

# Microphytoplankton assemblages in shallow waters at Admiralty Bay (King George Island, Antarctica) during the summer 2002–2003

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Received: 7 December 2006 / Revised: 8 May 2007 / Accepted: 8 May 2007 / Published online: 31 May 2007  
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**Abstract** Microphytoplankton populations were studied in shallow coastal water (<60 m) near the Brazilian Antarctic Station Comandante Ferraz (EACF) and three reference areas in Admiralty Bay in early and late summer (2002–2003). Phytoplankton was diverse (113 taxa), but not abundant ( $10^3$  cells  $l^{-1}$ ). The highest abundances ( $>10^4$  cells  $l^{-1}$ ) were caused by pennate benthic diatoms (*Fragilaria striatula* Lyngbye) that occurred mainly in early summer, associated with the presence of ice. In late summer, when the water temperature ( $-0.4$  to  $1.5^\circ\text{C}$ ), salinity (34 to 35), and phosphate ( $2.6$  to  $4.5$   $\mu\text{mol } l^{-1}$ ) were highest and the dissolved oxygen was lowest ( $6.4$  to  $2.9$   $\text{ml } l^{-1}$ ), centric diatoms (*Thalassiosira* spp.) were more abundant, suggesting an influence of oceanic waters. Phytoplankton abundance ( $\leq 10^2$  cells  $l^{-1}$ ) and chlorophyll *a* concentrations ( $0.22$   $\mu\text{g } l^{-1}$ ) were lowest close to EACF. Pennate diatoms were dominant close to shore and in surface waters elsewhere, probably because of ice melting or sediment resuspension caused by water mixing.

**Keywords** Microphytoplankton · Nearshore waters · Temporal and spatial variation · Antarctica

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## Introduction

Studies of nearshore antarctic phytoplankton are relatively scarce because of its apparent low contribution to the total primary production of the Southern Ocean (El-Sayed and Fryxell 1993). High phytoplankton abundance and biomass in coastal areas are usually related to resuspension of benthic microalgae (Brandini and Rebello 1994).

The first phytoplankton sample from Admiralty Bay was collected during the *Pourquoi-Pas?* Expedition in 1909 and the research on phytoplankton began in the 1970s with the establishment of research stations within the bay (Ligowski and Kopeczynska 1993). After that, other studies related to the Admiralty Bay phytoplankton have been made mainly during the 1980s by Polish scientists (Rakusa-Suszczewski 1980, Rakusa-Suszczewski et al. 1993), as well as during the 1990s by Polish and Brazilian scientists (Brandini and Rebello 1994).

Most anthropogenic activities in the bay have been related to four research stations: Henryk Arctowski (Poland) at Thomaz Point, the Comandante Ferraz Station—EACF (Brazil) at Keller Peninsula, Machu Picchu (Peru) at Crepin Point, and Copacabana (USA) at Llano Point (ATCM XXVIII).

The objective of this work was to evaluate the temporal and spatial distribution of microphytoplankton (cells  $> 20$   $\mu\text{m}$ ) during the austral summer of 2002–03, through the study of abundance and specific composition associated with hydrological features near EACF and three other reference areas at Admiralty Bay.

## Study area

Admiralty Bay ( $62^\circ 03' - 12'S$ ,  $58^\circ 18' - 38'W$ ) is located at King George Island, (area  $122 \text{ km}^2$ ) and has an estimated

water volume of 24 km<sup>3</sup>. It is a deep fjord-like embayment with 500 m maximum depth at its center (Rakusa-Suszczewski et al. 1993).

The water from the bay meets the oceanic deep waters from Bellingshausen and Weddell Seas at its southern opening, which connects to the Bransfield Strait (Rakusa-Suszczewski 1980; Lipski 1987).

According to Rakusa-Suszczewski (1980), the freshwater that flows to the sea at Admiralty Bay originates from the glaciers, covering more than 90% of King George Island. The melt water carries nutrients and organic matter to the sea, and lowers the seawater salinity. High values of sediment content and total phosphorus were found near the main terrestrial input areas. Strong winds and gusts are common in the coastal areas of the Antarctic Peninsula and South Shetland Islands. These winds, together with the tides and freshwater drainage, play an important role in the transport of organic matter, nutrients and trace metals to the sea, which influences primary production.

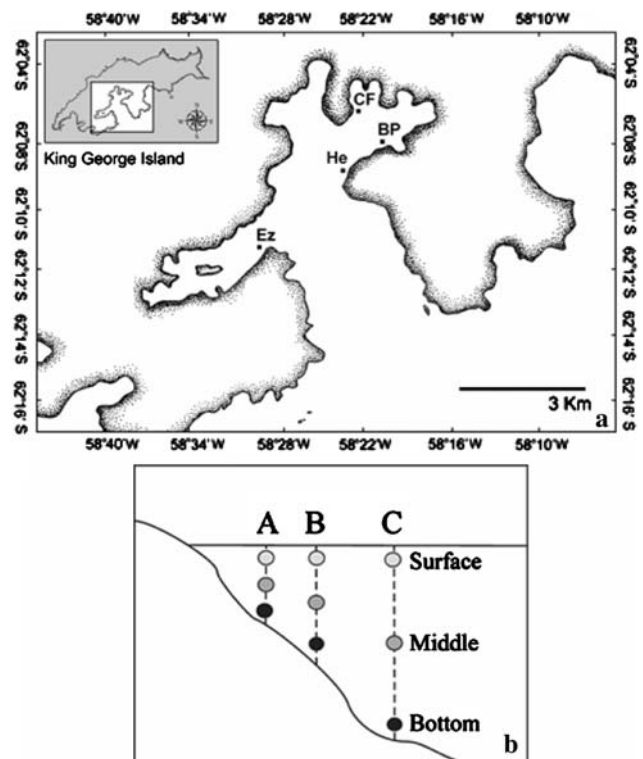
## Materials and methods

Admiralty Bay was surveyed during the early and late austral summer of 2002–03 (25 November–10 December, 2002 and 3–7 February, 2003). Samples were taken near EACF and three other reference areas distant from any anthropogenic activity using a Beyond BACI approach (Underwood 1994; Stark et al. 2003). The three other reference areas were Hennequin Point, Botany Point and an area about 300 m away from Thomaz Point within the Ezcurra Inlet (Fig. 1a). Three sampling stations were established in each area following the bathymetries of 20 m (St. A), 30 m (St. B) and 60 m (St. C) (Fig. 1b). At each station, samples were collected at surface, mid-water and at 1 m above the bottom.

