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

Analysis Results of ALPS Treated Water (Sampling at May 16, 2025, 9:46 JST)

Analysis was performed on the ALPS treated water in K4-C tank group in measurement and confirmation facility sampled at May 16, 2025, 9:46 JST.

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 (30 nuclides*¹) was 0.096(less than 1), confirming that regulatory standard is satisfied.

The nuclides that confirmed not to be significantly present in ALPS treated water (38 nuclides*2): All target radionuclides were confirmed not to be significantly present.

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

³H concentration was <u>2.5E+05 Bq/L*3</u>.

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

The nuclides which confirmed to be less than regulatory limit (30 nuclides)
charring and confidence of

14 C	90 Y	¹³⁴ Cs	²³⁴ U	²⁴¹ Am
⁵⁴ Mn	⁹⁹ Tc	¹³⁷ Cs	238U	²⁴⁴ Cm
⁵⁵ Fe	¹⁰⁶ Ru	¹⁴⁴ Ce	²³⁷ Np	
⁶⁰ Co	^{113m} Cd	¹⁴⁷ Pm	²³⁸ Pu	
⁶³ Ni	¹²⁵ Sb	¹⁵¹ Sm	²³⁹ Pu	
⁷⁹ Se	^{125m} Te	¹⁵⁴ Eu	²⁴⁰ Pu	
⁹⁰ Sr	¹²⁹ I	¹⁵⁵ Eu	²⁴¹ Pu	

The nuclides which confirmed not to be significantly present (38 nuclides)

5	⁵⁹ Fe	¹⁰³ Ru	¹²⁶ Sn	¹³⁵ Cs	¹⁴⁶ Pm	²⁴³ Am
5	⁵⁸ Co	^{103m} Rh	¹²⁴ Sb	¹³⁶ Cs	¹⁴⁸ Pm	²⁴² Cm
6	⁵⁵ Zn	¹⁰⁶ Rh	^{123m} Te	^{137m} Ba	^{148m} Pm	²⁴³ Cm
8	³⁶ Rb	^{110m} Ag	¹²⁷ Te	¹⁴⁰ Ba	¹⁵² Eu	
8	⁸⁹ Sr	^{115m} Cd	^{127m} Te	¹⁴¹ Ce	¹⁵³ Gd	
	⁹¹ Y	^{119m} Sn	¹²⁹ Te	¹⁴⁴ Pr	¹⁶⁰ Tb	
9	⁹⁵ Nb	¹²³ Sn	^{129m} Te	^{144m} Pr	^{242m} Am	

Fig.1 Classification of radionuclides other than ³H

- *1: The nuclides that are selected by TEPCO HD based on the flow defined in the implementation plan and should be confirmed to satisfy the discharge standard (below regulatory standards).
- *2: From the viewpoint 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 (30 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 <u>9.6E-02</u> (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)

Nuclide	Concentrations	Expanded Uncertainty*1	Detection Limit	Ratios to Regulatory Concentration Limit	Regulatory Concentration Limit*2	Measurement/ Evaluation method*3
	[Bq/L]	[Bq/L]	[Bq/L]	[-]	[Bq/L]	
¹⁴ C	3.3E+01	± 6.6E+00	3.5E-01	1.7E-02	2,000	Measurement
⁵⁴ Mn	ND	-	1.1E-02	1.1E-05	1,000	Measurement
⁵⁵ Fe	ND	-	7.7E-01	3.9E-04	2,000	Measurement
⁶⁰ Co	5.7E-01	± 9.2E-02	9.5E-03	2.9E-03	200	Measurement
⁶³ Ni	ND	-	1.1E+01	1.8E-03	6,000	Measurement
⁷⁹ Se	ND	-	2.0E+00	1.0E-02	200	Measurement
⁹⁰ Sr	3.1E-01	± 5.7E-02	6.2E-02	1.0E-02	30	Measurement
90Y	3.1E-01	-	6.2E-02	1.0E-03	300	Radiative equilibrium evaluation
⁹⁹ Tc	1.2E+00	± 2.9E-01	1.0E-01	1.2E-03	1,000	Measurement
¹⁰⁶ Ru	ND	-	1.1E-01	1.1E-03	100	Measurement
^{113m} Cd	ND	-	1.3E-01	3.3E-03	40	Measurement
¹²⁵ Sb	2.1E-01	± 4.5E-02	4.4E-02	2.6E-04	800	Measurement
^{125m} Te	5.1E-02	-	1.1E-02	5.7E-05	900	Radiative equilibrium evaluation
¹²⁹ I	3.4E-01	± 3.4E-02	7.0E-03	3.8E-02	9	Measurement
¹³⁴ Cs	ND	-	4.1E-02	6.8E-04	60	Measurement
¹³⁷ Cs	1.3E-01	± 2.2E-02	1.2E-02	1.4E-03	90	Measurement
¹⁴⁴ Ce	ND	-	1.4E-01	7.0E-04	200	Measurement
¹⁴⁷ Pm	ND	-	1.1E-01	3.7E-05	3,000	Relative ratio evaluation
¹⁵¹ Sm	ND	-	8.8E-03	1.1E-06	8,000	Relative ratio evaluation

