Electronic Supplementary Information

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1. Experimental section

a) General information

All chemicals, unless otherwise noted, were purchased from commercial sources and were used without further purification. Unless stated otherwise, all reactions were carried out under nitrogen atmosphere. The substrates were synthesized according to the literature methods.¹⁻⁴ Irradiation with visible light was performed using blue LEDs illumination instruments (The actual output power density of the LEDs at 0.5 cm distance is 33.70 mW/cm² detected by CEL-NP2000-10 (Beijing CeauLight Co. Ltd., China) light power meter).

The nuclear magnetic resonance spectra were recorded on the Bruker Ascend™ 400 MHz NMR spectrometer and the Bruker Ascend[™] 500 MHz NMR spectrometer with tetramethylsilane (TMS) as an internal standard. High resolution mass spectra were recorded using a Q Exactive mass spectrometer (Thermo Fisher Scientific, USA). X-Ray diffraction data of the single crystal were collected on XtaLAB PRO MM007HF DW.

b) Methods for the synthesis of substrates



i) To a solution of 2-((trimethylsilyl)ethynyl)aniline (3 mmol, 1.0 equiv) in AcOH (7.0 mL) was added benzaldehyde (3.6 mmol, 1.2 equiv). The reaction was stirred at room temperature for 1 h and then cooled to 0 °C. The resulting mixture was then warmed to ambient temperature and stirred for 1 h before the addition of NaBH4 (6 mmol, 2.0 equiv). NaOH (3 N, 30 mL) and ethyl acetate (30 mL) were added and the layers were separated. The aqueous phase was extracted twice with ethyl acetate. The combined organic phase was dried over Na₂SO₄, filtered and concentrated under reduced pressure to afford **S1**. The crude product **S1** were isolated by silica-gel column chromatography in excellent yield (64-82%).

ii) **S1** was treated with K₂CO₃ (0.83 g, 6 mmol) in MeOH (20 mL) at room temperature for 1 h. MeOH was removed under reduced pressure. The residue was diluted with ethyl acetate (30 mL) and water (20 mL). The phases were separated and the aqueous phase was extracted twice with ethyl acetate. The combined organic solution was dried over MgSO₄, filtered, and concentrated under reduced pressure to afford **S2** as a light yellow oil. The crude product **S2** was used in the following step without purification.

iii) To a solution of **S2** in CH₂Cl₂ (15 mL) was added Et₃N (0.84 mL, 6.0 mmol) and trifluoroacetic anhydride (0.51 mL, 3.6 mmol). The reaction mixture was stirred at room temperature for 2 h and then concentrated under reduced pressure. The residue was chromatographed through silica gel eluting with ethyl acetate/hexanes to give the product (47-85%).



Some of the substrates were synthesized using this method. To a solution of *N*-(2ethynylphenyl)-2,2,2-trifluoroacetamide (1.0 equiv) in CH₃CN (0.2 M) was added K_2CO_3 (1.2 equiv), followed by benzyl bromide (1.2 equiv). The reaction mixture was stirred at 80 °C for 2 h, cooled to RT, and concentrated under reduced pressure. The residue was chromatographed through silica gel eluting with ethyl acetate/hexanes to give the product (75-83%).



N-acetyl-*o*-ethynylaniline was synthesized using this method. To a solution of *o*ethynylaniline (1 mmol, 117 mg), Et₃N (1.5 mmol, 0.206 mL) in THF (2 mL) was added acetyl chloride (1.5 mmol, 118mg) dropwise at room temperature. After the addition was complete, the reaction mixture was stirred for a further 30 minutes. The reaction was quenched by water (10 mL) and the resulting mixture was extracted with Et₂O (3×10 mL). The combined extracts were washed with brine, dried over sodium sulfate and concentrated in vacuo. The residue was purified by flash column chromatography on silica gel to afford *N*-acetyl-*o*-ethynylaniline **S4** as a white solid (158.4 mg, 99% yield).



To a solid of **a23** (696 mg, 2 mmol, 1 equiv) in ethanol (8 mL) were added iron powder (448 mg, 8 mmol, 4 equiv) and concd HCl (2 mL, 2.4mmol, 1.2 equiv). After 2 h of reflux, the mixture was cooled to room temperature and Na₂CO₃ was added by

portions until gas evolution ceased. After filtration over Celite, the filtrate was extracted with Et_2O and the combined organic fractions were washed with H_2O and a saturated solution of NaCl. The organic layer was then dried over Na_2SO_4 , filtered, and concentrated in vacuo. The residue was purified by flash column chromatography on silica gel to afford **a** (217 mg, 35 % yield).⁵

	c)	Table S	1.0	ptimizati	on of the	reaction	time ^a
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	+ (* * *	F ₃ C CF ₃	$\frac{3\% \operatorname{Ru}(\operatorname{bpy})_3 \operatorname{Cl}_2^{\bullet} \operatorname{6H}_2 \operatorname{O}}{\operatorname{CH}_2 \operatorname{Cl}_2, \operatorname{hv}}$	
a1	b	c1		0 d1
1 equiv.	2.5 equiv.	2.5 equiv.		
Entry	Reaction	time (h)	Yie	$\operatorname{Id}(\%)^b$
1	3			50
2	4			59
3	5			73
4	6			85
5	7			86
6	8			86

^{*a*} Reaction conditions: 0.1 mmol **a1**, 3% Ru(bpy)₃Cl₂·6H₂O, 0.25 mmol **b** and **c1** were dissolved in CH₂Cl₂, degassed for 15 min under N₂, 450 nm LEDs irradiation at room temperature; ^{*b*} isolated yields.

d) Scheme S1. Optimization of pyridine-*N*-oxide derivatives.



e) Crystal structure determination of d1

A suitable crystal of **d1** was mounted with glue at the end of a glass fiber. Data collection was performed with a XtaLAB PRO MM007-DW diffractometer system equipped with a RA-Micro7HF-MR-DW(Cu/Mo) X-ray generator and Pilatus3R-200K-A detector (Rigaku, Japan, Cu K α , $\lambda = 1.54178$ Å). The structure was solved by direct methods (SHELXTL-97) and refined by full-matrix least-squares (SHELXTL-97)⁶ refinements based on F^2 . Anisotropic thermal parameters were applied to all non-hydrogen atoms. The hydrogen atoms were generated geometrically. Crystal data and structure refinement parameters are summarized in Table S2. CCDC No. 1936747.

Single crystals suitable for X-ray diffraction were obtained by slow evaporation of a saturated solution of **d1** (methanol) in a loosely capped vial.

Identification code	d1
Empirical formula	C ₁₈ H ₁₁ F ₆ NO
Formula weight	371.28
Temperature/K	100
Crystal system	monoclinic
Space group	P2 ₁ /c
a/Å	7.8518(2)
b/Å	15.2537(5)
c/Å	26.0789(7)
<u>α/°</u>	90
β/°	92.227(3)
γ/°	90
Volume/Å ³	3121.07(16)
Z	8
$\rho_{calc}g/cm^3$	1.580

Table S2 Crystal data and structure refinement for d1

µ/mm ⁻¹	1.305
F(000)	1504.0
Crystal size/mm ³	0.3 imes 0.1 imes 0.05
Radiation	$CuK\alpha (\lambda = 1.54184)$
2Θ range for data collection/°	6.714 to 148.6
Index ranges	$-7 \le h \le 9, -18 \le k \le 18, -31 \le l \le 32$
Reflections collected	18741
Independent reflections	$6153 [R_{int} = 0.0419, R_{sigma} = 0.0286]$
Data/restraints/parameters	6153/3/558
Goodness-of-fit on F ²	1.055
Final R indexes [I>= 2σ (I)]	$R_1 = 0.0521, wR_2 = 0.1340$
Final R indexes [all data]	$R_1 = 0.0587, wR_2 = 0.1378$
Largest diff. peak/hole / e Å-3	0.40/-0.22

f) General procedure for the photoreactions



The substrate (0.1 mmol, 1 equiv.), trifluoroacetic anhydride (0.25 mmol, 2.5 equiv.), pyridine *N*-oxide (0.25 mmol, 2.5 equiv.) and 3 mol% $Ru(bpy)_3Cl_2 \cdot 6H_2O$ were dissolved in 6.0 mL CH₂Cl₂ in a 15 mL reaction tube equipped with magnetic stirring bar, the reaction tube was sealed and the resulting mixture was deaerated with nitrogen for 15 min, then the reaction tube was irradiated by blue LEDs for 9 h. After reaction, the solvent was removed by rotary evaporation and purified by column chromatography on silica gel using petroleum ether/ethyl acetate as the eluent.



