

Article

Identifying karst aquifers recharge area using environmental isotopes: a case study in Central Italy

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Abstract: Water resources management is one of the most important challenges worldwide because water represents a vital resource for sustaining life and environment. In the aim of sustainable groundwater management, the identification of aquifer recharge areas is a useful tool for water resources protection. In a well-developed karst aquifer, environmental isotopes provide support for identifying aquifer recharge areas, residence time and interconnections between aquifer systems. This study deals the use of environmental isotopes to identify the main recharge area of a karst aquifer in the Upper Valley of Aniene River (Central Italy). The analysis of $^{18}\text{O}/^{16}\text{O}$ and $^2\text{H}/\text{H}$ values and their spatial distribution in the aquifer, make it possible to trace back groundwater recharge areas based on average isotope elevations. The Inverse Hydrogeological Balance Method was used to validate spring recharge elevations obtained by the use of stable isotopes. Areas impacted by direct and rapid rainfall recharge into the study area were delineated, showing groundwater flowpaths from the boundaries to the core of the aquifer. The results of this study demonstrate the contribution that spatial and temporal isotope changes can provide to the identification of groundwater flowpaths in a karst basin, taking into account the hydrogeological setting.

Keywords: recharge area; karst aquifers; stable isotope, environmental tracer

1. Introduction

Karst aquifers protection is, worldwide, the most important environmental factor for water sustainable management, as groundwater coming from karst aquifers is the largest source of freshwater for human supply.

Due to the impacts of climate change and the increasing of anthropogenic activities, groundwater exploitation in karst aquifers requires special strategies to prevent their quality and quantity depletion and to support water resources management [1,2]. Karst aquifers are characterized by spatial and temporal complexity of karst flowpaths, caused by high heterogeneity of the rock matrix, large voids, and high flow velocities [3]. Water flow velocities into a well-developed karst system are extremely fast and contaminants can quickly reach the saturated zone, making these hydrogeological systems particularly vulnerable to contamination. This system may receive localized inputs from sinking surface streams and as storm runoff through sinkholes, dolines and karst features in general. For this reason, in a karst setting, the identification of these fast recharge areas is an important tool in order to protect groundwater resources [4].

In hydrogeological studies of karst aquifers, the application of tracers can be considered as an advanced method that allows for an integrative investigation, in the aim of an effective management of water resources [5]. Environmental and artificial tracers are used more and more often in hydrogeology to investigate water flows inside the hydrological cycle, from infiltration, influenced by chemical and isotopic precipitations, to discharge due to water-rock interactions and geochemical processes [6,7,8,9,10,11,12]. At present, there are more than 1000 known isotopes of about 92 chemical elements. Most of these isotopes are either stable or unstable [5]. Naturally occurring stable isotope of water have been used extensively in hydrological investigations over the past few decades [13,14,15]. In particular, stable isotopes of hydrogen and oxygen ($^2\text{H}/\text{H}$ and $^{18}\text{O}/^{16}\text{O}$) are recognizable

tracers that are related to the water cycle and have been widely used in hydrology and water resources for decades [16]. These isotopes are often applied in hydrogeological investigations to study precipitations, groundwater recharge, groundwater-surface water interactions, delineation of flow systems and quantification of mass-balance relationships [13]. The spatial and temporal variations of stable hydrogen and oxygen isotopes in precipitations are affected by the moisture sources, elevation, temperature, rainfall amount, latitude, and distance from oceans [17-18]. The seasonal cycles of stable isotope tracers in water can be used to characterize the mean transit time of catchments, which is the average travel time for water parcels to enter as precipitation and leave as streamflow within catchments, but recent studies showed that most of these calculations have errors of several hundred percent because of aggregation bias [18-19]. Spatial and temporal variations in stable isotopes of rainfalls are related to the isotopic fractionation process, accompanying evaporation from the ocean and condensation during the atmospheric transport of water vapour [20-21]. Other factors that influence the water isotopic content, several of which are closely related to temperature, are the following: atmospheric moisture, rainfall quantity, latitude, elevation, continental and seasonal effects [17,20,21,22]. In hydrology, fractionation of $\delta^{18}\text{O}$ and $\delta^2\text{H}$ is driven by kinetic processes during evaporation and condensation [20]. During evaporation processes of oceanic water, a sequence of isotope fractionations involves variations in the precipitation isotopic composition, which are reflected in continental meteoric water. Since this fractionation process is based on the equilibrium of the isotopes ratio between vapour and liquid state of water, a specific relationship represents the distributions of isotope values of oxygen and hydrogen in rainfall [5,13]. Light variations of isotopic concentrations are usually measured by mass spectrometry. In general, the isotopic abundance ratios are expressed as parts per mil of their deviations, as given by (Vienna Standard Mean Ocean Water, VSMOW) Eq. (1):

$$\delta = \frac{(R_{\text{sample}} - R_{\text{SMOW}})}{R_{\text{SMOW}}} \cdot 10^3 \quad (1)$$

where $R = {}^2\text{H}/\text{H}$ for hydrogen and ${}^{18}\text{O}/{}^{16}\text{O}$ for oxygen, the ratio of the heavy to light isotope [23], due to the significant geographical and temporal variations in the isotopic composition of precipitation [20,24].

The application of stable isotopes ratios of hydrogen and oxygen of groundwater can provide information about springs recharge elevation, only in terms of relative values. Only the comparison between the oxygen and hydrogen isotopic compositions of precipitation and groundwater allows the evaluation of recharge elevation values and consequently the identification of aquifer recharge areas. Thus, in this paper, it has been applied a combined approach, based on the application of the Inverse Hydrogeological Balance Method [25] and the natural occurring of oxygen and hydrogen stables, to identify the boundaries of the main infiltration areas of a karst aquifer in Central Italy.

2. Geological and Hydrogeological Setting

The Simbruini Mountains karst aquifer is located in southern Latium Region (Central Italy), 60 km southeast of Rome and 65 km inland from the Tyrrhenian Sea, in the Upper Valley of the Aniene River, within the Simbruini Mountains Regional Park. Karst springs are numerous along the first part of the Aniene and are the most important water resources in the southeast part of Latium Region, supplying drinking water to the city of Rome. In this study, nine springs supplying the Simbrivio Aqueduct (Table 1), in the Simbruini Mountains karst aquifer, based on data availability, were examined. The hydrogeological basin of these karst springs, of about 15 km², is shown in Figure 1.

The lithological sequence outcropping in the Upper Valley of the Aniene River includes the North-West part of the Lower-Middle Miocene Latium-Abruzzi Carbonate Platform [26,27,28,29].

The stratigraphic succession of dolomite, dolomitic limestone and limestone is distributed homogeneously from North to South and from East to West in the study area [30]. The Triassic series outcrops among Filettino, Aniene River Springs and Faito Plateau. Dolomite is the dominant lithofacies, characterized by white and grey crystalline dolomite, with some breccia levels. Over this geological formation, limestones and dolomites, of Upper Cretaceous age are present, and their

immersion is concordant with the Triassic dolomite [29,30]. In fact, the Aniene basin is composed almost entirely of bare Mesozoic, highly fractured, karstified carbonate rocks of the central Apennine range [31]. This area is mostly made of highly permeable Cretaceous carbonate rocks, deeply fractured and mostly soluble. The base of the stratigraphic series is made of Upper Cretaceous carbonates, represented by the alternation of granular limestone and dolomites layers (Figure 1).

