

Geochemical Characterization of Shallow Lagoon Sediments

Omoregbe, O.A^{*}, Oloyede, D.A^{**}, Ekuerugbe, L.O^{**}, and Nkim, A.G^{*}

^{*} Resemco Limited, Port Harcourt
^{**} Geology Department, University of Benin,

DOI: 10.29322/IJSRP.10.05.2020.p10148
<http://dx.doi.org/10.29322/IJSRP.10.05.2020.p10148>

Abstract- Geochemical characterization of the Epe lagoon sediments was carried out in order to determine their depositional environment, provenance and weathering history. A total of Nineteen (19) samples were obtained from the upper 2-3m of the lagoon base and analyzed for Elemental composition using the Atomic Absorption Spectrophotometry (AAS) method. Laboratory analysis revealed values of the major oxides as follows: SiO₂ (52.99%-98.87%), Al₂O₃ (0.71%-35.46%), Fe₂O₃ (0.04-2.96%), MgO (0.01%-0.11%), CaO (0.01-0.42), Na₂O (0.01%-0.08%), K₂O (0.03%-0.91%), TiO₂ (0.01%-4.56%), P₂O₅ (0.01%-0.33%), MnO (0.01%-0.45%). Very low average values of Al₂O₃/SiO₂ (0.08932) and relatively high average value of SiO₂/Al₂O₃ (64.33) indicated quartz enrichment which was corroborated by the high SiO₂ values recorded. TiO₂/Al₂O₃ values (0.004-0.8604) indicated that the sediments are chemically matured. While geochemical analysis using specialized ternary plots revealed that the sediments were fine to medium grained clayey sands deposited in both continental and transitional environments with intensive weathering at the source area.

Index Terms- Geochemical, Lagoon, Provenance, Sediments,

I. INTRODUCTION

Geochemical studies have been used extensively in determining concentrations of heavy metals and other harmful elements within sediments. The Lagos lagoon which consists of three main segments (Lagos harbor, the Metropolitan and Epe division) has attracted a lot of geochemical studies arguably because of its location within the commercial city of Lagos which harbours over 75% of the Industrial outfits of Nigeria (Olatunji and

Abimbola, 2010). However, since geologic processes affect the chemical composition of sediments, geochemical studies have also been widely used in characterizing sediments. Modern approaches in monitoring sediment provenance especially through discriminatory diagrams based on major and trace elements relationship has proved effective (Bauluz *et.al*, 2000). The geochemical behaviour of elements within sediments indicates substantial changes in the physical, chemical and biological conditions of the environment (Avramidis *et. al*, 2010). Although various studies have been carried out by several workers in the Lagos Lagoon (Badejo *et.al*, 2014; Olatunji and Abimbola, 2010; Balogun *et.al*, 2011; Aderinola *et.al*, 2009; Ayola and Kuton 2009; Don-pedro *et.al*, 2004; Okoye *et.al*, 1991; Ogunsua *et.al*, 1993; Osibanjo and Bamgbose, 1990; Oyewo *et.al*, 1982), not much emphasis have been made on the Epe division segment. Also much of the previous studies have focused on using geochemical data to ascertain the impact of both Industrial and anthropogenic waste on the sediments as well as aquatic life.

This current study would focus on using geochemical data in characterizing sediments of the Epe division segment of the Lagos Lagoon in order to determine their depositional environment, provenance, weathering history and also update on existing available geochemical information.

The study Area

The Epe division segment of the Lagos lagoon (also called the Epe lagoon) lies within Latitude 6°29'N to 6°38'N and Longitude 3°30'E to 4°05'E. It is located in Lagos, one of Africa's densely populated cities. The lagoon is fed by river Oshun and opens into

the gulf of guinea through the Lagos harbour. It also has a surface area of about 225km² with maximum depth of about 6m. However, a large area of the lagoon is relatively shallow with minimum depth of about 1m. Geologically, the Epe lagoon lies within the eastern axis of the Dahomey sedimentary Basin. The regional geologic setting, basin evolution, tectonic elements, stratigraphy and structural styles of the Eastern Dahomey Basin has been dealt with by various authors (Storey, 1995; Mpanda, 1997; Omatsola and Adegoke, 1981; Ojeda, 1982; Adediran and Adegoke, 1987; Slansky, 1962; Billman, 1992; Adegoke, 1969; Ogbe, 1972; Agagu 1985).

II. MATERIALS AND METHODS

Sample collection

A total of nineteen (19) samples were obtained from the upper 2-3m of the lagoon base (*i.e., the sediment-water interface*) with sample points appropriately located using the Global Positioning System (Fig.1) (Table 1).

Sample preparation and sieving

After collection, the samples were drained of water and air-dried for one week. The dried samples were then disaggregated and the unwanted materials subsequently removed. In order to determine the granulometric characteristics of the samples, 100g of air-dried soil sample was soaked in 1 litre solution of distilled water with 2g of calgon and the whole content was shaken regularly and agitated to ensure effective disaggregation. The soaked sample was then left for 24 hours and thereafter washed through sieve 2.36mm opening a little at a time. The washing continued until the water passing through the sieve was clear. The soil retained on the sieve was oven dried at a temperature of 100⁰c to 110⁰c for 24 hours and then subjected to mechanical analysis.

The mechanical analysis of the samples involved the determination of particle size distribution for coarse grained soil *i.e.* grains coarser than 0.063mm using a set of sieves. The sieves were carefully cleaned and their corresponding weight recorded. The collecting pan was placed at the bottom and the sieves arranged in order of increasing mesh size with the finest at the bottom and the coarsest at the top. The oven dried soil was then poured into the sieve from the top, covered with a lid and the whole stack taken to the sieve shaker which shook for 10 minutes

thereafter the stack of sieves was removed from the sieve shaker and the amount retained on each sieve was weighed and the weight recorded. The percentage on each sieve was then computed by dividing the weight retained on each sieve with the original soil weight.

Laboratory Analysis

Inorganic geochemical analysis which involved digestion and determination of major and trace element abundances using the Atomic Absorption Spectrophotometry (AAS) method was carried out on the sediments.

III. RESULTS AND DISCUSSIONS

Lithologic Description

The sediments of the Epe lagoon were mainly fine to medium grained clayey-sands ranging from light to dark grey in colour with some organic debris as well as some fossil shells. Although, some very fine clay materials with preponderant organic debris were also found (Table 2). Majority of the sediments plotted within the clayey-sand region very close to the sand axis indicating a dominance of sandy materials within the sediments (Fig. 2), as confirmed by the high SiO₂ values (Table 3). The presence of organic debris within the sediments (Table 2) can be attributed to decay of both dead plant and animal matter which may have been transported from adjoining rivers, deposited within the lagoon and accumulated within the sediments.

