

LUND UNIVERSITY

Isomeric Decay Spectroscopy of the 217Bi Isotope

Gottardo, A.; Valiente-Dobón, J. J.; Benzoni, G.; Lunardi, S.; Gadea, A.; Algora, A.; Al-Dahan, N.; de Angelis, G.; Ayyad, Y.; Bazzacco, D.; Benlliure, J.; Boutachkov, P.; Bowry, M.; Bracco, A.; Bruce, A. M.; Bunce, M.; Camera, F.; Casarejos, E.; Cortes, M. L.; Crespi, F. C. L.; Corsi, A.; Denis Bacelar, A. M.; Deo, A. Y.; Domingo-Pardo, C.; Doncel, M.; Engert, T.; Eppinger, K.; Farrelly, G. F.; Farinon, F.; Farnea, E.; Geissel, H.; Gerl, J.; Goel, N.; Górska, M.; Grebosz, J.; Gregor, E.; Habermann, T.; Hoischen, Robert; Janik, R.; John, P. R.; Klupp, S.; Kojouharov, I.; Kurz, N.; Lenzi, S. M.; Leoni, S.; Mandal, S.; Menegazzo, R.; Mengoni, D.; Million, B.; Modamio, V.

Published in:

Physical Review C (Nuclear Physics)

DOI:

10.1103/PhysRevC.90.034317

2014

Link to publication

Citation for published version (APA):

Gottardo, A., Valiente-Dobón, J. J., Benzoni, G., Lunardi, S., Gadea, A., Algora, A., Al-Dahan, N., de Angelis, G., Ayyad, Y., Bazzacco, D., Benlliure, J., Boutachkov, P., Bowry, M., Bracco, A., Bruce, A. M., Bunce, M., Camera, F., Casarejos, E., Cortes, M. L., ... Wollersheim, H.-J. (2014). Isomeric Decay Spectroscopy of the 217Bi Isotope. *Physical Review C (Nuclear Physics)*, *90*(3), Article 034317. https://doi.org/10.1103/PhysRevC.90.034317

Total number of authors:

General rights

Unless other specific re-use rights are stated the following general rights apply:

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights

· Users may download and print one copy of any publication from the public portal for the purpose of private study

· You may not further distribute the material or use it for any profit-making activity or commercial gain

You may freely distribute the URL identifying the publication in the public portal

Read more about Creative commons licenses: https://creativecommons.org/licenses/

Take down policy If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Download date: 17. Dec. 2025

