

Broadening the Catalyst and Reaction Scope of Regio- and Chemoselective C-H Oxygenation: A Convenient and Scalable Approach to 2-Acylphenols by Intriguing Rh(II) and Ru(II) Catalysis

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SUPPORTING INFORMATION

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Materials and Methods

All commercial materials (Alfa Aesar, Aladdin, J&K Chemical LTD.) were used without further purification. All solvents were analytical grade. The ^1H -NMR and ^{13}C -NMR spectra were recorded on a Bruker 400 MHz spectrometer in CDCl_3 using TMS or solvent peak as a standard. All ^{13}C -NMR spectra were recorded with complete proton decoupling. Low-resolution mass spectral analyses were performed with a Waters AQUITY UPLCTM/MS. All reactions were carried out in oven-dried sealed tube. Analytical TLC was performed on Yantai Chemical Industry Research Institute silica gel 60 F254 plates and flash column chromatography was performed on Qingdao Haiyang Chemical Co. Ltd silica gel 60 (200-300 mesh). The rotavapor was BUCHI's Rotavapor R-3. Aryl ketones were commercially available or easily synthesized from acyl chlorides and Grignard reagents or from Friedel-Crafts reaction.

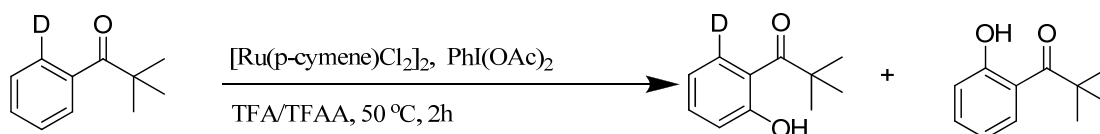
I General procedure for ruthenium and rhodium catalyzed ortho hydroxylation of aryl ketones

To a 15ml sealed-tube were added ketone (1.0 equiv), oxidants $\text{PhI}(\text{OAc})_2$, $\text{K}_2\text{S}_2\text{O}_8$ or Selectfluor (2.0 equiv), $[\text{Ru}(\text{p-cymene})\text{Cl}_2]_2$ or $\text{Rh}_2(\text{OAc})_4$ (0.025 equiv), TFAA (0.6 ml) and TFA (1.4ml). The tube was sealed and heated at 50°C or 80°C . The reaction was monitored by TLC (petroleum ether: toluene = 3:1). After completion of the reaction, dichloromethane was added to dilute the reaction mixture and saturated aqueous NaHCO_3 was added to neutralize TFA and TFAA. Then the organic layer was dried over anhydrous Na_2SO_4 and concentrated on rotavapor under reduced pressure. Finally, the residue was purified by silica gel column chromatography to give the desired product.

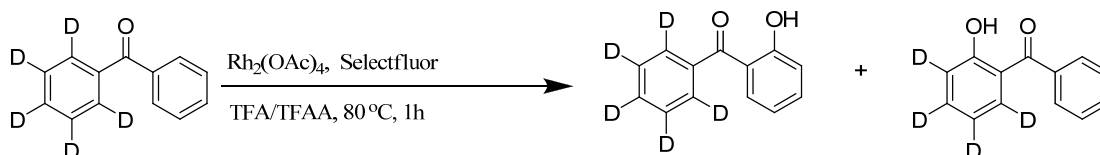
For $\text{Rh}_2(\text{OAc})_4$, 5% of catalyst was used. Selectfluor is the best oxidant for $\text{Rh}_2(\text{OAc})_4$.

II Procedure for preliminary mechanism study

1. Intramolecular Experiments



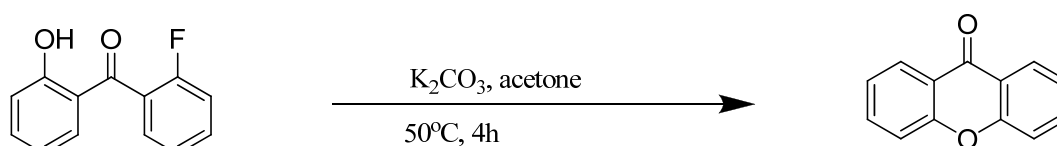
To a 15ml sealed-tube were added 2'-deuterio-2,2-dimethylpropiophenone (24mg, 0.15mmol), $\text{PhI}(\text{OAc})_2$ (100mg, 0.30 mmol), $[\text{Ru}(\text{p-cymene})\text{Cl}_2]_2$ (2mg, 0.0038mmol), TFAA (0.6 ml) and TFA (1.4 ml). The tube was sealed and heated at 50°C for 2h. The reaction mixture was cooled to room temperature. Then dichloromethane was added to dilute the reaction mixture and saturated aqueous NaHCO_3 was added to neutralize TFA and TFAA. The organic layer was dried over anhydrous Na_2SO_4 and concentrated on rotavapor under reduced pressure. The residue was eluted with DCM through a short pad of silica gel. Then the clear solution was concentrated on rotavapor under reduced pressure. ^1H -NMR shows $k_{\text{H}}/k_{\text{D}} = 2.7:1$.



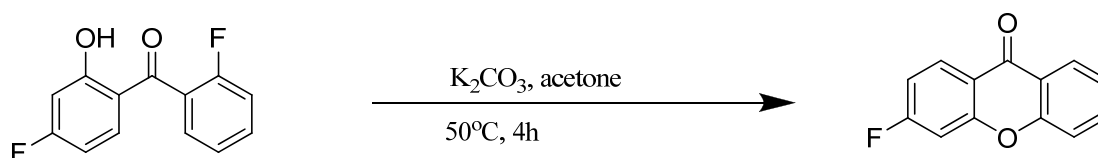
To a 15ml sealed-tube were added benzophenone – d₅ (19mg, 0.10mmol), Selectfluor (71mg, 0.20 mmol), Rh₂(OAc)₄ (2mg, 0.005mmol), TFAA (0.2 ml) and TFA (1.8 ml). The tube was sealed and heated at 80 °C for 1h. The reaction mixture was cooled to room temperature. Then dichloromethane was added to dilute the reaction mixture and saturated aqueous NaHCO₃ was added to neutralize TFA and TFAA. The organic layer was dried over anhydrous Na₂SO₄ and concentrated on rotavapor under reduced pressure. The residue was eluted with DCM through a short pad of silica gel. Then the clear solution was concentrated on rotavapor under reduced pressure. ¹H-NMR shows k_H/k_D = 2.1:1.

III Further transformations of 2-hydroxy aromatic ketones

1. Synthesis of 9H-xanthen-9-one

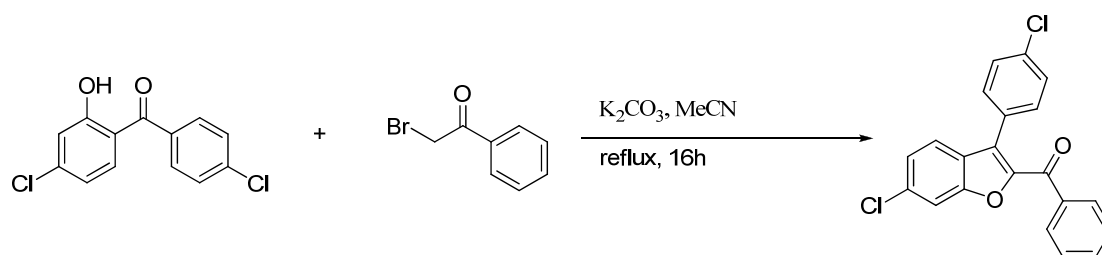


To a 15ml sealed-tube were added (2-fluorophenyl)(2-hydroxyphenyl)methanone (33 mg, 1 equiv), K₂CO₃ (42 mg, 2 equiv) and 2ml of acetone at room temperature. The tube was sealed and heated at 50 °C for 4h. Then the resulting mixture was allowed to cool to room temperature and extracted with CH₂Cl₂. The organic layer was dried over anhydrous Na₂SO₄ and concentrated on rotavapor under reduced pressure. Finally, the residue was purified by silica gel column chromatography to give the desired product (**38**) in 98% (29 mg, white solid) yield.



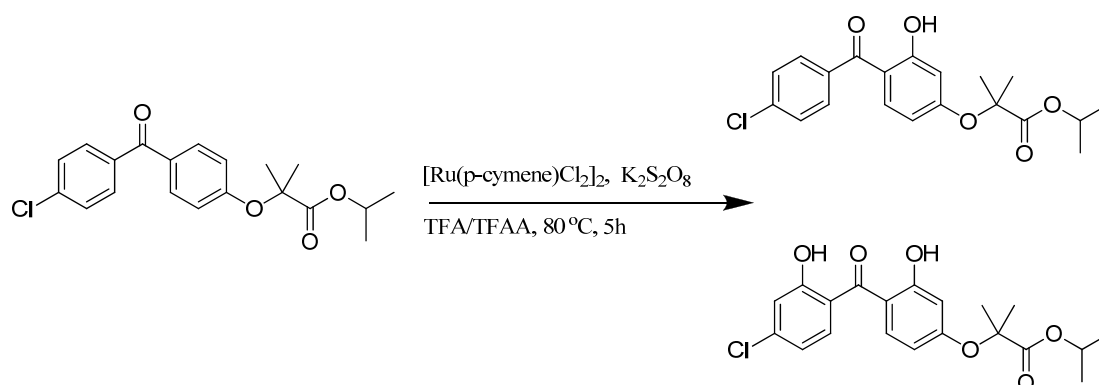
To a 15ml sealed-tube were added (4-fluoro-2-hydroxyphenyl)(2-fluorophenyl) Methanone (47 mg, 1 equiv), K₂CO₃ (55 mg, 2 equiv) and 2ml of acetone at room temperature. The tube was sealed and heated at 50 °C for 4h. Then the resulting mixture was allowed to cool to room temperature and extracted with CH₂Cl₂. The organic layer was dried over anhydrous Na₂SO₄ and concentrated on rotavapor under reduced pressure. Finally, the residue was purified by silica gel column chromatography to give the desired product compound (**39**) in 93% (40 mg, white solid) yield.

2. Synthesis of (6-chloro-3-(4-chlorophenyl)benzofuran-2-yl)(phenyl)methanone



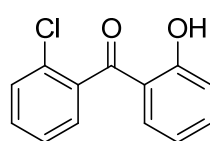
To a solution of (4-chloro-2-hydroxyphenyl)(4-chlorophenyl)methanone (80 mg, 0.30 mmol) in CH₃CN (5 mL) was added K₂CO₃ (124 mg, 0.90 mmol) and 2-bromoacetophenone (72 mg, 0.36 mmol) and the mixture refluxed for 16 h. The solvents were removed under reduced pressure and the residue dissolved in EtOAc and washed with water. The organic layer was dried over MgSO₄, concentrated, and the residue was purified by column chromatography on silica gel (hexane/EtOAc: 20/1) to give (6-chloro-3-(4-chlorophenyl)benzofuran-2-yl)(phenyl)methanone (**36**) in 95% (104 mg, pale yellow solid) yield.

IV Drug modification: Hydroxylation of Fenofibrate

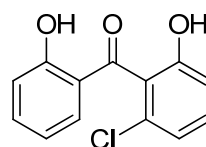


To a 15ml sealed-tube were added Fenofibrate (73mg, 0.20mmol), K₂S₂O₈ (108mg, 0.40 mmol), [Ru(p-cymene)Cl₂]₂ (3.1mg, 0.005mmol), TFAA (0.6 ml) and TFA (1.4 ml). The tube was sealed and heated at 80 °C for 5h. The reaction mixture was cooled to room temperature. Then dichloromethane was added to dilute the reaction mixture and saturated aqueous NaHCO₃ was added to neutralize TFA and TFAA. Then the organic layer was dried over anhydrous Na₂SO₄ and concentrated on rotavapor under reduced pressure. Finally, the residue was purified by silica gel column chromatography (Petroleum ether: Ethylacetate = 200:1) to give a mixture of Mono-OH- Fenofibrate (**41a**) and Di-OH- Fenofibrate (**41b**) (41.3mg, yellow oil) in 54% yield. ¹H-NMR shows the ratio is 1:0.55. Mono-OH- Fenofibrate (**41a**) LRMS (ESI) calcd for C₂₀H₂₂ClO₅ [M+H]⁺: 377.11, found 377.04; Di-OH- Fenofibrate (**41b**) LRMS (ESI) calcd for C₂₀H₂₂ClO₆ [M+H]⁺: 393.10, found 393.03;

Data of products



(2a)



(2b)

(2-Chlorophenyl)(2-hydroxyphenyl)methanone (**2a**)

(2-Chloro-6-hydroxyphenyl)(2-hydroxyphenyl)methanone (**2b**)

1) Small-scale

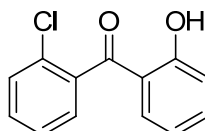
Following the general procedure I , 2-chlorobenzophenone (44 mg, 0.20 mmol), $\text{PhI}(\text{OAc})_2$ (129 mg, 0.40 mmol), $[\text{Ru}(\text{p-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 3h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound (**2a**) (34 mg, yellow oil) was isolated in 72% yield.

2) One-gram-scale

Following the general procedure I , 2-chlorobenzophenone (1g, 4.6 mmol), $\text{PhI}(\text{OAc})_2$ (2.2 g, 6.9 mmol), $[\text{Ru}(\text{p-cymene})\text{Cl}_2]_2$ (28.3 mg, 0.046 mmol), 12ml TFA and 18ml TFAA were used. The reaction mixture was stirred at 80 °C for 14h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1 or Petroleum ether: Ethylacetate = 100:1). Finally, compound (**2a**) (606 mg, yellow oil) and compound (**2b**) (178 mg, yellow solid) were isolated in 56% and 15% yield respectively.

3) Two-gram-scale

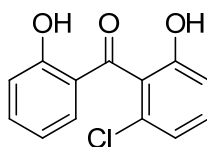
Following the general procedure I , 2-chlorobenzophenone (2g, 9.2 mmol), $\text{PhI}(\text{OAc})_2$ (4.5 g, 13.6 mmol), $[\text{Ru}(\text{p-cymene})\text{Cl}_2]_2$ (56.5 mg, 0.092 mmol), 12ml TFA and 18ml TFAA were used. The reaction mixture was stirred at 80 °C for 14h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1 or Petroleum ether: Ethylacetate = 100:1). Finally, compound (**2a**) (1.2 g, yellow oil) and compound (**2b**) (548 mg, yellow solid) were isolated in 56% and 24% yield respectively.



(2a)

(2-Chlorophenyl)(2-hydroxyphenyl)methanone (**2a**)

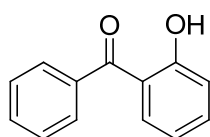
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 11.95 (s, 1H), 7.53 – 7.34 (m, 5H), 7.25 (d, J = 8.44 Hz, 1H), 7.07 (d, J = 8.44 Hz, 1H), 6.83 (t, J = 7.60 Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 200.7, 163.4, 137.5, 137.3, 133.7, 131.4, 131.0, 130.2, 128.7, 126.9, 119.5, 119.3, 118.5; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{ClO}_2$ $[\text{M}+\text{H}]^+$: 233.04, found 232.97



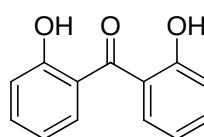
(2b)

(2-Chloro-6-hydroxyphenyl)(2-hydroxyphenyl)methanone (2b)

^1H -NMR (400 MHz, CDCl_3) δ (ppm) 11.73 (s, 1H), 7.51 (t, $J = 8.16$ Hz, 1H), 7.35 (d, $J = 8.00$ Hz, 1H), 7.29 (t, $J = 8.16$ Hz, 1H), 7.06 – 7.03 (m, 2H), 6.88 – 6.83 (m, 2H), 6.16 (s, 1H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 200.0, 163.0, 154.7, 137.5, 133.2, 131.97, 131.93, 124.4, 122.1, 120.0, 119.4, 118.4, 115.4; HRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{ClO}_3$ $[\text{M}+\text{H}]^+$: 249.0313, found 249.0315



(3a)



(3b)

2-Hydroxybenzophenone (3a)

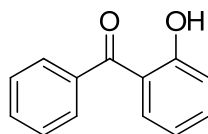
2, 2'-Dihydroxybenzophenone (3b)

1) $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$

Following the general procedure I, benzophenone (36.4 mg, 0.20 mmol), $\text{K}_2\text{S}_2\text{O}_8$ (108 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4 ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 3h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound (3a) (25 mg, yellow oil) and compound (3b) (13 mg, yellow solid) were isolated in 62% yield and 31% yield respectively.

2) $\text{Rh}_2(\text{OAc})_4$

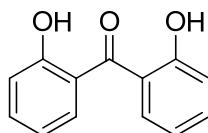
5% $\text{Rh}_2(\text{OAc})_4$, 1.1eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture were stirred at 70 °C for 10h. Compound (3a) was isolated in 62% yield.



(3a)

2-Hydroxybenzophenone (3a)

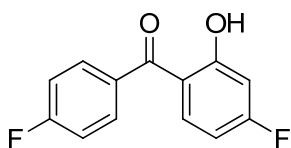
^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.03 (s, 1H), 7.68 (d, $J = 7.08$ Hz, 2H), 7.61 – 7.57 (m, 2H), 7.50 (m, 3H), 7.07 (d, $J = 8.36$ Hz, 1H), 6.87 (t, $J = 7.96$ Hz, 1H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 201.8, 163.4, 138.1, 136.5, 133.8, 132.1, 129.3, 128.5, 119.3, 118.8, 118.6; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{11}\text{O}_2$ $[\text{M}+\text{H}]^+$: 199.08, found 199.02;



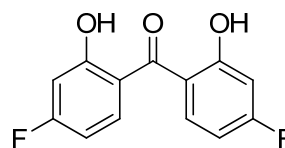
(3b)

2, 2'-Dihydroxybenzophenone **(3b)**

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 10.59 (s, 2H), 7.63 (d, $J = 7.96$ Hz, 2H), 7.52 (t, $J = 7.88$ Hz, 2H), 7.10 (d, $J = 8.36$ Hz, 2H), 6.95 (t, $J = 7.58$ Hz, 2H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 202.6, 161.9, 136.1, 133.2, 120.0, 119.0, 118.8; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{11}\text{O}_3$ $[\text{M}+\text{H}]^+$: 215.06, found 215.01



(4a)



(4b)

(4-Fluoro-2-hydroxyphenyl)(4-fluorophenyl)methanone **(4a)**

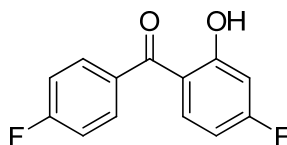
Bis(4-fluoro-2-hydroxyphenyl)methanone **(4b)**

1) $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$

Following the general procedure I, 4,4'-difluorobenzophenone (44 mg, 0.20 mmol), $\text{K}_2\text{S}_2\text{O}_8$ (108 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 6h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound **(4a)** (26 mg, yellow solid) was isolated in 56% yield and **(4b)** (16mg, yellow solid) in 33% yield.

2) $\text{Rh}_2(\text{OAc})_4$

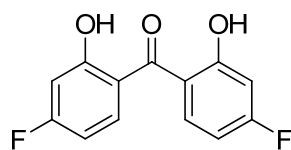
5% $\text{Rh}_2(\text{OAc})_4$, 1.1eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture were stirred at 70 °C for 10h. Compound **(4a)** and compound **(4b)** were isolated in 68% yield and 17% yield respectively.



(4a)

(4-Fluoro-2-hydroxyphenyl)(4-fluorophenyl)methanone **(4a)**

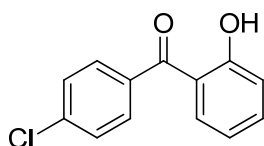
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 12.28 (s, 1H), 7.69 (dd, $J = 8.44$ Hz, $J = 5.48$ Hz, 2H), 7.59 (dd, $J = 8.84$ Hz, $J = 6.60$ Hz, 1H), 7.20 (t, $J = 8.52$ Hz, 2H), 6.76 (dd, $J = 8.84$ Hz, $J = 10.32$ Hz, 1H), 6.61 (td, $J = 10.68$ Hz, $J = 2.36$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 199.1, 167.7 (d, $J = 247$ Hz, 1C), 165.1 (d, $J = 243$ Hz, 1C), 166.4 (d, $J = 15$ Hz, 1C), 135.8 (d, $J = 16$ Hz, 1C), 134.0 (d, $J = 3.2$ Hz, 1C), 131.8 (d, $J = 9$ Hz, 2C), 116.2 (d, $J = 2.1$ Hz, 1C), 115.9 (d, $J = 22$ Hz, 2C), 107.2 (d, $J = 23$ Hz, 1C), 105.4 (d, $J = 24$ Hz, 1C); LRMS (ESI) calcd for $\text{C}_{13}\text{H}_9\text{F}_2\text{O}_2$ $[\text{M}+\text{H}]^+$: 235.06, found 234.94;



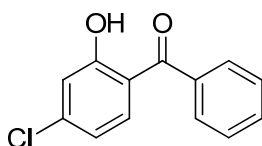
(4b)

Bis(4-fluoro-2-hydroxyphenyl)methanone **(4b)**

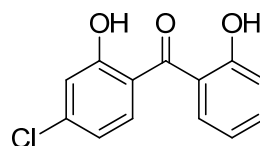
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 10.86 (s, 2H), 7.60 (dd, $J = 8.84$ Hz, $J = 6.44$ Hz, 2H), 6.78 (dd, $J = 10.20$ Hz, $J = 2.40$ Hz, 2H), 6.67 (td, $J = 8.38$ Hz, $J = 2.40$ Hz, 2H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 200.2, 167.4 (d, $J = 266.0$ Hz), 164.5 (d, $J = 14.0$ Hz), 135.1 (d, $J = 11.6$ Hz), 116.7 (d, $J = 2.5$ Hz), 107.4 (d, $J = 22.6$ Hz), 105.7 (d, $J = 23.9$ Hz); LRMS (ESI) calcd for $\text{C}_{13}\text{H}_9\text{F}_2\text{O}_3$ $[\text{M}+\text{H}]^+$: 251.05, found 251.03



(5a)



(5b)



(5c)

(4-Chlorophenyl)(2-hydroxyphenyl)methanone **(5a)**

(4-Chloro-2-hydroxyphenyl)(phenyl)methanone **(5b)**

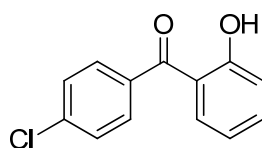
(4-Chloro-2-hydroxyphenyl)(phenyl)methanone **(5c)**

1) $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$

Following the general procedure I, 4-chlorobenzophenone (65 mg, 0.30 mmol), $\text{PhI}(\text{OAc})_2$ (200 mg, 0.60 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 15h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound **(5a)** (22 mg, yellow solid) and compound **(5c)** (45 mg, yellow solid) were isolated in 32% yield and 63% yield respectively.

2) $\text{Rh}_2(\text{OAc})_4$

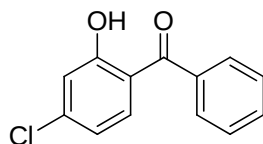
0.1mmol of 4-chlorobenzophenone, 5% $\text{Rh}_2(\text{OAc})_4$, 1.2eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture were stirred at 80 °C for 10.5h. The conversion ratio for compound **(5a)**, compound **(5b)** and compound **(5c)** were 44%, 11% and 7% respectively.



(5a)

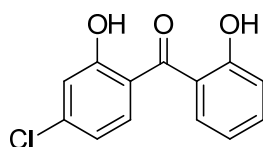
(4-Chlorophenyl)(2-hydroxyphenyl)methanone **(5a)**

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 11.87 (s, 1H), 7.64 (d, $J = 7.96$ Hz, 2H), 7.55 – 7.48 (m, 4H), 7.08 (d, $J = 8.48$ Hz, 1H), 6.89 (t, $J = 7.72$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 200.4, 163.4, 138.6, 136.7, 136.3, 133.4, 130.8, 128.9, 119.1, 118.9, 118.7; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{ClO}_2$ $[\text{M}+\text{H}]^+$: 233.04, found 232.95



(4-Chloro-2-hydroxyphenyl)(phenyl)methanone (**5b**)

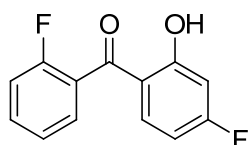
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 12.18 (s, 1H), 7.67 – 7.59 (m, 3H), 7.54 – 7.50 (m, 3H), 7.10 (s, 1H), 6.86 (d, $J = 8.56$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 201.0, 164.1, 142.4, 137.7, 134.6, 132.3, 129.2, 128.6, 119.5, 118.7, 117.9; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{ClO}_2$ $[\text{M}+\text{H}]^+$: 233.04, found 232.95



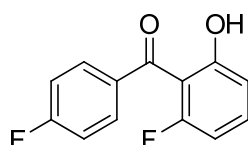
(**5c**)

(4-Chloro-2-hydroxyphenyl)(phenyl)methanone (**5c**)

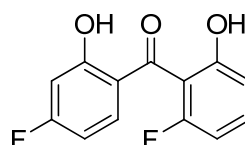
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 10.81 (s, 1H), 10.43 (s, 1H), 7.58 – 7.53 (m, 3H), 7.12 – 7.09 (m, 2H), 6.95 – 6.92 (m, 2H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 201.6, 162.7, 161.8, 142.1, 136.3, 134.2, 132.8, 119.8, 119.7, 119.2, 118.9, 118.4; HRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{ClO}_3$ $[\text{M}+\text{H}]^+$: 249.0313, found 249.0315



(**6a**)



(**6b**)



(**6c**)

(4-Fluoro-2-hydroxyphenyl)(2-fluorophenyl)methanone (**6a**)

(2-Fluoro-6-hydroxyphenyl)(4-fluorophenyl)methanone (**6b**)

(4-Fluoro-2-hydroxyphenyl)(2-fluoro-6-hydroxyphenyl)methanone (**6c**)

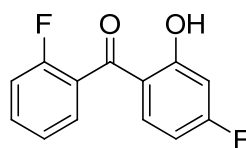
1)[$\text{Ru}(\text{p-cymene})\text{Cl}_2$] $_2$

Following the general procedure I, 2,4'-difluorobenzophenone (44 mg, 0.20 mmol), $\text{PhI}(\text{OAc})_2$ (133 mg, 0.40 mmol), [$\text{Ru}(\text{p-cymene})\text{Cl}_2$] $_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 6h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 4 : 1). Finally, compound (**6a**) (14mg, white solid) and (**6b**) (24 mg, white solid) were isolated in 30% and 51% yield.

2) $\text{Rh}_2(\text{OAc})_4$

0.1mmol of 2,4'-difluorobenzophenone, 5% $\text{Rh}_2(\text{OAc})_4$, 1.2eq. of Selectfluor, 1.8ml

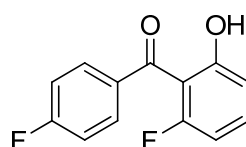
TFA and 0.2ml TFAA were used. The reaction mixture were stirred at 80 °C for 8h. The conversion ratio for compound (**6a**), compound (**6b**) and compound (**6c**) were 16%, 40% and 16% respectively.



(**6a**)

(4-Fluoro-2-hydroxyphenyl)(2-fluorophenyl)methanone (**6a**)

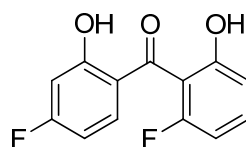
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 12.30 (s, 1H), 7.60 – 7.50 (m, 1H), 7.48 – 7.41 (m, 2H), 7.30 (t, $J = 7.52$ Hz, 1H), 7.20 (t, $J = 9.00$ Hz, 1H), 6.73 (dd, $J = 10.28$ Hz, $J = 2.40$ Hz, 1H), 6.58 (dd, $J = 8.48$ Hz, $J = 2.40$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 197.4, 168.0 (d, $J = 256.7$ Hz), 165.7 (d, $J = 14.54$ Hz), 159.1 (d, $J = 250.0$ Hz), 136.1 (d, $J = 2.6$ Hz), 136.0 (d, $J = 2.3$ Hz), 133.2 (d, $J = 8.2$ Hz), 129.9 (d, $J = 2.7$ Hz), 126.3 (d, $J = 15.6$ Hz), 124.6 (d, $J = 3.6$ Hz), 117.1 (d, $J = 2.2$ Hz), 116.5 (d, $J = 21.2$ Hz), 107.6 (d, $J = 21.9$ Hz), 105.1 (d, $J = 23.9$ Hz); LRMS (ESI) calcd for $\text{C}_{13}\text{H}_9\text{F}_2\text{O}_2$ $[\text{M}+\text{H}]^+$: 235.06, found 234.97



(**6b**)

(2-Fluoro-6-hydroxyphenyl)(4-fluorophenyl)methanone (**6b**)

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 10.86 (s, 1H), 7.76 – 7.73 (m, 2H), 7.44 (m, 1H), 7.14 (t, $J = 8.48$ Hz, 2H), 6.89 (d, $J = 8.44$ Hz, 1H), 6.63 (t, $J = 8.48$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) 196.5, 165.7 (d, $J = 253.1$ Hz), 162.4 (d, $J = 3.7$ Hz), 161.6 (d, $J = 254.0$ Hz), 136.1 (d, $J = 11.6$ Hz), 135.92 (d, $J = 3.4$ Hz), 135.89 (d, $J = 3.4$ Hz), 131.6 (d, $J = 3.7$ Hz), 131.5 (d, $J = 3.6$ Hz), 115.6 (d, $J = 22.0$ Hz), 114.2 (d, $J = 3.3$ Hz), 110.5 (d, $J = 14.7$ Hz), 106.9 (d, $J = 22.8$ Hz); LRMS (ESI) calcd for $\text{C}_{13}\text{H}_9\text{F}_2\text{O}_2$ $[\text{M}+\text{H}]^+$: 235.06, found 234.96

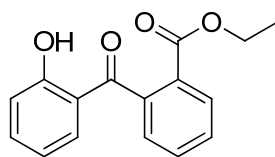


(**6c**)

(4-Fluoro-2-hydroxyphenyl)(2-fluoro-6-hydroxyphenyl)methanone (**6c**)

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 11.66 (s, 1H), 8.49 (s, 1H), 7.55 – 7.53 (m, 1H), 7.45 – 7.39 (m, 1H), 6.88 (d, $J = 8.36$ Hz, 1H), 6.74 – 6.69 (m, 2H), 6.62 (td, $J = 8.46$ Hz, $J = 2.08$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) 197.9, 168.17 (d, $J = 257.3$ Hz), 165.3 (d, $J = 14.6$ Hz), 160.0 (d, $J = 251.3$ Hz), 159.0 (d, $J = 17.6$ Hz), 136.0 (d, $J = 7.0$ Hz), 135.9 (d, $J = 7.2$ Hz), 135.1 (d, $J = 11.1$ Hz), 117.8, 114.1 (d, $J = 3$ Hz),

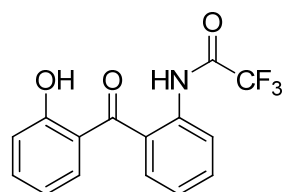
112.0 (d, $J = 17.3$ Hz), 107.9 (d, $J = 22.8$ Hz), 107.6 (d, $J = 22.6$ Hz), 105.1 (d, $J = 24$ Hz); LRMS (ESI) calcd for $C_{13}H_9F_2O_3$ $[M+H]^+$: 251.05, found 250.91



(7)

Ethyl 2-(2-hydroxybenzoyl)benzoate (7)

Following the general procedure I , ethyl 2-benzoylbenzoate (76 mg, 0.30 mmol), $K_2S_2O_8$ (162 mg, 0.60 mmol), $[Ru(p\text{-cymene})Cl_2]_2$ (3.1 mg, 0.005 mmol), 0.8ml TFA and 1.2 ml TFAA were used. The reaction mixture was stirred at 80 °C for 18h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (7) (38 mg, white solid) was isolated in 47% yield. 1H -NMR (400 MHz, $CDCl_3$) δ (ppm) 11.97 (s, 1H), 8.11 (dd, $J = 7.64$ Hz, $J = 0.84$ Hz, 1H), 7.66 (td, $J = 7.48$ Hz, $J = 1.24$ Hz, 1H), 7.59 (td, $J = 8.52$ Hz, $J = 1.32$ Hz, 1H), 7.46 (td, $J = 7.80$ Hz, $J = 1.60$ Hz, 1H), 7.38 (dd, $J = 7.56$ Hz, $J = 1.04$ Hz, 1H), 7.09 (dd, $J = 7.96$ Hz, $J = 1.56$ Hz, 1H), 7.04 (d, $J = 8.12$ Hz, 1H), 6.76 (td, $J = 7.58$ Hz, $J = 0.88$ Hz, 1H), 4.14 (q, $J = 7.16$ Hz, 2H), 1.09 (t, $J = 7.16$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ (ppm) 203.0, 165.6, 162.6, 140.0, 136.5, 132.7, 132.6, 130.5, 130.0, 129.0, 127.4, 120.5, 119.0, 118.3, 61.8, 13.6; LRMS (ESI) calcd for $C_{16}H_{15}O_4$ $[M+H]^+$: 271.10, found 271.06



(8)

2,2,2-Trifluoro-N-(2-(2-hydroxybenzoyl)phenyl)acetamide (8)

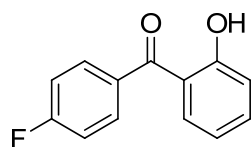
1) $[Ru(p\text{-cymene})Cl_2]_2$

Following the general procedure I , N-(2-benzoylphenyl)-2,2,2-trifluoroacetamide (88 mg, 0.30 mmol), $K_2S_2O_8$ (162 mg, 0.60 mmol), $[Ru(p\text{-cymene})Cl_2]_2$ (6 mg, 0.01 mmol), 1.0ml TFA and 1.0 ml TFAA were used. The reaction mixture was stirred at 80 °C for 15h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1 or Petroleum ether: Ethylacetate = 10:1). Finally, compound (8) (71 mg, yellow solid) was isolated in 77% yield.

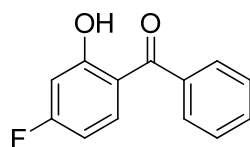
2) $Rh_2(OAc)_4$

0.1mmol of 2-aminobenzophenone , 5% $Rh_2(OAc)_4$, 1.2eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture was stirred at 80 °C for 7h. The conversion ratio for compound (8) was 61%. 1H -NMR (400 MHz, $CDCl_3$) δ (ppm) 11.27 (s, 1H), 10.62 (s, 1H), 8.46 (d, $J = 8.32$ Hz, 1H), 7.66 (t, $J = 7.68$ Hz, 1H), 7.61 (d, $J = 7.48$ Hz, 1H), 7.56 (t, $J = 7.32$ Hz, 1H), 7.52 (d, $J = 8.00$ Hz, 1H), 7.32 (d, $J =$

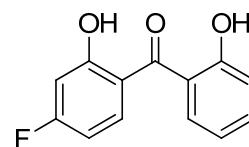
7.64 Hz, 1H), 7.10 (d, $J = 8.36$ Hz, 1H), 6.92 (d, $J = 7.44$ Hz, 1H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 201.81, 163.56, 155.34 (q, $J = 38$ Hz), 137.4, 136.1, 133.87, 133.78, 132.7, 126.0, 124.7, 122.8, 119.6, 119.2, 119.0, 115.8 (q, $J = 286.8$ Hz); ^{19}F -NMR (376 MHz, CDCl_3) δ (ppm) -76.14; LRMS (ESI) calcd for $\text{C}_{15}\text{H}_{11}\text{F}_3\text{NO}_3$ $[\text{M}+\text{H}]^+$: 310.07, found 310.09



(9a)



(9b)



(9c)

(4-Fluorophenyl)(2-hydroxyphenyl)methanone **(9a)**

(4-Fluoro-2-hydroxyphenyl)(phenyl)methanone **(9b)**

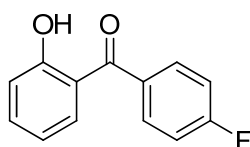
(4-Fluoro-2-hydroxyphenyl)(2-hydroxyphenyl)methanone **(9c)**

1) $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$

Following the general procedure I, 4-fluorobenzophenone (60 mg, 0.30 mmol), $\text{PhI}(\text{OAc})_2$ (200 mg, 0.60 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 60 °C for 15h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound **(9a)** (17.5 mg, yellow solid) and compound **(9c)** (43.5 mg, yellow solid) was isolated in 27% yield and 65% yield respectively.

2) $\text{Rh}_2(\text{OAc})_4$

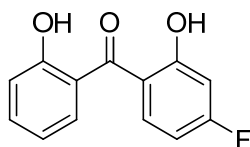
0.1mmol of 4-fluorobenzophenone, 5% $\text{Rh}_2(\text{OAc})_4$, 1.2eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture was stirred at 80 °C for 7h. The conversion ratio for compound **(9a)**, compound **(9b)** and compound **(9c)** were 51%, 8% and 13% respectively.



(9a)

(4-Fluorophenyl)(2-hydroxyphenyl)methanone **(9a)**

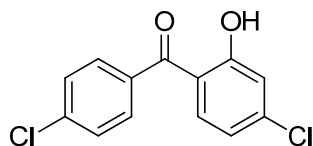
^1H -NMR (400 MHz, CDCl_3) δ (ppm) 11.88 (s, 1H), 7.74-7.71 (m, 2H), 7.56 (d, $J = 8.0$ Hz, 1H), 7.52 (t, $J = 8.0$ Hz, 1H), 7.19 (t, $J = 8.4$ Hz, 2H), 7.08 (d, $J = 8.4$ Hz, 1H), 6.89 (t, $J = 7.2$ Hz, 1H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 200.1, 165.2 (d, $J = 252.1$ Hz, 1C), 163.3, 136.5, 134.2 (d, $J = 3.1$ Hz, 1C), 133.4, 131.9 (d, $J = 8.9$ Hz, 1C), 119.2, 118.9, 118.7, 115.7 (d, $J = 21.7$ Hz, 1C); LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{FO}_2$ $[\text{M}+\text{H}]^+$: 217.06, found 216.90



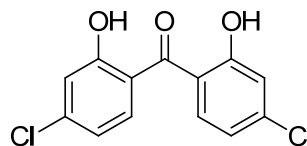
(9c)

(4-Fluoro-2-hydroxyphenyl)(2-hydroxyphenyl)methanone **(9c)**

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 11.12 (s, 1H), 10.33 (s, 1H), 7.67-7.63 (m, 1H), 7.56-7.50 (m, 2H), 7.09 (d, $J = 8.3$ Hz, 1H), 6.95 (t, $J = 7.6$ Hz, 1H), 6.77 (dd, $J = 10.2$ Hz, $J = 2.0$ Hz, 1H), 6.68-6.64 (m, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 201.3, 167.4 (d, $J = 256$ Hz, 1C), 164.8 (d, $J = 15$ Hz, 1C), 161.6, 136.1, 135.7 (d, $J = 11.4$ Hz, 1C), 132.7, 120.0, 119.1, 118.8, 116.7 (d, $J = 2.2$ Hz, 1C), 107.3 (d, $J = 23$ Hz, 1C), 105.6 (d, $J = 24$ Hz, 1C); HRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{FO}_3$ $[\text{M}+\text{H}]^+$: 233.0608, found 232.0605



(10a)



(10b)

(4-Chloro-2-hydroxyphenyl)(4-chlorophenyl)methanone **(10a)**

Bis(4-chloro-2-hydroxyphenyl)methanone **(10b)**

1) $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$

Following the general procedure I, 4,4'-dichlorobenzophenone (51 mg, 0.20 mmol), $\text{K}_2\text{S}_2\text{O}_8$ (108 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 5.5h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 4:1). Finally, compound **(10a)** (30 mg, yellow solid) was isolated in 56% yield and **(10b)** (17mg, yellow solid) in 29% yield.

2) $\text{Rh}_2(\text{OAc})_4$

0.1mmol of 4-fluorobenzophenone, 5% $\text{Rh}_2(\text{OAc})_4$, 1.2eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture was stirred at 80 °C for 11h. The conversion ratio for compound **(10a)** and compound **(10b)** were 62% and 20% respectively.

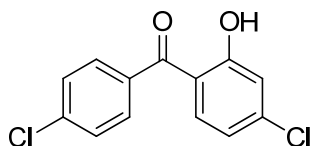
3) one-gram-scale

Following the general procedure I, 4,4'-dichlorobenzophenone (1 g, 4.0 mmol), $\text{K}_2\text{S}_2\text{O}_8$ (2160 mg, 8.0 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (12 mg, 0.020 mmol), 14ml TFA and 6ml TFAA were used. The reaction mixture was stirred at 80 °C for 24h. The residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 4:1 or Petroleum ether: Ethylacetate = 100:1). Finally, compound **(10a)** (650 mg, yellow solid) was isolated in 65% yield. Some starting material remained.

4) two-gram-scale

Following the general procedure I, 4,4'-dichlorobenzophenone (2 g, 7.96 mmol), $\text{K}_2\text{S}_2\text{O}_8$ (4300 mg, 15.9 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (48.8 mg, 0.080 mmol), 21ml TFA and 9ml TFAA were used. The reaction mixture was stirred at 80 °C for 24h.

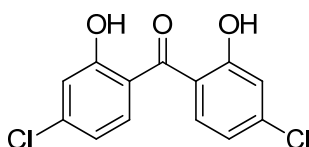
After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 4:1 or Petroleum ether: Ethylacetate = 100:1). Finally, compound (**10a**) (650 mg, yellow solid) and (**10b**) (435mg, yellow solid) were isolated in 51% and 23% yield respectively.



(**10a**)

(4-Chloro-2-hydroxyphenyl)(4-chlorophenyl)methanone (**10a**)

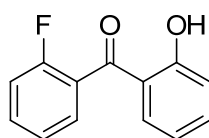
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 12.03 (s, 1H), 7.62 (d, $J = 8.40$ Hz, 2H), 7.51 – 7.47 (m, 3H), 7.10 (s, 1H), 6.87 (d, $J = 8.60$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 199.6, 164.0, 142.7, 138.9, 136.0, 134.2, 130.7, 129.0, 119.7, 118.9, 117.6; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_9\text{Cl}_2\text{O}_2$ $[\text{M}+\text{H}]^+$: 264.99, found 265.01



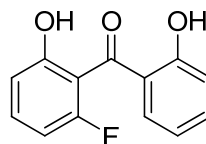
(**10b**)

Bis(4-chloro-2-hydroxyphenyl)methanone (**10b**)

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 10.65 (s, 2H), 7.50 (d, $J = 8.60$ Hz, 2H), 7.11 (s, 2H), 6.93 (d, $J = 8.56$ Hz, 2H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 200.6, 162.5, 142.3, 133.5, 119.8, 118.9, 118.3; HRMS (ESI) calcd for $\text{C}_{13}\text{H}_9\text{Cl}_2\text{O}_3$ $[\text{M}+\text{H}]^+$: 282.9923, found 282.9923



(**11a**)



(**11b**)

(2-Fluorophenyl)(2-hydroxyphenyl)methanone (**11a**)

(2-Fluoro-6-hydroxyphenyl)(2-hydroxyphenyl)methanone (**11b**)

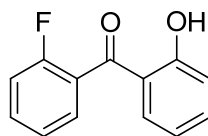
1) $[\text{Ru}(\text{p-cymene})\text{Cl}_2]_2$

Following the general procedure I , 2-fluorobenzophenone (60 mg, 0.30 mmol), $\text{PhI}(\text{OAc})_2$ (200 mg, 0.60 mmol), $[\text{Ru}(\text{p-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 2h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound (**11a**) (20 mg, yellow solid) and compound (**11b**) (40 mg, yellow solid) were isolated in 30% yield and 57% yield respectively.

2) $\text{Rh}_2(\text{OAc})_4$

0.1mmol of 2-fluorobenzophenone , 5% $\text{Rh}_2(\text{OAc})_4$, 1.2eq. of Selectfluor, 1.8ml TFA

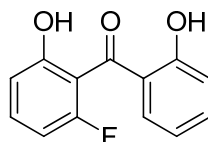
and 0.2ml TFAA were used. The reaction mixture was stirred at 80 °C for 10h. The conversion ratio for compound (**11a**) and compound (**11b**) were 10% and 42% respectively.



(**11a**)

(2-Fluorophenyl)(2-hydroxyphenyl)methanone (**11a**)

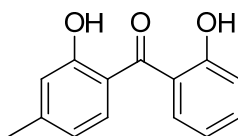
¹H-NMR (400 MHz, CDCl₃) δ (ppm) 11.96 (s, 1H), 7.54-7.45 (m, 3H), 7.41 (d, *J* = 8.0 Hz, 1H), 7.28 (t, *J* = 7.5 Hz, 1H), 7.19 (t, *J* = 9.0 Hz, 1H), 7.06 (d, *J* = 8.4 Hz, 1H), 6.86 (t, *J* = 7.5 Hz, 1H); ¹³C-NMR (100 MHz, CDCl₃) δ (ppm) 198.7, 163.2, 159.2 (d, *J* = 250 Hz, 1C), 137.2, 133.5 (d, *J* = 2.1 Hz, 1C), 133.0 (d, *J* = 8 Hz, 1C), 130.0 (d, *J* = 2.8 Hz, 1C), 126.5 (d, *J* = 15 Hz, 1C), 124.5 (d, *J* = 3.6 Hz, 1C), 119.9, 119.2, 118.4, 116.4 (d, *J* = 21 Hz, 1C); LRMS (ESI) calcd for C₁₃H₁₀FO₂ [M+H]⁺: 217.06, found 216.93



(**11b**)

(2-Fluoro-6-hydroxyphenyl)(2-hydroxyphenyl)methanone (**11b**)

¹H-NMR (400 MHz, CDCl₃) δ (ppm) 11.23 (s, 1H), 8.62 (s, 1H), 7.54-7.48 (m, 2H), 7.45-7.39 (m, 1H), 7.05 (d, *J* = 8.4 Hz, 1H), 6.92-6.87 (m, 2H), 6.71 (t, *J* = 8.8 Hz, 1H); ¹³C-NMR (100 MHz, CDCl₃) δ (ppm) 199.3, 162.6, 160.2 (d, *J* = 252 Hz, 1C), 159.3 (d, *J* = 4.5 Hz, 1C), 137.4, 135.0 (d, *J* = 11 Hz, 1C), 133.2 (d, *J* = 7.1 Hz, 1C), 120.8, 119.3, 118.4, 114.0 (d, *J* = 3 Hz, 1C), 112.0 (d, *J* = 17 Hz, 1C), 107.5 (d, *J* = 23 Hz, 1C); HRMS (ESI) calcd for C₁₃H₁₀FO₃ [M+H]⁺: 233.0608, found 233.0609

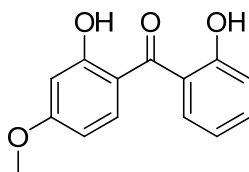


(**12**)

(2-Hydroxy-4-methylphenyl)(2-hydroxyphenyl)methanone (**12**)

Following the general procedure I, 4-methylbenzophenone (59 mg, 0.30 mmol), PhI(OAc)₂ (200 mg, 0.60 mmol), [Ru(*p*-cymene)Cl₂]₂ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 4h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound (**12**) (41 mg, yellow solid) was isolated in 65% yield. ¹H-NMR (400 MHz, CDCl₃) δ (ppm) 10.84 (s, 1H), 10.50 (s, 1H), 7.61 (d, *J* = 8.00 Hz, 1H), 7.53 – 7.48 (m, 2H), 7.08 (d, *J* =

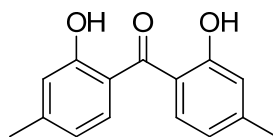
8.36 Hz, 1H), 6.95 – 6.90 (m, 2H), 6.74 (d, $J = 8.12$ Hz, 1H), 2.39 (s, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 202.0, 162.4, 161.6, 148.0, 135.7, 133.2, 133.0, 120.3, 120.2, 118.92, 118.88, 118.7, 117.5, 22.1; LRMS (ESI) calcd for $\text{C}_{14}\text{H}_{13}\text{O}_3$ $[\text{M}+\text{H}]^+$: 229.08, found 228.94



(13)

(2-Hydroxy-4-methoxyphenyl)(2-hydroxyphenyl)methanone (13)

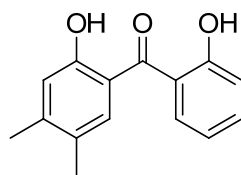
Following the general procedure I , (2-hydroxy-4-methoxyphenyl)(phenyl)methanone (69 mg, 0.30 mmol), $\text{K}_2\text{S}_2\text{O}_8$ (162 mg, 0.20 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.0ml TFA and 1.0 ml TFAA were used. The reaction mixture was stirred at 80 °C for 20h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound **(13)** (37 mg, pale yellow solid) was isolated in 51% yield ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 11.63 (s, 1H), 10.28 (s, 1H), 7.60 – 7.55 (m, 2H), 7.48 (t, $J = 7.32$ Hz, 1H), 7.07 (d, $J = 8.32$ Hz, 1H), 6.93 (t, $J = 7.60$ Hz, 1H), 6.53 (d, $J = 2.40$ Hz, 1H), 6.47 (dd, $J = 8.96$ Hz, $J = 2.44$ Hz, 1H), 3.88 (s, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 200.6, 166.4, 165.8, 161.0, 135.3, 135.2, 132.6, 120.5, 118.9, 118.6, 113.4, 107.7, 101.6, 55.8; LRMS (ESI) calcd for $\text{C}_{14}\text{H}_{13}\text{O}_4$ $[\text{M}+\text{H}]^+$: 245.08, found 245.11



(14)

Bis(2-hydroxy-4-methylphenyl)methanone (14)

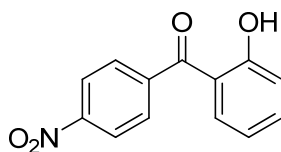
Following the general procedure I , 4,4'-dimethylbenzophenone (63 mg, 0.30 mmol), $\text{PhI}(\text{OAc})_2$ (300 mg, 0.90 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 12h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound **(14)** (45 mg, yellow solid) was isolated in 62% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 10.75 (s, 2H), 7.49 (d, $J = 8.12$ Hz, 2H), 6.87 (s, 2H), 6.73 (d, $J = 8.12$ Hz, 2H), 2.37 (s, 6H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 201.6, 162.1, 147.5, 133.0, 120.2, 118.8, 117.7, 22.0; LRMS (ESI) calcd for $\text{C}_{15}\text{H}_{15}\text{O}_3$ $[\text{M}+\text{H}]^+$: 243.09, found 242.93



(15)

(2-Hydroxy-4,5-dimethylphenyl)(2-hydroxyphenyl)methanone (15)

Following the general procedure I , 3,4-dimethylbenzophenone (42 mg, 0.20 mmol), $\text{PhI}(\text{OAc})_2$ (129 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 3h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound **(15)** (30 mg, yellow solid) was isolated in 61% yield. $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 10.62 (s, 1H), 10.54 (s, 1H), 7.62 (d, $J = 7.92$ Hz, 1H), 7.50 (t, $J = 7.68$ Hz, 1H), 7.36 (s, 1H), 7.08 (d, $J = 8.28$ Hz, 1H), 6.94 (t, $J = 7.56$ Hz, 1H), 6.89 (s, 1H), 2.31 (s, 3H), 2.21 (s, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 202.0, 161.6, 160.5, 146.8, 135.6, 133.4, 133.0, 127.4, 120.3, 119.4, 118.9, 118.6, 117.7, 20.6, 19.1; HRMS (ESI) calcd for $\text{C}_{15}\text{H}_{15}\text{O}_3$ $[\text{M}+\text{H}]^+$: 243.1016, found 243.1019



(16)

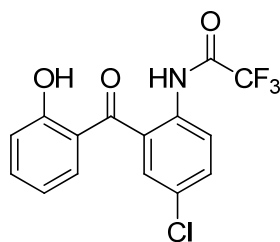
(2-Hydroxyphenyl)(4-nitrophenyl)methanone (16)

1) $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$

Following the general procedure I , 4-nitrobenzophenone (68 mg, 0.30 mmol), $\text{PhI}(\text{OAc})_2$ (200 mg, 0.60 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 80 °C for 14h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 1:1). Finally, compound **(16)** (61 mg, pale yellow solid) was isolated in 77% yield.

2) $\text{Rh}_2(\text{OAc})_4$

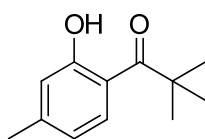
0.1mmol of 2-fluorobenzophenone , 5% $\text{Rh}_2(\text{OAc})_4$, 1.2eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture were stirred at 80 °C for 10h. The conversion ratio for compound **(16)** was 55%. $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 11.77 (s, 1H), 8.37 (d, $J = 8.68$ Hz, 2H), 7.83 (d, $J = 8.68$ Hz, 2H), 7.57 (td, $J = 7.80$ Hz, $J = 1.48$ Hz, 1H), 7.45 (dd, $J = 8.00$ Hz, $J = 1.44$ Hz, 1H), 7.11 (d, $J = 8.36$ Hz, 1H), 6.91 (td, $J = 7.60$ Hz, $J = 0.68$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 199.7, 163.6, 149.7, 143.4, 137.5, 133.2, 130.0, 123.8, 119.3, 119.0, 118.7; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{10}\text{NO}_4$ $[\text{M}+\text{H}]^+$: 244.06, found 243.95



(17)

N-(4-chloro-2-(2-hydroxybenzoyl)phenyl)-2,2,2-trifluoroacetamide (17)

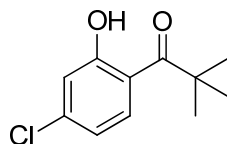
Following the general procedure I , 2-amino-5-chlorobenzophenone (46 mg, 0.20 mmol), $K_2S_2O_8$ (108 mg, 0.40 mmol), $[Ru(p\text{-cymene})Cl_2]_2$ (4 mg, 0.007 mmol), 1.0ml TFA and 1.0 ml TFAA were used. The reaction mixture was stirred at 80 °C for 16-23h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1 or Petroleum ether: Ethylacetate = 10:1). Finally, compound (17) (56 mg, yellow solid) was isolated in 81% yield. 1H -NMR (400 MHz, $CDCl_3$) δ (ppm) 11.15 (s, 1H), 10.41 (br s, 1H), 8.41 (d, J = 8.8 Hz, 1H), 7.62-7.57 (m, 3H), 7.50 (d, J = 8.0 Hz, 1H), 7.11 (d, J = 8.4 Hz, 1H), 6.96 (t, J = 7.6 Hz, 1H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ (ppm) 200.3, 163.7, 155.3 (q, J = 38 Hz), 137.9, 134.4, 133.5, 133.4, 132.0, 130.4, 127.4, 124.2, 119.5, 119.2, 115.7 (q, J = 286 Hz); HRMS (ESI) calcd for $C_{15}H_8ClF_3NO_3$ $[M+H]^+$: 344.0296, found 342.0296.



(18)

1-(2-Hydroxy-4-methylphenyl)-2,2-dimethylpropan-1-one (18)

Following the general procedure I , 2,2-dimethyl-1-p-tolylpropan-1-one (54 mg, 0.10 mmol), $PhI(OAc)_2$ (200 mg, 0.60 mmol), $[Ru(p\text{-cymene})Cl_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 60 °C for 4h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (18)(35 mg, yellow oil) was isolated in 60% yield. 1H -NMR (400 MHz, $CDCl_3$) δ (ppm) 12.78 (s, 1H), 7.90 (d, J = 8.28 Hz, 1H), 6.82 (s, 1H), 6.66 (d, J = 8.20 Hz, 1H), 2.33 (s, 3H), 1.44 (s, 9H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ (ppm) 211.7, 164.0, 146.9, 130.9, 119.4, 119.2, 115.3, 44.6, 28.9, 21.9; LRMS (ESI) calcd for $C_{12}H_{17}O_2$ $[M+H]^+$: 193.12, found 193.02



(19)

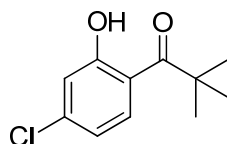
1-(4-Chloro-2-hydroxyphenyl)-2,2-dimethylpropan-1-one (19)

1)[Ru(*p*-cymene)Cl₂]₂

Following the general procedure I , 1-(4-chlorophenyl)-2,2-dimethylpropan-1-one (40 mg, 0.20 mmol), PhI(OAc)₂ (133 mg, 0.40 mmol), [Ru(*p*-cymene)Cl₂]₂ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 50 °C for 4h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (19) (22 mg, pale yellow solid) was isolated in 51% yield.

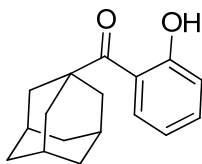
2) Rh₂(OAc)₄

0.1mmol of 1-(4-chlorophenyl)-2,2-dimethylpropan-1-one ,5% Rh₂(OAc)₄, 2.0eq. of Selectfluor, 1.8ml TFA and 0.2ml TFAA were used. The reaction mixture were stirred at 110 °C for 10h. The conversion ratio for compound (19) was 24%.



