Pedo-Chemical Studies on Saline Playas in the Arabian Shelf

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دراسات بيدو كيميانية على بعض البلايا الملحية (السبخات) في الرصيف (الساحلي) العربي

الملخص: يهدف هذا البحث إلى دراسة بعض الصفات البيدوكيمائية للبلايا الداخلية والساحلية الموجودة في تكوينات الرصيف (الساحلي) العربي. لقد أوضحت نتائج الدراسة أن معظم البلايا ذات اسطح مستوية إلى شبه مستوية مع تواجد قشور ملحية متباينة السمك . كما أظهرت النتائج أن القطاعات الممثلة للسبخات ذات ملوحة مرتفعة مقارنة بالمناطق المحيطة بها مسع سيادة أملاح الكلوريدات والكبريتات الذائبة للصوديوم متبوعا بالكالسيوم أو المغنيسيوم . ويتشابه تركيب ملوحة الطبقات السبخات السنخات المختلفة لقطاعات السبخات السبخات السبخات الساحلية مع تركيب مياه الخليج بينهما يكون تركيب ملوحة الطبقات المختلفة لقطاعات السبخات الداخلية أكثر تأثرا بالصرف من المناطق المجاورة لها والظروف البيئية المحيطة بها . وكان ذلك واضحا من نسبب بعصض الكاتيونات إلى بعضها الأخر وكذلك معامل الارتباط بين درجة التوصيل الكهربائي (ملوحة التربة) وكسلا مسن السورون والمغنيسيوم الذائب وكان معامل الارتباط كبيرا في حالة السبخات الساحلية مقارنة بمثيلتها في السبخات الداخلية. ولذلك نجد أن من أكثر العمليات البيدولوجية تأثيرا على تكوين السبخات الساحلية هي حركة المهاه الجوفية الغنية بسالأملاح و المتساشرة أن من أكثر العمليات البيدولوجية تأثيرا على تكوين السبخات الساحلية مقارنة بالعمليات الساخلية بها مثل عمليات الغسيل وتركيز الأملاح نتيجة للتبخر . وأوضحت نتائج تحليل القشرة الملحية سيادة أملاح الكلوريدات والكبريتات الذائبة للصوديوم والمغنيسيوم والكالسيوم ثم البوتاسيوم . لقد أكنت التحليلات باستخدام الأشعة السينية تواجد معادن الساحلية والجبس والكربونات بالاضافة الى معدني الابسومايت والبلودايت . وتعزى الاختلافات الرئيسية بيسن السبخات الساحلية والدخلية إلى مصدرها وتكوينها وعمق الماء الأرضي وبالتالي التركيب الكيميائي للقشرة الملحية لكل منهما.

ABSTRACT: This research was undertaken to study the pedo-chemical characteristics of selected playas (sabkhahs) in the Arabian Shelf. Results indicated that most of the sabkhahs are flat to almost flat having salt crusts with variable thickness. Profiles representing the sabkhah surface are extremely saline compared with that of the surrounding landscapes. The dominant soluble salts are Cl⁻ and SO_4^{-2-} of Na^+ followed by Ca^{2+} or Mg^{2+} . The compositions of the coastal sabkhahs profile layers are closely related to the Gulf water composition while the inland playas are more associated with local conditions. This was confirmed from the similarity in salt levels, ion molar ratios and relatively high correlation coefficients between the electrical conductivity of soluble salts (EC_e) and either soluble B^{3+} ($r^2 = 0.80$) or soluble Mg^{2+} ($r^2 = 0.60$) in coastal compared with that of the inland sabkhahs ($F^2 = 0.17$) and ($F^2 = 0.37$), respectively. The main pedological process affecting the formation of coastal sabkhahs is the upward movement of the water table which is rich in salts originating from the Gulf waters either directly or through wind action. Leaching, accumulation of sediments and evaporative concentrations of the solutes are the processes prevailing in the inland sabkhahs. Salt crusts are dominated by Cl⁻ and SO_4^{2+} of Na^+ , Mg^{2+} and Ca^{2+} salts while K^+ salts are relatively low. X-ray diffraction patterns confirm the presence of halite, gypsum and carbonate minerals. Epsomite and bloedite minerals were identified in both coastal and inland sabkhahs. Specific minerals and relative abundance was quite variable between the salt crusts due to the characteristic conditions of each sabkhah. The major differences between the inland and coastal sabkhahs are related to the source, composition and depth of the water table and consequently the composition and mineralogy of the surface salt crusts.

Playa is the geological term for the flat and generally barren lower portions of arid basins of internal drainage that periodically flood and accumulate sediments (Neal, 1965). This is an important characteristic that makes playa the flattest of all land forms. Sabkhah is the term applied to playas in many Middle East countries. Moreover, Al-Sayari and Zotl (1978) reported that sabkhahs are saline flats underlain

by clay, silt and sand, and often encrusted with salt. The Arabian Shelf is one of the two major land masses forming the Arabian Peninsula. This shelf lies to the east of the Arabian shield, where it forms about two thirds of the peninsula and is composed mainly of sedimentary rocks among which scattered dry lakes are present in depressional areas. A common feature of the deserts in this region is the topographic depressions

