

Cluster Radioactivity Study of Pt Isotopes

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Introduction

The cluster radioactivity is a spontaneous emission of clusters of nucleons whose mass is heavier than alpha particles and lighter than the lightest fission fragments. It was first observed by Sandulescu et al. in 1980 [1]. After few years, Rose and Jones first time experimentally observed the cluster decay ¹⁴C from ²²³Ra in 1984 [2]. Till now so many cluster decay has been verified with the emission of ¹⁴C, ^{18,20}O, ²³F, ^{22,24-26}Ne, ²⁸⁻³⁰Mg from heavy mass nuclei. So many theoretical models has been proposed to examine the cluster radioactivity [3,4,5] leading to the stable daughter nuclei around ²⁰⁸Pb.

The cluster decay of platinum element is not yet calculated experimentally. However, the authors of [6,7] give some productive information about the decay modes of Pt isotopes. Here we re-examine the decay modes of Pt isotopes within an axially deformed relativistic mean field (RMF) theory with NL3 force parameter set. The Q-value is obtained from the binding energies of the nuclei. With this same Q-value we study the α-decay as well as cluster decay half-life of Pt isotopes using the Viola-Seaborg [8] and Universal formula of Poenaru [9] et al.

Theoretical Formalism

The relativistic Lagrangian density for a nucleon-meson system is [10,11],

$$\begin{aligned}
 L = & \bar{\psi}_i(i\gamma^\mu \partial_\mu - M)\psi_i + \frac{1}{2}\partial_\mu \sigma \partial^\mu \sigma + \frac{1}{2}m_\sigma^2 \sigma^2 - g_\sigma \bar{\psi}_i \sigma \psi_i \\
 & - \frac{1}{4}\Omega_{\mu\nu} \Omega^{\mu\nu} \\
 & + \frac{1}{2}m_\omega^2 \vartheta_\mu \vartheta^\mu - g_\omega \bar{\psi}_i \gamma^\mu \psi_i \vartheta_\mu - \frac{1}{4}\vec{B}_{\mu\nu} \vec{B}^{\mu\nu} + \frac{1}{2}m_\rho^2 \vec{R}_\mu \vec{R}^\mu \\
 & - g_\rho \bar{\psi}_i \gamma^\mu \vec{\tau} \psi_i \cdot \vec{R}^\mu \\
 & \dots\dots\dots (1)
 \end{aligned}$$

All the terms have their usual meaning. The Q-value is calculated from the binding energies of parent nuclei, daughter nuclei and emitted nuclei as;

$$Q = M(A, Z) - M(A_1, Z_1) - M(A_2, Z_2) \dots\dots\dots(2)$$

Where $M(A, Z), M(A_1, Z_1), M(A_2, Z_2)$ are the atomic masses of parent, daughter and emitted nuclei respectively. The possibility to have a cluster decay method is that the decay energy of the Q-value should be greater than zero.

The expression for the α-decay half life of Viola-Seaborg is given by,

$$\log_{10} T_\alpha = \frac{aZ-b}{\sqrt{Q_\alpha}} - (cZ + d) + h_{log} \dots\dots\dots(3)$$

The details are found in Ref. [8]. The Q value and half lives for the emission of various clusters from the $Pt^{166-172}$ parent nucleus are tabulated in Table I. The half life calculations are also calculated by using Universal formula [9] for the cluster decay is given as,

$$\log_{10} T_{1/2}(s) = -\log_{10} P - \log_{10} S + [\log_{10}(h\nu) - \log_{10} \nu] \dots\dots\dots(4)$$

Where ν is a constant and S is the preformation probability of the cluster at the nuclear surface which depends only on the mass number of the emitted cluster.

Conclusion

In conclusion, the alpha decay, ⁸Be, ¹²C, ¹⁶O, ²⁰Ne, ²⁴Mg decay half lives are calculated from neutron-rich platinum isotopes by using RMF formalism. The calculated half-lives of alpha and other characteristics pertaining to possible cluster emissions with the Q-values obtained from RMF model have been computed and tabulated. The present study of the exotic decays of Pt isotopes may be helpful for future experiments.

Table –I : Cluster decay half-lives of Pt isotopes

Parent Nuclei	Binding Energy	Daughter Nuclei	Binding Energy	Emitted Cluster	Binding Energy	Q-Value (MeV)	T1/2 (V-S)	T1/2 (Univ.)	Expt.
¹⁶⁶ Pt	1289.133	¹⁶² Os	1267.051	⁴ He	28.14	6.058	-0.483	0.863	
¹⁶⁶ Pt		¹⁵⁸ W	1244.429	⁸ Be	51.507	6.803		55.23	
¹⁶⁶ Pt		¹⁵⁴ Hf	1221.997	¹² C	90.604	23.468		21.652	
¹⁶⁶ Pt		¹⁵⁰ Yb	1197.044	¹⁶ O	129.223	37.134		20.089	
		¹⁴⁶ Er	1171.203	²⁰ Ne	156.243	38.313		42.063	
		¹⁴² Dy	1145.168	²⁴ Mg	194.565	50.6		39.885	
¹⁶⁸ Pt	1310.482	¹⁶⁴ Os	1288.635	⁴ He	28.14	6.293	-1.410	-0.115	
¹⁶⁸ Pt		¹⁶⁰ W	1265.982	⁸ Be	51.507	7.007		-2.467	
¹⁶⁸ Pt		¹⁵⁶ Hf	1244.074	¹² C	90.604	24.196		19.808	
¹⁶⁸ Pt		¹⁵² Yb	1219.833	¹⁶ O	129.223	38.574		17.559	
¹⁶⁸ Pt		¹⁴⁸ Er	1194.571	²⁰ Ne	156.243	40.332		37.490	
		¹⁴⁴ Dy	1168.039	²⁴ Mg	194.565	52.122		37.006	
¹⁷⁰ Pt	1330.731	¹⁶⁶ Os	1309.192	⁴ He	28.14	6.601	-2.550	-1.303	-2.22
¹⁷⁰ Pt		¹⁶² W	1286.752	⁸ Be	51.507	7.528		46.599	
¹⁷⁰ Pt		¹⁵⁸ Hf	1263.332	¹² C	90.604	23.205		22.104	
¹⁷⁰ Pt		¹⁵⁴ Yb	1239.339	¹⁶ O	129.223	37.831		18.660	
		¹⁵⁰ Er	1216.469	²⁰ Ne	156.243	41.981		34.006	
		¹⁴⁶ Dy	1190.306	²⁴ Mg	194.565	54.14		33.456	
¹⁷² Pt	1350.163	¹⁶⁸ Os	1328.801	⁴ He	28.14	6.778	-3.169	-1.964	-0.96
¹⁷² Pt		¹⁶⁴ W	1306.485	⁸ Be	51.507	6.829		52.769	
¹⁷² Pt		¹⁶⁰ Hf	1283.130	¹² C	90.604	23.571		21.100	
¹⁷² Pt		¹⁵⁶ Yb	1259.158	¹⁶ O	129.223	38.218		17.900	
		¹⁵² Er	1234.665	²⁰ Ne	156.243	40.745		36.349	
		¹⁴⁸ Dy	1211.765	²⁰ Ne	194.565	56.167		30.119	

This work is supported by SERB, DST, Govt. of India, vi-de Grant No-SR/FTP/PS-106/2013.

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