### Physical and chemical analyses

The temperature was obtained using protected reversion thermometers, and the salinity was measured using a Beckman inductive salinometer. Dissolved oxygen was determined by the Winkler method using a Mettler DL 21 automatic titrator, following the Grasshoff et al. (1983) protocol.

The nutrient samples were collected using Go-flo bottles (General Oceanic®), filtered in Whatman GF/F membrane and stored in polyethylene flasks pre-washed with HCl 1:1, rinsed with distilled water. Then, the filtered water samples were frozen (−20°C) for further analysis. The analyses of nitrate and nitrite were performed using an automatic system—AutoAnalyzer II—Bran-Luebbe®. The silicate and phosphate analyses were processed by a spectrophotometric



**Fig. 1** Admiralty Bay (summer 2002/03): Sampling areas (a) (CF Brazilian Antarctic Research Station ‘Comandante Ferraz’; BP Botany Point; He Hennequin Point; Ez Ezcurra Inlet) and sampling stations with different water column depths (b) (A 20 m; B 30 m; C 60 m) at each sampling area

method and measured in a Genesys II spectrophotometer, Bausch & Lomb®. All these analyses followed the recommendations of Grasshoff et al. (1983).

### Phytoplankton and chlorophyll a analyses

For these analyses, the water samples were collected using Nansen bottles. Chlorophyll *a* was extracted from 1 l filtered samples (Whatman GF/F membrane), frozen (−20°C), and determined by fluorimetry after extraction in 90% acetone, using a Turner TD-7000 fluorometer (Parsons et al. 1984).

Phytoplankton samples were stored in 250 ml dark bottles and fixed with 2% borax-buffered formaldehyde, and analyses were taken according to the Utermöhl (1958) method.

The cells were allowed to settle in 50–100 ml chambers for 48–72 h. Using an inverted microscope (Olympus®IX70) equipped with phase contrast at a magnification of 200× (ocular 10×, objective 20×), organisms larger than 20 μm were analyzed throughout the whole counting chamber. A higher magnification (400×) was used to identify the smallest taxa, such as some diatoms.

Species composition was determined based on the following works: Peragallo and Peragallo (1921), Cupp (1943), Medlin and Priddle (1990), Round et al. (1990) and Hasle and Syvertsen (1997).

The diatom taxa that could not be identified were grouped according to their morphotypes in: Group 1 (pennate diatoms, free cells), Group 2 (pennate diatoms, chain-forming), Group 3 (centric diatoms, free cells).

The occurrence of taxa was considered *frequent* (more than 40% of samples) or *constant* (more than 80% of samples). In terms of abundance, they were categorized according to Lobo and Leighton (1986) as *abundant* (one particular species being higher than the mean abundance of all species that appear in the sample) or *dominant* (one species constituting more than 50% of the abundance of the sample).

### Statistics

In order to normalize distributions and eliminate zero values, the biological data were transformed using the log factor:  $\log_{10}(x + 1)$ . The Correspondence Analysis considered the biological data to distinguish patterns of distribution of phytoplankton populations. Results were compared using a One-Way ANOVA with a Kruskal–Wallis test ( $P < 0.05$ )

for multiple comparisons. All these analyses were performed using Statistica® version 6.0 (Statsoft).

### Results

The values of hydrological and biological data for the austral summer of 2002–03 are listed in Table 1. During the early summer, the water was relatively cold ( $-0.4 \pm 0.2^\circ\text{C}$ ) compared to late summer ( $1.5 \pm 0.3^\circ\text{C}$ ,  $P < 0.01$ ), while the salinity showed little variation (34.3 to 34.8) (Fig. 2a).

Dissolved oxygen showed highest concentrations ( $6.4 \pm 1.2 \text{ ml l}^{-1}$ ) during early summer (Fig. 2b). Phosphate, as well as temperature, increased in late summer.

The estimated ratios N:P showed three groups of samples: (1) Early summer, N:P ratio between 10 and 15, associated with the lowest temperatures; (2) Late summer, N:P ratio  $< 10$  influenced by highest phosphate concentrations; (3) Late summer, N:P ratio  $> 10$  because of a high inorganic nitrogen availability associated with low phosphate concentrations (Fig. 2c).

Chlorophyll *a* concentrations were constantly low ( $0.44 \pm 0.29 \mu\text{g l}^{-1}$ ), except for two peaks ( $> 1 \mu\text{g l}^{-1}$ ) from the surface waters at Hennequin Point (St. A). The silicate values did not vary.

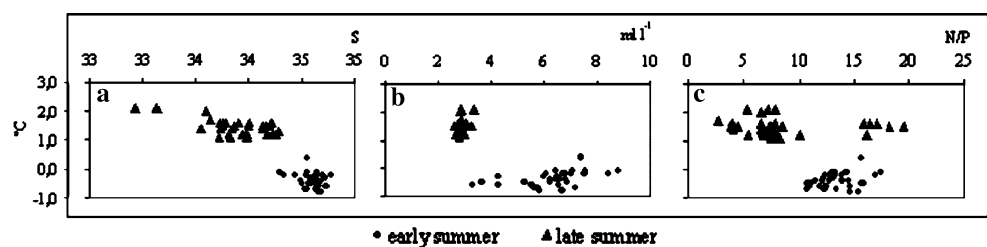
**Table 1** Admiralty Bay (summer 2002/03)