¹⁵⁴ Eu	ND	-	3.6E-02	9.0E-05	400	Measurement					
¹⁵⁵ Eu	ND	-	7.8E-02	2.6E-05	3,000	Measurement					
²³⁴ U					20	Measurement, Total α (U, Np group)					
²³⁸ U	ND	-	1.2E-02	3.0E-03*4	20	Measurement, Total α (U, Np group)					
²³⁷ Np					9	Measurement, Total α (U, Np group)					
²³⁸ Pu					4	Measurement, Total α (Pu, Am, Cm group)					
²³⁹ Pu	ND -			4		4	Measurement, Total α (Pu, Am, Cm group))				
²⁴⁰ Pu		ND	ND	ND	ND	-	8.2E-03	2.1E-03 ^{**4}	2.1E-03 ^{**4} 4	4	Measurement, Total α (Pu, Am, Cm group)
²⁴¹ Am					5	Measurement, Total α (Pu, Am, Cm group)					
²⁴⁴ Cm					7	Measurement, Total α (Pu, Am, Cm group)					
²⁴¹ Pu	ND	-	2.3E-01	1.2E-03	200	Relative ratio					
Sun	ns of the Ratios to Reg	gulatory Concentrati	ion Limits	9.6E-02		less than 1					

- $\bigcirc.\bigcirc E\pm\bigcirc$ means $\bigcirc.\bigcirc\times 10^{\pm\bigcirc}$
- The results are shown in two significant digits.
- Due to rounding of values, the sums 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: 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.

**4: 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.

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>2.5E+05 Bq/L</u>.

Table 2 Analysis results of ³H in ALPS treated water

	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**3
³ H	2.5E+05	±2.2E+04	9.4E+01	4.2E+00	60,000	Measurement

- $\bigcirc.\bigcirc$ E± \bigcirc means $\bigcirc.\bigcirc\times10^{\pm\bigcirc}$
- 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 (38 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)

Nuclide	Activity Concentrations [Bq/L]	Regulatory Concentration Ratio [-]	Regulatory Limit*1 [Bq/L]	Evaluation*2	Measurement/ Evaluation method**3
⁵⁹ Fe	2.0E-02	5.0E-05	400	0	Measurement
⁵⁸ Co	1.1E-02	1.1E-05	1,000	0	Measurement
⁶⁵ Zn	2.1E-02	1.1E-04	200	0	Measurement
⁸⁶ Rb	1.4E-01	4.7E-04	300	0	Measurement
⁸⁹ Sr	6.6E-02	2.2E-04	300	0	Measurement
⁹¹ Y	5.2E+00	1.7E-02	300	0	Measurement
⁹⁵ Nb	1.5E-02	1.5E-05	1,000	0	Measurement
¹⁰³ Ru	2.1E-02	2.1E-05	1,000	0	Measurement
^{103m} Rh	2.1E-02	1.1E-07	200,000	0	Radiative equilibrium evaluation
¹⁰⁶ Rh	1.1E-01	3.7E-07	300,000	0	Radiative equilibrium evaluation
^{110m} Ag	1.2E-02	4.0E-05	300	0	Measurement
^{115m} Cd	6.1E-01	2.0E-03	300	0	Measurement
^{119m} Sn	5.0E-03	2.5E-06	2,000	0	Relative ratio evaluation
¹²³ Sn	1.8E+00	4.5E-03	400	0	Measurement
¹²⁶ Sn	3.6E-01	1.8E-03	200	0	Measurement
¹²⁴ Sb	2.2E-02	7.3E-05	300	0	Measurement
^{123m} Te	2.1E-02	3.5E-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	1.8E-01	1.8E-05	10,000	0	Measurement
^{129m} Te	3.7E-01	1.2E-03	300	0	Measurement
¹³⁵ Cs	8.3E-08	1.5E-09	600	0	Relative ratio evaluation
¹³⁶ Cs	1.3E-02	4.3E-05	300	0	Measurement
^{137m} Ba	1.1E-02	1.5E-07	800,000	0	Radiative equilibrium evaluation

