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Uptake and elimination of brevetoxin in the invasive green mussel, *Perna viridis*, during natural *Karenia brevis* blooms in southwest Florida

Katherine McFarland^{a,b} kmcfarland@fgcu.edu, Fred Jean^b fjean@univ-brest.fr, Philippe Soudant^b philippe.soudant@univ-brest.fr, Aswani K. Volety^{a,c} voletya@uncw.edu

^a Department of Marine and Ecological Sciences, Florida Gulf Coast University, 10501 FGCU Blvd. South, Fort Myers, FL 33965, USA

^b LEMAR UMR 6539, Institute Universitaire Européen de la Mer, Université de Bretagne Occidentale, Rue Dumont d'Urville, Place Nicolas Copernic, 29280 Plouzané, France

^c Department of Biology and Marine Biology, University of North Carolina Wilmington, 601 South College Rd., Wilmington, NC, 28403, USA (current address)

Correspondence: Katherine McFarland; email: mcfarland.316@gmail.com; phone: 440-567-7335

Abstract

Perna viridis is a recently introduced species to US coastal waters and have vigorously spread throughout the southeastern seaboard since their invasion. Little information regarding their response to local environmental factors has been reported including responses to the local HAB species, *Karenia brevis*. This study monitored the tissue toxin concentration of brevetoxins in *P. viridis* from existing populations throughout two consecutive natural *K. brevis* blooms. The results showed *P. viridis* to rapidly accumulate PbTx upon exposure to the bloom, far exceeding the peak tissue concentrations of oysters, *Crassostrea virginica*, sampled during the same period, $57,653 \pm 15,937$ and $33,462 \pm 10,391$ ng g⁻¹ PbTx-3 equivalent respectively. Further, *P. viridis* retained high PbTx concentrations in their tissues post bloom remaining above the regulatory limit for human consumption for 4 – 5 months, significantly longer than the depuration time of 2 – 8 weeks for native oyster and clam species. In the second year, the bloom persisted at high cell concentrations resulting in prolonged exposure and higher PbTx tissue concentrations indicating increased bioaccumulation in green mussels. While this species is not currently harvested for

human consumption, the threat for post bloom trophic transfer could pose negative impacts on other important fisheries and higher food web implications.

Key Words: red tide, oyster, bioaccumulation, trophic transfer, harmful algal bloom, depuration

Abbreviations: PbTx, brevetoxin; HAB, harmful algal bloom; STX, saxitoxin; NSP, neurotoxic shellfish poisoning

1. Introduction

Red tide events occur regularly in the Gulf of Mexico, with the first recorded event in 1844 (Lasker and Smith, 1954) and identified as *Karenia brevis* in 1946 (Davis, 1948). This unarmored dinoflagellate produces a suite of neurotoxins known as brevetoxins (PbTx), which cause massive fish kills, marine mammal and seabird mortalities and neurotoxic shellfish poisoning (NSP) in humans through consumption of toxic shellfish (Ray and Wilson, 1957; McFarren et al. 1965; Adams et al., 1968; Forrester et al., 1977; Baden, 1989; Stiedinger, 1973; Flewelling et al. 2005; Naar et al., 2007; Landsberg et al., 2009). Over the past several decades outbreaks of *K. brevis* blooms have increased in frequency and duration due to eutrophication of coastal waters through anthropogenic input (Brand and Compton, 2007).

Bivalve molluscs bioaccumulate PbTx through normal filter feeding behavior with little to no mortality and, as a consequence, pose a threat to both human consumers and natural marine predators including whelks, crab, and fish (Tester et al., 2000; Brand et al., 2012). Several studies have demonstrated the transfer of PbTx from bivalves to whelks and fish (Ingham et al., 1986; Peirce et al., 2002; Naar et al., 2007; Bricelj et al., 2012). Shellfish beds are monitored regularly for both *K. brevis* cell counts and tissue toxin concentration, particularly during and

following bloom events. Closure of shellfish harvesting is required if *K. brevis* cell counts exceed 5,000 cells L⁻¹ or shellfish tissue concentrations exceed 800 ng g⁻¹ PbTx-3 equivalent brevetoxin (Plakas et al., 2008).

