

Granite is an Effective Helium Source Rock: Insights from the Helium Generation and Release Characteristics in Granites from the North Qinling Orogen, China

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Abstract: Global helium (He) shortage is a challenging problem; however, the types of helium source rock and the mechanisms of He generation and release therein remain still poorly understood. In this study, in order to evaluate the potential of granite as an effective helium source rock, we collected granitic samples from the North Qinling Orogen, Central China, in the south of the helium-rich Weihe Basin. The helium generation and release behaviors in granite were studied through analysis of U and Th concentrations, EMPA images, and He and Ar concentrations and isotopic ratios extracted by crushing and stepwise heating. The results indicate that Ar has a better retention and a lower mobility than He. $^3\text{He}/^4\text{He}$ ratios released by crushing and stepwise heating are 0.016–0.056 R_A and 0.003–0.572 R_A , respectively, where R_A is the atmospheric $^3\text{He}/^4\text{He}$ of 1.4×10^{-6} , reflecting a crustal and radiogenic source. Helium concentrations extracted by the two ways are 0.13–0.95 u cm^3 STP/g and 7.82–115.62 u cm^3 STP/g, respectively, suggesting that matrix-sited He accounts for more than 98% of total helium preserved in granite. In addition, the total generated He amounts in granites are calculated based on the measured U and Th concentrations in granitic samples. Dividing the preserved He quantities by the generated He amounts, it turned out that less than 10% of He produced since the formation of the granite is preserved in the rock over geological time, suggesting that more than 90% generated He can be transferred to the Weihe Basin. Temperature and fracture are the two critical factors controlling He release. Based on the relationship between He diffusivity of granites and temperature and the He closure temperatures of a variety of U- and Th-rich minerals (27–250°C), we estimate that He can be partially released out of granite at the depths <400 m and totally released at the depths >7800 m. Fractures provide effective transfer of free He from deep source rocks to shallow reservoirs. Finally, a model on granite as an effective helium source rock is established. We suggest exploring He resources in hydrocarbon basins with granitic basement (or adjacent to granite bodies), high geothermal field, and young active fractures.

Key words: granite, helium source rock, helium diffusivity, crushing, stepwise heating

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1 Introduction

Helium is a strategic resource that has been widely used in military, nuclear industry, cryogenic superconductor, and scientific research (Cai et al., 2010). It is often extracted as a by-product of natural gas processing or during the production of liquefied natural gas (LNG). At present, the economically recoverable He resources in the world have low $^3\text{He}/^4\text{He}$ ratios, suggesting that most of them were derived from the radioactive decay of U and Th (Ballentine and Lollar, 2002). At the end of 2006, He resources and reserves of the world were estimated to be about 51.9 and 7.4 billion cubic meters, respectively. The annual He demand is approximately 170 million cubic meters (Hamak, 2016). The He reserve and supply of the USA both account for more than 50% of the global amounts. At present, the world supply cannot meet the He demand, causing a “Liquid Helium Crisis” (Simon et al., 2016) and a “MRI Helium Crisis” (Mahesh and Barker, 2016). This issue has attracted broad attention on He reserves in many countries including China (Li et al., 2011; Zhang et al., 2019a; Zhang et al., 2019b; Tao et al., 2019), the USA (Brown, 2010), Qatar (Flower, 2012), Algeria (Reinoehl, 2012), Russia (Yakutseni V.P., 2014) and Tanzania (Danabalan et al., 2016; Ballentine et al., 2017).

China is a He-poor country, and He has been mainly exploited from a small gas field in the Sichuan Basin for more than 50 years (Wang et al., 2011). In recent years, extensive helium resources have been discovered from geothermal wells in the Weihe Basin, Central China (Liu et al., 2009; Li et al., 2011, 2015, 2016; Han et al., 2014; Zhang, 2015; Zhang et al., 2018; Zhang et al., 2019a; Zhang, 2019). The average He concentration in 101 wells is 1.50% and the maximum value is up to 9.23%, being the highest reported He concentration in China (Zhang

et al., 2018). The He isotopic ratios ($^3\text{He}/^4\text{He} = 2.2\text{--}78 \times 10^{-8}$, Zhang, 2015) reflect a primarily crustal and radiogenic source except for a few outliers. The correct estimation and economic recovery of He resources in the Weihe Basin are vital to the He market in China and around the world.

The urgent demand and uneven distribution of He around the world leads to an interesting question: where does the He come from? By analyzing the geologic settings of other He-rich gas fields, such as the Hugoton-Panhandle giant gas field in the USA (Ballentine and Lollar, 2002), the Weiyuan gas field in China (Zhang et al., 2015) and the Hassi R'Mel gas fields in Algeria (Sabaou, et al., 2009), it has been suggested that the accumulation of He is closely related to the granitoid basement or intrusions (Fig. 1). However, although the generation and release / preservation of He in sedimentary rocks and U- and Th-rich minerals have been studied in the field of groundwater modeling (Fourré et al., 2011; Tolstikhin et al., 2011) and U-Th-He dating (Wolf et al., 1998; Reiners, et al., 2002), little research on the system of He generation and release in granite has been carried out (Hussain, 1997).

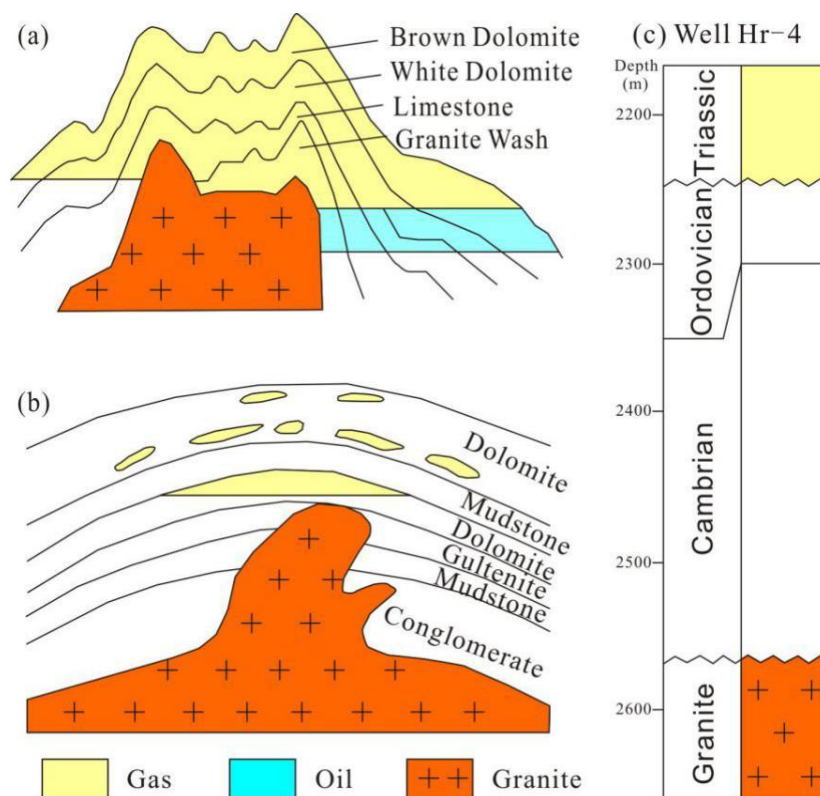


Fig. 1. Correlation between He-rich gas fields and granite.

