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# Synthesis and biological evaluation of PET tracers designed for imaging of calcium activated potassium channel 3.1 ( $K_{Ca}3.1$ ) channels *in vivo*<sup>†</sup>

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Expression of the  $Ca^{2+}$  activated potassium channel 3.1 ( $K_{Ca}3.1$ ) channel (also known as the Gárdos channel) is dysregulated in many tumor entities and has predictive power with respect to patient survival. Therefore, a positron emission tomography (PET) tracer targeting this ion channel could serve as a potential diagnostic tool by imaging the  $K_{Ca}3.1$  channel *in vivo*. It was envisaged to synthesize [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]**1**) since senicapoc (**1**) shows high affinity and excellent selectivity towards the  $K_{Ca}3.1$  channels. Because problems occurred during <sup>18</sup>F-fluorination, the [<sup>18</sup>F]fluoroethoxy senicapoc derivative [<sup>18</sup>F]**28** was synthesized to generate an alternative PET tracer targeting the  $K_{Ca}3.1$  channel. Inhibition of the  $K_{Ca}3.1$  channel by **28** was confirmed by patch clamp experiments. *In vitro* stability in mouse and human serum was shown for **28**. Furthermore, biodistribution experiments in wild type mice were performed. Since [<sup>18</sup>F]fluoride was detected *in vivo* after application of [<sup>18</sup>F]**28**, an *in vitro* metabolism study was conducted. A potential degradation route of fluoroethoxy derivatives *in vivo* was found which in general should be taken into account when designing new PET tracers for different targets with a [<sup>18</sup>F]fluoroethoxy moiety as well as when using the popular prosthetic group [<sup>18</sup>F]fluoroethyl tosylate for the alkylation of phenols.

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## 1. Introduction

Dysregulation of the  $K_{Ca}3.1$  channel is directly associated with many different diseases such as asthma,<sup>1,2</sup> sickle cell anemia,<sup>3–5</sup> coronary restenosis after angioplasty,<sup>6</sup> kidney fibrosis<sup>7,8</sup> and atherosclerosis.<sup>9,10</sup> It has a place in corneal<sup>11</sup> and conjunctival fibrosis<sup>12</sup> and was shown to be vastly overexpressed in Alzheimers disease.<sup>13–15</sup> Furthermore, the  $K_{Ca}3.1$  channel is a modulator in hepatic injury.<sup>16</sup> The  $K_{Ca}3.1$  channel plays an important role in cancer diseases: Along with other ion channels it contributes to

features of essential “cancer hallmarks”, such as proliferation, migration, invasion and extravasation.<sup>17–19</sup> Consequently, the ion channel plays a critical role in the metastatic cascade. Inhibition of the  $K_{Ca}3.1$  channel reduces proliferation, migration and metastasis in different tumor entities.<sup>20–22</sup> The direct correlation between overexpression of the  $K_{Ca}3.1$  channel and tumor grade as well as metastatic status highlights the importance of this ion channel in cancer diseases.<sup>23</sup> Furthermore, overexpression of  $K_{Ca}3.1$  channels is often related to poor prognosis of tumor patients and high lethality rates.<sup>24,25</sup> Therefore, the  $K_{Ca}3.1$  channel could serve as predictive marker with respect to patient survival.<sup>23–25</sup> Additionally, pro-tumor functions of endogenously expressed  $K_{Ca}3.1$  channels have been shown in a mouse model.<sup>18,26</sup>

The impact of the potassium channel on various diseases is rather outstanding, please also refer to the available comprehensive reviews.<sup>27–29</sup>

Our aim is to enable *in vivo* visualization of the  $K_{Ca}3.1$  channel *via* PET technology to offer the opportunity of a potential diagnostic tool in different diseases. We decided to focus on  $K_{Ca}3.1$  channels in cancer diseases since essentially all cells of the tumor stroma express  $K_{Ca}3.1$  channels.<sup>30</sup> A signal from a potential PET tracer binding to this ion channel should therefore be very robust. Thus,  $K_{Ca}3.1$  channels in tumor cells are particularly suitable as a first model for testing PET tracers targeting this ion channel.

Herein, we present the design and radiosynthesis of potential PET tracers for imaging of the  $K_{Ca}3.1$  channel *in vivo*.

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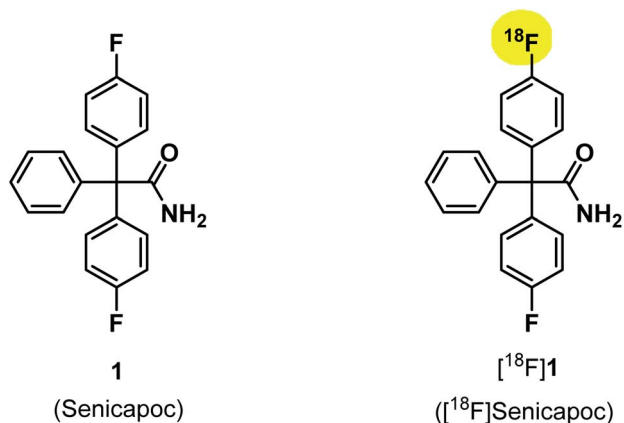
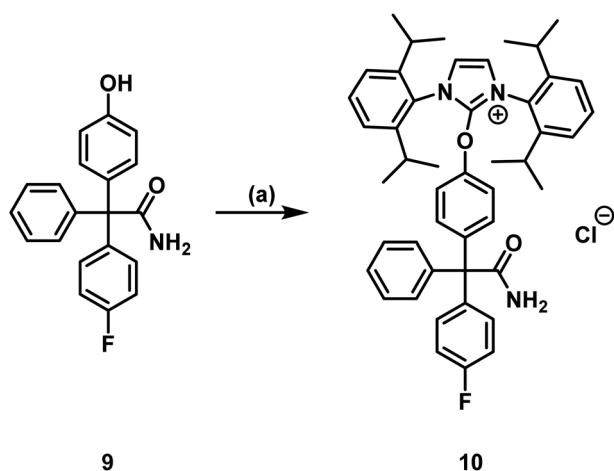



Fig. 1 Structures of senicapoc (**1**) and [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]1).



Scheme 1 Synthesis of the uronium salt precursor **10** from phenol **9**: reagents and reaction conditions: (a) 2-chloro-1,3-bis(2,6-diisopropylphenyl)-1H-imidazol-3-ium chloride (**8**), Ag<sub>2</sub>CO<sub>3</sub>, CH<sub>2</sub>Cl<sub>2</sub>, 60 °C, 4 h, 68%.

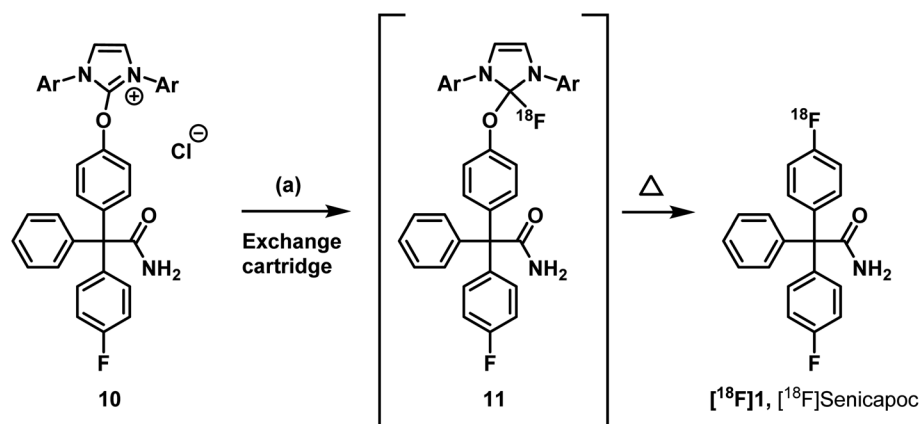
The K<sub>Ca</sub>3.1 channel inhibitor senicapoc (**1**) served as starting point for the development of PET tracers for *in vivo* imaging of the ion channel. Senicapoc (**1**) shows a high affinity towards the

K<sub>Ca</sub>3.1 channel with an IC<sub>50</sub> value of 11 ± 2 nM (measured on human erythrocytes) and excellent selectivity over related ion channels.<sup>31</sup> These properties are promising for selective imaging of the K<sub>Ca</sub>3.1 channel *in vivo*. Senicapoc (**1**) contains two fluorine atoms. Replacement of one fluorine atom by fluorine-18 results in [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]1) bearing an authentic label (Fig. 1). Moreover, senicapoc (**1**) was already evaluated in phase III clinical trials for potential therapy of sickle cell anemia, so data regarding toxicity and tolerability are available reducing the effort for first *in vivo* studies using the new tracer in humans. Herein, different strategies for the radiosynthesis of [<sup>18</sup>F]senicapoc, the synthesis of suitable precursors as well as advantages and disadvantages of the chosen routes are presented.

