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A study of heavy metals pollution in the coastal marine sediment of Ondo State, Nigeria

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ABSTRACT

The present study was carried out to assess the pollution status of heavy metals in the marine sediment in Ondo coastal area. A total of 36 sediment samples were collected from the three locations (Awoye, Abereke and Ayetoro) once in a month in 2015 covering both dry and wet season. The sediment samples were subjected to digestion and Atomic Absorption Spectrophotometer (AAS) was used to measure the concentrations of As, Cd, Cr, Cu, Fe, Mn, Ni, Pb, V and Zn. The measured concentrations data were used for evaluating the contamination level and accumulation status of heavy metals in sediment by employing several pollution indices: contamination factor (CF), degree of contamination (C_{deg}), pollution load index (PLI), enrichment factor (EF) and index of geoaccumulation (I_{geo}). The results showed that Fe had the highest average concentration values of 248.00 and 324.33 mg/kg at Awoye sampling site for wet and dry season, respectively while the highest mean concentration of Mn were 271.77±9.50 and 295±10.06 mg/kg at Abereke and Ayetoro, respectively. The contamination factor (CF) values for As, Cd and Mn were 1 < CF < 3, indicating a moderate contamination. The results of C_{deg} were ranged from 8.6 to 12.5 for both dry and wet season except in Ayetoro site that had 7.8 in the wet season. The PLI values were 0.47, 0.50 and 0.54 (<1) for Abereke, Awoye and Ayetoro sites, respectively, denoting unpolluted conditions. The EF values for Pb and Ni obtained from all the sites ranged from 20.8 to 38.9, showing their significantly enriched in sediment. The I_{geo} values of As, Cr, Fe, V, Zn and Cu were less than one ($I_{geo} < 1$) for the three sampling locations in both seasons, implying that the sediments in the study area are practically uncontaminated by these metals. Pb had the I_{geo} values of 2.80 and 2.85 during dry and wet season at Awoye sampling site while Ni had 2.57 at Abereke only in the dry season. The heavy metals in sediment might be derived from the upstream rivers from the top soil, mechanically weathered rock materials and anthropogenic activities.

Keywords: Sediment, Heavy metals, Pollution, Indices.

1. INTRODUCTION

Sediment is a mixture of weathered materials and mineral species from the parent rocks. It contains organic debris that are transported by detrital process from the upstream and deposited at the river bed [1]. Sediment, because of their variable physical and chemical properties acts as repository for heavy metals in the aquatic environment [2]. Naturally, sediment pollution due to heavy metals are caused by geologic weathering of the bedrock and direct atmospheric deposition. The interactions between water and crustal materials with which water is in contact is a major process by which heavy metals content in the sediment can be increased [3]. The anthropogenic activities such as sea transportation, energy generation and utilization, dredging, fishing, oil exploration, farming, infrastructural oil exploration, farming, infrastructural development and mining are crucial for social-economic reasons [4], however, they are probable sources of sediment pollution when their wastes are discharged into the river body by urban surface water runoff [5]. The

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34 increased concentration and accumulation of heavy metals alter sediment quality as well as
35 the food stuff available for the aquatic organism, leading to loss of aquatic biodiversity [6].

36 In recent years, heavy metals pollution of the aquatic environment has become a
37 worldwide problem due to their persistence in nature and their capacity to accumulate in
38 living organism [7]. Ahmadipour et al. [8] reported that heavy metals have negative health
39 implications due to their toxicity potential and tendency to accumulate in environmental
40 media such as water, soil, sediment and other biological media. Several heavy metals
41 persist in the sediment with varied concentrations most especially in fine grain [9]. The
42 increased concentration of the heavy metals in the sediment has adverse health implication
43 due to possibility of entering the food chain through aquatic organism uptake and direct
44 consumption of aquatic organism by man [10]. Polluted sediment can act as a metal pool,
45 thereby release metals to the overlying water column via natural or anthropogenic
46 processes, causing potential adverse health effects to the ecosystem and degrade the
47 sediment quality [11]. Heavy metals are not destructible and non-biodegradable, thus they
48 may exist in an environmental medium for a long period of time [12]. Moreover, heavy metals
49 in the sediment can undergo sorption and complexation, ion exchange, dissolution and
50 precipitation reactions which influence their behavior and bioavailability during transportation
51 [13].