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known as playa lakes, which usually have no drainage outlets for runoff. When the lakes dry up, mineral salts are left behind. There are two types of sabkhahs. The first type is the coastal sabkhah which is a super tidal surface produced by depositional overlap of marine sediments. In some cases, eolin deposits are also present. The associated brines are derived primarily from sea water. The second type is the inland sabkhah, which lies away from the coast, and represents areas of equilibrium between eolian deposits and deflation, controlled by the local water table. Evaporation through the surface causes the formation of brine and the precipitation of evaporite minerals (Al-Sayari and Zotl, 1978). Saudi Arabia has large numbers of both coastal and inland sabkhahs. These sabkhahs result from the varying influences of geological structure. surface features, climatological conditions and the location from the saline water sources. Sabkhahs are common features along the shoreline from Kuwait to the southern end of the Arabian Gulf. Sabkhat Matti, southeast of Qatar, is the largest of these with an area of about 6,000 km2 (Al-Sayari and Zotl, 1978). El-Farra (1978) reported on the number of sabkhahs existing in the different regions of the Arabian shelf,

focusing mainly on the eastern, western and Najd regions. The eastern regions include sabkhahs identified in the Gulf coast, Al-Hofuf and east Al-Rubh Alkhali. El-Farra reported that there were 150 sabkhahs along the Arabian Gulf (i.e. Sikak, 800 km²; ar Riyas, 900 km²; Salwah and Mashakhil sabkhahs) and about 30 sabkhahs east of Al-Hofuf. It was clear after reviewing the literature that there is insufficient data published on the chemical and mineralogical characteristics of the sabkhahs in the Arabian Shelf.

The main objective of our study was to examine the pedo-chemical characteristics of selected inland and coastal sabkhahs formed under the Arabian Shelf formations. A second objective was to evaluate the effect of saline conditions on the mineralogical composition of selected surface salt crusts of sabkhahs.

Materials and Methods

DESCRIPTION OF THE STUDIED SABKHAHS: Six sabkhahs were selected for this study to represent the coastal and inland sabkhahs in eastern and central regions of the Arabian Shelf (Figure 1). Rumadan sabkhah is located in the eastern region near

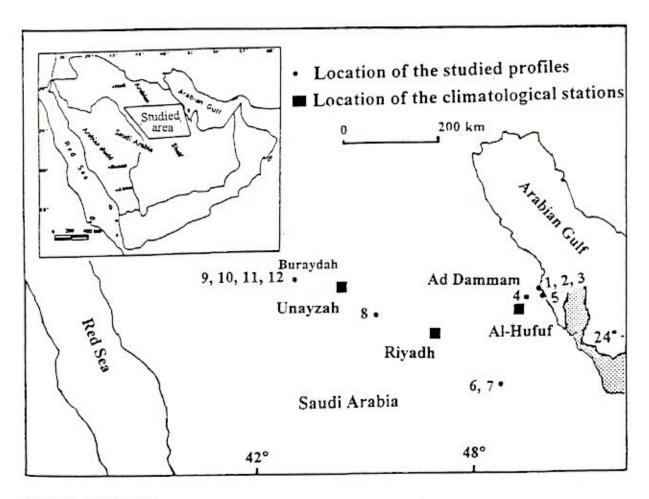


Figure 1. Location map

Abqaiq and is connected to the Arabian Gulf through sabkhat Dannan. It is one of the coastal and dry sabkhahs and lies at an elevation of 100 m above sea level (asl). The water table level was found at 45 to 60 cm below the surface. Profiles 1, 2 and 3 were taken from the sabkhah surface, the foot slope and the top of a surrounding sand dune, respectively. The dominant natural vegetation in the sabkhah surfaces are Zygophyllum qattarense, Zygophyllum Coccineum and Halopeplis Perfoliataa, while Hammada elegans Sp dominates higher on the sand dune.

Sihat sabkhah is located near the coast of the Arabian Gulf. It is considered to be a wet coastal sabkhah and lies at 20 m asl. The dominant natural vegetation is mainly extensive shrubs. Profile 4 was taken from the sabkhah surface.

Ad-Dammam sabkhah is located on the Gulf near the Dammam-Riyadh road 20 km from Dammam and is considered as one of the coastal sabkhahs. It lies at 20 m asl. Profile 5 was collected from the sabkhah surface sediments. The profile depth is limited by the presence of a very hard layer at 80 cm deep.

Yibrin sabkhah is located 7.5 km west of Yibrin. It is one of the dry inland sabkhahs and lies at 160 m asl. Many Wadis are associated with this sabkhah (e.g. Wadi Wabzah). Profiles 6 and 7 were collected from the sabkhah surfaces.

Al-Milh "Mamlahat Al-Qasab" sabkhah is located in the central region 60 km Northwest of Riyadh. It lies at an elevation of 600 m asl. The sabkhah is surrounded by the Twaiq mountains from the north and east, elevation 1022 m asl, and sand dunes of Uraq Al-Buldan from south and west, 700 m asl. This sabkhah is connected with Wadi Bu'aythiran and Wadi Wirat through a drainage system. Scattered natural vegetation such as Halopeplis perf is present. Profile 8 was collected from the sabkhah surface.

Al-Awasjiah sabkhah is located in the central region 5 km from Al-Awasjiah. It lies at 600 m asl and is surrounded by sand dunes from east Nafud ath Thuwayrat and from west Naas Sirr. These lie at 800 m asl. Scattered farms exist in the area at elevations higher than the sabkhah level. Profiles 9, 10 and 12 were taken from the sabkhah surface, while profile 11 was collected to represent the surrounding dunes.

SAMPLING AND ANALYSIS: The location of the profiles was selected to represent the surface features dominant in each of the sabkhahs (i.e. the physiographic position, wetness conditions, slope and presence or absence of natural vegetation). Profiles were dug, deep, morphologically described and then representative samples were collected from the surface salt crust when present. Various analyses were carried out as follows: calcium carbonate (CaCO₃) content was determined by

calcimeter according to US Salinity Laboratory Staff (1954). Gypsum (CaSO₄•2H₂O) content was determined according to Berigari and Al-Any (1994). Soluble salts (EC.) were assessed in the saturation paste extract using an electrical conductivity meter capable of measuring a relatively wide range of salt levels. Soluble cations and anions were determined in the saturation paste extracts, after the appropriate dilutions, according to the standard methods outlined by Page et al. (1982). Na+ and K+ were measured by flame photometry, Ca2+ and Mg2+ by titration with versenate, Cl by titration with a standard silver nitrate solution, CO32 and HCO3 by titration with HCl and SO42 by turbidimetry. Water soluble B3+ was measured in the saturation paste extract using the Azomethen-H method (Bingham, 1982). Water samples were collected from the standing water table in the auger hole of each profile followed by chemical analysis according to the standard methods (Page et al., 1982).

