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Fluorination of pyrrole derivatives by Selectfluor™

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ABSTRACT

Fluorination of a range of pyrrole substrates bearing various electron donating and withdrawing substituents at the 1-, 2- and 3-positions using Selectfluor™ has been assessed in order to develop effective methodology for the synthesis of corresponding fluoropyrrole derivatives. The synthesis of some novel fluorinated pyrrole derivatives were achieved in reasonable quantities although many pyrrole substrates were oxidatively polymerised by the fluorinating reagent, reducing the scope of a selective fluorination approach for the preparation of fluoropyrrole products.

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1. Introduction

Since the discovery of the bioactivity of 9 α -fluorocortisoid derivatives by Fried and Sabo in 1954,¹ fluorine and fluorinated groups have been used extensively in drug design for many reasons including changing lipophilicity, enhancing metabolic stability and controlling pH profiles.^{2–6} Consequently, fluorine-containing drugs are now prevalent within the life science industries and a considerable number of blockbuster pharmaceuticals and agrochemicals now contain at least one fluorine atom.^{7–9}

In particular, fluorinated azaheterocycles are very important subunits within life science products and a growing number of pharmaceuticals that contain fluoroheteroaromatic motifs such as Voriconazole (antifungal, Pfizer), Capecitabine (anticancer, Roche) and Diclosulam (herbicide, Dow) have now reached the commercial market^{10–12} with many others in clinical trials such as Abemaciclib (anticancer, Eli Lilly), Riociguat (heart failure, Bayer) and Verubecestat (Alzheimers, Merck).¹³ Notably, these pharmaceuticals all contain a fluorinated six-membered azaheterocycle, such as fluoro-pyridine or -pyrimidine structural subunit, and effective synthetic procedures at both discovery and manufacturing scales are well reported in the literature.^{14–21} In contrast, it is particularly striking that pharmaceuticals bearing related five-membered azaheterocycles, such as fluorinated pyrroles, furans and thiophenes, are very rarely found in life science products and this is, in part, due to the lack of available, efficient and regioselective methodology for

the synthesis of, for example, appropriate fluoropyrrole derivatives both for medicinal chemistry programmes and large scale synthesis.

Of relevance to this study, the few reported methods for the synthesis of fluorinated pyrroles²² that have been published involve the use of fluorinated building blocks in cyclisation reactions such as the cyclocondensation of α,α -difluoro- γ -iododicarbonyl compounds with ammonia²³ and various primary amines,²⁴ related cyclisation reactions of α,α -difluoro- γ -iodotrimethylsilyl ketones,^{25,26} rhodium catalysed intramolecular N–H insertion of 5-amino-4,4-difluoro-2-diazo-3-ketesters,²⁷ silver catalysed aminofluorination of activated allenes,²⁸ 1,3-dipolar cycloaddition of DMAD with azomethine ylides formed by reaction of imines and difluorocarbene,^{29,30} the reaction of *gem*-difluorocyclopropyl ketones with nitriles in the presence of trifluoromethanesulfonic acid³¹ and ring closing metathesis of fluoroamidoalkene derivatives followed by alkylation-aromatisation.³² All cyclisation methods provide access to fluoropyrrole products but the range of systems possible are limited by the structural complexity of the fluorinated building blocks and so corresponding synthesis of libraries of functional fluoro-pyrroles for drug screening programs are therefore limited. Other approaches that require pre-functionalisation of pyrrole derivatives and reaction with a suitable fluorinating agent include a photochemical modification of the Balz–Schiemann reaction,³³ fluorodecarboxylation of highly substituted derivatives using Selectfluor™,³⁴ and also the reaction of Grignard³⁵ or lithiated³⁶ intermediates with NFSI, formed from corresponding brominated pyrrole precursors. These methods rely on multi-step access to appropriately functionalised pyrroles which may be

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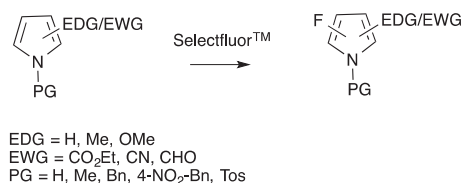
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difficult to prepare efficiently and, consequently, have not been developed to any great extent.

The most direct method for the synthesis of fluoropyrroles is the transformation of C–H to C–F bonds by reaction of the parent pyrrole with an electrophilic fluorinating agent. Electrophilic bromination and chlorination reactions of pyrroles are very well established where NBS, NCS, Br₂ or Cl₂ give simple access to a variety of halogenated pyrrole scaffolds.³⁷ However, related fluorination processes for the preparation of fluoropyrroles are very rare with only two reports published in which electrophilic fluorinating agents XeF₂^{38,39} and, for a limited number of ester derivatives, SelectfluorTM are employed.⁴⁰ Similarly, fluorination of pyrrole substrates utilising a nucleophilic source of fluorine is also very limited with only one example in which Et₃N·2HF is used in anodic fluorination reactions but these generally result in low yields and significant amounts of polyfluorinated by-products.⁴¹ This is despite the obvious potential utility of late stage fluorination of pyrroles for incorporation into drug discovery programmes.

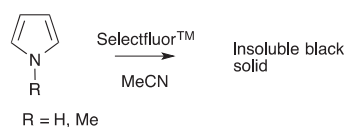
In this paper, we describe our attempts to develop general selective fluorination strategies for the synthesis of functionalised fluoropyrrole derivatives for use in life-science discovery laboratories. For this purpose, SelectfluorTM, a shelf stable, commercially available fluorinating agent of the N–F class was used.⁴² We aimed to assess a model range of pyrrole substrates, bearing electron-donating and -withdrawing substituents and protecting groups attached to the pyrrole ring nitrogen (Scheme 1), that could be efficiently and regioselectively fluorinated using SelectfluorTM, with a view to establishing the regioselectivity of electrophilic fluorination processes and providing access to a range of polyfunctional fluoropyrroles for incorporation into pharmaceutical screening libraries (Scheme 1).



Scheme 1. Proposed synthesis of fluoropyrrole derivatives.

