

Study on the element geochemical characteristics of the Shazi large-sized anatase ore deposit in Qinglong, Guizhou Province

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Abstract The Shazi anatase ore deposit in Qinglong, Guizhou Province, is a large-sized anatase deposit that has been recently explored. The characteristics of major oxides in the ore are similar to those of modern laterite weathering crust and laterite in the laterite-type gold deposits in the western part of Guizhou Province. Studies on the REE characteristics of basalts and anatase ores in the study region showed that both of them do have extremely strong affinities. There are two groups of trace elements in the ores, i.e., Au-Ag-As-Sb-Hg-Tl association and Sc-TiO₂-Cu-Fe-Mn association, reflecting that the formation of anatase ore is related to the formation of siliceous claystone at the early stage of eruption of the Emeishan basaltic magma. The siliceous claystones are the major country rocks for the formation of laterite-type gold ores and anatase ores. In the region anatase ores are rich in Sc and the basalts enriched in Fe, Mn, Ti and Sc are the material source of metallogenesis.

Key words anatase ore deposit; Emeishan basalt; trace element association; element geochemical characteristic; Qinglong; Guizhou

1 Introduction

The Shazi anatase ore deposit was discovered in 2007 by Profs. Nie Aiguo in Qinglong, Guizhou Province. From 2007 to 2011 a general survey and detailed geological work had been conducted on this deposit. It has been basically verified that the ore reserves of this anatase ore deposit are 2306.09×10^5 ton and the quantity of TiO₂ resource is 102.48×10^5 ton, with an average TiO₂ grade of 4.39%. It is indicated that the Shazi anatase ore deposit belongs to a large-scale eluvial-type anatase ore deposit. Before the discovery of the Shazi anatase ore deposit, there had been found the metasedimentary-type and hydrothermal reworking-type anatase ore deposits only in the Yangtizishan-Moshishan area, Inner Mongolia, China (Zhao et al., 2008, 2006). This is the first discovery of anatase

ore in the Emeishan basalt-weathering eluvial soil profile in Guizhou Province, marking the first discovery of a large-sized eluvial-type anatase ore deposit, which makes such a history come into an end that there is no anatase ore resource in Guizhou Province. This discovery is of great practical significance for the reevaluation of ore resources in Southwest Guizhou Province and has also made a certain contribution to exploration of anatase resources and research on their origin both at home and abroad.

Anatase ore is one of the ore types of critical shortage in China and one of the 14 kinds of strategic ore reserves on which China relies seriously (Wu and Zhang, 2006; Cao, 1996; Gao and Wang, 2008). Theoretical research on the newly discovered Shazi anatase ore deposit in Qinglong has just begun and there still exist many blind zones in research. To carry

out studies on the geochemical characteristics of this anatase ore deposit will promote deep-going investigations into the rules of transport, enrichment and metallogenesis of anatase ore and also are of great significance in constraining their origins.

Ten ore-exposed drill core assemblage samples were collected randomly from the Shazi anatase ore deposit in Qinglong for bulk oxide analysis, 13 ore-exposed drill core assemblage samples were collected from the same deposit for REE and trace element analysis, and 7 basalt samples were taken from the mining district for REE and trace element analysis. Then, analysis and discussion were conducted on the above analytical results. It is concluded that this anatase ore deposit is rich in Sc and is of great economic significance.