The green mussel, *Perna viridis*, is a recently introduced species to southwest Florida. The extent to which PbTx accumulates and persists in their tissues remains to be elucidated and is not monitored. If PbTx remains sequestered in tissues after bloom dissipation this poses a threat to predators including commercially important crab and fish, subsequently transferring toxins through the trophic web even long after a red tide event has subsided. Stomach content analyses in post bloom fish and dolphin mortalities have revealed fish with high levels of PbTx (Flewelling et al., 2005) and seabirds with toxic clams in the gut (Forrester et al., 1977) indicating a lethal dose through trophic transfer (Landsberg et al., 2009). Likewise, dead burrfish (*Chilomycterus schoepfi*) collected during a red tide bloom had the remains of toxic bivalves in their stomachs (Naar et al., 2007). While green mussels are not harvested locally for human consumption, they are an edible species and may pose potential human health concerns.

Accumulation of PbTx has been reported in the green lipped mussel, *Perna canaliculus*, during exposure to *Karenia selliformis* blooms (Morohashi et al., 1995; Ishida et al., 2004a, 2004b), however no published information directly addresses the accumulation of PbTx from *K. brevis* blooms in *P. viridis*. Therefore, the goal of this study was to assess the rate of PbTx uptake and elimination in *P. viridis* during a naturally occurring *K. brevis* bloom. This was accomplished through monthly collections from a population, which had become established in southwest Florida, before, during and after a red tide event in Estero Bay, Florida over a two year study period.

2. Material and Methods

2.1 Collection of *Perna viridis*

Perna viridis of an average length of 74.3 ± 7.3 mm were collected by SCUBA diving from Estero Bay, Florida at New Pass Bridge (26°22'40.89"N; 81°51'39.69"W) or Big Carlos Bridge (26°24'15.44"N; 81°52'49.64"W) once a month from October 2011 through October 2013 (N = 5 / month). Immediately following collections mussels were cleaned of epiphytic growth, soft tissue was dissected from the shell and individually frozen at -80°C. Whole individuals were homogenized cryogenically to a powder using a MixerMill 400 (Retsch® Solutions in Milling and Sieving, Hann, Germany) under liquid nitrogen and one gram wet tissue was weighed for PbTx extraction. For samples from October 2011 through December 2012 archived freeze-dried tissue was used following the same extraction method using wet to dry weight ratios to determine weight for each individual (Dr. Leanne Flewelling, personal communication).

2.2 Bloom formation

Karina brevis cell counts were obtained from Florida Wildlife Research Institute monitoring program. More frequent sampling was conducted in the Sanibel / Captiva area than Estero Bay due to the presence of several commercial shellfish beds. Because it is located less than 30 km north of the study site, monitoring data from Sanibel / Captiva was used to supplement data from the study site.

Onset of the bloom in local waters first occurred in the Sanibel / Captiva area in September 2011 and was observed in Estero Bay (study site) in November 2011 accompanied by a large fish kill consisting primarily of mullet, *Mugil cephalus*. The bloom persisted into January 2012, but cell counts diminished by the end of the month. This cycle was observed again in

2012-2013 with an increase in cell concentrations in the Sanibel / Captiva area in late September 2012 and Estero Bay by mid-October 2012. The boom persisted into February 2013 at which time a fish kill comprising several species was observed on Ft. Myers Beach. The bloom peaked in February and diminished by the end of April 2013.

2.3 Analysis of PbTx in tissues

All samples were extracted in methanol according to Naar et al. (2002); briefly the protocol is as follows. One gram homogenized wet tissue (or its equivalent in dry tissue) was vortexed well in 80% methanol, heated in a water bath at 60°C for 20 minutes, then transferred to an ice bath for 10 minutes and centrifuged. The supernatant was poured off and stored on ice and these steps repeated on the remaining precipitate. The combined supernatants were brought to a total volume of 10 mL 80% methanol, washed with hexane and stored at -20°C until further analysis.

