HYDROGEOCHEMICAL STUDY OF GROUNDWATER FROM DEEPER AQUIFER OF WRE-2 WATERSHED OF CHANDRAPUR AND BALLARPUR TALUKA, CHANDRAPUR DISTRICT, MAHARASHTRA

*Murkute Y.A.¹ and Solanki V.V.²

¹Department of Geology, R.T.M. Nagpur University, Law College Square, Nagpur-440001, India ²Groundwater Survey and Development Agency, Chandrapur-442401, India *Author for Correspondence

ABSTRACT

In the present study, evaluation of geochemistry of 33 water representative deeper aquifer groundwater samples from WRE2 watershed of Chandrapur and Ballarpur taluka of Chandrapur District Maharashtra is carried out. The groundwater from the deeper aquifer of watershed WRE2 area is alkaline in nature. The higher conductivity in the study area is due to high mineral concentration in groundwater. It indicates that the ion exchange and solublization processes have taken place within the aquifer. The interrelationship among $Ca^{++} + Mg^{++} vs SO_4 + HCO_3$ exhibits the calcium dissolution and silicate weathering for the source for Na⁺ in study area. Besides, the interrelationship among HCO₃ and Cl *vs* Na also point out calcium dissolution and silicate weathering. Considering the TDS concentration the 27% groundwater samples from borewell from the study area is fresh water. The major plots of groundwater sample fall in C4-S2 and C3-S2 with minor plots in C3-S1 in US Salinity diagram (1954), which indicate groundwater is of medium to high salinity-medium sodium type and thus can be utilized for irrigation with little danger of exchangeable sodium.

Keywords: Hydrogeochemistry, Deeper Aquifer Groundwater, Water Quality, WRE2 Watershed, Chandrapur District, Maharashtra

INTRODUCTION

Groundwater is a most vital natural resources required for drinking and irrigation. The quality of groundwater is largely controlled by discharge-recharge pattern, nature of host and associated rocks as well as contaminated activities. Moreover, the nature and amount of dissolved species in natural water is strongly influenced by mineralogy and solubility of rock forming minerals (Raymahasay, 1996). The quality of groundwater is function of various parameters which determines its suitability for industrial use as well as irrigation practices and most importantly drinking purposes (WHO 1984; Trivedy and Goel 1986; ISI 1991; APHA 1998)

The requirement of water for all living beings is utmost important. The ultimate source of water is only one i.e. rainfall, there are two phases of water, namely surface water and groundwater. The groundwater occurrence and accessibility has been inconsistent all over the India. This may be ascribed to precipitation percentage, physiography and diverse hydrogeological conditions. Both, the unmanaged utilization as well as excessive groundwater exploitation are exclusively responsible for lowering of groundwater levels, de-saturation of aquifers. The groundwater from shallow aquifer occurs in dugwells and shallow tubewells whereas from deep aquifers it is obtained by means of deep borewells.

The availability of good quality of water in high quantity is linked with food security. However, there are number of places on the earth surface where quality of groundwater is deteriorated, either by nature or due to human interventions and groundwater resources are moreover not suitable for drinking purpose, The various studies on groundwater quality, geochemical characterization and its utility of various purposes have been carried out in different parts of India (Saxsena and Ahmed, 2001; Shivanna and Mahokar, 2003; Subba Rao, 2006; Raju, 2007; Gupta *et al.*, 2008; Subba and Surya, 2009; Bharadwaj *et al.*, 2010; Marghade *et al.*, 2010). The high content of arsenic (Pal and Mukharji, 2008) and fluoride (Madhnure and Malpe, 2007; Subba, 2009) have been reported from groundwaters from different parts of India. In the Central India, particularly in east parts of Maharashtra State the districts like Yavatmal and

Chandrapur have high concentrations of fluoride content (Murkute and Badhan, 2011, Murkute, 2014; Dharashivkar *et al.*, 2014). The present study is also an attempt to understand the hydrogeochemical characteristics of deeper aquifer of groundwater in WRE2 watershed of Chandrapur District (Maharashtra), India.

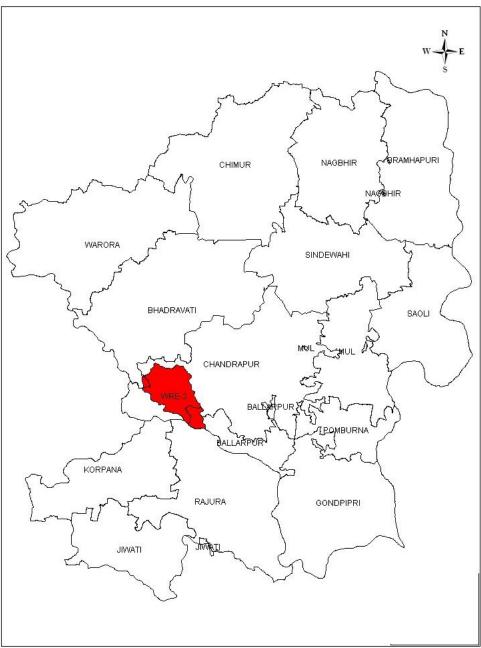


Figure 1: Location and Geological map of the study area

Geological Setting

The shale and sandstone of the Mesoproterozoic age belonging to Penganga Group form the basement along with the underlying by the Archaean metamorphics (Table 1). These rocks are unconformably overlain by the sediments of Gondwana Supergroup of Permo-carboniferous to Jurassic age. Geologically the area is dominated by Talchir shale and attained almost 80 to 100 m thickness, as noticed from

