

Human Health Risk Assessment of Some Trace Metals Through Fish Consumption From Qua Iboe River, Oruk Anam, Nigeria

Abstract

Food consumption is an important route of human exposure to organic chemicals. To assess the health risks associated with these chemicals due to fish consumption, the levels of trace metals (Cd, Cr, Ni, Cu, V, Pb and Sn) in water and fish from Qua Iboe River were assessed using atomic absorption spectrometry (AAS). Physicochemical parameters in water were determined using standard methods. The levels of physicochemical parameters in water were within permissible limits set by World Health Organisation (WHO). The levels of trace metals in water samples varied from one sampling location to another and ranged from 0.01 ± 0.01 to 0.10 ± 0.00 mg/L in wet season and from 0.01 ± 0.01 to 0.08 ± 0.01 mg/L in dry season. In fish the levels ranged from 0.01 ± 0.01 to 0.61 ± 0.01 mg/kg in wet season and from 0.02 ± 0.01 to 0.51 ± 0.01 mg/kg in dry season. Results of physicochemical parameters showed variations across sampling locations and seasons. Human health risk of trace metals contamination due to consumption of the fish revealed that the hazard quotient (HQ) values were all less than unity ($HQ < 1$). It therefore means that there is no health implication for consumers of commercial fish from the river. It is recommended that the studied river be monitored regularly through researches to check escalation in levels of these contaminants while the riverine communities should be educated by government on danger associated with indiscriminate disposal of waste into aquatic environment.

Keywords: Trace metals, Physicochemical, Fish, Analysis, Statistical, Oruk Anam, Nigeria.

Introduction

Environmental sustainability is a major consideration in environmental management. It is majorly achieved through routine monitoring to assess the levels of contaminants with a view to prevent escalation (Akpan *et al.*, 2024). The process of monitoring the environment to assess the possible presence of some contaminants enhances environmental protection. Monitoring of environmental media including water and fish gives information on the pollution status of the aquatic ecosystem and it is an important step in exposure control. The analysis of aquatic biota such as fish gives information on the bioavailability of contaminants as well as bio-magnification along the food chain.

Assessment of human health risk is essential in the maintenance of sustainable environment and safety. Lifetime exposure to contaminants such as trace metals and other chemicals in the environment through ingestion can pose risks to human health (Akpan *et al.*, 2024). Higher levels of chemical agents in water, food, air and soil are potentially harmful to humans as a result of exposure (Ido *et al.*, 2023). Trace metals and other chemicals find their ways into the aquatic environment from a variety of sources including the naturally occurring biogeochemical cycles such as carbon, nitrogen, water, oxygen (Burger *et al.*, 2002). Researches reveal that low levels of some trace metals are essential for aquatic animals and may also become toxic when they accumulate in different organs, damage tissues and interfere with normal growth and proliferation (Udosen, 2019).

Biological samples such as fish, among others have higher concentrations of trace metals than water making them more suitable for routine monitoring and more relevant in the context of exposure of humans and wildlife (Muir *et al.*, 2003). Fish accumulates some trace

metals in its fatty tissues due to its inability to metabolise them in the food they eat and from the intake of particulates in water and sediment onto which these chemicals have been adsorbed (Gbeddy *et al.*, 2015). The region of accumulation of these metals within fish varies with the route of uptake. The gills are directly in contact with water. Therefore the concentration of metals in gills reflects their concentration in water where the fish live (Rejomon *et al.*, 2010). The average concentration of these trace metals increases dramatically across a food chain. The bioaccumulation of trace metals in fish and other animals is the reason most of the human daily intake of such chemicals come from our food supply rather than from water (Butu and Iguisi, 2013).

The potential use as bio-monitors is therefore significant in the assessment of bioaccumulation and bio-magnification of contaminants within the ecosystem (Rahman *et al.*, 2012).

Possible sources of trace metals into rivers abound. These include urban activities within the area of study. The presence of such foreign chemicals in the river, could readily affect the vital fish population. This is because fish has the tendency to bioaccumulate these hydrophobic chemicals in its tissues and organs and is a suitable indicator for environmental pollution monitoring (Udosen *et al.*, 2012).

