

Biochemical assessment of some heavy metals on enzyme activities of *Oreochromis niloticus* fish species in the White Nile River at south of Khartoum City, Sudan

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Abstract

Background and objective: *Oreochromis niloticus* is a fish species locally known as *Bulti*, and greatly demanded in local markets. The present study was to assess the levels of heavy metals and their effects on enzyme activities in type of fish species.

Materials and Methods: Ethanol extract was taken for the determination of cholesterol, total lipid contents in fish muscles. While saline extract was used for the estimation of glucose, total protein, albumin, globulin and enzyme like GOT, GPT, LDH, CPK and amylase. The concentrations of heavy metals were carried out by the flame atomic absorption spectrophotometer (UNICAM 929).

Results: There were no significant differences between enzyme activities in both liver and muscle extracts except for ALP found to be lower in tested samples comparing to the control. The cadmium (Cd) level in the liver was found to be higher than that in the muscle, whereas the zinc (Zn) level in the muscle was found to be higher than that in the liver.

Conclusions: the higher levels of heavy metals found in the contaminated fish samples compared to control could poses risks to human health during time.

Keywords: heavy metals, hematology, biochemical, *Bulti*, pollution

Introduction

There are important precedents on the serious consequences of heavy metals contaminated fish consumption has on human health, being worth mentioning: Minamata, Japan tragedy, where over 900 people lost their lives and two million suffered health problems as a consequence of having eaten Hg contaminated fish, Minamata disease (McCurry, 2006) [15] and the event at Jinzu River basin, at Toyama prefecture, where one of the most serious Cd prolonged intake intoxication cases takes place following the mining extractions that contaminated those waters (Inaba *et al.* 2005) [11]. However, fish are exposed to heavy metals in polluted waters. Heavy metals come into the water from the anthropogenic activities, and sources that continually released into the aquatic environment create serious health risks due to their toxicity, long persistence, bioaccumulation and bio-magnifications in the food chain (Bat *et al.*, 2013) [4]. It is vital to determine heavy metal levels in seafood, because heavy metal ions can accumulate more easily in fish as compared with other foodstuffs (Bat *et al.*, 2014) [3]. Aquatic systems especially lakes are more sensitive to heavy metal pollutants and are contaminated by chemical substances, industrial, domestic and agriculture wastes in the form of particles, metal ions, organic and inorganic compounds (Tarra-Wahlberg *et al.*, 2001) [28]. River Nile and its main tributaries White Nile and Blue Nile Rivers are the most important rivers as a source of water supply and catfish area that supplies most of Khartoum City residents. The aquatic life of the White Nile River is very rich in fish species diversity. The area under the study is located south of

Khartoum City nearby Jabal Awlia Dam. This area was selected for the study purpose because the river over there receives sewage effluent from Soba Treatment Plant and from the Military Factory. A recent study conducted by Nogod *et al.* (2020, 2021a, 2021b) [16, 17, 18] reported that the Suba Treatment Plant is negatively impacts the water quality of the White Nile River and poses hazard to the aquatic life and the public health at the surrounding environment. Therefore, the study aimed to evaluate the effect of the accumulation of heavy metals on the biological activity of *Oreochromis niloticus* fish species.

Materials and methods

Collection of fish samples

A total of 120 *Oreochromis niloticus* fish species, that commonly known as *Bulti*, (Figure 1) were collected, of which, a 60 tested fish were collected from the targeted polluted area; the first 30 fish of them were collected from the discharge zone of the White Nile River site (1), whereas the second 30 fish were collected from the main ponds of sewage site (2).



Fig 1: *Oreochromis niloticus* fish species, popular known as *Bulti*.



Fig 2: Fish sampling points, Jebel Awlia Fisheries Sites 3 (control) and Site 1 at White Nile River (test sample). Modified from www.Googleearth.com, (2016).

The rest (120 fish) were collected from the upstream non-polluted area of Jebel Awlia Dam (site 3), as control (Figure 2). The collected *Oreochromis niloticus* fish species were well washed with clean water and kept in polyethylene packages containing ice, then transported to the Central Veterinary Research Laboratory of the Omdurman University pending identification and analysis

Processing of liver and muscle tissues

Liver and muscle were cut with razor, washed with distilled water and blotted with blotting paper. A weighted portion (about 3 grams) of both liver and muscle was homogenized in 3 ml ice-cold saline (0.89% NaCl) solution for saline extract and 3ml ethanol for ethanol extract in a motor driven Teflon glass homogenizer. The homogenate centrifuged rpm at 4000 rpm ($3.500\times g$) for 45 minutes at 5°C in a refrigerated centrifuge to get a clear saline supernatant and for 15 minutes at 5°C at the same speed for ethanol supernatant. Digestion methods described by Van Loon (1980) and Du Preez and Steyn (1992) were used for the estimation of heavy metals, e.g., Cr^{3+} , Cu^{2+} , Zn^{2+} , Cd^{2+} , Fe and Pb^{2+} (Ali, 2004).

