



Evaluation of seasonal variation of water quality using multivariate statistical analysis and irrigation parameter indices in Ajakanga area, Ibadan, Nigeria

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Abstract

The variation of groundwater quality across different regions is of great importance in the study of groundwater so as to ascertain the sources of contaminants to available water sources. Geochemical assessment of groundwater samples from hand-dug wells were done within the vicinity of Ajakanga dumpsite, Ibadan, Southwestern, Nigeria, with the aim of assessing their suitability for domestic and irrigation purposes. Ten groundwater samples were collected both in dry and wet seasons for analysis of physicochemical parameters such as: pH, EC, TDS, Na⁺, K⁺, Ca²⁺, Mg²⁺, HCO₃⁻, Cl⁻, SO₄²⁻, NO₃²⁻ principal component analysis (PCA) and cluster analysis (CA) were used to determine probable sources of groundwater contamination. The results of the analyses showed the groundwater samples to be within permissible limits of WHO/NSDWQ, while elevated values of concentrations of most analyzed chemical constituents in water samples were noticed in S₁ and S₁₀ due to their nearness to the dumpsite and agricultural overflow, respectively. Groundwater in the study area is of hard, fresh and alkaline nature. There are very strong associations between EC and TDS, HCO₃⁻ and CO₃²⁻ in both seasons. PCA identified five and three major factors accounting for 95.7 and 88.7% of total variation in water quality for dry and wet seasons, respectively. PCA also identified factors influencing water quality as those probably related to mineral dissolution, groundwater–rock interaction, weathering process and anthropogenic activities from the dumpsite. Results of CA show groups based on similar water quality characteristics and on the extent of proximity to the dumpsite. Assessment for irrigation purpose showed that most of the water samples were suitable for agricultural purpose except in a few locations.

Keywords Physicochemical · Dumpsite · Freshwater · Anthropogenic · PCA · CA

Introduction

The assessment of groundwater quality is as important as its quantity for various purposes ranging from domestic, industrial and agricultural uses all over the globe (Subramani and Damodarasamy 2005). The quality of groundwater in a particular region is a function of physical, chemical and biological parameters. The variation of groundwater quality in a particular area is a function of physical and chemical

parameters that are greatly influenced by geological formations and anthropogenic activities (Subramani and Damodarasamy 2005). Pollution of groundwater is a major threat posed by leachate which is formed by anaerobic decomposition of waste and may infiltrate the aquifer (Tesfaye 2007). Groundwater contamination has become a great problem due to rapid growth rate of population, industrialization and urbanization in the metropolitan city all over the world. The quality of groundwater is normally characterized by different physicochemical parameters level. These parameters change widely due to various types of pollution, seasonal variation and groundwater extraction (Ramakrishnaiah et al. 2009). Siting of open dumpsite near the residential areas can have undesirable effect on nearby water sources if the leachate emanated from decomposed solid waste penetrate and contaminate the water table. The use of polluted groundwater for drinking and consumption purposes can cause major health problem. According to WHO, about 80% of

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all diseases in human beings are caused by water (Ramakrishnaiah et al. 2009). Therefore, a periodic assessment of groundwater quality is necessary in order to ascertain the quality for human consumption purpose as well as to provide an overall scenario about the sources of groundwater contamination, thereby open an avenue for better planning for sustainable management of groundwater.

Hydrochemical study reveals the quality of water suitable for domestic and agricultural purpose. Further, it is possible to understand the change in quality due to rock–water interaction or any type of anthropogenic influence (Wilcox 1948). Several environmental researchers have identified contamination plumes from disposal sites (Matias et al. 1994; Ikem et al. 2002; Tijani et al. 2002) with most of these published studies focusing on defining the spatial extent of groundwater pollution based on geochemical analysis results. The suitability of groundwater resources for irrigation purpose was also studied (Sujatha and Reddy 2003; Sadashivaiah et al. 2008; Ramesh and Bhuvana 2012). Several published research studies have employed the use of multivariate statistical analysis in the interpretation of groundwater quality data obtained from various sources (Sundaray 2010; Singh et al., 2008; Uddamari et al. 2014; Oketola et al. 2013; Molla et al. 2015; Arslan 2013; Zhang et al. 2014; Majolagbe et al. 2016; Markic et al. 2015; Razmkhah et al. 2010). Scientists have also employed the use of principal component analysis (PCA) to study soil physicochemical properties and its geochemical constituents, identification of heavy metals pollutants in soil, analysis of heavy metals presence in dust and evaluation of influence of seasons on air pollution (Adhikari et al. 2003; Ma et al. 2016; Satyanarayanan et al. 2016; Gergen and Harmanescu 2012; Iwara et al. 2014; Lu et al. 2010; Burt et al. 2014; Benhaddya and Hadjel 2014; Abdul Raheem et al. 2008). However, the study of irrigation suitability of groundwater samples within dumpsite and their interpretation using multivariate statistical analysis has not been efficient.

The present study was carried out during dry and wet seasons from close by hand-dug wells neighboring Ajakanga solid waste disposal site for better understanding of spatial and seasonal variability of physicochemical parameters, hydro-geochemical facies of groundwater and identification of contamination sources that may affect the groundwater samples using multivariate statistical approach.

Site description and geological setting

Ibadan is located approximately within the squares of longitude $3^{\circ} 35^1-4^{\circ} 10^1$ east of the Greenwich meridian and latitude $7^{\circ} 20^1-7^{\circ} 40^1$ north of the equator. Solid wastes are dumped indiscriminately on open grounds and along road networks in so many places within Ibadan metropolis. There are several collection points from which refuse

are cleared by government trucks at regular intervals and deposited at the central dump sites managed by the government. The city generates about 1,618,293 kg of solid waste daily. There are four designated dumpsites in Ibadan namely: Aba-Eku, Ajakanga, Awotan and Lapite. For this study, the area is Ajakanga dumpsite in southwestern part of Ibadan. Ajakanga dumpsite lies between latitude of $3^{\circ} 50 187-3^{\circ} 50 696E$ and longitude $7^{\circ} 18 021-7^{\circ} 18 979N$. It was opened in 1998 and still in operation till date. The general overview of the dumpsite is shown in Fig. 1. The study area falls within the humid and sub humid tropical climate of southwestern Nigeria with a mean annual rainfall of about 1230 mm and mean maximum temperature of $32^{\circ}C$. The soil type of the study area belongs to Orthic Luvisol (FAO 2015). The mean value of the water retention capacity of the experimental soil within the dumpsite is 37.25%.

The geology of the area is a basement complex formation of southwestern Nigeria and are mainly the metamorphic rocks of Precambrian age with few intrusions of granites and porphyries of Jurassic age. The dominant rock types are: quartzite of meta-sedimentary series, banded gneiss, augen gneisses and migmatites which constitute the gneiss–migmatite complex. Other minor rock types include pegmatite, quartz, aplites, amphibolites and xenolith (Adeigbe and Oluwatoke 2009). Banded gneiss constitutes over 75% of the rocks in and around Ibadan while augen gneisses and quartzites share the remaining in about equal percentages (Adeigbe and Oluwatoke 2009). The basement complex rocks in their unchanged form are characterized by low porosity and permeability which determines the hydrogeological properties of the rocks depending on the grain size and mineralogy of the rocks. The topsoil has been disturbed due to dumping activities in the study area and hence, constitutes the waste dump and the leachate derived from its decomposition processing as shown in Fig. 2.

