Original Research Article

2 IMPACT OF POLUTION ON HAEMATOLOGICAL HAEMATOLOGY AND 3 HISTOLOGICAYL ASSESSMENT OF JUVENILES OF Chrysichthys nigrodigitatus IN 4 OGBESE RIVER, ONDO STATE, NIGERIA

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6 ABSTRACT

The silver catfish *Chrysichthys nigrodigitatus* catfish is a fish of economic importance in sub-7 sahara Africa. In Ogbese town, and its environs, it constitutes a means of income and food for 8 fisherfolks and community members. Hence, this study was undertaken to assess health status of 9 Chrysichthys nigrodigitatus using heamatology and histological assessment of the fish specie 10 due to the anthropogenic activities that takes place around the river body. A total 120 live fish 11 samples of *Chrysichthys nigrodigitatus* were collected by the assistance of fisherfolks using fish 12 cage at Ogbese River from May to August, 2018. Some water parameters measurements were 13 taken: temperature, pH, DO, Turbidity and Conductivity. Morphometric measurement: Weight 14 (g) and length (cm) of fish were taken. Heamatolotgy and histology of fish gills, liver and 15 intestine were determined. Mean water temperature (27.70±0.18°C), pH (7.36±0.22), DO 16 (6.98±0.15 mg/l), Turbidity (78.50±13.53 NTU) and Conductivity (148.35±27.98) of the river 17 determined respectively. Mean body weight of fish was 148.15 ± 36.53 g, and mean length was 18 19 25.64 ± 2.86 cm. The gills, liver and intestines of the fish specie were examined to assess the architecture of the organs. rResults of haematology studies of C. nigrodigitatus revealed high 20 values in the parameters measured. Red Blood Cell was higher than the White Blood Cell with 21 mean value of $(225.63\pm10.45 \ 10^3/\text{mm}^3)$ while Eosinophils recorded lowest parameters with 22 mean value of $(1.75 \pm 0.52 \%)$. Results of histology of gills, liver and intestines showed that the 23 gill filaments are were eroded with a deformation of the cartilage core and also hyperplasia of 24 the secondary lamellae. The intestines showed atrophy in a mucosal layer, hemorrhage and 25 dilation within blood vessels and within serosa of mucosa and for liver, picnotic nucleus are 26 were shattered, the hepatocytes are were ruptured and there is was increased kupffer cell count 27 as a result of exposure to pollutants. The results indicated that pollution level of the environment 28 have significant impact on health status of fish. 29

30 **KEYWORDS**: Chrysichthys nigrodigitatus, Ogbese River, Haematology, Histology.

32 INTRODUCTION

31

Fish is one of the most important animal protein sources that are widely consumed by all races and classes of people (Abolude and Abdullahi, 2005). It compares favorably with milk, meat, pork and poultry (James, 1984). Fish and fishery products are highly nutritious and are excellent sources of other dietary essentials like vitamins and minerals. Fish fat contains a high proportion of polyunsaturated fatty acids which may help to decrease the incidence of atherosclerosis and heart related diseases (Akande, 2011). Fish also provide an important complement to the predominantly carbohydrate based diet of many people in Nigeria (Akande, 2011).

The silver catfish *Chrysichthys nigrodigitatus* (Lacepede, 1803) is a highly valued food-fish
included among the dominant commercial catches exploited in Ogbese river, Ondo State,
Nigeria. It is restricted to the bottom of deep water, omnivorous; consume bivalves, detritus,

chironomids, crustaceans and vegetable matter (Bankole *et al.*, 2011). This fish can be raised in
both fresh and brackish water environments.