Table 1. Sampling location of the study area.

| Sample Codes | Spring | Lng (m) | Lat (m) | Elevation (m a.s.l.) |
|--------------|-------------------|---------|---------|----------------------|
| GW1 | Cardellina Alta | 4645069 | 355391 | 1057 |
| GW2 | Cardellina Media | 4644978 | 355274 | 989 |
| GW3 | Cardellina Bassa | 4644703 | 355176 | 939 |
| GW4 | Cesa degli Angeli | 4644606 | 355359 | 940 |
| GW5 | Cornetto | 4645520 | 352959 | 945 |
| GW6 | Carpinetto | 4646065 | 353431 | 960 |
| GW7 | Pantano Alta | 4645896 | 354187 | 952 |
| GW8 | Pantano Bassa | 4645547 | 354012 | 901 |
| GW9 | Pantano Presa | 4645714 | 353999 | 830 |

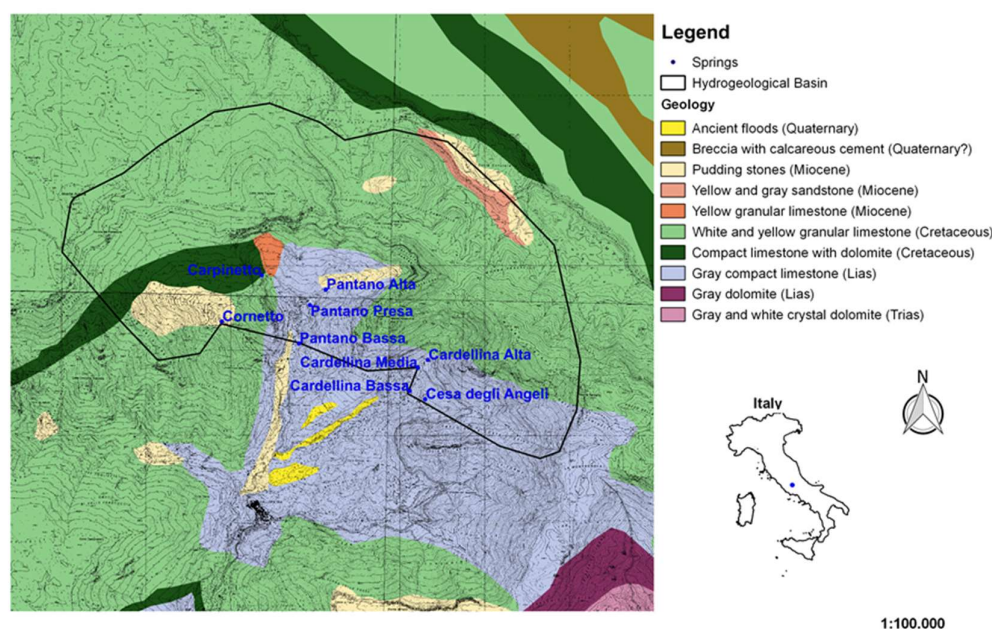


Figure 1. Geological map of the study area and location of karst basin.

Above these ones, Quaternary fluvial and alluvial deposits, conglomerate, Miocene clay and shale lie [31,32]. The carbonate deposits lying that forms the structure of the hills is typically monoclinical with immersion of the strata to N-NE and inclinations of approximately 40-45° [32,33]. Due to chemical weathering, distinctive surface and underground karst formations are developed in this area at small and large scales. The surface karst activity led to the formation of a typical karst landscape with rutted fields, sinkholes and red soils, while the underground activity has given rise to cavities, ponor and cave systems [29,30]. The karst surface is very permeable and enables the rapid infiltration of rainfall into the underground system, where the carbonate dissolution makes cavities [29,30]. Dissolution conduits strongly influence groundwater flow and evolve into complex networks, often crossing several kilometers throughout the limestone matrix [34,35,36]. The basin is characterized by both point and diffuse recharge. Recharge to the groundwater system is primarily dispersed over the basin from precipitation and secondarily concentrated at sinkholes and losing

streams. The springs discharge responses immediately to decrease and increase in precipitation especially after heavy rainfalls (from October to April).

3. Materials and Methods

Groundwater samples were collected during the wet season (June 1997) and the dry season (October 2002) to investigate seasonal variations. Eighteen water samples coming from nine of karst springs across the study area were taken. Analysis of the stable isotopic composition ($\delta^2\text{H}$ and $\delta^{18}\text{O}$) was undertaken using an Isotope Ratio Mass Spectrometer Instrumentation (IRMS) of the Isotope Geochemistry Laboratory of the University of Parma (Italy). The results (Table 2) are expressed as per mill deviations from the internationally accepted standard V-SMOW [14]. The analytical error is $\pm 1\text{‰}$ for $\delta^{18}\text{O}$ and $\pm 2\text{‰}$ for $\delta^2\text{H}$.

Table 2. Isotope composition of spring water samples collected in October 2002 and in June 1997.

| Sample Codes | June 1997 | | | October 2002 | | |
|--------------|-----------------------|--------------------|---------------------|-----------------------|--------------------|---------------------|
| | $\delta^{18}\text{O}$ | $\delta^2\text{H}$ | d_{excess} | $\delta^{18}\text{O}$ | $\delta^2\text{H}$ | d_{excess} |
| | (‰VSMOW) | (‰VSMOW) | (‰) | (‰VSMOW) | (‰VSMOW) | (‰) |
| GW1 | -8.44 | -56.89 | 10.63 | -8.67 | -50.14 | 19.22 |
| GW2 | -8.96 | -62.55 | 9.13 | -8.74 | -52.71 | 17.21 |
| GW3 | -8.60 | -59.38 | 9.42 | -8.82 | -51.87 | 18.69 |
| GW4 | -9.10 | -61.95 | 10.85 | -9.12 | -52.24 | 20.72 |
| GW5 | -8.20 | -56.10 | 9.50 | -8.38 | -47.55 | 19.49 |
| GW6 | -9.11 | -62.00 | 10.88 | -8.74 | -49.91 | 20.01 |
| GW7 | -8.63 | -59.60 | 9.44 | -8.84 | -51.56 | 19.16 |
| GW8 | -8.60 | -59.56 | 8.93 | -8.90 | -50.30 | 19.72 |
| GW9 | -8.66 | -60.35 | 9.24 | -8.82 | -50.84 | 20.90 |

4. Results and Discussions

4.1. Oxygen And Hydrogen Isotope Composition

Groundwater samples of June 1997 have $\delta^{18}\text{O}$ contents inside a range from a minimum of -8.2‰ VSMOW (GW5) and a maximum of -9.11‰ VSMOW (GW6), and with an average of -8.70‰ VSMOW. For ^2H , the average value is -59.82‰ VSMOW with a maximum of -62.55‰ VSMOW (GW2) and a minimum of -56.10‰ VSMOW (GW5) (Table 2). In October 2002 samples, the $\delta^{18}\text{O}$ values vary from -9.12‰ VSMOW (GW4) to -8.38‰ VSMOW (GW5) with an average of -8.78‰ VSMOW, and $\delta^2\text{H}$ values range from -52.71‰ VSMOW (GW2) to -47.55‰ VSMOW (GW5) (Table 2). These changes, as above mentioned, are controlled by hydrological parameters, including temperature, precipitation, altitude, latitude and seasonal effects [16,17,20,21,22]. A seasonal oscillation of stable isotope ratios is often observed as a result of temperature [5].

In order to study the overall stable isotopic characteristics of the spring samples, the isotope concentration values for June 1997 and October 2002 were plotted together with the Global Meteoric Water Line (GMWL) [17,23], the Mediterranean Meteoric Water Line (MMWL) [35] and the Central Italy Meteoric Water Line (CIMWL) [38] (Figure 2).