(a) Hugoton-Panhandle gas field, USA (after Ballentine and Lollar, 2002); (b) Weiyuan gas field, China (after Zhang et al., 2015); (c) Hassi R'Mel, Algeria (after Sabaou et al., 2009).

The term ‘helium source rock’ has been used by Burwash and Cumming (1974) and Brown (2010) but without clear definition. In this study, as analogue to the definition of hydrocarbon source rock, we define helium source rock as “a kind of rock that can generate and release large amounts of helium”. In order to provide scientific basis for He exploration and reserve estimation, we focus herein on granite and evaluate its potential to be an effective helium source rock, based on analysis of He generation and release behaviors using the granites from the North Qinling Belt as an example. U, Th concentrations were analyzed and EPMA (Electro Probe Microanalysis) images were used to evaluate the characteristics of U- and Th-rich minerals and the productivity of He. He in granites was extracted by crushing and stepwise heating, respectively, for assessing the He release behaviors, including the He release ratios and the influence of fracture and temperature on the release processes. In addition, K and Ar were also analyzed to assist with the interpretation of the results.

2 Geological Setting and Sampling

The Weihe Basin is a Cenozoic graben basin with a large number of faults and fractures within the basin and the basement. The Cenozoic strata are several kilometers thick and up to over 6 km. There are a large number of granitic bodies distributed or exposed in the Qinling Orogenic Belt (QOB), which is in fault contact with the Weihe Basin to the north (Fig. 2). The helium resource in the Weihe Basin has been assumed to be mainly derived from the QOB (Li et al., 2011; Han et al., 2014).

The QOB is situated between the North China Block (NCB) and the South China Block (SCB). Detailed studies suggest that the QOB has experienced multistage subduction, accretion, and collision of these two blocks (Dong and Santosh, 2016). After experiencing a significant collision in the Middle to Late Triassic, the QOB evolved into an intra-continental evolutionary process (Zhang et al., 2001), accompanied with magmatism, metamorphism, and deformation (Dong and Santosh, 2016). Most of the granites in the QOB were formed in the Mesozoic and Paleozoic (Wang et al., 2015).

In 2012-2016, 88 granite samples were collected from the 7 rock masses that make up the NQB (North Qinling Belt) to evaluate the average concentrations of He source elements, mainly U and Th, in the rock masses. In addition, we selected 2 Mesozoic granite samples (14-ZQ and 14-HNP) with high U and Th concentrations to study the He generation and release behaviors within these units. The two samples were collected from Zhuque forest park (33°49'25"N, 108°31'48"E) and Huangniupu (34°15'27"N, 106°54'04"E), respectively. Based on previous investigations on the ages of the adjacent rock masses (shown in Fig. 2), the average ages of the granite samples were calculated to be 212 Ma and 213 Ma for 14HNP and 14ZQ, respectively. This indicates that the plutons formed during the Late Triassic, which corresponds to the timing of the Indosinian movement (Lu, 2000; Zhang et al., 2001).

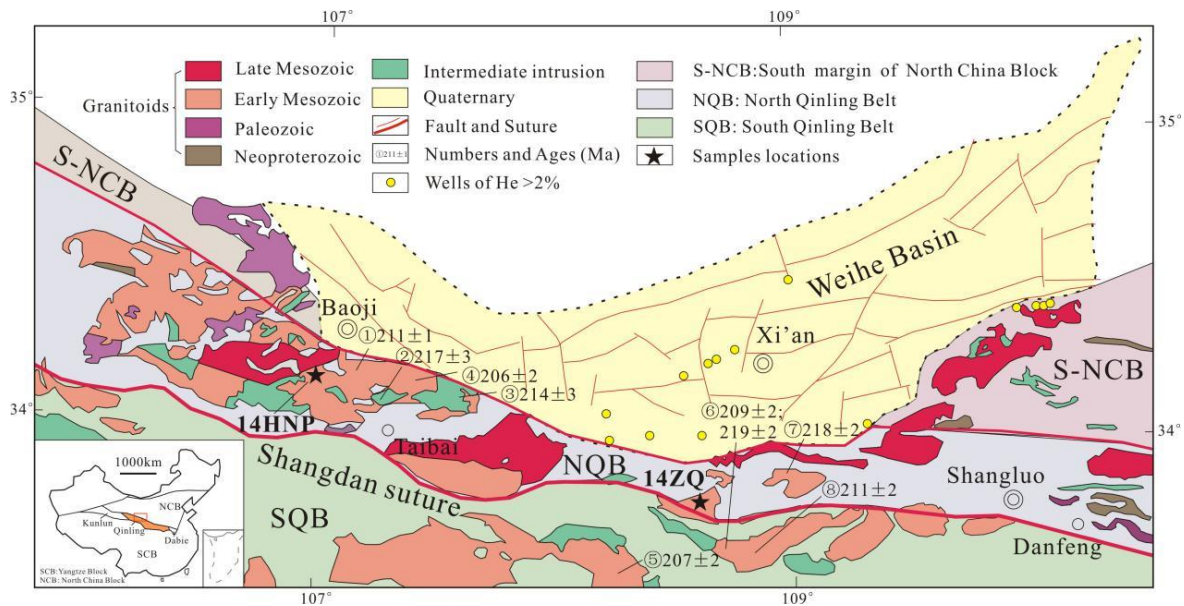


Fig.2. Distribution of granitoids in the Qinling Orogenic Belt (after Wang, et al., 2013).

The age data are cited from Zhang et al. (2006) for ①④; Lu et al. (2000) for ②③; Dong et al. (2011) for ⑤; Yang et al. (2009) for ⑥; Jiang et al. (2010) for ⑦; and Sun (2002) for ⑧. China basemap after China National Bureau of Surveying and Mapping Geographical Information.

3 Analytical Methods

U and Th concentrations were measured for 88 granitic samples, in which two samples with high U and Th contents were selected to do further detailed work, including EPMA and the analysis of He and Ar concentrations and isotopic ratios. The reason why we choose U- and Th-rich samples is to evaluate the processes in samples with elevated He quantities and thus a higher precision in the subsequent noble gas testing.

U, Th, and K concentration analysis and EPMA were performed in the Xi'an Center, China Geology Survey. The instruments used to measure the concentrations of U, Th and K were the Laser Fluorescence Spectrum, AAS (Atomic Absorption Spectrum) and ICP-AES (Inductively Coupled Plasma-Atomic Emission Spectrometry), respectively. To observe morphological features and identify distinguishable phases, SEI (secondary electron images) and BSE (back-scattered electron images) images from EPMA were used. Furthermore, part of U-Th-rich mineral compositions and phase bands were analyzed using a beam spot that was 1–5 μm in diameter.