borehole data (DGM, 2000). The Talchir Formation mainly comprises light green shale and sandstone. The Talchir Formation exhibits conformable relationship with the succeeding Barakar Formation. The rock types of Barakar formations are feldspathic sandstone and shale and are overlain by sandstones of Kamthi Formation. In turn, Deccan Trap basaltic flows cover the area and local alluvial patches can be noted from Wardha and Irai rivers which form the southern and eastern boundary of the watershed.

Age	Formation	Rock Type
Recent to subrecent	Alluvium, Soils	Sandy Clay silty soil
Lower Eocene to Upper cretaceous	Deccan Traps	Basalts, weathred, vesicular and massive basalt
	Unconfermity	
Lower Triassic to	Kamthis	Reddish Brown Sandstone,
Upper Carboniferous	Unconfermity	Shale, Clay
	Barakar	Light grey to white feldspathic
	shales,	Sandstones, carbonaceous
	Talchirs	Coal Seams and Clay Greenish to dark Olive green Colored shales and coarse
	Unconfermity	grained sandstones.
Precambrian	Vindhyan	Shales, sandstones, flaggy and massive limestone and sandstones of variegated colours
	Unconfermity	colours
Archeans	Crystalline and older metamorphic	Gneisses, quartzites, schists with acid and basic intrusive

Table 1: Geological	succession of	the area is	σiven	helow ((DGM. 2000)
Table 1. Ocological	succession of	the area is	gruun		

Study Area

Location and Climate

The watershed WRE-2 covers 25 villages of Chandrapur and Ballarpur tahsil of Chandrapur district of Maharashtra, (Central India). The study area falls in latitude 19⁰53'00" to 20⁰01'35" N and longitude 79⁰10'40" to 79⁰17'35" E of Survey of India toposheet no. 56M/1, 56M/5, 55P/8 and 55P/4 covering 16231 ha area. Total annual groundwater recharge in study area is 769.85 ham with net annual groundwater availability as 731.36 ham and represents the 'Safe' watershed (GSDA, 2011). The study area is mainly covered by major part of Chandrapur MIDC that includes small scale to large scale industries comprising cement product plants, chemical plants, fly ash depot, soybean oil plant etc. The

climate of the area is hot. The maximum temperature in summer reaches up to 47 0 C and in winter about 28 0 C. The average annual rainfall of the area is 1030 mm.



Figure 2: Showing watershed map of wre2 watershed with drainage pattern and village from water samples collected

Physiography and Drainage

The area exhibits moderately dissected topography with general slope towards southeast direction. The watershed is mainly drained by Erai and Wardha rivers and their tributaries. The streams of order I and III creates dendritic drainage pattern and higher order drainages run roughly parallel. The eastern boundary of watershed is marked by southerly flowing Erai River and southern boundary is marked by SW flowing Wardha River.

Hydrogeological Framework

In the area, the groundwater mainly occurs under phreatic conditions; however the deeper confined aquifers are also common. The dugwells penetrating Penganga limestones range in depth between 5 to 15 mbgl. The diameters of the dugwells range between 3 to 5 m. The deeper aquifers vary in depth from 25 to 60 mbgl upholding a discharge of 50 to 300 m³/ day (GSDA, 2005, 2009).

© Copyright 2014 | Centre for Info Bio Technology (CIBTech)

From Talchir, Barakar and Kamthi formations, the groundwater mainly occurs under phreatic conditions; however the confined aquifers are also not uncommon. The borewells from the Gondwana sedimentary province vary in depth stuck between 60 m and uphold a discharge of 50 to 300 m³/ day (GSDA, 2005). Murkute *et al.*, (2010) have established the relationship between aquifer parameters and the petrological characteristics of Kamthi Sandstone, exposed near Minjhari village, Chandrapur District. According to them the values of transmissivity and specific yield for the arenites range from 104.85 to 412.72 m²/day and 20 to 29 % respectively. These high values of transmissivity and specific yield in arenites aquifer are accountable for higher percentage of detrital grains, lesser amount of matrix and the moderate sorting of the grains. The values of transmissivity and specific yield for the graywackes grade from 59.82 to 146.43 m²/day and 11 to 16 % respectively. The lower percentage of detrital grains, higher amount of matrix and the poor or very poor sorting of the grains are responsible for low values of transmissivity and specific yield in graywacke aquifer.

The Basaltic lava flows are vesicular in nature and possess deep weathering as well as joints. The average depth of dugwells varies from 9 to 15 mbgl and the yield ranges from 75 to 100 m³/ day. The borewells penetrating deeper aquifers have good yields ranging from 150 to 250 m³/ day. The wells piercing deeper aquifers in alluvial areas have discharge in between 100 to 300 m³/ day, though in a few cases higher yields are observed (GSDA, 2005; 2009).

