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Transient release of radioactive iodine from the fission of UF₄ in 2LiF–BeF₂ salt

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In this study, the behavior of fission product iodine released from the melting process of a mixture consisting of UF₄ irradiated with neutrons and 2LiF–BeF₂ (FLiBe) salt was studied. The experiment showed that a large amount of iodine was released immediately during melting and captured by Ni metal foils. The transient release of iodine observed in this experiment is attributed to the redox reaction between the hot atoms of the fission product iodine that cumulated due to long-time irradiation. The effect of the redox status of the molten salt on the transient release of iodine was also investigated. Based on this investigation, it was proposed that the activity ratios of ¹³¹I to salt-seeking fission products in the fuel salt, as an effective diagnostic criterion, may be used for the surveillance of the redox potential of fuel salts in a molten salt reactor.

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Introduction

Nuclear fission produces a variety of fission products. In a solid fuel reactor, these fission products are restricted to fuel elements together with fissile and fertile materials. Differently from the solid fuel reactor, the fission products and fission materials circulate in the primary loop of the reactor, together with FLiBe molten salt, used as coolants in the molten salt reactor (MSR).¹ Thus, the effect of the fission products on MSR such as neutron economy, delayed neutron fraction, control of the reactor and corrosion of structure materials has received considerable attention.^{2,3} The study of the behavior and distribution of the fission products in MSR becomes a significant research project in the development of MSR.^{4,5}

Up to now, the chemistry of fission products in aqueous solutions has been investigated quite well.⁶ These research studies have provided strong scientific and technical supports for the development and optimization of spent fuel reprocessing. Similarly, the study of the chemistry of fission products in molten salts should be important not only for the processing of irradiated fuels but also for the design, operation, and safety control of the MSR. Unfortunately, reports on the chemistry of fission products in molten salts rarely exist. In the 1960s, Oak Ridge National Laboratory (ORNL) started the research and development of the molten salt reactor. The 8 MWt Molten Salt

Reactor Experiment (MSRE) facility was built in 1966 and successfully operated for nearly 4 years.^{7,8} During the operation of the MSRE, the chemistry of fission products in molten salts was one of the important research subjects conducted by the chemistry division of ORNL. Based on their behavior and distribution, in the research at ORNL, the fission products were divided into three categories: (1) noble gas fission products (Kr and Xe), which are slightly soluble in salts and easily escapable from the fuel salt at the MSRE operating temperature, (2) salt-seeking fission products (Rb, Cs, Sr, Ba, Y, Zr and the lanthanides), which could exist stably in the fuel salt in the form of fluorides and (3) noble metal fission products (Nb, Mo, Tc, Ru, Rh, Pd, Ag, Sb, and Te), which are unstable in salt and whose behavior was very complicated. In most cases, they tend to be reduced into metals and deposited on various surfaces of the MSRE materials such as graphite as the neutron moderator and Hastelloy-N as the fuel container. The research at ORNL found that their behavior and distribution are often affected by the redox potential of the fuel salt. Thus, their behavior and dependence on the redox potential, especially for ⁹⁵Nb, has become one of the key research objects.^{9,10}

Iodine isotopes are significant fission products because of their high yield and wide mass numbers involved in mass distribution. Moreover, they are also the precursor nuclei both for delayed neutron emitters and neutron poison ¹³⁵Xe. Their behavior was correlated closely with the operation and control of the MSR. Compared with the typical salt-seeking and noble metal fission products, however, less iodine-related investigations were reported by ORNL. Using γ -ray spectrometry, ORNL obtained some information about the behavior and distribution of the fission product iodine *via* detection of the fuel salt, gas