The concentrations and isotopic ratios of He and Ar were analyzed using a Noblesse noble gas mass spectrometer at the Lanzhou Center for Oil and Gas Resources, Institute of Geology and Geophysics, Chinese Academy of Sciences. The noble gases in rocks were extracted by crushing in the first place, followed by stepwise heating. During the first stage, bulk sample (~10 g and ~3–8 mm in diameter) was placed into a high-vacuum degassing tank sealed by six sets of bolts, which were fixed to an electromagnetic breaker. The pre-processing crushing system was pumped to lower than 10^{-5} Pa, eliminating air contamination to the maximum extent. Then, the sample was stroke 3000 times in one minute in vacuum with an electromagnetically controlled device, in order to release the gases. The released

gases were purified first using a spongy titanium furnace to remove active gases, such as C₁-C₄, H₂O, N₂ and CO₂ etc., and then using two linked Zr-Al getters to remove H₂ at ambient temperature. The purified gases were then absorbed in a cryogenic trap at the temperature of 8 degree K for 20 min. Then He and Ar were released under 15 degree K and 100 degree K respectively and introduced into the Noblesse SFT noble gas mass spectrometer separately for He and Ar concentrations and isotopic ratios measurement (Cao et al., 2018).

The crushed sample was then wrapped in Al foil and placed into a sample disk, which was connected to the double-vacuum high-temperature furnace. The whole system was evacuated for over 24 hours to eliminate air contamination. Noble gases were extracted using stepwise heating in a double-vacuum furnace at 250°C, 700°C, 1100°C and 1600°C successively, with the duration of each extraction step being 40 minutes. The purification and measuring processes of gases released by heating were the same as those extracted by crushing. During the whole process, air collected from the top of Gaolan Mountain in Lanzhou City, China was used as an internal standard. An air standard was analysed for each sample and the air standard results are reproducible, suggesting that the whole analysis system is stable and the testing results are credible. The experimental uncertainties for both He and Ar were <10%.

4 Results

4.1 U, Th and K concentrations in granites

The average U and Th concentrations in 7 granitic plutons (88 samples) and 2 granitic samples are listed in Table 1. The parameter of [U+0.24Th] is applied to evaluate the helium production capacity, where 0.24 is the ratio of the He production rate of Th to that of U (Zartman et al., 1961). It can be seen that the granites in NQB record high concentrations of U and Th. The average [U+0.24Th] values of 7 plutons range from 7.27 to 16.82 ppm, which are 1.5–3.4 times of the average value in the crust (5.00 ppm). The average [U+0.24Th] value in the crust is calculated based on the Clark values of U and Th (2.7 and 9.6ppm, Taylor and McLennan, 1985). The two samples with the highest [U+0.24Th] values, i.e., 14ZQ (U+0.24Th = 45.4 ppm) and 14HNP (U+0.24Th = 17.5 ppm), were used for further He and Ar concentrations and isotopic ratios analysis.

The concentrations of K in samples 14ZQ and 14HNP are 2.46% and 3.44% respectively, slightly higher than the Clark Value of K (2.09%, Taylor and McLennan, 1985). It suggests that there is not much difference in the Ar productivity between granite and other types of rocks.

Table 1 U, Th and K concentrations in granitic samples from North Qinling Orogen, China

Sample ID	Number of samples	U (ppm)	Th (ppm)	U+0.24Th ^a (ppm)	K (wt%)
Baoji Pluton ^b	19	3.94	18.32	8.33	-
Taibai Pluton	12	3.97	21.63	9.16	-
Cuihuashan Pluton	12	9.41	15.77	13.20	-
Lantian Pluton	17	6.29	17.67	16.82	-
Muhuguan Pluton	6	3.45	15.92	7.27	-
Huashan Pluton	7	4.07	11.37	6.80	-
Laoniushan Pluton	11	4.17	20.35	9.06	-
14ZQ	1	38.5	28.7	45.38	2.46
14HNP	1	13.6	16.4	17.54	3.44
Clark Value	-	2.7	9.6	5.00	2.09

^a 0.24 is the ratio of the He production rate of Th to that of U (Zartman et al., 1961).

^b The listed U and Th concentrations are the average ones for the 7 plutons.

^c Clark Values are quoted from Taylor and McLennan (1985).

4.2 EPMA results

EPMA images indicate that U- and Th-rich minerals are widely distributed throughout the samples (Fig. 3), as is consistent with the high U and Th contents of these rocks, according to our chemical analysis shown above. Grain sizes of those minerals range from 10 μm to 200 μm, except for a single elongate grain (~600 μm, shown in Fig. 3f).

The numbered dots in Fig. 3 show the components that were analyzed. Due to the ubiquitous isomorphic replacement in accessory minerals, it is often difficult to unambiguously identify the minerals. Therefore, we tentatively identified the minerals by their main elements (>10 wt% for its oxide) when it is uncertain. As we can see in Fig. 3, U and Th are mostly concentrated in uranothorite (No. 1) and thorite (No. 4), or are associated with Ti, Nb and Y (No. 2, 6). Part of U and Th are present in monazite (No. 3, 5), apatite (Fig. 3a), xenotime (Fig. 3c), magnetite (Fig. 3c), and zircon (Fig. 3d). These U- and Th-rich minerals are distributed within magmatic minerals, such as K-feldspar (Fig. 3a), albite (Fig. 3d) and quartz (Fig. 3e) or between them (Figs. 3b, e, f).

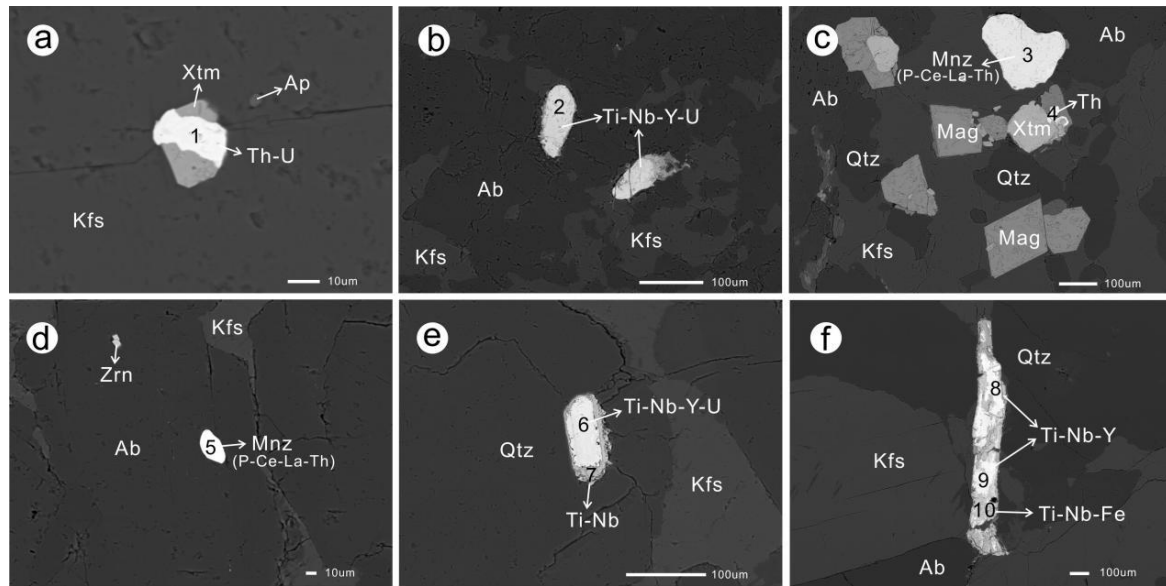


Fig.3. Representative BSE (backscattered electron) images of granites from the Qinling Orogen, China.

