**[18F]fluoro-benziodoxole: a no-carrier-added electrophilic fluorinating reagent. Rapid, simple radiosynthesis, purification and application for fluorine-18 labelling**

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Operational simplicity is the major advantage of using [18F]fluoro-benziodoxole as a fluorinating reagent. This electrophilic reagent is based on the ability of the palladium(IV) complex to accept fluorine (Fig. 1b). In the applied ligand exchange process the K–[18F]F interaction was transformed into Pd(IV)–[18F]F bonding. Change of the potassium counterion to a palladium(IV) center rendered the electron-rich fluorine-18 to an electron deficient one. Thus, the subsequent reductive elimination from the Pd(IV) complex created a new [18F]F–carbon bond. A similar process was developed using Ni(III) complexes.

[18F]fluoro-benziodoxole was developed starting from a cyclotron produced [18F]F− precursor, [18F]TBAF, and tosyl-benziodoxole. The synthetic utility of [18F]fluoro-benziodoxole was demonstrated by electrophilic fluorocyclization of α-styrylamides proceeding with high RCC (typically 50–90%) and high molar activity (up to 396 GBq μmol−1).

Organofluorine compounds are widely used in pharmaceutical and agrochemical industries as well as in medical diagnostics. The foremost radionuclide in Positron Emission Tomography (PET) is fluorine-18 because of its half-life of 109.8 minutes, the strong C–F bonding and the usually high metabolic stability of organofluorines. The optimal conditions for synthesis of organofluorines with the natural isotope fluorine-19 and those containing the radioactive isotope fluorine-18 are often fundamentally different. An important challenge in radiochemistry is to obtain [18F]-labelled organofluorine species in high radiochemical conversion (RCC) and at high molar activity within the lifetime of fluorine-18. Traditionally, high molar activity could only be achieved by nucleophilic [18F]-fluorination reactions. Electrophilic fluorination reactions target the realm of electron-rich functional groups, such as alkenes and aromatics with electron donating substituents. A characteristic feature of an electrophilic fluorinating reagent is the ability to accept an electron pair to their low-lying LUMO orbital having a substantial AO contribution from the fluorine atom, i.e. the fluorine-18 functional group in the reagent is electron deficient. A trivial example for such an electrophilic fluorination reagent is [18F]F2. However, this reagent is difficult to handle in radiosynthesis, very reactive (thus unselective) and it cannot be obtained with high molar activity.

Unfortunately, most of the selective electrophilic fluorination reagents, such as [18F]NFSI (Fig. 1a), [18F]Selectfluor and related species are also prepared from [18F]F2, which usually leads to unsatisfactory molar activities via rather cumbersome radiosynthetic procedures. On the other hand [18F]F− based reagents, such as [18F]KF or [18F]TBAF can be generated with high fluorine-18 molar activity.

However, these [18F]F− based reagents have an explicit nucleophilic reactivity, as the electron rich F− has an electron donating character. Construction of an electrophilic fluorinating reagent from [18F]F− requires the so-called “umpolung” of the reactivity. Ritter and co-workers have constructed an electrophilic fluorination reagent from a Pd(nI) complex and [18F]KF (Fig. 1b). In the applied ligand exchange process the K–[18F]F interaction was transformed into Pd(nI)–[18F]F bonding. Change of the potassium counterion to a palladium(nI) center rendered the electron-rich fluorine-18 to an electron deficient one. Thus, the subsequent reductive elimination from the Pd(nI) complex created a new [18F]F–carbon bond. A similar process was developed using Ni(nI) complexes as well. However, the widespread use of these methods has been limited by the complexity of the synthesis of the Pd(nI) and Ni(nI) complexes.

Safe, easy to handle and stable fluor-benziodoxole reagent was first reported by Prévost and Legault in 2012, and since then it has become a privileged fluorinating reagent. Recent DFT modelling studies discussed the mechanistic aspects of...
the electrophilic fluorination reactions by this hypervalent iodine reagent. Notably, fluoro-iodane \( \text{I} \) can easily be synthesized from anionic fluoride precursors.\(^{12c,14}\) Considering the importance of bridging the gap between the recent methodological advances in fluorine-19 chemistry and the high demand to access new types of fluorine-18 PET tracers for clinical diagnostics,\(^5\) we decided to develop a new methodology for rapid radiosynthesis of electrophilic \([^{18}\text{F}]\)fluoro-benziodoxole \([^{18}\text{F}]\text{I} \) reagent from standard \([^{18}\text{F}]\text{F} \) precursors (Fig. 1c).

