



Integrated stratigraphical, hydrogeochemical, and geophysical studies to characterize the shallow groundwater aquifers in Wadi Araba area, western Gulf of Suez, Egypt

Zenhom E. Salem¹ · Maha Abdelazeem² · Mohamed Mostafa Khalifa² · Mohamed Sobhy Fathy¹

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Abstract

The construction of the New Galala City besides great mining (e.g., konoozmisr) and Seawater Reverse Osmosis Desalination Plant projects in the semiarid Gulf of Suez region of Egypt increases the demand for groundwater resources. So, the current research provides integrated stratigraphical, hydrogeochemical, and geophysical studies to investigate the shallow groundwater aquifers in Wadi Araba area on the western side of the Gulf of Suez. The stratigraphic column starts with pre-Cambrian basement rocks which are non-conformably overlain by sedimentary succession of Phanerozoic Erathem. A detailed land magnetic profile has been conducted and processed to estimate the aquifer's geometrical parameters. This includes the subsurface spatial extension and thickness to delineate the prevailed subsurface structures in the study area. Lithostratigraphically, these aquifers are represented by the Paleozoic- lower Cretaceous siliciclastic-dominated sediments of the Nubia Group and fractured carbonate-dominated facies of the upper Cretaceous Galala Formation. Lithological characters, sedimentary facies (fluvial to shallow and open marine), microfacies types, and diagenesis processes (e.g., compaction, cementation, dissolution, and fracturing) are the main factors that affect the quality of the shallow aquifers in the study area. The main microfacies of the studied aquifers are quartz arenite, subarkose, sublitharenite, mudstone/wackestone, and packstone/grainstone. These facies are characterized by different primary and secondary porosities. According to the water sampling standard methods, nineteen groundwater samples were collected from the available hand-dug wells and springs from the Nubian sandstone aquifer and the Upper Cretaceous aquifer, respectively. Inductively coupled plasma-mass spectrometry (ICP-MS), titration, spectrophotometer and a flame photometer were used to obtain a complete chemical analysis. The results of hydrochemical analyses indicated that total dissolved solids (TDS) and concentrations of major ions as well in the lower Cretaceous aquifer increase toward the Gulf of Suez and in general higher than that of springs of the Upper Cretaceous Aquifer. The predominant factors influencing groundwater chemistry include seawater intrusion, evaporation, mineral alteration, dissolution of fossils, and ion exchange. Groundwater in the study area is unsuitable for drinking except in one sample and moderately suitable for irrigation with caution for higher salinity in samples located near the Gulf of Suez.

Keywords Wadi Araba · Nubian sandstone aquifer · Carbonate sediments · Aquifer quality · Magnetic tomography · Gulf of Suez

✉ Maha Abdelazeem
maazeem03@nriag.sci.eg

Zenhom E. Salem
zenhomsalem@yahoo.com

Mohamed Mostafa Khalifa
moh.moustafa92@gmail.com

Mohamed Sobhy Fathy
msfathy@science.tanta.edu.eg

¹ Geology Department, Faculty of Science, Tanta University, Tanta, Egypt

² Geomagnetism and Geoelectricity Department, National Research Institute of Astronomy and Geophysics, Helwan, Cairo, Egypt

Introduction

Egypt’s current principal challenge is the dire need for better development and management of the country’s finite water, land, and energy resources to meet the country’s population growth needs (Ashour et al. 2009). Groundwater represents the main source of freshwater supply for the local inhabitants. As tourism activities, quarrying, and the population of the area increase, spatial attention should be focused on the future development of the water supplies of Wadi Araba, west of the Gulf. In this respect, ambitious plans for groundwater exploration and evaluation are required (Aggour 1990). Nowadays, the demand for groundwater increased in the studied area because of the construction of the New Galal City close to Wadi Araba.

The area of study is located on the western side of the Gulf of Suez. It is bounded by latitudes 28° 30` and 29° 30` N and longitudes 31° 45` and 32° 45` E (Fig. 1) and

extends in an E–W direction between Northern Galala and Southern Galala plateaus with a total length and area of about 40 km and 1100 Km², respectively. The geological map of the area under study is shown (Fig. 2), which is modified after Conoco (1987). Since Schweinfurth (1885) recorded the Paleozoic fossils in Wadi Araba, the area attracted many geologists to investigate the stratigraphical and structural settings of the area. Most of the rock units of the Paleozoic–Mesozoic sequences of the studied area (e.g., Rod El Hamal, Qiseib, and Malha formations) were introduced by Abdallah and El Adendani (1963) and Abdallah et al. (1965). A revision of the lithostratigraphic classification of the Permo-Carboniferous outcrops of the Gulf of Suez region was carried out by Kora and Mansour (1992). Bandel and Kuss (1987) delineated the depositional environment of Palaeozoic–Eocene sediments at the Northern and Southern Galala plateau, which ranged from fluvial to open marine environments. Petrophysically, Nabawy et al. (2020) noted that the porosity and

Fig. 1 Location map of the study area, west of the Gulf of Suez, Egypt with position of the studied boreholes, collected rock samples and land magnetic measurements on the Landsat image of Wadi Araba, west Gulf of Suez, Egypt

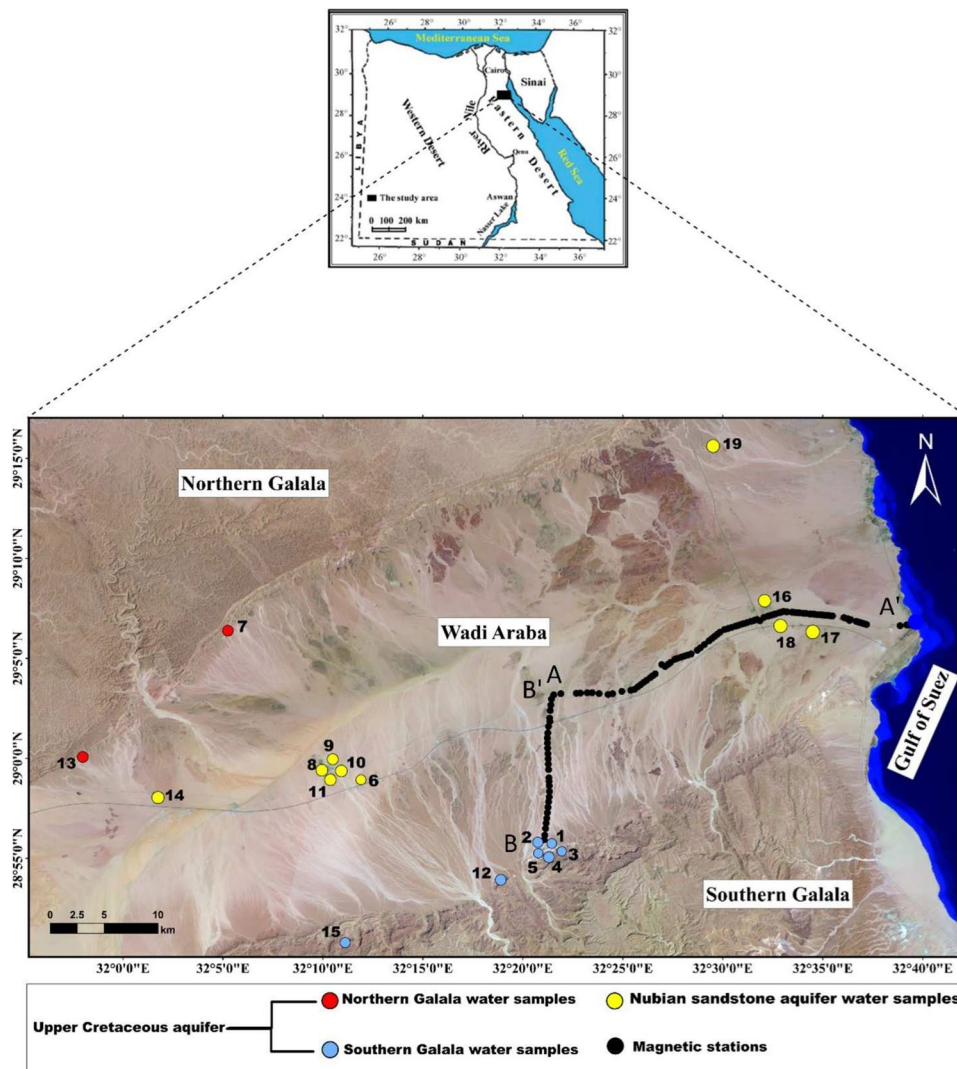
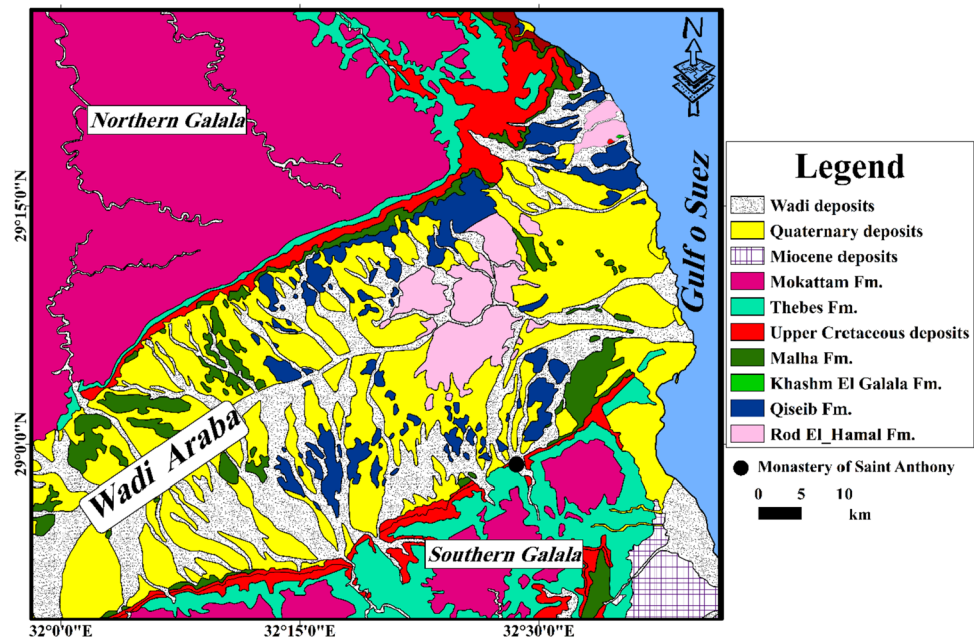


Fig. 2 Geological map of the studied Wadi Araba area (modified after Conoco 1987)



permeability of the Nubian microfacies of quartz arenite, sublitharenite and arkose are good to excellent, whereas the ferruginous subarkose to greywacke is characterized by relatively less petrophysical potential than the others.

Two structural phases were detected by Moustafa and Khalil (1994) in the Gulf of Suez region. In the Late Cretaceous phase of deformation, folds with NE–SW oriented axes were formed in northern Egypt as a result of convergence between Africa and Eurasia and the closure of the Neotethys. In the second early Miocene phase, deformation NW-oriented normal faults were formed as a result of the opening of the Suez Rift. The tertiary carbonate platform in the studied area formed as a result of the initial topography that was controlled by the tectonic uplift of the Northern Galala/Wadi Araba Syrian Arc structure (Scheibner et al. 2001).

Aggour (1990) elucidated the hydrogeologic framework of Wadi Araba. He classified Wadi Araba as a high Wadi according to its relative groundwater possibility. Rocks belonging to Carboniferous and Lower Cretaceous sandstone (Nubian aquifer) represent the main source of water in the Wadi Araba depression. The water is tapped from springs, shallow wells, and occasionally deep wells. The quaternary alluvium deposits constitute an insignificant water-bearing horizon. Gomaa and Aggour (1999) studied the hydrogeological and hydrogeochemical conditions of carbonate aquifers in the Gulf of Suez region. They considered that the carbonate rocks represent productive aquifers on both sides of the Gulf of Suez. These rocks constitute extensive watershed areas, e.g., Northern Galala, Southern Galala, and Abu Sha'r plateau on the western side. Some recommendations were proposed in this study for the development of groundwater

resources. Hand-dug wells are suggested along the scraps of the plateaus at the favorable sites and more attention should be focused on the utilization of the flowing water such as the construction of surface reservoirs to control the discharge and use. Younis et al. (2019) used VES/TEM techniques at the North Galala plateau to assess the present situation of groundwater in the Quaternary and Pliocene aquifers. They found groundwater in the shallow aquifer might be of good quality than the water in the deep aquifer. They also found the most appropriate areas for burrowing new water wells.

On the other hand, the magnetic method is a geophysical technique that proved its efficiency when applied to hydrogeological investigations. Of all geophysical techniques, the magnetic method is the most versatile since it may be used for both shallow and deep research (Dobrin and Savit 1988). Additionally, it is affordable and includes both local and regional investigations. The magnetic method has primarily been used for regional surveys in hydrocarbon exploration (Abdelazeem et al. 2020, 2021; De Smet et al. 2020; Abdelrahman et al. 2019; Gobashy and Al-Garni 2008; Al Garni et al. 2006); engineering geophysics (Araffa et al. 2021; Azeem et al. 2014); groundwater exploration (Gobashy et al. 2021b); mineral exploration (Gobashy et al. 2022, 2021a; El-Sawy et al. 2018; Abdelrahman et al. 2019); and mapping geological structures and igneous intrusions (Al Garni et al. 2012; Wang et al. 2017); in addition, estimating sedimentary basin thickness and identifying groundwater contamination (Nazih et al. 2022; Rehman et al. 2019) and geologic hazard and risk assessments (Fergany et al. 2014; Al Garni and Gobashy 2010) are examples of a such wide range of applications.

Based on the magnetic method, Abdelazeem et al. (2019) have delineated the main structures in this area as NW–SE, and E–W trends, with minor trends as the N–S, NE, ENE, WNW, NNW, NNE, and ENE, and noted that the carbonate rocks suffered from various diagenetic processes for a long time and have different caves, joints, sinkholes, and karst surfaces.

With the remarkable variations in the Wadi Araba vertical and lateral facies, the goal of this study is to explore and identify qualitatively the shallow groundwater aquifer from the detailed integrated petrographic, stratigraphic, geochemical, and geophysical point of view. The study aims to support groundwater sustainability and conservation. To identify the subsurface structure geometries, magnetic tomography is essentially used. The findings should help control the area's subterranean water resources to prevent the region's groundwater levels and quality from declining and facilitate upcoming upgrades within the region. The workflow presented helps globally to preserve and develop groundwater resources.

Methodology

Data collection

In the study area, 115 rock samples (WA1 to WA115) have been collected and described from the exposed Carboniferous–Cretaceous successions, and 47 blue-dyed thin sections are investigated and photographed under the polarized microscope for microfacies analysis, diagenesis, and for rock porosity estimation. The classification of Pettijohn et al. (1987) was used to classify the sandstones of the Nubia Group, whereas the classification of Dunham (1962) and Folk (1962) for carbonate rocks is adopted in the present work. Moreover, a total of 112 magnetic stations were measured along geologically selected localities. The measurements were carried out by using an Over-Houser magnetometer (GSM-19 v7.0) with a sensitivity of 0.022 nT@1 Hz. El Misallat station in Fayoum is considered as the base station (used for diurnal correction). Land magnetic data were conducted using GSM-19 v7.0 with the same sensitivity along a selected profile running from west to east as shown in Fig. 1. The total number of stations is 112 measured according to the accessibility of the area. The measurements are carried out on stations marked every 200–500 m, which is suitable to the objective of the study, to explore the aquifer and the effect of the structure on it.

There are no distributed drilled water wells (except one well) in the study area because there are no agriculture, urban or industrial activities. Therefore, the hydrogeochemical study was undertaken by collecting 16 groundwater samples during April from the available dug wells, one borehole,

and springs. The dug wells and the drilled well are tapped from the Nubian sandstone aquifer (from the mainstream of Wadi Araba) and the springs are tapped from the Upper Cretaceous limestones (Southern escarp of the Northern Galala and Northern escarp of the Southern Galala plateaus). Samples 17, 18, and 19 chemical analyses were obtained from Ezzeldin (2010) because it was not possible to sample them as the weathering products covered the spring itself. Water was sampled in new—additionally, rinsed—plastic bottles, and chemical analysis were done a few days after sampling.

Hydrogeochemical analyses

Parameters such as pH (Portable Consort pH Meter, Model P 314), electrical conductivity (EC), and TDS (Hach's Portable Conductivity/Total dissolved solids TDS Meter) were measured for the collected samples in the field (Hach 1990). Major and trace ions were analyzed in the central laboratory of the Desert Research Centre (DRC). ICP–MS (inductively coupled plasma–mass spectrometry) which applies to the determination of sub- $\mu\text{g/L}$ concentration of a large number of elements in water samples (Hoaghia et al. 2021) was used for trace elements analyses. Digital Titration method was applied for determining the concentration of CO_3^{2-} and HCO_3^- against a standard sulfuric acid solution, Ca^{++} and Mg^{++} using a total hardness test and chloride using a standard solution of mercuric nitrate and diphenylcarbazone reagent powder. SO_4^{2-} concentration was measured by spectrophotometer (Hach's Direct Reading (DR)/2000) precipitation with barium chloride. A flame photometer was used for Na^+ and K^+ determination (Banerjee and Prasad 2020). The obtained hydrochemical analysis has a charge balance error of less than 5%. The sample's locations map is shown in Fig. 1, and the statistical summary of concentration of the measured hydrochemical parameters in the two aquifers are listed in Tables 1 and 2.

