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# Accumulation of nine metals and one metalloid in the tropical scallop *Comptopallium radula* from coral reefs in New Caledonia

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This study investigates metal accumulation behaviour in the tropical scallop Comptopallium radula and preliminary risk assessment for consumers.

# Abstract

Uptake of waterborne Cd, Co, Mn and Zn was determined in laboratory experiments using radiotracer techniques (<sup>109</sup>Cd, <sup>57</sup>Co, <sup>54</sup>Mn and <sup>65</sup>Zn). Labelled Zn was mainly accumulated in the digestive gland (65%) and Co in kidneys (81%); Cd and Mn were similarly distributed in digestive gland and gills. In a complementary field study, Ag, As, Cd, Co, Cr, Cu, Fe, Mn, Ni, and Zn were analysed in scallops collected at two stations showing different contamination levels. Digestive gland and kidneys displayed the highest concentrations. Ag, As, Cd, and Fe differed in soft tissues from the two stations, suggesting that *Comptopallium radula* could be a valuable local biomonitor species for these elements. Low Mn and Zn concentrations found in kidneys suggest that their content in calcium-phosphate concretions differs from the other pectinids. Preliminary risk considerations suggest that As would be the only element potentially leading to exposure of concern for seafood consumers.

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# 1. Introduction

The urban and industrial expansion in tropical coastal zones increases the release of contaminants which can constitute a threat to local marine ecosystems and globally affect marine diversity (Peters et al., 1997). The SW lagoon of New Caledonia (South Pacific Ocean) represents a tropical case study as it is subject to large inputs of heavy metals mainly due to intense land-based mining activities (Ambastian et al., 1997). New Caledonia is the third largest producer of nickel (Ni) in the

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world and metal contamination mainly concerns Ni and its mining by-products such as cobalt (Co), chromium (Cr) and manganese (Mn) which occur at elevated concentrations in Ni ores. Furthermore, the city of Noumea produces sewage sludge, which can also lead to metal contamination in the surrounding lagoon. Nevertheless, published information about the contamination status of the New Caledonia marine environment is extremely scarce (Labrosse et al., 2000) and limited to a narrow range of species (Bustamante et al., 2000; Hédouin et al., 2007; Monniot et al., 1994). Environmental studies are therefore needed to understand the behaviour and fate of metals in this area in order to develop a programme of coastal zone monitoring and improve local marine resource management.

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The use of biomonitor species to examine metal contamination is a powerful tool to reveal the bioavailability of the considered contaminants (e.g. Rainbow and Phillips, 1993). In this respect, bivalves have been extensively used as biomonitor species because of their high capacity to bioaccumulate various contaminants (see e.g. Goldberg, 1995; Rainbow, 1995). Among bivalves, scallops have been shown to concentrate various trace elements to a large extent (e.g. Bryan, 1973; Bustamante and Miramand, 2005; Uthe and Chou, 1987), even in remote areas such as Antarctica that are not subject to direct anthropogenic inputs (Berkman and Nigro, 1992; Mauri et al., 1990; Viarengo et al., 1993).

Within the huge bivalve stock in New Caledonia, exhibiting around 1200 species among the whole 8000 bivalve species estimated in the world (Bouchet, 1979), 30 pectinid species have been described to date (Dijkstra et al., 1990). In the SW lagoon, *Comptopallium radula* appears to be relatively common and constitutes, with *Mimachlamys gloriosa* and *Brachtechlamys vexillum* the major species targeted for human consumption. Whereas information on stock, reproduction and taxonomy of these scallops is available (Dijkstra et al., 1990; Lefort, 1994), to the best of our knowledge, no study on the bioaccumulation of trace elements in the soft tissues of New Caledonian scallops has been published to date.

The aim of this study was therefore to examine the bioaccumulation and tissue distribution of a wide range of trace elements in the scallop *C. radula* using both laboratory experiments under controlled conditions and in situ investigations. Firstly, the uptake kinetics of four selected radiotracers (<sup>109</sup>Cd, <sup>57</sup>Co, <sup>54</sup>Mn, and <sup>65</sup>Zn) and their body distribution were determined following seawater exposure. Secondly, one metalloid (As) and nine metals (Ag, Cd, Co, Cr, Cu, Fe, Mn, Ni, and Zn) were analysed in the tissues of scallops from reference and contaminated sites in order to evaluate the bioaccumulation capacities of *C. radula* and its potential as a biomonitor species for trace element contamination. Finally, the human risk link to scallop consumption is discussed.

## 2. Material and methods

## 2.1. Radiotracer experiments

Twenty scallops (*C. radula* [Linnée, 1758]) were collected in the vicinity of Noumea City (i.e. Sainte Marie Bay) by SCUBA diving in October 2004. The organisms (76 ± 3 mm length) were experimentally exposed to four radio-tracers ( $^{109}$ Cd,  $^{57}$ Co,  $^{54}$ Mn, and  $^{65}$ Zn) via seawater. Seawater was spiked with low nominal activities of each selected radiotracer: 1.76 kBq l<sup>-1</sup>  $^{109}$ Cd, 0.91 kBq l<sup>-1</sup>  $^{57}$ Co, 0.65 kBq l<sup>-1</sup>  $^{54}$ Mn, 0.65 kBq l<sup>-1</sup>  $^{65}$ Zn. In terms of stable metal concentration, these additions corresponded to 9 × 10<sup>-3</sup> ng Cd l<sup>-1</sup>,  $6 \times 10^{-3}$  ng Co l<sup>-1</sup>,  $5 \times 10^{-2}$  ng Mn l<sup>-1</sup> and,  $4 \times 10^{-3}$  ng Zn l<sup>-1</sup>, which are one to five orders of magnitude lower than the background concentrations of these metals in open seas (Bruland, 1983).

