#### **ORIGINAL PAPER**



# Mineral chemistry aspects of radioactive mineralization associated with Zr-, Nb-, and REE-bearing minerals from felsic dikes at Abu Hawis, North Eastern Desert, Egypt

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#### Abstract

The exposed rocks in Abu Hawis area, North Eastern Desert (NED), Egypt, consist of tonalite-granodiorite and monzogranite, dissected by post-granite felsic (microgranite and rhyolite) and mafic (basaltic-andesite) dikes. The investigated radioactive minerals and Zr-, Nb-, and REE-bearing minerals were restricted to felsic dikes having E–W and NE–SW trends. Uraninite, uranothorite, and thorite occur as the main radioactive minerals in microgranite dikes, while thorite is represent in rhyolite dikes. Y<sub>2</sub>O<sub>3</sub> and HREE are recorded in zircon crystals from rhyolite dikes whereas HREEs in zircon grains from microgranite dikes are below detection limit. Zircon crystals from microgranite dikes contain high values of HfO<sub>2</sub> with up to 9.08 wt % owing to the effect of hydrothermal activity. Columbite from microgranite dikes has Ta/(Ta + Nb) and Mn/(Mn + Fe) ratios ranging between 0.0052–0.0164 and 0.0549–0.7010, respectively, which point to manganocolumbite composition, except for two spots that show a ferrocolumbite composition. Fergusonite is recorded in microgranite dikes, with average values of Nb<sub>2</sub>O<sub>5</sub>, Y<sub>2</sub>O<sub>3</sub>, and HREE<sub>2</sub>O<sub>3</sub> reaching 50.3, 22.93, and 17.68 wt%, respectively. Monazite is recorded in both microgranite and rhyolite dikes, with marked enrichment of ThO<sub>2</sub>, which reaches up to 12.52 wt% in the first one, while the total  $\Sigma LREE_2O_3$  reached up to 68.5 wt% in the latter. Parisite and chevkinite are confined to rhyolite dikes with clear enrichment in LREE with averages ranging between 53.53 and 43.75 wt%  $\Sigma LREE_2O_3$ , respectively.

Keywords Abu Hawis · Egypt · Felsic dikes · E-W and NE-SW trends · Radioactive minerals · Rare metal mineralization

### Introduction

The Arabian-Nubian Shield (ANS) is composed of basement rocks of Precambrian age along the Red Sea two flanks, located in western Arabia and northeastern Africa (Johnson and Woldehaimanot 2003; Liégeois and Stern 2010). The ANS was evolved in three stages, as follows: (a) Intra-ocean ridge subduction and arc magmatism <sup>2</sup>750 Ma, (b) collisional events and terrane amalgamation (<sup>2</sup>750 Ma-<sup>6</sup>20 Ma), (c) tectonic escape and extension <sup>6</sup>620 Ma (Farahat et al. 2011; Fathy et al. 2018). It consists of a large orogenic belt that extends from western Arabia to East African Orogen (EAO) (Fig. 1a), through Saudi Arabia, Yemen,

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Ahmed E. Abdel Gawad ahm.elsay@hotmail.com; gawadnma@gmail.com Oman, Jordan, Egypt, Sudan, Eritrea, Ethiopia, and Somalia, whereas the southern belt occupies Mozambique segment.

Rare metal granites (RMG) are those enriched in one or more mineralization of the elements Zr, Nb, Ta, Rb, Cs, W, Sn, Li, F, Be, U, Th, REEs, and Y. RMG can be attributed either to magmatic and/or metasomatic processes (Abdalla et al. 1998; Gaafar et al. 2014; Dessouky et al. 2020; Ali et al. 2021; Abdel Gawad 2021). RMG intrusion occurred between 530 and 620 Ma during post-orogenic magmatism in the Egyptian Nubian Shield (Abdel Karim and Sos 2000; Moussa et al. 2008; Eliwa et al. 2014; Ali 2015; Skublov et al. 2021; Abdel Gawad et al. 2021a). However, rare metal mineralizations are mainly restricted to granites, pegmatites, acidic volcanics, and mylonite rocks as well as quartz and jasper veins widely distributed in the Eastern Desert of Egypt (Fig. 1b) (Saleh 2007; Raslan et al. 2010; Abdel Wahed et al. 2012; Dawoud et al. 2018; Heikal et al. 2001; 2019; Fawzy et al. 2020; Ghoneim et al. 2020; Surour and Omar 2020; Dessouky et al. 2020; Abdel Gawad et al.

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Fig. 1 a Geologic map of the Arabian Nubian Sheild (ANS); b geologic map of the basement rocks of Neoproterozoic age, Eastern Desert (ED), Egypt (Liégeois and Stern 2010)



2021b, c; Mahdy 2021; Surour 2021; Abdellah et al. 2021; Abdel Gawad et al. 2022; Surour and Omar 2022).

The need of rare metals has amplified intensely in the last few decades. Many of rare metals represent strategic

economies, which are essential in the high technology, electronics, automobile industries, and renewable energy applications (Bourzac 2001; Eggert et al. 2010; Hoatson et al. 2011; Linnen et al. 2012).

Rare earth elements (REE) are considered as critical and vital material to future development. They are having a great importance, such as cell phones, computer memories, DVDs, rechargeable batteries, cerium oxide used for polishing gemstones such as granite and marble as well as glass, medical devices, high-efficiency motors in hybrid electric vehicles, catalytic converters, metallurgy, FeB magnets, wind power turbines, catalysts, and metal alloys as well as high demand for green energy and military defense systems (German and Elderfield 1990; Mariano and Mariano 2012; Gambogi 2013; Zepf 2013; Liu et al. 2013). Life demand is expected to increase dramatically over the next 10 years, without increase in the total REE reserves, thus highlighting the need to exploit for more REE deposits to be discovered (Hoatson et al. 2011; Jaireth et al. 2014). According to the high economical use of REEs, global request for REE will rise continuously, which will put great pressure on the current REE supply chain.

Several studies were carried out on Abu Hawis area (NED), Egypt (Ahmed and Moharem 2002; Al-Boghdady et al. 2005; Ali 2007; El-Bialy and Omar 2015; Dawoud et al. 2017). This paper presents in terms of novel data and interpretations about the mineralization especially U-, Th-, Zr-, Nb-, and REE-bearing minerals from the studied felsite dikes at Abu Hawis. To achieve this goal, micro-chemical analyses of these mineralization were first carried out in

order to obtain informative data about radioactive minerals as well as Zr-, Nb-, and REE-bearing minerals.

## **Geologic setting**

Field evidence shows that the exposed rock types are represented by tonalite-granodiorite, and monzogranite, which are cut by post-granite dikes and quartz veins (Figs. 2 and 3a–e).

The exposed rocks of Abu Hawis area are mostly composed of tonalite-granodiorite that have a huge batholith extending along the border of Qena–Safaga Road. They are medium- to coarse-grained rocks, form low relief, and vary from gray to whitish-gray colors. The pluton outcrops of these granitoids show marked intensive fracturing, jointing, exfoliation, and oval-like shapes, owing to the effect of weathering processes (Fig. 3f). They are strongly foliated and metamorphosed and occur as gneissose tonalite-granodiorite. Meanwhile, the other parts of tonalite-granodiorite occur as massive rocks. They are presented as smallsized plutons, without any deformed signatures. They are composed of quartz, plagioclase, K-feldspars, biotite, and hornblende.

The monzogranite pluton possesses circular to elliptical outlines and intrudes into tonalite-granodiorite of the studied area (Fig. 3f). It is frequently forming hard massive rocks,



Fig. 2 Geologic map of Abu Hawis area, North Eastern Desert, Egypt (Ali 2007; El-Bialy and Omar 2015; Dawoud et al. 2017)

high relief, pink to reddish brown in color, and medium- to coarse-grained. It is mainly composed of K-feldspar, quartz, plagioclase, muscovite, and biotite. Granitoids were dissected by basaltic-andesite dikes and quartz veins, striking in E–W, NW–SE, NE–SW, and N–S directions (Figs. 2 and 3d, e).

Pegmatites occur as pockets and/or dike-like forms. They are very coarse-grained, red, buff to reddish brown in color, and essentially composed of K-feldspar, quartz, plagioclase, biotite, and muscovite. They occur in the latest magmatic stages in Abu Hawis monzogranite and/or along the contacts between tonalite-granodiorite and monzogranite. Some exposures of these rocks are affected by hematitization and considered as good sources of strategic rare metal mineralization.