Our fluorine-19 experiments have shown that the synthesis of \( \text{I} \) from commonly used benziodoxole precursors, such as OH, OAc and halogenide derivatives requires relatively long reaction times (0.5–4 h). Therefore, the highly reactive (yet bench-stable) tosyl derivative \( \text{II} \) was selected as precursor for the radiosynthesis. Reagent \( \text{I} \) partially or completely decomposes under chromatography due to the high affinity of the iodine-bond fluorne to Si, Al and alkali metal ions. Indeed, over 95% of \([^{18}\text{F}]\text{I} \) was lost, when we attempted to purify it on silica or alumina adsorbents (ESI,\(^\dagger\) S12). Therefore, the analysis and purification of \([^{18}\text{F}]\text{I} \) cannot be achieved by chromatographic methods, as fluoro-iodines \( \text{I} \) and \([^{18}\text{F}]\text{I} \) decomposed when subjected to reverse-phase HPLC, TLC or paper chromatography. The alternative purification method was based on the relatively good solubility of \( \text{I} \) in hexane. According to our fluorine-19 experiments the starting materials tosyl-benziodoxole \( \text{II} \) and TBAF, as well as the inorganic compounds formed in the synthesis of \( \text{I} \) are insoluble in hexane. Thus, after the allotted times (Table 1) we evaporated the solvent (DCM) of the reaction mixture of \( \text{II} \) and \([^{18}\text{F}]\text{Bu}_4\text{NF} \) and, then the solid residue was extracted with hexane. We assumed that the entire amount of the radioactivity of the hexane solution arose from \([^{18}\text{F}]\text{I} \), and this radioactivity was the basis of the calculation of RCC (Table 1).

To get further evidence for the formation of \([^{18}\text{F}]\text{I} \) we studied the chemical composition of a decayed sample of \([^{18}\text{F}]\text{I} \) by \( ^{19}\text{F}-\text{NMR} \) and HRMS. Such analysis takes advantage of the inevitable co-production of unlabeled \( \text{I} \), which is formed by the reaction of \( \text{II} \) with the minute amounts of fluorine-19 that is present in the reaction mixture. Indeed, our analysis of the decayed sample of \([^{18}\text{F}]\text{I} \) confirmed both by \( ^{19}\text{F}-\text{NMR} \) and HRMS the presence of \([^{19}\text{I}]\text{I} \) in the hexane extract (ESI,\(^\dagger\) S9–S12), which is diagnostic of the formation of \([^{18}\text{F}]\text{I} \) and its extraction to hexane. Interestingly, TBAF was not detected in the \( ^{19}\text{F}-\text{NMR} \) spectrum of the decayed sample indicating that the extracted sample did not contain significant contamination of \([^{18}\text{F}]\text{Bu}_4\text{NF} \).

We have carefully optimized the reaction temperature, time and concentration of the reaction of \( \text{II} \) and \([^{18}\text{F}]\text{Bu}_4\text{NF} \) and the temperature for the extraction with hexane (Table 1). Initially, we carried out the reactions in DCM (typically 500 \( \mu \)L at 70 °C (entries 1–5). Using 5 mg of \( \text{II} \) the RCC of \([^{18}\text{F}]\text{I} \) was similar (40–47%) after 40 and 20 minutes of reaction time (entries 1 and 2). Increasing the amount of \( \text{II} \) (entry 3) or increasing the concentration of the reactants (by reducing the amount of solvent, entry 4) did not change significantly the RCC (42% and 37% respectively). Contrarily, decreasing the concentration (entry 5) led to increasing of RCC (56%), albeit an elongated time for the evaporation of DCM (entry 5). Therefore, we kept the solvent volume at 500 \( \mu \)L in the subsequent experiments. It was found that the RCC was nearly unchanged when the reaction was carried out at room temperature instead of 70 °C (entry 6). Moreover, the reaction time could be shortened to five minutes without decreasing the RCC (41%, entry 7) thus reducing significantly the duration of the radiosynthesis of \([^{18}\text{F}]\text{I} \).

