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New us Isomers in the Neutron-rich 210Hg Nucleus

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New μ s isomers in the neutron-rich ²¹⁰Hg nucleus

A. Gottardo^{1,2}, J.J. Valiente-Dobón¹, G. Benzoni³, A. Gadea⁴, S. Lunardi^{2,5}, P. Boutachkov⁶, A.M. Bruce⁷, M. Górska⁶, J. Grebosz⁸, S. Pietri⁶, Zs. Podolyák⁹, M. Pfützner¹⁰, P.H. Regan⁹, H. Weick⁶, J. Alcántara Núñez¹¹, A. Gorska⁶, J. Grebosz⁶, S. Pietr⁶, Z.S. Podolya⁷, M. Prutzne^{1,6}, P.H. Regan⁷, H. Weick⁶, J. Alcantara Nunez^{1,1}, A. Algora⁴, N. Al-Dahan⁹, G. de Angelis¹, Y. Ayyad¹¹, N. Alkhomashi¹², P.R.P. Allegro¹³ D. Bazzacco⁵, J. Benlliure¹¹, M. Bowry⁹, A. Bracco^{3,14}, M. Bunce⁷, F. Camera^{3,14}, E. Casarejos¹⁵, M.L. Cortes⁶, F.C.L. Crespi³, A. Corsi^{3,14}, A.M. Denis Bacelar⁷, A.Y. Deo⁹, C. Domingo-Pardo⁶, M. Doncel¹⁶, Zs. Dombradi¹⁷, T. Engert⁶, K. Eppinger¹⁸, G.F. Farrelly⁹, F. Farinon⁶, E. Farnea⁵, H. Geissel⁶, J. Gerl⁶, N. Goel⁶, E. Gregor⁶, T. Habermann⁶, R. Hoischen^{6,19}, R. Janik²⁰, P. R. John^{2,5}, S. Klupp¹⁸, I. Kojouharov⁶, N. Kurz⁶, S.M. Lenzi^{2,5}, S. Leoni^{3,14}, S. Mandal²¹, R. Menegazzo⁵, D. Mengoni⁵, B. Million³, V. Modamio¹, A.I. Morales³, D.R. Napoli¹, F. Naqvi^{6,22}, R. Nicolini^{3,14}, C. Nuclése⁶, A. Den le le le ⁶, B. Millon³, V. Modamio¹, A.I. Morales³, D.R. Napoli¹, F. D. P. P. Let¹⁴, C. S. Leoni^{3,14}, C. Nociforo⁶, A. Prochazka⁶, W. Prokopowicz⁶, F. Recchia⁵, R.V. Ribas¹³, M.W. Reed⁹, D. Rudolph¹⁹, E. Sahin¹, H. Schaffner⁶, A. Sharma⁶, B. Sitar²⁰, D. Siwal²¹, K. Steiger¹⁸, P. Strmen²⁰, T.P.D. Swan⁹, I. Szarka²⁰, C.A. Ur⁵, P.M. Walker⁹, O. Wieland³, H-J. Wollersheim⁶. ¹Istituto Nazionale di Fisica Nucleare, Laboratori Nazionali di Legnaro, Legnaro, 35020, Italy ²Dipartimento di Fisica dellUniversità degli Studi di Padova, Padova, 35131, Italy ³Istituto Nazionale di Fisica Nucleare, Sezione di Milano, Milano, 20133, Italy ⁴Instituto de Física Corpuscular, CSIC-Universitat de València, València, E-46980, Spain ⁵Istituto Nazionale di Fisica Nucleare, Sezione di Padova, Padova, 35131, Italy ⁶GSI Helmholtzzentrum für Schwerionenforschung, Darmstadt, D-64291, Germany ⁷School of Computing, Engineering and Mathematics, University of Brighton, Brighton, BN2 4GJ, United Kingdom ⁸Niewodniczanski Institute of Nuclear Physics, Polish Academy of Science, Krakow, PL-31-342, Poland ⁹Department of Physics, University of Surrey, Guildford, GU2 7XH, United Kingdom ¹⁰Faculty of Physics, University of Warsaw, Warsaw, PL-00681, Poland ¹¹Universidade de Santiago de Compostela, Santiago de Compostela, E-175706, Spain ¹²KACST, Riyadh, 11442, Saudi Arabia ¹³Instituto de Fisica, Universidade de São Paulo, São Paulo, 05315-970, Brazil ¹⁴Dipartimento di Fisica dell'Università degli Studi di Milano, Milano, 20133, Italy ¹⁵EEI, Universidade de Vigo, Vigo, E-36310, Spain ¹⁶Grupo de Física Nuclear, Universidad de Salamanca, Salamanca, E-37008, Spain ¹⁷Institute of Nuclear Research of the Hungarian Academy of Sciences, Debrecen, H-4001, Hungary ¹⁸ Physik Department, Technische Universität München, Garching, D-85748, Germany ¹⁹Department of Physics, Lund University, Lund, S-22100, Sweden ²⁰Faculty of Mathematics and Physics, Comenius University, Bratislava, 84215, Slovakia ²¹Department of Physics and Astrophysics, University of Delhi, Delhi, 110007, India ²²Institut für Kernphysik, Universität zu Köln, Köln, D-50937, Germany

