

**Visible light mediated oxidative lactonization of 2-methyl-1,1'-biaryls
for the synthesis of benzocoumarins**

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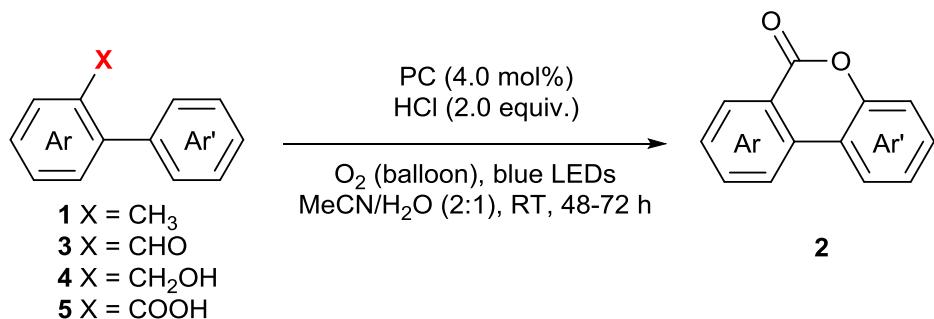
Part I Experimental part.....	S2
1. General information	S2
2. Photocatalyzed synthesis of benzocoumarins 2 (Scheme 2 & 3)	S2
3. Control experiments (Scheme 4).....	S7
4. References	S9
Part II NMR Spectra.....	S11

Part I Experimental part

1. General information

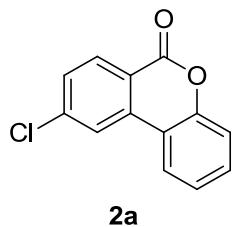
Unless otherwise indicated, all reagents were purchased from commercial suppliers and used without purification. 2-Methyl-1,1'-biphenyl¹, 2-phenylbenzaldehyde², and 2-phenylbenzyl alcohol³ derivatives were prepared according to literatures. Column chromatography was performed on silica gel 200~300 mesh. ¹H NMR, ¹³C NMR, and ¹⁹F NMR spectra were recorded at room temperature in CDCl₃ with tetramethylsilane (TMS) as internal standard and reported in parts per million (ppm, δ). ¹H NMR Spectroscopy splitting patterns were designated as singlet (s), doublet (d), triplet (t), quartet (q). Splitting patterns that could not be interpreted or easily visualized were designated as multiplet (m) or broad (br).

2. Photocatalyzed synthesis of benzocoumarins 2 (Scheme 2 & 3)



General procedure: To an oven-dried 10 mL Schlenk tube equipped with a stir bar was charged with substrate **1**, **3**, **4** or **5** (0.3 mmol) and Acr⁺MesClO₄⁻ (5.0 mg, 0.012 mmol). The tube was closed with a septum, evacuated, and back-filled with oxygen (balloon). To this mixture was added 1.0 mL solvent (MeCN/H₂O = 2:1) and HCl (150 μ m, 0.6 mmol, 4M in 1,3-dioxane). The reaction mixture was stirred at room temperature under blue LEDs until the full consumption of the starting materials (typically, 48 h). The reaction mixture was concentrated under reduced pressure, and the residue was purified by column chromatography on silica gel.

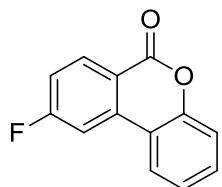
(petroleum ether/EtOAc = 70:1 - 20:1) to furnish the benzocoumarins **2**.



2a

9-chloro-6*H*-benzo[*c*]chromen-6-one⁴

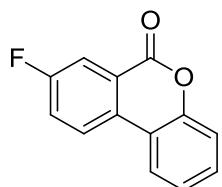
45.1 mg, 65% yield. White solid, ¹H NMR (400 MHz, Chloroform-*d*) δ 8.34 (d, *J* = 8.5 Hz, 1H), 8.08 (d, *J* = 1.9 Hz, 1H), 8.00 (dd, *J* = 8.0, 1.5 Hz, 1H), 7.53 (ddd, *J* = 9.4, 7.9, 1.8 Hz, 2H), 7.41 – 7.32 (m, 2H). ¹³C NMR (101 MHz, Chloroform-*d*) δ 160.4, 151.6, 141.9, 136.4, 132.3, 131.2, 129.3, 124.8, 122.9, 121.8, 119.6, 118.0, 117.0.



2b

9-fluoro-6*H*-benzo[*c*]chromen-6-one⁵

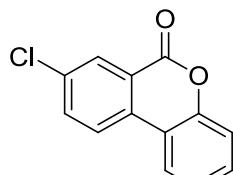
38.6 mg, 64% yield. White solid, ¹H NMR (400 MHz, Chloroform-*d*) δ 8.43 (dd, *J* = 8.8, 5.7 Hz, 1H), 7.96 (dd, *J* = 8.0, 1.5 Hz, 1H), 7.73 (dd, *J* = 9.7, 2.4 Hz, 1H), 7.52 (ddd, *J* = 8.5, 7.2, 1.5 Hz, 1H), 7.41 – 7.31 (m, 2H), 7.32 – 7.23 (m, 1H). ¹³C NMR (101 MHz, Chloroform-*d*) δ 167.0 (d, *J* = 256.7 Hz), 160.2, 151.6, 137.7 (d, *J* = 10.0 Hz), 133.9 (d, *J* = 10.2 Hz), 131.2, 124.7, 123.0, 117.9, 117.8, 117.34 (d, *J* = 2.9 Hz), 117.0 (d, *J* = 23.0 Hz), 108.2 (d, *J* = 23.6 Hz). ¹⁹F NMR (377 MHz, Chloroform-*d*) δ -101.1.



2c

8-fluoro-6*H*-benzo[*c*]chromen-6-one⁶

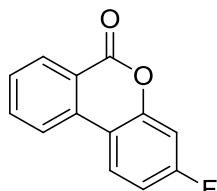
35.3 mg, 59% yield. White solid, ^1H NMR (400 MHz, Chloroform-*d*) δ 8.13 (dd, *J* = 8.9, 4.8 Hz, 1H), 8.08 – 7.95 (m, 2H), 7.62 – 7.42 (m, 2H), 7.41 – 7.30 (m, 2H). ^{13}C NMR (101 MHz, Chloroform-*d*) δ 162.4 (d, *J* = 252.5 Hz), 160.0 (d, *J* = 256.7 Hz), 150.8, 131.2 (d, *J* = 2.5 Hz), 130.4, 124.8, 124.3 (d, *J* = 7.6 Hz), 123.2 (d, *J* = 8.9 Hz), 122.9 (d, *J* = 22.7 Hz), 122.6, 117.8, 117.4, 116.2 (d, *J* = 22.7 Hz). ^{19}F NMR (377 MHz, Chloroform-*d*) δ -110.2.



2d

8-chloro-6*H*-benzo[*c*]chromen-6-one⁶

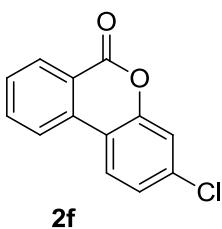
42.2 mg, 61% yield. White solid, ^1H NMR (400 MHz, Chloroform-*d*) δ 8.45 (d, *J* = 8.0 Hz, 1H), 8.19 (t, *J* = 8.1 Hz, 2H), 7.96 – 7.86 (m, 1H), 7.69 (t, *J* = 7.6 Hz, 1H), 7.65 – 7.56 (m, 2H). ^{13}C NMR (101 MHz, Chloroform-*d*) δ 160.3, 151.0, 135.2, 133.4, 130.9, 130.2, 123.6, 122.2, 121.7, 121.1, 121.1, 115.3, 115.3.



2e

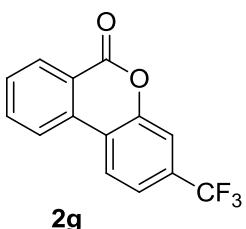
3-fluoro-6*H*-benzo[*c*]chromen-6-one⁷

44.1 mg, 74% yield. White solid, ^1H NMR (400 MHz, Chloroform-*d*) δ 8.37 (d, *J* = 7.9 Hz, 1H), 8.03 (dd, *J* = 9.2, 5.7 Hz, 2H), 7.83 (t, *J* = 7.7 Hz, 1H), 7.57 (t, *J* = 7.6 Hz, 1H), 7.08 (ddd, *J* = 7.7, 4.5, 1.9 Hz, 2H). ^{13}C NMR (101 MHz, Chloroform-*d*) δ 163.4 (d, *J* = 251.5 Hz), 160.8, 152.2 (d, *J* = 12.1 Hz), 135.1, 134.2, 130.7, 128.7, 124.3 (d, *J* = 10.1 Hz), 121.5, 120.4, 114.6, 112.4 (d, *J* = 22.2 Hz), 105.1 (d, *J* = 25.2 Hz). ^{19}F NMR (377 MHz, Chloroform-*d*) δ -108.4.



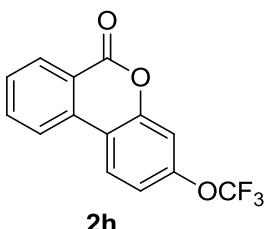
3-chloro-6*H*-benzo[*c*]chromen-6-one⁷

40.8 mg, 60% yield. White solid, ¹H NMR (500 MHz, Chloroform-*d*) δ 8.40 (dd, *J* = 8.0, 1.3 Hz, 1H), 8.08 (d, *J* = 8.1 Hz, 1H), 7.99 (d, *J* = 8.6 Hz, 1H), 7.84 (ddd, *J* = 8.3, 7.3, 1.4 Hz, 1H), 7.61 (ddd, *J* = 8.1, 7.3, 1.1 Hz, 1H), 7.39 (d, *J* = 2.0 Hz, 1H), 7.32 (dd, *J* = 8.5, 2.1 Hz, 1H). ¹³C NMR (126 MHz, Chloroform-*d*) δ 160.6, 151.6, 136.0, 135.1, 134.0, 130.8, 129.2, 125.0, 123.8, 121.7, 121.0, 118.0, 116.7.



3-(trifluoromethyl)-6*H*-benzo[*c*]chromen-6-one⁷

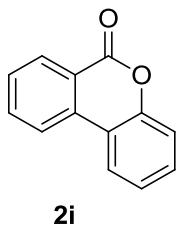
39.3 mg, 50% yield. White solid, ¹H NMR (400 MHz, Chloroform-*d*) δ 8.40 (dd, *J* = 7.9, 1.3 Hz, 1H), 8.15 (t, *J* = 7.7 Hz, 2H), 7.88 (td, *J* = 7.7, 1.4 Hz, 1H), 7.72 – 7.63 (m, 1H), 7.57 (d, *J* = 9.0 Hz, 2H). ¹³C NMR (101 MHz, Chloroform-*d*) δ 160.3, 150.9, 135.2, 132.2 (q, *J* = 33.5 Hz), 130.8, 130.1, 123.6, 123.3 (q, *J* = 272.5 Hz), 122.2, 121.6, 121.1 (q, *J* = 3.6 Hz), 121.06, 115.2 (q, *J* = 4.0 Hz). ¹⁹F NMR (377 MHz, Chloroform-*d*) δ -62.8.



3-(trifluoromethoxy)-6*H*-benzo[*c*]chromen-6-one⁸

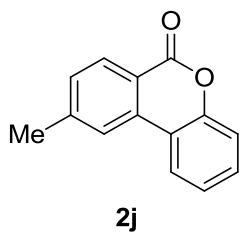
45.2 mg, 54% yield. White solid, ¹H NMR (500 MHz, Chloroform-*d*) δ 8.40 (dd, *J* = 8.0, 1.4 Hz, 1H), 8.08 (dd, *J* = 8.4, 3.0 Hz, 2H), 7.86 (td, *J* = 7.7, 1.4 Hz, 1H), 7.62 (t, *J* = 7.6 Hz, 1H), 7.31 – 7.14 (m, 2H). ¹³C NMR (126 MHz, Chloroform-*d*) δ 160.5,

151.8, 150.2, 135.2, 133.8, 130.8, 129.3, 124.2, 121.7, 120.9, 120.3 (q, $J = 259.6$ Hz), 117.0, 116.8, 110.2. ^{19}F NMR (377 MHz, Chloroform-*d*) δ -57.8.



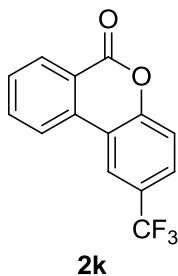
6H-benzo[c]chromen-6-one⁶

28.2 mg, 48% yield. White solid, ^1H NMR (400 MHz, Chloroform-*d*) δ 8.39 (dd, $J = 7.9, 1.4$ Hz, 1H), 8.11 (d, $J = 8.1$ Hz, 1H), 8.05 (dd, $J = 8.0, 1.5$ Hz, 1H), 7.82 (td, $J = 7.8, 1.5$ Hz, 1H), 7.64 – 7.54 (m, 1H), 7.47 (ddd, $J = 8.5, 7.1, 1.5$ Hz, 1H), 7.41 – 7.28 (m, 2H). ^{13}C NMR (101 MHz, Chloroform-*d*) δ 161.1, 151.2, 134.8, 134.7, 130.5, 130.4, 128.8, 124.5, 122.7, 121.6, 121.2, 118.0, 117.7.



9-methyl-6H-benzo[c]chromen-6-one⁷

39.1 mg, 62% yield. White solid, ^1H NMR (400 MHz, Chloroform-*d*) δ 8.28 (d, $J = 8.1$ Hz, 1H), 8.05 (dd, $J = 8.0, 1.5$ Hz, 1H), 7.90 (s, 1H), 7.47 (ddd, $J = 8.5, 7.2, 1.5$ Hz, 1H), 7.41 – 7.29 (m, 3H), 2.56 (s, 3H). ^{13}C NMR (101 MHz, Chloroform-*d*) δ 161.3, 151.4, 145.9, 134.7, 130.6, 130.3, 130.2, 124.4, 122.7, 121.8, 118.8, 118.1, 117.8, 22.3.

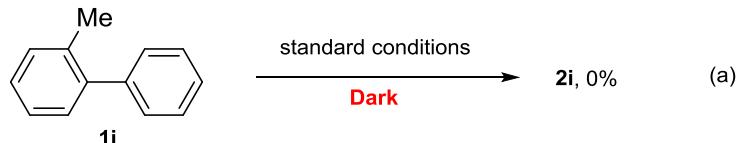


2-(trifluoromethyl)-6H-benzo[c]chromen-6-one⁹

51.9 mg, 66% yield. White solid, ^1H NMR (500 MHz, Chloroform-*d*) δ 8.43 (dd, $J = 7.9, 1.4$ Hz, 1H), 8.32 (d, $J = 2.0$ Hz, 1H), 8.17 (d, $J = 8.0$ Hz, 1H), 7.90 (td, $J = 7.8,$

1.5 Hz, 1H), 7.73 (dd, J = 8.7, 2.1 Hz, 1H), 7.70 – 7.62 (m, 1H), 7.48 (d, J = 8.6 Hz, 1H). ^{13}C NMR (126 MHz, Chloroform-*d*) δ 160.2, 153.1, 135.3, 133.5, 130.8, 129.9, 127.0 (q, J = 3.5 Hz), 126.9 (q, J = 32.8 Hz), 123.8 (q, J = 270.5 Hz), 121.9, 121.3, 120.4 (q, J = 3.8 Hz), 118.6, 118.4. ^{19}F NMR (377 MHz, Chloroform-*d*) δ -62.0.