52 The capacity of the sediment to accumulate heavy metals make them relevant
53 indicators for monitoring purposes as well as the evaluation of their pollution levels and
54 patterns of the aquatic system [14]. Based on these, several studies had been conducted on
55 degradation of sediment quality due heavy metals using several pollution indices [15, 16, 17,
56 18] while few studies were based on heavy metals speciation and potential bioavailability in
57 the sediment. In the study area, quite a few studies had been undertaken on the distribution
58 of heavy metals in the sediment [19]. In this study, the concentrations of heavy metals in the
59 sediment samples were determined using Atomic Absorption Spectrophotometer (AAS)
60 technique. This was with a view to assessing the pollution status due to heavy metals in the
61 sediment. The main objective of this work was to assess the level of heavy metal pollution in
62 the sediment samples by employing the combination of pollution indices: contamination
63 factor (CF), degree of contamination (D_{deg}), pollution load index (PLI), index of geo-
64 accumulation (I_{geo}) and enrichment factor (EF) analysis.

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66 **2. MATERIALS AND METHODS**

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68 **2.1 Study area**

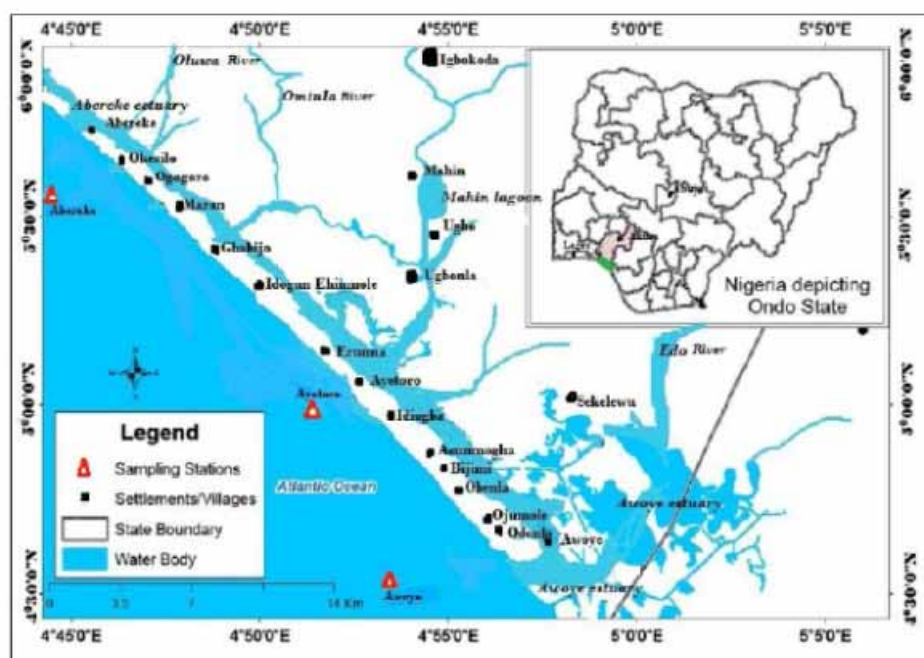
69 Ondo State coast is strategically located along the gulf of guinea as the
70 Transgressive Mud Beach (TMB) lying in the east of the West African lagoon system and on
71 the west of the Benin Flank of the Niger Delta Basin. Most of the creeks and rivers (Omila
72 and Edo) in the inland area are drained into the Atlantic ocean through Awoye and Abereke
73 estuaries. Also, the sandy beach materials of the Nigeria coastal plain are replaced by mud
74 and lacustrine deposits in Ondo state coast. Other unique features of this coast are overlying
75 mud of about 60 m and massive incursion of the sea into the inland swamps. It is one of the
76 longest coastlines in Nigeria (more than 10 % of the country coastline) which favors fishing
77 activities in the riverine areas. The major anthropogenic activities around the study area are
78 transportation services, seismic investigations, crude oil exploration, commercial and
79 agricultural activities. The study area lies within the tropical rain forest zone and bounded by
80 the latitude 5.56 - 6.30°N and longitude 4.40 - 5.43°E with 21.38 m above the sea level. The
81 study area is characterized with two distinct seasons namely: dry and wet season. The dry
82 season spans through the month of November to March while wet season occurs between
83 April to October with the annual rainfall of about 3000 mm [20].

