The effect of saxitoxin and non-saxitoxin extracts of *Cylindrospermopsis raciborskii* (Cyanobacteria) on cyanobacteria and green microalgae

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Abstract The effect of saxitoxins (STX) on phytoplankton species is poorly understood. To date, no correlation between STX concentrations and phytoplankton physiology has been reported. We investigated the effect of STX (STX+, 0.5–10 µg L^{-1} total STX) and non-STX (STX−, 0.5–10 µg L^{-1} total STX biomass equivalent) extracts of *Cylindrospermopsis raciborskii* on *Microcystis wesenbergii* BCCUSP11, *Microcystis aeruginosa* BCCUSP232 (microcystin producing), *Scenedesmus acuminatus* UFSCar036, and *Monoraphidium convolutum* CMEA/UFF0201 under controlled laboratory conditions. Both STX+ and STX− extracts inhibited the cell density and specific growth rate of *M. wesenbergii*, *M. aeruginosa*, and *S. acuminatus*. However, the effect of STX+ extract on the phytoplankton strains was significantly higher than that of STX− extract. *M. convolutum*, on the other hand, was tolerant as both STX+ and STX− extracts did not significantly reduce its cell density and specific growth rate (day^{-1}). The exposure of *M. aeruginosa* to STX+ and STX− resulted in higher total (intracellular and extracellular) microcystin concentration than the control. STX concentrations had a significant negative correlation with cell density and growth response of the phytoplankton strains investigated in this study. Conclusions can be made that although both STX+ and STX− extracts of *C. raciborskii* inhibited the growth of some phytoplankton species, the STX+ extracts were more toxic.

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Introduction

The potential of cyanobacteria to produce toxins makes their excessive proliferation a serious environmental problem (Chia et al. 2009a, b). Among the toxins produced by cyanobacteria, saxitoxins (STX) are alkaloids associated with paralytic shellfish poisoning (PSP) in freshwater ecosystems (Al-Tebrineh et al. 2010). The toxins affect cellular homeostasis by acting on sodium channels in both prokaryotes and eukaryotes (Neilan et al. 2013). STX are of high ecotoxicological risk due to the toxicity they induce in fish, mussels, copepods, mammals, and birds (Landsberg 2002). A number of cyanobacteria such as *Aphanizomenon flos-aquae*, *Cylindrospermopsis raciborskii*, *Lyngbya* spp., and *Anabaena* spp. synthesize STX in freshwater ecosystems. However, the production of this group of toxins tends to be strain specific (Molica et al. 2002; Bernard et al. 2003; Wiegand and Pflugmacher 2005; Al-Tebrineh et al. 2010). The discovery that *C. raciborskii* produces STX and cylindrospermopsin has made it an important cyanobacterium to be monitored in aquatic ecosystems. This is due to the increased frequency of its occurrence, scientific interest, improved water quality monitoring, availability of suitable habitat, and in some cases, its invasion of new habitats (Figueredo et al. 2007; Kling 2009; Sukenik et al. 2012). The cosmopolitan and increasing distribution of *C. raciborskii* is easily traced to its ecophysiological adaptation and tolerance to changing physicochemical

conditions in water bodies (Carneiro et al. 2013). The strains isolated from South America have been implicated for only STX production (Lagos et al. 1999; Molica et al. 2002; Costa et al. 2013; Hoff-Risseti et al. 2013).

Although a number of studies have shown that STX-producing cyanobacteria and dinoflagellates exhibit allelopathic effects on phytoplankton species, no direct correlation with STX has been reported (Tillmann and John 2002; Fistarol et al. 2004; Tillmann et al. 2008; Hattenrath-Lehmann and Gobler 2011). To the best of our knowledge, only one study using *Chlamydomonas reinhardtii* as a model organism showed that the microalga suffered oxidative stress after exposure to STX (Melegari et al. 2012). Therefore, generalizations cannot be made for microalgal response to STX based on the findings with *C. reinhardtii* alone (Perreault et al. 2011; Melegari et al. 2012). Ecotoxicological investigations that employ microalgae are of great significance because they can demonstrate the toxicity of bioactive compounds on different levels of the food chain. Studies of the effect of STX are important because microalgae possess characteristics such as voltage-dependent ion channels, photosynthetic ability, cellular metabolism, and structural specializations that are different from those found in animals (Wheeler and Brownlee 2008; Taylor 2009). This implies that STX interaction with algal cells may be very different from that with animal cells (Taylor 2009; Melegari et al. 2012).

To date, the ecological role of cyanotoxins remains generally unknown; however, according to one of the hypotheses, they may be associated with phytoplankton competition (Bar-Yosef et al. 2010; Rzymski et al. 2014). Microcystins (MC) are among the most studied cyanotoxins, and their production has been established to be geared toward giving the producer(s) a competitive advantage over neighboring species (Bittencourt-Oliveira et al. 2015). Furthermore, as a defense against predation, cyanobacteria (e.g., *Microcystis aeruginosa*) upregulate the synthesis of MC and other bioactive secondary metabolites (Pineda-Mendoza et al. 2014). However, a number of studies exclude this hypothesis due to lack of correlation between the production of MC and the rate of predation (Van Gremberghe et al. 2009; Wilken et al. 2010). The production of MC has also been shown to be regulated by physicochemical factors as well as the presence of chemical exudates from toxigenic and nontoxigenic algae (Downing et al. 2005; Deblois and Juneau 2010). Hence, we hypothesize that when threatened by the presence of cyanotoxins like STX, cyanobacteria can respond by upregulating MC synthesis as a defense mechanism.

