

## GOLD MINING IN IGUN-IJESHA, SOUTHWEST NIGERIA: IMPACTS AND IMPLICATIONS FOR WATER QUALITY

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### ABSTRACT

This study was carried out in Igun-Ijesha, Osun State, Nigeria to determine the likely hotspots of water contamination for future treatment trials. Water quality analyses were based on physico-chemical and heavy metal parameters of surface and ground water collected from the study community. A total of thirty-eight water samples were collected between September 2012 and February 2013 and analyzed using standard procedures. Concentrations of heavy metals were determined using Atomic Absorption Spectrophotometer. Results showed that water samples within the gold mining community are contaminated and the hydrochemistry varied with seasons. The values of magnesium, turbidity, total dissolved solids, electrical conductivity and pH during the period of sampling ranged 3.1-42.1 mg/L, 0-150 NTU, 30-560 mg/L, 80-1192  $\mu$ S/cm and 5.95-8.55 respectively. Chloride, nitrate, phosphate, sulphate and sodium data were within the stipulated limits set by the Nigerian Standards for Drinking Water Quality (NSDWQ). Heavy metal contents of the groundwater were generally higher than those from surface water sampled within the mining district. The values of arsenic, cadmium, chromium, copper, lead, manganese, nickel and zinc, ranged from 0.01-1.20, 0.05-0.52, 0.80-34.80, 0.09-4.30, 0.09-8.30, 0.05-3.94, 0.05-19.6 and 1.80-29.90 mg/L respectively. Most of the listed metals have values exceeding the international and national recommended limits. The daily intake of water in the study area poses a potential health threat from long-term exposure to heavy-metals. The study suggests that water safety plans should be developed to safeguard water resource and public health within the mining community.

**Keywords:** Gold Mining, Heavy Metal Pollution, Igun-Ijesa, Surface and Groundwater, Water Quality, Water Safety

### 1. INTRODUCTION

Water is one of the essentials that supports all forms of plant and animal life and it is generally obtained from two principal natural sources; groundwater and surface water such as fresh water lakes, rivers, streams (Awomeso *et al.*, 2010). However, one of the most important environmental issues today is water contaminated with heavy metals because of strong toxicity even at low concentrations (Marcovecchio *et al.*, 2007). Water contamination with light and heavy metals are mainly determined by natural (i.e., weathering,

erosion of bed rocks and ore deposits) and anthropogenic (e.g., mining, smelting, industries, agriculture and waste water irrigation) processes (Ettler *et al.*, 2011; García-Lorenzo *et al.*, 2011; Muhammad *et al.*, 2011; Sracek *et al.*, 2011). These processes degrade water quality and impair their use for drinking, domestic, agricultural and industrial purposes (Krishna *et al.*, 2009). The common heavy metals that have been identified in polluted water include arsenic, copper, cadmium, lead, chromium, nickel, mercury and zinc. The release of the metals without proper treatment poses significant danger to

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water and public health. The danger lies in two aspects. Firstly, heavy metals have the ability to persist in natural ecosystems for an extended period. Secondly, the metals have the ability to accumulate in successive levels of the biological chain, thereby causing acute and chronic diseases (Nomanbhay and Palanisamy, 2005; Lone *et al.*, 2008). Recently, pollution of general environment has increasingly gained a global interest. Contamination of surface and groundwater in gold mining communities is a serious environmental problem in many countries, Nigeria inclusive (Aslibekian and Moles, 2003). Heavy metal contamination of soil and water and related health impact on residents is a persistent social issue and several studies have identified human health risks subject to abandoned mines (Chung *et al.*, 2005; Bada *et al.*, 2012).

In view of the various challenges of pollution, conformation with drinking water quality standards is a concern because of the ability of water to spread diseases within a large population. Although the standards vary from country to country, the objective anywhere is to reduce the possibility of spreading water borne diseases to the barest minimum in addition to being pleasant to drink, which implies that water must be wholesome and palatable in all respects (Edema *et al.*, 2001; Awomeso *et al.*, 2010). In Igun Ijesha gold city, the community depend on surface and groundwater contaminated from mining activities and so far, there is no evidence of any water quality assessment. Therefore, the aim of this study is to evaluate the physicochemical parameters and trace metals in drinking water sources in Igun-Ijesha gold mine area. The assessment is expected to determine water quality status of drinking water in the study area to ensure source and water safety.

## 2. MATERIALS AND METHODS

### 2.1. The study Area

Igun-Ijesha gold city lies between latitudes 7°30' and 7°35' N and between longitudes 4°38' and 4°42' E in Atakumosa West Local Government Council southwestern Nigeria (**Fig. 1**). The study area is a rural community of about 2,400 to 2,600 people that engage in predominantly in subsistence farming and cocoa plantation. Igun Ijesha is a community with many dilapidated buildings. The city is accessible through a poorly erected bridge. Mapping of the community was done with assistance of the community members. The mapping exercise reveals local knowledge of resources, land use and settlement patterns.

The mine locations are within one of the six (6) classes of the Basement Complex rock that is from slightly migmatized to non-migmatized, meta-

sedimentary and meta-igneous rock or simply called the Schist belt. The study area is a part of Ilesa-Ife schist belt (Ademeso *et al.*, 2013). The belt is one of the 11 schist belts documented by Ademeso *et al.* (2013). The area has two contrasting lithologies separated by NNE trending Ifewara fault zone. The west of the fault is occupied by the amphibole schist, amphibolites, talc-tremolite and pelitic rocks (TML, 1996). The eastern part has quartzite, quartz schist and amphibole schist, The gold deposit occur in the eastern area, thus, the three Local government areas lie on the east of Ifewara fault zone. Gold occurs with ores such as: Pyrite, pyrrhotite and minor chalcopyrite, galena, sphalerite, magnetite and ilmenite. Adjacent to the gold bearing veins the host granite-gneiss has been hydrothermally altered to a sericitechlorite epidote assemblage (with also hematite and pyrite) (NMC, 1987).