## 2. Synthesis

To identify the [<sup>18</sup>F]labeled compounds by co-injection on the radio-HPLC system after radiosynthesis, non-radioactive [<sup>19</sup>F]1 reference compound had to be synthesized first. Therefore, senicapoc (**1**) was prepared on the basis of previously described procedures.<sup>32</sup> (see ESI, Scheme S1†). The literature-known synthesis of senicapoc (**1**) is not suitable for radiosynthesis as a multi-step protocol is following the introduction of the fluorine atoms. In PET tracer synthesis the fluorine has to be introduced in the last steps due to the short half-life of [<sup>18</sup>F]fluorine.

In 2016, Neumann *et al.* published the deoxyfluorination of phenols *via* concerted nucleophilic aromatic substitution (CSNAr), which was also performed with [<sup>18</sup>F]fluoride.<sup>33</sup> Uronium salts were used as precursors for the radiosynthesis. We started our efforts to prepare [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]1) using this method, which required the uronium salt **10** *via* the phenolic derivative **9**. (Scheme 1): Synthesis of the uronium salt precursor **10** from phenol **9**: reagents and reaction conditions: (a) 2-chloro-1,3-bis(2,6-diisopropylphenyl)-1H-imidazol-3-ium chloride (**8**), Ag<sub>2</sub>CO<sub>3</sub>, CH<sub>2</sub>Cl<sub>2</sub>, 60 °C, 4 h, 68%. The phenol **9** was synthesized in five steps as recently published in connection with the synthesis of fluorescently labeled probes for *in vitro* imaging of K<sub>Ca</sub>3.1 channels.<sup>34–37</sup>



Scheme 2 Radiosynthesis of [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]1). Reagents and reaction conditions: (a) [<sup>18</sup>F]F<sup>-</sup>, butan-2-one : EtOH : Bu<sub>3</sub>N (100 : 10 : 1), 130 °C, 20 min. RCY (d.c.) = 0.4% (n = 1).



Table 1 Reagents and reaction conditions for the  $^{18}\text{F}$ -labelling of trimethylarylammonium salt precursor **18**

Eluent solution (in $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ )	Solvent	Temperature [ $^\circ\text{C}$ ]	Time [min]	Product
$\text{K}_2\text{CO}_3$ (aq), $\text{K}_{222}$	Acetonitrile ( $\text{CH}_3\text{CN}$ )	120	15	—
$\text{K}_2\text{CO}_3$ (aq), $\text{K}_{222}$	Dimethylformamide (DMF)	150	15	—
$\text{K}_2\text{CO}_3$ (aq), $\text{K}_{222}$	DMF	155	15	—
$\text{K}_2\text{CO}_3$ (aq), $\text{K}_{222}$	Dimethylsulfoxide (DMSO)	165	30	—
$\text{K}_2\text{C}_2\text{O}_4$ (aq), $\text{K}_{222}$	DMF	155	30	—
$\text{Bu}_4\text{NHCO}_3$ (aq)	DMF	150	15	?
$\text{Bu}_4\text{NHCO}_3$ (aq)	DMSO	150	15	?
$\text{Bu}_4\text{NHCO}_3$ (aq)	DMSO	140	15	?

Neumann *et al.* used the uronium salts prepared by this method directly for  $^{18}\text{F}$ -deoxyfluorination without further purification.<sup>33</sup> Since the purity of the prepared uronium salt **10** was only 78% (HPLC), we decided to purify **10** by column chromatography. Thus, uronium salt **10** was isolated with 99.2% purity (HPLC) and was fully characterized. The purified as well as the unpurified uronium salts **10** were used in radiosynthesis test reactions in order to obtain  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**) (Scheme 2).

The radiosynthesis was performed semiautomatically according to Neumann *et al.*<sup>33</sup> An aqueous solution of  $^{18}\text{F}$  fluoride was flushed through an anionic exchange cartridge. Different cartridges were used during evaluation, starting with Chromabond® PS- $\text{HCO}_3$  cartridges containing 45 mg or 10 mg sorbent identical with the sorbent of the Chromafix PS- $\text{HCO}_3$  cartridges used in the original publication.<sup>33</sup> After trapping of  $^{18}\text{F}$  fluoride, the cartridge (45 mg) was flushed with butan-2-one : ethanol (1 : 1, 1 mL). Thereafter, the cartridge was reversed manually using a luerlock® system as suggested in the original procedure. Since this step is highly critical in terms of radiation protection, we only used maximum starting activities of  $<1$  GBq. After reversing the cartridge it was eluted with  $5 \pm 1$  mg of the uronium salt **10** in butan-2-one:ethanol:tributylamine (100 : 10 : 1, 1 mL). In an average synthesis  $40 \pm 10\%$  of the starting activity remained on the cartridge. After a reaction time of 20 min at  $130^\circ\text{C}$ , only a small fraction of the theoretical activity was found in the reactor solution. This is a second critical step in the protocol, as the only explanation for this reproducible result is the formation of volatile  $^{18}\text{F}$ -labeled compounds. The protocol led to  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**). Unfortunately, the yield was much lower than in comparable published labelling approaches (0.4% RCY d.c.). Moreover, the radiochemical purity was insufficient (87%) (for chromatogram see ESI: Fig. S1†).

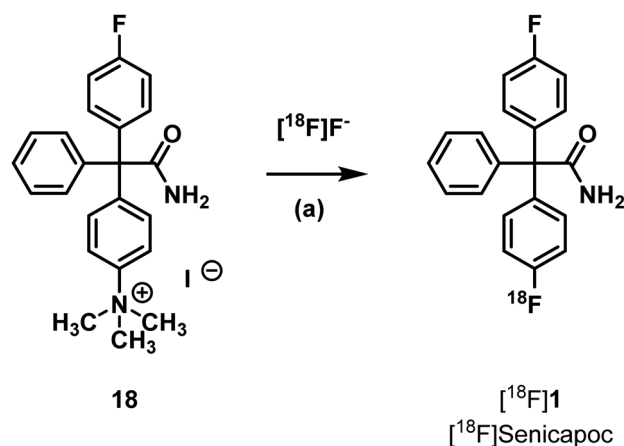
The very low radiochemical yield in combination with the very low starting activity (due to radiation protection regulations) and the technical problems mentioned above rendered *in vivo* experiments impossible. The use of cartridges with lower sorbent amounts (10 mg) resulted in even lower elution efficacy combined with considerably lower radiochemical yields ( $<0.1\%$  RCY d.c.). The use of our standard cartridges Sep-Pak® Light QMA Fa. Waters® produced even worse results. However, in these experiments the cartridges were not reversed in the elution process to avoid the risk of contamination. 67% of the starting activity remained on the cartridge and it was not

possible to isolate  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**). Further experiments combining the elution with potassium hydrogencarbonate ( $\text{KHCO}_3$ ) (5 mg) as well as different elution solutions did not improve the initial result. For these reasons, the synthesis of  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**) using the uronium salt precursor **10** was no longer pursued.