3. Control experiments (Scheme 4)



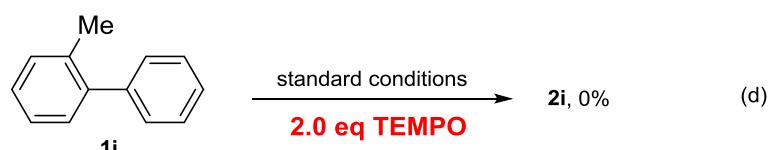
Reaction (a): To an oven-dried 10 mL Schlenk tube equipped with a stir bar was charged with substrate **1i** (50.4 mg, 0.3 mol) and $\text{Acr}^+\text{MesClO}_4^-$ (5.0 mg, 0.012 mmol). The tube was closed with a septum, evacuated, and back-filled with oxygen (balloon). To this mixture was added 1.0 mL solvent ($\text{MeCN}/\text{H}_2\text{O}$ = 2:1) and HCl (150 μL , 0.6 mmol, 4M in 1,3-dioxane). The reaction mixture was stirred at room temperature under dark for 3 days. No benzocoumarin **2i** was detected by TLC.



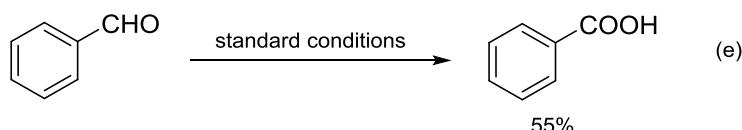
Reaction (b): To an oven-dried 10 mL Schlenk tube equipped with a stir bar was charged with substrate **1i** (50.4 mg, 0.3 mol). The tube was closed with a septum, evacuated, and back-filled with oxygen (balloon). To this mixture was added 1.0 mL solvent ($\text{MeCN}/\text{H}_2\text{O}$ = 2:1) and HCl (150 μL , 0.6 mmol, 4M in 1,3-dioxane). The reaction mixture was stirred at room temperature under blue LEDs for 3 days. No benzocoumarin **2i** was detected by TLC.



Reaction (c): To an oven-dried 10 mL Schlenk tube equipped with a stir bar was charged with substrate **1i** (50.4 mg, 0.3 mol) and $\text{Acr}^+\text{MesClO}_4^-$ (5.0 mg, 0.012 mmol). The tube was closed with a septum, evacuated, and back-filled with oxygen (balloon). To this mixture was added 1.0 mL solvent ($\text{MeCN}/\text{H}_2\text{O} = 2:1$). The reaction mixture was stirred at room temperature under blue LEDs until the full consumption of the starting materials. The reaction mixture was concentrated under reduced pressure, and the residue was purified by column chromatography on silica gel (petroleum ether/EtOAc = 70:1 - 20:1) to furnish the benzocoumarins **2i** (8.2 mg, 14% yield).

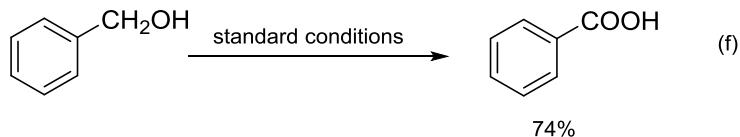


Reaction (d): To an oven-dried 10 mL Schlenk tube equipped with a stir bar was charged with substrate **1i** (50.4 mg, 0.3 mol), $\text{Acr}^+\text{MesClO}_4^-$ (5.0 mg, 0.012 mmol) and TEMPO (93.6 mg, 0.6 mmol). The tube was closed with a septum, evacuated, and back-filled with oxygen (balloon). To this mixture was added 1.0 mL solvent ($\text{MeCN}/\text{H}_2\text{O} = 2:1$) and HCl (150 μm , 0.6 mmol, 4M in 1,3-dioxane). The reaction mixture was stirred at room temperature under blue LEDs for 2 days. No benzocoumarin **2i** was detected by TLC.



Reaction (e): To an oven-dried 10 mL Schlenk tube equipped with a stir bar was charged with benzaldehyde (31.8 mg, 0.3 mol) and $\text{Acr}^+\text{MesClO}_4^-$ (5.0 mg, 0.012 mmol). The tube was closed with a septum, evacuated, and back-filled with oxygen (balloon). To this mixture was added 1.0 mL solvent ($\text{MeCN}/\text{H}_2\text{O} = 2:1$) and HCl (150 μm , 0.6 mmol, 4M in 1,3-dioxane). The reaction mixture was stirred at room temperature under blue LEDs for 2 days. The reaction mixture

was concentrated under reduced pressure, and the residue was purified by column chromatography on silica gel (petroleum ether/EtOAc = 10:1 - 5:1) to furnish the benzoic acid (20.0 mg, 55% yield).



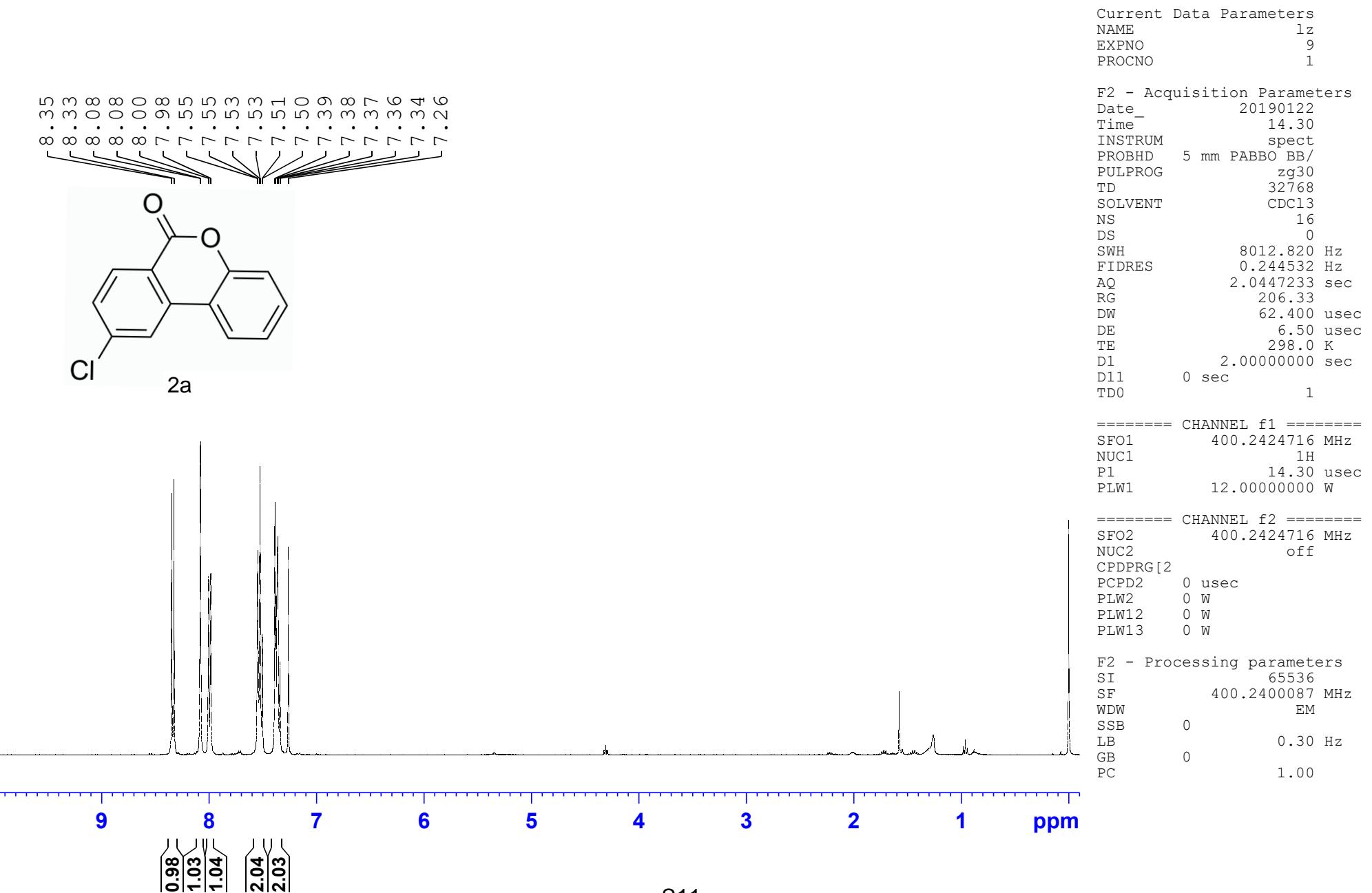
Reaction (f): To an oven-dried 10 mL Schlenk tube equipped with a stir bar was charged with benzyl alcohol (32.4 mg, 0.3 mol) and $\text{Acr}^+\text{MesClO}_4^-$ (5.0 mg, 0.012 mmol). The tube was closed with a septum, evacuated, and back-filled with oxygen (balloon). To this mixture was added 1.0 mL solvent ($\text{MeCN}/\text{H}_2\text{O} = 2:1$) and HCl (150 μm , 0.6 mmol, 4M in 1,3-dioxane). The reaction mixture was stirred at room temperature under blue LEDs for 2 days. The reaction mixture was concentrated under reduced pressure, and the residue was purified by column chromatography on silica gel (petroleum ether/EtOAc = 10:1 - 5:1) to furnish the benzoic acid (27.2 mg, 74% yield).

4. References

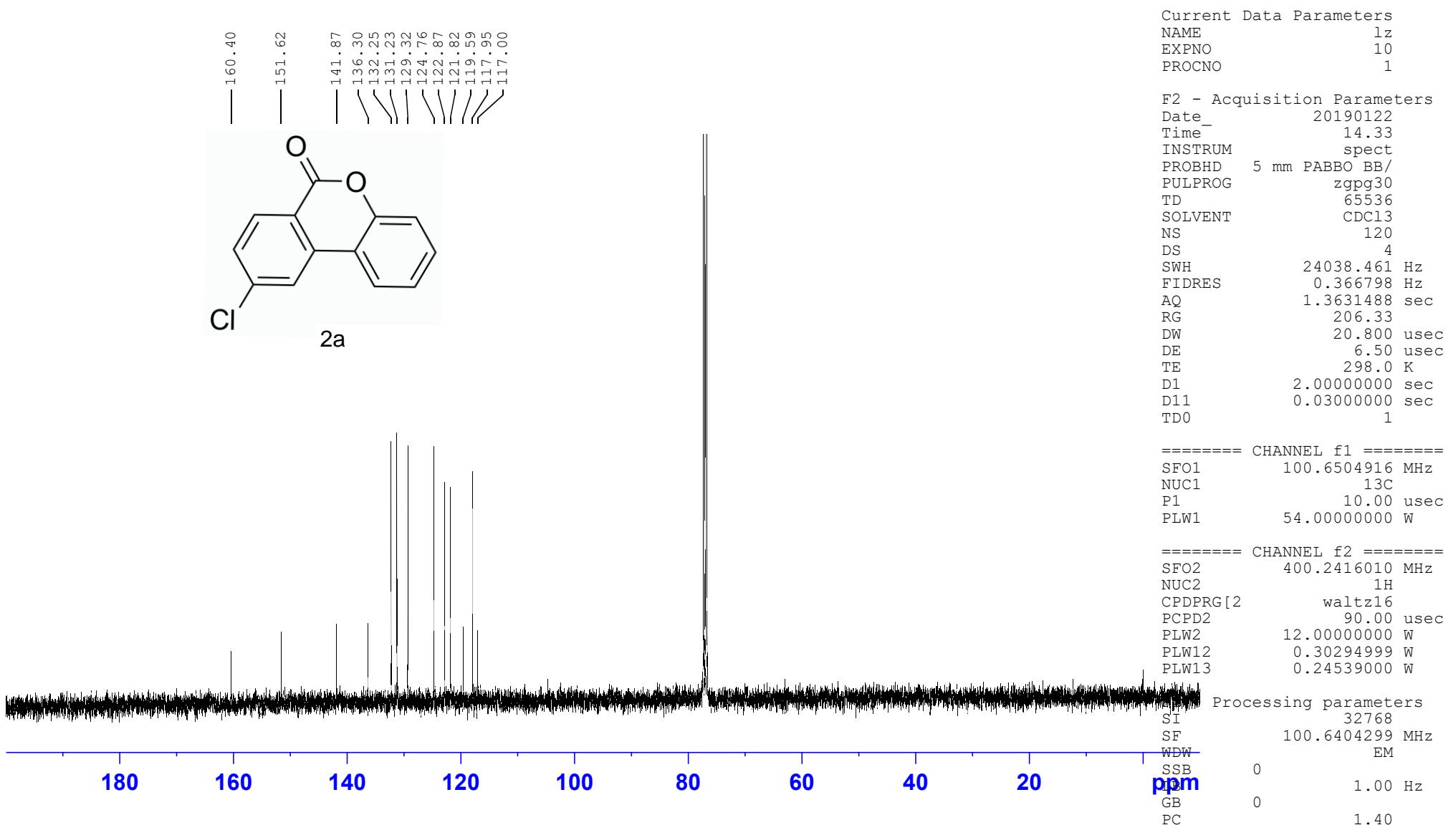
- 1 J. K. Laha, K. P. Jethava, S. Patel and K. V. Patel, *J. Org. Chem.*, 2017, **82**, 76.
- 2 M. Saifuddin, P. K. Agarwal and B. Kundu, *J. Org. Chem.*, 2011, **76**, 10122.
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- 5 J. Zhang, D. Shi, H. Zhang, Z. Xu, H. Bao, H. Jin and Y. Liu, *Tetrahedron*, 2017, **73**, 154.