The uptake of the radiotracers was followed during four days using gamma spectrometric techniques. Uptake kinetics were expressed as change in concentration factor (CF; ratio of activity of the radiotracer in the whole soft tissues or in a body compartment  $-Bq g^{-1}$  wet wt - to time-integrated activity of radiotracer in seawater  $-Bq g^{-1}$ ) according to time. Radiotracer uptake kinetics were described using a simple linear regression model

At various time intervals (12, 30, 48, 72, and 96 h), four individuals were randomly sampled and dissected in order to determine the tissue distribution of the elements. Shells, digestive gland, kidneys, gills, gonad, adductor muscle and the remaining soft tissues were separated and counted using a high-resolution  $\gamma$ -spectrometry system consisting of four coaxial Ge (N- or P-type) detectors (EGNC 33-195-R, Canberra<sup>®</sup> and Eurysis<sup>®</sup>) connected to a multi-channel analyzer and a computer equipped with a spectra analysis software (Interwinner<sup>®</sup> 6). The radioactivity of the samples was determined by comparison with standards of known activities and appropriate geometries and was corrected for background and physical decay of the radiotracers. Counting times were adapted to obtain counting rates with relative propagated errors lower than 5%.

## 2.2. Trace element analyses

Ten scallops were collected by SCUBA diving in October 2004 in the southern New Caledonian lagoon in two bays showing different physicochemical specificities (Fig. 1): Maa Bay (n = 5;  $76 \pm 6$  mm length) and Sainte Marie Bay (n = 5;  $77 \pm 2$  mm length). This last site, located to the East of Noumea, is characterised by elevated inputs of sewage sludge (Jaquet, 2005). Moreover terrigenous inputs due the open casting mining are released by the Coulée River and deposited along the shoreline, reaching areas as distant as Sainte Marie Bay (Fernandez et al., 2006). Conversely, Maa Bay is located 20 km North-West of Noumea and is considered to be exempt from direct anthropogenic inputs of metals and the area is sparsely inhabited. Moreover, Maa Bay is not surrounded by lateritic soils (viz. a typical source of metal contamination by natural erosion) and is therefore considered as a non-contaminated "reference" site in this study.

In order to verify the contamination level of the two locations, sediments were collected by SCUBA diving in parallel to organisms. Sediments were stored in acid-washed and hermetically sealed plastic bags until return to the laboratory.

Upon arrival to the laboratory, organisms were depurated for 24 h in clean seawater aquaria to remove pseudo-faecal and faecal material from the digestive tract. Each individual was measured and weighed prior to dissection: the digestive gland, kidneys, gills, gonads and adductor muscle were carefully removed from each individual. The remaining tissues were also taken into account in order to calculate the whole metal content of the organisms.

All samples (scallop tissues and sediments) were dried for several days at 60 °C until they reached a constant weight. Then, sediments were sieved on a 1 mm meshed sieve in order to eliminate heterogeneous materials (e.g. stones, fragments of corals) prior to elemental analysis. Aliquots of the samples ranging from 50 to 300 mg were digested using a 3:1 (v:v) nitric—hydrochloric acid mixture with 65% HNO<sub>3</sub> (Merck, suprapur quality) and 70% HCl (Merck, suprapur quality). Acidic digestion was performed overnight under ambient temperature and then the samples were heated in a microwave during 30 min with increasing temperature until 105 °C, and 15 min at 105 °C (1200 W). After the mineralization process, each sample was diluted to 30 or 50 ml with milli-Q quality water, according to the volume of acid added to the mineralization (3 or 4.5 ml).

Ag, As, Cd, Co, Cr, Cu, Fe, Mn, Ni and Zn were analysed either by ICP-OES (Varian<sup>®</sup> Vista-Pro) or ICP-MS (Varian<sup>®</sup> Ultra Mass 700). Reference tissues (dogfish liver DOLT-3, NRCC, and lobster hepatopancreas TORT-2, NRCC) were treated and analysed in the same way as the samples. Results were in good agreement with the certified values, and the standard deviations were low, proving good repeatability of the method. The results for standard reference materials displayed recoveries of the elements ranging from 79% to 122% (n = 10). For each set of analyses, blanks were included in each analytical batch. The detection limits ( $\mu g g^{-1}$  dry wt) for ICP-OES were 10.1 (As), 0.8 (Cr), 0.5 (Cu), 0.3 (Fe), 0.04 (Mn), 1.1 (Ni) and 0.7 (Zn) and for ICP-MS, they were 0.1 (Ag), 0.3 (Cd), 0.03 (Co). All metal concentrations are given on a dry weight basis ( $\mu g g^{-1}$  dry wt).

#### 2.3. Preliminary risk assessment for consumers

A maximum recommended consumption of scallops was evaluated on the basis of the Provisional Maximum Tolerable Daily Intake (PMTDI) or

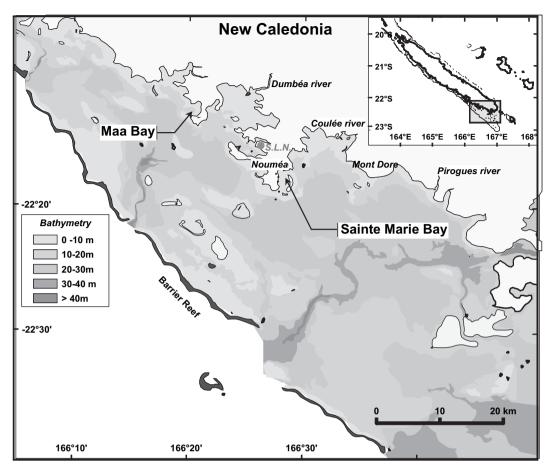


Fig. 1. Sampling locations along the SW coast of New Caledonia.

