

Original Research Articles

Copper toxicity in juvenile largemouth bass (*Micropterus salmoides*): acute toxicity bioassay and oxidative stress response in gut and kidney

Junhao Zhang¹, Na Zhao², Zihao Meng¹, Mengkang You¹, Ke Wang¹, Songlin Dai¹, Zhenyang Zhang¹, Yuchao Huang¹, Weijun Chen¹, Shiyang Gao^{1a}

¹ College of Animal Science and Technology, Henan University of Science and Technology, ² College of Agricultural Equipment Engineering, Henan University of Science and Technology

Keywords: Toxicity test, Copper sulfate, Largemouth bass, Oxidative stress, Static bioassay

https://doi.org/10.46989/001c.116377

Israeli Journal of Aquaculture - Bamidgeh

Vol. 76, Issue 2, 2024

Aquaculture intensification has resulted in serious disease outbreaks in largemouth bass production. Compounds containing copper (Cu) are widely used as therapeutic agents in aquaculture. Currently, Cu misuse has been a severe issue in largemouth bass farming. However, few investigations have been performed on Cu toxicity in largemouth bass so far. In this study, an acute and a chronic toxicity test was carried out to determine the toxicity and the recommended dose of waterborne Cu in largemouth bass. In the acute toxicity bioassay, fish $(2.58 \pm 0.03 \text{ g})$ were exposed to 0 (control), 3, 6, 9, 18, or 30 mg/L Cu, and the results showed that the 96-h LC₅₀ of waterborne Cu was 12.78 mg/L. Then a 30-day chronic toxicity test containing six treatments (i.e., 0, 51.3, 164, 513, 1640, and 5130 µg/L Cu) was conducted to investigate the influence of Cu on intestinal and renal health in terms of oxidative stress in juvenile largemouth bass (2.69 \pm 0.02 g). The results showed that Cu concentrations at and above 51.3 µg/L significantly increased the malondialdehyde contents (in the intestine) and simultaneously decreased total superoxide dismutase activity levels (in the intestine and kidney), glutathione peroxidase activity levels (in the kidney), and reduced glutathione contents (in the kidney), compared to control. In contrast to control, fish exposed to high Cu concentrations (at and above 1640 µg/L) demonstrated lower catalase activity levels in the intestine and kidney. Based on the findings in the study, waterborne Cu content for largemouth bass farming was recommended to be less than 51.3 µg/L.

INTRODUCTION

Largemouth bass is a freshwater carnivorous fish that is widely preferred by the farmers and consumers because of its high economic value and delicious meat. According to the statistics of FBMA, the production of largemouth bass in China has increased from 185941 tons in 2010 to 802486 tons in 2022. With the intensification of aquaculture, the immune system of largemouth bass has been severely weakened, resulting in outbreaks of bacterial, parasitic, and fungal diseases. To combat these problems, many therapeutic agents are applied to the water, and used in largemouth bass farming, including copper (Cu). A.5 Cu is widely used as an algaecide, fungicide, and parasiticide in aquaculture. Waterborne Cu has been widely used to treat external bacteria (e.g., columnaris disease), parasites (e.g.,

white spot disease), and fungal (e.g., water mold infection) diseases in largemouth bass. ^{3,4} However, this has led to Cu overuse, resulting in excessive Cu build-up in largemouth bass aquaculture. ³ It is well known that excessive Cu in water often causes various negative effects on aquatic animals, such as growth retardation, fish mortality, and tissue damage. ⁵ Based on this, it is particularly important to determine the toxicity and appropriate level of Cu in largemouth bass aquaculture water.

The toxicity test is an important method to determine the tolerance of animals to certain toxic substances and the appropriate dose of toxic substances in fish farming.⁷ The 96-h median lethal concentration (LC₅₀) trial is the most common approach in toxicity tests.⁸ The intestine and kidneys are important organs for the fish to maintain normal metabolism and growth. However, at present, there are

few studies on Cu toxicity and its effects on the intestinal and renal health of largemouth bass juveniles. Therefore, in this study, we intended to use the 96-h $\rm LC_{50}$ trial to assess the acute toxicity of Cu. Additionally, the relationship of Cu to intestinal and renal health of largemouth bass, as well as the recommended dose of Cu in aquaculture water, were studied from the perspective of oxidative stress using a chronic toxicity experiment. The results of this investigation will offer theoretical support for the healthy development of largemouth bass culture.

