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Electrochemical dissolution of the neutron irradiated ^{193}Ir metal powder

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Due to its nuclear characteristics, the Auger electron emitter isomer $^{195\text{m}}\text{Pt}$ ($T_{1/2} = 4.02$ d) is a promising radionuclide for radioimmunotherapy. We performed a test experiment to produce the isomer $^{195\text{m}}\text{Pt}$ by double neutron capture by the ^{193}Ir target nucleus with consequent population of $^{195\text{m}}\text{Pt}$ through β^- decay, and provided theoretical estimates for the $^{195\text{m}}\text{Pt}$ yield [1]. One of the advantages of the proposed approach is the possibility to isolate $^{195\text{m}}\text{Pt}$ from irradiated Ir target (in the form of metal powder) using radiochemical methods. It is known that iridium is a very hard, refractory, difficult to dissolve transition metal of the platinum group. Currently used methods of dissolving iridium do not meet the requirements for isolation of therapeutic radionuclides. We utilized anodic corrosion using alternating currents and solutions of hydrochloric and nitric acids that are used as electrolytes to study dissolution of Ir metal. We determined optimum conditions for electro dissolution of irradiated ^{193}Ir metal powder. In this work we also described and discussed details of the procedure.

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[1] S.A.Karamian, Aksenov N.V., Albin Yu.V., Belov A.G., Bozhikov G.A., Dmitriev S.N., Starodub G.A. Methods for production of $^{195\text{m}}\text{Pt}$ isomer. Bulletin of the Russian Academy of Sciences: Physics. 2014, in press; JINR Preprint P15-2013-84; Dubna, 2013.

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