Geochemical Analysis

Whole rock geochemistry of the study area is discussed mainly under major elements through the use of various ternary and bivariate plots which help in classification as well as deducing the depositional environment and weathering history of the sediments of the Epe lagoon.

Major Element Geochemistry

The abundance of major elements in the sediments of the Epe lagoon as well as their tabulated range of values and calculated mean values as obtained from the major element geochemical analysis are listed in Table's 3, 4, 5 and 6. The dominance of SiO₂ over all other oxides is a confirmation of the greater proportions of sandy materials within the sediments. The sediments have average SiO₂ values higher than the average

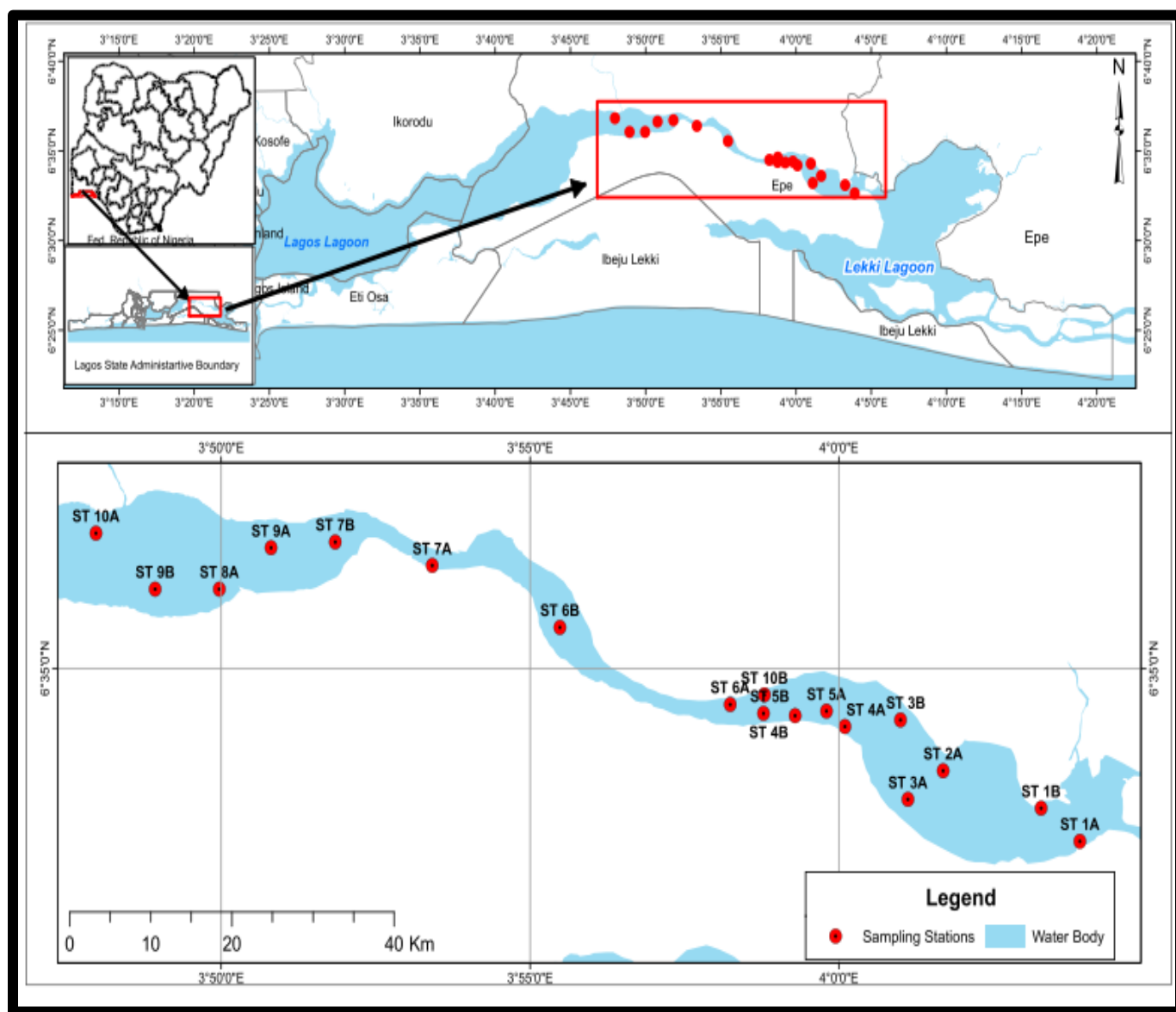


Fig. 1: Map of study area showing sample locations

TABLE 1: GPS Coordinates of sample points and locations within the study area

S/N	SAMPLE NUMBER	LOCATION	LATITUDE	LONGITUDE
1	Epe/st/1b	Epe lagoon	6°33'27.74°N	4°03'21.51°E
2	Epe/st/10a	Epe lagoon	6°37'00.10°N	3°46'32.02°E
3	Epe/st/7a	Epe lagoon	6°36'38.63°N	3°51'50.86°E
4	Epe/st/4b	Epe lagoon	6°34'18.29°N	3°59'45.66°E
5	Epe/st/1a	Epe lagoon	6°32'39.88°N	4°03'54.85°E
6	Epe/st/4a	Epe lagoon	6°34'49.34°N	3°59'23.79°E
7	Epe/st/8b	Epe lagoon	6°36'02.41°N	3°49'17.74°E
8	Epe/st/3a	Epe lagoon	6°33'15.64°N	4°01'07.48°E
9	Epe/st/5b	Epe lagoon	6°34'32.26°N	3°58'17.06°E

10	Epe/st/7b	Epe lagoon	6°36'04.42°N	3°49'57.97°E
11	Epe/st/3b	Epe lagoon	6°34'11.56°N	4°00'06.70°E
12	Epe/st/9a	Epe lagoon	6°36'48.28°N	3°47'58.18°E
13	Epe/st/9b	Epe lagoon	6°36'40.01°N	3°48'38.14°E
14	Epe/st/5a	Epe lagoon	6°34'18.59°N	3°58'47.62°E
15	Epe/st/2a	Epe lagoon	6°33'39.23°N	4°01'43.03°E
16	Epe/st/8a	Epe lagoon	6°36'36.01°N	3°50'48.30°E
17	Epe/st/6a	Epe lagoon	6°35'36.11°N	3°55'27.72°E
18	Epe/st/6b	Epe lagoon	6°36'19.03°N	3°53'32.66°E
19	Epe/st/10b	Epe lagoon	6°34'33.45°N	3°58'42.63°E

