모나자이트 침출액으로부터 Primene-JM-T를 이용한 토륨의 용매추출

전중혁 · 김홍인 · 이진영 · §조티 라제쉬 쿠마

한국지질자원연구원 DMR융합연구단

Solvent Extraction of the Thorium from Monazite Leaching Solution by Primene-JM-T

Jong-Hyuk Jeon, Hong-In Kim, Jin-Young-Lee and §Rajesh Kumar Jyothi

*Convergence Research Center for Development of Mineral Resources(DMR),
Korea Institute of Geoscience and Mineral Resources (KIGAM), Daejeon, Korea*

요 약

본 연구는 모나자이트 침출액으로부터 용매추출을 통한 토륨의 추출 및 분리 가능성에 대해 연구하였다. 토륨 용매추출 공정을 위해 1차 아민계 추출제인 Primene-JM-T가 선택되었다. 토륨의 액-액 추출 공정을 위해 다양한 조건의 실험이 수행되고 조건이 확립되었다. 연구 결과로부터 추출제 검증, pH 조건, 추출제의 변화 및 추출 등온선의 작성 및 최종적으로 추출제로부터의 탈거 조건을 확립하였다.

주제어 : 모나자이트, 용매추출, Primene-JM-T, 토륨

Abstract

Thorium extraction and possible separation from monazite leaching solution was studied. Primary amine Primene JM-T was selected for thorium extraction processing. Various experiments were tested and established for the thorium liquid -liquid extraction process. The screening of extractant, lower pH conditions, extractant variation and extraction isotherms construction, and finally, stripping studies were established.

Key words : Monazite, solvent extraction, Primene-JM-T, Thorium

1. Introduction

Six major issues are most concern in new millennium fourth industrial revolution development. Those are global energy resources (Fig. 1), global electricity production, global accessibility and non-accessibility of electricity,

lifecycle of greenhouse gas emissions, global non-renewable energy resources and global nuclear reactors scenario. Primary energy supply was predicting 82 to 76% reduction, increasing the renewables from 10 to 16% hydro and nuclear energy utilization was not much change in year 2020¹⁾. By using lignite, coal, oil natural

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§ Corresponding Author : Rajesh Kumar Jyothi (E-mail : rkumarphd@kigam.re.kr)

Convergence Research Center for Development of Mineral Resources, Korea Institute of Geoscience and Mineral Resources, 124 Gwahak-ro, Yuseong-gu, Daejeon 34132, Korea

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Fig. 1. World energy resources total proved reserves¹⁾.

gas etc generating the high carbon emissions whereas biomass, nuclear, hydro and wind energy sectors will generate very low carbon emissions when compared with fossil fuel energy sources¹⁾.

Nuclear power mainly depends on the uranium and thorium resources. A uranium nuclear reactor functions by flowing water over the surface of the zirconium metal tubes with uranium fuel sealed inside the fuel rods. Much radioactive material is possible to contaminate in air. Humans and all living animals get effected by this radiation. Whereas liquid fluoride thorium reactor (LFTR) are generate 0.01% of waste only as well as generate 1,000 MW of electricity from 0.9 MT of natural ThO₂. Moreover, proliferation material is zero, fission products such as iodine etc can chemical process at on-site. In addition, actinides such as neptunium etc will generated up to 1 kg, which can be separated and recycled onsite. Three major factors are exposes that, thorium is better nuclear sources; those are thorium is more abundant in nature than uranium, it is a fertile rather than fissile and chemically distinct (easy to removal the radioactive materials). Finally, molten salt reactors are well suited to thorium fuel as normal fuel fabrication is avoided²⁻⁵⁾.

For thorium processing from monazite, one of the better technique is hydrometallurgy. Various reports are available in past literature on thorium extraction from monazite or synthetic liquors⁶⁻¹²⁾. The present study is focused mainly on as much as thorium extraction with

less interfere metals and experimental conditions for liquid-liquid extraction processing was optimized.

2. Experimental

2.1. Reagents and apparatus

Aqueous sample analysis of metals such as thorium, cerium and others were analyzed by using in ICP-OES manufactured by Thermo Scientific, USA model iCAP 6000 Series. The commercial grade amine based extractant such as Primene JM-T amine was supplied by Cognis Corporation USA and used as it is without purification. The present study we were used diluent as A150 (commercial form of kerosene) supplied by Samsung, Korea and all other reagents used were analytical reagent grade. The aqueous feed of monazite was used.

2.2. Liquid-liquid extraction process

Metal distribution ratios were determined by shaking equal volumes (30 mL each) of aqueous and organic phases for 30 min in a separating funnels using a mechanical shaker at 25±0.5°C. Preliminary experiments showed that the extraction equilibrium was attained within 5 min for thorium. The solutions were then allowed to settle, the phases separated, and thorium and other metals in the aqueous phases determined by ICP-OES. The concentration of metal ion in the organic phase was then obtained by mass balance. The distribution ratio, D was defined as the ratio of the concentration of metal ion in organic phase to that in the aqueous phase ($D = \text{metal in organic phase} / \text{metal in aqueous phase}$). The percentage extraction (% E) calculated by the following equation: % $E = (D \times 100 / D + 1)$ A:O. The general agreement between the distribution ratios values obtained was within ± 5%. The present investigations for thorium separations using amine based extractants, following experimental conditions are constant for the all experiments: time = 5 min, temperature = 25°C and aqueous: organic (A:O) = 1:1 that means unit phase ratio.