Mineralogical analysis of the salt crust was carried out on the collected samples by X-ray diffraction, following grinding and powder mounting of the fine materials (<2.0 mm) on aluminum holders. Selected samples were heated in an oven at 105°C for 2 h before examining by X-ray diffraction. A Philips difractometer equipped with a Cu tube was used to obtain the X-ray diffraction data. Scans were run at 2 degrees theta per minute from 6 to 50. Identification of the evaporite minerals was carried out according to Braitsch (1971), Gumuzzio et al. (1982), Whittig et al. (1982) and Doner and Lynn (1989).

Results and Discussions

The relief of sabkhah surfaces was level. The texture of the Rumadan sabkhah surface profile and the surrounding dunes was sandy. Sihat and Ad-Dammam sabkhahs had sandy clay loam and loamy sand texture throughout their profiles, respectively. Yibrin sabkhah had a loamy texture below the surface crust and a sandy loam texture in the underlying layers. The texture of Al-Malh sabkhah was clay loam in the surface layer underlaid by sandy loam and clay in the deep layers. Al-Awasjiah sabkhah had sandy to sandy loam texture in the layers. The observed variations in texture within and between the studied sabkhahs were due to the sediment sources and sedimentation pattern of each sabkhah. Salt crusts having variable thickness were present in both inland (2-15 cm) and coastal (2-3 cm) sabkhahs. They had slight to strong effervescence and similar soft to hard fine crystals. Water table levels were shallow in most of the studied profiles, particularly those representing the sabkhah surfaces. There were very hard layers close to the surface in the Silhate and Ad-Dammam sabkhahs.

Climatic data for the coastal sabkhahs were collected from the nearest meteorological station in Al-Hassa for which the mean annual precipitation was 78.9 mm. The highest monthly evaporation occurs in June and July with values of about 450 mm. The Riyadh meteorological station was the nearest station to Al-Malh inland sabkhah. The mean annual precipitation was 83.3 mm while the monthly evaporation ranged from 102.7 mm in January to 354 mm in June. The Unayzah meteorological station was the nearest to the Al-Awasjiah inland sabkhah where the mean annual precipitation was 69.5 mm. Accordingly, long periods of moisture deficit occur. The upward movement of water and accumulation of surface salt crusts was abundant throughout most of the year. Under such climatic conditions certain mineralogical species are likely to be found in the surface salt crust (Gumuzzio et al., 1982).

COASTAL SABKHAHS: Data of profiles 1, 2 and 3 show relatively low saturation percentage, SP, values reflecting the sandy texture of all layers, except the deepest layer of profile 3 (Table 1). Calcium carbonate contents were quite low, while the pH values ranged from 7.14 to 8.25. Relatively low gypsum contents were present in the different profile layers with the highest amount occuring at the surface. Variable Ec. values were detected with the highest values appearing in the sabkhah surface profile. These variations could be due to downward movement of soluble salts from the higher elevation (profile 3) and their accumulation in the lower elevation (sabkhah surface, profile 1). This was also clear from the level of salinity and its distribution within and between profiles 2 and 3. Salt crust was present only in the sabkhah surface profile reflecting the higher salinity level and the upward movements of salts and their accumulation on the surface. Sodium ions were the dominant cation in the saturation extracts, followed by Mg2+ in the higher salinity samples and Ca2+ in the relatively low salinity samples. Further, the Mg2+/Ca2+ ratios increased with an increase in salinity values. This trend may have been due to the existence of saline water rich in Mg2+ from the Gulf. Analysis of the water table samples of profile 1 (Table 2) supports this trend since the salinity level was quite high and Mg2+ was the second dominant In addition, the values of Na+/Mg2+ and Na+/Cl ratios were quite similar in both water table and Gulf water samples. Chloride ions were the dominant anions followed by SO₄² with HCO₃ being the lowest. Accordingly, the dominant soluble salts could be in the form of chlorides and sulfates of Na+, Mg2+ and or Ca2+. Water-soluble B3+ was relatively high in the sabkhah surface profile. Chemical analysis of the surface salt crust (Table 3) indicated the

dominance of chloride and sulfate salts of Na+ followed by Mg2+ and Ca2+, while K+ was relatively low. Salt distribution and compositions which resulted from the upward movements through the profile can be explained by the chemical solubility, particularly in the sabkhah surface profile (Hardie and Eugster, Precipitation of Ca2+ in the form of calcite or gypsum depletes the solution in Ca2+ ions. Next calcite-gypsum and mixed Na-Mg-(SO4) minerals may occur as saline water migrates upward through the profile (Timpson et al., 1986). Similar patterns were found by Butler (1969) who reported constant Mg2+/Ca2+ values with increasing Mg2+ concentrations for the Trucial Coast near Abu Dhabi. This suggested equilibrium conditions between brines and evaporite-carbonate minerals. Data for Sihat Sabkhah (profile 4) showed that the calcium carbonate content was quite high whereas the gypsum was relatively low. Levels of salinity and water-soluble B were particularly high in the surface layers. This trend may reflect the long term accumulation of salts in the surface layers via evaporation. Sodium is the dominant cation followed by Mg2+ and Ca2+ ions. Considerable concentration of K+ ions were also present. Chloride was the dominant anion followed by SO₄ and HCO₃. The dominance of Na⁺ and Mg²⁺ cations and Cl anions indicated that the main source of salinity was the sea water coming from the Gulf through direct seepage and wind action. This trend agreed well with the analysis of Gulf water and the water table samples (Table 2) which were closely similar in salinity levels, ion compositions and Na+/Mg2+ and Na+/Cl molar ratios. Data of profile 1, which represent the Ad-Dammam sabkhah (Table 1), show that gypsum was present in appreciable amounts in the deepest two layers of the profile whereas CaCO3 contents were quite low with no trend in distribution with depth. Salinity levels were quite high and decreased with depth. Salts were dominated by Na+, followed by Ca2+ in the salt crust, whereas Mg2+ was the second dominant ion in the other layers. Regarding the anions, Cl dominated followed by SO42- then HCO3. Therefore, the dominant salts were chlorides and sulfates of Na+ followed by Ca2+ or Mg2+ and K+.