Figure S1. Reaction setup for general photoreactions.



The substrate (0.1 mmol, 1 equiv.), ethyldifluoroiodoacetate (0.2 mmol, 2 equiv.), Potassium carbonate (0.15 mmol, 1.5 equiv.) and the **Pt-I** complex (0.003 mmol, 3 mol%) were dissolved in 5.0 mL (CH₂Cl₂:CH₃OH=1:1) in a 15 mL reaction tube equipped with magnetic stirring bar, the reaction tube was sealed and the resulting mixture was deaerated with nitrogen for 15 min, then the reaction tube was irradiated by blue LEDs for 12 h. After reaction, the solvent was removed by rotary evaporation and purified by column chromatography on silica gel using petroleum ether/ethyl acetate as the eluent.



The substrate (0.1 mmol, 1 equiv.), diethyl bromodifluoromethanephosphonate (0.2 mmol, 2 equiv.), sodium dicarbonate (0.15 mmol, 1.5 equiv.) and the **Pt-I** complex (0.003 mmol, 3 mol%) were dissolved in 5.0 mL CH_2Cl_2 in a 15 mL reaction tube equipped with magnetic stirring bar, the reaction tube was sealed and the resulting mixture was deaerated with nitrogen for 15 min, then the reaction tube was irradiated by blue LEDs for 12 h. After reaction, the solvent was removed by rotary evaporation and purified by column chromatography on silica gel using petroleum ether/ethyl acetate as the eluent.

g) Procedure for the gram-scale reaction

The substrate (3.4 mmol, 1 equiv.), trifluoroacetic anhydride (8.5 mmol, 2.5 equiv.), pyridine *N*-oxide (8.5 mmol, 2.5 equiv.) and 3 mol% $Ru(bpy)_3Cl_2 \cdot 6H_2O$ were dissolved in 80 mL CH₂Cl₂ in a 250 mL reaction flask equipped with magnetic stirring bar, the flask was sealed and the resulting mixture was deaerated with nitrogen for 30 min, then the flask was irradiated by blue LEDs for 48 h. After reaction, the solvent was removed by rotary evaporation and purified by column chromatography on silica gel using petroleum ether/ethyl acetate as the eluent.



Figure S2. Gram scale reaction setup and product d1.

h) Scheme S2. Trifluoroacetic anhydride induced oxidation of triple bond.



i) Procedures for further transformation of d1



A white solid of **d1** (1.1 g, 2.8 mmol) in MeOH (150 mL) was cooled to 0 °C, and then 5 N NaOH solution (25 mL) was added slowly. This mixture was stirred for 1 h at room temperature. The residue mixture was concentrated under reduced pressure. This solution was then extracted with CH_2Cl_2 (2 × 20 mL). The combined organic phases were washed with brine and dried over sodium sulphate. The solvent was removed by rotary evaporation and purified by column chromatography on silica gel by using petroleum ether/ethyl acetate as eluent.⁷



The substrate **f** (0.1 mmol, 1 equiv) and **Pt 1** (0.00025 mmol, 0.0025 equiv) were dissolved in DMF (2 mL) in a 10 mL reaction tube equipped with magnetic stirring bar, and the resulting mixture was irradiated using blue LEDs at the ambient condition. After the substrate was completely converted (monitored by TLC), the reaction mixture was evaporated under reduced pressure until DMF was gone. Then diethyl ether (25 mL) and water (25 mL) were added to the residue. The organic layer was extracted with diethyl ether (3 × 25 mL). The combined organic phases were washed with brine and dried over sodium sulphate. The solvent was removed by rotary evaporation and purified by column chromatography on silica gel by using petroleum ether/ethyl acetate as eluent.⁸



To a solution of **f** (0.21 mmol, 1.0 equiv) in CHCl₃ (5 mL) was added MnO₂ (1.0 mmol, 5.0 equiv). The reaction mixture was heated for 1 h at 60 °C and cooled to room temperature. The solvent was removed by rotary evaporation and purified by column chromatography on silica gel by using petroleum ether/ethyl acetate as eluent.

j) Reaction Quantum Yield (Φ) Measurement⁹⁻¹¹

1) Determination of the light intensity at 450 nm

The photon flux of the LED ($\lambda_{max} = 450$ nm) was determined by standard ferrioxalate actinometry. By dissolving potassium ferrioxalate hydrate (0.737 g) in H₂SO₄ (10 mL of a 0.05 M solution), a 0.15 M solution of ferrioxalate was prepared. By dissolving 1,10-phenanthroline (25 mg) and sodium acetate (5.63 g) in H₂SO₄ (25 mL of a 0.5 M solution), a buffered solution of 1,10-phenanthroline was prepared. Both solutions were stored in the dark. To determine the photon flux of the LED, a cuvette containing 1.0 mL ferrioxalate solution was irradiated for 90 s at $\lambda_{max} = 450$ nm. After irradiation, 0.175 mL phenanthroline solution was added to the cuvette and the mixture was allowed to stir in the dark for 1.5 h to allow the ferrous ions to completely coordinate to the phenanthroline. The absorbance of the solution was measured at 510 nm. A non-irradiated sample was also prepared and measured the absorbance at 510 nm. The results were shown as below:



Figure S3. UV-vis spectrum of irradiation and non-irradiation sample. Conversion was calculated using equation 1.

$$\operatorname{mol} \operatorname{Fe}^{2+} = \frac{V \cdot \Delta A(510 \ nm)}{1 \cdot \varepsilon}$$
(1)

Where V is the total volume (0.001175 L) of the solution after addition of phenanthroline, ΔA is the difference in absorbance at 510 nm between the irradiated and non-irradiated solutions, 1 is the path length (1.0 cm), and ε is the molar absorptivity of the ferrioxalate actinometer at 510 nm (11100 Lmol⁻¹cm⁻¹). The photon flux can be calculated using equation 2.

$$=\frac{mol Fe2 +}{\Phi \bullet t \bullet f}$$
(2)

Where Φ is the quantum yield for the ferrioxalate actinometer at $\lambda_{max} = 450$ nm, t is the irradiation time, and f is the fraction of light absorbed at $\lambda_{max} = 450$ nm by the ferrioxalate actinometer. This value is calculated using equation 3 where A is the absorbance of the ferrioxalte solution at 450 nm. An absorption spectrum gave an A value of >3, indicating that the fraction of absorbed light (f) is > 0.999.

$$f = 1 - 10^{-A(450 \text{ nm})}$$

The photon flux was thus calculated to be 1.05×10^{-9} einstein s⁻¹.

2) Determination of the reaction quantum yield

The degassed solution containing **a1** (0.1 mmol, 1 equiv.), trifluoroacetic anhydride (0.25 mmol, 2.5 equiv.), pyridine *N*-oxide (0.25 mmol, 2.5 equiv.) and 3 mol% $Ru(bpy)_3Cl_2 \cdot 6H_2O$ with magnetic stirring bar was constantly irradiated for 30 min. After irradiation, the yield of product **d1** was determined to be 9 mol% by ¹H NMR. The reaction quantum yield was determined using equation 4, where photon flux was determined as above described, t is the reaction time, f is the fraction of incident light absorbed by the reaction mixture. This value is calculated using equation 3 where A is the absorbance of the reaction mixture at 450 nm. An absorption spectrum of the reaction mixture shown as Figure S4 gave an A value of >3, indicating that the fraction of absorbed light (f) is > 0.999.

$$\Phi = \frac{mol \ of \ product \ formed}{photon \ flux \bullet t \bullet f}$$
(4)



Figure S4. UV-vis spectrum of reaction mixture.