LUND UNIVERSITY

PO Box 117 221 00 Lund +46 46-222 00 00

Isomeric decay spectroscopy of the ²¹⁷Bi isotope

```
A. Gottardo, <sup>1,2,*</sup> J. J. Valiente-Dobón, <sup>1</sup> G. Benzoni, <sup>3</sup> S. Lunardi, <sup>2,4</sup> A. Gadea, <sup>5</sup> A. Algora, <sup>5,6</sup> N. Al-Dahan, <sup>7</sup> G. de Angelis, <sup>1</sup> Y. Ayyad, <sup>8</sup> D. Bazzacco, <sup>4</sup> J. Benlliure, <sup>8</sup> P. Boutachkov, <sup>9</sup> M. Bowry, <sup>7</sup> A. Bracco, <sup>3,10</sup> A. M. Bruce, <sup>11</sup> M. Bunce, <sup>11</sup> F. Camera, <sup>3,10</sup>
      E. Casarejos, <sup>12</sup> M. L. Cortes, <sup>9</sup> F. C. L. Crespi, <sup>3,10</sup> A. Corsi, <sup>3,10</sup> A. M. Denis Bacelar, <sup>11</sup> A. Y. Deo, <sup>7,13</sup> C. Domingo-Pardo, <sup>9</sup>
  M. Doncel, <sup>14</sup> T. Engert, <sup>9</sup> K. Eppinger, <sup>15</sup> G. F. Farrelly, <sup>7</sup> F. Farinon, <sup>9</sup> E. Farnea, <sup>4</sup> H. Geissel, <sup>9</sup> J. Gerl, <sup>9</sup> N. Goel, <sup>9</sup> M. Górska, <sup>9</sup> J. Grebosz, <sup>16</sup> E. Gregor, <sup>9</sup> T. Habermann, <sup>9</sup> R. Hoischen, <sup>9,17</sup> R. Janik, <sup>18</sup> P. R. John, <sup>2,4</sup> S. Klupp, <sup>15</sup> I. Kojouharov, <sup>9</sup> N. Kurz, <sup>9</sup> S. M. Lenzi, <sup>2,4</sup> S. Leoni, <sup>3,10</sup> S. Mandal, <sup>19</sup> R. Menegazzo, <sup>4</sup> D. Mengoni, <sup>4</sup> B. Million, <sup>3</sup> V. Modamio, <sup>1</sup> A. I. Morales, <sup>3</sup> D. R. Napoli, <sup>1</sup> F. Naqvi, <sup>9,20</sup> R. Nicolini, <sup>3,10</sup> C. Nociforo, <sup>9</sup> M. Pfützner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>21</sup> S. Pietri, <sup>9</sup> Zs. Podolyák, <sup>7</sup> A. Prochazka, <sup>9</sup> M. Pittzner, <sup>9</sup> S. Pitti, 
    W. Prokopowicz, F. Recchia, P. H. Regan, M. W. Reed, D. Rudolph, T. E. Sahin, H. Schaffner, A. Sharma, B. Sitar, R. D. Siwal, K. Steiger, L. P. Strmen, R. T. P. D. Swan, I. Szarka, R. C. A. Ur, P. M. Walker, H. Weick,
                                                                                                                               O. Wieland,<sup>3</sup> and H-J. Wollersheim<sup>9</sup>
                                                           <sup>1</sup>Istituto Nazionale di Fisica Nucleare, Laboratori Nazionali di Legnaro, 35020 Legnaro, Italy
                                                        <sup>2</sup>Dipartimento di Fisica e Astronomia dell'Università degli Studi di Padova, 35131 Padova, Italy
                                                                               <sup>3</sup>Istituto Nazionale di Fisica Nucleare, Sezione di Milano, 20133 Milano, Italy
                                                                               <sup>4</sup>Istituto Nazionale di Fisica Nucleare, Sezione di Padova, 35131 Padova, Italy
                                                                 <sup>5</sup>Instituto de Física Corpuscular, CSIC-Universitat de València, E-46980 València, Spain
                                                        <sup>6</sup>Institute of Nuclear Research of the Hungarian Academy of Sciences, 4026 Debrecen, Hungary
                                                                      <sup>7</sup>Department of Physics, University of Surrey, Guildford, GU2 7XH, United Kingdom
                                                                      <sup>8</sup>Universidade de Santiago de Compostela, E-175706 Santiago de Compostela, Spain
                                                                         <sup>9</sup>GSI Helmholtzzentrum für Schwerionenforschung, D-64291 Darmstadt, Germany
                                                                         <sup>10</sup>Dipartimento di Fisica dell'Università degli Studi di Milano, 20133 Milano, Italy
                               <sup>11</sup>School of Computing, Engineering and Mathematics, University of Brighton, Brighton, BN2 4GJ, United Kingdom
                                                                                                                 <sup>12</sup>EEI, Universidade de Vigo, E-36310 Vigo, Spain
                                                                  <sup>13</sup>Department of Physics, Indian Institute of Technology Roorkee, Roorkee 247667, India
                                                                         <sup>14</sup>Grupo de Física Nuclear, Universidad de Salamanca, E-37008 Salamanca, Spain
                                                                      <sup>15</sup>Physik Department, Technische Universität München, D-85748 Garching, Germany
                                               <sup>16</sup>Niewodniczanski Institute of Nuclear Physics, Polish Academy of Science, PL-31-342 Krakow, Poland
                                                                                               <sup>17</sup>Department of Physics, Lund University, S-22100 Lund, Sweden
                                                                  <sup>18</sup>Faculty of Mathematics and Physics, Comenius University, 84215 Bratislava, Slovakia
                                                                         <sup>19</sup>Department of Physics and Astrophysics, University of Delhi, Delhi 110007, India
                                                                                        <sup>20</sup>Institut für Kernphysik, Universität zu Köln, D-50937 Köln, Germany
                                                                                        <sup>21</sup> Faculty of Physics, University of Warsaw, PL-00681 Warsaw, Poland
```

The structure of the neutron-rich bismuth isotope 217 Bi has been studied for the first time. The fragmentation of a primary 238 U beam at the FRS-RISING setup at GSI was exploited to perform γ -decay spectroscopy, since μs isomeric states were expected in this nucleus. Gamma rays following the decay of a $t_{1/2}=3$ μs isomer were observed, allowing one to establish the low-lying structure of 217 Bi. The level energies and the reduced electric quadrupole transition probability B(E2) from the isomeric state are compared to large-scale shell-model calculations.