Abstract

Neutron-rich nuclei in the lead region, beyond N = 126, have been studied at the FRS-RISING setup at GSI, exploiting the fragmentation of a primary uranium beam. Two isomeric states have been identified in ²¹⁰Hg: the 8⁺ isomer expected from the seniority scheme in the $vg_{9/2}$ shell and a second one at low spin and low excitation energy. The decay strength of the 8⁺ isomer confirms the need of effective three-body forces in the case of neutron-rich lead isotopes. The other unexpected low-lying isomer has been tentatively assigned as a 3⁻ state, although this is in contrast with theoretical expectations.

Keywords:

Atomic nuclei are complex many-body systems with many degrees of freedom; nevertheless their spectral properties often show very regular features due to the symmetries of the nuclear hamiltonian. A remarkable example of this is offered by the occurrence of the seniority excitation scheme in spherical, semi-magic nuclei [1]. A deviation from this regular behaviour suggests a change in the underlying nuclear structure, as for example a sudden onset of deformation. The neutron-rich regions around double shell closures have been studied for light and medium-mass nuclei, using fission and deep-inelastic reactions. However, the neutron-rich region around ²⁰⁸Pb has not been thoroughly explored so far, due to its high mass and neutron richness. Pioneering work has been reported in Ref. [2, 3]. A deeper knowledge would be desirable, since ²⁰⁸Pb is a benchmark for the study of nuclear structure thanks to its double-shell closure character. For semi magic neutron-rich ²¹⁰⁻²¹⁶Pb isotopes, a standard seniority structure has been found, as it is expected from neutrons in the $2g_{9/2}$ shell [4]. A 8⁺ isomer was measured in each isotope (with $t_{1/2}$ in the range ~ 0.1-6 μ s), decaying via the 6⁺ \rightarrow 4⁺ \rightarrow 2⁺ \rightarrow 0⁺ yrast cascade, with the levels spaced with decreasing energies as the spin increases. The analysis of the transition rates from the isomeric states allowed one to asses the role played by the usually neglected effective three-body forces, arising from core excitations outside the valence space [4]. Generally, the seniority scheme provided by the coupling of the valence neutrons in the $g_{9/2}$ shell may also hold when few protons are added to the ²⁰⁸Pb core. In fact, they can act as spectators, while the angular momentum of the excited states is given by the coupling of the neutrons. This is indeed the case if two protons are added to the Z = 82 core: the resulting polonium isotopes show a $g_{9/2}$ seniority scheme [5, 6, 7], apart from ²¹⁴Po whose known level scheme is limited to low-spin states.

Similarly, when two protons are removed from the Z = 82 core, leading to mercury isotopes, one would expect to observe the same $g_{9/2}$ seniority scheme. The isotope ²⁰⁸Hg has indeed a 8⁺ isomer attributed to the maximally-aligned configuration $vg_{9/2}^2$ [8]. In this nucleus, the two proton holes are in the $s_{1/2}$ and $d_{3/2}$ orbits, the less bound ones below Z = 82. They appear to be inactive spectators with respect to the neutron valence space. A similar behaviour is, in principle, expected also in the more neutron-rich isotope ²¹⁰Hg, whose level structure was completely unknown up to now.

This Letter reports the first experimental study of excited states in ²¹⁰Hg providing evidence of two isomers in the μ s range. One is interpreted as the 8⁺ isomer expected from the $g_{9/2}$ seniority scheme, while the other one, for which we tentatively suggest a 3⁻ assignment, is located at an unexpected low energy, remaining therefore a challenge for future experiments and theoretical models.