	1	I	I	1	1
¹⁴⁰ Ba	6.8E-02	2.3E-04	300	0	Measurement
¹⁴¹ Ce	5.9E-02	5.9E-05	1,000	0	Measurement
¹⁴⁴ Pr	1.4E-01	7.0E-06	20,000	0	Radiative equilibrium
11	1.4L-01	7.0L-00	20,000	O	evaluation
144m p r	1.25.02	2.25.09	40.000		Radiative equilibrium
·····PT	1.3E-03	3.3E-08	40,000	0	evaluations
¹⁴⁶ Pm	2.1E-02	2.3E-05	900	0	Measurement
¹⁴⁸ Pm	2.2E-01	7.3E-04	300	0	Measurement
^{148m} Pm	1.4E-02	2.8E-05	500	0	Measurement
¹⁵² Eu	5.7E-02	9.5E-05	600	0	Measurement
¹⁵³ Gd	4.6E-02	1.5E-05	3,000	0	Measurement
¹⁶⁰ Tb	3.4E-02	6.8E-05	500	0	Measurement
^{242m} Am	3.8E-05	7.6E-06	5	0	Relative ratio evaluation
²⁴³ Am			5	0	Measurement, Total α
Alli			3	U	(Pu, Am, Cm group)
²⁴² Cm	8.2E-03	2.1E-03 ^{*4}	60	0	Measurement, Total α
CIII	6.2E-03	2.1E-03	00	O	(Pu, Am, Cm group)
²⁴³ Cm			6		Measurement, Total α
CIII				0	(Pu, Am, Cm group)

- $\bigcirc.\bigcirc$ E± \bigcirc means $\bigcirc.\bigcirc$ × $10^{\pm\bigcirc}$
- 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: If it is not significantly present, it is marked with a "○"; if it is significantly present, it is marked with an "×".
 The evaluation is made as not significantly present ("○") 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	8.7E-07	1.5E-09	600
^{137m} Ba	1.2E-01	1.5E-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.
- *3: 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.

**4: 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.

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

	1		the third-party analysis of AETS treated water
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
		p iuj	Liquid scintillation counter
2	¹⁴ C	β-ray	Isolation by collection on adsorbent, mixing sample and scintillator
		p-ray	Liquid scintillation counter
3	⁵⁴ Mn	W-r93/	A single sample was prepared in a Marinelli beaker.
3	10111	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
4	⁵⁵ Fe	X-ray	Isolation by resin and sedimentation
4	T'C	A-1ay	Low-energy photon detector (Ge-LEPS)
5	⁵⁹ Fe		A single sample was prepared in a Marinelli beaker.
3	re	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
6	⁵⁸ Co		A single sample was prepared in a Marinelli beaker.
6	1300	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
7	⁶⁰ Co		A single sample was prepared in a Marinelli beaker.
7	**C0	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
0	⁶³ Ni	0	Isolation by resin, mixing sample and scintillator
8	⁶⁵ N1	β-ray	Liquid scintillation counter
0	⁶⁵ Zn		A single sample was prepared in a Marinelli beaker.
9	Zn	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
10	⁷⁹ Se	ICP-MS	Isolation by resin
10	Se	ICF-IVIS	Inductively coupled plasma mass spectrometer
11	⁸⁶ Rb		A single sample was prepared in a Marinelli beaker.
11	**Kb	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
12	⁸⁹ Sr	0	Isolation by resin and sedimentation
12	Sr	β-ray	Beta-ray spectrometer (plastic scintillator)
12	⁹⁰ Sr	0	Isolation by resin and sedimentation
13	Sr	β-ray	Beta-ray spectrometer (plastic scintillator)
14	⁹⁰ Y	Evaluation	Radiative equilibrium evaluation
1.5	⁹¹ Y		A single sample was prepared in a Marinelli beaker.
15	, Y	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
1.6	⁹⁵ Nb	a. ====	A single sample was prepared in a Marinelli beaker.
16	IND	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.
			Gamma-ray spectrometry using high purity germanium (HPGe) detector
20	^{103m} Rh	Evaluation	Radiative equilibrium evaluation
21	¹⁰⁶ Rh	Evaluation	Radiative equilibrium evaluation
22	$^{110\mathrm{m}}\mathrm{Ag}$	γ-ray	A single sample was prepared in a Marinelli beaker.
	715	1 Iuy	Gamma-ray spectrometry using high purity germanium (HPGe) detector
23	^{113m} Cd	β-ray	Isolation by resin, mixing sample and scintillator
23	Cu	p-ray	Liquid scintillation counter
24	^{115m} Cd	A #01/	A single sample was prepared in a Marinelli beaker.
24	Cu	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
25	^{119m} Sn	Evaluation	Relative ratio evaluation:
26	¹²³ Sn		A single sample was prepared in a Marinelli beaker.
26	Sil	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
27	¹²⁶ Sn		A single sample was prepared in a Marinelli beaker.
27	120Sn	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
20	¹²⁴ Sb		A single sample was prepared in a Marinelli beaker.
28	12186	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
20	¹²⁵ Sb		A single sample was prepared in a Marinelli beaker.
29	12386	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
20	^{123m} Te		A single sample was prepared in a Marinelli beaker.
30	1231e	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
31	^{125m} Te	Evaluation	Radiative equilibrium evaluation
22	¹²⁷ Te		A single sample was prepared in a Marinelli beaker.
32	12/16	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
33	^{127m} Te	Evaluation	Radiative equilibrium evaluation
2.4	¹²⁹ Te		A single sample was prepared in a Marinelli beaker.
34	123 Te	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
2.5	^{129m} Te		A single sample was prepared in a Marinelli beaker.
35	123m1e	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
26	1291	ICD MG	Isolation by resin
36	¹²⁹ I	ICP-MS	Inductively coupled plasma mass spectrometer
27	¹³⁴ Cs		A single sample was prepared in a Marinelli beaker.
37	is iCs	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
38	¹³⁵ Cs	Evaluation	Relative ratio evaluation:
20	1360		A single sample was prepared in a Marinelli beaker.
39	¹³⁶ Cs	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector
	127 ~		A single sample was prepared in a Marinelli beaker.
40	¹³⁷ Cs	γ-ray	Gamma-ray spectrometry using high purity germanium (HPGe) detector