### 2.2. Field Investigation and Water Sampling

Igun-Ijesha area was selected for this study primarily due to the presence of gold mining activities in the community. Three surface water and three groundwater sampling points were selected and their coordinates located using a Global Positioning System GARMING 45XLS. The sampling points are shown in **Fig. 1**.

Sampling was done between September 2012 and February 2013. A total of thirty-eight water samples were collected from both surface and ground water samples in the study area. Water samples were collected with 1.5 L plastic bottles, which had been soaked in 70% nitric acid for 24 h and rinsed thoroughly with double distilled water. Samples for trace metal analyses were put into 250 mL plastic bottles and 2 mL concentrated Nitric acid added to it. Collected samples were preserved and stored in an ice-chest at a temperature of 4°C and transported to the laboratory for analyses. Samples were taken in separate containers for physicochemical and trace metal analysis respectively. Samples for trace metal analysis were each preserved with 0.5 mL of concentrated nitric acid before transporting to the Central laboratory, University of Ibadan for analysis. During sampling, relevant information like the ambient temperature (31°C), date of Sampling, time of sampling and seasons of the year were recorded.

### 2.3. Sample Analysis

The methods of laboratory analysis used were those specified in International analytical standards such as APHA for water quality. All equipments were duly calibrated with standards and samples were analysed in two replicates. All tests and laboratory analyses were carried out at the ACEME Analytical Laboratory Limited Canada and International Institute for Tropical Agriculture (IITA) Laboratory, Ibadan.

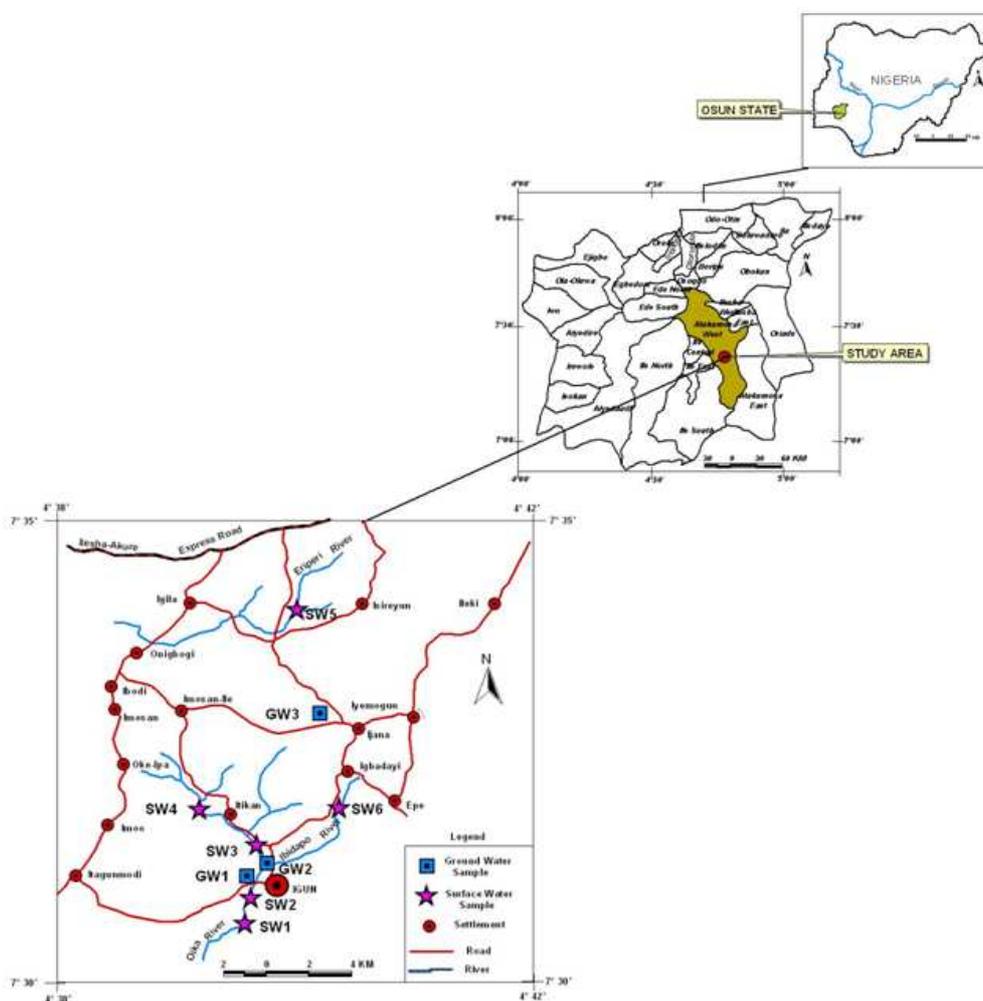


Fig. 1. Map of the study area showing the sampling locations

#### 2.4. Determination Physico-Chemical Parameters

Water pH, temperature, Electrical Conductivity (EC), TDS were analyzed in-situ during sampling using pH/TDS/Conductivity meter. Turbidity was determined using a Spectrometer (HACH DR/2000). Samples for water soluble anions (sulphate, nitrate, phosphate and chloride) were determined with Ion Chromatography System (ICS) model Dionex ICS 2000. The element used was potassium hydroxide. The ICS analytical column and guard column are AS II HC (2×250 mm) and AG II HC (2×250 mm), respectively (Thermo Fischer Scientific Inc, 2012). Samples for cationic water soluble constituents (calcium, magnesium and potassium) were analysed with Dionex DX 500 with CS 12A Analytical column (4×250 mm) and CG 12A guard column. Details of analytical

procedures of both anionic and cationic species can be found in (Taiwo, 2013; Gashi *et al.*, 2013). Standard calibration curves were generated with series of known salt concentrations in the series of 5, 10, 20, 50 and 100 ppm.