The most significant U mineralization occupies about 5 to 10 m width and 500 m length, subvertical, and highly altered basic dikes, striking in the NE–SW direction along the main shear zone of Abu Hawis plutons. Argillization, hematitization, and silicification are the main alterations affecting the sheared basic dikes. Relatively, high radio-active anomalies recorded in the main shear zone reached

**Fig.3 a** Microgranite dikes having E–W trend cross cut tonalitegranodiorite; **b** NE–SW strike-slip fault (right lateral) cross cut E–W microgranite dikes; **c** rhyolite dikes having NE–SW cross cut tonalite-granodiorite; **d** and esite dikes having NW–SE trend cross

cut tonalite-granodiorite; **e** basaltic dikes having N–S trend cross cut tonalite-granodiorite; **f** tonalite-granodiorite rocks were intruded by the monzogranite (oval shape) with sharp intrusive contacts



about 80 ppm for eU and about 45 ppm for eTh, with an eU/eTh ratio reaching about 1.75 (Ammar et al. 2007).

Felsite dikes are cut through the tonalite-granodiorite. They range between 1 and 6 m width and are buff, pale pink to reddish brown color, are mostly fine-grained textures, and dominated by microgranite and rhyolite dikes. These dikes are mainly composed of quartz, K-feldspars, and plagioclase with mica as well as iron oxides. They are striking in the E–W and NE–SW directions with steeply dipping or nearly vertical (Fig. 3a–c). These dikes were jointed and fractured, highly altered, and affected by hematitization in addition to dendritic manganese oxides.

#### Materials and analytical methods

Twelve samples were selected and prepared in order to identify the petrographic characteristics of felsite dikes. Thin polished sections and slaps were prepared for microchemical analyses in order to investigate the rare metal mineralization especially in microgranite and rhyolite dikes. Mineral chemistry was obtained using an electron microprobe equipped with CAMECA SX 100, using five automated wavelength-dispersive spectrometers (WDS) and an energy-dispersive spectrometer (EDS). All performed analyses of minerals were made using the following conditions: the diameter of the electron beam on the specimen surface is under 1 µm, an acceleration voltage 15 kV, beam current 15 nA, counting time 10 s for peak, and 5 s for background. The standards used for calibration include natural standards such as albite for Al (K $\alpha$ ), orthoclase for Si (K $\alpha$ ), wollastonite for Ca (K $\alpha$ ), hematite for Fe (K $\alpha$ ), olivine for Mg (K $\alpha$ ), and monazite for Yb  $(L\alpha)$ , whereas the synthetic compounds UO<sub>2</sub>, ThO<sub>2</sub>, ZrO<sub>2</sub>, HfO<sub>2</sub>, PbCrO<sub>4</sub>, YPO<sub>4</sub>, LaPO<sub>4</sub>, CePO<sub>4</sub>, PrPO<sub>4</sub>, NdPO<sub>4</sub>, SmPO<sub>4</sub>, EuRu<sub>2</sub>Ge<sub>2</sub>, GdTiGe, TbPO<sub>4</sub>, DyRu<sub>2</sub>Ge<sub>2</sub>, HoP<sub>5</sub>O<sub>4</sub>, ErNi<sub>2</sub>Si<sub>2</sub>, TmPO<sub>4</sub>, YbPO<sub>4</sub>, and LuPO<sub>4</sub> are used as standards for U (M $\beta$ ), Th (M $\alpha$ ), Zr (L $\alpha$ ), Hf (M $\alpha$ ), Pb (M $\alpha$ ), Y (L $\alpha$ ), La (L $\alpha$ ), Ce (L $\alpha$ ), Pr(L $\beta$ ), Nd (L $\alpha$ ), Sm (L $\alpha$ ), Eu  $(L\alpha)$ , Gd  $(L\alpha)$ , Tb  $(L\alpha)$ , Dy  $(L\alpha)$ , Ho  $(L\beta)$ , Er  $(L\alpha)$ , Tm  $(L\alpha)$ , Yb  $(L\alpha)$ , and Lu  $(L\alpha)$  respectively. Pure metals of Nb and Ta were used for Nb-Ta-bearing minerals.

For an argillic sample, heavy liquids separation technique using bromoform of specific gravity 2.85 gm/cm<sup>3</sup> was used to concentrate the clay minerals. The clay minerals were picked under a binocular microscope to obtain pure mineral samples for X-ray diffraction (XRD) investigation. The X-ray diffraction (XRD) technique was used to identify the clay mineral sample using a PHILIPS PW 3710/31 diffractometer. These analyses were carried out in the laboratories of the Nuclear Materials Authority (NMA) of Egypt.

#### Results

#### Petrography of felsic dikes

#### Microgranite

Microscopically, microgranite shows equigranular texture and is composed mainly of K-feldspar, quartz, plagioclase, and muscovite (Fig. 4a). Zircon, pyrite, galena, sphalerite, and iron oxides are the most common accessory minerals, while sericite and kaolinite are the main alteration products. K-feldspar consists of microcline- and orthoclase microperthites. Microcline microperthite occurs as subhedral to euhedral crystals showing cross-hatched twinning (Fig. 4a). Orthoclase microperthite occurs as anhedral to subhedral crystals having simple twinning. Quartz occurs as interstitial grains between feldspar crystals as anhedral to subhedral crystals and fine grained. Plagioclase occurs as euhedral to subhedral tabular crystals. It exhibits simple and lamellar twinning (Fig. 4a, d). Muscovite occurs as anhedral crystals, corroded peripheries, and encloses opaques. Zircon crystals are coated by iron oxides, and occur as prismatic crystals enclosed in quartz and feldspars (Fig. 4b-d). Microgranite affected by hydrothermal alterations include kaolinization. chloritization, and hematitization (Fig. 4a-f).

#### Rhyolite

Rhyolite dikes are fine-grained and buff to reddish brown color and composed mainly of K-feldspars, quartz, and muscovite embedded in fine groundmass, showing porphyritic and spherulitic textures (Fig. 4g–j). Accessories are zircon, iron oxides, and opaque minerals, whereas sericite and kaolinite occur as secondary minerals. K-feldspar occurs either as subhedral to anhedral phenocrysts of orthoclase or as finegrained groundmass in the matrix. Kaoline and sericite are the common alteration products of K-feldspar. Quartz occurs as subrounded to subangular phenocrysts and shows undulose extinction. Muscovite occurs as anhedral crystals and encloses zircon and iron oxides (Fig. 4g, h). Both kaolinization and hematitization were affected rhyolite (Fig. 4h–j).

#### **Mineral chemistry**

Uraninite, thorite, uranothorite, zircon, columbite, fergusonite, monazite, and parisite as well as pyrite, galena, sphalerite, apatite, and fluorite are well investigated from Abu Hawis felsic dikes, by using an electron probe microanalyses (EPMA) technique, as shown on Figs. 5–9 and Tables 1–5. Fig. 4 Photomicrographs in felsic dikes of Abu Hawis area, North Eastern Desert, Egypt. a Microgranite is composed of plagioclase, microcline, quartz, and muscovite in addition to zircon and opaques, crossed nicols (C.N.); b-d zircon crystals are coated by iron oxides in microgranite (C.N.); e zircons surrounded by chlorite and iron oxides in microgranite (C.N.); f hematitization affected microgranite (C.N.); g muscovite in rhyolite (C.N.); h muscovite encloses zircon and iron oxides (hematitization) in rhyolite (C.N.); i spherulitic texture in rhyolite (C.N.); j K-feldspar shows porphyritic texture in rhyolite (C.N.). Abbreviations: plg, plagioclase; mic, microcline; K-fs, K-feldspar; ms, muscovite; zr, zircon; qz, quartz; ir, iron; sp, spherulitic



Fig. 5 SEM-BSE images of U- and Th-bearing minerals associated with other accessory minerals from felsic dikes, Abu Hawis area, North Eastern Desert, Egypt. a Micro-inclusions of uraninite, uranothorite, monazite, and opaques enclosed in large zircon crystals in microgranite; b fine thorite grains occurring along the peripheries of zircon in microgranite; c fine grained of thorite associated with parisite and fluorite in rhyolite; d thorite occurring along the peripheries of parasite in rhyolite. Abbreviations: ura, uraninite; uth, uranothorite; thr, thorite; Zir, zircon; mon, monazite; par, parisite; flu, fluorite; apt, apatite; qz, quartz; ilm, ilmenite; hem, hematite



#### **Radioactive minerals**

Uraninite is considered the main radioactive U-bearing mineral, while thorite and uranothorite are the main radioactive Th-bearing minerals in the studied microgranite and rhyolite dikes (Fig. 5a–d and Table 1).