(Received 5 June 2014; revised manuscript received 14 August 2014; published 23 September 2014)

DOI: 10.1103/PhysRevC.90.034317 PACS number(s): 23.35.+g, 21.10.Tg, 23.20.-g, 27.80.+w

I. INTRODUCTION

The study of nuclei far from stability is a major research field in modern nuclear physics, a field that has grown substantially with the advent of radioactive ion beams. Various regions of the nuclide chart have been explored with stable beams using mainly fusion-evaporation, deep-inelastic, or fission reactions. However, the neutron-rich isotopes around lead, Z=82, have always been difficult to populate with

the aforementioned reactions. In the last fifteen years their study has been made gradually possible by the use of fragmentation reactions combined with in-flight mass separators and advanced setups for decay spectroscopy. For example, the fragmentation of a uranium beam was used to produce 212 Pb and 211 Bi and measure their isomeric decay [1]. Similarly, the adjacent elements beyond N=126 and below Z=82, such as thallium and mercury, have been studied [2,3]. For the elements beyond N=126 but well above Z=82 the situation is very different, as they can be populated with comparatively large cross sections with spallation or fragmentation reactions on uranium. The α decay from these heavy isotopes can also populate lighter nuclei toward the lead region, enabling their spectroscopic study. However, the bismuth isotopes, one

^{*}andrea.gottardo@lnl.infn.it; present address: Institut de Physique Nucléaire, CNRS/IN2P3, and Université Paris Sud, Orsay, 91406, France.

proton above lead, are not all reachable via decay of easy-to-produce heavy nuclei. The 211 Bi isotope has been studied via transfer and fragmentation reactions [1,4,5] and via the β and α decay of 211 Pb and 215 At, respectively [6,7], leading to the discovery of two isomeric states in the same decay sequence. The 213 Bi nucleus has also been studied by α decay of 217 At, and its first excited state, $I^{\pi}=13/2^{-}$, was observed [8]. A long-lived isomer has recently been observed in 213 Bi, produced via 238 U fragmentation, using the technique of mass measurement in a storage ring [9]. Besides 211 Bi, the only other known isomeric state in the neutron-rich odd-even bismuth isotopes is in 215 Bi, six neutrons above N=126, which was populated with a spallation reaction at ISOLDE [10]. Such reactions at ISOLDE were also successfully employed for the production of 215 Pb [11] and 218 Bi [12].

Recent improvements in experimental devices and beam intensities presently allow exploring more extensively the exotic neutron-rich nuclei in this region, via fragmentation reactions from a 238 U beam [13]. For example, the neutron-rich lead isotopes were studied, by taking advantage of long-lived (μ s range) 8⁺ seniority isomers, predicted by theory and indeed identified up to 216 Pb [13]. Nearby isotopes were also effectively populated for the first time, up to mass number 219 for the bismuth nuclei and to mass 210 for the mercury isotopes [14–16].

In more detail, in this mass region neutrons are filling the $2g_{9/2}$ shell beyond the N = 126 shell closure, and this gives rise in the even-even lead isotopes to a low-lying level structure which agrees with the predictions of the seniority scheme [13]. A very basic expectation is that the level structure of ^{211–217}Bi follows the same pattern, since the unpaired proton outside the Z = 82 shell should couple with the excited levels of the corresponding even-even lead core. The situation can be complicated by the fact that the coupling gives rise to a multiplet of states with different spins, which can allow the decay of the seniority isomer to proceed through several branchings. It is also possible that the high-spin members of a multiplet lower the energy, thus creating spin traps with long half-lives ($t_{1/2} \sim \text{ms}$). Experimentally, the two known nuclei ^{211,215}Bi have basically the same low-spin structure [4,10], apart from a level inversion at high spin that forms the measured yrast trap in ²¹⁵Bi [10]. The long-lived isomer observed in ²¹³Bi is probably also coming from a level inversion. Knowledge of the ²¹⁷Bi nucleus, which is the object of the present study, is limited to its ground state, which $\beta^$ decays to excited states of ²¹⁷Po with a half-life of 92(3) s [17]. The spin and parity of the ground state are not known but from systematics one can safely assume that it is $9/2^-$, due to the occupation by the 83rd proton of the $h_{9/2}$ single-particle state.

The present paper, which reports the first spectroscopic study of the neutron-rich nucleus 217 Bi, is organized as follows. In the first section the experimental setup is described in detail. The second section provides the results from ion-mass identification and γ spectroscopy. The final section deals with the theoretical interpretation of the observed level scheme, in the framework provided by shell-model calculations and by taking into account the recent results on effective three-body forces in lead nuclei [13].