Variables	Range		Average		SD		N	
	ES	LS	ES	LS	ES	LS	ES	LS
Temperature—T ( $^\circ\text{C}$ )	-0.8 to 0.4	1.1–2.1	-0.4	1.5	0.2	0.3	36	36
Salinity—S	34.3–34.8	32.9–35.8	34.6	33.9	0.1	0.5	36	35
Dissolved oxygen—DO ( $\text{ml l}^{-1}$ )	3.3–8.8	2.7–3.4	6.4	2.9	1.2	0.1	36	36
Dissolved inorganic nitrogen—DIN ( $\mu\text{mol l}^{-1}$ )	23.7–46.9	16.6–38.6	35.0	33.5	5.0	4.1	36	36
Silicate—Si-SiO <sub>4</sub> ( $\mu\text{mol l}^{-1}$ )	38.0–91.3	20.8–99.2	67.2	66.0	13.5	20.2	36	33
Phosphate—P- PO <sub>4</sub> ( $\mu\text{mol l}^{-1}$ )	1.2–3.0	0.2–9.9	2.6	4.5	0.3	2.5	36	36
Chlorophyll <i>a</i> —Chl <i>a</i> ( $\mu\text{g l}^{-1}$ )	0.1–1.7	0.2–0.9	0.4	0.5	0.4	0.2	12	36
Phytoplankton abundance ( $\text{cells l}^{-1}$ )	$5.1 \times 10^1$ – $1.6 \times 10^4$	$2.4 \times 10^2$ – $6.9 \times 10^3$	$1.4 \times 10^3$	$1.5 \times 10^3$	$3.5 \times 10^3$	$1.5 \times 10^3$	36	36

Hydrochemical and biological data (ranges, averages, standard deviations and number of samples)

ES early summer, LS late summer

**Fig. 2** Admiralty Bay (summer 2002/03): Relationships between seawater temperature and salinity (a), dissolved oxygen (b) and the ratio N/P (c)



## Microphytoplankton

The microphytoplankton assemblage was diverse, mostly represented by diatoms (87 taxa), dinoflagellates (12 taxa), cyanobacteria (8 taxa—mainly Nostocales and Oscillatoriales) and flagellates (6 taxa—mainly unidentified cryptophyceans).

In general, the microphytoplankton abundance was low ( $1.5 \times 10^3$  cells  $l^{-1} \pm 2.7 \times 10^3$  cells  $l^{-1}$ ) with a high standard deviation because of the temporal and spatial variations. Diatoms were dominant in terms of the number of taxa (77%) and abundance (90%).

### Temporal variation

Composition, number of taxa (Table 2) and abundance of microphytoplankton differed during the study periods. In early summer, the highest number of taxa ( $P > 0.05$ ) occurred due to the contribution of pennate tycho planktonic diatoms in the water column. However, the abundance ( $6.2 \times 10^2 \pm 6.9 \times 10^2$  cells  $l^{-1}$ ) was lower than in late summer ( $1.5 \times 10^3 \pm 1.5 \times 10^3$  cells  $l^{-1}$ ) (Fig. 3), except for the surface waters at station A, at Hennequin Point, where, the chain-forming pennate benthic diatoms (cell linear length  $\cong 60 \mu m$ ) were responsible for the high

abundance ( $>1.5 \times 10^4$  cells  $l^{-1}$ ). Pennate diatoms reached 85% of abundance during early summer.

In late summer, centric (55%) and pennate diatoms (42%) dominated the phytoplankton assemblage, and the highest abundance was caused by centric diatoms.

Dinoflagellates showed highest abundance ( $P < 0.01$ ) in late summer, and the other groups (cyanobacteria and flagellates) were not well represented during early or late summer.

### Spatial variation

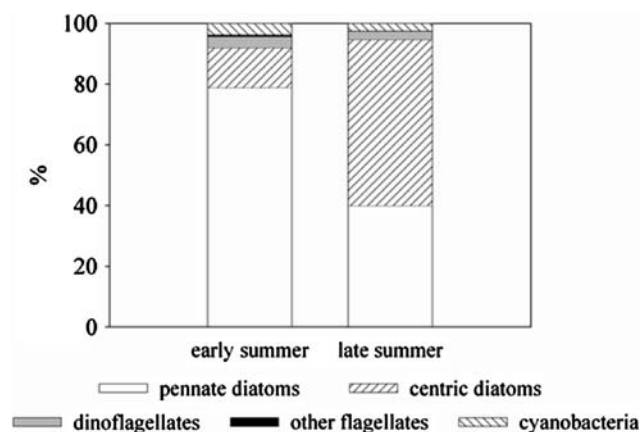
The phytoplankton abundance (Fig. 4a) was different between sampling areas ( $P = 0.03$ ). The EACF ( $5.2 \times 10^2 \pm 3.7 \times 10^2$  cells  $l^{-1}$ ) and Ezcurra Inlet ( $1.2 \times 10^3 \pm 0.8 \times 10^3$  cells  $l^{-1}$ ) sample areas had the lowest ( $<1.5 \times 10^3$  cells  $l^{-1}$ ) and most homogeneous ( $SD < 10^3$  cells  $l^{-1}$ ) values. Conversely, the highest ( $>1.5 \times 10^3$  cells  $l^{-1}$ ) and most heterogeneous ( $SD \geq 10^3$  cells  $l^{-1}$ ) values were found at Botany Point ( $1.8 \times 10^3 \pm 2.1 \times 10^3$  cells  $l^{-1}$ ) and Hennequin Point ( $2.3 \times 10^3 \pm 4.8 \times 10^3$  cells  $l^{-1}$ ) (Fig. 4a) due to the high abundance ( $>10^4$  cells  $l^{-1}$ ) of chain-forming pennate diatoms (*Fragilaria striatula* Lyngbye, *Fragilariopsis* Hustedt) (Fig. 4b).

There were no significant differences between the sampling stations in regard to the microphytoplankton

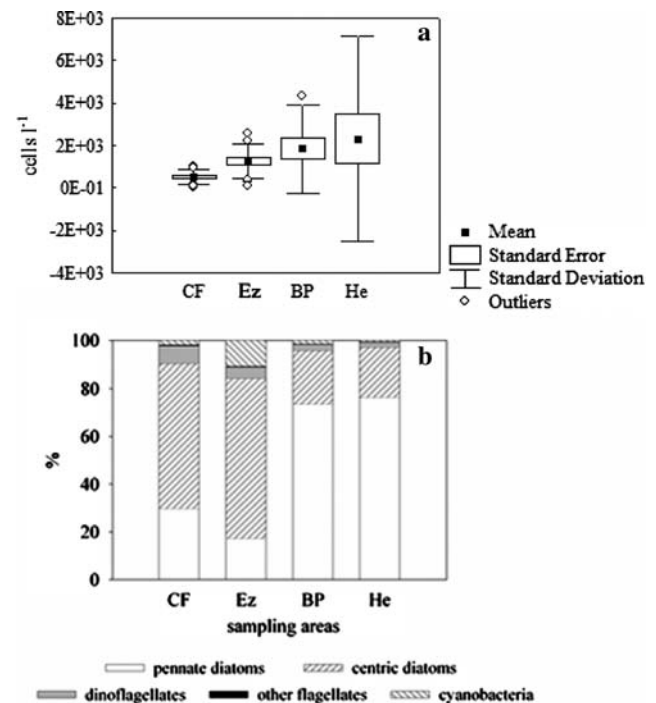
**Table 2** Admiralty Bay (summer 2002/03)

