Okuma Analysis and Research Center Fukushima Research and Engineering Institute Japan Atomic Energy Agency

Analysis Results of ALPS Treated Water (Sampling at February 29, 2024, 10:08 JST)

Analysis was performed on the ALPS treated water in K4-C tank group in measurement and confirmation facility sampled at <u>February 29, 2024, 10:08 JST</u>.

The analysis results of ³H and other nuclides are as follows:

• The confirmation of nuclides other than ³H which have been purified to below the regulatory limit by ALPS treatment:

The sum of ratios to the regulatory concentration limits of nuclides other than ³H (29 nuclides*¹) was 0.28 (less than 1), confirming that regulatory standard is satisfied.

The nuclides that confirmed not to be significantly present in ALPS treated water (39 nuclides*2): All targete radionuclides were confirmed not to be significantly present.

• The confirmation of ³H concentration in ALPS treated water:

³H concentration was <u>1.9E+05 Bq/L*3</u>.

The analyzed 68 radionuclides other than ³H are shown in Fig.1

²⁴¹Am

The nuclides which confirmed to be less than regulatory limit (29 nuclides)								
¹⁴ C	90 Y	¹³⁷ Cs	238 U	²⁴⁴ Cm				
⁵⁴ Mn	⁹⁹ Tc	¹⁴⁴ Ce	²³⁷ Np					
⁵⁵ Fe	¹⁰⁶ Ru	¹⁴⁷ Pm	²³⁸ Pu					
⁶⁰ Co	¹²⁵ Sb	¹⁵¹ Sm	²³⁹ Pu					
⁶³ Ni	^{125m} Te	¹⁵⁴ Eu	²⁴⁰ Pu					
⁷⁹ Se	¹²⁹ I	¹⁵⁵ Eu	²⁴¹ Pu					

234[]

⁹⁰Sr

134Cs

The nuclides which confirmed not to be significantly present (39 nuclides)							
⁵⁹ Fe	¹⁰³ Ru	¹²³ Sn	^{129m} Te	^{144m} Pr	^{242m} Am		
⁵⁸ Co	^{103m} Rh	¹²⁶ Sn	¹³⁵ Cs	¹⁴⁶ Pm	²⁴³ Am		
⁶⁵ Zn	¹⁰⁶ Rh	¹²⁴ Sb	¹³⁶ Cs	¹⁴⁸ Pm	²⁴² Cm		
⁸⁶ Rb	^{110m} Ag	^{123m} Te	^{137m} Ba	^{148m} Pm	²⁴³ Cm		
⁸⁹ Sr	^{113m} Cd	¹²⁷ Te	¹⁴⁰ Ba	¹⁵² Eu			
91 Y	^{115m} Cd	^{127m} Te	¹⁴¹ Ce	¹⁵³ Gd			
⁹⁵ Nb	^{119m} Sn	¹²⁹ Te	¹⁴⁴ Pr	¹⁶⁰ Tb			

Fig.1 Classification of radionuclides other than ³H

- *1: The nuclides that should be confirmed to satisfy the discharge standards (below the regulatory standards), as defined in the implementation plan.
- *2: From the view point of preventing the adverse impacts on reputation, the nuclides which independently confirmed by TEPCO HD not to significantly present in ALPS treated water.
- *3: Dilute more than 100 times so that the ³H concentration after seawater dilution is less than the maximum tritium concentration, 1,500 Bq/L/

1. The confirmation of nuclides other than ³H which have been purified to below the regulatory limit by ALPS treatment

Analysis results for the sum of ratios to the regulatory concentration limits of nuclides other than ³H (29 nuclides) that are confirmed to be less than regulatory limit (sum total of 1) are shown in Table 1. As the result of analysis, the sum of ratios to the regulatory concentration limits of nuclides other than ³H was 2.8E-01 (less than 1), confirming that regulatory standard is satisfied.

