

Supporting Information

Ligand and Counteranion Enabled Regiodivergent C-H Bond Functionalization of Naphthols with α -Aryl- α -Diazoesters

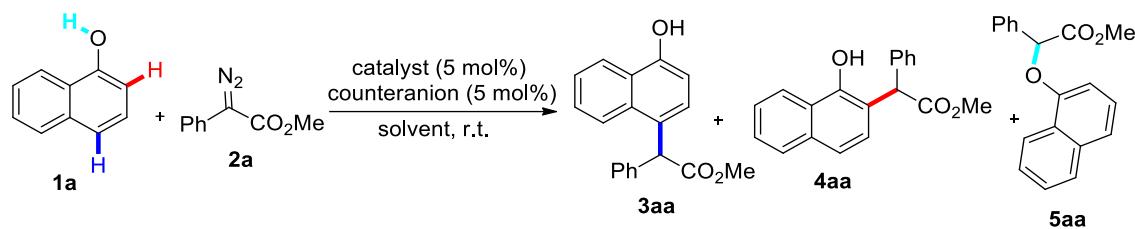
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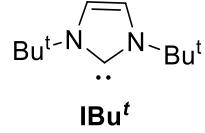
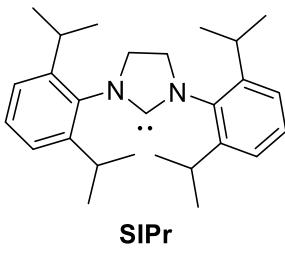
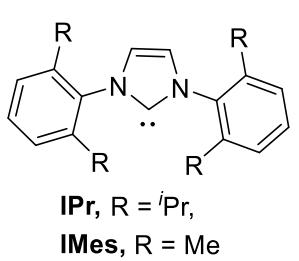
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1. Table S1: Optimization of Reaction Conditions.



Entry	Catalyst	Counteranion	Solvent	Yield(%) ^a
				3aa/4aa/5aa
1	Ph ₃ P AuCl	AgSbF ₆	CH ₂ Cl ₂	57/24/0
2	IPrAuCl	AgSbF ₆	CH ₂ Cl ₂	32/43/0
3	(2,4- <i>t</i> Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgSbF ₆	CH ₂ Cl ₂	43/15/0
4	(C ₆ F ₅) ₃ PAuCl	AgSbF ₆	CH ₂ Cl ₂	14/14/0
5	(4-CF ₃ C ₆ H ₄) ₃ PAuCl	AgSbF ₆	CH ₂ Cl ₂	52/18/0
6	JohnphosAuCl	AgSbF ₆	CH ₂ Cl ₂	53/19/0
7	X-PhosAuCl	AgSbF ₆	CH ₂ Cl ₂	50/31/0
8	'BuXPhosAuCl	AgSbF ₆	CH ₂ Cl ₂	49/27/0
9	PicAuCl ₂	AgSbF ₆	CH ₂ Cl ₂	NR/NR/NR
10	JohnphosAuCl	AgBF ₄	CH ₂ Cl ₂	68/8/0
11	JohnphosAuCl	AgOTf	CH ₂ Cl ₂	70/0/0
12	JohnphosAuCl	AgNTf ₂	CH ₂ Cl ₂	71/25/0
13	JohnphosAuCl	NaBAR _F	CH ₂ Cl ₂	26/54/0
14	JohnphosAuCl	AgOTf	DCE	73/7/0
15	JohnphosAuCl	AgOTf	Toluene	67/0/0
16	JohnphosAuCl	AgOTf	THF	90 (87)/0/0
17	JohnphosAuCl	AgOTf	CH ₃ CN	54/0/0
18	JohnphosAuCl	AgOTf	Ether	80/0/0
19	JohnphosAuCl	AgOTf	1,4-dioxane	47/21/0
20	JohnphosAuCl	-	CH ₂ Cl ₂	NR/NR/NR
21	-	AgOTf	CH ₂ Cl ₂	0/0/0
22	SIPrAuCl	AgSbF ₆	CH ₂ Cl ₂	32/48/0

23	IBu ^t AuCl	AgSbF ₆	CH ₂ Cl ₂	42/30/0
24	IMesAuCl	AgSbF ₆	CH ₂ Cl ₂	30/26/0
25	IPrAuCl	AgOTf	CH ₂ Cl ₂	52/18/0
26	IPrAuCl	AgBF ₄	CH ₂ Cl ₂	45/28/0
27	IPrAuCl	AgNTf ₂	CH ₂ Cl ₂	25/49/0
28	IPrAuCl	NaBAr _F	CH ₂ Cl ₂	13/90(82) /0
29	IPrAuCl	NaBAr _F	DCE	28/64/0
30	IPrAuCl	NaBAr _F	Toluene	28/65/0
31	IPrAuCl	NaBAr _F	CH ₃ CN	NR/NR/NR
32	IPrAuCl	-	CH ₂ Cl ₂	NR/NR/NR
33	-	NaBAr _F	CH ₂ Cl ₂	NR/NR/NR
34	Ph ₃ PAuCl	AgOTf	CH ₂ Cl ₂	66/16/0
35	(2,4-'Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgOTf	CH ₂ Cl ₂	60/12/0
36	Ph ₃ PAuCl	NaBAr _F	CH ₂ Cl ₂	42/42/0
37	(2,4-'Bu ₂ C ₆ H ₃ O) ₃ PAuCl	NaBAr _F	CH ₂ Cl ₂	19/14/0
38	Cu(CH ₃ CN) ₄ PF ₆	-	CH ₂ Cl ₂	0/12 ^b /0
39	Cu(OTf) ₂	-	CH ₂ Cl ₂	0/48 ^b /0
40	Rh(OAc) ₂	-	CH ₂ Cl ₂	0/54 ^b /9
41	Pd(OAc) ₂	-	CH ₂ Cl ₂	NR/NR/NR
42	(C ₆ F ₅) ₃ B	-	CH ₂ Cl ₂	0/24/0



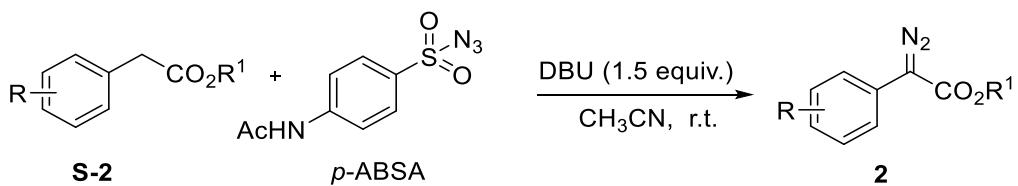
Reaction Conditions: **1a** (0.6 mmol), **2a** (0.4 mmol), catalyst (5 mol%), solvent (5mL), r.t, 15 min. ^aNMR yield, the number in parenthesis is isolated yield.

^bCorresponding lactone was obtained.

2. General Information:

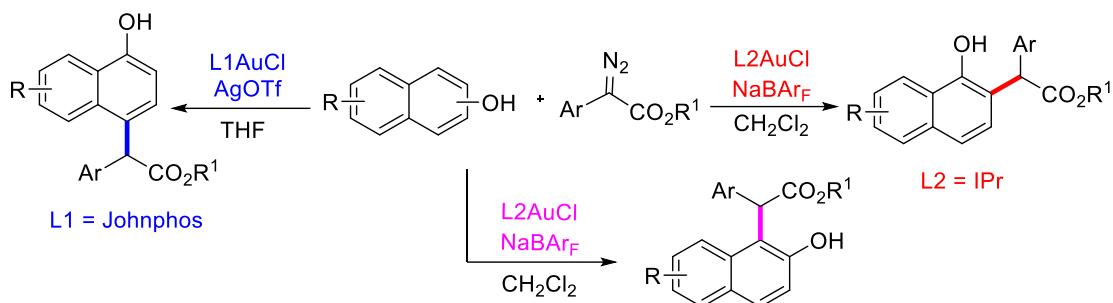
All air- and moisture-sensitive manipulations were carried out with standard Schlenk techniques under nitrogen or in a glove box under nitrogen. ^1H NMR, ^{13}C NMR spectra were measured at 400 MHz and 100 MHz in CDCl_3 . Data for ^1H NMR spectra are reported as follows: chemical shift (ppm, referenced to TMS; s = singlet, d = doublet, t = triplet, dd = doublet of doublets, dt = doublet of triplets, m = multiplet), coupling constant (Hz), and integration. Data for ^{13}C NMR are reported in terms of chemical shift (ppm) relative to residual solvent peak (CDCl_3 : 77.0 ppm). THF, Toluene, 1,4-dioxane, Ether were distilled from sodium and benzophenone prior to be used. CH_2Cl_2 , CH_3CN were distilled from CaH_2 prior to be used.

3. General procedure for synthesis of α -aryl- α -diazoesters



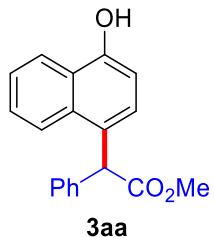
The arylacetate was dissolved in CH_3CN (10 mL/g) and *p*-ABSA (1.5 equiv) and DBU (1.5 equiv.) was added at r.t. The reaction was stirred at r.t. for 12 h. The reaction was diluted with water extracted with ether and dried over Na_2SO_4 . The reaction was concentrated and purified by silica chromatography (PE/EA = 200:1) to obtain the desired diazo compound.

4. General procedure for the C-H bond functionalization of naphthalenols.



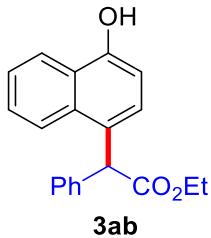
In a dried glass tube, a mixture of LAuCl (0.02 mmol), AgOTf or NaBAr_F (5 mol%) in solvent (4 mL) was stirred at room temperature for 15 mins. Subsequently, naphthalenol (0.6 mmol) was added to the reaction mixture at room temperature. Then a solution of diazo compounds (0.4 mmol) in solvent (1 mL) was introduced into the reaction mixture by a syringe in 15 mins. The mixture was concentrated under reduced pressure and the residue was purified by column chromatography on silica gel (PE/EA = 10:1 to 5:1 or PE/EA = 20:1) to afford the desired product.

1) The synthesis of 3aa



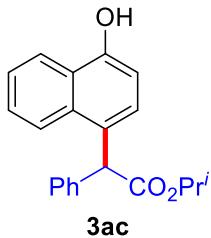
3aa, light red oil, (102.1 mg, 87% yield). ¹H-NMR (400 MHz, CDCl₃) δ 8.24 (d, *J* = 7.6 Hz, 1H), 7.92 (d, *J* = 8.0 Hz, 1H), 7.45-7.55 (m, 2H), 7.27-7.38 (m, 5H), 7.05 (d, *J* = 8.0 Hz, 1H), 6.57 (d, *J* = 7.6 Hz, 1H), 6.14 (bs, 1H), 5.73 (s, 1H), 3.79 (s, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 52.58, 53.17, 107.93, 122.67, 122.89, 124.91, 124.95, 126.39, 126.56, 126.99, 127.33, 128.64, 128.91, 132.61, 138.02, 151.45, 174.14. MS(EI): m/z (%): 292 (M⁺, 17.19), 231 (100); HRMS (EI) calcd. for C₁₉H₁₆O₃: 292.1099, found: 292.1096.

2) The synthesis of 3ab



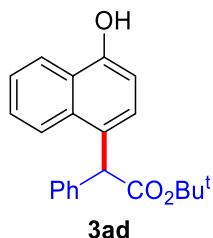
3ab, light red oil, (106.2 mg, 87% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.23 (dd, $J = 8.0$ Hz, 1.2 Hz, 1H), 7.94 (d, $J = 8.0$ Hz, 1H), 7.45-7.55 (m, 2H), 7.25-7.35 (m, 5H), 7.05 (d, $J = 8.0$ Hz, 1H), 6.54 (d, $J = 8.0$ Hz, 1H), 6.21 (bs, 1H), 5.71 (s, 1H), 4.25-4.30 (m, 2H), 1.27 (d, $J = 7.2$ Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 14.10, 53.29, 61.50, 107.96, 122.66, 122.95, 124.85, 124.96, 126.50, 126.55, 126.87, 127.26, 128.60, 128.93, 132.64, 138.12, 151.41, 173.72. MS(EI): m/z (%): 306 (M^+ , 27.00), 233 (100); HRMS (EI) calcd. for $\text{C}_{20}\text{H}_{18}\text{O}_3$: 306.1256, found: 306.1259.

3) The synthesis of 3ac



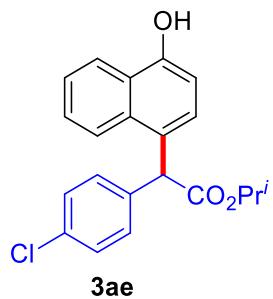
3ac, colorless oil, (117.1 mg, 91% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.22 (d, $J = 8.0$ Hz, 1H), 7.95 (d, $J = 8.0$ Hz, 1H), 7.25-7.55 (m, 7H), 7.03 (d, $J = 7.6$ Hz, 1H), 6.53 (d, $J = 8.0$ Hz, 1H), 5.85-5.95 (m, 1H), 5.66 (s, 1H), 5.14 (hept, $J = 6.4$ Hz, 1H), 1.26 (d, $J = 6.4$ Hz, 3H), 1.23 (d, $J = 6.4$ Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 21.64, 53.47, 68.93, 107.98, 122.61, 123.05, 124.84, 126.51, 126.81, 127.20, 128.57, 128.94, 132.71, 138.22, 151.29. MS(EI): m/z (%): 320 (M^+ , 22.79), 233 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{20}\text{O}_3$: 320.1412, found: 320.1407.

4) The synthesis of 3ad



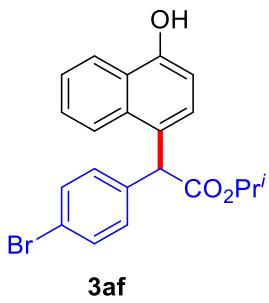
3ad, white solid, (99.5 mg, 74% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.21 (d, $J = 8.0$ Hz, 1H), 7.96 (d, $J = 8.4$ Hz, 1H), 7.45-7.55 (m, 2H), 7.25-7.40 (m, 5H), 6.98 (d, $J = 8.0$ Hz, 1H), 6.49 (d, $J = 8.0$ Hz, 1H), 6.02 (s, 1H), 5.60 (s, 1H), 1.47 (s, 9H); ^{13}C -NMR (100 MHz, CDCl_3) δ 27.95, 54.23, 81.71, 108.02, 122.60, 123.09, 124.77, 124.93, 126.47, 126.69, 127.09, 127.20, 128.54, 128.96, 132.75, 138.48, 151.18, 172.91. MS(EI): m/z (%): 334 (M^+ , 12.89), 233 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{22}\text{O}_3$: 334.1569, found: 334.1575.

5) The synthesis of 3ae



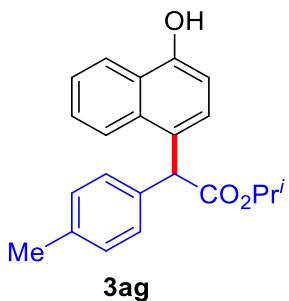
3ae, colorless oil, (134.5 mg, 95% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.21 (d, $J = 7.6$ Hz, 1H), 7.86 (d, $J = 8.0$ Hz, 1H), 7.40-7.50 (m, 2H), 7.20-7.30 (m, 4H), 7.00 (d, $J = 7.6$ Hz, 1H), 6.51 (d, $J = 7.6$ Hz, 1H), 6.19 (s, 1H), 5.61 (s, 1H), 5.13 (hept, $J = 6.4$ Hz, 1H), 1.23 (d, $J = 6.4$ Hz, 3H), 1.22 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.60, 21.63, 52.86, 69.21, 107.89, 122.69, 122.91, 124.96, 124.99, 126.16, 126.33, 126.92, 128.73, 130.28, 132.53, 133.12, 136.75, 151.51, 172.89. MS(EI): m/z (%): 354 (M^+ , 21.99), 356 ($[\text{M}+2]^+$, 6.88), 267 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{19}\text{O}_3\text{Cl}$: 354.1023, found: 354.1024.

6) The synthesis of 3af



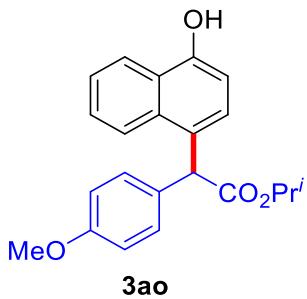
3af, colorless oil, (150.0 mg, 94% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.20-8.25 (m, 1H), 7.87 (d, J = 8.0 Hz, 1H), 7.42-7.52 (m, 4H), 7.20 (d, J = 8.4 Hz, 2H), 7.02 (d, J = 7.6 Hz, 1H), 6.53 (d, J = 8.0 Hz, 1H), 6.10 (s, 1H), 5.60 (s, 1H), 5.14 (hept, J = 6.4 Hz, 1H), 1.25 (d, J = 6.4 Hz, 3H), 1.24 (d, J = 6.4 Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 21.61, 21.64, 52.93, 69.23, 107.90, 121.28, 122.69, 122.90, 124.98, 126.11, 126.35, 126.94, 130.64, 131.68, 132.52, 137.27, 151.48, 172.80. MS(EI): m/z (%): 398 (M^+ , 21.50), 400 ($[\text{M}+2]^+$, 21.42), 231 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{19}\text{O}_3\text{Br}$: 398.0518, found: 398.0520.

7) The synthesis of 3ag



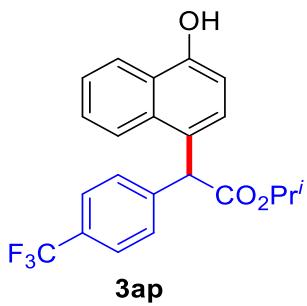
3ag, white solid, (112.9 mg, 84% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.21 (d, J = 8.4 Hz, 1H), 7.95 (d, J = 8.4 Hz, 1H), 7.40-7.55 (m, 2H), 7.23 (d, J = 8.0 Hz, 2H), 7.16 (d, J = 8.0 Hz, 2H), 7.00 (d, J = 8.0 Hz, 1H), 6.47 (d, J = 8.0 Hz, 1H), 6.21 (s, 1H), 5.64 (s, 1H), 5.15 (hept, J = 6.4 Hz, 1H), 2.35 (s, 3H), 1.27 (d, J = 6.4 Hz, 3H), 1.24 (d, J = 6.4 Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 21.05, 21.63, 21.65, 53.11, 68.92, 108.02, 122.63, 123.00, 124.75, 124.95, 126.46, 126.73, 126.83, 128.80, 129.30, 132.68, 135.10, 136.83, 151.27, 173.48. MS(EI): m/z (%): 334 (M^+ , 20.02), 247 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{22}\text{O}_3$: 334.1569, found: 334.1573.

8) The synthesis of 3ao



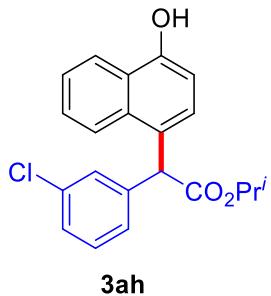
3ao, yellow oil, (112.9 mg, 84% yield). $^1\text{H-NMR}$ (500 MHz, CDCl_3) δ 8.20 (d, $J = 8.0$ Hz, 1H), 7.91 (d, $J = 8.0$ Hz, 1H), 7.40-7.50 (m, 2H), 7.24 (d, $J = 8.5$ Hz, 2H), 7.00 (d, $J = 7.5$ Hz, 1H), 6.86 (d, $J = 8.5$ Hz, 2H), 6.50 (d, $J = 8.0$ Hz, 1H), 5.58 (s, 1H), 5.12 (hept, $J = 6.0$ Hz, 1H), 3.77 (s, 3H), 1.24 (d, $J = 6.0$ Hz, 3H), 1.21 (d, $J = 6.0$ Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 21.64, 52.68, 55.20, 68.88, 107.92, 113.96, 122.64, 123.00, 124.77, 124.94, 126.30, 126.73, 126.88, 129.97, 130.19, 132.61, 151.31, 158.61, 173.52. HRMS (EI) calcd. for $[\text{C}_{22}\text{H}_{22}\text{O}_4+\text{Na}]^+$: 373.1410, found: 373.1401.

9) The synthesis of 3ap



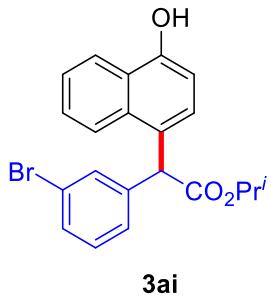
3ap, yellow oil, (125.9 mg, 81% yield). $^1\text{H-NMR}$ (500 MHz, CDCl_3) δ 8.25 (d, $J = 8.5$ Hz, 1H), 7.89 (d, $J = 8.5$ Hz, 1H), 7.40-7.65 (m, 6H), 7.01 (d, $J = 8.0$ Hz, 1H), 6.60-6.70 (m, 1H), 6.50 (d, $J = 8.0$ Hz, 1H), 5.74 (s, 1H), 5.19 (hept, $J = 6.5$ Hz, 1H), 1.28 (d, $J = 6.5$ Hz, 3H), 1.27 (d, $J = 6.5$ Hz, 3H); $^{13}\text{C-NMR}$ (125 MHz, CDCl_3) δ 21.54, 21.59, 53.25, 69.58, 107.87, 122.74, 122.80, 124.08 (q, $J = 268.8$ Hz, 1C), 124.99, 125.05, 125.41 (q, $J = 3.8$ Hz, 1C), 125.51 (q, $J = 3.8$ Hz, 1C), 126.48, 126.73, 129.29, 129.47 (q, $J = 32.5$ Hz, 1C), 132.47, 142.20, 151.77, 172.91. HRMS (ESI) calcd. for $[\text{C}_{22}\text{H}_{19}\text{F}_3\text{O}_3+\text{Na}]^+$: 411.1178, found: 411.1177.

10) The synthesis of 3ah



3ah, white solid, (120.3 mg, 85% yield). ¹H-NMR (400 MHz, CDCl₃) δ 8.20-8.25 (m, 1H), 7.87 (d, *J* = 7.6 Hz, 1H), 7.45-7.55 (m, 2H), 7.15-7.35 (m, 4H), 7.05 (d, *J* = 7.6 Hz, 1H), 6.57 (d, *J* = 7.6 Hz, 1H), 5.83 (s, 1H), 5.61 (s, 1H), 5.13 (hept, *J* = 6.4 Hz, 1H), 1.24 (d, *J* = 6.4 Hz, 6H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.60, 21.66, 53.06, 69.22, 107.95, 122.67, 122.92, 124.95, 125.03, 126.08, 126.38, 127.01, 127.13, 127.50, 129.08, 129.76, 132.56, 134.43, 140.29, 151.43, 172.52. MS(EI): m/z (%): 354 (M⁺, 23.69), 356 ([M+2]⁺, 8.41), 267 (100); HRMS (EI) calcd. for C₂₁H₁₉O₃Cl: 354.1023, found: 354.1028.

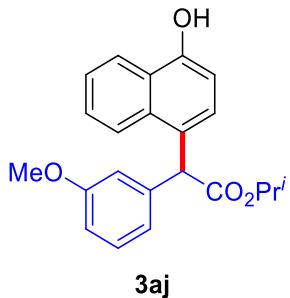
11) The synthesis of 3ai



3ai, white solid, (136.1 mg, 85% yield). ¹H-NMR (400 MHz, CDCl₃) δ 8.22-8.26 (m, 1H), 7.87 (d, *J* = 8.0 Hz, 1H), 7.45-7.55 (m, 3H), 7.40 (d, *J* = 8.0 Hz, 1H), 7.16-7.26 (m, 2H), 7.07 (d, *J* = 8.0 Hz, 1H), 6.60 (d, *J* = 8.0 Hz, 1H), 5.87 (bs, 1H), 5.60 (s, 1H), 5.12 (hept, *J* = 6.4 Hz, 1H), 1.23 (d, *J* = 6.4 Hz, 6H). ¹³C-NMR (100 MHz, CDCl₃) δ 21.60, 21.67, 53.03, 69.14, 107.96, 122.66, 122.98, 124.99, 125.06, 126.19, 126.40, 127.03, 127.59, 130.05, 130.41, 131.96, 132.60, 140.67, 151.45, 172.32. MS(EI): m/z (%): 398 (M⁺, 17.19), 400 ([M+2]⁺, 16.89), 231 (100); HRMS (EI) calcd. for

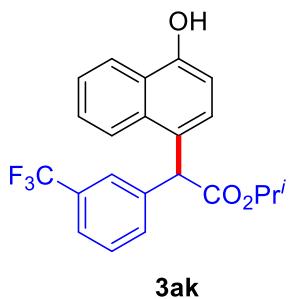
$C_{21}H_{19}O_3Br$: 398.0518, found: 398.0521.

12) The synthesis of 3aj



3aj, white solid, (122.6 mg, 88% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.19 (d, $J = 8.0$ Hz, 1H), 7.91 (d, $J = 8.4$ Hz, 1H), 7.40-7.50 (m, 2H), 7.21-7.25 (m, 1H), 6.99 (d, $J = 8.0$ Hz, 1H), 6.85-6.95 (m, 2H), 6.81 (dd, $J = 8.4$ Hz, 2.4 Hz, 1H), 6.45 (d, $J = 8.0$ Hz, 1H), 6.22 (s, 1H), 5.61 (s, 1H), 5.12 (hept, $J = 6.4$ Hz, 1H), 3.74 (s, 3H), 1.24 (d, $J = 6.4$ Hz, 3H), 1.21 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 21.63, 21.65, 53.44, 55.16, 69.01, 107.97, 112.59, 114.86, 121.43, 122.66, 122.92, 124.77, 124.93, 126.43, 126.52, 126.77, 129.53, 132.67, 139.67, 151.38, 159.67, 173.13. MS(EI): m/z (%): 350 (M^+ , 23.36), 263 (100); HRMS (EI) calcd. for $C_{22}H_{22}O_4$: 350.1518, found: 350.1524.

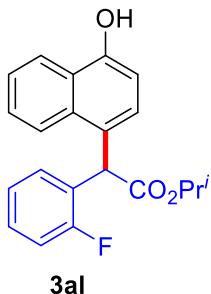
13) The synthesis of 3ak



3ak, colorless oil, (136.5 mg, 88% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.24 (d, $J = 8.0$ Hz, 1H), 7.90 (d, $J = 8.0$ Hz, 1H), 7.64 (s, 1H), 7.40-7.58 (m, 5H), 7.02 (dd, $J = 8.0$ Hz, 5.2 Hz, 1H), 6.52-6.58 (m, 1H), 6.15 (bs, 1H), 5.72 (s, 1H), 5.16 (hept, $J = 6.0$ Hz, 1H), 1.25 (d, $J = 6.0$ Hz, 3H), 1.24 (d, $J = 6.0$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 172.62, 151.61, 139.25, 132.53, 132.34, 130.88 (q, $J = 32$ Hz, 1C), 129.02, 127.06,

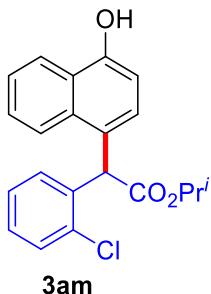
126.47, 125.84, 125.70 (q, $J = 4$ Hz, 1C), 125.05, 125.01, 124.19 (q, $J = 3$ Hz, 1C), 124.04 (q, $J = 271$ Hz, 1C), 122.80, 122.75, 107.93, 69.42, 53.20, 21.59, 21.56. MS(EI): m/z (%): 388 (M^+ , 20.94), 301 (100); HRMS (EI) calcd. for $C_{22}H_{19}O_3F_3$: 388.1286, found: 388.1280.

14) The synthesis of 3al



3al, white solid, (122.7 mg, 91% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.20-8.25 (m, 1H), 7.85-7.90 (m, 1H), 7.45-7.52 (m, 2H), 7.20-7.28 (m, 1H), 6.95-7.15 (m, 4H), 6.60-6.68 (m, 1H), 5.90-6.10 (m, 1H), 5.93 (s, 1H), 5.16 (m, 1H), 1.26 (d, $J = 6.0$ Hz, 3H), 1.24 (d, $J = 6.0$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 21.53, 21.62, 46.30 (d, $J = 3.0$ Hz, 1C), 69.28, 107.94, 115.16 (d, $J = 22$ Hz, 1C), 122.81 (d, $J = 32$ Hz, 1C), 124.06, 124.09, 124.95, 125.00 (d, $J = 8$ Hz, 1C), 125.75 (d, $J = 14$ Hz, 1C), 126.16, 126.93, 128.98 (d, $J = 8$ Hz, 1C), 130.29, 130.32, 132.62, 151.66, 160.54 (d, $J = 45$ Hz, 1C), 172.76. MS(EI): m/z (%): 338 (M^+ , 23.17), 251 (100); HRMS (EI) calcd. for $C_{21}H_{19}O_3F$: 338.1318, found: 338.1323.

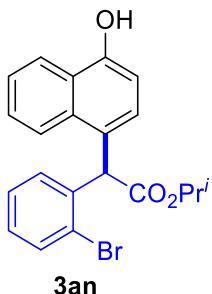
15) The synthesis of 3am



3am, white solid, (105.6 mg, 75% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.20-8.25 (m, 1H), 7.70-7.80 (m, 1H), 7.40-7.50 (m, 3H), 7.19 (dd $J = 7.6$ Hz, 1.6 Hz, 1H), 7.07-

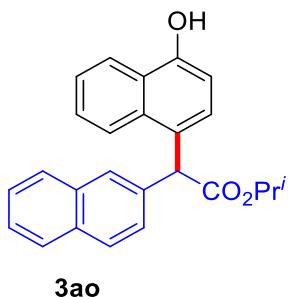
7.13 (m, 2H), 6.98-7.03 (m, 1H) 6.65-6.75 (m, 1H), 6.03 (s, 1H), 5.60-5.90 (m, 1H), 5.18 (hept, $J = 6.4$ Hz, 1H), 1.28 (d, $J = 6.4$ Hz, 3H), 1.24 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.56, 21.69, 50.80, 69.27, 107.91, 122.61, 123.14, 125.01, 125.13, 125.18, 126.12, 126.87, 126.99, 128.58, 129.42, 130.32, 132.62, 134.20, 136.43, 151.66, 172.75. MS(EI): m/z (%): 354 (M^+ , 26.77), 356 ($[\text{M}+2]^+$, 8.84), 267 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{19}\text{O}_3\text{Cl}$: 354.1023, found: 354.1023.

16) The synthesis of 3an



3an, white solid, (115.0 mg, 72% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.17-8.23 (m, 1H), 7.68-7.74 (m, 1H), 7.60 (dd, $J = 7.6$ Hz, 1.6 Hz, 1H), 7.40-7.46 (m, 2H), 7.00-7.15 (m, 4H), 6.64 (d, $J = 8.0$ Hz, 1H), 6.24 (s, 1H), 6.02 (s, 1H), 5.20 (hept, $J = 6.4$ Hz, 1H), 1.30 (d, $J = 6.4$ Hz, 3H), 1.25 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.57, 21.73, 53.49, 69.21, 107.88, 122.60, 123.23, 125.01, 125.05, 125.14, 125.45, 126.10, 127.00, 127.49, 128.83, 130.53, 132.62, 132.78, 138.15, 151.67, 172.56. MS(EI): m/z (%): 398 (M^+ , 26.76), 400 ($[\text{M}+2]^+$, 27.02), 231 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{19}\text{O}_3\text{Br}$: 398.0518, found: 398.0521.

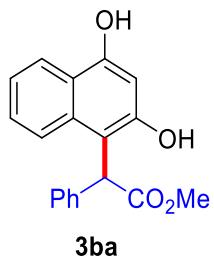
17) The synthesis of 3ao



3ao, white solid, (146.2 mg, 99% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.20-8.25 (m,

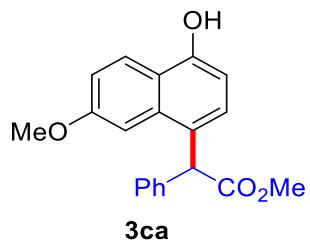
1H), 8.00 (d, J = 8.4 Hz, 1H), 7.75-7.85 (m, 4H), 7.40-7.50 (m, 5H), 7.00-7.05 (m, 1H), 6.45-6.55 (m, 1H), 6.03 (bs, 1H), 5.83 (s, 1H), 5.19 (hept, J = 6.4 Hz, 1H), 1.29 (d, J = 6.4 Hz, 3H), 1.25 (d, J = 6.4 Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.64, 21.65, 53.61, 69.17, 108.03, 122.70, 122.90, 124.78, 124.99, 125.92, 126.10, 126.38, 126.79, 126.82, 127.19, 127.55, 127.58, 127.93, 128.25, 132.55, 132.69, 133.38, 135.60, 151.46, 173.43. MS(EI): m/z (%): 370 (M^+ , 23.62), 283 (100); HRMS (EI) calcd. for $\text{C}_{25}\text{H}_{22}\text{O}_3$: 370.1569, found: 370.1571

18) The synthesis of 3ba



3ba, colorless oil, (71.5 mg, 58% yield). ^1H -NMR (400 MHz, CD_3OD) δ 8.11 (dd, J = 8.4 Hz, 0.8 Hz, 1H), 7.67 (d, J = 8.8 Hz, 1H), 7.26-7.33 (m, 3H), 7.11-7.23 (m, 4H), 6.63 (s, 1H), 5.65 (s, 1H), 3.65 (s, 3H), 2.12 (s, 2H); ^{13}C -NMR (100 MHz, CDCl_3) δ 30.66, 52.54, 101.28, 110.19, 122.23, 122.31, 123.59, 123.74, 127.50, 127.78, 128.90, 130.38, 135.31, 140.27, 154.26, 155.25, 177.27. MS(EI): m/z (%): 308 (M^+ , 3.12), 247 (100); HRMS (EI) calcd. for $\text{C}_{19}\text{H}_{16}\text{O}_4$: 308.1049, found: 308.1047.

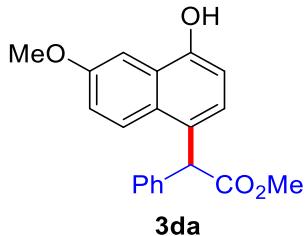
19) The synthesis of 3ca



3ca, colorless oil, (120.3 mg, 93% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.14 (d, J = 9.2 Hz, 1H), 7.25-7.38 (m, 5H), 7.17 (d, J = 2.4 Hz, 1H), 7.12 (dd, J = 9.2 Hz, 2.4 Hz, 1H), 7.03 (d, J = 8.0 Hz, 1H), 6.48 (d, J = 7.6 Hz, 1H), 5.93 (bs, 1H), 5.60 (s, 1H), 3.83 (s, 3H), 3.79 (s, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 52.49, 53.74, 55.13, 102.35,

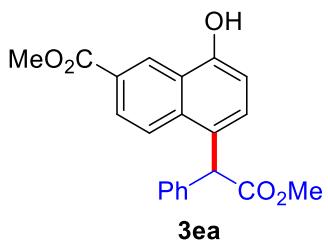
106.03, 116.95, 120.03, 124.38, 125.49, 127.21, 127.34, 128.63, 128.94, 134.06, 137.89, 151.55, 158.41, 173.89. MS(EI): m/z (%): 322 (M^+ , 6.61), 195 (100); HRMS (EI) calcd. for $C_{20}H_{18}O_4$: 322.1205, found: 322.1207.

20) The synthesis of 3da



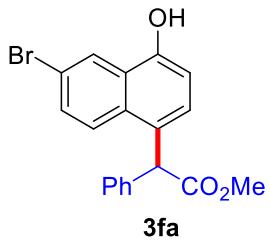
3da, colorless oil, (124.6 mg, 97% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 7.82 (d, J = 9.2 Hz, 1H), 7.50 (d, J = 2.8 Hz, 1H), 7.25-7.38 (m, 5H), 7.16 (dd, J = 9.2 Hz, 2.4 Hz, 1H), 6.91 (d, J = 8.0 Hz, 1H), 6.54 (d, J = 8.0 Hz, 1H), 6.32 (s, 1H), 5.70 (s, 1H), 3.90 (s, 3H), 3.80 (s, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 52.61, 53.31, 55.28, 101.06, 108.51, 119.42, 124.07, 124.61, 126.08, 126.33, 127.30, 127.99, 128.61, 128.88, 138.00, 150.52, 156.87, 174.30. MS(EI): m/z (%): 322 (M^+ , 39.31), 84 (100); HRMS (EI) calcd. for $C_{20}H_{18}O_4$: 322.1205, found: 322.1210.

21) The synthesis of 3ea



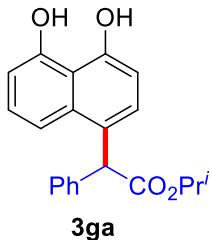
3ea, colorless oil, (112.4 mg, 80% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.98 (s, 1H), 8.06 (d, J = 8.8 Hz, 1H), 7.92 (d, J = 8.8 Hz, 1H), 7.25-7.40 (m, 5H), 7.13 (d, J = 8.0 Hz, 1H), 6.97 (s, 1H), 6.66 (d, J = 8.0 Hz, 1H), 5.71 (s, 1H), 3.97 (s, 3H), 3.80 (s, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 52.29, 52.72, 53.11, 108.80, 123.13, 124.29, 125.99, 126.10, 126.27, 127.48, 128.75, 128.87, 129.41, 134.66, 137.58, 152.90, 167.48, 174.09. MS(EI): m/z (%): 350 (M^+ , 32.26), 43 (100); HRMS (EI) calcd. for $C_{21}H_{18}O_5$: 350.1154, found: 350.1156.

22) The synthesis of 3fa



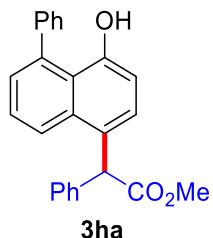
3fa, colorless oil, (130.1 mg, 88% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.34 (d, $J = 2.0$ Hz, 1H), 7.74 (d, $J = 8.8$ Hz, 1H), 7.55 (dd, $J = 8.8$ Hz, 2.0 Hz, 1H), 7.28-7.38 (m, 5H), 6.99 (d, $J = 8.0$ Hz, 1H), 6.50 (d, $J = 8.0$ Hz, 1H), 6.14 (s, 1H), 5.65 (s, 1H), 3.80 (s, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 52.78, 53.18, 108.96, 119.22, 124.67, 125.27, 126.14, 126.43, 127.04, 127.52, 128.76, 128.85, 130.24, 131.02, 137.49, 150.57, 174.15. MS(EI): m/z (%): 370 (M^+ , 34.87), 372 ($[\text{M}+2]^+$, 35.18), 311 (100); HRMS (EI) calcd. for $\text{C}_{19}\text{H}_{15}\text{O}_3\text{Br}$: 370.0205, found: 370.0202.

23) The synthesis of 3ga



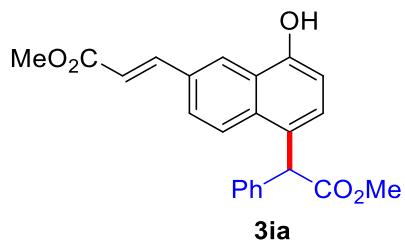
3ga, colorless oil, (70.2 mg, 52% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 7.20-7.40 (m, 8H), 6.81 (d, $J = 8.0$ Hz, 1H), 6.65 (d, $J = 7.6$ Hz, 1H), 6.41 (d, $J = 8.0$ Hz, 1H), 5.56 (s, 1H), 5.17 (hept, $J = 6.0$ Hz, 1H), 1.31 (d, $J = 6.0$ Hz, 3H), 1.24 (d, $J = 6.0$ Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 21.52, 21.67, 54.15, 69.61, 108.21, 109.53, 114.70, 114.94, 125.83, 127.39, 127.42, 127.64, 128.72, 128.96, 134.74, 137.53, 153.16, 154.26, 174.60. MS(EI): m/z (%): 336 (M^+ , 1.62), 84 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{20}\text{O}_4$: 336.1362, found: 336.1369.

24) The synthesis of 3ha



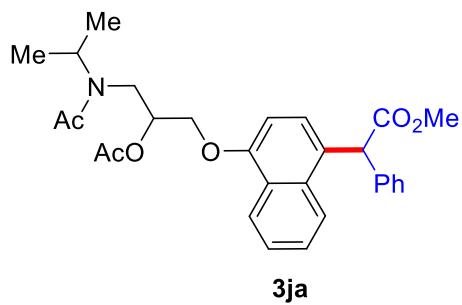
3ha, colorless oil, (130.2 mg, 88% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.03 (d, $J = 8.4$ Hz, 1H), 7.45-7.56 (m, 6H), 7.25-7.40 (m, 6H), 7.23 (dd, $J = 7.2$ Hz, 1.2 Hz, 1H), 6.88 (d, $J = 8.0$ Hz, 1H), 5.78 (s, 1H), 5.49 (s, 1H), 3.80 (s, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 52.46, 53.77, 111.19, 121.82, 123.68, 125.47, 126.69, 127.35, 127.72, 128.50, 128.60, 128.65, 128.94, 129.35, 129.40, 133.45, 137.02, 138.12, 141.23, 152.83, 173.56. MS(EI): m/z (%): 368 (M^+ , 38.32), 309 (100); HRMS (EI) calcd. for $\text{C}_{25}\text{H}_{20}\text{O}_3$: 368.1412, found: 368.1411.

25) The synthesis of 3ia



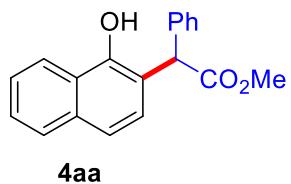
3ia, yellow solid, (136.6 mg, 91% yield). $^1\text{H-NMR}$ (400 MHz, Acetone- d_6) δ 9.40 (s, 1H), 8.48 (s, 1H), 8.03 (d, $J = 8.8$ Hz, 1H), 7.70-7.80 (m, 2H), 7.25-7.40 (m, 6H), 6.95 (d, $J = 7.6$ Hz, 1H), 6.64 (d, $J = 16.0$ Hz, 1H), 5.79 (s, 1H), 3.77 (s, 3H), 3.72 (s, 3H); $^{13}\text{C-NMR}$ (100 MHz, Acetone- d_6) δ 51.76, 52.52, 53.55, 109.28, 118.73, 125.15, 126.17, 127.02, 127.98, 129.04, 129.30, 129.84, 131.57, 134.33, 139.68, 145.31, 154.37, 167.57, 173.79. HRMS (ESI) calcd. for $\text{C}_{23}\text{H}_{21}\text{O}_5$, $[\text{M}+1]^+$: 377.1384, found: 377.1389.

26) The synthesis of 3ja



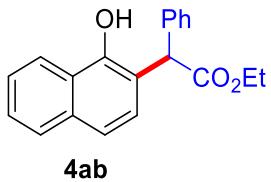
3ja, colorless oil, (184.2 mg, 94% yield, d.r. = 3:1). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.25-8.35 (m, 1H), 7.85-8.00 (m, 1H), 7.45-7.60 (m, 2H), 7.20-7.40 (m, 6H), 6.70-6.80 (m, 1H), [5.72 (s, 0.27), 5.70 (s, 0.73)], [5.60-5.65 (m, 0.73), 5.45-5.55 (s, 0.27)], [4.45-4.55 (m, 0.27), 4.00-4.10 (s, 0.73)], 4.20-4.35 (m, 2H), [3.80-3.90 (m, 0.73), 3.60-3.70 (s, 0.27)], [3.80-3.90 (m, 0.27), 3.35-3.45 (s, 0.73)], 3.75 (s, 3H), [2.24 (s, 0.81H), 2.16 (s, 2.19)], [2.11 (s, 0.81H), 2.08 (s, 2.19)], 1.20-1.35 (m, 6H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ [20.09, 20.86], [20.51, 21.00], 21.61, [21.95, 22.63], [41.33, 45.42], [47.21, 49.40], 52.27, 53.10, [66.91, 68.33], [70.79, 71.44], 104.17, [122.17, 122.58], [122.90, 123.07], [125.07, 125.36], [125.74, 125.93], [126.20, 126.33], [126.83, 126.93], [127.18, 127.27], [128.49, 128.54], 128.79, [132.38, 132.47], 138.06, [153.33, 153.88], [170.05, 170.38], 171.21, [173.24, 173.34]. MS(EI): m/z (%): 491 (M^+ , 0.25), 200 (100); HRMS (EI) calcd. for $\text{C}_{29}\text{H}_{33}\text{NO}_6$: 491.2308, found: 491.2303.

27) The synthesis of 4aa



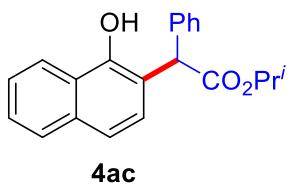
4aa, colorless oil, (95.6 mg, 82% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.77 (s, 1H), 8.25-8.35 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.15-7.30 (m, 6H), 5.21 (s, 1H), 3.84 (s, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 53.26, 55.86, 116.04, 120.28, 122.55, 125.39, 126.32, 126.61, 127.23, 127.52, 128.74, 129.03, 134.42, 136.62, 151.35, 176.55. MS(EI): m/z (%): 292 (M^+ , 15.46), 231 (100); HRMS (EI) calcd. for $\text{C}_{19}\text{H}_{16}\text{O}_3$: 292.1099, found: 292.1108.

28) The synthesis of 4ab



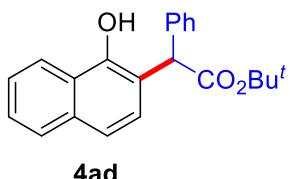
4ab, colorless oil, (105.8 mg, 87% yield). ¹H-NMR (400 MHz, CDCl₃) δ 9.04 (s, 1H), 8.25-8.35 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.15-7.30 (m, 6H), 5.17 (s, 1H), 4.20-4.40 (m, 2H), 1.33 (t, *J* = 7.2 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 14.04, 56.29, 62.59, 115.96, 120.16, 122.69, 125.33, 126.41, 126.59, 127.18, 127.44, 127.47, 128.71, 129.20, 134.44, 136.75, 151.55, 176.25. MS(EI): m/z (%): 306 (M⁺, 14.98), 231 (100); HRMS (EI) calcd. for C₂₀H₁₈O₃: 306.1256, found: 306.1262.

29) The synthesis of 4ac



4ac, colorless oil, (117.2 mg, 91% yield). ¹H-NMR (400 MHz, CDCl₃) δ 9.26 (s, 1H), 8.30-8.40 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.15-7.30 (m, 6H), 5.15 (hept, *J* = 6.4 Hz, 1H), 5.12 (s, 1H), 1.35 (d, *J* = 6.4 Hz, 3H), 1.27 (d, *J* = 6.4 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.59, 21.69, 56.68, 70.60, 115.89, 120.07, 122.78, 125.28, 126.48, 126.57, 127.15, 127.36, 127.42, 128.69, 129.34, 134.45, 136.87, 151.69, 175.88. MS(EI): m/z (%): 320 (M⁺, 15.59), 231 (100); HRMS (EI) calcd. for C₂₁H₂₀O₃: 320.1412, found: 320.1415.

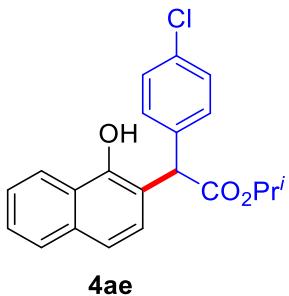
30) The synthesis of 4ad



4ad, colorless oil, (120.7 mg, 90% yield). ¹H-NMR (400 MHz, CDCl₃) δ 9.43 (s, 1H), 8.30-8.40 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.15-7.30 (m, 6H), 5.05 (s,

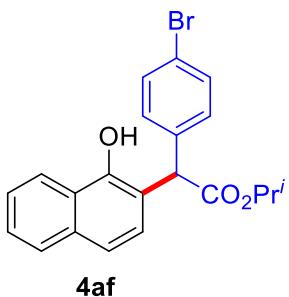
1H), 1.52 (s, 9H); ^{13}C -NMR (100 MHz, CDCl_3) δ 27.94, 57.61, 83.94, 116.12, 119.92, 122.84, 125.20, 126.51, 127.11, 127.30, 128.63, 129.48, 134.41, 137.17, 151.78, 175.85. MS(EI): m/z (%): 334 (M^+ , 6.23), 231 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{22}\text{O}_3$: 334.1569, found: 334.1576.

31) The synthesis of 4ae



4ae, light yellow oil, (125.2 mg, 88% yield). ^1H -NMR (400 MHz, CDCl_3) δ 9.18 (s, 1H), 8.30-8.40 (m, 1H), 7.80-7.85 (m, 1H), 7.40-7.55 (m, 3H), 7.15-7.30 (m, 5H), 5.21 (hept, $J = 6.4$ Hz, 1H), 5.13 (s, 1H), 1.39 (d, $J = 6.4$ Hz, 3H), 1.31 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.53, 21.66, 55.89, 70.82, 115.66, 120.30, 122.65, 125.42, 126.41, 126.70, 127.20, 128.80, 128.82, 129.00, 133.33, 134.47, 135.40, 151.53, 175.49. MS(EI): m/z (%): 354 (M^+ , 14.58), 356 ($[\text{M}+2]^+$, 5.02), 231 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{19}\text{O}_3\text{Cl}$: 354.1023, found: 354.1018.

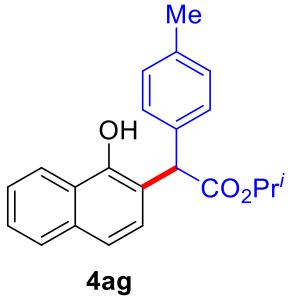
32) The synthesis of 4af



4af, colorless oil, (130.0 mg, 82% yield). ^1H -NMR (400 MHz, CDCl_3) δ 9.15 (s, 1H), 8.32-8.38 (m, 1H), 7.78-7.84 (m, 1H), 7.48-7.55 (m, 2H), 7.40-7.46 (m, 3H), 7.23 (d, $J = 8.4$ Hz, 1H), 7.15 (d, $J = 8.4$ Hz, 2H), 5.20 (hept, $J = 6.4$ Hz, 1H), 5.10 (s, 1H), 1.39 (d, $J = 6.4$ Hz, 3H), 1.30 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.54,

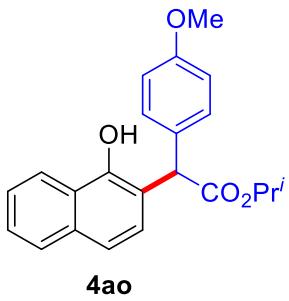
21.67, 55.96, 70.85, 115.58, 120.31, 121.48, 122.65, 125.43, 126.41, 126.72, 127.20, 129.00, 129.17, 131.76, 134.48, 135.94, 151.53, 175.42. MS(EI): m/z (%): 398 (M^+ , 7.77), 400 ($[M+2]^+$, 7.90), 231 (100); HRMS (EI) calcd. for $C_{21}H_{19}O_3Br$: 398.0518, found: 398.0532.

33) The synthesis of 4ag



4ag, colorless oil, (117.7 mg, 88% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 9.25 (s, 1H), 8.25-8.35 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.15-7.30 (m, 5H), 5.15 (hept, $J = 6.4$ Hz, 1H), 5.08 (s, 1H), 2.28 (s, 3H), 1.34 (d, $J = 6.4$ Hz, 3H), 1.27 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 20.97, 21.58, 21.68, 56.34, 70.51, 116.07, 120.00, 122.79, 125.22, 126.47, 126.51, 127.12, 127.21, 129.33, 129.38, 133.84, 134.40, 137.09, 151.64, 176.03. MS(EI): m/z (%): 334 (M^+ , 16.49), 231 (100); HRMS (EI) calcd. for $C_{22}H_{22}O_3$: 334.1569, found: 334.1563.

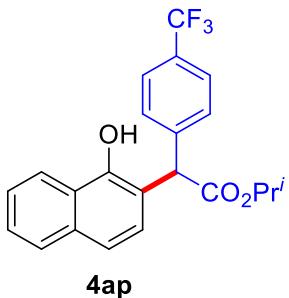
34) The synthesis of 4ao



4ao, yellow oil, (73.8 mg, 53% yield). 1H -NMR (500 MHz, $CDCl_3$) δ 9.31 (s, 1H), 8.32-8.36 (m, 1H), 7.75-7.82 (m, 1H), 7.45-7.52 (m, 2H), 7.40 (d, $J = 8.5$ Hz, 1H), 7.21 (d, $J = 8.5$ Hz, 1H), 7.16 (d, $J = 8.5$ Hz, 2H), 6.82 (d, $J = 8.5$ Hz, 2H), 5.17 (hept, $J = 6.5$ Hz, 1H), 5.06 (s, 1H), 3.76 (s, 3H), 1.36 (d, $J = 6.5$ Hz, 3H), 1.28 (d, $J = 6.5$ Hz,

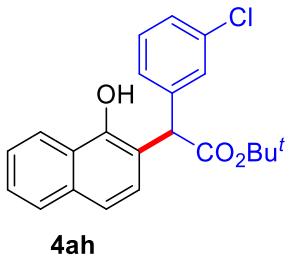
3H); ^{13}C -NMR (125 MHz, CDCl_3) δ 21.61, 21.72, 55.22, 55.99, 70.56, 114.05, 116.06, 120.02, 122.79, 125.26, 126.44, 126.55, 127.13, 128.47, 128.80, 129.32, 134.38, 151.62, 158.79, 176.17. HRMS (ESI) calcd. for $[\text{C}_{22}\text{H}_{22}\text{O}_4 + \text{Na}]^+$: 373.1410, found: 373.1416.

35) The synthesis of 4ap



4ap, yellow oil, (80.2 mg, 52% yield). ^1H -NMR (500 MHz, CDCl_3) δ 9.17 (s, 1H), 8.45-8.52 (m, 1H), 7.30-7.35 (m, 1H), 7.58 (d, $J = 8.5$ Hz, 2H), 7.50-7.58 (m, 2H), 7.49 (d, $J = 8.5$ Hz, 1H), 7.40 (d, $J = 8.5$ Hz, 2H), 7.26 (d, $J = 8.5$ Hz, 1H), 5.23 (hept, $J = 6.0$ Hz, 1H), 5.21 (s, 1H), 1.41 (d, $J = 6.0$ Hz, 3H), 1.32 (d, $J = 6.0$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.52, 21.65, 56.21, 71.06, 115.41, 120.47, 122.60, 123.98 (q, $J = 270$ Hz, 1C), 125.53, 125.61 (q, $J = 3.8$ Hz, 1C), 126.40, 126.82, 127.26, 127.83, 128.95, 129.65 (q, $J = 32.5$ Hz, 1C), 134.53, 140.89, 151.56, 175.30; HRMS (ESI) calcd. for $[\text{C}_{22}\text{H}_{19}\text{F}_3\text{O}_3+\text{Na}]^+$: 411.1178, found: 411.1189.

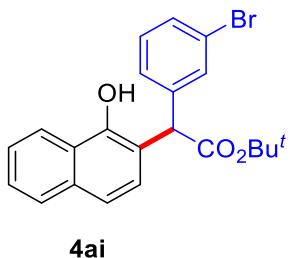
36) The synthesis of 4ah



4ah, colorless oil, (116.9 mg, 79% yield). ^1H -NMR (400 MHz, CDCl_3) δ 9.26 (s, 1H), 8.25-8.35 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.10-7.25 (m, 5H), 5.01 (s, 1H), 1.52 (s, 9H); ^{13}C -NMR (100 MHz, CDCl_3) δ 27.91, 57.04, 84.33, 115.66, 120.19,

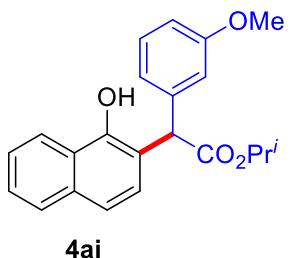
122.77, 125.35, 125.56, 126.50, 126.67, 127.18, 127.55, 127.64, 129.17, 129.83, 134.48, 134.50, 139.25, 151.68, 175.20. MS(EI): m/z (%): 368 (M^+ , 0.93), 370 ($[M+2]^+$, 1.29), 231 (100); HRMS (EI) calcd. for $C_{22}H_{21}O_3Cl$: 368.1179, found: 368.1182.

37) The synthesis of 4ai



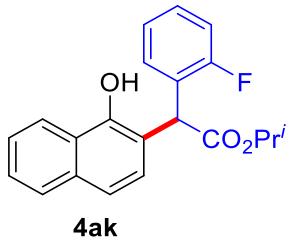
4ai, colorless oil, (133.2 mg, 81% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 9.27 (s, 1H), 8.30-8.40 (m, 1H), 7.75-7.85 (m, 1H), 7.35-7.55 (m, 5H), 7.10-7.25 (m, 3H), 5.03 (s, 1H), 1.55 (s, 9H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 27.92, 56.99, 84.35, 115.65, 120.20, 122.70, 122.77, 125.36, 126.02, 126.50, 126.68, 127.18, 129.15, 130.13, 130.49, 130.53, 134.48, 139.51, 151.66, 175.17. MS(EI): m/z (%): 412 (M^+ , 2.00), 414 ($[M+2]^+$, 1.84), 231 (100); HRMS (EI) calcd. for $C_{22}H_{21}O_3Br$: 412.0674, found: 412.0668.

38) The synthesis of 4aj



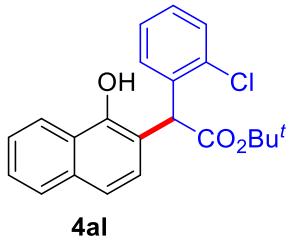
4aj, colorless oil, (105.6 mg, 75% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 9.23 (s, 1H), 8.30-8.35 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.15-7.25 (m, 2H), 6.70-6.85 (m, 3H), 5.16 (hept, $J = 6.4$ Hz, 1H), 5.09 (s, 1H), 3.70 (s, 3H), 1.35 (d, $J = 6.4$ Hz, 3H), 1.27 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 21.56, 21.67, 55.09, 56.51, 70.58, 112.44, 113.63, 115.88, 119.71, 120.06, 122.74, 125.25, 126.43, 126.54, 127.13, 129.26, 129.64, 134.43, 138.40, 151.63, 159.77, 175.67. MS(EI): m/z (%): 350 (M^+ , 29.71), 231 (100); HRMS (EI) calcd. for $C_{22}H_{22}O_4$: 350.1518, found: 350.1526.

39) The synthesis of 4ak



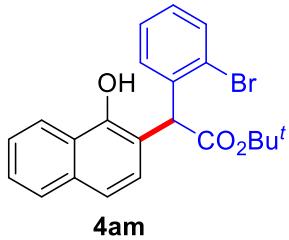
4ak, colorless oil, (103.3 mg, 76% yield). ¹H-NMR (400 MHz, CDCl₃) δ 8.75 (s, 1H), 8.25-8.35 (m, 1H), 7.70-7.80 (m, 1H), 7.35-7.50 (m, 3H), 7.20-7.30 (m, 2H), 6.95-7.15 (m, 3H), 5.24 (s, 1H), 5.13 (hept, *J* = 6.4 Hz, 1H), 1.33 (d, *J* = 6.4 Hz, 3H), 1.21 (d, *J* = 6.4 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.37, 21.47, 50.82, 70.62, 113.79, [115.36, *J* = 22 Hz, 1C], 120.43, 122.63, [123.65, *J* = 15 Hz, 1C], [124.22, *J* = 4 Hz, 1C], 125.40, 126.20, 126.71, 127.25, [129.32, *J* = 25 Hz, 1C], 129.37, 134.53, 151.85, 159.46, 161.91, 174.94.

40) The synthesis of 4al



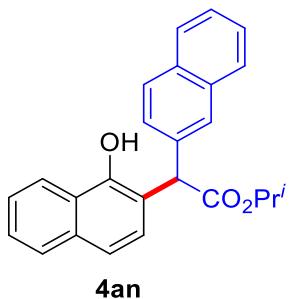
4al, colorless oil, (110.1 mg, 75% yield). ¹H-NMR (400 MHz, CDCl₃) δ 8.25-8.35 (m, 1H), 8.26 (s, 1H), 7.75-7.85 (m, 1H), 7.40-7.50 (m, 4H), 7.10-7.30 (m, 4H), 5.34 (s, 1H), 1.46 (s, 9H); ¹³C-NMR (100 MHz, CDCl₃) δ 27.76, 54.98, 83.61, 114.63, 120.44, 122.52, 125.37, 126.03, 126.64, 127.00, 127.29, 128.89, 129.17, 129.81, 130.00, 134.18, 134.31, 134.39, 151.58, 174.03. MS(EI): m/z (%): 368 (M⁺, 2.63), 370 ([M+2]⁺, 0.96), 231 (100); HRMS (EI) calcd. for C₂₂H₂₁O₃Cl: 368.1179, found: 368.1187.

41) The synthesis of 4am



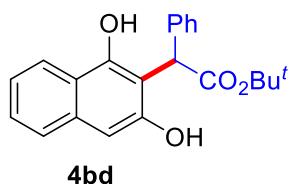
4am, yellow oil, (113.3 mg, 69% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.25-8.35 (m, 1H), 8.07 (s, 1H), 7.75-7.85 (m, 1H), 7.60 (d, $J = 8.0$ Hz, 1H), 7.45-7.55 (m, 2H), 7.41 (d, $J = 8.4$ Hz, 1H), 7.10-7.30 (m, 4H), 5.39 (s, 1H), 1.47 (s, 9H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 27.80, 56.85, 83.58, 115.26, 120.47, 122.42, 124.78, 125.40, 125.97, 126.62, 127.33, 127.66, 128.96, 129.12, 130.32, 133.22, 134.34, 136.22, 151.28, 173.69. MS(EI): m/z (%): 412 (M^+ , 1.49), 414 ($[\text{M}+2]^+$, 1.50), 231 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{21}\text{O}_3\text{Br}$: 412.0674, found: 412.0677.

42) The synthesis of 4an



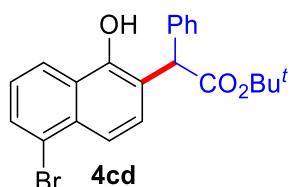
4an, colorless oil, (123.5 mg, 83% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 9.25 (s, 1H), 7.35-7.45 (m, 1H), 7.70-7.90 (m, 5H), 7.45-7.55 (m, 5H), 7.41 (d, $J = 8.4$ Hz, 1H), 7.34 (d, $J = 8.4$ Hz, 1H), 5.35 (s, 1H), 5.26 (hept, $J = 6.4$ Hz, 1H), 1.43 (d, $J = 6.4$ Hz, 3H), 1.35 (d, $J = 6.4$ Hz, 3H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 21.59, 21.72, 56.66, 70.64, 115.96, 120.14, 122.77, 125.30, 125.56, 126.07, 126.11, 126.21, 126.46, 126.60, 127.18, 127.49, 128.06, 128.48, 129.29, 132.55, 133.26, 134.22, 134.46, 151.71, 175.65. MS(EI): m/z (%): 370 (M^+ , 11.25), 281 (100); HRMS (EI) calcd. for $\text{C}_{25}\text{H}_{22}\text{O}_3$: 370.1569, found: 370.1561.

43) The synthesis of 4bd



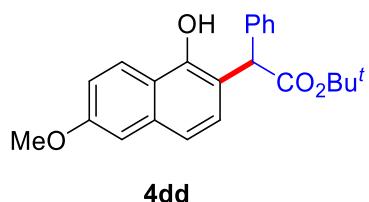
4bd, colorless oil, (104.4 mg, 75% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.98 (s, 1H), 8.17 (d, $J = 8.0$ Hz, 1H), 7.92 (d, $J = 8.8$ Hz, 1H), 7.47 (t, $J = 8.0$ Hz, 1H), 7.31 (t, $J = 7.6$ Hz, 1H), 7.15-7.25 (m, 5H), 6.52 (s, 2H), 5.80 (s, 1H), 1.50 (s, 9H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 27.94, 49.43, 83.77, 103.12, 107.63, 120.96, 121.63, 122.31, 122.71, 127.03, 127.37, 127.60, 128.48, 134.23, 137.51, 153.07, 154.33, 176.32. MS(EI): m/z (%): 350 (M^+ , 0.83), 247 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{22}\text{O}_4$: 350.1518, found: 350.1511.

44) The synthesis of 4cd



4cd, yellow oil, (137.5 mg, 83% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 9.71(s, 1H), 8.34 (d, $J = 8.4$ Hz, 1H), 7.70-7.80 (m, 2H), 7.20-7.35 (m, 7H), 5.06 (s, 1H), 1.52 (s, 9H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 27.92, 57.65, 84.28, 117.04, 119.06, 122.08, 122.93, 125.39, 127.19, 127.43, 127.91, 128.71, 130.56, 130.78, 132.95, 136.83, 151.97, 175.89. MS(EI): m/z (%): 412 (M^+ , 3.90), 414 ($[\text{M}+2]^+$, 4.03), 311 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{21}\text{O}_3\text{Br}$: 412.0674, found: 412.0668.

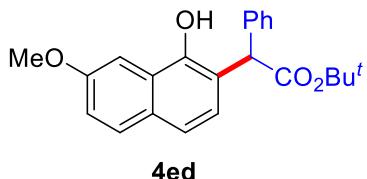
45) The synthesis of 4dd



4dd, colorless oil, (124.9 mg, 86% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 9.46 (s, 1H), 8.30 (d, $J = 9.2$ Hz, 1H), 7.10-7.40 (m, 9H), 5.07 (s, 1H), 3.94 (s, 3H), 1.57 (s, 9H);

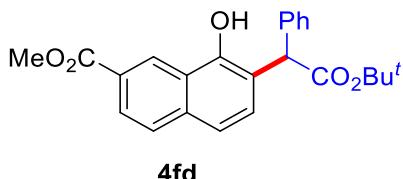
¹³C-NMR (100 MHz, CDCl₃) δ 27.93, 55.24, 57.50, 83.87, 105.26, 114.24, 117.65, 118.81, 121.76, 124.64, 127.26, 128.60, 130.26, 135.86, 137.36, 152.00, 158.31, 176.01. MS(EI): m/z (%): 364 (M⁺, 8.73), 261 (100); HRMS (EI) calcd. for C₂₃H₂₄O₄: 364.1675, found: 364.1680.