It appears that the chemical composition of the coastal sabkhahs was affected mainly by the salt water of the Gulf either through direct seepage or through wind action which carried the saline water droplets to the sabkhah or the surrounding dunes and then to the sabkhah surface through leaching. This finding was consistent with the ion ratios Na⁺/Mg²⁺ and Na⁺/Cl⁻ of the Gulf water and the water table of the coastal sabkhahs. Further, the values of water soluble B³⁺ increased with the increase in salinity of the different layers. Therefore, the main pedo-chemical process taking place in the coastal sabkhah was the upward

TABLE 1

Profile	Depth	SP	Hd	EC				Soluble ions	ions						CaCO,	CaSO, 2H, O	B3+
No.	. 15							cmol·kg	1.87			Na ⁺ /Mg ²⁺	Na ⁺ /Cl	Mg2+/Ca2+	g-kg	g.kg.	mg-kg
		38		ds m.	Cart	Mg1+	Na+	K	HCO3.	C	SO,						
-	2-10	22.5	7.14	27.9	0.98	1.40	5.87	0.22	0.21	6.20	1.18	4.8	0.95	1.43	47	44.6	1.34
	10-25	20.4	7.73	28.1	1.09	1.73	5.45	0.21	0.24	5.59	1.49	6.3	0.97	1.59	35	12.9	0.79
	25-45	20.5	7.47	25.8	1.00	1.48	5.60	0.17	0.19	5.15	1.41	2.6	1.09	1.48	4	11.2	0.47
7	0-15	17.5	7.29	28.7	1.24	1.27	4.26	0.14	0.19	5.23	0.95	6.7	0.82	1.02	38	15.9	1.10
	15-30	19.0	7.52	8.6	0.73	0.40	1.36	0.07	0.19	1.62	0.82	8.9	0.84	0.55	35	3.0	0.39
	30-45	20.5	7.44	6.6	0.70	0.40	1.65	0.07	0.22	1.70	0.81	8.3	0.97	0.57	29	10.8	0.55
	45-60	19.5	7.68	13.8	0.77	0.50	2.37	80.0	0.13	2.47	0.87	9.5	96.0	0.65	36	7.5	0.55
3	0-15	18.3	8.25	2.9	0.40	60.0	0.19	0.02	80.0	0.28	0.31	4.2	89.0	0.20	57	1.5	90.0
	15-30	19.5	7.65	3.6	0.34	0.11	0.26	0.02	0.04	0.49	0.33	4.7	0.53	0.32	14	1.8	0.04
	30-45	19.0	7.94	6.57	0.59	0.16	0.83	90.0	90.0	0.82	0.51	10.4	10.1	0.27	39	3.0	0.28
	45-60	20.1	7.54	7.0	0.54	0.25	1.03	0.05	0.05	86.0	0.59	8.2	1.05	0.46	28	3.0	0.32
	60-110	28.5	7.47	17.3	1.38	92.0	4.71	0.13	0.33	5.70	1.68	12.4	0.83	0.55	34	11.2	0.74
4	2-7	32.5	7.17	0.091	2.57	29.30	61.90	2.23	0.94	94.30	2.97	4.2	99.0	11.40	308	124.0	5.39
	7-12	33.1	7.47	133.0	3.38	25.90	44.30	2.00	0.63	67.10	3.64	3.4	99.0	7.65	400	20.2	2.91
	12+	32.0	7.28	103.0	3.46	12.30	20.50	96.0	0.67	37.40	3.10	3.3	0.55	3.54	300	32.9	1.13
2	3-15	19.8	7.09	200.0	2.26	2.62	74.20	0.29	0.49	72.30	1.98	56.7	1.03	1.16	77	8.0	2.78
	1540	18.4	6.75	180.0	3.16	20.40	37.60	2.91	0.44	00.09	1.20	3.7	0.63	6.46	3	16.2	3.71
	40-60	19.2	6.94	157.0	2.76	13.20	39.50	1.02	0.40	53.80	1.56	0.9	0.73	4.79	34	194.0	3.45
	08.09	37.5	1631	117 0	4 58	13.10	54 00	1 18	99 0	05 29	2 70	8.3	0.80	2.87	19	129.0	2 55

TABLE 2

Sample No.	Depth	Hd	T.S.S. *				Soluble ions mmol·L·l				Na Mg2	Mg2*/Ca2*	Na*/CI
				Ca2+	Mg2+	Na.	Κ.	HCO,	CI	SO,			
A.G.	1	7.77	4.90	14.5	71.0	848	0.53	5.25	7.3	44.0	9.13	4.90	0.8
Prof.1	45+	7.28	2.90	30.0	40.3	351	0.55	5.00	385.0	50.0	8.71	1.34	0.91
Prof. 4	12+	7.76	4.84	19.0	72.8	629	0.35	5.75	705.0	48.5	2.6	3.83	8.0
Prof. 8	100+	7.72	20.30	62.0	315.0	2150	1.70	3.50	2800.0	45.0	6.83	5.08	0.7
Prof. 9	30+	06.9	36.50	11.0	1070.0	3724	1.90	11.0	5250.0	314.0	3.48	97.30	0.7
Prof. 10	50+	6.95	36.10	11.0	1023.0	3628	1.80	11.5	5110.0	284.0.	3.55	93.00	0.7
Prof. 12	150+	6.83	21.50	21.0	750.0	1959	1.30	9.25	3080.0	205.0	2.61	35.70	9.0