The reaction quantum yield (Φ) was determined to be 5.5, which is above unity, indicating that a radical chain propagation might be operative in this reaction.

k) References

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2. Characterization data of the products



¹H NMR (400 MHz, CDCl₃) δ ppm = 7.49 – 7.41 (m, 4H), 7.39 – 7.34 (m, 2H), 7.30 (t, *J* = 7.1 Hz, 1H),
7.16 (d, *J* = 7.5 Hz, 1H), 6.12 (q, *J* = 8.0 Hz, 1H), 5.98 (d, *J* = 17.1 Hz, 1H), 4.34 (d, *J* = 17.1 Hz, 1H).
¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.83 (q, *J* = 36.7 Hz), 148.56 (q, *J* = 5.0 Hz), 136.28 (s), 135.85 (s), 134.75 (s), 133.69 (s), 130.07 (s), 129.86 (s), 129.39 (s), 129.31 (s), 128.82 (q, *J* = 2.9 Hz), 128.05 (s), 127.93 (s), 126.69 (d, *J* = 2.5 Hz), 122.14 (q, *J* = 270.9 Hz), 120.18 (q, *J* = 34.4 Hz), 116.11 (q, *J* = 288.5 Hz), 50.62 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.18 (s, 3F), -67.90 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₁F₆NONa: 394.0643, found: 394.0631.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.40 (dd, *J* = 7.6, 1.3 Hz, 1H), 7.35 (td, *J* = 7.4, 1.5 Hz, 1H), 7.31 - 7.28 (m, 1H), 7.24 (s, 2H), 7.20 (s, 1H), 7.14 (d, *J* = 7.5 Hz, 1H), 6.09 (q, *J* = 8.1 Hz, 1H), 5.96 (d, *J* = 17.1 Hz, 1H), 4.31 (d, *J* = 17.1 Hz, 1H), 2.39 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.97 (q, *J* = 36.5 Hz), 148.78 (q, *J* = 5.6 Hz), 139.67 (s), 136.09 (s), 134.95 (s), 133.76 (s), 133.19 (s), 130.60 (s), 129.78 (s), 129.26 (s), 129.20 (d, *J* = 2.5 Hz), 128.06 (s), 127.86 (s), 126.43 (d, *J* = 1.7 Hz), 122.20 (q, *J* = 272.2 Hz), 119.91 (q, *J* = 34.0 Hz), 116.16 (q, *J* = 288.5 Hz), 50.69 (s), 21.17 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.21 (s, 3F), -67.90 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₃F₆NONa: 408.0799, found: 408.0786.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.41 (dd, *J* = 7.6, 1.2 Hz, 1H), 7.35 (td, *J* = 7.4, 1.5 Hz, 1H), 7.30 (d, *J* = 7.7 Hz, 2H), 7.23 (d, *J* = 7.9 Hz, 1H), 7.18 (s, 1H), 7.15 (d, *J* = 7.5 Hz, 1H), 6.11 (q, *J* = 8.1 Hz, 1H), 5.96 (d, *J* = 17.1 Hz, 1H), 4.33 (d, *J* = 17.0 Hz, 1H), 2.41 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.82 (q, *J* = 36.6 Hz), 148.66 (q, *J* = 5.6 Hz), 140.55 (s), 135.73 (s), 134.98 (s), 133.73 (s), 133.22 (s), 130.06 (s), 129.74 (s), 129.26 (s), 128.55 (q, *J* = 2.8 Hz), 128.05 (s), 127.86 (s), 127.08 (d, *J* = 1.7 Hz), 122.24 (q, *J* = 272.2 Hz), 120.00 (q, *J* = 34.0 Hz), 116.15 (d, *J* = 288.5 Hz), 50.67 (s), 21.16 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -57.10 (s, 3F), -67.86 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₃F₆NONa: 408.0799, found: 408.0789.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.38 (dd, J = 13.2, 7.3 Hz, 3H), 7.31 (t, J = 7.4 Hz, 1H), 7.17 – 7.12 (m, 3H), 6.13 (q, J = 8.0 Hz, 1H), 5.97 (d, J = 17.1 Hz, 1H), 4.32 (d, J = 17.1 Hz, 1H). ¹³**C NMR** (126 MHz, CDCl₃) δ ppm = 163.25 (s), 161.26 (s), 156.85 (q, J = 36.7 Hz), 147.37 (q, J = 5.0 Hz), 138.42 (d, J = 9.0 Hz), 134.14 (s), 133.57 (s), 131.84 (d, J = 2.6 Hz), 130.12 (s), 129.44 (s), 128.72 (d, J = 9.3 Hz), 128.08 (d, J = 3.4 Hz), 121.93 (q, J = 271.5 Hz), 120.70 (q, J = 34.6 Hz), 116.90 (q, J = 23.0 Hz), 116.06 (ddd, J = 24.5, 6.0, 3.0Hz), 116.04 (q, J = 289.2 Hz), 50.63 (s) ¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.30 (s, 3F), -67.92 (s, 3F), -110.42 (s, 1F). **HRMS** (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₀F₇NONa: 412.0548, found: 412.0533.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.44 – 7.39 (m, 3H), 7.37 (dd, J = 7.5, 1.2 Hz, 1H), 7.31 (t, J = 8.0

Hz, 2H), 7.16 (d, *J* = 7.6 Hz, 1H), 6.14 (q, *J* = 8.0 Hz, 1H), 5.97 (d, *J* = 17.1 Hz, 1H), 4.32 (d, *J* = 17.1 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.70 (q, *J* = 36.8 Hz), 147.22 (q, *J* = 5.6 Hz), 137.87 (s), 135.34 (s), 134.34 (s), 134.13 (s), 133.46 (s), 130.14 (s), 129.39 (s), 128.82 (q, *J* = 2.9 Hz), 128.15 (d, *J* = 2.5 Hz), 128.10 (d, *J* = 3.8 Hz), 121.95 (q, *J* = 270.9 Hz), 120.72 (q, *J* = 34.5 Hz), 116.04 (q, *J* = 289.8 Hz), 50.52 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.29 (s, 3F), -67.89 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₀ClF₆NONa: 428.0253, found: 428.0229.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.61 – 7.56 (m, 2H), 7.38 (ddd, *J* = 8.9, 5.6, 1.8 Hz, 2H), 7.33 – 7.29 (m, 1H), 7.24 (s, 1H), 7.16 (d, *J* = 7.4 Hz, 1H), 6.13 (q, *J* = 8.0 Hz, 1H), 5.97 (d, *J* = 17.2 Hz, 1H), 4.31 (d, *J* = 17.1 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.64 (q, *J* = 36.8 Hz), 147.07 (q, *J* = 5.6 Hz), 138.10 (s), 134.85 (s), 134.13 (s), 133.43 (s), 133.16 (s), 131.71 (q, *J* = 2.9 Hz), 130.14 (s), 129.37 (s), 128.36 (d, *J* = 1.8 Hz), 128.10 (d, *J* = 4.7 Hz), 123.23 (s), 121.93 (q, *J* = 272.2 Hz), 120.74 (q, *J* = 35.3 Hz), 116.00 (q, *J* = 288.5 Hz), 50.47 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.31 (s, 3F), -67.90 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₀BrF₆NONa: 471.9748, found: 471.9740.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.75 (d, *J* = 8.2 Hz, 1H), 7.70 (s, 1H), 7.53 (d, *J* = 8.2 Hz, 1H), 7.45 - 7.38 (m, 2H), 7.36 - 7.32 (m, 1H), 7.18 (d, *J* = 7.5 Hz, 1H), 6.19 (q, *J* = 7.9 Hz, 1H), 6.01 (d, *J* = 16.9 Hz, 1H), 4.34 (d, *J* = 16.6 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.50 (q, *J* = 37.3 Hz), 147.11 (q, *J* = 5.5 Hz), 139.00 (s), 137.14 (s), 133.83 (s), 133.29 (s), 131.76 (q, *J* = 32.8 Hz), 130.28 (s), 129.39 (s), 128.26 (s), 128.10 (s), 127.53

(s), 127.16 (dd, J = 7.0, 3.5 Hz), 126.20 (t, J = 3.8 Hz), 123.20 (q, J = 273.4 Hz), 121.87 (q, J = 272.2