Experimentally, nuclei around the ²⁰⁸Pb region, more

than two or three neutrons from stability, have been produced with relatively cold fragmentation reactions from a primary ²³⁸U beam. The results on ²¹⁰Hg have been obtained by exploiting the uniqueness of the FRS-RISING setup [9, 10, 11, 12] and the UNILAC-SIS accelerator facilities at GSI by using a 1 GeV A ²³⁸U beam at an intensity of around 1.5×10^9 ions/spill. The ~ 1 s spills were separated by ~ 2 s without beam. The beam impinged on a 2.5 g/cm² Be target followed by a 223 mg/cm² Nb stripper. The isotopes resulting from the fragmentation reaction were separated and identified with the double-stage magnetic spectrometer FRS [9]. The FRS allows one to discriminate the magnetic rigidities of the fragments with a resolution sufficient to distinguish the masses of adjacent isotopes even at the high masses of interest ($A \sim 210 - 220$). The identification in mass over charge ratio (A/q) is achieved through time of flight and focal-plane position measurements, while the atomic number is obtained from two ionization chambers placed in the final focal plane. These measurements are sufficient to provide a complete identification of the isotopes event by event. 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 so they could be implanted in a composite double-sided siliconstrip (DSSSD) detector system [12, 13]. The DSSSD detector system was surrounded by the RISING γ spectrometer [10, 11], with a full-energy γ -ray peak detection efficiency of 15% at 662 keV [10]. Further experimental details can be found in Ref. [4, 14].

Figure 1, upper panel, shows the γ -ray spectrum following the isomeric decays of ²¹⁰Hg. After an implantation event a delayed gate of 5 μ s length was used to detect isomers. In total, 936 ions were identified as ²¹⁰Hg. Three γ lines are clearly visible at 663, 643 and 553 keV. A forth line is also identified at 170 keV. The spectra obtained from the $\gamma\gamma$ coincidence data are also presented in Fig. 1. The background in these highly-selective spectra is almost non existent and, when two γ -transitions are not in coincidence (see for example the bottom spectrum in Fig. 1), the few counts are distributed randomly in the entire energy range, never exceeding one count per channel. If two lines are in coincidence, as it is the case for those at 553 and 643 keV, this is proven clearly by the concentration of even a small amount of counts at the expected channels (see second and third spectra of Fig. 1). By summing the spectra in coincidence with the 643and 553-keV lines, the transition at 170 keV is also evident, thereby strengthening its identification as the third member of a γ -ray cascade composed by the 170-, 553and 643-keV transitions. As stated above, no γ rays are



Figure 1: The upper panel shows the γ spectrum, time-gated ($\Delta t = 0.1 - 5.0 \ \mu s$) for the ²¹⁰Hg isotope. The spectra below show the $\gamma\gamma$ coincidences for the different γ rays.

observed in coincidence with the 663-keV line, despite its higher intensity with respect to the 170- and 553-keV transitions. The statistical significance of the absence of coincidences between the 663- and the 643-keV γ rays can be estimated with a binomial distribution, the use of a Poisson distribution being less justified due to the low statistics. The calculation is performed considering the absolute γ efficiency at 663 keV, estimated to be 12.7%, and the missing feeding of the 643-keV state, which results to be 33(8) counts. As a consequence, if the 663-keV γ ray feeds with such intensity the 2⁺ state, the probability of not observing any coincidence between the 643-keV and 663-keV transitions in cascade is only $1.1^{+2.3}_{-0.7}$ %. Therefore, the absence of coincidence between the 643- and the 663-keV γ rays is a clear indication that the large intensity of the 643-keV line cannot be attributed to the feeding from the 663keV γ ray.

Figure 2 shows the time distributions of the three most intense γ transitions assigned to ²¹⁰Hg together with the results of an exponential fit of the data. Within errors, the half lives of the 553- and 643-keV transitions are equal and, being in mutual coincidence, they are assigned to follow the decay of the same isomer. The 663-keV transition shows also a similar half life, but it is not in coincidence with the other γ -rays. Because of intensity reasons (see below), we will conclude that it belongs to the decay of another, low-lying isomer with a half-life similar to the one de-excited by the 170-, 553and 643-keV sequence.

Table 1 summarises the intensities of the γ rays as-



Figure 2: (Colour online) Time distributions and the exponential fit in red for the 553-, 643- and 663-keV transitions assigned to ²¹⁰Hg.