2 Geology of the ore deposit

There have been already explored three anatase orebodies of industrial value, which are arranged in a N-E-S direction and are numbered consecutively: No. 1 anatase orebody, No. 2 anatase orebody and No. 3 anatase orebody (Nie et al., 2011). Through phase analysis it can be seen that the independent minerals of this anatase ore deposit do not occur as commonly observed ilmenite and rutile, but occur in the form of anatase which is not usually observed. The discovered anatase orebodies are distributed in the Quaternary cumulates derived from weathering of basalts in the karst negative-relief fluid bowls on the Maokou Formation limestones at the altitude of 1450–1350 m above sea level, forming eluvial claystone-type ores. Ore minerals are mainly anatase and ilmenite; opaque minerals are mainly sericite, chlorite, kaolinite, quartz and secondly are mica, plagioclase and ferrous mud. Ore structures are: as observed by naked eyes, the ores are earthy-yellow, pale brown, grey in color and loosely earthy. The ores exhibit mainly earthy massive structure, secondly massive, faviform, brecciated structures. As observed under microscope, some limonite is distributed as fine veins along ore fissures, thus making the ores exhibiting fine-vein structure either. The ore textures are dominated by muddy texture, micro-scaly texture, blastoporphyritic texture, micro-crystalline texture, pseudomorphic texture, alteration intergradular-intersertal texture, fine sand texture, and alteration aleuritic texture. It is primarily considered that this anatase ore deposit belongs to the eluvial slope-type anatase ore deposits associated with Emeishan basaltic magma eruption.

Through process mineralogical studies it has been found that the contents of anatase (the molecular formula is TiO_2) are about 4.42%. Electron microprobe compositional analysis revealed that anatase

contains as much Ti, O and Si as 51.37%, 47.72%, and 0.91%, respectively. Microscopic observation shows that anatase displays yellowish-green, yellowish-brown, adamantine and port oily lusters, and they are cubic dipyramid-shaped and xenomorphic granular. As observed under polarizing microscope, anatase is distributed mainly in seriously altered basalts, it is xenomorphic granular, dentritic and muddy in shape and is of close intergrowth with quartz, feldspar and muddy material and are enclosed in gangue minerals or are distributed in quartz and feldspar fissures. It also intergrows with ferruginous material and is probably derived from the decomposition of melanocratic minerals during the process of rock alteration. Therefore, its grains are fine. In the artificial heavy concentrates an extremely small amount of granular anatase (about 0.15%) can be observed under stereomicroscope, but when observed under polarizing microscope anatase grains which are less than 47.2 μm in size account for 96.35% and the grains which are less than 23.6 μm account for 75.37%. It can be seen that the dissemination extent of anatase is relatively small. This kind of anatase occurs as micro-fine particles in soils derived from weathering of the Emeishan basalts.

3 Major element geochemical characteristics

Oxide bulk analysis was conducted of ten ore-exposed drill-core composite samples from this ore deposit and the analytical results are listed in Table 1. The major characteristics are presented as follows:

(1) The total amount of major oxides such as SiO_2 , Al_2O_3 , Fe_2O_3 and TiO_2 in the ore is within the range of 81.64%–88.15% and this feature is approximate to that of laterite in the laterite-type gold ores in the western part of Guizhou and also to that of the laterite weathering crust in Guizhou (Wang et al., 2000; Gao et al., 2002), though the content of TiO_2 is relatively high.

(2) Iron in the ores is all present in the form of Fe_2O_3 , indicating that the ores were strongly weathered and completely oxidized.

(3) The content of SiO_2 is all less than 55%, indicating that the ores are clayey ores.

(4) There is a positive correlation between TiO_2 and Fe_2O_3 and the correlation coefficient is 91.06%, reflecting that anatase has a relationship of coexistence (association) with Fe-bearing minerals in the protolith.

(5) There is a positive correlation between TiO_2 and Al_2O_3 and the correlation coefficient is 86.21%, reflecting that anatase has a relationship of coexistence (association) with clay minerals in the protolith.

Table 1 Analytical results for the major elements and relevant coefficients of the Shazi anatase ore deposit, Qinglong, Guizhou Province