Successional studies show that the occurrence of *Cylindrospermopsis* blooms is usually accompanied by the presence of chlorophytes of the genera *Monoraphidium* and *Scenedesmus* (Bouvy et al. 1999, 2006; Kokociński et al. 2010). Furthermore, there is evidence of the alternation between the dominance of green microalgae and cyanobacteria in different aquatic ecosystems (Zalocar de Domitrovic et al. 2007; Chia et al. 2012). However, the chemical interactions involved in this successional process are still not comprehensively understood. Previous studies have shown that some green algae are tolerant to MC (Bártová et al. 2011; Campos et al. 2013; Bittencourt-Oliveira et al. 2015), while not much is known about how STX affect green microalgae.

The objective of this study was to determine the effect of STX and non-STX extracts of *C. raciborskii* on the growth of green microalgae (*Scenedesmus acuminatus* and *Monoraphidium convolutum*) and cyanobacteria (*Microcystis wesenbergii* and *M. aeruginosa*). Furthermore, total (intracellular and extracellular) MC production by *M. aeruginosa* after exposure to STX and non-STX extracts of *C. raciborskii* was investigated. The results of the present study demonstrate for the first time the effect of STX on different phytoplankton species.

Materials and methods

The phytoplankton strains used in this study were *Microcystis wesenbergii* BCCUSP11 and *Microcystis aeruginosa* BCCUSP232 obtained from the Brazilian Cyanobacteria Collection of University of São Paulo (BCCUSP), *Scenedesmus acuminatus* UFSCar036 from the Microalgal Collection of Federal University of São Carlos (UFSCar), and *Monoraphidium convolutum* CMEA/UFF0201 from the Elizabeth Aidar Collection of Microalgae (CMEA/UFF). The *M. aeruginosa* BCCUSP232 strain is a MC-producing strain (Bittencourt-Oliveira 2003), while *M. wesenbergii* BCCUSP11 is a non-MC-producing strain (Bittencourt-Oliveira et al. 2011). All strains were maintained in ASM-1 liquid medium (Guillard 1973) at pH 7.4 using climatic chambers with controlled light intensity (40 μmol photons m^{-2} s^{-1}, measured with a LI-COR model LI-250 photometer equipped with an underwater spherical sensor), photoperiod (14:10 h, light/dark), and temperature (24±1 °C).

Preparation and quantification of cyanobacterial extracts

The STX and non-STX extract sources were *C. raciborskii* ITEPA1 and *C. raciborskii* ITEP31, respectively. Both *C. raciborskii* strains were obtained from the Cyanobacterial Collection of the Technological Institute of Pernambuco (ITEP), Brazil. The biomass required for extract preparation was obtained by culturing approximately 20 L of each *C. raciborskii* strain under the controlled conditions stated above. At exponential growth phase, the cultures were centrifuged at 9500 rcf for 15 min at 20 °C, and the resulting pellet was lyophilized and stored at −80 °C until needed.

Total STX concentration in the STX+ crude extract was determined following the procedures outlined in Oshima (1995). Briefly, the lyophilized material was extracted in

0.05 N acetic acid. The extract was analyzed using high-performance liquid chromatography (HPLC) equipped with an online postcolumn fluorescence derivatization system (Shimadzu, Japan). The different STX variants were identified and quantified by comparison with the retention time and the areas of standard peaks. The standards (≥98 % purity) were purchased from the National Research Council, Canada—Certified Reference Materials Program. The HPLC analysis results showed that dc-saxitoxin (0.41 µg mg^{-1}, 62.58 %), neo-saxitoxin (0.21 µg mg^{-1}, 32.50 %), and saxitoxin (0.03 µg mg^{-1}, 4.90 %) made up the total STX produced by *C. raciborskii* ITEPA1 (STX+).

Preparation and addition of aqueous crude extract to the cultures

Crude aqueous extracts were prepared from 76 mg of lyophilized biomass. The biomass was resuspended in 8 mL autoclaved deionized water and ultrasonicated at 15 W and 22.5 kHz for 5 min (Microson Ultrasonic Cell Disruptor, USA), to obtain an initial concentrated total STX solution. Subsequently, this initial concentrated total STX solution was serially diluted to arrive at the desired final exposure concentrations of 0.5, 1.0, 5.0, and 10.0 µg L^{-1} total STX, which corresponded to 0.75, 1.5, 7.5, and 15 mg L^{-1} *C. raciborskii* wet weight, respectively. For the non-STX (STX−) treatments, the lyophilized biomass was extracted following the same procedures as the STX+-containing biomass, and the extracts were added to the cultures at equivalent biomass of the STX+ extracts.