Fish health can be adversely affected by temperature changes, habitat deterioration and aquatic 45 pollution (Skouras et al., 2003). Hematological parameters are considered an important indicator 46 of fish health status, and provide valuable information to assess the fish welfare (Azevedo et al., 47 48 2006). Hematology is also used as an indicator of physiological and pathological changes in fish (Chekrabarty and Banerjee 1988, Martins et al., 2008). It can be affected by several factors 49 including gonad maturation (Ranzani-Paiva and Godinho, 1985), dissolved oxygen alterations 50 (Ranzani-Paiva et al., 2000), gender (Lusková, 1998), spawning and water temperature (Joshi 51 1982), lotic or lentic environment (Val et al., 1985), handling stress and transportation (Gbore et 52 al., 2006), fish inflammation (Martins et al., 2006), size, feeding and stocking density (Rey 53 Vázquez and Guerrero, 2007), microbial infection and parasitism (Martins et al., 2004, Azevedo 54 55 et al., 2006. Jamalzadeh et al., 2009). Ogbese region comprises Ogbese community and some neighboring agrarian settlements that 56 sustain it with agricultural produce. The location of Ogbese in the rain forest zone in South 57 Western Nigeria gives it a natural tendency of wood, timber and food production in the region. 58 The community serves as an economic life wire of Akure North Local Government Area of 59 Ondo State that produces food crops in large quantities. With Dispite these economic potentials, 60 the town still remains a remote rural settlement in the State. 61 Pollution of the rivers examined in this study is mainly through run-off activities from 62 agricultural practices and commercial activities. Many studies have shown that very large 63 64 quantities of heavy metals are found in run-off associated with the operation of motor vehicles, 65 atmospheric fallout and road surface materials (Harper, 1985). To the environmental scientists, the ultimate concern of trace metal contaminants in receiving water is their toxic impact on 66 aquatic organisms and including fish species (Sutherland and Tolosa, 2000; De Carlo et al., 67 2004). Assessing pollutants in different components of the ecosystem is an important task in 68 preventing risk to natural life and public health. Pollutants entering these receiving waters by 69 way of run-off conveyance systems, indiscriminate dumping of wastes e.t.c, may adversely 70 impact many of the desired uses. The Ogbese community has undergone great economic 71 development in recent years. In fact, it is notably one of the fastest growing, economically 72 important communities in Ondo State and handles a considerable number of micro- industries. 73 The very popular market (Ogbese market) and the timber business coupled with unequalled 74 agricultural practices have drawn people from several cultural backgrounds in the country to 75 76 make the settlement inter-tribal. This increase in anthropogenic activities surrounding the area has lead to an increase in environmental degradation. These multiple sources make it especially 77 78 difficult to identify and isolate the risks associated with this contaminated water. Unfortunately,

records of water quality parameters are non-existing and no known monitoring programmes
 onfish health due to the water quality have been initiated within the state.

81

82 MATERIALS AND METHODS

83 Study Area

84 The study site was Ayede, Ogbese River along Akure-Benin expressway in Ondo State. The area

lies between $E6^{0}SE8^{0}$ and longitude $N4^{0}N6^{0}E$. The river has its source from Ayede-Ekiti in Ekiti

state and flows through Ogbese in Ondo State to Edo State. The Ogbese community is about

87 10km east of Akure, the Ondo state capital.

88 Collection of Water Samples

89 Water samples were collected using water samplers at 10 cm depth at three points locations from

90 the river body, and parameters were determined using multi- parameter machine Model No: for

91 dsissolved oxygen, temperature, turbidity, conductivity, and pH.

92

93 Collection of Fish

120 live *Chrysichthys nigrodigitatus* fish samples were collected by the assistance of fisherfolks
using fish cage at Ogbese River from May to August, 2018. They were then transported alive in
buckets containing water to the Marine Biology Laboratory of the Department of Fisheries and

97 Aquaculture Technology, Federal University of Technology, Akure.

98

99 Length-weight Measurement

The weight in grams (g) of each specimen was taken using a digital weighing balance, which was wiped dry between samples. Standard length was measured in centimeters (cm) using a meter ruler.