| Sample No. | Analytical result (10^{-2}) | | | | | | | | | | |
|--|---------------------------------|--------------------------------|---------------------------------|------------------|---------|---------|------------------|-------------------|------------------|-------------------------------|-------|
| | SiO ₂ | Al ₂ O ₃ | TFe ₂ O ₃ | TiO ₂ | CaO | MgO | K ₂ O | Na ₂ O | MnO ₂ | P ₂ O ₅ | LOSS |
| ZK5501 | 34.26 | 21.08 | 23.10 | 4.71 | 0.20 | 0.32 | 0.52 | 0.02 | 0.29 | 0.35 | 12.78 |
| ZK5902 | 30.26 | 24.23 | 23.95 | 4.97 | 0.06 | 0.17 | 0.35 | 0.04 | 0.32 | 0.49 | 13.62 |
| ZK21-2 | 32.16 | 21.40 | 23.74 | 4.71 | 0.90 | 1.39 | 1.30 | 0.46 | 0.16 | 0.27 | 12.02 |
| ZK903 | 26.60 | 24.19 | 25.82 | 5.03 | 0.07 | 0.13 | 0.44 | 0.03 | 0.25 | 0.37 | 14.04 |
| ZK104 | 41.08 | 19.70 | 19.45 | 3.93 | 0.26 | 0.51 | 0.61 | 0.01 | 0.21 | 0.35 | 11.80 |
| ZK1103 | 51.22 | 16.09 | 16.94 | 3.90 | 0.20 | 0.34 | 1.48 | 0.02 | 0.07 | 0.52 | 7.78 |
| ZK1701 | 46.10 | 15.26 | 18.18 | 3.42 | 1.35 | 1.30 | 0.87 | 1.02 | 0.12 | 0.47 | 10.34 |
| ZK23-1 | 36.96 | 21.45 | 20.89 | 3.80 | 0.25 | 0.47 | 0.67 | 0.05 | 0.26 | 0.39 | 12.68 |
| ZK5003 | 29.62 | 23.38 | 25.52 | 5.16 | 0.08 | 0.20 | 0.52 | 0.04 | 0.32 | 0.39 | 13.74 |
| ZK5401 | 43.28 | 18.34 | 16.31 | 3.80 | 0.82 | 0.92 | 0.64 | 0.02 | 0.22 | 0.40 | 12.32 |
| SiO ₂ | 1 | | | | | | | | | | |
| Al ₂ O ₃ | -0.9538 | 1 | | | | | | | | | |
| TFe ₂ O ₃ | -0.9519 | 0.8915 | 1 | | | | | | | | |
| TiO ₂ | -0.8732 | 0.8621 | 0.9106 | 1 | | | | | | | |
| CaO | 0.4397 | -0.6426 | -0.4822 | -0.5812 | 1 | | | | | | |
| MgO | 0.3583 | -0.5439 | -0.4025 | -0.5013 | 0.9518 | 1 | | | | | |
| K ₂ O | 0.5877 | -0.6138 | -0.4414 | -0.3802 | 0.3961 | 0.5054 | 1 | | | | |
| Na ₂ O | 0.2632 | -0.4969 | -0.1892 | -0.3921 | 0.8563 | 0.7733 | 0.3264 | 1 | | | |
| MnO ₂ | -0.7782 | 0.8309 | 0.6643 | 0.6443 | -0.5577 | -0.5574 | -0.8700 | -0.4905 | 1 | | |
| P ₂ O ₅ | 0.4967 | -0.3826 | -0.4181 | -0.3050 | -0.0719 | -0.2736 | 0.0773 | 0.0619 | -0.2446 | 1 | |
| LOSS | -0.8982 | 0.8727 | 0.7472 | 0.6637 | -0.3519 | -0.3269 | -0.8262 | -0.3070 | 0.9062 | -0.4715 | 1 |
| Laterite in the laterite type gold deposits in western Guizhou | 51.50 | 22.39 | 10.93 | 1.56 | 0.22 | 0.13 | 0.97 | 0.16 | 0.13 | 0.02 | 9.50 |
| Laterite weathering crust, Guizhou | 52.35 | 22.02 | 8.55 | 1.16 | 0.20 | 1.38 | 1.97 | 0.10 | 0.02 | 0.04 | 12.07 |

Note: Laterite in the laterite-type gold deposits in the western part of Guizhou (Wang et al., 2000); laterite weathering crust in Guizhou (Liao et al., 2004).

