

Article

Sources of Salinization of Groundwater in the Lower Yarmouk Gorge, East of the River Jordan

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Abstract: In the Lower Yarmouk Gorge the chemical composition of regional, fresh to brackish, mostly thermal groundwater reveals a zonation in respect to salinization and geochemical evolution, which is seemingly controlled by the Lower Yarmouk fault (LYF) but does not strictly follow the morphological Yarmouk Gorge. South of LYF, the artesian Mukeihbeh well field region produces in its central segment groundwaters, an almost pure basaltic-rock type with a low contribution (<0.3 vol-%) of Tertiary brine, hosted in deep Cretaceous and Jurassic formations. Further distal, the contribution of limestone water increases, originating from the Ajloun Mountains in the South. North of the LYF, the Mezar wells, the springs of Hammat Gader and Ain Himma produce dominantly limestone water, which contains 0.14–3 vol-% of the Tertiary brine, and hence possesses variable salinity. The total dissolved equivalents, TDE, of solutes gained by water/rock interaction (WRI) and mixing with brine, $TDE_{WRI+brine}$, amount to 10%–70% of total salinity in the region comprising the Mukheibeh field, Ain Himma and Mezar 3 well; 55%–70% in the springs of Hammat Gader; and 80%–90% in wells Mezar 1 and 2. The type of salinization indicates that the Lower Yarmouk fault seemingly acts as the divide between the Ajloun and the Golan Heights-dominated groundwaters.

Keywords: Hauran Plateau; Golan Heights; Ajloun Dome; Yarmouk basin; salinization; mixing of water types; statistical modelling

1. Introduction

This study aims at the identification and quantification of sources of salinization in different geological formations in the well field region of the Lower Yarmouk Gorge (LYG) shared by Israel, Jordan and Syria (Figure 1). The productive water resource in the LYG is vitally important for Jordan and Israel because of water shortage in both countries [1]. Like elsewhere in all semi-arid regions in the world, the ground and surface-water resources are over-exploited, leading to water shortages due to increasing demand and to consequences of climate change [2]. In such regions, water supply is a serious socio-economic and political issue, if transboundary flow is involved. For sustainable management and protection of groundwater resources, water authorities need detailed knowledge

of (i) the respective recharge areas of distinct wells or well fields, (ii) the flow characteristics and (iii) possible inter-aquifer flow [3]. In areas with transboundary water resources, rules of equitability and no-harm also have to be obeyed in order to prevent political interference [4].

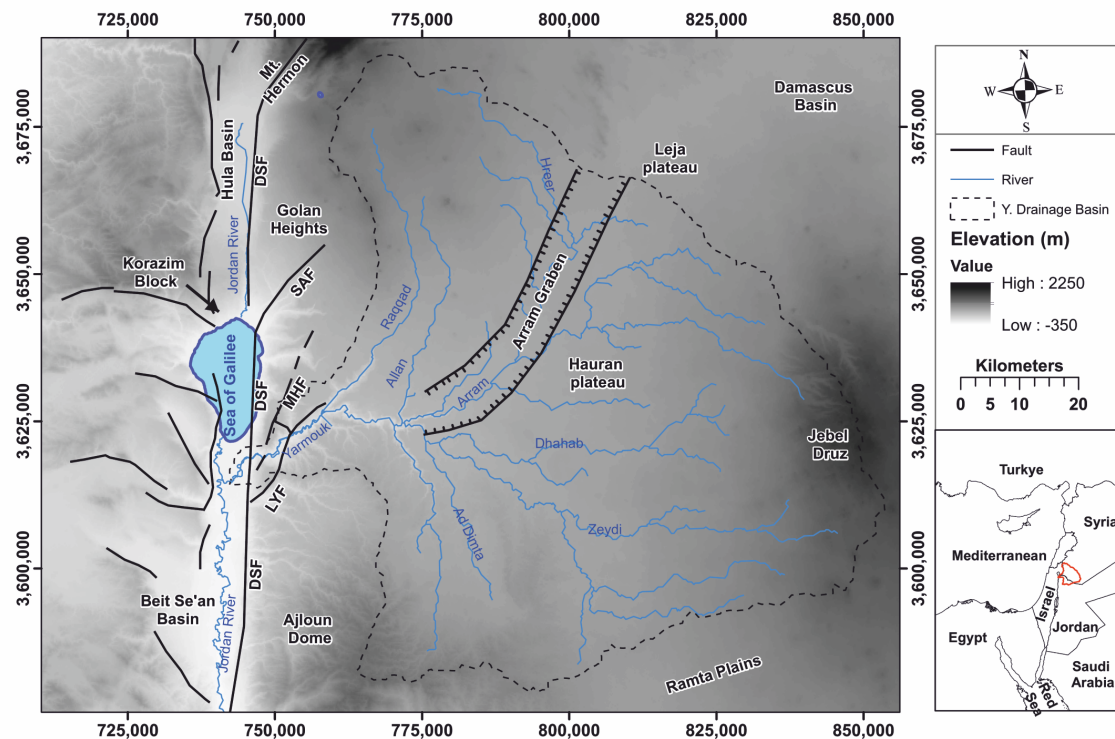


Figure 1. Overview of the Yarmouk drainage basin and its environment. The basin is outlined in red in the lower insert. Note the graben structure with the perennial Arram River in the Yarmouk basin. DSF = Dead Sea Transform fault; SAF = Sheik Ali fault; MHF = Mevo Hama fault; LYF = Lower Yarmouk fault.

The study of the hydrochemical compositions of groundwater and brines reveals the origin of water and its salinization (e.g., [2,5–15]). The inorganic composition of groundwater depends on the weathering of minerals in the catchment rocks, the water/rock interaction (WRI) along the flow paths and the amounts of atmospheric deposition [16]. Processes together yield typical ionic ratios in groundwater ([17] and references therein).

In contrast, trace elements such as the rare earth elements (REE) and uranium behave differently and thus are useful to highlight other aspects [15]. Their extremely low concentrations ($< \mu\text{mol/L}$) are controlled by adsorption onto mineral surfaces and co-precipitation with alteration minerals [18–26]. The suite of REE and yttrium, Y, henceforth combined to REY, is widely used to identify the origin of groundwaters [8–10,12,13,15,27]. REY immediately achieve steady state conditions in the infiltrating water due to their high charge and affinity to build surface complexes. For instance, more than 99% of REY released from dissolving calcite in limestones are immediately adsorbed onto calcite surfaces [17]. During migration of groundwater, REY are continuously subjected to exchange with surface-adsorbed REY. Thus, after some time the REY patterns of groundwater resemble those gained during weathering of the catchment rocks [28]. Therefore, REY patterns characterize the lithology of the recharge area, whereas major and minor elements reflect the solubility of minerals of the entire assemblage of catchment and aquifer rocks.

A previous conjoint application of major and trace elements in the ground water yielded insights into its salinization and flow paths in the study area, which will be briefly summarized in Section 2 [2, 15,27,29]. However, the issue of potential transboundary flow of water between the Ajloun in the

south [30,31] and the Golan Heights in the north [2,32–34] underneath the Yarmouk River was and still is debated. It has been considered whether or not the Gorge delineates a fault or represents an anisotropy zone which possibly prevents transboundary flow of groundwater between Jordan and Israel [35,36]. Seismic lines crossing this border have not been shot up to now but several ones in the southern Golan Heights are available [37]. Shallow faults in northwest Jordan are described in [38]. Based on these surveys and additional evidence [37], a strike-slip-flower-structured fault system and numerous buried faults crossing the Gorge at an acute angle are plausible.

The purpose of this contribution is to analyse and interpret the available hydrogeochemical data concerning major ions, the REY and the hydrogeological setting (Section 2) in order to discern the possible mixing paths and sources of salinization for the groundwaters of the Lower Yarmouk Gorge, discriminating between water/rock interactions and mixing with relic brines of different natures (Section 3). In particular, two different conceptual models are employed to estimate the fractions of limestone and basaltic-rock waters and brines which can be observed in the data (Section 3.3). Multivariate statistical analysis in the form of clustering (Section 3.4) is also employed to support and ultimately validate the interpretation and to trace the discrepancies between geographically contiguous water samples in view of the identified processes (Section 4.4).

2. Hydrogeological Setting

The Yarmouk basin comprises the eastern Golan Heights and south-eastern flanks of the Hermon Massif, shared by Israel and Syria; the northern plunges of the Ajloun Dome (Jordan); and the Hauran Plateau, including the western flank of the Jebel Druz (Syria) (Figure 1). The LYG is the major outlet of surface and groundwater from the Yarmouk basin. Morphologically, the Gorge separates the Ajloun Mountains and the Jordanian Ramta Plains from the Golan Heights and the Hauran Plateau, respectively.

The anticlinal structure of the Ajloun is built of Lower Cretaceous Kurnub sandstones and Upper Cretaceous, marine, strongly karstified, fractured and silicified lime and dolostones, forming the A7/B2 aquifer in Jordan (Figure 2), which descends northward. For easement and shortness, the Jordanian nomenclature of formations is preferred in this contribution. Groundwaters in the A7/B2 are confined by the overlying bituminous Senonian B3 aquiclude, which contains phosphorite, chert and chalk, and separates the A7/B2 from the locally exploited limy B4 aquifer.

The Cretaceous and Tertiary sedimentary aquifers, which crop out in the Ajloun anticline, descend northward into the Golan syncline and surface later, again at the foothills of the Hermon anticline, which consist of thick Jurassic lime and dolostone aquifers with abundant basaltic intrusions [39]. As a consequence, groundwaters south and north of the Gorge migrate through the same aquiferous formations. Contrasting the Ajloun, the Golan Heights are unconformably covered by up to 700 m thick Plio-Pleistocene cover basalt [40,41]. A marly sequence at its base, the highly fractured basalt, serve as a regional aquifer, which itself is annually directly recharged by 500–1200 mm of precipitation [7]. During the pre-Quaternary phase, the Golan Heights were subjected to tectonic stress documented in a highly faulted and deformed subsurface [42] with intense and deep karstification of the Upper Cretaceous and Tertiary lime and dolostones [43]. A meridional ridge in these formations acts as a subsurface water divide in the covering basalt aquifer [7], leading to groundwater drainage in the latter either W–SW to the Hula and the Sea of Galilee basin, or E–SE into the Hauran Plateau and the Upper Yarmouk Gorge. Hydraulic connections between the basaltic cover and the underlying aquiferous Cretaceous carbonate formations may exist throughout the Golan Heights [44].

The eastward continuation of the Golan Heights is the flat and southward dipping volcanic area of the Hauran Plateau, which passes in the SE into the enormous accumulation of Neogene-Quaternary basalts of the Jebel Druze [45,46]. Precipitation infiltrates directly into Upper Quaternary basalts exposed all over the Plateau and drains towards the LYG, partially feeding perennial springs in the arcuated Wadi Arram (Figure 1) which was a major contributor to the Yarmouk River in the past [47].

Age		Group		Formation		Hydrogeological properties		
Period	Epoch/Stage	Golan	Ajloun	Golan	Ajloun	Golan	Ajloun	
Quaternary	Holo-/Pleistocene			Yarmouk Basalt			basalt	
Neogene	Pliocene	Kefar Giladi		Cover Basalt			basalt	
				Bira/Gesher			silicified limest., dolomite	
				Hordos	Waqas Conglomerate (WC) - Jordan Valley		marl, sand	
	Lower Basalt					alkali olivine basalt		
	Oligocene			Susita/Fiq			marlst., sandy dolomite	
Paleogene	Eocene		Belqa		B5 Wadi Shallalah Chalk (WSC)		chalk, bituminous	
		Avedat		Maresha/Adulam	B4 Umm Rijam (URC)	marl, chalk, limestone	chalky limest., chert beds	
Upper Cretaceous	Upper Maastricht-Paleocene	Mt. Scopus		Taqiye	B3 Muwaqqar Chalk (MCM)	bituminous, chert, phosphorite	micritic limest., bituminous (oilshale)	
	Masstrichtian			Ghareb	B2b Al Hasa Phosphorite (AHP)		calcareous, phosphorite beds, limest., chalk, marl	
	Campanian			Mishash	B2a Amman silicified limestone (ASL)	marly limestone,	limest., chalky dolomite, phosphate, chert	
	Santonian			Menuha	B1 Wadi Umm Ghudran (WUG)		massive chalk, limest., phosphatic sandst., chert	
	Turonian	Judea		Ajloun	Bina	A7 Wadi Es Sir Limest. (WSL)	limestone, dolomite, marly limest.	dolomitic limest., sandst., cherts
	Cenomanian-Turonian					F/H/S-undifferentiated		marl and gypsum
								A5-6 Shueib (S)
Period	Epoch/Stage	Hermon	Ajloun	Hermon	Ajloun	Hermon	Ajloun	
Upper Jurassic	Tithonian	Arad	Zarqa	J6-J7 (Nahal Saar)	Azab	limestone	sandstone, siltstone, limestones	
	Kimmeridgian			J5 (Kidod)		marl, shale		
Middle Jurassic	Oxfordian			J4 (Hermon/Zohar)		limestone		
Lower Jurassic	Hettangian-Toarcian			J1-J3		dolomite, limestone		

Figure 2. Stratigraphic table comparing the Israeli and Jordanian nomenclature. Aquiferous units are colored in blue after [15].