MATERIALS AND METHODS

Sr.No	Parameters	Characteristics	Unit
1	General	Tempreture	⁰ C
2		pH	
3		Electrical conductity	µs/cm
4		Total dissolved solids	mg/L
5		Total hardness(as CaCo ₃)	mg/L
6	Major Cations	Calcium (Ca ²⁺)	mg/L
7		Magnesium(Mg ²⁺)	mg/L
8		Sodium(Na ⁺)	mg/L
9		Potassium(K ⁺)	mg/L
10	Major anions	Bicarbonate(HCO ³⁻)	mg/L
11		Nitrate(NO ₃ ⁻)	mg/L
12		Sulphate($SO_4^{2^-}$)	mg/L
13		Chloride(Cl ⁻)	mg/L
14		Flouride(F)	mg/L
15	Irrigation Water use (by calculation)	Sodium Absorption Ratio(SAR)	meq/L
16		Percent Sodium(%Na)	%
17		Residual Sodium Carbonate(RSC)	meq/L
18		Mg Ratio(MR)	
19		Corrosivity Ratio (CR)	

Table 2: Analytical methods followed in present study

© Copyright 2014 / Centre for Info Bio Technology (CIBTech)

Table 3: Analytical data of water samples collected from study area

Se. No.	Sample No.	Forma- tion	Aquifer/ Litholog Y	Source	рН	EC μs/cm	TDS	тн	Ca++	Mg++	Na+	K+	NO3-	HCO3-	SO4	CI-	F-
1	WRE2-1	Talchir	Shale	Borewell	7.5	2080.00	1331.2	179.95	44.10	17.00	190.00	1.50	5.46	632.00	285.80	160.00	0.44
2	WRE2-3	Talchir	Shale	Borewell	7.2	2180.00	1395.2	500.08	132.30	41.30	210.00	49.00	160.87	500.00	276.00	179.90	0.54
3	WRE2-5	Talchir	Shale	Borewell	8.2	2120.00	1356.8	378.98	28.10	75.30	260.00	109.00	92.83	380.00	258.90	249.90	1.10
4	WRE2-8	Alluvium	Gravely sand	Borewell	8.0	1718.00	1099.5	339.67	60.10	46.20	222.00	20.00	145.61	280.00	124.60	194.90	1.24
5	WRE2-9	Talchir	Shale	Borewell	7.8	2250.00	1440	1196.8	80.20	243.00	372.00	9.00	88.70	580.00	160.00	359.90	1.08
6	WRE2-10	Talchir	Shale	Borewell	7.6	2760.00	1766.4	1197.5	132.30	211.40	350.00	10.00	90.25	500.00	224.70	329.90	0.40
7	WRE2-11	Talchir	Shale	Borewell	7.8	2050.00	1312	420.15	108.20	36.50	217.00	27.00	193.30	288.00	166.10	229.90	0.77
8	WRE2-12	Alluvium	Gravely sand	Borewell	7.6	2970.00	1900.8	439.9	120.20	34.00	330.00	14.50	681.95	480.00	269.90	409.90	0.83
9	WRE2-15	Alluvium	Gravely sand	Borewell	7.6	3140.00	2009.6	399.83	92.20	41.30	460.00	8.50	126.22	300.00	258.90	599.80	0.66
10	WRE2-17	Barakar	Fels pathic	Borewell	7.6	1071.00	685.44	480.24	128.30	38.90	82.00	7.00	51.19	230.00	289.50	84.00	0.49
11	WRE2-18	Talchir	Shale	Borewell	7.6	2390.00	1529.6	698.72	92.20	114.20	198.00	8.50	69.15	340.00	434.80	209.90	0.30
12	WRE2-20	Talchir	Shale	Borewell	7.9	1260.00	806.4	279.72	68.10	26.70	120.00	20.00	29.01	430.00	64.70	89.00	0.71
13	WRE2-23	Talchir	Shale	Borewell	7.9	1800.00	1152	559.46	116.20	65.60	240.00	5.50	116.04	330.00	114.80	304.90	0.57
14	WRE2-24	Alluvium	Gravely sand	Borewell	8.0	3130.00	2003.2	798.67	100.20	133.70	390.00	37.50	209.04	360.00	414.10	489.80	0.73
15	WRE2-26	Alluvium	Gravely sand	Borewell	7.8	5710.00	3654.4	1298.1	192.40	199.30	1280.00	42.00	131.94	360.00	978.30	1524.50	0.73
16	WRE2-28	Barakar	Fels pathic sst	Borewell	7.9	3780.00	2419.2	918.67	152.30	131.20	550.00	72.00	213.41	440.00	447.00	699.80	0.26
17	WRE2-29	Talchir	Shale	Borewell	7.8	5030.00	3219.2	1438.2	264.50	189.50	880.00	117.00	353.69	500.00	647.30	999.70	0.28
18	WRE2-30	Talchir	Shale	Borewell	8.0	3140.00	2009.6	638.63	72.10	111.80	490.80	60.50	224.54	360.00	263.80	599.80	0.50
19	WRE2-31	Talchir	Shale	Borewell	7.7	7160.00	4582.4	2240.2	601.20	179.80	1430.00	115.50	225.73	400.00	1027.20	1824.40	0.61
20	WRE2-32	Talchir	Shale	Borewell	8.2	2410.00	1542.4	839.59	188.40	89.90	260.00	16.00	17.88	520.00	505.70	189.90	0.97
21	WRE2-34	Talchir	Shale	Borewell	7.7	5910.00	3782.4	1299.3	280.60	145.80	2160.00	40.00	61.20	460.00	1023.50	2724.20	0.96
22	WRE2-36	Talchir	Shale	Borewell	8.6	947.00	606.08	139.54	24.00	19.40	292.50	18.00	5.40	212.00	57.40	230.00	0.54
23	WRE2-39	Talchir	Shale	Borewell	8.0	4400.00	2816	880.6	264.50	53.50	640.00	132.50	74.32	380.00	621.70	849.70	0.61
24	WRE2-40	Barakar	Fels pathic sst	Borewell	8.1	1476.00	944.64	479.67	120.20	43.70	200.00	21.50	19.08	300.00	73.30	135.00	0.30
25	WRE2-42	Alluvium	Gravely	Borewell	8.2	824.00	527.36	340.29	100.20	21.90	90.00	20.50	7.15	196.80	31.80	54.00	0.18
26	WRE2-44	Barakar	Fels pathic sst	Borewell	8.2	2500.00	1600	839.68	180.40	94.80	349.50	31.00	0.99	420.00	377.40	249.90	0.35
27	WRE2-45	Barakar	Fels pathic sst	Borewell	8.3	1240.00	793.6	1158	148.30	192.00	227.00	28.50	15.28	272.00	67.20	214.90	0.23