II. EXPERIMENTAL SETUP

The results of this work have been obtained by exploiting the advanced features of the state-of-the-art FRS-RISING setup [18-21] and the UNILAC-SIS-18 accelerator facilities at GSI by using a $1 \text{ GeV}/A^{238} \text{U}$ beam with an intensity of around 1.5×10^9 ions/spill. The \sim 1-s spills were separated by a \sim 2-s period without a beam. The uranium ions were fragmented on a 2.5 g/cm² Be target, followed by a 223 mg/cm² Nb stripper. The reaction products were separated and identified in mass and atomic number with the double-stage magnetic spectrometer FRS [18]. This is a mass spectrometer suitable for discriminating the different magnetic rigidities of relativistic beams, from light to heavy ions. The information gathered from its detectors (see later) allows one to unambiguously identify masses in the heavy region of interest ($A \sim 210-220$). The first particle detectors along the spectrometer were located at the second focal plane, where different charge states from the primary beam as well as other heavy ions can arrive, since their magnetic rigidity is similar to one of the isotopes of interest. These detectors cannot sustain the resulting high counting rate $(\sim 10^8 \text{ Hz})$ coming from the aforementioned contaminations. Therefore, a homogenous 2 g/cm² Al degrader was placed after the first dipole in order to exclude the heavy fragments above polonium from the acceptance of the FRS. The wedgeshaped Al degrader at the intermediate focal plane, after the second dipole, had a thickness of 758 mg/cm², and its angle was set to produce a monochromatic beam.

The identification in magnetic rigidity $(B\rho)$ is achieved through focal-plane position measurements with respect to the position of a beam with a well-known $B\rho$. The plastic scintillators at the intermediate and final focal planes allow extracting the time of flight (TOF). The mass-to-charge ratio (A/q) of the fragments is calculated from the TOF and the $B\rho$, measured on an event-by-event basis. The atomic number of the fragments is obtained from two ionization chambers placed in the final focal plane. Finally, the comparison of the $B\rho$ before and after the Al wedge-shaped degrader allows one to discriminate a possible change in the ion charge state. These measurements are sufficient to provide a complete identification of the isotopes event by event. Figure 1 shows a typical identification plot obtained. The different isotopes are clearly separated in both Z and A/q ratio (or better A/Z), since only the fully stripped ions were selected.

At the final focal plane, the ions were slowed down in a thick Al degrader in order to reduce the energy of the fragments of interest before they were implanted in a double-sided siliconstrip (DSSSD) detector system comprising three layers, each with three DSSSD pads [21,22]. The monochromatic beam ensured that the implantation depth in the active stopper was the same for all the fragments of a certain A/q and Z. The DSSSD detector system was surrounded by the RISING γ spectrometer, consisting of 105 germanium crystals arranged in 15 clusters with 7 crystals each [19,20]. The full-energy γ -ray peak detection efficiency of the array was measured to be 15% at 662 keV [19]. In the present experiment, due to the presence of the active stopper with its casing, the absolute efficiency of the array was \sim 13% at 662 keV. The time correlation between the γ rays and the ions detected with the

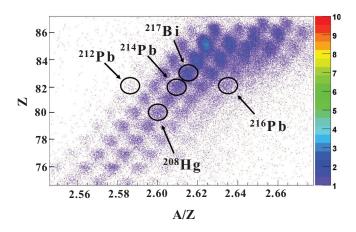


FIG. 1. (Color online) Ion identification plot at the final focal plane of the FRS. The ²¹⁷Bi products have been circled, as have some neighboring nuclei to provide a reference.

active stopper allowed one to perform at the same time isomer spectroscopy and β -delayed γ -ray spectroscopy [14,15].

III. EXPERIMENTAL RESULTS

Figure 2 shows the γ -ray spectrum following the detected isomeric decay of ²¹⁷Bi. Four transitions are clearly visible and their intensities are reported in Table I. The peak at 77 keV corresponds to the K_{α} x rays from bismuth.

Figure 3 presents the results of $\gamma\gamma$ coincidence analysis. The γ rays at 744, 492, and 200 keV are in mutual coincidence, and a coincidence relationship is also evident between the 744-and the 685-keV lines. As already outlined above, one can expect that the structure of bismuth nuclei is determined by the coupling of the single proton outside the Z=82 shell closure to the excited levels in 216 Pb. Since the lowest single-proton orbital above Z=82 is $h_{9/2}$, one expects that the low-lying levels in 217 Bi have the configuration $\pi h_{9/2} \otimes \nu g_{9/2}^2$ ($^{0^+...8^+}$). The isomeric state should have a spin-parity $25/2^-$, with

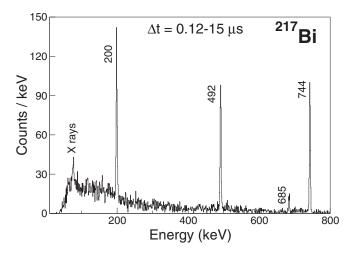


FIG. 2. Gamma-ray spectrum from the decay of the isomeric state in $^{217}{\rm Bi}$. The spectrum has been obtained by gating on the time window 0.12–15 $\mu{\rm s}$.