Uraninite is recorded in peraluminous granites and their pegmatites as well as dolerite dikes (Gaafar et al. 2014; Ali et al. 2021; Ghoneim et al. 2021). It occurs as fine micro-inclusions that are enclosed in zircon crystals (Fig. 5a). Uraninite composed of UO<sub>2</sub> reached 69.1 and 69.57 wt%. PbO, Y<sub>2</sub>O<sub>3</sub>, and ThO<sub>2</sub> are present in uraninite and their contents reached up to 6.85, 8.2, and 4.02 wt%, respectively. Electron probe micro-analyzer (EPMA) data show that uraninite contains lower contents of SiO<sub>2</sub> and HREE, especially Dy<sub>2</sub>O<sub>3</sub>, Er<sub>2</sub>O<sub>3</sub>, and Yb<sub>2</sub>O<sub>3</sub> (Table 1). Thorite is a radioactive mineral. It is one of the most common Th-silicate minerals in felsic dikes of the area under investigation. It is enclosed as fine bright microinclusions in zircon crystals and shows intergrowth with zircon along its peripheries (Fig. 5b). It occurs as fine individual grains, subhedral to anhedral crystals (Fig. 5c). Other minute grains of thorite occur adjacent to parisite crystals (Fig. 5d). The chemical composition of thorite shows that the means of ThO<sub>2</sub> concentrations attain 57.89 and 61.69 wt% in both microgranite and rhyolite dikes, respectively. The means of SiO<sub>2</sub> contents reach 18.76 and 17.11 wt% in thorite from microgranite and rhyolite dikes, respectively. Thorite in microgranite dikes shows high mean values of UO<sub>2</sub>,  $Y_2O_3$ , and  $ZrO_2$  reaching 3.05, 3.39, and 0.51 wt%, when compared with thorite from rhyolite dikes.  $Dy_2O_3$ ,  $Er_2O_3$ , and  $Yb_2O_3$  were recorded as minor amounts in the studied thorite, but with higher mean contents in microgranite than those in rhyolite dikes (Table 1). CaO,  $P_2O_5$ , and  $Fe_2O_3$  were recorded as minor constituents in the analyzed thorite (Table 1).

Uranothorite is a common Th-silicate mineral in microgranite dikes. It occurs as subrounded, subhedral grains, and micro-inclusions in zircon crystals (Fig. 5a). According to EPMA, the chemical composition of uranothorite reveals that ThO<sub>2</sub> concentration ranges between 49.89 and 51.17 wt%, UO<sub>2</sub> from 27.01 to 27.42 wt%, and SiO<sub>2</sub> from 17.25 to 18.04 wt% (Table 1). Small amounts of PbO, CaO, P<sub>2</sub>O<sub>5</sub>, Y<sub>2</sub>O<sub>3</sub>, ZrO<sub>2</sub>, and Fe<sub>2</sub>O<sub>3</sub> were detected, with means reaching 0.32, 0.89, 0.79, 0.33, 0.65, and 0.44 wt%, respectively.

#### Zircon

Zircon occurs as prismatic, subhedral to anhedral crystals, and varies from fine-grained to large crystals, reaching up to 200  $\mu$ m in microgranite and rhyolite. It is associated with columbite, fergusonite, barite, rutile, ilmenite, hematite, and opaques (Fig. 6a–d). The presence of fine-grained zircon as micro-inclusion in large zircon crystals indicates the presence of two zircon generations, which are encountered in the studied microgranite dikes.

 Table 1
 Representative EPMA data (oxides, in wt%) of U and Th minerals from felsic dikes, Abu Hawis area, North Eastern Desert, Egypt

Min. in dike/oxid	Uraninite in microgranite		Thorite	Thorite in microgranite			Aver	Thorite in rhyolite			Aver	Uranot microg	horite in ranite		
SiO <sub>2</sub>	0.83	0.71	18.97	17.13	18.72	20.22	18.76	16.64	17.45	16.92	17.43	17.11	18.04	17.61	17.25
UO <sub>2</sub>	69.1	69.57	4.32	5.33	1.28	1.28	3.05	1.27	1.23	1.08	1.22	1.20	27.42	27.21	27.01
ThO <sub>2</sub>	3.19	4.02	56.72	57.38	58.93	58.51	57.89	61.67	61.32	62.59	61.18	61.69	50.17	49.95	49.89
PbO	6.85	6.83	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	0.35	0.32	0.29
CaO	n.d	n.d	1.41	0.99	1.24	1.45	1.27	0.98	1.51	1.19	1.57	1.31	1.03	0.87	0.76
$P_2O_5$	n.d	n.d	1.22	1.18	0.92	1.28	1.15	1.37	1.03	1.83	1.67	1.48	0.82	0.78	0.77
$Y_2O_3$	8.2	7.66	3.57	3.24	3.18	3.58	3.39	3.46	4.12	2.43	3.19	3.30	0.35	0.32	0.31
$ZrO_2$	n.d	n.d	0.25	0.58	0.97	0.24	0.51	0.63	0.26	0.21	0.22	0.33	0.71	0.67	0.58
Fe <sub>2</sub> O <sub>3</sub>	n.d	n.d	1.78	2.43	1.46	1.62	1.82	2.46	1.95	1.99	1.73	2.03	0.43	0.46	0.44
Dy <sub>2</sub> O <sub>3</sub>	2.72	2.78	0.62	0.58	0.65	0.63	0.62	0.57	0.45	0.53	0.61	0.54	n.d	n.d	n.d
Er <sub>2</sub> O <sub>3</sub>	3.24	3.08	0.55	0.45	0.44	0.51	0.49	0.49	0.47	0.48	0.51	0.49	n.d	n.d	n.d
Yb <sub>2</sub> O <sub>3</sub>	3.52	2.9	2.84	0.98	3.13	0.75	1.93	0.72	0.78	1.12	0.73	0.84	n.d	n.d	n.d
Total	97.65	97.55	92.25	90.27	90.92	90.07	90.88	90.26	90.57	90.37	90.06	90.32	99.32	98.19	97.30
Cation															
Si	0.04	0.03	0.99	0.94	1.00	1.04	0.99	0.91	0.95	0.92	0.94	0.93	0.96	0.95	0.94
U	0.66	0.67	0.05	0.06	0.02	0.01	0.04	0.02	0.01	0.01	0.01	0.01	0.32	0.33	0.33
Th	0.03	0.04	0.67	0.71	0.71	0.69	0.70	0.77	0.76	0.78	0.75	0.76	0.60	0.61	0.62
Pb	0.08	0.08											0.00	0.00	0.00
Ca			0.08	0.06	0.07	0.08	0.07	0.06	0.09	0.07	0.09	0.08	0.06	0.05	0.04
Р			0.05	0.05	0.04	0.06	0.05	0.06	0.05	0.08	0.08	0.07	0.04	0.04	0.04
Y	0.19	0.18	0.10	0.09	0.09	0.10	0.10	0.10	0.12	0.07	0.09	0.10	0.01	0.01	0.01
Zr			0.01	0.02	0.03	0.01	0.01	0.02	0.01	0.01	0.01	0.01	0.02	0.02	0.02
Fe			0.07	0.10	0.06	0.06	0.07	0.10	0.08	0.08	0.07	0.08	0.02	0.02	0.02
Dy	0.04	0.04	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01			
Er	0.044	0.042	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01			
Yb	0.05	0.04	0.05	0.02	0.05	0.01	0.03	0.01	0.01	0.02	0.01	0.01			
∑cations	1.12	1.11	2.08	2.07	2.08	2.07	2.08	2.07	2.09	2.06	2.07	2.07	2.03	2.03	2.02

Min., mineral; oxid., oxides; Aver., average

Calculated chemical formulae are based on 2 oxygen atoms (apfu) for uraninite

Calculations are based on 4 oxygen atoms per formula unit (apfu) for thorite and uranothorite

n.d., not determined

According to EPMA (Table 2), zircon crystals from microgranite dikes are composed of  $ZrO_2$  that ranges from 56.06 to 62.76 wt% and a mean reaching 60.09 wt%, SiO<sub>2</sub> from 30.76 to 34.29 wt% and a mean reaching 32.19 wt%, and HfO<sub>2</sub> from 1.03 to 9.08 wt% and a mean reaching 3.81 wt%. HfO<sub>2</sub> shows higher concentrations in the analyzed zircon crystals, reaching to 7.41, 9.08, and 6.12 wt% (Table 2). ThO<sub>2</sub> and UO<sub>2</sub> were recorded in zircon crystals and show low concentrations reaching up to 2.16 and 0.54 wt%, respectively. Only Yb<sub>2</sub>O<sub>3</sub> from HREE is recorded with low contents. Minor amounts of CaO, Fe<sub>2</sub>O<sub>3</sub>, and Al<sub>2</sub>O<sub>3</sub> were also recorded (Table 2).

