

HYDROCHEMICAL CHARACTERISTICS AND WATER QUALITY ASSESSMENT IN ABU-ZAABAL AREA, EASTERN NILE DELTA, EGYPT

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ABSTRACT

Objective: The study presents simple tools for water resources quality classification based on its chemical compositions in Abu Zaabal area, eastern Nile Delta, Egypt and assess the water quality for different uses.

Methods: 31 water samples were collected from different water resources in the study area and analyzed for physicochemical parameters. Hydrochemical relations, contour maps and statistical methods were used to estimate the contamination indices and evaluate the water resources for different purposes.

Results: 83.3% of groundwater samples is fresh water and 16.7% are brackish water. 85.7% of surface water samples are fresh and 14.3% is saline. 92% of groundwater samples and 71.5% of surface water samples are very hard water. According to HPI values, 8% of the quaternary groundwater samples are good, 4% are poor, 4% are very poor and 84% of the samples are unsuitable. All groundwater samples and 71% of surface water samples are contaminated with respect to ammonia.

Conclusion: Higher concentrations of TDS and heavy metal may be due to the clay nature of the soil, the marine sediments in the aquifer matrix together with the dissolution and leaching of minerals from agricultural, anthropogenic and industrial activities. The groundwater in the polluted zones is considered unsuitable for human drinking.

Keywords: Water resources, Hydrochemistry, Water quality indices

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INTRODUCTION

Surface and Groundwater have an associated hydrological relationship affected by different factors related to geological; hydrological and climatic conditions where these factors control the circumstances of groundwater movement in shallow aquifers as well as the quantity of water can be gained or lost from the aquifer and river [1].

Due to industrial and agricultural activities, large amounts of untreated urban, industrial wastewater and rural household waste discharge into the Nile River, canals or agricultural drain, which become an easy dumping site for all types of wastes [2]. Ismailia Canal is the most distal downstream of the principal Nile River. And the water contains all the toxins that are discharged into the Nile. The Ismailia Canal has many pollution sources which potentially affect and deteriorate the canal's water quality [3]. Heavy metals are considered to be a serious pollution of aquatic ecosystems due to

their environmental persistence and toxicity effects on living organisms [4]. In the aquatic environment, trace elements are partitioned between different environmental components (water, suspended solids, sediments and biota [5]. Water resources chemistry is due to long-term interaction between the water systems and the surrounding environment, which can indicate the water formation and migration [6, 7].

The objective of this present study is to highlight the chemical compositions of different water resources in Abu Zaabal area, Eastern Nile Delta, Egypt. Assess the water quality for different uses.

The study area lies in the eastern portion of the River Nile delta in Qalyoubiya governorate, northeast Cairo city bounded by longitudes 31.320 and 31.440 E and latitudes 30.240 and 30.320 N, (fig. 1) occupies about 20 km². The study area is bounded by Cairo ring road from the North, Belbis city from the south, Shebin El Qanater city from the west and Cairo–Belbeis desert road from the east.

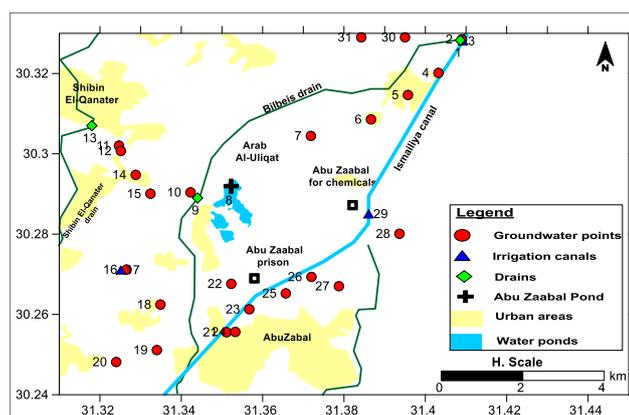


Fig. 1: The location map and sampling points of the area under investigation

Abu Zaabal is considered as a plain area with an average elevation of 27 m above the mean sea level [8]. The area under investigation is characterized by cultivating lands surrounded by urban localities. The urban area is served by freshwater pipelines coming from a Mostorod water station is situated in the northern part of the study area as well as many shallow-private wells have been drilled for water extraction.

Geologically The Pleistocene and Holocene quaternary deposits cover most of the study area; the Basaltic rocks belonging to an Upper Oligocene age are exposed at Abu Zaabal area, while the Pliocene and Miocene sediments outcrops at the eastern part of Ismailia canal. The Holocene Nile silt and clay cover the majority of the study area with different thickness varies from 0 to 20 m, the sand dune unites belongs to the Holocene age found in the eastern part of the study area.

The hydrological conditions and the groundwater aquifers of the eastern portion of the Nile delta were discussed by many authors [9-13].

The surface water infrastructure in the study area consists of a network of the surface water system (Ismailia canal, Belbies drain and Shebin El Qanater drain). The surface water systems are passing through Holocene deposits (Nile silt and clay deposits) and the Pleistocene sediments after the disappearance of the Holocene deposits. The contaminated liquids are directly discharged into canals, drain and on the land surface. The Pleistocene aquifer is influenced by the contaminated water infiltrates due to the small thickness of the clay cap.

The Quaternary aquifers are discriminated into the upper unit (Holocene aquitard) and the lower one (the Pleistocene aquifer) [11, 13 and 14]. The Pleistocene aquifer is overlain by the Holocene unit and underlain by the Pliocene clay in the majority of the area. Around Abu Zaabal Quarries, it is underlain by Miocene sediments or the Oligocene Basaltic sheet. The Holocene aquifer is composed of the Nile silt and clay, with thickness ranges between 0 m at the eastern portions of 20 m at the southwestern part of the study area. The Pleistocene aquifer consists of sand and gravel with clay lenses with thickness ranges between 0 to nearly 50 m, while at the northwestern part of the investigated area, they may reach 200 m. The groundwater movements in the Pleistocene aquifer are mainly due north and northwest reflected that Ismailia canal is the main recharging source as the surface water level in the canal is higher than the groundwater level. Besides, the recharges from irrigation canals and return flow after irrigation. Septic tanks and sewer systems are considered a local source of recharge. The main discharge of the Pleistocene aquifer takes place artificially through pumping wells used for irrigation and domestic uses.

MATERIALS AND METHODS

Sampling procedures

Thirty-one water samples were collected from different water resources (24 samples from groundwater wells and 7 samples represents surface water systems) in August 2019; Surface water samples were collected using an autosampler and polyvinyl chloride Van Dorn bottle

Field measurements

The location (longitudes and latitudes) of the water points was recorded using global positioning system (GPS) model etrex 10 (Germany).