Brevetoxin ELISA assays were run within seven days of the tissue extraction using a competitive ELISA kit prepared at the University of North Carolina Wilmington according to Griffith et al. (2013). The assay was completed in a 96-well microplate (Nunc-Immunoplate with Maxisorp surface, Thermo Scientific™, Waltham, Massachusetts). The plate was first coated with 100µL PbTx-3-bovine serum albumin conjugate and incubated on a plate shaker for one hour. The plate was then washed with Phosphate Buffered Saline (PBS) followed by adding a blocking buffer (Superblock™ dry blend) (200µL) and incubated for 30 minutes on a plate shaker. Following the incubation period the plate was washed with PBS-Tween™ (buffered detergent). Samples and standards were loaded (100µL) onto the plate in duplicates with the addition of a primary antibody (100µL) and incubated on a plate shaker for 1 hour. The plate was

washed again with PBS-TweenTM, then a secondary antibody was loaded into the wells (100 μ L) and again incubated for one hour. Following this incubation the plate was washed with PBS-TweenTM and a final PBS wash. Tetramethylbenzidine (TMB) was added as an indicator and the reaction stopped after approximately three minutes using 0.5 M H₂SO₄. The absorbance was measured at 450 nm on a TECAN Genois Pro® plate reader (TECAN Group Ltd., Männedorf, Switzerland). The antibody used detects type-2 brevetoxins which make up 90 – 95% of the toxins produced during blooms and include PbTx-2, PbTx-3 and PbTx-9 (Naar et al., 2002). Results are presented as PbTx-3 equivalent to reflect the standard curve used in calculations of PbTx concentrations.

2.4 Statistical Analysis

To assess PbTx concentrations throughout each bloom, paired t-tests were performed to assess differences across time within each collection group. Due to non-normality of the data, the intensity of post-bloom tissue PbTx accumulation was assessed through integrated trapezoidal area-under-the-curves regarding increase from baseline (AUC_i) according to the previously published formula by Pruessner et al. (2003). AUC_i was calculated each month separately to compare cumulative increase in PbTx in relation to onset of the bloom between years. The differences in the intensity of PbTx accumulation (AUC_i) during corresponding months following *K. brevis* blooms (2012 and 2013) were evaluated using independent t-tests. All data are presented as means \pm S.D. with statistical significance being defined as $p \leq 0.05$.

3. Results

Green mussels accumulated high levels of PbTx in their tissues during a natural exposure to *K. brevis*. Tissue PbTx concentrations reached a peak in February 2013 (137,000 \pm 40,000 ng

g⁻¹ PbTx-3 equivalent) following a bloom phase which lasted approximately four months. Peak tissue PbTx concentrations corresponded with peaks in cell counts and duration of bloom (Fig. 1). Tissue toxin concentration remained above the regulatory limit (800 ng g⁻¹ PbTx-3 equivalent) four to five months post bloom dissipation (Fig. 1A).

The second year of exposure showed a higher peak in toxin concentration compared to the first year (Fig. 1A), coinciding with a more intense and prolonged bloom during the second year (Fig. 1B). Intensity of PbTx accumulation (AUCi) was significantly different between the two bloom years ($p \leq 0.001$) in which a slower accumulation of PbTx was observed during the second bloom with a peak in PbTx tissue concentrations at the end of the bloom after four months of exposure (Fig. 2). During both bloom years, mussels eliminated PbTx while *K. brevis* was present in the water column, during the waning phase of the bloom in late January 2012 and March – April 2013 (Fig. 1), however this process was slow; PbTx concentration in tissue remained above the regulatory limit until September 2013 and never fell below regulatory limit in 2012.

In March 2012, a population-wide mass mortality event was observed in the green mussels that decimated the population, leaving only sporadic individuals. During the time of the mortality event the average PbTx tissue concentration in the green mussels was $15,694 \pm 3,885$ ng g⁻¹ PbTx-3 equivalent. This mortality event made the collection of adults impossible in April and July 2012 and difficult throughout the summer months. Smaller mortality events continued to be observed throughout the remainder of the monitoring period with the highest mortality observed in the fall of 2012 and 2013 with the return of bloom status in *K. brevis*.