Table 1: Area, transition intensities corrected for efficiency and internal conversion and their half-life $t_{1/2}$ for ²¹⁰Hg. The line at 170 keV, with its low statistics, is in a too high background region to fit its time distribution.

E_{γ} (keV)	Area	Intensity (%)	$t_{1/2}$ (μs)
170	13 (7)	22 (12)	-
553	12 (4)	23 (8)	2(1)
643	45 (7)	100 (16)	2.0(4)
663	25 (6)	65 (13)	2.1(7)

signed to ²¹⁰Hg and the half-lives derived from their time distributions. The 170- and 553-keV lines have comparable intensities, while the 643- and 663-keV γ rays have much larger intensity. The combination of this experimental information with the expected systematics of neutron-rich even-even Hg nuclei [8] allows one to assign the three transitions in mutual coincidence of 170, 553, and 643 keV to the de-excitation of the 8^+ isomer, similarly to what is observed in ²⁰⁸Hg [8] or in the even-even Pb isotopes [4]. The three γ rays are then attributed to the $6^+ \rightarrow 4^+ \rightarrow 2^+ \rightarrow 0^+$ decay sequence. The $8^+ \rightarrow 6^+$ is not detected because of its low energy and, as discussed in Ref. [4, 8], it is assumed to be between 20 and 80 keV. In fact, the characteristic K_{α} X rays from mercury at 71 keV are compatible with their origin from the 170-keV transition only: therefore, it is inferred that the $8^+ \rightarrow 6^+$ energy must be below the 78 keV binding energy of the mercury K electrons, and hence the given upper limit. From systematics, a conservative lower limit of 20 keV is adopted.

Figure 3 shows the level scheme deduced for ²¹⁰Hg compared with the one of ²⁰⁸Hg [8]. The sequence $6^+ \rightarrow 4^+ \rightarrow 2^+ \rightarrow 0^+$ from the decay of the 8⁺ isomer follows closely the systematics of the $g_{9/2}$ seniority isomers and it is in good agreement, within 100 keV, with

state-of the-art shell-model calculations, shown in the same figure. Theoretical calculations were performed using the shell-model codes ANTOINE and NATHAN [15, 16], and the Kuo-Herling interaction (KH) [17], slightly modified as indicated in Ref. [8]. The experimental spectrum of ²⁰⁹Pb was used for the single-particle energies. The neutron valence space considered was $(g_{9/2}i_{11/2}j_{15/2})^4$ above the closed N = 126 core and the proton valence space was $(2d_{5/2}2d_{3/2}h_{11/2}s_{1/2})^{-2}$ below the Z = 82 shell closure (it is a two proton-hole space). Assuming that the decay curve of the $4^+ \rightarrow 2^+$ transition is due to the lifetime of the feeding 8^+ isomer, and with the aforementioned energy limits on the this level, the B(E2) from the isomeric state ranges from 5(2) to 6(2) $e^2 fm^4$. The value predicted by shell model calculations is 55 $e^2 f m^4$, using the effective charges $e_{\nu} = 1.0e$ and $e_{\pi} = 1.5e$ as in the case of ²⁰⁸Hg [8], where the agreement with the measured value was very good. In ²¹⁰Hg, on the contrary, the shell-model calculation overestimates the B(E2) by a factor nine, similarly to what observed in the analogous four-valenceneutrons ²¹²Pb [4]. This would require the use of effective charges for neutrons and protons much lower than the standard ones. The state dependence of the effective charges is most probably related to the neglect of $0\hbar\omega$ excitations across the 208Pb core and of the associated effective three-body components of the nuclear hamiltonian, as has been discussed extensively in Ref. [4]. In that publication it was shown how the particle-hole excitations, coming from the lead core, play a major role in determining the values of the effective charge and how this contribution cannot be renormalized via the use of a constant effective charge. Actually, the same mechanism is expected to take place in ²¹⁰Hg, because the neutrons and the protons occupy the same $\Delta \hbar \omega = 0$ relevant orbitals $(v0i_{13/2} - v1g9/2 \text{ and } \pi 0h_{11/2} - \pi 1f_{7/2})$ as ²¹²Pb. The proton-hole wave function is mainly in the $\pi d_{3/2}$ and $\pi s_{1/2}$ orbitals below Z=82: this implies a very weak contribution to the $\Delta J=2$ particle-hole jumps responsible for quadrupole coherence [4]. Exact calculations are not possible at the moment but the two proton holes are not expected to play a role in the structure of the seniority isomer. The isomer should thus have the same structure of the one of neutron-rich lead isotopes. The overestimation of the B(E2) in ²¹⁰Hg of roughly the same amount as in the isotone ²¹²Pb appears as a confirmation of the importance of effective three-body forces in this region.