4 REE geochemical characteristics

Thirteen ore-exposed drill -core composite samples randomly were selected from this deposit for REE analysis, and 7 basalt samples were taken from the mining district for REE analysis. The analytical results are listed in Table 2. The REE patterns of basalt samples and ores in three orebodies in the mining district are established (Figs. 1–2) and their characteristics are described as follows:

(1) The total REE amount ($\sum\text{REE}$) of basalts in the mining district is $(183.53\text{--}215.86)\times 10^{-6}$ and that of basalts in the western part of Guizhou is $(144.73\text{--}265.500)\times 10^{-6}$ (Mao et al., 1992), and their abundances vary within the range of abundances of basalts in the western part of Guizhou. The chondrite-normalized REE patterns of basalts in the mining district are of right-inclined type. The LREE values of the samples are $(125.15\text{--}144.40)\times 10^{-6}$, their HREE values are $(58.38\text{--}71.46)\times 10^{-6}$, and their LREE/HREE

ratios vary within the range of 1.92–2.25, indicative of relative LREE enrichment. Basalts in the western part of Guizhou are sodium-modified basalts, and in the more alkaline media the LREE tend to separate from the HREE, thus leading to relative LREE enrichment.

(2) The total REE amount ($\sum\text{REE}$) of ores in the three orebodies in the mining district is relatively high, mostly within the range of $(163.56\text{--}370.26)\times 10^{-6}$. The total REE amount of some samples is $(402.26\text{--}702.67)\times 10^{-6}$, indicating REE enrichment to different degrees in the ores. The chondrite-normalized REE patterns of basalts in the mining district are of right-inclined type and most samples have much similarity to the basalts in the mining strict, indicating that both of them have strong affinities (Wang et al., 1989). The LREE values are within the range of $(112.57\text{--}529.15)\times 10^{-6}$ and the HREE values are within the range of $(35.52\text{--}244.26)\times 10^{-6}$; the LREE/HREE ratios vary between 1.41 and 4.01 with an average value of 2.71, indicative of LREE enrichment. That is

to say, anatase was formed in a more alkaline environment, thus leading to further separation of LREE from HREE.

(3) The δEu values of basalts in the mining district are 0.86–0.95, all less than unity, and the basalts show weak negative Eu anomalies. Their δCe values are 0.92–0.97, all less than unity, indicating that the major minerals plagioclase and pyroxene in the basalts simultaneously crystallized in proportion. The δEu values of ores in the three orebodies are 0.79–0.93, all less than unity, indicative of weak negative Eu anomalies. Their δCe values are 0.48–1.44, mostly within the range of 0.48–0.88 and a few are within the range of 1.01–1.44. It can be seen that the values vary over a relatively wide range. It is indicated that ore-forming materials of the deposit sourced from basalts, but in the process of metallogenesis, the decomposition of pyroxene took place in the low-temperature and low-pressure weakly alkaline waterbody (Winchell, 1953; Doucet and Synthese, 1967) and in the process of weathering the depletion extent is different very obviously.

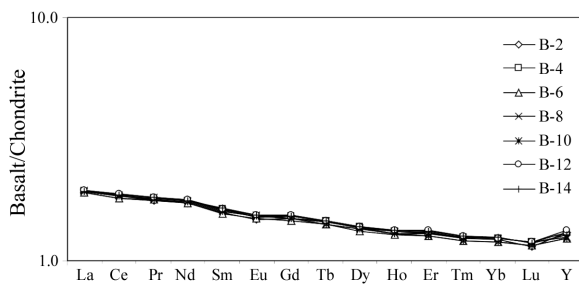


Fig. 1. REE distribution patterns of basalts in the Shazi anatase ore-field, Qinglong.

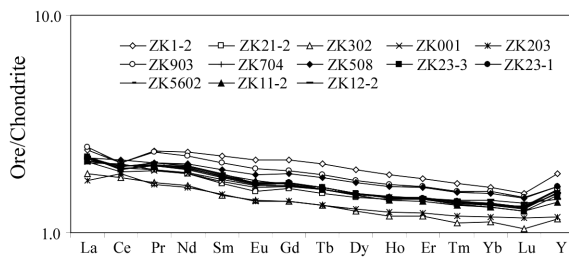


Fig. 2. REE distribution patterns of ores from the Shazi anatase ore deposit, Qinglong.

