

Half-lives of neutron-rich $^{128-130}\text{Cd}$

R. Dunlop,^{1,*} V. Bildstein,¹ I. Dillmann,^{2,3,†} A. Jungclaus,⁴ C. E. Svensson,¹ C. Andreoiu,⁵ G. C. Ball,² N. Bernier,^{2,6} H. Bidaman,¹ P. Boubel,¹ C. Burbadge,¹ R. Caballero-Folch,² M. R. Dunlop,¹ L. J. Evitts,^{2,7} F. Garcia,⁵ A. B. Garnsworthy,² P. E. Garrett,¹ G. Hackman,² S. Hallam,^{2,7} J. Henderson,² S. Ilyushkin,⁸ D. Kisliuk,¹ R. Krücken,^{2,6} J. Lassen,^{2,9} R. Li,² E. MacConnachie,² A. D. MacLean,¹ E. McGee,¹ M. Moukaddam,² B. Olaizola,¹ E. Padilla-Rodal,¹⁰ J. Park,^{2,6} O. Paetkau,² C. M. Petrache,¹¹ J. L. Pore,⁵ A. J. Radich,¹ P. Ruotsalainen,² J. Smallcombe,² J. K. Smith,² S. L. Tabor,¹² A. Teigelhöfer,^{2,9} J. Turko,¹ and T. Zidar¹

¹*Department of Physics, University of Guelph, Guelph, Ontario N1G 2W1, Canada*

²*TRIUMF, 4004 Wesbrook Mall, Vancouver, British Columbia V6T 2A3, Canada*

³*Department of Physics and Astronomy, University of Victoria, Victoria, British Columbia V8P 5C2, Canada*

⁴*Instituto de Estructura de la Materia, CSIC, E-28006 Madrid, Spain*

⁵*Department of Chemistry, Simon Fraser University, Burnaby, British Columbia V5A 1S6, Canada*

⁶*Department of Physics and Astronomy, University of British Columbia, Vancouver, British Columbia V6T 1Z4, Canada*

⁷*Department of Physics, University of Surrey, Guildford GU2 7XH, United Kingdom*

⁸*Department of Physics, Colorado School of Mines, Golden, Colorado 80401, USA*

⁹*Department of Physics and Astronomy, University of Manitoba, Winnipeg, Manitoba R3T 2N2, Canada*

¹⁰*Universidad Nacional Autónoma de México, Instituto de Ciencias Nucleares, AP 70-543, México City 04510, DF, México*

¹¹*Centre de Sciences Nucléaires et Sciences de la Matière, CNRS/IN2P3, Université Paris-Saclay, 91405 Orsay, France*

¹²*Department of Physics, Florida State University, Tallahassee, Florida 32306, USA*

(Received 29 March 2016; revised manuscript received 16 May 2016; published 16 June 2016)

The β -decay half-lives of $^{128-130}\text{Cd}$ have been measured with the newly commissioned GRIFFIN γ -ray spectrometer at the TRIUMF-ISAC facility. The time structures of the most intense γ rays emitted following the β decay were used to determine the half-lives of ^{128}Cd and ^{130}Cd to be $T_{1/2} = 246.2(21)$ ms and $T_{1/2} = 126(4)$ ms, respectively. The half-lives of the $3/2^+$ and $11/2^-$ states of ^{129}Cd were measured to be $T_{1/2}(3/2^+) = 157(8)$ ms and $T_{1/2}(11/2^-) = 147(3)$ ms. The half-lives of the Cd isotopes around the $N = 82$ shell closure are an important ingredient in astrophysical simulations to derive the magnitude of the second r -process abundance peak in the $A \sim 130$ region. Our new results are compared with recent literature values and theoretical calculations.

DOI: [10.1103/PhysRevC.93.062801](https://doi.org/10.1103/PhysRevC.93.062801)

I. INTRODUCTION

The β -decay properties (half-lives and β -delayed neutron-branching ratios) of nuclei below doubly magic $^{132}\text{Sn}_{82}$ (i.e., $N \approx 82$, $Z < 50$) are key input parameters for any astrophysical r -process scenario because they play an important role in the formation and shape of the second abundance peak at $A \sim 130$ [1]. This is despite the fact that the astrophysical site, or sites, where rapid neutron capture nucleosynthesis [2–6] takes place remain(s) elusive.