Water samples were collected in a 1000 ml clean polyethylene bottle which was used for major ions measurements, whereas a 50 ml clean polyethylene bottles was acidified with concentrated HNO₃ to pH<2 for heavy metals detections. E. C and pH were measured in situ using portable meters (AD 310 and 3510, Jenway, UK).

Laboratory measurements

The chemistry of water samples was detected in Hydrogeochemistry laboratories, Desert Research Center, Cairo, Egypt. The measuring of the major, minor constituents of the water samples (total dissolved solids (TDS), major ions as Ca²⁺, Mg²⁺, Na⁺, K⁺, CO₃²⁻, HCO₃⁻, SO₄²⁻, Cl⁻, NH₄⁺, NO₂⁻, NO₃⁻ and PO₄³⁻) were carried out according to the methods adopted by Bufflap SE and Allen HE (1995), Onken and Sunderman (1977), Fishman and Friedman (1985), Barer et al. (2000) and American Public Health Association (2005) [15-19] table 1.

Table 1: Methods adopted for water quality analysis

Quality parameter	Method used
PH	Potentiometric (1:2.5 H ₂ O, v/v)
Electrical Conductivity EC	Conductometry (1:2.5 H ₂ O, v/v)
Calcium Ca ²⁺	EDTA (0.05 N) titrimetric
Magnesium Mg ²⁺	EDTA (0.05 N) titrimetric
Sodium Na ⁺	Flame photometric
Potassium K ⁺	Flame photometric
Chloride Cl ⁻	Titration using 0.05 N AgNO ₃
Carbonate CO ₃ ²⁻	Titration (with 0.01 N H ₂ SO ₄)
Bicarbonate HCO ₃ ⁻	Titration (with 0.01 N H ₂ SO ₄)
Sulphate SO ₄ ²⁻	Spectrophotometric

After physiochemical analysis, the accuracy of the analysis results (% Balance error (%E)) was checked. Generally speaking, the relative error should be within±5%.

Heavy metals and trace components (Al, B, Cd, Co, Cr, Cu, Fe, Pb, Mn, Mo, Ni, Sr, V and Zn) were detected by plasma optical emission mass spectrometer (ICP) (POEMSIII, thermo Jarrell elemental company USA), using 1000 mg/l (Merck) Stock solution for standard preparation. The water quality parameters were estimated to evaluate the water resources in the study area (tables 2, 3).

RESULTS AND DISCUSSION

Physicochemical parameters of water resources

The physical and chemical analyses of water samples in Abu Zaabal area are summarized in (tables 4, 5).

Hydrogen ion concentration (pH)

The pH value reflects the acidic or alkaline material present in the water. The decrease of pH less than 7 reflects an increase in hydrogen ion concentration. Where the increase in pH more than 7 is reflects an increase in the hydroxyl ion. In the study area, the pH values range from 7.8 to 8.6 and from 8.0 to 8.7 for the ground and surface water, respectively, which indicates that the water resources in the study area are generally alkaline in nature.

Table 2: Water quality parameter estimation methods from measured parameters

Quality parameters	Formula adopted	Reference/source
Total dissolved solids (TDS)	$TDS = (Ca^{2+} + Mg^{2+} + Na^{+} + K^{+} + CO_3 + (HCO_3/2) + SO_4^{2-} + Cl^{-})$	[20]
Total hardness (TH)	$TH = (Ca + Mg) \times 50$	[21]
Heavy metal pollution index (HPI)	$HPI = (\sum W_i \times Q_i) / \sum W_i$ (1) W _i is the unit weightage of the heavy metal (i), n is the number of heavy metals, Q _i is the sub-index of the heavy metal. $W_i = \frac{K}{S_i}$ (2) K is the proportionality constant; S _i is the standard permissible limit of the heavy metal. $K = \frac{1}{\sum_{i=1}^n \frac{1}{S_i}} = \frac{1}{\frac{1}{S_1} + \frac{1}{S_2} + \dots + \frac{1}{S_n}}$ (3) Where, S ₁ , S ₂ , S ₃ , and S _i represent standards for different heavy metals in the groundwater samples. $Q_i = 100 \times \frac{V_i}{S_i}$ (4) V _i is the monitored value of the i parameter in mg/l, HPI is classified into five classes, excellent (0–25), good (26–50), poor (51–75), very poor (76–100) and unsuitable (100).	[22, 23]

Quality parameters	Formula adopted	Reference/source
Nitrate pollution index (NPI)	$NPI = \frac{Cs-HAV}{HAV}$ Where Cs: The analytical concentration of nitrate. HAV: The threshold value of anthropogenic source (human affected value) taken as 20 mg/l. The water quality according to NPI values was classified into five types: clean (unpolluted)(NPI<0), light pollution (0<NPI<1), moderate pollution (1<NPI<2), significant pollution (2<NPI<3), very significant pollution (NPI>3).	[24]
Drinking water quality index (DWQI)	The relative weight (Wi) is computed from the following equation: $Wi = \frac{wi}{\sum_{i=1}^n wi}$ where Wi is the relative weight wi is the weight of each parameter n is the number of parameters $qi = \frac{Ci}{Si} \times 100$ where qi is the quality rating Ci is the concentration of each chemical parameter in each water sample in milligrams per liter Si is the Egyptian drinking water standard for each chemical parameter in milligrams per liter according to the guidelines of the (Egyptian Higher Committee, 2007; WHO, 2011). For computing the WQI, the SI is first determined for each chemical parameter, which is then used to determine the WQI as per the following equation $Sli = Wi \times qi$ $WQI = \sum Sli$ where Sli is the sub-index of ith parameter qi is the rating based on the concentration of ith parameter n is the number of parameters The standard is the standard of the water quality parameter. The water samples were classified according to WQI rate as excellent, good, poor, very poor and unfit for human consumption (table 4).	[25]
Sodium Adsorption Ratio (SAR)	$SAR = \frac{Na}{\sqrt{(Ca + Mg)/2}}$	[26]
Residual Sodium Carbonate (RSC)	$RSC = (CO_3^{2-} + HCO_3^-) - (Ca^{2+} + Mg^{2+})$	[27]
Sodium percentage (Na%)	$\%Na = \frac{[Na+K^+]}{[Na+K^++Ca^{2+}+Mg^{2+}]} \times 100$	[28]
Magnesium ratio (MAR)	$MAR = \frac{[Mg^{2+}]}{[Ca^{2+}+Mg^{2+}]} \times 100$	[29]
% Balance error (%E)	$\%E = \frac{[\sum cation - anion]}{[\sum cation + anion]} \times 100$	[20]

Table 3: Water quality parameters, their standard values, their ideal values and the assigned weighting factors

Parameter	Standard value (Si)	Weight (wi)	Realative weight Wi
TDS	1000	5	0.161
Ca2+	200	3	0.097
Mg2+	150	3	0.097
Na+	200	4	0.129
HCO3-	500*	1	0.032
SO42-	250	5	0.161
Cl-	250	5	0.161
NO3	45	5	0.161
		$\sum = 31$	$\sum = 1$
The values according to Egyptian standards (2007) and WHO (2011) [30, 31]			$\sum = 0.146$

Total dissolved solids (TDS)

The water salinity of groundwater ranges of 243 mg/l to 3390 mg/l and in surface water of 240 mg/l to 5600 mg/l, as shown in (fig. 2). 83.3% of groundwater samples are fresh water and 16.7% are brackish water. 85.7% from surface water samples are fresh and 14.3% is saline. Higher concentrations of TDS may be credited to the impact of evaporation and the marine sediments in the aquifer matrix together with the dissolution and leaching of minerals from agricultural, anthropogenic and industrial activities [32, 33].