Result and discussions

Magnetic data

Land magnetic profile

Regular data reduction is considered including diurnal reduction (subtracting base station readings from the measured field) and denoising filtering (omitting the odd readings). The reduced-to-pole filter is then applied (using ZondGM2D) to overcome the geomagnetic field polarity shift. The resulting RTP data are then interpreted. Euler deconvolution has also been applied (using ZondGM2D) to ensure the expected faults/contacts.

Table 1 Statistical summary of concentration of the measured major ions

| | Aquifers | pH | EC | TDS | TH | CATIONS | | | | ANIONS | | | |
|---------|-------------|----|--------|--------|------|---------|----------------|-----------------|------------------|------------------|-----------------|-------------------------------|-------------------------------|
| | | | | | | mg/l | | | | | | | |
| | | | | | | μS/cm | K ⁺ | Na ⁺ | Mg ⁺⁺ | Ca ⁺⁺ | Cl ⁻ | SO ₄ ⁻⁻ | HCO ₃ ⁻ |
| MAX | Galala Fm | 9 | 13,646 | 8734 | 886 | 8 | 2900 | 144 | 189 | 2055 | 2650 | 1327 | 292 |
| MIN | | 7 | 1987 | 951 | 508 | 3 | 180 | 41 | 21 | 397 | 166 | 159 | 0 |
| STDEV | | 1 | 3721 | 2442 | 133 | 2 | 871 | 34 | 45 | 521 | 788 | 380 | 96 |
| Average | | 8 | 4021 | 2336 | 629 | 5 | 609 | 77 | 125 | 755 | 590 | 316 | 36 |
| MAX | Nubia Group | 8 | 38,520 | 24,770 | 7064 | 264 | 7200 | 751 | 2010 | 13,947 | 3001 | 325 | 26 |
| MIN | | 5 | 3266 | 1745 | 525 | 3 | 420 | 41 | 142 | 590 | 277 | 31 | 0 |
| STDEV | | 1 | 13,998 | 8625 | 2337 | 80 | 2523 | 249 | 589 | 5259 | 962 | 87 | 12 |
| Average | | 8 | 14,441 | 8995 | 2612 | 52 | 2354 | 274 | 594 | 4627 | 1197 | 127 | 11 |

Table 2 Statistical summary of concentration of the measured trace elements

| | Aquifers | mg/l | | | | | | | | | | | | |
|---------|-------------|-------|-------|-------|-------|-------|--------|--------|-------|-------|-------|-------|-------|--------|
| | | Al | Cd | Co | Cr | Cu | Fe | Mn | Mo | Ni | Pb | V | Zn | Si |
| Max | Galala Fm | 0.779 | 0.001 | 0.001 | 0.000 | 0.032 | 0.245 | 0.084 | 0.013 | 0.003 | 0.215 | 0.057 | 0.088 | 32.646 |
| Min | | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 4.175 |
| STDEV | | 0.248 | 0.000 | 0.000 | 0.000 | 0.011 | 0.085 | 0.027 | 0.004 | 0.001 | 0.083 | 0.020 | 0.031 | 9.229 |
| Average | | 0.155 | 0.000 | 0.000 | 0.000 | 0.004 | 0.039 | 0.013 | 0.004 | 0.000 | 0.041 | 0.011 | 0.022 | 8.434 |
| Max | Nubia Group | 0.456 | 0.001 | 0.003 | 0.047 | 0.029 | 60.350 | 10.890 | 0.006 | 0.013 | 0.235 | 0.054 | 1.337 | 29.548 |
| Min | | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.000 | 0.271 |
| STDEV | | 0.180 | 0.000 | 0.001 | 0.015 | 0.011 | 18.904 | 3.425 | 0.002 | 0.004 | 0.093 | 0.022 | 0.426 | 10.572 |
| Average | | 0.192 | 0.000 | 0.001 | 0.005 | 0.005 | 7.108 | 1.337 | 0.003 | 0.002 | 0.062 | 0.019 | 0.187 | 10.271 |

Quantitative interpretation

Quantitative magnetic interpretation is an essential step for the delineation of subsurface structures and the distribution of rock units. The surface magnetic anomalies have resulted from subsurface variations (either lateral or vertical) in the magnetic susceptibilities of rock units. Modern methodologies and efficient rapid algorithms are used for inverting surface magnetic anomalies to geologically plausible models. In this work, along measured two profiles, AA” and BB’, the subsurface is assumed to be subdivided into a large number of cells or blocks, each with unknown magnetic susceptibility (*k*). This model is subjected to a tomographic inversion process using the Occam inversion procedure (Constable, 1987) to retrieve the possible distribution of the magnetic susceptibilities along the 2D profile as shown in Fig. 3a, b. The equation used for the inversion (uses the depth weighting to overcome erroneous shallow depth bodies, also covariance matrix for smoother model) is given as:

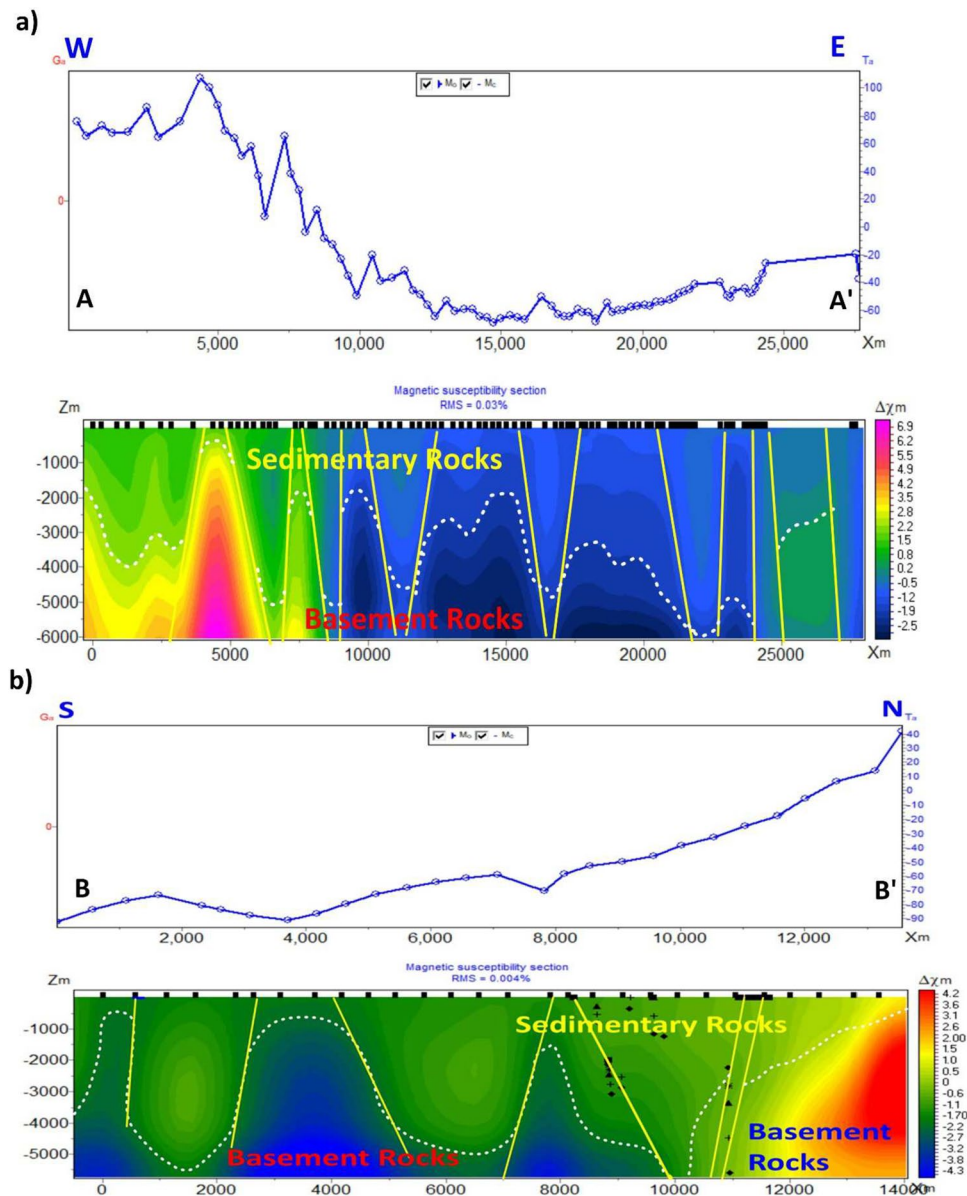
$$(A^T W^T W A + \mu C^T C) \Delta m = A^T W^T \Delta f - \mu C^T C m \tag{1}$$

where *A* is the kernel matrix that is related directly to the mesh design or the constructed subsurface model, *W* is the weighting matrix that minimizes the distortion of *A* with increasing depth, *C* smoothing operator, *m* the model parameter vector, μ —regularizing parameter, and Δf is the vector of residuals between the observed and calculated values. The Occam inversion algorithm is used to model the subsurface and minimize the difference between the observed and the modeled field based on the formula in Eq. (1). Moreover, the Euler deconvolution (Thompson 1982) is applied along the profiles to estimate possible locations and depths to magnetized targets, which are posted on the tomographic model as black dots, indicating the different depths of sources. The Euler’s homogeneity equation in two dimensions (2D) (Reid et al. 1990) is expressed as:

$$(x - x_o) \frac{\partial T}{\partial x} + (z - z_o) \frac{\partial T}{\partial z} = N(B - T) \tag{2}$$

where *T* (*x*, *z*) is the total magnetic field in Cartesian coordinates, (*x*_o, *z*_o) the coordinates of the source, *N* is the structural index, and *B* is the regional field (Thompson 1982).

Fig. 3 Tomographic inversion results for the land magnetic profiles. **a** Profile A-A', and **b** Profile B-B'. Black dots indicate calculated Euler deconvolution depths, yellow lines are the expected faults, and white dotted line indicates the possible basement surface



The inverted models of the profiles are shown in Fig. 3. The total length of the profile AA' reaches 28 km, while that of BB' is 14 km cutting the different structural units. The Zondgm2d software is used for the inversion process. The resulting tomographic sections represent the possible distribution of the subsurface magnetic susceptibilities along the two profiles. Each profile is projected to a central straight line that best fits the original one. This process is necessary for preparing the data for inversion along a regular array of data. Blue colors are used to indicate low susceptibility contrast ratings, whereas red and pink colors indicate high values. Using a K-20 magnetic susceptibility meter, surface magnetic susceptibilities are also measured along the profile. The inverted profile is constrained by the observed magnetic susceptibilities, which are employed as *a priori* information.

Profile AA' (Fig. 3a) is running parallel to the Wadi axis. Two types of basement rocks with different magnetic susceptibilities can be recognized. The depth to the basement varies from 2 km at the west to 5 km close to the Gulf of Suez. Tomographic inversion shows an increase in effective magnetic susceptibilities in the western direction in the first 7 km from the measured profile than the eastern direction. The increase in the magnetic anomalies and hence the subsurface magnetic susceptibilities may be related to the ferruginous characteristics of the Nubian Sandstone aquifer as the stratigraphical and microfacies analyses indicate. Possible contact is indicated as a dotted line separating the sedimentary rocks from the basement rocks. This is determined according to the large difference in the susceptibility contrast observed at this level. Consequently, the expected aquifer

is geometrically controlled by the basement topography as indicated by the solid lines representing possible faults along the profile. The thickness of the aquifer increases toward the east and decreases toward the west. Magnetic contacts are observed with the sedimentary succession. Fractures may descend the basement and/or the sedimentary cover.

Profile BB' (Fig. 3b) is similar to AA'. It runs from south to north crossing the Wadi. It reflects the same features as profile AA'. The sedimentary rocks to the north are characterized by relatively high magnetic susceptibility contrast in comparison with the low susceptible basement which may be composed of granitic rocks. The depth to the basement is shallow toward the south and increases to the north. This is indicated by Euler solution depths that is represented as black dots. The section is controlled by many faults that are nearly striking north–south.

In summary, the magnetic inversion shows that the aquifer is highly structurally controlled. The thickness of the aquifer increases toward the east and decreases toward the west. Magnetic contacts are observed within the sedimentary succession.

Stratigraphical and microfacies analyses

Stratigraphic setting

The lithostratigraphy of the studied Phanerozoic Erathem is established based on field investigations, rock description, sedimentary structures, stratigraphic contacts, and fossil contents. The oldest exposed rocks in the Wadi Araba are represented by Rod El Hamal Formation of Carboniferous age, which is unconformably overlain by the Qiseib Formation (Permo-Triassic), Khashm El Galala Formation (Jurassic) and Malha Formation (Lower Cretaceous). These rock units are equivalent to the subsurface Nubia Group in the Gulf of Suez province. The upper Cretaceous Galala Formation is exposed along the two borders of the Wadi Araba (Fig. 2).

The Carboniferous Rod El Hamal Formation (Carboniferous age) forms the main bedrock of the eastern part of Wadi Araba. It consists of five succeeded blocks of about 250 m (Fig. 4). The lower 70m is composed of sandstone, yellow, moderately hard cross-bedded, and pebbly, with the thin lamina of shale and siltstone intercalations (Fig. 5A), followed by fossiliferous mixed carbonate-siliciclastic sediments of about 40 m. Pebbly, cross-bedded sandstones with mudstone and limestone intercalation, is the main constituent of the upper part of the formation (Fig. 5B). Many fossils of Carboniferous age are encountered in the field and under a microscope with the sediments of this formation, e.g., corals (*Amygdalophylloides ivanovi* and *Bothrophyllum pseudoconicum*), brachiopods, echinoderms, and foraminifers (Fig. 6G–H). This formation is coeval to Um Bogma

Formation in Sinai, the Gilf Formation at Aswan, the sub-surface Desouqy Formation of Egypt, the M'rar Formation in Libya and the Berwath Formation of Saudia Arabia (Kora 1998).

The Permo-Triassic red beds of the Qiseib Formation unconformably overlain the Rod El Hamal Formation. This formation is widely distributed in the Wadi (Fig. 2), which is represented by about 100m, from reddish yellow, cross-bedded pebbly sandstones with varicolored mudstone intercalations (Figs. 4 and 5C). Yellow orange dolostone thin beds with badly preserved fossils are noted in the middle part of the formation in some places. The Qiseib Formation is widely distributed in the Easter Desert and Sinai of Egypt, the Amanus Sand of Syria and the Unayzah Formation of Saudi Arabia (Kora 1998).

The Jurassic Khashm El Galala Formation was exposed due to faulting as a small block (20m) at the eastern entrance of the Wadi. It consists mainly of marine sediments greenish shale and marl with limestone intercalations (Fig. 4).

The best exposures of the lower Cretaceous Malha Formation appear at the base of the borders of Wadi Araba (Fig. 2). It consists of about 140m, from non-marine friable, cross-bedded, coarse-to-fine-grained sandstones and kaolinitic mudstone interbeds with occasional marine layers at the upper part (Figs. 4 and 5D). It is widely distributed in Sinai, Eastern and Western Deserts of Egypt and equal to Abu Ballas Formation in Aswan.