MATERIALS AND METHODS

EXPERIMENTAL FISH AND THE STOCK SOLUTION

Juvenile largemouth bass were provided by a fish farm in Guangzhou, China. Prior to the experiment, fish were exposed to laboratory settings for two weeks and were fed a commercial pelleted floating diet (Dongyufeng No.0, 48% crude protein, 5% crude fat, and 3 mg/kg Cu) (Zhejiang Dongyu Biological Technology Co., Ltd., Huzhou, China).

To create a stock solution of Cu (31.77 g/L), CuSO $_4$ ·5H $_2$ O (Sinopharm Chemical Reagent Co., Ltd, Shanghai, China) (124.84 g) was dissolved in distilled water using a volumetric flask (1000 mL). The Cu concentrations required for the experiment were achieved by diluting the stock solution with distilled water.

EXPERIMENTAL DESIGN

The 96-h acute waterborne Cu toxicity test was carried out in accordance with OECD guideline 203.9 180 experimental fish (2.58 ± 0.03 g) were evenly distributed to 18 tanks (20 L) and then subjected to different Cu concentrations: 0, 3, 6, 9, 18, and 30 mg/L (estimated Cu contents were 0.0 \pm 0.0, 2.8 \pm 0.2, 5.4 \pm 0.6, 10.2 \pm 0.8, 16.3 \pm 1.7, and 27.2 \pm 2.4 mg/L, respectively). Each treatment (Cu concentration) had three duplicates. To maintain the waterborne Cu concentrations, fresh water that has been aerated and has the appropriate Cu content was added to the system on a regular basis. Throughout the acute toxicity test, the fish were not fed, and dead fish were taken out of the fish tank every day. Death was defined as the absence of gill movement and a reaction to mild prodding. The 96-h LC₅₀ was calculated using Finney's Probit analysis method¹⁰ based on the fish death rates in each treatment.

In the exposure study, six Cu concentrations (0, 51.3, 164, 513, 1640, and 5130 µg/L) were selected, with the highest concentration being 40% of the 96-hour LC₅₀. ¹¹ To keep the Cu concentration as consistent as possible, each tank's water was refreshed and re-dosed on a daily basis using the stock solution. Mean Cu concentrations in tank water samples of 0, 51.3, 164, 513, 1640, and 5130 µg Cu/L treatments were 0.030 ± 0.011 , 50.5 ± 2.8 , 161.0 ± 3.1 , 519.3 ± 8.4 , 1630 ± 10.8 , and 5142.0 ± 42.6 µg/L, respectively. 270 largemouth bass (2.69 \pm 0.02 g) were randomly and evenly distributed among 18 tanks (20 L) (three tanks for each Cu concentration). The fish were fed by hand to apparent satiety twice (10:00 and 17:00) daily with the commercial pelleted float-

ing feed for 30 days. The photoperiod and water parameters throughout the experiment were as follows: photoperiod 12 h light/12 h dark, dissolved oxygen 8.0 \pm 1.2 mg/L, water temperature 25.0 \pm 0.5 °C, pH 8.2 \pm 0.4, ammonia less than 0.1 mg/L, alkalinity 157 \pm 17 mg/L, total hardness 166 \pm 6 mg/L, and conductivity 450 \pm 30 μ S/cm.

SAMPLING

On the 30th day of the experiment, all fish were starved for 24 h. Then 6 fish from each tank were sampled and anesthetized with 50 mg/L benzocaine. Intestine and kidney samples were collected on ice-cold plates, rapidly frozen in liquid nitrogen, and stored at -80 °C until analysis of oxidative-related parameters.

All of the studies were carried out in accordance with China's official protocol for the care and use of laboratory animals. The Institutional Animal Care and Use Committee at Henan University of Science and Technology approved both the study protocol and all experimental techniques.