In both the high- and low-entropy hot neutrino-driven-wind scenarios, the most important nuclei in this mass region are the $N = 82$ isotones with $Z = 40$ –50 because the enhanced neutron binding energy compared with their isotopic neighbors leads to a barrier for the r -process reaction flow toward heavier masses. After the breakout of the $N = 82$ shell, isotopes with $N = 84$, 86, and 88 also become important, such as $^{134,136,138}\text{Sn}$, $^{133,135}\text{Ag}$, $^{132,134,136}\text{Cd}$, $^{131,133}\text{Rh}$, and ^{130}Pd [1].

At the so-called “waiting-point nuclei,” an accumulation of r -process material occurs (under given astrophysical conditions) and material can be transferred to the next elemental chain via β decay. The half-lives of these waiting points thus determine how much material is accumulated and, therefore,

the amplitude and shape of the resulting r -process abundance peaks after decay back to stability. The prominent r -process abundance peaks at $A \sim 80$, 130, and 195 correspond to waiting-point isotopes at the closed neutron shells $N = 50$, 82, and 126 where, due to nuclear-shell-structure effects, the reaction flow is hindered.

More neutron-rich “cold” r -process scenarios, such as neutron-star mergers [7,8], drive the reaction path towards the neutron dripline into regions that will only be partially accessible to experiments at the new generation of radioactive-beam facilities. Since most nuclei involved in r -process calculations are currently experimentally inaccessible, one has to rely heavily on the predictive power of theoretical models for the β decay of these nuclei. The relative r -process abundances of nuclei around neutron shell closures are particularly sensitive to the half-lives, and it is thus critical to have models that can accurately reproduce these decay properties.

In particular, shell-model calculations for the waiting-point nuclei near the $N = 82$ neutron shell closure [9,10] have been performed by adjusting the quenching of the Gamow–Teller (GT) operator to reproduce the ^{130}Cd half-life reported in Ref. [11] and are known to yield systematically large values for the half-lives of other nuclei in the region [10]. A new, shorter, half-life for ^{130}Cd as measured by the EURICA collaboration [12] would resolve this discrepancy by scaling the GT quenching by a constant factor for all nuclei in this region.

*rdunlop@uoguelph.ca

†dillmann@triumf.ca

Distinguishing between these discrepant half-life measurements for ^{130}Cd [11,12] is thus of critical importance since the as-yet-unknown half-lives of other $N = 82$ waiting-point nuclei with $40 \leq Z \leq 44$ play a key role for the reproduction of the second abundance peak in r -process calculations.

A recent experimental campaign with the EURICA detector [13] at RIKEN measured the β -decay half-lives of 110 neutron-rich isotopes between Rb and Sn; among them, $^{128-130}\text{Cd}$ [12]. While the previously reported half-life value for ^{128}Cd of $T_{1/2} = 280(40)$ ms [14] was in agreement with the much more precise value of $T_{1/2} = 245(5)$ ms reported in Ref. [12], large discrepancies were found for $^{129,130}\text{Cd}$.

^{129}Cd β decays from both a $3/2^+$ and an $11/2^-$ state but it is presently unknown which of the two states is the ground state and which is the isomeric state. In an experiment at ISOLDE, Arndt *et al.* [15,16] measured the half-lives of both states by using β -delayed neutrons. They reported $T_{1/2} = 104(6)$ ms for the “ $11/2^-$ ground state” and $T_{1/2} = 242(8)$ ms for the “ $3/2^+$ isomer” but gave no explanation of how the ground state was assigned.

The ground-state-spin assignments for many neutron-rich odd- A Cd isotopes have recently been confirmed via laser spectroscopy [17]. The odd- A Cd isotopes $^{121,123,125,127}\text{Cd}$ have a well-established ground-state spin of $3/2^+$ but the exact position of the $11/2^-$ isomer is not known for $^{125,127}\text{Cd}$. Shell-model calculations [18] suggest that this order is inverted at ^{129}Cd compared with the lighter odd- A Cd isotopes; however, there is no direct experimental evidence for this inversion. We thus label the two states only according to their spin and parity.

The recent measurements of the EURICA collaboration [18] did not resolve the issue of the ground-state spin-parity assignment in ^{129}Cd . However, the half-lives for the $3/2^+$ and $11/2^-$ states were determined separately via the γ transitions at 1423 and 1586 keV in the daughter nucleus to be $T_{1/2}(3/2^+) = 146(8)$ ms, and via the γ -ray transitions at 359, 995, 1354, 1796, and 2156 keV to be $T_{1/2}(11/2^-) = 155(3)$ ms. These results are in clear contradiction with the previous measurements [15,16].