Fig. 2. Screening of the extractant (Experimental conditions: time 30 min, temperature 25°C, phase ratio, one; aqueous feed contains: 2720 mg/L of Th, 150 mg/L of Fe, 1603 mg/L of La and 1913 mg/L of Ce).

3. Results and discussions

3.1. Influence of extractant concentration on thorium extraction

Monazite leaching solution contained 2720 mg/L of thorium, 150 mg/L of iron, 1603 mg/L of lanthanum and 1913 mg/L of cerium. The first screening test was conducted with commercial extractant Primene JM-T; it was varied from 0.1 mol/L to 1.0 mol/L. The other experimental conditions such as phase ratio (Aqueous (A) / Organic (O)) was one and temperature is 25°C. The obtained extraction was presented in Fig. 2. The data was make known that, with 0.1 mol/L of Prime JM-T 48.3 % of Th was extracted, after that both the cases 0.5 and 1.0 mol/L of Prime JM-T, Th was quantitatively complete extracted (100%). Whereas, other associated metals were not extracted with Prime JM-T up to 0.5 mol/L concentration and with 1.0 mol/L Prime JM-T 11.2 % of La and 23.5 % of Ce were co-extracted with thorium.

Based on the first experiment, Primene JM-T was selected for further thorium extraction process development. The extractant concentration was varied from 0.1 to 1.0 mol/L concentration. The significant extraction

Fig. 3. Effect of extractant (Primene JM-T) (Experimental conditions: time 30 min, temperature 25°C, phase ratio, one; aqueous feed contains: 3926 mg/L of Th, and 15080 mg/L of Ce for pH -0.49 and pH 0.12 4096 mg/L of Th and 15750 mg/L of Ce).

was observed mainly of thorium and cerium (Fig. 3). With 0.1 mol/L of Primene JM-T up to 68% of thorium was extracted, whereas cerium was extracted less than 1.2%. On the other hand, from 0.5 mol/L to 1.0 mol/L of Primene JM-T concentration thorium was 100% extracted, whereas cerium was extracted up to 9.7% at initial pH of feed was -0.49, at pH 0.12 cerium was extracted 26.2% with 0.5 mol/L of Primene JM-T and 58.7% with 1.0 mol/L of Primene JM-T. The data indicates that lower acidic region (-0.49 pH) is the much promising to thorium, in addition cerium co-extraction was very nominal (less than 1.2%).

3.2. Primene JM-T treated with 50% sulfuric acid

Primene JM-T was treated with 50% sulfuric acid to neutralize the extractant basic nature. Further lower concentration of the Primene JM-T and 50% H_2SO_4 -Primene JM-T were tested. Both the extractants were varied from 0.01 to 0.1 mol/L of concentration, at two different initial pH conditions such as -0.49 and 0.12, the obtained data was presented in Fig. 4. Cerium extraction was more favorable with acidified extractant and pH 0.12 feed condition.

Fig. 4. Effect of extractant (Primene JM-T) treated with sulfuric acid (Experimental conditions: time 30 min, temperature 25°C, phase ratio, one; aqueous feed contains: 3926 mg/L of Th, and 15080 mg/L of Ce for pH -0.49 and for pH 0.12, 4096 mg/L of Th and 15750 mg/L of Ce).

Fig. 5. Extraction isotherm for thorium extraction from monazite leaching solution.

3.3. Extraction isotherm (McCabe Thiele diagrams) for thorium

Two different concentration of the extractant was tested for extraction isotherm construction for thorium extraction. With 0.3 mol/L of Primene JM-T with phase ratio (A/O) 4 needs three extraction stages to load the above 9 g/L of Th from aqueous feed containing 2301 mg/L. Whereas with same feed, above 11 g/L needs to load the two extraction stages at phase ratio 5 with 0.5 mol/L of Primene JM-T (Fig. 5).

3.4. Stripping of thorium from loaded organic phase

The stripping of the targeted metal is most significant process in hydrometallurgical process development. The present study the loaded organic phase was water washed by two or three times then followed by acidified water treatment to remove un-wanted metals from thorium then finally applied the thorium stripping by using 1 mol/L of hydrochloric acid was able to recover the thorium more than 90% at single stripping stage. It was indicated that

multiple stripping stages could able to strip the loaded thorium.

4. Conclusions

The present study was concluded that, Primene JM-T is the best extractant for extract the thorium from monazite leach liquors. Moreover, very low acidified leach liquor (-0.49 pH) condition is more suitable for thorium complete extraction with very nominal cerium co-extraction. The water washings and diluted sulfuric acid solutions could able to remove the all impurities from loaded organic phase. The most suitable reagent is HCl to recover the loaded thorium, successfully.

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전 종 혁

- 광운대학교 화학공학과 공학박사
- 한국지질자원연구원 박사 후 연구생
- 당 학회지 제29권 2호 참조

김 홍 인

- 한국과학기술연합대학원대학교 자원순환공학과 공학박사
- 현재 한국지질자원연구원 DMR융합연구단 책임연구원
- 당 학회지 제24권2호 참조

이진영

- 현재 한국지질자원연구원 책임연구원
 - 당 학회지 제22권 1호 참조
-

라제쉬 쿠마 조티

- Sri Venkateswara University(India) 이학박사
 - 현재 한국지질자원연구원 DMR융합연구단 책임연구원
 - 당 학회지 제29권 4호 참조
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