Experimental design

The cyanobacteria and green algae were exposed separately to STX+ and STX− extracts for 10 days. The extracts were added to the cultures on day 4 of the experiment. The cultures were maintained in climatic chambers set to luminous intensity of 40 µmol photons m^{-2} s^{-1}, photoperiod 14:10 h (light/dark), and temperature 24±1 °C. For each experiment, 1000 mL Erlenmeyer flasks containing 600 mL of ASM-1 culture medium were used. All experiments were carried out in triplicates. The initial cell density for all experimental treatments was 1.0×10^5 cells mL^{-1}.

Determination of cell density and specific growth rates

Aliquots (2 mL) were taken every 2 days throughout the experiment and preserved in 10 % Lugol's solution for microscopic analysis. Estimation of cell density was done microscopically using a Fuchs-Rosenthal counting chamber according to Guillard (1973). In order to obtain reliable results, a minimum of 400 cells were counted to maintain the error within ±10 % (Lund et al. 1958).

Specific growth rate (day^{-1}) was calculated according to the equation:

$$\mu = \mathrm{Ln}(N_t/N_0)/t \tag{1}$$

where μ is the growth rate, N_t is the cell number at time t, and N_0 is the cell number at time zero (right after the inoculation).

Cyanotoxin analyses during culture experiments

Immediately after the addition of the STX+ extract to the cultures, 1 mL aliquots were taken to confirm the concentrations of neo-STX and STX added. In addition, at the end of the experiment, 1 mL aliquots were collected to obtain final neo-STX and STX concentrations. For MC quantification, 1 mL aliquots were taken on days 4 and 14 of the experiment to determine the effect of the different STX+ and STX− treatments on total (intracellular and extracellular) MC production by *M. aeruginosa* BCCUSP232. All samples for cyanotoxin analyses were stored at −80 °C. Just before analyses, the samples were thawed and ultrasonicated for 3 min (15 W and 22.5 kHz) in an ice bath to lyse the cells.

The determination of neo-STX, STX, and total MC concentrations was carried out using Beacon ELISA plate kits (Beacon Analytical Systems, USA) specific to each cyanotoxin. All analyses were done following the manufacturer's instructions. The analyses were performed in triplicate and the results were expressed in µg L^{-1}. STX analyses revealed that neo-STX had the highest degradation (>90 %) at the end of the experiment, while that of STX was lower than 50 % of the initial concentrations in most cases (Table 1).

Statistical analyses

Biomass production and specific growth rate data were first subjected to normality and homoscedasticity tests. Repeated-measures ANOVA and Tukey's tests were carried out to determine significant differences between cell densities and specific growth rates under the different treatment conditions. A correlation-based principal components analysis was used to determine possible relationships between the treatments and the response variables. All analyses were done at 5 % significance level.

Results

The exposure of *M. aeruginosa* and *M. wesenbergii* to STX+ extracts resulted in a significant decrease in the cell density of both strains. The STX− extracts caused a slight reduction in the cell density of both cyanobacterial strains (Fig. 1a–d). Among the green microalgae, only *S. acuminatus* was significantly affected by both STX+ and STX− crude

Table 1 Changes in total saxitoxins and saxitoxin variant concentrations in the beginning and at the end of the experiments with selected cyanobacterial and green microalgal strains. Decay (%) of initial concentration of neo-STX and STX

Target strain	Total STX concentration	Initial concentrations		Final concentrations		Decay (%)	
		Neo-STX	STX	Neo-STX	STX	Neo-STX	STX
M. wesenbergii BCCUSP11	0.5	0.694±0.140	0.185±0.002	0.133±0.030	0.136±0.001	81 %	26 %
	1.0	1.269±0.069	0.716±0.040	0.094±0.011	0.293±0.021	93 %	59 %
	5.0	3.192±0.164	1.093±0.062	0.113±0.011	0.503±0.059	96 %	53 %
	10.0	3.900±0.383	1.886±0.177	0.289±0.024	1.067±0.069	93 %	43 %
M. aeruginosa BCCUSP232	0.5	0.609±0.140	0.173±0.010	0.132±0.009	0.070±0.005	78 %	59 %
	1.0	0.580±0.045	0.338±0.005	0.176±0.047	0.206±0.015	70 %	39 %
	5.0	1.426±0.316	0.956±0.047	0.358±0.026	0.855±0.064	75 %	10 %
	10.0	4.811±0.482	1.950±0.115	0.217±0.043	1.280±0.071	95 %	34 %
S. acuminatus UFSCar036	0.5	0.483±0.069	0.153±0.016	0.111±0.015	0.149±0.010	77 %	2 %
	1.0	0.743±0.015	0.507±0.016	0.106±0.022	0.267±0.007	86 %	47 %
	5.0	2.315±0.162	1.126±0.106	0.122±0.022	0.710±0.075	95 %	36 %
	10.0	4.293±0.328	1.740±0.082	0.255±0.029	1.073±0.013	94 %	38 %
M. convolutum CMEA/UFF0201	0.5	0.406±0.057	0.202±0.011	0.178±0.015	0.147±0.014	56 %	27 %
	1.0	0.652±0.008	0.230±0.025	0.311±0.012	0.216±0.019	52 %	6 %
	5.0	1.693±0.331	1.123±0.069	0.053±0.007	0.101±0.009	97 %	90 %
	10.0	3.329±0.416	1.945±0.022	0.095±0.011	1.191±0.099	97 %	38 %

extracts, while *M. convolutum* showed no sensitivity to the extracts (Fig. 2a–d).