5 Trace element geochemical characteristics

Trace element analysis was carried out on 13 ore-exposed drill-core samples which had been selected randomly from the ore deposit and the analytical results are listed in Table 3. The average trace element values were statistically calculated, compared with those of the continental upper crust to work out

the element concentration clark values. By comparison of 24 trace elements, Au, Ag, As, Hg, Sb, Tl, V, Ti, Fe, Mn, Sc, Th, Cu, Cr and Co are obviously enriched. Correlation analysis was conducted of those obviously enriched trace elements in the ores. Their correlativities are listed in Table 4.

As viewed from the above statistical results, it can be seen that anatase ores have the following essential characteristics:

(1) Coherent elements such as Au, Ag, As, Hg, Sb, Tl, V, U, Pb, Fe, Mn, Th, Cu, Cr and Co are relatively enriched in the ores, but are not enriched to such an extent that they can be comprehensively utilized.

(2) According to the results of correlativity analysis for trace elements in the ores ($N=13$), with the correlation coefficient of 0.6 at the positive correlation level, the coherent elements can obviously be divided into two groups:

Group I: Au-Ag-As-Sb-Hg-Tl association. This element association is the same as the trace element association in the laterite-type gold ores in the southwestern part of Guizhou Province, reflecting that the formation of anatase is consistent with that of the siliceous claystones vastly distributed at the top of the Maokou Formation limestones at the early stage of eruption of regional-background Emeishan basaltic magmas. Those siliceous claystones are the major country rocks for the formation of the laterite-type gold ores and anatase ores.

Group II: Sc-TiO₂-Cu-Fe-Mn association, reflecting the characteristic geochemical environment of a confined waterbody under regional background condition, i.e., the confined waterbody in the strong oxidation environment on the earth's surface, basaltic eruptive material rich in Fe, Mn, Sc and Ti fell into the waterbody and hydrolyzed to form a low-temperature and low-pressure and weakly alkaline water environment (Nie, 2008; Chen et al., 2003; Cao, 1991). Ferrous iron in the original basalts was oxidized as trivalent iron to form limonite, divalent manganese in the original basalts was oxidized as trivalent or tetravalent manganese to form psilomelane, and titanium was involved in the formation of anatase under sufficient oxygen supply and low-temperature and low-pressures conditions (Jackson et al., 2006; Chen and Ji, 1985).

6 The element (Sc) in the ores

The element Sc is obviously enriched in the ores of the three orebodies and its contents vary over a range of $(24.7-53.8) \times 10^{-6}$, with an average value of 40.3×10^{-6} , 3.7 times those of the continental upper crust (Table 4). The atomic number of Sc is 21 and its outer electronic configuration is $3d^1 4s^2$. In nature Sc

Table 2 Results of REE analysis and characteristic parameters for anatase ore and basalt in the Shazi anatase ore deposit, Qinglong ($\times 10^6$)

