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Half-life of ²²⁸Pu and α decay of ²²⁸Np

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Two α -decay chains starting from ²²⁸Pu produced in the reaction of ³⁴S+¹⁹⁸Pt were observed, and the half-life was measured to be $1.1^{+2.0}_{-0.5}$ s. The half-life follows the Geiger-Nuttall curve for even-even Pu isotopes, which shows that α decay is the dominant decay mode. In addition, five α -decay events of ²²⁸Np were observed for the first time. The evaporation residue cross sections of ²²⁸Pu, ²²⁸Np, and ²²⁵U are reproduced by a statistical model calculation.

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I. INTRODUCTION

Plutonium-228 was produced and identified for the first time by Andreyev *et al.* [1] in the fusion reaction of 24 Mg $+^{208}$ Pb. However, they were not able to determine the half-life of 228 Pu.

In the present paper we report on the first measurement of half-life of ^{228}Pu produced in the fusion reaction $^{34}\text{S}+^{198}\text{Pt}$ (compound nucleus ^{232}Pu). Our experimental method is similar to that in Ref. [1]. The evaporation residues (ERs) were separated by the JAERI recoil mass separator (JAERI-RMS) [2], and then implanted into a focal plane detector. The identification of the nucleus was made by observing the α -decay chain. The lifetime of the ER is measured by the time interval between the recoil implantation and its α decay. In-flight separation allows us to measure the nucleus with a short half-life.

For the actinide nuclei with proton number of Z=90–94 and neutron number of N=134–138, α -particle emission and electron capture (EC) are two competing decay modes. For 228 Pu, a calculation suggests that the α -decay partial half-life, $T_{\alpha,1/2}$ =0.42 s [3], is two orders of magnitude shorter than that of EC decay $T_{\rm EC,1/2}$ =44 s [4]. It is known that the Geiger-Nuttall curve shows the relation between the partial half-life $T_{\alpha,1/2}$ and Q value Q_{α} of α decay. If the experimental half-life follows this curve, one can estimate that the decay is dominated by α decay.

In addition, the α decay of 228 Np was observed in the present measurement. The first production of this nucleus was made by Kuznetsov *et al.* [5], who determined the half-life of fission activity as 60 s in the reaction 22 Ne+ 209 Bi. The EC-delayed fission properties of 228 Np was studied by Kreek *et al.* [6] in detail. So far there is no report on the measurement of the α decay of 228 Np.

The ER cross sections for ²²⁸Pu, ²²⁸Np, and ²²⁵U were also determined. The results were compared with a statistical model calculation, and the discussions are given in this paper.

II. EXPERIMENT

The fusion reaction $^{34}S + ^{198}Pt$ was used to produce the neutron deficient nuclei $^{228}Pu(4n)$, $^{228}Np(p3n)$, and

²²⁵U(α3n). The ³⁴S ions were accelerated to $E_{\rm beam}$ =170 and 172 MeV by the JAERI-tandem accelerator and irradiated a ¹⁹⁸Pt target. Typical beam current was 40–50 pnA. The target with thickness 390 μg/cm² was made by sputtering an enriched ¹⁹⁸Pt material (98%) on a 1.2-μm- thick aluminum (Al) foil. The above bombarding energy corresponds to the center-of-mass energy ($E_{\rm c.m.}$) of 141 and 143 MeV, respectively, at the half depth of the target layer, which was determined by calculating the energy loss of the beam in the Al foil and target.

The evaporation residues emitted in the beam direction were separated in flight from the primary beams and other reaction products by the JAERI-RMS. Before entering the recoil mass separator, the ER's charge state was reset by passing through a 30 μ g/cm² carbon foil (charge reset foil). The detection efficiencies of ERs by the JAERI-RMS were calculated by the method described in Ref. [7], where we obtained good agreement between the measured detection efficiency and that calculated by using a program code GIOS [8]. The angular distributions of ERs were estimated by using the statistical model code PACE2 [9]. The multiple scattering of the ERs in the target and the charge reset foil were estimated by the TRIM code [10]. The JAERI-RMS was set to transport particles of 16⁺ charge state, which has the probability of 0.18 [11] in the charge state distribution. This probability was multiplied by the transport efficiency through the JAERI-RMS to yield the detection efficiency. Thus, the detection efficiencies for 4n, p3n, and $\alpha 3n$ channels are determined to be 0.086, 0.074, and 0.050, respectively. The same procedure was also adopted in the previous measurement of ER cross sections [12].