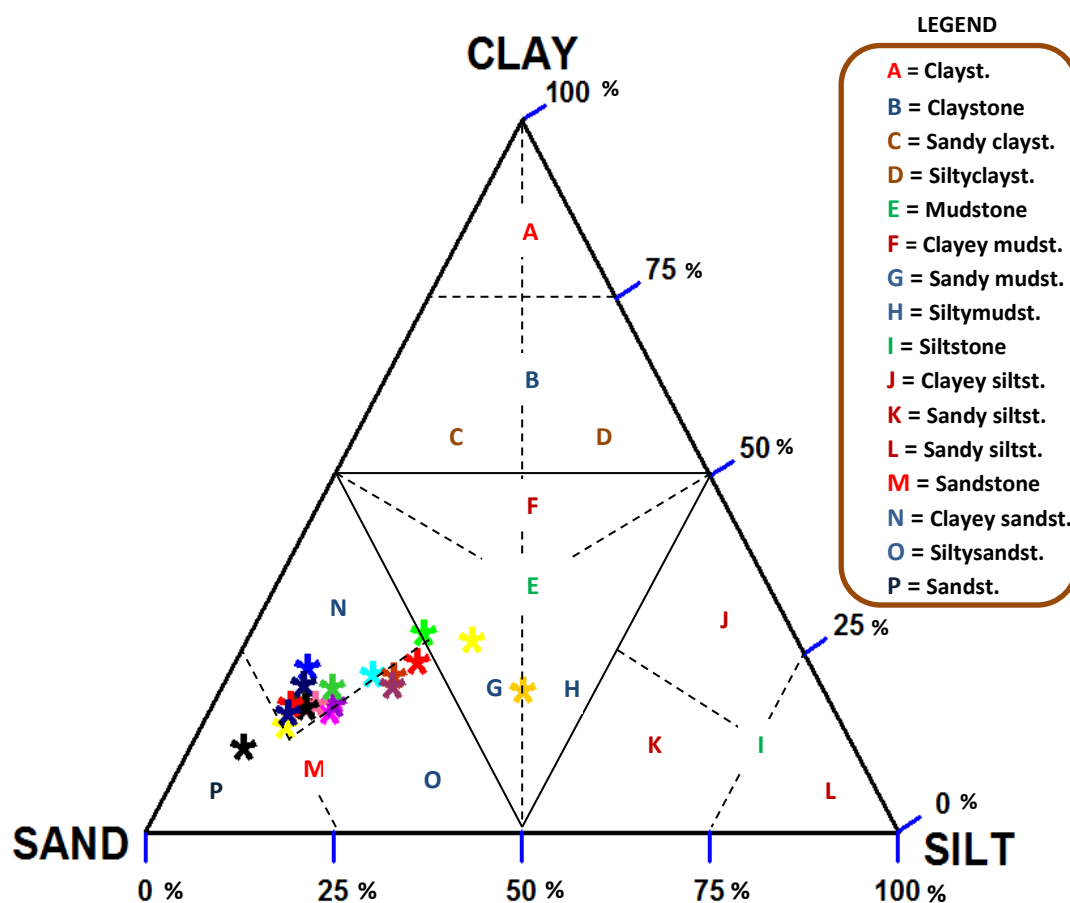


Fig. 2: Ternary plot showing Lithologic distribution of sediments

Table 2: Lithologic Description of the Epe Lagoon Sediments

S/N	SAMPLE NUMBER	FORMATION	LITHOLOGY	TEXTURE	COLOUR	ORGANIC MATTER
1	Epe/st/1b	Coastal-plain sands	Clayey-sand	Fine grain	Dark Grey	Presence of fossil shells
2	Epe/st/10a	Coastal-plain sands	Clayey-sand	Fine	Grey	No visible organic material found
3	Epe/st/7a	Coastal-plain sands	Clayey-sand	Fine-medium	Grey	Presence of fossil shells
4	Epe/st/4b	Coastal-plain sands	Clayey-sand	Fine	Light grey	Presence of fossil shells
5	Epe/st/1a	Coastal-plain sands	Clayey-sand	Fine	Dark grey	Presence of fossil shells
6	Epe/st/4a	Coastal-plain sands	Clayey-sand	Fine	Dark grey	Presence of fossil shells and organic debris
7	Epe/st/8b	Coastal-plain sands	Clayey-sand	Fine	Light	Presence of fossil shells
8	Epe/st/3a	Coastal-plain sands	Clayey-sand	Fine-medium	Grey	Presence of fossil shells
9	Epe/st/5b	Coastal-plain sands	Clayey-sand	Fine-medium	Grey	presence of fossil shells
10	Epe/st/7b	Coastal-plain sands	Clayey-sand	Fine-medium	Grey	Presence of fossil shells
11	Epe/st/3b	Coastal-plain sands	Clayey-sand	Fine-medium	Grey	Presence of fossil shells
12	Epe/st/9a	Coastal-plain sands	Clayey-sand	Fine-medium	Dark grey	Presence of fossil shells and organic debris
13	Epe/st/9b	Coastal-plain sands	Clayey-sand	Fine	Dark grey	Presence of fossil shells and organic debris
14	Epe/st/5a	Coastal-plain sands	Clayey-sand	Fine-medium	Grey	Presence of fossil shells
15	Epe/st/2a	Coastal-plain sands	Clayey-sand	Fine	Light grey	No visible organic material found
16	Epe/st/8a	Coastal-plain sands	Clayey-sand	Fine	Grey	Presence of fossil shells
17	Epe/st/6a	Coastal-plain sands	Clayey-sand	Fine	Light grey	Presence of organic debris
18	Epe/st/6b	Coastal-plain sands	Clayey-sand	Fine	Grey	Presence of organic debris
19	Epe/st/10b	Coastal-plain sands	Clayey-sand	Fine	Light grey	Presence of organic debris

of 66% by weight of the upper continental crust (Taylor and McLennan, 1985). Corresponding high average value of $Al_2O_3/(Na_2O+CaO)$ recorded as 138.099 (Table 5) which is a ratio of the most immobile element to the most mobile element confirms the high content of clastic materials over carbonate materials within the sediment. Also, $Al_2O_3/(Fe_2O_3+MgO)$ values which ranged from (1.65-29.54) (Table 5) indicates relatively moderate to high diagenetic alteration. Furthermore, K_2O/Al_2O_3 values were found to range from 0.002-0.4973 suggesting sedimentary recycling or increase in the degree of source area weathering (Bauluz *et.al*, 2000); it also indicates weathering