The exposure of the green microalgal and cyanobacterial strains to STX− and STX+ resulted in a significant reduction in their specific growth rates especially at the highest concentration (10 µg L^{-1}) (Fig. 3). Among the four strains investigated, *S. acuminatus* and *M. wesenbergii* were the most sensitive to the STX+ and STX− extracts. However, the toxic effect of STX+ on the investigated strains was higher than that of the STX− extract.

MC concentrations in *M. aeruginosa* BCCUSP232 cultures at the end of the experiment were significantly higher than those observed at the point of STX− and STX+ extract addition (Fig. 4). The higher the STX+ or STX− concentrations, the higher the MC content that was observed.

Principal components analysis (PCA) results of the STX+ and STX− experiments are given in Figs. 5 and 6, respectively. The PCA for STX+ treatments showed that the first two components accounted for over 90 % of the total variation (Fig. 5). Cell density had a significant negative correlation with neo-STX and STX concentrations, while MC production by *M. aeruginosa* had a significant positive correlation with neo-STX and STX concentrations. On the second principal component, a negative relationship of total STX (treatments), neo-STX, and STX concentrations with specific growth rates of the two cyanobacterial and two green microalgal strains was obtained. For the STX− treatments (Fig. 6), MC production by *M. aeruginosa* was significantly positively correlated with the different STX− extract concentrations. Furthermore,

the first two principal components accounted for over 98 % of the total variations.