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vapor and the alloy/graphite surveillance specimens inserted into the pump bowl.^{7,11} According to the reports from ORNL, the fission product iodine was categorized sometime in the halogen group, like bromine, but in most cases, categorized as a subgroup together with Sb and Te in the noble metal fission product group. Regardless of how the fission product iodine was grouped, the results from the analysis for γ -spectra appeared very often to be contradictory. ORNL pointed that fission product iodine seemed to exist steadily, in the form of I^- , in fuel salts at the redox potential of the MSRE.¹¹ However, the radioactive measurements of the fuel salt samples indicated that the ^{131}I radioactivity relative to the inventory of irradiated UF_4 was between 8% and 113%, with most values ranging from 30% to 60%.⁷ The phenomenon of the iodine loss from the fuel salt was very similar to that of the noble metal fission product. However, the analysis of the γ -spectra for the surfaces of alloy and graphite specimens indicated no noticeable deposition of fission product iodine to be found except for the great deposition of typical noble metal fission products such as Mo, Te, Nb and Ru. Considering the high volatility of the iodine element, the gas samples from the pump bowl and the off-gas system were carefully measured many times with γ -ray detectors; however, the existence of fission product iodine was not found except for occasionally detectable in the pump bowl gas samples.^{11,12} It can be seen that, iodine is indeed an elusive fission product in MSR, and it is interesting and necessary to understand the reason for iodine loss up to about one-fourth to one-third.

In this work, the UF_4 powder was irradiated by a photo neutron source driven by a 15 MeV electron linear accelerator. The behaviors of iodine and some fission products in the molten salt compound consisting of irradiated UF_4 and FLiBe were investigated under experimental conditions. To understand the reason for the loss of fission product iodine, as observed at MSRE operation, the emphasis of this work was placed on the unexpected release of iodine from the molten salts at the beginning of melting. Moreover, a new way to examine the redox potential of the molten salt was proposed on the basis of the activity measurement of ^{131}I .

Experimental

Reagents and materials

UF_4 was purchased from China National Nuclear Corporation. Li metal (99.9% purity), LiI (99.99% purity) and CeF_4 (99.99% purity) were purchased from Alfa Aesar Co., Ltd. All the reagents were used without further purification. Ar gas with 99.999% purity was purchased from Shanghai Louyang Gas Co. Ltd. The 2LiF-BeF₂ (FLiBe) eutectic salt of 99.9% purity was prepared according to the procedures described in our previous work, and its oxygen content was determined to be less than 400 ppm. Graphite and Hastelloy C276 specimens were used as the surveillance specimens, and the size was $20 \times 5 \times 2$ mm for both of them. The reason for choosing the Hastelloy-C276 alloy as the surveillance specimen in this work was its similarity in components (Ni-58.10%, Mo-15.82%, Cr-15.98%, Fe-5.95%, and W-3.41%) to the Hastelloy-N alloy used in the MSRE. A ^{60}Co

standard source and a ^{152}Eu standard source used to calibrate the gamma-ray spectrometer were purchased from Eckert & Ziegler BEBIG GmbH, Germany.

Equipment

A self-designed reaction apparatus placed in a glove box in Ar atmosphere was used to investigate the behavior of iodine, and the detailed description has been reported in our previous works.^{10,13,14} A graphite crucible with an inner diameter of 25 mm and a height of 60 mm was used as the reaction crucible.

A photo neutron source driven by a 15 MeV electron linear accelerator was self-designed and constructed by Shanghai Institute of Applied Physics, CAS (SINAP);¹⁵ its maximum neutron output of the photo neutron source was about $1.2 \times 10^{11} \text{ n s}^{-1}$, and the average energy of the neutrons was about 1 MeV.¹⁶ The detector used for γ -spectrum measurement (count the nuclides) was an n-type ORTEC GEM304P Hyper Pure Germanium (HPGe) detector, and the energy resolution was 1.85 keV at 1.33 MeV γ -ray. Before measurement of the γ -spectra, the energy calibration and the efficiency calibration of the detectors were obtained using the ^{60}Co and ^{152}Eu standard sources.

Experimental method

First, 20 g UF_4 powder was loaded into a customized PMMA box and then installed on the photo neutron source. The neutron flux was estimated to be about $1 \times 10^8 \text{ n s}^{-1} \text{ cm}^{-2}$ on the target, with total fluence being about $3 \times 10^{13} \text{ n}$. After four-day irradiation and 24 hour cooling, irradiated UF_4 was well mixed, and about 0.5 g powder was sampled accurately and measured using a calibrated γ -ray spectrometer to obtain the original activities (denoted as inventory) of ^{237}U and fission products.