2. Results and discussion

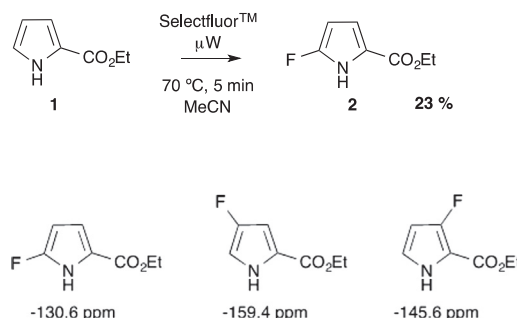
We began our investigations by attempting the fluorination of pyrrole and *N*-methylpyrrole using SelectfluorTM by reaction in acetonitrile but all attempts utilising various conditions gave an insoluble black solid which we attribute to the formation of appropriate poly(pyrrole) derivatives. Pyrrole and related systems are readily polymerised when in contact with oxidising systems such as H₂O₂, *m*CPBA, O₃, AgNO₃, and FeCl₃,^{43,44} and it appears that SelectfluorTM is sufficiently oxidising to allow the formation of polymer products to occur. SelectfluorTM has been reported to oxidise benzyl alcohols to corresponding aldehydes,⁴⁵ consistent with these observations (Scheme 2).



Scheme 2. Attempted synthesis of fluoropyrrole derivatives using SelectfluorTM.

In light of this, we switched our focus to the fluorination of pyrrole derivatives bearing an electron withdrawing substituent to

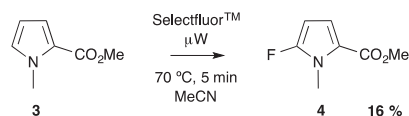
lower the oxidation potential of the pyrrole substrate in order to limit competing oxidation reactions. Reaction of pyrrole-2-carboxylate **1** was attempted using microwave irradiation which resulted in the formation of a number of fluorinated products and some insoluble polymeric material with an appreciable amount of starting material remaining. From the crude product mixture, however, it was possible to isolate monofluorinated pyrrole derivative (**2**) in 23% yield (Scheme 3) by purification using column chromatography.



Scheme 3. Synthesis of ethyl 5-fluoropyrrole-2-carboxylate **2** and DFT predicted ¹⁹F chemical shifts of fluoropyrrole isomers.

DFT computations⁴⁶ were performed to predict the ¹⁹F chemical shifts of the three possible fluorinated pyrrole isomers and, therefore, determine the regioselectivity of the fluorination reaction. The calculations (Scheme 3) show that the predicted shift for fluorine attached to the carbon atom adjacent to the ring nitrogen (–130.6 ppm) is in excellent agreement with the measured value (–130.73 ppm). In addition, the ¹⁹F chemical shifts of the three possible difluorinated isomers were also calculated, confirming that the 4,5-difluorinated derivative is the minor product formed.

Subsequently, *N*-methyl protected pyrrole-2-carboxylate derivative (**3**) was employed as the substrate but, under the same conditions as above, gave a much more complex mixture of fluorinated products due to the increased nucleophilicity of the substrate. Nevertheless, it was possible to isolate the monofluorinated pyrrole derivative (**4**), in 16% yield (Scheme 4). The lower yield was attributed to a combination of loss of material during solvent removal due to the high volatility of the product and the greater number of by-products observed by ¹⁹F NMR spectroscopy. The reaction was also performed at room temperature and using conventional heating in an attempt to decrease the formation of by-products but, unfortunately, many fluorinated products were observed at reasonable conversion.



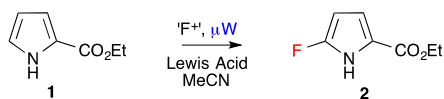
Scheme 4. Synthesis of methyl 1-methyl-5-fluoropyrrole-2-carboxylate.

In attempts to improve the regioselectivity of these processes, fluorination of pyrrole-2-carboxylate derivative (**1**) was screened using a range of Lewis acid catalysts (Table 1). However, the addition of all Lewis acids screened lowered the observed yields of fluoropyrrole product, when either SelectfluorTM or related electrophilic fluorinating agent *N*-fluoro-benzene sulfonimide (NFSI)⁴⁷ were used.

Consequently, due to the highly nucleophilic nature of pyrrole derivatives and the oxidising ability of SelectfluorTM, we chose to

Table 1

Reaction of **1** with electrophilic fluorinating agent in the presence of a Lewis acid under microwave conditions^a



Entry	Fluorinating agent	Lewis acid	NMR Yield/% ^b
1	Selectfluor™	—	21
2	Selectfluor™	ZrCl ₄	8
3	Selectfluor™	AgNO ₃	2
4	Selectfluor™	Ga(OTf) ₃	10
5	Selectfluor™	BF ₃ ·Et ₂ O	6
6	Selectfluor™	HfCl ₄	9
7	Selectfluor™	InCl ₃	12
8	NFSI	—	4
9	NFSI	ZrCl ₄	<1

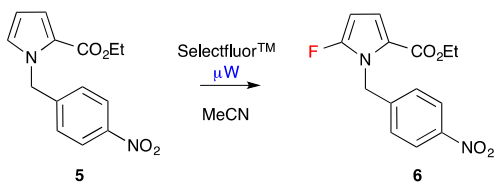
^a Conditions: **1** (1.5 mmol), 'F⁺' reagent (1 equiv), Lewis acid (0.5 equiv), MeCN (15 mL), 70 °C, 5 min.

^b NMR yield determined using an α,α,α -trifluorotoluene reference.

assess fluorination reactions of pyrrole-2-carboxylates bearing electron withdrawing protecting groups attached to ring nitrogen, in an attempt to improve the regioselectivity and yield of fluoropyrrole product. Therefore, the synthesis of 4-nitrobenzyl protected pyrrole derivative (**5**) was carried out by reaction of nitrobenzyl bromide and pyrrole **1** and the product successfully isolated in 51% yield. Fluorination of nitrobenzyl pyrrole **5** was

Table 2

Reaction of **5** with Selectfluor™ under microwave conditions^a



Entry	Temp/°C	Time/min	Yield/% ^b
1	70	5	27 (15)
2	70	10	21
3	70	15	20
4	70	2	23
5	100	5	22
6	100	10	20
7	100	15	20
8	60	5	20
9	60	10	25
10	60	15	25

^a Conditions: **5** (0.5–0.6 mmol), Selectfluor™ (1 equiv), MeCN (15 mL).