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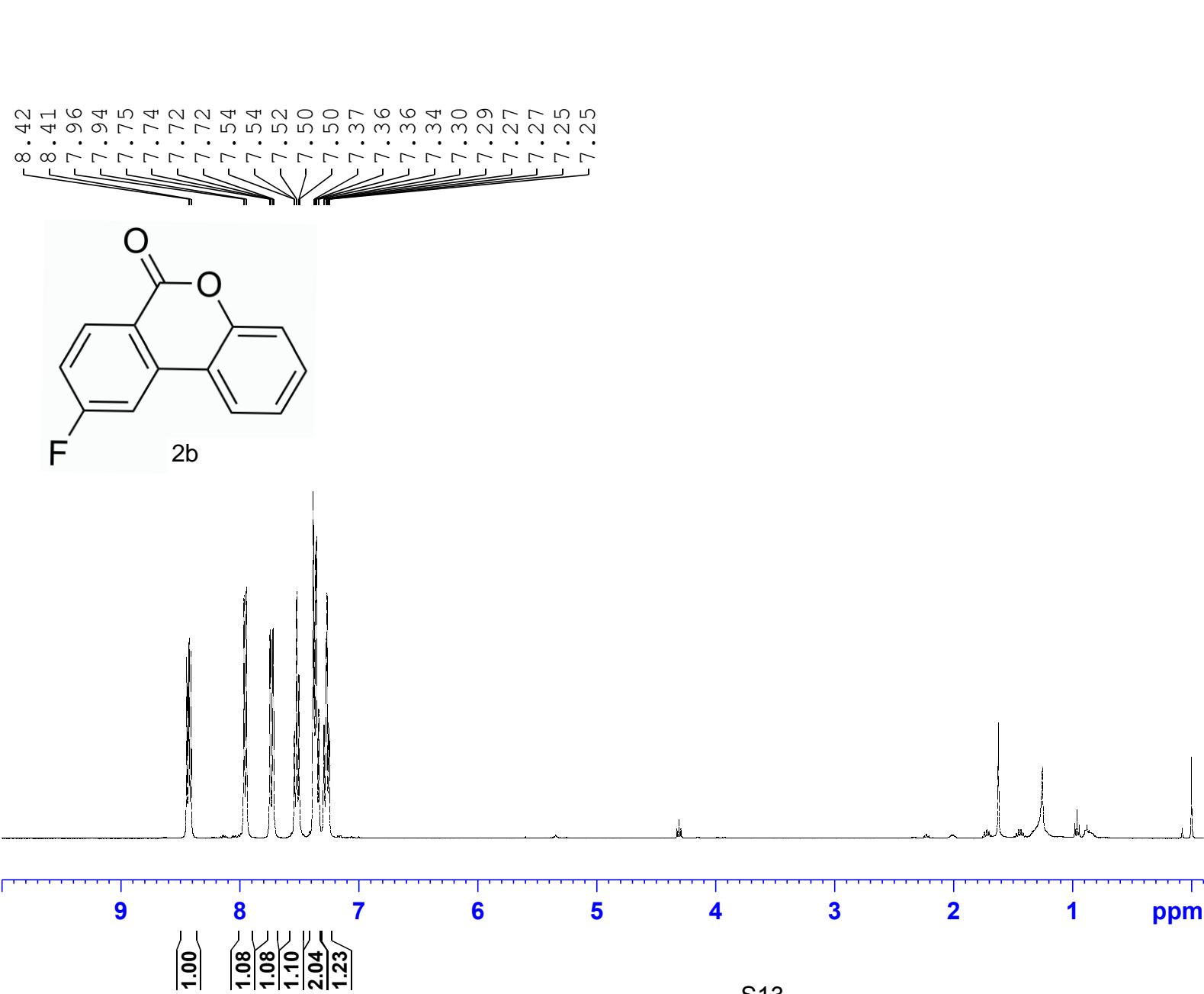
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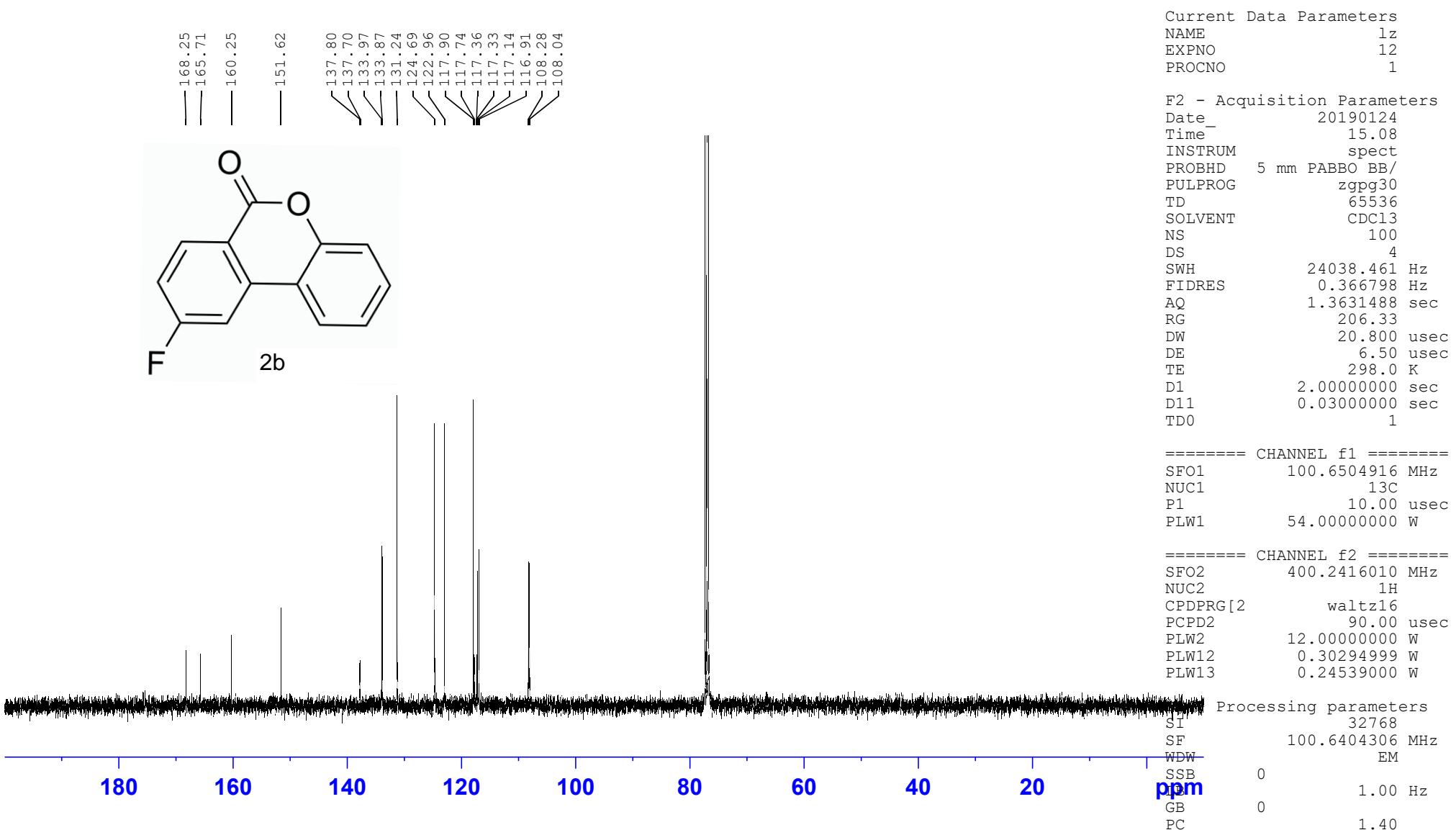
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lz-1065C

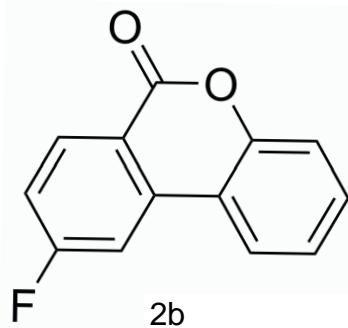


lz-1065C



1065C

-101.08



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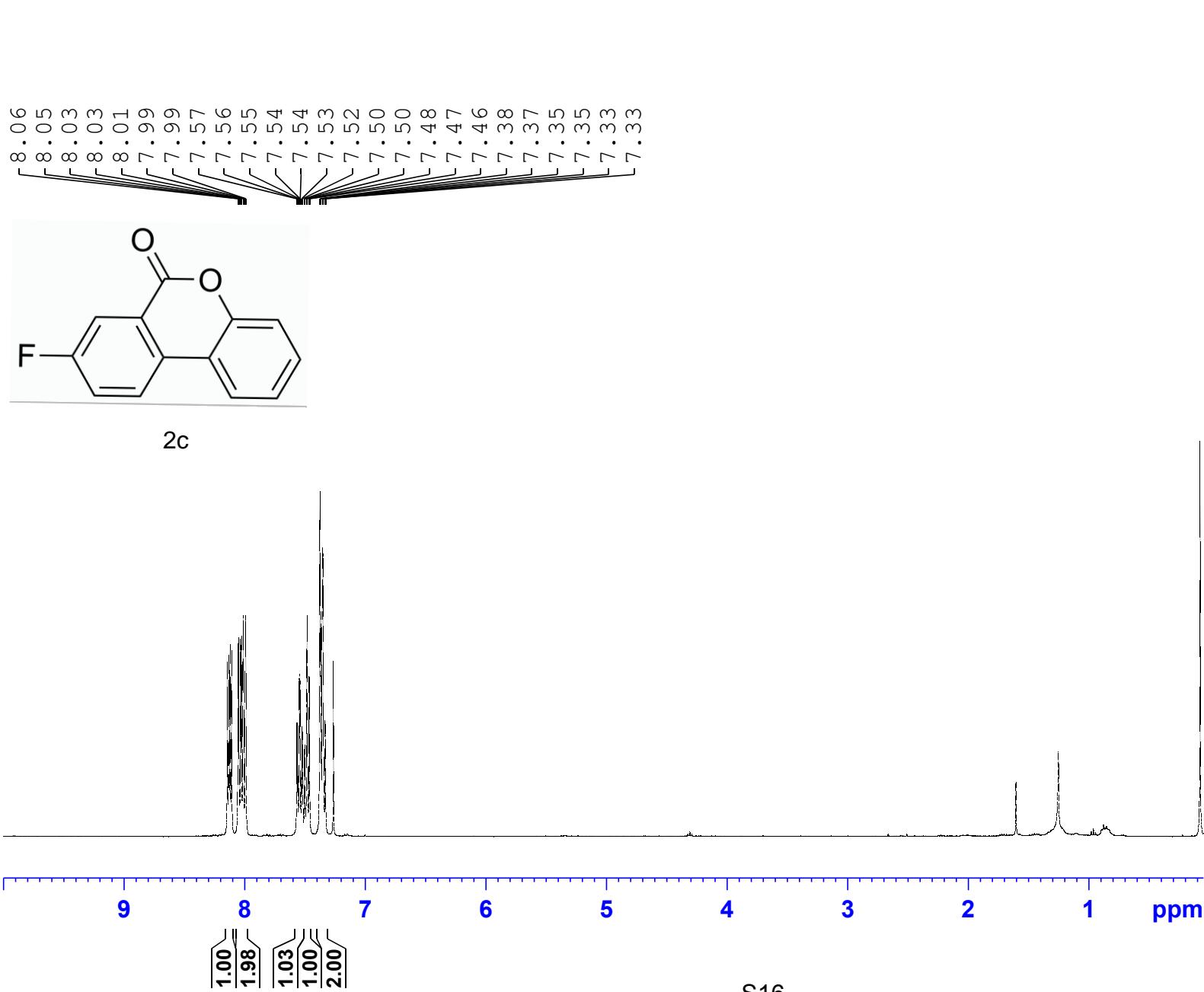
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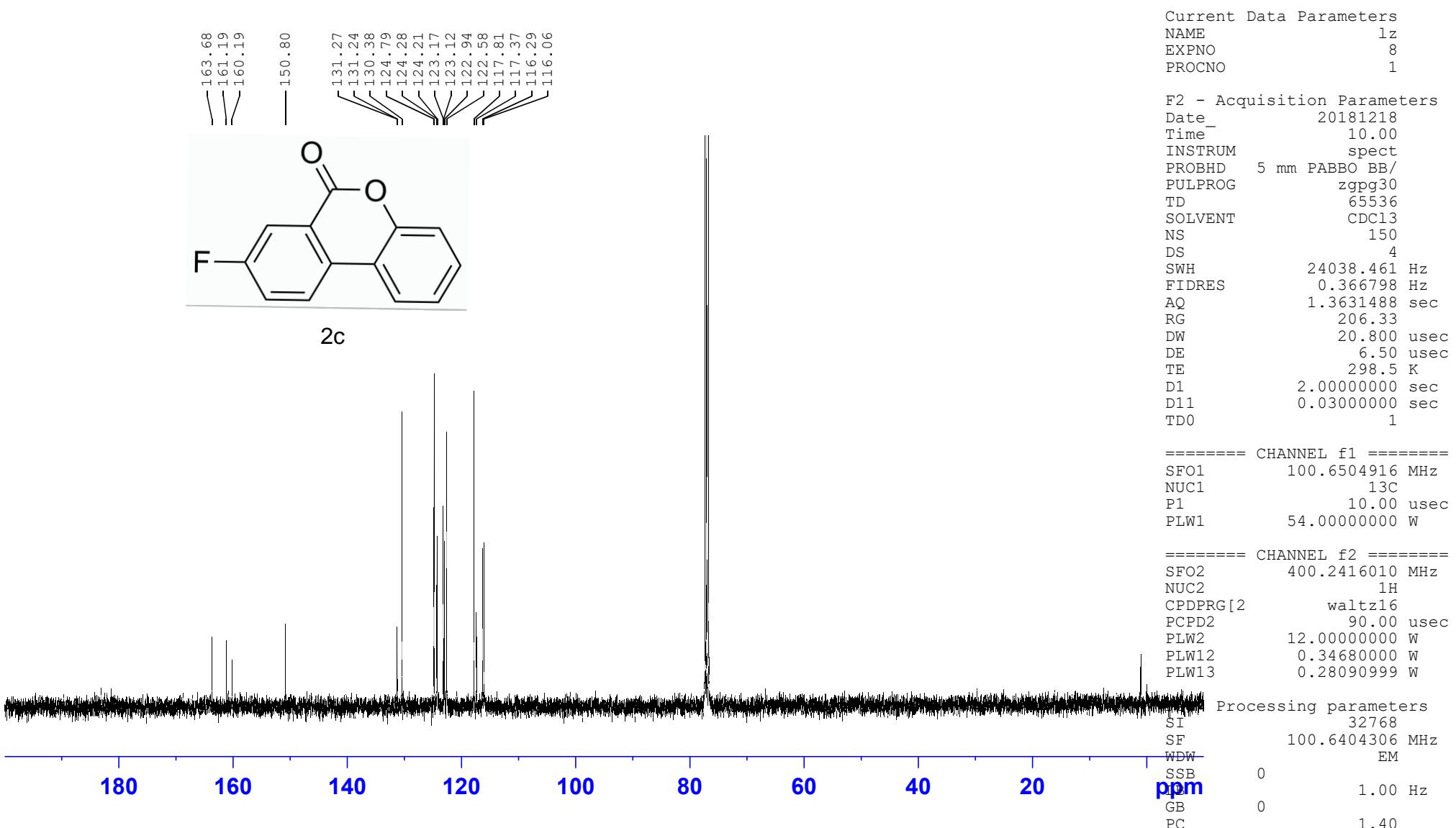
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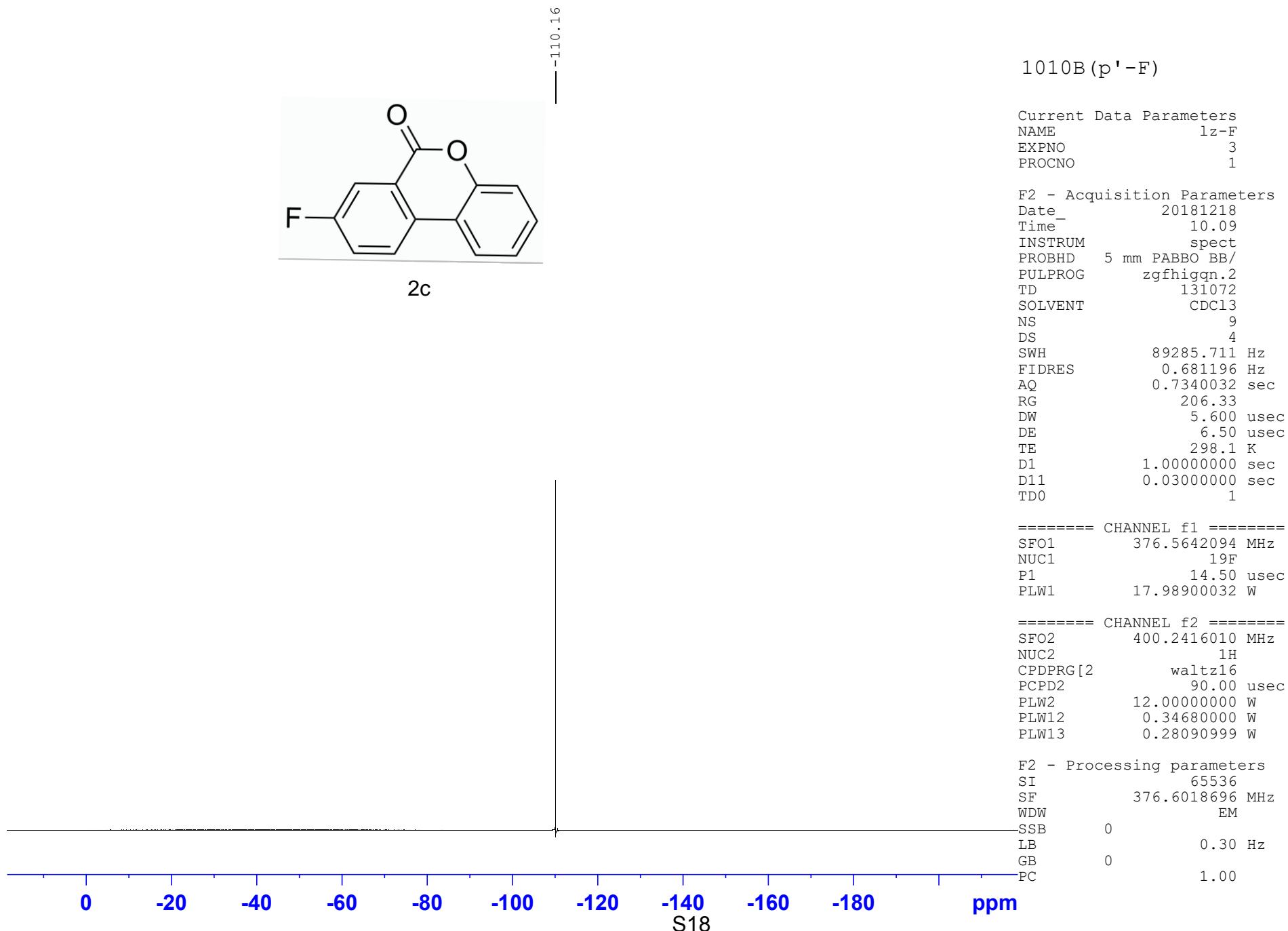
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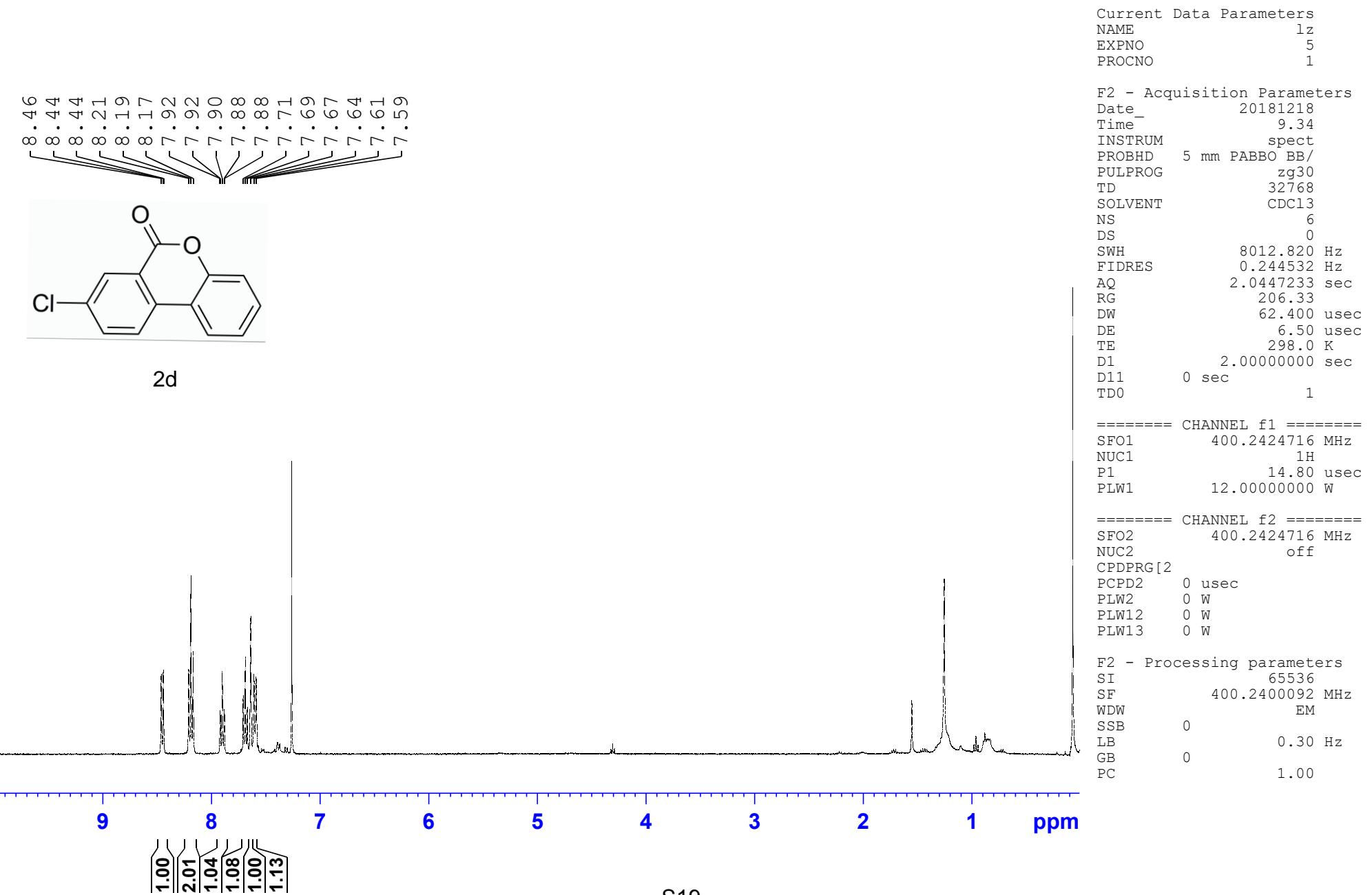
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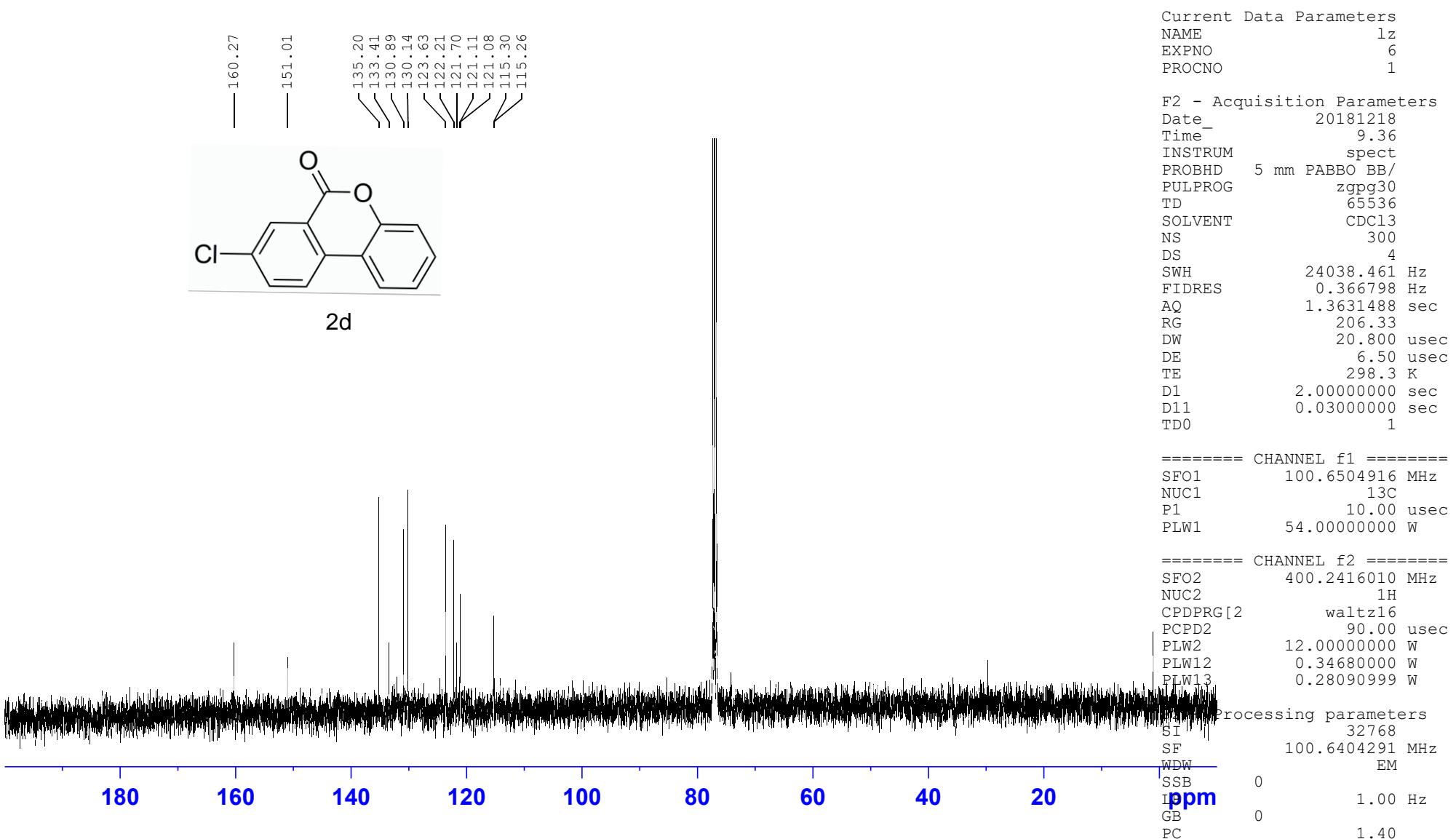
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1010C(p'-Br)



