



## Isotopic geochemistry utilization in the determination of the origin recharge of the water springs of the Algerian upper plains

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### Abstract

Chemical and isotopic properties of water springs have been studied in Aures massif, which is located South of Algerian upper plains, city of Khenchela, Northeast of Algeria. The physico-chemical and isotopic characteristics of water springs samples have been studied in order to assess the origin of groundwater and determine the factors involved in its geochemical composition. The water springs of Algerian upper plains, initially weakly mineralized, can become salty by interaction with surrounding basin rocks and successive evaporation. The salt concentrations can increase up to the precipitation of evaporites. The use of chemical and isotopic tools has shown that Water springs are of bicarbonate-calcium facies. The origin of waters mineralization is linked to water-rock interactions, by mineral dissolution and cations exchange, and shows that the waters springs are undersaturated with respect to calcite, dolomite, aragonite, gypsum, anhydrite and halite. The isotopic tool has demonstrated that the aquifer recharge is ensured by meteoric mediterranean waters, with altitudinal influence.

## 1. Introduction

Water springs represent an important source of drinking water in many countries around the world where the importance of water quality in human health has recently attracted a great deal of interest [1-3]. In the developing world, 80 % of all diseases are directly related to poor drinking water and unsanitary conditions [4,5]. These water springs are also used for irrigation and for producing electricity (hydropower plants). For all these reasons, water resources must be managed in a sustainable way. Since the significance of water springs as important water resources and valuable ecosystems is growing worldwide, these hydrological systems are receiving rapidly increasing attention from the scientific, engineering, and regulatory communities. Literature shown that groundwater plays a dominant role in this part of Algeria received various studies [6-10]. Because of the lack of permanent surface water reservoirs owing to the hard climatic conditions, it constitutes the most widely available source of fresh water. In this region, groundwater is used for domestic and agricultural purposes. The present article aims to identify the origin of the water springs recharge of the Algerian upper plains, and to distinguish the salinity origin of the groundwater masses.

## 2. Experimental details

### 2.1. Location and climate

The study area is located in the Algerian upper plains, between Tell Atlas to the North and Saharan Atlas to the South, between 35°15' and 35°45'N of latitude and between 6 and 7° of longitudes (Figure 1). The climate of the study area is semi-arid, the average annual rainfall is 470 mm and the average annual temperature is around 16 °C.

The study area is a hydrogeological entity of the steppe upper plains, surrounded by landforms having the geological aspects of the Algerian Atlas, formed by a very pleated field, fractured and karstified, having a Cretaceous frame and a very thick tertiary recoverment composed of gypseous red clay and sandstone, and leans directly and discordantly on the lower Cretaceous. The marly cretaceous, marlo-calcareous and sole compact limestone of gypseous and salifere trias recover the powerful series of Miocene sandstone that have a poor hydrogeological interest. The center of the plain is a collapse basin filled by detrital deposits, where conglomerates, gravels, fine sand and red clay have been deposited alternatively in an aquifer multilayer system of Quaternary age. The South and South-West borders of the basin are formed by the mountains of Aures (Figure 1). These latter are constituted of the karstic formations, that are the bottom of a deep aquifer having a limited extension toward the plain center and that sink slightly to the North, which would favour the flow and drainage of these waters of karstic origin toward the basin of Chemora (Figure 2) [11, 12]. Hydrogeologically, the region has two types of aquifers: the multilayer aquifers of the sedimentary basin constituted of the aquifers of Plio-Quaternary, and the aquifers of the carbonate fissured formations of Cretaceous.

