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Chapter

Subtropical Coastal Lagoon from Southern Brazil: Environmental Conditions and Phytobenthic Community Structure

Leticia Donadel and Lezilda Torgan

Abstract

The chapter is about the study of environmental conditions and the structure of the benthic diatoms community in Peixe Lagoon, which is inserted in a National Park in southern Brazil. The study was carried out covering four seasons from 2011 to 2012. The system is shallow (<60 cm) located parallel to the coastline, and it is connected to the ocean through a single channel, which occurs naturally or through human action. In this lagoon, during the study, the water temperature ranged between 15.3 and 32.1°C, and the dissolved oxygen presented higher value in the winter (12.5 mg.L⁻¹) and lower value in the summer (7.5 mg.L⁻¹). The lagoon ranged from mesotrophic to hypereutrophic conditions. The salinity varied between 1.3 and 36.2%, and these variations were mainly related to meteorological conditions. The community of diatoms in Peixe Lagoon is composed by 62 taxa distributed in 30 genera composed largely of marine, brackish, and few freshwater species. Among the attributes of the community, composition better reflects the environmental variations. The opening and closing of the channel, salinity, temperature, and the action and direction of the wind are variables influencing the dynamics of the microphytobenthic community.

Keywords: environmental variables, diatoms, community attributes, microphytobenthos, system dynamic

1. Introduction

The shallow coastal lagoons are low depth water column mixing systems in which phytoplankton and microphytobenthos communities, microscopic eukaryotic photosynthetic algae, and cyanobacteria which live on the seabed [1] play a key role in the primary production and recycling of matter and nutrients. The role of microphytobenthos is quite important where macrophytes are absent and light radiation penetrates down to the bottom [2].

Microphytobenthos are composed of a set of microorganisms distributed in very diversified taxonomic groups, among which the diatoms are an important and often dominant component in estuarine and shallow coastal environments. These algae have varied adaptive strategies for adhesion and migration on different substrates, and there is a very large number of species sensitive to environmental changes. Traditionally, benthic diatoms are classified according to the substrate in which they live. Those that live on thin sediment are called epipelic and those that live on sandy substrate are called epipsammic [3].

Diatoms studies in coastal lagoon were mainly concentrated in the world's largest water bodies, the Baltic, Black, and Caspian seas, which are ecosystems impacted by the anthropogenic actions and global climate changes. The eutrophication happened due to the increased nitrogen and phosphorus loads during the last century, and the increase in water temperature related to climate was detected by the changes in subfossil diatom assemblages. The accumulation of heavy metals from surrounded waters can be monitored due to the capacity of these algae to accumulate metals attached to the outside of the cell wall. There is an excellent literature review about these and other impacts; see the Snoeijs and Weckström chapter [4]. We can also find excellent information about the composition, spatial distribution of modern diatom assemblages, diversity, production, and ecology of the sediment-inhabiting diatoms in the estuaries [5–11]. In the smaller shallow lagoon from the east coast of Uruguay (South America) the diatoms studies were used to infer the paleosalinity, trophic and climate changes in relation to the sea level variation [12–16].

So far, most studies concentrated on phytoplankton at Patos Lagoon [17–24], Tramandaí-Armazém Lagoon [25–29], and Peixe Lagoon [30–34]. Regarding microphytobenthic, studies were limited to salt marshes and to the Patos Lagoon estuary [35–43].

The knowledge of the diatoms at Peixe Lagoon began with investigations on diatom assemblages in current and fossil sediments that allowed paleoenvironmental reconstruction. It demonstrated that the lagoon behaved as a deeper and more extensive lagoon system connected to the ocean by one or more permanent linking channels during the Holocene [44]. Later, studies were carried out on the taxonomic composition of diatoms in the marginal sediment of the lagoon. One study emphasizes the genus *Diploneis* Ehrenb. ex Cleve, rich in species [45]. Another investigation highlights the occurrence of *Cocconeis sawensis* Al-Handal et Riaux-Gobin, recently described for saline lakes in southern Iraq as an epiphyte in *Chara* sp. Linnaeus (1753: 1156). It was also recorded on an island in the South Pacific and epizoic on manatee in Florida Bay, USA [46]. The other species of the community were described, illustrated, and compiled with information on ecology and distribution in these coastal systems [47].

Studies about phytoplankton in subtropical coastal lagoon from south of Brazil showed that the structure and dynamic of the phytoplanktonic community were regulated by hydrological factors (inflow-outflow of continental and coastal waters in the system) as well as by meteorological conditions (wind and rainfall) and limnological variables (temperature and salinity) [18, 19, 33]. We have a set of factors that can act simultaneously while being difficult to recognize a main factor. Our question is to know if the structure and dynamic of the benthic diatoms in the Peixe Lagoon are related with these same factors. In order to answer this question, the study objectives were: (1) to know the composition of the diatoms community; (2) to verify the community structure and its spatial and temporal variation; and (3) to relate the variations of the community to environmental variables over an annual cycle.

In this chapter, firstly, we present information about the geographic, environmental, and climatic features where the Peixe Lagoon is situated. To be a case study, the methods are also included. Secondly, we describe the physical and chemical conditions of the lagoon and the benthic diatoms composition. Thirdly, we present and discuss the environmental variables related to the composition and spatial and temporal variation of the community attributes. Finally, we review the relationships of organisms occurring in plankton and sediment that should not be overlooked in studies in shallow coastal lagoons.

2. Study area

Peixe Lagoon is the only intermittent lagoon of the extreme south of Brazil and it is situated in the Lagoa do Peixe National Park (31°00′46″ S; 51°09′51″ W and 31°29′00″ S; 50°46′31″ W). This park is recognized by the Ramsar Convention as a Wetlands site, as well as an area of the UNESCO Atlantic Forest Biosphere Reserve, an Important Bird and Biodiversity Area (IBA) and a designated a site of international importance by the Western Hemisphere Shorebird Reserve Network (WHSRN). The coast is characterized by a microtidal regime, with a mean amplitude of 0.45 m [48].

Peixe Lagoon is a shallow, elongated system (35 km long and 1 km wide), parallel to the coastline (Figure 1) connected to the Atlantic Ocean through a single narrow channel (chocked lagoon) (Figure 2). The channel occlusion occurs due to deposition of sand caused by the predominance of the north and northeast winds [49, 50]. The connection with the ocean usually occurs during winter and spring, when the precipitation becomes more pronounced and the marshes and fields marginal to the lagoon are flooded. During these periods, an artificial opening of the channel is carried out by means of machines, since a natural opening only occurs sporadically [50]. The margins of the lagoon are covered by salt marshes vegetation dominated by *Paspalum vaginatum* Sw., Cotula coronopifolia L., Sporobolus montevidensis (Arechavaleta) P.M. Peterson & Saarela (= Spartina densiflora Brong), Hydrocotyle bonariensis Lam., Androtrichum trigynum (Spreng.) H. Pfeiff., Bacopa monnieri (L.) Wettst., and Juncus acutus L. [51]. The surface sediments at the bottom are essentially sandy. On the sites with greater depth of the lagoon, the sediments are thinner, with addition of silt and clay [49].