© Copyright 2014 | Centre for Info Bio Technology (CIBTech)

Research Article

28	WRE2-47 Barakar	Fels pathic	Borewell	7.8	2900.00	1856	941.47	328.70	29.20	398.00	103.00	12.27	540.00	247.90	369.90	0.28
		sst Fels														
29	WRE2-50 Barakar	pathic sst	Borewell	8.0	2350.00	1504	819.73	204.40	75.30	388.50	24.00	13.43	360.00	173.40	369.90	0.20
		Fels														
30	WRE2-52 Barakar	pathic sst	Borewell	8.1	1698.00	1086.7	479.69	100.20	55.90	300.00	24.00	6.91	270.00	171.00	239.90	0.40
31	WRE2-54 Talchir	Shale	Borewell	8.4	1460.00	934.4	239.81	44.10	31.60	110.00	24.00	5.56	300.00	276.00	89.00	0.44
			D 11												~~ ~~	
32	WRE2-55 Talchir	Shale	Borewell	8.4	1230.00	787.2	219.79	56.10	19.40	88.00	24.00	4.93	288.00	235.70	69.00	0.65
22		Chala	D	0.1	776.00	105.54	200.40	60.10	7 20	02.00	10 50	F 00	256.00	45.20	64.00	0.72
33	WRE2-57 Talchir	Shale	Borewell	8.1	776.00	490.64	200.18	68.10	7.30	92.00	19.50	5.88	256.90	45.20	64.00	0.72

Cation Chemistry

Among the cations, Na⁺ and Ca⁺⁺ are the major constituents in the groundwater of the study area and ranges from 82 to 2160 mg/l and 24 to 601.20 mg/l respectively. The concentration of Mg⁺⁺ from borewell samples ranges from 7.30 to 243 mg/l. The principal source of Mg⁺⁺ in the natural waters is the magnesium bearing mineral present in the rocks. The domestic and industrial wastes also higher up the Mg⁺⁺ concentration. Generally percentage of Mg⁺⁺ content remains less than the Ca⁺⁺ in groundwater samples. The K⁺ contents in deeper aquifer range between 1.50 to 132.50 mg/l. The Na⁺ and K⁺ in groundwater samples from borewells related to weathered rock-forming minerals like sodium plagioclase, potash plagioclase and anthropogenic sources like domestic and animal waste (Marghade *et al.*, 2010). Handa (1975); Jacks *et al.*, (2005); Murkute (2014) have pointed out increased Na⁺ content with decreasing Ca⁺⁺ concentration in alkaline water conditions. However, in the present study negates such behaviour of these cations (Figure 1). Such behaviour of the Na⁺ and Ca⁺⁺ probably points out some another chemical mechanism which releases Na⁺ in groundwater along with release of Ca⁺⁺ and Mg⁺⁺.