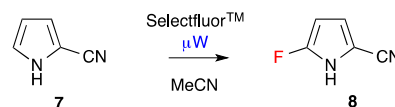
^b NMR yield determined using an α,α,α -trifluorotoluene reference, isolated yield in parentheses.

assessed and a range of reaction conditions were screened (Table 2). From the initial starting point (entry 1), the time of reaction was increased but no improvement in yield was observed despite significant quantities of starting material remaining. Hence, the reaction time was lowered (entry 4) but, again, no improvement in yield was observed. The reaction temperature was varied but neither an increase or decrease in temperature, over a range of reaction times, gave increased yields of fluoropyrrole product and, in all cases, the yield of 5-fluoropyrrole product remained low (~20%). Despite this, the desired product (**6**) could be isolated by column chromatography in 15% yield.

We next studied reaction of a pyrrole derivative bearing a nitrile substituent which could be transformed to a number of substrates of synthetic versatility post fluorination. Initially, the unprotected pyrrole-2-carbonitrile derivative (**7**) was used as substrate for a similar screen of fluorination conditions (Table 3)

Table 3

Reaction of **7** with Selectfluor™ under microwave conditions^a



Entry	Temp/°C	Time/min	Yield/% ^b
1	70	5	62 (30)
2	70	10	46
3	70	2	38
4	60	5	40
5	60	10	53
6	60	15	45
7	100	2	45
8	100	5	47

^a Conditions: **7** (2–10 mmol), Selectfluor™ (1 equiv), MeCN (15–25 mL).

^b NMR yield determined using an α,α,α -trifluorotoluene reference, isolated yield in parentheses.

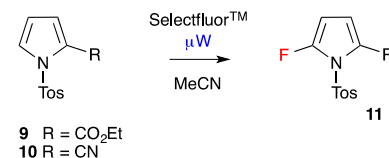
and in this case, conditions shown in entry 1 gave a much-improved yield of 62%.

However, variations in the reaction time and temperature (entries 2 and 3) resulted in reduced yields. The desired product (**8**) could however, be purified by column chromatography and was isolated in a 30% yield. Similarly, the 4-nitrobenzyl substituted pyrrole-2-carbonitrile derivative was again synthesised but subsequent related fluorination reactions gave no improvement in yield.

Due to the limited success observed when employing the 4-nitrobenzyl group, the tosyl group was selected as ring nitrogen protecting group due to its strong electron-withdrawing ability, stability to oxidising conditions and ease of introduction and re-

Table 4

Reaction of **9** with Selectfluor™ under microwave conditions^a



Entry	R	Temp/°C	Time/min	Yield/% ^b
1	CO ₂ Et	70	5	<1
2	CO ₂ Et	70	60	2
3	CO ₂ Et	100	60	5
4	CN	70	5	0
5	CN	100	15	<1
6	CN	150	15	11
7	CN	100	60	6
8	CN	100	180	10

^a Conditions: **9** or **10** (1 mmol), Selectfluor™ (1 equiv), MeCN (15 mL).

^b NMR yield determined using an α,α,α -trifluorotoluene reference.

moval. Both tosyl protected pyrroles **9** and **10** were synthesised and subsequent fluorination conditions were screened (Table 4).

Initial fluorination reactions of **9** and **10** demonstrated the ability of the tosyl substituent to reduce the nucleophilicity of the pyrrole ring compared to substrates attempted previously but, in these cases, negligible conversion to fluorinated products was observed (entries 1 and 4). Unfortunately, upon increasing the reaction time and temperature of the fluorination reactions, the major product observed was an unwanted fluorosulfonyl derivative (**12**) (Fig. 1), which was identified by ¹⁹F NMR and GC–MS of the crude reaction mixture. The formation of this species presumably

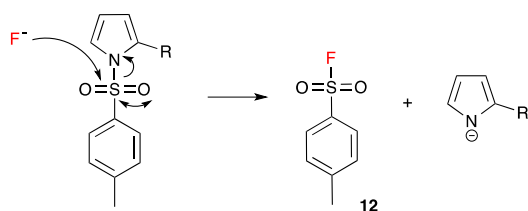


Fig. 1. Possible reaction mechanism resulting in the formation of fluorinated by-product **12**.

occurs via nucleophilic attack of fluoride at the sulfur atom of the tosyl group and we suspect that the fluoride ion originates from the tetrafluoroborate counterion of the Selectfluor™ reagent, due to prolonged exposure to high temperatures under microwave conditions.

An additional substituent present on the pyrrole ring will prevent the formation of any 4,5-difluorinated by-product which can be difficult to separate and could provide useful functionality for subsequent synthetic transformations. Consequently, the synthesis of **13** was carried out, as a bromine substituent allows a range of possible synthetic transformations, particularly palladium catalysed cross coupling chemistry. 4-Bromopyrrole-2-carboxaldehyde (**13**) was obtained in good yield⁴⁸ and optimised fluorination conditions (Table 5) provided the desired product in 29% yield. Furthermore, the structure of **14** was confirmed by X-ray crystallography (Fig. 2) and the regiochemistry of the fluorination reaction was consistent with DFT predicted ¹⁹F NMR data.

The reactivity of 3-substituted pyrrole derivatives bearing an electron withdrawing substituent, such as methyl pyrrole-3-carboxylate (**15**), was also investigated (Table 6). Based on steric and electronic factors, the major product predicted was the 5-fluoro product **16** and the initial conditions applied gave

Table 5
Reaction of **13** with Selectfluor™ under microwave conditions^a



Entry	Time/min	Yield/% ^b
1	5	29
2	10	31
3	15	12
4	7.5	34 (29)

^a Conditions: **13** (1–6 mmol), Selectfluor™ (1 equiv), MeCN (15–25 mL).

^b NMR yield determined using an α,α,α -trifluorotoluene reference, isolated yield in parentheses.

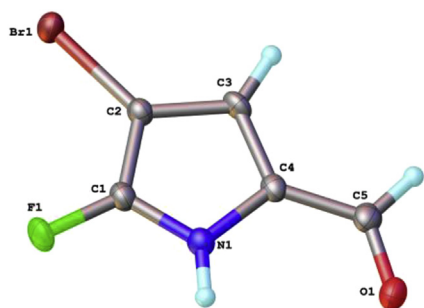
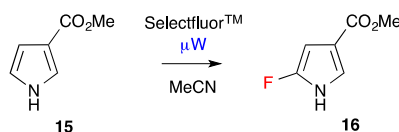


Fig. 2. Molecular structure of **14**.