Materials and methodology

Collection of groundwater samples

Ten water samples were collected from nearby hand-dug wells within Ajakanga solid waste disposal site during the months of March and August, 2013 using 2-L polyethylene bottles. The distance of the hand-dug wells to the study area as well as the latitude and longitude of each sampling point was taken with the aid of Hand held Garmin Etrex GPS (Azim et al. 2011) is shown in Table 1. The groundwater flow direction of the sampling points is shown in Fig. 3. Groundwater flow divergent zones were around sampling points S6, S7 and S8 while the convergent zones were



Fig. 1 General overview of Ajakanga dumpsite

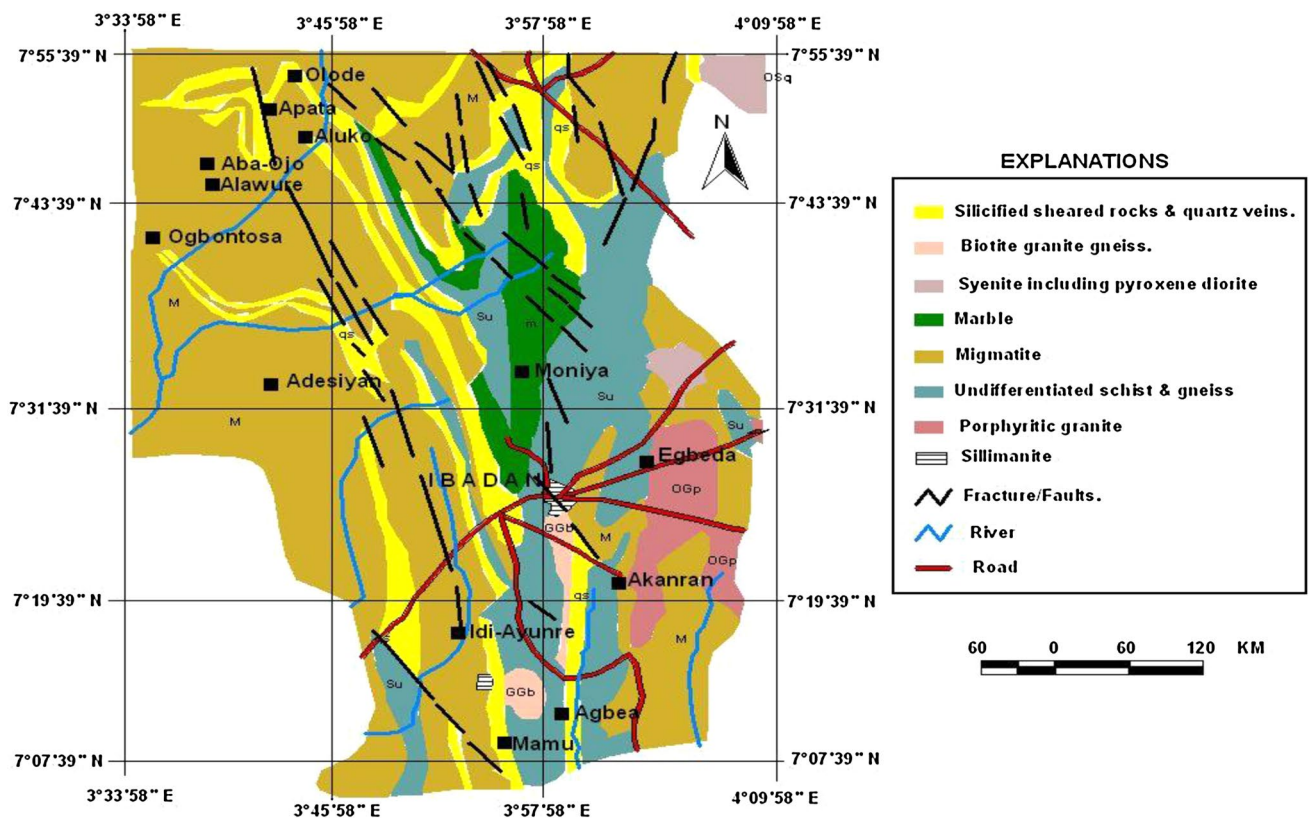


Fig. 2 Generalized geological map of Ibadan (JPEG)

around sampling points S1, S4 and S10. Samples S1, S2, and S10 were at downside of the dumpsite while samples S7 and S8 were at a distance of more than 250 m from the dumpsite.

Water samples were collected by lowering the bottle at depth of about one foot below the surface, rinsed the bottle three times with the water to be collected before the actual

Table 1 Water sampling location parameters during dry and wet seasons

Sample code	Depth to bottom (m)	Dry season depth to water level (m)	Wet season depth to water level (m)	Distance to landfill (m)	Coordinates
S ₁	9.1	3.7	2.7	90	7° 18' 40.53"N 3° 50' 32.22"E
S ₂	2.7	2.0	2.1	110	7° 18' 38.57"N 3° 50' 26.00"E
S ₃	3.5	3.5	3.2	100	7° 18' 44.22"N 3° 50' 32.71"E
S ₄	6.4	5.8	2.7	200	7° 18' 44.03"N 3° 50' 40.84"E
S ₅	5.5	5.2	2.7	220	7° 18' 47.81"N 3° 50' 41.94"E
S ₆	5.5	4.6	4.3	200	7° 18' 49.23"N 3° 50' 40.01"E
S ₇	5.8	5.5	3.2	270	7° 18' 56.15"N 3° 50' 45.07"E
S ₈	8.2	7.2	6.5	520	7° 18' 04.45"N 3° 50' 53.57"E
S ₉	–	–	–	120	7° 18' 49.46"N 3° 50' 53.57"E
S ₁₀	3.7	1.8	1.8	120	7° 18' 41.10"N 3° 50' 34.89"E

collection of the samples. After collection, the cap of each sampling bottle was screwed on tightly to avoid leakage (Asef Iqbal and Gupta 2009; Reza and Singh 2010; Odukoya and Abimbola 2010). A 0.45- μ m membrane filter was used to remove unwanted materials from collected water samples.

The collected water samples were transferred into 2-L sterilized polyethylene bottles and kept at 4 °C before chemical analyses at the laboratories. Water samples of approximately 125 mL were used for elemental analysis. Some of water sampling bottles for the analysis of cations and heavy metals were acidified with concentrated nitric acid to bring water acid solution to pH below 2 while the other un-acidified water samples were analyzed for anions concentration. Chemical analyses were carried out for major anions, cations and heavy metal concentrations using the standard procedure recommended by APHA (1998). The qualitative chemical analyses were carried out at the analytical laboratory of department of Environmental Management and Toxicology (EMT) and Central Biotechnology laboratory, both of Federal University of Agriculture, Abeokuta (FUNAAB), Ogun State, Nigeria. Total dissolved solids (TDS), electrical conductivity (EC) and pH were measured in situ with the aid of multi-purpose conductivity meter.