Discussion

During interspecific competition, the success of phytoplankton species is dependent on their physiological adaptation to changes in the environment or the production of bioactive substances to directly interfere with the growth of competing species (Dunker et al. 2013; Zhang et al. 2013). The effect of these compounds is contingent on their mode of action and the specific particularities of the species or strain under investigation (Perreault et al. 2011; Melegari et al. 2012). Our results indicated potential inhibitory effects of STX− and STX+ extracts on phytoplankton species, when the exposure concentrations were between 5 µg L^{-1} (or 16.7 nM) STX and above. In agreement with our findings, Fistarol et al. (2004) and Hattenrath-Lehmann and Gobler (2011) observed that filtrates of STX- and non-STX-producing strains of *Alexandrium* (dinoflagellate) inhibited the growth of competing species. Our results are slightly different because the STX-containing extracts were more toxic to three out of the four phytoplankton species investigated. Furthermore, Mello et al. (2012) showed that filtrates of STX-producing *C. raciborskii* not only inhibited growth but also retarded colony formation of *M. aeruginosa*, while Melegari et al. (2012) demonstrated that purified STX (0.4–3.0 nM) induced oxidative stress in *C. reinhardtii* after 72 h incubation. Contrary to our findings,
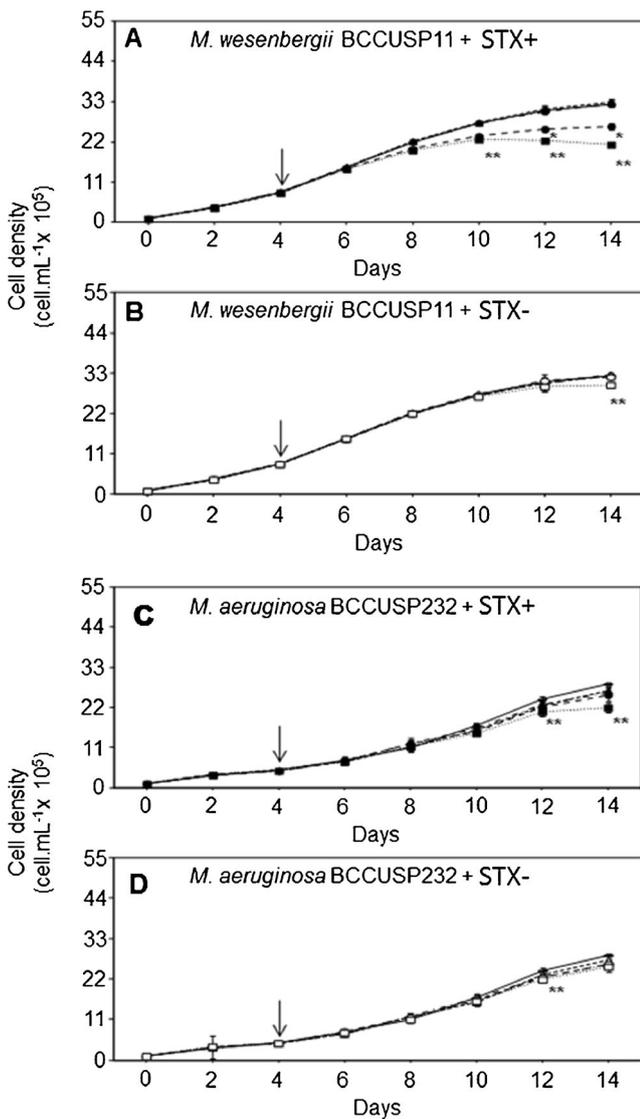
Fig. 1 Cell density of **a** *M. wesenbergii* BCCUSP11 exposed to *C. raciborskii* ITEPA1 STX+. **b** *M. wesenbergii* exposed to *C. raciborskii* ITEP31 STX−. **c** *M. aeruginosa* BCCUSP232 exposed to *C. raciborskii* ITEPA1 STX+. **d** *M. aeruginosa* BCCUSP232 exposed to *C. raciborskii* ITEP31 STX−. Control (*solid line*), 0.5 STX+ μg L^{-1} (*filled diamond* and *dashed line*), 0.5 STX− μg L^{-1} (*empty diamond* and *dashed line*), 1.0 STX+ μg L^{-1} (*filled triangle* and *dashed line*), 1.0 STX− μg L^{-1} (*empty triangle* and *dashed line*), 5.0 STX+ μg L^{-1} (*filled circle* and *dashed line*), 5.0 STX− μg L^{-1} (*empty circle* and *dashed line*), 10.0 STX+ μg L^{-1} (*filled square* and *dotted line*), 10.0 STX − μg L^{-1} (*empty square* and *dotted line*). *Arrows* represent addition of the extracts. Significant differences in cell density between treatments and control 5.0 μg L^{-1} STX+ (*) and 10.0 μg L^{-1} STX+ (**)
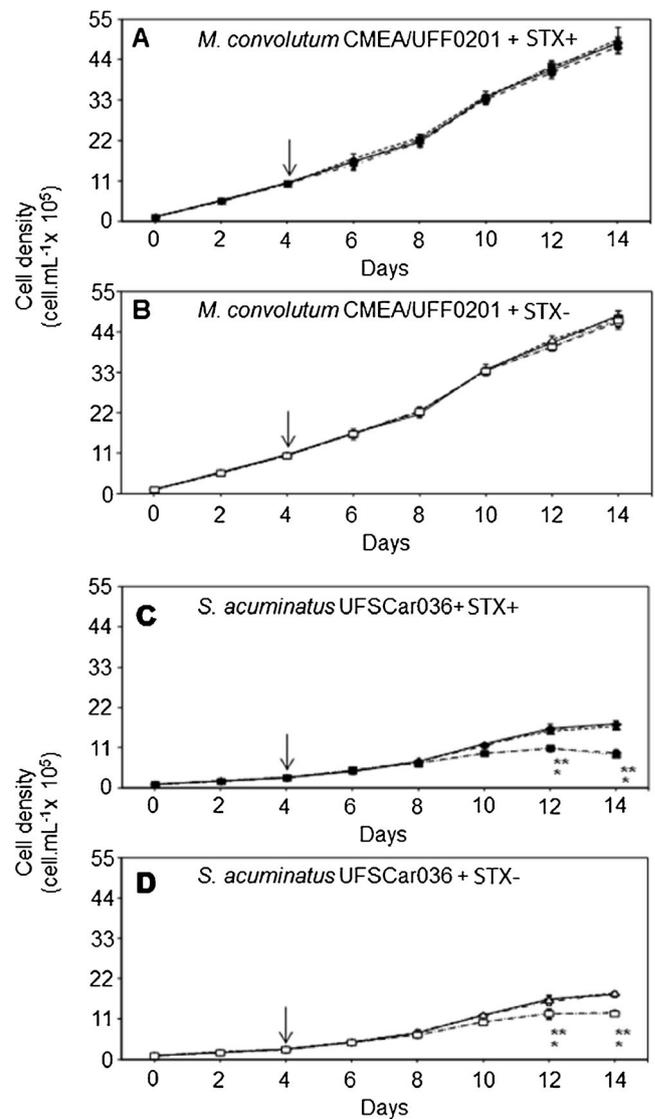
Fig. 2 Cell density of **a** *M. convolutum* CMEA/UFF0201 exposed to *C. raciborskii* ITEPA1 STX+. **b** *M. convolutum* CMEA/UFF0201 exposed to *C. raciborskii* ITEP31 STX−. **c** *S. acuminatus* UFSCar036 exposed to *C. raciborskii* ITEPA1 STX+. **d** *S. acuminatus* UFSCar036 exposed to *C. raciborskii* ITEP31 STX. Control (*solid line*), 0.5 STX+ μg L^{-1} (*filled diamond* and *dashed line*), 0.5 STX− μg L^{-1} (*empty diamond* and *dashed line*), 1.0 STX+ μg L^{-1} (*filled triangle* and *dashed line*), 1.0 STX− μg L^{-1} (*empty triangle* and *dashed line*), 5.0 STX+ μg L^{-1} (*filled circle* and *dashed line*), 5.0 STX− μg L^{-1} (*empty circle* and *dashed line*), 10.0 STX+ μg L^{-1} (*filled square* and *dotted line*), 10.0 STX− μg L^{-1} (*empty square* and *dotted line*). *Arrows* represent addition of the extracts. Significant differences in cell density between treatments and control 5.0 μg L^{-1} STX+ (*) and 10.0 μg L^{-1} STX+ (**)

Perreault et al. (2011) reported no inhibitory effect of purified STX (2–128 nM) on *C. reinhardtii* after 96 h exposure. It is important to note that this difference may be because in addition to STX, there were other secondary metabolites present in the extracts used in our study, which probably interacted with STX to influence their effect on the phytoplankton species.