Table 6
Reaction of **15** with Selectfluor™ under microwave conditions^a



Entry	Temp/°C	Time/min	Yield/% ^c
1	70	5	10
2	70	10	9
3	70	20	8
4	70	2	9
5	100	2	9
6 ^b	70	5	—

^a Conditions: **15** (2 mmol), Selectfluor™ (1 equiv), MeCN (15 mL).

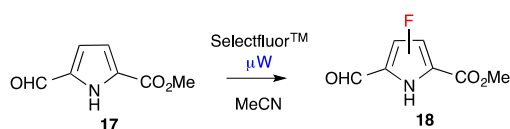
^b Selectfluor (2 equiv).

^c NMR yield determined using an α,α,α -trifluorotoluene reference.

a promising yield of around 10% as measured by ¹⁹F NMR spectroscopy since the other major component in the crude reaction mixture was unreacted starting material. However, upon increasing the reaction time, a larger quantity of insoluble polymer product was observed alongside a slight decrease in yield. Therefore, a shorter reaction time was employed both with the temperature unchanged and at an elevated temperature of 100 °C but, in both cases, comparable yields were obtained. Finally, two equivalents of Selectfluor™ were used in an attempt to increase conversion but a large amount of polymeric product was obtained and a negligible yield of the 5-fluorinated product observed by NMR.

Finally, the reactivity of a 2,5-disubstituted pyrrole derivative was assessed as it was envisaged that two electron withdrawing substituents would help to prevent competing oxidation reactions. Methyl 5-formylpyrrole-2-carboxylate (**17**) was employed and early investigations of reactions conducted at 70 °C (Table 7) did not give rise to any polymeric product. However, even after a period of 2.5 h at this temperature (entry 3), the yield of fluorinated product was disappointingly low and hence the reaction temperature was increased. The results obtained at a reaction temperature of 100 °C were somewhat similar to that seen at lower temperatures and again a low yield of 17% was obtained even after 2.5 h although a large proportion of starting material still remained. Further increases in reaction temperature, however, showed no significant benefit in terms of obtained yield, but appreciable amounts of polymer by-product began to form.

Table 7
Reaction of **17** with Selectfluor™ under microwave conditions^a



Entry	Temp/°C	Time/min	Yield/% ^b
1	70	5	4
2	70	30	11
3	70	150	16
4	100	10	5
5	100	60	16
6	100	150	17
7	125	10	11
8	125	30	17
9	125	60	15
10	150	10	18

^a Conditions: **17** (1 mmol), Selectfluor™ (1 equiv), MeCN (15 mL).

^b NMR yield determined using an α,α,α -trifluorotoluene reference.

3. Conclusions

In conclusion, the effect of various substituents on the fluorination of pyrrole substrates using Selectfluor™ has been assessed but, unfortunately, general methodology for the ready synthesis of fluoropyrroles could not be achieved despite assessment of a range of pyrrole derivatives in electrophilic fluorination reactions. In general, competing oxidation and subsequent polymerisation of the pyrrole substrates by Selectfluor™, which is established as a reasonably strong oxidising agent as well as a fluorinating agent, was a recurring problem for all substrates encountered. However, the synthesis of fluorinated pyrrole derivatives (**2**, **4**, **6**, **8** and **14**), which possess suitable functionality for further synthetic transformations to produce structures relevant for application in the life science industries, was achieved in synthetically useful reactions.

4. Experimental

4.1. General

Chemicals were purchased from Alfa Aesar, Apollo Scientific, Fluorochem or Sigma Aldrich and, unless otherwise stated, were used without any further purification. Dry solvents were obtained using an Innovative Technology Inc. Solvent Purification System. All column chromatography was carried out using Silicagel LC60A (40–63 micron) purchased from Fluorochem. Microwave reactions were performed in a Biotage Initiator microwave synthesiser (0–400 W) in a sealed vessel. Proton, carbon and fluorine nuclear magnetic resonance spectra (^1H NMR, ^{13}C NMR and ^{19}F NMR) were recorded on a Bruker 400 Ultrashield (^1H NMR at 400 MHz; ^{13}C NMR at 101 MHz; ^{19}F NMR at 376 MHz) spectrometer or a Varian VNMRs-700 (^1H NMR at 700 MHz; ^{13}C NMR at 176 MHz) with residual solvent peaks as the internal standard. ^1H , ^{13}C and ^{19}F spectroscopic data are reported as follows: chemical shift (ppm), integration, multiplicity (s=singlet, d=doublet, t=triplet, q=quartet, m=multiplet), coupling constant (Hz). Accurate mass analysis was achieved with a QToF Premier mass spectrometer (Waters Ltd., UK) or an LCT Premier XE mass spectrometer (Waters Ltd., UK) equipped with an accurate solids analysis probe (ASAP). Infra-red (IR) spectra were recorded on a Perkin Elmer FTIR Spectrum Two™ fitted with an ATR probe. Melting points were measured with a Gallenkamp apparatus at atmospheric pressure and are uncorrected.

4.2. Fluorination of pyrroles. General procedures

4.2.1. Reaction of ethyl pyrrole-2-carboxylate (1**) with electrophilic fluorinating agent in the presence of a Lewis acid under microwave conditions (Table 1).** To a solution of **1** (0.21 g, 1.5 mmol) and Selectfluor™ (0.53 g, 1.5 mmol) in dry MeCN (15 mL) was added Lewis acid (50 mol %) before heating with microwave irradiation at 70 °C for 5 min. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3×30 mL) before the solvent was removed in vacuo; analysis by ^{19}F NMR relative to an α,α,α -trifluorotoluene reference gave the yield of 5-fluorinated product.

4.2.2. Reaction of ethyl N-(4-nitrobenzyl)-pyrrole-2-carboxylate (5**) with Selectfluor™ under microwave conditions (Table 2).** A solution of **5** (0.15 g, 0.55 mmol) and Selectfluor™ (0.20 g, 0.55 mmol) in MeCN (15 mL) was heated with microwave irradiation. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3×30 mL) before the solvent

was removed in vacuo; analysis by ^{19}F NMR relative to an α,α,α -trifluorotoluene reference gave the yield of 5-fluorinated product.

4.2.3. Reaction of pyrrole-2-carbonitrile (7**) with Selectfluor™ under microwave conditions (Table 3).** A solution of **7** (0.18 g, 2 mmol) and Selectfluor™ (0.71 g, 2 mmol) in MeCN (15 mL) was heated with microwave irradiation. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3×30 mL) before the solvent was removed in vacuo; analysis by ^{19}F NMR relative to an α,α,α -trifluorotoluene reference gave the yield of 5-fluorinated product.