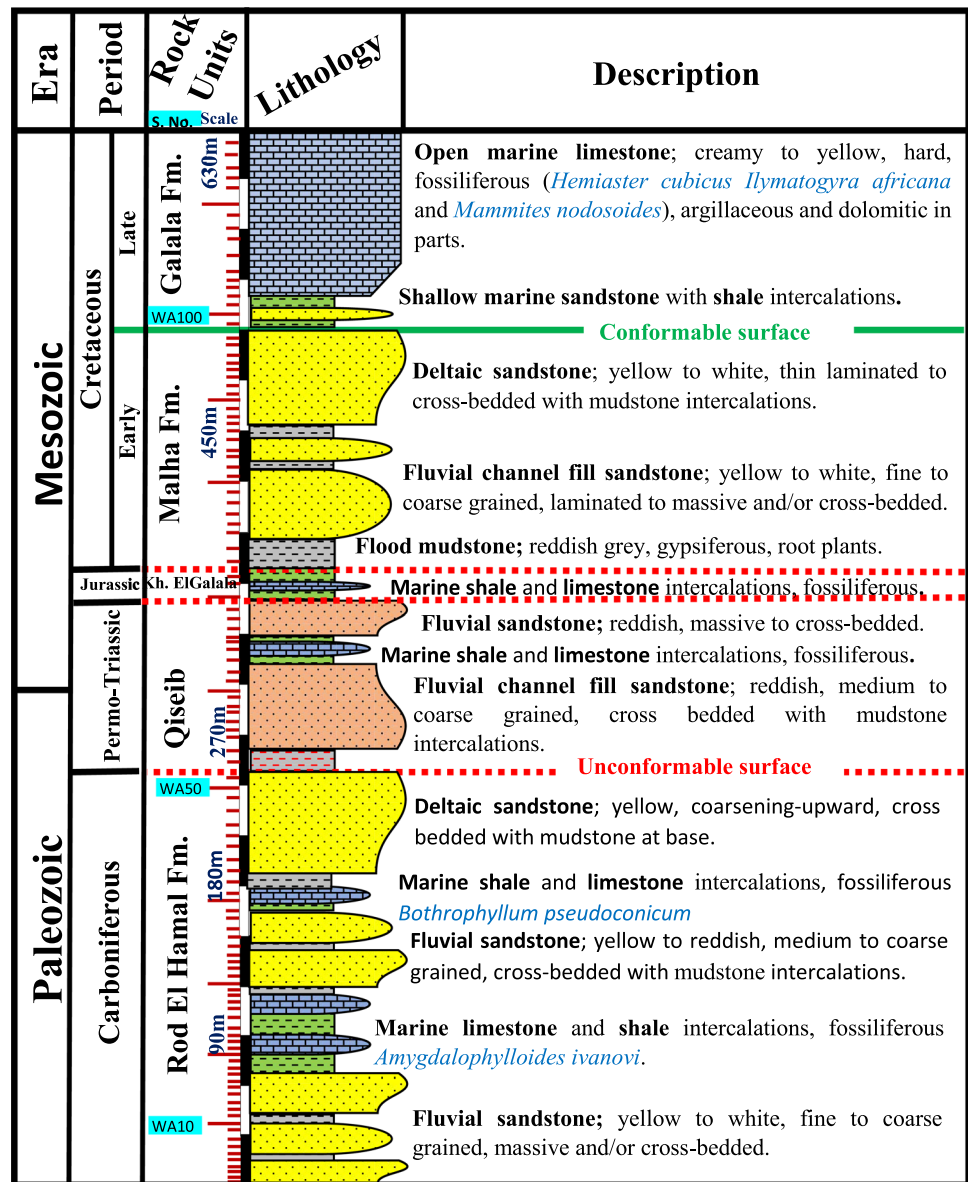
The upper Cretaceous Galala Formation represents the carbonate-dominated facies (120m) in the studied area (Fig. 4). Its basal part is characterized by yellowish-green clastic sediments of sandstone, siltstone, and shales. Most succession is composed mainly of fossiliferous, thin-bedded to massive, dolomitic limestones and marls (Fig. 5F–H). These sediments encountered many shallow marine faunas of upper Cretaceous age from echinoderms (*Hemias-ter cubicus*), pelecypods (*Ilymatogyra Africana*), cephalopods (*Mammites nodosoides*), corals, and foraminifers (Fig. 7A–F). Galala Formation was recorded in different localities in the Eastern Desert and is equal to the Raha Formation in Sinai of Egypt (Kora et al. 2001). Unconformably, it succeeded by the Eocene carbonate-dominated rocks of Thebes and Mokattam formations.

A thin layer of the quaternary deposits covers most parts of the Wadi Araba (Fig. 2). The grain size of these siliciclastic sediments varies from gravel-, sand-, silt- and mud-sized. It is represented by Wadi deposits, Aeolian dunes and sheets.

Microfacies analysis

In addition to stratigraphic features and sedimentary structures, petrographic examination and microfacies analysis are carried out to delineate the characteristics and quality of the shallow aquifers in the Wadi Araba area. The encountered

Fig. 4 Composite lithostratigraphic chart of the exposed Carboniferous–Cretaceous systems with prevailed sedimentary facies and depositional environments in the Wadi Araba area, western Gulf of Suez, Egypt



siliciclastic and carbonate microfacies and their diagenetic processes are briefly described and discussed below:

Quartz arenite The sandstones of the quartz arenite microfacies are characterized by white to pale yellow color, friable to soft, and massive to planar cross-bedding. Microscopically, it is composed mainly of quartz grains representing 96% of the rock materials. The quartz grains are medium to coarse-sized, rounded to subrounded, and moderately to well-sorted and corroded and cracked in parts (Fig. 6A–C). Most of the quartz grains are monocrystalline, but there is polycrystalline undergoing undulate extinction. About 4% of the rock constituents consist of iron oxides, glauconite, chert, stable heavy minerals, zircon, and tourmaline. As shown in Fig. 5C silica overgrowth, calcareous and fer-

ruginous coatings are the main cement materials. The blue dye in the thin sections represents the intergranular porosity which reached about 18–20% of the rock volume of the shallow aquifers.

Subarkose Petrographically, subarkose microfacies are made up of quartz grains which form about 85% of the rock materials and more than 10% of different types of feldspars and other rock fragments (Fig. 6D, E). Other grains are also recorded as iron oxides, opaque minerals, glauconite, mica chlorite, and zircon. The subangular and subrounded quartz grains are mono- to polycrystalline, cracked, and moderately sorted. Most alkali feldspar, plagioclase, biotite, chlorite, and glauconite minerals are subjected to alterations and partial dissolution (Fig. 6E) causing secondary porosity

Fig. 5 Field photographs of the Carboniferous–Cretaceous systems in the Wadi Araba area, west Gulf of Suez, Egypt; **A, B**—shallow marine and deltaic sandstones of the Carboniferous Rod El Hamal Fm., yellow triangular refer to fining- and coarsening-upward, respectively, blue arrows refer to cross-bedding, red arrows refer to NE fracture system, hammer for scale; **C**—Fluvial (flood plain and channel fill) sandstones of the Permo-Triassic Qiseib Fm., red arrows refer to NW fracture system; **D**—Exposed coarsening-upward fluvial sandstones of lower Cretaceous Malha Fm. near RIGWA boreholes west Wadi Araba; **E**—RIGWA boreholes; **F**—Fractured open marine limestone of upper Cretaceous Galala Fm.; **G, H**—St. Antony and Enaba springs respectively within fractured limestones of Galala Fm



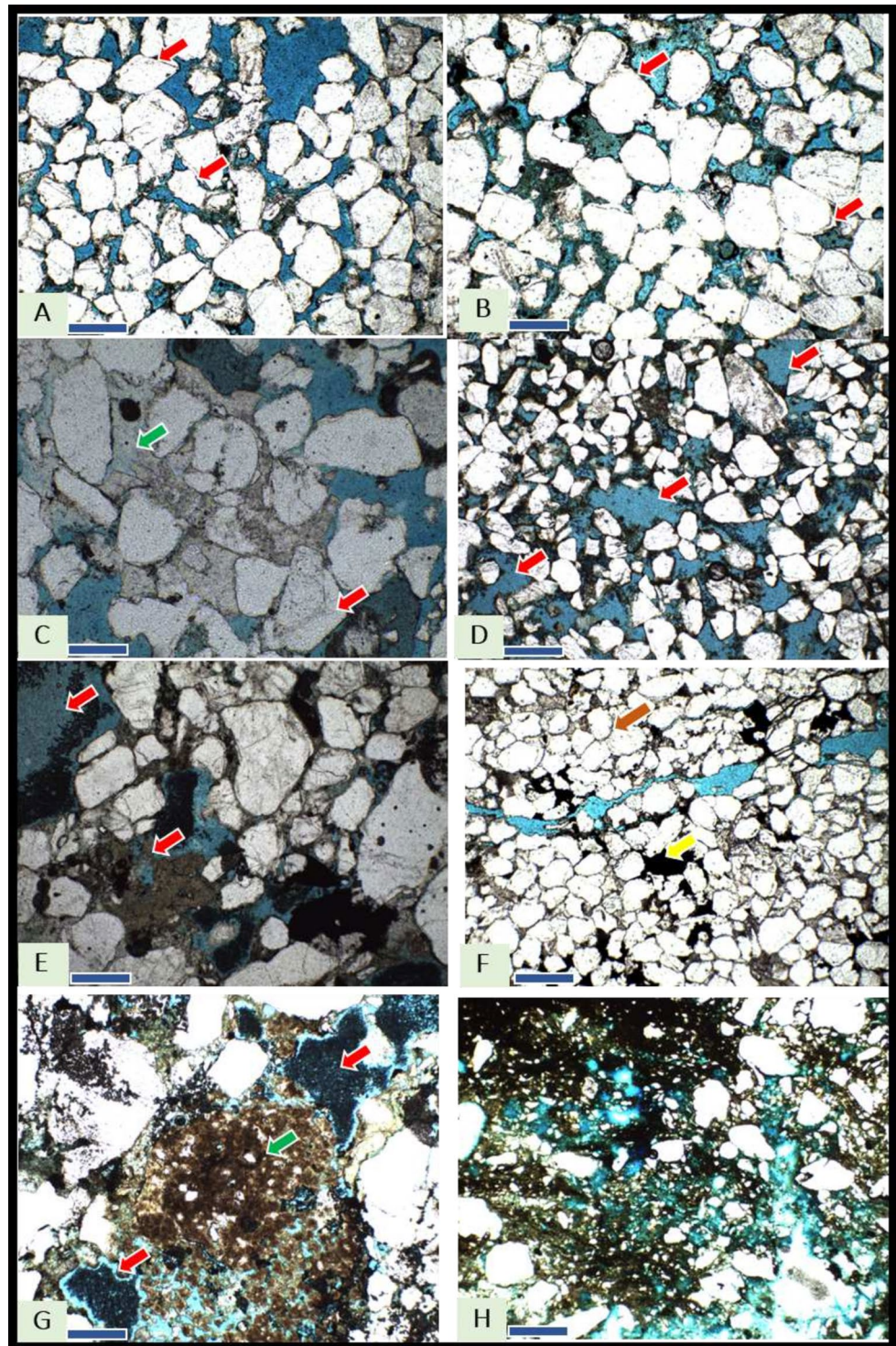
in Wadi Araba aquifers. Scattered altered biotite and hornblende flakes are also observed. All these rock materials are cemented by ferruginous and/or calcareous matrices.

Sublitharenite These microfacies are characterized by gray to brownish-yellow cross-bedded pebbly sandstone, which is recorded in all studied rock units of the shallow Nubian aquifers. It consists of quartz grains (75–83%) and rock fragments (about 20%) as shown in Fig. 6F. The mono- and polycrystalline quartz grains are poorly to moderately sorted and angular to subrounded in shape. The recorded rock fragments consist mainly of old sedimentary sediments, e.g., shales, limestones, and cherts. Some iron oxides, glauco-

nites, and heavy minerals are observed within the facies. The different types of rock materials are embedded in the fine matrix and ferruginous or calcareous cement. The porosity within such facies resulted from different diagenetic processes such as compaction, ferruginous cement, fracturing (Fig. 6F), dissolution, and dolomitization (Fig. 6E).

Greywackes The greywackes microfacies characterize the clastic sediments of the floodplain fluvial deposits of the Nubian sandstone aquifers. These immature, poorly sorted microfacies are composed mainly of pebbly to sand-sized fragments ranging from 70 to 75% of rock materials embedded in a fine-grained matrix (Fig. 6H). These

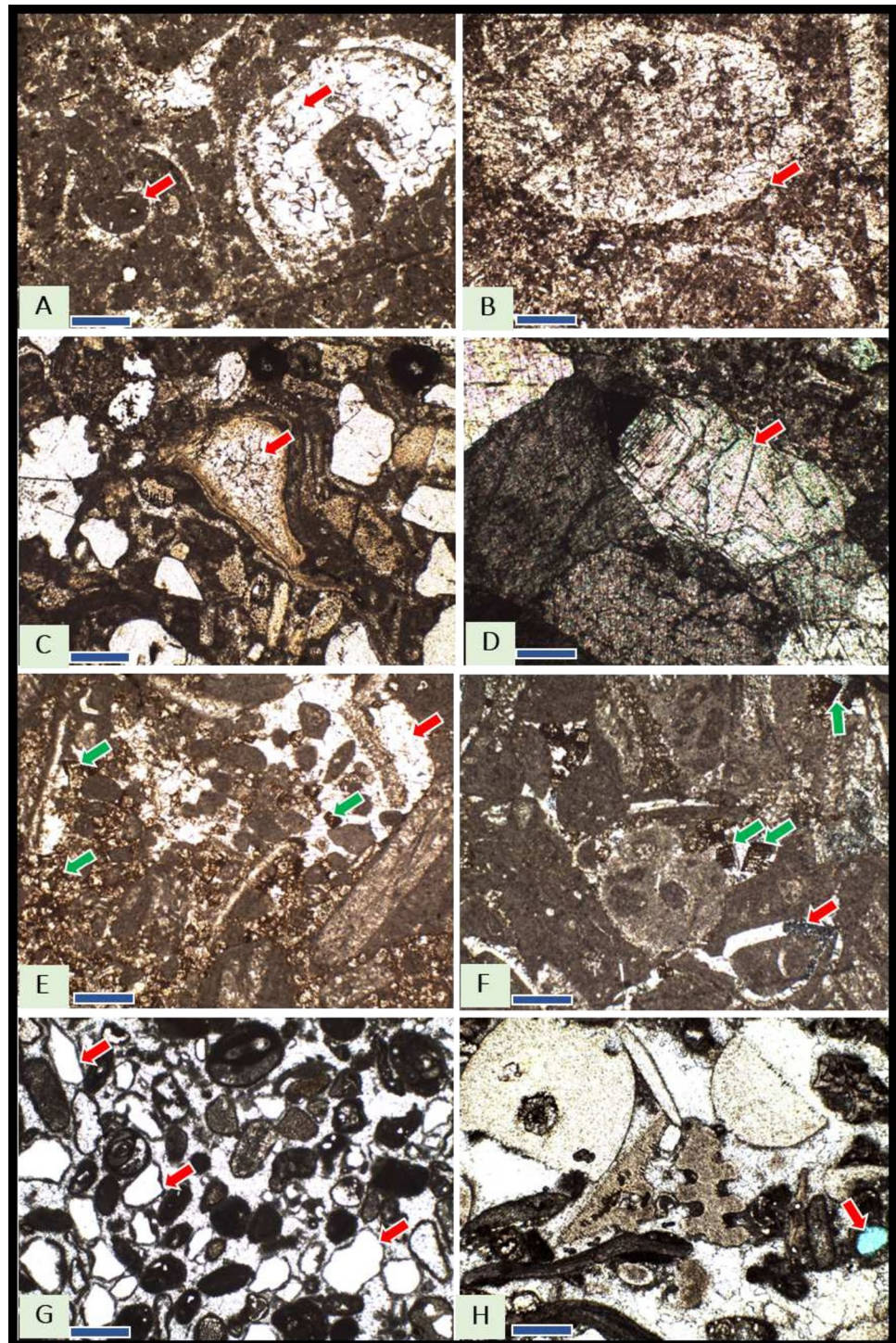
Fig. 6 Microscopic photographs of sandstone microfacies of the Nubia Group at Wadi Araba area; **A–C**—Quartz arenite of Rod El Hamal (WA7), Malha (WA77) and Qiseib (WA55) formations, respectively, red arrows refer to quartz overgrowth and green one refers to carbonate cement; **D, E**—Subarkose of Rod El Hamal Fm. (WA3, WA45), moderately sorted, red arrows refer to partial to complete dissolution of mafic minerals; **F, G**—Sublitharenite of Rod El Hamal Fm. (WA21, WA32), poor to moderately sorted, brown arrow refers to concavo-convex contacts between quartz grains, yellow to iron cement, red arrows to partial dissolution of rock fragments, and green to dolomitization. **H**—Greywacke microfacies of Qiseib Fm. (WA67), the blue color indicates porosity. All figs under PPL. Blue bar = 1mm



mixed facies consist of angular to subrounded polycrystalline quartz, altered feldspars, and cracked rock fragments. Besides the clayey matrix, the greywacke fragments are cemented by ferruginous and/or calcareous material. The blue color of the Fig. 6H refers to porosity within aquifer rocks.

Mudstone/Wackestone microfacies It is a common carbonate microfacies that constitutes about 90% of the shallow carbonate aquifers of the studied area. The limestones of these facies are white to yellow in color, thin-bedded, fossiliferous, dolomitic, and fractured. Microscopically, it is composed of several types of allochems (up to 40%)

Fig. 7 Microscopic photographs of limestone microfacies of the Galala Fm. (A to F) and Rod El Hamal Fm. (G and H) at Wadi Araba area; A, B—mudstone (biomicrite) red arrows refer to recrystallization of shell fragments (WA109). C, D—sandy wackestone (biomicrite), shell fragments filled with sparry calcite (red arrows) (WA103). E, F—Packstone (Dolomitic intra-biosparite) allochems imbedded in sparite (red arrows), which subjected to dolomitization in most parts (green arrows) (WA114). G—Grainstone (sandy pelbiosparite) (WA27) quartz grain coated by thin micritic film (red arrows). H—Grainstone (biosparite) (WA41) red arrow refers to partial dissolution of shell fragments (intragranular porosity). D and H under PLX, other under PPL. Blue bar = 1mm



from bioclasts (e.g., pelecypods, gastropods, corals), peloids, and aggregated grains (Fig. 7A–D). Some detrital silt to sand-sized quartz grains is observed within the facies. All these carbonate grains are embedded in a micritic and/or calcitic cement. However, the shell fragments of these microfacies were replaced and/or filled by calcite

minerals. Some calcite minerals recrystallized to form sparry calcite as shown in Fig. 7D

Packstone/Grainstone microfacies In the field, the carbonate sediments of these facies are characterized by massive to planar cross-bedded, dolomitic with fenestra and/or porous

fabrics. Petrographically, it consists mainly of well-sorted different allochems from bioclasts, ooids, peloids, intra-clasts, and aggregated grains with detrital quartz grains (Fig. 7E–H). The allochems represent about 90% of the rock constituent and are tightly packed and cemented by sparry calcite. The bioclasts are mainly represented by molluscs, bryozoans, foraminifera, algae, and echinoid plates. The main diagenetic processes recorded in the present facies are dolomitization, recrystallization (Fig. 7E, F), micritization (Fig. 7G), and dissolution (Fig. 7H) which formed a secondary type of porosity with the carbonate aquifers.