(19)

¹H-NMR (400 MHz, CDCl₃) δ (ppm) 12.86 (s, 1H), 7.94 (d, *J* = 8.80 Hz, 1H), 7.02 (d, *J* = 2.08 Hz, 1H), 6.82 (dd, *J* = 8.80 Hz, *J* = 2.04 Hz, 1H), 1.44 (s, 9H); ¹³C-NMR (100 MHz, CDCl₃) δ (ppm) 211.6, 164.7, 141.1, 132.0, 119.4, 118.6, 116.1, 44.8, 28.8; LRMS (ESI) calcd for C₁₁H₁₄ClO₂ [M+H]⁺: 213.07, found 213.03

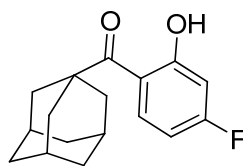


(20)

Adamantan-1-yl-(2-hydroxy-phenyl)-methanone (20)

Following the general procedure I , adamantan-1-yl-phenyl-methanone (48 mg, 0.20 mmol), PhI(OAc)₂ (133 mg, 0.40 mmol), [Ru(*p*-cymene)Cl₂]₂ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 50 °C for 4h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (20) (34 mg, pale yellow solid) was isolated in 67% yield. ¹H-NMR (400 MHz, CDCl₃) δ (ppm) 12.76 (s, 1H), 8.21 (dd, *J* = 8.24 Hz, *J* = 1.16 Hz, 1H), 7.40 (td, *J* = 7.78 Hz, *J* = 1.24 Hz, 1H), 7.00 (d, *J* = 8.28 Hz, 1H), 6.84 (t, *J* = 7.96 Hz, 1H), 2.17 – 2.13 (m, 9H), 1.81 (s, 6H); ¹³C-NMR (100 MHz, CDCl₃) δ (ppm) 211.7, 163.6, 135.3, 130.7, 119.5, 118.2, 117.7, 48.0, 39.9, 36.8, 28.5; LRMS (ESI) calcd for C₁₇H₂₁O₂ [M+H]⁺: 257.15,

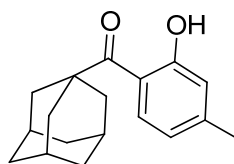
found 257.12



(21)

Adamantan-1-yl-(4-fluorophenyl-2-hydroxy-phenyl)-methanone (21)

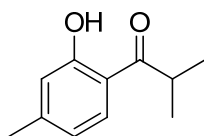
Following the general procedure I , adamantan-1-yl-(4-fluorophenyl)-methanone (77 mg, 0.30 mmol), PhI(OAc)₂ (133 mg, 0.40 mmol), [Ru(*p*-cymene)Cl₂]₂ (3.1 mg, 0.005 mmol), 0.8ml TFA and 1.2ml TFAA were used. The reaction mixture was stirred at 50 °C for 6h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1 or Petroleum ether: Ethylacetate = 100:1). Finally, compound (21) (32 mg, pale yellow solid) was isolated in 59% yield. ¹H-NMR (400 MHz, CDCl₃) δ (ppm) 13.19 (s, 1H), 8.23 (dd, *J* = 9.16 Hz, *J* = 2.56 Hz, 1H), 6.66 (dd, *J* = 10.40 Hz, *J* = 2.60 Hz, 1H), 6.55 (td, *J* = 8.50 Hz, *J* = 2.60 Hz, 1H), 2.14 (s, 9H), 1.80 (m, 6H); ¹³C-NMR (100 MHz, CDCl₃) δ (ppm) 210.5, 166.4 (d, *J* = 254 Hz, 1C), 166.6 (d, *J* = 14 Hz, 1C), 133.0 (d, *J* = 11 Hz, 1C), 115.1 (d, *J* = 2.4 Hz, 1C), 105.9 (d, *J* = 22 Hz, 1C), 105.71 (d, *J* = 15 Hz, 1C), 47.9, 39.9, 36.7, 28.4; LRMS (ESI) calcd for C₁₇H₂₀FO₂ [M+H]⁺: 275.14, found 275.08



(22)

Adamantan-1-yl-(4-methyl-2-hydroxy-phenyl)-methanone (22)

Following the general procedure I , adamantan-1-yl-(4-methyl-phenyl)-methanone (51 mg, 0.20 mmol), PhI(OAc)₂ (133 mg, 0.40 mmol), [Ru(*p*-cymene)Cl₂]₂ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 50 °C for 5.5h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (22) (31 mg, white solid) was isolated in 56% yield. ¹H-NMR (400 MHz, CDCl₃) δ (ppm) 12.88 (s, 1H), 8.09 (d, *J* = 8.40 Hz, 1H), 6.81 (s, 1H), 6.65 (d, *J* = 8.36 Hz, 1H), 2.33 (s, 3H), 2.16 – 2.13 (m, 9H), 1.81 (s, 6H); ¹³C-NMR (100 MHz, CDCl₃) δ (ppm) 211.1, 164.0, 146.7, 130.5, 119.5, 119.0, 115.8, 47.9, 39.9, 36.8, 28.5, 21.9; LRMS (ESI) calcd for C₁₈H₂₃O₂ [M+H]⁺: 271.17, found 271.10

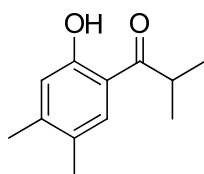


(23)

1-(2-Hydroxy-4-methylphenyl)-2-methylpropan-1-one (23)

Following the general procedure I , 2-methyl-1-p-tolylpropan-1-one (33 mg, 0.20

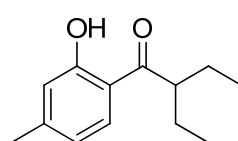
mmol), $\text{PhI}(\text{OAc})_2$ (133 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 50 °C for 5.5h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (**23**) (15 mg, colourless oil) was isolated in 40% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.55 (s, 1H), 7.66 (d, J = 8.20 Hz, 1H), 6.80 (s, 1H), 6.71 (d, J = 8.20 Hz, 1H), 3.62 – 3.52 (m, 1H), 2.35 (s, 3H), 1.23 (d, J = 6.84 Hz, 6H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 210.3, 163.4, 147.9, 129.9, 120.2, 118.9, 116.1, 34.9, 22.0, 19.5; LRMS (ESI) calcd for $\text{C}_{11}\text{H}_{15}\text{O}_2$ $[\text{M}+\text{H}]^+$: 179.11, found 179.09



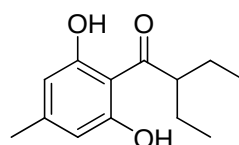
(**24**)

1-(2-Hydroxy-4,5-dimethylphenyl)-2-methylpropan-1-one (**24**)

Following the general procedure I , 1-(3,4-dimethylphenyl)-2-methylpropan-1-one (35 mg, 0.20 mmol), $\text{PhI}(\text{OAc})_2$ (133 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 50 °C for 5.5h.. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (**24**) (20 mg, colourless oil) was isolated in 52% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.37 (s, 1H), 7.50 (s, 1H), 6.79 (s, 1H), 3.62 – 3.55 (m, 1H), 2.26 (s, 3H), 2.22 (s, 3H), 1.23 (d, J = 6.84 Hz, 6H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 210.3, 161.6, 146.8, 130.1, 127.1, 119.4, 116.2, 34.8, 20.6, 19.5, 19.2; LRMS (ESI) calcd for $\text{C}_{12}\text{H}_{17}\text{O}_2$ $[\text{M}+\text{H}]^+$: 193.12, found 193.10



(**25a**)

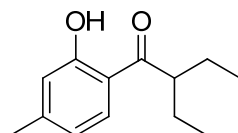


(**25b**)

2-Ethyl-1-(2-hydroxy-4-methylphenyl)butan-1-one (**25a**)

1-(2,6-Dihydroxy-4-methylphenyl)-2-ethylbutan-1-one (**25b**)

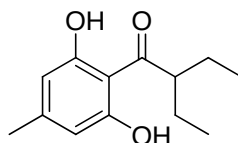
Following the general procedure I , 2-ethyl-1-p-tolylbutan-1-one (19 mg, 0.10 mmol), $\text{PhI}(\text{OAc})_2$ (65 mg, 0.20 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.5ml TFA and 0.5ml TFAA were used. The reaction mixture was stirred at 50 °C for 12h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1 or Petroleum ether: Ethylacetate = 100:1). Finally, compound (**25a**) (4 mg, colourless oil) and compound (**25b**) (12mg, white solid) were isolated in 13% and 65% yield respectively.



(25a)

2-Ethyl-1-(2-hydroxy-4-methylphenyl)butan-1-one (25a)

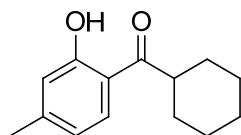
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 12.76 (s, 1H), 7.68 (d, $J = 8.24$ Hz, 1H), 6.80 (s, 1H), 6.71 (d, $J = 8.20$ Hz, 1H), 3.30 – 3.20 (m, 1H), 2.35 (s, 3H), 1.83 – 1.76 (m, 2H), 1.62 – 1.55 (m, 2H), 0.88 (t, $J = 7.44$ Hz, 6H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 210.4, 163.3, 148.0, 130.0, 120.2, 118.8, 117.9, 48.5, 25.4, 22.0, 12.0; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_{19}\text{O}_2$ $[\text{M}+\text{H}]^+$: 207.12, found 207.17



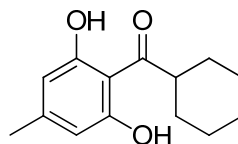
(25b)

1-(2,6-Dihydroxy-4-methylphenyl)-2-ethylbutan-1-one (25b)

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 9.59 (br s, 2H), 6.21 (s, 2H), 3.75 (m, 1H), 2.23 (s, 3H), 1.85-1.72 (m, 2H), 1.57-1.43 (m, 2H), 0.89 (t, $J = 7.20$ Hz, 6H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 211.4, 161.2, 147.5, 109.5, 108.7, 53.2, 24.7, 21.9, 12.0; HRMS (ESI) calcd for $\text{C}_{13}\text{H}_{19}\text{O}_3$ $[\text{M}+\text{H}]^+$: 223.1329, found 223.1332



(26a)



(26b)

Cyclohexyl(2-hydroxy-4-methylphenyl)methanone (26a)

Cyclohexyl(2,6-dihydroxy-4-methylphenyl)methanone (26b)

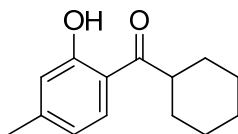
1) small-scale

Following the general procedure I, cyclohexyl(2-hydroxy-4-methylphenyl)Methanone (21 mg, 0.10 mmol), $\text{PhI}(\text{OAc})_2$ (65 mg, 0.20 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.5ml TFA and 0.5ml TFAA were used. The reaction mixture was stirred at 50 °C for 11h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1 or Petroleum ether: Ethylacetate = 100:1). Finally, compound (26a) (3.3 mg, yellow oil) and compound (26b) (13 mg, white solid) were isolated in 15% and 56% yield respectively.

2) Two gram-scale

Following the general procedure I, cyclohexyl(2-hydroxy-4-methylphenyl)methanone (2 g, 9.9 mmol), $\text{PhI}(\text{OAc})_2$ (6.4 g, 19.3 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (121 mg, 0.20 mmol), 21ml TFA and 7ml TFAA were used. The reaction mixture was stirred at 50 °C for 14h. The residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1 or Petroleum ether: Ethylacetate = 100:1). Finally,

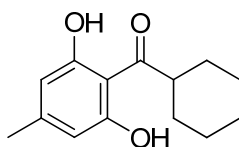
compound (**26a**) (1080 mg, yellow oil) and compound (**26b**) (139 mg, white solid) were isolated in 50% and 6% yield respectively.



(**26a**)

Cyclohexyl(2-hydroxy-4-methylphenyl)methanone (**26a**)

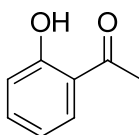
^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.61 (s, 1H), 7.65 (d, J = 8.20 Hz, 1H), 6.79 (s, 1H), 6.70 (d, J = 8.20 Hz, 1H), 3.29 – 3.23 (m, 1H), 2.34 (s, 3H), 1.88 – 1.34 (m, 10H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 209.6, 163.4, 147.8, 129.8, 129.1, 118.9, 116.2, 45.2, 29.7, 26.0, 25.9, 22.0; LRMS (ESI) calcd for $\text{C}_{14}\text{H}_{19}\text{O}_2$ $[\text{M}+\text{H}]^+$: 219.14, found 219.09



(**26b**)

Cyclohexyl(2,6-dihydroxy-4-methylphenyl)methanone (**26b**)

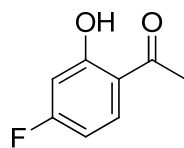
^1H -NMR (400 MHz, CDCl_3) δ (ppm) 9.41 (br s, 2H), 6.21 (s, 2H), 3.60 (m, 1H), 2.23 (s, 3H), 1.96-1.93 (m, 2H), 1.84-1.81 (m, 2H), 1.73-1.70 (m, 1H), 1.47-1.23 (m, 5H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 210.6, 161.2, 147.3, 109.5, 107.5, 50.3, 29.6, 26.3, 21.9; HRMS (ESI) calcd for $\text{C}_{14}\text{H}_{19}\text{O}_3$ $[\text{M}+\text{H}]^+$: 235.1329, found 235.1324



(**27**)

1-(2-Hydroxyphenyl)ethanone (**27**)

Following the general procedure I , acetophenone (36 mg, 0.30 mmol), $\text{PhI}(\text{OAc})_2$ (200 mg, 0.60 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAC were used. The reaction mixture was stirred at 50 °C for 4h. After completion of the reaction, the residue was purified by silica gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (**27**) (13 mg, pale yellow oil) was isolated in 30% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.26 (s, 1H), 7.73 (d, J = 8.00 Hz, 1H), 7.47 (t, J = 8.08 Hz, 1H), 6.98 (d, J = 8.40 Hz, 1H), 6.90 (t, J = 7.76 Hz, 1H), 2.63 (s, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 204.7, 162.5, 136.6, 130.9, 119.9, 119.1, 118.6, 26.8; LRMS (ESI) calcd for $\text{C}_8\text{H}_9\text{O}_2$ $[\text{M}+\text{H}]^+$: 137.06, found 137.00



(28)

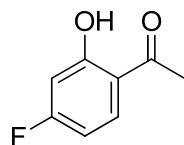
1-(4-Fluoro-2-hydroxyphenyl)ethanone (28)

1) [Ru(*p*-cymene)Cl₂]₂

Following the general procedure I, 4'-Fluoroacetophenone (41 mg, 0.30 mmol), PhI(OAc)₂ (200 mg, 0.60 mmol), [Ru(*p*-cymene)Cl₂]₂ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at rt for 24h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (28) (12 mg, yellow oil) was isolated in 26% yield.

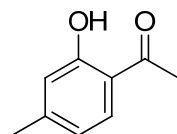
2) Rh₂(OAc)₄

0.1mmol of 4'-Fluoroacetophenone, 5% Rh₂(OAc)₄, 2.0eq. of Selectfluor, 0.9ml TFA and 0.1ml TFAA were used. The reaction mixture were stirred at 75 °C for 11h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (28) was isolated in 41% yield.



(28)

¹H-NMR (400 MHz, CDCl₃) δ (ppm) 12.59 (s, 1H), 7.47 (dd, *J* = 6.60 Hz, *J* = 8.72 Hz, 1H), 6.68 – 6.60 (m, 2H), 2.61 (s, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ (ppm) 203.4, 167.6 (d, *J* = 255 Hz, 1C), 165.1 (d, *J* = 14 Hz, 1C), 133.2 (d, *J* = 12 Hz, 1C), 117.03 (d, *J* = 2.2 Hz, 1C), 107.03 (d, *J* = 23 Hz, 1C), 105.11 (d, *J* = 22 Hz, 1C), 26.75; LRMS (ESI) calcd for C₈H₈FO₂ [M+H]⁺: 155.05, found 155.08

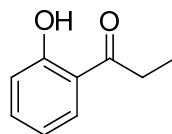


(29)

1-(2-Hydroxy-4-methylphenyl)ethanone (29)

Following the general procedure I, 4'-methylacetophenone (41 mg, 0.30 mmol), PhI(OAc)₂ (200 mg, 0.60 mmol), [Ru(*p*-cymene)Cl₂]₂ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 50 °C for 4h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (29) (25 mg, pale yellow oil) was isolated in 54% yield. ¹H-NMR (400 MHz, CDCl₃) δ (ppm)

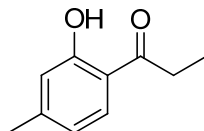
12.28 (s, 1H), 7.61 (d, $J = 8.16$ Hz, 1H), 6.78 (s, 1H), 6.71 (d, $J = 8.16$ Hz, 1H), 2.60 (s, 3H), 2.35 (s, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 204.0, 162.6, 148.2, 130.7, 120.3, 118.5, 117.7, 26.6, 22.1; LRMS (ESI) calcd for $\text{C}_9\text{H}_{11}\text{O}_2$ $[\text{M}+\text{H}]^+$: 151.08, found 151.07



(30)

1-(2-Hydroxyphenyl)propan-1-one (30)

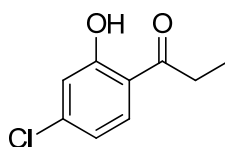
Following the general procedure I , propiophenone (27 mg, 0.20 mmol), Selectfluor (106 mg, 0.30 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.5ml TFA and 0.5ml TFAA were used. The reaction mixture was stirred at 50 °C for 6h and then at 60 °C for 5h . After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (30) (16 mg, colourless oil) was isolated in 54% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.35 (s, 1H), 7.77 (dd, $J = 1.12$ Hz, $J = 8.00$ Hz, 1H), 7.46 (td, $J = 1.32$ Hz, $J = 7.78$ Hz, 1H), 6.98 (d, $J = 8.40$ Hz, 1H), 6.89 (t, $J = 7.96$ Hz, 1H), 3.04 (q, $J = 7.28$ Hz, 2H), 1.25 (t, $J = 7.28$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 207.2, 162.5, 136.3, 130.0, 119.4, 119.0, 118.7, 31.7, 8.4; LRMS (ESI) calcd for $\text{C}_9\text{H}_9\text{O}_2$ $[\text{M}-\text{H}]^-$: 149.07, found 149.08



(31)

1-(2-Hydroxy-4-methylphenyl)propan-1-one (31)

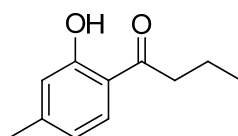
Following the general procedure I , 4'-methylpropiophenone (45 mg, 0.30 mmol), $\text{PhI}(\text{OAc})_2$ (200 mg, 0.60 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (4.5 mg, 0.0074 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 35 °C for 12h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound (31) (25 mg, colourless oil) was isolated in 50% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.37 (s, 1H), 7.64 (d, $J = 8.16$ Hz, 1H), 6.79 (s, 1H), 6.70 (d, $J = 8.12$ Hz, 1H), 3.10 – 2.90 (m, 2H), 2.34 (s, 3H), 1.21 (t, $J = 7.20$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 206.7, 162.7, 147.8, 129.9, 120.3, 118.7, 117.2, 31.5, 22.1, 8.5; LRMS (ESI) calcd for $\text{C}_{10}\text{H}_{13}\text{O}_2$ $[\text{M}+\text{H}]^+$: 165.08, found 164.92



(32)

1-(4-chloro-2-hydroxyphenyl)propan-1-one (32)

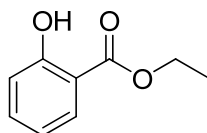
Following the general procedure I , 4'-chloropropiophenone (34 mg, 0.20 mmol), $\text{PhI}(\text{OAc})_2$ (133 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.4ml TFA and 0.6ml TFAA were used. The reaction mixture was stirred at 50 °C for 5.5h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound **(32)** (20 mg, colourless oil) was isolated in 55% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.47 (s, 1H), 7.67 (d, J = 8.60 Hz, 1H), 6.99 (d, J = 2.04 Hz, 1H), 6.86 (dd, J = 8.56 Hz, J = 2.00 Hz, 1H), 2.99 (q, J = 7.28 Hz, 2H), 1.24 (t, J = 7.28 Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 206.5, 163.2, 142.0, 130.9, 119.7, 118.7, 117.9, 31.8, 8.3; LRMS (ESI) calcd for $\text{C}_9\text{H}_8\text{ClO}_2$ $[\text{M}-\text{H}]^-$: 183.03, found 182.89



(33)

1-(2-Hydroxy-4-methylphenyl)butan-1-one (33)

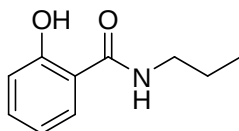
Following the general procedure I , 4'-methylbutyrophenone (33 mg, 0.20 mmol), $\text{PhI}(\text{OAc})_2$ (133 mg, 0.40 mmol), $[\text{Ru}(p\text{-cymene})\text{Cl}_2]_2$ (3.1 mg, 0.005 mmol), 1.5ml TFA and 0.5ml TFAA were used. The reaction mixture was stirred at rt for 21h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 3:1). Finally, compound **(33)** (16 mg, colourless oil) was isolated in 45% yield. ^1H -NMR (400 MHz, CDCl_3) δ (ppm) 12.43 (s, 1H), 7.64 (d, J = 8.16 Hz, 1H), 6.79 (s, 1H), 6.70 (d, J = 8.16 Hz, 1H), 2.93 (t, J = 7.32 Hz, 2H), 2.34 (s, 3H), 1.82 – 1.73 (m, 2H), 1.06 (t, J = 7.40 Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 206.4, 162.8, 147.9, 130.0, 120.2, 118.7, 117.4, 40.2, 22.0, 18.2, 14.0; LRMS (ESI) calcd for $\text{C}_{11}\text{H}_{15}\text{O}_2$ $[\text{M}+\text{H}]^+$: 179.11, found 179.17



Ethyl 2-hydroxybenzoate (34)

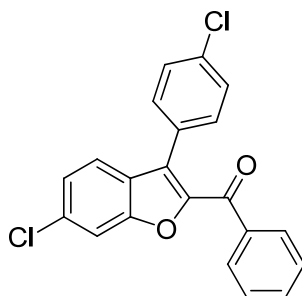
Following the general procedure I , ethyl benzoate (15 mg, 0.10 mmol), $\text{K}_2\text{S}_2\text{O}_8$ (30mg, 0.11 mmol), $\text{Rh}_2(\text{OAc})_4$ (2 mg, 0.005 mmol), 0.9 ml TFA and 0.1 ml TFAA were used. The reaction mixture was stirred at 90 °C for 9h. After completion of the reaction, the residue was purified by silical gel column chromatography (Petroleum ether: Toluene = 5:1). Finally, compound **(34)** (10.4 mg, colourless oil) was isolated

in 59% yield. $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 10.84 (s, 1H), 7.85 (dd, $J = 1.68$ Hz, $J = 8.00$ Hz, 1H), 7.42 - 7.47 (m, 1H), 6.98 (dd, $J = 0.72$ Hz, $J = 8.40$ Hz, 1H), 6.85 – 6.89 (m, 1H), 4.41 (q, $J = 7.16$, 2H), 1.42 (t, $J = 7.16$ Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 170.3, 161.8, 135.7, 130.0, 119.2, 117.7, 112.8, 61.5, 14.32; LRMS (ESI) calcd for $\text{C}_9\text{H}_{11}\text{O}_3$ $[\text{M}+\text{H}]^+$: 167.06, found 166.71



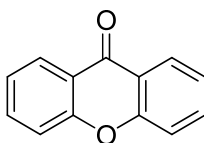
2-Hydroxy-N-propylbenzamide (35)

Following the general procedure I, N-propylbenzamide (17 mg, 0.10 mmol), Selectfluor (56mg, 0.22 mmol), $\text{Rh}_2(\text{OAc})_4$ (2 mg, 0.005 mmol), 0.9 ml TFA and 0.1 ml TFAA were used. The reaction mixture was stirred at 60 °C for 5.5h. $^1\text{H-NMR}$ shows the conversion ratio is 31%. $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 12.41 (s, 1H), 7.40 – 7.34 (m, 2H), 6.98 (d, $J = 8.32$ Hz, 1H), 6.83 (t, $J = 7.56$ Hz, 1H), 6.39 (s, 1H), 3.44 – 3.39 (m, 2H), 1.66 – 1.64 (m, 2H), 0.99 (t, $J = 7.40$ Hz, 1H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 170.1, 161.7, 134.2, 125.3, 118.8, 118.7, 114.5, 41.5, 22.9, 11.5; LRMS (ESI) calcd for $\text{C}_{10}\text{H}_{14}\text{NO}_2$ $[\text{M}+\text{H}]^+$: 180.09, found 179.98



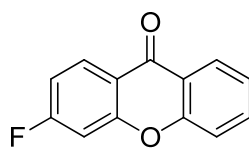
(6-Chloro-3-(4-chlorophenyl)benzofuran-2-yl)(phenyl)methanone (36)

$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 7.89 (d, $J = 7.88$ Hz, 2H), 7.65 (s, 1H), 7.58 – 7.53 (m, 2H), 7.45 – 7.34 (m, 7H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 185.2, 154.6, 147.9, 137.0, 134.9, 134.5, 133.2, 131.3, 130.0, 129.0, 128.9, 128.4, 128.0, 126.7, 125.3, 122.9, 113.0; LRMS (ESI) calcd for $\text{C}_{21}\text{H}_{13}^{35}\text{Cl}_2\text{O}_2$ $[\text{M}+\text{H}]^+$: 367.0287, found 367.0288



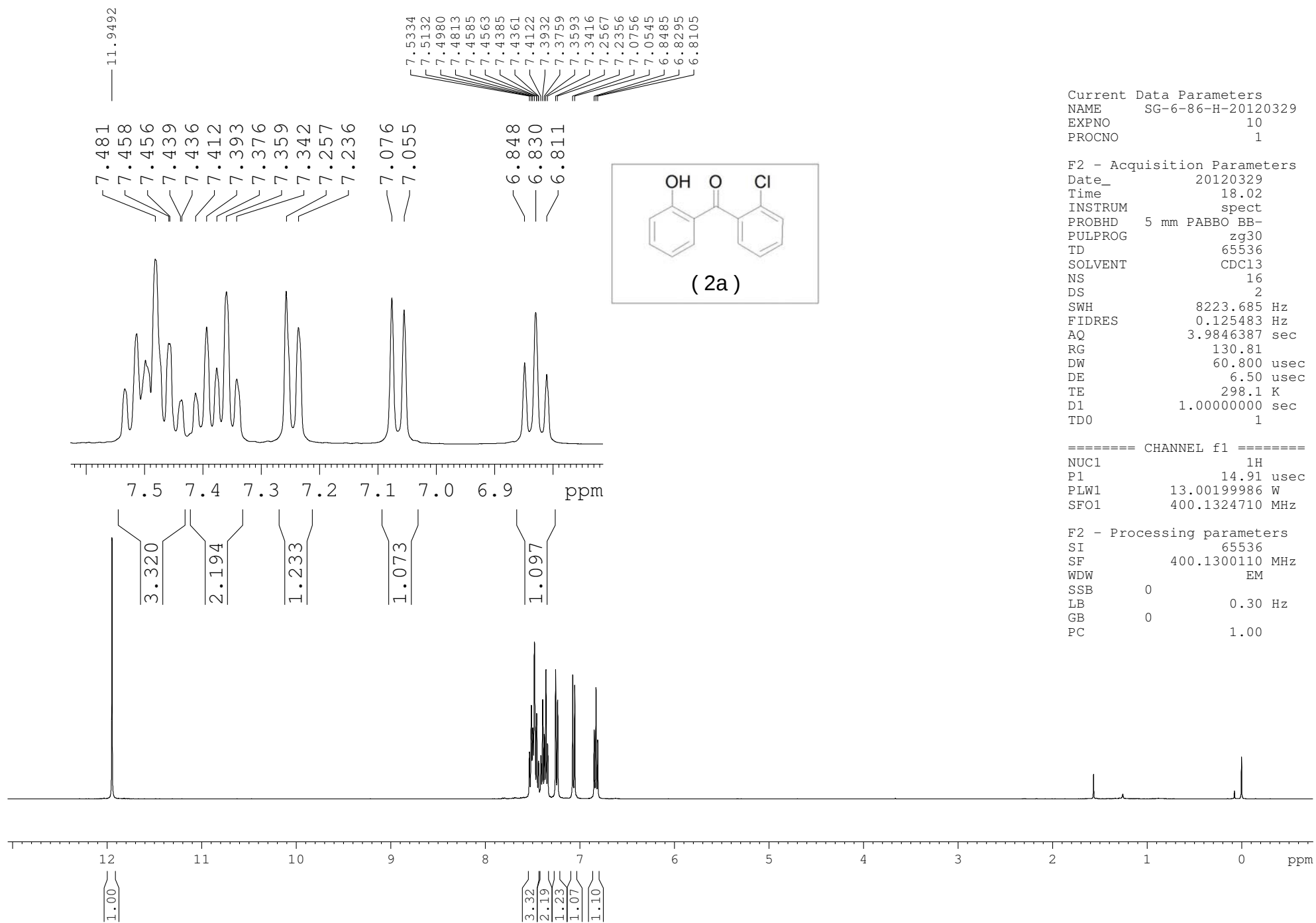
Xanthone (38)

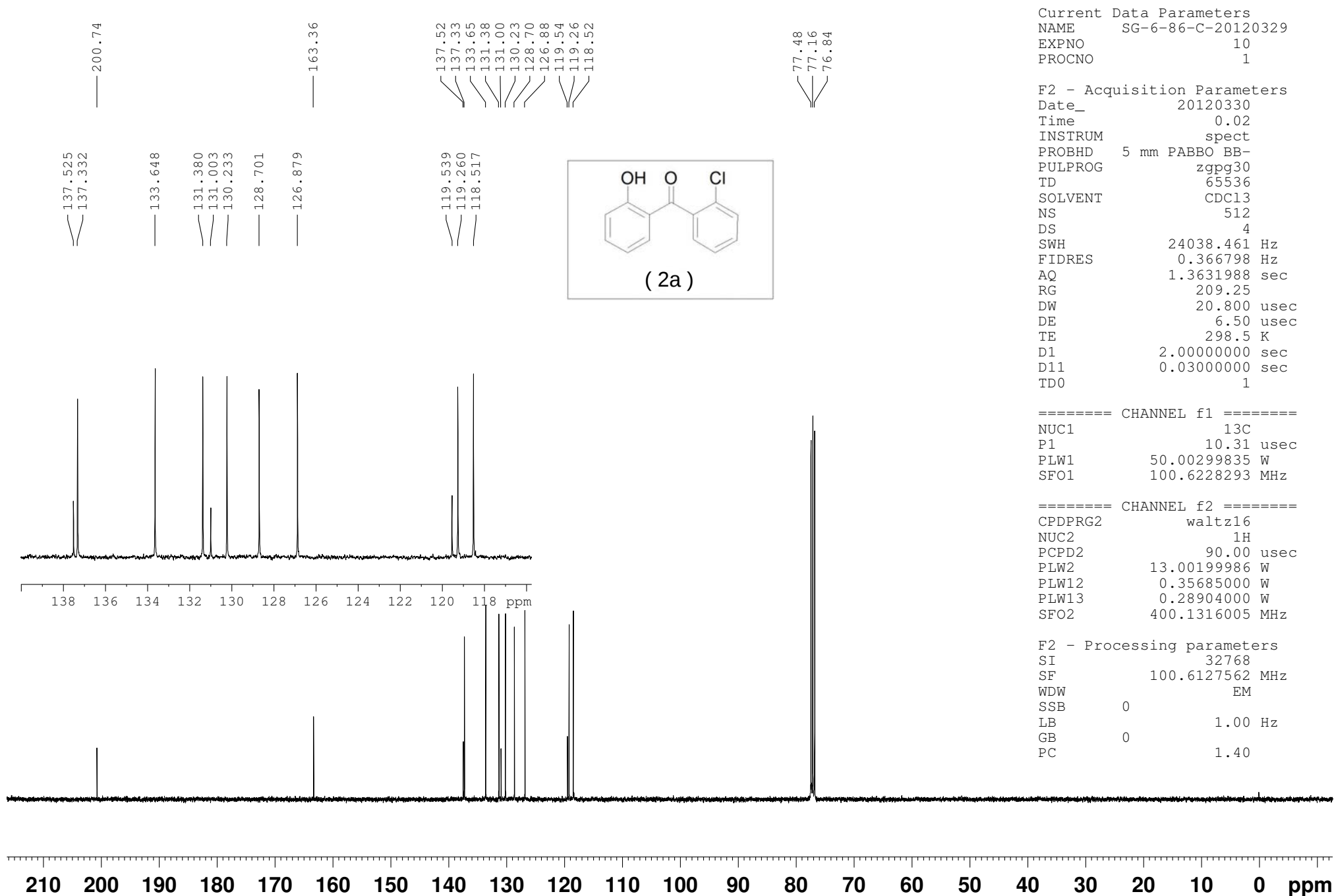
$^1\text{H-NMR}$ (400 MHz, CDCl_3) δ (ppm) 8.35 (d, $J = 7.88$ Hz, 2H), 7.73 (t, $J = 7.88$ Hz, 2H), 7.50 (dd, $J = 2.16$ Hz, $J = 8.36$ Hz, 2H), 7.38 (d, $J = 7.88$ Hz, 2H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ (ppm) 177.4, 156.3, 135.0, 126.9, 124.1, 122.0, 118.1; LRMS (ESI) calcd for $\text{C}_{13}\text{H}_9\text{O}_2$ $[\text{M}+\text{H}]^+$: 197.06, found 197.09



3-Fluoro-9H-xanthen-9-one (39)

^1H -NMR (400 MHz, CDCl_3) δ (ppm) 8.38 – 8.33 (m, 2H), 7.73 (t, $J = 7.44$ Hz, 1H), 7.49 (d, $J = 8.44$ Hz, 1H), 7.40 (t, $J = 7.52$ Hz, 1H), 7.17 (dd, $J = 9.32$ Hz, $J = 1.80$ Hz, 1H), 7.11 (td, $J = 8.46$ Hz, $J = 1.96$ Hz, 1H); ^{13}C -NMR (100 MHz, CDCl_3) δ (ppm) 176.3, 166.7 (d, $J = 254$ Hz, 1C), 157.5 (d, $J = 13$ Hz, 1C), 156.4, 135.04, 129.5 (d, $J = 11$ Hz, 1C), 126.9, 124.5, 121.9, 118.9, 118.0, 112.9 (d, $J = 22$ Hz, 1C), 104.7 (d, $J = 25$ Hz, 1C); LRMS (ESI) calcd for $\text{C}_{13}\text{H}_8\text{FO}_2$ $[\text{M}+\text{H}]^+$: 215.04, found 214.91





Current Data Parameters
NAME HXS-Cl-DI-OH-H
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters

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D1 1.00000000 sec
TD0 1

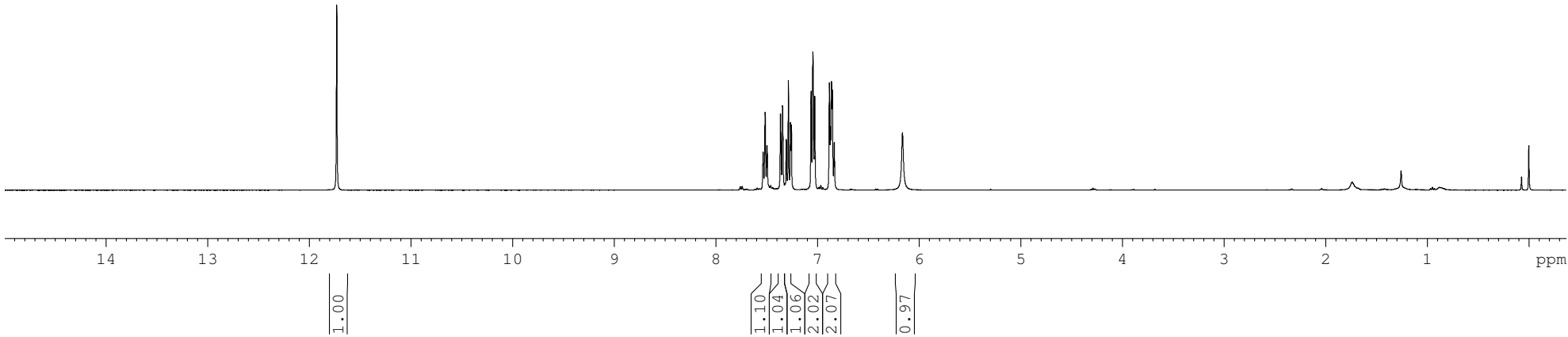
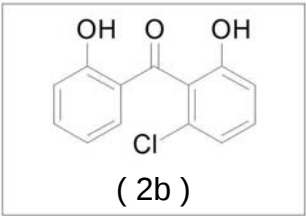
===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters

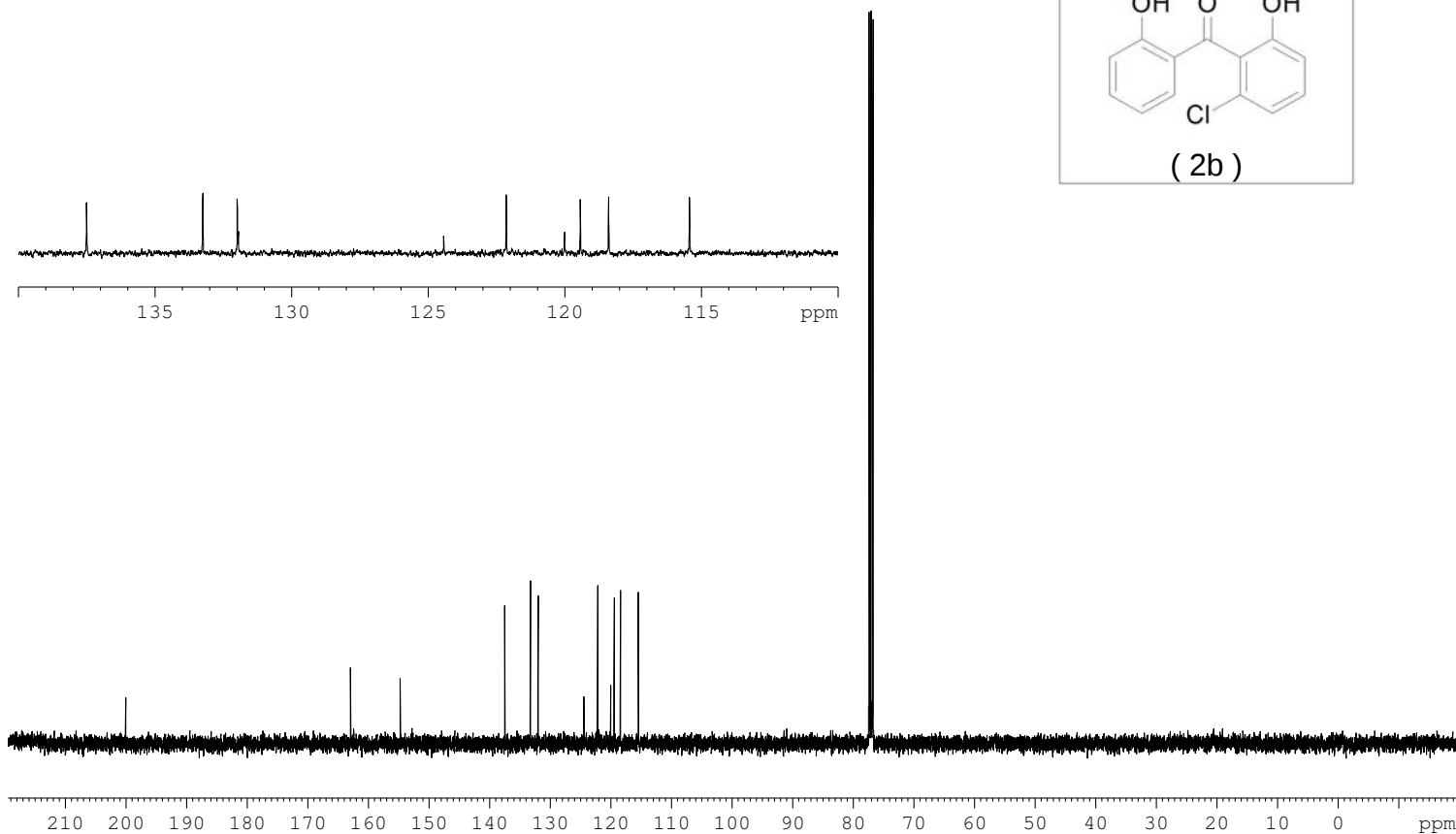
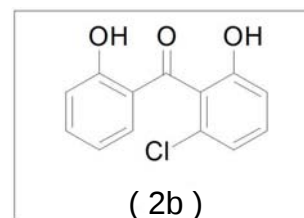
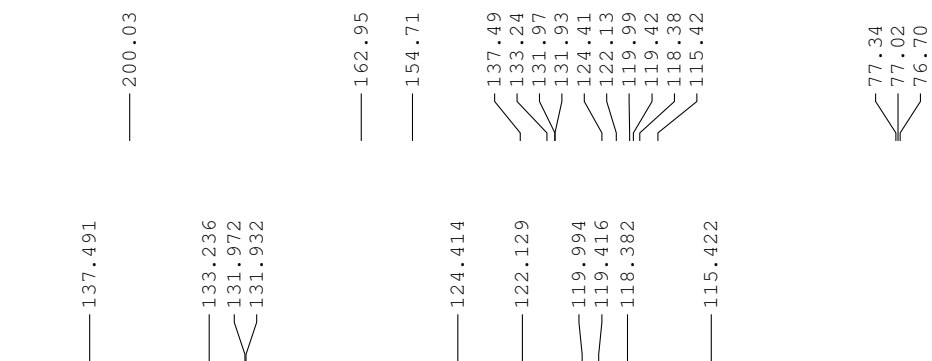
SI 65536
SF 400.1300116 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

11.7305

7.5353
7.5149
7.4965
7.3635
7.3435
7.3062
7.2858
7.2653
7.2564
7.0619
7.0450
7.0265
6.8828
6.8724
6.8619
6.8534
6.8344
6.1632



C13CPD CDC13 {D:\NMR_DATA} RY 10



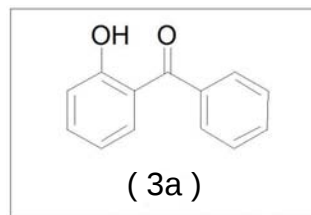
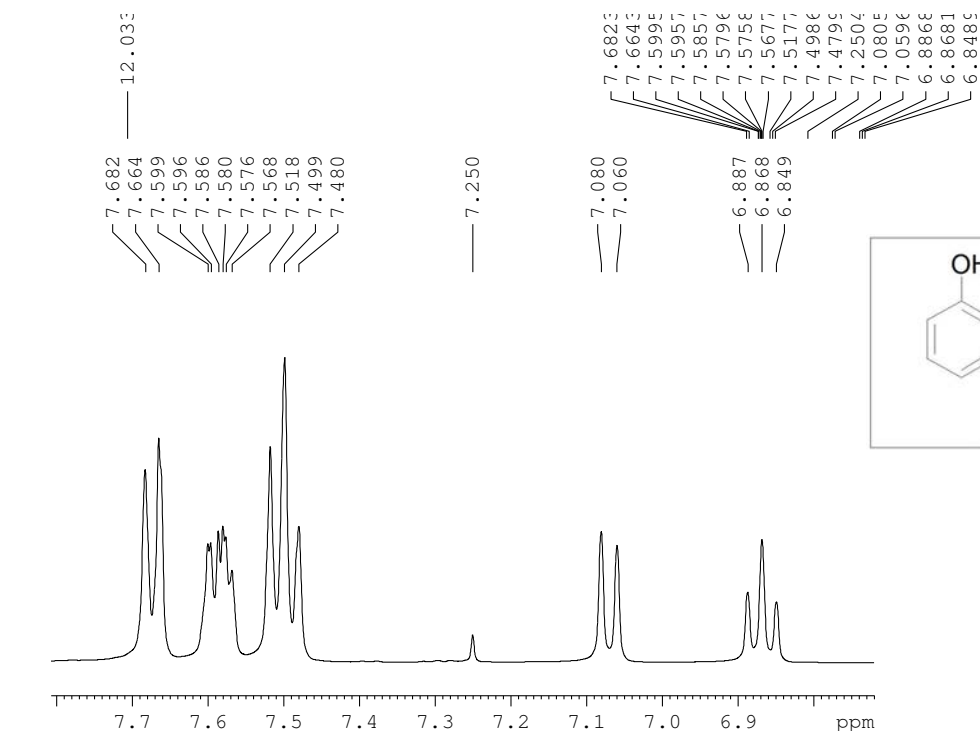
Current Data Parameters
NAME HXS-Cl-DI-OH-C
EXPNO 20
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121120
Time 14.25
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 86
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.0000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127690 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

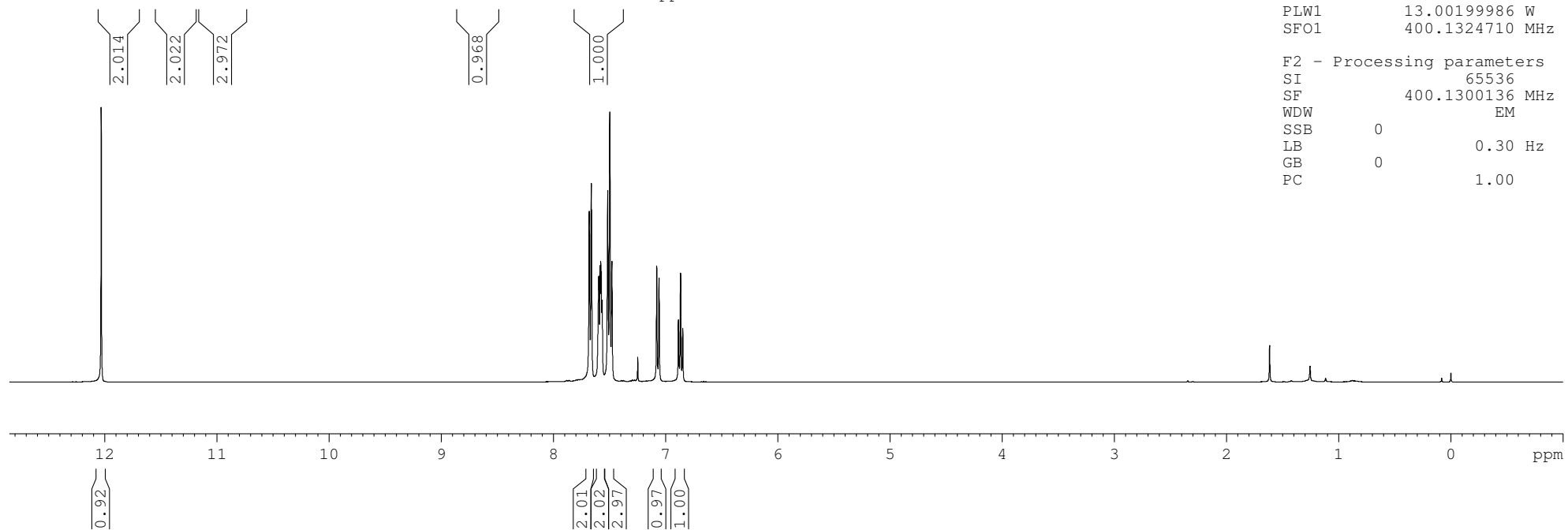


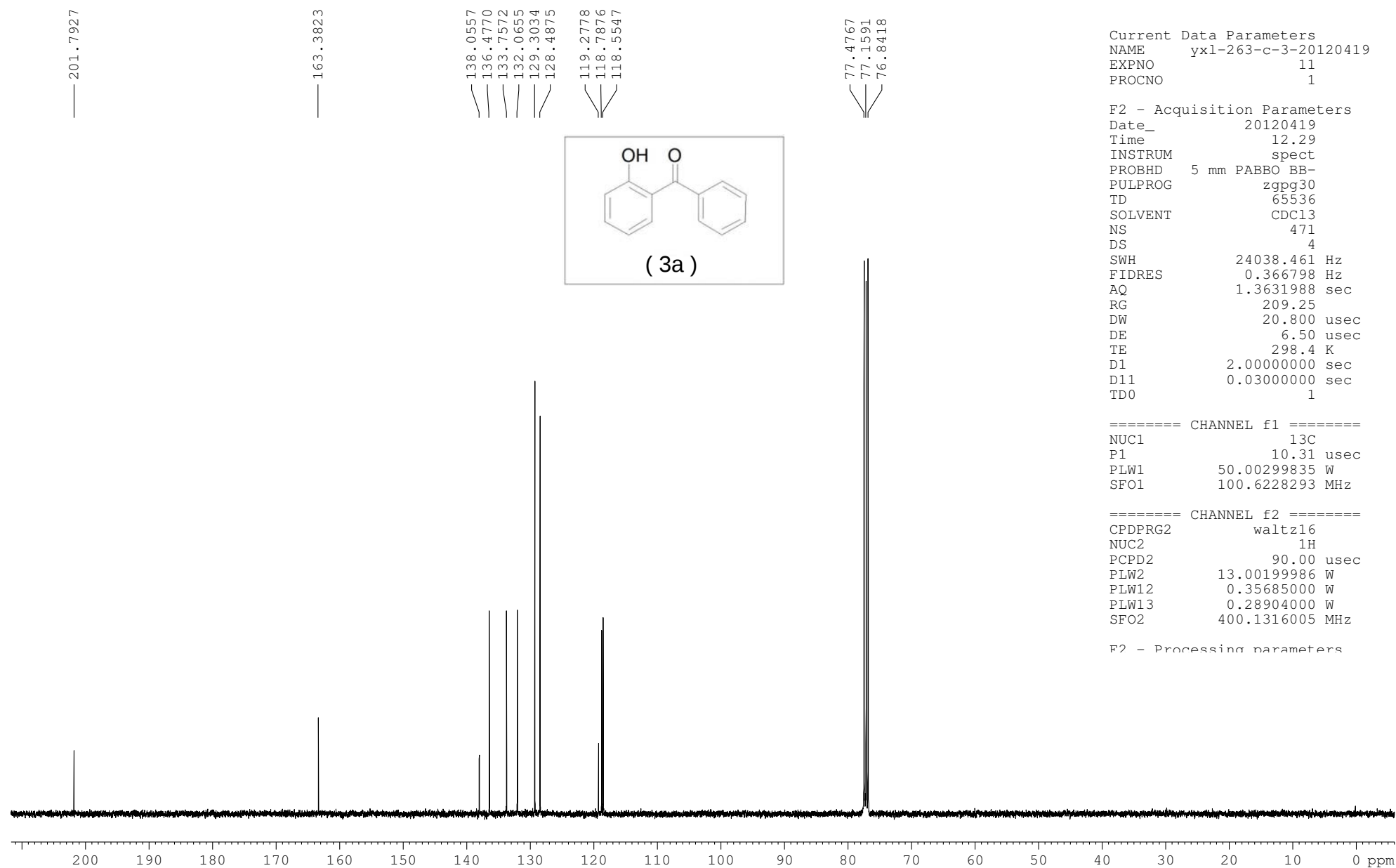
Current Data Parameters
NAME YXL-263-H-C-20120418
EXPNO 30
PROCNO 1

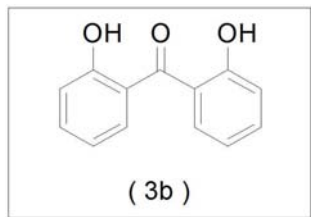
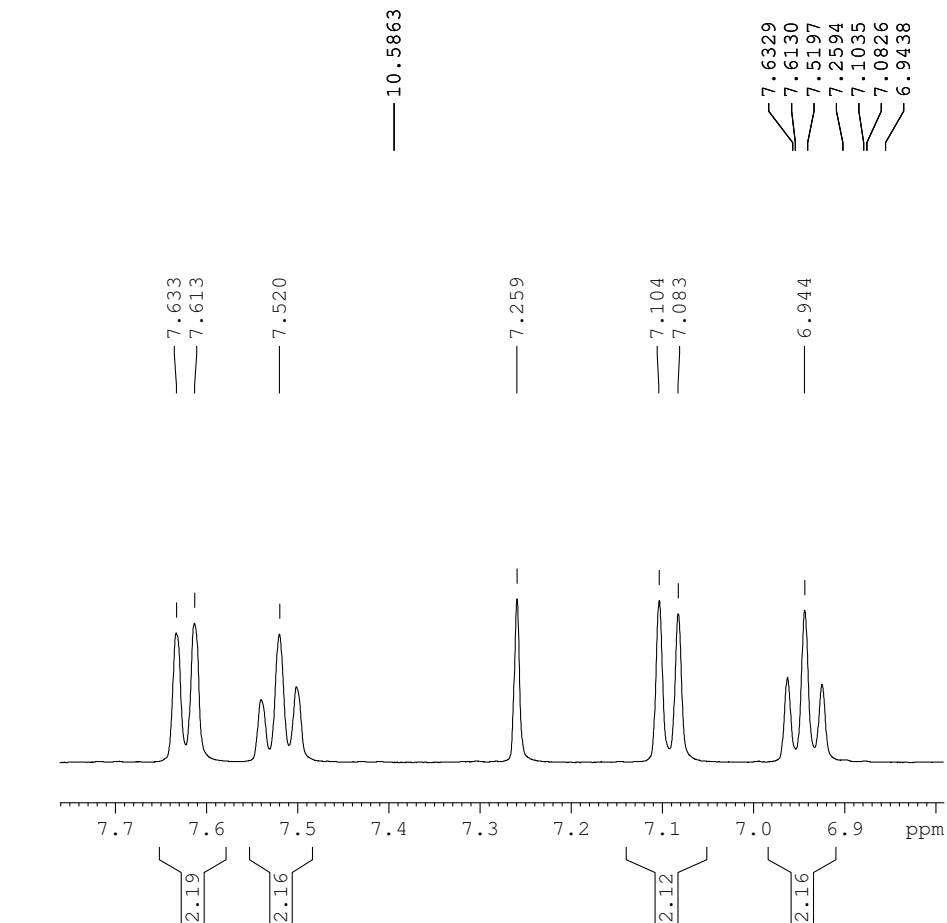
F2 - Acquisition Parameters
Date_ 20120418
Time 17.47
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 70.24
DW 60.800 usec
DE 6.50 usec
TE 298.1 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.91 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300136 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00





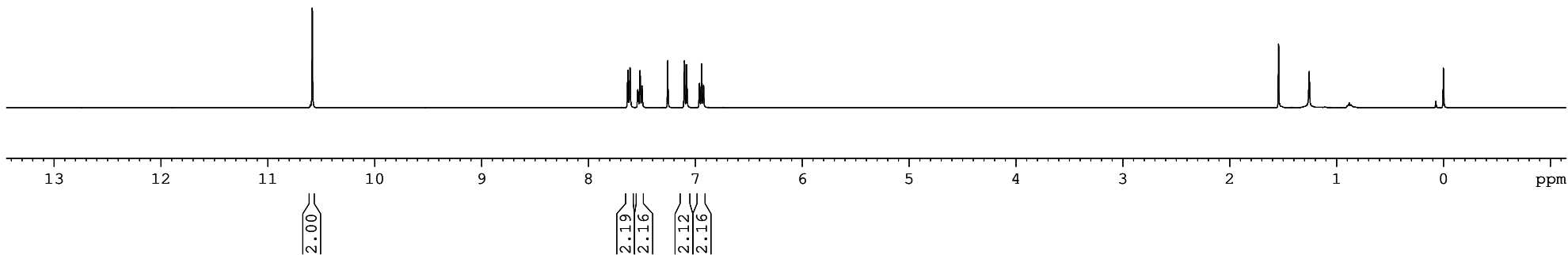


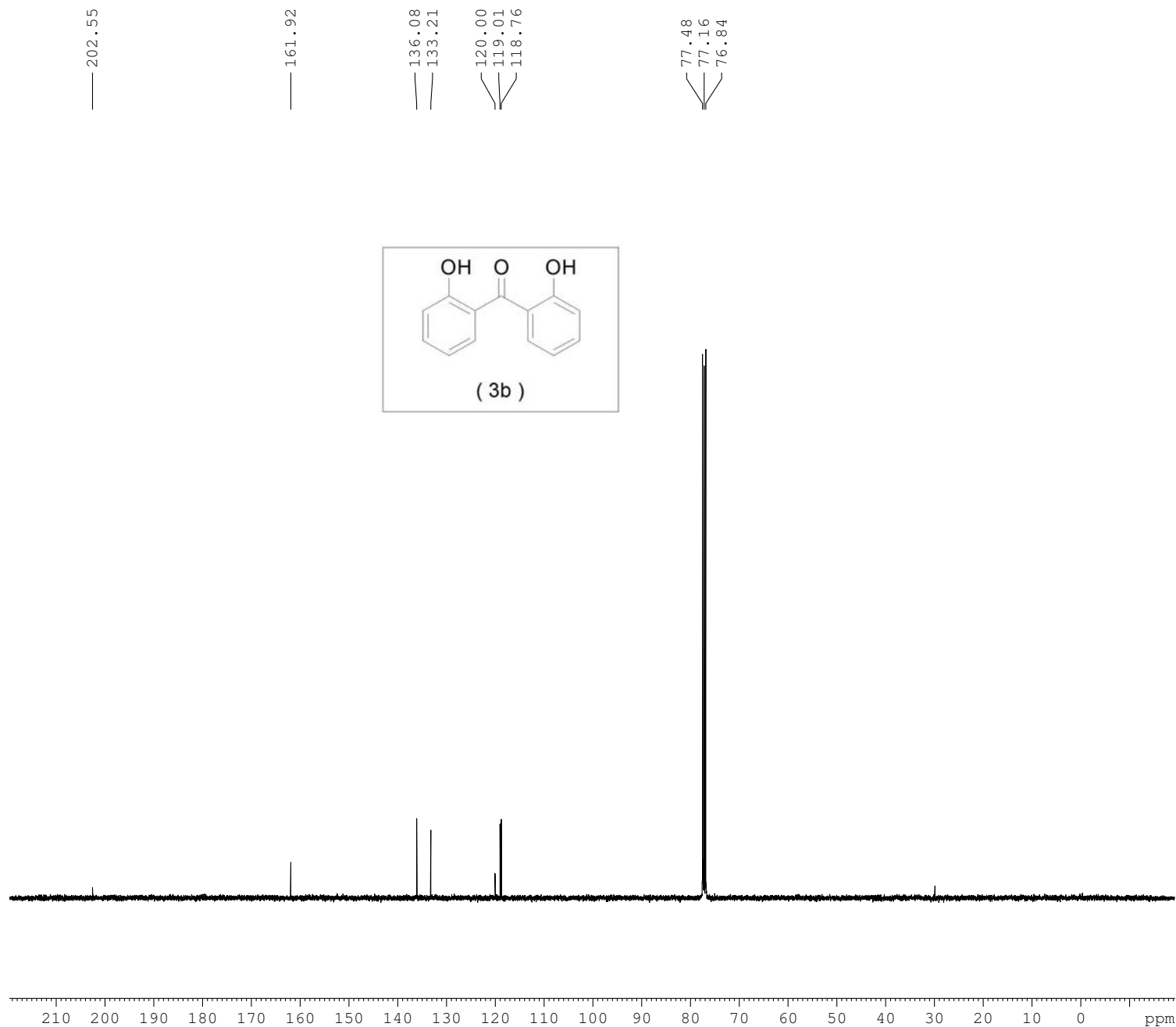
Current Data Parameters
NAME 20121108-yu 1-143
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121108
Time 15.28
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 209.25
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300105 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00





Current Data Parameters

NAME 20121108-yu 1-143
EXPNO 11
PROCNO 1

F2 - Acquisition Parameters

Date_ 20121108
Time 16.03
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 488
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====

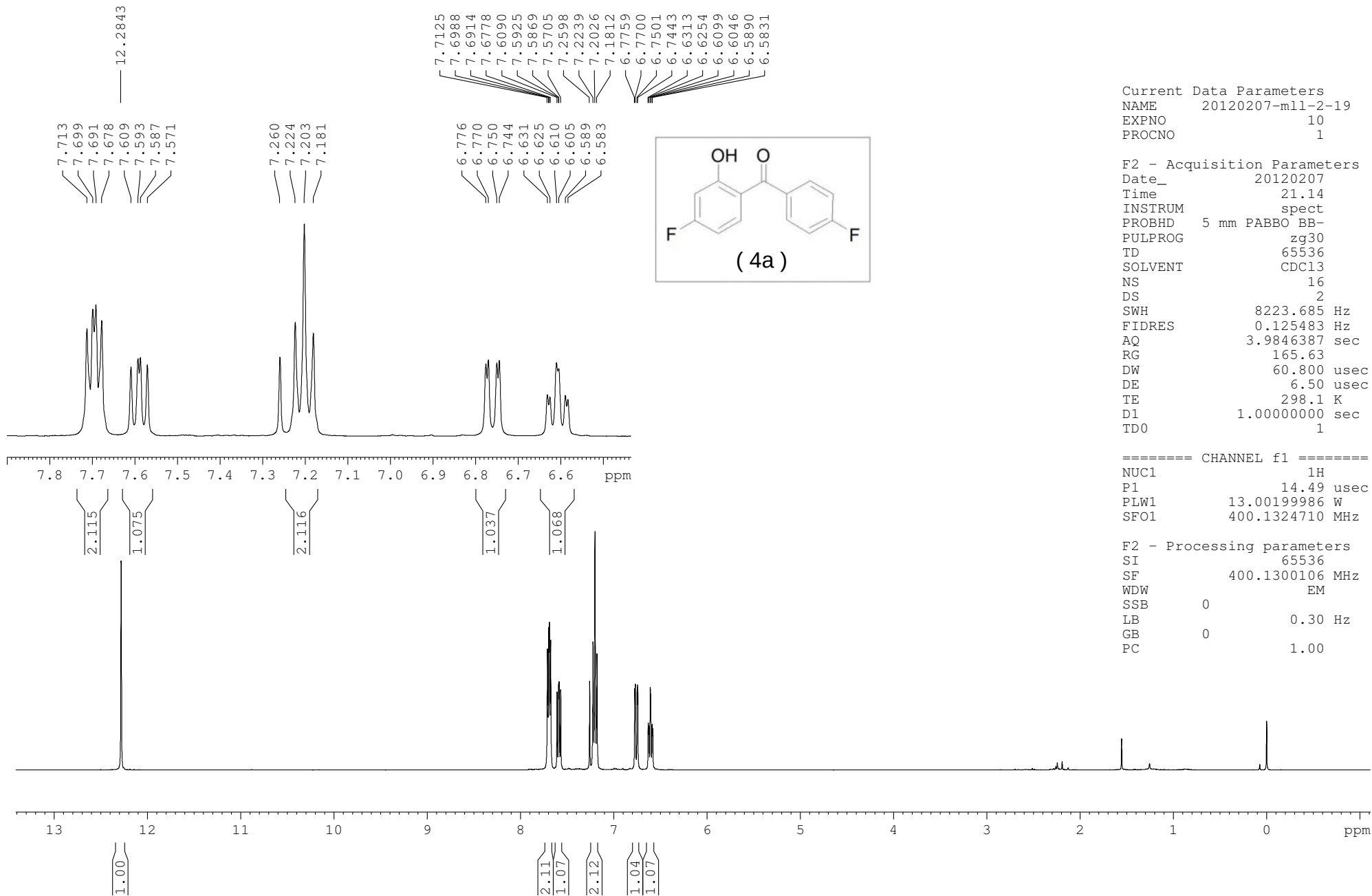
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