Along the northern flank of the LYG, hot groundwater emerges at Hammat Gader springs (codes EM, ER and EB in Table 1; 38–43 °C) from the B3 aquitard [48] and ascends at 41–60 °C in the artesian wells of Mezar from A7 (Mezar 2) and B2 (Mezar 3) aquifers (Figure 3).

South of the Yarmouk River the hot water from Ain (Arabic term for spring) Himma (42 °C) emerges from the B3 aquitard, ascending along faults from the B2 aquifer [6]. In the nearby artesian Mukheibeh well field, the groundwater is exploited from A7/B2 discharging with temperatures of 29–46 °C probably heated by volcanic intrusions at depths of 3–4 km [49]. Hot groundwater from the A7 aquifer is also known to come from the western Ajloun escarpment within the Lower Jordan Valley [30,50,51]. In the Ajloun the temperature of groundwater from A7/B2 is only slightly enhanced (23–31 °C) and cool if draining the shallow basaltic aquifer or the B4.

With few exceptions, the groundwaters of the LYG are saturated with respect to calcite but not with respect to gypsum or halite [2]. The groundwaters from Ain Himma, the springs of Hammat Gader (enclosing codes ES, EB, ER and EM in Table 1) and Mezar well 3 are seemingly mixtures of local groundwater, relic seawater evaporation brine(s) and leached evaporites and dissolved calcite from limestone [2,15].

Table 1. Analyses of spring and well waters from the Lower Yarmouk Gorge. Grouping according the geographical and chemical proximity (code). In the two last columns the normalized Tb/Lu ratios and the fraction of basaltic-rock water, ε_{bw} , in mixture with limestone water are given as derived from Figure 4.

Code	Source	Sampling	East	North	pH	Eh *	Temp	EC	Ca ²⁺	Mg ²⁺	K ⁺	Na ⁺	Cl [−]	SO ₄ ^{2−}	HCO ₃ [−]	TDE	Tb/Lu	ε_{bw}
Group		Year	(UTM, WGS84, Z36N)			mV	°C	μS/cm					meq/L §				normalized	
U1	Mukheibeh 2	2016	753241	3622342	6.13	17.90	28.90	830	3.90	2.14	0.07	1.70	1.59	1.17	4.50	15.07	0.23	0.95
U1	Mukheibeh 4	2016	753212	3622331	7.04	272.71	29.10	827	4.20	2.30	0.07	1.74	1.59	1.19	4.69	15.79	1.07	0
U1	Mukheibeh 4	2013	753209	3622333	7.10	62.99	28.80	807	4.80	2.39	0.08	1.67	1.56	1.14	5.11	16.75	0.27	0.83
U1	Mukheibeh 2	2013	753243	3622340	7.12	64.61	28.14	809	4.80	2.47	0.08	1.68	1.65	1.19	5.11	16.98	0.21	0.99
U1	Mukheibeh 1	2001	753119	3622154	7.00	66.71	29.10	797	4.57	2.47	0.08	1.70	1.61	1.13	5.86	17.42	0.29	0.80
U2	Mukheibeh 6	2016	753018	3622417	7.06	−40.08	31.00	667	4.30	2.39	0.09	2.13	2.07	1.25	5.01	17.23	0.19	1.00
U2	Mukheibeh 7	2016	754257	3623142	7.17	−17.13	38.50	774	4.20	2.39	0.07	1.74	1.71	0.74	4.73	15.57	0.46	0.47
U3	Mukheibeh 10	2016	753267	3622856	6.96	−116.60	39.00	710	3.60	2.06	0.09	2.04	1.69	0.82	4.27	14.58	0.7	0.20
U3	Mukheibeh 5	2016	747753	3618570	7.32	161.62	40.90	876	3.55	1.98	0.10	2.04	1.61	0.76	5.00	15.04	0.78	0.12
U3	Mukheibeh 11	2016	753845	3622588	6.92	200.18	31.90	821	3.80	3.21	0.07	1.91	1.59	1.11	4.73	16.43	0.53	0.35
U3	Mukheibeh 9	2016	756312	3624296	7.20	−67.10	28.90	1157	2.70	2.39	0.41	5.22	2.33	0.44	7.52	21.00	0.35	0.66
U3	Mukheibeh 8	2013	755490	3624127	7.45	−123.14	44.90	701	3.40	1.65	0.13	2.32	1.82	0.58	4.64	14.53	1.1	0
U3	Mukheibeh 8	2016	755495	3624134	7.16	−62.14	44.90	723	2.95	1.65	0.13	2.43	1.88	0.26	4.03	13.33	0.73	0.17
U3	Mukheibeh 13	2016	754268	3623212	7.12	−46.50	38.90	778	3.60	2.06	0.09	2.00	1.74	0.71	4.33	14.53	0.69	0.24
U3	Mukheibeh 13	2013	754280	3623202	7.38	−76.13	38.50	752	4.05	2.06	0.10	1.90	1.67	0.51	5.04	15.32	0.67	0.35
ES	Ein Sahina	2016	750190	3619926	7.04	453.74	28.00	844	4.35	2.30	0.08	1.91	1.88	1.02	4.93	16.48	0.75	0.15
AH	Ein Himma	2007	751665	3621722	7.02	−27.54	40.00	1433	5.67	2.80	0.35	5.31	6.88	4.43	6.88	32.32	0.98	0.33
AH	Ein Himma	2001	751665	3621722	7.06	−144.95	41.50	1418	5.38	2.87	0.36	5.40	5.86	3.09	5.52	28.47	0.94	0.01
AH	Ein Himma	2013	751600	3621710	7.10	71.63	37.70	1130	5.90	2.63	0.23	3.67	4.23	2.60	4.77	24.04	0.5	0.40
AH	Ein Himma	2016	751661	3621700	7.03	85.46	40.00	499	5.50	2.47	0.24	4.09	4.54	2.69	4.00	23.53	0.32	0.70
AS	Ain Saraya	2016	750429	3619424	6.83	−15.94	38.30	1655	6.50	3.13	0.31	6.96	9.65	2.50	4.47	33.51	0.29	0.79
ER	Ein Balsam	2016	749705	3619324	6.76	10.68	41.90	1600	6.00	2.80	0.31	5.65	9.17	2.77	4.73	31.43	0.56	0.37
ER	Ein Reach	2016	750014	3619198	6.69	−75.73	43.40	1860	6.50	3.05	0.36	6.96	10.61	2.67	4.78	34.91	0.66	0.24
ER	Ein Reach	2000	750348	3619399	6.85	−95.66	38.00	1728	7.12	3.25	0.31	6.23	8.71	2.39	5.93	33.95	0.68	0.21
ER	Ein Reach	2004	749984	3618816	6.81	−146.95	41.50	1759	6.72	3.25	0.32	6.79	9.75	2.48	5.79	35.11	0.55	0.36
EM	Ein Makla	2016	749859	3619141	6.59	−116.47	49.50	2160	7.50	3.21	0.44	9.13	13.40	3.12	3.58	40.38	0.46	0.47
EM	Ein Makla	2000	749909	3619091	6.64	−131.40	47.30	2190	8.91	3.64	0.46	9.23	13.77	3.30	5.57	44.88	0.57	0.33
EM	Ein Makla	2004	749811	3618793	6.73	−166.56	49.60	2160	7.58	3.46	0.44	9.12	13.13	3.14	5.35	42.23	0.57	0.33
ME1	Meizar 1	2016	752652	3625884	7.38	−79.02	35.20	1630	2.60	1.40	0.33	10.00	10.89	0.26	3.56	29.05	0.748	0.13
ME2	Meizar 2	2016	752706	3622894	6.40	−76.90	60.60	2080	8.50	3.05	0.62	9.13	11.06	7.04	3.62	43.01	1.44	0
ME2	Meizar 2	2001	752700	3622914	6.63	−102.33	60.00	1650	7.12	2.86	0.56	7.74	8.93	5.80	4.00	37.01	1.4	0
ME3	Meizar 3	2016	752725	3622926	6.81	−119.60	42.20	810	3.55	2.06	0.10	2.30	2.22	0.40	5.41	16.05	0.48	0.43
ME3	Meizar 3	2001	752707	3622922	7.09	−129.23	41.80	664	3.27	1.72	0.12	2.11	1.74	0.08	5.17	14.20	1.39	0
YR	Yarmouk	2016	756255	3624301	8.40	190.97	23.50	961	2.75	2.47	0.15	4.22	3.10	3.06	3.02	18.77	0.47	0.46

* Eh-values: expressed relative to standard hydrogen electrode; § relative standard deviation for analytes: ≤1% for Ca, Mg, K, Na; ≤2% for Cl, SO₄ and ≤5% for HCO₃.

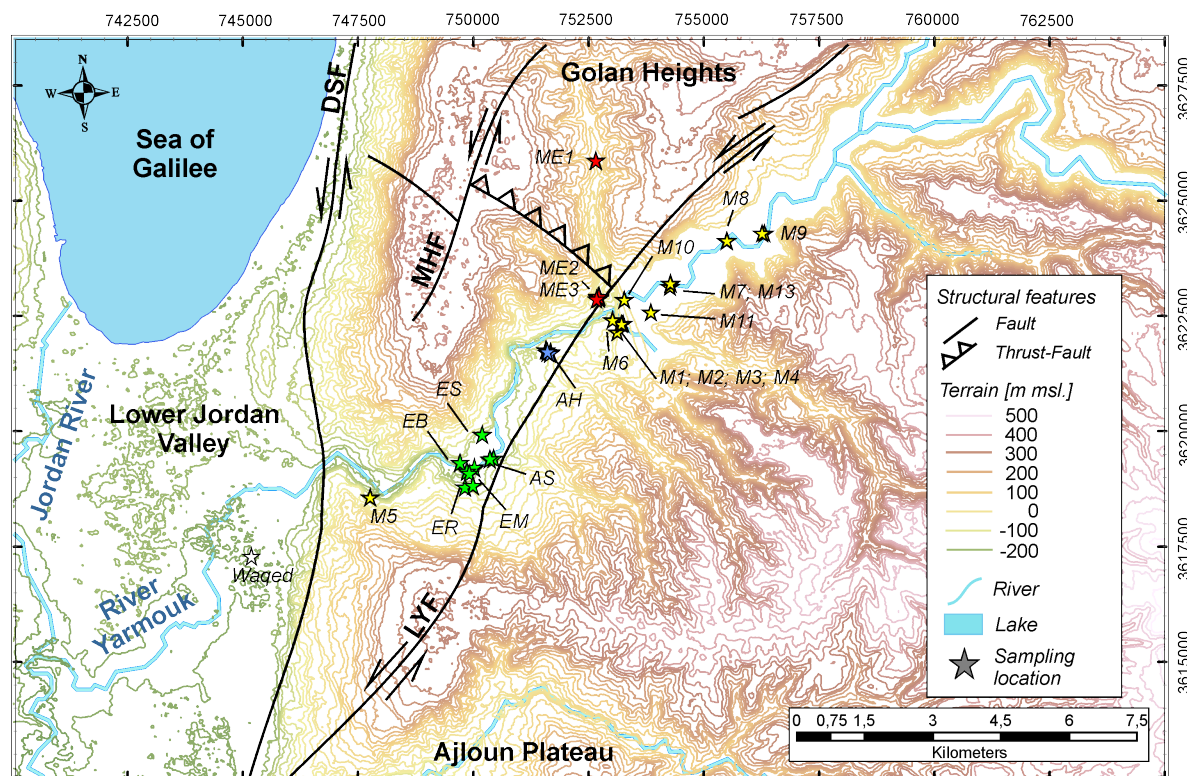


Figure 3. A detailed view of the sampling locations in the Lower Yarmouk Gorge. Very important are the Lower Yarmouk (LYF) and Mevo Hamma faults (MHF) and the rise of the Hammat Gader block in between. DST= Dead Sea fault. The region of Hammat Gader encloses codes, EB, ER and EM (Table 1). Color codes of stars: yellow—Mukheibeh well field; red—Mezar well field; green—Hammat Gader; blue—Ain Himma.

