Histopathological and biochemical evidence of hepatopancreatic toxicity caused by cadmium and zinc in the white shrimp, *Litopenaeus vannamei*

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**Abstract**

The white shrimp, *Litopenaeus vannamei*, a globally important cultured prawn species, is an ideal animal for studying the impairment caused by the effects of heavy metals that are often detected in coastal areas. In this study, *L. vannamei* was exposed to different concentrations of cadmium (Cd) and zinc (Zn) for up to 28 d. Histopathological alterations in the hepatopancreas were observed in *L. vannamei* after long-term exposure to Cd and Zn. Hepatopancreatic injury was further confirmed by the inductions of two biochemical markers, hemolymphatic glutamate-oxalacetate transaminase (GOT) and glutamate-pyruvate transaminase (GPT). It was notable that *L. vannamei* was able to repair its hepatopancreas from the damage caused by Zn, which was evidenced by the results of the histopathological observations, determinations of tissue metal concentrations, and examination of GOT and GPT levels.

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1. Introduction

The hepatopancreas, being analogous to the liver and combining many of the functions of the liver, pancreas, and intestine of vertebrates, plays important roles in several metabolic processes in crustaceans (Caceci et al., 1988; Bhavan and Geraldine, 2000). Previous studies on the hepatopancreas at different biological levels such as the structure, development, physiology, metabolism, and biochemistry concluded that this digestive organ possesses several functions, including absorption, digestion, storage, and secretion (Dall and Moriarty, 1983; Caceci et al., 1988). In addition, several specific enzyme systems responsible for biotransformation, e.g., the cytochrome p450 system and glutathione S-transferase, as well as proteins such as metallothioneins appear in the hepatopancreas, so that this digestive organ also possesses the ability to biotransform, sequester, and detoxify many kinds of xenobiotics; some biotransformations, however, may in fact actually increase the toxicity of certain xenobiotics (Martinez et al., 1993; Pederson et al., 1997; James and Boyle, 1998; Snyder, 2000; Ahearn et al., 2004). Therefore, the hepatopancreas, or so-called mid-gut gland, is a very important organ for crustaceans, and topics related to it have consistently attracted the attention of many researchers. Although the hepatopancreas is responsible for the major portion of detoxification activities in crustaceans when exposed to toxicants and pollutants, in aquatic crustaceans, its functions and structure are likely to be affected by certain xenobiotics, such as pesticides and aflatoxin (Lightner et al., 1982; Bautista et al., 1994; Bhavan and Geraldine, 2000). These materials have been reported and demonstrated to have hepatopancreatic toxicities, which resulted in histological alterations to the hepatopancreas of studied organisms. Surprisingly, there are relatively few related studies on heavy metals even though they are common pollutants. We previously reported that the heavy metals, cadmium (Cd) and zinc (Zn), caused alterations and the unavailability of biochemical and nutritional materials within the hepatopancreas of the white shrimp, *L. vannamei*; these changes might have been the cause of growth retardation we observed after exposure to these metals (Wu and Chen, 2005a). However, the effects of these metals on the histological structure of the hepatopancreas of this species have never been reported.

Therefore, the major objective of the present paper was to study histopathological alterations in the hepatopancreas of *L. vannamei*, a globally important cultured prawn species, after exposure to Cd and Zn. Furthermore, the activities of two enzymes, glutamate-oxalacetate transaminase (GOT) and glutamate-pyruvate transaminase (GPT), were also measured in this study to serve as biochemical markers of hepatopancreatic toxicities, which resulted in histological alterations to the hepatopancreas of studied organisms. Surprisingly, there are relatively few related studies on heavy metals even though they are common pollutants. We previously reported that the heavy metals, cadmium (Cd) and zinc (Zn), caused alterations and the unavailability of biochemical and nutritional materials within the hepatopancreas of the white shrimp, *L. vannamei*; these changes might have been the cause of growth retardation we observed after exposure to these metals (Wu and Chen, 2005a). However, the effects of these metals on the histological structure of the hepatopancreas of this species have never been reported.