According to Tao *et al.* (2012) the exposure assessment is a function of quantitative determination of the extent of exposure of the population to the hazardous agent. The assessment gives the real knowledge of the state of pollution of an area and dates obtained in environmental monitoring are commonly used as a starting point. Risk characterisation looks at the consequences of exposure to chemical contaminants in the environment. It supports how well the data support the conclusion about the nature and extent of risk from the contaminants.

Water bodies are continually polluted with trace metals due to domestic, urban and agricultural practice. Of certainty is the fact that when it rains, trace metals are washed into the aquatic environment, and due to their low solubility in water, they become strongly bonded to particulate matter in aquatic sediment and in turn become available to biota such as fish and humans that eat the fish (Ekeanyanwu *et al.*, 2010). Though trace metals are usually present in water at low levels, aquatic organisms have been shown to accumulate them in their tissues and also transfer to secondary feeders. Since water, sediment and fish are the ultimate reservoirs of these chemicals, they often accumulate them in such a way that can cause health problems such as kidney and liver damage, central nervous system damage, thyroid and bladder damage as well as kidney and liver cancer in animals that depend directly or indirectly on the fish and the water bodies (Akpan *et al.*, 2024). In recent times, Oruk Anam has witnessed tremendous anthropogenic activities culminating in the overall development of the area. These activities include roads construction, industrialisation transportation of all kinds and all forms of agricultural practices. The end products and the after effects of all these activities are usually washed into the aquatic environment when it rains, polluting the ecosystem with all kinds of trace metals. Many studies have been carried out on the concentration of trace metals in different portions of Qua Iboe River (Uwah *et al.*, 2013 and Udosen *et al.*, 2005). However, information on the emerging trends of these chemicals in Qua Iboe River, Oruk Anam, especially the information on fish samples are non-existent, therefore, there is every need to quantify the levels of trace metals in fish from Qua Iboe River, Oruk Anam and also to assess the health risk on humans associated with the consumption of the fish.

Quality Control and Assurance

The implementation of laboratory quality assurance and quality laboratory methods, including the use of standard operating procedures, calibrations with standards, analysis of reagent blanks and analysis of replicates guaranteed the quality of the analytical data. Each sample was analysed in triplicates. All chemicals and reagents used in the study were of analytical

grade. Standard solutions prepared were performed under clean laboratory environment. All samples were digested alongside with blanks

Statistical Analysis:

The generated data were processed using the statistical package for social sciences (SPSS).

2. Materials and Methods

2.1. Study Area

The area of this study is Qua Iboe River (QIR), Oruk Anam, Nigeria. The Map of the study area is located between latitude $04^{\circ} 28'31.0''N$ and $07^{\circ} 10'12.4''N$ and longitude $06^{\circ}52'41.2''E$ and $06^{\circ}52' 50.5''E$ of Greenwich Meridian. The river flows from Etim Ekpo through Ikot Okoro in Oruk Anam via Ibagwa in Abak to Ekpene Obom and Ekpene Ukpa in Etinan Local Government Area, then through Ndiya in Nsit Ubium local government area before flowing into Atlantic Ocean via Beight of Bonny (Udosen, 2015). It is one of the major sources of water for the inhabitants of these local government areas. The distances between sampling locations were about 200 meters apart. The sampling locations in this study as contained in the map of the study area were:

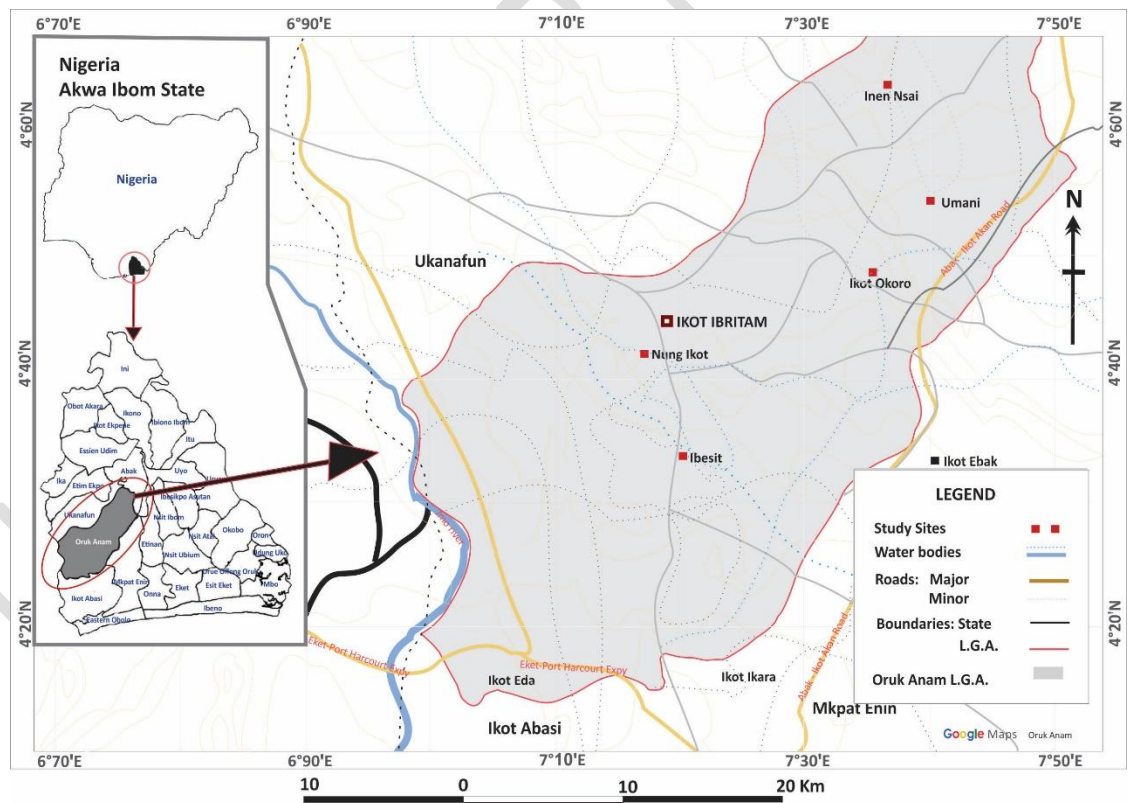


Figure1: Map of study area

Samples Collection and Preservation

Samples were collected between November and October covering wet and dry seasons from the five sampling stations namely: InenNsai, Umani, Ikot Okoro, Nung Ikot and Ibesit using InenNsai as a control station.

Water samples: The bottles were pre-rinsed several times with the water from the different sampling locations. Water samples were collected according to Udosen *et al.* (2019) by dipping each mouth of the container against the flow direction to avoid trapping air bubbles in the bottle. Water samples were collected using pre-rinsed 1-litre (1L) polyethylene bottles with caps for the analysis of water quality parameters. Water samples for dissolved oxygen determination were fixed on site with 2 mL of Winkler solutions 1 and 11. The samples for trace metals analysis were collected with HNO₃ pre-rinsed (1L) polyethylene containers and 5 mL of concentrated nitric acid (HNO₃) to a pH < 2 was added to maintain the oxidation state of the metals and to reduce microbial action. The samples collected were transported to the laboratory and preserved in a refrigerator to 4°C (Akpan *et al.*, 2024).

Fish samples: Fish samples were collected from the local fishermen fishing in the river using local fishing traps (Ikpa) set adjacent to the sampling locations at dusk and inspected at dawn (Udosen *et al.*, 2012). Five medium sized fish samples were obtained very early in the morning and were put in a locally made aquarium and transported immediately to laboratory for further analysis (Ido *et al.*, 2023). Prior to analysis, the fish samples were washed with distilled water to remove loosely held particles before they were gutted and decapitated using stainless steel kitchen knife. The decapitated fish samples were filleted on both sides, placed in a porcelain crucible and dried in an oven. The dried muscles were homogenized using mortar and pestle. The resultant homogenized fish samples were stored in a well-labeled plastic bag before digestion.