Another strategy is the direct nucleophilic substitution of a suitable precursor with  $^{18}\text{F}$  fluoride to produce  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**). A well-established leaving group to achieve aromatic nucleophilic substitution is the trimethylammonium group.<sup>38,39</sup> Although the aromatic system is not specially activated by electron withdrawing functional groups, the  $^{18}\text{F}$ -labelling of the trimethylammonium precursor **18** was envisaged. The synthesis of such a precursor is similar to the synthesis of senicapoc (**1**) itself (see ESI, Scheme S3†).<sup>32</sup>

Radiosynthesis by nucleophilic aromatic substitution with  $^{18}\text{F}$  fluoride was performed using various conditions summarized in Table 1 (Scheme 3 and Table 1).

The investigated reaction conditions using carbonate or oxalate as base did not lead to  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**). The use of tetrabutylammonium hydrogen carbonate as base led to the formation of tetrabutylammonium- $^{18}\text{F}$ -fluoride ( $^{18}\text{F}$ TBAF) as fluorination agent. All reactions with  $^{18}\text{F}$ TBAF provided a product with very similar polarity compared to  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**). The retention times on the semipreparative as well as on the analytical radio-HPLC were very similar to those



Scheme 3 Radiosynthesis of  $^{18}\text{F}$ senicapoc ( $^{18}\text{F}$ **1**). Reagents and reaction conditions: (a) cartridge: Sep-Pak® Light QMA Fa. Waters®, conditions see Table 1.



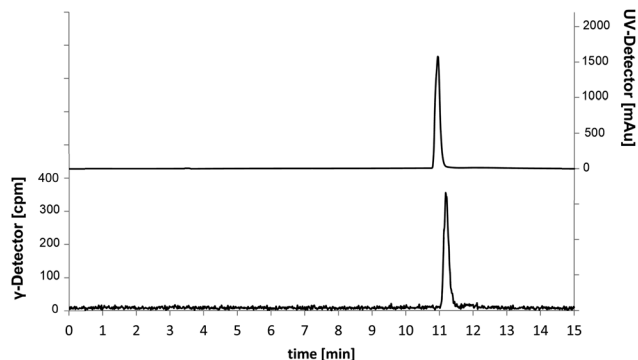
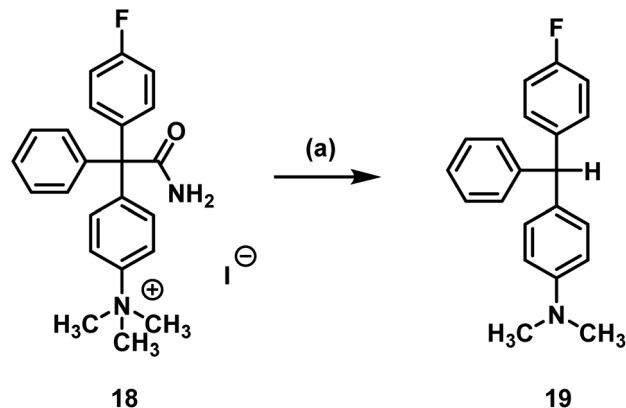


Fig. 2 Chromatograms of the analytical radio-HPLC. Traces for  $^{18}\text{F}$ -labelled ( $\gamma$ -detector [cpm], bottom) compound and co-injection of [ $^{19}\text{F}$ ]senicapoc (**1**) as reference compound (UV-detector [mAu], top).

determined for senicapoc (**1**) (Fig. 2). Due to the high similarity, we initially assumed a successful transformation.

In order to prepare a solution for injection after HPLC purification, the solvents were removed under reduced pressure. Afterwards, only a small fraction of the theoretical radioactivity was left. It was postulated that the reduced amount of radioactivity is due to removal of a volatile  $^{18}\text{F}$ -labelled compound. As [ $^{18}\text{F}$ ]senicapoc ([ $^{18}\text{F}$ ]**1**) should not be able to release a volatile  $^{18}\text{F}$ -species, the result of the synthesis was questioned. To get a better insight into the decomposition of the  $^{18}\text{F}$ -labelled compound obtained during radiosynthesis, samples were taken during the removal of the solvent and analyzed by analytical radio-HPLC and radio-TLC (Fig. 3; for further chromatograms see ESI: Fig. S2 and S3<sup>†</sup>).

It was shown that decomposition occurred leading to a less polar compound. The peak with a retention time of 3 min (analytical radio-HPLC; Fig. 3), which increased over time, as well as the baseline peak in the radio-TLC (see ESI: Fig. S4<sup>†</sup>) are a strong hint towards the formation of [ $^{18}\text{F}$ ]fluoride. Since senicapoc (**1**) is stable even at high temperature and under reduced pressure, it was suggested that a different product had been formed. To get an idea of the formed product, the fluorination of the quaternary ammonium compound **18** was performed with [ $^{19}\text{F}$ ]fluoride under the conditions of the



Scheme 4 Decarboxylation and demethylation of trimethylarlammonium precursor **18**. Reagents and reaction conditions: (a) TBAF, DMF, 155 °C, 10 min, 78%.

radiosynthesis. Indeed, the reaction of **18** with TBAF in DMF at 155 °C for 10 min resulted in the less polar compound **19**, which was purified and fully characterized. Spectroscopic analysis showed the formation of triarylmethane **19**, which was formed by decarboxylation and demethylation of the quaternary ammonium precursor **18** (Scheme 4).

The same reaction occurred in dry DMF used in the radiosynthesis. The formation of [ $^{18}\text{F}$ ]fluoride during radiochemistry may be explained by radical formation of the  $^{18}\text{F}$ -substituted triarylmethane, which would lead to loss of fluoride upon dimerization forming a quinoidic system. The dimerization would form a structure that is a substituted Gomberg's dimer.

The titanium nanoparticle catalyzed nucleophilic substitution of aryl tosylates to obtain [ $^{18}\text{F}$ ]aryl compounds was successfully established by Sergeev *et al.*<sup>40</sup> The required tosylate precursor **20** was obtained by reaction of phenol **9** with tosyl chloride (Scheme 5).

Since stability issues occurred using the trimethylarlammonium salt **18** as precursor in  $^{18}\text{F}$ -labelling reactions, the stability of the prepared tosylate precursor **20** was evaluated before starting radiosynthesis. Usually, the titanium nanoparticle catalyzed nucleophilic substitution takes place at

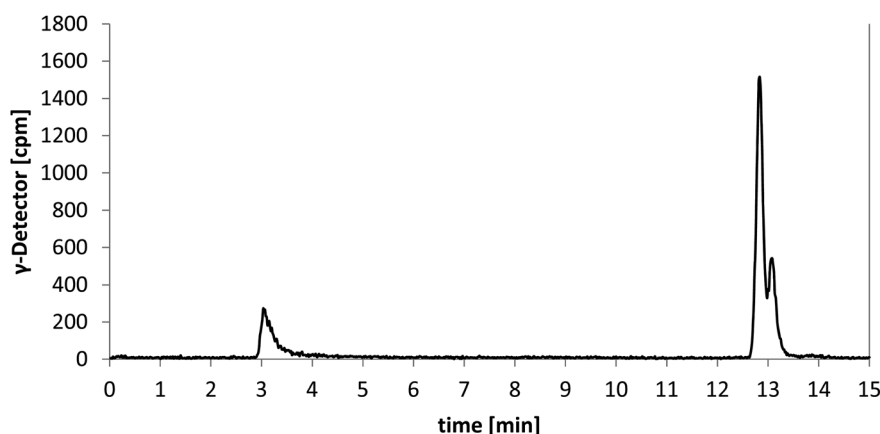
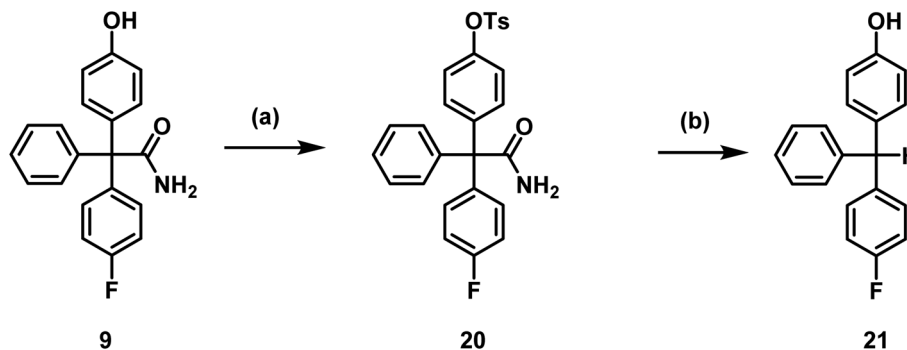


Fig. 3 Chromatogram of the analytical radio-HPLC after 30 min of removal of the solvent.