About 4 g of irradiated UF_4 was mixed with 26 g FLiBe eutectic salt in a graphite crucible placed into the reaction apparatus. Two pieces of surveillance specimens (one for graphite and one for Hastelloy C276 alloy) were inserted into the molten salt to examine the possible deposition of fission products on the surface of the Hastelloy C276 alloy and crucible. Before heating, the reaction apparatus was covered with a Ni foil to collect the fission products released from the molten salt. The temperature of the apparatus was then raised up to 650 °C in 2 hours and maintained at 650 °C throughout the experiment. In the process of heating and holding, the Ni foil on the top of the apparatus was removed at intervals as required. At the same time, a small portion of salt was taken out from the crucible using a quartz tube with an inner diameter of 5 mm and weighted accurately. After sampling, a new Ni foil was covered again on the reaction apparatus instead. The Ni foils caught fission products and salt samples were subjected to measure the γ -ray spectra. In order to investigate the effect of the redox potential on the release of iodine, Li metal and CeF_4 were used to change the redox condition of the FLiBe molten salt. Sampling inspection of salt and Ni surveillance specimens was carried out after 24 hours of addition of the Li metal and CeF_4 .



Table 1 Data used for nuclide identification and specific activity calculation and the initial activities in 1 g irradiated UF₄

Nuclides	T _{1/2} (day)	Energy (keV)	Emission intensity	Activity (Bq g ⁻¹ UF ₄)
²³⁷ U	6.75	208.0	0.212	5.50 × 10 ⁴
¹⁴³ Ce	1.377	293.3	0.428	2.26 × 10 ³
¹⁴⁰ Ba	12.75	537.3	0.2439	8.72 × 10 ²
⁹⁵ Zr	64.02	756.7	0.5446	2.43 × 10 ²
¹³¹ I	8.021	364.5	0.817	9.00 × 10 ²
¹³² I	0.096	667.7	0.987	2.88 × 10 ³
¹³³ I	0.867	529.9	0.87	4.21 × 10 ³
⁹⁹ Mo	2.748	181.1	0.0599	2.51 × 10 ³
¹⁰³ Ru	39.26	497.1	0.91	3.11 × 10 ²
¹³² Te	3.204	228.2	0.88	1.46 × 10 ³
¹³⁵ Xe	0.381	249.8	0.9	3.12 × 10 ³
¹⁴⁷ Nd	10.98	531.02	0.1337	3.33 × 10 ²

Measurement and data analysis

All the samples of Ni foils and salt samples were counted using a calibrated γ -ray spectrometer. The counting times varied from 1 hour to 24 hours, depending on the activity of interesting nuclides. The γ -ray spectra were analyzed using the Gamma Vision Program provided by ORTEC Co. Assignments of radioactive nuclides were made mainly on the basis of their characteristic energies. If necessary, half-life and concordance with other γ -rays emitted by the presumed nuclide were used to help the assignments of nuclides. The basic nuclear data used for nuclide assignment and activity calculation were provided by ENDF/B-VII.1 (USA, 2011), and they are listed in Table 1. The uncertainties in activities experimentally determined in this work were standard deviations, including mainly the statistic errors in the γ -ray counts, a 5–6% error in detector efficiency and a 5% error in sample geometry. The typical uncertainties for ¹³¹I were about 10–15%.

Results and discussions

After complete dissolution of irradiated UF₄ in the FLiBe molten salt, a certain amount of molten salt was sampled using a quartz

tube, weighed, and then counted using a γ -ray spectrometer. The initial specific activities at 4 day irradiation and 1 day cooling for over all 12 nuclides were obtained from the measurement and calculation of γ -ray spectra, and the results are also listed in Table 1. In addition, the specific activities of ⁹⁵Zr, ¹³¹I, ¹⁴³Ce, ¹⁴⁰Ba and ²³⁷U, after decay correction, are shown in Fig. 1, as a function of the melting time.