Total hardness

The total hardness (TH) is caused primarily by the presence of cations such as calcium and magnesium and anions such as carbonate, bicarbonate, chloride and sulfate in water. The total hardness values of groundwater samples range from 150 mg/l to 1300 mg/l reflected that 8% of these samples are hard and 92% are very hard water. The total hardness values in surface water range from 140 mg/l to 1480 mg/l reflected that 28.5% of samples are hard and 71.5% are very hard.

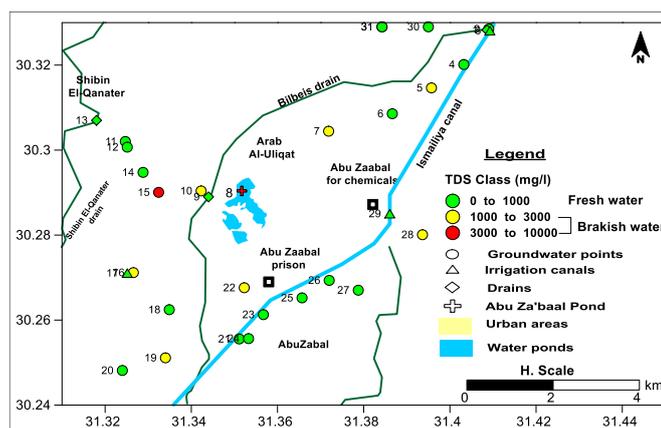


Fig. 2: TDS classification map for the quaternary groundwater in the study area

Soluble anions

Bicarbonate ion (HCO_3^-) source is from the dissolution of carbonate rocks (dolomite, limestone, magnesites etc.). HCO_3^- is mainly formed due to the action of CO_2 from the atmosphere and that released from organic decomposition [34, 35]. Bicarbonate concentration in groundwater of the Quaternary (Pleistocene) aquifer varies from 138 mg/l to 520 mg/l and in surface water from 134 mg/l to 420 mg/l. The bicarbonate distribution in groundwater indicates that high content and the presence of local variations advocates the existence of local pollution sources. The distribution of bicarbonate salts increased from west to east. This direction may be due to the recharge of the quaternary aquifer from the Ismailia canal (fig. 3. a).

Sulfate ion (SO_4^{2-}) is naturally formed due to rock weathering, input from volcanoes and biochemical process [36]. The oxidation and decomposition of substances containing sulfur (fossil fuels and dissolution of sulfur-bearing minerals such as gypsum and pyrite) and anthropogenic activities are other sources of SO_4 ions [35]. The sulfate

content in groundwater of the Quaternary aquifer varies from 23.1 mg/l to 780 mg/l and from 28.9 mg/l to 1080 mg/l in surface water. The groundwater distribution of sulfate indicates the presence of local zones of high concentrations at Abu Zaabal, reflecting that the effect of the saline pond from the west and the influence of the sulfate fertilizers in the new reclaimed land in the east (fig. 3. b).

The Cl-ion form in nature is usually of chlorine salts (CaCl_2 , MgCl_2 and NaCl). The main source is due to the leaching and dissolution of sedimentary rocks; common evaporates minerals and saline deposits. Industrial, municipal wastes and irrigated agricultural activities are other main sources of chloride salts [37]. The chloride content varies from 32 mg/l to 970 mg/l in the quaternary groundwater samples and from 35 mg/l to 1750 mg/l in surface water. The chloride content distribution in groundwater shows the presence of local zones of high concentrations at Abu Zaabal. The local variations in the chloride concentrations are attributed to local recharge from the saline ponds in the study area; this also confirms the existence of local pollution sources (fig. 3. c).

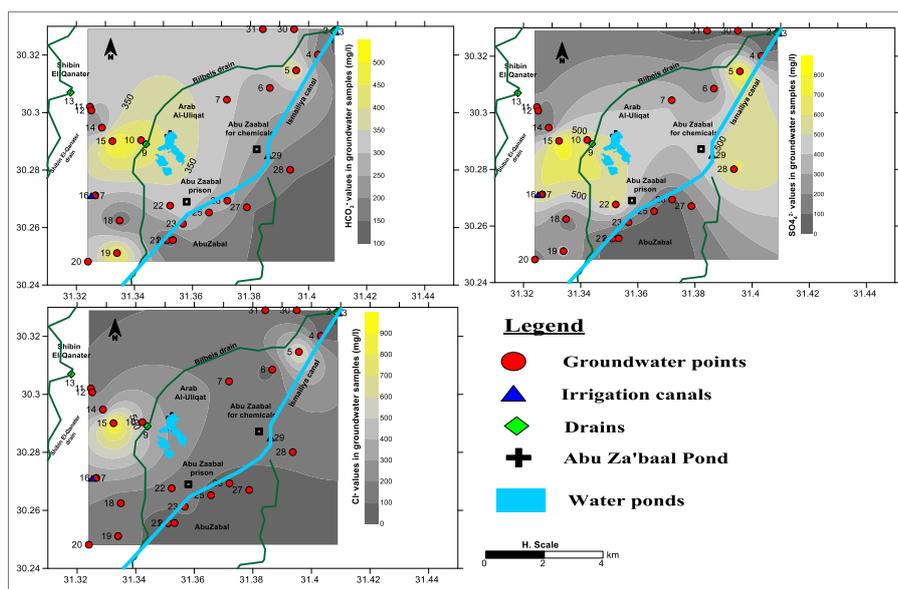


Fig. 3: Spatial distribution map of anions concentrations in the quaternary groundwater in the study area

Soluble cations

Calcium plays an important role in the health of water bodies which reduces the toxicity of chemical compounds in natural water [38]. The removal of Ca^{2+} ion from the water resources is due to an ion-exchange or calcite (CaCO_3) precipitation. Calcite precipitation occurs when CO_2 content is low, causing the chemical reaction process in the reverse direction [35]. The calcium content in groundwater of the Quaternary aquifer varies from 32.8 mg/l to 392 mg/l and from 32.8 mg/l to 384 mg/l in the surface water samples. The calcium distribution in groundwater confirms the presence of local zones of high concentrations that occurred at Abu Zaabal (fig. 4. a). Hardness of water is attributed to the presence of calcium and magnesium ions; the water in the study area varied from hard to very hard.