46) The synthesis of 4ed



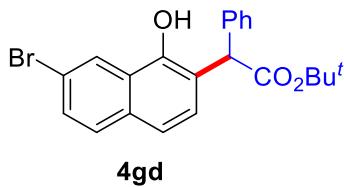
4ed, colorless oil, (126.8 mg, 86% yield). ¹H-NMR (400 MHz, CDCl₃) δ 9.46 (s, 1H), 7.60-7.70 (m, 2H), 7.05-7.40 (m, 8H), 5.03 (s, 1H), 3.91 (s, 3H), 1.53 (s, 9H); ¹³C-NMR (100 MHz, CDCl₃) δ 27.94, 55.28, 57.71, 83.96, 100.81, 116.77, 119.50, 119.75, 127.11, 127.23, 127.27, 127.52, 128.62, 128.70, 129.90, 137.22, 150.73, 157.45, 176.05. MS(EI): m/z (%): 364 (M⁺, 8.02), 261 (100); HRMS (EI) calcd. for C₂₃H₂₄O₄: 364.1675, found: 364.1671.

47) The synthesis of 4fd



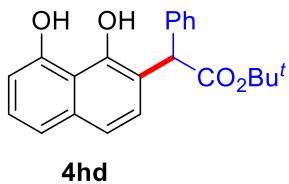
4-4fd, light yellow oil, (133.3 mg, 85% yield). ¹H-NMR (400 MHz, CDCl₃) δ 9.76 (s, 1H), 9.12 (s, 1H), 8.06 (dd, *J* = 8.4 Hz, 1.6 Hz, 1H), 7.79 (d, *J* = 8.8 Hz, 1H), 7.40 (d, *J* = 8.4 Hz, 1H), 7.20-7.35 (m, 6H), 5.07 (s, 1H), 3.94 (s, 3H), 1.53 (s, 9H); ¹³C-NMR (100 MHz, CDCl₃) δ 27.90, 52.04, 57.52, 84.21, 117.10, 119.61, 125.70, 125.94, 126.34, 126.68, 127.26, 127.37, 127.42, 128.70, 132.05, 136.41, 136.89, 152.99, 167.37, 175.72. MS(EI): m/z (%): 392 (M⁺, 3.96), 289 (100); HRMS (EI) calcd. for C₂₄H₂₄O₅: 392.1624, found: 392.1628.

48) The synthesis of 4gd



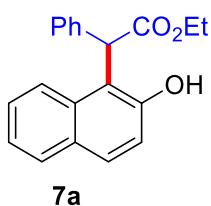
4gd, colorless oil, (138.5 mg, 84% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 9.57 (s, 1H), 8.43 (s, 1H), 7.53 (d, $J = 8.8$ Hz, 1H), 7.44 (dd, $J = 8.8$ Hz, 1.6 Hz, 1H), 7.10-7.30 (m, 7H), 4.95 (s, 1H), 1.44 (s, 9H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 27.92, 57.73, 84.27, 117.16, 119.37, 119.74, 125.48, 127.20, 127.43, 127.71, 128.71, 128.77, 129.86, 129.96, 132.78, 136.86, 151.05, 175.85. MS(EI): m/z (%): 412 (M^+ , 4.17), 414 ($[\text{M}+2]^+$, 4.08), 311 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{21}\text{O}_3\text{Br}$: 412.0674, found: 412.0666.

49) The synthesis of 4hd



4hd, colorless oil, (92.7 mg, 66% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 10.97 (s, 1H), 9.81 (s, 1H), 7.20-7.35 (m, 8H), 7.11 (d, $J = 8.4$ Hz, 1H), 6.77 (dd, $J = 6.8$ Hz, 1.6 Hz, 1H), 5.08 (s, 1H), 1.54 (s, 9H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 27.92, 56.74, 84.46, 109.87, 115.96, 116.68, 118.77, 120.72, 127.35, 127.38, 127.50, 128.66, 129.09, 136.57, 136.97, 152.06, 154.84, 176.72. MS(EI): m/z (%): 350 (M^+ , 6.39), 247 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{22}\text{O}_4$: 350.1518, found: 350.1514.

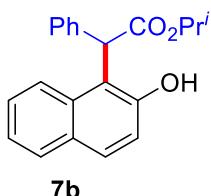
50) The synthesis of 7a



7a, colorless oil, (117.0 mg, 95% yield). $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.45 (s, 1H), 8.01 (d, $J = 8.4$ Hz, 1H), 7.80 (d, $J = 8.0$ Hz, 1H), 7.75 (d, $J = 8.8$ Hz, 1H), 7.49 (t, $J = 7.6$ Hz, 1H), 7.34 (t, $J = 7.6$ Hz, 1H), 7.10-7.30 (m, 6H), 5.99 (s, 1H), 4.36 (q, $J = 7.2$

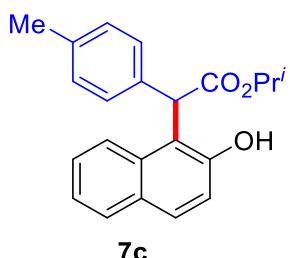
Hz, 1H), 4.23 (q, $J = 7.2$ Hz, 1H), 1.30 (t, $J = 7.2$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 14.05, 48.51, 62.49, 114.62, 120.53, 121.70, 123.20, 127.12, 127.30, 127.48, 128.58, 128.91, 129.39, 130.34, 133.30, 136.61, 154.08, 176.22. MS(EI): m/z (%): 306 (M^+ , 5.40), 231 (100); HRMS (EI) calcd. for $\text{C}_{20}\text{H}_{18}\text{O}_3$: 306.1256, found: 306.1258.

51) The synthesis of 7b



7b, colorless oil, (121.3 mg, 95% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.75 (s, 1H), 8.02 (d, $J = 8.8$ Hz, 1H), 7.81 (d, $J = 8.0$ Hz, 1H), 7.76 (d, $J = 9.2$ Hz, 1H), 7.49 (t, $J = 8.4$ Hz, 1H), 7.35 (t, $J = 7.2$ Hz, 1H), 7.10-7.30 (m, 6H), 5.96 (s, 1H), 5.17 (hept, $J = 6.4$ Hz, 1H), 1.37 (d, $J = 6.4$ Hz, 3H), 1.21 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.51, 21.77, 48.81, 70.52, 114.47, 120.69, 121.65, 123.14, 127.07, 127.26, 127.33, 128.58, 128.90, 129.36, 130.31, 133.37, 136.72, 154.29, 175.88. MS(EI): m/z (%): 320 (M^+ , 26.30), 233 (100); HRMS (EI) calcd. for $\text{C}_{21}\text{H}_{20}\text{O}_3$: 320.1412, found: 320.1416.

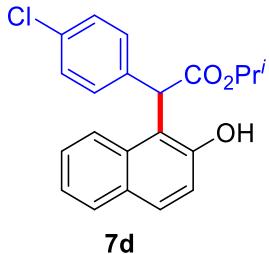
52) The synthesis of 7c



7c, colorless oil, (107.7 mg, 81% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.80 (s, 1H), 8.05 (d, $J = 8.8$ Hz, 1H), 7.84 (d, $J = 7.6$ Hz, 1H), 7.79 (d, $J = 8.8$ Hz, 1H), 7.53 (t, $J = 8.4$ Hz, 1H), 7.38 (t, $J = 7.2$ Hz, 1H), 7.22 (d, $J = 8.8$ Hz, 1H), 7.00-7.10 (m, 4H), 5.96 (s, 1H), 5.20 (hept, $J = 6.4$ Hz, 1H), 2.32 (s, 3H), 1.41 (d, $J = 6.4$ Hz, 3H), 1.24 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 20.98, 21.52, 21.78, 48.52, 70.46, 114.57,

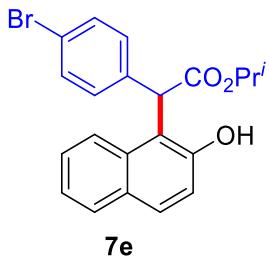
120.76, 121.66, 123.10, 127.04, 127.18, 128.89, 129.32, 129.36, 130.24, 133.39, 133.66, 136.94, 154.33, 176.03. MS(EI): m/z (%): 334 (M^+ , 11.86), 231 (100); HRMS (EI) calcd. for $C_{22}H_{22}O_3$: 334.1569, found: 334.1563.

53) The synthesis of 7d



7d, colorless oil, (140.2 mg, 99% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.49 (s, 1H), 7.96 (d, J = 8.8 Hz, 1H), 7.81 (d, J = 8.0 Hz, 1H), 7.75 (d, J = 8.8 Hz, 1H), 7.49 (t, J = 8.4 Hz, 1H), 7.35 (t, J = 7.6 Hz, 1H), 7.10-7.25 (m, 5H), 5.88 (s, 1H), 5.16 (hept, J = 6.4 Hz, 1H), 1.35 (d, J = 6.4 Hz, 3H), 1.19 (d, J = 6.4 Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 21.47, 21.75, 48.25, 70.65, 114.47, 120.45, 121.60, 123.28, 127.19, 128.68, 128.96, 129.41, 130.47, 133.10, 133.18, 135.37, 153.94, 175.42. MS(EI): m/z (%): 354 (M^+ , 6.74), 231 (100); HRMS (EI) calcd. for $C_{21}H_{19}O_3Cl$: 354.1023, found: 354.1019.

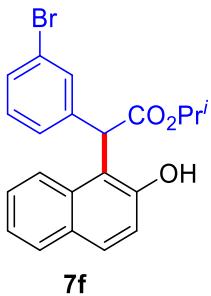
54) The synthesis of 7e



7e, colorless oil, (159.5 mg, 99% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.50 (s, 1H), 7.99 (d, J = 8.8 Hz, 1H), 7.84 (d, J = 8.0 Hz, 1H), 7.78 (d, J = 8.8 Hz, 1H), 7.50-7.55 (m, 1H), 7.35-7.40 (m, 3H), 7.18 (d, J = 8.8 Hz, 1H), 7.09 (d, J = 8.4 Hz, 2H), 5.89 (s, 1H), 5.19 (hept, J = 6.4 Hz, 1H), 1.38 (d, J = 6.4 Hz, 3H), 1.21 (d, J = 6.4 Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 21.46, 21.75, 48.32, 70.65, 114.42, 120.43, 121.23, 121.59, 123.29, 127.19, 128.96, 129.32, 129.41, 130.48, 131.62, 133.17, 135.92, 153.92, 175.31. MS(EI): m/z (%): 398 (M^+ , 5.08), 400 ($[M+2]^+$, 5.56), 231 (100); HRMS (EI)

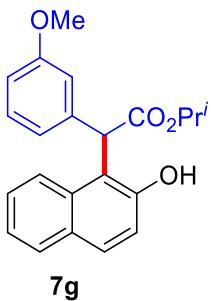
calcd. for C₂₁H₁₉O₃Br: 398.0518, found: 398.0517.

55) The synthesis of 7f



7f, colorless oil, (130.9 mg, 82% yield). ¹H-NMR (400 MHz, CDCl₃) δ 8.47 (s, 1H), 7.98 (d, *J* = 8.4 Hz, 1H), 7.84 (d, *J* = 8.0 Hz, 1H), 7.79 (d, *J* = 8.8 Hz, 1H), 7.50-7.55 (m, 1H), 7.30-7.40 (m, 3H), 7.10-7.20 (m, 3H), 5.91 (s, 1H), 5.19 (hept, *J* = 6.4 Hz, 1H), 1.38 (d, *J* = 6.4 Hz, 3H), 1.22 (d, *J* = 6.4 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.46, 21.76, 48.44, 70.70, 114.22, 120.42, 121.58, 122.65, 123.30, 126.28, 127.22, 128.97, 129.42, 130.03, 130.43, 130.55, 130.66, 133.17, 139.19, 153.93, 175.13. MS(EI): m/z (%): 398 (M⁺, 7.14), 400 ([M+2]⁺, 7.35), 231 (100); HRMS (EI) calcd. for C₂₁H₁₉O₃Br: 398.0518, found: 398.0514.

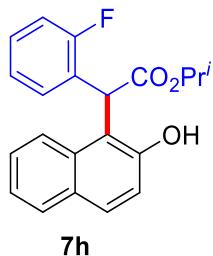
56) The synthesis of 7g



7g, colorless oil, (116.7 mg, 83% yield). ¹H-NMR (400 MHz, CDCl₃) δ 8.78 (s, 1H), 8.04 (d, *J* = 8.4 Hz, 1H), 7.83 (d, *J* = 8.0 Hz, 1H), 7.78 (d, *J* = 8.8 Hz, 1H), 7.45-7.55 (m, 1H), 7.37 (t, *J* = 7.2 Hz, 1H), 7.15-7.25 (m, 2H), 6.70-6.85 (m, 3H), 5.96 (s, 1H), 5.19 (hept, *J* = 6.4 Hz, 1H), 3.72 (s, 3H), 1.40 (d, *J* = 6.4 Hz, 3H), 1.24 (d, *J* = 6.4 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.51, 21.77, 48.81, 55.07, 70.53, 112.17, 113.71, 114.33, 119.72, 120.69, 121.62, 123.12, 127.06, 128.89, 129.37, 129.55, 130.34, 133.38,

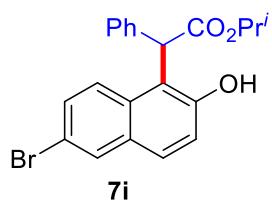
138.31, 154.36, 159.71, 175.69. MS(EI): m/z (%): 350 (M^+ , 10.68), 231 (100); HRMS (EI) calcd. for $C_{22}H_{22}O_4$: 350.1518, found: 350.1515.

57) The synthesis of 7h



7h, white solid, (127.1 mg, 94% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.15 (s, 1H), 7.96 (d, $J = 8.8$ Hz, 1H), 7.70-7.85 (m, 2H), 7.40-7.50 (m, 1H), 7.30-7.35 (m, 1H), 6.90-7.25 (m, 5H), 6.02 (s, 1H), 5.13 (hept, $J = 6.4$ Hz, 1H), 1.33 (d, $J = 6.4$ Hz, 3H), 1.15 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 21.40, 21.43, 43.63, 70.46, 112.26, [115.22, $J = 22$ Hz, 1C], 120.07, 121.85, 123.30, [124.14, $J = 4$ Hz, 1C], 127.13, 128.87, [129.31, $J = 8$ Hz, 1C], [129.55, $J = 9$ Hz, 1C], 129.63, 130.53, 133.21, 154.24, 159.93, 162.38, 174.92. MS(EI): m/z (%): 338 (M^+ , 9.61), 249 (100); HRMS (EI) calcd. for $C_{21}H_{19}O_3F$: 338.1318, found: 338.1320.

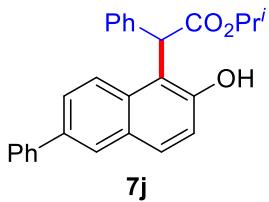
58) The synthesis of 7i



7i, colorless oil, (123.4 mg, 77% yield). 1H -NMR (400 MHz, $CDCl_3$) δ 8.74 (s, 1H), 7.94 (d, $J = 2.0$ Hz, 1H), 7.87 (d, $J = 9.2$ Hz, 1H), 7.65 (d, $J = 8.8$ Hz, 1H), 7.54 (dd, $J = 9.2$ Hz, 2 Hz, 1H), 7.10-7.30 (m, 6H), 5.88 (s, 1H), 5.17 (hept, $J = 6.4$ Hz, 1H), 1.37 (d, $J = 6.4$ Hz, 3H), 1.21 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, $CDCl_3$) δ 21.50, 21.75, 48.90, 70.69, 114.83, 116.79, 121.82, 123.65, 127.31, 127.40, 128.66, 129.36, 130.17, 130.58, 130.72, 131.97, 136.39, 154.53, 175.56. MS(EI): m/z (%): 398 (M^+ , 1.86), 400 ($[M+2]^+$, 1.91), 222 (100); HRMS (EI) calcd. for $C_{21}H_{19}O_3Br$: 398.0518,

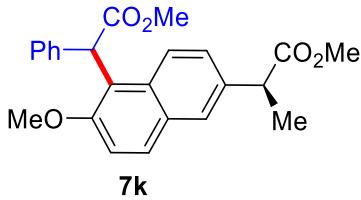
found: 398.0523.

59) The synthesis of 7j



7j, colorless oil, (157.0 mg, 99% yield). ^1H -NMR (400 MHz, CDCl_3) δ 8.76 (s, 1H), 8.00-8.10 (m, 1H), 7.81 (s, 1H), 7.60-7.80 (m, 4H), 7.10-7.50 (m, 9H), 5.98 (s, 1H), 5.18 (hept, $J = 6.4$ Hz, 1H), 1.37 (d, $J = 6.4$ Hz, 3H), 1.21 (d, $J = 6.4$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 21.51, 21.77, 48.96, 70.55, 114.52, 121.11, 122.35, 126.58, 126.69, 127.09, 127.15, 127.29, 127.40, 128.61, 128.82, 129.64, 130.59, 132.58, 135.84, 136.70, 140.63, 154.36, 175.80. MS(EI): m/z (%): 396 (M^+ , 9.76), 307 (100); HRMS (EI) calcd. for $\text{C}_{27}\text{H}_{24}\text{O}_3$: 396.1725, found: 396.1713.

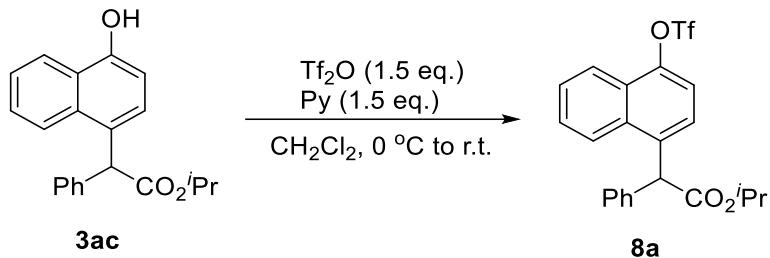
60) The synthesis of 7k



7k, colorless oil, (131.0 mg, 87% yield). ^1H -NMR (400 MHz, CDCl_3) δ 7.75-7.85 (m, 2H), 7.69 (s, 1H), 7.15-7.40 (m, 7H), 5.73 (s, 1H), 3.92 (s, 3H), 3.82 (q, $J = 7.2$ Hz, 1H), 3.69 (s, 3H), 3.63 (s, 3H), 1.54 (d, $J = 7.2$ Hz, 3H); ^{13}C -NMR (100 MHz, CDCl_3) δ 18.41, 45.06, 48.39, 51.99, 52.10, 56.54, 113.95, 120.53, 123.61, 126.86, 128.18, 129.12, 129.57, 131.84, 135.54, 135.57, 137.77, 154.45, 173.89, 174.90. MS(EI): m/z (%): 392 (M^+ , 17.56), 84 (100); HRMS (EI) calcd. for $\text{C}_{24}\text{H}_{24}\text{O}_5$: 392.1624, found: 392.1626.

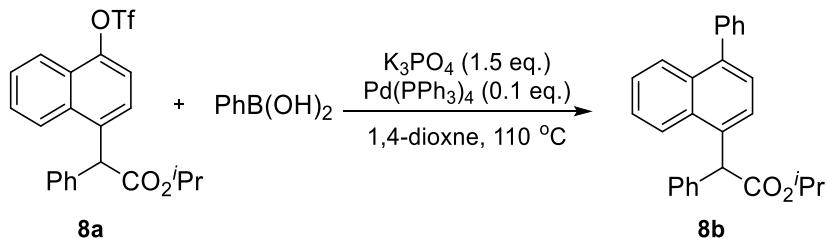
5. The transformations of products

1) the synthesis of **8a**



To a solution of **3ac** (64 mg, 0.2 mmol), pyridine (24 μ L, 0.3 mmol) in CH_2Cl_2 (4 mL) was added Tf_2O (42 μ L, 0.3 mmol) at 0 °C. The reaction was warmed to room temperature after 5 mins and stirred for another 10 mins. The reaction was extracted with ethyl acetate and dried with Na_2SO_4 and filtered. The filtrate was concentrated and purified by silica chromatography ($\text{PE}/\text{EA} = 20:1$) to obtain the desired product **8a** (90.4 mg, 99%) as a colorless oil. $^1\text{H-NMR}$ (400 MHz, CDCl_3) δ 8.05-8.15 (m, 2H), 7.60-7.70 (m, 2H), 7.20-7.45 (m, 7H), 5.71 (s, 1H), 5.12 (hept, $J = 6.4$ Hz, 1H), 1.25 (d, $J = 6.4$ Hz, 3 H), 1.21 (d, $J = 6.4$ Hz, 3 H); $^{13}\text{C-NMR}$ (100 MHz, CDCl_3) δ 21.57, 21.63, 53.71, 69.14, 117.13, 118.71 [q, $J = 18$ Hz, 1C], 121.64, 123.74, 125.77, 126.65, 127.54, 127.67, 127.87, 128.85, 128.89, 133.10, 135.80, 137.22, 145.20, 171.53. MS(EI): m/z (%): 452 (M^+ , 22.74), 365 (100); HRMS (EI) calcd. for $\text{C}_{22}\text{H}_{19}\text{O}_5\text{SF}_3$: 452.0905, found: 452.0900.

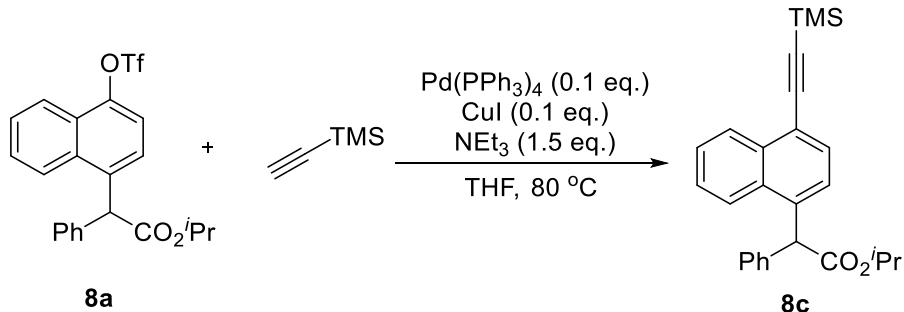
2) the synthesis of **8b**



The mixture of **8a** (90 mg, 0.2 mmol), benzeneboronic (37 mg, 0.3 mmol), K_3PO_4 (64 mg, 0.3 mmol), $\text{Pd}(\text{PPh}_3)_4$ (23 mg, 0.02 mmol) in 1,4-dioxane (4 mL) was heated to 110 °C for 8 hours. The reaction was cooled to room temperature and extracted with ethyl acetate and dried with Na_2SO_4 . The filtrate was concentrated and the residue was

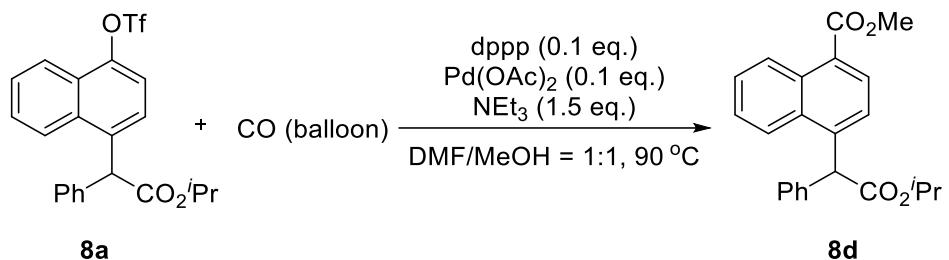
purified by silica chromatography (PE/EA = 20:1) to obtain the desired product **8b** (65.3 mg, 86%) as a colorless oil. ¹H-NMR (400 MHz, CDCl₃) δ 8.12 (d, *J* = 8.4 Hz, 1H), 7.96 (d, *J* = 8.4 Hz, 1H), 7.25-7.60 (m, 14H), 5.83 (s, 1H), 5.19 (hept, *J* = 6.4 Hz, 1H), 1.30 (d, *J* = 6.4 Hz, 3H), 1.28 (d, *J* = 6.4 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.67, 53.90, 68.79, 123.46, 125.65, 125.76, 126.20, 126.42, 127.06, 127.22, 127.31, 128.19, 128.63, 129.01, 130.10, 131.91, 132.15, 134.23, 138.07, 140.09, 140.66, 172.27. MS(EI): m/z (%): 380 (M⁺, 39.27), 293 (100); HRMS (EI) calcd. for C₂₇H₂₄O₂: 380.1776, found: 380.1774.

3) the synthesis of **8c**



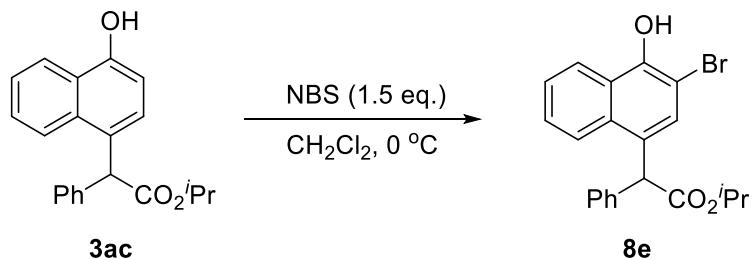
The mixture of **8a** (90 mg, 0.2 mmol), trimethylsilylacetylene (29 μL, 0.3 mmol), CuI (3.8 mg, 0.02 mmol), Pd(PPh₃)₄ (23 mg, 0.02 mmol), NEt₃ (43 μL, 0.3 mmol) in THF (4 mL) was heated to 80 °C for 12 hours. The reaction was cooled to room temperature and extracted with ethyl acetate and dried with Na₂SO₄. The filtrate was concentrated and the residue was purified by silica chromatography (PE/EA = 20:1) to obtain the desired product **8c** (73.1 mg, 91%) as a colorless oil. ¹H-NMR (400 MHz, CDCl₃) δ 8.26 (d, *J* = 8.4 Hz, 1H), 7.86 (d, *J* = 8.4 Hz, 1H), 7.51 (d, *J* = 7.2 Hz, 1H), 7.30-7.45 (m, 2H), 7.00-7.40 (m, 6H), 5.57 (s, 1H), 4.96 (hept, *J* = 6.4 Hz, 1H), 1.05 (d, *J* = 6.4 Hz, 3H), 1.04 (d, *J* = 6.4 Hz, 3H), 0.18 (s, 9H); ¹³C-NMR (100 MHz, CDCl₃) δ 0.06, 21.57, 21.62, 53.92, 68.83, 99.79, 102.98, 120.70, 123.60, 125.54, 126.52, 126.84, 127.17, 127.37, 128.63, 128.91, 130.33, 131.41, 133.77, 135.86, 137.78, 171.89. MS(EI): m/z (%): 400 (M⁺, 35.88), 313 (100); HRMS (EI) calcd. for C₂₆H₂₈O₂Si: 400.1859, found: 400.1861.

4) the synthesis of 8d



The mixture of **8a** (90 mg, 0.2 mmol), Pd(OAc)₂ (4.5 mg, 0.02 mmol), dppp (8.2 mg, 0.02 mmol), NEt₃ (43 µL, 0.3 mmol) in DMF (2 mL) and MeOH (2 mL) was heated to 90 °C under CO atmosphere (balloon) for 12 hours. The reaction was cooled to room temperature and extracted with ethyl acetate and dried with Na₂SO₄. The filtrate was concentrated and the residue was purified by silica chromatography (PE/EA = 20:1) to obtain the desired product **8d** (54.0 mg, 74%) as a colorless oil. ¹H-NMR (400 MHz, CDCl₃) δ 8.95-9.00 (m, 1H), 8.10-8.20 (m, 2H), 7.55-7.65 (m, 2H), 7.25-7.45 (m, 6H), 5.79 (s, 1H), 5.14 (hept, *J* = 6.4 Hz, 1H), 4.00 (s, 3H), 1.24 (d, *J* = 6.4 Hz, 3H), 1.21 (d, *J* = 6.4 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.57, 52.15, 52.19, 54.13, 68.99, 123.58, 125.04, 126.57, 126.68, 127.01, 127.30, 127.49, 128.70, 128.91, 129.46, 131.68, 131.97, 137.46, 139.95, 167.91, 171.69. MS(EI): m/z (%): 362 (M⁺, 37.84), 275 (100); HRMS (EI) calcd. for C₂₃H₂₂O₄: 362.1518, found: 362.1514.

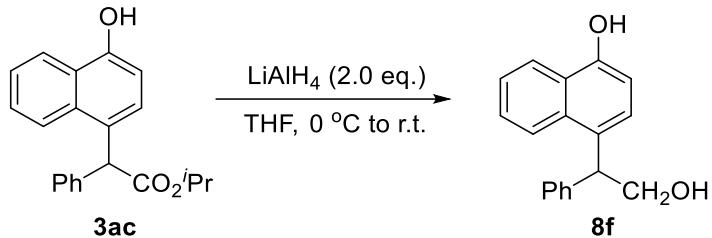
5) the synthesis of 8e



To a solution of **7a** (64 mg, 0.2 mmol) in CH₂Cl₂ (4 mL) was added NBS (53 mg, 0.3 mmol) at 0 °C. The reaction was stirred at 0 °C for 15 min and quenched with saturated Na₂S₂O₃ solution and extracted with ethyl acetate, dried with Na₂SO₄. The filtrate was concentrated and the residue was purified by silica chromatography (PE/EA = 10:1) to obtain the desired product **8e** (77.2 mg, 97%) as a white solid. ¹H-NMR (400

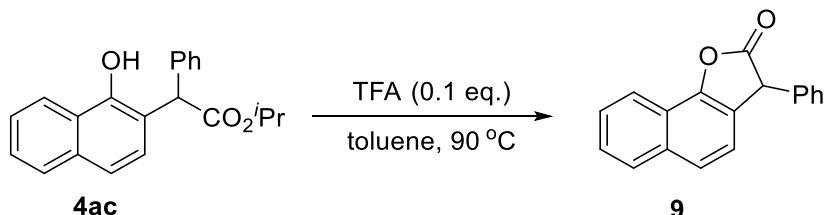
MHz, CDCl₃) δ 8.25-8.35 (m, 1H), 7.90-8.00 (m, 1H), 7.50-7.60 (m, 2H), 7.25-7.40 (m, 6H), 6.00 (s, 1H), 5.63 (s, 1H), 5.14 (hept, *J* = 6.4 Hz, 1H), 1.26 (d, *J* = 6.4 Hz, 3H), 1.23 (d, *J* = 6.4 Hz, 3H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.60, 21.62, 53.16, 68.90, 103.55, 123.12, 123.34, 124.80, 125.94, 127.22, 127.46, 128.49, 128.72, 128.83, 131.90, 137.60, 147.88, 171.99. MS(EI): m/z (%): 398 (M⁺, 28.52), 400 ([M+2]⁺, 28.84), 311 (100); HRMS (EI) calcd. for C₂₁H₁₉O₃Br: 398.0518, found: 398.0522.

6) the synthesis of 8f



To a solution of **8f** (64 mg, 0.2 mmol) in THF (4 mL) was added LiAlH₄ (15 mg, 0.4 mmol) at 0 °C. The reaction was warmed to room temperature after 15 min and stirred for another 1 hour. The reaction was quenched with saturated NaHCO₃ solution and extracted with ethyl acetate, dried with Na₂SO₄. The filtrate was concentrated and the residue was purified by silica chromatography (PE/EA = 2:1) to obtain the desired product **8f** (51.3 mg, 97%) as a white solid. ¹H-NMR (400 MHz, Acetone-*d*₆) δ 8.94 (s, 1H), 8.25-8.35 (m, 1H), 8.05-8.10 (m, 1H), 7.10-7.45 (m, 8H), 6.93 (d, *J* = 8.0 Hz, 1H), 4.88 (t, *J* = 7.2 Hz, 1H), 4.10-4.30 (m, 2H), 3.75-3.85 (m, 1H); ¹³C-NMR (100 MHz, Acetone-*d*₆) δ 49.65, 66.54, 108.30, 123.55, 124.52, 124.96, 126.05, 126.46, 126.90, 129.00, 129.41, 129.69, 134.26, 144.40, 152.84. MS(EI): m/z (%): 264 (M⁺, 17.79), 233 (100); HRMS (EI) calcd. for C₁₈H₁₆O₂: 264.1150, found: 264.1159.

7) the synthesis of 9



The solution of **4ac** (64 mg, 0.2 mmol) and TFA (2 μL, 0.02 mmol) in toluene (4 mL)

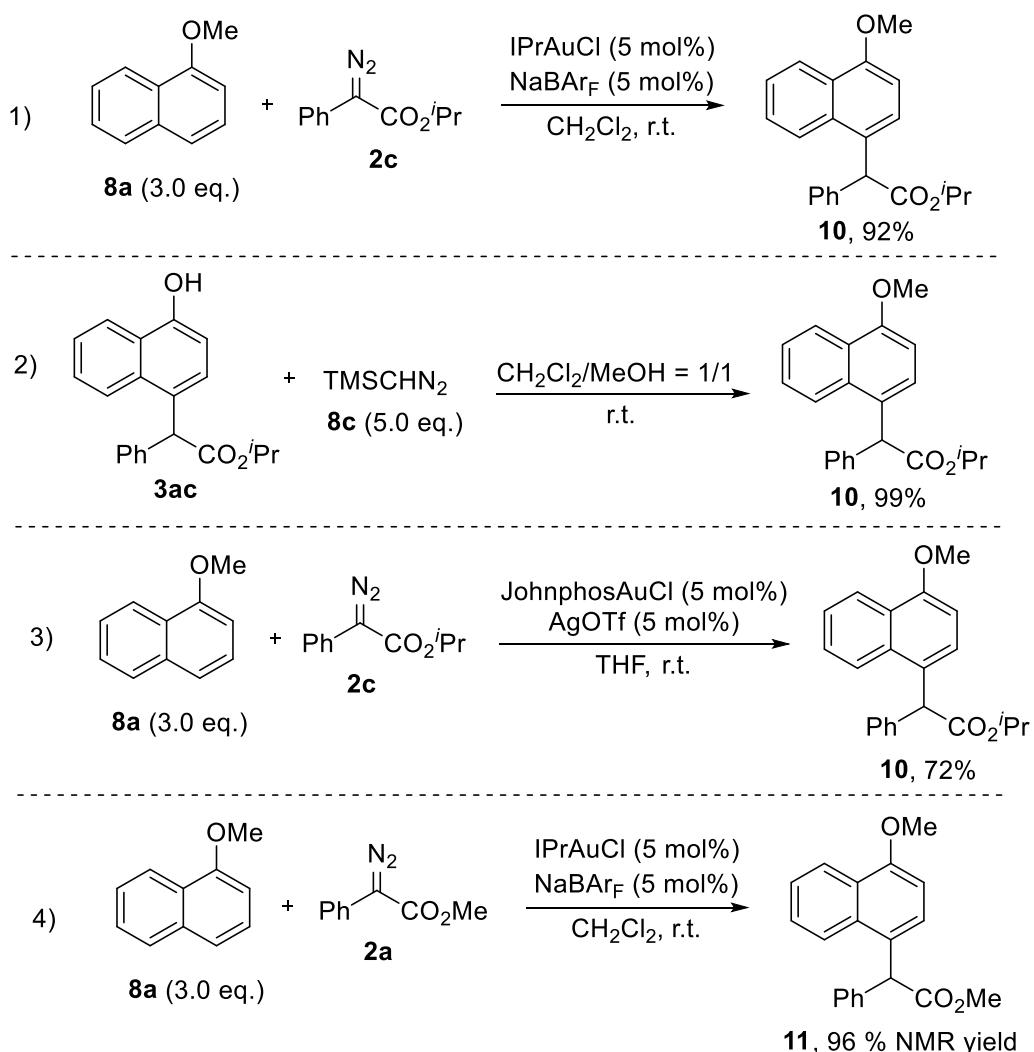
was heated to 90 °C for 12 h. Then the reaction was cooled to room temperature and purified by silica chromatography (PE/EA = 20:1) directly to obtain the desired product **9** (51.5 mg, 99%) as a white solid. ¹H-NMR (400 MHz, CDCl₃) δ 8.00 (d, *J* = 8.0 Hz, 1H), 7.80 (d, *J* = 8.0 Hz, 1H), 7.58 (d, *J* = 8.4 Hz, 1H), 7.45-7.55 (m, 2H), 7.10-7.30 (m, 6H), 4.96 (s, 1H); ¹³C-NMR (100 MHz, CDCl₃) δ 50.73, 119.79, 121.17, 121.54, 124.40, 126.93, 128.21, 128.24, 128.31, 129.13, 134.14, 135.05, 149.58, 175.58. MS(EI): m/z (%): 360 (M⁺, 27.90), 231 (100); HRMS (EI) calcd. for C₁₈H₁₂O₂: 260.0837, found: 260.0841.

6. Preliminary mechanism studies

We performed a series of control experiments to confirm *ortho* selectivity stemmed from the coordination between hydroxyl group and gold. First, the reaction between 1-methoxynaphthalene and diazo compound under standard condition was conducted, delivering the *para* C-H bond functionalization product in 92% yield (scheme 1, eq 1). The structure was confirmed by additional experiment (scheme 1, eq 2, eq 3). In order to exclude the *para* selectivity arise from steric hindrance, we switch diazo compound to methyl 2-diazo-2-phenylacetate with less steric hindrance, affording only *para* C-H bond functionalization product (scheme 1, eq 4). These results indicated that the weak interaction between naphthol and gold catalyst played a key role in the *ortho*-selectivity.

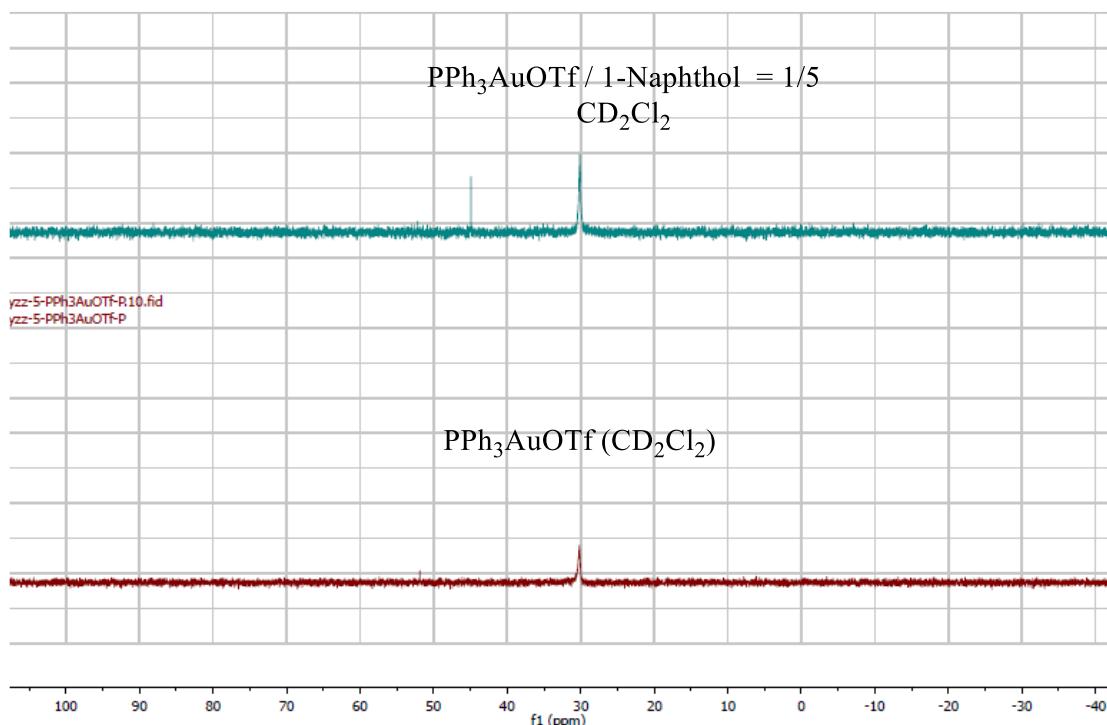
The date of **10** was illustrated as following:

¹H-NMR (400 MHz, CDCl₃) δ 8.31 (d, *J* = 7.6 Hz, 1H), 7.93 (d, *J* = 7.6 Hz, 1H), 7.40-7.55 (m, 2H), 7.20-7.35 (m, 6H), 6.74 (d, *J* = 8.4 Hz, 1H), 5.64 (s, 1H), 5.11 (hept, *J* = 6.4 Hz, 1H), 3.96 (s, 3H), 1.21 (t, *J* = 6.8 Hz, 6H); ¹³C-NMR (100 MHz, CDCl₃) δ 21.66, 53.43, 55.41, 55.44, 68.56, 103.15, 122.70, 123.08, 124.94, 126.03, 126.31, 126.81, 127.12, 128.50, 128.91, 132.54, 138.49, 154.98, 172.52. MS(EI): m/z (%): 334 (M⁺, 24.12), 247 (100); HRMS (EI) calcd. for C₂₂H₂₂O₃: 334.1569, found: 334.1566.



Scheme 1 Weak interaction study

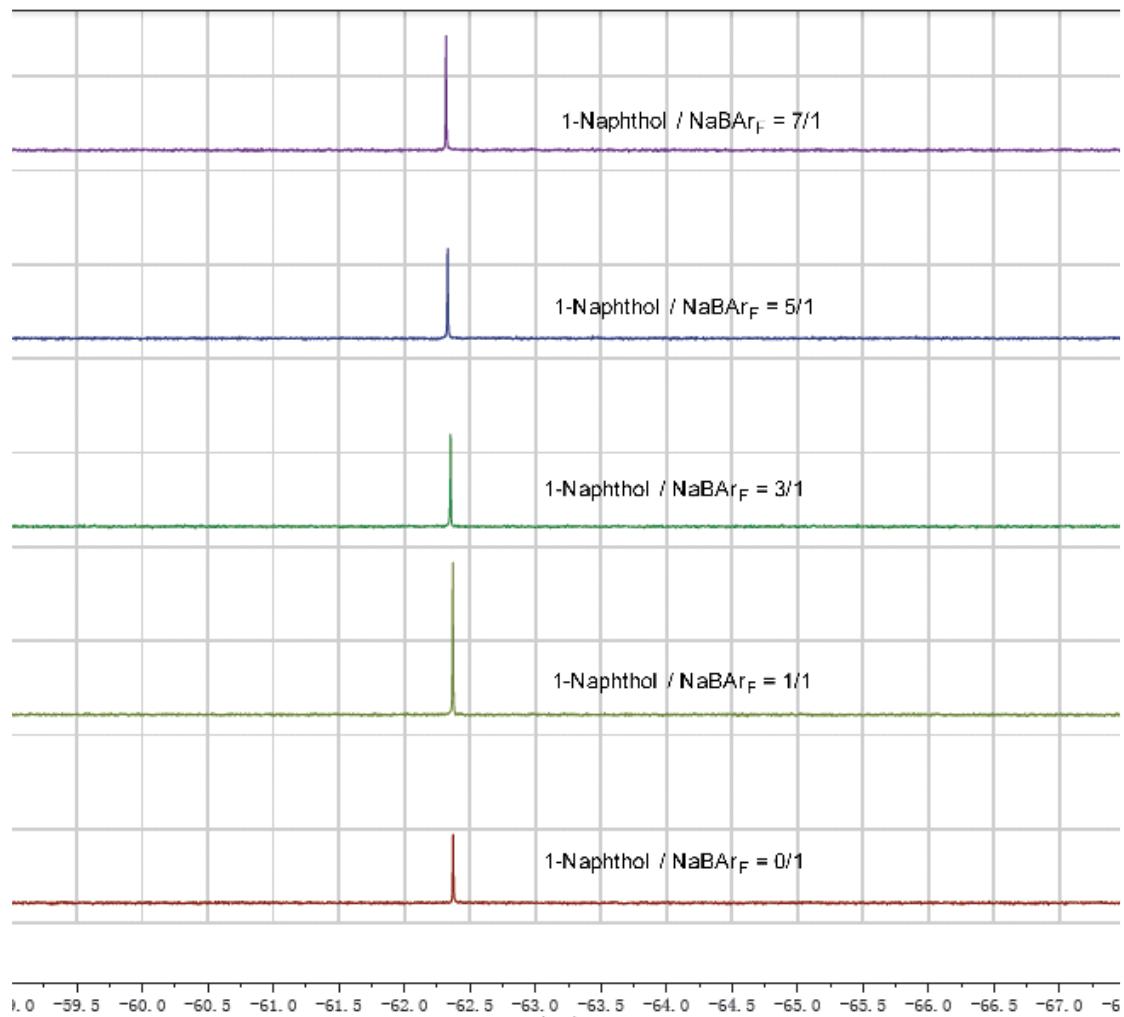
Subsequently, we further conducted a series of related control experiments to identify the type of weak interaction. First, ^{31}P -NMR experiment shown that the coordination between gold catalyst and hydroxyl group was indeed existed (scheme 2).



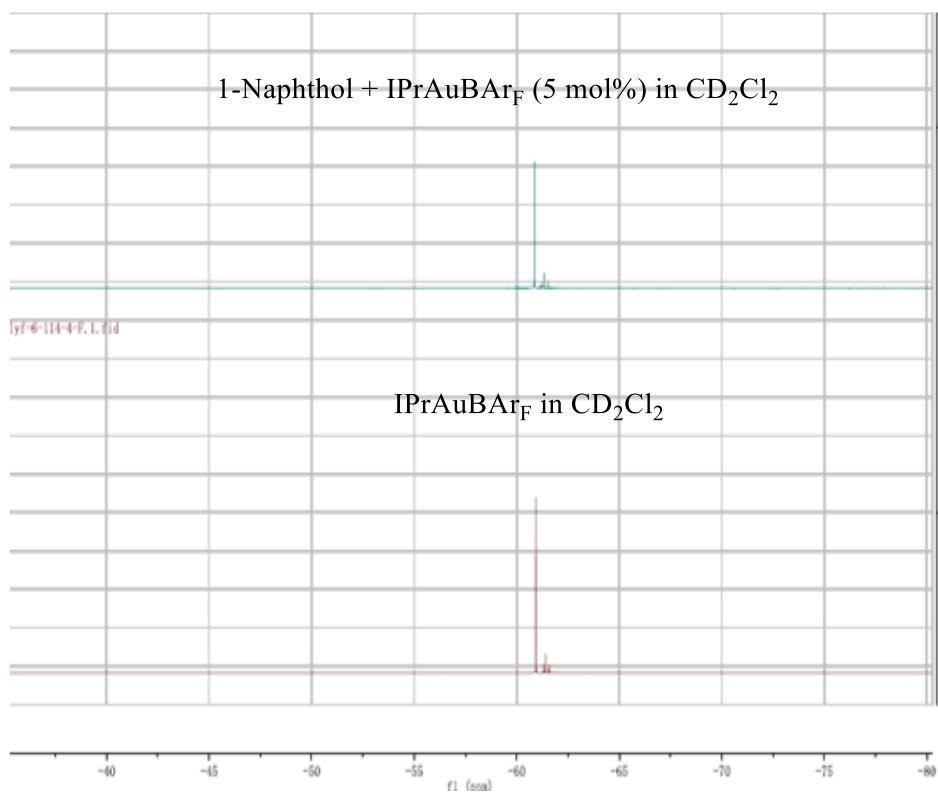
Scheme 2 ^{31}P -NMR experiments

Meanwhile, NMR titration disclosed that no hydrogen bonding interaction was existed between naphthol and gold catalyst (scheme 3).

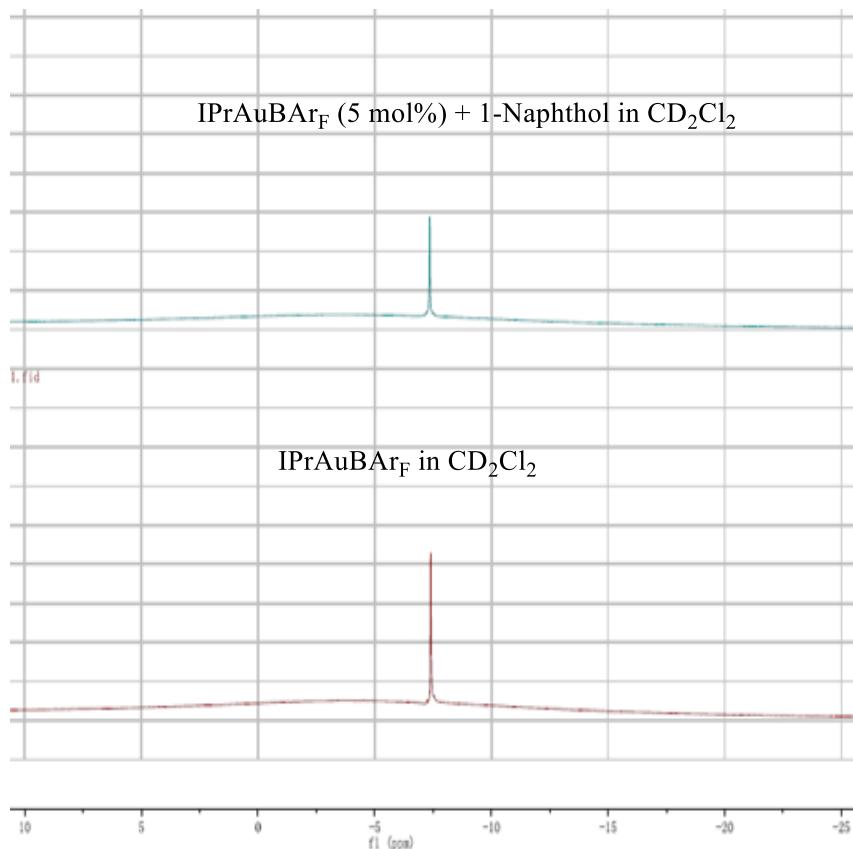
Allowing for the fact the tetrasubstituted boron compounds could undergo SN₂ reaction (*Organometallics* **1998**, *17*, 4155; *J. Fluor. Chem.* **2014**, *168*, 111; *Inorg. Chem.* **2004**, *43*, 5789) and has certain lewis acidity (*Chem. Eur. J.* **2017**, *23*, 5219), we wanted to know whether the interaction between hydroxyl group and boron was the key for *ortho*-selectivity. ^{19}F -NMR experiment shown that no new compound was generated when 1-naphthol, IPrAuCl and NaBAr_F were mixed. Meanwhile, the ^{11}B -NMR experiment revealed that no coordination interaction between hydroxyl group and boron atom. Based on these results, the possibility that the *ortho*-selectivity stem from the interaction between hydroxyl group and boron atom could be excluded (scheme 6).



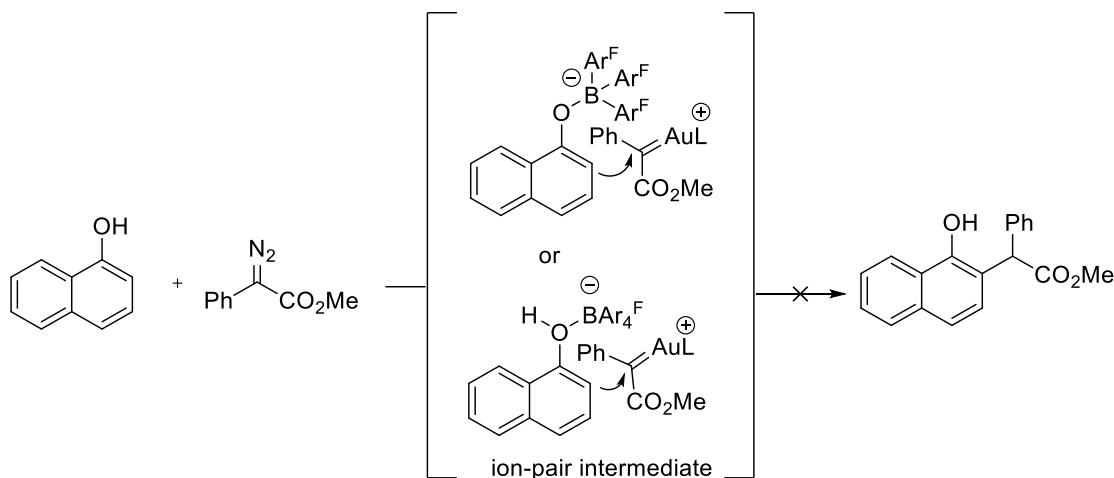
Scheme 3. Titration experiments



Scheme 4. ¹⁹F-NMR experiments

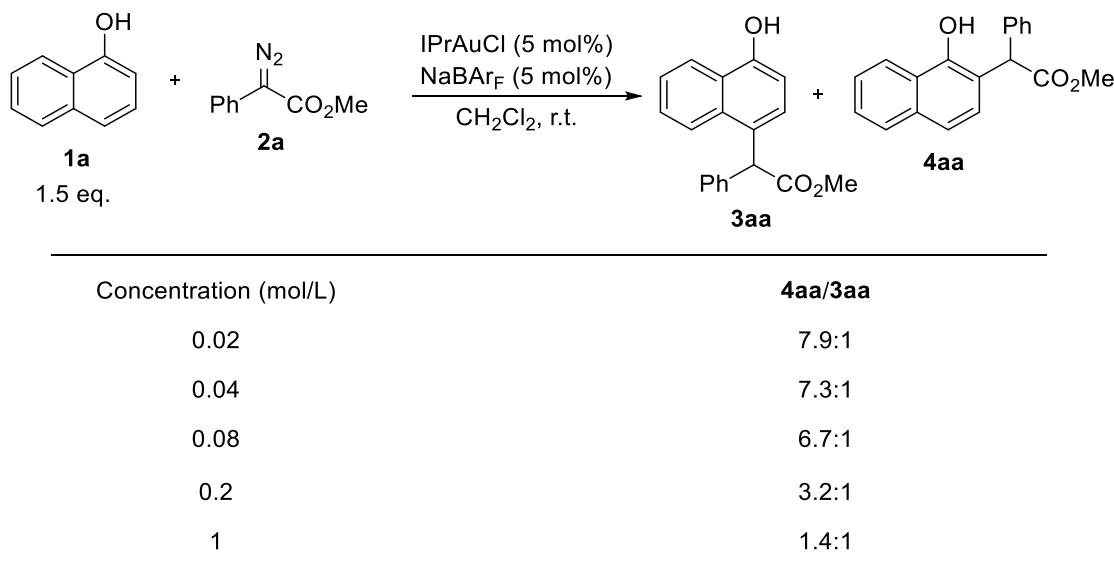


Scheme 5. ¹¹B-NMR experiments



Scheme 6. Impossible mechanism for *ortho*-selectivity

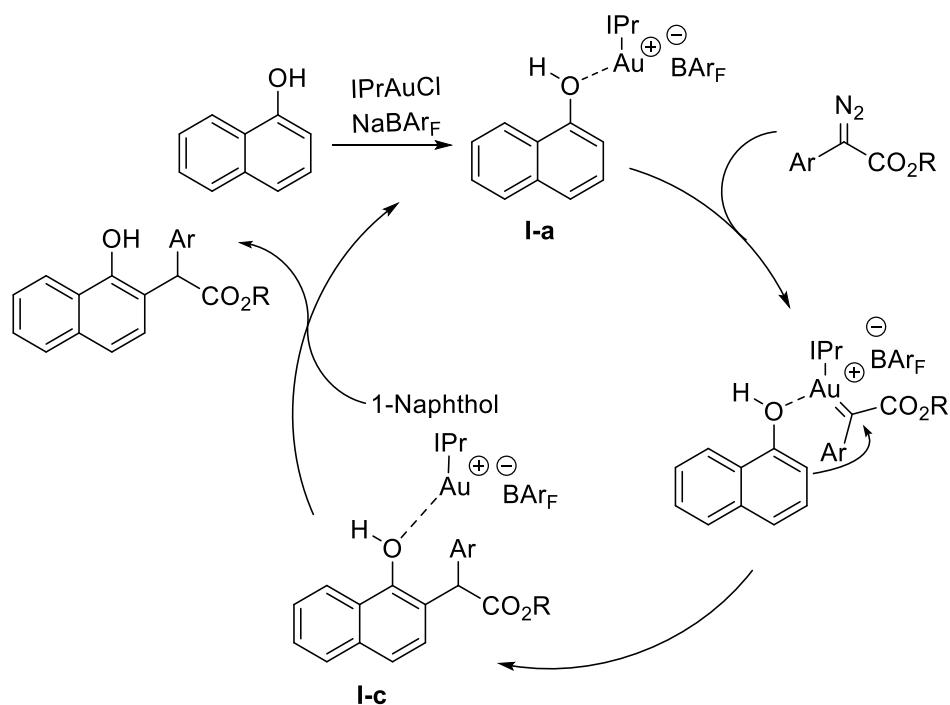
In order to confirm our conclusion that the *ortho* C-H bond functionalization is an intramolecular reaction and *para* C-H bond functionalization is an intermolecular reaction, we performed the reactions under different concentration. As we know that the low concentration is good for the intramolecular reaction while high concentration would facilitate the intermolecular reaction. Thus, the *para* C-H bond functionalization product should become more with increasing of concentration under standard conditions. To my excitement, the reactions were indeed proceeded as our expectation (scheme 7).



Scheme 7. Concentration experiments

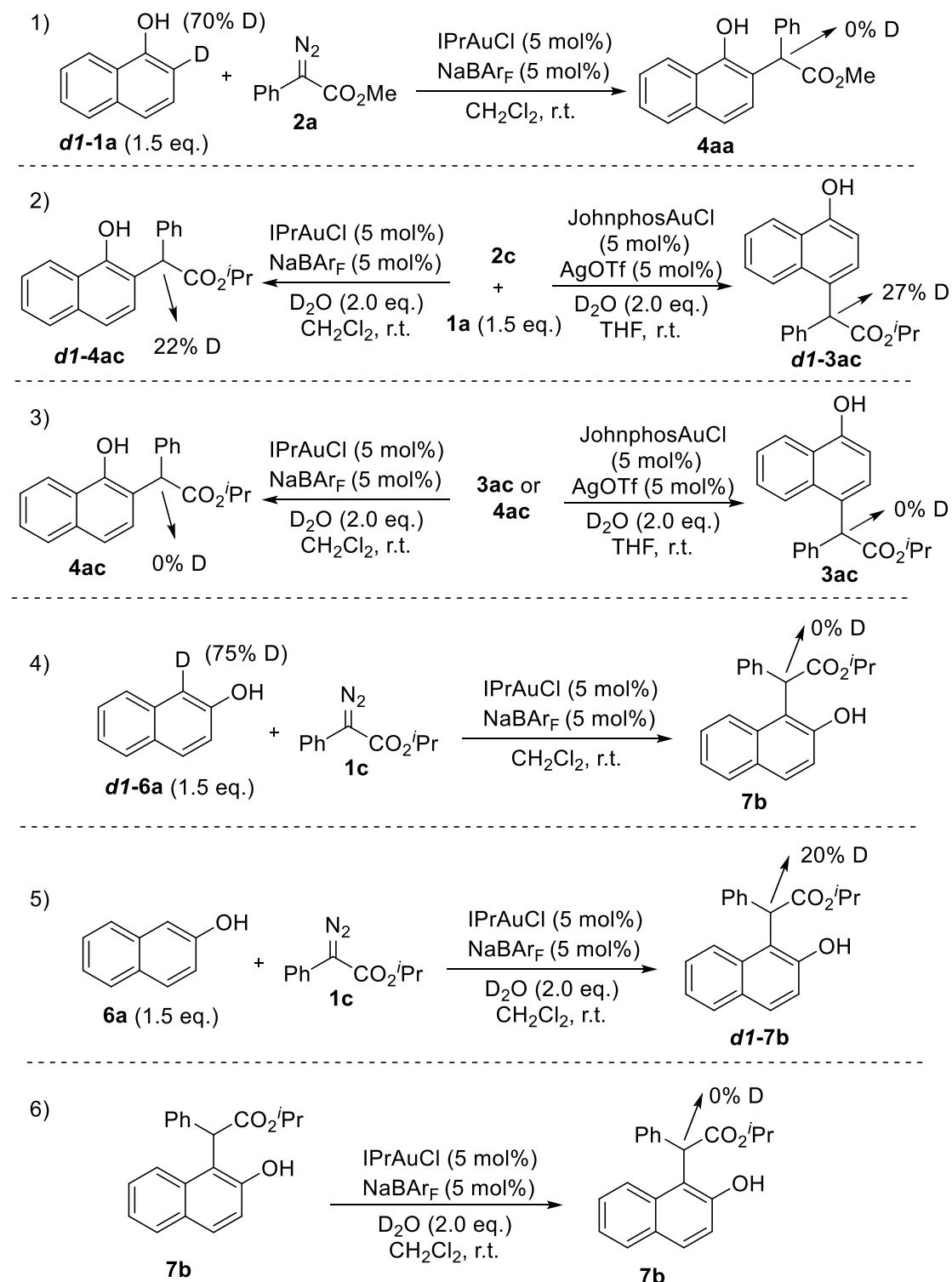
Based on above results, we could conclude that the coordination between hydroxyl

group and gold was the key for the *ortho*-selectivity. The proposed mechanism was illustrated as following (scheme 8).



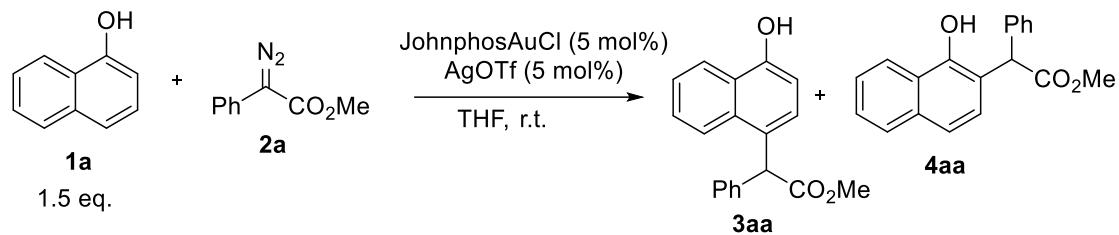
Scheme 8. Proposed mechanism for *ortho*-selectivity

Besides, related deuterium labeling experiments were also conducted to get more insights into mechanism (scheme 9). The results indicated that the reaction did not proceeded via a [1,2]-H shift but a possible water assisted hydrogen. The results of control experiments also demonstrated that no enolization took place in our reaction.



Scheme 9. Deuterium labeling experiments

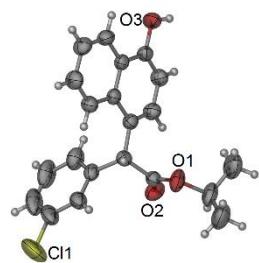
Finally, we studied the silver effect, the results disclosed that no silver effect in para C-H bond functionalization (scheme 10).