The samples were collected in both wet and dry seasons. Preservation of water samples and chemical analyses were carried out as using standard methods of APHA (20th edition, 1998). The groundwater sampling locations and dumpsite are depicted in Fig. 4. The predominant rock type in the study area is migmatite gneiss (as shown in Fig. 4). Sodium and potassium were determined using flame photometric

method while calcium and magnesium concentrations were analyzed using absorption mode of atomic absorption spectrometric (AAS) method. Sulphate and nitrate were analyzed by turbidimetric and UV spectrophotometric method, respectively, chloride, carbonate and bicarbonate by titration method while total hardness (TH) was determined by ethylene diamine tetra acetic acid (EDTA) titration method using Eriochrome black-T as an indicator.

Multivariate statistical analysis

Two different multivariate statistical analyses were used to analyze the groundwater geochemical data. These are principal component analysis (PCA) and cluster analysis (CA). PCA is a multivariate statistical procedure which is used to diminish the dimensionality of the original data set consisting of a large number of interrelated variables while still retaining the inherent dependencies existed in the data set (Jianqin et al. 2010). Cluster analysis (CA) is a statistical technique that classifies water samples quality parameters into cluster whereby samples/variables within a particular cluster are similar to each other, but dissimilar from other clusters (Zhang et al. 2014; Sundaray 2010). CA was performed based on agglomerative schedule using a combination of Ward's linkage method (Ward 1963) and squared Euclidean distances as a measure of similarity between samples and/or parameters (Zhang et al. 2014) while PCA extract factor with eigenvalue > 1 which explained more total variation in the data set. Only component (factor) with eigenvalue > 1 were retained and later subjected to varimax

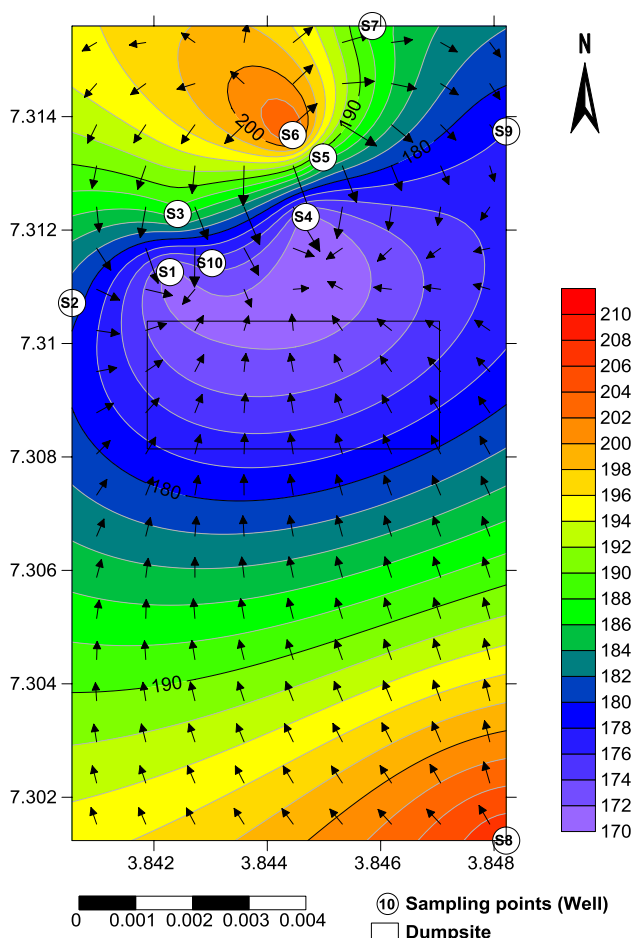


Fig. 3 Potentiometric map showing the groundwater flow direction with respect to dumpsite location

rotation (Kaiser 1958; Vega et al. 1998; Usman et al. 2014) before being used for interpretation.

Results and discussion

The results of water quality parameter analyses on collected water samples during dry and wet seasons sampling periods are presented in Table 2. The table shows the variation in the concentration level of analyzed parameters during wet and dry seasons.

Groundwater quality for drinking purposes

The pH values of groundwater samples during dry and wet seasons ranged from 6.9 to 7.8 and 6.7 to 7.3, respectively. The pH values for the two seasons lie within the permissible limit (Kamble and Saxena 2016; Chavan and Zambare 2014; Ariyo and Enikanoselu 2007). The total dissolved solids (TDS) concentrations during dry and wet seasons varied

from 88 to 299 mg/L and 95 to 351 mg/L, respectively. All TDS values lie below 500 mg/L specified by WHO (2007) and NSDWQ (2007) limit. Based on TDS results, all the analyzed water samples can be classified as freshwater since their TDS values is less than 1000 mg/L (Subramani and Damodarasamy 2005; Adebayo et al. 2015). Highest TDS value of 299 mg/L was noticed in S₁, 90 m away from the dumpsite during dry season. The result agrees with similar work by Adeolu et al. 2011. Electrical conductivity (EC) values ranged from 176 to 598 $\mu\text{s}/\text{cm}$ in dry season and from 191 to 705 $\mu\text{s}/\text{cm}$ during wet season. The EC values in both seasons lie within the standard limit of 1000 $\mu\text{s}/\text{cm}$ specified by WHO (2007) and NSDWQ (2007). The average concentration of total hardness (TH) varies from 46 to 406 mg/L and 116 to 432 mg/L during dry and wet sampling periods, respectively. Based on Sawyer and McCarthy (1967) classification for total hardness, 20% fall under “soft class”, 40% under “Hard class”, 30% under “moderate hard” class while the remaining 10% falls under “very hard” class during dry season. However, during the wet season, none of the samples falls under “soft” class of hardness, 10% falls under “moderate hard” class, 60% fall under “Hard” class while the remaining 30% fall under “Very Hard” Class. It was observed that in all the sampling locations, TH values were higher in wet than in dry season. The Cl^- concentration of water samples during dry and wet seasons ranged from 16 to 113 mg/L and 10 to 53 mg/L, respectively. The observed values Cl^- in both seasons were within the permissible limit of 250 mg/L.

The NO_3^- concentration in groundwater ranged from 1.5 to 15.9 mg/L during dry season and 0–3.9 mg/L during wet season. The concentration of NO_3^- in groundwater and surface water is normally low (Azim et al. 2011). The NO_3^- values for both seasons were found to be within the limit of 50 mg/L specified by WHO (2007). The low concentrations of NO_3^- in analyzed groundwater samples agree with similar studies by Chavan and Zambare (2014), Ariyo and Enikanoselu (2007) and Subramani and Damodarasamy (2005). The values of SO_4^{2-} in the groundwater samples ranged from 14.4 to 127.7 mg/L and 7.6 to 52.3 mg/L during dry and wet seasons, respectively. However, sulphate values in both seasons lie below 250 mg/L specified by WHO (2007) and NSDWQ (2007). For the anions (HCO_3^- and CO_3^{2-}), CO_3^{2-} concentrations in dry and wet seasons ranged from 60 to 288 mg/L and 60 to 300 mg/L; while HCO_3^- values ranged from 122 to 586 mg/L and 122 to 610 mg/L in dry and wet seasons, respectively.