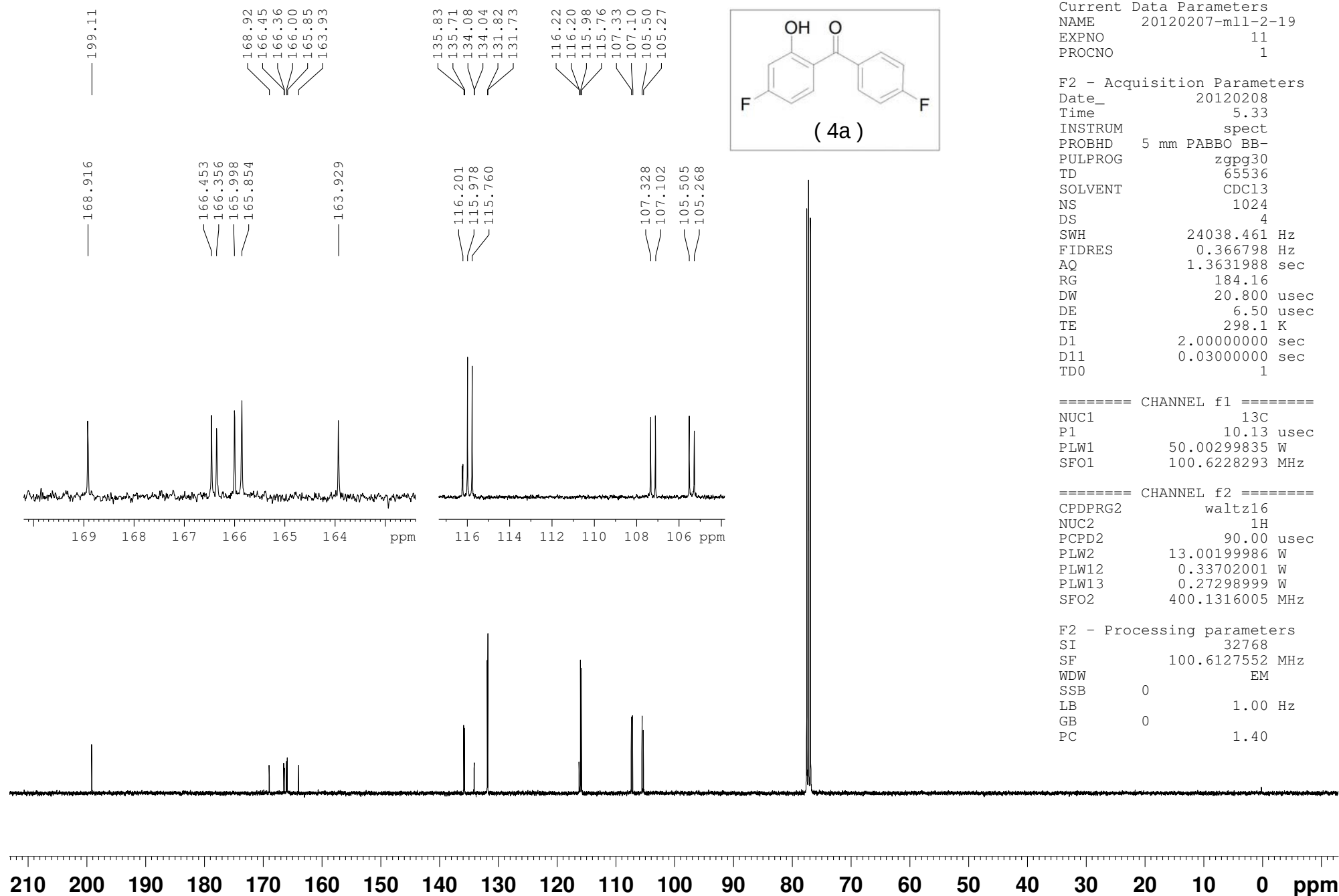
===== CHANNEL f2 =====

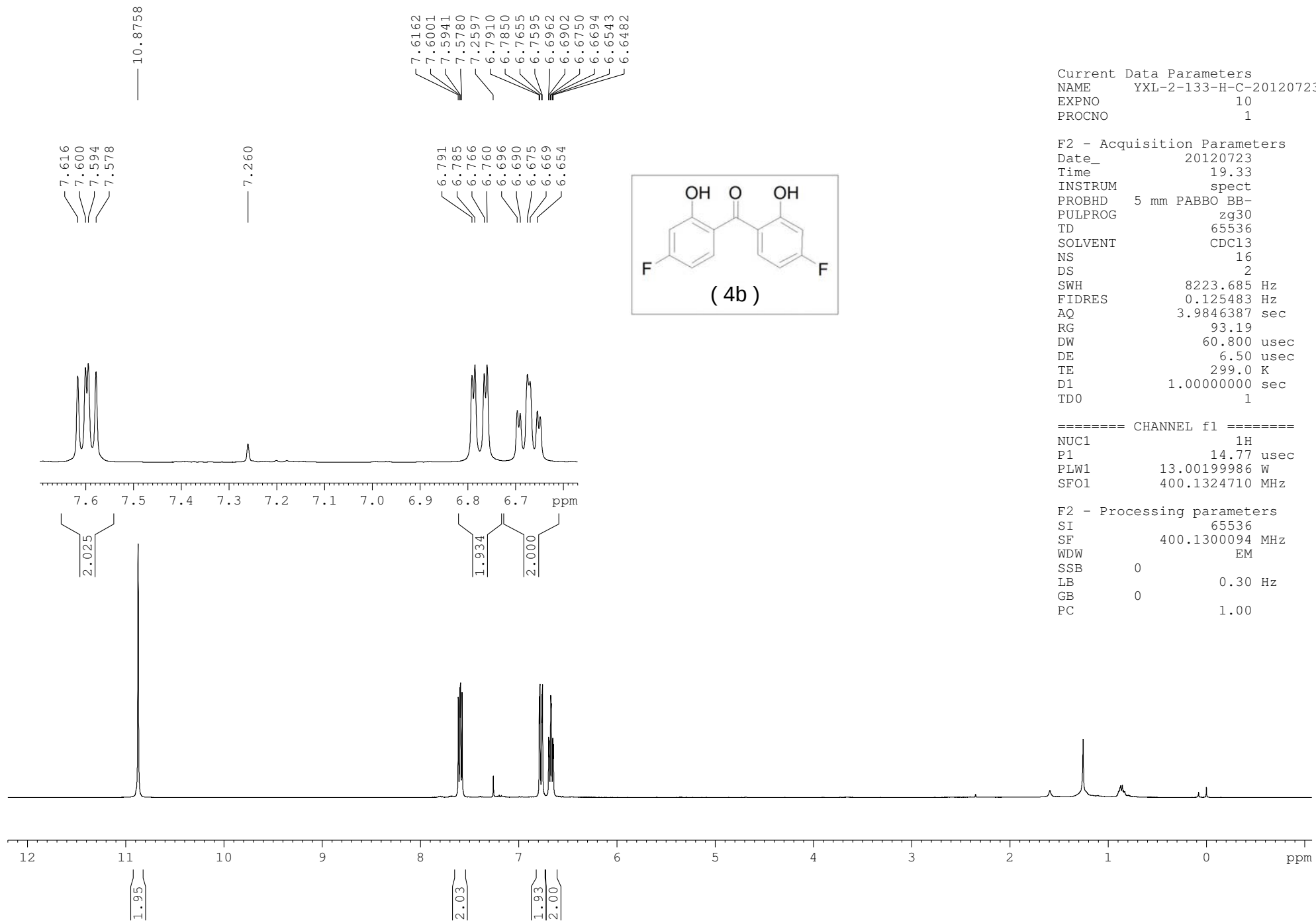
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

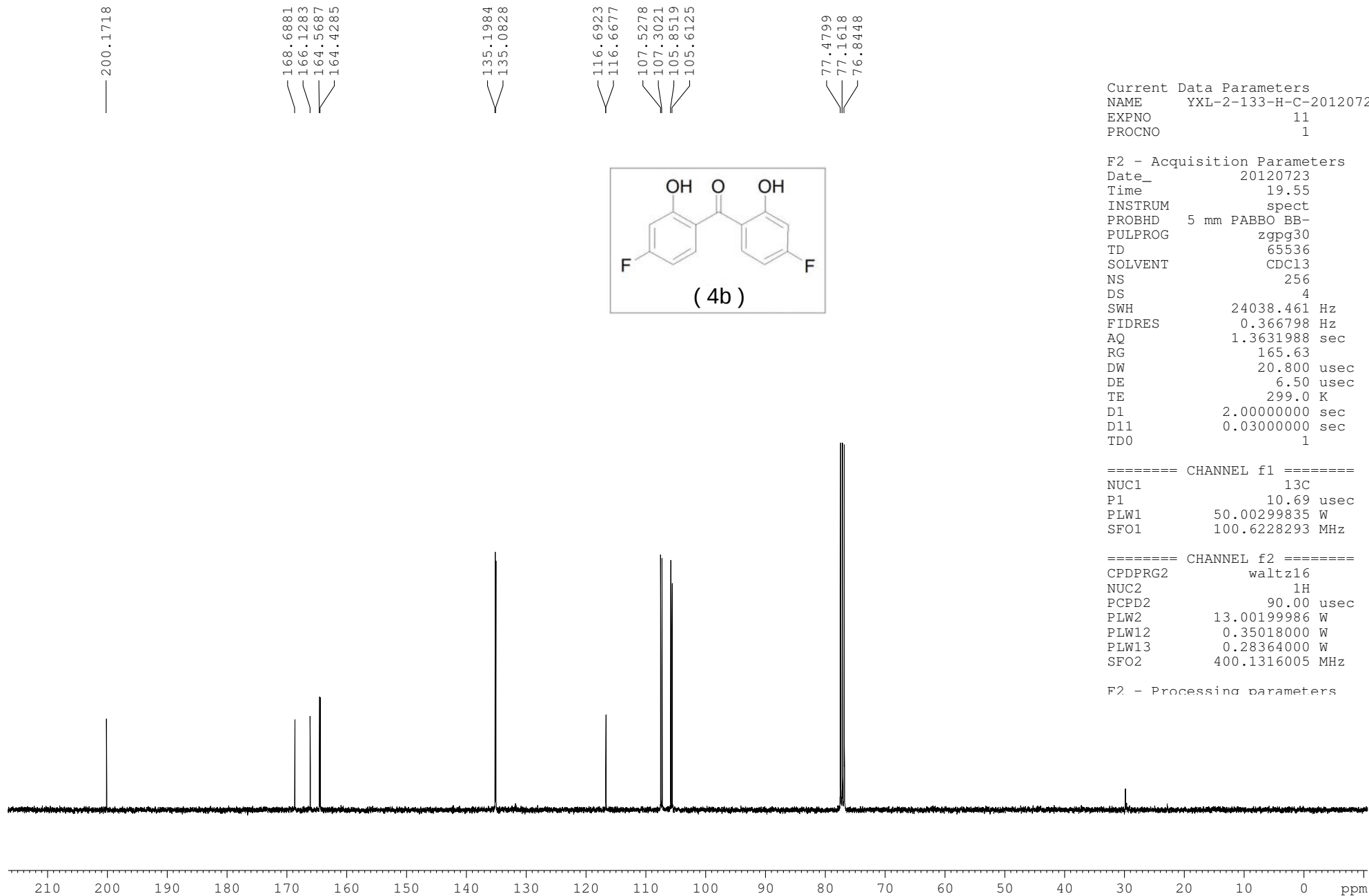
F2 - Processing parameters

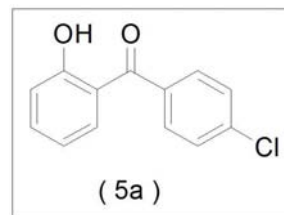
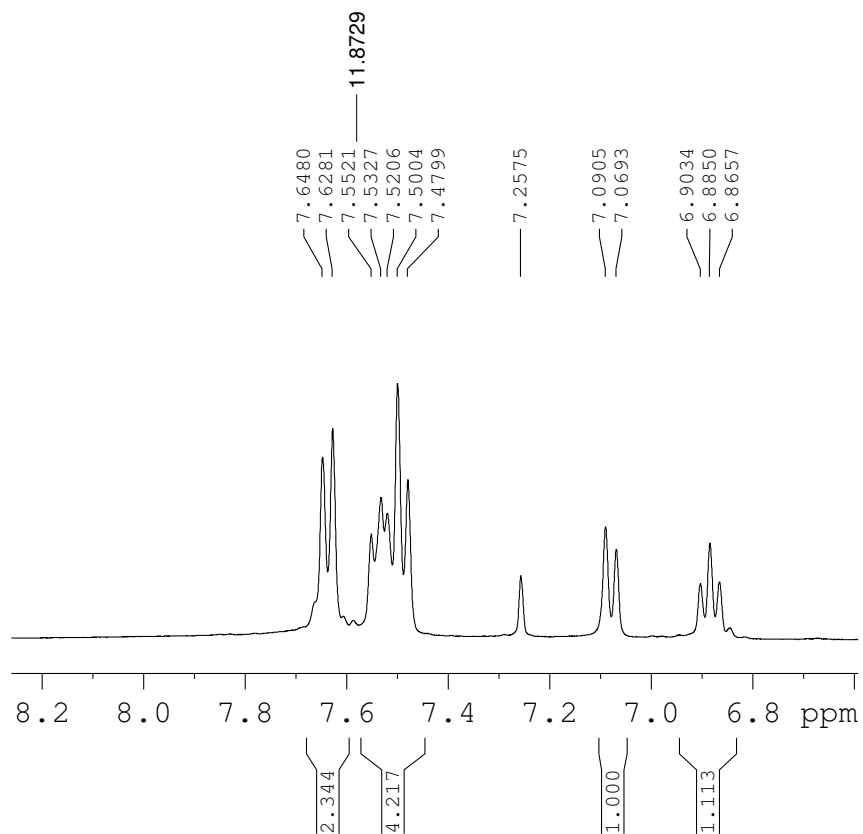
SI 32768
SF 100.6127539 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40









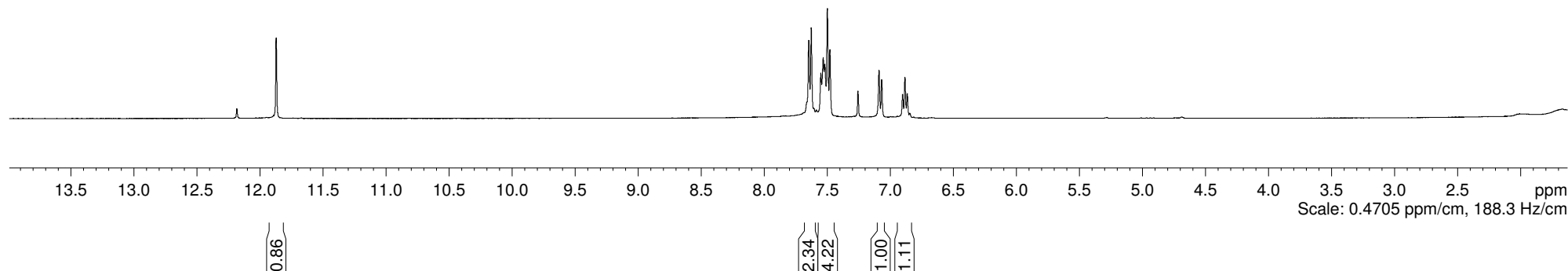


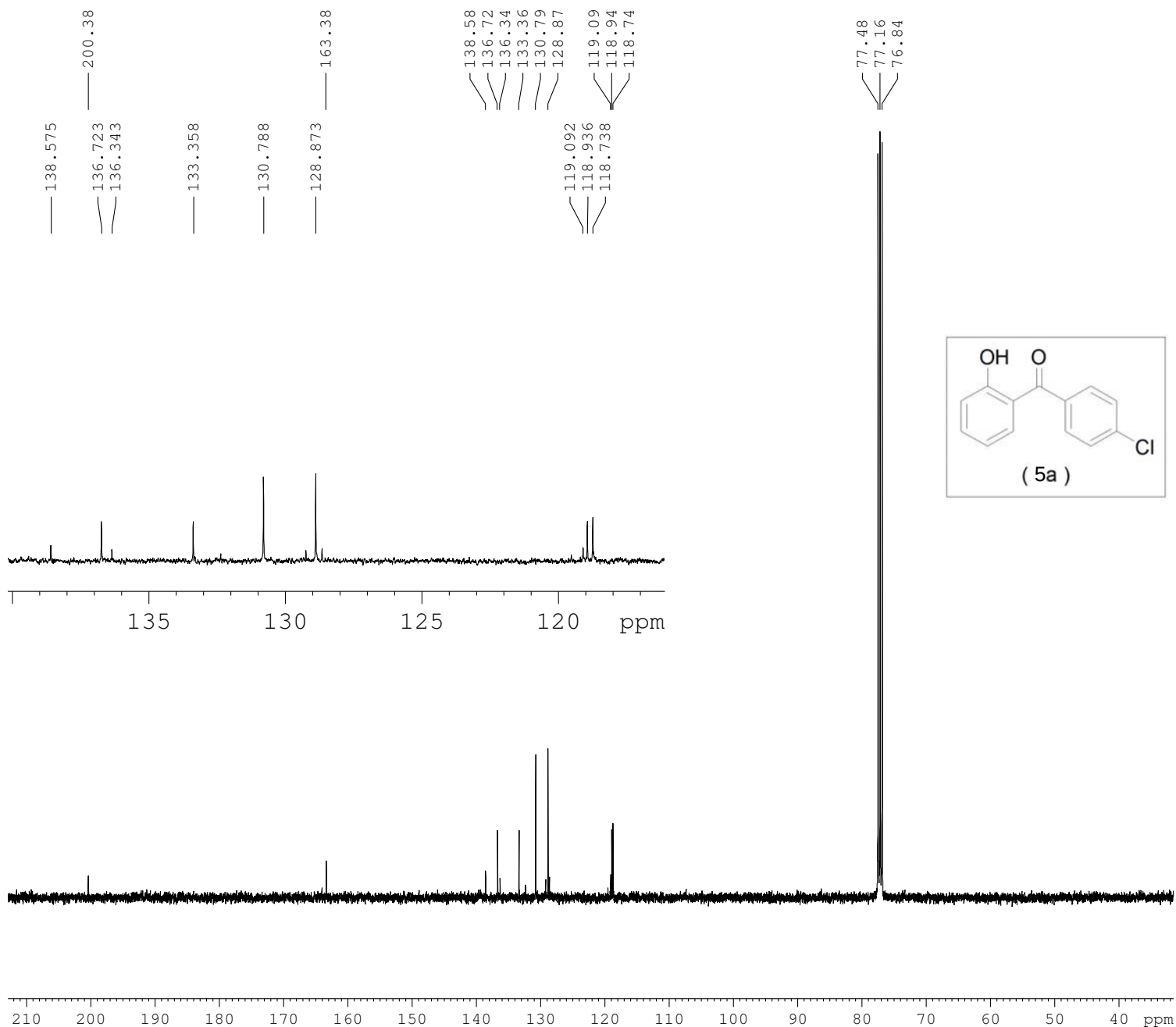
Current Data Parameters
NAME 20121127-ysy---1--mono
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121127
Time 17.31
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 102.67
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SF01 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300117 MHz
WDW EM





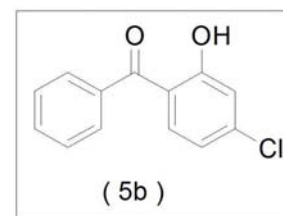
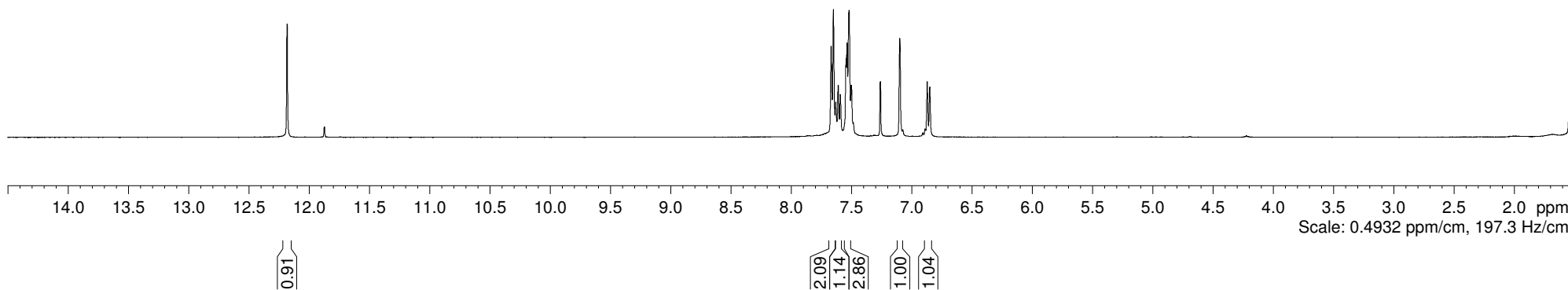
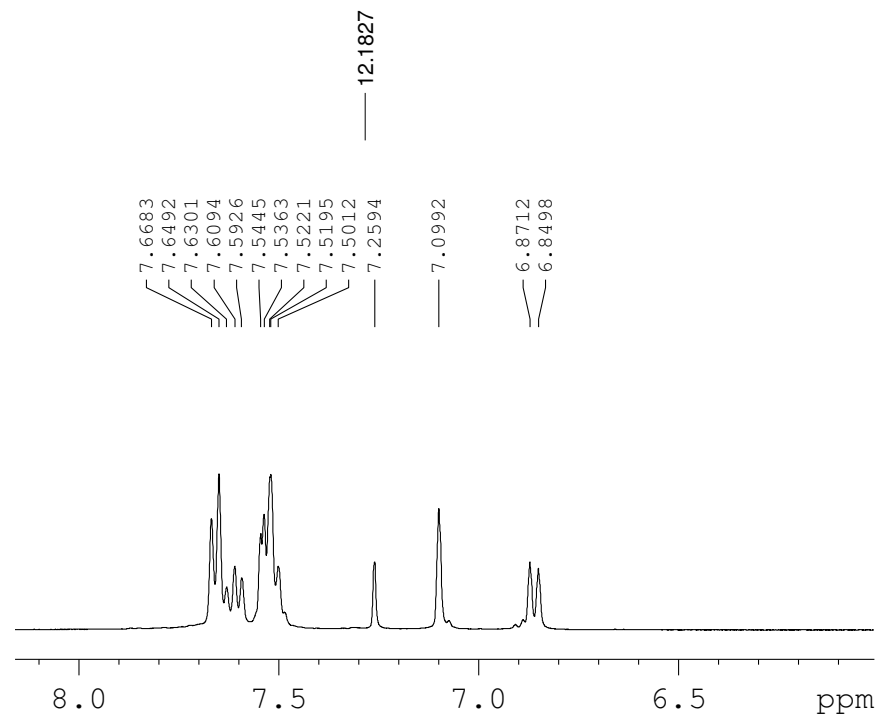
Current Data Parameters
NAME 20121128-ysy---1--4-lv
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121128
Time 18.17
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127539 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



Current Data Parameters
NAME 20121128-ysy--1-lv--er
EXPNO 10
PROCNO 1

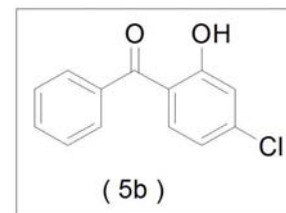
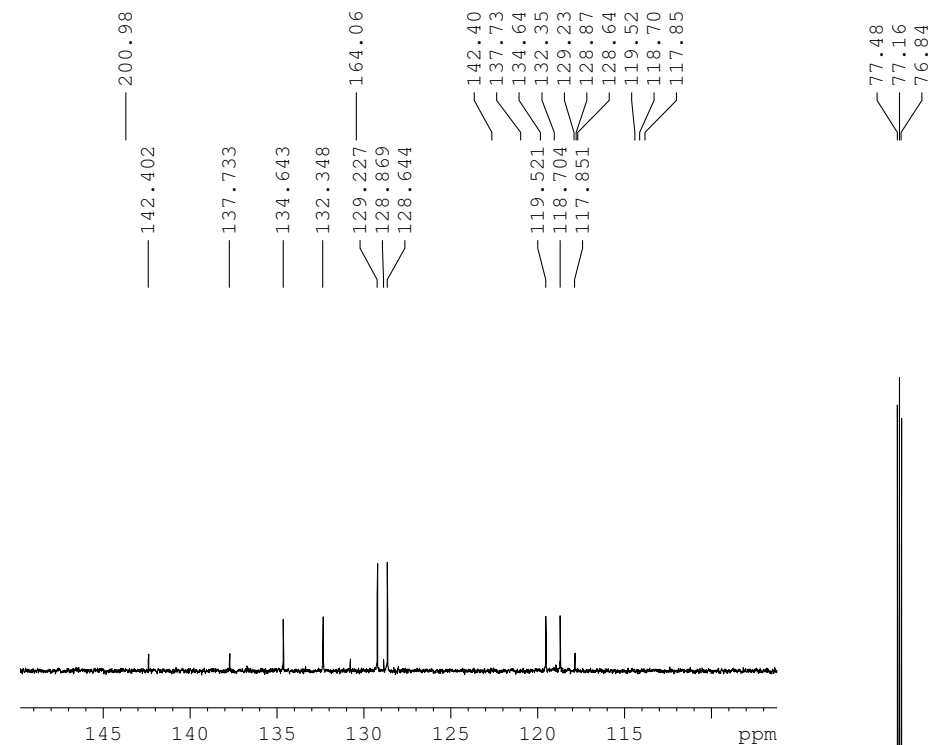
F2 - Acquisition Parameters
Date_ 20121128
Time 19.51
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 165.63
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300103 MHz
WDW EM
SSB 0
GB 0
PC 1.00

Scale: 0.4932 ppm/cm, 197.3 Hz/cm

C13CPD CDC13 {D:\NMR_DATA} RY 32



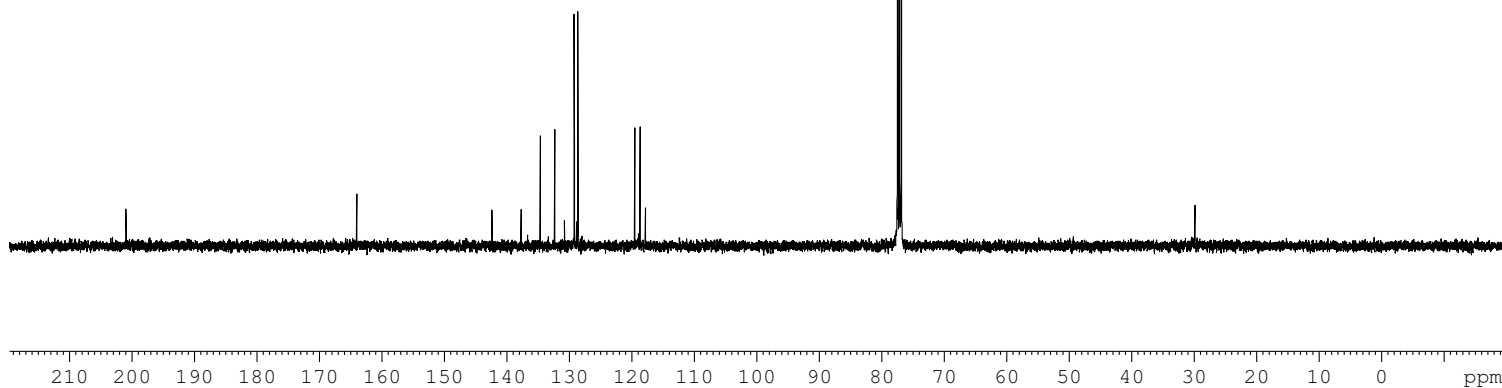
Current Data Parameters
NAME 20121128-ysy--1-lv--erbenjiaotong
EXPNO 11
PROCNO 1

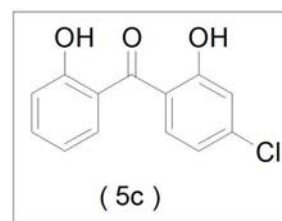
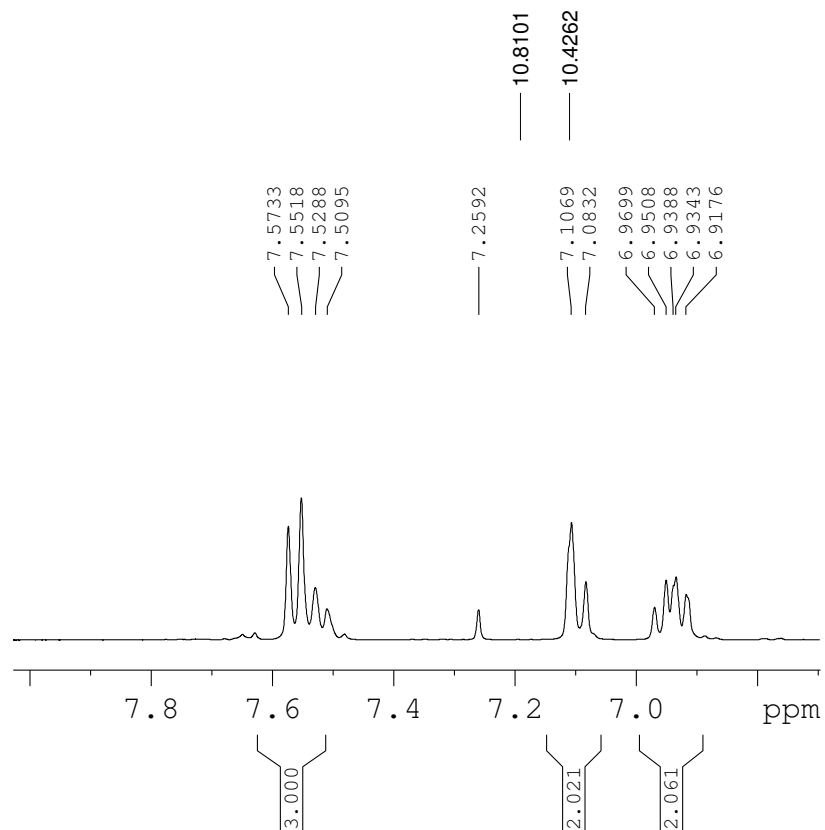
F2 - Acquisition Parameters
Date_ 20121129
Time 0.19
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127541 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





Current Data Parameters
NAME 20121123-ysy---duilv erbenjiat
EXPNO 10
PROCNO 1

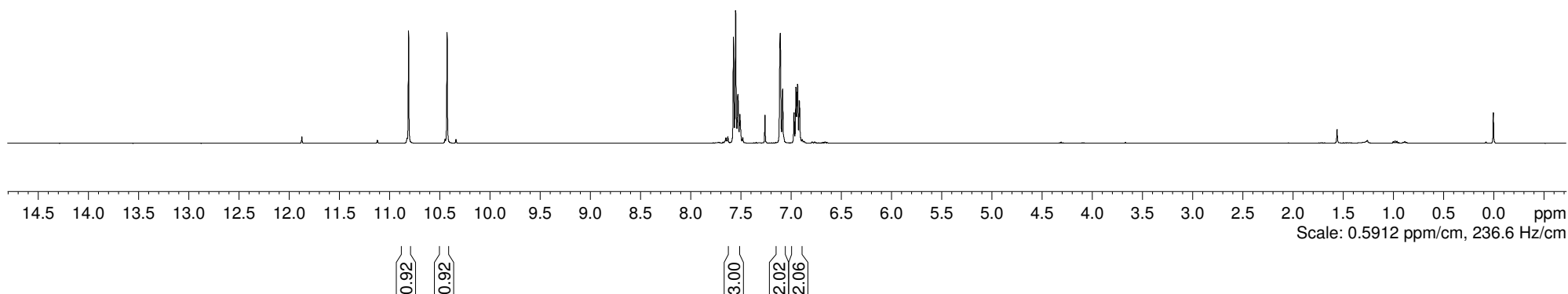
F2 - Acquisition Parameters

Date_ 20121123
Time 11.08
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 144.49
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

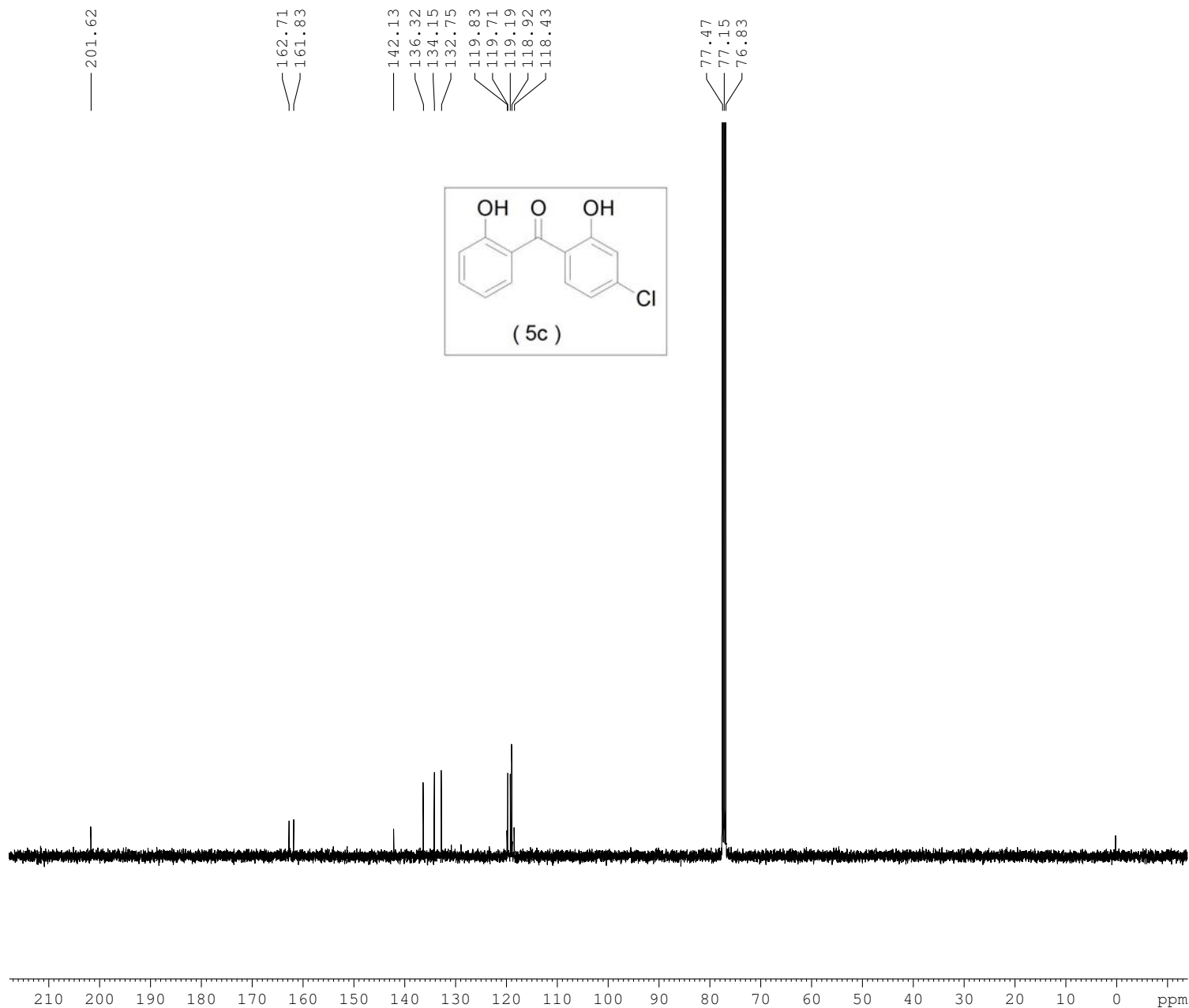
===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters

SI 65536
SF 400.1300105 MHz
WDW EM



C13CPD CDC13 {D:\NMR_DATA} RY 36



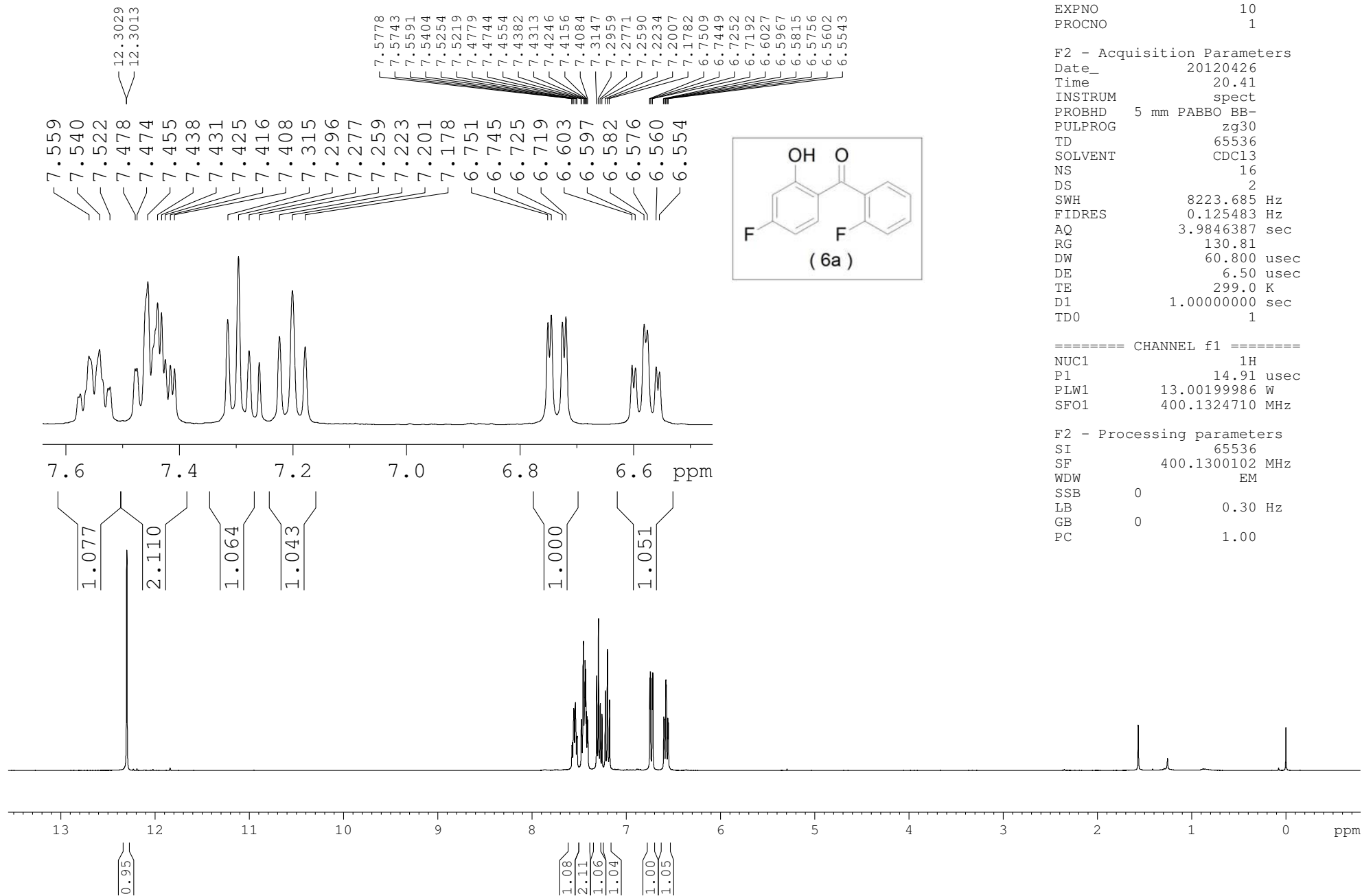
Current Data Parameters
NAME ysy-1----16220121120-
EXPNO 10
PROCNO 1

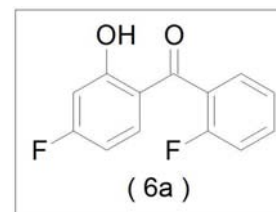
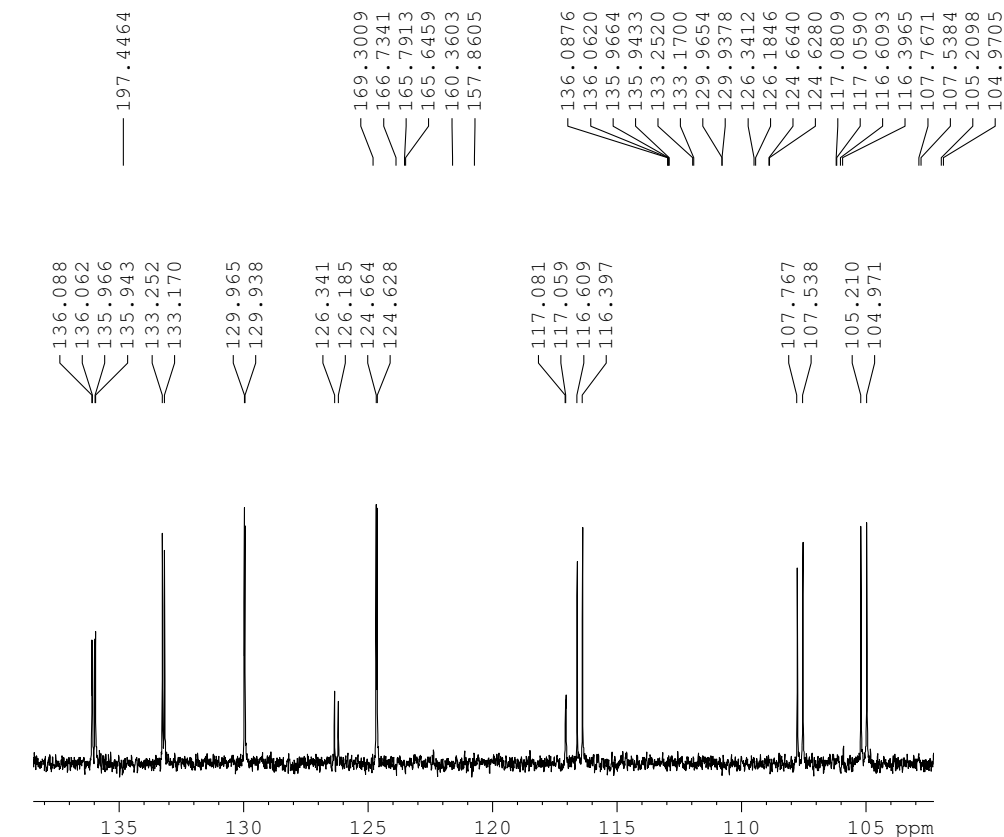
F2 - Acquisition Parameters
Date_ 20121120
Time 19.53
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127547 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





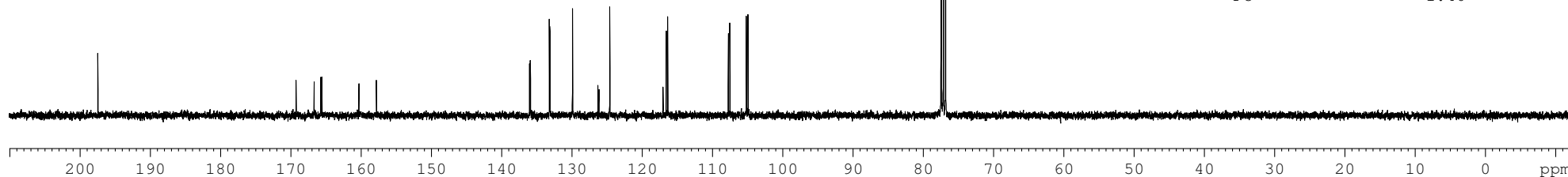
Current Data Parameters
NAME YXL-1-281-1-H-C-20120428
EXPNO 10
PROCNO 1

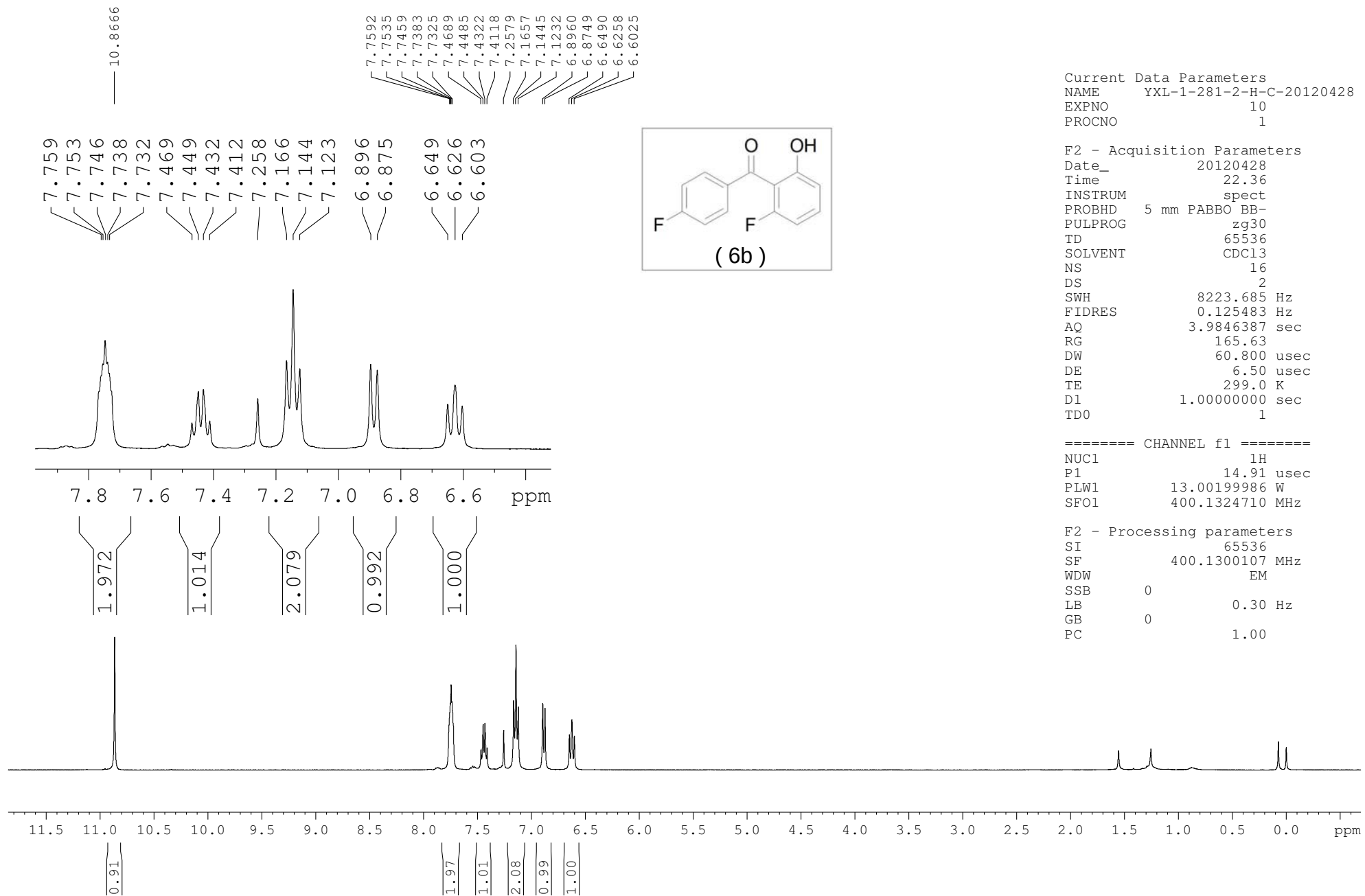
F2 - Acquisition Parameters
Date_ 20120428
Time 22.48
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 256
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

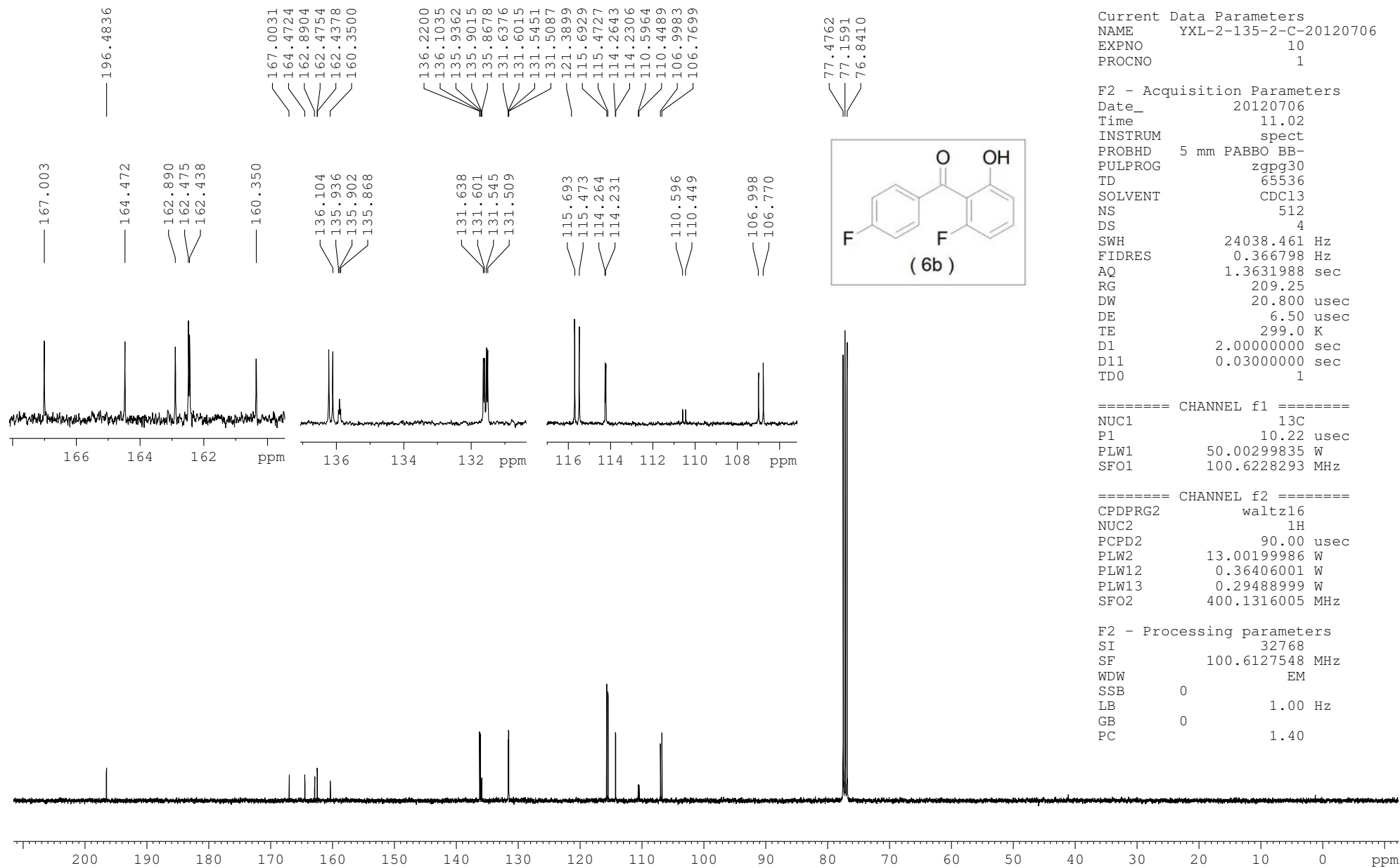
===== CHANNEL f1 =====
NUC1 13C
P1 10.31 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

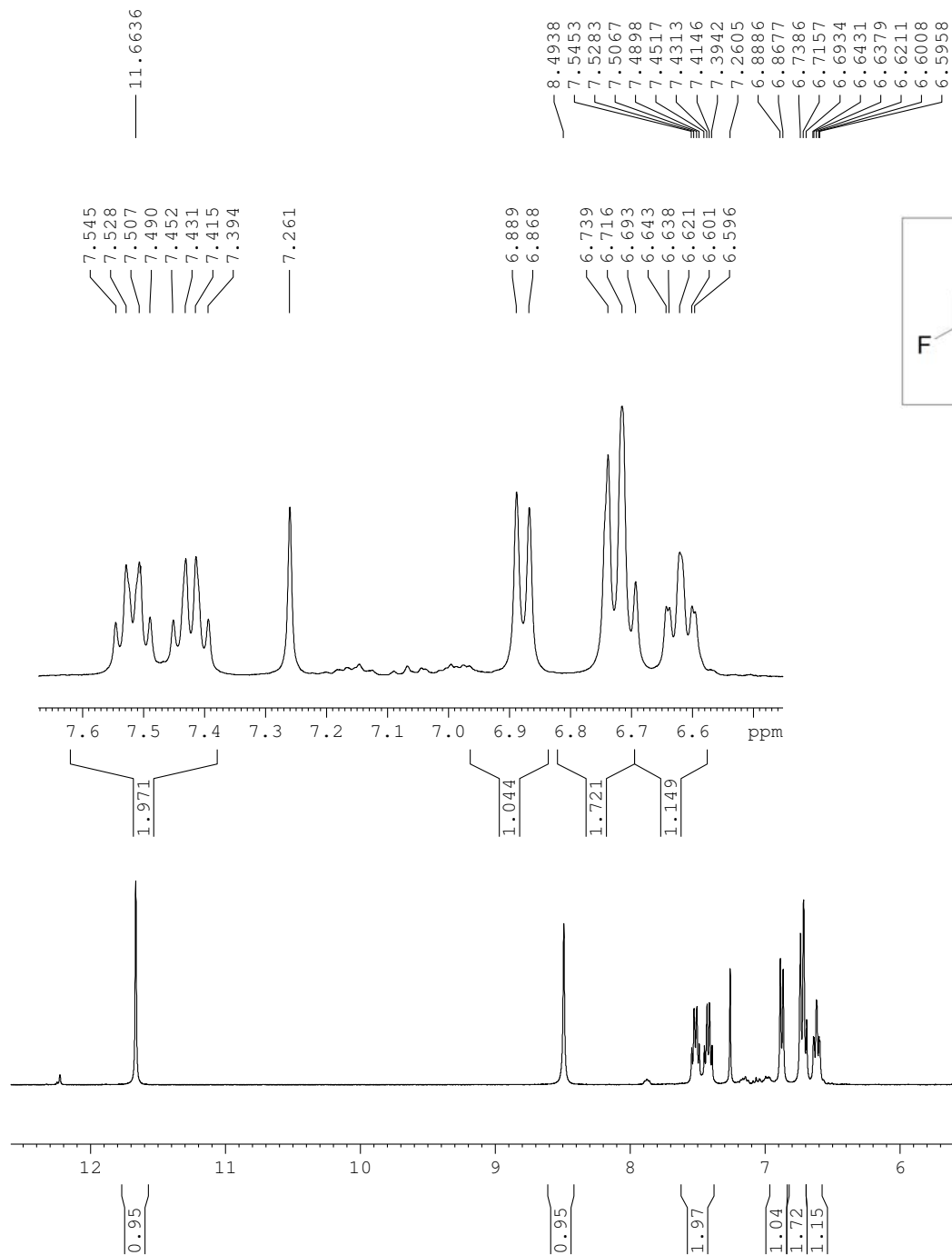
===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35685000 W
PLW13 0.28904000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127554 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40







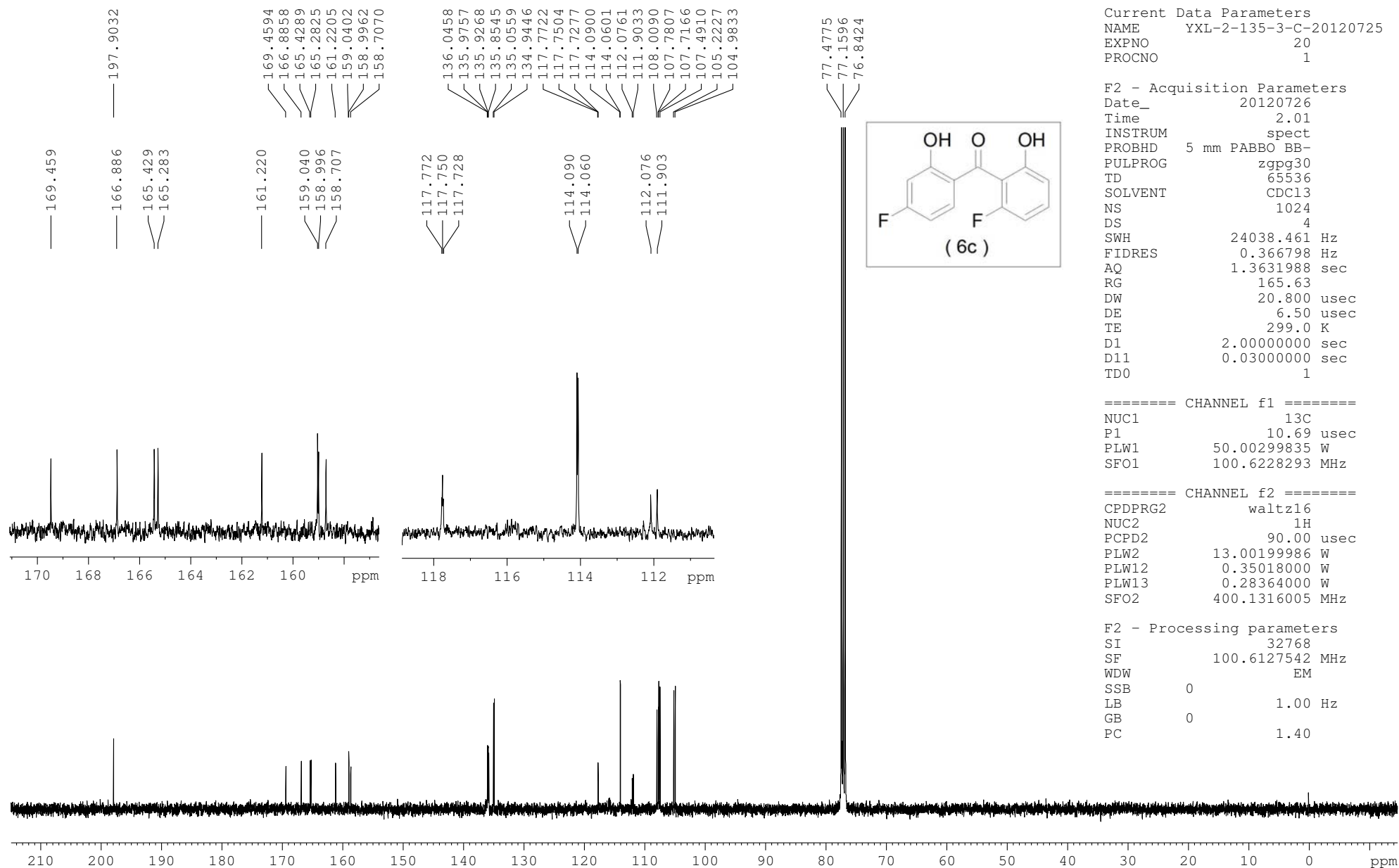


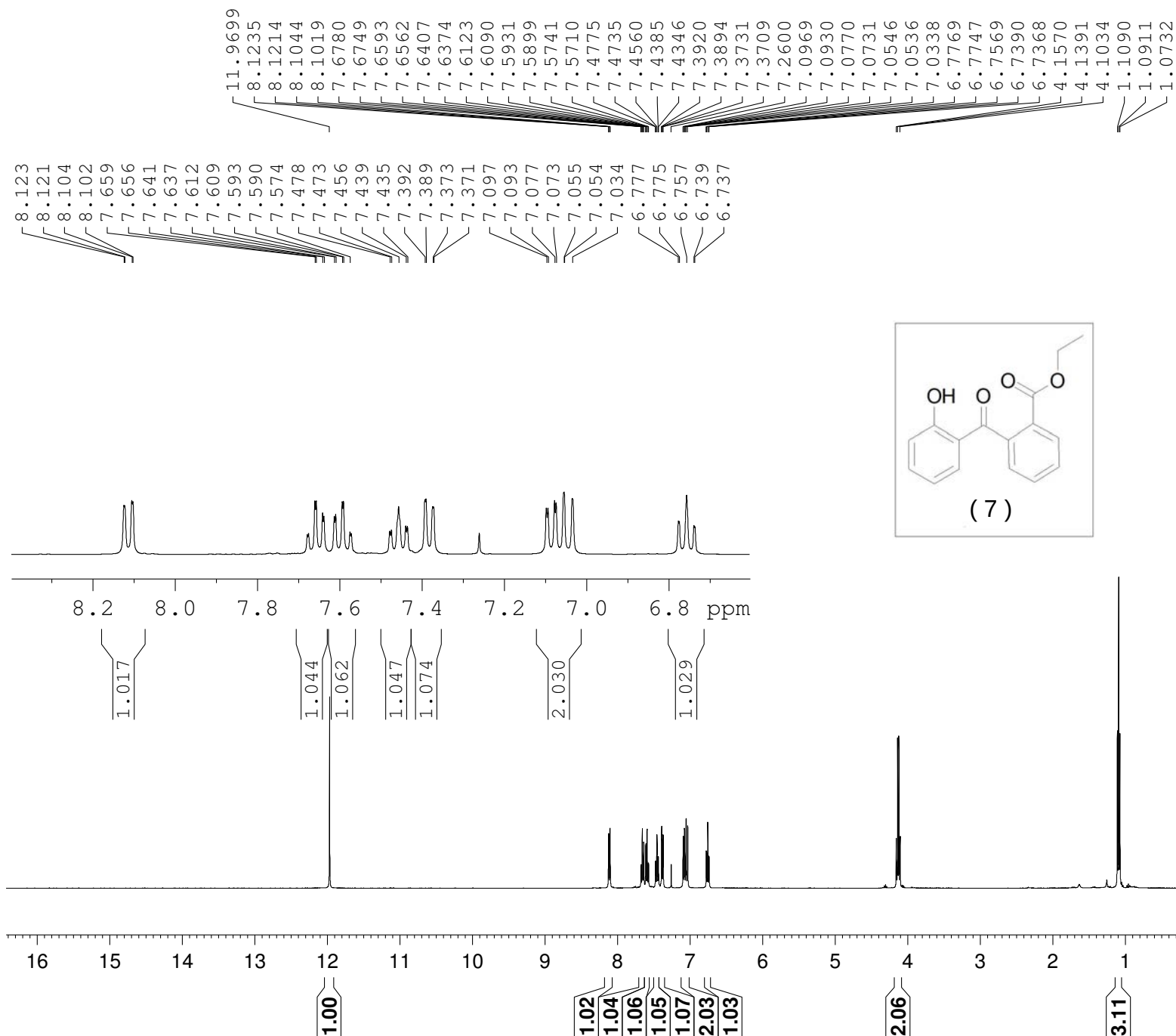
Current Data Parameters
NAME YXL-2-135-3-H-C-20120725
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120725
Time 21.36
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 184.16
DW 60.800 usec
DE 6.50 usec
TE 299.0 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300096 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