Hz), 121.16 (q, J = 35.3 Hz), 115.95 (q, J = 288.5 Hz), 50.40 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.36 (s, 3F), -62.90 (s, 3F), -67.94 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₀F₉NONa: 462.0516, found: 462.0508.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.42 – 7.35 (m, 3H), 7.31 (t, *J* = 7.2 Hz, 1H), 7.18 – 7.12 (m, 3H), 6.15 (q, *J* = 8.0 Hz, 1H), 5.96 (d, *J* = 17.0 Hz, 1H), 4.34 (d, *J* = 17.0 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 163.80 (s), 161.80 (s), 156.57 (q, *J* = 37.0 Hz), 147.62 (q, *J* = 5.7 Hz), 137.06 (d, *J* = 9.8 Hz), 134.44 (s), 133.39 (s), 132.38 (d, *J* = 3.7 Hz), 132.30 (d, *J* = 5.0 Hz), 130.02 (s), 129.30 (s), 128.08 (d, *J* = 1.3 Hz), 122.07 (q, *J* = 272.2 Hz), 120.71 (q, *J* = 34.0 Hz), 116.72 (d, *J* = 21.4 Hz), 115.96 (q, *J* = 288.5 Hz), 114.46 (d, *J* = 21.7 Hz), 50.51 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.21 (s, 3F), -68.01 (s, 3F), -109.54 (s, 1F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₀F₇NONa: 412.0548, found: 412.0526.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.46 (ddd, *J* = 22.7, 7.9, 0.8 Hz, 2H), 7.37 (t, *J* = 7.9 Hz, 2H), 7.31 (dd, *J* = 18.5, 7.7 Hz, 2H), 7.14 (d, *J* = 7.6 Hz, 1H), 6.07 (q, *J* = 8.0 Hz, 1H), 5.99 (d, *J* = 17.2 Hz, 1H), 4.27 (d, *J* = 17.2 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.58 (q, *J* = 36.8 Hz), 143.16 (q, *J* = 5.5 Hz), 137.01 (s), 135.72 (s), 134.32 (s), 133.11 (s), 132.85 (d, *J* = 2.5 Hz), 130.49 (s), 130.15 (s), 129.94 (s), 129.38 (s), 128.03 (s), 127.71 (s), 125.18 (s), 121.75 (q, *J* = 271.5 Hz), 121.12 (q, *J* = 34.7 Hz), 116.02 (q, *J* = 288.5 Hz), 50.29 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -62.76 (s, 3F), -67.96 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₀ClF₆NONa: 428.0253, found: 428.0233.



¹H NMR (400 MHz, CDCl₃) δ ppm = 8.10 (dd, *J* = 8.0, 1.6 Hz, 1H), 8.05 (s, 1H), 7.49 (d, *J* = 8.0 Hz, 1H), 7.41 (d, *J* = 8.9 Hz, 1H), 7.37 (dd, *J* = 7.5, 1.5 Hz, 1H), 7.31 (t, *J* = 7.0 Hz, 1H), 7.16 (d, *J* = 7.6 Hz, 1H), 6.15 (q, *J* = 8.0 Hz, 1H), 6.00 (d, *J* = 17.2 Hz, 1H), 4.33 (d, *J* = 17.2 Hz, 1H), 3.95 (s, 3H).
¹³C NMR (126 MHz, CDCl₃) δ ppm = 165.31 (s), 156.69 (q, *J* = 36.9 Hz), 147.71 (q, *J* = 6.3 Hz), 140.82 (s), 136.06 (s), 134.01 (s), 133.52 (s), 132.16 (s), 130.49 (s), 130.17 (s), 129.40 (s), 129.07 (d, *J* = 3.0 Hz), 128.11 (d, *J* = 3.8 Hz), 121.96 (q, *J* = 272.2 Hz), 120.68 (q, *J* = 34.7 Hz), 116.00 (q, *J* = 288.5 Hz), 52.64 (s), 50.48 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -57.30 (s, 3F), -67.85 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₀H₁₃F₆NO₃Na: 452.0697, found: 452.0681.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.48 – 7.40 (m, 3H), 7.37 – 7.34 (m, 1H), 7.22 (s, 1H), 7.17 (d, *J* = 7.9 Hz, 1H), 7.03 (d, *J* = 7.9 Hz, 1H), 6.11 (q, *J* = 8.1 Hz, 1H), 5.94 (d, *J* = 16.9 Hz, 1H), 4.29 (d, *J* = 17.0 Hz, 1H), 2.36 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.79 (d, *J* = 36.5 Hz), 148.69 (dd, *J* = 11.3, 5.6 Hz), 137.72 (s), 136.37 (s), 135.83 (s), 134.55 (s), 130.65 (s), 130.60 (s), 129.97 (s), 129.72 (s), 129.32 (s), 128.75 (d, *J* = 3.0 Hz), 127.99 (s), 126.70 (d, *J* = 1.8 Hz), 122.19 (q, *J* = 270.9 Hz), 119.87 (q, *J* = 34.7 Hz), 116.11 (q, *J* = 288.5 Hz), 50.44 (s), 20.91 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.13 (s, 3F), -67.90 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₃F₆NONa: 408.0799, found: 408.0783.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.49 – 7.41 (m, 3H), 7.36 (d, *J* = 6.2 Hz, 1H), 7.08 – 7.04 (m, 1H), 6.91 (dq, *J* = 5.3, 2.7 Hz, 2H), 6.13 (q, *J* = 8.0 Hz, 1H), 5.90 (d, *J* = 16.7 Hz, 1H), 4.26 (d, *J* = 16.7 Hz, 1H), 3.84 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 158.89 (s), 156.74 (q, *J* = 36.7 Hz), 148.56 (q, *J* = 5.5 Hz), 136.03 (s), 135.84 (s), 135.66 (s), 130.05 (s), 129.31 (d, *J* = 4.9 Hz), 128.77 (d, *J* = 2.9 Hz), 126.71 (d, *J* = 1.7 Hz), 125.46 (s), 122.12 (q, *J* = 270.9 Hz), 120.20 (q, *J* = 34.7 Hz), 116.11 (q, *J* = 288.5 Hz), 115.62 (s), 114.28 (s), 55.55 (s), 50.15 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.18 (s, 3F), -67.89 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₃F₆NO₂Na: 424.0748, found: 424.0736.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.44 (ddd, *J* = 13.8, 6.9, 1.7 Hz, 2H), 7.37 (t, *J* = 7.1 Hz, 2H), 7.26 (t, *J* = 8.1 Hz, 1H), 6.99 (d, *J* = 7.8 Hz, 1H), 6.88 (d, *J* = 8.2 Hz, 1H), 6.13 (q, *J* = 8.1 Hz, 1H), 6.03 (d, *J* = 18.0 Hz, 1H), 4.11 (d, *J* = 18.0 Hz, 1H), 3.83 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.88 (q, *J* = 36.4 Hz), 156.84 (s), 148.42 (q, *J* = 5.6 Hz), 136.62 (s), 135.99 (s), 135.77 (s), 129.89 (s), 129.26 (s), 128.45 (d, *J* = 3.8 Hz), 128.31 (s), 126.66 (d, *J* = 1.8 Hz), 122.35 (s), 122.17 (q, *J* = 271.5 Hz), 121.21 (s), 119.73 (q, *J* = 34.3 Hz), 116.15 (q, *J* = 290.4 Hz), 110.82 (s), 55.63 (s), 46.88 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.34 (s, 3F), -67.85 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₃F₆NO₂Na: 424.0748, found: 424.0730.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.46 – 7.41 (m, 2H), 7.41 – 7.33 (m, 3H), 7.05 (d, *J* = 16.4 Hz, 2H), 6.11 (q, *J* = 8.1 Hz, 1H), 5.91 (d, *J* = 17.2 Hz, 1H), 4.09 (d, *J* = 17.2 Hz, 1H), 2.33 (s, 3H), 2.23 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.94 (q, *J* = 36.5 Hz), 149.22 (q, *J* = 5.6 Hz), 137.20 (s), 136.96 (s), 136.26 (s), 135.53 (s), 135.00 (s), 132.75 (s), 129.80 (s), 129.36 (s), 128.61 (s), 128.31 (d, *J* = 2.9 Hz), 128.06 (s), 126.48 (s), 122.25 (q, *J* = 271.5 Hz), 119.23 (q, *J* = 34.2 Hz), 116.13 (q, *J* = 288.5 Hz), 49.03 (s), 20.74 (s), 19.39 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.31 (s, 3F), -67.84 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₀H₁₅F₆NONa: 422.0956, found: 422.0945.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.46 – 7.41 (m, 2H), 7.38 (d, *J* = 8.1 Hz, 2H), 7.14 (dd, *J* = 22.9, 8.0 Hz, 2H), 6.09 (q, *J* = 8.1 Hz, 1H), 5.98 (d, *J* = 17.1 Hz, 1H), 4.15 (d, *J* = 17.1 Hz, 1H), 2.30 (s, 3H), 2.16 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.97 (q, *J* = 36.6 Hz), 149.21 (dd, *J* = 11.3, 5.7 Hz), 138.82 (s), 137.09 (s), 135.51 (s), 134.74 (s), 133.09 (s), 131.48 (s), 129.80 (s), 129.34 (d, *J* = 12.3 Hz), 128.23 (d, *J* = 2.8 Hz), 127.03 (s), 126.33 (s), 122.29 (q, *J* = 270.9 Hz), 118.81 (q, *J* = 34.2 Hz), 116.13 (q, *J* = 288.5 Hz), 49.83 (s), 20.63 (s), 14.71 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.20 (s, 3F), -67.86 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₀H₁₅F₆NONa: 422.0956, found: 422.0940.