In the case of the ^{130}Cd half-life, the value of 127(2) ms reported in Ref. [12] also differed from the previously accepted value of 162(7) ms [11] by more than 5σ . The measurement of Ref. [11] was performed with the same technique using β -delayed neutrons as the ^{129}Cd measurements of Refs. [15,16]. In an earlier paper, the ^{130}Cd half-life was reported as 195(35) ms [19].

In this paper we report an independent determination of the half-lives of $^{128-130}\text{Cd}$ which, in general, confirm the recent EURICA results [12,18,20] but disagree with the previous measurements [11,15,16]. We report an improved precision for the ^{128}Cd half-life, and revised half-lives for the two β -decaying states of ^{129}Cd based on more detailed γ -ray spectroscopy.

II. EXPERIMENT

The half-lives of $^{128-130}\text{Cd}$ were measured with the newly commissioned GRIFFIN γ -ray spectrometer [21,22] at the TRIUMF-ISAC facility [23]. Many of the nuclei in this neutron-rich region below doubly magic ^{132}Sn have complicated decay chains, including significant β -delayed neutron-

emission branches, as well as the presence of β -decaying isomeric states. A measurement of the temporal distribution of characteristic γ rays emitted from the excited states of the daughter nucleus following β decay of the parent isotope is a powerful method to reduce the complex background contributions to the measurement. This method requires the use of a high-efficiency γ -ray spectrometer because of the low production rates and short half-lives.

The isotopes of interest were produced by using a 500 MeV proton beam with $9.8 \mu\text{A}$ intensity from the TRIUMF main cyclotron incident on a UC_x target. The ion-guide laser ion source (IG-LIS) [24] was used to suppress surface-ionized isobars such as In and Cs, while the neutral Cd atoms of interest were extracted and selectively laser ionized in a three-step-excitation scheme. The Cd isotopes of interest were then accelerated to 28 keV, selected by a high-resolution mass separator, and delivered to the GRIFFIN spectrometer. GRIFFIN is comprised of 16 high-purity germanium (HPGe) clover detectors [21,22]. The radioactive-ion beam (RIB) was implanted into an aluminized mylar tape of the moving tape collector at the mutual centers of SCEPTAR, an array of 20 thin plastic scintillators for tagging β particles [25], and GRIFFIN. The longer-lived background activity, either from isobaric contaminants in the beam or from daughters following the decay of the Cd isotopes, could be removed by moving the tape following a measurement. A typical cycle for the $^{128-130}\text{Cd}$ runs consisted of a background measurement for 0.5 s, followed by a collection period (beam-on) with the beam being implanted into the tape for 10 s, followed by a collection period (beam-off) with the beam blocked by the ISAC electrostatic beam kicker downstream of the mass separator. The beam-off period consisted of a decay time of typically two to three half-lives, the movement of the tape for 1 s to a shielded position outside of the array, and the start of the new cycle with the background measurement.

The high efficiency of the GRIFFIN array coupled with the SCEPTAR β detector allowed for the sensitive detection of the γ rays following the β decay of interest. All of the analyses reported here were performed by using add-back algorithms in which all of the detected energy in a clover within a 400 ns coincidence timing window was summed in order to increase the photopeak efficiency of GRIFFIN, as well as reduce the contribution of Compton background to the γ -ray spectrum [21,22].

III. DATA ANALYSIS

The data were analyzed by using β - γ coincidences requiring a β particle to be detected in SCEPTAR within a coincidence window of 400 ns of a γ ray detected in GRIFFIN, resulting in a strong suppression of room-background γ rays. Cycles in which the RIB dropped out for a portion of the cycle were rejected in order to increase the signal-to-background ratio. In the case of ^{128}Cd , this resulted in the removal of 30% of the cycles, but only 1% of the total data.