The ERs transported through the JAERI-RMS were implanted into a double-sided position-sensitive silicon detector (PSD; 73×55 mm²) located in the focal plane. The identification of the nucleus is made by constructing an α -decay chain and finding the known α -particle energies (and also lifetimes) of descendants, whose decay position agrees with that of the recoil implantation. The front side of PSD contains 15 position-sensitive strips with resistive layer. The signals induced on both sides of each strip were amplified with a shaping time of 0.3 μ s, and the pulse heights give the energy and position of X. The backside is divided into eight sections, and the position Y in each section can be deter-

mined. The pulse heights were digitized by an analog-to-digital converter with the gate width of 1 μ s.

The α decay in the PSD is distinguished from the transported particle by determining that no time-of-flight (TOF) signal is present, which is measured by two timing detectors separated by the distance of 30 cm and placed upstream the PSD. In the off-line data analysis, the ERs were separated from the scattered beam particles on the two-dimensional spectrum of the TOF versus energy. This process considerably reduces the chance coincidence in finding a recoil- α correlation. Here, recoil means the event of the ER to hit the PSD. The dead time arising from the data acquisition process is $200~\mu s$, and an α decay whose lifetime is shorter than the period will be missed.

The energy calibration of the PSD was made using α lines from the known nuclei [13], 216 Th(E_{α} =7921 keV), ²¹²Ra(6899), ²¹⁵Ac(7604), ²¹¹Fr(6534), ²¹¹Ra(6910). ²¹⁵Th(7524, 7395), ²⁰⁷Rn(6131), 210 Ra(7019), 206 Rn(6258), produced in the reaction 34 S+ 186 W. In this measurement, we constructed the spectra of position difference $(\Delta X \text{ and } \Delta Y)$ between the recoil implantation and α decay for the chains of $recoil-\alpha(^{216}Th)-\alpha(^{212}Ra)$ and $recoil-\alpha(^{215}Th)$ $-\alpha(^{211}\text{Ra})$. The widths of ΔX and ΔY spectra were 0.15 mm and 0.26 mm [full width at half maximum (FWHM)], respectively. In the data analysis we searched for the correlated α -decay event whose position agreement is $(|\Delta X|, |\Delta Y|)$ <(0.2, 0.4) mm.

The gain stability of the detection system was checked by exposing the PSD to α particles from an ²⁴¹Am source. Typical energy resolution of the PSD was 75 keV (FWHM). In the data analysis, the discrimination level was set at about 2 MeV.

III. EXPERIMENTAL RESULTS AND DISCUSSIONS

Figure 1 shows the energy spectrum obtained from the PSD. The spectrum (a) shows the events which do not generate TOF signals and includes all events taken during a 41-h run at the reaction energy of $E_{c.m.}$ =141 MeV. The large peak at 5.4 MeV represents α particles from the external α source (241Am) which irradiated the silicon detector during the measurement. The broad 2-6 MeV spectrum is formed by scattered beam particles transported through the JAERI-RMS. As the detection efficiency of the timing detectors was not 100%, such background particles are not fully rejected and appear in the spectrum (a). There are several α lines in Fig. 1(a) including α decays of 216 Th(7921 keV, 28 ²¹⁵Ac(7604, 0.17 s), ²¹⁵Th(7524:7395, 1.2 s), ²¹²Ra(6899, 13 s), ²¹¹Ra(6.910, 13 s), ²¹¹Fr(6534, 3.1 min), ²⁰⁷Rn(6131, 9.3 min), and ²⁰⁷At(5758, 1.8 h). These nuclei are produced by the reaction of the beam with ¹⁸⁶W nucleus which is contained in the ¹⁹⁸Pt target.