Scheme 5 Tosylation of phenol **9** and decomposition of tosylate **20**. Reagents and reaction conditions: (a) TsCl, CH<sub>2</sub>Cl<sub>2</sub>, RT, 2 h, 98%, (b) TBAF, DMF, 130 °C, 20 min, 80%.

temperatures between 110 °C and 140 °C with reaction times between 7 and 12 min.<sup>40</sup> The conditions required for the radiosynthesis were tested and are summarized in Table 2.

Tosylate **20** was not stable in the presence of carbonate or tetrabutylammoniumfluoride (TBAF), as a less polar compound was formed. To characterize the decomposition product, the tosylate **20** was heated up to 130 °C for 20 min with TBAF in DMF and the product **21** was isolated by flash chromatography. The obtained decomposition product **21** indicated that hydrolysis of the tosylate and loss of the carbamoyl moiety at the quaternary C-atom had taken place. Hydrolysis and decarboxylation of the amide **20** could lead to the triaryl methane derivative **21** (Scheme 5).

Since the titanium nanoparticle catalyzed nucleophilic substitution of aryl tosylates requires activation of [<sup>18</sup>F]fluoride as [<sup>18</sup>F]TBAF, it was concluded that the tosylate **20** is not a useful precursor for <sup>18</sup>F-labelling. Furthermore, it is impossible to use the heterogeneous system in the automatic synthesis modules, since the solids would clog the thin tubing resulting in limitations in the further transfer of the radiosynthesis into clinical environment.

Literature protocols for <sup>18</sup>F-labelling of aryl pinacol boronic esters use [<sup>18</sup>F]TBAF or tetraethylammonium [<sup>18</sup>F]fluoride ([<sup>18</sup>F]TEAF) and temperatures above 100 °C.<sup>41</sup> It is very unlikely that these conditions are tolerated by the triaryl acetamide structure of the required precursor, since similar conditions led to

decomposition of both precursors **18** and **20**. Because the solvent system differs and the reaction time is relatively short, the pinacol boronic ester **24** should nevertheless be tested as precursor. Therefore, aryl bromide **22** was prepared in four steps analogous to the synthesis of the trimethylammonium precursor **18** using 1,4-dibromobenzene instead of 4-bromo-*N,N*-dimethylaniline (**12**) as starting material (see ESI, Scheme S4†). Finally, the boronic ester **24** was prepared by reaction of aryl bromide **22** with diboronate **23** and ferrocene-1,1'-(diyl) bis(diphenylphosphane)palladium(II)chloride (Pd(dppf)Cl<sub>2</sub>) (Scheme 6).

For Cu(II)-catalyzed transformations of areneboronates into aryl fluorides, azeotropic drying was not reported.<sup>41</sup> Thus, a solution of precursor **24** and tetrakis(pyridine)copper(II)triflate (Cu(OTf)<sub>2</sub>py<sub>4</sub>) in DMF was added to a solution of [<sup>18</sup>F]TEAF in butan-1-ol and the mixture was heated to 110 °C for 10 min. The copper catalyst was removed by a C18+ cartridge and the reaction mixture was analyzed by semipreparative and analytical radio-HPLC. However, a peak showing the formation of [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]**1**) could not be detected. It was concluded that this strategy is not useful for the synthesis of [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]**1**).

Senicapoc (**1**) contains the same triarylacetamide structure as the unstable precursors **18**, **20** and **24**. Since it is essential that senicapoc (**1**) is stable under the conditions used to prepare [<sup>18</sup>F]senicapoc ([<sup>18</sup>F]**1**), different reaction conditions reasonable for <sup>18</sup>F-labelling were evaluated. The results are summarized in Table 3.

Senicapoc (**1**) showed very similar reactivity under the conditions tested as the precursors **18** and **20**. As reported for **18** and **20**, triaryl methane **25** was formed as decomposition product. Removal of the carbamoyl moiety, presumably by hydrolysis and decarboxylation, is an obvious inherent instability of this type of triarylacetamide derivatives under basic conditions (K<sub>2</sub>CO<sub>3</sub>, TBAF) and elevated temperature (Scheme 7).

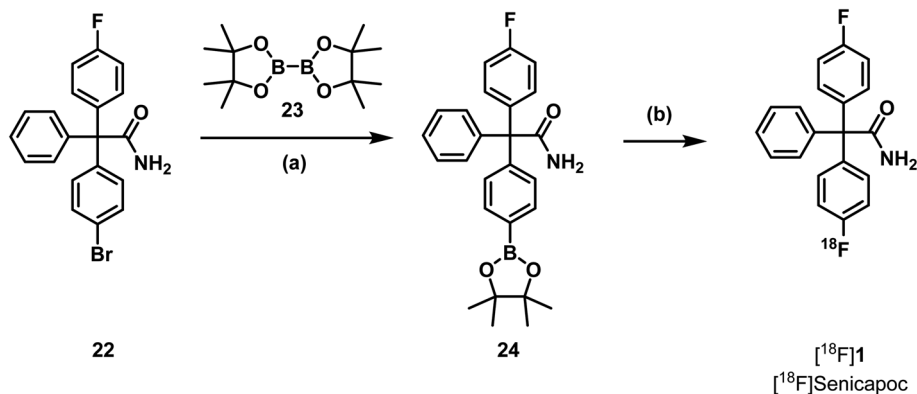
Analysis of the data summarized in Table 3 showed that oxalate is the preferred base to perform a direct nucleophilic fluorination, as decomposition of senicapoc to the methane derivative **25** could not be observed. The base carbonate can be used if the reaction temperature does not exceed 90 °C.

Since aliphatic nucleophilic substitution with [<sup>18</sup>F]fluoride is possible using rather mild conditions, the [<sup>18</sup>F]fluoroethoxy derivative [<sup>18</sup>F]**28** of senicapoc (**1**) was envisaged for imaging

Table 2 Conditions used to evaluate the stability of aryl tosylate precursor **20**; identical conditions for all reactions: reaction time: 10 min; solvent: DMF

Base (5 eq.)	Temperature [°C]	Stability (TLC)
—	130	Stable
TBAF	130	Full conversion
TBAF	110	Full conversion
TBAF	90	Full conversion
K <sub>2</sub> CO <sub>3</sub>	130	Full conversion
K <sub>2</sub> CO <sub>3</sub>	110	Partial conversion
K <sub>2</sub> CO <sub>3</sub>	90	Partial conversion
(NH <sub>4</sub> ) <sub>2</sub> C <sub>2</sub> O <sub>4</sub>	130	Stable
(NH <sub>4</sub> ) <sub>2</sub> C <sub>2</sub> O <sub>4</sub>	110	Stable
(NH <sub>4</sub> ) <sub>2</sub> C <sub>2</sub> O <sub>4</sub>	90	Stable



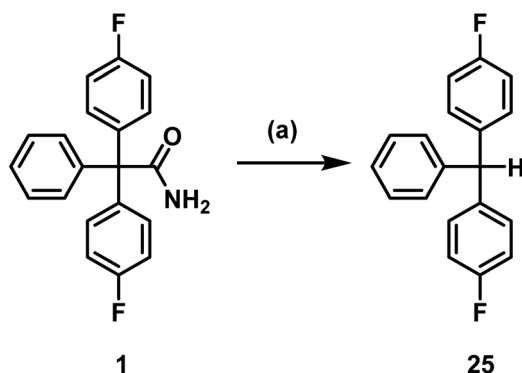


**Scheme 6** Last step of the synthesis of the pinacol boronic ester **24** and radiosynthesis of [ $^{18}\text{F}$ ]senicapoc ( $^{18}\text{F}$ 1). Reagents and reaction conditions: (a) Pd(dppf)Cl<sub>2</sub>, KOAc, dioxane, RT- 50 °C, 5 h, 15%. (b) cartridge: Sep-Pak® Light QMA Fa. Waters®, [ $^{18}\text{F}$ ]TEAF, butan-1-ol, Cu(OTf)<sub>2</sub>py<sub>4</sub>, DMF, 110 °C, 10 min, 0%.

