



HAL
open science

Neutron-induced cross sections of short-lived nuclei via the surrogate reaction method

G. Boutoux, B. Jurado, V. Méot, O. Roig, M. Aïche, L. Mathieu, G. Barreau, N. Capellan, I. Companis, S. Czajkowski, et al.

► **To cite this version:**

G. Boutoux, B. Jurado, V. Méot, O. Roig, M. Aïche, et al.. Neutron-induced cross sections of short-lived nuclei via the surrogate reaction method. 5th International Conference FUSION11, 2011, Saint Malo, pp.06005, 10.1051/epjconf/20111706005 . in2p3-00634855

HAL Id: in2p3-00634855

<https://hal.in2p3.fr/in2p3-00634855>

Submitted on 25 Oct 2011

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Neutron-induced cross sections of short-lived nuclei via the surrogate reaction method

G. Boutoux¹, B. Jurado¹, V. Méot², O. Roig², M. Aïche¹, L. Mathieu¹, G. Barreau¹, N. Capellan¹, I. Companis¹, S. Czajkowski¹, J. T. Burke³, E. Bauge², J.M. Daugas², T. Faul², L. Gaudefroy², P. Morel², N. Pillet², P. Romain², J. Taieb², C. Théroine², X. Derckx⁴, O. Sérot⁵, I. Matea⁶, L. Tassan-Got⁶ and F. Gunsing⁷

¹ CENBG, CNRS/IN2P3, Université Bordeaux I, Chemin du Solarium, B.P. 120, 33175 Gradignan, France

² CEA DAM DIF, 91297 Arpajon, France

³ LLNL, Livermore, CA 94550, USA

⁴ GANIL, 14076 CAEN, France

⁵ CEA-Cadarache, DEN/DER/SPRC/LEPh, 13108 Saint Paul lez Durance, France

⁶ IPN, CNRS/IN2P3, Univ. Paris-Sud, 91405 Orsay, France

⁷ CEA Saclay, DSM/DAPNIA/SPhN, 91191 Gif-sur-Yvette cedex, France

Abstract. The measurement of neutron-induced cross sections of short-lived nuclei is extremely difficult due to the radioactivity of the samples. The surrogate reaction method is an indirect way of determining cross sections for nuclear reactions that proceed through a compound nucleus. This method presents the advantage that the target material can be stable or less radioactive than the material required for a neutron-induced measurement. We have successfully used the surrogate reaction method to extract neutron-induced fission cross sections of various short-lived actinides. In this work, we investigate whether this technique can be used to determine neutron-induced capture cross sections in the rare-earth region.

1 Introduction

Neutron-induced radiative-capture cross sections of short-lived nuclei are crucial for fundamental nuclear physics and also for applications such as reactor physics and astrophysics. In particular, these data are needed to test s- and r-process models. The latter are used to understand the synthesis of the elements between iron and uranium in astrophysical environments. However, very often the high radioactivity of the samples makes the direct measurement of these cross sections extremely difficult. The surrogate reaction method is an indirect way of determining cross sections for compound nuclear reactions. This method was first proposed by J.D.Cramer and H.C.Britt [1] in the seventies and is schematically represented in figure 1. The left part of figure 1 illustrates a neutron-induced reaction on target $A-1$, which leads to the compound-nucleus A at an excitation energy E^* . The nucleus A^* can decay through different exit channels: fission, gamma-decay, neutron emission, etc... On the right part of figure 1, in the surrogate reaction method, the same compound nucleus A^* is produced by a transfer reaction between a projectile y (a light charged particle)

and a target X . The transfer reaction ($y+X \rightarrow A+w$) leads to a heavy recoil nucleus A^* and an ejectile w . The identification of the ejectile permits to determine the mass A and charge Z of the decaying nucleus. In addition, we can deduce the excitation energy E^* of the compound nucleus A by measuring the kinetic energy and the emission angle of the ejectile w . The measurement of the number of coincidences between the ejectiles and the decay products normalised to the total number of detected ejectiles allows one to extract the decay probability $\overline{P}_{decay}^{A,exp}$ for the corresponding decay channel. According to the surrogate reaction method, the neutron-induced cross section for the nucleus $A-1$ is then given by the equation:

$$\sigma_{decay}^{A-1}(E_n) \cong \sigma_{CN}^A(E_n) \cdot P_{decay}^{A,exp}(E^*) \quad (1)$$

where σ_{CN}^A is the calculated compound nuclear formation cross section in the desired reaction (formation of the nucleus A after a neutron absorption with an energy

E_n). In our case, σ_{CN}^A is obtained from optical model calculations performed with the code TALYS [2]. The relation between incident neutron energy E_n and excitation energy E^* of the compound nucleus A can be written as:

$$E^* = S_n + E_n \cdot \frac{A-1}{A} \quad (2)$$

where S_n is the one-neutron separation energy in the nucleus A . The interest of this method is that in some cases the target X is stable or less radioactive than the target $A-1$.

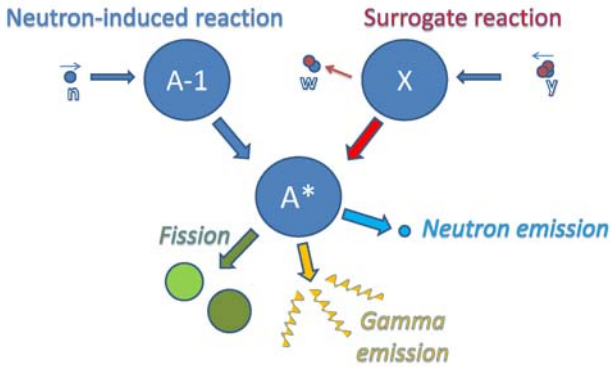


Fig. 1: Schematic representation of the surrogate reaction method. The surrogate reaction is here a transfer reaction $X(y,w)A^*$. Three possible exit channels (fission, gamma emission and neutron emission) are also represented.

Recently, we used the surrogate reaction method to determine the neutron-induced fission cross sections of ^{242}Cm ($T_{1/2}=162.8$ d), ^{243}Cm ($T_{1/2}=29.1$ y) and ^{241}Am ($T_{1/2}=432.2$ y). To reach and study these nuclei we employed few-nucleon transfer reactions using a ^3He projectile on a ^{243}Am ($T_{1/2}=7370$ y) target. All details are given in [3]. A remarkable good agreement was observed between our results and the existing neutron-induced data at the lowest neutron energies. In the present work, we investigate whether the surrogate reaction technique can be used to infer neutron-induced capture cross sections.

2 Validity of the surrogate reaction method

The neutron-induced reaction and the transfer reaction permit to produce the same compound nucleus in Z , A and E^* . However, the angular momentum (J) and parity (π) distributions populated by a transfer reaction may not be the same as the ones populated in a neutron-induced reaction. Since at low E^* the decay probabilities may strongly depend on J^π , the decay probability obtained in surrogate experiments can be very different from the one measured in neutron-induced experiments.

Assuming that the nucleus A^* is in a compound state, its formation and decay are independent and the decay probabilities are given by:

$$P_n^{decay}(E^*) = \sum_{J^\pi} P_n^{form}(E^*, J^\pi) \cdot G_{decay}(E^*, J^\pi) \quad (3)$$

$$P_t^{decay}(E^*) = \sum_{J^\pi} P_t^{form}(E^*, J^\pi) \cdot G_{decay}(E^*, J^\pi) \quad (4)$$

where the indices n and t stand for neutron and transfer reactions, respectively, $P_n^{form}(E^*, J^\pi)$ and $P_t^{form}(E^*, J^\pi)$ correspond to the probability that the compound nucleus is formed in the state J^π by the neutron-induced and the transfer reaction, respectively. $G_{decay}(E^*, J^\pi)$ is the branching ratio for a given decay channel. The two decay probabilities of eqs. (3) and (4) are equal in two cases:

1) The J^π distributions populated in both reactions are similar:

$$P_n^{form}(E^*, J^\pi) \approx P_t^{form}(E^*, J^\pi) \quad (5)$$

Unfortunately, the experimental and/or theoretical determination of the angular momentum populated in transfer reactions represents a big challenge. An important effort from theoreticians and experimentalists should be done in order to determine these distributions. We will see below how the present work can provide very valuable information on this issue.