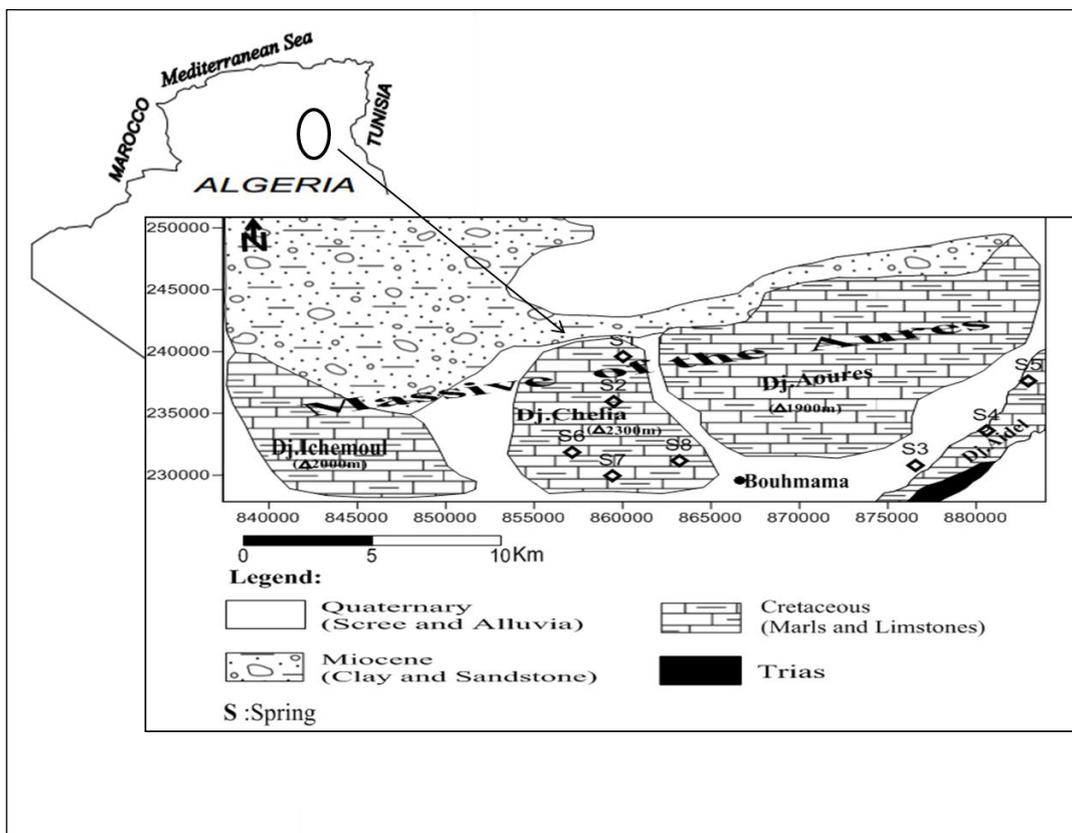


Figure 1: Location and geological features of the study area.

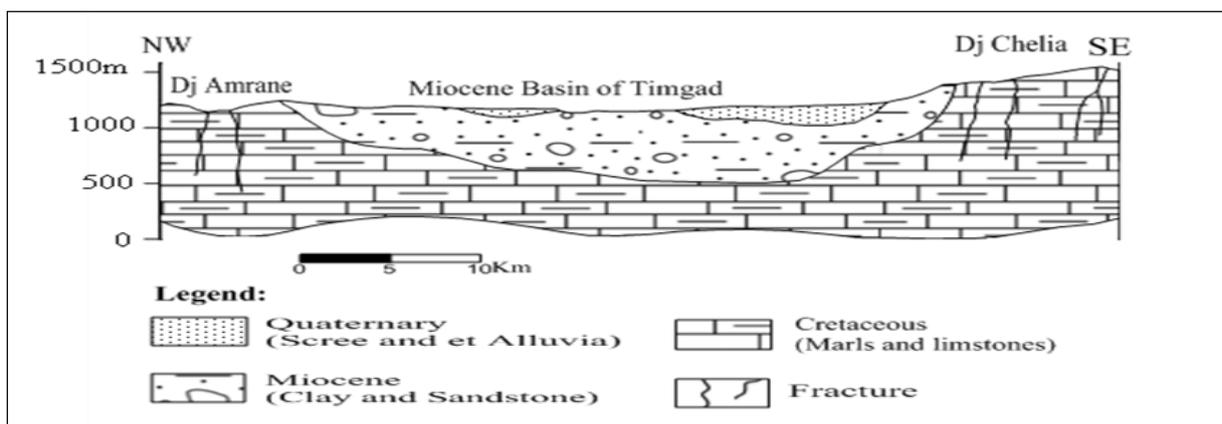


Figure 2: Geological cross section.