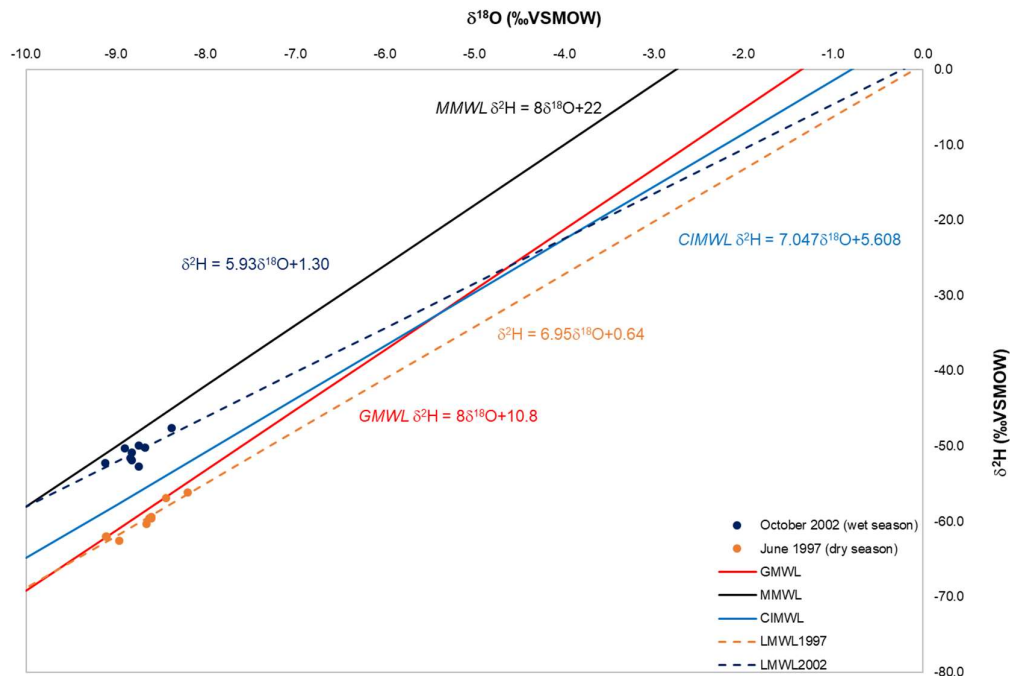


Figure 2. $\delta^{18}\text{O}$ versus $\delta^2\text{H}$, with [GLMM]: Global Meteoric Water Line [17,23]; [MLML]: Mediterranean Meteoric Water Line [37], [CILML]: Central Italy Meteoric Water Line [38].

Groundwater samples of June 1997 are located along the Global Meteoric Water Line (GMWL) of Craig (1961) revised by Rozanski (1993) ($\delta^2\text{H}\text{‰} = 8\delta^{18}\text{O}\text{‰} + 10.8$) indicating that rainfall is the primary source of groundwater. The plot of $\delta^2\text{H}$ values versus $\delta^{18}\text{O}$ for spring samples of October 2002 shows a distribution, which tends to converge with the MMWL ($\delta^2\text{H}\text{‰} = 8\delta^{18}\text{O}\text{‰} + 22$) [37] showing more $\delta^{18}\text{O}$ enrichment in comparison with $\delta^2\text{H}$ and suggesting input from local rainfall belonging to weather fronts, coming from the Mediterranean Sea. The regression lines of the samples have different slopes, which either means evaporation effect or water-rock interaction [23]. This difference is related to samples collected in two different seasons, June and October. In wet season (2002 samples), the hydrogen and oxygen isotopes showed a depleted composition compared to ones of dry season (1997 samples). The amplitude of these isotopic variations is directly related to the changing of the seasonal temperature. The distribution of hydrogen and oxygen isotopic composition in groundwater samples of Figure 2 provides an initial understanding of the origin of groundwater in the study area. The examination of the relationship $^2\text{H}/^{18}\text{O}$ shows that the values of June 1997 and October 2002 samples measured align right equations (2) and (3), respectively.

$$\delta^2\text{H}\text{‰} = 6.95\delta^{18}\text{O}\text{‰} + 0.64 \quad (2)$$

$$\delta^2\text{H}\text{‰} = 5.93\delta^{18}\text{O}\text{‰} + 1.30 \quad (3)$$

Several processes are responsible of water samples deviation from local meteoric water line. Evaporation from surface bodies is a nonequilibrium process, that enriches the residual water body such than the $\delta^2\text{H}/\delta^{18}\text{O}$ slope is less than 8, and often between 3 and 6 [13]. The slope is a function of humidity, salt concentration, and other factors [21]. In both, the slope is low (<8) as that determined by the GMWL, defined by Craig (1961) the equation $\delta^2\text{H}\text{‰} = 8\delta^{18}\text{O}\text{‰} + 10$, which indicates a meteoric groundwater's origin. So, it appears that this slope shows that waters are affected by slight phenomenon of evaporation, before infiltration [16,23]. The 2002 evaporation line (3) has a slope of 5.93, which is less than the GMWL slope of 8.17 [17], less than the MMWL slope of 8 [37], less than the CILML gradient of 7.047 [38] and also less than the 1997 evaporation line (2) gradient of 6.95.

The ^2H excess value is a function of several factors, such as the temperature, humidity, and the isotopic characteristics of the environment water vapor and the evaporating water. The value of d may differ significantly from area to area and over geologic time. Gat and Carmi (1970) report value of d greater than +24‰ for the Mediterranean region. The d -excess factor has been shown to be a

diagnostic tool for measuring the contribution of evaporated moisture to the downwind atmosphere. High d -excess values generally indicate that more evaporated moisture has been added to the atmosphere [39,40], and low values of it are associated with samples fractionated by evaporation. Previous studies showed that the western part of the Mediterranean basin has a d -excess of +14‰, whereas the eastern part shows an excess of +22‰ and this variation reflects a mixture between the Mediterranean and the Atlantic air masses [41]. These high values are probably related to a strong isotopic kinetics occurred in the summer, during evaporation process above the Mediterranean Sea and due to the low relative humidity of the atmosphere. Groundwater samples in the study area present d -excess values in the range of +8.93 and 10.88‰ in June 1997 and of +17.21 and +20.90‰ in October 2002, with an average of 9.78 and 19.46‰ respectively (Table 2). These values include the range from 10 for global precipitation to 22 for the eastern Mediterranean area indicating the significance of the Mediterranean as a moisture source for Italy [15,41]. Most of the spring samples with high deuterium excess values (October 2002) suggest that the precipitation in the groundwater comes from the Mediterranean sector. On the contrary, groundwater sampled in June 1997 show low d -excess values (ranges from 8.93 to 20.90‰). Based on that isotopic diagram of Figure 3, the first meaningful assessments were made regarding which springs were fed from the highest elevation areas and which ones from the lowest ones. First, GW4 water samples are the poorest in heavy isotopes, suggesting that Cesa degli Angeli spring has a higher infiltration elevation than all the other ones (Figure 3). The plot of $\delta^2\text{H}$ values versus $\delta^{18}\text{O}$ for water samples from the Pantano group (GW7, GW8 and GW9) shows a similar isotopic composition, suggesting the same recharge area. The same approach can be applied to GW1, GW2 and GW3 springs, whose isotopic values are depicted as a group. Lastly, water samples coming from Carpinetto spring (GW6) show different isotopic behavior referring to the sampling season. As a matter of fact, samples collected in June seem to be depleted in heavy isotopes, while this fact is not clear in ones taken in October. These different behaviors could be related to the different flow paths of groundwater coming out from Carpinetto spring. GW06 sample collected in June has been infiltrated in winter, coming from ocean reservoir. In fact, the related sample lays down along the GMWL, showing a long residence time in the aquifer. On the other hand, the sample collected in October is rich in heavy isotope, because it is referred to groundwater, infiltrated in summer, with short residence time.