2) The branching ratios are independent of J^π :

$$G_{decay}(E^*, J^\pi) = G_{decay}(E^*) \quad (6)$$

Then the branching ratios can be taken out of the summation signs in eqs. (3) and (4). Since

$$\sum_{J^\pi} P_n^{form}(E^*, J^\pi) = 1 \quad (7)$$

$P_n^{decay}(E^*, J^\pi) \approx P_t^{decay}(E^*, J^\pi)$ and the cross section for the desired reaction takes the simple product form of eq. (1). This second hypothesis is known as the Weisskopf-Ewing approximation [4] and is justified for high excitation energies where the decay of the compound-nucleus is dominated by statistical level densities. At lower excitation energies, the decay probabilities strongly depend of the J^π of discrete states, whose population depends on the reaction mechanism used to produce the compound nucleus A^* .

In Ref. [3] we showed that our results for the fission cross sections obtained with the surrogate method are in very good agreement with the neutron-induced data at low excitation energies. The reason is that for all the

reactions considered in [3] the excitation energy of the decaying nuclei was high enough for the Weisskopf-Ewing approximation to be valid. In radiative capture reactions, however, we have to consider the competition with neutron emission to the ground or first excited states which is much more sensitive to the differences between the spin-parity distributions [5]. In addition, in the case of actinides, one may need to distinguish between gamma rays originating from the fission fragments and radiative capture gamma rays. This can make radiative capture measurements extremely complicated. Therefore, as a first step we have chosen to investigate the validity of the surrogate method for radiative capture reactions on rare earth nuclei. In particular, our aim is to study the transfer reactions $^{174}\text{Yb}({}^3\text{He},p\gamma){}^{176}\text{Lu}$ and $^{174}\text{Yb}({}^3\text{He},{}^4\text{He}\gamma){}^{173}\text{Yb}$ as surrogate for the $^{175}\text{Lu}(n,\gamma)$ and $^{172}\text{Yb}(n,\gamma)$ reactions, respectively. We have considered the $^{175}\text{Lu}(n,\gamma)$ and $^{172}\text{Yb}(n,\gamma)$ cross sections because they present the advantage to be very well known, see for example [6, 7, 8, 9].

3 Experimental set-up

The measurement was performed at the Tandem accelerator of the IPN Orsay. We used an incident ${}^3\text{He}$ beam with an energy of 24 MeV. The beam intensity was 20 particle nA. The stable ^{174}Yb target was fabricated at the SIDONIE facility of the CSNSM laboratory. The ^{174}Yb sample had a thickness of $250\ \mu\text{g}/\text{cm}^2$ and was deposited onto a C foil of $50\ \mu\text{g}/\text{cm}^2$. Figure 2 illustrates our experimental set-up. To infer the radiative capture probability, gamma rays were detected in coincidence with the ejectiles. The latter were fully identified by two large area ΔE -E telescopes placed symmetrically at 130° with regard to the ${}^3\text{He}$ beam. The ΔE detectors were two $300\ \mu\text{m}$ silicon position sensitive detectors, whose $16*16$ X-Y strips provided the angle of the detected particle with an angular coverage of 108° to 152° . The E detectors were two Si(Li) detectors of 3 mm thickness. We shielded the Si detectors against delta electrons coming from the target with a thin Mylar(Al) foil polarized at -300V.