3. Materials and Methods

3.1. Sampling and Analytical Procedure

During a synchronous sampling campaign in 2016, wells and springs on both sides of the LYG were chemically analyzed. To allow for a regionally comprehensive elaboration, selected analytical results from earlier local campaigns were included (Table 1). During all campaigns, onsite parameters (pH, temperature, redox potential and electric conductivity) have been measured in the discharging water jets of springs and pumping wells using WRW 350i and respective probes (SenTix41, Sentix ORP, Tetracon325). When parameters were stable, samples were derived from jets applying a peristaltic pump and pre-cleaned silicon tubes connected to 0.22 μm filter before storage in pre-cleaned HD-PE bottles. Contrasting the anion samples, cation samples were later acidified using one drop of 6 N HCl to ensure conservation. The alkalinity (expressed as HCO_3^-) was determined by titrating 10 mL of sample with 1.6 N H_2SO_4 to pH 4.3. To determine REY, pre-concentration was required. Therefore, about 4 L of sample were filtered (0.22 μm), acidified by sub-boiled (index sbb) 6 N HCl to pH of about 2–2.5 and spiked with 1 mL of Tm solution, by which the recovery of REY in subsequent pre-concentration procedure could be checked. Usually the recovery was between 95% and 103%. At the same day, the samples were passed through C_{18} Sep-Pak cartridges, loaded with ethylhexylphosphate liquid ion exchanger. In the lab, each cartridge was eluted with 6 N HCl_{sbb} ; eluates were evaporated to incipient dryness and taken up with $\text{HNO}_{3\text{sbb}}$; and the resulting solution was analyzed applying ICP-MS (Elan DRC-e). Independently, Ca^{2+} and Mg^{2+} were determined by ICP-MS. K^+ and Na^+ were analyzed by ICP-AES (Spectro Arcos) using matrix adjusted standard solution for calibration. Cl^- , Br^- , SO_4^{2-} were determined by Dionex ICS (AS18 column).

3.2. Selection of End Member Fluids in the Yarmouk Basin

The suggested quantification of salinity contributions is based on defined end members of water types in the LYG:

- Infiltration of precipitation over basaltic catchments, particularly in the Hauran plateau;
- Infiltration of precipitation over a limestone catchment mainly in the Ajloun Mountains and Mount Hermon/Golan Heights;
- Relics of brines residing in Jurassic-Cretaceous limestones;
- Dissolution of evaporites and water/rock interaction (WRI) along flow paths.

Basaltic rock and limestone waters with lowest Cl^- concentrations are suggested as end members because enhanced Cl^- concentrations suggest dissolution of halite from evaporites and/or leaching of seawater brines enclosed in limestones. The lowest Cl^- concentrations of basaltic-rock water from the Golan and Hauran Plateau were 0.84 meq/L (Tables 1 and A1).

The averages of limestone water of two well waters from each the Golan Heights and the Ajloun Mountains were selected. Their Cl^- concentrations ranged between 0.89 and 0.80 meq/L (Table 2), suggesting that the samples with values >0.80 meq/L might have leached either evaporites or seawater brines.

Table 2. Averages of water types in the Lower Yarmouk Gorge. For more details refer to Table A1.

	Ca^{2+}	Mg^{2+}	K^+	Na^+	Cl^-	SO_4^{2-}	HCO_3^-
	meq/L						
Basaltic-rock water	1.49	1.31	0.10	1.19	0.84	0.25	2.69
Ajloun limestone water	4.93	1.02	0.04	0.71	0.80	0.22	4.66
Golan limestone water	0.39	0.31	0.20	2.49	0.89	0.24	6.41
Ha'On brine	45.1	123	8.79	255	421	4.31	8.42
Rosh Pinna brine, 2486–2586 m	240	59.6	9.18	348	643	24.2	6.06

Two particular brines have to be considered:

1. The Late Tertiary brine was generated by evaporation of intruded Tethys seawater into the Jordan–Dead Sea Rift [52]. This evaporation brine infiltrated the Cretaceous and Jurassic aquifers east and west of the Rift. This type of Mg^{2+} - Cl^- brine was identified at Ha'On in the 1960s [53] along the SE shore of Lake Tiberias. The variations in composition of two wells at Ha'On between 1961 and 2004 are averaged (Table 2). For more detail refer to Table A1.
2. The Late Triassic to Early Jurassic brine of Rosh Pinna is hosted at depths of 2500 m in limestones of the Korazim block north of Lake Tiberias (Figure 1).

3.3. Estimation of Fractions of Brine, Basaltic Rock and Limestone Water

The fractions of basaltic-rock water, ε_{bw} , in mixtures of both pure basaltic-rock water and limestone water (Figure 4a), are derived from interpretation of REY distribution patterns showing the variation of mixtures of both types of groundwater. Each of the mixtures of REY patterns is characterized by C1 chondrite-normalized Tb/Lu values decreasing with increasing ε_{bw} values (Figure 4). Determining the Tb/Lu ratio of each individual groundwater and brine a reliable ε_{bw} for each groundwater in the study area is derived from Figure 4b. The REY distribution patterns of all groundwaters and brines are discussed in [15].

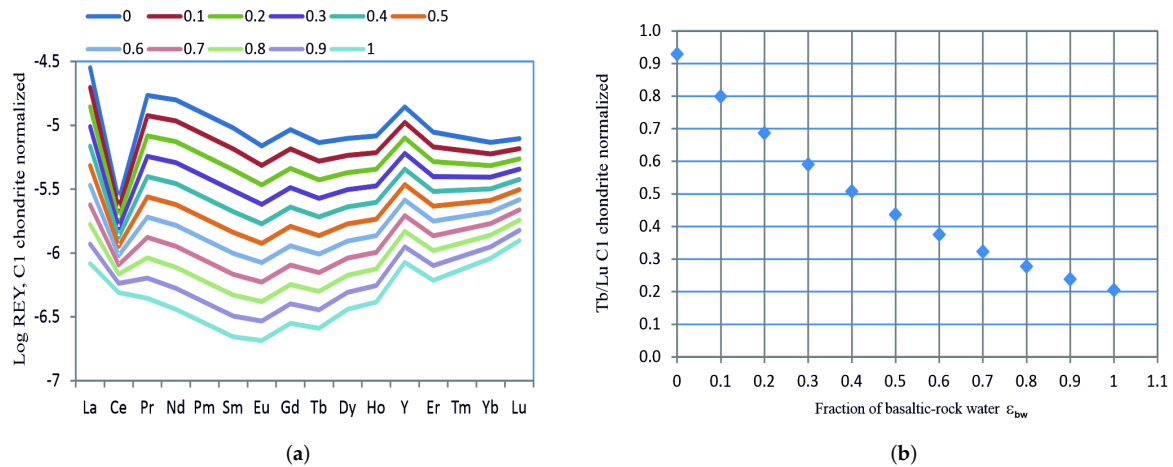


Figure 4. Variation of rare earth element distribution patterns in mixtures of basaltic-rock and limestone water. (a) Estimates of mixing of basaltic-rock-(1) and limestone water (0); (b) resulting C1 chondrite normalized Tb/Lu values as a function of ϵ_{bw} . REY data are taken from [15] and are normalized to C1 chondrite to get smooth curves. The average of Ein Dan and Ein Banyas in Mount Hermon Massif are used as pure limestone water; as basaltic-rock water the analysis of Mukheibeh 2/2013 is used. Data are taken from [2].

This approach of ϵ_{bw} assumes that the REY patterns are not significantly varied by dissolution of evaporites or by WRI, applying ϵ_{bw} and $\epsilon_{lmst} = 1 - \epsilon_{bw}$, and the analyzed concentrations of species i . Two different estimates are presented by assuming either presence or absence of halite in evaporites. The concentration of $c_{i,agw}$ is approached by summation of the end member composition of basaltic-rock and limestone water (Table 2), and contributions of leached brine and WRI (Equation (1)). The summation of contribution of Cl^- from basaltic-rock and limestone water is given as "estimated" $c_{Cl,est}$ by Equation (2). Equation (1) yields the sum of $c_{i,est} + \epsilon_{brine} \cdot c_{i,brine} + c_{i,WRI}$ for each species i . If halite is absent, the maximum fraction of brine is derived from Equation (3) which is probably an overestimation. Assuming that Cl^- is only a component of the brine and not of WRI, ϵ_{brine} is given by Equation (4).

$$c_{i,agw} = \epsilon_{bw} \cdot c_{i,bw} + (1 - \epsilon_{lmst}) \cdot c_{i,bw} + \epsilon_{brine} \cdot c_{i,brine} + c_{i,WRI} \quad (1)$$

$$c_{i,est} = \epsilon_{bw} \cdot c_{i,bw} + (1 - \epsilon_{lmst}) \cdot c_{i,bw} \quad (2)$$

$$c_{i,agw} = c_{i,est} + \epsilon_{brine} \cdot c_{i,brine} \quad (3)$$

$$\epsilon_{brine} = (c_{Cl,agw} - c_{Cl,est}) / c_{brine} \quad (4)$$

Another way to characterize the salinization of groundwater is achieved by estimating the total dissolved equivalents, TDE (in meq/L), which is independent on processes such as ion exchange with clay minerals; albitization and associated calcite precipitation; and dolomitization of limestones. TDE, however, varies due to dissolution and precipitation of minerals and mixing of fresh and saline waters. TDE_{bw} and TDE_{lmst} are estimated for the contributions of corresponding waters (Equation (5)). TDE of the analyzed groundwater, TDE_{agw} , is given by summation over all dissolved species i (Equation (6)). Assuming that Cl^- is only contributed by WRI, the sum of $TDE_{WRI} + TDE_{brine}$ is estimated according to Equation (7) with $TDE_{brine} = 0$.

$$TDE_{est} = TDE_{bw} + TDE_{lmst} \quad (5)$$

$$TDE_{agw} = TDE_{est} + TDE_{brine} + TDE_{WRI} \quad (6)$$

$$TDE_{WRI+brine} = TDE_{WRI} + TDE_{brine} = \sum_i [c_{i,agw} - (\epsilon_{bw} \cdot c_{i,bw} + (1 - \epsilon_{bw}) \cdot c_{i,lmst})] \quad (7)$$

3.4. Multivariate Statistical Analysis: K-Medoids Clustering

Multivariate statistical analysis provides a powerful set of tools to leverage the growing amount of available data and has been successfully used to discriminate past and active geochemical processes, especially in complex hydrogeochemical settings [54–56]. To fully explore the possible outcomes of such analyses, however, is outside the scope of this work, mostly because of the relatively low number of samples ($n = 35$) and their inhomogeneous covering of the region of study. Furthermore, the possible end members of the groundwaters and brines in the region of interest have already been identified in Section 3.2. A clustering algorithm is applied to the data of Table 1, limited to the elemental concentrations. This means grouping the samples into a specified number of categories based on the dissimilarities between their chemical components, in a data-driven approach, and thus, irrespective of the locations of the samples. The robust method called partitioning around medoids or *k-medoids* [57] in the implementation of [58] was hereby chosen instead of the classical *k-means*. The *k-medoids* algorithm chooses one actual data point as the center of each cluster instead of the center of mass, as in the *k-means*. The L1 norm or Manhattan distance (sum of the absolute distances of all components) was retained as a metric for the computation of distances for the concentration vectors. The L2 norm or Euclidean distance produced the same clustering but less readable results. The number of nine clusters was retained as a trade-off between the statistical significance of the partitioning, evaluated through the silhouette method, and the interpretability of the results. The added value of this analysis is presented in Section 4.4.

4. Results

4.1. Correlations of Solutes in Yarmouk Groundwater

The cross plots of dissolved species in groundwater reveal relationships between end members of saline and fresh water. The fresh water end members of basaltic rock and limestone water (Table 2) are implemented in Figure 5. The correlation of various elements with Cl^- reveals several things: Mukheibeh concentrations either tightly cluster (Figure 5a–d) or spread in one direction (Figures 5e–f). Waters from Hammat Gader and Mezar/Himma seem to represent dilution lines with different saline end members and the assumed basaltic rock and limestone water both plotting near the Mukheibeh cluster. The same may be true for Figure 5d, where the Br^- concentrations of the low-salinity end members are close to zero. Values of Ca^{2+} and Mg^{2+} in fresh water show a wide spread, with diverging (Figure 5e) or monotonous (Figure 5f) increase with increasing Cl^- . The Mukheibeh waters show enhanced Ca^{2+} and Mg^{2+} concentrations compared with the basaltic rock water and limestone water.