A magnesium source water resources is due to chemical weathering and dissolution of dolomite, marls and other rocks [39]. Magnesium content in the Quaternary aquifer samples varies from 13.92 mg/l to 114 mg/l and from 11.5 mg/l to 124.8 mg/l in surface water. The magnesium distribution in groundwater shows the presence of local zones of high concentrations, but the magnesium contents are still below the excessive limits for drinking (fig. 4. b).

A sodium source in the water resources is due to weathering of Na bearing minerals/rocks (halite, feldspar and montmorillonite), cation-exchange process (displacement from absorbing complex of

rocks and soils by Ca and Mg), and anthropogenic activities (pollution from industrial effluent, domestic sewage, and agricultural activities). Sodium content in groundwater of the Quaternary (Pleistocene) aquifer varies from 18 mg/l to 442.9 mg/l and from 20 mg/l to 1265 mg/l in surface water. The distribution of sodium ions in the study area reflects local variations may be attributed to local recharge from the saline ponds in the study area. The groundwater in the polluted zones is considered unsuitable for drinking (fig. 4. c). Potassium is slightly less common than sodium in igneous rocks, but more abundant in all sedimentary rocks. In igneous rocks, potassium is present as feldspars (orthoclase and microcline (KAlSi_3O_8)), wherein sediments it is present in clay minerals. Potassium is slightly less common than sodium in igneous rocks, but more abundant in all sedimentary rocks. In igneous rocks, potassium is present as feldspres (orthoclase and microcline (KAlSi_3O_8)), wherein sediments it is present in clay minerals. The concentration of potassium in natural water is generally less than 10 mg/l as much as 100 mg/l in hot springs and about 25000 mg/l in brines.

Minor, trace and heavy metals

Nitrate concentrations in the groundwater samples ranges between 12 mg/l to 42 mg/l and from 8 mg/l to 75 mg/l in the surface water samples. Nitrite concentration in the groundwater samples ranged between 0.05 mg/l to 0.51 mg/l and from 0.01 mg/l to 0.61 mg/l in the surface water samples. Ammonia concentration in the groundwater samples ranges between 0.5 mg/l to 3.7 mg/l and from

0.1 mg/l to 8 mg/l in the surface water samples. From the previous data, the groundwater samples are contaminated with ammonia. This shows that groundwater samples is mixed with sewage and the presence of *Escherichia coli* bacteria from bacteriological analysis of some groundwater samples proved that.

Iron content, 83.3% of groundwater samples in the Pleistocene aquifers of Abu Zaabal area are unsuitable for human drinking, while the rest of the samples (16.7%) are suitable for drinking. On the other hand, 71.4% of surface water samples are unsuitable for human drinking, while the rest of the samples (28.6%) are suitable for drinking. Iron values of groundwater ranges from 0.004 mg/l to

5.39 mg/l and in surface water from 0.03 mg/l to 6.96 mg/l. This is due to the clay nature of the soil.

Manganese content (33.3%) of the groundwater samples and 28.6% of surface water samples are unsuitable for drinking. Lead content (12.5%) of the groundwater samples is unsuitable for drinking. Cadmium content, (83.3%) of the groundwater samples and 71.4% of surface water samples are unsuitable for drinking. Aluminum content (8.3%) of the groundwater samples and 71.4% of surface water samples are unsuitable for drinking. Nickel content (12.5%) of the groundwater samples and 42.8% of surface water samples are unsuitable for drinking.

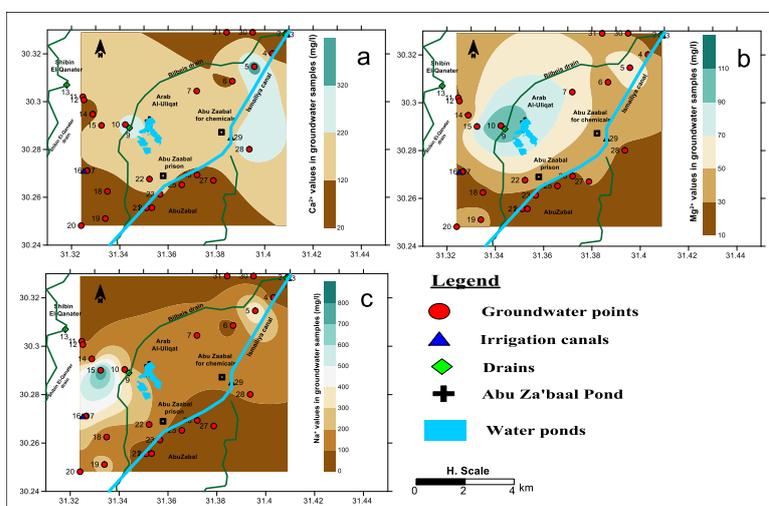


Fig. 4: Spatial distribution map of cations concentrations in the quaternary groundwater in the study area

Table 4: Major and minor element concentrations of water samples in Abu Zaabal area

