EUROPEAN ORGANIZATION FOR NUCLEAR RESEARCH

Addendum to the ISOLDE and Neutron Time-of-Flight Committee

Simultaneous spectroscopy of γ - rays and conversion electrons: Systematic study of E0 transitions and intruder states in close vicinity of mid-shell point in odd-Au isotopes

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Abstract: We propose to perform the study of β decay of mass-separated samples of the ¹⁸⁵Hg isotope. Samples will be collected by the deposition of low-energy radioactive-ion beam on the rapidly quenched metallic tape of the TATRA system. Conversion electrons will be detected with windowless Si(Li) detector cooled with liquid nitrogen. Gamma rays will be detected with the BE2020 Broad Energy Germanium detector. Level scheme of the ¹⁸⁵Au isotope will be constructed and electric monopole transitions will be identified.

Requested shifts: 7 shifts, (in 1 run together with remaining 5 shifts)

1 Motivation for study of ¹⁸⁵Hg

The Au isotopes play a unique role in our understanding of shape coexistence in that strongly-deformed structures intrude to become the ground state at mid shell (¹⁸³Au) and to exhibit a classic "parabolic" trend in excitation energy [1]. Present addendum is a part of our focused experimental programme on the study of nuclear structure of odd-Au isotopes. The programme involves in-beam γ -ray and conversion electron studies (performed at University of Jyvaskyla and iThemba Labs) [2, 3], β -decay experiments (ISOLDE) [4, 5] and isomer spectroscopy [6].

Our previous achievements at ISOLDE were reported to the INTC in 2019 and we refer the committee to the INTC-SR-061 status report. In this report, we suggested a construction of the new detection system based on rapidly-quenched metals with longer and wider tape. This would allow to extend studies to isotopes with half lives of few seconds. However in 2019, Ministry of Education, Science, Research and Sport of Slovak Republic stopped completely the funding of our activities at ISOLDE. This decision, which had purely political nature, included also the payments of CERN and ISOLDE member fees. After massive campaign in the Slovak media, which was closely followed also by CERN management and created public pressure to authorities, the funding was returned at very end of 2019 and in early 2020. Due to these reasons, the group at the Institute of Physics, Slovak Academy of Sciences, could not construct the prototype of the system. The only system that is available in the moment is the original TATRA. However, the development of new system is still planned and it will be discussed with representatives of the new Slovak government.

Therefore, as a continuation of the program we propose to carry out dedicated study of the β decay of ¹⁸⁵Hg. It can be performed with existing setup and there are further reasons for this choice (it was our plan to study this decay anyway).

An interest in systematic study of the Au isotopes can be traced back to before the emergence of shape coexistence as a feature of heavy nuclei. Experiments at the UNISOR [7] and the ISOCELE facilities [8, 9] revealed a remarkable constancy in excitation energies for many low-lying excited states in the odd-Au isotopes. However, "intruder" states, i.e., "unexpected" states that appeared at low energy, were established in the most neutrondeficient Au isotopes accessible at the time [8, 9]. This led to the first review of shape coexistence in nuclei [10]. The structure of the neutron-deficient Au isotopes became clearer with the summary provided in a paper by Kortelahti *et al.* in 1988 [11]. It revealed a complex situation of four coexisting structures in ^{185,187}Au. Some details of the 185 Hg^{m,g} decay to 185 Au are summarized in an unpublished thesis [12]. It pointed to serious errors in an earlier study of the ¹⁸⁵Hg decay. Most critical was a failure to identify parity-changing E1 transitions, which led to wrong parity assignments for a series of excited states and thus to misinterpretation of the underlying nuclear structure. These errors occurred due to missing of doublets (or even higher multiplets) in measured spectra. Analysis of γ -gated electron spectra and application of a running gates technique solved this problem.

A very detailed investigation of the decays of ${}^{187}\text{Hg}^{m,g}$ to ${}^{187}\text{Au}$ followed [13]. These studies involved measurements of conversion electrons using γ -e⁻ coincidences and identification of electric monopole transitions (*E*0), which provide a model independent fingerprint of

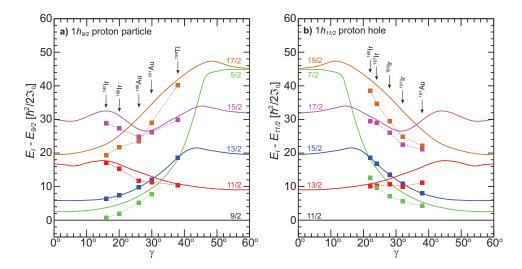


Figure 1: Comparison of calculations based on the PTRM approach, see the text for details, with experimental data. Energy of excited states is given as a function of the γ deformation parameter for **a**) $1h_{11/2}$ proton-hole and **b**) $1h_{9/2}$ proton-particle configurations.

