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Activation cross sections of deuteron-induced reactions on niobium up to 24 MeV



BEAM INTERACTIONS WITH MATERIALS AND ATOMS

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ABSTRACT

We measured cross sections of deuteron-induced reactions on a niobium target up to 24 MeV using the stacked-foil activation technique and off-line γ -ray spectrometry. The production cross sections of 93m Mo, 92m Nb and 89 Zr were determined. The results were compared with the experimental data studied earlier and the predictions of theoretical calculations.

1. Introduction

There are several radioisotopes of Mo, Nb and Zr suitable for medical use. One of the potential radioisotopes for nuclear medicine among these elements is ^{93m}Mo [1]. This radioisotope has features of a moderate half-life ($T_{1/2} = 6.85$ h), γ -lines ($E_{\gamma} = 263.049$, 684.693 and 1477.138 keV with $I_{\gamma} = 57.4$, 99.9 and 99.1%, respectively), conversion electrons ($E_e = 243.050$ keV with $I_e = 29.7\%$) and Auger electrons ($E_e = 2.27$ keV with $I_e = 37.9\%$) [2], which were summarized in NuDat 2.7 [3]. With these decay parameters, ^{93m}Mo has great potential for use in medical diagnosis and/or therapy.

Several reactions to produce 93m Mo were studied earlier, such as proton- (see e.g. [4] and references therein) and deuteron-induced reactions on Nb [5–8], α -induced reactions on Zr [9] and ⁷Li-induced reactions on Y [10,11]. The experimental literature was surveyed in the EXFOR database [12]. In this paper, we focused on the deuteron-induced reactions on Nb because the cross sections are about four times larger than those of the proton-induced reactions [8]. In addition, the ^{93m}Mo production cross sections at around the peak energy of 17 MeV are scattered over several tens of mb [6,7]. Therefore, in order to obtain an accurate and smooth curve as a function of projectile energy, an experiment was performed to measure the cross sections of 93m Mo, 92m Nb and 89 Zr. Based on the result, the integral yield of 93m Mo was also deduced.

2. Method

We used the standard methods, such as the stacked-foil activation

method and off-line γ -ray spectrometry. The stacked target was composed of ^{93}Nb (99.9% purity, Nilaco Corp., Japan) and ^{nat}Ti (99.6% purity, Nilaco Corp., Japan) foils for monitoring the beam parameters. Niobium is a mono isotopic element, ^{93}Nb . The thicknesses of the Nb and Ti foils were 27.11 and 9.13 mg/cm², respectively, which were estimated from measurement of surface area and weight of larger sheets. After the thickness measurement, we cut the large sheets into $10 \times 10 \text{ mm}^2$ rectangular pieces to fit into our target holder.

The target was irradiated by a 23.6 MeV deuteron beam at the AVF cyclotron of the RIKEN RI Beam Factory. The irradiation lasted for 30 min with an average intensity of 200.3 nA. The intensity was measured by a Faraday cup. The incident energy was 23.6 ± 0.1 MeV, which was determined by the time-of-flight method using a plastic scintillator monitor [13]. The degradation of the beam energy in the stacked target was calculated using the SRIM code [14]. The propagation of the initial energy uncertainty is also estimated through the target. These beam parameters can be assessed by the ^{nat}Ti(d,x)⁴⁸V monitor reaction.

The irradiated foils were separated from each other shortly after the end of bombardment. The γ -lines from the foils were measured by a high-resolution HPGe detector (ORTEC GMX-25190-P) and analyzed by dedicated software, Gamma Studio (SEIKO EG&G). The detector was calibrated using a multiple standard γ -ray point source consisting of ^{57,60}Co, ⁸⁸Y, ¹⁰⁹Cd, ¹¹³Sn ¹³⁷Cs, ¹³⁹Ce and ²⁴¹Am radioisotopes. The reaction and decay data used for the γ -ray spectrometry were taken from NuDat 2.7 [3], LiveChart [15] and QCalc [16] (Table 1).

We derived the cross sections of the $^{nat}Ti(d,x)^{48}V$ monitor reaction and compared the result with the IAEA recommended values [17] to assess the beam parameters. The cross sections were obtained from

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Table 1

Reactions and decay data for reaction products.