103 Condition factor of the fish was assessed to know the state of <u>health</u> being of the fish.

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 $K = \frac{100 \text{ X W}}{L^3}$

106 K = Condition Factor

107 W = Body Weight of Fish in gram (g)

108 L = Standard Length of Fish in centimetreers (cm)

110 3.4 Haematological Analysis

Blood samples were taken from the caudal vein of each fish using a syringe and transferred to 5ml of Ethylene Diamine Tetraacetic Acid (EDTA) bottles. After blood collection in the laboratory, the samples were maintained on ice and sent to the laboratory of Animal Production and Health Technology, Federal University of Technology, Akure for hematological analysis.

115 The haematological parametres analysed were; Erythrocyte Sedimentation Rate Count (ESR),

Packed Cell Volume Count (PCV), Red Blood Cell Count (RBC), Haemoglobin Concentration
 (Hgb), White Blood Cell Count (WBC), Lymphocyte Count, Neutrophils Count, Monocytes

- 117 (figb), white Blood Cen Count (WBC), Eynphocyte Count, Neurophils Count, Monocytes 118 Count, Basophils Count, Eusonophils Count._Mean Corpuscular Volume (MCV), Mean
- 119 Corpuscular Haemoglobin (MCH) And Mean Corpuscular Haemoglobin Concentration (MCHC)
- 120 were calculated according to (Houston, 1990).
- 121 The Haemoglobin was calculated as: Hb (g/100ml) = Absorbance of test x Concentration of
- 122 standard Absorbance of standard Absorbance of standard of Total erythrocyte (RBC)
- 123 Red Blood Cell and White Blood Cell <u>counts</u> were calculated thus; = C x D x 4000
 124 Where:

Volume of whole blood

- 124 V 125
 - C = dilution factor (20)

D = number of cells counted Hematocrit/ PCV = Volume of packed red blood cell X 100

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- 129 130

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- White blood cell (WBC) = %WBC X total WBC + thrombocytes counts
- 132 The red cell indices MCHC, MCH and MCV were derived thus;

Comment [U1]: Include the model number or delete Model No:

133 134 135 136	Mean	Cell Hemoglobin Concer	ntration (MCHC) = <u>Hemog</u> l	obin (g/100ml) X 100 PCV(%)		
130 137 138	Mean	Corpuscular Haemoglobi		<u>globin (g/100ml)</u> X 100 C (x10,000rbc/mm ³)		
139 140 141 142	$\begin{array}{l} \text{Mean Cell Volume (MCV)} = & \underline{PCV \times 100} \\ \text{RBC} (x10,000_rbc/mm^3) \\ \text{RBC} (x10,000_rbc/mm^3) \end{array}$					
143 144 145 146 147	 3.5 Histological Analysis The fish specimen was dissected using a dissecting set. The gills, liver and intestines were then removed and rinsed in distilled water to remove blood stains. The organs were then placed in a 10ml sample bottle with 10% formalin for preservation and transported to the Anatomy and 					
148 149 150 151 152	 148 149 3.6. Statistical Analysis 150 Data collected were analyszed using one-way ANOVA. Further tests were done using Duncan 151 Multiple Range Test. And test of significance were done at P ≥ 0.05. 				((Comment [U2]: How were the histological parameters measured? Comment [U3]: Is it > or < ?
153	4.0.	Results and Discussion				
154	4.1.	Physico-Chemical Par	ameters of <u>water from</u> Riv	ver Ogbese		
155	The physicochemical properties of water obtained from River Ogbese are presented in					
156	Table	1.				
157		Table 1: Physico_chemical parameters of <u>water from</u> River Ogbese.				Comment [U4]: Include a Colum of WHO or EPA standards for comparison
		Parameters	Range	Mean±SD		
		DO (mg/l)	5.80 - 7.99	6.98 ± 0.15		
		Turbidity (NTU)	67.00-97.00	78.50 ± 13.53		

 27.70 ± 0.18

 148.35 ± 27.98

 7.36 ± 0.22

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Temperature (°C)

Ph

Conductivity (µohm's/cm)

