

Spatial and seasonal variation of microphytoplankton community and the correlation with environmental parameters in a hypereutrophic tropical estuary - Maranhão - Brazil

Ana Karoline Duarte dos Santos¹, Amanda Lorena Lima Oliveira², Jordana Adorno Furtado², Francinara Santos Ferreira², Bethânia de Oliveira Araújo², James Jordan Marques Corrêa², Lisana Furtado Cavalcanti², Andrea Christina Gomes de Azevedo Cutrim¹, Marco Valerio Jansen Cutrim^{2*}

¹ Universidade Estadual do Maranhão

(Cidade Universitária Paulo VI - Travessa Paulo VI, s/n - 65057-630 - São Luís - MA, Brazil)

² Universidade Federal do Maranhão

(Cidade Universitária Dom Delgado - Avenida dos Portugueses s/n - 65080-805 - São Luís - Maranhão - Brazil)

*Corresponding author: cutrim@ufma.br

ABSTRACT

The Bacanga River Estuary has a hydrodynamic behavior and its tidal flow is limited by a dam. It is considered as a hypertrophic environment that receives daily high loads of domestic sewage without treatment. This study aimed to evaluate the spatial and temporal variation of phytoplankton community and its relationship with environmental parameters. Bi-monthly sampling campaigns were carried out at six fixed sites between 2012 and 2013. Physical-chemical and biological parameters were collected (chlorophyll a, phytoplankton composition and abundance) to perform the statistical correlations. The results indicate that phytoplankton community is mostly represented by diatoms, with *Skeletonema costatum* being the dominant species responsible for bloom in April and June of 2012. The dominance of this species is related to the high silicate concentrations, pH and turbidity. Other blooms events as well as the *Euglena gracilis* and *Chlamydomonas* sp. were recorded in February 2013, when the total phosphorus concentrations were high and the dissolved oxygen concentrations were higher. Dinoflagellates, cyanobacteria and diatom *Thalassiosira* sp. were widely distributed in the dry period and highly correlated with salinity, water transparency and nutrients. Hence, the distribution of phytoplankton community is more defined seasonally, rather than spatially.

Descriptors: Algal bloom, Microalgae, Nutrients.

RESUMO

O estuário do rio Bacanga apresenta um comportamento hidrodinâmico com fluxo de marés limitado por uma barragem. Ele é considerado como um ambiente hipereutrótico que recebe diariamente altas cargas de esgoto doméstico sem tratamento. Este trabalho teve como objetivo avaliar a variação espaço-sazonal da comunidade fitoplanctônica e suas relações com parâmetros ambientais. Amostragens bimestrais foram realizadas em seis pontos fixos entre 2012 e 2013, obtendo valores dos parâmetros físico-químicos e biológicos (clorofila a, composição e abundância do fitoplâncton) para realização das análises estatísticas. Os resultados indicam que a comunidade fitoplanctônica é representada por diatomáceas, sendo *Skeletonema costatum* a espécie dominante responsável por pulsos de florações em abril e junho de 2012. O predomínio dessa espécie está relacionado aos elevados teores de silicato, pH e turbidez da água. Outros eventos de florações como da *Euglena gracilis* e *Chlamydomonas* sp. foram registrados em fevereiro de 2013, quando os teores de fósforo total estiveram elevados e as taxas de oxigênio dissolvido foram superiores. Os dinoflagelados, cianobactérias e a diatomácea *Thalassiosira* sp. apresentaram ampla distribuição no período de estiagem e estão altamente correlacionados com a salinidade, transparência da água e nutrientes. Desta forma, a distribuição da comunidade fitoplanctônica é mais definida sazonalmente que espacialmente.

Descritores: Florações algas, Microalgas, Nutrientes.

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INTRODUCTION

Estuaries are hydrodynamic coastal environments, where the freshwater flow of rivers is mixed with the salt water from the oceans causing extensive environmental fluctuations (NASCIMENTO et al., 2003; MASUDA et al., 2011). Those conditions create a biologically productive system, considered “hotspots” in the cycling and export of nutrients, increasing the rates of biological metabolism and the local primary production (GREGO et al., 2009; CALLAWAY et al., 2014; CLOERN et al., 2014).

Those environments are such good places for ecological studies, not only because of the spatial and temporal variations of biotic and abiotic processes, but also because they are considered as the most degraded environments due the coastal urban development, which leads to aquatic eutrophication (MATOS et al., 2011; CAREY et al., 2013; WILD-ALLEN et al., 2013).

The high load of nutrients in coastal habitats promotes an increase in the magnitude and frequency of phytoplankton blooms, affecting the biogeochemistry and structure of the trophic web being a challenge for the continued activity of these systems (PAERL et al., 2014; LU; GAN, 2015; SIN et al., 2015). Thus, the phytoplankton acts as an outstanding environmental indicator (GUENTHER et al., 2015), once it is the transfer link and energy in estuarine trophodynamics. The phytoplankton community is capable of changing its composition and structure in response to different modifications of the environmental conditions caused by humans (REYNOLDS, 2006; GREGO et al., 2009; SEGURO et al., 2015).

This study was conducted in a hypereutrophic tropical estuary (SILVA et al., 2014), located in a densely populated area of the São Luís Island. The area receives daily high loads of domestic sewage without treatment (Instituto da Cidade, Pesquisa e Planejamento Urbano e Rural - INCID). The estuary of the Bacanga River has a hydrodynamic behavior of raising and lowering of tides limited by a transverse dam installed in its mouth. The opening and closing of the dam gates is irregular, which hinders the water level control up to 4 meters (PITOMBEIRA; MORAIS, 1977). Studies about the microphytoplankton community are scarce, unlike the ones about the use and occupation of soil, degradation and availability of water resources, geomorphology and rehabilitation of degraded areas (CARVALHO et al., 2000; POTAZIO et al., 2004; RIBEIRO et al., 2005; COELHO; DAMÁZIO, 2006; ANDRADE; CASTRO, 2007; ROJAS

et al., 2007; BEZERRA et al., 2009; SILVA et al., 2014). In this context, this study was established as the first record of the phytoplankton in that area, in order to evaluate the spatial and temporal variation of that community and its relationship with environmental parameters over an annual cycle.

MATERIAL AND METHODS

STUDY AREA

The Bacanga River Estuary is located in the northwest portion of the São Luís Island, Maranhão (2°32'26" 2°38'07"S and 44°16'00" and 44°19'16"W) and from its source to its mouth it is about 22 Km covering an area of 11,030 hectares and then it flows into São Marcos Bay. It is featured as a shallow system formed by two tributaries (Bicas River and Gapara River), which together are part of an Environmental Protection Area called Bacanga State Park (Figure 1). The area has a tropical humid climate with high temperatures throughout the year, represented by two well-defined seasonal periods: rainy season (January to July) and dry season (August to December) (AZEVEDO et al., 2008).

Samples were collected at six sites (Figure 1) along the estuary, following the annual cycle between 2012 and 2013. Six bimonthly samples were taken, three of which occurred in the rainy season (April/12, June/12 and February/13) and three in the dry season (August/12, September/12 and December/12). The sampling followed the normal inflow of seawater (downstream-upstream) during the ebb tide in compliance with the lunar cycle.