Diagenetic processes and rock porosity

As soon as the sediments of the studied area were deposited, they were subjected to different diagnostic physical, chemical, and biological processes. The most common diagenetic processes are compaction, cementation, recrystallization, dolomitization, dissolution, pressure solution, fracturing, and leaching out. Diagenetic processes played an important role in the aquifer porosity, and permeability as well as groundwater chemistry and quality.

Compaction, cementation, recrystallization, micritization, and pressure solution are the principal processes leading to porosity reduction in the shallow aquifers of the studied area. Concavo-convex contacts, cracking and corrosion of the quartz grains refer to compaction (Fig. 6F). According to Graton & Fraser, (1935), compaction can reduce the sandstone porosity to 26%. In most cases, calcareous (Fig. 6C), ferruginous (Fig. 6F), and micritic cement filled the pore spaces, vugs, and fractures. Moreover, silica overgrowth (Fig. 6A, B) and micrite and/or clay mineral coating (Fig. 7G) are recorded in many microfacies. However, all these processes lead to reducing rock porosity. The main source of silica and clay minerals resulted from the alteration and pressure solution of feldspar minerals (Bjorkum et al. 1993). Dolomitization is a process whereby limestone or its precursor sediment is completely or partly converted to dolomite by the replacement of the original CaCO_3 with magnesium carbonate, through the action of Mg-bearing water.

Several fractures and joint sets were observed in the field (Fig. 5B, C) and in microscopic scales (Fig. 6F) in the sediments which accompanied the tectonic events in the Wadi Araba area. However, these fractures and joints enhanced and increased the porosity and permeability of the studied aquifer forming potential pathways for groundwater movement. Dissolution of many unstable minerals within subarkose (Fig. 6D, E) and sublitharenite (Fig. 6G) of sandstone microfacies and fossil fragments of the carbonate microfacies (Fig. 7H) leads to increasing the inter- and intra-porosity of the studied shallow aquifers. According to Selly (1998), the dissolution process is a very important diagenetic factor for reservoir quality. In most cases, unsaturated groundwater

is considered the cause of dissolving the unstable minerals and shell fragments. Moreover, the dissolution of authigenic minerals or grains can enhance porosity as a secondary type (Ehrenberg 1990).

Hydrogeochemistry

Groundwater chemistry is greatly significant in determining the type of water, origin, quality, and suitability for different purposes. It is mainly based on the chemical analysis of the collected water samples from different aquifers and springs in the area (Fig. 1 and Tables 1 and 2). Groundwater at all sites is brackish to saline water according to the classification based on TDS (fresh: < 1000 mg/l, slightly saline: 1000–3000 mg/l, moderately saline: 3000–10,000 mg/l, highly saline: 10,000–35,000 mg/l) suggested by the USGS (2000). The integrated classification of groundwater quality based on TDS and TH is demonstrated in Fig. 8. It can be seen that all groundwater samples belong to the hard brackish and very hard saline categories.

In the histogram chart depicting the distribution of total dissolved solids (TDS) in the two aquifers (Fig. 9a), it is evident that the salinity of groundwater is higher in samples obtained from Southern Galala compared to those from Northern Galala (Upper Cretaceous aquifer). This increase could be attributed to the leaching and dissolution processes of the aquifer limestone occurring during the flow of groundwater. Additionally, the salinity shows a further escalation toward the Gulf of Suez, which is associated with seawater intrusion in the Nubian sandstone aquifer.

The concentration of major cations, including Ca^{++} , Mg^{++} , Na^+ and K^+ , in the examined groundwater are detailed in Table 1. Calcium and magnesium ions

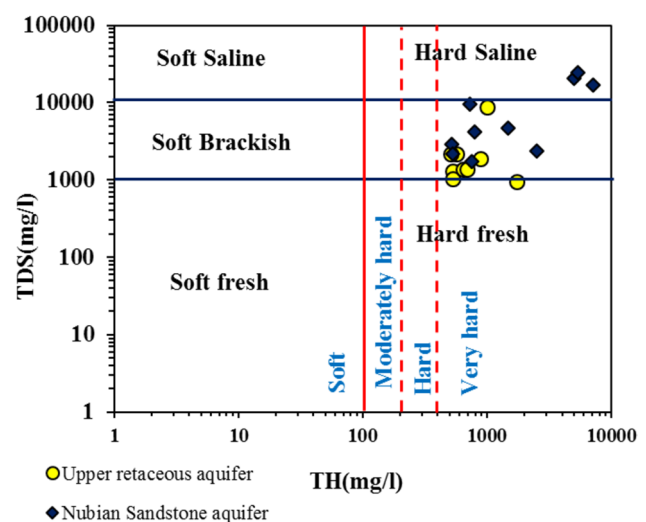


Fig. 8 a Distribution of collected water samples and b scatter plots of TDS versus TH showing groundwater quality

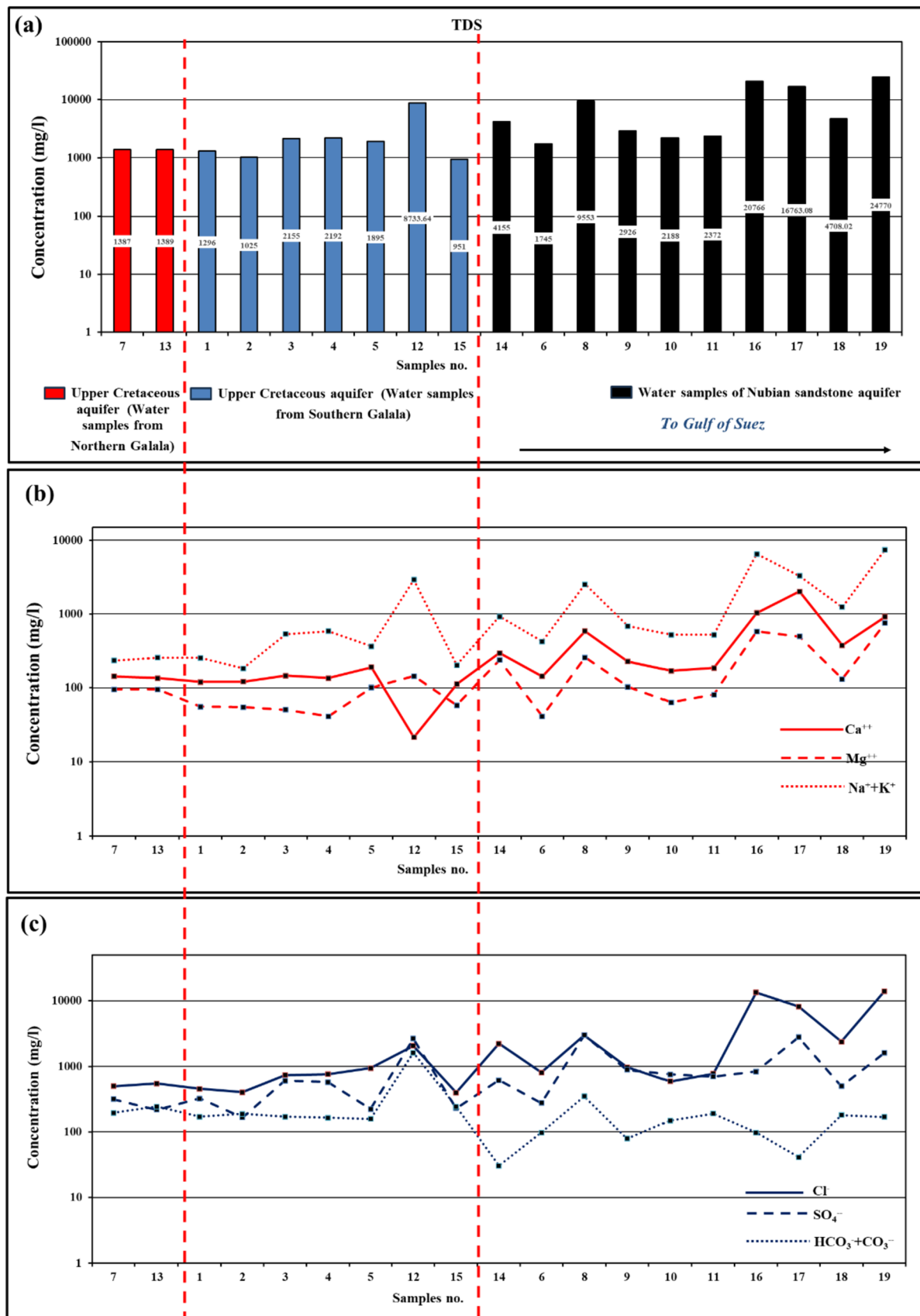


Fig. 9 a TDS distribution of collected water samples based on their presence also, b Major cations and c Major anions

predominantly originate from processes such as calcareous leaching, dolomites, gypsum, and anhydrites. Additionally, calcium ions result from the cation exchange cycle (Garrels 1976). The Ca^{++} concentration exhibits variability, ranging from 21.4 mg/l for sample 12 to 2010.3 mg/l for sample 17, with an average of 371.6 mg/l (Table 1 and Fig. 9b). Magnesium concentration varies from 41.1 mg/l in sample 6 to 751.3 mg/l in sample 19, averaging 181.1 mg/l (Table 1 and Fig. 9b).

Sodium, a crucial ion in natural water due to its impact on soil structure and its direct association with salinity primarily occurs in clay minerals and evaporates. Sodium concentration spans from 180 mg/l in sample 2 to 7200 mg/l in sample 19, averaging 1527.4 mg/l (Table 1 and Fig. 9b). Potassium, derived from K-feldspar weathering and aquifer matrix clay minerals, has lower concentration in groundwater compared to sodium, as potassium minerals exhibit greater resistance to weathering (El-Aziz 2017). The concentration of potassium ranges from 3 mg/l in samples 1, 2, 7, 10, 11, and 15 to 264 mg/l in sample 19, with an average of 29.3 mg/l (Table 1 and Fig. 9b).

The primary anion, chloride, originates from the dissolution of sodium chloride in rocks and soil within water. It serves as a significant indicator of groundwater quality, with its concentration increasing in groundwater when combined with wastewater or seawater (Deshpande and Aher 2012). The chloride concentration in the investigated groundwater ranges from 13946.8 mg/l in sample 19 to 397 mg/l in sample 15, averaging 2793.2 mg/l (Table 1 and Fig. 9c).

Sulfate is chiefly derived from sedimentary rocks, particularly gypsum, anhydrite, and certain types of shales. The measured sulfate concentration varies from 166.3 mg/l in sample 2 to 3001.2 mg/l in sample 8, with an average of 909.6 mg/l (Table 1 and Fig. 9c). Bicarbonates, likely originating from the dissolution of carbonate sediments, play a role in groundwater composition. Another potential source is the rapid dissolution during initial infiltration, as soil produces CO_2 , which dissociates into carbonic acid, impacting the groundwater's ability to dissolve calcium carbonate (Dyke 1999). The presence of bicarbonates ranges from 30.5 mg/l in sample 14 to 1327.2 mg/l in sample 12, with an average of 216.6 mg/l (Table 1 and Fig. 9c).

In general, the spatial trend of the concentration changes of ions like TDS. Ion concentration in carbonate aquifer is higher in southern Galala Springs compared to Northern Galala. Ion concentration of the Nubian Sandstone Aquifer is higher than that of the carbonate and increase toward the Gulf of Suez. The higher concentration in sandstone aquifers could be related to seawater intrusion and or evaporation (Salem and Osman 2017a, b; Hasan et al. 2023 and Salem et al. 2021a and 2022a). Carbonate concentrations carry the aquifer lithology impact. Although the higher TDS and ions concentrations in the sandstone aquifer, the higher

concentration of carbonate ions were recorded in the carbonate aquifer which might related to dissolution processes (Abdelazeem et al. 2021).

Samples classification

Statistical classification The studied groundwater samples were classified into five groups to describe the characteristics of each group of samples according to the degree of similarity between the groundwater samples with respect to each other which depends on the major ion concentration in meq/l (Figs. 10 and 11). Table 3 presents the average values of hydrochemical data for each group.

Samples from group A (samples 16, 17 and 19) have the highest TDS average value (20,766 mg/l) and $\text{Na}^+ - \text{Cl}^-$ groundwater type ($\text{Na}^+ + \text{K}^+ > \text{Ca}^{++} > \text{Mg}^{++}$ and $\text{Cl}^- > \text{SO}_4^{--} > \text{HCO}_3^-$) (Table 3, 4 and Fig. 11). These samples belong to Nubian Sandstone aquifer and of high salinity as it is affected by the seawater intrusion from Gulf of Suez. Samples from group D (samples 1, 2, 7, 13, and 15) have the lowest TDS average value (2127.4 mg/l) and $\text{Na}^+ - \text{Cl}^-$ groundwater type ($\text{Na}^+ + \text{K}^+ > \text{Ca}^{++} > \text{Mg}^{++}$ and $\text{Cl}^- > \text{SO}_4^{--} > \text{HCO}_3^-$) (Tables 3, 4 and Fig. 11). These samples belong to Upper Cretaceous aquifer and its low salinity could be related to rainwater source. Samples from groups B, C, and E (samples 3, 4, 5, 6, 8, 9, 10, 11, 12, 14, and 18) have TDS average values of 9143.3, 4431.5, 2210.4 mg/l, respectively, and of $\text{Na}^+ - \text{Cl}^-$ groundwater type ($\text{Na}^+ + \text{K}^+ > \text{Ca}^{++} > \text{Mg}^{++}$ and $\text{Cl}^- > \text{SO}_4^{--} > \text{HCO}_3^-$) for groups C and E but $\text{Na}^+ - \text{SO}_4^{--}$ groundwater type ($\text{Na}^+ + \text{K}^+ > \text{Ca}^{++} > \text{Mg}^{++}$ and $\text{SO}_4^{--} > \text{Cl}^- > \text{HCO}_3^-$) for group B (Tables 3,4 and Fig. 11).

Higher silicon concentration in samples of groups A, B, and C could be related to the clastic facies of the aquifer where most samples of the three groups belong to the Nubian sandstone aquifer. Al, Fe, and Mn are the most abundant trace elements in the investigated groundwater, and their average concentration (mg/l) in the four groups, respectively, are (0.19, 0.11, 0.21, 0.25, and 0.12), (3.53, 0.103, 1.05, 0.05, 0.01) and (0.73, 0.11, 5.45, 0.02 and 0.01). Encountered higher Fe and Mn could be related to the ferruginous characteristics of the Nubian Sandstone aquifer.