Provisional Tolerable Weekly Intake (PTWI) given by the Joint Expert Committee on Food Additives (http://www.inchem.org/pages/jecfa.html).

In this calculation, metal sources supplied by other meals or by drinking water on the same day were not taken into account, i.e. only metal intake coming from the scallop meal has been considered. The PMTDIs for Cu, Fe, and Zn are respectively 500, 800 and 1000  $\mu$ g kg<sup>-1</sup> d<sup>-1</sup> and PTWI for As and Cd are 15 and 7  $\mu$ g kg<sup>-1</sup> wk<sup>-1</sup> (JECFA, 2006; WHO, 1989). The calculation was done for As, Cd, Cu, Zn and Fe by dividing the load ( $\mu$ g) of each element measured in whole soft parts or in the adductor muscle (for both sites) by the respective PMTDI or PTWI and by a consumer average body weight (viz. 50 and 80 kg body weight for female and male humans, respectively). In this manner, results indicated the maximum number of consumed *C. radula* not to be exceeded. Although *C. radula* is generally eaten whole, both whole soft parts and the adductor muscle alone were considered in the risk calculations for comparison purpose (viz. the adductor muscle is the only tissue eaten in the most usually marketed scallop species).

## 2.4. Data treatment

Statistical analyses of the data were performed by one- or two-way AN-OVA, followed by the multiple comparison test of Tukey. The variability explained by each factor was derived from the sum of squares (Warnau et al., 1998). After verification of the variance homogeneity, two-way analysis of variance (ANOVA) was used with sampling location and body compartment as fixed factors. When the homogeneity hypothesis was rejected by ANOVA, a multiple comparison test of Tukey was carried out to assess the most significant impact levels in the examined stations, and then the most significant metal concentrations in the body compartments. Comparison of data expressed as percentages (body distribution) was carried out after arcsine transformation of the data with respect to basic normality requirements of parametric tests (Zar, 1996).

Concerning radiotracer data analyses, kinetics of metal uptake in scallops were fitted using linear regression routines (Statistica<sup>®</sup> 6). The best fit of the different regressions was selected by examination of residuals and  $R^2$ . Linearity of the kinetics was tested by the linearity test for regression with replication (Zar, 1996).

The level of significance for statistical analyses was always set at  $\alpha = 0.05$ .

# 3. Results

## 3.1. Laboratory experiments: radiotracer uptake study

The bioaccumulation kinetics of the four radiotracers in the whole soft parts of *C. radula* were best fitted by a linear model (Fig. 2). The uptake rate constants ( $k_u$ ) found for the studied radiotracers rank as follows: Mn = Co < Cd < Zn. After four days of exposure, the soft parts reached concentration factors (CFs) of  $62 \pm 14$  for Cd,  $15 \pm 8$  for Co,  $15 \pm 4$  for Mn, and  $149 \pm 34$  for Zn (Fig. 2). In addition, CFs were calculated for each organ (Table 1). The latter ones clearly show the contrasting efficiencies of metal accumulation by scallop tissues. Indeed, the highest CF was found in the kidneys for <sup>109</sup>Cd ( $391 \pm 69$ ) and <sup>57</sup>Co ( $603 \pm 298$ ) and in the digestive gland for <sup>65</sup>Zn ( $549 \pm 92$ ). In the case of <sup>54</sup>Mn, the CFs of gills, kidneys and digestive gland approximately reached the same value ( $\sim 30$ ) after four days of exposure.

At the end of the exposure period, the distribution of the radiotracers was compared among the scallop body

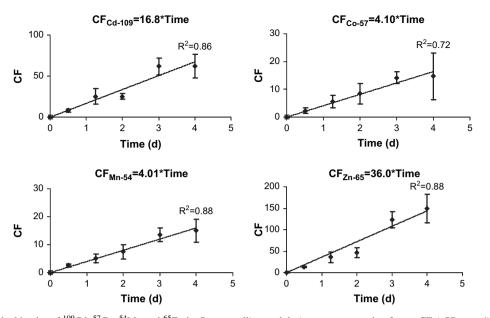


Fig. 2. Whole-body uptake kinetics of <sup>109</sup>Cd, <sup>57</sup>Co, <sup>54</sup>Mn and <sup>65</sup>Zn in *Comptopallium radula* (mean concentration factor,  $CF \pm SD$ ; n = 4) and the corresponding linear model (p < 0.0001).

compartments following two steps (Table 2): 1) between the shell and the soft parts and 2) among the different tissues and organs constituting the soft parts. Considering the whole scallops, <sup>57</sup>Co and <sup>54</sup>Mn were mainly (i.e. 92% and 96%, respectively) contained in the shell. Among the soft tissues, <sup>54</sup>Mn was mainly associated with the digestive gland (33%) and the gills (33%), whereas kidneys contained most of the <sup>57</sup>Co (65%). The two other radioisotopes were mainly present in the soft parts (69% for <sup>109</sup>Cd and 58% for <sup>65</sup>Zn), with 65% of the soft part <sup>65</sup>Zn activity being in the digestive gland and <sup>109</sup>Cd being mainly present in the gills (38%) and the digestive gland (25%).