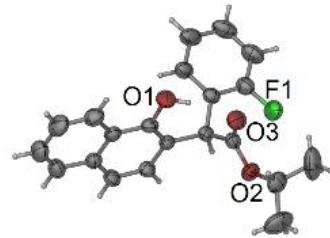
- 1) with AgCl in reaction system, **3aa** (90% NMR yield), **4aa** (0% NMR yield)
 2) without AgCl in reaction system, **3aa** (86% NMR yield), **4aa** (0% NMR yield)

Scheme 10. Silver effect studies

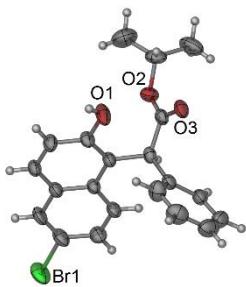
7. X-ray crystal date for 3ah, 4al, 7i and 9



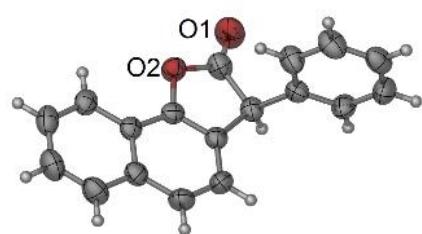
3ah



4al

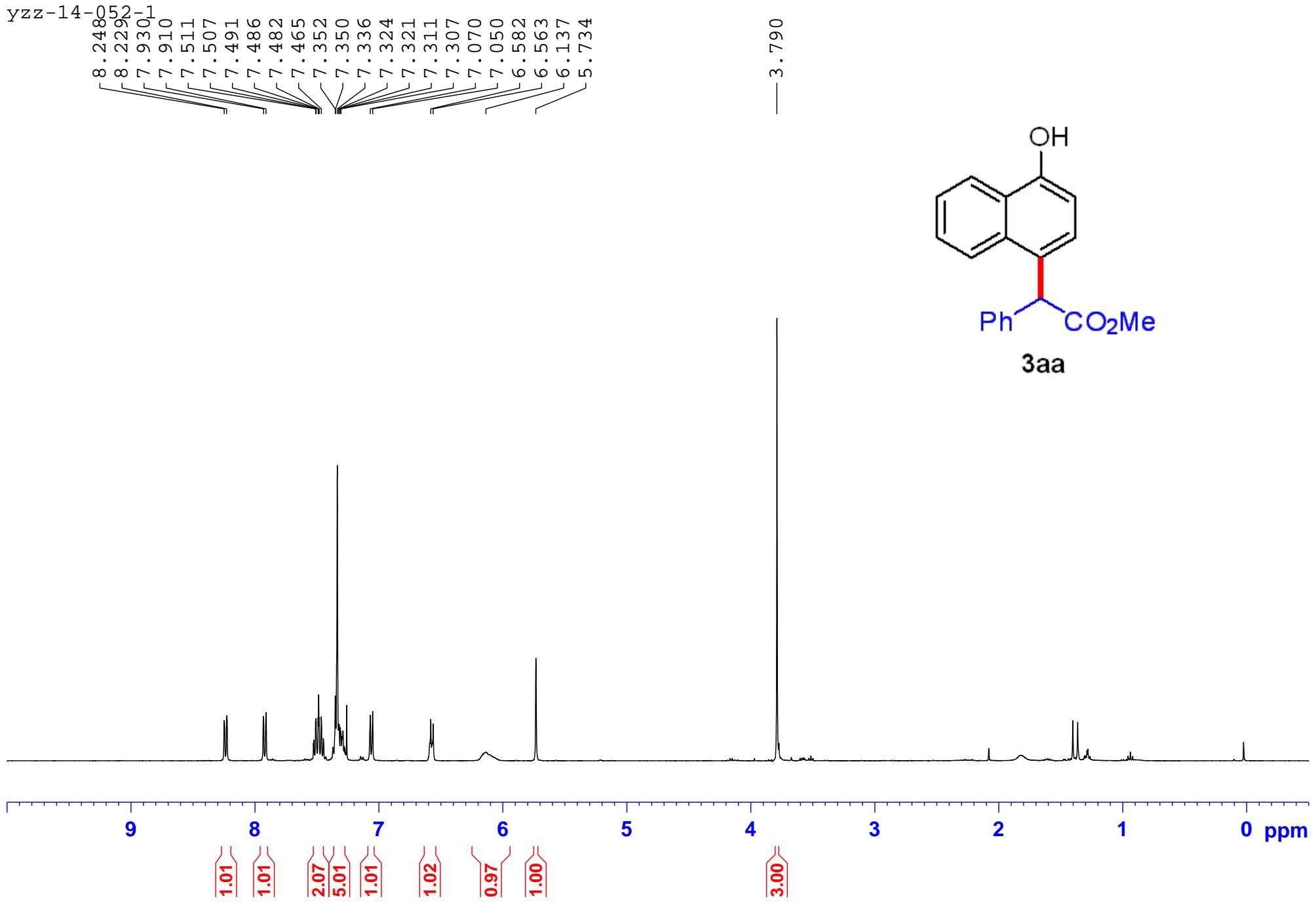


7i

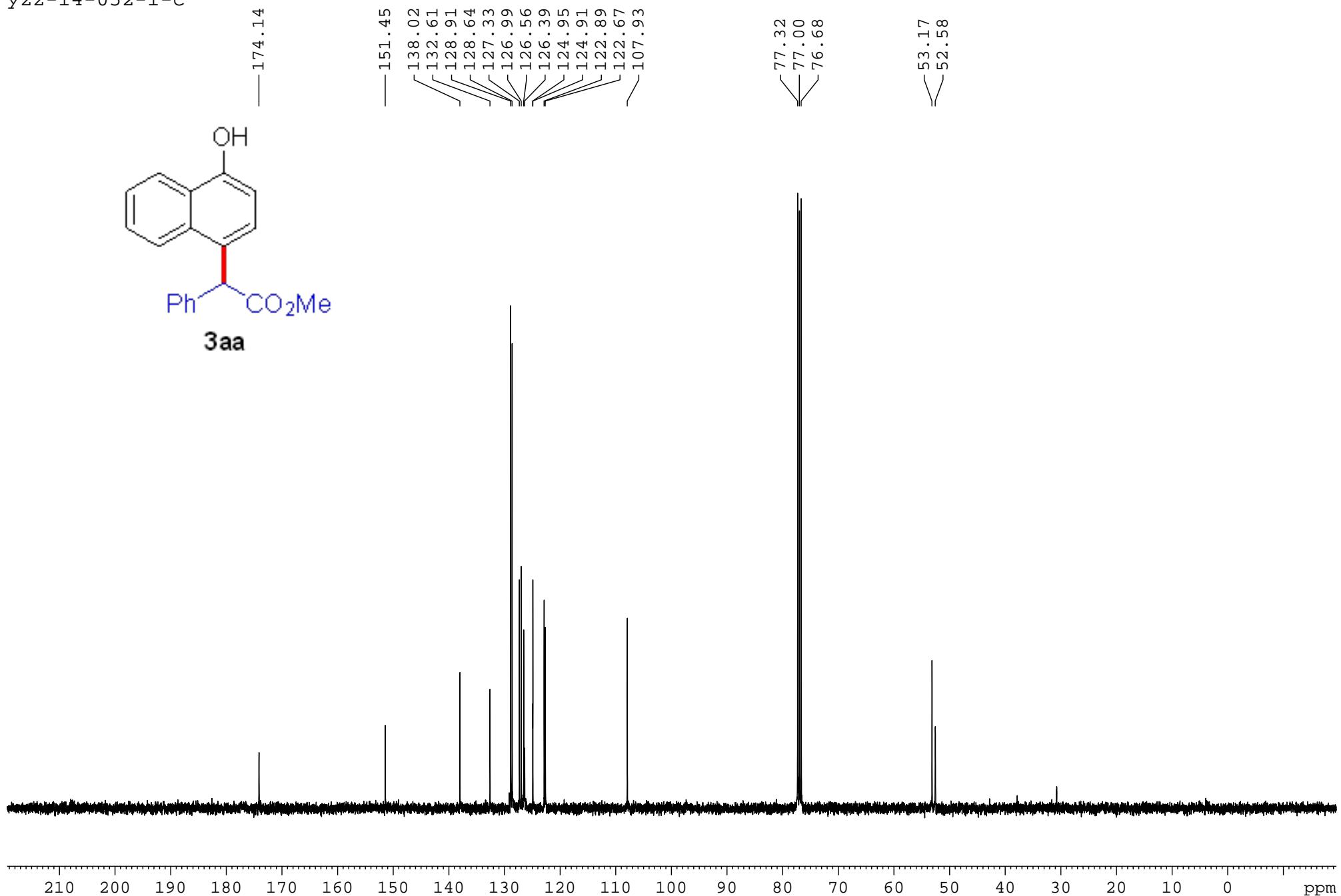


9

8. NMR Spectra of new compounds



yzz-14-052-1-c



8.242

8.222

8.219

7.950

7.930

7.508

7.504

7.487

7.483

7.479

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7.459

7.456

7.352

7.342

7.299

7.288

7.260

7.060

7.040

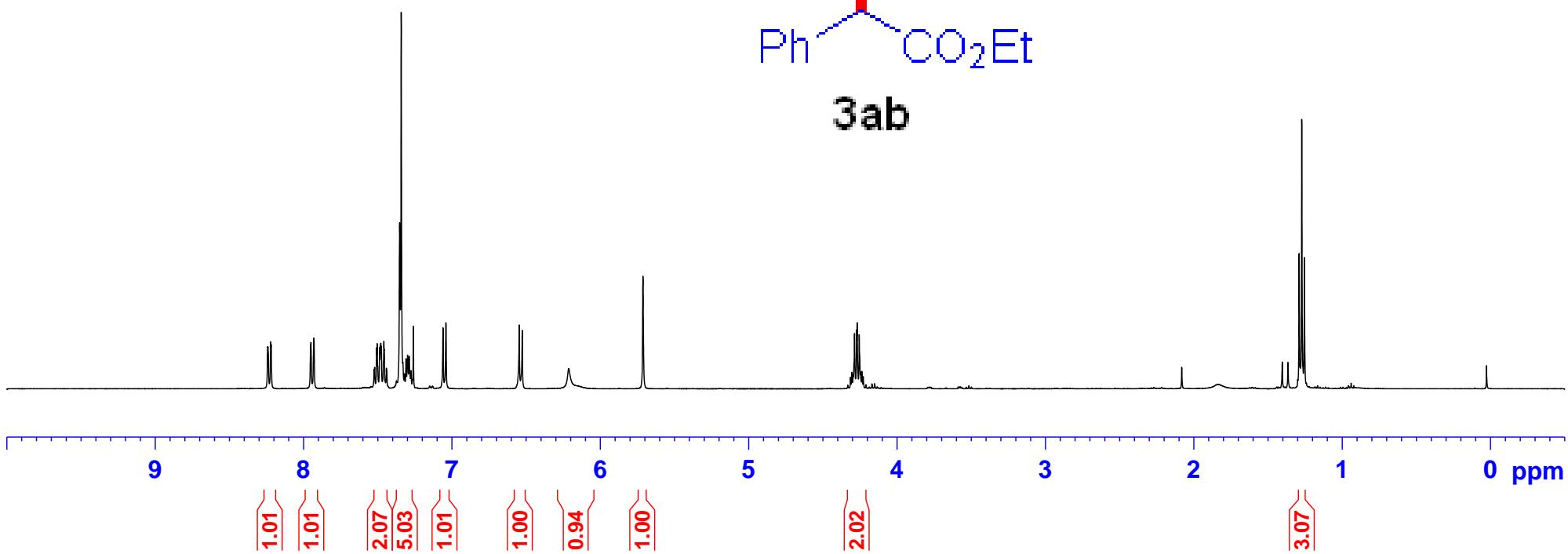
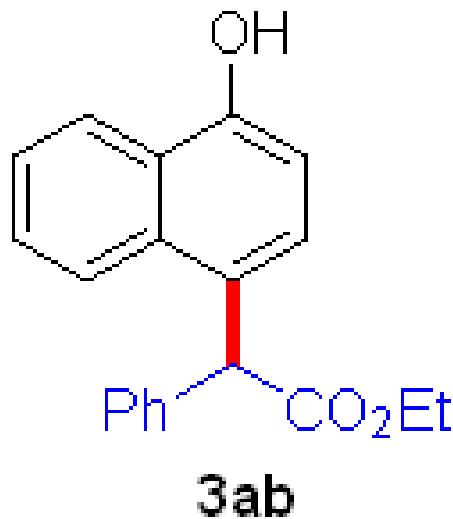
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6.527

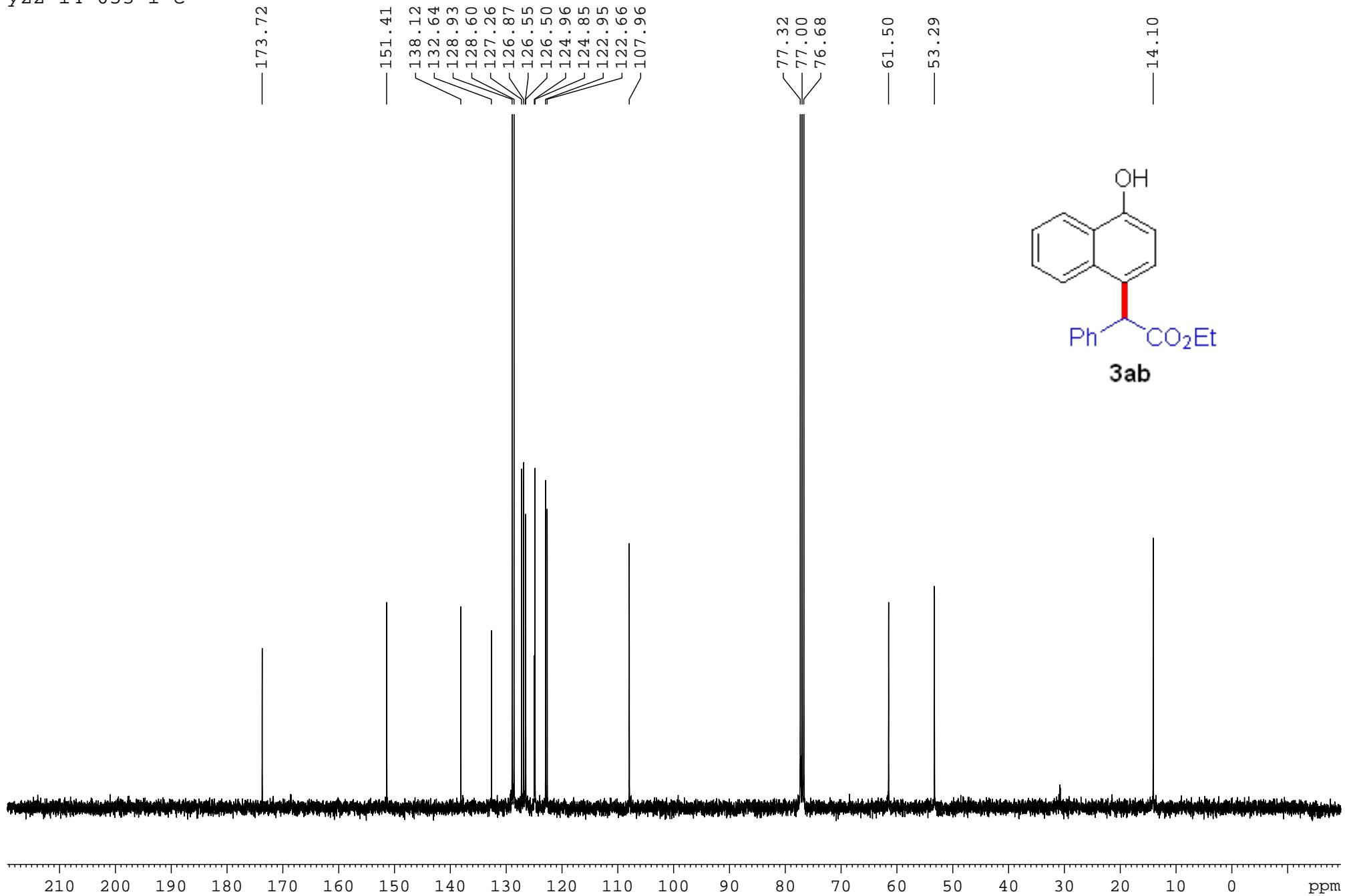
6.214

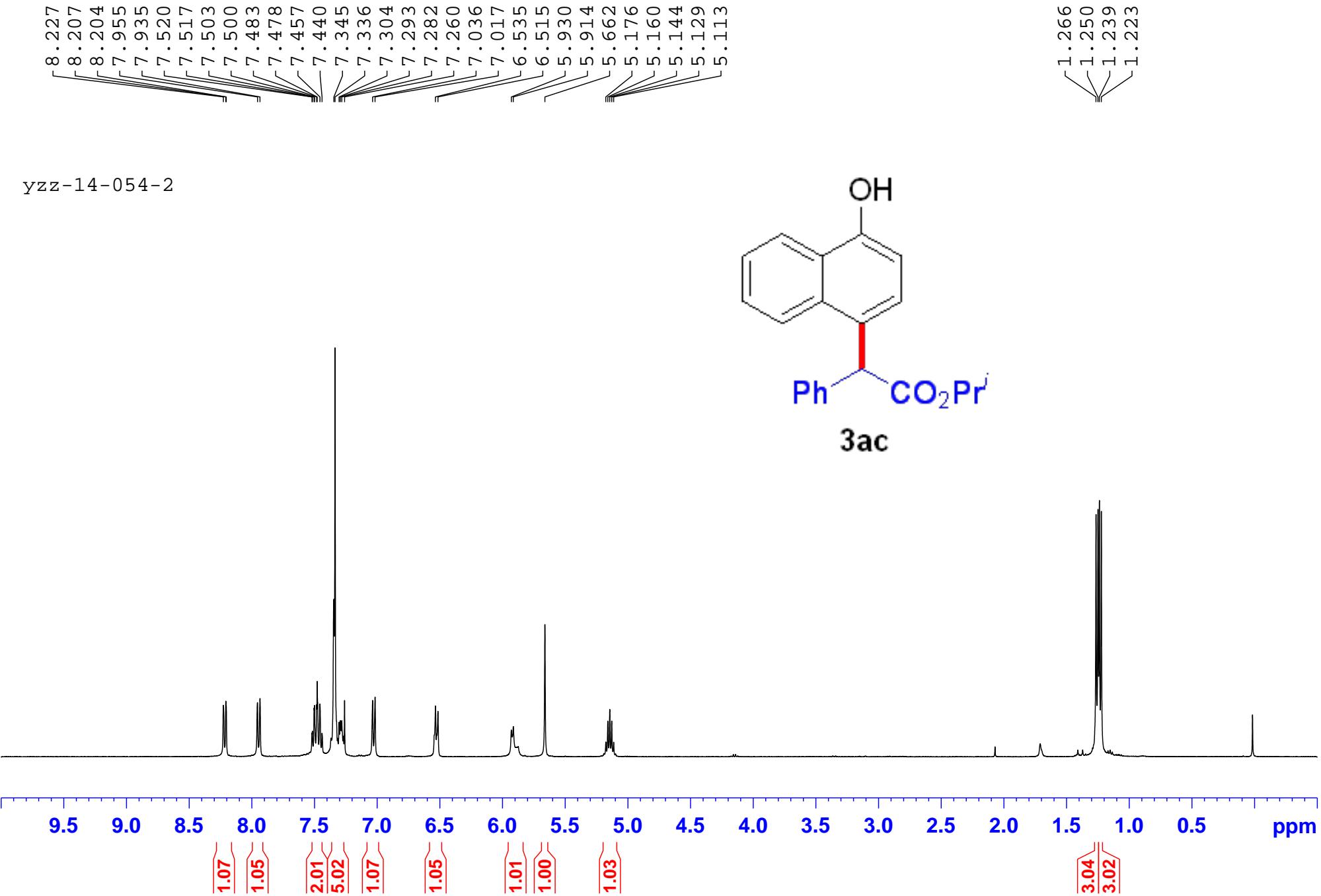
5.713

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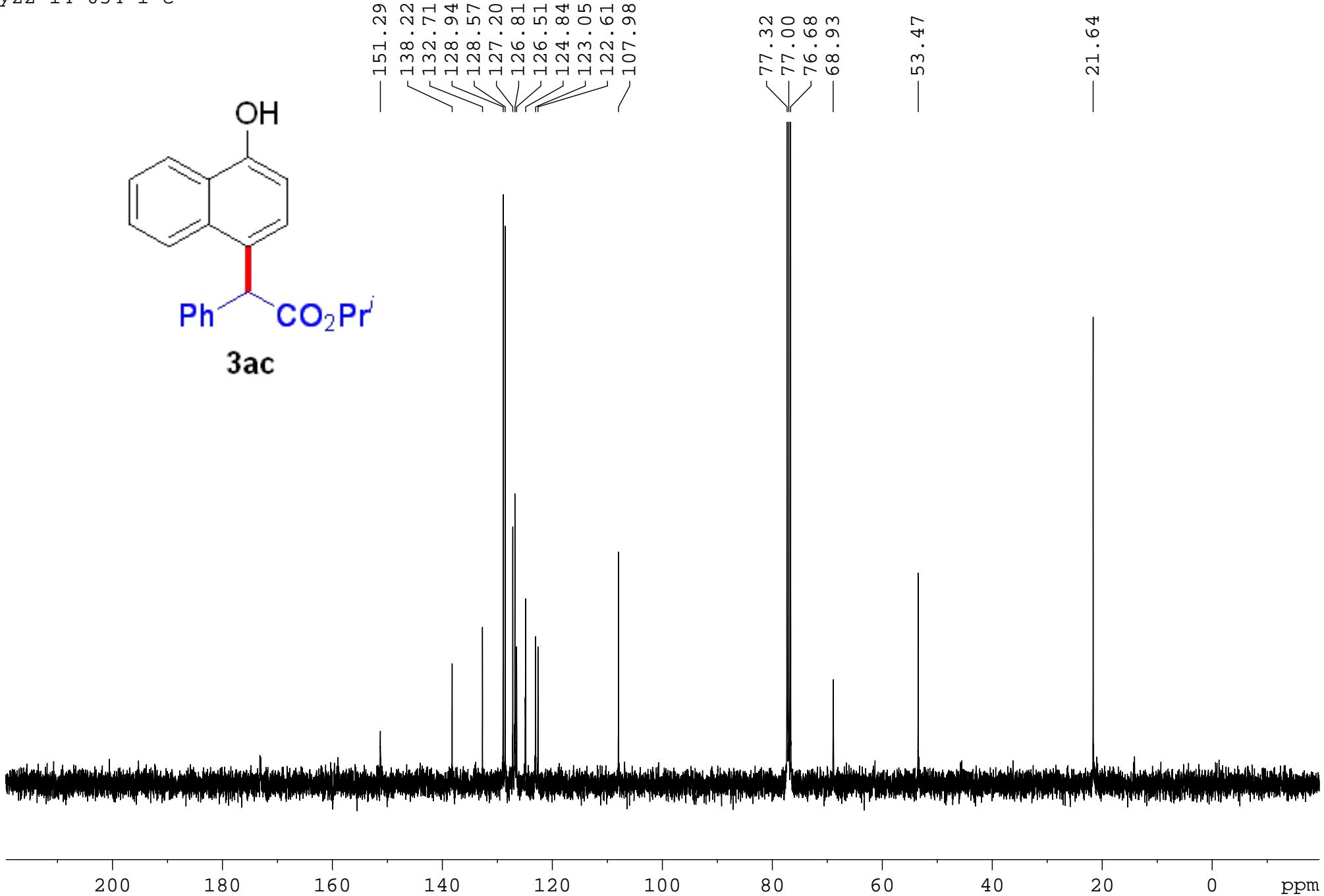
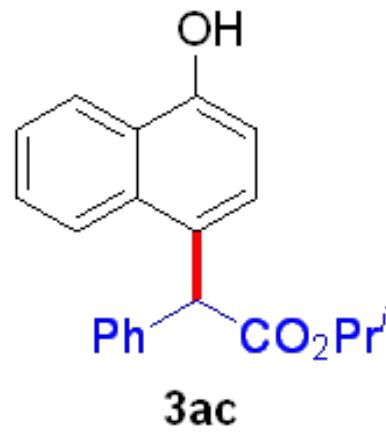
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4.2551.291
1.273
1.255

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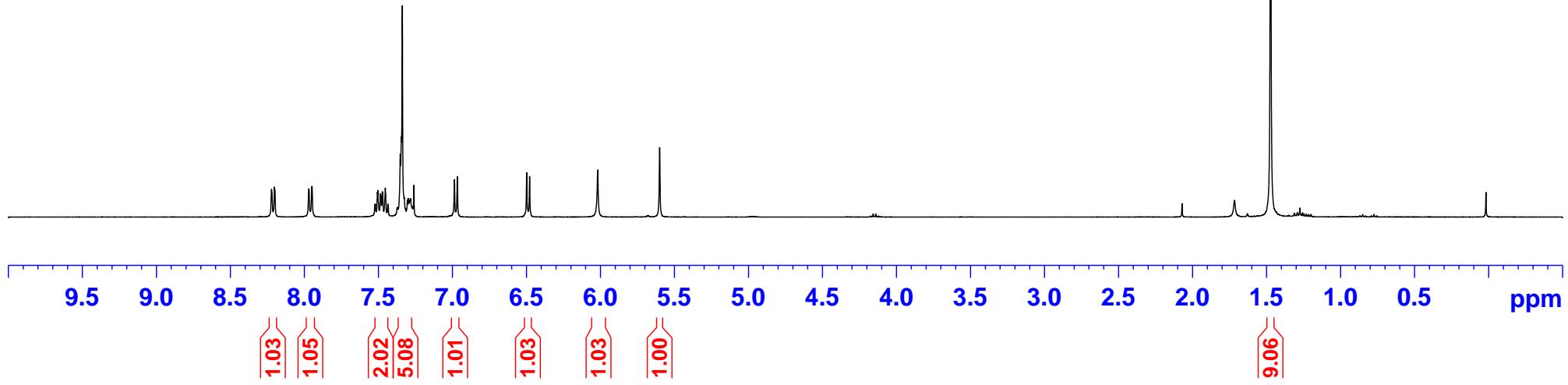
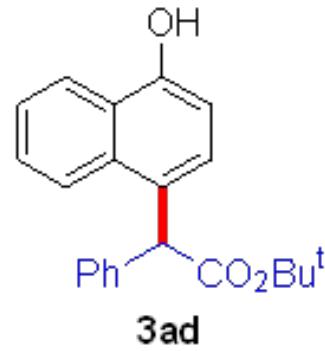


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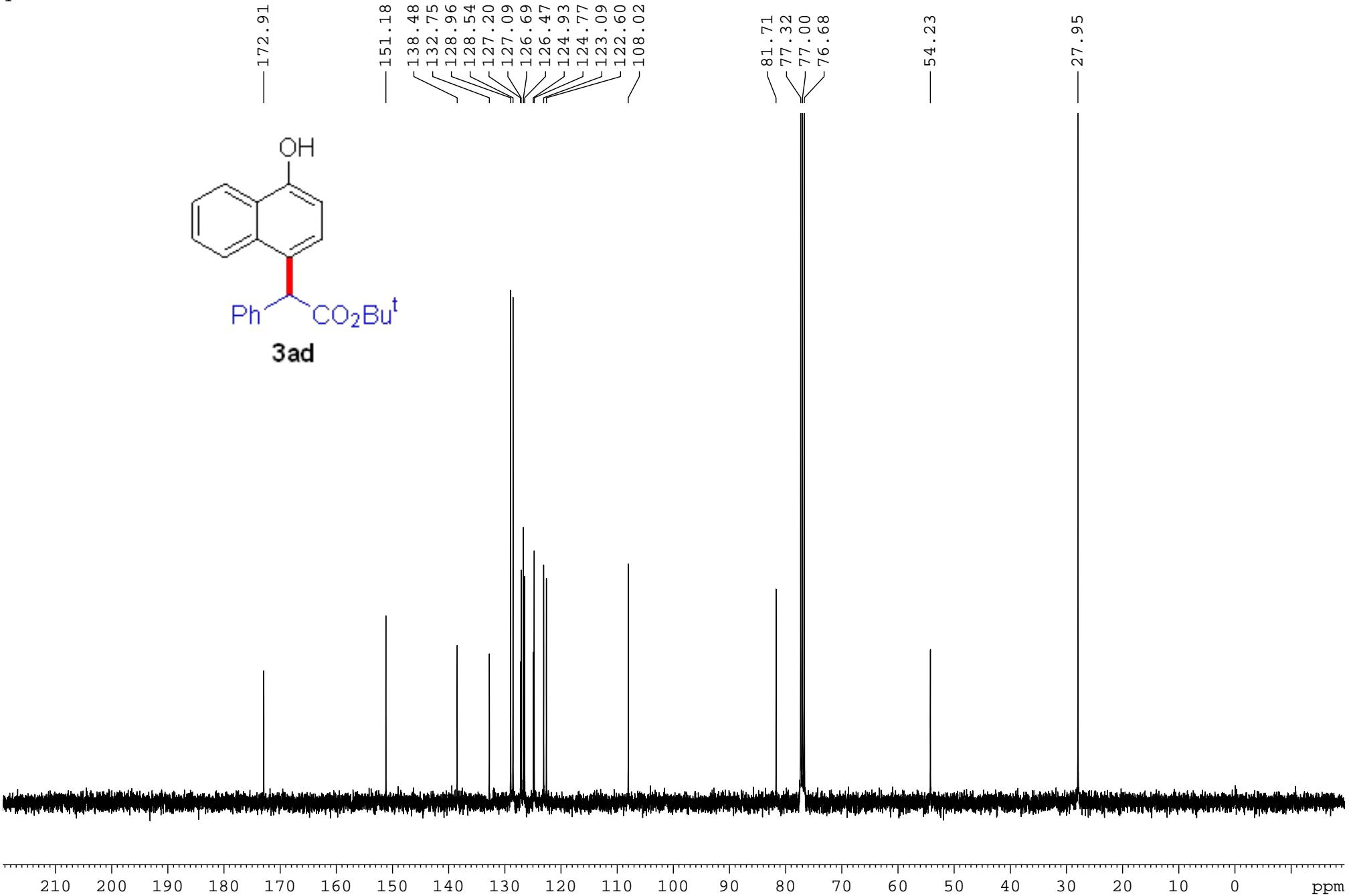


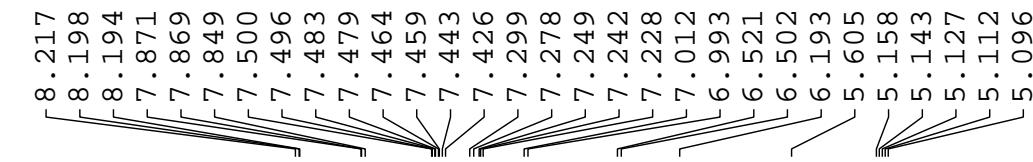
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1.473

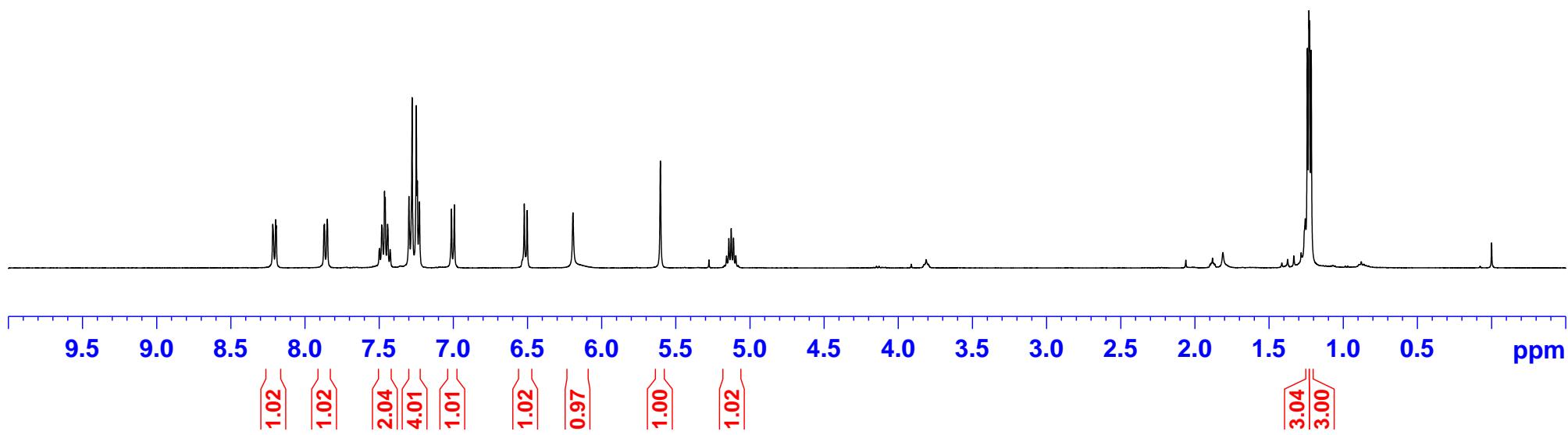
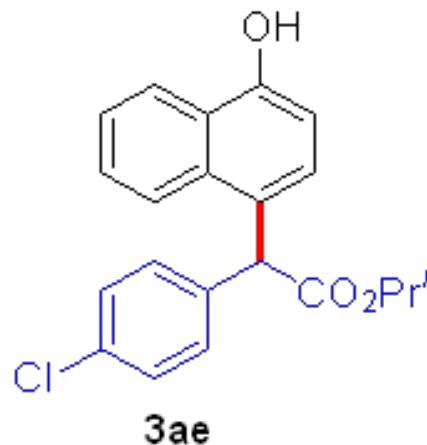


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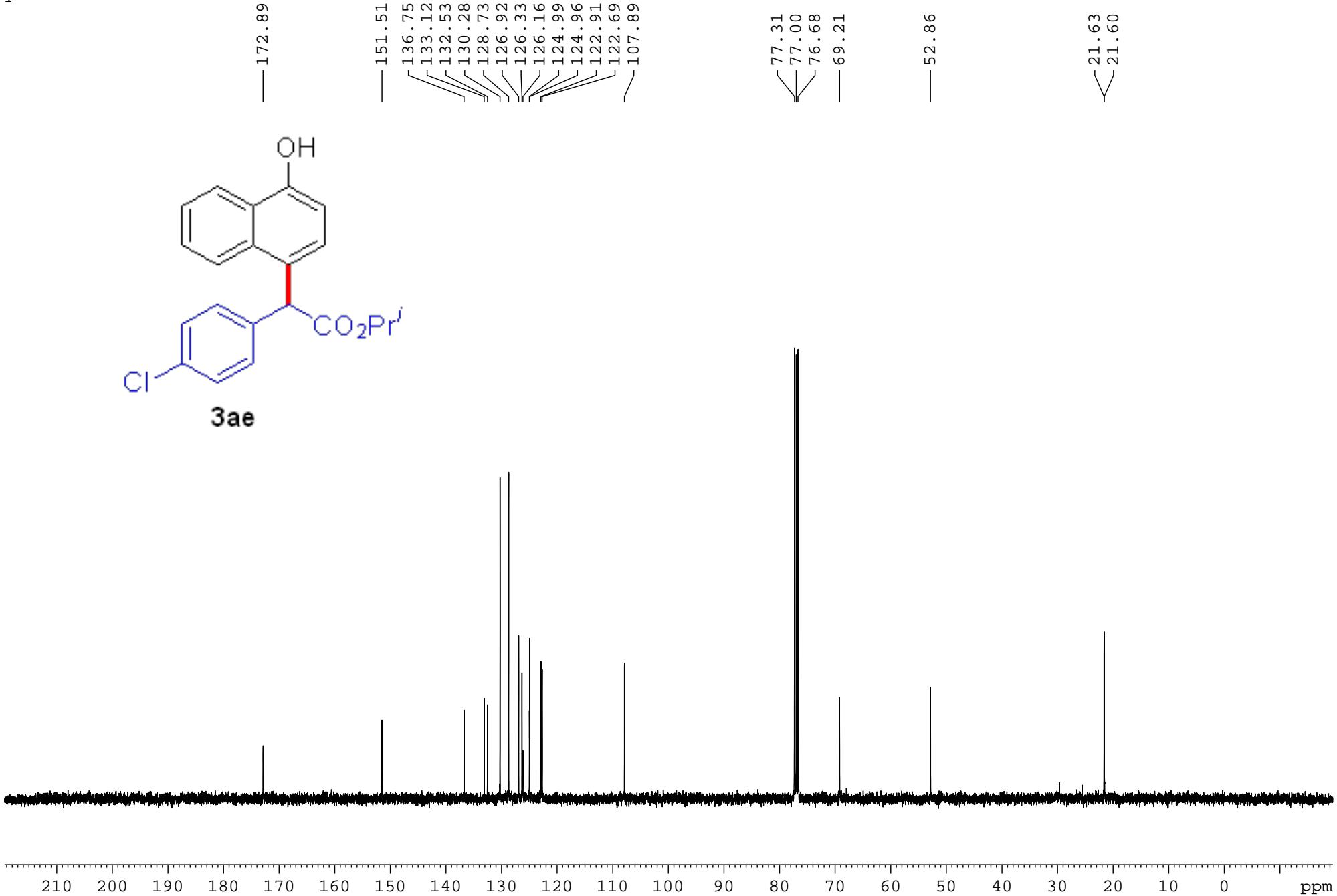
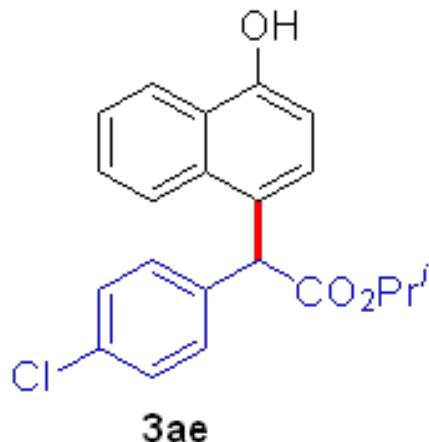




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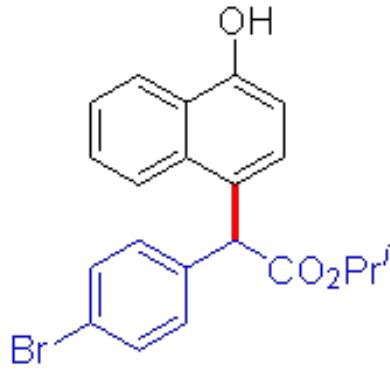
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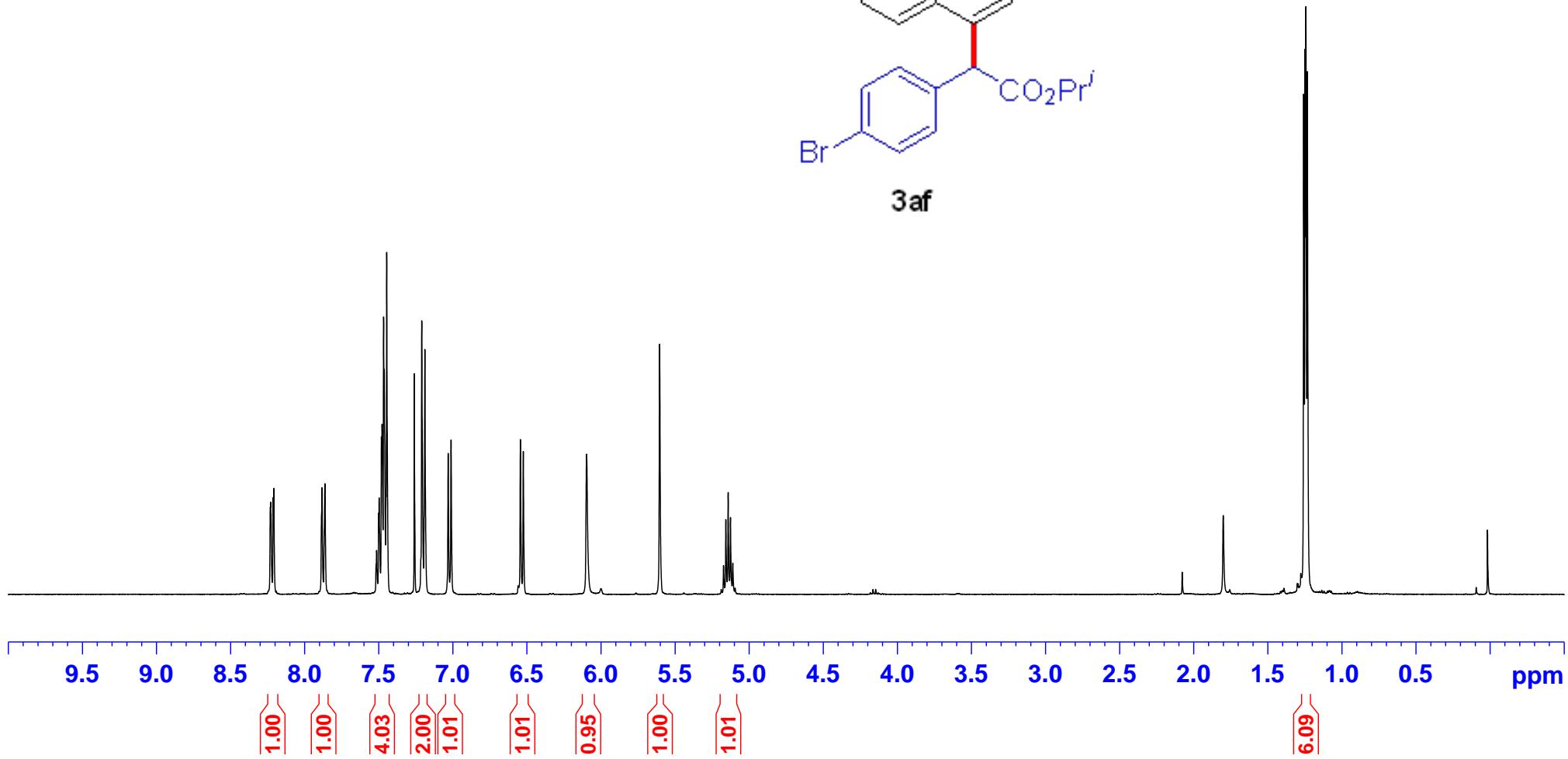
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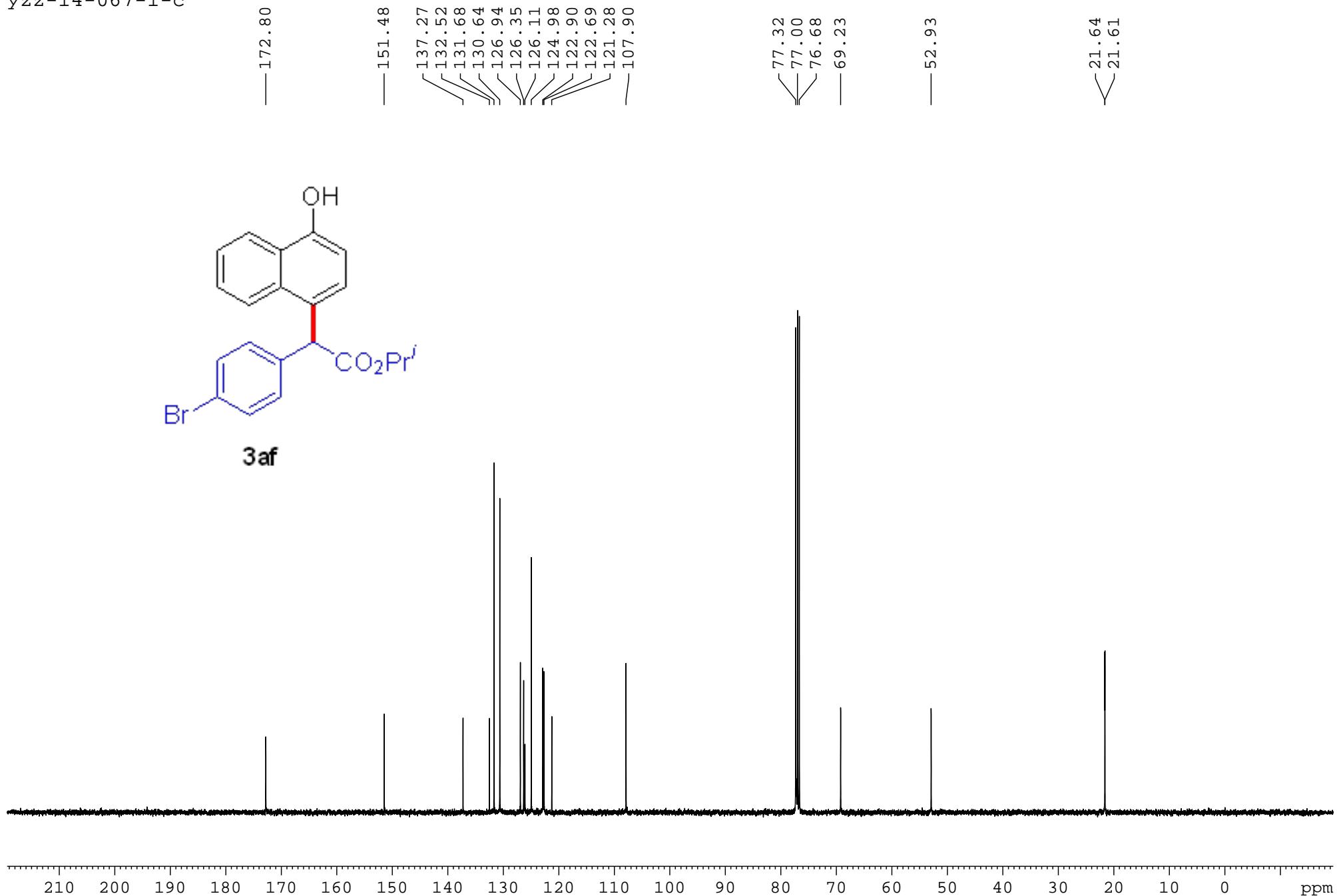
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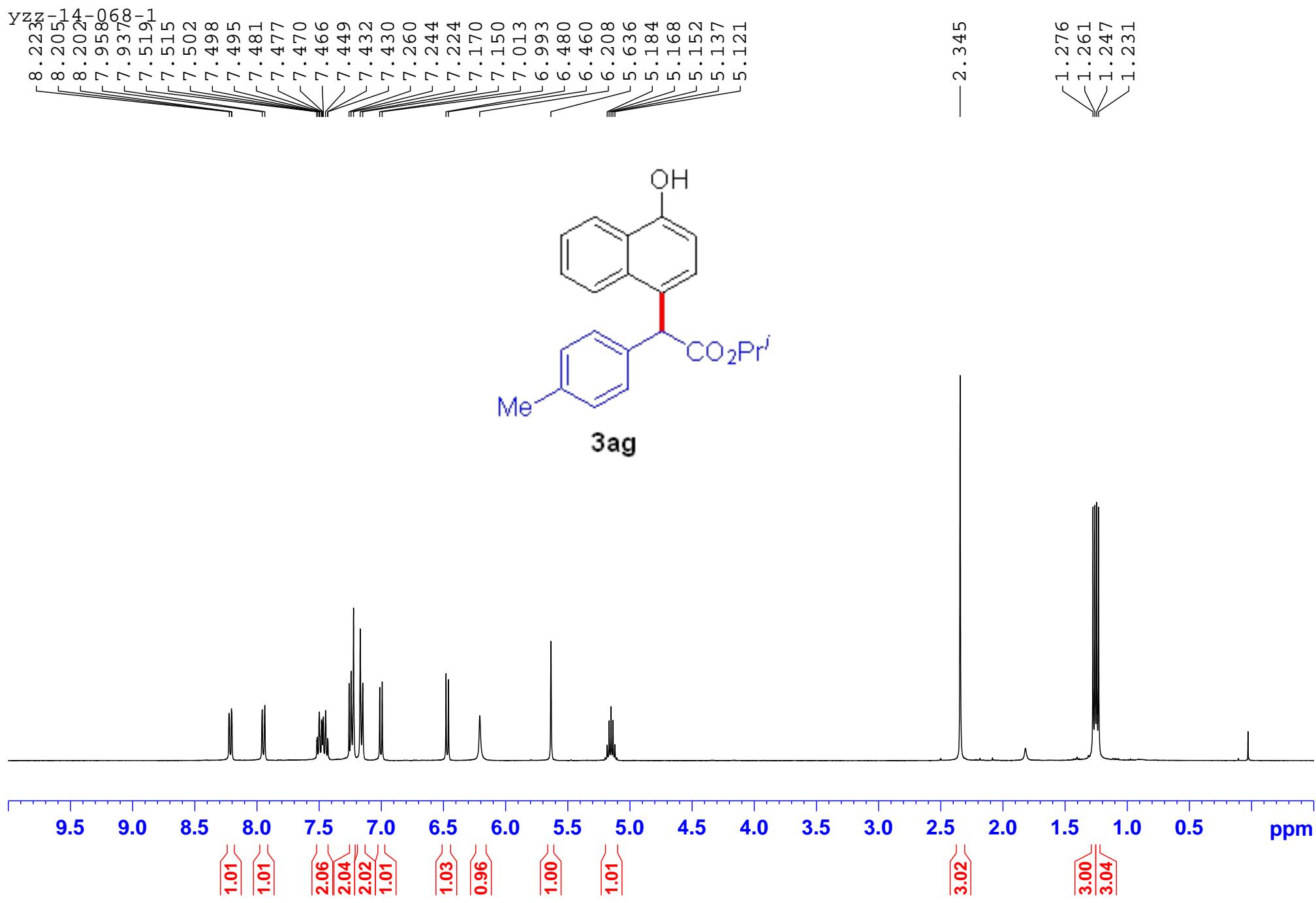


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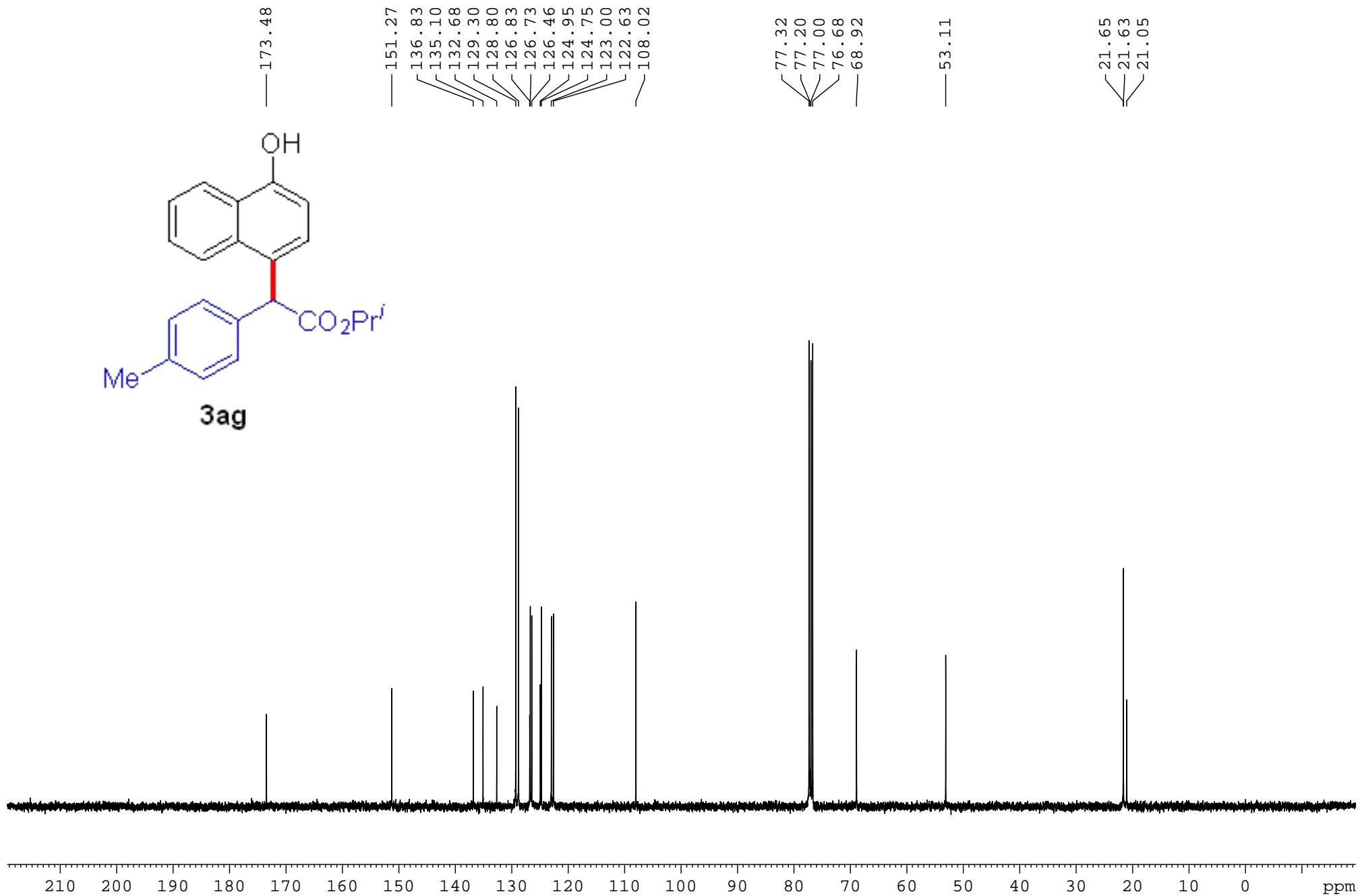


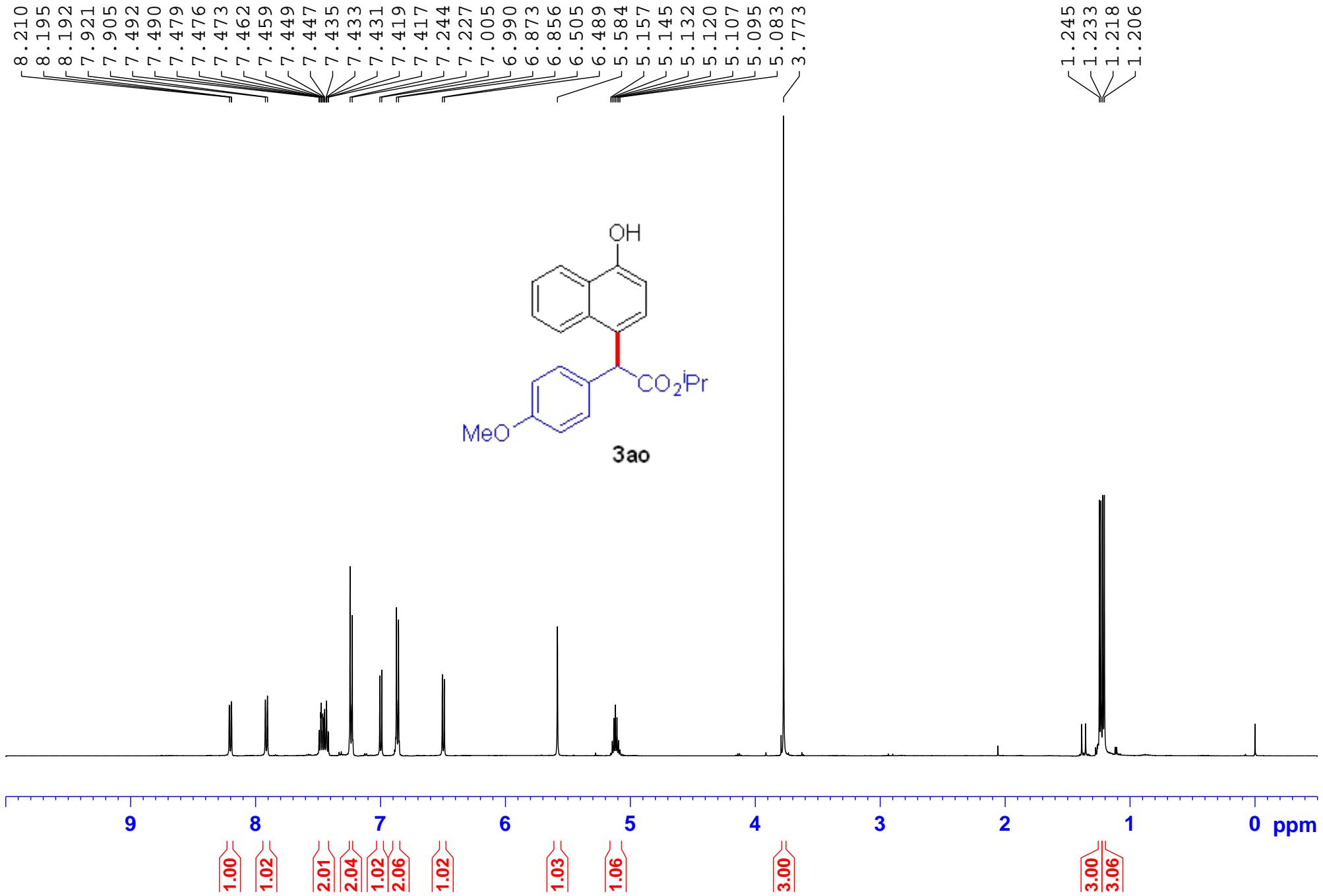
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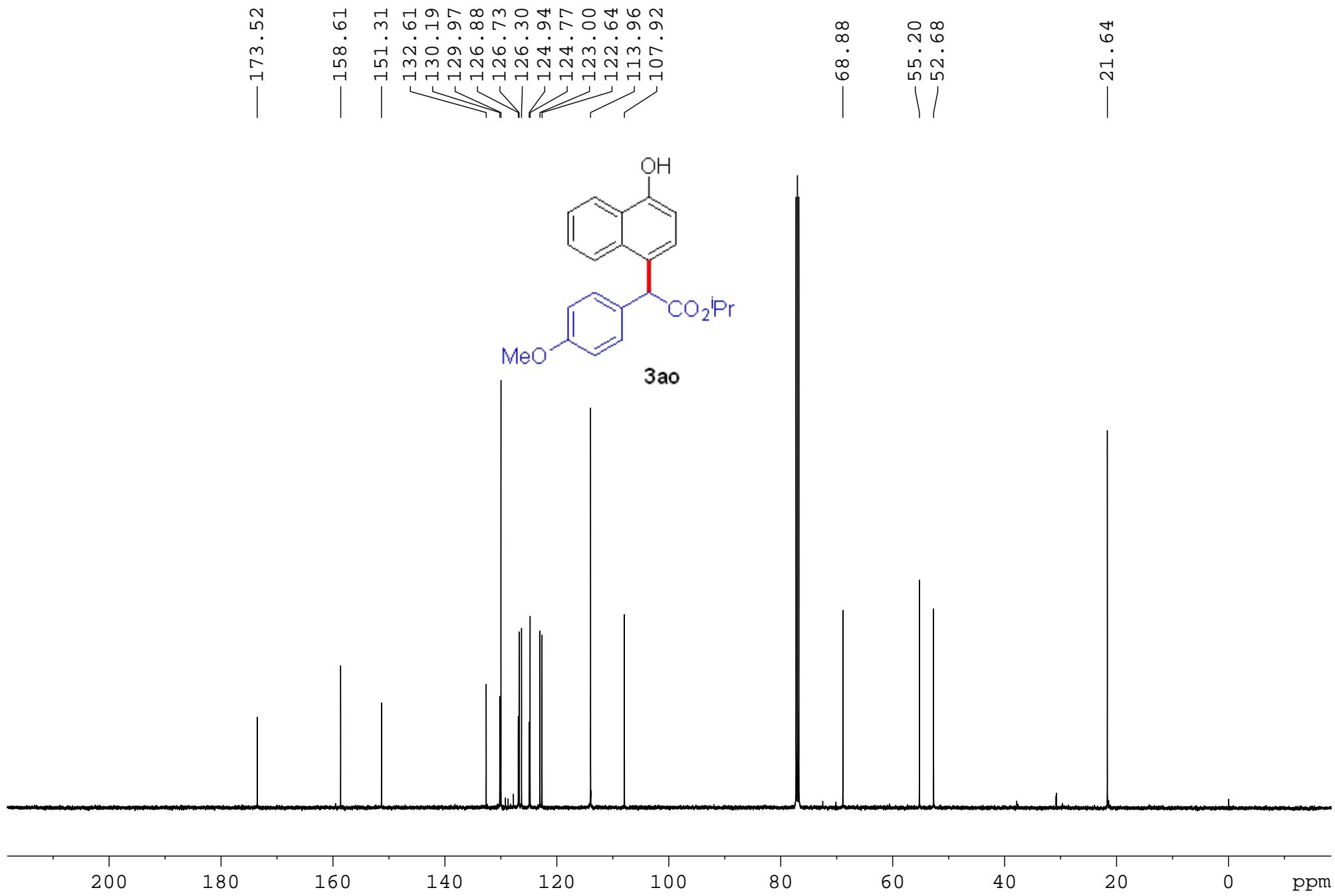


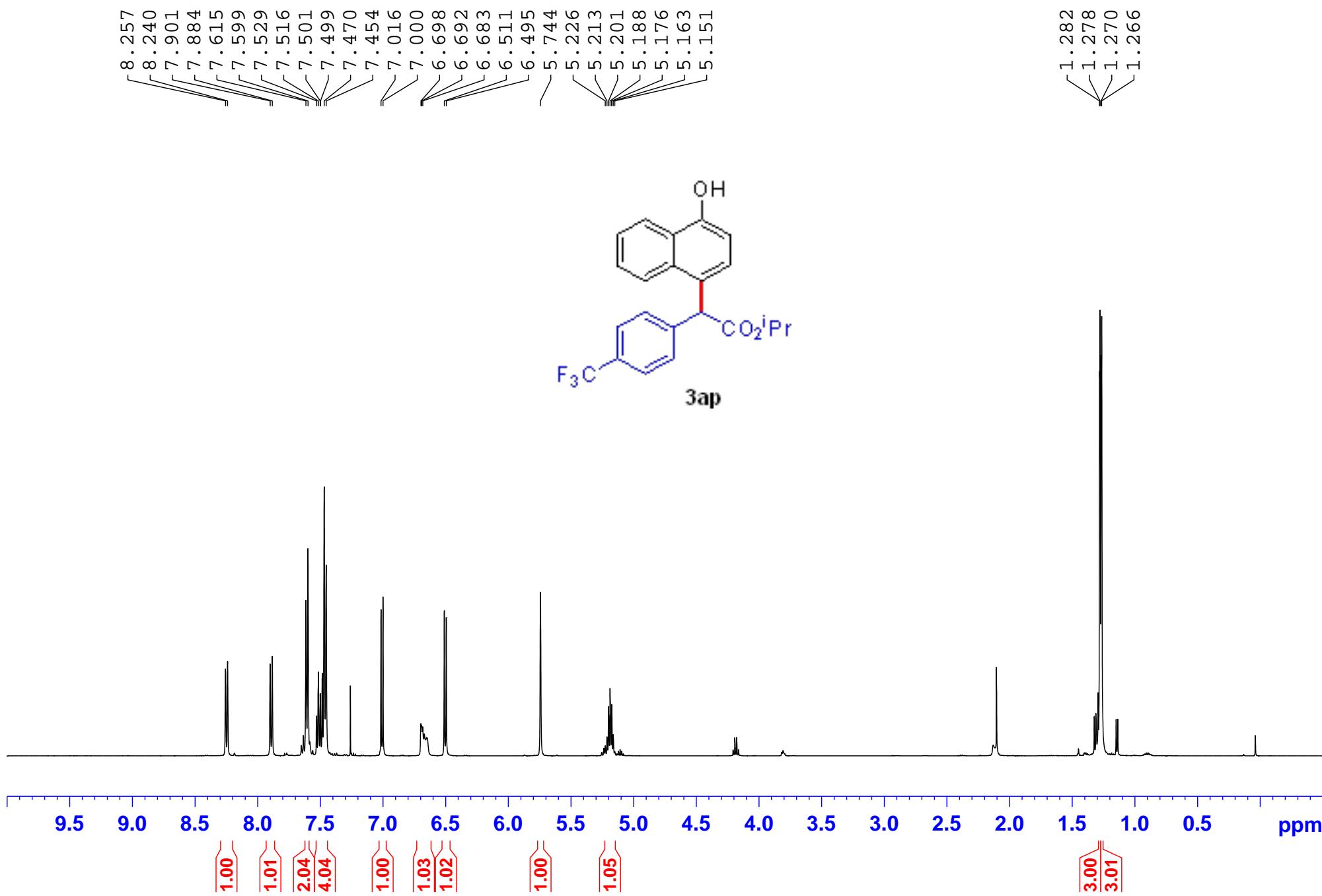


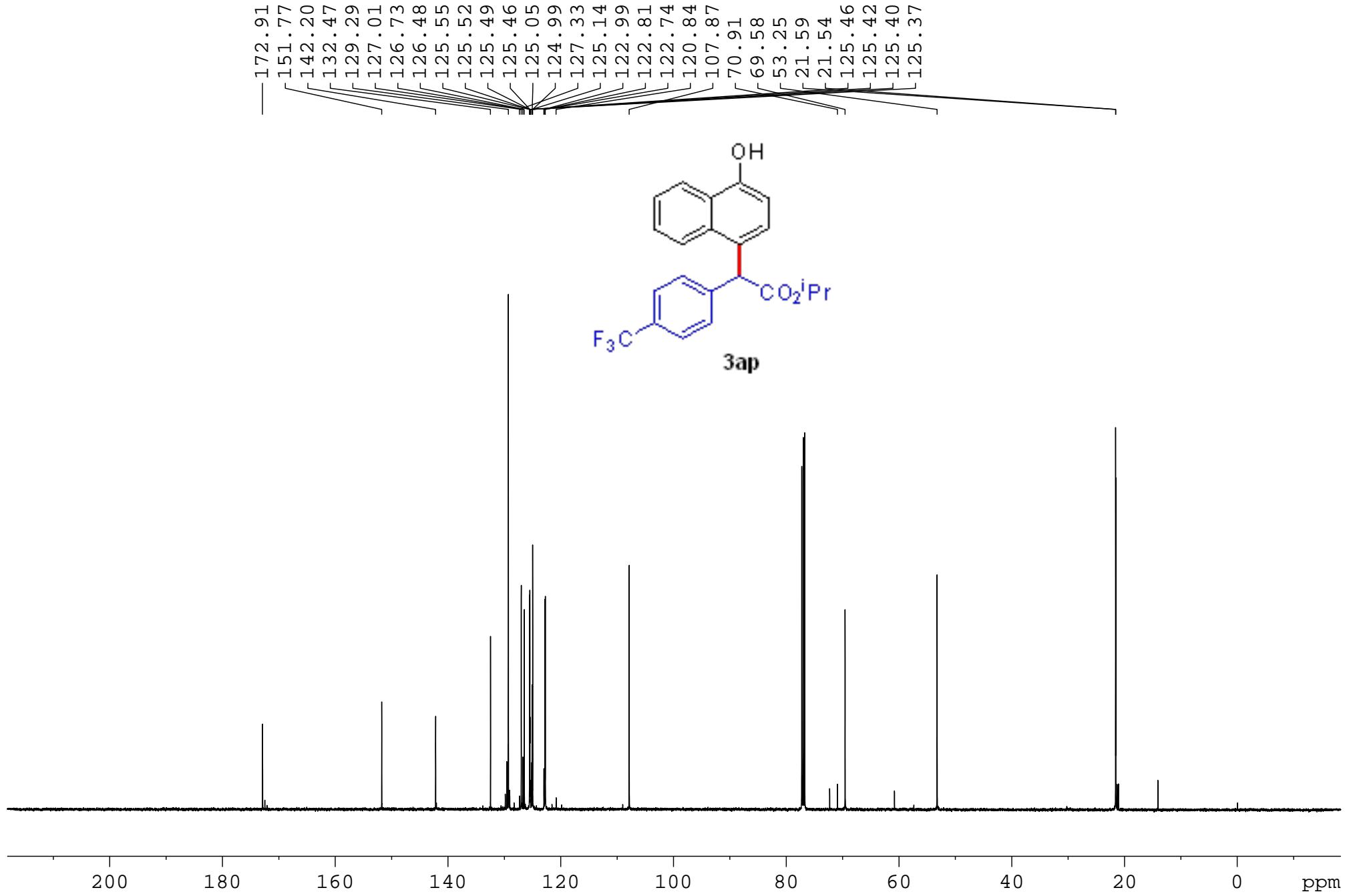
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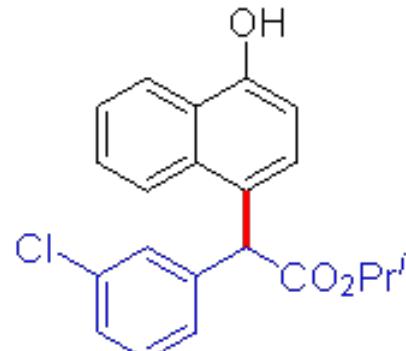