and alteration (Table 6). Very low average values of Al_2O_3/SiO_2 recorded as 0.08932 is an indication of quartz enrichment (Bhatia, 1983), this supports the relatively high average value of SiO_2/Al_2O_3 recorded as 64.33 (Table 5) which also confirms the high SiO_2 values previously observed within the sediments. Values of Al_2O_3 , Fe_2O_3 and MgO were observed to decrease with increasing SiO_2 , while others such as CaO , Na_2O , K_2O , TiO_2 , P_2O_5 and MnO had very low concentrations (Table 3). More so, TiO_2/Al_2O_3 values ranged from 0.004-0.8604, indicating that the sediments are chemically matured (Jenner *et.al*, 1981). The correlation results of TiO_2 against Al_2O_3 and Fe_2O_3 with Al_2O_3

were positive ($r = 0.98$ and 0.60), while Na_2O with Al_2O_3 was negative ($r = 0.24$) suggesting partial association with phyllosilicates (Fig. 3). The correlation results with K_2O were also positive ($r = 0.22$, 0.14 and 0.74) suggesting the input of aluminosilicates and feldspar within the sediments. The positive correlation between K_2O and Al_2O_3 ($r = 0.19$) (Fig. 3) implies that the concentration of potassium bearing minerals have significant influence on the distribution of aluminum (Al) and thus suggest that the abundance of these elements is controlled by the content of clay minerals (McLennan *et. al*, 1983).

Table 3: Geochemical Composition of Epe lagoon sediments (Major elements expressed in percentage)

S/N	SAMPLE	LOCATION	SiO_2	Al_2O_3	Fe_2O_3	MgO	CaO	Na_2O	K_2O	TiO_2	P_2O_5	MnO	Total
1	Epe/st/1b	Epe lagoon	60.9	30.3	1.43	0.07	0.04	0.02	0.33	3.42	0.08	0.01	96.6
2	Epe/st/10a	Epe lagoon	81.73	11.52	0.35	0.04	0.03	0.03	0.16	1.25	0.05	0.01	95.17
3	Epe/st/7a	Epe lagoon	52.99	35.46	1.55	0.05	0.02	0.01	0.17	4.56	0.1	0.01	94.92
4	Epe/st/4b	Epe lagoon	97.31	1.07	0.25	0.02	0.03	0.04	0.04	0.26	0.02	0.01	99.0
5	Epe/st/1a	Epe lagoon	98.37	0.71	0.04	0.01	0.03	0.04	0.04	0.06	0.01	0.01	99.32
6	Epe/st/4a	Epe lagoon	92.86	3.56	0.8	0.02	0.01	0.02	0.03	0.32	0.01	0.01	97.64
7	Epe/st/8b	Epe lagoon	80.19	10.17	2.96	0.03	0.02	0.01	0.03	0.9	0.06	0.01	94.38
8	Epe/st/3a	Epe lagoon	96.87	0.86	0.51	0.01	0.01	0.01	0.06	0.05	0.01	0.01	98.40
9	Epe/st/5b	Epe lagoon	98.87	0.71	0.04	0.01	0.01	0.03	0.04	0.05	0.01	0.01	99.78
10	Epe/st/7b	Epe lagoon	94.35	2.56	0.05	0.05	0.06	0.01	0.05	0.84	0.02	0.01	98.0
11	Epe/st/3b	Epe lagoon	97.46	0.86	0.34	0.04	0.05	0.01	0.04	0.74	0.01	0.01	99.56
12	Epe/st/9a	Epe lagoon	96.65	1.7	0.29	0.01	0.02	0.03	0.08	0.12	0.01	0.01	98.92
13	Epe/st/9b	Epe lagoon	97.62	0.87	0.11	0.01	0.01	0.02	0.04	0.12	0.01	0.01	98.82
14	Epe/st/5a	Epe lagoon	97.77	0.87	0.1	0.01	0.02	0.01	0.04	0.12	0.01	0.01	98.96
15	Epe/st/2a	Epe lagoon	97.30	1.05	0.22	0.02	0.03	0.03	0.04	0.25	0.01	0.01	98.96
16	Epe/st/8a	Epe lagoon	96.40	1.75	0.27	0.04	0.30	0.03	0.09	0.43	0.06	0.45	99.82
17	Epe/st/6a	Epe lagoon	93.36	1.83	0.67	0.04	0.42	0.08	0.91	0.58	0.21	0.08	98.18
18	Epe/st/6b	Epe lagoon	94.97	2.41	0.40	0.11	0.07	0.02	0.05	0.01	0.22	0.04	98.30
19	Epe/st/10b	Epe lagoon	92.70	4.20	0.59	0.05	0.07	0.05	0.30	0.35	0.33	0.07	98.71

Table 4: Summary of major oxide composition of the Epe-lagoon sediments

Oxides	Range (%)	Mean (N=19)	Standard Error	Standard Deviation	Variance
SiO_2	52.99-98.87	90.46	2.97	12.93	167.20
Al_2O_3	0.71-35.46	5.92	2.30	10.01	100.26
Fe_2O_3	0.04-2.96	0.58	0.16	0.72	0.52
MgO	0.01-0.11	0.03	0.006	0.03	0.001
CaO	0.01-0.42	0.099	0.04	0.18	0.033
Na_2O	0.01-0.08	0.026	0.004	0.02	0.000
K_2O	0.03-0.91	0.13	0.048	0.20	0.043
TiO_2	0.01-4.56	0.76	0.28	1.20	1.45
P_2O_5	0.01-0.33	0.07	0.021	0.09	0.008
MnO	0.01-0.45	0.04	0.23	0.10	0.010

Table 5: Major element ratios for sediments of the Epe lagoon, Eastern Dahomey basin, Nigeria

S/NO	SAMPLE	SiO ₂ /Al ₂ O ₃	Fe ₂ O ₃ /K ₂ O	Na ₂ O+CaO	Al ₂ O ₃ /(Na ₂ O+CaO)	Fe ₂ O ₃ +MgO	Al ₂ O ₃ /(Fe ₂ O ₃ +MgO)
1	Epe/st/1b	2.01	4.33	0.06	505	1.5	20.2
2	Epe/st/10a	7.10	2.19	0.06	192	0.39	29.54
3	Epe/st/7a	1.49	9.12	0.03	1182	1.6	22.16
4	Epe/st/4b	95.40	6.25	0.07	14.57	0.27	3.78
5	Epe/st/1a	138.55	1	0.07	10.14	0.05	14.2
6	Epe/st/4a	26.08	26.67	0.03	118.67	0.82	4.34
7	Epe/st/8b	7.89	98.67	0.03	339	2.99	3.40
8	Epe/st/3a	112.64	8.5	0.02	43	0.52	1.65
9	Epe/st/5b	139.25	1	0.04	17.75	0.05	14.2
10	Epe/st/7b	36.86	1	0.07	36.57	0.1	25.6
11	Epe/st/3b	113.33	8.5	0.06	14.33	0.38	2.26
12	Epe/st/9a	56.86	3.63	0.05	34	0.3	5.67
13	Epe/st/9b	112.21	2.75	0.03	29	0.12	7.25
14	Epe/st/5a	112.38	2.5	0.03	29	0.11	7.91
15	Epe/st/2a	92.67	5.5	0.06	17.5	0.24	4.38
16	Epe/st/8a	55.09	3	0.33	5.30	0.31	5.65
17	Epe/st/6a	51.02	0.74	0.5	3.66	0.71	2.58
18	Epe/st/6b	39.41	8	0.09	26.78	0.51	4.73
19	Epe/st/10b	22.07	1.97	0.75	5.6	0.64	6.56