Graphical classification The groundwater samples labeled according to their respective group of HCA (Fig. 11) were presented in the Piper diagram (Fig. 11). Na^+ is the major cation in the majority of the groundwater samples and Cl^- is the main anion. In the right triangle, SO_4^{--} increases to reach its maximum concentration in groups B and E (samples 3, 4, 5, 6, 8, 9, 10, 11, and 12). In the left triangle, Ca^{++} and Mg^{++} increase, while Na^+ decreases indicating ion exchange where Ca^{++} and Mg^{++} replace Na^+ , especially in group D (samples 1, 2, 7,

Fig. 10 Dendrogram shows the grouping of the groundwater samples into five groups depending on the concentration of the ions in mg/l

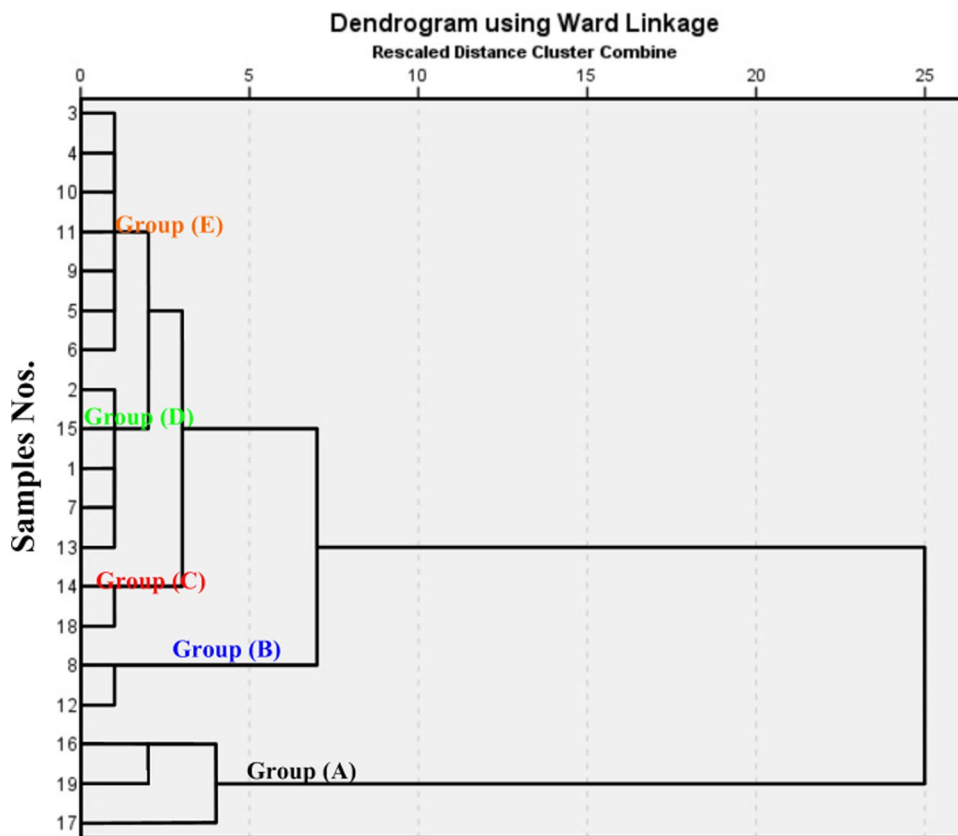
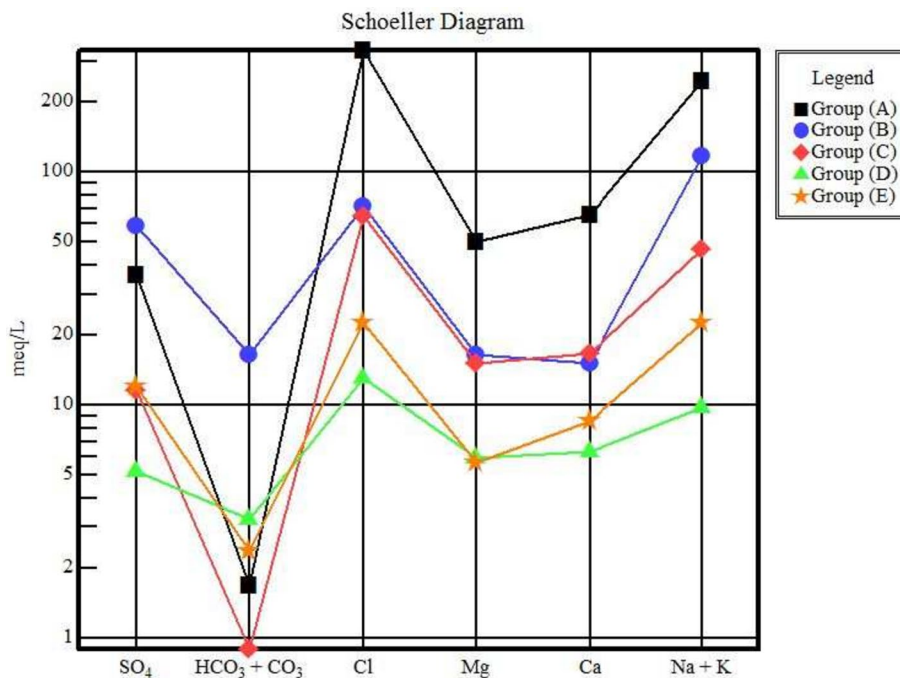


Fig. 11 Schoeller diagram of the obtained groundwater sample groups



13, and 15). Based on general patterns of plotted data of groundwater samples on the diamond field (Fig. 18), the majority of the groundwater samples (68%; samples 1, 3, 4, 6, 8, 9, 10, 11, 12, 14, 16, 18 and 19) occupy the sub-

zone (7) in which the water is characterized by primary salinity where Na⁺ and K⁺ exceed SO₄ and NaCl salt prevails compared to seawater (Fig. 12) (Rajendra et al. 2009). The other groundwater samples (32%; samples 2,

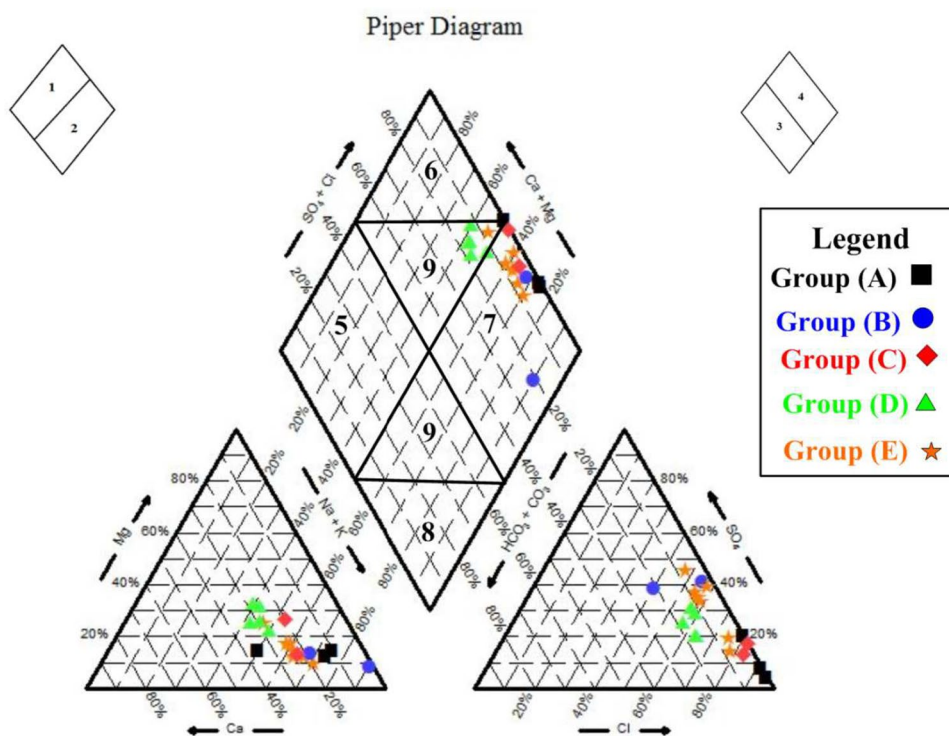
Table 3 Average composition of the major hydrochemical parameters of the classified groundwater sample groups

| Groups | T | pH | EC | TDS | Ca ⁺⁺ | Mg ⁺⁺ | Na ⁺ | K ⁺ | CO ₃ ⁻⁻ | HCO ₃ ⁻ | Cl ⁻ | SO ₄ ⁻⁻ |
|--|------|-----|----------|----------|------------------|------------------|-----------------|----------------|-------------------------------|-------------------------------|-----------------|-------------------------------|
| Units | °C | | µS/cm | | | | | | | | | |
| A (samples, 16, 17, and 19) | 26.0 | 8.0 | 33,503.7 | 20,766.4 | 1320.0 | 609.7 | 5600.0 | 138.7 | 6.8 | 95.7 | 11,838.1 | 1744.3 |
| B (samples, 12 and 8) | 23.7 | 8.4 | 13,952.2 | 9143.3 | 303.0 | 200.6 | 2700.0 | 14.5 | 159.0 | 825.8 | 2527.7 | 2825.6 |
| C (samples, 14 and 18) | 30.8 | 6.3 | 7588.5 | 4431.5 | 335.5 | 183.9 | 1050.0 | 31.5 | 0.0 | 105.9 | 2303.7 | 558.0 |
| D (samples, 1, 2, 7, 13, and 15) | 22.4 | 7.6 | 2127.4 | 1209.6 | 125.9 | 72.1 | 222.0 | 3.8 | 4.8 | 203.7 | 461.8 | 251.3 |
| E (samples, 3, 4, 5, 6, 9, 10, and 11) | 20.0 | 8.0 | 3767.6 | 2210.4 | 170.6 | 68.8 | 515.7 | 4.3 | 10.5 | 135.1 | 797.9 | 575.1 |

Table 4 Average composition of the trace elements of the classified groundwater sample groups

| Groups | Al | Cd | Co | Cr | Cu | Fe | Mn | Mo | Ni | Pb | V | Zn | Si |
|---------------------------------------|------|------|------|------|------|-------|------|------|------|------|------|------|-------|
| Units | mg/l | | | | | | | | | | | | |
| A (samples, 16, 17 and 19) | 0.19 | 0.00 | 0.00 | 0.00 | 0.01 | 3.53 | 0.73 | 0.00 | 0.00 | 0.00 | 0.00 | 0.60 | 16.61 |
| B (samples, 12 and 8) | 0.11 | 0.00 | 0.00 | 0.00 | 0.00 | 0.10 | 0.11 | 0.01 | 0.00 | 0.08 | 0.03 | 0.01 | 18.73 |
| C (samples, 14 and 18) | 0.21 | 0.00 | 0.00 | 0.02 | 0.00 | 31.05 | 5.45 | 0.00 | 0.01 | 0.00 | 0.00 | 0.69 | 14.72 |
| D (samples, 1, 2, 7, 13 and 15) | 0.25 | 0.00 | 0.00 | 0.00 | 0.01 | 0.05 | 0.02 | 0.01 | 0.00 | 0.07 | 0.02 | 0.02 | 5.94 |
| E (samples, 3, 4, 5, 6, 9, 10 and 11) | 0.12 | 0.00 | 0.00 | 0.00 | 0.00 | 0.01 | 0.01 | 0.00 | 0.00 | 0.07 | 0.01 | 0.01 | 4.54 |

Fig. 12 Piper diagram of the obtained groundwater sample groups

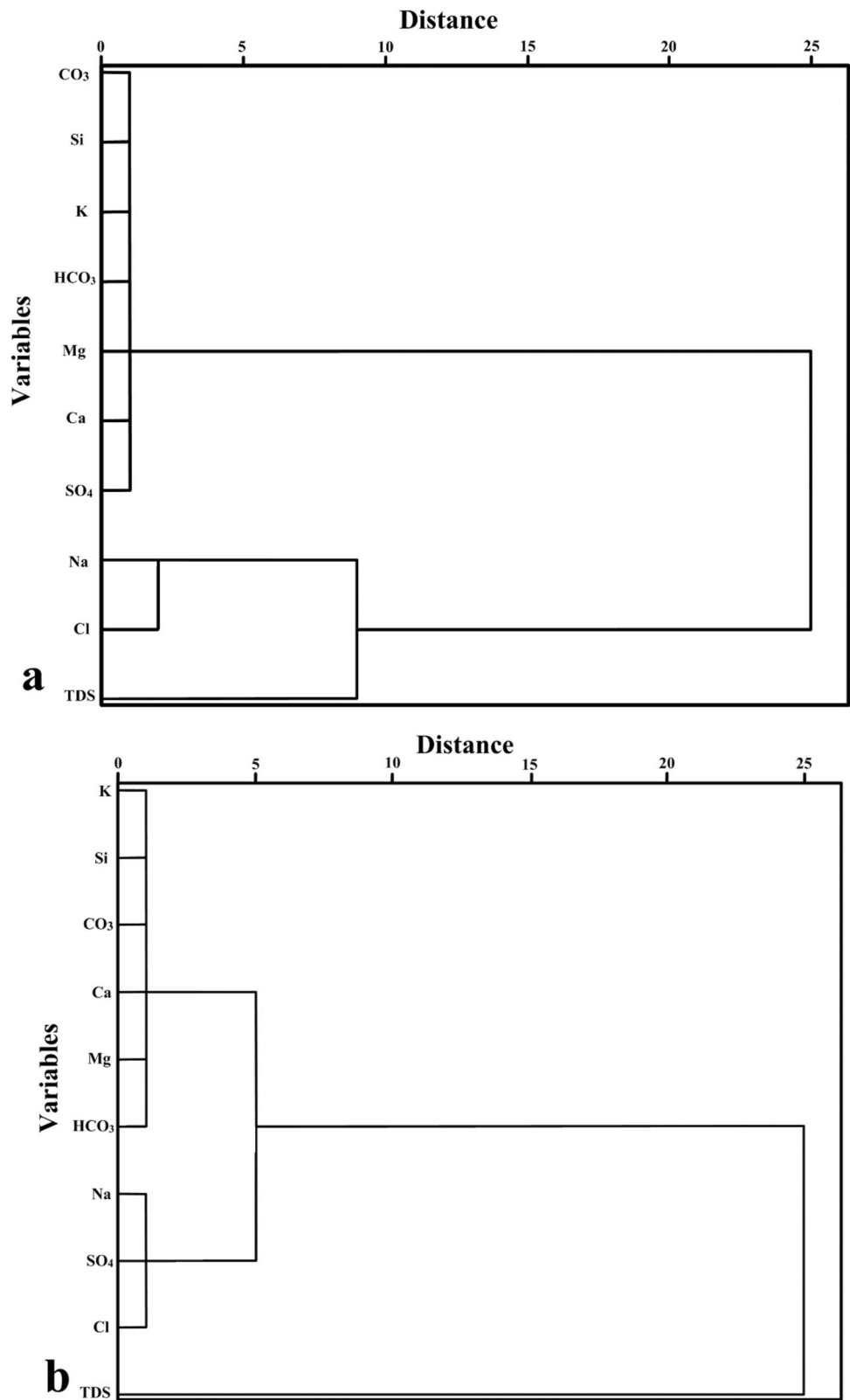


5, 7, 13, 15, and 17) occupy the subzone (9), in which none of the cation–anion pairs exceeds 50% (Rajendra et al. 2009). The plot displays that the chemistry of the groundwater belongs essentially to CaMgSO₄ and NaCl.

Ionic relationships

Statistical analysis The dendrograms used for classifying the ion association in the studied groundwater for the two

Fig. 13 Dendrograms show the major ion grouping in the Nubian Sandstone groundwater aquifer (a) and the Upper Cretaceous aquifer (b)



aquifers are shown in Fig. 13a and b. There are two groups of chemicals in the Nubian Sandstone groundwater aquifer system: SO_4 -Ca-Mg- HCO_3 -K-Si- CO_3 and Cl-Na. The

first group could be related to water-rock interaction, and the second is related to either seawater intrusion or evaporation. The SO_4 -Ca association is located close to Na-Cl

because either seawater intrusion or evaporation is accompanied by an increase in Ca and SO₄, while the HCO₃-K-Si-CO₃ subgroup is located far from Na-Cl because it is just related to water-rock interaction. Compared to the Nubian Sandstone Aquifer, the Upper Cretaceous aquifer shows two different ion associations: Na-Cl-SO₄ and Ca-Mg-HCO₃-K-Si-CO₃. The association between Na and SO₄ indicates ion exchange between Na of rocks and Ca of dissolved gypsum, while K-HCO₃-Ca-Mg indicates a freshwater source (Salem and Osman 2017a).

Based on Pearson’s method, the correlation between the groundwater’s major chemical variables was determined as shown in Table 5. A strong correlation ($r \geq 0.75$) was recognized between TDS and Ca, Mg, Na, K, and Cl; Na and Cl and K; Mg and Na, Cl and K; CO₃ with HCO₃⁻, while moderate correlation (0.75–0.5) was noticed between several water specifications such as between TDS and SO₄ and Si; Ca and Na, K and SO₄; Mg and SO₄; Na and SO₄ and Si; K and Si; CO₃ and SO₄ and Si; HCO₃ and Si. The rest of the relationships are weak (<0.5). From this analysis, we can conclude that Ca, Mg, Na, K, Cl, and Si are the main component affecting groundwater salinity.

Binary relationships TDS versus major ions: The relationships of TDS versus major ions (Ca, Mg, Na, HCO₃, SO₄, and Cl) are shown in Fig. 13. TDS versus Ca relationship (Fig. 14a) is strong ($r=0.784$, Table 3). Samples related to the Upper Cretaceous carbonate aquifer and the low salinity samples of the Nubian sandstone aquifer are plotted just below the rainwater-seawater mixing line indicating the meteoric origin of Ca in such samples. On the other hand, Ca concentration in the Nubian aquifer samples with higher salinity showed higher values than that of the mixing line which means the addition of Ca either by weathering of silicate minerals or ion exchange with Na (Ibrahim and Lyons 2017). TDS versus Mg, Cl, and Na + K relationships (Fig. 14b, e, and f) are significant with a higher correlation

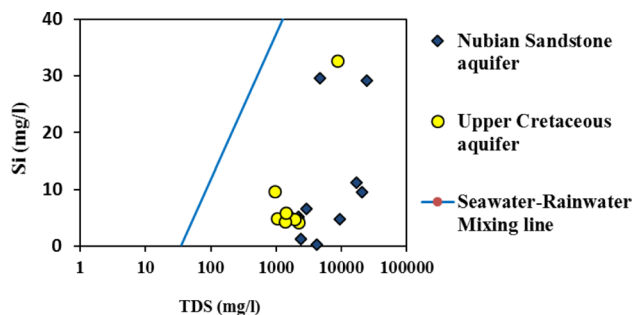


Fig. 14 TDS-Si relationship

coefficient (Table 3). All the samples were plotted below and parallel to the mixing line. This means the increase in Na + K, Cl, and Mg might be related to seawater intrusion or evaporation, especially in the Nubian aquifer samples. Also, ion exchange processes might be responsible for the lower concentration compared to that of the mixing line where Na and K were consumed from the groundwater by ion exchange with calcium (Kumar et al. 2009). The TDS versus HCO₃ and SO₄ relationships (Fig. 14c and d) are weak and intermediate, respectively (Table 5). The samples in the TDS-HCO₃ relationship are mostly clustered around the mixing line while in the TDS-SO₄ relationship sample is plotted mostly under the mixing line. This indicates that the dissolution of gypsum is accompanied by carbonate precipitation (Salem and Elhoriény, 2014). TDS versus Si relationship (Fig. 15) is intermediate ($r=0.509$, Table 5). If TDS would have resulted from water-rock interaction, a positive correlation between SiO₂ and TDS would have been logically expected (Khan and Umar, 2010).