Table 1 Analysis results of nuclides other than ³H in ALPS treated water (The nuclides which confirmed to be less than regulatory limit)

(Sampled at February 29, 2024, 10:08 JST)

Nuclide	Concentrations [Bq/L]	Expanded Uncertainty*1 [Bq/L]	Detection Limit [Bq/L]	Ratios to Regulatory Concentration Limit [-]	Regulatory Concentratio n Limit*2 [Bq/L]	Measurement/ Evaluation method*4
¹⁴ C	2.0E+01	± 4.2E+00	4.7E-01	1.0E-02	2,000	Measurement
⁵⁴ Mn	ND	-	1.1E-02	1.1E-05	1,000	Measurement
⁵⁵ Fe	ND	-	6.0E-01	3.0E-04	2,000	Measurement
⁶⁰ Co	3.7E-01	± 4.9E-02	9.9E-03	1.9E-03	200	Measurement
⁶³ Ni	ND	-	1.2E+01	2.0E-03	6,000	Measurement
⁷⁹ Se	ND	-	2.0E+00	1.0E-02	200	Measurement
⁹⁰ Sr	3.5E-01	± 6.4E-02	7.1E-02	1.2E-02	30	Measurement
⁹⁰ Y	3.5E-01	-	7.1E-02	1.2E-03	300	Radiative equilibrium evaluation
⁹⁹ Tc	4.0E+00	± 4.2E-01	1.0E-01	4.0E-03	1,000	Measurement
¹⁰⁶ Ru	ND	-	1.0E-01	1.0E-03	100	Measurement
¹²⁵ Sb	5.0E-02	± 3.0E-02	4.4E-02	6.3E-05	800	Measurement
^{125m} Te	1.2E-02	-	1.1E-02	1.3E-05	900	Radiative equilibrium evaluation
¹²⁹ I	2.0E+00	± 3.1E-01	1.3E-01	2.2E-01	9	Measurement
¹³⁴ Cs	ND	-	4.2E-02	7.0E-04	60	Measurement
¹³⁷ Cs	3.8E-01	± 4.9E-02	1.3E-02	4.2E-03	90	Measurement
¹⁴⁴ Ce	ND	-	1.4E-01	7.0E-04	200	Measurement
¹⁴⁷ Pm	ND	-	1.8E-01	6.0E-05	3,000	Relative ratio
¹⁵¹ Sm	ND	-	8.6E-03	1.1E-06	8,000	Relative ratio evaluation

¹⁵⁴ Eu	ND	-	4.1E-02	1.0E-04	400	Measurement
¹⁵⁵ Eu	ND	-	8.2E-02	2.7E-05	3,000	Measurement
²³⁴ U					20	Measurement, Total
					20	α (U, Np group)
²³⁸ U	ND	_	1.5E-02	3.8E-03 ^{**3}	20	Measurement, Total
	TVD		1.52 02	3.02 03	20	α (U, Np group)
²³⁷ Np					9	Measurement, Total
МР					,	α (U, Np group)
						Measurement,
²³⁸ Pu					4	Total α (Pu, Am,
						Cm group)
					4	Measurement,
²³⁹ Pu						Total α (Pu, Am,
						Cm group))
						Measurement,
²⁴⁰ Pu	ND	ND -	8.7E-03	2.2E-03*3	4	Total α (Pu, Am,
						Cm group)
						Measurement,
²⁴¹ Am					5	Total α (Pu, Am,
						Cm group)
						Measurement,
²⁴⁴ Cm					7	Total α (Pu, Am,
						Cm group)
²⁴¹ Pu	ND		2.7E 01	1 4E 02	200	Relative ratio
² T'Pu	ND	-	2.7E-01	1.4E-03	200	evaluation
Sums o	f the Ratios to Reg	gulatory Concenti	ration Limits	2.8E-01		less than 1

