in the upper layers (< 18 cm) is noted at sites P2 and P3 (Fig. 3).

463

#### 464 4.3. Carbon and nutrients accumulation rates and stocks

465

466 The TOC, TN and TP accumulation rates and stocks showed differences between the sites  
467 (Table 2, Fig. 5). The highest stocks and accumulation rates were found in the mudflat sediment from  
468 mangrove tidal creek (P2 and P3) compared to the MRE margin (P1). Two factors may cause these  
469 differences. Firstly, although site P1 presented the highest sedimentation rate (Fig. 2) and highest  
470 density (Fig. 3), it is composed of >50% sand content with lower TOC, TN and TP contents compared  
471 to the sites P2 and P3. Previous studies indicated that OM preservation is often enhanced by the large  
472 surface area of fine-grained sediments, and the low energy associated with slack water deposits

473 (Mayer, 1994; Keil et al. 1994). Sites P2 and P3 are protected from the direct impact of tidal energy  
474 and waves, as compared to site P1 that is located in the MRE margin and close to a sand bar.

475 The other factor is that the efficiency of OM storage in the mudflat sediments is also dependent  
476 on the origin of OM (Saintilan et al., 2013; Watanabe and Kuwae, 2015; Kusumaningtyas et al.,  
477 2019). OM proportions in our simple end-member mixing model indicated that 49-95% of OM  
478 contribution is from marine OM. The slight increase in fluxes of terrestrial OM along the mangrove  
479 tidal creek sites (P2 and P3, Table 1) contributed to higher TOC, TN and TP accumulation rates and  
480 stocks than those seen along the MRE margin (P1). These results were corroborated with previous  
481 studies, which indicated the terrestrial OM is preserved more efficiently than phytoplankton-derived  
482 OM in estuarine sediments (Hedges et al., 1997).

483 The average TOC accumulation rate (TOC AR) calculated in the studied mudflats was slightly  
484 higher ( $192.5 \pm 43.5 \text{ g C m}^{-2} \text{ yr}^{-1}$ ) than the global average in conserved mangrove ( $170 \text{ g C m}^{-2} \text{ yr}^{-1}$ ,  
485 Perez et al., 2018) and the current global average for mangrove ecosystems ( $179.6 \text{ g C m}^{-2} \text{ yr}^{-1}$ ,  
486 Alongi, 2020). Higher rates of  $555 \text{ g C m}^{-2} \text{ yr}^{-1}$  were measured in other tropical mudflat in Piraquê-  
487 Açú estuary, Brazil, supported by high SAR ( $1.8 \text{ cm yr}^{-1}$ , Bernadino et al., 2020). In contrast, lower  
488 carbon accumulation rates were reported in subtropical microtidal creeks systems. For example,  
489 Marchio et al. (2016) evidenced TOC AR of  $162 \text{ g C m}^{-2} \text{ yr}^{-1}$  in Southwest Florida, and Santos et al.  
490 (2019) observed TOC AR of  $63 \text{ g C m}^{-2} \text{ yr}^{-1}$  in a tidal creek in Evans Head, Australia. However,  
491 according Perez et al. (2018), the distribution of TOC AR within the forests and adjacent sites (margin  
492 and mudflat environments) of mangrove ecosystems exhibited non-significant differences among  
493 regions, as variations are influenced by a combination of many local factors (e.g. geomorphology,  
494 vegetation cover, flooding frequency, hydrological regime and anthropogenic influence).

495 The average TN and TP accumulation rates were also high (TN:  $15.3 \text{ g m}^{-2} \text{ yr}^{-1}$  and TP:  $3.2 \text{ g}$   
496  $\text{m}^{-2} \text{ yr}^{-1}$ ; Table 4) when compared to the global average of anthropogenically non-impacted  
497 mangroves (TN:  $8.9 \text{ g m}^{-2} \text{ yr}^{-1}$  and TP:  $0.5 \text{ g m}^{-2} \text{ yr}^{-1}$ , Breithaupt et al., 2014). Similar to our study,

498 Bernadino et al. (2020) measured high TN accumulation rates of  $27.9 \text{ g m}^{-2} \text{ yr}^{-1}$  in a conserved mudflat  
499 in Piraquê-Açu estuary. OM enriched in nutrients may be expected to decompose faster and to a larger  
500 extent than nutrient-depleted OM (Kristensen and Hansen, 1995; McGlathery et al., 2007). Thereby,  
501 the capacity to sequester OC may decrease with the increase of TN and TP burial rates (Breithaupt et  
502 al., 2014). However, in our study area TOC AR were higher in the sites with higher TN and TP AR.  
503 Similarly, TOC stocks were higher in the sites with higher TN and TP stocks (Table 4).

504 Due to the relatively shallow sediment depth (30 cm) used in our study, it is difficult to directly  
505 compare our results with those from other studies which usually assess sediment stocks based on  
506 deeper cores (e.g.,  $\geq 100$  cm, Howard et al., 2014). However, we found some studies similar to ours  
507 with more superficial sampling. Our TOC stocks ranged from 2333 to 5218  $\text{g C m}^{-2}$  (average of 3811  
508  $\text{g C m}^{-2}$ ), which is similar those mudflat of West Papua, Indonesia (in the top 50 cm, 6200  $\text{g C m}^{-2}$ ;  
509 Sasmito et al., 2020), Araçá Bay, Brazil (in the top 20 cm, range from 1700 to 2200  $\text{g C m}^{-2}$ ; Gorman  
510 et al., 2020), and for unvegetated mudflats of China (in the top 50 cm, 4808  $\text{g C m}^{-2}$ ; Feng et al.,  
511 2019). Comparing our stocks results with the values found in other Amazon mangroves sediments (in  
512 the top 30 cm) (supplementary information in Kauffman et al., 2018), our values are within the range  
513 of 3430 – 6230  $\text{g C m}^{-2}$ , except at site P1 that was relatively lower ( $2337 \pm 279 \text{ g C m}^{-2}$ ), which can  
514 be attributed to coarse textured sediments. These results suggest that mudflat along tidal creeks also  
515 have a high potential for organic carbon storage compared to vegetated habitats such as mangrove  
516 forests.

517 Few studies have assessed nitrogen and phosphorus stocks in mangrove and mudflat  
518 sediments. Feng et al. (2017) measured TN and TP stocks in an unvegetated mudflat of China (in the  
519 top 40 cm). Their average of TN stock of  $\sim 290 \text{ g N m}^{-2}$  is comparable to  $296 \pm 109 \text{ g N m}^{-2}$  (Table 4)  
520 in our study, but TP stock of  $\sim 150 \text{ g P m}^{-2}$  is higher than what we found of  $58 \pm 15 \text{ g P m}^{-2}$  (Table 4).  
521 Saderne et al. (2020) measured TN and TP stocks in mangrove sediments (in the top 20 cm) in the  
522 central Red Sea ( $99 \text{ g N m}^{-2}$  and  $70 \text{ g P m}^{-2}$ ) and along the Gulf coast of Saudi Arabia ( $223 \text{ g N m}^{-2}$

523 and 32.8 g P m<sup>-2</sup>), where TN and TP stocks in the Gulf were quite comparable to what is presented  
524 here, but TN stocks were at least twofold higher in the Gulf compared to the Red Sea, while TP stocks  
525 were a maximum of 1.2 times higher in Red Sea. Perez et al. (2017) observed in a conserved  
526 mangrove forest in New Zealand estuary TN stock between 400 and 500 g N m<sup>-2</sup>, in the top 40 cm,  
527 which is slightly higher than our values. In contrast, Ray et al. (2014) and Ray et al. (2017) measured  
528 the TN and TP stocks, respectively, in the top 60 cm sediment in the Indian Sundarban mangrove and  
529 found an average of 4.2 g N m<sup>-2</sup> and 0.4 g P m<sup>-2</sup>, which are lower than we found (58 ± 15 g P m<sup>-2</sup>,  
530 Table 4), like as a result of the TN and TP conserved in the living biomass.