More details are revealed by the trend lines in the cross plots of $1000 \cdot \text{Br}^- / \text{Cl}^-$ and $\text{Na}^+ / \text{Cl}^-$ in Figure 6. For orientation, the trend of evaporated seawater is given as a red line [59,60]. The groundwater from springs of Hammat Gader and Himma and from well Mezar 2 define vertical trends, which are only explainable by leaching of Br^- from the organic-rich limestones of the B3 aquitard. Mezar 1 and 3 and the low $\text{Br}^- / \text{Cl}^-$ samples of all vertical trends suggest a mixing line between Mukheibeh groundwater and evaporated seawater such as the Ha'On brine [61]. A second mixing line is indicated by Ein (Hebrew term for spring) Sahina (ES) and the wells Mukheibeh 1 and 6. The water of Mukheibeh 9 well shows an extreme position.

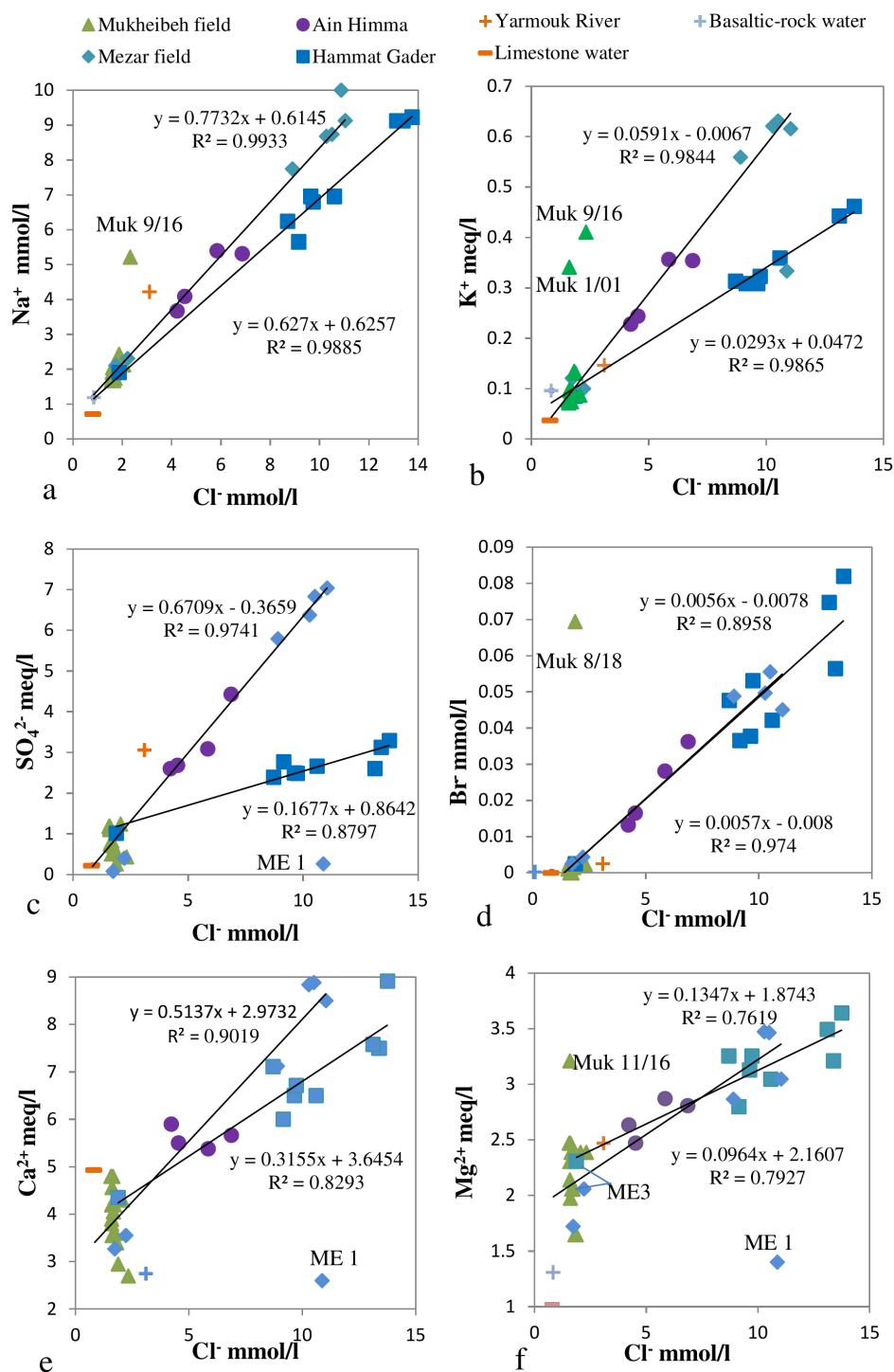


Figure 5. Cross plots of dissolved species in groundwater. Note that the high-salinity groundwaters are related to either basaltic or limestone water (a–e). This is not the case in (f). Averages of low- Cl^- -containing water from the Ajloun Mountains and from the Golan Heights are used for limestone water. Averages of low- Cl^- -containing water from the Hauran Plateau are used for basaltic-rock water (code names are explained in Table 1). x/yy indicates the sampling ID and the year of sampling in the 21st century. Regression lines and R^2 are estimated with Microsoft excel 2010.

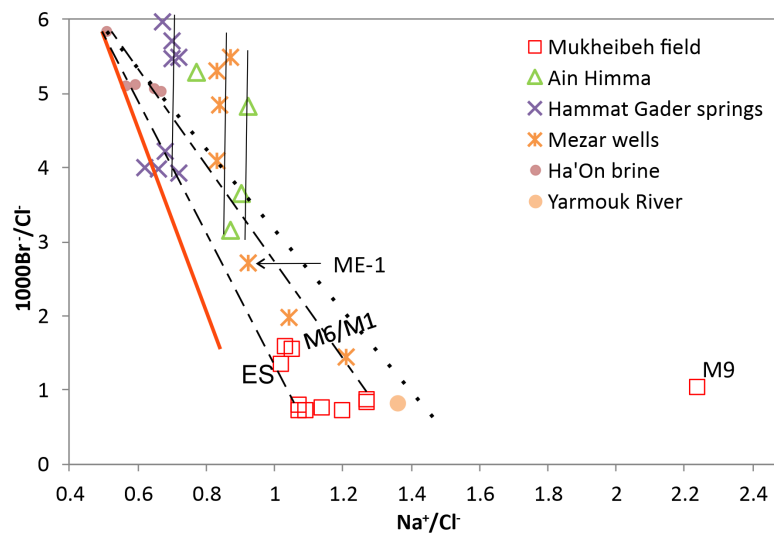


Figure 6. Cross plots of Na^+/Cl^- vs. $1000 \cdot \text{Br}^-/\text{Cl}^-$ of Ha'On brine analyses in the years 1961–2004 and of evaporated seawater (red line) [59,60]. Present-day Ha'On brine is a dilution product of the original Ha'On brine and basaltic-rock groundwater with $\text{Na}^+/\text{Cl}^- > 1$ (dashed line). The intersection of the dilution line with the trend line of the evaporated seawater approach: the values of Na^+/Cl^- and $1000 \text{ Br}^-/\text{Cl}^-$ of the original Tortonian Ha'On brine are 0.43 and 5.9, which significantly deviate from the ones measured in the years of 1961 and 2004 (Table 1). The vertical lines indicate leaching of Br^- from B2. The solid line represents mixing of Ha'On brine and basaltic-rock water in the Mukheibeh field. Code names are explained in Table 1.

4.2. Mixing of Basaltic Rock and Limestone Water

The cross plots of ε_{bw} and $\varepsilon_{\text{lmst}}$ show the distributions of the various types of water mixtures along the diagonal line (Figure 7). The red cross marks the arrays of either dominantly limestone or basaltic rock groundwater. The pure limestone water is presented by Mezar 2 and Mukheibeh 8 in the year 2013; the most basaltic-rock groundwater is among the Mukheibeh ones. Hammat Gader, Ain Himma and Mezar 3 cover the range of ε_{bw} between zero and 0.5. Most of the Mukheibeh waters (U1 and U2) are of the basaltic-water type, whereas the Mukheibeh subgroup U3 (with one exception) and the remaining groundwaters are of limestone water type.

The Mukheibeh field is characterized by mixing of basaltic rock and Ajloun limestone water with $\varepsilon_{\text{brine}}$ of 0.0019–0.004 of Ha'On brine (Table 3); in Ain Himma water $\varepsilon_{\text{brine}}$ varies between 0.0086 and 0.015. When fitting Hammat Gader and the Mezar waters to mixtures of Golan limestone and basaltic rock water and Ha'On brine, $\varepsilon_{\text{brine}}$ ranges between 0.019 and 0.031 for Hammat Gader and ME1 and ME 2. In contrast, ME3 reveals $\varepsilon_{\text{brine}}$ between 0.0022 and 0.0039, resembling Mukheibeh water. After substituting the Ha'on brine by Rosh Pinna brine in Hammat Gader and Mezar 1 and 2, the $\varepsilon_{\text{brine}}$ declines to 0.013–0.028 (Table A2a) as the result of the enhanced chlorinity of Rosh Pinna which is 36% higher than in Ha'On brine (Table 2). The maximum of volume of brine fraction is 0.03. The cross plots of ε_{bw} and $\varepsilon_{\text{brine}}$ suggest three different trends (Figure 8a). The low $\varepsilon_{\text{brine}}$ values of Mukheibeh water slightly increase with ε_{bw} . Although Ein Sahina and Ain Saraya discharge in the area of Hammat Gader, they plot together with Mezar 3 and the Mukheibeh data. The trends of Hammat Gader and Mezar wells 1 and 2 show the highest $\varepsilon_{\text{brine}}$ fractions, while Ain Himma plots slightly lower $\varepsilon_{\text{brine}}$.

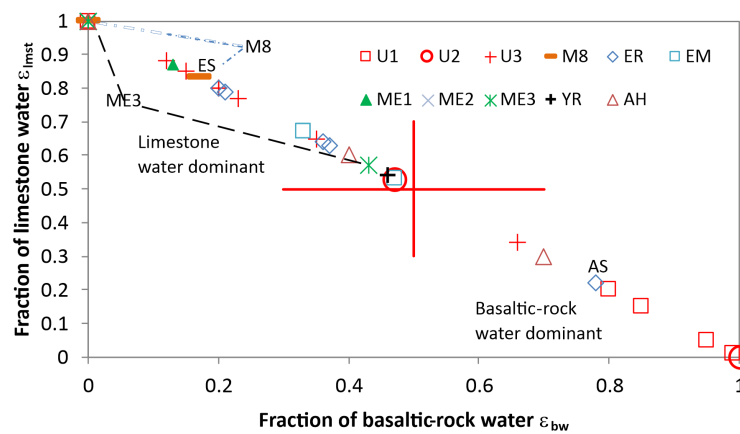


Figure 7. Cross plots of fractions of basaltic rock and limestone water in the Lower Yarmouk Gorge (LYG). Hammat Gader and Mezar are based on Golan limestone water; Mukheibeh water is related to Ajloun limestone water. M = Mukheibeh sample.

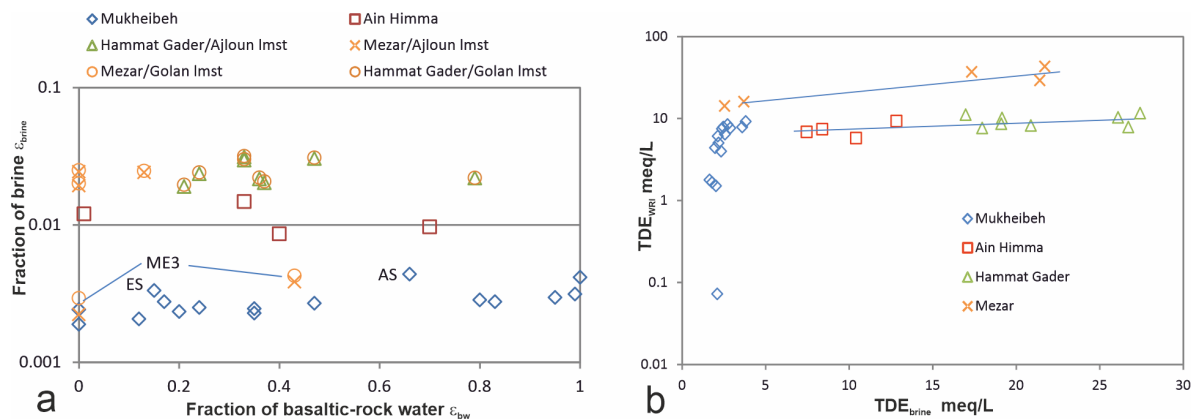


Figure 8. Cross plots of (a) ϵ_{bw} and ϵ_{brine} and (b) TDE_{brine} and TDE_{WRI} . These data are obtained under the assumption that ϵ_{brine} can be estimated after Equation (2). Hammat Gader and Mezar data are based on Golan limestone water and Ha'On brine. Mukheibeh water is related to Ajloun limestone water.

