

Production of an alpha-emitting radionuclide At-211 for medical use at JAEA tandem accelerator

タンデム加速器施設での核医学利用 α 放射性核種 At-211 生成

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In general, the ^{211}At nuclide, a prospective candidate for targeted alpha radiotherapy, has been produced through the $^{208}\text{Bi}(^4\text{He},2n)^{211}\text{At}$ reaction. In contrast, our project is focused on the production in the $^{208}\text{Bi}(^7\text{Li},2n)^{211}\text{At}$ reaction using the JAEA tandem accelerator [1,2]. This enables us to supply ^{211}At in a $^{211}\text{Rn}/^{211}\text{At}$ generator system. The daughter ^{211}At of 7.2 h in half-life is generated through EC decay of the parent ^{211}Rn of 14.7 h, expanding time-frame for transportation and use of ^{211}At . In this project, chemical procedures based on dry- and wet-chemistry have been studied to develop the $^{211}\text{Rn}/^{211}\text{At}$ generator system. In addition, research subjects relating to the development of the $^{211}\text{Rn}/^{211}\text{At}$ generator system, namely, production of astatine and iodine radioisotopes [3, 4], astatine chemistry [5, 6, 7] as well as the analytical method of ^{211}At using an alpha-scintillation camera and thin-layer chromatography [8] have been studied using the JAEA tandem accelerator. In the presentation, some experimental results in the project, e.g., production of astatine and iodine radioisotopes [2, 3, 4], the chemical procedure based on dry-chemistry for the $^{211}\text{Rn}/^{211}\text{At}$ generator system, and astatine chemistry [5, 6], will be presented.

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Primary author: Dr NISHINAKA, Ichiro (Tokai Quantum Beam Science Center, Takasaki Advanced Radiation Research Institute, National Institutes for Quantum Science and Technology)

Presenter: Dr NISHINAKA, Ichiro (Tokai Quantum Beam Science Center, Takasaki Advanced Radiation Research Institute, National Institutes for Quantum Science and Technology)

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