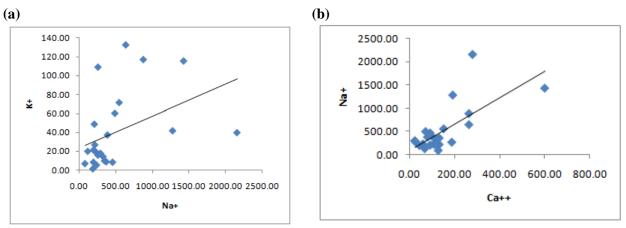


Figure 3: Interrelationship between cations a)Na⁺ vs K⁺ and b) Ca⁺⁺ vs Na⁺

Anion Chemistry

In groundwater samples from the study area HCO_3^- and SO_4^- are the leading ions. HCO_3^- content from the study area grades from 196.80 to 632 mg/l. The dissolution of CO_2 gas through anoxic biodegradation of organic matter derived from industrial and domestic waste in shallow aquifers is the source of HCO_3^- in groundwater (Canter, 1997; Jeong, 2001). Hitherto no health hazards due to HCO_3^- have been identified; however it should not exceed 300mg/l (WHO, 1997). However low values of HCO_3^- content is noted from the Alluvium and Barakar formation than the Talchir formation. The SO_4^- concentration grades from 31.80 to 1027.20 mg/l showing that 73 % of sample have SO_4^- concentration within potable limit (BIS,

1991). NO_3^- content varies between 0.99 to 681.95 mg/l indicating that 58.6% of groundwater samples have concentration of NO_3^- more than desirable limit (BIS, 1991). The Cl⁻ concentration in the water samples of the study area ranges between 54 to 2724.20 mg/l pointing that 45.5% of water samples have concentration above desirable limit (BIS, 1991). The higher value of Cl⁻ in the water is usually considered as pollution and indicative of groundwater contamination (Loizidou and Kapetanions, 1993). According to them the high Cl⁻ concentration in groundwater comes from weathering of minerals as well as from the percolation of domestic effluents and fertilizers. The F⁻ content in borewells of study area range between 0.18 to 1.24mg/l indicating that the groundwater from the study area have F⁻ concentration within permissible limit (BIS,1991).

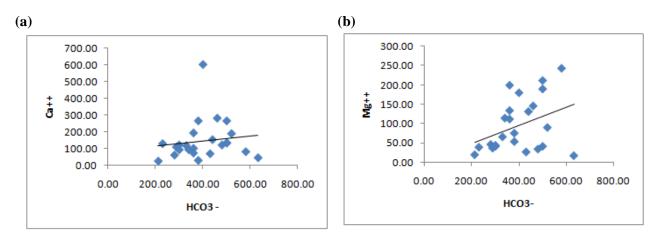
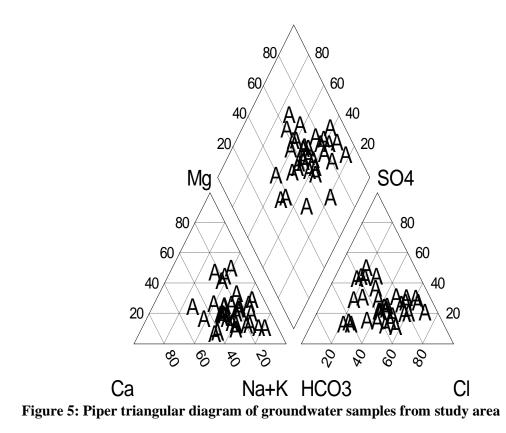


Figure 4: Relationship of cations and anions a) HCO₃⁻ vs Ca⁺⁺ and b) HCO₃⁻ vs Mg⁺⁺



© Copyright 2014 | Centre for Info Bio Technology (CIBTech)

Rock-Water Interaction

The Ca⁺⁺ and HCO₃⁻ as well as Mg⁺⁺ and HCO₃⁻ show positive correlation between these ions because these cations have common tendency to combine with HCO₃⁻ as noted by Todd (1982). The interrelationship among Ca⁺⁺+ Mg⁺⁺ vs SO₄ + HCO₃ exhibits the calcium dissolution and silicate weathering for the source for Na⁺ in study area (Figure 3). The interrelationship among HCO₃ and Cl vs Na also point out calcium dissolution and silicate weathering (Figure 4). The feldspars, calcite and clay minerals are the probable sources of Ca⁺⁺ from sedimentary rocks (Hem, 1970; Todd, 1982; Murkute, 2014). The chief sources of Mg⁺⁺ in natural water are the magnesium bearing minerals like pyroxenes, olivine and amphiboles (Singh *et al.*, 2010; Marghade *et al.*, 2010) while the feldspar and clay minerals are the probable sources for Na⁺ and K⁺. SO₄⁻ is generally derived from the oxidative weathering of sulphate bearing minerals like pyrite (FeS₂) which is very common secondary minerals in the Gondwana formations and associated sediments (Singh *et al.*, 2010). The F⁻ concentration in groundwater depends upon the degree of weathering and leaching of fluoride bearing minerals from the rocks and soils (Ramesam and Rajagopalan, 1985; Murkute and Badhan, 2011). Besides, the source of F⁻ is also expected to be the amphibolites and biotite (Madhnure and Malpe, 2007).