¹H NMR (400 MHz, CDCl₃) δ ppm = 7.49 (tt, *J* = 7.5, 5.9 Hz, 2H), 7.42 (t, *J* = 8.1 Hz, 3H), 7.35 (dd, *J* = 7.8, 1.1 Hz, 1H), 7.29 (d, *J* = 7.7 Hz, 1H), 6.19 – 6.11 (m, 2H), 4.24 (d, *J* = 17.8 Hz, 1H).
¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.98 (q, *J* = 36.7 Hz), 147.69 (q, *J* = 5.6 Hz), 137.16 (s), 136.19 (s), 135.39 (s), 133.92 (s), 131.33 (s), 130.96 (s), 130.29 (s), 129.60 (s), 128.59 (s), 128.45 (d, *J* = 2.5 Hz), 128.14 (s), 126.72 (d, *J* = 1.8 Hz), 121.95 (q, *J* = 271.5 Hz), 120.50 (q, *J* = 34.6 Hz), 116.03 (q, *J* = 288.5 Hz), 49.45 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.59 (s, 3F), -67.92 (s, 3F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{18}H_{10}ClF_6NONa$: 428.0253, found: 428.0243.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.51 – 7.45 (m, 2H), 7.42 (s, 1H), 7.38 (d, *J* = 7.5 Hz, 1H), 7.16 – 7.12 (m, 2H), 7.07 (td, *J* = 8.2, 2.6 Hz, 1H), 6.14 (q, *J* = 7.9 Hz, 1H), 5.94 (d, *J* = 17.0 Hz, 1H), 4.29 (d, *J* = 16.9 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 161.69 (d, J = 248.2 Hz), 156.77 (q, J = 36.8 Hz), 147.49 (q, J = 5.0 Hz), 136.35 (d, J = 7.1 Hz), 135.82 (s), 135.53 (s), 130.37 (s), 129.91 (d, J = 8.2 Hz), 129.52 (s), 129.43 (d, J = 3.3 Hz), 128.90 (q, J = 2.9 Hz), 126.75 (d, J = 1.8 Hz), 121.95 (q, J = 272.2 Hz), 121.07 (q, J = 34.0 Hz), 116.93 (d, J = 21.4 Hz), 116.07 (q, J = 288.5 Hz), 115.90 (d, J = 22.9 Hz), 50.11 (s).
¹⁹F NMR (376 MHz, CDCl₃) δ ppm = -57.33 (s, 3F), -67.94 (s, 3F), -114.11 (s, 1F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{18}H_{10}F_7NONa$: 412.0548, found: 412.0531.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.49 – 7.44 (m, 2H), 7.42 (d, *J* = 5.0 Hz, 2H), 7.38 (d, *J* = 7.4 Hz,

1H), 7.32 (d, *J* = 8.3 Hz, 1H), 7.10 (d, *J* = 8.3 Hz, 1H), 6.15 (q, *J* = 7.9 Hz, 1H), 5.94 (d, *J* = 17.2 Hz, 1H), 4.29 (d, *J* = 17.3 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.81 (q, *J* = 36.9 Hz), 147.31 (q, *J* = 5.6 Hz), 136.21 (s), 135.74 (s), 135.60 (s), 133.49 (s), 132.23 (s), 130.38 (s), 129.78 (s), 129.54 (d, *J* = 7.4 Hz), 128.99 (s), 128.91 (q, *J* = 2.9 Hz), 126.78 (d, *J* = 1.7 Hz), 121.92 (q, *J* = 271.5 Hz), 121.06 (q, *J* = 34.7 Hz), 116.05 (q, *J* = 288.5 Hz), 50.15 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.36 (s, 3F), -67.94 (s, 3F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{18}H_{10}ClF_6NONa$: 428.0253, found: 428.0239.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.57 (d, *J* = 1.8 Hz, 1H), 7.49 – 7.44 (m, 3H), 7.41 (d, *J* = 7.3 Hz, 1H), 7.37 (d, *J* = 7.5 Hz, 1H), 7.03 (d, *J* = 8.3 Hz, 1H), 6.14 (q, *J* = 7.9 Hz, 1H), 5.92 (d, *J* = 17.2 Hz, 1H), 4.26 (d, *J* = 17.2 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.83 (q, *J* = 36.9 Hz), 147.15 (q, *J* = 5.6 Hz), 136.50 (s), 135.71 (s), 135.61 (s), 132.75 (s), 132.71 (s), 131.84 (s), 130.38 (s), 129.70 (s), 129.58 (s), 128.89 (q, *J* = 2.7 Hz), 126.77 (d, *J* = 1.7 Hz), 121.89 (q, *J* = 272.2 Hz), 121.34 (s), 121.10 (q, *J* = 35.3 Hz), 116.02 (q, *J* = 288.5 Hz), 50.20 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -57.34 (s, 3F), -67.94 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₀BrF₆NONa: 471.9748, found: 471.9733.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 8.32 (d, *J* = 2.2 Hz, 1H), 8.19 (dd, *J* = 8.5, 2.3 Hz, 1H), 7.53 – 7.48 (m, 2H), 7.47 – 7.45 (m, 1H), 7.41 (d, *J* = 7.5 Hz, 1H), 7.37 (d, *J* = 8.5 Hz, 1H), 6.26 (q, *J* = 7.7 Hz, 1H), 6.08 (d, *J* = 17.9 Hz, 1H), 4.41 (d, *J* = 17.8 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.97 (q, *J* = 37.1 Hz), 147.20 (s), 146.46 (q, *J* = 5.5 Hz), 141.08 (s), 136.16 (s), 135.58 (s), 135.05 (s), 130.80 (s), 129.89 (s), 129.52 (s), 129.08 (d, *J* = 2.9 Hz), 126.87

(d, J = 1.7 Hz), 124.38 (s), 124.15 (s), 122.41 (q, J = 35.3 Hz), 121.68 (q, J = 272.2 Hz), 115.94 (q, J =

288.5 Hz), 50.47 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.57 (s, 3F), -68.00 (s, 3F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{18}H_{10}F_6N_2O_3Na$: 439.0493, found: 439.0480.