#### 2.5. Sample Digestion for Heavy Metal Analysis

Samples for the determination of arsenic, cadmium, chromium, copper, lead, manganese, nickel and zinc were collected with 500 mL plastic bottles, since such metal may be adsorbed on the wall of glass bottles. About 3 mL of concentrated Nitric acid was added and the samples were refrigerated at 4°C before digestion. The water samples (100 mL) were digested with 10 mL concentrated HNO<sub>3</sub>. The mixture was then heated on a hotplate for 30 min (USEPA, 1989). The extracts were filtered and made to 100 mL with distilled deionised water. The ready

digests were sent to ACEME analytical laboratory Canada for heavy metal determination using Inductively Coupled Plasma-Mass Spectrometer (ICPMS).

## 2.6. Metal Analysis

Heavy metals were determined with ICPMS (Agilent 7500ce) at the ACEME Analytical Laboratory in Canada. The ICPMS was equipped with octopole reaction system, which is effective in removing interfering species. Standards were prepared from VWR standard soluble prepared in the series of 5, 10, 20, 50 and 100 ppb. The procedures of ICPMS can be read elsewhere in (Taiwo, 2013; Gashi *et al.*, 2013).

## 3. RESULTS

### 3.1. Physical and Chemical Parameters

**Table 1 (a and b)** show the data collected for individual surface and ground water samples while **Table 2 (a and b)** summarizes the physico-chemical parameters with respective WHO/NSDWQ standards. The measured pH gives the general indication that the water samples range from neutral to alkaline for wet season and slightly acidic during dry season. The highest desirable level for pH stipulated for drinking and domestic purposes is within the range of 6.5 to 8.5 (WHO, 2004). Electrical conductivity values in all the water samples varied from 103  $\mu\text{S}/\text{cm}$  (SW5) to 1192  $\mu\text{S}/\text{cm}$  (GW3) for dry season and 80-620  $\mu\text{S}/\text{cm}$  in GW2 and GW1 samples respectively, for wet season. With the exception of GW3 sample with the value of 1192  $\mu\text{S}/\text{cm}$ , all other water samples are within the permissible limit of 1000  $\mu\text{S}/\text{cm}$  for EC in drinking water (WHO, 2004).

The turbidity values in the water samples varied between 5 FTU (GW1 and GW2) to 150 FTU (SW6) in rainy season and 0 FTU (GW1 and GW3) to 33 FTU (SW6) in dry season. The turbidity values for the groundwater for all the water samples except GW3 (at dry season) are within the recommended WHO limit of 5 FTU.

TDS values in the sampled water bodies range from 43 mg/L (SW5) to 560 mg/L (GW2) for rainy season samples while the values in the dry season, concentrations of TDS varied from 30 mg/L (SW5) to 270 mg/L (GW1). The TDS values recorded for ground and surface water samples in both seasons are within the WHO limit of 500 mg/L (WHO, 2004) with the exception of GW3 groundwater recording the value of 560 mg/L.  $\text{Mg}^{2+}$  concentration varied from 5.1 mg/L (GW2) to 42.12 mg/L (GW1) during the rainy season and from 3.1 mg/L (GW2) to 38.21 mg/L (GW1) during the dry season exceeding the recommended limit of 0.2 mg/L set by the Nigerian Standard for Drinking Water Quality (SON, 2007). Phosphate concentrations in the water samples range from 0.09 mg/L in SW3 to 0.53 mg/L in SW2 (dry season) and 0.11 mg/L in GW1, GW2 and SW6 to 0.51 mg/L in SW2 (wet season). There is no WHO guideline value to compare the measured phosphate values.

The summary of both surface and ground water physico-chemical parameters during the wet and dry seasons are presented in **Table 2(a and b)**. Most of the observed parameters have shown higher mean values at the wet season compared to dry season. The mean values of potassium have shown an exception, with higher values obtained in the dry season.

**Table 1a.** Physico-chemical parameter values in both surface and ground water samples during the rainy season in Igun-Ijesha, Osun state, Nigeria