(a)-(c) are from sample 14ZQ; (d)-(f) are from sample 14HNP. The grain sizes range from 10 μm to 200 μm , except for a single elongate accessory mineral ($\sim 600 \mu\text{m}$), which is shown in (f). U and Th are mostly concentrated in thorite (No. 1) and uranothorite (No. 4), and associated with Ti, Nb, Y (No. 2, 6). U- and Th-bearing minerals include monazite (No. 3, 5), apatite (a), xenotime (c), magnetite (c), and zircon (d). Th-U: uranothorite; Th: thorite; Qtz: Quartz; Kfs: K-feldspar; Ab: Albite; Xtm: Xenotime; Ap: Apatite; Mag: Magnetite; Zrn: Zircon; Mnz: Monazite.

4.3 He and Ar geochemistry in granites

4.3.1 He concentrations and $^3\text{He}/^4\text{He}$ ratios

The concentrations and isotopic ratios of He are presented in Table 2. The He contents of samples 14ZQ and 14HNP measured by crushing are 0.949 and 0.129 $\text{ucm}^3 \text{STP/g}$, respectively. The $^3\text{He}/^4\text{He}$ ratios in the two samples are 0.016 R_A and 0.056 R_A , respectively; thus, both exhibit characteristics of a crustally-derived source (0.008 R_A , Ballentine and Burnard, 2002).

The stepwise heating results (Table 2, Fig. 4) show that the total amounts of He released from 14ZQ and 14HNP are 114.74 and 7.76 $\text{ucm}^3 \text{STP/g}$, respectively. In addition, most of the He (80.47% for 14ZQ and 78.68% for 14HNP) was released at a temperature of 700°C. The residual He in the rock was mainly released at 1100°C before final heating to 1600°C. These $^3\text{He}/^4\text{He}$ ratios are similar to those obtained by crushing, and both record typical features of crust-derived He: ranging from 0.003 R_A to 0.572 R_A (average 0.033 R_A) for 14ZQ and from 0.009 R_A to 0.036 R_A (average 0.030 R_A) for 14HNP. However, due to the low He concentrations extracted from 14ZQ at 250°C and from 14HNP at 250°C and 1600°C, the experimental errors of their corresponding $^3\text{He}/^4\text{He}$ ratios are too large for these results to be meaningful. We can see that the minimum $^3\text{He}/^4\text{He}$ ratios appeared at 1100°C, while the maximum He concentrations occurred at 700°C (Fig. 4), possibly reflecting the better retention of ^3He than ^4He in granite rocks.

Table 2 He and Ar concentrations, isotopic ratios in granites from Qinling Orogen, China

Sample ID	Extraction method	^4He (1 σ) $\text{ucm}^3 \text{STP/g}$	R/R_A^a (1 σ)	^{40}Ar (1 σ) $\text{ucm}^3 \text{STP/g}$	$^{40}\text{Ar}/^{36}\text{Ar}$ (1 σ)	
14ZQ	Crushing	0.949 \pm 0.001	0.016 \pm 0.002	12.79 \pm 0.01	294.00 \pm 0.92	
	Stepwise heating	250°C	0.001 \pm 0.0002	-	0.82 \pm 0.003	244.45 \pm 30
		700°C	92.336 \pm 0.11	0.039 \pm 0.002	5.77 \pm 0.002	348.14 \pm 2.8
		1100°C	22.203 \pm 0.01	0.003 \pm 0.0006	15.65 \pm 0.004	1811.48 \pm 50
		1600°C	0.200 \pm 0.0007	0.572 \pm 0.07	1.97 \pm 0.001	334.86 \pm 13
		Total	114.740	0.033	24.20	702.71
14HNP	Crushing	0.129 \pm 0.001	0.056 \pm 0.013	4.09 \pm 0.004	304.41 \pm 1.70	
	Stepwise heating	250°C	0.001 \pm 0.0001	-	0.72 \pm 0.0003	201.81 \pm 14
		700°C	6.108 \pm 0.016	0.036 \pm 0.003	4.06 \pm 0.0087	518.44 \pm 6.7
		1100°C	1.653 \pm 0.0006	0.009 \pm 0.004	28.70 \pm 0.013	4121.55 \pm 140
		1600°C	0.001 \pm 0.0001	-	2.00 \pm 0.0001	482.19 \pm 14
		Total	7.761	0.030	35.49	1575.07

^a R is the ratio of $^3\text{He}/^4\text{He}$ for sample; R_A is the atmospheric ratio of $^3\text{He}/^4\text{He} = 1.4 \times 10^{-6}$.

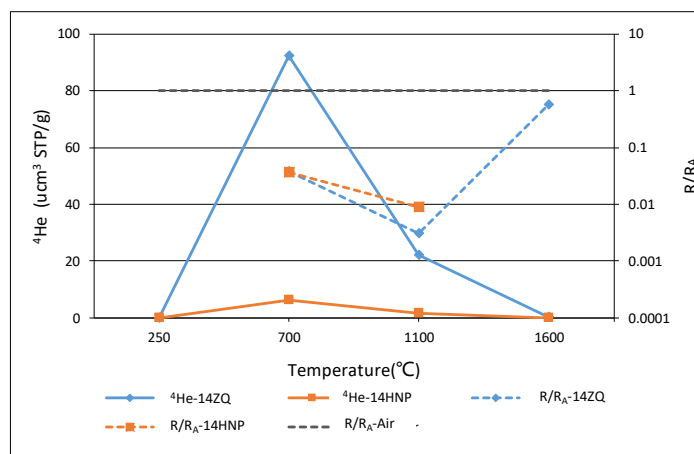


Fig.4. He concentrations and isotopic ratios released by stepwise heating.

Most He (80.47% for 14ZQ and 78.68% for 14HNP) is released at a temperature of 700°C, whereas the R/R_A reaches its peak at 1100°C, potentially suggesting better retention of ^3He than ^4He . Isotopic data at 250°C (14ZQ and 14HNP) and 1600°C (14HNP) are absent due to their unacceptable errors produced by the low He concentrations released at these temperatures. The $^3\text{He}/^4\text{He}$ ratios ranging from $0.003R_A$ to $0.572R_A$ are typical of crust-derived He.

4.3.2 Ar contents and $^{40}\text{Ar}/^{36}\text{Ar}$ ratios

The ^{40}Ar concentrations and $^{40}\text{Ar}/^{36}\text{Ar}$ isotopic ratios of these samples were also measured following crushing and stepwise heating. In the crushing experiments, 14ZQ released higher ^{40}Ar content (16.25 ucm^3 STP/g), with a $^{40}\text{Ar}/^{36}\text{Ar}$ isotopic ratio (294.00) similar to that of the atmosphere (295.99, Sano et al., 2013). Gas released from 14HNP has a lower ^{40}Ar content (5.15 ucm^3 STP/g), but a higher $^{40}\text{Ar}/^{36}\text{Ar}$ ratio (304.41), indicating that partial radiogenic ^{40}Ar was released during 14HNP crushing.