¹**H** NMR (400 MHz, CDCl₃) δ ppm = 7.67 (s, 1H), 7.61 (d, *J* = 8.1 Hz, 1H), 7.51 – 7.43 (m, 3H), 7.39 (d, *J* = 7.4 Hz, 1H), 7.30 (d, *J* = 8.1 Hz, 1H), 6.19 (q, *J* = 7.8 Hz, 1H), 6.04 (d, *J* = 17.5 Hz, 1H), 4.37 (d, *J* = 17.5 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.93 (q, J = 36.9 Hz), 147.30 (q, J = 6.3 Hz), 137.81 (s), 135.68 (s), 135.53 (s), 135.45 (s), 130.52 (s), 130.44 (q, J = 32.8 Hz), 129.69 (s), 128.97 (d, J = 2.5 Hz), 128.86 (s), 126.79 (d, J = 1.7 Hz), 126.35 (q, J = 3.8 Hz), 126.21 (d, J = 2.5 Hz), 123.50 (q, J = 272.2 Hz), 121.81 (q, J = 272.2 Hz), 121.59 (q, J = 34.0 Hz), 115.99 (q, J = 288.5 Hz), 50.43 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.43 (s, 3F), -62.63 (s, 3F), -67.97 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₀F₉NONa: 462.0516, found: 462.0501.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.94 (d, *J* = 8.2 Hz, 1H), 7.79 (d, *J* = 8.0 Hz, 1H), 7.47 (t, *J* = 7.8 Hz, 4H), 7.43 – 7.36 (m, 2H), 6.31 (q, *J* = 8.0 Hz, 1H), 6.19 (d, *J* = 17.4 Hz, 1H), 4.38 (d, *J* = 17.4 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.82 (q, *J* = 36.8 Hz), 140.69 (q, *J* = 5.7 Hz), 138.64 (s), 137.34 (d, *J* = 15.7 Hz), 136.87 (s), 135.21 (s), 130.03 (s), 129.61 (s), 128.81 (s), 128.16 (s), 127.58 (s), 125.28 (d, *J* = 4.7 Hz), 122.71 (s), 122.46 (q, *J* = 272.2 Hz), 122.15 (s), 121.11 (q, *J* = 34.5 Hz), 116.07 (q, *J* = 288.5 Hz), 47.41 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -56.66 (s, 3F), -67.75 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₀H₁₁F₆NOSNa: 450.0363, found: 450.0355.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.45 (ddd, *J* = 7.6, 6.1, 3.0 Hz, 1H), 7.41 – 7.38 (m, 2H), 7.32 (ddd, *J* = 7.5, 6.6, 5.6 Hz, 3H), 7.23 (d, *J* = 6.8 Hz, 1H), 7.14 (d, *J* = 7.6 Hz, 1H), 6.08 (q, *J* = 8.1 Hz, 2H), 4.15 (d, *J* = 18.8 Hz, 1H), 1.93 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 170.42 (s), 150.91 (q, J = 5.8 Hz), 138.88 (s), 137.16 (s), 135.81 (s), 135.12 (s), 130.34 (s), 129.54 (s), 128.83 (s), 128.64 (s), 128.52 (q, J = 2.5 Hz), 128.06 (s), 127.25 (s), 127.05 (s), 122.50 (q, J = 272.2 Hz), 119.36 (q, J = 34.0 Hz), 48.10 (s), 21.49 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -55.64 (s, 3F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{18}H_{14}F_3NONa$: 340.0925, found: 340.0913.



¹H NMR (500 MHz, CDCl₃) δ ppm = 7.45 (ddd, J = 18.5, 10.9, 4.1 Hz, 4H), 7.38 (dd, J = 11.3, 3.6 Hz, 2H), 7.31 (t, J = 7.3 Hz, 1H), 7.16 (d, J = 7.6 Hz, 1H), 6.08 – 5.99 (m, 2H), 4.35 (d, J = 17.1 Hz, 1H).
¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.85 (q, J = 36.7 Hz), 151.20 (t, J = 4.5 Hz), 136.63 (s), 135.58 (s), 135.37 (s), 133.55 (s), 130.03 (s), 129.94 (s), 129.33 (s), 128.61 (t, J = 4.2 Hz), 128.04 (s), 127.94 (s), 126.68 (d, J = 1.7 Hz), 118.97 (m), 117. 20 (m), 116.11 (m), 113.8 (m), 50.48 (s).

¹⁹F NMR (376 MHz, CDCl₃) δ ppm = -67.95 (s, 3F), -85.22 (s, 3F), -106.10 (dt, J = 278.2, 7.5 Hz, 1F),
-108.85 (dq, J = 278.2, 11.3 Hz, 1F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₉H₁₁F₈NONa: 444.0611, found: 444.0594.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.46 (dd, *J* = 7.1, 2.7 Hz, 1H), 7.40 (dd, *J* = 15.6, 6.8 Hz, 4H), 7.32 (dd, *J* = 15.1, 7.9 Hz, 2H), 7.16 (d, *J* = 7.4 Hz, 1H), 6.06 (dd, *J* = 32.2, 16.1 Hz, 2H), 4.35 (d, *J* = 17.2 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.86 (q, J = 36.7 Hz), 151.11 (t, J = 4.6 Hz), 136.68 (s), 135.52 (s), 133.53 (s), 130.02 (s), 129.93 (s), 129.34 (s), 129.27 (s), 128.47 (s), 128.03 (s), 127.95 (s), 126.69 (d, J = 1.7 Hz), 118.14 (m), 117.15 (m), 115.75 (m), 114.74 (m), 111.71 (m), 50.43 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -67.96 (s, 3F), -80.21 (s, 3F), -106.10 (tt, *J* = 7.5, 3.2 Hz, 1F). -108.84 (ddd, *J* = 23.9, 10.8, 10.7 Hz, 1F), -126.98 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₀H₁₁F₁₀NONa: 494.0579, found: 494.0568.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.50 (dd, *J* = 12.9, 6.1 Hz, 2H), 7.44 (dd, *J* = 15.7, 7.5 Hz, 2H), 7.38 – 7.30 (m, 3H), 7.16 (d, *J* = 7.3 Hz, 1H), 6.23 – 6.17 (m, 1H), 6.02 – 5.79 (m, 2H), 4.37 (d, *J* = 16.9 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.25 (q, *J* = 36.2 Hz), 147.07 (t, *J* = 12.8 Hz), 136.62 (d, *J* = 15.1 Hz), 134.55 (s), 133.28 (s), 130.25 (s), 129.76 (s), 129.54 (s), 129.34 (s), 128.00 (s), 127.93 (s), 127.30 (d, *J* = 1.5 Hz), 125.26 (dd, *J* = 31.8, 22.4 Hz), 116.24 (q, *J* = 288.5 Hz), 112.81 (t, *J* = 230.7 Hz), 50.80 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -68.12 (s, 3F), -104.62 (ddd, *J* = 319.6, 56.4, 3.8 Hz, 1F), -103.14 (ddd, *J* = 319.6, 54.5, 9.4 Hz, 1F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₂F₅NONa: 376.0737, found: 376.0725.



¹H NMR (500 MHz, CDCl₃) δ ppm = 7.46 – 7.41 (m, 4H), 7.37 – 7.33 (m, 2H), 7.30 (t, *J* = 7.3 Hz, 1H),
7.15 (d, *J* = 7.5 Hz, 1H), 6.30 – 6.25 (m, 1H), 5.98 (d, *J* = 17.1 Hz, 1H), 4.33 (d, *J* = 17.1 Hz, 1H).
¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.83 (q, *J* = 36.6 Hz), 143.95 (d, *J* = 8.1 Hz), 136.48 (s), 135.82 (s), 134.80 (s), 133.78 (s), 129.88 (s), 129.74 (s), 129.30 (s), 128.83 (t, *J* = 3.8 Hz), 128.05 (s), 127.90 (s), 126.70 (s), 125.97 (s), 125.76 (d, *J* = 6.8 Hz), 121.55 (s), 116.11 (q, *J* = 288.5 Hz), 50.55 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -45.50 (dd, *J* = 159.8, 5.6 Hz, 1F), -47.31 (dt, *J* = 157.9, 13.2 Hz,

1F). -67.79 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₁₈H₁₁ClF₅NONa: 410.0347, found: 410.0338.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.35 – 7.29 (m, 2H), 7.27 (d, *J* = 6.1 Hz, 1H), 7.23 (d, *J* = 7.8 Hz, 1H), 7.18 (d, *J* = 6.6 Hz, 1H), 7.09 (t, *J* = 7.7 Hz, 1H), 6.69 (t, *J* = 8.0 Hz, 1H), 6.51 (d, *J* = 8.2 Hz, 1H), 5.71 (q, *J* = 8.6 Hz, 1H), 4.81 (s, 1H), 3.94 (s, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 154.10 (q, *J* = 5.8 Hz), 144.26 (s), 142.18 (s), 135.54 (s), 131.12 (q, *J* = 3.6 Hz), 130.41 (s), 129.03 (s), 128.31 (s), 126.44 (s), 125.42 (d, *J* = 0.5 Hz), 122.89 (q, *J* =

271.5 Hz), 119.00 (s), 117.55 (s), 117.25 (s), 117.15 (q, J = 33.4 Hz), 46.99 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -56.17 (s, 3F).

HRMS (ESI) (m/z): [M+H]⁺ called. for C₁₆H₁₃F₃N: 276.1000, found: 276.0992.



¹**H** NMR (400 MHz, CDCl₃) δ ppm = 8.76 (s, 1H), 7.60 – 7.54 (m, 2H), 7.52 – 7.47 (m, 2H), 7.42 (ddd, J = 8.0, 4.5, 1.8 Hz, 2H), 7.33 (dt, J = 14.7, 7.0 Hz, 2H), 5.76 (q, J = 8.0 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 159.44 (s), 148.79 (q, *J* = 5.7 Hz), 143.40 (s), 141.84 (s), 132.18 (s), 130.44 (s), 129.38 (s), 129.23 (s), 128.81 (s), 127.80 (q, *J* = 3.0 Hz), 127.32 (s), 125.72 (d, *J* = 0.5

Hz), 122.23 (q, *J* = 271.2 Hz), 119.78 (q, *J* = 34.3 Hz).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -56.51 (s, 3F).