Biochemical analysis of fish liver and muscle

Saline and ethanol extracts were analyzed for the following biochemical parameters, including some enzymes to evaluate the possible effect of water pollution on them. Ethanol extract was taken for the determination of cholesterol, total lipid contents in fish muscles. While saline extract was used for the estimation of glucose, total protein, albumin, globulin and enzyme like GOT, GPT, LDH, CPK and amylase. Detailed procedures for estimation of all above parameters were the same as given under the biochemical analysis of liver and blood (Ali, 2004).

Determination of heavy metals in fish tissues

Frozen tissue samples of liver and muscle were thawed, rinsed in distilled water and blotted in blotting paper. A known weight of liver and muscle of the fish was shifted to 250 ml volumetric flasks for digestion. Samples digestion done according to methods described by Van Loon (1980) and Du Preez and Steryn (1992). By this method digestion was completed in about 20 minutes instead of 3 to 4 hours as

stated by Van Loon (1980). Sample after digestion were cooled and diluted to 10 ml with distilled water by proper rinsing of digestion flasks. Atomic absorption spectrophotometer was used to determine the concentration of Cr^{3+} , Cu^{2+} , Zn^{2+} , Cd^{2+} , Fe^{2+} and Pb^{2+} in the tissues sample of the fish. A range of analytical standards for each metal was prepared from Merck Stock solution (Van Loon, 1980 and Du Preez and Steyn, 1992).

Statistical analysis

Student t-test analysis of variance, mean differences, correlation and chart were done by using SPSS (Statistical Package for Social Sciences) program version.

Result and discussion

Biochemical evaluation of liver and muscle of *Oreochromis niloticus* fish species with respect to GOT, GPT, LDH, ALP, CPK and amylase

With regard to liver extract, table 1 showed that the level of GOT in the sample caught from contaminated river water site 1 (84.5) was higher but not significantly different comparing to that caught from the control site 3 (81.9). No significant difference was observed regarding GPT and LDH levels in both of the samples under the study. Whereas ALP level in site 1 (93.7) was found to be lower and significantly different comparing to the control (96.3). Amylase in site 1 sample (63.6) was found to be higher but not significantly different comparing to the control (62.4). For muscle extract, table 1 showed that all of the parameters of GOT, GPT, LDH and CPK of the tested liver extract were found to be not significantly different comparing to the control. However, GOT level was found to be lower in the control (71.8) comparing to contaminated site 1 which was both found to be (73.9). As well, GPT level in control (23.7) was lower than that of contaminated site 1 which was both found to be (26.4). CPK level was found to be higher in the control (36.4) comparing to contaminated site 1 which was found to be (34.1). Generally, GOT, GPT and LDH levels were found to be in the liver extract higher than that of the muscle extract. The hematology and biochemical reference values for aquaculture fish are varied according to a specific situation (i.e., malnutrition, stress, infection/diseases) and for a

specific species (Kerr, 2008) [12]. As previously stated, the activity of serum enzyme in this study found to be varied according to the organ from which the sample was extracted. This observation confirmed the result that was observed by a previous study conducted by Krajnovic-Ozretic and Ozretic (1987) [13] which stated that the LDH activity was found to be varied and it was primarily related to the organ from which the blood sampled. As well, similar result was observed by an earlier study conducted by Panteghini *et al.* (1984) [20] which reported that the determination of the basic enzymes provides the opportunity to identify their origin and also permits estimation of the severity of injury at the tissue/cellular level. The findings of this study were in accordance with the results obtained by the study conducted by Dorcas and Solomon (2014) [7], which reported both significant and insignificant differences between the levels of ALT, ALP and AST of fishes that studied and the levels of the standards. Ali (2004) reported that the concentration levels of blood serum enzyme of GOT, GPT and CPK was found to be much higher in blood tissues of *Tur-putitora* fish species from polluted waters River Kabul, Pakistan, when compared with that in control dams waters. Sagar *et al.* (2015) [26] reported that in liver cell injury, a considerable increase in the serum level of both GOT and GPT enzymes was observed, whereas serum CPK activity is raised in all varieties of muscular dystrophy. While raised LDH values are found in renal disease, liver disease, disseminated malignancy and certain hematological disorders. The increased serum levels of ALP are found in both skeletal and hepatic disorders, and amylase enzyme raised in acute abdominal conditions, e.g. perforated peptic ulcer, cholecystitis, common bile duct and intestinal obstruction, but the higher levels are found in acute pancreatitis. Moreover, Sagar *et al.* (2015) [26] stated reference range values of 5 - 50 U/L for both GOT and GPT, and for LDH to be 230-400U/L, ALP to be 70-230U/L, CPK to be 24-105 U/L, and for amylase is to be 60-160U/L. Previous studies conducted by Lemarie *et al.* (1991) [14]; De La Torre *et al.* (2000) [6]; Panigrahi *et al.*, 2010 [19] and Tahmasebi-Kohyani *et al.* (2012) [27] stated that the alterations of blood enzymes' activities were associated with liver damage, induced by pathological or stress situations or with feeding nutrients with hepatic protective effects. Perez-Jimenez *et al.* (2013) and Hari Krishnan *et al.* (2011) [10] reported that the increase of serum LHD activity may indicate hypoxia conditions, limiting water temperature or toxins, whereas the increase of serum CPK activity which is particularly active in heart and skeletal muscle, may indicate damage to these tissues. As well, Almeida *et al.* (2002) [1] and Rehulka and Minari'k, (2007) [4] declared that pollutants and stress conditions were reported to increase CPK activity, probably due to muscle injury.