QUALITY WATER AND METEOROLOGICAL ANALYSIS

The evaluation of water quality was based on physical and chemical parameters, such as water temperature, pH, salinity and total dissolved solids (TDS) were obtained with the aid a multi-parameter probe (Hanna-9878). The water transparency was obtained using the Secchi disk and turbidity was measured using a turbidimeter (Lamotte 2020), these parameters were analyzed *in situ*. For nutrient analysis 2 liters of water were collected at the surface layer (50 cm depth) using a Von Dorn bottle. Quantification of DIN ($\text{NH}_4^+ + \text{NO}_2^- + \text{NO}_3^-$), DIP (PO_4^{3-}), total phosphorus (TP) and inorganic silicate (SiO_2^{3-}) followed the methodology described in “Standard Methods for Water and Wastewater” (APHA, 2012). Total phosphorus and nitrate were determined using the methodology described

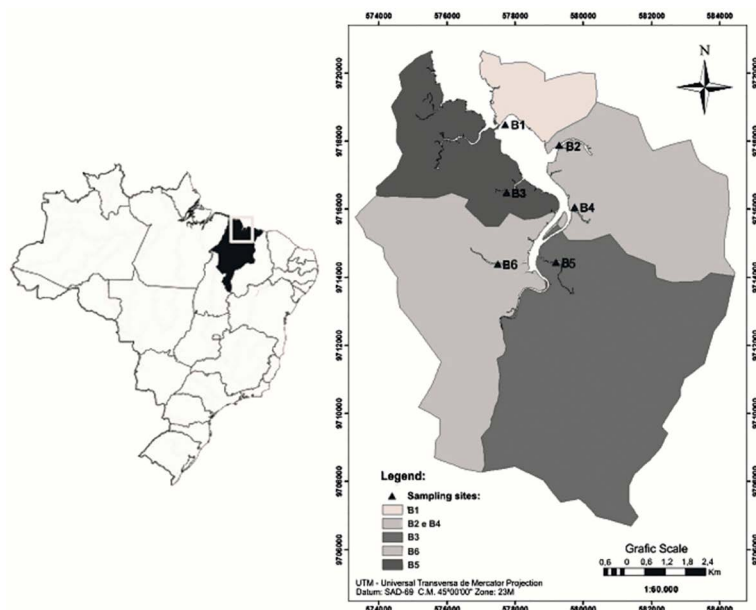


Figure 1. Estuary of Bacanga River and sampling sites (B1, B2, B3, B4, B5 e B6).

by KOROLEFF (1983). Nitrite determination was based on the method of STRICKLAND and PARSONS (1972), silicate and orthophosphate (GRASSHOFF et al., 1983). Dissolved oxygen was determined by the chemical method of Winkler modified by GOLTERMAN et al. (1978). The historical mean (2003-2013) and total monthly rainfall were obtained by the Laboratory of Meteorology of the Maranhão State University (LabMet/UEMA).

PHYTOPLANKTON COMPOSITION

For qualitative analysis of microphytoplankton, horizontal hauls were taken in the euphotic zone (20 cm depth). The samples were stored in plastic bottles and kept with buffered formalin 4%. In the laboratory, the identification was made through mounting of temporary and permanent slides (MÜLLER-MELCHERS; FERRANDO, 1956), using the recommended bibliography. For classification and updating of the taxa the website AlgaeBase was used (<http://www.algaebase.org>).

The frequency of occurrence of taxa was calculated according to MATEUCCI and COLMA (1982), using the ratio between the numbers of samples in which each taxon occurred, and the total number of samples tested. The following categories were obtained: very frequent ($\geq 75\%$), frequent ($< 75\%$ and $\geq 50\%$) occasional ($< 50\%$ and $\geq 25\%$) and rare ($< 25\%$).

CHLOROPHYLL A AND MICROPHYTOPLANKTON DENSITY

To obtain the chlorophyll data, two liters of water were collected in the surface and placed in matte vials and then they were vacuum filtered through glass-fiber filters Whatman GF/C (0.48 μm pore size and 24 mm diameter). Two total filtrations and two fractionated filtrations were performed resulting in the largest and in the smallest fraction of the phytoplankton, microphytoplankton ($> 20\mu\text{m}$) and nano/pico phytoplankton ($< 20\mu\text{m}$), respectively. For the samples fractioning, it was used a screen with a mesh aperture of 20 μm . The filtrate volume ranged from 0.10 to 0.25 liters. After drying, the filters were kept in a freezer (-18°C) until the extraction of the chlorophyll pigments using the method of spectrophotometry (UNESCO, 1966) and it was used the calculation suggested by STRICKLAND and PARSONS (1972).

The sampling for phytoplankton density followed the same methodology used to obtain chlorophyll. However, 250 mL aliquots in Lugol's solution were kept until cell counting through the Utermöhl technique (FERRARIO et al., 1995). Based on the cell counts per liter, ecological diversity indexes were calculated (SHANNON; WIENNER, 1963) according to the classification to VALENTIN (2000), species richness (MARGALEF, 1958) and species evenness (PIELOU, 1966).

STATISTICAL ANALYSIS

PERMANOVA and Two-way ANOVA were performed in order to determine the significant differences ($p < 0.05$) of the physical-chemical factors among seasonal periods and sampled sites. Principal Component Analysis was applied to standardized data in order to obtain the correlation among the environmental variables. To see the correlation of environmental variables among dominant species groups in the counting of cell per liter, it was used the Canonical Correspondence Analysis not rectified (CCA). Pearson's Correlation Analysis was performed to correlate the phytoplankton density in relation to different physical-chemical factors analyzed. SIMPER analysis was used to determine the contribution of taxa between seasonal periods. All statistical analyzes were performed with the aid of the software STATISTIC 10.0 and Past 2.16.

RESULTS

PHYSICAL AND CHEMICAL WATER PARAMETERS AND METEOROLOGICAL CHARACTERISTICS

The characteristics of the estuarine waters of the Bacanga River are summarized in Table 1, with significant spatial and temporal differences of environmental variables according to the PERMANOVA analysis ($F=6.51$ and $p=0.0001$).

The pH (overall average 8.35 ± 0.56 ; $F=1.59$, $p > 0.05$) and the dissolved oxygen (6.06 ± 0.56 mg L⁻¹; $F=0.65$, $p > 0.05$) showed no significant temporal and spatial differences. However, the salinity values (20.43 ± 0.04 ; $F=34.79$, $p < 0.001$) were significant regarding the seasonal and spatial variations, showing lower values in the rainy period and increasing in the downstream direction (Table 1).

Turbidity (10.99 ± 4.79 NTU $F=5.77$, $p < 0.05$) and TDS (19.00 ± 6.26 mg L⁻¹; $F=5.51$, $p < 0.05$) were significant in the rainy season, whereas the water transparency (0.76 ± 0.26 m; $F=0.34$; $p > 0.05$) and depth (2.39 ± 2.34 m; $F=0.99$; $p > 0.05$) were significant only spatially. The water temperature (29.99 ± 1.03 °C; $F=0.13$; $p > 0.05$) was significant in regard to the spatial variation with higher values in section 5 (Table 1).

The concentrations of NO₃⁻ (0.13 ± 0.81 μmol L⁻¹; $F=0.65$, $p > 0.05$) showed no significant spatio-temporal differences. However, the concentrations of NO₂⁻ (0.47 ± 0.36 μmol L⁻¹; $F=5.03$, $p < 0.05$) were significant considering the seasonal periods and the spatial variation,

showing higher values in the dry season at site 3 (Table 1). The NH₄⁺ ion (84.92 ± 100.44 μmol L⁻¹; $F=5.56$, $p < 0.05$) was significant considering both seasonal and spatial variations, showing higher values in the dry season, specifically at the sites 2 and 4. The DIN showed an overall average of 100.44 ± 85.58 μmol L⁻¹ ($F=5.66$, $p < 0.05$) and had the NH₄⁺ as its main component, indicating a recent pollution of the environment.