159 Length, Weight, Condition Factor (K) and LWR of Chrysichthys nigrodigitatus

160 Length (cm), Weight (g), Length / Weight Relationship and Condition factor (K) of C. 161 *nigrodigitatus* obtained at River Ogbese are shown in (Table 2). The average body weight of 162 *Chrysichthys nigrodigitatus* used was 148.15 ± 36.53 g which ranged from 106g - 185g, while 163 the average body length was 25.64 ± 2.86 cm ranging between 23cm - 30cm. The condition 164 factor was 0.88. The "b" values of the fish were not equal to 3, hence growth in the individual

26.44 - 30.64

119.0-178.0

6.81-8.12

species was allometric (i.e. b values were less/greater than 3) showing that the rate of increase in 165 body length is not proportional to the rate of increase in body weight. 166

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Table 2: Morphometric Characteristic of Chrysichthys nigrodigitatus_obtained from River Ogbese

Length / Weight Relationship	Measurement
Length (cm)	25.64 ± 2.09
Weight (g)	148.15 ± 28.56
Condition Factor (K)	0.88
Intercept (a)	2.08
Slope (b)	2.29
Coefficient of determination (r ²)	0.64

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Haematological Parameters of Chrysichthys nigrodigitatus obtained from River Ogbese 171

Tables 3 and 4 showed haematology characteristics of the Chrysichthys nigrodigitatus. The 172 173

result showed high values in parameters measured. Red Blood Cell was higher than the White Blood Cell count with mean value of (225.63±10.45). Eosinophils recorded the lowest

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parameters count with mean value of (1.75 ± 0.52) . 175 176

Table 3: Haematological Profile of Chrysichthys nigrodigitatus from River Ogbese. 177

Parameters	MAY	JUNE	JULY	AUGUST
ESR	3.50±0.71 ^a	4.00±0.78 ^a	3.75 ± 0.42^{a}	4.00±0.00 ^a
PCV (%)	24.50±0.71 ^a	22.50±0.41 ^a	23.50±1.41 ^a	24.50±0.28 ^a
RBC (µL)	237.00±8.49 ^a	218.00 ± 4.24^{b}	219.50±9.19 ^b	228.00±11.31 ^c
WBC (µL)	123.00±7.07 ^a	113.50±2.12 ^b	115.50±13.44 ^b	113.50±10.61 ^b
Hb (gdL-1)	8.15±0.21 ^a	$7.80{\pm}0.42^{a}$	$8.00{\pm}0.28^{a}$	8.50±0.21 ^a
Lymphocytes	59.00±1.41 ^a	50.00 ± 0.00^{a}	55.00±1.41 ^a	59.50±2.12 ^a
Neutrophils	25.00 ± 0.00^{a}	34.00±2.83 ^a	22.50±2.12 ^{ab}	23.00±4.24 ab
Monocytes	12.50±1.41 ^a	12.00±2.83 ^a	13.50 ± 2.12^{a}	13.00±1.41 ^a
Basophils	2.00±0.71 ^a	2.50±0.91 ^a	2.00±0.41 ^a	2.50±0.71 ^a
Eosinophils	$1.50{\pm}0.71^{a}$	$1.00{\pm}0.71^{a}$	2.50±0.71 ^a	$2.00{\pm}0.00^{a}$
MCHC (gdL-1)	33.27 ± 0.09^{a}	33.19±0.21 ^a	33.19±0.29 ^a	33.27±0.16 ^a
MCH	3.44±0.03 ^a	3.58±0.06 ^a	3.56±0.02 ^a	3.50±0.10 ^a
MCV (pg)	$10.34{\pm}0.07^{a}$	10.78 ± 0.11^{a}	10.71 ± 0.13^{a}	10.75±0.23 ^a

Values on the same row with the same superscript alphabet are not significantly different. N = 30 178

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Comment [U5]: Compared to which values? May be values of fish from a non polluted source?