# 3.2. In situ analyses

Arsenic and metal concentrations in the sediment and in the tissues and organs of *C. radula* collected from the two sampling sites are given in Tables 3 and 4. Except for Ag which concentrations were below the detection limit, sediments from Maa Bay displayed significantly higher elemental concentrations compared to the supposedly more contaminated site of Sainte Marie Bay (Table 3). In contrast, scallops from

Table 1

Concentration factor (mean  $\pm$  SD; n = 4) of the radiotracers in the different body compartments of *Comptopallium radula* after four days of seawater exposure

Compartment	<sup>109</sup> Cd	<sup>57</sup> Co	<sup>54</sup> Mn	<sup>65</sup> Zn
Digestive gland	$107\pm29$	$9.7\pm4.1$	$29\pm11$	$549\pm92$
Kidneys	$391\pm69$	$603\pm298$	$33\pm12$	$184\pm52$
Gonad	$76\pm75$	$3.1\pm1.8$	$14\pm 6$	$55\pm14$
Muscle	$7.0 \pm 1.9$	$0.6\pm0.3$	$2.0\pm0.4$	$18\pm5.3$
Gills	$142\pm29$	$3.2\pm0.5$	$24\pm5.3$	$96\pm25$
Remaining tissues	$3.2\pm0.6$	$0.2\pm0.1$	$0.6\pm0.2$	$4.2\pm0.9$

Maa Bay showed higher concentrations of As, Cd, and Fe in their tissues whereas Ag was far more concentrated in scallops from Sainte Marie Bay. Independently of the sampling site, the digestive gland and kidneys of *C. radula* generally showed the highest element concentrations (Table 4). The distribution patterns of elemental concentrations were similar between scallops from both sampling sites except for Ag which specifically accumulated in the digestive gland in Maa Bay and in the kidneys in Sainte Marie Bay.

The body distributions of the different elements among the tissues and organs of the scallops from both sites are compared in Fig. 3. The digestive gland clearly exhibited the highest proportion of Ag, As, Cr, Fe and Zn in both areas. In the case of Cd and Ni, the digestive gland and kidneys contained most of the metal burden. Regarding Co and Cu, the major part of the metals was contained in kidneys in Sainte Marie Bay whereas in Maa Bay, the digestive gland also stored a large fraction of Co and contained the major part of Cu. In the case of Mn, distribution among tissues and organs is rather homogeneous in scallops from the two bays.

Table 2

Metal distribution (mean  $\% \pm$  SD, wet wt basis; n = 4) between shell and soft parts; and within the soft parts of *Comptopallium radula* after four days of seawater exposure

water exposure					
Compartment	<sup>109</sup> Cd	<sup>57</sup> Co	<sup>54</sup> Mn	<sup>65</sup> Zn	
Shell	$31\pm 6$	$92\pm4$	$96\pm1$	$42\pm9$	
Soft parts	$69\pm 6$	$8\pm4$	$4\pm1$	$58\pm9$	
Digestive gland	$25\pm7$	$9.2\pm2.4$	$33\pm7$	$65\pm3$	
Kidneys	$13\pm1$	$81\pm4.9$	$5.6\pm1.7$	$3.2\pm0.6$	
Gonad	$14 \pm 11$	$2.5\pm0.6$	$14\pm4$	$5.8\pm1.2$	
Muscle	$8.3\pm1.0$	$2.8\pm1.2$	$12 \pm 1$	$11\pm1.9$	
Gills	$38\pm7$	$3.9\pm1.7$	$33\pm7$	$13\pm1.8$	
Remaining tissues	$2.2\pm0.3$	$0.8\pm0.5$	$2.2\pm0.4$	$1.5\pm0.3$	

Table 3 Element concentrations (mean  $\pm$  SD;  $\mu$ g g<sup>-1</sup> dry wt; n = 5) in sediment from the two sampling locations

	Maa Bay	Sainte Marie Bay		
Ag	$< dl^a$	$< dl^a$		
As	$6.4 \pm 0.3$	$5.8 \pm 0.2$		
Cd	$1.0 \pm 0.2$	<d.1. <sup>a</sup>		
Со	$4.6 \pm 2.3$	$1.9 \pm 0.1$		
Cr	$44\pm 8$	$15 \pm 1$		
Cu	$11 \pm 3$	$0.8 \pm 0.1$		
Mn	$132\pm8$	$33 \pm 3$		
Ni	$64 \pm 13$	$12 \pm 1$		
Zn	$15\pm3$	$3.9\pm0.1$		

<sup>a</sup> dl: Detection limit.

## 3.3. Risk assessment

The element content in the whole soft parts of the scallops allowed the computation of the number of bivalves to be eaten by male and female humans to reach the PMTDI or, in the case of As and Cd, the PTWI (Table 5). Overall, As appears to be the main element of concern regarding the consumption of *C. radula* from both locations since the consumption of four (for women) or eight (for men) scallops would result in exceeding the PTWI threshold. For the other elements, the consumption of more than 40 scallops is necessary to reach the recommended PMTDI or PTWI.

When only the adductor muscle was considered, the associated risk was lower as expected (element concentrations in the muscle were always lower than in whole soft tissues). If any, As was the only element of concern; however, 27 adductor muscles for women and 44 for men should be eaten before reaching the PTWI threshold (data not shown). For the other elements, the consumption of more than 374 scallops is necessary to reach the recommended PMTDI or PTWI.

## 4. Discussion

Several studies have pointed out the ability of various scallop species to accumulate high levels of trace elements in their tissues (e.g. Bryan, 1973; Bustamante and Miramand, 2005; Uthe and Chou, 1987). Such a bioaccumulation ability does not appear to be linked to specific anthropogenic contamination of the environment since scallops from remote areas such as the North Atlantic or Antarctica were shown to contain elevated Cd concentrations compared to related or even identical species from temperate regions (Berkman and Nigro, 1992; Bustamante and Miramand, 2004; Mauri et al., 1990; Viarengo et al., 1993). Conversely, very few studies have focused on tropical scallops (Francesconi et al., 1993) and the current literature globally lacks related baseline information.