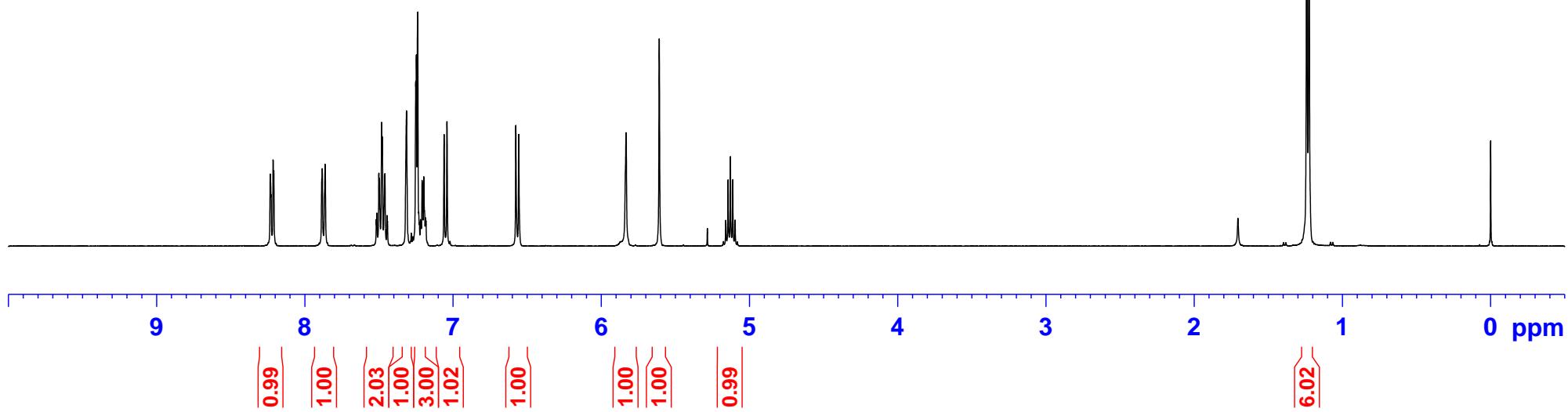
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5.098

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1.225



3ah



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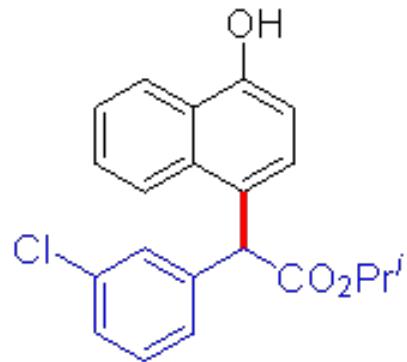
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— 151.43
— 140.29
— 134.43
— 132.56
— 129.76
— 129.08
— 127.50
— 127.13
— 127.01
— 126.38
— 126.08
— 125.03
— 124.95
— 122.92
— 122.67
— 107.95

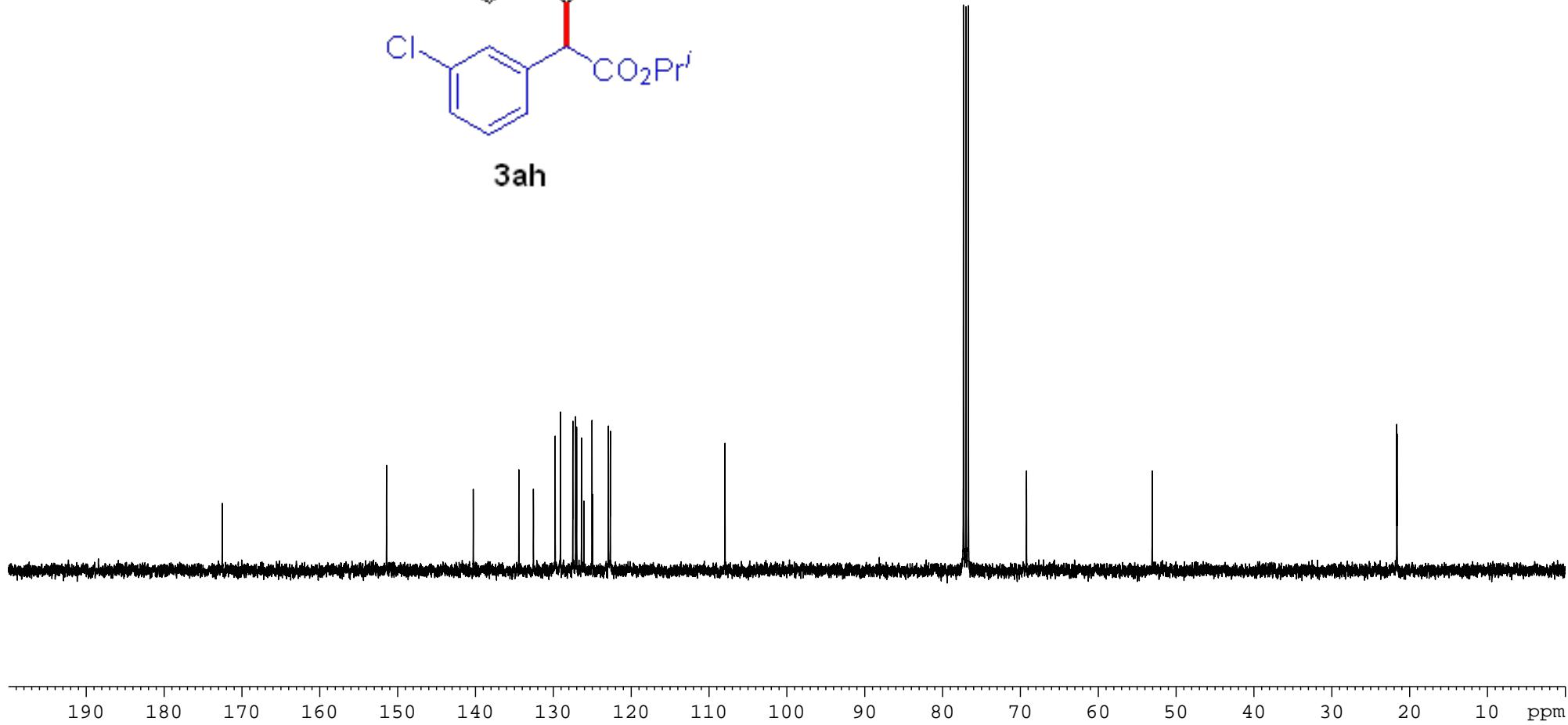
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— 77.00
— 76.68
— 69.22

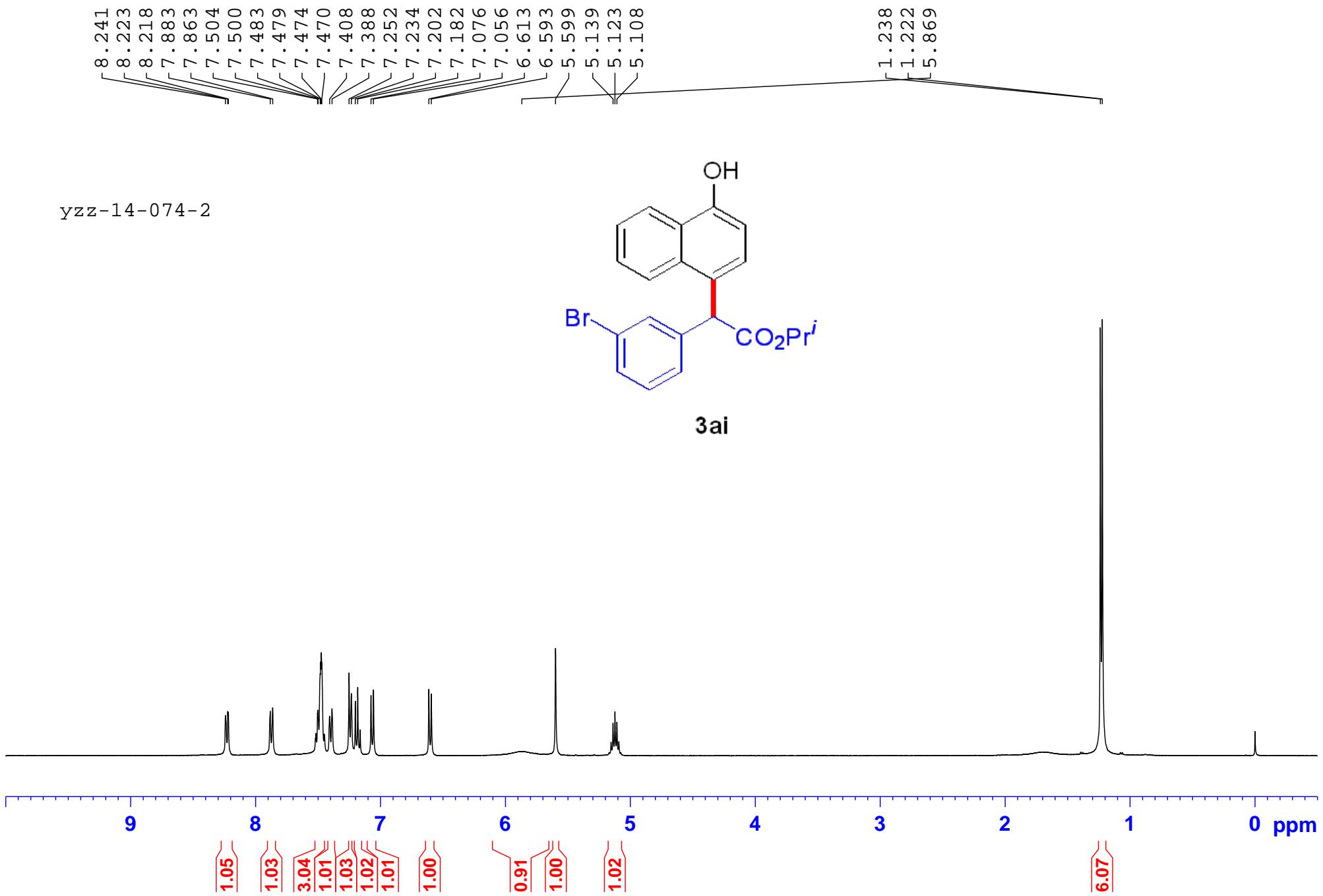
— 53.06

— 21.66
— 21.60

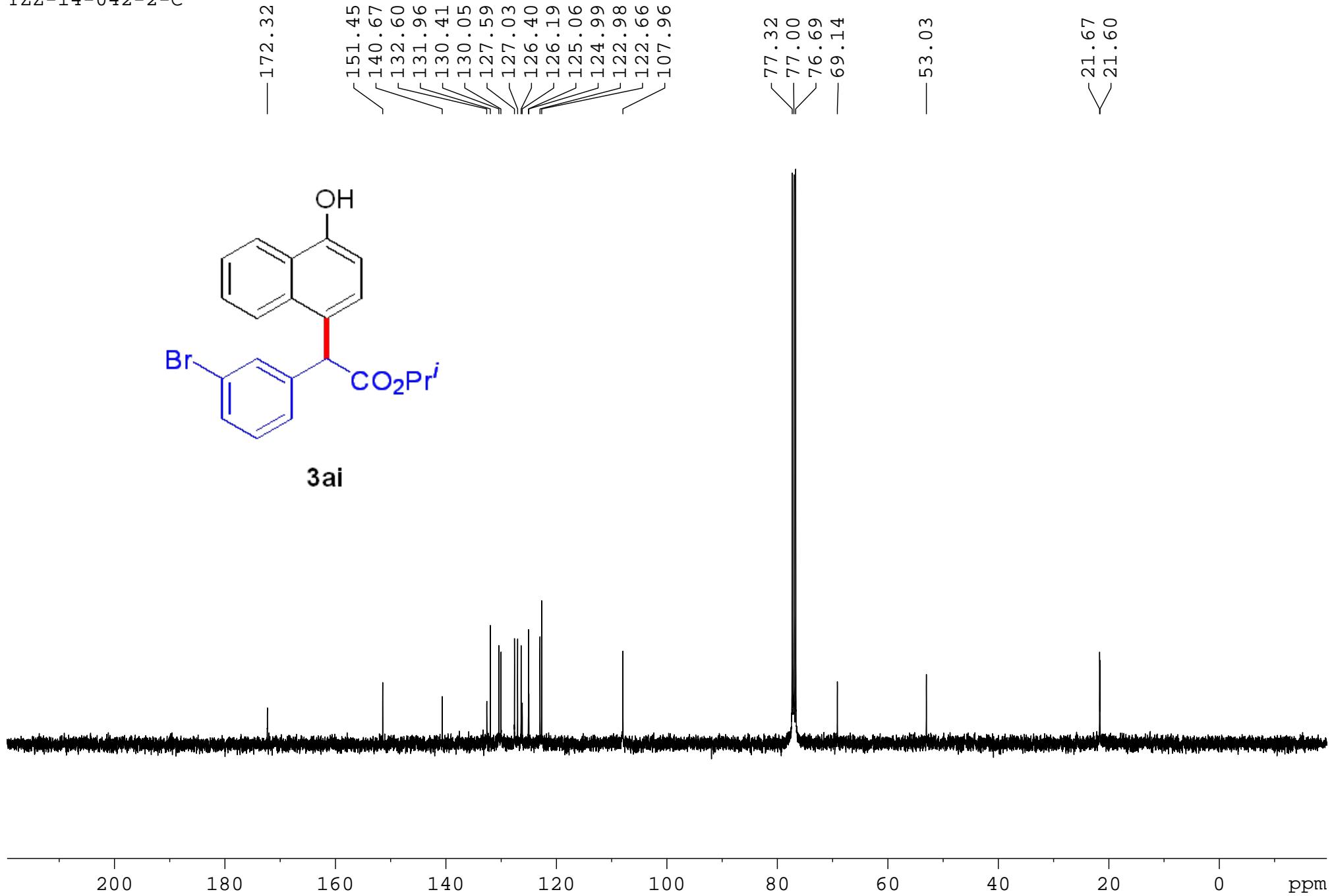


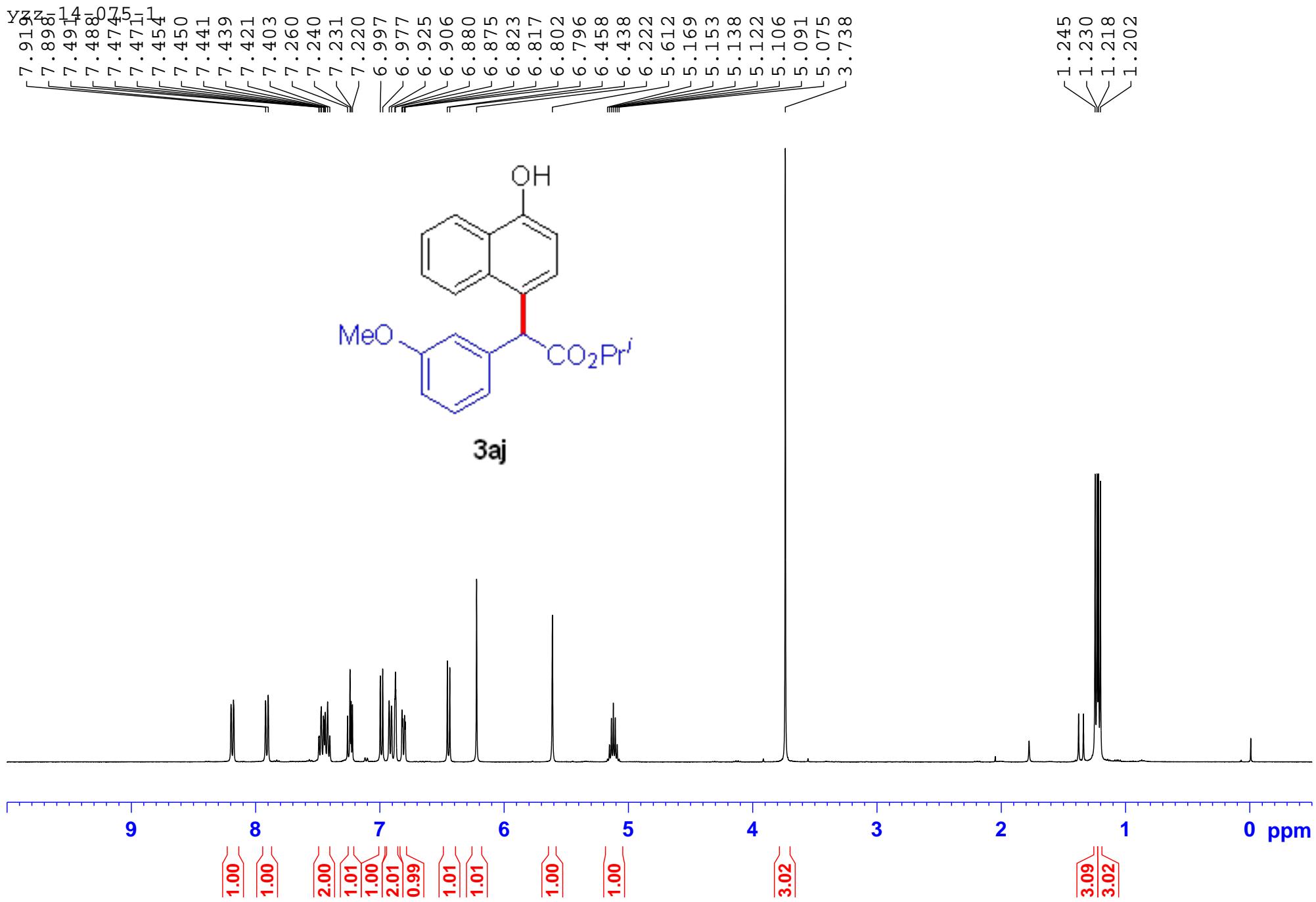
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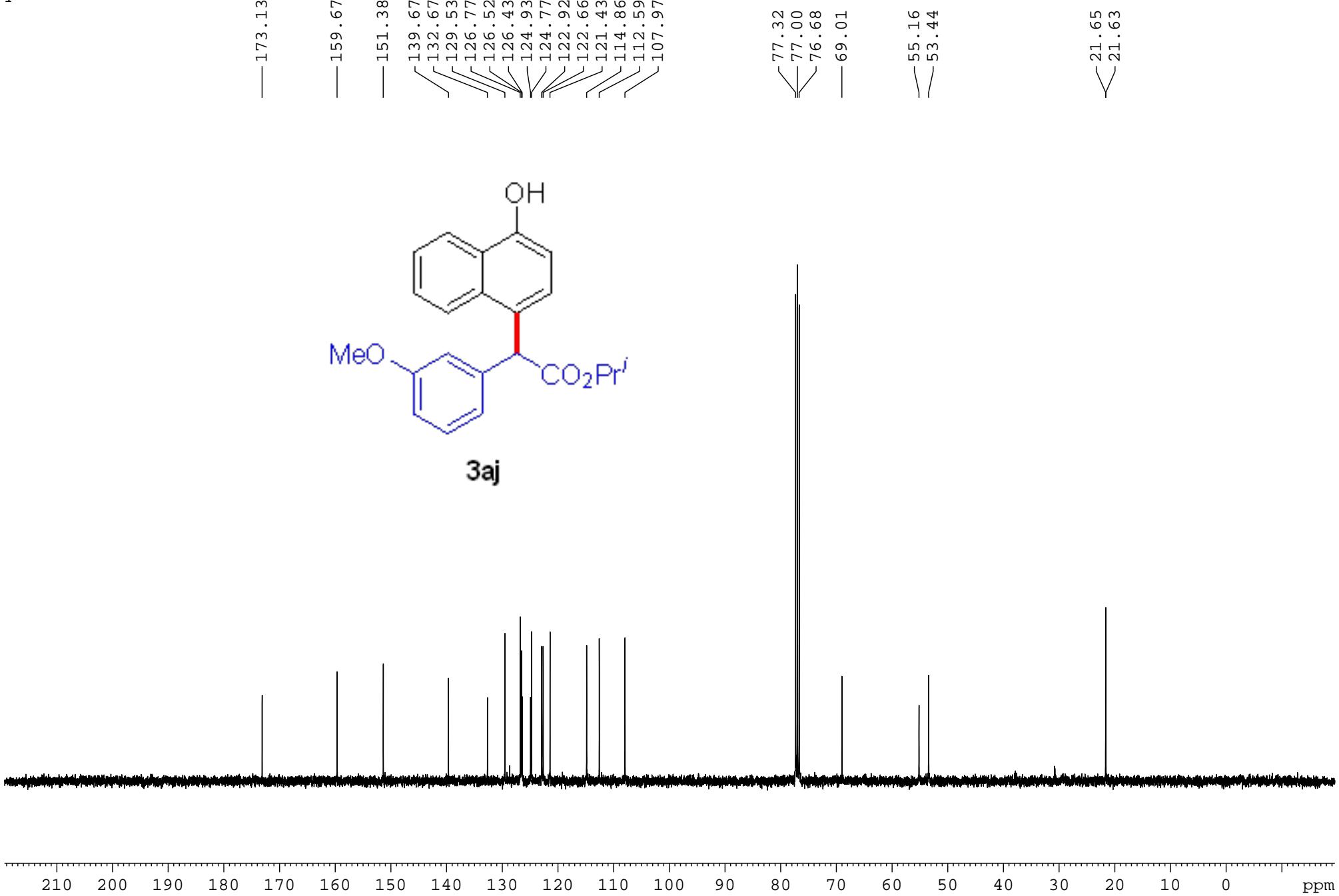


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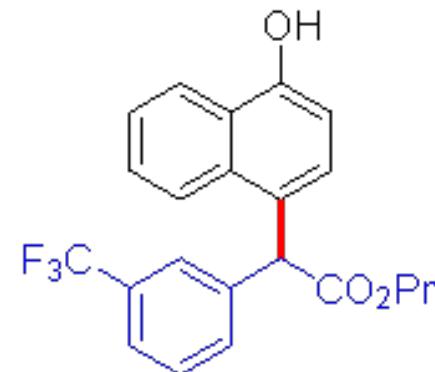
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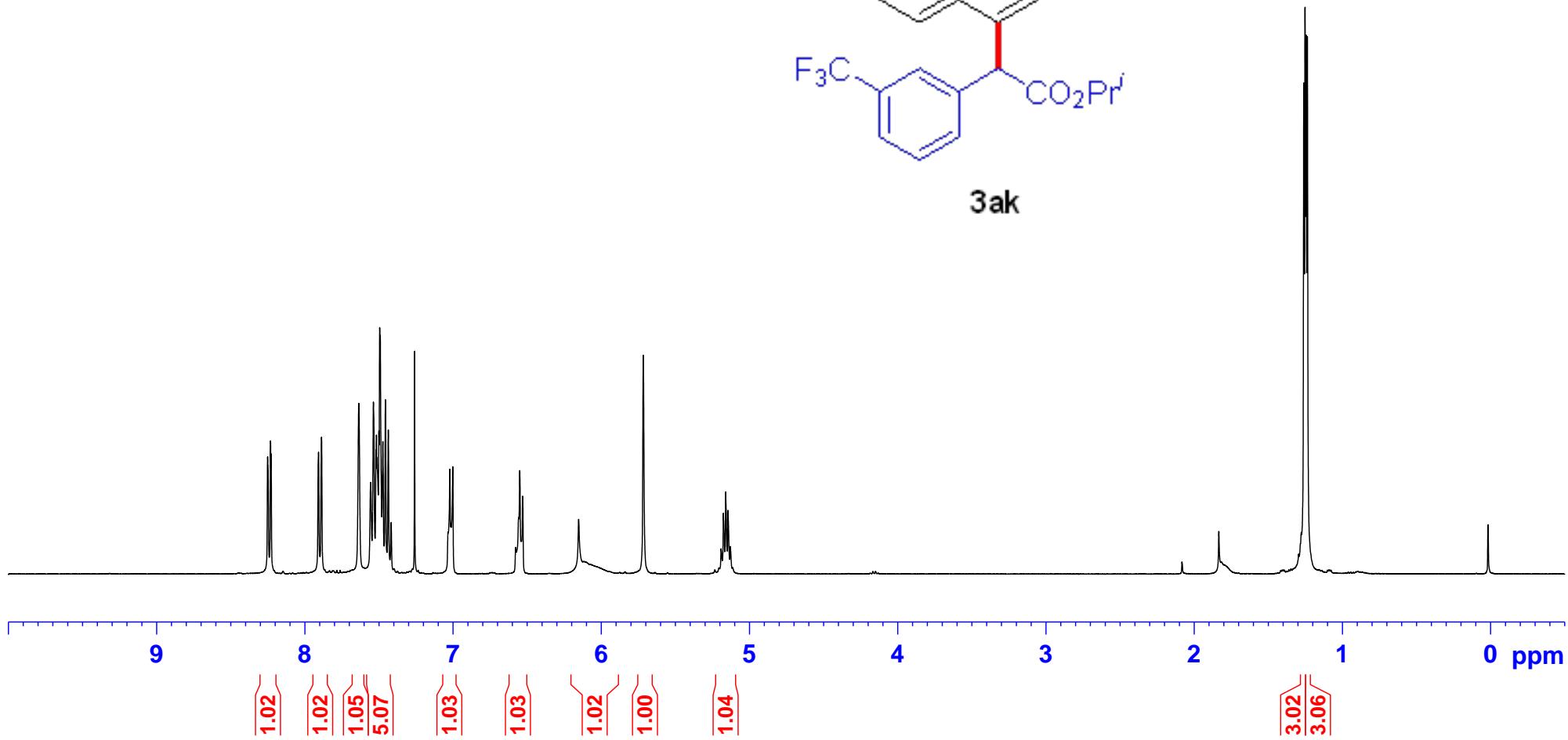
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7.260
7.035
7.023
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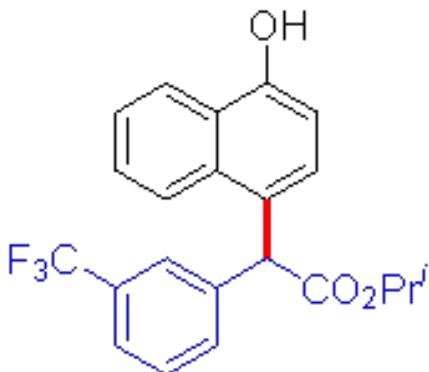
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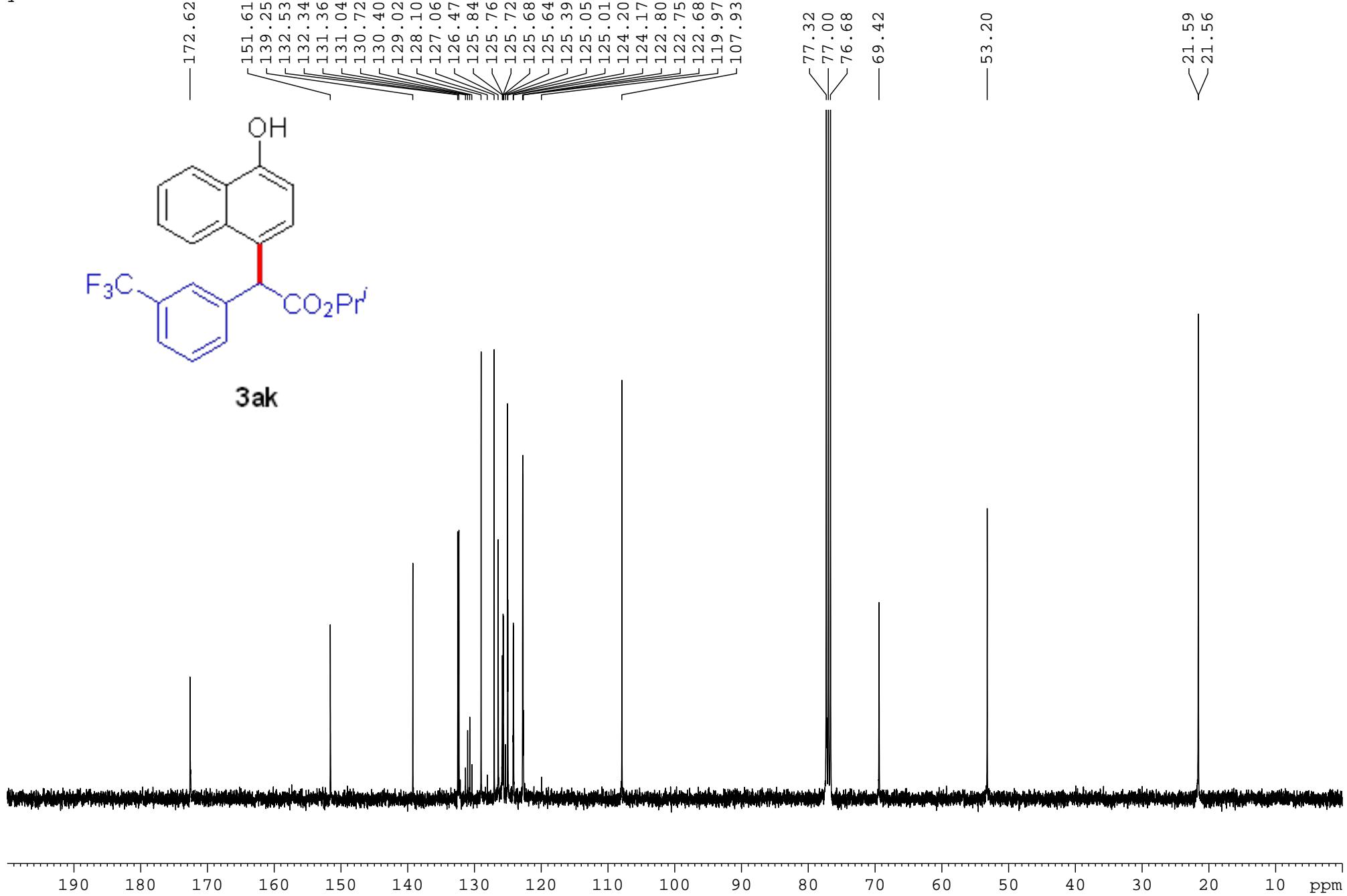
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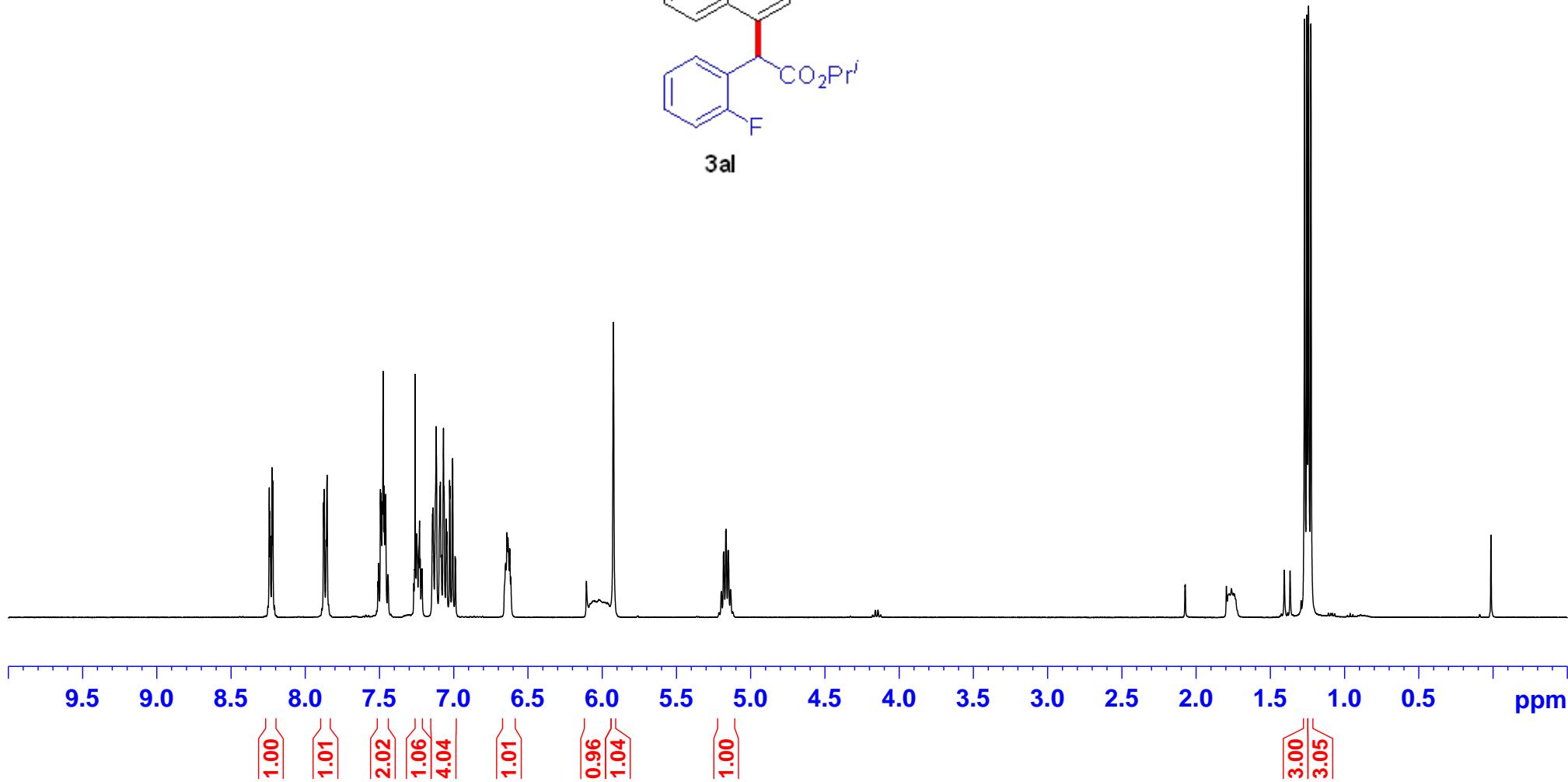
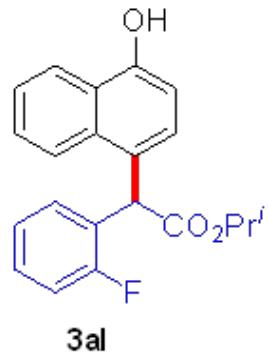
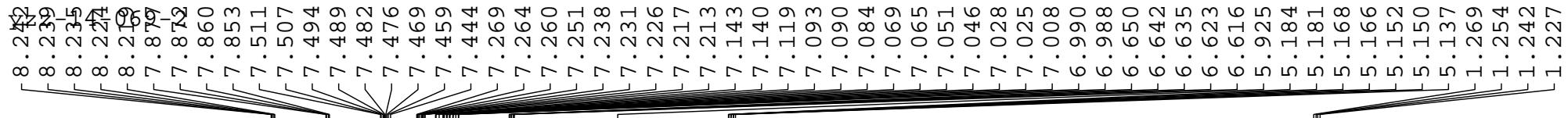


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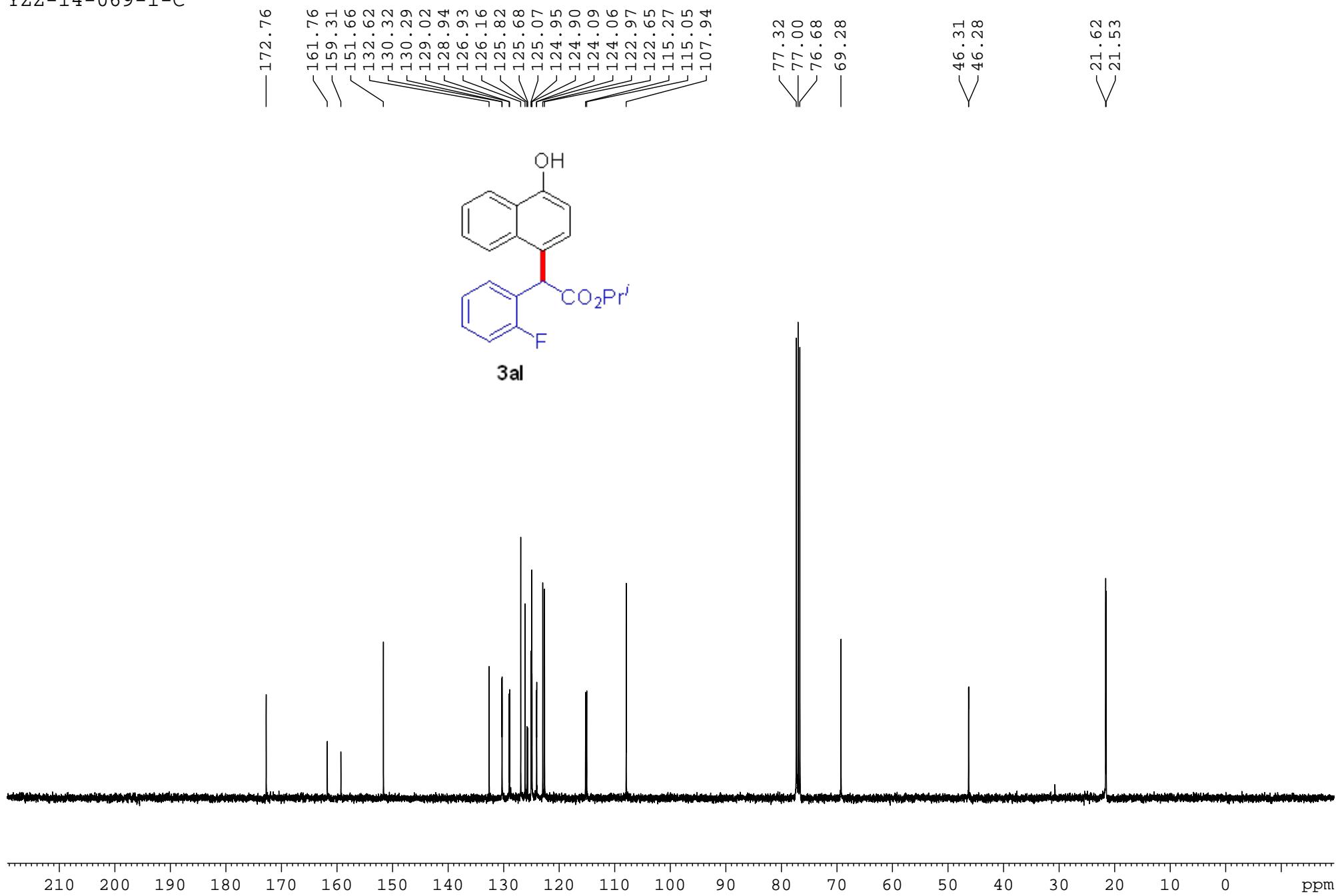


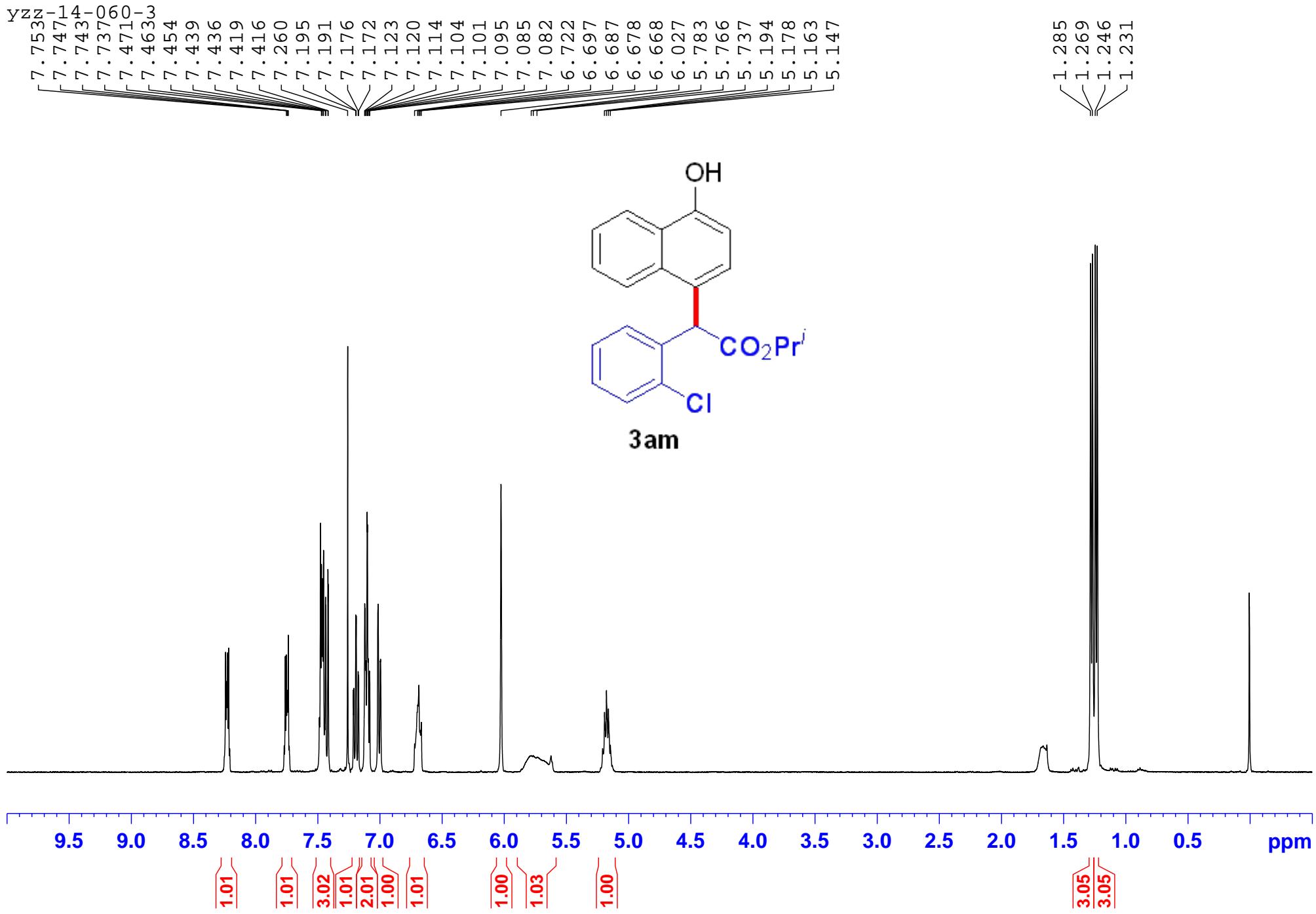
3ak



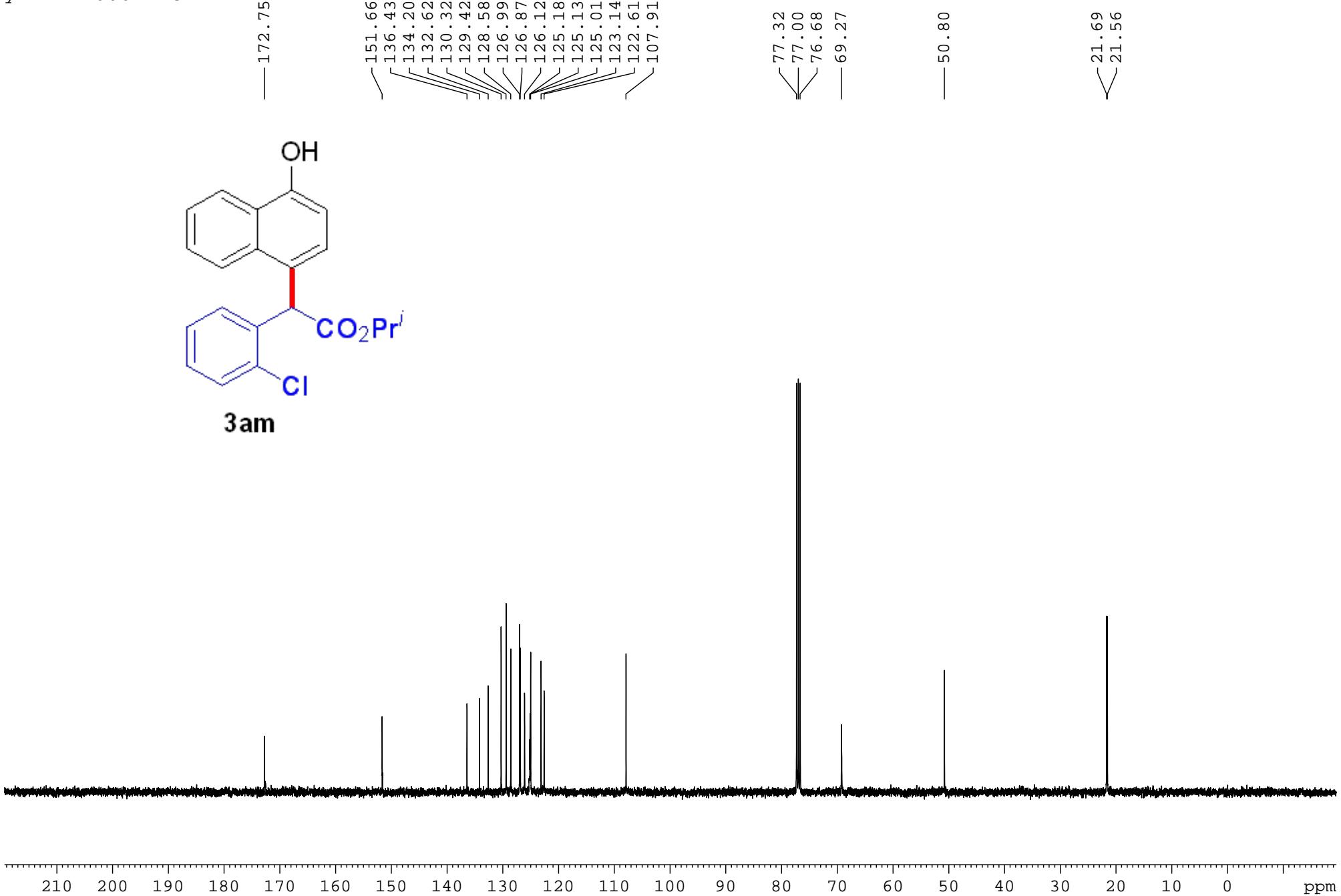


YZZ-14-069-1-C





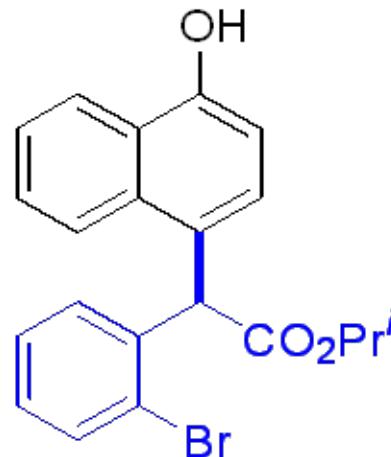
yzz-14-060-2-c



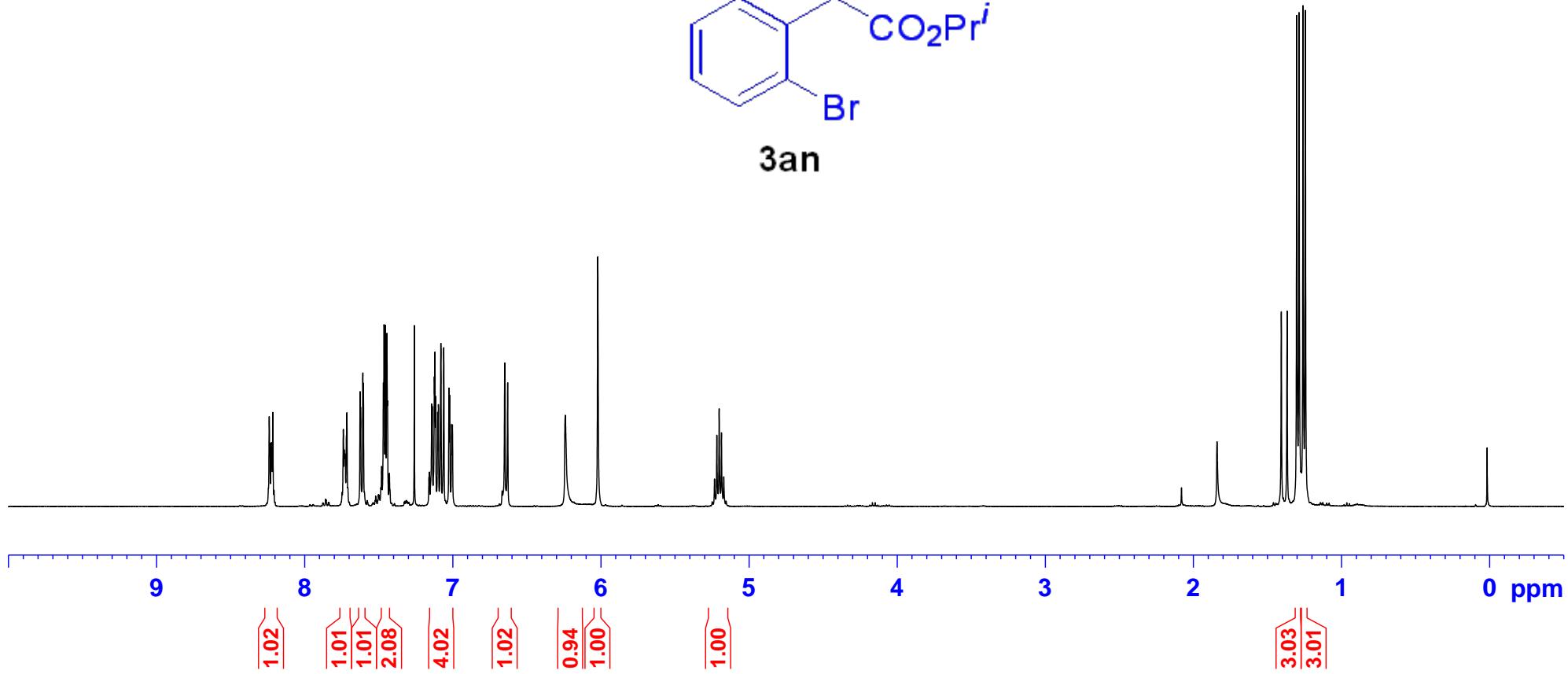
YZZ-14-070-1

7.734
7.730
7.727
7.719
7.716
7.704
7.627
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7.608
7.604
7.469
7.465
7.455
7.445
7.441
7.260
7.143
7.140
7.125
7.122
7.116
7.103
7.098
7.081
7.021
7.008
7.003
6.650
6.630
6.241
6.022
5.234
5.218
5.202
5.187
5.171

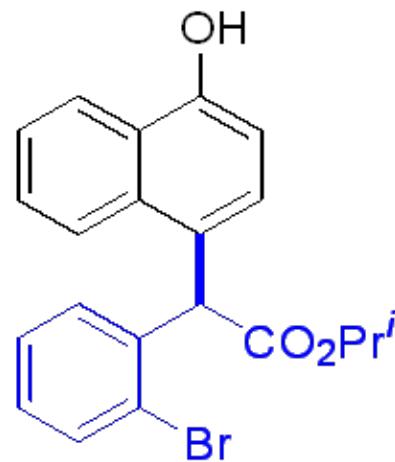
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1.288
1.261
1.245



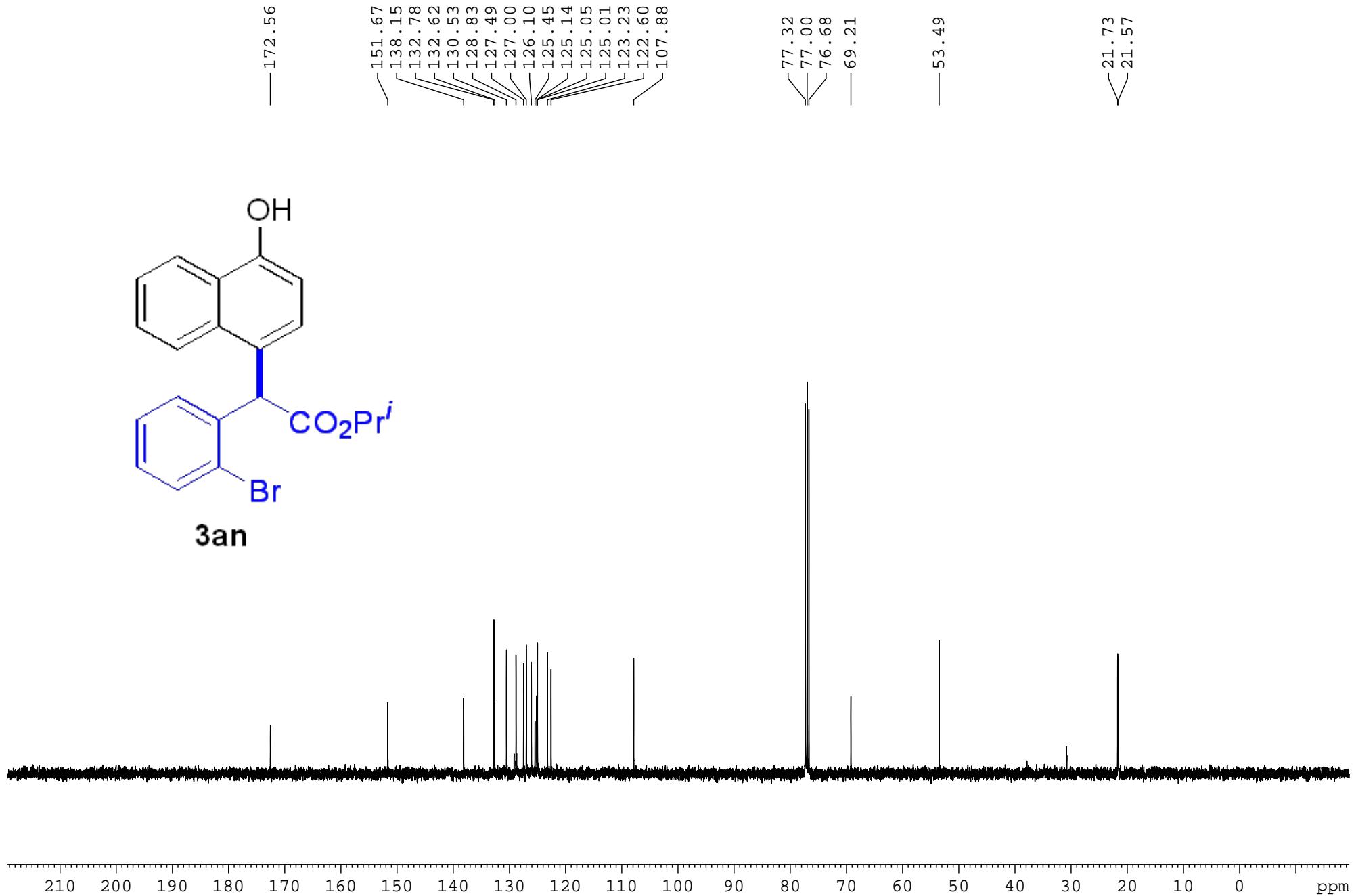
3an

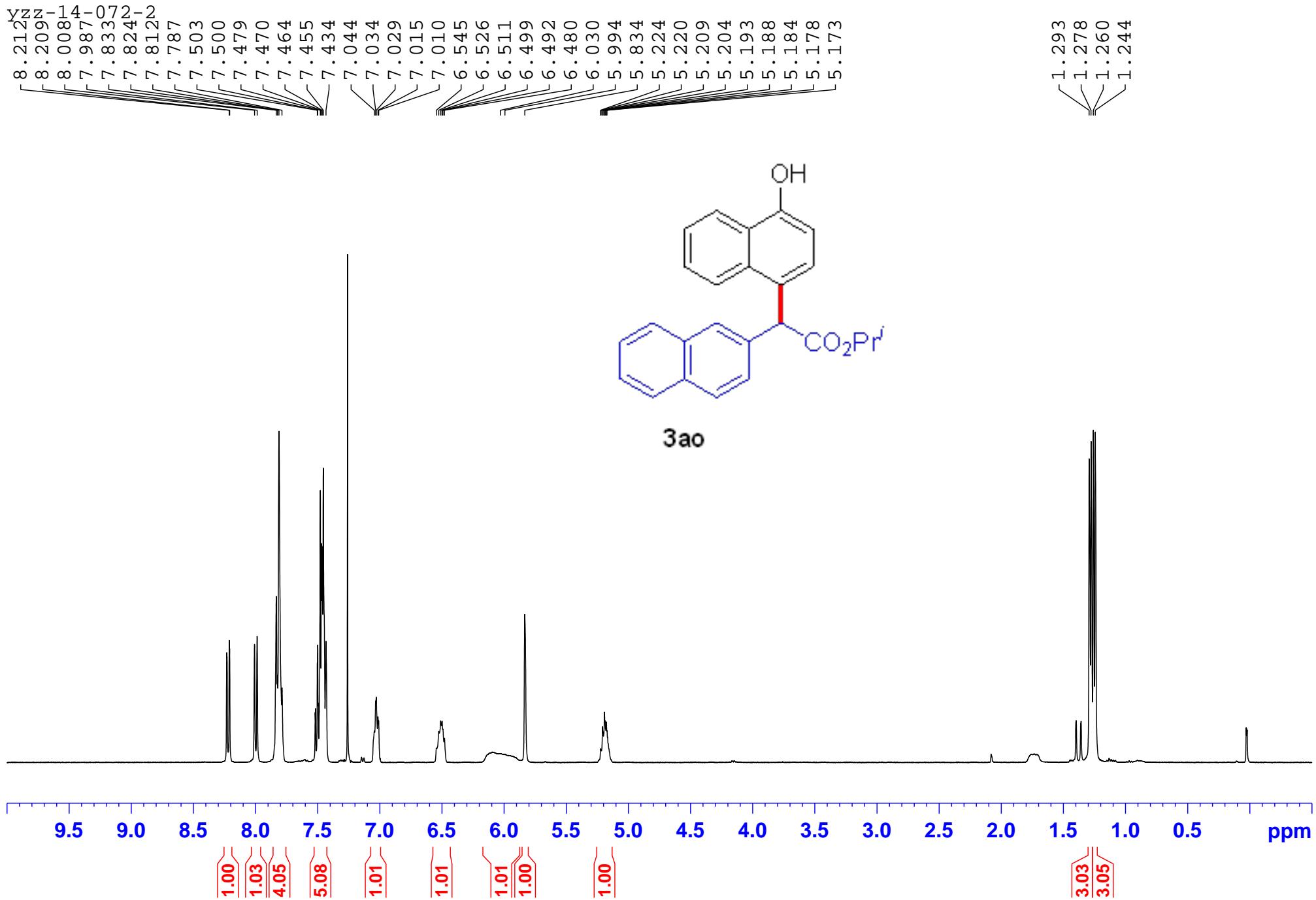


YZZ-14-070-1-C



3an

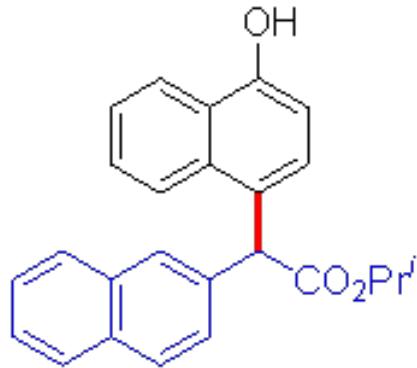




yzz-14-072-1-c

— 173.43

151.46
135.60
133.38
132.69
132.55
128.25
127.93
127.58
127.55
127.19
126.82
126.79
126.38
126.10
125.92
124.99
124.78
122.90
122.70
108.03