M. convolutum revealed some tolerance to both STX− and STX+ extracts. Unlike the other phytoplankton species investigated, the cell density and specific growth rates of *M. convolutum* CMEA/UFF0201 were not significantly affected by STX+ and STX− extracts. Recent studies have demonstrated that green microalgae appear to be resistant to a number of cyanobacterial toxins at environmentally relevant

Fig. 3 Specific growth of
S. acuminatus, *M. convolutum*,
M. wesenbergii, and
M. aeruginosa exposed to
extracts of *C. raciborskii* ITEPA1
STX+ and *C. raciborskii* ITEP31
STX−. *Sac* represents
S. acuminatus; *Mono*,
M. convolutum; *M232*,
M. aeruginosa; *M11*,
M. wesenbergii. *$p<0.05$, means
are significantly different from the
control
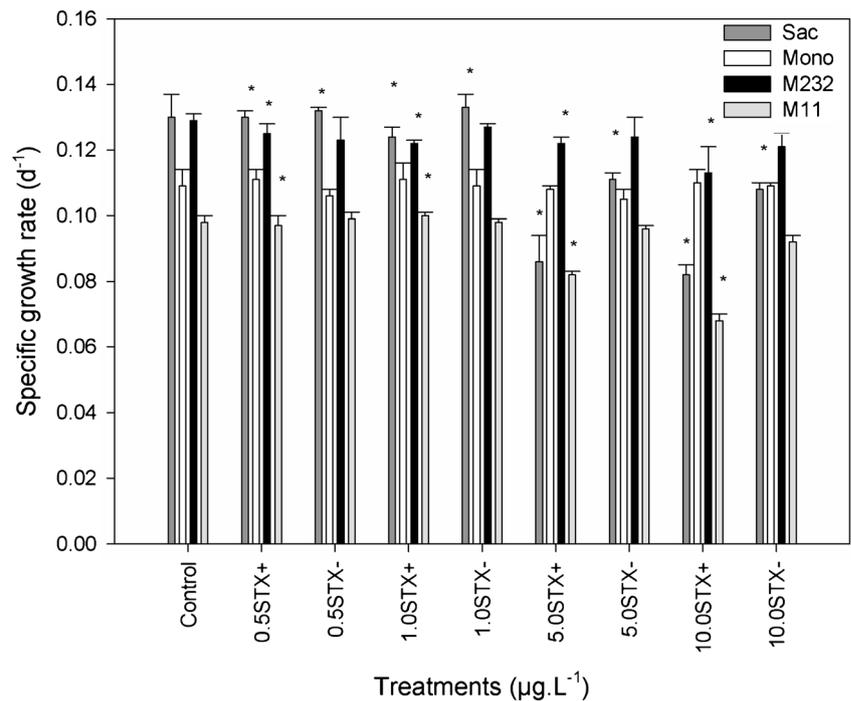
concentrations (Babica et al. 2007; Bártová et al. 2011; Campos et al. 2013; Bittencourt-Oliveira et al. 2015). In addition, Leão et al. (2009) reported that cell-free filtrates of "nontoxic" *C. raciborskii* had no significant effect on *Ankistrodesmus falcatus* and *Chlorella vulgaris*.

Non-STX-containing extracts of *C. raciborskii* also inhibited the growth of some of the investigated phytoplankton species.

Different "nontoxic" strains of this cyanobacterium have been reported to inhibit the growth of phytoplankton species (Suikkanen et al. 2004; Figueredo et al. 2007; Mello et al. 2012). However, there is growing evidence that the "nontoxic" strains of *C. raciborskii* may actually produce toxic (or bioactive) but unknown compounds with allelopathic potential (Acs et al. 2013; Rzymski and Poniedzialek 2014; Poniedzialek et al.

Fig. 4 Total microcystin
production by *Microcystis
aeruginosa* after exposure to
saxitoxin (STX+) and
non-saxitoxin (STX−) extracts of
Cylindrospermopsis raciborskii.
*$p<0.05$, means are significantly
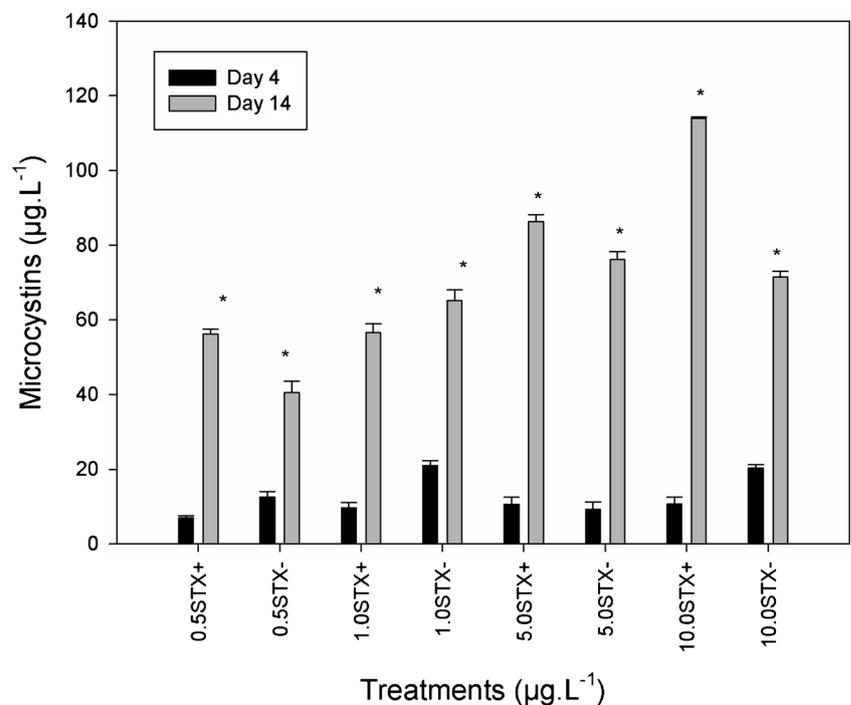different from the control