Na versus Cl: The parameter Na/Cl reflects the inequity of sodium chloride in water. This differs greatly from the difference in maximum dissolved solid concentration. Water is meteoric when this ratio exceeds unity and marine when it is less than unity (about 0.8) (Salem and Osman

Table 5 Correlation coefficient between groundwater parameters

| Variables | TDS | Ca ⁺⁺ | Mg ⁺⁺ | Na ⁺ | K ⁺ | CO ₃ ⁻⁻ | HCO ₃ ⁻ | Cl ⁻ | SO ₄ ⁻⁻ | Si |
|-------------------------------|---------|------------------|------------------|-----------------|----------------|-------------------------------|-------------------------------|-----------------|-------------------------------|----|
| TDS | 1 | | | | | | | | | |
| Ca ⁺⁺ | 0.784** | 1 | | | | | | | | |
| Mg ⁺⁺ | 0.976** | 0.798** | 1 | | | | | | | |
| Na ⁺ | 0.981** | 0.657** | 0.945** | 1 | | | | | | |
| K ⁺ | 0.858** | 0.570* | 0.890** | 0.852** | 1 | | | | | |
| CO ₃ ⁻⁻ | 0.114 | -0.183 | -0.029 | 0.178 | -0.056 | 1 | | | | |
| HCO ₃ ⁻ | 0.036 | -0.261 | -0.112 | 0.11 | -0.123 | 0.965** | 1 | | | |
| Cl ⁻ | 0.975** | 0.761** | 0.976** | 0.966** | 0.866** | -0.037 | -0.112 | 1 | | |
| SO ₄ ⁻⁻ | 0.598** | 0.577** | 0.502* | 0.534* | 0.322 | 0.506* | 0.452 | 0.414 | 1 | |
| Si | 0.509* | 0.194 | 0.412 | 0.547* | 0.532* | 0.577** | 0.570* | 0.427 | 0.41 | 1 |

** Correlation is significant at the 0.01 level (two-tailed)

* Correlation is significant at the 0.05 level (two-tailed)

Fig. 15 TDS–major ions relationships

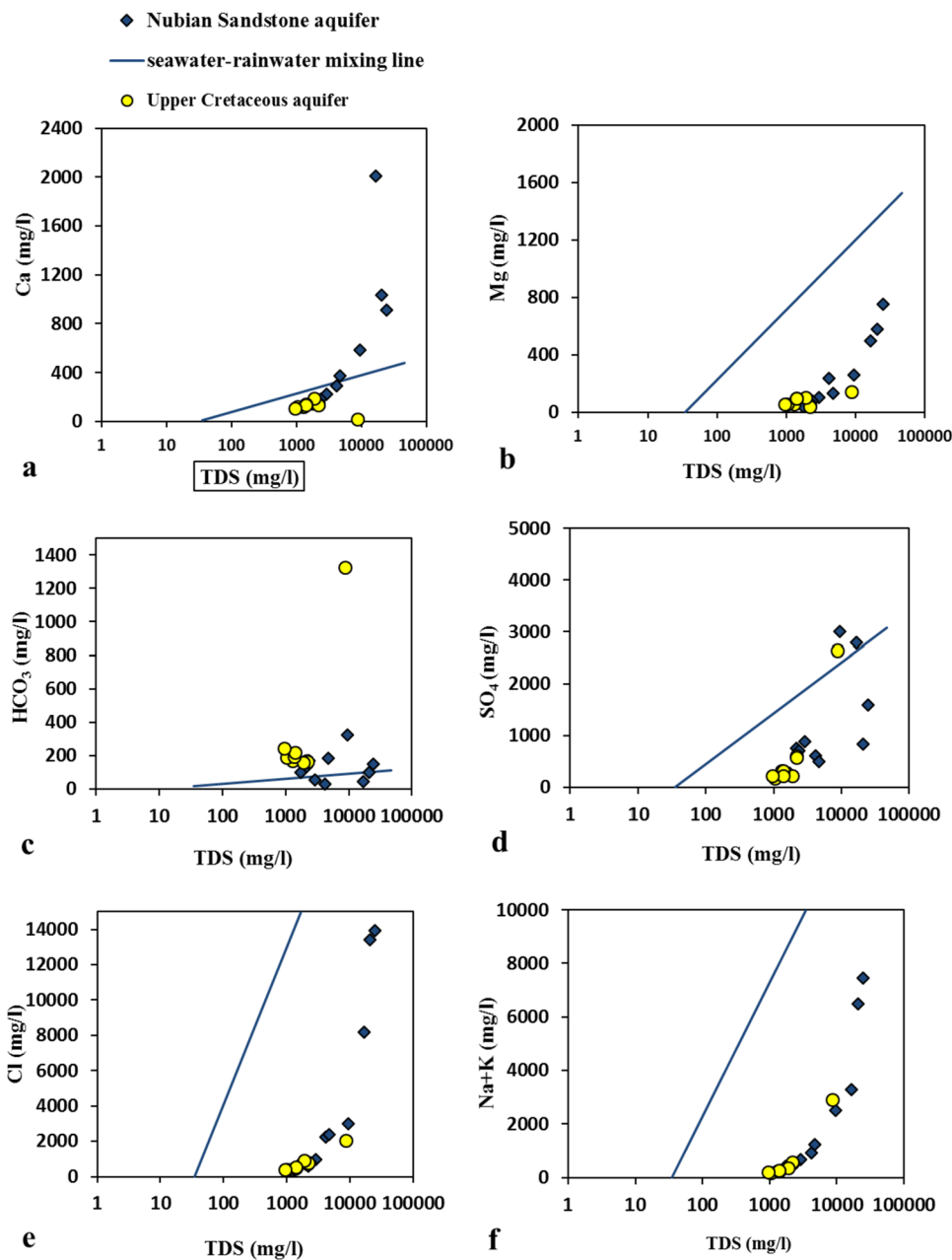
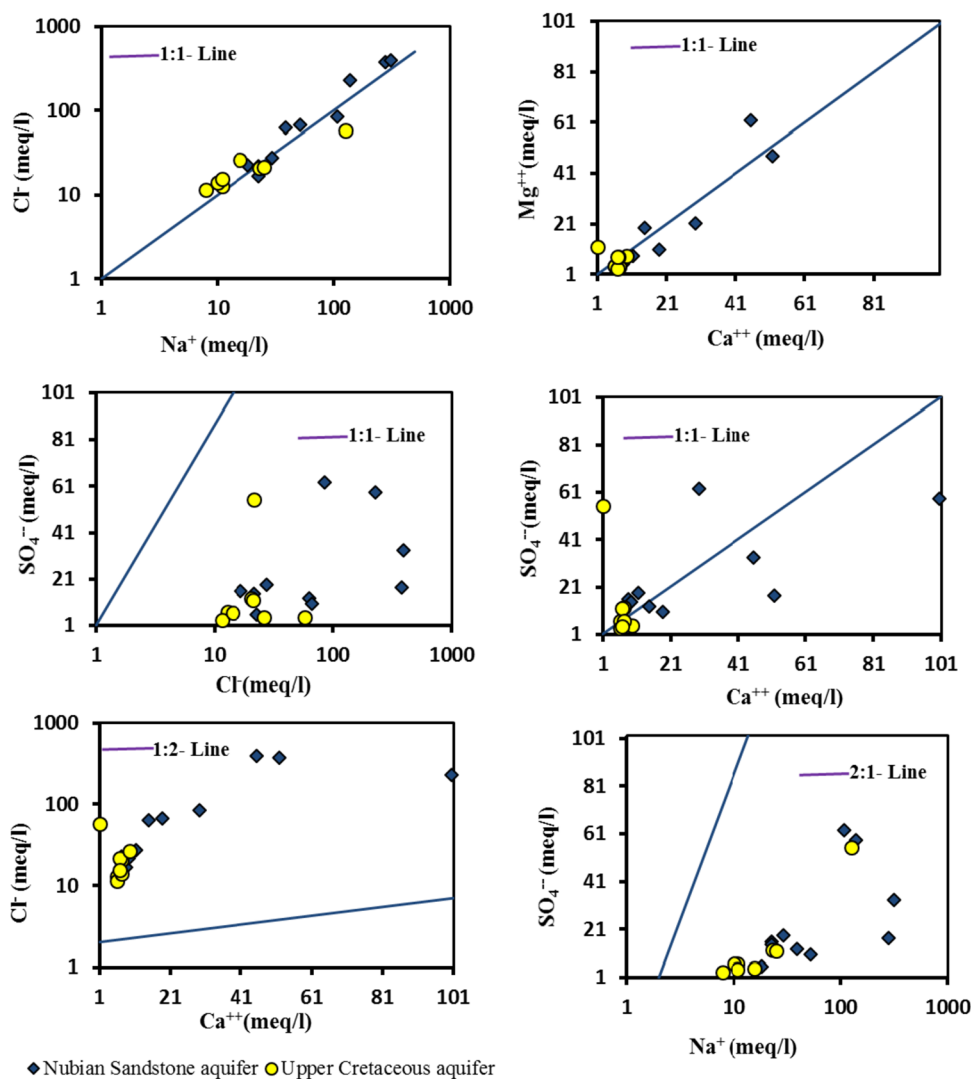


Table 6 Statistical summary of the proposed ionic ratios of the collected samples and that of seawater and rainwater

| Sample no | Na/Cl | Ca/Mg | SO ₄ /Cl | Ca/SO ₄ | Ca/Cl | Ca/HCO ₃ | Mg/HCO ₃ | SO ₄ /HCO ₃ |
|------------|-------|-------|---------------------|--------------------|-------|---------------------|---------------------|-----------------------------------|
| Max | 2.26 | 2.46 | 0.95 | 3.12 | 0.52 | 151.17 | 61.56 | 85.74 |
| Min | 0.60 | 0.09 | 0.05 | 0.02 | 0.02 | 0.05 | 0.56 | 1.12 |
| STDV | 0.39 | 0.55 | 0.28 | 0.72 | 0.14 | 34.32 | 16.89 | 19.35 |
| Average | 0.95 | 1.33 | 0.44 | 1.17 | 0.35 | 15.17 | 10.38 | 11.01 |
| > 1 | 7 | 14 | 0 | 11 | 0 | 18 | 18 | 19 |
| > 1% | 37 | 74 | 0 | 58 | 0 | 95 | 95 | 100 |
| < 1 | 12 | 5 | 19 | 8 | 19 | 1 | 1 | 0 |
| < 1% | 63 | 26 | 100 | 42 | 100 | 5 | 5 | 0 |
| Seawater | 0.88 | 0.19 | 0.08 | 0.39 | 0.03 | 13.74 | 72.09 | 35.53 |
| Rain water | 1.04 | 5.65 | 0.73 | 4.25 | 3.10 | 1.05 | 0.19 | 0.25 |

Fig. 16 Ionic relationships to estimate the hydrogeochemical processes that occurred in the studied groundwater



2017b). Most samples (63%) are less than 1 (Table 6) reflecting the marine origin and/or evaporation effect in the groundwater. The samples are plotted in Fig. 16a above and below the unit line. The correlation coefficient (r) is 0.966 (Table 5). Sodium excess could be related to either ion exchange and/or addition from the terrestrial origin (Fisher and Mullican 1997; Lee and Song 2007; Salem and Osman 2017a).

Ca versus Mg: The Ca/Mg ratio benefits from any pollution of seawater. Indication of dissolution of calcium minerals and rocks, contamination of surface water, and contact between CO_2 and CaCO_3 depends on the existence of high values of this ratio (Salem and Osman 2017b). Nevertheless, the operation of the base-exchange processes is strongly influenced by this ratio. 74% of water samples have Ca/Mg ratio higher than 1 (Table 6) reflecting anhydrite and gypsum dissolution. Samples with a lower-unit Ca/Mg ratio (Table 6) may be a feature of intrusion of seawater (Salem and Osman 2017b). The collected samples were usually distributed

with a high correlation ($r=0.798$) in the dispersed diagram (Fig. 16b, Table 5).

SO₄ versus Cl: The SO_4/Cl ratio assists in determining the dissolution effect of halite, gypsum, or anhydrite. The SO_4/Cl ratios above unity reflect the excess of sulfate than chloride and this may reflect the presence of sulfate minerals in the formation that contains water samples. All the collected water samples have SO_4/Cl ratios below unity (Table 6), reflecting the excess of chloride over sulfate which may indicate the abundance of halite in water-bearing formation deposits, seawater intrusion, and/or evaporation effect (Salem and Osman 2017b). The collected samples are scattered in the binary diagram (Fig. 16c) with a weak correlation ($r=0.414$, Table 5), revealing different sources of both SO_4 and Cl.

Ca versus SO₄: Most samples collected (58%) have Ca/ SO_4 ratios higher than the unit (Table 6). Ca excess in these samples indicates additional calcium by weathering minerals other than Ca-rich gypsum (Salem and Osman

2017a; Desbarats 2009; Jalali 2009). Samples with lower Ca/SO₄ ratios than 1 (42%) suggest the removal of calcium either through precipitation as calcium carbonates in and/or by ion exchange processes (Desbarats 2009; Jalali 2009; Hamzaoui-Azaza et al. 2011). The plotted samples (Fig. 16d) are scattered with moderate correlation ($r=0.577$, Table 5).

Ca versus Cl: The Ca/Cl ratio helps to change the composition of water samples by leaching and dissolving various salts such as calcite, dolomite, gypsum, and anhydrite (Fisher and Mullican 1997; Salem and Osman 2017a and Salem et al. 2015). All the collected water samples have Ca/Cl ratios below unity (Table 6). The Ca/Cl relationship (Fig. 16e) shows a scattered pattern with high correlation coefficients ($r=0.761$, Table 5). Plotted water samples are

scattered mostly over the 1:2 line and this ratio confirms the seawater intrusion, especially in samples with high salinity.

HCO₃ versus ions: HCO₃ has a weak relation to Ca and Mg ($r=-0.261, -0.112$, respectively) as shown in Table 5 and Fig. 17a and b. HCO₃ is moderately related to SO₄, and weakly related to Cl⁻ and Na⁺, where r equals 0.452, -0.112 , and 0.11, respectively (Table 5 and Fig. 17c, d, and e). All the collected water samples have SO₄/HCO₃ ratios above unity but most of the collected samples (95%) have Ca/HCO₃ and Mg/HCO₃ ratios above unity (Table 6). Such low concentration of HCO₃ might be related to either calcite precipitation or reduction of carbonate minerals dissolution due to water enrichment with Ca and Mg due to silicate or/and sulfate minerals dissolution (Fisher and Mullican 1997). As shown in Fig. 18a, Ca + Mg concentration has no obvious