shape coexistence. Decay schemes have been constructed incorporating 99% of the decay intensities assigned to the high-spin and low-spin decays. The detection limit was pushed down nearly to 0.1% if the intensity of the strongest γ ray. The γ -ray gated conversionelectron spectra permitted determination of 367 conversion coefficients. In total 9 E0 transitions were identified between both negative- and positive-parity states. Compared to that, the existing information on ¹⁸⁵Au is incomplete. Counterparts of only few E0 transitions that are known in ¹⁸⁷Au were observed. Knowledge of systematic evolutions of these coexisting structures will provide major constraints for nuclear models. Known E0 transitions were observed in the ¹⁸⁵Hg decay with intensities of 5-10% of the strongest γ ray (which is not assigned into the level scheme!). Therefore, the spectroscopy at least at the level of 1% of the strongest γ ray is demanding.

Other motivation for study of excited states ¹⁸⁵Au is the evolution of deformation as midshell point is approached. In general, deformation parameters, both axial and triaxial can be deduced from the spectra of excited states of the odd-mass nuclei. The odd particle or hole acts as a probe of the even-even core. The particle-core coupling models [14] suggest a very strong dependence of excitation energies of states with various spins. Fig. 2 gives a spectra of excited states associated with the $1h_{11/2}$ odd particle and $1h_{9/2}$ odd hole as a function of the triaxial deformation parameter (with fixed axial parameter). It is evident that the spectrum changes gradually from decoupled at $\gamma = 0^{\circ}$ to stronglycoupled structure at $\gamma = 0^{\circ}$, when passing through the triaxial plane. Comparison with the experimental data for several isotopes is also given. Different regions of the triaxial deformation have a typical signature. This allows to deduce the γ parameter from measured spectra of the odd-mass isotopes. This approach was successfully used in the study of ¹⁸⁷Au - see a comparison of the calculation with various deformation parameters given in Fig. 2. Present proposal aims to extension of this towards the lighter isotope.

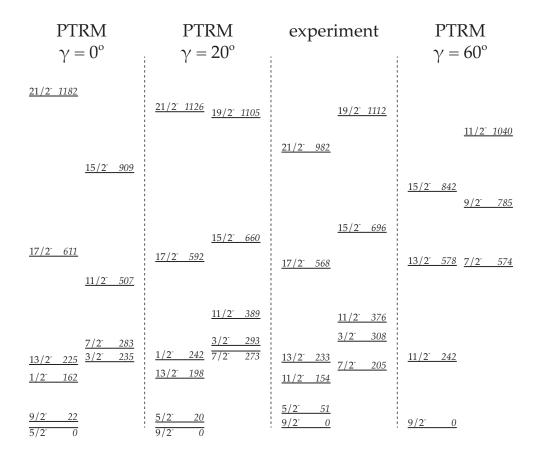


Figure 2: Excited states of the $1h_{9/2}$ proton-intruder configuration in ¹⁸⁷Au calculated for axially symmetric, prolate, oblate and triaxial ($\gamma = 20^{\circ}$) shape. For a comparison, experimental data is shown. Figure is adapted from [13].

Therefore, multipolarities of transitions need to be determined.

2 Detection system

The TATRA tape system [16] will be used for collection and transportation of samples of the ¹⁸⁵Hg isotope. The operation principle of the system is given in Fig. 3. The tape is made of rapidly quenched metal and it is designed at the Institute of Physics. To solve the problems with welding of the quenched material that complicated runs in 2014 and 2016, new tape with two layers was prepared and tested.

Conversion electrons will be detected with the 5 mm thick Si(Li) detector with 80 mm^2 surface. The detector is windowless, cooled with liquid nitrogen and it is housed in the retractable cryostat. It detects conversion electrons approximately up to 2.5 MeV with full width at half maximum of 1.6 keV. The TATRA system reaches vacuum below 10^{-7} mbar, which allows operation of such detector. The system is equipped with cryogenic module to ensure safe cooling and warming of the detector. Conversion electrons are detected both in singles and coincidence modes. It is important to note, that due to large density

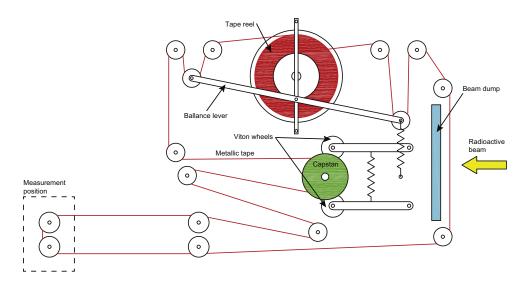


Figure 3: Operation principle of the TATRA tape transportation system. Figure is adopted from [?].