The system is located in subtropical climate where the rainfall is distributed throughout the year. In the period of studies, the highest rainfall (145.8 mm) occurred at the end of the fall (June 2011), decreasing in the following months and then increasing (131.2 mm) in early spring (October 2011). November had the lowest cumulative precipitation (23.0 mm). The average monthly temperature varied between 12.6 and 18°C in the autumn/winter seasons and between 18.3 and 24.4°C in spring/summer (**Figure 3**).

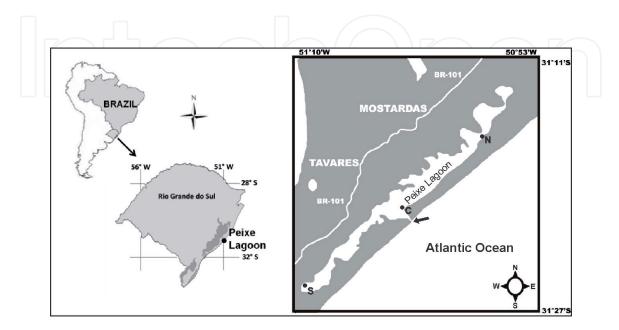


Figure 1.

Location of Peixe Lagoon area in the state of Rio Grande do Sul, southern Brazil, the sampling stations (North = N, Center = C, South = S) and the channel of connection with ocean (arrow).



Figure 2.

Aerial view of the Peixe Lagoon channel. Source: Lagoa do Peixe National Park (PNLP).

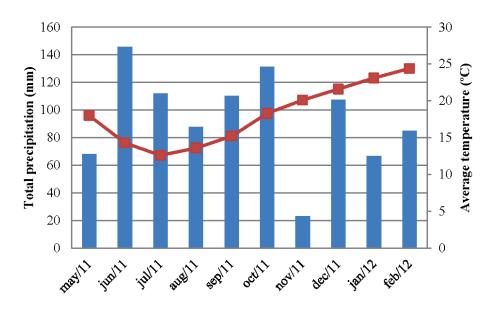


Figure 3.

Total precipitation (mm) and average monthly temperature (°C) recorded by the meteorological station of Mostardas/RS. Source: National Institute of Meteorology (INMET).

3. Methods

3.1 Sampling

The study was based on samples collected at three sampling stations in the lagoon. North (**Figure 4**) is close to a narrow channel that interconnects the northernmost sealed bodies with a central portion of the lagoon; Center (**Figure 5**) is next to the channel that connects to the ocean; and South (**Figure 6**) is at the south end of the lagoon. Sampling occurred in the four seasons, fall (June 2011), winter (August 2011), spring (November 2011), and summer (February 2012). During the first sampling, in the fall, the channel was closed. It was open days before winter sampling and remained open for the remainder of the sampling period. For the diatoms analysis, sediment samples were collected at depths of 2 cm with a spatula, at the lagoon margin, and packed in glass pods for transport to the laboratory.

3.2 Abiotic variable

Conductivity (mS.cm⁻¹), salinity, pH, water temperature (°C), dissolved oxygen—DO (mg.L⁻¹), and oxidation-reduction potential—ORP (mg.L⁻¹) were



Sampling station: North.



Figure 5. Sampling station: Center.



Figure 6. *Sampling station: South.*

measured with a HORIBA U52 probe. Depth and water transparency (cm) were measured with a Secchi disk. Precipitation, wind velocity, and wind direction data were obtained from the National Institute of Meteorology—INMET.

Laboratory analyses were performed as follows: total phosphorus—TP (mg.L⁻¹) by absorptiometry reduction of ascorbic acid, total nitrogen—TN (mg.L⁻¹), according to the Kjeldahl method (NBR 10560-1988, 13796-1997), and total silicate (mg.L⁻¹) with the silicomolybdate method [52]. The classification of salinity was based on the Venice System [53]. Trophic level was determined by the modified system of Vollenweider [54].

3.3 Diatom analysis

The sediment samples (1g) were dried in an oven and cleaned with potassium permanganate and hydrochloric acid according to the Simonsen technique [55]. For light microscopy (LM) analyzed, a Zeiss Axioplan Microscope (Carl Zeiss, Oberkochen, Germany) was used. The relative abundance of the taxa was carried out in slides seeking the minimum sample efficiency of 80% [56]. Species richness was estimated by the number of taxa present in the samples. The specific diversity was assessed using Shannon index (H') [57] and Evenness equitability (E). The analysis of variance (ANOVA) was applied to test the significance among the community attributes, since the data presented a normal distribution. The PAST® software was used for these analyzes. The relationship between biotic and abiotic variables with canonical correspondence analysis (CCA), PC-ORD® version 6.08 was used. For the construction of the biotic matrix, only species with a frequency equal to or >5% were considered in at least one sample unit and for the abiotic matrix, 10 environmental variables were included (Table 1). The data were transformed into log10(x + 1) in order to normalize the variances [58]. The Monte Carlo permutation test was carried out to verify the significance of the ordination axes.

		Fall			Winter			Spring	5		Summer				
	N	С	S	N	С	S	N	С	S	N	С	S			
Depth	65	35	25	90	20	40	8o	30	30	50	25	42			
Secchi	65	35	25	30	20	40	60	25	30	50	25	42			
Temp	15.3	17.2	17.6	17.0	17.6	17.4	28.3	26.1	29.3	29.8	32.1	31.9			
pН	7.9	7.9	8.4	7.6	8.o	7.7	7.9	8.2	8.8	8.1	8.3	8.2			
ORP	275	282	263	290	291	283	235	205	202	214	198	214			
DO	10.1	10.3	12.3	12.3	11.3	12.5	10.7	9.8	9.3	7.5	9.0	9.4			
Salin	31.2	28.3	10.8	4.5	17.0	1.3	7.1	36.2	10.9	34.9	36.0	29.5			
Cond	32.2	30.4	12.1	8.1	27.8	2.5	12.4	34.6	18.5	52.9	54.6	45.5			
Sil	3.8	1.7	1.1	17.0	18.0	17.4	1.0	2.7	6.6	7.8	0.0	9.1			
PΤ	0.03	0.07	0.08	0.15	0.12	0.06	0.08	0.13	0.07	0.07	0.05	0.06			
NT	0.61	0.16	0.43	o.87	0.68	0.90	0.01	0.09	0.02	0.55	0.65	0.60			

Table 1.

Physical and chemical variables analyzed in Peixe Lagoon in the four seasons, from June 2011 to February 2012, in the North (N), Center (C), and South (S) sampling stations. Depth (cm); Secchi = Secchi transparency (cm); temp = temperature (°C); ORP = oxide-reduction potential (mV); cond = conductivity (mS.cm⁻¹); DO = dissolved oxygen, salin = salinity (ppt); sil = silica (mg.L⁻¹); PT = total phosphorus (mg.L⁻¹); NT = total nitrogen (mg.L⁻¹).

4. Environmental conditions

The lagoon has a mean depth (<60 cm) and the Secchi disk depth generally coincides with the total depth. The water temperature varies between 15.3 and 17.6°C in the colder seasons (fall and winter) and from 26.1 to 32.1°C in the hottest seasons (spring and summer). The pH varies from 7.6 to 8.8 and the oxidation-reduction potential as well as the dissolved oxygen present similar trends, with higher values in the cold seasons and lower in the hot seasons (**Table 1**).