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85 **2.2 Sample collection and preparation**

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87 Sediment samples were collected at approximately 2.0 nautical mile from the Ondo
 88 state coastline at 10 m depth using a VanVeen grab sampler. A representative samples
 89 were taken with plastic spatula from the middle of the grab bulk sediment. Samples were
 90 collected once in a month from January to December, 2015. Three sampling locations were
 91 chosen based on their proximity to various anthropogenic activities such as oil exploration,
 92 ship breaking yard, market activities and fishing and the areas of incursion of the estuaries
 93 into the Atlantic ocean. Figure 1 shows the map of the Ondo section of the southwest coast
 94 of Nigeria with the study locations (Abereke, Ayetoro and Awoye) and other settlements
 95 along the coast. After sampling, each sample was transported to the laboratory and air dried
 96 at room temperature for two weeks, then grounded in a pre-washed agate mortar and pestle.
 97 In order to remove coarse debris, the samples were screened through a 100 µm stainless
 98 steel sieve. The samples of the fine-grained sediment were retained for chemical analysis.
 99 All the sampling materials were washed with water and clean with acetone after each
 100 sampling to avoid possible contamination.



101 Figure 1: Map of the study area showing the sampling sites
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104 2.3 Elemental analysis

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 106 The samples were subjected to wet digestion in Foss Tecator digestion vessel. One
 107 (1) gram of sediment sample was added in a 15 mL mixture of nitric (HNO₃) and perchloric
 108 acid (HClO₄) in ratio 1:3. The whole content was placed on a heating electric plate at about
 109 334 - 350°C for 2 hours. The mixed solution was boiled until the evaporation of the acid
 110 solution and a clear solution was obtained. Upon cooling, the solution was filtered to a 100
 111 mL volumetric flask and then diluted to 50 mL with distilled de-ionized water and kept in the
 112 sample bottle at 4°C before analysis. The concentrations of As, Cd, Cr, Cu, Fe, Mn, Ni, Pb,
 113 V and Zn were determined using Atomic Absorption Spectrophotometer (Buck Scientific
 114 VGP 210 Model) at the Central Science Laboratory (CSL), Obafemi Awolowo University, Ile-
 115 Ife, Nigeria. All reagent were of analytical grade. The operating conditions consisting of

116 maximum wavelength λ_{max} , energy, current and slit width of the ASS instrumental analytical
 117 set up are presented in table 1. The detection limits of As, Cd, Cr, Cu, Fe, Mn, Ni, Pb and Ni
 118 were 0.05,0.001, 0.005, 0.001, 0.005, 0.001, 0.003, 0.003 and 0.002 mg/kg, respectively.
 119 The heavy metal concentrations data were subjected to descriptive statistic using the
 120 statistical package for social science software (SPSS 16.0).

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122 **2.3.1 Contamination factor and Degree of contamination**

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124 The contamination factor (CF) is a single pollution index which indicates the
 125 contamination level of a particular toxic substance or elements in a given environmental
 126 medium [21]. In this study, the sediment contamination due to As, Cd, Cr, Cu, Fe, Mn, Ni,
 127 Pb, V and Zn were assessed using the contamination factor (CF). The CF of i^{th} heavy metal
 128 was calculated as:

$$129 \quad C_f^i = \frac{C_s^i}{C_m^i} \quad (1)$$

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131 where C_s^i is the mean concentration of the element i^{th} . C_m^i is the background concentration
 132 which is the maximum level of that metal in a given, beyond which the medium is said to be
 133 contaminated [22]. In this study, Taylor and McLennan [23] continental crustal average data
 134 was used as the background concentration. The CF is classified into four main groups: CF
 135 <1 means low contamination; $1 \leq CF < 3$ means moderate contamination; $3 \leq CF \leq 6$ implies
 136 considerate contamination and $CF > 6$ indicates very high contamination [10]. The degree of
 137 contamination (C_{deg}) is an indicator that shows the extent to which the environmental media
 138 is polluted. The C_{deg} is estimated as the sum of the contamination factors ($C_{deg} = CF_1 + CF_2$
 139 $+ CF_3 + CF_n$) of all the heavy metals in the sediment for a site. Four major category of the
 140 degree of contamination have been identified: < 8, 8-16, 16-32 and > 32 indicate low,
 141 moderate, considerable and very high degree of contamination, respectively [22].