of excited states, the cryogenic Si(Li) detector is the only option for this study. Silicon detector cooled with, e.g., alcohol does not have sufficient resolution - see details in ??. A general complication related to studies of odd-Au isotopes is a large density of excited states at low energies, see e.g. [13]. This leads to the complexity of odd-mass decay schemes (several hundreds of γ rays), that renders the Rydberg-Ritz technique too ambiguous to be useful, unless γ -ray energies are measured to a precision better than 50 eV. Therefore, advanced coincidence analysis has to be performed, which includes, e.g, running gates technique. This makes the analysis procedure very complicated a leaves a risk of serious mistakes to be made, see [11]. Further, the use of coincidence spectroscopy to reliably sequence decay paths can be difficult or impossible because of isomerism (coincidence delay) occurring for some of the low-lying excited states. In the face of such challenges, we have developed and applied a dedicated experimental technique [15], which is based on novel Broad Energy Germanium (BEGe) detector, for these studies.

A BE2020 BEGe detector operated at ultra-high gain was successfully used to construct the level scheme of ^{181,183}Au [4, 5], which are nuclei with large densities of excited states at low energies. The advantage of the BEGe detector is not only excellent resolution, and nearly ideal gaussian peak shape, but also the ability to detect high-energy γ rays. Using this detector, γ -ray energies with a precision below 50 eV (in most cases even down to 10 eV) could be determined. To reach such precision, it is critical to operate the detector at ultra-high gains and also to ensure the stability of the electronics. With precisely determined γ -ray energies, the Rydberg-Ritz combination principle at the level of 30 eV precision could be used, which makes the process of complex level scheme construction much more simple. Properties of the BEGe detector, particularly the smooth background continuum (which helps normalisation and deconvolution), allowed the peaks of interest to be distinguished from other processes such as from the decay of daughter activities. This is very important since it simplifies the analysis and reduces the risk of misinterpreting observed transitions. It also provides additional information on the daughter isotopes, which can be analysed separately. Even with the precision of the BEGe detector, the γ - γ coincidence analysis cannot be omitted. Therefore the BE2020 BEGe detector will be used in a combination with coaxial germanium detectors.

3 Clarification of amount of requested beam time

We aim on the spectroscopy at the level 1% of the intensity of the strongest observed γ ray. Such sensitivity guarantees that expected E0 transitions between both negativeand positive-parity states will be identified and assigned to the level scheme. Our previous study of the ¹⁸³Hg decay, presented [4], was based on the data collected during approximately 2 days of measurement. Weakest assigned γ rays have intensity 6% of the strongest transition. Compared with study reported in [4], the γ -ray detection efficiency of the TATRA system was increased by installation of third large (90% relative efficiency) coaxial germanium detector. This increased γ - γ coincidence efficiency by a factor of approximately 3 and γ -electron coincidence efficiency by a factor of approximately 1.5. Another increase of the statistical quality of the data will come from increased activity of samples of ¹⁸⁵Hg. The data acquisition system of the TATRA has been optimised and presently it can accept count rates of approximately factor of 2 larger, then previously. The yield for the ¹⁸⁵Hg is not a limitation, since it is so high that the beam gate needs to be manipulated carefully, to keep the activity of the each sample and thus counting rate of detectors sufficiently low to not overload the electronics.

Therefore, within 4 days of beam time (i.e., twice more than it was used for the study of ¹⁸³Hg) a factor of 12 more γ - γ coincidences and a factor of 6 more γ -electron coincidences will be detected. This will allow to fulfil the above goal.

Since the TATRA system is transportable and needs to be delivered to CERN by ground means, the economy aspect of the experiment needs to carefully planned. We consider 4 days of measurements as acceptably long time that pays off the costs of transportation, installation with the final tuning of the system and uninstallation after the experiment. Study of ¹⁸⁹Hg could be proposed, within present document. However, we still have data from previous measurements and we prefer to finish their analysis. After this will completed, if needed, we will apply for dedicated experiment on the ¹⁸⁹Hg decay. Proposals to study ^{179,181}Hg decays will come as soon as the new high-vacuum tape system is ready, see the discussion above. Therefore we propose to add 7 shifts to already approved 5 and use 12 shifts to study ¹⁸⁵Hg decay.

Summary of requested shifts: 7

References

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Appendix

DESCRIPTION OF THE PROPOSED EXPERIMENT

The experimental setup comprises: (name the fixed-ISOLDE installations, as well as flexible elements of the experiment)

Part of the	Availability	Design and manufacturing
TATRA tape transportation sys-	\boxtimes Existing	\boxtimes To be used without any modification
tem (needs to be transported		
from Bratislava)		

HAZARDS GENERATED BY THE EXPERIMENT (if using fixed installation:) Hazards named in the document relevant for the fixed [COLLAPS, CRIS, ISOLTRAP, MINIBALL + only CD, MINIBALL + T-REX, NICOLE, SSP-GLM chamber, SSP-GHM chamber, or WITCH] installation.