In relation to nutrients, total phosphorus varies between 0.03 and 0.08 mg.L⁻¹ in fall and summer at 0.12–0.15 mg.L⁻¹ in winter and spring, from eutrophic to hypereutrophic conditions. Total nitrogen presented higher values (0.68–0.90 mg.L⁻¹) in winter (mesoeutrophic conditions), with a decline in spring (0.01–0.09 mg.L⁻¹) and elevation in the summer (0.55–0.65 mg.L⁻¹), changing to mesotrophic conditions. Silica concentrations are higher in winter sampling (mean of 17.5 mg.L⁻¹). When the

channel of connection with ocean was open, the total nitrogen concentrations at all stations elevated as well as the total phosphorus in the north and center in winter. This may have been due to the water runoff from the land around the lagoon, used for livestock (**Table 1**).

The salinity demonstrated outstanding spatial and seasonal variations. These variations were mainly related to the meteorological conditions. Spatially, the salinity varies between 1.3% in the South station in the winter (oligohaline zone) and 36.2% in the Center during the summer (euhaline zone). The station with the highest salinity variation is the North (4.5% min./winter and 34.9% max./summer), followed by the South station (1.3% min./winter and 29.5% max./summer). The Center station maintained higher values of salinity in all the climatic seasons due to its proximity with the ocean. Seasonally, salinity has the highest values in summer and the lowest in winter. These low values can be attributed to the action of the wind, predominantly northeast, that propelled the waters from the Ruivo Lake which are less saline, to the Peixe Lagoon [33]. In the summer, the decrease of the precipitation and intensity of the wind causes the outstanding increase of the salinity. This dynamic was also observed in the system from 1991 to 1996 [50]. The South station of the lagoon presents less marine influence, therefore, lower salinity.

The variation of water levels of the lagoon is also strongly controlled by the winds regime, both intensity and direction, as well as precipitation. In the periods of predominant south wind (fall) and low precipitation (spring), the lowest levels of depth were observed. The Center is located next to the connection channel with the ocean and has a low average depth (30 cm). It is constantly saline (poly to euhaline zone). Due to the predominant northeasterly winds for most of the year, the water body of the lagoon is pushed to the west bank. The variation of the intensity of the winds can vary in the periods of day and night, causing great extensions of marginal sediment to be exposed and to be submerged again in a matter of hours [50].

The wind velocity during the period of studies had the lowest averages in the fall. It intensified in the following months of winter, with a peak in August

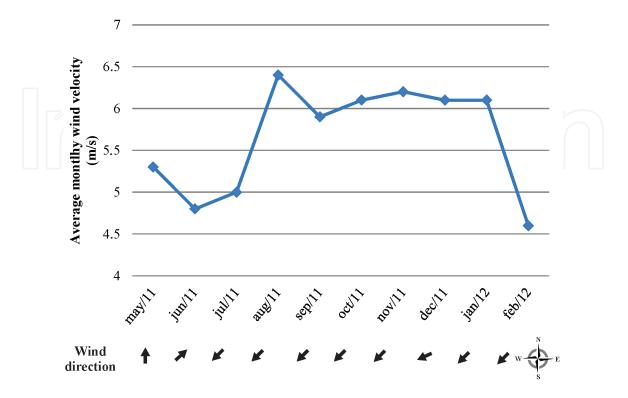


Figure 7.

Average monthly wind velocity (m/s) and predominant monthly wind direction (arrows indicate direction). Data recorded by the Meteorological Station of Mostardas/RS. Source: INMET.

(6.4 m/s). A sharp decrease occurred in February 2012 (4.6 m/s). The predominant direction of the wind in the fall was south and southwest, shifting northeast in July and in the following months (**Figure 7**).

5. Benthic diatoms composition

The diatoms community in Peixe Lagoon is composed by 62 taxa distributed in 30 genera composed largely of marine, brackish, and few freshwater species (**Table 2**). Similar results were recorded in an area adjacent to this study [28], where a total of 73 predominantly benthic and brackish taxa were found.

The genera with the greatest number of taxa are *Amphora* Ehrenberg ex Kützing, *Nitzschia* Hassall, and *Diploneis* Ehrenberg ex Cleve. The freshwater species that probably tolerate the wide variation of salinity are *Amphora ectorii*, *Cocconeis neodiminuta*, *C. euglypta*, *Chamaepinnularia truncate*, *Diploneis aestuari*, *D. didyma*, *Nitzschia palea*, *N. scalpelliformis*, *N. frustulum*, *N. vitrea* var. *salinarum*, and *Planothidium delicatulum*. More than 50% of taxa are cosmopolitan, and the remaining are restricted to a large extent to South America. An aspect to be highlighted is