Fig. 5 PCA biplot showing the relationship of saxitoxin variants (neo-STX and STX) with cell density, microcystin production, and specific growth rates of the different phytoplankton species. *MC232* represents microcystin production by *M. aeruginosa*; *S. acu*, *S. acuminatus* cell density; *Mono*, *M. convolutum* cell density; *M232*, *M. aeruginosa* cell density; *M11*, *M. wesenbergii* cell density. *SGR* represents specific growth rates for S. acu (*S. acuminatus*), Mono (*M. convolutum*), M232 (*M. aeruginosa*), and M11 (*M. wesenbergii*)
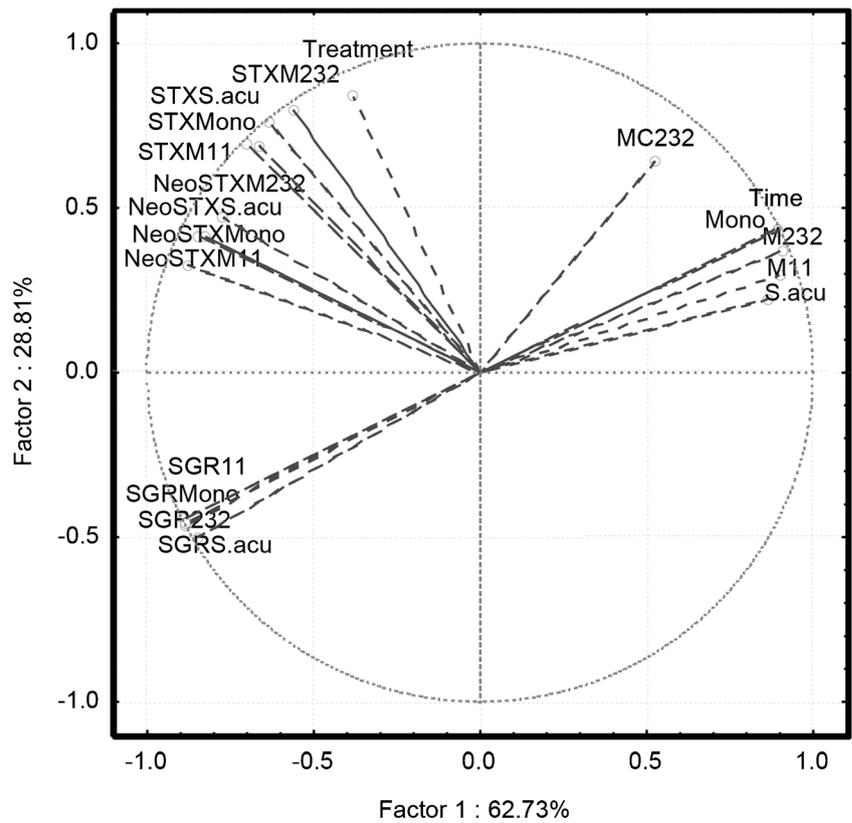