In this work, the bioaccumulation efficiency of trace elements of the tropical scallop *C. radula* has been evaluated by comparing individuals collected from two contrasting coastal areas. To assess the difference in the contamination status of both stations, trace elements were analysed in sediments (Table 3). Surprisingly, element concentrations in sediment were high in the reference site (Maa Bay) compared to those from the site close to Noumea City and subject to higher terrigenous inputs (Sainte Marie Bay). The only exception to this was Ag which was below the detection limit of the method whereas Co, Cr, Cu, Mn, Ni and Zn concentrations were at least two times higher in the sediment from Maa Bay compared to Sainte Marie Bay. Even if these results are somewhat surprising, they demonstrate that the two sampling sites are actually contrasting from the point of view of the relative bioavailability of the former elements.

Regarding the levels in *C. radula*, scallop tissues highly concentrated the analysed elements irrespective of the environment, which is consistent with reported data for other pectinid species (e.g. Bryan, 1973; Bustamante and Miramand, 2004; Mauri et al., 1990; Uthe and Chou, 1987). Interestingly, the bioconcentration ability of *C. radula* allowed measuring relatively high levels of elements, even when sediment analysis showed concentrations below detection limit, as it was the case for Ag (see Tables 3 and 4). Out of the 10 analysed elements, Ag levels were significantly higher (p < 0.0001) in *C. radula* collected in Sainte Marie Bay whereas As, Cd and Fe concentrations were higher in scallops from Maa Bay; the other elements did not show significant differences (p > 0.05) between the two bays.

Silver is a proxy for anthropogenic input in coastal waters due to its high enrichment in sewage sludge from coastal cities (Andren and Bober, 2002; Luoma et al., 1995). Hence, higher levels of Ag in scallop tissues from Sainte Marie Bay indicated that this area is subject to urban pollution which supports the results of the modelling approach recently published by Fernandez et al. (2006). Specifically, exposure to high Ag levels in Sainte Marie Bay lead to the accumulation of the metal in the kidneys of C. radula rather than in the digestive gland as observed in Maa Bay. In other Pectinidae species, the digestive gland generally exhibits the highest Ag concentrations among soft tissues (Bryan, 1973; Bustamante and Miramand, 2004, 2005; Segar et al., 1971). To the best of our knowledge, these very different Ag storage strategies of C. radula between two sites have never been reported for the Pectinidae family and deserve further investigation.

In contrast to Ag, As has received little attention in pectinids. The levels of this element in the soft parts of C. radula were quite high in both bays (i.e.  $45 \pm 6$  and  $87 \pm 12 \ \mu g \ g^{-1}$ dry wt in Sainte Marie Bay and Maa Bay, respectively) which rather contrasts with published data in Pectinidae from temperate zones which display generally much lower concentrations (Bustamante and Miramand, 2005; Lai et al., 1999). For example, the variegated scallop Chlamys varia from the Bay of Biscay (France) reached maximum As concentrations of  $25 \pm 5 \ \mu g \ g^{-1}$  dry wt in the whole soft parts (Bustamante and Miramand, 2005). Bivalves generally display a limited capacity in accumulating As from seawater (e.g. Hédouin, 2006; Unlü and Fowler, 1979) and available literature indicates that the high As levels measured in tropical pectinids probably originate from their diet (see e.g. Neff, 1997). Hence, higher As levels in C. radula soft tissues could indicate a high dietary source and/or an elevated bioavailability of As in their

Table 4 Element concentrations (mean  $\pm$  SD and min-max values,  $\mu g g^{-1}$  dry wt; n = 5) in tissues and organs of the scallop *Comptopallium radula* from two stations in the South-West lagoon of New Caledonia