Water Samples	Temp (°C)	pH	EC ( $\mu\text{S}/\text{cm}$ )	TDS (mg/L)	Turb. (FTU)	$\text{Cl}^-$ (mg/L)	$\text{NO}_3^-$ (mg/L)	$\text{PO}_4^{3-}$ (mg/L)	$\text{K}^+$ (mg/L)	$\text{SO}_4^{2-}$ (mg/L)	$\text{Ca}^{2+}$ (mg/L)	$\text{Mg}^{2+}$ (mg/L)	$\text{Na}^+$ (mg/L)
SW1	26.1	8.14	100	100	9	8.71	0.04	0.21	4.12	13	16.19	10.14	11.02
SW2	26.3	8.05	110	100	18	6.16	0.03	0.51	3.08	22	16.32	12.25	11.41
SW3	26.1	8.06	100	98	21	5.23	0.04	0.20	3.10	13	15.51	12.24	10.52
SW4	26.2	7.90	110	91	22	6.38	0.03	0.12	2.98	15	20.06	13.10	10.15
SW5	26.3	8.53	160	43	29	6.23	0.02	0.24	4.78	4	9.30	6.72	7.24
SW6	26.7	7.71	90	86	150	6.43	0.02	0.11	3.11	4	10.41	9.61	10.01
GW1	25.7	8.53	620	68	5	71.02	0.02	0.11	150.70	80	65.05	42.12	24.62
GW2	25.6	7.50	80	390	5	6.23	0.02	0.11	3.15	4	25.16	5.10	6.05
GW3	25.2	8.55	270	560	9	30.40	0.04	0.13	57.63	53	39.60	15.89	15.35
WHO STANDARD		6.5-8.5	1000	500	5	250.00	50.00	3.00	n.a	100	n.a	0.2.00	200.00

Note: SW1: Oika River 1, SW2: Oika River 2, SW3: Oika River 3, SW4: Oika River 4, SW5: Eriper River, SW6: Justice Ibidapo River, GW1: Igunwell 1, GW2: Igun well 2, GW3: Ijana well, n.a: Not available, Temp-temperature, Turb-Turbidity

**Table 1b.** Physico-chemical parameter values in both surface and ground water samples at dry season in Igun-Ijesha, Osun state, Nigeria

Water Samples	Temp (°C)	pH	EC (µS/cm)	TDS (mg/L)	Turb. (FTU)	Cl <sup>-</sup> (mg/L)	NO <sub>3</sub> <sup>-</sup> (mg/L)	PO <sub>4</sub> <sup>3-</sup> (mg/L)	K <sup>+</sup> (mg/L)	SO <sub>4</sub> <sup>2-</sup> (mg/L)	Ca <sup>2+</sup> (mg/L)	Mg <sup>2+</sup> (mg/L)	Na <sup>+</sup> (mg/L)
SW1	27.1	6.54	218	40	6	4.81	0.05	0.21	4.29	11	15.19	9.27	10.97
SW2	28.2	6.22	218	50	14	5.57	0.03	0.53	4.05	21	15.48	10.19	11.14
SW3	28	6.5	210	40	21	4.60	0.05	0.09	3.23	13	14.51	10.13	10.28
SW4	27.9	6.37	205	50	14	4.38	0.03	0.1	3.23	11	19.07	11.13	10.25
SW5	27.3	6.62	103	30	19	4.17	0.01	0.24	4.88	2	8.95	5.59	6.72
SW6	26.7	6.04	183	40	33	4.43	0.01	0.1	3.28	2	9.73	8.36	9.99
GW1	26.1	6.64	138	270	0	65.59	0.01	0.12	158.66	65	63.02	38.21	24.18
GW2	26.4	6.18	818	40	1	4.25	0.01	0.12	4.81	2	21.08	3.10	6.55
GW3	26.3	5.95	1192	120	0	28.76	0.04	0.13	56.11	43	37.63	13.99	14.85
WHO		6.5-8.5	1000	500	5	250.00	50.00	3.00	n.a	100	n.a	0.20	200.00

STANDARD

Note: SW1: Oika River 1, SW2: Oika River 2, SW3: Oika River 3, SW4: Oika River 4, SW5: Eriper River, SW6: Justice Ibidapo River, GW1: Igunwell 1, GW2: Igun well 2, GW3: Ijana well, n.a: Not available, Temp-temperature, Turb-Turbidity

