

BIOACCUMULATION OF AUTO-MECHANIC WASTES IN THE TISSUES OF *Pellonula afzeliusi* (Johnels, 1954) COLLECTED FROM AGBA STREAM ILORIN, KWARA STATE, NIGERIA

Adeyemi-Ale, O. A.^{1*}, Oladipo, S. O.² and Abdulkareem, S. I.¹

¹Department of Zoology, Faculty of Life Sciences, University of Ilorin, Ilorin, Kwara State, Nigeria.

²Department of Biosciences and Biotechnology, Kwara State University, Malete, Kwara State, Nigeria.

* E-mail of corresponding author: adewoyin.aa@unilorin.edu.ng

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ABSTRACT

Hydrocarbons and heavy metals (HM) present in automobile wastes get into water bodies close to automobile mechanic villages. They deteriorate the physico-chemical qualities of such water bodies and exert toxic effects on fish and other aquatic fauna. They also pose risks to humans that consume fish from streams near such auto-mechanic villages. There is paucity of information on the HM contents of Agba stream, Ilorin and no information on the bioaccumulation of HM in the tissues of *Pellonula afzeliusi* is currently available. The purpose of this study was to assess the physico-chemical qualities of water and bioaccumulation of HM in the tissues of *Pellonula afzeliusi* fish collected from Agba stream, Ilorin, Kwara State, Nigeria. Water and samples of *Pellonula afzeliusi* were collected from the upstream (station A), midstream (station B) of Agba stream located close to an auto-mechanic village and Oyun dam (control site, station C) located within the University of Ilorin. Standard methods were used to analyse oil and grease (OG), total petroleum hydrocarbon (TPH) and HM in water samples. The concentrations of TPH and HM were analysed in the skin, bone, muscle, gills, gastrointestinal tract, liver and kidney of the fish. The order of water HM concentrations indicated Zn > Cd > Ba > Fe > Pb at station A; Cd > Zn > Fe > Ba > Pb at station B, and Zn > Ba > Pb > Fe > Cd at station C. TPH and HM bioaccumulated in the tissues of fish while lead was not detected in all tissues from station C. Bioaccumulation factor varied with the highest values at station C. The result of this study indicated heavy pollution burden in station B with lower pollution burden in station C (the control site). These call for attention of government in ensuring ecological restoration and prevention of further discharge of automobile wastes into the stream.

Keywords: Automobile wastes, Heavy metals, Hydrocarbons, Bioaccumulation, *Pellonula afzeliusi*

INTRODUCTION

In recent times, there is an increase in the discharge of automobile wastes through motor servicing centers popularly known as auto-mechanic workshops. Petrol, diesel, solvents, grease, and lubricants are accidentally or deliberately discharged from the auto-mechanic workshops on to the land (Adewoyin *et al.*, 2013). The release of these fossil fuel products on the land, leads to the accumulation of heavy metals in the soil and leaching to nearby wells and streams, thus bringing about biological threats and adverse environmental impacts to the nearby ecosystem (Adelekan and Abegunde, 2011; Utang *et al.*, 2013).

The large-scale spilling of petroleum products has been recognized as a serious environmental problem because used engine oil picks up a number of additional compounds like heavy metals from engine wear (Horsfall, 2011; Popoola and Ayodele, 2016). These additives and contaminants make used oil disposal to be more

environmentally damaging than crude oil (Abioye *et al.*, 2012). Due to the bioaccumulation and toxic tendency of heavy metals (El-Nagger *et al.*, 2009), they are considered to be the most hazardous environmental pollutants (Al-Attar, 2005).

The increase in anthropogenic activities such as the auto-mechanic servicing activities brings a growing concern over the ecological effects of accumulation of heavy metals in the environment bringing about significant effects on the trophic chain, plants, animals and man (Campbell *et al.*, 2005). Hence, deterioration of water quality and a steep degradation of freshwater biodiversity habitats are experienced (Nevoh *et al.*, 2015).

