

RESEARCH ARTICLE

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4-Nitrophenyl activated esters are superior synthons for indirect radiofluorination of biomolecules†

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Indirect radiolabelling has for a long time been the mainstay strategy for radiofluorination of biomolecules. Acylation of biomolecules through the use of an ^{18}F -labelled activated ester is a standard method for indirect radiolabelling. However, the preparation of ^{18}F -labelled activated esters is typically a complex and multistep procedure. Herein, we describe the use of 4-nitrophenyl (PNP) activated esters to rapidly prepare ^{18}F -labelled acylation synthons in one step. Furthermore, we present a comparative study of PNP activated esters and the commonly utilised 2,3,5,6-tetrafluorophenyl (TFP) activated esters under direct radiofluorination conditions and demonstrate their relative acylation behaviour. We demonstrate the superiority of PNP esters under direct radiofluorination conditions with favourable acylation kinetics.

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Molecular imaging is transforming modern medicine through the use of radiopharmaceuticals that allow specific physiological processes occurring within the body to be detected at the cellular level.¹ Positron emission tomography (PET) is a common molecular imaging technique that has had a pronounced influence on personalized healthcare.² Radiopharmaceuticals utilised for PET imaging are typically labelled with short lived radionuclides including fluorine-18 (^{18}F), carbon-11 (^{11}C) and gallium-68 (^{68}Ga).^{3,4} In particular, the use of fluorine-18 to radiolabel peptides combines favourable pharmacology with the ideal physicochemical and imaging characteristics of fluorine-18.^{5–8} However, the preparation of ^{18}F -labelled peptides remains challenging and time consuming with multistep and laborious processes often required. Typically, sensitive biomolecules are radiolabelled with fluorine-18 indirectly, necessitating the use of a pre-labelled synthon (prosthetic groups) as they cannot tolerate the harsh radiofluorination conditions and are often thought to be inert to direct radiofluorination.⁹ [^{18}F]Fluoroacylation is a common method used for indirect radiolabelling of biomolecules.¹⁰ [^{18}F]Fluoroacylation synthons are typically prepared in multiple steps through the radiofluorination of

an appropriate stable ester, hydrolysis of the ester, activation of the resulting acid to form an acylating agent and final HPLC purification of the synthon.^{8,10–12} This multi-step process is then followed by the reaction of the isolated [^{18}F] fluoroacylation synthon with a biomolecule followed by a second HPLC purification to prepare the corresponding radiopharmaceutical. The use of TFP pre-activated diaryliodoniumbenzoate (**4**) was recently employed as a direct precursor to the TFP activated 4-[^{18}F]fluorobenzoate synthon [^{18}F]**6** (Fig. 2), though the radiosynthesis proceeded in less than 1% yield when standard radiofluorination conditions (containing KHCO_3 /Kryptofix mixture) were employed.¹³ The yield of ester [^{18}F]**6** was enhanced to 24% under less basic conditions (no addition of KHCO_3 /Kryptofix).¹³ Recently, we demonstrated the direct, one-step radiofluorination of peptides using ^{18}F -labelled pre-activated acylation agent 4-nitrophenyl 2-[^{18}F]fluoropropionate ([^{18}F]**1**).^{14–16} A similar approach has been reported for the preparation of 2,3,5,6-tetrafluorophenyl 6-[^{18}F]fluoronicotinate ([^{18}F]**2**) (Fig. 1).^{17,18}

Herein, we evaluate the stability of PNP and TFP esters to direct radiofluorination conditions for the preparation of 4-[^{18}F]fluorobenzoate and 6-[^{18}F]fluoronicotinate synthons. Furthermore, we report the relative stability and acylation

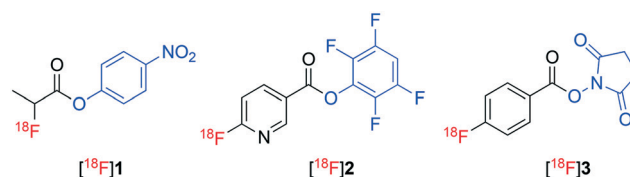


Fig. 1 [^{18}F]-labelled acylating agents [^{18}F]**1**, [^{18}F]**2** and [^{18}F]**3**.

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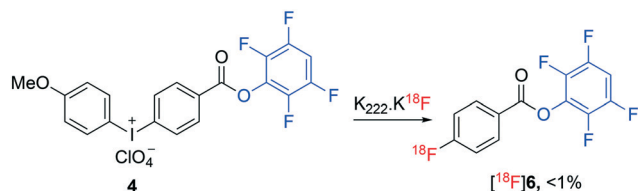


Fig. 2 Reported radiosynthesis of ester $[^{18}\text{F}]6$ by direct radiofluorination using $\text{K}_{222}\cdot\text{K}[^{18}\text{F}]\text{F}$ complex.¹³

activities of PNP and TFP pre-activated synthons. This study demonstrates the application of PNP active esters of ^{18}F -labelled synthons for subsequent radiolabelling of biomolecules.