Sam ple	pH	EC (µS/cm)	TDS (mg/l)	T. Hardness	Ca ²⁺ (mg/l)	Mg ²⁺ (mg/l)	Na ⁺ (mg/l)	CO ₃ ²⁻ (mg/l)	HCO ₃ ⁻ (mg/l)	SO ₄ ²⁻ (mg/l)	Cl ⁻ (mg/l)	%E	NH ₄ ⁺ (mg/l)	NO ₂ ⁻ (mg/l)	NO ₃ ⁻ (mg/l)	PO ₄ ⁻ (mg/l)
Surface water																
1S	8	1375	880	355	82	36	130	8	220	170.2	225	-3.0	8	0.61	75	5.5
3S	8.1	375	240	140	36.8	11.5	20	2	136	29.3	37	-3.0	0.1	0.01	10	0
8S	8.1	8750	5600	1480	384	124.8	1265	10	420	1080	1750	3.5	5.5	0.6	70	18
9S	8.2	1390.6	890	360	88	33.6	125	9	223	171.1	218.7	-3.1	6.3	0.61	58	24
13S	8.7	806.3	516	239	62	20.2	76.7	2	134	108	156	-4.5	0.9	0.12	28	7.4
17S	8.5	406.3	260	196	52	15.8	21.5	4	140	32.9	58	2.3	0.9	0.14	14	3.8
29S	8.1	400	256	156	32.8	17.8	21.7	2	146	28.9	35	0.7	0.1	0.01	8	0.01
Groundwater																
2G	8.5	392.2	251	161	32.85	18.9	18.2	4	138	32.3	32	2.0	0.9	0.05	16	2.7
4G	8.3	695.3	445	290	61.6	32.6	35	4	172	110.1	53	4.9	0.8	0.24	14	0.01
5G	7.8	4312.5	2760	1300	392	76.8	442.9	6	396	820	680	2.7	3.7	0.51	42	0.08
6G	7.8	1073.4	687	400	104	33.6	64	4	258	183.8	65	4.1	0.9	0.13	24	0.18
7G	7.9	1995.3	1277	643	144	68	170	4	316	392.4	187.8	3.9	1.1	0.15	25	0.27
10G	7.7	3031.3	1940	1075	240	114	225	4	440	547.1	395.8	2.3	1.5	0.19	26	0.01
11G	8.3	1378.1	882	525	164.8	27.1	95	8	318	217.5	131	4.1	1.4	0.17	25	0.05
12G	8	1315.6	842	436	96	47	115	6	234	158	200	3.4	0.7	0.08	19	0.01
14G	8	1226.6	785	346	96	25.7	115	2	240	98.3	238	-3.2	0.9	0.15	24	0.07
15G	7.7	5296.9	3390	780	172	84	860	8	520	780	970	0.7	2.3	0.51	30	0.03
16G	8.3	2625	1680	383	112	24.8	420	7	312	592.8	312	-0.7	1.1	0.16	19	0.27
18G	8.3	875	560	240	53.6	25.4	83.3	6	150	134.8	123	-2.0	1.8	0.22	28	0.26
19G	8.2	1953.1	1250	394	99.2	35	260	6	480	340.8	174	-1.9	1.2	0.15	24	0.37
20G	8.4	1054.7	675	336	76.8	34.6	116	4	292	140.8	120	2.7	1	0.13	18	0.05
21G	8.6	412.5	264	152	37.6	13.9	25	4	138	42	38.7	-1.4	1.6	0.09	27	0.01
22G	7.9	2229.7	1427	662.5	194	42.6	190	4	322	620.9	156	-2.6	1.1	0.16	23	0.01
23G	8	1129.7	723	422.5	130	23.4	80	4	298	140.6	136	1.0	0.9	0.09	18	0.028
24G	8.2	379.7	243	150	32.8	16.3	24.6	2	144	22.5	52.4	-3.1	0.6	0.08	16	0.01
25G	7.9	812.5	520	294	78.8	23.3	46.7	8	249	86.3	65.7	0.9	1.3	0.16	25	0.05
26G	7.8	721.9	462	366	102.4	26.4	35	4	280	69.2	72.9	4.2	0.5	0.07	12	0.01
27G	7.8	837.5	536	313	72	32	50	4	158	170.7	76	0.7	1.2	0.13	25	0.05
28G	8.1	1937.5	1240	760	258.4	27.4	110	4	162	620.6	117	2.7	2.5	0.17	26	0.01
30G	8.2	864.1	553	346.6	92	28	40	4	318	34.8	68	4.7	0.8	0.21	21	0.01
31G	8.1	760.9	487	356	72.8	41.8	42.9	8	296	23.1	103	4.0	2.5	0.15	32	0.01

Table 5: Trace and heavy metal concentrations of water samples in Abu Zaabal area in mg/l unit

Sample	Al	Co	Cu	Cd	Fe	Pb	Sr	Mn	Mo	Ni	Ba
Surface water											
1S	1.29	0.035	0.050	0.042	2.368	<0.008	1.556	0.192	0.090	0.082	0.092
3S	1.26	0.097	<0.006	<0.0006	0.030	<0.008	0.400	0.090	0.107	<0.002	0.071
8S	14.29	<0.001	0.065	0.047	6.960	<0.008	12.290	0.819	0.036	<0.002	0.097
9S	1.04	<0.001	0.078	0.027	2.836	<0.008	1.578	0.158	<0.001	0.072	0.098
13S	2.01	0.007	0.039	0.028	1.430	<0.008	0.584	0.118	<0.001	<0.002	0.067
17S	0.01	<0.001	<0.006	0.015	1.827	<0.008	1.444	0.661	0.075	0.028	0.041
29S	0.15	0.010	0.034	<0.0006	0.061	<0.008	0.599	<0.002	0.025	<0.002	0.082
Groundwater											
2G	0.10	0.023	<0.006	0.028	0.695	<0.008	0.348	0.035	0.043	<0.002	0.0720
4G	<0.01	<0.001	<0.006	0.029	0.495	<0.008	0.645	0.192	0.027	<0.002	0.070
5G	0.08	0.008	0.006	0.050	0.463	<0.008	3.808	0.902	0.051	0.020	0.070
6G	0.16	0.010	0.022	0.026	0.518	0.052	1.909	0.421	0.003	0.011	0.069
7G	0.12	<0.001	0.012	0.009	0.445	<0.008	2.954	0.016	0.065	<0.002	0.060
10G	0.06	0.003	0.020	0.027	4.730	0.019	2.282	0.848	<0.001	0.028	0.135
11G	0.00	0.000	<0.006	0.024	3.980	<0.008	1.281	0.509	0.018	0.014	0.131
12G	0.11	<0.001	<0.006	0.025	0.500	<0.008	1.433	0.200	<0.001	0.010	0.111
14G	0.0738	0.033	0.021	0.009	0.304	<0.008	0.857	0.206	0.112	<0.002	0.030
15G	<0.01	0.039	<0.006	0.045	0.159	0.156	4.524	0.843	0.064	0.035	0.0469
16G	0.03	0.043	0.019	0.003	5.390	<0.008	4.795	0.921	0.051	<0.002	0.049
18G	0.33	0.072	<0.006	<0.0006	2.220	<0.008	0.332	0.041	<0.001	<0.002	0.034
19G	<0.01	0.021	0.014	0.011	2.190	<0.008	1.394	0.115	0.003	0.013	0.138
20G	0.16	<0.001	0.001	0.018	0.940	<0.008	0.463	0.044	0.061	<0.002	0.067
21G	0.05	0.001	0.057	0.056	0.218	<0.008	7.011	0.221	<0.001	<0.002	0.144
22G	<0.01	<0.001	<0.006	0.038	1.277	<0.008	1.564	0.139	0.000	<0.002	0.497
23G	0.04	0.062	<0.006	0.025	0.777	<0.008	0.337	0.102	<0.001	<0.002	0.073
24G	<0.01	0.049	0.024	0.011	0.028	<0.008	1.315	0.008	<0.001	<0.002	0.089
25G	<0.01	0.021	0.032	0.037	0.004	<0.008	0.728	0.088	<0.001	0.015	0.143
26G	0.18	0.047	0.085	0.071	0.400	<0.008	1.112	0.881	0.159	<0.002	0.307
27G	<0.01	<0.001	0.001	<0.0006	2.670	<0.008	4.574	0.061	0.067	0.054	0.083
28G	1.20	0.019	0.034	0.070	2.259	<0.008	0.339	0.108	0.171	0.010	0.041
30G	0.07	<0.001	<0.006	<0.0006	1.151	<0.008	1.243	0.189	<0.001	0.015	0.080
31G	0.18	0.008	0.004	0.023	0.436	<0.008	0.400	0.726	0.040	<0.002	0.078