TABLE I. Areas, intensities corrected for efficiency and internal conversion [23], and $t_{1/2}$ of the isomeric state gated on the different γ -ray transitions measured in 217 Bi.

E_{γ} (keV)	Area	Intensity (%)	$t_{1/2} (\mu \mathrm{s})$
200	401 (38)	92 (9)	3.1 (2)
492	331 (32)	87 (8)	2.9 (2)
685	45 (13)	14 (4)	3.3 (7)
744	307 (35)	100 (11)	2.8 (1)

a $\pi h_{9/2} \otimes (\nu 2g_{9/2})^{8^+}$ configuration, corresponding to the 8^+ isomer of lead nuclei. Therefore, following the systematics from lighter odd-even bismuth isotopes, the 200-, 492-, and 744-keV γ rays are assigned to the cascade $21/2^- \rightarrow$ $17/2^- \to 13/2^- \to 9/2^-$. The $25/2^- \to 21/2^-$ transition is expected to have a low energy (in ²¹¹Bi it is only 30 keV), which makes it highly converted and unfeasible to measure with the present experimental setup. The 685-keV γ ray is in coincidence only with the 744-keV transition (see Fig. 3). It is hence assigned to a decay from a state located 685 keV above the 13/2 level and 7 keV lower than the 21/2 state. Figure 4 shows the exponential χ^2 fit to the decay curves of the four transitions. Within errors, the four fits give the same decay constant, which suggests that the four transitions might be following the decay of the same isomer, namely, the expected 25/2 state. Moreover, since one cannot exclude completely that the 685-keV γ ray follows the decay of a second isomer (with a very similar half-life) we have preferred to extract the half-life of the 25/2⁻ isomer from the error-weighted average of the decay constants of the 200- and 492-keV transitions. The isomer half-life deduced in this way is $3.0 \pm 0.2 \mu s$.

The characteristic K_{α} x rays from bismuth at 77 keV are observed with an intensity compatible with the internal conversion of the other four γ rays. Given that the binding energy of the K electrons in bismuth is 90.5 keV, this means that the transition directly depopulating the isomer must be below \sim 90 keV. From systematics in bismuth, lead, and mercury isotopes [2,13], we assume a lower limit

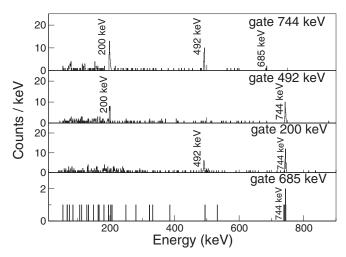


FIG. 3. Gamma-ray prompt coincidence spectra for the decay from the isomeric state in ²¹⁷Bi, with gates on the four transitions following the isomer.

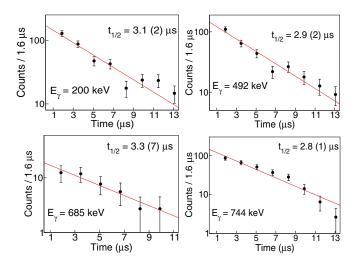


FIG. 4. (Color online) Time distributions and exponential fits (in red) for the 200-, 492-, 685-, and 744-keV transitions assigned to the $^{217}\mathrm{Bi}$ nucleus.

of 20 keV for this transition. Although the energy of the $25/2^- \rightarrow 21/2^-$ transition is not known, the fact that it has to be between 20 and 90 keV implies that it is highly converted, and consequently the energy dependence of the E2 transition rate is compensated by the opposite energy dependence of the total E2 conversion coefficient [23]. As a result, the B(E2) value from the isomeric state is only weakly dependent on the transition energy, making it possible to have an estimate of the B(E2) strength that ranges from $6.2 \pm 0.3~e^2~{\rm fm}^4$ for $20~{\rm keV}$ to $4.4 \pm 0.2~e^2~{\rm fm}^4$ for $90~{\rm keV}$.