TP rates (6.08 ± 5.56 μmol L⁻¹; $F=1.08$, $p > 0.05$) were high all the year, and the concentrations of DIP (PO₄⁻³) had an average of 26.53 ± 46.73 μmol L⁻¹ ($F=3.19$, $p > 0.05$). The SiO₂⁻ had an average of 66.24 ± 86.37 μmol L⁻¹ ($F=3.70$, $p > 0.05$) showing no significant difference with DIN: DIS ratio (02.04 ± 1.29 μmol L⁻¹; $F=2.88$, $p < 0.05$). The DIN: DIP molar ratio was lower than 16:1 in both seasonal periods, showing nitrogen limitation. However, the months of February (at the sites 5 and 6) showed high Redfield ratio, having the phosphate as the limiting factor.

The historical mean rainfall (2003 - 2013) showed an average of 1790.43 mm, ranging from 995 mm in 2012 to 2536.10 mm in 2011. It was observed a seasonal cycle well defined with an intensification of the rainy season from January to July and April as the month that presents the highest rainfall events, and a dry period from August to December, in which September is the driest month. The heaviest rainfall occurred in February/2013 (296 mm) and the lowest in October/2012 (2.8) (Figure 2).

PHYTOPLANKTON COMPOSITION

With the qualitative analysis of phytoplankton composition at the investigated area of the estuary of the Bacanga River, 133 phytoplankton species have been distributed in five divisions: Bacillariophyta (64.66%), Dinophyta (12.03%), Cyanobacteria (28.11%), Euglenophyta (27.8%) and Chlorophyta (3.76%) (Figure 3).

Regarding the seasonal distribution, diatoms predominated in the rainy season with a percentage of 67.96% and in the dry season with 61.61%. Dinoflagellates were more representative in the dry season with 14.29% and cyanobacteria in the rainy season with 12.62% (Figure 3). The most representative class was Mediophyceae with 41 species, especially the Chaetocerotaceae family, in which *Chaetoceros* was the most expressive genus with 23 species. Within the dinoflagellate group, eight species of the genus *Protoperidinium* were identified. For the spatial distribution, the diatoms (site 1, 78.31%), dinoflagellates (site 2, 3.93%), and cyanobacteria (site 2, 16.79%) showed

Table 1. Statistic summary of environmental parameters collected from surface water in the estuary of the Bacanga River.

	Unit	SPATIAL						SEASONAL	
		1	2	3	4	5	6	Rainy	Dry
DO	mg L ⁻¹	6.60±1.967	5.14±2.05	4.66±1.94	5.36±3.48	6.25±2.29	8.39±5.69	6.52±3.36	5.60±3.06
pH	---	8.75±0.592	8.44±0.39	8.30±0.35	8.17±0.53	8.35±0.53	8.13±0.86	8.48±0.72	8.22±0.30
Sal.	---	24.73±8.536	22.71±6.85	23.01±8.99	23.34±8.36	14.68±8.95	14.16±9.44	14.47±7.73	24.40±5.78
Turb.	NTU	7.01±3.01	10.22±4.13	11.28±3.34	13.01±3.58	10.63±3.70	13.83±7.91	12.69±5.25	9.30±3.69
Sec.	M	1.07±0.30	0.65±0.19	0.69±0.07	0.69±0.14	0.77±0.22	0.68±0.35	0.73±0.28	0.78±0.23
Depth	M	2.04±1.975	4.21±2.28	1.71±1.88	4.25±3.16	0.69±0.66	1.48±1.43	2.75±2.29	2.04±2.39
TDS	mg L ⁻¹	21.25±5.902	17.96±5.76	19.25±7.10	19.97±6.18	17.03±6.20	18.58±8.15	23.97±2.79	21.52±4.27
T ^o W	°C	28.80±0.66	29.50±1.05	29.65±1.25	31.13±0.76	30.47±1.34	30.63±1.15	30.18±0.77	29.81±1.29
SiO ₂ ⁻	µmol L ⁻¹	63.94±123.76	52.48±44.15	39.28±46.44	86.16±101.87	69.99±71.64	85.63±123.47	96.16±108.39	36.32±41.41
NO ₃ ⁻ -N	µmol L ⁻¹	0.81±1.97	0.00002±0.00003	0.00002±0.00003	0.00002±0.00003	0.00003±0.00003	0.00002±0.00003	0.28±0.00003	0.14±0.00003
NO ₂ ⁻ -N	µmol L ⁻¹	0.29±0.17	0.38±0.09	0.92±0.61	0.39±0.16	0.28±0.13	0.56±0.36	0.57±0.35	0.43±0.24
NH ₄ ⁺ -N	µmol L ⁻¹	45.85±33.447	195.23±163.61	44.37±37.60	160.83±89.40	19.33±14.22	44.28±24.78	111.48±58.47	113.93±79.39
TP	µmol L ⁻¹	3.36±2.22	7.78±2.78	3.34±1.81	10.04±9.79	5.38±4.03	6.59±6.93	0.22±0.14	0.16±0.19
DIN	µmol L ⁻¹	46.94±34.48	195.61±163.78	45.29±37.91	161.22±89.34	19.61±14.31	44.85±25.08	112.33±113.0	58.83±79.38
DIP	µmol L ⁻¹	1.77±2.62	1.82±2.35	0.91±0.97	1.56±2.23	3.33±5.28	4.35±8.14	3.22±5.58	1.36±1.72
Chl-a	µg L ⁻¹	49.87±47.86	31.03±31.09	24.15±17.03	32.00±30.65	27.67±38.61	38.73±56.20	49.52±46.14	18.28±13.79

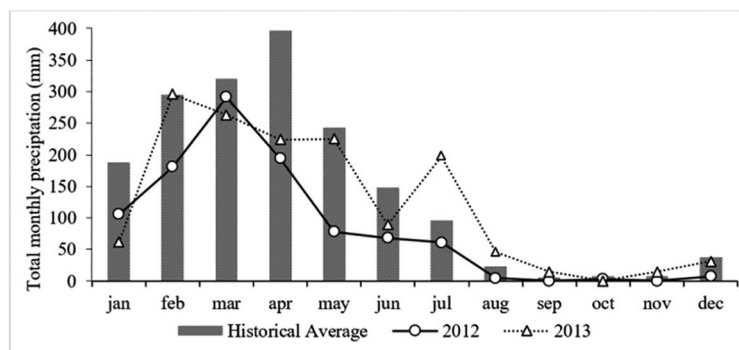


Figure 2. Historical mean (2003-2013) and total monthly rainfall during the study period (April 2012 to February 2013).

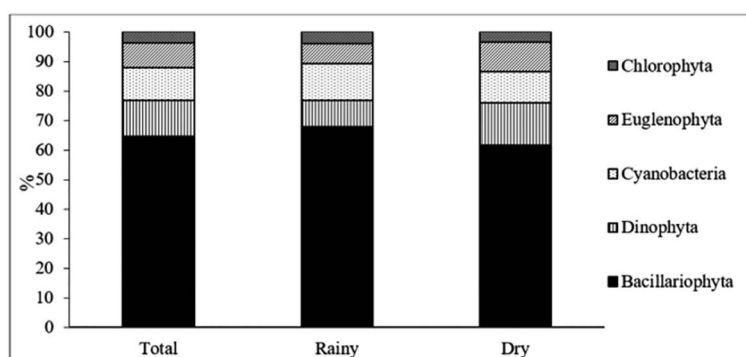


Figure 3. Seasonal variation of phytoplankton distribution by representative group in the estuary of the River Bacanga.

higher contribution at the sites closer to the dam; however, euglenophytes (site 4, 13.74%) and chlorophytes (site 5, 2.27%) at the sites further from the dam (Table 2).