142

143 Table 1: The operating conditions of the AAS instrumental set up

144

Elements	λ_{max} (nm)	Current	Energy	Slith width
As	193.00	10	75	0.7
Cd	228.80	4	69	2.7/0.80
Cr	357.87	25	97	2.7/0.80
Cu	324.75	15	80	2.7/0.80
Fe	248.33	30	66	1.8/1.35
Mn	279.48	20	62	1.8/1.30
Ni	232.00	25	67	1.8/1.35
Pb	283.31	10	75	2.7/1.05

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153 **2.3.2 Pollution load index**

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155 The pollution load index (PLI), as proposed by Tomlinson et al. [24], is a simple
 mathematical model use to determine the pollution status of a particular site taking into

156 consideration the metal concentrations of that site as a single value. In this study, the PLI
157 was evaluated as:

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$$159 \quad PLI = \sqrt[n]{CF_1 * CF_2 * CF_3 * \dots * CF_n} \quad (2)$$

160

161 CF is the contamination factor of each of the heavy metals obtained from equation 1 while n
162 is the number of heavy metals under consideration. The PLI values could be < 0, > 0 and 1
163 implying unpolluted condition, progressive degradation in the sediment quality and the
164 occurrence of baseline pollution level, respectively [24].

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166 **2.3.3 Enrichment factor (EF) analysis**

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168 The Enrichment Factor (EF) is used to evaluate the magnitude of the elements in air, water,
169 soil and sediment samples. It provides information on the relative abundance of species in a
170 given medium to the background values. It is also employed to assess the degree of
171 pollution and to differentiate the elements of anthropogenic and natural sources [25]. Its
172 approach is based on the standardization of the measured element against a reference
173 element. The EF_x was estimated using the equation:

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$$175 \quad EF_x = \frac{\left[\frac{X_s}{E_{s(ref)}} \right]}{\left[\frac{X_c}{E_{c(ref)}} \right]} \quad (3)$$

176 Where EF_x is the enrichment values for the element x. X_s and X_c are concentrations of the
177 element of interest in the sample and in the crust while $E_{s(ref)}$ and $E_{c(ref)}$ are the
178 concentrations of the reference element used for normalization in the sample and in the
179 crust. A reference element is often characterized by low occurrence variability and stable
180 chemical properties. The common reference elements for EF analysis are Sc, Mn, Ti, Al, Si
181 and Fe. In this study, Fe was selected as the reference element due to its lithogenic origin
182 and abundance. Taylor and McLennan [23] continental crustal average data was adopted as
183 the background concentration. The EF values were categorized as < 2, 2-5, 5-20, 20-40 and
184 ≥ 40 and considered as deficiency to minimal, moderate, significant, very high and extremely
185 high enrichment, respectively [26].

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187 **2.3.4 Index of Geoaccumulation**

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189 the index of geoaccumulation (I_{geo}) was originally proposed by Muller [28]. It is a
190 quantitative index which indicates the metal contamination in soil, water and sediment [1].
191 The (I_{geo}) compares the measured concentration C_m of the elements in a given sample with
192 the geochemical background concentrations B_m for the element m in the sample [27]. The
193 I_{geo} was calculated using the logarithmic function:

$$194 \quad I_{geo} = \log \left[\frac{C_m}{1.5 B_m} \right] \quad (4)$$

195 Taylor and McLennan [23] continental crustal average data was used as the background
196 elemental concentrations. The factor 1.5 was introduced to minimize the effect of possible
197 variations in the background values, B_m , which may be attributed to the lithogenic variations
198 in the sediment [1, 27, 22]. In this study, I_{geo} was estimated using the seasonal average
199 values of the measured elemental concentrations. The categories of I_{geo} values are <0, 0-1,

200 1-2, 2-3, 2-4, 4-5 and >5 and their respective interpretations are practically unpolluted,
201 unpolluted to moderately polluted, moderately polluted, moderately to strongly polluted,
202 strongly polluted, strongly to extremely polluted and extremely polluted [28].