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2. Materials and methods

2.1. Animal maintenance

Postlarval *L. vannamei* shrimp were obtained from a commercial shrimp hatchery in Pingtung, southern Taiwan and maintained in...
the laboratory for over 2 months until they reached the juvenile stage (2.15 ± 0.18 g; 9.83 ± 0.59 cm long). Water conditions during shrimp rearing and the experimental period were a temperature of 25 °C, a salinity of 15 p.s.u., dissolved oxygen (DO) of 5.8–6.5 mg L⁻¹, and a pH of 7.15–7.87, under a 12:12-h light–dark regime with continuous aeration and filtration.

2.2. Histopathological studies of the hepatopancreas and determination of metal concentrations

*Ln. vannamei* shrimp in the intermolt stage were exposed to Cd or Zn and then sacrificed for the histological study of the hepatopancreas. Animals were divided into seven groups, each of which contained 10 shrimp that were exposed to concentrations of either 0.1, 0.2, or 0.4 mg Cd L⁻¹ as CdSO₄, or 0.05, 0.2, or 0.6 mg Zn L⁻¹ as ZnSO₄, and one control group (with no exposure to either metal).

Samples were taken after 7, 14, 21, and 28 d of exposure. The hepatopancreas samples were very carefully dissected out for both histopathological studies and determination of metal concentrations. For the histopathological studies, samples were fixed in 4% buffered formalin, embedded in paraffin, sectioned on an 8-μm thickness microtome on a microton (Microm, HM330, Heidelberg, Germany), stained with hematoxylin and eosin (H&E), and examined with an Olympus microscope (Tokyo, Japan). Metal concentrations in the hepatopancreas were determined by atomic absorption spectrophotometry as previously described (Wu and Chen, 2005b).

2.3. Determination of GOT and GPT activities

To examine the enzymatic activities of GOT and GPT, *Ln. vannamei* shrimp were exposed to concentrations of either 0.2 mg Cd L⁻¹ or 0.2 mg Zn L⁻¹, in addition to the control set, with each treatment containing at least 15 shrimp in a 20-L tank. Each treatment was repeated three times. Four to six shrimp in each treatment were randomly taken on days 7, 14, and 28. In all cases, hemolymph was extracted by heart puncture using a 0.5-mL syringe pre-rinsed with 9 g L⁻¹ NaCl before detection as suggested in the kit instructions.

2.4. Statistical analyses

Statistical analysis was performed with t-test to determine the difference between results of treated and control animals, and a p < 0.05 level was considered statistically significant.

3. Results

3.1. Histopathological studies of the hepatopancreas of *L. vannamei*

The hepatopancreas of control shrimp exhibited the well-organized glandular tubular structure normally seen in prawn species (Bell and Lightner, 1988; Caceci et al., 1988; Lightner et al., 1996; Bhavan and Geraldine, 2000). A longitudinal section of the apical region of a hepatopancreatic tubule showed that the cell surface facing the lumen was covered with a microvillous brush border, and the tubule apex contained undifferentiated embryonic (embryonalzellen) cells (E-cells; Fig. 1A). Moving away from the apex, the cells began to differentiate into developing absorptive, storage (restzellen) cells (R-cells). A transverse section of the middle-proximal region of the tubules showed that the tubules were empty of food material and appeared in a hexagonal arrangement or “star shape” in the lumen, and the basal lamina outlined each tubule (Fig. 1B). Also, in this section, different cell types could clearly be observed under a higher magnification (Fig. 1C). Developing R-cells are those in which the cytoplasm characteristically contains numerous vacuoles and lipid droplets, while fibrous (fibrillenzellen) cells (F-cells) are more basophilic, have larger nuclei than R-cells, and typically contain one prominent nucleoli. This region also contained the large distinctive secretory (blasenzellen) cells (B-cells), each of which contained one large apical secretory vacuole. Immediately surrounding each tubule was a network of myoepithelial cells with prominent nuclei and associated contractile fibers.