Water resources contamination indices

Nitrate pollution index

The source of nitrate in the groundwater is classified to nonpoint sources such as intensive agricultural activities and point sources such as irrigation of land by sewage effluents [40]. The surface water samples in the Abu Zaabal area are classified according to NPI values as follows: 43 % of the samples are cleaned (unpolluted), 14% are light-polluted, 14% of the samples are moderately polluted and 28% are significant pollution. Where the groundwater samples are 33% of samples are clean (unpolluted), 63% are samples are light polluted and 4% of the samples are moderate polluted table 6. The distribution of the NPI values presented that the majority of the study area located under light-polluted zone may be due to the influence of agricultural activities (nitrification of synthetic fertilizers and soil organic nitrogen). Where is the moderate

pollution is located close to Bilbeis drain reflected the influence of groundwater recharge from the drain (fig. 5).

Heavy metal pollution index (HPI)

The Heavy metal pollution index (HPI) for water resources in the study area was calculated based on the concentration of Al, Cu, Cd, Fe, Pb, Mn, Mo and Ni use the permissible limits according to WHO, 2011. HPI of surface water samples ranged between 31.8 and 993.2 reflected wide variation in the surface water resources in the study area (table 6). Ismailia canal samples (3S and 29 S) are classified as good samples according to HPI values. The Quaternary groundwater samples can be classified according to HPI values as: 8% of the quaternary groundwater samples are good, 4% are poor, 4% are very poor and 84% of the samples are unsuitable. The distribution map of HPI values (fig. 6) reflects the increasing of the HPI values in the majority of the study area may be due to wider sources of pollution.

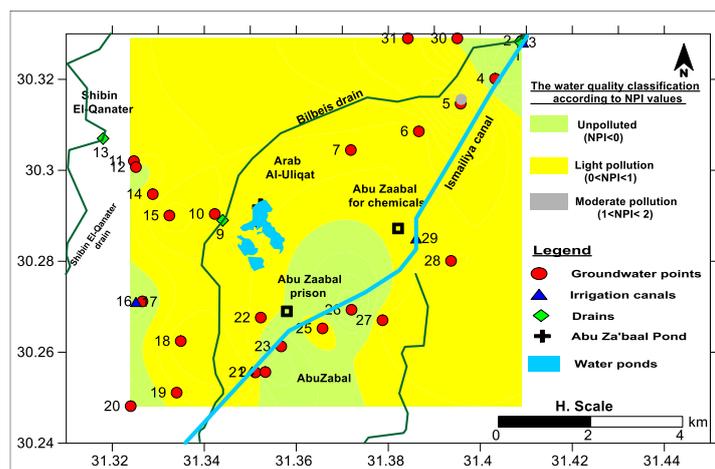


Fig. 5: Spatial distribution of the NPI values for the quaternary water samples in Abu Zaabal area

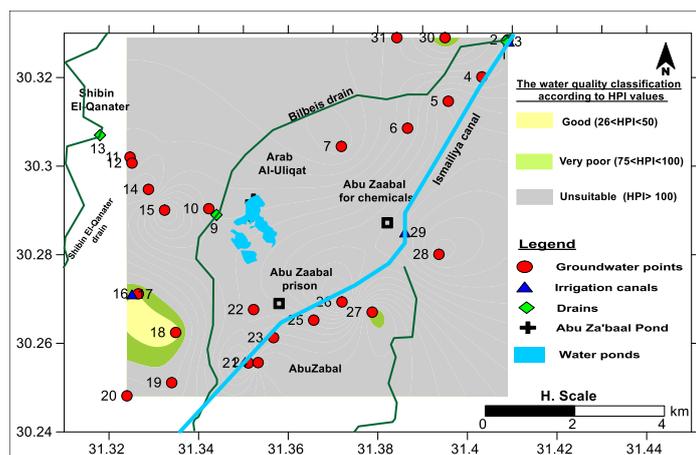


Fig. 6: Spatial distribution of the HPI values for the quaternary water samples in Abu Zaabal area

Table 6: The contamination indices, water quality indices for drinking and evaluation for water resources in the study area

Sample	NPI	HPI	DWQI	SAR	% Na	MR	RSC
Surface water samples							
1S	2.75	993.2	82.6	3	58.5	72.4	-3.2
3S	-0.5	40.7	16.4	0.7	35.8	51.6	-0.5
8S	2.5	1136.2	408.4	14.3	78.4	53.6	-22.2
9S	1.9	655.3	76.1	2.9	62.9	62.9	-3.2
13S	0.4	644.3	45.4	2.2	57.9	53.6	-2.5
17S	-0.3	366.6	20.9	0.7	35.4	50.2	-1.5
29S	-0.6	31.8	16.2	0.8	35.7	89.2	-0.6
Groundwater samples							
2G	-0.2	634.9	18.8	0.6	33.2	95.1	-0.8
4G	-0.3	655.5	31.1	0.9	32.9	87.3	-2.8
5G	1.1	1127.2	211.1	5.4	59	32.3	-19.2
6G	0.2	681.1	48.7	1.4	41.3	53.2	-3.6
7G	0.25	218.4	91.2	2.9	53.8	77.8	-7.5
10G	0.3	655.5	137.6	3	46.7	78.3	-14
11G	0.25	559.3	63.5	1.8	44.6	27.1	-5
12G	-0.05	571.1	60.0	2.4	52.8	80.8	-4.6
14G	0.2	220	58.2	2.7	59.2	44.1	-2.9
15G	0.5	1316.6	250.6	13.4	81.6	80.5	-6.7
16G	-0.05	97.8	128.2	9.4	81.2	36.5	-2.3
18G	0.4	36.5	46.2	2.3	59.6	78.2	-2.1
19G	0.2	268.7	88.8	5.7	74.2	58.2	0.2
20G	-0.1	417.1	49.4	2.8	60.6	74.2	-1.8
21G	0.35	1246.4	24.3	0.9	42	61	-0.6
22G	0.15	853.5	107.7	3.2	54.3	36.2	-7.8
23G	-0.1	567.3	50.8	1.7	43.9	29.7	-3.4
24G	-0.2	258.3	19.6	0.9	39.8	82	-0.6
25G	0.25	835.2	37.0	1.2	39.6	48.7	-1.5
26G	-0.4	1583.7	31.6	0.8	27.9	42.5	-2.6
27G	0.25	64.3	43.3	1.2	40.2	73.2	-3.5
28G	0.3	1574.4	99.2	1.7	37.2	17.5	-12.4
30G	0.05	39.5	33.9	0.9	31.3	50.2	-1.5
31G	0.6	525.7	38.3	1	29.6	94.5	-1.9