### 3. Material and Methods

A sampling of eight water springs capturing the Cretaceous aquifer (Figure 1) has been carried out during the rainy period in April 2015. The electrical conductivity, the temperature and the pH have been measured in the field. The results of the chemical analyses of major elements, the ratio of stable isotopes  $^2\text{H}$  and  $^{18}\text{O}$ , carried out in the laboratory of isotopic hydrology of atomic research center of Algiers (CRNA) are provided in Table 1.

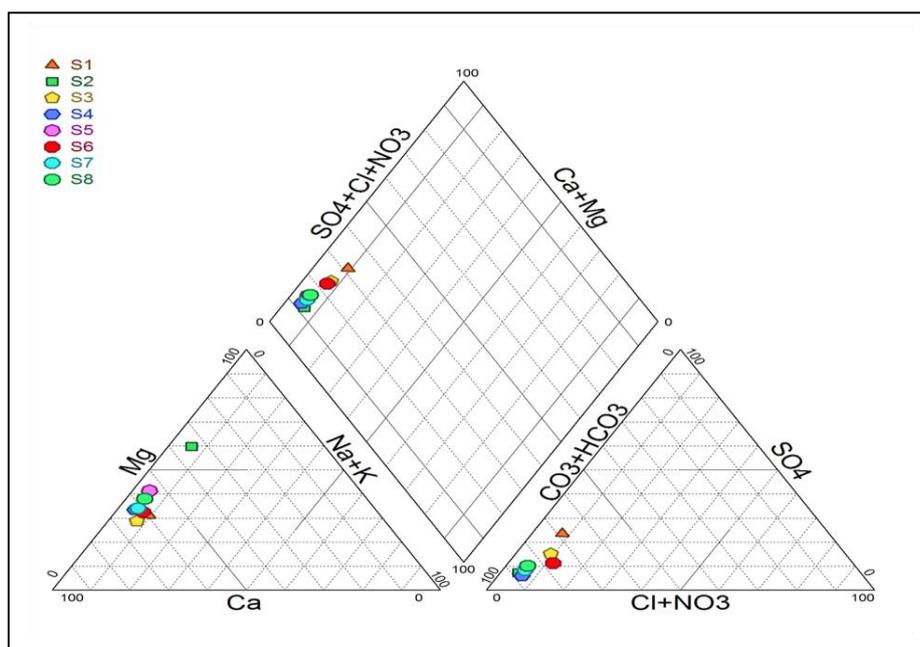
**Table 1:** Results of physico-chemical and isotopic analyses of the of the water springs of the study area

Samples	T (°C)	EC (µs/cm)	pH	Ca <sup>2+</sup> (mg/l)	Mg <sup>2+</sup> (mg/l)	Na <sup>+</sup> (mg/l)	K <sup>+</sup> (mg/l)	HCO <sub>3</sub> <sup>-</sup> (mg/l)	Cl <sup>-</sup> (mg/l)	SO <sub>4</sub> <sup>2-</sup> (mg/l)	δ <sup>18</sup> O ‰	δ <sup>2</sup> H ‰
S1	16	491	6.9	95.4	30.3	15.4	2.6	295.9	16.5	79.9	-7.77	-48.3
S2	14	394	6.8	34.2	36.3	5.6	2.3	308.1	7.7	19.8	-8.88	-55.7
S3	14	492	6.8	91.6	25.1	11.1	1.8	320.3	17.2	50.4	-8.16	-50.2
S4	14	382	6.8	83.6	27.4	5.8	2.1	318.7	10.4	16.9	-8.72	-55.3
S5	12	675	6.67	110.9	51.4	8.5	2.8	536.8	18.4	50	-8.58	-53.4
S6	14	200	7.04	32.7	10.7	3.4	1.6	114.4	5.7	13.1	-9	-55.9
S7	14	288	7	50.4	17.1	3.9	1.7	204.4	6.5	15.5	-9.53	-60.7
S8	15	511	6.8	91.8	36.8	7.5	2.5	382.8	13.9	36.6	-9.33	-58.3

### 4. Results and discussion

#### 4.1. Hydrochemistry

The water temperatures of the water springs are comprised between 12 and 16°C, with a mean value of 14°C. The groundwater pH values range from 6.67 to 7.04. The electrical conductivity varies between 200 and 675 µS.cm<sup>-1</sup>. The report of major elements contents of different sampled points of water on Piper diagram (Figure 3) shows that the cretaceous waters exhibit a bicarbonate-calcium facies because calcite is abundant in these aquifers [13, 14].