After exposure to Cd, alterations to the hepatopancreatic tissue of treated *Ln. vannamei* were observed. In shrimp exposed to lower concentrations of 0.1 and 0.2 mg Cd L⁻¹, large numbers of vacuoles appeared in the tubular epithelial cells of the hepatopancreas of animals exposed to 0.2 mg Cd L⁻¹ for 28 d, as well as one deeply affected individual exposed to 0.1 mg Cd L⁻¹ for 21 d (Fig. 2A and B). In addition, the star shape of the lumen was partially lost due to morphological changes of the tubular epithelial cells, because some cells decreased in height from a normal columnar height to a low cuboidal form. Also, a slight thickening of the basolateral lamina increased the distance between adjacent tubules in shrimp exposed to lower concentrations of Cd. When *Ln. vannamei* was exposed to a higher concentration of Cd (0.4 mg L⁻¹), severe lesions in the hepatopancreas were very obvious. In shrimp exposed to 0.4 mg Cd L⁻¹ for 14 d, tubular epithelial cells were heavily vacuolated and even ruptured (Figs. 2C and D). Thickening of the basal lamina and a decrease in cell height were also conspicuous with this treatment. Cellular hypertrophy and vacuolization resulting in a scrabby lumen surface were obvious, when the section was examined longitudinally in comparison to the hepatopancreas of normal animals (Fig. 2E). Furthermore, after exposure for 21 d, the arrangement of the tubules and shape of the lumens had substantially been altered (Fig. 2F). Infiltration of hemocytes began in the intertubular hemocoel between the middle-proximal portions of the hepatopancreas tubules. Generally, the distance between adjacent tubules was longer than that of normal shrimp. Atrophy and necrosis of tubules were conspicuous. Necrotic tubules of the hepatopancreas were rounded off and began to slough off their basement membrane (Fig. 2G). In this study, no shrimp survived after exposure to 0.4 mg Cd L⁻¹ for 28 d, and thus no histological results from this treatment are available.

Similarly, exposure to Zn caused some histological effects on the hepatopancreas of *Ln. vannamei*, especially those exposed to 0.2 and 0.6 mg Zn L⁻¹. Histological differences between the hepatopancreas of 0.2 mg Zn L⁻¹–treated shrimp and normal ones included vacuolization resulting in hypertrophy in a large number of tubular epithelial cells and an increase in the distance between adjacent tubules, probably due to detachment of tubules from their basement membrane (Fig. 3A and B). It is very interesting that symptoms of treated shrimps exposed to 0.2 mg Zn L⁻¹ for both 14 and 28 d were almost the same, and showed no time-dependence with these two treatments. In other words, the hepatopancreas of shrimp exposed to 0.2 mg Zn L⁻¹ for 28 d showed no more-serious or -substantial alterations than those individuals exposed to the same concentration of Zn for 14 d. However, when shrimp were exposed to a higher concentration of Zn (0.6 mg Zn L⁻¹), different alterations were apparent.
in shrimp at different exposure times. In the hepatopancreatic tubular systems of shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 14 and 21 d, slight to moderate vacuolization was apparent in tubular epithelial cells (Fig. 3C and D). In some regions of the hepatopancreatic tubules from shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 21 d, a decrease in the epithelial cell height, cell rupture, and formation of an abnormal lumen shape were observed (Fig. 3E and F). The most impressive symptoms appeared in the section from shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 28 d (Fig. 3G and I). Tubular epithelial cells had detached from the basal lamina. The tubules were seriously atrophied, and some epithelial cells were hypertrophied and vacuolated, so that the tubules looked like they were compressed and the lumen-facing surface was oppilated and irregular (Fig. 3H). Furthermore, melanization was observed in the epithelial cells, especially in the periphery of the tubules (Fig. 3I).

No obvious alterations were observed by light microscopy of the hepatopancreatic tissue of treated \(L.\) \textit{vannamei} after exposure to 0.1 mg Cd L\(^{-1}\) for 14 or 21 d, 0.2 mg Cd L\(^{-1}\) for 14 or 21 d, or 0.05 mg Zn L\(^{-1}\) for 14, 21, or 28 d in this study, except for one deeply affected individual exposed to 0.1 mg Cd L\(^{-1}\) for 21 d, which was moribund when sampled.