4.3. Contributions by Water/Rock Interaction

Following the two suggested approaches of salinization in Section 3.2, two results are obtained depending on the origin of Cl^- either from leaching of only evaporites or from leaching of evaporites (WRI) and brines. Both ways of estimations are documented in Table A2 and summarized in Table 3. The second approach yields the sum of TDE_{brine} and TDE_{WRI} and a maximum for TDE_{brine} (Table A2), whereas $\epsilon_{brine} = 0$ yields $TDE_{WRI+brine}$ in Table 3. TDE_{WRI} increases over two orders of magnitude in Mukheibeh groundwater. Contrastingly, the increase of TDE_{WRI} in each of Mezar, Ain Himma and Hammat Gader is less than a factor of two (Figure 8b). The contributions of TDE from water/rock interactions (TDE_{WRI}) are less those by brine (TDE_{brine}) in Hammat Gader, Mezar and most of the Ain Himma samples.

Table 3. Compilation of TDE values; brine fraction ϵ_{brine} ; and mineralogical compositions of water/rock interaction (WRI) and WRI+brine.

Code Group	Location	Sampling Date Year	Sum of TDE	TDE _{BW}	TDE _{est}	TDE _{brine} meq/L	TDE _{WRI}	Basaltic Water Fraction	Brine Water Fraction	ϵ_{brine} after Equation (4) diss(+)/precip(-)		$\epsilon_{\text{brine}} = 0$ after Equation (7) diss(+)/precip(-)		
								ϵ_{bw}	ϵ_{brine}	Gypsum	Calcite	Gypsum	Calcite	Halite
mmol/L														
Estimates based on basaltic-rock-, Ajloun limestone water and Ha'On brine														
U1	Mukheibeh 2	2016	15.07	5.47	6.09	2.57	6.41	0.95	0.00	0.47	0.56	0.47	0.62	1.25
U1	Mukheibeh 4	2016	15.79	0.00	12.37	1.63	1.79	0.00	0.00	0.48	−0.89	0.49	−0.85	0.79
U1	Mukheibeh 4	2013	16.75	4.78	6.88	2.39	7.48	0.83	0.00	0.45	0.83	0.46	0.88	1.16
U1	Mukheibeh 2		16.98	5.70	5.83	2.72	8.43	0.99	0.00	0.48	1.07	0.49	1.13	1.32
U1	Mukheibeh 1	2001	17.42	4.61	7.08	2.46	7.87	0.80	0.00	0.45	0.66	0.46	0.72	1.20
U2	Mukheibeh 6	2016	17.23	5.76	5.76	3.60	7.87	1.00	0.00	0.50	0.79	0.51	0.87	1.75
U2	Mukheibeh 7	2016	15.57	2.71	9.27	2.33	3.98	0.47	0.00	0.26	0.12	0.26	0.17	1.13
U3	Mukheibeh 10	2016	14.58	1.15	11.05	2.03	1.50	0.20	0.00	0.30	−0.68	0.30	−0.63	0.99
U3	Mukheibeh 5	2016	15.04	0.69	11.58	1.79	1.66	0.12	0.00	0.27	−0.80	0.27	−0.76	0.87
U3	Mukheibeh 11	2016	16.43	2.02	10.06	1.97	4.40	0.35	0.00	0.44	−0.46	0.45	−0.42	0.96
U3	Mukheibeh 9	2016	21.00	3.80	8.01	3.80	9.19	0.66	0.00	0.10	−0.19	0.11	−0.10	1.85
U3	Mukheibeh 8	2013	14.53	0.00	12.36	2.10	0.07	0.00	0.00	0.18	−0.99	0.18	−0.95	1.02
U3	Mukheibeh 8	2016	13.33	0.98	11.24	2.39	−0.29	0.17	0.00	0.02	−0.77	0.02	−0.73	1.16
U3	Mukheibeh 13	2016	14.53	1.38	7.33	2.17	5.02	0.24	0.00	0.24	1.17	0.25	−0.50	1.06
U3	Mukheibeh 13	2013	15.32	2.02	7.11	2.13	6.08	0.35	0.00	0.18	1.39	0.15	0.01	1.04
U3	Ein Sahina	2016	16.48	0.86	5.93	2.88	7.66	0.15	0.00	0.44	1.38	0.40	−0.44	1.15
AH	Ain Himma	2007	32.32	1.90	10.19	12.83	9.30	0.33	0.01	2.07	−1.48	2.10	−1.18	6.24
AH	Ain Himma	2001	28.47	0.06	12.31	10.41	5.75	0.01	0.01	1.41	−1.44	1.44	−1.19	5.06
AH	Ain Himma	2013	24.04	2.30	9.73	7.45	6.85	0.40	0.01	1.18	−0.20	1.19	−0.03	3.62
AH	Ain Himma	2016	23.53	4.03	7.74	8.39	7.40	0.70	0.01	1.21	0.04	1.23	0.24	4.08
Estimates based on basaltic-rock-, Golan limestone water and Ha'On brine														
AS	Ain Saraya	2016	33.51	4.55	5.78	19.11	8.63	0.79	0.02	1.11	1.00	1.15	1.45	9.29
EB	Ein Balsam	2016	31.43	2.13	5.82	17.96	7.65	0.37	0.02	1.27	0.86	1.31	1.28	8.73
ER	Ein Reach	2016	34.91	1.38	5.83	20.87	8.21	0.24	0.02	1.22	1.16	1.27	1.65	10.15
ER	Ein Reach	2000	33.95	1.21	5.83	16.97	11.15	0.21	0.02	1.09	1.71	1.13	2.11	8.25
ER	Ein Reach	2004	35.11	2.07	5.82	19.15	10.14	0.36	0.02	1.12	1.34	1.17	1.79	9.31
EM	Ein Makla	2016	40.38	2.71	5.81	26.70	7.87	0.47	0.03	1.42	1.17	1.48	1.80	12.98
EM	Ein Makla	2000	44.88	1.90	5.82	27.41	11.65	0.33	0.03	1.51	1.85	1.58	2.50	13.33
EM	Ein Makla	2004	42.23	1.90	5.82	26.10	10.32	0.33	0.03	1.44	1.29	1.50	1.91	12.69
ME1	Mezar 1	2016	29.05	0.75	5.84	21.41	1.80	0.13	0.02	0.02	0.45	0.07	0.96	10.41
ME2	Mezar 2	2016	43.01	0.00	5.85	21.71	15.45	0.00	0.02	3.42	0.07	3.47	0.59	10.56
ME2	Mezar 2	2001	37.01	0.00	5.85	17.34	13.82	0.00	0.02	2.80	0.11	2.85	0.52	8.43
ME3	Mezar 3	2016	16.05	2.48	5.81	3.70	6.54	0.43	0.00	0.12	1.12	0.09	−0.05	1.63
ME3	Mezar 3	2001	14.20	0.00	5.85	2.54	5.81	0.00	0.00	−0.01	1.39	−0.07	−0.77	0.94

Table 3. Cont.

Code Group	Location	Sampling Date Year	Sum of TDE	TDE _{BW}	TDE _{est}	TDE _{brine} meq/L	TDE _{WRI}	Basaltic	Brine	ϵ_{brine} after Equation (4)		$\epsilon_{\text{brine}} = 0$ after Equation (7)		
								Water	Water	diss(+)/precip(-)		diss(+)/precip(-)		
								Fraction	Fraction	Gypsum	Calcite	Gypsum	Calcite	Halite
								ϵ_{bw}	ϵ_{brine}	mmol/L		mmol/L		
Estimates based on basaltic-rock-, Golan limestone water and Rosh Pinna brine														
AS	Ain Saraya	2016	33.51	4.55	5.78	19.21	8.52	0.79	0.01	0.98	−0.11	1.15	1.45	9.29
EB	Ein Balsam	2016	31.43	2.13	5.82	18.06	7.55	0.37	0.01	1.15	−0.18	1.31	1.28	8.73
ER	Ein Reach	2016	34.91	1.38	5.83	20.99	8.09	0.24	0.02	1.08	−0.05	1.27	1.65	10.15
ER	Ein Reach	2000	33.95	1.21	5.83	17.06	11.06	0.21	0.01	0.98	0.73	1.13	2.11	8.25
ER	Ein Reach	2004	35.11	2.07	5.82	19.25	10.04	0.36	0.01	1.00	0.23	1.17	1.79	9.31
EM	Ein Makla	2016	40.38	2.71	5.81	26.85	7.72	0.47	0.02	1.24	−0.37	1.48	1.80	12.98
EM	Ein Makla	2000	44.88	1.90	5.82	27.57	11.50	0.33	0.02	1.33	0.26	1.58	2.50	13.33
EM	Ein Makla	2004	42.23	1.90	5.82	26.24	10.17	0.33	0.02	1.26	−0.22	1.50	1.91	12.69
ME1	Mezar 1	2016	29.05	0.75	5.84	21.53	1.68	0.13	0.02	−0.12	−0.79	0.07	0.96	10.41
ME2	Mezar 2	2016	43.01	0.00	5.85	21.83	15.33	0.00	0.02	3.27	−1.18	3.47	0.59	10.56
ME2	Mezar 2	2001	37.01	0.00	5.85	17.43	13.73	0.00	0.02	2.69	−0.90	2.85	0.52	8.43
ME3	Mezar 3	2016	16.05	2.48	5.81	3.72	6.52	0.43	0.00	0.09	0.91	0.13	1.21	1.80
ME3	Mezar 3	2001	14.20	0.00	5.85	2.56	5.80	0.00	0.00	−0.03	1.24	−0.01	1.45	1.24

From the estimated species i of WRI or WRI+brine (Table A2), the amounts of dissolved gypsum and calcite are given by $\text{SO}_4^{2-}/2$ and $(\text{Ca}^{2+} - \text{SO}_4^{2-})/2$ in mmol/L in Table 3. The amount of halite equals the amount of Cl^- in meq/L; (-) signs indicate precipitation; (+) values show dissolution. Calcite shows precipitation when fitting Hammat Gader and Mezar waters with Ajloun limestone water, which is not the case when using Golan limestone water. The compositions of brines from Hammat Gader and groundwater from Mezar 1 and 2 are estimated for various combinations of brines and fresh waters. The results of these mixing estimates are compiled in the lower parts of Tables A2 and 3. The differences in mixing either Ajloun or Golan limestone water with either Ha'On or Rosh Pinna brines yield similar results for gypsum dissolution but significantly different ones for the dissolution of calcite. In the presence of brines, calcite is precipitated from Ajloun limestone water, whereas in Golan limestone, calcite dissolves. In the absence of brine, some Hammat Gader waters dissolve calcite, and Mezar groundwater precipitates calcite when fitted to Golan limestone water. The estimates reveal dissolution of significant amounts of gypsum and calcite in waters from Ain Himma, springs of Hammat Gader and well Mezar 2, whereas Mukheibeh waters dissolve much smaller amounts of both minerals (Table 3). The dissolution of calcite and gypsum leads to enhancement of Ca^{2+} in Mukheibeh groundwater (Figure 5e). The increase of Mg^{2+} in groundwater (Figure 5f) is caused by high Mg^{2+} concentration in the admixed Ha'On brine.

The cross plots of calcite and gypsum reveal that their amounts are very similar and independent on the absence or presence of brine Equation (2). Gypsum is always dissolved, but calcite is both dissolved in Hammat Gader, Mezar and part of the Mukheibeh waters, and precipitated in the other part of Mukheibeh and Himma water (Figure 9a). The cross plots of halite and gypsum dissolution only reveal two trends between Mukheibeh at one end and either Hammat Gader waters or Mezar and Himma waters at the other end (Figure 9b).

