

Precise Measurement of Decay Half Lives of n-Rich Iodine Isotopes after Radiochemical Separation

D. Banerjee¹, D. Kumar^{2,3}, T. Bhattacharjee^{2,3}, S. Basak^{2,3}, S. S. Alam^{2,3}

¹Radiochemistry Division (BARC), VECC, 1/AF, Bidhan Nagar, Kolkata-700064, INDIA,

²Variable Energy Cyclotron Centre, 1/AF Bidhan Nagar, Kolkata – 700 064

³Homi Bhabha National Institute, Mumbai, India, PIN - 400094

Introduction

The half-life of a radionuclide is an indispensable parameter both for fundamental nuclear physics and the application of any particular radionuclide. The measurement of half-life provides a direct route to determine the nuclear matrix element involved in the corresponding β -decay process. Again the half-life determines the application of a radionuclide both in nuclear physics (as standard source for calibration of detector) and in nuclear medicine. In general, a radionuclide with a long half-life (~few years) is preferred to be used for instrument calibration and a radionuclide with intermediate half-life (few hours) is preferred to be used in nuclear diagnostics purpose.

Hence a precise database of half-life is always desirable and the same for radioisotopes having several application is significantly important. Iodine (I)-isotopes have several such application, viz., ¹²³⁻¹²⁵I & ¹³¹I are used in medicine and biology, ¹²⁵I is important in nuclear reactor physics as its decay product ¹³⁵Xe has a very high thermal n-absorption cross section etc., to name a few. Present work aims at the half-life measurement of n-rich I-isotopes (¹³⁰⁻¹³⁵I) as these isotopes are difficult to populate by fusion-evaporation reaction and most viable route is through nuclear fission. As fission reaction produces a large number of fission products involving a wide range of elements, it is always recommended to radio chemically separate the element of interest. A clean radiochemical separation of I-isotopes from the rest of the fission products within a limited time period considering the shortest half-life of ¹³⁵I (52.5m) is the major challenge of the present work. Although there exists the half-lives of the above n-rich I-isotopes in the literature [1], there are several values reported for ¹³⁰I [2-5]. For other I-isotopes also, it is worth performing a precise

measurement of half-life with radio chemically separated I-isotopes. The production of n-rich I-isotopes by α -induced fission of ²³⁸U and measurement of decay half-lives after radiochemical separation has been attempted in the present work.

Experiment

The α -irradiation of ²³⁸U target was performed in K=130 AVF Cyclotron at VECC with 40MeV α -beam. The U-target was electro-deposited on 1mil Al-backing and thickness was kept between 500-600 $\mu\text{g}/\text{cm}^2$. Two irradiations with different duration were performed. One irradiation was done for 1h to attempt shorter-lived I-isotopes and other was performed for 40h to address the longer-lived ones.

The irradiated target and Al-catcher foils were dissolved in sodium hydroxide solution and I-tracer was added to the reaction mixture. All iodide was oxidized to Iodine and extracted in CCl₄ medium. The Iodine was then back-extracted from organic phase in the form of Iodide and used for subsequent counting with HPGe detector. Both 50% HPGe and Compton-suppressed Clover-HPGe detector was used for counting the sample. Here, data from 50% HPGe detector are presented and the data from Clover detector will be presented during the conference if the paper is accepted for presentation.

Results and Discussion

The γ -ray spectrum obtained after radiochemical separation of I-isotopes from the rest of the fission products is shown in Fig. 1. The figure represents the γ -spectrum acquired immediately after the radiochemical separation performed for short-irradiation of 1h. All the peaks could be assigned only due to the decay of I-isotopes to the corresponding Xe-isotopes.