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204 **3 RESULTS AND DISCUSSION**

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206 **3.1 Average elemental concentrations of heavy metal results**

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208 Table 2 shows the results of the seasonal average concentrations of As, Cd, Cr, Cu,
209 Fe, Mn, Ni, Pb, V and Zn. Fe had the highest average concentration values of 248.00 ± 20.00
210 and 324.33 ± 5.80 mg/kg at Awoye sampling site for wet and dry season, respectively. The
211 highest mean concentration of Mn is 271.77 ± 9.50 and 295 ± 10.06 mg/kg at Abereke and
212 Ayetoro, respectively. The high concentrations of Fe and Mn are most likely related to the
213 local mineralogy and lithogenic origin, rather than the anthropogenic activities [35]. Fe had
214 been reported as being an elements whose origin is fundamentally natural and one of the
215 most common elements in the earth's crust [29]. The mean concentrations of Fe and Mn
216 were similar to high values of 463.0 and 321.4 mg/kg for both wet and dry season reported
217 by Iqbal and Shah [30] in the sediment samples obtained from Khanpur Lake, Pakistan. A
218 few difference in seasonal concentrations of Cd, Cr, Ni, V and Zn could be related to their
219 regional deposition and the heavy metal accumulation rate in the sediment. The erosion
220 activities, runoff by the action of water and land based sources might introduced heavy metal
221 into the water body. Atmospheric deposition of metals from anthropogenic sources has been
222 reported to be the probable sources for the variation in metal concentrations in the sediment
223 in study area [19]. Most industries such as oil exploration which are located around the sites
224 discharged industrial waste directly into the water without treatment. This might contain
225 heavy metals such Pb, Ni, V, Cr and Cd. The commercial ship and industrial flying boat for
226 the transportation of the workers during oil exploration and seismic investigation could also
227 release significant amount of Pb, Cd, V and Ni-containing contaminants into the sediment. In
228 each site, the average values of Cd, Cu and Cd were lower than the World Health
229 Organization [31] sediment quality guidelines values of 6, 25 and 123 mg/kg, respectively.
230 These show the unpolluted conditions of the sediment by Cu, Cr and Cd. The Ni and Cr
231 mean concentration values in each site exceeded the [31] values of 20 and 25 mg/kg,
232 suggesting that a minute pollution due to Ni and Cr.

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235 **3.2 Contamination factor, degree of contamination and pollution load** 236 **index results**

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238 Table 3 shows the results of contamination factor, degree of contamination and
239 pollution load index for the wet and dry season of the three studied areas. In the dry season,
240 Cr, Cu, Fe, Ni, Pb, V and Zn had a CF values to be < 1 , signifying the low contamination
241 status of the sediment. The CF values for As Cd and Mn were $1 < CF < 3$, indicating a
242 moderate contamination. The results of the CF showed similar trend in the three sites both
243 seasons indicating that CF values were less influenced by the seasons. The results of C_{deg}
244 ranged from 8.6 to 12.5 for both dry and wet season except in Ayetoro site that had 7.8. This
245 classified the sediment from the three sites as a moderate degree of contamination. The PLI
246 values were 0.47, 0.50 and 0.54 (< 1) for Abereke, Awoye and Ayetoro sites, respectively,
247 denoting unpolluted (perfection) conditions [24]. The little seasonal variations in PLI values
248 might be associated with upstream discharged into the water and the alteration in various
249 phases of elements in solution. In the wet season, a similar pattern of CF, C_{deg} and PLI
250 Table 2: Average elemental concentrations of the heavy metals results (mg/kg)

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	Dry			Wet		
	Abereke	Awoye	Ayetoro	Abereke	Awoye	Ayetoro
As	2.67±0.12	4.33±0.23	2.47±0.06	5.27±0.15	6.97±0.32	3.3±0.20
Cd	0.57±0.06	1.02±0.03	0.31±0.03	0.53±0.03	0.67±0.03	0.32±0.03
Cr	0.32±0.02	0.33±0.06	0.02±0.01	0.71±0.02	1.15±0.01	0.45±0.01
Cu	2.10±0.10	1.26±0.29	1.17±0.06	2.73±0.06	1.30±0.18	2.40±0.10
Fe	185.67±5.80	324.33±5.80	126.67±5.80	153.00±10.00	248.0±20.00	108.33±11.50
Mn	271.77±9.80	265.77±14.00	215.53±10.00	273.57±9.50	288.2±9.20	295.00±10.60
Ni	7.09±0.02	7.24±0.03	3.07±0.06	7.63±0.04	8.44±0.13	4.39±0.12
Pb	0.71±0.02	2.05±0.01	0.05±0.01	0.63±0.02	2.08±0.08	0.10±0.02
V	3.02±0.02	4.73±0.06	2.33±0.06	4.13±0.02	5.03±0.02	3.07±0.03
Zn	11.23±0.23	28.30±0.20	1.27±0.06	17.43±0.06	22.0±0.10	3.60±0.09

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Table 3: Contamination Factor, degree of contamination and pollution load index results