As shown in Fig. 1, the specific activities of the nuclides ⁹⁵Zr, ¹⁴⁰Ba, ¹⁴³Ce and ²³⁷U in molten salt increased gradually and reached a steady state after about 40 hours. The result was consistent with ORNL's study that ⁹⁵Zr, ¹⁴⁰Ba and ¹⁴³Ce fell into the group of salt-seeking fission products and could stably exist in the fuel salt.⁷ ¹³¹I appeared to reach dissolution equilibrium more quickly in the molten salt, which may be attributed to its lower melting point. According to the report of ORNL, iodine was grouped as a noble metal fission product. However, iodine could form iodide and exist in a steady form in the molten salt of the MSRE.

After the dissolution equilibrium, total activities of each radioactive nuclide in the FLiBe molten salt could be calculated on the basis of their specific activities measured in the salt sample. When comparing the total activities with the data of inventories in irradiated UF₄, some interesting information



Fig. 1 Specific activities of ¹³¹I, ⁹⁵Zr, ¹⁴⁰Ba, ¹⁴³Ce and ²³⁷U as a function of melting times.



Fig. 2 Comparison of total activities in the salt with the inventories for ²³⁷U, ¹⁴³Ce, ¹⁴⁰Ba and ⁹⁵Zr.



could be obtained. Fig. 2 shows the total activities of ^{95}Zr , ^{143}Ce , ^{140}Ba and ^{237}U in the molten salt and the relative data of inventories in irradiated UF_4 . ^{95}Zr , ^{143}Ce and ^{140}Ba , belonging to the salt-seeking fission products, should completely dissolve and stably exist in the fuel salt. Consequently, the total activity of each nuclide in the FLiBe molten salt, like that of ^{237}U , was in agreement with those of the inventories in irradiated UF_4 , as expected.

However, the noble metal fission products (^{99}Mo , ^{103}Ru and ^{132}Te), iodine and noble gas fission products (^{131}I , ^{132}I , ^{133}I and ^{135}Xe) exhibited different behaviors, and their total activities in the FLiBe molten salt were much lower than the inventories. Due to little solubility in the fluoride salt and high volatility, noble gas ^{135}Xe was easily released from the molten salt. Hence, no ^{135}Xe fission product existed in the molten salt any longer, as shown in Fig. 3. According to the report of ORNL,⁹ the noble metal fission products tended to deposit, in a form of metal, on the surface of structural materials at the redox potential of the MSRE. Thus, the deposition of ^{99}Mo , ^{103}Ru and ^{132}Te on the wall of graphite crucible would be responsible for the decrease in total activities observed in this work. It is interesting to note that the total activities of ^{131}I , ^{132}I and ^{133}I in salt, similarly to these of ^{99}Mo , ^{103}Ru and ^{132}Te , also became low when compared to the inventories. Although iodine was grouped under noble metal fission products, the decrease in the total activities of the fission product iodine in the salt could not be explained by their deposition on the surface of solid materials.

To investigate the underlying cause for the decrease in iodine activity in the molten salt, two pieces of graphite and Hastelloy C276, used as the surveillance specimens, were inserted into the salt for several hours. After taken out, these specimens were measured using a γ -ray spectrometer. After correction for the additional activity induced by the nuclides in the salt adsorbed on specimens, the real activity of the nuclides deposited on the surface of the specimen was examined. The result indicated that all the activities were due to noble metal fission products ^{99}Mo , ^{103}Ru , ^{132}Te and its daughter ^{132}I , with no activities of ^{131}I and ^{133}I being observed. Moreover, three samples of salt, after cooling, were taken out from the upper,

middle and lower layers of the molten salt in the crucible. The measurements for γ -activity indicated that there was no obvious difference in the specific activity of ^{131}I in the samples from three different regions, implying homogeneous distribution of ^{131}I in the molten salt. Since no deposition on the surface of the specimens and no precipitation at the bottom of the molten salt for fission product ^{131}I were observed, unique possible explanation for iodine loss observed by ORNL should be their release, in the form of volatile gas I_2 from the molten salt.