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movement of saline water through the profile and accumulation of salts at the sabkhah surface as a result of evaporation. At higher elevations, particularly in the surrounding dunes, downward movement of solutes was an effective process in leaching accumulated salts. Periodic flooding of the sabkhah surfaces could also disturb the chemical balance in the profile thus affecting the salt level, composition and distribution within the profiles and the crust (Butler 1969). The difference in salinity between the sabkhah surface profiles may have been due to the periodic flooding or other processes which reduced the salt levels of Rumadan (100 m asl) relative to those of Ad-Dammam (20 m asl) and Sihat (20 m asl) sabkhahs. In addition, the surrounding dunes of Rumadan sabkhah, which had very low salinity levels, constitute an important land resource for agricultural development, provided that irrigation water can be secured.

INLAND SABKHAHS: Data in Table 4 indicate that relatively high contents of gypsum were present in Yiprin sabkhah (profiles 6 and 7) with an increasing content with depth as in profile 6. Levels of soluble salts were quite high in the surface layers but decreased drastically with depth in both profiles. Soluble Na+ and Cl were the dominant ions followed by Mg2+ in the very high salinity samples and Ca2+ in the relatively low salinity counterparts. Relatively high contents of soluble K+ were present in the upper layers of profile 6. Therefore, NaCl was the main soluble salt followed by Mg2+ or Ca2+ chlorides. Sulfate anion was relatively low and may be controlled by the presence of gypsum. Ratios of Na+/Mg2+ and Na+/Cl- showed no specific distribution trend with depth, while the Mg2+/Ca2+ ratio increased with increasing salinity. The rise in the Mg²⁺/Ca²⁺ ratio can be attributed to the precipitation of Ca²⁺ ions as carbonate and sulfate minerals. On the other hand, relatively low salinity levels were present in the dunes surrounding Yibrin sabkhah (profile 7), even through it had a thick surface salt crust rich in gypsum and depleted in CaCO3. Therefore, the main process affecting the formation of the Yibrin sabkhah was the accumulation of salts and gypsum through evaporative concentration as well as accumulation of gypsum and salt-rich sediments from the surrounding wadis. Data from the Al-Milh "Mamlahat Al-Qasab" sabkhah show that CaCO3 and CaSO₄2H₂O contents were relatively high in the different layers of the profile (profile 8). The amounts of soluble salts were quite high particularly in the surface layers and were dominated by NaCl followed by Ca2+ or Mg2+ chlorides. Relatively low levels of sulfate ions were present which could be controlled by the presence of gypsum. Similarly, no specific trend was observed for Na+/Mg2+ ratios in the different layers. Data from Al-Awasjiah sabkhah showed high salinity levels and variable gypsum contents in the profiles representing the sabkhah surface, whereas contents in the surrounding dunes were very low. Relatively high quantities of CaCO3 were present in the wet sabkhahs of profiles 9 and 10. The surrounding dunes generally had sandy textures throughout the studied layers. The variation in texture of profiles and layers could be due to the lack of homogeneity amongst the parent materials from which the sabkhah surface was formed. Sodium chloride was the dominant soluble salt followed by Mg2+ or Ca2+ salts in the higher and lower salinity samples respectively. Salinity of the surrounding dunes was low with high Ca2+ concentrations relative to Na+ and Mg2+. Relatively high Na+/Mg2+ and Na+/Cl ratios were present in the dry sabkhah (profile 12) as compared with those of the wet sabkhah (profiles 9 and 10). This could be related to the existing conditions of drying which may have precipitated Mg2+ in slightly soluble forms. A similar trend was observed for the Mg2+/Ca2+ ratios, which showed relatively higher values in the wet sabkhah compared to that of the dry one. Concentrations of the total soluble salts were extremely high in the water table samples of the inland sabkhah (profiles 8,9, 10 and 12). Dominant salts were mainly Cl followed by SO₄² of Na⁺ and Mg²⁺.

Correlation coefficients (r2) between ECe values and saturation paste extract compositions were B3+ (0.17), Na+ (0.91), Cl (0.88), and Mg2+ (0.37). Values of (r2) indicated that Na+ and Cl- contributed more to EC, values than B and Mg2+. The ion ratios Na⁺/Mg²⁺, Mg²⁺/Ca²⁺and Na⁺/Cl⁻ of the water table (Table 2) were quite similar to profiles 9 and 10 while profiles 8 and 12 were different. Under the existing conditions of high salinity, evaporation will cause precipitation of slightly soluble compounds in the profile and at the surface. This process will deplete the solution in some ions and change the concentration of ions, ion ratios and chemical composition of the profile layers. The high values of the Mg2+/Ca2+ ratio of the water table samples and the profile layers of Al-Awasjiah sabkhah could have been due to the precipitation of calcite and gypsum and the depletion of the solution in Ca2+ ions. This process is consistent with the presence of relatively high gypsum and calcite contents. Further, the content of water-soluble B3+ in the studied inland sabkhahs was relatively low compared with that of the coastal sabkhahs. This was due mainly to the source of the sabkhah sediments as well as the source and composition of the water table and salts.