Biochemical evaluation of liver and muscle of *Oreochromis niloticus* fish species with respect to total protein, albumin, globulin, glucose, cholesterol and total lipids

Table 2 revealed that all of the parameters of total protein, globulin and total lipids in the tested liver samples were found to be not significantly different comparing to the control. While albumin, glucose and cholesterol levels in tested samples were found to be significantly different comparing to the control. In muscle extracts, the total cholesterol level in contaminated site 1 (52.4) was found to

be higher and significantly different comparing to the control (47.1). For the rest parameters in muscle extract, no significant differences were found to be between the tested samples and the control. However, albumin and cholesterol levels in the liver were found to be higher than that in muscles. Globulin level in liver was found to be much lower than that in muscles. Sandnes and Waagb (1988) considered the range for total protein of normal healthy fish was to be 41.6–56.6 g l^{-1} , for albumin: 18.3–24.3 g l^{-1} , for cholesterol: 9.3 - 12.8 μmol , for triglycerides: 2.53–4.98 μmol , and for creatinine: 26–46 μmol . It was reported that plasma protein levels are often associated with fish nutritional and physiological status (Rehulka *et al.* 2005; Maita 2007). Basal plasma protein values averaged was stated by Polakof *et al.* (2012) to be 4.9 g dl-1. Whereas basal blood glucose levels are known to differ considerably among fish species. Basal plasma cholesterol levels reported by this study are lower comparing to the previously reported values by Peres *et al.* (2014). It also confirmed by Peres *et al.*, 2014 that, in seabass, liver has a pivotal role in the maintenance of glucose homeostasis.

Biochemical evaluation of liver and muscle of *Oreochromis niloticus* fish species with respect to Cd, Pb, Cu, Cr, Fe and Zn

Table 3 summarizes that the heavy metals levels of Cd, Pb, Cu, Cr and Zn of the tested liver extracts for treatment site 1 sample were found to be not significantly different comparing to the control sample, while Fe level in tested sample was found to be the only metal that significantly different comparing to the control.

In muscle extract, Cd, Fe and Zn levels exhibited no significant differences between contaminated sample site 1 and the control. Whereas the levels of Pb, Cu and Cr in contaminated site 1 were found to be significantly different comparing to the control. The level of Fe in the liver was found to be higher than that found in the muscle. Whereas the level of Zn in the tested site 1 sample in both liver and muscle were found to be higher than that found in the control. However, no remarkable increase of heavy metals in the liver has been observed comparing to their levels in the muscle. It was stated that the liver is the most organs accumulates pollutants of various kinds, as reported earlier by Galindo *et al.* (1986) and recently by Bat *et al.* (2015) [2]. On contrast, the Cu level in the muscle was found to be higher in control site 3 than that in the liver, this result disagree with that reported by Dugo *et al.* (2006). Pb levels in the liver were found to be much higher than that found for other metals. However, other studies stated that gills uptake certain metals such as Pb (Coban *et al.*, 2009). The accumulation of metals in the liver found to be followed the order: Fe, Pb, Cu, Cd, Cr and Zn, while in the muscle followed the order of Fe, Cu, Cd, Pb, Cr and Zn. Topçuoğlu *et al.* (2002) found that the accumulation of metals in muscle in wild Seabass taken from Perşembe, eastern Black sea showed the order of: Cd, Cr, Cu, Fe, Pb, Zn. Catsiki *et al.* (1999). Various studies suggest that metal uptake by fish performed via food that constitutes the major pathway for accumulation in liver and then muscle. However, when contamination occurs, concentration of metals in certain tissues increases in proportion with that of seawater (Coban *et al.*, 2009). The results of this study showed that the Fe concentrations were the highest in both liver and muscle and Zn concentrations were found to be the lowest. This result may be comparable to that reported by Bat

