Histopathological changes in liver and gill epithelium of Nile tilapia, *Oreochromis niloticus*, exposed to waterborne copper

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**ABSTRACT.** Figueiredo-Fernandes A., Ferreira-Cardoso J.V., Garcia-Santos S., Monteiro S.M., Carrola J., Matos P. & Fontainhas-Fernandes A. 2007. Histopathological changes in liver and gill epithelium of Nile tilapia, *Oreochromis niloticus*, exposed to waterborne copper. *Pesquisa Veterinária Brasileira* 27(3):103-109. University of Trás-os-Montes and Alto Douro, Center of Studies on Technological of Environmental and Life Sciences (CETAV), Apartado 1013, Vila Real 5001-801, Portugal. E-mail: fontain@utad.pt

Nile tilapia, *Oreochromis niloticus*, of both sexes were reared in freshwater and exposed to 0.5, 1.0 and 2.5 mg L⁻¹ of waterborne copper for a period of 21 days. Liver and gill samples were collected after 21 days of exposure to copper and lesions were analyzed by light microscopy. The main histopathological changes observed in gills exposed to the highest concentration were edema, lifting of lamellar epithelia and an intense vasodilatation of the lamellar vascular axis. Although less frequent, lamellar fusion caused by the filamentar epithelium proliferation and some lamellar aneurisms were also found. The liver of control group exhibited a quite normal architecture, while the fish exposed to copper showed vacuolization and necrosis. These hepatic alterations were more evident in fish exposed to 1.0 and 2.5 mg L⁻¹ copper concentrations. The number of hepatocytes nucleus per mm² of hepatic tissue decreased with the increase of copper concentration. In contrast, the hepatic somatic index was high in fish exposed at 2.5 mg L⁻¹ of copper. In short, this work advance new knowledge as influence of copper in the gill and liver histology of *O. niloticus* and demonstrated that their effects could be observed at different concentrations.

**INDEX TERMS:** Copper, gill, liver, histopathological changes, *Oreochromis niloticus*, Teleostei.
INTRODUCTION
Copper is an essential trace metal in small concentrations for several fish metabolic functions. Essentiality of copper arises from its specific incorporation into a variety of enzymes which play important roles in physiological processes (e.g. enzymes involved in cellular respiration, free radical defence, neurotransmitter function, connective tissue biosyntheses and other functions), as well as, into some structural proteins (WHO 1998).

Although the crucial role of copper in several enzymatic processes (Baker 1969, Li et al. 1998), this heavy metal can exert adverse toxicological effects, when present in high concentrations in water (Pelgrom et al. 1995). In fact, it is potentially toxic when the internal available concentration exceeds the capacity of physiological detoxification processes.

Increasing agricultural production has resulted in increasing number of freshwater systems being impacted by the contaminants present in wastewater releases. In Portugal copper has been used in viticulture to control fungal diseases in vineyard plants. High concentrations of this heavy metal were detected in some aquatic ecosystems collecting vineyard runoff water and it is also highly concentrated in ground water (Gerbe 1996, Teisseire 1999). There are also anthropogenic sources of environmental contamination by copper including mining, smelting, foundries, municipal waste incinerators, burning of coal for power generation and a variety of copper-based products used in building and construction (Nor 1987, WHO 1998).

Heavy metal contamination has been reported in aquatic organisms (Adham et al. 2002, Olojo et al. 2005). These pollutants build up in the food chain and are responsible for adverse effects and death in the aquatic organisms (Farkas et al. 2002). Fish are widely used to evaluate the health of aquatic ecosystems and physiological changes serve as biomarkers of environmental pollution (Kock et al. 1996). Nile tilapia, Oreochromis niloticus, is one of the most common freshwater fish used in toxicological studies (Figueiredo-Fernandes et al. 2006a,b, Garcia-Santos et al. 2006), because it present a number of characteristics that may make it an appropriate model that can be used as indicator species in biomonitoring programmes (Gadagbui et al. 1996).