CHEMICAL COMPOSITION AND MINERALOGY OF SALT CRUSTS: Analysis of salt crusts samples was carried out

TABLE 3

EC Soluble ions CaCO ₃ CaSO ₄ 2H ₁ O Na ⁺ / ₂					Chen	nical an	alysis an	num pi	ralogica	dimoo n	osttons	of salt c.	Chemical analysis and mineralogical compositions of salt crust samples		
Ca ²⁺ Mg ²⁺ Na ²⁺ HCO ₃ Cl SO ₃ ²⁻ 0-2 7.30 0.39 6.0 3.6 29.4 1.4 19.8 53.3 11.1 29 47.9 16.3 0-2 7.10 1.15 17.5 7.5 94.0 1.2 38.1 169.0 36.0 35 52.5 25.1 0-30 7.10 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 0-30 7.10 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 0-10 8.20 1.5 130.0 2.1 30.5 22.0 132.0 9.0 174.0 0-10 8.20 1.3 168.0 3.0 0.5 0.1 30.5 22.0 132.0 9.0 25 9.3 14.5 0-1 3.20 1.3 10.0 5.7 <t< th=""><th>Profile No.</th><th>Depth</th><th>Hd</th><th>EC dS m-1</th><th></th><th></th><th>S</th><th>oluble io</th><th>su</th><th></th><th></th><th>CaCO,</th><th>CaSO₄·2H₂O g·kg⁻¹</th><th>Na⁺/ Mg²*</th><th>Mineralogy *.b</th></t<>	Profile No.	Depth	Hd	EC dS m-1			S	oluble io	su			CaCO,	CaSO ₄ ·2H ₂ O g·kg ⁻¹	Na ⁺ / Mg ² *	Mineralogy *.b
0-2 7.30 0.39 6.0 3.6 29.4 1.4 19.8 53.3 11.1 29 47.9 16.3 0-2 7.10 1.15 17.5 7.5 94.0 1.2 38.1 169.0 36.0 35 52.5 25.1 0-30 7.10 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 0-30 7.10 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 0-10 8.20 1.90 65.0 1.5 130.0 2.1 30.5 225.0 174.0 32.7 0-10 8.20 1.35 168.0 3.0 0.5 0.1 30.5 26.6 396.0 47 500.0 0.3 0-2 0.11 1.50 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.					Ça,	Mg2+		K	HCO,	c	SO,				
0-2 7.30 0.39 6.0 3.6 29.4 1.4 19.8 53.3 11.1 29 47.9 16.3 0.2 0.2 7.10 1.15 17.5 7.5 94.0 1.2 38.1 169.0 36.0 35 52.5 25.1 1.11 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 17.0 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 174.0 0.10 8.20 1.35 168.0 3.0 0.5 0.1 30.5 22.0 132.0 91 257.0 174.0 0.2 0.1 150 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.5 0.15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 0.3 0.3 37.0 0.7 30.5 488 55.2 57 199.0 205.0 0.5 0.5 0.5 0.5 0.7 22.5 298 29.2 32 177.0 85.5							Coa	istal Sabi	chah						
0-2 7.10 1.15 17.5 7.5 94.0 1.2 38.1 169.0 36.0 35 52.5 25.1 11.0 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 11.0 0.35 5.0 1.8 29.4 0.2 19.8 47.9 7.6 61 4.0 32.7 11.0 0.3 6.60 1.90 65.0 1.5 130.0 2.1 30.5 222.0 132.0 91 257.0 174.0 0.2 0.1 1.50 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.5 0.15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 0.3 0.3 7.20 25.0 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5 0.5 0.5 0.7 22.5 298 29.2 32 177.0 85.5	-	0-5	7.30	0.39	0.9		29.4	1.4	19.8	53.3	11.1	53	47.9	16.3	Q, H, D, B, G, E, C
0-30 7.10 0.35 5.0 1.8 29,4 0.2 19.8 47.9 7.6 61 4.0 32.7 Inland Sabkhah 0-3 6.60 1.90 65.0 1.5 130.0 2.1 30.5 222.0 132.0 91 257.0 174.0 0-10 8.20 1.35 168.0 3.0 0.5 0.1 30.5 26.6 396.0 47 500.0 0.3 0-2 0.11 1.50 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.5 0-15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 0-3 7.20 3.20 25.0 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5	4	0-2	7.10	1.15	17.5		0.40	1.2	38.1	169.0	36.0	35	52.5	25.1	Q, H, G, E, B
0-3 6.60 1.90 65.0 1.5 130.0 2.1 30.5 222.0 132.0 91 257.0 174.0 0-10 8.20 1.35 168.0 3.0 0.5 0.1 30.5 220.0 132.0 91 257.0 174.0 0-2 0.11 1.50 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.5 0-15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 0-3 7.20 3.20 25.0 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5	· v	0-30	7.10	0.35	5.0		29.4	0.2	19.8	47.9	7.6	61	4.0	32.7	Q, H, G, E, B
0-3 6.60 1.90 65.0 1.5 130.0 2.1 30.5 222.0 132.0 91 257.0 174.0 0-10 8.20 1.35 168.0 3.0 0.5 0.1 30.5 26.6 396.0 47 500.0 0.3 0.2 0.1 1.50 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.5 0-15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 0-3 7.20 3.20 25.0 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5							III	and Sabk	thah						
0-10 8.20 1.35 168.0 3.0 0.5 0.1 30.5 26.6 396.0 47 500.0 0.3 0.2 0.2 0.11 1.50 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.5 0.15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 0.3 7.20 3.20 25.0 3.0 307.0 0.7 30.5 488 55.2 57 199.0 205.0 0.5 7.09 2.20 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5	9	0-3	9.60	8.1	65.0		130.0	2.1		222.0	132.0	91	257.0	174.0	H, G, A, D, B, T, C, An
0-2 0.11 1.50 17.5 16.5 120.0 2.7 41.9 229.0 9.0 25 9.3 14.5 H 0-15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 H, 0-3 7.20 3.20 25.0 3.0 307.0 0.7 30.5 488 55.2 57 199.0 205.0 H, 0-5 7.09 2.20 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5 H,	1	0-10	8.20	1.35	168.0		0.5	0.1	30.5	26.6	396.0	47	500.0	0.3	G, B, E, D
0-15 7.10 3.60 10.0 6.0 371.0 0.9 26.7 577 31.2 27 37.0 129.0 H, 0-3 7.20 3.20 25.0 3.0 307.0 0.7 30.5 488 55.2 57 199.0 205.0 H, 0-5 7.09 2.20 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5 H,	00	0-2	0.11	1.50	17.5		120.0	2.7	41.9	229.0	0.6	25	9.3	14.5	H, B, A
0-3 7.20 3.20 25.0 3.0 307.0 0.7 30.5 488 55.2 57 199.0 205.0 H, 0-5 7.09 2.20 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5 H,	6	0-15	7.10	3.60	10.0		371.0	6.0	26.7	577	31.2	27	37.0	129.0	H, T, E, G
0-5 7.09 2.20 30.2 7.3 312.0 0.7 22.5 298 29.2 32 177.0 85.5 H _s	10	0-3	7.20	3.20	25.0		307.0	0.7	30.5	488	55.2	57	199.0	205.0	G, B,
	12	0-5	7.09	2.20	30.2		312.0	0.7	22.5	298	29.5	32	177.0	85.5	H, G, B, E