MEASUREMENT OF CU CONTENT IN THE FEED AND WATER SAMPLES

The microwave digestion method was used to assess Cu contents in feed and water. 12 Samples (approximately 0.5 g for feed and 2 mL for water) were first digested by HNO_3 (70%) in the Berghof mws-3 microwave system. All digested samples were cooled to room temperature before being diluted to 10 mL with distilled water in 10 mL volumetric flasks. An atomic absorption spectrometer (PerkinElmer, Inc. Waltham, USA) was used to determine the Cu content at 324.8 nm.

ANALYSIS OF OXIDATIVE-RELATED PARAMETERS

Samples of intestine and kidney were homogenized with 0.68% physiological saline (tissue weight:saline weight = 1 g:9 mL). The homogenate was centrifuged at 1000 g for 10 min. Then the supernatant was collected and stored at 4 $^{\circ}$ C until the analysis of oxidative-related parameters.

Both contents of malondialdehyde (MDA), reduced glutathione (GSH), and total soluble protein and activity levels of total superoxide dismutase (T-SOD), catalase (CAT), and glutathione peroxidase (GPx) were determined by commercial kits (Nanjing Jiancheng Bioengineering Research Institute, Nanjing, China).

STATISTICAL ANALYSIS

The probit analysis in IBM SPSS Statistics 20.0 software was used to calculate the 96-h LC_{50} based on the fish mortality at the acute toxicity test. For the 30-day exposure test, all data were expressed as means \pm standard errors of means (SEM) and analyzed with one-way analysis of variance using SPSS 20.0. Duncan's multiple range test was used to examine mean differences at a significance level of 5%.

Table 1. Data of mortality rates of largemouth bass juveniles at the 96-h acute waterborne copper exposure test

Concentration (mg/L)	Number of fish exposed	Number of dead fish	Percentage mortality (%)
0	10	0.00	0
3.0	10	0.33	3.33
6.0	10	2.00	20.00
9.0	10	4.67	46.67
18.0	10	5.33	53.33
30.0	10	8.00	80

Table 2. Probit analysis of juvenile largemouth bass mortality at the acute waterborne Cu exposure test

	Number	Concentration (mg/L)	Number of subjects	Observed responses	Expected responses	Residual	Probability
	1	3.0	10	0	.532	202	.053
	2	6.0	10	2	1.798	.202	.180
Probit	3	9.0	10	5	4.052	.618	.405
	4	18.0	10	5	6.020	690	.602
	5	30.0	10	8	7.890	.110	.789

Table 3. The 96-h estimated lethal concentration of copper in juvenile largemouth bass

LC values	95% confidence level for concentration at 96 h				
	Estimate	Lower limit	Upper limit		
LC ₁₀	3.830	.980	6.222		
LC ₂₀	5.792	2.282	8.609		
LC ₃₀	7.806	4.031	11.330		
LC ₄₀	10.073	6.189	15.176		
LC ₅₀	12.783	8.599	21.427		
LC ₆₀	16.222	11.178	32.339		
LC ₇₀	20.932	14.102	52.714		
LC ₈₀	28.209	17.904	96.544		
LC ₉₀	42.665	24.248	229.717		

RESULTS

LETHAL CONCENTRATION OF COPPER IN LARGEMOUTH BASS

Table 1 shows the relationship between Cu concentrations and fish mortality at the 96-h acute waterborne Cu exposure test. In the control group, no dead fish were found. With the increase of waterborne Cu concentration, dead fish number and fish mortality rate increased. 80% of fish were dead when waterborne Cu concentration was 30 mg/L.

<u>Table 2</u> demonstrates the probit analysis of fish mortality at different Cu concentrations. Observed responses were close to expected responses (<u>Table 2</u>).

Table 3 shows the estimated LC values of waterborne Cu and their 95% upper and lower confidence limits at the 96-h acute toxicity test. The 96-h LC_{50} of waterborne Cu was 12.78 mg/L at the 95% confidence level, with upper and lower limits of 8.60 mg/L and 21.43 mg/L, respectively.

The regression line between fish mortality and the log values of waterborne Cu generated by the software SPSS 20.0 is shown in <u>Fig. 1</u>. R² linear =0.976 suggested that experimental data fitted the predicted regression very well.