In the above procedures (entries 1–7), we employed 70 °C for the extraction of the solid residue that remained after the evaporation of DCM. Decreasing the extraction temperature to 40 °C or room temperature (entries 8 and 9) led to significant decrease of the RCC (20–23%). The optimal procedure (entry 7) involves the reaction of \( \text{II} \) and \([^{18}\text{F}]\text{Bu}_4\text{NF} \) in 500 \( \mu \)L DCM (5 minutes at room temperature); then evaporation of DCM and extraction of the solid residue by hexane for one minute at 70 °C.

As mentioned above, the fluorine-19 experiments showed that TBAF is insoluble in hexane. To confirm this observation, we made a further control experiment using \([^{18}\text{F}]\text{Bu}_4\text{NF} \). We carried out the above optimized three-step procedure without employing \( \text{II} \) (entry 10). The radioanalysis indicated 7% RCC for the extracted inorganic fluoride. The extracted amounts of \([^{18}\text{F}]\)fluoride were about the same by varying the extraction temperature (entries 11 and 12). These experiments (together with the fluorine-19 results) confirmed that the above three-step procedure results in high purity sample of \([^{18}\text{F}]\text{I} \) with very low (if at all) contamination of \([^{18}\text{F}]\text{Bu}_4\text{NF} \). Notably, the chemical stability of \( \text{II} \) allows for centralized large-scale production and subsequent shipment of \([^{18}\text{F}]\text{I} \) to PET centers that lack an on-site cyclotron.

In order to demonstrate the radiosynthetic utility of purified \([^{18}\text{F}]\text{I} \) for \( ^{18}\text{F} \)-labelling we performed electrophilic fluorocyclization reactions of o-styrylamides 3. Gulder and co-workers\(^{12b,14}\) have shown that using equimolar amounts of 3 and 1 a rapid fluorocyclization can be performed at room temperature resulting in fluoro-benzoxazepines bearing a quaternary fluoride.
Radiosynthesis of $[^{18}F]4$ was performed by adding the hexane solution of $[^{18}F]1$ to a solution of precursor 3 in acetonitrile followed by evaporation of the solvents. Subsequently, the solid residue was dissolved in acetonitrile (500 μL) and the fluorocyclization reaction was studied at different temperatures and reaction times (Table 2). Using the optimal fluorocyclization conditions reported by Gulder for the reaction with fluoro-1912b (using 3a and 1), we obtained low but encouraging (9% RCC) levels of the fluoro-benzoxazepines product $[^{18}F]4$a (Table 2, entry 1). Increasing the temperature from room temperature to 90 °C and decreasing the reaction time from 40 minutes to 7 minutes led to a significant increase of the RCC to 76% (entries 2–4). Further elongation of the reaction time did not improve the RCC (74%, entry 5). The bromo analog $[^{18}F]3b$ and the isomeric chloro o-styrilamide 3c reacted under identical conditions affording the corresponding fluoro-benzoxazepines $[^{18}F]4$b and $[^{18}F]4$c in 57 and 88% RCC (entries 6 and 7). Using these conditions, Bu amide $[^{18}F]4$d was obtained in 54% RCC along with an unidentified $[^{18}F]$-labeled impurity (28%). This impurity was probably a decomposition product resulting from the chemical instability of $[^{18}F]4$d. Decreasing the reaction temperature to 70 °C led to an RCC of 50% (entry 10) with reduction of the labeled impurity to 20%. When 3e, having a more electron rich nitrogen than 3c, was reacted with $[^{18}F]1$ under identical conditions, the RCC of $[^{18}F]4$e was much lower (10%, entry 11) than for $[^{18}F]4$c (88%, entry 7). A possible reason can be the relatively low chemical stability of $[^{18}F]4$e at elevated temperatures. When the reaction time was decreased from seven to two minutes, the RCC increased to 90% (entry 12). The electrophilic fluorination of the electron-rich alkene in 3f is probably faster than for its electron-poor counterparts in 3a–b (entries 4 and 6), but product $[^{18}F]4$f has low thermostability compared to $[^{18}F]4$a–b. Accordingly, the RCC for the formation of $[^{18}F]4$f was relatively low at 90 °C (46%, entry 13). However, when the reaction temperature was decreased to 70 °C the RCC was increased to 74% (entry 14). The parent o-styrilamide 3g reacted also smoothly at 70 °C in 2 minutes affording $[^{18}F]4$g with 80% of RCC. A final confirmation that the nucleophilic fluorination reagent $[^{18}F]$TBAF is not able to perform fluorocyclization is demonstrated by entry 16. When 3g was reacted with $[^{18}F]$TBAF instead of $[^{18}F]1$ formation of $[^{18}F]4$g was not observed. This control experiment also proved that even if small amounts of $[^{18}F]$TBAF would have contaminated the purified sample of $[^{18}F]1$, it was not able to perform fluorocyclization of o-styrilamides 3.

