Solvent Extraction of Actinide Elements in New Extraction System for TRU Recovery (NEXT)

Development of Advanced Reactor Fuel Reprocessing

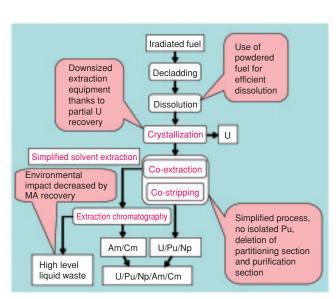


Fig.1-13 Outline of the NEXT Process

The NEXT process is a simplified conventional PUREX process aiming for economic competitiveness. MAs are recovered and loaded as MOX fuel of the advanced reactor for efficient utilization of resources and low environmental impact.

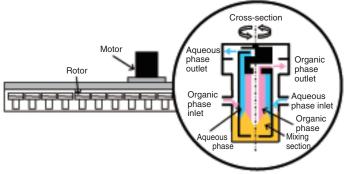


Fig.1-14 Type CPF centrifugal contactor

Centrifugal contactors separate organic phase from aqueous phase by centrifugal force. These have large throughput and low degradation of solvent in comparison with mixer-settlers because residence time is very short.

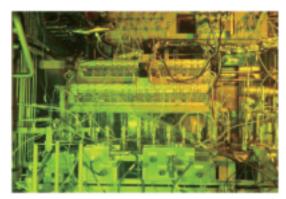


Fig.1-15 Centrifugal contactors in hot cell

We have been studied an advanced aqueous reprocessing process, named New Extraction System for TRU Recovery (NEXT) (Fig.1-13). The NEXT process basically consists of Uranium (U) crystallization, simplified solvent extraction of U by the extractant tri-n-butylphosphate (TBP), Plutonium (Pu) and Neptunium (Np) co-recovery, and Americium (Am) and Curium (Cm) recovery with extraction chromatography technology.

In the NEXT process, it is desirable to recover not only U and Pu but also minor actinides (MAs) for loading them in MOX fuel of the advanced reactor. This will be effective in decreasing environmental impact because it brings about the reduction of long-lived radioactive elements in the wastes.

Among the MAs, Am and Cm are poorly extracted by TBP. Therefore, these elements need to be separated from high level waste using a new extractant which has stronger affinity to MAs than TBP. On the contrary, by proper control of Np valence, it is possible to recover Np with U and Pu by TBP. Assuming that a rather low decontamination factor of the reprocessing product is made possible by the fast reactor fuel cycle, partitioning section and purification section are deleted, and U, Pu and Np co-recovery flow sheet is designed by single cycle.

Experimental studies using irradiated fuel have been carried out at the Chemical Processing Facility (CPF) with centrifugal contactors, whose residence time is considerably smaller than that of mixer-settler (Fig.1-14, Fig.1-15). The feed solution is adjusted to have high nitric acid concentration ([HNO₃]). This condition adjusts the Np valence so that it is extractible by TBP. As expected, about 99% of Np was recovered with U and Pu in this process. Through this series of studies, a U, Pu and Np co-recovery process using high [HNO₃] feed solution was successfully demonstrated.

Reference

Nakahara, M. et al., U, Pu and Np Co-recovery in Simplified Solvent Extraction Process—The Extraction Behavior of Np at the Condition of High HNO3 Concentration Feed Solution and Scrubbing Solution -, 2006, JAEA-Research 2006-030, 43p. (in Japanese).