For the state decaying via the 685-keV γ ray to the 13/2⁻ level in ²¹⁷Bi, the most straightforward assumption from the measured decay constant is that it belongs to a second decay branch of the same isomer feeding the other states. The γ rays connecting the isomer to this level may not be observed due to their low energy. Given that the x rays observed are compatible with the internal conversion of the four transitions, the energy of these connecting transitions has to be below 90 keV. The most probable scenario is that the 685-keV transition has an M1 or E2 multipolarity, leading to a $15/2^$ or 17/2 assignment for the new state at 1429 keV. Such an assignment would imply at least two (or three if the spin is $15/2^-$) transitions connecting the $25/2^-$ isomer to the 1429-keV state. All such transitions will be well below 90 keV, of E2 or M1 character, and thus almost completely converted. The new states will be isomeric, and the combination of the 25/2 isomer half-life with these intermediate states should lead to the measured half-life of the 1429-keV level. This is compatible only with half-lives of the order of 100 ns or shorter for the intermediate states, which are inside the range expected from systematics in the decay sequence of a seniority isomer of this region [5]. If the 1429-keV state is fed by the $25/2^-$ seniority isomer, this branching ratio of 14(4)% must be considered when estimating $B(E2; 25/2^- \rightarrow$ $21/2^{-}$), resulting in a value which ranges from 5.6 ± 0.3 to $3.6 \pm 0.2 \ e^2 \, \text{fm}^4$. Nevertheless, one cannot exclude that a second isomer exists in ²¹⁷Bi and one possibility could be that the 1429-keV state itself is isomeric. An isomeric transition of

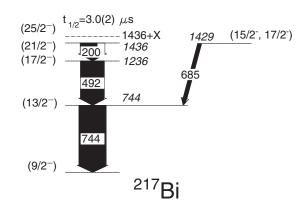


FIG. 5. Level scheme of ²¹⁷Bi deduced from the present data.

685 keV would be well compatible with an E3 multipolarity, thus giving a $19/2^+$ isomeric nature for the 1429-keV state. In view of systematics, it is highly unlikely that this state is a second isomer located at such low energy.

Figure 5 shows the level scheme of ²¹⁷Bi proposed from the present work. Considering the most likely scenario of a second decay branch from the seniority isomer, the two possible spin-parity assignments are indicated for the 1429-keV state, while the 19/2⁺ hypothesis is disregarded.

IV. THEORETICAL DISCUSSION

The new states observed in 217 Bi should be formed by coupling a valence proton in the $h_{9/2}$ orbital to the core-excited states in ²¹⁶Pb. Since the excited states up to 8⁺ in ²¹⁶Pb are understood within the seniority scheme $(2g_{9/2})^2$, the same structure is expected in ²¹⁷Bi with the yrast states forming the sequence $9/2^-$, $13/2^-$, $17/2^-$, $21/2^-$, and $25/2^-$. Figure 6 shows, for the $^{211-217}$ Bi nuclei, the results of shell-model calculations with the Kuo-Herling (KH) interaction [24] compared with the experimentally known level schemes. The valence space to describe these nuclei is constituted by the neutron shells $(g_{9/2}i_{11/2}d_{3/2}d_{5/2}g_{7/2}s_{1/2}j_{15/2})^8$ and by the proton shells $(h_{9/2}f_{7/2}i_{13/2}f_{5/2}p_{3/2}p_{1/2})^1$. A full calculation in this space is feasible, using state-of-the-art large-scale shell-model codes such as ANTOINE or NATHAN [25,26], only up to ²¹³Bi. For ^{215,217}Bi a reduction in the model space is needed. The calculations for these latter two nuclei were performed by restricting the proton valence space to $h_{9/2} f_{7/2}$ and by allowing up to six neutrons in the $i_{11/2}$ shell and up to four neutrons in the $j_{15/2}$ shell, while no restriction is given to the occupancy of the other neutron orbitals $g_{9/2}$, $d_{3/2}$, $d_{5/2}$, $g_{7/2}$, and $s_{1/2}$. The fact that the energy of the first excited state, $13/2^-$, is well reproduced shows that, even with this truncation, the pair scattering from the $\nu g_{9/2}$ orbital to the shells above is properly described.

The agreement between the calculated and experimental level energies is very good, being of the order of 100 keV (see Fig. 6). The analysis of the nuclear wave function confirms the above-mentioned simple scheme where the single proton in the $h_{9/2}$ orbital couples to the excited, seniority-two, neutron states of the corresponding even-even Pb isotopes.