The Ca^{2+} , and Mg^{2+} status level in analyzed water samples during dry and wet seasons ranged from 1.3 to 49.2 mg/L and 2.0 to 173.4 mg/L; 1.1 to 14.2 mg/L and 3.3 to 49.3 mg/L, respectively. Na^+ concentration value in groundwater samples ranged from 12 to 30 mg/L and 11 to 24 mg/L during dry and wet seasons, respectively.

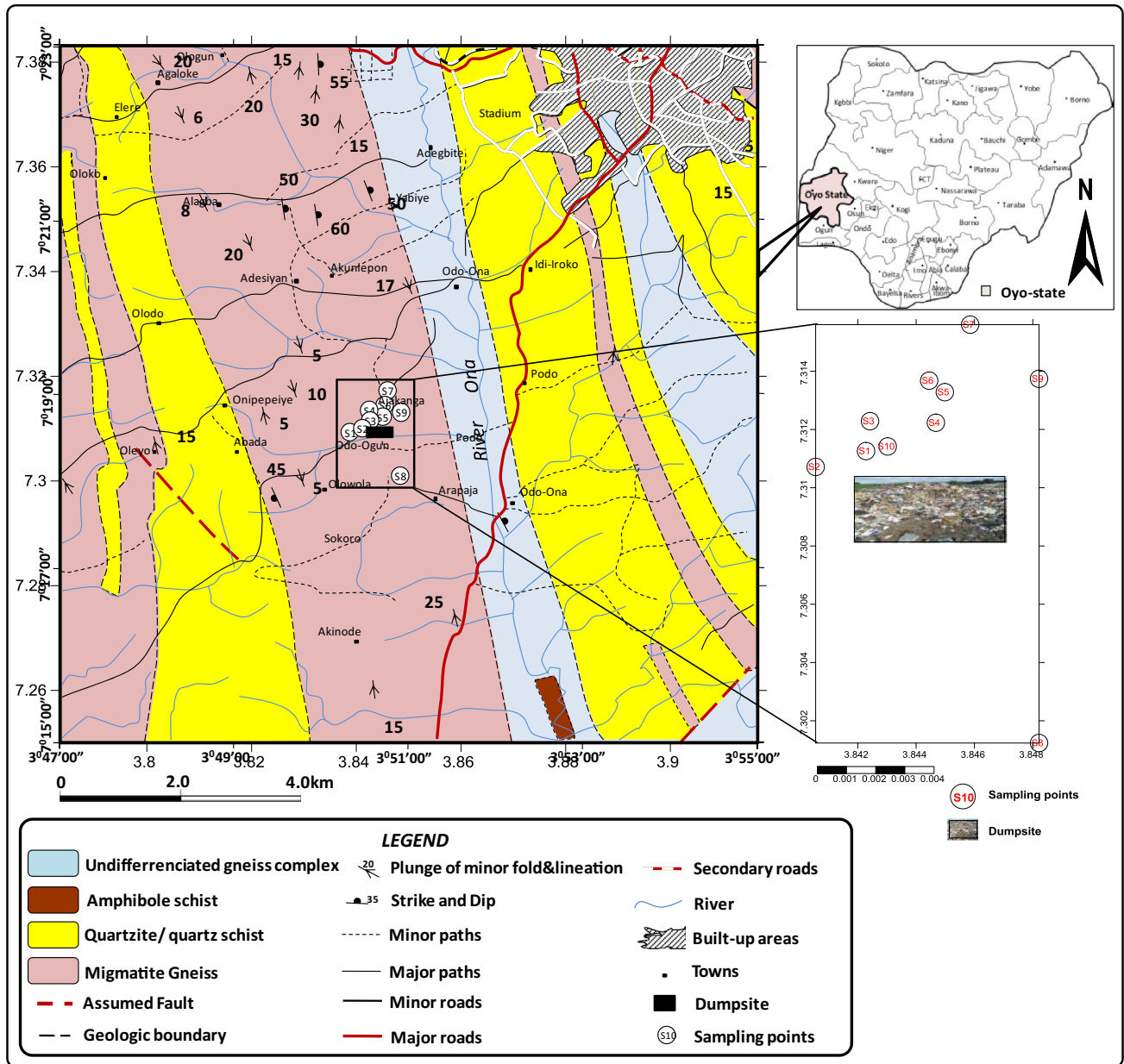


Fig. 4 Geological map showing the rock type that underlies the sampled area, dumpsite and layout of the sampling points (reproduced with permission from Jones and Hockey 1964)

There is no significant seasonal variations of K^+ . The values in groundwater samples ranged from 2 to 6 mg/L and 1 to 6 mg/L during dry and wet seasons, respectively (Kamble and Saxena 2016; Udayalaxmi et al. 2010; Odukoya and Abimbola 2010). The lowest and highest concentration of K^+ in groundwater may be due to the fact that most potassium-bearing minerals are resistant to decomposition by weathering processes and fairly low concentrations of ionic potassium in groundwater (Scheytt 1997; Sravanthi and Sudarshan 1998). However, higher concentration of some water quality parameters were noticed in Wells 1 and

10 which may be due to effect of leachate migration in the southern part of the dumpsite; nearness to dumpsite; agricultural run-off and fertilizer application on the nearby farm settlement.

Result of statistical analyses

Table 3 shows the details of the descriptive statistics of the analyzed water quality parameters from ten sampling points within the vicinity of the dumpsite. The degree of a linear association between any two of the analyzed variables

Table 2 Physicochemical parameters during dry and wet seasons for analyzed water samples

Sample	pH	EC (µs/cm)		TDS (mg/L)		Cl ⁻ (mg/L)		HCO ₃ ⁻ (mg/L)		CO ₃ ²⁻ (mg/L)		TH (mg/L)		Na ⁺ (mg/L)		K ⁺ (mg/L)		SO ₄ ²⁻ (mg/L)		NO ₃ ⁻ (mg/L)		Mg ²⁺ (mg/L)		Ca ²⁺ (mg/L)		
		Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry
S ₁	7.1	7.0	598	465	299	237	96	52	414.8	317.2	204	156	180	432	30	22	2	1	26.45	25.49	1.8	.1	13.88	18.02	24.01	20.76
S ₂	7.1	7.3	420	425	210	214	24	33.5	195.2	366	96	180	276	404	17	18	5	3	14.36	16.45	2.2	0	13.61	23.32	18.26	22.63
S ₃	7.4	6.9	367	377	184	185	24	13	292.8	414.8	144	204	178	350	13	17	1	1	19.68	7.58	1.5	0	12.69	20.07	8.12	14.27
S ₄	7.8	7.3	275	259	138	128	54	10.5	585.6	268.4	288	132	100	234	16	17	4	6	38.71	15.32	2.7	0	10.35	5.89	5.38	23.05
S ₅	7.4	7.1	176	205	88	101	16	10	219.6	219.6	108	108	46	180	18	11	2	1	57.74	18.23	10.2	2.9	1.12	3.29	.41	9.66
S ₆	7.2	6.7	530	411	264	205	113	52.5	170.8	219.6	84	108	200	260	18	24	6	1	127.74	27.74	11.9	3.2	12.97	14.69	7.28	4.51
S ₇	7.2	6.9	299	191	150	95	32	16.5	366	122	180	60	96	116	16	14	4	1	88.07	24.68	16.0	3.0	5.03	5.32	1.32	2.01
S ₈	7.6	7.0	225	242	112	121	28	15	122	170.8	60	84	70	206	15	16	2	1	45.81	14.20	5.1	.9	5.68	5.42	2.19	8.43
S ₉	7.3	7.2	273	251	137	125	26	13.5	268.4	219.6	132	108	118	210	12	15	3	2	21.77	14.20	5.8	3.9	11.63	9.64	9.02	10.88
S ₁₀	6.9	7.2	568	705	284	351	39	40.5	536.8	610	264	300	406	190	19	24	2	2	29.19	52.26	3.4	2.6	14.23	49.34	49.18	173.42