OXIDATIVE RESPONSE IN THE INTESTINE

Cu concentrations over 51.3 μ g/L substantially increased MDA levels while decreased T-SOD activity levels when compared to the control (P < 0.05). GPx and GSH demonstrated similar changes. Compared to that in the unexposed group, GPx activity levels and GSH contents were significantly reduced when Cu concentrations were at and above 164 μ g/L (P < 0.05). In addition, fish in the groups with Cu levels greater than 513 μ g/L showed lower CAT levels than the control (P < 0.05) (Fig. 2).

Probit Transformed Responses 1.0 R² Linear = 0.976 0.5 0.0 -1.5 -2.0 0.50 0.75 1.00 1.25 1.50 Log of copper concentration

Figure 1. Regression line between the log concentration of copper and the probit mortality of juvenile largemouth bass

OXIDATIVE RESPONSE IN THE KIDNEY

Compared to the control, fish exposed to Cu concentrations at and above 51.3 μ g/L demonstrated lower T-SOD and GPx activity levels and GSH contents (P < 0.05). However, only high Cu concentrations induced significant changes in MDA contents (Cu at and above 513 μ g/L) and CAT activity levels

(Cu at and above 1640 μ g/L) in the kidney (P < 0.05) (Fig. 3).

DISCUSSION

In aquaculture, Cu is often used as algaecides, fungicides, and parasiticides. 13 However, excess Cu in water bodies has been shown to significantly reduce fish growth and health.¹⁴ Therefore, it is necessary to determine the appropriate Cu level in water for the healthy development of aquatic animals. The 96-h LC50 test is often used to assess the potential susceptibility and mortality of a species to toxic substances.^{6,9} In this study, the 96-h LC₅₀ assay was used to assess the tolerance of largemouth bass to waterborne Cu. Cu toxicity in water is affected by a variety of factors, such as the acclimation period of fish to Cu, water chemical parameters (e.g., water hardness, alkalinity, and pH), species and growth stage of the tested aquatic animals, and Cu types (e.g., copper nanoparticles and copper sulfate). 6 In general, an increase in the acclimation period to waterborne Cu, water alkalinity or hardness, and fish size will reduce the toxicity of Cu.⁶ In this experiment, the 96-h LC₅₀ for juvenile largemouth bass was 12.8 mg/ L, which was consistent with the results in juvenile Nile tilapia (Oreochromis niloticus) (12.0 mg/L)¹⁵ and juvenile Indian knife fish (Notopterus notopterus) (12.0 mg/L). However, in butterfish (Poronotus triacanthus) (0.5 mg/L for 96-h LC₅₀), ¹⁶ channel catfish (*Ictalurus punctatus*) (0.05 mg/L for 96-h LC₅₀),¹⁷ and rohu (*Labeo rohita*) (0.80 mg/L for 96-h LC_{50}), ¹⁸ the 96-h LC_{50} of waterborne Cu is ranged from 0.05 to 0.80 mg/L, which is much smaller than the results of

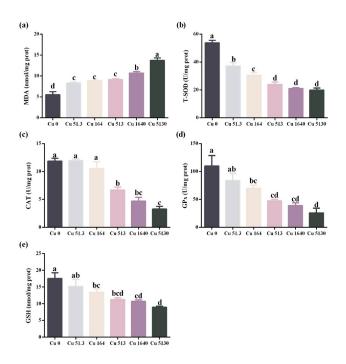


Figure 2. The impact of waterborne copper exposure on the intestinal oxidative state in juvenile largemouth bass.

(a) MDA, malondialdehyde; (b) T-SOD, total superoxide dismutase; (c) CAT, catalase; (d) GPx, glutathione peroxidase; (e) GSH, reduced glutathione. Means of exposures with different lowercase letters are significantly different at P < 0.05

the present study. Differences in water chemical parameters and fish species may account for this phenomenon.