Streams and reservoirs are invaluable ecological resources that serve many human needs domestically, industrially and ecosystem wise (Oladipo *et al.*, 2018). Oyun and Agba reservoirs in Ilorin were constructed to serve the purposes of irrigation, domestic water supply and fish farming.

Fishes are aquatic inhabitants that are unable to get rid of the lethal effects of various pollutants in natural waters (Olaiya *et al.*, 2004). They have been recognized as suitable bio-indicator for the assessment of heavy metal contamination in aquatic systems due to the ease of accumulation through uptake of metals in water and diet (Papagiannis *et al.*, 2004; Izegaegbe and Oloye, 2017). These metals can bioaccumulate in the tissues of fish. The beneficial effects of fish consumption can be compromised by the presence of toxic heavy metals in them (Bawuro *et al.*, 2018).

Pellonula afzeliusi are small-toothed, herring-like fishes belonging to the family Clupidae, and distributed in West African rivers and man-made lakes (Ikusemiju *et al.*, 1983). They are pelagic (Froese and Pauly, 2019), and their diet include crustaceans, Ephemeropteran nymphs, chironomids, fishes, plant detritus and sand grains (Ikusemiju *et al.*, 1983; Ezenwaji and Offiah, 2004). *Pellonula afzeliusi* is the dominant fish in Agba stream and residents around the stream

collect and consume the fish. There is paucity of information on the heavy metal contents of Agba stream, Ilorin and no information on the bioaccumulation of heavy metals in the tissues of *Pellonula afzeliusi*. Therefore, there is need to assess the level of heavy metals in Agba stream, their concentrations and bioaccumulation levels in *Pellonula afzeliusi* inhabiting the stream.

MATERIALS AND METHODS

Description of the Study Sites

Agba Stream (8°47'0" N, 4°56'0" E) located in Ilorin South Local Government Area, Kwara State, Nigeria (Figure 1) flows out from Agba dam. It flows through the side of an auto-mechanic village. The upstream with no influence of auto-mechanic activities was designated as station A, while the point where auto-mobile wastes are directly dumped or sipped into the stream, located about 500 m from station A was designated as station B. Oyun dam (8°28'10" N, 4°39'50" E) located at the University of Ilorin, Kwara State was taken as the control site (station C) because it is apparently free from the influx of pollutants.