TÁXON	N	FALL C	S	WINTER S N C S			SPRING N C S			SUMMER N C S		
TAXON Achnanthes curvirostrumBrun		-	X	-	-	-	- -	-	-	X	X	X
A. intermedia Kützing	-	x	-	-	-	-	x	-	-	-	-	Λ
	-									-		-
Imphora aff. baicalensisSkvortzow & Meyer	-	-	Х	-	-	-	X	-	-	-	-	-
1. ectorii Levkov	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х
1. maracaiboensis Levkov	Х	-	Х	-	-	-	Х	-	Х	Х	Х	-
1. <i>fusca</i> Schmidt	Х	-	-	Х	-	-	Х	-	-	Х	Х	-
Imphora sp.1	-	-	-	-	-	-	Х	-	-	-	-	-
Amphora sp. 2	-	-	-	-	-	-	-	-	-	-	Х	-
Amphora sp. 3	-	-	Х	-	-	-	-	-	-	-	-	-
Caloneis permagna (Bailey) Cleve	х	-	_	-	-	-	-	-	-	-	-	-
Caloneis sp.1	-	-	-	-	-			-	-	Х	-	-
Campylosira cymbelliformis(Schmidt) Grunow ex Van Heurck		Х	-		х					-	-	
Catenulaadhaerens (Mereschkowsky) Mereschkowsky	-	-	x	-	-	-	-	x	2	-	x	-
	-	x		-	-	-	-	-	-			-
Chamaepinnularia truncata (König)Witk.,Lange-Bert. & Metz.	-			-	-	-	-			Х		-
Cocconeis euglypta Ehrenberg	-	-	Х	-	-	Х	-	-	Х	-	-	-
C. sawensis Al-Handal et Riaux-Gobin	-	-	Х	-	-	Х	-	-	-	-	-	-
C.neodiminuta Krammer	-	-	-	-	-	-	-	Х	-	-	Х	-
Dimeregramma minus (Gregory) Ralfs	-	-	-	-	-	-	-	Х	-	-	Х	-
Diploneis aestuariHustedt	-	-	-	-	х	-	-	-	-	-	-	-
D. didyma (Ehrenberg) Cleve	Х	-	Х	Х	x	-	-	Х	-	-	Х	-
D. interrupta var. interrupta (Kützing) Clave	-	х	-	x	x	-	-	-		-	-	
D. litoralis var. clathrata (Østrup) Cleve		-		-	x	-	x		-	x	-	
D. moraus var. cramrata (Ostrup) Cleve D. smithii (Brébisson) Cleve	x	2	x	x		2		2	-		-	-
					-		X	-	-	-		-
Diploneis sp. 1	X	-	-	-		-	Х	-	-	-	-	-
Ehrenbergia granulosa	Х	-	-	Х	-	-	-	-	-	Х	-	-
Fallacia florinae (Moeller) Witkowski	-	Х	Х	-	-	х	Х	-	Х	-	-	Х
F. subforcipata (Hustedt) Mann	Х	-	-	Х	Х	-	Х	Х	-	Х	Х	-
F. tenera (Hustedt) Mann	-	-	-	Х	-	-	-	-	-	-	-	-
Fragilaria eichhornii Witkowski & Lange-Bertalot	-	-	Х	-	-	Х	-	-	Х	Х	-	Х
Halamphora coffeaeformis (Agardh) Kützing	-	_	-	-	-	-	-	-	Х	х	Х	Х
H. subholsatica (Krammer) Levkov	-		-	-	Х	-	_	_	-	-	-	-
H. turgida (Gregory) Levkov	x	x	x	-	-	x	-	-	-	x	-	-
Halamphora sp. 1	~	-	x	-	-		-	-	-	x	-	-
	-				-	-	-	-	-			
Luticola simplex Metzeltin, Lange-Bertalot & García-Rodríguez	Х	-	-	Х	-	-	-	-	-	-	-	-
Mastogloia braunii Grunow	-	-	-	-	-	-	-	-	-	-	-	Х
M. pusilla (Grunow) Cleve	-	-	Х	Х	-	-	-	-	-	-	-	-
Navicula flagellifera Hustedt	Х	-	-	-	-	-	-	-	-	Х	Х	-
N. juanitalinda Metzeltin, Lange-Bertalot & García-Rodríguez	-	Х	-	-	-	-	-	-	-	-	-	-
N. phylleptosomaformis Seddon et Witicowski	-	х	-	-	х	-	Х	-	-	х	-	-
Vitzschia dissipatoides Archibald	Х	-			-	-	-	_	-	-	-	-
N. frustulum (Kützing) Grunow	~	x	-	x	-	x	x	-	-	x	x	-
	-		-		-			-				-
N.palea (Kützing) Smith		-	-	-	-	-	Х	-	Х	-	-	-
N. scalpelliformis Grunow	Х	-	-	Х	-	-	-	-	-	-	-	-
N. spathulata Brébisson ex W.Smith	-	-	-	-	-	-	-	-	-	-	-	Х
V. vítrea var. salinarum (Grunow) Grunow	Х	-	-	-	-	-	-	-	-	-	-	-
Vitzschiasp. 1	Х	-	-	-	-	-	-	-	-	-	-	-
Opephora aff. mutabilis (Grunow) Sabbe & Vyverman	-	-	Х	-	-	-	Х	Х	-	Х	Х	-
0. pacifica (Grunow) Petit	-	_	-	-	-	-	-	x	-		X	-
Paralia sulcata (Ehrenberg) Cleve	х		-	Х				-	-	Х	-	
Petroneis marina (Ralfs) Mann	Λ	-	-		-	-	-	x	-	x	x	-
	-				-	-	-					-
Placoneis elegantula Metzeltin, Lange-Bertalot & García-Rodríguez	-	Х	Х	-	Х	-	-	-	Х	-	-	-
Planothidium delicatulum (Kützing) Round & Bukhtiyarova	-	-	Х	-	-	-	Х	Х	-	-	-	-
Pseudostaurosiropsis geocollegarumm (Witkowski) Morales	-	-	-	Х	-	-	Х	-	-	Х	-	-
Rhopalodia musculus (Kützing) Müller	-	-	Х	-	-	-	-	-	-	-	-	-
R. runrichiae Krammer	х	-	-	Х	-	-	-	-	-	-	-	-
Seminavis eulensteinii (Grunow) Danielidis, Ford & Kennett	-	-	х	-	-	-	-	-	-	-	-	-
Seminavis pusilla (Grunow) Cox & Reid	-	-	x	-	-	x	-	-	-	-	-	-
Seminuors pasma (Granow) - Cor & Reia Robinson (These do Donial dia - Romanna Anailti	-	x		-			-	-	-			-
S.strigosa (Hustedt) Danielidis e Economou-Amilli	-		-	-	Х	-	-	-	-	X	Х	-
Stauroforma sp.2	-	-	-	-	-	-	-	-	-	Х	-	-
Staurophora soodensis (Krasske) L.Bahls	-	-	Х	-	-	-	-	-	-	-	-	-
Terpsinoë americana (Bailey) Grunow	Х	-	-	Х	-	-	Х	-	-	Х	-	-
Tryblionella hungarica (Grunow) Frenguelli	_	-	-	-	-	х	-	-	-	_	-	-
									8			

Table 2.

Distribution of the diatoms at the North—N, Center—C, and South—S in the four seasons of the year in Peixe Lagoon, from June 2011 to February 2012.

the predominance of birraphid and monoraphid diatoms, which are organisms that have raphe. This structure is a selective characteristic of the epipelic species [59–63] because it promotes the movement of organisms in search for better light and humidity conditions, since it allows the secretion of polymeric substances produced by their cells.

6. Structure of the community versus environmental conditions

6.1 Spatial and temporal variation

The community attributes (richness, diversity, and evenness) showed a decreasing spatial gradient from the North to the South stations. The specific diversity ranged between 2.3 bits/ind. at North and 0.4 bits/ind. at South and the evenness varied between 74 and 20% at North and South stations.

Seasonally, in fall, without connection with the ocean, the community attributes presented the highest values. After the channel opening, the richness was similar in winter and spring, rising in the summer. The values of evenness and diversity increased from winter (0.6–1.7 bits/ind.) to summer (0.8–2.3 bits/ind.) (**Figure 8**). However, these attributes did not differ significantly between the seasons and the station sampling.

6.2 Diatoms composition related to environmental variables

The composition of the diatoms and the physical and chemical variables of the water in the canonical correspondence analysis (**Figure 9**) of the abundant species (25 species with more than 5% abundance) can better demonstrate the community dynamics in the system.

The sampling units of the South station are grouped on the negative side of axis 1. They were related to the lower values of conductivity and salinity. The species associated with this axis were *Cocconeis sawensis*, *Fragilaria eichhornii*, *Cocconeis euglypta*, *Fallacia florinae*, and *Halamphora coffeaeformis*. In this axis, it is also possible to observe the separation of the sampling units from the North, mainly due to the difference in temperature between hot and cold seasons, in fall and winter months. The related species were *Nitzschia scalpelliformis*, *Luticola simplex*, *Ehrenbergia granulosa*, *Rhopalodia runrichiae*, *Diploneis smithii*, and *D. didyma*. The sampling units of the Center station are grouped on the positive side of axis 2, where higher values of salinity and temperature were observed in the hotter seasons, as well as the lower values of silica and ORP. The species *Opephora pacifica*, *Catenula adhaerens*, and *Opephora* aff. *mutabilis* were related to this axis (**Figure 10**).

During the study, we observed periods with higher and lower marine influence, due to the opening of the channel. In fall, the only season in which the channel was closed, the composition of diatom species was distinct in the north and south of the lagoon. The south and south-west quadrant wind might also have been an influence factor for the distinction of community composition.