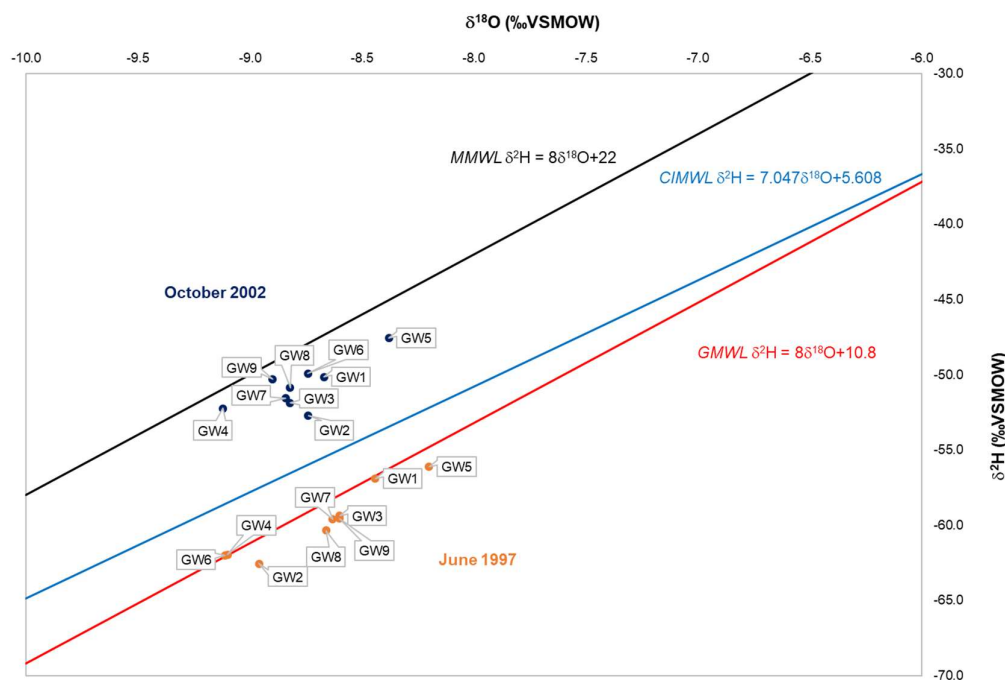


Figure 3. Plot of $\delta^2\text{H}$ vs. $\delta^{18}\text{O}$ for spring water samples collected in June 1997 and October 2002.

GW5 water samples show the heaviest content in heavy isotopes in both seasons, highlighting that the Cornetto spring is indeed fed at a lower elevation than all the other ones. However, it is impossible to define the groundwater infiltration elevations based on the samples positions on the isotopic diagram, without having precipitation markers as reference.

In order to better identify the above-mentioned recharge periods, average monthly precipitation for the twenty-year period 1951–1999 was calculated for the Vallepia pluviometric station. The evolution of the rainfalls and the outflow diagram of the Carpinetto spring (GW6) is reported in Figure 4. The other springs, which have been analyzed in this study, show similar behavior.

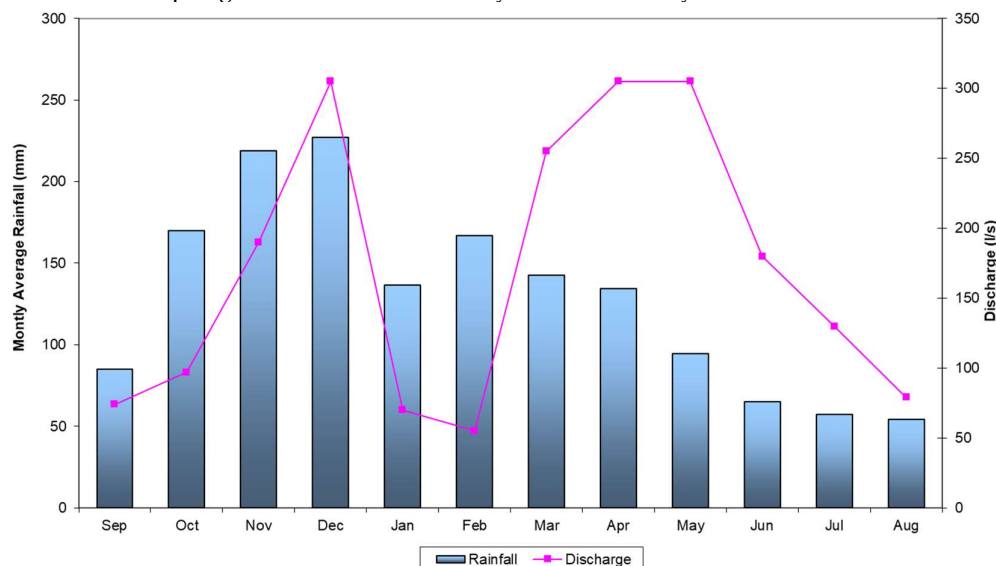


Figure 4. Monthly rainfall and Carpinetto spring (GW6) average rate.

Based on fifty-year rainfall data, Figure 4 shows that the maximum and the minimum precipitation average values occur in November/December and in August, respectively. The evolution of the outflow diagram of the Carpinetto spring (GW6), as compared to the monthly average rainfall shows a delay between the maximum precipitations, reached in November/December and the maximum flow coming from GW6 in May. In general, not only for the represented GW6 in Figure 4, regarding to the hydrological regimes of the springs under study, there is an average delay of about six months, between the meteoric inflow and the discharge of the springs.

Spring water samples, collected in June, refer to the main recharge period of the year (November and December), when rainfalls drop from clouds coming from the Atlantic, and are therefore considerably less rich in heavy isotopes. On the contrary, samples collected in October refer to waters infiltrated during summer months, when rainfalls come from clouds formed in the Mediterranean basin. These considerations have been useful in supposing, with acceptable reliability, that the aquifer residence time is about six months.

4.2. Springs Recharge Areas Identification Using Isotope Analysis

The isotopic composition of groundwater in relation to elevation is an indicator for locating the groundwater recharge area [41]. In a given groundwater basin, groundwater sources may be fingerprinted using isotopes, like ^2H and ^{18}O , which depend on infiltration altitude of rainfalls. Higher altitude will have precipitation that is depleted in ^2H and ^{18}O . If the recharge source end members at different altitudes can be identified, it is possible to partition the amount of recharge to groundwater sampled from springs or in lowland areas accordingly [13]. As a matter of fact, fissured-karstic aquifers receive the recharge from the surrounding higher elevated areas. The high altitude contributes to more depletion in heavy isotopes in rainwater, and the joints and fractures of these aquifers supply good paths for transporting this infiltrated rainwater to the lower area. The distribution of the isotopic composition shows that, in the mountain areas, heavy isotopes are more

depleted than in the plain area [13]. At the highest elevations, where the average temperatures are lower, meteoric precipitation is characterized by depletion in heavy isotopes. For $\delta^{18}\text{O}$ the depletion varies from -0.15 to -0.5‰ for each 100 m increase in elevation, with a corresponding decrease in $\delta^2\text{H}$ oscillating between -1 and -4‰. The altitude effect is used in hydrogeological studies because it allows one to determine which recharge zones are located at higher or lower altitudes. As a result, due to the fact that isotopic tracers are conservative, the spring water that is supplied at higher elevations will be low in heavy isotopes, whereas an enrichment, in heavy isotopes, will be found in water that, once infiltrated into the aquifer, supplies the springs located at lower altitude. Many studies on this topic have emphasized that the value assumed by the gradient of isotopic content varies according to meteoroclimatic characteristics [36]. The altitude effect is linked to the relation between rainfall isotope composition and elevation in meters. In this study, it has been used the average value of the vertical isotopic gradient measured in Italy of about -0.15‰/100 m elevation for $\delta^{18}\text{O}$ [38] and -1.2‰/100 m elevation for $\delta^2\text{H}$. Vertical isotopic gradients evaluation, however, requires information about precipitation hydrogen and oxygen isotopic composition and their sampling altitudes. In this study, as the isotopic content of rainfall at specific elevations were not available, in the aim of calculating the mean recharge elevation of spring water samples, we applied two different methods to verify the reliability of the results obtained.