- \bigcirc . \bigcirc E \pm \bigcirc means \bigcirc . \bigcirc × 10^{\pm} \bigcirc
- The results are shown in two significant digits.
- · Due to rounding of values, the sums may not exactly match with the actual value.
- ND (Not Detected) in the table indicates that the value is below the detection limit.
- The reference value for the decay correction is the date and time of sampling.
- %1: The uncertainty is the degree of variation in analytical value. Uncertainty is determined by combined all the variations of each step of the analytical procedure from sample collection to measurement. Here, the expanded uncertainty (U = 2 x u) is attached to the analyzed value by doubling the combined standard uncertainty (u).
- ※2: Legally required activity concentration limit established in the Ordinance for Operational Safety and Protection of Specified Nuclear Fuel Materials of the Nuclear Reactors at TEPCO's Fukushima Daiichi (Appendix of first and sixth column: The concentration limit in water outside the peripheral surveillance zone [In this table, Bq/cm³ was converted to Bq/L]).
- **3: The ratio to the regulatory concentration limits of α nuclides (U-234, U-238, Np-237, Pu-238, Pu-239, Pu-240,

Am-241, and Cm-244) was calculated by dividing the total alpha value by the lowest required activity concentration limit (4 Bq/L) of the selected alpha nuclides.

*4: Details of the measurement and evaluation methodology are described below.

Measurement: The radiation of samples is measured then the results are converted into the concentrations of each nuclide.

Total α (U, Np group): U and Np derived α rays of the samples were measured and the results are converted to total α concentration.

Total α (Pu, Am, Cm group): Pu, Am, Cm derived α rays of the samples were measured and the results are converted to total α concentration.

Radiative equilibrium evaluations: The state in which the ratio of the number of atoms of the parent nuclide to that of its progeny nuclides in a decay series is nearly constant is called radiative equilibrium. The concentration of each nuclide was evaluated based on this radiative equilibrium relationship and the measurement results of the parent (or progeny) nuclide.

Relative ratio evaluation: The ratio of each nuclide present in the reactor is evaluated by taking into account the generation, decay, impairment loss, etc. of the nuclides. The concentration of each nuclide is calculated by multiplying the measurement results of the reference nuclide by its existence ratio.

2. Analysis results of ³H in ALPS treated water

Analysis results for the ³H concentration in ALPS treated water are shown in Table 2. As the result of analysis, ³H concentration was confirmed to be <u>1.9E+05 Bq/L</u>.

	Activity	Expanded	Detection	Regulatory	Regulatory	Measurement/
Nuclide	Concentrations	Uncertainty*1	Limit	Concentration	Limit**2	Evaluation
	[Bq/L]	[Bq/L]	[Bq/L]	Ratio	[Bq/L]	method*4
³ H	1.9E+05	±2.0E+04	7.7E+01	3.2E+00	60,000	Measurement

Table 2 Analysis results of ³H in ALPS treated water (sampled at February 29, 2024 10:08 JST)

- ○.○E±○ means ○.○×10^{±○}
- The results are shown in two significant digits.
- The reference value for the decay correction is the date and time of sampling.
- \times 1: The uncertainty is the degree of variation in analytical value. Uncertainty is determined by combined all the variations of each step of the analytical procedure from sample collection to measurement. Here, the expanded uncertainty (U = 2 x u) is attached to the analyzed value by doubling the combined standard uncertainty (u).
- **2: Legally required activity concentration limit established in the Ordinance for Operational Safety and Protection of Specified Nuclear Fuel Materials of the Nuclear Reactors at TEPCO's Fukushima Daiichi (Appendix of first and sixth column: The concentration limit in water outside the peripheral surveillance zone [In this table, Bq/cm³ was converted to Bq/L]).

- ※3: Measurement: The radiation of samples is measured then the results are converted into the concentrations of each nuclide.
- 3. Nuclides other than ³H to be confirmed as not significantly present in ALPS treated water Analysis results for the target nuclides (39 nuclides) other than ³H that were confirmed to be as not significantly present in ALPS treated water are shown in Table 3. As the result of analysis, <u>all target radionuclides were confirmed not to be significantly present</u>. Furthermore, the sum of the ratios to the regulatory concentration limits for the 68 nuclides in Tables 1 and 3 is also less than 1.