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- 182 183

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Table 4: Range and Mean Haematological Profile of Chrysichthys nigrodigitatus from River
Ogbese

Parameter	Range	Mean±SD	SR
ESR (mm)	3.00-4.00	3.81±0.35	4-10
PCV (%)	23.00-25.00	23.75±0.76	21-26
RBC $(10^{3}/mm^{3})$	213.0-243.0	225.63±0.45	200-250
WBC $(10^{3}/mm^{3})$	106.0-128.0	116.38±8.19	100-150
Hb (g/100ml)	7.60 - 8.30	8.11 ±0.27	5-10
Lymphocytes	58.00-61.00	55.88±1.19	64-80
Neutrophils (%)	20.00 - 26.00	26.13±2.33	25-30
Monocytes (%)	10.00-15.00	12.75±1.69	10-20
Basophils (%)	2.00-3.00	2.25±0.53	2-5
Eosinophils (%)	1.00-2.00	1.75±0.52	1-2
MCHC (gdL-1)	33.04 - 33.33	33.23±0.13	30-45
MCH (pg)	3.40 - 3.60	3.52±0.07	5-10
MCV (pg)	10.20 - 10. 90	10.65±0.22	10-15

185 Data are presented as Means \pm S.D. ESR =Erythrocyte Sedimentation Rate, PCV =Packed Cell

Volume, HB =Haemoglobin, RBC =Red Blood Cell, WBC =White Blood Cell, MCV =Mean
Corpuscular Volume, MCHC =Mean Cell Haemoglobin Concentration, MCH =Mean Cell

188 Haemoglobin. S.R = Standard Range

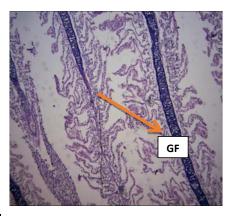
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190 Histology of Chrysichthys nigrodigitatus

Results of histology of gills, liver and intestines of *Chrysichthys nigrodigitatus* are given in the
plates 1 - 13 below. The gill filaments are were eroded with a deformation of the cartilage core
and also hyperplasia of the secondary lamellae. The intestines showed atrophy in a mucosal
layer, hemorrhage and dilation within blood vessels and within serosa of mucosa. Liver histology
revealed shattered picnotic nucleus, ruptured hepatocytes and increased kupffer cells.

196 Histology of the Gills

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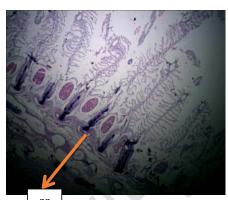
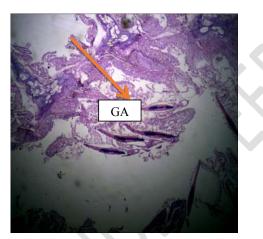


PLATE 2: The gill filaments are <u>showed</u> eroded cartilage<u>.</u> Magnification; x 100

199 **cartilage. Magnificatio** 200

cc TLATE3: There is a deformation of the core. Magnification x 100



- PLATE 4: The gill arch and gill filaments areshowing visible signs of lesions
- 204 Magnification; x400
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206 4.4.2 Histology of the Intestines

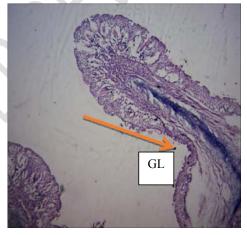
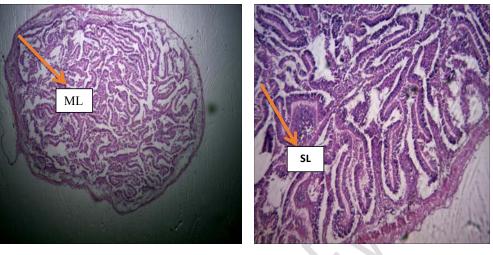