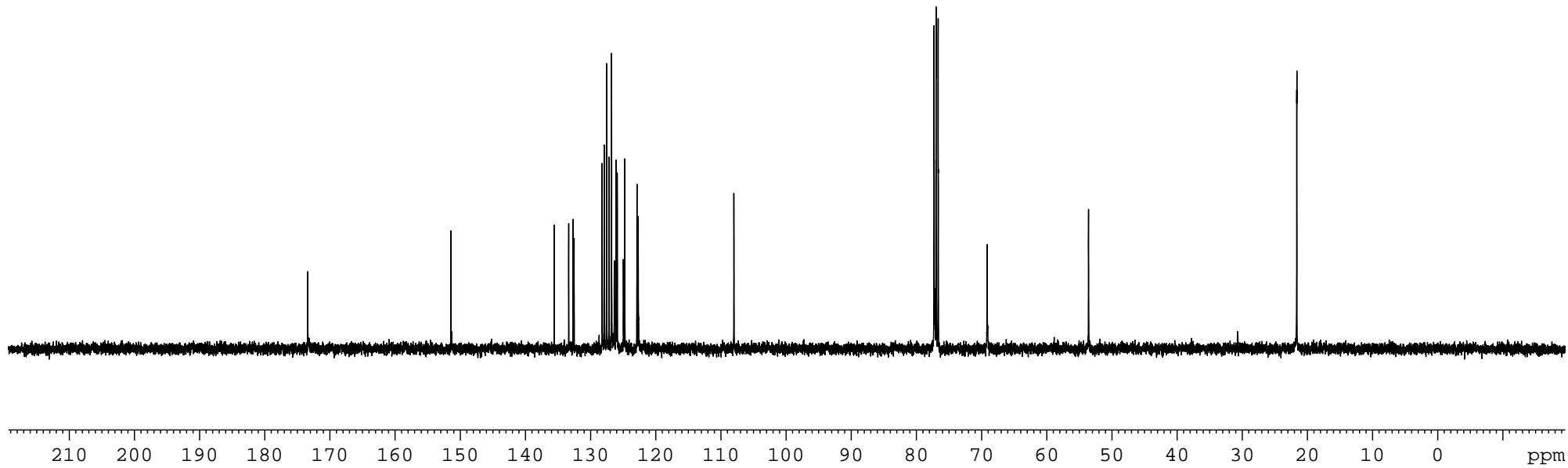


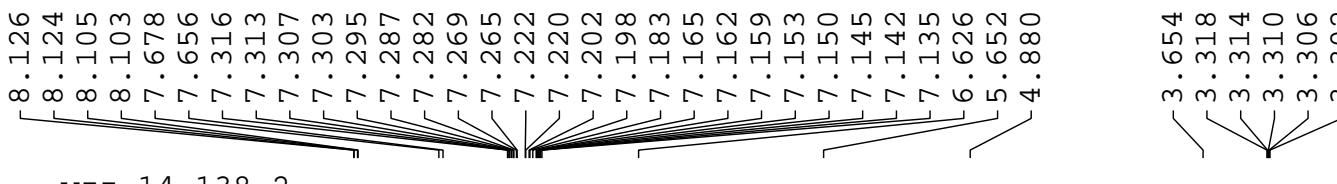
3ao

— 53.61

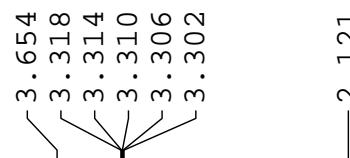
77.32
77.00
76.68
— 69.17

21.65
21.64

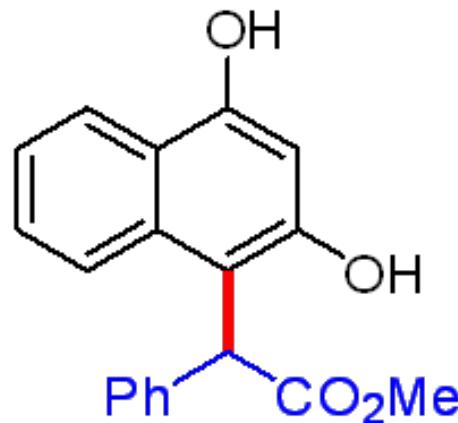




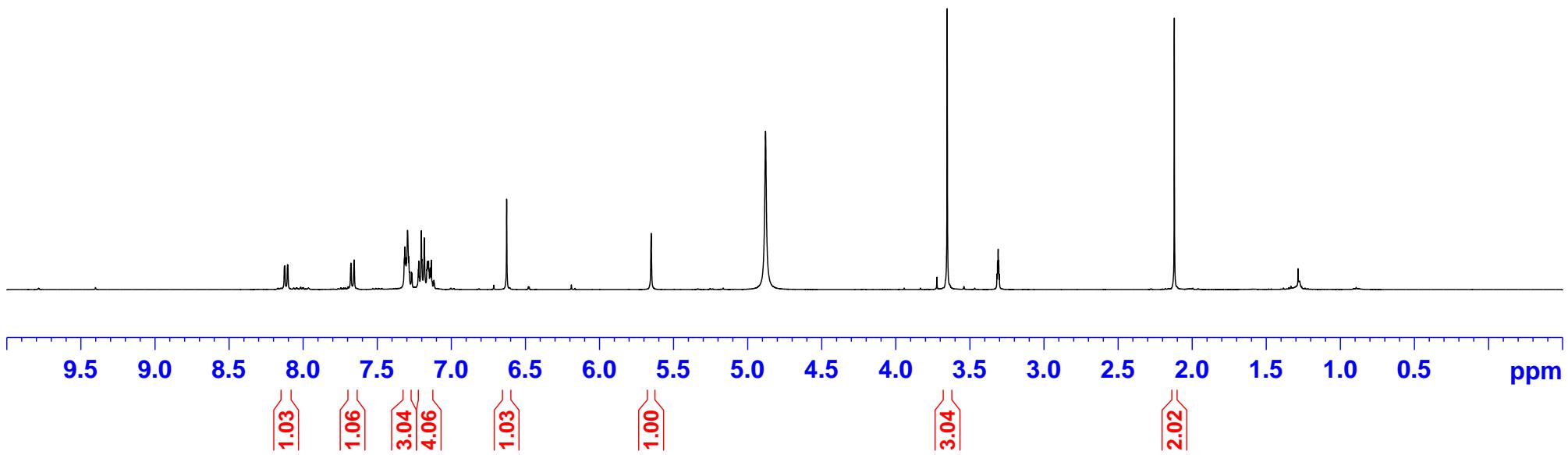
yzz-14-138-2



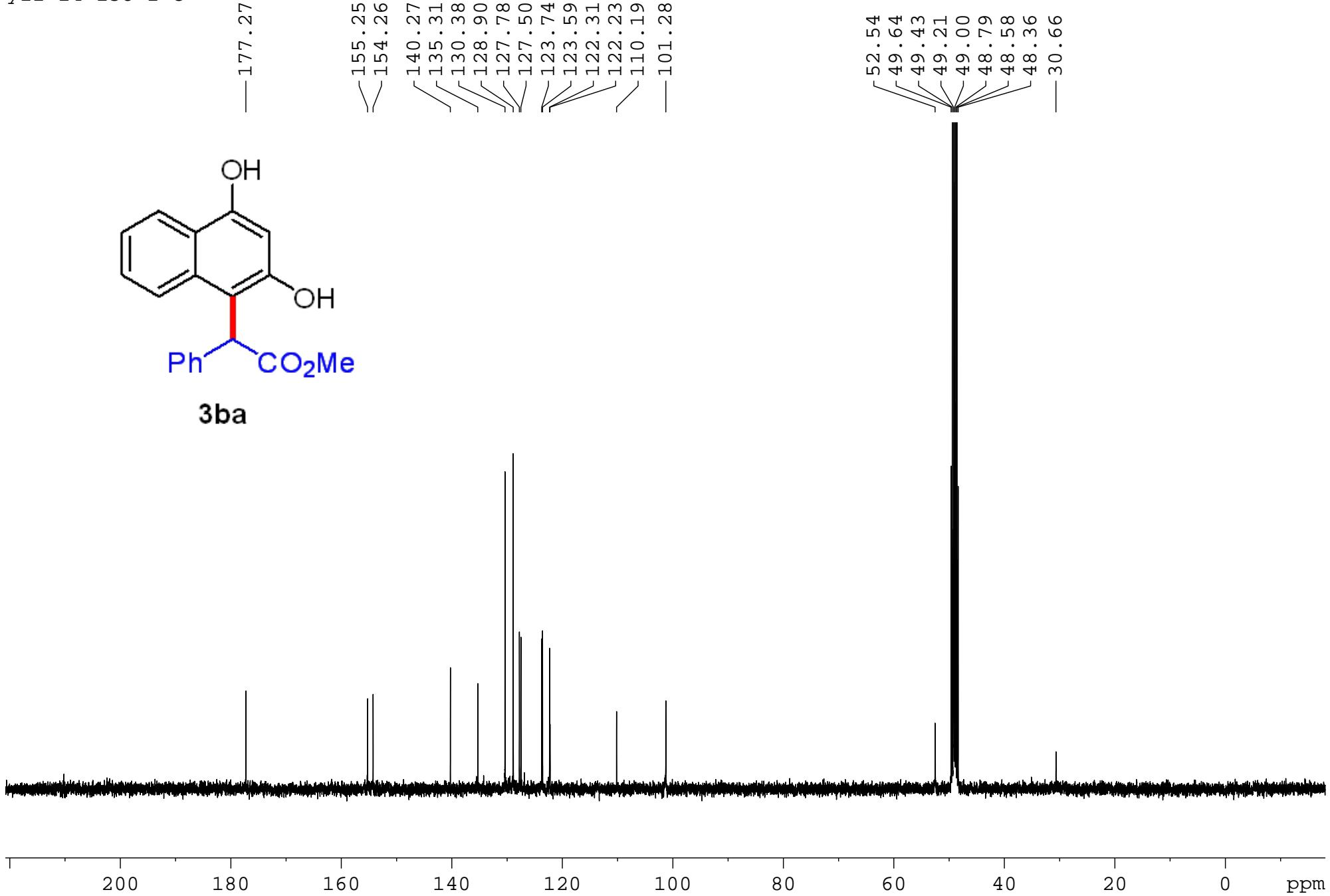
— 2.121



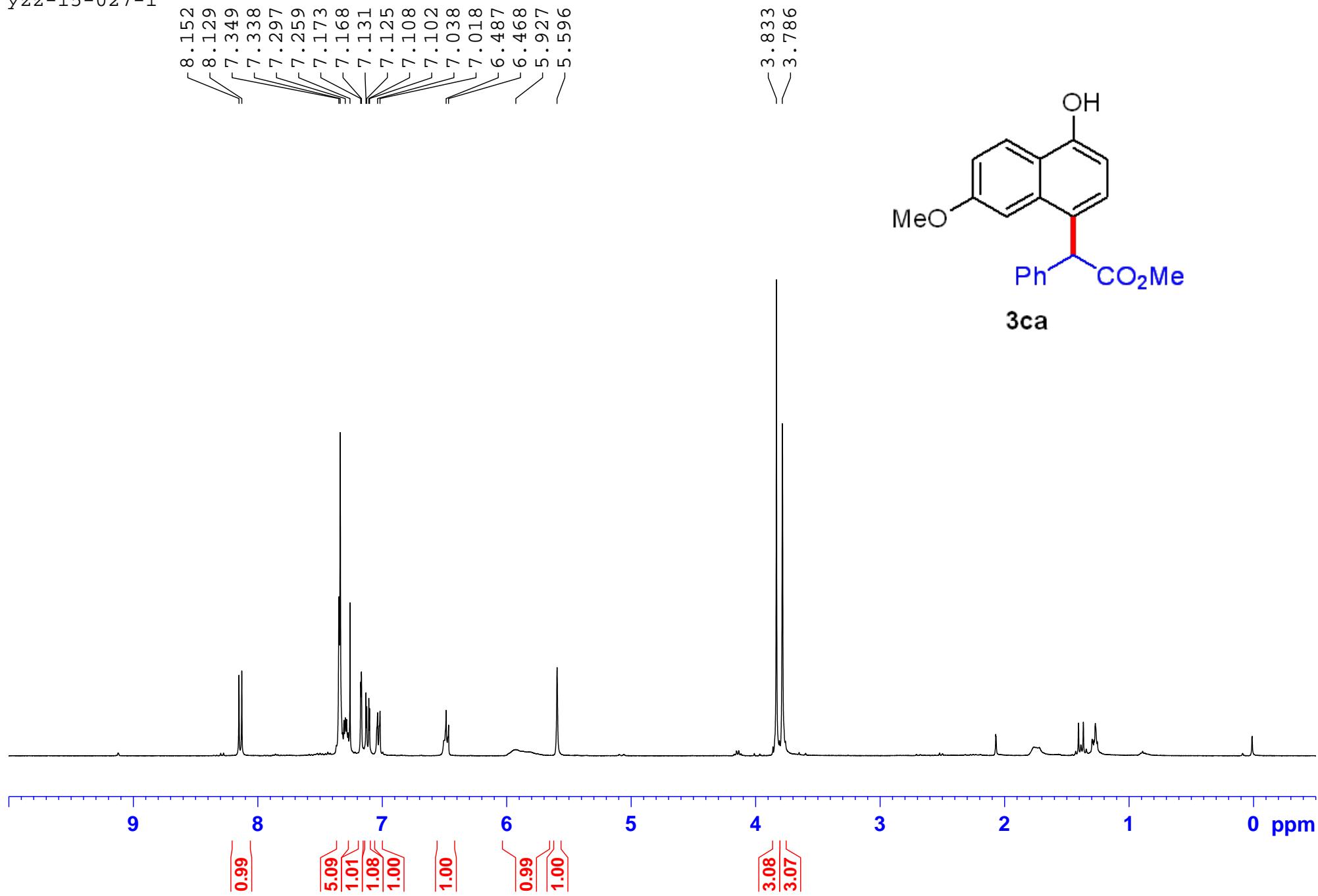
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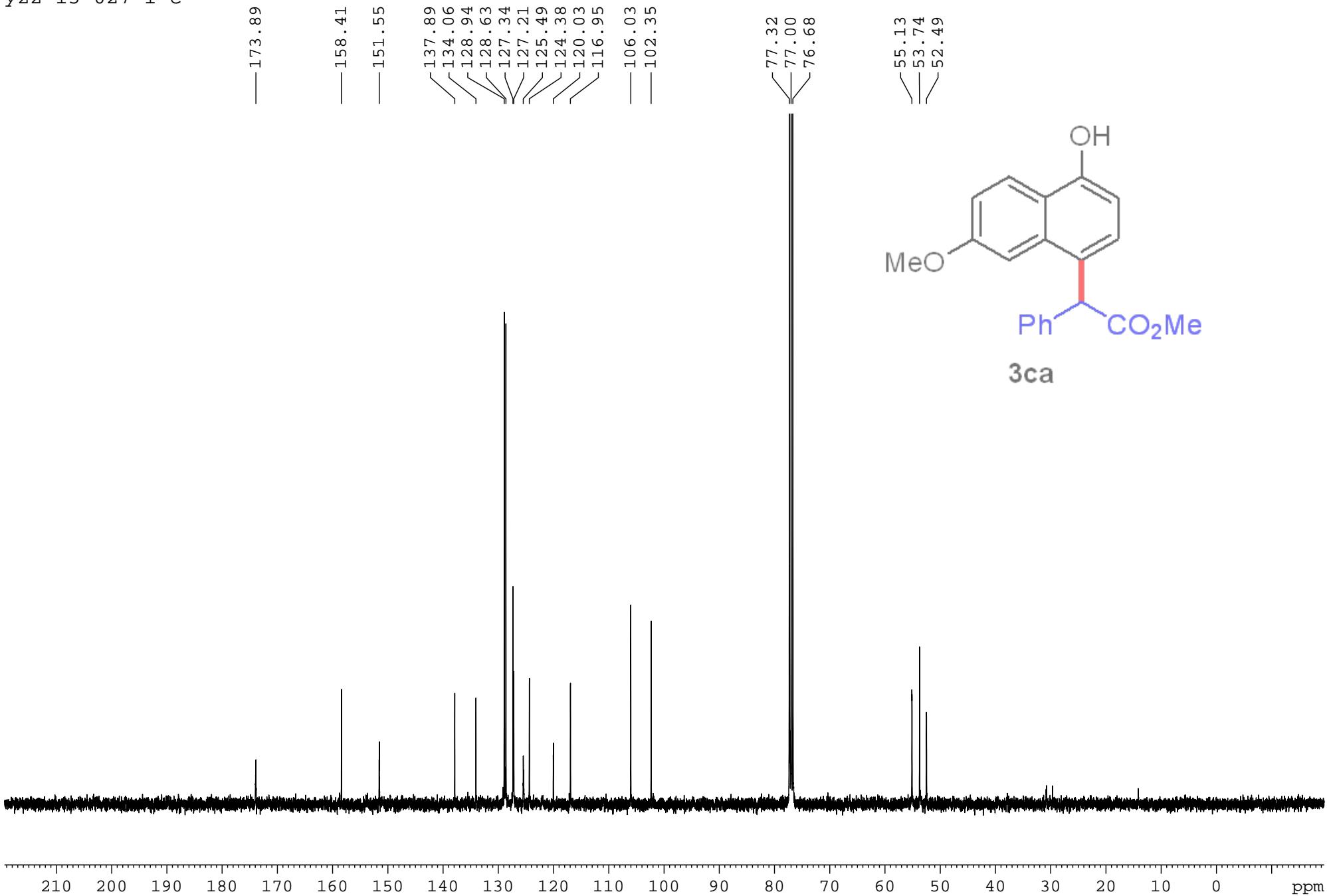
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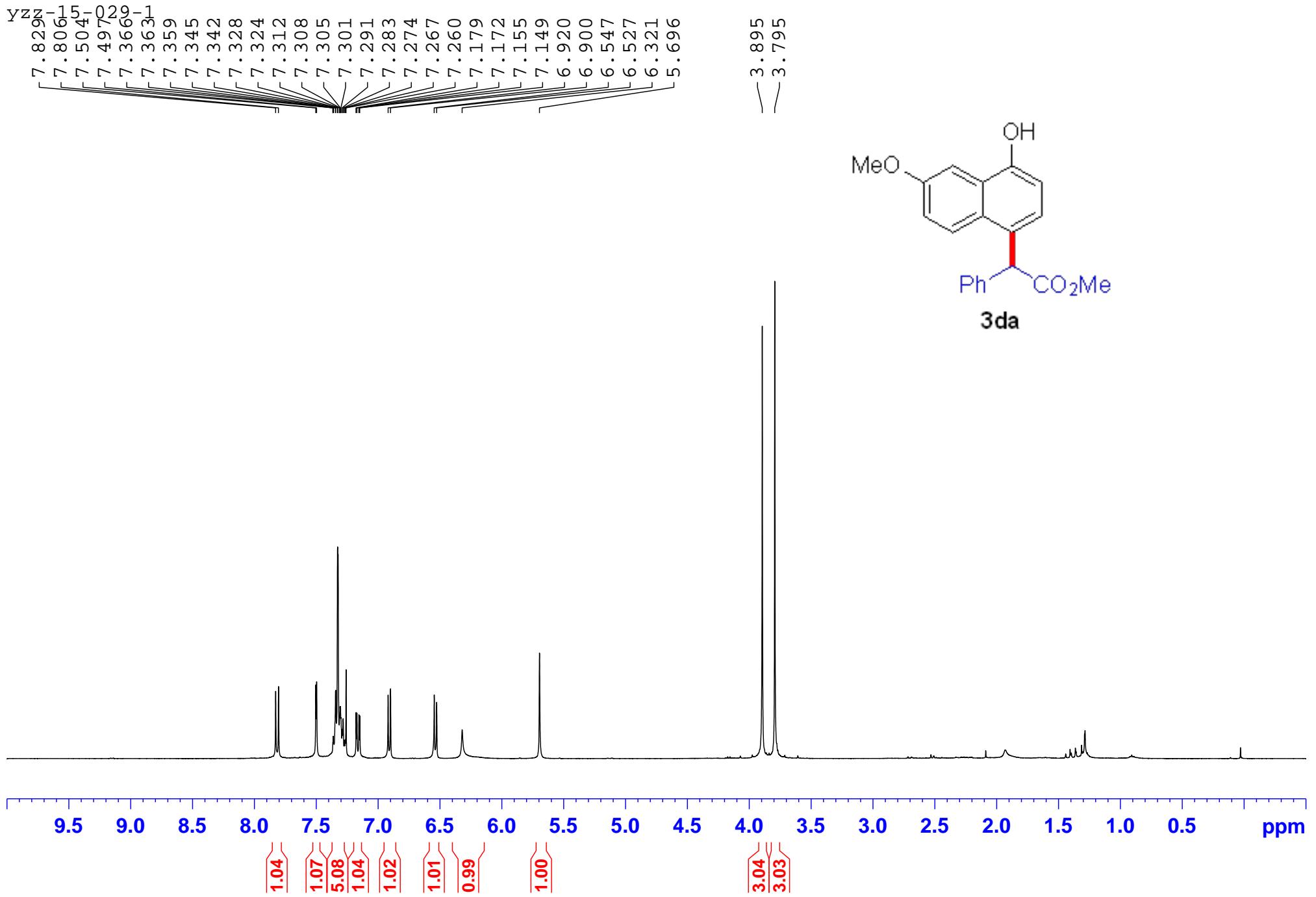


yzz-15-027-1

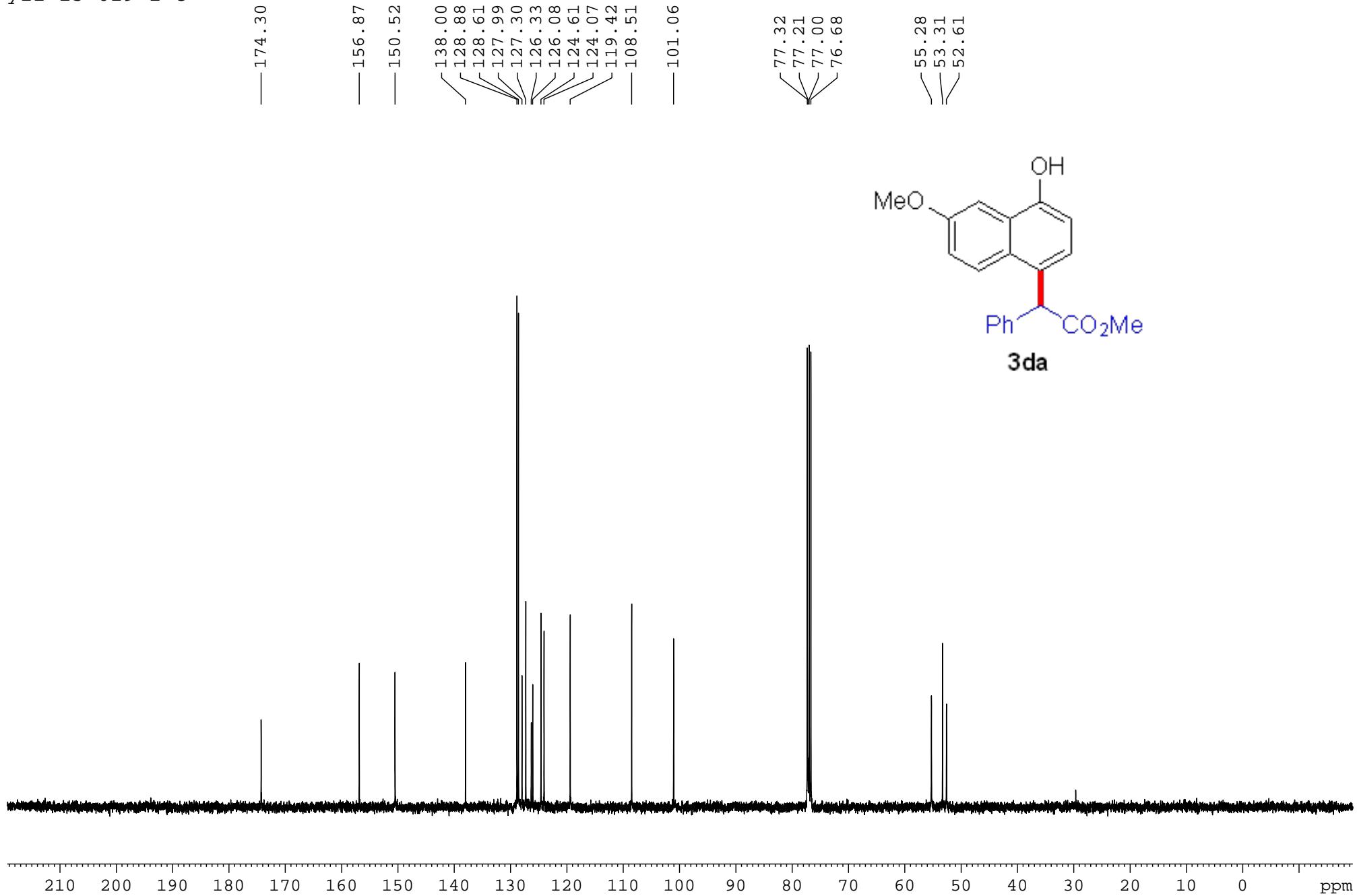


yzz-15-027-1-c

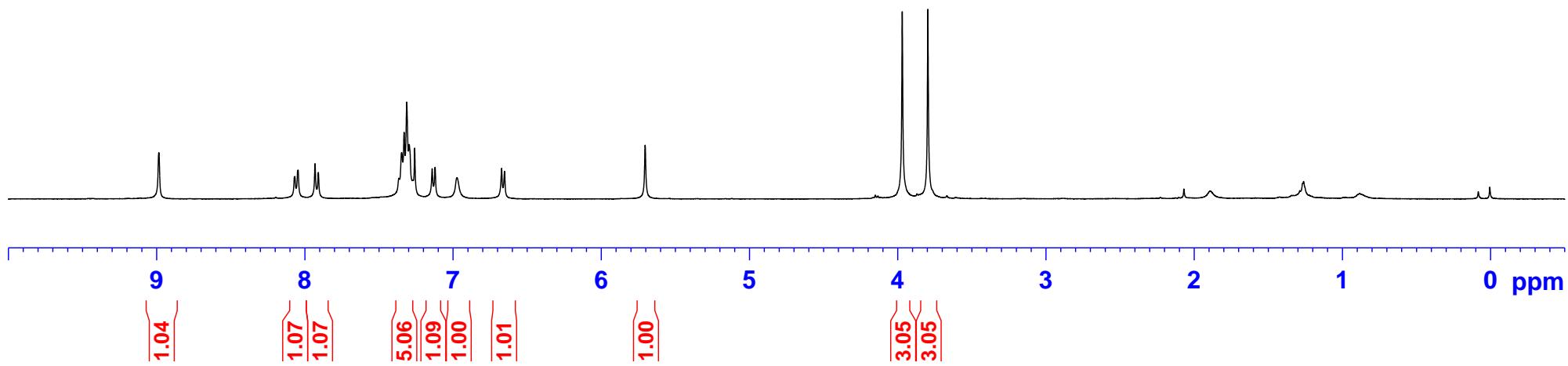
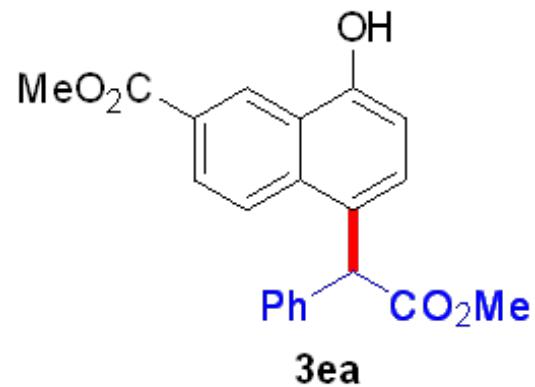
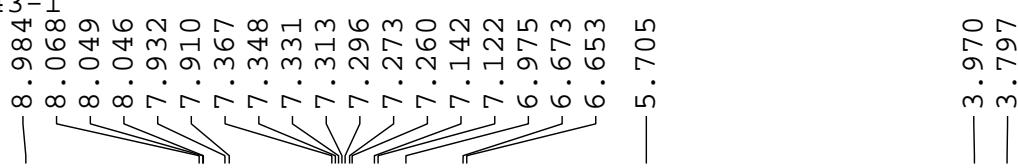




yzz-15-029-1-c



yzz-15-043-1



yzz-15-043-1-c

— 174.09

— 167.48

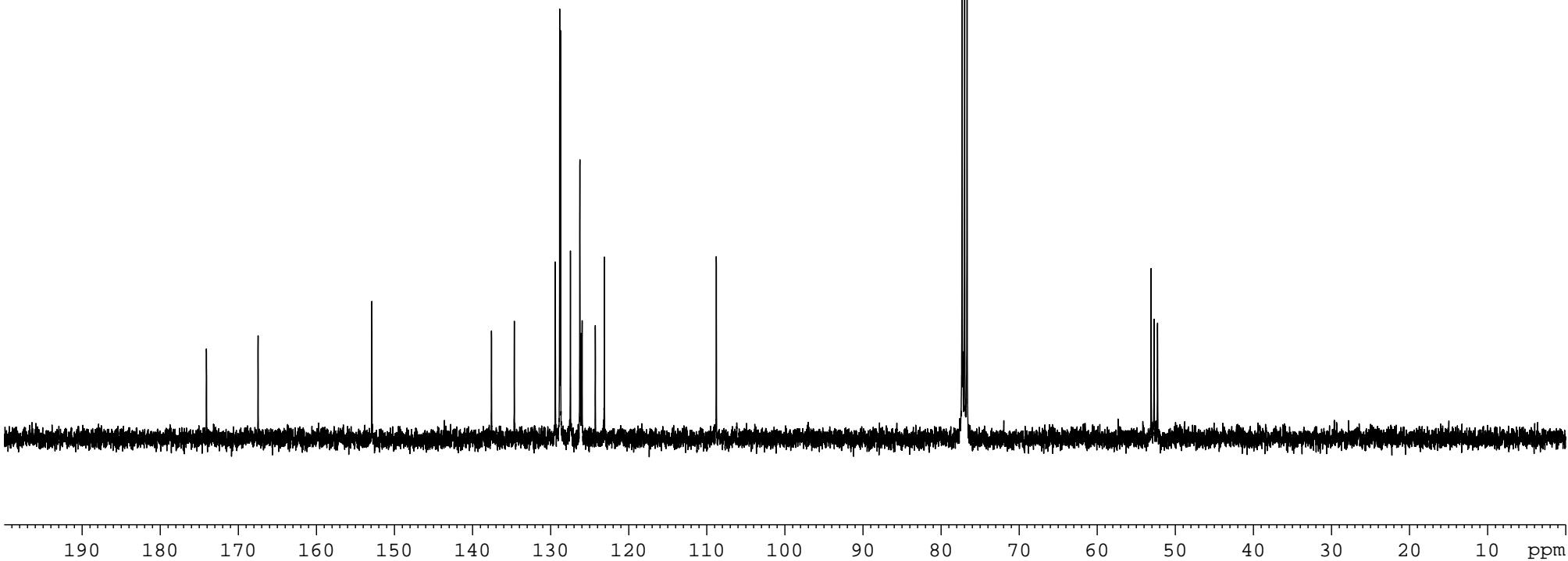
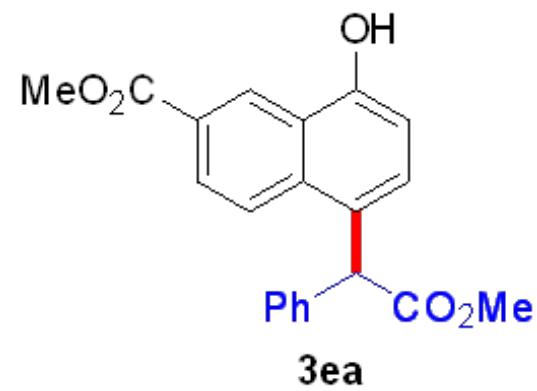
— 152.90

137.58
134.66
129.41
128.87
128.75
127.48
126.27
126.10
125.99
124.29
123.13

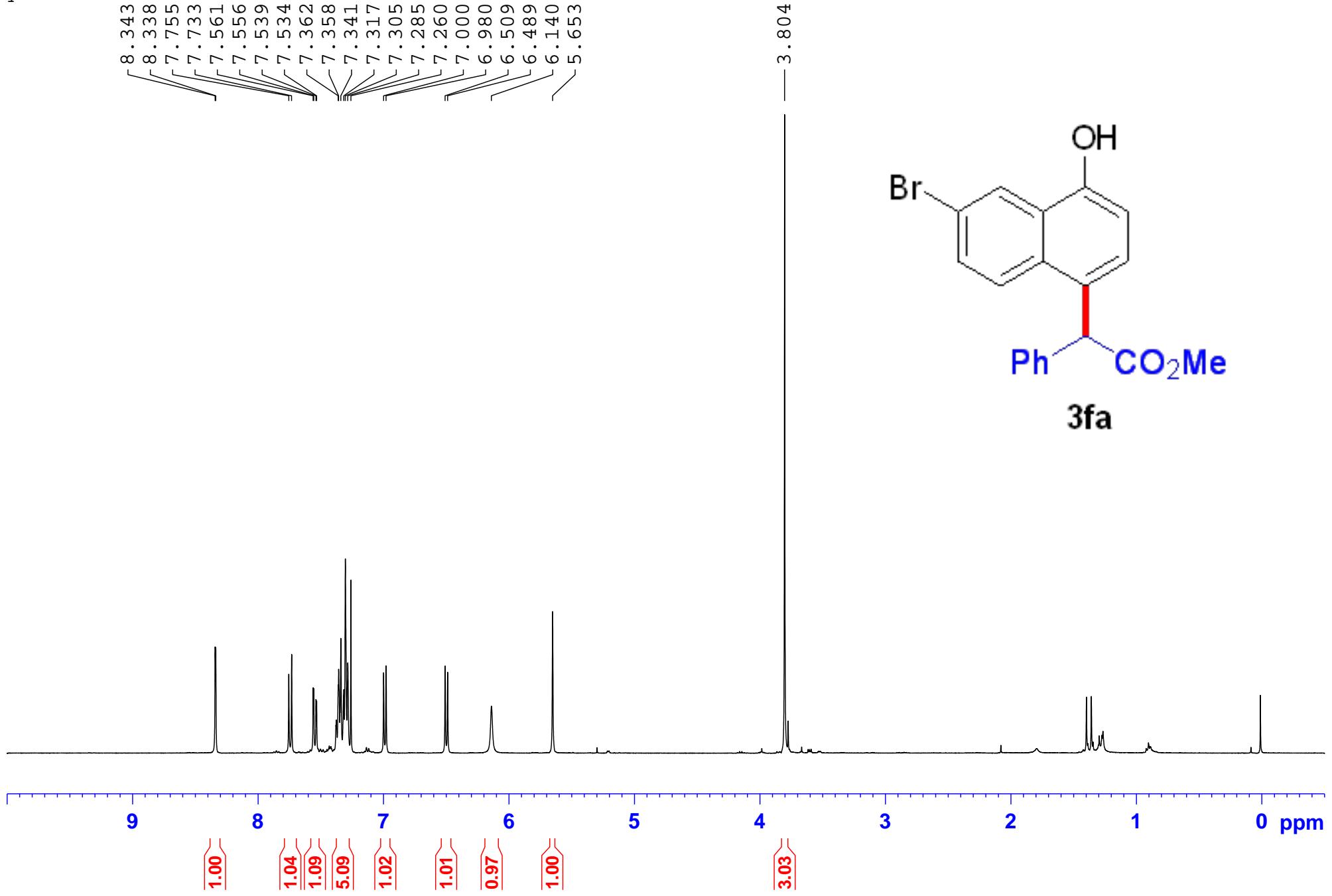
— 108.80

77.32
77.00
76.68

53.11
52.72
52.29



yzz-14-137-1



yzz-14-137-1-**16**

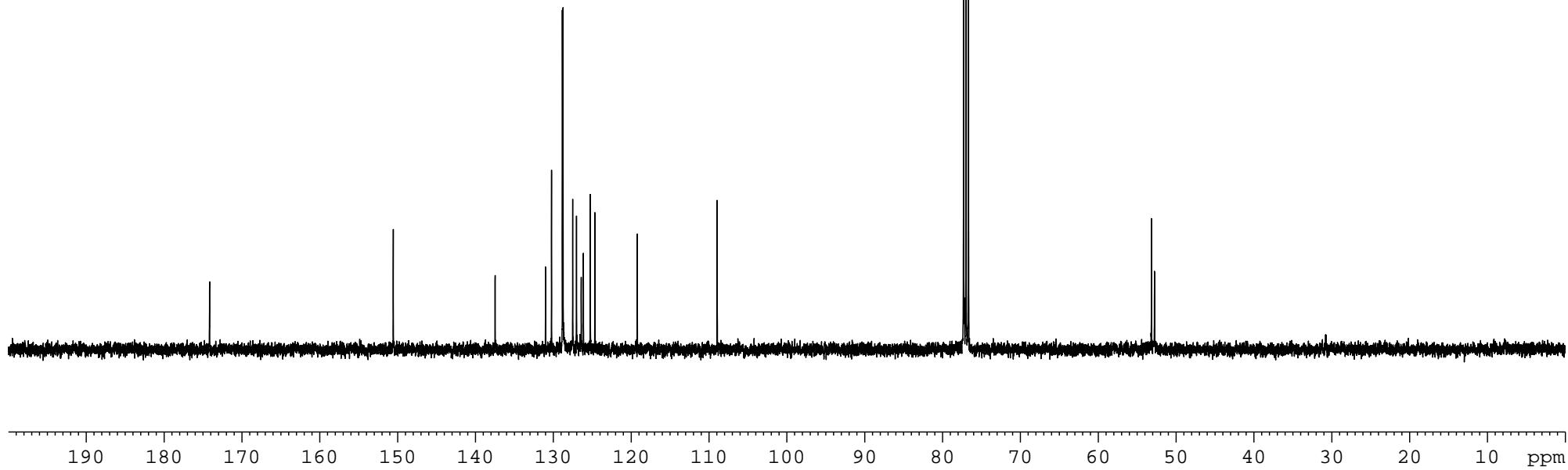
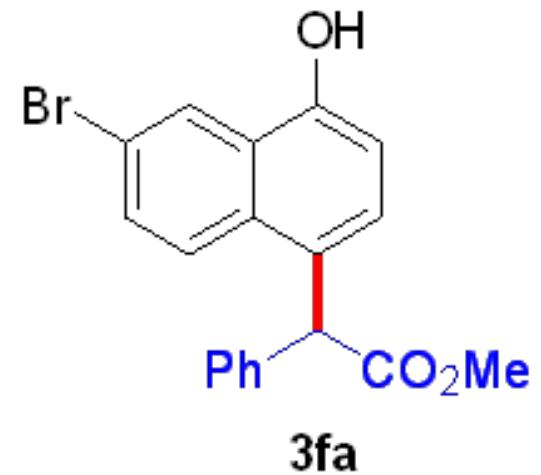
— 174.16

— 150.57

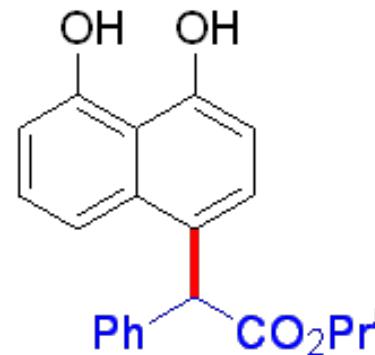
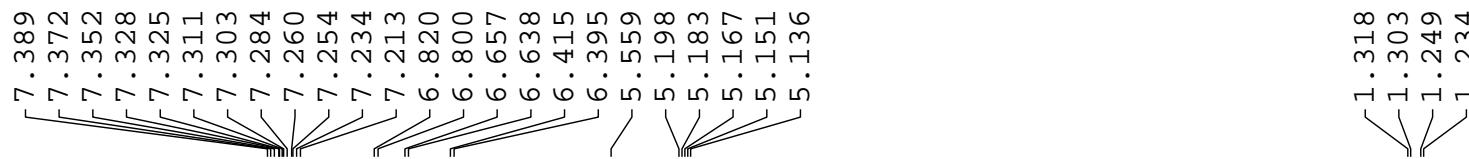
137.49
131.02
130.24
128.85
128.76
127.52
127.04
126.43
126.14
125.27
124.67
119.22
108.96

77.32
77.00
76.68

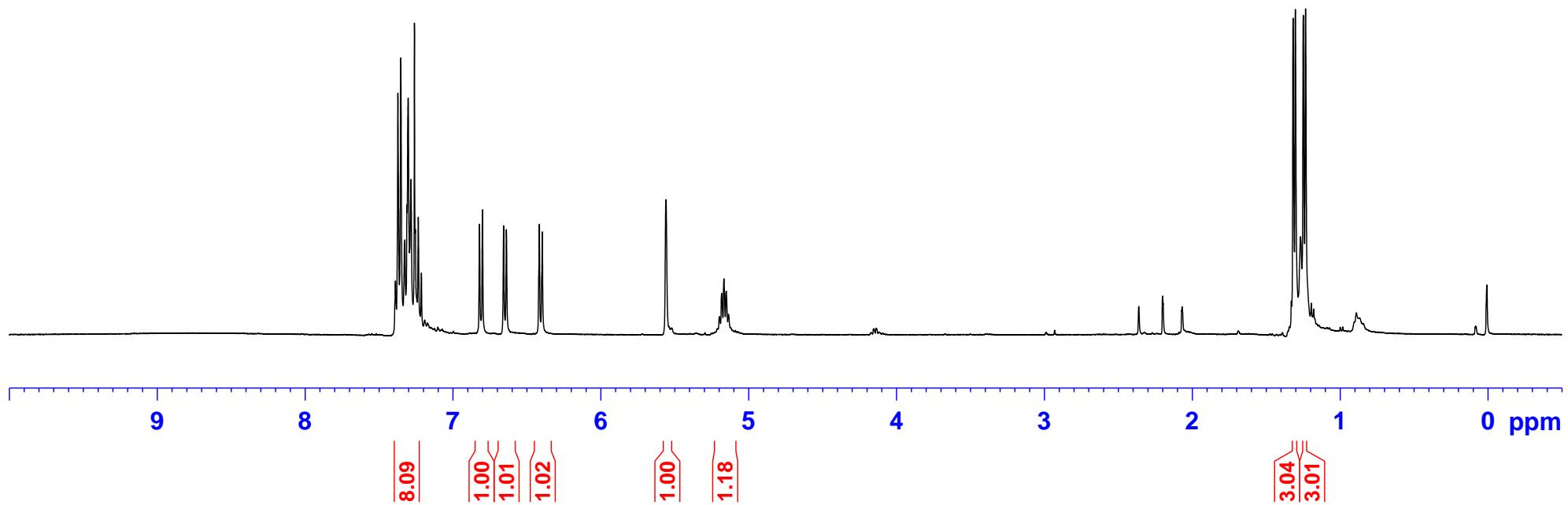
53.18
52.78



yzz-14-056-2



3ga



yzz-14-056-1

— 174.60

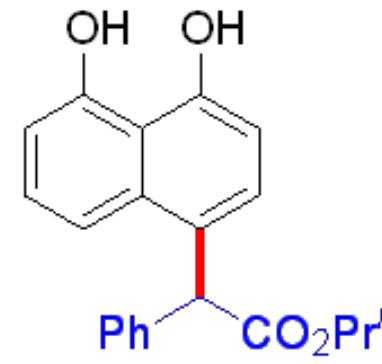
— 154.26
— 153.16

— 137.53
— 134.74
— 128.96
— 128.72
— 127.64
— 127.42
— 127.39
— 125.83
— 114.94
— 114.70
— 109.53
— 108.21

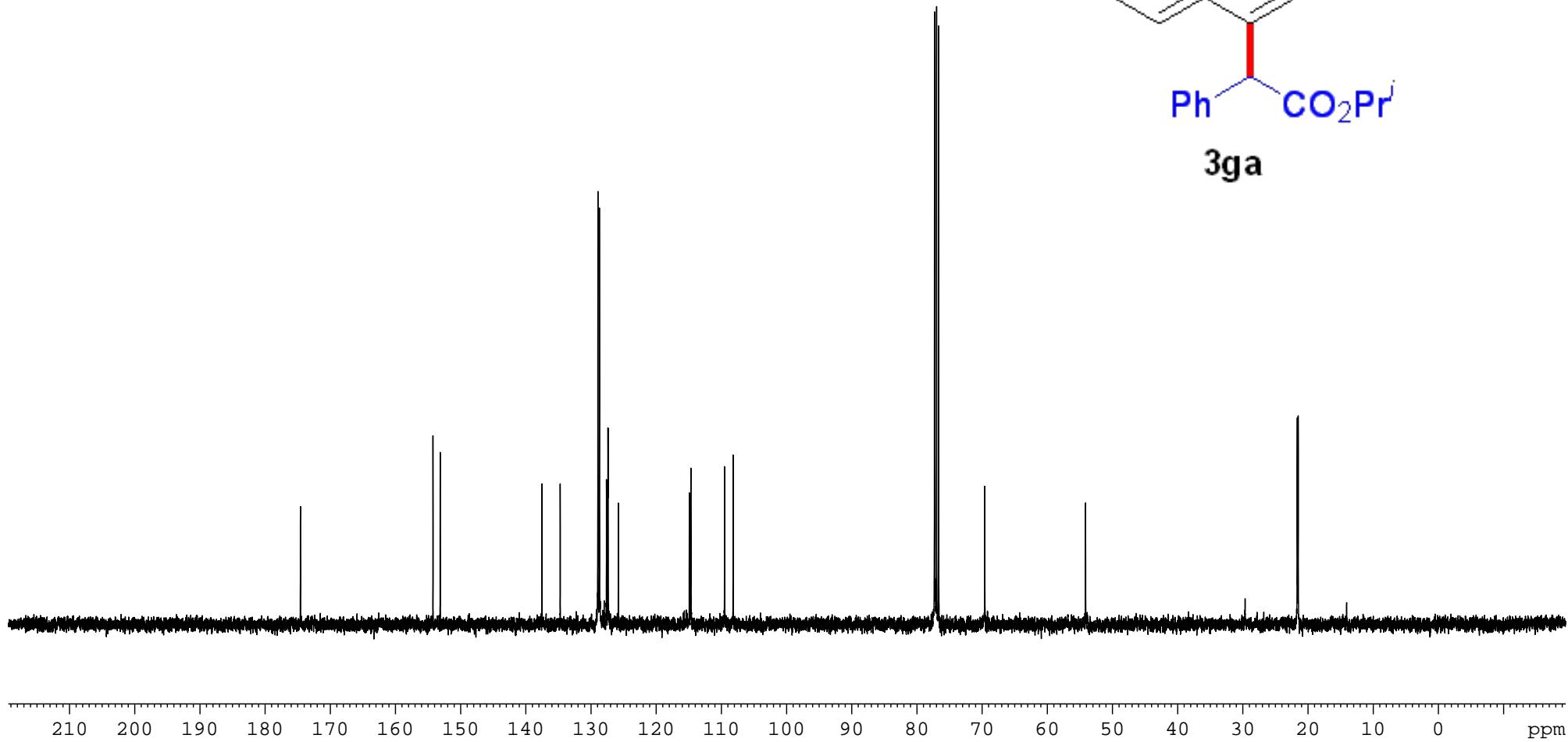
— 77.32
— 77.00
— 76.68
— 69.61

— 54.15

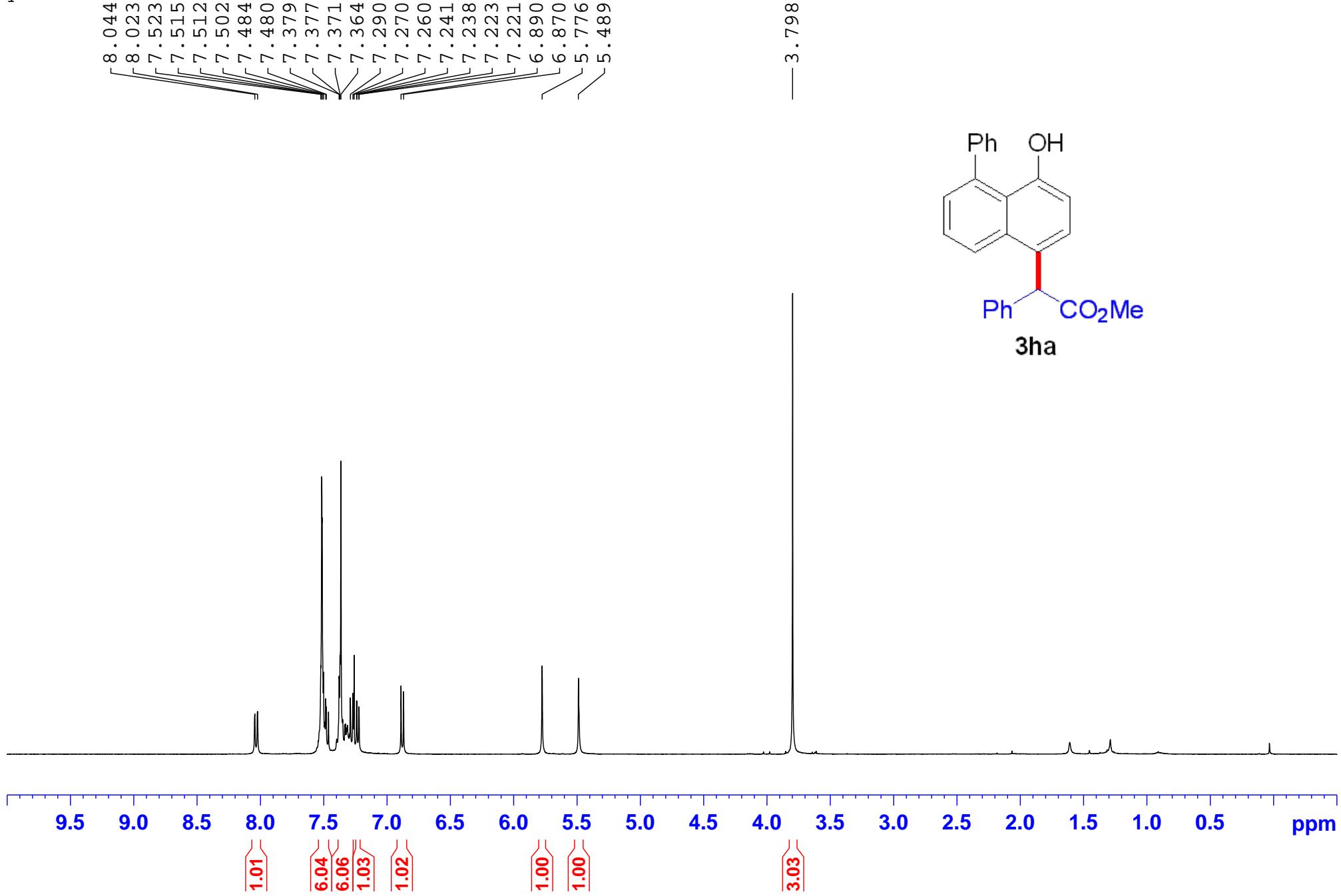
— 21.67
— 21.52



3ga

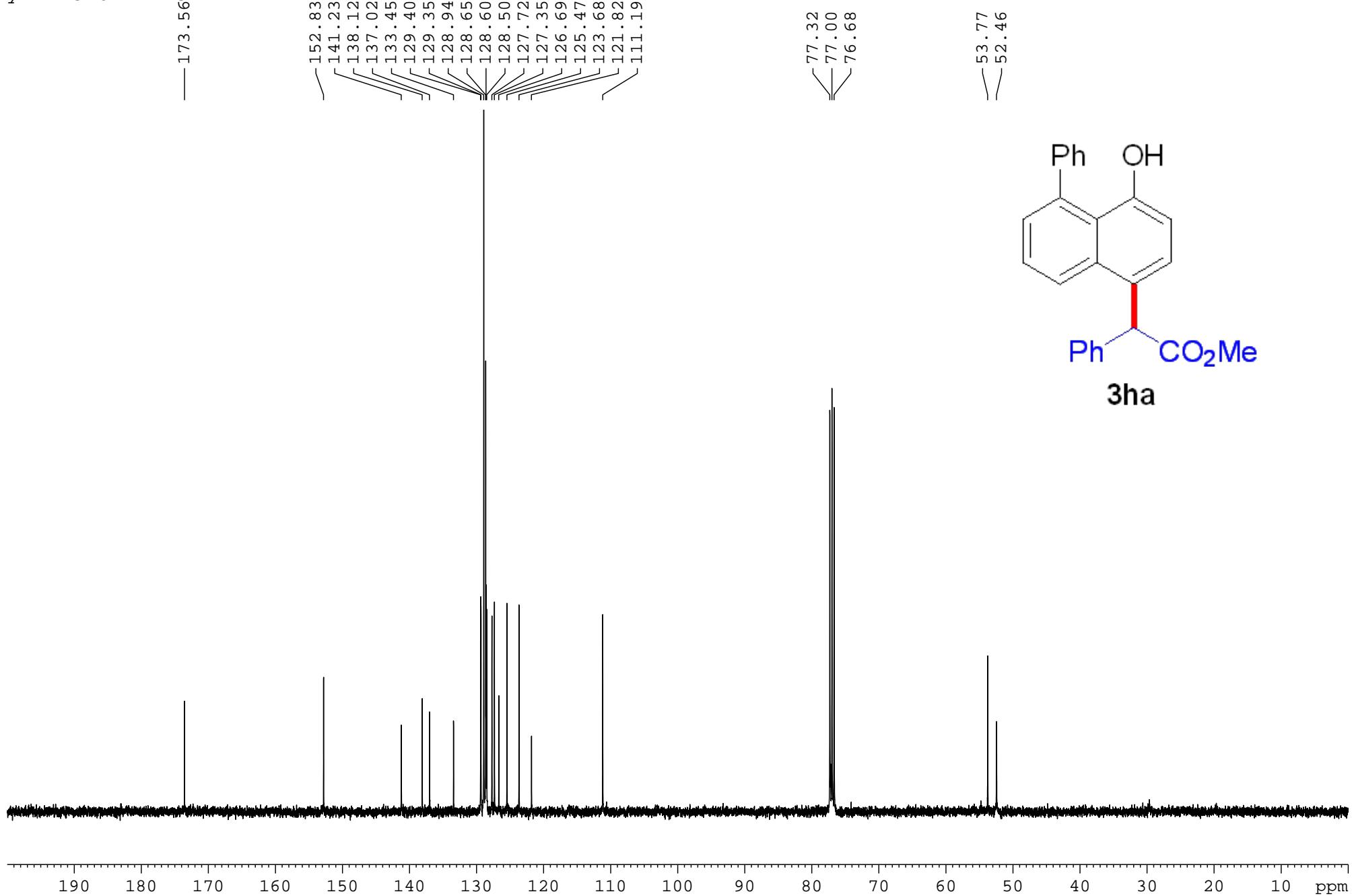


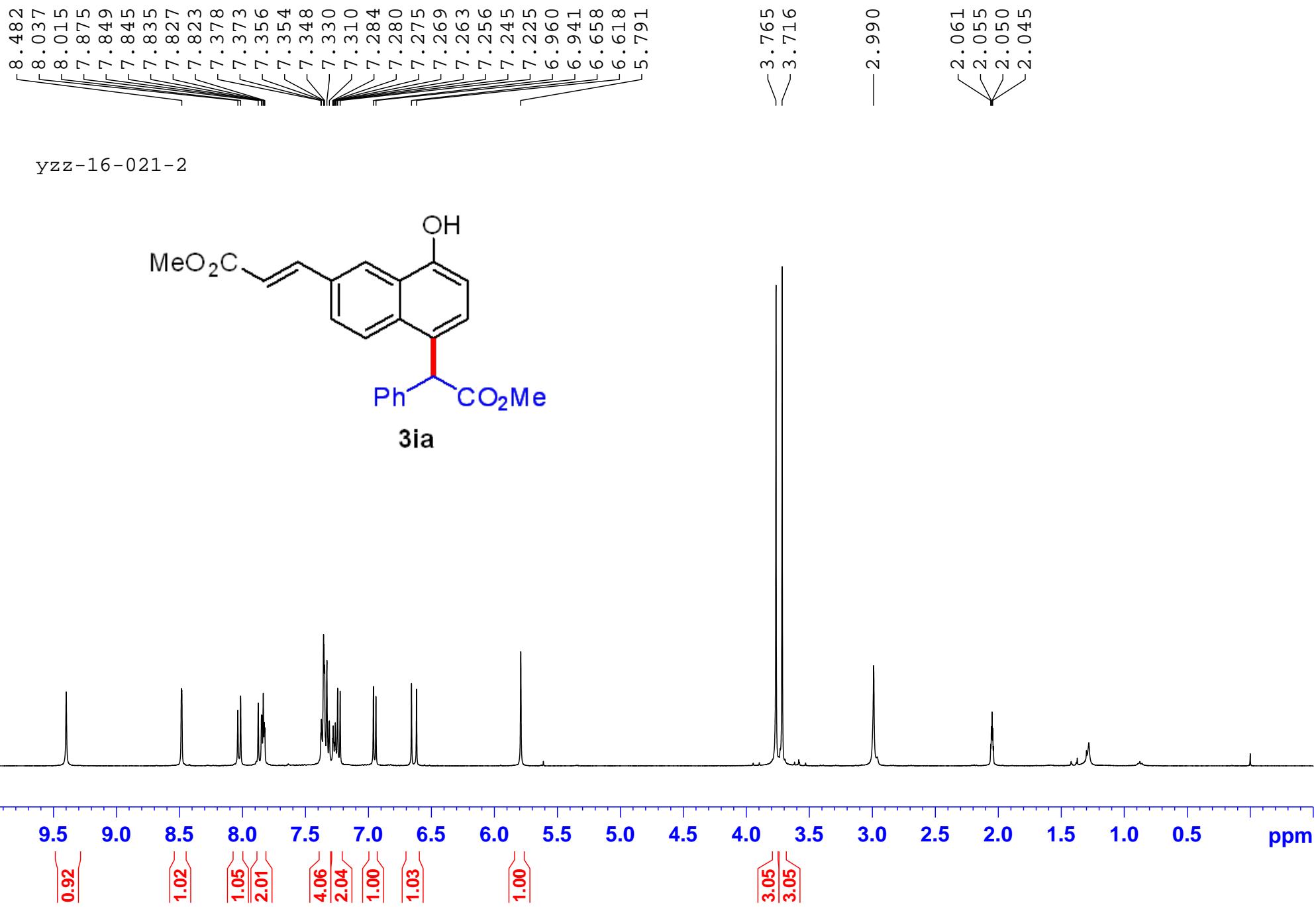
yzz-15-024-1



yzz-15-024-1-C

— 173.56C

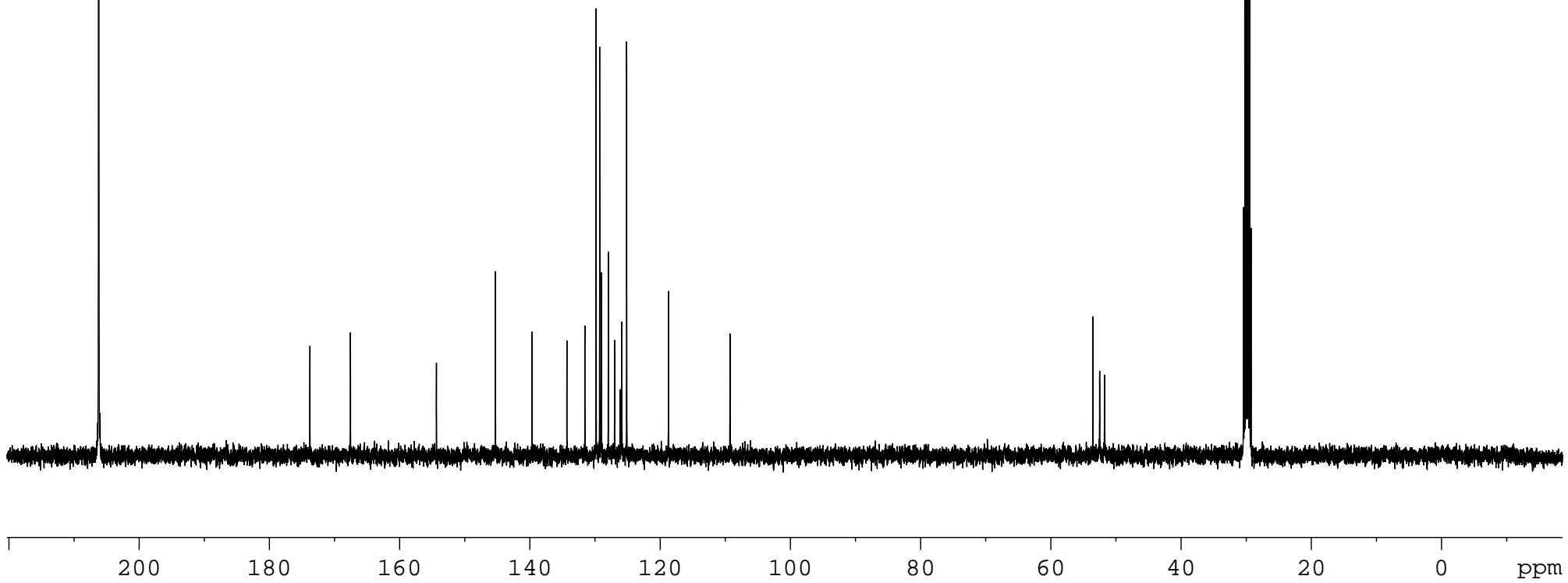
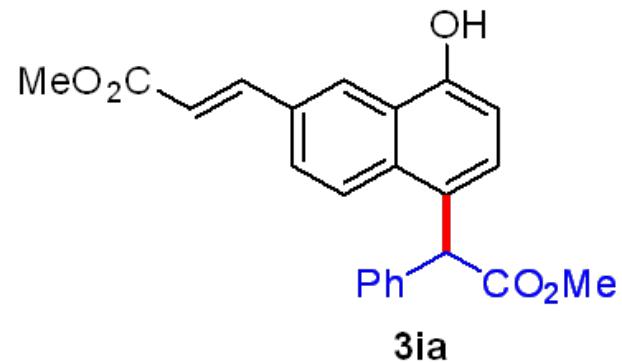


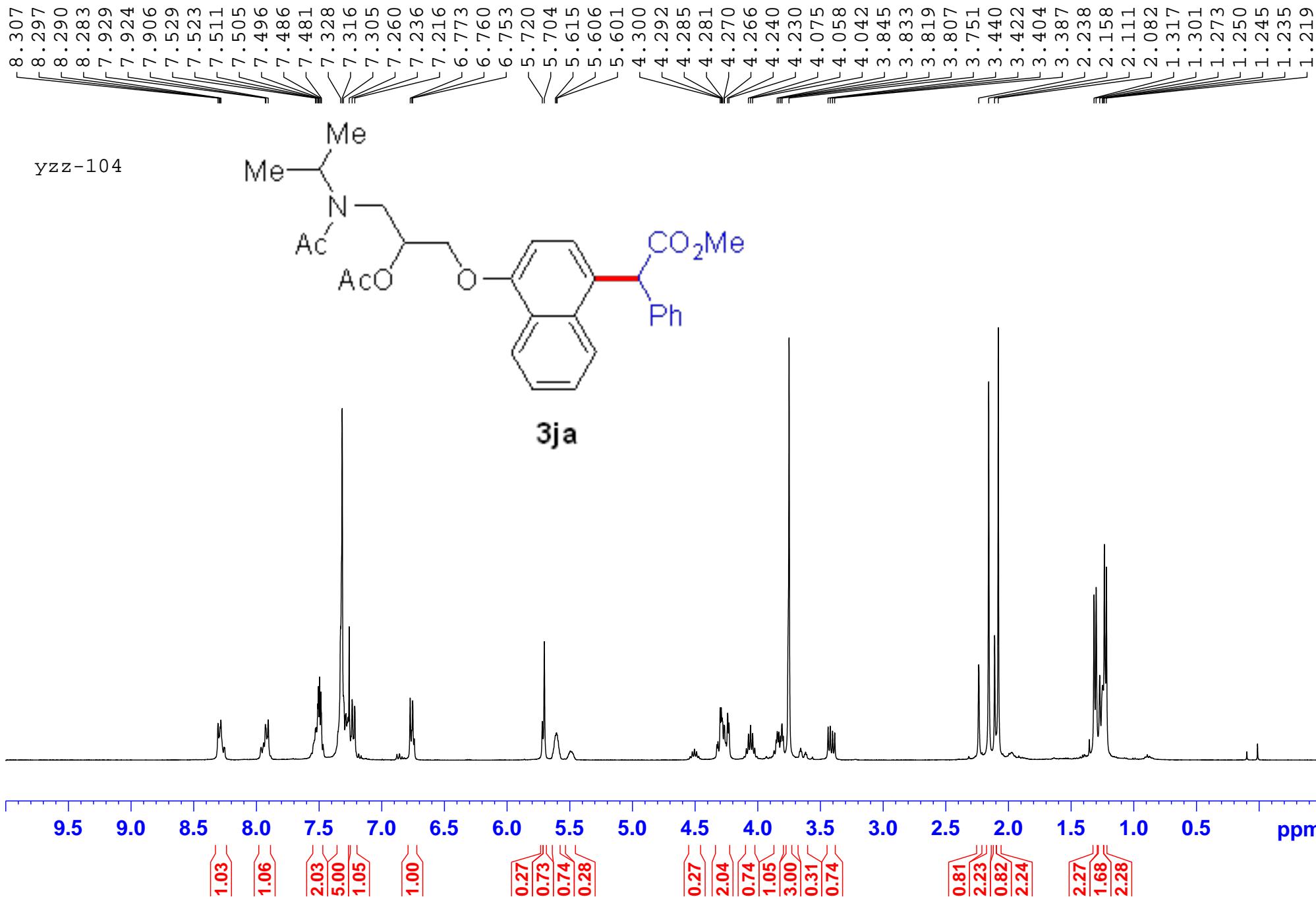


yzz-16-021-2-c
3

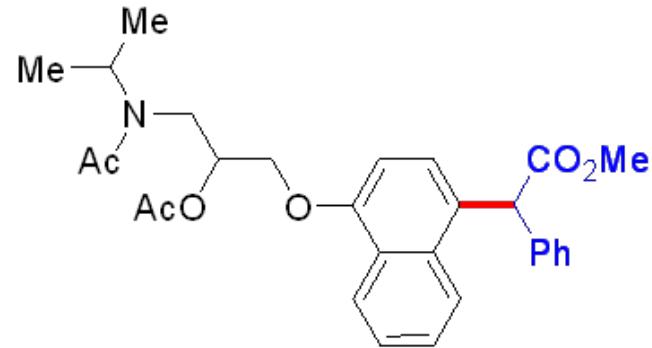
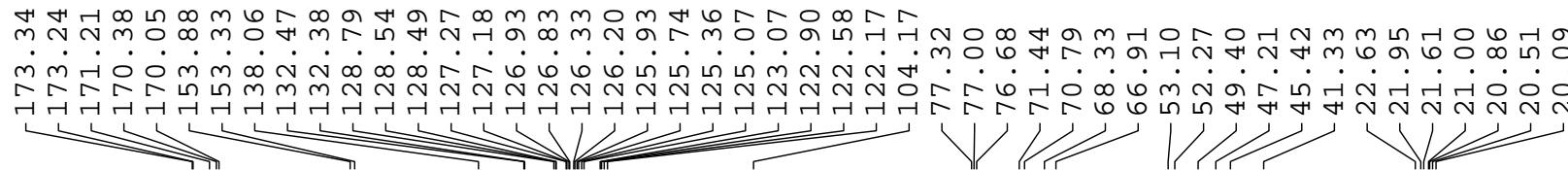
— 206 . 23 —

— 173.79
— 167.57

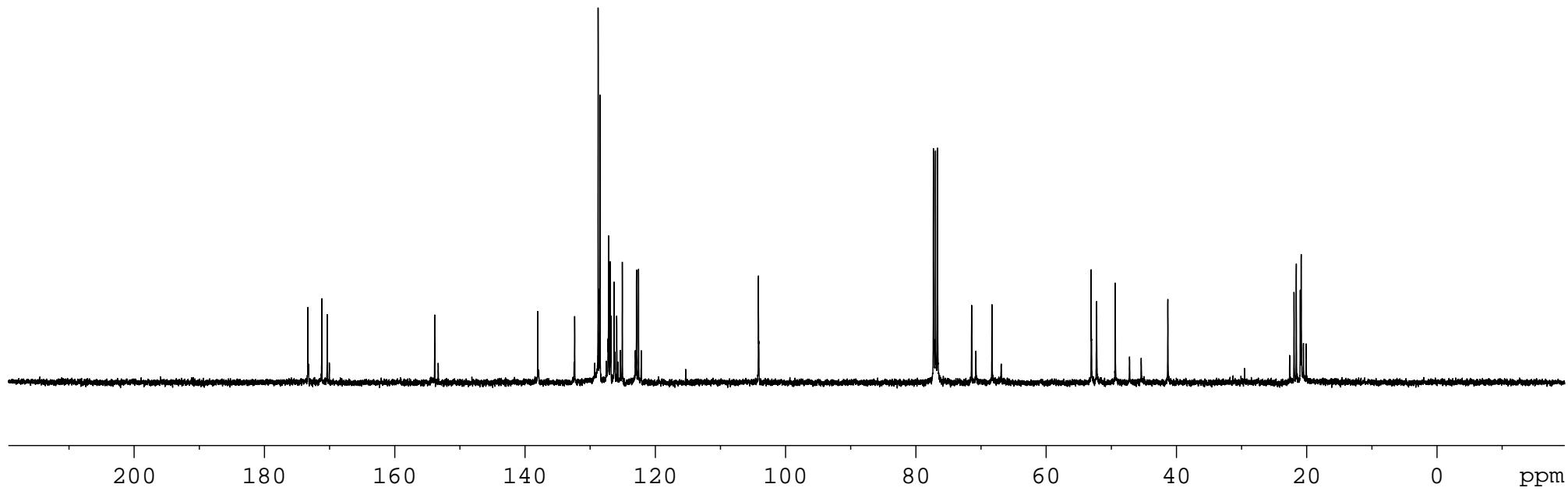


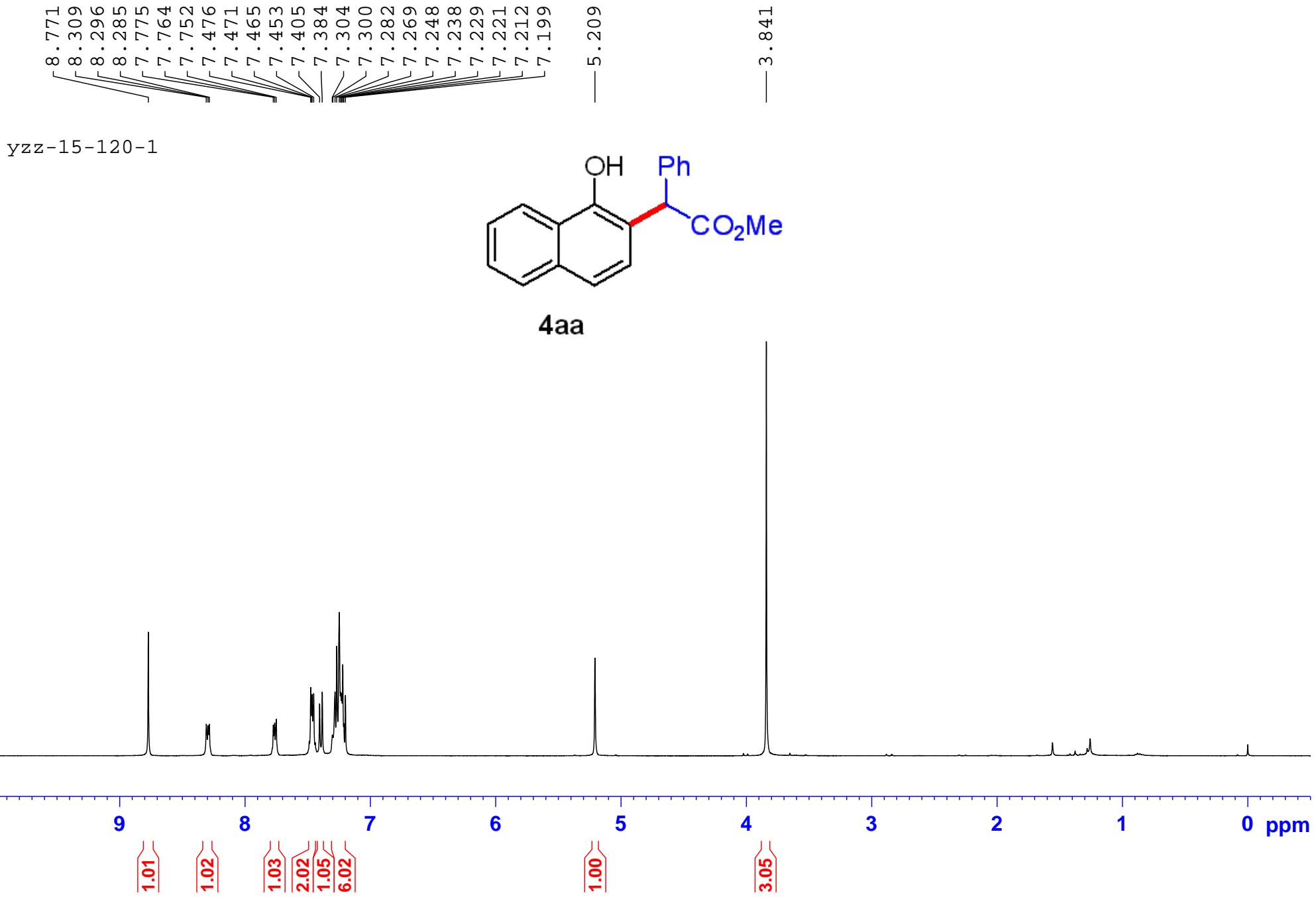


yzz-15-054-2-c



3ja





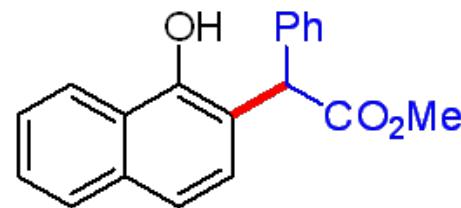
yzz-15-120-1-c

— 176.55

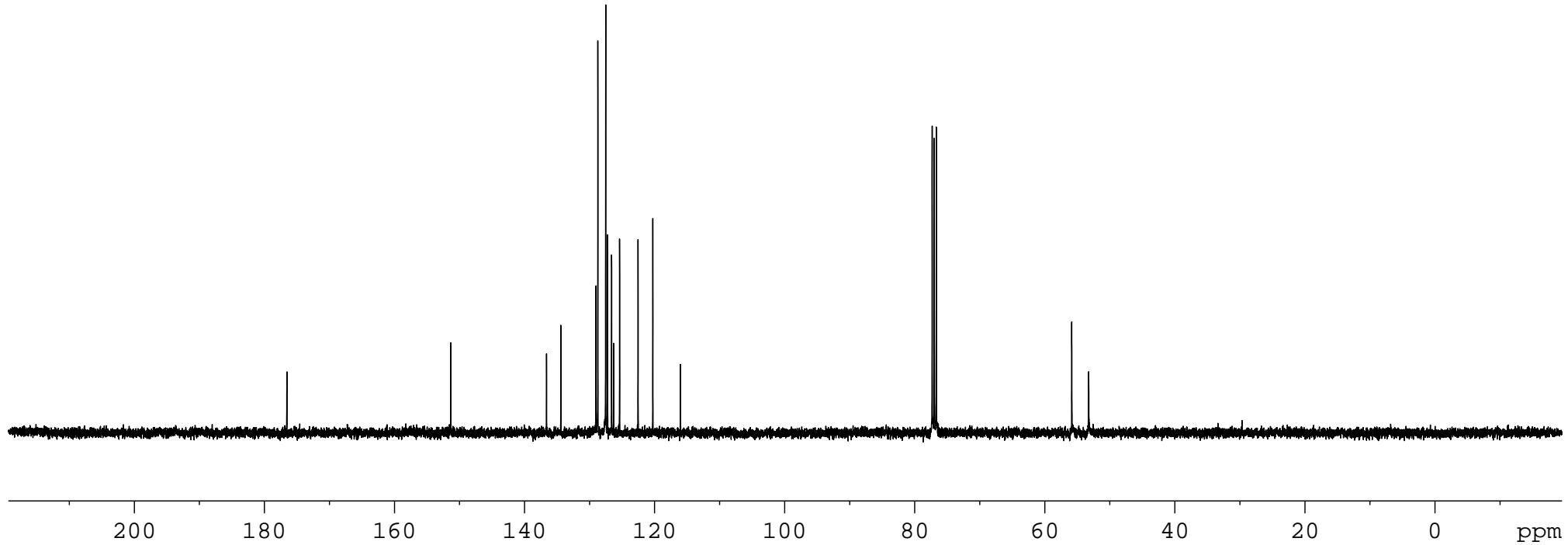
— 151.35
— 136.62
— 134.42
— 129.03
— 128.74
— 127.52
— 127.23
— 126.61
— 126.32
— 125.39
— 122.55
— 120.28
— 116.04

