Determination of the gamma emission probabilities of $^{239}$Np

Shang Jian-bo$^a$, Du Li-li, Bai Tao, Dai Yi-hua, Zhong Zhen-yuan, Liu Jie, and Shi Quan-lin

Northwest Institute of Nuclear Technology, Xi’an, China

Abstract. $^{239}$Np is an important nuclide as the decay daughter of $^{239}$U and it decays to $^{239}$Pu by emitting beta particles and gamma rays with a half life of 2.356 days. The data of the emission probabilities of its gamma-rays in the open references are consistent except for the main gamma-ray of 106.1 keV, the emission probability of which varies from 25.9% to 27.2%. To verify the emission probability of 106.1 keV gamma-ray of $^{239}$Np, a N-type coaxial HPGe detector was calibrated using $^{241}$Am, $^{133}$Ba, $^{60}$Co, $^{152}$Eu and $^{155}$Eu reference gamma sources to get the accurate efficiency of the 106.1 keV gamma-ray. $^{239}$Np was purified from solution containing $^{243}$Am, where $^{239}$Np is the alpha decay daughter of $^{243}$Am. The specific activity of $^{239}$Np solution was determined by a $4\pi$$\beta$($PC$)-$\gamma$ coincidence counting device. There were 6 gamma sources prepared to measure with the HPGe detector, and the activity of $^{239}$Np in each gamma source was calculated with the weights of the solution contained in it. The emission probability of 106.1 keV of $^{239}$Np is measured to be $(25.4 \pm 0.3)\%$, which is consistent with 25.34%, the value evaluated in 2014.

1. Introduction

The neptunium isotope $^{239}$Np is a short-lived $\gamma$ and $\beta$ particle emitting radionuclide, with a half-life about 56.5 hours. It decay towards $^{239}$Pu with $\beta$ and $\gamma$ particle emission. It can be produced by disintegration of $^{243}$Am. $^{239}$Np is of great importance as a chemical yield tracer for the radiochemical determination of $^{237}$Np in samples [1]. The absence of standardized solutions and the poor quality of the associated decay scheme data inhibited the use of $^{239}$Np. For this purpose, gamma emission probabilities of $^{239}$Np were determined in this work.

2. Comparison of $^{239}$Np half-life and decay scheme

Table 1 shows the half-life and $\gamma$-ray emission probabilities of $^{239}$Np given by several different nuclear databases [2–5]. It can be seen that there is obvious differences between these results. In this paper, we focus on the $\gamma$-ray emission probability determination.

3. Sample chemical separation

The original $^{243}$Am solution contains both $^{243}$Am and $^{239}$Np. In order to get pure $^{239}$Np solution, chemical separation should be done by the following steps:

1) Appropriately mix $^{243}$Am solution and 10 ml HNO$_3$ solution of 0.1 M concentration;
2) Transfer the solution mentioned above to a CL-P204 cylinder, collect the solution outflow from this cylinder which should only contains $^{239}$Np, and record the time;
3) Add 10 ml HNO$_3$ solution of 1.0 M concentration, and collect the solution outflow which should only contains $^{243}$Am.

After getting a solution of $^{239}$Np, six sources were prepared to determined the specific activity of $^{239}$Np solution with the coincidence method, and three sources were measured by a HPGe detector whose full-energy peak efficiency curve had already been obtained.

4. Measurement of $^{239}$Np activity

Measurement of specific activity of $^{239}$Np concentration was achieved by $4\pi$$\beta$($PC$)-$\gamma$ coincidence counting system which is a standard activity counting system. The system consists of a beta counting channel constructed from a proportional counter(PC) with P10 gas used as the counting medium and a gamma counting channel from a NaI detector.

Corrections for background, dead time and accidental coincidence were carried out using classical formulae. A linear extrapolation is usually needed in this method to estimate source activity when the counting efficiency of beta channel approaches 100%. The description of linear extrapolation is as follow [6]

$$\frac{N_\beta N_\gamma}{N_c} = N_0 \left( 1 + K \frac{1 - \epsilon_\beta}{\epsilon_\beta} \right)$$

(1)

$$\epsilon_\beta = \frac{N_c}{N_\gamma}$$

(2)

Where $N_\beta$, $N_\gamma$ and $N_c$ are the counting rates of the beta channel, gamma channel and coincidence channel.