	Early summer	Late summer	Both periods
Cyanobacteria	5	7	8
Diatoms	75	56	87
Dinoflagellates	11	7	12
Other flagellates	6	5	6
Total	97	75	113

Number of taxa from each taxonomic group



**Fig. 3** Admiralty Bay (summer 2002/03): Percentage abundance of the major taxonomic groups during early and late summer



**Fig. 4** Admiralty Bay (summer 2002/03): Phytoplankton abundance (a) (mean, SE and SD) and percentage abundance of the major taxonomic groups (b) at the sampling areas: CF Brazilian Antarctic Research Station ‘Comandante Ferraz’; Ez Ezcurra Inlet; BP Botany Point and He Hennequin Point

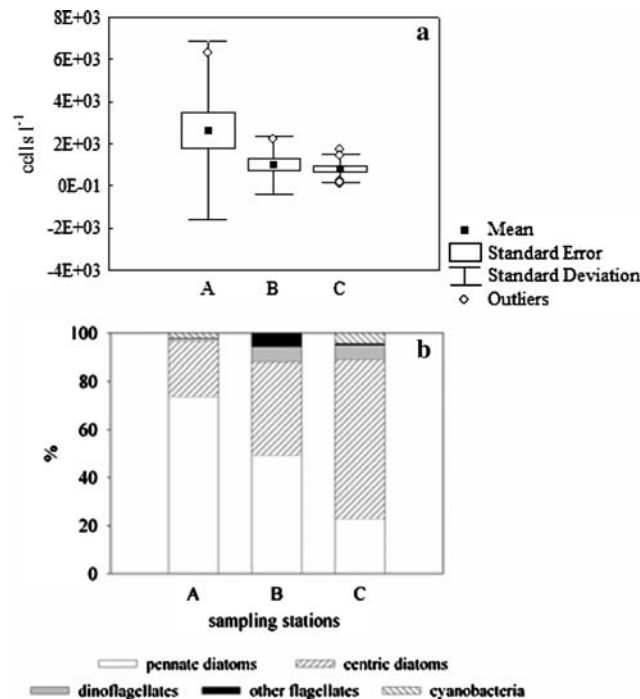
abundance (Fig. 5a). However, there were differences ( $P = 0.02$ ) in the taxonomic groups' representation. The highest contribution was from the pennate diatoms in shallow water (80%) at St. A, whereas the centric diatoms were more abundant (65%) at St. C. The cyanobacteria and flagellates were sparse at St. A, while at the deepest stations they represented more than 5% of the total abundance (Fig. 5b).

Different sampling areas had distinct vertical distribution patterns (Fig. 6), with highest abundance in surface water at Hennequin Point in early summer ( $>5 \times 10^3$  cells  $l^{-1}$ ) and Botany Point in late summer ( $>4 \times 10^3$  cells  $l^{-1}$ ). There was a decreasing vertical distribution gradient in early summer, but a homogeneous gradient in late summer at Ezcurra Inlet. No vertical distribution variation was observed at EACF.

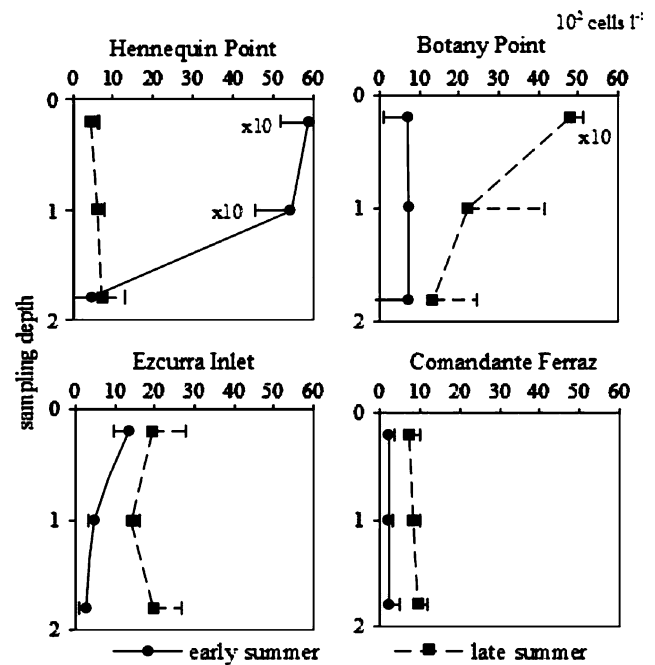
*Species composition and Correspondence Analysis*

Considering the 113 taxa identified in Admiralty Bay's flora, and according to the criteria of abundance and occurrence, 31 taxa were selected for the Correspondence Analysis, which shows the temporal and spatial distribution of the taxa.

The I  $\times$  II factorial plan (Fig. 7a, b) represented 42% of the total variance and isolated the samples from late summer in the positive portion of Axis I (23%). These samples



**Fig. 5** Admiralty Bay (summer 2002/03): Phytoplankton abundance (a) (mean, SE and SD) and percentage abundance of the major taxonomic groups (b) at stations with different water column depths (A 20 m; B 30 m; C 60 m)



**Fig. 6** Admiralty Bay (summer 2002/03): Phytoplankton abundance at three depths in the water column (0 surface, 1 midpoint of water column and 2 close to the bottom), at the four sampling areas

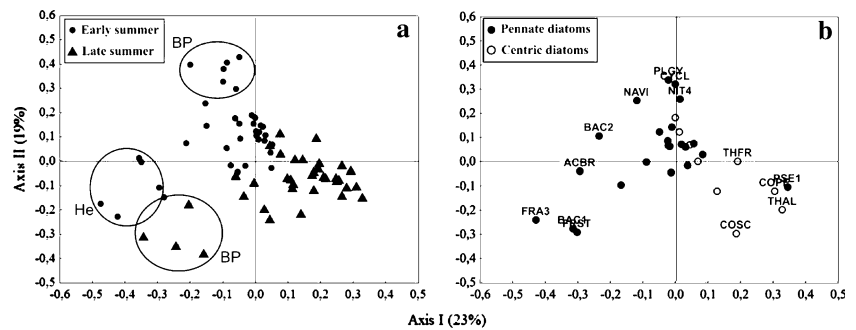
were characterized by the dominance of centric diatoms, such as *Corethron pennatum* (Grunow) Ostenfeld (39%), *Thalassiosira anguste-lineata* (Schmidt) Fryxell & Hasle, *Thalassiosira cf. frenguellii* Kozlova, *Thalassiosira* spp., and some species identified as "Group 3", besides the pennate diatom *Pseudo-nitzschia "delicatissima"* (Cleve) Heiden & Kolbe complex.