Results of stepwise heating (Table 2, Fig. 5) show that in both samples, the released ^{40}Ar contents reach their peak values at 1100°C (15.65 ucm^3 STP/g for 14ZQ and 28.70 ucm^3 STP/g for 14HNP), which correspond to the temperature that released Ar with maximum $^{40}\text{Ar}/^{36}\text{Ar}$ isotopic ratios (1811.48 for 14ZQ and 4121.55 for 14HNP). It is clear that ^{40}Ar is mainly released at 1100°C and thus at a higher temperature than ^4He (700°C), indicating the higher mobility of He relative to Ar, which could be related to the smaller molecular radius of He than Ar.

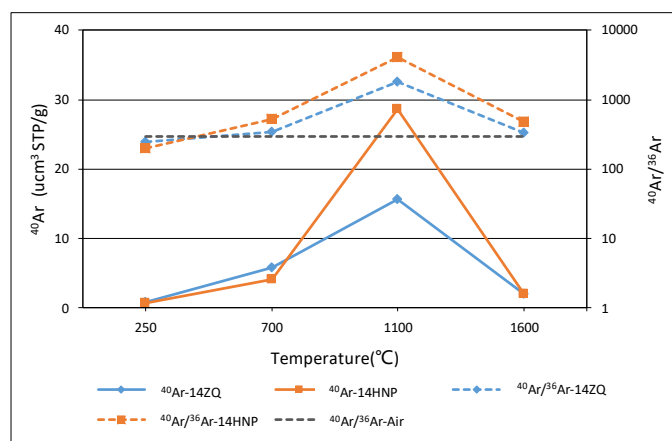


Fig.5. Ar concentrations and isotopic ratios extracted by stepwise heating.

A large proportion of Ar was released at 1100°C (64.64% for 14ZQ and 82.55% for 14HNP), corresponding to the $^{40}\text{Ar}/^{36}\text{Ar}$ peak. Compared to the temperature range that released the most He in Fig. 4 (250–700°C), it is obvious that Ar has a better retention capacity than He.

4.3.3 Comparison of the crushing and stepwise heating results

The He contents extracted by crushing are only 0.82% and 1.65% of the total released He quantities for 14ZQ and 14HNP, respectively. These results are slightly lower than those of Matsumoto et al. (2002), who suggested that ~4.37% of the total He content was released by crushing in Archean komatiites. Furthermore, Tolstkhin et al. (2016) calculated that this proportion was ~3% for amphibole in alkaline granites. The differences between these results are likely caused by the diverse U and Th contents used in the samples. When the U and Th contents are higher, the proportion of radiogenic He sited in matrix is higher, and the proportion of matrix-sited He released by crushing is lower. Comparing the two samples in this work, sample 14ZQ has higher U and Th concentrations than 14HNP and yielded a lower proportion of He released by crushing. The $^3\text{He}/^4\text{He}$ ratios released by crushing and stepwise heating are both

typically radiogenic.

Based on the same calculation method, the proportions of radiogenic $^{40}\text{Ar}^*$ (where $^{40}\text{Ar}^* = ^{40}\text{Ar} - 295.5 \times ^{36}\text{Ar}$) released by crushing are 0.00% and 0.41% for 14ZQ and 14HNP, respectively. The proportions of $^{40}\text{Ar}^*$ released by crushing are lower than those of He. This can be explained by the smaller molecule and higher mobility of He than those of Ar. The $^{40}\text{Ar}/^{36}\text{Ar}$ ratios obtained by stepwise heating are 2–5 times larger than those extracted by crushing, thus suggesting that more radiogenic $^{40}\text{Ar}^*$ was released during stepwise heating.

4.3.4 He and Ar diffusion coefficients in granites

Based on the He and Ar extraction results by stepwise heating, their diffusion coefficients (Table 3) can be calculated using the spherical diffusion equation presented by Fechtig and Kalbitzer (1966). Since the mean particle size in stepwise heating experiments is $\sim 200 \mu\text{m}$, the radius (r) of the spherical diffusion domain is assumed to be $100 \mu\text{m}$. The diffusivity of Ar in this study is assessed with radiogenic $^{40}\text{Ar}^*$.

The He diffusion coefficient in fine-grained ($r = \sim 100 \mu\text{m}$) granite particles are $4.46\text{--}4.82 \times 10^{-9} \text{ cm}^2/\text{s}$ at 700°C . At 1100°C , the coefficient of He rises to $2.00\text{--}3.05 \times 10^{-8} \text{ cm}^2/\text{s}$. The diffusion coefficients of Ar are $1.30\text{--}13.7 \times 10^{-11} \text{ cm}^2/\text{s}$ at 700°C , lower than those of He by two orders of magnitude. At 1100°C , the diffusion coefficients of Ar ($1.51\text{--}1.71 \times 10^{-8} \text{ cm}^2/\text{s}$) are lower than those of He by approximately 1.5 times.

Table 3 Results of stepwise heating and calculated He and Ar diffusion coefficients

Sample	T °C	^4He ucm ³ STP/g	Fractional loss	D cm ² /s	$^{40}\text{Ar}^*$ ucm ³ STP/g	Fraction	D cm ² /s
14ZQ	250	0.001	0.00%	–	0.00	0.00%	–
	700	92.335	80.48%	4.82×10^{-9}	0.87	6.15%	1.37×10^{-11}
	1100	22.203	99.83%	2.00×10^{-8}	13.09	98.37%	1.71×10^{-8}
	1600	0.200	100.00%	–	0.23	100.00%	–
14HNP	250	0.001	0.01%	–	0.00	0.00%	–
	700	6.108	78.69%	4.46×10^{-9}	1.75	5.99%	1.30×10^{-11}
	1100	1.653	99.98%	3.05×10^{-8}	26.65	97.34%	1.51×10^{-8}
	1600	0.001	100.00%	–	0.78	100.00%	–

5 Discussions

5.1 He sited in fluid inclusions and minerals matrix in granite

It is generally agreed that fluid inclusions (FIs) and mineral matrix are the two main reservoirs for He and other noble gases in rocks (Scarsi, 2000; Barry et al., 2015). Common extraction techniques include crushing, the stepwise heating of powders following crushing (Graham et al., 1992), and laser extraction (Scarsi, 2000). Although crushing under high-vacuum conditions is commonly used to extract He in FIs to trace processes that occurred before the formation of minerals and rocks (Day et al. 2015), it has been demonstrated that matrix-sited He may also be released during this process (Hilton et al., 1993; Matsumoto et al., 2002; Moreira et al., 2003). Furthermore, previous step-crushing studies have suggested that a negligible amount of gas from the matrix can be released by crushing during a short period (Scarsi, 2000; Hilton et al., 2011; Barry et al., 2015; Tolstikhin et al., 2016).

