pharmaceuticals

MDPI

Supporting information

Supporting information: Fully automated GMP compliant synthesis of ^{18}F-FE-PE2I

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Chromatograms and data from stabillity studies of OTsE-PE2I

Stabillity studies were conducted on the precursor (OTsE-PE2I) and reference (FE-PE2I) by adding solutions of the respective compounds to dried eluates from the QMA. The same experiments were performed for elutions using K_2CO_3/K_{222}, Et_4NHCO_3 and $Bu_4NH_2PO_4$ Some examples of the analytical HPLC chromatograms can be seen in Figure S1.

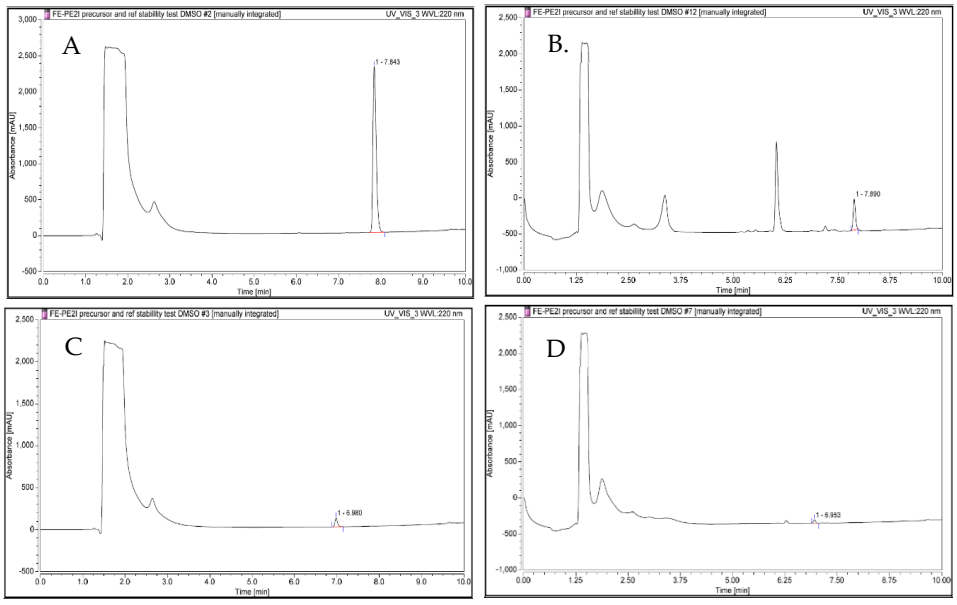

Figure S1. Examples of anlytical HPLC chromatograms to evaluate the stability of the tosylate precursor (OTsE-PE2I) and reference (FE-PE2I) dissolved in DMSO at 120 °C added to dried QMA eluates using $Bu_4NH_2PO_4$ (20 mM, 1 mL) for elution. The integrated peaks correspond to the identfied respective compound and remaining concentration at later time points (2-15 min) where calculated as a percentage of t = 0 min. The chromatograms display OTsE-PE2I dissolved in DMSO at t = 0 min (A.) and t = 10 min (B.) as well as FE-PE2I at t = 0 min (C.) and t = 2 min (D.).

Synthesis reports for synthesis using the K_2CO_3/K_{222} and $Bu_4NH_2PO_4$-elution methods starting from 45 GBq starting activity on Synthera® + synthesis module.