We first attempted the synthesis of PNP and TFP esters of both 4-methoxyphenyliodonium benzoate and 6-trimethylammonium nicotinate, **16–19**, as precursors for direct radiofluorination. Esterification of 4-iodobenzoic acid **8** and 6-bromonicotinic acid **9** with 4-nitrophenol **10** and 2,3,5,6-tetrafluorophenol **11** afforded the corresponding PNP and TFP esters **12–15**, respectively (Fig. 3). Notably, the PNP esters **12** and **13** were formed in very high yield and were readily recrystallized immediately after work up to afford pure products. In comparison, the TFP esters **14** and **15** lacked the crystallinity of the PNP analogues making them less convenient to handle. Iodo-benzoates **12** and **14** were oxidized by mCPBA and converted to the corresponding (4-methoxyphenyl)-(phenyl)iodonium tosylates **16** and **17**. Bromo-nicotinates **13** and **15** were treated with 1 M trimethylamine in THF to form the corresponding ammonium species **18** and **19**. The formation of esters **16–19** proceeded readily in high yields and on gram scale, making them highly accessible precursors. However, TFP ester **19** required careful handling and storage as it was found to be particularly sensitive to hydrolysis.

Esters **16–19** were treated with $[^{18}\text{F}]$ fluoride in the form of $\text{K}_{222}\cdot\text{K}[^{18}\text{F}]\text{F}$ complex at 100 °C in DMSO:amyl alcohol (4:6) using potassium bicarbonate (for **18** and **19**) or potassium oxalate (for **16** and **17**) as accompanying bases. PNP esters **16** and **18** were readily soluble in the DMSO:*t*-amyl alcohol (TAA) mixture and were stable to the radiofluorination conditions leading to $94 \pm 2\%$ ($n = 3$) (58% isolated yield, decay corrected to SOS) and $85 \pm 3\%$ ($n = 3$) (42% isolated

yield, decay corrected to SOS) of the corresponding fluoronicotinate and fluorobenzoate PNP esters, $[^{18}\text{F}]20$ and $[^{18}\text{F}]21$, respectively. The yields of esters $[^{18}\text{F}]20$ and $[^{18}\text{F}]21$ ($94 \pm 2\%$ and $85 \pm 3\%$) were determined based on HPLC analysis of the crude product. The recovery of radioactivity from the analytical column was always above 85%. The formation of $[^{18}\text{F}]21$ demonstrates the first one-step synthesis of an activated ester of $[^{18}\text{F}]$ fluorobenzoate in high and reproducible yield. The yields of fluorobenzoate PNP ester $[^{18}\text{F}]21$ were strongly dependent on the mass of precursor **16** used; furnishing 0%, 68% and 88% yields when employing 5 mg, 10 mg and 20 mg of diaryliodonium precursor **16**, respectively. No such correlation was observed for the nicotinate PNP ester $[^{18}\text{F}]20$, with the greater reactivity of the nicotinate **18** towards nucleophilic aromatic substitution resulting in high yields of **20** on all scales. With excellent yields of esters $[^{18}\text{F}]20$ and $[^{18}\text{F}]21$ obtained, no further optimisation was undertaken.¹⁹

The performance of the analogous TFP esters **17** and **19** was then investigated under similar conditions. Nicotinate TFP ester **19** afforded $[^{18}\text{F}]2$ in variable yields (33–75%), with significant amounts of decomposition observed. This variability suggested the possible instability of the TFP activated ester $[^{18}\text{F}]2$. Benzoate TFP ester **17** was insoluble in the DMSO:amyl alcohol mixture, and was sparingly soluble in neat DMSO. Subjecting benzoate TFP ester **17** to direct radiofluorination in neat DMSO resulted in <1% yield of $[^{18}\text{F}]6$, a similar result to that reported by Richarz *et al.*¹³ However, when the insoluble mixture of precursor **17** in DMSO:amyl alcohol was heated with $\text{K}_{222}\cdot\text{K}[^{18}\text{F}]\text{F}$ complex, ester $[^{18}\text{F}]6$ was formed in 31% after 10 minutes. No further studies were conducted with ester **17** or product $[^{18}\text{F}]6$ due to their relative intolerance to direct radiofluorination conditions and low solubility (Fig. 4).