	Ag	As	Cd	Co	Cr	Cu	Fe	Mn	Ni	Zn
Maa Bay										
Digestive gland	$0.14\pm0.12$	$340\pm58.4$	$7.11 \pm 3.93$	$3.23\pm0.62$	$12.3\pm3.37$	$12.6\pm3.63$	$1327\pm79$	$8.22 \pm 1.62$	$45.8 \pm 13.6$	$787\pm398$
	<dl<sup>a-0.32</dl<sup>	293-420	4.50-13.0	2.31-3.61	7.95-15.8	8.77-17.3	1219-1409	7.11-10.6	27.7 - 60.8	412-1341
Kidneys	$< dl^a$	$297 \pm 109$	$80.1\pm51.7$	$84.5\pm36.4$	$9.47\pm0.48$	$7.90 \pm 1.05$	$147\pm73$	$18.9\pm4.34$	$343\pm138$	$344\pm175$
·		181-424	36.7-149	46.3-117	8.88-10.0	6.43-8.72	95-255	12.9-22.3	216-477	142-508
Gills	$< dl^a$	$75.1\pm23.5$	$2.78\pm2.13$	$1.51\pm0.23$	$1.75\pm0.34$	$5.10\pm0.81$	$239 \pm 109$	$5.82\pm0.36$	$7.48 \pm 2.24$	$84 \pm 15$
		49.3-99.2	1.54 - 5.95	1.27-1.79	1.29-2.04	4.30-5.89	156-386	5.42-6.29	5.58-10.6	71-105
Gonad	$< dl^a$	$25.2\pm13.9$	$2.02\pm1.32$	$0.24\pm0.18$	$1.96 \pm 1.12$	$4.86\pm0.62$	$92\pm60$	$14.1\pm9.31$	$1.74 \pm 1.67$	$38\pm20$
		11.2-44.4	1.13-3.99	0.10-0.47	0.73-3.45	4.28-5.73	50-181	1.30-23.6	<dl<sup>a-4.11</dl<sup>	13-61
Muscle	$< dl^a$	$30.5\pm5.08$	$1.32\pm1.69$	$0.11\pm0.03$	$< dl^a$	$0.60\pm0.16$	$11 \pm 2.2$	$0.58\pm0.12$	$< dl^a$	$70 \pm 22$
		25.0-36.2	0.39-3.86	0.08 - 0.15		$< dl^{a} - 0.81$	8.7-67	0.44-0.73		56-102
Remaining tissues	$< dl^a$	$43.6 \pm 14.7$	$1.67 \pm 1.64$	$0.62\pm0.12$	$2.17\pm0.58$	$1.63\pm0.21$	$155\pm55$	$4.90 \pm 1.35$	$3.08\pm0.82$	$37\pm8$
-		27.4-57.7	0.76-4.13	0.56 - 0.80	1.41-2.64	1.47-1.93	116-236	3.59-6.29	1.86-3.65	29-47
Whole soft parts	$< dl^a$	$86.9 \pm 11.6$	$3.57 \pm 2.69$	$1.98\pm0.54$	$3.17 \pm 1.25$	$3.91\pm0.38$	$289\pm80$	$5.96 \pm 2.44$	$13.8\pm4.39$	$176\pm62$
•		72.4-100	1.92-7.56	1.35-2.66	1.78-4.38	3.60-4.45	221-375	3.60-9.30	10.2-19.8	90-235
Sainte Marie Bay										
Digestive gland	$16.9\pm4.3$	$74.1 \pm 16.0$	$0.75\pm0.20$	$0.58\pm0.39$	$9.24 \pm 4.62$	$9.82 \pm 3.51$	$650\pm344$	$5.87 \pm 3.17$	$15.9\pm7.85$	$580\pm356$
	11.3-22.2	56.2-94.3	0.56-1.09	0.29-1.20	4.72-16.8	6.17-15.0	367-1195	3.03-11.3	7.85-28.7	243-1115
Kidneys	$104\pm35$	$103 \pm 18$	$31.8\pm21.2$	$104\pm49.7$	$11.0\pm1.78$	$11.0\pm3.00$	$122 \pm 21$	$99.5\pm81.7$	$555\pm224$	$687\pm383$
•	72.3-160	72.6-116	4.92-57.8	58.6-179	9.05-13.2	7.50-15.2	108-158	13.5-187	331-876	138-1094
Gills	$0.78\pm0.37$	$59.9 \pm 7.37$	$1.72\pm0.66$	$1.37\pm0.21$	$2.51\pm0.33$	$4.35\pm0.27$	$325\pm62$	$7.14\pm0.48$	$6.54 \pm 1.56$	$99 \pm 14$
	0.55-1.43	54.1-72.7	1.14 - 2.70	1.22-1.73	2.30-3.09	4.12-4.80	264-405	6.40-7.72	4.57-8.15	81-120
Gonad	$0.67\pm0.28$	$21.2\pm3.39$	$0.49\pm0.20$	$0.24\pm0.15$	$2.05\pm1.08$	$3.80 \pm 1.01$	$118\pm65$	$12.0\pm12.9$	$2.90\pm2.30$	$33\pm16$
	0.28 - 0.94	18.0-25.9	0.28-0.81	0.08 - 0.45	<dla-3.37< td=""><td>2.95-5.01</td><td>56-217</td><td>2.06-29.0</td><td><dl<sup>a-6.49</dl<sup></td><td>20-51</td></dla-3.37<>	2.95-5.01	56-217	2.06-29.0	<dl<sup>a-6.49</dl<sup>	20-51
Muscle	$< dl^a$	$26.6\pm6.98$	$0.45\pm0.29$	$0.06\pm0.02$	$< dl^a$	$< dl^a$	$13 \pm 10$	$0.62\pm0.14$	$< dl^a$	$57\pm8$
		15.7-32.9	0.26-0.96	<dl<sup>a-0.09</dl<sup>			5.1-26	0.45 - 0.78		49-67
Remaining tissues	$0.43\pm0.22$	$47.3 \pm 19.6$	$0.89\pm0.63$	$0.38\pm0.12$	$2.80\pm0.55$	$1.54\pm0.08$	$178\pm63$	$4.07\pm0.95$	$3.79 \pm 1.58$	$47\pm7$
e	0.16-0.74	21.2-67.1	0.40 - 1.87	0.25-0.58	2.02 - 3.28	1.43-1.63	118-283	2.56 - 5.00	1.66-5.29	36-56
Whole soft parts	$5.24 \pm 1.22$	$44.7\pm5.8$	$1.12\pm0.53$	$1.79\pm0.7$	$3.26 \pm 1.15$	$3.70\pm0.97$	$221\pm88$	$6.06\pm2.46$	$12.8\pm3.45$	$183\pm90$
*	3.92-6.48	39.3-53.6	0.54 - 1.92	0.99 - 2.79	2.12-4.97	2.88-5.14	162-375	2.81-8.70	8.65-17.2	109-308

<sup>a</sup> dl: Detection limit.