4.2.4. Reaction of ethyl N-tosyl-pyrrole-2-carboxylate (9**) with Selectfluor™ under microwave conditions (Table 4).** A solution of **9** or **10** (1 mmol) and Selectfluor™ (0.35 g, 1 mmol) in MeCN (15 mL) was heated with microwave irradiation. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3×30 mL) before the solvent was removed in vacuo; analysis by ^{19}F NMR relative to an α,α,α -trifluorotoluene reference gave the yield of 5-fluorinated product. The presence of fluorinated by-product **12** was confirmed by a combination of ^{19}F NMR and GC–MS analysis. ^{19}F NMR (376 MHz, Chloroform- d) δ 66.31 (1 F, s, SF). GC–MS (EI) m/z 174.0.

4.2.5. Reaction of 4-bromopyrrole-2-carboxaldehyde (13**) with Selectfluor™ under microwave conditions (Table 5).** A solution of **13** (0.17 g, 1 mmol) and Selectfluor™ (0.35 g, 1 mmol) in MeCN (15 mL) was heated with microwave irradiation at 70 °C. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3×30 mL) before the solvent was removed in vacuo; analysis by ^{19}F NMR relative to an α,α,α -trifluorotoluene reference gave the yield of 5-fluorinated product.

4.2.6. Reaction of methyl pyrrole-3-carboxylate (15**) with Selectfluor™ under microwave conditions (Table 6).** A solution of **15** (0.25 g, 2 mmol) and Selectfluor™ (0.71 g, 2 mmol) in MeCN (15 mL) was heated with microwave irradiation. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3×30 mL) before the solvent was removed in vacuo; analysis by ^{19}F NMR relative to an α,α,α -trifluorotoluene reference gave the yield of 5-fluorinated product.

4.2.7. Reaction of methyl 5-formylpyrrole-2-carboxylate (17**) with Selectfluor™ under microwave conditions (Table 7).** A solution of **17** (0.15 g, 1 mmol) and Selectfluor™ (0.35 g, 1 mmol) in MeCN (15 mL) was heated with microwave irradiation. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3×30 mL) before the solvent was removed in vacuo; analysis by ^{19}F NMR relative to an α,α,α -trifluorotoluene reference gave the yield of 5-fluorinated product.

4.3. Synthesis of pyrrole substrates

4.3.1. Ethyl N-(4-nitrobenzyl)-pyrrole-2-carboxylate (5**).** To a solution of ethyl pyrrole-2-carboxylate **1** (2.09 g, 15 mmol) in dry DMF (25 mL) at 0 °C, sodium hydride (0.54 g, 23 mmol) was added. The reaction mixture was stirred for 20 min, before the dropwise addition of 4-nitrobenzyl bromide (4.80 g, 23 mmol) in dry DMF (10 mL). After a further 25 min, any excess hydride was decomposed by the addition of ethanol (15 mL) and the reaction mixture poured into distilled water (50 mL). The aqueous solution was

extracted with DCM (4×50 mL) and washed with distilled water (9×100 mL) and brine (100 mL) before being dried (Na₂SO₄) and concentrated in vacuo. The crude product was purified by column chromatography on silica gel using a gradient of hexane/ethyl acetate (0–20% ethyl acetate) as the eluent to give *ethyl N-(4-nitrobenzyl)-pyrrole-2-carboxylate* **5** (2.10 g, 51%) as a yellow solid; Mp 92–94 °C. ¹H NMR (400 MHz, Chloroform-*d*) δ 8.21–8.11 (2 H, m, C3'H), 7.21–7.13 (2 H, m, C2'H), 7.05 (1 H, dd, ³J_{HH} 4.0, ⁴J_{HH} 1.8, C5H), 6.93 (1 H, dd, ³J_{HH} 2.6, ⁴J_{HH} 1.8, C3H), 6.25 (1 H, dd, ³J_{HH} 4.0, ³J_{HH} 2.6, C4H), 5.65 (2 H, s, ArCH₂), 4.19 (2 H, q, ³J_{HH} 7.1, CH₂CH₃), 1.28 (3 H, t, ³J_{HH} 7.1, CH₂CH₃). ¹³C NMR (176 MHz, Chloroform-*d*) δ 161.1 (s, C=O), 147.4 (s, C4'), 146.1 (s, C1'), 129.2 (s, C3), 127.2 (s, C2'), 124.0 (s, C3'), 122.4 (s, C2), 118.9 (s, C5), 109.2 (s, C4), 60.2 (s, CH₂CH₃), 51.8 (s, ArCH₂), 14.5 (s, CH₂CH₃). HRMS (ESI) *m/z* calcd for [M+H]⁺ C₁₄H₁₅N₂O₄ 275.1032; found 275.1043.

4.3.2. N-(4-Nitrobenzyl)-pyrrole-2-carbonitrile. The same procedure as for the synthesis of **5** was employed with pyrrole-2-carbonitrile **7** (0.36 g, 4 mmol), sodium hydride (0.14 g, 6 mmol) and 4-nitrobenzyl bromide (1.28 g, 6 mmol). The crude product obtained was purified by column chromatography on silica gel using a gradient of hexane/ethyl acetate (0–10% ethyl acetate) as the eluent to yield *N-(4-Nitrobenzyl)-pyrrole-2-carbonitrile* (0.22 g, 24%) as a yellow solid; Mp 97–99 °C. IR (neat, cm⁻¹) 3142, 3127, 3077, 2212, 1507, 1340, 1072, 734. ¹H NMR (700 MHz, Chloroform-*d*) δ 8.22–8.19 (2 H, m, C3'H), 7.30–7.27 (2 H, m, C2'H), 6.91 (1 H, dd, ³J_{HH} 2.7, ⁴J_{HH} 1.6, C5H), 6.88 (1 H, dd, ³J_{HH} 4.0, ⁴J_{HH} 1.6, C3H), 6.28 (1 H, dd, ³J_{HH} 4.0, ³J_{HH} 2.7, C4H), 5.33 (2 H, s, ArCH₂). ¹³C NMR (176 MHz, Chloroform-*d*) δ 148.0 (s, C1'), 143.2 (s, C4'), 127.9 (s, C2'), 127.1 (s, C5), 124.4 (s, C3'), 121.1 (s, C3), 113.5 (s, CN), 110.8 (s, C4), 104.5 (s, C2), 51.7 (s, ArCH₂). HRMS (ASAP) *m/z* calcd for [M]⁺ C₁₂H₉N₃O₂ 227.0693; found 227.0695.