Table 6: Major element ratios for Epe lagoon sediments

S/NO	SAMPLE	K ₂ O/Al ₂ O ₃	TiO ₂ /Al ₂ O ₃	K ₂ O+CaO+MgO	Fe ₂ O ₃ +MgO	Al ₂ O ₃ /SiO ₂
1	Epe/st/1b	0.011	0.11	0.44	1.5	0.498
2	Epe/st/10a	0.014	0.11	0.23	0.39	0.141
3	Epe/st/7a	0.005	0.13	0.24	1.6	0.669
4	Epe/st/4b	0.039	0.25	0.09	0.27	0.010
5	Epe/st/1a	0.056	0.08	0.08	0.05	0.007
6	Epe/st/4a	0.008	0.09	0.06	0.82	0.038
7	Epe/st/8b	0.003	0.09	0.08	2.99	0.127
8	Epe/st/3a	0.070	0.06	0.08	0.52	0.009
9	Epe/st/5b	0.056	0.07	0.06	0.05	0.007
10	Epe/st/7b	0.020	0.33	0.16	0.1	0.027
11	Epe/st/3b	0.047	0.86	0.13	0.38	0.009
12	Epe/st/9a	0.047	0.07	0.11	0.3	0.018
13	Epe/st/9b	0.046	0.14	0.06	0.12	0.009
14	Epe/st/5a	0.046	0.14	0.07	0.11	0.009
15	Epe/st/2a	0.038	0.24	0.09	0.24	0.011
16	Epe/st/8a	0.051	0.25	0.43	0.31	0.018
17	Epe/st/6a	0.050	0.32	1.37	0.71	0.020
18	Epe/st/6b	0.021	0.004	0.23	0.51	0.025
19	Epe/st/10b	0.071	0.08	1.05	0.64	0.045

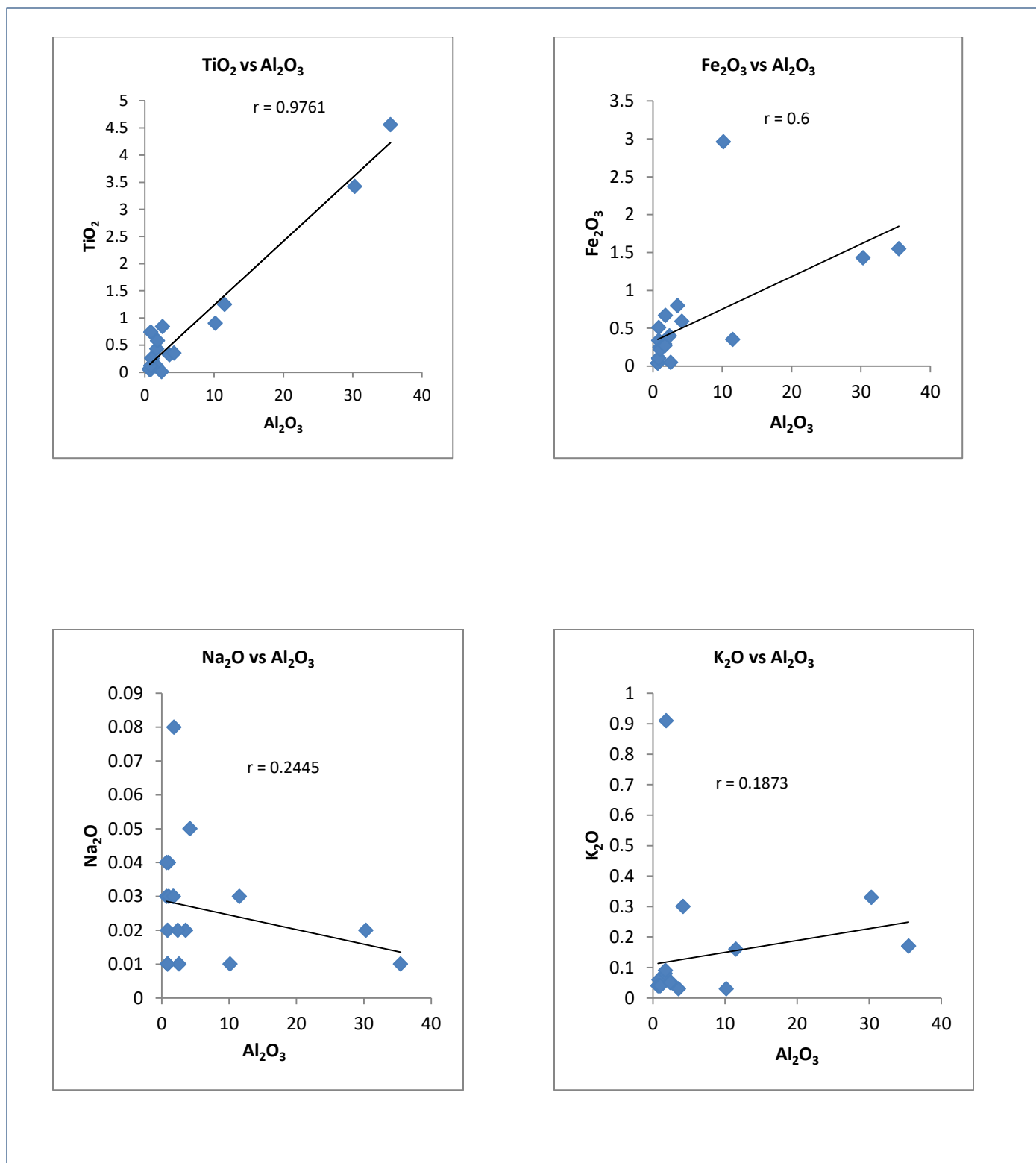


Fig. 3: Correlation plot of major oxides

Classification of the Epe lagoon sediments

Standard plot of Herron (1988) using $\log (\text{Fe}_2\text{O}_3/\text{K}_2\text{O})$ against $\log (\text{SiO}_2/\text{Al}_2\text{O}_3)$ which is the modified version of Pettijohn *et.al*, (1972) was employed to classify the sediments of the Epe lagoon. However, it was observed that about one-third of the sediments plotted within the quartz arenite region which is an indication of quartz enrichment and a confirmation of the high SiO_2 values recorded from the major element geochemistry results (Table 3). However, a notable fraction of the sediments also plotted within the Sublitharenite and Subarkose region indicating a change in the level of sediment maturity. Minor fractions of the Epe lagoon sediments also plotted randomly within the Fe-shale, Fe-sand, shale and wacke regions (Fig. 4).