On the other hand, EPMA data show that zircon crystals from rhyolite dikes are composed essentially of  $ZrO_2$ , ranging from 58.46 to 59.21 wt% and a mean reaching 58.72 wt%, SiO<sub>2</sub> from 29.73 to 30.29 wt% and a mean attaining

30.0 wt%, and  $HfO_2$  from 0.98 to 3.09 wt% and a mean reaching 1.59 wt%. ThO<sub>2</sub> and UO<sub>2</sub> display low concentrations in the analyzed zircon crystals with mean values of 0.74 and 0.85 wt%, respectively. HREE (Dy<sub>2</sub>O<sub>3</sub>, Er<sub>2</sub>O<sub>3</sub>, and Yb<sub>2</sub>O<sub>3</sub>) and Y<sub>2</sub>O<sub>3</sub> were recorded in the analyzed zircon crystals, with mean values reaching 0.54, 0.63, 1.06, and 4.03, respectively (Table 2). CaO and Fe<sub>2</sub>O<sub>3</sub> were also recorded in zircon crystals as minor constituents.

#### Nb-rich minerals

Columbite occurs as anhedral to subhedral crystals, up to 100  $\mu$ m in microgranite, commonly enclosed in plagioclase, K-feldspar, and quartz crystals. Other columbite crystals are dispersed as fine- to medium-grained crystals enclosed in fergusonite (Fig. 6c). Some crystals of Fig. 6 SEM-BSE images of Zr- and Nb-bearing minerals, associated with other accessory minerals from felsic dikes, Abu Hawis area, North Eastern Desert, Egypt. a Faint zoning of zircon enclosing microinclusions of zircon, barite, and opaques in microgranite; b micro-inclusions of fergusonite enclosing zircon in microgranite; c fergusonite enclosing columbite, adjacent to zircon in microgranite; d zircon crystals associated with rutile, ilmenite, and hematite in rhyolite; e micro-inclusions of galena enclosed in a large crystal of columbite that is associated with fergusonite in microgranite; f columbite and fergusonite adjacent to the peripheries of ilmenite in microgranite. Abbreviations: zir, zircon; bar, barite; fer, fergusonite; col, columbite; pyt, pyrite; gal, galena; ilm, ilmenite; hem, hematite; rut, rutile



columbite enclose micro-inclusions of galena and opaque minerals (Fig. 6e). EPMA analyses reveal that the typical composition of manganocolumbite, according to the quadrilateral diagram of Černý and Ercit (1985), except two spots is ferrocolumbite composition (Fig. 7). EPMA data reveal that manganocolumbite is composed of Nb<sub>2</sub>O<sub>5</sub>, which ranges between 65.26 and 73.34 wt% with a mean value of 73.24 wt%. MnO shows high concentrations that reached 14.95 wt%, with marked lower contents in two spots 1.1 and 1.52 wt%. Meanwhile, Fe<sub>2</sub>O<sub>3</sub> reached up to 18.11 wt%. Y<sub>2</sub>O<sub>3</sub>, Ta<sub>2</sub>O<sub>5</sub>, TiO<sub>2</sub>, and CaO are represented in low concentrations, with means attaining 0.53, 1.48, 2.88, and 028, respectively. UO<sub>2</sub> and ThO<sub>2</sub> have quite low concentration in the analyzed samples, reaching up to 1.43 and 1.75 wt%, respectively.

Ta/(Ta + Nb) ratios of columbite range from 0.0052 to 0.0164, whereas Mn/(Mn + Fe) ratios show a marked wide range, from 0.0549 to 0.7010. These two ratios

could indicate enrichment of manganese in the analyzed manganocolumbite composition except the two spots enriched in iron (Table 3 and Fig. 7).

Fergusonite occurs as subhedral to anhedral and finegrained, while some crystals are strongly deformed and corroded along their peripheries. It is enclosed in zircon crystals as micro-inclusions and occurs along the peripheries of columbite as thin inclusions (Fig. 6b, f). Other fergusonite crystals poikilitically enclose columbite (Fig. 6c). EPMA analyses of fergusonite show that Nb<sub>2</sub>O<sub>5</sub> ranges between 48.6 and 51.95 wt%, a mean of 50.3 wt%, and Y<sub>2</sub>O<sub>3</sub> between 22.13 and 23.64 wt%, a mean of 22.93 wt%. It represents one of the main resources which carry heavy rare earth elements, where the total HRE<sub>2</sub>O<sub>3</sub> (Gd<sub>2</sub>O<sub>3</sub>-Lu<sub>2</sub>O<sub>3</sub>) ranges between 16.2 and 19.08 wt%, a mean of 17.68 wt%. Ta<sub>2</sub>O<sub>5</sub> and TiO<sub>2</sub> show low concentration, with means attaining 0.77 and 1.02 wt%, respectively. UO<sub>2</sub> and ThO<sub>2</sub> reached up to 3.82 and 2.94

Table 2 Representative EPMA data (oxides, in wt%) of zircon from felsic dikes, Abu Hawis area, North Eastern Desert, Egypt

Min. in dike/oxid	Zircon	in microg	ranite			Aver	Zircon in rhyolite				Aver		
SiO <sub>2</sub>	31.27	31.02	32.97	30.76	31.41	33.64	34.29	32.19	30.24	29.73	30.29	29.75	30.00
ZrO <sub>2</sub>	60.44	61.21	62.53	56.06	60.46	57.19	62.76	60.09	59.21	58.46	58.64	58.56	58.72
HfO <sub>2</sub>	7.41	2.28	1.73	9.08	6.12	2.26	1.26	4.31	3.09	1.23	1.06	0.98	1.59
CaO	n.d	1.29	0.89	2.04	n.d	2.52	n.d	1.69	1.24	n.d	n.d	0.92	1.08
Fe <sub>2</sub> O <sub>3</sub>	n.d	0.72	0.77	0.63	0.47	1.32	0.86	0.80	1.12	1.54	0.86	1.44	1.24
$Al_2O_3$	n.d	0.86	0.34	n.d	0.38	0.61	n.d	0.55	n.d	n.d	n.d	n.d	n.d
ThO <sub>2</sub>	0.34	2.16	0.33	0.24	0.41	1.10	0.35	0.70	1.19	0.49	0.54	0.75	0.74
UO <sub>2</sub>	0.54	0.46	0.38	0.53	0.39	0.42	0.43	0.45	0.82	0.87	0.92	0.77	0.85
Y <sub>2</sub> O <sub>3</sub>	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	1.31	5.09	5.12	4.59	4.03
$Dy_2O_3$	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	0.45	0.59	0.62	0.48	0.54
Er <sub>2</sub> O <sub>3</sub>	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	0.31	0.75	0.77	0.67	0.63
Yb <sub>2</sub> O <sub>3</sub>	n.d	n.d	n.d	0.63	0.36	0.58	n.d	n.d	1.02	1.07	1.18	0.95	1.06
Total	100	100	99.94	99.97	100	99.64	99.95	99.93	100	99.82	100	99.86	99.92
Cation	Calcula	ted chemi	cal formu	lae are ba	sed on 4 c	oxygen ato	ms (apfu)						
Si	0.99	0.97	1.01	0.99	0.99	1.03	1.04	0.99	0.96	0.95	0.97	0.95	0.95
Zr	0.93	0.93	0.93	0.88	0.93	0.85	0.93	0.90	0.92	0.91	0.91	0.91	0.91
Hf	0.07	0.02	0.02	0.08	0.06	0.02	0.01	0.04	0.03	0.01	0.01	0.01	0.01
Ca	0.00	0.04	0.03	0.07	0.00	0.08	0.00	0.06	0.04	0.00	0.00	0.03	0.04
Fe	0.00	0.02	0.02	0.02	0.01	0.03	0.02	0.02	0.03	0.04	0.02	0.03	0.03
Al	0.00	0.03	0.01	0.00	0.01	0.02	0.00	0.02	0.00	0.00	0.00	0.00	0.00
Th	0.00	0.02	0.00	0.00	0.00	0.01	0.00	0.00	0.01	0.00	0.00	0.01	0.01
U	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.01	0.01	0.01	0.01
Y	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.02	0.09	0.09	0.08	0.07
Dy	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.01	0.00	0.01
Er	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.01	0.01	0.01
Yb	0.00	0.00	0.00	0.01	0.00	0.01	0.00	0.00	0.01	0.01	0.01	0.01	0.01
$\sum$ cations	2.00	2.03	2.02	2.04	2.01	2.06	2.00	2.04	2.04	2.04	2.03	2.05	2.05

Min, mineral; oxid, oxides; Aver., average; n.d., not determined

wt%, respectively. Minor amounts of CaO, MnO,  $Fe_2O_3$ , and SiO<sub>2</sub> were recorded (Table 3). The chemical formula of fergusonite is  $A^{3+}B^{5+}O_4$ . A-site is dominated by Y and REE, whereas B-site is occupied by Nb, Ta, and Ti. Other cations could occur in A-site. The calculated chemical formula of fergusonite is mentioned as follows:

$$\begin{split} & \left(Y_{0.53}Gd_{0.01}Dy_{0.04}Ho_{0.01}Er_{0.06}Tm_{0.02}Yb_{0.08}Lu_{0.01}U_{0.02}Fe_{0.04}Ca_{0.04}Th_{0.02}\;\right) \\ & \left(Nb_{0.99}\;Ta_{0.01}Ti_{0.03}\right)O4 \end{split}$$

