

Internal Dose Estimation for Radiopharmaceuticals labelled with Accelerator Produced Technetium-99m

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INTRODUCTION

^{99m}Tc is the radionuclide most widely used in diagnostic nuclear medicine. It is available from ⁹⁹Mo/^{99m}Tc generators, where ⁹⁹Mo is obtained by nuclear-fission reaction on HEU-WP (²³⁵U) material in nuclear reactors. An alternative production route which can fulfill the shortage of this isotope in case of reactors shutdown exploits proton-irradiation of ¹⁰⁰Mo-enriched target [e.g. ¹⁰⁰Mo(p,2n)^{99m}Tc]. The TECHN_OSP/LARAMED research project at LNL-INFN has yet experimentally proved that, as expected from theoretical analyses [1], the accelerator-produced ^{99m}Tc (AP-^{99m}Tc) contains small quantities of several other technetium radioisotopes (⁹³Tc, ^{93m}Tc, ⁹⁴Tc, ^{94m}Tc, ⁹⁵Tc, ^{95m}Tc, ⁹⁶Tc and ^{97m}Tc) [2].

The aim of this work was to estimate the total contribution of technetium radioisotopes to the patient radiation absorbed dose after administration of four radiopharmaceuticals prepared with AP-^{99m}Tc, produced by irradiation of an Isoflex Mo target enriched with ¹⁰⁰Mo (99.05 %) with a proton beam of 15.7 MeV, accelerated by a PETtrace GE cyclotron.

METHODS

Four radiopharmaceuticals in use at present in the nuclear medicine clinic have been studied: Pertechnetate (used in clinical diagnostic of thyroid function and morphology), Sestamibi (widely used on cardiac scans for diagnosis of heart disease), hexamethylpropyleneamine oxime (HMPAO) (used as tracers of brain function) and disodium etidronate (HEDP) (a phosphonate commonly used for defining bone metastasis in cancer patients). The biokinetics models reported by the International Commission on Radiological Protection (ICRP) for each radiopharmaceutical [3-4] were used to determine the main source organs and to calculate the number of disintegrations that have occurred in each source organ (N_{source}), for each Tc-radioisotope found in the AP-^{99m}Tc. Then equivalent dose in the main organs of the adult male phantom was calculated for each Tc-radioisotope with

OLINDA/EXM software version 1.1 and 2.0 [5]. Finally the total effective dose produced by the AP-^{99m}Tc solution was calculated at 5 different times after the end of bombardment (EOB) and compared with the effective dose produced by the Generator-Produced ^{99m}Tc (GP-^{99m}Tc).

RESULTS

The amounts of other technetium radioisotopes in the AP-^{99m}Tc-radiopharmaceuticals produce a low (below 10%) increase of the effective doses with respect to the same radiopharmaceuticals marked with GP-^{99m}Tc, up to 12 h after EOB. ⁹⁶Tc is the radioisotope with higher contribution to the effective dose, followed by ⁹⁵Tc and ⁹⁴Tc, while other Tc-radioisotopes contribution is quite low, being at least 3 orders of magnitude smaller than ^{99m}Tc contribution.

CONCLUSIONS

The effective dose increase remains inside the 10 % limit from 6 to 12 h after EOB, for all the studied radiopharmaceuticals labelled with AP-^{99m}Tc. The percent of dose increase is different for each radiopharmaceutical, because the biological half-life in each source organ differs among compound. Therefore dose calculation must be done for each radiopharmaceutical in order to determine the range of time after the EOB in which the Tc-radiopharmaceutical can be administered to remain under the 10 % limit of dose increase. More details about this study will be published in reference [6].

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