After the channel opening, it is possible to observe the difference in the composition of the community at the southern portion in relation to the north and center portions of the lagoon. The species highlighted in the south (*Cocconeis sawensis*, *C. euglypta*, *Fallacia florinae*, and *Halamphora coffeaeformis*) are found in brackish and marine waters, with the exception of *C. euglypta*, a characteristic species of freshwater, but it supports high conductivity water [64]. So, the marine influence appeared as one of the main factors affecting spatial diatom composition and spatial distribution in the lagoon.

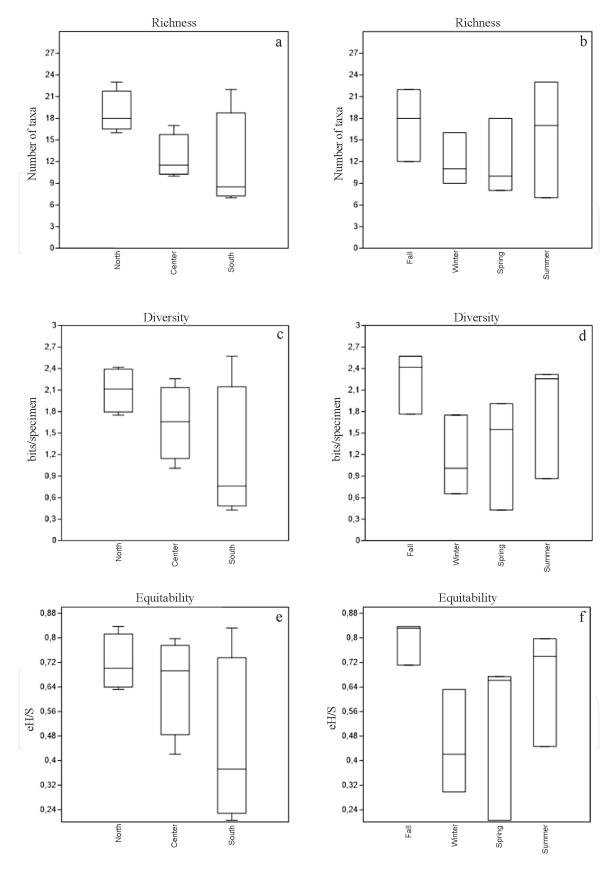
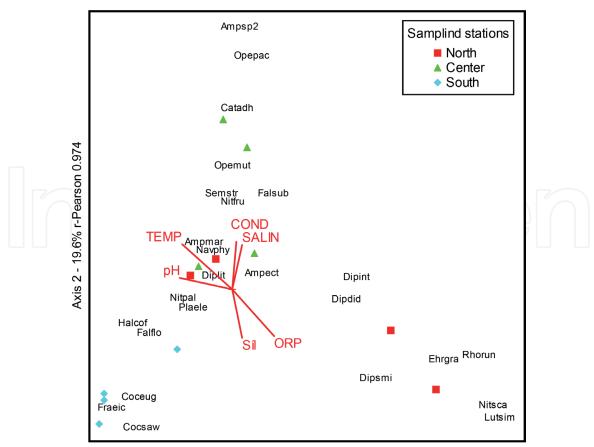


Figure 8.

(a-f) Distribution of community attributes related to sampling stations and the seasons of the year in Peixe Lagoon from June 2011 to February 2012.

However, the salinity cannot be considered as the only driving force that determines the composition of diatom species in environments with marine influence in subtropical and temperate regions. Temperature is also considered a very important environmental factor [11]. In Peixe Lagoon, the temperature difference between hot seasons (spring and summer) and cold seasons (fall and winter) also differentiated



Axis 1 - 26.5% r-Pearson 1.000

Figure 9.

Canonical correspondence analysis (CCA) of the abundant species in the sampling stations and seasons fall, winter, spring, summer in Peixe Lagoon. For legends of the variables, see **Table 2**. Amphora ectorii (Ampect), A. maracaiboensis (Ampmar), Amphora sp.2 (Ampsp2), Catenula adhaerens (Catadh), Cocconeis euglypta (Coceug), C. sawensis (Cocsaw), Diploneis didyma (Dipdid), D. interrupta (Dipint), D. litoralis (Diplit), D. smithii (Dipsmi), Ehrenbergia granulosa (Ehrgra), Fallacia florinae (Falflo), F. subforcipata (Falsub), Halamphora coffeaeformis (Halcof), Luticula simplex (Lutsim), Navicula phylleptosomaformis (Navphy), Nitzschia frustulum (Nitfru), N. palea (Nitpal), N. scalpelliformis (Nitsca), Opephora aff. mutabilis (Opemut), O. pacifica (Opepac), Placoneis elegantula (Plaele), Rhopalodia runrichiae (Rhorun), Seminavis strigosa (Semstr), Fragilaria eichhornii (Fraeic).

the composition of the species. *Diploneis interrupta*, *D. didyma*, *D. smithii*, *N. scalpelliformis*, and *Luticola simplex* were related to the colder seasons. This was also observed for species of *Diploneis* in the sediment of sublittoral zone of the Gulf of Trieste [65]. *Catenula adhaerens*, *Nitzschia frustulum*, *Opephora aff. mutabilis*, *O. pacifica*, and *Seminavis strigosa* were related to the sampling units with higher temperatures and salinities.

6.3 Diatoms related to sediment

Another important factor regarding diatom distribution is the sediment characteristic [66, 67]. In the Center and South stations, the surface of the Peixe Lagoon is essentially covered by sandy sediments, in which we find *Campylosira cymbelliformis*, *Catenula adhaerens*, *Dimeregramma minus*, and *Staurophora soodensis* species usually associated with sand grains.

In deeper sites of the lagoon, such as near the North, the sediments are thinner, with addition of silt and clay [49]. In this station, where muddy sand is present, we observed more clearly the seasonal variation of the diatom community. This site also showed highest diversity (1.7–2.4 bits/ind.) and richness (16–26 táxons) and the presence of more exclusive epipelic species; among these are the following: *Caloneis permagna*, *Luticola simplex*, *Nitzschia dissipatoides*, *N. scapelliformis*,

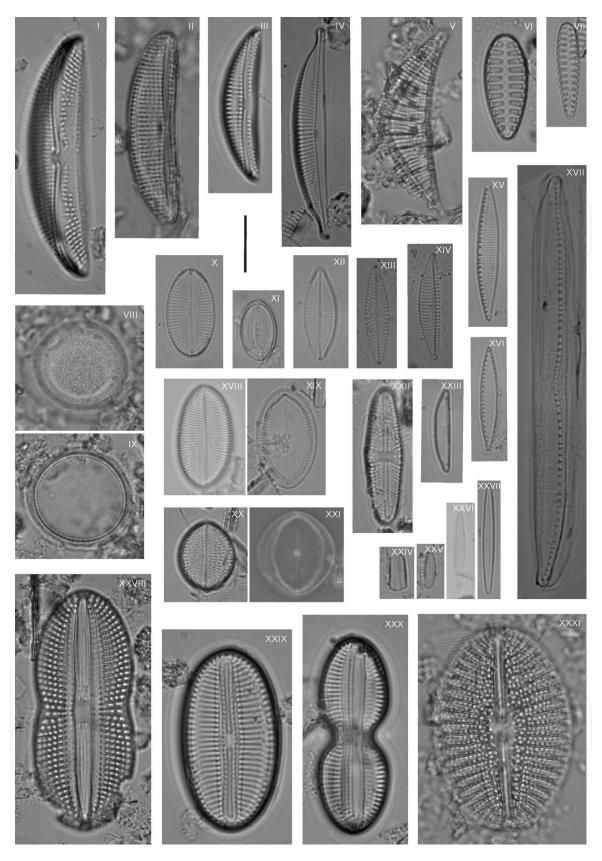


Figure 10.