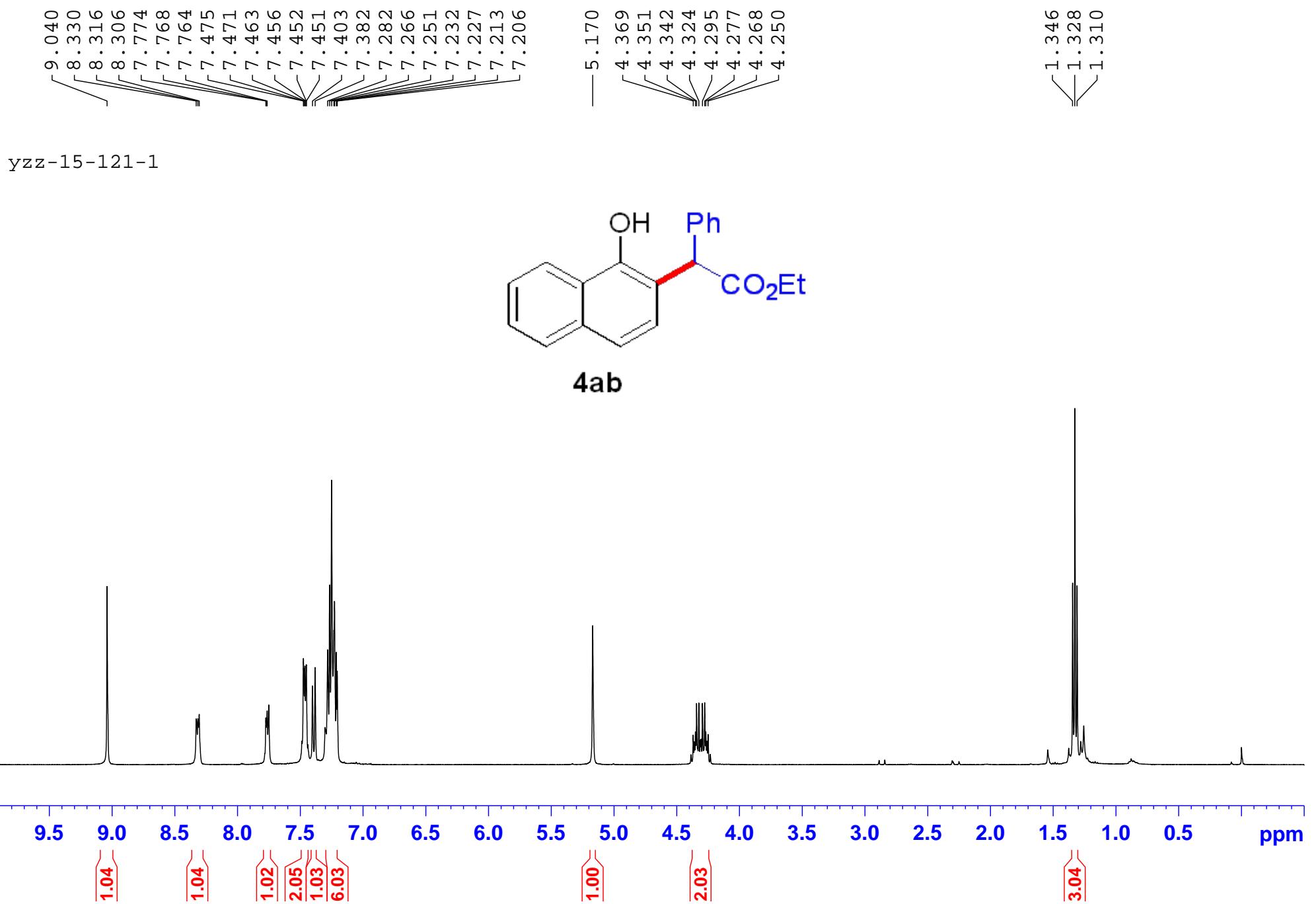
— 77.32
— 77.00
— 76.68

— 55.86
— 53.26



4aa





yzz-15-121-1-c

— 176.25

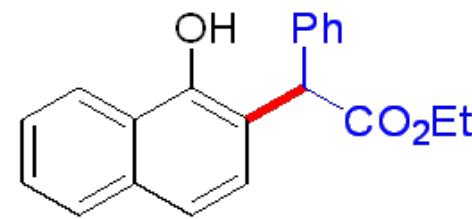
— 151.55
— 136.75
— 134.44
— 129.20
— 128.71
— 127.47
— 127.44
— 127.18
— 126.59
— 126.41
— 125.33
— 122.69
— 120.16
— 115.96

— 77.32
— 77.00
— 76.68

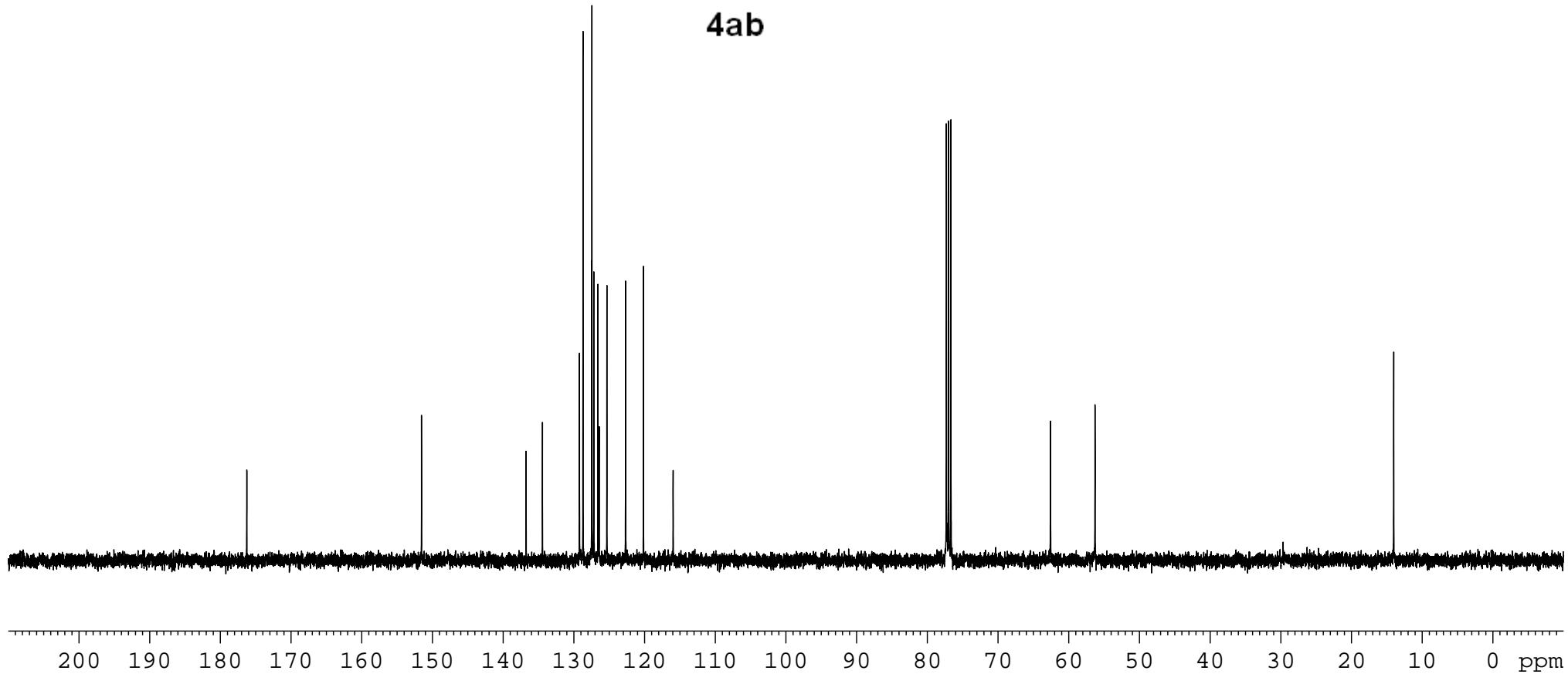
— 62.59

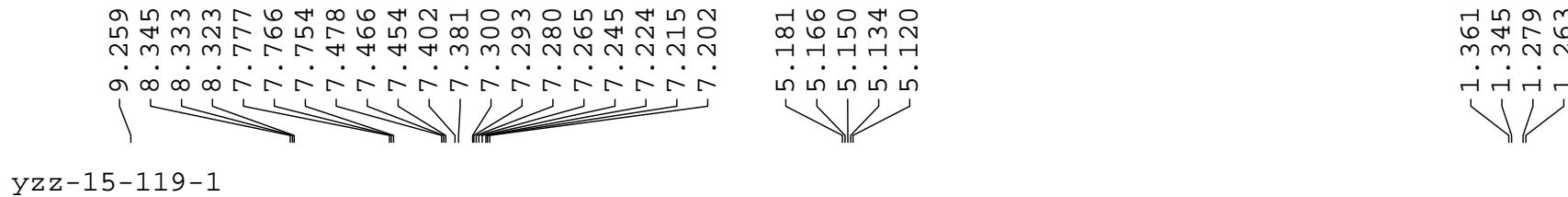
— 56.29

— 14.04

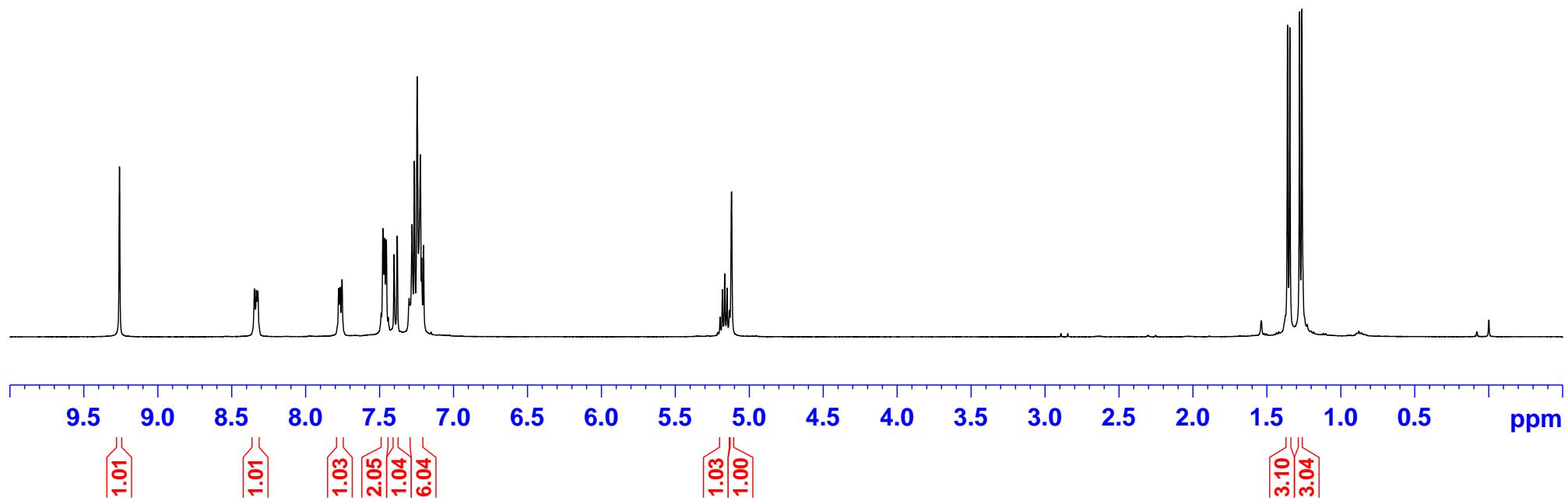
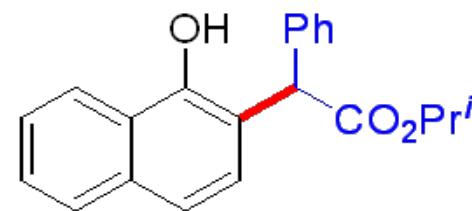


4ab

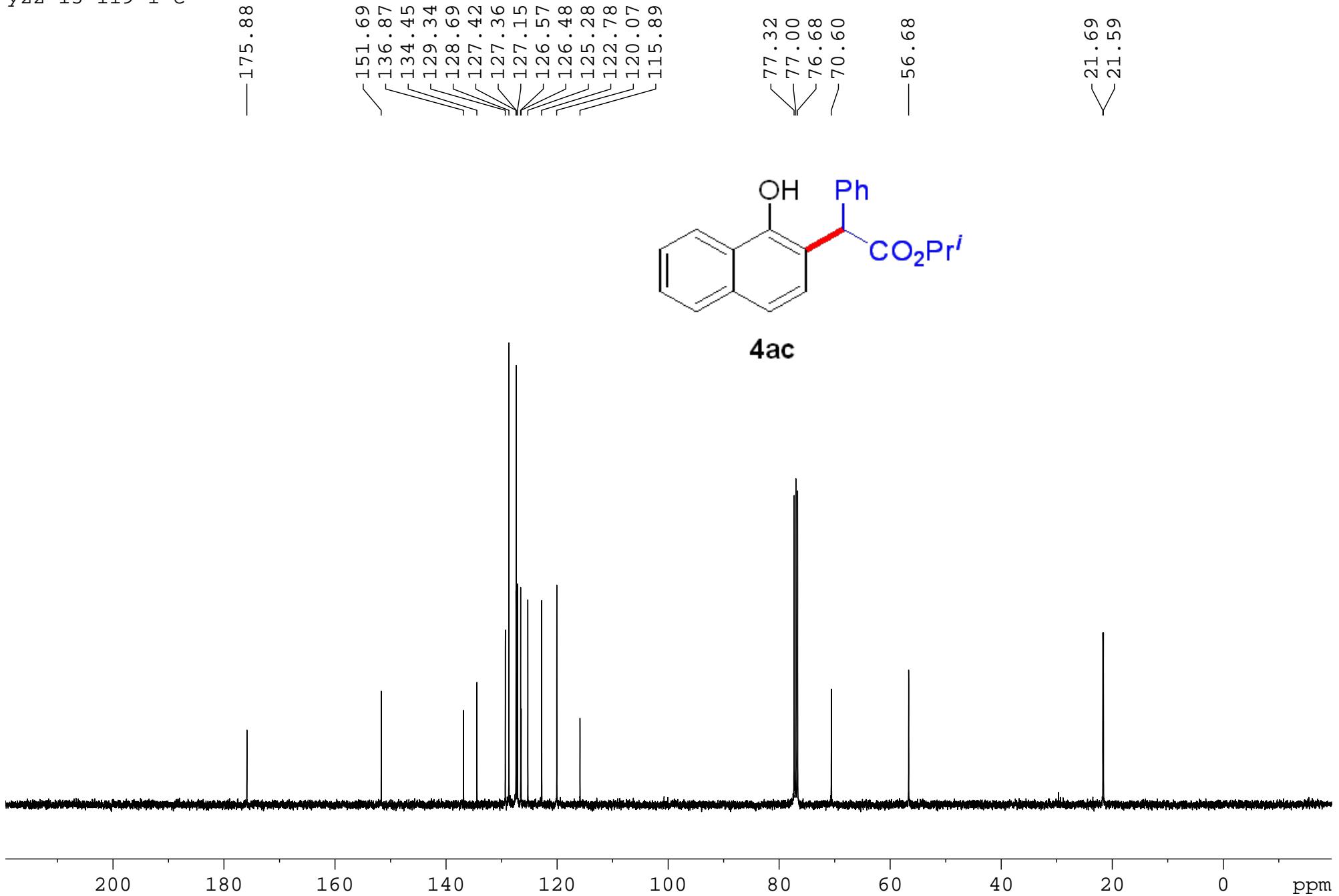


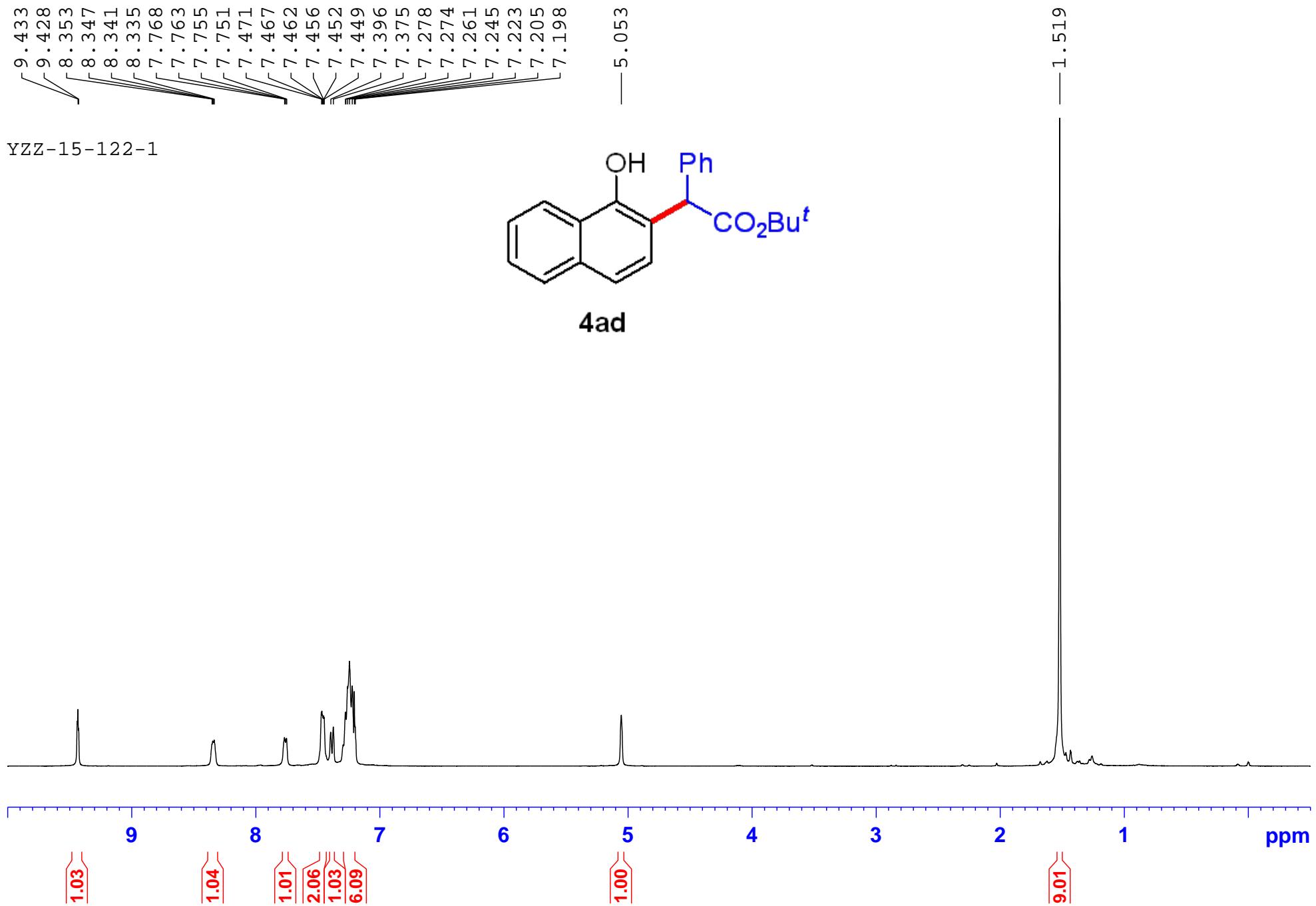


yzz-15-119-1

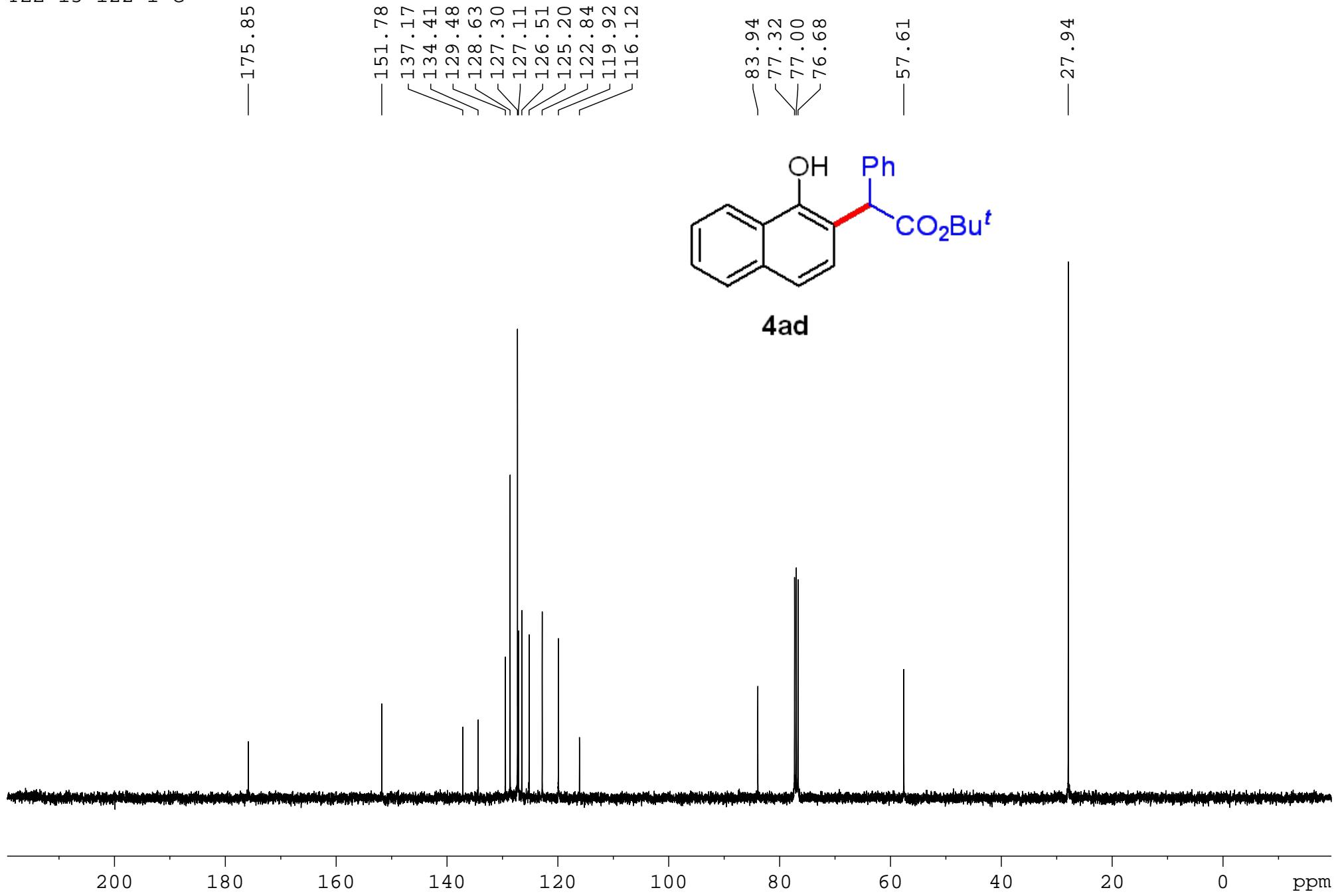


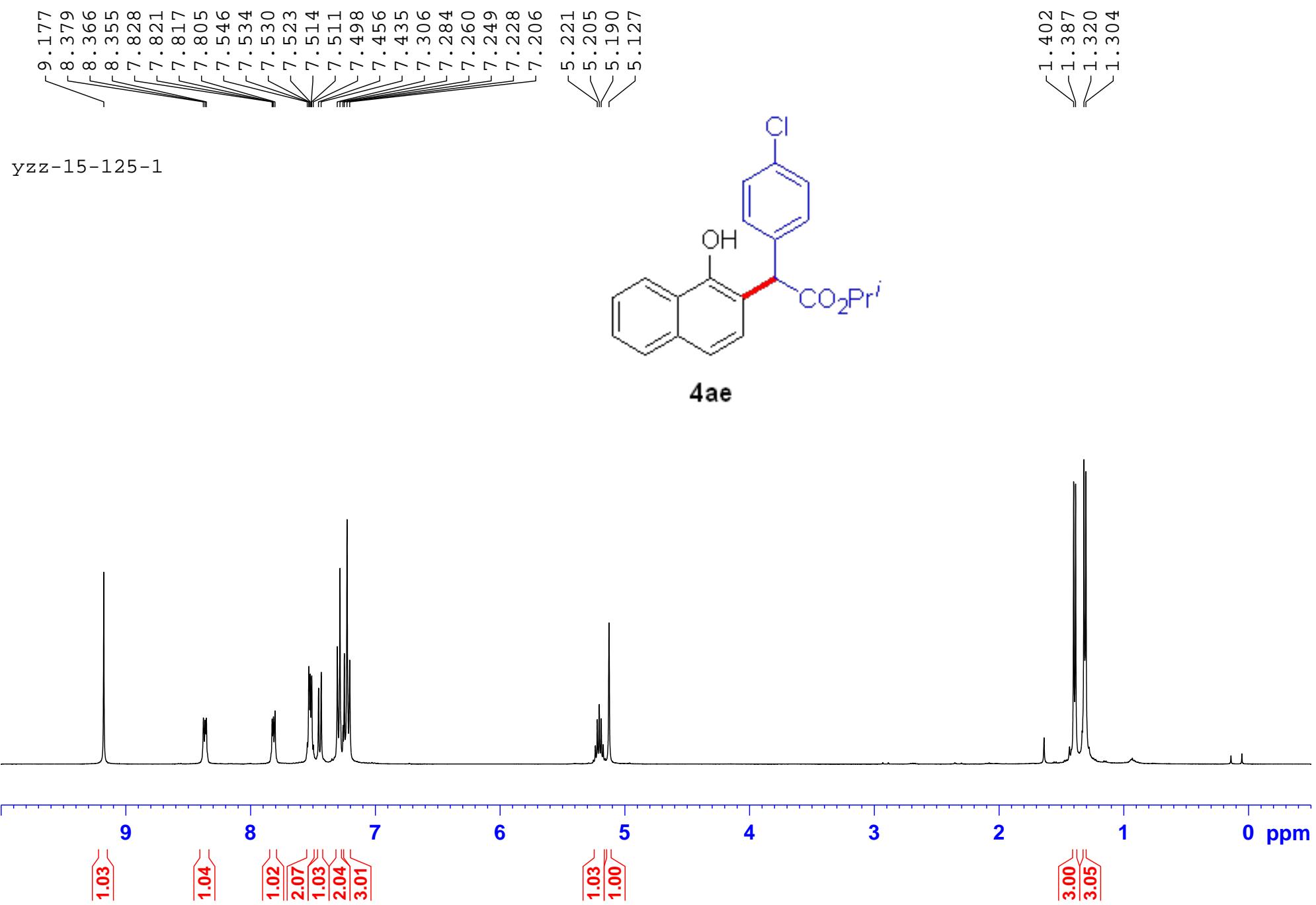
yzz-15-119-1-c



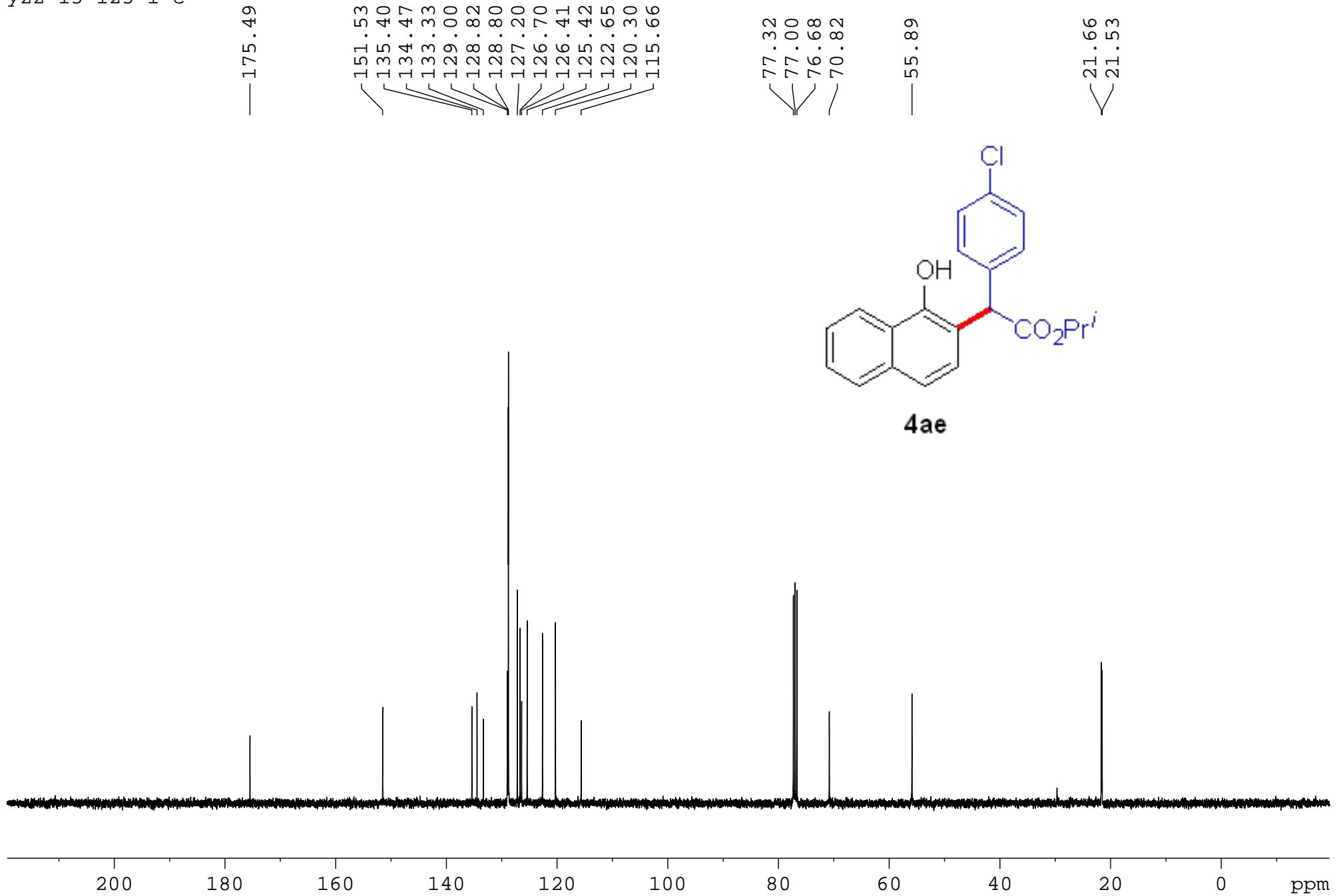


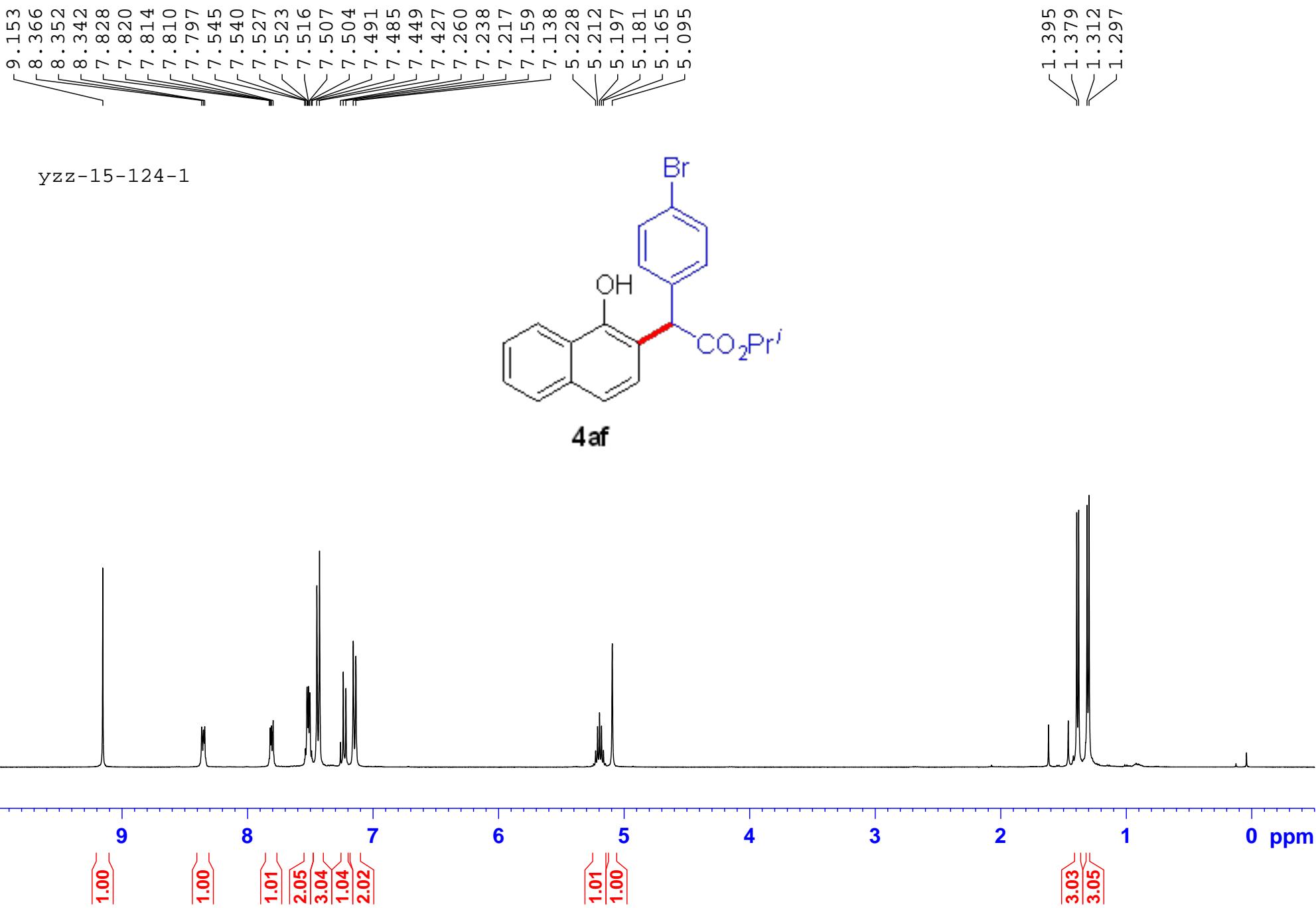
YZZ-15-122-1-C



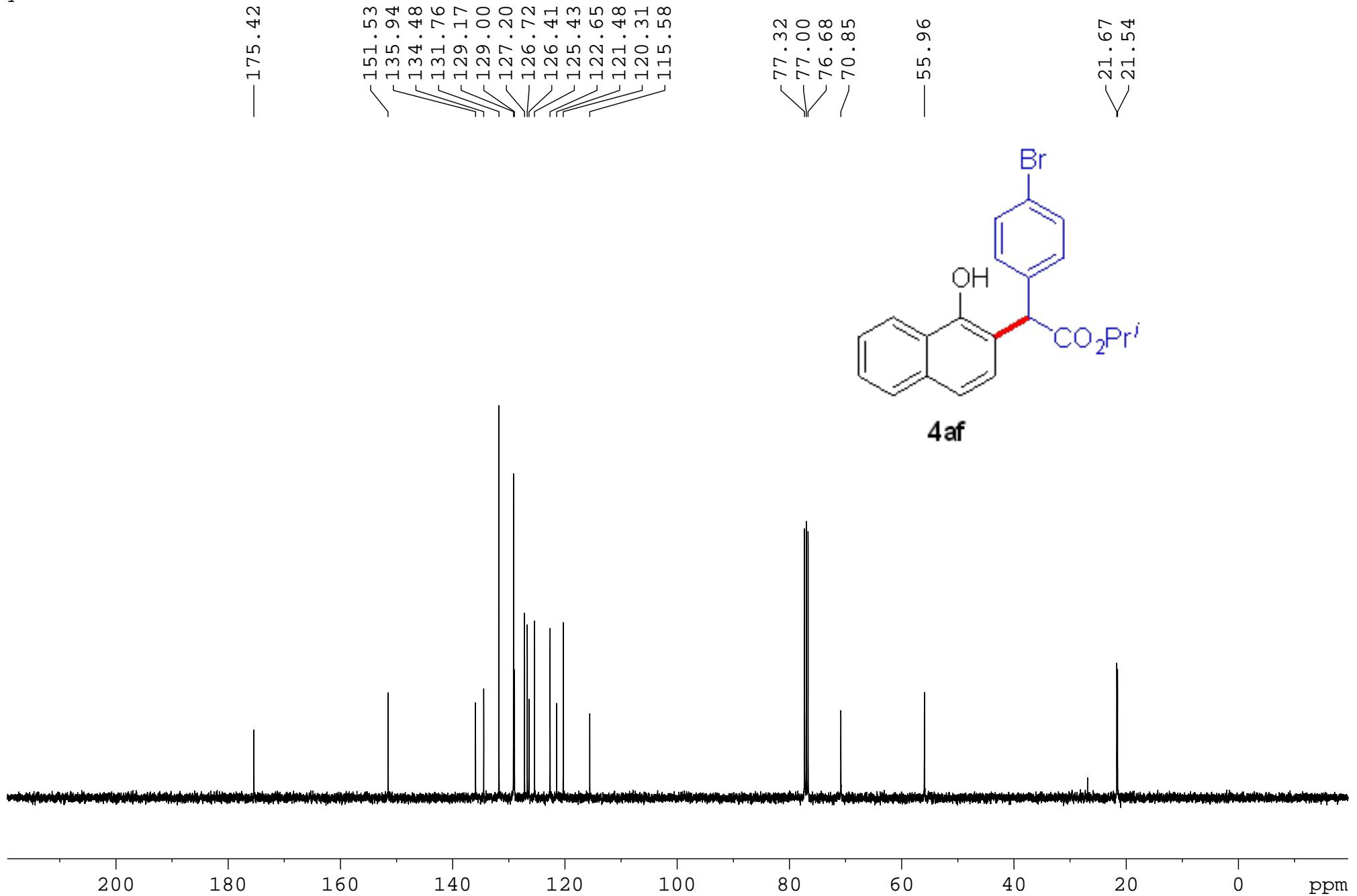


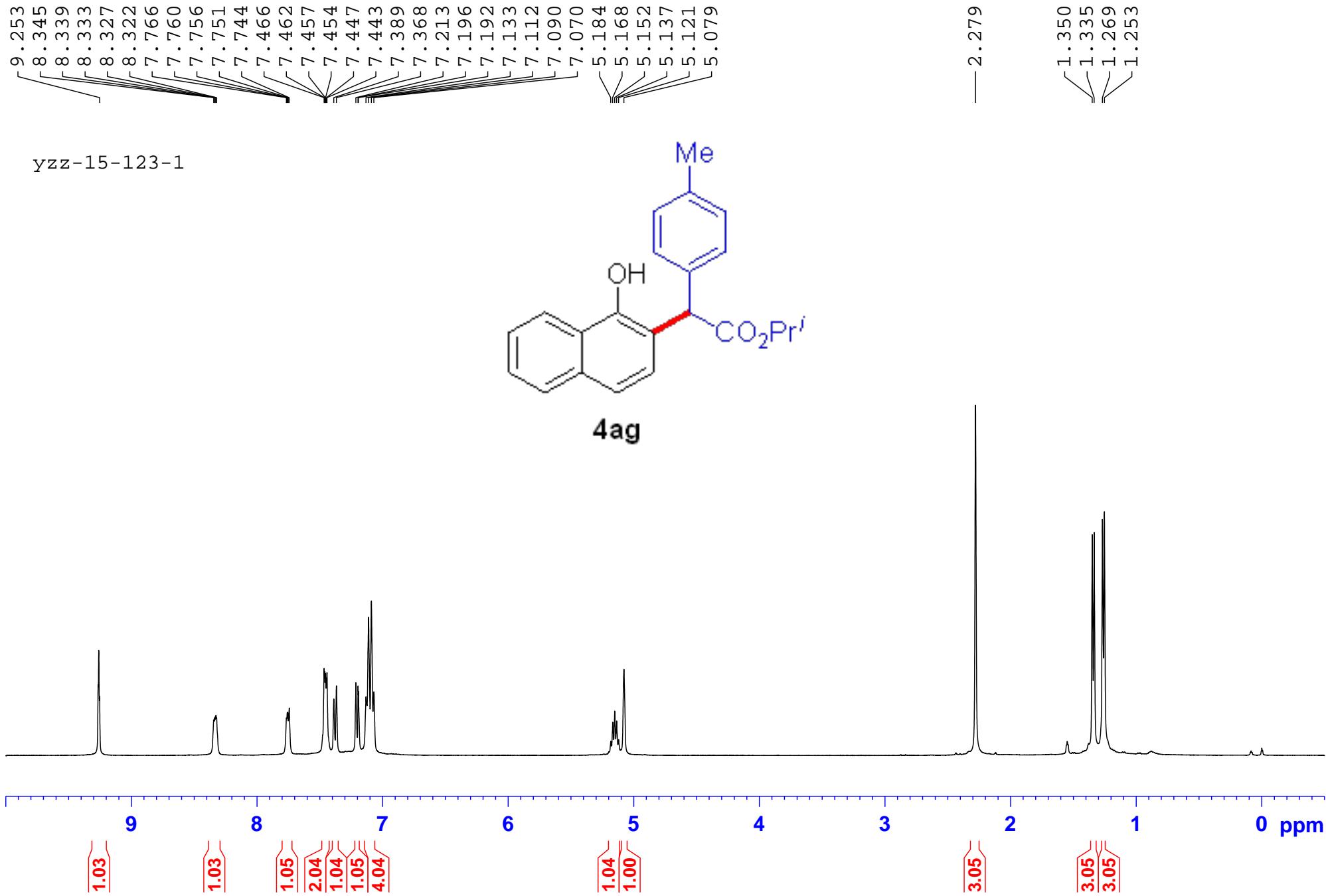
yzz-15-125-1-c





yzz-15-124-1-c





yzz-15-123-1-c

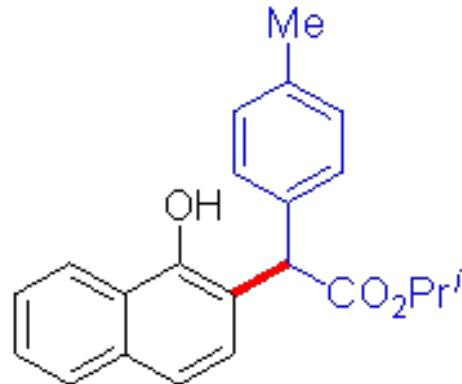
— 176.03

151.64
137.09
134.40
133.84
129.38
129.33
127.21
127.12
126.51
126.47
125.22
122.79
120.00
116.07

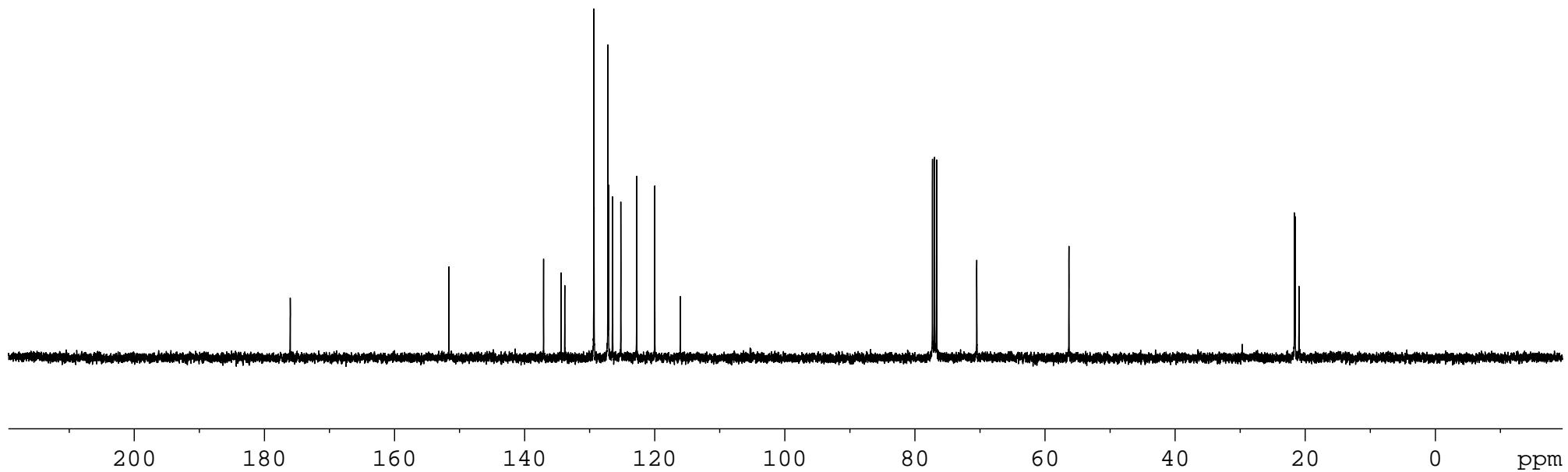
77.32
77.00
76.68
70.51

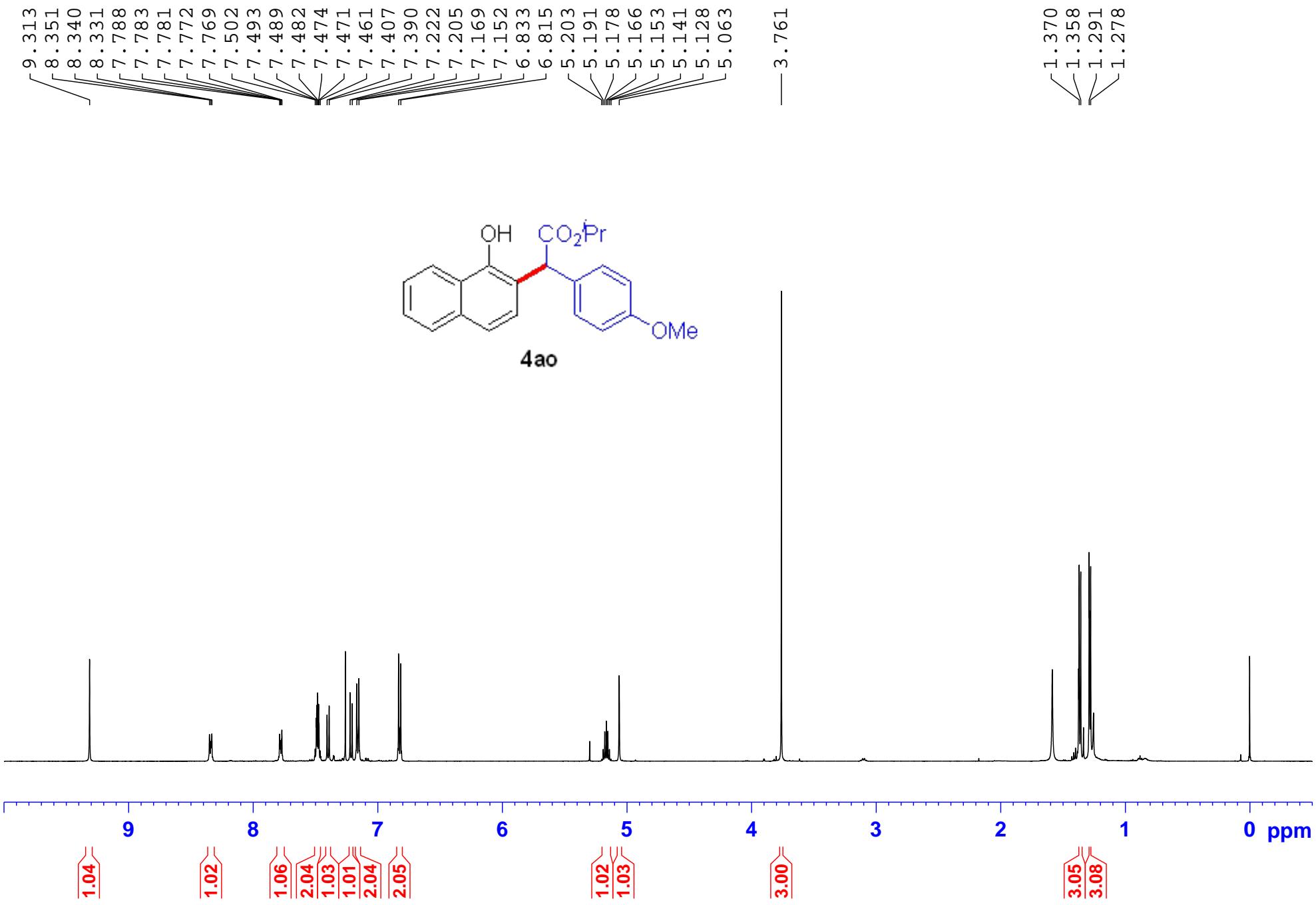
— 56.34

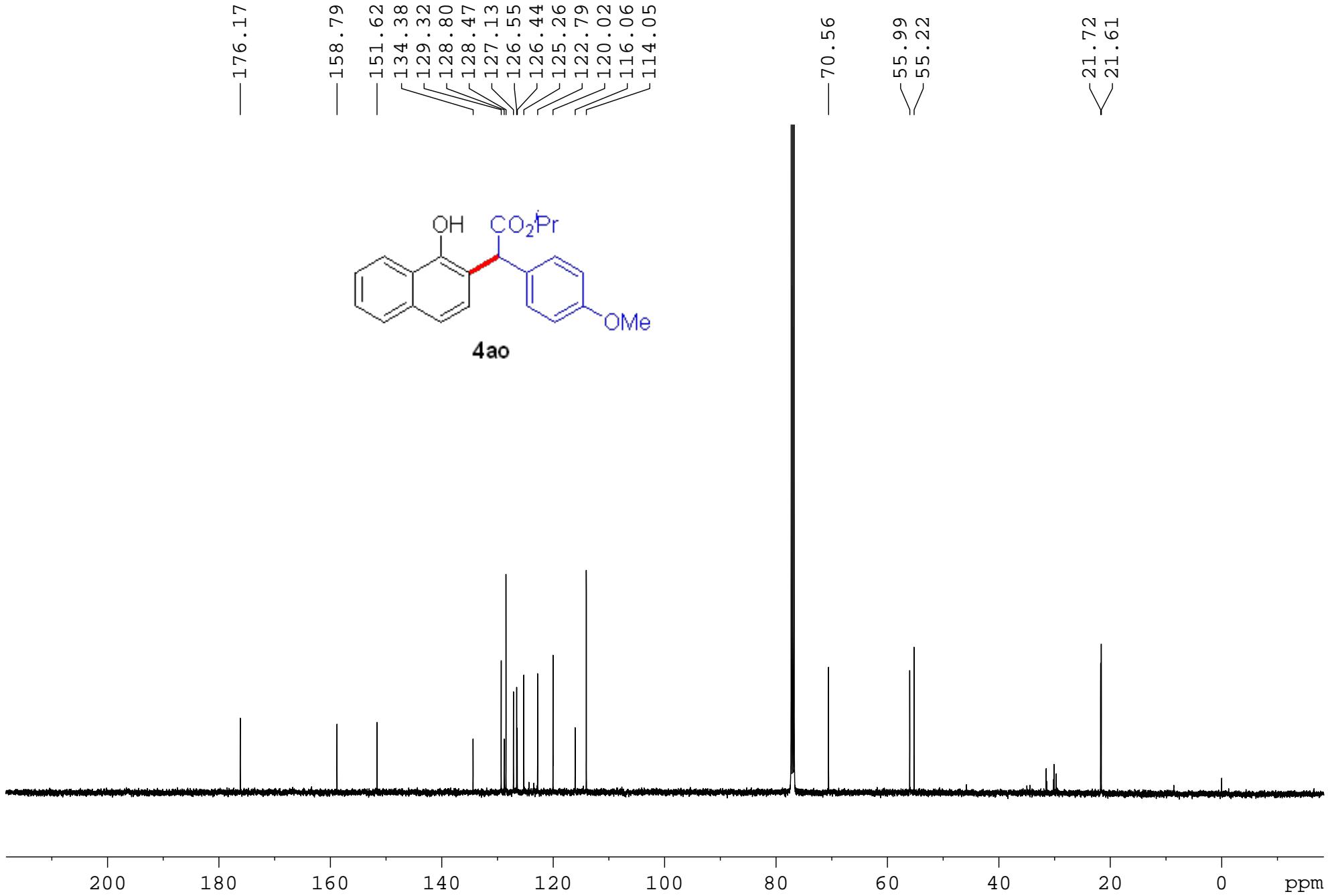
21.68
21.58
20.97

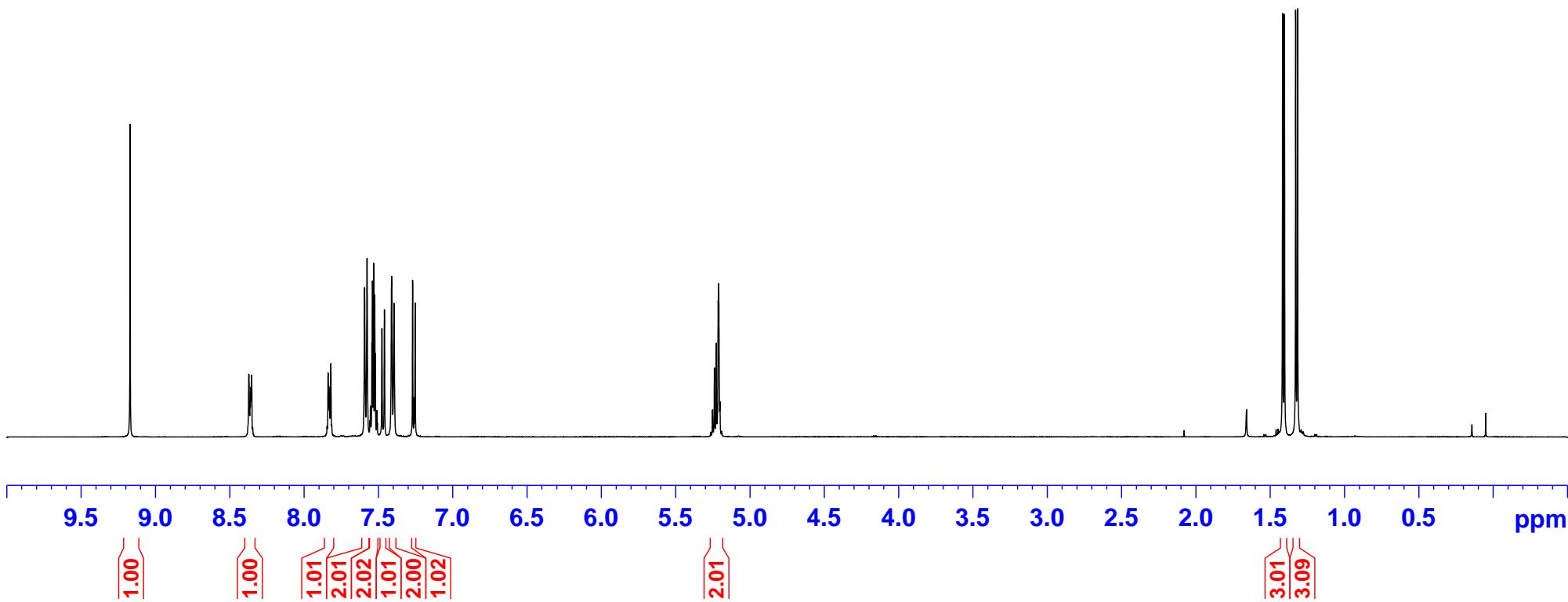
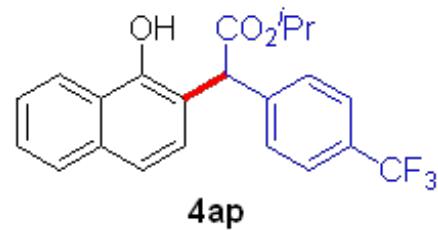
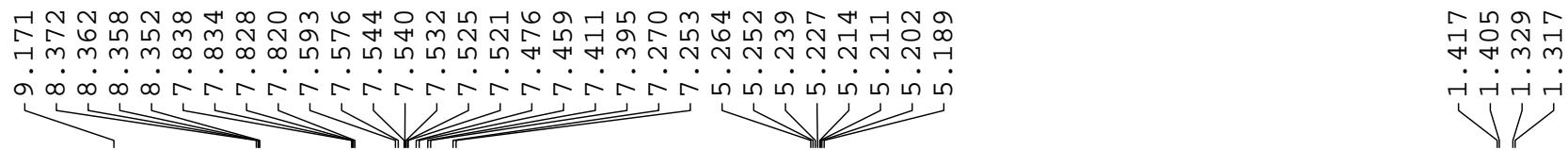


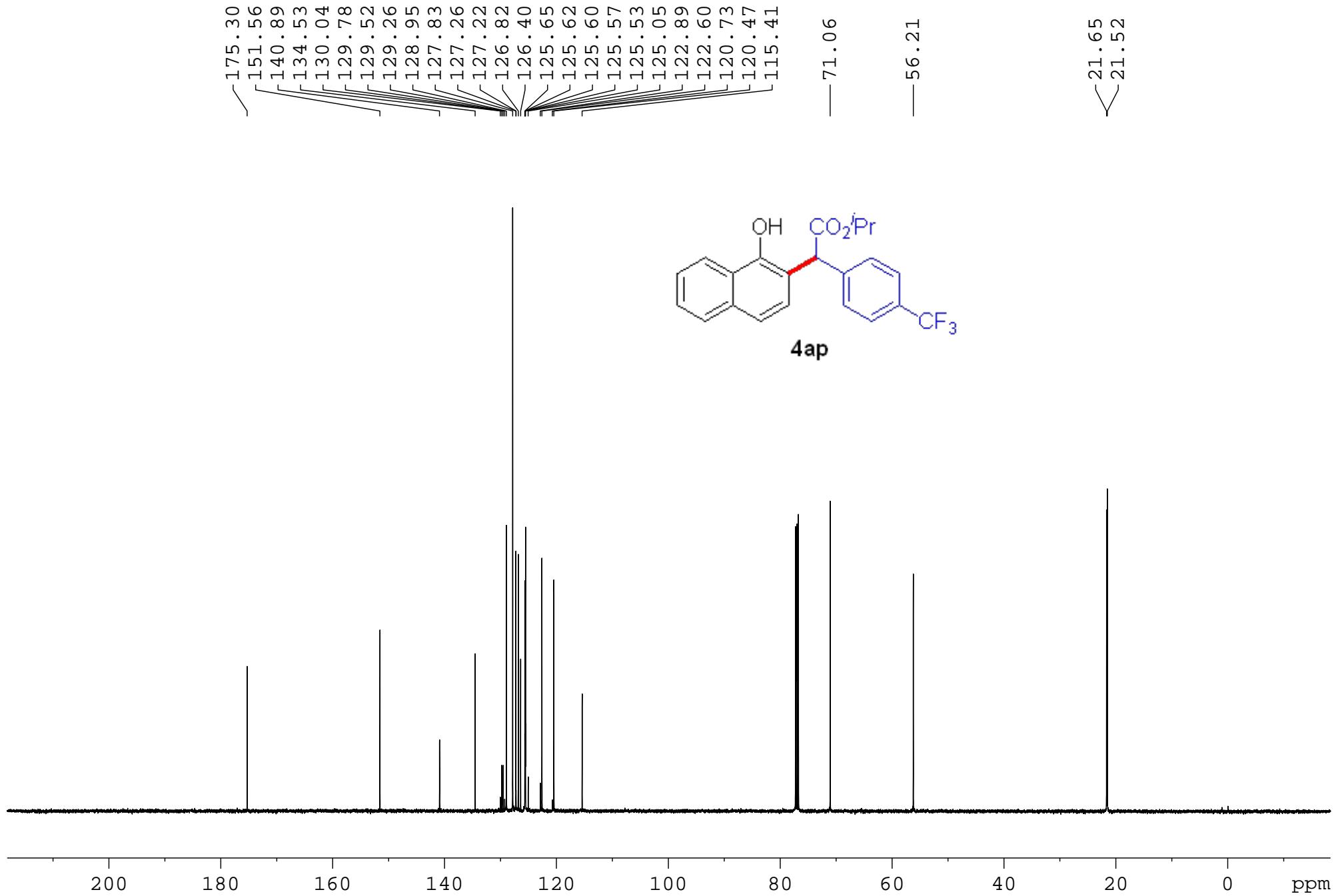
4ag

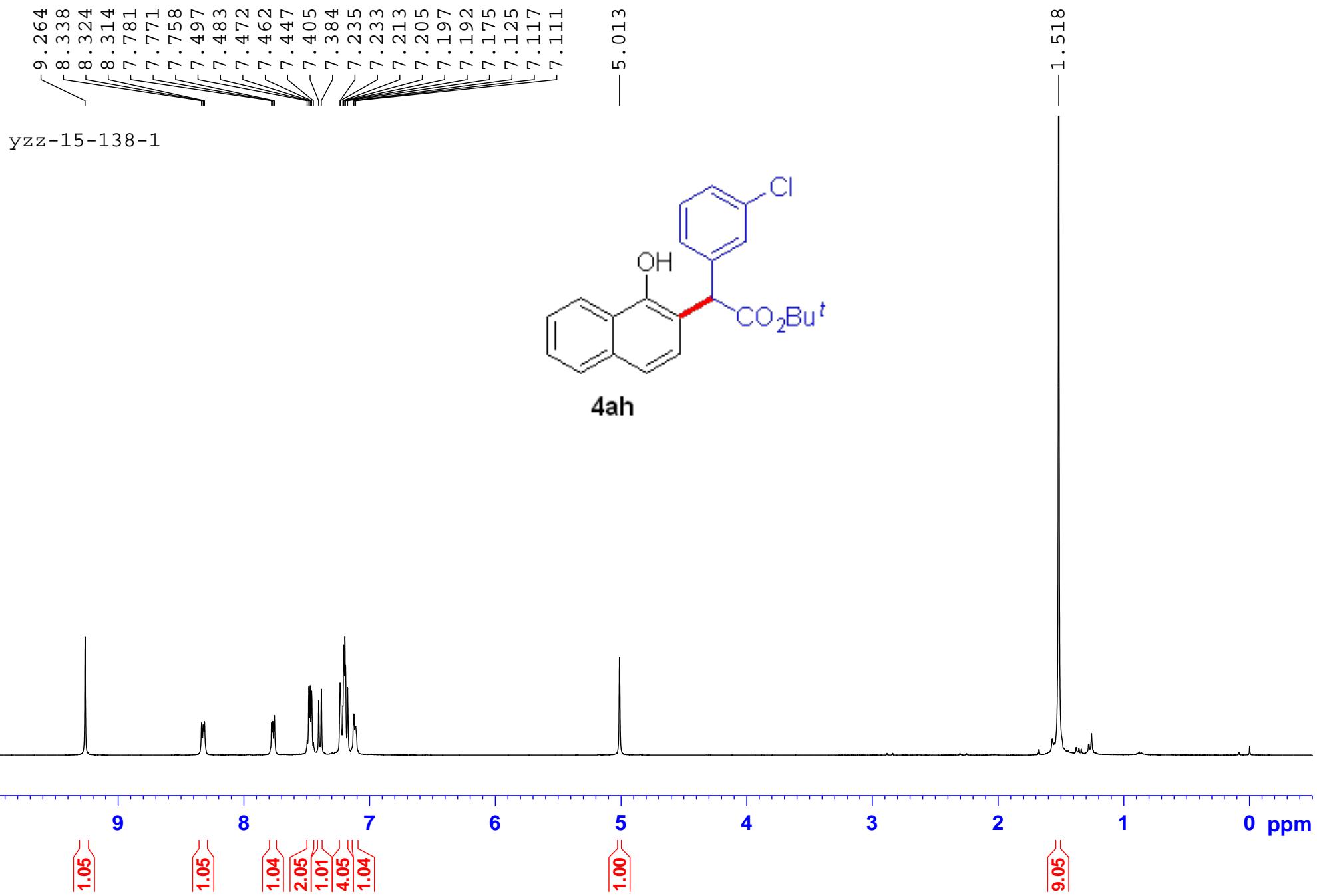












yzz-15-138-1-c

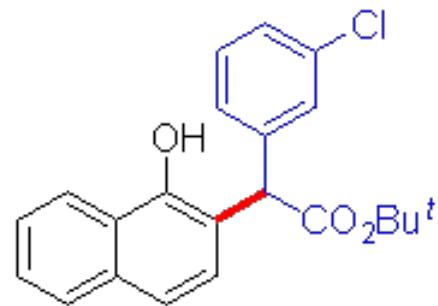
— 175.20

151.68
139.25
134.50
134.48
129.83
129.17
127.64
127.55
127.18
126.67
126.50
125.56
125.35
122.77
120.19
115.66