When exposed to toxic concentrations, organs of aquatic animals may accumulate copper (Pelgrom et al. 1995, Grosell et al. 1996, Mazon et al. 2002), which can lead to redox reactions generating free radicals and, therefore, may cause biochemical and morphological alterations (Varanka et al. 2001, Monteiro et al. 2005). Gills are the first target of waterborne pollutants due to the constant contact with the external environment, as well as the main place for copper uptake (Campbell et al. 1979, Perry & Laurent 1993). It is well known that changes in fish gill are among the most commonly recognized responses to environmental pollutants (Mallatt 1985, Laurent & Perry 1991, Au 2004).

The liver was examined because it plays a primary role in the metabolism and excretion of xenobiotic compounds with morphological alterations occurring in some toxic conditions (Rocha & Monteiro 1999). Metals can either increase or decrease hepatic enzyme activities and can lead to histopathological hepatic changes, depending on the metal type and concentration, fish species, length of exposure and other factors (Paris-Palacios et al. 2000). The monitorization of histological changes in fish liver is a highly sensitive and accurate way to assess the effects of xenobiotic compounds in field and experimental studies.

Hence, this study was undertaken to examine the effect of different sublethal copper sulphate concentrations on histological aspects of gill and liver of Nile tilapia, Oreochromis niloticus.

MATERIALS AND METHODS
Fish and experimental system
Nile tilapia, Oreochromis niloticus, Teleostei (Bouaké strain) were originally obtained from the Instituto Nacional de Recherche Agronomique (Rennes, France) and raised in the Aquaculture Station of the University of Trás-os-Montes and Alto Douro (UTAD, Vila Real, Portugal) for three generations. The fish used for this experiment were maintained in 100 L recirculating tanks, filled with dechlorinated tap water (pH 6.5-7.5; alkalinity 60mg L⁻¹ as HCO₃⁻; conductivity 63µS/cm⁻¹; Na⁺, 14mg L⁻¹; K⁺, 2.3mg L⁻¹; Ca²⁺, 4.1mg L⁻¹; Mg²⁺, 6.5mg L⁻¹; Cl⁻, 19.5mg L⁻¹; NO₃⁻, 27mg L⁻¹; NO₂⁻, 0.5mg L⁻¹). Fish were fed daily to satiation with a previous tested diet (Fontainhas-Fernandes et al. 1999), kept at a constant temperature of 25 ± 1°C and controlled photoperiod (12D: 12L). Supplemental aeration was provided to maintain dissolved oxygen near saturation.

Sexually mature tilapia O. niloticus (35.3 ± 5.4g of mean body weight) were randomly distributed through 12 tanks of 100 L. There were 3 fish per tank and 3 tanks for each treatment in a total of n = 36. Fish from 3 tanks containing water without copper served as the control group. Fish from the remaining tanks were exposed to water copper concentrations of 0.5, 1.0 and 2.5mg L⁻¹, supplied as copper sulphate (CuSO₄; MERCK, Lisbon, Portugal), during 21 days. The copper concentrations were selected based on preliminary results, shown to be sublethal after a 21 day period of exposure. The experiments were carried out under constant temperature (25 ± 1°C), controlled photoperiod (12D: 12L) and constant filtration. The water had identical physical and chemical characteristics of the acclimation tanks and the experiments described comply with the Guidelines of the European Union Council (86/609/EU). Both control and experimental tanks were submitted to a rate of water renewal of 1/3 every two days. The water quality parameters mentioned above were assessed in the experimental period, with no significant changes being observed. During the experimental period fish were once daily fed the referred diet to visual satiation and were fasted for 24 h before sampling.

No fish mortality was observed during the experiment. Six fish per treatment (2 fish per tank) were anaesthetized with 2-phenoxyethanol (Sigma, Barcelona, Spain) (1ml L⁻¹ water), weighed and sampled. Gills and livers were collected and weighed at 21 days of exposition.