We also performed one-pot reactions without isolation and purification of $[^{18}F]1$ (Fig. 2). When the one-pot reaction was performed in two sequential steps: (1) in situ formation of $[^{18}F]1$ followed by (2) fluorocyclization of 3a the reaction time was shorter by about 5 minutes than for the version with purified $[^{18}F]1$ (Table 2, entry 4, RCC = 76%). However, the RCC in the one-pot reaction was poor: 14% and 34% based on $[^{18}F]$Bu$_4$NF or $[^{18}F]1$ as limiting reagent, respectively (Fig. 2). These results confirm that radiosynthetic attempts for using in situ generated $[^{18}F]1$ proceed with inferior results compared to the above method with purified $[^{18}F]1$ reagent.12

The optimized protocol was employed for the determination of the isolated yield (RCY) and the molar activity. In this reaction 3.93 GBq of $[^{18}F]$Bu$_4$NF were reacted with 2 affording

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**Table 2** Electrophilic fluorocyclization affording $[^{18}F]$fluoro-benzoxazepines

<table>
<thead>
<tr>
<th>Entry</th>
<th>Precursor</th>
<th>T (°C)</th>
<th>t (min)</th>
<th>Product</th>
<th>RCC (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3a</td>
<td>RT</td>
<td>40</td>
<td>$[^{18}F]4$a</td>
<td>9 ± 2 (n = 2)</td>
</tr>
<tr>
<td>2</td>
<td>3a</td>
<td>50</td>
<td>40</td>
<td>$[^{18}F]4$a</td>
<td>7 ± 2 (n = 2)</td>
</tr>
<tr>
<td>3</td>
<td>3a</td>
<td>50</td>
<td>7</td>
<td>$[^{18}F]4$a</td>
<td>12 ± 1 (n = 2)</td>
</tr>
<tr>
<td>4</td>
<td>3a</td>
<td>90</td>
<td>7</td>
<td>$[^{18}F]4$a</td>
<td>76 ± 2 (n = 2)</td>
</tr>
<tr>
<td>5</td>
<td>3a</td>
<td>90</td>
<td>15</td>
<td>$[^{18}F]4$a</td>
<td>74 ± 4 (n = 2)</td>
</tr>
<tr>
<td>6</td>
<td>3b</td>
<td>90</td>
<td>7</td>
<td>$[^{18}F]4$a</td>
<td>57 ± 1 (n = 2)</td>
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<tr>
<td>7</td>
<td>3b</td>
<td>90</td>
<td>15</td>
<td>$[^{18}F]4$a</td>
<td>88 ± 6 (n = 2)</td>
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<tr>
<td>8</td>
<td>3b</td>
<td>90</td>
<td>7</td>
<td>$[^{18}F]4$a</td>
<td>54 ± 4 (n = 2)</td>
</tr>
<tr>
<td>9</td>
<td>3d</td>
<td>90</td>
<td>2</td>
<td>$[^{18}F]4$a</td>
<td>19 ± 6 (n = 2)</td>
</tr>
<tr>
<td>10</td>
<td>3d</td>
<td>70</td>
<td>7</td>
<td>$[^{18}F]4$a</td>
<td>50 ± 6 (n = 2)</td>
</tr>
<tr>
<td>11</td>
<td>3e</td>
<td>90</td>
<td>2</td>
<td>$[^{18}F]4$a</td>
<td>10 ± 1 (n = 2)</td>
</tr>
<tr>
<td>12</td>
<td>3e</td>
<td>90</td>
<td>7</td>
<td>$[^{18}F]4$a</td>
<td>90 ± 2 (n = 2)</td>
</tr>
<tr>
<td>13</td>
<td>3e</td>
<td>90</td>
<td>15</td>
<td>$[^{18}F]4$a</td>
<td>46 ± 3 (n = 2)</td>
</tr>
<tr>
<td>14</td>
<td>3f</td>
<td>90</td>
<td>2</td>
<td>$[^{18}F]4$a</td>
<td>74 ± 5 (n = 2)</td>
</tr>
<tr>
<td>15</td>
<td>3e</td>
<td>90</td>
<td>7</td>
<td>$[^{18}F]4$a</td>
<td>80 ± 1 (n = 2)</td>
</tr>
</tbody>
</table>

