α -decay branching ratio of ¹⁸⁰Pt

J. G. Cubiss^{1,*} R. D. Harding,^{1,2} A. N. Andreyev,^{1,3} N. Althubiti,^{4,5} B. Andel^{6,6,7} S. Antalic^{6,6} A. E. Barzakh,⁸ T. E. Cocolios^{6,7,4} T. Day Goodacre,^{4,2,†} G. J. Farooq-Smith,^{4,7} D. V. Fedorov,⁸ V. N. Fedosseev^{6,2} L. P. Gaffney^{6,9,7,‡} L. Ghys,^{10,7} M. Huyse,⁷ K. M. Lynch,² B. A. Marsh,² Y. Martinez Palenzuela,⁷ P. L. Molkanov,⁸ R. E. Rossel,^{2,11} S. Rothe,² M. D. Seliverstov^{6,8} S. Sels^{6,7,8} P. Spagnoletti^{6,9} C. Van Beveren,⁷ P. Van Duppen^{6,7}

M. Veinhard,² E. Verstraelen,⁷ and A. Zadvornaya^{7,||}

¹Department of Physics, University of York, York YO10 5DD, United Kingdom

²CERN, CH-1211 Geneve 23, Switzerland

³Advanced Science Research Center (ASRC), Japan Atomic Energy Agency (JAEA), Tokai-mura, Ibaraki 319-1195, Japan

⁴School of Physics and Astronomy, The University of Manchester, Manchester M13 9PL, United Kingdom

⁵*Physics Department, Faculty of Science, Jouf University, Aljouf, Saudi Arabia*

⁶Department of Nuclear Physics and Biophysics, Comenius University in Bratislava, 84248 Bratislava, Slovakia

⁷KU Leuven, Instituut voor Kern - en Stralingsfysica, B-3001 Leuven, Belgium

⁸Petersburg Nuclear Physics Institute, NRC Kurchatov Institute, Gatchina 188300, Russia

⁹School of Engineering and Computing, University of the West of Scotland, Paisley PA1 2BE, United Kingdom

¹⁰Belgian Nuclear Research Center SCK•CEN, Boeretang 200, B-2400 Mol, Belgium

¹¹Institut für Physik, Johannes Gutenberg-Universität, 55122 Mainz, Germany

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A study of the ¹⁸⁰Hg decay chain performed at the CERN-ISOLDE facility has allowed the ground-state–to– ground-state α decay of ¹⁸⁰Pt to be investigated. A more precise α -decay branching ratio of $b_{\alpha}(^{180}\text{Pt}) = 0.52(5)\%$ has been deduced. The reduced α -decay width calculated using the new value provides a more consistent picture of the systematics for $J^{\pi} = 0^+ \rightarrow 0^+$ ground-state–to–ground-state state α decays of neutron-deficient, eveneven platinum isotopes.

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I. INTRODUCTION

Alpha decay is a useful probe for studying the underlying structures of nuclei involved in the process. For example, reduced α -decay widths (δ_{α}^2) are particularly sensitive to the overlap in wave function between the initial and final states connected by the decay [1]. These may be calculated, for instance, using the Rasmussen approach [2], which requires experimental α -decay energies and partial half-lives. The latter are dependent on α -decay branching ratios (b_{α}), which are often challenging to measure in nuclei with small b_{α} values

due to only low statistics and/or the presence of more intense α decays.

In this work, we report on a more precise b_{α} for the ¹⁸⁰Pt ground state (g.s.). This value was extracted from decay data recorded during the same experiment as described in Refs. [3,4].

The currently accepted value $b_{\alpha}(^{180}\text{Pt}) \approx 0.3\%$ came from a study by Siivola [5], in which ¹⁸⁰Pt was produced in ¹⁶O + ^{170,172}Yb, ¹⁹F + ¹⁶⁹Tm, and ²⁰Ne + ¹⁶²Er fusion-evaporation reactions. The b_{α} values for several platinum isotopes were deduced by comparing measured α -decay intensities to expected production yields, based on similar heavy-ion reactions studied in the rare-earth region. Due to this approach, the extracted value of $b_{\alpha}(^{180}\text{Pt}) \approx 0.3\%$ had a large uncertainty factor of 3–5 [5].