Histology
A gill arch of the right side of each fish was collected and fixed in Bouin’s fluid for 24 h, dehydrated in graded ethanol concentrations and embedded in paraffin wax. Sagittal sections (5µm of thickness) were cut and mounted on glass slides. Sections were deparaffinized in xylene, hydrated in ethanol and stained with hematoxylin-eosin (HE). The liver was quickly dissected, sliced into
Histopathological changes in liver and gill epithelium of Nile tilapia, Oreochromis niloticus, exposed to waterborne copper

3 mm thick slabs, and immersed in Bouin’s fixative for 24 h, dehydrated, and embedded in paraffin; a minimum of 5 pieces resulted. Histological sections (5 µm of thickness) were cut and stained with H&E. Changes induced by treatment in the gill and liver tissues were photographed and analyzed by light microscopy (Nikon® Labophot).

The hepatosomatic index (HSI) was calculated and the number of hepatocytes per mm² of hepatic tissue was obtained in 12 microscopic fields (100x). Means ± standard deviation (SD) were calculated for each experimental group. Statistical differences between exposed groups and respective control group were analysed using ANOVA and multiple comparison by Student-Newman-Keuls test, at a 5% significant level. All tests were performed using the software STATISTICA, version 6.0 (StatSoft Inc. 2001).

RESULTS

The gill morphology of the control tilapia is similar to that of other teleost fish species (Wilson & Laurent 2002). The gill is made up of double rows of filaments from which arise perpendicularly the lamellae. The lamellae are lined by a squamous epithelium composed by pavement and non-differentiated cells. Below that epithelium are lamellar blood sinuses separated by pillar cells. Between the lamellae, the filament is lined by a thick stratified epithelium constituted by several cellular types, such as chloride, mucous and pavement cells (Fig. 1A). Fish showed some signs of epithelial lesions when exposed to the highest concentration, 2.5 mg L⁻¹ CuSO₄ (Fig. 1B, C, D, E). The main changes observed after 21 days included:

- Intense lamellar epithelium lifting (Fig. 1B).
- Proliferation of filamentar epithelium with fusion of adjacent lamellae (Fig. 1D).
- Vascular congestion or lamellar aneurisms (Fig. 1E). cc = chloride cell, cvs = central venous sinus, fe = filament epithelium, pc = pillar cell, pv = pavement cell.

Fig. 1. Representative light micrographs of gills in control and copper treated (2.5 mg L⁻¹ CuSO₄, 21 days) tilapia, Oreochromis niloticus. (A) Control fish, showing normal appearance of gill filaments (F) and lamellae (L). (B) Gills from exposed fish showing an intense lamellar epithelium lifting (Lf). Note the epithelium proliferation in the above filament (Fp). (C) Section of gill with lamellar axis vasodilation (v) and evident epithelium interstitial edema (**) in the filament near the lamellar axis. (D) Proliferation of filamentar epithelium (Fp) with fusion of adjacent lamellae (Lfu). (E) Gill epithelium of treated fish showing vascular congestion or lamellar aneurisms (A). cc = chloride cell, cvs = central venous sinus, fe = filament epithelium, pc = pillar cell, pv = pavement cell. HE, bars = 20 µm.
days of exposure were accentuated lifting of the lamellar epithelium (Fig.1B), edema in the filamentar epithelium and an intense lamellar vasodilation (Fig.1C). The gills of some fish also exhibited lamellar fusion in numerous areas (Fig.1D) as a result of filamentar epithelium proliferation (Fig.1B, D). In addition, a few aneurisms were also observed at gill lamellae (Fig.1E).