4.2.1. Center of Mass Method

The center of mass method is a concept of weighted average, used in solids mechanics. The center of mass is the point in an object or system of objects at which the whole mass may be considered as concentrated. The distribution of mass is balanced around the center of mass and the average of the weighted position coordinates of the distributed mass defines its coordinates.

In this study, it has been considered the center of mass as a statistical and virtual point of the basin, where the whole infiltration of rainfall occurs. The Inverse Hydrogeological Balance Water Method [1] was used to evaluate the distribution of the average annual active recharge (i.e. the effective infiltration) within a GIS environment, discretizing the basin into Finite Squared Elements (FSE). Using this method it can be possible to identify the center of mass of the basin to which refers the hydrogeological characteristics of the basin. To apply the vertical isotopic gradients of oxygen and hydrogen, previously described, it has been chosen the procedure described in the following.

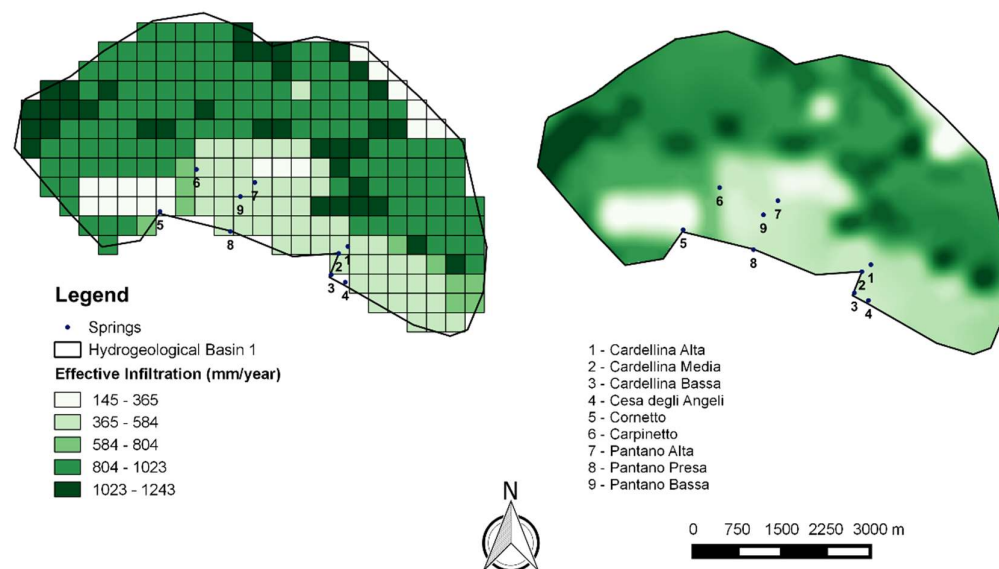


Figure 5. Effective infiltration distribution: (a) FSE, (b) interpolation.

Water stable isotopes are conservative tracers of groundwater, related to rainfalls isotopic composition before infiltration. In the aim of identifying the recharge areas of karst springs under study [20], it has been chosen GW1 spring, which has the highest outcropping elevation in the hydrogeological basin (1057 m a.s.l.), that is reasonably closer than other ones, in its groundwater isotopic content, to the rainfall value [13]. Thus, for GW1 spring, according to equation (4) it has been calculated the average infiltration elevation, weighted according to the effective infiltration values, obtained by the application of the Inverse Hydrogeological Balance Water Method (Figure 5):

$$Z_{average} = \frac{\sum_{i=1}^n z_i \cdot I_i}{I_{tot}}, \quad (4)$$

where z_i is the average elevation of the i^{th} FSE, I_i is the effective infiltration for the i^{th} FSE, n is the number of FSE into which, the hydrogeological basin has been, previously, discretized and I_{tot} is effective infiltration calculated for the whole basin. Mean recharge elevation of any spring has been calculated by the application of the vertical isotopic gradient, as for ^{18}O as for ^2H , using as end member 2002 isotope values of $\delta^{18}\text{O}$ (−8.67) and $\delta^2\text{H}$ (−50.14), referred to the GW1 (Cardellina Alta) sample, which is the closest to the recharge area in the basin (Table 3).

Table 3. Centre of Mass Method – Average elevation of Cardellina Alta Spring (GW1)

| Sample Codes | Spring | Average Elevation (m a.s.l.) | $\delta^{18}\text{O}$ (‰VSMOW) | $\delta^2\text{H}$ (‰VSMOW) |
|--------------|-----------------|------------------------------|--------------------------------|-----------------------------|
| GW1 | Cardellina Alta | 1400 | −8.67 | −50.14 |

Identified recharge elevations for each spring samples are presented in Table 4. The recharge elevation values obtained by the application of the vertical isotopic gradient of $\delta^{18}\text{O}$ (−0.15‰/100 m) and $\delta^2\text{H}$ (−1.2‰/100 m) [38] are very similar, an indication of the accuracy of the relations used. Calculated average recharge elevations for the springs is 1431 m.a.s.l., showing a gradual decrease with increasing recharge elevation.

Table 4. Recharge elevation obtained by the application of the Center of Mass Method

| Sample Codes | Recharge Elevation (m a.s.l.) | Recharge Elevation (m a.s.l.) | Average Recharge Elevation (m a.s.l.) |
|--------------|--------------------------------|-------------------------------|---------------------------------------|
| | $\delta^{18}\text{O}$ (‰VSMOW) | $\delta^2\text{H}$ (‰VSMOW) | |
| GW1 | 1400 | 1400 | 1400 |
| GW2 | 1423 | 1503 | 1463 |
| GW3 | 1448 | 1469 | 1459 |
| GW4 | 1545 | 1484 | 1515 |
| GW5 | 1306 | 1296 | 1301 |
| GW6 | 1423 | 1391 | 1407 |
| GW7 | 1455 | 1457 | 1456 |
| GW8 | 1448 | 1428 | 1438 |
| GW9 | 1474 | 1406 | 1440 |

4.2.2. The Arithmetic Average Method

In order to verify the reliability of the results obtained by Center of Mass Method, the Arithmetic Average Method was applied to identify the recharge areas of the karst aquifer (Eq. 5).

$$AAM = \frac{\sum_{i=1}^n h_i}{h_a} \quad (4)$$

where h_i is the elevation of the i^{th} FSE, located at higher elevations, than GW1 spring, and h_a is the average elevation of all FSE, located at higher elevations, than GW1 spring. For the calculation of average recharge elevation of spring water samples, we used, all the same, the 2002 isotope values of $\delta^{18}\text{O}$ (−8.67) and $\delta^2\text{H}$ (−50.14) referred to the end member sample GW1, whose recharge elevation topographically does not exceed 1412 (Table 5). Identified springs recharge elevations are presented in Table 6. Calculated average recharge elevations for the springs is 1443 m.a.s.l..