In relation to the occurrence frequency, 81.20% of the species were classified as rare, 10.53% as occasional, 2.06% as frequent and only 2.26% as very frequent. Within this group, *Gyrosigma balticum*, *Navicula* sp. and *Nitzschia sigma* occur in 100% of the samples. As for seasonal variation, *Oscillatoria principes* occurred in 100% of the samples collected in the rainy season, whereas *Coscinodiscus rothii* and *Protoperdinium* sp₁ occurred in 100% of the dry season samples.

CHLOROPHYLL A AND MICROPHYTOPLANKTON DENSITY

CHLOROPHYLL A

Chlorophyll *a* concentrations ranged from $10.06 \pm 5.44 \mu\text{g L}^{-1}$ (dry season - site 5) to $73.29 \pm 61.42 \mu\text{g L}^{-1}$ (rainy season - site 1) and it was more significant in the rainy season ($F=5.99$ and $p<0.05$) (Figure 4).

The microphytoplankton was the main contributor to the photosynthetic process in 47.22% of the samples, ranging from 1.69 ± 0.16 at site 3 during the dry season, to $54.97 \pm 54.71 \mu\text{g L}^{-1}$ at site 1 during the rainy season with a higher level of significance ($F=4.35$ and $p<0.05$). However, the nano/pico phytoplankton ($< 20\mu\text{m}$) was not significant between seasonal periods and the sampled sites. It was considered the most representative portion of the sample 52.78%, with the lowest value of $7.67 \pm 5.40 \mu\text{g L}^{-1}$ in the dry season and the higher value of $41.60 \pm 51.18 \mu\text{g L}^{-1}$ ($F=3.54$ and $p>0.05$) in the rainy season, both at site 5 (Figure 5).

DENSITY OF MICROPHYTOPLANKTON

The phytoplankton density was lower during the dry season ($369.024 \pm 203.786 \text{ cell L}^{-1}$) at site 3 and higher in the rainy season ($15,671,680 \pm 27,039,949 \text{ cell L}^{-1}$) at site 6 ($F=1.47$; $p>0.05$) (Figure 6), and in the rainy season occurred the major algal blooms.

Some algal bloom events were recorded over the study period. The increase in algae recorded at sites 1 and 2 was

Table 2. Composition, taxonomic classification and occurrence of microphytoplankton identified in the estuary of the Bacanga River (1, 2, 3, 4, 5 and 6 sites).

TAXON	RAINY	DRY	TAXON	RAINY	DRY
CYANOBACTERIA			<i>Protoperidinium</i> sp ₁	1, 2, 3, 4 e 5	ALL
<i>Chroococcus turgidus</i>		1,3,4,5	<i>Protoperidinium</i> sp ₂	2, 3, 4 e 6	1 e 2
<i>Lyngbya</i> sp.	2,3,5 e 6	ALL	<i>Protoperidinium</i> sp ₃	1, 3, e 5	1, 4 e 5
<i>Oscillatoria limosa</i>	2, 3, 4 e 5	2, 3, 4 e 5	<i>Protoperidinium</i> sp ₄		1, 2 e 3
<i>Oscillatoria princeps</i>	ALL	ALL	<i>Protoperidinium</i> sp ₅		1
<i>Oscillatoria</i> sp ₁	2, 4, e 6	1,2 e 4	<i>Protoperidinium</i> sp ₆		1, 5 e 6
<i>Oscillatoria</i> sp ₂	3	2 e 3	<i>Pyrophacus steinii</i>		4 e 5
<i>Spirulina</i> sp.	2 e 6		<i>Tripos lineatus</i>	1	1
<i>Planktothrix agardhii</i>		2 e 5	<i>Tripos macroceros</i>		3
<i>Geitlerinema amphibium</i>	1, 2, 3 e 6	2 e 6	BACILLARIOPHYTA		
<i>Microcystis botrys</i>	5		<i>Achnanthes brevipes</i>	1	
<i>Microcystis viridis</i>	2	5	<i>Amphora proteus</i>	2 e 3	6
<i>Thricodesmium</i> sp ₁	3	2, 3 e 6	<i>Amphora</i> sp.	ALL	1, 3, 4, 5 e 6
<i>Limnococcus limneticus</i>	5		<i>Bellerochea malleus</i>	1, 2, 3 e 4	4
<i>Limnococcus limneticus</i>	5		<i>Chaetoceros affinis</i>	1, 3, 4, 5 e 6	1, 3 e 4
<i>Merismopedia elegans</i>	2,4 e 5	2,3 e 5	<i>Chaetoceros atlanticus</i> f. <i>audax</i>	1 e 3	
<i>Leptolyngbya tenuis</i>	5	1, 2, 3, 5, e 6	<i>Chaetoceros brevis</i>	1 e 3	2
DINOPHYTA			<i>Chaetoceros compressus</i>		1
<i>Dinophysis</i> sp.	1 e 3	1, 2, 3, 5 e 6	<i>Chaetoceros constrictus</i>	3 e 4	1, 2 e 3
<i>Gonyaulax</i> sp.		5	<i>Chaetoceros curvisetus</i>	1, 2, 3 e 4	1, 2 e 4
<i>Gymnodinium</i> sp.	1 e 3	2, 3, 4 e 5	<i>Chaetoceros danicus</i>	1 e 3	
<i>Oxytoxum</i> sp.	1 e 3	1, 4 e 5	<i>Chaetoceros debilis</i>	3	
<i>Prorocentrum micans</i>		4	<i>Chaetoceros decipiens</i>	1 e 3	1 e 2
<i>Protoperidinium divergens</i>	1 e 3	1, 2, 3 e 5	<i>Chaetoceros densus</i>	6	
<i>Protoperidinium steinii</i>	2	ALL	<i>Chaetoceros didymus</i>	1, 4 e 6	1, 3 e 4
<i>Chaetoceros diversus</i>	3		<i>Entomoneis alata</i>	ALL	ALL
<i>Chaetoceros gracilis</i>	1		<i>Gyrosigma balticum</i>	ALL	ALL
<i>Chaetoceros holsaticus</i>	ALL	3	<i>Hantzschia amphioxys</i>	5	5
<i>Chaetoceros lorenzianus</i>	1, 2, 3 e 6	1, 2, 3 e 4	<i>Helicotheca tamesis</i>	3	1, 2, 3, 4 e 5
<i>Chaetoceros pendulus</i>	1		<i>Lithodesmium undulatum</i>		1
<i>Chaetoceros peruvianus</i>		3	<i>Lyrella lyra</i>	3 e 5	
<i>Chaetoceros pseudocurvisetus</i>	1 e 4	1, 2, 3 e 4	<i>Melosira nummuloides</i>	1, 3 e 4	
<i>Chaetoceros rostratus</i>	1 e 2		<i>Navicula</i> sp.	ALL	ALL
<i>Chaetoceros simplex</i>	ALL	ALL	<i>Navicula pennata</i>	3, 4 e 5	1 e 3
<i>Chaetoceros subtilis</i>	1, 2 e 4	6	<i>Nitzschia fasciculata</i>	1 e 5	3, 5 e 6
<i>Chaetoceros subtilis</i> var. <i>abnormis</i>	1, 3, 4 e 5	1 e 3	<i>Neostreptotheca subindica</i>		2 e 3
<i>Chaetoceros teres</i> Cleve	1, 3, 4 e 5	ALL	<i>Pleurosigma normanii</i>	5	2
<i>Coscinodiscus curvatus</i>	1, 2, 3 e 4	ALL	<i>Pleurosigma</i> sp ₁	ALL	ALL
<i>Coscinodiscus centralis</i>	1, 3, 4 e 5		<i>Pleurosigma</i> sp ₂	1, 2, 3 e 4	ALL
<i>Coscinodiscus granii</i>	1 e 2		<i>Pseudo-nitzschia pungens</i>	1	1 e 2
<i>Coscinodiscus radiatus</i>	4 e 6	2	<i>Paralia sulcata</i>		3
<i>Coscinodiscus rothii</i>	ALL	ALL	<i>Pleurosigma angulatum</i>	2, 3 e 6	