I. A. maracaiboensis. II. Amphora sp. 2. III. A. ectorii. IV. Halamphora coffeaeformis. V. Rhopalodia runrichiae. VI. Opephora pacifica. VII. O. aff. mutabilis. VIII,IX. Erhembergia granulosa. X. Fallacia subforcipata. XI. F. florinae. XII. Placoneis elegantula. XIII. Navicula phylleptosomaformis. XIV. Seminavis strigosa. XV. Nitzschia frustulum. XVI. N. palea. XVII. N. scalpelliformis. XVIII,XIX. Cocconeis euglypta. XX,XXI. C. sawensis. XXII. Luticola simplex. XXIII. Catenula adhaerens. XXIV-XXVII. Fragilaria eichhornii. XXVIII. Diploneis didyma. XXIX. D. litoralis var. clathrata. XXX. D. interrupta. XXXI. D. smithii. Scale bar = 10 µm.

N. vitrea var. *salinarum*, *Rhopalodia runrichiae*, and *Terpsinöe americana* (**Table 2**). In agreement with other studies, the epipsammic fraction appeared to be much more stable than epipelic assemblage [9, 67].

7. Sediment and water interaction

We expected to find planktonic forms in the sediment due to the low depth of the lagoon and because of the fact that the sediment usually integrates planktonic and periphytic taxa [3]. The absence of planktonic forms could be explained by the hydrographic processes that tend to transport nonliving, unattached forms out of the system, similar to estuaries [67]. Furthermore, the location of the sampling stations, since the material was collected on the lagoon margin, was outside the water surface. However, it is known that in periods with decreasing wind intensity, water tend to return flooding areas that had been exposed [50]. A few planktonic species in the sediment were also recorded in a study of microphytobenthos in the Gulf of Trieste, Europe, although the collections were made in submerged sediment [65].

Comparing with an earlier study about phytoplankton at Peixe Lagoon, with sampling performed during the same period, *Asterionellopsis glacialis* (Castracane) Round, *Chaetoceros gracillis* Pantocsek, and *Skeletonema potamos* (Weber) Hasle were found in abundance in plankton samples. In this study, however, these species were not found in the sediment; whereas *Diploneis didyma*, a highlighted species found in benthos, was also present in the plankton. The species *Cocconeis sawensis* was recorded at the southern benthos of the lagoon, and it was also observed in the plankton and epiphyton in association with the macroalgae *Cladophora* sp. in the fall and winter seasons [33, 46]. This suggests that in shallow environments, the plankton receives a greater contribution of benthic species than the opposite. Similar results were found in shallow estuarine zone of Patos Lagoon [19].

Estuary and shallow coastal waters develop the process of resuspension whereby sediment particles with diatoms enter the water column. Examination of diatoms in the water revealed that 75% of frustules belonged to pennate forms and we concluded that flooding tides were responsible for a net transport of epipelic diatoms from the mudflat to a salt marsh. The resuspension of the diatoms can be the source of the chl *a* peak in the plankton [68]. So, this organism may greatly augment the primary production in water [69, 70]. Other investigations have showed large number of benthic diatoms in the water column [71, 72]. The wind, flooding tides, and tidal inducing waves and currents are the causes of this process.

8. Conclusions

In the Peixe Lagoon, the benthic diatoms were present in high diversity. Among the attributes of the community, the taxonomic composition best responded to the environmental variables. The quantitative attributes did not show significant relationships. The connection with the ocean, salinity, rainfall, wind action, and temperature were strongly related to the spatial and seasonal variation of the composition of the diatom community in this lagoon system. These organisms substantiate their use as indicators of environmental variations, mainly regarding salinity and temperature in subtropical coastal systems.

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References

[1] MacIntyre HL, Geider RJ, Miller DC.
Microphytobenthos: The ecological role of the "Secret Garden" of unvegetated, shallow-water marine habitats.
I. Distribution abundance and primary production. Estuaries. 1996;19:186-201

[2] Facca C, Sfriso A, Socal G. Temporal and spatial distribution of diatoms in the surface sediments of the Venice Lagoon. Botanica Marina. 2002;**45**:170-183

[3] Round F, Crawford R, Mann D. The Diatoms. Biology and Morphology of the Genera. 1st ed. Cambridge: Cambridge University Press; 1990. p. 747

[4] Snoeijs P, Weckström K. Diatoms environmental change in large brackishwater ecosystems. In: Stoermer EF, Smol JP, editors. The Diatoms: Applications for the Environmental and Earth Sciences. 2nd ed. United Kingdom: Cambridge University Press; 2010. pp. 287-308

[5] Admiraal W. The ecology of estuarine sediment-inhabiting diatoms. In: Round FE, Chapman D J, editors. Progress in Phycological Research. Bristol: Biopress. 1984;**3**:269-322

[6] Cahoon LB, Cooke JE. Benthic microalgal production in Onslow Bay, North Carolina, USA. Marine Ecology Progress Series. 1992;**84**:185-196

[7] Cahoon LB, Beretich GR, Thomas CJ, McDonald AM. Benthic microalgal production at Stellwagen Bank, Massachusetts Bay, USA. Marine Ecology Progress Series. 1993;**102**:179-185

[8] Trobajo R, Sullivan MJ. Applied diatom studies in estuaries and shallow coastal environments. In: Stoermer EF, Smol JP, editors. The Diatoms: Applications for the Environmental and Earth Sciences. 2nd ed. United Kingdom: Cambridge University Press; 2010. pp. 309-323

[9] Ribeiro LLCS. Intertidal benthic diatoms of the Tagus estuary: Taxonomic composition and spatialtemporal variation [thesis]. Lisboa: Universidade de Lisboa; 2010

[10] Hassan GS, Espinosa MA, Isla FI. Modern diatom assemblages in surface sediments from estuarine systems in the southeastern Buenos Aires Province, Argentina. Journal of Paleolimnology. 2006;**35**:39-53. DOI: 10.1007/ s10933-005-6444-8

[11] Espinosa MA, Isla FI. Modern diatom assemblages in surface sediments from meso-macrotidal estuaries of Patagonia, Argentina. Pan-American Journal of Aquatic Sciences. 2015;**10**:20-43

[12] García-Rodrígues F, Metzeltin D, Sprechmann P, Trettin R, Stams G, Beltrán-Morales LF. Upper and Holocene paleosalinity and trophic state changes in relation to sea level variation in Rocha Lagoon, southern Uruguay. Journal of Paleolimnology. 2004;**32**:117-135

[13] García-Rodrígues F, Sprechmann P, Metzeltin D, Scafati L, Melendi DL, Volkheimer W, et al. Holocene trophic state changes in relation to sea level variation in Lake Blanca, S E Uruguay. Journal of Paleolimnology. 2004;**31**: 99-115. DOI: 10.1023/B:JOPL.00000132 81.31891.8e