Fig. 17 Ionic relationships between HCO₃⁻ and other major ions

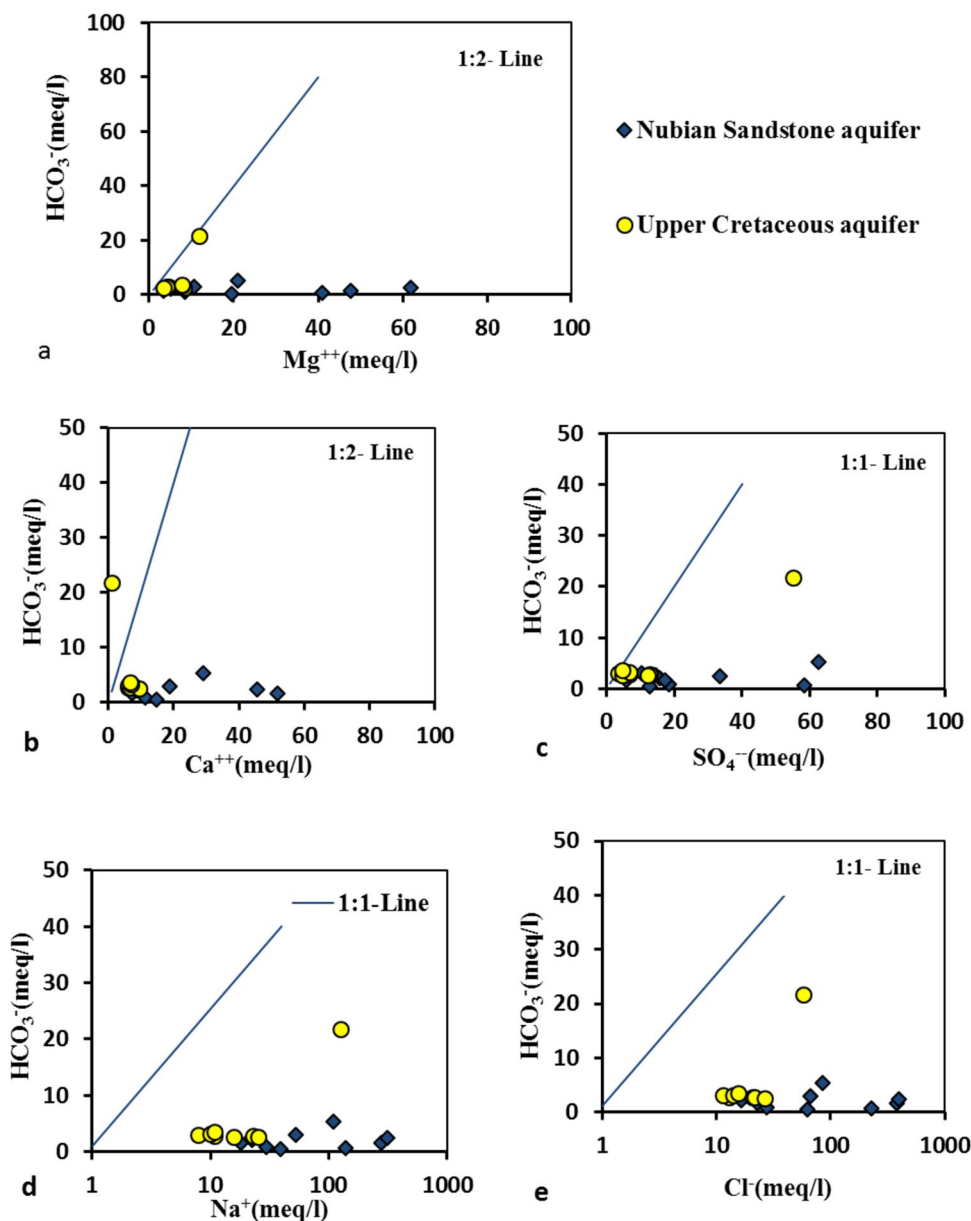
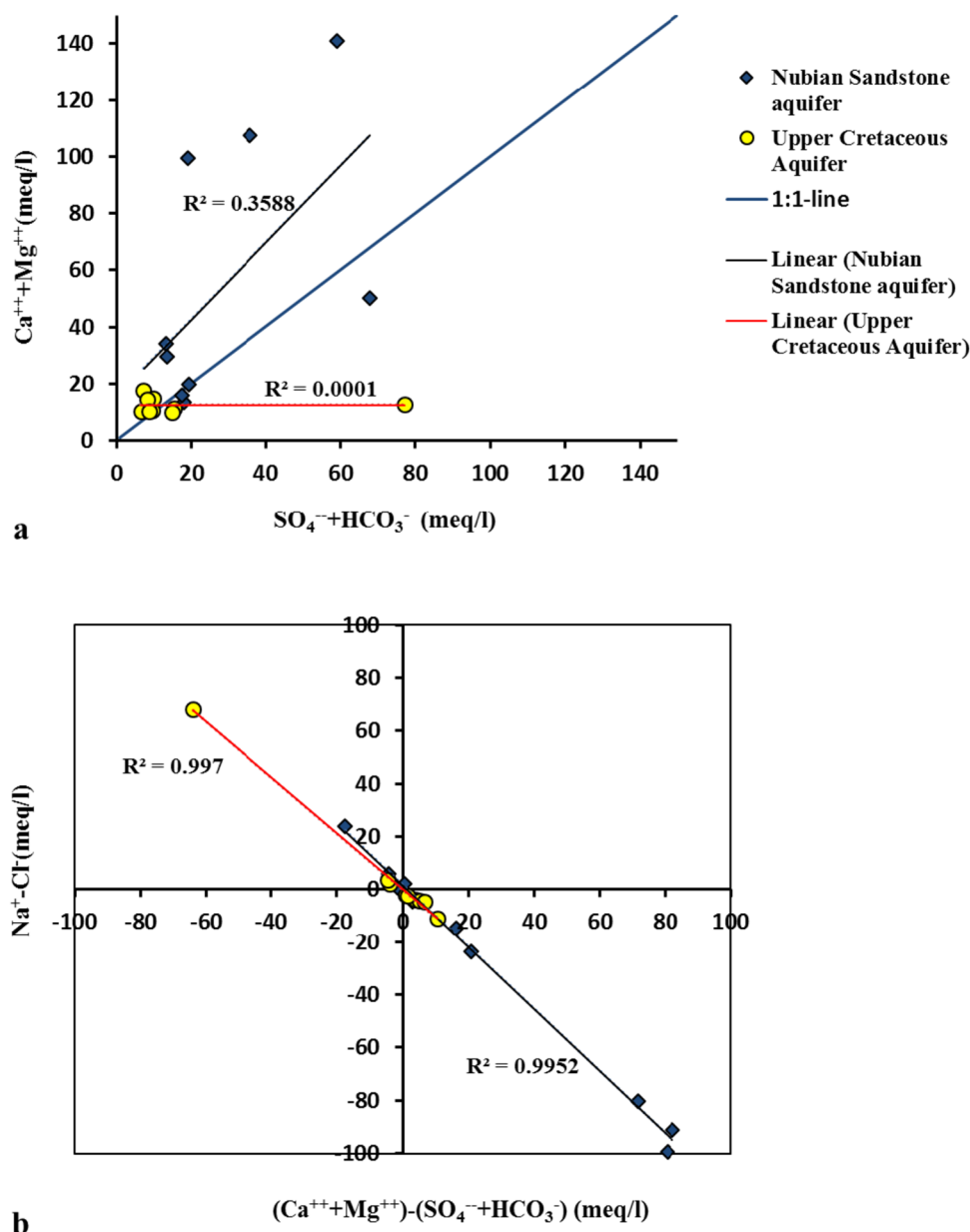


Fig. 18 Relationship between $\text{SO}_4 + \text{HCO}_3$ and $\text{Ca} + \text{Mg}$ in the dissolved sulfate and carbonate minerals **a** and a Binary diagram illustrates the ion exchange processes in the studied groundwater **b**.

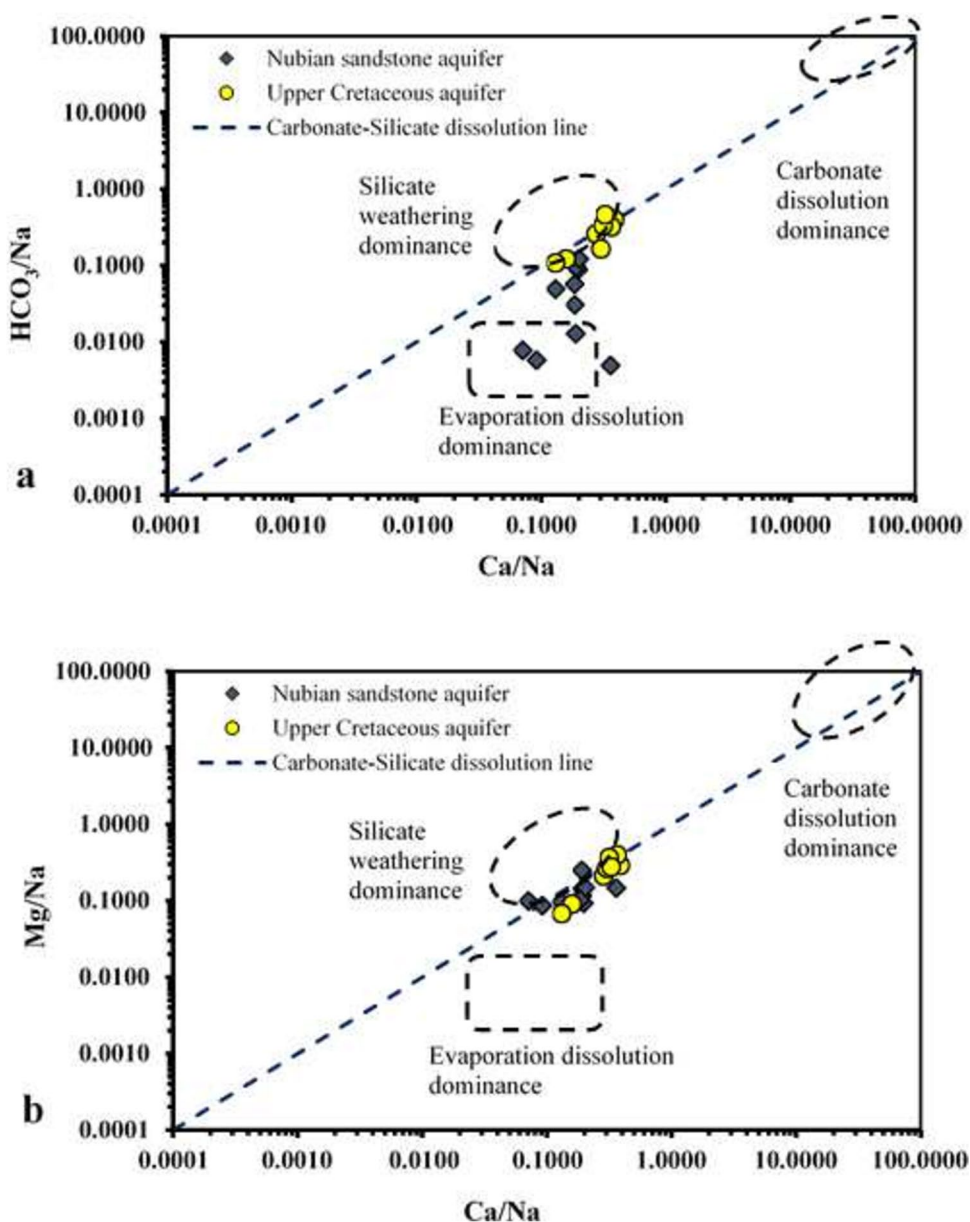


relationships with $\text{HCO}_3 + \text{SO}_4$ ($r^2 = 0.3588$ and 0.0001 for the Nubian Sandstone aquifer and the Upper Cretaceous aquifer, respectively). This relationship means that Ca and Mg are not solely derived from the dissolution of gypsum, calcite, dolomite, and anhydrite, but may also be associated with weathering of silicate minerals. The ion exchange of Na and calcite and dolomite precipitations due to sulfate dissolution often extract Ca and Mg from the solution (Fisher and Mullican 1997; Salem and Osman 2017a).

Figure 18b indicates that ion exchange is occurring in the systems with a plot of $(\text{Ca} + \text{Mg}) - (\text{SO}_4 + \text{HCO}_3)$ versus $\text{Na} - \text{Cl}$. By subtracting Cl from Na and assuming that all the Cl comes from NaCl, groundwater samples not affected by ion exchange will plot close to zero on the y-axis. The x-axis

of $(\text{Ca} + \text{Mg}) - (\text{SO}_4 + \text{HCO}_3)$ shows the amount of Ca and Mg given by congruent calcite, dolomite, and gypsum dissolution. If there are, groundwater is plotted on a straight line with a high correlation coefficient (exchange reactions in the process Jankowski et al. 1998). Data from the study area (Fig. 18b) have r^2 values of 0.9952 and 0.997 for the Nubian Sandstone aquifer and the Upper Cretaceous aquifer, respectively. From this information, it could be estimated that ion exchange reaction is a predominant process in both aquifers (Salem and Osman 2017a). To identify the relative contributions of the major weathering/dissolution mechanisms (silicate, carbonate, and evaporite) to the ion concentration in groundwater, the Na-normalized molar ratio model ($\text{Ca}/\text{Mg}/\text{HCO}_3$) suggested by Gaillardet et al. (1999) was employed

Fig. 19 Molar ratio bivariate plots of **a** Na-normalized Ca and HCO_3^- and **b** Na-normalized Ca and Mg. The dashed circles represent the ranges of approximate compositions of the silicate and carbonate end members, and a dashed box represents the evaporite end member. Most groundwater samples fell in the area of silicate end member area, indicating that the majority of ions originated from silicate mineral weathering



(Fig. 19). The bivariate mixing plots (Fig. 19a and b) of Na-normalized HCO_3^- versus Ca and Na-normalized Mg versus Ca showed that most of the groundwater samples of the two aquifers were within or very close to the silicate weathering domain, which indicates a significant contribution of silicate mineral weathering dissolution to the groundwater chemistry of these samples. However, some of the Nubian Sandstone groundwater samples plotted in Fig. 19b have been affected by evaporation dominance and might be seawater intrusion.

Equilibrium modeling (saturation indices)

The degree of mineral saturation in waters is determined by the percentage of saturation, defined as the saturation index (SI) = $\log(\text{IAP}/K_s)$, where IAP is the result of the ionic

activity of the appropriate ions, and K_s is the product of mineral solubility. The hydrochemical program PHREEQC Interactive Program (USGS 2011) has been used to measure the distribution of species equilibrium in an aqueous solution and the saturation level of related minerals. The composition of groundwater is determined by the composition of sedimentary rocks in the basin and by hydrological characteristics such as the direction and time of residence. Table 7 and Fig. 20 show the calculated mineral saturation indices revealed that most of the collected groundwater samples are undersaturated with respect to carbonate minerals (aragonite, calcite, dolomite, and siderite), silicate minerals (albite, K-feldspar and anorthite), and chlorite, and all the obtained water samples are undersaturated for evaporates minerals (anhydrite, gypsum, halite, and sylvite) and melanterite.

Table 7 Statistical summary of the saturation indices of minerals of the collected water samples of two aquifers

| Groups | Minerals/Statistics | Aquifer | Minimum | Maximum | Average | Standard deviation | Aquifer | Minimum | Maximum | Average | Standard deviation |
|---------------------|---------------------|--|---------|---------|---------|--------------------|--|---------|---------|---------|--------------------|
| Silicate minerals | Albite | Upper Cretaceous aquifer water samples | -1.89 | 1.83 | -0.65 | 1.15 | Nubian sandstone aquifer water samples | -4.47 | 1.15 | -1.32 | 1.69 |
| | Alunite | | 0.65 | 3.30 | 2.06 | 1.33 | | -1.16 | 3.70 | 2.50 | 1.75 |
| | Anorthite | | -2.53 | -0.01 | -1.50 | 1.03 | | -3.61 | 0.16 | -1.88 | 1.44 |
| | K-feldspar | | -1.47 | 1.60 | -0.25 | 0.89 | | -3.68 | 1.54 | -0.97 | 1.63 |
| | Quartz | | -0.17 | 0.74 | 0.01 | 0.29 | | -1.35 | 0.74 | -0.03 | 0.62 |
| | Talc | | -5.55 | -1.06 | -4.00 | 1.30 | | -8.09 | 1.36 | -3.30 | 2.97 |
| | Sepiolite | | -6.58 | -3.29 | -5.49 | 0.96 | | -11.57 | -1.69 | -5.32 | 2.81 |
| Clay minerals | Ca-Montmorillonite | 2.50 | 5.88 | 4.20 | 2.47 | -0.52 | 5.42 | 2.77 | 2.11 | | |
| | Chlorite | -5.40 | -2.08 | -3.35 | 2.00 | -7.22 | 0.95 | -3.69 | 2.86 | | |
| | Illite | 1.41 | 5.05 | 3.21 | 2.03 | -0.78 | 5.13 | 2.06 | 2.00 | | |
| | K-mica | 8.01 | 11.81 | 10.00 | 5.39 | 6.41 | 12.36 | 9.07 | 4.22 | | |
| | Kaolinite | 4.50 | 7.03 | 5.72 | 3.11 | 3.32 | 6.51 | 4.82 | 2.28 | | |
| Carbonate minerals | Aragonite | -0.55 | -0.19 | -0.34 | 0.11 | -1.05 | 0.23 | -0.31 | 0.37 | | |
| | Calcite | -0.40 | -0.05 | -0.19 | 0.11 | -0.91 | 0.38 | -0.16 | 0.37 | | |
| | Dolomite | -0.72 | 0.35 | -0.24 | 0.34 | -1.57 | 0.73 | -0.33 | 0.75 | | |
| | Goethite | 6.37 | 7.15 | 6.76 | 2.99 | 5.15 | 9.49 | 7.18 | 3.67 | | |
| Iron oxide minerals | Hematite | 14.75 | 16.31 | 15.53 | 6.86 | 12.30 | 20.98 | 16.36 | 8.26 | | |
| | Jarosite-K | -3.85 | -3.52 | -3.69 | 1.63 | -8.25 | 4.76 | -1.69 | 3.83 | | |
| | Melanterite | -6.62 | -6.57 | -6.60 | 2.91 | -8.51 | -4.10 | -6.26 | 3.26 | | |
| | Siderite | -0.67 | -0.67 | -0.67 | 0.30 | -8.67 | -0.29 | -2.47 | 2.63 | | |
| Evaporate minerals | Halite | -5.75 | -3.97 | -5.25 | 0.56 | -5.16 | -2.79 | -4.17 | 0.91 | | |
| | Sylvite | -7.10 | -6.11 | -6.76 | 0.32 | -6.97 | -3.83 | -5.62 | 1.11 | | |
| | Anhydrite | -1.81 | -1.18 | -1.49 | 0.20 | -1.45 | -0.07 | -0.87 | 0.40 | | |
| Gibbsite minerals | Gypsum | -1.51 | -0.88 | -1.18 | 0.20 | -1.15 | 0.23 | -0.57 | 0.40 | | |
| | | 1.94 | 2.92 | 2.32 | 1.25 | 1.35 | 2.61 | 2.21 | 1.01 | | |