PLATE 5: There is hyperplasia of the eroded secondary lamellae

Magnification; x 400

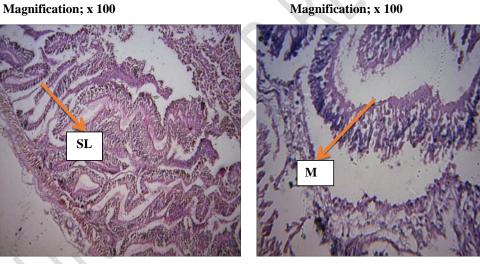


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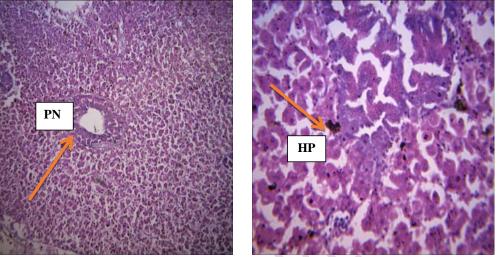
PLATE 6: shows atrophy in a mucosal layer PLATE 7: Intestine shows sign of haemorrhage 208

Magnification; x 100



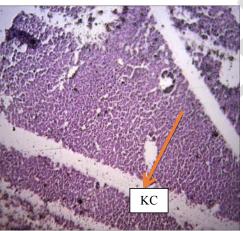
211 PLATE 8: shows hemorrhage and dilation PLATE 9: shows severe degeneration and 212 within blood vessels and within serosa of mucosa. necrosis of mucosal membrane of intestine 213 Magnification; x400 Magnification; x400

Histology of the Livers 214



215 216

PLATE 10: The picnotic nucleus are shattered PLATE 11: The hepatocytes are ruptured 217 Magnification; x 100 Magnification; x 100



218 219 PLATE 12; There is increased kupffer cells

220 Magnification; x400

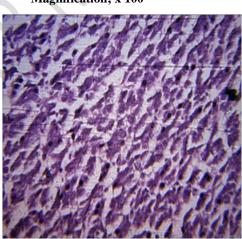


PLATE 13; Visible lesions seen

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x400

Magnification;

- GF= Gill Filaments, CC= Cartilage Core, GA= Gill Arch, GL= Gill Lamellae, ML= Mucosa 221
- Layer, SL= Serosa Layer, PN= Picnotic Nucleus, KC= Kkupffer Cell. 222

DISCUSSION 223

- Results of physico_chemical parameters of water obtained in this study were within the tolerable 224
- range of fish as recommended by WHO (2001 and 2006) except for DO. The result was similar 225
- to the reports of Ansa (2004) on the benthic macrofauna of the Andoni flats in the Niger Delta 226

Area of Nigeria, Chindah *et al.*,(1998) on effect of municipal waste discharge on the physico_
chemical and phytoplankton in a brackish wetland in Bonny Estuary and Ladipo *et al.*, (2011) on
seasonal variations in physico-chemical properties of water in some selected locations of Lagos
Lagoon who opined that waters with little change in physico-chemical parameters are generally
more conducive to aquatic life. Most organisms including *C. nigrodigitatus* do not tolerate wide
variations in physico-chemical parameters and if such conditions persist_death may occur. High
oxygen demand experienced in theis study is in line with Adebayo *et al.*, (2007) observation.

Ujjania et al., (2012) opined that condition factor greater or equal to one is good, indicating a 234 good level of feeding, and proper environmental condition. Mean K-values gotten from this 235 study (0.88) were less than one (1) in samples, hence revealing that the species fell slightly from 236 beening unhealthy. This support the report of Gesto et al., (2017) who worked on the Length-237 Weight Relationship and Condition factor of C. gariepinus and O. niloticus of Wudil River, 238 Kano, Nigeria, and obtained condition factor less than one (1). Also feeding intensity, 239 availability of food, fish-size, age, sex, season, stage of maturation, fullness of the gut, degree of 240 muscular development and amount of reserved fat (Gupta and Banerjee, 2015) also have 241 influence on also K factor of fish 242

The observation of absolute Isometric growth (b = 3) in nature is occasional (Bagenal 1978; Bassey and Ricardo, 2003), and deviation from isometric growth is often observed in most aquatic organisms which changes shape as they grow (Thomas *et al.*, 2003). The differences in the length-weight relationship also agrees with the report of Olurin and Aderibigbe (2006) who stated that the differences may be due to sex and developmental stages of fish.