Table 5. Arithmetic Average Method – Average elevation of Cardellina Alta Spring (GW1)

| Sample Codes | Spring | Average Elevation (m a.s.l.) | $\delta^{18}\text{O}$ (‰VSMOW) | $\delta^2\text{H}$ (‰VSMOW) |
|--------------|-----------------|---------------------------------|-----------------------------------|--------------------------------|
| GW1 | Cardellina Alta | 1412 | −8.67 | −50.14 |

Table 6. Recharge elevation obtained by the application of the Arithmetic Average Method

| Spring | Recharge Elevation (m a.s.l.) $\delta^{18}\text{O}$ (‰VSMOW) | Recharge Elevation (m a.s.l.) $\delta^2\text{H}$ (‰VSMOW) | Average Recharge Elevation (m a.s.l.) |
|--------|--|---|---|
| GW1 | 1412 | 1412 | 1412 |
| GW2 | 1435 | 1515 | 1475 |
| GW3 | 1460 | 1481 | 1471 |
| GW4 | 1557 | 1496 | 1527 |
| GW5 | 1318 | 1308 | 1313 |
| GW6 | 1435 | 1403 | 1419 |
| GW7 | 1467 | 1469 | 1468 |
| GW8 | 1460 | 1440 | 1450 |
| GW9 | 1486 | 1418 | 1452 |

4.2.3. Recharge Areas Identification Methods Comparison

Figures 6 and 7 show the distribution of $\delta^{18}\text{O}$ and $\delta^2\text{H}$ values of groundwater samples with respect to recharge elevation, obtained by the application of the Center of Mass and Arithmetic Average methods.

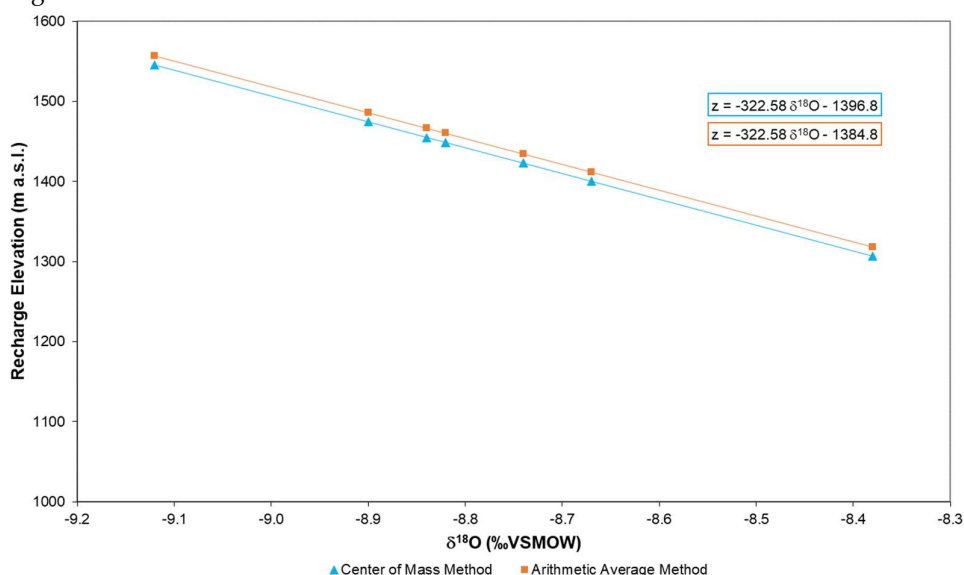


Figure 6. $\delta^{18}\text{O}$ values of spring samples plotted versus recharge elevations (October 2002).

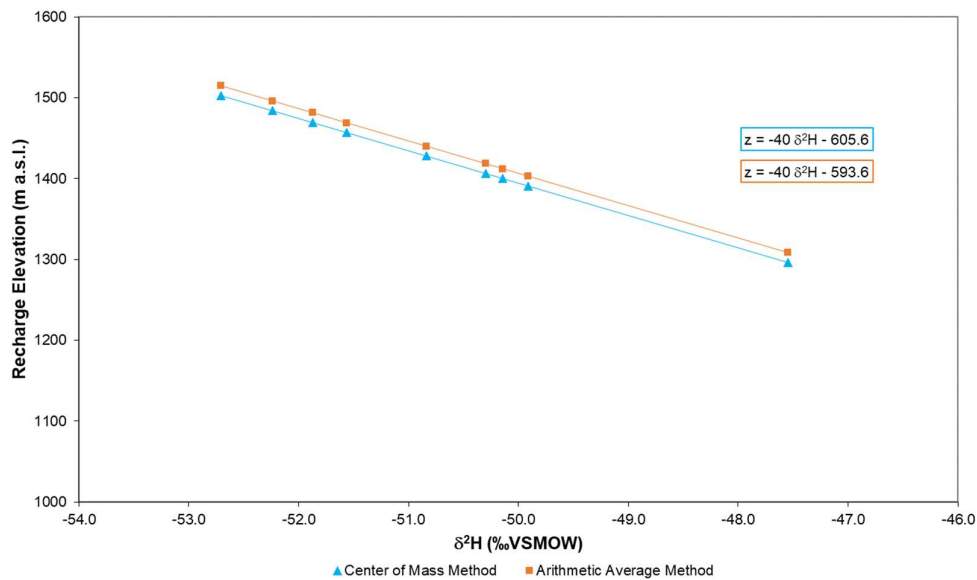


Figure 7. $\delta^2\text{H}$ values of spring samples plotted versus recharge elevations (October 2002).

Average recharge elevations calculated for the springs show a gradual decrease with increasing recharge elevation. The recharge elevation values obtained by the application of both methods are very similar, and they indicate the accuracy of the relations used.

The obtained recharge elevation values appear to be congruent with the elevations of the hydrogeological basin. In the aim of better verifying the accuracy, the spring recharge elevations have been compared with the results of the application of the Inverse Hydrogeological Balance Method, reported in Figure 5. For this reason, the basin has been divided into six elevation classes of 200 m each and the effective infiltration percentage in each class was calculated (Figure 8).

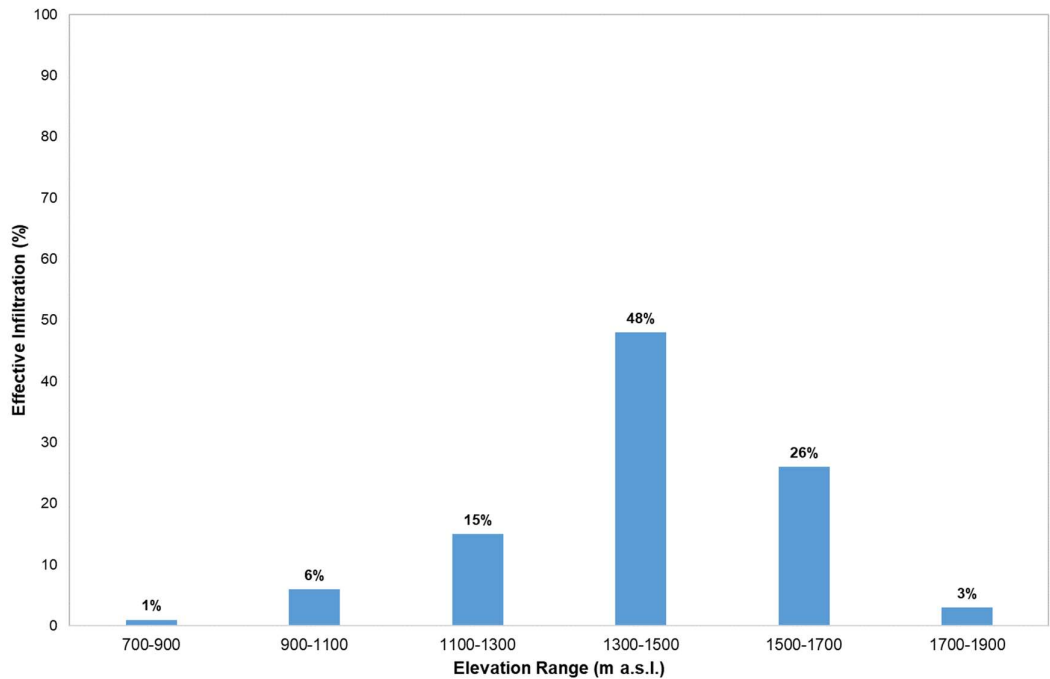


Figure 8. Bar plot of effective infiltration (%).