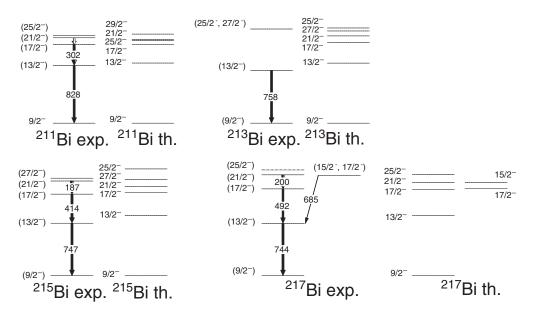


FIG. 6. Experimental and calculated partial level schemes for the odd-mass bismuth isotopes. The calculations were performed using the KH interaction. The ²¹⁷Bi level scheme results are from the present work. The experimental data are taken from Refs. [4,8–10].

Concerning the 1429-keV level in ²¹⁷Bi, for which we propose a $15/2^-$ or $17/2^-$ assignment, the shell-model calculations predict both a $15/2^-$ and a $17/2^-$ state close by in energy. A sensitive test of the nuclear wave function is given by the B(E2) values of the transitions depopulating the isomeric states. The half-lives of the 25/2⁻ levels in ²¹¹Bi and 217 Bi are $1.4 \pm 0.3 \ \mu s$ [1] and $3.0 \pm 0.2 \ \mu s$ (this work), which yield a reduced transition probability B(E2) of 8(2) $e^2 \, \text{fm}^4$ for ²¹¹Bi and from 6.2 ± 0.3 to 4.4 ± 0.2 $e^2 \, \text{fm}^4$ for ²¹⁷Bi, as discussed before. The 25/2⁻ seniority isomer is not known in ²¹³Bi, while in ²¹⁵Bi the presence of the 27/2 spin trap does not allow a seniority isomer. It is worth noting that the spin inversion in 215 Bi between $25/2^-$ and $27/2^$ is well reproduced by shell-model calculations, as shown in Fig. 6. We have calculated the B(E2) values using the same valence space and interaction already employed for the level energies and adopting the standard effective charges for this region: $e_{\pi}=1.5e$ and $e_{\nu}=0.8e$ [24]. The results are 92 and $1.0~e^2~{\rm fm}^4$ for $^{211}{\rm Bi}$ and $^{217}{\rm Bi}$, respectively. The discrepancy with the experimental results is large. What is most disturbing is that, while the experimental B(E2) values are close to each other, as happens for the corresponding B(E2) values in the core nuclei ^{210}Pb and ^{216}Pb , the theoretical B(E2)values, which for the lead cores were comparable, differ here by a factor of 100. As mentioned above, for the ²¹⁷Bi calculations a restricted shell-model space had to be used and this could be a possible cause of the large difference in the calculated B(E2) values. With the intent to further understand this behavior, we have applied to the Bi isotopes the same approach successfully adopted in Ref. [13], where effective three-body forces have been included. In the bismuth case, however, since there is a proton in the valence space, it is difficult to perform a diagonalization in a space which includes all the neutron (and proton) shells as in Ref. [13]. On the other hand, it was shown that the relevant renormalization for the quadrupole operator is provided by particle-hole excitations

across the $\Delta J=2$, $0\hbar\omega$ shells $vi_{13/2}-vg_{9/2}$ and $\pi h_{11/2}-\pi f_{7/2}$, partners in the quasi-SU(3) scheme [13]. They are responsible for quadrupole coherence [27] and their inclusion allows one to evaluate the possible effect of effective three-body forces.

For the bismuth isotopes we have then performed the calculations including only these relevant shells, plus the $\pi h_{9/2}$ orbital occupied by the unpaired proton, where the Kahana-Lee-Scott (KLS) interaction and the effective charges e_{π} ~ 1.5e and $e_v \sim 0.5e$ have been used. The calculated level energies are in agreement with the ones obtained with the full space once the paring matrix elements of the $vg_{9/2}$ shell are renormalized to reproduce the energy of the $13/2^+$ state. The B(E2) strengths calculated are 38 and 28 e^2 fm⁴ for ²¹¹Bi and ²¹⁷Bi, respectively. The disagreement between the measured and the calculated B(E2) values remains large for both ²¹¹Bi and ²¹⁷Bi. However, with the inclusion of effective three-body forces the calculated B(E2) values for ²¹¹Bi and ²¹⁷Bi become similar, as they are experimentally. The fact that the ratio between the 211 Bi and 217 BiB(E2) values is reproduced is significant since it shows that the inclusion of core excitations, equivalent to considering effective threebody forces, is restoring the symmetry in the B(E2) values relatively. What remains to be understood is the discrepancy of the absolute value, which in both nuclei is experimentally lower by a factor of 4–5. Explanations for this behavior are not straightforward and may be found when more refined shell-model calculations in such large spaces become possible or when more dedicated experiments are performed to look into this problem.