et al. (2015) [2]. However, it was stated that fish contents of heavy metals varies depending on the zone, environmental conditions, the contamination level of the fishing site, and the characteristics of the fish (Roderiguez et al., 2015). The maximum Cd concentrations (0.071) that detected in the liver tissues were remarkably higher than the maximum level (0.05 mg/g wet weights) set by the Commission Regulation (2006) whereas the Cd concentrations in the muscle tissues (0.0572)

were slightly above that recommended maximum level. Similarly, the maximum Pb concentrations (0.439) found in the liver tissues were slightly higher than the maximum level (0.30 mg/ g wet weights) set by the Commission Regulation. This findings in accordance of that reported by Bat et al. (2015) [2]. Zuluaga et al. (2015) [31] reported that Cd and Pb pose the highest risks for human health due to causing toxicity and possible carcinogenic effect.

Table 1: Biochemical parameters of blood serum of *Oreochromis niloticus* fish species (commonly known as *Bulti*) caught from Jebel Awlia Dam site₃ (control) and White Nile River site₁ (contaminated).

Locations of sample	Parameters													
	SGOT IU/L	SGPT IU/L	CPK IU/L	ALP IU/L	Total protein (g/dl)	Albu-min (g/dl)	Globu-lin (g/dl)	glucos-e mg/dl	Choles-terol mg/dl	Chlori-de mg/dl	Na mg/l	K mg/l	Ca mg/l	Fe mg/l
Site ₃	67.7	32.5	30.5	92.3	7.0	3.3	3.7	72.8	50.4	44.2	133.8	4.3	8.2	62.0
Site ₁	82.0	33.5	35.7	90.5	6.8	3.4	3.4	68.5	58.8	41.9	137.0	4.7	8.4	61.2
d.f.(N-2)	28	28	28	28	28	28	28	28	28	28	28	28	28	28
S.E±	3.77	1.94	2.31	1.42	0.08	0.08	0.13	1.66	2.26	1.04	1.91	0.07	0.12	0.12
P-value	0.001	0.598	0.033	0.223	0.016	0.097	0.015	0.001	0.032	0.001	0.000	0.125	0.480	0.028
Sig-level	**	Ns	*	Ns	*	Ns	*	**	*	Ns	**	Ns	Ns	*

Ns = No significant difference, * = Significant difference at 5%, ** = Significant difference at 1%. GOT = glutamate oxaloacetate transaminase, GPT = glutamate pyruvate transaminase, CPK = creatin phosphor kinase, ALP = alkaline phosphatase, Na = sodium, K = potassium, Ca = calcium, Fe = iron.

Table 2: Biochemical parameters of blood serum of *Oreochromis niloticus* fish species (commonly known as *Bulti*) caught from Jebel Awlia Dam site₃ (control) and White Nile River site 1 and treatment pond site 2 (contaminated).

Locations of sample	Parameters													
	SGOT IU/L	SGPT IU/L	CPK IU/L	ALP IU/L	Total protein (g/dl)	Albu-min (g/dl)	Globu-lin (g/dl)	glucos-e mg/dl	Choles-terol mg/dl	Chlori-de mg/dl	Na mg/l	K mg/l	Ca mg/l	Fe mg/l
Site ₃	67.7 ^c	23.0 ^c	32.2 ^a	87.4 ^b	6.6 ^b	3.14 ^b	3.18 ^b	53.2 ^c	43.8 ^b	42.8 ^a	131.2 ^b	3.98 ^b	7.4 ^c	55.6 ^a
Site ₂	87.8 ^a	35.6 ^a	35.1 ^a	92.3 ^a	6.8 ^a	3.41 ^a	3.31 ^a	69.3 ^a	60.3 ^a	42.4 ^a	137.0 ^a	4.36 ^a	8.3 ^a	60.3 ^a
Site ₁	79.4 ^b	30.6 ^b	33.1 ^a	88.6 ^b	6.5 ^b	3.23 ^b	3.07 ^b	58.0 ^b	45.1 ^b	41.9 ^a	129.3 ^b	4.20 ^b	8.0 ^b	59.6 ^a
P-value	0.000	0.000	0.185	0.001	0.000	0.000	0.008	0.000	0.000	0.520	0.000	0.001	0.000	0.539
Sig-level	**	**	Ns	**	**	**	**	**	**	Ns	**	**	**	Ns
S.E±	4.19	2.16	2.29	1.77	0.15	0.16	0.14	2.39	2.75	1.04	1.87	0.12	0.13	4.48