Evaluation of groundwater for human drinking

The comparison between the maximum permissible limits major, minor, trace and heavy metals for human drinking (table 7) with the concentrations of these constituents in groundwater and surface water of the investigated samples in the study area leads to the following conclusions:

a) Evaluation of water in human drinking suitability according to physical properties reflected that 100% of water samples (ground and surface) are suitable in respect to pH. 66.7% of groundwater samples and 85.7% of surface water samples are suitable for human drinking in respect to TDS. 70.8% of groundwater samples and 85.7% of surface water samples are suitable for human drinking in respect to TH.

b) Evaluation of water for human drinking suitability according to major constituents reflected that 100%, 91.6%, 66.7%, 83.3% and 79.2% of groundwater samples are suitable for drinking purposes in respect to Mg, Ca, SO₄, Cl and Na, respectively. 100% and 85.7% of surface water samples are suitable for drinking purposes in respect (Mg, Ca) and (SO₄, Cl, Na), respectively.

c) Evaluation of water for human drinking suitability according to minor-trace constituents and heavy metals reflected that:

- 91.7% of groundwater samples and 57.1% of surface samples are suitable in respect to Mo.
- 87.5% of groundwater samples and 57.1% of surface samples are suitable in respect to Ni.

- 16.7% of groundwater samples and 28.6% of surface samples are suitable in respect to Cd.
- 100% of water samples (ground and surface) are suitable in respect to Cu and Zn.
- 87.5%, 91.7%, 100%, 16.7% and 66.7% of groundwater samples and 100%, 28.6%, 42.8%, 28.6%, 71.4% of surface water samples is suitable in respect to Pb, Al, NO₃, Fe and Mn, respectively.
- The evaluation of the water resources for drinking purposes according to quality parameters can be estimated water quality index values (WQI). The water quality index in this study is

calculated according to 8 parameters (TDS, HCO₃, Cl, SO₄, NO₃, Ca, Mg, and Na) has been assigned a weight (wi) according to its relative importance in the overall quality of water for drinking purposes table 3 [25]. The results of the drinking water quality index (DWQI) reflected that 57% of surface water samples are excellent, 28.5% are good and 14.% of samples tapping to Belbis drain are very poor for drinking purposes (table 6 and table 8). The evaluation of the groundwater samples is classified as: 50% of the groundwater samples are excellent, 29% are good, 13% are poor and 8% are very poor. The distribution of drinking water quality index values for the groundwater samples in Abu Zaabal (fig. 7) area reflected the effect of the Belbis drain on the groundwater quality.

Table 7: Water quality guidelines used in evaluation for human drinking water quality index

Parameter	Egyptian1 maximum permissible limit in mg/l (2007) [30]	WHO guidelines for human drinking 2003 [41]
pH-value	6.5-8.5	6.5-9.5
Na (mg/l)	200	200
Mg (mg/l)	150	150
K (mg/l)		12
Ca (mg/l)	350	200
Cl (mg/l)	250	250
SO ₄ (mg/l)	250	250
NO ₃ (mg/l)	45	50
TDS (mg/l)	1000 (at 120 C)	1000
Hardness as CaCO ₃ (mg/l)	500	500
Al (mg/l)	0.2	0.2
Fe (mg/l)	0.3	0.3
Mn (mg/l)	0.4	0.4
Cu (mg/l)	2	2
Zn (mg/l)	3	3
Pb (mg/l)	0.01	0.01
Cr (mg/l)	-	0.05
Cd (mg/l)	0.005	0.003
Ni (mg/l)	0.02	0.02
B (mg/l)	0.5	0.5

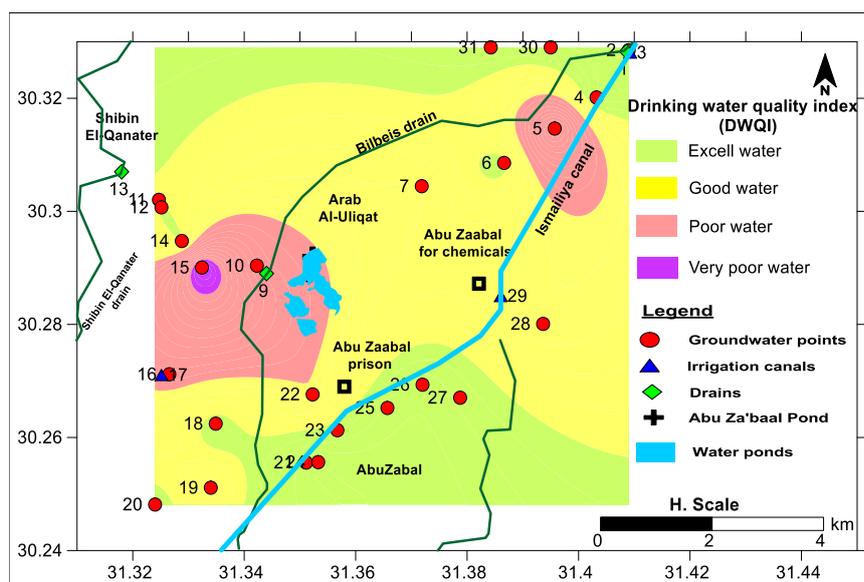


Fig. 7: Spatial distribution of the DWQI values for the quaternary water samples in Abu zaabal area

Table 8: Water quality index scale

Range	Type of water
<50	Excellent
50-100	Good water
100.1-200	Poor water
200.1-300	Very poor water
>300	Water unsuitable for drinking purposes

The evaluation of water resources for irrigation purposes

The suitability of water for irrigation is determined by its mineral constituents and the type of the plant and soil to be irrigated. Water quality used for irrigation is well recognized as an important factor in the productivity of crops. The suitability of water for irrigation is determined not only by the total amount of salt present but also by

the kind of salt. Different chemical factors affecting the suitability of water for irrigation and its effect on crop production and soil quality. Among these are:

- Salinity hazard (EC)-total soluble salt content
- Sodium hazard (SAR)
- Sodium percentage (Na %):

-Magnesium ratio (MR)

-Residual sodium carbonate (RSC)

Salinity hazard (EC)

Based on the EC, irrigation water can be classified into four categories [42] as shown in table 9.