4.3.3. Ethyl N-tosyl-pyrrole-2-carboxylate (9). Sodium hydride (0.43 g, 18 mmol) was suspended in anhydrous DMF (10 mL) and cooled to 0 °C before a solution of ethyl pyrrole-2-carboxylate **1** (2.09 g, 15 mmol) in anhydrous DMF (15 mL) was added over a period of 20 min. The reaction mixture was then allowed to return to room temperature and stirred for 1 h before the addition of tosyl chloride (3.43 g, 18 mmol) in anhydrous DMF (10 mL). After 17 h the reaction mixture was poured into distilled water (100 mL) and the aqueous solution extracted with ethyl acetate (3×50 mL). The combined organic extracts were washed with distilled water (4×100 mL) and brine (100 mL) before being dried (Na₂SO₄) and concentrated in vacuo. The crude product was filtered through a silica plug using ethyl acetate as the eluent and the solvent removed in vacuo to yield **9** (3.27 g, 88%) as a pale yellow solid; Mp 43–45 °C. IR (neat, cm⁻¹) 2993, 2980, 1725, 1705, 1681, 1543, 1359, 1258, 1171. ¹H NMR (700 MHz, Chloroform-*d*) δ 7.90–7.84 (2 H, m, C2'H), 7.70 (1 H, dd, ³J_{HH} 3.2, ⁴J_{HH} 1.9, C5H), 7.35–7.28 (2 H, m, C3'H), 7.04 (1 H, dd, ³J_{HH} 3.7, ⁴J_{HH} 1.9, C3H), 6.30 (1 H, t, ³J_{HH} 3.4, C4H), 4.19 (2 H, q, ³J_{HH} 7.1, CH₂CH₃), 2.42 (3 H, s, ArCH₃), 1.26 (3 H, t, ³J_{HH} 7.1, CH₂CH₃). ¹³C NMR (176 MHz, Chloroform-*d*) δ 158.9 (s, C=O), 145.0 (s, C4'), 136.1 (s, C1'), 129.6 (s, C3'), 129.2 (s, C5), 128.3 (s, C2'), 125.4 (s, C2), 123.2 (s, C3), 110.4 (s, C4), 60.9 (s, CH₂CH₃), 21.8 (s, ArCH₃), 14.3 (s, CH₂CH₃). HRMS (ESI) *m/z* calcd for [M–H][–] C₁₄H₁₄NO₄S 292.0644; found 292.0640.

4.3.4. N-Tosyl-pyrrole-2-carbonitrile (10). The same procedure as for the synthesis of **9** was employed with pyrrole-2-carbonitrile **7** (1.38 g, 15 mmol), sodium hydride (0.43 g, 18 mmol) and tosyl chloride (3.43 g, 18 mmol). The crude product was recrystallized from hexane/ethyl acetate to give *N-tosyl-pyrrole-2-carbonitrile* **10** (3.02 g, 82%) as a pale yellow solid; Mp 110–112 °C (lit. 114–1150 °C⁴⁹). IR (neat, cm⁻¹) 2989, 1726, 1705, 1681, 1582, 1354, 1283, 1134. ¹H NMR (400 MHz, Chloroform-*d*) δ 8.01–7.85 (2 H, m,

C2'H), 7.47 (1 H, dd, ³J_{HH} 3.2, ⁴J_{HH} 1.6, C5H), 7.40–7.34 (2 H, m, C3'H), 6.95 (1 H, dd, ³J_{HH} 3.7, ⁴J_{HH} 1.6, C3H), 6.32 (1 H, t, ³J_{HH} 3.5, C4H), 2.44 (3 H, s, ArCH₃). ¹³C NMR (176 MHz, Chloroform-*d*) δ 146.7 (s, C4'), 134.3 (s, C1'), 130.5 (s, C3'), 128.0 (s, C2'), 126.7 (s, C5), 126.7 (s, C3), 112.4 (s, C4), 111.8 (s, CN), 103.9 (s, C2), 21.9 (s, ArCH₃). HRMS (ESI) *m/z* calcd for [M+H]⁺ C₁₂H₁₁N₂O₂S 247.0541; found 247.0547.

4.3.5. 4-Bromopyrrole-2-carboxaldehyde (13). A solution of pyrrole-2-carboxaldehyde **12** (3.80 g, 40 mmol) in dry THF (40 mL) was cooled to 0 °C under Argon. NBS (7.12 g, 40 mmol) was added and the reaction mixture stirred for 15 min before the solvent was removed in vacuo. The crude product was dried under high vacuum for 30 min before the addition of distilled water (20 mL) and the resulting suspension filtered. The resulting solid was dissolved in a minimum amount of hot ethanol/water solution (9:1) before the addition of activated charcoal and filtration through a Celite plug. Upon cooling, the product recrystallised to give *4-bromopyrrole-2-carboxaldehyde* **13** (3.54 g, 51%) as an off white solid; Mp 119–122 °C. IR (neat, cm⁻¹) 3204, 3108, 3982, 2861, 1653, 1378, 1354, 918, 769. ¹H NMR (400 MHz, Acetone-*d*₆) δ 11.39 (1 H, s, NH), 9.52 (1 H, d, ⁴J_{HH} 1.0, CHO), 7.34–7.31 (1 H, m, C5H), 7.09–7.05 (1 H, m, C3H). ¹³C NMR (101 MHz, Acetone-*d*₆) δ 179.4 (s, C=O), 134.3 (s, C2), 126.8 (s, C5), 121.5 (s, C3), 98.4 (s, C4). HRMS (ESI) *m/z* calcd for [M+H]⁺ C₅H₄BrNO 173.9554; found 173.9553.