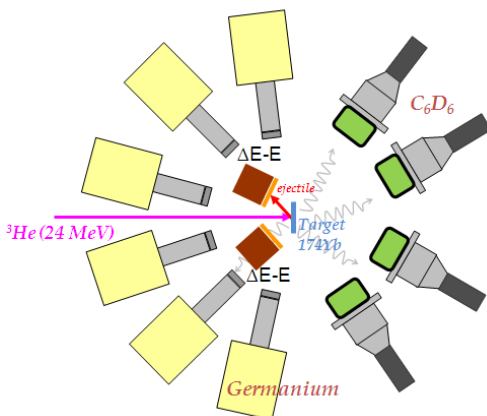


Fig. 2 : Top view of the experimental set-up for radiative capture probability measurements. The four C_6D_6 liquid scintillators were placed at forward angles with respect to the beam direction, whereas the two Si telescopes and the

six germanium detectors were placed at backward angles.

Four C_6D_6 liquid scintillators were used for gamma detection to infer the radiative capture probability. The use of C_6D_6 liquid scintillators has the important advantage that the coupling of these detectors to a pulse shape discriminator permits to separate between photons and neutrons interacting within the scintillators. The ^{174}Yb target was also surrounded by six high-volume germanium detectors. They were used to measure low-lying γ -ray transition intensities as a function of the compound nucleus excitation energy, which is an additional way to investigate the radiative capture probability and the difference in spin distributions between transfer and neutron-induced reactions.

The ${}^3\text{He}$ -induced transfer reactions on the ^{174}Yb target lead to the production of various heavy residues. In this work we consider only the $({}^3\text{He},p)$ and the $({}^3\text{He},{}^4\text{He})$ channels because they lead to nuclei for which there exist neutron-induced data to compare with. The advantage of using transfer reactions is clear: the simultaneous access to two transfer channels allowed us to study two nuclei from a single projectile-target combination. Moreover, since there are two bodies in the outgoing reaction channel, the excitation energy of the decaying nucleus E^* follows a broad probability distribution.

4 Results

The experimental radiative capture probability $P_\gamma(E^*)$ can be obtained in the following way:

$$P_\gamma(E^*) = \frac{N_{\text{coinc}}(E^*)}{N_{\text{single}}(E^*)} \cdot \frac{1}{\varepsilon(E^*)} \quad (8)$$

where $N_{\text{coinc}}(E^*)$ is the number of ejectiles detected in coincidence with the C_6D_6 detectors, $N_{\text{single}}(E^*)$ the total number of ejectiles, i.e. the total number of decaying nuclei formed, and $\varepsilon(E^*)$ represents the C_6D_6 cascade detection efficiency.

4.1 Preliminary results for the $^{174}\text{Yb}({}^3\text{He},p){}^{176}\text{Lu}$ reaction

In this transfer channel the $N_{\text{single}}(E^*)$ and $N_{\text{coinc}}(E^*)$ spectra have to be corrected for the ejectiles coming from transfer reactions between the ${}^3\text{He}$ beam and the carbon backing. Since the ^{176}Lu is formed by a transfer reaction, it is possible to extend our investigation below the neutron separation energy S_n where only γ -rays can be emitted and consequently the measured radiative capture probability should be 1. The probability to detect with our set-up various gamma rays coming from the same cascade is less than 3%. Therefore, below S_n the ratio $N_{\text{coinc}}(E^*)/N_{\text{single}}(E^*)$ gives the total efficiency of the C_6D_6 detectors for detecting a gamma cascade. Applying a 500 keV threshold to the gamma-energy in order to suppress the contribution of the $({}^3\text{He}, p n \gamma){}^{175}\text{Lu}^*$ channel, this ratio is of around 6.5% ($\pm 0.3\%$) and remains essentially constant from $E^*=5.5$ MeV to S_n , see figure 3.