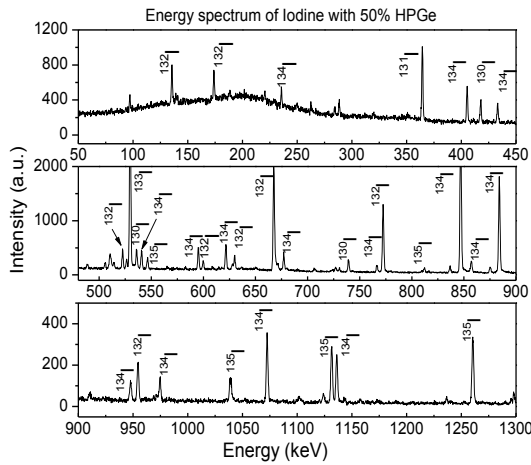


Fig 1: γ -ray spectrum taken with 50% HPGe detector after radiochemical separation of I-isotopes from other fission products

From the above spectrum, it may be realized that the radiochemical separation of I-isotopes was very clean and no other contamination peak was observed. The decay plots for some of the I-isotopes are shown in Fig. 2

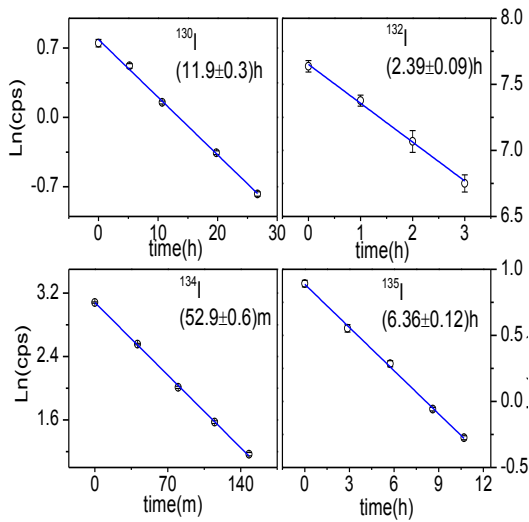


Fig. 2: Decay plots of separated I-isotopes

The decay half-lives of ¹³⁰I and ¹³²⁻¹³⁵I isotopes were measured. Only the measurement for ¹³¹I (8d) could not be given in the abstract as data for sufficient duration could not be accumulated till date. However, same will be presented during

the conference along with measurement for all I-isotopes with Clover HPGe detector as already mentioned. The half-lives obtained from the present measurement are tabulated in Table 1:

I-isotope	Half life	
	This work	Lit.
¹³⁰ I	11.9(3)h	12.36(1)h
¹³¹ I	-	8.0252(6)d
¹³² I	2.39(9)h	2.29(1)h
¹³³ I	20.8(1)h	20.83(8)h
¹³⁴ I	52.9(6)m	52.5(2)m
¹³⁵ I	6.36(12)h	6.58(3)h

The half-lives obtained from the present measurement agree well with the existing literature.

Conclusions

From the present work, it may be realized that the radiochemical separation of n-rich I-isotopes could be done successfully from the rest of the fission products. The γ -ray spectrum shows only the peaks originating from the decay of I-isotopes only. A precise measurement of half-lives could also be performed.

Acknowledgements:

The authors sincerely acknowledge Shri Ashim Kumar Biswas, RCD, VECC for his active participation in acquiring spectra and necessary help in data analysis. The effort of cyclotron staffs is acknowledged for providing the high quality beam. DB sincerely acknowledge Dr. R. Acharya, Head, NA&ACS, RCD, BARC and Dr. P. K. Pujari, AD, RC&I Group, BARC for their kind support for the work.

References

- [1] <https://www.nndc.bnl.gov/>
- [2] P. K. Hopke et al., Phys. Rev. C **8**, 745 (1973).
- [3] G. Andersson, G. Rudstam, and G. Sorensen, Ark. Fys. **28**, 37 (1965).
- [4] P. Aagaard et al., J. Inorg. Nucl. Chem. **5**, 105 (1957)
- [5] J. J. Livingood and G. T. Seaborg, Phys. Rev. **54**, 775 (1958).