531 The capacity to sequester and store OM in the studied creek mudflats likely reflects the SAR  
532 (1.5 ± 0.3 cm yr<sup>-1</sup>), which is higher than the global average in conserved mangrove (0.36 ± cm yr<sup>-1</sup>;  
533 Perez et al., 2018) and the current global average (0.77 cm yr<sup>-1</sup>; Breithaupt et al., 2012). This high  
534 SAR is supported by strong interaction between the river stream and the tides that characterize this  
535 environment, being close to the highest global accumulation rates in the mangrove forest (Breithaupt  
536 et al. 2012; Kusumaningtyas et al., 2019) and mudflat (Bernadino et al., 2020). In general, the  
537 sedimentation rate is considered as a driving factor controlling the OM burial efficiency (Canfield,  
538 1994). Although the sedimentary OM with a high proportion of marine origin tends to be more  
539 susceptible to decomposition, sites with high sedimentation rates contribute significantly to preserve  
540 OM (Hedges and Keil, 1995; Canfield et al., 2005). In addition, it should be noted that the studied  
541 mangrove area is densely inhabited by *Rhizophora*, *Avicennia* and *Laguncularia* trees, with heights  
542 up to 33 m. This probably reflects a high productivity that provided a large supply of mangrove litter  
543 and dead roots to the mangrove creeks through tidal exchange.

544 Recently, Call et al. (2019) demonstrated that a mangrove tidal creek in Amazon presented  
545 large CO<sub>2</sub> and CH<sub>4</sub> fluxes from a nearby mangrove tidal creek. However, our results indicate that the  
546 Amazon mangrove creeks are also sites that accumulate considerable amounts of organic carbon (Fig.  
547 7). For example, the total mudflat area in the tidal creek catchment of this study was near 0.5 km<sup>2</sup>

548 (Fig. 1c), and considering an average of the TOC accumulation rates between the three study sites,  
549 we estimate that a total of 96 kg TOC yr<sup>-1</sup> is sequestered in these creeks catchment. This suggests a  
550 previously unaccounted carbon sink in creek environments that are relevant to the carbon budgets  
551 along mangrove-colonized coastal zones. Therefore, mangrove tidal creeks can play an important role  
552 in the carbon cycling in tropical land-sea interfaces more than just acting as a conduit for exchanges  
553 between mangrove forests and coastal waters.

554         Considering global average in conserved mangrove of 170 g C m<sup>-2</sup> yr<sup>-1</sup> (Perez et al., 2018)  
555 and the mangrove forest area within the catchment of this study was 7,638,368 m<sup>2</sup>, we estimate that  
556 a total of 1299 t TOC yr<sup>-1</sup> is sequestered in these mangrove soils. Thus, the creek mudflat area  
557 accumulates around 0.007% of what would be estimated for the mangrove forest soils. Carbon  
558 dioxide emissions at the water–air interface were 2794 ± 2072 g m<sup>-2</sup> yr<sup>-1</sup> (174 ± 129 mmol m<sup>-2</sup> d<sup>-1</sup>)  
559 from a nearby mangrove tidal creek (Call et al., 2019). Using our creek mudflat area (0.5 km<sup>2</sup>), CO<sub>2</sub>  
560 emissions would be 1397 kg C-CO<sub>2</sub> yr<sup>-1</sup>, while total mudflat sediment carbon burial would be 96.3  
561 kg yr<sup>-1</sup>. Therefore, sediment carbon sequestration in this system may offset < 7.0 % of the aquatic  
562 CO<sub>2</sub> emissions.

563

## 564 **5. Conclusion**

565

566         This study investigated the recent (24 ± 4 years) carbon and nutrient accumulation rates and  
567 stocks in intertidal mudflats situated along mangrove creeks within the Brazilian Amazon coast. Even  
568 though marine OM was predominant along our study sites (49% to 95%), fine-grained sediments and  
569 higher terrestrial OM input, a product of a dense and tall mangroves trees, contributed to higher rates  
570 of TOC, TN, and TP accumulation and stocks in the mangrove tidal creeks (P2 and P3) compared to  
571 the Marapanim River estuary margin (P1). The capacity to sequester and store TOC also increased  
572 with increases in TN and TP burial rates and stocks. Due to the high sedimentation rates (1.5 ± 0.3



573 cm yr<sup>-1</sup>), the TOC (192.5 ± 43.5), TN (15.3 ± 4.1), and TP (3.2 ± 0.8) accumulation rates found here  
574 were slightly higher than the global averages estimated for mangrove forest sediments. The potential  
575 of intertidal mudflats to sequester OC suggests that the creek environments are unaccounted carbon  
576 sinks and are relevant in terms of carbon budgets in mangrove-colonized coastal zones.

577

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579

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583

## 584 **References**

585

586 AgisoftPhotoScan, 2018. AgiSoft PhotoScan Professional. Version 1.4.5. Agisoft LLC. St.  
587 Petersburg, Russia. Retrieved from <http://www.agisoft.com/downloads/installer/>.

588

589 Alongi, D.M., Boto, K.G., Robertson, A.I., 1992. Nitrogen and phosphorus cycles, in: Robertson,  
590 A.I., Alongi, D.M. (Eds.), *Tropical Mangrove Ecosystems. Coastal and Estuarine Studies*. American  
591 Geophysical Union, Washington, DC, pp. 251–292. <https://doi.org/10.1029/ce041p0251>.

592

593 Alongi, D.M., 2014. Carbon cycling and storage in mangrove forests. *Annu. Rev. Mar. Sci.* 6, 195–  
594 219. <https://doi.org/10.1146/annurev-marine-010213-135020>.

595

596 Alongi, D.M., 2020. Global Significance of Mangrove Blue Carbon in Climate Change Mitigation  
597 (Version 1). *Sci*, 2, 57.

598

599 Appleby, P.G., Oldfield, F., 1992. Application of lead-210 to sedimentation studies, in:  
600 Ivanovich, M., Harmon, S. (Eds.), *Uranium Series Disequilibrium: Application to*  
601 *Earth, Marine and Environmental Science*, second ed. Oxford Science Publications, pp. 731-783.

602

603 Arias-Ortiz, A., Masqué, P., Garcia-Orellana, J., Serrano, O., Mazarrasa, I., Marbà, N., Lovelock,  
604 C.E., Lavery, P.S., Duarte, C.M., 2018. Reviews and syntheses: <sup>210</sup>Pb-derived sediment and carbon  
605 accumulation rates in vegetated coastal ecosystems – setting the record straight. *Biogeosciences* 15,  
606 6791–6818. <https://doi.org/10.5194/bg-15-6791-2018>.

607

608 Aspila, K.L., Agemian, H., Chau, A.S.Y., 1976. A semi-automated method for the determination of  
609 inorganic, organic and total phosphate in sediments. *Analyst* 101, 187–197. <https://doi.org/10.1039/AN9760100187>.

610

611

612 Bernardino, A.F., Sanders, C.J., Bissoli, L.B., de O. Gomes, L.E., Kauffman, J.B., Ferreira, T.O.,  
613 2020. Land use impacts on benthic bioturbation potential and carbon burial in Brazilian mangrove  
614 ecosystems. *Limnol. Oceanogr.* 9999, 1-11. <http://dx.doi.org/10.1002/lno.11458>.