The constancy of this ratio is to be expected as TALYS calculations show that the multiplicity of the gamma cascade and the average gamma energy vary only very weakly between S_n and 8 MeV excitation energy. Thus, it is reasonable to assume this value for the efficiency also above S_n . The independence of the cascade detection efficiency with the excitation energy and its absolute value have been confirmed by using the total-energy detection principle in combination with the pulse-height weighting technique. This technique allows one to extract the cascade detection efficiency independently from the measured data. It was used in previous surrogate studies using the $^{232}\text{Th}(^3\text{He},p)$ reaction [10, 11].

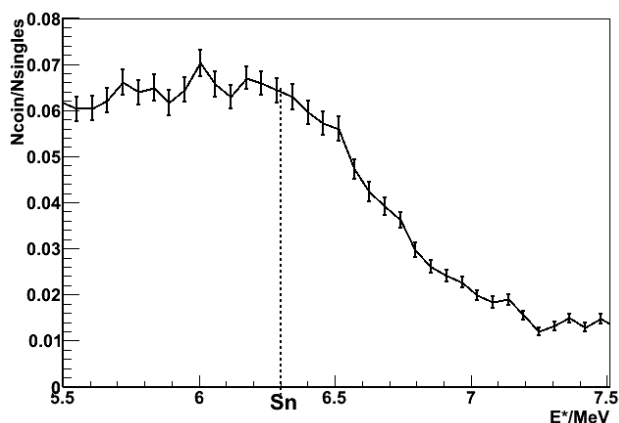


Fig. 3: Ratio $N_{\text{coinc}}/N_{\text{singles}}$ as a function of excitation energy for the $^{174}\text{Yb}(^3\text{He}, p)^{176}\text{Lu}$ reaction.

Once the detection efficiency is determined, we apply equation (8) to determine the radiative capture probability of ^{176}Lu as a function of E^* , see figure 4. Our preliminary results are compared with TALYS calculations for the neutron-induced capture probability of ^{175}Lu and for the gamma-induced capture reaction of ^{176}Lu . The parameters of the TALYS code have been tuned to be in excellent agreement with the experimental data for the $^{175}\text{Lu}(n,\gamma)$ cross sections. The results of figure 4 show that our surrogate data present big discrepancies with respect to the neutron-induced data, while we observe a very good agreement with the $^{176}\text{Lu}(\gamma,\gamma)$ calculation at low energies. This indicates that the J^π distribution populated in the $(^3\text{He},p)$ surrogate reaction is close to the one populated in the photon-induced reaction. The ground state J^π of ^{175}Lu and ^{176}Lu are respectively $7/2^+$ and 7^- . Therefore, the angular momentum of ^{176}Lu populated by low energy neutrons is mainly centered between 3 and 4 \hbar , while the angular momentum populated in the gamma-induced reaction is centered around 6,7 and 8 \hbar . Consequently, the big discrepancies found at low E^* can be explained by the differences between the spin distributions populated in transfer and neutron-induced reactions. For excitation energies after neutron emission below the first excited state of ^{175}Lu (113keV, $9/2^+$), the (n,γ) decay channel is only in competition with the compound-elastic channel (n,n) , where the residual nucleus is left in its ground state after neutron emission. Since the nucleus can only decay to one state with a well-defined J^π ($7/2^+$) and the angular

momentum carried by the emitted neutron is very small, this particular exit channel is extremely sensitive to the spin of the compound nucleus $^{176}\text{Lu}^*$. TALYS calculations show that the (n,n) channel remains rather strong up to about 7MeV. As said above, the angular momentum induced by the $(^3\text{He},p)$ transfer reaction seems to be also centered around 7 or 8 \hbar , which is about two times higher than the angular momentum of the ^{175}Lu ground state $7/2$, leading to a suppression of the compound-elastic channel. In conclusion, the high selectivity of the (n,n) decay channel is most probably at the origin of the large discrepancies observed between surrogate and neutron-induced measurements. Since the gamma decay probability of ^{176}Lu is very sensitive to J^π , in the future we plan to extract the populated J^π from a fit to the experimental decay probability using the branching ratios (eqs. 3 and 4) calculated by TALYS. Note that since the J^π of the ground state of the target used is 0^+ and we are considering high E^* , this fit provides direct information on the angular momentum transferred in a $(^3\text{He},p)$ reaction. High discrepancies have been also observed for ^{155}Gd and ^{157}Gd radiative capture cross sections obtained via the surrogate method involving (p,p') inelastic reactions [12]. They have also been attributed to the J^π mismatch.