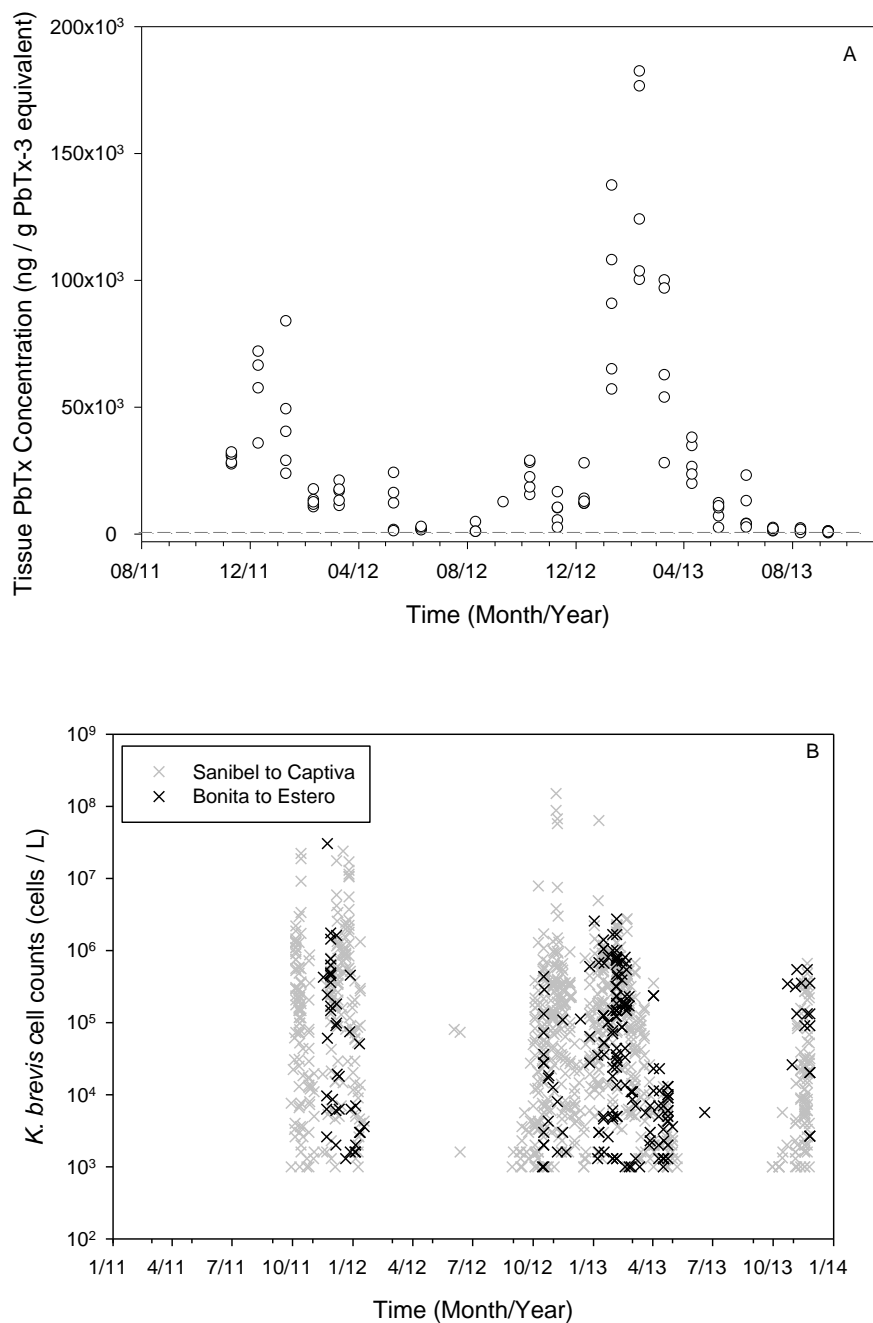


Figure 1: Brevetoxin concentration in individual mussels plotted over time (A). Dashed line indicates the regulatory limit (800 ng g⁻¹ PbTx-3 equivalent). *Karenia brevis* cell counts (log scale) by location over time (B).