**Figure 3:** Piper diagram for the water samples

#### 4.2. Mineralization acquisition

To better distinguish the salinization process of groundwater, we have employed the cross diagrams of major elements with chloride ion (Figure 4). This latter is a conserved element, is not involved in water-rock interactions, characterizes the origin of water salinity and constitutes a mixing tracer [5]. The disposition of the different points of waters analyzed with respect to the mixing line freshwater- saltwater (rainwater- seawater) can be of great interest to identify other phenomena, which are annexed to the mixing process [15, 16]. The relationship between  $\text{Na}^+$  and  $\text{Cl}^-$  (Figure 4) reveals that all the points are below the mixing line. Given that  $\text{Na}^+$  content should balance the  $\text{Cl}^-$  content, the  $\text{Na}^+$  deficit is explained by the exchange phenomenon of ionic base between water and the aquifer, and is reflected by  $\text{Na}^+$  absorption and  $\text{Ca}^{2+}$  release. The graph  $\text{Ca}^{2+}$  versus  $\text{Cl}^-$  illustrates well this, showing that the points are often over the mixing line of freshwater- saltwater.

The relationship between  $\text{Mg}^{2+}$  and  $\text{Cl}^-$  reveals that the majority of the points are located on the mixing line with a tendency to the  $\text{Mg}^{2+}$  enrichment which might be related to the dolomite dissolution [15-18]. The graph  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  demonstrates that all the points are over the mixing line of freshwater- saltwater. The sulphate enrichment might be caused by the gypsum dissolution, the leaching of evaporites [15].