The intestine and kidneys play critical roles in maintaining fish health. The intestine plays an important role in the digestion and absorption of nutrients, salt and water balance, and barrier function and immunology in fish (Dawood 2021). The kidney mainly functions in osmoregulation, acid-base regulation, and immunity in fish.¹⁹ Previous studies have shown that oxidative stress can damage intestinal and renal functions and threaten the health of aquatic animals.^{20,21} MDA is a typical marker of oxidative stress.²² In this study, MDA contents in the gut and kidney were significantly higher when Cu concentration was at and above 513 µg/L, indicating the occurrence of oxidative stress in these tissues. To avoid the negative effects of oxidative stress, fish develop antioxidant defense systems, including enzymatic (e.g., T-SOD, CAT, and GPx) and non-enzymatic antioxidants (e.g., GSH).²³ However, these indexes decreased significantly when Cu concentration was above 513 µg/L in the intestine and 1640 µg/L in the kidney, indicating that high waterborne Cu concentration reduced the antioxidant capacity of the gut and kidney. Consistent with our results, studies in Rhynchocypris lagowski²⁴ and grouper (Acrossocheilus fasciatus)²⁵ showed that high waterborne Cu exposure caused oxidative stress and decreased antioxidant capacity in the intestine and kidney. The causes of oxidative stress induced by Cu can be attributed to the following three points. Firstly, Cu is a transition metal and can act as a catalyst for the production of reactive oxygen species (e.g., superoxide anion and hydroxyl radical) through the Fenton reaction.²⁶ Secondly, Cu weakens fish antioxidant

defense systems by inhibiting antioxidant enzymes and depleting cellular GSH.²⁷ Thirdly, Cu inhibits the mitochondria electron-transfer chain and stimulates the formation of reactive oxygen species.²⁸ Based on the results of renal and intestinal oxidative stress, it is recommended that the Cu concentration in the cultured water for largemouth bass farming should be less than 51.3 µg/L.

CONCLUSION

The 96-h LC $_{50}$ of Cu in the form of CuSO $_4$ for largemouth juveniles (2.6 g) was 12.8 mg/L. Waterborne Cu exposure can increase oxidative stress and decrease the antioxidant capacity of the intestine and kidney in largemouth bass when Cu concentration is at and above 513 μ g/L. Based on the results of the present study, Cu content in the water for largemouth bass culture is recommended to be less than 51.3 μ g/L.

ACKNOWLEDGMENTS

This work was supported by the Doctoral Scientific Research Foundation of Henan University of Science and Technology (13480087; 13480088), the National Natural Science Foundation of China (NSFC, no. 32202952), Henan Provincial Science and Technology Research Project (222102320144), and the Innovation and Entrepreneurship Training Program for College Students in Henan University of Science and Technology (2023438, 2023442).

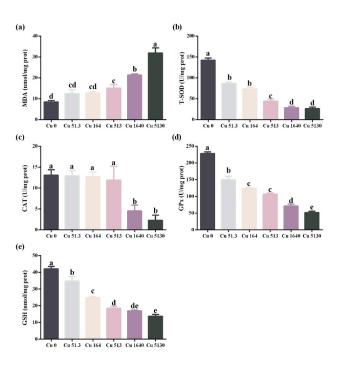


Figure 3. The impact of waterborne copper exposure on the oxidative state in the kidney of juvenile largemouth bass.

(a) MDA, malondialdehyde; (b) T-SOD, total superoxide dismutase; (c) CAT, catalase; (d) GPx, glutathione peroxidase; (e) GSH, reduced glutathione. Means of exposures with different lowercase letters are significantly different at P < 0.05

AUTHORS' CONTRIBUTION

Conceptualization: Junhao Zhang (Equal), Shiyang Gao (Lead). Methodology: Junhao Zhang (Equal), Na Zhao (Equal), Zihao Meng (Equal), Mengkang You (Equal). Writing – original draft: Junhao Zhang (Equal), Yuchao Huang (Equal). Resources: Na Zhao (Lead). Formal Analysis: Ke Wang (Equal), Songlin Dai (Equal), Zhenyang Zhang (Equal). Investigation: Ke Wang (Equal), Songlin Dai (Equal), Zhenyang Zhang (Equal). Writing – review & editing: Weijun Chen (Equal), Shiyang Gao (Lead). Funding acquisition: Weijun Chen (Equal), Shiyang Gao (Lead). Supervision: Shiyang Gao (Lead).