#### **REE minerals**

A. Monazite-Ce is a common REE-phosphatic mineral. It is always present in the studied felsic dikes. It occurs as anhedral to subhedral, fine-grained, micro-veinlets, and is observed as micro-inclusions in zircon and parisite crystals (Figs. 5a and 8a, b). The chemical composition of monazite using EPMA is shown on Table 4. EPMA data of the analyzed monazite from the microgranite reveal that  $P_2O_5$ ranges from 26.66 to 30.98 wt% and a mean reaching 29.35 wt%. The total REE oxides oscillate from 55.97 to 60.19 wt% and a mean reaching 58.38 wt%. Ce<sub>2</sub>O<sub>3</sub> is often a common REE that varies from 28.76 to 31.95 wt% and a mean reaching 30.42 wt%, which designates monazite-Ce. The abundance of Ce<sub>2</sub>O<sub>3</sub> is followed by Nd<sub>2</sub>O<sub>3</sub>, La<sub>2</sub>O<sub>3</sub>, Pr<sub>2</sub>O<sub>3</sub>, and Sm<sub>2</sub>O<sub>3</sub>, with a mean attaining values 11.19, 9.17, 4.35, and 2.05 wt%, respectively. Gd<sub>2</sub>O<sub>3</sub> is the only heavy rare earth element, recorded in monazite, reaching up to 1.37 wt%. Monazite grains are enriched in ThO<sub>2</sub> concentrations ranging from 10.11 to 11.98 wt% and a mean reaching 11.29 wt%. SiO<sub>2</sub>, CaO, and Fe<sub>2</sub>O<sub>3</sub> occur in minor concentrations (Table 4).

On the other hand, EPMA data show that the chemical composition of monazite crystals from rhyolite is composed mainly of  $P_2O_5$  that ranges from 28.48 to 29.32 wt% and a mean reaching 29.08 wt%. The total  $\sum REE_2O_3$  content ranges from 67.82 to 68.50 wt% and a mean reaching 68.14



**Fig. 7** Quadrilateral diagram according to Černý and Ercit (1985) illustrating the chemical composition of columbite-tantalite from the microgranite, Abu Hawis area, North Eastern Desert, Egypt

wt%. Ce<sub>2</sub>O<sub>3</sub> is the most abundant REE element. It ranges from 32.24 to 33.06 wt% and a mean reaching 32.62 wt%, followed by La<sub>2</sub>O<sub>3</sub> with a mean attaining 15.85 wt%, then Nd<sub>2</sub>O<sub>3</sub> with a mean attaining of 14.17 wt%, Pr<sub>2</sub>O<sub>3</sub> with a mean reaching 3.12 wt%, and Sm<sub>2</sub>O<sub>3</sub> with a mean attaining 1.60 wt%. Gd<sub>2</sub>O<sub>3</sub> is the only recorded HREE in the analyzed monazite, with a mean reaching 0.79 wt%. ThO<sub>2</sub> shows low concentration in the monazite grains, with a mean of 1.53 wt%. SiO<sub>2</sub>, CaO, and Fe<sub>2</sub>O<sub>3</sub> are present at lesser concentrations, with mean values of 0.22, 0.31, and 0.35 wt%, respectively (Table 4).

The chemical formula of monazite could be expressed as follows: (Ce, La, Nd, Sm, Th)PO<sub>4</sub>. The fundamental chemical formulae of average monazite grains in the microgranite and the rhyolite are as follows:  $(Ce_{0.44}Nd_{0.16}La_{0.14}Pr_{0.06}Sm_{0.03}Gd_{0.02}Th_{0.10})(P_{0.99}Si_{0.01})O_4$ and  $(Ce_{0.47}Nd_{0.20}La_{0.23}Pr_{0.05}Sm_{0.02}Gd_{0.01}Th_{0.01})(P_{0.98}Si_{0.01})O_4$ , respectively. It is clear that the analyzed monazite shows a predominance of Ce, up to (0.47 apfu), over the other REEs designating monazite-Ce.

B. Parisite-(Ce) is always present only in rhyolite. It is a calcium REE-fluorocarbonate mineral and occurs as anhedral to subhedral large grains up to 100  $\mu$ m (Figs. 5d and 8b, c). Some corroded parisite crystals are in parallel intergrowth with monazite. Parisite occurs in association with fluorite, hematite, and thorite while the latter mineral, in some cases, occurs along the peripheries of parisite crystals (Fig. 5d). From the EMP analyses (Table 5), parisite shows a strong enrichment in LREE. It is essentially composed of CaO that ranges from 12.84 to 14.87 wt%, with a mean of 13.8 wt%. The total  $\sum \text{REE}_2\text{O}_3$ (La<sub>2</sub>O<sub>3</sub> + Ce<sub>2</sub>O<sub>3</sub> + Pr<sub>2</sub>O<sub>3</sub> + Nd<sub>2</sub>O<sub>3</sub> + Sm<sub>2</sub>O<sub>3</sub> + Gd<sub>2</sub>O<sub>3</sub>) concentrations range from 52.28 to 55.21 wt%, with a mean of 53.53 wt%. Ce<sub>2</sub>O<sub>3</sub> is considered the most predominant LREE in parisite. It ranges from 22.09 to 23.91 wt% with a mean of 22.99 wt%. La<sub>2</sub>O<sub>3</sub> and Nd<sub>2</sub>O<sub>3</sub> show high concentrations in parisite, with mean values of 14 and 11.71 wt%, respectively, then Pr<sub>2</sub>O<sub>3</sub> followed by Sm<sub>2</sub>O<sub>3</sub> and then Gd<sub>2</sub>O<sub>3</sub>, in ascending magnitudes (Table 5). ThO<sub>2</sub> and SiO<sub>2</sub> are present in minor amounts, with mean values of 0.64 and 0.12 wt%, respectively. The empirical chemical formula of average parisite grains in rhyolite is as follows: Ca<sub>1.02</sub>(Ce<sub>0.58</sub> La<sub>0.36</sub>Nd<sub>0.29</sub>Pr<sub>0.08</sub>Sm<sub>0.04</sub>Gd<sub>0.01</sub>)<sub>2</sub>(CO<sub>3</sub>)<sub>3</sub>F<sub>1.84</sub>.

C. Chevkinite-(Ce) occurs as fine-grained crystals, up to 5 µm in length (Fig. 8d), subrounded to subhedral, and enclosed in quartz. It is recorded only in the rhyolite of the study area and has a limited abundance. It is a titanosilicate REE mineral. As a result of EPMA data, chevkinite crystals from the rhyolite are mainly composed of SiO<sub>2</sub> TiO<sub>2</sub>, and  $Fe_2O_3$  that reached up to 19.97, 20.35, and 13.11 wt%, respectively (Table 5). The analyzed EPMA data indicate that chevkinite shows considerably higher  $\sum LREE_2O_3$  $(La_2O_3-Sm_2O_3)$  concentrations, reaching up to 44.21 wt% (Table 5). Chevkinite is enriched in LREE and  $Ce_2O_3$  is the main predominant REE that reaches 21.52 and 21.85 wt% corresponding to chevkinite-(Ce) in chemical composition. La<sub>2</sub>O<sub>3</sub> shows high concentrations reaching to 11.38 and 11.56 wt% and Nd<sub>2</sub>O<sub>3</sub> 7.87 and 8.24 wt%, while  $Pr_2O_3$ and Sm<sub>2</sub>O<sub>3</sub> show low concentrations in the analyzed samples (Table 5). ThO<sub>2</sub> and UO<sub>2</sub> are present in minor amounts, up to 2.11 and 0.22 wt%, respectively. CaO and P2O5 are recorded also in minor amounts, up to 1.65 and 0.09 wt%, respectively and/or not recorded.

The chemical composition of chevkinite has the formula  $A_4BC_2D_2(Si_2O_7)_2O_8$ , where the cations are represented in each site as follows:  $A = REE^{3+}$ ,  $Ca^{2+}$ ;  $B = Fe^{2+}$ ;  $C = Fe^{2+}$ ,  $Fe^{3+}$ ,  $Ti^{4+}$ ,  $Al^{3+}$ ; D,  $Ti^{4+}$  according to Macdonald et al. (2012) and Bagiński et al. (2015).

#### Other accessory minerals

Sulfide minerals including pyrite, galena, and sphalerite are recorded in the microgranite, whereas fluorite and fluorapatite are presented in the rhyolite.