A. ^{128}Cd decay

Approximately 7 h of ^{128}Cd data were collected with a beam intensity of ~ 1000 pps. The 857 keV [$I_{\gamma,\text{rel}} = 95(10)\%$] and

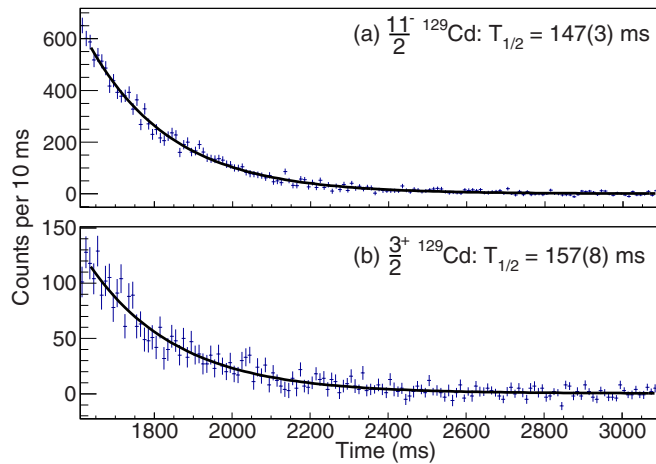


FIG. 4. (a) Fitted activity of the $11/2^-$ state using the 358.9, 1796.5, and 2155.6 keV γ rays. (b) Fitted activity of the $3/2^+$ state using the 1422.9 and 1586.2 keV γ rays. Note that the time represents the amount of time that has elapsed since the start of a cycle.

found to have negligible influence compared with the statistical uncertainties.

Unlike Ref. [18], in this work we do not use the strong 995 keV γ ray in the analysis of the half-life of the $11/2^-$ state of ^{129}Cd due to the previously observed feeding from the $3/2^+$ state of ^{129}Cd via the 1222 keV transition in ^{129}In [18]. We also do not include the 1354 keV γ ray in this analysis because it is contaminated by a γ ray of the same energy from the decay of the 611 (5) ms ground state of ^{129}In [30]. Based on the relative intensities of the observed γ rays from ^{129}In and ^{129}Sn , we estimate that approximately 20% of the total 1354 keV photopeak intensity in our experiment was from ^{129}In decay, which, if included, would bias the measured half-life to longer values.

TABLE I. Half-life of ^{129}Cd deduced from individual γ rays in this work and the corresponding results from the measurements of Ref. [20] (see text for details).

E_γ (keV)	Half-life (ms)		Parent state J^π
	This work	Ref. [20]	
358.9	148(3)	155.8(42)	$11/2^-$
1796.5	143(6)	157.9(99)	$11/2^-$
2155.6	136(12)	144(24)	$11/2^-$
Weighted average	146.5(26)	155.8(38)	
Summed value	147(3)	155(3) ^a	
1422.9	158(8)		$3/2^+$
1586.2	157(19)		$3/2^+$
Weighted average	157.9(75)		
Summed value	157(8)	146(8)	
995.1 ^b	151.8(23)	152.0(27)	$11/2^- + 3/2^+$
1354.2 ^c	192(12)	158.6(81)	Doublet

^aThe value published in Ref. [20] includes the transitions at 995 and 1354 keV.

^b γ ray was not used in the current analysis due to a potential contamination between the ^{129}Cd and $^{129}\text{Cd}^m$ decays.

^cDoublet with a γ ray from the decay of ^{129}In .

The half-lives for the $11/2^-$ and $3/2^+$ β -decaying states of ^{129}Cd measured in this work agree with the general conclusion of Ref. [18] that the half-lives of the two states are very similar and do not differ by a factor of ≈ 2 , as reported in Ref. [15,16,31]. A direct comparison of the results from the individual γ -ray transitions between this work and those of Ref. [20] is given in Table I. For statistical reasons, in Ref. [20] the counts in the 1423 and 1586 keV photopeaks were summed, and the fit of the summed decay curve resulted in the published value of 146 (8) ms [18] for the $3/2^+$ state which is consistent with the value of 157 (8) ms reported here. The weighted average of these two independent measurements is 151.5 (57) ms. For the half-life of the $11/2^-$ state we do not average with Ref. [18] but recommend the value of 147 (3) ms reported here due to the exclusion of contaminant γ -ray photopeaks in the current work.

C. ^{130}Cd decay

For ^{130}Cd , approximately 38 h of data were collected with a beam intensity of 15–30 pps. Figure 5 shows a portion of the β -coincident γ -ray spectrum obtained during the ^{130}Cd experiment.