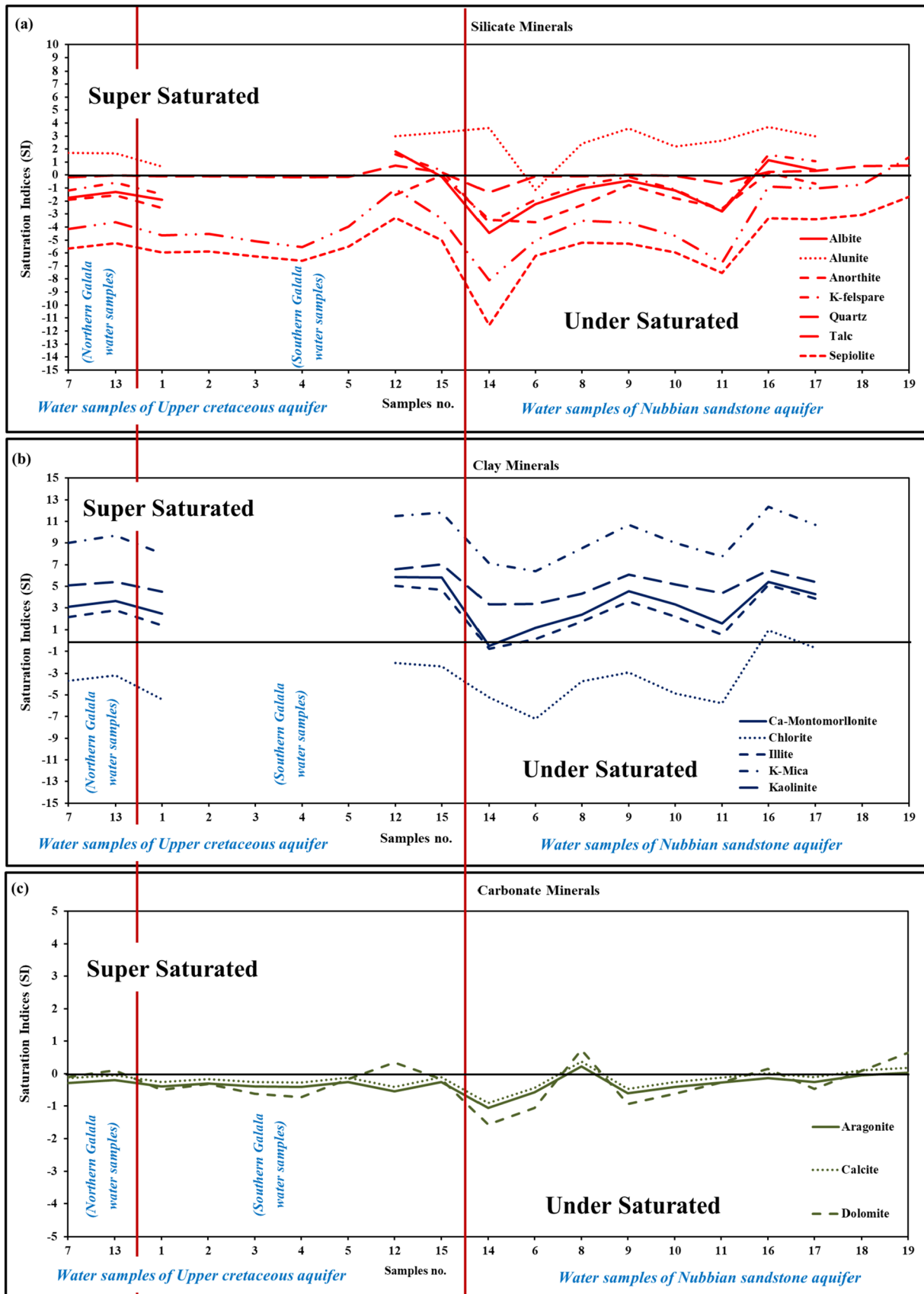


Fig. 20 Groundwater saturation indices (SI) with respect to silicate, clay, and carbonate minerals. The groundwater saturation indices (SI) with respect to iron oxide, evaporates, and Gibbsite minerals

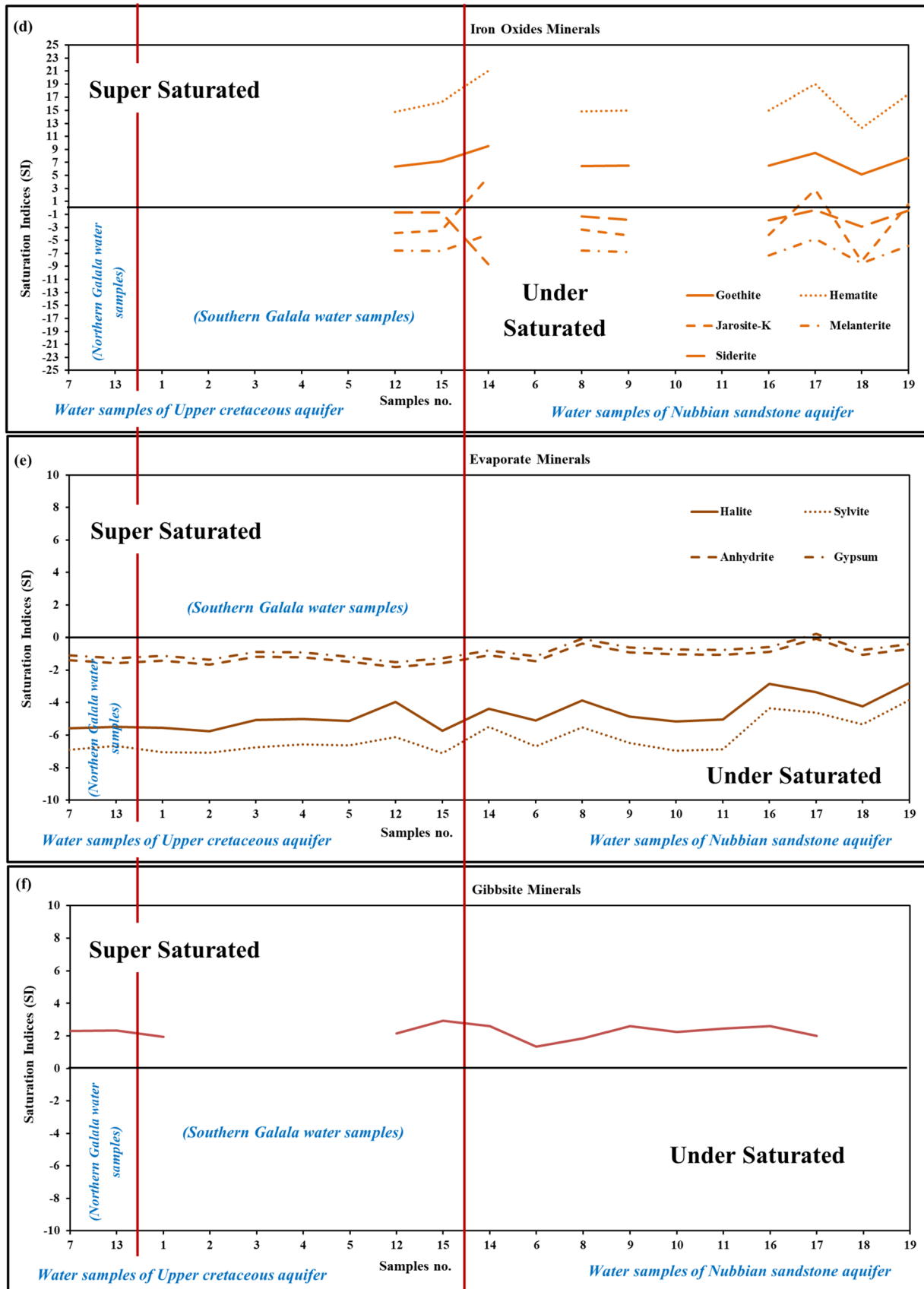


Fig. 20 (continued)

Such conditions reveal that the dissolution of these minerals along the flow system is expected. The water samples, on the other hand, were oversaturated with regard to most of the clay minerals, gibbsite and iron oxides (goethite, and hematite) and with SI close to zero for quartz. This high SI of quartz, clay, and gibbsite minerals could be due to silicate weathering. The high iron concentration could be attributed to the ferruginous mineralization in the water-bearing facies (Aly 2015). It could therefore lead to the precipitation of Fe-rich minerals (Fig. 20d).

Groundwater quality

Evaluating water in general for different uses especially irrigation and drinking is extremely important to make the most benefits of the available water or to treat it to suit a specific use (Salem et al. 2017). As for the groundwater under study, compared to WHO 2011 standards, the concentration of most elements in most samples fall below the permissible limits for drinking water, with the exception of TDS. Only sample 15 which representing the Upper Cretaceous reservoir fall below the TDS permissible limit for drinking water, and the rest of the samples are not suitable for drinking water. Therefore, desalination must be done if it is needed as drinking water.

Long-term production water management requires an understanding of irrigation water quality data. The influence and management of crops and soils are correlated with the quality of irrigation water (Salem et al. 2018, 2021b and 2022b). High-quality crops will also be produced by using high-quality irrigation water to hold other supplies at their best. The qualities of irrigation water are determined by many factors depending on the water source as APHA (2005).

An intuitive way to evaluate the overall irrigation water quality was employed in this investigation using the Irrigation Water Quality (IWQ) index (Simsek and Gunduz 2007). The IWQ index is a linear combination of several factors, including sodium ions, EC, SAR, and boron chloride. It

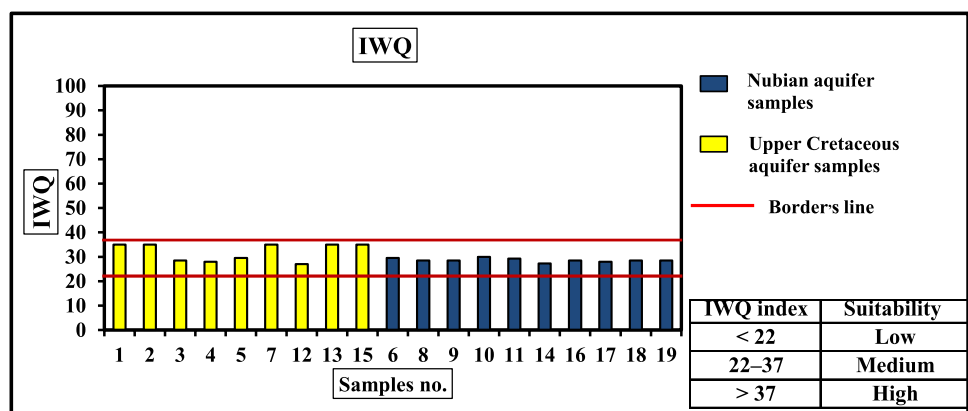
also takes into account the toxicity of the trace element and necessitates the measurement of calcium and magnesium concentration. Since not every location may report all of these metrics, a weighted trace element average is utilized. Lastly, the index takes into account the impact of various influences on sensitive crops by combining nitrate-nitrogen, bicarbonate, and pH linearly (Simsek and Gunduz 2007). The collected water samples were subjected to a suitability analysis based on IQW according to the steps mentioned in Simsek and Gunduz, (2007). All the collected groundwater samples of the studied aquifers are plotted in the medium suitability category (Fig. 21). Despite the quality of the studied groundwater, which is beneficial for irrigation water, caution must be taken against using the water from samples 8, 12, 14, 16, 17, 18 and 19 due to its higher salinity (> 3000 mg/l) due to its proximity to the Gulf of Suez.

Conclusions

A potential contact surface between the sedimentary strata and the basement rocks at Wadi Araba area, western Gulf of Suez, Egypt can be magnetically detected. Therefore, the predicted aquifer is geometrically controlled by the basement topography. The thickness of the aquifer increases toward the east and decreases toward the west. Some magnetic contacts are observed within the sedimentary succession. The petrographic investigations suggest that the rise in magnetic anomalies and, consequently, the subsurface magnetic susceptibility contrasts, may be connected to the ferruginous properties of the aquifer deposits. Minor fractures may descend the basement and/or the sedimentary cover.

The shallow aquifers in the study area consist mainly of siliciclastic-dominated facies of Carboniferous- lower Cretaceous systems (Nubia Group) and upper Cretaceous fractured carbonate-dominated sediments. Lithostratigraphically, the exposed rocks in the studied area are represented by sandstones with thin shale and limestone bed intercalations of Carboniferous Rod El Hamal Formation, which

Fig. 21 Histograms show the IWQ values in the collected water samples



are unconformably overlaid by Permo-Triassic red beds of Qiseib Formation, and cross-bedded sandstones of lower Cretaceous Malha Formation. Two sides of the Wadi resemble fractured limestones of the upper Cretaceous Galala Formation.

The stratigraphical and microfacies investigations suggest that the rise in magnetic anomalies and, consequently, the subsurface magnetic susceptibilities may be connected to the ferruginous properties of the Nubian Sandstone aquifer. A potential contact surface between the sedimentary strata and the basement rocks can be magnetically detected. Therefore, the predicted aquifer is geometrically controlled by the basement topography. The thickness of the aquifer increases toward the east and decreases toward the west. No major magnetic contacts are observed with the sedimentary succession. Minor fractures may descend the basement and/or the sedimentary cover.

Quartz arenite, subarkose, sublitharenite, and greywacke are the main constituents microfacies of the siliciclastic sediments, whereas mudstone/wackestone, packstone/Grainstone resemble the carbonate rocks of the shallow aquifers. The main diagenetic processes that have affected the studied shallow aquifer rocks include mechanical and chemical compaction, the precipitation of calcite, dolomite, and silica (in the form of quartz overgrowths), as well as the dissolution of unstable minerals and shell fragments. These minerals result from the chemical interaction between rocks and groundwater of the studied aquifer. However, some of the diagenetic processes reduced the primary porosity of the aquifers, e.g., compaction, cementation, and recrystallization, while others led to the formation of secondary porosity and changed the groundwater chemistry as dissolution and fracturing.

The groundwater samples under investigation are categorized as hard brackish and very hard saline based on total dissolved solids (TDS) and total hardness (TH). Statistically, the collected samples are classified into five groups (A, B, C, D, and E). All groups exhibit NaCl-type water, except for group B, which has Na-SO₄ type water. Higher TDS values are associated with the Nubian Sandstone aquifer in groups A, B, and C (with average values of 20,766, 9143, and 4431 mg/l, respectively). Conversely, lower TDS values are observed in the D and E groups (with average values of 1209 and 2210 mg/l, respectively), which are primarily sourced from fractured limestone springs. Elevated TDS, Si, Al, Fe, and Mg concentration may be indicative of seawater intrusion and/or water-rock interaction in the Nubian sandstone aquifer, while lower salinity values in fractured limestone samples may result from direct recharge from rainwater. According to the Piper diagram, most samples exhibit NaCl type, signifying primary salinity, with some samples categorized as CaMgSO₄ type.

The dendrograms illustrate two distinct ion associations in Nubian Sandstone groundwater:

SO₄-Ca-Mg-CO₃-K-Si-CO₃ and Cl-Na, suggesting water-rock interaction and seawater intrusion, respectively. In contrast, the Upper Cretaceous aquifer displays two different ion associations: Na-Cl-SO₄ and Ca-Mg-HCO₃-K-Si-CO₃. The Na-SO₄ association indicates ion exchange between Na from rocks and Ca from dissolved gypsum, while the presence of K-HCO₃-Ca-Mg points to a freshwater source. Pearson's correlation analysis suggests that Ca, Mg, Na, K, Cl, and Si are the primary components influencing groundwater salinity.

The relationships among total dissolved solids (TDS), ionic ratios, and other associations imply that seawater intrusion, ion exchange, and silicate weathering are the predominant hydrogeochemical processes affecting the examined groundwater. Samples with low carbonate concentration may be linked to an excess of dissolved calcium resulting from silicate weathering and/or gypsum dissolution. Mineral saturation indices indicate that most collected groundwater samples are undersaturated with carbonate minerals (aragonite, calcite, dolomite, and siderite) and silicate minerals (albite, feldspar, and anorthite), as well as chlorite. Additionally, all samples show insufficient saturation with evaporite minerals (anhydrite, gypsum, halite, and sylvite) and melanterite.

Conversely, water samples are supersaturated with respect to most clay minerals, gypsite, and iron oxides (goethite and hematite), with a saturation index close to zero for quartz. The elevated saturation indices for quartz, clay, and gibbsite minerals may be attributed to silicate weathering. The higher iron concentration is likely linked to ferrous mineralization in the water-bearing facies, potentially leading to the precipitation of iron-rich minerals. Studied groundwater is unsuitable for drinking except one sample, while it is moderately suitable for irrigation with caution for higher salinity in samples located near to the Gulf of Suez.

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Data availability The data were collected and analyzed primarily by us.

Declarations

Conflict of interest The authors have no conflicts of interest to declare that are relevant to the content of this article.

Consent to participate The authors consent to participate in this research study.

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