	Dry			Wet		
	Abereke	Awoye	Ayetoro	Abereke	Awoye	Ayetoro
As	1.3	2.2	1.2	2.6	3.5	1.7
Cd	3.2	5.7	1.7	2.9	3.7	1.8
Cr	0.3	0.1	0.1	0.1	0.2	0.5
Cu	0.1	0.2	0.2	0.4	0.3	0.4
Fe	0.3	0.6	0.6	0.3	0.4	0.2
Mn	2.7	2.7	2.2	2.7	2.8	2.9
Ni	0.1	0.1	0.4	0.2	0.2	0.6
Pb	0.2	0.2	0.5	0.1	0.2	0.1
V	0.8	0.4	0.4	0.2	0.3	0.2
Zn	0.3	0.3	0.5	0.2	0.3	0.4
C _{deg}	9.3	12.5	7.8	9.7	11.9	8.6
PLI	0.47	0.50	0.54	0.43	0.56	0.52

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values were observed in the three sites. This shows that the pollution of sediment is most likely less affected by the season. Although, there might be progressive accumulation of the heavy metals in the sediment which depends on the increased man-made activities such as oil exploration and waste discharge into the river from the upstream location. The wet season in 2015 was a period of heavy rainfall, leading to high fluvial inputs. The transport of the industrial waste alongside with the soil materials by the erosion activities into the ocean should increase the pollution of the sediment by the heavy metals concentration in the sediment during wet season. However, in this study, the results of CF, C_{deg} and PLI showed that the sediment quality was less affected by wet season.

3.3 Enrichment factor and index of geoaccumulation results

The results of the enrichment factor (EF) analysis for the three locations are presented in the Table 4. The EF values of As and Cr were <2 for the three sites, signifying deficiency to minimal enrichment. Zn and V had EF between $2 < EF < 5$ and $5 < EF < 10$ for dry and wet season, respectively, indicating moderate and significant enrichment while Cd and Mn had the EF values of $5 < EF < 20$ in the three sites, classifying them as significant enrichment in both seasons. The moderate and significant enrichment of Zn and V could also be related to the anthropogenic sources such as boat exhaust systems, antifouling paints and fossil fuel from mechanized boat used in fishing and transportation which were common in the study area.

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Table 4: Enrichment factor results

The EF values for Pb and Ni were > 20 in all the sampling sites at dry and wet season, indicating that the sites were highly enriched with Pb and Ni. The high enrichment of Pb and Ni might be linked to the fact that Pb and Ni are mostly anthropogenic elements associated with oil pollution which is prevalent in the study area. The enrichment of Pb might also be

	Dry			Wet		
	Abereke	Awoye	Ayetoro	Abereke	Awoye	Ayetoro
As	1.6	1.5	1.4	2.4	1.5	2.2
Cd	12.1	12.4	9.6	13.6	8.3	11.6
Cr	1.2	0.1	1.9	3.2	2.9	2.7
Cu	10.0	5.7	0.9	6.8	2.1	4.0
Mn	7.4	5.2	8.6	9.0	5.5	13.8
Ni	25.7	21.4	26.3	33.6	20.7	27.3
Pb	33.8	38.9	35.0	36.4	22.5	28.2
V	5.0	4.2	4.8	8.3	5.1	8.7
Zn	5.3	3.7	4.9	10.1	6.2	9.2

286 attributed stable isotopic nature of Pb in crustal materials and it takes a considerable period
287 of time before an appreciable depletion can occur. The difference in the EF values of heavy
288 metals in the coastal sediment might be related to the difference in the magnitude of input for
289 each metal in the sediment as well as difference in the removal rate of each metal from the
290 sediment [32, 33]. Largely, the low EF values of As and Cr indicating that they are mostly
291 derived from soil. The high EF values of Zn, V, Pb and Ni might be attributed to contribution
292 from anthropogenic activities such as oil exploration in the study area. Table 4 shows the
293 results index of geoaccumulation (I_{geo}) of As, Cd, Cr, Cu, Fe, Mn, Ni, Pb, V and Zn. The I_{geo}
294 values of As, Cr, Fe, V, Zn and Cu were negative for the three sampling sites in the both
295 seasons. These indicated that the sediment in the study area are practically uncontaminated
296 by As, Cr, Fe, V, Zn and Cu. This is in agreement with the study of Salah et al. [1] that
297 obtained negative I_{geo} values for most metals in the sediment samples of Euphrates River in
298 Iraq. Also, the negative I_{geo} 261 values signify that the average concentrations of heavy
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Table 5: Index of geoaccumulation results