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"To a solution of 3 in MeCN (0.4 μmol, 25 μL) was added $[^{18}F]1$ in hexane (50–100 μL). The mixture was evaporated and re-dissolved in MeCN (500 μL) before heating at the indicated temperatures and times.\(^{2}\) Estimated by radio-HPLC of the crude reaction mixture starting from $[^{18}F]1$.\(^{3}\)Using $[^{18}F]$Bu$_4$NF instead of $[^{18}F]1$.\(^{4}\) RCC based on $[^{18}F]$Bu$_4$NF as limiting reagent, \(^{5}\) RCC based on $[^{18}F]1$ assuming that it formed with 41% RCC (Table 1, entry 7).
904 MBq of $^{[18}F]1$, which was used for the labelling of 3a in MeCN at 90 °C for 7 minutes. The reaction resulted in 92.4 MBq of $^{[18}F]1\text{a}$ (with >99% radiochemical purity) corresponding to 10% isolated activity yield (RCY) and a molar activity of 396 GBq μmol$^{-1}$. This molar activity is in the range of the values obtained for tracers by nucleophilic fluorination in our laboratories and orders of magnitude higher than the corresponding values reported for the electrolytic fluorination methods based on application of $^{[18}F]F_2$.$^{6–8}$ The overall time of the reaction from the bombardment to isolation of the $^{[18}F]F$benzoxazepine product was about two hours, of which the radio-synthesis and purification of $^{[18}F]1\text{a}$ from $^{[18}F]1\text{ABF}$ was achieved in about 20 minutes. This method was not automatized, and thus proper automatization may further improve the efficiency of fluorine-18 labelling by $^{[18}F]1$ in clinical applications.

In summary, we have presented a new radiosynthetic method for the preparation of the electrophilic fluorination reagent $^{[18}F]1$ from a nucleophilic $^{[18}F]F^-$ precursor, $^{[18}F]1\text{ABF}$. The reaction and purification of $^{[18}F]1$ could be performed by an operationally simple three-step process within about 20 minutes. The purified $^{[18}F]1$ reagent was free from the excess of its precursor 2 and may contain only traces of $^{[18}F]1\text{ABF}$. Since 1 is chemically stable for an indefinite time, the purified $^{[18}F]1$ is even transportable in hexane solution to PET facilities that lack on-site cyclotrons. We have also demonstrated the synthetic utility of $^{[18}F]1$ for electro-philic fluorocyclization of o-stryrylamides. The reactions could be performed in 2–7 minutes with typically high RCC. The preparative scale experiments led to a product with 396 GBq μmol$^{-1}$ of molar activity, which is higher than the molar activity typically obtained in electrophilic $^{18}F$-labelling experiments. We will continue in our laboratories the studies of exploitation of $^{[18}F]1$ in further $^{18}F$-labelling applications and development of further hypervalent iodine based $^{18}F$-fluorination/alkylfluorination reagents.

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Conflicts of interest

There are no conflicts to declare.

Notes and references