Besides the seniority-scheme isomer, it is evident from the upper panel of Fig. 1 that the intensity of the $2^+ \rightarrow 0^+$ transition is four times larger than the intensity of the γ ray (553 keV) directly feeding the 2^+ . This requires the presence of another isomeric state whose decay feeds, for three quarters of its population, the 2^+ state. This must be due to the other isomer that, as seen experimentally, exists in ²¹⁰Hg. A single transition that could explain the intensity missing in the 2^+ level should have three times the intensity of the 553keV γ ray. Therefore, it should be clearly visible in the data, unless it is of low energy where internal conversion becomes dominant. No further transitions, besides the 170- and the 553-keV ones, are however observed in coincidence with the 643-keV line, pointing to a nonmeasurable transition connecting the other isomer to the 2^+ state. If an observational limit of five counts is set, this transition must be below 40 keV, 150 keV, 200 keV and 260 keV for E1, E2, M1, and E3 multipolarities, respectively. The fact that the quite intense delayed transition at 663 keV is not in coincidence with any other line could suggest that it connects directly the other isomer to the 0⁺ ground state. This scenario, illustrated in Fig. 3, is also compatible with a direct decay of the new isomer to the 2⁺ state through a strongly converted 20-keV transition, as required by the high intensity of the $2^+ \rightarrow 0^+$ transition. The proposed state at 663 keV, because of its decay to both the 0^+ ground state and to the 2^+ state, can assume only spins from 1 to 3. The low spin of this isomer is also supported by the analysis of the isomeric ratio, performed using the method described in Ref. [18].

The isomeric ratio is, for each isotope and each isomer, the ratio between the number of nuclei populated in their excited isomeric state and the total number of ions of that nuclide produced after a fragmentation reaction. The final value takes into account the loss of ions in excited states due to the time of flight in the FRS and the loss in RISING γ efficiency due to the prompt γ flash [18]. In the scenario proposed in Fig. 3 for ²¹⁰Hg, the values obtained are 0.11 (4) for the 8⁺ isomer and 0.6(1) for the isomer at 663 keV, supposing it feeds the 2^+ state. If only the yield of the 663 keV-line is used, then the isomeric ratio of this level would be 0.25(6). Therefore, in any case the relative population of the two isomers is suggesting that the spin of the 663-keV state is quite low, in particular lower than 8^+ , and that it is an yrast state [19]. In fact, the schematic sharp cutoff model [18] predicts an increasing population with decreasing spin.

The lifetime derived from the decay curve of the 663keV line (see Fig. 1) will help to discriminate among the possible spin-parities of the proposed low-spin isomer by deducing the transition probabilities for the different possible multipolarities involved. Table 2 shows the Weisskopf strengths of the various multipolarities



Figure 3: Experimental and calculated level schemes of 208,210 Hg. The 663-keV state in 210 Hg is tentatively identified. The 3⁻ state in 210 Hg, calculated via particle-vibration models, is thicker to distinguish it from the other levels calculated with the shell-model. The experimental data for 208 Hg are taken from Ref. [8].

for the proposed 20- and the 663-keV transitions. The resulting transition rates appear in most cases very much hindered, at variance with systematics, with the exception of the E3 case for the 663-keV energy and the E1 case for the 20-keV energy. In fact, E3 transitions are normally of the order of one to several W.u., while a suppression factor of 10^6 for E1 transitions is not uncommon [20]. Therefore, a 3^- assignment for the 663-keV state seems the most reasonable based on the reduced transition probabilities. This in fact would give a 663-keV E3 transition to the ground state and a 20-keV E1 transition to the 2⁺ level.

Other possibilities are discussed in the following. A 1⁺ assignment would give a 663-keV M1 transition hindered by a factor 10⁸, which is not realistic, even for an ℓ -forbidden M1. For example, in the one-proton hole ²⁰⁷Tl isotope, there is a 3/2⁺ state (configuration $(\pi d_{3/2})^{-1}$) decaying to the ground state 1/2⁺ (configuration $(\pi s_{1/2})^{-1}$), with a half life of 30 ps. This is a ℓ -forbidden M1 transition, of ~ 10⁻² W.u., 10⁶ times higher than in the present case. A 2⁺ assignment would give an E2 transition suppressed by a factor 10⁵, which is is against all the systematics in literature. Finally, the large hindrance, 10⁻¹⁰ required to justify an isomeric 1⁻ 663-keV state rules out also this possibility.