	Dry			Wet		
	Abereke	Awoye	Ayetoro	Abereke	Awoye	Ayetoro
As	-1.73	1.40	0.79	-1.05	0.50	0.50
Cd	0.75	1.33	0.17	0.67	0.14	0.14
Cr	-0.56	-1.53	-1.22	-0.76	-4.34	-4.34
Cu	0.62	0.12	-1.54	0.88	-2.26	-2.26
Fe	-1.74	-0.18	-2.28	-0.93	-2.12	-2.12
Mn	0.26	0.26	0.34	0.26	0.03	0.03
Ni	2.57	1.55	1.03	1.57	1.67	1.67
Pb	1.78	2.84	1.08	1.65	2.85	1.88
V	-0.13	0.32	-1.12	0.18	0.37	-0.39
Zn	-0.07	0.86	-1.20	0.37	0.61	-2.25

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metals in sediment are lower than the average crustal concentration [1]. The I_{geo} values for Cd, Mn, Ni and V were $0 < I_{geo} < 1$ implying uncontaminated to moderately contaminated [27]. Pb has the I_{geo} values of 2.8 and 2.85 during dry and wet season at Awoye sampling site while Ni has 2.57 at Abereke only in the dry season. The accumulation of heavy metal in sediment might be derived from the discharge of Omila and Edo rivers into the ocean and the mechanically weathered surface materials.

309 Generally, the results of all the pollution indices employed in this study agreed well.
310 However, few differences exist among the pollution indices which could be attributed to their
311 various approach of computation since the same background concentrations data were
312 used. The CF, C_{deg} and PLI results all confirmed the uncontaminated status of most heavy
313 metal in all the sites. The results of EF revealed that sediment were enriched by Pb, Cd and
314 Ni but the I_{geo} values showed that the the sediments were uncontaminated to moderately
315 contaminated with respect to Pb and moderately contaminated with Cd. Also, the EF values
316 of Cd and Mn were $5 < EF < 20$, classifying the sites as significant enriched by Cd and Mn in
317 both seasons. Meanwhile, Cd and Mn had the I_{geo} values in between 0 and 1 ($0 < I_{geo} < 1$)
318 indicating uncontaminated to moderately contaminated. This implies that the calculations of
319 I_{geo} had more reliability than those of EF. The differences in the results of the I_{geo} and EF in
320 this study might be associated with the methods of their computations. The nature of the I_{geo}
321 computation, which involved the logarithmic function and matrix correction factor of 1.5 was
322 quite different from enrichment factor which normalized heavy metal concentration as the
323 ratio to another constituents in the sediment. The difference in classification and
324 interpretation of the each class might also contribute to little differences in the results.
325

326 **4. CONCLUSION**

327
328 In this study, the bottom sediment samples were obtained in Abereke, Awoye
329 and Ayetoro sites in Ondo state coastal marine area, Southwestern Nigeria in order to
330 study the pollution status of heavy metals contents. The elemental concentration of heavy
331 metals analysis of the sediment samples was performed using Atomic Absorption
332 Spectrophotometer (AAS). The heavy metals concentrations data were processed using
333 statistical analysis and pollution indices: contamination factor (CF), degree of contamination
334 (D_{deg}), pollution load index (PLI), enrichment factor (EF) and index of geoaccumulation (I_{geo}).
335 Fe had the highest average concentration values of 248.00 ± 20.00 and 324.33 ± 5.80 mg/kg
336 at Awoye sampling site for wet and dry season, respectively. Cr, Cu, Fe, Ni, Pb, V and Zn
337 had a CF value to be < 1 , signifying the low contamination status of the sediment. The PLI
338 values were < 1 for Abereke, Awoye and Ayetoro sites, respectively. The EF values of As
339 and Cr were < 2 for the three sites. The I_{geo} values of As, Cr, Fe, V, Zn and Cu were
340 negative for the three sampling sites in the both season, implying practically uncontaminated
341 condition. Generally, the results of pollution indices due to heavy metals showed low
342 pollution status. Despite the low pollution level, continuous monitoring of the heavy metals is
343 necessary to provide information on the alteration in the sediment quality due to the
344 possibility of gradual increase in heavy metals loading in the sediment.
345

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