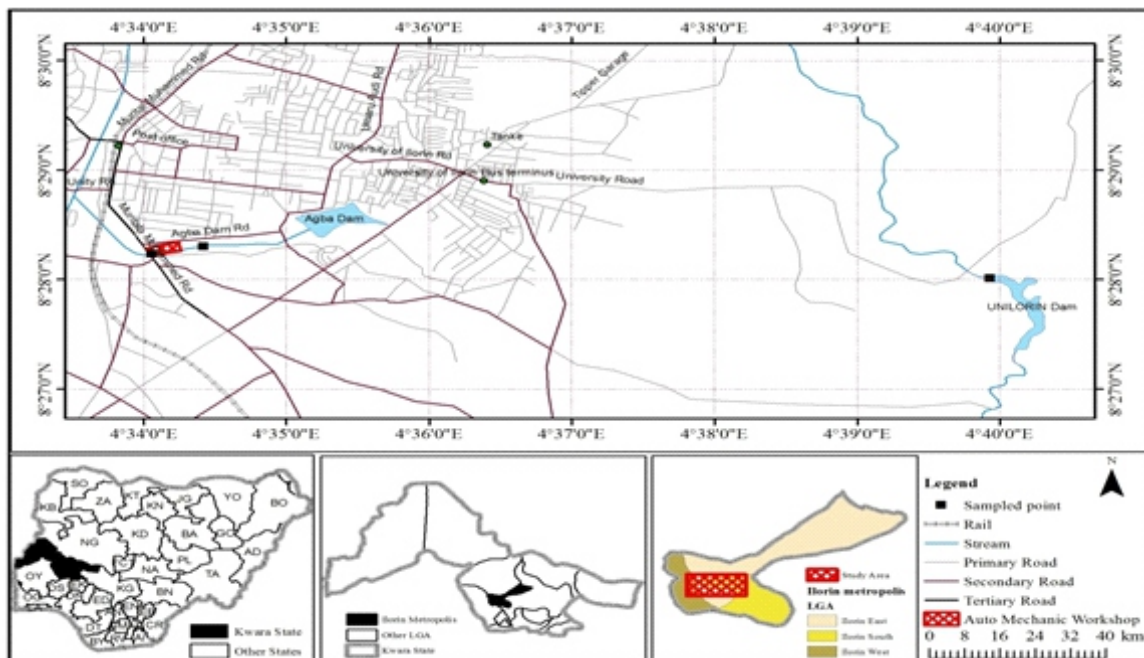


Figure 1: Map of the Study Sites Showing Agba stream and Oyun (UNILORIN) Dam

Water Sampling and Analysis

Water samples were collected in the early hours between 7:00 am to 9:00 am in the month of March 2018. The samples were analysed for physico-chemical qualities such as oil and grease (OG), total petroleum hydrocarbon (TPH) and heavy metals (HM) such as iron (Fe), zinc (Zn), cadmium (Cd), barium (Ba) and lead (Pb).

Water samples for the analysis of OG and TPH were collected in one-litre amber glass bottles with Teflon-lined screw caps. The samples were immediately filtered and preserved with concentrated HCl and held at 4 °C until extracted within 14 days of collection. The water samples for HM analysis were collected in 500 ml reagent bottles and were immediately preserved with 0.75 ml of concentrated nitric acid (HNO₃) prior to digestion and further analysis (APHA, 2012). The hexane extractable gravimetric method, (EPA 1664B extraction technique, 2010) was used for the determination of OG and TPH while samples for HM digestion and analysis were according to the standard procedures of American Public Health Association (APHA, 2012).

Preparation, Extraction and Analyses of OG and TPH

In the laboratory, the samples were adjusted to a pH < 2 with concentrated HCl. The samples were brought out to room temperature. In a 500 ml separating funnel, 350 ml of sample was added. Five (5) boiling chips were added to 125 ml distillation flask and the flask was weighed. For extraction, 20 ml n-hexane was added to the separating funnel; the stopper was put and inverted. This was vigorously shaken for 2 minutes. It was left to stand for 10 minutes to separate the lower water layer and the upper solvent. The lower water layer was drained into a 500 ml volumetric flask for about 4 min. Ten grams (10 g) of anhydrous sodium sulphate (Na₂SO₄) was added to a folded filter paper in the filtering funnel. The Na₂SO₄ was rinsed with a small amount of n-hexane and the n-hexane was discarded. Through the funnel that contains filter paper and 10 g Na₂SO₄, the solvent layer was drip-drained into the pre-weighed boiling flask and the water layer was returned to the separating funnel. The extraction procedure was repeated two more times.

The separating funnel was rinsed with three different 5 ml aliquots of fresh n-hexane to remove oil film that stayed on the funnel walls. Each aliquot was drained into the distillation flask through the funnel that contains Na₂SO₄. After this, distillation process was carried out and completed within 28 min. A vacuum connector was attached and vacuum applied for 2 min to remove the remaining n-hexane solvent vapors from the distillation flask. The distillation flask was put in a desiccator until the flask temperature decreases to room temperature and subsequently the flask was weighed.