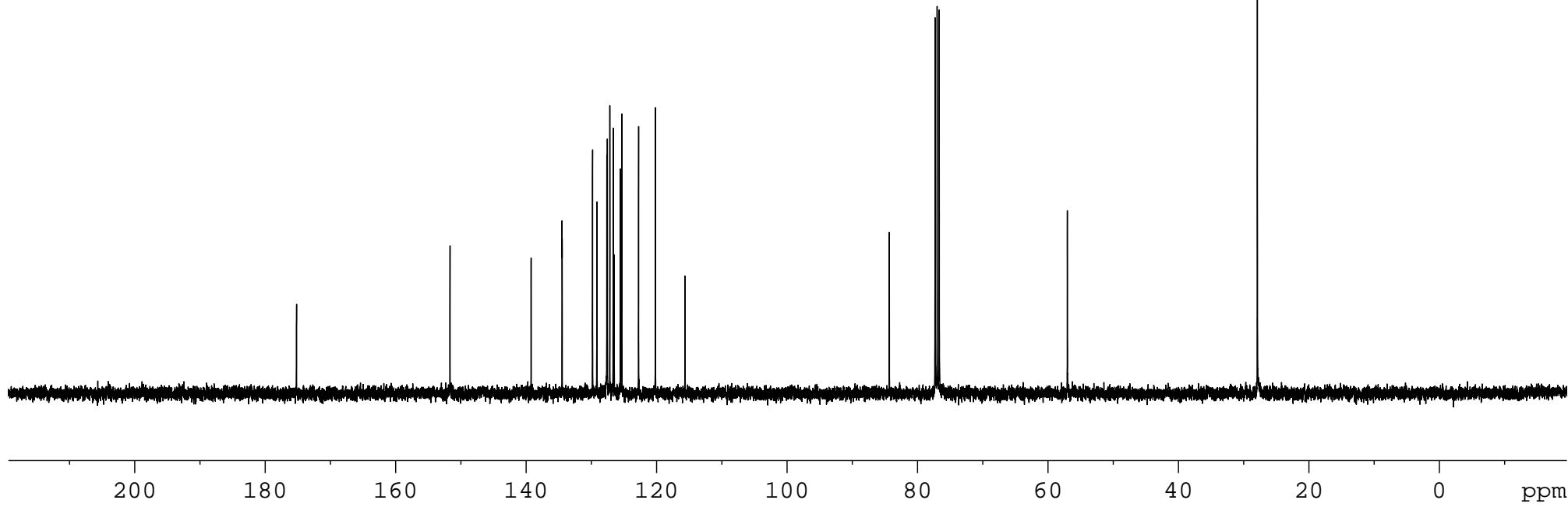
84.33
77.32
77.00
76.68

— 57.04

— 27.91



4ah

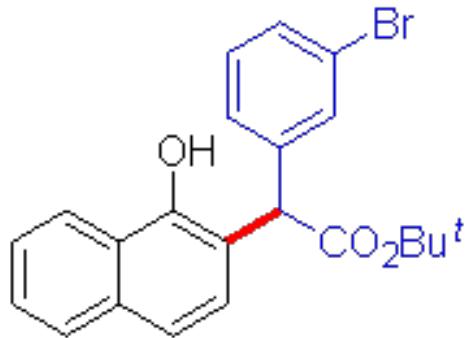


yzz-15-137-1

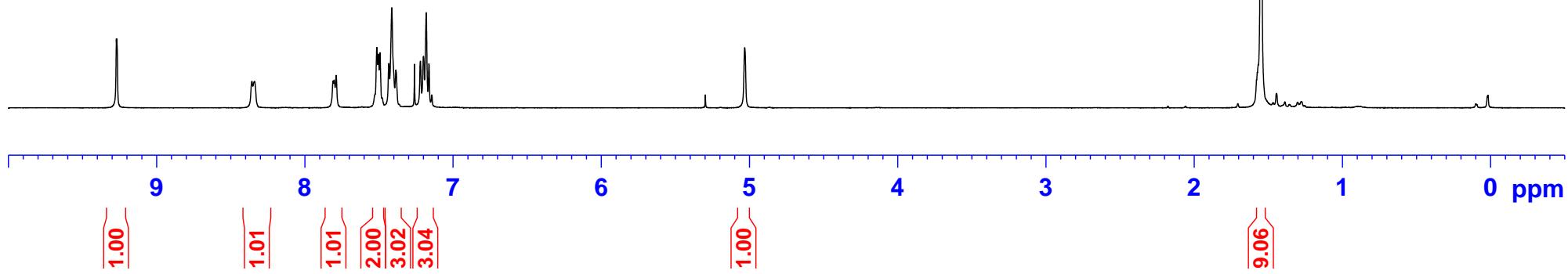
9.272
9.269
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8.338
7.809
7.802
7.789
7.526
7.514
7.503
7.492
7.480
7.433
7.413
7.385
7.260
7.220
7.199
7.181
7.162
7.143

— 5.033

— 1.548

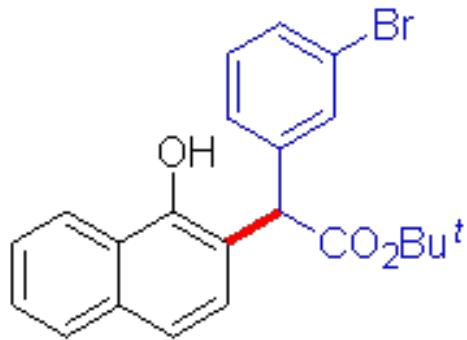


4ai

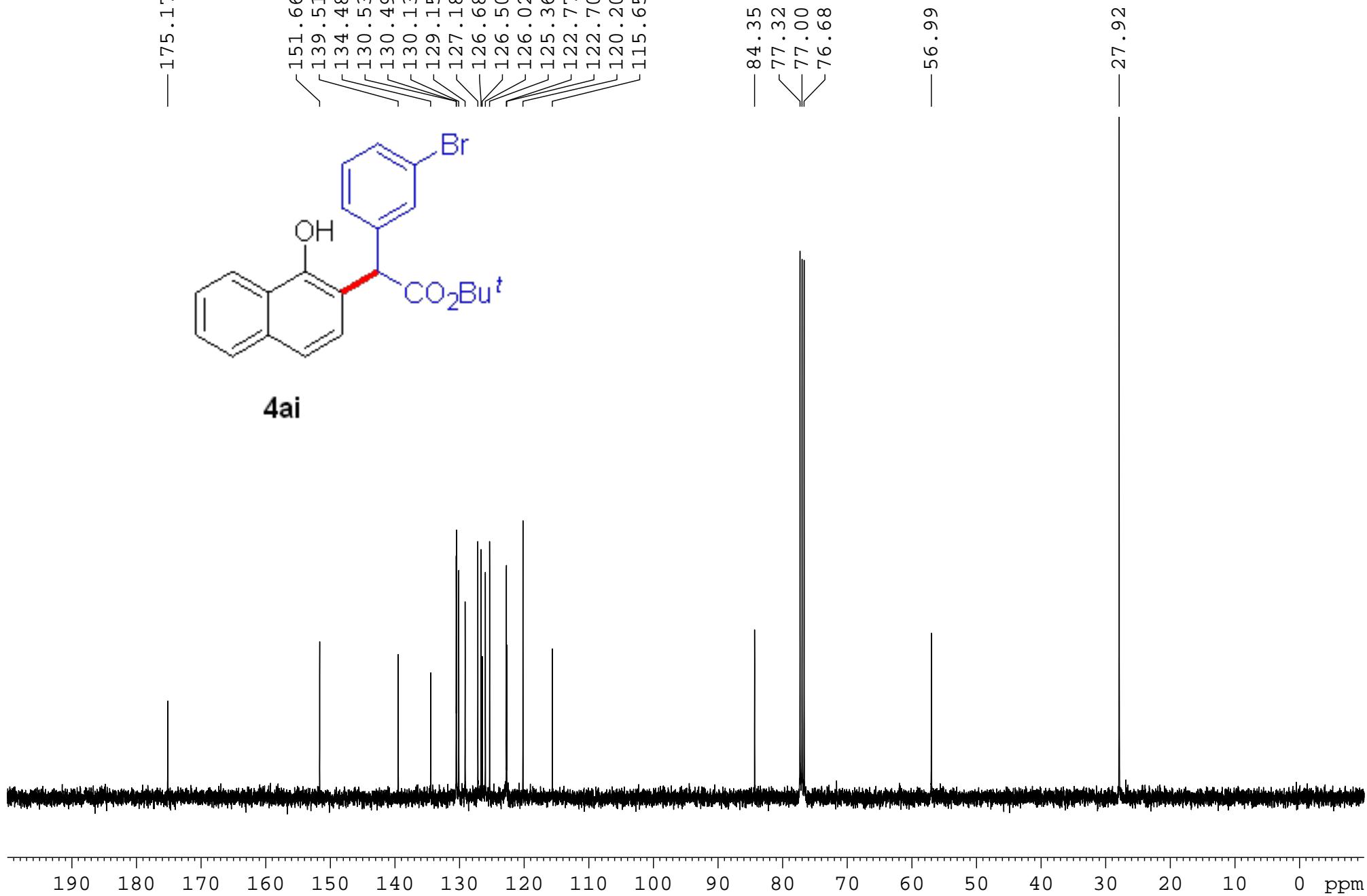


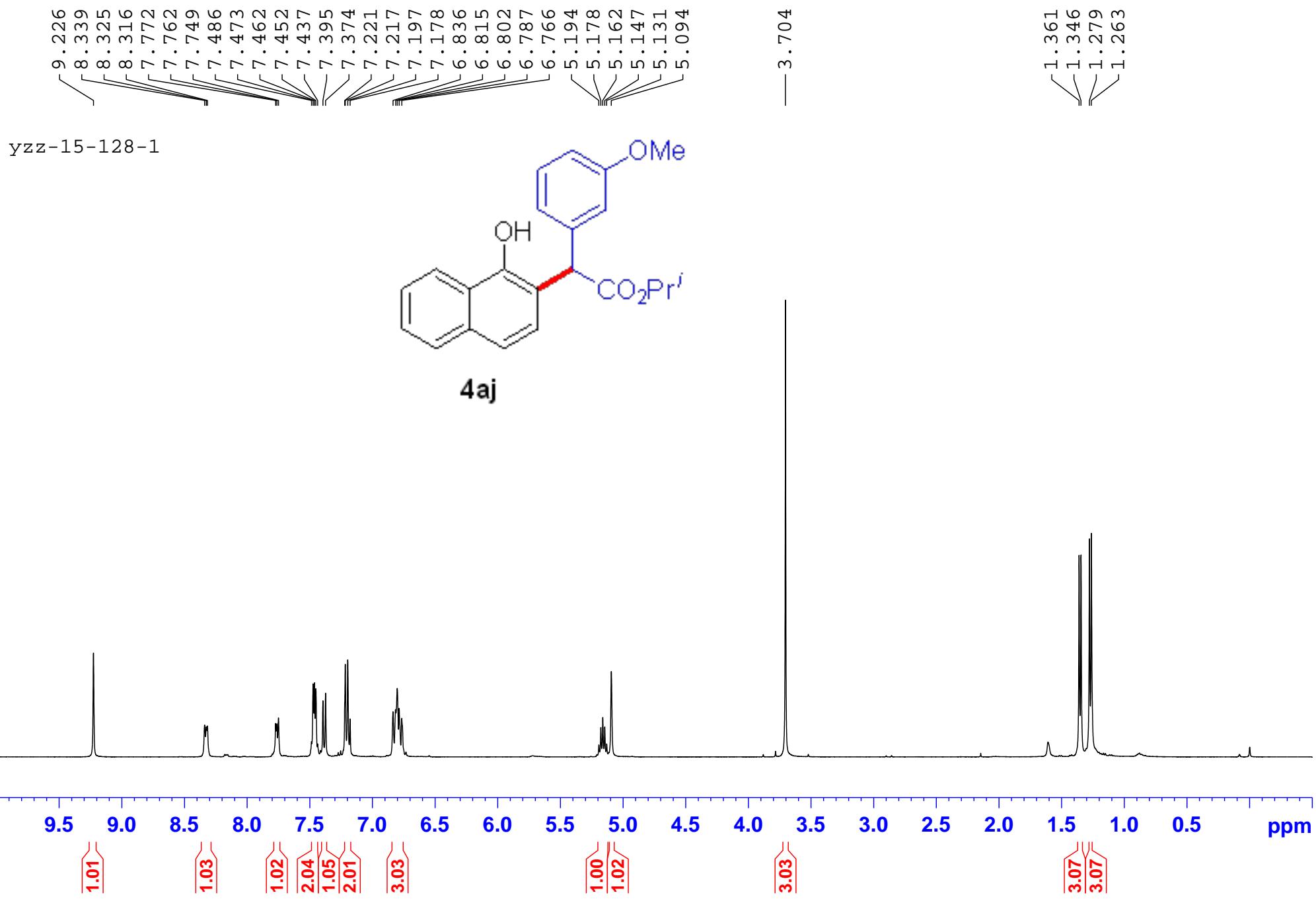
yzz-15-137-1-c

— 175.17



4ai





yzz-15-128-1-c

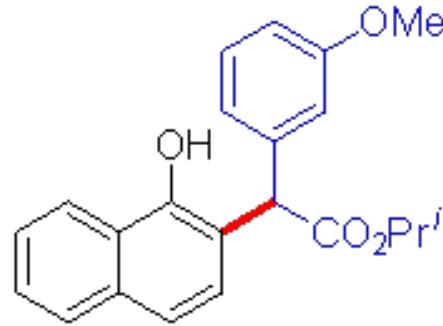
— 175.67

— 159.77
— 151.63
— 138.40
— 134.43
— 129.64
— 129.26
— 127.13
— 126.54
— 126.43
— 125.25
— 122.74
— 120.06
— 119.71
— 115.88
— 113.63
— 112.44

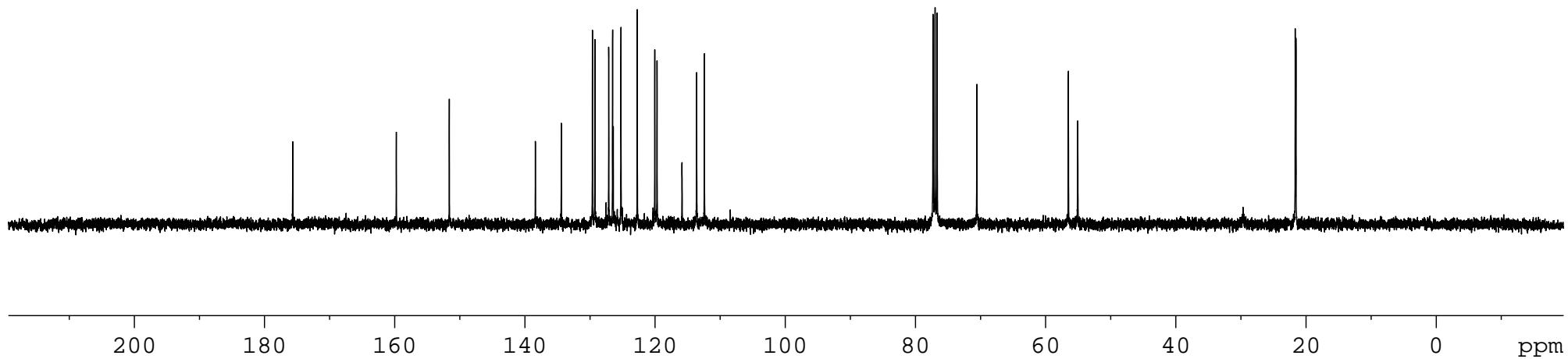
— 77.32
— 77.00
— 76.68
— 70.58

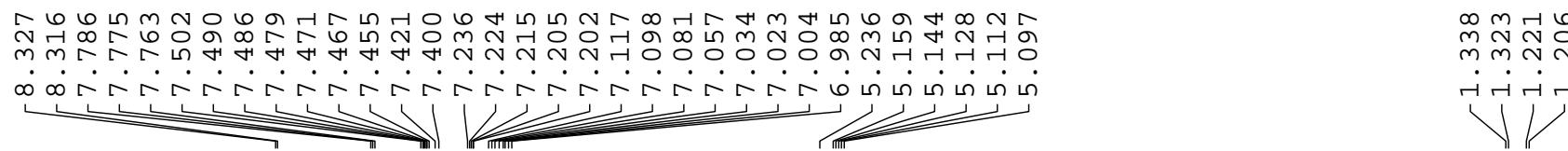
— 56.51
— 55.09

— 21.67
— 21.56

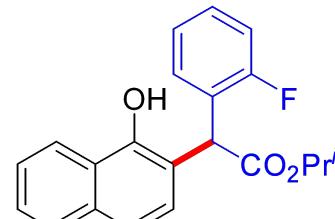


4aj

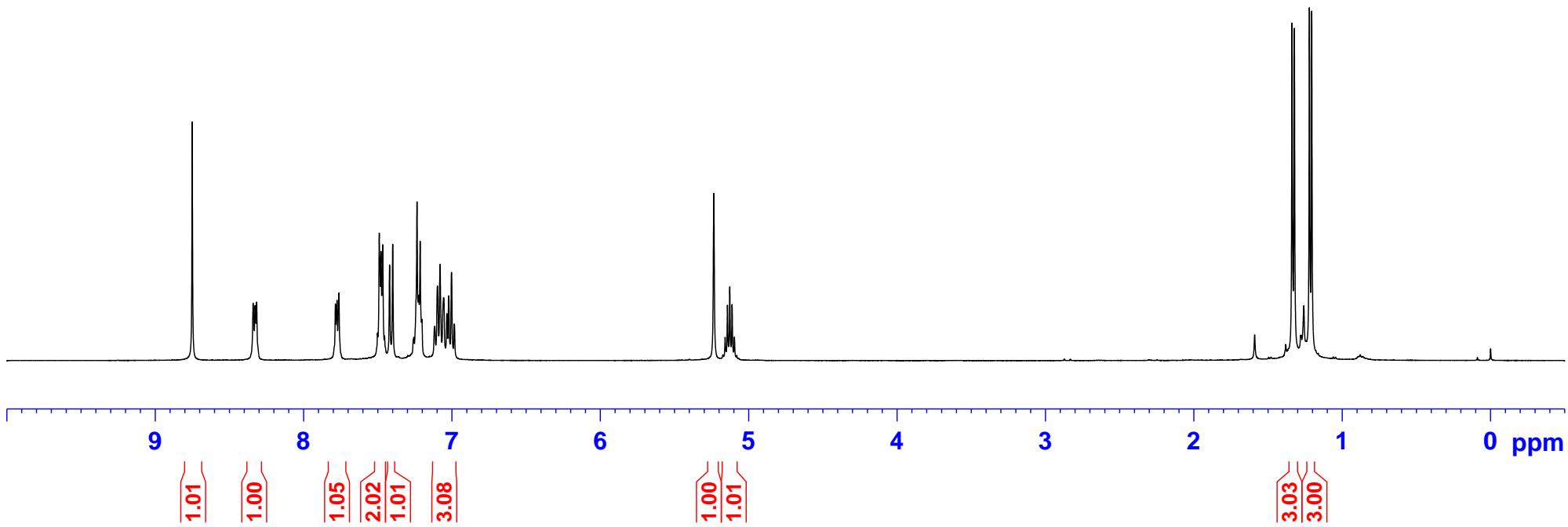




yzz-15-130-1



4ak



yzz-15-130-1-c

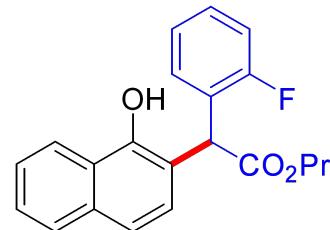
— 174.94

— 161.91
— 159.46
/ 151.85
/ 134.53
— 129.44
— 129.37
— 129.19
— 127.25
— 126.71
— 126.20
— 125.40
— 124.24
— 124.20
— 123.72
— 123.57
— 122.63
— 120.43
— 115.47
— 115.25
— 113.79

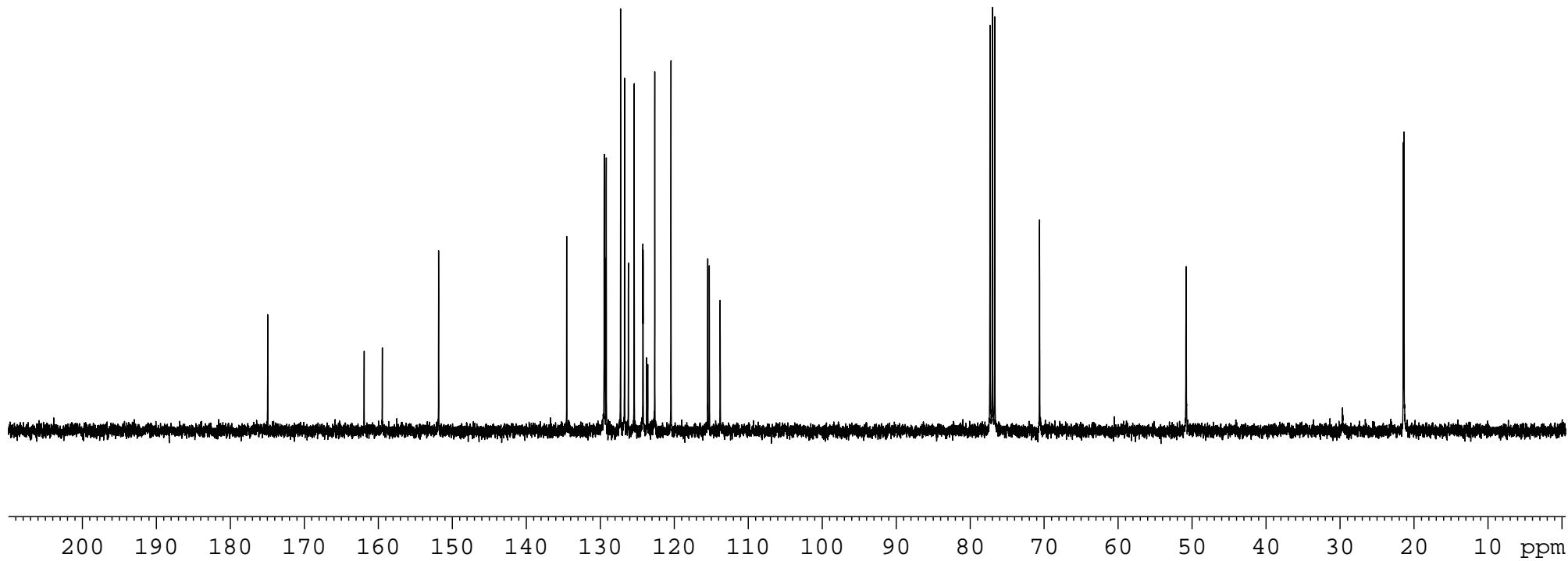
— 77.32
— 77.00
— 76.68
— 70.62

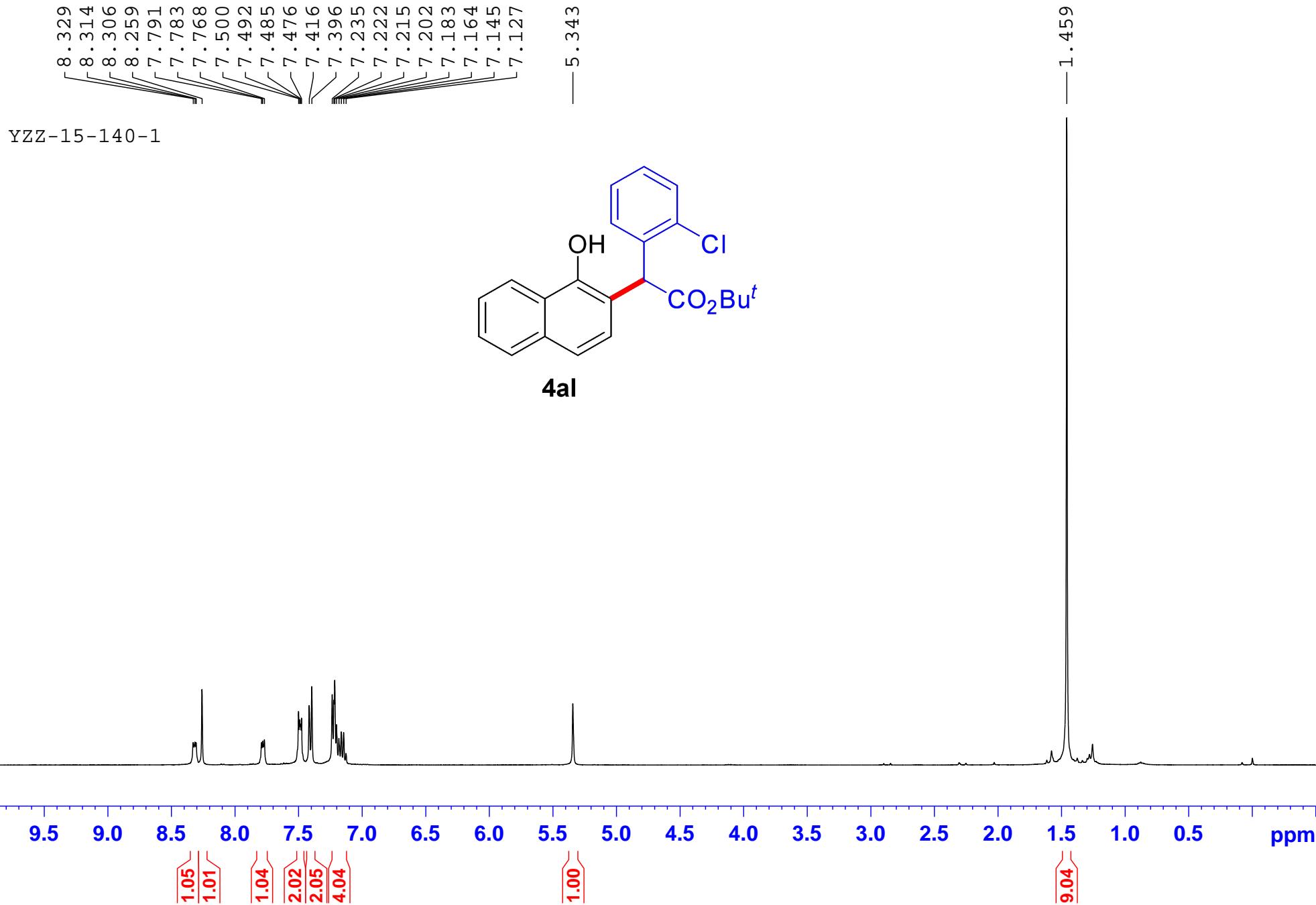
— 50.82

— 21.47
— 21.37

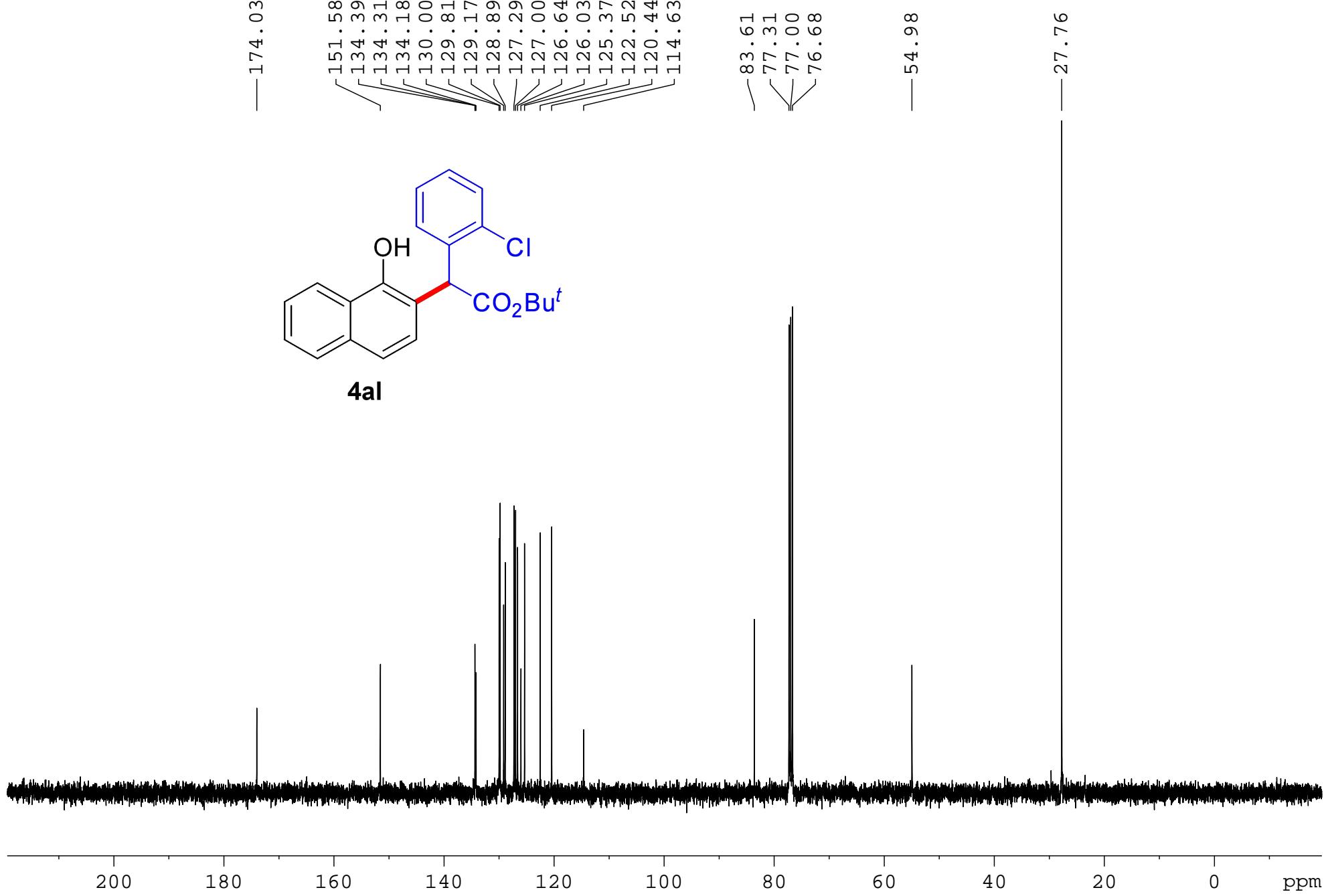


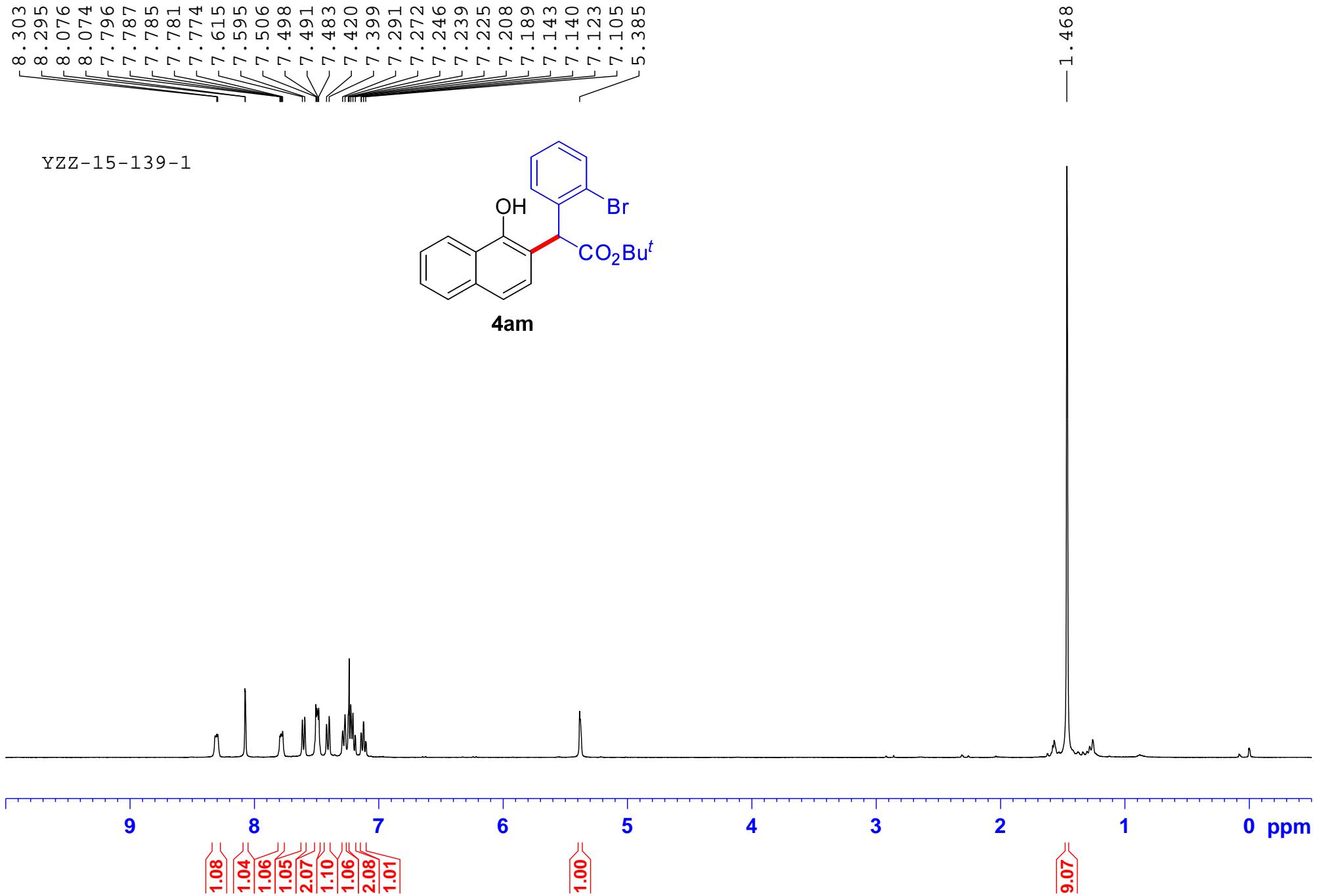
4ak





YZZ-15-140-1-C





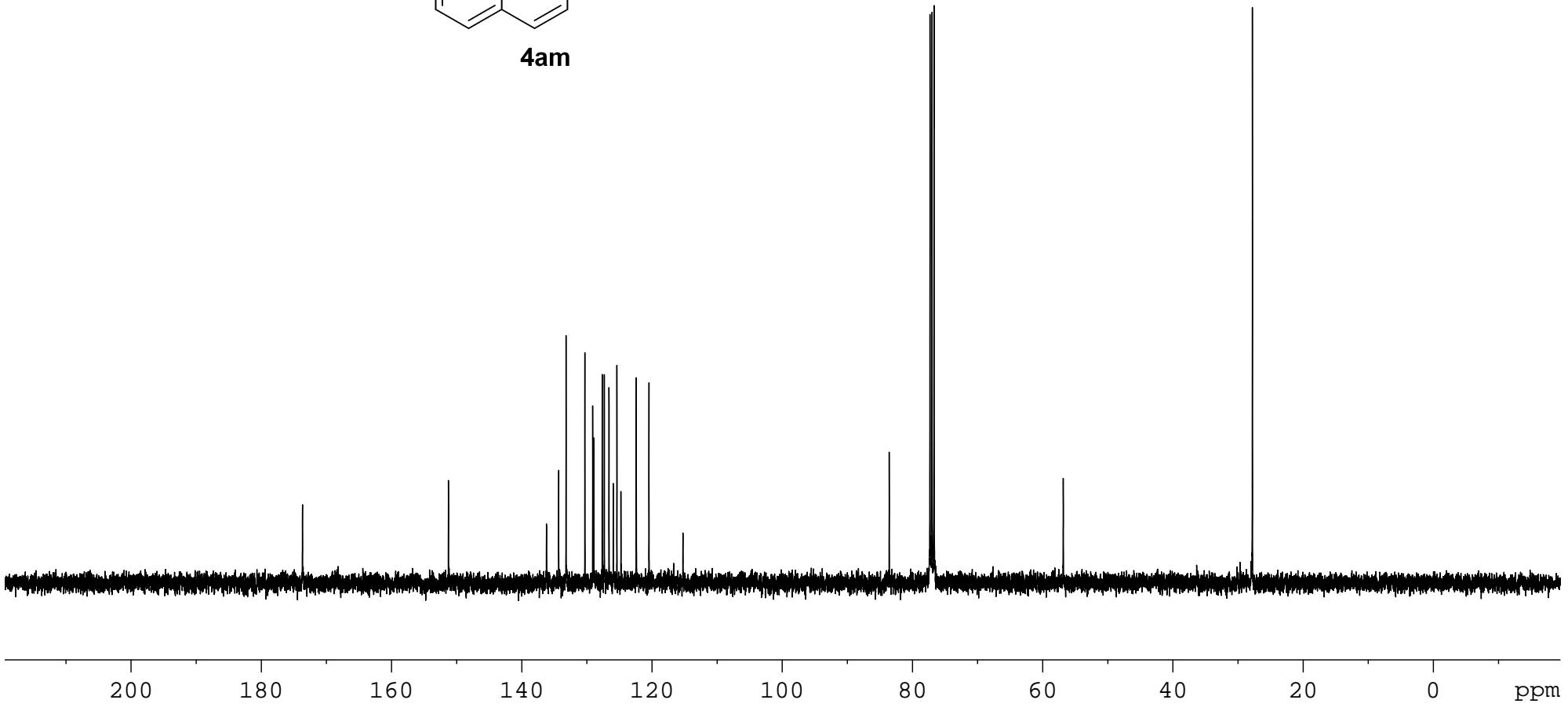
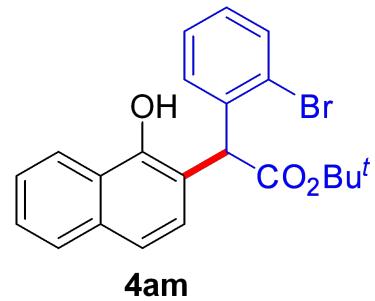
yzz-15-139-1-c

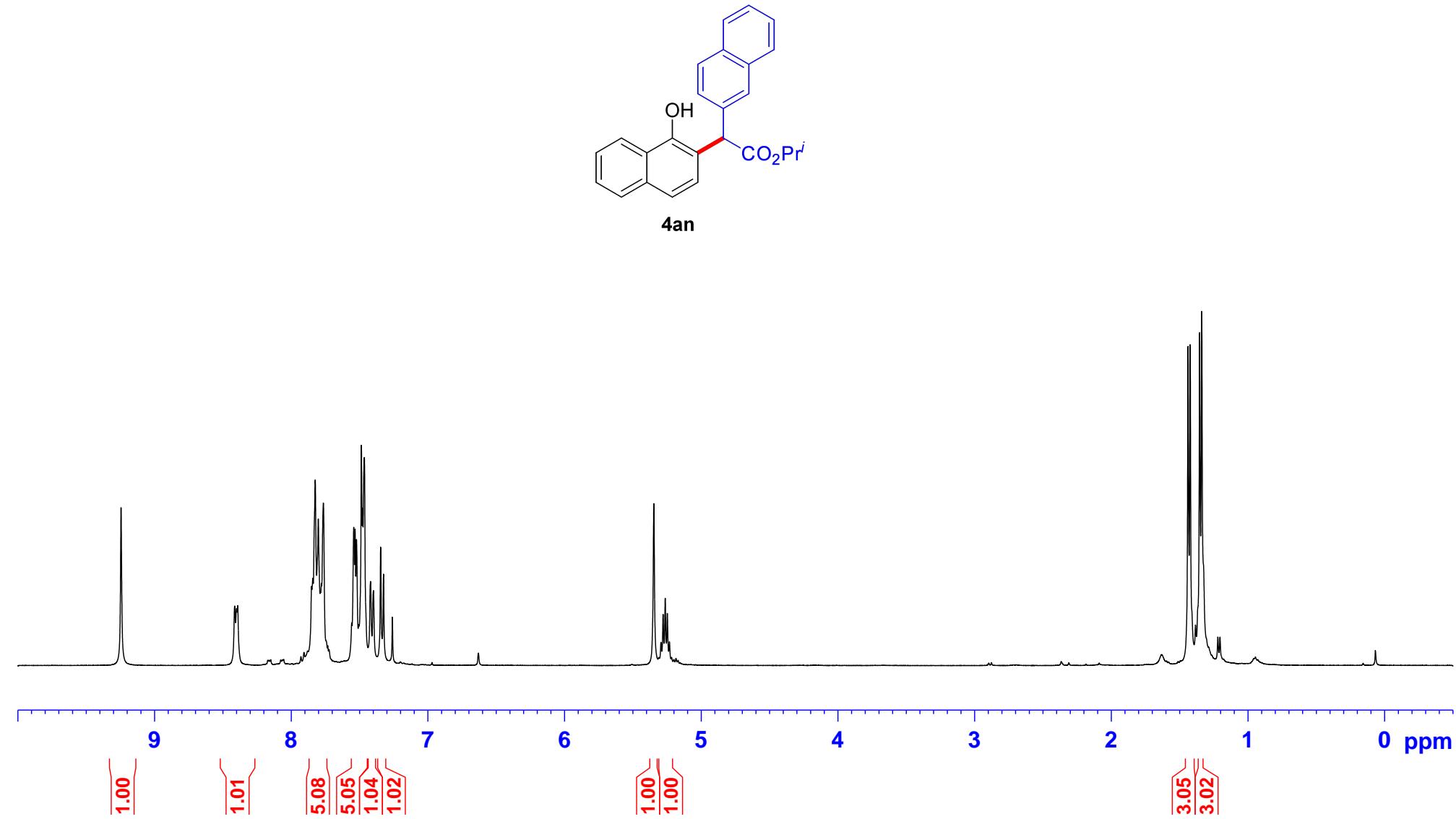
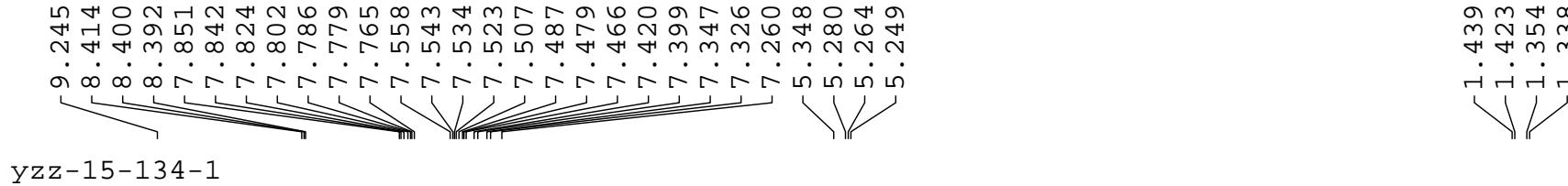
— 173.69

151.28
136.22
134.34
133.22
130.32
129.12
128.96
127.66
127.33
126.62
125.97
125.40
124.78
122.42
120.47
115.26

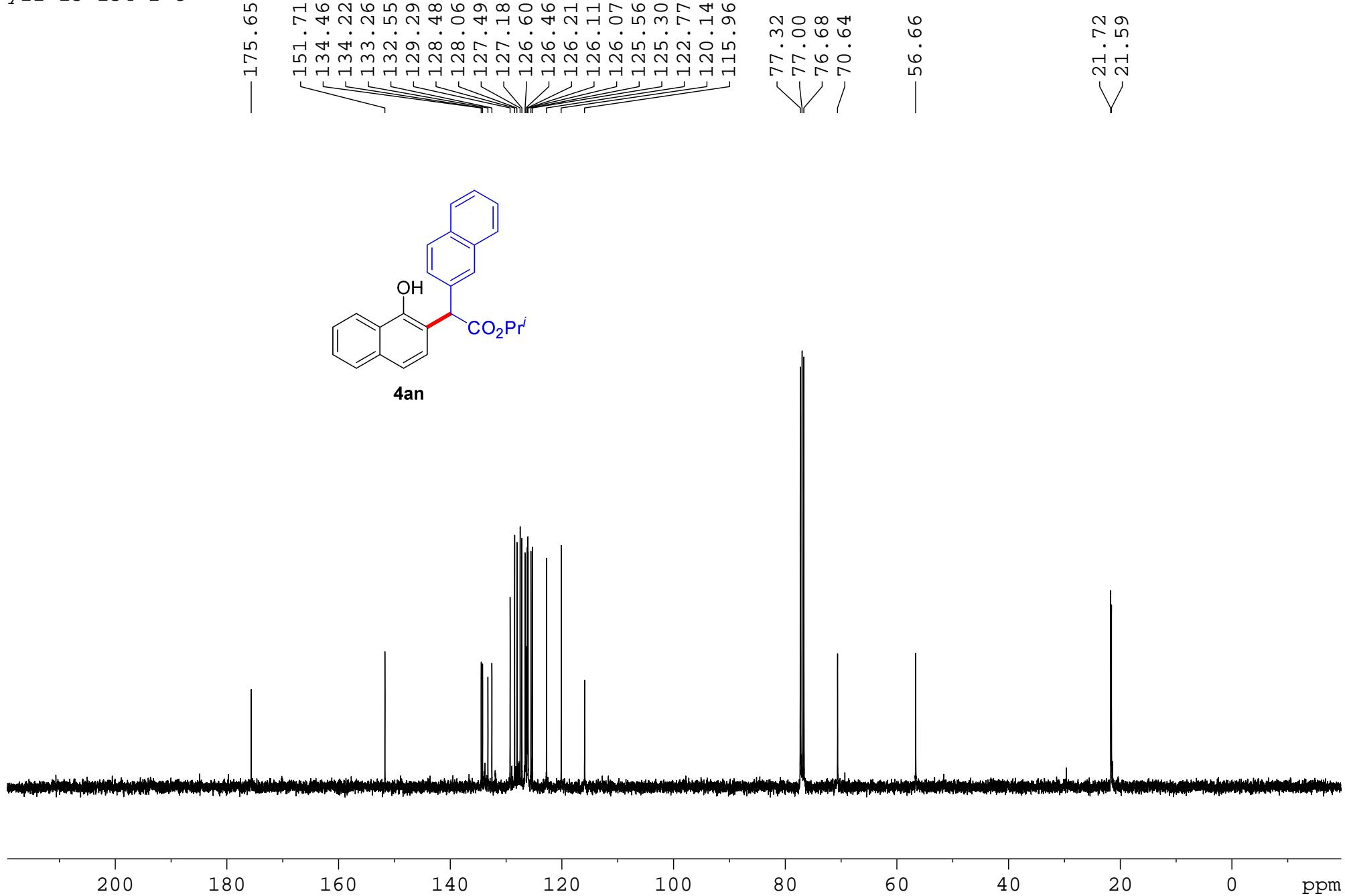
— 56.85

— 27.80

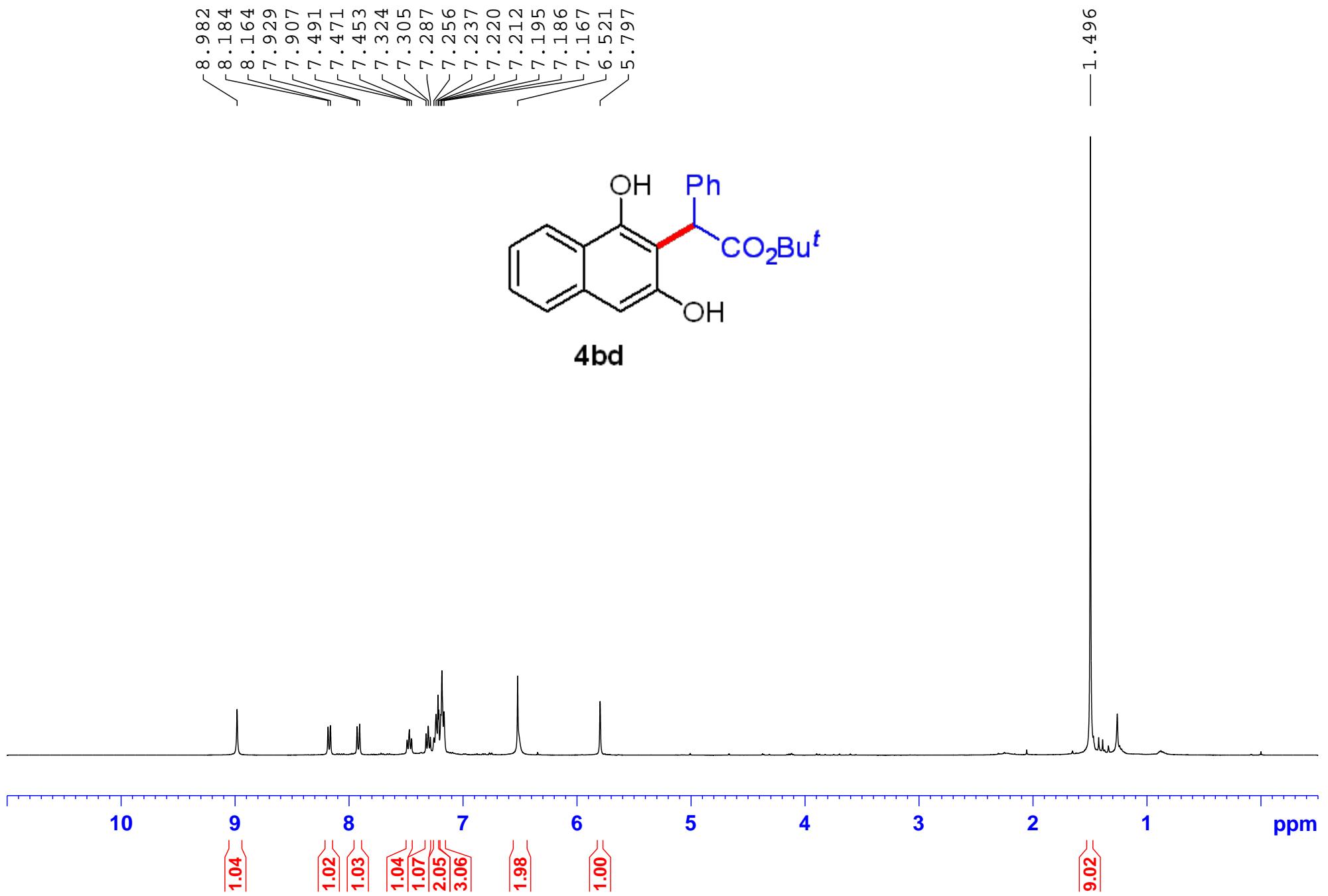




yzz-15-134-1-c



YZZ-16-001-3



yzz-16-001-3-c

— 176.32

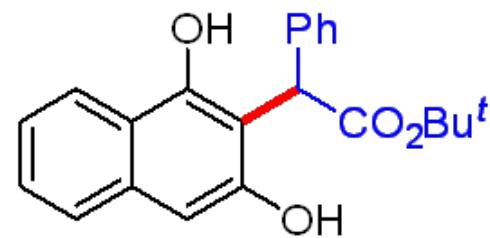
— 154.33
— 153.07

— 137.51
— 134.23
— 128.48
— 127.60
— 127.37
— 127.03
— 122.71
— 122.31
— 121.63
— 120.96
— 107.63
— 103.12

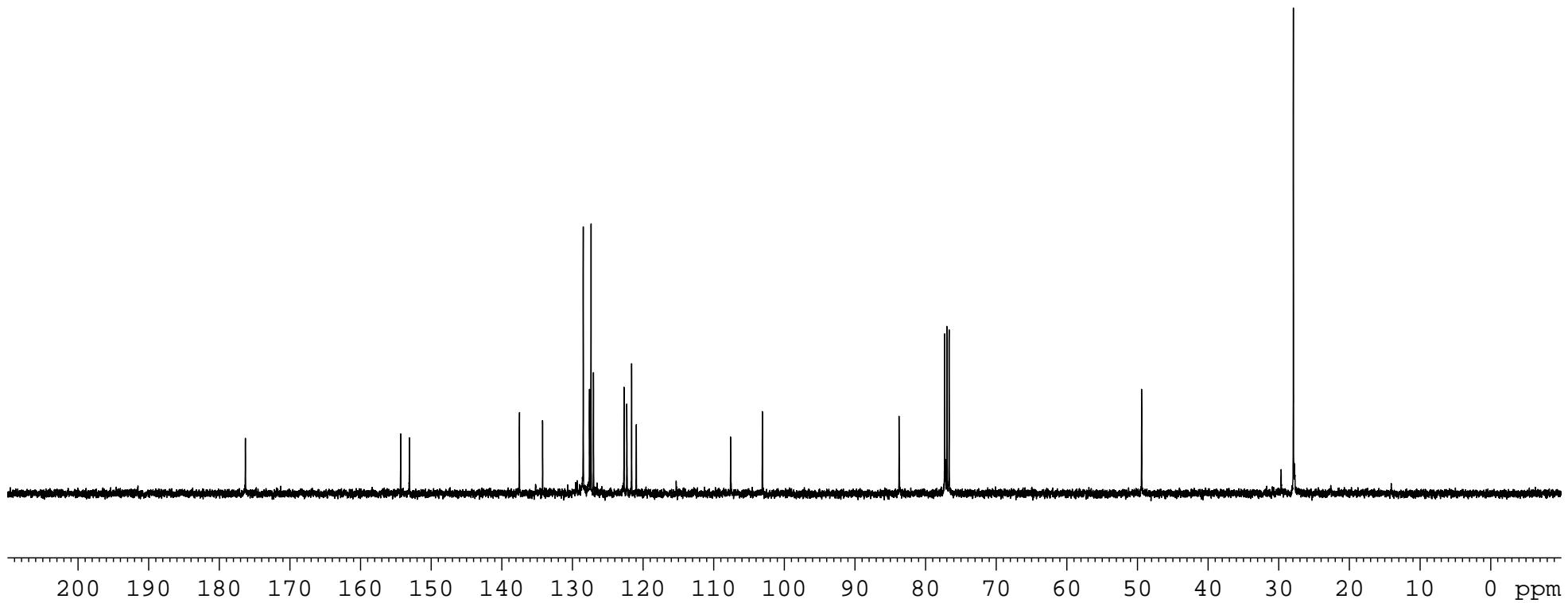
— 83.77
— 77.32
— 77.20
— 77.00
— 76.68

— 49.43

— 27.94



4bd



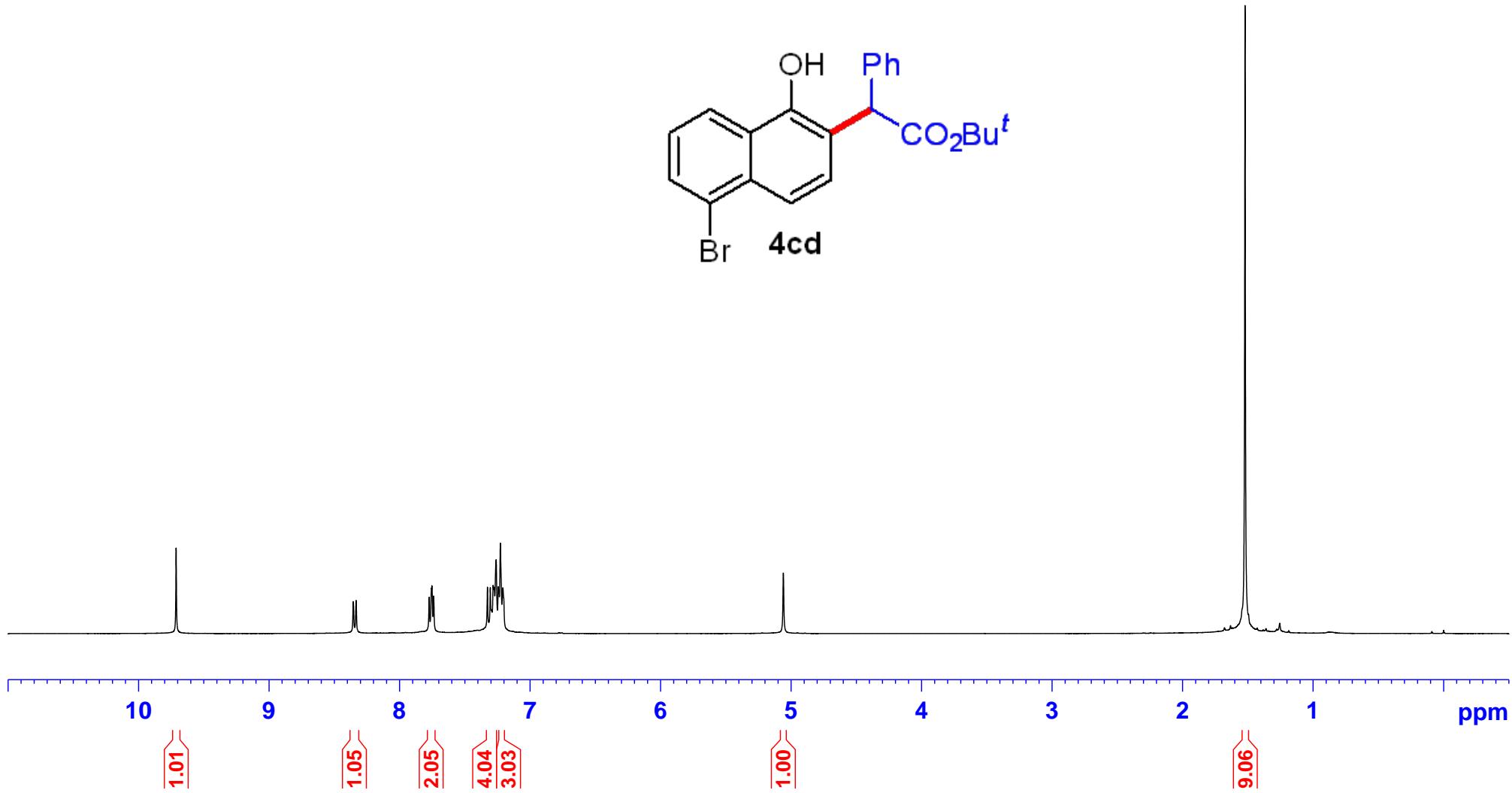
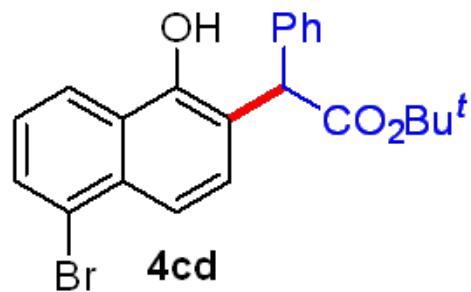
YZZ-15-012-1

— 9.713

8.355
8.334
7.774
7.756
7.753
7.740
7.327
7.305
7.284
7.277
7.261
7.244
7.240
7.228
7.209

— 5.060

— 1.522



YZZ-15-012-1-C

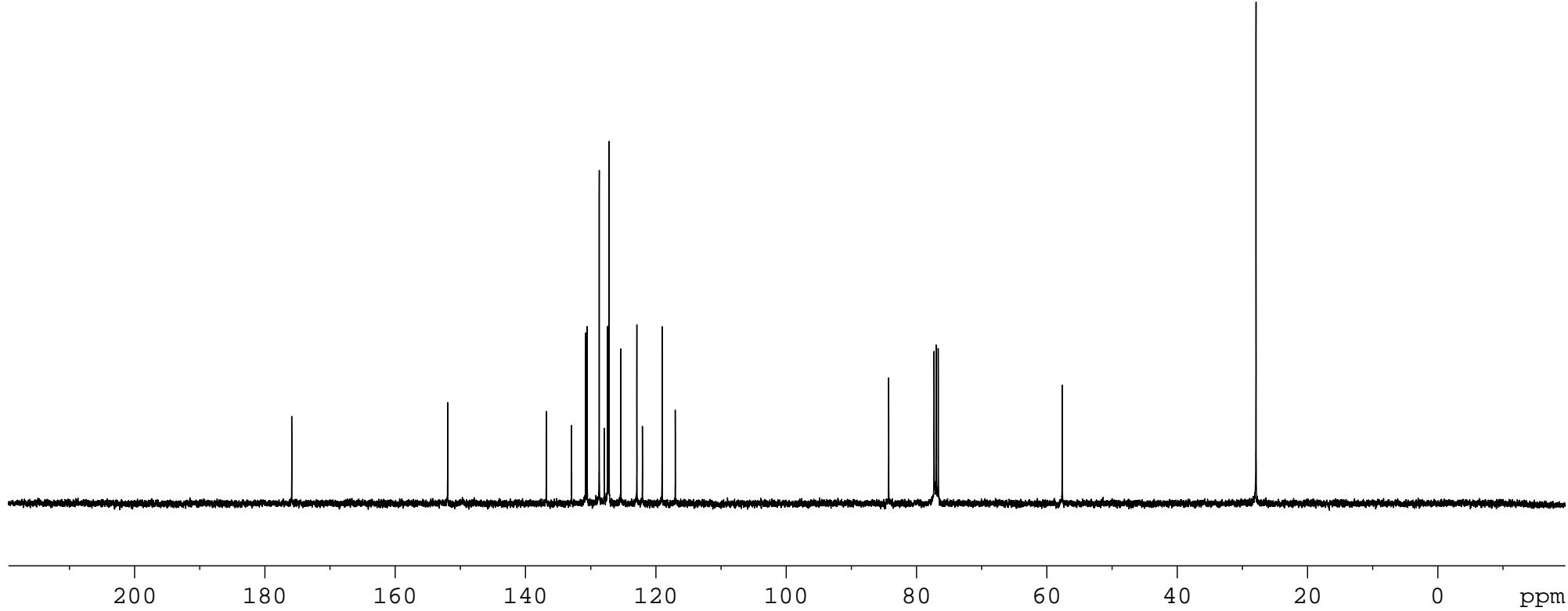
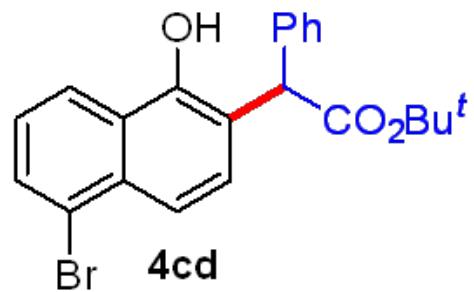
— 175.89