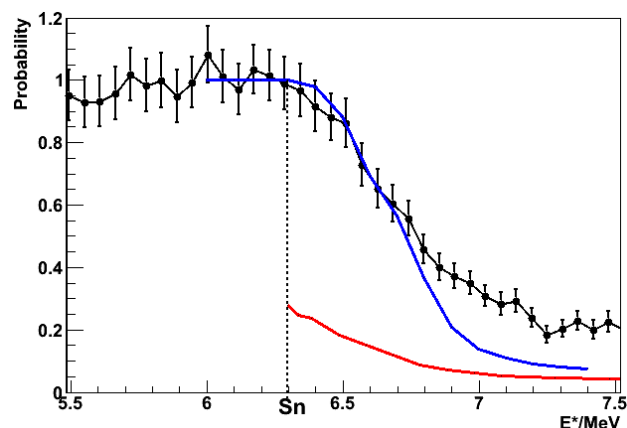


Fig. 4: Preliminary results for the radiative capture probability measured in the $^{174}\text{Yb}(^3\text{He}, p)^{176}\text{Lu}$ reaction. The red line is the result of a calculation performed with TALYS for the neutron-induced capture probability of ^{176}Lu . The blue line is TALYS result for the photon-induced capture probability of ^{176}Lu .

Our previous conclusion is reinforced by the measurement of low-lying γ -ray transition intensities performed with the germanium detectors. Figure 5 shows the Germanium gamma-ray spectrum in coincidence with protons for various excitation energies. The gamma-ray transitions coming from de-excitation of the 8^+ level of ^{176}Lu , located at 424.9 keV, are clearly observed at 424.9 keV and 241 keV. In the $^{175}\text{Lu}(n,\gamma)$ reaction at thermal neutron energies [13] these gamma-ray transitions are observed with an intensity 10 times lower than the 139.3 keV transition coming from the 4^+ level ($E^*=372.5$ keV), which is not observed in our surrogate data. For 1 MeV neutron energy, the gamma-ray intensity ratio (139.3

keV/424.9 keV) is predicted to be around 4 in our (n, γ) TALYS calculation. The comparison of the three spectra shown in figure 5 reflects very clearly the contamination due to the decay of ^{175}Lu produced in the reaction ($^3\text{He},\text{pn}\gamma$). The bottom panel of figure 5 shows that above $E^*=6.9$ MeV the decay is dominated by this channel. As mentioned before, a threshold of 500 keV in the gamma energy measured by the C_6D_6 is set for all excitation energies to remove the contribution of the ($^3\text{He},\text{pn}\gamma$) reaction. This value of the threshold comes from the fact that for ^{175}Lu all the gamma-rays energies that have been measured between $E^*=0$ and 1.5 MeV are below 500 keV.