Table 3 Analysis results of nuclides other than ³H in ALPS treated water (Nuclides which confirmed not to be significantly present) (sampling time: February 29, 2024 10:08 JST)

Nuclide	Activity Concentrations [Bq/L]	Regulatory Concentration Ratio [-]	Regulatory Limit*1 [Bq/L]	Evaluation**3	Measurement/ Evaluation method**4
⁵⁹ Fe	1.9E-02	4.8E-05	400	0	Measurement
⁵⁸ Co	1.1E-02	1.1E-05	1,000	0	Measurement
⁶⁵ Zn	1.9E-02	9.5E-05	200	0	Measurement
⁸⁶ Rb	1.6E-01	5.3E-04	300	0	Measurement
⁸⁹ Sr	7.4E-02	2.5E-04	300	0	Measurement
⁹¹ Y	4.2E+00	1.4E-02	300	0	Measurement
⁹⁵ Nb	1.6E-02	1.6E-05	1,000	0	Measurement
¹⁰³ Ru	1.6E-02	1.6E-05	1,000	0	Measurement
^{103m} Rh	1.6E-02	8.0E-08	200,000	0	Radiative equilibrium evaluation
¹⁰⁶ Rh	1.0E-01	3.3E-07	300,000	0	Radiative equilibrium evaluation
^{110m} Ag	1.2E-02	4.0E-05	300	0	Measurement
^{113m} Cd	1.4E-01	3.5E-03	40	0	Measurement
^{115m} Cd	5.5E-01	1.8E-03	300	0	Measurement
^{119m} Sn	3.0E-02	1.5E-05	2,000	0	Relative ratio evaluation
¹²³ Sn	1.7E+00	4.3E-03	400	0	Measurement
¹²⁶ Sn	3.9E-01	2.0E-03	200	0	Measurement
¹²⁴ Sb	2.0E-02	6.7E-05	300	0	Measurement
^{123m} Te	2.0E-02	3.3E-05	600	0	Measurement
¹²⁷ Te	1.4E+00	2.8E-04	5,000	0	Measurement
^{127m} Te	1.4E+00	4.7E-03	300	0	Relative ratio evaluation
¹²⁹ Te	1.9E-01	1.9E-05	10,000	0	Measurement
^{129m} Te	3.7E-01	1.2E-03	300	0	Measurement

			1		1
¹³⁵ Cs	8.3E-08	4.2E-09	600	0	Relative ratio evaluation
¹³⁶ Cs	1.4E-02	4.7E-05	300	0	Measurement
^{137m} Ba	1.2E-02	4.5E-07	800,000	0	Radiative equilibrium evaluation
¹⁴⁰ Ba	6.6E-02	2.2E-04	300	0	Measurement
¹⁴¹ Ce	6.5E-02	6.5E-05	1,000	0	Measurement
¹⁴⁴ Pr	1.4E-01	7.0E-06	20,000	0	Radiative equilibrium evaluation
^{144m} Pr	1.3E-03	3.3E-08	40,000	0	Radiative equilibrium evaluations
¹⁴⁶ Pm	1.9E-02	2.1E-05	900	0	Measurement
¹⁴⁸ Pm	9.6E-02	3.2E-04	300	0	Measurement
^{148m} Pm	1.3E-02	2.6E-05	500	0	Measurement
¹⁵² Eu	5.8E-02	9.7E-05	600	0	Measurement
¹⁵³ Gd	4.6E-02	1.5E-05	3,000	0	Measurement
¹⁶⁰ Tb	3.6E-02	7.2E-05	500	0	Measurement
^{242m} Am	4.9E-05	9.8E-06	5	0	Relative ratio evaluation
²⁴³ Am			5	0	Measurement, Total α (Pu, Am, Cm group)
²⁴² Cm	8.7E-03	2.2E-03**2	60	0	Measurement, Total α (Pu, Am, Cm group)
²⁴³ Cm			6	0	Measurement, Total α (Pu, Am, Cm group)