Mineralogy; Q: Quartz; H: Halite; G: Gypsum; D: Dolomite; C: Calcire; E: Epsomite; B: Bloedite; T: Thenardite; A: Anhydrite b Minerals present in sequences (descending order).

TABL

Profile No.	Depth	S %	Ħ	EC dSm-1			vă -	Soluble ions cmol·kg ⁻¹	8_			Na*/Mg2+	Na+/CI	Mg ²⁺ /Ca ²⁺	CaCO, g-kg	CaSO,2H,O	B ¹⁺ mg·kg ⁻¹
					Ça,	Mg2+	Na.	¥.	HCO.	CI	20°		CHANGE WASHINGTON				
9	3-15	27.4	7.10	214.0	09.9	20.40	102.9	4.44	0.34	118.0	0.38	10.1	0.87	3.09	105.0	242	0.75
,	15-25	24.5	7.20	199.0	2.87	8.72	78.2	2.58	0.49	94.3	1.32	17.9	0.83	3.04	0.68	486	0.46
	25-40	25.0	8.10	51.8	2.15	1.60	14.5	0.79	0.29	18.0	0.43	18.1	0.81	0.74	160.0	552	0.28
	40-50	35.0	8.20	19.7	1.75	0.84	6.31	0.30	0.40	6.5	1.87	15.0	1.02	0.48	128.0	592	0.18
7	10-30	22.1	7.65	7.2	2.06	0.28	0.91	0.03	0.23	1.2	1.12	6.5	92.0	0.14	0.69	200	0.60
00	2-10	29.4	6.50	220.0	29.0	5.50	118.5	0.04	0.10	153.0	0.40	43.1	0.78	0.19	368.0	80	0.47
,	10-25	27.5	6.74	99.5	9.63	10.50	23.9	0.93	0.52	41.3	96.0	4.6	0.58	1.09	297.0	908	0.85
	25.40	27.2	7.52	36.6	2.90	1.55	9.82	0.26	0.11	11.2	1.43	12.7	0.88	0.53	216.0	260	99.0
	40-70	29.4	7.50	38.2	3.16	1.65	10.9	0.22	0.88	12.8	1.66	13.2	0.85	0.52	201.0	285	0.35
	70.85	34.6	7.20	41.9	7.29	6.39	11.2	0.30	0.83	20.8	1.31	3.5	0.54	0.88	273.0	170	0.37
	85-100	47.2	7.02	34.7	2.93	2.56	16.0	0.31	0.42	16.9	2.34	12.5	0.95	0.87	314.0	82	0.34
0	15-30	35.8	98.9	185.0	1.36	40.20	101	2.53	0.82	137.8	8.48	5.0	0.74	29.50	367.0	104	1.50
101	3-10	33.0	7.48	165.0	7.26	48.80	67.4	3.05	0.79	113.5	10.40	2.8	0.59	6.72	168.0	387	6.12
2	10-25	33.0	7.24	168.0	1.32	51.40	67.4	2.74	1.16	118.8	7.39	2.6	0.57	38.90	254.0	312	2.46
	25-50	37.1	7.11	117.0	2.45	27.20	40.3	1.66	1.19	9.89	4.01	3.0	0.59	11.09	296.0	286	9.
=	0-50	18.5	8.35	1.6	0.20	0.07	0.13	0.02	0.07	0.1	0.16	3.7	0.93	0.35	4.9	36	0.05
	50-100	19.0	8.13	2.1	0.23	0.11	0.16	0.01	0.05	0.2	0.19	2.9	0.70	0.48	3.7	38	0.04
	100-150	18.5	7.90	398.0	0.46	0.39	0.38	0.03	0.05	9.0	0.08	2.0	0.60	0.85	6.3	31	40.0
12	5-15	22.3	6.58	145.0	2.38	10.30	43.8	1.24	0.79	51.3	3.03	8.5	0.85	4.31	19.5	23	0.54
	15-50	22.0	6.73	140.0	1.98	5.89	39.6	0.98	0.42	47.3	3.34	13.5	0.84	2.97	18.1	56	0.55
	50-70	28.7	7.05	116.0	2.35	5.27	39.9	0.84	0.60	47.4	4.08	15.2	0.84	2.24	15.6	28	0.67
	70-100	22.6	7.49	86.0	1.24	2.40	21.7	0.41	0.43	26.0	3.01	18.1	0.84	1.9	18.1	23	0.50
	100-150	23.5	7.22	106.0	2.54	08.9	29.8	0.55	0.52	35.3	3.29	8.78	0.85	2.68	15.6	41	0.46