Paleo-Weathering Indices and Maturity

The Chemical Index of Alteration (CIA) and the Plagioclase Index of Alteration (PIA) are the most widely used indices for quantitative estimation of the degree of chemical weathering undergone by rocks of the provenance area of clastic sediments (Fedo *et.al*, 1995). CIA records the alkali, and alkali earth elements (Nesbitt and Young, 1982); this is because alteration of igneous rocks during weathering results in depletion of alkali and alkaline earth elements and preferential enrichment of Al_2O_3 in sediments. Some other indices which may also be used include Chemical Indices of Weathering (CIW) (Harnois, 1988). Summary of CIA, PIA and PIW values are shown in Table 7.

$$\text{CIA} = 100\{ \text{Al}_2\text{O}_3 / (\text{Al}_2\text{O}_3 + \text{CaO} + \text{Na}_2\text{O} + \text{K}_2\text{O}) \}$$

$$\text{PIA} = 100\{ (\text{Al}_2\text{O}_3 - \text{K}_2\text{O}) / (\text{Al}_2\text{O}_3 + \text{CaO} + \text{Na}_2\text{O} - \text{K}_2\text{O}) \}$$

$$\text{CIW} = 100\{ \text{Al}_2\text{O}_3 / (\text{Al}_2\text{O}_3 + \text{CaO} + \text{Na}_2\text{O}) \}$$

Nesbitt and young (1982) suggested that high values (76-100) indicate intensive chemical weathering at the source areas while

low values (50 or less) indicate unweathered source areas. Furthermore, Osae *et.al*, (2006) also suggested that high CIA and PIA values (75-100) indicate intensive weathering in the source area whereas low values (<60) indicates low weathering in source area. Generally, the sediments of the Epe lagoon has CIA, PIA and CIW values ranging from (56.48-99.44, 64.79-99.92, and 78.54-99.92) with an average of (90.44, 93.53, and 94.47) respectively (Table 7). This results indicates high weathering at source areas as suggested by Nesbitt and young (1982) and Osae *et.al*, (2006). Also the CIA values were plotted in the (A-CN-K) diagram (Fig. 5), in the diagram the sediments plotted mostly within the kaolinite-chlorite and illite region. The fact that the samples did not plot very close to the plagioclase and K-feldspar line confirms also that the Epe lagoon sediments have undergone high (intensive) weathering at the source area.

Depositional Environment of the Epe lagoon sediments

The depositional environment of the Epe lagoon sediments was obtained using the ternary plots of Englund and Jorgensen (1973) which was done on the basis of the (Al_2O_3) - $(\text{K}_2\text{O} + \text{Na}_2\text{O} + \text{CaO})$ - $(\text{Fe}_2\text{O}_3 + \text{MgO})$ contents. As observed from the plots, most of the sediments deposited within the continental zone and are argillaceous (Fig. 6). However, there is a fairly gradual transition of some of the sediments from the continental environments into the transitional environment. More so, some of the sediments within the continental zone also plotted within the region of ferruginous argillites.

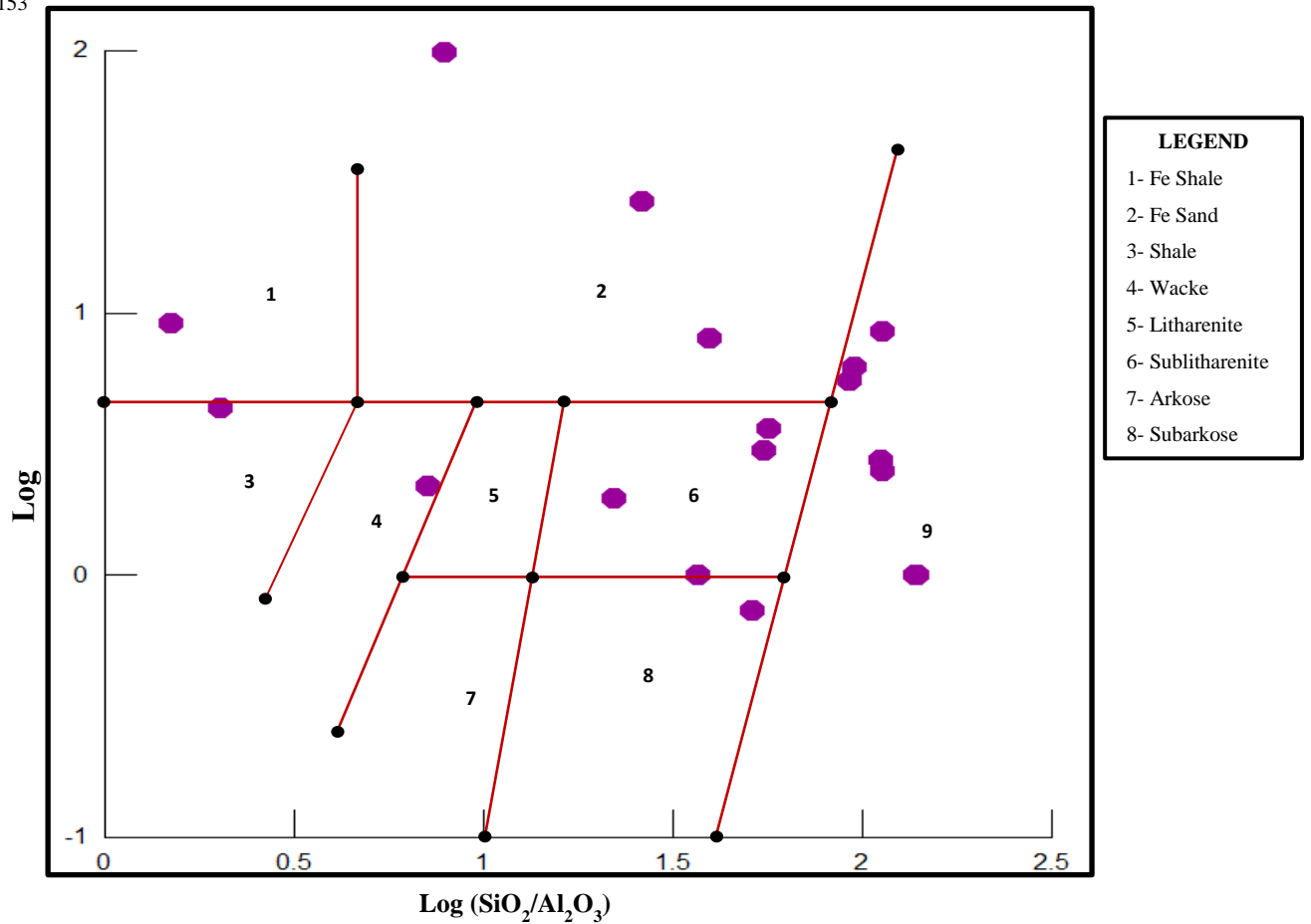


Fig. 4: Classification of Epe lagoon sediments (modified after Herron, 1988)