II. EXPERIMENT

A detailed description of the experiment can be found in Refs. [3,4], while only the information pertinent to the present work is provided here. In our study, ¹⁸⁰Pt was produced in the ¹⁸⁰Hg \rightarrow ¹⁸⁰Au \rightarrow ¹⁸⁰Pt β -decay chain, shown in Fig. 1. An isotopically pure ion beam of ¹⁸⁰Hg was produced at the ISOLDE facility [13,14] through spallation reactions induced by a 1.4-GeV proton beam impinged upon a molten lead target, followed by a three-step, resonance laser ionization

^{*}james.cubiss@york.ac.uk

[†]Present address: TRIUMF, 4004 Wesbrook Mall, Vancouver, BC, Canada V6T 2A3.

[‡]Present address: Oliver Lodge Laboratory, University of Liverpool, Liverpool L69 7ZE, United Kingdom.

[§]Present address: CERN, CH-1211 Geneve 23, Switzerland.

^{II}Present address: Department of Physics, University of Jyväskylä, P.O. Box 35, SF-40351, Finland.

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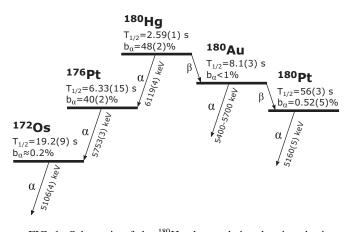


FIG. 1. Schematic of the ¹⁸⁰Hg decay chain, showing the isotopes and α -decay energies important to the present work. The $b_{\alpha}(^{180}\text{Pt}) = 0.52(5)\%$ value is taken from the results of our study (see Sec. III), and $E_{\alpha}(^{180}\text{Pt}) = 5160(5)$ keV is from Ref. [6]. All other data are taken from Refs. [7-12].

process in the VADLIS [15,16] in order to selectively ionize the mercury atoms of interest. The ions were extracted and accelerated by a 30-kV potential difference, and separated according to their mass-to-charge ratio by the ISOLDE general purpose separator.

The ¹⁸⁰Hg ion beam was then delivered to the Windmill system [17,18] for decay measurements. The beam entered the Windmill through the central hole of an annular silicon detector (Si1), and was implanted into one of ten, $20 - \mu g \text{ cm}^{-2}$ thick carbon foils mounted upon a rotatable wheel. A second silicon detector (Si2) was placed a few millimeters behind the foil being irradiated. The data presented in the current work were taken in runs where the wheel was not rotated to avoid loss of activity during the dedicated b_{α} measurements. Furthermore, due to the specific conditions during this measurement, only events recorded in Si2 were used in the following analysis. The full width at half maxima of the α decay peaks recorded by Si2 within the $E_{\alpha} = 5000-6200$ -keV region of interest were \approx 30 keV.

The energy calibration for Si2 was performed using $E_{\alpha}(^{180}\text{Hg}) = 6119(4) \text{ keV} [19] \text{ and } E_{\alpha}(^{180}\text{Pt}) = 5160(5) \text{ keV}$ for the g.s.-to-g.s. decay of ^{180}Pt . The latter was deduced in our recent study of gold isotopes and will be discussed in Ref. [6]. Our new value differs significantly from $E_{\alpha}(^{180}\text{Pt}) =$ 5140(10) keV reported by Siivola [5] but has a higher precision. In addition to the energy calibration, our $E_{\alpha}(^{180}\text{Pt})$ value will be used in the δ_{α}^2 calculations presented in Sec. IV.

III. RESULTS

Figure 2(a) shows the singles α -decay spectrum recorded by Si2. The spectrum readily illustrates the purity of the ¹⁸⁰Hg beam, as only the decays of ¹⁸⁰Hg, its daughter and its granddaughter nuclei are seen.