In this study, we assume that a small portion of matrix-sited He can be extracted by crushing. In addition, the ^4He obtained by crushing is composed of the ^4He in both FIs and the matrix, as follows:

$$^4\text{He}_{\text{crushing}} = ^4\text{He}_{\text{FIs}} + X_4 \cdot ^4\text{He}_{\text{matrix}} \quad (1)$$

where X_4 is the proportion of the matrix-sited ^4He released by crushing. Then, the ^4He extracted by stepwise heating is the residual radiogenic He:

$$^4\text{He}_{\text{heating}} = (1 - X_4) \cdot ^4\text{He}_{\text{matrix}} \quad (2)$$

Combining the above two equations, we can deduce that:

$$^4\text{He}_{\text{crushing}} = ^4\text{He}_{\text{FIs}} + X_4 / (1 - X_4) \cdot ^4\text{He}_{\text{heating}} \quad (3)$$

An analogous approach can be applied to calculate the ^3He in FIs:

$$^3\text{He}_{\text{crushing}} = ^3\text{He}_{\text{FIs}} + X_3 / (1 - X_3) \cdot ^3\text{He}_{\text{heating}} \quad (4)$$

Due to the similar geological backgrounds, petrological features and rock-forming ages (Fig. 2) of the two samples, their formation environments are assumed to be the same, which implies that the He trapped in the primary fluid inclusions ($^4\text{He}_{\text{FIs}}$) of the two samples are similar. Additionally, the experimental procedures for the two samples are identical, which allows us to assume the same proportions of matrix-sited He released by crushing (X).

Based on Equation (1–4), X_4 equals 0.76% (Table 4) and X_3 equals 0.23%, suggesting better retention of ^3He than ^4He , which is in accordance with our results in section 4.3.1 and the He isotopes retention behaviors in amphibole and some chemical sediments (Tolstikhin, et al., 1996; Tolstikhin, et al., 2016). In addition, the $^4\text{He}_{\text{FIs}}$ and $^3\text{He}_{\text{FIs}}$ are calculated to be $0.07 \text{ ucm}^3 \text{ STP/g}$ and $0.94 \times 10^{-8} \text{ ucm}^3 \text{ STP/g}$ respectively, thus $^3\text{He}/^4\text{He}$ in FIs is calculated to be 1.33×10^{-7} (Table 4). Based on a simple two-phase mixing model,

where the $^3\text{He}/^4\text{He}$ ratios of mantle and crust endmembers are taken as 1.1×10^{-5} and 2×10^{-8} respectively (Mamyrin and Tolstikhin, 1984), the percentage of mantle-derived He is calculated as 1.03%, which suggests a mainly crustal source for granites from QOB. These results are consistent with those of Wang et al. (2015), who maintained that the granites exposed in the western Qinling Orogen are primarily I-type and I-A-type rocks, i.e., they were formed from crust rather than mantle. In addition, the matrix-sited He are calculated based on Equation (2), suggesting that most of matrix-sited He (99.24%) are released by stepwise heating (Table 4).

The calculation above is proposed for the first time in this study to evaluate the relationship between matrix-sited gas and those in fluid inclusions during crushing. However, the calculation in this study has large uncertainties due to the small amounts of samples used here. It is recommended to use more samples (>3, enough for statistical analysis) to reduce errors. In this work, the high amounts of radiogenic ^4He have a big influence on the crushing results. It is recommended that U- and Th-poor minerals and rocks should be fully characterised when discussing the noble gases of FIs.

Table 4 Calculated He concentrations and isotopic ratios in the FIs and matrix

	14ZQ		14HNP	
	^4He ucm ³ STP/g	$^3\text{He}/^4\text{He}$ 10 ⁻⁸	^4He ucm ³ STP/g	$^3\text{He}/^4\text{He}$ 10 ⁻⁸
FIs ^a (Extracted by crushing)	0.07	13.30	0.07	13.30
Matrix ^b	115.62	4.57	7.82	4.25
Extracted by crushing ^c	0.88 (X ₄ =0.76%)	-	0.06 (X ₄ =0.76%)	-
Extracted by heating	114.74 (99.24%)	-	7.76 (99.24%)	-

^a $^4\text{He}_{\text{crushing}} = ^4\text{He}_{\text{FIs}} + X_4/(1-X_4) \cdot ^4\text{He}_{\text{heating}}$, assuming that the He concentrations and isotopic compositions in the FIs of the two samples are the same, where X_4 is the proportion of the matrix-sited ^4He released by crushing; the Ar concentrations and isotopic compositions in the FIs are uncertain, see section 5.1 for details.

^b $^4\text{He}_{\text{matrix}} = ^4\text{He}_{\text{heating}}/(1-X_4)$, He isotopes are the same as the results of stepwise heating.

^c $^4\text{He}_{\text{Matrix, extracted by crushing}} = ^4\text{He}_{\text{matrix}} - ^4\text{He}_{\text{heating}}$

5.2 He and Ar production and preservation

Based on the law of radioactive decay, the production of ^4He and $^{40}\text{Ar}^*$ (in ucm³ STP/g) can be calculated as follows (after Ballentine and Burnard, 2002):

$$[^4\text{He}]_{\text{calculated}} = (12.06 [\text{U}] + 2.87 [\text{Th}]) \times 10^{-8} \times t \quad (5)$$

$$[^{40}\text{Ar}^*]_{\text{calculated}} = 3.80 \times 10^{-8} \times [\text{K}] \times t \quad (6)$$

where $[U]$ and $[Th]$ are the concentrations of U and Th in ppm, respectively; $[K]$ is the K content in wt%; t is the formation age of granites in year. Dividing the preserved (matrix-sited) He and Ar* contents by the generated He and Ar*, their apparent retention coefficients can be calculated as $L(^4\text{He})$ and $L(^{40}\text{Ar}^*)$.

The results show that $L(^4\text{He})$ is 9.93% and 1.74% for 14ZQ and 14HNP, respectively (Table 5). The lower $L(^4\text{He})$ value of 14HNP is likely due to the strong weathering and microfissures that are present around its U- and Th-rich minerals (Fig. 3). These results indicate that more than 90% of the He that has been generated since the formation of the granite has been released during the geological time. However, the values of $L(^{40}\text{Ar}^*)$ are far larger than those of $L(^4\text{He})$, as they are 71.26% and 105.81% for 14ZQ and 14HNP, respectively (Table 5). The high retention coefficients may indicate that the granites have experienced temperatures that fall above the He closure temperatures of the U- and Th-rich minerals but fall below the Ar closure temperatures of the K-rich minerals, which favours Ar retention.