**Table 3** Conditions used to evaluate the stability of senicapoc (**1**); identical conditions for all reactions: reaction time: 10 min; solvent: DMF

Base (5 eq.)	Temperature [°C]	Stability (TLC)
—	130	Stable
TBAF	130	Complete decomposition
TBAF	110	Complete decomposition
TBAF	90	Complete decomposition
K <sub>2</sub> CO <sub>3</sub>	130	Complete decomposition
K <sub>2</sub> CO <sub>3</sub>	110	Partial decomposition
K <sub>2</sub> CO <sub>3</sub>	90	Partial decomposition
(NH <sub>4</sub> ) <sub>2</sub> C <sub>2</sub> O <sub>4</sub>	130	Stable
(NH <sub>4</sub> ) <sub>2</sub> C <sub>2</sub> O <sub>4</sub>	110	Stable
(NH <sub>4</sub> ) <sub>2</sub> C <sub>2</sub> O <sub>4</sub>	90	Stable

of K<sub>Ca</sub>3.1 channels *in vivo*. At first, the non-radioactive  $^{19}\text{F}$ -containing reference compound **28** was synthesized to perform *in vitro* experiments addressing the desired interaction of **28** with the K<sub>Ca</sub>3.1 channel and to identify unequivocally the

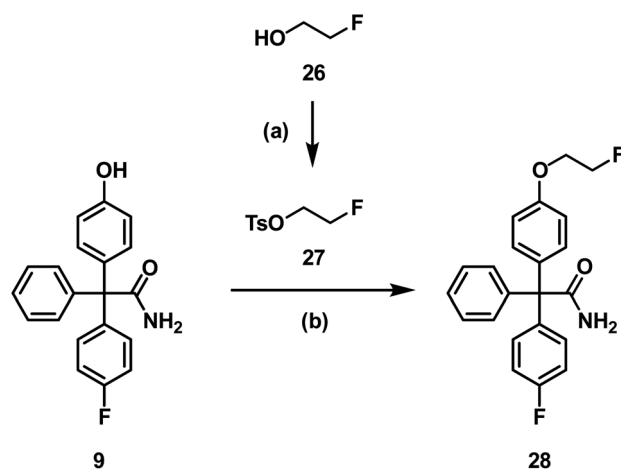


**Scheme 7** Decomposition of senicapoc (**1**). Reagents and reaction conditions: (a) TBAF, DMF, 130 °C, 10 min, 75%.

labelled compound by co-injection on the radio-HPLC system. **28** was synthesized by alkylation of phenol **9** with (2-fluoroethyl) tosylate (**27**),<sup>42</sup> which was prepared from 2-fluoroethanol (**26**) and tosyl chloride (Scheme 8).

At first, the interaction of **28** with the K<sub>Ca</sub>3.1 channel had to be analyzed. For this purpose, patch clamp experiments were performed using the non-small-cell lung cancer cell line A549-3R, which expresses a high number of the K<sub>Ca</sub>3.1 channels (Fig. 4).

A549-3R cells were activated with 1-EBIO leading to an increased outward current in the patch clamp experiments. Then, the fluoroethoxy derivative **28** was added, which led to reduced ionic current. Finally, senicapoc was added resulting in a complete inhibition of the current increase induced by 1-EBIO. The experiment was performed in a paired fashion and showed that **28** was able to inhibit the K<sub>Ca</sub>3.1 channel, although not as efficiently as senicapoc (**1**). The effect of **28** on the ionic current was regarded as interaction with the ion channel. Therefore, imaging of the K<sub>Ca</sub>3.1 channel *in vivo* using [ $^{18}\text{F}$ ]**38** was assumed to be possible.



**Scheme 8** Synthesis of the fluoroethoxy derivative **28**. Reagents and reaction conditions: (a) TsCl, Et<sub>3</sub>N, CH<sub>2</sub>Cl<sub>2</sub>, RT, 24 h, 88%. (b) DMF, Cs<sub>2</sub>CO<sub>3</sub>, 100 °C, 10 °C/h, 84%.



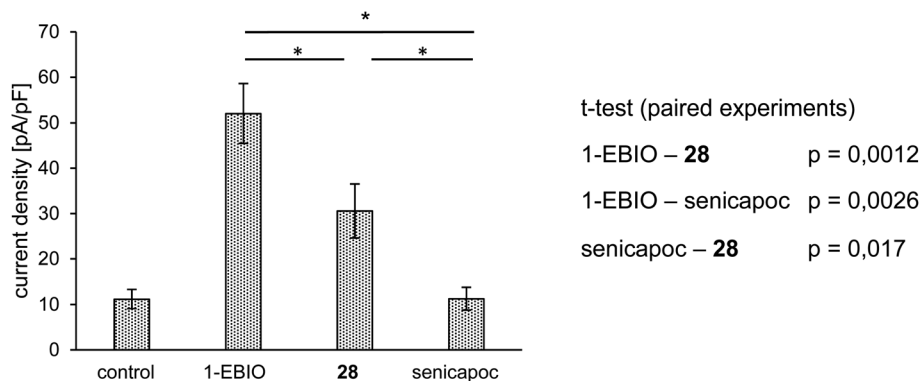


Fig. 4 Summary of patch-clamp experiments. Average current density [pA/pF]  $\pm$  SEM ( $n = 5$ ).  $K_{Ca3.1}$  mediated current is activated with 1-EBIO (50  $\mu$ M). It is inhibited by the consecutive application of 28 (10  $\mu$ M) and senicapoc (1  $\mu$ M). All compounds were dissolved in DMSO and added to the standard extracellular solution and applied for a period of 2–5 min (1-EBIO = 1-ethyl-2-benzimidazolone).

Consequently, radiosynthesis of [ $^{18}$ F]28 using precursor 31 bearing a tosyloxy leaving group for aliphatic nucleophilic substitution with [ $^{18}$ F]fluoride under mild conditions was planned. Thus, phenol 9 was directly alkylated with ethylene ditosylate (30),<sup>43</sup> which was obtained by ditosylation of ethylene glycol (29) with an excess of tosyl chloride (Scheme 9).