Pyrite occurs as fine-grained, anhedral grains associated with fergusonite, columbite, and zircon in the microgranite (Fig. 6c). Other pyrite grains, associated with galena, are disseminated throughout plagioclase (Fig. 9a). EPMA data show that the chemical composition of pyrite is Fe reaching 47.03 wt% and S attaining 52.74 wt%.

 Table 3
 Representative EPMA data (oxides, in wt%) of Nb-rich minerals from microgranite dikes, Abu Hawis area, North Eastern Desert, Egypt

Min./oxid	Columbi	te							Fergus	onite			
Y <sub>2</sub> O <sub>3</sub>	0.84	0.42	0.45	0.43	0.98	0.62	n.d	n.d	22.16	23.15	22.13	23.58	23.64
$Nb_2O_5$	72.07	73.03	74.68	73.53	71.31	71.8	74.57	74.93	48.6	51.95	50.77	51.19	48.97
Ta <sub>2</sub> O <sub>5</sub>	1.89	1.62	1.64	1.67	1.81	1.79	0.73	0.72	0.78	0.81	0.75	0.79	0.74
$\mathrm{Gd}_2\mathrm{O}_3$	n.d	1.15	0.63	0.64	0.62	0.58							
Dy <sub>2</sub> O <sub>3</sub>	n.d	3.24	2.64	3.27	2.43	2.75							
Ho <sub>2</sub> O <sub>3</sub>	n.d	0.89	1.02	1.1	0.73	1.13							
Er <sub>2</sub> O <sub>3</sub>	n.d	4.37	4.44	4.86	4.38	5.22							
Tm <sub>2</sub> O <sub>3</sub>	n.d	0.75	1.53	0.78	1.02	1.93							
Yb <sub>2</sub> O <sub>3</sub>	n.d	6.72	6.08	5.11	6.48	5.75							
$Lu_2O_3$	n.d	0.73	0.57	0.61	0.54	1.72							
TiO <sub>2</sub>	3.37	2.94	1.71	3.05	4.62	3.1	2.14	2.09	1.62	1.53	0.93	0.33	0.67
CaO	0.28	0.28	n.d	n.d	0.07	0.22	0.36	0.45	0.71	0.41	0.97	0.99	0.92
MnO	14.21	14.03	14.95	13.75	14.22	14.15	1.10	1.52	0.61	0.71	0.55	0.53	0.49
Fe <sub>2</sub> O <sub>3</sub>	6.21	6.61	6.51	6.85	6.12	6.98	18.11	17.25	0.79	0.95	2.85	1.23	0.85
ThO <sub>2</sub>	n.d	n.d	n.d	n.d	n.d	n.d	1.75	1.31	2.15	1.14	2.06	2.94	2.33
UO <sub>2</sub>	1.07	0.87	n.d	0.71	0.85	1.43	1.24	1.04	3.82	1.22	1.33	0.73	0.71
SiO <sub>2</sub>	n.d	0.69	1.17	1.04	1.49	1.60							
Total	99.94	99.8	99.94	99.99	99.98	100.09	100	99.31	99.78	99.95	99.75	100	100
$\Sigma$ HREE <sub>2</sub> O <sub>3</sub>									18.85	16.91	17.37	16.2	19.08
Cation													
Y	0.03	0.01	0.01	0.01	0.03	0.02	0.00	0.00	0.52	0.53	0.51	0.54	0.55
Nb	1.84	1.86	1.90	1.87	1.80	1.83	1.91	1.93	0.97	1.00	0.99	1.00	0.97
Та	0.03	0.02	0.03	0.03	0.03	0.03	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Gd									0.02	0.01	0.01	0.01	0.01
Dy									0.05	0.04	0.05	0.03	0.04
Но									0.01	0.01	0.02	0.01	0.02
Er									0.06	0.06	0.07	0.06	0.07
Tm									0.01	0.02	0.01	0.01	0.03
Yb									0.09	0.08	0.07	0.09	0.08
Lu									0.01	0.01	0.01	0.01	0.02
Ti	0.14	0.12	0.07	0.13	0.19	0.13	0.09	0.09	0.05	0.05	0.03	0.01	0.02
Ca	0.02	0.02	0.00	0.00	0.00	0.01	0.02	0.03	0.03	0.02	0.04	0.05	0.04
Mn	0.68	0.67	0.71	0.65	0.67	0.68	0.05	0.07	0.02	0.03	0.02	0.02	0.02
Fe	0.29	0.31	0.31	0.32	0.29	0.33	0.86	0.82	0.03	0.03	0.09	0.04	0.03
Th	0.00	0.00	0.00	0.00	0.00	0.00	0.02	0.02	0.02	0.01	0.02	0.03	0.02
U	0.01	0.01	0.00	0.01	0.01	0.02	0.02	0.01	0.04	0.01	0.01	0.01	0.01
Si									0.03	0.05	0.04	0.06	0.07
$\sum$ Cations	3.03	3.03	3.03	3.02	3.03	3.05	2.99	2.98	1.98	1.96	1.99	1.98	2.00
Ta/(Ta + Nb)	0.01604	0.01064	0.01554	0.01579	0.01639	0.01613	0.00521	0.00515					
Mn/(Mn + Fe)	0.7010	0.6837	0.6961	0.6701	0.6979	0.6733	0.0549	0.0787					

Min., mineral; oxid., oxides

Calculations are based on 6 and 4 oxygen atoms per formula unit (apfu) for columbite and fergusonite, respectively

n.d., not determined

Galena is a lead sulfide mineral and occurs as very fine grains, thin films along the peripheries of sphalerite grains, and/or fine grains along zircon in microgranite (Fig. 9a–d). EPMA data show that Pb concentration reached 85.22 wt% and S 12.73 wt%. Fe and Si are pre-

sented as minor constituents, with mean values of 0.51 wt% and 0.89 wt%, respectively.

Sphalerite is a zinc sulfide mineral and occurs as anhedral to subhedral grains and contains thin films of galena along its peripheries in the microgranite (Fig. 9d). EPMA data

Fig. 8 Back-scattered images of REE-bearing minerals from felsic dikes, Abu Hawis area, North Eastern Desert, Egypt. **a** Micro-veinlet filled with monazite and hematite in microgranite; **b** corroded parisite crystal enclosing monazite in rhyolite; **c** anhedral crystal of parasite in rhyolite; **d** fine-grained chevkinite in rhyolite. Abbreviations: mon, monazite; par, parisite; chv, chevkinite; hem, hematite



show that sphalerite is composed mainly of Zn reaching 61.59 wt% and S 34.91 wt%. Si and Fe occurred as minor constituents, reaching 0.69 and 2.22 wt%, respectively. Fluorite occurs as anhedral to subhedral grains in the rhyolite. It occurs also as fine-grained and is associated with thorite, parisite, and hematite (Fig. 5c). EPMA data show that fluorite is composed mainly of Ca 52.23 wt% and F 45.72 wt%, with minor amounts of Si 0.19 wt%.

Apatite occurs as anhedral grains in the rhyolite and is associated with parisite and thorite (Fig. 5d). EPMA data show that the main constituents of apatite are CaO 55.03 wt% and  $P_2O_5$  42.07 wt%. SiO<sub>2</sub> occurs as minor constituent 0.41 wt% whereas F reached to 5.34 wt% designating fluorapatite.

## Discussion

# The E–W and NE–SW control of rare metal mineralization

Felsite dike swarms are almost abundant and numerous in the North Eastern Desert of Egypt, having E–W and NE–SW (Ali 2007, Dessouky et al. 2020; Waheeb and EL Sundoly 2020). The distribution of felsite dikes confirms that a gradual transition from N-S extension trend to NE–SW and NW–SE extension trends was occurred in Abu Hawis area (Ali 2007; Waheeb and EL Sundoly 2020). Ali (2009) states that a marked positive correlation was taken place between the extensional events and the NE–SW, NW–SE, and E–W trends. These events led to U-migration from Abu Hawis monzogranite to be deposited along the main shear zones trending in the NE–SW direction. These trends dominate on the mobility map (eU-eTh/3.5), and show NW–SE, NE–SW, and N–S extensional trends, respectively.

Abu Hawis granitoids are located to the northeast of El-Erediya granites enriched in U mineralization especially in jasper veins having NE–SW to ENE–WSW-striking fractures that dated 130–160 Ma for hydrothermal event (Abu-Deif 1992). Besides, the studied area is located in the western part of Ras Abda syenogranite. Age dating of the magmatic crystallization for Ras Abda mineralized syenogranite is 610 Ma (Dessouky et al. 2020; Abdel Gawad et al. 2021a, b, c), whereas the age of the hydrothermal event is  $126.8 \pm 2.0$  Ma (Abdel Gawad et al. 2021a, b, c). These trends could have been reactivated many times at different ages. Therefore, they may be used as good indicators for the existence U mineralization that is associated with other rare metal mineralization.