### 3.2. Metal concentrations in the hepatopancreas of \(L.\) \textit{vannamei}

Metal concentrations accumulating in the hepatopancreas of \(L.\) \textit{vannamei} were determined (Tables 1 and 2). The hepatopancreatic Cd concentrations were undetectable in control animals at each sampling time (Table 1). After exposure to 0.1 mg Cd L\(^{-1}\), however, the hepatopancreatic Cd concentration was 0.07 ± 0.03 mg g\(^{-1}\) after 7 d of exposure and significantly increased to 0.19 ± 0.04 mg g\(^{-1}\) after 14, 21, and 28 d, respectively. A similar increasing trend was also observed in 0.2 mg Cd L\(^{-1}\)-treated animals, and the hepatopancreatic Cd concentrations were 0.18 ± 0.09, 0.37 ± 0.07, 0.38 ± 0.05, and 0.52 ± 0.18 mg g\(^{-1}\), respectively, after 7, 14, 21, and 28 d of exposure.

The hepatopancreatic zinc concentration in control shrimp was 0.19 ± 0.04 mg g\(^{-1}\) (Table 2). No significant difference was apparent in hepatopancreatic zinc concentrations of 0.05 mg Zn L\(^{-1}\)-treated animals compared to the control. In the 0.2 mg Zn L\(^{-1}\)-treated shrimp, the hepatopancreatic zinc concentration significantly increased to 0.31 ± 0.04 and 0.34 ± 0.06 mg g\(^{-1}\) (\(p < 0.05\)) after exposure for 7 and 14 d, but then declined to levels that showed no significant difference compared to those of the control after exposure for 21 and 28 d.

### 3.3. Effects of Cd and Zn on hemolymphatic GOT and GPT activities of \(L.\) \textit{vannamei}

According to the results of the enzyme activity determinations, hemolymphatic GOT and GPT activities of control \(L.\) \textit{vannamei} were 19.8 ± 4.4 and 14.7 ± 4.9 U L\(^{-1}\), respectively. Exposure of \(L.\) \textit{vannamei} to Cd for 28 d resulted in increased hemolymphatic GOT and GPT activities, according to our observations (Table 1). After exposure for 7, 14, and 28 d, the average levels of GOT activities of Cd-treated \(L.\) \textit{vannamei} were 180.3%, 264.4%, and 218.7% higher compared to the average level of the control animals. Similarly, the average levels of GPT activities of individuals exposed to Cd for 7, 14, and 28 d were 352.4%, 337.1%, and 381.0% higher, respectively. Although both hemolymphatic GOT and GPT activities also increased after shrimp were exposed to Zn for 7 d and were 361.4% and 521.4% higher than those of control individuals, declines in both enzyme activities were observed and no significant differences were evident from those of the control group after exposure for 14 and 28 d (Table 2).

### 4. Discussion

The liver and hepatopancreas are known to be very sensitive to different diets and water-borne pollutants; thus these organs are often used to monitor the effects of various toxicants (Bautista et al., 1994). The hepatopancreas is essentially composed
of branched tubules and different types of epithelial cells lining the tubules. Therefore, it is likely that exposure to noxious chemicals or xenobiotics would be reflected in alterations to the structures of tubules and epithelial cells (Bhavan and Geraldine, 2000). The effects of exposure to various toxicants on histological and cellular changes to the liver and hepatopancreas of several aquatic organisms have been investigated (Lightner et al., 1982; Doughtie and Rao, 1984; Förlin et al., 1986; Khangarot, 1992; Bautista, 1994; Lightner et al., 1996; Bhavan and Geraldine, 2000). The effects on the hepatopancreas of several prawn species of aflatoxin, naturally produced by the fungi, Aspergillus flavus and Aspergillus parasiticus, have been studied (Lightner et al., 1982; Bautista, 1994), and it is thought to possess toxicity to the liver of higher organisms. Aflatoxin-induced structural changes in the hepatopancreas of prawns included a decrease in the cellular height of the tubular epithelium, a reduction in the abundance of secretory and lipid vacuoles, infiltration of hemocytes, atrophy of epithelial cells, development of pyknotic nuclei, cytolysis, the formation of fibrosis, and the melanized encapsulation of necrotic tissues. Likewise, similar studies have