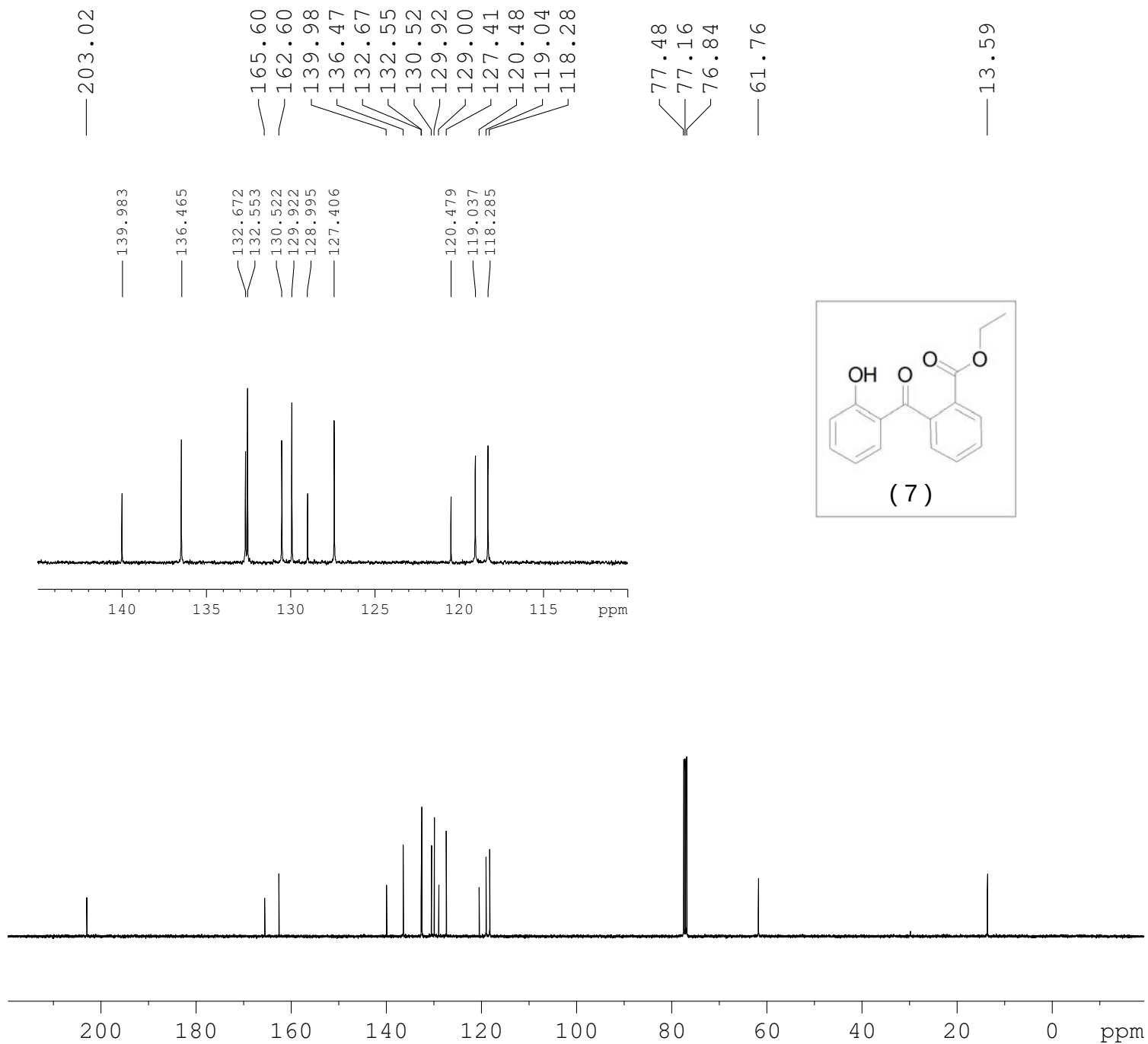


Current Data Parameters
NAME 20120526-ml1-3-3-p1
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120526
Time 16.58
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 93.19
DW 60.800 usec
DE 6.50 usec
TE 299.0 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 10.83 usec
PLW1 13.00199986 W
SF01 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300095 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



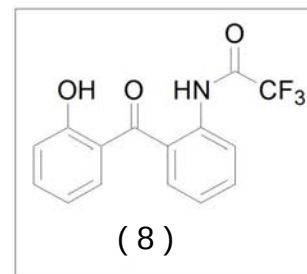
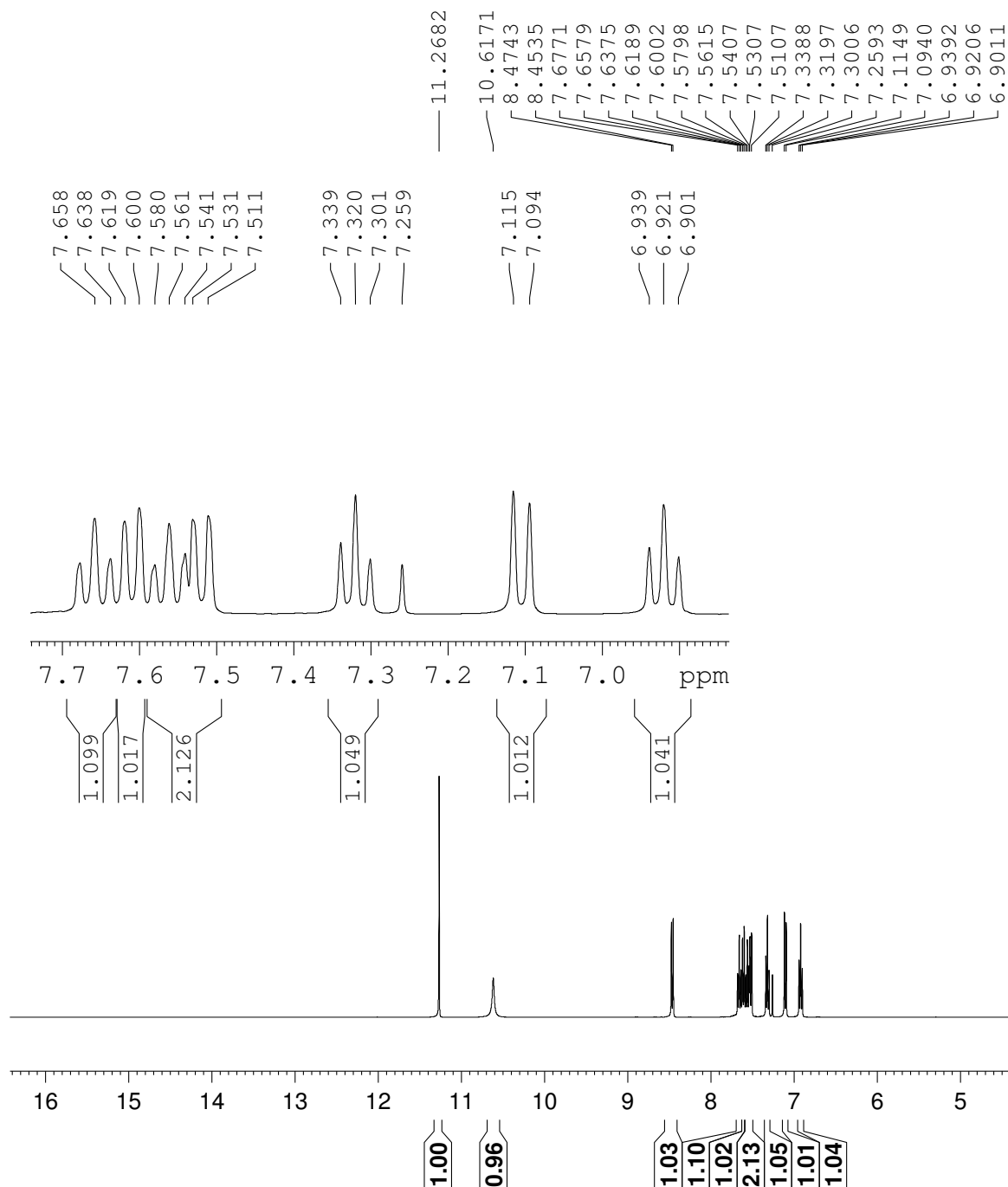
Current Data Parameters
NAME 20120530-ml1-3-3-pl
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120530
Time 17.28
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 300
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.0 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.38 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.18827000 W
PLW13 0.15250000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127584 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

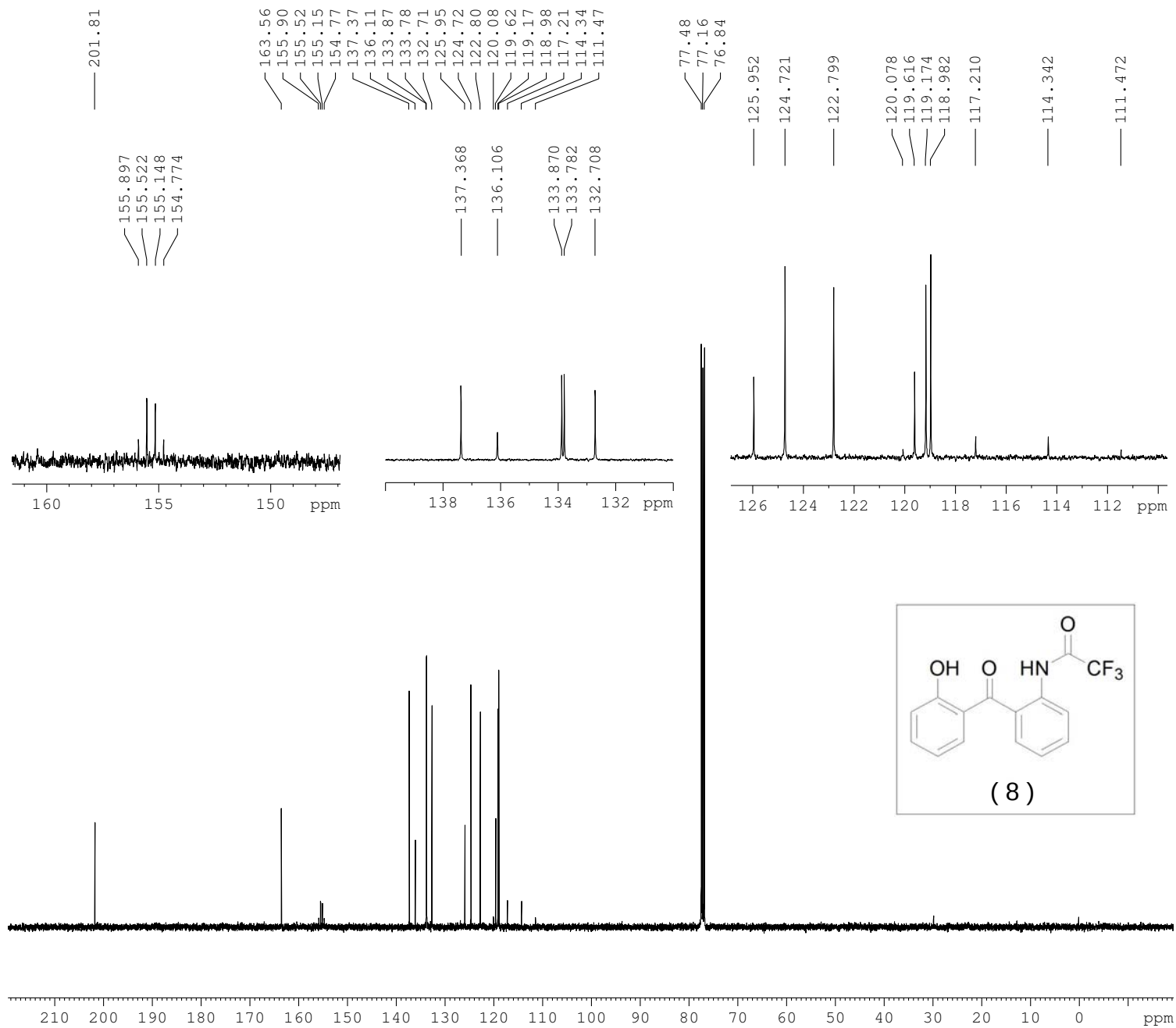


Current Data Parameters
NAME SG-7-32-h-20120704
EXPNO 20
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120705
Time 9.32
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 115.38
DW 60.800 usec
DE 6.50 usec
TE 299.0 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 15.06 usec
PLW1 13.00199986 W
SF01 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300098 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



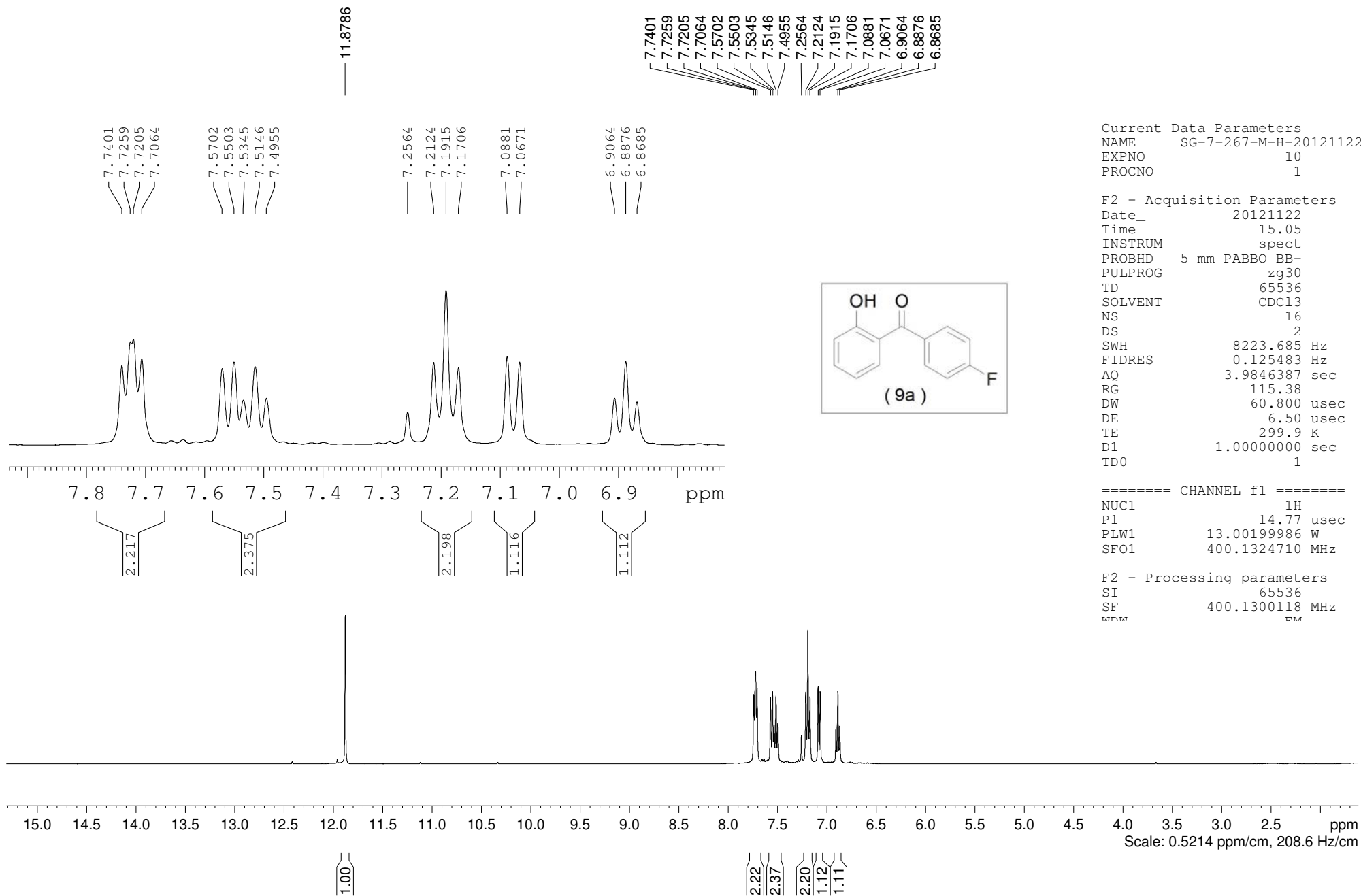
Current Data Parameters
NAME SG-7-32-C-20120704
EXPNO 10
PROCNO 1

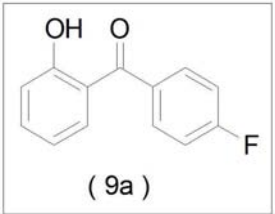
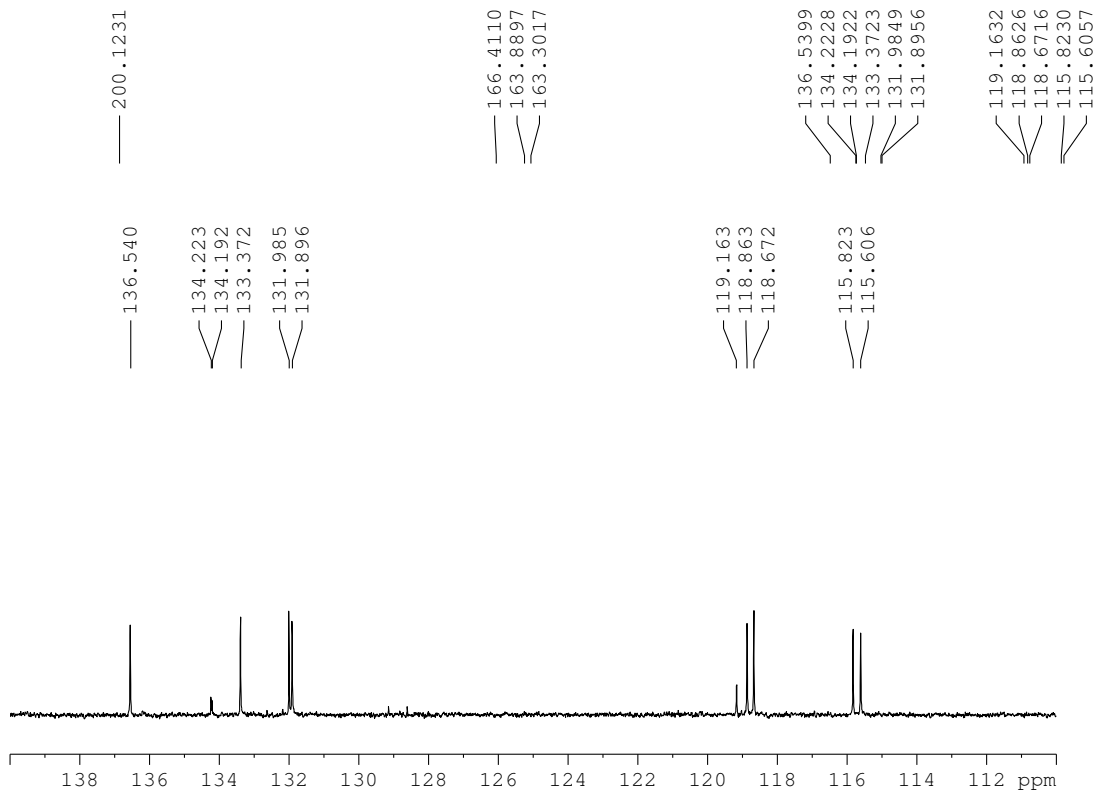
F2 - Acquisition Parameters
Date_ 20120705
Time 0.03
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl₃
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.22 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.36406001 W
PLW13 0.29488999 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127550 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





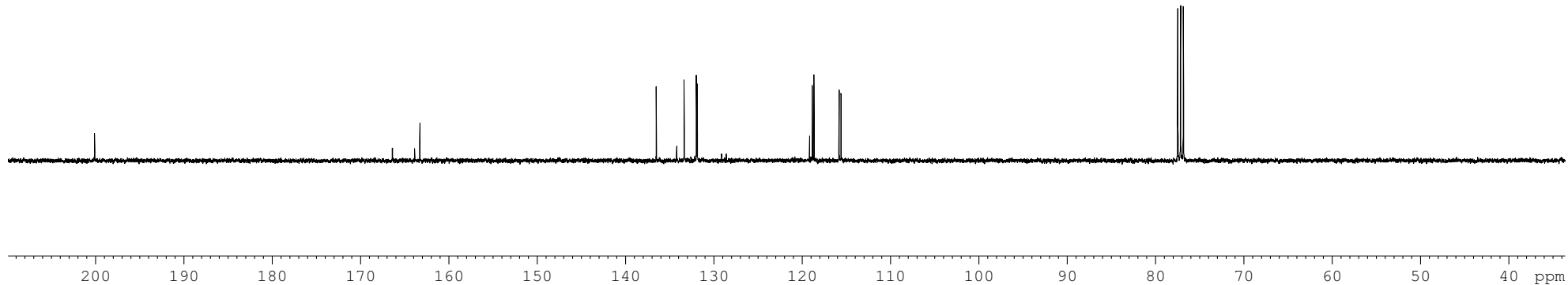
Current Data Parameters
NAME HXS-4-F-C
EXPNO 20
PROCNO 1

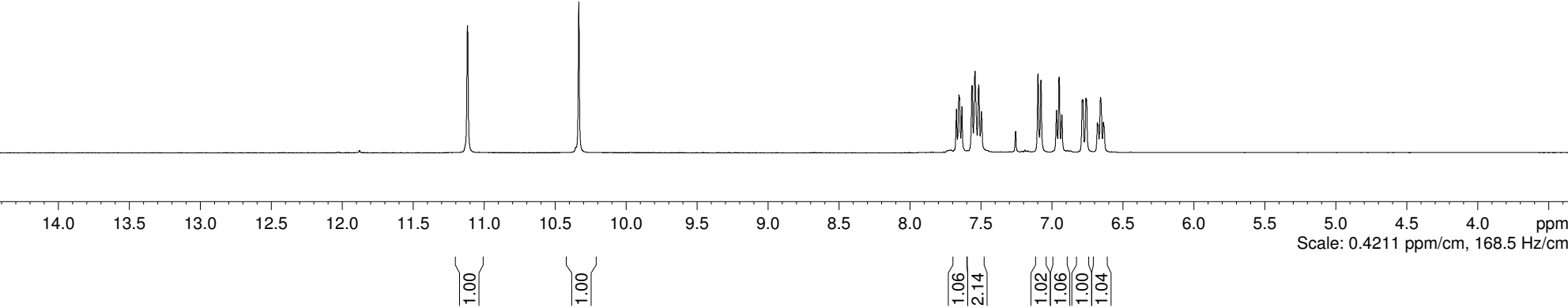
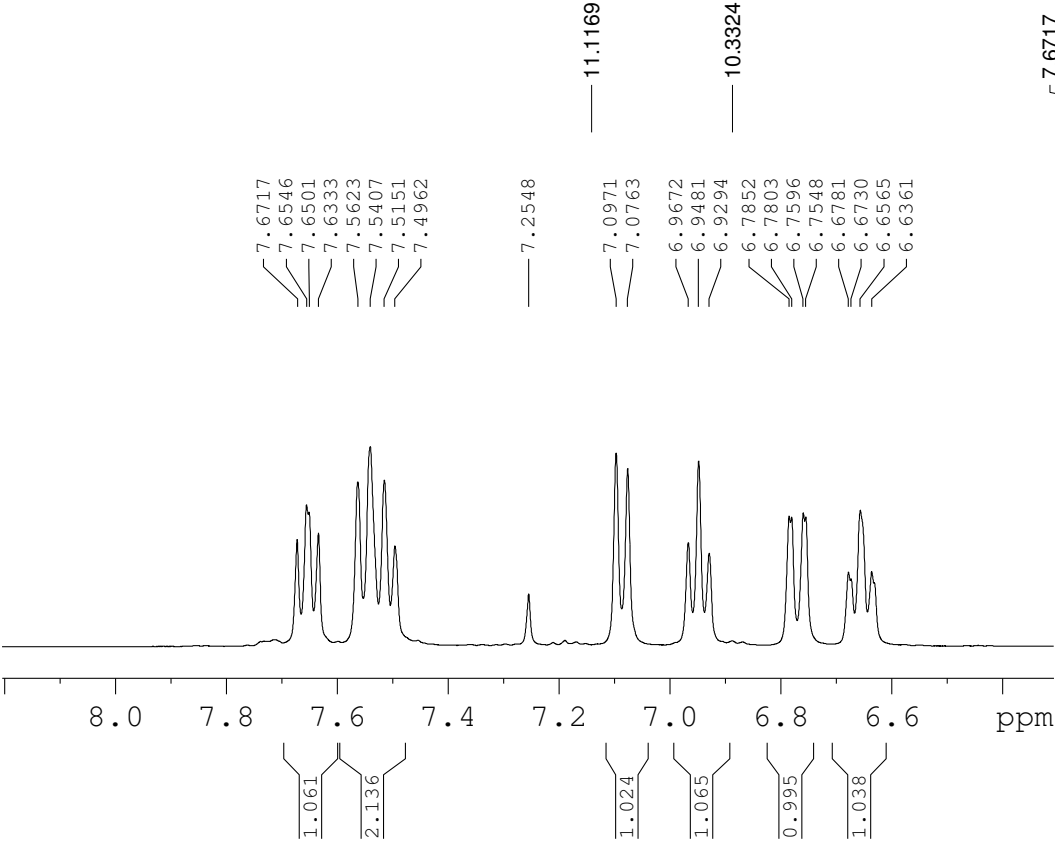
F2 - Acquisition Parameters
Date_ 20121127
Time 22.54
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 75
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters





7.6717
7.6546
7.6501
7.6333
7.5623
7.5407
7.5151
7.4962
7.2548
7.0971
7.0763
6.9672
6.9481
6.9294
6.7852
6.7803
6.7596
6.7548
6.6781
6.6730
6.6565
6.6361

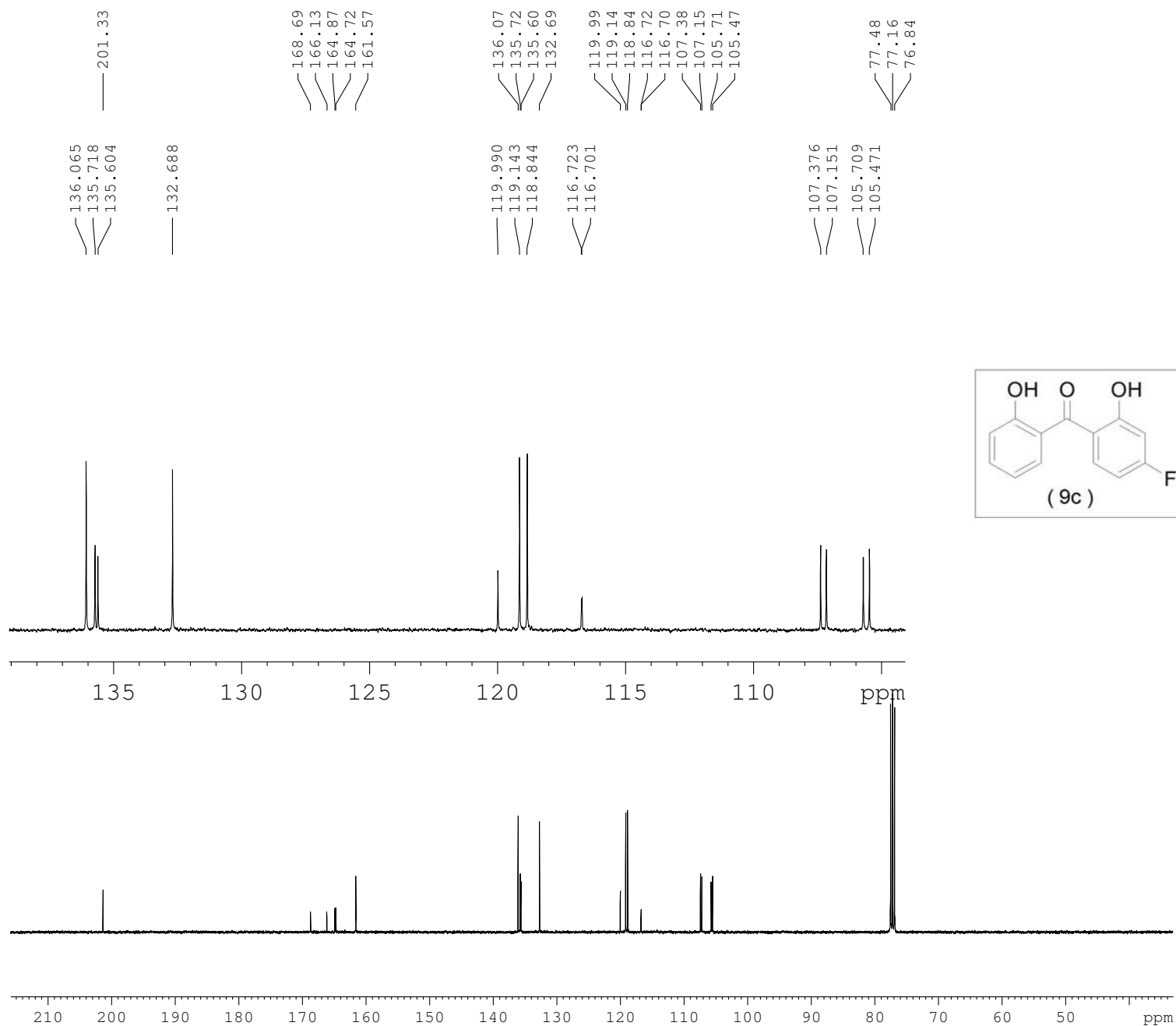
Current Data Parameters
NAME SG-7-267-D-H-20121122
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121122
Time 15.10
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 93.19
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300124 MHz
WDW EM

Scale: 0.4211 ppm/cm, 168.5 Hz/cm



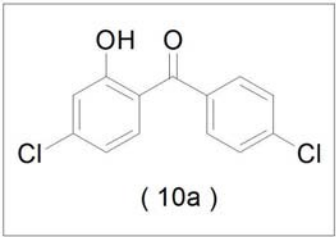
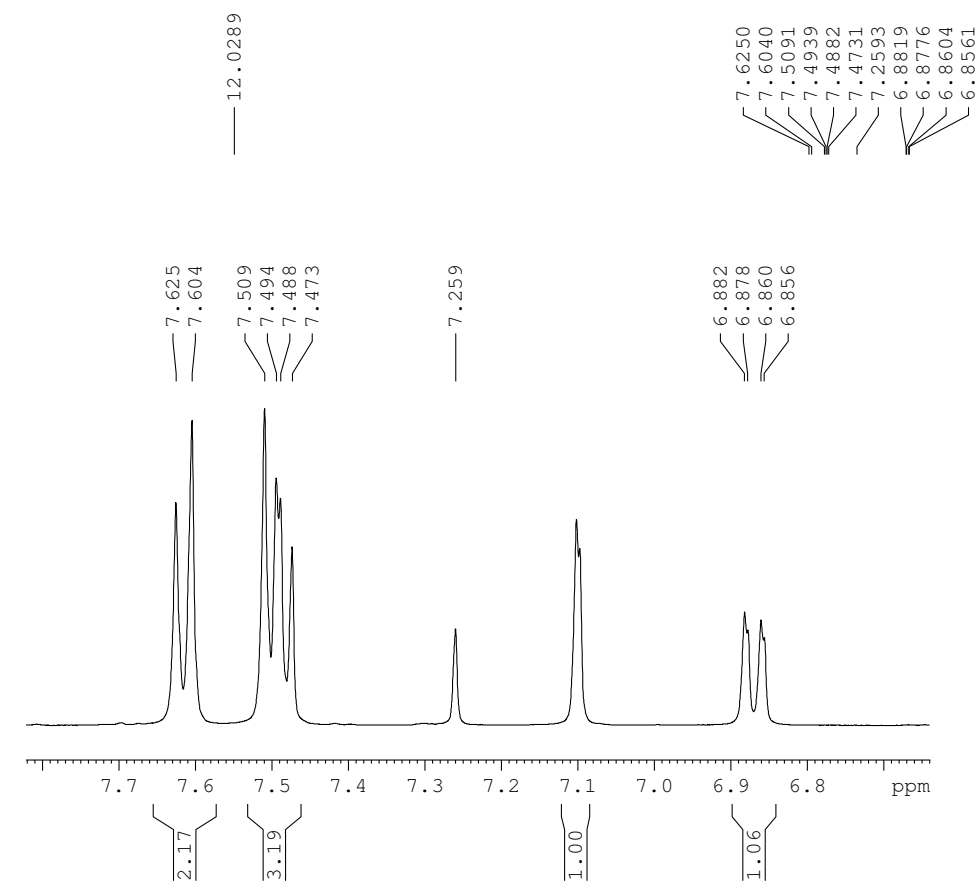
Current Data Parameters
NAME SG-7-267-M-C-20121122
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121123
Time 5.33
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 ¹³C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 ¹H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127556 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

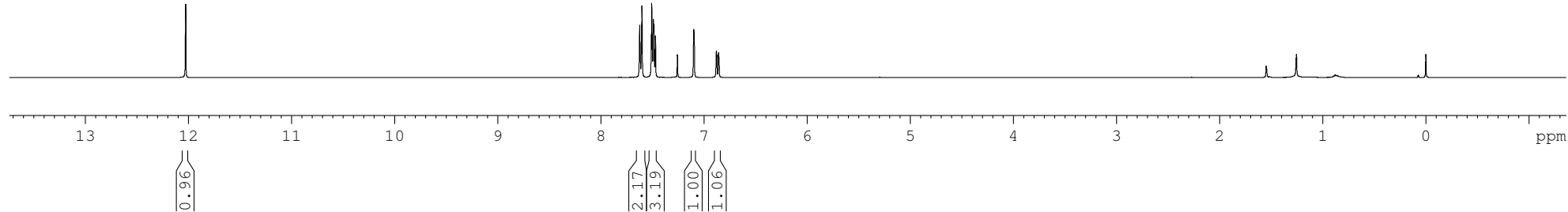


Current Data Parameters
NAME 20121112-ysy-1-150
EXPNO 20
PROCNO 1

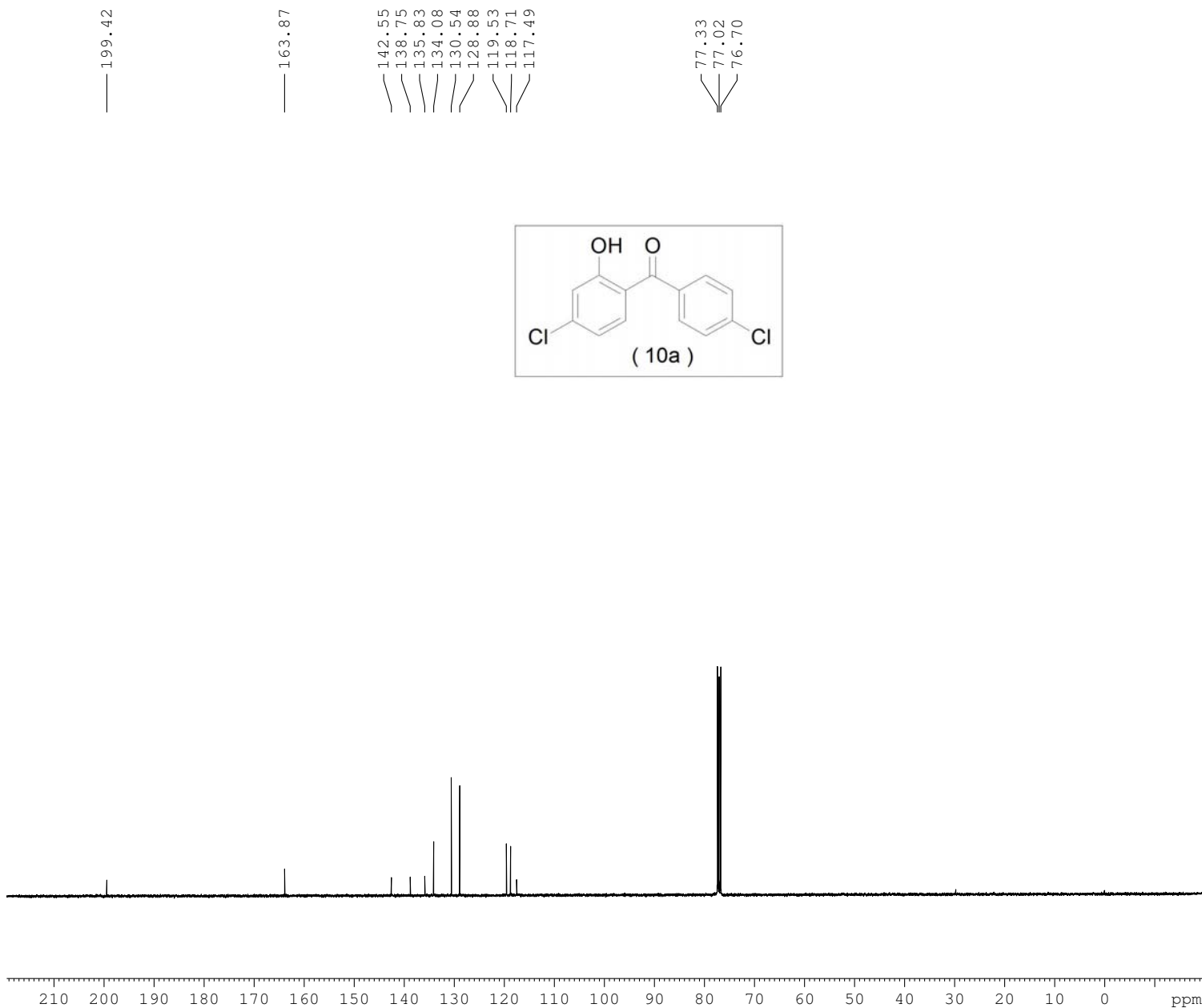
F2 - Acquisition Parameters
Date_ 20121112
Time 20.29
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 165.63
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300104 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



C13CPD CDC13 {D:\NMR_DATA} RY 20



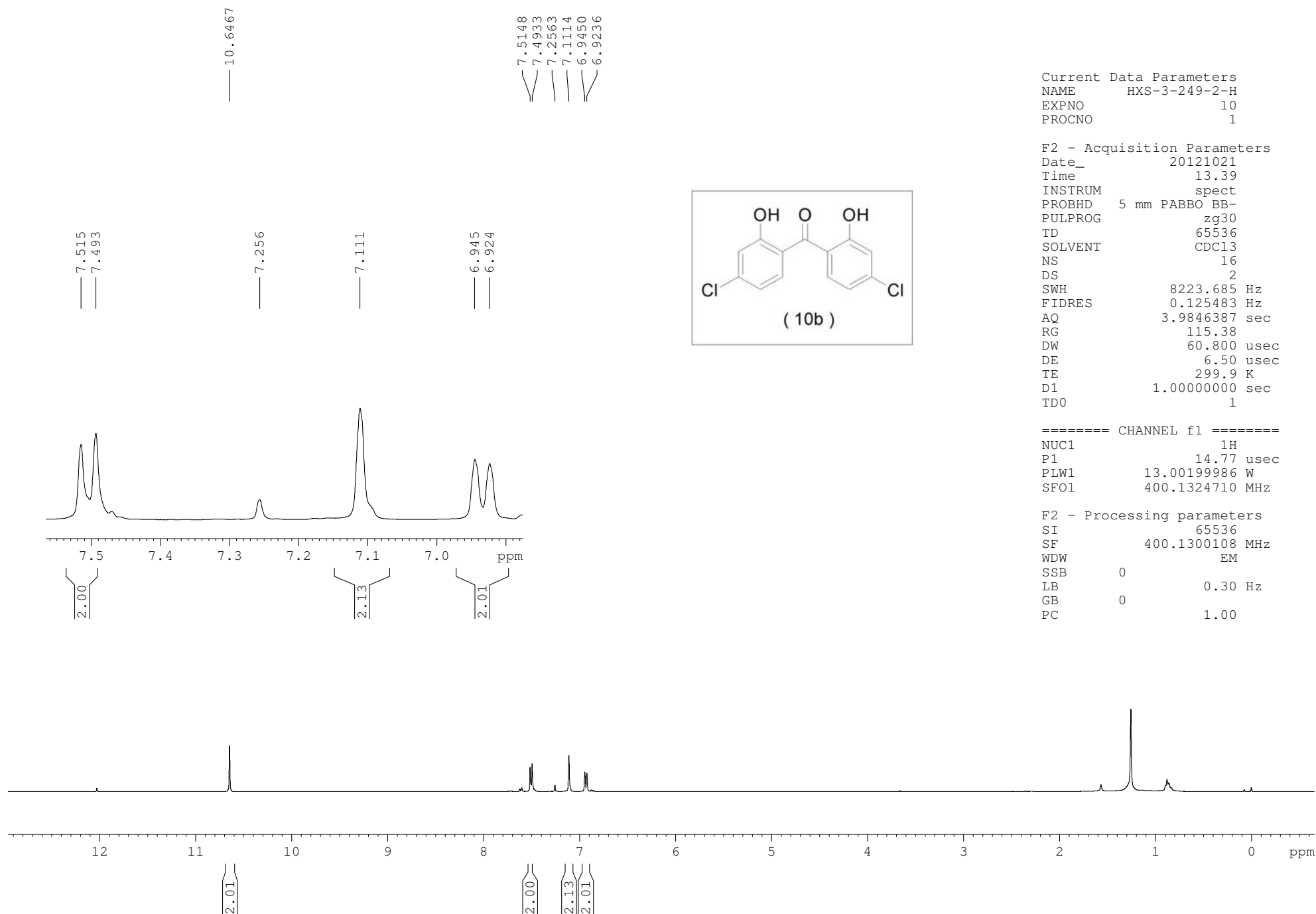
Current Data Parameters
NAME 20121112-YSY-1-151
EXPNO 20
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121112
Time 21.51
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

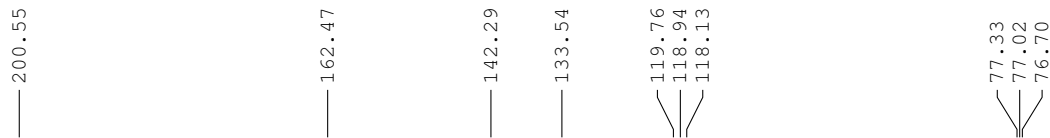
===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127690 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



C13CPD CDC13 {D:\NMR_DATA} RY 60



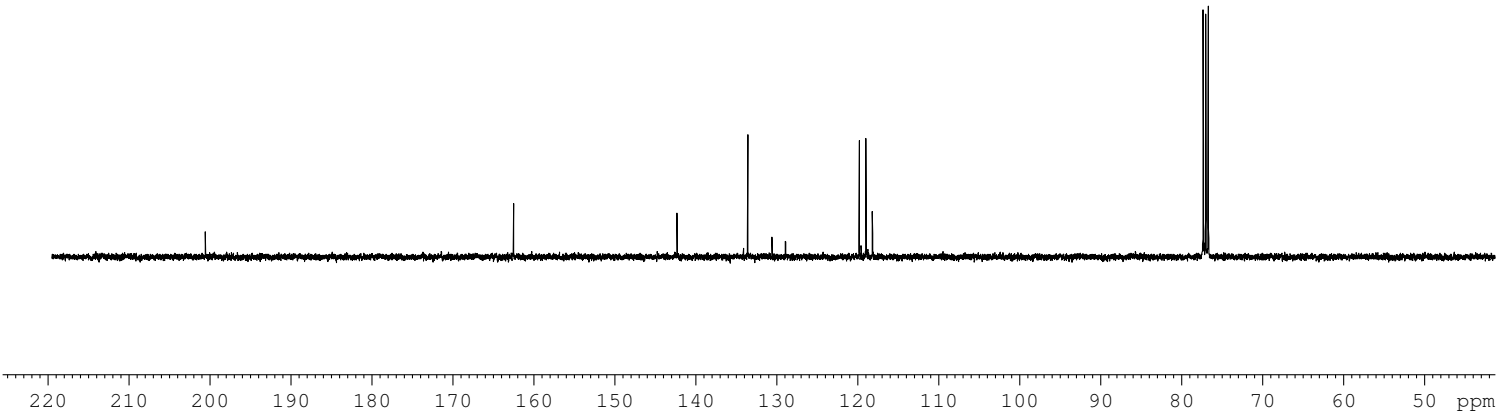
Current Data Parameters
NAME HXS-3-249-2-C
EXPNO 20
PROCNO 1

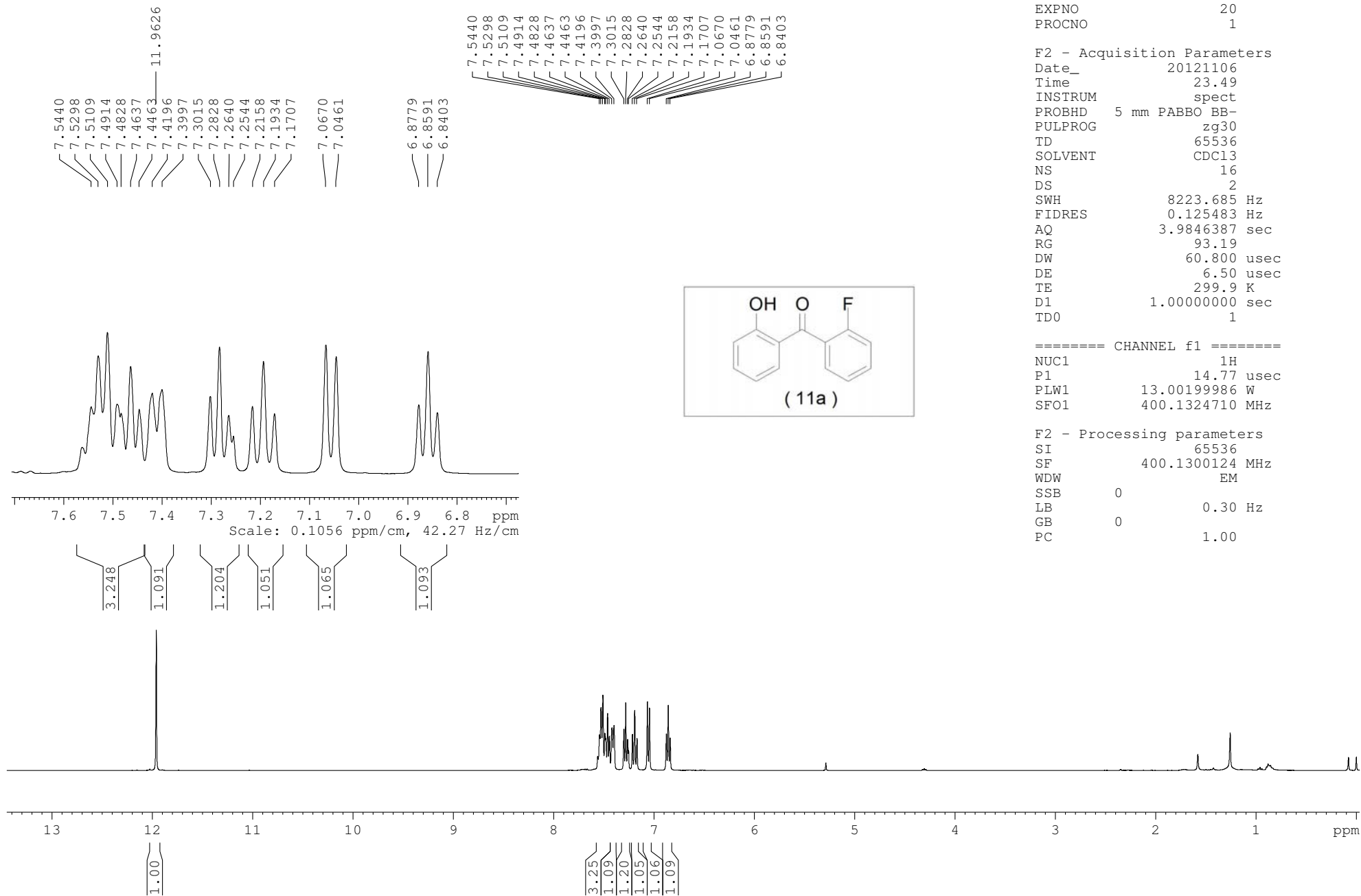
F2 - Acquisition Parameters
Date_ 20121021
Time 13.59
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 100
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

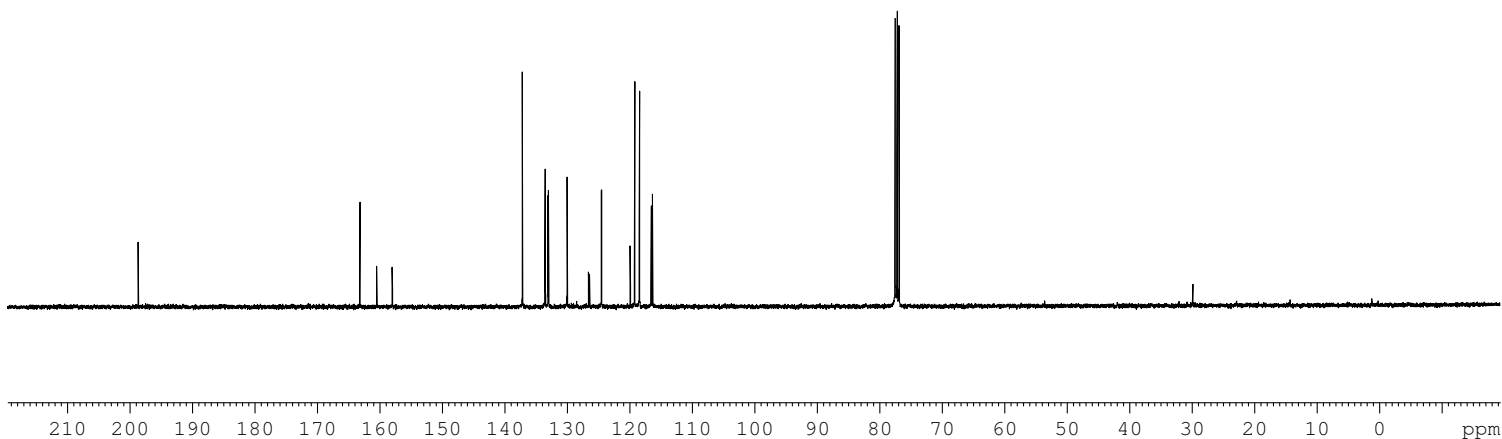
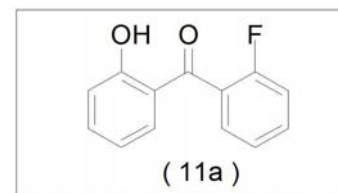
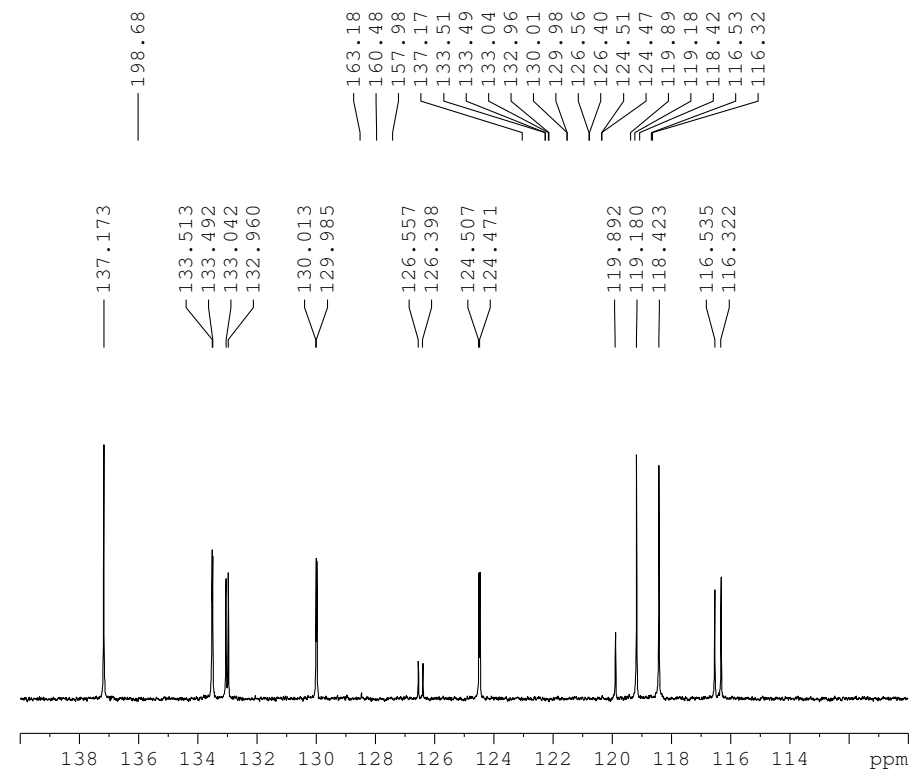
===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127690 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





C13CPD CDC13 {D:\NMR_DATA} RY 5



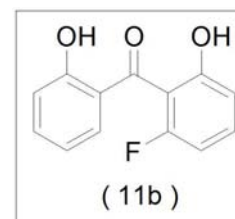
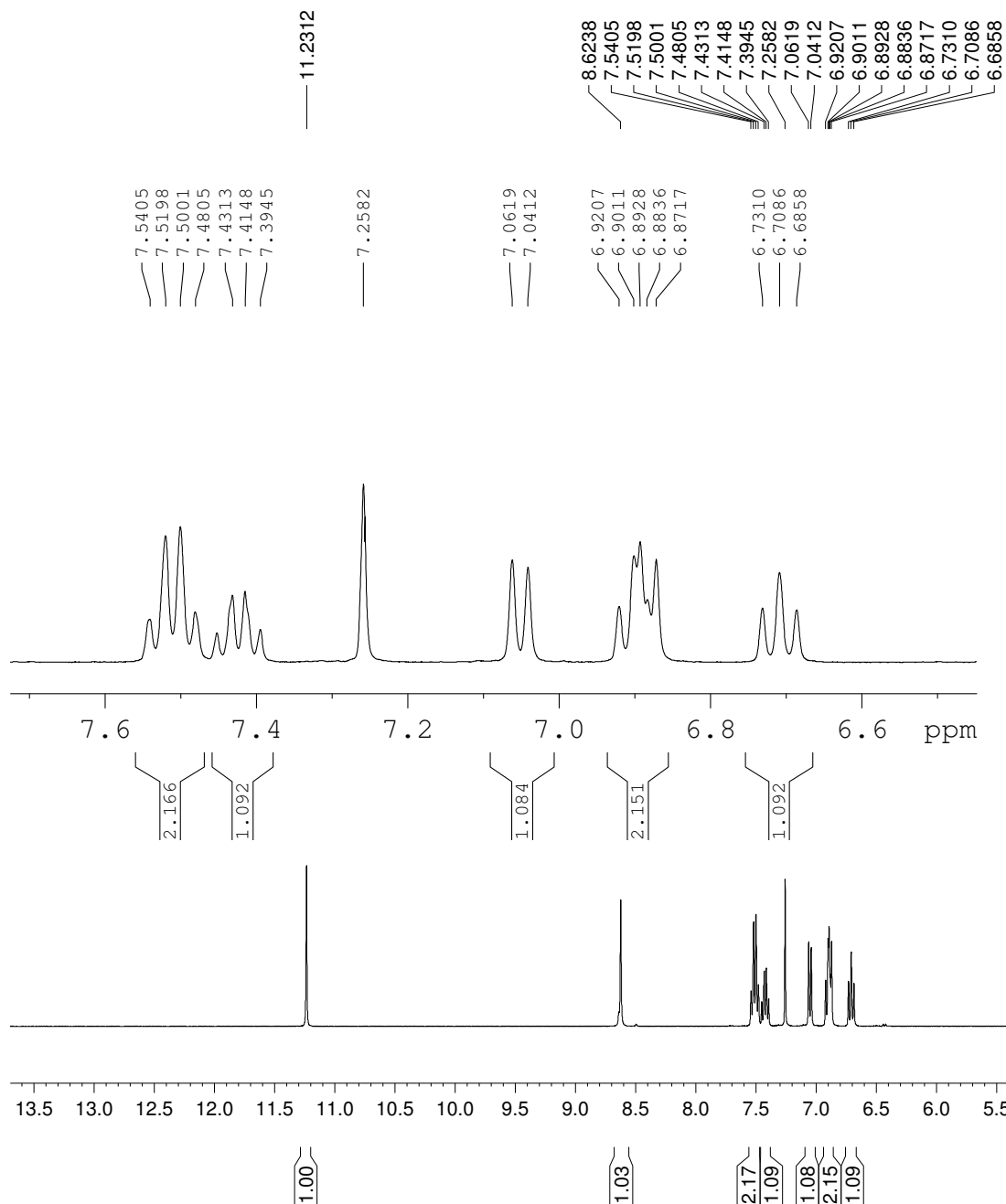
Current Data Parameters
NAME SG-7-275-M-H-20121113
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121113
Time 23.33
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127572 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