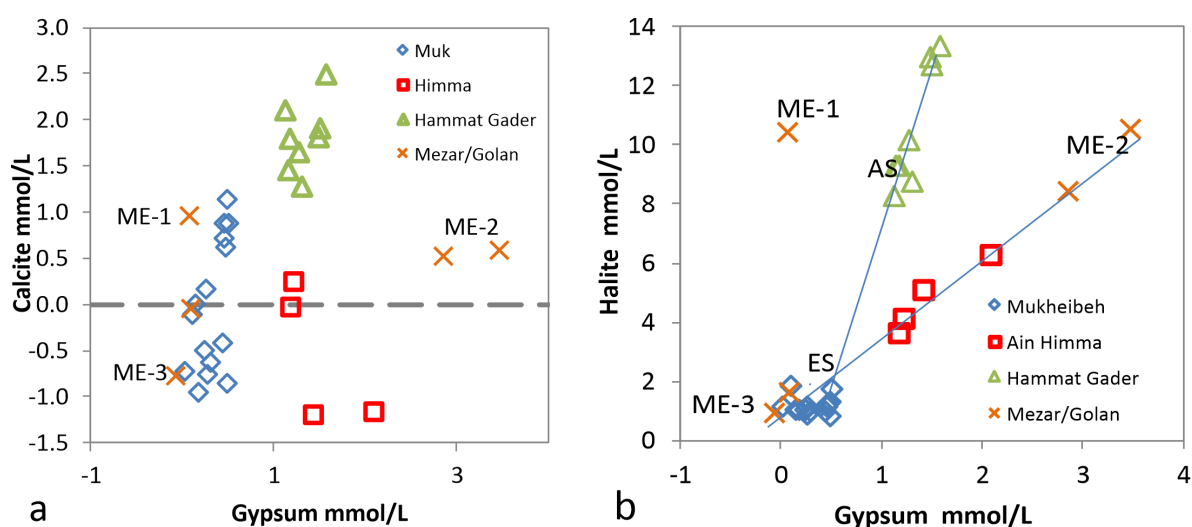


Figure 9. Cross plots of minerals. (a) Amounts of gypsum and calcite in WRI based on the assumption that ϵ_{brine} can be estimated after Equation (2). (b) Amounts of halite and gypsum in WRI+brine. Here ϵ_{brine} is assumed to be zero and its contribution appears together with those of the WRI.

4.4. Data Interpretation Through Multivariate Statistical Analysis

The results of the k -medoid clustering are visualized in terms of principal coordinates in the three-dimensional plot of Figure 10. Hereby a classical multidimensional scaling (otherwise known as *principal coordinates analysis* [62,63]) was performed on the dissimilarity matrix already computed by the clustering algorithm described above. These three first coordinates cumulatively explain 83% of the variance of the samples of Table 1. The spheres represent the positions of samples in the vectorial space of the principal coordinates; the similarly colored small dots indicate the corresponding projections on the three faces of the cube. In the C1/C2 plane Mukheibeh waters (code U1, U2, U3 in Table 1) yield

a curve which is far away from the projection of Hammat Gader samples (codes ER and EM). ME2 waters show some relationship to ME1. The projections onto C1/C2 and C1/C3 planes reveal that ME3 waters are closely associated with code U3 in plane C1/C2. Only in the plane C2/C3 are ME3 and U3 well separated. Ain Himma is well separated from Hammat Gader and Mukheibeh (U1-U3) in the C1/C2 plane.

To summarize, Figure 10 visualizes and validates the grouping of waters and brines based on the similarities in chemical analyses (restricted to major ions) compiled in Table 1. The different code groups in Figure 10 form either clusters or strings in space, thereby indicating constant or variable mixtures, respectively.

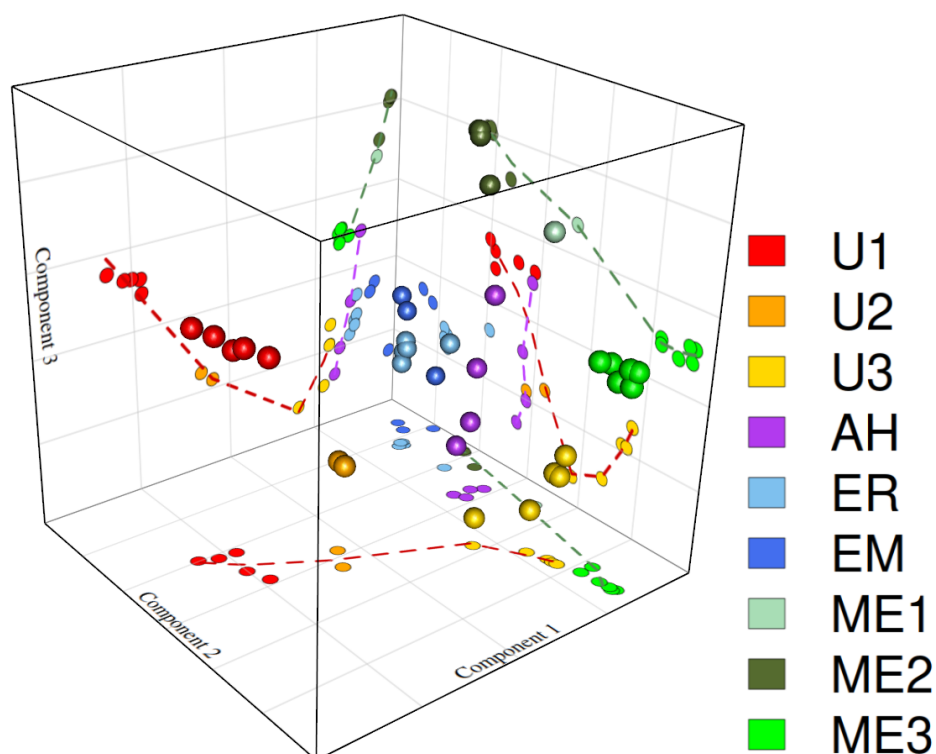


Figure 10. 3D visualization of the clustering of the water samples along the first three principal coordinates. The different colors of large spheres in three-dimensional space and the corresponding colored dots on the projection planes visualize the differences in composition of the groundwater in the Yarmouk Gorge subdivided into nine code groups (Table 1). This plot is based on concentrations in meq/L.

5. Discussion

5.1. Sources of Salinization

Because the fractions of basaltic rock and limestone water are based on interpretations of REY patterns, it should be kept in mind that the end member limestone water may have already dissolved some gypsum and halite. This may have led to overly high brine and limestone water fractions, due to which the fraction of basaltic-rock water was lowered. For similar reasons the true contribution of WRI may be slightly higher than derived in Table 3. Possible atmospheric contributions are minimized by selecting basaltic rock and limestone water with the lowest Cl^- concentrations.

The triplot visualizes the differences of the various local groundwaters and brines (Figure 11). The contributions $\text{TDE}_{\text{WRI}+\text{brine}}$, TDE_{bw} and TDE_{lmst} in groundwaters show a narrow cluster of Mezar wells 1 and 2, Hammat Gader and Ain Saraya samples, whereas water from Mukheibeh well field, Ain Himma and Mezar 3 covers a wide field between the dashed lines. The contributions in TDE from

brine and WRI are between 10% and 70%; contributions are 80% and 90% in Mukheibeh, Ain Himma and Mezar 3 and Hammat Gader and Mezar wells 1 and 2. These estimates do not really differ, if the sources of limestones water or brines are varied.

The Mukheibeh groundwater originates from an aquifer with a constant contribution of brine but increasing dissolution of calcite (Figure 9a). Calcite in code group U3 and Ain Himma is always precipitated (Table 3) contrasting the mixing in Hammat Gader, and Mezar wells 1 and 2. The mixture of Mount Hermon/Golan limestone water and Ha'On brine in Mezar 2 distinctly differs from Hammat Gader by enhanced contributions by WRI (Figures 8, 10 and 11). The correlation of gypsum and halite suggests dissolution of evaporites (Figure 9b).

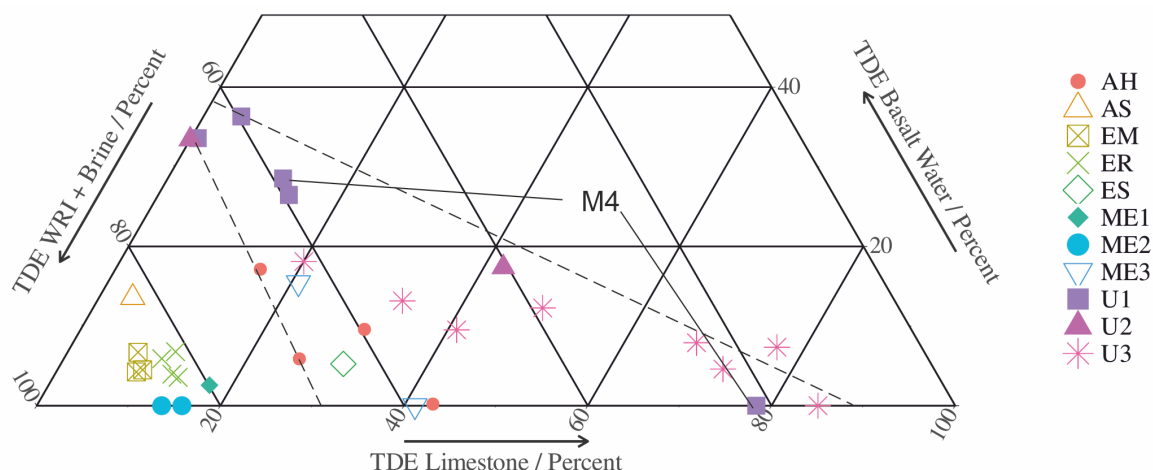


Figure 11. Ternary plot of contributions in %TDE of basaltic-rock-water, limestone water and WRI+brine. Several trends evolve. All trends seem to culminate in the Yarmouk River water. The dashed lines fix the array of Mukheibeh, Ain Himma, Mezar 3 and Ein Sahina. Clearly separated, the waters from Hammat Gader (EM, ER); Mezar 1 and 2; and Ain Saraya cluster in the lower left corner. With M4, a subgroup is indicated within the Mukheibeh U1 group (code refers to Table 3) visualizing the possibility of high variation in composition of groundwater from one and the same well.

In the assumed absence of brine the dissolution of halite amounts to about 1 mmol/L for Mukheibeh and Mezar 3; about 4–6 mmol/L in Ain Himma; and between 8 and 13 mmol/L in Hammat Gader and Mezar wells 1 and 2 (Table 3; Figure 11). Independent on the type of estimate, the dissolution of gypsum varies between 0 and 0.5 mmol/L in Mukheibeh and Mezar well 1 and 3 waters. It ranges from 1 to 3.5 mmol/L in Ain Himma, Hammat Gader and Mezar 2.

In the absence of deep brines, gypsum and calcite are dissolved in Hammat Gader and Mezar 1 and 3 in Golan limestone water. In the presence of Rosh Pinna brine instead of Ha'On brine, calcite often has to be precipitated, making the former less reasonable because the limestone water is already saturated with respect to calcite. Mezar 3 does not dissolve gypsum but calcite, particularly in the presence of Rosh Pinna brine (Table 3). Taking Ajloun limestone water and Ha'On brine, calcite is precipitated from groundwater of Hammat Gader and Mezar (Table A2), suggesting that Ajloun water is not present in any in these waters.

All groundwater mixes with brine present in aquifer rocks and interacts with aquifer rocks. The contribution of brine dominates the salinity of groundwater. The Tortonian Ha'On brine was identified in the study area. It is reasonable to assume that this brine infiltrated the Cretaceous (and probably Jurassic) limestone aquifers and is therefore omnipresent in the surroundings of the Yarmouk Gorge [29]. Estimates based on the contributions of Rosh Pinna brine abundantly lead to dissolution of calcite when applying Equation 2, which is unreasonable because in limestone aquifers calcite saturation should be attained.

5.2. Groundwater Divide between the Ajloun and the Golan Heights

Chemical similarities suggest that Mezar 3 on the northern Yarmouk River bank, but located very near to the LYF, produces groundwater of the Mukheibeh type (Figure 3). Ein Sahina and M5, both north of LYF, produce water of the Mukheibeh type (Table 1). Ain Saraya south of the Yarmouk River, just opposite of Hammat Gader but north of LYF, produces water typical for Hammat Gader (Table 1). Ain Himma, located southwest of the Yarmouk River but north of the LYF, is seemingly related to Hammat Gader brines (Figure 5). The thermohaline water of Hammat Gader seems to ascend along faults from greater depth. These examples of distribution of salinized groundwater indicate that probably not the Yarmouk River, but instead, the LYF delineates the groundwater divide between the Ajloun and the Golan Heights. LYF clearly separates the Mukheibeh well field with $0.002 < \varepsilon_{\text{brine}} < 0.004$ from the Mezar well field ($0.02 < \varepsilon_{\text{brine}} < 0.04$), Ain Himma ($0.009 < \varepsilon_{\text{brine}} < 0.013$) and Hammat Gader region ($0.02 < \varepsilon_{\text{brine}} < 0.04$).