OG was calculated using the formula:

$$\frac{A - B}{\text{Sample volume}} \times 100 = \text{mg/L O\&G}$$

A – Weight (mg) of residue

B – Weight (mg) of flask with boiling chips

TPH Analysis

The residue was dissolved with 25 ml of fresh n-hexane and heated slightly. The dissolved residue was poured into 100 ml volumetric flask and made up to volume with n-hexane and properly mixed. Silica gel (3 g) was added and the solution was stirred on a magnetic stirrer for 5 min. A funnel was put on a clean dry distillation flask with 5 boiling chips. A filter paper was pre-moistened with fresh n-hexane, placed in the funnel and the solution was filtered. The volumetric flask with the remaining silica gel was rinsed with 5 ml aliquots of fresh n-hexane and the aliquots was poured into the distillation flask, then the distillation process was carried out. The remaining product at the bottom of the flask was weighed and TPH was calculated using the formula:

$$\frac{A - B}{\text{Sample volume}} = \text{mg/L TPH}$$

A – Weight (mg) of residue

B – Weight (mg) of flask with boiling chips

Heavy Metals Analysis

Ten millilitres (10 ml) concentrated nitric acid and 5 ml concentrated perchloric acid were added to 20 ml of water sample in a 100 ml Erlenmeyer flask. This was allowed to mix thoroughly and the mixture was evaporated to 2 to 5 ml on a hot plate until a clear solution was obtained. It was made up

to volume with double distilled water. Each heavy metal was analysed with graphite furnace Atomic Absorption Spectrophotometer (AAS Model Agilent Aa55).

Fish Sampling and Analysis

Fish samples ($n = 30$; average length = 12.03 ± 0.32 cm; average weight = 16.3 ± 0.5 g) were collected in the early hours between 7:00 am to 9:00 am in the month of March, 2018. Fish samples were collected using hook and line, and cast net (3/8 inches mesh size) with the help of local fishermen. All the specimens of *Pellonula afzeliusi* were kept in a water container and transported to the laboratory of the Department of Zoology, University of Ilorin, Nigeria. Specimens were identified following the identification keys provided by Olaosebikan and Raji (1998) and Idodo-Umeh (2003). Five fish were randomly picked for analysis. The length of the fish selected ranged from 10 – 15 cm while their weights measured between 13.90 – 20.10 g.

In the laboratory, the fish was dissected and the skins, bones, muscles, gills, gastrointestinal tract, liver and kidney removed. They were then transferred into different bottles containing 0.25 ml sucrose solution (1:5 wt/vol) and then homogenized with mortar and pestle. After homogenization, they were transferred back into the bottles containing sucrose solution.

Fish TPH Analysis

TPH concentrations were determined following the method of Ikpe *et al.* (2016). TPH extracting solvent (1:1 v/v of acetone and dichloromethane) was prepared. Twenty-five millilitres (25 ml) each of acetone and dichloromethane were measured into 100 ml volumetric flask and mixed properly. Then, 0.5 g of each homogenized tissue sample was weighed into a 50 ml beaker and 30 ml of the extracting solvent was added. The fish TPH content was extracted by shaking method according to Schwab *et al.* (1999). The beaker with the content was placed on a magnetic stirrer and shaken for about 10 min at 70 °C. The extract was decanted into a clean round bottom flask, 10 ml of fresh solvent was added, and the process repeated. The extracts were combined, and 1 g of anhydrous sodium sulphate was added to remove water. The extract was concentrated to 3 ml with rotary evaporator maintained at 20 °C (Webster *et al.*,

1997).

Analysis of Heavy Metals

The concentrations of heavy metals in the tissues were digested and analysed following the method described by Du Preez and Steyn (1992). Briefly, 0.1 g of each fish tissue sample was weighed into 100 ml Erlemenyer flask into which 10 ml concentrated nitric acid and 5 ml perchloric acid were added. Digestion was done on a hotplate (200 to 250 °C) for about 4 hours during which there was total digestion and clear samples obtained. Each digested sample was filtered through filter paper. The filter system was rinsed with double distilled water to remove traces of the dissolved metal. Each sample was then made up to volume with the distilled water. An analytical blank was also prepared in a similar manner. Each heavy metal was analysed using a graphite furnace Atomic Absorption Spectrophotometer (AAS Model Agilent AA55).