Current Data Parameters
NAME SG-7-276-D-H-20121119
EXPNO 10
PROCNO 1

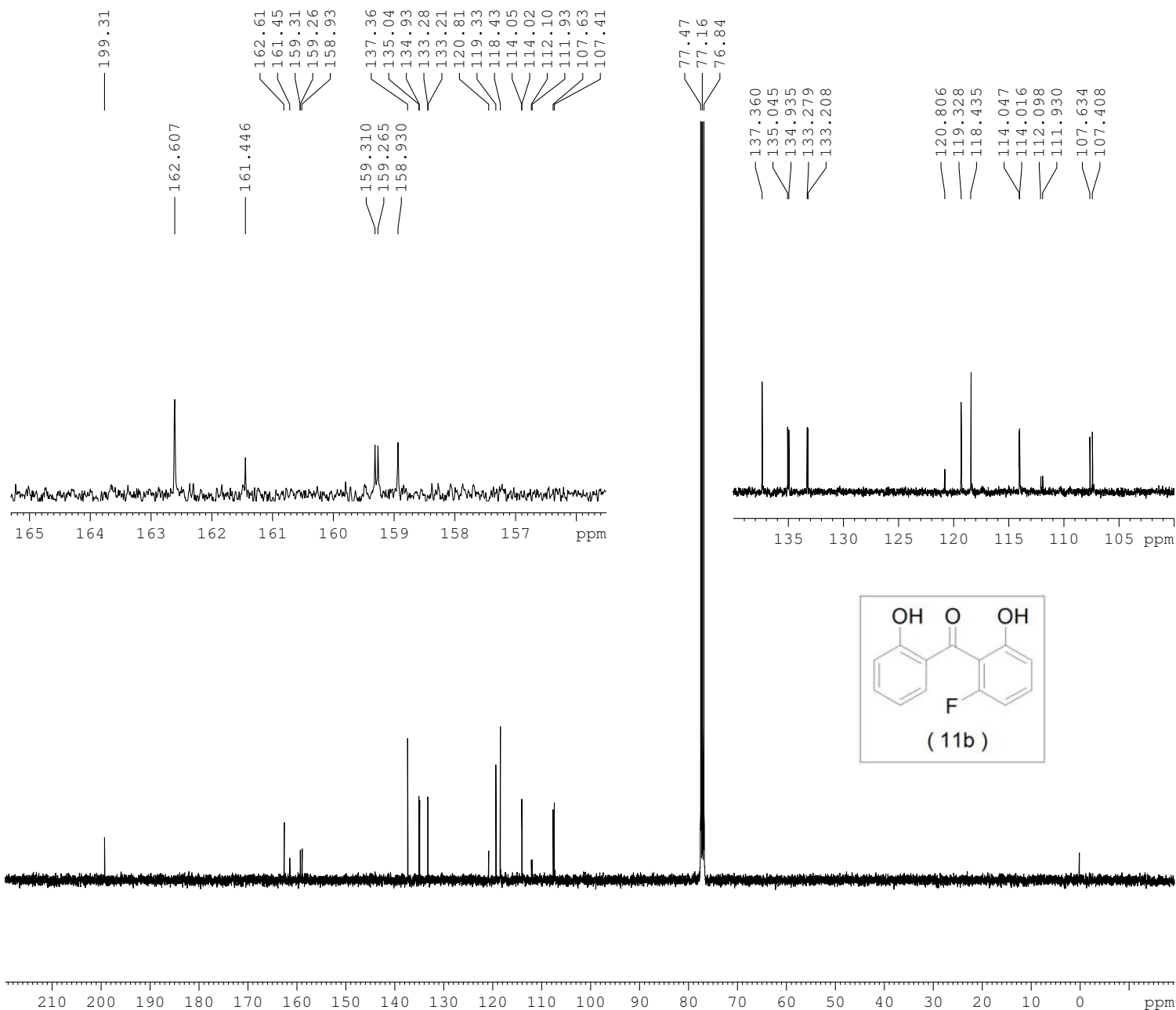
F2 - Acquisition Parameters
Date_ 20121119
Time 18.04
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl₃
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 209.25
DW 60.800 usec
DE 6.50 usec
TE 299.8 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SF01 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300109 MHz
WDW EM

Scale: 0.5747 ppm/cm, 230 Hz/cm

C13CPD CDC13 {D:\NMR_DATA} RY 41



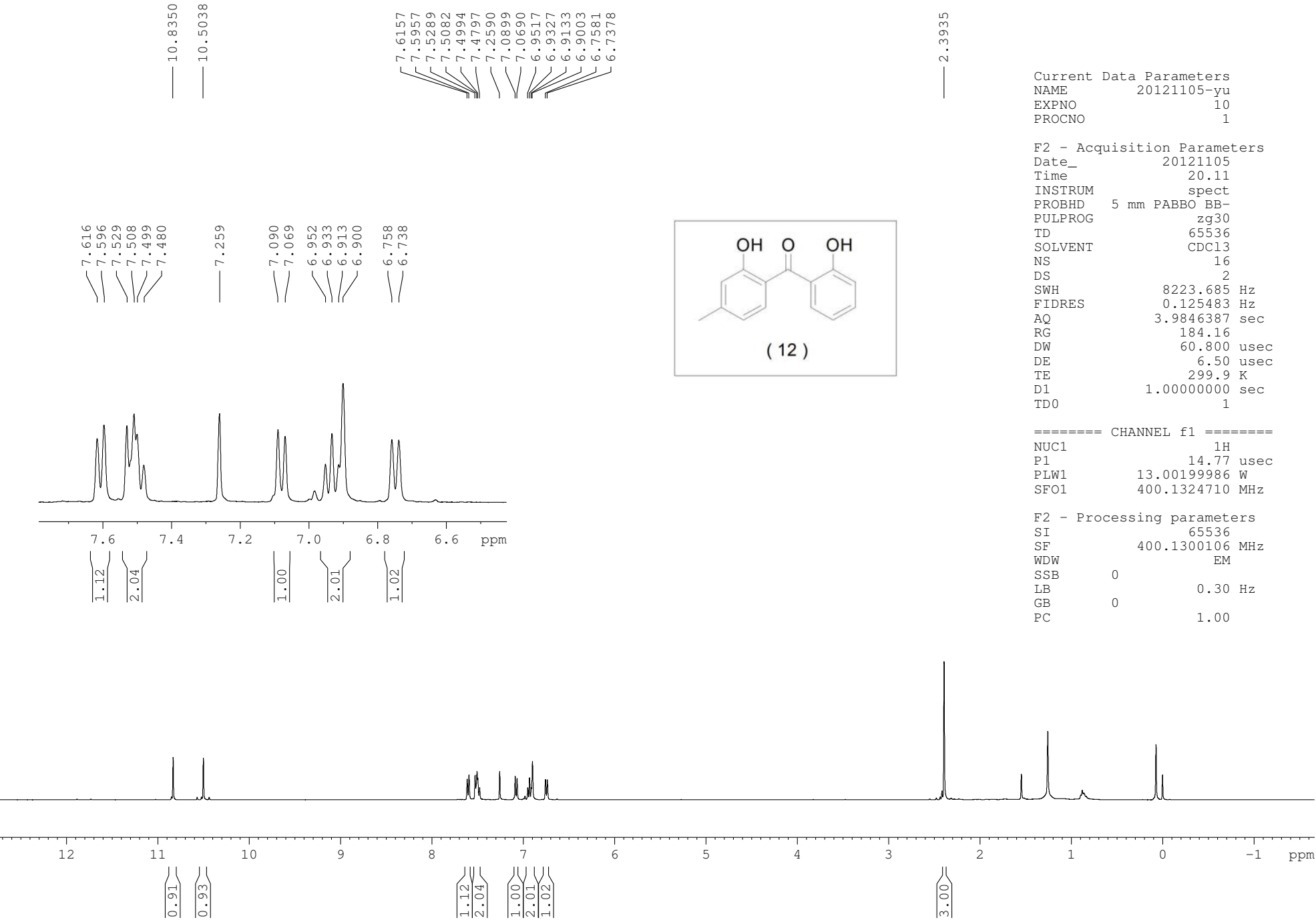
Current Data Parameters
NAME SG-7-275-D-C-20121121
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121121
Time 6.42
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 1600
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127542 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



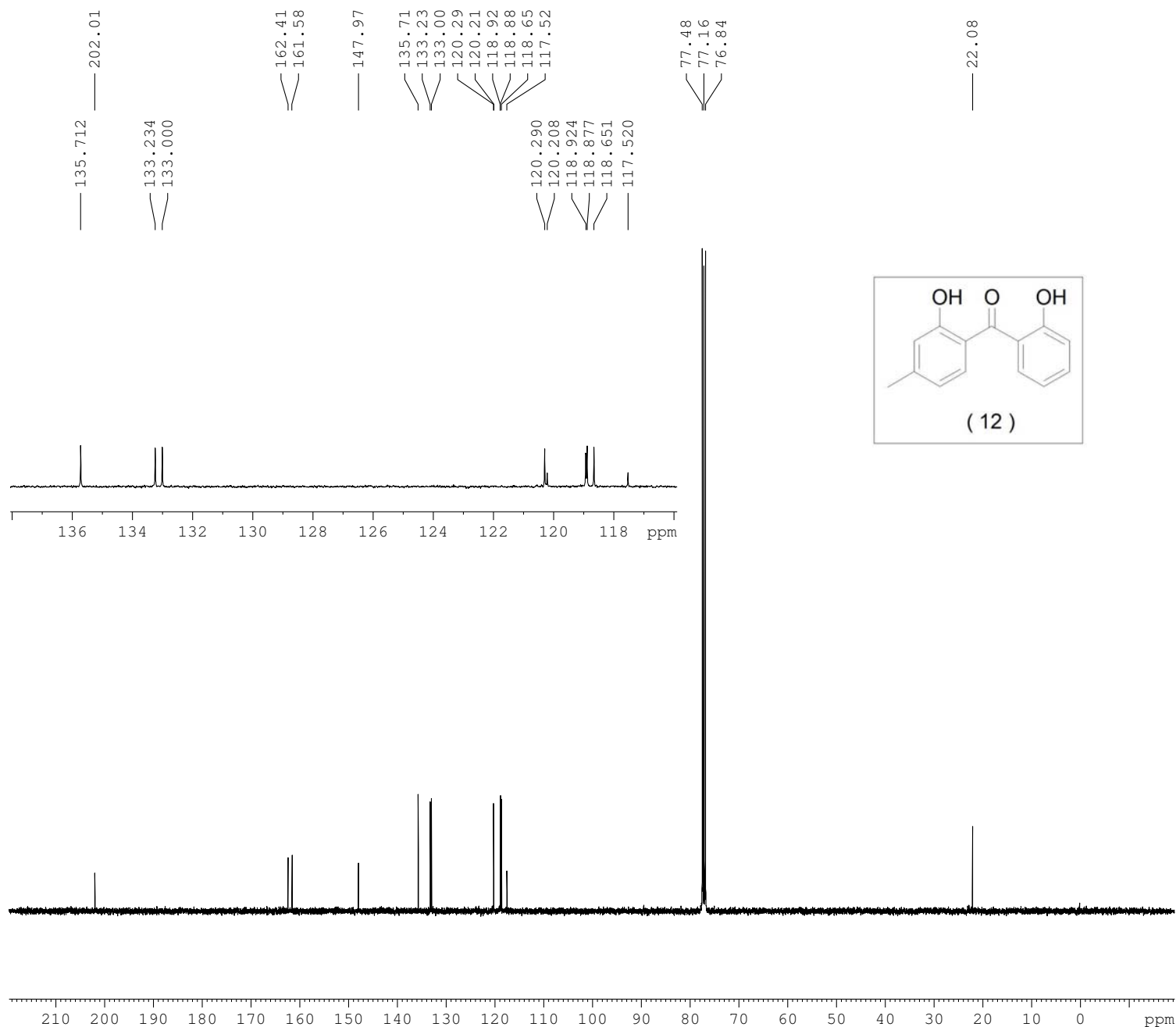
Current Data Parameters
NAME 20121105-yu
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121105
Time 20.11
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 184.16
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300106 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

C13CPD CDC13 {D:\NMR_DATA} RY 55



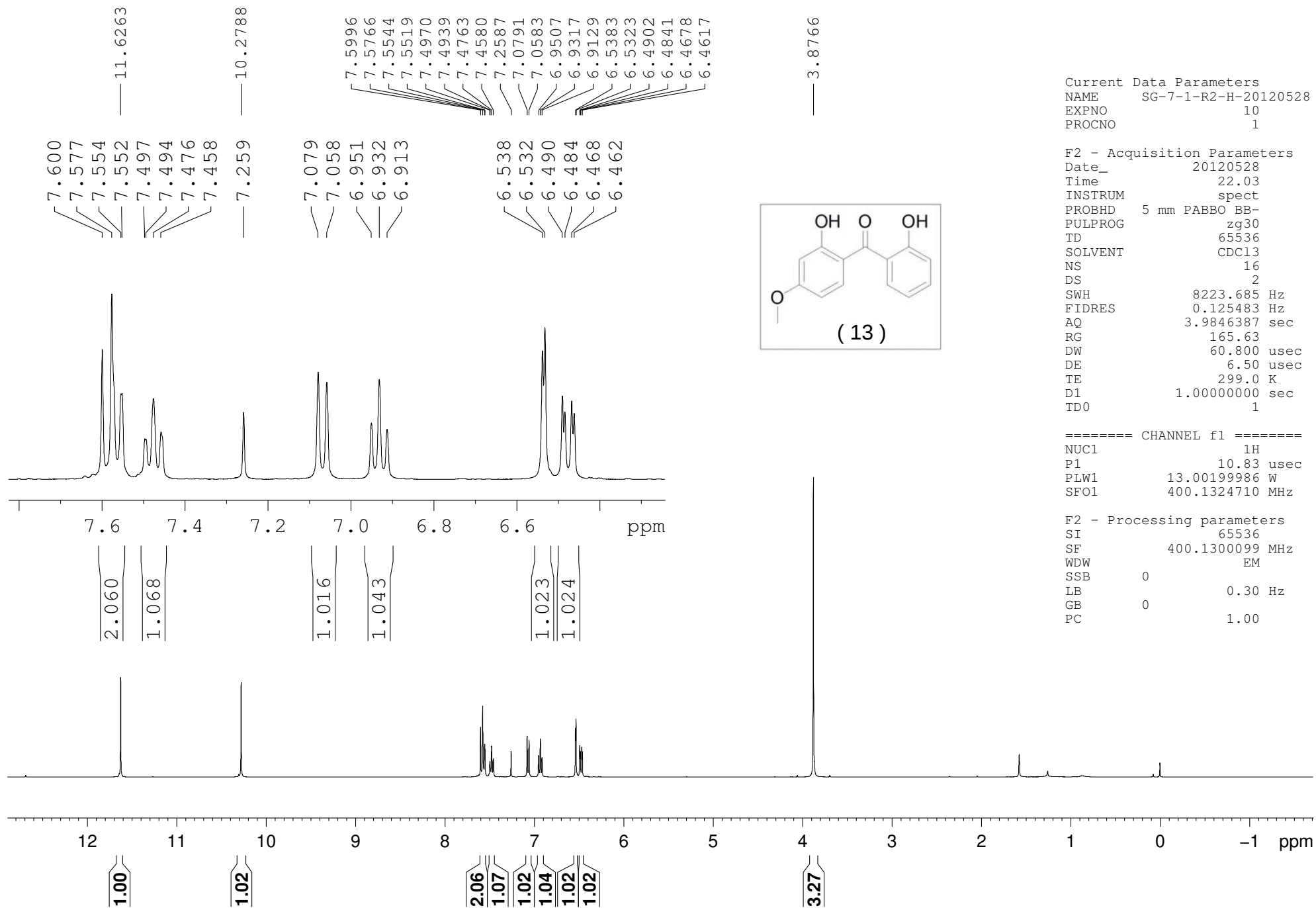
Current Data Parameters
NAME 20121107-yu-c
EXPNO 20
PROCNO 1

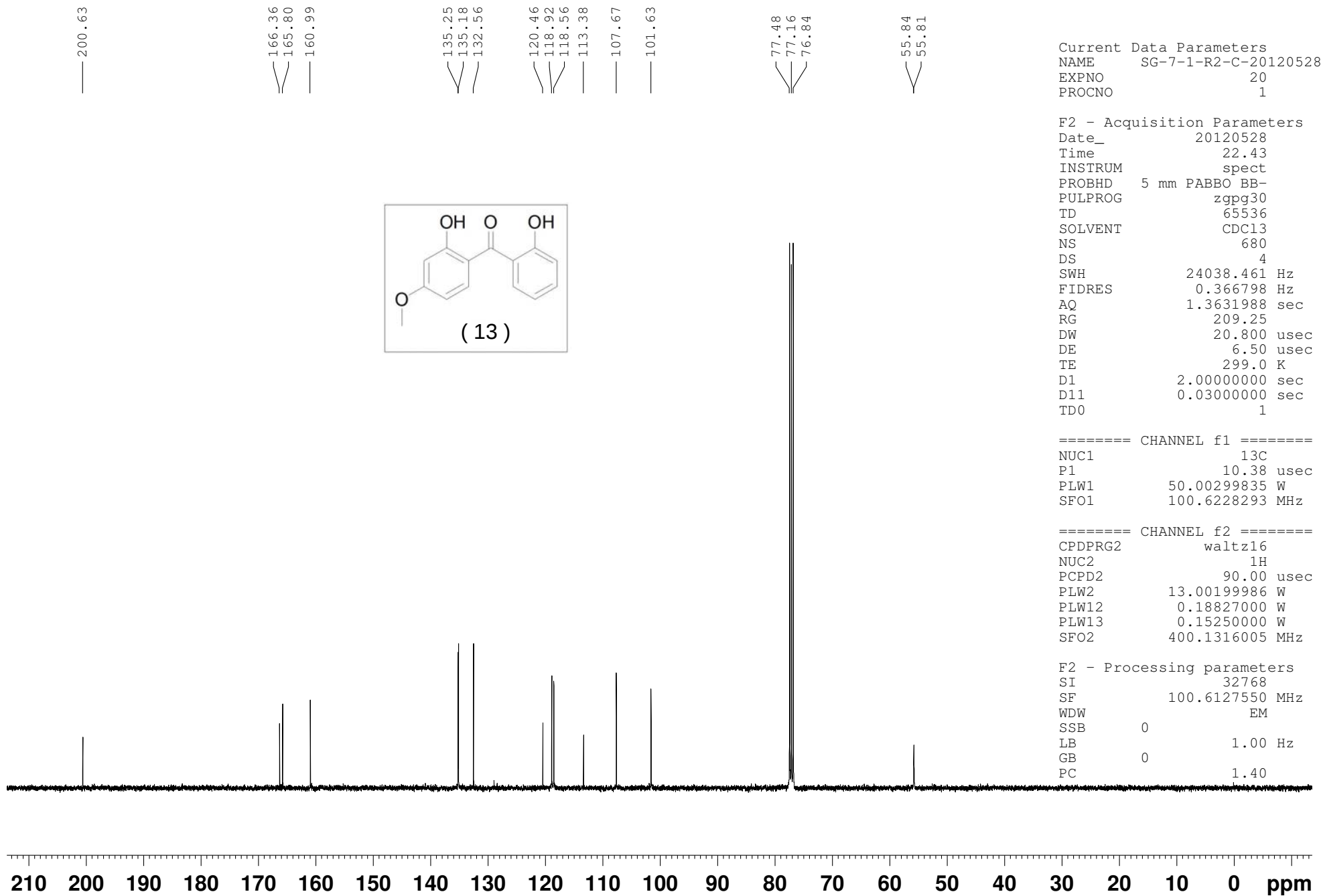
F2 - Acquisition Parameters
Date_ 20121107
Time 12.33
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

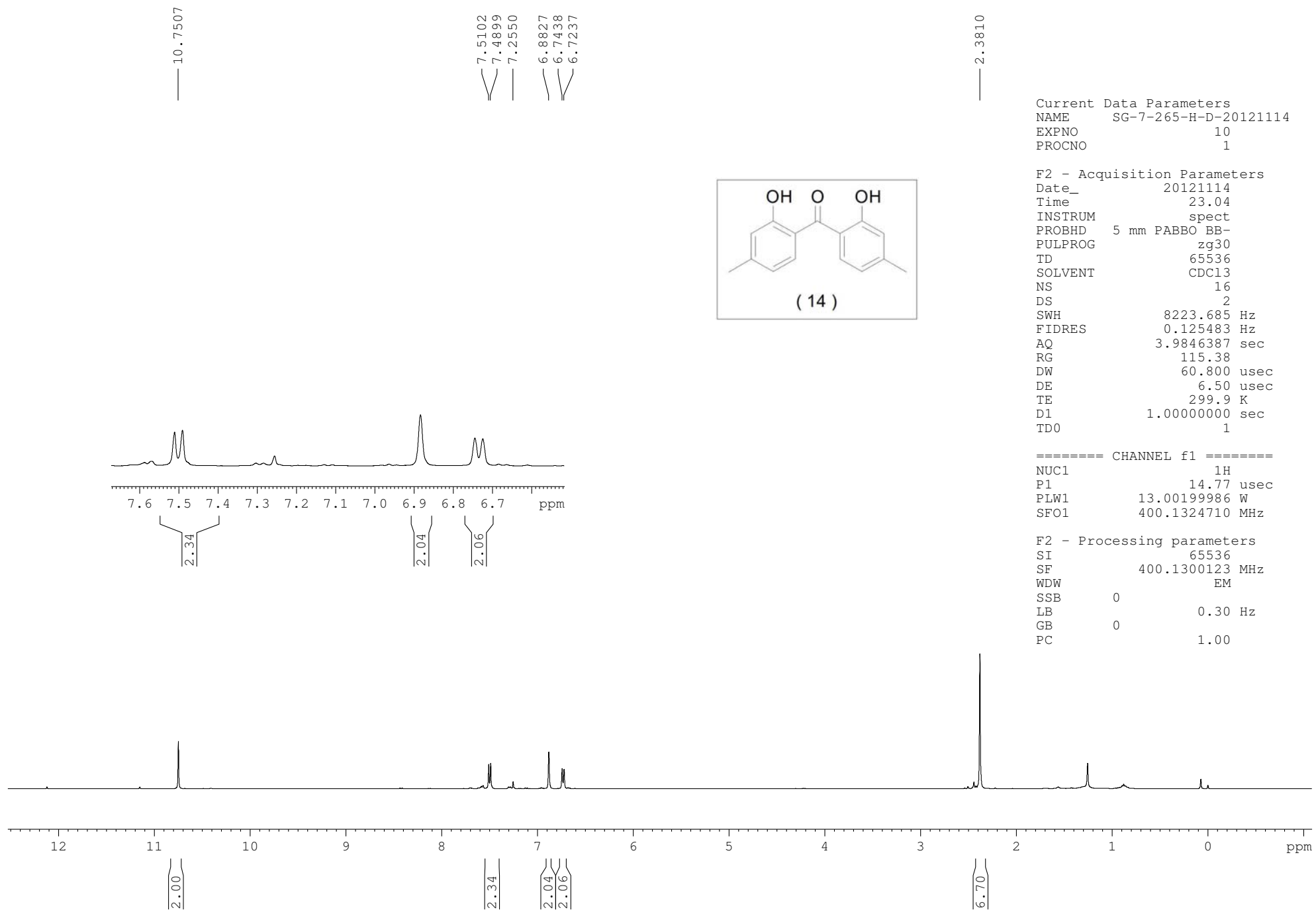
===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127546 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

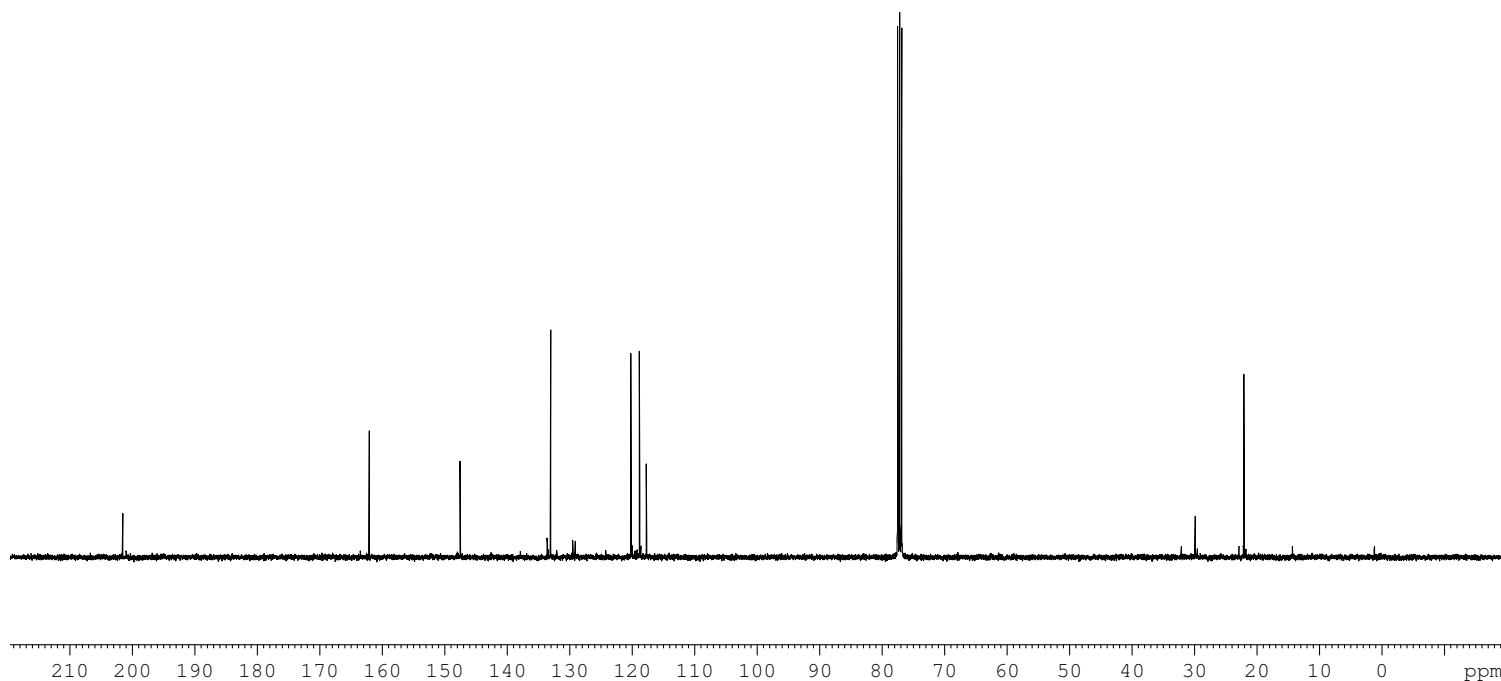
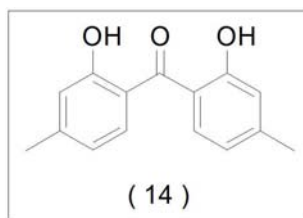






C13CPD CDC13 {D:\NMR_DATA} RY 11

— 201.55 — 162.08 — 147.52 — 133.04 — 120.17 — 118.80 — 117.69 — 77.48 — 77.16 — 76.84 — 22.03



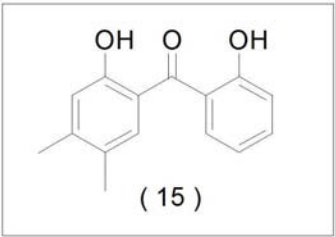
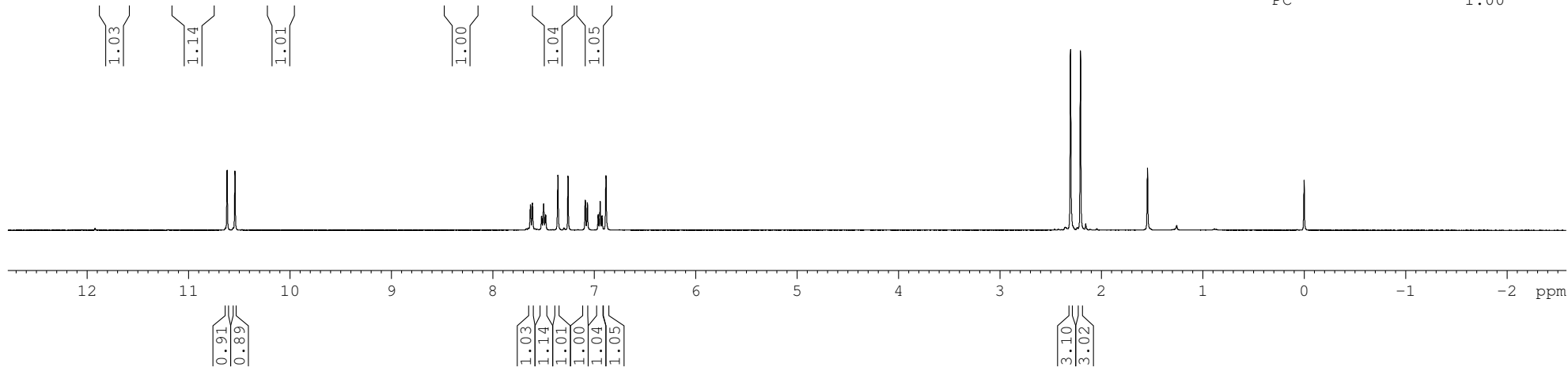
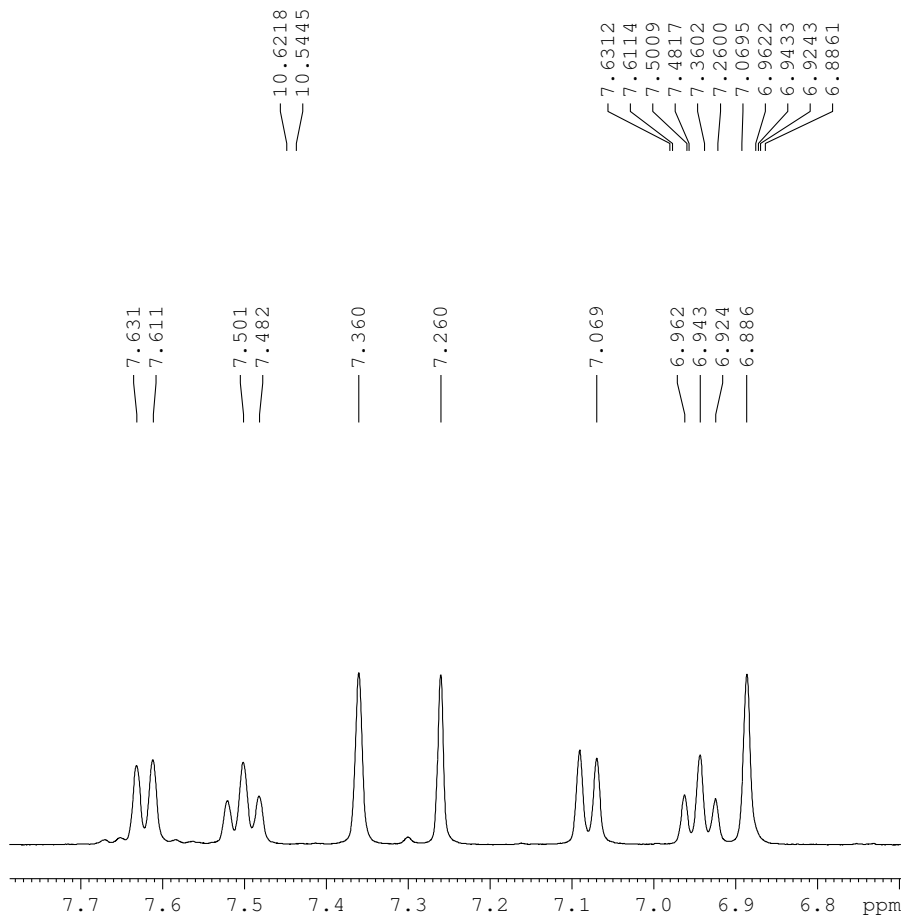
Current Data Parameters
NAME SG-7-265-D-H-20121115
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121115
Time 15.03
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 184.16
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127552 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



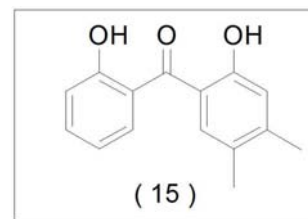
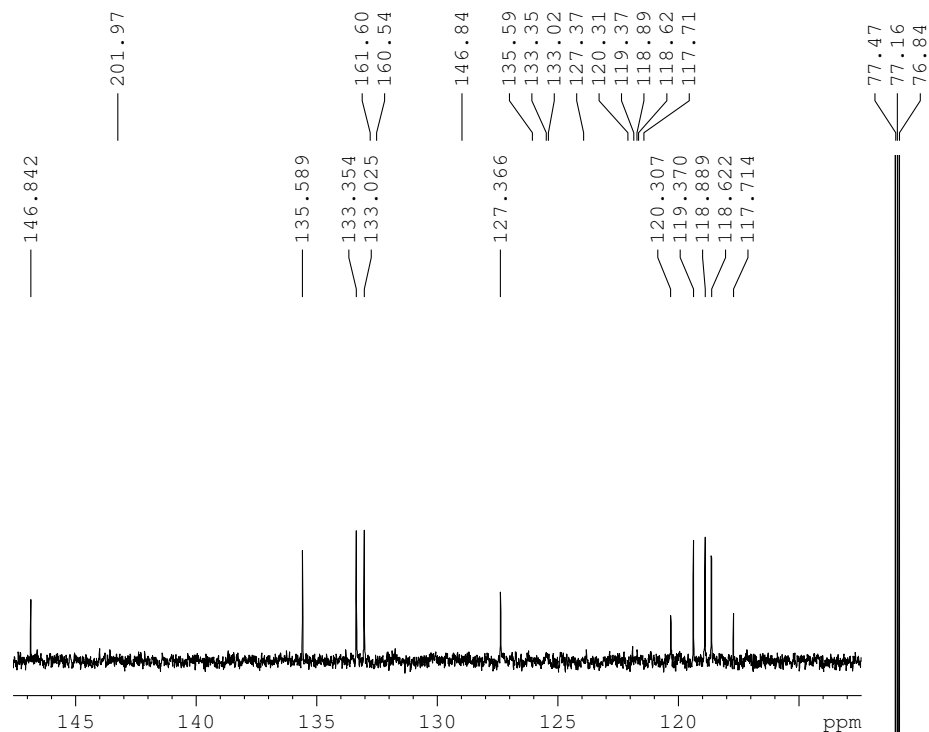
Current Data Parameters
NAME 20121107-yu
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121107
Time 22.24
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 184.16
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300103 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

C13CPD CDC13 {D:\NMR_DATA} RY 49



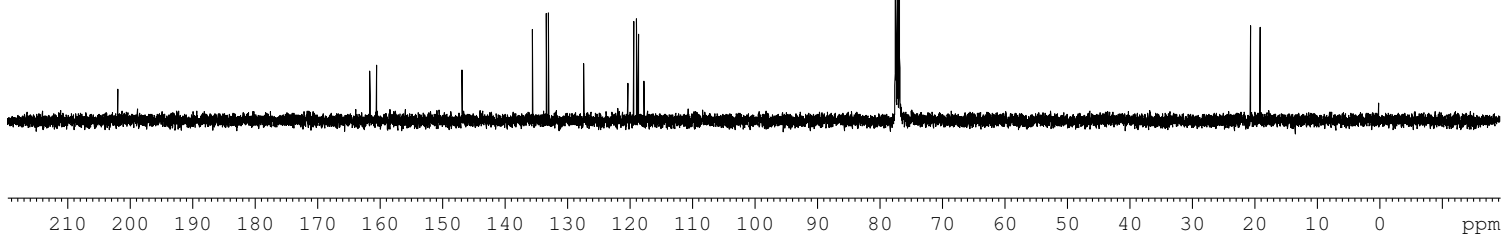
Current Data Parameters
NAME 20121107-yu
EXPNO 11
PROCNO 1

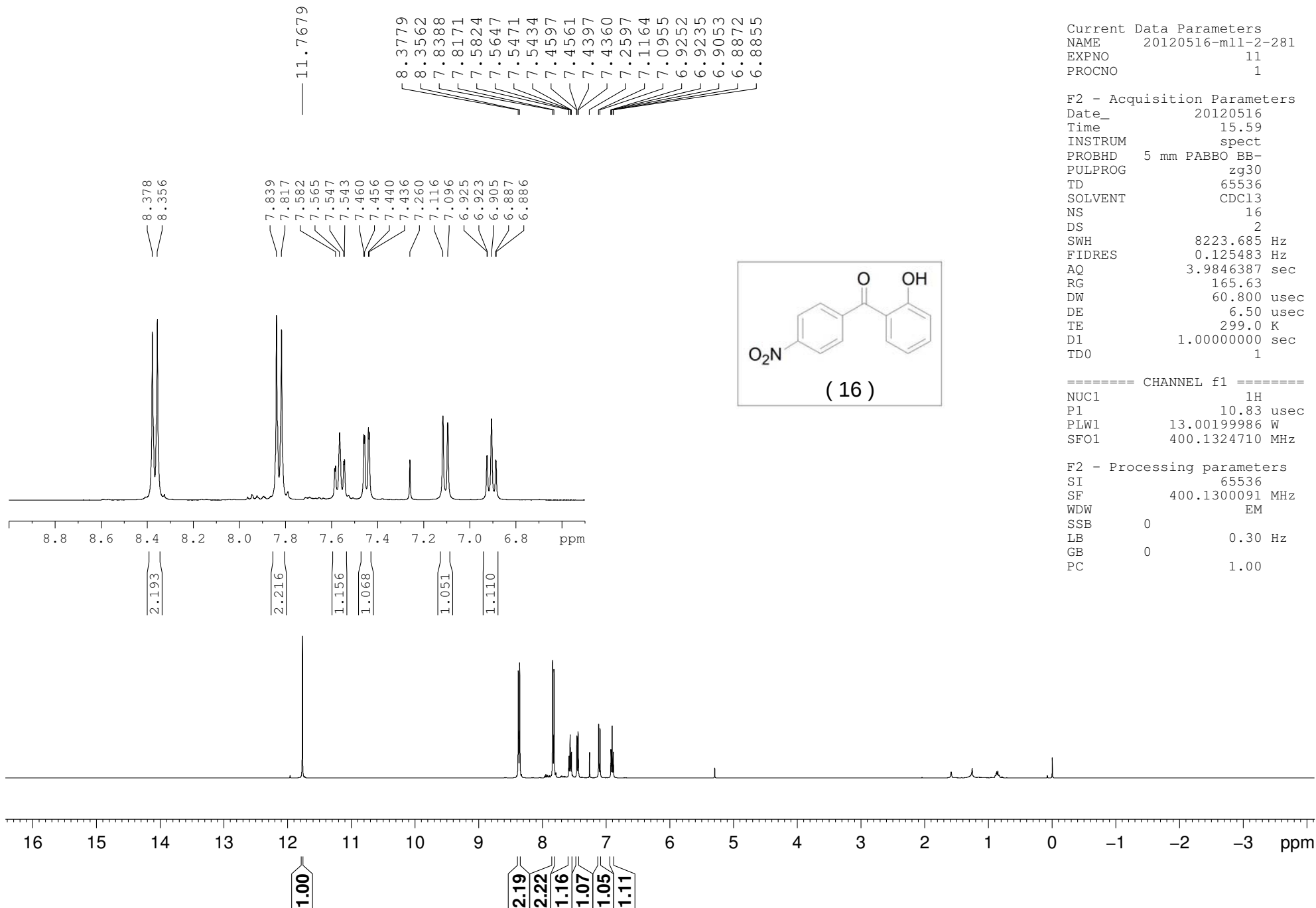
F2 - Acquisition Parameters
Date_ 20121108
Time 2.10
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

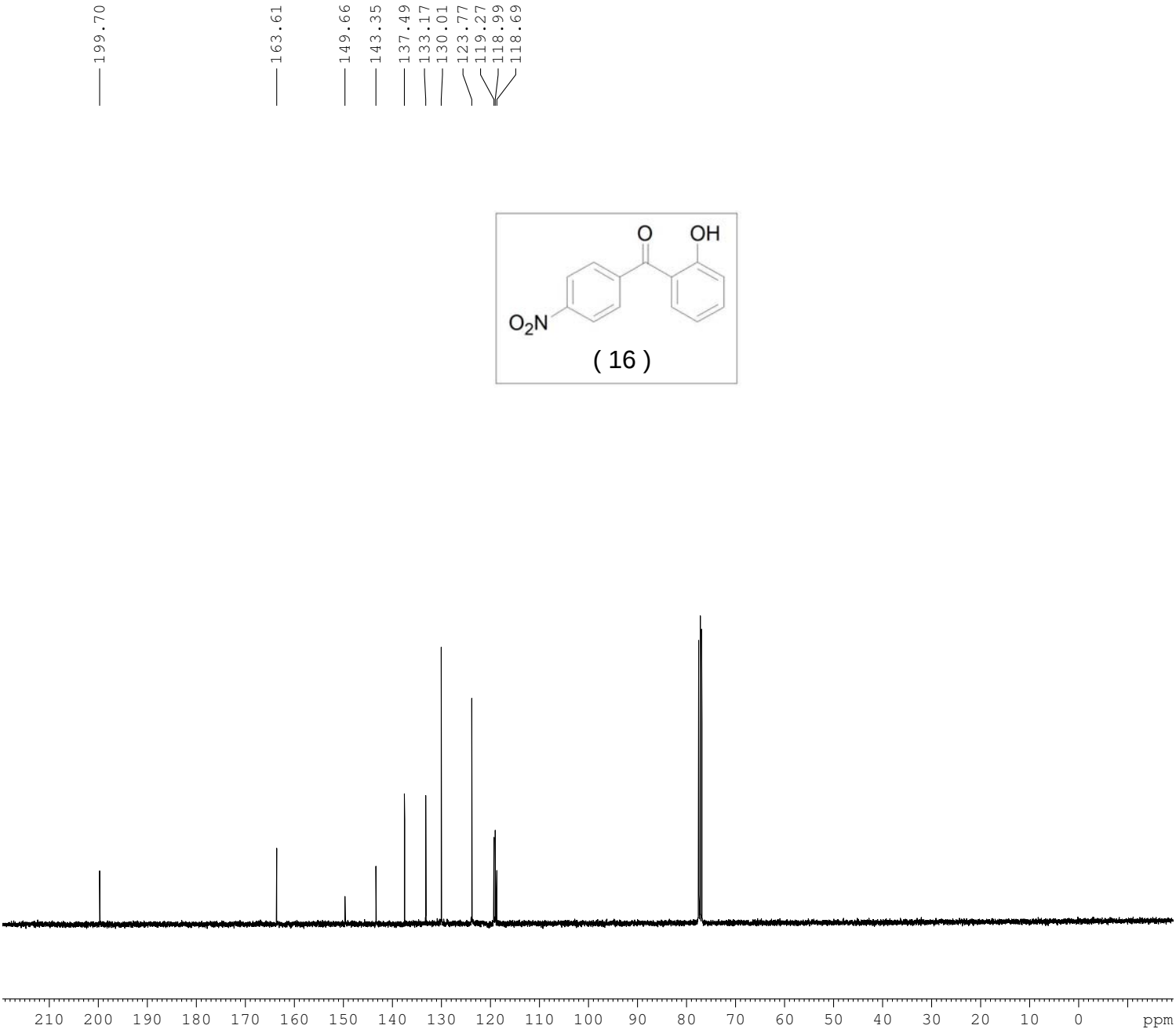
===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127541 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40







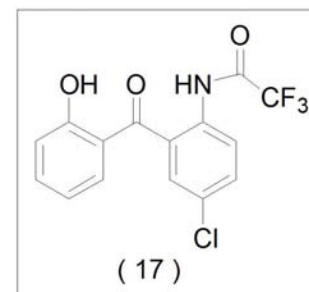
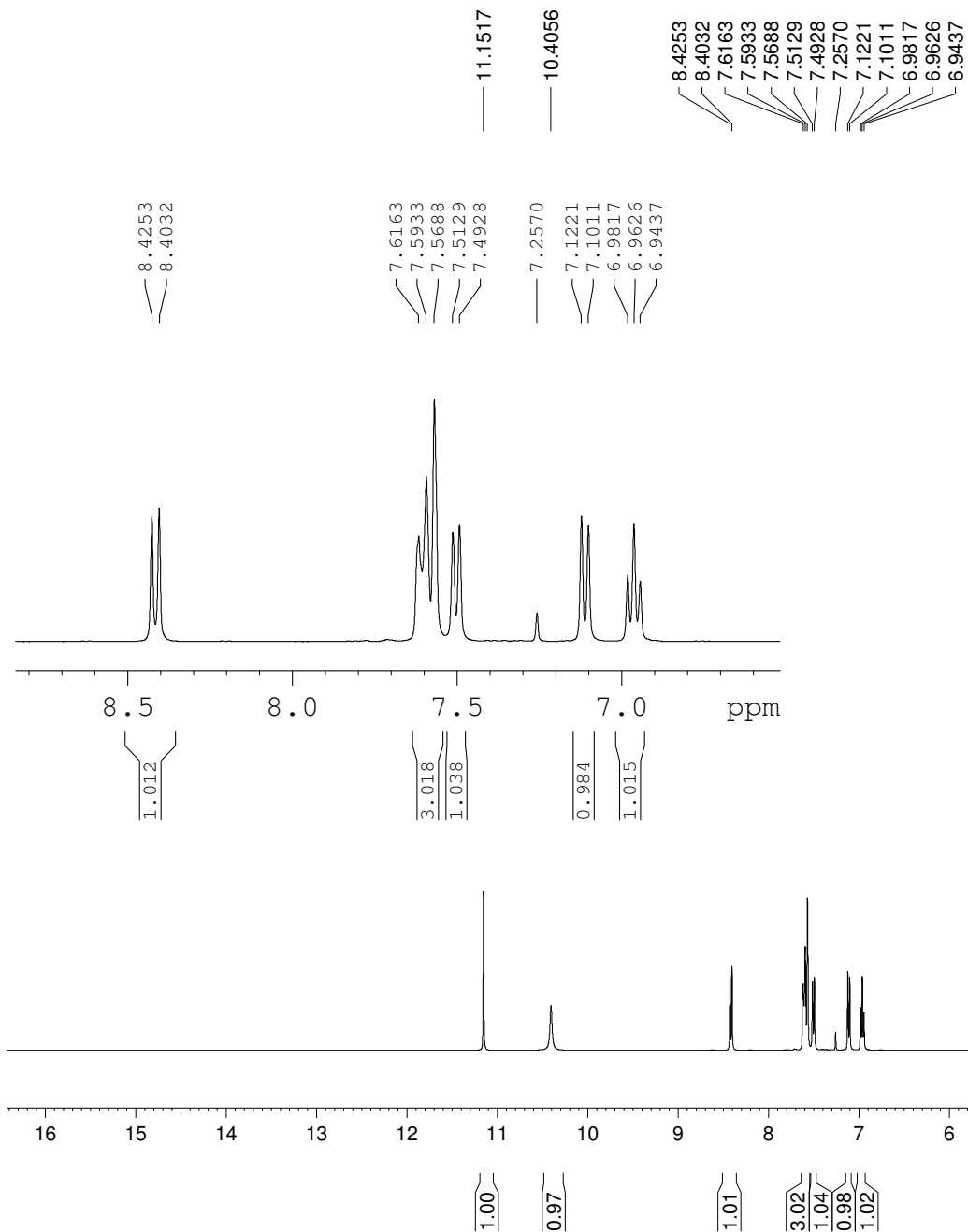
Current Data Parameters
NAME 20120516-ml1-2-281-c
EXPNO 12
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120516
Time 16.50
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 203
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.0 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.38 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.18827000 W
PLW13 0.15250000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127564 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



Current Data Parameters
NAME SG-7-261-H-20121106
EXPNO 20
PROCNO 1

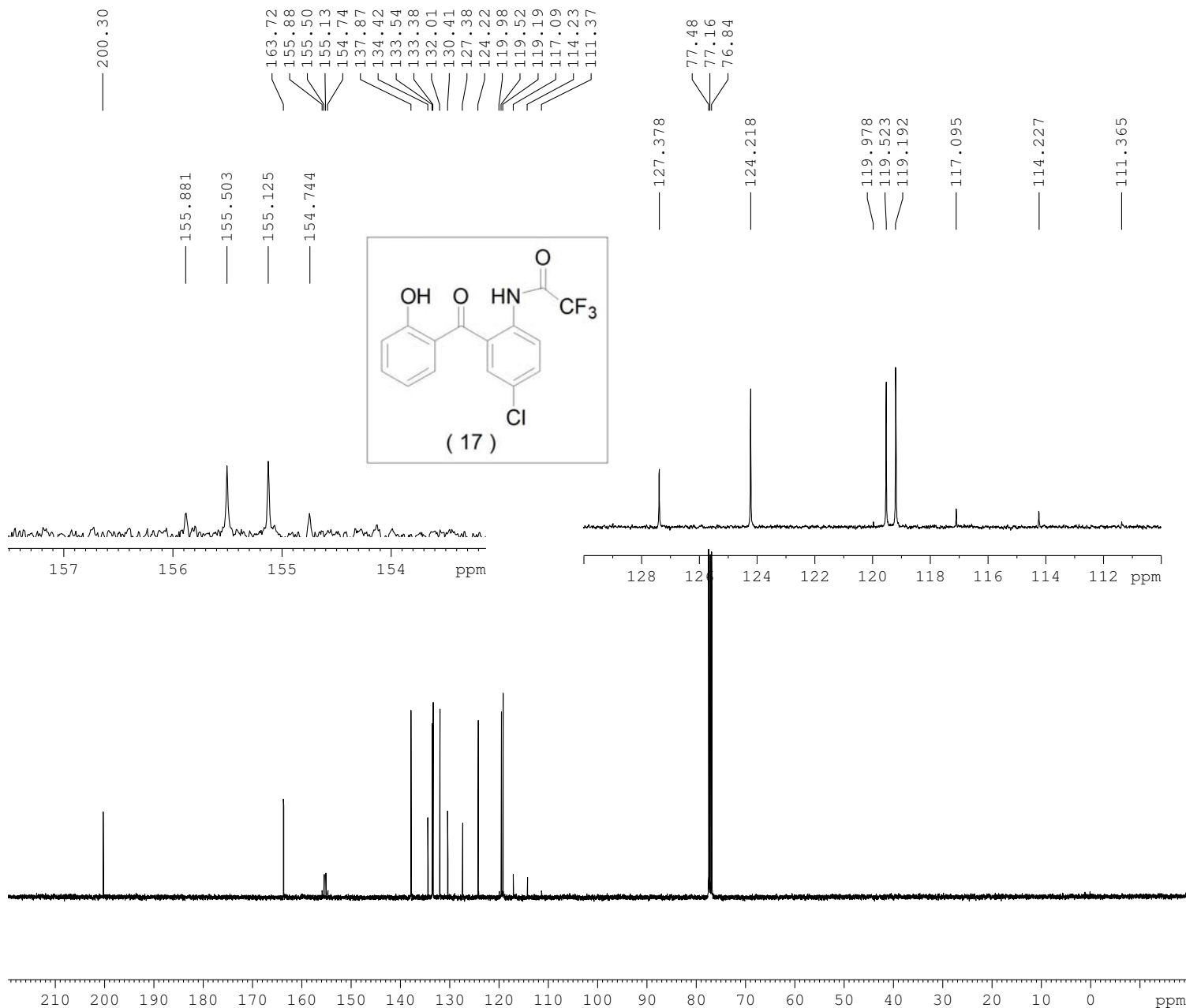
F2 - Acquisition Parameters
Date_ 20121106
Time 22.21
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDC13
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 93.19
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SF01 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300114 MHz
WDW EM

Scale: 0.7824 ppm/cm, 313 Hz/cm

C13CPD CDC13 {D:\NMR_DATA} RY 36



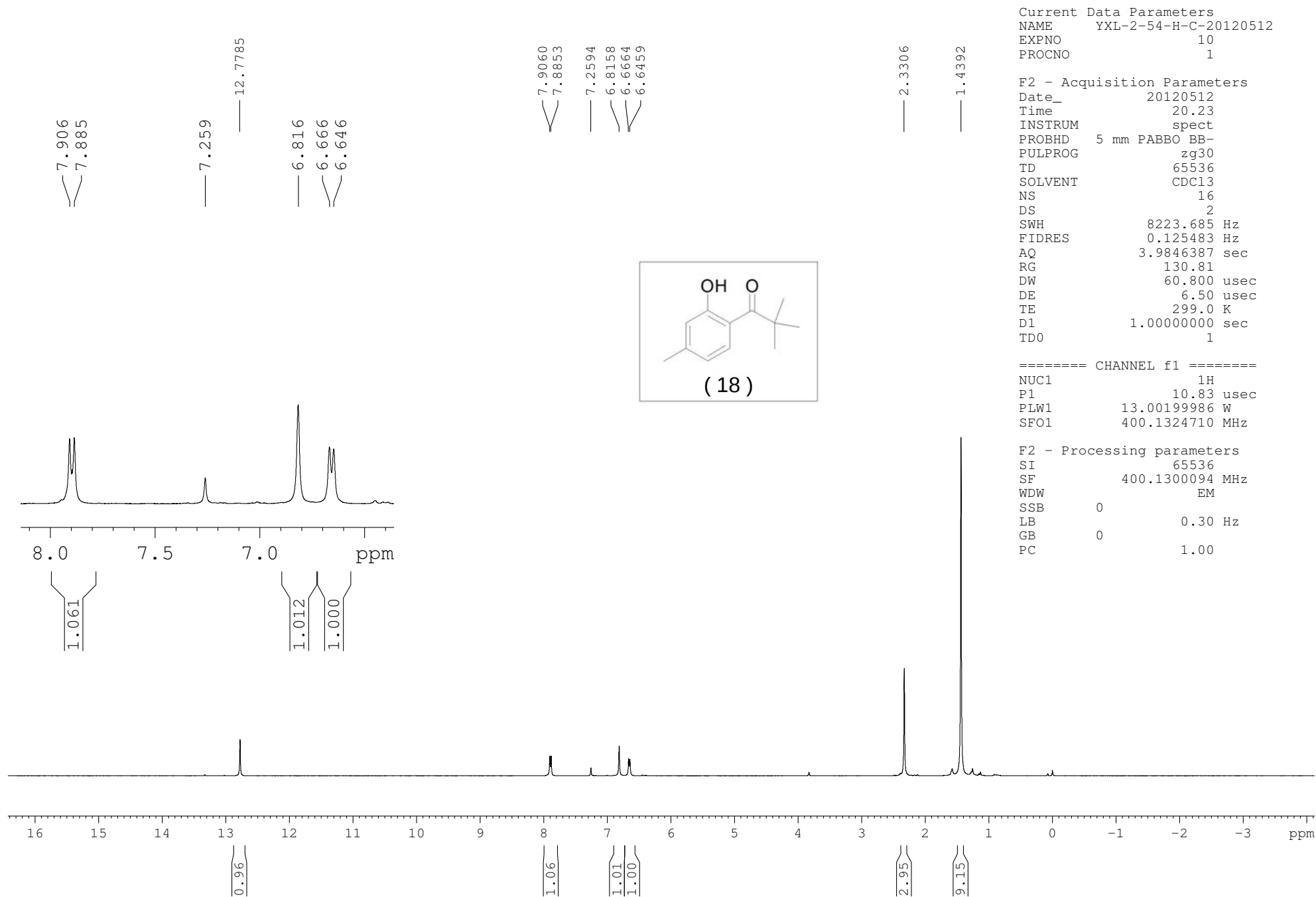
Current Data Parameters
NAME SG-7-261-C-20121108
EXPNO 10
PROCNO 1

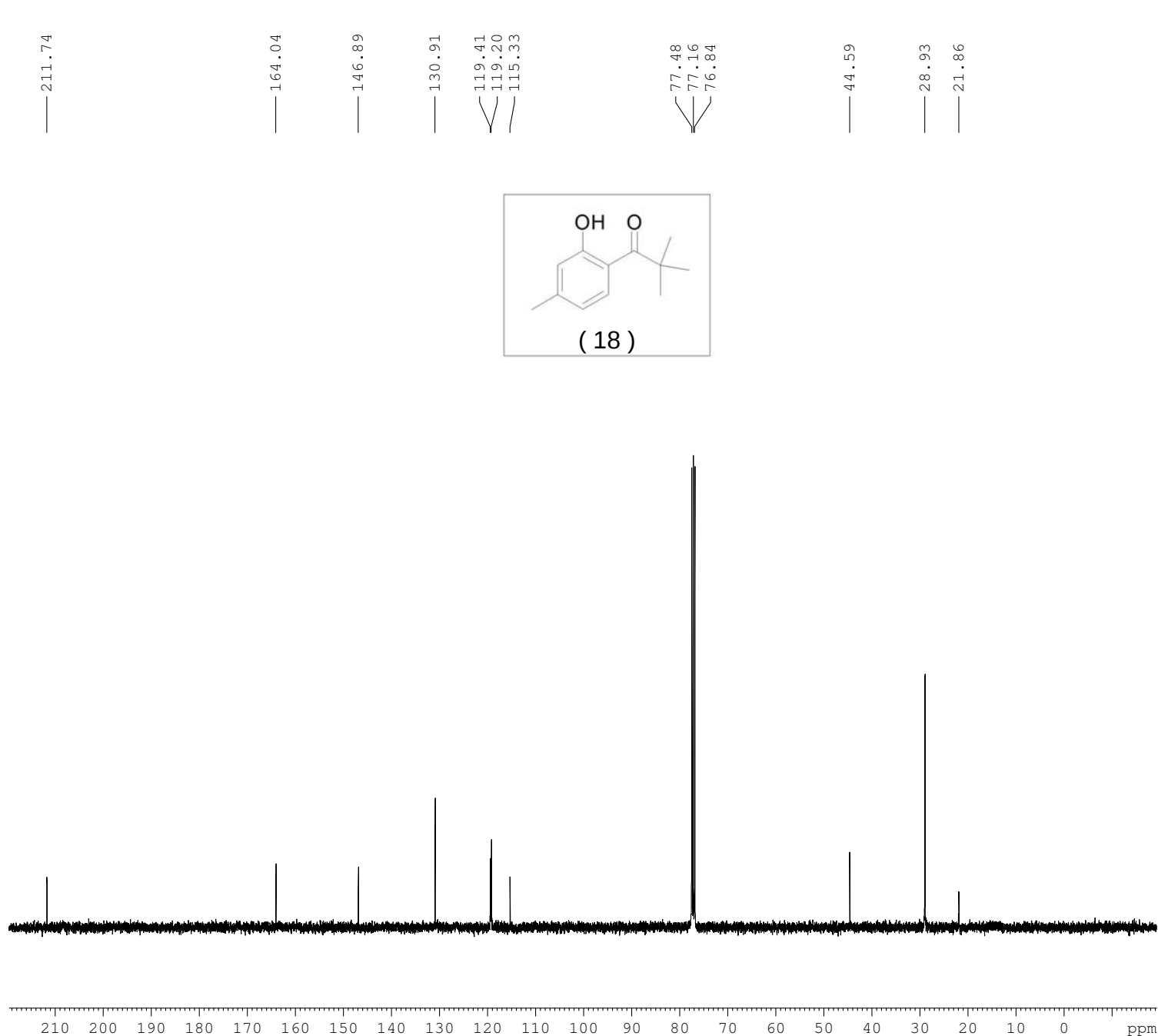
F2 - Acquisition Parameters
Date_ 20121108
Time 23.18
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 256
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127556 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