Fig. 6 PCA biplot showing the relationship of non-saxitoxin extracts (STX−) with cell density, microcystin production, and specific growth rates of the different phytoplankton species. *MC232* represents microcystin production by *M. aeruginosa*; *S. acu*, *S. acuminatus* cell density; *Mono*, *M. convolutum* cell density; *M232*, *M. aeruginosa* cell density; *M11*, *M. wesenbergii* cell density. *SGR* represents specific growth rates for S. acu (*S. acuminatus*), Mono (*M. convolutum*), M232 (*M. aeruginosa*), and M11 (*M. wesenbergii*)
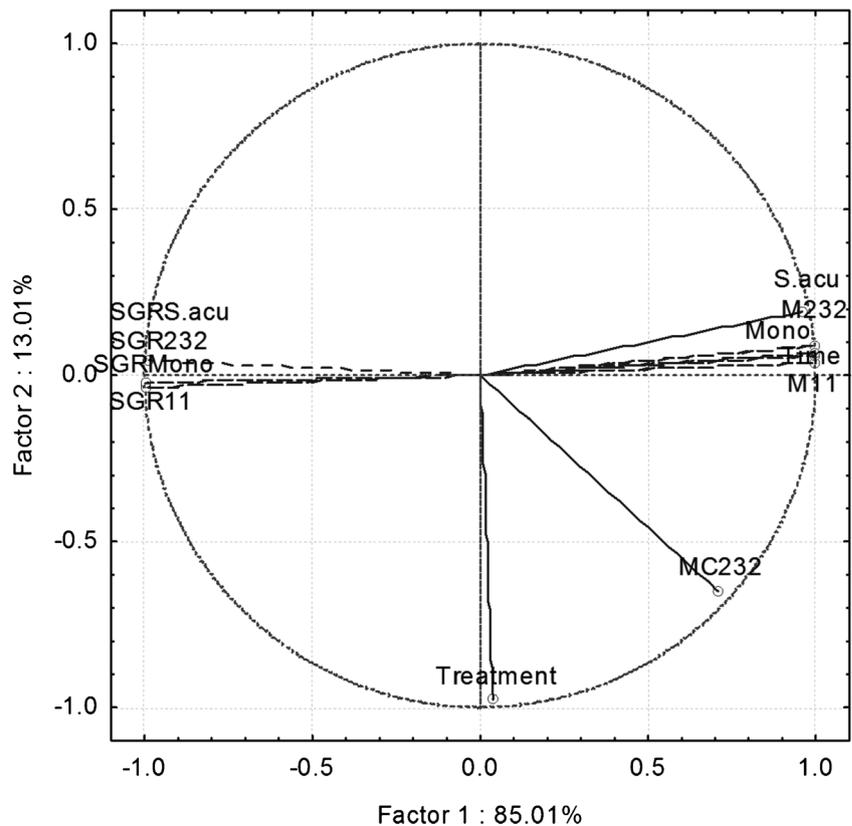
2015). This probably explains why the STX− strain was able to inhibit the growth of the phytoplankton species investigated in our study. However, the inhibition of phytoplankton growth by STX− extracts in our study was lower than that of the STX+ extracts. The greater inhibition by STX+ suggests that there is a STX concentration-dependent toxicity with respect to cell density and specific growth rates. This supports the significant negative correlation found between the concentrations of the different STX variants (neo-STX and STX) with specific growth rate and cell densities of the sensitive phytoplankton strains. Thus, the effects of STX+ extract on the phytoplankton species may be linked to a decrease in the number of cell divisions (Hattenrath-Lehmann and Gobler 2011).

Recent studies have demonstrated that MC are responsible for the formation and maintenance of *M. aeruginosa* cell colonies (Schatz et al. 2007; Gan et al. 2012; Rzymski et al. 2014). Increased synthesis of MC has been reported to correlate with increased growth rate in *M. aeruginosa*, which promotes its competitiveness with other phytoplankton (Orr and Jones 1998; Kaplan et al. 2012) and defense against predation (Pineda-Mendoza et al. 2014). An inhibition of MC synthesis can lead to a decrease in the fitness of *M. aeruginosa* population. In this study, the exposure of *M. aeruginosa* BCCUSP232 to exogenous STX caused an increase in the production of MC. This explains the significant correlation observed between MC production by *M. aeruginosa* and the concentrations of the different STX variants observed in this study. The increased MC production observed in the present study may have been an adaptive strategy by the cyanobacterium to maintain and possibly increase its growth, under the influence of STX+ and STX− extracts. Another interesting phenomenon was observed with the STX− extracts, where the treatments only had a significant positive correlation with the production of MC by *M. aeruginosa* but not with its specific growth rates and cell density. Our previous study (Cordeiro-Araújo and Bittencourt-Oliveira 2013) showed that the active release of MC by *M. aeruginosa* BCCUSP232 was controlled by an endogenous rhythm, which explains the increased concentration of the toxin recorded at the end of the present investigation in all treatments and control. This means the active release of MC by *M. aeruginosa* was stimulated by the presence of bioactive substances produced by both strains of *C. raciborskii*.

It is important to note that in vitro studies cannot be directly extrapolated to field conditions. However, the STX concentrations used in the present study are comparable to those obtained under field conditions (Velzeboer et al. 2000). In addition, the effect of MC on STX-producing and non-STX-producing species remains unknown at the moment, which will form an important subject for future investigations. This is because *C. raciborskii* co-occurs or alternates with the studied green algae and cyanobacteria species in aquatic ecosystems (Moustaka-Gouni et al. 2007). Therefore, there may be several direct interference interactions taking place between these phytoplankton species that are not known at the moment.

In conclusion, our experiments showed that STX− and STX+ crude extracts of *C. raciborskii* were capable of inhibiting the growth of different phytoplankton species. This study provided for the first time a direct correlation of the presence and concentrations of STX variants with biomass production and specific growth rates of three out of the four phytoplankton species investigated. Furthermore, the STX+ extracts were more toxic to the phytoplankton species than the STX− extracts. In response to the presence of STX+ and STX− extracts, total (intracellular and extracellular) MC production by *M. aeruginosa* increased as a means maintaining its growth and probably combating the effects of *C. raciborskii* extracts.

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