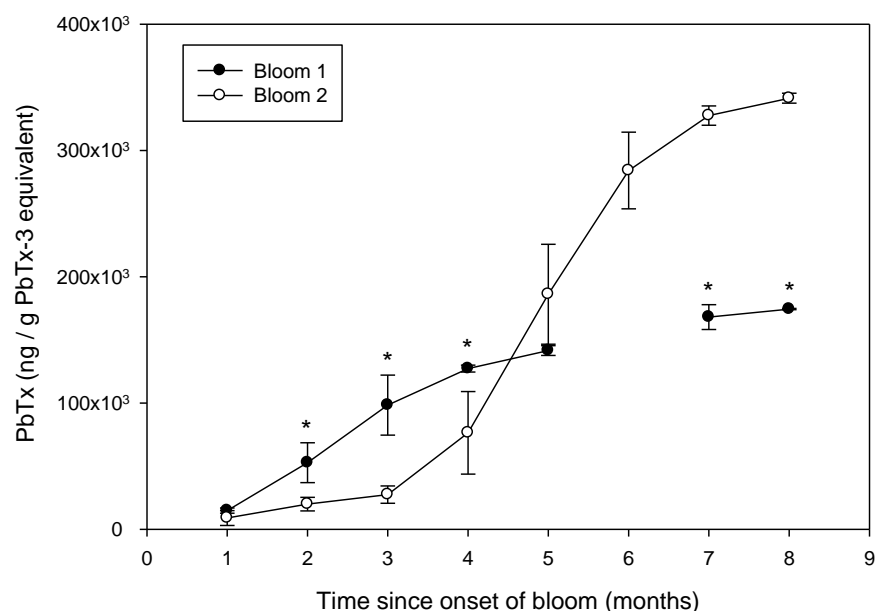


Figure 2: Relative concentrations of PbTx in green mussel tissues in relation to onset of the bloom for comparison between the two bloom cycles as determined by AUCi analyses. Error bars represent standard deviation and * indicate significant differences between blooms.

Table 1: Southwest Florida bivalve species and the reported time required to depurate PbTx to levels below the regulatory (800 ng g^{-1} PbTx-3 equivalent) limit following exposure to *K. brevis*.

Species	Depuration Time	Exposure	Source
<i>Crassostrea virginica</i>	2 weeks	lab	Plakas et al., 2002; Griffith et al., 2013
<i>Crassostrea virginica</i>	2 weeks	field	Plakas et al., 2008
<i>Crassostrea virginica</i>	2 - 8 weeks	field	Morton and Burklew, 1969
<i>Crassostrea virginica</i>	10 weeks	field	Dickey et al., 1999
<i>Crassostrea virginica</i>	6 weeks	field	Pierce et al., 2002
<i>Mercenaria mercenaria</i>	6 weeks	field	Pierce et al., 2002
<i>Mercenaria mercenaria</i>	2 weeks	lab	Griffith et al., 2013
<i>Perna viridis</i>	16 - 20 weeks	field	This Study

Table 2: Comparison of green mussel and local oyster average tissue brevetoxin concentrations during the 2011 – 2012 *K. brevis* bloom. PbTx concentrations expressed as ng g⁻¹ PbTx-3 equivalent.

Month	<i>Perna viridis</i> (this study)		<i>Crassostrea virginica</i> (Volety et al., unpublished results)		Red Tide Status (FWRI)
	PbTx	SD	PbTx	SD	
Oct. 2011	N.D.				
Nov. 2011	29,696	2,051			bloom
Dec. 2011	57,653	15,937	33,462	10,391	bloom
Jan. 2012	44,982	23,786	7,583	2,516	waning
Feb. 2012	12,913	2,709	6,406	1,010	gone
Mar. 2012	15,694	3,885	4,159	309	
April 2012	no sampling *		810	317	
May 2012	10,806	9,815	no sampling		

N.D. indicates that brevetoxin levels were below the detection limit

*April collections could not be completed for *P. viridis* due to a mass mortality event in March 2012 severely reducing the population in addition to poor visibility during SCUBA diving making collections impossible

4. Discussion

Through monthly monitoring of PbTx concentrations in the soft tissue of *P. viridis* established in southwest Florida before, during and after two consecutive *K. brevis* blooms, we were able to characterize the uptake and elimination under natural conditions. The data presented here demonstrate that *P. viridis* rapidly accumulated large quantities of PbTx in their tissues and concentrations remained above the regulatory limit (800 ng g⁻¹ PbTx-3 equivalent) for an extended period even after bloom dissipation, suggesting the potential for trophic transfer up to five months post bloom.