The surface samples from shallow stations (St. A and B) appear in the negative portion of Axis I. These were characterized by high concentrations ( $>1.5 \times 10^3$  cells  $l^{-1}$ ) of pennate diatoms, mainly: Group 1, benthic species such as the chain-forming *Fragilariopsis ritscheri*,—especially at Hennekin Point ( $>30\%$ )—*Fragilaria striatula*, *Achnanthes brevipes* (Kützing) Cleve and others; along with Group 2, the single-celled *Licmophora gracilis* (Ehrenberg) Grunow and unidentified single pennate diatoms.

Axis II (19%) established a separation between early and late summer of the Botany Point sampling area. The early summer samples (positive portion) were characterized by the highest abundance ( $>20\%$ ) of species included in *Pleurosigma*/*Gyrosigma* and *Cylindrotheca closterium* (Ehrenberg) Lewin & Reimann/*Nitzschia longissima* (Brébisson) Ralfs complexes. In samples from late summer (negative portion) the pennate diatom *Fragilaria striatula* was dominant (55% of abundance) (Fig. 7b).

The centric *Corethron pennatum* was the only species that was present in all four sampling areas and during both sampling periods ( $>88\%$  of the samples). Moreover, this

**Fig. 7** Admiralty Bay (summer 2002/03): correspondence analysis of microphytoplankton species: sampling areas (a); *He* Hennequin Point, *BP* Botany Point; taxa (b). see Table 3



species was considered *abundant* in 68% of the samples and *dominant* in 31% (mostly in late summer).

The pennate diatoms, such as *Licmophora gracilis*, *Pseudogomphonema kamtschaticum* (Grunow) Medlin, *Cocconeis* cf. *costata* (Gregory) Cleve and the *Cylindrotheca closterium/Nitzschia longissima* complex, had an important role in Admiralty Bay's phytoplankton population. They were considered *frequent* (>50% of the samples) and *abundant* (Table 3).

## Discussion

In the context of water column production, the Admiralty Bay nearshore water can be considered a high nutrient—low chlorophyll (HNLC) area (Platt et al. 2003). The pattern of temporal variation of the inorganic nutrients distribution is correlated with the biogeochemical processes and physical dynamics of the Bay (light incidence, melting ice, oceanic input, resuspended sediments, and wind stress generating advective processes). This can be seen in the N:P ratios, which showed the lowest values ( $8.5 \pm 4.5$ ) in late summer as a result of the highest phosphate concentrations ( $4.5 \pm 2.5 \mu\text{mol l}^{-1}$ ). During the 2002–03 summer, the microphytoplankton abundance was low ( $10^3 \text{ cells l}^{-1}$ ) when compared to values reported in other Antarctic areas like the Weddell Sea (Kang et al. 2001, Moisan and Fryxell 1993, Estrada and Delgado 1990) and Prydz Bay (Kang and Fryxell 1991), but similar to Maxwell Bay (Ahn et al. 1997), Bransfield Strait and Bellingshausen Sea (Bidigare et al. 1996) (Table 4).

The abundance values shown by different authors should be compared with caution, as different methods may have been used. In this work, cells smaller than  $20 \mu\text{m}$  were disregarded. Conversely, the authors cited above took into account micro (> $20 \mu\text{m}$ ) and nanoplankton ( $2\text{--}20 \mu\text{m}$ ) (Brandini et al. 1997). This may be the cause of the higher values found by these authors compared to those found in this work. Another reason may be that the sampling was carried out before and after but not within the high primary productivity season in the mid-summer...

The absence of strong water currents that resuspend the sediment allow the microalgae to sink to the bottom of the euphotic zone, causing low phytoplankton abundance ( $\leq 10^3 \text{ cells l}^{-1}$ ) in the water column at Admiralty Bay (Rakusa-Suszczewski 1980). Conversely, strong turbulence may have been present only at the shallowest station (St. A) at Hennequin Point during the 2002–03 early summer, and probably caused the highest phytoplankton abundance ( $10^4 \text{ cells l}^{-1}$ ) and chlorophyll *a* concentrations ( $>1 \mu\text{g l}^{-1}$ ). These high phytoplankton abundance and chlorophyll *a* concentrations have also been reported previously by Brandini and Rebelló (1994) at the same area.

With the exception of Hennequin Point, Admiralty Bay showed the highest phytoplankton abundance in late summer with a direct relation to the water temperature. El-Sayed and Fryxell (1993), Kang et al. (1997), Kopczyńska et al. (1998) and Kang et al. (2002) reported similar results from other areas in the Southern Ocean. The lowest phytoplankton abundance ( $<10^2 \text{ cells l}^{-1}$ ) and chlorophyll *a* concentrations ( $0.22 \pm 0.05 \mu\text{g l}^{-1}$ ) associated with the highest ammonia values ( $5.7 \pm 2.9 \mu\text{mol l}^{-1}$ ) near the EACF area suggest a response from the environment to either water turbulence (Brandini and Rebelló 1994) or human presence at the Brazilian station along with mammals and penguins that often use the Keller Peninsula throughout the summer. Further study is necessary to distinguish between these two influences.

## Dominant species

The centric diatom *Corethron pennatum* and several species of the pennate diatom genus *Fragilariopsis* were dominant in Admiralty Bay in the summer of 2002/03, as found previously by Kopczyńska (1993) in Admiralty Bay, by Kang and Fryxell (1991), and Kopczyńska et al. (1995) in Prydz Bay.