HRMS (ESI) (m/z): [M+H]⁺ called. for C₁₆H₁₁F₃NO: 290.0793, found: 290.0782.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 8.76 (s, 1H), 7.57 (ddd, *J* = 7.9, 7.1, 2.0 Hz, 2H), 7.52 – 7.48 (m, 2H), 7.42 (ddd, *J* = 8.0, 4.5, 1.7 Hz, 2H), 7.33 (dt, *J* = 13.5, 7.1 Hz, 2H), 5.76 (q, *J* = 8.0 Hz, 1H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 159.44 (s), 148.77 (q, J = 5.7 Hz), 143.37 (s), 141.84 (s), 132.20 (s), 130.43 (s), 129.39 (s), 129.24 (s), 128.82 (s), 127.80 (q, J = 3.0 Hz), 127.33 (d, J = 2.9 Hz), 125.73 (s), 122.22 (q, J = 271.3 Hz), 119.79 (q, J = 34.4 Hz).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -56.52 (s, 3F).

HRMS (ESI) (m/z): $[M+H]^+$ called. for $C_{16}H_{11}F_3N$: 274.0844, found: 274.0829.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.60 – 7.48 (m, 3H), 7.42 (qd, *J* = 7.3, 1.7 Hz, 2H), 7.35 (dtt, *J* = 6.5, 4.9, 1.5 Hz, 2H), 7.23 (dt, *J* = 17.9, 7.5 Hz, 4H), 7.10 (t, *J* = 7.8 Hz, 1H), 6.88 (s, 1H), 6.74 (d, *J* = 7.9 Hz, 1H), 6.25 (q, *J* = 8.3 Hz, 2H), 3.68 (qd, *J* = 10.3, 9.8, 4.1 Hz, 1H), 1.80 (dd, *J* = 11.6, 4.8 Hz, 1H), 1.75 – 1.52 (m, 4H), 1.32 (ddt, *J* = 22.2, 11.3, 5.8 Hz, 2H), 1.23 – 0.96 (m, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 172.70 (s), 167.60 (s), 150.52 (s), 137.10 (s), 135.38 (s), 134.68 (s), 134.14 (s), 133.69 (s), 130.90 (s), 130.63 (s), 129.87 (s), 129.52 (s), 129.50 (s), 129.24 (s), 128.73 (s), 128.37 (s), 127.96 (s), 127.77 (s), 122.50 (q, *J* = 270.9 Hz), 119.02 (q, *J* = 34.0 Hz), 61.69 (s), 47.96 (s), 33.00 (s), 32.74 (s), 26.91 (s), 25.39 (s), 24.53 (s), 24.50 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -56.02 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₃₀H₂₇F₃N₂O₂Na: 527.1930, found: 527.1923.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.47 – 7.40 (m, 4H), 7.35 (t, *J* = 7.3 Hz, 2H), 7.31 (d, *J* = 7.4 Hz, 1H), 7.15 (d, *J* = 7.5 Hz, 1H), 6.18 (t, *J* = 14.3 Hz, 1H), 6.00 (d, *J* = 16.9 Hz, 1H), 4.35 (d, *J* = 17.0 Hz, 1H), 3.93 (s, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 164.22 (t, J = 34.4 Hz), 156.85 (q, J = 36.5 Hz), 147.44 (t, J = 5.3 Hz)

Hz), 136.96 (s), 135.72 (d, *J* = 8.6 Hz), 133.37 (s), 129.71 (s), 129.52 (s), 129.44 (s), 129.15 (s), 129.92 (s), 129.79 (s), 126.47 (s), 122.50 (t, *J* = 23.4 Hz), 116.11 (q, *J* = 288.0 Hz), 111.65 (t, *J* = 251.0 Hz), 53.73 (s), 50.55 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -67.81 (s, 3F), -94.81 (dd, *J* = 270.7, 7.5 Hz, 1F), -102.17 (dd, *J* = 270.7, 7.5 Hz, 1F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₀H₁₄F₅NO₃Na: 434.0792, found: 434.0774.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.65 – 7.62 (m, 1H), 7.42 – 7.38 (m, 3H), 7.30 (dd, *J* = 13.4, 6.1 Hz, 3H), 7.11 (d, *J* = 7.4 Hz, 1H), 6.12 (dd, *J* = 17.0, 13.9 Hz, 1H), 5.97 (d, *J* = 17.1 Hz, 1H), 4.35 – 4.26 (m, 5H), 1.40 (t, *J* = 7.0 Hz, 6H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.88 (q, *J* = 36.6 Hz), 147.28 (dd, *J* = 12.0, 8.2 Hz), 137.48 (s), 136.00 (s), 135.58 (s), 133.34 (s), 129.59 (s), 129.45 (s), 129.41 (s), 129.19 (s), 129.14 (s), 127.85 (s), 127.75 (s), 126.36 (d, *J* = 1.6 Hz), 122.00 (d, *J* = 14.2 Hz), 121.69 (d, *J* = 14.1 Hz), 116.78 (d, *J* = 220.0 Hz), 116.14 (q, *J* = 288.54 Hz), 64.88 (dd, *J* = 29.3, 6.9 Hz), 50.52 (s), 16.44 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -67.77 (s, 3F), -104.29 (ddd, *J* = 303.3, 108.5, 5.8 Hz, 1F), -106.89 (ddd, *J* = 303.1, 112.4, 8.2 Hz, 1F).

³¹**P** NMR (162 MHz, CDCl₃) δ ppm = 6.44 (t, *J* = 110.16 Hz, 1P).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₂H₂₁F₅NO₄PNa: 512.1026, found: 512.1045.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.51 (dd, *J* = 7.8, 1.5 Hz, 1H), 7.30 (tt, *J* = 5.1, 2.3 Hz, 4H), 7.20 (dd, *J* = 6.5, 3.0 Hz, 2H), 7.14 (td, *J* = 7.7, 1.6 Hz, 1H), 6.72 (d, *J* = 7.9 Hz, 1H), 5.66 (d, *J* = 14.1 Hz, 1H), 4.27 (d, *J* = 14.0 Hz, 1H), 2.43 (t, *J* = 7.0 Hz, 2H), 1.69 – 1.61 (m, 2H), 1.07 (t, *J* = 7.4 Hz, 3H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 157.17 (q, J = 35.8 Hz), 139.56 (s), 135.30 (s), 132.94 (s), 129.89 (s), 129.51 (s), 128.95, 128.51 (s), 128.07 (s), 127.67 (s), 123.91 (s), 116.37 (q, J = 288.6 Hz), 97.06 (s), 76.17 (s), 53.80 (s), 22.07 (s), 21.51 (s), 13.52 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -68.36 (s, 3F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{20}H_{18}F_3NONa$: 368.1540, found: 368.1523.



¹H NMR (400 MHz, CDCl₃) δ ppm = 7.65 (dd, J = 7.8, 1.6 Hz, 1H), 7.55 (ddd, J = 5.1, 2.8, 1.6 Hz, 2H), 7.43 - 7.36 (m, 4H), 7.32 - 7.27 (m, 3H), 7.26 - 7.19 (m, 3H), 6.83 (d, J = 7.9 Hz, 1H), 5.72 (d, J = 14.0 Hz, 1H), 4.39 (d, J = 14.1 Hz, 1H).
¹³C NMR (126 MHz, CDCl₃) δ ppm = 157.33 (q, J = 36.0 Hz), 139.67 (s), 135.12 (s), 132.76 (s),

131.75 (s), 130.02 (s), 129.53 (s), 129.14 (s), 129.02 (s), 128.58 (s), 128.52 (s), 128.16 (s), 123.30

(s), 122.31 (s), 116.41 (q, *J* = 288.5 Hz), 95.61 (s), 84.56 (s), 54.07 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -68.28 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₃H₁₆F₃NONa: 402.1085, found: 402.1083.