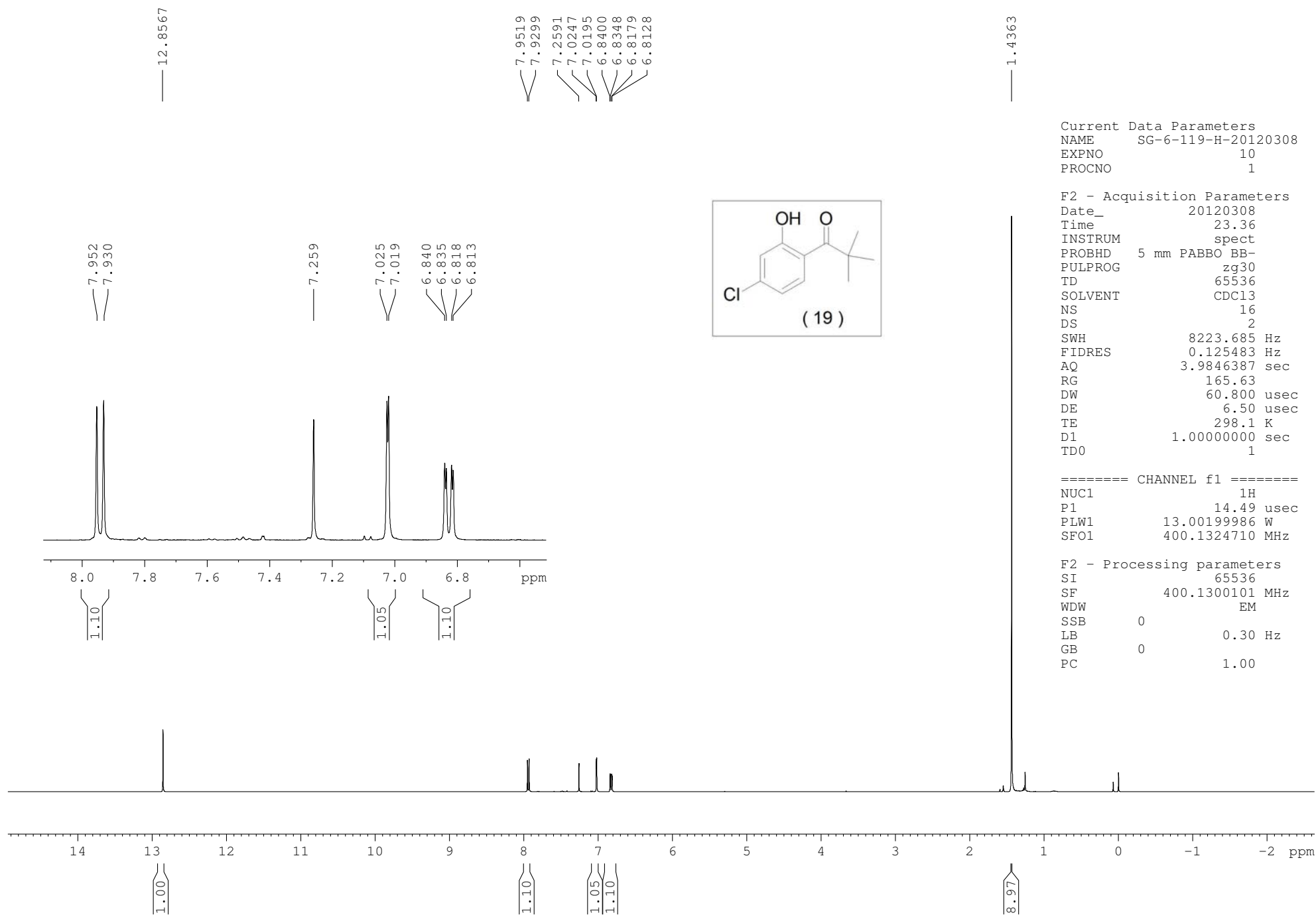
Current Data Parameters
NAME YXL-2-21-C-20120512
EXPNO 20
PROCNO 1

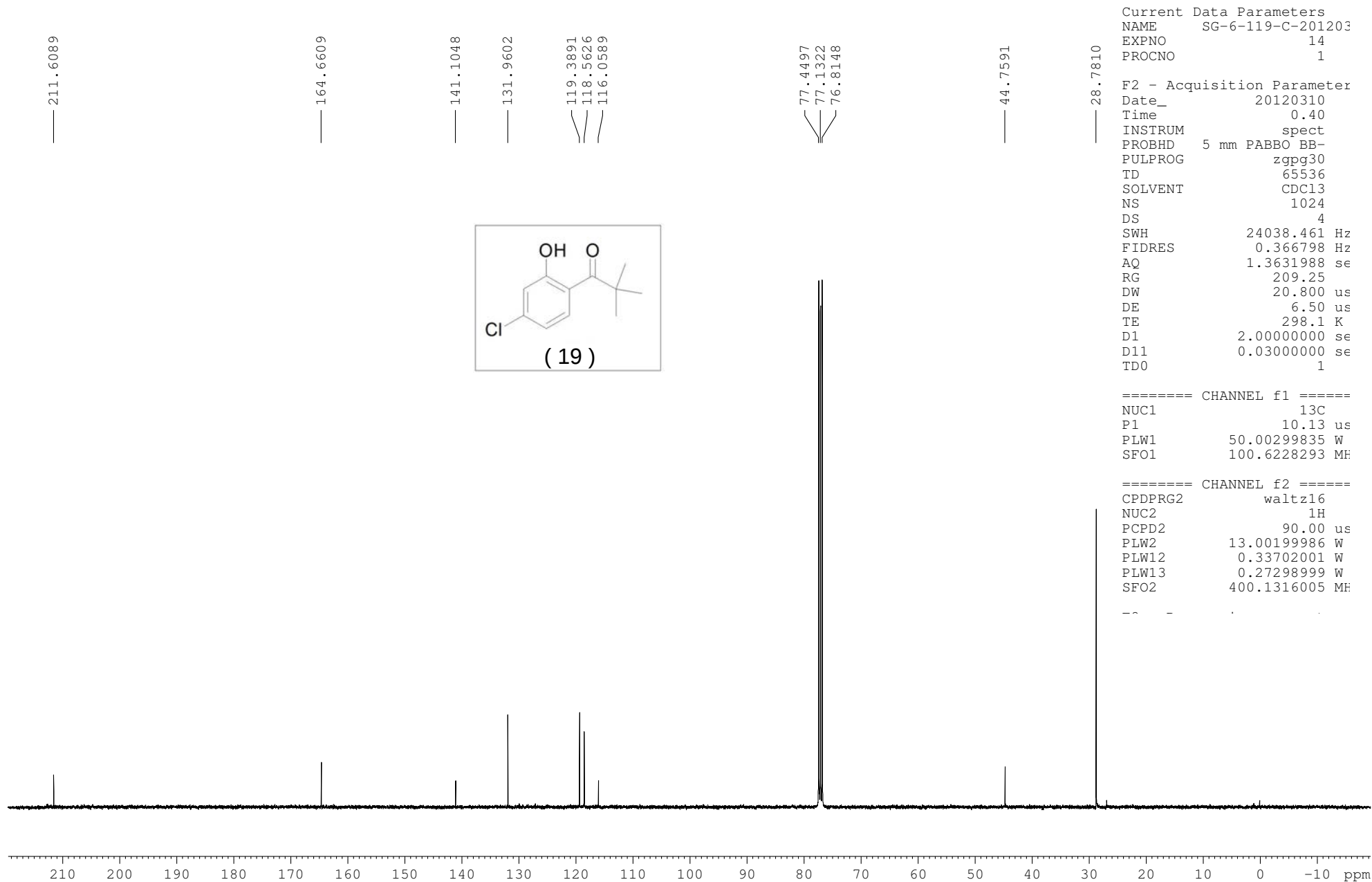
F2 - Acquisition Parameters
Date_ 20120512
Time 20.53
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 256
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.0 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

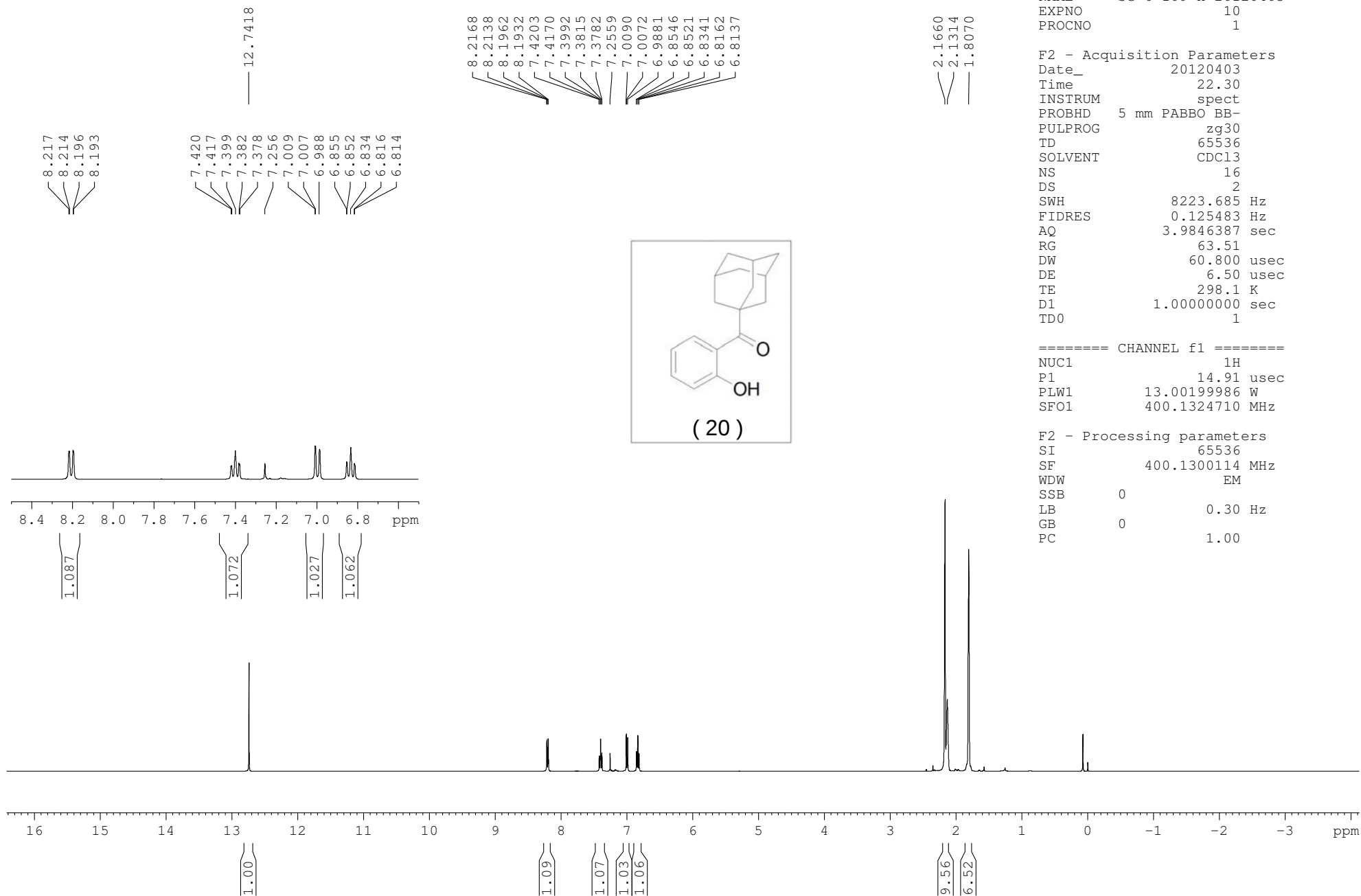
===== CHANNEL f1 =====
NUC1 13C
P1 10.38 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.18827000 W
PLW13 0.15250000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127547 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





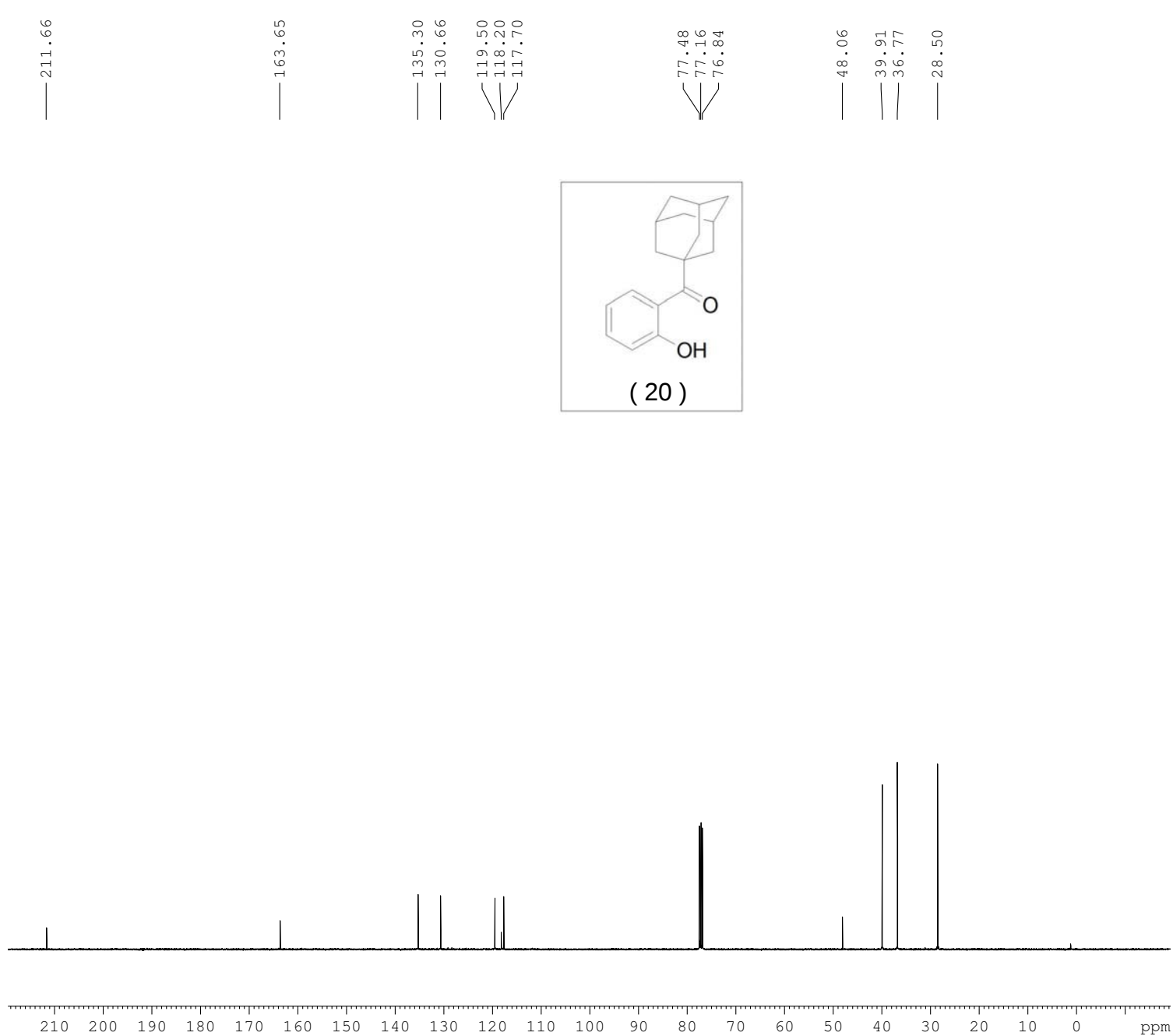


Current Data Parameters
NAME SG-6-109-H-20120403
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120403
Time 22.30
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDC13
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 63.51
DW 60.800 usec
DE 6.50 usec
TE 298.1 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.91 usec
PLW1 13.0019986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300114 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



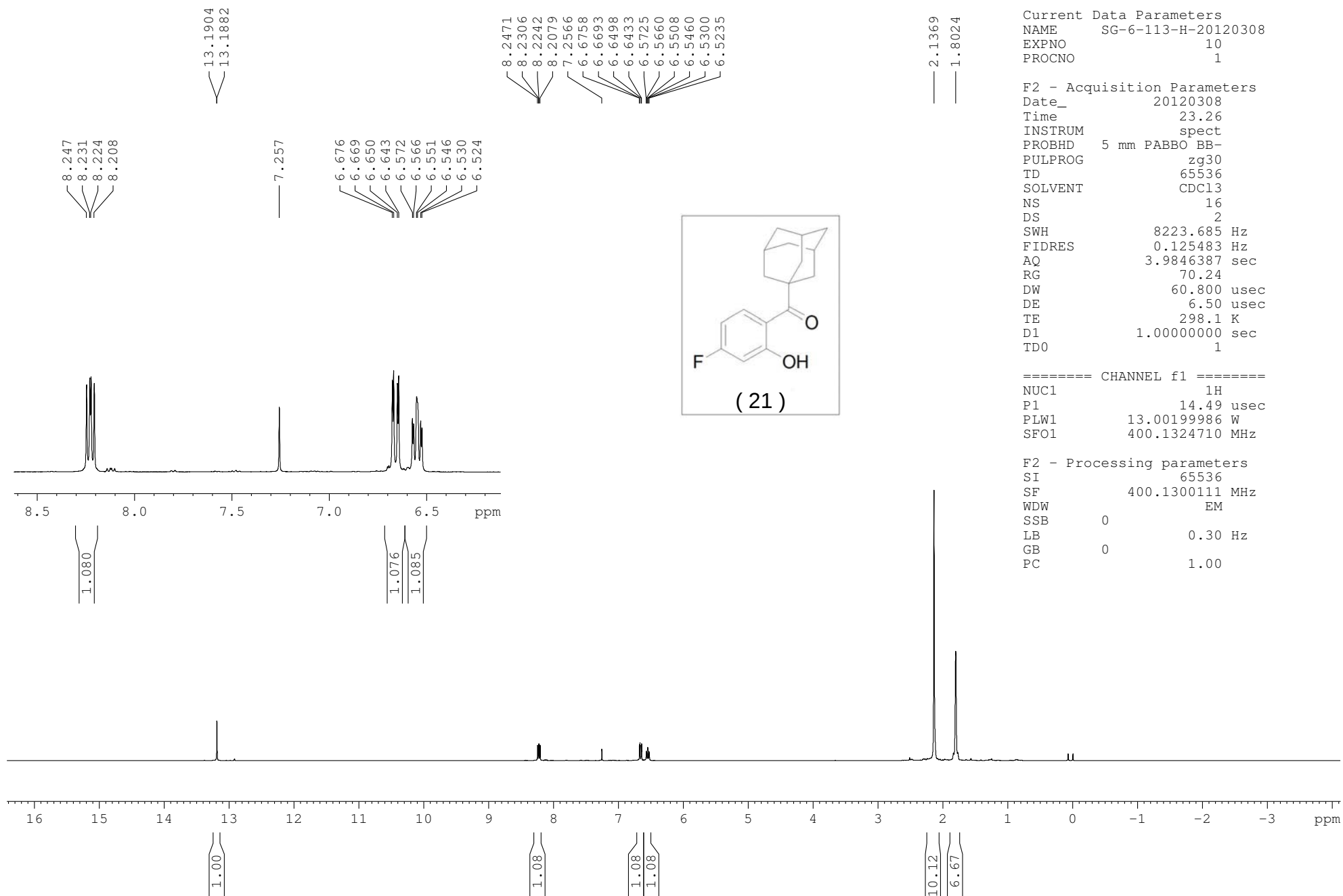
Current Data Parameters
NAME SG-6-109-C-20120404
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120404
Time 22.12
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 184.16
DW 20.800 usec
DE 6.50 usec
TE 298.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.31 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35685000 W
PLW13 0.28904000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127565 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

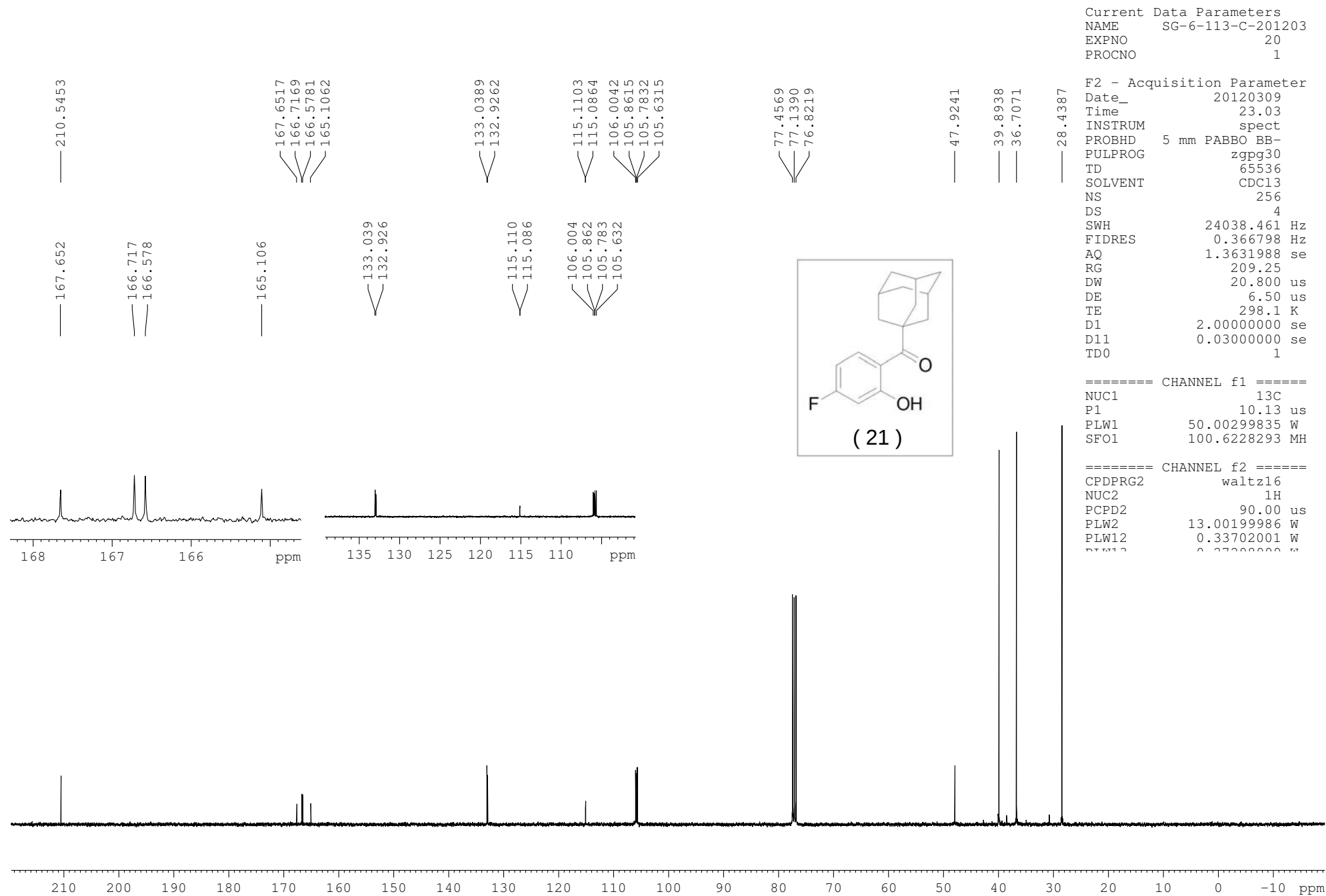


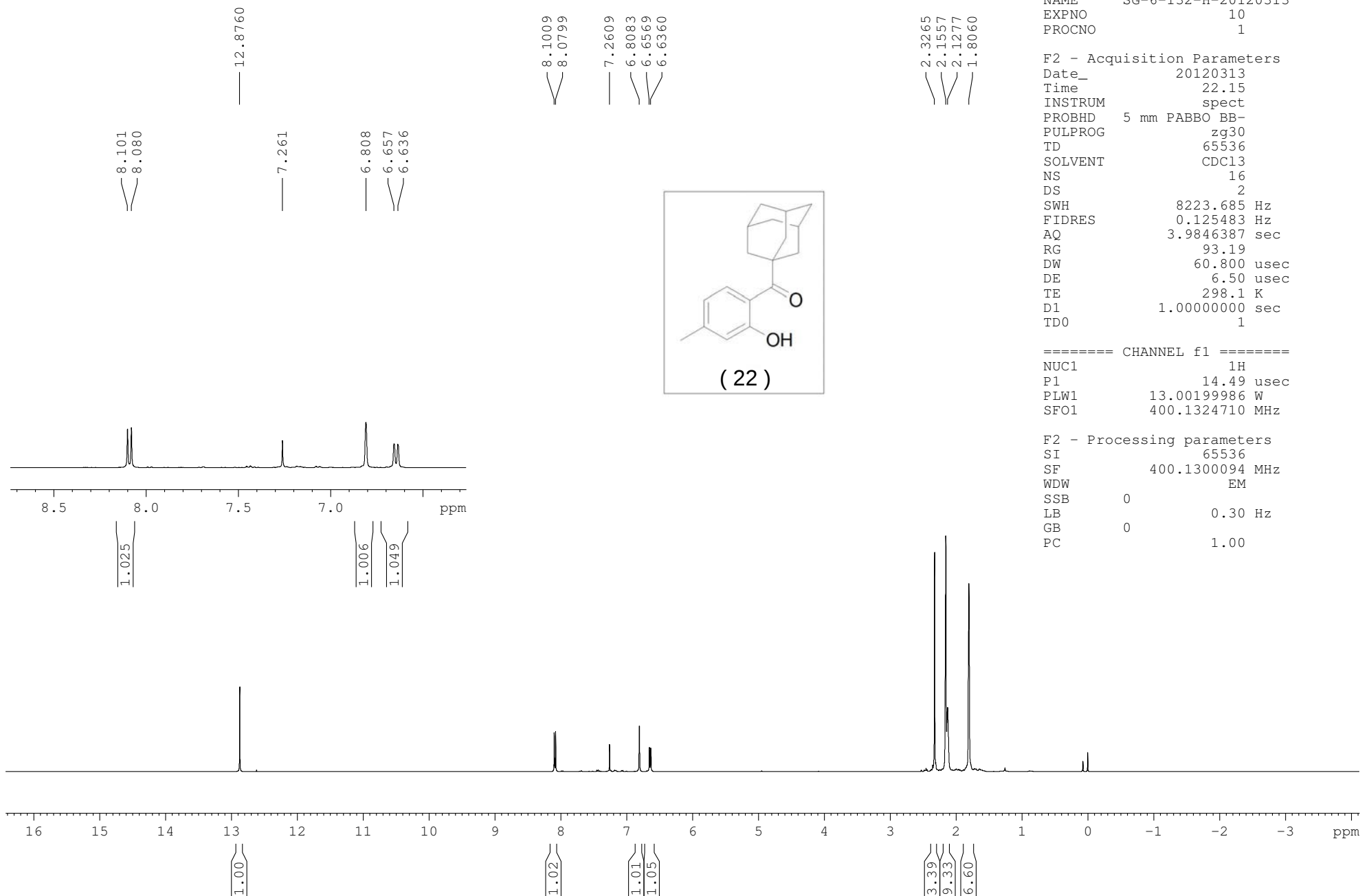
Current Data Parameters
NAME SG-6-113-H-20120308
EXPNO 10
PROCNO 1

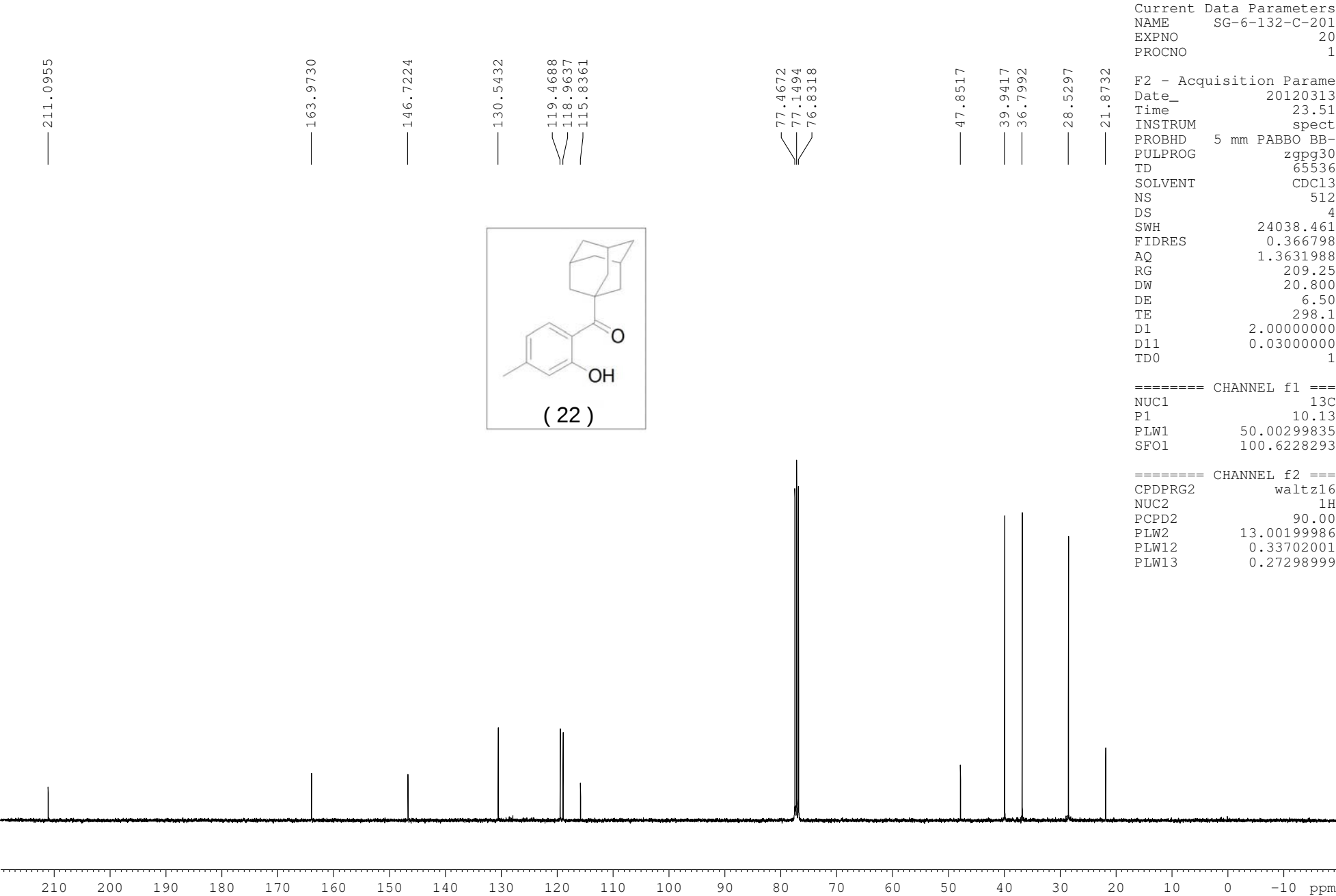
F2 - Acquisition Parameters
Date_ 20120308
Time 23.26
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 70.24
DW 60.800 usec
DE 6.50 usec
TE 298.1 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.49 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300111 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00





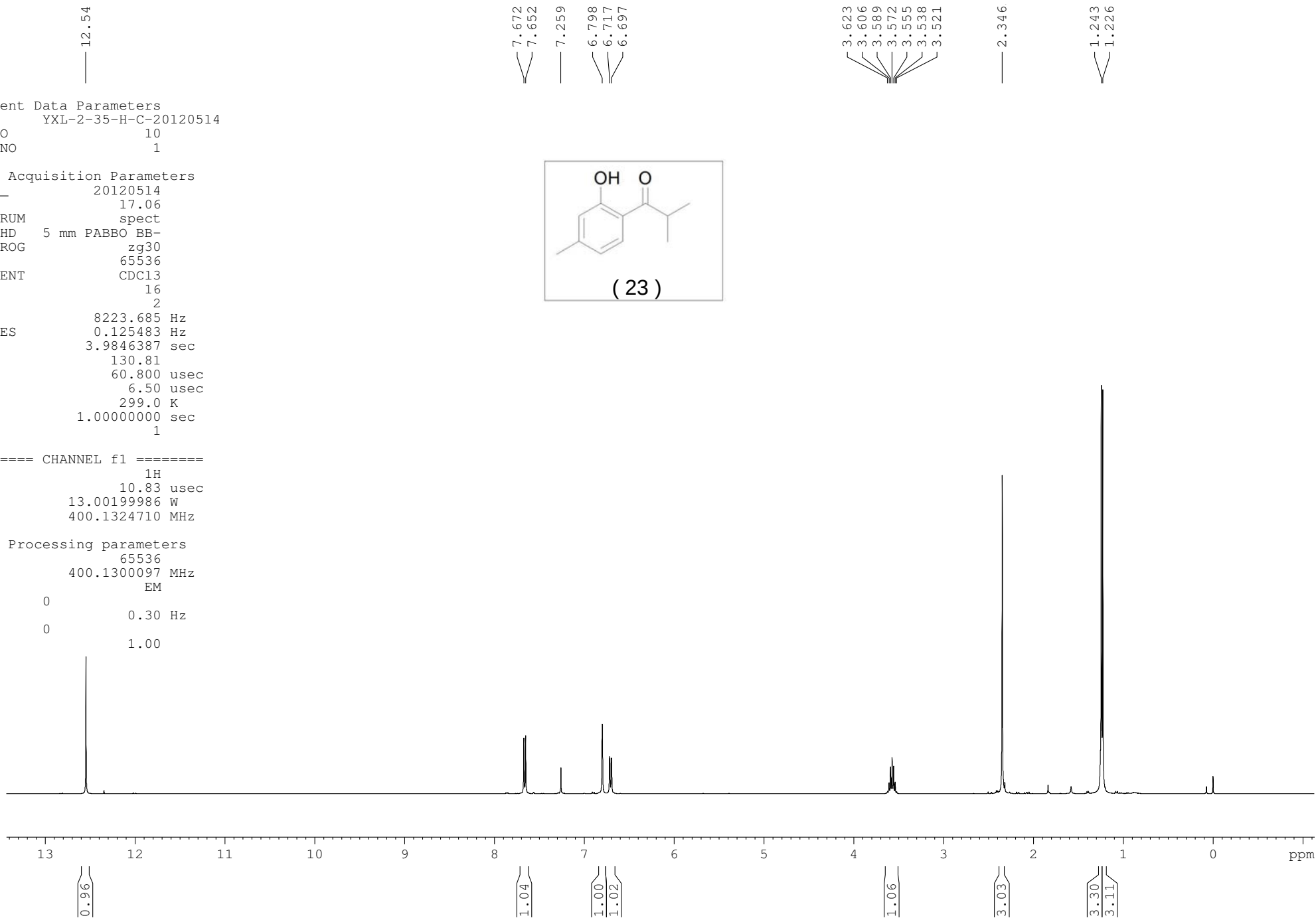
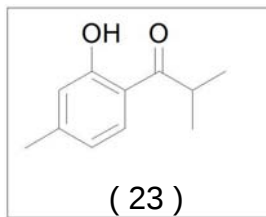


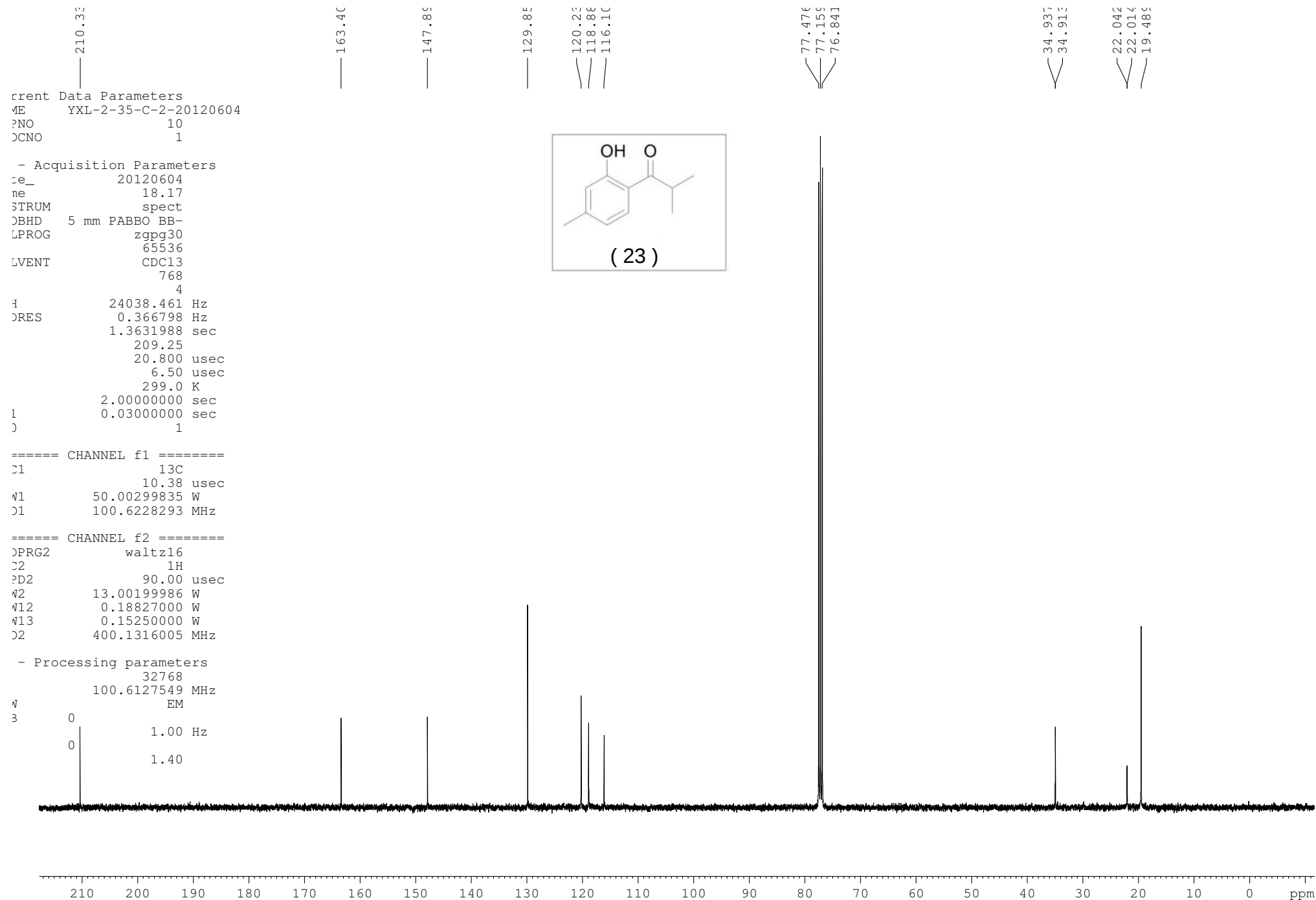
Current Data Parameters
NAME YXL-2-35-H-C-20120514
CPRO 10
PROCNO 1

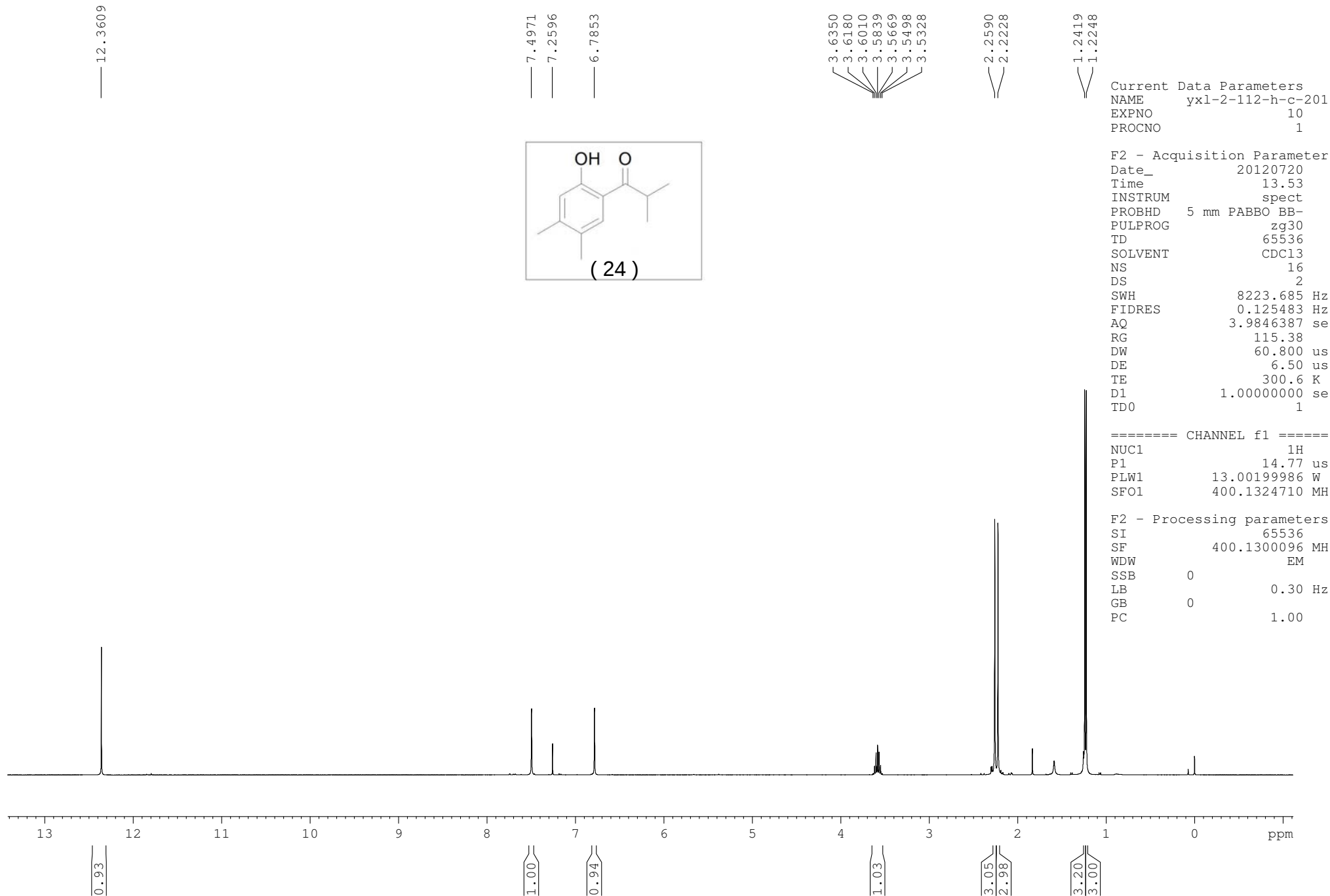
2 - Acquisition Parameters
Date_ 20120514
Time 17.06
INSTRUM spect
PROBHD 5 mm PABBO BB-
JLPROG zg30
NUC1 65536
PULPROG zgpg30
PCVENT CDC13
SOLVENT 16
SOLVENT 2
FID 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
SFO 130.81
V 60.800 usec
E 6.50 usec
T 299.0 K
L 1.00000000 sec
DO 1

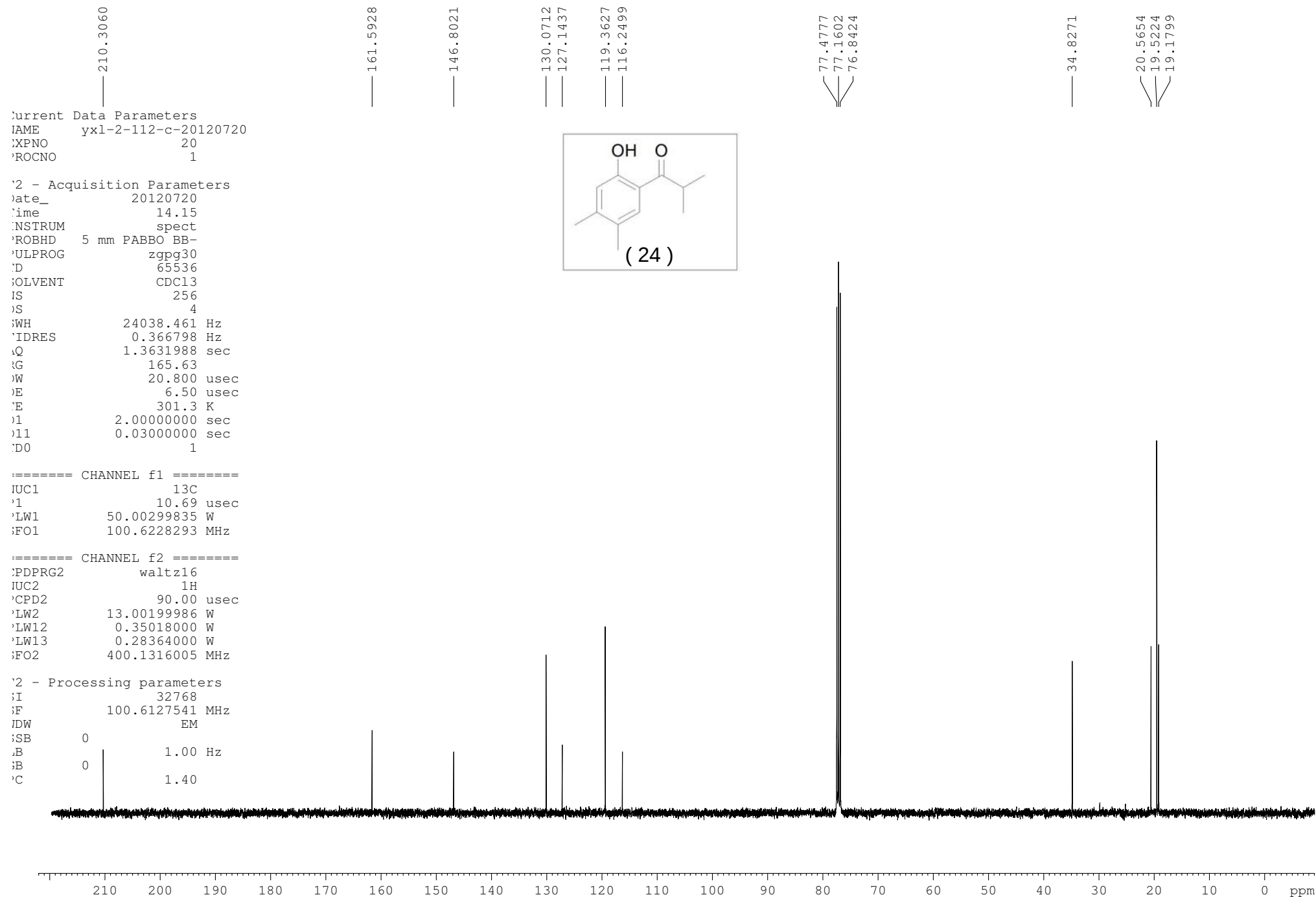
===== CHANNEL f1 =====
JC1 1H
L 10.83 usec
P1 13.00199986 W
PC1 400.1324710 MHz

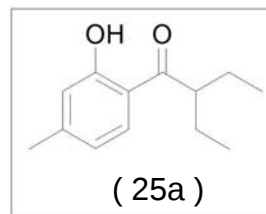
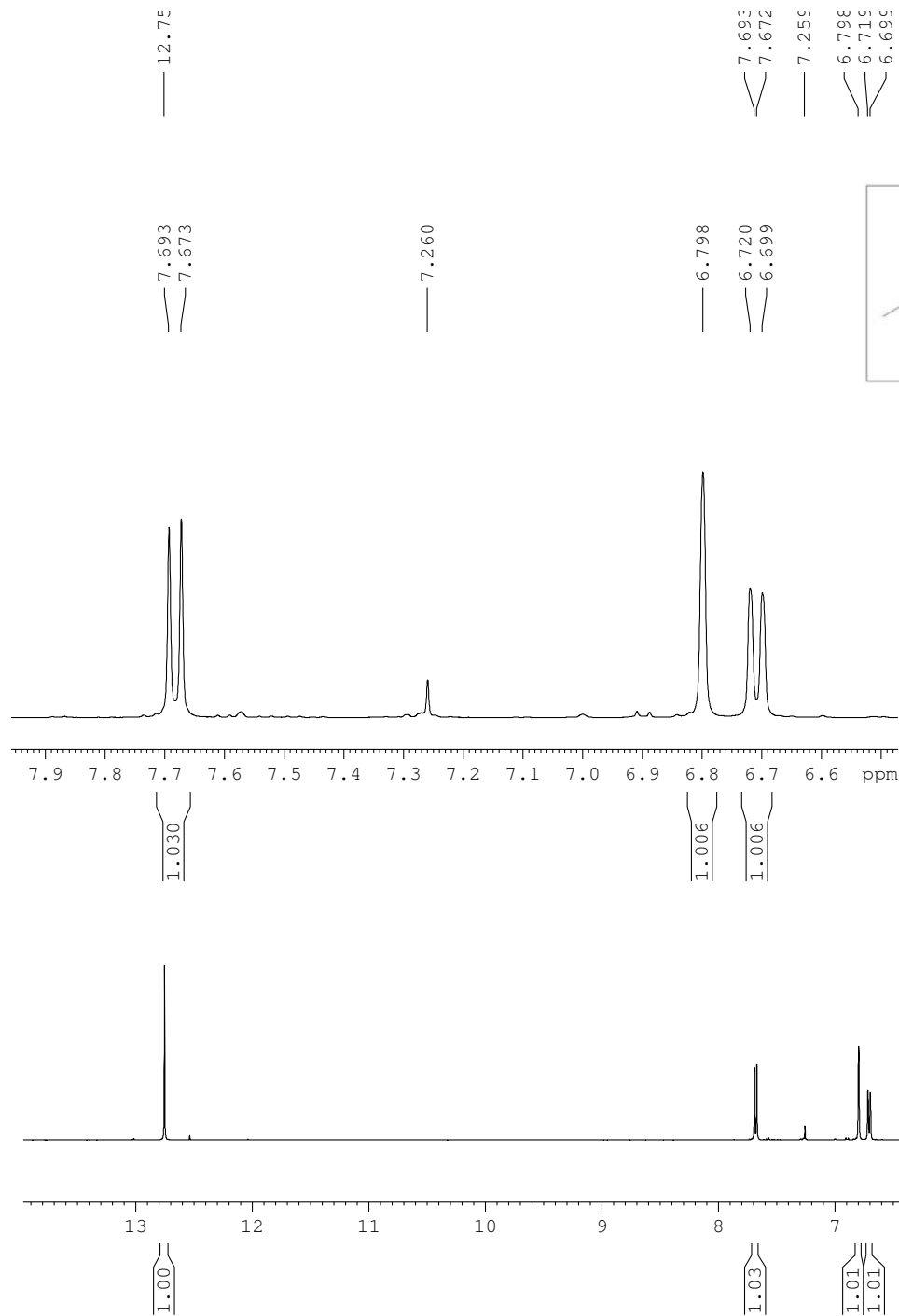
2 - Processing parameters
F 65536
P 400.1300097 MHz
W EM
S 0
B 0.30 Hz
C 0
G 1.00









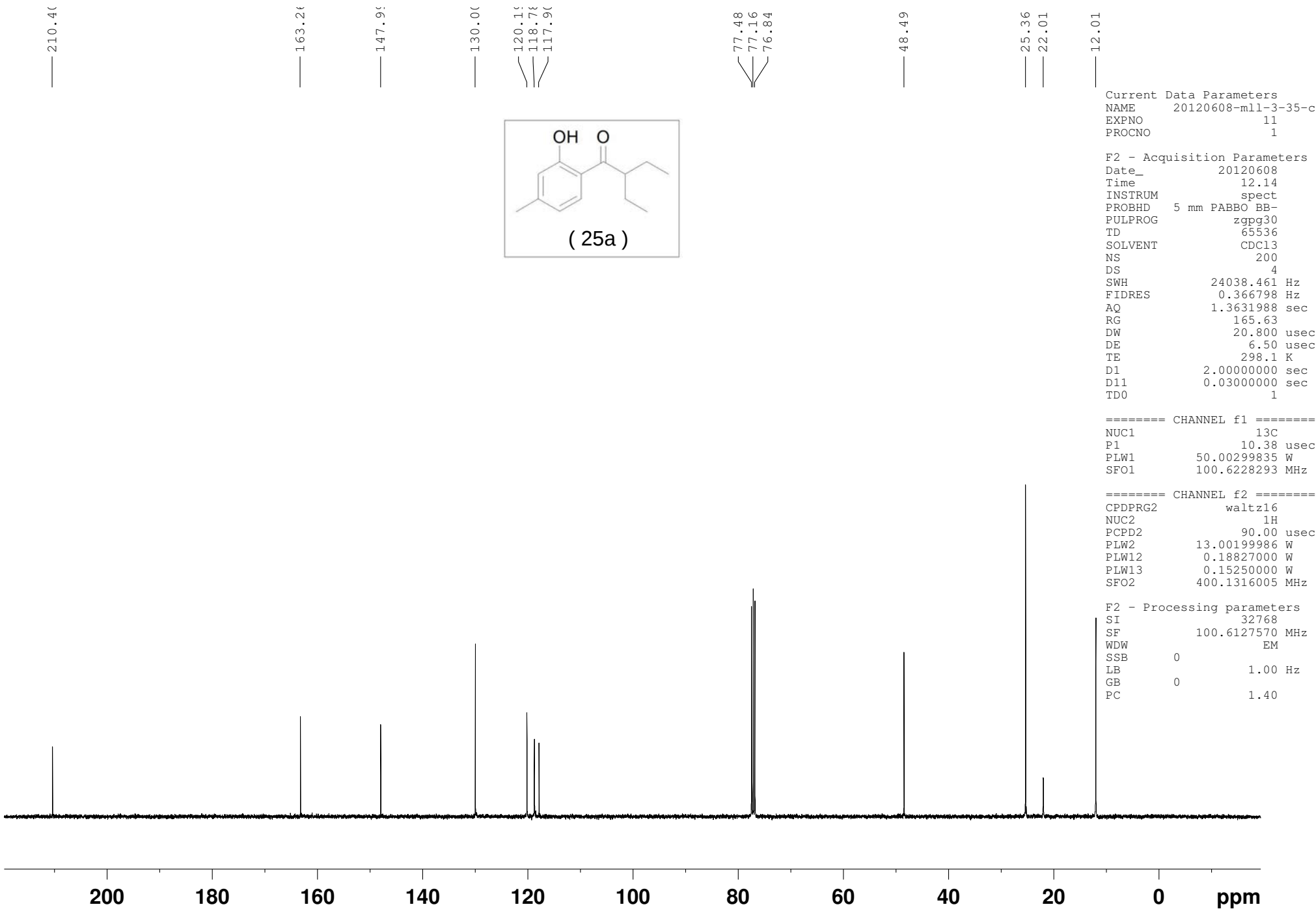


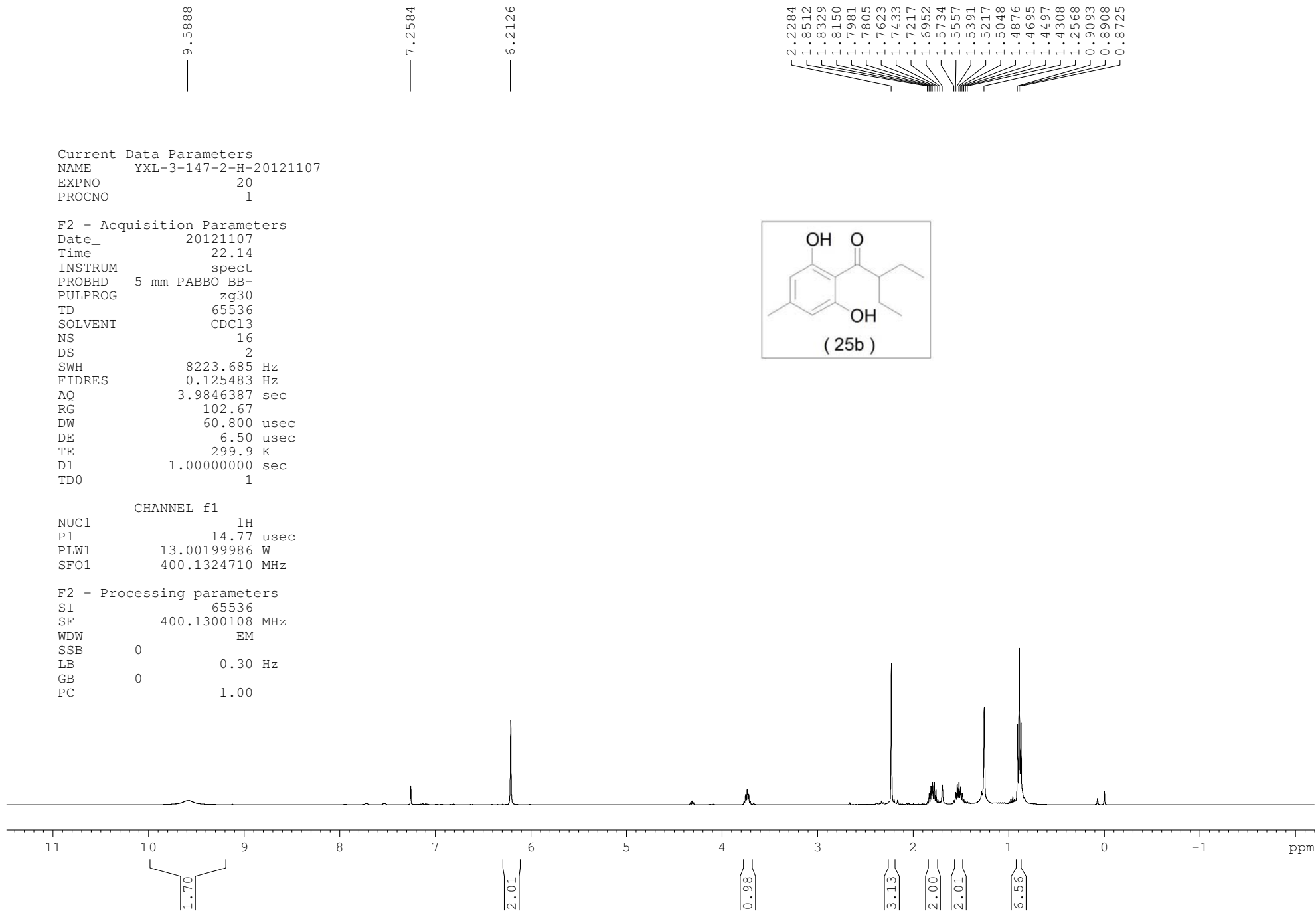
Current Data Parameters
NAME 20120608-ml1-3-35
EXPNO 10
PROCNO 1

F2 - Acquisition Parameter
Date_ 20120608
Time 10.52
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 se
RG 56.81
DW 60.800 us
DE 6.50 us
TE 298.1 K
D1 1.00000000 se
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 10.83 us
PLW1 13.00199986 W
SFO1 400.1324710 MH

F2 - Processing parameters
SI 65536
SF 400.1300094 MH
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00





C13CPD CDC13 {D:\NMR_DATA} RY 32

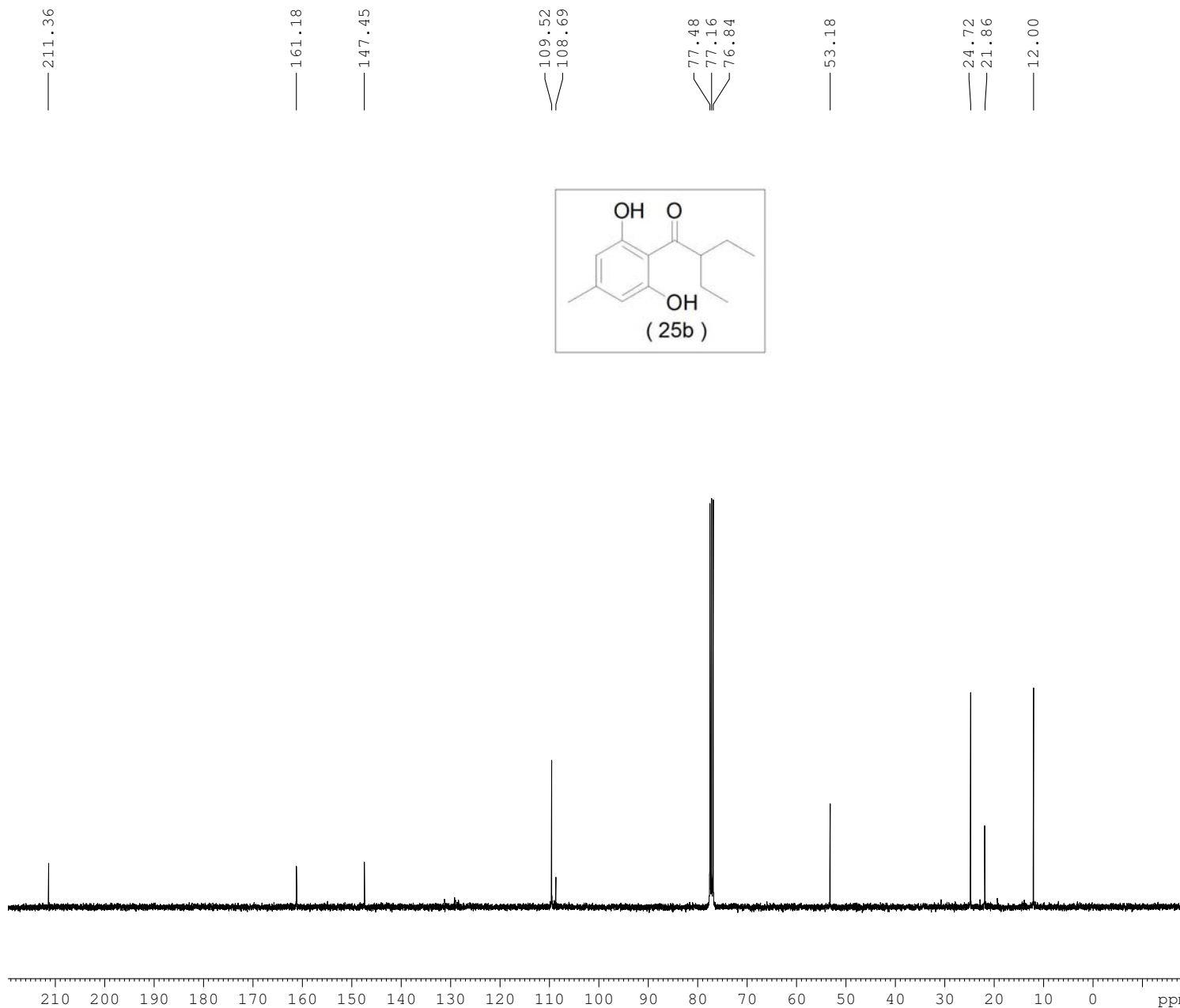
Current Data Parameters
NAME 20121116-YSY-1---157
EXPNO 11
PROCNO 1