Table 9: Classification of irrigation water based on salinity (EC) values

Level	EC ($\mu\text{S}/\text{cm}$)	Total dissolved salts (mg/l)	Hazard and limitations
C1	<250	<200	Low hazard; no detrimental effects on plants, and no soil buildup expected.
C2	250-750	200-500	Sensitive plants may show stress; moderate leaching prevents salt accumulation in soil.
C3	750-2250	500-1500	Salinity will adversely affect most plants; requires selection of salt-tolerant plants, careful irrigation, good drainage, and leaching.
C4	>2250	>1500	Generally unacceptable for irrigation, except for very salt tolerant plants, excellent drainage, frequent leaching, and intensive management.

Based on this classification, it should be noted that 20.8% of groundwater samples (samples Nos. 2 G, 4G, 21G, 24G and 26G) and 42.85% of surface water samples (samples Nos. 3S, 17S, and 29S) are classified as class C2. 62.5% of groundwater samples (Samples Nos. 6G, 7G, 11G, 12G, 14G, 18G, 19G, 20G, 22G, 23G, 25G, 27G, 28G, 30G and 31G) and 42.85% of surface water samples (Samples Nos. 1S, 9S and 13S,) are classified as class C3 which are saline and then require selection of salt-tolerant plants, careful irrigation, good drainage, and leaching. 16.7% of groundwater samples (samples Nos. 5G, 10G, 15G and 16G) and 14.3% of surface water samples (Sample No. 8S) are classified as class C4.

Sodium adsorption ratio (SAR)

Continued use of water having a high SAR leads to a breakdown in the physical structure of the soil. The sodium replaces calcium and magnesium sorbed on clay minerals and causes dispersion of soil particles. This dispersion results in the breakdown of soil aggregates and causes cementation of the soil under drying conditions as well as preventing infiltration of rainwater. Classification of irrigation water based on SAR values is shown in table 10.

Based on this classification, it should be noted that all samples are classified as class S1 except samples 8S and 15G are classified S2

table 6. 20.8% of groundwater samples (samples Nos.2G, 4G, 21G, 24G and 26G) and 42.85% of surface water samples (samples Nos. 3S, 17S, and 29S) in the study area lie in the fields C2-S1. 62.5% of groundwater samples (Samples Nos. 6G, 7G, 11G, 12G, 14G, 18G, 19G, 20G, 22G, 23G, 25G, 27G, 28G, 30G and 31G) and 42.85% of surface water samples (Samples Nos. 1S, 9S and 13S,) lie in the fields C3-S1. 16.7% of groundwater samples (samples Nos. 5G, 10G and 16G) lie in the fields C4-S1. 4.2% of groundwater samples (samples No. 15G) and 14.3% of surface water samples (Samples No. 8S) lie in the fields C4-S2, which reflect unacceptable for irrigation, except for very salt-tolerant plants, excellent drainage, frequent leaching, and intensive management and problems on fine texture soils and sodium-sensitive plants, especially under low-leaching conditions, but could be used on sandy soils with good permeability.

Sodium percentage (Na %)

The groundwater samples are suitable for irrigation in 33.3% of samples and 42.8% of surface water samples according to Na% values. 50% and 28.6% from groundwater samples and surface water samples, respectively, were Permissible. 8.3% and 28.6% from groundwater samples and surface water samples, respectively, were doubtful. 8.3% and zero% from groundwater samples and surface water samples, respectively, were unsuitable (table 11).

Table 10: Classification of irrigation water based on SAR values [42]

Level	SAR	Quality	Hazard
S1	<10	Low sodium	No harmful effects from sodium.
S2	10-18	Medium sodium	Problems on fine texture soils and sodium-sensitive plants, especially under low-leaching conditions, but could be used on sandy soils with good permeability.
S3	18-26	High sodium	Harmful effects could be anticipated in most soils and amendments such as gypsum would be necessary to exchange sodium ions.
S4	>26	Very high sodium	Generally unsatisfactory for irrigation.

Table 11: Suitability for irrigation based on sodium percent

Na%	Suitability for irrigation	No. of samples	Percentage (%)
<20	Excellent	-	-
20-40	Good	8 G-3S	33.33-42.8
40-60	Permissible	12G-2S	50-28.6
60-80	Doubtful	2G-2S	8.33-28.6
>80	Unsuitable	2G	8.33

Magnesium ratio (MR)

Calcium and magnesium maintain equilibrium in most waters, in equilibrium. Mg^{2+} in the waters will adversely affect crop yield; magnesium impact on irrigated water is expressed as magnesium ratio (MR) ($\text{MR}>50\%$ is suitable for irrigation and $\text{MR}<50\%$ is unsuitable).

M. R values reflected that 45.8% of investigating groundwater samples are unsuitable for irrigation table 6.

Residual sodium carbonate (RSC)

An excess of sodium bicarbonate and carbonate is considered to be detrimental to the physical properties of soils as it causes dissolution of organic matter in the soil, which in turn leaves a black stain on the soil surface on drying; this excess amount is denoted by Residual Sodium Carbonate (RSC).

All samples (ground and surface) is good for irrigation table 6. (R. S. C>2.5 meq/l is unsuitable for irrigation, RSC values from 1.25 to 2.5

meq/l are doubtful and R. S. C<1.25 meq/l are good for irrigation) [43].

CONCLUSION

The water resources in the study area are generally alkaline in nature. 83.3% of groundwater samples are fresh water and 16.7% are brackish water. 85.7% of surface water samples are fresh and 14.3% is saline. The groundwater distribution of sulfate indicates the presence of local zones of high concentrations at Abu Zaabal, reflecting the effect of the saline pond from the west and the influence of the sulfate fertilizers in the new reclaimed land in the east. The NPI values presented that the majority of the study area located under light polluted zone due to the influence of agricultural activities (nitrification of synthetic fertilizers and soil organic nitrogen) and moderate pollution zone is located closed to Bilbeis drain reflected that the influence of groundwater recharge from the drain According to WQI values, the distribution of drinking water quality index values for the groundwater samples in Abu Zaabal area reflected the effect of the Belbis drain on the groundwater quality. 33.3% of groundwater and 42.8% of surface water samples are suitable for irrigation according to Na% values. 45.8% of investigated groundwater samples are unsuitable for irrigation according to M. R values.

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AUTHORS CONTRIBUTIONS

Prof. Dr. Ragaa El-Sheikh; Prof. Dr. Ayman A. Gouda and Dr. Ehab Zaghlool have been generated the research idea and interpreted the data and helped to draft the manuscript. Dr. Mohamed E. A. Ali has suggested the research idea and participated in the design of the study. Mr. Ibrahim Hegazy was prepared the solutions, carried out the experiments, interpreted the data and helped to draft the manuscript.

CONFLICTS OF INTERESTS

The authors confirm that this article's content has no conflict of interest.

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