To verify the explanation for the iodine loss, the irradiated UF_4 and FLiBe salt were mixed in the crucible. Prior to heating, a piece of Ni foil was placed on the top of the reaction apparatus. In the experimental process, the Ni foil covered on the apparatus was replaced in turn at certain time intervals. Within the 25 hour experiment, six Ni foils were used and activities on these Ni foils were measured. The measurements showed that a great amount of ^{131}I and ^{133}I activities existed on the Ni foils. According to their activities on the Ni foil and collection time, the collection rates of ^{131}I and ^{133}I , expressed as Bq min^{-1} , are given in Fig. 4. Surprisingly, more or less, a large amount of iodine was released within the initial two hours and captured by the first piece of the Ni foil, with the collection rates of ^{131}I and ^{133}I being up to 4.4 Bq min^{-1} and 18.3 Bq min^{-1} , respectively. With the continuing experimental process, the collection rates of ^{131}I and ^{133}I decreased rapidly. After 8 hours, the collection rates decreased to 0.2 Bq min^{-1} and 0.8 Bq min^{-1} , about 4% of the peak values. Then, the collection rates of ^{131}I and ^{133}I were maintained at a stable level close to the background for the rest time of the experiment. Exemplified by ^{131}I , the activity of ^{131}I on the first two Ni foils accounted for more than 80% of the total collection activity on six Ni foils, and about 43% of ^{131}I activity lost in the molten salt, indicating that the release of iodine from the molten salt occurred mainly in the first 4 hours from the beginning of the melting process. It could be seen clearly that the release of fission product iodine exhibited typically transient characteristics. In other words, the release of a part of iodine occurred intensively within a short time at the beginning of melting. After the transient release, the remaining iodine was able to exist stably in the molten salt in the form of iodide, with

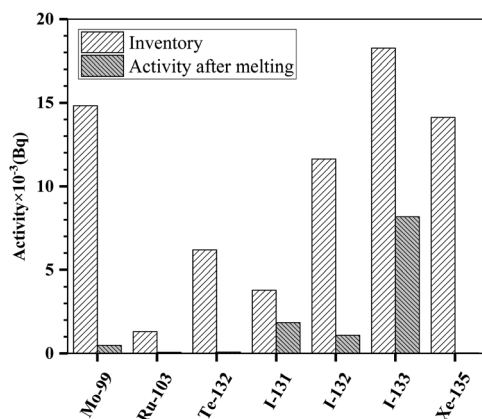


Fig. 3 Activities after melting and the inventories of ^{99}Mo , ^{103}Ru , ^{132}Te , ^{131}I , ^{132}I , ^{133}I and ^{135}Xe .

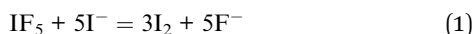


Fig. 4 Collection rates of ^{131}I and ^{133}I on the Ni foils.



no distinct decrease in its activity being observed during the continuous heating of the molten salt. The latter has been confirmed for times by ORNL.¹⁷

The transient release of a large amount of fission product iodine could be explained by hot atom chemistry.^{18,19} It is well known that, after capturing a neutron, the fissioning nucleus will split into two high-speed fission fragments. During the interaction of the primary fission fragments with the surrounding medium, various number of extranuclear electrons of these fission fragments would be striped sequentially, and form the highly exciting ions with various charges. When the fission fragments were retarded by the medium, the highly exciting iodine ions would, *via* relaxing, form the relatively stable iodine ions with a variety of chemical valence ranging from -1 to $+7$. The coexistence of fission product iodine with different chemical valence has been confirmed in aqueous solution systems.²⁰ Therefore, the chemical valence for the iodine carrier added in the aqueous solution must be adjusted to be identical to that of fission product iodine in the classical chemistry procedure of separation and yield determination. It was reasonable to believe that the fission product iodine generated in this work also had different chemical valences in the molten salt. When the mixture of irradiated UF_4 and FLiBe eutectic salt was heated, a chemical interaction between the iodine isotopes with different valences would occur to produce high-volatile elemental iodine as follows:



Then the released iodine was captured by the Ni foils and easily detected using a γ -ray spectrometer.

To demonstrate capture of the elemental iodine by a Ni foil, an additional experiment using natural iodine was carried out. CeF_4 , as an oxidation reagent, was added into the FLiBe molten salt containing 10 wt% of LiI, and then the mixture was heated at 650°C for 24 hours. After cooling, the collections on the Ni foil were characterized by XRD, and the XRD pattern is shown in Fig. 5. As can be seen, the existence of NiI_2 could be certainly verified, although the peaks located at 38 and 82 have not yet been matched. The demonstrating experiment indicated the

ability of the Ni foil to capture the released I_2 , which was formed by the reaction as follows:



Consequently, the additional experiment provided further support for the transient release of fission product iodine, as a form of I_2 .