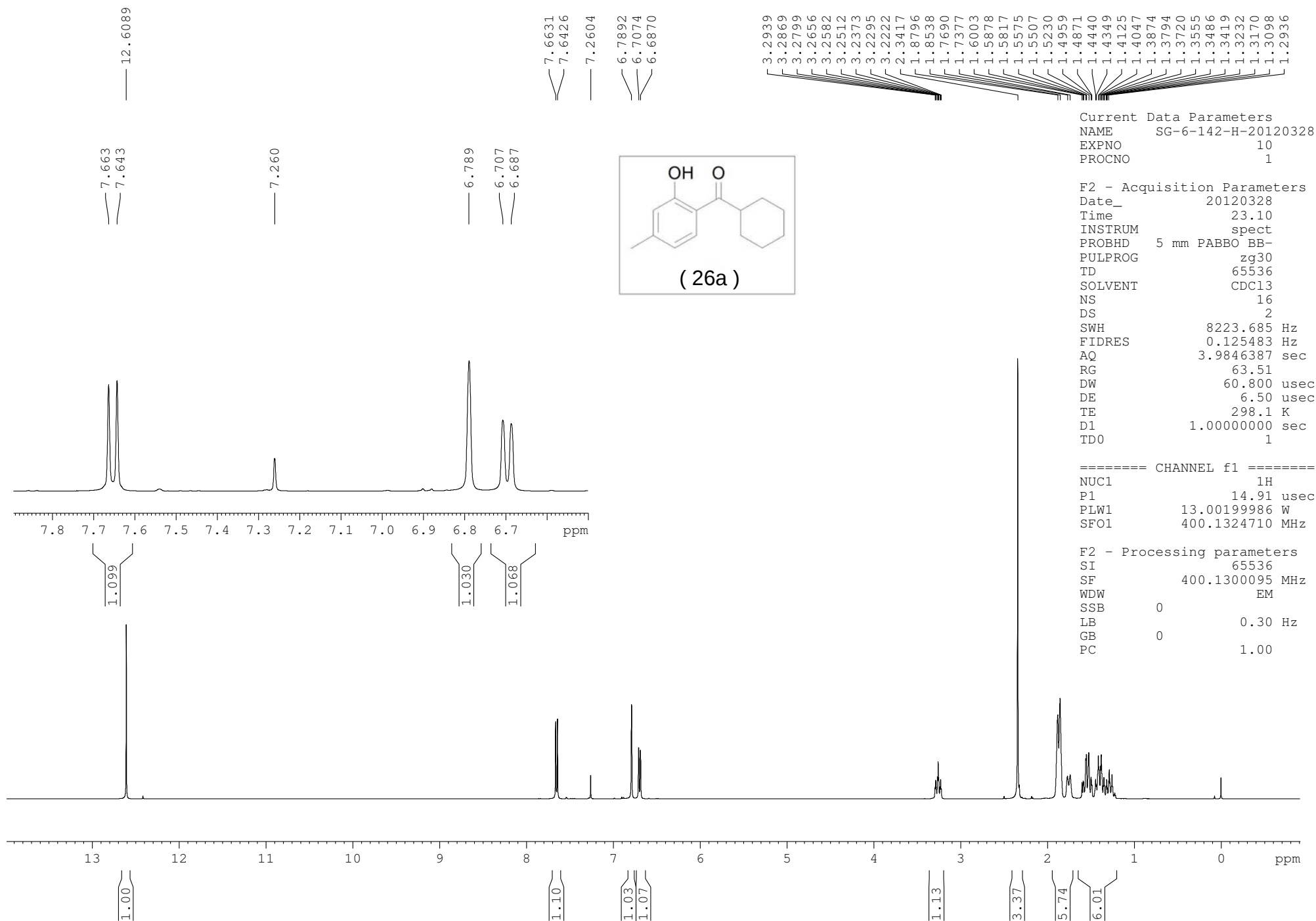
F2 - Acquisition Parameters
Date_ 20121116
Time 13.55
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 453
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

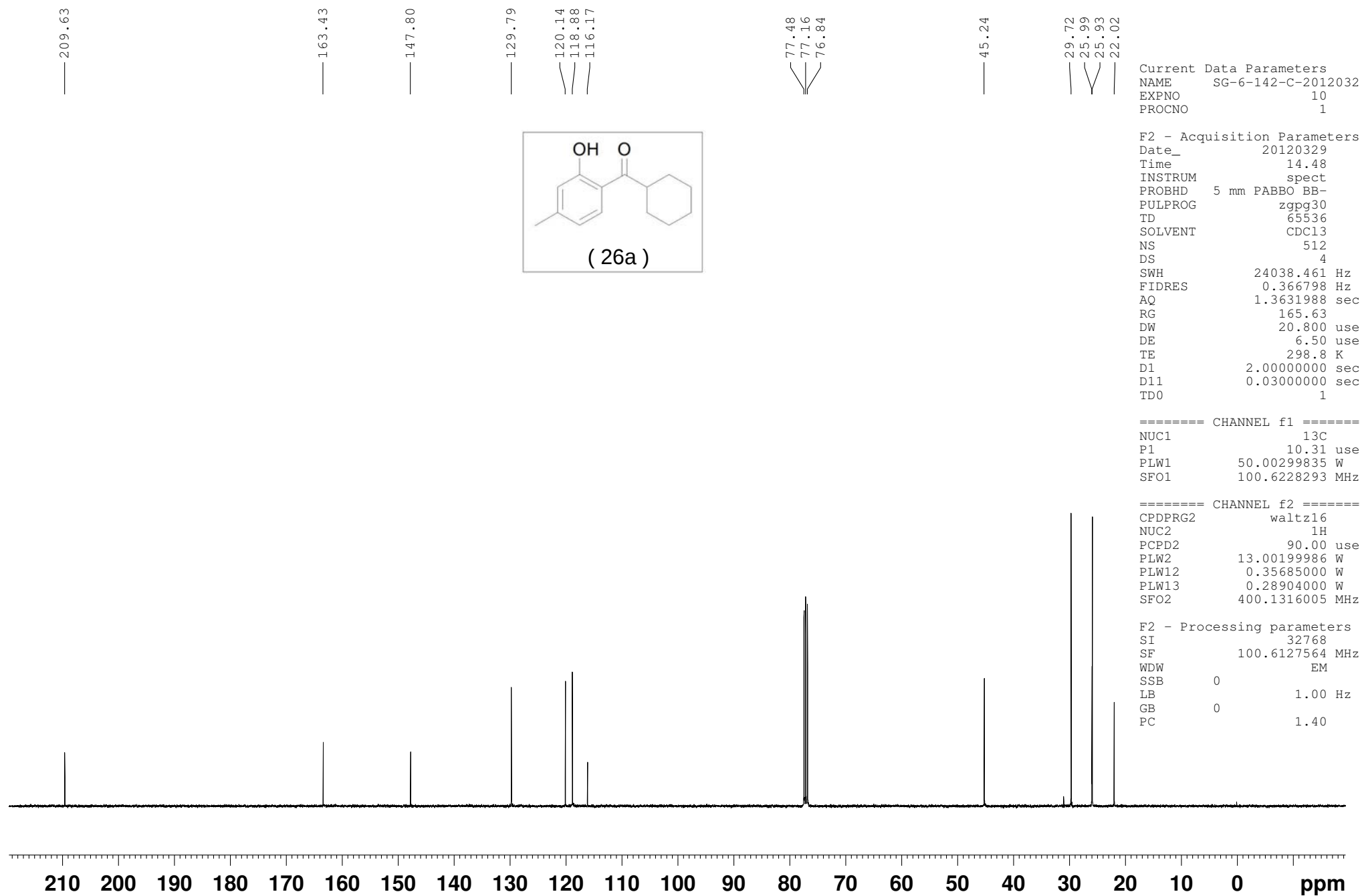
===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

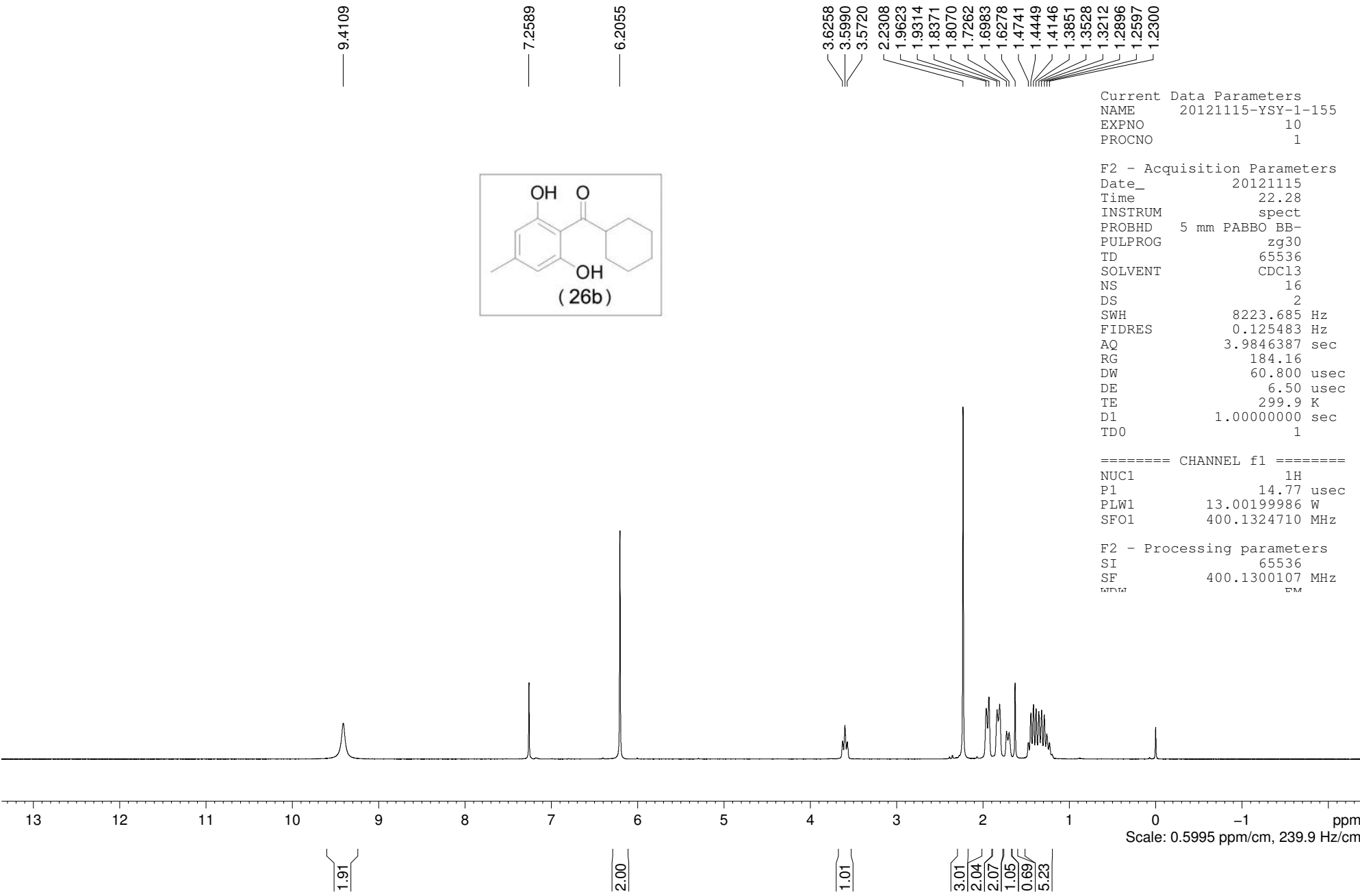
===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127548 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

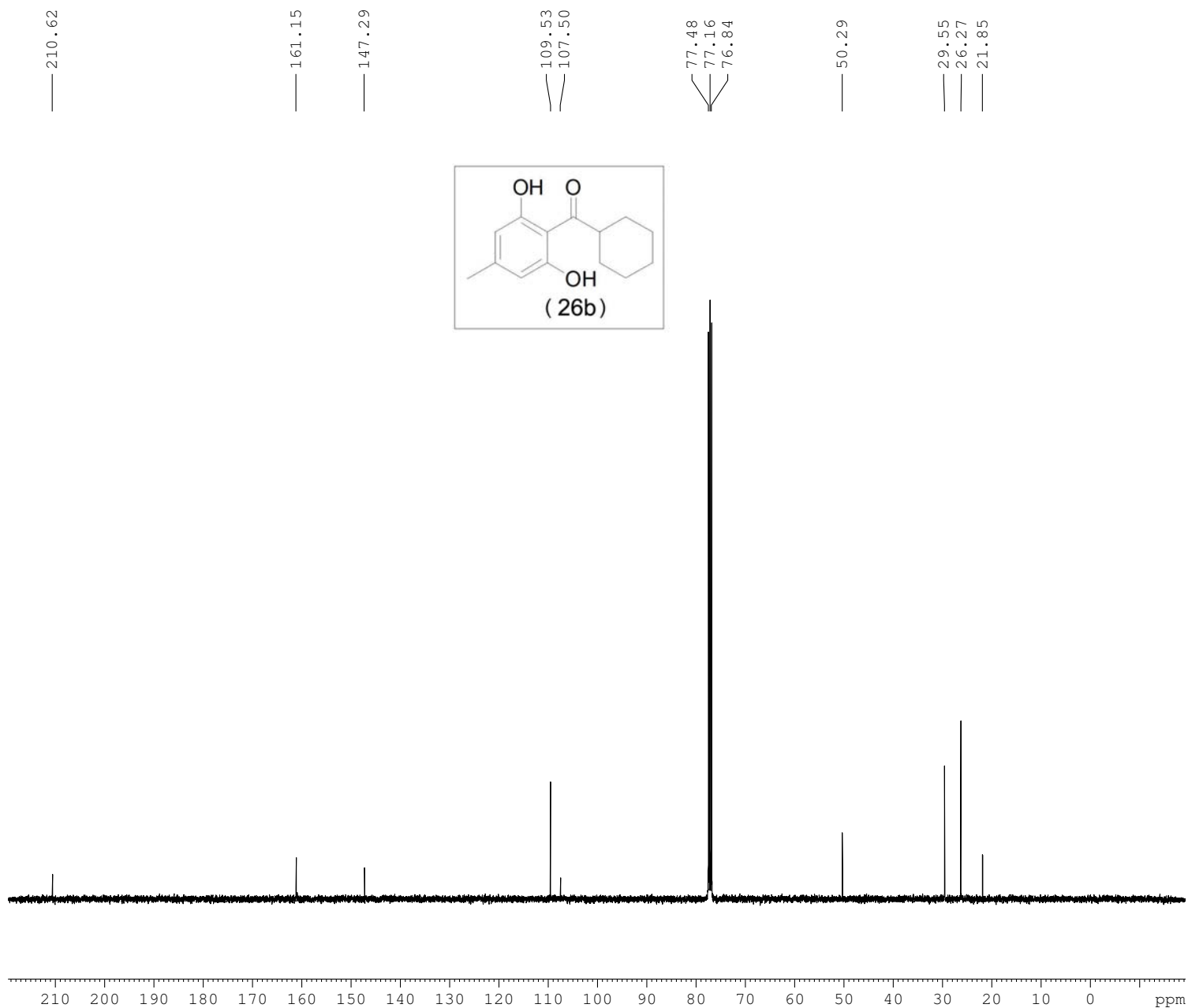








C13CPD CDC13 {D:\NMR_DATA} RY 19



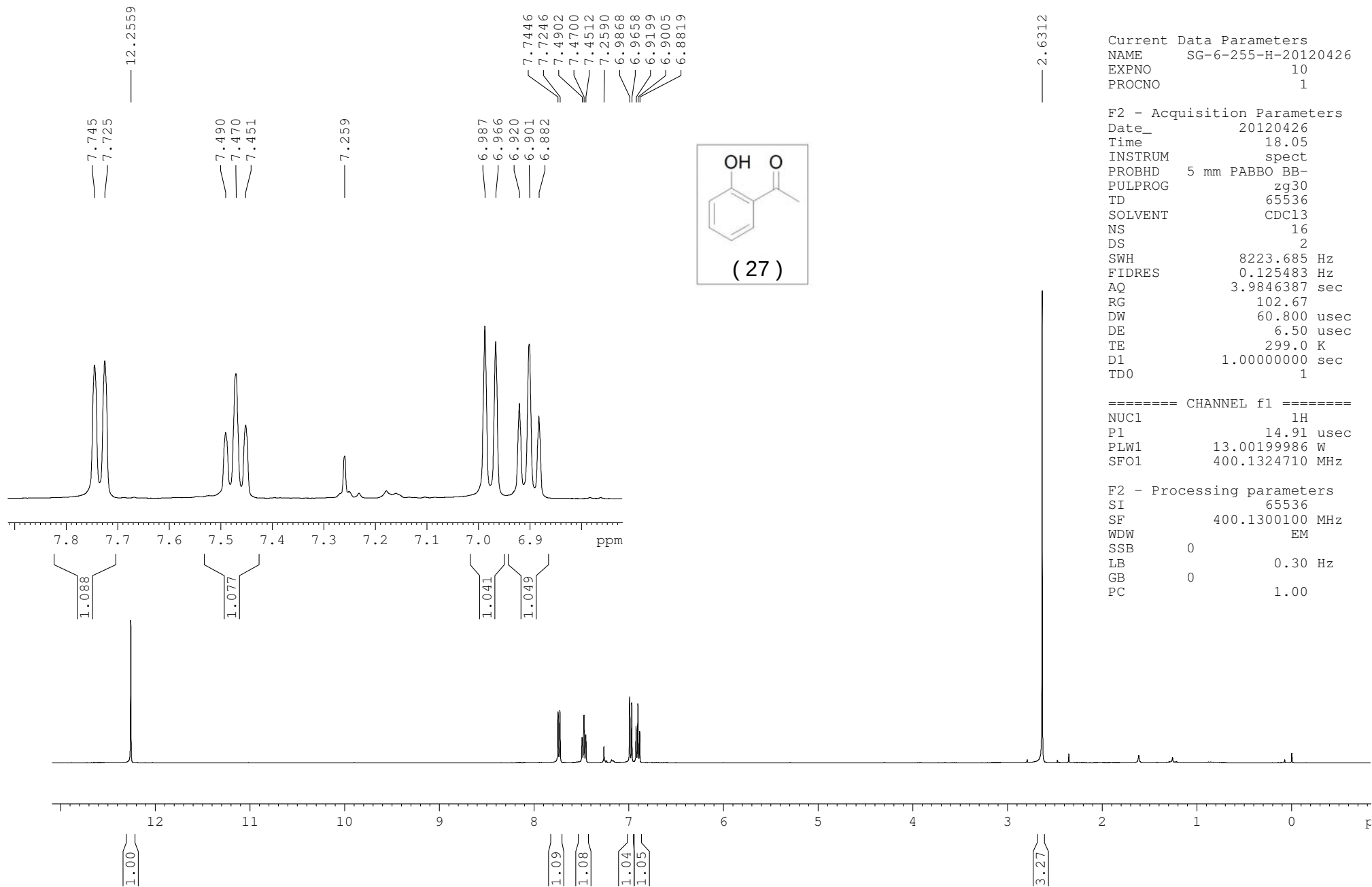
Current Data Parameters
NAME 20121115-YSY-1-155
EXPNO 11
PROCNO 1

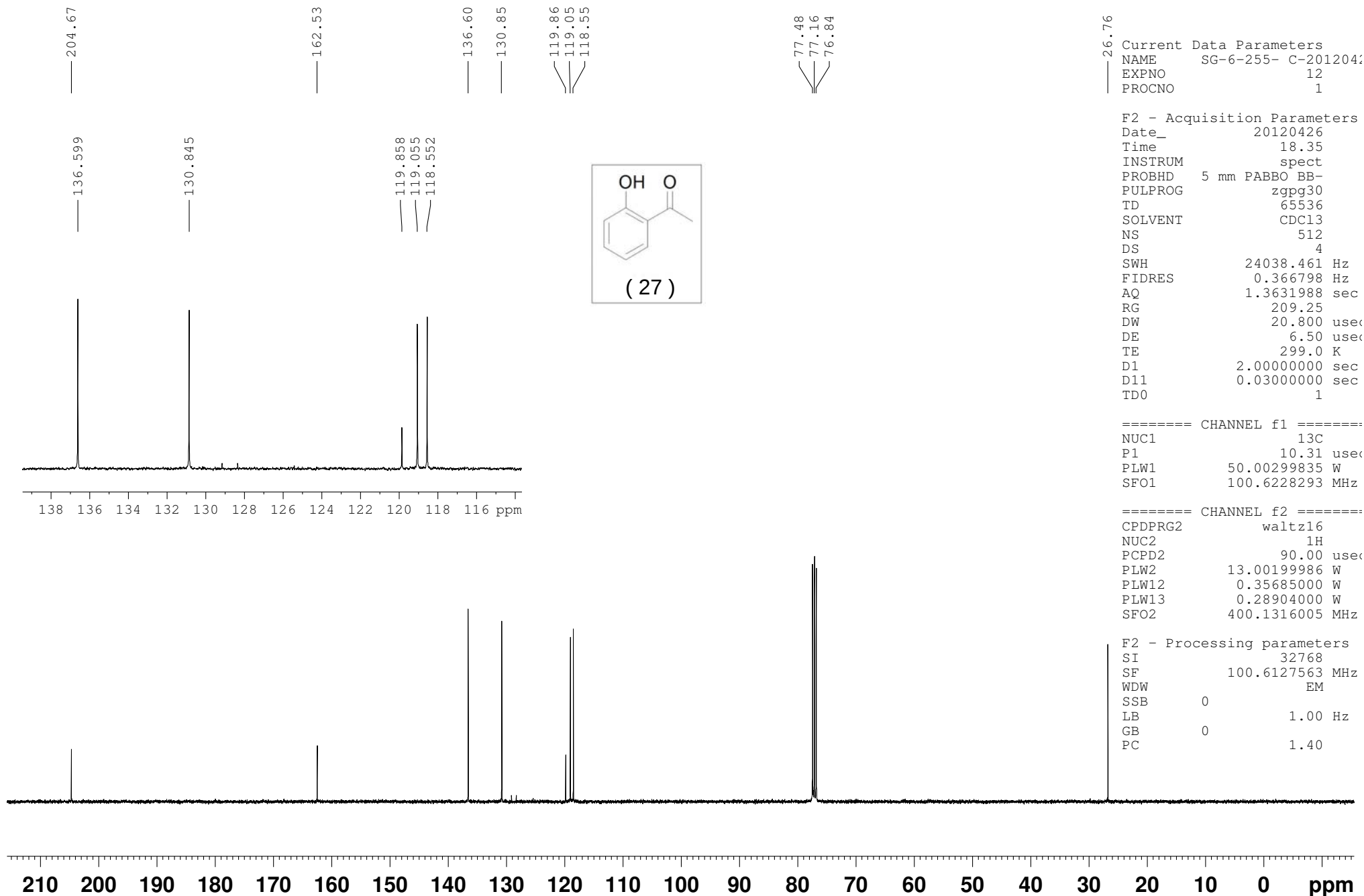
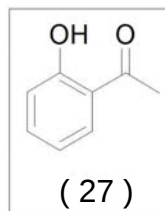
F2 - Acquisition Parameters
Date_ 20121116
Time 7.05
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 184.16
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

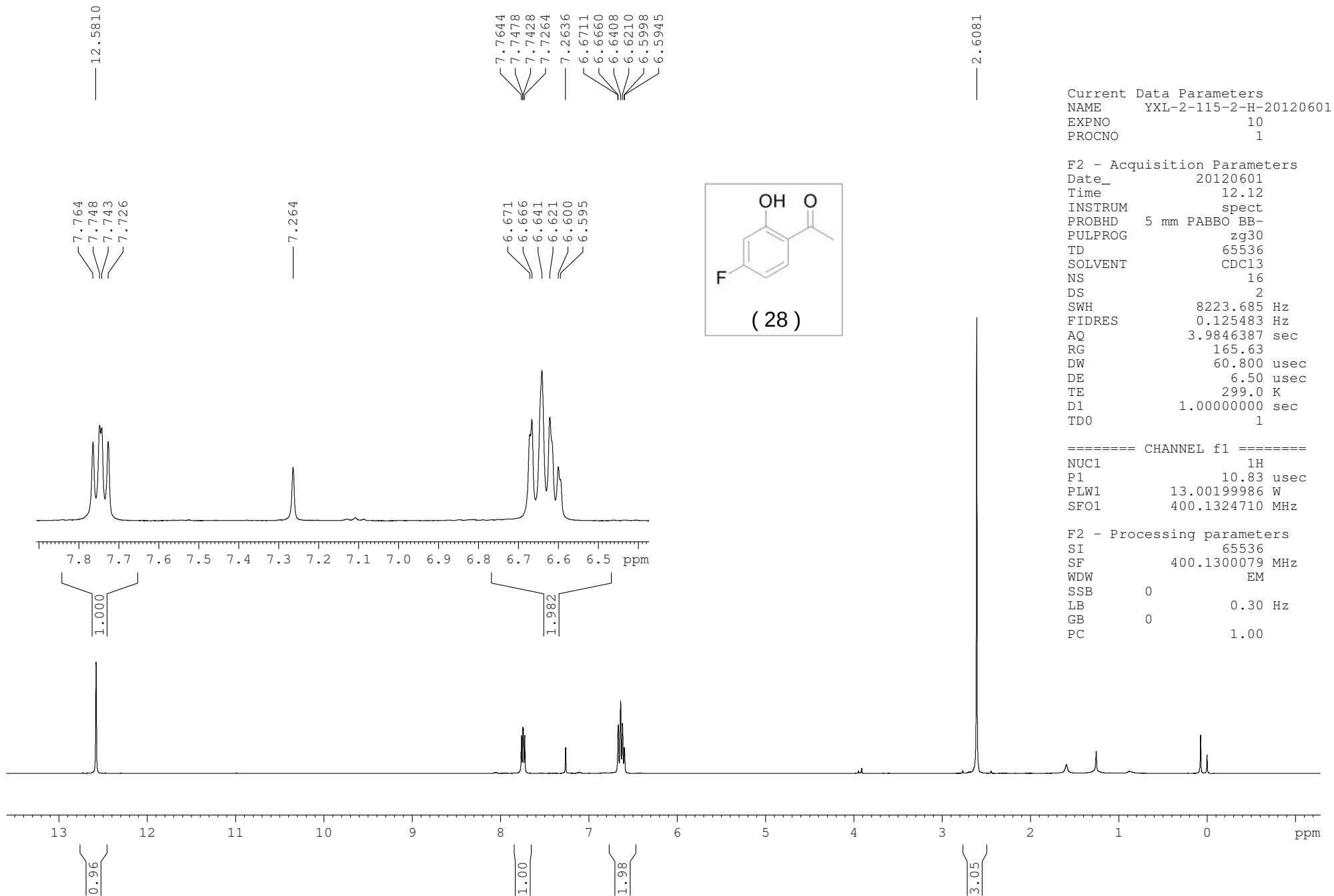
===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

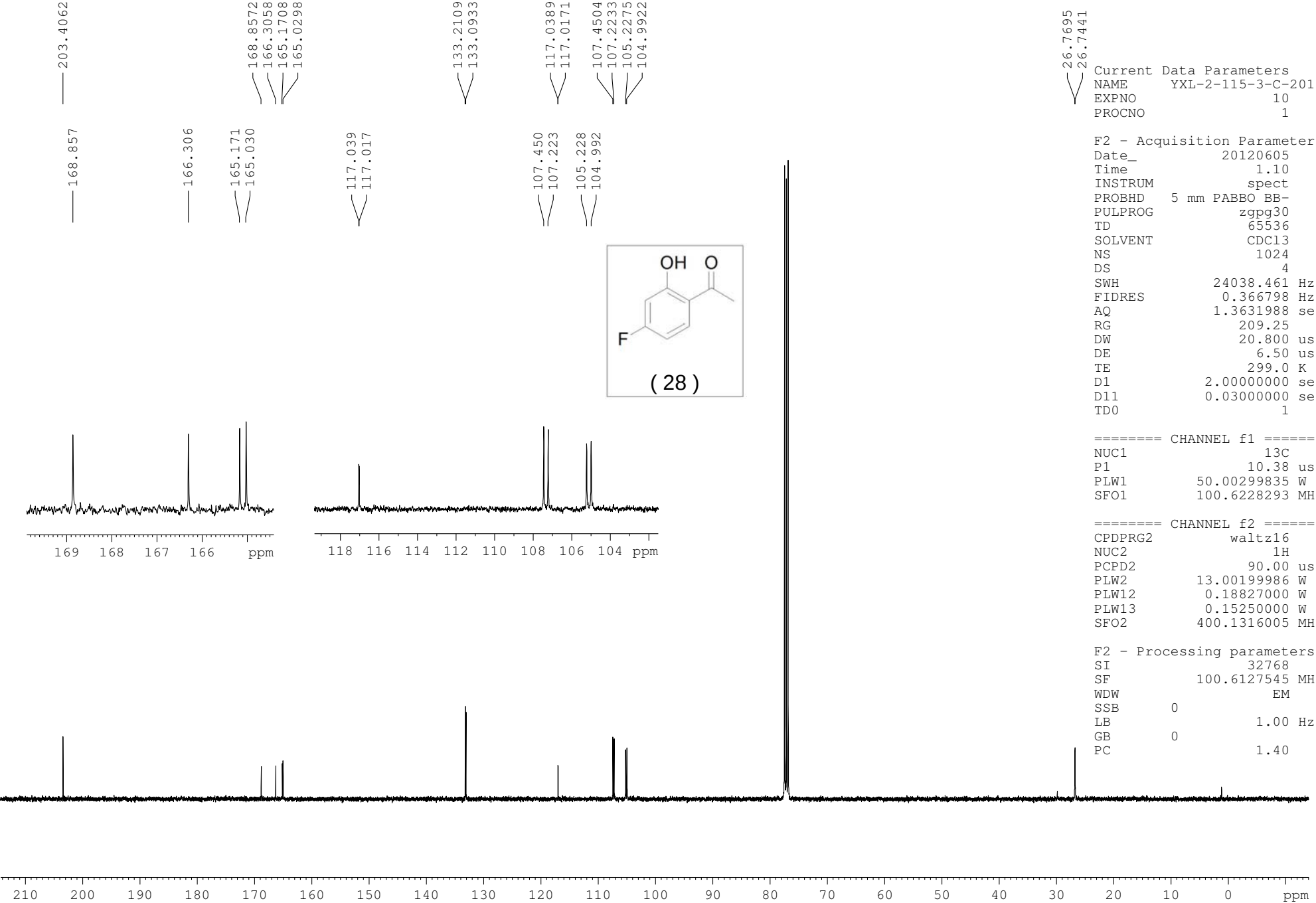
===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

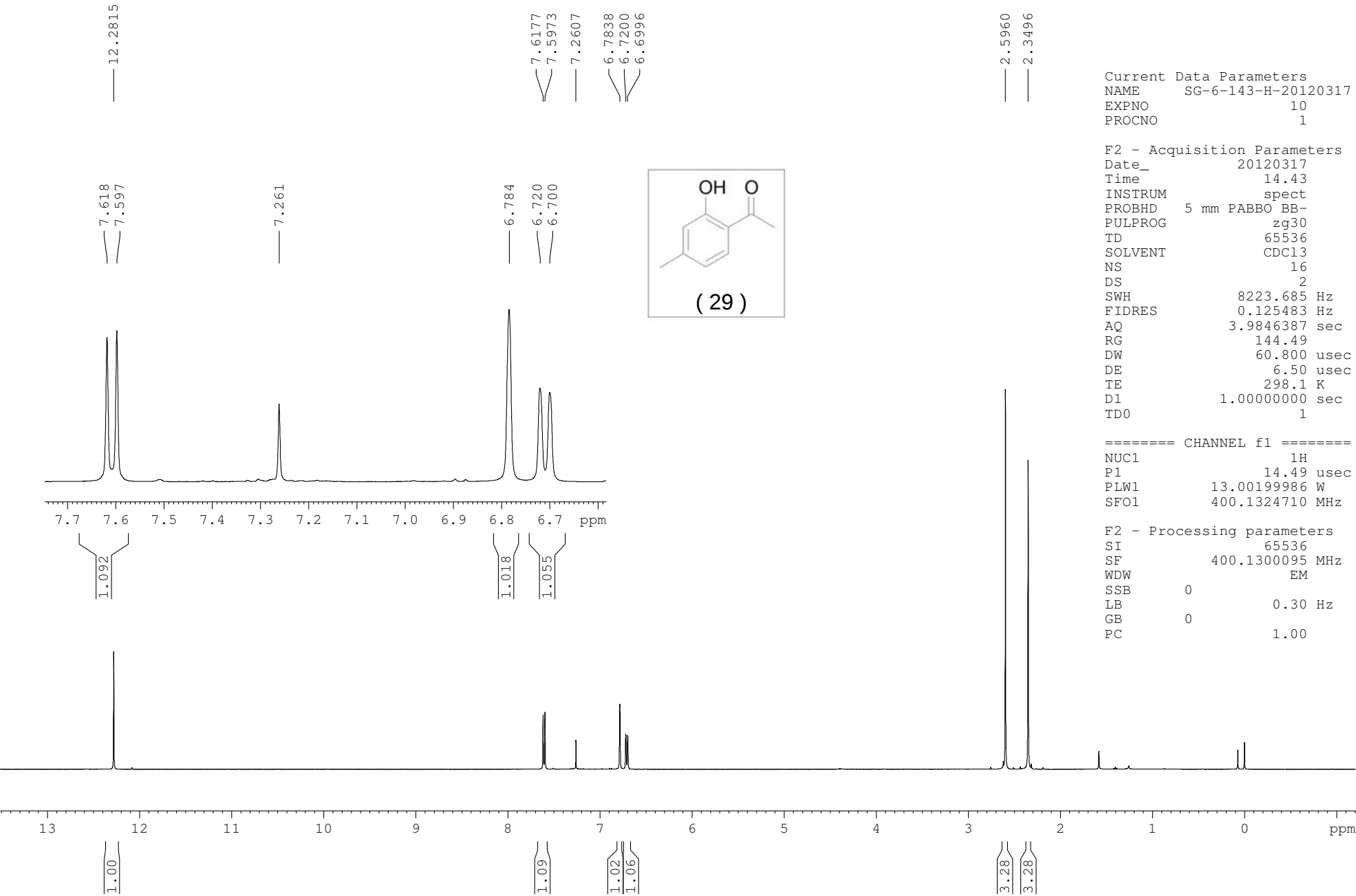
F2 - Processing parameters
SI 32768
SF 100.6127541 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

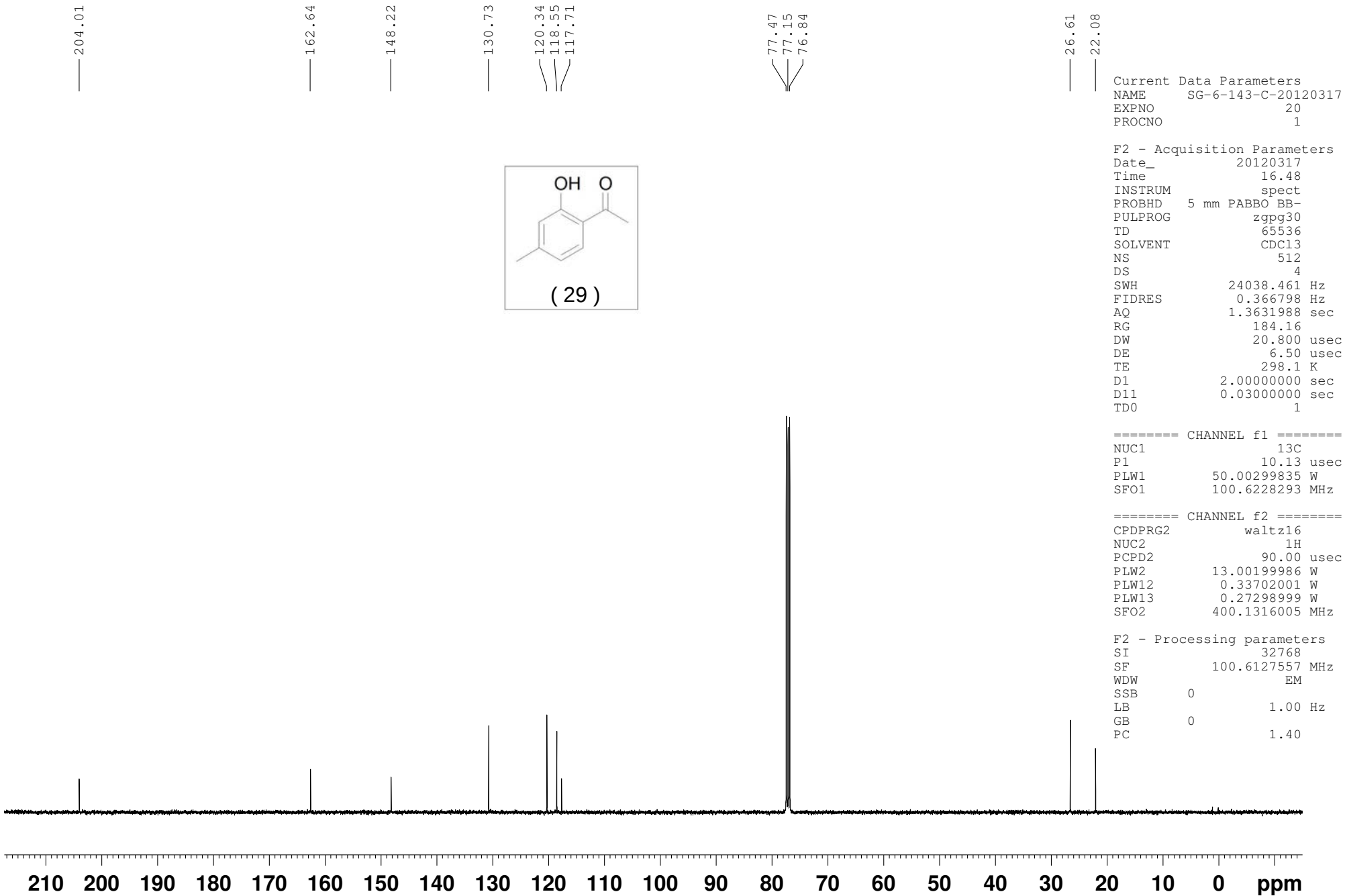


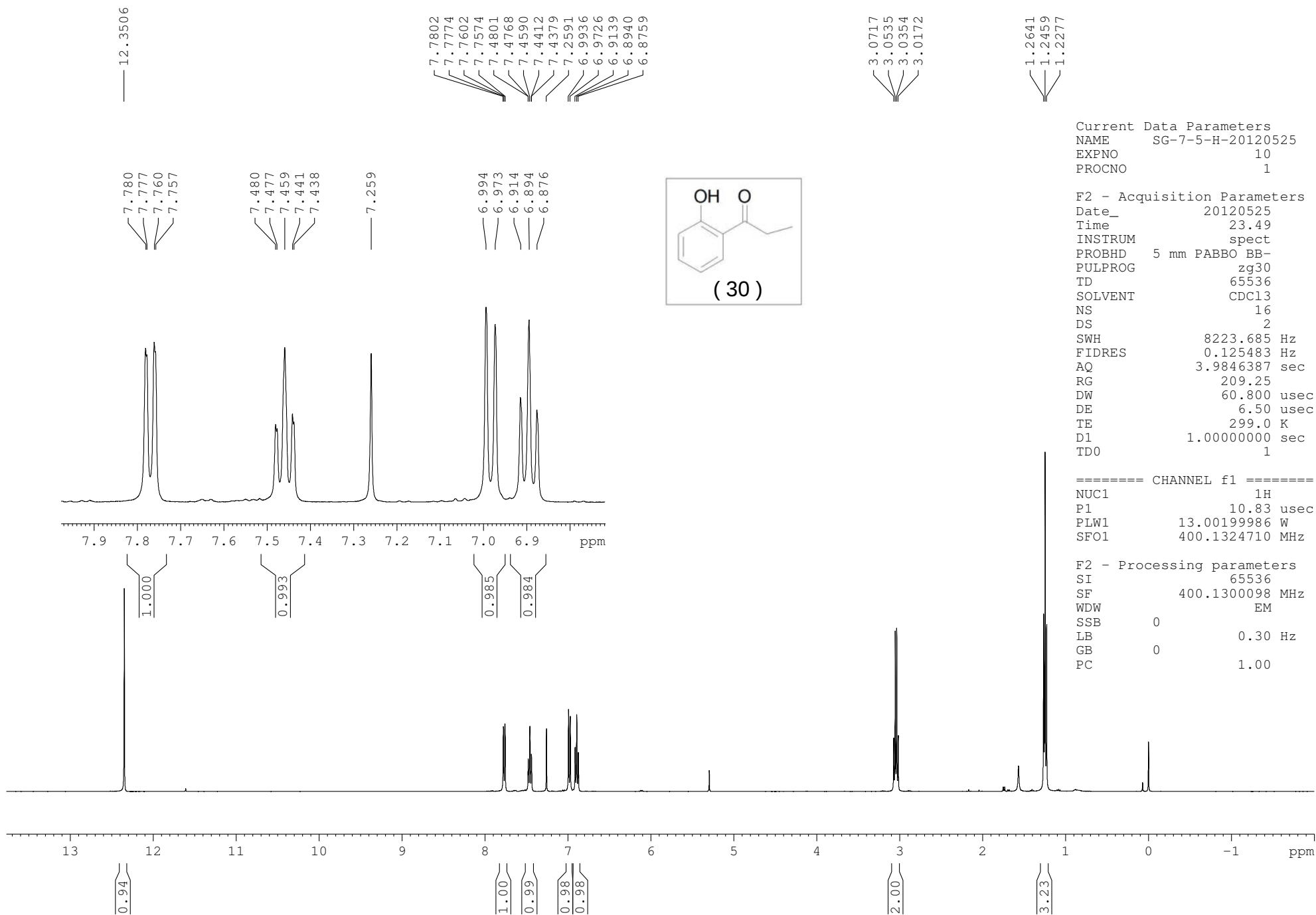


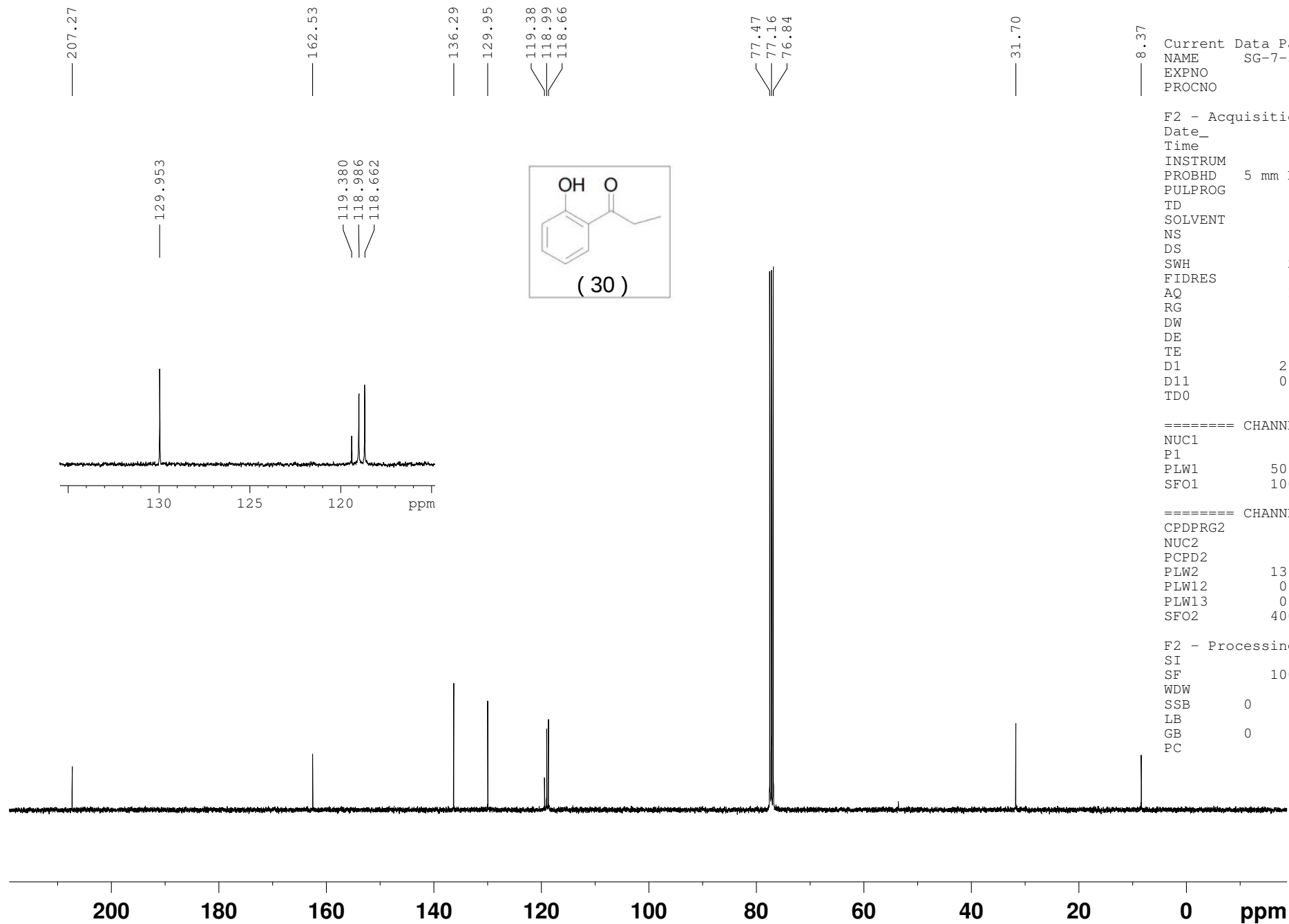












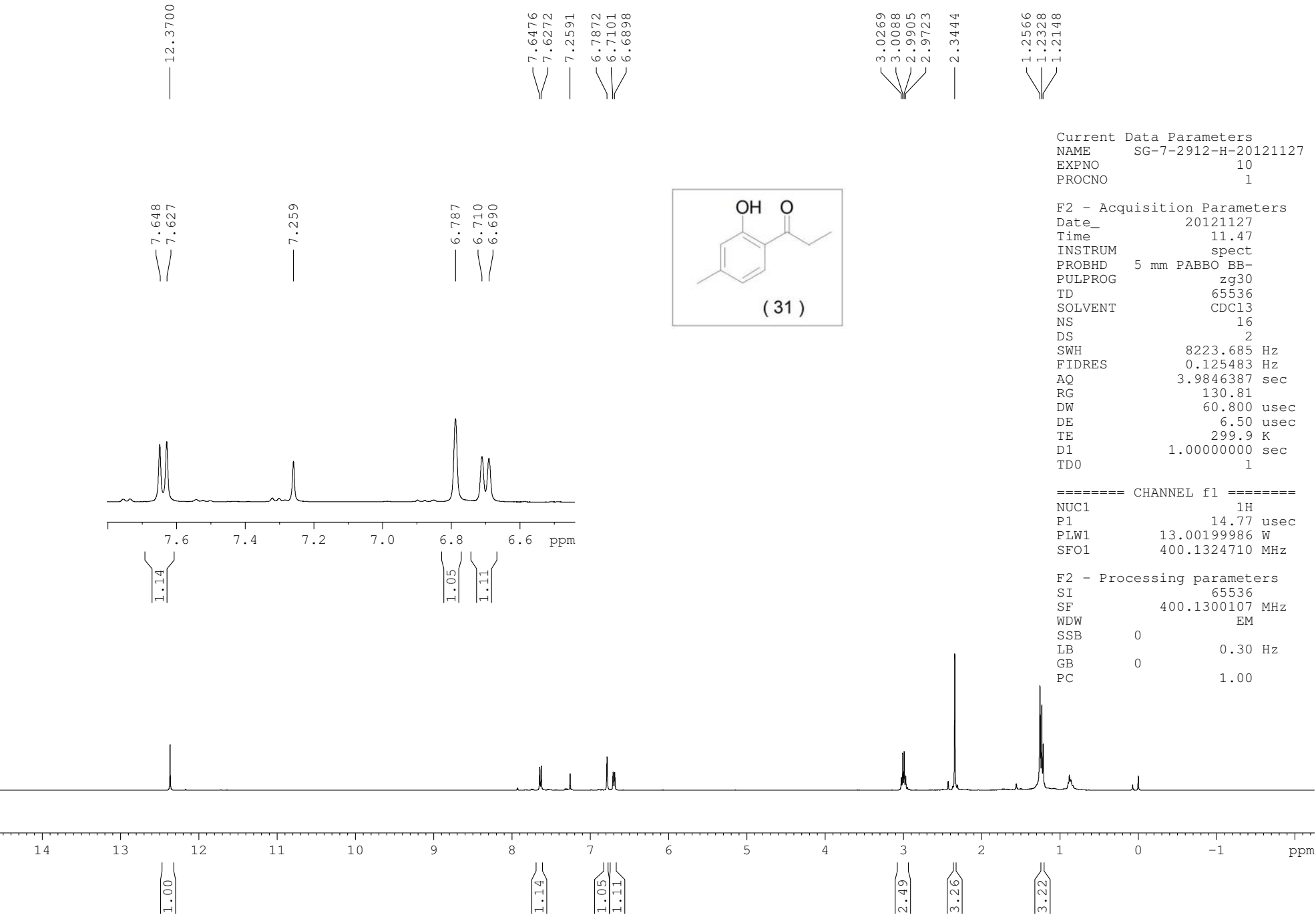
Current Data Parameters
NAME SG-7-5-C-20120526
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120526
Time 13.00
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl₃
NS 750
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 184.16
DW 20.800 usec
DE 6.50 usec
TE 299.0 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

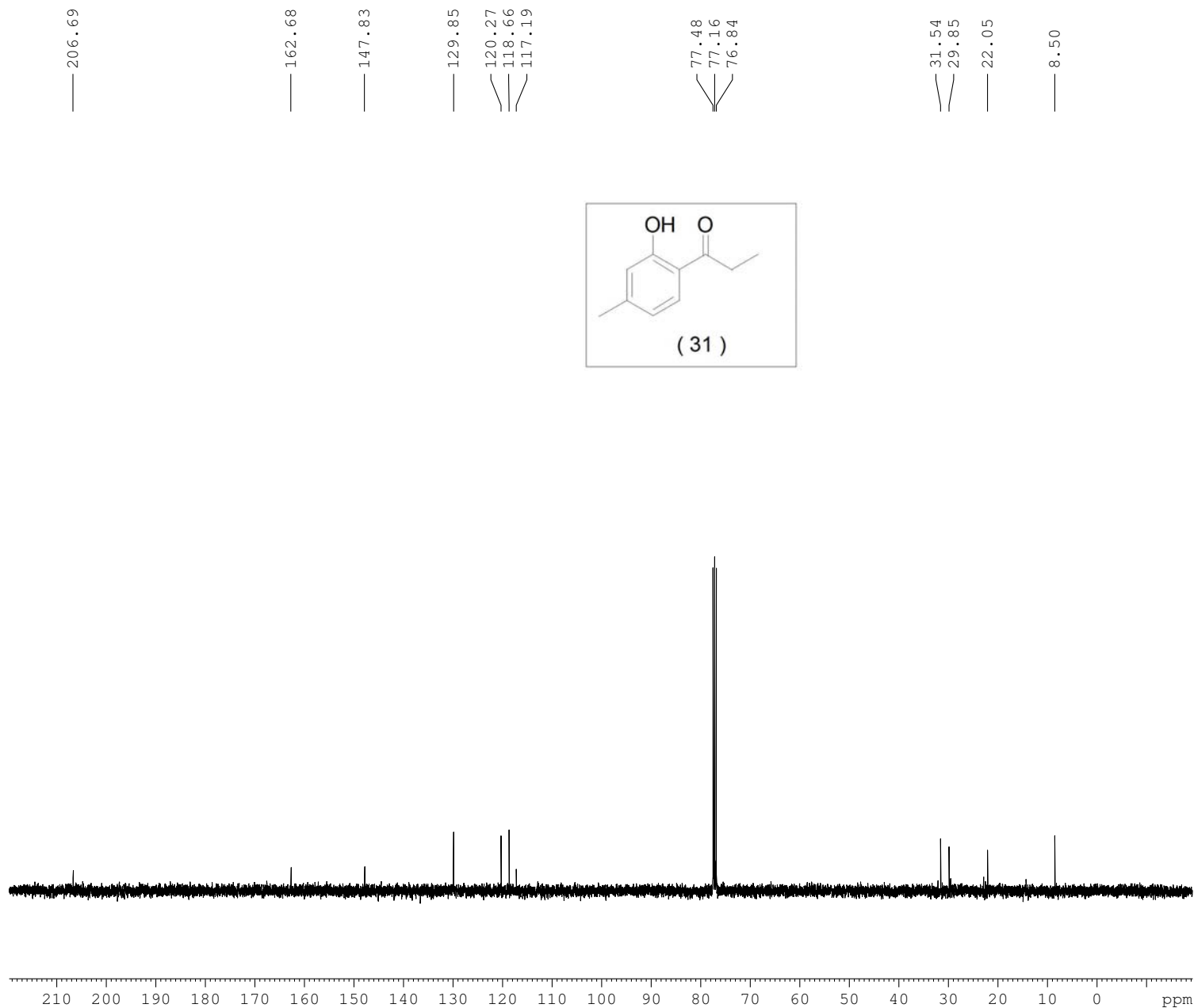
===== CHANNEL f1 =====
NUC1 13C
P1 10.38 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.18827000 W
PLW13 0.15250000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127548 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



C13CPD CDC13 {D:\NMR_DATA} RY 20



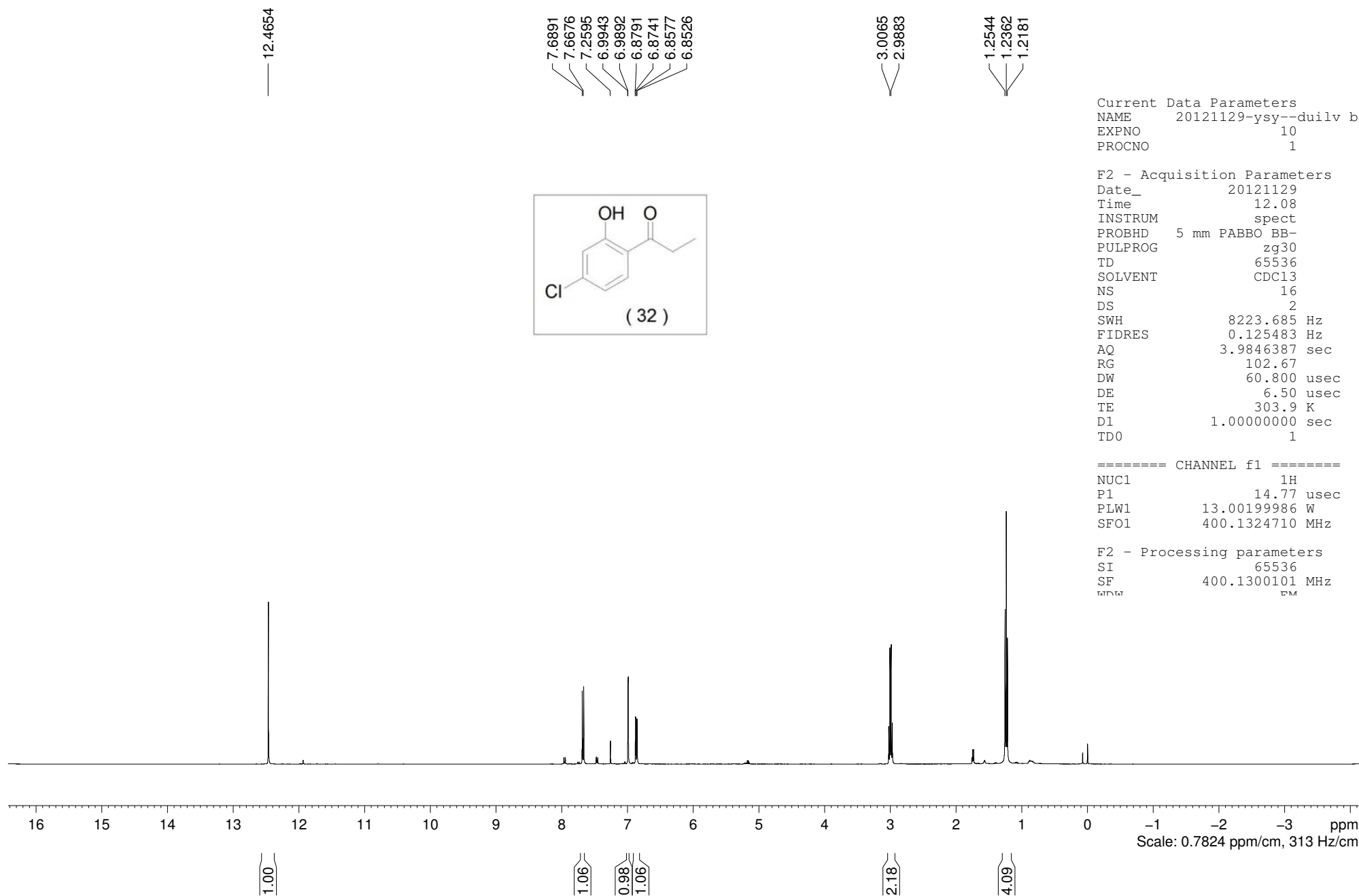
Current Data Parameters
NAME SG-7-2912-C-20121127
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121127
Time 15.31
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 184.16
DW 20.800 usec
DE 6.50 usec
TE 299.8 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