lz-1066A

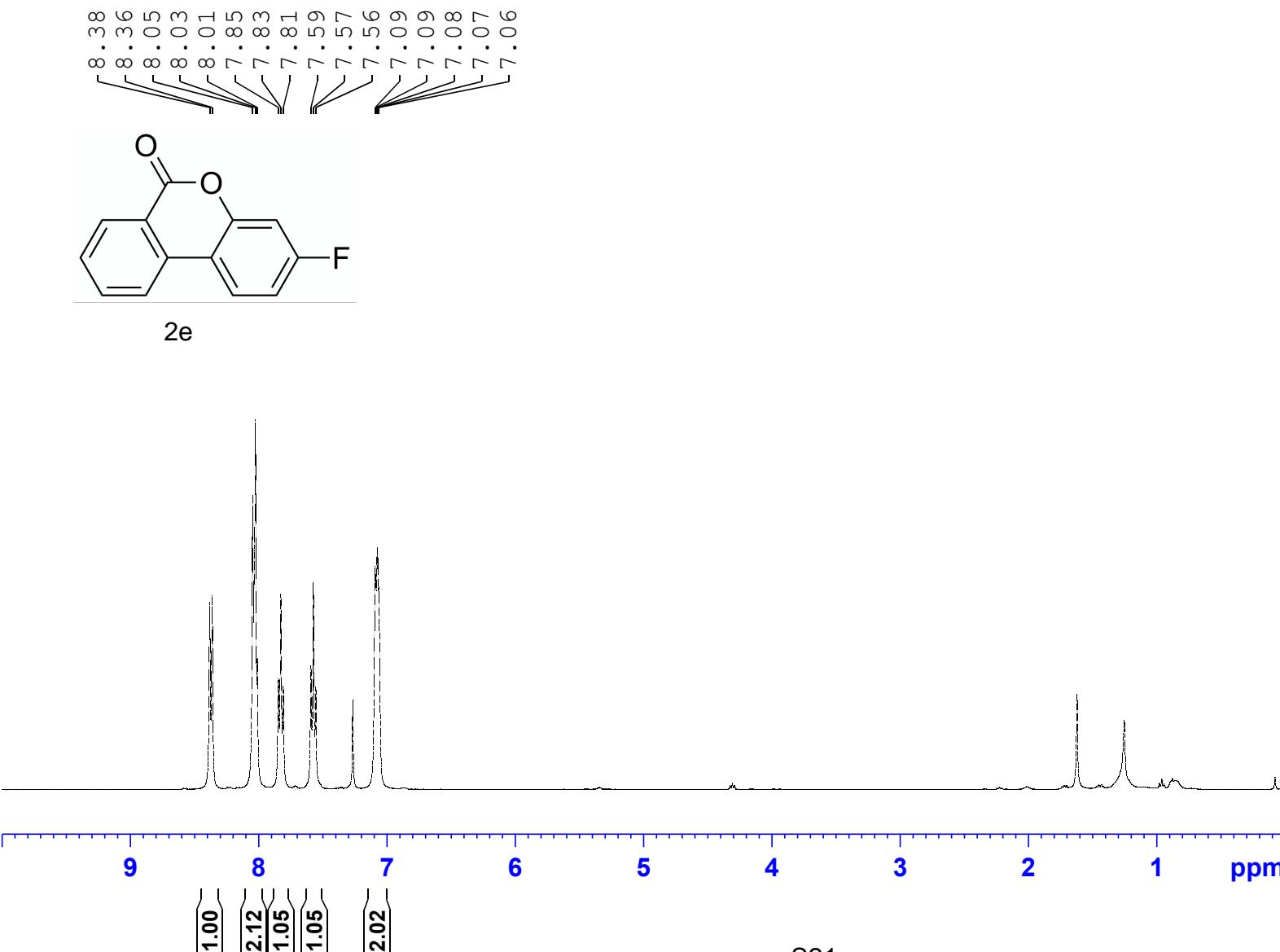
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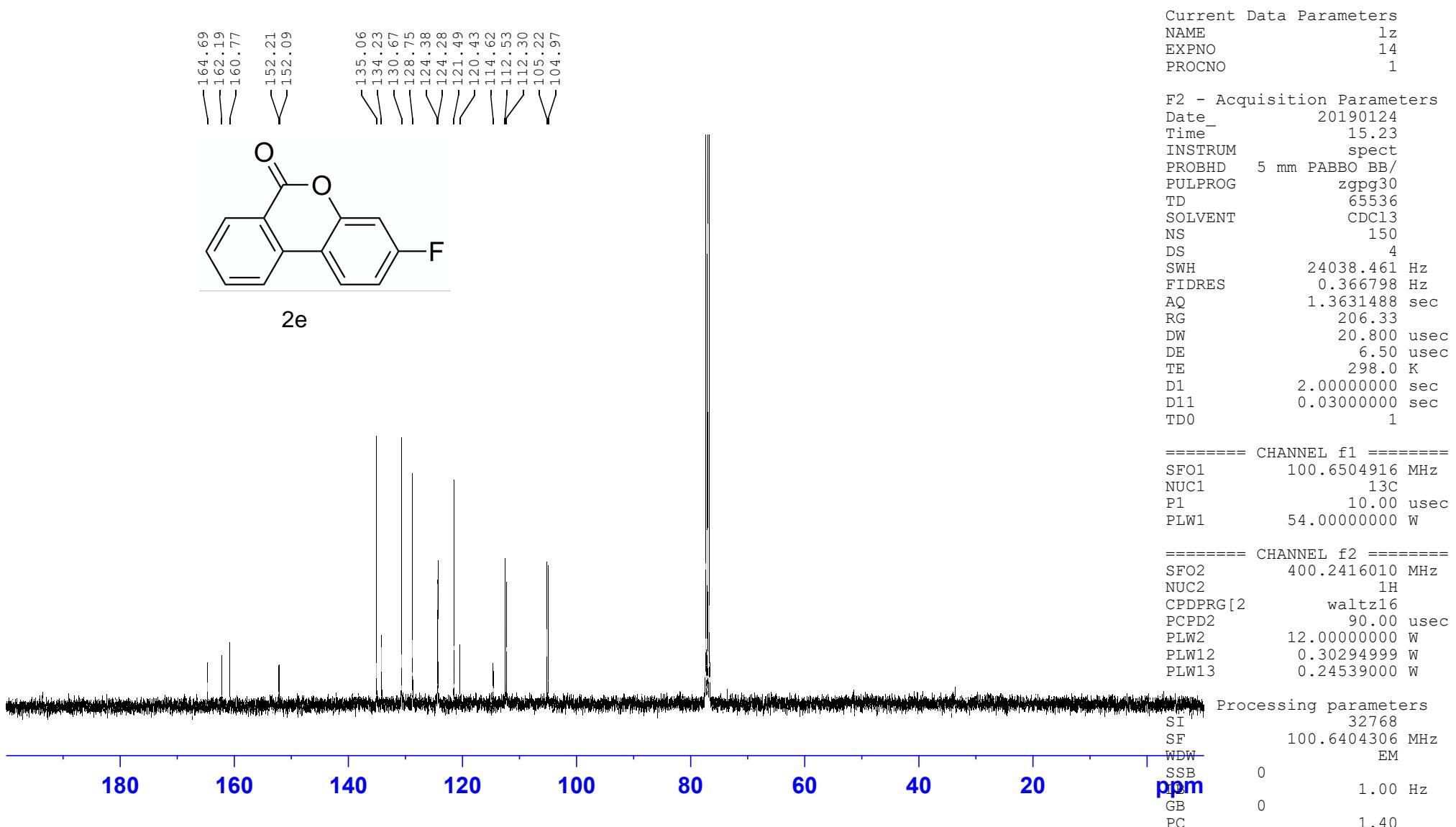
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PLW1 12.00000000 W