| Sample No. | Analytical result | | | | | | | | | | | | | | LREE/HREE | δ Eu | δ Ce | (La/Yb) _N | (La/Sm) _N | (Gd/Yb) _N | | | | | |
|--------------|-------------------|-------|--------|-------|--------|-------|-------|-------|------|-------|------|-------|------|------|-----------|-------------|-------------|----------------------|----------------------|----------------------|------|--------------|------|------|------|
| | La | Ce | Pr | Nd | Sm | Eu | Gd | Tb | Dy | Ho | Er | Tm | Yb | Lu | | | | | | | Y | Σ REE | HREE | | |
| No.1 orebody | ZK1-2 | 78.35 | 103.00 | 28.60 | 136.00 | 35.70 | 10.70 | 38.00 | 5.56 | 28.70 | 5.08 | 12.60 | 8.67 | 1.11 | 143.00 | 636.61 | 392.35 | 244.26 | 1.61 | 0.88 | 0.52 | 6.09 | 1.98 | 1.38 | 3.54 |
| | ZK21-2 | 46.71 | 79.23 | 10.88 | 44.95 | 9.60 | 2.57 | 10.31 | 1.55 | 9.02 | 1.88 | 5.54 | 4.59 | 0.69 | 74.20 | 302.48 | 193.94 | 108.54 | 1.79 | 0.79 | 0.82 | 6.86 | 1.54 | 3.06 | 1.81 |
| | ZK302 | 22.97 | 50.96 | 6.13 | 26.43 | 6.01 | 1.89 | 6.51 | 1.02 | 5.79 | 1.12 | 3.26 | 4.42 | 0.37 | 27.88 | 163.56 | 114.40 | 49.16 | 2.33 | 0.92 | 1.31 | 5.57 | 1.17 | 2.41 | 1.89 |
| | ZK001 | 40.47 | 63.91 | 10.52 | 44.56 | 10.32 | 3.05 | 11.23 | 1.79 | 10.06 | 2.06 | 5.67 | 4.63 | 0.65 | 65.76 | 275.48 | 172.84 | 102.64 | 1.68 | 0.86 | 0.73 | 5.89 | 1.65 | 2.47 | 1.96 |
| No.2 orebody | ZK203 | 17.16 | 59.18 | 5.62 | 24.63 | 6.27 | 1.81 | 6.38 | 1.03 | 6.18 | 1.25 | 3.60 | 3.19 | 0.49 | 29.54 | 166.84 | 114.67 | 52.17 | 2.20 | 0.87 | 1.44 | 3.63 | 0.76 | 1.72 | 1.62 |
| | ZK303 | 94.12 | 102.19 | 26.81 | 110.49 | 23.85 | 6.80 | 21.65 | 3.35 | 17.61 | 3.27 | 8.89 | 7.27 | 0.95 | 84.74 | 513.12 | 364.26 | 148.86 | 2.45 | 0.90 | 0.48 | 8.73 | 2.40 | 2.48 | 2.40 |
| | ZK304 | 53.66 | 75.26 | 12.95 | 54.07 | 12.17 | 3.47 | 12.56 | 1.97 | 10.75 | 2.09 | 5.86 | 4.80 | 0.68 | 70.69 | 321.74 | 211.58 | 110.16 | 1.92 | 0.85 | 0.57 | 7.54 | 1.86 | 2.77 | 2.11 |
| | ZK308 | 48.73 | 117.29 | 15.46 | 69.77 | 17.14 | 5.17 | 18.98 | 2.88 | 16.29 | 3.11 | 8.55 | 1.11 | 6.83 | 0.91 | 85.24 | 417.46 | 273.56 | 143.90 | 1.90 | 0.87 | 1.32 | 4.81 | 1.08 | 1.79 |