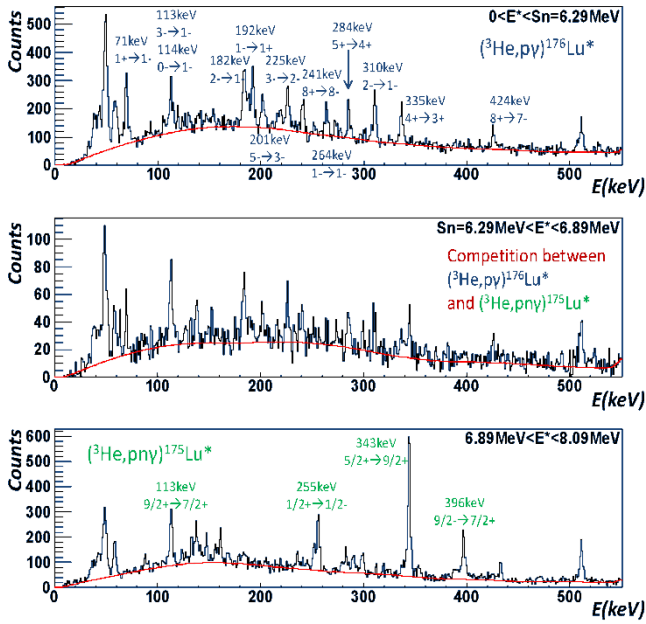


Fig. 5 : Germanium gamma-ray spectra for three ranges of E^* . One can identify the main γ -ray transitions related to the ($^3\text{He},\text{py}$) $^{176}\text{Lu}^*$ and ($^3\text{He},\text{pn}\gamma$) $^{175}\text{Lu}^*$ reactions. The background is shown in red.

4.2 Preliminary results for the $^{174}\text{Yb}(^3\text{He}, ^4\text{He})^{173}\text{Yb}$ reaction

The same analysis as the one presented in the previous section was performed to extract the radiative capture probability associated to the $^{174}\text{Yb}(^3\text{He}, ^4\text{He})^{173}\text{Yb}$ reaction. In this case a gamma-energy threshold of 400 keV for the C_6D_6 detectors was necessary to remove the contribution from the $^{174}\text{Yb}(^3\text{He}, ^4\text{He})\gamma$ reaction. The preliminary results for the gamma decay probability are shown in figure 6. The data are compared with TALYS results for the neutron and photon-induced radiative capture probabilities. The parameters of the code have been fixed to best reproduce the existing neutron-induced data. Again, the transfer-induced results are much higher (by a factor 10 at the lowest energies!) than the neutron-induced data. In this case, the photon-induced capture probability is also below the transfer-induced data. The two TALYS calculations show clear changes of slope at S_n and at energies that correspond to the first and second excited state of ^{172}Yb . They indicate the reduction of the gamma decay probability caused by the competition with

neutron emission leaving the residual nucleus ^{172}Yb in the ground state, in the first or the second excited state. These changes in slope can also be observed at similar energies for the $^{174}\text{Yb}(^3\text{He}, ^4\text{He})^{173}\text{Yb}$ reaction although in this case the changes due to higher states are also observed. Note that the excitation-energy resolution for these data is 85 keV. It is interesting to see that for the transfer reaction the decay probability remains almost equal to 1 until the E^* of the first excited state is reached, indicating that the emission of neutrons to the ground state of ^{172}Yb is forbidden. Also the drops in the radiative capture probability are less intense than for the (n, γ) and (γ , γ) reactions. This suggests that the angular momentum populated in the transfer reaction is significantly bigger than the one populated in the neutron and photon-induced reactions. The ground state of ^{173}Yb is $5/2^-$, therefore the angular momentum of ^{173}Yb after photon absorption will be centered around $3/2$ and $7/2$. These values are smaller than the ones populated for ^{176}Lu whose ground-state spin is 7^+ . This explains why in this case the photon-induced reaction is clearly below the transfer reaction. As for the ($^3\text{He}, \text{p}$) reaction, we will extract rather direct information on the transferred angular momentum in the ($^3\text{He}, ^4\text{He}$) reaction from a fit to the experimental radiative capture probability of figure 6 using eqs. (3-4) and the branching ratios from TALYS.