Calculation of bioaccumulation factor (BAF):

$$AF = \frac{[X]_{organism}}{[X]_{water}} \quad (\text{USEPA, 2003})$$

Where

$[X]_{organism}$ is the concentration of the metal X in the tissue of the organism

$[X]_{water}$ is the concentration of the metal X in the water

BAF values less than 250 were scored 1 (low concern), values from 250 - 1000 were assigned 2 (medium concern) and 3 (high concern) was assigned to values greater than 1000 (waste minimization prioritization tool (WMPT) (USEPA, 2003).

Statistical Analysis

All statistical analysis was carried out using IBM SPSS version 21. The means and standard error of the means of the physico-chemical parameters of water were calculated. The bioaccumulation of TPH and HM in the fish tissues were also computed. ANOVA (Analysis of variance) and Duncan's multiple range tests were used to test for statistical ($P < 0.05$) differences of physico-chemical parameters of water, TPH and HM bioaccumulation in fish samples of the stations.

RESULTS AND DISCUSSION

The assessment of TPH, OG and HM in this study showed high contributions of auto-mechanic wastes to the environment. Concentrations of heavy metals in the water samples from Agba stream at station A and station B, and the reference site (station C) reflected the order of metal concentration as follows: Zn > Cd > Ba > Fe > Pb at station A, Cd > Zn > Fe > Ba > Pb at station B, and Zn > Ba > Pb > Fe > Cd at station C. All the heavy metal contents of the station C were significantly ($P < 0.05$) different from stations A and B, except that of zinc of station A. The TPH and OG contents of water samples from stations A and B exceeded the standard permissible limits of 10 mg/L and 0.1 mg/L by Environmental Guidelines and

Standards for the Petroleum Industry in Nigeria (EGASPIN, 2002) and National Environmental Standards and Regulatory Enforcement Agency (NESREA, 2011) respectively. All the HM concentrations of the water samples from station B exceeded the NESREA (2011) allowable limits (Table 1). Out of all the metals measured in the three stations, Zn concentrations were mostly elevated and Cd concentration obtained from station B. The increase in mean concentration of heavy metals in the samples could be attributed to metal concentration from the auto-mechanic wastes together with reduction in water volume during the period of sampling which coincided with the peak period of dry season. Similar observation in Bradford Beck catchment, West Yorkshire was reported by Goodwin *et al.* (2003).

Table 1: Total Petroleum Hydrocarbon, Oil and Grease and Heavy Metals Concentrations (mg/L) of Water Samples from Agba stream and Oyun dam, Ilorin, Nigeria

Parameters	Station A	Station B	Station C	NESREA Limit
TPH	138.6000±0.0577 ^b	153.4000±0.0577 ^c	9.0820±0.0006 ^a	-
Oil and G	350.0000±0.5774 ^b	1000.0000±0.0000 ^c	0.0070±0.5774 ^a	0.1
Iron (Fe)	0.1300±0.0000 ^b	0.8067±0.0088 ^c	0.0227±0.0001 ^a	0.5
Zinc (Zn)	0.6700±0.0056 ^{ab}	1.0000±0.0000 ^b	0.1200±0.0058 ^a	0.2
Cadmium (Cd)	0.0900±0.0056 ^b	1.2000±0.0577 ^c	0.0130±0.0000 ^a	0.01
Barium (Ba)	0.3000±0.0056 ^b	0.5200±0.0058 ^c	0.1000±0.0000 ^a	-
Lead (Pb)	0.0800±0.0006 ^b	0.3900±0.0058 ^c	0.0260±0.0006 ^a	0.1

Means with different superscript along same row are significantly different at $P < 0.05$
NESREA – National Environmental Standards and Regulatory Enforcement Agency