[14] García-Rodriguez F. Inferring paleosalinity trends using the chrysophyte cyst to diatom ration in coastal shallow temperate/subtropical lagoons influenced by the sea level changes. Journal of Paleolimnology.
2006;36:165-1173. DOI: 10.1007/ s10933-006-0011-9 [15] del Puerto L, García-Rodrígues
F, Inda H, Bracco R, Castineira C, Adams JB. Paleolimnological evidence of Holocene climate changes in Lake
Blanca, southern Uruguay. Journal of
Paleolimnology. 2006;**36**:151-163. DOI: 10.1007/s10933-006-0012-8

[16] García-Rodrigues F, Stutz S, Inda H, Puerto Ld, Bracco R, Panario D. A multiprox approach to inferring Holocene paleobotanical changes linked to sea-level variation, paleosalinity levels, and shallow lake alternative states in Negra Lagoon, SE Uruguay. Hydrobiology. 2010;**646**:5-20. DOI: 10.1007/s10750-010-0184-0

[17] Torgan L, Garcia-Baptista M,
Odebrecht C, Möller O. Distribuição
vertical do fitoplâncton na Laguna dos
Patos, Rio Grande do Sul, Brasil (verão,
1986). Acta Limnologica Brasiliensia.
1995;7:67-77

[18] Torgan L. Estrutura e dinâmica da comunidade fitoplanctônica na Laguna dos Patos, Rio Grande do Sul, Brasil, em um ciclo anual [thesis]. São Paulo: Universidade Federal de São Carlos; 1997

[19] Torgan L, Tundisi J, Niencheski L. Seasonal variation of planktonic diatoms in Patos Lagoon, Southern Brazil. In: Jacob J editor. Proceedings of the 15th Diatom Symposium. Tuggell: A.R.G. Gantner Verlag; 2002. pp. 459-470

[20] Torgan L, Odebrecht C, Niencheski L. Variação espacial da estrutura de tamanho do fitoplâncton na Laguna dos Patos, Sul do Brasil. Atlantica. 2000;**22**:95-111

[21] Torgan L, Raupp S. Morfologia externa de *Melosira moniliformis*(O.F. Müller) C. Agardh var. *moniliformis* (Bacillariophyta) do estuário da laguna dos Patos, Rio Grande do sul, Brasil. Iheringia, Série Botânica.
2001;56:185-196 [22] Torgan L, Pillar V, Niencheski F. Phytoplankton associations of a coastal lagoon in south of Brazil. Journal of Coastal Research. 2006;**39**:1149-1151

[23] Torgan L, Becker V, Santos C. *Skeletonema potamos* (Bacillariophyta) in Patos Lagoon, southern Brazil: Taxonomy and distribution. Revista Peruana de Biología. 2009;**16**(1):93-96

[24] Odebrecht C, Bergesch M, Medeanic S, Abreu P. A comunidade de microalgas. In: Seeliger U, Odebrecht C, editors. O estuário da Lagoa dos Patos: um século de transformações. Editora da FURG: Rio Grande; 2010. pp. 49-63

[25] Rosa Z, Callegaro V. Diatomáceas da Lagoa Tramandaí e da Lagoa do Armazém, Rio Grande do Sul, Brasil: I— Gênero *Navicula* Bory. Iheringia, Série Botânica. 1988;**37**:17-32

[26] Rosa Z, Miranda-Kiesslich A. O gênero *Pediastrum* Meyen (Chlorococcales-Hydrodictyaceae) no sistema lagunar da Região Litoral do Rio Grande do Sul, Brasil. Iheringia, Série Botânica. 1988;**38**:149-169

[27] Werner V. Cianofíceas planctônicas da Lagoa de Tramandaí e da Lagoa do Armazém, Rio Grande do Sul, Brasil. Iheringia, Série Botânica. 1988;**37**:33-70

[28] Rosa Z, Werner V. Diatomáceas da Lagoa Tramandaí e da Lagoa do Armazém, Rio Grande do Sul, Brasil: II—Gêneros *Gyrosigma* Hassal, *Pleurosigma* W. Smith e *Mastogloia* Thwaites. Iheringia, Série Botânica. 1993;**43**:67-87

[29] Rosa Z, Werner V, Dacroce L. Diatomáceas da Lagoa de Tramandaí e da Lagoa Armazém, Rio Grande do Sul, Brasil: III—Ordem Centrales. Iheringia, Série Botânica. 1994;**45**:29-55

[30] Werner V. Cyanophyceae/ Cyanobacteria no sistema de lagoas e lagunas da planície costeira do Estado do

Rio Grande do Sul, Brasil [thesis]. São Paulo: Universidade Estadual Paulista; 2002

[31] Werner V, Sant'anna C. A new species of *Aphanothece* (*Cyanophyceae*, *Chroococcales*) from a shallow coastal lagoon, south Brazil. Nova Hedwigia. 2000;**70**:113-125

[32] Werner V, Sant'anna C. Occurrence of the rare genus *Microcystis* P. Richter (*Chroococcales*, *Cyanobacteria*) in a coastal lagoon from southern Brazil. Revista Brasileira de Botânica. 2006;**29**:183-186. DOI: 10.1127/nova. hedwigia/70/2000/113

[33] Donadel L, Cardoso L, Torgan L. Plankton community dynamics in a subtropical lagoonal system and related factors. Anais da Academia Brasileira de Ciências. 2016;**88**:249-267. DOI: 10.1590/0001-3765201520150022

[34] Donadel L, Torgan L. *Falcula hyalina* (*Fragilariaceae*, *Bacillariophyta*) from a coastal lagoon, southern Brazil: An additional approach on its morphology. Phytotaxa. 2016;**243**:185-189. DOI: 10.11646/phytotaxa.243.2.10

[35] Bergesch M, Odebrecht C, Abreu P. Microalgas do estuário da Lagoa dos Patos: Interação entre o sedimento e a coluna de água. Oecologia Brasiliensis. 1995;**1**:273-289

[36] Garcia M. The transfer of *Fragilaria obtusa* Hustedt to the genus *Staurosira* Ehrenberg (*Bacillariophyceae*). Phycological Research. 2006;**54**:87-93. DOI: 10.1111/j.1440-1835.2006.00412.x

[37] Garcia M, Talgatti D. The diatom *Anorthoneis dulcis* Hein from southern Brazil: Morphology and ecology. Research Letters in Ecology. 2008:1-5. DOI: 10.1155/2008/140245

[38] Torgan L, Donadel L, Silva J. A transferência de *Navicula sovereignae* Hustedt para o gênero *Placoneis* Mereschkowsky (*Bacillariophyta*). Iheringia, Série Botânica. 2010;**65**:107-114

[39] Silva J, Torgan L, Cardoso L. Diatomáceas (*Bacillariophyceae*) em marismas no sul do Brasil. Acta Botânica Brasílica. 2010;**24**:935-947

[40] Talgatti D. Diatomáceas (Bacillariophyta) em marismas do sul do Brasil: Estudo da comunidade bentônica [thesis]. Porto Alegre: Universidade Federal do Rio Grande do Sul; 2014