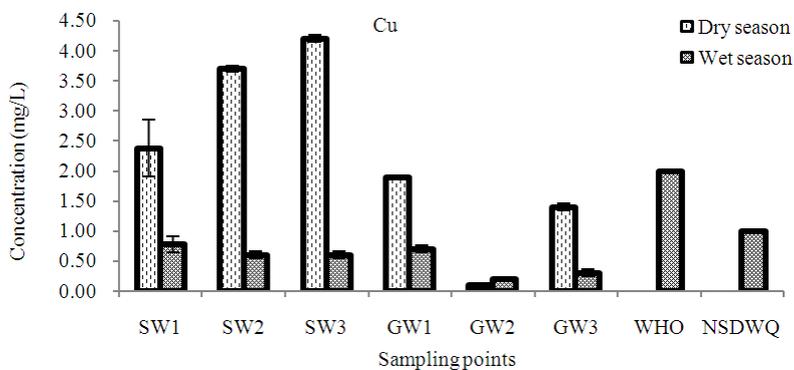
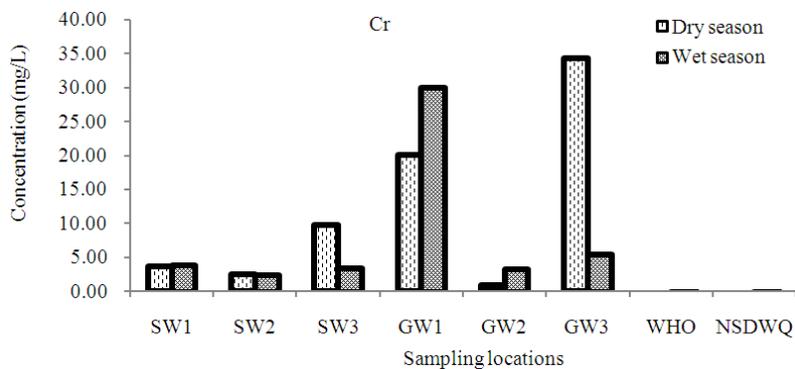
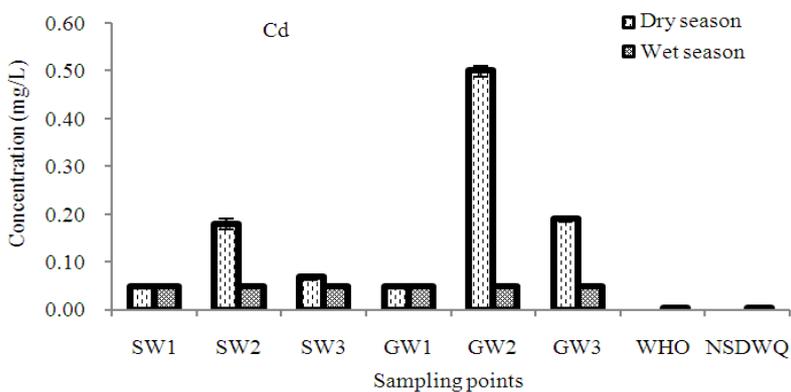
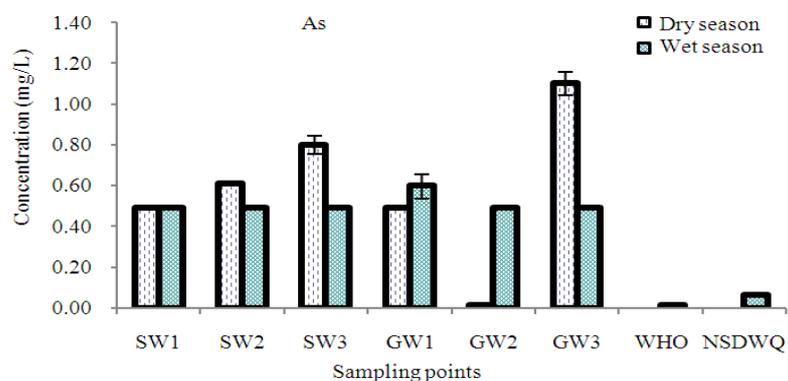
Note: Data in bold were higher than the WHO standard in drinking water

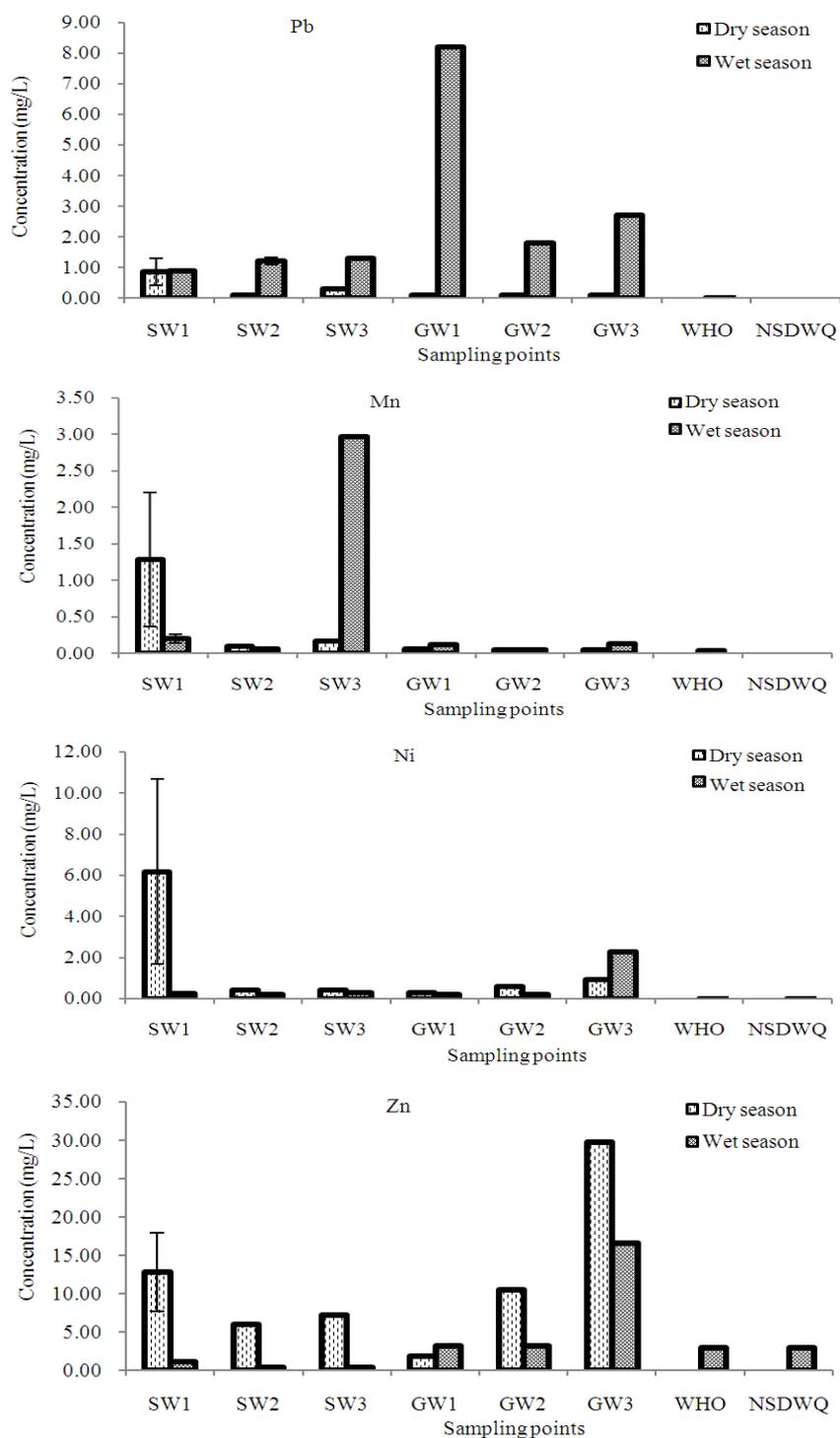
**Table 2a.** The summary of surface water parameters during the wet and dry seasons for Igun-Ijesha, Osun state, Nigeria (n = 38)