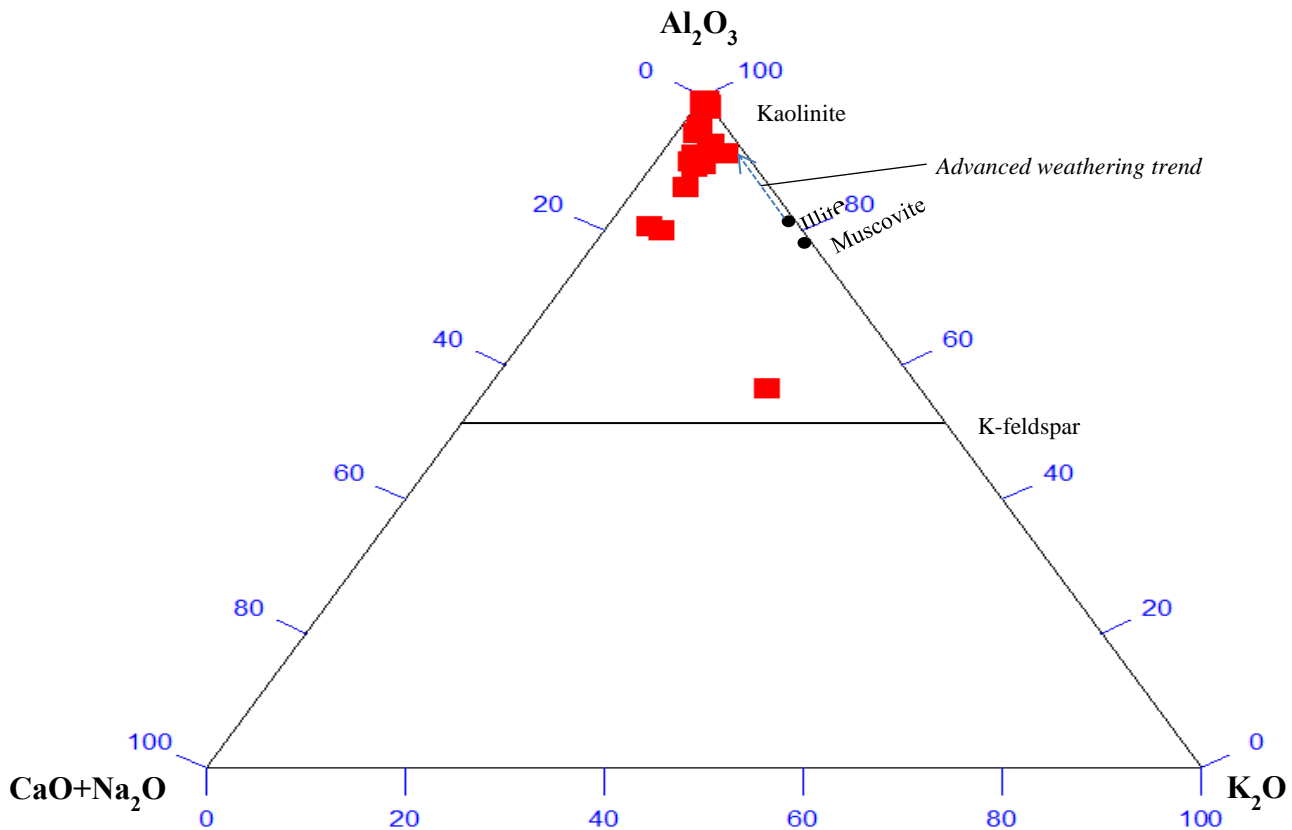


Fig. 5: CIA ternary diagram (Al_2O_3)-(CaO+Na₂O)-(K₂O) for the Epe lagoon sediments (after Nesbitt and young, 1982)

Table 7: CIA, PIA and CIW values for sediments of the Epe lagoon

S/N	SAMPLE NUMBER	CIA	PIA	CIW
1	Epe/st/1b	98.73	99.80	99.80
2	Epe/st/10a	98.13	99.47	99.48
3	Epe/st/7a	99.44	99.92	99.92
4	Epe/st/4b	90.27	93.33	93.58
5	Epe/st/1a	86.59	90.54	91.03
6	Epe/st/4a	98.34	99.16	99.16
7	Epe/st/8b	99.41	99.71	99.71
8	Epe/st/3a	91.49	97.56	97.73
9	Epe/st/5b	89.87	94.37	94.67
10	Epe/st/7b	95.52	97.29	97.34
11	Epe/st/3b	89.58	93.18	93.48
12	Epe/st/9a	92.90	97.01	97.14
13	Epe/st/9b	92.55	96.51	96.67
14	Epe/st/5a	92.55	96.51	96.67
15	Epe/st/2a	91.30	94.39	94.59
16	Epe/st/8a	80.65	83.42	84.13
17	Epe/st/6a	56.48	64.79	78.54
18	Epe/st/6b	94.51	96.33	96.40
19	Epe/st/10b	80.0	83.87	84.85