Fig. 2. Hepatopancreas from cadmium (Cd)-treated Litopenaeus vannamei. (A) The hepatopancreas from a shrimp exposed to 0.2 mg Cd L⁻¹ for 28 d. Bar = 50 μm. (B) The hepatopancreas from one deeply affected and moribund individual exposed to 0.1 mg Cd L⁻¹ for 21 d. Bar = 50 μm. Both (A) and (B) show that large numbers of vacuoles appeared in the tubular epithelial cells, and the lumina are abnormal. The slight thickening of the basal lamina is obvious. (C) The hepatopancreas from a shrimp exposed to 0.4 mg Cd L⁻¹ for 14 d showing that the tubular epithelial cells are heavily vacuolated; and some are ruptured. Bar = 200 μm. (D) The same section shown in (C), but at a higher magnification. Bar = 50 μm. (E) Longitudinal section of the hepatopancreas from a shrimp exposed to 0.4 mg Cd L⁻¹ for 14 d. In this section, the lumen appears scraggly due to cellular hypertrophy as indicated by the arrow. Bar = 50 μm. (F) The hepatopancreas from a shrimp exposed to 0.4 mg Cd L⁻¹ for 21 d. Arrangement of the tubules and the shape of the lumina have substantially been altered. Infiltration of hemocytes and necrosis of tubules appeared. Bar = 200 μm. (G) Further examination at a higher magnification showing that necrotic tubules and cells appeared in the hepatopancreas of a shrimp exposed to 0.4 mg Cd L⁻¹. Necrotic tubules are rounded off and have sloughed off the basement membrane. Bar = 50 μm. ALU, abnormal lumen; BL, basal lamina; IHc, infiltration of hemocytes; NC, necrosis of epithelial cells; NT, necrosis of tubules; REc, ruptured epithelial cells.
Fig. 3. Hepatopancreas from zinc (Zn)-treated *Litopenaeus vannamei*. (A) The hepatopancreas from a shrimp exposed to 0.2 mg Zn L\(^{-1}\) for 14 d. Bar = 50 \(\mu\)m. (B) The hepatopancreas from a shrimp exposed to 0.2 mg Zn L\(^{-1}\) for 28 d. Bar = 50 \(\mu\)m. Both (A) and (B) show that large numbers of vacuoles appeared in the tubular epithelial cells, and the lumens are abnormal. (C) The hepatopancreas from a shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 14 d. Bar = 200 \(\mu\)m. (D) The hepatopancreas from a shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 21 d. Bar = 200 \(\mu\)m. In sections shown in (C) and (D), slight to moderate vacuolization of the tubular epithelial cells is apparent. (E) The hepatopancreas from a shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 21 d. In this section, a decrease in epithelial cell height as well as cell rupture resulting in the formation of an abnormal lumen are clearly evident. Bar = 50 \(\mu\)m. (F) A longitudinal section of the hepatopancreas from a shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 21 d. Note that the lumen of the tubules is abnormal due to vacuolization and hypertrophy of the epithelial cells. Bar = 200 \(\mu\)m. (G) The hepatopancreas from a shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 28 d. Tubular epithelial cells have detached from the basal lamina with this treatment, and the tubules have atrophied. Bar = 200 \(\mu\)m. (H) A longitudinal section of the hepatopancreas from a shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 28 d. Note that the lumen-facing surface is almost completely opiplated and irregular. Bar = 200 \(\mu\)m. (I) The hepatopancreas from a shrimp exposed to 0.6 mg Zn L\(^{-1}\) for 28 d. Melanization of the epithelial cells has appeared. Bar = 50 \(\mu\)m. ALU, abnormal lumen; BL, basal lamina; Mel, melanization of cells; REc, ruptured epithelial cells.
also examined the effects of pesticides and fungicides on other decapods (Lightner et al., 1996; Bhavan and Geraldine, 2000).