Major Ions	Min	Max	Mean	WHO (199	WHO (1997)BIS)500
				Desirable	Permissible	Desirable	Permissible
pН	7.4	8.5	7.95	7.0-8.5	6.5-9.2	6.5-8.5	8.5-9.2
EC	565	1723	1144	750	1500		
TDS	362	1103	732.5	500	1500	500	2000
TH	289	1302	795.5	100	500	300	600
Ca++	62.4	431.3	246.85	75	200	75	200
Mg++	6.1	244.4	125.25	30	150	30	100
Na+	3.2	101.2	52.2	50	200		
K+	2.1	61.3	31.7	100	200		
NO3-	7.6	134.3	70.95		50	45	100
HCO3-	74	273	173.5	200	600	200	600
SO4	5.6	356.7	181.15	200	600	200	400
Cl-	3.8	126.5	65.15	250	600	250	1000
F-	0.3	0.8	0.55	0.6-0.9	1.5	1	1.5

 Table 4: Geochemical parameters of the study area in comparision to WHO(1997) and BIS(1991)

The geochemical evaluation of groundwater can be understood by plotting the concentration of major cations and anions in the piper diagram (Piper, 1953). In this diagram the relative abundance of cations *i.e.* Na+K, Ca and Mg in % meq/l is plotted on the cation triangle followed by the relative abundance of anions *i.e.* Cl+NO₂=NO₃, SO₄ and HCO₃+CO₃ in % meq/l is plotted on the anion triangle. The central diamond-shaped field (quadrilateral field) is used to show overall chemical character of the water (Hill, 1940; Piper, 1944). Thus, the piper diagram directly deciphers the groundwater quality, and as a result it is commonly used as an effective tool to specify different water-types. In general, piper diagram reveals six types with nine combinations, according to ionic placement in the diamond field. The Piper's trilinear diagram (Piper, 1953) prepared for the samples from the study area is presented in Figure 5. The plots of the chemical data on Piper's trilinear diagram show that non carbonate alkalis exceed 50% in the groundwater samples of the study area. In addition, some plots on Piper's trilinear diagram also suggest no one cation-anion pair exceeds 50%. In general, 26 water facies are noted from the study area, out of which Na-Ca-HCO3-Cl water-type is the dominant facies (34%). *Water Quality Assessment*

Se. No.	Sample No.	Formation	SAR	% Na	RSC	MR
1	WRE2-1	Talchir	6.158	8.587	27.991	38.813
2	WRE2-3	Talchir	4.083	15.277	14.984	33.936
3	WRE2-5	Talchir	5.802	24.156	11.397	81.514
4	WRE2-8	Alluvium	5.235	12.670	7.192	55.849
5	WRE2-9	Talchir	4.671	16.751	4.996	83.294
6	WRE2-10	Talchir	4.395	15.873	0.992	72.447
7	WRE2-11	Talchir	4.603	13.163	5.986	35.695
8	WRE2-12	Alluvium	6.841	15.931	15.190	31.762
9	WRE2-15	Alluvium	10.001	20.780	6.991	42.433
10	WRE2-17	Barakar	1.627	4.907	1.884	33.285
11	WRE2-18	Talchir	3.255	9.565	2.993	67.085
12	WRE2-20	Talchir	3.119	9.733	15.895	39.216
13	WRE2-23	Talchir	4.411	11.086	5.292	48.159
14	WRE2-24	Alluvium	5.997	19.791	1.989	68.708
15	WRE2-26	Alluvium	15.439	56.978	-8.016	63.025
16	WRE2-28	Barakar	7.887	28.093	3.590	58.636
17	WRE2-29	Talchir	10.086	42.549	-3.816	54.106
18	WRE2-30	Talchir	8.439	25.683	5.195	71.844
19	WRE2-31	Talchir	13.137	64.890	-24.848	32.982
20	WRE2-32	Talchir	3.901	12.744	9.182	43.984
21	WRE2-34	Talchir	26.049	94.805	-3.025	46.092
22	WRE2-36	Talchir	10.762	15.604	9.402	57.084
23	WRE2-39	Talchir	9.379	34.775	1.373	24.972
24	WRE2-40	Barakar	3.970	11.616	5.393	37.432
25	WRE2-42	Alluvium	2.122	8.575	3.131	26.452
26	WRE2-44	Barakar	5.243	17.619	4.179	46.373
27	WRE2-45	Barakar	2.899	12.030	-9.344	68.055
28	WRE2-47	Barakar	5.642	24.104	8.161	12.754
29	WRE2-50	Barakar	5.899	18.709	1.584	37.741
30	WRE2-52	Barakar	5.954	15.687	3.890	47.863
31	WRE2-54	Talchir	3.087	10.800	11.526	54.110
32	WRE2-55	Talchir	2.581	10.769	11.063	36.267
33	WRE2-57	Talchir	2.828	9.864	8.938	14.994

The data obtained by geochemical analyses are evaluated in terms of suitability for drinking, irrigation, livestock and industrial uses.

Drinking and General Domestic Use

The prescribed limits of WHO (1997) and Indian drinking water standards (BIS, 1991) were used to assess the suitability for drinking and public health purposes (Table 4). The major objectionable physicochemical parameters are TDS, total hardness (TH) and sulphate. Carrol (1962) has propounded the relationship of TDS and salinity as 0-1000 (mg/l), fresh water; 1000-10000 (mg/l), brakish water; 10000-100000 (mg/l), saline water and >100000 (mg/l) it is brine. According to his classification scheme the 27% groundwater samples from borewell from the study area is fresh water.