¹**H NMR** (500 MHz, CDCl₃) δ ppm = 7.83 (d, *J* = 1.1 Hz, 1H), 7.78 (dd, *J* = 7.7, 1.2 Hz, 1H), 7.51 – 7.34 (m, 4H), 7.18 (d, *J* = 7.7 Hz, 1H), 6.21 (q, *J* = 8.0 Hz, 1H), 6.02 (d, *J* = 17.3 Hz, 1H), 4.36 (d, *J* = 17.4 Hz, 1H), 1.39 (s, 12H).

¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.86 (q, *J* = 36.6 Hz), 148.26 (q, *J* = 5.7 Hz), 136.67 (s), 136.43 (s), 135.95 (s), 135.71 (s), 135.52 (s), 129.96 (s), 129.37 (s), 128.79 (d, *J* = 3.1 Hz), 127.38 (s), 126.63

(d, *J* = 2.1 Hz), 122.14 (q, *J* = 271.1 Hz), 120.30 (q, *J* = 34.3 Hz), 116.04 (q, *J* = 287.7 Hz), 84.23 (s), 50.72 (s), 24.86 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -57.05 (p, J = 7.6 Hz, 3F), -67.88 (q, J = 7.4 Hz, 3F). **HRMS** (ESI) (m/z): [M+Na]⁺ called. for C₂₄H₂₂F₆NO₃BNa: 520.1500, found: 520.1513.



¹H NMR (400 MHz, CDCl₃) δ ppm = 8.60 (dd, J = 4.8, 1.6 Hz, 1H), 7.36-7.26 (m, 3H), 7.20-7.12 (m, 3H), 7.06-7.00 (m, 1H), 5.76 (d, J = 14.2 Hz, 1H), 4.26 (d, J = 14.2 Hz, 1H), 3.46 (s, 1H).
¹³C NMR (126 MHz, CDCl₃) δ ppm = 156.78 (q, J = 36.4 Hz), 150.41 (s), 142.05 (s), 137.80 (s), 137.14 (s), 134.44 (s), 129.48 (s), 128.84 (s), 128.54 (s), 123.20 (s), 116.10 (q, J = 288.5 Hz), 83.29 (s), 78.65 (s), 53.67 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -68.29 (s, 3F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{16}H_{11}F_3N_2ONa$: 327.0725, found: 327.0721.



¹**H NMR** (400 MHz, CDCl₃) δ ppm = 7.81 (dd, *J* = 7.7, 1.5 Hz, 1H), 7.49 (td, *J* = 7.6, 1.2 Hz, 1H), 7.41 (td, *J* = 7.7, 1.6 Hz, 1H), 7.29 – 7.25 (m, 3H), 7.15 (dd, *J* = 6.8, 2.6 Hz, 2H), 6.85 (d, *J* = 7.8 Hz, 1H), 5.43 (d, *J* = 14.3 Hz, 1H), 4.33 (d, *J* = 14.3 Hz, 1H), 2.45 (s, 3H).

¹³C NMR (101 MHz, CDCl₃) δ ppm = 198.14 (s), 156.65 (dd, J = 69.9, 34.4 Hz), 136.54 (s), 135.67 (s),
135.14 (s), 132.41 (s), 131.56 (s), 130.16 (s), 129.84 (s), 129.42 (s), 128.57 (s), 128.13 (s), 117.93 (s),
115.06 (s), 55.51 (s), 28.73 (s).

¹⁹**F NMR** (376 MHz, CDCl₃) δ ppm = -68.30 (s, 3F).

HRMS (ESI) (m/z): $[M+Na]^+$ called. for $C_{17}H_{14}F_3NO_2Na$: 344.0874, found: 344.0892.


¹**H NMR** (400 MHz, CDCl₃) δ ppm = 8.07 (s, 1H), 7.73 (d, *J* = 2.2 Hz, 1H), 7.53 (td, *J* = 8.2, 2.0 Hz, 1H), 7.46 (ddd, *J* = 14.7, 7.4, 1.7 Hz, 2H), 7.42 – 7.36 (m, 2H), 7.17 (d, *J* = 8.4 Hz, 1H), 6.15 (q, *J* = 7.9 Hz, 1H), 5.95 (d, *J* = 17.4 Hz, 1H), 4.32 (d, *J* = 17.1 Hz, 1H).

¹³C NMR (101 MHz, CDCl₃) δ ppm = 156.91 (q, *J* = 37.0 Hz), 154.96 (q, *J* = 38.0 Hz), 147.54 (q, *J* = 5.6 Hz), 135.82 (s), 135.60 (s), 134.68 (s), 131.69 (s), 130.37 (s), 129.58 (s), 129.23 (s), 128.91 (d, *J* = 2.8 Hz), 126.74 (d, *J* = 1.8 Hz), 123.22 (s), 121.68 (s), 121.24 (q, *J* = 34.6 Hz), 120.94 (s), 116.02 (d, *J* = 287.7 Hz), 115.57 (q, *J* = 288.9 Hz), 100.03 (s), 50.28 (s).

¹⁹**F** NMR (376 MHz, CDCl₃) δ ppm = -57.34 (s, 3F), -67.92 (s, 3F), -75.67 (s, 3F).

HRMS (ESI) (m/z): [M+Na]⁺ called. for C₂₀H₁₁F₉N₂O₂Na: 505.0575, found: 505.0591.

3. NMR spectra for the products

























165.25 165.71 165.72 165.71 155.70 156.71 156.71 156.71 156.71 153.35 13



4.33344.2991

































~-57.36 ---62.90 ~-67.94





156.94 156.63 156.63 156.63 156.63 156.63 174.73 147.73 174.73 174.73 173.74 172.73 173.83 173.74 173.73 173.74 172.75 172.74 172.74 172.75 172.74 172.75 172.74 172.75 175.75 17







65.380 65.721 65.721 65.721 65.721 65.642 65.642 65.642 65.642 65.642 137.052 137.













67.20 66.29 66.24 66.24 66.24 66.24 66.29 66.29 60.29 60.29 66.29 66.29 66.29 66.29 66.29 66.29 66.29 66.29 66.29 66.20 66.74 66.74 71.25.68 72.58



8,1155 8,1116 8,1116 8,0543 8,0543 8,0543 8,0543 7,7,3574 6,11778 6,11778 6,11778 6,11778 6,11778 6,11778 6,11778 6,11778 5,9775 5,9775 5,9775 5,9775 5,9775 5,9775







-165.31 -166.31 -166.34 -166.54 -16







































































162.67 165.32 165.32 165.32 165.32 165.32 165.32 165.32 165.32 165.32 165.32 165.34 165.34 175.55 1135.55 1739.52 1739.52 1739.52 1729.52 1729.55 1720.55 1720














6157.25 156.637 156.637 156.637 156.637 135.74 135.74 135.73 135.74 135.74 135.74 135.74 135.74 132.85 135.74 132.85 1

















157.27 156.39 156.39 156.39 147.18















157.41 155.52 156.552 146.552 146.552 144.655 144.655 144.655 144.655 144.655 144.655 145.656 145.656 145.656 172.656 172.650 172.455







~-57.43 --62.63 --67.97































10 0 -10 -20 -30 -40 -50 -60 -70 -80 -90 -100 -120 -140 -160 -180 -200 f1 (ppm)









157.29 156.67 156.67 156.67 156.67 156.67 156.67 156.67 156.67 135.56 137.56 13









-67.96 -6





157.29 155.70 155.64 155.64 155.64 151.15 151.15 151.15 151.15 151.15 151.15 151.15 151.15 151.15 113.29 113.56 113.56 113.56 115.75 115.75 115.75 115.57 115.53 115.55



7,5205 7,4848 7,4848 7,4846 7,4846 7,4846 7,4850 7,4850 7,7329 7,





-68.12 -68.12 -68.12 -68.12 -7104.12 -7104.92 -7105.72 -7113.73 -7113.73 -7113.55 -7











45.28 45.70 45.70 45.72 47.07 47.07 47.07 47.79 67.79 67.79 67.79



















F₃C



F₃C



159.44 148.61 148.61 148.77 148.77 148.77 148.72 148.72 148.72 123.34 1227.88 1227.88 1227.88 1227.88 1227.88 1227.88 1227.88 1227.88 1227.89 1227.70 1227.70 1227.89 1227.70 1227.89 1227.70 127.70 127



F₃C



F₃C



- 159,44 148,86 148,87 148,75 148,75 148,75 148,75 143,37 143,37 143,37 143,37 143,37 143,37 143,37 143,37 1132,43 1132,73 1127,73 1137,73 113


















