COMPETING OF INTEREST

No competing interests were disclosed.

ETHICAL CONDUCT APPROVAL - IACUC

All experiments and handling of the animals were conducted according to the research protocols approved by the Institutional Animal Care and Use Committee, Henan University of Science and Technology.

INFORMED CONSENT STATEMENT

All authors and institutions have confirmed this manuscript for publication.

DATA AVAILABILITY STATEMENT

Data are available upon reasonable request.

Submitted: March 04, 2024 CST. Accepted: March 26, 2024 CST. Published: April 29, 2024 CST.



This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International License (CCBY-4.0). View this license's legal deed at http://creativecommons.org/licenses/by/4.0 and legal code at http://creativecommons.org/licenses/by/4.0/legalcode for more information.

REFERENCES

- 1. Hussein GHG, Chen M, Qi PP, et al. Aquaculture industry development, annual price analysis and out-of-season spawning in largemouth bass Micropterus salmoides. *Aquaculture*. 2020;519:734901. doi:10.1016/j.aquaculture.2019.734901
- 2. FBMA. *China Fishery Statistical Yearbook*. China Agriculture Press; 2023.
- 3. Mitchell KM, Durborow RM. Common diseases of largemouth bass. In: *Largemouth Bass Aquaculture*. 5M Publishing Ltd; 2019:306-338.
- 4. Straus DL, Ledbetter CK, Heikes DL. Inhibiting fungus on largemouth bass eggs with copper sulfate and its toxicity to fry and juveniles. *J World Aquacult Soc.* 2020;51(1):214-223. doi:10.1111/jwas.12631
- 5. Gao S, Sun P, Zhao X, Chang K, Chen W. Waterborne copper exposure decreases fish growth and survival by promoting gills and liver impairments in largemouth bass (Micropterus salmoides). *Environ Sci Pollut Res.* 2023;30(56):119204-119216. doi:10.1007/s11356-023-30755-7
- 6. Tavares-Dias M. Toxic, physiological, histomorphological, growth performance and antiparasitic effects of copper sulphate in fish aquaculture. *Aquaculture*. 2021;535:736350. doi:10.1016/j.aquaculture.2021.736350
- 7. Sprague JB. Measurement of pollutant toxicity to fish I. Bioassay methods for acute toxicity. *Water Res.* 1969;3(11):793-821. doi:10.1016/0043-1354(69)90050-5
- 8. Su T, Lian D, Bai Y, et al. The feasibility of the zebrafish embryo as a promising alternative for acute toxicity test using various fish species: A critical review. *Sci Total Environ*. 2021;787:147705. doi:10.1016/j.scitotenv.2021.147705
- 9. OECD. Test No. 203: Fish, Acute Toxicity Test. In: *OECD Guidelines for the Testing of Chemicals, Section* 2. OECD Publishing; 2019. doi:10.1787/9789264069961-en
- 10. Finney DJ. *Probit Analysis*. Cambridge University Press; 1952.
- 11. Wong PPK, Chu LM, Wong CK. Study of toxicity and bioaccumulation of copper in the silver sea bream Sparus sarba. *Environ Int*. 1999;25(4):417-422. doi:10.1016/s0160-4120(99)00008-2