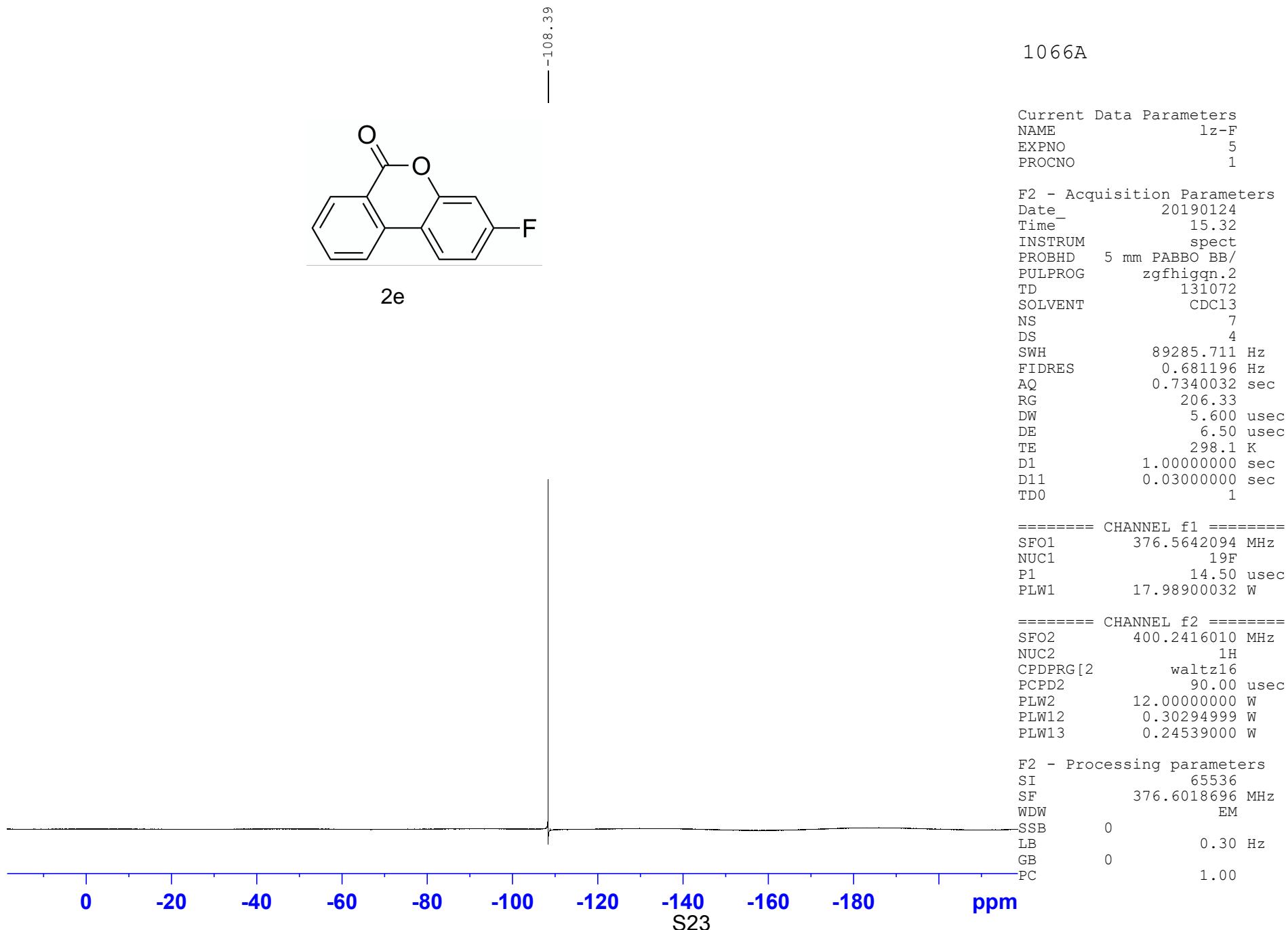
===== CHANNEL f2 =====
SFO2 400.2424716 MHz
NUC2 off
CPDPRG[2]
PCPD2 0 usec
PLW2 0 W
PLW12 0 W
PLW13 0 W

F2 - Processing parameters
SI 65536
SF 400.2400071 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

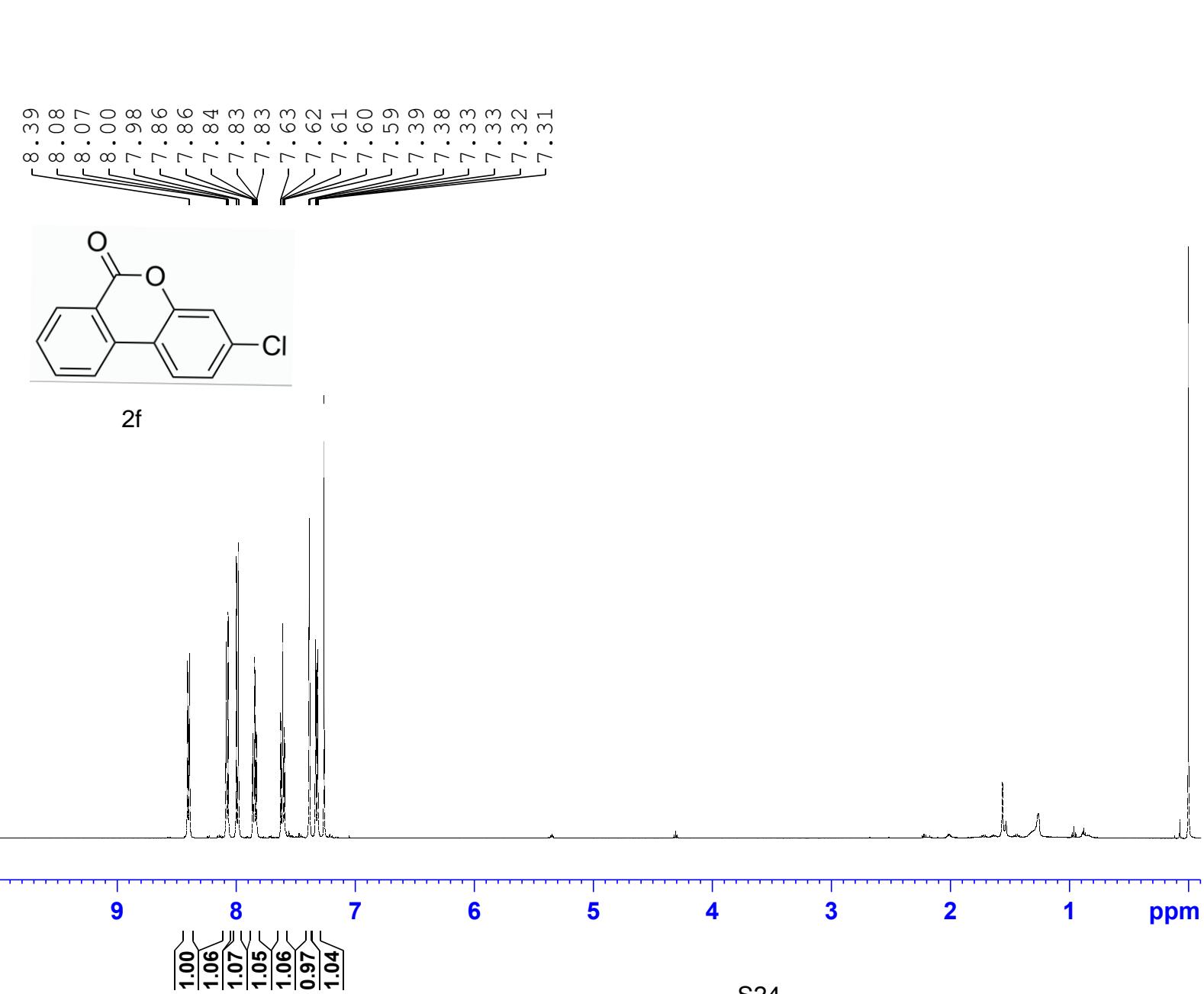


lz-1066A

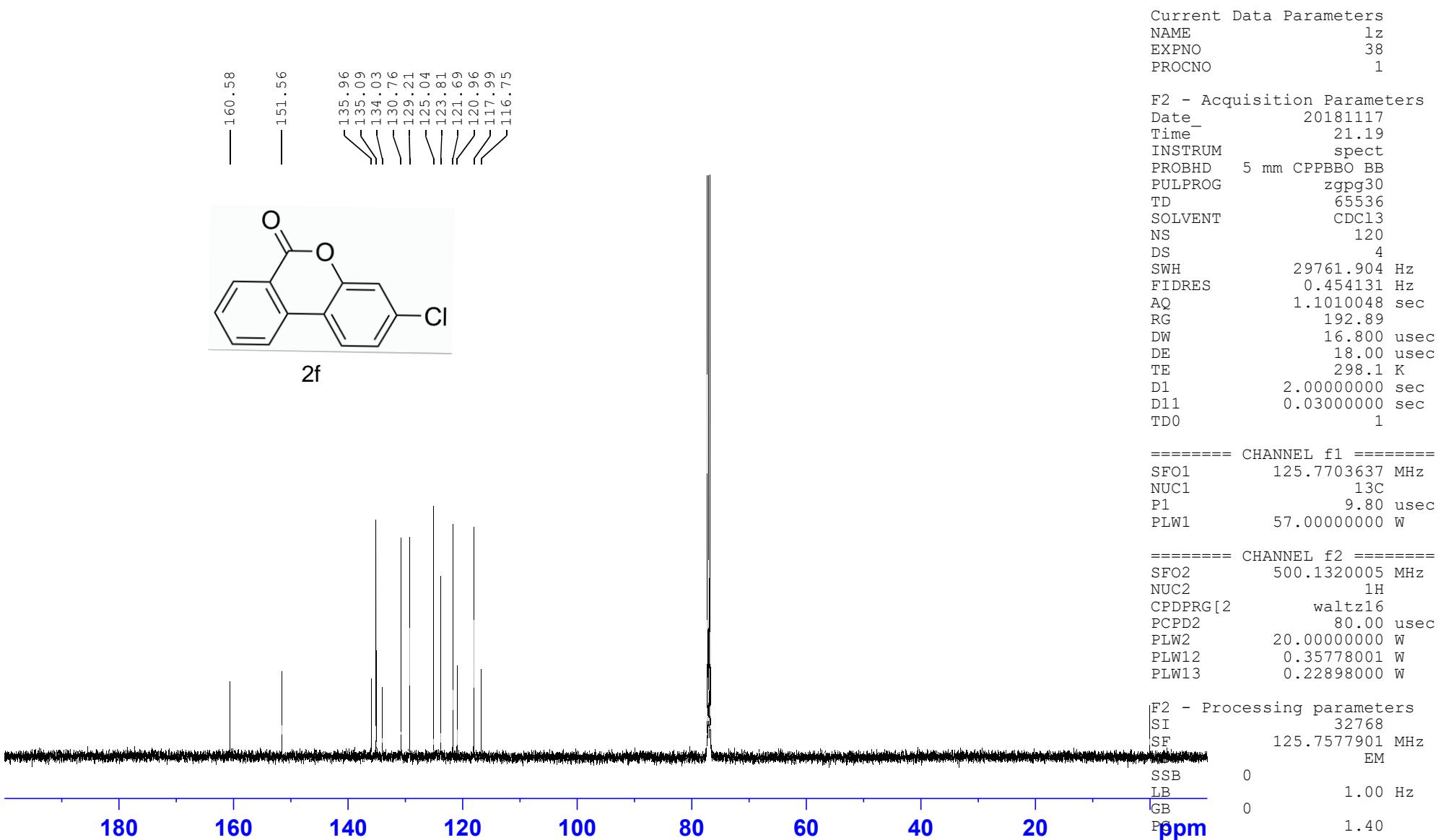




lz-976A

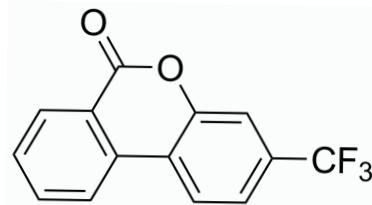


lz-976A

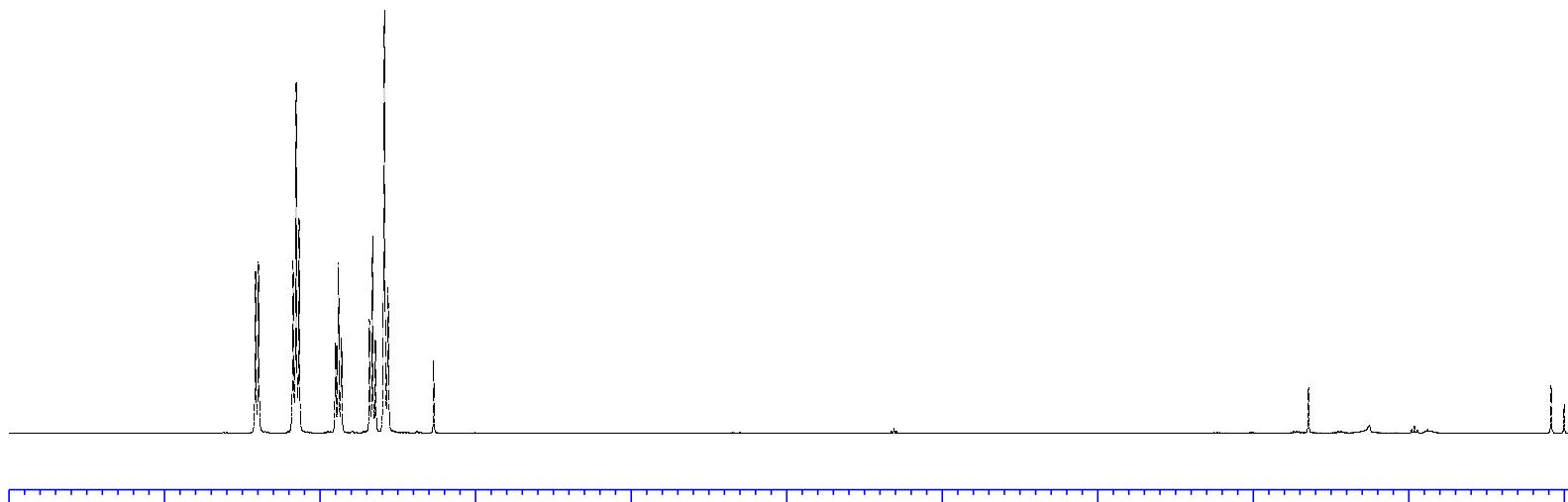


lz-p-*CF*3

4.2 .41
3.9 .17
1.5 .13
1.3 .90
1.1 .90
8.8 .86
7.8 .86
7.6 .68
7.6 .68
7.4 .64
7.3 .64
7.2 .59
7.1 .56