41	^{137m} Ba	Evaluation	Radiative equilibrium evaluation
42	¹⁴⁰ Ba	γ-ray	A single sample was prepared in a Marinelli beaker.
			Gamma-ray spectrometry using high purity germanium (HPGe) detector
43	¹⁴¹ Ce	γ-ray	A single sample was prepared in a Marinelli beaker.
			Gamma-ray spectrometry using high purity germanium (HPGe) detector
44	¹⁴⁴ Ce	γ-ray	A single sample was prepared in a Marinelli beaker.
44			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	γ-ray	A single sample was prepared in a Marinelli beaker.
47			Gamma-ray spectrometry using high purity germanium (HPGe) detector
48	¹⁴⁷ Pm	Evaluation	Relative ratio evaluation:
40	¹⁴⁸ Pm	γ-ray	A single sample was prepared in a Marinelli beaker.
49			Gamma-ray spectrometry using high purity germanium (HPGe) detector
50	^{148m} Pm	γ-ray	A single sample was prepared in a Marinelli beaker.
50	· ······Pm		Gamma-ray spectrometry using high purity germanium (HPGe) detector
51	¹⁵¹ Sm	Evaluation	Relative ratio evaluation:
52	¹⁵² Eu	γ-ray	A single sample was prepared in a Marinelli beaker.
52			Gamma-ray spectrometry using high purity germanium (HPGe) detector
53	¹⁵⁴ Eu	γ-ray	A single sample was prepared in a Marinelli beaker.
33			Gamma-ray spectrometry using high purity germanium (HPGe) detector
54	¹⁵⁵ Eu	γ-ray	A single sample was prepared in a Marinelli beaker.
34			Gamma-ray spectrometry using high purity germanium (HPGe) detector
55	¹⁵³ Gd	γ-ray	A single sample was prepared in a Marinelli beaker.
33			Gamma-ray spectrometry using high purity germanium (HPGe) detector
56	¹⁶⁰ Tb	γ-ray	A single sample was prepared in a Marinelli beaker.
30			Gamma-ray spectrometry using high purity germanium (HPGe) detector
57	²³⁴ U	α-ray	Separated by resin and evaporated and solidified on a stainless plate
57			α-ray scintillation counter (ZnS scintillator)
50	²³⁸ U	α-ray	Separated by resin and evaporated and solidified on a stainless plate
58			α-ray scintillation counter (ZnS scintillator)
50	²³⁷ Np	α-ray	Separated by resin and evaporated and solidified on a stainless plate
59			α-ray scintillation counter (ZnS scintillator)
60	²³⁸ Pu	α-ray	Separated by resin, collected as sedimentation, and evaporated to dryness
			α-ray scintillation counter (ZnS scintillator)
61	²³⁹ Pu	α-ray	Separated by resin, collected as sedimentation, and evaporated to dryness
			α-ray scintillation counter (ZnS scintillator)
		1	

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