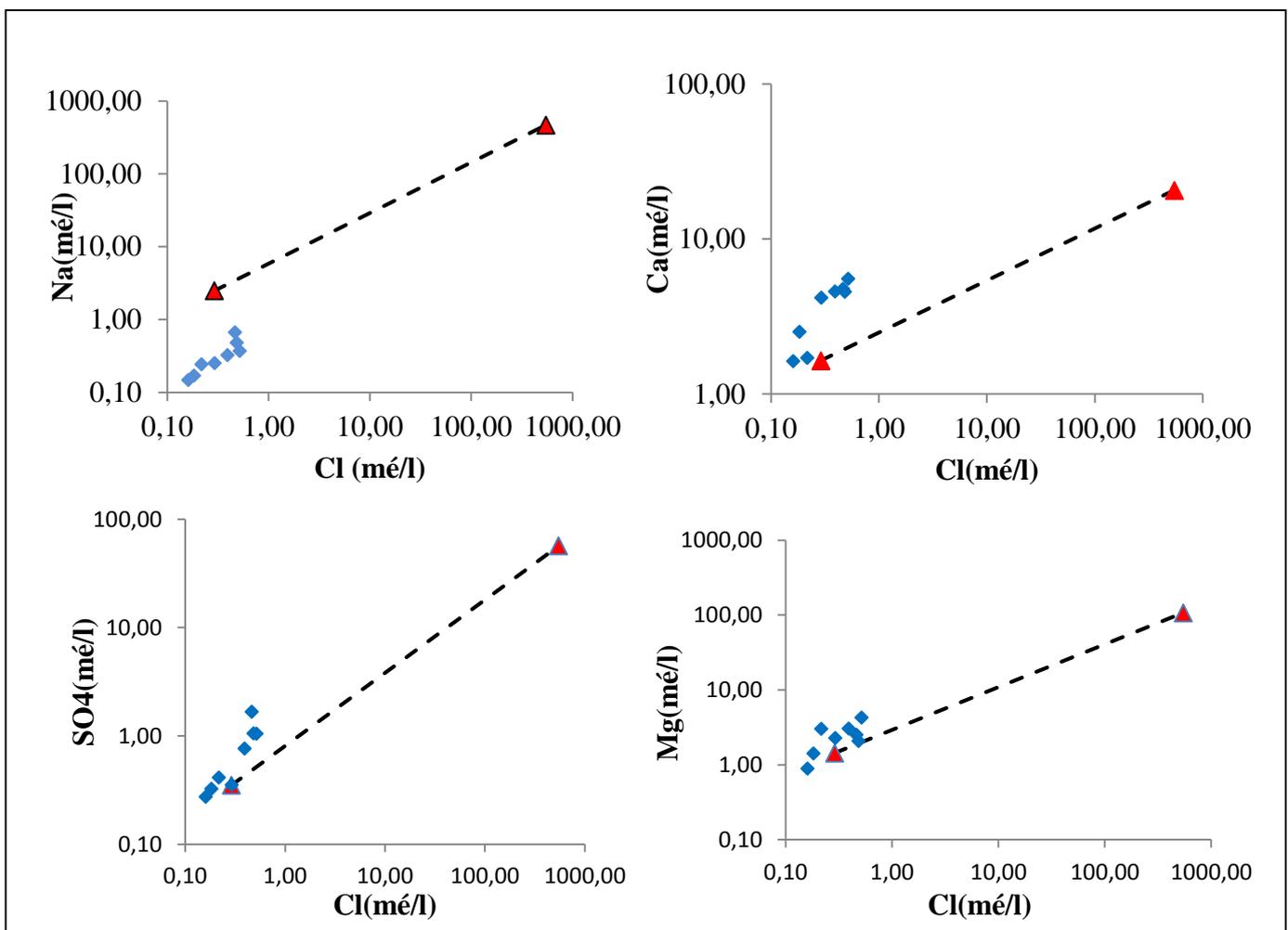


Figure 4: Relationship between major elements and chloride.

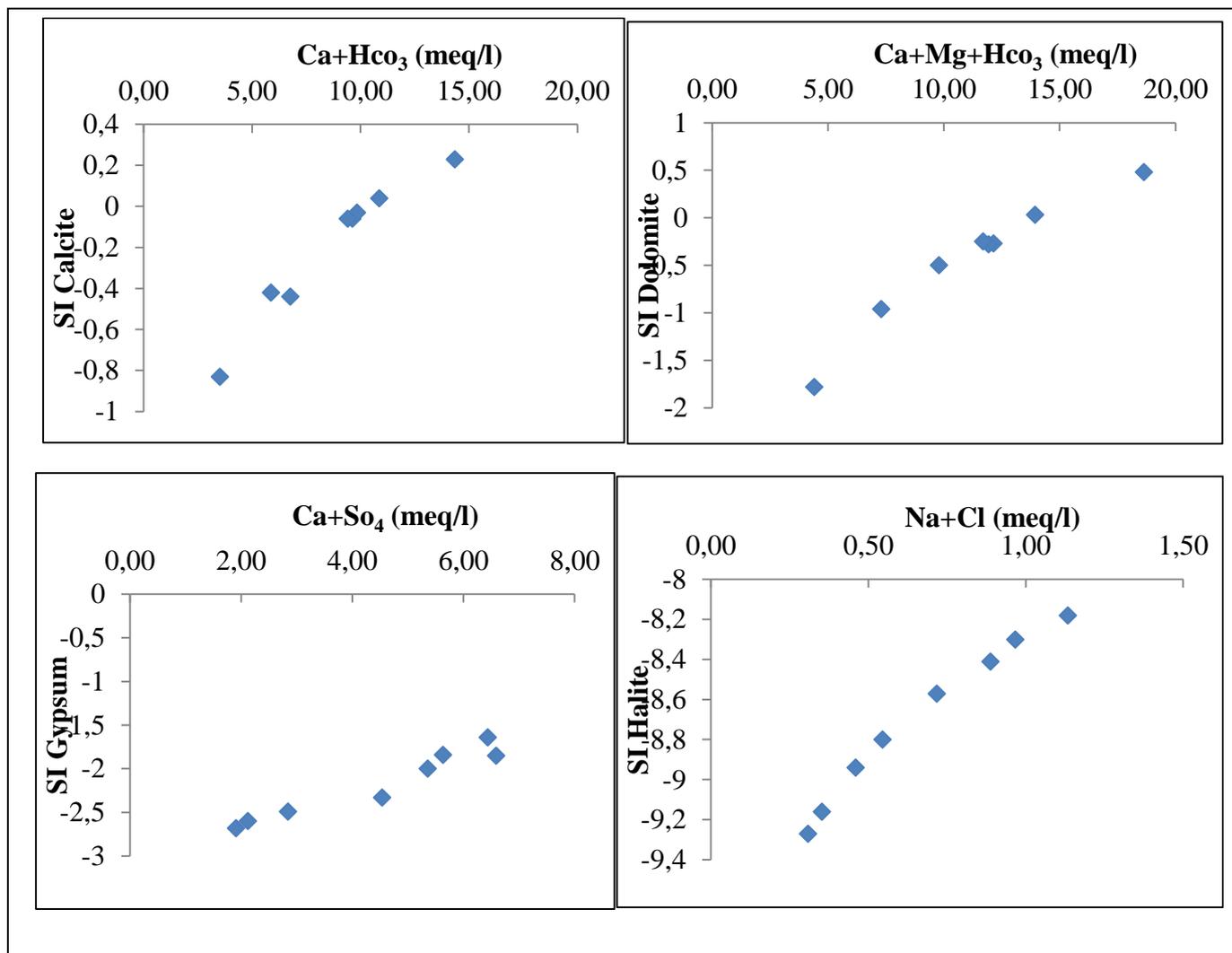
#### 4.3. Saturation index of water samples