2g



1.00
2.13
1.05
1.05
2.04

Current Data Parameters
NAME lz
EXPNO 18
PROCNO 1

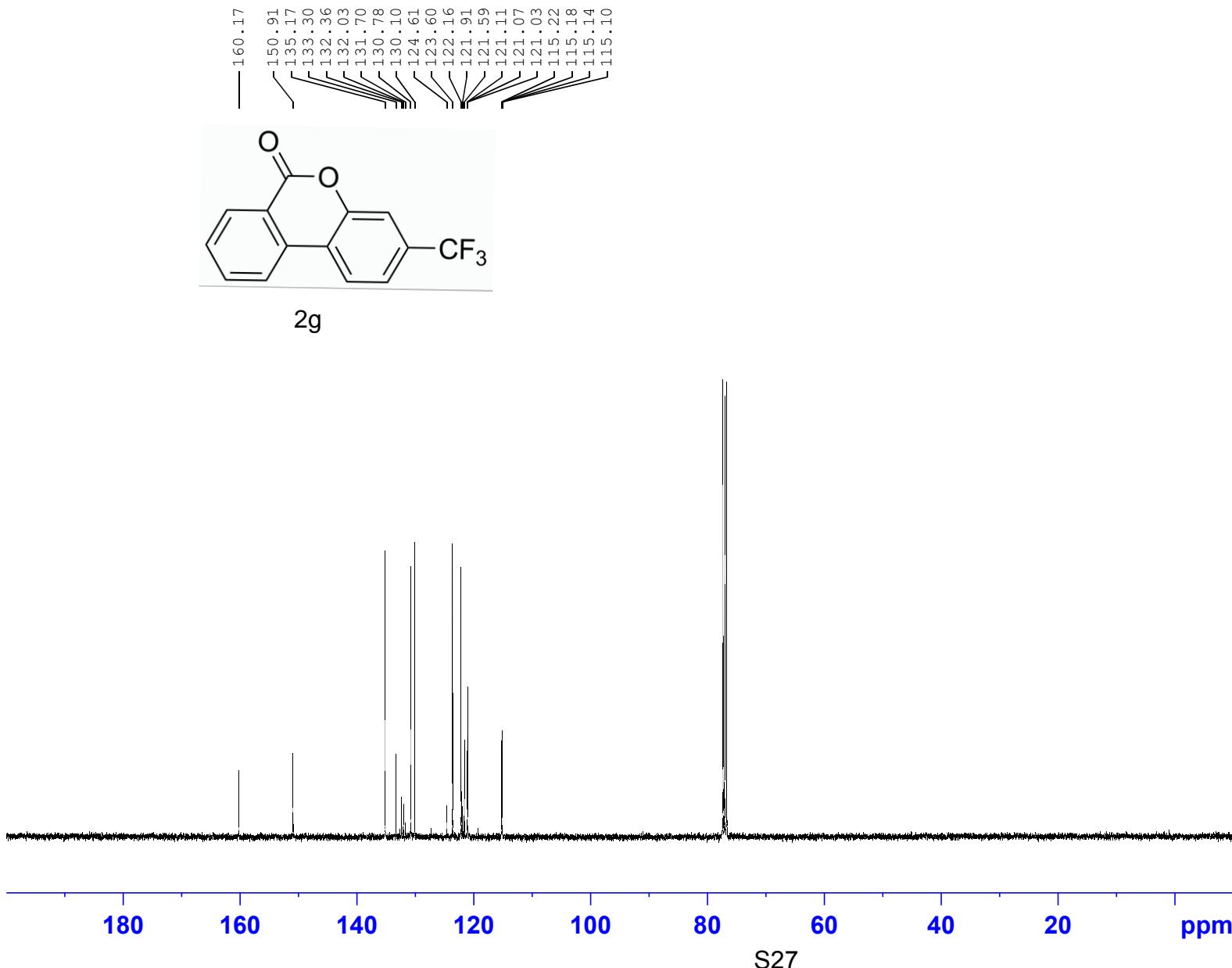
F2 - Acquisition Parameters
Date_ 20190227
Time 14.33
INSTRUM spect
PROBHD 5 mm PABBO BB/
PULPROG zg30
TD 32768
SOLVENT CDCl3
NS 6
DS 0
SWH 8012.820 Hz
FIDRES 0.244532 Hz
AQ 2.0447233 sec
RG 206.33
DW 62.400 usec
DE 6.50 usec
TE 297.8 K
D1 2.00000000 sec
D11 0 sec
TD0 1

===== CHANNEL f1 =====
SFO1 400.2424716 MHz
NUC1 1H
P1 14.30 usec
PLW1 12.00000000 W

===== CHANNEL f2 =====
SFO2 400.2424716 MHz
NUC2 off
CPDPRG[2
PCPD2 0 usec
PLW2 0 W
PLW12 0 W
PLW13 0 W

F2 - Processing parameters
SI 65536
SF 400.2400062 MHz
WDW EM
SSB 0 0.30 Hz
LB 0 1.00
GB PC

lz-p-CF₃



Current Data Parameters
NAME lz
EXPNO 17
PROCNO 1

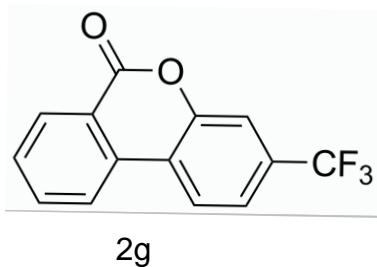
F2 - Acquisition Parameters
Date 20190227
Time 14.20
INSTRUM spect
PROBHD 5 mm PABBO BB/
PULPROG zgpg30
TD 65536
SOLVENT CDCl₃
NS 230
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631488 sec
RG 206.33
DW 20.800 usec
DE 6.50 usec
TE 298.2 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
SFO1 100.6504916 MHz
NUC1 ¹³C
P1 10.00 usec
PLW1 54.00000000 W

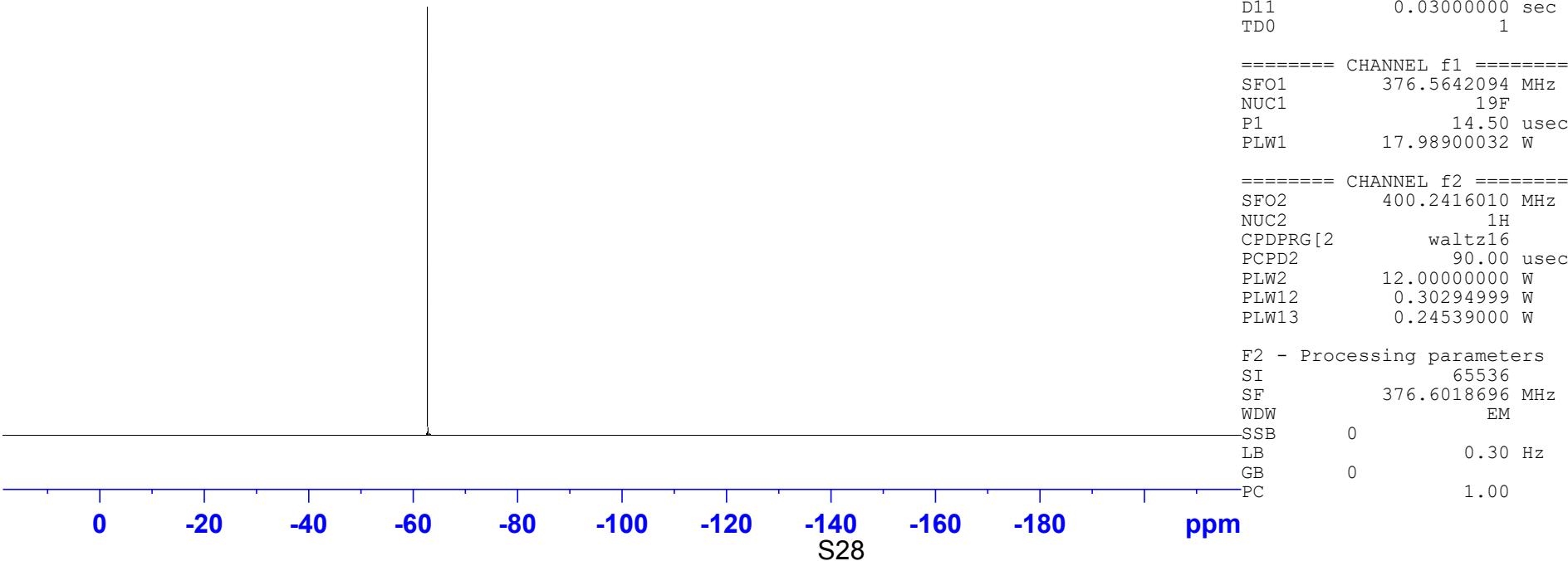
===== CHANNEL f2 =====
SFO2 400.2416010 MHz
NUC2 ¹H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 12.00000000 W
PLW12 0.30294999 W
PLW13 0.24539000 W

F2 - Processing parameters
SI 32768
SF 100.6404306 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