Table 5 Calculated He and Ar retention coefficients

	^4He (ucm ³ STP/g)		$^{40}\text{Ar}^*$ (ucm ³ STP/g)	
	14ZQ	14HNP	14ZQ	14HNP
Preserved	115.62	7.82	14.20	29.17
Generated	1164.30	447.72	19.93	27.73
L (apparent retention coefficients)	9.93%	1.74%	71.26%	105.18%

5.3 He extracted by stepwise heating: implication of temperature on He release

Under upper crustal conditions (<150°C), He diffusion coefficients in crustal minerals of the same size range from 10^{-18} cm²/s to 10^{-22} cm²/s (Lippolt and Weigel 1988; Trull et al. 1991). In this study, the stepwise heating results suggest that the He diffusion coefficients in fine-grained granite particles ($r = \sim 100 \mu\text{m}$) are $4.46 - 4.82 \times 10^{-9}$ cm²/s and $2.00 - 3.05 \times 10^{-8}$ cm²/s at 700°C and 1100°C respectively (Table 3). Based on the approximate calculation formula proposed by Lagerwall and Zimen (1963), the time required for a grain to lose 90% of its He were calculated, which is 1 Ma for minerals under lower temperature (<150°C) when the He diffusion coefficient is approximately 10^{-18} cm²/s, and is 0.001 year (8.64 h) for granite grains ($r = \sim 100 \mu\text{m}$) at 700°C (Fig. 6), suggesting that He can be closed in minerals under lower temperatures but cannot be trapped under high temperature over geological time.

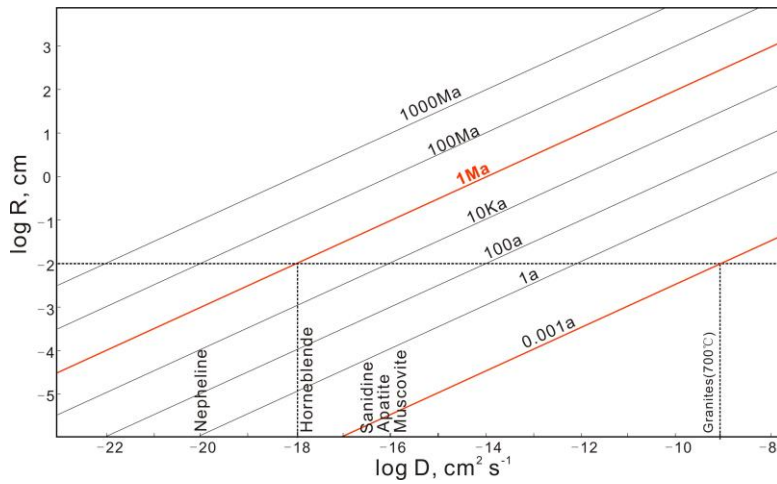


Fig.6. Relationship between grain size, He diffusion coefficient and time required for a grain to lose 90% of its He (after Ballentine and Burnard 2002).

The diffusivities of several minerals are measured at 100°C (Lippolt and Weigel 1988; McDougall and Harrison 1988; Wolf et al., 1996). The diffusivity of granites at 700°C is from this study. The horizontal dashed line indicates that the time required for a granitic grain (~100 μm) to lose 90% of its He is only 0.001 year (8.64 h) at 700°C.

In the past few decades, the influence of temperature on He retention, diffusion, and closure temperature in U- and Th-rich minerals have been researched thoroughly in the field of U-Th-He dating (Wolf et al., 1996; Farley et al., 1996; Flowers et al., 2009). In our samples, U and Th are present in independent minerals (uranothorite, thorite, minerals rich in Ti, Nb, Y and U), or isomorphously, in accessory minerals (monazite, apatite, xenotime, magnetite and zircon). Besides, other U- and Th-rich minerals, such as titanite and allanite, are also present in granites. We summarized the He closure temperatures in a variety of U and Th bearing minerals, which are concentrated between 27–250°C (Fig.7). Uranium oxide has the lowest He closure temperature (27–76°C, calculated from Roudil et al., 2008) and it can only conserve 2.1% (on average) of the He produced since its formation (Roudil et al., 2008). General minerals and U and Th-bearing minerals have variably higher He closure temperatures, which are lower than 250°C: fluorite (47–143°C, Wolff et al., 2016), carbonates (70±10°C, Copeland et al., 2007), apatite (75±7°C, Wolf et al., 1996), hematite (>90–180°C, Wernicke and Lippolt, 1997), zircon (170–190°C, Reiners, et al., 2002), titanite (191–218°C, Reiners, et al., 1999), monazite (241°C on average, Boyce et al., 2005) and magnetite (250°C, Blackburn et al., 2007). Thus, it is harder to retain He in these U- and Th-bearing minerals at temperatures above 250°C on the geological time scale.

On the other hand, the Ar closure temperatures are higher than those of He by over 100°C for general mineral grain sizes in crustal rocks (Ballentine and Burnard, 2002). For example, the Ar closure temperatures of hornblende and biotite are ~700°C and 350–400°C, respectively (Berger et al., 1981). All the evidence indicates that Ar has better retention and lower mobility than He, which can be readily explained by the larger atomic radius of Ar than that of He.

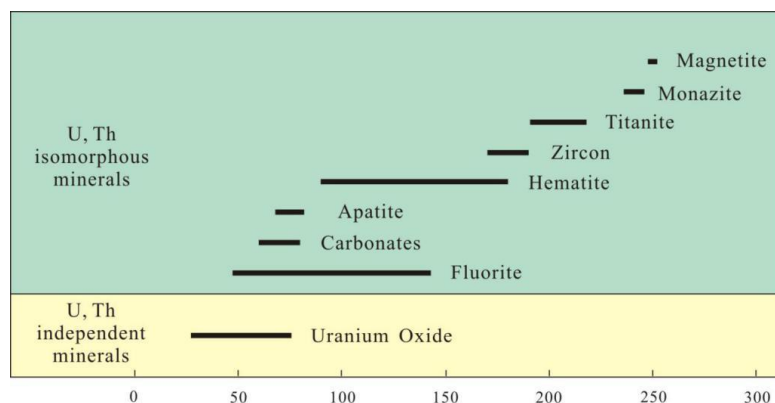


Fig.7. He closure temperatures of U-Th-bearing minerals.

Uranium oxide (27–76°C, calculated from Roudil et al., 2008); fluorite (47–143°C, Wolff et al., 2016); carbonates (70±10°C, Copeland et al., 2007); apatite (75±7°C, Wolf et al., 1996); hematite (>90–180°C, Wernicke and Lippolt, 1997); zircon (170–190°C, Reiners, et al., 2002); titanite (191–218°C, Reiners, et al., 1999); monazite (241°C on average, Boyce et al., 2005); magnetite (250°C, Blackburn et al., 2007).

5.4 Implication of fracture on He release

Many phenomena observed in nature suggest that He release is closely associated with fractures: the geothermal wells with high He concentrations are distributed along main faults in the Weihe Basin (Fig. 1, Li et al., 2011; Han et al., 2014); release of He is applied to monitor the stress state of coal seam during coal mining (Zhu et al., 2003); He is also a sensitive indicator for earthquake prediction and activity (Sano et al., 1998; Hong et al., 2010; Umeda et al., 2013; Wei et al., 2016). Since granite has no retention ability for helium under high temperature (Section 5.3), helium is mobilizable rather than closed in minerals in the depth of granitic pluton. Fractures extended to deep geological bodies can further transfer free He from deep source rocks to shallow reservoirs.

Research at Yellowstone National Park in the U.S.A has shown that He accumulated in Archaean cratonic rocks (> 2.5 billion years old) was liberated over the past two million years by intense crustal metamorphism (Lowenstern et al., 2014). Generally, crustal metamorphism is accompanied by the formation of fractures. Young active fractures can mobilize He to adjacent gas fields.

