

Medically Important Radionuclides Obtained by Alpha Particle Irradiation

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Some radionuclides used in nuclear medicine have unique properties. These include the α -emitter ^{211}Ac and the β -emitter ^{61}Cu used in therapy, and the PET radionuclide ^{43}Sc used in PET diagnostics. Unfortunately, they can be effectively produced only on α particle-accelerating cyclotrons in α, n , and α, p reactions. ^{211}At seems to be the most promising candidate for targeted α -radiotherapy because its 7.2 h half-life assures sufficient time for its transportation, synthetic chemistry, multistep labeling, quality control, and clinical application without the problems caused by the relatively long-lived daughters emitting the α -particle. Recently, significant interest in ^{44}Sc ($T_{1/2} = 3.87$ h) as a tracer for PET imaging has been observed. Unfortunately, the co-emission by ^{44}Sc of high-energy γ rays ($E_{\gamma} = 1157$ keV, 99%) causes a dangerous increase in the radiation dose to the patients and clinical staff. Also, the co-production of longer-lived ^{44m}Sc ($T_{1/2} = 58.6$ h) increases the radiation dose for patients. However, it is possible to produce another radionuclide of scandium, ^{43}Sc which has properties similar to ^{44}Sc but is characterized by much lower energy of the concurrent gamma emissions. One of the most promising ways of production of ^{43}Sc are $^{40}\text{Ca}(\alpha, p)^{43}\text{Sc}$ and $^{40}\text{Ca}(\alpha, n)^{43}\text{Ti} \rightarrow ^{43}\text{Sc}$ nuclear reactions on natural calcium carbonate running simultaneously during the bombardment of targets made of natural calcium carbonate.

Unfortunately, only few cyclotrons in the world are currently able to accelerate α -beams with adequate for ^{211}At production energy and intensity consequently, the availability of ^{211}At and ^{43}Sc is limited to a few nuclear medical centers. In Europe, ^{211}At was produced only in France and Denmark. We believe that the use of the p- ^{11}B fusion to produce a flux of alpha particles as proposed in the PROBONO project can solve the problem with the availability of these radionuclides and radically change the availability of radiopharmaceuticals, especially for targeted alpha therapy.

The lecture will present the latest achievements in diagnostics and radionuclide therapy. Next, the methods of obtaining medical radionuclides and the production of radiopharmaceuticals will be presented, with particular emphasis on the radiopharmaceuticals based on ^{43}Sc , ^{211}At , and ^{61}Cu radionuclides.