The similarity between this flora and the one found in Bellingshausen Sea (Bidigare et al. 1996) and Bransfield Strait (Burkholder and Sieburth 1961) strengthens the hypothesis that the offshore waters from Bellingshausen Sea inflow through Bransfield Strait, as suggested by Lipski (1987) and Madejski and Rakusa-Suszczewski (1990). The Weddell Sea presents some similarities to Admiralty Bay as

**Table 3** Admiralty Bay (summer 2002/2003)

Group	Taxa	Code	Early summer		Late summer	
			Abundance (cells l <sup>-1</sup> )	Criteria	Abundance (cells l <sup>-1</sup> )	Criteria
Pennate diatoms	<i>Achnanthes brevipes</i> (Kützing) Cleve	ACBR	3.3 × 10 <sup>3</sup>	A	3.5 × 10 <sup>1</sup>	–
	<i>Achnanthes</i> cf. <i>taeniata</i> Grunow	ACTA		–	1.1 × 10 <sup>3</sup>	A
	<i>Amphora</i> cf. <i>proteus</i> Gregory	AMPR	8.1 × 10 <sup>1</sup>	A	1.2 × 10 <sup>1</sup>	–
	<i>Amphora</i> spp		4.0 × 10 <sup>1</sup>	–	7.0 × 10 <sup>0</sup>	–
	<i>Banquisia</i> sp			–	6.0 × 10 <sup>0</sup>	–
	<i>Cocconeis costata</i> (Gregory) Cleve	COCO	2.4 × 10 <sup>2</sup>	A	2.5 × 10 <sup>2</sup>	F
	<i>Cocconeis</i> cf. <i>pinnata</i> (Gregory) Greville		3.6 × 10 <sup>1</sup>	–	7.0 × 10 <sup>0</sup>	–
	<i>Cocconeis gautieri</i> Van Heurck			–	5.0 × 10 <sup>0</sup>	–
	<i>Cocconeis orbicularis</i> Frenguelli & Orlando	COOR	6.3 × 10 <sup>1</sup>	A		–
	<i>Cocconeis</i> spp	COCC	1.7 × 10 <sup>2</sup>	A	6.6 × 10 <sup>1</sup>	–
	<i>Cylindrotheca closterium</i> (Ehrenberg) Lewin & Reimann/ <i>Nitzschia longissima</i> (Brébisson) Ralfs complex	CYCL	2.2 × 10 <sup>3</sup>	D/F	2.2 × 10 <sup>2</sup>	A
	<i>Diploneis</i> sp			–	3.0 × 10 <sup>0</sup>	–
	<i>Fragilaria striatula</i> Lyngbye	FRST	8.5 × 10 <sup>2</sup>	A	1.2 × 10 <sup>4</sup>	D
	<i>Fragilariopsis</i> cf. <i>doliolus</i> (Wallich) Medlin & Sims		2.9 × 10 <sup>1</sup>	–		–
	<i>Fragilariopsis cylindrus</i> (Grunow) Krieger			–	8.0 × 10 <sup>0</sup>	–
	<i>Fragilariopsis kerguelensis</i> (O'Meara) Hustedt		1.8 × 10 <sup>1</sup>	–		–
	<i>Fragilariopsis obliquecostata</i> (Van Heurck) Hasle		3.5 × 10 <sup>1</sup>	–		–
	<i>Fragilariopsis rhombica</i> (O'Meara) Hustedt		3.6 × 10 <sup>1</sup>	–	3.5 × 10 <sup>1</sup>	–
	<i>Fragilariopsis ritscheri</i> Hustedt	FRAG	1.5 × 10 <sup>4</sup>	D	4.0 × 10 <sup>3</sup>	A
	<i>Licmophora belgicae</i> Peragallo		2.4 × 10 <sup>2</sup>	–	3.9 × 10 <sup>1</sup>	–
	<i>Licmophora</i> cf. <i>plana</i> Heiden & Kolbe			–	3.7 × 10 <sup>1</sup>	–
	<i>Licmophora gracilis</i> (Ehrenberg) Grunow	LIGR	2.8 × 10 <sup>3</sup>	A/F	3.9 × 10 <sup>2</sup>	F
	<i>Manguinea</i> sp		2.5 × 10 <sup>1</sup>	–	3.0 × 10 <sup>0</sup>	–
	<i>Mastogloia</i> cf. <i>smithii</i> Smith		2.4 × 10 <sup>1</sup>	–		–
	<i>Membraneis challengerii</i> Grunow		8.4 × 10 <sup>1</sup>	–	2.4 × 10 <sup>1</sup>	–
	<i>Navicula algida</i> Grunow		8.4 × 10 <sup>1</sup>	–	7.9 × 10 <sup>1</sup>	–
	<i>Navicula</i> cf. <i>septentrionalis</i> (Grunow) Gran.		3.0 × 10 <sup>2</sup>	–	1.1 × 10 <sup>1</sup>	–
	<i>Navicula</i> cf. <i>transitans</i> Grunow	NATR	2.2 × 10 <sup>2</sup>	A		–
	<i>Navicula</i> cf. <i>vanhoeffenii</i> Gran.		1.2 × 10 <sup>1</sup>	–		–
	<i>Navicula directa</i> (Smith) Ralfs		2.3 × 10 <sup>2</sup>	–	1.3 × 10 <sup>1</sup>	–
	<i>Navicula</i> sp		3.6 × 10 <sup>1</sup>	–		–
	Naviculaceae	NAVI	1.3 × 10 <sup>3</sup>	A/F	1.7 × 10 <sup>2</sup>	–
	<i>Neodenticula</i> sp		4.0 × 10 <sup>0</sup>	–		–
	<i>Nitzschia braarudii</i> Hasle		1.1 × 10 <sup>2</sup>	–		–
	<i>Nitzschia</i> cf. <i>longa</i> Grunow		3.0 × 10 <sup>0</sup>	–		–
	<i>Nitzschia lecointei</i> Van Heurck	NILE	2.3 × 10 <sup>2</sup>	A	1.2 × 10 <sup>1</sup>	–
<i>Nitzschia</i> sp1	NIT1	1.8 × 10 <sup>2</sup>	A		–	
<i>Nitzschia</i> sp2	NIT2	9.9 × 10 <sup>1</sup>	A		–	
<i>Parlibellus delognei</i> (Van Heurck) Cox.		6.0 × 10 <sup>0</sup>	–	8.0 × 10 <sup>0</sup>	–	
Pennate diatoms, free cells (group 1)	GROUP 1	8.8 × 10 <sup>3</sup>	A	1.9 × 10 <sup>3</sup>	A	
Pennate diatoms, chain-forming (group 2)	GROUP 2	2.2 × 10 <sup>3</sup>	A/F	2.3 × 10 <sup>1</sup>	–	
<i>Pinnularia quadratarea</i> (Schmidt) Cleve		2.7 × 10 <sup>1</sup>	–		–	
<i>Pleurosigma</i> / <i>Gyrosigma</i> complex	PLGY	1.7 × 10 <sup>3</sup>	A/F	1.0 × 10 <sup>2</sup>	–	
<i>Pseudogomphonema kamtshaticum</i> (Grunow) Medlin	PSKA	1.1 × 10 <sup>3</sup>	A/F	1.2 × 10 <sup>3</sup>	A/C	