615

616 Berrêdo, J.F., Costa, M.L., Progene, M.P.S., 2008. Efeitos das variações sazonais do clima tropical  
617 úmido sobre as águas e sedimentos de manguezais do estuário do rio Marapanim, costa nordeste do  
618 Estado do Pará. *Acta Amazônica* 38, 473-482. [http://dx.doi.org/10.1590/S0044-](http://dx.doi.org/10.1590/S0044-59672008000300012)  
619 [59672008000300012](http://dx.doi.org/10.1590/S0044-59672008000300012).

620

621 Blott, S.J., Pye, K., 2001. GRADISTAT: a grain size distribution and statistics package for the  
622 analysis of unconsolidated sediments. *Earth Surface Processes and Landforms* 26, 1237-1248.  
623 <https://doi.org/10.1002/esp.261>.

624

625 Bouillon, S., Connolly, R.M., Lee, S.Y., 2008. Organic matter exchange and cycling in mangrove  
626 ecosystems: Recent insights from stable isotope studies. *Journal of Sea Research*, 59, 44-58,  
627 <https://doi.org/10.1016/j.seares.2007.05.001>.

628

629 Breithaupt, J.L., Smoak, J.M., Smith, T.J., Sanders, C.J., Hoare, A., 2012. Organic carbon burial rates  
630 in mangrove sediments: strengthening the global budget. *Global Biogeochem. Cycles* 26.  
631 <https://doi.org/10.1029/2012gb004375>.

632

633 Breithaupt, J.L., Smoak, J.M., Smith, T.J., Sanders, C.J., 2014. Temporal variability of carbon and  
634 nutrient burial, sediment accretion, and mass accumulation over the past century in a carbonate

635 platform mangrove forest of the Florida Everglades. *J. Geophys. Res. Biogeosci.* 119, 2032–2048.  
636 <https://doi.org/10.1002/2014JG002715>.

637

638 Call, M., Santos, I.R., Dittmar, T., Rezende, C.E., Asp, N.E., Maher, D.T., 2019. High pore-water  
639 derived CO<sub>2</sub> and CH<sub>4</sub> emissions from a macro-tidal mangrove creek in the Amazon region.  
640 *Geochimica et Cosmochimica Acta* 247, 106–120. <https://doi.org/10.1016/j.gca.2018.12.029>.

641

642 Canfield, D.E., 1994. Factors influencing organic carbon preservation in marine  
643 sediments. *Chem. Geol.* 114, 315-329. [https://doi.org/10.1016/0009-2541\(94\)90061-2](https://doi.org/10.1016/0009-2541(94)90061-2).

644

645 Canfield, D.E., Kristensen, E., Thamdrup, B., 2005. Heterotrophic Carbon Metabolism. *Aquatic  
646 geomicrobiology: Advances Marine Biology* 48, 129-166. [https://doi.org/10.1016/S0065-  
647 2881\(05\)48005-0](https://doi.org/10.1016/S0065-2881(05)48005-0).

648

649 Chambers, L.G., Davis, S.E., Troxler, T., Boyer, J.N., Downey-Wall, A., Scinto, L.J., 2013.  
650 Biogeochemical effects of simulated sea level rise on carbon loss in an Everglades mangrove peat  
651 soil. *Hydrobiologia* 726, 195–221. <https://doi.org/10.1007/s10750-013-1764-6>.

652

653 Cohen, M.C.L., de Souza, A.V., Rossetti, D.F., Pessenda, L.C.R., França, M.C., 2018. Decadal-scale  
654 dynamics of an Amazonian mangrove caused by climate and sea level changes: inferences from  
655 spatial-temporal analysis and Digital Elevation Models. *Earth Surface Processes and Landforms* 43,  
656 2876–2888. <https://doi.org/10.1002/esp.4440>.

657

658 Cohen, M.C.L., Figueiredo, B.L., Oliveira, N.N., Fontes, N.A., França, M.C., Pessenda, L.C.R.,  
659 Lima, P., 2019. Impacts of Holocene and modern sea- level changes on estuarine mangroves from  
660 Northeastern Brazil. *Earth Surface Processes and Landforms*, esp.4737.  
661 <https://doi.org/10.1002/esp.4737>.

662

663 Darrow, E.S., Carmichael, R.H., Calci, K.R., Burkhardt, W., 2017. Land-use related changes to  
664 sedimentary organic matter in tidal creeks of the northern Gulf of Mexico. *Limnol. Oceanogr.* 62,  
665 686–705. <https://doi.org/10.1002/lno.10453>.

666

667 Deines, P., 1980. The isotopic composition of reduced organic carbon, in: Fritz, P., Fontes, J.C.  
668 (Eds.), Handbook of Environmental Isotope Geochemistry. The Terrestrial Environment, vol. 1. A.  
669 Elsevier, Amsterdam, pp. 329–406. <https://doi.org/10.1016/B978-0-444-41780-0.50015-8>.  
670

671 Dittmar, T., Hertkorn, N., Kattner, G., Lara, R.J., 2006. Mangroves, a major source of dissolved  
672 matter sources to the oceans. *Global Biogeochemical Cycles* 20, GB1012.  
673 <https://doi.org/10.1029/2005GB002570>.  
674

675 Feng, J., Zhou, J., Wang, L., Cui, X., Ning, C., Wu, H., Zhu, X., Lin, G., 2017. Effects of short-term  
676 invasion of *Spartina alterniflora* and the subsequent restoration of native mangroves on the soil  
677 organic carbon, nitrogen and phosphorus stock *Chemosphere* 184, 774–83.  
678 <https://doi.org/10.1016/j.chemosphere.2017.06.060>.  
679

680 Feng, J., Wang, S., Wang, S., Ying, R., Yin, F., Jiang, L., Li, Z., 2019. Effects of Invasive *Spartina*  
681 *alterniflora* Loisel. and Subsequent Ecological Replacement by *Sonneratia apetala* Buch.-Ham. on  
682 Soil Organic Carbon Fractions and Stock. *Forests* 10, 171. <https://doi.org/10.3390/f10020171>.  
683

684 Fernandes, M.E.B., Oliveira, F.P., Eyzaguirre, I.A.L., 2018. Mangroves on the Brazilian Amazon  
685 Coast: Uses and Rehabilitation, in: Makowski C., Finkl C. (eds) Threats to Mangrove Forests. Coastal  
686 Research Library, vol 25. Springer, Cham. [https://doi.org/10.1007/978-3-319-73016-5\\_29](https://doi.org/10.1007/978-3-319-73016-5_29).  
687

688 Giri, C., Ochieng, E., Tieszen, L.L., Zhu, Z., Singh, A., Loveland, T., Masek, J., Duke, N., 2011.  
689 Status and distribution of mangrove forests of the world using earth observation satellite data. *Global*  
690 *Ecol. Biogeogr.* 20, 154–159. <https://doi.org/10.1111/j.1466-8238.2010.00584.x>.  
691

692 GlobalMapper, 2017. (Version 19). Blue Marble Geographics. Hallowell, Maine, United State.  
693

694 Goñi, M., Teixeira, M., Perkey, D., 2003. Sources and distribution of organic matter in  
695 a river-dominated estuary (Winyah Bay, SC, USA). *Estuar. Coast Shelf Sci.* 57, 1023–1048.  
696 [https://doi.org/10.1016/S0272-7714\(03\)00008-8](https://doi.org/10.1016/S0272-7714(03)00008-8).  
697

698 Gontharet, S., Mathieu, O., Lévêque, J., Milloux, M.-J., Lesourd, S., Philippe, S., Caillaud, J., Gardel,  
699 A., Sarrazin, M., Proisy, C., 2014. Distribution and sources of bulk organic matter (OM) on a tropical

700 intertidal mud bank in French Guiana from elemental and isotopic proxies. *Chemical Geology*, 376,  
701 1-10, <https://doi.org/10.1016/j.chemgeo.2014.03.009>.