Liver histology from control and exposed fish is briefly illustrated in Fig.2. In the control group, the liver exhibited a normal architecture and there were no pathological abnormalities, with hepatocytes presenting a homogenous cytoplasm, and a large central or subcentral spherical nucleus (Fig.2A). The hepatic parenchyma of fish exposed to copper showed lower eosinophilia and an increase of cytoplasmatic vacuolation (Fig.2B, C). Additionally, tilapias exposed to copper also showed hepatocellular necrosis (Fig.2C). These liver histological alterations were more evident in fish exposed to 1.0 and 2.5mg L\(^{-1}\) copper concentrations. The HSI increased with the copper concentration (Table 1). However, the number of hepatocytes nucleus per mm\(^2\) of hepatic tissue decreased with the increase of copper concentration (Table 1).

**DISCUSSION**

Histological study of the gills shows a typical structural organization of the lamella in the untreated fish. However, fish exposed to copper shows several histological alterations, namely lamellar epithelium lifting, epithelium proliferation, lamellar axis vasodilation, edema in the filament, fusion of lamellae and lamellar aneurisms.

Some studies revealed that interstitial edema is one of the more frequent lesions observed in gill epithelium of fish exposed to heavy metals (Mallatt 1985). The results of this study confirm the occurrence of edema independently of copper levels, as in other fish species (Sola et al. 1995, Bury et al. 1998). The lifting of lamellar epithelium is other histological change observed, probably induced by the incidence of severe edema (Arellano et al. 1999, Pane et al. 2004, Schwaiger et al. 2004).

Edema with lifting of lamellar epithelium could be serve as a mechanism of defense, because separation epithelial of the lamellae increases the distance across which waterborne pollutants must diffuse to reach the bloodstream (Arellano et al. 1999). These gill histological alterations has been observed by several authors in fish submitted to copper (Karan et al. 1998, Chen & Lin 2001, De Boeck et al. 2001). However, these changes also can be due to the exposition to different kinds of pollutants, such as endosulfan (Nowak 1992), arsenic (Hwang & Tsai 1993), drugs (Schwaiger et al. 2004) and other.

<table>
<thead>
<tr>
<th>[CuSO(_4)] (mg.L(^{-1}))</th>
<th>HSI</th>
<th>Hepat.nucl.mm(^{-2})</th>
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<tr>
<td>0</td>
<td>0.98 ± 0.0008(^a)</td>
<td>15484.76 ± 1023.97(^a)</td>
</tr>
<tr>
<td>0.5</td>
<td>1.44 ± 0.0011(^b)</td>
<td>5476.29 ± 1376.93(^b)</td>
</tr>
<tr>
<td>1.0</td>
<td>1.52 ± 0.0023(^b)</td>
<td>3096.82 ± 747.07(^c)</td>
</tr>
<tr>
<td>2.5</td>
<td>2.02 ± 0.0023(^c)</td>
<td>1482.27 ± 669.93(^d)</td>
</tr>
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\(^a\) Values are expressed as means ± SE (n=6). Means in the same column with different letters are significantly different (ANOVA, P<0.05).

**Fig.2.** Photomicrographs of Nile tilapia *Oreochromis niloticus* liver tissue. (A) Control group showing hepatocytes (he) and pancreatic area (pa) that corresponds to the acini of exocrine pancreas. (B) Liver of fish exposed to copper (1 mg.l\(^{-1}\), showing alterations in hepatocytes and vacuolation (black arrows); bv, blood vessel. (C) Liver of fish exposed to copper (2.5 mg.l\(^{-1}\), showing vacuolation (black arrows) and necroses area (*) and picnotic nucleus (black arrow).HE, bars = 50μm
heavy metals, as aluminium (Karlsson-Norggren et al. 1986), cadmium (Reid & McDonald 1988), and nickel (Pane et al. 2004). Thus, this signifies that these alterations are not specifically induced by copper or other heavy metals.