4.4. Preparative scale fluorination of pyrrole derivatives

4.4.1. Ethyl 5-fluoropyrrole-2-carboxylate (2). A solution of ethyl pyrrole-2-carboxylate **1** (0.56 g, 4 mmol) and SelectfluorTM (1.42 g, 4 mmol) in MeCN (25 mL) was heated by microwave irradiation at 70 °C for 5 min. The reaction was quenched by the addition of H₂O (50 mL) and CHCl₃ (50 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl₃ (3×25 mL). The combined organic phases were dried (Na₂SO₄) and concentrated in vacuo. The crude product was purified by column chromatography on silica gel using hexane/diethyl ether (5:1) as the eluent to give *ethyl 5-fluoropyrrole-2-carboxylate* **2** (0.15 g, 23%) as a yellow solid; *R*_f (3:1, hexane:Et₂O) 0.40. IR (neat, cm⁻¹) 3218, 2984, 1676, 1586, 1237. ¹H NMR (400 MHz, Chloroform-*d*) δ 9.85 (1 H, s, NH), 6.75 (1 H, ddd, ⁴J_{HF} 4.7, ³J_{HH} 4.0, ⁴J_{HH} 2.9, C3H), 5.58 (1 H, ddd, ³J_{HF} 4.0, ³J_{HH} 4.0, ⁴J_{HH} 2.7, C4H), 4.31 (2 H, q, ³J_{HH} 7.1, CH₂CH₃), 1.34 (3 H, t, ³J_{HH} 7.2, CH₂CH₃). ¹⁹F NMR (376 MHz, Chloroform-*d*) δ –130.64 (1 F, ddd, ⁴J_{HF} 4.7, ³J_{HF} 4.0, ³J_{HF} 2.5). ¹³C NMR (176 MHz, Chloroform-*d*) δ 161.0 (s, C=O), 149.4 (d, ¹J_{CF} 267.4, CF), 115.2 (s, C3), 114.2 (s, C2), 89.2 (d, ²J_{CF} 11.4, C4), 60.6 (s, CH₂CH₃), 14.6 (s, CH₂CH₃). HRMS (ESI) *m/z* calcd for [M+H]⁺ C₇H₉FNO₂ 158.0617; found 158.0598.

4.4.2. Methyl 1-methyl-5-fluoropyrrole-2-carboxylate (4). A solution of methyl 1-methyl-pyrrole-2-carboxylate **3** (0.56 g, 4 mmol) and SelectfluorTM (1.42 g, 4 mmol) in MeCN (25 mL) was heated by microwave irradiation at 70 °C for 5 min. The reaction was quenched by the addition of H₂O (50 mL) and CHCl₃ (50 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl₃ (3×25 mL). The combined organic phases were dried (Na₂SO₄) and concentrated in vacuo. The crude product was purified by column chromatography on silica gel using hexane/diethyl ether (5:1) as the eluent to yield *methyl 1-methyl-5-fluoropyrrole-2-carboxylate* **4** (0.10 g, 16%) as a colourless oil; *R*_f (3:1, hexane:Et₂O) 0.42. IR (neat, cm⁻¹) 2955, 1704, 1565, 1472, 1252, 1103. ¹H NMR (400 MHz, chloroform-*d*) δ 6.81 (1 H, dd, ⁴J_{HF} 6.3, ³J_{HH} 4.3, C3H), 5.56 (1 H, dd, ³J_{HH} 4.3, ³J_{HF} 4.3, C4H), 3.79 (3 H, s, CO₂CH₃), 3.76 (3 H, d, ⁴J_{HF} 1.2, NCH₃). ¹⁹F NMR (376 MHz, Chloroform-*d*) δ –131.89 (1 F, ddq, ⁴J_{HF} 6.3, ³J_{HF} 4.2, ⁴J_{HF} 1.2). ¹³C NMR (101 MHz, Chloroform-*d*) δ 161.4 (d, ⁴J_{CF} 2.2, C=O), 150.5 (d, ¹J_{CF} 267.1, CF),

116.0 (d, $^3J_{\text{CF}}$ 4.3, **C3**), 114.5 (s, **C2**), 87.6 (d, $^2J_{\text{CF}}$ 12.2, **C4**), 51.1 (s, OCH_3), 30.6 (d, $^3J_{\text{CF}}$ 2.8, NCH_3). HRMS (ESI) m/z calcd for $[\text{M}+\text{H}]^+$ $\text{C}_7\text{H}_9\text{FNO}_2$ 158.0617; found 158.0615.

4.4.3. Ethyl N-(4-nitrobenzyl)-5-fluoropyrrole-2-carboxylate (6). A solution of ethyl N-(4-nitrobenzyl)-pyrrole-2-carboxylate **5** (0.15 g, 0.55 mmol) and Selectfluor™ (0.20 g, 0.55 mmol) in MeCN (15 mL) was heated with microwave irradiation at 70 °C for 5 min. The reaction was quenched by the addition of H_2O (30 mL) and CHCl_3 (30 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3 × 30 mL). The combined organic extracts were dried (Na_2SO_4) and concentrated in vacuo. The crude product was purified by column chromatography on silica gel using a gradient of hexane/diethyl ether (0–10% diethyl ether) as the eluent to give ethyl N-(4-nitrobenzyl)-5-fluoropyrrole-2-carboxylate **6** (25 mg, 15%) as a pale yellow solid; Mp 52–54 °C. R_f (3:1, hexane:Et₂O) 0.26. IR (neat, cm^{-1}) 3217, 2932, 1751, 1667, 1523, 1344, 1256. ^1H NMR (400 MHz, Chloroform-*d*) δ 8.21–8.12 (2 H, m, **C3'H**), 7.31–7.27 (2 H, m, **C2'H**), 6.93 (1 H, dd, $^4J_{\text{HF}}$ 6.2, $^3J_{\text{HH}}$ 4.2, **C3H**), 5.69 (1 H, dd, $^3J_{\text{HH}}$ 4.2, $^3J_{\text{HF}}$ 4.2, **C4H**), 5.59 (2 H, s, ArCH_2), 4.21 (2 H, q, $^3J_{\text{HH}}$ 7.1, CH_2CH_3), 1.29 (3 H, t, $^3J_{\text{HH}}$ 7.1, CH_2CH_3). ^{19}F NMR (376 MHz, Chloroform-*d*) δ –131.65 (1 F, dd, $^4J_{\text{HF}}$ 6.2, $^4J_{\text{HF}}$ 4.2). ^{13}C NMR (176 MHz, Chloroform-*d*) δ 160.8 (d, $^4J_{\text{CF}}$ 2.1, $\text{C}=\text{O}$), 150.3 (d, $^1J_{\text{CF}}$ 268.7, CF), 147.5 (s, **C4'**), 144.7 (s, **C1'**), 127.7 (s, **C2'**), 124.1 (s, **C3'**), 117.0 (d, $^3J_{\text{CF}}$ 4.0, **C3**), 114.3 (s, **C2**), 88.3 (d, $^2J_{\text{CF}}$ 11.6, **C4**), 60.2 (s, CH_2CH_3), 46.2 (d, $^3J_{\text{CF}}$ 2.0, ArCH_2), 14.5 (s, CH_2CH_3). HRMS (ESI) m/z calcd for $[\text{M}+\text{H}]^+$ $\text{C}_{14}\text{H}_{14}\text{FN}_2\text{O}_4$ 293.0938; found 293.0944.