Research Article

However, 77% of groundwater samples are brakish water and such high salinity in few samples is attributed to concentration of soluble salts within the geological formation (Todd, 1980). Based on TH, Sawyer and Mccarty (1967) have categorized the water as soft (< 75mg/l) moderately hard (75-150mg/l), hard (150-300 mg/l) and very hard water (> 300mg/l). The analyzed data indicate that only 18% groundwater samples have TH value lesser than 300 mg/l, which is the desirable limit as per BIS (1991), and remaining 82% are of very hard type.

The long term consumption of very hard water leads in increase incidences of urolithiasis, some types of cancer, permanent mortality, anecephaly and cardio-vascular disorders (Agrawal and Jagetia, 1997; Durvey *et al.*, 1991). 73% groundwater samples in the study area have the SO_4^- concentration less than 400 mg/l which is the permissible limit of BIS (1991). However, the water with 200-400 mg/l SO_4^- has a bitter taste and it may also cause corrosion of metals particularly in water with low alkalinity (Singh *et al.*, 2010).

Irrigation Use

In the present study, the suitability of groundwater has been checked for irrigation purpose using Sodium Absorption Ratio (SAR), Percent Sodium (%Na), Residual Sodium Carbonate (RSC), Mg Ratio (MR), Corrosivity Ratio (CR) (Table 5).

Sodium Absorption Ratio (SAR): The Sodium Absorption Ratio (SAR) is defined by the expression: SAR $= Na^+ / \sqrt{[(Ca^{++}+Mg^{++})/2]}$. Water with SAR value ≤ 10 is considered as of excellent quality, 10 to 18 is good, from 18 to 26 is fair and above 26 is said to be unsuitable for irrigation. The calculated SAR values in study area range from 1.627 to 26.049 meq/l. The major plots of groundwater sample fall in C4-S2 and C3-S2 with minor plots in C3-S1 in US Salinity diagram (1954), which indicate groundwater is of medium to high salinity-medium sodium type (Figure 6) and thus can be utilized for irrigation with little danger of exchangeable sodium.

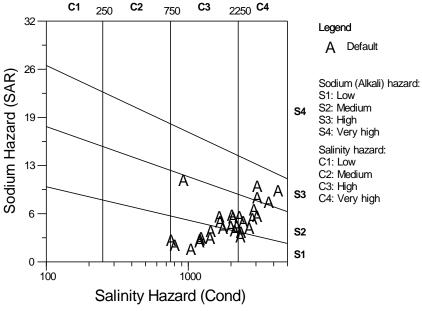


Figure 6: US salinity diagram of groundwater samples from study area

Percent Sodium (%Na): The % Na is widely used for evaluating the suitability of water quality for irrigation (Wilcox, 1995). It is defined by the expression: %Na= Na⁺+K⁺/(Ca⁺⁺+Mg⁺⁺+Na⁺+K⁺) x 100. High % Na in irrigation water causes exchange of sodium in water, and exchange of calcium and magnesium contents in soil having poor internal drainage. The %Na in study area ranges from 4.907 to 94.805 meq/l. out of total samples 94% samples have the % Na < 60 represents safe water while 6% of samples have > 60 that is unsafe (Eaton, 1950) from study area for irrigation purpose.

Residual Sodium Carbonate (RSC): A high value (in meq/l) in water leads to an increase in the adsorption of sodium on soil (Eaton 1950). It is calculated by following as RSC = $(HCO_3^++CO_3^-)$ $(Ca^{++}+Mg^{++})$. The RSC values < 1.25 meq/l are considered as suitable for the irrigation and in study area only 18% groundwater sample are suitable for irrigation.

Mg Ratio (MR): Magnesium ratio (MR) is calculated as MR = $(Mg^{++} \times 100)/(Ca^{++}+Mg^{++})$. The MR values >50 are unsuitable. Excess amount of magnesium can affect the quality of soil reduces the yield of crops. About 39 % samples from the study area are not suitable for irrigation while 61% samples are suitable for irrigation.

Corrosivity Ratio (CR): Corrosivity Ratio (in mg/l) is calculated by the following formula: CR= [(Cl $(35.5) + 2(SO_4^{-1}/96)]/2(HCO_3^{-1}+CO_3^{-1}/100)$. The CR values ≤ 1 is considered as good and > 1 indicates corrosive nature of water, hence water should not be transported through the metal pipes (Ryner, 1994 and Raman, 1985). In the study area only the 6% samples have CR values ≤ 1 and remaining 94% of samples have CR value >1 which indicates that only 6% water in these areas is good for irrigation.

Conclusion

The groundwater from deeper aquifer of the watershed WRE2 area is alkaline in nature. The higher conductivity in the study area is due to high mineral concentration in groundwater. It indicates that the ion exchange and solublization processes have taken place within the aquifer. The Ca^{++} and HCO_3^{-} as well as Mg^{++} and HCO_3 show positive correlation between these ions. The interrelationship among $Ca^{++} + Mg^{++}$ vs SO₄ + HCO₃ as well as HCO₃ and Cl vs Na exhibits the calcium dissolution and silicate weathering for the source for Na⁺ in study area. The feldspars, calcite and clay minerals are the probable sources of Ca⁺⁺ from sedimentary rocks. The chief sources of Mg⁺⁺ in natural water are the magnesium bearing minerals like pyroxenes, olivine and amphiboles while the feldspar and clay minerals are the probable sources for Na^+ and K^+ . SO_4^- is generally derived from the oxidative weathering of sulphate bearing minerals like pyrite (FeS₂) which is very common secondary minerals in the Gondwana formations and associated sediments. The F⁻ concentration in groundwater depends upon the degree of weathering and leaching of fluoride bearing minerals from the rocks and soils. The 27% groundwater samples from the study area are fresh water and can be suitable for drinking and domestic use. The major plots of groundwater sample in C4-S2 and C3-S2 with minor plots in C3-S1 class of US salinity diagram indicate groundwater is of medium to high salinity-medium sodium type and thus can be utilized for irrigation with little danger of exchangeable sodium.