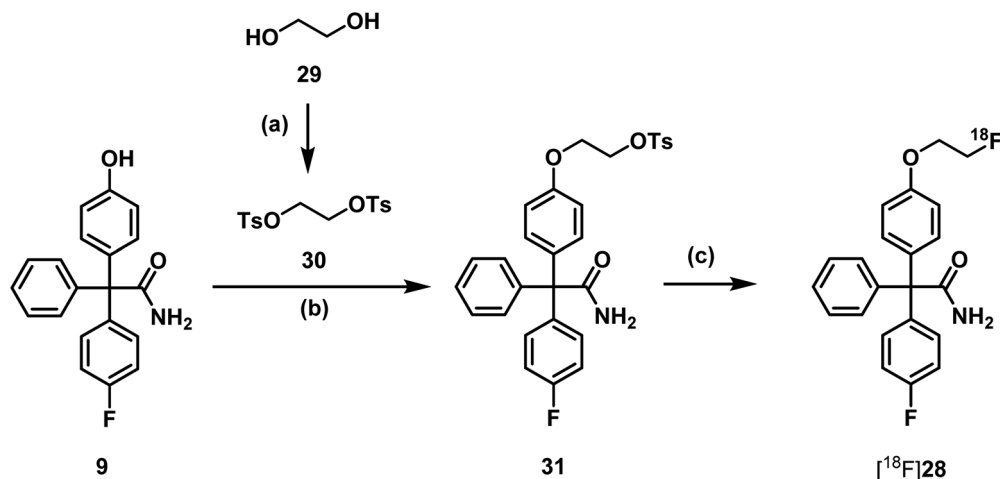
Since it had been found that carbonate leads to degradation of triarylacetamides, oxalate was used as base in the first  $^{18}$ F-labelling experiments. The radiosynthesis was carried out in a semiautomated radiosynthesizer. The aqueous solution of [ $^{18}$ F]fluoride ions was rinsed through a preconditioned QMA-light cartridge (Waters®). The trapped [ $^{18}$ F]fluoride ions were eluted with a solution of  $K_{222}$  and  $K_2CO_3$  in  $CH_3CN$  and  $H_2O$ . After azeotropic drying a solution of the precursor 31 in DMF was added to the radiosynthesizer. The mixture was heated to 90  $^{\circ}C$  for 15 min in the closed reactor (Scheme 9).

The desired product [ $^{18}$ F]28 was isolated by semi-preparative radio-HPLC. Identity was confirmed by coinjection of the corresponding reference compound 28 using the analytical HPLC-system (for chromatogram see ESI: Fig. S5<sup>†</sup>).

After removal of the solvent under reduced pressure, the product [ $^{18}$ F]28 was reconstituted with ethanol. PBS was added to achieve an ethanol concentration below 10%. This ready-to-inject solution was controlled again by radio-HPLC as well as radio-TLC. The decay corrected radiochemical yield of the formulated PET tracer was  $4.0 \pm 1.5\%$ , radiochemical purity was found to be  $>96\%$ .

*In vitro* serum stability was tested using murine and human blood [ $^{18}$ F]28 showed high stability over the complete test period of 120 min. The corresponding chromatograms are shown in the ESI (Fig. S6 and S7<sup>†</sup>). The  $\log D_{7,4}$ -value was determined to be  $1.84 \pm 0.5$  ( $\pm$ SEM) using the shake-flask method by Prante *et al.*<sup>44</sup>

Before performing biodistribution experiments *in vivo*, the plasma protein binding (PPB) of senicapoc (1) and its fluoroethoxy derivative 28 were determined *in vitro* using a LC-MS method with external calibration. The PPB has a considerable influence on both distribution and metabolic stability of a drug *in vivo*. A high PPB ensures good distribution and long circulation in the blood as well as low metabolic degradation *in vivo*. On the other side, only the unbound fraction of the drug can



Scheme 9 Synthesis of precursor 31 and radiosynthesis of [ $^{18}$ F]fluoroethoxy senicapoc [ $^{18}$ F]28. Reagents and reaction conditions: (a) TsCl,  $Et_3N$ ,  $CH_2Cl_2$ , RT, 12 h, 97%. (b) DMF,  $Cs_2CO_3$ , RT, 24 h, 49%. (c) [ $^{18}$ F]KF· $K_{222}$ ,  $CH_3CN$ , 90  $^{\circ}C$ , 15 min, RCY (d.c.) =  $4 \pm 1.5\%$ .



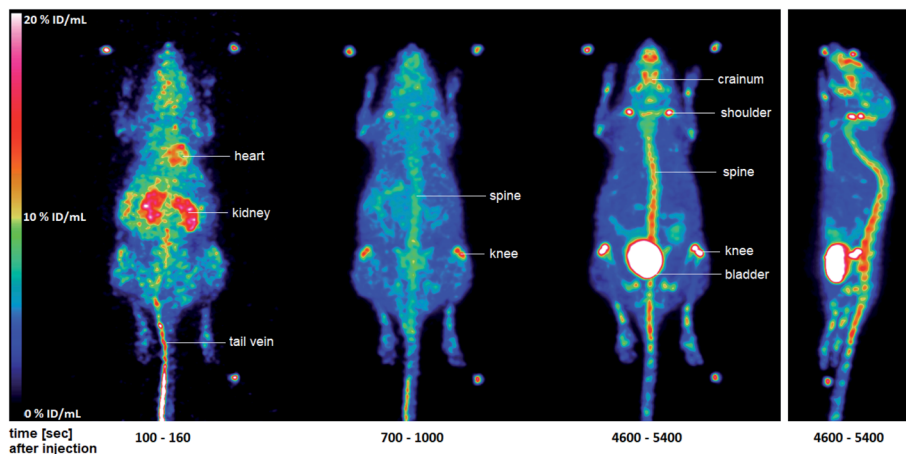


Fig. 5 Maximum-intensity-projection of the activity distribution of selected time intervals in a C57BL/6 wild-type mouse after injection of  $[^{18}\text{F}]28$ . The two additional images of mice measured may be found in the ESI (Fig. S8†).

enter the tissue and cells and thus bind to the respective target. It was found that both senicapoc (**1**) and its fluoroethoxy derivative **28** have a PPB of >99%.

Due to the high PPB, a good biodistribution of  $[^{18}\text{F}]28$  *in vivo* was expected. Biodistribution studies with the PET tracer  $[^{18}\text{F}]28$  in mice will show undesired possible accumulation in different tissues, which *e.g.* also physiologically express the target of the PET tracer, or non-specific binding to other proteins. Moreover, biodistribution studies show the route of excretion of the tracer. Altogether, analysis of the *in vivo* biodistribution of a PET tracer provides important data, which are crucial for the further development of a PET tracer for imaging studies.

In this study, three 12 week-old C57BL/6 wild-type mice ( $n = 3$ ) were used. The experiments were approved by the State Office for Nature, Environment and Consumer Protection North Rhine-Westphalia under license Az. 84–02.04.2015.A410. The mice were anesthetized with isoflurane and placed on a warmed PET scanner bed. Important parameters such as breathing were constantly monitored. The formulated PET tracer  $[^{18}\text{F}]28$  was administered *via* a tail vein catheter and with the aid of an injection pump. After 90 min acquisition time in the quadHIDAC PET scanner, a CT scan of the mouse was performed. The data and images were evaluated with the in-house developed software MEDgical (Fig. 5; please also refer to Fig. S8 in the ESI†).

Fig. 5 shows the distribution of radioactivity expressed as percentage of the injected dose per mL (% ID/mL) in whole-body images after injection of  $[^{18}\text{F}]28$  in a representative mouse. After a few seconds, a large amount of the injected dose (ID) was still found in the tail vein  $[^{18}\text{F}]28$  also reached the heart *via* the blood. A significant amount of ID was already seen in the kidneys at this early stage. After 700–1000 s, accumulation in the bones, especially in the spine and knees, was detected. This accumulation increased substantially over the time. After 90 min, the highest amount of the ID was found in the bladder indicating renal elimination of the radioactive compound. However, large amounts of ID were also found in the spine, knees, shoulders, and cranium. The accumulation in the bones is probably due to the formation of  $[^{18}\text{F}]$ fluoride. This finding indicates a metabolic instability of the tracer  $[^{18}\text{F}]28$  *in vivo*, which was not observed during the *in vitro* serum stability tests. Due to this metabolic instability, the PET tracer  $[^{18}\text{F}]28$  was not suitable for further tumor studies.