702

703 Gorman, D., Sumida, P.Y.G., Figueira, R.C.L., Turra, A., 2020. Improving soil carbon estimates of  
704 mudflats in Araçá Bay using spatial models that consider riverine input, wave exposure and  
705 biogeochemistry. *Estuarine, Coastal and Shelf Science*, 238, 106734,  
706 <https://doi.org/10.1016/j.ecss.2020.106734>.

707

708 Grasshoff, K., Kremling, K., Ehrhardt, M., 1999. *Methods of Seawater Analysis*. third ed. Wiley,  
709 New York. <https://doi.org/10.1002/9783527613984>.

710

711 Hammer, O., Harper, D.A.T., Ryan, P.D., 2001. PAST: Paleontological statistics software package  
712 for education and data analysis. *Palaeontologia Electronica* 4(1): 9 pp. [https://palaeo-](https://palaeo-electronica.org/2001_1/past/issue1_01.htm)  
713 [electronica.org/2001\\_1/past/issue1\\_01.htm](https://palaeo-electronica.org/2001_1/past/issue1_01.htm).

714

715 Hedges, J.I., Keil, R.G., 1995. Sedimentary organic matter preservation: an assessment and  
716 speculative synthesis. *Marine Chemistry* 49, 81–115. [https://doi.org/10.1016/0304-4203\(95\)00008-](https://doi.org/10.1016/0304-4203(95)00008-F)  
717 [F](https://doi.org/10.1016/0304-4203(95)00008-F).

718

719 Hedges, J.I., Keil, R.G., Benner, R., 1997. What happens to terrestrial organic matter in the ocean?  
720 *Org. Geochem.* 27, 195–212. [https://doi.org/10.1016/S0146-6380\(97\)00066-1](https://doi.org/10.1016/S0146-6380(97)00066-1).

721

722 Howard, J., Hoyt, S., Isensee, K., Pidgeon, E., and Telszewski, M., 2014. Coastal Blue Carbon:  
723 Methods for Assessing Carbon Stocks and Emissions Factors in Mangroves, Tidal Salt Marshes, and  
724 Seagrass Meadows, *Conserv. Int. Intergov. Oceanogr. Comm. UNESCO, Int. Union Conserv. Nature*,  
725 Arlington, Virginia, USA, 1–180.

726

727 ICMBio. Atlas dos Manguezais do Brasil. 2018. 1st ed. Brasília: Instituto Chico Mendes de  
728 Conservação da Biodiversidade; 176 p. [https://doi.org/10.1016/S0146-6380\(97\)00066-1](https://doi.org/10.1016/S0146-6380(97)00066-1).

729

730 Jennerjahn, T.C., Ittekkot, V., 2002. Relevance of mangroves for the production and deposition of  
731 organic matter along tropical continental margins, *Naturwissenschaften* 89(1), 23–30.  
732 <https://doi.org/10.1007/s00114-001-0283-x>.

733

734 Jennerjahn, T.C., 2012. Biogeochemical response of tropical coastal systems to present and past  
735 environmental change. *Earth Sci. Rev.* 114, 19–41. <https://doi.org/10.1016/j.earscirev.2012.04.005>.

736

737 Kauffman, J.B., Bernardino, A.F., Ferreira, T.O., Giovannoni L.R., de O. Gomes L.E.,  
738 Romero D.J., Jimenez L.C.Z., Ruiz F., 2018. Carbon stocks of mangroves and salt marshes of the  
739 Amazon region, Brazil. *Biol. Lett.* 14: 20180208. <http://dx.doi.org/10.1098/rsbl.2018.0208>.

740

741 Keil, R.G., Montluçon, D.B., Prahl, F.G., Hedges, J.I., 1994. Sorptive preservation of labile organic  
742 matter in marine sediments. *Nature* 370, 549–552. <https://doi.org/>

743

744 Kjerfve, B., Lacerda, L.D., 1993. Mangroves of Brazil. In: *Conservation and Sustainable Utilization*  
745 *of Mangrove Forests in Latin America and Africa Regions. Part 1: Latin America. ITTO/ISME*  
746 *Project PD114/90(F)*, pp. 245-272.

747

748 Kristensen, E., Hansen, K., 1995. Decay of plant detritus in organic-poor marine sediment:  
749 production rates and stoichiometry of dissolved C and N compounds. *J. Mar. Res.* 53, 675–702.  
750 [https://doi: 10.1357/0022240953213115](https://doi:10.1357/0022240953213115).

751

752 Kristensen, E., Bouillon, S., Dittmar, T., Marchand, C., 2008. Organic carbon dynamics in mangrove  
753 ecosystems: a review. *Aquatic botany* 89(2), 201-219.  
754 <https://doi.org/10.1016/j.aquabot.2007.12.005>.

755

756 Kusumaningtyasa, M.A., Hutahaean, A.A., Fischer, H.W., Mayod, M.P., Ransby, D., Jennerjahn  
757 T.C., 2019. Variability in the organic carbon stocks, sources, and accumulation rates of Indonesian  
758 mangrove ecosystems. *Estuarine, Coastal and Shelf Science* 218, 310–323.  
759 <https://doi.org/10.1016/j.ecss.2018.12.007>.

760

761 Lamb, A.L., Wilson, G.P., Leng, M.J., 2006. A review of coastal palaeoclimate and relative sea-level  
762 reconstructions using  $\delta^{13}\text{C}$  and C/N ratios in organic material. *Earth-Sci. Reviews* 75, 29–57.  
763 <https://doi.org/10.1016/j.earscirev.2005.10.003>.

764

765 Liu, D., Li, X., Emeis, K.C., Wang, Y., Richard, P., 2015. Distribution and sources of organic matter  
766 in surface sediments of Bohai Sea near the Yellow River Estuary, China. *Estuarine, Coastal and Shelf*  
767 *Science* 165, 128-136. <https://doi.org/10.1016/j.ecss.2015.09.007>.  
768

769 Loring, D.H., Rantala, R.T.T., 1992. Manual for the geochemical analyses of marine sediments and  
770 suspended particulate matter. *Earth-Sci. Rev.* 32, 235–283. [https://doi.org/10.1016/0012-](https://doi.org/10.1016/0012-8252(92)90001-A)  
771 [8252\(92\)90001-A](https://doi.org/10.1016/0012-8252(92)90001-A).  
772

773 Marchio, D., Savarese, M., Bovard, B., Mitsch, W., 2016. Carbon sequestration and sedimentation in  
774 mangrove swamps influenced by hydrogeomorphic conditions and urbanization in Southwest  
775 Florida. *Forests* 7(12), 116. <https://doi.org/10.3390/f7060116>.  
776

777 Martorano, L.G., Pereira, L.C., Cesar, E.G.M., Pereira, I.C.B., 1993. Estudos climáticos do Estado  
778 do Pará, classificação climática (Köppen) e deficiência hídrica (Thornthwhite, Mather). Belém,  
779 SUDAM/EMBRAPA, SNLCS, pp. 53.  
780