Continued **Table 2.**

<i>Cyclotella stelligera</i>	1 e 5		<i>Pleurosigma elongatum</i>	5	
<i>Cyclotella striata</i>	3	5 e 6	<i>Pleurosigma formosum</i>		1, 2, 3 e 5
<i>Cyclotella meneghiniana</i>	ALL	3, 5 e 6	<i>Pseudo-nitzschia</i> sp.	2	1, 3 e 5
<i>Cylindrotheca closterium</i>	1, 3, 4, 5 e 6	2, 3, 5 e 6	<i>Rhizosolenia setigera</i>		3
<i>Cymbella</i> sp.	3, 4 e 6	1, 2, 4 e 6	<i>Skeletonema costatum</i>	ALL	ALL
<i>Diploneis gruendleri</i>		3	<i>Skeletonema tropicum</i>	1, 2, 3, 4 e 5	2 e 6
<i>Diploneis ovalis</i>		3, 5 e 6	<i>Synedra</i> sp.		1
<i>Diploneis weissflogii</i>		2	<i>Tabularia fasciculata</i>	1	6
<i>Ditylum brightwellii</i>		2	<i>Thalassiosira eccentrica</i>	ALL	1, 3 e 6
<i>Thalassiosira leptopus</i>	1, 2, 4 e 5	1, 2 e 6	EUGLENOPHYTA		
<i>Thalassiosira nanolineata</i>	4	4	<i>Euglena gracilis</i>	4, 5 e 6	5
<i>Thalassionema nitzschioides</i>	1	1	<i>Euglenaformis proxima</i>	4, 5 e 6	3
<i>Triceratium favus</i>	1, 2 e 4	2	<i>Euglena</i> sp ₁	4, 5 e 6	1, 4, 5 e 6
<i>Trieres mobiliensis</i>		1	<i>Euglena</i> sp ₂	4	6
<i>Trieres regia</i>	1, 2, 4 e 5	1, 2, 3 e 4	<i>Lepocinclis acus</i>		3
<i>Tryblionella granulata</i>	4 e 5	3	<i>Lepocinclis oxyuris</i>	2, 3 e 4	1, 2 e 6
<i>Tryblionella punctata</i>	4	5 e 6	<i>Peranema</i> sp.		2
<i>Tryblioptychus cocconeiformis</i>		5	<i>Phacus longicauda</i>		1 e 2
<i>Ulnaria ulna</i>	1 e 3	2 e 5	<i>Phacus</i> sp.	3	4
CHLOROPHYTA			<i>Trachelomonas armata</i>	4 e 5	5 e 6
<i>Chlamydomonas</i> sp.	3	6	<i>Trachelomonas</i> sp.		3
<i>Desmodesmus quadricaudatus</i>	5	2			
<i>Pediastrum duplex</i>	3				
<i>Mougeotia</i> sp.	5	1			
<i>Spirogyra</i> sp.		5			

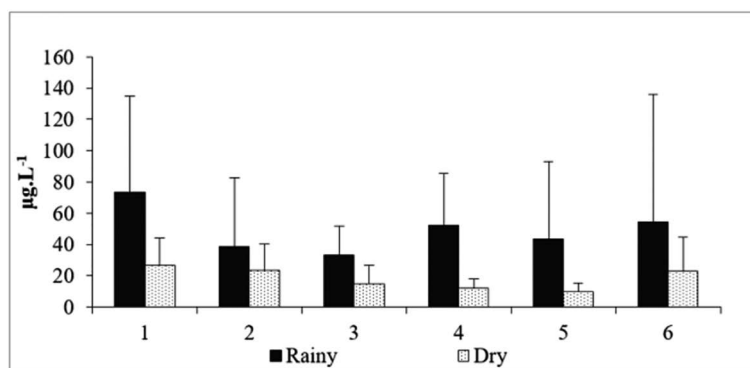


Figure 4. Spatial-temporal variation of the total chlorophyll concentrations in the estuary of the River Bacanga.

due to the specie *Skeletonema costatum* ($814,775 \pm 1,968,950$ cells L^{-1}) in April and June 2012. While the proliferation of *Euglena gracilis* ($19,144,334 \pm 7,523,378$ cells L^{-1}) and *Chlamydomonas* sp. ($731,154 \pm 310,224$ cells L^{-1}) occurred at site 6 in February 2013. In the rainy season, the number of *Thalassiosira* sp. ($F=5.12$ and $p=0.03$) and dinoflagellates ($F=4.10$ and $p=0.005$) decreased significantly (Table 3).

Regarding ecological indexes, the Bacanga River estuary showed a low species diversity (H') with values ranging from 0.90 ± 0.58 bits. $cell^{-1}$ at site 6 during the dry season to 1.87 ± 0.56 bits. $cell^{-1}$ at site 5 in the rainy season. The richness was very low throughout the year, with the lowest value of 1.33 ± 0.58 in the rainy season at site 6 and the highest value of 2.60 ± 0.35 in the dry season at site

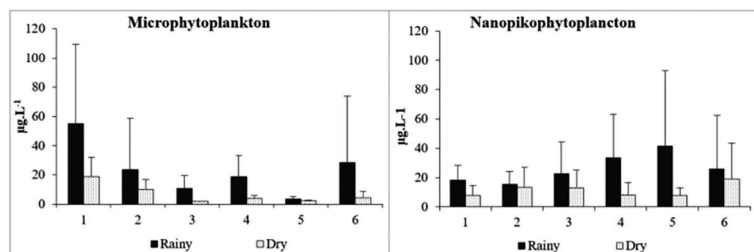


Figure 5. Spatial-temporal variation of chlorophyll concentrations ($\mu\text{g L}^{-1}$) of microphytoplankton and nanophytoplankton in the estuary of the River Bacanga.

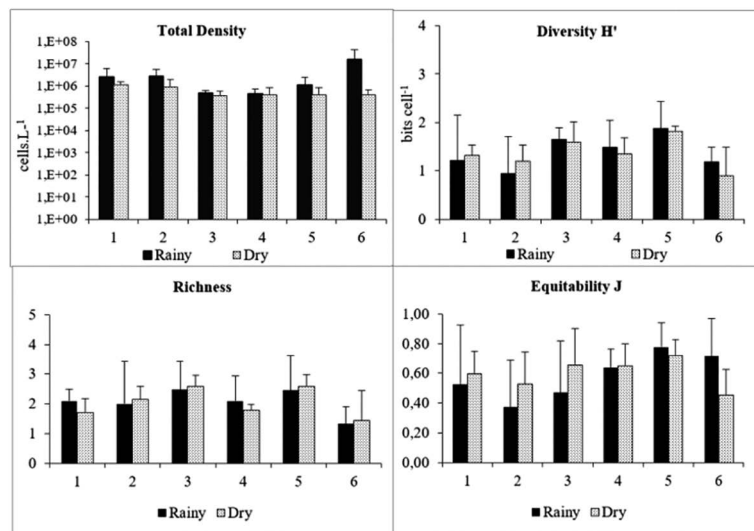


Figure 6. Variation spatial-temporal density of phytoplankton and ecological index in the estuary of the River Bacanga.

Table 3. Mean values of phytoplankton density ($10^3 \text{ cells L}^{-1}$) in the estuary of the Bacanga River. (-) absent.