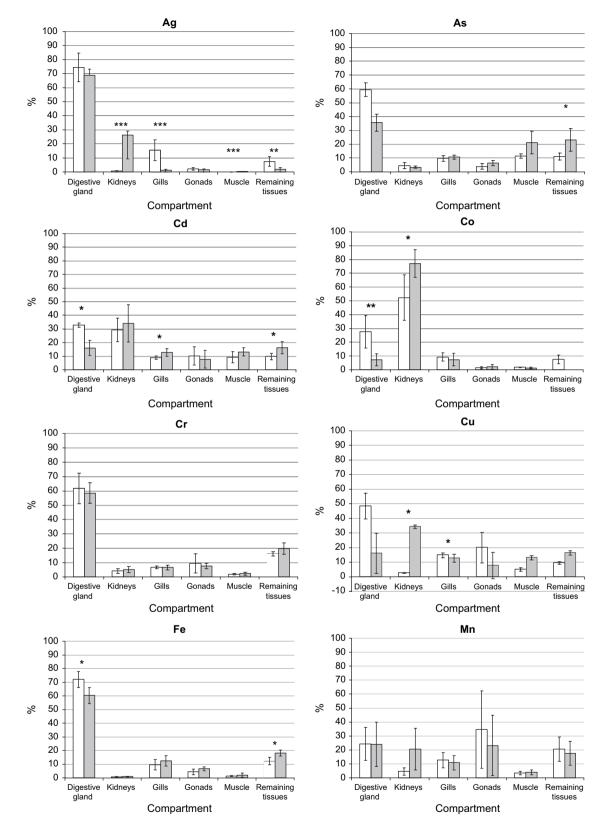


Fig. 3. Comparison of trace element distribution (mean  $\% \pm$  SD; dry wt basis; n = 5) in tissues and organs of *Comptopallium radula* from Maa Bay (white bars) and Sainte Marie Bay (dark grey bars). Significant differences between stations are indicated by \* (p < 0.05), \*\* (p < 0.01) or \*\*\* (p < 0.005).

environment. Such an hypothesis would be consistent with coastal organic contamination due to agricultural activities and waste discharges from shrimp aquaculture that has been reported for the two bays where *C. radula* was collected

(Labrosse et al., 2000; Lemonnier and Faninoz, 2006). Indeed, important discharges of N-enriched compounds could locally modify the N:P ratio and as phytoplankton metabolises As more easily in a phosphate-depleted environment Table 5

Element content in C. radula (µg)		PMTDI or PTWI* $(\mu g k g^{-1})$	Number of scallops to reach PMTDI or PTWI				
Element Maa Bay	Maa Bay	Sainte Marie Bay		Maa Bay		Sainte Marie Bay	
				50 kg	80 kg	50 kg	80 kg
As	214	144	15*	4	6	5	8
Cd	8.5	3.6	7*	41	66	97	155
Cu	9.4	12.0	500	2653	4244	2080	3328
Fe	720	688	800	56	89	58	93
Zn	450	602	1000	111	178	83	133

Mean element content ( $\mu$ g) in scallops from the two sampling stations and number of individuals to be eaten to reach the PMTDI and the PTWI for women (50 kg) and men (80 kg)

\* Refers to PTWI values (the other ones in the column being PMTDI).

(e.g. Smedley and Kinniburgh, 2002), this might result in As transfer to filter-feeders and enhanced As bioaccumulation in their tissues (Benson and Summons, 1981). Similarly, the higher As concentrations observed in scallops from Maa Bay than in Sainte Marie Bay would likely to be due to difference in dietary bioavailability of As between the two areas.

Higher Cd and Fe concentrations in C. radula from Maa Bay also support this hypothesis. Indeed, authors generally agree that high Cd concentrations in scallop tissues are related to their intense feeding rates (e.g. Berkman and Nigro, 1992; Palmer and Rand, 1977) and Fe tends to complex with organic matter or colloids, leading to a high bioavailability to phytoplankton (Chen et al., 2003). Other studies also reported similar results with higher As and/or Cd concentrations in scallop from clean sites compared to contaminated ones (viz. C. varia from France, Bustamante and Miramand, 2005, and Placopecten magellanicus from Canada, Uthe and Chou, 1987). In both cases, these atypical levels were attributed to feeding and nutritional inadequacy of the diet. A further characterization of food composition (stomach content analyses) and/or metal concentrations in the food of the scallop in the sampling areas would be the next step to validate such an assumption. Even if the food seems to be the major accumulation pathway, the contribution of dissolved compartment cannot be neglected. Indeed, radiotracer experiments revealed the relatively high uptake of waterborne <sup>109</sup>Cd in scallops, clearly showing that the high accumulation of Cd would not be only restricted to dietary inputs.

Although not different between the two studied stations, Co, Cr and Ni concentrations measured in the tissues of C. radula appear relatively elevated when compared with data from the literature (Bryan, 1973; Bustamante and Miramand, 2005; Pesch et al., 1977). Some of the variations could account for the specific characteristics of metal accumulation, but high Co. Cr and Ni concentrations might rather be due to natural and anthropogenic inputs of these metals in the context of the local geology of New Caledonia since lateritic soils are particularly rich in these three very elements (Labrosse et al., 2000). However, the lack of differences in tissues between the two sampling sites was unexpected in regard to sediment results and suggests that the scallops readily accumulate these three metals until a threshold level above which a higher contamination does not affect any longer the levels within C. radula tissues. Field experiments such as transplantations from relatively clean to strongly contaminated sites would be interesting to validate this hypothesis.

Studies on metal concentrations in the tissues of scallops from various locations around the world have highlighted their common ability to concentrate Mn and Zn in the kidneys up to very high levels, typically exceeding  $1000 \ \mu g \ g^{-1}$  dry wt