PbTx tissue concentrations corresponded well with the *K. brevis* cell counts showing peaks in tissue concentrations during and following peaks in water column cell concentrations, suggesting that rapid uptake likely occurred through filtering of whole *K. brevis* cells. Free PbTx in the water column may have contributed to high tissue concentrations. Water column PbTx concentrations range from 0.4 – 0.12 µg L⁻¹ when cell counts are ≤ 2.4 x 10³ cells L⁻¹ (Pierce et al., 2005) and frequently exceed 24 µg L⁻¹ during bloom concentrations of 10⁶ cells L⁻¹ (Pierce

et al., 2008). Likewise, Rolton et al. (2015) detected PbTx in *K. brevis* culture filtrate to range from 5 – 51 $\mu\text{g L}^{-1}$. The peak in tissue concentration occurred near the end of the bloom indicating continued bioaccumulation of the toxin in mussels, which did not die, leading to increased tissue concentrations through continuous exposure. Tissue PbTx concentrations were higher in the second year and persisted at elevated levels for an extended period due to the prolonged bloom conditions in 2013 versus 2012 as both elevated exposure intensities and duration affect the accumulation and depuration rate (Morton and Burklew, 1969; Bricelj and Shumway, 1998). However, this peak in tissue concentration occurred late in the bloom cycle after four months of exposure suggesting bioaccumulation and an inability to eliminate toxins during prolonged exposure. Previous exposure history is also a contributing factor to the increased toxin levels in 2013 as tissue toxin concentrations never fell below the regulatory limit following the 2011 – 2012 bloom. In June 2012, *K. brevis* cell counts reached bloom concentrations for several days at which time it appears the uptake rate exceeded the elimination rate resulting in a small spike in tissue PbTx concentration, prolonging the elimination period.

While PbTx accumulation and elimination has not previously been studied in *P. viridis*, evidence for metabolism / biotransformation of PbTx has been found in several bivalve species including the closely related New Zealand green-lipped mussel *Perna canaliculus* (Ishida et al., 2004a, 2004b; Ishida et al., 1995; Morohashi et al., 1995; Murata et al., 1998; Nozawa et al., 2003). In both field and controlled laboratory exposures, *P. canaliculus*, showed both an accumulation of PbTx and metabolism from PbTx-2 (most prevalent in the *K. brevis* cell) to PbTx-3 and BTX-5 at rates dependent upon the intensity and duration of exposure (Ishida et al., 2004a, 2004b).

When compared with native Florida bivalves, oysters and clams, *C. virginica* and *M. mercenaria*, *P. viridis* appear to accumulate higher PbTx concentrations and demonstrate slower elimination rates. These local bivalves have been shown to accumulate PbTx with high survival and rapid depuration rates of 2 – 8 weeks (Table 1) (Morton and Burklew, 1969; Steidinger and Ingle, 1972; Plakas et al., 2002; Plakas et al., 2004; Plakas et al., 2008; Bricelj et al., 2012; Griffith et al., 2013). *Crassostrea virginica* sampled from Pine Island Sound, just 30 km north of Estero Bay, during the bloom of 2011 – 2012 were shown to reach a peak in PbTx levels averaging approximately 35,000 ng g⁻¹ PbTx-3 equivalent but exhibited a rapid initial elimination rate, decreasing to 7,500 ng g⁻¹ PbTx-3 equivalent just two weeks after bloom dissipation and dropped below the regulatory limit within 2 ½ months (Volety et al., unpublished results). *Perna viridis* sampled from Estero Bay during this period exhibited nearly double (~58,000 ng g⁻¹ PbTx-3 equivalent) the PbTx concentrations of *C. virginica* (Table 2) and did not fall below the regulatory limit before the return of bloom conditions in October 2012 and a second exposure period.