Species	SPATIAL						SEASONAL	
	1	2	3	4	5	6	RAINY	DRY
<i>Skeletonema costatum</i>	1178±2558	1201±2371	51±83	11±19	16±20	10±18	815±1969	7±13
Dinoflagellates	292±504	66±115	40±88	69±169	42±90	4±8	12±36	159±313
<i>Euglena gracilis</i>	50±88	247±590	25±39	68±151	413±893	5341±13054	1914±7523	134±343
<i>Chlamydomonas</i> sp.	-	-	-	-	-	2193±5373	731±3102	-
Cyanobacteria	12±14	41±69	7±15	2±4	5±9	2±4	4±9	20±42
Group <i>Chaetoceros</i>	153±127	37±35	37±35	35±32	56±97	219±533	130±301	49±92
Group <i>Thalassiosira</i>	93±202	93±192	146±151	93±101	128±164	141±178	55±85	176±190
TOTAL	1178±2558	1201±2371	51±83	11±19	16±20	10±18	815±1969	7±13

3. The evenness (J') ranged from 0.37 ± 0.31 (site 2) to 0.78 ± 0.16 (site 5), both in the dry season (Figure 6). The moderate evenness is common to both seasonal periods, with no significant difference between rain and drought.

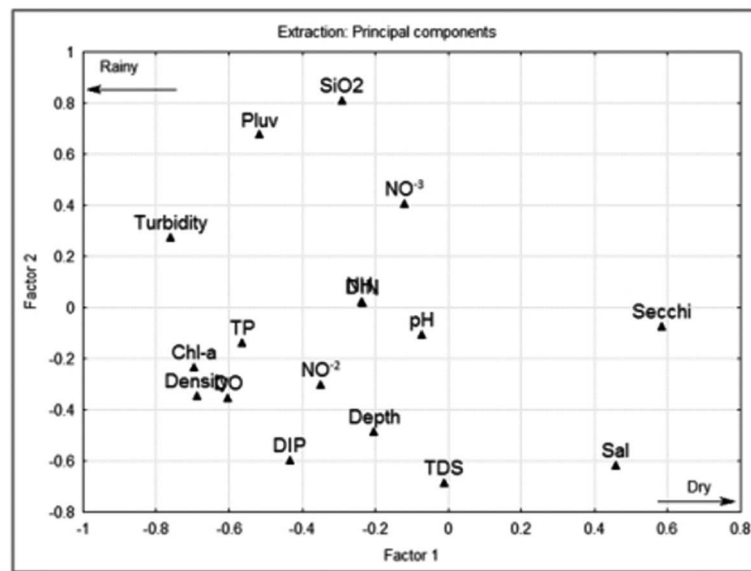
STATISTICAL ANALYSIS

The PCA explained 55.46% of the total variance (Factor 1=21.15%, Factor 2=18.99% and Factor 3=15.31%)

(Table 4) showing a seasonal scenario well defined in relation to environmental variables in factor 1, where the phytoplankton density (-0.68) and chlorophyll a (-0.69) were directly correlated with turbidity (-0.76) and dissolved oxygen (-0.60). Rainfall (0.67) was directly correlated with SiO_2 (0.81) and inversely correlated with salinity (-0.62) in the factor 2. The NH_4^+ (-0.92) and NID (-0.92) showed strong correlation in factor 3 (Figure 7).

Table 4. Principal taxon contributing to the mean dissimilarities between rainy and dry seasons, as determined by the SIMPER analysis.

Taxon	Av. dissim	Contrib. %	Cumulative %	Mean abund. 1	Mean abund. 2
<i>Skeletonema costatum</i>	21.48	25.96	25.96	8.15E+05	7.49E+03
<i>Thalassiosira</i> sp.	19.12	23.10	49.07	5.50E+04	1.76E+05
<i>Euglena gracilis</i>	15.98	19.31	68.38	1.91E+06	1.34E+05
Dinoflagellates	13.16	15.90	84.28	1.23E+04	1.59E+05
<i>Chaetoceros</i> sp.	8.726	10.54	94.82	1.30E+05	4.85E+04
Cyanobacteria	2.731	3.30	98.12	3.61E+03	1.95E+04
<i>Chlamydomonas</i> sp.	1.556	1.88	100	7.31E+05	0.01

**Figure 7.** Principal components analysis performed on environmental parameters, where: Secchi - water transparency; TDS - total dissolved solids; Pluv - pluviosity; DO - dissolved oxygen concentrations; Den - density; T°C - water temperature; NO³⁻ - nitrate concentrations; NO²⁻ - nitrite concentrations; NH₄⁺ - ammonium concentrations; DIN - dissolved inorganic nitrogen concentrations; DIP - dissolved inorganic phosphorus concentrations; SiO₂ - silicate concentrations; TP - total phosphorus concentrations; Chl-a - chlorophyll *a* concentrations.

The Canonical Correspondence Analysis (CCA) explained 80.72% of the relationships between species and environmental variables. The first axis (54.96%) showed that rain favors blooms of *Skeletonema costatum* at sites 1 and 2 and it was directly related to silicate concentrations, pH, and turbidity, causing an increase of chlorophyll concentrations. Also in the first axis, the proliferation of *Euglena gracilis* and *Chlamydomonas* sp. in February/2013 at sites 5 and 6, was strongly influenced by the high levels of total phosphorus that contributed to an increase in dissolved oxygen rates. The second axis 2 (26.45%) showed that the dinoflagellates, cyanobacteria and *Thalassiosira* sp. have a wide distribution in the dry season and are highly correlated with salinity, transparency of water and nutrients (DIP and DIN) (Figure 8).

The SIMPER analysis showed dissimilarity of 82.74% in the distribution of dominant groups of phytoplankton between rainy season and dry season. The species *Skeletonema costatum* (25.96%), *Euglena gracilis* (19.31%), *Chaetoceros* sp. (10.54%) and *Chlamydomonas* sp. (1.88%) were dominant in the rainy season. Whereas the dry season had *Thalassiosira* sp. (23.10%), the dinoflagellates group (15.90%) and cyanobacteria (3.30%) as the most representative (Table 5).

Regarding the linear correlations, the phytoplankton density showed a strong positive correlation with NH₄⁺ (0.80), rain (0.80), DIN (0.81) and NO₃⁻ (0.86), and the DIP showed strong correlation with the rain (0.98). The blooms of *Skeletonema costatum* (0.86), *Euglena gracilis* (0.83) and *Chlamydomonas* sp. (0.86) showed