5.5 A model: granite as a Helium Source Rock

Based on the U, Th distribution and He concentrations extracted by crushing and stepwise heating obtained from this study, a model on granite as an effective helium source rock is constructed here (Fig. 8). We propose that the extensively distributed granites in the NQB are the most likely source rocks of the He resources accumulated in the Weihe Basin.

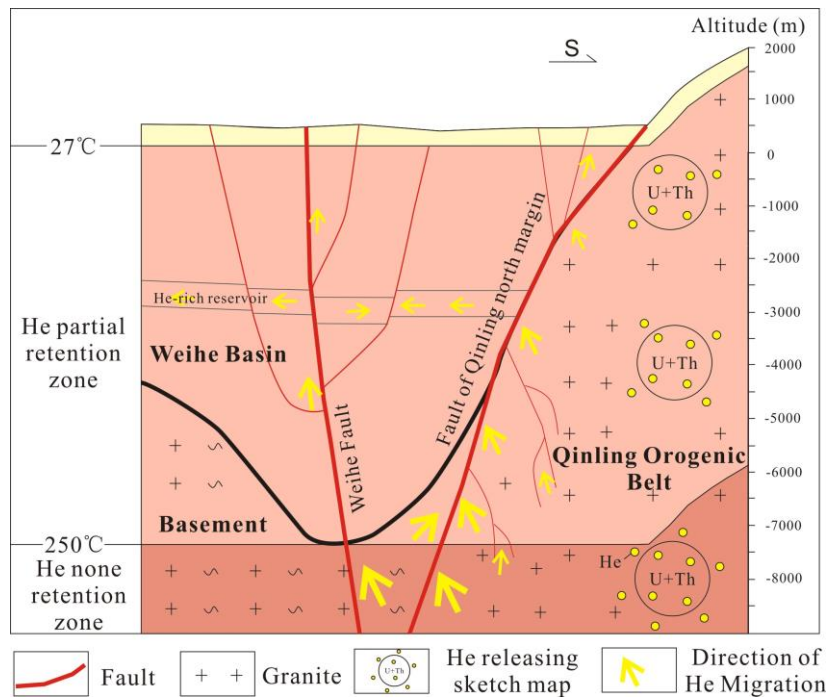


Fig.8. Model of granite as an effective helium source rock in the Weihe Basin.

(1) U- and Th-rich granites can slowly but constantly generate He; (2) Temperature has a pivotal impact on He release from granite. Part of He is retained in minerals and rocks under 27–250°C, which correspond to ~400m – ~7800m assuming a 15°C surface temperature and a 3°C/100m geothermal gradient (He partial retention zone). Geologic bodies deeper than 7800m are classified as He none retention zone. (3) Fracture system facilitates He release from the source rocks and provides channels to transfer free He from deep source rocks to shallow reservoirs.

Firstly, it has been demonstrated that granites in the NQB are rich in uranium and thorium. The [U+0.24Th] values of 7 granitic plutons from the NQB range from 7.27 to 16.82 ppm, which are 1.5–3.4 times that of the Clark value. U and Th are present in independent minerals (uranthorite, thorite, minerals rich in Ti, Nb, Y, and U), or isomorphously, in accessory minerals (monazite, apatite, xenotime, magnetite, zircon, titanite and allanite). Most of the granites in NQB were formed during the Paleozoic and Mesozoic. Based on the radioactive decay law, these granites can slowly but constantly generate radiogenic He. Results show that only 9.93% and 1.74% of the He generated since their formation are preserved in the rocks for the two samples. This suggests that more than 90% of the generated He has been released out of these granitic bodies, some of which may have been accumulated in the Weihe Basin.

Additionally, temperature has a pivotal impact on He release from granite. The He close temperatures of many U- and Th-rich minerals are concentrated between 27°C to 250°C, which correspond to ~400 – ~7800m in depth, assuming a 15°C surface temperature and a 3°C/100m geothermal gradient. This suggests that variable amounts of He can be retained in the granite at depths from 400m to 7800m (He partial retention zone), and part of it was released. At the depths of greater than 7800m, He may have been completely released, herein termed the He none retention zone. Furthermore, He diffusion experiments under high pressure for olivine

yield a similar result with those under 1-atm (Cherniak and Watson, 2012), suggesting that high pressure has little impact on He diffusion in rocks at depth. Since He mobilization is controlled by temperature, He is easier to release in regions with a high geothermal gradient.

Fractures further facilitate release of He from the source rocks under high temperatures. They provide channels to the effectively transfer free He from deep source rocks to shallow reservoirs. A fracture system consists of faults, fractures and microfissures. The migration of He in a fracture system makes the He concentrations in the fractures lower than the source rocks, further accelerating the He release process from the granites. Relatively young granitic source rocks and active fracture systems can provide large amounts of He to nearby sedimentary basins, such as the Weihe Basin.

Therefore, it is advantageous to explore He resources in hydrocarbon basins with granitic basement (or adjacent to granite), high geothermal field, and young active fractures.

6 Conclusions

The characteristics of granites as an effective helium source rock were evaluated using the granites from the Northern Qinling Belt near the Weihe Basin as an example. The following conclusions are made.

(1) The 7 granitic plutons in NQB are rich in U and Th. The [U+0.24Th] values of these granites range from 7.27 to 16.82 ppm, which are 1.5–3.4 times that of the Clark value. U and Th are present in independent minerals, or isomorphously, in accessory minerals. These granites can slowly but constantly generate radiogenic He.

(2) Temperature has a pivotal impact on He release from granite. Almost all of the He (99.24%) generated by radioactive decay was released in the stepwise heating experiments. He diffusivity (D_{He}) of granites are $4.46\text{--}4.82\times 10^{-9}$ cm²/s at 700°C and $2.00\text{--}3.05\times 10^{-8}$ cm²/s at 1100°C. The He closure temperatures of many U- and Th-rich minerals are concentrated between 27°C to 250°C, which corresponds to ~400m to ~7800 m in depth assuming a 15°C surface temperature and a 3°C/100 m geothermal gradient. It means that part of the He generated by radioactive decay is retained in minerals and rocks from 400m to 7800 m depths (He partial retention zone), and part of it is released. At the depths of greater than 7800 m, He is potentially completely mobilized (He none retention zone).

(3) Fractures facilitate release of He from source rocks under high temperatures. They provide channels for the effective transfer of the free He from deep source rocks to shallow reservoirs.

(4) It is advantageous to explore He resources in hydrocarbon basins with granitic basement (or adjacent to granite), high geothermal field, and young active fractures.

(5) Ar has better retention and lower mobility than He, which can be readily explained by the larger atomic radius of Ar than that of He. More ⁴⁰Ar (~71% and ~105%) are retained in granitic samples, far higher than He (9.93% and 1.74%). The diffusion coefficients of Ar are $1.30\text{--}1.37\times 10^{-11}$ cm²/s and $1.51\text{--}1.71\times 10^{-8}$ cm²/s at 700°C and 1100°C, respectively, lower than those of He.

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