- 12. Sastre J, Sahuquillo A, Vidal M, Rauret G. Determination of Cd, Cu, Pb and Zn in environmental samples: microwave-assisted total digestion versus aqua regia and nitric acid extraction. *Anal Chim Acta*. 2002;462(1):59-72. doi:10.1016/s0003-2670(02)00307-0
- 13. Buley RP, Adams C, Belfiore AP, et al. Field evaluation of seven products to control cyanobacterial blooms in aquaculture. *Environ Sci Pollut Res.* 2021;28(23):29971-29983. doi:10.1007/s11356-021-12708-0
- 14. Qian D, Xu C, Chen C, Qin JG, Chen L, Li E. Toxic effect of chronic waterborne copper exposure on growth, immunity, anti-oxidative capacity and gut microbiota of Pacific white shrimp Litopenaeus vannamei. *Fish Shellfish Immunol*. 2020;100:445-455. doi:10.1016/j.fsi.2020.03.018
- 15. Khan A, Rehman H, Awais S, et al. Investigation of acute toxicity and LC50 value of Cu for a fish Oreochromis niloticus. *J Entomol Zool Stud*. 2016;4:605-607.
- 16. Jiraungkoorskul W, Sahaphong S, Kangwanrangsan N. Toxicity of copper in butterfish (*Poronotus triacanthus*): tissues accumulation and ultrastructural changes. *Environ Toxicol*. 2007;22(1):92-100. doi:10.1002/tox.20238
- 17. Straus DL, Tucker C s. Acute toxicity of copper sulfate and chelated copper to channel catfish Ictalurus punctatus. *J World Aquacult Soc.* 1993;24(3):390-395. doi:10.1111/j.1749-7345.1993.tb00170.x
- 18. Latif A, Ali M, Sayyed AH, et al. Effect of copper sulphate and lead nitrate, administered alone or in combination, on the histology of liver and kidney of Labeo rohita. *Pak J Zool*. 2013;45:913-920.
- 19. Kiron V. Fish immune system and its nutritional modulation for preventive health care. *Animal Feed Science and Technology*. 2012;173(1-2):111-133. doi:10.1016/j.anifeedsci.2011.12.015
- 20. Shi Y, Hu Y, Wang Z, et al. The protective effect of taurine on oxidized fish-oil-induced liver oxidative stress and intestinal barrier-function impairment in juvenile Ictalurus punctatus. *Antioxidants*. 2021;10(11):1690. doi:10.3390/antiox10111690. PMID:34829560

- 21. Wu X, Xu B, Chen H, et al. Crosstalk of oxidative stress, inflammation, apoptosis, and autophagy under reactive oxygen stress involved in difenoconazole-induced kidney damage in carp. *Fish Shellfish Immunol.* 2023;132:108508. doi:10.1016/j.fsi.2022.108508
- 22. Del Rio D, Stewart AJ, Pellegrini N. A review of recent studies on malondialdehyde as toxic molecule and biological marker of oxidative stress. *Nutr Metab Carbiovasc Dis.* 2005;15(4):316-328. doi:10.1016/j.numecd.2005.05.003
- 23. Hoseinifar SH, Yousefi S, Van Doan H, et al. Oxidative stress and antioxidant defense in fish: the implications of probiotic, prebiotic, and synbiotics. *Rev Fish Sci Aquac*. 2020;29(2):198-217. doi:10.1080/23308249.2020.1795616
- 24. Yu Z, Dai ZY, Qin GX, Li MY, Wu LF. Alleviative effects of dietary microbial floc on copper-induced inflammation, oxidative stress, intestinal apoptosis and barrier dysfunction in Rhynchocypris lagowski Dybowski. *Fish Shellfish Immunol*. 2020;106:120-132. doi:10.1016/j.fsi.2020.07.070

- 25. Wang L, Wang C, Huang C, et al. Dietary berberine against intestinal oxidative stress, inflammation response, and microbiota disturbance caused by chronic copper exposure in freshwater grouper (Acrossocheilus fasciatus). *Fish Shellfish Immunol*. 2023;139:108910. doi:10.1016/j.fsi.2023.108910
- 26. Gaetke LM, Chow-Johnson HS, Chow CK. Copper: toxicological relevance and mechanisms. *Arch Toxicol*. 2014;88(11):1929-1938. doi:10.1007/s00204-014-1355-y. PMID:25199685
- 27. Lushchak VI. Contaminant-induced oxidative stress in fish: a mechanistic approach. *Fish Physiol Biochem*. 2016;42(2):711-747. doi:10.1007/s10695-015-0171-5
- 28. Sappal R, MacDougald M, Fast M, et al. Alterations in mitochondrial electron transport system activity in response to warm acclimation, hypoxia-reoxygenation and copper in rainbow trout, Oncorhynchus mykiss. *Aquat Toxicol*. 2015;165:51-63. doi:10.1016/j.aquatox.2015.05.014