Six sources with masses between 68.5 mg and 92.7 mg were prepared onto metalized VYNS foils. Each source was measured with linear extrapolation by varying the threshold of beta channel to change the counting efficiency.

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Table 1. The half-life and γ-ray emission probabilities of 239Np.

| Data Base | ENDF 7.1 | TOI | BIPM | Nuclear data sheet |
|-----------|---------|-----|------|-------------------|
| T1/2      | 2.356d  | 2.356d | 2.356d(3d) | 2.356d |
| Eγ/keV    |         |      |       |                   |
| 106.123   | 26.3(10)| 27.2(4) | 25.9(3) | 25.3(4) |
| 209.753   | 3.42(3) | 3.42(5) | 3.42(3) | 3.36(3,20) |
| 228.183   | 11.14(11) | 10.76(16) | 11.32(22) | 10.73(9) |
| 277.599   | 14.4(10) | 14.38(21) | 14.4(1) | 14.51(8) |
| 315.88    | 1.60(2)  | 1.60(3) | 1.59(1) | 1.60(12) |
| 334.31    | 2.06(2)  | 2.07(3) | 2.04(2) | 2.056(13) |

Table 2. Specific activities determined by each source.

| Number | Mass/g | Specific Activity (Bq/g) |
|--------|--------|--------------------------|
| 1      | 0.0685 | 31220.72                 |
| 2      | 0.07425 | 31438.01                |
| 3      | 0.07679 | 30733.17                |
| 4      | 0.09266 | 31050.74                |
| 5      | 0.07643 | 31089.89                |
| 6      | 0.08329 | 31196.49                |
| Average|        | 31157.51                 |

Table 3. Uncertainty components in the standardization of 239Np solution.

| Components                          | Value(%) (k = 1) |
|-------------------------------------|-----------------|
| Extrapolation of efficiency and Counting statistics(A) | 0.77            |
| Resolving time(B)                   | 0.1             |
| Dead time(B)                        | 0.12            |
| Background(B)                       | 0.1             |
| weighing(B)                         | 0.1             |
| Half time(B)                        | 0.02            |
| Combined (k = 1)                    | 0.80            |

The gamma ray emission rate of 239Np was calculated from the gamma-ray spectrum, and then the gamma ray emission probabilities of 239Np can be determined by Eq. (3) [8].

\[
P = \frac{N_r}{\varepsilon_r A T_L} \lambda T_R (1 - e^{-\lambda T_R}) e^{\lambda T}
\]

Where \(N_r\) is the is the full-energy peak areas of each gamma-ray, \(\varepsilon_r\) is the full-energy efficiency of each gamma-ray, \(A\) is the activity of 239Np at time zero, \(\lambda\) is the decay constant of 239Np, \(T_R\) is the real time of measurement, \(T_L\) is the live time of measurement, \(T\) is the time from time zero to the beginning of measurement.

5. Gamma ray emission probabilities

Gamma ray emission rate of 239Np was determined by a N-type coaxial HPGe detector. The detector was calibrated by 241Am, 133Ba, 60Co, 152Eu and 155Eu. Activity of all these sources had already been determined by standardized measurement system with the relative standard uncertainties less than 1%. The decay data of radioactive nuclides was selected from gamma ray decay data standards for detector calibration recommended by the IAEA in 2007. The full-energy efficiency function of the HPGe is shown in Fig. 1. The full-energy peak efficiencies of gamma-ray from decay of 239Np was obtained by intercepting the absolute full-energy peak efficiency curve also with the relative standard uncertainties less than 1% [7].

6. Conclusions

In this work, the principal γ emission of six energies in 239Np decay was experimentally determined as shown in Table 4. The use of absolute counting techniques to determine the activity and calibrated HPGe γ detector has been of great importance in this work. There are still some differences between the values proposed in this work and the currently evaluated data. The emission probability of 106.1 keV of 239Np is measured to be (25.4 ± 0.3)%γ, which is consistent with 25.34%, the value evaluated in 2014.

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