Nuclide	Half-life	Decay mode (%)	E _γ (keV)	I _γ (%)	Contributing reactions	Q-value (MeV)	Refs.
^{93m} Mo	6.85 h	IT (99.88) β ⁻ (0.12)	263.049	57.4(11)	⁹³ Nb(d,2n)	-3.4	[2]
^{92m} Nb ⁸⁹ Zr	10.15 d 78.41 h	$\varepsilon + \beta^+ (1 \ 0 \ 0)$ $\varepsilon + \beta^+ (1 \ 0 \ 0)$	934.44 909.15	99.15(4) 99.04(3)	⁹³ Nb(d,t) ⁹³ Nb(d,α2n)	-2.6 -7.8	[18] [19]
⁴⁸ V	15.97 d	- F ()	983.5	99.98(4)	^{nat} Ti(d,x)		[20]



Fig. 1. Excitation function of the $^{\rm nat} Ti(d,x)^{48} V$ monitor reaction compared with the recommended values.

measurements of the 983.5-keV γ ray (I $_{\gamma}$ = 99.98%) emitted from the ⁴⁸V decay (T_{1/2} = 15.97 d) after an adequate cooling time for decay of the co-produced ⁴⁸Sc (T_{1/2} = 43.67 h). The result of the comparison with the IAEA recommended values is shown in Fig. 1, which shows good agreement without any adjustments. According to the comparison, the correctness of the measured beam intensity and the energy calculation in the targets was guaranteed.

3. Result and discussion

The derived cross sections are presented in Table 2 and graphically in Figs. 2–4. Only direct reactions in Table 2 are energetically possible for the three products and hence the reported cross sections are independent. The results are compared with the data measured earlier [5–7] and the TENDL-2017 data [21]. The data in the literature [8] were excluded because their energy range of 30–50 MeV was not

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Energy (MeV)	⁹³ Nb(d,2n) ^{93m} Mo (mb)	⁹³ Nb(d,x) ^{92m} Nb (mb)	⁹³ Nb(d,x) ⁸⁹ Zr (mb)
$\begin{array}{c} 22.7 \pm 0.4 \\ 22.1 \pm 0.4 \\ 21.5 \pm 0.4 \\ 19.6 \pm 0.4 \\ 19.0 \pm 0.5 \\ 18.3 \pm 0.5 \\ 16.2 \pm 0.5 \\ 15.4 \pm 0.5 \\ 15.4 \pm 0.5 \\ 14.6 \pm 0.6 \\ 12.1 \pm 0.6 \\ 11.2 \pm 0.7 \end{array}$	$\begin{array}{c} 64.5 \pm 5.3 \\ 70.6 \pm 5.8 \\ 81.5 \pm 6.7 \\ 100.3 \pm 8.2 \\ 108.3 \pm 8.9 \\ 114.4 \pm 9.4 \\ 113.2 \pm 9.3 \\ 107.3 \pm 8.8 \\ 98.6 \pm 8.1 \\ 62.8 \pm 5.1 \\ 48.7 \pm 4.0 \end{array}$	(mb) 53.9 ± 4.4 45.6 ± 3.8 41.2 ± 3.4 24.0 ± 2.0 16.4 ± 1.4 13.9 ± 1.2 7.0 ± 0.6 6.5 ± 0.6 5.5 ± 0.5 3.9 ± 0.4 3.3 ± 0.3	(mb) 9.1 \pm 0.7 6.4 \pm 0.6 4.5 \pm 0.4 0.83 \pm 0.11 0.36 \pm 0.09
11.2 ± 0.7 10.2 ± 0.7	48.7 ± 4.0 33.6 ± 2.7	3.3 ± 0.3 2.4 ± 0.3	
10.2 ± 0.7 6.8 ± 0.9	33.6 ± 2.7 0.94 ± 0.08	2.4 ± 0.3	
5.3 ± 1.1 3.5 ± 1.4	0.016 ± 0.001 0.0038 ± 0.0003		



Fig. 2. The excitation function of the ⁹³Nb(d,2n)^{93m}Mo reaction.



Fig. 3. The excitation function of the ${}^{93}Nb(d,x){}^{92m}Nb$ reaction.

overlapped with our data.

The estimated total uncertainty of the cross sections was 8.0–24.4%. It was the square root of the quadratic summation of each component; statistical uncertainty (< 23.1%), target thickness (1%), target purity (1%), beam intensity (5%), detector efficiency (5%), γ -intensity (< 1.1%) and peak fitting (3%).

3.1. The ${}^{93}Nb(d,2n){}^{93m}Mo$ reaction

The γ -line at 263.049 keV (I $_{\gamma} = 57.4\%$) from the ^{93m}Mo IT decay (T $_{1/2} = 6.85$ h) was measured after a cooling time of about 10 h. The derived excitation function of the ⁹³Nb(d,2n)^{93m}Mo reaction is shown in Fig. 2 together with the earlier experimental data [5–7] and the TENDL-2017 data [21]. Our result shows a very smooth curve and good agreement with the other experimental data in the whole energy region. The theoretical calculation overestimates the experimental data.

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