Fish meat is an important protein source to man (Buikema *et al.*, 1982). The accumulation of heavy metals in fish tissues is dependent upon physicochemical and biological variables such as pH, temperature, hardness, salinity, exposure duration, feeding habits of species and habitat complexity (Zeitoun and Mehana, 2014; Putshaka *et al.*, 2015). The result of the concentrations of TPH and HM in all the fish organs collected from the sampling stations are presented in table 2. Through bioaccumulation and bio-magnification, large volumes of HM get concentrated in the body of fish since they live primarily in water (Velez and Montoro, 1998) and they are at higher trophic level. The concentrations of all the metals investigated in all the organs of fish at station B were significantly ($P < 0.05$) higher compared to stations A and C. TPH was not accumulated in the bone of fish samples from station C while it had

the highest and lowest concentrations in the gills and kidney of samples from stations B and C, respectively. The TPH was higher in the muscle than in the skin and bone. This is similar to the reports of Enuneku *et al.* (2015) and Al-Ali *et al.* (2016) who observed higher values of TPH in fish muscles. Samples from station B showed that the liver had the highest concentrations of Fe and Zn while the lowest were recorded in the muscle and kidney, respectively; Cd had highest concentration in the kidney while the liver recorded the lowest concentration. However, Cd was not bioaccumulated in the gills, GIT and liver of fish sampled from stations A and C. Fish samples from station C did not have any trace of Ba in the gill, GIT and liver while Pb was not detected in all the organs except a very trace amount in the liver. The fish were able to regulate the bioaccumulation of these metals because of the chronically elevated

levels of Zn (USEPA, 2003). This is due to the fact that there are no known disorders associated with the excessive accumulation of zinc (Vallee and Falchuk, 1993) and its general stable association with macromolecules and coordination flexibility makes it highly adaptable to meet with the needs of proteins and enzymes that carry out various biological functions (Vallee and Auld, 1989, 1990a and 1990b).

The accumulation of the metals in the liver is likely linked to the role of liver in metabolism (Zhao *et al.*, 2012). The physiological role of the liver in blood cells and hemoglobin synthesis gives iron the tendency of accumulating in hepatic tissues (Korkmaz *et al.*, 2012). The pattern of bioaccumulation in the liver has also been observed in a number of other reports on fish species (Rashed, 2001; Dural *et al.*, 2006; Pyle *et al.*, 2005; Storelli *et al.*, 2006; Ploetz *et al.*, 2007; Agah *et al.*, 2009). The resultant effects of these accumulated TPH and HM in humans consuming this fish from Agba stream over time could result into vomiting and diarrhea (Dudkar and Miller, 1999). Similarly, pollutants in fish can severely reduce some vital nutrients in the body of consumers and this may be accountable for declining immune defense, growth delay, reduced physiological abilities, incapacities related with malnutrition and greater occurrence of upper

gastro-intestinal cancer (Iyengar and Nair, 2000; Arora *et al.*, 2008).

The result of the bioaccumulation factors (BAF) of total petroleum hydrocarbon and heavy metals in all the fish organs collected from the sampling stations are presented in table 3. The BAF computed in the organs of fish samples collected from station A ranged from 0 for Cd in the gills, GIT and liver to 687.69 for iron observed in the kidney. For station B, the BAF ranged from 0.01 for TPH in the bone and kidney to 219.4 for Zn in the liver. The BAF for station C ranged from 0 for TPH in the bone; Cd and Ba in the gills, GIT and liver; and Pb in all the organs to 2832.6 for iron in the liver. The highest BAF values were gotten for tissues from station C (Table 3) which is the control station that had the lowest concentrations of TPH and HM (Table 2). The values obtained at the control site confirmed that BAF values were highest at the lowest exposure concentrations and decreased as concentrations increased (McGeer *et al.*, 2003). Similar reports of inverse relationship of metal concentrations and BAFs were given by different authors (Yehia and Sebae, 2012; Asante *et al.*, 2014; Muir *et al.*, 2017). Body burdens tend to increase (i.e., bioaccumulation occurs) as metal exposure concentration increases, but at a relatively slower rate (USEPA, 2003).