In the study area, the highest effective infiltration percentage has been obtained for the elevation class from 1300 to 1500 m a.s.l. (Figure 8), corresponding very well to the results of the isotopic analysis carried out for the basin. This result is due to the presence of the karst cretaceous limestone outcropping. As to be expected, in a karst framework as this hydrogeological basin is, the presence of highly fractured rocks and most of all the specific permeability conditions, due to the local karst processes development, make spread and high the infiltration, as the general isotopic depletion of groundwater sampled confirms

The springs average recharge elevations, obtained by the application of the Center of Mass Method (1431 m a.s.l.) and the Arithmetic Average Method (1443 m a.s.l.) fall within the class for which the infiltration percentage is highest according to the Inverse Hydrogeological Balance Method (Figure 8). The recharge elevations, obtained for the hydrogeological basin using the two methods described above are very similar. There is a 1% difference between the values obtained in the two cases.

5. Conclusions

This paper deals with the identification of aquifer recharge area in a karst basin in Central Italy (Figure 9), using the stable isotope ratios of hydrogen and oxygen, combined with the application of the Inverse Hydrogeological Balance Method.

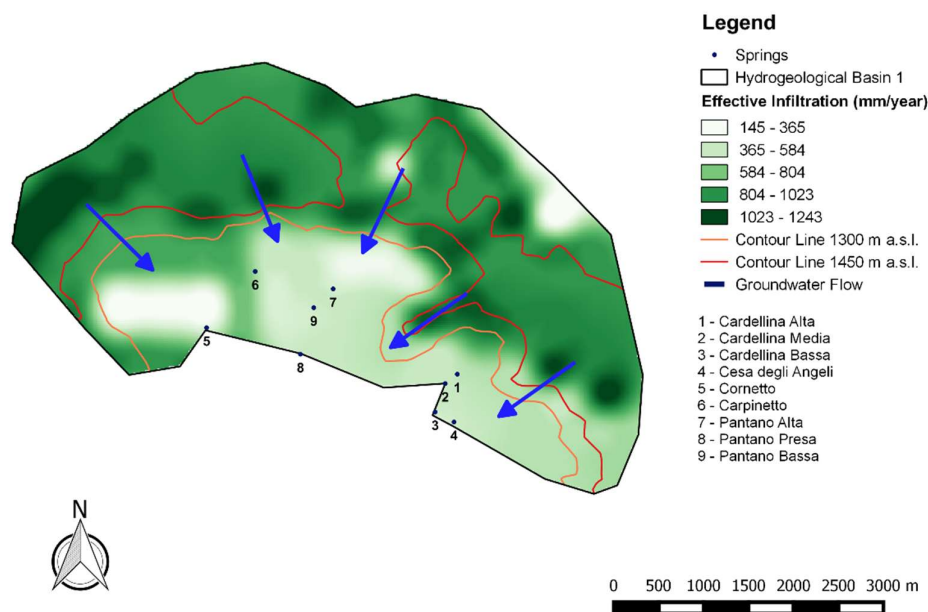


Figure 9. Hydrogeological basin recharge areas and groundwater flowpaths.

For the Cardellina springs group (GW1, GW2 and GW3), the results show an infiltration area, located in the southeast part of the basin at an average elevation of about 1450 m a.s.l., in the Upper Cretaceous carbonates. On the contrary, Cesa degli Angeli spring (GW4) has the highest infiltration elevation than all the other ones. This second infiltration area is located in Upper Cretaceous limestone at an elevation of about 1520 m a.s.l.. The recharge area of Cornetto (GW5) and Carpinetto (GW0) are located in the northeast part of the basin, at an average elevation of about 1300 and 1400 m a.s.l., respectively. The Pantano springs (GW7, GW8 and GW9) are fed by rainfall, which infiltrates in the western slope of the Assalone Mt, at an elevation of about 1450 m a.s.l.. The groundwater isotopic composition of Pantano springs (GW7, GW8 and GW9) is very similar, and as a result, their recharge elevations are very close to each other, highlighting the same infiltration area at an average elevation of about 1450 m a.s.l., in the permeable fractured outcropping of limestone and dolomite.

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References

1. Sappa, G.; Ferranti, F.; Ergul, S.; Ioanni, G. Evaluation of the groundwater active recharge trend in the coastal plain of Dar Es Salaam (Tanzania). *J. Chem. Pharm. Res.* **2013**, *5*(12), 548–552.
2. Foster, S.; Hirata, R.; Andreo, B. The aquifer pollution vulnerability concept: Aid or impediment in promoting groundwater protection? *Hydrogeol. J.* **2013**, *21*, 1389–1392.
3. Bakalowicz, M. Karst groundwater: A challenge for new resources. *Hydrogeol. J.* **2005**, *13*, 148–160.
4. Cozma, A.I.; Baciuc, C.; Moldovan, M.; Pop, I.C. Using natural tracers to track the groundwater flow in a mining area. *Procedia Environ. Sci.* **2016**, *32*, 211–220.
5. Leibundgut, C.; Maloszewski, P.; Kull, C. *Tracer in hydrology*, 1st ed.; Wiley-Blackwell, Chichester, U.K. 2009; pp. 432, ISBN 978-0-470-51885-4.
6. Barbieri, M.; Nigro, A.; Petitta, M. Groundwater mixing in the discharge area of San Vittorino Plain (Central Italy): geochemical characterization and implication for drinking uses. *Environ Earth Sci.* **2017**, *76*(11):393, DOI:10.1007/s12665-017-6719-1.
7. Massmann, G.; Sültenfuß, J.; Dünnebier, U.; Knappe, A.; Taute, T.; Pekdeger, A. Investigation of groundwater residence times during bank filtration in Berlin: A multi-tracer approach. *Hydrol. Process.* **2008**, *22*(6), 788–801, DOI:10.1002/hyp.6649.
8. Petitta, M.; Scarascia Mugnozza, G.; Barbieri, M.; Bianchi Fasani, G.; Esposito, C. Hydrodynamic and isotopic investigations for evaluating the mechanisms and amount of groundwater seepage through a rockslide dam. *Hydrol. Process.* **2010**, *24*, 3510–3520, DOI: 10.1002/hyp.7773.
9. Gasser, G.; Pankratov, I.; Elhanany, S.; Glazman, H.; Lev, O. Calculation of wastewater effluent leakage to pristine water sources by the weighted average of multiple tracer approach. *Water Resour. Res.* **2014**, *50*(5), 4269–4282, DOI: 10.1002/2013WR014377.
10. Moeck, C.; Radny, D.; Popp, A.; Brennwald, M.; Stoll, S.; Auckenthaler, A.; Berg, M.; Schirmer, M. Characterization of a managed aquifer recharge system using multiple tracers. *Sci. Total Environ.* **2017**, *609*, 701–714, DOI:10.1016/j.scitotenv.2017.07.211.
11. Nigro, A.; Sappa, G.; Barbieri, M. Application of boron and tritium isotopes for tracing landfill contamination in groundwater. *J. Geochem Explor* **2017**, *172*, 101–108, DOI: 10.1016/j.gexplo.2016.10.011.
12. Sappa, G.; Ferranti, F.; De Filippi, F.M.; Cardillo, G. Mg²⁺-based method for the Pertuso spring discharge evaluation. *Water (Switzerland)* **2017**, *9*(1), DOI:10.3390/w9010067.
13. Coplen, T.B.; Herczeg, A.L.; Barnes, C. Isotope engineering – Using stable isotopes of the water molecule to solve practical problems. In *Environmental tracers in subsurface hydrology*; Cook, P., Herczeg, A.L. (Eds.); Kluwer Academic Publishers, 2000, pp. 79–110, ISBN 0-7923-7707-9.
14. Barbieri, M.; Boschetti, T.; Petitta, M.; Tallini, M. Stable Isotopes (²H, ¹⁸O and ⁸⁷Sr/⁸⁶Sr) and Hydrochemistry Monitoring for Groundwater Hydrodynamics Analysis in a Karst Aquifer (Gran Sasso, Central Italy). *Appl. Geochem.* **2005**, *20*, 2063–2081, DOI:10.1016/j.apgeochem.2005.07.008.
15. Sappa, G.; Ergul, S.; Ferranti, F. Water Quality Assessment Of Carbonate Aquifers In Southern Latium Region, Central Italy: A Case Study For Irrigation And Drinking Purposes. *Applied Water Science* **2014**, *4*(2), 115–128, DOI:10.1007/s13201-013-0135-9.
16. Gat, J.R. Oxygen and hydrogen isotopes in the hydrological cycle. *Annu. Rev. Earth Planet. Sci.* **1996**, *24*(2–3), 225–262.
17. Rozanski, K.; Araguas-Araguas, L.; Gonfiantini, R. Isotopic patterns in modern global precipitation. In *Climate change in continental isotopic records*; Swart, P.K.; Lohmann, K.C.; McKenzie, J.; Savin, S. (Eds.); American Geophysical Union, Washington DC, 1993, Volume 78, pp. 1–36.
18. Song, C.; Wang, G.; Liu, G.; Mao, T.; Sun, X.; Chen, X. Stable isotope variations of precipitation and streamflow reveal the young water fraction of a permafrost watershed. *Hydrol. Process.* **2017**; *31*, 935–947.
19. Kirchner, J. W. Aggregation in environmental systems-Part 1: Seasonal tracer cycles quantify young water fractions, but not mean transit times, in spatially heterogeneous catchments. *Hydrol. Earth Syst. Sci.* **2016**, *20*(1), 279–297, DOI:10.5194/hess-20-279-2016.
20. Dansgaard, W. Stable isotopes in precipitation. *Tellus* **1964**, *XVI*, 436–468.