All parameters are in unit of mg/L except pH (no unit) and EC in µs/cm

Table 3 Descriptive statistics of the analyzed water quality for dry and wet season

	Dry		Wet		Analysis N
	Mean	Std. deviation	Mean	Std. deviation	
pH	7.3000	.26247	7.0600	.19551	10
EC	373.1000	149.56860	353.1000	158.21255	10
TDS	186.6000	74.62082	176.2000	79.46180	10
Cl ⁻	45.2000	33.13877	25.7000	17.24529	10
HCO ₃ ⁻	317.2000	156.02439	292.8000	142.27523	10
CO ₃ ²⁻	156.0000	76.73330	144.0000	69.97142	10
TH	167.0000	108.71063	258.2000	103.55653	10
Na ⁺	17.4000	4.94862	17.8000	4.31535	10
K ⁺	3.1000	1.59513	1.9000	1.59513	10
SO ₄ ²⁻	46.9520	35.86853	21.6150	12.40855	10
NO ₃ ⁻	6.0600	4.98045	1.6600	1.59457	10
Mg ²⁺	10.1190	4.55695	15.5000	13.80501	10
Ca ²⁺	12.5170	14.89881	28.9620	51.28535	10

measured by Pearson’s correlation coefficients for dry and wet seasons are presented in Tables 4, 5, respectively. There are very strong associations between EC and TDS, HCO₃⁻ and CO₃²⁻ during both seasons. Highly significant correlation between EC and TDS buttress the fact that EC depends largely on the quality of the dissolved salts present in water sample. There is negative correlation between Na⁺ and K⁺, TH and NO₃⁻ during dry and wet seasons. The negative correlations between TH and NO₃⁻ and between Na⁺ and K⁺ were expected because the effect of nitrogen-fixing bacteria decreases with increasing hardness of water (Fabiyyi 2008) while ion is normally less than and Na⁺ in igneous rock typical of basement complex formation (Scheytt 1997).

Both PCA and CA were performed on the normalized data set of 13 physicochemical parameters during dry and wet seasons. Tables 6, 7 show the factor loading and eigenvalues of extracted components during dry and wet seasons, respectively, while Fig. 5a, b shows the dendrograms of analyzed parameters for dry and wet seasons while Fig. 6a, b depicts dendrograms for groundwater sampling points.

PCA, CA and ANOVA results during dry season

PC analysis identified five principal components accounting for 95.7% of the total variation in the original water quality data set during dry season.

PCI (Factor 1) accounts for 44.9% of the total variance, showing strong positive loading on EC, TDS, Ca²⁺, Mg²⁺ and TH, moderate loading for Na⁺ The strong positive loading factor of EC, TDS and TH may be interpreted as the influence of anthropogenic pollution from solid waste on the dumpsite while high loading of Ca²⁺, Mg²⁺ and TH may be

Table 4 Correlation coefficient of Ajakanga water samples parameters during dry season

	pH	EC	TDS	Cl ⁻	HCO ₃ ⁻	Hardness	CO ₃ ²⁻	SO ₄ ²⁻	NO ₃ ⁻	Na ⁺	K ⁺	Mg ²⁺	Ca ²⁺
pH	1												
EC	-.735 ^a	1											
TDS	-.736 ^a	1.000 ^b	1										
Cl ⁻	-.255	.680 ^a	.678 ^a	1									
HCO ₃ ⁻	-.001	.282	.286	.121	1								
Hardness	-.687 ^a	.784 ^b	.784 ^b	.165	.308	1							
CO ₃ ²⁻	-.001	.282	.286	.121	1.000 ^b	.308	1						
SO ₄ ²⁻	-.049	.032	.029	.504	-.256	-.229	-.256	1					
NO ₃ ⁻	-.106	-.264	-.266	.078	-.271	-.381	-.271	.837 ^b	1				
Na ⁺	-.481	.640 ^a	.640 ^a	.614	.270	.226	.270	-.004	-.162	1			
K ⁺	-.124	.154	.152	.422	-.153	.079	-.153	.569	.409	-.076	1		
Mg ²⁺	-.425	.795 ^b	.796 ^b	.448	.289	.746 ^a	.289	-.302	-.588	.230	.170	1	
Ca ²⁺	-.656 ^a	.748 ^a	.749 ^a	.148	.490	.907 ^b	.490	-.376	-.460	.421	-.185	.642 ^a	1

^aCorrelation is significant at the .05 level (2-tailed)

^bCorrelation is significant at the .01 level (2-tailed)

Table 5 Correlation coefficient of water samples parameters during wet season

	pH	EC	TDS	Cl ⁻	CO ₃ ²⁻	HCO ₃ ⁻	Hardness	SO ₄ ²⁻	NO ₃ ⁻	Na ⁺	K ⁺	Mg ²⁺	Ca ²⁺
pH	1												
EC	.075	1											
TDS	.077	1.000 ^b	1										
Cl ⁻	-.321	.717 ^a	.729 ^a	1									
CO ₃ ²⁻	.333	.889 ^b	.882 ^b	.344	1								
HCO ₃ ⁻	.333	.889 ^b	.882 ^b	.344	1.000 ^b	1							
Hardness	.164	.379	.391	.473	.338	.338	1						
SO ₄ ²⁻	-.104	.738 ^a	.738 ^a	.595	.541	.541	-.235	1					
NO ₃ ⁻	-.376	-.130	-.135	-.020	-.253	-.253	-.704 ^a	.360	1				
Na ⁺	-.361	.375	.387	.775 ^b	.055	.055	.626	.069	-.296	1			
K ⁺	.719 ^a	-.034	-.037	-.238	.131	.131	.036	-.104	-.367	-.020	1		
Mg ²⁺	.171	.961 ^b	.957 ^b	.538	.934 ^b	.934 ^b	.243	.730 ^a	-.057	.155	-.022	1	
Ca ²⁺	.277	.816 ^b	.809 ^b	.311	.840 ^b	.840 ^b	-.129	.836 ^b	.103	-.145	.114	.887 ^b	1

^aCorrelation is significant at the .05 level (2-tailed)

^bCorrelation is significant at the .01 level (2-tailed)

due to calcite and dolomite dissolution, weathering process, groundwater geological interaction and mineral precipitation. This positive loading on Ca²⁺, Mg²⁺ and TH suggest Ca²⁺, Mg²⁺ that probably contribute mostly to hardness of water in the study area.