This work clearly revealed the release of a large amount of iodine that resulted from the effect of the hot atom chemistry. The UF_4 used in this work has been irradiated as long as about 70 hours, and an abundant amount of fission product iodine with different chemical valences was generated. During the melting process, iodine accumulated in the long time was released in a short time from the salt, making it easy to be detected. A similar process should also occur in the MSRE. Differently from the observations of this study, however, both the generation of fission product iodine and the redox reactions between iodine species with different chemical valences would continuously occur to follow the nuclear fission. Because of no opportunity for the transient release of fission product iodine, the slow release certainly made its detection very difficult. This should be one of the main reasons responsible for the unknown fate of about one-fourth to one-third of fission product iodine observed in the MSRE operation by ORNL.⁷

The significant release of iodine from the fuel salt of the MSR might affect the neutron economics and the fraction of delayed-neutron precursor. Therefore, the behavior of iodine and its surveillance should be taken into account for the design and operation of MSRs. Furthermore, information about the release of fission product iodine from the reactor might have potential applications in the operation and control of the MSR. In this work, a Li metal reductant and a CeF_4 powder oxidant were added into the FLiBe molten salt containing irradiated UF_4 respectively. The effect of the variation in the redox potential of the molten salt on the iodine release from the salt was examined (see Table 2). Compared with 54.7% of release rate of iodine for the salt without addition of any reductant or oxidant, the transient release of iodine decreased to 23.1% when the Li metal was added into the salt. Conversely, the addition of the CeF_4 oxidant made the transient release increase to 87.2%. The reason was easily intelligible. The addition of the Li metal reductant consumed directly a part of fission product iodine with high valence, resulting in more iodine, in the form of I^- , remain in the salt. When the CeF_4 oxidant was added into the system, the I^- ions would be oxidized into elemental iodine I_2 followed by release from the salt. It could be inferred that the amount of iodine that remained in the salt, or that released from the salt as well, was correlated closely with the variation in the redox potential that the molten salt underwent.

According to the operation experience of the MSRE in ORNL, the fission process would gradually increase the redox potential of the fuel salt,^{7,21–23} leading to the corrosion of the structural material of the MSR. In order to mitigate the material corrosion, the beryllium metal was required to be added into the fuel salt periodically to control the redox potential. Recently, the study of

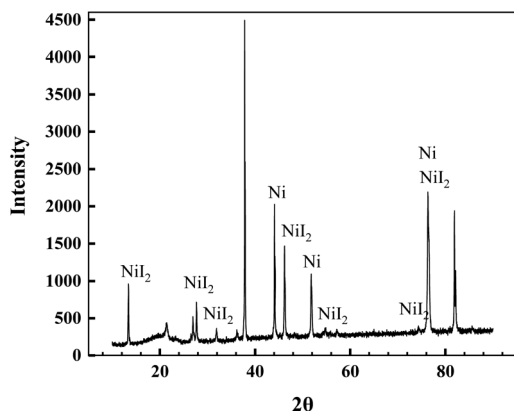


Fig. 5 XRD pattern of the deposits on the Ni foil.



Table 2 Release rate of ^{131}I and the amount of ^{131}I remaining in the salt under different redox conditions

Molten salt system	Additives (mg)	Release rate of ^{131}I	Ratio of activities		
			$^{131}\text{I}/^{140}\text{Ba}$	$^{131}\text{I}/^{143}\text{Ce}$	$^{131}\text{I}/^{95}\text{Zr}$
UF_4	—	—	0.83	0.41	3.56
$\text{UF}_4 + \text{FLiBe}$	—	54.7%	0.40	0.17	1.82
$\text{UF}_4 + \text{FLiBe}$	Li (20 mg)	23.1%	0.61	— ^a	2.43
$\text{UF}_4 + \text{FLiBe}$	CeF_4 (20 mg)	87.2%	0.106	0.035	0.444