The correlation between the recoil implantation and the subsequent α decays were searched for within the time interval $\Delta t(recoil-\alpha)$ of 10 s. We only selected chains in which the recoil implantation is followed by two or more α decays. We obtained 18 chains and the corresponding α -particle energy spectrum is shown in Fig. 1(b). In this process, the condition was imposed that the recoil event generates the

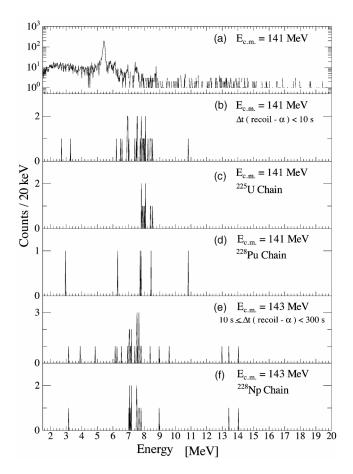


FIG. 1. Energy spectra obtained from the PSD in the reaction of $^{34}\mathrm{S}+^{198}\mathrm{Pt}$. Spectra in (a)–(d) include all data for measurements where $E_{\mathrm{c.m.}}=141$ MeV. Spectra in (e) and (f) are obtained in the measurements where $E_{\mathrm{c.m.}}=143$ MeV.

TOF and energy signal corresponding to the ERs on the twodimensional spectrum to reject chance events associated with the scattered particle.

Among the 18 chains in Fig. 1(b), we found four chains starting from the ²²⁵U implantation based on detected α -particle energies and half-lives. The E_{α} spectrum of the ²²⁵U chain is shown in Fig. 1(c). Here, the events of full energy absorption for the α decay of ²²⁵U itself and ²²¹Th are selected. Table I shows the α -decay character (kinetic energy and lifetime) for the 225 U chains. When the α -particle energy of ²²¹Th is around 8472 keV instead of 8146 keV, the identification of the ²²⁵U chain requires the detection of ²¹³Rn. This is because the α -particle energies for $^{225}\mathrm{U}(E_{\alpha}$ =7879,7821 keV)- $^{221}\text{Th}(8472)$ are close to that of 228 Pu(7810[1])- 224 U(8466) and the lifetimes of 225 U(95 ms) and 221 Th(1.68 ms) are also close to that of 228 Pu(1.1 s; see below) and 224 U(0.9 ms), respectively. The short-lived nucleus ²¹⁷Ra does not appear in Table I because of the experimental limitation. The other chains in Fig. 1(b) include those starting from ²¹⁶Th (two chains) and ²¹⁵Th (three chains) produced in the ³⁴S+¹⁸⁶W reaction. The same process was adopted for the measurement at $E_{c.m.}$ =143 MeV, in which two chains starting from the ²²⁵U implantation were found as listed in Table I.

TABLE I. α -Decay energy (in keV) and lifetime (given in parentheses) starting from the recoil implantation of 225 U ($\alpha 3n$ channel). Kinetic energy of α particle and half-life (in square brackets) taken from the literature [13] is shown in the first line. Intensities of α lines are shown in % for 225 U and 221 Th. The second column shows the center-of-mass energy $E_{c.m.}$ in MeV.

No.	$E_{ m c.m.} \ m (MeV)$	²²⁵ U 7879 ^{85%} , 7821 ^{15%} [95 ms]	²²¹ Th 8472 ^{39%} , 8146 ^{56%} [1.68 ms]	²¹⁷ Ra 8992[1.6 μs]	²¹³ Rn 8088[25 ms]
I-1	141	7828(3.2 ms)	8415(4.6 ms)		8024(31 ms)
I-2	141	7898(47 ms)	8071(1.7 ms)		
I-3	141	7823(210 ms)	8548(1.3 ms)		7976(15 ms)
I-4	141	7873(613 ms)	8400(1.8 ms)		8074(8.1 ms)
I-5	143	7875(198 ms)	8164(4.0 ms)		
I-6	143	7973(99 ms)	8238(7.2 ms)		8177(4.3 ms)

The half-life of 225 U was determined to be 135^{+93}_{-39} ms. This agrees with 80 ± 40 ms [14] and 95 ± 15 ms [15], where 225 U is produced in the reactions of 180 Hf(48 Ca, $_3n$) [14] and 209 Bi(19 F, $_3n$) [15]. There are reports of a slightly shorter half-life of 30^{+20}_{-10} ms [16] and 59^{+5}_{-2} ms [17], both of which used the reaction 208 Pb(22 Ne, $_5n$) to produce 225 U.