Ns = No significant difference, * = Significant difference at 5%, ** = Significant difference at 1%. GOT = glutamate oxaloacetate transaminase, GPT = glutamate pyruvate transaminase, CPK = creatin phosphor kinase, ALP = alkaline phosphatase, Na = sodium, K = potassium, Ca = calcium, Fe = iron.

Table 3: Biochemical parameters of blood serum of *Oreochromis niloticus* fish species (commonly known as *Bulti*) caught from Jebel Awlia Dam site₃ (control) and White Nile River site 1 and treatment pond site 2 (contaminated).

Locations of sample	Parameters													
	SGOT IU/L	SGPT IU/L	CPK IU/L	ALP IU/L	Total protein (g/dl)	Albu-min (g/dl)	Globu-lin (g/dl)	glucos-e mg/dl	Choles-terol mg/dl	Chlori-de mg/dl	Na mg/l	K mg/l	Ca mg/l	Fe mg/l
Site ₃	77.2	27.6	28.3	89.4	6.2	3.02	3.22	56.6	41.2	39.5	130.5	4.18	8.14	59.0
Site ₁	71.4	27.6	32.5	88.4	6.3	3.12	3.17	55.9	47.6	41.2	127.3	4.27	8.21	56.2
P-value	0.048	0.992	0.0055	0.233	0.494	0.167	0.652	0.758	0.050	0.51	0.074	0.335	0.386	0.052
Sig-level	*	Ns	**	Ns	Ns	Ns	Ns	Ns	**	Ns	Ns	Ns	Ns	Ns
S.E±	2.83	1.90	1.35	0.88	0.10	0.07	0.10	2.32	1.60	0.82	9.78	0.10	0.08	1.40

Ns = No significant difference, * = Significant difference at 5%, ** = Significant difference at 1%. GOT = glutamate oxaloacetate transaminase, GPT = glutamate pyruvate transaminase, CPK = creatin phosphor kinase, ALP = alkaline phosphatase, Na = sodium, K = potassium, Ca = calcium, Fe = iron.

Table 4: Biochemical parameters of blood serum of *Oreochromis niloticus* fish species (commonly known as *Bulti*) caught from Jebel Awlia Dam site₃ (control) and White Nile River site 1 and treatment pond site 2 (contaminated).

Locations of sample	Parameters													
	SGOT IU/L	SGPT IU/L	CPK IU/L	ALP IU/L	Total protein (g/dl)	Albu-min (g/dl)	Globu-lin (g/dl)	glucos-e mg/dl	Choles-terol mg/dl	Chlori-de mg/dl	Na mg/l	K mg/l	Ca mg/l	Fe mg/l
Site ₃	70.9	16.6	27.4	87.9	6.7	3.22	3.44	51.2	48.4	39.2	130.2	4.10	8.08	62.4
Site ₁	76.9	30.2	30.2	85.8	6.4	3.19	3.19	57.6	54.4	40.3	131.6	4.26	8.25	57.5
P-value	0.004	0.000	0.065	0.042	0.029	0.741	0.110	0.003	0.006	0.370	0.815	0.100	0.177	0.187
Sig-level	**	**	Ns	*	*	Ns	Ns	**	**	Ns	Ns	Ns	Ns	Ns
S.E±	1.87	2.80	1.48	1.00	0.12	0.09	0.15	1.93	2.03	1.21	6.10	0.09	0.12	5.47

Ns = No significant difference, * = Significant difference at 5%, ** = Significant difference at 1%. GOT = glutamate oxaloacetate transaminase, GPT = glutamate pyruvate transaminase, CPK = creatin phosphor kinase, ALP = alkaline phosphatase, Na = sodium, K = potassium, Ca = calcium, Fe = iron.

Conclusion

In conclusion, the higher levels of Fe metal found in the contaminated samples compared to control accumulate them in the liver more than in muscles of the tested fish species, and have greatly affected enzymes' activities. Among these enzymes, GOT, GPT, LDH and CPK of the tested liver extract were found to be higher comparing to that level in the muscles. Albumin and cholesterol levels in the liver were found to be higher than that in muscles. The concentration levels of Fe were found to be the highest in both liver and muscle, and Zn concentration levels were found to be the lowest.

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