REE characteristics of anhydrite in the Nihe magnetite-apatite deposit and its relation to mineralization, Middle-Lower Yangtze River Valley metallogenic belt

Yu Fan*, Yinan Liu, Taofa Zhou, Noel C White, and Lejun Zhang

School of Resources and Environmental Engineering, Hefei University of Technology, Hefei, Anhui, China, *e-mail, fanyu@hfut.edu.cn

Nihe is a major magnetite-apatite deposit located in the northwestern part of the Lu-Zong volcanic basin in the Middle-Lower Yangtze River Valley Metallogenic Belt. Anhydrite is extensively developed in the deposit. The anhydrite can be divided into three types on the basis of mineral assemblage and stage: Type I is tabular violet crystals coexisting with diopside, magnetite, pyrite, and apatite; Type II is tabular white crystals coexisting with pyrite; and Type III is equant pink or white crystals in carbonate-anhydrite veins. We present *in situ* LA-ICP-MS geochemical data for the three different types of anhydrite in the Nihe deposit. From fluid inclusion analysis, the three anhydrite forming environments range from high temperature, highmedium salinity, to low temperature, low salinity. Type I and type II anhydrite was precipitated under near-equilibrium conditions. Type I anhydrite coexists with magnetite, and type II coexists with type I pyrite, indicating that fS_2 in the system increased with decreasing temperature. Type III anhydrite formed at less than 200°C with no associated oxide and sulfide minerals, LA-ICP-MS data indicate that type I anhydrite contains a much higher REE concentration than the other two types, and the REE patterns and Eu anomalies are also quite different (LaN/YbN=1.1-466.3, $\delta Eu=0.3-2.4$). The main factor controlling the pattern of REEs in anhydrite is deposition of the REE-rich minerals (apatite, titanite, epidote). After precipitating the REE-rich minerals, the REE content greatly decreased in the fluid, resulting in a lack of REEs in types II and III, and their REE patterns are flat. In addition, decreasing fluid salinity caused lower Cl⁻/SO₄²⁻ ratios in later stages. Furthermore, some type III anhydrite has elevated LREEs, suggesting some late-stage dissolution and reprecipitation of anhydrite due to fluctuating fluid compositions. Addition of abundant anhydrite to the ore-forming fluid increases fO₂, allowing iron to be transported as FeCl₄⁻ to the magnetite deposition site.