Variables	Dry season				Wet season			
	Min.	Max.	Mean	Std Dev	Min.	Max.	Mean	Std Dev
Temp (°C)	26.70	28.20	27.53	0.59	26.10	26.70	26.28	0.22
pH	6.04	6.62	6.38	0.22	7.71	8.53	8.07	0.27
EC (µS/cm)	90.00	160.00	111.67	24.83	103.00	218.00	189.50	44.29
TDS (mg/L)	30.00	50.00	41.67	7.53	43.00	100.00	86.33	21.95
Turbidity (mg/L)	6.00	33.00	17.83	9.06	9.00	150.00	41.50	53.55
Chloride (mg/L)	4.17	5.57	4.66	0.50	5.23	8.71	6.52	1.16
Nitrate (mg/L)	0.01	0.05	0.03	0.02	0.02	0.04	0.03	0.01
Phosphate (mg/L)	0.09	0.53	0.21	0.17	0.11	0.51	0.23	0.15
Potassium (mg/L)	3.23	4.88	3.83	0.69	2.98	4.78	3.53	0.75
Sulphate (mg/L)	2.00	21.00	10.00	7.21	4.00	22.00	11.83	6.91
Calcium (mg/L)	8.95	19.07	13.82	3.82	9.30	20.06	14.63	4.04
Magnesium (mg/L)	5.59	11.13	9.11	1.96	6.72	13.10	10.68	2.36
Sodium (mg/L)	6.72	11.14	9.89	1.62	7.24	11.41	9.89	1.62

**Table 2b.** The summary of ground water parameters during the wet and dry seasons for Igun-Ijesha, Osun state, Nigeria (n = 38)

Variables	Dry season				Wet season			
	Min.	Max.	Mean	Std Dev	Mini.	Max.	Mean	Std Dev
Temp (°C)	26.10	26.40	26.27	0.15	25.20	25.70	25.50	0.26
pH	5.95	6.64	6.26	0.35	7.50	8.55	8.19	0.60
EC (µS/cm)	80.00	620.00	323.33	273.92	138.00	1192.00	716.00	534.35
TDS (mg/L)	40.00	270.00	143.33	116.76	68.00	560.00	339.33	249.88
Turbidity (mg/L)	0.00	1.00	0.33	0.58	5.00	9.00	6.33	2.31
Chloride (mg/L)	4.25	65.59	32.87	30.88	6.23	71.02	35.88	32.74
Nitrate (mg/L)	0.01	0.04	0.02	0.02	0.02	0.04	0.03	0.01
Phosphate (mg/L)	0.12	0.13	0.12	0.01	0.11	0.13	0.12	0.01
Potassium (mg/L)	4.81	158.66	73.19	78.33	3.15	150.70	70.49	74.61
Sulphate (mg/L)	2.00	65.00	36.67	31.97	4.00	80.00	45.67	38.53
Calcium (mg/L)	21.08	63.02	40.58	21.12	25.16	65.05	43.27	20.20
Magnesium (mg/L)	3.10	38.21	18.43	17.97	5.10	42.12	21.04	19.04
Sodium (mg/L)	6.55	24.18	15.19	8.82	6.05	24.62	15.19	8.82





**Fig. 2.** Mean values of heavy metals in (mg/L) of both the surface and groundwater bodies in Igun-Ijesha, Osun state, Nigeria. The bars indicate the standard deviation. Note: SW1: Oika River, SW2: Eriperi River, SW3: Justice Ibadapo River, GW1: Igun well 1, GW2: Igun well 2, GW3: Ijana well

### 3.2. Heavy Metal Concentrations

The mean values of heavy metals determined in the water samples at both seasons are shown in **Fig. 2**. The concentration of arsenic ranges 0.49-0.70 mg/L during the rainy season to 0.01-1.20 mg/L in dry season samples; cadmium concentration is between 0.05 and 0.50 µg/L, with the highest value observed in GW2 sample during the dry season and the lowest concentration in SW1 during the wet season. The concentrations of Cr range from 2.30-30.50 mg/L during the rainy season and 0.80-34.80 mg/L during the dry season, which exceeded the recommended limit set by the WHO and NSDWQ. The values of Cu range from 0.18-1.10 mg/L (rainy season) to 0.09-4.30 mg/L (dry season); nickel values are 0.10-2.40 mg/L (rainy season) and 0.05-9.60 mg/L (dry season); while zinc range 0.49-16.70 mg/L (rainy season) to 1.8-29.90 mg/L (dry season). The concentration of Pb ranged from 0.80-8.30 mg/L (rainy season) to 0.09-0.31 mg/L (dry season).