The two most intense peaks in Fig. 2(a) belong to the well-known decays of ¹⁸⁰Hg [$E_{\alpha} = 6119(4)$ keV] [19] and ¹⁷⁶Pt [$E_{\alpha} = 5753(3)$ keV] [8]. The low-intensity $E_{\alpha}(^{180}\text{Hg}) =$ 5862(5) keV fine-structure (f.s.) decay is also visible [20]. The

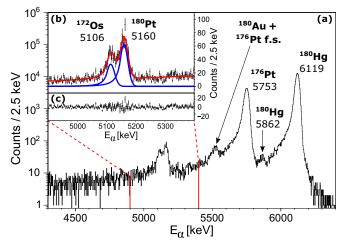


FIG. 2. (a) Energy spectrum of singles α -decay events measured in Si2 at A = 180, (b) zoomed view of the 4900–5400-keV region, fitted with a linear background plus two Crystal Ball functions (red line), the individual contributions from the α decays of ¹⁷²Os and ¹⁸⁰Pt are shown by the blue lines, (c) the residual between the fit and the data shown in panel (b). The main α -decay peaks are labeled with the corresponding isotope and α -particle energies.

structure seen in the $E_{\alpha} = 5460-5560$ -keV region is due to the f.s. α line of ¹⁷⁶Pt [$E_{\alpha} = 5530(3)$ keV] [12] and the complex f.s. decay of ¹⁸⁰Au [6,20,21]. The α -decay peak of ¹⁸⁰Pt is seen to be partially overlapping with the 5106(4)-keV ¹⁷²Os α -decay peak [12]. The two peaks lie on top of a significant background from the low-energy tails of the higher-energy and higher-intensity α decays of ¹⁷⁶Pt and ¹⁸⁰Hg.

The g.s.-to-g.s. $b_{\alpha}(^{180}\text{Pt})$ value was deduced by using the number of ¹⁸⁰Hg α decays in Fig. 2, N_{α} (¹⁸⁰Hg), to calculate the number of β decays feeding to ¹⁸⁰Au and then to ¹⁸⁰Pt (see decay scheme in Fig. 1). This approach treats the number of ¹⁸⁰Au and ¹⁸⁰Hg β decays as approximately equal, as the correction for the small ¹⁸⁰Au α -decay branch [b_{α} (¹⁸⁰Au) \approx 0.6% taken from Ref. [6]]¹ is negligible compared to the statistical error on N_{α} ⁽¹⁸⁰Pt) extracted from Fig. 2 ($\approx 10\%$). Thus, $N_{\alpha}(^{180}\text{Hg})$ and $N_{\alpha}(^{180}\text{Pt})$ may be directly compared in order to calculate $b_{\alpha}(^{180}\text{Pt})$, such that

$$b_{\alpha}(^{180}\text{Pt}) = \frac{N_{\alpha}(^{180}\text{Pt})}{\frac{N_{\alpha}(^{180}\text{Hg})}{b_{\alpha}(^{180}\text{Hg})}(1 - b_{\alpha}(^{180}\text{Hg}))},$$
(1)

where $b_{\alpha}({}^{180}\text{Hg}) = 48(2)\%$ [11,12]. To evaluate $N_{\alpha}({}^{180}\text{Pt})$, the $E_{\alpha} = 4900-5900$ -keV region of Fig. 2(b) was fitted with the ROOT Minuit minimizer [22], using a binned-likelihood method. A linear function was used to model the background and Crystal Ball functions [23-25], which shared the same set of parameters to describe the width and tails of the peaks were used for the ¹⁸⁰Pt and

¹A study at SHIP [21] deduced a lower limit of b_{α} (¹⁸⁰Au) > 1.8%. However, this limit was calculated using the $b_{\alpha}(^{180}\text{Pt}) \approx 0.3\%$ value with the factor of 3–5 uncertainty [5]. Furthermore, the expression used to calculate $b_{\alpha}(^{180}\text{Au})$ was incorrect (see Table 2 in Ref. [21]), as confirmed in private communications with the authors of the study.

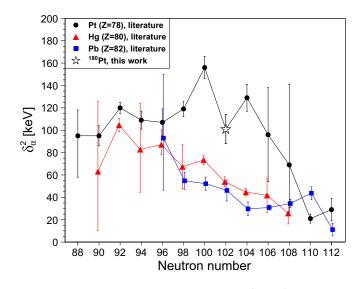


FIG. 3. Reduced α -decay widths for $J^{\pi} = 0^+ \rightarrow 0^+$, g.s.-to-g.s. decays of even-*A* platinum (•), mercury (**A**), and lead (**B**) isotopes, calculated using the Rasmussen approach [2] with data taken from Refs. [8–12,26,27]. The open star is the δ^2_{α} ⁽¹⁸⁰Pt) calculated using the results from the present work and Ref. [6].