Among the nine chains in Fig. 1(b) that were not identified in the above process, we searched for one that is followed by a long-lived nucleus with the help of position agreement. The searching time is extended to 120 min, long enough to observe the α decay of ²¹²Rn ($T_{1/2}$ =23.9 min [13]) as the descendant of ²²⁸Pu. In this process, we found two decay chains, and the E_{α} spectrum is shown in Fig. 1(d). They are both attributed to the ²²⁸Pu chain based on their decay character as shown in Table II. The event II-1 has the α decay of 10817 keV, which may be a partial pileup of 220 Th and 216 Ra decays. The chain is terminated by the α decay of ²¹²Rn. The event II-2 is determined to be a ²²⁸Pu-chain from the decay character of ²²⁴U and the subsequent α decay after 11 min as the candidate for ²¹²Rn decay. The α particle from ²¹²Rn of II-2 is considered to have escaped from the PSD, and part of the energy deposition is detected in the PSD. We have estimated the probability of a chance coincidence signal of 2-20 MeV to be detected as the α decay in the time interval of 30 min. This depends on the counting rate of the PSD and the detection efficiency of the timing detector placed upstream the PSD. The obtained probability is 0.013. The same procedure was adopted to the $E_{\rm c.m.}$ =143 MeV run. We detected two chains having decay character of $\alpha_1(E_{\alpha}=7853 \text{ keV}, \tau=4.5 \text{ ms}) - \alpha_2(6893, 7.37 \text{ s})$ $-\alpha_2(6116, 53 \text{ s})$ and $\alpha_1(7617, 0.13 \text{ s}) - \alpha_2(7048, 3.2 \text{ s})$ $-\alpha_3(6316, 13 \text{ min})$. The former and the latter were

determined to be decay chains of 216 Th(E_{α} =7921, $T_{1/2}$ =28 ms) $-^{212}$ Ra(6899, 13 s) $-^{208}$ Rn(6144, 24 min) and 214 Th(7678, 0.1 s) $-^{210}$ Ra(7019, 3.7 s) $-^{206}$ Rn(6258, 5.7 min), respectively.

The half-life of 228 Pu was measured to be $1.1^{+2.0}_{-0.5}$ s. Here, the error is estimated by the method in Ref. [18]. The obtained average α -particle energy of ²²⁸Pu, 7772±35 keV, reasonably agrees with the data [1]. The present E_{α} and halflife are compared with the prediction of the Geiger-Nuttall law. In order to determine the α -decay Q-value Q_{α} for ²²⁸Pu, we assumed that the detected α decay is from ground-state to ground-state transition, and the electron screening effect [19] is corrected for. The $Q_{\alpha}(7948\pm36 \text{ keV})$ and $T_{1/2}$ are plotted in Fig. 2 together with the other data [13]. In this figure, the Geiger-Nuttall curves for elements Th, U, and Pu are drawn by using the expression and constants in Ref. [19]. Our data of ²²⁸Pu match the characteristics of Pu isotopes. This means that the 228 Pu decay is dominated by the α decay. The dominance of α decay is supported by the theoretical calculation that the partial half-life for α decay $T_{\alpha,1/2}$ =0.42 s [3] is two orders of magnitude shorter than that for EC decay $T_{\rm EC,1/2}$ =44 s [4], as described in Sec. I. Even ²³⁰Pu, which has two more neutrons and so may be expected to have a larger EC branch than ²²⁸Pu, has a dominant α -decay probability of 84% [20]. The ²³⁰Pu data shown in Fig. 2 refers to the experimental half-life of 102 s [21], α -decay probability (84%), and Q_{α} value of 7213 keV obtained by using E_{α} =7050 keV

In order to find the α decay of ²²⁸Np with half-life of 61.4 s [6], the recoil- α chain was searched for in the time span of 10 s $\leq \Delta t (recoil - \alpha) < 300$ s. We set the condition that the *recoil* implantation is followed by two or more α decays in this

TABLE II. α -Decay energy (in keV) and lifetime (given in parentheses) starting from the recoil implantation of 228 Pu (4n channel). The literature value of kinetic energy of α particle and half-life (in square brackets) for 228 Pu [1] and the other nuclei [13] is shown in the first line. The time with symbol "<" represents the time interval relative to the preceding α decay. The signal with escaped event is indicated by "esc." The signal of pileup, "pil," is caused by the α decays of a mother and a short-lived daughter indicated by " \leftarrow ."