| No.3 orebody | ZK23-3 | 51.46 | 70.07 | 13.91 | 58.18 | 12.77 | 3.51 | 12.59 | 1.94 | 10.59 | 2.01 | 5.66 | 4.23 | 0.61 | 60.26 | 308.50 | 209.89 | 98.61 | 2.13 | 0.84 | 0.52 | 8.21 | 1.91 | 2.54 | 2.40 |
| | ZK23-1 | 47.44 | 87.04 | 12.95 | 56.87 | 13.26 | 3.75 | 12.81 | 1.97 | 10.74 | 2.08 | 5.86 | 4.84 | 0.69 | 64.22 | 325.31 | 221.31 | 104.00 | 2.13 | 0.87 | 0.83 | 6.60 | 1.42 | 2.25 | 2.13 |
| | ZK3602 | 53.88 | 83.78 | 14.94 | 63.33 | 14.30 | 3.61 | 12.79 | 1.82 | 10.22 | 2.05 | 5.92 | 4.63 | 0.67 | 63.68 | 336.39 | 233.84 | 102.55 | 2.28 | 0.80 | 0.70 | 7.84 | 1.68 | 2.37 | 2.23 |
| | ZK11-2 | 43.38 | 87.34 | 13.91 | 61.86 | 14.01 | 3.96 | 12.12 | 1.81 | 10.03 | 1.88 | 5.20 | 4.32 | 0.59 | 46.84 | 307.96 | 224.46 | 83.49 | 2.69 | 0.91 | 0.85 | 6.77 | 1.29 | 1.95 | 2.26 |
| Basalt | ZK12-2 | 38.78 | 94.07 | 11.02 | 46.98 | 11.24 | 3.26 | 11.46 | 1.78 | 10.22 | 2.06 | 5.84 | 5.35 | 0.77 | 52.68 | 296.37 | 205.36 | 91.02 | 2.26 | 0.87 | 1.38 | 4.89 | 1.07 | 2.17 | 1.73 |
| | B-2 | 25.30 | 52.10 | 7.19 | 32.30 | 7.34 | 2.22 | 8.15 | 1.31 | 7.42 | 1.42 | 4.11 | 3.45 | 0.46 | 39.10 | 192.41 | 126.45 | 65.96 | 1.92 | 0.87 | 0.92 | 4.94 | 1.27 | 2.17 | 1.91 |
| | B-4 | 26.60 | 55.60 | 7.82 | 34.40 | 8.26 | 2.42 | 8.76 | 1.36 | 7.84 | 1.52 | 4.29 | 3.61 | 0.52 | 40.30 | 203.79 | 135.06 | 68.74 | 1.96 | 0.86 | 0.92 | 4.97 | 1.25 | 2.02 | 1.96 |
| | B-6 | 24.70 | 52.40 | 7.33 | 31.40 | 7.05 | 2.28 | 7.54 | 1.22 | 6.74 | 1.34 | 3.81 | 3.29 | 0.48 | 33.50 | 183.53 | 125.15 | 58.38 | 2.14 | 0.95 | 0.93 | 5.07 | 1.23 | 2.21 | 1.85 |
| Basalt | B-8 | 27.90 | 58.50 | 8.02 | 34.20 | 8.12 | 2.54 | 8.08 | 1.35 | 7.53 | 1.49 | 4.08 | 3.57 | 0.51 | 34.70 | 201.38 | 139.50 | 61.89 | 2.25 | 0.95 | 0.93 | 5.28 | 1.24 | 2.17 | 1.83 |
| | B-10 | 26.20 | 56.60 | 7.51 | 32.50 | 7.67 | 2.43 | 7.94 | 1.24 | 7.10 | 1.41 | 3.88 | 3.26 | 0.48 | 33.30 | 192.07 | 132.96 | 59.11 | 2.25 | 0.95 | 0.96 | 5.42 | 1.21 | 2.15 | 1.97 |
| | B-12 | 28.10 | 61.30 | 8.07 | 36.20 | 8.25 | 2.53 | 8.90 | 1.38 | 7.85 | 1.58 | 4.55 | 3.60 | 0.53 | 42.50 | 215.86 | 144.40 | 71.46 | 2.02 | 0.90 | 0.97 | 5.26 | 1.19 | 2.14 | 1.99 |
| | B-14 | 26.50 | 58.70 | 8.04 | 34.80 | 8.61 | 2.58 | 8.62 | 1.39 | 7.75 | 1.56 | 4.42 | 3.67 | 0.51 | 36.80 | 204.52 | 139.16 | 65.36 | 2.13 | 0.91 | 0.96 | 4.87 | 1.18 | 1.94 | 1.89 |