REFERENCES

Agrawal V and Jagetia M (1997). Hydrogeochemical assessment of groundwater quality in Udaypur city Rajasthan, India. In: Proceeding on national conference on dimension of environmental stress in India, University of Baradoa, India 151-154.

Allen SKJM and Lucas S (1996). Concentration of containments in surface water samples collected in west-central impacted by acid mine drainage. *Environmental Geology* 27 34-37.

APHA (1995). Standard Method for Examination of the Water and Waste Water, 19th edition, Americal Health Association, Washington DC, USA.

BIS (1991). Bureau of Indian Standard specification for drinking water. IS: 10500, Indian Standard Institute 1-5.

Canter LW (1997). Nitrates in Groundwater (Lewis publisher) New York.

Carrol D (1962). Rainwater as chemical agent of geologic process-a review. USGS water supply paper 1535-G 18.

Durvey VS, Sharma LL, Saini VP and Sharma BK (1991). Handbook on methodology of water quality assessment. Rajasthan Agricultural University, India 156.

Eaton FM (1950). Significance of carbonates in irrigation waters. Soil Science 39 23-133.

Gibbs RJ (1970). Mechanism controlling world water chemistry. Science 17 1088-1090.

GSI (2008). Geology and Mineral Resources of Maharashtra, Geological Survey of India (Miscellaneous publication) **30**(II) 120.

Handa BK (1975). Geochemistry and genesis of fluoride containing groundwater in India. *Groundwater* 13(3) 275-281.

Hem JD (1970). Study and interpretation of the chemical characteristics of the natural water. USGS Water Supply Paper 1459 286.

Jeong CH (2001). Effects of landuse and urbanisation on hydrochemistry and contamination of groundwater from Taejon area, Korea. *Journal of Hydrology* 253 194-210.

Loizidou M and Kapetanious EG (1993). Effect of leachate from landfills on underground water quality. *Science of the Total Environment* 128 69-81.

Madhnure P and Malpe DB (2007). Flouride contamination of groundwater in rural parts of Yavatmal District, Maharashtra: Causes and Remedies. *Gond. Geol. Mag. Spl. Pub.* **11** 127-135.

Marghade D, Malpe DB and Zade AB (2010). Geochemical characterization of groundwater from north eastern part of Nagpur urban, Central India. *Environmental Earth Sciences*.

Meybeck (1987). Global chemical weathering of surfacial rocks estimated from river dissolved loads. *American Journal of Science* 287 401-428.

Murkute YA and Badhan PP (2011). Fluoride contamination in groundwater from Bhadravati tahsil, Chandrapur District, Maharashtra. *Nature Environment and Pollution Technology* **10**(2) 255-260.

Murkute YA (2014). Hydrogeochemical Characterization and Quality Assessment of Groundwater around Umrer Coal Mine Area Nagpur District, Maharashtra, India. *Environmental Earth Sciences*.

Piper AM (1953). A graphical procedure in the geochemical interpretation of water analysis. *Transactions - American Geophysical Union* **25** 914-923.

Raman V (1985). Impact of corrosion in the conveyance and distribution of water. *Journal of the American Water Resources Association* **15**(11) 115-121.

Ramesam V and Rajgopalan K (1985). Fluoride ingestion into the natural waters of hard-rock areas of peninsular India. *Journal of Geological Society of India* 26 125-132.

Ravikumar P, Venkatesharaju K, Prakash KL and Somashekhar RK (2010). Geochemistry of groundwater and groundwater prospects evaluation, Anekal Taluk, Banbalore urban district, Karnataka, India. *Environmental Monitoring and Assessment*.

Richards LA (1954). Diagnosis and improvement of saline and alkali soils, US department of agricultural handbool. Washington DC, USA.

Sanhez-Perez JM and Tremolieres M (2003). Change in groundwater chemistry as a consequences of suppression of floods: The case of Rhine floodplains. *Journal of Hydrology* 270 89-104.

Sawyer GN and McCarty DL (1967). Chemistry of Sanitary Engineers (McGraw Hill) New York.

Todd DK (1995). *Groundwater Hydrology* (John Wiley and Sons) Singapore New York Chichester Brisbane Toronto.

US Salinity Laboratory Staff (1954). Diagnosis and improvement of saline and alkali soils, U.S. Dept Agriculture. Agriculture Handbook.

WHO (1997). Guidlines for drinking water quality. Geneva World health organization.

Wilcox LV (1955). The quality of water for irrigation use, U.S. Department Agriculture, *Technical Bulletin* 962 40.