For the level scheme proposed for ²¹⁰Hg in Fig. 3, the analysis of the transition probabilities favours therefore a 3⁻ spin-parity assignment to the state at 663 keV. All the other scenarios are incompatible with the measured lifetimes and intensities. In summary, the experimental data, i.e. $\gamma\gamma$ coincidences, half-life and branching ratio, suggest a possible level scheme as the one shown in Fig. 3, where the 663-keV transition has most probably a 3⁻ character.

Table 2: Experimental reduced transition strengths $B(E,M\lambda)$ in Weisskopf units. The branching ratio is 0.43(9) and 0.57(9) for the 663and 20-keV transitions, respectively.

E_{γ} (keV)	E,Mλ	$B(E,M\lambda)$ (W.u)
663	E1	$3(1) \cdot 10^{-10}$
663	E2	$3(1) \cdot 10^{-5}$
663	E3	4(2)
663	M1	$4(1) \cdot 10^{-8}$
20	M1	$8(3) \cdot 10^{-6}$
20	E1	$1.5(6) \cdot 10^{-6}$

In the following theoretical predictions for the 663keV transition in ²¹⁰Hg will be discussed, pointing out the difficulty to explain the lower energy isomer. The shell model calculations, discussed previously and successful in reproducing the sequence of states populated by the 8⁺ isomer and the isomer lifetime (once the problem of neglecting effective three-body forces is considered), predict the second 0^+ at 1.1 MeV, the 1^+ state at 1.5 MeV, the 1⁻ level at 2.7 MeV and the second 2⁺ at 1.34 MeV (see Fig. 3), but cannot reliably predict the location of a 3⁻ state. This is because the present shell-model calculations do not allow core excitations and it is known that the 3⁻ state in the lead region is very fragmented [21], involving many particlehole excitations across the 208Pb core. As a consequence, shell model calculations do not predict any excited state below 1.1 MeV, apart from the first excited 2^+ state. By looking at the systematics of 3^- excitations in this region, one observes that the 3⁻ level is at 2.6 MeV in ²⁰⁸Pb and drops to 1.9 MeV in ²¹⁰Pb due to mixing between the collective ²⁰⁸Pb octupole phonon and the 3⁻ originating from the octupole coupling of the $\Delta J = \Delta \ell = 3$ neutron shells $g_{9/2} - j_{15/2}$ [21]. In more neutron-rich nuclei further reduction of the excitation energy is expected [21]. For example, in ²¹⁴Po, which has the same neutron number of ²¹⁰Hg and two protons above Z=82 shell closure (²¹⁰Hg has two protons below), the energy of the 3⁻ state is at around 1.3 MeV [22], the value which is predicted also in ²¹⁰Hg by particle-vibration coupling models [21], as shown in Fig. 3. These calculations are very reliable, since they reproduce quite accurately the energy of the 3⁻ states in the region. A 3⁻ state at only 663 keV is therefore in contrast with basic theoretical expectations.

In conclusion, the very exotic ²¹⁰Hg nucleus was investigated for the first time via isomer decay spectroscopy. Its production and study were made possible thanks to state-of-the-art improvements in experimental techniques and to the intense uranium beams provided by the accelerator complex at GSI. A sequence of positive parity states 0^+ , 2^+ , 4^+ , 6^+ is proposed to belong to the decay of the 8⁺ seniority isomer. This sequence, very similar to that of the neighbouring ²⁰⁸Hg, is well reproduced by shell-model calculations. The decay strength of the isomer is not well reproduced by shell-model calculations, confirming the important role played by effective three-body forces in this region [4]. The experimental data indicate also the presence of another isomer at 663 keV. From its lifetime it was tentatively assigned as a $J^{\pi} = 3^{-}$ state, which is not theoretically expected at such low energy by particlevibration coupling models. In fact, in almost spherical nuclei, not too far from the doubly-magic ²⁰⁸Pb, 3⁻ excitations are predicted at higher energy. Such a large drop of the 3⁻ excitation in ²¹⁰Hg, if proven by more sophisticated and high statistics experiments, will be a real challenge for present theoretical models: ad augusta per angusta. An important future experimental development will be to directly measure the mass of the isomeric state feeding the 2^+ state with storage rings [23].

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