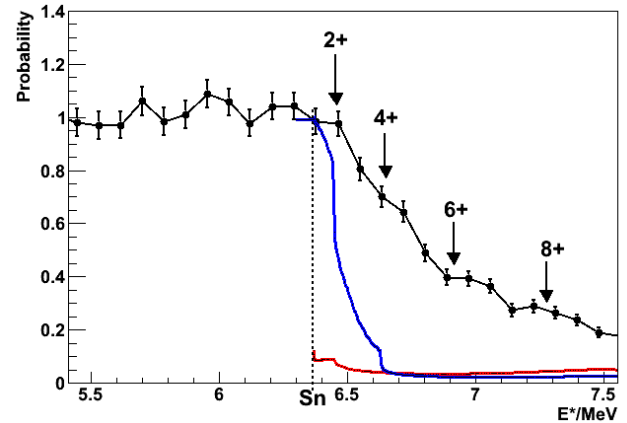


Fig. 6: Preliminary results for the radiative capture probability measured in the $^{174}\text{Yb}(^3\text{He}, ^4\text{He})^{173}\text{Yb}$ reaction. The red line is the result of a calculation performed with TALYS for the neutron-induced capture probability of ^{173}Yb . The blue line is TALYS result for the photon-induced capture probability of ^{173}Yb . The arrows indicate the position of the first excited states of ^{172}Yb .

5. Conclusion

We have performed an experiment to study the validity of the surrogate method for extracting neutron-induced capture cross sections. We have used the well known $^{175}\text{Lu}(n,\gamma)$ and $^{172}\text{Yb}(n,\gamma)$ cross sections to study the $^{174}\text{Yb}(^3\text{He},\text{p})^{176}\text{Lu}^*$ and $^{174}\text{Yb}(^3\text{He},^4\text{He})^{173}\text{Yb}$ surrogate reactions. Our experimental results indicate that the angular momentum populated in the transfer reactions used is significantly higher than the one populated in neutron-induced reactions. These differences explain the

big discrepancies observed between the surrogate capture measurements and the neutron-induced data. At low excitation energy, the compound elastic (n,n) decay channel is predominant and has the particularity to be extremely sensitive to the spin and the parity of the decaying nucleus. Our experimental data clearly reflect that this decay channel is not accessible in the transfer reactions we have considered. This study is extremely important in view of the application of the surrogate method to infer capture cross sections of actinides. In the future, we plan to use the $^{238}\text{U}(n,\gamma)$ and fission cross sections to investigate $^{238}\text{U}(d,p)^{239}\text{U}$ surrogate reaction. Note that the (d,p) reaction is very important for future experiments in inverse kinematics.

Acknowledgments

We would like to express our gratitude to M.-G. Porquet and the SIDONIE facility of the CSNSM for providing the ^{174}Yb targets. This work is supported by the CNRS program PACEN/GEDEPEON and the EURATOM programs EFNUDAT (European Facilities for Nuclear Data Measurements) contract n° FP6-036434 and ANDES (Accurate Nuclear Data for nuclear Energy Sustainability) contract n° FP7-249671.

References

1. J.D. Cramer, H.C. Britt, Nucl. Sci. Eng. **41**, 177 (1970)
2. A. J. Koning, S. Hilaire et al., User Manual TALYS, www.talys.eu
3. G. Kessedjian et al., Phys. Lett. B **692**, 297 (2010)
4. V. Weisskopf and D. H. Ewing, Phys. Rev. **57**, 472 (1940)
5. J.E. Escher and F.S. Dietrich, Phys. Rev. C **81**, 024612 (2010)
6. R.L. Macklin et al., Nucl Sci. Eng. **95**, 189 (1987)
7. K. Wisshak et al., Phys. Rev. C **73**, 015807 (2006)
8. M.V. Borkhovko et al., Vop. At. Nauki i Tekhn., Ser. Yadernye Konstanty, 44 (1985)
9. K. Wisshak et al., Phys. Rev. C **61**, 065801 (2000)
10. S. Boyer et al., Nucl. Phys. A **775**, 175 (2006)
11. J. N. Wilson et al., Nucl. Instr. Meth. A **511**, 388 (2003)
12. N.D. Scielzo et al., Phys. Rev. C **81**, 034608 (2010)
13. W. Andrejtscheff, Nucl. Phys. A **226**, 142 (1974)