- $\bigcirc.\bigcirc E\pm\bigcirc$ means $\bigcirc.\bigcirc\times 10^{\pm\bigcirc}$
- The results are shown in two significant digits.
- Due to rounding of values, the sums of shown results in the table may not match.
- The reference value for the decay correction is the date and time of sampling.
- ** 1: Legally required activity concentration limit established in the Ordinance for Operational Safety and Protection of Specified Nuclear Fuel Materials of the Nuclear Reactors at TEPCO's Fukushima Daiichi (Appendix of first and sixth column: The concentration limit in water outside the peripheral surveillance zone [In this table, Bq/cm³ was converted to Bq/L]).
- **2: The ratio to the regulatory concentration limits of α nuclides (Am-243, Cm-242, and Cm-243) was calculated by dividing the total alpha value by the lowest required activity concentration limit (4 Bq/L) of the selected alpha nuclides.
- *3: If it is not significantly present, it is marked with a "O"; if it is significantly present, it is marked with an "x".

 The evaluation is made as not significantly present ("O") if any of the following are satisfied:
 - The measured nuclide concentration is below the detection limit.
 - Nuclides evaluated by radiative equilibrium, etc.: When the evaluated nuclide is detected, the concentration is extremely low compared to the regulatory concentration limit, i.e., the evaluated value is less than the set

detection limit (less than 1/100 of the regulatory concentration limit), and can be judged to be equivalent to less than the detection limit.

Evaluated Nuclide	Evaluated Value [Bq/L]	Regulatory Concentration Ratio	Regulatory Limit ^{*1} [Bq/L]
^{103m} Rh	ND	-	200,000
¹⁰⁶ Rh	ND	-	300,000
^{119m} Sn	ND	-	2,000
^{127m} Te	ND	-	300
¹³⁵ Cs	2.5E-06	4.2E-09	600
^{137m} Ba	3.6E-01	4.5E-07	800,000
¹⁴⁴ Pr	ND	-	20,000
^{144m} Pr	ND	-	40,000
^{242m} Am	ND	-	5

- ND (Not Detected) in the table indicates that the value is below the detection limit.
- *4: Details of the measurement and evaluation methodology are described below.

Measurement: The radiation of samples is measured then the results are converted into the concentrations of each nuclide.

Total α (U, Np group): U and Np derived α rays of the samples were measured and the results are converted to total α concentration.

Total α (Pu, Am, Cm group): Pu, Am, Cm derived α rays of the samples were measured and the results are converted to total α concentration.

Radiative equilibrium evaluations: The state in which the ratio of the number of atoms of the parent nuclide to that of its progeny nuclides in a decay series is nearly constant is called radiative equilibrium. The concentration of each nuclide was evaluated based on this radiative equilibrium relationship and the measurement results of the parent (or progeny) nuclide.

Relative ratio evaluation: The ratio of each nuclide present in the reactor is evaluated by taking into account the generation, decay, impairment loss, etc. of the nuclides. The concentration of each nuclide is calculated by multiplying the measurement results of the reference nuclide by its existence ratio.

Reference: Measurement and evaluation methods for each nuclide in the third-party analysis of ALPS treated water