Next, the stability of $[^{18}\text{F}]20$, $[^{18}\text{F}]21$ and $[^{18}\text{F}]2$ in DMSO at room temperature was assessed. TFP ester $[^{18}\text{F}]2$ was found to undergo complete hydrolysis in less than one hour. In contrast, both PNP esters $[^{18}\text{F}]20$ and $[^{18}\text{F}]21$ esters were found to be remarkably stable, with no hydrolysis detected

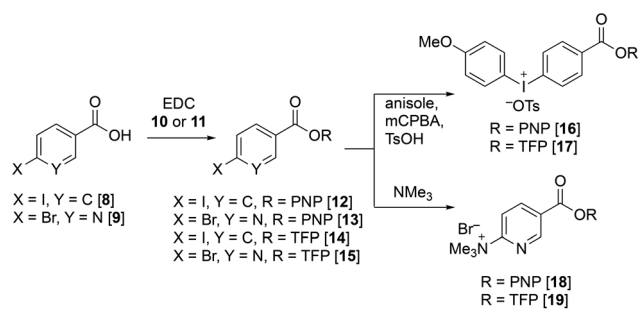


Fig. 3 The synthesis of precursors **16–19** from benzoic acid **8** and nicotinic acid **9**.

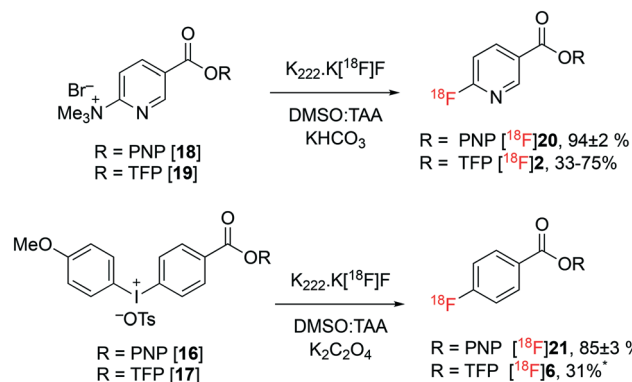


Fig. 4 Radiofluorination of esters **16–19** to form the corresponding synthons $[^{18}\text{F}]2$, $[^{18}\text{F}]6$, $[^{18}\text{F}]20$ and $[^{18}\text{F}]21$. Yields of $[^{18}\text{F}]20$ and $[^{18}\text{F}]21$ presented as average of $n = 3 \pm$ standard deviation. *Labelling attempted only once.



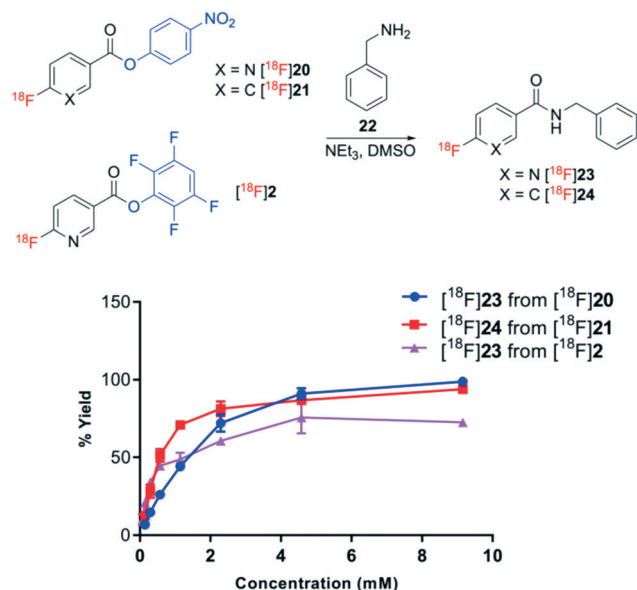


Fig. 5 Acylation of esters $[^{18}\text{F}]\text{20}$, $[^{18}\text{F}]\text{21}$ and $[^{18}\text{F}]\text{2}$ with benzylamine **22** to afford amidated products $[^{18}\text{F}]\text{23}$ and $[^{18}\text{F}]\text{24}$. % yield is the average of $n = 3$ with standard deviation bars shown.

over nine hours. Indeed, the susceptibility of $[^{18}\text{F}]\text{2}$ to hydrolysis and subsequent formation of by-products is well established.^{17,18}