Table 2: Concentration of Heavy Metals and Total Petroleum Hydrocarbon in the Tissues of *Pellonula afzeliusi* from the Agba Stream and Oyun Dam

Parameters	Tissues	Sampling Locations		
		Station A	Station B	Station C
TPH (mg/g)	Skin	0.0063±0.0001 ^a	0.0079±0.0001 ^c	0.0042±0.0002 ^b
	Bone	0.0011±0.0017 ^b	0.0015±0.000 ^b	0.0000±0.0000 ^a
	Muscle	0.0107±0.0020 ^b	0.0123±0.00051 ^c	0.0092±0.0004 ^a
	Gills	0.0122±0.0007 ^b	0.0145±0.0003 ^c	0.0099±0.0005 ^a
	GIT	0.0113±0.0009 ^{ab}	0.0128±0.0004 ^b	0.0107±0.0002 ^a
	Liver	0.0069±0.0006 ^b	0.0071±0.0006 ^b	0.0049±0.0001 ^a
	Kidney	0.0008±0.0001 ^b	0.0008±0.0001 ^b	0.0006±0.0000 ^a
Fe (mg/g)	Skin	0.0798±0.0034 ^b	0.0906±0.0004 ^c	0.0337±0.0004 ^a
	Bone	0.0325±0.0009 ^{ab}	0.0649±0.0001 ^b	0.0213±0.0001 ^a
	Muscle	0.0273±0.0029 ^b	0.0408±0.0003 ^c	0.0139±0.0001 ^a
	Gills	0.0435±0.0007 ^b	0.0926±0.0004 ^c	0.0280±0.0002 ^a
	GIT	0.0352±0.0016 ^b	0.0918±0.0012 ^c	0.0231±0.0012 ^a
	Liver	0.0841±0.0021 ^a	0.1532±0.0006 ^c	0.0643±0.0003 ^b
	Kidney	0.0894±0.0015 ^b	0.0966±0.0002 ^c	0.0425±0.0005 ^a
Zn (mg/g)	Skin	0.0835±0.0018 ^b	0.1443±0.0004 ^c	0.0718±0.0003 ^a
	Bone	0.0883±0.0021 ^b	0.1495±0.0021 ^c	0.0668±0.0002 ^a
	Muscle	0.0882±0.0026 ^a	0.1460±0.0028 ^b	0.0875±0.0004 ^a
	Gills	0.0977±0.0027 ^a	0.1418±0.0004 ^b	0.0967±0.0003 ^a
	GIT	0.0937±0.0046 ^a	0.1416±0.0008 ^b	0.0847±0.0025 ^a
	Liver	0.0773±0.0095 ^a	0.2194±0.0020 ^b	0.0709±0.0008 ^a
	Kidney	0.0687±0.0013 ^a	0.1224±0.0004 ^b	0.0685±0.0005 ^a
Cd (mg/g)	Skin	0.0005±0.0005 ^a	0.0088±0.0006 ^b	0.0001±0.0009 ^a
	Bone	0.0005±0.0003 ^a	0.0119±0.0002 ^b	0.0001±0.0002 ^a
	Muscle	0.0003±0.0007 ^a	0.0098±0.0006 ^b	0.0001±0.0003 ^a
	Gills	0.0000±0.0000 ^a	0.0007±0.0001 ^a	0.0000±0.0000 ^a
	GIT	0.0000±0.0000 ^a	0.0006±0.0000 ^a	0.0000±0.0000 ^a
	Liver	0.0000±0.0000 ^a	0.0005±0.0000 ^a	0.0000±0.0000 ^a
	Kidney	0.0002±0.0000 ^a	0.0131±0.0007 ^b	0.0001±0.0000 ^a
Ba (mg/g)	Skin	0.0006±0.0002 ^b	0.0100±0.0003 ^c	0.0001±0.0002 ^a
	Bone	0.0009±0.0002 ^a	0.0280±0.0007 ^b	0.0001±0.0007 ^a
	Muscle	0.0004±0.0009 ^a	0.0090±0.0006 ^b	0.0001±0.0007 ^a
	Gills	0.0002±0.0000 ^a	0.0091±0.0002 ^b	0.0000±0.0000 ^a
	GIT	0.0001±0.0000 ^a	0.0011±0.0000 ^b	0.0000±0.0000 ^a
	Liver	0.0001±0.0000 ^a	0.0034±0.0003 ^b	0.0000±0.0000 ^a
	Kidney	0.0005±0.0000 ^a	0.0109±0.0009 ^b	0.0001±0.0000 ^a
Pb (mg/g)	Skin	0.0002±0.0006 ^a	0.0009±0.0003 ^b	0.0000±0.0000 ^a
	Bone	0.0007±0.0002 ^a	0.0087±0.0002 ^b	0.0000±0.0000 ^a
	Muscle	0.0001±0.0003 ^a	0.0001±0.0002 ^a	0.0000±0.0000 ^a
	Gills	0.0003±0.0000 ^b	0.0013±0.0000 ^c	0.0000±0.0000 ^a
	GIT	0.0002±0.0002 ^b	0.0009±0.0000 ^c	0.0000±0.0000 ^a
	Liver	0.0001±0.0000 ^a	0.0059±0.0007 ^b	0.0001±0.0000 ^a
	Kidney	0.0000±0.0000 ^a	0.0008±0.0000 ^b	0.0000±0.0000 ^a