## 4. DISCUSSION

Based on the guidelines, the pH of water sources from the study area may not adversely affect its suitability for drinking and domestic purposes, as well as aquatic ecosystem. On the contrary, groundwater pH at location GW3 during the dry season is below the WHO limit suggesting possible acidity. Schafer *et al.* (2009) and Ayantobo *et al.* (2013) in previous studies reported that low pH in groundwater may be due to natural geochemical and biochemical processes within the aquifers. The GW3 sampling point is within the mining and processing facilities, it is thus possible that seepage of effluent discharges, as well as surface and agricultural run-off may contribute to the increased concentration of ions. The elevated EC content may also be due to the weathering of sulphide-bearing rocks, which is facilitated by the on-going mining activities (Marcovecchio *et al.*, 2007; Ayantobo *et al.*, 2013). Turbidity is a major concern for surface water in particular during the rainy seasons (Schafer *et al.*, 2009). Elevated turbidity values are expected during the wet season due to erosion of organic materials and solids into the water bodies ([http://water.epa.gov/scitech/wastetech/guide/stormwater/upload/2006\\_10\\_31\\_guide\\_stormwater\\_usw\\_b.pdf](http://water.epa.gov/scitech/wastetech/guide/stormwater/upload/2006_10_31_guide_stormwater_usw_b.pdf)). Water turbidity can also be attributed to the presence of clays in surface water. Pathogens and bacteria can be measured in turbidity (Schafer *et al.*, 2009).

According to Karikari and Ansa-Asare (2006), the palatability of water with TDS level less than 600 mg/L is generally considered to be good whereas water with TDS

above 1,200 mg/L becomes increasingly unpalatable. The highest TDS concentration recorded at GW3 may be due to seepage of effluent discharges, as well as agriculture and domestic waste substances (Ayantobo *et al.*, 2013).

This study has found higher concentration of  $Mg^{2+}$  than that reported by Muhammad *et al.* (2011) for drinking water in the Kohistan region, Northern Pakistan.  $Mg^{2+}$  occurs naturally in water after passage through mineral deposits and rock strata and suggests that  $Mg^{2+}$  contributes to total hardness of water in the study area.

Comparing the values of phosphate concentration obtained in this study with that of natural water (usually between 0.005 and 0.02 mg/L (Taiwo *et al.*, 2010; DebRoy *et al.*, 2012); it can be concluded that, this study has recorded elevated phosphate values in groundwater samples. High concentrations of phosphate could indicate the presence of pollution and are largely responsible for eutrophication conditions in water bodies (Taiwo *et al.*, 2012). The high phosphate loads recorded at the sampling points may be attributed to inflows from eroded materials carried from waste rock dumpsite (Taiwo *et al.*, 2010; DebRoy *et al.*, 2012). Elevated Phosphate level could also be related to the weathering of calcium-bearing minerals largely facilitated by mining activities in the area (Singh *et al.*, 2008).

Calcium and potassium are not a health concern and are not regulated (Schafer *et al.*, 2009). Although, the reported values may not implicate the public health, however, the presence of multivalent cations such as Ca may pose negative treatment's implications for certain processes (Schafer *et al.*, 2009).

Arsenic concentrations are well above the recommended WHO limit of 0.01 mg/L (except GW2 sample). The Arsenic concentration could be due to mineral dissolutions such as pyrite oxidation. Previous study of Kortatsi (2004) has reported Arsenic concentrations of 0.05 and 0.14 mg/L in ground and surface water. Elevated amount of Arsenic observed in this can be damaging to public health on persistent exposures through consumption. Arsenic is a known carcinogen (Roy and Saha, 2002). Skin cancer has been associated with long-term exposure to arsenic through drinking water (Kortatsi, 2007). Early clinical symptoms of acute intoxication of Arsenic include abdominal pain, vomiting, diarrhoea, muscular pain and body weakness (Tchounwou *et al.*, 2003). Cadmium occurs naturally with zinc and lead in sulphide ores. Cd concentrations in unpolluted natural waters are usually below 1.0 µg/L. In this study, Cd concentrations in surface and groundwater at both seasons are higher than the permissible limit. The guideline value for cadmium is given as 0.003 mg/L in

drinking water by both the World Health Organization (WHO, 2004) and the Nigerian standard for drinking water quality (SON, 2007). Previous studies show maximum levels in groundwater to be 0.003 mg/L (Kortatsi, 2004; Armah *et al.*, 2010) and 0.06 mg/L (Oluwasanya and Martins, 2006). Maximum levels in surface water were less than 0.05 mg/L (Kuma and Younger, 2004; Yem *et al.*, 2013). The observed cadmium values show that water quality in the study area is questionable and unfit for human consumption. As a practical measure, the guideline is set as 0.05 mg/L, which is considered to be unlikely to give rise to significant risks to health (WHO, 2004). Maximum levels in groundwater has been shown to be 0.07 mg/L (Kortatsi, 2004; Marcovecchio *et al.*, 2007) and 0.06 mg/L (Oluwasanya and Martins, 2006) and in surface water to be 0.49 mg/L (Kuma and Younger, 2004, Marcovecchio *et al.*, 2007). The observed elevated chromium concentrations represent an identified hazard, which calls for necessary control measures to mitigate consequent health impacts.