781 Mayer, L.M., 1994. Surface area control of organic carbon accumulation in continental shelf  
782 sediments. *Geochim. Cosmochim. Acta* 58, 1271–1284. [https://doi.org/10.1016/0016-](https://doi.org/10.1016/0016-7037(94)90381-6)  
783 [7037\(94\)90381-6](https://doi.org/10.1016/0016-7037(94)90381-6).  
784

785 McGlathery, K. J., Sundback, K., and Anderson, I. C., 2007. Eutrophication in shallow coastal bays  
786 and lagoons: the role of plants in the coastal filter. *Mar. Ecol. Prog. Ser.* 348, 1–18.  
787 <https://doi/10.3354/meps07132>.  
788

789 Mcleod, E., Chmura, G.L., Bouillon, S., Salm, R., Björk, M., Duarte, C.M., Lovelock, C.E.,  
790 Schlesinger, W.H. and Silliman, B.R., 2011. A blueprint for blue carbon: toward an improved  
791 understanding of the role of vegetated coastal habitats in sequestering CO<sub>2</sub>. *Frontiers in Ecology and*  
792 *the Environment*, 9: 552-560. <https://doi.org/10.1890/110004>.  
793

794 Meyers, P.A., 1994. Preservation of elemental and isotopic source identification of sedimentary  
795 organic matter. *Chem. Geol.* 114, 289–302. [https://doi.org/10.1016/0009-2541\(94\)90059-0](https://doi.org/10.1016/0009-2541(94)90059-0).  
796

797 Meyers, P.A., 1997. Organic geochemical proxies of paleoceanographic, paleolimnologic, and  
798 paleoclimatic processes. *Organic Geochemistry* 27, 213 – 250. <https://doi.org/10.1016/S0146->  
799 6380(97)00049-1.  
800



801 Meyers, P.A., 2003. Applications of organic geochemistry to paleolimnological reconstructions: a  
802 summary of examples from the Laurentian Great Lakes. *Org. Geochem.* 34, 261–289.  
803 [https://doi.org/10.1016/S0146-6380\(02\)00168-7](https://doi.org/10.1016/S0146-6380(02)00168-7).

804

805 Nascimento, W.R., Souza-Filho, P.W.M., Proisy, C., Lucas, R.M., Rosenqvist, A., 2013 Mapping  
806 changes in the largest continuous Amazonian mangrove belt using object-based classification of  
807 multisensor satellite imagery. *Estuarine Coastal and Shelf Sci.* 117, 83–93.  
808 <https://doi.org/10.1016/j.ecss.2012.10.005>.

809

810 Nellemann, C., Corcoran, E., Duarte, C. M., Valdrés, L., Young, C. D., Fonseca, L., Grimsditch, G.,  
811 2009. *Blue Carbon - The Role of Healthy Oceans in Binding Carbon*. UN Environment, GRID-  
812 Arendal. ISBN: 978-82-7701-060-1.

813

814 Paiva, R.S., Eskinazi-Leça, E., Passavante, J.Z.O., Silva-Cunha, M.G.G., Melo, N.F.A.C., 2006.  
815 Ecological considerations on phytoplankton from the Guajará bay and from the Guamá river estuary  
816 in Para, Brazil. *Bol. Mus. Para. Emílio Goeldi. Ciências Naturais* 1, 133–146.

817

818 Pérez, A., Machado, W., Gutierrez, D., Stokes, D., Sanders, L., Smoak, J.M., Santos, I., Sanders, C.J.,  
819 2017. Changes in organic carbon accumulation driven by mangrove expansion and deforestation in a  
820 New Zealand estuary. *Est. Coast. Shelf Sci.* 192, 108–116.  
821 <https://doi.org/10.1016/j.ecss.2017.05.009>.

822

823 Pérez, A., Libardoni, B.G., Sanders, C.J., 2018. Factors influencing organic carbon accumulation in  
824 mangrove ecosystems. *Biol. Lett.* 14 (10). [https://doi.org/10.1098/rsbl.](https://doi.org/10.1098/rsbl.2018.0237)  
825 [2018.0237](https://doi.org/10.1098/rsbl.2018.0237).

826

827 Peterson, B.J., Howarth, R.W., 1987. Sulfur, carbon, and nitrogen isotopes used to trace organic  
828 matter flow in the salt-marsh estuaries of Sapelo Island, Georgia. *Limnology and*  
829 *oceanography* 32(6), 1195-1213. <https://doi.org/10.4319/lo.1987.32.6.1195>.

830

831 Prahl, F.G., De Lange, G.T., Scholten, S., Cowie, G.L., 1997. A case of post-depositional aerobic  
832 degradation of terrestrial organic matter in turbidite deposits from the Madeira Abyssal Plain. *Org.*  
833 *Geochem.* 27, 141–152. [https://doi.org/10.1016/S0146-6380\(97\)00078-8](https://doi.org/10.1016/S0146-6380(97)00078-8).

834

835 Prasad, M.B.K., Kumar, A., Ramanathan, A.L., Datta, D.K., 2017. Sources and dynamics of  
836 sedimentary organic matter in Sundarban mangrove estuary from Indo-Gangetic delta. *Ecological*  
837 *Processes* 6, 8. <https://doi.org/10.1186/s13717-017-0076-6>.

838

839 Prasad, M.B.K., Ramanathan, A.L., 2009. Organic matter characterization in a tropical estuarine-  
840 mangrove ecosystem of India: preliminary assessment by using stable isotopes and lignin phenols.  
841 *Estuarine Coastal Shelf Science* 84(4): 617–624. <https://doi.org/10.1016/j.ecss.2009.07.029>.

842

843 Ranjan, R.K., Routh, J., Ramanathan, A.L., Klump, J.V., 2011. Elemental and stable isotope records  
844 of organic matter input and its fate in the Pichavaram mangrove–estuarine sediments (Tamil Nadu,  
845 India). *Marine Chemistry* 126, 163–172. <https://doi.org/10.1016/j.marchem.2011.05.005>.

846

847 Ray, R., Majumder N., Das, S., Chowdhury, C., Jana, T.K., 2014. Biogeochemical cycle of nitrogen  
848 in a tropical mangrove ecosystem, east coast of India. *Marine Chemistry*, 167, 33–43.  
849 <https://doi.org/10.1016/j.marchem.2014.04.007>.

850

851 Ray, R., Majumder, N., Chowdhury, C., Das, S., Jana, T.K., 2017. Phosphorus Budget of the  
852 Sundarban Mangrove Ecosystem: Box Model Approach. *Estuaries and Coasts* 41, 1036–1049.  
853 <https://doi.org/10.1007/s12237-017-0332-0>.

854

855 Routh, J., Meyers, P.A., Choudhury, P., Kumar, B., 2009., A sediment record of recent  
856 nutrient loading and trophic state change in Lake Norrviken, Sweden. *J. Paleolimnol.* 42, 325–341.  
857 <https://doi.org/10.1007/s10933-008-9279-2>.

858

859 Redfield, A.C., Ketchum, B.H. and Richards, F.A., 1963. The influence of organisms on the  
860 composition of sea-water, in: Hill, M.N. (Ed.) *The composition of seawater: Comparative and*  
861 *descriptive oceanography. The sea: ideas and observations on progress in the study of the seas*, 2: pp.  
862 26-77.