¹⁷²Os α -decay peaks. The red line in Fig. 2(b) shows the result of the fitting procedure, the blue lines represent the contributions from the ¹⁸⁰Pt and ¹⁷²Os α decays. Figure 2(c) shows the residual between the result of the fitting procedure and the data shown in Fig. 2(b). The N_{α} (¹⁸⁰Hg) value was assessed in a similar way, however we assumed background-free conditions. Using the results from the fitting procedure a value of b_{α} (¹⁸⁰Pt) = 0.52(5)% was deduced.

IV. DISCUSSION

The reduced widths of $J^{\pi} = 0^+ \rightarrow 0^+$ g.s.-to-g.s. decays of even-*A* platinum isotopes calculated using the Rasmussen approach [2] are shown in Fig. 3, along with those for even-*A* mercury and lead isotopes. The open star represents $\delta_{\alpha}^2(^{180}\text{Pt}) = 101(13)$ keV, calculated using the b_{α} value from the current work and $E_{\alpha} = 5160(5)$ keV from Ref. [6].

In general, Fig. 3 displays the expected behavior (see Fig. 3 in Ref. [28] and Fig. 4 in Ref. [29]), whereby the $\delta_{\alpha}^2(0_{g.s.}^+ \rightarrow 0_{g.s.}^+)$ values, or equivalently the α -particle preformation probabilities, decrease as the proton number approaches Z = 82. This trend was interpreted in Ref. [28] as an effect of the Z =82 shell closure on the α -decay process. As further shown in Ref. [29], the N = 126 shell closure displays a similar influence, whereby δ_{α}^2 values at N = 126 are the smallest along an isotopic chain and an increase followed by a saturation in δ_{α}^2 is observed as N reduces towards and beyond the N = 104midshell.

Our new $\delta_{\alpha}^2({}^{180}\text{Pt})$ value is first of all in better agreement with these systematics than $\delta_{\alpha}^2({}^{180}\text{Pt}) = 74$ keV calculated using the data from Ref. [5]. Furthermore, the new value

reveals that as expected [29] the δ_{α}^2 for platinum isotopes is saturated in the $88 \le N \le 104$ region, with near-constant values of $\delta_{\alpha}^2 \approx 113$ keV.

One noticeable feature in Fig. 3 is the $\delta_{\alpha}^2 = 156(10)$ -keV value for ¹⁷⁸Pt (N = 100) which is $\approx 30\%$ larger than those of other platinum isotopes in the saturation region. The $T_{1/2}(^{178}$ Pt) [30–32] and $b_{\alpha}(^{178}$ Pt) [33,34] values from different studies are consistent with one another, which suggests that the experimental $\delta_{\alpha}^2(^{178}$ Pt) value is reliable. Therefore the jump in δ_{α}^2 could possibly be related to the change in deformation when going from ¹⁷⁸Pt to ¹⁸⁰Pt [35], and related to the possible change in the configuration mixing in the corresponding ground states [20,36,37]. Alternatively, this could be a sign of evolving nuclear structures between the α -decay daughter nuclei, ^{174,176}Os.

In addition to the large δ_{α}^2 ⁽¹⁷⁸Pt) value, the other noticeable features of the platinum chain in Fig. 3 are the sizable error bars on δ_{α}^2 ^{(184,186}Pt) (N = 106, 108). More precise measurements of b_{α} ^{(184,186}Pt) are required in order to determine whether there is a smooth transition towards the saturated δ_{α}^2 values, as would usually be expected.

V. CONCLUSION

Decay data recorded at the CERN-ISOLDE facility have been used to deduce a more precise value of $b_{\alpha}(^{180}\text{Pt}) = 0.52(5)\%$. This value has been used to calculate the reduced width of the ¹⁸⁰Pt g.s.-to-g.s. α decay, which is in better agreement with the δ_{α}^2 systematics in the region than the value calculated using the current literature value.