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No.	Nuclide		Measurement and evaluation methods for each nuclide
			in the third-party analysis of ALPS treated water
1	³ H	β-ray	Purify tritiated water by distillation and mixing sample and scintillator
_		F,	Liquid scintillation counter
2	¹⁴ C	β-ray	Isolation by collection on adsorbent, mixing sample and scintillator
		p 14.5	Liquid scintillation counter
3	⁵⁴ Mn	γ-ray	A single sample was prepared in a Marinelli beaker.
	14111	7 Tay	Gamma-ray spectrometry using high purity germanium (HPGe) detector
4	⁵⁵ Fe	X-ray	Isolation by resin and sedimentation
	10	71 Tay	Low-energy photon detector (Ge-LEPS)
5	⁵⁹ Fe	W- r 937	A single sample was prepared in a Marinelli beaker.
<i>J</i>	T'C	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
6	⁵⁸ Co	\/_ * 937	A single sample was prepared in a Marinelli beaker.
U		γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
7	⁶⁰ Co	W- r 937	A single sample was prepared in a Marinelli beaker.
,	Co	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
8	⁶³ Ni	β-ray	Isolation by resin, mixing sample and scintillator
0	INI	p-ray	Liquid scintillation counter
9	⁶⁵ Zn	0/ #03/	A single sample was prepared in a Marinelli beaker.
9	ZII	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
10	⁷⁹ Se	ICP-MS	Isolation by resin
10	Se	101-1015	Inductively coupled plasma mass spectrometer
11	⁸⁶ Rb	0/ #03/	A single sample was prepared in a Marinelli beaker.
11	Ku	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
12	⁸⁹ Sr	0	Isolation by resin and sedimentation
12	o ⁵ Sr	β-ray	Beta-ray spectrometer (plastic scintillator)
13	⁹⁰ Sr	0	Isolation by resin and sedimentation
13	Si	β-ray	Beta-ray spectrometer (plastic scintillator)
14	⁹⁰ Y	Evaluation	Radiative equilibrium evaluation
15	⁹¹ Y	A. #0**	A single sample was prepared in a Marinelli beaker.
13	У- У	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
16	95 N TL	A. #0**	A single sample was prepared in a Marinelli beaker.
10	IND	⁹⁵ Nb γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
17	⁹⁹ Tc	ICD MC	Isolation by resin
17	1C	ICP-MS	Inductively coupled plasma mass spectrometer
10	103 D		A single sample was prepared in a Marinelli beaker.
18	¹⁰³ Ru	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
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19	¹⁰⁶ Ru	γ-ray	A single sample was prepared in a Marinelli beaker.
		7 1.03	Gamma-ray spectrometry using high purity germanium (HPGe) detector
20	^{103m} Rh	Evaluation	Radiative equilibrium evaluation
21	¹⁰⁶ Rh	Evaluation	Radiative equilibrium evaluation
22	^{110m} Ag	M rox	A single sample was prepared in a Marinelli beaker.
22	Ag	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
23	^{113m} Cd	0	Isolation by resin, mixing sample and scintillator
23	Ca	β-ray	Liquid scintillation counter
2.4	^{115m} Cd		A single sample was prepared in a Marinelli beaker.
24	· · · · · · · · · · · · · · · · · · ·	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
25	^{119m} Sn	Evaluation	Relative ratio evaluation:
26	122.0		A single sample was prepared in a Marinelli beaker.
26	¹²³ Sn	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
25	1260		A single sample was prepared in a Marinelli beaker.
27	¹²⁶ Sn	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
20	12461		A single sample was prepared in a Marinelli beaker.
28	¹²⁴ Sb	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
• •	125 ~ 1		A single sample was prepared in a Marinelli beaker.
29	¹²⁵ Sb	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
2.0	122mm		A single sample was prepared in a Marinelli beaker.
30	^{123m} Te	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
31	^{125m} Te	Evaluation	Radiative equilibrium evaluation
	1277		A single sample was prepared in a Marinelli beaker.
32	¹²⁷ Te	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
33	^{127m} Te	Evaluation	Radiative equilibrium evaluation
2.4	129m		A single sample was prepared in a Marinelli beaker.
34	¹²⁹ Te	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
2.5	129m/F		A single sample was prepared in a Marinelli beaker.
35	^{129m} Te	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
26	120*	37	A single sample was prepared in a measurement cell.
36	^{129}I	X-ray	Low-energy photon detector (Ge-LEPS)
	124 ~		A single sample was prepared in a Marinelli beaker.
37	¹³⁴ Cs	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
38	¹³⁵ Cs	Evaluation	Relative ratio evaluation:
20	126 ~		A single sample was prepared in a Marinelli beaker.
39	¹³⁶ Cs	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
4.0	127~		A single sample was prepared in a Marinelli beaker.
40	¹³⁷ Cs	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
	i		