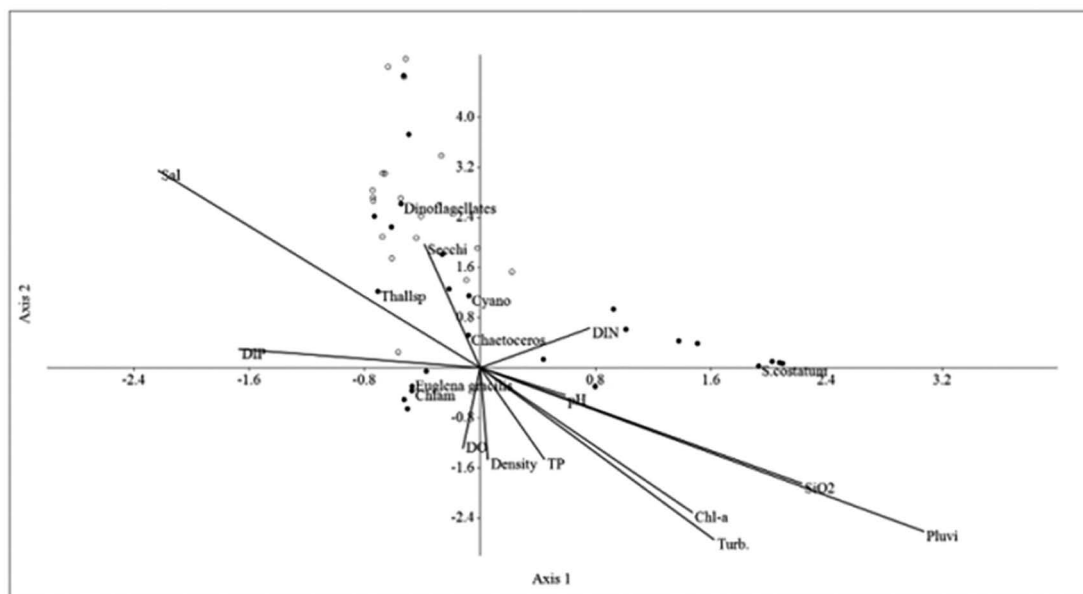


Figure 8. Canonical correspondence analysis performed with environmental parameters and phytoplankton groups, where: Secchi - water transparency; Turb - Turbidity, Pluv - pluviosity, DO - Dissolved oxygen concentrations, DIN - dissolved inorganic nitrogen concentrations, DIP - dissolved inorganic phosphorus concentrations, SiO_2 - silicate concentrations, TP - total phosphorus concentrations, Chl-a - chlorophyll *a* concentration, Sal - salinity, Dino - Dinoflagellates, Ciano - Cyanobacteria, Euglena - *Euglena gracilis*, Chlam - *Chlamydomonas* sp., *S.costatum* - *Skeletonema costatum*, Chaetoceros - *Chaetoceros* group and *Thallsp* - *Thalassiosira* sp.

Table 5. Eigenvalues, proportion of variation explained by each factor and variable loads of Principal Component Analysis (PCA).

	Factor 1	Factor 2	Factor 3
Eigenvalue	3.59	3.22	2.60
% Total variance	21.15	18.99	15.31
DO	-0.60	-0.35	0.42
pH	-0.07	-0.10	0.19
Sal	0.45	-0.62	-0.18
Turbidity	-0.76	0.27	-0.01
Secchi	0.58	-0.07	0.11
Depth	-0.20	-0.48	-0.59
DIS	-0.29	0.81	-0.12
NO-3	-0.12	0.40	0.16
NO-2	-0.33	-0.30	0.02
NH4	-0.23	0.01	-0.92
DIN	-0.23	0.02	-0.92
DIP	-0.43	-0.59	-0.13
TP	-0.56	-0.13	-0.05
Chl-a	-0.69	-0.23	0.36
TDS	-0.01	-0.68	-0.03
Pluv	-0.51	0.67	-0.11
Density	-0.68	-0.34	0.25

high correlation with NO_3^- concentrations. *Thalassiosira* sp. showed a strong relationship with water transparency (0.98), dinoflagellates (0.97) and cyanobacteria (0.82) groups were correlated with pH (Table 6).

DISCUSSION

These results show that environmental variations in the Bacanga River estuary are strongly affected by rainfall, which during the study showed a seasonal cycle well defined. However, it was atypical, according to the historical series (AZEVEDO et al., 2008). That is a typical scenario for the Amazon regimes (MONTEIRO et al., 2016). The following diagram shows the interrelationships among those variables involved and the phytoplankton community, in which the rain was the enhancing agent of nutrient concentrations and the increase in local blooms.

These rainfall variations cause changes in the dynamics of the estuary such as reduced water transparency and salinity (GREGO et al., 2004; MOSER et al., 2005; LEÃO et al., 2008; SANTIAGO et al., 2010). The Bacanga River Estuary was characterized as a shallow system, low water transparency and high turbidity. According to CONLEY et al. (1993) and CLOERN (2001), the increasing of

Table 6. Matrix of the Pearson correlation between environmental parameters and phytoplankton groups, where: Sec - water transparency; NTU - Turbidity, Pluv - pluviosity, DO - Dissolved oxygen concentrations, NO₃ - nitrate concentrations, NO₂ - nitrite concentrations, NH₄ - ammon concentrations, DIN - dissolved inorganic nitrogen concentrations, DIP - dissolved inorganic phosphorus concentrations, Si - silicate concentrations, TP - total phosphorus concentrations, pH - pH, Chla - chlorophyll a concentration, Sal - salinity, Dino - Dinoflagellates, Cyan - Cyanobacteria, Eug- Euglena gracilis, Chlam - Chlamydomonas sp., S.cost - Skeletonema costatum, and Thall - Thallasiosira sp. Significant correlations are indicated in bold text.

	DO	pH	Sal	NTU	Sec	Dep	Si	NO ₃	NO ₂	NH ₄	DIN	DIP	TP	Chla	TDS	Pluv	Den	S.cost	Dino	Eug	Chla	Cyan	Thall
DO	0.36	0.87	0.03	0.42	0.98	0.80	0.55	0.29	0.28	0.28	0.28	0.00	0.17	0.00	0.03	0.65	0.00	0.50	0.13	0.00	0.00	0.80	0.39
pH		0.69	0.07	0.12	0.63	0.09	0.00	0.45	0.93	0.91	0.54	0.93	0.03	0.10	0.96	0.45	0.32	0.97	0.57	0.60	0.82	0.69	
Sal			0.01	0.22	0.28	0.01	0.46	0.58	0.77	0.78	0.41	0.71	0.09	0.00	0.00	0.29	0.24	0.01	0.35	0.35	0.45	0.03	
NTU				0.00	0.96	0.00	0.82	0.55	0.59	0.59	0.88	0.01	0.04	0.88	0.00	0.05	0.59	0.03	0.05	0.06	0.29	0.07	
Sec				0.67	0.18	0.61	0.19	0.49	0.49	0.71	0.04	0.46	0.51	0.29	0.07	0.66	0.20	0.04	0.06	0.95	0.98		
Dep				0.03	0.36	0.21	0.00	0.00	0.03	0.49	0.29	0.24	0.39	0.43	0.09	0.50	0.61	0.57	0.73	0.41			
Si				0.00	0.17	0.28	0.27	0.17	0.82	0.52	0.11	0.00	0.46	0.62	0.25	0.53	0.57	0.61	0.03				
NO ₃				0.50	0.95	0.98	0.58	0.48	0.28	0.23	0.04	0.86	0.87	0.71	0.85	0.87	0.58	0.46					
NO ₂				0.98	1.00	0.01	0.63	0.28	0.99	0.71	0.11	0.86	0.14	0.09	0.09	0.46	0.87						
NH ₄				0.00	0.19	0.39	0.54	0.86	0.86	0.15	0.81	0.74	0.94	0.77	0.81	0.64	0.08						
DIN				0.19	0.39	0.54	0.85	0.85	0.15	0.81	0.74	0.93	0.77	0.82	0.64	0.08							
DIP				0.26	0.03	0.00	0.99	0.04	0.34	0.44	0.02	0.03	0.41	0.46									
TP				0.05	0.91	0.79	0.00	0.65	0.22	0.00	0.01	0.88	0.09										
Chla				0.90	0.90	0.27	0.00	0.00	0.15	0.00	0.00	0.94	0.48										
TDS				0.01	0.26	0.21	0.09	0.18	0.21	0.09	0.18	0.21	0.75	0.03									
Pluv				0.80	0.47	0.10	0.88	0.85	0.26	0.01													
Den				0.48	0.75	0.00	0.00	0.72	0.42														
S.cost				0.53	0.74	0.78	0.89	0.24															
Dino				0.70	0.71	0.65	0.44																
Eug				0.00	0.70	0.54																	
Chla				0.71	0.46																		
Cyan				0.94																			
Thall																							

phytoplankton biomass contributes to reduce water transparency and decreased availability of light energy.