Mean heamotocrit value of C. nigrodigitatus was 23.75±0.76% which did not differ considerably 248 from those found by Badawi and Said 1971 and Etim et al., 1999. The Red Blood Cell counts 249 hasd a mean value of 225.63 x 10^6 mm³ ± 10.45 x 10^6 mm³. The Packed cell volume (PCV) hasd 250 a mean value of 23.75±0.76%. Heamoglobin concentration hasd a mean value of 8.11±0.27g/dl. 251 The mean haemoglobin value is low which may be due to the exposure of fish to pollutants 252 resulting in inhibitory effect of those substances on the enzyme system responsible for the 253 synthesis of haemoglobin according to Pamila et al., 1991. The low hb value in the water body 254 may also be associated with less active fishes. Similar results were reported by Engel and Davis, 255 (1964) and Rambhaskar and Rao, (1987). Eisler suggested that there was a correlation between 256 haemoglobin concentration and the activity of the fish. The more active fishes tend to have 257 258 higher haemoglobin values than the more sedentary ones (Pradan et al., 2012). The high erythrocyte number was associated with fast movement, predaceous nature and high activity 259 streamlined body (Satheeshkumaretal., 2011). A fall in hematological parameters 260 with count, Hb% and PCV%, in the fishes, due to water pollution, has been reported along with acute 261 anemia (Singh, 1995). According to Singh et al., 2002), the discharge of waste may cause serious 262 263 problems as they impart odour and can be toxic to aquatic animals. The organic wastes present in 264 Ogbese river seem to cause stress in the fish and as such seem to be responsible for the changes in the hematological parameters. The PCV or haematocrit is an important tool for determining 265 266 the amount of plasma and corpuscles in the blood (measurement of packed erythrocytes) and is used to determine the oxygen carrying capacity of blood (Larsson et al., 1985). Hematocrit or 267 PCV in the present study is low compared to the works of (Joshi et al., 2002) and (Banerjee and 268 Banerjee, 1988) have suggested that pollutant exposure decreases the TEC count, Hb content and 269 270 PCV value due to impaired intestinal absorption of iron.

Comment [U8]: Not clear

Comment [U9]: Year of publication?

There were variations in WBC quantity and leukocyte cell proportions (neutrophil, monocyte) in 271 the fish specimens. The implication of this result is that the fish has been able to defend itself 272 from invading pathogens both by cell and antibody-mediated responses (Kumar et al., 1999). 273 Similar results were obtained by Sahan and Cengizler, (1894) on carp caught from different 274 regions of Seyhan River. Leukocytosis is directly proportional to severity of stress condition in 275 276 maturing fish and is a result of direct stimulation of immunological defense due to the presence of pollutants in water bodies. This is in conformity with the report of Saravanan and 277 Harikrishnan, (1999) in freshwater fish, Sarotherodon mossambicus, when exposed to sublethal 278 concentration of copper and endosulfan and by Nanda, (1997) in respect of *Heteropneustes* 279 fossilis during nickel intoxication. This may be attributed to alteration in blood parameters and 280 direct effects of various pollutants. The lymphocytes are reported to be responsible for immune 281 response (Cazenave et al., 2005), while neutrophils are reported to show the greatest sensitivity 282 to change in the environment. Their characterization and identification is therefore, of 283 significance for assessing the changes in the physiological state of fishes 284