Determination of Trace Metals in Water

The concentrations of trace metals were determined using UNICAM solar 969 Atomic Absorption Spectrophotometer (AAS). The water samples (100 mL each) were filtered using filter paper No.1. The filtrate was acidified with HNO₃ (10 mL) and 50% HCl solution (10 mL). It was evaporated to near dryness on an electric hot plate. The solution was transferred to a 100 mL volumetric flask and made up to mark with deionized water. A blank was also prepared the same way with the omission of the sample using deionized water. The samples were aspirated into the air/acetylene flame.

Determination of Trace Metals in Fish

Fish samples from each sampling location were dried in an oven at 80°C for 24 hours after which they were homogenized to powder form. Exactly 2 g of each were weighed and transferred to a beaker. Into each of the beakers, trioxonitrate (v) acid (6 mL) and perchloric acid (2mL) were added and stirred. Distilled water (30mL) was also added. Each beaker was placed on the hot plate and heated for digestion to take place. After heating, the samples were allowed to cool, filtered and the volume was made up to 50 mL using deionized water. Then the digested samples were used for the analysis. The digests were prepared in triplicates and a blank determination was also made using 25 mL volumetric flask and made up to the mark with distilled water. These geochemical fractions were analysed using Atomic Absorption spectrophotometer (Udosen *et al.*, 2012).

Human Health Risk Assessment of Trace Metals Through Fish Consumption

The exposure assessment via consumption of fish was evaluated using Equations 1 and 2.

$$\text{Intake}_{(\text{seafood})} (\text{mg/kg/day}) = \frac{(\text{CF} \times \text{IR} \times \text{FI} \times \text{EF} \times \text{ED})}{(\text{BW} \times \text{AT})} \quad \text{Equation 1}$$

$$\text{HQ} = \frac{\text{Intake}}{\text{RfDo}} \quad \text{Equation 2}$$

Here, CF is the chemical concentration in fish(mg/kg), IR is ingestion rate (kg/day), FI is the fraction ingested (unitless), ED is exposure duration, EF is exposure frequency (365 days/year), BW is body weight and AT is the averaging time (period over which exposure is average in days). RfDo is the oral reference doses of the trace metals. The actual risk was characterized based on hazard quotient (HQ) as shown in equations 1 and 2. HQ below one (1) implies that the level of exposure is not likely to cause any obvious adverse effects (USEPA, 2000; Zabinet *et al.*, 2008).

Quality Assurance

The implementation of laboratory quality assurance and quality laboratory methods, including the use of standard operating procedures among others guaranteed the quality of the analytical data (Ido *et al.*, 2023).

Statistical Analysis:

The generated data were processed using the statistical package for social sciences (SPSS).

3. Results and Discussion

The results of analysis of trace metals in fish and physicochemical parameters from five sampling locations are discussed in this unit. Table 1 shows the levels of physicochemical parameters in surface water in wet and dry seasons; Table 2 shows the levels of trace metals in water in wet and dry seasons; Table 3 shows the levels of trace metals in fish in both wet and dry seasons; Table 4 shows estimated risk (in terms of HQ) of trace metals via fish consumption.

Table 1: Levels of physicochemical parameters in surface water from the sampling locations for wet and dry seasons

Sampling Locations	Temp (°C)		pH		TDS (µs/cm)		EC (mg/L)		DO (mg/L)		BOD (mg/L)		Cl ⁻ (mg/L)		NO ₃ ⁻ (mg/L)		SO ₄ ²⁻ (mg/L)		PO ₄ ³⁻ (mg/L)		NO ₂ -Salinity (mg/L)	
	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry
IN	27.01	29.01	6.50	7.20	70.50	86.00	119.13	133.33	6.13	5.53	2.05	2.04	19.01	18.20	1.32	1.32	5.13	5.18	0.23	0.23	0.01	0.02
UM	29.02	29.67	6.90	7.13	73.50	92.67	110.40	132.70	6.10	6.00	2.60	2.50	19.43	18.41	1.26	1.45	5.01	5.91	0.22	0.23	0.02	0.02
IO	30.08	30.90	6.80	7.30	96.80	10.700	119.63	145.00	8.90	7.30	3.09	3.02	21.92	19.50	1.63	1.85	5.23	6.05	0.24	0.30	0.02	0.02
NI	30.10	30.01	6.30	7.40	96.88	23.00	121.18	167.33	10.01	9.50	3.10	3.05	26.50	25.30	1.67	1.97	5.26	6.14	0.26	0.32	0.02	0.02
IB	30.02	32.11	6.40	7.41	97.73	24.00	123.36	167.67	11.50	10.40	4.50	4.01	35.20	34.60	1.74	2.04	5.29	6.15	0.31	0.33		

season and ranged between 2.03 and 4.01 mg/L with mean level of 2.92 ± 0.01 mg/L in dry season