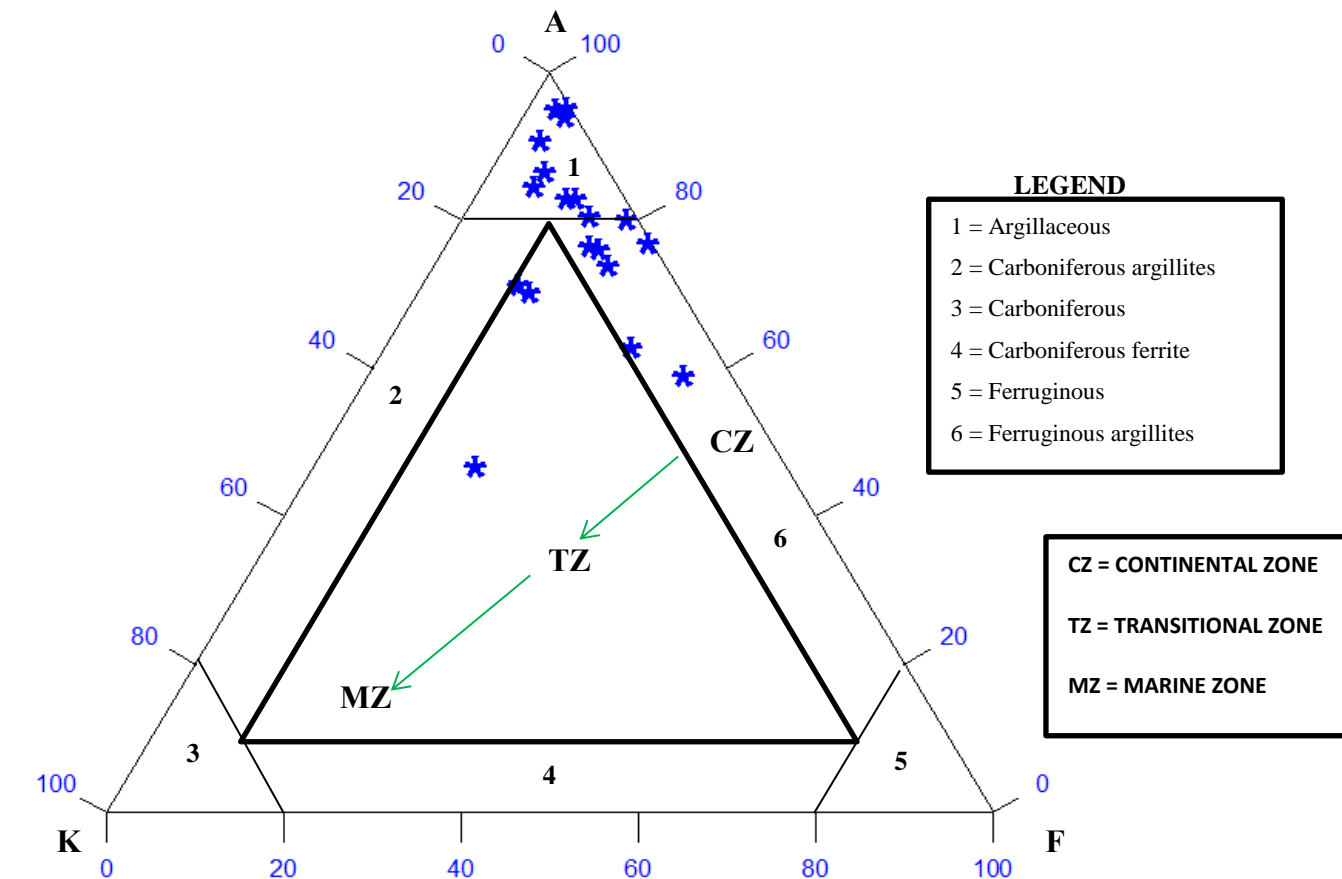


Fig. 6: $\text{Al}_2\text{O}_3\text{-(K}_2\text{O+CaO+MgO)-(Fe}_2\text{O}_3\text{+MgO)}$ {AKF} Ternary plot for Epe lagoon sediments (after Englung and Jorgensen, 1973)

IV CONCLUSION

The inorganic geochemical analysis of the Epe lagoon revealed that SiO_2 is the dominant oxide, followed by Al_2O_3 and Fe_2O_3 which constitutes over 90% of the total oxides. Others are MgO , CaO , Na_2O , K_2O , TiO_2 and P_2O_5 . The $\text{SiO}_2/\text{Al}_2\text{O}_3$ ratios for the sediments were appreciably high indicating that the sediments have been heavily weathered and transported; this is also corroborated by the enrichment of quartz and depletion feldspar in most of the samples. The fairly high $\text{K}_2\text{O}/\text{Na}_2\text{O}$ ratios may be attributed to the presence of plagioclase within the sediments. From the major element geochemistry, the application of source area weathering using CIA, CIW and PIA with mean values of 90.44, 93.53 and 94.47 respectively indicates that the sediments of the Epe lagoon have been subjected to intense weathering at the source area as well as high levels of chemical weathering and transportation. In the A-CN-K diagram the sediments of the Epe lagoon plotted mostly within the kaolinite-chlorite and illite region; also an indication of high weathering at the source.

The use of heron's model classified the sediments as mainly quartz arenite. However, a notable fraction of the sediments were also as Sublitharenite and Subarkose with very minor fractions classified as Fe-shale, Fe-sand and Wacke. AKF ternary plots revealed that the sediments were deposited mainly in continental environment and dominated by argillaceous materials with some ferruginous argillites, although some fractions of the sediments were also found to have been transported from the continental environment to the transition zone.

V. APPENDIX

INORGANIC GEOCHEMISTRY

Determination of Major element abundances using the Atomic Absorption Spectrophotometry (AAS) Method

Atomic absorption spectrophotometry is based upon the observation that atoms of an element can absorb electromagnetic radiation. This occurs when the element is atomized and the wavelength of light absorbed is specific to each element. Thus, the atomic absorption spectrophotometer comprises an atomizing device, during the atomization of a sample in a beam of light, as a consequence of atomic absorption, can be calibrated and is

sensitive at the ppm level. Each sample was first pulverized and 0.5gram digested with a mixture of prepared solution of Nitric acid and Hydrochloric acid (ratio 1:3); also known as Aqua regia solution. The sample was stirred and heated inside a fume cupboard. The digested sample was diluted with distilled water and made up to 20ml mark. About 1ml was taken from the solution and further diluted with distilled water to 10ml mark, which represents the stock solution, x 10-dilution factor. The major elements Si, Al, Fe, Mn, Mg, Ca, Na, K and Ti were then determined from the digested samples using a Perkin Elmer 400 atomic absorption spectrophotometer. These elements were further converted from ppm level to their respective oxides in percentage. The AAS is a widely used technique and comparatively cheap. However, there are two limitations to the routine use of AAS in silicate analysis. Firstly, the sample must be prepared as a solution, and secondly it is element-specific, i.e. one element can usually be analyzed at a time, although this latter limitation has in part been overcome by fitting instruments with multiple-turret lamp holders.

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AUTHORS

First Author – OMOREGBE Osagie Aret, OND, HND (Petroleum Training Institute), B.Sc., (University of Ibadan), M.Sc., (University of Benin) Pg Dip (Laser Petroleum Geoscience Centre). Omoregbeosagie88@gmail.com

Second Author – OLOYEDE, David Abiodun, B.Sc., (University of Ilorin), M.Sc., (University of Benin)

Third Author – EKUERUGBE, Lucky Orezimena, B.Sc., (Federal University of Petroleum Resources), M.Sc., (University of Benin)

Fourth Author – NKIM, Akere Godson, B.Sc., (University of Port Harcourt), M.Sc., (University of Port Harcourt)

Correspondence Author – OMOREGBE Osagie Aret, Omoregbeosagie88@gmail.com, 08138820324.