863

864 Rosentreter, J.A., Maher, D.T., Erler, D.V., Murray, R., Eyre, B.D., 2018. Seasonal and temporal CO<sub>2</sub>  
865 dynamics in three tropical mangrove creeks – a revision of global mangrove CO<sub>2</sub>  
866 emissions. *Geochim. Cosmochim. Acta* 222, 729–745. <https://doi.org/10.1016/j.gca.2017.11.026>.

867

868 Saintilan, N., Rogers, K., Mazumder, D., Woodroffe, C., 2013. Allochthonous and autochthonous  
869 contributions to carbon accumulation and carbon store in southeastern Australian coastal wetlands.  
870 *Estuar. Coast Shelf Sci.* 128, 84–92. <https://doi.org/10.1016/j.ecss.2013.05.010>.

871

872 Sanders, C.J., Santos, I.R., Maher, D.T., Breithaupt, J.L., Smoak, J.M., Ketterer, M., Call,  
873 M., Sanders, L.M., Eyre, B.D., 2016. Examining  $^{239+240}\text{Pu}$ ,  $^{210}\text{Pb}$  and historical  
874 events to determine carbon, nitrogen and phosphorus accumulation in mangrove  
875 sediments of Moreton Bay, Australia. *J. Environ. Radioact.* 151, 623–629.  
876 <https://doi.org/10.1016/j.jenvrad.2015.04.018>.

877

878 Santos, I.R., Maher, D.T., Larkin R., Webb, J.R., Sanders, C.J., 2019. Carbon outwelling and  
879 outgassing vs. burial in an estuarine tidal creek surrounded by mangrove and saltmarsh wetlands.  
880 *Limnol. Oceanogr.* 64, 996–1013. <https://doi.org/10.1002/lno.11090>.

881

882 Sasmito, S.D., Kuzyakov, Y., Lubis, A.A., Murdiyarso, D., Hutley, L.B., Bachri, S., Friess, D.A.,  
883 Martius, C., Borchard, N. 2020. Organic Carbon Burial and Sources in Soils of Coastal Mudflat and  
884 Mangrove Ecosystems. *Catena*, 187, 104414. <https://doi.org/10.1016/j.catena.2019.104414>.

885

886 Schultz, D.J., Calder, J.A., 1976. Organic carbon  $^{13}\text{C}/^{12}\text{C}$  variations in estuarine sediments.  
887 *Geochim. Cosmochim. Acta* 40, 381–385. [https://doi.org/10.1016/0016-7037\(76\)90002-8](https://doi.org/10.1016/0016-7037(76)90002-8).

888

889 Silva, C.A., Souza-Filho, P.W.M., Rodrigues, S.W.P., 2009. Morphology and modern sedimentary  
890 deposits of the macrotidal Marapanim Estuary (Amazon, Brazil). *Continental Shelf Research* 29,  
891 619–631. <https://doi.org/10.1016/j.csr.2008.09.018>.

892

893 Smoak, J.M., Breithaupt, J.L., Smith, T.J., Sanders, C.J., 2013. Sediment accretion and  
894 organic carbon burial relative to sea-level rise and storm events in two  
895 mangrove forests in Everglades National Park. *Catena* 104, 58–66.

896

897 Steinmuller, H.E., Foster, T.E., Boudreau, P., Ross Hinkle, C., Chambers, L.G., 2020. Tipping Points  
898 in the Mangrove March: Characterization of Biogeochemical Cycling Along the Mangrove–Salt  
899 Marsh Ecotone. *Ecosystems* 23, 417–434. <https://doi.org/10.1007/s10021-019-00411-8>.

900

901 Udden, J.A., 1914. Mechanical composition of clastic sediments. *Bulletin of the Geological Society*  
902 *of America* 25, 655–744.

903

904 Vilhena, M.P.S.P., Costa, M.L., Berrêdo, J.F., 2013. Accumulation and transfer of Hg, As, Se, and  
905 other metals in the sediment-vegetation-crab-human food chain in the coastal zone of the northern  
906 Brazilian state of Para (Amazonia). *Environ Geochem Health*, 35, 477–494.  
907 <https://doi.org/10.1007/s10653-013-9509-z>.

908

909 Vilhena, M.P.S.P., Costa, M.L., Berrêdo, J.F., Paiva, R.S., Moreira M.Z., 2018. The sources and  
910 accumulation of sedimentary organic matter in two estuaries in the Brazilian Northern coast. *Regional*  
911 *Studies in Marine Science* 18, 188–196. <https://doi.org/10.1016/j.rsma.2017.10.007>.

912

913 Watanabe, K., Kuwae, T., 2015. How organic carbon derived from multiple sources contributes to  
914 carbon sequestration processes in a shallow coastal system? *Global Change Biol.* 21, 2612–2623.  
915 <https://doi.org/10.1111/gcb.12924>.

916

917 Wentworth, C.K., 1922. A scale of grade and class terms for clastic sediments. *Journal of Geology*  
918 30, 377–392.

919

920 Zonneveld, K.A.F., Versteegh, G.J.M., Kasten, S., Eglinton, T.I., Emeis, K.-C., Huguet, C.,  
921 Koch, B.P., de Lange, G.J., de Leeuw, J.W., Middelburg, J.J., Mollenhauer, G., Prahl,  
922 F., Rethemeyer, J., Wakeham, S., 2010. Selective preservation of organic matter in  
923 marine environments; processes and impact on the fossil record. *Biogeosciences*  
924 7, 483–511. <https://doi.org/10.5194/bg-7-483-2010>.

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## 926 **Figure Captions**

927 **Fig. 1.** (A) and (B) Map of the study area in Marapanim estuarine mangrove, Brazil. (B) Sampling  
928 location (P1, P2 and P3). (C) Catchment and mudflat areas related to the sampling sites. (D) Digital  
929 vegetation height model. (E) The diagrammatic mangrove forest distribution and positioning of the  
930 sampled cores in the mud tidal flat mangrove sediments. Black dashed lines represent the height of  
931 the mangrove.

932 **Fig. 2.**  $\ln^{210}\text{Pb}_{\text{ex}}$  ( $\text{Bq kg}^{-1}$ ) activities versus depth in the sediment cores (P1, P2 and P3), sedimentation  
933 rates were calculated using selected points of  $^{210}\text{Pb}_{\text{ex}}$  profiles (filled symbols). Open symbols  
934 correspond to result interpreted as biologically and physically disturbed.

935 **Fig. 3.** Depth profiles of grain size, DBD, TOC, TN, TP,  $\delta^{15}\text{N}$ ,  $\delta^{13}\text{C}$  and TOC/TN (molar ratio) for  
936 the sediment cores (P1, P2 and P3). Including proportions of terrestrial ( $\text{OM}_{\text{terr}}$ ) and marine ( $\text{OM}_{\text{mar}}$ )  
937 sedimentary OM, during wet (filled circles) and dry (open circles) seasons.

938 **Fig. 4.** (a) Principal Component Analysis (PCA) plot showing the multivariate variation among three  
939 sites in terms of environmental variables. (b) Vectors indicate the direction and strength of each  
940 environmental variable to the overall distribution. The first two principal axes explained 74.5 % of  
941 the variance.

942 **Fig. 5.** TOC, TN and TP stocks (solid bars,  $\text{g m}^{-2}$ ) and accumulation rates (striped bars,  $\text{g m}^{-2} \text{yr}^{-1}$ ) in  
943 the sediment cores (P1, P2 and P3). The error bars are based on the standard deviation from the  
944 average between the wet and dry season dataset.

945 **Fig. 6.** Origin of sedimentary OM as indicated by  $\delta^{13}\text{C}$  against TOC/TN (molar ratio). Fields are  
946 defined from the compilation of coastal sediments by Lamb et al. (2006). The phytoplankton (white  
947 diamond symbol) and Mangrove plants (black diamond symbol) endmember data are taken from the  
948 original data in this study.