The 451.0 [$I_{\gamma,\text{rel}} = 88.6(36)\%$], 1170.3 [$I_{\gamma,\text{rel}} = 20.0(2)\%$], and 1669.2 keV ($I_{\gamma,\text{rel}} = 100\%$) γ rays following the decay of ^{130}Cd [32] were used to measure the half-life, yielding 123 (5), 138 (20), and 126 (6) ms, respectively. The transition at 951 keV [$I_{\gamma,\text{rel}} = 22.1(33)\%$] [32] was not used because it is a doublet with a γ ray from the decay of ^{130}In [33]. Fitting the sum of the time distributions of these three γ rays yields a half-life of 126 (4) ms for the decay of ^{130}Cd (Fig. 6), in excellent agreement with the value of 127 (2) ms recently reported in Ref. [12] and in strong disagreement with the previous half-life measurement of 162 (7) ms [11]. The study of systematic uncertainties was performed as discussed above and did not reveal any statistically significant effects on the measured half-life.

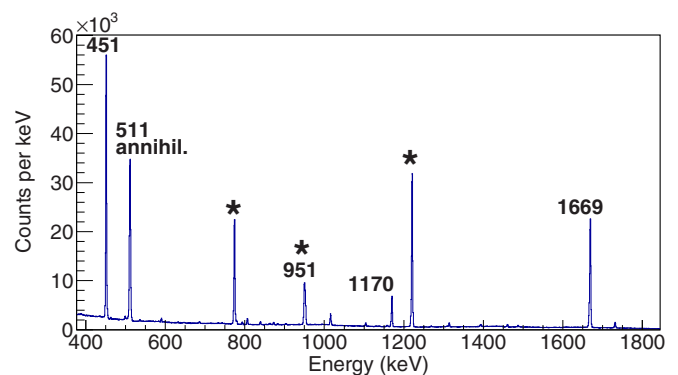


FIG. 5. A portion of the β -gated γ -ray energy spectrum for the ^{130}Cd experiment. The strongest peaks in the spectrum are labeled, including the doublet at 951 keV. The three strong γ rays at 451, 1170, and 1669 keV were used for the half-life analysis. γ rays following the β decay of ^{130}In are labeled with *.

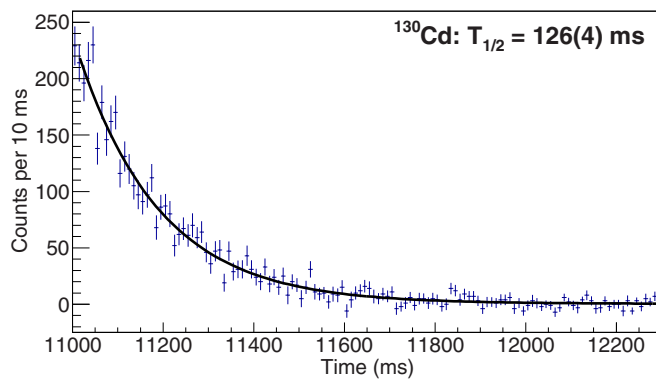


FIG. 6. Sum of the 451, 1170, and 1669 keV γ -ray time distributions. The half-life obtained from the fit is 126 (4) ms. Note that the time represents the amount of time that has elapsed since the start of a cycle.

IV. DISCUSSION AND CONCLUSION

The half-lives of ^{128}Cd , of the $11/2^-$ and the $3/2^+$ states of ^{129}Cd , and of the $N = 82$ isotope ^{130}Cd were measured at TRIUMF-ISAC by using the GRIFFIN γ -ray spectrometer. The ^{128}Cd half-life measured in this work of 246.2 (21) ms is in excellent agreement with the previous measurement of Ref. [12], but a factor of 2.4 more precise. The measured half-lives of the two known β -decaying states in ^{129}Cd , 147 (3) ms for the $11/2^-$ state and 157 (8) ms for the $3/2^+$ state, are found to be similar, in agreement with the recent work of Ref. [18], but in disagreement with the results of Refs. [15,16,31]. We recommend the revised value for the $11/2^-$ state reported here rather than averaging with Ref. [18] due to the exclusion of potential contaminants in the current analysis. Finally, the half-life of the $N = 82$ waiting point nucleus ^{130}Cd was measured to be 126 (4) ms, in excellent agreement with the value of 127 (2) ms reported in Ref. [12] but in strong disagreement with the measurements of 162 (7) and 195 (35) ms from Refs. [11,19].