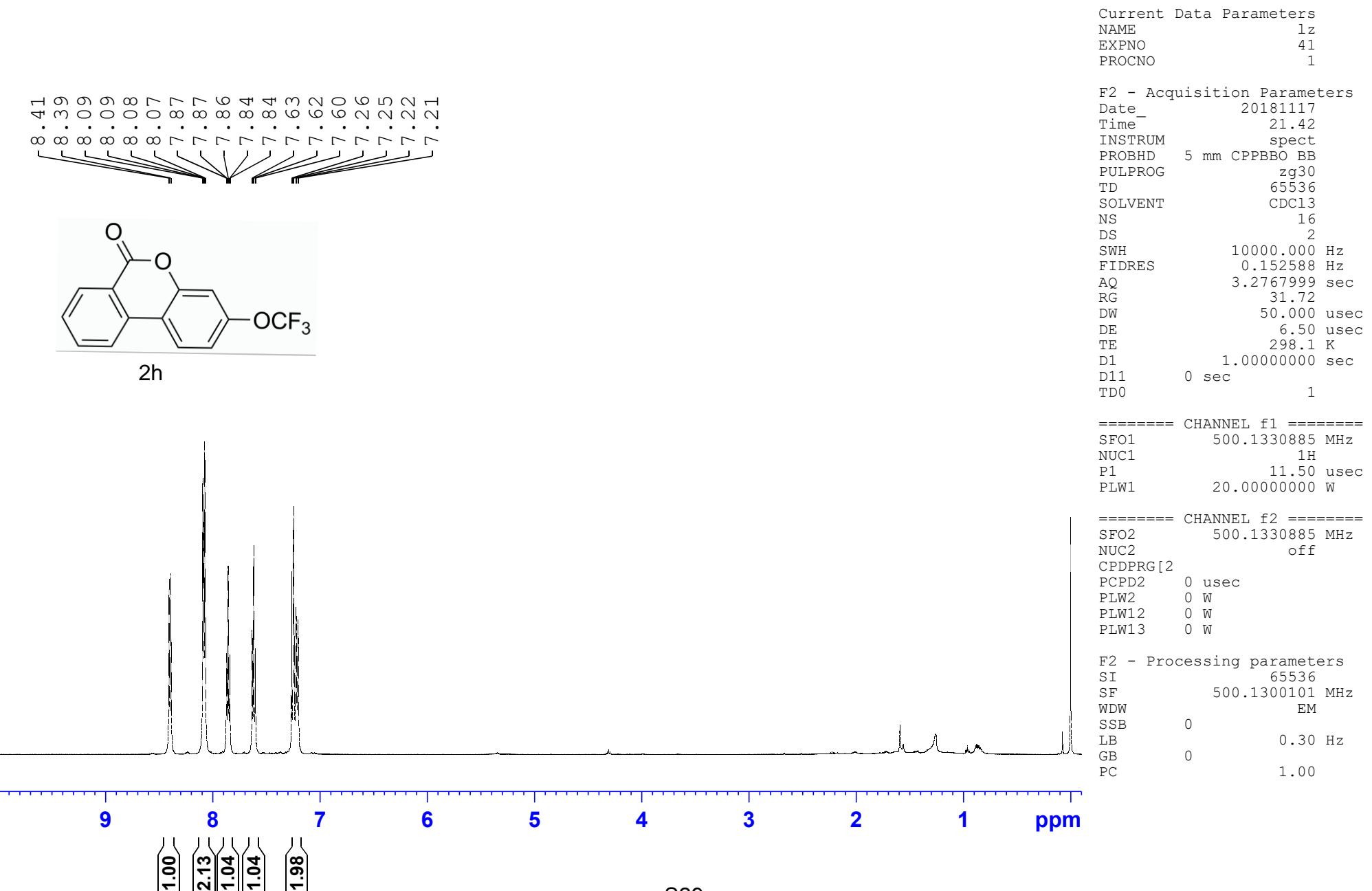
1z-p-cf3



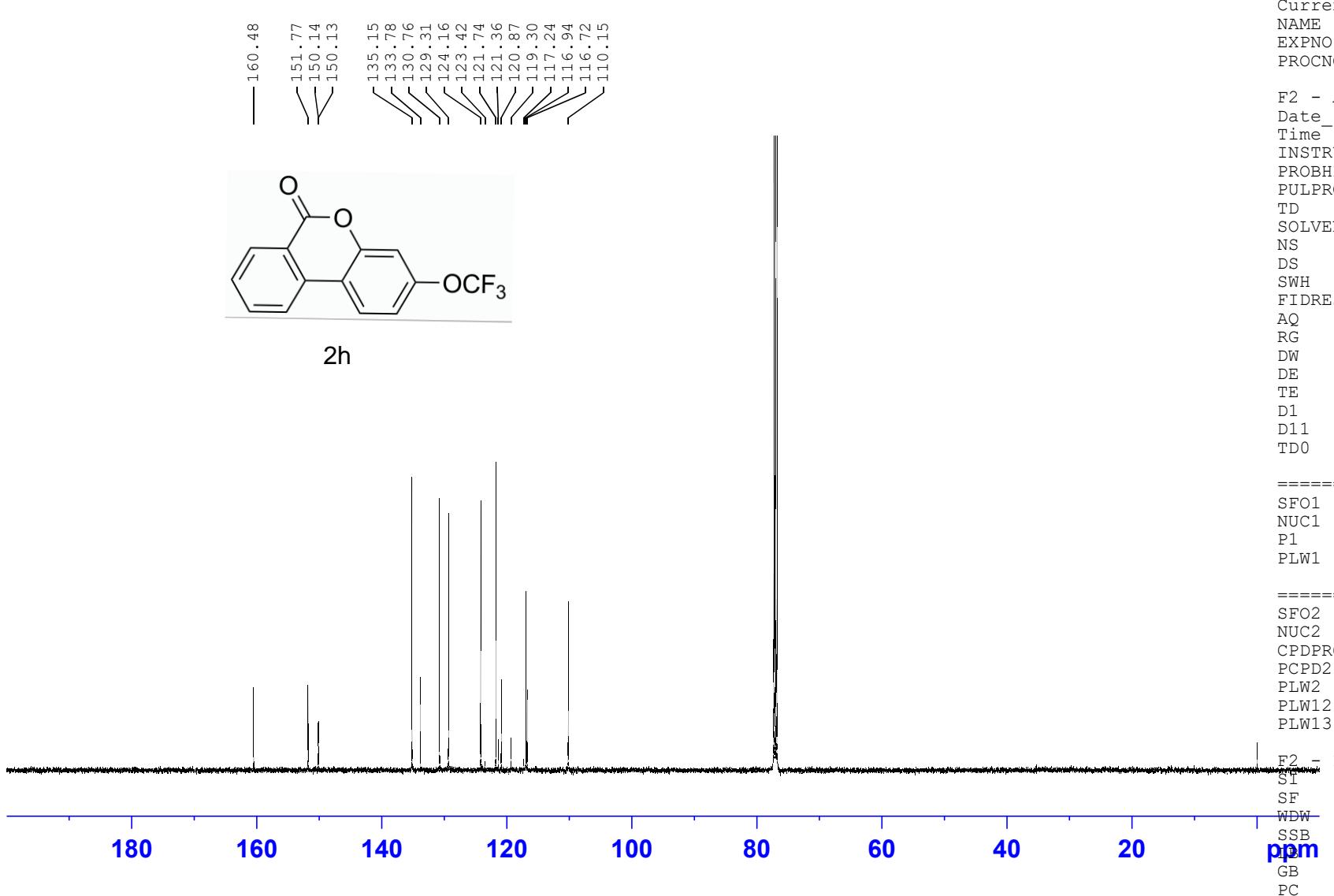
-62.82



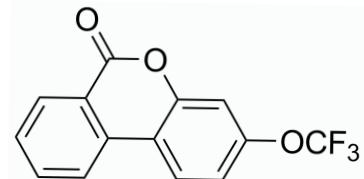
lz-976C



lz-976C

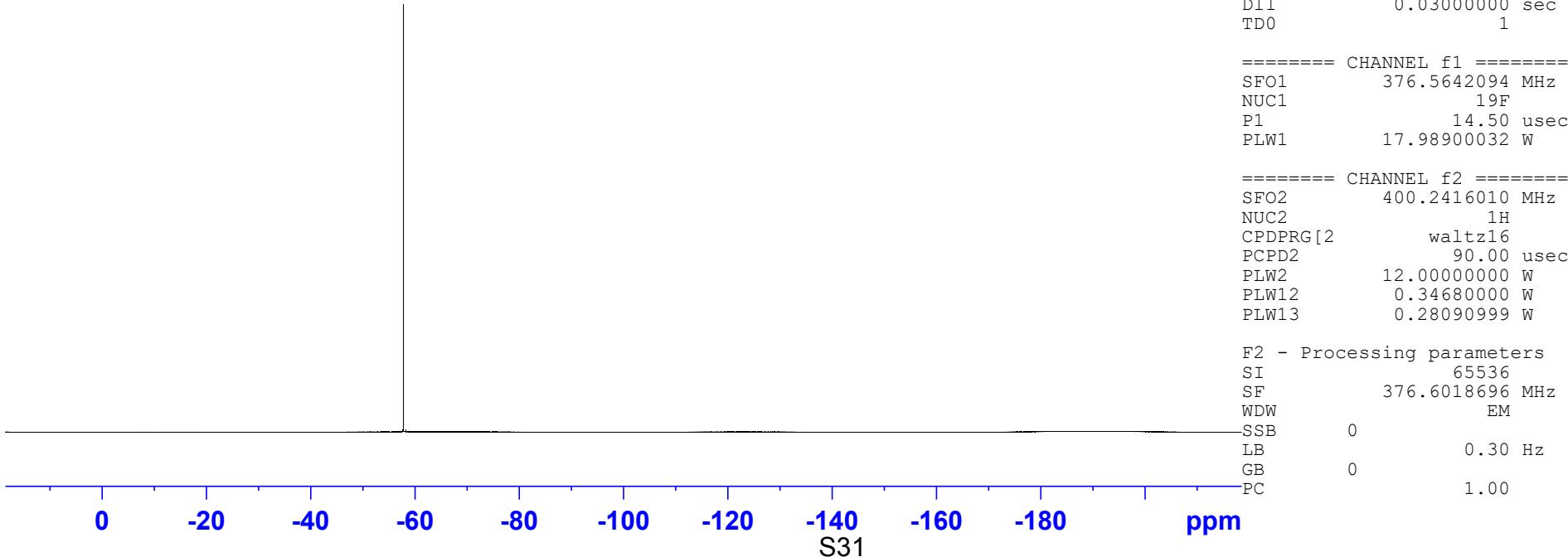


lz-976C



2h

-57.84



Current Data Parameters
NAME lz-F
EXPNO 4
PROCNO 1

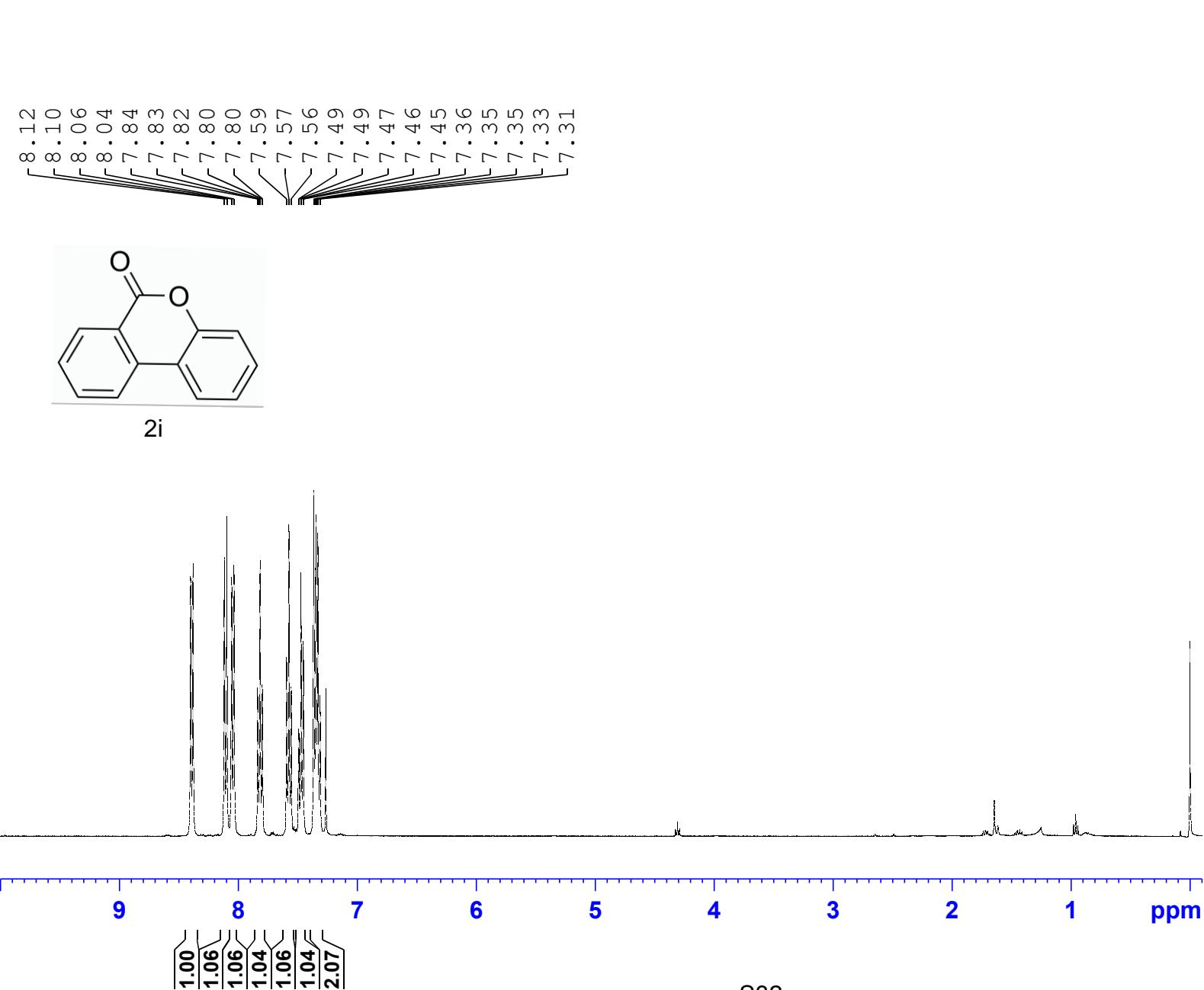
F2 - Acquisition Parameters
Date 20181118
Time 13.45
INSTRUM spect
PROBHD 5 mm PABBO BB/
PULPROG zgfhiggqn.2
TD 131072
SOLVENT CDCl3
NS 7
DS 4
SWH 89285.711 Hz
FIDRES 0.681196 Hz
AQ 0.7340032 sec
RG 206.33
DW 5.600 usec
DE 6.50 usec
TE 298.1 K
D1 1.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
SFO1 376.5642094 MHz
NUC1 19F
P1 14.50 usec
PLW1 17.98900032 W

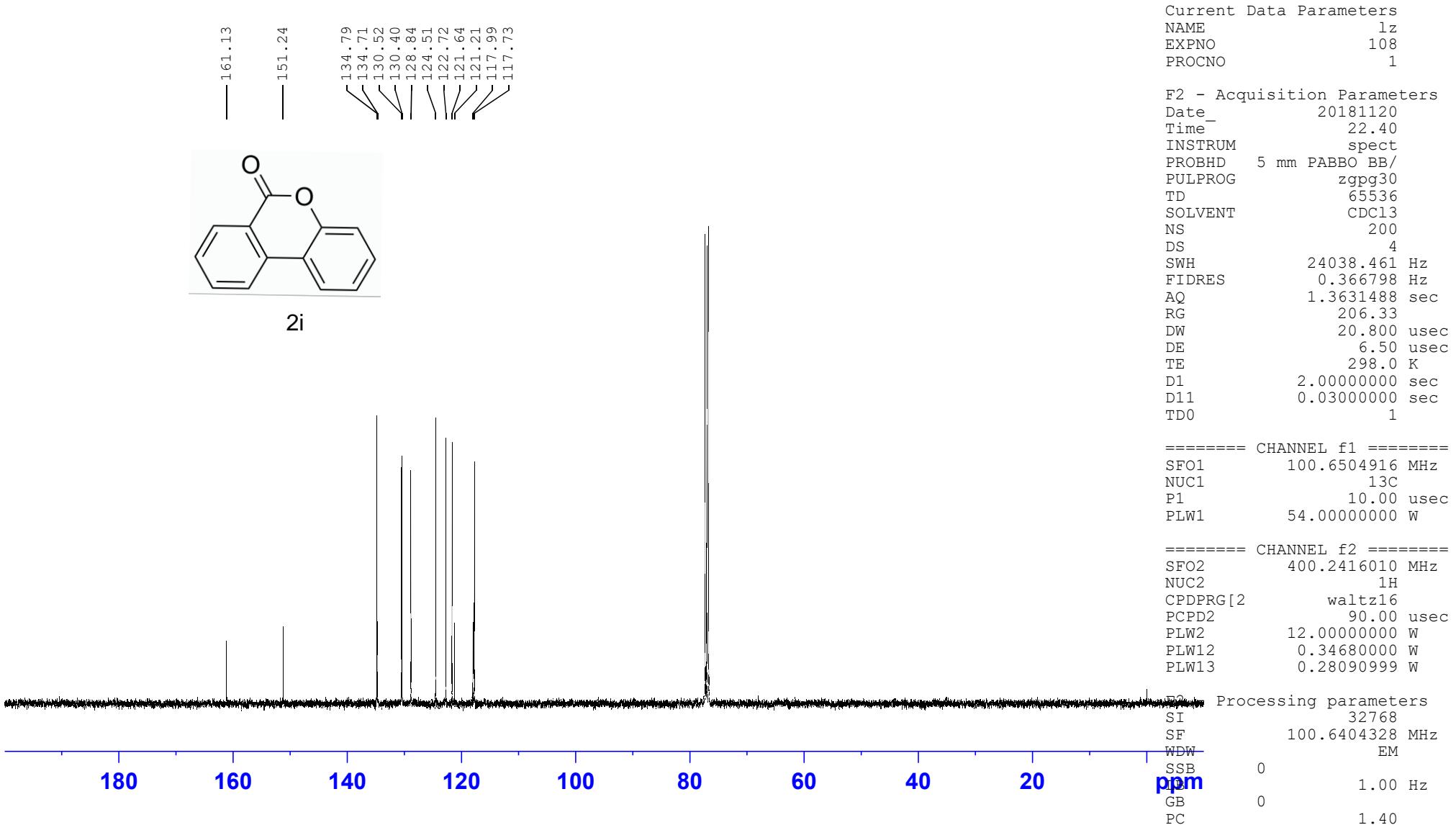
===== CHANNEL f2 =====
SFO2 400.2416010 MHz
NUC2 1H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 12.00000000 W
PLW12 0.34680000 W
PLW13 0.28090999 W

F2 - Processing parameters
SI 65536
SF 376.6018696 MHz
WDW EM
SSB 0
LB 0 0.30 Hz
GB 0
PC 1.00