Field monitoring of bivalves during red tide events in Tampa Bay, Florida have reported tissue concentrations in *C. virginica* ranging from 2,800 – 16,500 ng g⁻¹ (Pierce et al., 2002; Plakas et al., 2008) to as high as 28,660 – 80,000 ng g⁻¹ (Dickey et al., 1999; Weidner et al., 2002) and *M. mercenaria* ranging from 1,800 – 6,600 ng g⁻¹ (Poli et al., 2000; Pierce et al., 2002; Weidner et al., 2002). In controlled exposures, Griffith et al. (2013) found *C. virginica* and *M. mercenaria* to reach 2,000 and 1,000 ng g⁻¹, respectively after eight days of exposure with acute daily dosing of 5 x 10⁵ cells L⁻¹, whereas Plakas et al. (2004) reported tissue concentrations in *C. virginica* of 740 ng g⁻¹ after only two doses of 10⁴ cells L⁻¹ over 48 hours. Concentrations observed in *P. viridis* tissues are higher than those observed in local species. This is likely due to

a combination of several factors including differences in metabolic activity (uptake, ingestion, and assimilation rates), growth rates, energy turnover, and / or reduced mechanism for “detoxifying” / metabolizing PbTx which could lead to increased tissue damage and physiological impairment. Additionally, PbTx’s are lipophilic, thus both lipid composition and lipid turnover rate will play a role in the accumulation and elimination rates and will vary between species and within species at different times of the year or reproductive cycle (Svensson and Förlin, 2004)

Previous work has shown several mussel species to rapidly accumulate high concentrations of toxins produced by HAB’s compared to many oyster and clam species (Ingham et al., 1986; Shumway and Cucci, 1987; Shumway et al., 1988, 1990, 1995; Bricelj and Shumway, 1998; Lesser and Shumway, 1993). In the Philippines, *P. viridis* was found to be highly toxic with low survival following a *Pyrodinium bahamense* var. *compressa* red tide event in 1983 reaching 9,620 MU 100 g⁻¹ saxitoxin (STX) (~18,000 ng g⁻¹) leading to an eight month shellfish ban due to prolonged toxicity (Gacutan et al., 1984) and in more recent blooms, *P. viridis* reached concentrations as high as 90,000 ng g⁻¹ STX, far exceeding the regulatory limit while other local bivalves showed minimal toxicity (Montejo et al., 2010, 2012). Jaafar et al. (1989) reported high tissue STX concentrations in *P. viridis* during a *P. bahamense* bloom in Malaysia reaching 50, 354 ng g⁻¹, but contrary to blooms in the Philippines showed rapid elimination rates.

Species specific metabolism of PbTx has been reported widely throughout the literature (Cummins et al., 1971; Steidinger et al., 1998; Nozawa et al., 2003; Ishida et al., 2004a, 2004b; Abraham et al., 2012; Echevarria et al., 2012) offering a likely explanation to observable differences between *P. viridis* in this study and those reported for *C. virginica* and *M.*

mercenaria. Indeed, Ishida et al. (2004b) found a significant difference in PbTx metabolism in *P. canaliculus* and *Crassostrea gigas*, two similar species to those compared in this study, with different concentrations of metabolites found in each indicating similar pathways, but different reaction rates. *Crassostrea gigas* showed a more rapid transformation of PbTx-2 to PbTx-3 compared to *P. canaliculus*. While both species had PbTx-3, BTX-B1 and BTX-B5, *P. canaliculus* also produced BTX-B2, BTX-B3 and BTX-B4, but *C. gigas* did not (Ishida et al., 2004b; Morohashi et al., 1999; Nozawa et al., 2003). Concentrations of metabolites present may effect tissue retention time between species as different derivatives may be stored differently and in various tissue compartments, thus effecting elimination rates (Echevarria et al., 2012). Detoxification is partially dependent upon the peak tissue concentration (Bricelj and Shumway, 1998) and previous exposure history can affect the uptake kinetics of biotoxins (Shumway and Cucci, 1987). Feeding behavior and metabolic demands play a significant role in differences in accumulation rates between species. *Perna viridis* has been shown to have high clearance rates compared to *C. virginica* (McFarland et al., 2013) suggesting increased exposure due to increased filtration of *K. brevis* cells. Additionally, *P. viridis* has high growth and reproductive rates leading to increased metabolic demands (Rajagopal et al., 2006; Siddall, 1980; Vakily, 1989) and increased nutrient requirements which may increase filtration behavior causing increased exposure to the toxin.

As a recently introduced species, *P. viridis* may lack the needed adaptations for rapid PbTx elimination observed in *C. virginica* and *M. mercenaria*, which have a long history of *K. brevis* exposure during which PbTx tolerance and metabolism may have developed, existing in coastal waters of the eastern US seaboard since the 1800's (NOAA, 2005; Volety et al., 2014).