Although the LYF follows the trend of the Yarmouk River, the chemical composition of local groundwater and brines is oriented according to the LYF and not the political border between Jordan and Israel given by the Yarmouk River. According to the regional differences, the transboundary flow may be influenced by local pumping on the Israeli side, the artesian outflow on the Jordanian side and recharge of the common aquifer on both sides of the LYF.

In well and spring water of the Mezar field and Hammat Gader region, significant changes in REY patterns [15] indicate variation in groundwater flow and mixing of basaltic rock and limestone waters (Figure 4b). Mezar 3 in 2008 produced water with the same REY pattern of Mezar 2, which definitely originates from the deep aquifer in the Golan. In Figure 11 Mukheibeh 2, 4 and 8 show high variations in composition within the Mukheibeh array of dashed lines. This behavior suggests that their flow system probably depends on pumping and recharge. The most extremely different compositions reveal waters from the wells Mezar 2 (depth –807 m) and Mezar 3 (depth –102 m), drilled few tens of meters apart. Mezar 3 water in 2001 and 2016 was of the Mukheibeh type (Figure 2). In 2008, Mezar 2 and 3 showed the same type of REY patterns which do not fit into Figure 4a [15].

Figures 5, 6, 8, 10 and 11 suggest different aquifers. The uppermost fresh water aquifer producing the Mukheibeh type is dominantly recharged by either basaltic-rock or limestone water. Some of the infiltrated water penetrates into deeper aquifers and leaches along their flow paths, evaporites and relics of brine. The deepest aquifer is that of Mezar 2. Hammat Gader originates from an aquifer which enables much less contact with gypsum but slightly more with halite, whereas Mezar 2 and Himma water had more contact with gypsum and less halite.

6. Conclusions

The combined use of REY, multivariate statistical analysis, chemical features and geological features represents a new approach with which to semiquantitatively determine the contributions of various sources of salinization in groundwater, such as basaltic-rock and limestone water, water/rock interactions and leaching fossil brines. The contribution of atmospheric precipitation is considered as part of the recharge water or to be negligible in water with lowest Cl^- concentrations.

The basaltic-rock-dominated waters show the lowest salinities. The basaltic-rock waters show higher TDE_{WRI} than $\text{TDE}_{\text{brine}}$. The basaltic-rock groundwaters from the Hauran Plateau mix with limestone water from either the Ajloun or the Golan Heights depending on the positions of springs and wells south or north of the Lower Yarmouk Fault, respectively. South of LYF, the artesian Mukheibeh well field produces in its central segment, groundwaters of almost pure basaltic-rock type with a low contribution of limestone water and <0.3 vol-% of Tertiary brine, hosted in deep Cretaceous and Jurassic formations. In the Mukheibeh field the total dissolved equivalents of solutes gained by water/rock interaction and mixing with brine, $\text{TDE}_{\text{WRI}+\text{brine}}$, amounts to 10%–65%. Further away from this center, the contribution of limestone water originating from the Ajloun Mountains increases. The limestone waters manifest the highest salinities. These waters occur in the region of Mezar and Hammat Gader. North of the LYF, the Mezar wells, the springs of Hammat Gader and Ain Himma,

produce dominantly limestone water, which contains 0.14–3 vol-% of the Tertiary brine and hence possess variable salinity.

The water of Ain Himma composition sometimes resembles that of Mezar 2, suggesting groundwater from great depth. Ain Saraya south of the Yarmouk River produces a similar type of saline water as in Hammat Gader north of the LYF. The total dissolved equivalents of solutes gained by water/rock interaction and mixing with brine, $TDE_{WRI+brine}$, amount to 60%–70% of total salinity in Ain Himma and Mezar 3 well; 55%–70% in the springs of Hammat Gader; and 80%–90% in wells Mezar 1 and 2. The uniform trend of Mg^{2+} with Cl^{-} in all groundwaters except the Mukeheibeh ones suggest leaching of the Tertiary Ha'On brine which is of $Mg^{2+}-Cl^{-}$ type. The different dilution trends of other dissolved species such as Na^{+} , K^{+} , Ca^{2+} , Br^{-} and SO_4^{4-} of either Hammat Gader or Mezar/Ain Himma indicate differences in occurrences of evaporite minerals in the respective aquifers. The salinity of groundwater in the study region is mainly due to (i) leaching of remnants of Tertiary Rift brine and not to mixing with relicts of the Triassic brine, and (ii) water/rock interactions, such as dissolution of halite, gypsum and calcite. Running sub-parallel to the Yarmouk River, the LYF seems to be the actual groundwater divide between the Ajloun and the Golan Heights. The Lower Yarmouk fault seemingly acts as the divide between the Ajloun and the Golan Heights-dominated groundwater. Since the Yarmouk River represents the international border between Jordan and Israel, these examples suggest transboundary flow over short distances, possibly through local N–S-trending faults and fissures beneath the river.

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Conflicts of Interest: The authors declare that they have no conflicts of interest.

Abbreviations

The following abbreviations are used in this manuscript:

LYG	Lower Yarmouk Gorge
WRI	Water/rock interaction
ϵ_{brine}	Fraction of brine
ϵ_{bw}	Fraction of basaltic-rock water
ϵ_{lmst}	Fraction of limestone water
$c_{i,agw}$	Concentration of species i in analyzed groundwater
$c_{i,bw}$	Concentration of species i in basaltic-rock water
$c_{i,brine}$	Concentration of species i in brine
$c_{i,est}$	Concentration of mixed basaltic-rock and limestone water
$c_{i,WRI}$	Concentration of species i due to WRI
TDE_{bw}	Total dissolved equivalents due to weathered basalt
TDE_{lmst}	Total dissolved equivalents due to dissolved limestones
TDE_{agw}	Total dissolved equivalents in analyzed groundwater
TDE_{brine}	Total dissolved equivalents in brine
TDE_{est}	Total dissolved equivalents of estimated mixture of basaltic rock and limestone water
TDE_{WRI}	Total dissolved equivalents due to water/rock interaction
$TDE_{WRI+brine}$	Total dissolved equivalents due to water/rock interaction and mixing with brine

Table A1. Analyses of Ha'On brine, Triassic brines of Rosh Pinna, basaltic spring waters from the Golan Heights and the Hauran Plateau and limestone waters from the Ajloun Mountains and Golan Heights. Because the analyses of Ha'On brine are from different wells for each group, the average and standard deviation (%) are given. The extrapolated concentrations of Mg^{2+} , K^+ , Na^+ , Cl^- and Br^- of the original Ha'On are derived from data on evaporated seawater [59,60].

Reference	Well	Sampling Date	Ca^{2+} mg/L	Mg^{2+} mg/L	K^+ mg/L	Na^+ mg/L	Cl^- mg/L	SO_4^{2-} mg/L	Br^- mg/L	HCO_3^- mg/L	Na/Cl eq.ratio	1000Br/Cl eq.ratio	Mg/Ca eq.ratio
Extrapolated original Ha'On (Figure 6)				47,500	14,200	50,000	184,000		2250		0.42	5.43	
Late Tertiary Ha'On brine													
Mandel 1965 [53]	D-1071	1961	1476	2100	317	8633	21971	32.5	266.6	384	0.61	5.38	2.34
Mandel 1965 [53]	D-1071	1964	1427	2141	285	8990	22624	tr	153.2	341	0.61	3.00	2.47
Mandel 1965 [53]	D-1071	1965	1285	2258	296	9093	23045	3.7		186	0.61		2.89
	Average		1396	2166	299	8906	22547	18	210	304	0.61	4.13	2.55
	Dilution factor			22	47	6	8		11				
Bergelson et al. 1999 [64]	D-1071	1993	593	971	409	4040	9600	112	109	427	0.65	5.04	2.69
Bergelson et al. 1999 [64]	D-1071	1993	589	1000	414	4209	10188	98.6		537	0.64		2.79
	Average		591	985	411	4125	9894	106	109	482	0.64	4.89	2.74
	Dilution factor			48	35	12	19		21				
Mandel 1965 [53]	D-1072	1961	653	1645	185	6103	14670	117	179	1195	0.64	5.41	4.15
Bergelson et al. 1999 [64]	D-1072	1993	599	1140	262	4382	11394	27.4	131	439	0.59	5.10	3.13
Bergelson et al. 1999 [64]	D-1072	2000	1078	1657	442	5212	15800	59.8	207	504	0.51	5.81	2.53
Siebert et al. 2014 [2]	D-1072	2004	758	1169	394	4276	11668	644	133	576	0.56	5.07	2.54
	Average		772	1403	321	4993	13383	212	163	678			
	Dilution factor			34	44	10	14		14				
Klein BenDavid et al. 2004 [65]	D-1072	2003	559	910	425	3687	8486	765	95.6	548	0.67	5.00	2.68
	Dilution factor			52.2	33.4	13.6	21.7		23.5				
	Ha'On average		902	1499	343	5863	14945	207	159	514	0.60	4.73	2.74
	Dilution factor			31.7	41.4	8.5	12.3		14.1				
Triassic brine													
		depth											
Starinski 1974 [66]	Rosh Pinna 1	3845-3864	26946	2918	1157	33415	105349	347		171	0.49		0.18
Starinski 1974 [66]	Rosh Pinna 1	2486-2586	4794	724	358	8004	22819	1161		370	0.54		0.25
Limestone water from Mount Hermon Massif													
Sie 786/08	Ein Dan	2008	65.9	5.4	0.7	4.3	9.8	9.0			0.68		0.14
Sie 787/08	Ein Banyas	2008	72.3	12.4	1.5	9.9	12.9	57.3			1.18		0.28
	Average		69.1	8.9	1.1	7.1	11.3	33.1			0.97		0.21
Limestone water from the Ajloun													
Siebert et al. 2014 [2]	Ain Al Azal		96.0	16.8	0.6	16.1	32.4	14.6		291			
Siebert et al. 2014 [2]	Ain Al Murarar		101	8.04	2.29	16.6	24.3	6.06		277			
	Average		98.6	12.4	1.43	16.4	28.3	10.337		284			

Table A1. Cont.

Reference	Well	Sampling Date	Ca ²⁺ mg/L	Mg ²⁺ mg/L	K ⁺ mg/L	Na ⁺ mg/L	Cl ⁻ mg/L	SO ₄ ²⁻ mg/L	Br ⁻ mg/L	HCO ₃ ⁻ mg/L	Na/Cl eq.ratio	1000Br/Cl eq.ratio	Mg/Ca eq.ratio
Limestone water from the Golan													
Siebert et al. 2014 [2]	Alonei HaBashan 3		9.9	8	4.6	43.3	17.2	4.9		132			
Siebert et al. 2014 [2]	Allonei HaBashan 8		5.63	3.97	3.31	56.2	18.4	4.695		124			
	Average		7.77	5.99	3.96	49.8	17.8	4.7975		128			
Basaltic-rock water from Golan Heights and Hauran Plateau													
Dafny et al. 2006 [7]	Average (n = 60)		30.8	17.2	3.8	27.8	29.7	10.8	0.022	177	1.44	0.33	0.92
Kattan 1997 [11]	Average (n = 21)		26.8	12.1	3.6	25.9	29.6	15		126			
	Weighted average		29.8	15.9	3.75	27.3	29.7	11.9	0.022	164			

Table A2. Estimation of brine and water/rock contributions to the salinity of groundwaters. (a) The contribution of the basaltic-rock water, the sum of basaltic rock and limestone water; the contributions of brine are estimated based on brine after Equation (2). The difference between analyzed and estimated concentrations yield the contribution of water/rock interaction here presented as gypsum and calcite dissolution. (b) Similar to (a) but brine is set to zero. Consequently, only the sum of water/rock interaction and brine is obtained.