Cell proliferation with thickening of gill filament epithelium is one histological change found in fish exposed to copper by several authors (Arellano 1999, De Boeck et al. 2001, van Heerden et al. 2004), and may lead to the lamellar fusion observed in this study. These results also were found in fish exposed to other pollutants (Randi et al. 1996, Van den Heuvel et al. 2000, Rosety-Rodriguez et al. 2002). The edema, epithelial lifting as well as lamellar fusion also are defence mechanisms that reduce the branchial superficial area in contact with the external milieu. These mechanisms also increase the diffusion barrier to the pollutant (Lauren & McDonald 1985, Van Heerden et al. 2004).

Lamellar axis vasodilatation was also found in tilapia exposed to copper. Garcia-Santos et al. (2006) refer that this lesion can induce changes in pillar cell normal structure, with consequent loss of their support function and probably, and was responsible for the emergence of lamellar aneurysms in fish exposed to cadmium. Similar results are observed by Thophon et al. (2003) in Lates calcarifer exposed to cadmium. However, Mallat (1985) suggests that these lesions are rarely associated to metals exposition.

The biological parameters are sometimes indicative of toxicant effects (Mayer et al. 1992). Our results reveal that the HSI increased with copper concentration. Figueiredo-Fernandes et al. (2006b) also found an increase of HSI in male and female tilapia, Oreochromis niloticus, exposed to paraquat. In contrast, rainbow trout Oncorhynchus mykiss injected with paraquat showed a decrease in the HSI after 9 weeks, however histological results were not shown (Ækerman et al. 2003).

The qualitative liver histology showed an increase on the hepatocytes size in tilapia exposed to copper that may be due to the high content of lipids. Huuskozen & Lindstrom-Seppa (1995) and Stephensen et al. (2000) found that the high HSI observed in the perch (Perca fluviatilis) and sculpin (Myoxocephalus scorpius) can be indicative of increased activity of xenobiotic biotransformation enzymes. Figueiredo-Fernandes et al. (2006b) also suggested a positive relationship between the relative liver weight and the xenobiotic-metabolizing enzymes of tilapia exposed to paraquat.

The present work also shows that the number of hepatocytes nucleus in hepatic tissue decrease with the copper concentration. Several studies demonstrated that alterations in number, size and shape of the hepatocyte nucleus can be due to contaminants. Alterations in the size of nucleus have been previously regarded by Paris-Palacios et al. (2000) in Brachydanio rerio exposed to sublethal concentrations of copper sulphate. Braunbeck et al. (1990) referred that alterations in size and shape of nucleus have often been regarding as signs of increased metabolic activity but may be of pathological origin.

In a previous study the normal morphology of the liver of Nile tilapia, Oreochromis niloticus (Figueiredo-Fernandes et al. 2006c) was characterized, and it was shown that liver was made up of hepatocytes that were not oriented into distinct lobules but were arranged in branched laminae two cells thick, separated by sinusoids. Hepatocytes were polygonal cells with a central spherical nucleus and a densely stained nucleolus. The present study also demonstrates that the liver of control fish exhibits a normal architecture and there were no pathological abnormalities. The hepatocytes present a homogenous cytoplasm and a large central or subcentral spherical nucleus.

The histology showed that copper caused some alterations of the liver parenchyma, like vacuolization and necrosis. The liver histological changes observed were more evident in fish exposed to high copper concentrations. These alterations are often associated with a degenerative-necrotic condition (Myers et al. 1987). Several studies had shown a variety of changes in the liver of Oreochromis niloticus, resulting from exposure to different toxic chemicals (Visoottiviseth et al. 1999, Figueiredo-Fernandes et al. 2006a,b). Moreover, it was also reported by several studies that chronic copper accumulation in the liver of fish causes hepatocyte lysis, cirrhosis and ultimately death (Pourahamad & O’Brien 2000, Varanka et al. 2001).

In the present study gill and liver histological changes have been related to copper concentrations. It can be concluded that gill and hepatic alterations as a result of heavy metal exposition of fish may serve as a sensitive biomarker for the toxicity of sublethal concentrations of metals as well as other pollutants. However, complementary studies are necessary for a better understanding of its deleterious effects.

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