[41] Talgatti D, Bertolli L, Torgan L. *Seminavis recta* comb. nov. et stat. nov.: Morphology and distribution in salt marshes from southern Brazil. Fottea. 2014;**14**:141-148. DOI: 10.5507/ fot.2014.011

[42] Talgatti D, Sar E, Torgan L. *Haslea sigma (Naviculaceae, Bacillariophyta)* a new sigmoid benthic species from salt marshes of southern Brazil. Phytotaxa. 2014;**177**:231-238. DOI: 10.11646/ phytotaxa.177.4.4

[43] Talgatti D, Wetzel C, Morales E, Ector L, Torgan L. Transfer of *Fragilaria atomus* Hust. To the genus *Stauroforma* (*Bacillariophyta*) based on observation of type and newly collected material. Phytotaxa. 2014;**158**:43-56. DOI: 10.11646/phytotaxa.158.1.3

[44] Santos C. Assembléias de diatomáceas em sedimentos Holocênicos no extremo sul do Brasil: Reconstruções paleoambientais [dissertation]. Porto Alegre: Universidade Federal do Rio Grande do Sul; 2011

[45] Pacheco C, Bertolli L, Donaldel L, Torgan L. O gênero *Diploneis* Ehrenberg ex Cleve (*Bacillariophyceae*) em marismas do sul do Brasil. Iheringia, Série Botânica. 2016;**71**:331-355

[46] Donadel L, Torgan L, Al-Handal A. Additional morphological features of the epiphytic diatom *Cocconeis*

sawensis Al-Handal & Riaux-Gobin (*Cocconeidaceae*, *Bacillariophyta*) from a coastal lagoon, southern Brazil. Phytotaxa. 2018;**371**:217-229. DOI: 10.11646/phytotaxa.371.3.5

[47] Donadel L, Torgan L. Benthic diatoms from lagoon system in the National Park Lagoa do Peixe southern Brazil. Biota Neotropica. (forthcoming)

[48] Toldo EE Jr, Dillenburg SR, Almeida LESB, Tabajara LL, Martins RR, Cunha LOBP. Parâmetros morfodinâmicos da praia de Imbé, RS. Pesquisas em Geociencias. 1993;**20**:27-32

[49] Arejano T. Geologia e Evolução Holocênica do Sistema Lagunar da "Lagoa do Peixe", Litoral Médio do Rio Grande do Sul, Brasil [dissertation]. Porto Alegre: Universidade Federal do Rio Grande do Sul; 2006

[50] Knak R. Relatório Técnico Final. In: Knak R. (coord.) Projeto caracterização ambiental do Parque Nacional da Lagoa do Peixe. Rio Grande: Fundação Universidade de Rio Grande; 1998. p. 1-327

[51] Costa CS, Tagliani PR. Cobertura vegetal e uso preponderante do espaço. In: Tagliani PRA, editor. Ecologia da paisagem da restinga da Lagoa dos Patos: Uma contribuição para o manejo e conservação da Reserva da Biosfera. Rio Grande: Editora da Furg; 2011. p. 184

[52] American Public Health Association (APHA). Standard Methods for
Examination of Water and Wastewater.
20nd ed. Washington: American
Public Health Association and Water
Environment Federation; 1998. p. 964

[53] Anonymous. Final resolution of the symposium on the classification of brackish waters. Archivio di Oceanografia e Limnologia.1959;11(Supplement):243-245 [54] Wetzel RG. Limnologia. 2nd ed. Lisboa: Fundação Calouste Gulbenkian; 1993. p. 919

[55] Simonsen R. The diatom plankton of the Indian Ocean Expedition of R/V "Meteor" 1964-1965. Meteor Forschungen Ergebniss. 1974;**19**(Série D):1-107

[56] Pappas J, Stoermer E. Quantitative method for determining a representative algal sample count. Journal of Phycology. 1996;**32**:693-696

[57] Shannon CE, Weaver W. TheMathematical Theory of Communication.Urbana: The University of Illinois Press;1949. p. 125

[58] Ter Braak C. Canonical correspondence analysis: A new eigenvector technique for multivariate direct gradient analysis. Ecology. 1986;**67**:1167-1179

[59] Lund J. Observations on soil algae.I. The ecology size and taxonomy of British soil diatoms. New Phytologist.1945;44:56-110

[60] Stoermer EF. Notes on Iowa diatoms. II. Species distribution in a subaerial habitat. Proceedings of the Iowa Academy of Science. 1962;**69**:87-96

[61] Hay S, Maitland T, Paterson D. The speed of diatom migration through natural and artificial substrata. Diatom Research. 1993;**8**:371-384

[62] Underwood G, Paterson D. The importance of extracellular carbohydrate production by marine epipelic diatoms. Advances in Botanical Research. 2003;**40**:183-240. DOI: 10.1016/S0065-2296(05)40005-1

[63] Witkowski A, Brehm U, Palińska K, Rhiel E. Swarm-like migratory behaviour in the laboratory of a pennate diatom isolated from

North Sea sediments. Diatom Research. 2012;**27**:95-100. DOI: 10.1080/0269249X.2012.690204

[64] Romero OE, Jahn R. Typification of *Cocconeis lineata* and *Cocconeis euglyta* (Bacillariophyta). Diatom Research. 2013;**28**:175-184. DOI: 10.1080/0269249X.2013.770801

[65] Cibic T, Blasutto O, Falconi C, Umani S. Microphytobenthic biomass, species composition and nutrient availability in sublittoral sediments of the Gulf of Triste (northern Adriatic Sea). Estuarine, Coastal and Shelf Science. 2007;75:50-62. DOI: 10.1016/j. ecss.2007.01.020

[66] Ampsoker MC, McIntire CD. Distribution of intertidal diatoms associated with sediments in Yaquina estuary, Oregon. Journal of Phycology. 1978;**14**:387-395. DOI: 10.1111/j.1529-8817.1978.tb02457.x

[67] Sabbe K. Short-term fluctuation in benthic diatom numbers on an intertidal sandflat in the Westerschelde estuary (Zeeland, the Netherland). Hidrobiologia. 1993;**269/270**:275-284. DOI: 10.1007/BF00028026

[68] Baillie PW, Welsh BL. The effect of tidal resuspension on the distribution of intertidal epipelic algae in an estuary. Estuarine and Coastal Marine Science. 1980;**10**:165-180. DOI: 10.1016/ S0302-3524(80)80056-9

[69] Riaux-Gobin C. Phytoplankton, tripton et microphytobenthos: échanges au cours de la marée, dans un estuaire du Nord-Finistére. Cahiers du Biologie Marine. 1987;**28**:159-185

[70] Shaffer GP, Sullivan MJ. Water column productivity attributable to displaced benthic diatoms in well-mixed shallow estuaries. Journal of Phycology. 1988;**24**:132-140. DOI: 10.1111/j.1529-8817.1988.tb04226.x [71] Gallagher JB. The significance of the surface film in salt marsh plankton metabolism. Limnology and Oceanography. 1975;**29**:120-123. DOI: 10.4319/lo.1975.20.1.0120

[72] Lukatelich RJ, McComb AJ.
Distribution and abundance of benthic microalgae in a shallow southwestern Australian estuarine system. Marine Ecology Progress Series.
1986;27:287-297