# Post-magmatic and hydrothermal processes controlling mineralization

The radioactive mineralizations — including uraninite, thorite, and uranothorite — are well recorded in felsic dikes. They could be of a syn-genetic origin during their emplacement along E–W and NE–SW structural trends. The sulfide Table 4Representative EPMAdata (oxides, in wt%) ofmonazite from the studied felsicdikes, Abu Hawis area, NorthEastern Desert, Egypt

Min in dileo/orid	Monar	ita in m	200 000-	ta		Aver	Aver Monazite in rhyolite				
wini. in dike/oxid	wonaz	ite in mi	crogran			Aver	wonaz	ite in rh	yonte		Aver
SiO <sub>2</sub>	0.19	0.26	0.29	0.28	0.25	0.25	0.24	0.22	0.23	0.19	0.22
CaO	0.43	0.41	0.24	0.22	0.22	0.30	0.29	0.32	0.28	0.34	0.31
$P_2O_5$	26.66	29.18	30.91	30.98	29.02	29.35	29.32	28.48	29.31	29.21	29.08
Fe <sub>2</sub> O <sub>3</sub>	0.24	0.28	0.31	0.25	0.19	0.25	0.31	0.37	0.29	0.42	0.35
La <sub>2</sub> O <sub>3</sub>	9.23	9.86	10.25	8.69	7.81	9.17	16.12	15.88	16.14	15.25	15.85
$Ce_2O_3$	31.79	29.5	28.76	31.95	30.12	30.42	32.24	32.42	32.74	33.06	32.62
Pr <sub>2</sub> O <sub>3</sub>	4.27	4.58	3.99	4.19	4.73	4.35	3.44	2.67	2.87	3.48	3.12
Nd <sub>2</sub> O <sub>3</sub>	11.14	12.52	9.93	10.17	12.19	11.19	14.03	13.95	14.12	14.56	14.17
Sm <sub>2</sub> O <sub>3</sub>	2.39	2.11	1.95	1.60	2.18	2.05	1.83	2.16	1.19	1.23	1.60
$Gd_2O_3$	1.37	1.18	1.09	1.16	1.20	1.20	0.63	0.74	0.87	0.92	0.79
ThO <sub>2</sub>	11.96	10.11	11.98	10.51	11.87	11.29	1.55	2.16	1.19	1.23	1.53
Total	99.67	99.99	99.7	100	99.78	99.83	100	99.37	99.23	99.89	99.62
$\sum \text{REE}_2 O_3$	60.19	59.75	55.97	57.76	58.23	58.38	68.29	67.82	67.93	68.5	68.14
Cation	Calcul	ated che	mical for	rmulae a	ire based	l on 4 ox	ygen ato	oms (apf	u)		
Si	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Ca	0.02	0.02	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Р	0.94	0.99	1.02	1.02	0.99	0.99	0.98	0.97	0.99	0.98	0.98
Fe	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
La	0.14	0.15	0.15	0.12	0.12	0.14	0.24	0.24	0.24	0.22	0.23
Ce	0.49	0.43	0.41	0.45	0.44	0.44	0.47	0.48	0.48	0.48	0.47
Pr	0.06	0.07	0.06	0.06	0.07	0.06	0.05	0.04	0.04	0.05	0.05
Nd	0.17	0.18	0.14	0.14	0.18	0.16	0.20	0.20	0.20	0.21	0.20
Sm	0.03	0.03	0.03	0.02	0.03	0.03	0.02	0.03	0.02	0.02	0.02
Gd	0.02	0.02	0.01	0.01	0.02	0.02	0.01	0.01	0.01	0.01	0.01
Th	0.11	0.09	0.11	0.09	0.11	0.10	0.01	0.02	0.01	0.01	0.01
$\Sigma$ cations	2.00	1.98	1.95	1.96	1.97	1.97	2.01	2.02	2.01	2.01	2.01

Min., mineral; oxid., oxides; Aver., average

 $\sum \text{REE}_2\text{O}_3 (\text{La}_2\text{O}_3 + \text{Ce}_2\text{O}_3 + \text{Pr}_2\text{O}_3 + \text{Nd}_2\text{O}_3 + \text{Sm}_2\text{O}_3 + \text{Gd}_2\text{O}_3)$ 

n.d., not determined

minerals in the microgranite are pyrite, sphalerite, and galena as accessories associated with U and Th minerals.

The total analyses of thorite from both microgranite and rhyolite are often close to 90 wt% (Table 1) due to strong hydration or metamictization (Abd El-Naby 2009; Gaafar et al. 2014; Abdel Gawad 2021). Some grains of thorite are enclosed in hematite, in which could be a good indicator of strong alteration.

Zircon crystals incorporate micro-inclusions of uraninite, thorite, uranothorite, monazite, and zircon into their crystal lattice, which could enhance metamictization of zircon (Alekseev and Alekseev 2020; Levashova et al. 2021) (Fig. 4a). The presence of fine-grained zircon as inclusions in large zircon crystals indicates the occurrence of two zircon generations in the studied microgranite dikes. EPMA data show higher  $HfO_2$  that reached 9.08 wt% in the analyzed zircon crystals from microgranite dikes. This could be a good indicator for the hydrothermal activities, according to Wang et al. (2010). The calculated Ta/(Ta + Nb) and Mn/(Mn + Fe) ratios of the analyzed columbite from microgranite range between (0.0052 and 0.0164) and (0.0549 and 0.7010), respectively, which indicate manganocolumbite composition. Fergusonite is the most dominated HREE-bearing mineral recorded in the microgranite with a mean  $\Sigma$ HRE<sub>2</sub>O<sub>3</sub> 17.68 wt%. On the other hand, parisite and chevkinite are LREE minerals recorded in the rhyolite with means  $\Sigma$ LRE<sub>2</sub>O<sub>3</sub> 53.53 and 43.75 wt%. However, monazite is a dominated LREE mineral in both microgranite and rhyolite with means  $\Sigma$ LRE<sub>2</sub>O<sub>3</sub> 58.38 and 68.14 wt%.

High field strength elements (HFSE) Zr, Hf, Th, and Ti are generally mobile elements during magmatic stage, and/ or hydrothermal alterations containing complexing agents as F, S, and others (Keppler 1993; Ali 2012). Abu Hawis, Umm Tagher El Foqani, and Ras Barud monzogranites are enriched in HFSE (Zr, Hf, Th, and Ti), as well as rare metal mineralization. It is well known that F could have a prominent role during mobilization of Zr, Hf, Th, and Ti as well

Table 5 Representative EPMA data (oxides, in wt%) of parisite and chevkinite from rhyolite, Abu Hawis area, North Eastern Desert, Egypt

Min/oxid	Parisit	e	Aver	Chevkinite							
SiO <sub>2</sub>	0.13	0.16	0.11	0.13	0.08	0.12	0.13	0.09	0.12	19.79	19.97
CaO	13.8	14.87	14.08	13.89	12.84	13.75	13.55	13.64	13.80	1.21	1.65
$P_2O_5$	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	0.09	n.d
TiO <sub>2</sub>	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	20.35	19.25
Fe <sub>2</sub> O <sub>3</sub>	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	13.11	12.59
$La_2O_3$	14.58	13.64	14.05	13.9	14.72	14.21	13.62	13.27	14.00	11.38	11.56
Ce <sub>2</sub> O <sub>3</sub>	22.72	22.73	23.91	22.64	23.77	22.18	22.09	23.9	22.99	21.52	21.85
Pr <sub>2</sub> O <sub>3</sub>	3.08	3.17	3.03	2.89	3.01	2.94	3.06	2.98	3.02	1.78	1.85
Nd <sub>2</sub> O <sub>3</sub>	10.74	10.92	12.24	11.35	11.55	12.48	12.26	12.12	11.71	7.87	8.24
$Sm_2O_3$	1.41	1.48	1.61	1.61	1.31	1.42	1.53	1.54	1.49	0.73	0.71
$Gd_2O_3$	0.32	0.34	0.37	0.31	0.30	0.26	0.35	0.29	0.32	n.d	n.d
ThO <sub>2</sub>	0.49	0.62	0.79	0.45	0.44	0.90	0.75	0.65	0.64	1.98	2.11
UO <sub>2</sub>	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	n.d	0.18	0.22
F	8.56	7.81	8.45	8.72	8.69	8.73	8.19	8.09	8.41	n.d	n.d
Total	75.83	75.74	78.64	75.89	76.71	76.99	75.53	76.57	76.49	99.99	100
$\sum \text{REE}_2 O_3$	52.85	52.28	55.21	52.7	54.66	53.49	52.91	54.1	53.53	43.28	44.21
Cation S;	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	4.03	4 10
SI Co	1.02	1.11	1.02	1.03	0.01	1.01	1.02	1.02	1.02	4.05	4.10
Са Р	1.05	1.11	1.02	1.05	0.95	1.01	1.02	1.02	1.02	0.20	0.00
Ti										3.12	2.08
Fe										2.23	2.50
La	0.37	0.35	0.35	0.35	0.38	0.36	0.35	0 34	0.36	0.86	0.88
Ce	0.58	0.55	0.59	0.55	0.60	0.56	0.55	0.61	0.58	1.68	1.72
Pr	0.08	0.08	0.07	0.07	0.08	0.07	0.08	0.08	0.08	0.14	0.15
Nd	0.00	0.00	0.30	0.28	0.29	0.31	0.31	0.30	0.29	0.57	0.60
Sm	0.03	0.04	0.04	0.04	0.03	0.03	0.04	0.04	0.04	0.05	0.05
Gd	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.00	0.00