No.	E _{c.m.} (MeV)	²²⁸ Pu 7810	²²⁴ U 8466[0.9 ms]	²²⁰ Th 8790[9.7 μs]	²¹⁶ Ra 9349[0.18 μs]	²¹² Rn 6264[23.9 min]
II-1 II-2	141 141	7807(0.35 s) 7736(2.76 s)	8446(3.2 ms)	10817 _{pil} (<0.77 ms)	←	6309(34 min) 2960 _{esc} (11 min)

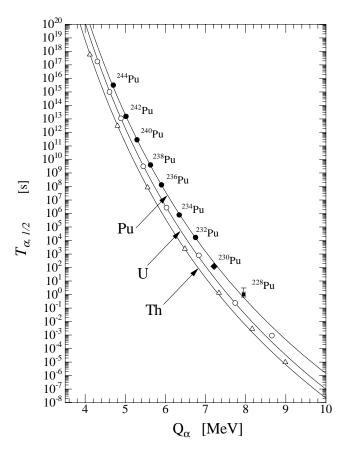


FIG. 2. α -Decay energy and half-life for ²²⁸Pu (solid square with error bar) is plotted on the map of $T_{\alpha,1/2}$ versus Q_{α} together with the other nuclei (solid circle is for Pu, open circle for U, and open triangle for Th [13]). The $T_{\alpha,1/2}$ value for ²³⁰Pu is shown by the solid diamond. Geiger-Nuttall curve for even-even Pu, U, and Th isotopes [19] are shown by the solid curves.

time region. Figure 1(e) shows the spectrum of E_{α} for the obtained 15 decay chains in a 29-h run of $E_{\rm c.m.}$ =143 MeV. In these chains, five chains starting from $^{228}{\rm Np}$ are obtained as shown in the E_{α} spectrum of Fig. 1(f), and the decay properties are listed in Table III. Here, we selected events in which the full energy of an α decay of $^{228}{\rm Np}$ is absorbed. The same procedure was adopted for the $E_{\rm c.m.}$ =141 MeV measurement, but we observed no candidate for the production of $^{228}{\rm Np}$.

The time interval between the recoil implantation and ²²⁸Np decay ranges from 15 s to 196 s. In finding the recoil implantation event of ²²⁸Np, the probability of a random event in 200 s is estimated to be about 17%. This probability decreases with decrease in the lifetime of the implanted ER. The obtained half-life 56^{+45}_{-17} s of 228 Np agrees with 61.4 s [6]. The daughter 224 Pa has the α -particle energy of E_{α} =7488 keV (70% abundance) and many other energies [13]. The detected four events of ²²⁴Pa are all close to the energy of the main branch. Events of III-1 and III-3 have the large energies 14048 and 13425 keV. This apparently comes from the pileup of the decays of 220 Ac and 216 Fr(0.7 μ s). In our data acquisition system, the pulse height generated by the sequential α decays depends sensitively on the time interval on the sub- μ s scale, and is given by the sum of the signal of the first pulse and a portion of the second one. When the second decay is delayed by $1 \mu s$, this signal is rejected. Thus, the reason for the large energies should be that the full energies of the α particles of ²²⁰Ac and ²¹⁶Fr are deposited in the PSD with the time difference less than 1 μ s, and part of the second signal is piled to the first one. The event III-5 does not have decay signals of ²²⁴Pa and ²²⁰Ac, but the full energy absorption of ²¹⁶Fr is obtained. The time interval of this event (3.39 s) is the sum of lifetimes of ²²⁴Pa, ²²⁰Ac, and ²¹⁶Fr. Astatine-212 with 0.134 s [13] has dominant groundstate to ground-state α decay of 7679 keV (82%) and many branches to the excited states of ²⁰⁸Bi. The two decays of ²¹²At (7687 and 7658 keV) in Table III are the ground-state to ground-state transition. The 7012 keV event of III-3 seems to correspond to the 7045-keV transition which has low frequency (0.45%).