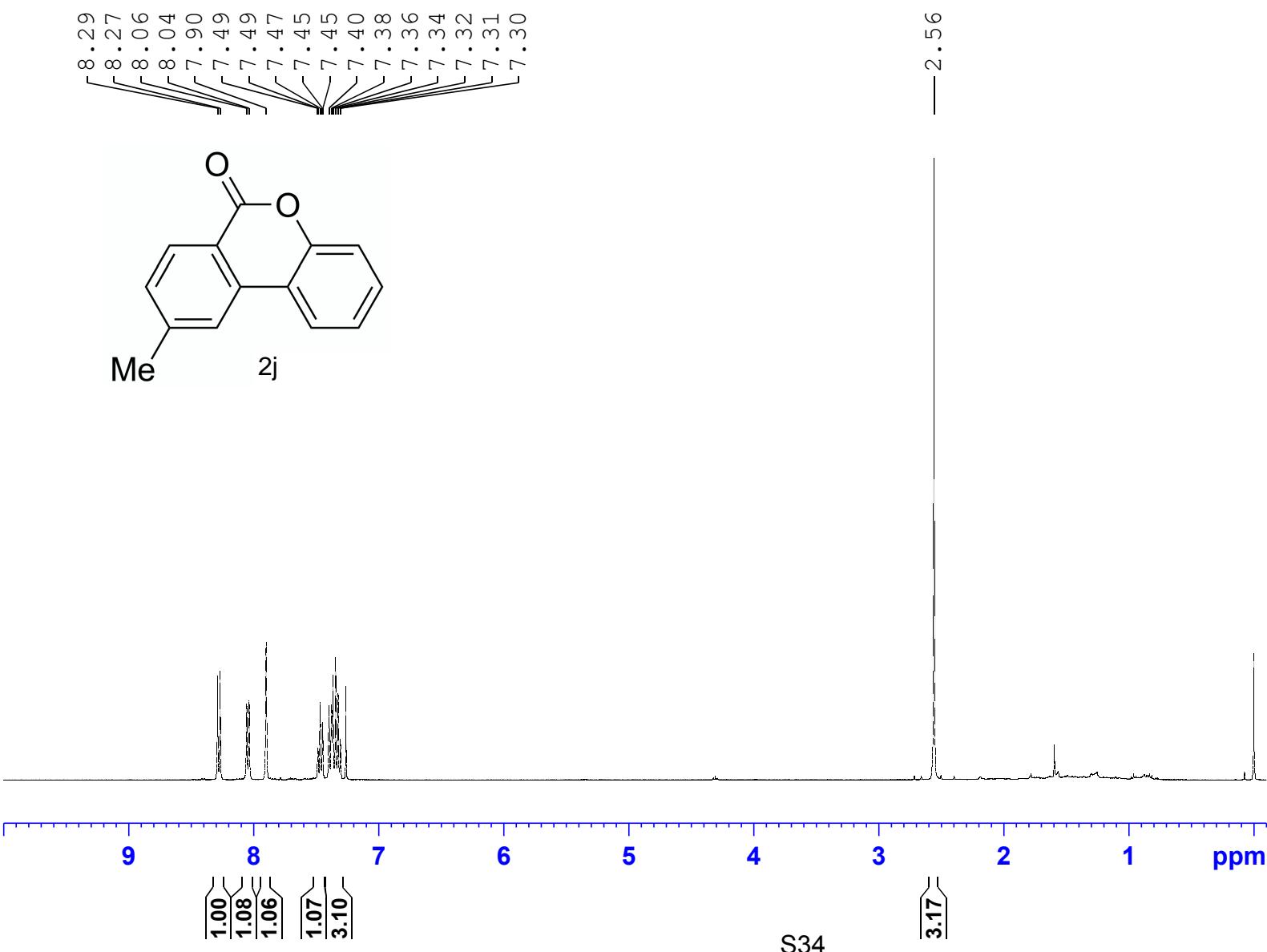
lz-975C



lz-975C



lz-979C



Current Data Parameters
NAME lz
EXPNO 106
PROCNO 1

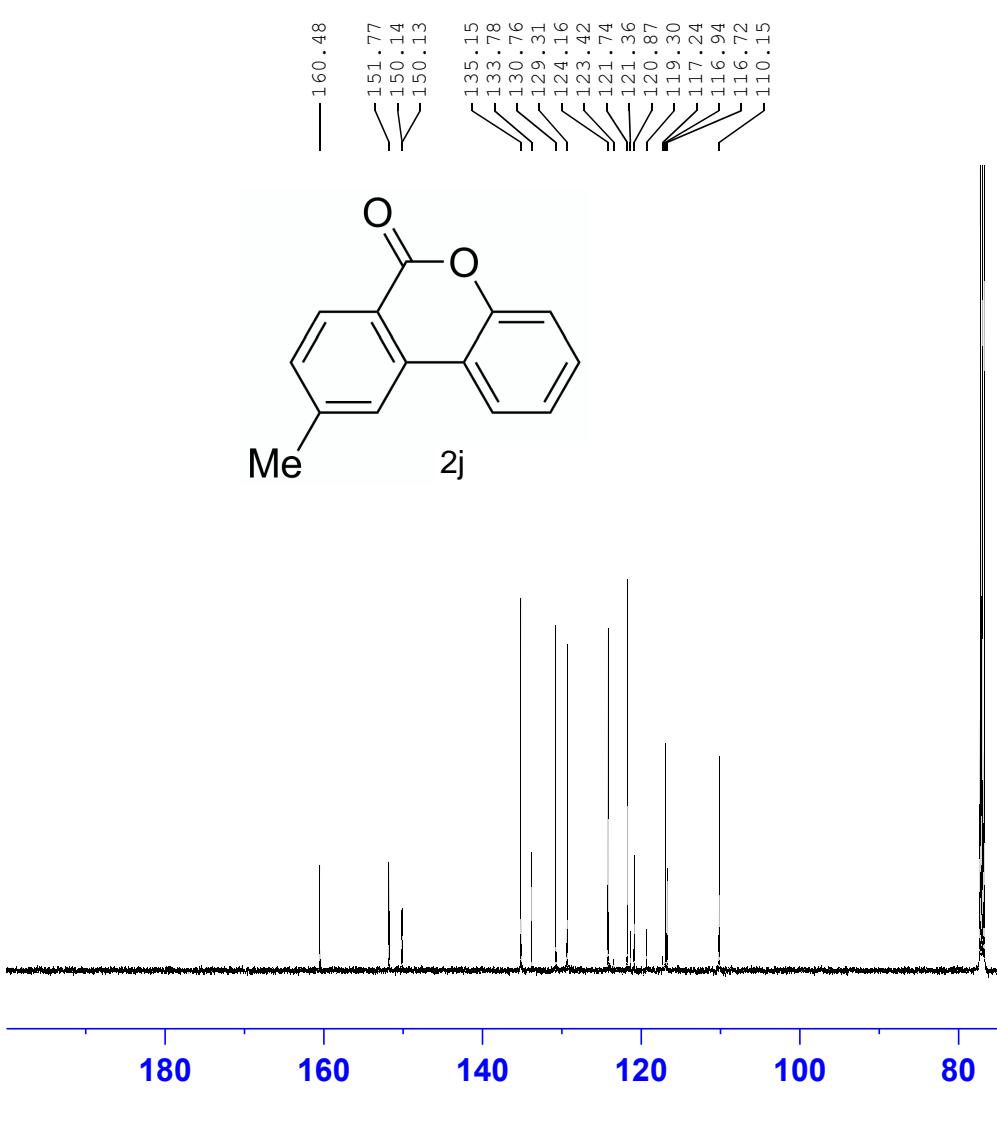
F2 - Acquisition Parameters
Date 20181120
Time 8.21
INSTRUM spect
PROBHD 5 mm PABBO BB/
PULPROG zg30
TD 32768
SOLVENT CDC13
NS 16
DS 0
SWH 8012.820 Hz
FIDRES 0.244532 Hz
AQ 2.0447233 sec
RG 206.33
DW 62.400 usec
DE 6.50 usec
TE 298.0 K
D1 2.00000000 sec
D11 0 sec
TD0 1

===== CHANNEL f1 =====
SFO1 400.2424716 MHz
NUC1 1H
P1 14.80 usec
PLW1 12.00000000 W

===== CHANNEL f2 =====
SFO2 400.2424716 MHz
NUC2 off
CPDPRG[2
PCPD2 0 usec
PLW2 0 W
PLW12 0 W
PLW13 0 W

F2 - Processing parameters
SI 65536
SF 400.2400087 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

lz-976C



Current Data Parameters
 NAME lz
 EXPNO 42
 PROCNO 1

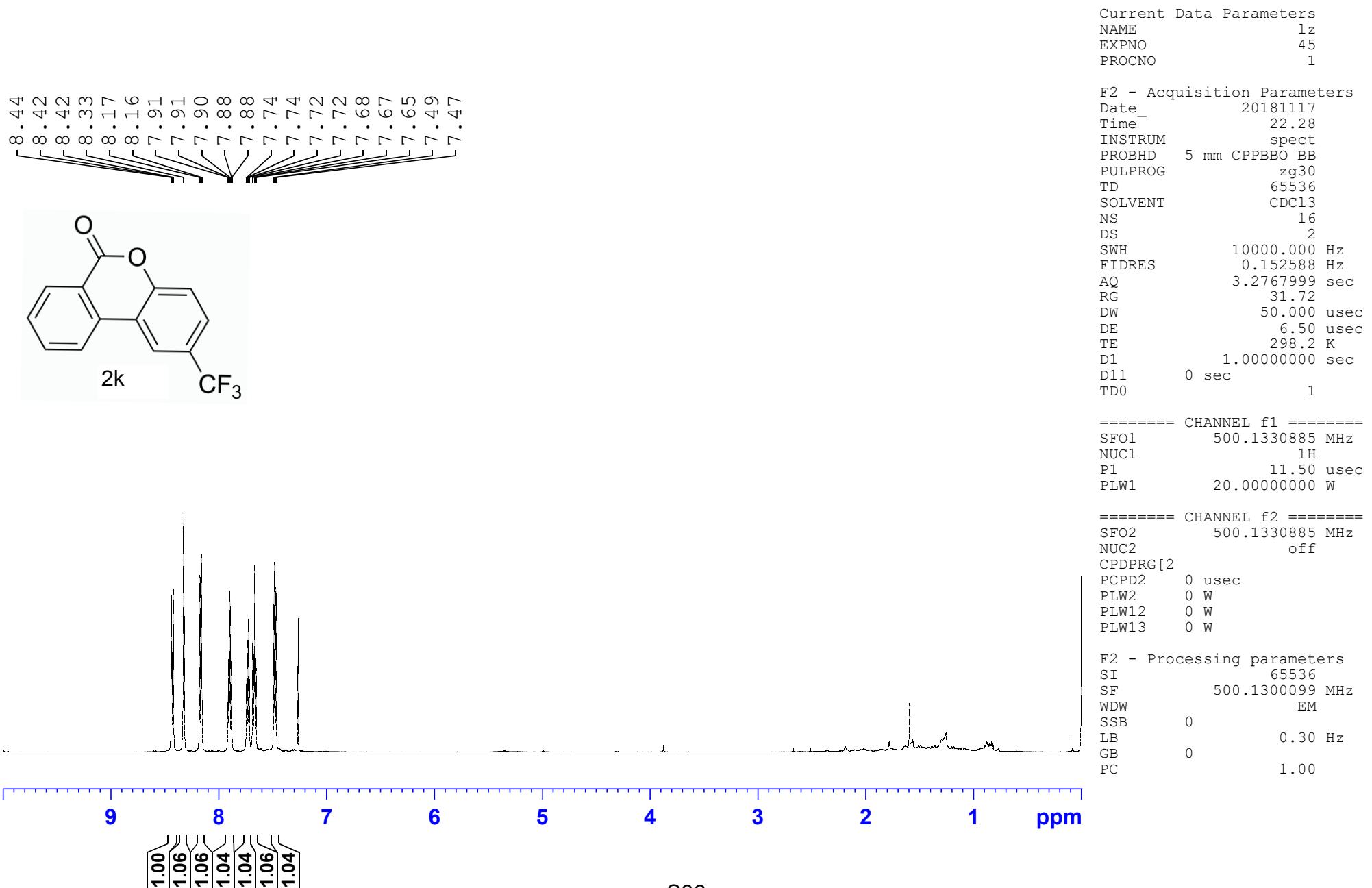
F2 - Acquisition Parameters
 Date_ 20181117
 Time_ 21.53
 INSTRUM spect
 PROBHD 5 mm CPPBBO BB
 PULPROG zgpg30
 TD 65536
 SOLVENT CDCl3
 NS 200
 DS 4
 SWH 29761.904 Hz
 FIDRES 0.454131 Hz
 AQ 1.1010048 sec
 RG 192.89
 DW 16.800 usec
 DE 18.00 usec
 TE 298.2 K
 D1 2.0000000 sec
 D11 0.0300000 sec
 TDO 1

===== CHANNEL f1 =====
 SFO1 125.7703637 MHz
 NUC1 13C
 P1 9.80 usec
 PLW1 57.00000000 W

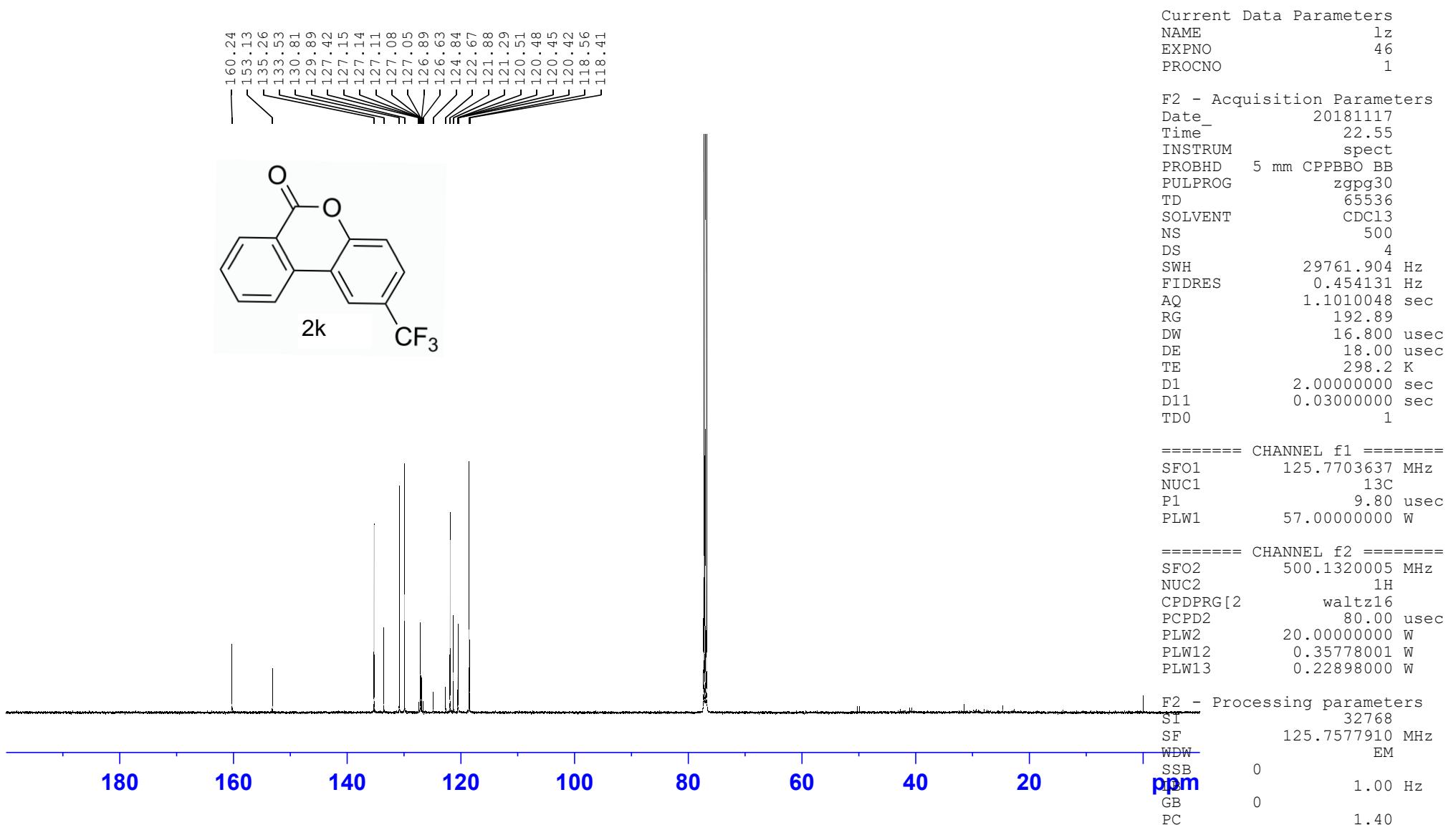
===== CHANNEL f2 =====
 SFO2 500.1320005 MHz
 NUC2 1H
 CPDPRG[2] waltz16
 PCPD2 80.00 usec
 PLW2 20.00000000 W
 PLW12 0.35778001 W
 PLW13 0.22898000 W

F2 - Processing parameters
 S1 32768
 SF 125.7577910 MHz
 WDW EM
 SSB 0
 GB 0
 PC 1.00 Hz
 1.40

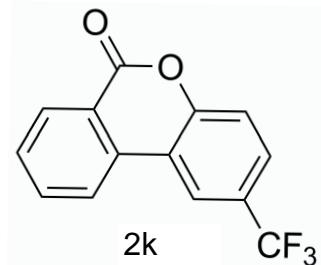
lz-977A1



lz-977A1



lz-977A1



-62.03



Current Data Parameters
NAME lz-F
EXPNO 5
PROCNO 1

F2 - Acquisition Parameters
Date 20181118
Time 13.41
INSTRUM spect
PROBHD 5 mm PABBO BB/
PULPROG zgfhiggqn.2
TD 131072
SOLVENT CDCl₃
NS 9
DS 4
SWH 89285.711 Hz
FIDRES 0.681196 Hz
AQ 0.7340032 sec
RG 206.33
DW 5.600 usec
DE 6.50 usec
TE 298.1 K
D1 1.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
SFO1 376.5642094 MHz
NUC1 ¹⁹F
P1 14.50 usec
PLW1 17.98900032 W

===== CHANNEL f2 =====
SFO2 400.2416010 MHz
NUC2 ¹H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 12.00000000 W
PLW12 0.34680000 W
PLW13 0.28090999 W

F2 - Processing parameters
SI 65536
SF 376.6018696 MHz
WDW EM
SSB 0
LB 0 0.30 Hz
GB 0
PC 1.00