Table 6

Co, Cu, Mn, and Zn concentrations (mean $\pm$ SD; $\mu$ g g <sup>-</sup>	<sup>1</sup> dry wt) in kidneys from different scallop species
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Species	Co	Cu	Mn	Zn	Sampling area	References
Aequipecten opercularis	_	$507 \pm 144$	_	$31053\pm8147$	Bay of Biscay	Bustamante and Miramand (2004)
A. opercularis	_	$106\pm49$	_	$6632\pm3664$	Faroe Island	Bustamante and Miramand (2004)
A. opercularis	$15.1\pm6.1$	$1285\pm591$	$17300\pm4570$	$40800\pm7290$	English channel	Bryan (1973)
Chlamys varia	_	$142\pm20$	_	$24107\pm3465$	Bay of Biscay	Bustamante and Miramand (2004)
Pecten jacobeus		$17.5\pm10.9$	$6390\pm3030$	$2790 \pm 1340$	Adriatic sea	Mauri et al. (1990)
Pecten novae-zelandiae	_	_	2660	2630	New Zealand	Brooks and Rumsby (1965)
Pecten maximus	_	$13.5\pm2.0$	_	$7278\pm3041$	Bay of Biscay	Bustamante and Miramand (2004)
P. maximus	$9.05\pm3.07$	$20.8\pm0.3$	$15300\pm4100$	$19300\pm4800$	English channel	Bryan (1973)
P. maximus	_	$117\pm100$	$17811 \pm 11978$	$32766\pm 11978$	Scotland	George et al. (1980)
Patinopecten yessoensis	$0.35\pm0.01$	$5.4 \pm 1.7$	$10\pm5$	$174 \pm 149$	Japan	Ishii et al. (1985)
Adamussium colbecki	_	$4.0\pm1.7$	$16.3\pm8.9$	$199\pm89$	Antarctic	Mauri et al. (1990)
Comptopallium radula	$84\pm36$	$7.9\pm1.0$	$18.9\pm4.3$	$344 \pm 175$	Maa Bay	Present study
C. radula	$104\pm50$	$11 \pm 3$	$100 \pm 82$	$687\pm383$	Sainte Marie Bay	Present study

(see Table 6). However, *C. radula* does not follow this trend (Table 4), and rather behaves as the southern and the northern scallops *Adamussium colbecki* and *Patinopecten yessoensis*, respectively (Ishii et al., 1985; Mauri et al., 1990). Although kidney cells of *A. colbecki* exhibit the same cytoarchitecture as the other Pectinidae, it has been shown that the low renal bioaccumulation capacity for Mn and Zn in this species was due to the low degree of mineralization of their nephrolithes, viz. the renal phosphate-rich concretions that are well known for their ability to trap and detoxify intracellular metals (George et al., 1980; Nigro et al., 1992). Hence, the atypical low levels of Mn and Zn in *C. radula* kidneys might be due to interspecific diversity in the amount and/or degree of mineralization of renal concretions.

The lower Zn accumulation capacity of the kidneys compared to the digestive gland was also observed in our laboratory experiment:  $^{65}$ Zn concentration factor was far higher in the digestive gland than in kidneys of *C. radula*. Interestingly, Bryan (1973) showed that the Queen scallop *Aequipecten opercularis* (a scallop presenting typical, highly mineralized renal concretions) concentrated  $^{65}$ Zn to a much higher extent in kidneys than in digestive gland. These observations would support the hypothesis that *C. radula* might have developed an alternative detoxification strategy for Zn in the digestive gland to compensate the supposedly low mineralization of its renal concretions.

In New Caledonia, C. radula is commonly fished and consumed whole. Therefore, we performed a preliminary risk assessment taking into account the trace element concentrations measured in scallops from the two selected bays. Accordingly, As appears to be the only limiting element for human consumption of these scallops. Indeed, consumption of the whole soft parts of four to eight C. radula during a single week would lead to exceeding the PTWI value recommended for As by WHO (viz.  $15 \,\mu\text{g}\,\text{kg}^{-1}\,\text{wk}^{-1}$ ; WHO, 1989). However, one has to keep in mind that As speciation has not been taken into account in the present computation. Now, it is well known that both bioavailability and toxicity of As largely depend on its chemical speciation (see e.g. Neff, 1997; Warnau et al., 2007). Inorganic As species (i.e. arsenite and arsenate), on which is based the PTWI, are the most toxic forms both in terms of acute toxicity and carcinogenicity whilst it is generally reported that marine organisms bioaccumulate the element mainly as organic (e.g. arsenobetaine and arsenosugars) compounds that are not or slightly toxic (Neff, 1997; Warnau et al., 2007). Since our preliminary computations were made using the concentrations of total As, it is obvious that the actual number of scallops needed to reach As PTWI will be higher than 4 (for women) to 8 (for men). However, it has been shown in several marine species that inorganic As can sometimes represents a high proportion of total As concentrations (see e.g. Fattorini and Regoli, 2004; Warnau et al., 2007). Hence, rather than speculating on these values, the present computations should be considered as a warning signal about the very possible threat due to eating large meals composed of these scallops. Although traditional habits are not easily changed, a solution could be to recommend the sole consumption of adductor muscles (as is the case for most other pectinid species) since results showed consuming muscle was a non-risk issue, even without taking into account As speciation, as a minimum of 27 muscles would be needed to reach the As PTWI.

# 5. Conclusions

This work has shown that C. radula could be used to biomonitor Ag contamination in the New Caledonian lagoon. For the other elements being accumulated differently between the two sampling sites (i.e. As, Cd, and Fe), the differences between sites seem to reflect the elemental variations in the scallop environment, supporting the use of C. radula as a valuable biomonitor species for As, Cd and Fe contamination. However, additional information has to be collected from the field to understand more precisely the reason for such a difference. Furthermore, the contribution of the different routes of intake (seawater vs food) needs to be characterised. In this respect, a set of laboratory experiments using radiotracer techniques has been planned to explore the "dietary contamination pathway" hypothesis. Our results also allowed speculating on the occurrence of different accumulation/detoxification strategies of various metals (i.e. Ag, Mn and Zn) within the Pectinidae family. Finally, a preliminary estimation of trace element exposure to humans through scallop consumption has warned consumers about the possible health risks linked to elevated As accumulation by C. radula.

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