Negative loading of pH with Ca²⁺ during dry season means a decrease in pH as the Ca²⁺ concentration in water rises (Mohapatra et al. 2011). PC2 accounted for 21.9% of the total variance, showing strong positive loading on SO₄²⁻ and this may be due to anthropogenic/organic wastes, atmospheric deposition, agricultural wastes, fertilizers and bacterial oxidation from dumpsite (Sidle et al. 2000) or atmospheric deposition (Wayland et al. 2003). Moderate

positive loading for K⁺ may be due to weathering of granite and magmatic rock, Cl⁻ may be due to anthropogenic waste from dumpsite or mineralization of groundwater while moderate loading of NO₃²⁻ may be an indication of livestock and municipal wastes from the dumpsite. PC3 accounts for 12.54% of the total variance while PC4 and PC5 accounted for 8.39 and 7.91% of the total variance.

During dry season, three clusters were identified on the dendrogram of physicochemical parameters. Cluster 1 showed a closed association between HCO₃⁻ and CO₃²⁻ and completely agrees with results of correlation coefficient analysis during dry season. Cluster 2 formed

Table 6 Factor loading and eigenvalues of extracted components during dry season

	Component				
	1	2	3	4	5
TDS	.945	.288	-.056	-.091	-.027
EC	.943	.292	-.058	-.093	-.027
Ca ²⁺	.892	-.235	-.125	.246	-.137
TH	.862	-.047	-.302	.336	.115
Mg ²⁺	.829	-.030	-.183	-.141	.466
pH	-.687	-.271	.386	-.363	.355
Na ⁺	.600	.202	.262	-.432	-.510
SO ₄ ²⁻	-.226	.884	.259	.167	-.030
K ⁺	.018	.707	.073	.184	.553
NO ₃ ⁻	-.451	.688	.189	.450	-.261
Cl ⁻	.497	.651	.335	-.425	.097
HCO ₃ ⁻	.495	-.411	.723	.242	.055
CO ₃ ²⁻	.495	-.411	.723	.242	.055
Initial eigenvalue	5.832	2.859	1.630	1.090	1.029
% of variance	44.858	21.991	12.538	8.386	7.912
Cumulative %	44.858	66.849	79.388	87.773	95.685

Table 7 Factor loading and eigenvalues of extracted components during wet season

	Component		
	1	2	3
EC	.996	-.048	-.071
TDS	.994	-.049	-.085
Mg ²⁺	.970	.002	.119
CO ₃ ²⁻	.910	.269	.108
HCO ₃ ⁻	.910	.269	.108
Ca ²⁺	.857	.008	.492
Na ⁺	.817	-.228	-.331
SO ₄ ²⁻	.757	-.421	.372
Cl ⁻	.672	-.398	-.475
pH	.157	.772	.457
K ⁺	.029	.753	.245
NO ₃	-.136	-.716	.553
TH	.337	.392	-.815
Initial eigenvalue	7.162	2.366	2.006
% of variance	55.096	18.203	15.428
Cumulative %	55.096	73.298	88.726

by Cl⁻, Na⁺, TH, Ca²⁺, Mg²⁺ and TDS are completely in accordance with correlation coefficient and PC1. This is an indication of common source. Cluster 3 during dry season comprises SO₄²⁻, NO₃⁻, K⁺ and pH. This is an indication of anthropogenic pollution from nearby dumpsite, agricultural wastes and effect of dumping wastes on pH.

On the basis of dendrogram of sampling points, two clusters were formed during dry season. Cluster 1 consists of samples S₂, S₃, S₄, S₅, S₇, S₈ and S₉ while cluster 2 consists of S₁, S₆ and S₁₀. These clusters of sampling points during dry season were grouping based on similar water quality characteristics. One-way ANOVA result shows significant difference at 5% level between the two clusters for EC, TDS, Cl⁻, Na⁺ and Ca²⁺. This is an indication that the EC, TDS, Cl⁻, Na⁺ and Ca²⁺ are physicochemical variables that differentiate the two identified clusters.

PCA, CA and ANOVA during wet season

Three principal components were extracted and accounted for 88.7% of the total variation in data set. Factor 1 accounts for 55.1% of the total variance and characterized by strong positive loading for EC, TDS, Ca²⁺, Mg²⁺, HCO₃⁻, CO₃²⁻, Na⁺ and SO₄²⁻, moderate loading on Cl⁻ and negative loading on NO₃⁻. The elements in PC1 probably show mineral components of groundwater, dissolution of carbonate minerals, rock–water geochemical reaction, dilution of groundwater, weathering and anthropogenic pollution. Dissolution of gypsum mineral could increase SO₄²⁻ concentration in groundwater (Yidana 2010).

Negative loading on NO₃⁻ may be due to action of denitrifying bacteria, laminar flow direction and diffusion process (Singh et al. 2008). Factor 2 has strong positive loadings for pH and K⁺ and accounts for 18.2% of the total variability in the data set while PC3 accounted for 15.4% of the total variance and has a strong negative loading for TH.

Negative loading of EC, TDS, Na⁺, SO₄²⁻, Cl⁻ and NO₃⁻ in PC 2 reflects their reduction due to dilution process during wet season.

On the basis of cluster analysis on physicochemical parameters, four clusters were identified during wet season. Cluster 1 comprises HCO₃⁻, CO₃²⁻, EC, TDS, Ca²⁺, Mg²⁺ and SO₄²⁻ and completely agrees with correlation coefficient analysis and PC1 during wet season. This is a cluster based on rock–water interaction, mineral dissolution and anthropogenic pollution source. Cluster 2 comprises Cl⁻, Na⁺ and TH, cluster 3 comprises pH and K⁺ and correlates very well with strong positive loading of pH and K⁺ during wet season while cluster 4 consists of only NO₃⁻ and corresponds with strong negative loading of NO₃⁻ on PC 2. The dendrogram schedule during wet season based on groundwater sampling sites depicts three (3) clusters. Cluster 1 comprises S₄, S₅, S₇, S₈ and S₉ which were scattered upstream of dumpsite and can be regarded as samples without influence of dumpsite. Cluster 2 comprises S₁, S₂, S₃ and S₆ samples on the southern part of the dumpsite. It should be noted that S₁, S₂, S₃ water samples in cluster 2 are within the vicinity of the dumpsite. Cluster 3 contains only sample S₁₀ which is an isolated hand-dug well within a