Min., mineral; oxid., oxides; Aver., average

0.01

1.72

4.18

0.01

1.81

4.20

0.01

1.88

4.25

Calculated formulae (apfu) of parasite are based on number of ions on the basis of two fluorine apfu on the basis of (CO<sub>3</sub>)<sub>3</sub>F<sub>2</sub>

0.01

1.90

4.25

0.01

1.89

4.26

0.01

1.82

4.21

0.01

1.78

4.19

0.01

1.84

4.23

0.09

0.01

13.07

0.10

0.01

13.11

Chevkinite formulae is calculated on the basis of 22 oxygen per formula unit (apfu)

0.01

1.90

4.27

n.d., not determined

Th

U

F

∑cations

as REE (Moine and Salvi 1999) during hydrothermal alterations (argillic and hematitization).

The reactivated structures are accompanied in association with succession fluid circulations leading to the injection of a series of microgranite, rhyolite, andesite, and basaltic dikes as well as quartz veins. Changes in host rock composition and mineralogy are used to decipher the type and extent of fluid-wall rock interactions during alteration. The physico-chemical conditions of these fluids were changing due to consequent reactions with the granitoids and post-granitic dikes with meteoric water.

The hydrolysis of K-feldspar and plagioclase into illite or Na-montmorillonite (K-alteration) according to the reactions (1, 2, and 3) (Pirajno 1992).

$$3\text{KAISi}_{3}\text{O}_{8 \text{ K-feldspar}} + 2\text{H}^{+}_{\text{acidicfluids}}$$

$$\rightarrow \text{KAI}_{2}(\text{Si}_{3}\text{AIO}_{10})(\text{OH})_{2 \text{ illite}} + 2\text{K}^{+} + 6\text{SiO}_{2 \text{ quartz}}$$
(1)

$$6NaAlSi_{3}O_{8 Na-plagioclase} + 2K^{+} + 4H^{+}$$
(2)

$$\rightarrow 2\text{KAl}_2(\text{Si}_3\text{AlO}_{10})(\text{OH})_{2 \text{ illite}} + 12\text{SiO}_{2 \text{ quartz}} + 6\text{Na}^+$$

Fig. 9 Back-scattered images of sulfide minerals from microgranite, Abu Hawis area, North Eastern Desert, Egypt. **a** Pyrite associated with very fine grains of galena, enclosed in plagioclase; b fine grains of galena, associated with microveinlets of hematite, enclosed in K-feldspar; c fine grain of galena occurs along the peripheries of zircon crystal; d thin film of galena occurring along the peripheries of sphalerite grain. Abbreviations: zir, zircon; pyt, pyrite; gal, galena; sph, sphalerite; hem, hematite



$$\begin{split} &1.17 NaAlSi_3O_{8\ Na-plagioclase} + H^+ \rightarrow 0.5 Na_{0.33}Al_{2.33} \\ &Si_{3.67}O_{10}(OH)_{2\ Na-montmorilloni}te + 1.67SiO_{2\ quartz} + Na^+ \end{split}$$

 $\begin{aligned} & \text{KAl}_{2}(\text{Si}_{3}\text{AlO}_{10})(\text{OH})_{2 \text{ illite}} + \text{H}^{+} + 1.5 \\ & \text{H}_{2}\text{O} \rightarrow 1.5\text{Al}_{2}\text{Si}_{2}\text{O}_{5}(\text{OH})_{4 \text{ kaolinite}} + \text{K}^{+} \end{aligned} \tag{4}$ 

The hydrolysis of illite and Na-montmorillonite produces kaolinite according to the following reactions (4 and 5), respectively (Pirajno 1992).

 $3Na_{0.33}Al_{2.33}Si_{3.67}O_{10}(OH)_{2\ Na-montmorillonite}$ 

(3)

 $+ H^{+} + 3.5H_{2}O \rightarrow 3.5Al_{2}Si_{2}O_{5}(OH)_{4 \text{ kaolinite}} + 4SiO_{2 \text{ quartz}} + Na^{+}$ (5)

Fig. 10 XRD for clay alteration products (montmorillonite, illite, and kaolinite), Abu Hawis area, North Eastern Desert, Egypt



Table 6X-ray diffraction dataof clay alteration products,Abu Hawis area, North EasternDesert, Egypt

Clay alteration products		Identifie	Identified minerals										
		Montmo	orillonite	Illite		Kaolinite							
		ASTM card no											
		13–0135		9–0343		06–221							
dÅ	I/I <sub>0</sub>	dÅ	I/I <sub>0</sub>	dÅ	I/I <sub>0</sub>	dÅ	I/I <sub>0</sub>						
15	99	15	100										
10.10	100			10	100								
7.20	72					7.18	100						
5.02	26			4.95	80								
4.52	8	4.5	80	4.48	90	4.48	80						
3.58	15					3.58	100						
2.5	2	2.5	40	2.5	10								
2.39	1					2.386	80						

Clay alteration products have been identified by using XRD analyses (Fig. 10 and Table 6).

Monazite, parisite, and chevkinite are associated with fluorapatite in the rhyolite dikes. These LREE minerals are poor in Th (Tables 4 and 5), with the absence of U and/or the presence with minor concentrations in chevkinite (up to 0.22 wt%). In this case of LREE minerals, the compositional features could suggest their formation as a result of fluorapatite metasomatism (Ziemann et al. 2005; Ali 2012). On the other hand, the chemical composition of monazite from the microgranite shows high Th contents, with a mean of ThO<sub>2</sub> 11.29 wt%, which could indicate magmatic monazite.

# Conclusions

- 1- The investigated felsic dikes having E–W and NE–SW trends at Abu Hawis area show a marked enrichment of rare metal mineralization, through the use of electron probe microanalysis.
- 2- Uraninite, uranothorite, and thorite are recorded in the microgranite dikes, while the latter one is only recorded in the rhyolite dikes.  $UO_2$  contents reached up to 69.57, 27.42, and 5.33 wt% in the analyzed uraninite, uranothorite, and thorite from the microgranite dikes. HREEs ( $Dy_2O_3$ ,  $Er_2O_3$ , and  $Yb_2O_3$ ) were recorded as minor amounts in uraninite and thorite, but absent in uranothorite.
- Both ThO<sub>2</sub> and UO<sub>2</sub> show low concentrations in zircon crystals, whereas HfO<sub>2</sub> shows high values up to 9.08 wt% from the microgranite dikes. This could be related to strong hydrothermal alteration (hematitization). HREE and Y<sub>2</sub>O<sub>3</sub> are well presented in zircon crystals from the rhyolite dikes.

- 4- Nb-bearing minerals are well represented by columbite and fergusonite from the microgranite dikes. Columbite is enriched in manganese relative to iron and this could indicate the formation of manganocolumbite composition, except in only two analyses that are ferrocolumbite composition. Fergusonite is enriched in Nb<sub>2</sub>O<sub>5</sub>, Y<sub>2</sub>O<sub>3</sub>, and HREE<sub>2</sub>O<sub>3</sub> that reached up to 51.95, 23.64, and 19.08 wt%.
- 5- REE minerals including monazite are well predominated in both the microgranite and the rhyolite dikes, with mean values of  $P_2O_5$  (29.35 to 29.08 wt%), ThO<sub>2</sub> (11.29 to 1.53 wt%), and LREE<sub>2</sub>O<sub>3</sub> (58.38 to 68.14 wt%). Parisite and chevkinite are recorded only in the rhyolite dikes. They are enriched in LREE, where Ce<sub>2</sub>O<sub>3</sub> is the most predominant REE that reaches up to 21.85–23.91 wt%, followed by La<sub>2</sub>O<sub>3</sub> 11.56– 14.00 wt%, then Nd<sub>2</sub>O<sub>3</sub> 8.24–11.71 wt%, with minor amounts of other element in both parisite and chevkinite, respectively.

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# Declarations

**Conflict of interest** The author declares that he has no competing interests.

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