In the measurement with the beam energy $E_{\rm c.m.}$ = 143 MeV, we detected one event of a α_1 - α_2 correlation with energies of $E_{\alpha,1}$ =6679 keV and $E_{\alpha,2}$ =7483 keV. The time difference between the two α decays was 0.49 s. They are identified to be ²²⁸U and ²²⁰Ra, respectively, in the chain [13],

228
U(E_{α} = 6680 keV, $T_{1/2}$ = 9.1 min)
 \rightarrow 224 Th(7170, 1.05 s) \rightarrow 220 Ra(7455, 18 ms)
 \rightarrow 216 Rn(8050, 45 μ s) \rightarrow 212 Po(8784, 0.3 μ s)
 \rightarrow 208 Pb.

Since the α particle from ²²⁴Th escaped from detection by

TABLE III. α -Decay energy (in keV) and lifetime starting from the recoil implantation of 228 Np (p3n channel). The literature value for 228 Np [6] and the other nuclei [13] is shown in the first line. See the captions of Tables I and II for the detailed explanation.

No.	$E_{\mathrm{c.m.}}$	²²⁸ Np [61.4 s]	²²⁴ Pa 7488 ^{70%} [0.79 s]	²²⁰ Ac 7855 ²⁶ %[26.4 ms]	²¹⁶ Fr 9005[0.7 μs]	²¹² At 7679 ^{82%} , 7045 ^{0.45%} [0.314 s]
III-1	143	7183(29 s)	7529(1.10 s)	14048 _{pil} (3.7 ms)	←	3170 _{esc} (0.070 s)
III-2	143	7062(128 s)	7543(0.20 s)	7794(58 ms)		7687(2.4 ms)
III-3	143	7126(196 s)	7495(0.56 s)	13425 _{pil} (21 ms)	←	7012(0.12 s)
III-4	143	7177(35 s)	7521(3.03 s)	•		7658(0.18 s)
III-5	143	7065(15 s)		8969(<3.39 s)		

the PSD, the measured time interval 0.49 s is the sum of lifetimes of ²²⁴Th and ²²⁰Ra. Radon-216 and ²¹²Po were not detected due to the dead time of our data acquisition system. The implantation event to the PSD was found at 18 min before the α decay of ²²⁸U. Because of the increased probability of a random event in such a long time span in searching for the recoil implantation event, we can only state that 18 min is the lower limit for the lifetime of ²²⁸U. The ²²⁸U is not directly produced as an ER in the present reaction but is produced as a daughter of the EC decay of 228 Np. The detected number of α -decay events for ²²⁸U (one event) and ²²⁸Np (five events) leads to the α -decay probability of $83^{+17}_{-36}\%$ for $^{228}{\rm Np}.$ This is comparable to 40^{+8}_{-6} % reported by Kreek et al. [6], which they determined by measuring the 216 Fr(9005 keV, 0.7 μ s) and ²¹²Po, which are the descendants of the α -decay chains starting from ²²⁸Np and ²²⁸U.

From the mass table of Audi and Wapstra [22], the α -decay Q-value Q_{α} of ²²⁸Np is obtained to be 7415 keV. This results in the α -decay energy of 7285 keV for the ground-state to ground-state α decay of ²²⁸Np. This is 100–220 keV larger than the experimental data. A possible reason is that the α decay predominantly produces the excited states of ²²⁴Pa.

We determined the ER cross sections for ²²⁸Pu, ²²⁸Np, and ²²⁵U. The results are shown in Fig. 3. The errors are in the margin of the statistical error. The data were compared to a statistical model calculation. For this purpose, the partial wave cross section for the fusion ³⁴S+¹⁹⁸Pt was calculated by using the CCDEF code [23], which was then inputted to the HIVAP code [24] to calculate the surviving probability and the ER cross section of the specific channel. In the CCDEF code, we took into account the couplings to inelastic channels of the projectile and target. For ³⁴S, deformation parameter (excitation energy) of the quadrupole and octupole vibrations are β_2 =0.252 (2.13 MeV) [25] and β_3 =0.330 (4.62 MeV) [26], respectively. $\beta_3 = 0.05$ (1.68 MeV) [26] was adopted for the octupole vibration of ¹⁹⁸Pt. We also took into account the static deformation of ¹⁹⁸Pt ($\beta_2 = -0.060$ [27], $\beta_4 = -0.030$ [28]). The calculated fusion cross section σ_{fus} is shown on the upper section of Fig. 3 by the dashed curve. The dotted curve is the result of the one-dimensional barrier penetration model, which gives the barrier height of 141.1 MeV. The ER cross sections calculated by the HIVAP code are shown by the solid curve in each section of the figure. We have to adjust the factor b_{fac} , by which the fission barrier height of the liquid-drop part [29] is multiplied to calculate the fission barrier $B_f = b_{fac}B_{LDM} - \delta W$ from 1.03 [12] to 1.00 so as to obtain reasonable agreement with the experimental data. The δW is the ground-state shell energy correction. The change of the factor reduces the $B_{\rm f}$ value by the amount of 0.11 MeV for ²³²Pu. This modification reduces the cross section by a factor of about 2 with the beam energy which yields the highest cross section for each channel being unchanged.