41	^{137m} Ba	Evaluation	Radiative equilibrium evaluation
42	¹⁴⁰ Ba		A single sample was prepared in a Marinelli beaker.
42	ъъва	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
42	¹⁴¹ Ce		A single sample was prepared in a Marinelli beaker.
43	Ce	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
4.4	¹⁴⁴ Ce		A single sample was prepared in a Marinelli beaker.
44	···Ce	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
45	¹⁴⁴ Pr	Evaluation	Radiative equilibrium evaluation
46	^{144m} Pr	Evaluation	Radiative equilibrium evaluation
47	¹⁴⁶ Pm		A single sample was prepared in a Marinelli beaker.
47	Pm	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
48	¹⁴⁷ Pm	Evaluation	Relative ratio evaluation:
40	1480		A single sample was prepared in a Marinelli beaker.
49	¹⁴⁸ Pm	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
5 0	148m D		A single sample was prepared in a Marinelli beaker.
50	^{148m} Pm	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
51	¹⁵¹ Sm	Evaluation	Relative ratio evaluation:
50	15215	γ-ray	A single sample was prepared in a Marinelli beaker.
52	¹⁵² Eu		Gamma-ray spectrometry using high purity germanium (HPGe) detector
53	¹⁵⁴ Eu		A single sample was prepared in a Marinelli beaker.
53	Eu	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
<i>5</i> 1	¹⁵⁵ Eu		A single sample was prepared in a Marinelli beaker.
54	Eu	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
55	¹⁵³ Gd		A single sample was prepared in a Marinelli beaker.
33	- Ga	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
5.6	¹⁶⁰ Tb		A single sample was prepared in a Marinelli beaker.
56	10	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
57	²³⁴ U	or mov.	Separated by resin and evaporated and solidified on a stainless plate
37		α-ray	α-ray scintillation counter (ZnS scintillator)
F 0	238 [J		Separated by resin and evaporated and solidified on a stainless plate
58	2300	α-ray	α-ray scintillation counter (ZnS scintillator)
50	237ът		Separated by resin and evaporated and solidified on a stainless plate
59	²³⁷ Np	α-ray	α-ray scintillation counter (ZnS scintillator)
60	²³⁸ Pu	0.6	Separated by resin, collected as sedimentation, and evaporated to dryness
60	-5°Pu	α-ray	α-ray scintillation counter (ZnS scintillator)
<i>(</i> 1	239 D		Separated by resin, collected as sedimentation, and evaporated to dryness
61	²³⁹ Pu	α-ray	α-ray scintillation counter (ZnS scintillator)
	1	1	

(2)	²⁴⁰ Pu		Separated by resin, collected as sedimentation, and evaporated to dryness
62	2.ºPu	α-ray	α-ray scintillation counter (ZnS scintillator)
63	²⁴¹ Pu	Evaluation	Relative ratio evaluation:
64	²⁴¹ Am		Separated by resin, collected as sedimentation, and evaporated to dryness
64	- Am	α-ray	α-ray scintillation counter (ZnS scintillator)
65	^{242m} Am	Evaluation	Relative ratio evaluation:
66	242 .		Separated by resin, collected as sedimentation, and evaporated to dryness
00	²⁴³ Am	α-ray	α-ray scintillation counter (ZnS scintillator)
67	²⁴² Cm		Separated by resin, collected as sedimentation, and evaporated to dryness
07	Cm	α-ray	α-ray scintillation counter (ZnS scintillator)
(0)	243.0		Separated by resin, collected as sedimentation, and evaporated to dryness
68	²⁴³ Cm	α-ray	α-ray scintillation counter (ZnS scintillator)
(0)	244.		Separated by resin, collected as sedimentation, and evaporated to dryness
09	69 ²⁴⁴ Cm	α-ray	α-ray scintillation counter (ZnS scintillator)