In order to analyze the metabolic instability of the tracer  $[^{18}\text{F}]28$  in more detail and provide information about possible degradation pathways, an *in vitro* metabolism study was conducted with the non-radioactive reference compound **28**. Therefore, **28** was incubated with mouse liver microsomes and NADPH for 90 min at 37 °C. The transformation was stopped by the addition of  $\text{CH}_3\text{CN}/\text{CH}_3\text{OH}$  (1 : 1), the solution was cooled

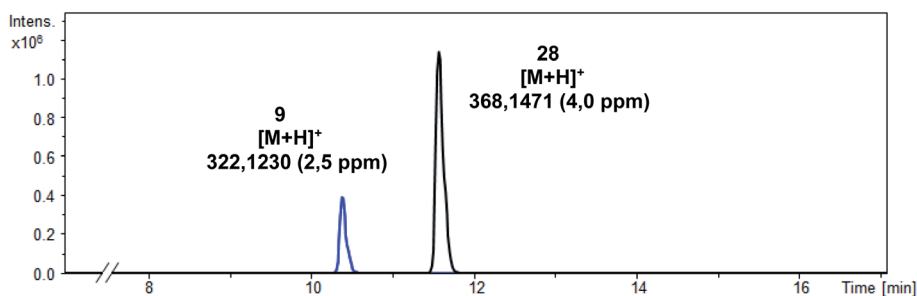
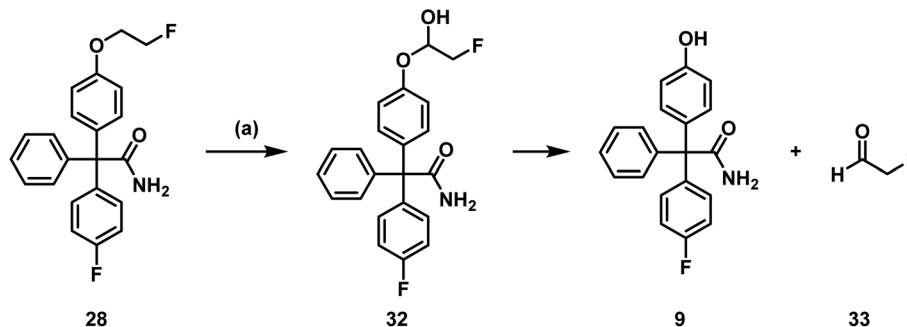


Fig. 6 LC-MS/MS chromatogram of the supernatant.







Scheme 10 Postulated metabolism pathway of **28**. (a) Incubation with NADPH and murine liver microsomes, 37 °C, 90 min.

and then centrifuged to remove the protein precipitate. The supernatant was analyzed by LC-MS/MS (qToF) (Fig. 6).

In addition to the intact parent compound **28**, a more polar metabolite with  $m/z = 322.1230$  was detected. Since ESI is a soft ionization method, only limited fragmentation reactions were expected. Therefore, the peak of the metabolite probably corresponds to  $[M + H]^+$  indicating the formation of phenol **9** (calculated exact mass of  $[M + H]^+$ : 322.1238) as main metabolite (Scheme 10).

Phenol **9** can be formed by oxidation of the fluoroethoxy side chain of **28** by CYP450 enzymes and NADPH resulting in the labile hemiacetal **32**, which releases the phenol **9** and 2-fluoroacetaldehyde (**33**) spontaneously. 2-Fluoroacetaldehyde or its oxidation product 2-fluoroacetic acid can release fluoride due to the high reactivity of the F-atom in  $\alpha$ -position of the carbonyl moiety.

### 3. Conclusion

Introduction of  $[^{18}\text{F}]$ fluoride in the aromatic ring resulting in  $[^{18}\text{F}]$ senicapoc ( $[^{18}\text{F}]\mathbf{1}$ ) required high temperature and strong bases. Unfortunately, the triarylamide moiety was not stable under these conditions leading to triarylmethane derivatives. Thus, only traces of  $[^{18}\text{F}]$ senicapoc ( $[^{18}\text{F}]\mathbf{1}$ ) were obtained by concerted nucleophilic substitution using the uronium salt precursor **10**.

As an alternative PET tracer targeting  $\text{K}_{\text{Ca}3.1}$  channels the  $[^{18}\text{F}]$ fluoroethoxy derivative  $[^{18}\text{F}]\mathbf{28}$  was designed for imaging *in vivo*. Affinity towards the ion channel was shown in patch clamp experiments using the non-radioactive reference compound **28**. Radiosynthesis of  $[^{18}\text{F}]\mathbf{28}$  was performed successfully by aliphatic nucleophilic substitution of the tosylated precursor **31** with  $[^{18}\text{F}]$ fluoride under mild conditions  $[^{18}\text{F}]\mathbf{28}$  was obtained with a decay corrected radiochemical yield of  $4 \pm 1.5\%$  ( $n = 4$ ) within  $106 \pm 11.5$  min ( $n = 4$ ) and a radiochemical purity of  $>96\%$  by analytical radio-HPLC and radio-TLC.

Over a period of 120 min,  $[^{18}\text{F}]\mathbf{28}$  was stable in both human and mouse serum. The  $\log D_{7.4}$  value of 1.84 lies in a promising range. More than 99% of the drug was bound to human serum albumin. During biodistribution studies with  $[^{18}\text{F}]\mathbf{28}$  *in vivo* using C57/BL6 wild-type mice, accumulation of activity in the bones was observed indicating the release of  $[^{18}\text{F}]$ fluoride from the tracer. In order to confirm the degradation of  $[^{18}\text{F}]\mathbf{28}$  *in vivo*, an *in vitro* metabolism study was conducted with **28**. The phenol

**9** was detected as main metabolite, which was formed upon release of  $[^{18}\text{F}]$ 2-fluoroacetaldehyde. The release of  $[^{18}\text{F}]$ 2-fluoroacetaldehyde could explain the enrichment of  $[^{18}\text{F}]$ fluoride in the bones. This finding is of general interest, since the  $[^{18}\text{F}]$ 2-fluoroethyl moiety is a well-known prosthetic group often used in PET tracers.<sup>45,46</sup> The potential degradation to afford  $[^{18}\text{F}]$ 2-fluoroacetaldehyde should be generally taken into account, when designing novel PET tracers with the  $[^{18}\text{F}]$ 2-fluoroethyl moiety or using the popular prosthetic group  $[^{18}\text{F}]$ fluoroethyl tosylate to alkylate phenols.

### Conflict of interests

The authors have no conflicts of interest to declare.

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### Notes and references

- Z. H. Yu, J. R. Xu, Y. X. Wang, G. N. Xu, Z. P. Xu, K. Yang, D. Z. Wu, Y. Y. Cui and H. Z. Chen, *Am. J. Respir. Cell Mol. Biol.*, 2013, **48**, 685–693.
- P. Bradding and H. Wulff, *Br. J. Pharmacol.*, 2009, **157**, 1330–1339.
- K. I. Ataga and J. Stocker, *Expert Opin. Invest. Drugs*, 2009, **18**, 231–239.
- K. I. Ataga, W. R. Smith, L. M. de Castro, P. Swerdlow, Y. Sauntharajah, O. Castro, E. Vichinsky, A. Kutlar,