949 **Fig. 7.** Schematic comparing the global carbon (Perez et al., 2018) and nitrogen and phosphorus burial  
950 rates (Breithaupt et al. 2014) for conserved mangrove forest with Marapanim mangrove creek (this  
951 study) and the global mangrove water-atmosphere  $\text{CO}_2$  and  $\text{CH}_4$  flux rates (Rosentreter et al., 2018)  
952 with “Furo do Meio” mangrove creek (Call et al. 2019). All rates are in  $\text{g m}^{-2} \text{yr}^{-1}$ .

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Figure 1

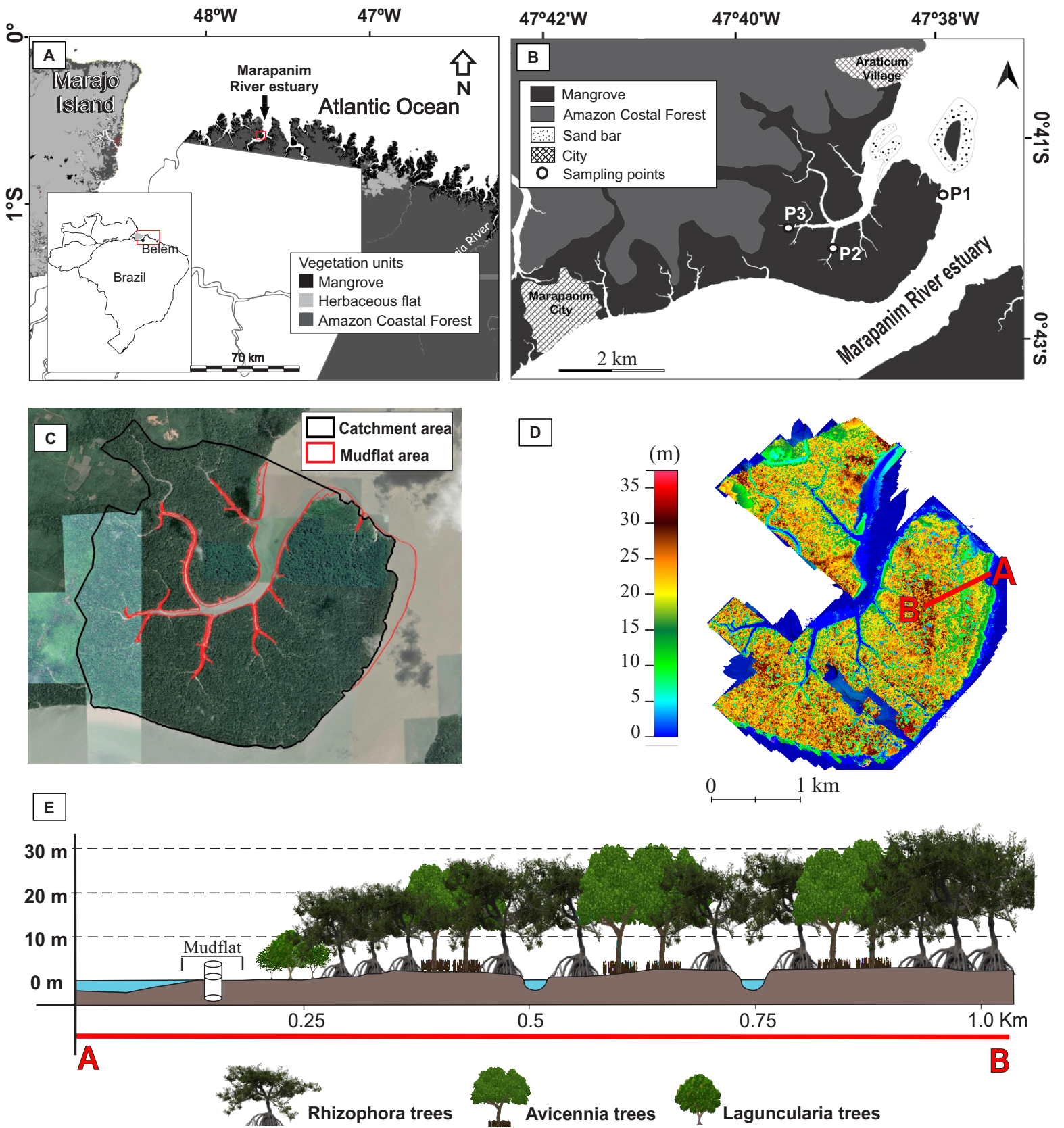


Figure 2

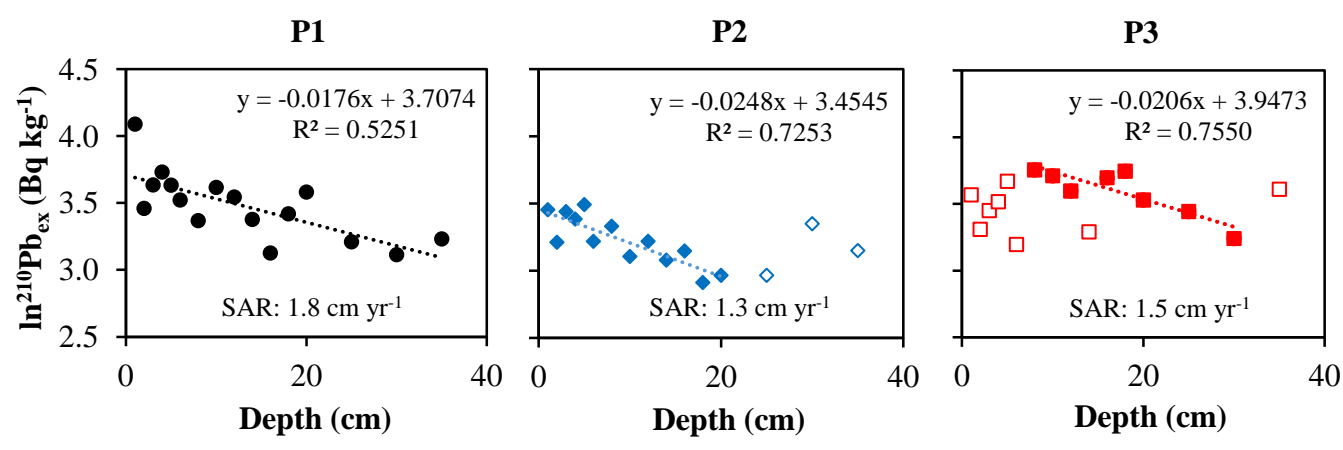
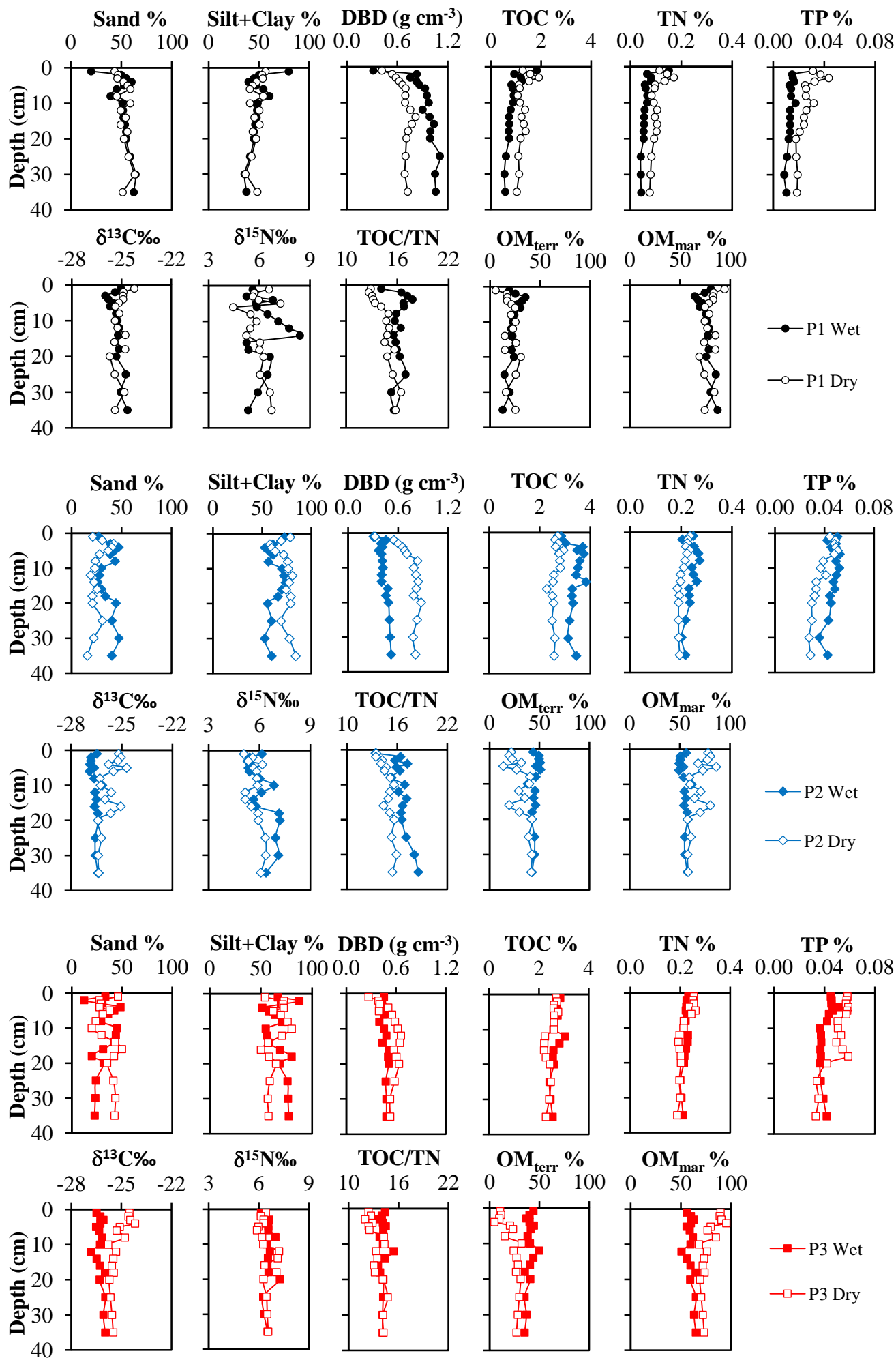


Figure 3





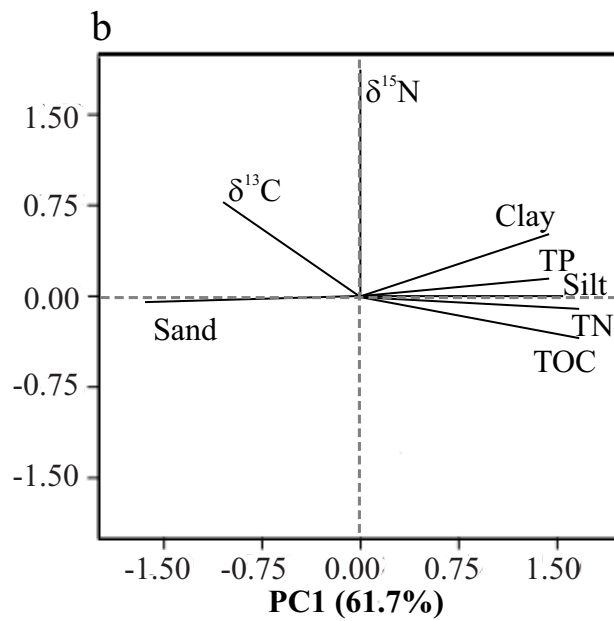
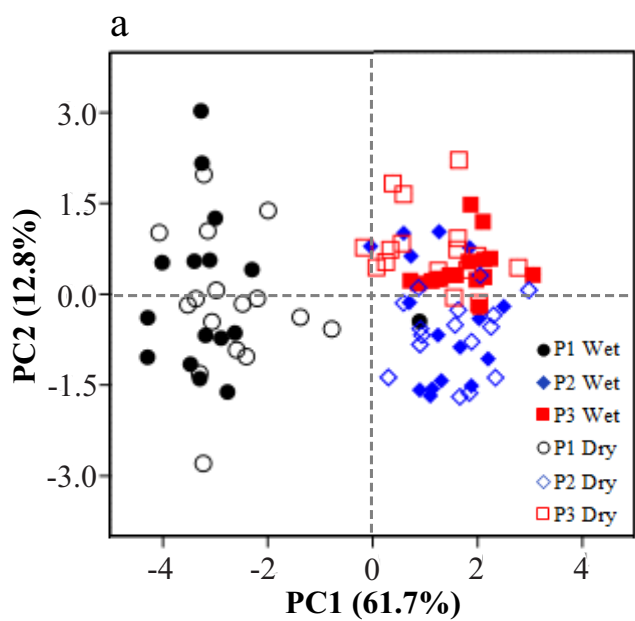


Figure 5

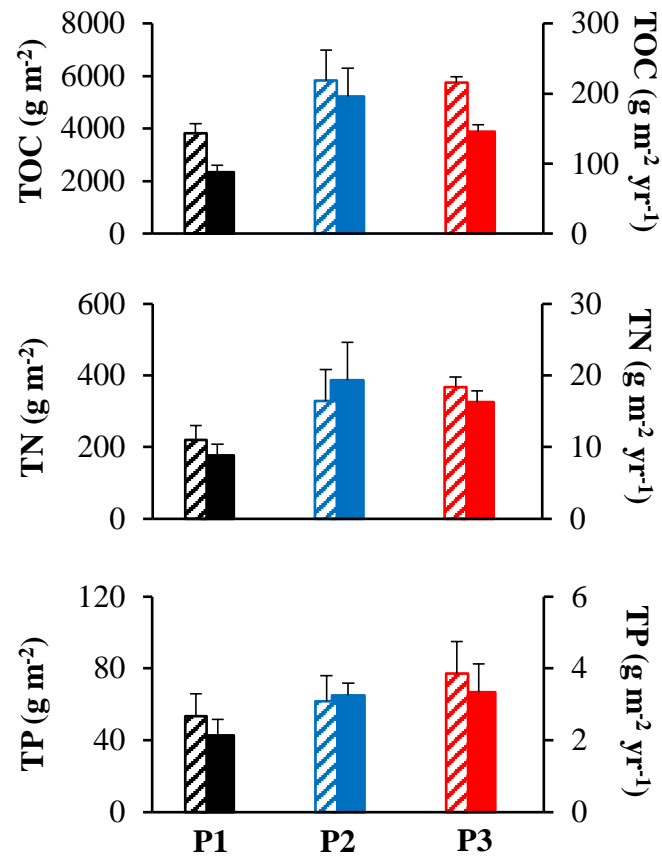


Figure 6