285 Marked variations like hyperplasia, vacuolation, deformation of cartilage core, bubbling of gill 286 filament, epithelial lifting, lamellar fusion; secondary lamellar damage, shorter secondary lamellae and erosion of secondary lamellae were noticed in the gill tissues of C. nigrodigitatus 287 collected from river Ogbese. Similar results were obtained by several works: Fernandes and 288 289 Mazon, (2003), Simonato et al., (2008), Rajeshkumar et al., (2015), as they revealed alterations like aneurysm, mucous deposition, hypertrophy, fusion of secondary lamellae, ruptured epithelial 290 291 layer, lifting of primary lamellae, lamellar swelling and necrosis. Through the gills, as the main 292 site of xenobiotic transfer, the toxins are distributed through their bodies accumulating in tissues and organs and may have deleterious effects Vasanthi, et al., (2015). 293

The extent of liver damage observed in the present investigation indicates that chronic exposure 294 always causes impairment to the architecture of the tissue. Since liver is involved in 295 detoxification of pollutants (Lagadic et al., 2000), it is susceptible to a greater degree of 296 disruption in its structural organization due to toxic stress. Some distinct changes like rupture of 297 hepatocytes, melanomacrophages, increased Kupffer cell, increased pycnotic nucleus, 298 vacuolation, ruptured nucleus, Blood congestion, cytoplasmatic vacuolation and nucleus 299 300 disorganization were observed in the liver of fish. Macrophage aggregates have been suggested as potentially sensitive histological biomarkers and or immunological biomarker of contaminant 301 exposure (Schmitt et al., 2000). Histological changes observed in various studies in liver taken 302 from the fishes exposed to pollutants include increased vacuoles in the cytoplasm, changes in 303 nuclear shapes, focal area of necrosis (death of cells in a localized area), ischemia (blockage of 304 capillary circulation), hepatocellular shrinkage, and regression of hepatocytic microvilli at the 305 bile canaliculi, fatty degeneration and loss of glycogen.(Marchand et al., 2012) reported that 306 histopathological changes of fish liver from polluted freshwater system shows structural 307 alterations in hepatic plates or cords, multiple focal areas of cellular alterations leading to a loss 308 309 of uniform hepatocyte structure, steatosis, cytoplasmic and nuclear alterations (hypertrophic and 310 pyknotic nuclei) of hepatocyte, increase in the size of melanomacrophage centers (MMCs), and focal areas of necrosis. The results from this study also agrees with the result of microscopic 311 examination of liver specimens from Lagos and Ologe Lagoon which were consistent with the 312 findings of Olarinmoye et al. (2009) in which liver of C. nigrodigitatus from Lagos lagoon 313

showed several alterations including vacuolar hepatocellular degeneration and hepatic necrosis.

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315 Histology of the Intestine in the study revealed visible sign of lesions. Although, uptake of metals occurs mainly through gills, it may also occur via intestinal epithelium. Histopathological 316 alterations in the intestine of C. nigrodigitatus included severe degenerative and necrotic changes 317 in the intestinal mucosa and sub mucosa, atrophy in the muscularis and sub mucosa and 318 319 aggregations of inflammatory cells in the mucosa and sub mucosa with edema between them. 320 These findings are in agreement with those of Hanna et al., (2005), Bashir (2010), Yousafzai et 321 al., (2010) and Soufy et al., (2007), who opined that pollutants and contaminants affects gills by epithelial lifting, hyperplasia of epithelial cells and blood congestion within filaments and in 322 liver tissue produced hemolysis between hepatocytes, cytoplasmic degeneration and necrosis. 323 Whereas an aggregation of inflammatory cells, edema in an intestinal mucosal layer and 324 325 hemorrhage between blood vessels were the main alterations observed in the intestine. The changes seemed to be more pronounced in the liver and gills rather than the intestine. 326

327 Conclusion

Human activities including industrialization and agricultural practices contributed immensely in no small measure to the degradation and pollution of aquatic environment which adversely has effects on the water bodies that is a necessity for life. Since water pollution has direct consequences on human well beings, an effective teaching strategy in the formal education sector is essential for aquatic health

Regulation and monitoring is an effective way of pollution management. There is need to enact legislation to regulate various types of pollution as well as to mitigate the adverse effects of pollution.

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