is present stably in the form of Sc^{3+} (Liu et al., 1984). Mainly unequivalently replacing Fe^{2+} and Mg^{2+} , Sc^{3+} is present in pyroxene, hornblende and olivine (Lü et al., 1992). In the mining district Sc^{3+} is present mainly in pyroxene minerals in basalts and coexists isomorphically with Fe, Mn and Ti. When the eruptive materials of basaltic magmas fell into the waterbody, Sc^{3+} would be released from rocks along with the weathering and disintegration of pyroxene and other minerals, forming $\text{Sc}(\text{OH})_3$ or Sc_2O_3 which would be adsorbed on colloidal clay minerals. That is why anatase in the mining district is enriched in Sc. As Fe, Mn, Ti and Sc are derived from the same basalt, the correlativities between Sc and Ti, Fe and Mn are relatively high. The coefficient of correlation between Sc and Fe is +0.9155, that between Sc and Mn is +0.7268, and that between Sc and Ti is +0.6568.

The nature of Sc is very close to that of REE. As compared to La and Y, the ionic radius of Sc is particularly small. The alkalinity of hydroxides is extremely weak. Therefore, when hydroxides are mixed with REE, it is possible to isolate Sc from ores by using the ammonia grading precipitation method or the nitrate grading decomposition method. Sc can play a fantastic role in metallurgical, electronic, chemical and optical power and other material industries and it has become a new star in the field of material science. Sc is very expensive. Sophisticated Sc-purification technologies are now available both at home and abroad. A great success has been achieved on an experimental basis in extracting Sc from specimens with a Sc content of 0.0017% (Feng and Zhang, 2005). The total REE amount of this anatase ore deposit does not reach the industrial grade of the weathering-crust ion absorption-type REE ores, but the total REE amount of some samples reaches $(402.26\text{--}702.67)\times 10^{-6}$, indicative of a certain degree of mineralization. So Sc is relatively high and can be comprehensively utilized. With the development of anatase ores and the comprehensive utilization of Sc, the economic value of Sc is different to estimate.

7 Conclusions

The following conclusions have been drawn from the REE and trace element geochemical studies of oxides in ores from the Shazi anatase ore deposit in Qinglong.

(1) The total amount of major oxides SiO_2 , Al_2O_3 , Fe_2O_3 and TiO_2 in the ores is close to that of oxides in laterite in the laterite-type gold deposits in the western part of Guizhou and in laterite weathering crust in Guizhou, though the content of TiO_2 is relatively high. The ores are strongly weathered and completely oxi-

dized. There is a positive correlation between TiO_2 and Fe_2O_3 and the correlation coefficients are 86.21%, respectively, reflecting that anatase is associated (coexists) with clay minerals in the protolith.

(2) According to the correlation analysis of trace elements in the ores, the correlative elements can be divided into two groups: Group I: Au-Ag-As-Sb-Hg-Tl association. This element association is identical to the trace element association in the laterite-type gold ores in the southwestern part of Guizhou, reflecting that the formation of anatase is consistent with the formation of siliceous claystones vastly developed at the top of the Maokou Formation limestones during the early stage of regional- background Emeishan basaltic magmas. Siliceous claystones of this type are the major country rocks for the formation of laterite-type gold ores and anatase ores. Group II: Sc-TiO₂-Cu-Fe-Mn association, reflecting the characteristic geochemical environment of the confined waterbody under regional background condition, i.e., in the confined waterbody in the strongly oxidized zone on the Earth's surface, eruptive materials from basalt magmas enriched in Fe, Mn and Ti fell into the waterbody, followed by hydrolysis, thus creating a low-temperature and low-pressure weakly alkaline water environment. Ferrous iron in the original basalts was oxidized as trivalent iron to form limonite, and divalent manganese in the original basalts was oxidized as trivalent or tetravalent manganese to form psilomelane. The formation of anatase took place in the low-temperature and low-pressure weakly alkaline water environment where oxygen gas supply is sufficient.

(3) Research on the REE geochemical characteristics of basalts and ores in the three orebodies in the study region indicate that both basalts and ores have extremely strong affinities. The ore-forming materials of the deposit come from basalts. In the study region the anatase ore deposit is enriched in Sc, Fe, Mn, Ti and Sc which all sourced from the same type of basalts, mainly from decomposition of pyroxenes in the basalts. Therefore, Sc possesses a very strong correlativity with Ti, Fe and Mn.

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