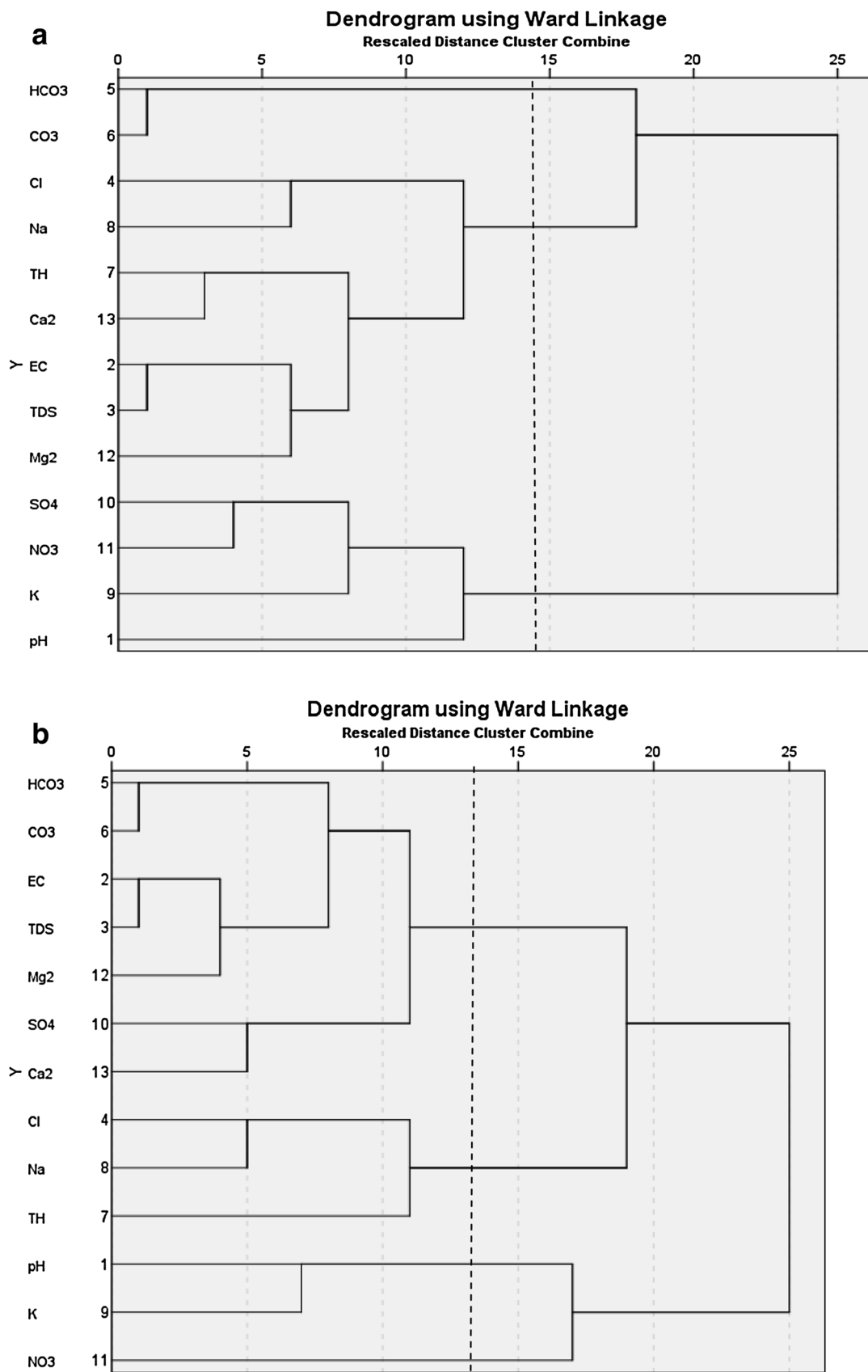


Fig. 5 a Dendrogram of analyzed parameters during dry season. b Dendrogram of analyzed parameters during wet season

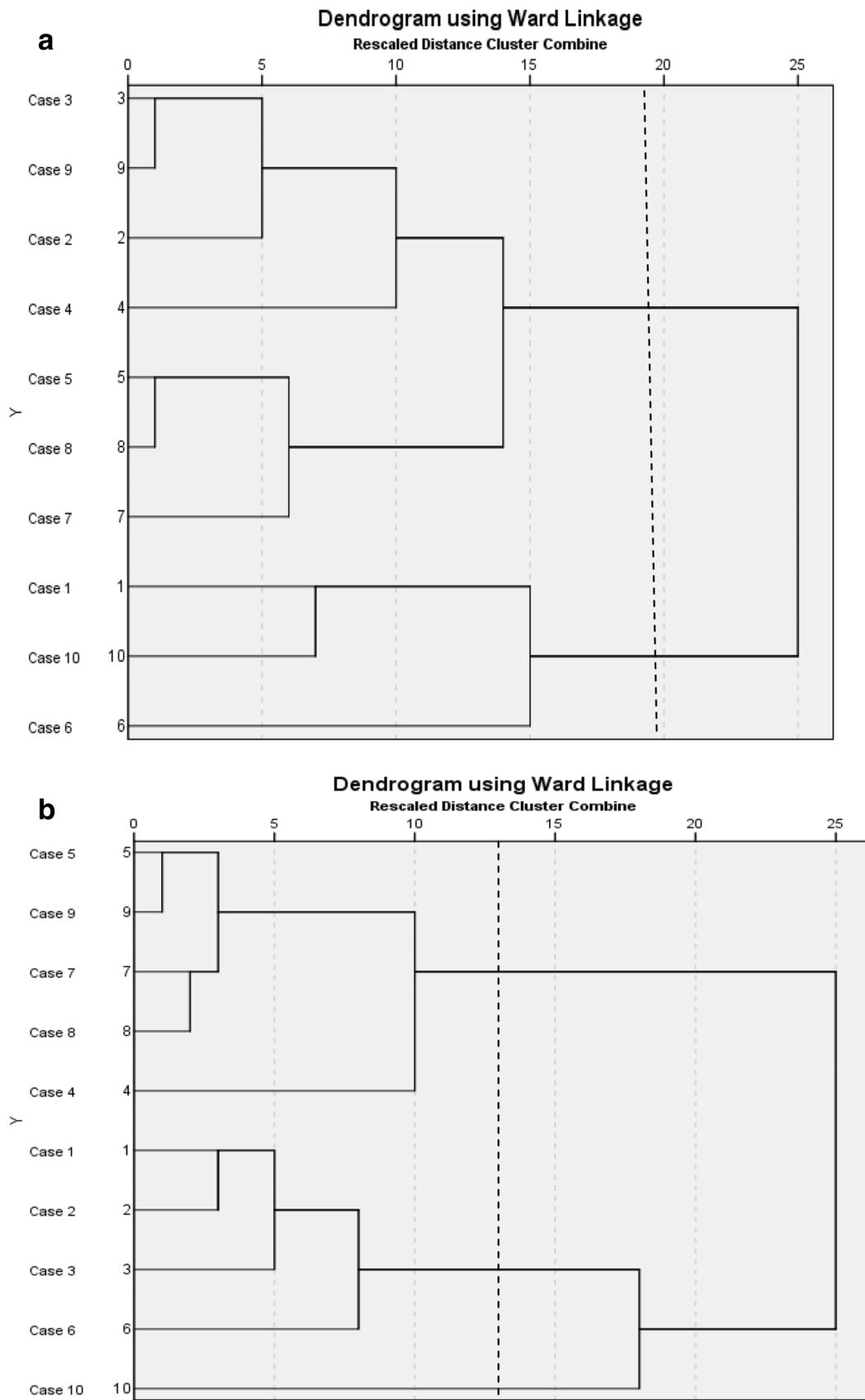


Fig. 6 a Dendrogram of water sampling locations during dry season. b Dendrogram of water sampling locations during wet season

