Preparation and quality control of [18F]NaF for clinical application

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Abstract

[¹⁸F]NaF is used for skeletal imaging. While some ^{99m}Tc-labeled pharmaceuticals such as [^{99m}Tc]methylene diphosphonate also have high affinity for bones and widely used, [¹⁸F]NaF is considered to be more preferable agent in some viewpoints. One benefit of using [¹⁸F]NaF is rapid blood clearance which serves shorter study time for patient convenience. [¹⁸F]NaF is a simple compound and easily prepared only by eluting ¹⁸F trapped on an anion exchange column (QMA) with normal saline or NaHCO₃ solution through a 0.20 μm membrane filter. However, [¹⁸F]NaF injection produced by this way is subject to the quality of ¹⁸O enriched water that is irradiated to produce ¹⁸F, and the product is likely accompanied by some impurities such as vanadium-48 (⁴⁸V), a radionuclide with half-life of 15.97 days derived from irradiation of titanium target chamber. To use [¹⁸F]NaF for clinical purpose, it is important to assure the quality. In this paper, content of impurities in samples taken at some points of [¹⁸F]NaF preparation is analyzed by using PIXE method and pure-Ge semiconductor detector to find optimum conditions (ion form of QMA, kind of eluent and its volume) for preparing [¹⁸F]NaF. Contamination is shown least in [¹⁸F]NaF eluted with 2 mL of 0.4% NaHCO₃ solution. This suggests that inorganic nuclides such as ⁴⁸V are oxidized in aqueous solution and held trapped on QMA when ¹⁸F is eluted with diluted NaHCO₃.