The confirmation of the shorter half-life for the $N = 82$ isotope ^{130}Cd has significant implications for nuclear structure calculations in this region, as well as for r -process nucleosynthesis simulations. As shown in Fig. 7, the Fayans energy-density functional (DF3) + continuum quasiparticle random-phase approximation (CQRPA) model [34] and the relativistic Hartree–Bogoliubov (RHB) + relativistic quasiparticle random-phase approximation (RQRPA) [35] in general do a reasonable job of describing the systematic trend of the half-lives of neutron-rich Cd isotopes but slightly overestimate the absolute values.

As discussed in Ref. [12], the systematic overestimate of the half-lives for the $N = 82$ isotones can be traced to the scaling of the Gamow–Teller quenching to the previously reported

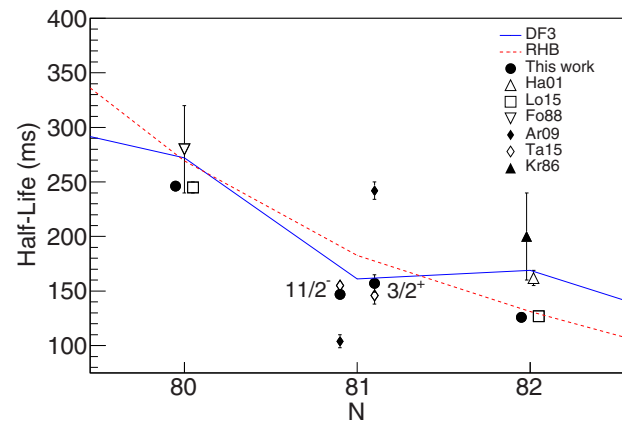


FIG. 7. Comparison between the calculated half-lives using the DF3 + CQRPA [34] and RHB + RQRPA [35] models to the measured half-lives of ^{128}Cd , ^{129}Cd , and ^{130}Cd . The measured half-lives in this work are in good agreement with Ref. [12,18] but do not agree with the previous measurements of Refs. [11,14–16,31]. Some error bars are not visible in the plot because they are smaller than the data points. Ha01: [11], Lo15: [12], Fo88: [14], Ar09: [16], Ta15 [18], Kr86 [19].

longer half-life for ^{130}Cd [11]. Increasing the GT quenching factor from $q = 0.66$ to $q = 0.75$ in order to reproduce the shorter half-life of ^{130}Cd reported in Ref. [12] and confirmed in the current work resolves this discrepancy. This directly affects the predicted half-lives for the yet-unmeasured $N = 82$ isotones ^{127}Rh , ^{126}Ru , and ^{125}Tc . As demonstrated in Refs. [1,12], the decrease in the calculated half-lives for these nuclei has a major influence on the shape of the rising wing of the r -process abundance peak at $A \sim 130$.

ACKNOWLEDGMENTS

This work has been partially supported by the Natural Sciences and Engineering Research Council of Canada (NSERC) and the Canada Research Chairs Program. I.D. and R.C.-F. are supported by NSERC Discovery Grants SAPIN-2014-00028 and RGPAS 462257-2014. A.J. acknowledges financial support by the Spanish Ministerio de Ciencia e Innovación under contract FPA2011-29854-C04 and the Spanish Ministerio de Economía y Competitividad under contract FPA2014-57196-C5-4-P. S.L.T. acknowledges financial support from the U.S. National Science Foundation under contract NSF-14-01574. E.P.-R. acknowledges financial support from the DGAPA-UNAM under the PASPA program. The GRIFFIN spectrometer was funded by the Canada Foundation for Innovation, TRIUMF, and the University of Guelph. TRIUMF receives federal funding via a contribution agreement with the National Research Council of Canada.

- [1] M. R. Mumpower, R. Surman, G. C. McLaughlin, and A. Aprahamian, *Prog. Part. Nucl. Phys.* **86**, 86 (2016).
 [2] E. M. Burbidge, G. R. Burbidge, W. A. Fowler, and F. Hoyle, *Rev. Mod. Phys.* **29**, 547 (1957).

- [3] A. G. W. Cameron, *Publ. Astron. Soc. Pac.* **69**, 201 (1957).
 [4] J. J. Cowan, F.-K. Thielemann, and J. W. Truran, *Phys. Rep.* **208**, 267 (1991).