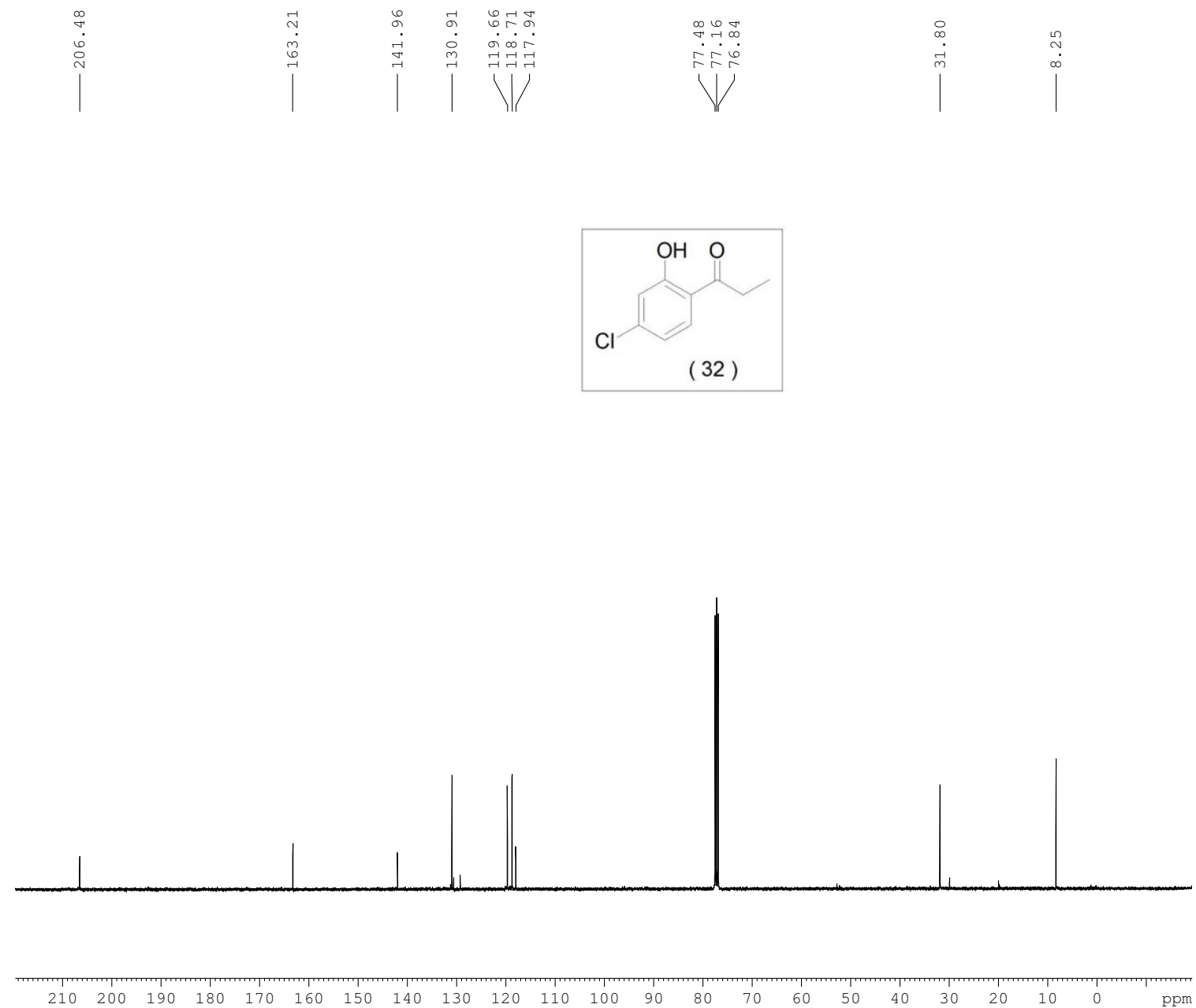
===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127541 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



C13CPD CDC13 {D:\NMR_DATA} RY 27



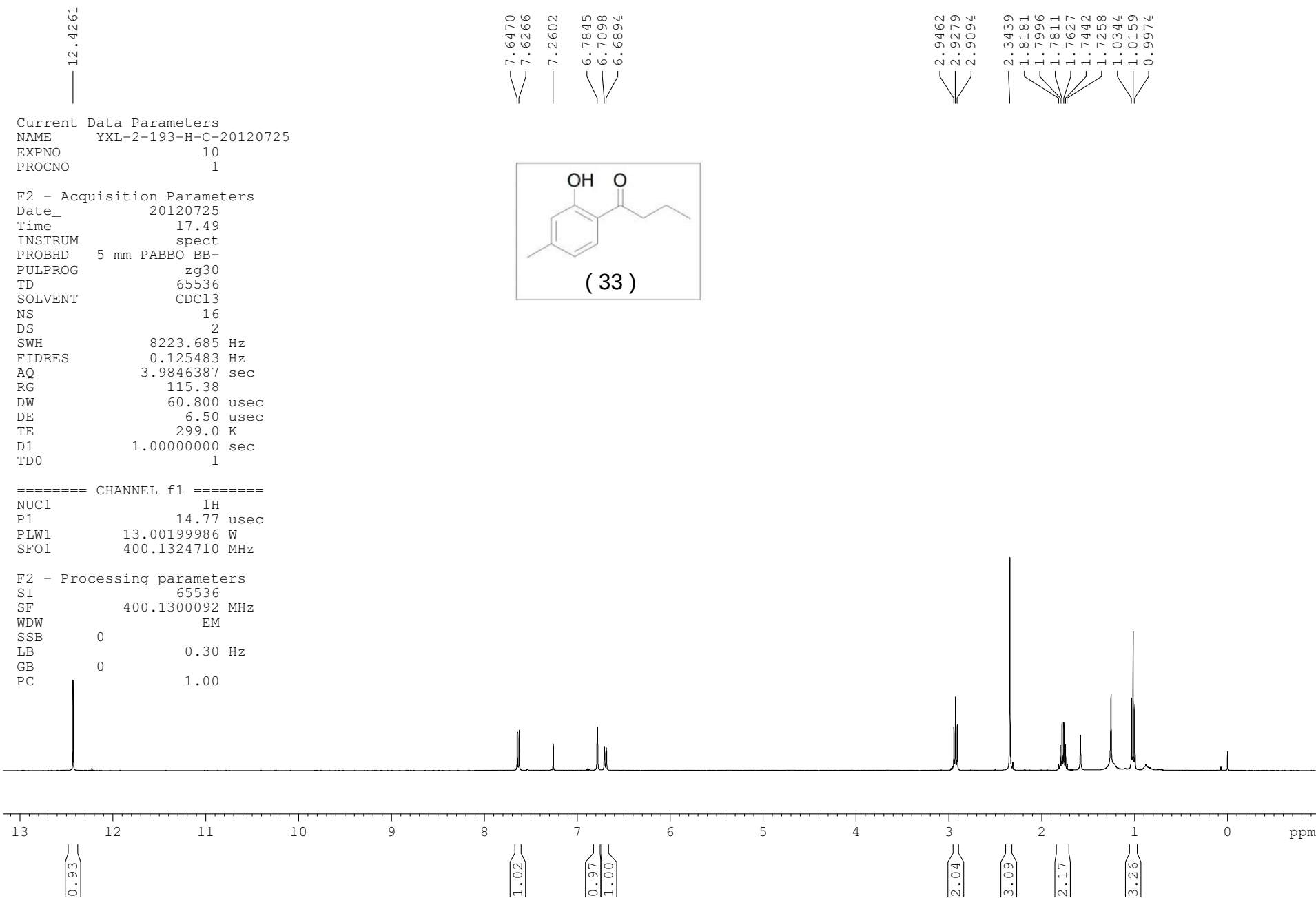
Current Data Parameters
NAME 20121129-ysy--duilv benbington--m-oh
EXPNO 11
PROCNO 1

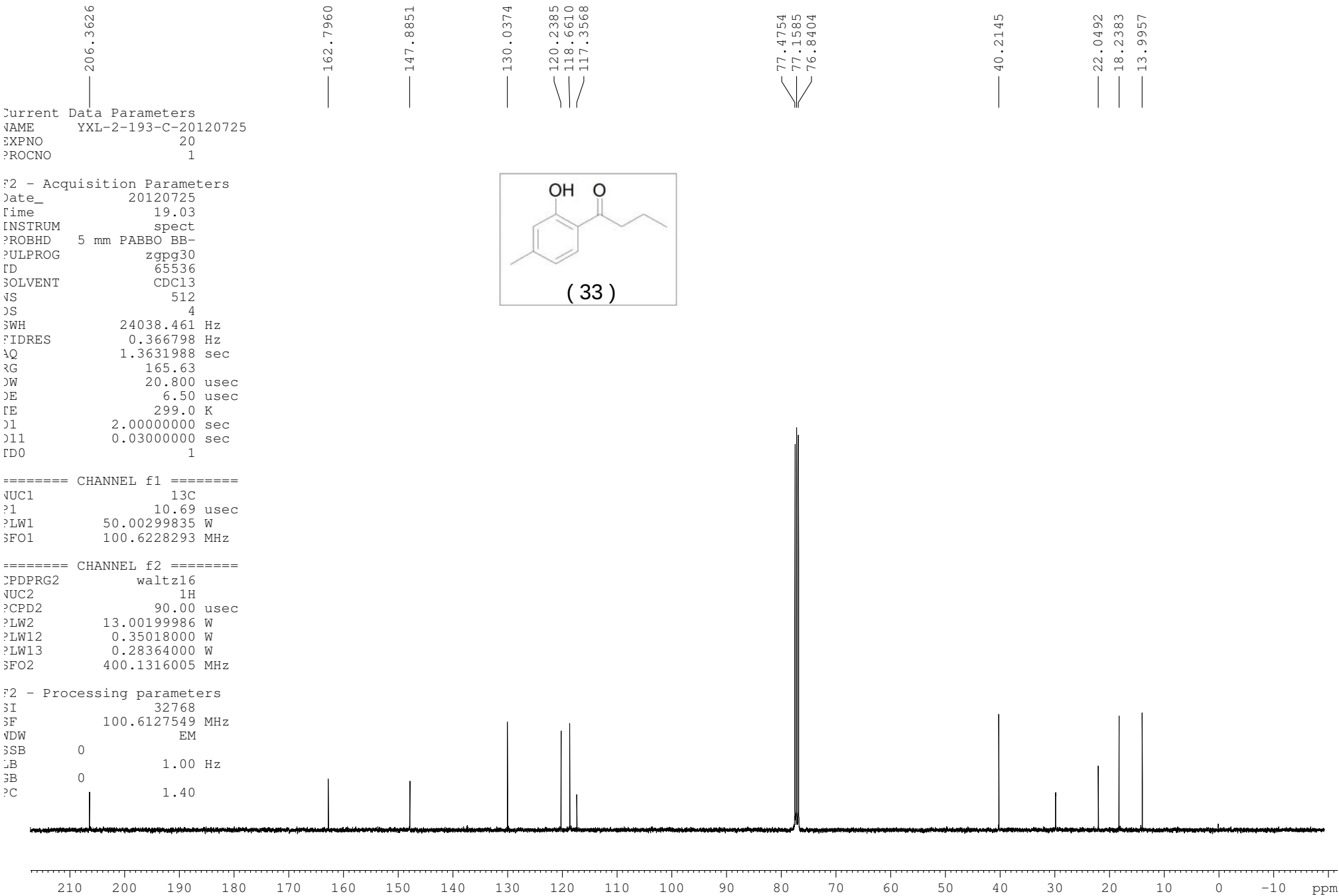
F2 - Acquisition Parameters
Date_ 20121129
Time 12.38
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

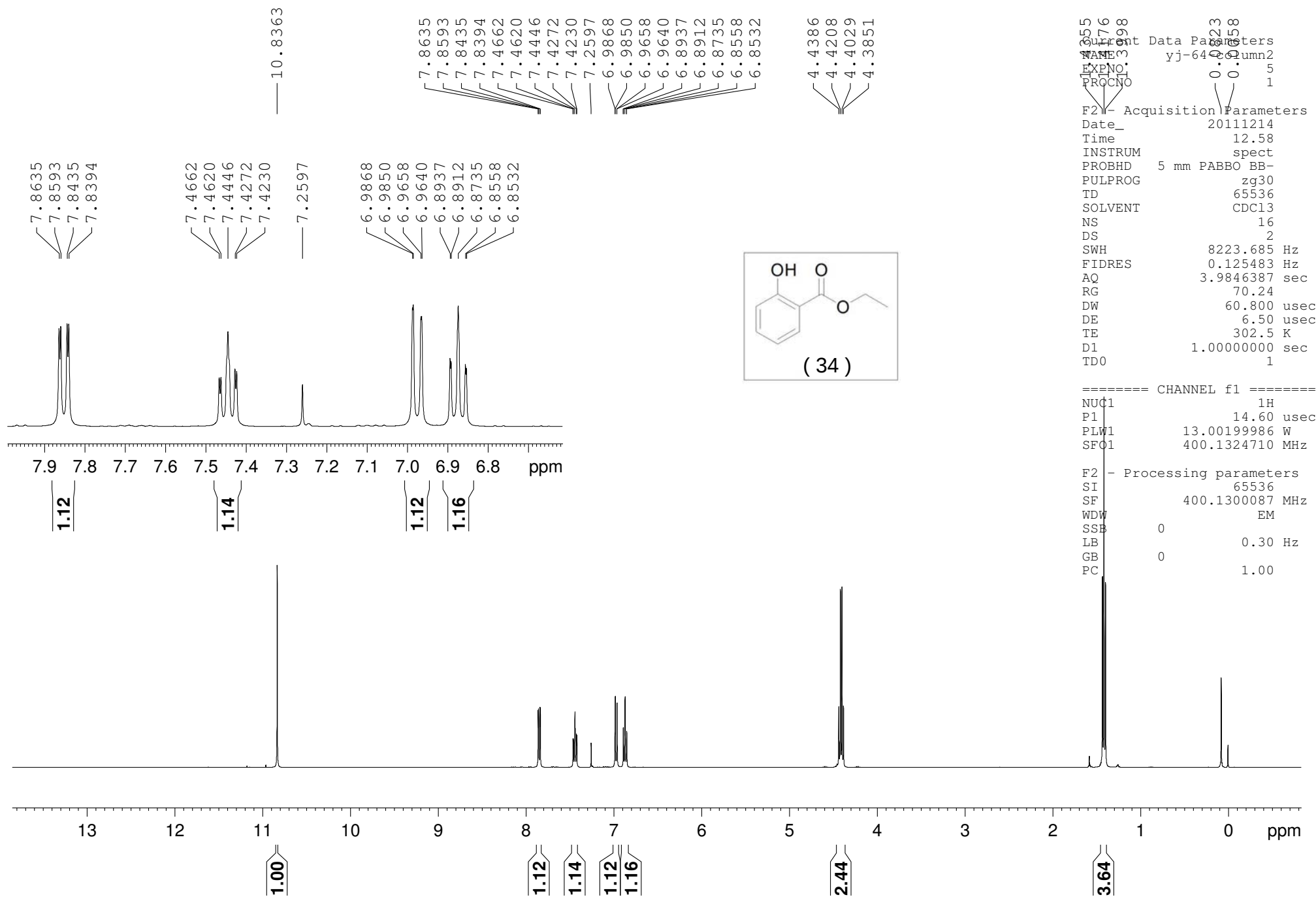
===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

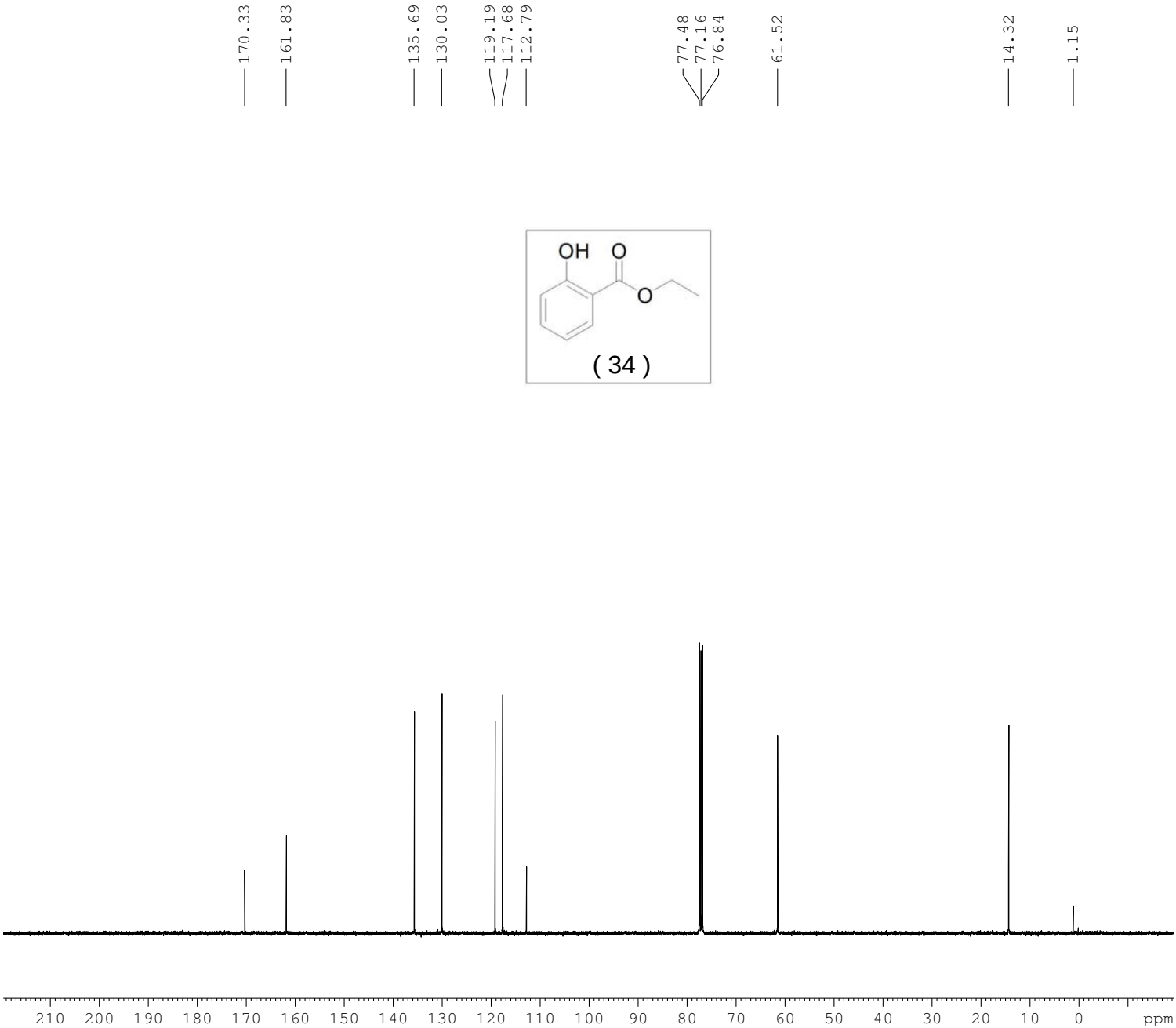
===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127547 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40









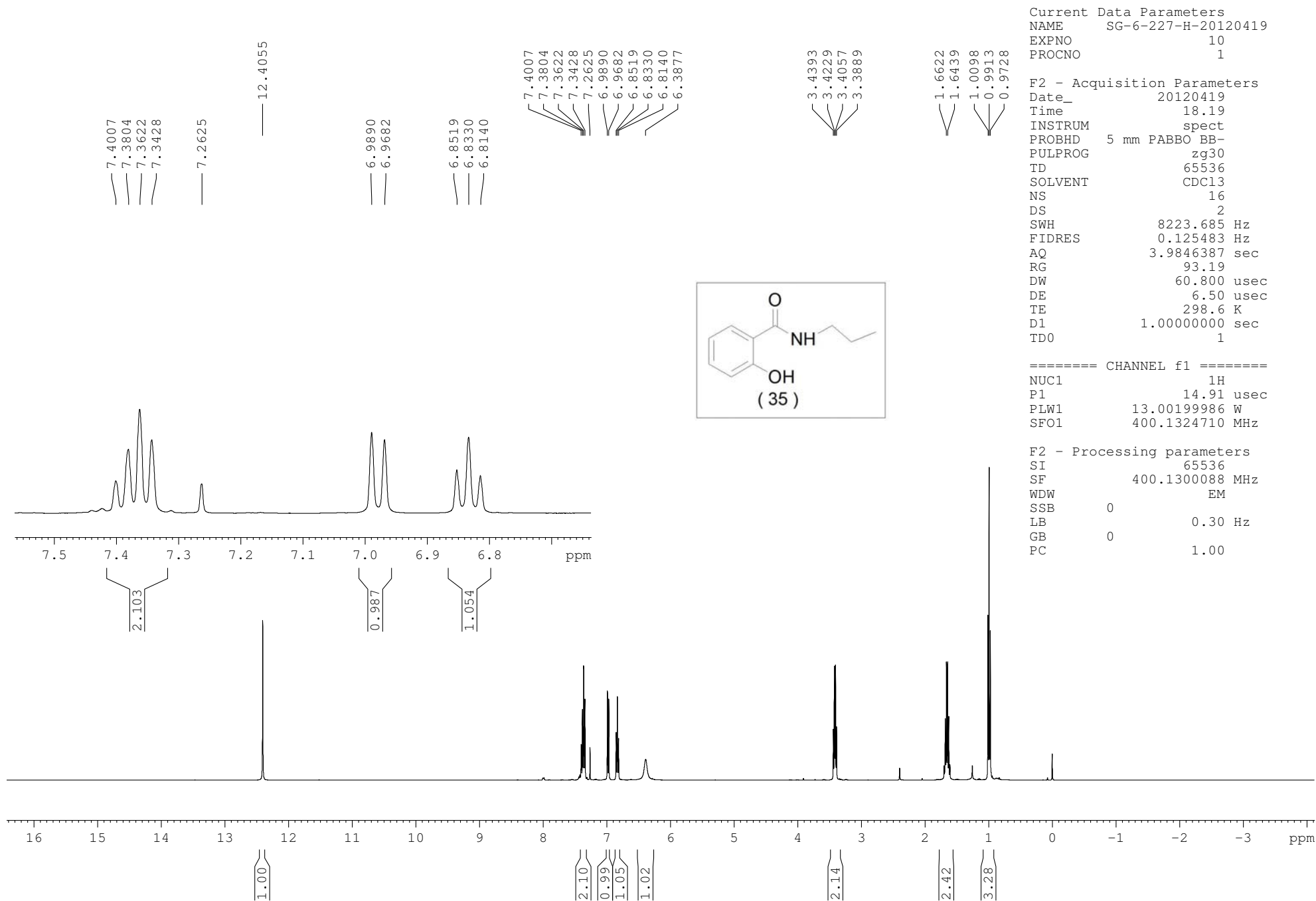
Current Data Parameters
NAME yj-64-column2-C
EXPNO 10
PROCNO 1

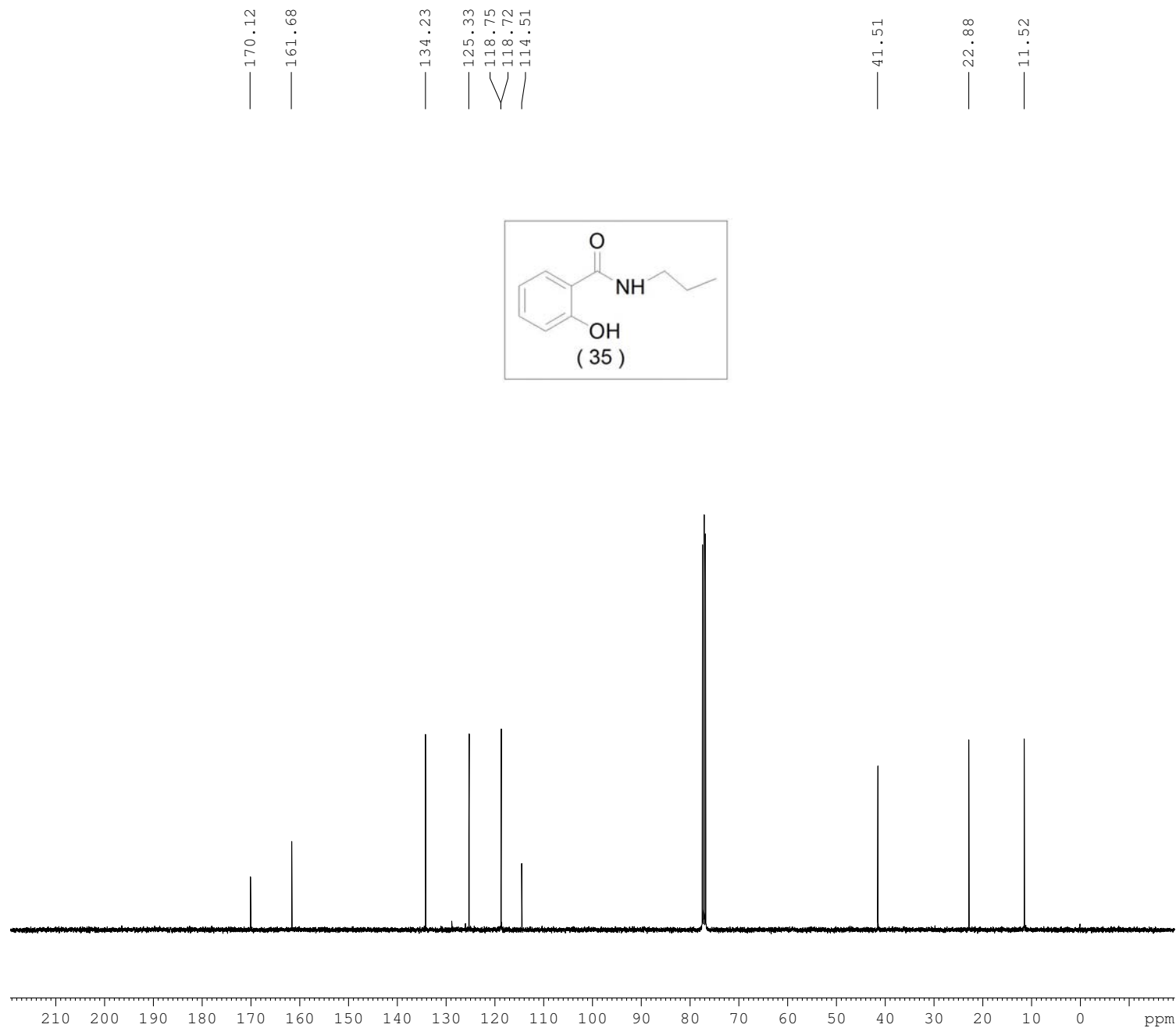
F2 - Acquisition Parameters
Date_ 20111214
Time 13.22
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 400
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 302.5 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.00 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.36116999 W
PLW13 0.29255000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127543 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





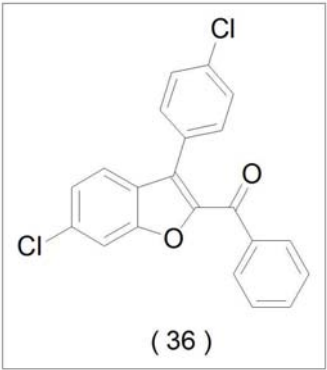
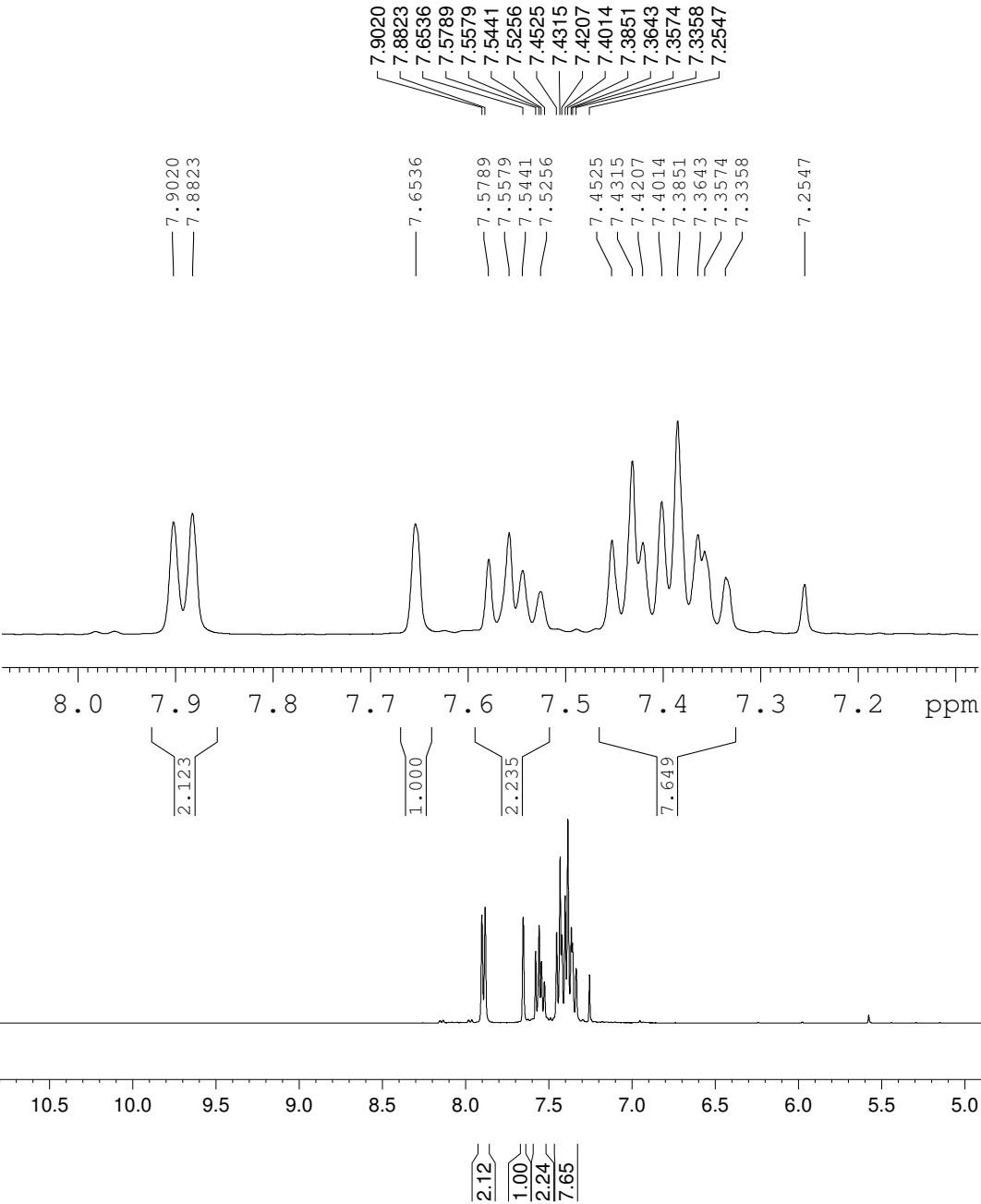
Current Data Parameters
NAME SG-6-227-C-20120419
EXPNO 20
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120419
Time 21.02
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.7 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.31 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35685000 W
PLW13 0.28904000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127563 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40



Current Data Parameters
NAME SG-8-1-H-20121126
EXPNO 10
PROCNO 1

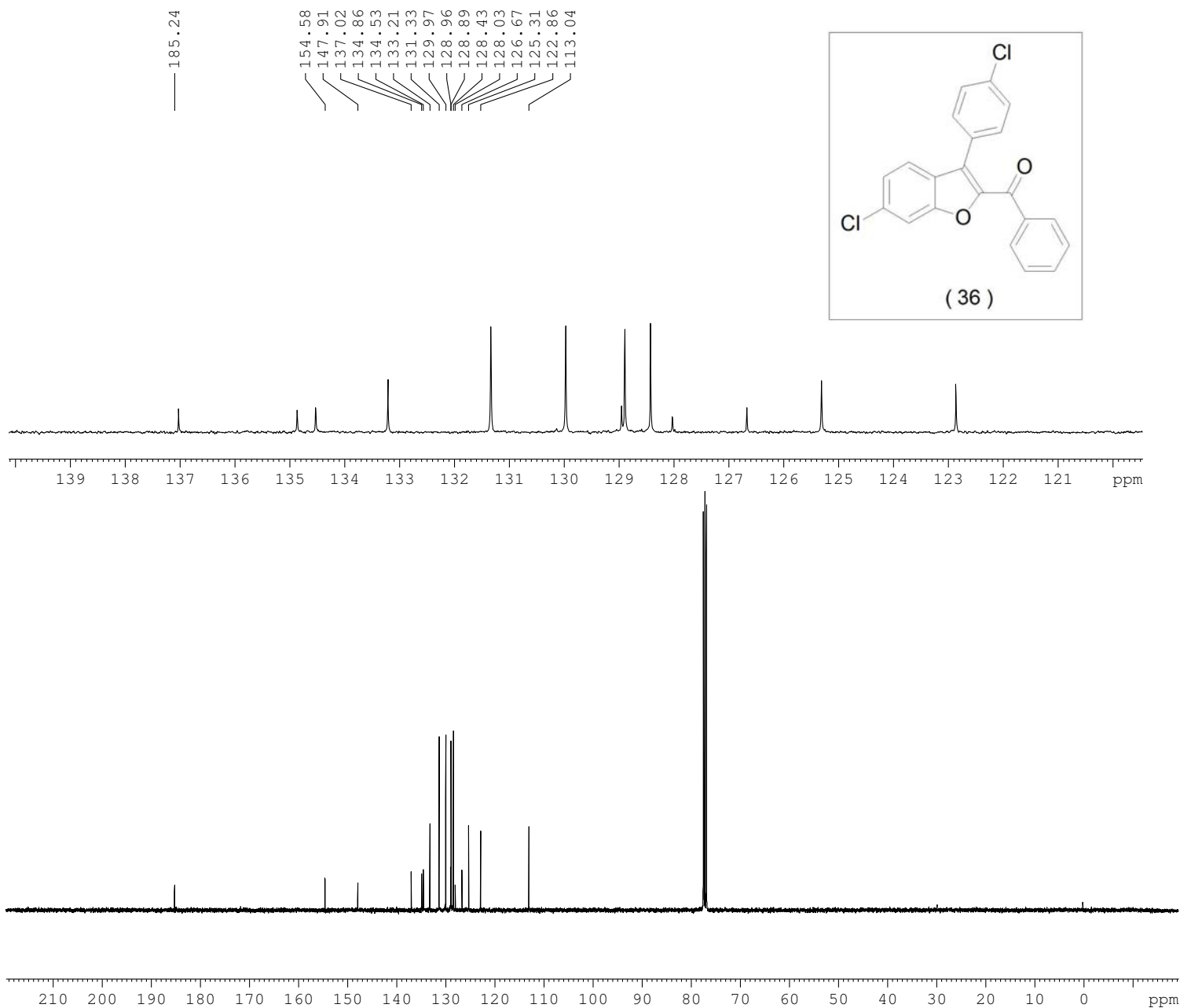
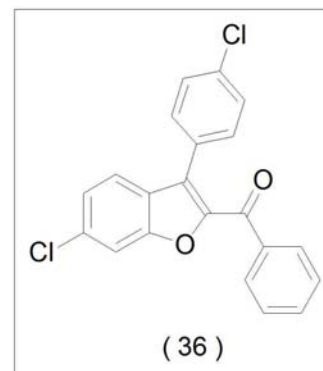
F2 - Acquisition Parameters
Date_ 20121127
Time 7.32
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 144.49
DW 60.800 usec
DE 6.50 usec
TE 299.9 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300123 MHz
WDW EM

Scale: 0.4197 ppm/cm, 167.9 Hz/cm

C13CPD CDC13 {D:\NMR_DATA} RY 19



Current Data Parameters

NAME SG-8-1-C-20121127
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters

Date_ 20121127
Time 11.32
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====

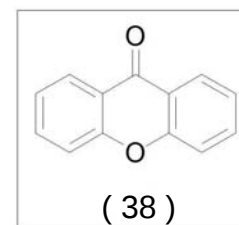
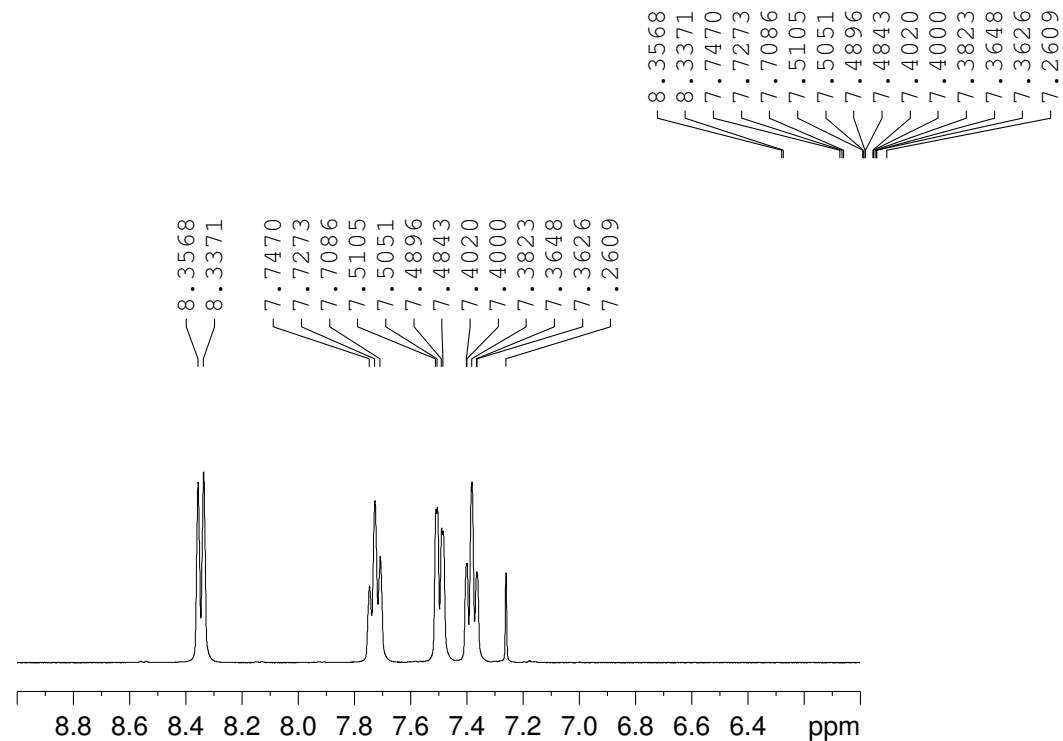
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====

CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters

SI 32768
SF 100.6127554 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

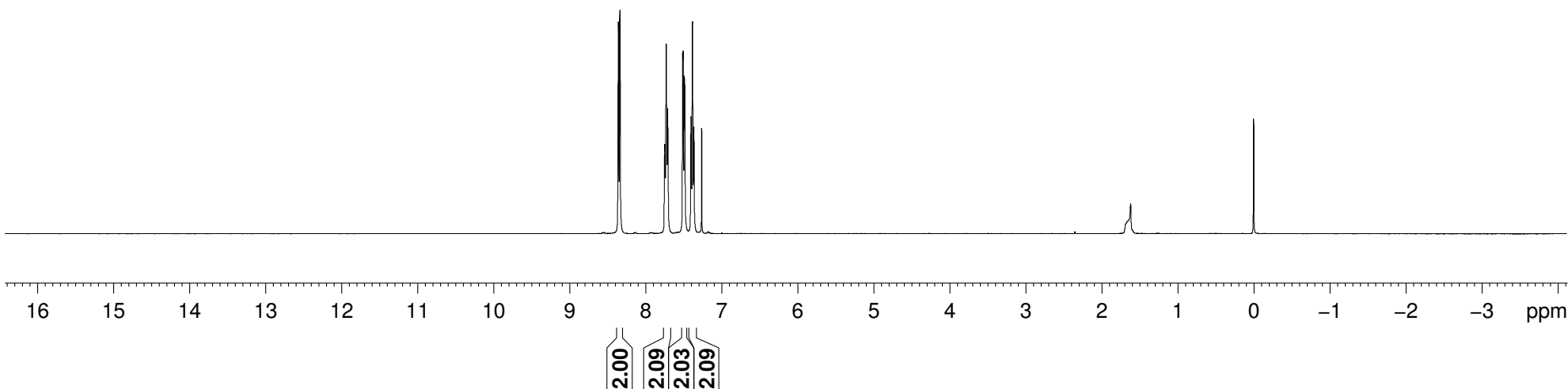


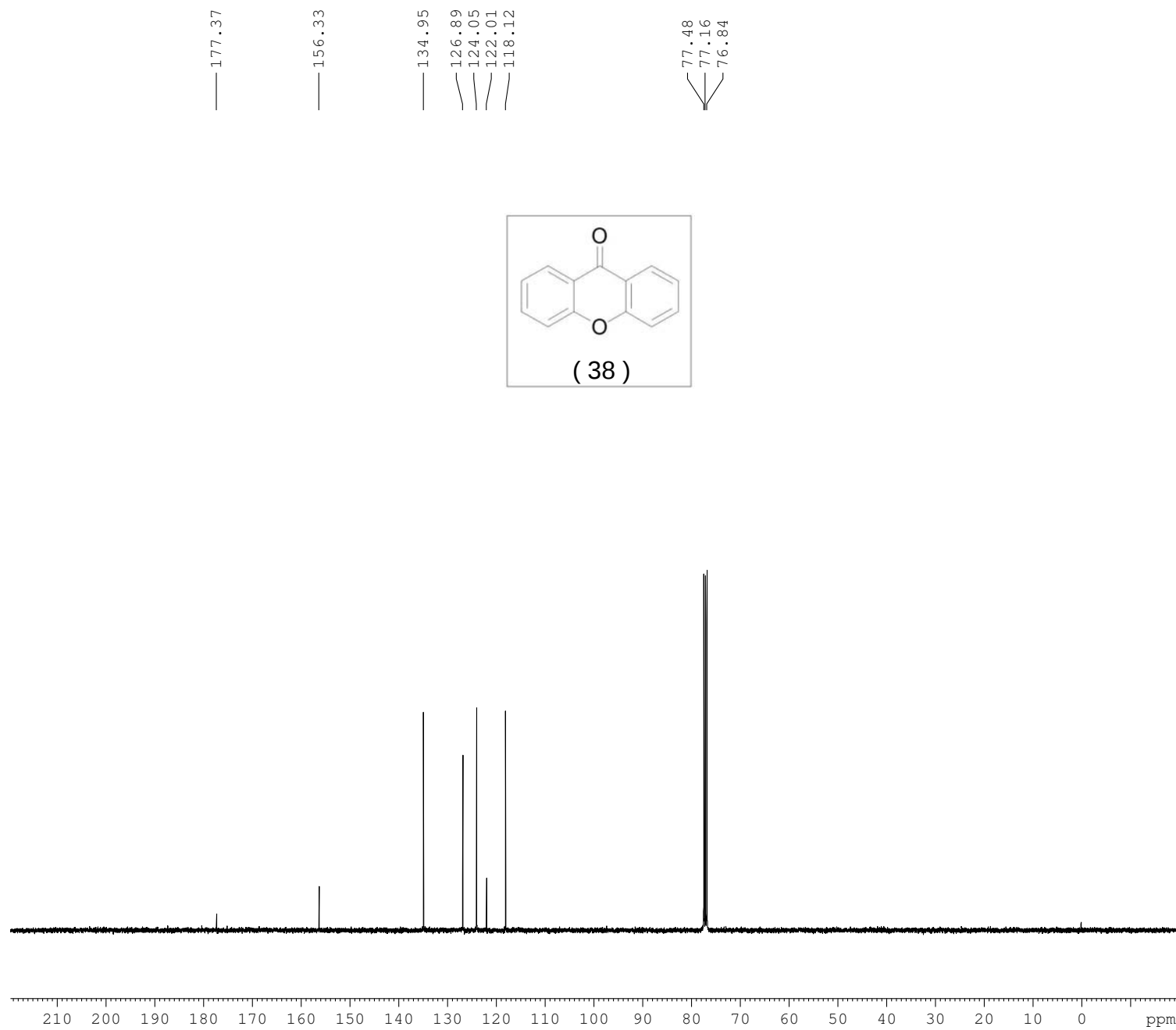
Current Data Parameters
NAME SG-7-Xanthone-H-2012
EXPNO 10
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120815
Time 2.34
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 165.63
DW 60.800 usec
DE 6.50 usec
TE 299.0 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300093 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00





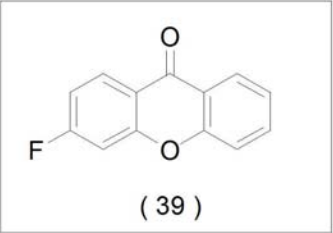
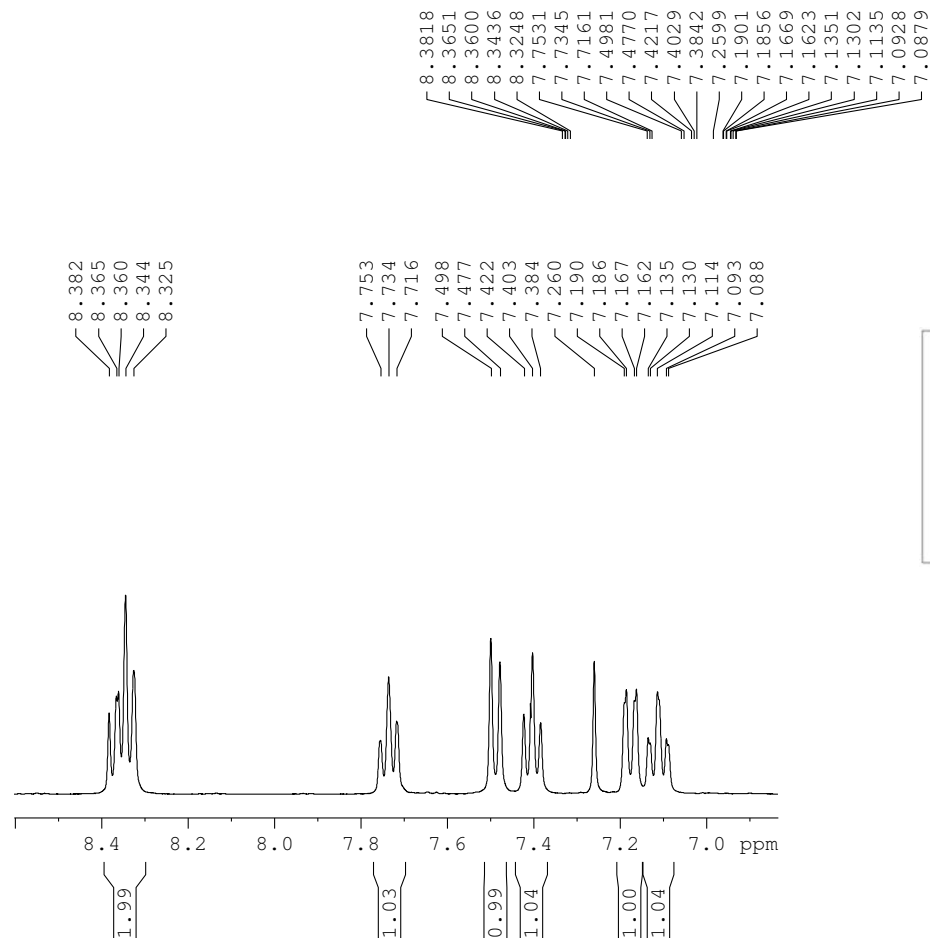
Current Data Parameters
NAME SG-7-Xanthone-C-20120814
EXPNO 12
PROCNO 1

F2 - Acquisition Parameters
Date_ 20120815
Time 3.04
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 512
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 299.0 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.69 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127554 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

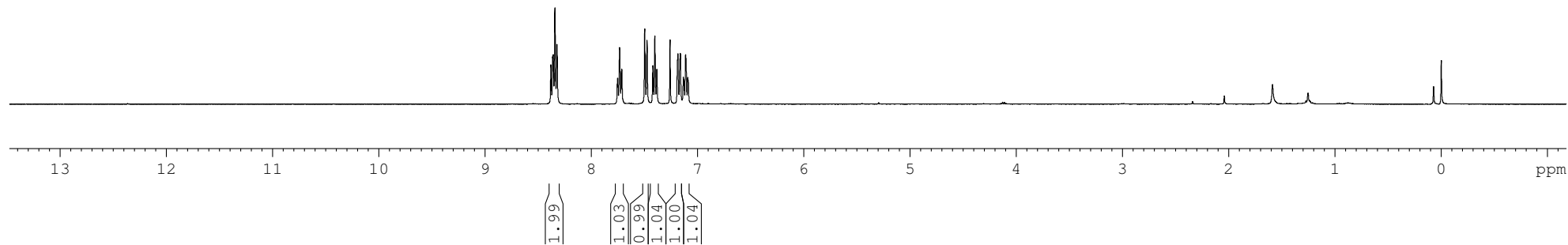


Current Data Parameters
NAME 20121124-ysy--1--171
EXPNO 10
PROCNO 1

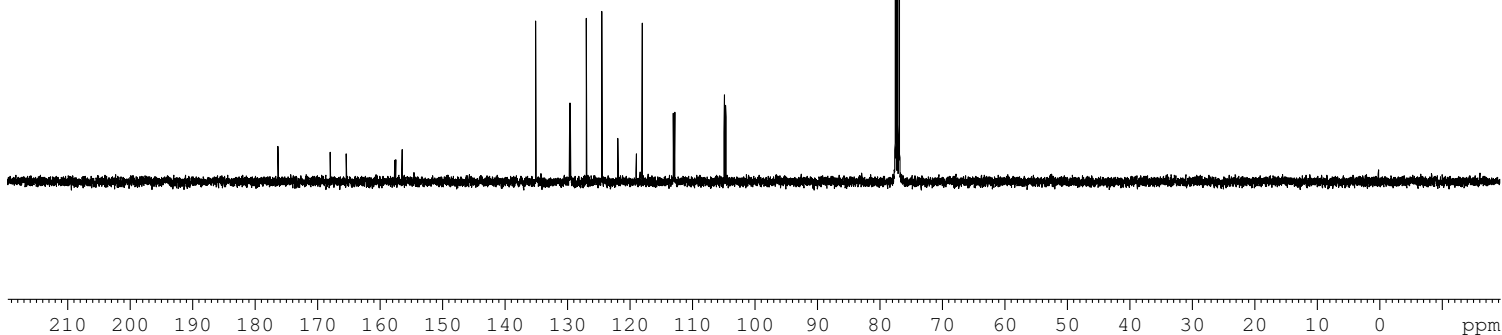
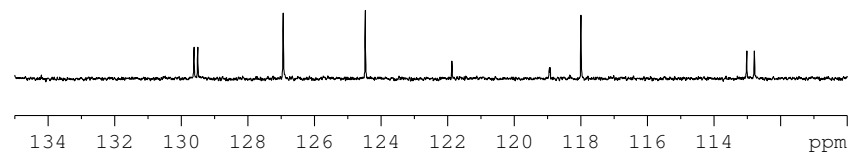
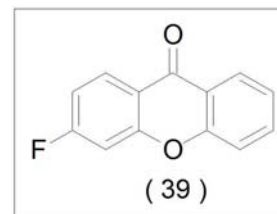
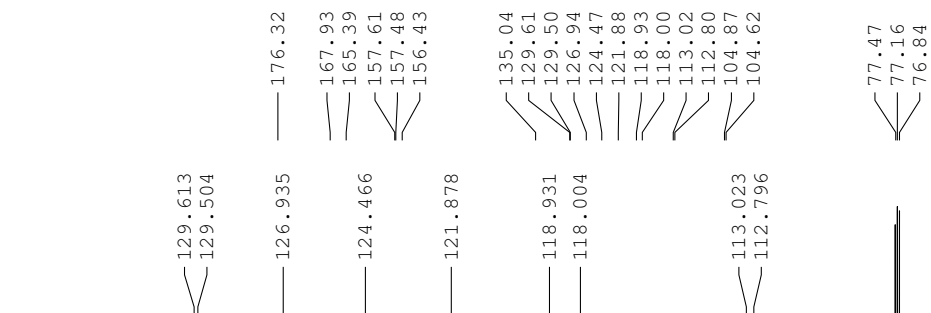
F2 - Acquisition Parameters
Date_ 20121124
Time 21.49
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 8223.685 Hz
FIDRES 0.125483 Hz
AQ 3.9846387 sec
RG 209.25
DW 60.800 usec
DE 6.50 usec
TE 299.8 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 1H
P1 14.77 usec
PLW1 13.00199986 W
SFO1 400.1324710 MHz

F2 - Processing parameters
SI 65536
SF 400.1300104 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



C13CPD CDC13 {D:\NMR_DATA} RY 15



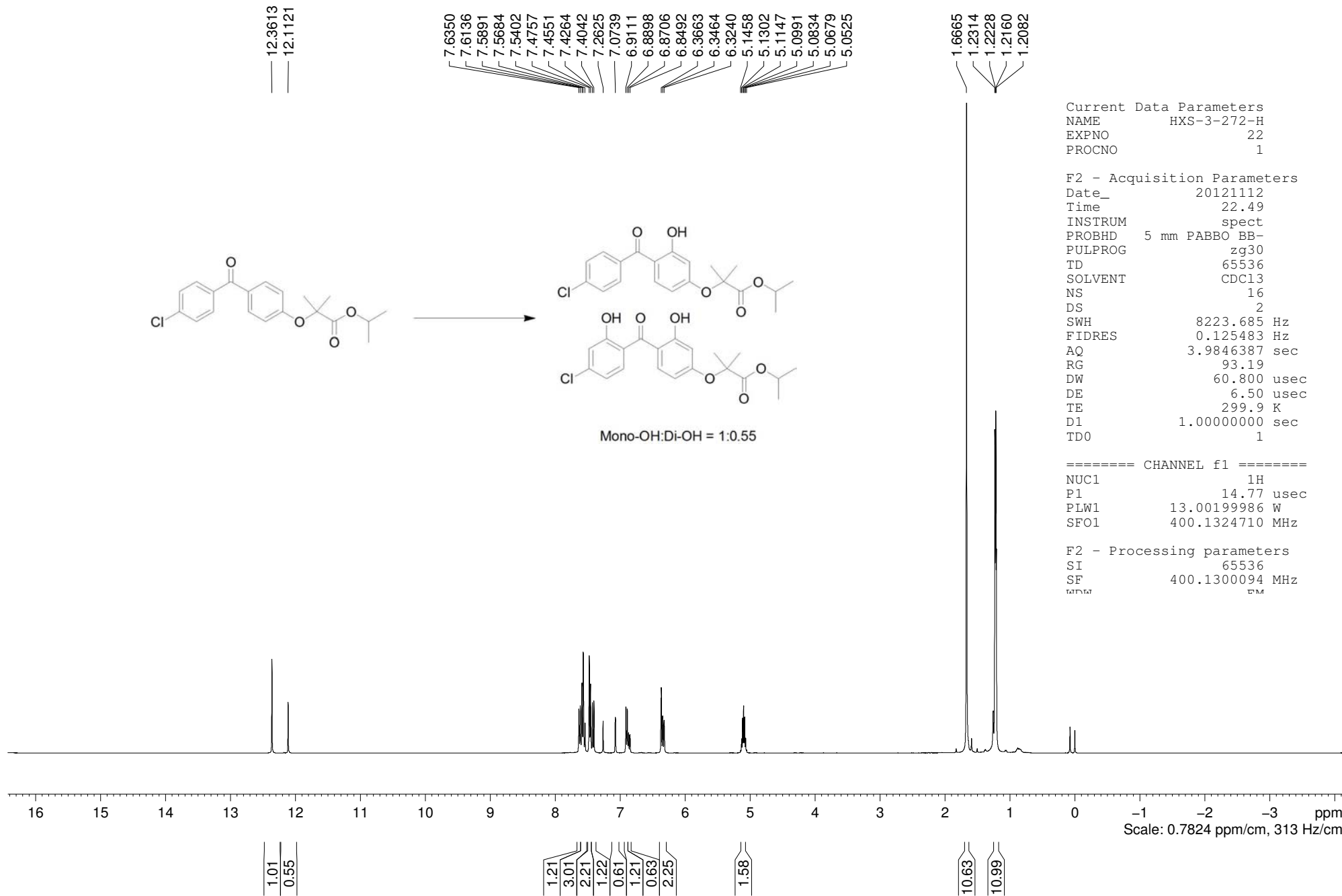
Current Data Parameters
NAME 20121124-ysy--1--171
EXPNO 11
PROCNO 1

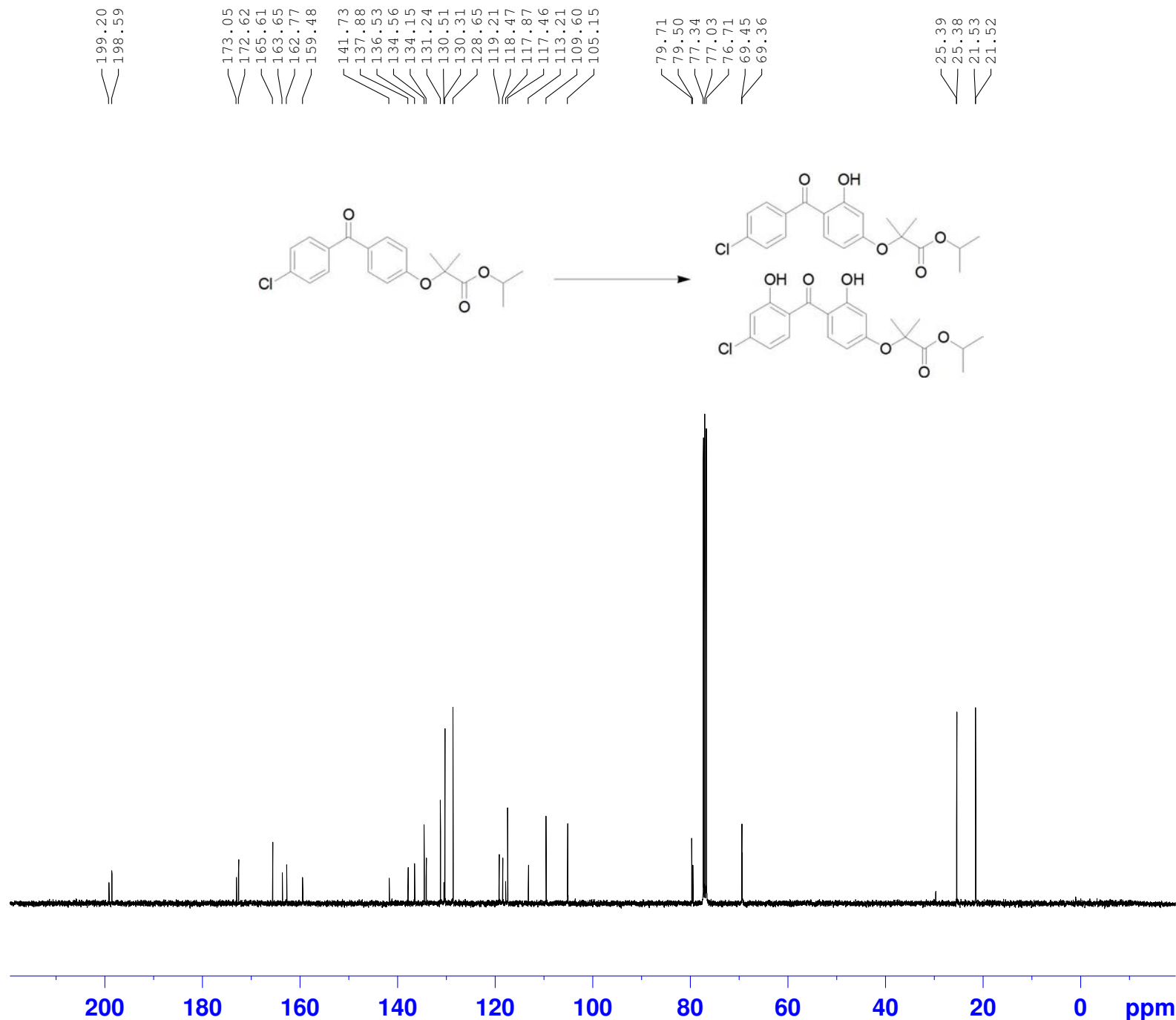
F2 - Acquisition Parameters
Date_ 20121126
Time 12.00
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 465
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 165.63
DW 20.800 usec
DE 6.50 usec
TE 673.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127549 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40





Current Data Parameters
NAME HXS-3-272-C
EXPNO 30
PROCNO 1

F2 - Acquisition Parameters
Date_ 20121112
Time 23.18
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDC13
NS 400
DS 4
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631988 sec
RG 209.25
DW 20.800 usec
DE 6.50 usec
TE 299.9 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 =====
NUC1 13C
P1 10.30 usec
PLW1 50.00299835 W
SFO1 100.6228293 MHz

===== CHANNEL f2 =====
CPDPRG2 waltz16
NUC2 1H
PCPD2 90.00 usec
PLW2 13.00199986 W
PLW12 0.35018000 W
PLW13 0.28364000 W
SFO2 400.1316005 MHz

F2 - Processing parameters
SI 32768
SF 100.6127690 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