**Table 3** continued

Group	Taxa	Code	Early summer		Late summer	
			Abundance (cells l <sup>-1</sup> )	Criteria	Abundance (cells l <sup>-1</sup> )	Criteria
	<i>Pseudo-nitzschia</i> “ <i>delicatissima</i> ” (Cleve) Heiden & Kolbe complex	PSEU	3.6 × 10 <sup>1</sup>	–	1.4 × 10 <sup>3</sup>	A/F
	<i>Rhopalodia</i> sp		6.0 × 10 <sup>0</sup>	–		–
	<i>Surirella</i> sp		3.0 × 10 <sup>0</sup>	–	7.0 × 10 <sup>0</sup>	–
	<i>Synedropsis</i> spp	SYNE	7.8 × 10 <sup>2</sup>	A	2.6 × 10 <sup>1</sup>	–
	<i>Thalassionema</i> sp		2.2 × 10 <sup>2</sup>	–		–
	<i>Thalassiothrix antarctica</i> Schimper & Karsten		3.0 × 10 <sup>0</sup>	–		–
	<i>Trichotoxon</i> sp	TRIC	8.0 × 10 <sup>0</sup>	A	1.1 × 10 <sup>1</sup>	–
	<i>Tropidoneis</i> cf. <i>antarctica</i> Cleve	TRAN	1.3 × 10 <sup>2</sup>	A	1.8 × 10 <sup>1</sup>	–
	<i>Tropidoneis</i> spp		3.8 × 10 <sup>1</sup>	–		–
Centric diatoms	<i>Actinocyclus actinochilus</i> (Ehrenberg) Simonsen			–	2.3 × 10 <sup>2</sup>	–
	<i>Actinoptychus senarius</i> (Ehrenberg) Ehrenberg			–	1.2 × 10 <sup>1</sup>	–
	<i>Asteromphalus</i> sp		3.0 × 10 <sup>0</sup>	–		–
	Centric diatoms, free cells (group 3)	GROUP 3	7.1 × 10 <sup>2</sup>	A	3.0 × 10 <sup>3</sup>	A/C
	<i>Corethron inerme</i> Karsten		5.7 × 10 <sup>1</sup>	–		–
	<i>Corethron pennatum</i> (Grunow) Ostenfeld	COPE	1.1 × 10 <sup>3</sup>	D/F	2.1 × 10 <sup>4</sup>	D/C
	<i>Coscinodiscus asteromphalus</i> Ehrenberg		1.6 × 10 <sup>1</sup>	–	2.2 × 10 <sup>2</sup>	–
	<i>Dactyliosolen antarcticus</i> Castracane			–	2.4 × 10 <sup>1</sup>	–
	<i>Melosira sol</i> (Ehrenberg) Kützing	MESO	1.3 × 10 <sup>2</sup>	A	9.0 × 10 <sup>0</sup>	–
	<i>Odontella litigiosa</i> (Van Heurck) Hoban	ODLI	3.4 × 10 <sup>2</sup>	A	1.2 × 10 <sup>1</sup>	–
	<i>Porosira pseudodenticulata</i> (Hustedt) Jousé	POPS	2.7 × 10 <sup>1</sup>	–	1.7 × 10 <sup>2</sup>	A
	<i>Stellarima microtrias</i> (Ehrenberg) Hasle & Sims		7.3 × 10 <sup>1</sup>	–	1.4 × 10 <sup>2</sup>	–
	<i>Thalassiosira anguste-lineata</i> (Schmidt) Fryxell & Hasle	THAN		–	4.5 × 10 <sup>2</sup>	F
	<i>Thalassiosira</i> cf. <i>frenguelli</i> Kozlova	THFR	2.5 × 10 <sup>1</sup>	A	2.4 × 10 <sup>2</sup>	–
	<i>Thalassiosira ritscheri</i> (Hustedt) Hasle in Hasle & Heimdal		3.0 × 10 <sup>0</sup>	–	1.3 × 10 <sup>2</sup>	–
	<i>Thalassiosira</i> spp	THAL	5.4 × 10 <sup>1</sup>	A	2.7 × 10 <sup>3</sup>	D/F

List of diatoms taxa with the code (for the Correspondence Analysis), abundance (cells l<sup>-1</sup>) and the criteria of abundance (A abundant, D dominant) and occurrence (F frequent, C constant)

well, where diatoms such as *Nitzschia lecointei* Van Heurck and several species of the genera *Fragilariopsis*, *Thalassiosira* and *Pseudo-nitzschia* are most frequently found (Kang et al. 2001).

Some areas inside Admiralty Bay such as Hennequin Point and Botany Point have similar flora to those found in Maxwell Bay (King George Island) which is highly influenced by ice and benthic diatoms (*Fragilaria striatula*, *Achnanthes brevipes*, *Fragilariopsis ritscheri*) (Ahn et al. 1997). However, the processes that allow benthic diatoms in the water column in Admiralty Bay may be different. Instead of being detached from the sea ice only, these algae also may be carried from the bottom to the water column by upwelling water currents (Brandini and Rebello 1994; Gilbert 1991), or by low saline meltwater-sediment suspension (Pichlmaier et al 2004).