^a No corresponding experimental data were obtained due to the short half-life of ^{143}Ce .

the redox potential and its control has received increasing attention.^{10,13} Zhang *et al.* pointed out the major phenomena that affect the redox potential and material corrosion, and main technologies for the redox potential measurement, including electro-chemical sensors and optical spectroscopy.²²

ORNL pointed out that the fission product ^{95}Nb in the fuel salt could be used as an in-line indicator of the redox condition of the fuel salt, because the transition in the chemical species of ^{95}Nb between soluble fluoride and the insoluble metal is dependent on the redox potential of the fuel salt.⁹ Our previous investigation indicated that, however, it was difficult to correctly distinguish ^{95}Nb existing as soluble fluoride from those in the form of insoluble metal particles,¹⁰ because the fuel salt was circulated at high velocity in the primary loop of MSR. Consequently, the experimental result given by ORNL was fragmentary, scattered, and even anomalous sometimes.⁹ Since the amount of iodine that remained in the salt was correlated closely with the variation in the redox potential, the activity of fission product iodine that remained in the fuel salt might be used as another indicator, to monitor the redox potential of the fuel salt in the MSR.

For a given reactor, the yields of fission products are known and fixed, and the ratio of activities for various fission products can be estimated correctly. Although ORNL classified the fission product iodine as a noble metal, in fact, after partly released in the form of elemental iodine from the molten salt, iodine that remained in the salt exhibited a similar behavior to the salt-seeking fission products and existed stably in the form of I^- in the salt. As a result, the redox condition of the molten salt in the MSR could be characterized by the activity ratios of ^{131}I to the salt-seeking fission products in the salt. As shown in Table 2, all of the activity ratios for ^{131}I to ^{140}Ba , ^{143}Ce and ^{95}Zr in salt increased with the addition of Li metal reductants when compared with the reference sample. On the contrary, these ratios decreased with the addition of the CeF_4 oxidant. The highly positive correlation between the activity ratios of ^{131}I to salt-seeking fission products in the salt illustrated the possible application of ^{131}I as an indicator for monitoring the redox potential of the molten salt.

Compared with the ^{95}Nb redox indicator proposed by ORNL, the application of ^{131}I as a redox indicator has several advantages. First, the fission product ^{131}I in the salt exists in the unique form of I^- , unlike the case of ^{95}Nb , where careful difference of soluble ^{95}Nb from insoluble ^{95}Nb is necessary but

very difficult. Second, the measurement of the activity ratios of ^{131}I to the salt-seeking fission products relies only on the relative measurement, which is more reliable and feasible than the absolute measurement involved in the determination of ^{95}Nb activity in the salt. Finally, the half-life of $^{131\text{m}}\text{Te}$, the precursor of ^{131}I , is about 25 min, much shorter than that of ^{95}Zr (65 days), the precursor of ^{95}Nb , thus the evaluations of the redox potential for the salt based on the measurement of activity ratios of ^{131}I to the salt-seeking fission products have less effect resulting from the radioactive growth-decay. In order to validate the application of ^{131}I in the surveillance of the redox potential of the salt in the MSR, it is certainly necessary to carry out further investigation.

Conclusions

The experiments indicated that a large amount of fission product iodine was released in a short time after melting the mixture of irradiated UF_4 and FLiBe eutectic salt. The behavior of transient release of iodine was attributed to the redox reaction between fission product iodine cumulated due to long-time irradiation. A similar process would occur continuously in the MSR, but with no condition of transient release. As a result, the detection of iodine released from the salt was not easy more or less, and this should be one of the main reasons responsible for unknown loss up to about one-fourth to one-third of fission product iodine observed by ORNL in the operation of MSRE.

Because of the close correlation between the release of iodine from the salt and the redox potential of the molten salt, it was pointed that the amount of ^{131}I remaining in the fuel salt may be used as another indicator for the redox potential of the MSR. The activity ratios of ^{131}I to the salt-seeking fission products such as ^{95}Zr , ^{140}Ba and ^{143}Ce would be preferable for the characterization of the redox potential of the MSR.

Conflicts of interest

There are no conflicts to declare.

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