Copper is both an essential nutrient and a drinking-water contaminant. During the rainy season, copper concentration in the surface and groundwater sampled are within the acceptable limits of 2.0 and 1.0 mg/L set by WHO and NSDWQ, respectively. Exceptions are Justice Ibidapo River (SW3), Eriperi River (SW2) and Oika River (SW1); the observed values are below the permissible level. The present findings show much lower values than the concentrations of copper in the Indus River, Pakistan (5.0 mg/L) and Mississippi, USA (River (4.0 mg/L). Lead is a possible human carcinogen and it is also a cumulative poison so that any increase in the lead burden should be avoided. The Pb values in this study revealed clear exceedencies relative to the permissible limit of 0.001 mg/L set by the WHO. Previous studies also show maximum levels in groundwater to be 0.03 mg/L (Kortatsi, 2004, Armah *et al.*, 2010) and in surface water to be <0.05 mg/L (Kuma and Younger, 2004; Yem *et al.*, 2013). A provisional tolerable daily intake is set as 3.5 µg of lead per kg of body weight for infants. Human health concerns associated with lead intoxication in children include brain damage, behavioral problems, anaemia, liver and kidney damage and hearing loss (Gohar and Mohammadi, 2010; Rajaganapathy *et al.*, 2011) whereas in adults poor muscle coordination, nerve damage to the sense organs, increased blood pressure, hearing and vision impairment, reproductive problems and retarded fetal development (Surendran and El-Fawal, 2008). In this respect, the lead content in the surface and groundwater within the mine area are dangerous for human health and aquatic life.

Manganese concentrations above 0.1 mg/L can impact an undesirable taste to drinking water. Effects from manganese are primarily to the central nervous system (Wasserman *et al.*, 2006; Bhattacharya *et al.*, 2008; Bada *et al.*, 2012). Surface water bodies investigated in this study had Mn concentrations (0.06-2.97 mg/L) above the WHO guideline value of 0.04 mg/L (WHO, 2004). Ground water sources sampled in the study area also recorded Mn concentrations (0.05-0.13 mg/L) above WHO guideline. The high Mn concentration level (2.97 mg/L) detected in Ibidapo River, which is within vicinity of the mines can be attributed to anthropogenic sources such as effluent discharges and acid-mine drainage from mining activities. It can also be linked to natural geochemical and biochemical processes such as weathering of manganese bearing minerals and rocks within the aquifers (Singh *et al.*, 2008). Nickel concentrations in drinking water are normally below 20 µg/L, although levels up to several hundred micrograms per litre in groundwater and drinking water have been reported (Obiri *et al.*, 2010). The concentrations of nickel observed in the present study are above the permissible limit of 0.07 mg/L for WHO standard and 0.02 mg/L of NSDWQ for domestic water (SON, 2007). The observed nickel values also exceed the findings of Kortatsi (2004), Oluwasanya and Martins (2006) who found maximum levels in groundwater to be 0.08 mg/L and 0.34 mg/L respectively. The presence of nickel in the study area is a chemical hazard to both aquatic biota of the river as well as for human consumption. Zn concentration of the surface water sampled during the rainy season are within the recommended limits of 3 mg/L set by the WHO and NSDWQ while value from sampling point GW1 during the dry season also are within the limit. Zinc is an essential trace element found in virtually all food and potable water in the form of salts or organic complexes (Edema *et al.*, 2001; WHO, 2003).

## 5. IMPLICATIONS FOR MINING MANAGEMENT POLICES

The findings of this study hold several implications for water quality management and policy. Previously, most mining communities depended on surface water as drinking water source. However, contamination of surface water particularly via mining activities made it imperative for government and other non-state stakeholders to resort to groundwater (Armah *et al.*, 2010). Results from this study and other studies (Obiri *et al.*, 2010) have shown that the quality of groundwater is similarly, questionable. The results identifying many water

quality hazards also revealed that the surface and groundwater extracted from the vicinity of the mine cannot be considered safe for particularly drinking and other domestic purposes. Policy makers need to be aware for appropriate regulations that would make it mandatory to analyze drinking water for physical and chemical parameters in mining communities on a regular basis. Where water sources have been tested, communities should be notified of contaminant levels so as to inform apt household or communal treatment solutions and daily decision-making regarding access to safe drinking water.

## 6. CONCLUSION

The main goal of this study was to assess the effect of mining activities on water quality sampled at Igun Ijesha gold mine district. The results from this study showed that water resources (surface and ground water) within the vicinity of the study area are contaminated. The hydrochemistry of both surface and groundwater show seasonal variation probably due to natural variations in geology and the mining activities. The results also indicated that the values of most of the observed physicochemical parameters of water samples are found within the standards set by the WHO. Elevated values are observed for most parameters in the wet season while  $K^+$  revealed higher concentrations in the dry season. The heavy metal concentrations are generally higher than the WHO recommended limit indicating threats to public health.

This study, therefore, recommends that activities of miners should be monitored to ensure that gold extraction and other mining activities do not occur close to drinking water sources. The miners and the entire community should be educated on the health risk associated with human exposure to trace metals to prevent pollution of water bodies. The recommended immediate control measures to the identified water supply and quality problem in the study area is provision of alternative water supply and treatment of metal contaminated water, in the short and medium term to protect health and well being of the rural people living in the vicinity of the mine. Igun-Ijesha is located within the humid tropics with 7-8 months of adequate rainfall. Rainwater harvesting for drinking purposes has been generally reported as a common practice (Oluwasanya, 2006). Rainwater harvesting may thus be considered as an alternative option for water supply in Igun-Ijesha. Piped water supply may be considered as the ultimate goal for safe

water provision in the study area such that safe water can be delivered in close proximity of the consumers. Apart from hand-dug wells and boreholes, other water supply options for the household level includes solar distillation, solar disinfection, sand house hold filters and engaging the services of water vendors. Overall, the results presented in this study indicate the critical need for a clearly laid out water safety planning to mitigate public health risk in the study area.

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