4.4.4. 5-Fluoropyrrole-2-carbonitrile (8). A solution of pyrrole-2-carbonitrile **7** (0.90 g, 10 mmol) and Selectfluor™ (3.55 g, 10 mmol) in MeCN (25 mL) was heated with microwave irradiation at 70 °C for 5 min. The reaction was quenched by the addition of H_2O (100 mL) and CHCl_3 (50 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3 × 30 mL). The combined organic extracts were dried (MgSO_4) and concentrated in vacuo. The crude product was purified by column chromatography on silica gel using a gradient of hexane/diethyl ether (0–10% diethyl ether) as the eluent to give 5-fluoropyrrole-2-carbonitrile **8** (0.33 g, 30%) as a white solid; Mp 55–57 °C. R_f (3:1, hexane:Et₂O) 0.25. IR (neat, cm^{-1}) 3196, 3152, 2227, 1588, 1007. ^1H NMR (400 MHz, Chloroform-*d*) δ 8.67 (1 H, s, NH), 6.72 (1 H, ddd, $^4J_{\text{HF}}$ 4.5, $^3J_{\text{HH}}$ 4.3, $^4J_{\text{HH}}$ 3.0, **C3H**), 5.64 (1 H, ddd, $^3J_{\text{HH}}$ 4.1, $^3J_{\text{HF}}$ 3.6, $^4J_{\text{HH}}$ 2.7, **C4H**). ^{19}F NMR (376 MHz, Chloroform-*d*) δ –128.90 (1 F, ddd, $^4J_{\text{HF}}$ 4.5, $^3J_{\text{HF}}$ 3.6, $^3J_{\text{HF}}$ 2.5). ^{13}C NMR (176 MHz, Chloroform-*d*) δ 148.7 (d, $^1J_{\text{CF}}$ 268.1, CF), 120.7 (d, $^3J_{\text{CF}}$ 2.7, **C3**), 114.1 (s, CN), 92.8 (d, $^3J_{\text{CF}}$ 5.2, **C2**), 89.3 (d, $^2J_{\text{CF}}$ 11.0, **C4**). HRMS (ESI) m/z calcd for $[\text{M}-\text{H}]^-$ $\text{C}_5\text{H}_2\text{FN}_2$ 109.0202; found 109.0196.

4.4.5. 4-Bromo-5-fluoropyrrole-2-carboxaldehyde (14). A solution of 4-bromopyrrole-2-carboxaldehyde **13** (1.04 g, 6 mmol) and Selectfluor™ (2.13 g, 6 mmol) in MeCN (25 mL) was heated with microwave irradiation at 70 °C for 7.5 min. The reaction was quenched by the addition of H_2O (50 mL) and CHCl_3 (50 mL) was subsequently added. The layers were separated and the aqueous phase extracted with CHCl_3 (3 × 30 mL). The combined organic extracts were dried (Na_2SO_4) and concentrated in vacuo. The crude product was purified by column chromatography on silica gel using a gradient of hexane/diethyl ether (0–20% diethyl ether) as the eluent to give 4-bromo-5-fluoropyrrole-2-carboxaldehyde **14** (0.33 g, 29%) as off white crystals. Mp 148–150 °C (with degradation). R_f (3:1, hexane:Et₂O) 0.20. IR (neat, cm^{-1}) 3115, 2950, 2783, 2676, 2553, 1627, 1576, 1409, 1143, 668. ^1H NMR (400 MHz, Acetone-*d*₆) δ 9.40 (1 H, d, $^4J_{\text{HH}}$ 3.5, CHO), 7.05 (1 H, d, $^4J_{\text{HF}}$ 4.5, **C3H**). ^{19}F NMR (376 MHz, Acetone-*d*₆) δ –132.72 (1 F, m). ^{13}C NMR (151 MHz, Acetone-*d*₆) δ 178.2 (d, $^4J_{\text{CF}}$ 2.7, $\text{C}=\text{O}$), 149.3 (d, $^1J_{\text{CF}}$ 268.0, CF), 124.3 (s, **C2**), 120.9 (s, **C3**), 76.1 (d, $^2J_{\text{CF}}$ 15.1, **C4**). HRMS (ESI) m/z calcd for

$[\text{M}+\text{H}]^+$ $\text{C}_5\text{H}_4\text{BrFNO}$ 191.9460; found 191.9463. Crystals suitable for X-ray diffraction were grown by slow evaporation from acetone.

Crystal data for **14**: $\text{C}_5\text{H}_3\text{BrFNO}$, $M=191.99$, orthorhombic, space group $P\ bca$, $a=7.1484(3)$, $b=8.5392(4)$, $c=19.5396(9)$ Å, $U=1192.73(9)$ Å³, $F(000)=736.0$, $Z=8$, $D_c=2.138$ mg m^{−3}, $\mu=6.817$ mm^{−1} (Mo-K α , $\lambda=0.71073$ Å), $T=120(1)$ K. 15,683 reflections were collected on a Bruker D8 Venture diffractometer (Photon100 CMOS detector, μS -microsource, focussing mirrors) yielding 1582 unique data ($R_{\text{merge}}=0.0669$). The structure was solved by direct method and refined by full-matrix least squares on F^2 for all data using SHELXTL and OLEX2 software. All non-hydrogen atoms were refined with anisotropic displacement parameters, H-atoms were placed in calcd positions and refined in riding mode. Final $wR_2(F^2)=0.0934$ for all data (82 refined parameters), conventional $R_1(F)=0.0399$ for 1201 reflections with $I \geq 2\sigma$, GOF=1.111. Crystallographic data for the structure have been deposited with the Cambridge Crystallographic Data Centre as supplementary publication CCDC-14449050.

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