The average concentrations of dissolved oxygen, classifies the estuary area as a low saturation zone, according to MACÊDO and COSTA (1978). However, in February/2013 oxygen super saturation zones were recorded at sites 5 and 6. These high concentrations of dissolved oxygen are related to the high chlorophyll rates (LU; GAN, 2015). According to GUENTHER et al. (2015), this scenario is common to restricted circulation environments and to the high primary production rates, exceeding the depletion of oxygen by heterotrophic organisms.

The pH is alkaline, showing a reduction in the water quality. However, it is within the limits for marine life (NASCIMENTO et al., 2003). The renewal time of Bacanga River Estuary waters depends on the seasonality. During the dry season, the gates remain closed most of the time. This residence time of freshwater in estuaries generates stabilization of the water column, and it is essential to add nutrients and allow faster growth of phytoplankton (ODEBRECHT et al., 2015).

The water temperature is stable in both seasons, which is common in tropical estuaries (MOSER et al., 2005; AZEVEDO et al., 2008; RODRIGUES; CUTRIM, 2010). The rain influences directly the salinity, which showed values significantly lower during the months of April and June, and it causes an increased salinity gradient toward the downstream, similar to that registered by PAMPLONA et al. (2013) in the Amazon estuary. The blooms of *Skeletonema costatum*, *Euglena gracilis* and *Chlamydomonas* sp occur when there is an intensification of the rainfall (LEÃO et al., 2008; BAZIN et al., 2014; JENDYK et al., 2014).

The combination of high nutrient concentrations may limit the growth of phytoplankton on a short time scale (WU; CHU, 2003) and the N: P ratio has been related to the occurrence of algal blooms in coastal waters (SOUZA et al., 2014). Based on that, the nitrogen was shown to be a limiting factor for the diatoms blooms and the phosphorus was a limiting factor for the proliferation of green algae and euglenoids. According to SIN et al. (2015) and SEGURO et al. (2015) it is common the nitrogen being the main limiting factor for marine phytoplankton and the phosphorus for continental phytoplankton. According to RODRIGUES and CUTRIM (2010), phosphorus is usually the main nutrient for microphytoplankton in the estuaries of Maranhão,

suggesting that this community is physiologically well adapted to the conditions of nutrients mixing.

This scenario shows a distribution of the phytoplankton community well-defined seasonally. However, it does not happen spatially, once the algal bloom was potentiated by high concentrations of nutrients. The entry of nutrients can occur naturally through runoff, or through the discharge of domestic sewage (THOMAS et al., 2005; PAERL et al., 2010; SOUZA et al., 2014), and when dominants (ammonium and phosphorus) the N:P ratio is less than the Redfield ratio 16:1 (COELHO et al., 2015; SILVA et al., 2015).

Based on that, the seasonal rainfall in tropical areas produce fluctuations in salinity, nutrient concentrations, turbidity and biological productivity (SANTIAGO et al., 2010) and the ammonium was the main inorganic form of DIN in the Bacanga River Estuary, corroborating with the data of MOSER et al. (2005); PAMPLONA et al. (2013) and SILVA et al. (2015).

According to WU and CHOU (2003) and SIN et al. (2015), high concentrations of nitrogen in the environment may reduce productivity in eutrophic estuaries due to the increase of urban pollution.

Regarding the microphytoplankton composition, diatoms were the most representative group, followed by dinoflagellates (SEGURO et al., 2015), which are transported by tidal currents, and are more frequent in the dry season (PAIVA et al., 2006). The dominance of diatoms is common in estuarine systems due to their euryhaline nature (SENA et al., 2015) and it is correlated directly with the silicate content, since the major source of silica is from the continent and the diatoms frustules have silica in their composition. In estuaries of Maranhão, the predominance of that group occurs in over 80% of the phytoplankton composition (FERREIRA-CORREIA et al., 2004; RODRIGUES; CUTRIM, 2010) and when eutrophication happens, blooms of diatoms become more frequent due to the wide availability of nutrients and high salinity (MATOS et al., 2011). Under these conditions, the local phytoplankton community consists mostly of centric taxa classified as rare and occasional. That is due to the high hydrodynamic environment that can reach areas far from the mouth, that are considered lentic zones (PAIVA et al., 2006; SOUSA et al., 2008; REZENDE et al., 2015; SENA et al., 2015).

Thalassiosira sp. was the most representative species in the dry season, and it is related to the marine influence.

In estuarine ecosystems it is common the events of biomass production by the microphytoplankton, specifically diatoms, which are highly adapted to rapid population growth opportunities in shallow, turbulent, turbid waters and in waters enriched by nutrients (CARSTENSEN et al., 2015).

Among the main representatives of phytoplankton blooms, the *Skeletonema costatum* is evident. It is a cosmopolitan and euryhaline species, one of the most abundant in estuaries, considered opportunistic in eutrophic environments (VILLAC; TENEBBAUM, 2010; MASUDA et al., 2011; REZENDE et al., 2015; BARRERA-ALBA et al., 2016). *Skeletonema costatum* is classified as a harmful species and it is associated with toxic microorganisms in some cases, causing economic losses for aquaculture (KHAN et al., 1998; UNESCO, 2004).

The euglenoids occurred in low salinity and high total phosphorus concentrations during the rainy season (LEÃO et al., 2008; HONORATO DA SILVA et al., 2009). *Chlamydomonas* sp. became dominant also on these same conditions (BAZIN et al., 2014; JENDYK et al., 2014.). This species tolerates a wide range of salinity. However, the highest numbers occur in low salinities, showing that there are other factors influencing the presence of this genus (NCHE-FAMBO et al., 2015).

Coastal estuarine waters are mainly composed of high density of diatoms and dinoflagellates, and green algae and cyanobacteria blooms are restricted to low salinity habitats. These blooms generate important ecological and biochemical implications for the food web and nutrient cycling (CARSTENSEN et al., 2015).

Based on the chlorophyll concentrations the estuary was classified as hypereutrophic, corroborating with SANTOS et al. (2009), SILVA et al. (2014), COTOVICZ JR. et al. (2012) in which the higher values of the pigment were associated with the period of highest rainfall events, unlike TEIXEIRA et al. (1988) and AZEVEDO et al. (2008) in their studies in other estuaries.

Thus, the Bacanga River estuary showed a rapid growth of microphytoplankton due to the discharge of domestic sewage and consequently affected the environmental changes, which conducted to ecological instability scenarios especially in the rainy season, defined by low diversity and species richness. According to VALENTIN et al. (1999), that is a response to the intense eutrophication process in large and sporadic nutrients pulsing favoring the dominance of larger cells. However, when the photosynthetic efficiencies are compared, the

nano/pico phytoplankton are superior and the synthesis is always of large scale (RODRIGUES; CUTRIM, 2010; GUENTHER et al., 2012).

CONCLUSION

The results confirm that the distribution of phytoplankton community is more defined seasonally, rather than spatially. Therefore, the rain is the factor that intensifies the concentrations of nitrogen compounds and phosphate, which favors blooms at different periods. During the rainy season, the *Skeletonema costatum* proliferates using nitrogen, whereas *Euglena gracilis* and *Chlamydomonas* sp. proliferates using phosphate. The *Thalassiosira* sp. is the dominant species in the dry season, which is the period when the residence time in the estuary is higher and domestic effluents are concentrated in the form of ammonium. The environmental impact on the community is evidenced by the low diversity, species evenness and species richness with high chlorophyll concentrations determined by the microphytoplankton. Thus, the eutrophication process promotes algal blooms events that affect the quality of the water and show ecological instability scenarios. This study contributes to a better understanding of the microphytoplankton and suggests the control of the local eutrophication as a mitigation measure for these blooms pulsing. In addition, this study can be used as a model for the characterization and monitoring of the Bacanga River Estuary.

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