Additional hazards:

Hazards	[Part 1 of experiment/ equipment]	[Part 2 of experiment/ equipment]	[Part 3 of experiment/ equipment]		
Thermodynamic and fluidic oquipment					
Pressure					
Vacuum	$10^{-7}\mathrm{mbar}$				
Temperature	100 K				
Heat transfer					
Thermal properties of materials					
Cryogenic fluid	[fluid], [pressure][Bar], [volume][l]				
Electrical and electromagnetic					
Electricity	$4~{\rm kV}, 1~{\mu}{\rm A}$ - bias of ger-				
	manium detectors				
Static electricity					
Magnetic field					
Batteries					
Capacitors					
Ionizing radiation	Ionizing radiation				
Target material [mate- rial]	molten Pb target				
Beam particle type (e,	$^{185}\mathrm{Hg}$				
p, ions, etc)					
Beam intensity	10^7 particles/s				
Beam energy	$30\mathrm{keV}$				

Cooling liquids	liquid nitrogen	
Gases	venting gas - nitrogen	
Calibration sources:		
• Open source		
Sealed source	\Box [ISO standard]	
Isotope	60 Co, 152 Eu, 241 Am - γ	
• ibotope	calibration sources	
• Activity	100 kBq	
Use of activated mate-		
rial:		
Description		
• Dose rate on contact	[dose][mSV]	
and in 10 cm distance		
Isotope		
Activity		
Non-ionizing radiatio	n	
Laser		
UV light		
Microwaves (300MHz-		
30 GHz)		
Radiofrequency (1-300		
MHz)		
Chemical		<u> </u>
Toxic	Pb bricks will be used	
TOXIC	for shielding of γ radi-	
	ation	
Harmful		
Harmful CMB (carcinogens		
CMR (carcinogens,		
CMR (carcinogens, mutagens and sub-		
CMR (carcinogens, mutagens and sub- stances toxic to repro-		
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction)		
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive		
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant		
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive	alcohol for cleaning of	
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant Flammable	alcohol for cleaning of vacuum chambers	
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant Flammable Oxidizing	alcohol for cleaning of vacuum chambers [chem. agent], [quant.]	
CMR(carcinogens, mutagensmutagensandstancestoxictorepro- duction)CorrosiveIrritantFlammableOxidizingExplosiveness	alcohol for cleaning of vacuum chambers [chem. agent], [quant.] [chem. agent], [quant.]	
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant Flammable Oxidizing Explosiveness Asphyxiant	alcohol for cleaning of vacuum chambers [chem. agent], [quant.] [chem. agent], [quant.] [chem. agent], [quant.]	
CMR(carcinogens, mutagensmutagensandstancestoxictorepro- duction)CorrosiveIrritantIrritantFlammableOxidizingExplosivenessAsphyxiantDangerous for the envi-	alcohol for cleaning of vacuum chambers [chem. agent], [quant.] [chem. agent], [quant.]	
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant Flammable Oxidizing Explosiveness Asphyxiant Dangerous for the envi- ronment	alcohol for cleaning of vacuum chambers [chem. agent], [quant.] [chem. agent], [quant.] [chem. agent], [quant.]	
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant Flammable Oxidizing Explosiveness Asphyxiant Dangerous for the envi- ronment Mechanical	alcohol for cleaning of vacuum chambers [chem. agent], [quant.] [chem. agent], [quant.] [chem. agent], [quant.] [chem. agent], [quant.]	
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant Flammable Oxidizing Explosiveness Asphyxiant Dangerous for the envi- ronment Mechanical Physical impact or me-	alcohol for cleaning of vacuum chambers [chem. agent], [quant.] [chem. agent], [quant.] [chem. agent], [quant.]	
CMR (carcinogens, mutagens and sub- stances toxic to repro- duction) Corrosive Irritant Flammable Oxidizing Explosiveness Asphyxiant Dangerous for the envi- ronment Mechanical	alcohol for cleaning of vacuum chambers [chem. agent], [quant.] [chem. agent], [quant.] [chem. agent], [quant.] [chem. agent], [quant.]	

Mechanical properties	[location]	
(Sharp, rough, slip-		
pery)		
Vibration	[location]	
Vehicles and Means of	[location]	
Transport		
Noise		
Frequency	[frequency],[Hz]	
Intensity		
Physical		
Confined spaces	[location]	
High workplaces	[location]	
Access to high work-	[location]	
places		
Obstructions in pas-	[location]	
sageways		
Manual handling	Vacuum chamber is	
	heavy and needs to be	
	manipulated carefully	
Poor ergonomics	[location]	

Hazard identification:

Average electrical power requirements (excluding fixed ISOLDE-installation mentioned above): [make a rough estimate of the total power consumption of the additional equipment used in the experiment]