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using sample water ratios ranging from 1:100 to 1:1000. Data are presented only for the 1:500 ratio (Table 3). Results indicated that the effect of increasing the dilution of the suspension was to reduce the EC No significant change in the ion content (g-kg-1) of samples was found as a result of dilution except for Ca2+ and sulfate ions in the high gypsum samples, e.g. at Yibrin. Soluble Na+ was the dominant cation followed by Ca2+, Mg2+ and K+ in all salt crust samples but one, namely, the Yibrin sabkhah (profile 7). Here Ca2+ dominated followed by Mg2+, Na+ and K⁺. Besides the Yibrin sabkhah profiles which were rich in sulfates, anions were dominated by Cl. Therefore, with the exception of the sabkhah of Yibrin, the dominant soluble salts in salt crust samples were mainly Cl and SO42 of Na+, Ca2+ and Mg2+. The former, however, was dominated by Ca2+ and SO4- in the surrounding dunes (profile 7) and SO₄2- of Na+ and Ca2+in the dry sabkhah surface (profile 6). X-ray diffraction patterns of the different salt crust samples (Figure 2) confirmed the presence of halite (NaCl) mineral as a major salt constituent in the coastal sabkhahs. Gypsum, epsomite (MgSO₄-7H₂O), bloedite (Na2MgSO4·4H2O) and dolomite (CaMg(CO3)2) were also present in relatively low contents (Table 3). Quartz was detected in almost all samples particularly in the coastal sabkhah sample which undoubtedly was

due to eolian action. The salt crust of Yibrin sabkhah showed different mineralogy. The dry crust contained almost entirely pure gypsum (profile 6) with the surrounding dunes being rich in gypsum and halite (profile 7). Other minerals were detected in both profiles e.g. calcite, bloedite, epsomite, dolomite and anhydrite (CaSO₄). Traces of the mineral analcime were detected in profile 6 from the characteristic peaks (d-spacing) at 0.342, 0.292 and 0.558 nm (Gude, Variations in mineralogy are undoubtedly 1981). related to the composition of the solution and sediments which appeared to be rich in gypsum and other sedimentary rocks. Surface crusts of Al-Awasjiah sabkhah were dominated by halite with minor quantities of gypsum, bloedite, thenardite (Na2SO4), epsomite and calcite minerals. It is worth mentioning that the identified minerals and in particular gypsum and the Na+ and Mg2+ sulfates were subject to seasonal changes due to temperature variation and to other prevailing climatic factors. Timpson et al. (1986) concluded from their study on the evaporite mineralogy that hydration state, mineral occurrences, and mineral solubility were seasonally dependent. Further, Driessen and Schoorl (1973) reported on the temperature stability of mirabilite and thenardite with respect to hydration Similarly, heating the samples at 105°C followed by examination using X-ray diffraction

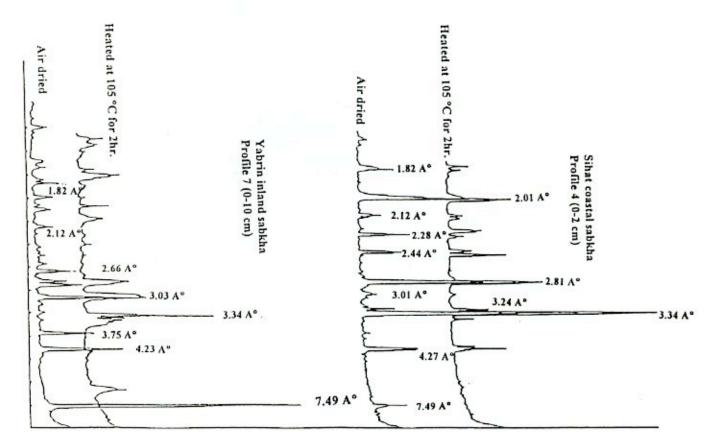


Figure 2. X-ray diffraction pattern of selected salt crust samples

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indicated the elimination of gypsum peaks and the appearance of new diagnostic peaks for bassanite (CaSO_{4*1/2}H₂O), (Figure 2). Therefore, the absence of gypsum peaks on heating confirms the conversion of gypsum crystals to bassanite in all gypsum containing samples. This mechanism might partially occur under the prevailing arid climate.

Conclusions

Very high quantities of water soluble salts were present in the sabkhah surfaces. The composition of the coastal sabkhahs was closely related to the composition of the Gulf water whereas inland sabkhahs were associated to local environmental conditions and seeps This was confirmed from the similarity in salt levels, ion ratios and the relatively high correlation coefficients between EC_e and water soluble B3+ (r2=0.8) or Mg2+(r2=0.6) in coastal sabkhahs as compared with the inland sabkhahs (r2=0.17 and 0.37) respectively. The main pedological process affecting formation of the coastal sabkhahs were the upward movement of the water tables rich in salts originating mainly from Gulf waters either directly or through wind action. Leaching, accumulation of sediments, and evaporative concentration of the solutes were the processes prevailing in the inland sabkhahs. Salt crusts were dominated by Cl and SO42 of Na+, Mg2+ and Ca2+ salts where K+ salts were present in low concentrations. X-ray diffraction patterns confirmed the presence of halite, gypsum and carbonate minerals associated with various other minerals. Epsomite and bloedite were identified in the coastal and inland playas. Specific minerals and relative abundance was quite variable between the evaporites and the characteristic conditions of each sabkhah. The major differences between the inland and coastal sabkhahs were related to the sources, composition, and depth of the water tables. This, of course, reflects the composition and mineralogy of the profile and the surface salt crusts.

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