21. Gat, J.R.; Gonfiantini, R. (Ed.). *Stable isotope hydrology. Deuterium and Oxygen-18 in the water cycle*. Technical Reports Series No. 210, International Atomic Energy Agency (IAEA), Vienna, 1981; pp. 356, ISBN 92-0-145281-0.
22. Gonfiantini, R.; Roche, M.-A.; Olivry, J.-C.; Fontes, J.-C.; Zuppi, G.M. The altitude effect on the isotopic composition of tropical rains. *Chem. Geol.* **2001**, *181*(1-4), 147-167, DOI:10.1016/S0009-2541(01)00279-0.
23. Craig, H. Isotopic variations in meteoric waters. *Science* **1961**, *133*, 1702-1703, DOI: 10.1126/science.133.3465.1702.
24. Clark, I.D.; Fritz, P. *Environmental Isotopes in Hydrogeology*; CRC Press/Lewis Publishers, Boca Raton, 1997; pp. 342, ISBN 9781566702492.
25. Civita, M.; De Maio, M. *Average groundwater recharge in carbonate aquifers: a GIS processed numerical model*. Proceedings of the VII Conference on Limestone Hydrology and Fissured media, Besancon Francia, 2001.
26. Devoto, G. Note geologiche sul settore centrale dei Monti Simbruini ed Ernici (Lazio nord-orientale). *Bollettino della Società dei Naturalisti in Napoli* **1967**, *76*, pp. 1-112.
27. Devoto, G. Sguardo geologico dei Monti Simbruini (Lazio nord-orientale). *Geol. Rom.* **1970**, *9*, pp. 127-136.
28. Devoto, G.; Parotto, M. Note geologiche sui rilievi tra Monte Crepacuore e Monte Ortara (Monti Ernici-Lazio nord-orientale). *Geol. Rom.* **1967**, *6*, pp. 145-163.
29. Accordi, G.; Carbone, F. Sequenze carbonatiche meso-cenozoiche. Note illustrative alla Carta delle Litofacies del Lazio-Abruzzo ed aree limitrofe. *Quad. Ric. Scient.* **1988**, *114*(5), 11-92.
30. Damiani, A.V. Studi sulla piattaforma laziale-abruzzese. Nota I. Considerazioni e problematiche sull'assetto tettonico e sulla paleogeologia dei Monti Simbruini. In *Memorie. Descrittive Carta Geologica D'Italia*, 1990; Volume 38, pp. 145-176.
31. Bono, P.; Percopo, C. Flow dynamics and erosion rate of a representative karst basin (Upper Aniene River, Central Italy). *Environ. Geol.* **1996**, *27*(3), pp. 210-218, DOI 10.1007/BF00770434.
32. Ventriglia, U. *Idrogeologia della Provincia di Roma, IV, Regione Orientale*. Amministrazione provinciale di Roma, LL.PP., Viabilità e Trasporti, 1990.
33. Penta, F. (Sapienza University of Rome, Rome, Italy). Indagini nella zona delle sorgenti del Simbrivio. Istituto di Giacimenti Minerari e Geologia Applicata, 1956.
34. Sappa, G.; Ferranti, F. An integrated approach to the Environmental Monitoring Plan of the Pertuso Spring (Upper Valley of Aniene River). *Ital. J. Groundw.* **2014**, *3*(2), 47-55, DOI: 10.7343/as-073-14-0099.
35. White, W.B. *Geomorphology and hydrology of karst terrains*; Oxford University Press, New York, 1988.
36. White, W.B. Karst hydrology: recent developments and open questions. *Eng. Geol.* **2002**, *65*, 85-105, DOI:10.1016/S0013-7952(01)00116-8.
37. Gat, J.R.; Carmi, I. Evolution of the isotopic composition of atmospheric waters in the Mediterranean Sea Area. *J. Geophys. Res.* **1970**, *75*, 3039-3048, DOI:10.1029/JC075i015p03039.
38. Longinelli, A.; Selmo, E. Isotopic Composition of Precipitation in Italy: A First Overall Map. *J. Hydrol.* **2003**, *270*(1-2), 75-88.
39. Cappa, C.D.; Hendricks, M.B.; DePaolo, D.J.; Cohen, R.C. Isotopic fractionation of water during evaporation. *J. Geophys. Res.* **2003**, *108*, DOI:10.1029/2003JD003597.
40. Gat, J.R.; Matsui, E. Atmospheric water balance in the Amazon Basin: an isotopic evapotranspiration model. *J. Geophys. Res. Atmos.* **1991**, *96*(D7), 13179-13188, DOI: 10.1029/91JD00054.
41. Bortolami, G.C.; Ricci, B.; Susella, G.F.; Zuppi, G.M. Isotope hydrology of the Val Corsaglia, Maritime Alps, Piedmont, Italy. In *Isotope Hydrology*, International Atomic Energy Agency IAEA, Vienna, 1978; Volume 1, pp. 27-350, ISBN 92-0-040079-5.