Sampling		Estimated groundwater $c_{i,est} = c_{i,bw} + c_{i,lmsst}$										Contribution of brine to estimated groundwater $c_{i,est}$										ε_{brine} after Eq. (4) Difference of analysed & estimated conc.: $c_{i,WRI} = c_{i,agw} - c_{i,est} - c_{i,brine}$										$\varepsilon_{brine} = 0$; TDE _{WRI} after Eq. (7) Difference of analysed & estimated conc.: $c_{i,WRI} = c_{i,agw} - c_{i,est}$									
		Ca Mg K Na Cl SO ₄ HCO ₃ TDE _{gw}										Ca Mg K Na Cl SO ₄ HCO ₃ TDE _{brine}										Ca Mg K Na Cl SO ₄ HCO ₃ TDE _{WRI}										Ca Mg K Na Cl SO ₄ HCO ₃ TDE _{WRI}									
		Year	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	Year	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	Year	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	Year	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	meq/l	Year	meq/l	meq/l	meq/l	meq/l	meq/l
Code	Source	Year	Ca	Mg	K	Na	Cl	SO ₄	HCO ₃	TDE _{gw}	Ca	Mg	K	Na	Cl	SO ₄	HCO ₃	TDE _{brine}	Ca	Mg	K	Na	Cl	SO ₄	HCO ₃	TDE _{WRI}	Gypsum	Calcite	Ca	Mg	K	Na	Cl	SO ₄	HCO ₃	TDE _{WRI}	Gypsum	Calcite	Halite		
Estimates based on basaltic-rock-, Ajloun limestone water and Ha'On brine																																									
U1	Mukheibeh 2	2016	1.70	0.58	0.82	0.07	0.34	0.22	2.36	6.09	0.13	0.37	0.03	0.76	1.25	0.01	0.02	2.57	2.06	1.20	-0.77	0.87	0.94	2.12	6.41	0.47	0.56	2.20	1.56	-0.74	1.62	1.25	0.95	2.14	8.98	0.47	0.62	1.25			
U1	Mukheibeh 4	2016	4.93	1.02	0.04	0.71	0.80	0.21	4.66	12.37	0.09	0.23	0.02	0.48	0.79	0.01	0.02	1.63	-0.81	1.05	0.02	0.55	0.97	0.02	1.79	0.48	-0.89	-0.73	1.28	0.04	1.03	0.80	0.97	0.04	3.42	0.49	-0.85	0.79			
U1	Mukheibeh 4	2013	2.11	0.63	0.72	0.15	0.40	0.22	2.65	6.88	0.12	0.34	0.02	0.70	1.16	0.01	0.02	2.39	2.57	1.41	-0.67	0.82	0.91	2.44	7.48	0.45	0.83	2.69	1.75	-0.64	1.52	1.16	0.92	2.46	9.87	0.46	0.88	1.16			
U1	Mukheibeh 2	2013	1.57	0.56	0.85	0.05	0.32	0.22	2.26	5.83	0.14	0.39	0.03	0.80	1.32	0.01	0.03	2.72	3.09	1.52	-0.80	0.83	0.96	2.82	8.43	0.48	1.07	3.23	1.91	-0.77	1.64	1.32	0.97	2.85	11.15	0.49	1.13	1.32			
U1	Mukheibeh 1	2001	2.21	0.65	0.69	0.17	0.41	0.22	2.72	7.08	0.13	0.35	0.03	0.73	1.20	0.01	0.02	2.46	2.23	1.47	-0.64	0.80	0.90	3.12	7.87	0.45	0.66	2.35	1.82	-0.62	1.52	1.20	0.91	3.14	10.33	0.46	0.72	1.20			
U2	Mukheibeh 6	2016	1.53	0.55	0.86	0.04	0.32	0.22	2.24	5.76	0.19	0.51	0.04	1.06	1.75	0.02	0.03	3.60	2.58	1.32	-0.81	1.03	1.01	2.74	7.87	0.50	0.79	2.77	1.83	-0.77	2.09	1.75	1.03	2.77	11.47	0.51	0.87	1.75			
U2	Mukheibeh 7	2016	4.93	1.02	0.04	0.71	0.80	0.21	4.66	12.37	0.12	0.33	0.02	0.69	1.13	0.01	0.02	2.33	0.75	1.25	-0.37	0.61	0.72	1.19	3.98	0.87	0.73	6.49	2.75	-0.55	3.44	1.13	0.82	1.94	6.31	0.26	-1.63	1.13			
U3	Mukheibeh 10	2016	4.25	0.93	0.20	0.58	0.70	0.22	4.17	11.05	0.11	0.29	0.02	0.60	0.99	0.01	0.02	2.03	-0.76	0.84	-0.13	0.87	0.60	0.08	1.50	0.30	-0.68	-0.65	1.13	-0.11	1.47	0.99	0.61	0.10	3.53	0.30	-0.63	0.99			
U3	Mukheibeh 5	2016	4.52	0.97	0.14	0.63	0.74	0.22	4.37	11.58	0.09	0.26	0.02	0.53	0.87	0.01	0.02	1.79	-1.06	0.75	-0.06	0.88	0.53	0.62	1.66	0.27	-0.80	-0.97	1.01	-0.04	1.41	0.87	0.54	0.63	3.46	0.27	-0.76	0.87			
U3	Mukheibeh 11	2016	3.74	0.86	0.33	0.48	0.63	0.22	3.81	10.06	0.10	0.28	0.02	0.58	0.96	0.01	0.02	1.97	-0.04	2.07	-0.27	0.85	0.88	0.90	4.40	0.44	-0.46	0.96	2.35	-0.25	1.44	0.96	0.89	0.92	6.37	0.45	-0.42	0.96			
U3	Mukheibeh 9	2016	2.69	0.71	0.58	0.27	0.48	0.22	3.06	8.01	0.20	0.54	0.04	1.12	1.85	0.02	0.04	3.80	-0.18	1.13	-0.21	3.83	0.20	4.43	9.19	0.10	-0.19	0.01	1.67	-0.17	4.95	1.85	0.22	4.46	12.99	0.11	-0.10	1.85			
ME3	Mukheibeh 8	2013	4.92	1.02	0.04	0.71	0.80	0.21	4.66	12.36	0.11	0.30	0.02	0.62	1.02	0.01	0.02	2.10	-1.63	0.32	0.08	0.99	0.35	-0.04	0.07	0.18	-0.99	-1.53	0.62	0.10	1.61	1.02	0.36	-0.02	2.16	0.18	-0.95	1.02			
U3	Mukheibeh 12	2016	4.34	0.94	0.18	0.60	0.72	0.22	4.25	11.23	0.12	0.34	0.02	0.70	1.16	0.01	0.02	2.39	-1.51	0.36	-0.07	1.13	0.94	-0.24	-0.29	0.02	-0.77	-1.40	0.70	-0.05	1.84	1.16	0.05	-0.22	2.09	0.02	-0.73	1.16			
U3	Mukheibeh 13	2016	0.66	0.91	0.23	0.55	0.68	0.22	4.08	7.33	0.11	0.31	0.02	0.64	1.06	0.01	0.02	2.17	-2.82	0.84	-0.16	0.81	0.48	0.23	5.02	0.24	1.17	-0.51	1.15	-0.14	1.45	1.06	0.49	0.25	3.75	0.25	-0.50	1.06			
U3	Mukheibeh 13	2013	0.79	0.51	0.37	1.42	0.44	0.14	2.15	5.82	0.13	0.36	0.03	0.74	1.23	0.01	0.02	2.53	3.13	1.18	-0.30	-0.27	0.36	2.87	6.97	0.18	1.39	0.31	1.20	-0.23	1.42	1.04	0.29	1.23	5.26	0.15	0.01	1.04			
ES	Ein Sahina	2016	0.56	0.50	0.21	1.84	0.47	0.12	2.12	5.84	0.15	0.41	0.03	0.85	1.40	0.01	0.03	2.88	3.64	1.39	-0.16	-0.78	0.89	2.78	7.76	0.44	1.38	-0.07	1.35	-0.08	1.30	1.15	0.80	0.64	5.10	0.40	-0.44	1.15			
AH	Ein Himma	2007	3.81	0.87	0.31	0.49	0.64	0.22	3.86	10.19	0.67	1.83	0.13	3.78	6.24	0.06	0.12	12.83	1.19	0.11	-0.08	1.05	4.15	2.89	9.30	2.07	-1.48	1.86	1.94	0.05	4.82	6.24	4.21	3.02	22.13	2.10	-1.18	6.24			
AH	Ein Himma	2001	4.89	1.02	0.04	0.70	0.79	0.21	4.63	12.31	0.54	1.48	0.11	3.07	5.06	0.05	0.10	10.41	-0.06	0.37	0.21	1.63	2.82	0.78	5.75	1.41	-1.44	0.48	1.85	0.31	4.69	5.06	2.87	0.88	16.16	1.44	-1.19	5.06			
AH	Ein Himma	2016	3.57	0.84	0.37	0.44	0.61	0.22	3.69	9.73	0.39	1.06	0.08	2.19	3.62	0.04	0.07	7.45	1.94	0.74	-0.21	1.03	2.35	1.01	6.85	1.18	-0.20	2.33	1.80	-0.14	3.23	3.62	2.39	1.08	14.31	1.19	-0.03	3.62			
AH	Ein Himma	2016	2.55	0.69	0.61	0.24	0.46	0.22	2.96	7.74	0.44	1.20	0.09	2.47	4.08	0.04	0.08	8.39	2.51	0.58	-0.45	1.38	2.43	0.96	7.40	1.21	0.04	2.95	1.77	-0.37	3.84	4.08	2.87	1.04	15.79	1.23	0.24	4.08			
AS	Ain Saraya	2016	2.25	0.65	0.69	0.18	0.42	0.22	2.75	7.15	0.99	2.70	0.19	5.59	9.23	0.09	0.18	18.98	3.27	-0.23	-0.57	1.19	2.19	1.54	7.38	1.09	0.54	4.25	2.48	-0.38	6.77	9.23	2.28	1.73	26.36	1.14	0.99	9.23			
ER	Ein Balsam	2016	3.67	0.85	0.34	0.46	0.62	0.22	3.76	9.93	0.92	2.50	0.18	5.17	8.55	0.09	0.17	17.58	1.41	-0.56	-0.21	0.01	2.47	0.80	3.93	1.23	-0.53	2.33	1.95	-0.03	5.19	8.55	2.55	0.97	21.51	1.28	-0.11	8.55			
ER	Ein Reach	2016	4.11	0.91	0.23	0.55	0.68	0.22	4.08	10.79	1.06	2.91	0.21	6.01	9.92	0.10	0.20	20.41	1.32	-0.77	-0.08	0.40	2.35	0.50	3.71	1.17	-0.51	2.39	2.13	0.12	6.41	9.92	2.45	0.70	24.12	1.23	-0.03	9.92			
ER	Ein Reach	2000	4.22	0.93	0.21	0.57	0.70	0.22	4.15	10.98	0.86	2.35	0.17	4.85	8.02	0.08	0.16	16.49	2.04	-0.02	-0.06	0.81	2.10	1.62	6.48	1.05	-0.03	2.90	2.33	0.10	5.66	8.02	2.18	1.78	22.97	1.09	0.36	8.02			
ER	Ein Reach	2004	3.71	0.85	0.33	0.47	0.63	0.22	3.79	9.99	0.98	2.67	0.19	5.52	9.12	0.09	0.18	18.76	2.04	-0.28	-0.20	0.80	2.17	1.82	6.36	1.09	-0.07	3.01	2.40	-0.01	6.32	9.12	2.27	2.01	25.12	1.13	-0.37	9.12			
EM	Ein Makla	2016	3.33	0.80	0.42	0.40	0.57	0.22	3.52	9.27	1.37	3.76	0.27	7.77	12.83	0.13	0.26	26.38	2.79	-1.35	-0.26	0.97	2.78	-0.20	4.73	1.39	0.01	4.17	2.41	0.01	8.73	12.83	2.91	0.06	31.11	1.45	0.63	12.83			
EM	Ein Makla	2000	3.81	0.87	0.31	0.49	0.64	0.22	3.86	10.19	1.41	3.85	0.27	7.95	13.13	0.13	0.26	27.00	3.70	-1.08	-0.12	0.79	2.95	1.45	7.69	1.47	0.38	5.11	2.77	0.15	8.74	13.13	3.08	1.72	34.69	1.54	1.01	13.13			
EM	Ein Makla	2004	3.81	0.87	0.31	0.49	0.64	0.22	3.86	10.19	1.34	3.66	0.26	7.56	12.49	0.13	0.25	25.69	2.44	-1.07	-0.13	1.07	2.80	1.25	6.35	1.40	-1.18	3.78	2.59	0.13	8.63	12.49	2.93	1.49	32.04	1.46	0.43	12.49			
ME1	Meizar 1	2016	4.49	0.96	0.14	0.62	0.74	0.22	4.34	11.51	1.09	2.98	0.21	6.15	10.15	0.10	0.20	20.88	-2.98	-2.54	-0.02	3.23	-0.05	-0.98	-3.34	-0.03	-1.46	-1.89	0.44	0.19	3.38	10.15									

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