Chloride: Chloride levels in surface water ranged between 19.01 and 35.20 mg/L in wet season and between 18.20 and 34.60 mg/L during dry season with mean levels of 24.41 ± 1.04 mg/L and 23.20 ± 5.06 mg/L in wet and dry seasons respectively.

Nitrate: Nitrate levels in surface water in this study ranged between 1.26 and 1.74 mg/L in wet season and between 1.32 to 2.04 mg/L in dry season with mean levels of 1.52 ± 0.20 mg/L and 1.61 ± 0.31 mg/L for wet and dry seasons respectively.

Sulphate: Sulphate levels in surface water in this study ranged between 5.01 and 5.29 mg/L in the wet season with mean levels of 5.18 ± 0.12 mg/L. During the dry season, the levels ranged between 5.18 and 6.15 mg/L with mean levels of 5.20 ± 0.55 mg/L

Phosphate: Phosphate levels in surface water ranged between 0.22 and 0.31 mg/L during wet season and 0.23 to 0.33 mg/L in the dry season with mean levels of 0.25 ± 0.25 and 0.28 ± 0.10 mg/L for wet and dry seasons respectively.

Nitrite: Nitrite levels in surface water in this study ranged between 0.01 and 0.05 mg/L with a mean value of 0.02 ± 0.14 mg/L in wet season and 0.02 mg/L in all locations with mean level of 0.02 ± 0.01 mg/L during dry season.

Turbidity: Turbidity levels in surface water ranged between 1.52 and 2.76 NTU with mean level of 1.85 ± 0.01 NTU in wet season and between 1.51 and 2.72 NTU with mean level of 1.80 ± 0.01 NTU for dry season

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Table 2: Levels (mg/L) of trace metals in surface water of Qua Iboe River for wet and dry seasons

Trace Metals (mg/L)		Cd	Cr	Ni	Cu	V	Pb	Sn
IN	Wet	0.02±0.01	0.01±0.01	0.01±0.01	0.02±0.01	0.01 ^a ±0.01	0.00±0.00	0.02±0.01
	Dry	0.03±0.01	0.01 ±0.01	0.02±0.01	0.03±0.01	0.02±0.01	0.00±0.00	0.00±0.01
UM	Wet	0.02±0.01	0.01±0.01	0.02±0.01	0.05±0.01	0.02±0.01	0.00±0.00	0.01±0.01
	Dry	0.04±0.01	0.0 ±0.01	0.03±0.01	0.07±0.01	0.03±0.01	0.00±0.00	0.02±0.01
IO	Wet	0.0±0.00	0.03 0.01	0.0±0.00	0.07±0.01	0.0 ±0.00	0.00±0.00	0.02±0.00
	Dry	0.06±0.01	0.04±0.01	0.01±0.01	0.08±0.01	0.04±0.01	0.01±0.00	0.02±0.01
NI	Wet	0.05±0.01	0.07±0.01	0.02±0.01	0.08±0.01	0.03±0.01	0.00±0.01	0.02±0.01
	Dry	0.07±0.00	0.08±0.00	0.04±0.01	0.10±0.00	0.04±0.01	0.01±0.00	0.03±0.00
IB	Wet	0.05±0.01	0.06±0.01	0.03±0.01	0.08±0.01	0.03±0.01	0.01±0.01	0.02±0.01
	Dry	0.08±0.00	0.09±0.00	0.04±0.00	0.10±0.01	0.04±0.01	0.01±0.00	0.03±0.01

Within column, different superscript letters mean significantly different ($p < .05$).