The study of the geochemical processes which have caused the salinization of the water springs of the Algerian upper plains and their saturation with respect to the minerals is based on the ionic activity calculation. The saturation index expresses the chemical balance degree between water and the mineral in the aquifer matrix, considered as a measure of the dissolution process and / or the precipitation regarding water-rock interaction.

The use of the geochemical programme of PHREEQ, integrated in the diagram programme of the hydrochemical calculations [19] has allowed us to calculate the saturation index of calcite, dolomite, aragonite, gypsum, anhydrite and halite. The significant results of all the analyzed water samples Table 2 show that calcite saturation index of sampled water (Figure 5) in relation to concerned minerals indicates an undersaturation state of most sampled waters, and this illustrates the close correlation between calcite saturation index and concerned ions; this correlation confirms carbonate minerals dissolution [13, 14].

**Table 2:** Calculated values of the saturations indexes

samples	Anhydrite	Aragonite	Calcite	Dolomite	Gypsum	Halite
S1	-1.86	-0.31	-0.16	-0,48	-1.64	-8.18
S2	-2.82	-0.79	-0.64	-0,91	-2.6	-8.94
S3	-2.06	-0.38	-0.23	-0,68	-1.84	-8.3
S4	-2.55	-0.4	-0.26	-0,65	-2.33	-8.8
S5	-2.07	-0.25	-0.1	-0,19	-1.85	-8.41
S6	-2.9	-0.93	-0.79	-1,71	-2.68	-9.26
S7	-2.71	-0.57	-0.43	-0,97	-2.49	-9.16
S8	-2.22	-0.31	-0.16	-0,37	-2	-8.57



**Figure 5:** Plots of  $(Ca^{+}+Hco_{3}^{-})$  versus SI of Calcite,  $(Ca^{+}+SO_{4}^{-})$  versus SI of gypsum,  $(Ca^{+}+Mg^{+}+Hco_{3}^{-})$  versus SI of Dolomite and  $(Na^{+}+Cl^{-})$  versus SI of Halite.

Dolomite saturation index of most samples is negative (figure 5), which also confirms the dissolution hypothesis of these minerals in water. Gypsum dissolution (figure 5) is confirmed through the proportional relationship between water saturation index with respect to this mineral and the sum of ions, here of probable dissolution. Sampled water is undersaturated with respect to halite, indicating the dissolution possibility of this mineral. This probable dissolution is corroborated by the close correlation between the saturation index of this mineral and ions  $\text{Na}^{++} \text{Cl}^-$  (figure 5).

#### 4.4. Isotopic hydrology

Measures of stable isotopes contents (oxygen 18 and deuterium) have been carried out on 8 water samples (Figure 1). The stable isotopes contents have been reported on diagram  $\delta^2\text{H}=f(\delta^{18}\text{O})$  (Figure 6), where also appear:

- The Global Meteoric Water Line (GMWL) of equation  $\delta^2\text{H}=8 \delta^{18}\text{O}+10$  [10].
- The Western Mediterranean Meteoric Water Line (WMMWL), of equations  $\delta^2\text{H}=8 \delta^{18}\text{O}+13.7$  [21].