- [5] M. Arnould, S. Goriely, and K. Takahashi, *Phys. Rep.* **450**, 97 (2007).
- [6] F.-K. Thielemann *et al.*, *J. Phys.: Conf. Ser.* **202**, 012006 (2010).
- [7] C. Freiburghaus, S. Rosswog, and F.-K. Thielemann, *Astrophys. J.* **525**, L121 (1999).
- [8] O. Korobkin, S. Rosswog, A. Arcones, and C. Winteler, *Mon. Not. R. Astron. Soc.* **426**, 1940 (2012).
- [9] J. J. Cuenca-García, G. Martínez-Pinedo, K. Langanke, F. Nowacki, and I. N. Borzov, *Eur. Phys. J. A* **34**, 99 (2007).
- [10] Q. Zhi, E. Caurier, J. J. Cuenca-García, K. Langanke, G. Martínez-Pinedo, and K. Sieja, *Phys. Rev. C* **87**, 025803 (2013).
- [11] M. Hannawald *et al.*, *Nucl. Phys. A* **688**, 578 (2001).
- [12] G. Lorusso *et al.*, *Phys. Rev. Lett.* **114**, 192501 (2015).
- [13] P.-A. Söderström *et al.*, *Nucl. Instrum. Methods Phys. Res., Sect. B* **317**, 649 (2013).
- [14] B. Fogelberg *et al.*, *Proc. Intern. Conf. Nuclear Data for Science and Technology*, Mito, Japan (Saikon Publishing Co., Tokyo, Japan, 1988), p. 837.
- [15] O. Arndt, I. Dillmann, U. Koester, K.-L. Kratz, A. N. Ostrowski, B. Pfeiffer, J. Shergur, W. B. Walters, and A. Woehr, *Annual Reports of the Institute for Nuclear Chemistry*, University of Mainz, Germany, <http://www.kernchemie.uni-mainz.de/downloads/jb2003>, A15 (2003).
- [16] O. Arndt *et al.*, *Acta Phys. Pol., B* **40**, 437 (2009).
- [17] D. T. Yordanov *et al.*, *Phys. Rev. Lett.* **110**, 192501 (2013).
- [18] J. Taprogge *et al.*, *Phys. Rev. C* **91**, 054324 (2015).
- [19] K.-L. Kratz, H. Gabelmann, W. Hillebrandt, B. Pfeiffer, K. Schlosser, and F.-K. Thielemann, *Z. Phys. A* **325**, 489 (1986).
- [20] J. Taprogge, Ph.D. thesis, Universidad Autónoma de Madrid, 2015 (unpublished).
- [21] C. E. Svensson and A. B. Garnsworthy, *Hyperfine Interact.* **225**, 127 (2014).
- [22] U. Rizwan *et al.*, *Nucl. Instrum. Methods Phys. Res., Sect. A* **820**, 126 (2016).
- [23] J. Dilling, R. Krücken, and G. Ball, *Hyperfine Interact.* **225**, 1 (2013).
- [24] S. Raeder, H. Heggen, J. Lassen, F. Ames, D. Bishop, P. Bricault, P. Kunz, A. Mjøos, and A. Teigelhöfer, *Rev. Sci. Instrum.* **85**, 033309 (2014).
- [25] G. C. Ball *et al.*, *J. Phys. G* **31**, S1491 (2005).
- [26] H. Göktürk, B. Ekström, E. Lund, and B. Fogelberg, *Z. Phys. A* **324**, 117 (1986).
- [27] G. F. Grinyer *et al.*, *Phys. Rev. C* **71**, 044309 (2005).
- [28] A. T. Laffoley *et al.*, *Phys. Rev. C* **88**, 015501 (2013).
- [29] G. F. Grinyer *et al.*, *Nucl. Instrum. Methods Phys. Res., Sect. A* **579**, 1005 (2007).
- [30] H. Gausemel, B. Fogelberg, T. Engeland, M. Hjorth-Jensen, P. Hoff, H. Mach, K. A. Mezilev, and J. P. Omtvedt, *Phys. Rev. C* **69**, 054307 (2004).
- [31] K.-L. Kratz *et al.*, *Eur. Phys. J. A* **25**, 633 (2005).
- [32] I. Dillmann *et al.*, *Phys. Rev. Lett.* **91**, 162503 (2003).
- [33] B. Fogelberg, K. Heyde, and J. Sau, *Nucl. Phys. A* **352**, 157 (1981).
- [34] I. N. Borzov, J. J. Cuenca-García, K. Langanke, G. Martínez-Pinedo, and F. Montes, *Nucl. Phys. A* **814**, 159 (2008).
- [35] T. Marketin, L. Huther, and G. Martínez-Pinedo, *Phys. Rev. C* **93**, 025805 (2016).