Table 8 Irrigation parameters during dry season

Sample	SAR	% Na	SSP	RSBC	KR	PI
S ₁	1.201193	35.13	36.50	5.599	.55	106.8
S ₂	.730645	25.37	29.76	2.287	.36	90.8
S ₃	.660378	27.49	28.76	4.394	.38	135.8
S ₄	.923797	36.01	41.35	9.331	.61	207.6
S ₅	3.279626	82.59	87.97	3.579	6.86	298.8
S ₆	.921176	32.87	39.33	2.436	.54	110.2
S ₇	1.411331	54.16	62.19	5.934	1.43	266.5
S ₈	1.208652	50.74	54.70	1.891	1.12	167.4
S ₉	.6195	25.85	29.66	3.949	.37	134.8
S ₁₀	.611852	18.26	19.39	6.341	.23	84.8

Table 9 Irrigation parameters during wet season

Sample	SAR	%Na	SSP	RSBC	KR	PI
S ₁	.84831	27.14	27.88	4.162	.38	92.6
S ₂	.63066	19.87	21.83	4.868	.25	83.8
S ₃	.67644	23.44	24.27	6.086	.31	107.1
S ₄	.81534	29.14	35.21	3.247	.45	119.1
S ₅	.77695	37.90	39.97	3.117	.63	192.3
S ₆	1.22494	41.40	42.44	3.375	.72	117.9
S ₇	1.16686	51.66	53.86	1.90	1.11	175.7
S ₈	1.05194	43.60	45.23	2.379	.79	151.0
S ₉	.79447	31.80	34.29	3.056	.48	127.5
S ₁₀	.25789	4.83	5.21	1.329	.05	28.4

cultivated farmland on the upstream of the dumpsite. It should be noted that clusters of sampling points were based on similar topography setting, characteristics location of the sampling sites and vicinity with respect to the dumpsite.

The one-way ANOVA result shows that all the analyzed parameters except pH, K⁺, NO₃⁻ differ significantly at 5% level among the three clusters of groundwater samples. This implies that all the variables in the data set except pH, K⁺, NO₃⁻ are factors that discriminate one cluster from the other.

Groundwater quality for irrigation purpose

The suitability of groundwater for irrigation purpose was evaluated by calculating SAR, %Na, soluble sodium percentage (SSP), Kelly's ratio (KR), permeability index (PI) and residual sodium bicarbonate (RSBC). The results of calculated irrigation parameters are presented in Tables 8, 9 for dry and wet seasons, respectively.

The %Na for the groundwater samples from the study area was estimated using the formula:

$$\%Na = \frac{(Na^+) \times 100}{(Ca^{2+} + Mg^{2+} + Na^+ + K^+)}, \quad (1)$$

where the concentrations are in meq/L.

Table 10 Quality of groundwater based on %Na values

%Na	Water class	% during dry season	% during wet season
< 20	Excellent	10	20
20–40	Good	60	50
40–60	Permissible	20	30
60–80	Doubtful	–	–
> 80	Unsuitable	10	–

The classification of groundwater samples based on %Na values is shown in Table 10.

SAR for the groundwater samples was estimated from the formula (Karanth 1987):

$$SAR = \frac{Na^+}{\sqrt{\frac{Ca^{2+} + Mg^{2+}}{2}}}. \quad (2)$$

The water samples having SAR values less than 10 are considered excellent, 10–18 as good, 18–26 as fair (doubtful) and above 26 as unsuitable for irrigation use (USDA 1954). In the present study, the SAR values for both seasons are less

than 10 and can thus be graded as “Excellent” for irrigation use (as shown in Tables 8 and 9).

Kelly’s ratio (KR) was calculated by using the numerical formula (Kelly 1963):

$$KR = \frac{Na^+}{Ca^{2+} + Mg^{2+}}, \tag{3}$$

where concentrations are expressed in meq/L.

The Kelly’s ratio of 1 or less than 1 is an indication of good quality water for irrigation purpose, whereas above one is suggestive of unsuitable for agricultural purpose due to alkali hazards (Karanth 1987). It is observed from Tables 8 and 9 that, 70% of the samples in the study area have KR values below 1, thus belonging to “Good” class while 30% belongs to “Unsuitable” class during dry season. However, in wet season, 90% belong to “Good” class while 10% of samples belong to “Unsuitable” class for irrigation need.

The residual sodium bicarbonate (RSBC) was determined using the formula (Gupta and Gupta 1987):

$$[(HCO_3^-) - (Ca^+)], \tag{4}$$

where the ion concentrations are in meq/L.

According to USDA (1954), RSBC values exceeding 2.5 meq/L is “Unsuitable for irrigation”, if the value of RSBC lies between 1.25 and 2.5 meq/L, it is “marginally suitable” while a value less than 1.25 meq/L indicate safe water quality. Based on this classification, during the dry season, 70% fall under “Unsuitable class and 30% fall under “marginally suitable”. However, in wet season, 70% of analyzed samples still fall under “Unsuitable” class while 30% fall under “Marginally suitable” class based on RSBC values (Tables 8 and 9).

Todd (1995) defines soluble sodium percentage (SSP) as:

$$SSP = \frac{(Na^+ + K^+) \times 100}{(Na^+ + K^+ + Ca^{2+} + Mg^{2+})}, \tag{5}$$

where the concentrations are in meq/L.

The classification of groundwater for irrigation purpose based on SSP value is shown in Table 11.

The permeability index is calculated by using the formula (Ragunath 1987):

$$PI = \frac{(Na^+ + \sqrt{HCO_3^-})}{(Ca^{2+} + Mg^{2+} + Na^+)} \times 100, \tag{6}$$

where the concentrations are expressed in meq/L.

The PI values > 75 indicate excellent quality water for irrigation. PI values less than 25 reflect “unsuitable” water for irrigation. On the basis of PI values in Tables 8 and 9, all the water samples from the study area during dry season can be classified as “Excellent” class for agricultural use. During

Table 11 Quality of groundwater based on SSP values

SSP values	Water class	% during dry season	% during wet season
< 60	Excellent	80	100
60–75	Good to permissible	10	–
> 75	Doubtful to unsuitable	10	–

wet season, 90% belong to “Excellent” class while only 10% falls under “Doubtful to unsuitable” class.

Conclusion

The study provides information about the quality of groundwater from hand-dug wells at several locations closed to Ajakanga dumpsite. The major ions in all analyzed groundwater samples were found to lie within the standard limits of WHO (2007) and NSDWQ (2007). However, high concentration of some water quality parameters were noticed in Wells 1 and 10, which may be due to effect of leachate migration towards the southern part of the dumpsite; nearness to dumpsite; agricultural run-off and fertilizer application.

Five principal components with three factors were responsible for 95.7 and 88.7% of the total variance in the data set during dry and wet seasons, respectively. PCA identified parameters influencing water quality were probably related to mineral dissolution, groundwater–rock interaction, weathering process and anthropogenic activities from the dumpsite while cluster analysis based on groundwater samples during dry and wet seasons showed 2 and 3 significant clusters, respectively.

The dendrogram also reflects variation of water quality with climatic season as shown in the differing number of clusters during both seasons. The analyzed physicochemical parameters that explained more than 40% of the total variance in the original data set during both seasons were: EC, TDS, Ca²⁺, Mg²⁺ and TH. Calculated irrigation parameters values indicate that, sizeable number of groundwater samples will neither cause salinity hazards nor have adverse effects on soil properties and thus suitable for irrigation needs.

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