The experimental ER cross section $17^{+2.2}_{-1.3}$ nb of the 4n channel at $E_{\rm c.m.}$ =141 MeV in the fusion reaction $^{34}{\rm S} + ^{198}{\rm Pt}$ was close to 4 ± 2 nb of the 4n channel of $^{24}{\rm Mg} + ^{208}{\rm Pb}$ reaction at $E_{\rm c.m.}$ =118 MeV [1] which forms the same compound

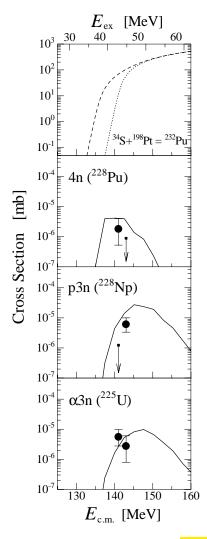


FIG. 3. Evaporation residue cross sections for ²²⁸Pu, ²²⁸Np, and ²²⁵U are shown together with the statistical model calculation (solid curve). The vertical bar with arrow shows the upper limit of the cross section. The fusion cross section calculated by the CCDEF code is shown by the dashed curve, and the fusion cross section of the one-dimensional barrier penetration model is shown by the dotted curve.

nucleus ^{232}Pu . For the $^{24}\text{Mg}+^{208}\text{Pb}$ reaction, we also calculated the cross sections of ^{228}Pu with the same procedure. In the CCDEF calculation, we used deformation parameters β_2 =0.606 (1.37 MeV) [25] and β_3 =0.250 (7.62 MeV) [26] to take into account the couplings to the excited states of ^{24}Mg . For the excitation of ^{208}Pb , parameters β_2 =0.054 (4.09 MeV) [25] and β_3 =0.110 (2.61 MeV) [26] are adopted. The calculated result is 6 nb at $E_{\rm c.m.}$ =118 MeV, reproducing the experimental data in Ref. [1]. Since the excitation energy, $E_{\rm ex}$ =43 MeV, of $^{34}\text{S}+^{198}\text{Pt}$ at $E_{\rm c.m.}$ =141 MeV is close to $E_{\rm ex}$ =44 MeV of $^{24}\text{Mg}+^{208}\text{Pb}$ at $E_{\rm c.m.}$ =118 MeV, the survival probabilities for both systems are nearly identical. The fusion cross section given by the CCDEF code is $\sigma_{\rm fus}$ =69 mb of $^{34}\text{S}+^{198}\text{Pt}$, which agrees with $\sigma_{\rm fus}$ =184 mb of $^{24}\text{Mg}+^{208}\text{Pb}$ within factor of \sim 3.

IV. SUMMARY

The fusion reaction $^{34}\mathrm{S}+^{198}\mathrm{Pt}$ was used to produce $^{228}\mathrm{Pu}$ as the evaporation residue and its half-life was measured to be $1.1^{+2.0}_{-0.5}$ s for the first time. The present half-life follows the Geiger-Nuttall curve for even-even Pu isotopes, which shows that α decay is the dominant decay mode. In this reaction, five α decays of $^{228}\mathrm{Np}$ were observed for the first time. The

ER cross sections of 228 Pu(4*n*), 228 Np(*p*3*n*), and 225 U(α 3*n*) were reproduced by a statistical model calculation.

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