Levels of Trace Metals in Surface Water

Table 2 shows the levels of trace metals in water in both wet and dry seasons. The distribution pattern of the levels of trace metals in surface water indicates $Cu > Cd > Cr > V > Ni > Sn > Pb$.

Levels of Cadmium: The levels of cadmium in water samples across sampling locations during wet season ranged between 0.02±0.01 and 0.05±0.01 mg/L and between 0.03±0.01 and 0.080±0.01 mg/L during dry season.

Levels of Chromium: The levels of Cr in surface water ranged between 0.01±0.01 and 0.068±0.01 mg/L in wet season and during dry season, the levels ranged between 0.02±0.09 and 0.087±0.004 mg/L.

Levels of Nickel: The levels of Ni across sampling locations ranged between 0.01±0.01 and 0.03±0.01 mg/L in wet season and between 0.02±0.01 and 0.04± 0.00 mg/L during dry season.

Levels of Copper: The mean levels of Cu in surface water across sampling locations during wet season ranged between 0.02±0.01 and 0.08±0.01 mg/L in wet season and between 0.03±0.01 and 0.10±0.01 mg/L during dry season.

Levels of Vanadium: The mean levels of V in surface water across all sampling locations during wet season ranged from 0.01±0.01 to 0.03± 0.01 mg/L and from 0.02±0.01 to 0.04±0.01 mg/L.

Levels of Lead: The levels of Pb in water across sampling locations ranged between 0.00±0.00 and 0.01± 0.00 mg/L in wet season and between 0.00±0.00 and 0.01±0.00 mg/L during dry season

Levels of Tin: The mean levels of Sn in water across sampling locations ranged between 0.00±0.01 and 0.02±0.01 mg/L in wet season and during dry season levels obtain ranged between 0.02±0.00 to 0.03±0.01 mg/L. Exposure to high levels of inorganic tin may cause symptoms including stomach aches, liver and kidney problems, and anemia (Ahmad *et al.*, 2010).

Table 3: Levels (mg/kg) of trace metals in fish for wet and dry seasons

Trace Metals		Cd	Cr	Ni	Cu	V	Pb	Sn
IN	Wet	0.06±0.01	0.87±0.02	0.06±0.01	0.06±0.01	0.23±0.01	0.01±0.01	0.21±0.01
	Dry	0.09±0.01	0.74±0.01	0.48±0.01	0.33±0.01	0.33±0.01	0.02±0.01	0.26±0.01
UM	Wet	0.07±0.00	0.78±0.01	0.08±0.01	0.23±0.01	0.12±0.01	0.23±0.01	0.27±0.01
	Dry	0.30±0.00	0.5±0.01	0.07±0.01	0.50±0.01	0.32±0.01	0.23±0.00	0.29±0.01
IO	Wet	0.48±0.01	0.8±0.01	0.03±0.01	0.41±0.01	0.45 ±0.01	0.31±0.01	0.37±0.01
	Dry	0.30±0.01	0.53±0.01	0.04±0.01	0.51±0.01	0.04±0.01	0.41±0.00	0.38±0.01
NI	Wet	0.48±0.01	0.56±0.01	0.03±0.01	0.61±0.01	0.33±0.01	0.43±0.01	0.41±0.01
	Dry	0.51±0.01	0.67±0.01	0.03 ^a ±0.01	0.51±0.01	0.4 ±0.01	0.45±0.01	0.51±0.01
IB	Wet	0.58±0.01	0.87±0.01	0.03±0.01	0.44±0.01	0.6 ±0.01	0.45±0.00	0.56±0.01
	Dry	0.53±0.01	0.71±0.01	0.03±0.01	0.50±0.01	0.4 ±0.01	0.45±0.01	0.40±0.01

Levels of Trace Metals in Fish

The levels of trace metals obtained in fish varied across the seasons as presented in Table 3. The distribution pattern was Cu > Cd > Cr > Sn > Ni > V > Pb. The level of Cu recorded was highest compared to other trace metals studied.

Levels of Cadmium: The levels of Cd in fish across the sampling locations ranged between 0.06±0.01 and 0.78±0.01 mg/kg in wet season and between 0.90±0.00 and 0.73±0.01 mg/kg during dry season.