The relative acylation propensities of esters $[^{18}\text{F}]\text{20}$, $[^{18}\text{F}]\text{21}$ and $[^{18}\text{F}]\text{2}$ were then examined with benzylamine **22** as a substrate. Varying concentrations of benzylamine **22** in DMSO (0.14–9.2 mM) containing 0.36 mM triethylamine were treated with $[^{18}\text{F}]\text{20}$, $[^{18}\text{F}]\text{21}$ and $[^{18}\text{F}]\text{2}$ for 10 min at room temperature (Fig. 5). In order to understand the relative acylation potential of these esters, it was essential to correct for their decomposition in neat DMSO (control reaction). Since only TFP-ester $[^{18}\text{F}]\text{2}$ underwent decomposition in the control reaction, only the resulting amidated product $[^{18}\text{F}]\text{23}$ was influenced by this correction affording slightly higher acylation yields. PNP esters $[^{18}\text{F}]\text{20}$ and $[^{18}\text{F}]\text{21}$ underwent rapid acylation of benzylamine affording 93–100% yields of amidated products $[^{18}\text{F}]\text{23}$ and $[^{18}\text{F}]\text{24}$, respectively, in the presence of 9.2 mM benzylamine **22**. The yields gradually decreased to 5–13% of amidated products $[^{18}\text{F}]\text{23}$ and $[^{18}\text{F}]\text{24}$ at 0.14 mM benzylamine **22**. The TFP ester $[^{18}\text{F}]\text{2}$ afforded the amidated product $[^{18}\text{F}]\text{23}$ in 73% yield when treated with 9.2

mM benzylamine **22**, which decreased to 10% with 0.14 mM benzylamine. Moreover, the yield of amides generated from TFP-benzoate $[^{18}\text{F}]\text{2}$ were less consistent than the yields generated from the PNP esters $[^{18}\text{F}]\text{20}$ and $[^{18}\text{F}]\text{21}$. The higher and more reproducible yields of amide products generated from the PNP esters $[^{18}\text{F}]\text{20}$ and $[^{18}\text{F}]\text{21}$ are presumed to be a direct consequence of their higher stability in solution.

These results highlight the superiority of PNP esters as activated intermediates for the direct $[^{18}\text{F}]\text{fluoroacylation}$ of biomolecules. This study reflects the importance of the relative stability of activated esters as synthons to facilitate efficient $[^{18}\text{F}]\text{fluoroacylation}$.

We next applied the PNP ester synthons $[^{18}\text{F}]\text{20}$ and $[^{18}\text{F}]\text{21}$ to the direct radiolabelling of RGD-peptide c(RGDyK) **25**. Both esters $[^{18}\text{F}]\text{20}$ and $[^{18}\text{F}]\text{21}$ were quantitatively consumed at room temperature over 10 min to form the corresponding radiolabelled peptides $[^{18}\text{F}]\text{26}$ and $[^{18}\text{F}]\text{27}$, respectively (Fig. 6). Radiolabelled peptides $[^{18}\text{F}]\text{26}$ and $[^{18}\text{F}]\text{27}$ were obtained in >98% radiochemical purity and with high molar activity (170 and 211 GBq μmol^{-1} , respectively) at end of synthesis as determined by analytical HPLC (using UV absorption at 220 nm).

Peptides $[^{18}\text{F}]\text{26}$ and $[^{18}\text{F}]\text{27}$ were prepared in $\geq 98\%$ conversion from the corresponding synthons $[^{18}\text{F}]\text{20}$ and $[^{18}\text{F}]\text{21}$. The overall reaction time, purification and formulation of peptides $[^{18}\text{F}]\text{26}$ and $[^{18}\text{F}]\text{27}$ was achieved in less than 90 minutes on an iPHASE Flexlab automated radiochemistry module. This is in stark contrast with previously reported conditions that employ multistep procedures (>3 radiochemical steps) for the preparation of synthons, leading to overall synthesis time of >130 minutes and ultimately affording fluorine-18 labelled c(RGDyK) in low overall yields and sub-optimal automation procedures.^{20,21}

Conclusions

In conclusion, PNP esters of $[^{18}\text{F}]\text{fluoronicotinate}$ $[^{18}\text{F}]\text{21}$ and $[^{18}\text{F}]\text{fluorobenzoate}$ $[^{18}\text{F}]\text{20}$ were formed under direct radiofluorination conditions at elevated temperatures. TFP-nicotinate $[^{18}\text{F}]\text{2}$ and TFP-benzoate $[^{18}\text{F}]\text{6}$ were found to be less tolerable of direct radiofluorination conditions employed. Moreover, the stability of the PNP esters renders them more effective at acylation in comparison to their TFP-counterparts.

Conflicts of interest

There are no conflicts to declare.

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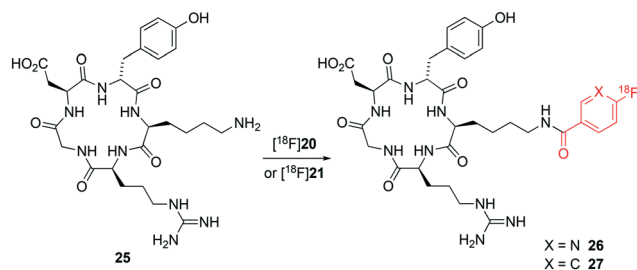


Fig. 6 Radiolabelling of c(RGDyK) peptide **25** with PNP esters $[^{18}\text{F}]\text{20}$ and $[^{18}\text{F}]\text{21}$.



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