Previously exposed bivalves have shown increased resistance and reduced sensitivity to HAB

exposure compared to unexposed populations (Shumway and Gainey, 1992). While *P. viridis* experiences harmful algal blooms in their native range (Gacutan et al., 1984, 1985; Choi et al., 2003; Li et al., 2005; Montojo et al., 2012), exposure to a new toxin may cause reduced physiological function and could cause the widespread mortality observed in southwest Florida. Leverone et al. (2007) observed decreased clearance rates in juvenile *P. viridis* during short term exposure to *K. brevis* indicating reduced feeding, which may turn lethal if conditions persist. Baker et al. (2012) documented a population-wide mortality event in Tampa Bay following a red tide bloom from which green mussel populations never fully recovered and Gacutan et al. (1984) reported >90% *P. viridis* mortality following a red tide event caused by *P. bahamense* in the Philippines. In the current study, a similar mortality event was observed two months post bloom dissipation, however tissue toxin concentrations were still relatively high (15,700 ng g⁻¹ PbTx-3 equivalent). The lag time between peak tissue PbTx concentration and the first mortality event indicates that other factors are likely involved. Indeed, it coincided with a major spawning period which may have inflicted additional stress on the animal. Galimany et al. (2008) showed a decreased ability to resume normal tissue function, including paralysis of the adductor muscle and prolonged histopathology in *Mytilus edulis* which had spawned during a laboratory exposure to *Alexandrium fundyense*. While the mortality events observed in this study cannot be directly related to *K. brevis* exposure, *P. viridis* had high levels of accumulated toxins in their tissues during the observed mortality indicating the potential for prolonged cumulative effects of high tissue toxin burden and physiological stress. Allelopathic, hemolytic, and other ichthyotoxic compounds produced by *K. brevis* may also contribute to the negative effects observed in marine organisms (Kubanek et al., 2005; Marshall et al., 2005). While prolonged elevated tissue toxin

levels are of concern for trophic transfer, if the mussels die the risk is reduced and invasive species kept at bay.

5. Conclusion

To date, the green mussel populations have not recovered from mortalities observed during red tide events in Estero or Tampa Bay, Florida (Baker et al., 2012; this study). However, not all green mussels died during *K. brevis* exposure as collections of adults were made in the months following the mortality event. Thus, seemingly normal mussels may contain high levels of PbTx for long periods post bloom posing a threat to both natural marine predators and human consumption. Landsberg et al. (2009) attributed post bloom mortality in several fish species and blue crab to trophic transfer from feeding on benthic fauna and post bloom dolphin mortalities in 2004 to trophic transfer from toxic fish. Brand and Compton (2007) noted an increase in the presence of sublethal or background concentrations of *K. brevis* which may be contributing to prolonged tissue toxins post bloom and several studies have noted the persistence of PbTx in marine sediments and seagrass epiphytes (Shumway and Cucci, 1987; Mendoza et al., 2008). Additionally, whole cell and temporary encystment of toxic dinoflagellates have been observed in bivalve biodeposits following exposure leading to increased risk for transport and increased exposure to benthic organisms via deposition into the sediments (Hégaret et al., 2008). Even at low tissue concentrations, this persistence allows for long term exposure to higher trophic levels through biomagnification post bloom (Rounsefell and Nelson, 1966; Landsberg, 2002; Flewelling et al., 2005). Animals that prey directly on benthic filter feeders including gastropods, crab, fish and water birds are at risk of exposure and may pose a threat to higher trophic levels through trophic transfer of the toxins by themselves becoming prey (Shumway et al., 1995; Pierce et al., 2002; Flewelling et al., 2005; Hégaret et al., 2008; Bricelj et al., 2012).

This study has demonstrated the natural accumulation and depuration rates of local *P. viridis* populations with PbTx concentrations and depuration periods exceeding that of native shellfish in southwest Florida. Further research on the potential trophic transfer from contaminated mussels to commercially important fish and crab is vital to monitoring the bloom and post bloom risks. Additionally, as *P. viridis* is an edible species, both monitoring and public awareness are essential to prevent cases of NSP during post bloom periods in which mussels may still be toxic.

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