Levels of Chromium: The levels of Cr in fish ranged between 0.87±0.02 and 0.87±0.01 mg/kg during wet season and between 0.74±0.00 and 0.71±0.01 mg/kg in dry season.

Levels of Nickel: The levels of Ni in fish ranged between 0.93±0.01 and 0.03±0.01 mg/kg in wet season while in dry season the levels ranged between 0.76±0.00 and 0.91±0.01 mg/kg.

Level of copper: The levels of Cu in fish ranged between 1.06±0.01 and 14.98±0.01 mg/kg in wet season and between 2.93±0.01 and 16.54±0.001 mg/kg in dry season.

Levels of Vanadium: The levels of V obtained in fish samples in this study ranged between 0.63±0.01 and 5.56±0.01 mg/kg in wet season and between 0.93±0.01 and 5.43±0.020 mg/kg during dry season.

Levels of Lead: The levels of Pb in fish in the study during wet season ranged between 0.01±0.01 and 0.98±0.01 mg/kg while in dry season the range was between 0.12±0.00 and 0.98±0.01 mg/kg.

Levels of Tin: The levels of Sn obtained in fish across all the sample locations ranged between 0.67±0.01 and 4.56 mg/kg in wet season and between 0.76±0.01 and 5.43±0.01 mg/kg during dry season. Highest mean level of tin 5.43 ±0.01 mg/kg was obtained at Ibesit sampling location

Table 4: Risks assessment in terms of hazard quotient (HQ) of trace metals via fish consumption

TM	HQ(wet)						HQ(dry)					
	Ibesit	Nung Ikot	Ikot Okoro	Umani	InenNsai	Ibesit	Nung Ikot	Ikot Okoro	Umani	InenNsai		
Cd	1E-2	0.31	0.31	0.25	0.24	0.22	0.12	0.12	0.24	0.57	0.11	
Cr	3E-2	0.20	0.20	0.09		0.07	0.33	0.04	0.03	0.20	0.28	0.05
Ni	2E-2	0.22	0.22	0.02		0.11	0.05	0.48	0.28	0.22	0.14	0.08
Cu	4E-2	0.42	0.32	0.25		0.15	0.07	0.43	0.42	0.20	0.11	0.08
V	1E-2	0.27	0.24	0.24		0.22	0.17	0.45	0.45	0.40	0.28	0.17
Pb	3E-2	0.05	0.05	0.05		0.19	0.19	0.05	0.05	0.05	0.03	0.19
Sn	1E-5	0.40	0.40	0.28		0.22	0.11	0.45	0.45	0.45	0.28	0.11

Human Health Risk of Trace Metals via Fish Consumption

Calculation of exposure (due to oral intake) of trace metals through fish consumption from Qua Iboe River are presented in Table 4. The result revealed the following trends of exposures across all locations and seasons: Cu > Cd > Cr > Ni > V > Sn > Pb. Also revealed was that levels were higher though not significantly around Ibesit, Nung Ikot and Ikot Okoro (downstream) compared to Umani and InenNsai due to intense anthropogenic activities downstream. The estimated risks of trace metals via fish consumption from the studied river as presented in Table 4 indicated no risk as hazard quotients for the metals determined in fish were less than unity (HQ <1) across all the sampling locations. It therefore means that there is no health risk implication for consumers of commercial fish from the river.

4. Conclusion

The following conclusions were drawn based on the statistical analyses of the results. The water and fish samples obtained from the five sampling locations of Qua Iboe River contain variable levels of the investigable parameters. The investigated parameters show seasonal variations in the samples. Correlation analysis revealed positive relationships between Cd with Cr, Ni, Cu, V and Pb in the water, fish and sediment in both the dry and wet seasons. There were also positive relationship between the trace metals in water and sediment, water and fish as well as sediment and fish in the two seasons. The levels of physicochemical parameters of the water and sediment were all within permissible limits set by WHO, FEPA and NIS. The levels of Cd, Cr, Ni, V, Pb and Sn in the water were all within WHO and NIS guidelines except for copper. . Risk assessment of the trace metals through fish consumption from the studied river indicated low risk.

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