The influence of temporal variation on sea ice cover and phytoplankton populations was identified by the alternate dominance of pennate and centric diatoms. After a heavy winter during which most of Admiralty Bay froze (INPE 2002), the sea ice within the bay melted in approximately 2 months during the spring (November and December). This fast melting may have caused an increase of benthic and ice diatoms, such as *Achnanthes brevipes*, *Cocconeis* spp., *Navicula directa* (Smith) Ralfs, *Fragilaria striatula* in the water column. However, during late summer, there was an increase in the abundance of the centric diatoms *Corethron pennatum*, *Actinocyclus actinochilus* (Ehrenberg) Simonsen, and *Thalassiosira ritscheri* (Hustedt) Hasle in Hasle & Heimdal derived from open waters, possibly inflowing from Bellingshausen Sea (Rakusa-Suszczewski 1980).



**Table 4** Microphytoplankton abundance (cells l<sup>-1</sup>) and dominant taxa in some areas of Antarctica

Area	Location	Cells l <sup>-1</sup>	Dominant taxa
Admiralty Bay <sup>a</sup>	Nearshore	10 <sup>1</sup> –10 <sup>4</sup>	<i>Corethron pennatum</i> (Grunow) Ostenfeld <i>Fragilariopsis ritscheri</i> Hustedt <i>Fragilaria striatula</i> Lyngbye
Weddell Sea <sup>b, c, d</sup>	Oceanic	10 <sup>3</sup> –10 <sup>7</sup>	<i>Chaetoceros socialis</i> Lauder <i>Corethron pennatum</i> (Grunow) Ostenfeld <i>Fragilariopsis</i> spp <i>Nitzschia lecointei</i> Van Heurck <i>Proboscia truncata</i> (Karsten) Nöthig & Ligowski <i>Pseudo-nitzschia</i> spp. <i>Thalassiosira</i> spp.
Prydz Bay <sup>e</sup>	Nearshore	10 <sup>6</sup>	<i>Corethron pennatum</i> (Grunow) Ostenfeld <i>Cylindrotheca closterium</i> (Ehrenberg) Lewin & Reimann <i>Fragilariopsis</i> spp
Maxwell Bay <sup>f</sup>	Nearshore	10 <sup>3</sup> –10 <sup>5</sup>	<i>Achnanthes brevipes</i> (Kützing) Cleve <i>Fragilaria striatula</i> Lyngbye <i>Licmophora</i> spp
Bransfield Strait <sup>g</sup>	Oceanic	10 <sup>3</sup> –10 <sup>5</sup>	<i>Fragilariopsis</i> spp <i>Rhizosolenia</i> spp
Bellingshausen Sea <sup>g</sup>	Oceanic	10 <sup>3</sup> –10 <sup>6</sup>	<i>Corethron pennatum</i> (Grunow) Ostenfeld <i>Fragilariopsis</i> spp <i>Pseudo-nitzschia</i> spp <i>Thalassiosira</i> spp

<sup>a</sup> This study<sup>b</sup> Kang et al. (2001)<sup>c</sup> Moisan and Fryxell (1993)<sup>d</sup> Estrada and Delgado (1990)<sup>e</sup> Kang and Fryxell (1991)<sup>f</sup> Ahn et al. (1997)<sup>g</sup> Bidigare et al. (1996)

This temporal variation of phytoplankton composition was very clear at Hennequin Point. During early summer, there was the dominance of the chain-forming pennate diatom *Fragilariopsis ritscheri*, suggesting the influence of sea ice melting or sediment resuspension, as this diatom is abundant at marginal ice zones (Kang et al. 2001), into the sea ice (Crosta et al. 2004), and in surface sediments (Mohan et al. 2006). Conversely, during late summer, *Corethron pennatum* was the most abundant species, as found at the other areas of the bay.

At Botany Point, the dominance of benthic pennate diatoms during early (*Pleurosigma*/*Gyrosigma* complex) and late summer (*Fragilaria striatula*) reflected the high influence of resuspended sediments (Pichlmaier et al 2004).

The dominance of centric diatoms (60%) at EACF and Ezcurra Inlet was caused basically by *Corethron pennatum*, which reached abundances up to 10<sup>3</sup> cells l<sup>-1</sup>, mainly in late summer.

*C. pennatum* is cosmopolitan and planktonic (Fryxell 1989), and it is dominant during low-productivity seasons in this area, especially in Bransfield (von Bodungen 1986), suggesting the influence of Bransfield waters in the bay. Also, *C. pennatum* usually shows high abundances (10<sup>7</sup> cells l<sup>-1</sup>) under ice cover and at ice edges (Moisan and Fryxell 1993; Kang and Fryxell 1991; Garrison 1991).

The dominance of *C. pennatum*, which was present at the four sampling areas (88% of all samples) with relatively

high abundance (*abundant* in 68% of all samples), was previously reported in Admiralty Bay (Kopcsynska 1993) and areas adjacent to King George Island, such as Marian Cove at Maxwell Bay (Kang et al. 1997).

For the reasons mentioned above, it is suggested that *C. pennatum* is an important component for primary production and maintenance of the food web at Admiralty Bay. Key organisms like krill (*Euphausia superba*) preferably feed on this microalga (Schultes et al. 2006; Granéli et al. 1993). Some diatom taxa identified in this study of Admiralty Bay, such as *Fragilariopsis*, *Licmophora*, and *Thalassiosira*, are commonly grazed by krill as well (Schultes et al. 2006; Ligowski 2000).

Admiralty Bay's phytoplankton is dominated by diatoms, and highly influenced by benthic species, derived from the sediment resuspension or from the sea ice melting.

**Acknowledgments** This project is part of a Research Network dedicated to local environmental impact studies in Admiralty Bay, funded primarily by the Brazilian Ministry of Environment (MMA), Ministry of Science and Technology (MCT), and National Research Council (CNPq). We are thankful to the logistical support provided by Interministerial Secretary for the Sea Resources (SECIRM). We thank Dr. Rosalinda Montone (IO-USP) for the oxygen analysis, Vitor Chiozzini and Glaucia Berbel (IOUS-USP) for the salinity and nutrients analysis, Dr. Rodolfo Paranhos (UFRJ) for the chlorophyll *a* analysis, and Rafael B de Moura for drawing the map of Admiralty Bay. We are also very grateful to Dr. Silvia Nascimento (UENF), Dr. Frederico Brandini (UFPR), Dr. Elzbieta Kopczynska and two other unidentified reviewers for their useful suggestions to this manuscript, and to Alberto Bezerril and Martha Villac for the English review.

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