Förlin et al. (1986) and Khangarot (1992) studied the effects of the heavy metals, copper (Cu) and Cd, on ultrastructural changes in the liver of a teleost. Major alterations of liver cells after fish were exposed to Cu or Cd included proliferation of smooth endoplasmic reticula (SER), fragmentation and dilation of cisternae of rough endoplasmic reticula (RER), detachment of ribosomes from endoplasmic reticula (SER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertrophy of Golgi complexes, swelling of mitochondria with an amorphous, dense and intramaterial plasmic reticula (ER), hypertroph...
that it might have been due to production of collagen and walling off by hemocytes and also represented a defensive reaction against the toxicity of xenobiotics (Bhavan and Geraldine, 2000). Among the different types of cells in the tubular epithelia of the hepatopancreas, R-cells have been found to be the most readily and severely affected (Vogt, 1990; Bautista, 1994). R-cells are characterized by numerous vacuoles and lipid droplets appearing within the cytoplasm, and they are considered to be the major lipid reserve in the hepatopancreas. We found that a common feature of the hepatopancreas of Cd- and Zn-exposed L. vannamei was the appearance of moderate to heavy vacuolization in tubular epithelial cells. Since xylene was used when we stained the tissue paraffin sections with hematoxylin and eosin, it is possible that lipids within the lipid droplets or lipid-containing vacuoles may have been dissolved away, and the resulting tissue section showed only the existence of hollow vacuoles of cells. It was reported in M. malcolmsonii that the number of R-cells in the tubular epithelium of the hepatopancreas of test prawns exposed to 10.6 ng L\(^{-1}\) colmsonii that the number of R-cells in the tubular epithelium of Cd- and Zn-exposed L. vannamei was the appearance of moderate to heavy vacuolization in tubular epithelial cells. Since xylene was used when we stained the tissue paraffin sections with hematoxylin and eosin, it is possible that lipids within the lipid droplets or lipid-containing vacuoles may have been dissolved away, and the resulting tissue section showed only the existence of hollow vacuoles of cells. It was reported in M. malcolmsonii that the number of R-cells in the tubular epithelium of the hepatopancreas of test prawns exposed to 10.6 ng L\(^{-1}\) colmsonii that the number of R-cells in the tubular epithelium of Cd- and Zn-exposed L. vannamei was the appearance of moderate to heavy vacuolization in tubular epithelial cells. Since xylene was used when we stained the tissue paraffin sections with hematoxylin and eosin, it is possible that lipids within the lipid droplets or lipid-containing vacuoles may have been dissolved away, and the resulting tissue section showed only the existence of hollow vacuoles of cells. It was reported in M. malcolmsonii that the number of R-cells in the tubular epithelium of the hepatopancreas of test prawns exposed to 10.6 ng L\(^{-1}\) colmsonii that the number of R-cells in the tubular epithelium of Cd- and Zn-exposed L. vannamei was the appearance of moderate to heavy vacuolization in tubular epithelial cells. Since xylene was used when we stained the tissue paraffin sections with hematoxylin and eosin, it is possible that lipids within the lipid droplets or lipid-containing vacuoles may have been dissolved away, and the resulting tissue section showed only the existence of hollow vacuoles of cells. It was reported in M. malcolmsonii that the number of R-cells in the tubular epithelium of the hepatopancreas of test prawns exposed to 10.6 ng L\(^{-1}\) colmsonii that the number of R-cells in the tubular epithelium of Cd- and Zn-exposed L. vannamei was the appearance of moderate to heavy vacuolization in tubular epithelial cells. Since xylene was used when we stained the tissue paraffin sections with hematoxylin and eosin, it is possible that lipids within the lipid droplets or lipid-containing vacuoles may have been dissolved away, and the resulting tissue section showed only the existence of hollow vacuoles of cells. It was reported in M. malcolmsonii that the number of R-cells in the tubular epithelium of the hepatopancreas of test prawns exposed to 10.6 ng L\(^{-1}\) colmsonii that the number of R-cells in the tubular epithelium of Cd- and Zn-exposed L. vannamei was the appearance of moderate to heavy vacuolization in tubular epithelial cells.