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- P. Van Duppen and M. Huyse, Hyperfine Interact. 129, 149 (2000).
- [2] J. O. Rasmussen, Phys. Rev. 113, 1593 (1959).
- [3] B. A. Marsh et al., Nat. Phys. 14, 1163 (2018).
- [4] S. Sels et al., Phys. Rev. C 99, 044306 (2019).
- [5] A. Siivola, Nucl. Phys. 84, 385 (1966).
- [6] R. D. Harding et al. (unpublished).
- [7] J. Husson, C. Liang, and C. Richard-Serre, J. Physique Lett. 38, 245 (1977).
- [8] B. Singh, Nucl. Data Sheets 75, 199 (1995).
- [9] M. Basunia, Nucl. Data Sheets 107, 791 (2006).
- [10] C. M. Baglin, Nucl. Data Sheets 111, 1807 (2010).
- [11] E. McCutchan, Nucl. Data Sheets 126, 151 (2015).
- [12] NNDC, Evaluated Nuclear Structure Data File, 2019.
- [13] E. Kugler, Hyperfine Interact. 129, 23 (2000).
- [14] R. Catherall, W. Andreazza, M. Breitenfeldt, A. Dorsival, G. J. Focker, T. P. Gharsa, T. J. Giles, J.-L. Grenard, F. Locci, P. Martins *et al.*, J. Phys. G: Nucl. Part. Phys. 44, 094002 (2017).
- [15] T. Day Goodacre *et al.*, Nucl. Instrum. Methods Phys. Res., Sect. B **376**, 39 (2016).
- [16] Y. Martinez Palenzuela *et al.*, Nucl. Instrum. Methods Phys. Res., Sect. B 431, 59 (2018).
- [17] J. G. Cubiss et al., Phys. Rev. C 97, 054327 (2018).
- [18] A. N. Andreyev et al., Phys. Rev. Lett. 105, 252502 (2010).
- [19] F. G. Kondev et al., Phys. Rev. C 62, 044305 (2000).
- [20] J. Wauters, P. Dendooven, M. Huyse, G. Reusen, P. Van Duppen, R. Kirchner, O. Klepper, and E. Roeckl, Z. Phys. A: Hadrons Nucl. 345, 21 (1993).

- [21] J. G. Keller, K.-H. Schmidt, F. P. Hessberger, G. Münzenberg, W. Reisdorf, H.-G. Clerc, and C.-C. Sahm, Nucl. Phys. A 452, 173 (1986).
- [22] F. James and M. Roos, Comput. Phys. Commun. 10, 343 (1975).
- [23] M. J. Oreglia, Ph.D. thesis, SLAC-R-236, 1980.
- [24] J. E. Gaiser, Ph.D. thesis, SLAC-R-255, 1982.
- [25] T. Skwarnicki, Ph.D thesis, DESY F31-86-02, 1986.
- [26] H. Badran et al., Phys. Rev. C 94, 054301 (2016).
- [27] J. Hilton et al., Phys. Rev. C 100, 014305 (2019).
- [28] A. N. Andreyev et al., Phys. Rev. Lett. 110, 242502 (2013).
- [29] C. Qi, A. N. Andreyev, M. Huyse, R. J. Liotta, P. Van Duppen, and R. Wyss, Phys. Lett. B 734, 203 (2014).
- [30] J. D. Bowman, R. E. Eppley, and E. K. Hyde, Phys. Rev. C 25, 941 (1982).
- [31] F. Meissner, H. Salewski, W. D. Schmidt-Ott, U. Bosch-Wicke, V. Kunze, and R. Michaelsen, Phys. Rev. C 48, 2089 (1993).
- [32] F. G. Kondev et al., Phys. Rev. C 61, 044323 (2000).
- [33] P. G. Hansen, H. L. Nielsen, K. Wilsky, M. Alpsten, M. Finger, A. Lindahl, R. A. Naumann, and O. B. Nielsen, Nucl. Phys. A 148, 249 (1970).
- [34] U. J. Schrewe et al., Phys. Lett. B 91, 46 (1980).
- [35] F. Le Blanc et al., Phys. Rev. C 60, 054310 (1999).
- [36] G. D. Dracoulis, B. Fabricius, A. E. Stuchbery, A. O. Macchiavelli, W. Korten, F. Azaiez, E. Rubel, M. A. Deleplanque, R. M. Diamond, and F. S. Stephens, Phys. Rev. C 44, R1246 (1991).
- [37] J. L. Wood, K. Heyde, W. Nazarewicz, M. Huyse, and P. van Duppen, Phys. Rep. 215, 101 (1992).