This graphic representation (Figure 6) shows that the water springs present an impoverished isotopic signature [22], the mean weighted  $\delta^{18}\text{O}$  value is  $-8.75 \text{‰}$  vs-SMOW. According to the diagram  $\delta^{18}\text{O}/\delta^2\text{H}$  (Figure 6), the water springs are located near the WMMWL; this indicates that the aquifer recharge of Cretaceous has been made from rainwaters of mediterranean origin in high altitudes [23, 24]. The impoverished feature of the water springs signifies that the isotopic composition of the precipitations evolves with the altitudes and becomes more and more impoverished in  $\delta^{18}\text{O}$  and  $\delta^2\text{H}$  when it increases (Figure 7) [26,22]. The effect of the altitudes is in good agreement with the Aures massif [15], whose relief is the most marked, and culminates at 2300 m [22, 23]. On average, the waters of these springs indicate a deuterium excess of 15 ‰, which is nearer to 13.7 ‰, which characterizes the mediterranean originated waters than 10 ‰ characterizing the oceanic originated waters [25, 26].

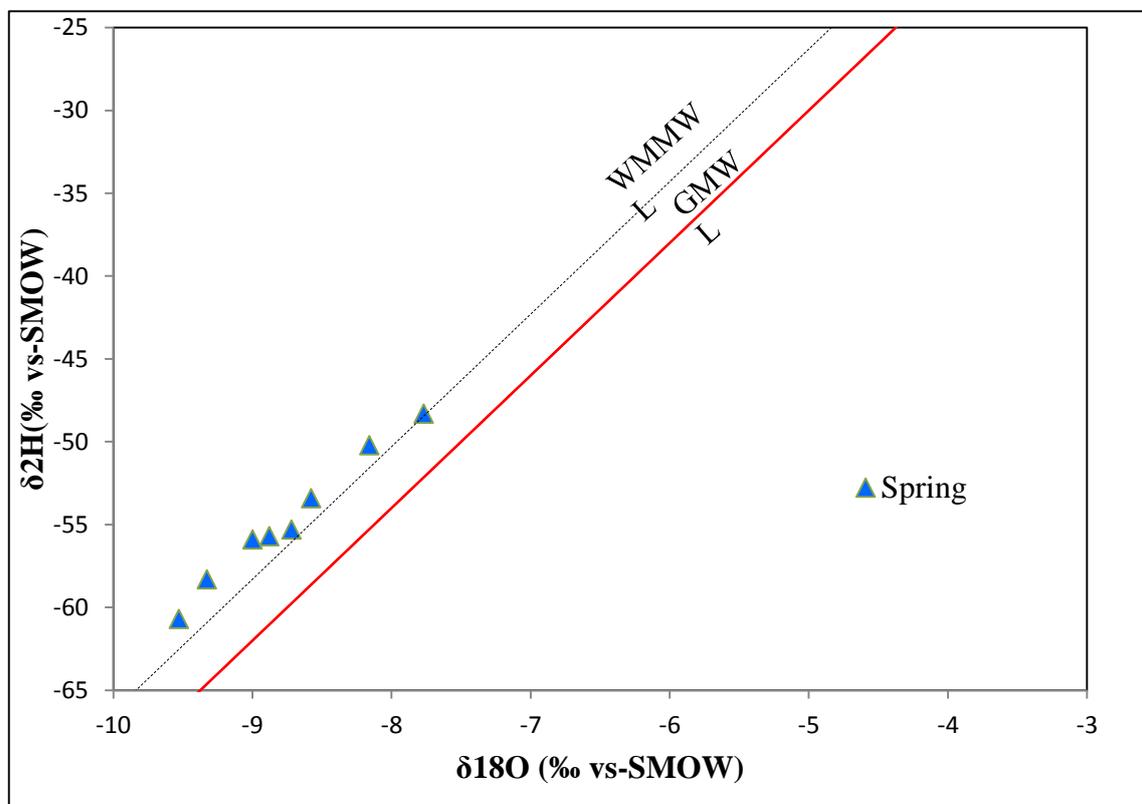
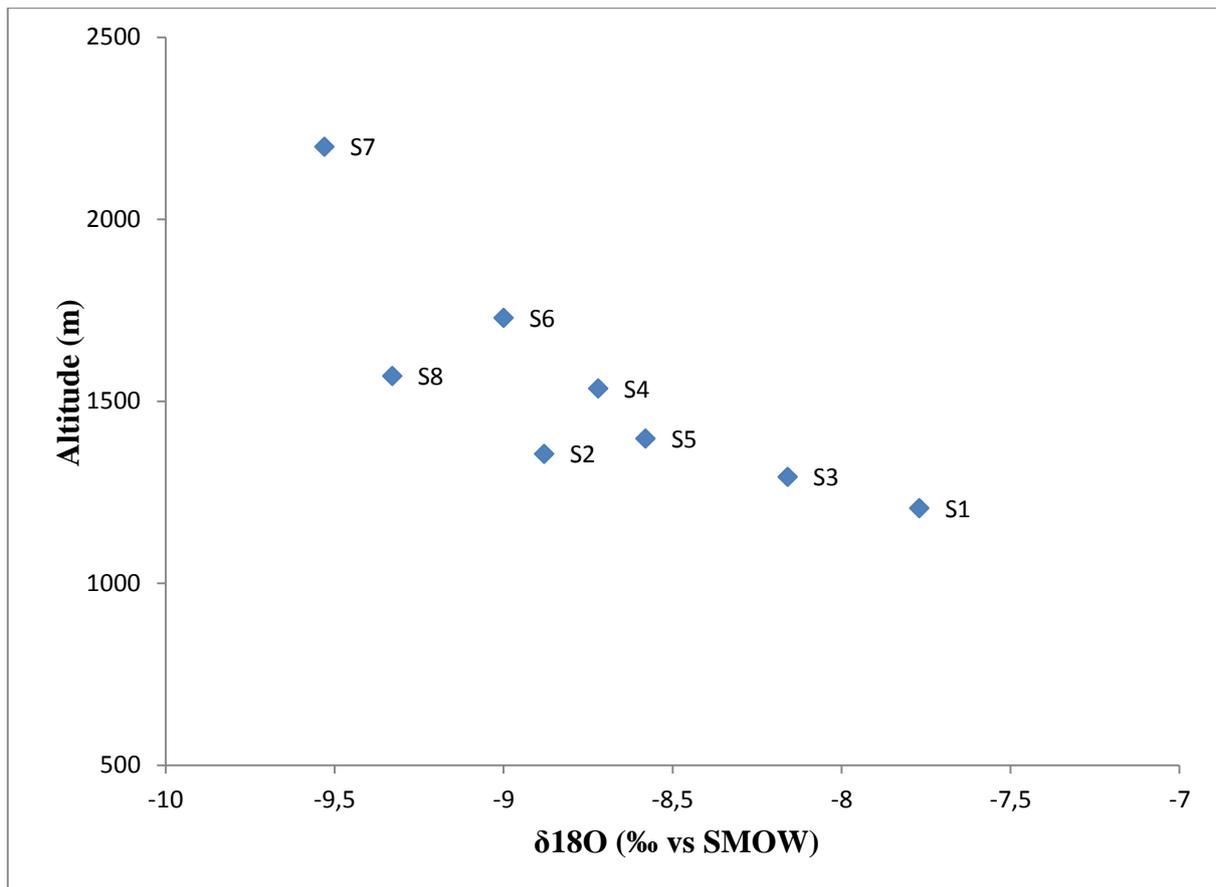


Figure 6:  $\delta^2\text{H}$  vs  $\delta^{18}\text{O}$  relationship for groundwater of the study area.



**Figure 7:** Altitude and  $\delta^{18}\text{O}$  relationship of water samples

## Conclusions

The water springs of Algerian upper plains are not yet enough mineralized, and can become salty by water-rock interaction. Water springs are of bicarbonate-calcium facies.

The use of chemical tool has shown that the origin of waters mineralization is linked to water-rock interactions by mineral dissolution and cation exchange, and shows that the water springs are undersaturated with respect to calcite, dolomite, aragonite, gypsum, anhydrite and halite. The isotopic tool utilization has shown that the origin recharge of the cretaceous aquifer is ensured by rainwaters of mediterranean origin. This is confirmed by a deuterium excess value of 15‰. The impoverished feature of the waters springs in oxygen 18 is attributed to the altitude effect.

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