V. CONCLUSIONS

The present paper reports on the first results on the excited states in the neutron-rich nucleus ²¹⁷Bi. The study of this exotic isotope was made possible by the presence of isomeric states,

which allowed one to perform decay γ spectroscopy using a radioactive beam produced from the uranium fragmentation. Four transitions were assigned to the decay from an isomeric state with a half-life of $3.0 \pm 0.2 \mu s$. The expected decay branch from the seniority isomer was observed, but there is evidence for another decay branch, probably from the same isomer. The derived level scheme was compared with systematics from lighter isotopes, as well as state-of-the-art shell-model calculations. Whereas the level energies of ²¹⁷Bi as well as of the other lighter odd-even Bi isotopes are well reproduced, the same calculations fail completely to predict the experimental B(E2) values from the $25/2^-$ seniority isomers in ²¹¹Bi and ²¹⁷Bi. When effective three-body forces are included, the correct ratio between the experimental B(E2) values is restored but not the absolute value. These experiments on heavy exotic nuclei are still at the limits in terms of statistics and sensitivity and may gain a lot from the expected improvements of experimental setups and beam intensities. Finally, from the theoretical point in view, in order to overcome the present difficulties, developments of codes able to perform a diagonalization in the full valence space are mandatory.

ACKNOWLEDGMENTS

The work of the GSI accelerator staff is acknowledged. AG, MD, and EF acknowledge the support of Istituto Nazionale di Fisica Nucleare (INFN), Italy, and MICINN, Spain, through the AIC10-D-000568 action. AG has been partially supported by MICINN, Spain, and the Generalitat Valenciana, Spain, under Grants No. FPA2008-06419 and No. PROMETEO/2010/101. The support of the UK STFC and of the DFG(EXC 153) is also acknowledged.

- [1] M. Pfützner et al., Phys. Lett. B 444, 32 (1998).
- [2] N. Al-Dahan et al., Phys. Rev. C 80, 061302(R) (2009).
- [3] S. J. Steer et al., Phys. Rev. C 78, 061302(R) (2008).
- [4] G. J. Lane et al., Nucl. Phys. A 682, 71 (2001).
- [5] K. H. Maier et al., Z. Phys. A 332, 263 (1989).
- [6] M. M. Hindi, E. G. Adelberger, S. E. Kellogg, and T. Murakami, Phys. Rev. C 38, 1370 (1988).
- [7] J. D. Bowman, R. E. Eppley, and E. K. Hyde, Phys. Rev. C 25, 941 (1982).
- [8] V. G. Chumin et al., Z. Phys. A 358, 33 (1997).
- [9] L. Chen et al., Nucl. Phys. A 882, 71 (2012).
- [10] J. Kurpeta et al., Eur. Phys. J. A 18, 31 (2003).
- [11] H. DeWitte et al., Phys. Rev. C 87, 067303 (2013).
- [12] H. DeWitte et al., Phys. Rev. C 69, 044305 (2004).
- [13] A. Gottardo et al., Phys. Rev. Lett. 109, 162502 (2012).

- [14] G. Benzoni et al., Phys. Lett. B 715, 293 (2012).
- [15] A. I. Morales et al., Phys. Rev. C 89, 014324 (2014).
- [16] A. Gottardo et al., Phys. Lett. B 725, 292 (2013).
- [17] K. Rykaczewski et al., AIP Conf. Proc. 455, 581 (1998).
- [18] H. Geissel et al., Nucl. Instrum. Methods B 70, 286 (1992).
- [19] S. Pietri et al., Nucl. Instrum. Methods B 261, 1079 (2007).
- [20] P. H. Regan et al., Nucl. Phys. A 787, 491 (2007).
- [21] R. Kumar et al., Nucl. Instrum. Methods A 598, 754 (2009).
- [22] P. H. Regan et al., Int. J. Mod. Phys. E 17, 8 (2008).
- [23] T. Kibédi et al., Nucl. Instrum. Methods A 589, 202 (2008).
- [24] E. K. Warburton and B. A. Brown, Phys. Rev. C 43, 602 (1991).
- [25] E. Caurier and F. Nowacki, Acta Phys. Pol. B 30, 705 (1999).
- [26] E. Caurier, G. Martinez-Pinedo, F. Nowacki, A. Poves, J. Retamosa, and A. P. Zuker, Phys. Rev. C 59, 2033 (1999).
- [27] E. Caurier et al., Rev. Mod Phys. 77, 427 (2005).