- E. P. Orringer, G. C. Rigdon and J. W. Stacker, *Blood*, 2008, **111**, 3991–3997.
- 5 K. I. Ataga, E. P. Orringer, L. Styles, E. P. Vichinsky, P. Swerdlow, G. A. Davis, P. A. DeSimone and J. W. Stocker, *Pharmacotherapy*, 2006, **26**, 1557–1564.
- 6 D. L. Tharp, B. R. Wamhoff, H. Wulff, G. Raman, A. Cheong and D. K. Bowles, *Arterioscler., Thromb., Vasc. Biol.*, 2008, **28**, 1084–1089.
- 7 I. Grgic, E. Kiss, B. P. Kaistha, C. Busch, M. Kloss, J. Sautter, A. Müller, A. Kaistha, C. Schmidt, G. Raman, H. Wulff, F. Strutz, H. J. Gröne, R. Köhler and J. Hoyer, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 14518–14523.
- 8 C. Huang, S. Shen, Q. Ma, J. Chen, A. Gill, C. A. Pollock and X. M. Chen, *Diabetes*, 2013, **62**, 2923–2934.
- 9 R. Xu, C. Li, Y. Wu, L. Shen, J. Ma, J. Qian and J. Ge, *Arterioscler., Thromb., Vasc. Biol.*, 2017, **37**, 226–236.
- 10 Y. R. Zhu, X. X. Jiang and D. M. Zhang, *J. Mol. Med.*, 2019, **97**, 1219–1229.
- 11 G. Anumanthan, S. Gupta, M. K. Fink, N. P. Hesemann, D. K. Bowles, L. M. McDaniel, M. Muhammad and R. R. Mohan, *PLoS One*, 2018, **13**(3), e0192145.
- 12 G. Anumanthan, P. J. Wilson, R. Tripathi, N. P. Hesemann and R. R. Mohan, *Exp. Eye Res.*, 2018, **167**, 140–144.
- 13 L. Jin, J. di Lucente, H. M. Nguyen, V. Singh, L. Singh, M. Chavez, T. Bushong, H. Wulff and I. Maezawa, *Ann. Clin. Transl. Neurol.*, 2019, **6**, 723–738.
- 14 I. Maezawa, D. P. Jenkins, B. E. Jin and H. Wulff, *Int. J. Alzheimer's Dis.*, 2012, 2012.
- 15 M. Yi, P. Yu, Q. Lu, H. M. Geller, Z. Yu and H. Chen, *Mol. Cell. Neurosci.*, 2016, **76**, 21–32.
- 16 L. Sevelsted Møller, A. D. Fialla, R. Schierwagen, M. Biagini, C. Liedtke, W. Laleman, S. Klein, W. Reul, L. Koch Hansen, M. Rabjerg, V. Singh, J. Surra, J. Osada, R. Reinehr, O. B. S. de Muckadell, R. Köhler and J. Trebicka, *Sci. Rep.*, 2016, **6**, 28770.
- 17 Z. Pethő, K. Najder, E. Bulk and A. Schwab, *Cell Calcium*, 2019, **80**, 79–90.
- 18 C. J. Mohr, D. Gross, E. C. Sezgin, F. A. Steudel, P. Ruth, S. M. Huber and R. Lukowski, *Cancers*, 2019, **11**, 1285.
- 19 N. Prevarskaya, R. Skryma and Y. Shuba, *Physiol. Rev.*, 2018, **98**, 559–621.
- 20 K. L. Turner, A. Honasoge, S. M. Robert, M. M. McFerrin and H. Sontheimer, *Glia*, 2014, **62**, 971–981.
- 21 M. Rabjerg, A. Oliván-Viguera, L. K. Hansen, L. Jensen, L. Sevelsted-Møller, S. Walter, B. L. Jensen, N. Marcussen and R. Köhler, *PLoS One*, 2015, **10**, e0122992.
- 22 M. Faouzi, F. Hague, D. Geerts, A. S. Ay, M. Potier-Cartereau, A. Ahidouch and H. Ouadid-Ahidouch, *Oncotarget*, 2016, **7**, 36419–36435.
- 23 N. Haren, H. Khorsi, M. Faouzi, A. Ahidouch, H. Sevestre and H. Ouadid-Ahidouch, *Histol. Histopathol.*, 2010, **25**, 1247–1255.
- 24 E. Bulk, A.-S. Ay, M. Hammadi, H. Ouadid-Ahidouch, S. Schelhaas, A. Hascher, C. Rohde, N. H. Thoennissen, R. Wiewrodt, E. Schmidt, A. Marra, L. Hillejan, A. H. Jacobs, H.-U. Klein, M. Dugas, W. E. Berdel, C. Müller-Tidow and A. Schwab, *Int. J. Cancer*, 2015, **137**, 1306–1317.
- 25 J. H. Ko, E. A. Ko, W. Gu, I. Lim, H. Bang and T. Zhou, *Mol. Cancer*, 2013, **12**, 106.
- 26 F. A. Steudel, C. J. Mohr, B. Stegen, H. Y. Nguyen, A. Barnert, M. Steinle, S. Beer-Hammer, P. Koch, W. Y. Lo, W. Schroth, R. Hoppe, H. Brauch, P. Ruth, S. M. Huber and R. Lukowski, *Mol. Oncol.*, 2017, **11**, 1172–1188.
- 27 H. Wulff and N. A. Castle, *Expert Rev. Clin. Pharmacol.*, 2010, **3**, 385–396.
- 28 C. Gross, *Epilepsy Currents*, 2020, **20**, 211–213.
- 29 S. Sugunan, S. Nampoothiri, T. Garg and R. Krishnamurthy, *CNS Neurol. Disord.: Drug Targets*, 2016, **15**, 1299–1305.
- 30 V. Hofschroer, K. Najder, M. Rugi, R. Bouazzi, M. Cozzolino, A. Arcangeli, G. Panyi and A. Schwab, *Front. Pharmacol.*, 2021, **11**, 1979.
- 31 J. W. Stocker, L. de Franceschi, G. A. McNaughton-Smith, R. Corrocher, Y. Beuzard and C. Brugnara, *Blood*, 2003, **101**, 2412–2418.
- 32 B. I. Mobebe, S. Venkatraman, G. McNaughton-Smith, C. Gibb, L. G. Ulysse, C. A. Lindmark, S. Shaw, B. Marron, K. Spear and M. J. Suto, *Org. Process Res. Dev.*, 2012, **16**, 1385–1392.
- 33 C. N. Neumann, J. M. Hooker and T. Ritter, *Nature*, 2016, **534**, 369–373.
- 34 T. Fujimoto, F. Becker and T. Ritter, *Org. Process Res. Dev.*, 2014, **18**, 1041–1044.
- 35 K. Brömmel, S. Maskri, I. Maisuls, C. P. Konken, M. Rieke, Z. Pethő, C. A. Strassert, O. Koch, A. Schwab and B. Wünsch, *Angew. Chem., Int. Ed.*, 2020, **59**, 8277–8284.
- 36 M. Lasch, A. Caballero Martinez, K. Kumaraswami, H. Ishikawa-Ankerhold, S. Meister and E. Deindl, *Cells*, 2020, **9**(4), 913.
- 37 S. A. Nicolaou, L. Neumeier, Y. Q. Peng, D. C. Devor and L. Conforti, *Am. J. Physiol.: Cell Physiol.*, 2007, **292**(4), 1431–1439.
- 38 E. Cole, M. Stewart, R. Littich, R. Hoareau and P. Scott, *Curr. Top. Med. Chem.*, 2014, **14**, 875–900.
- 39 P. W. Miller, N. J. Long, R. Vilar and A. D. Gee, *Angew. Chem., Int. Ed.*, 2008, **47**, 8998–9033.
- 40 M. E. Sergeev, F. Morgia, M. Lazari, C. Wang and R. M. van Dam, *J. Am. Chem. Soc.*, 2015, **137**, 5686–5694.
- 41 M. Tredwell, S. M. Preshlock, N. J. Taylor, S. Gruber, M. Huiban, J. Passchier, J. Mercier, C. Génicot and V. Gouverneur, *Angew. Chem., Int. Ed.*, 2014, **53**, 7751–7755.
- 42 F. Dollé, F. Hinnen, A. Damont, B. Kuhnast, C. Fookes, T. Pham, B. Tavitian and A. Katsifis, *J. Labelled Compd. Radiopharm.*, 2008, **51**, 435–439.
- 43 S. Comagic and R. Schirmacher, *Synthesis*, 2004, **2004**, 885–888.
- 44 O. Prante, C. Hocke, S. Löber, H. Hübner, P. Gmeiner and T. Kuwert, *Nuklearmedizin*, 2006, **45**, 41–48.
- 45 W. Mohnike, G. Hör, A. Hertel, H. Schelbert, *PET/CT-Atlas*, Springer, Berlin, Heidelberg, 2016.
- 46 Z. Li and P. S. Conti, *Adv. Drug Delivery Rev.*, 2010, **62**, 1031–1051.