Means with different superscript along same row are significantly different at P<0.05.

Table 3: Bioaccumulation Factors of Total Petroleum Hydrocarbon and Heavy Metals in the Tissues of *Pellonula afzeliusi* Collected from the Agba Stream and Oyun Dam

Tissues	Stations	Parameters					
		TPH	Fe	Zn	Cd	Ba	Pb
Skin	A	0.05	613.85	124.63	5.56	2	2.5
	B	0.05	112.31	144.3	7.33	10.92	2.31
	C	0.46	1484.58	598.33	7.69	1	0
Bone	A	0.01	250	131.79	5.56	3	8.75
	B	0.01	80.45	149.5	9.92	53.85	22.31
	C	0	938.33	556.67	7.69	1	0
Muscle	A	0.08	210	131.64	3.33	1.33	1.25
	B	0.08	50.58	146	8.17	17.31	0.26
	C	1.01	612.34	729.17	7.69	1	0
Gills	A	0.09	334.62	145.82	0	0.67	3.75
	B	0.09	114.79	141.8	0.58	17.5	3.33
	C	1.09	1233.48	805.83	0	0	0
GIT	A	0.08	270.77	139.85	0	0.33	2.5
	B	0.08	113.8	141.6	0.5	2.12	2.31
	C	1.18	1017.62	705.83	0	0	0
Liver	A	0.05	646.92	115.37	0	0.33	1.25
	B	0.05	189.91	219.4	0.42	6.54	15.13
	C	0.54	2832.6	590.83	0	0	3.85
Kidney	A	0.01	687.69	102.54	2.22	1.67	0
	B	0.01	119.75	122.4	10.92	20.96	2.05
	C	0.07	1872.25	570.83	7.69	1	0

According to waste minimization prioritization tool (WMPT) scoring (USEPA, 2003), the BAF values obtained from this study indicated that total petroleum hydrocarbon, cadmium, barium and lead were of low concern, zinc ranged from low to medium concern, and iron ranged from low, medium to high concern.

CONCLUSION

The environmental quality assessment of the stream indicated heavy pollution burden in station B as a result of auto-mechanic workshop along the stream with lower pollution burden in station C (the control site). This calls for attention in ensuring ecological restoration and prevention of further discharge of auto-mobile wastes into the stream through management. This will ensure preservation and sustainability of the ecosystem, and fish in this stream will pose less health risk to humans consuming them.

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