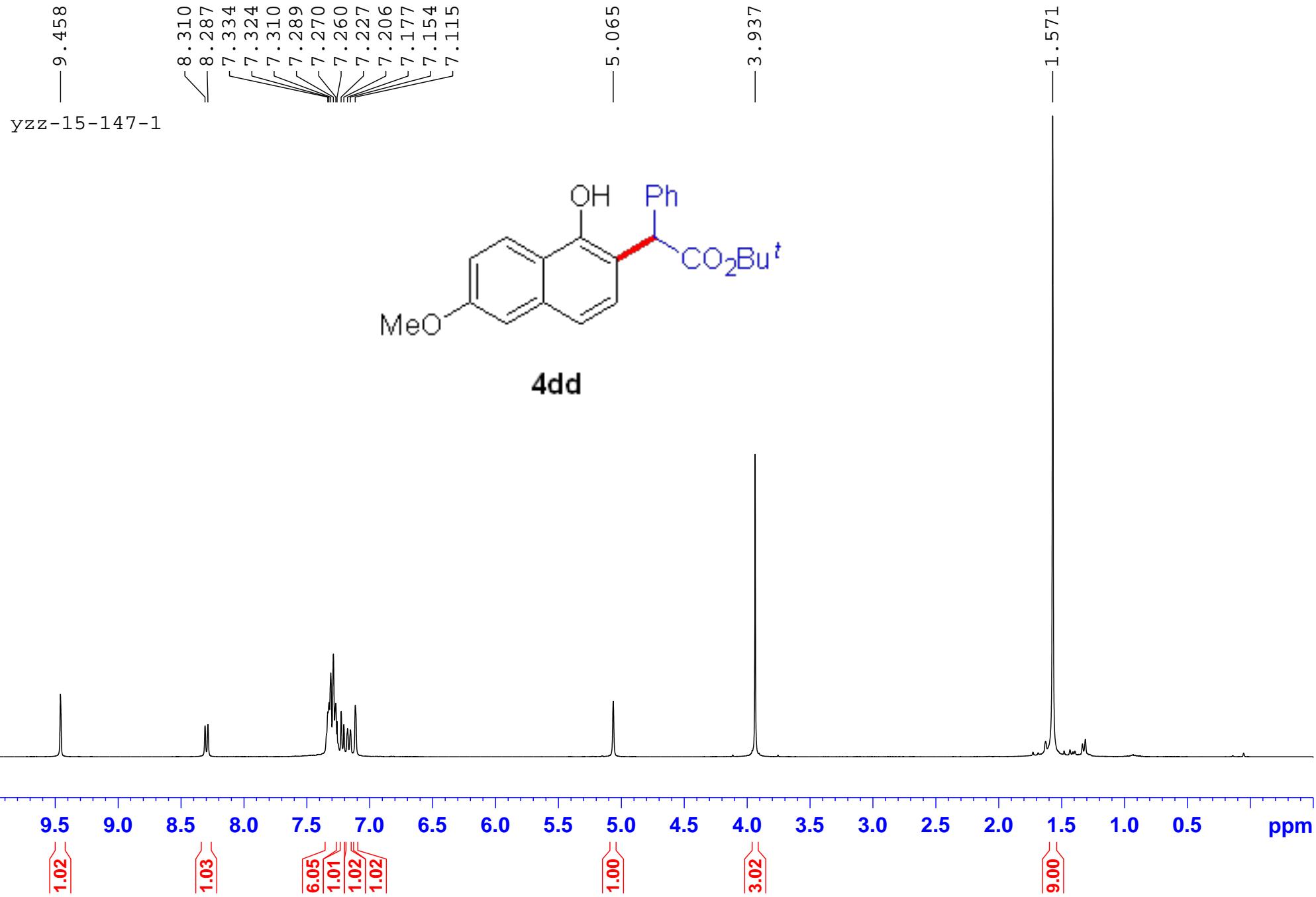
151.97
136.83
132.95
130.78
130.56
128.71
127.91
127.43
127.19
125.39
122.93
122.08
119.06
117.04

84.28
77.32
77.00
76.68

57.65

27.92





yzz-15-147-1-c

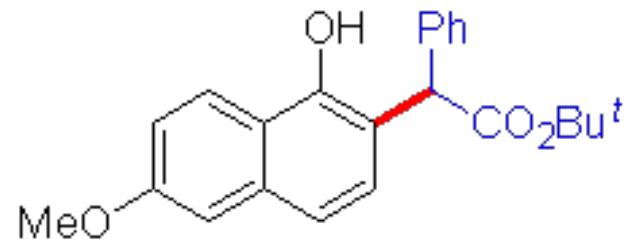
— 176.01

— 158.31
— 152.00
/ 137.36
/ 135.86
/ 130.26
/ 128.60
/ 127.26
/ 124.64
/ 121.76
/ 118.81
/ 117.65
/ 114.24
/ 105.26

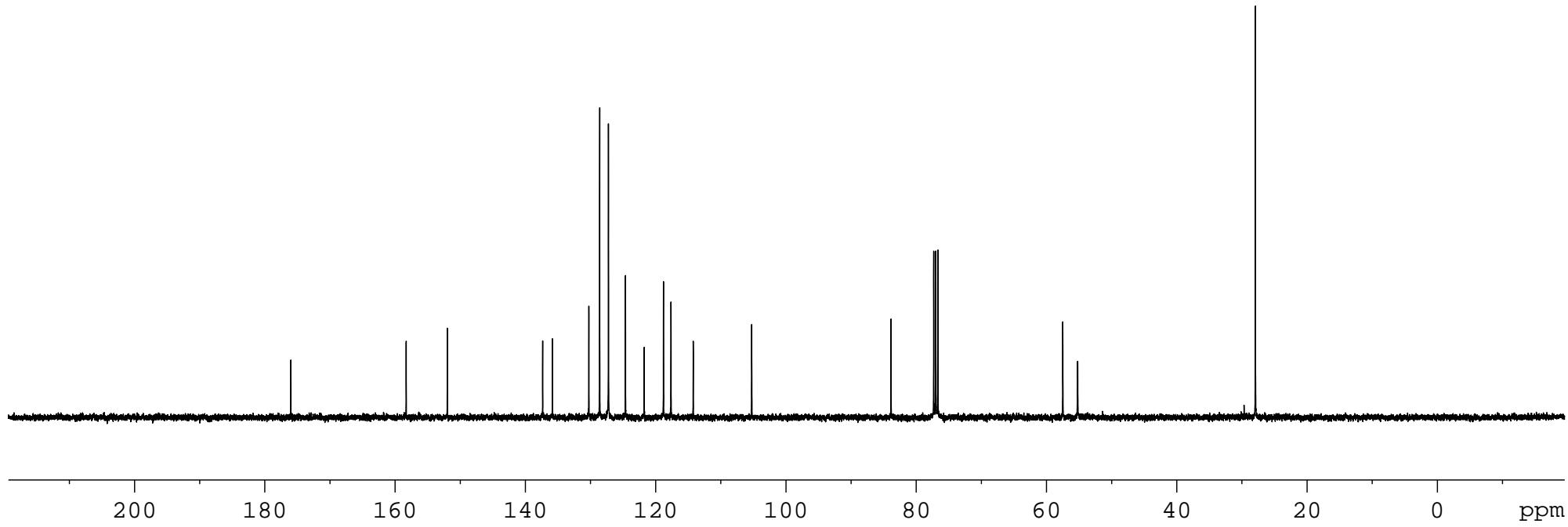
— 83.87
— 77.32
— 77.00
— 76.68

— 57.50
— 55.24

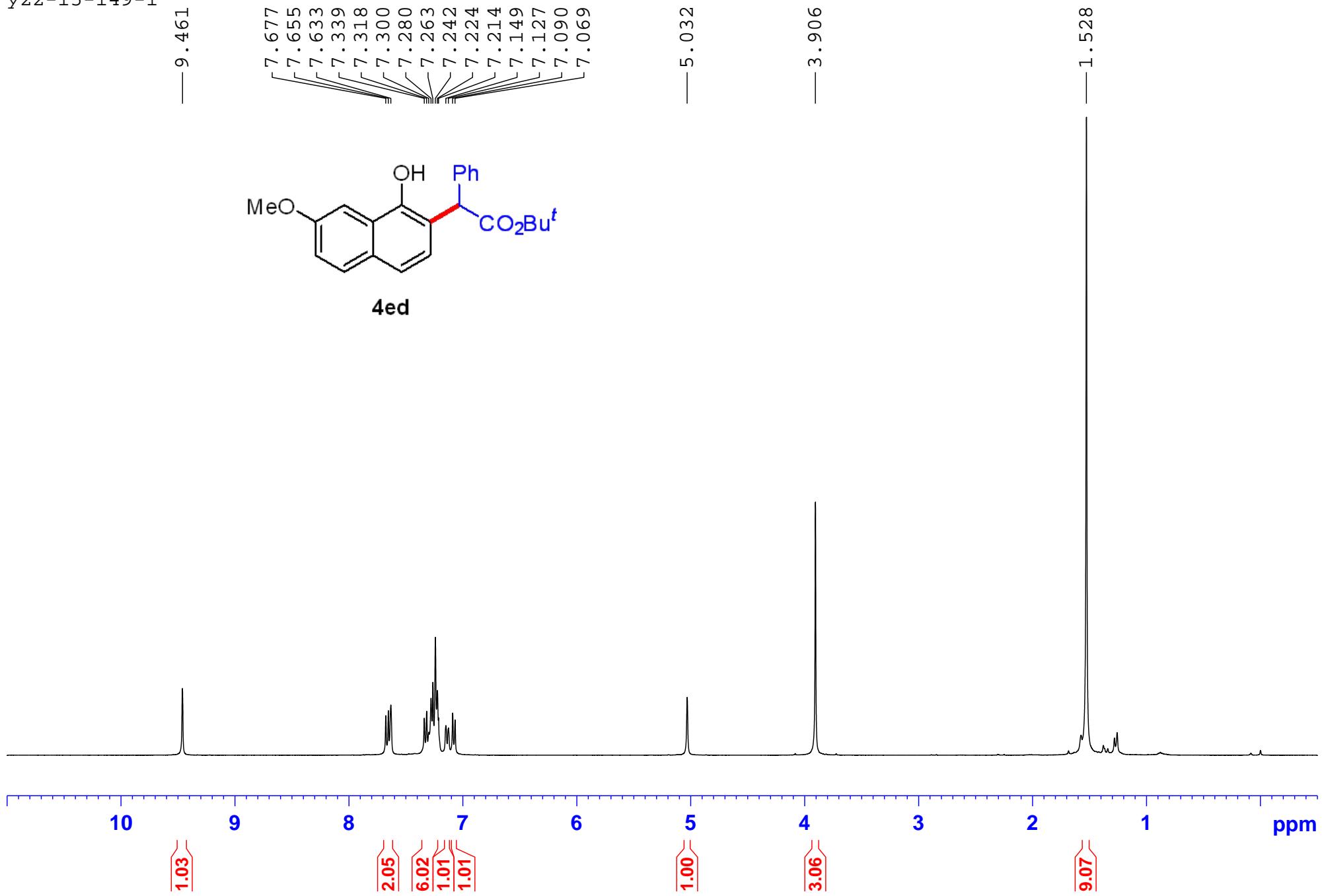
— 27.93



4dd



yzz-15-149-1



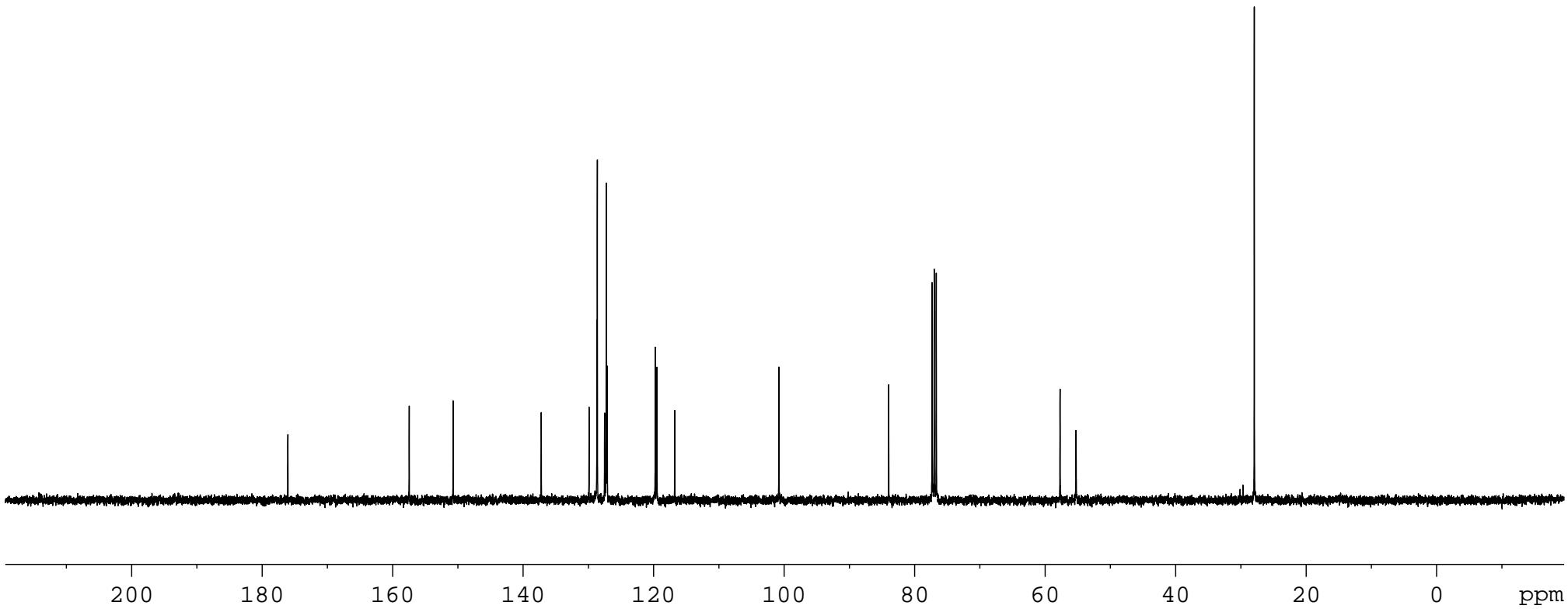
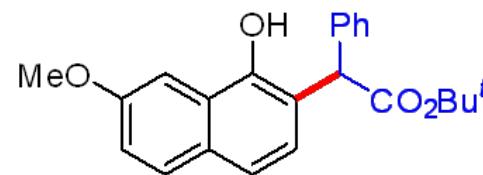
yzz-15-149-1-c

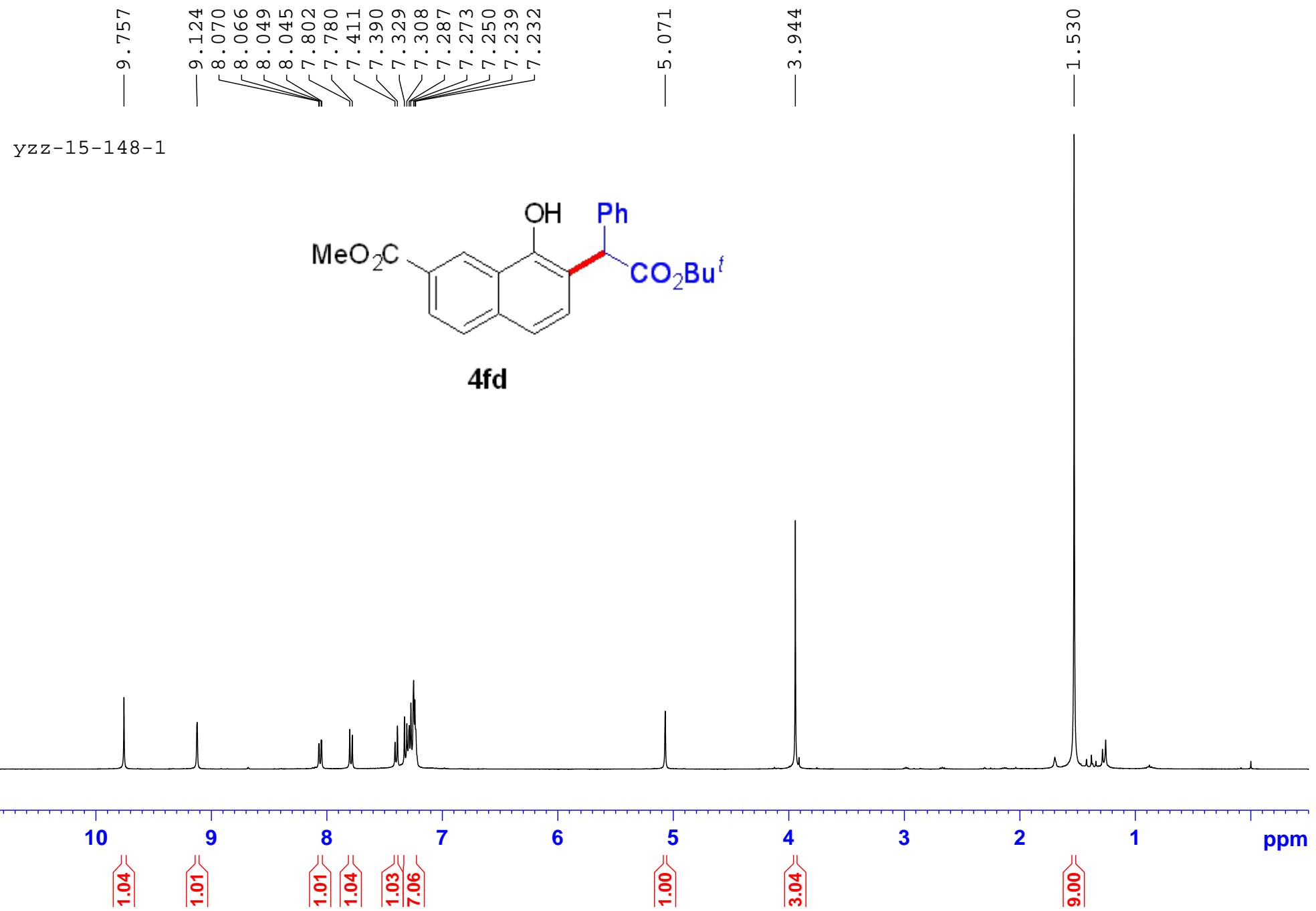
— 176.05

— 157.45
— 150.73
— 137.22
— 129.90
— 128.70
— 128.62
— 127.52
— 127.27
— 127.23
— 127.11
— 119.75
— 119.50
— 116.77
— 100.81

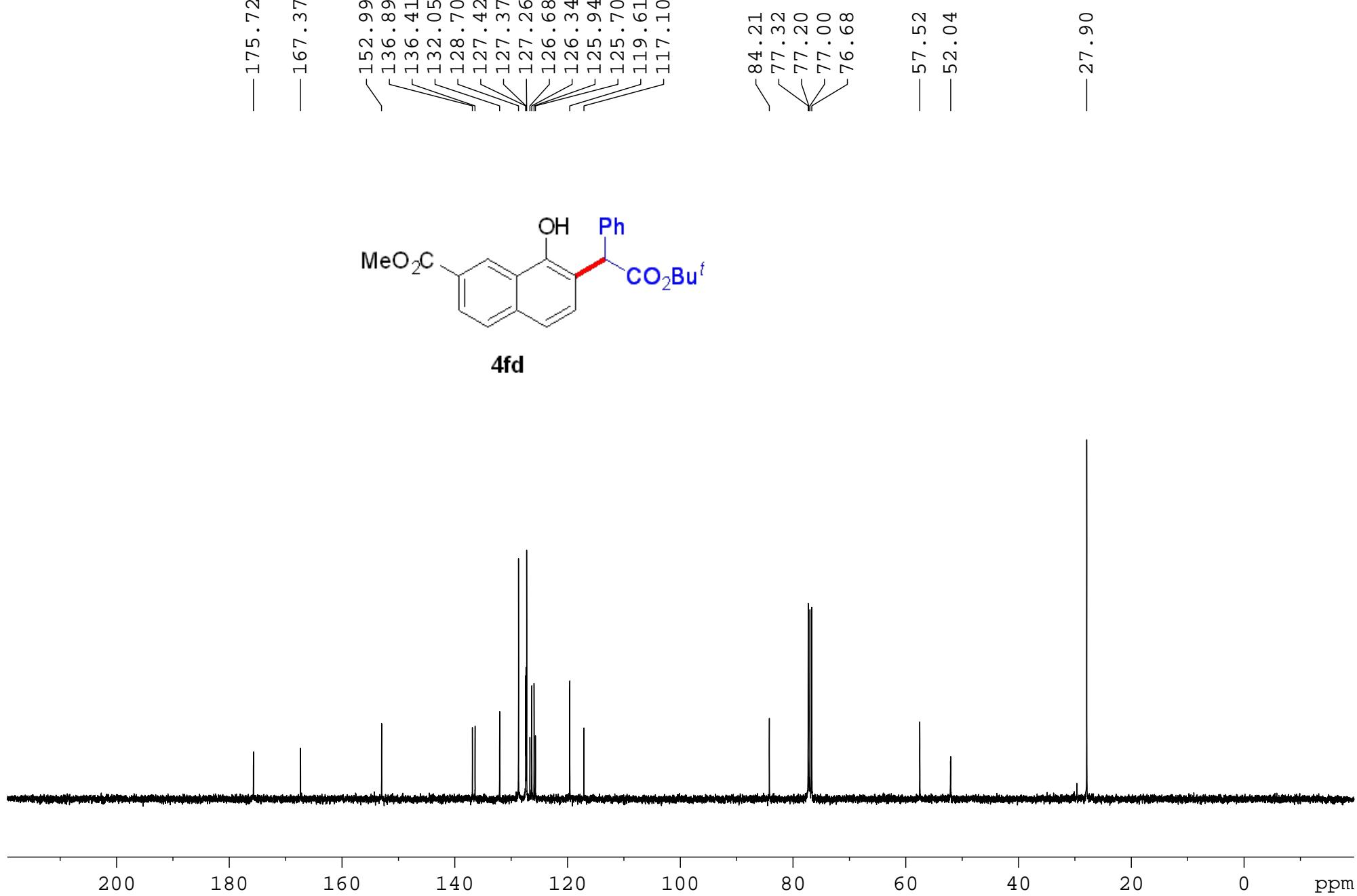
— 83.96
— 77.32
— 77.00
— 76.68
— 57.71
— 55.28

— 27.94





yzz-15-148-1-c



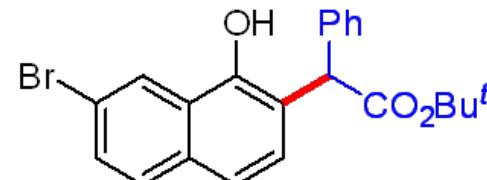
yzz-15-150-1

— 9.568

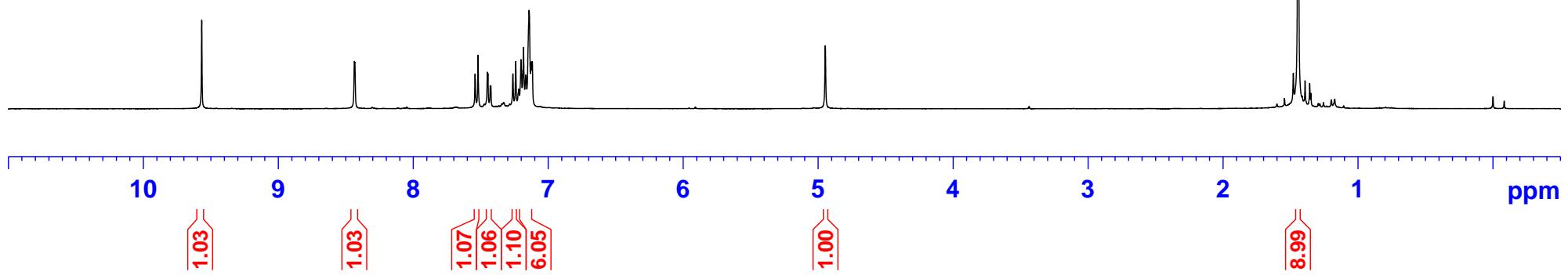
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7.543
7.521
7.452
7.448
7.431
7.426
7.263
7.242
7.220
7.203
7.184
7.167
7.165
7.142
7.126
7.118

— 4.948

— 1.444



4gd



yzz-15-150-1-c

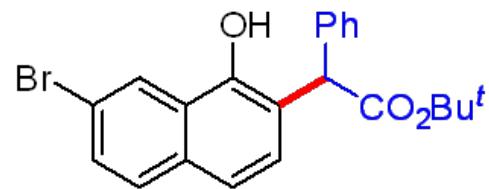
— 175.85

151.05
136.86
132.78
129.96
129.86
128.77
128.71
127.71
127.43
127.20
125.48
119.74
119.37
117.16

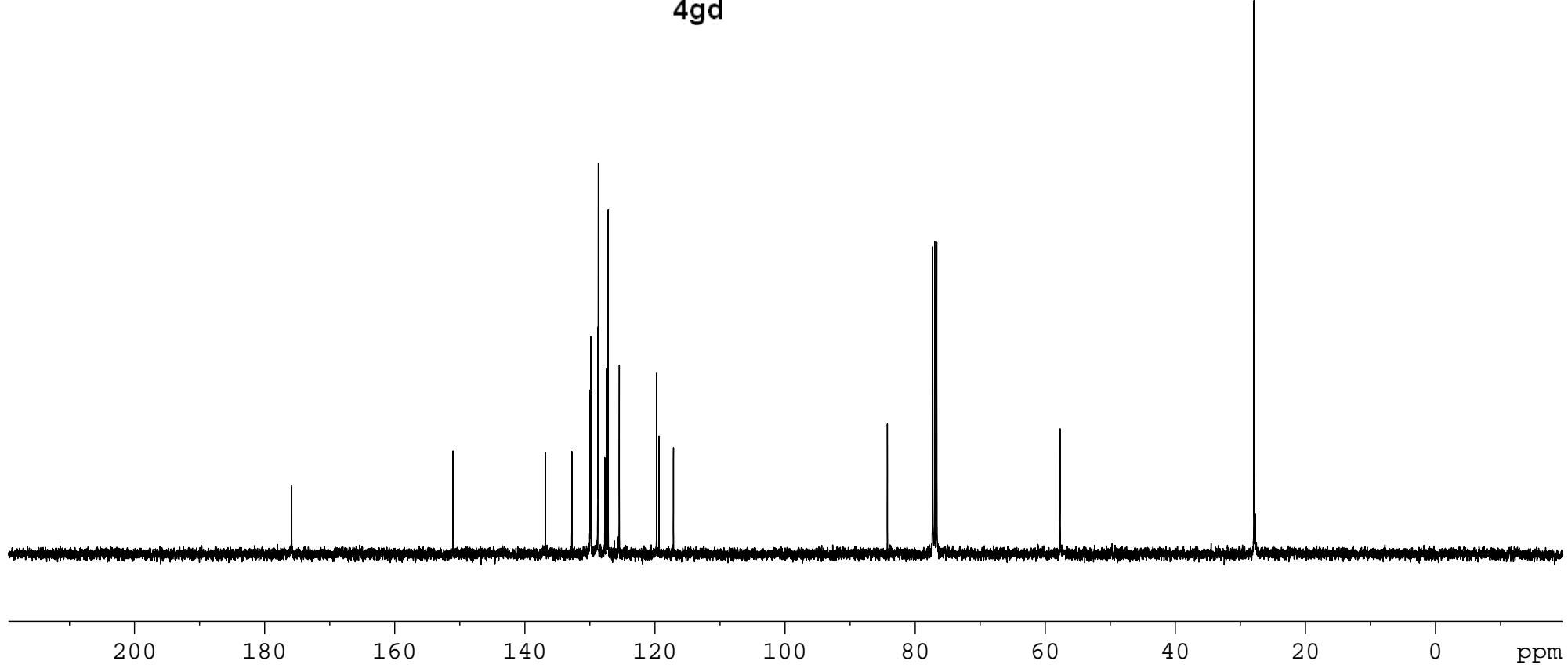
84.27
77.32
77.00
76.68

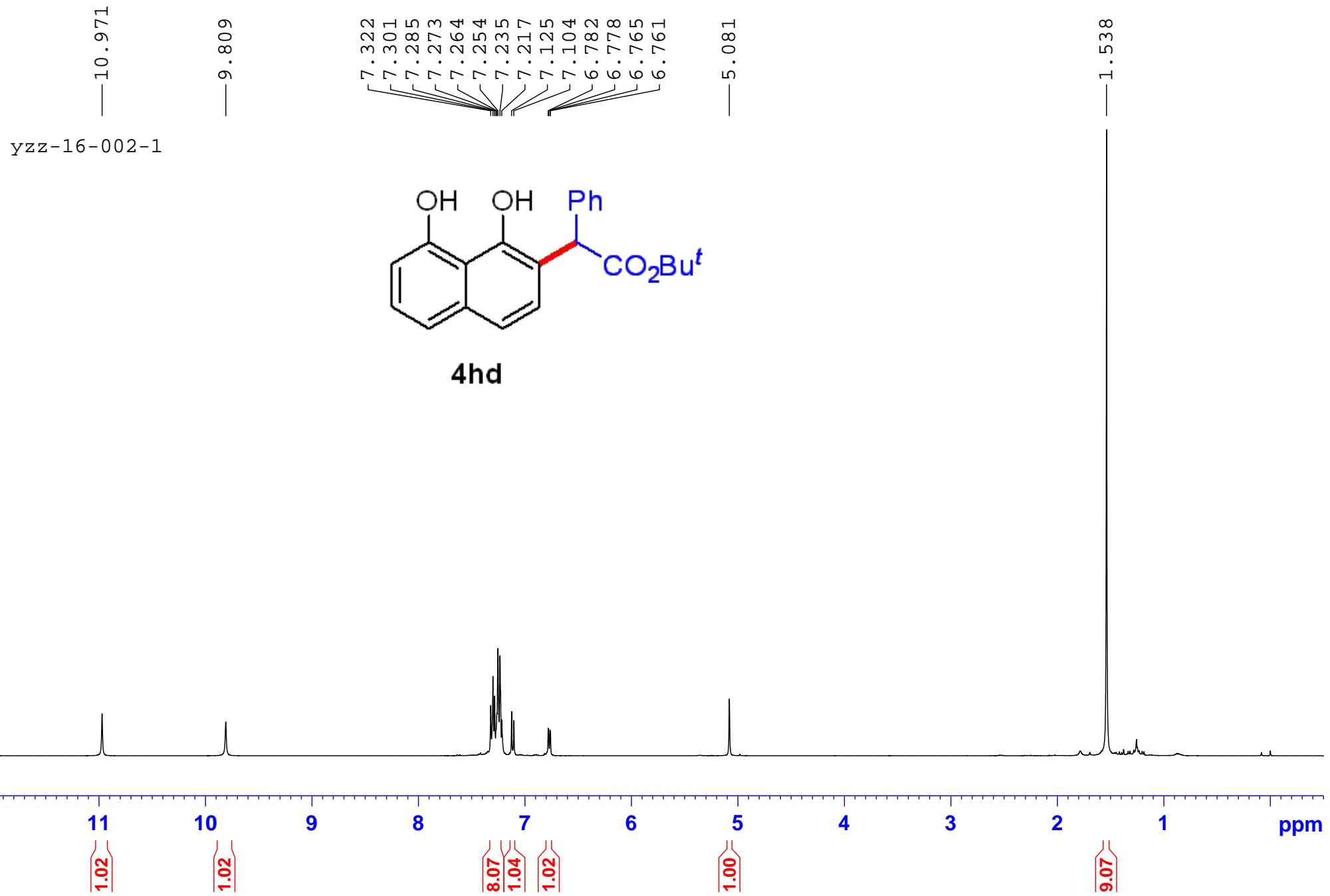
— 57.73

— 27.92



4gd





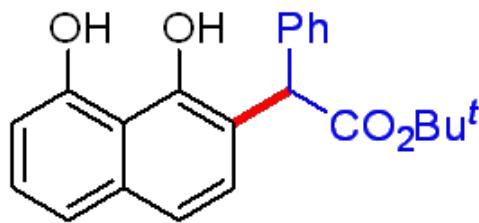
yzz-16-002-1-c

— 176.72

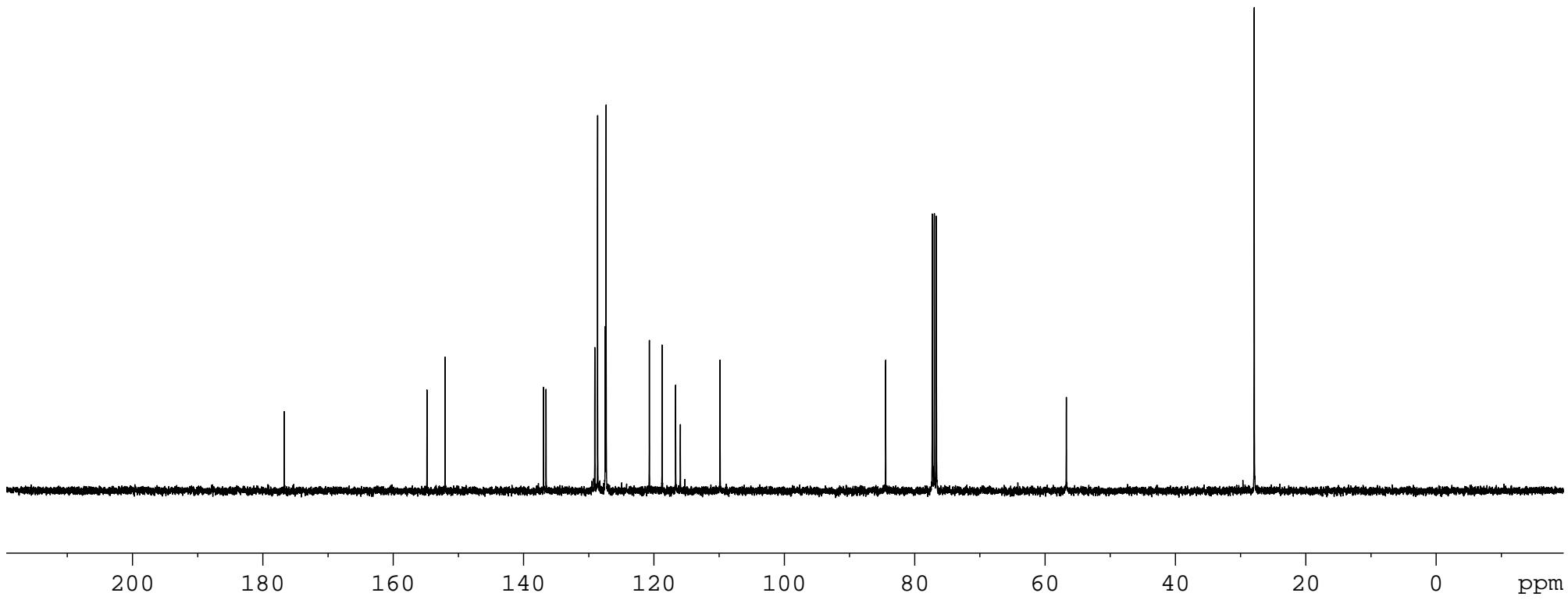
154.84
152.06
136.97
136.57
129.09
128.66
127.50
127.38
127.35
120.72
118.77
116.68
115.96
109.87

— 56.74

— 27.92



4hd

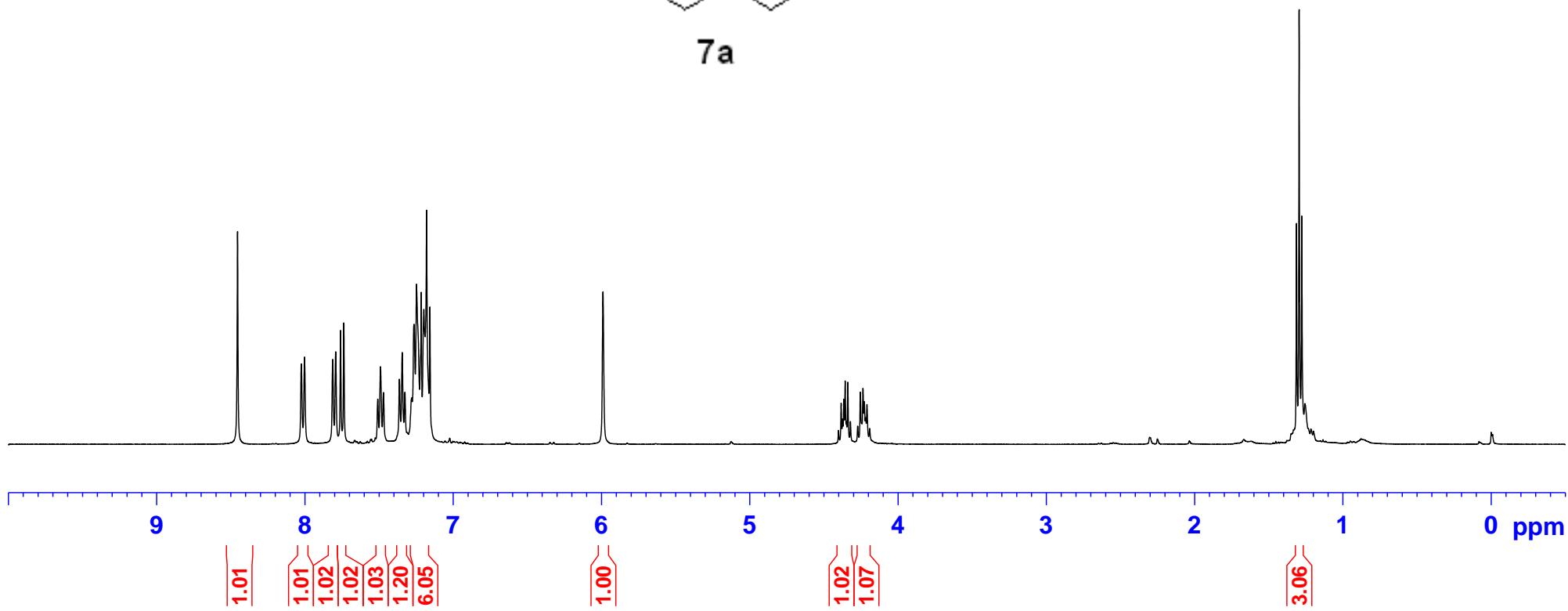
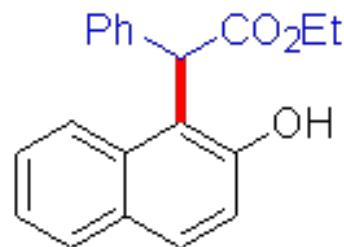


yzz-15-060-2

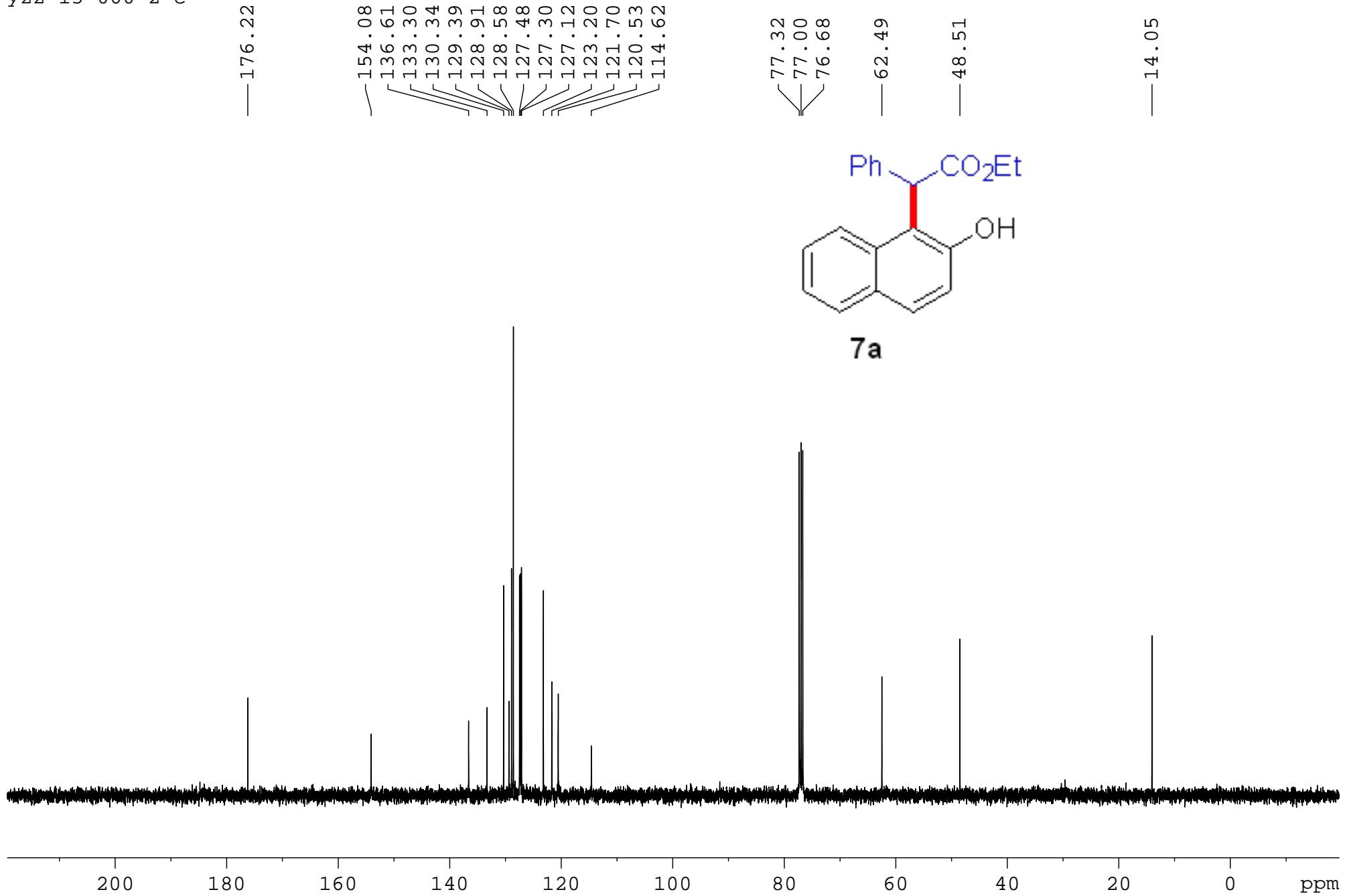
8.454
8.024
8.003
7.813
7.793
7.760
7.738
7.508
7.490
7.470
7.363
7.344
7.326
7.264
7.215
7.247
7.198
7.180
7.157
5.990

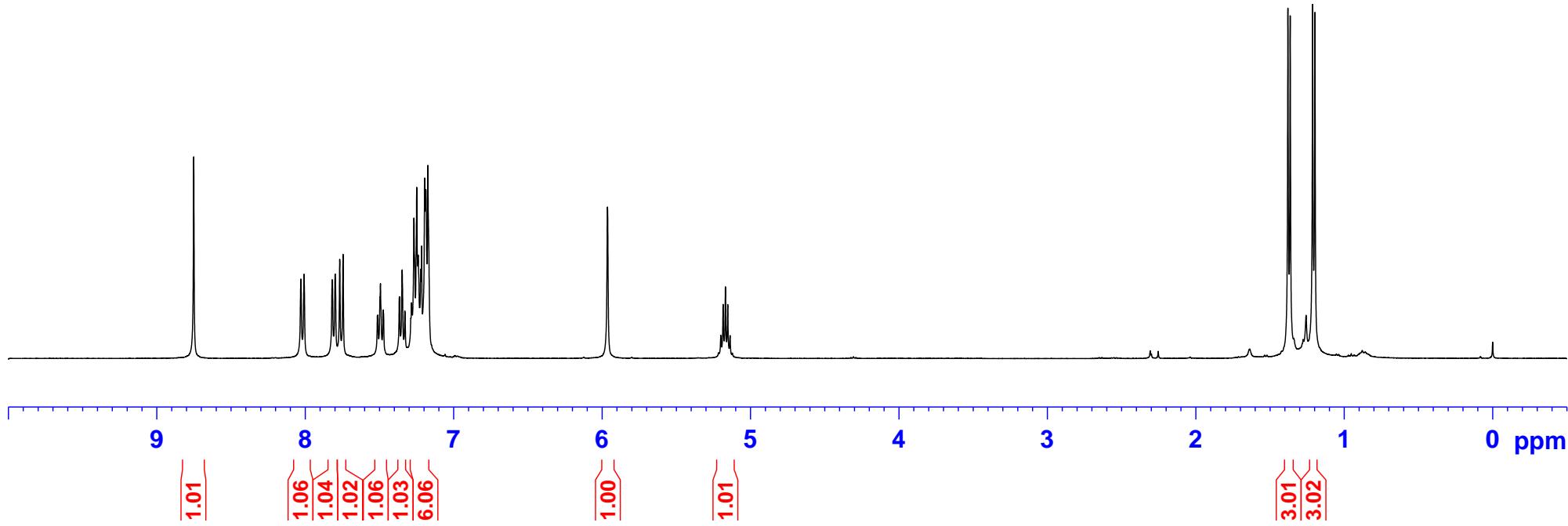
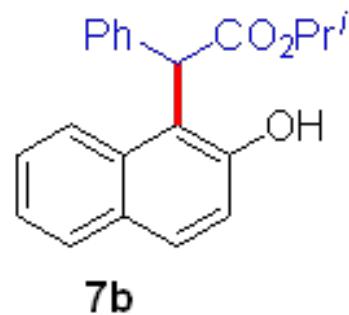
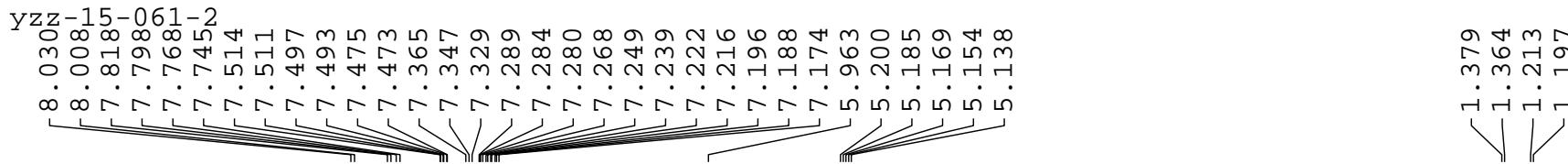
4.384
4.366
4.357
4.339
4.254
4.237
4.228
4.219
4.210

1.313
1.295
1.277

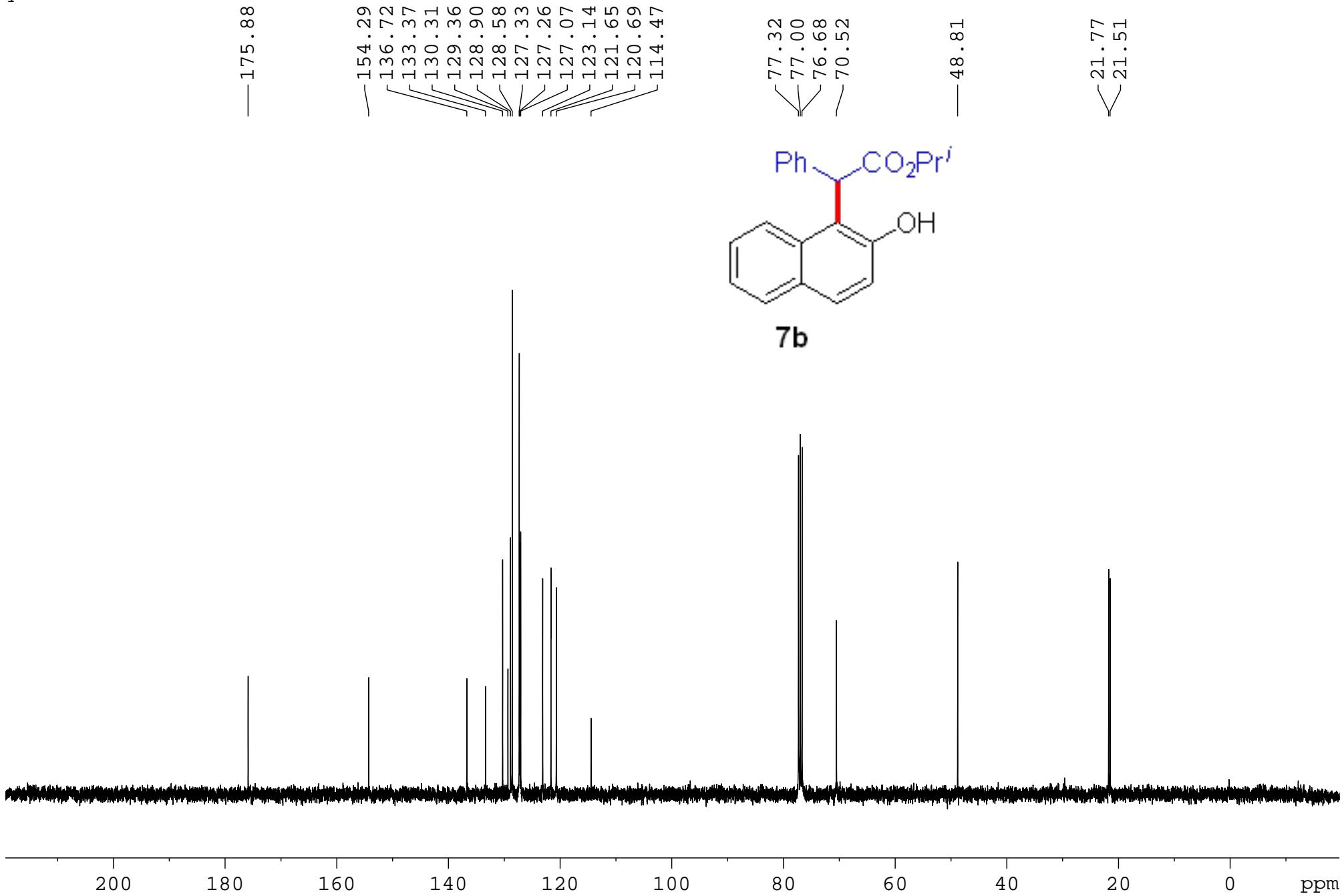


yzz-15-060-2-c

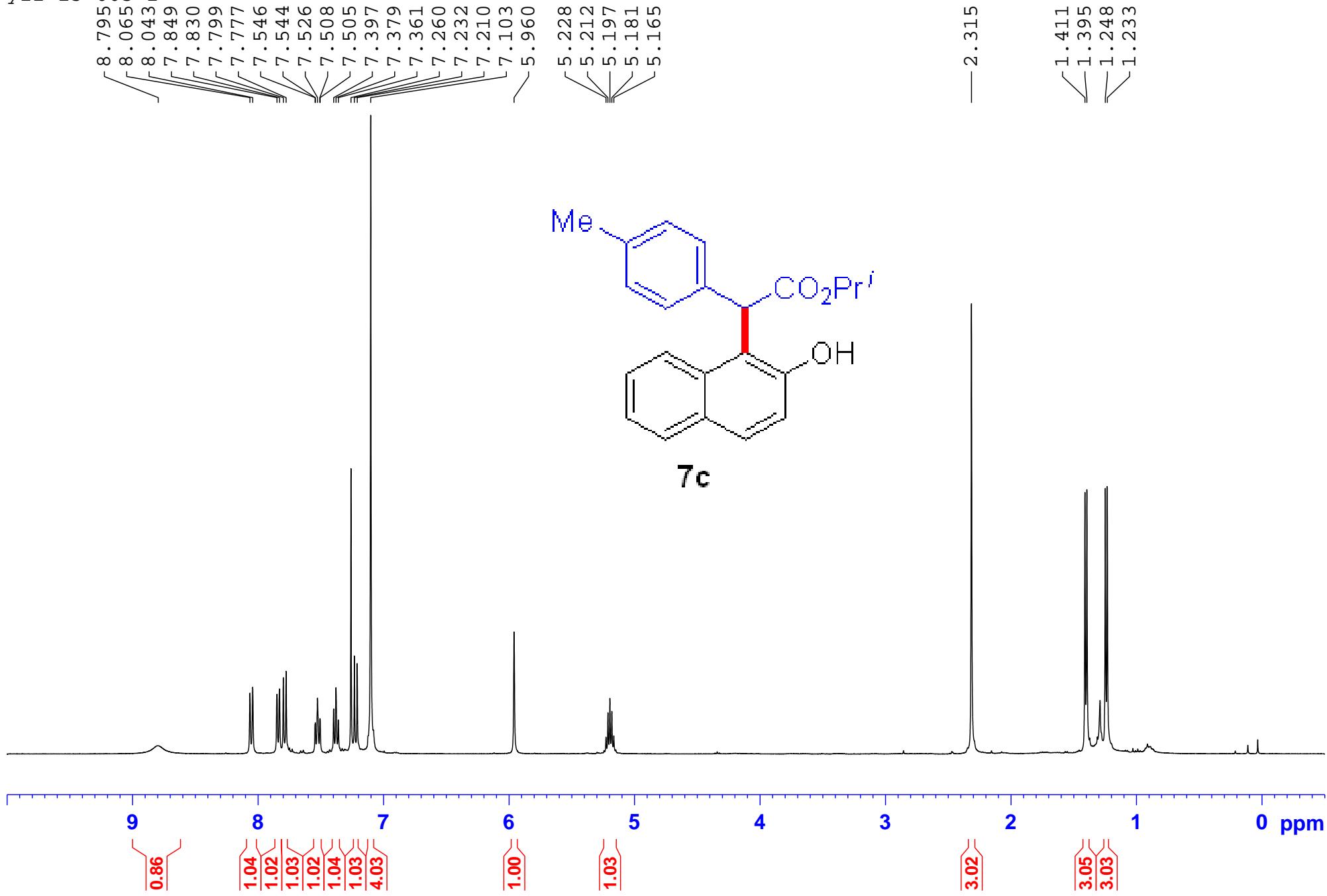




yzz-15-061-2-c



yzz-15-063-2



yzz-15-063-2-c

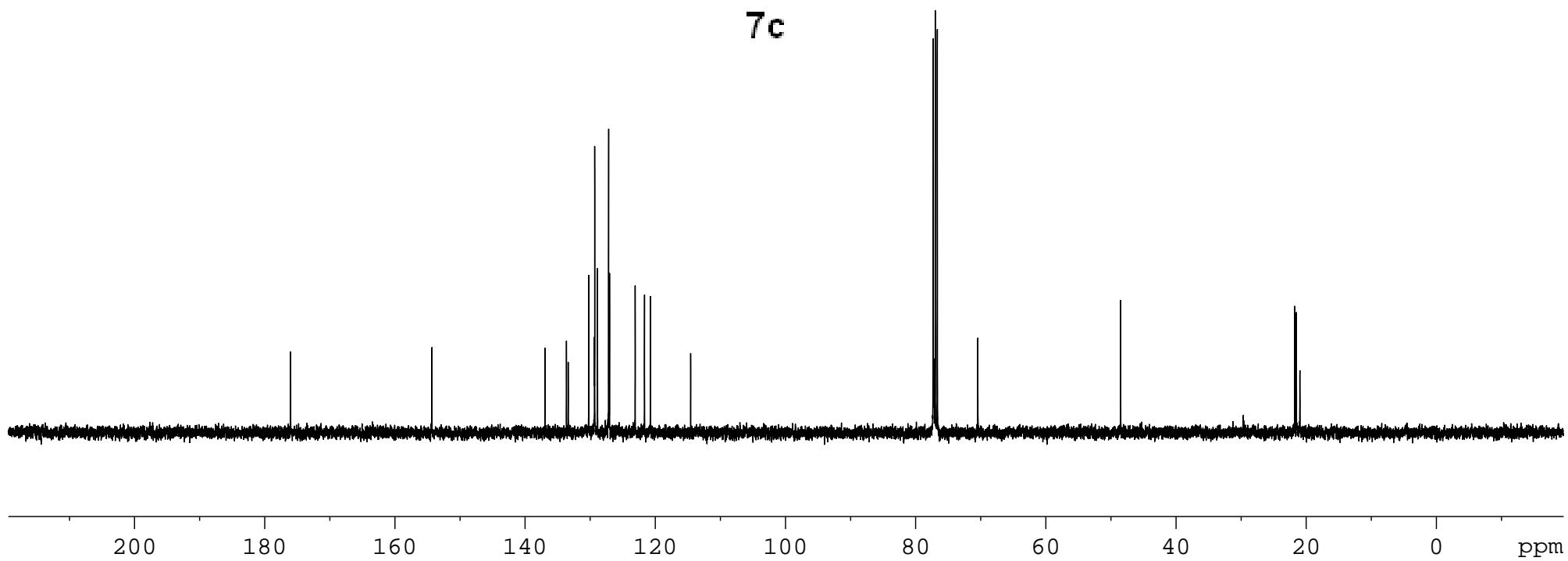
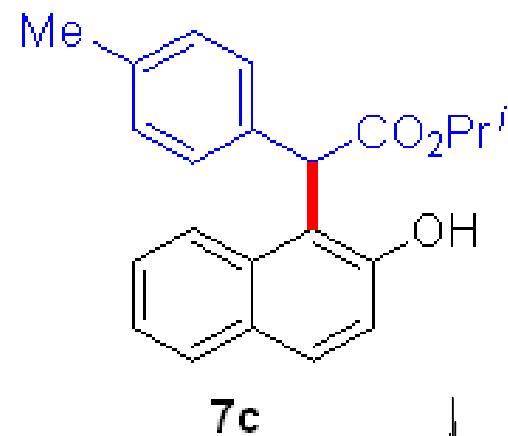
— 176.03

154.33
136.94
133.66
133.39
130.24
129.36
129.32
128.89
127.18
127.04
123.10
121.66
120.76
114.57

77.32
77.00
76.68
70.46

— 48.52

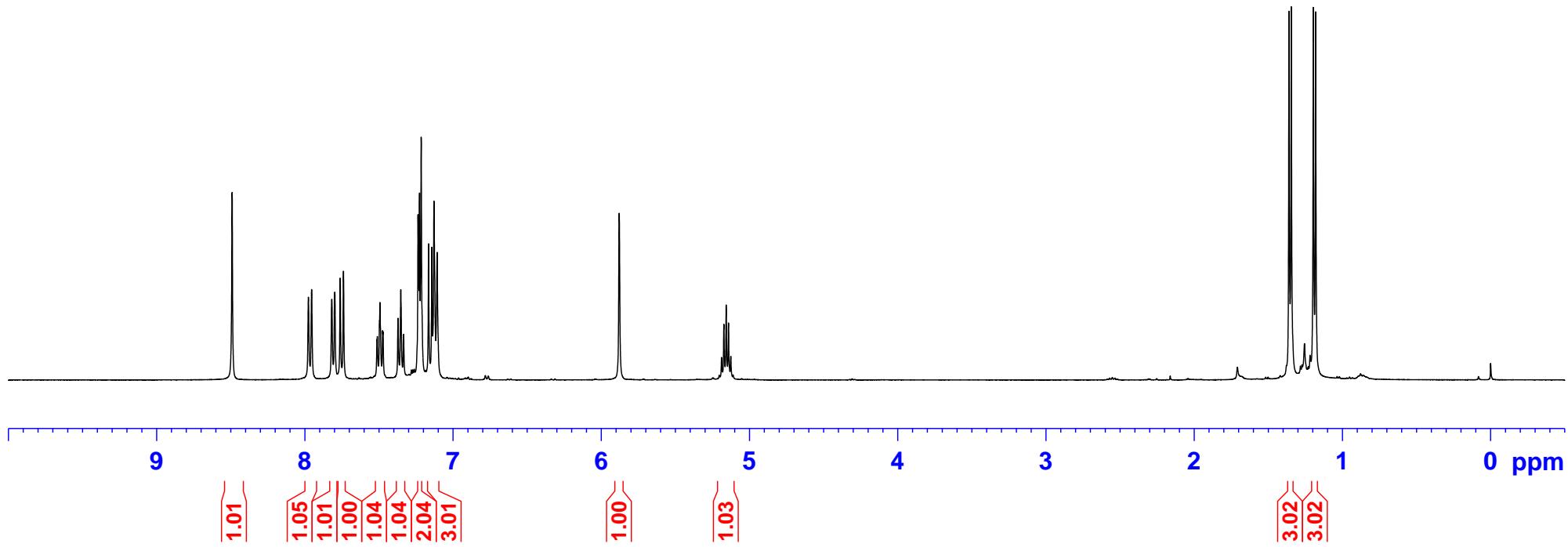
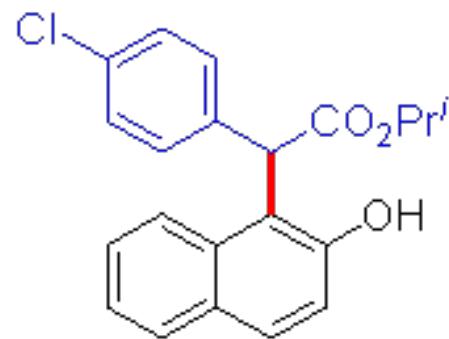
21.78
21.52
20.98



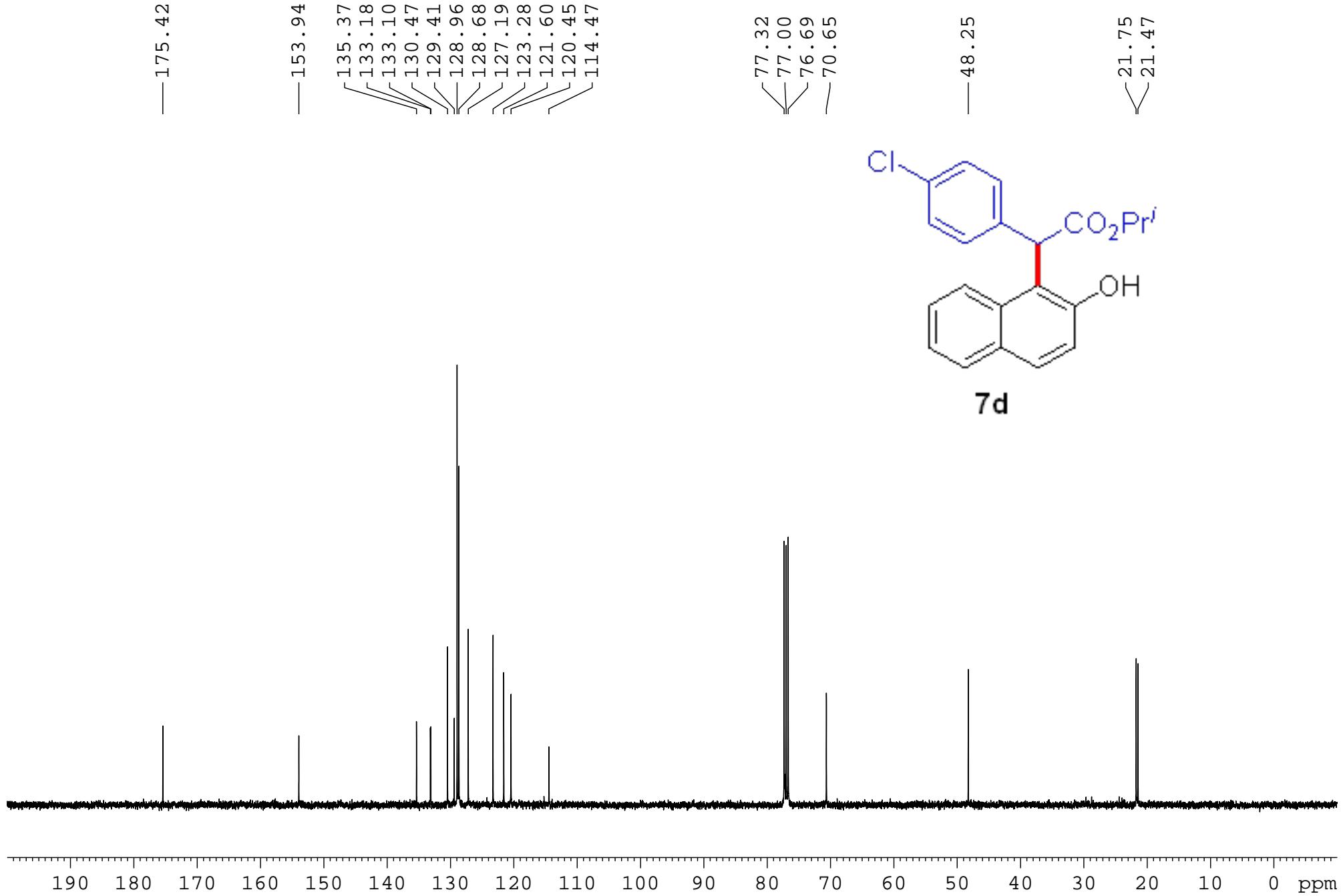
yzz-15-064-2

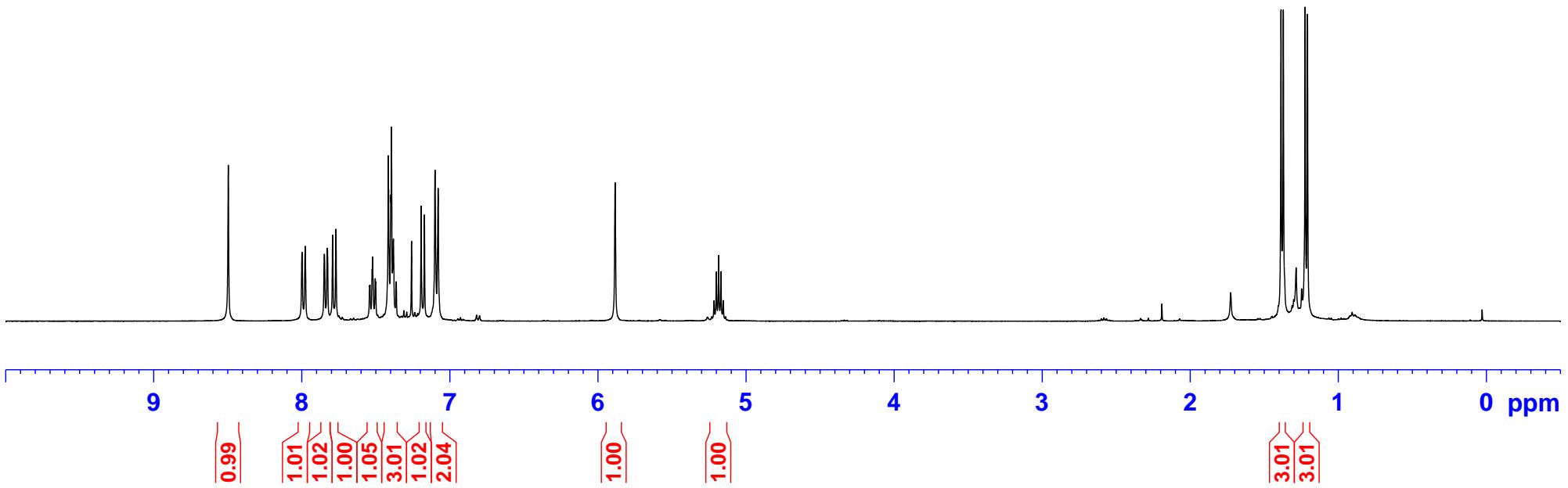
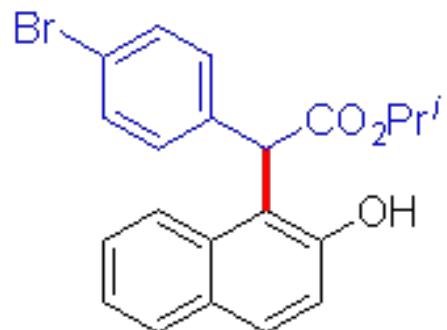
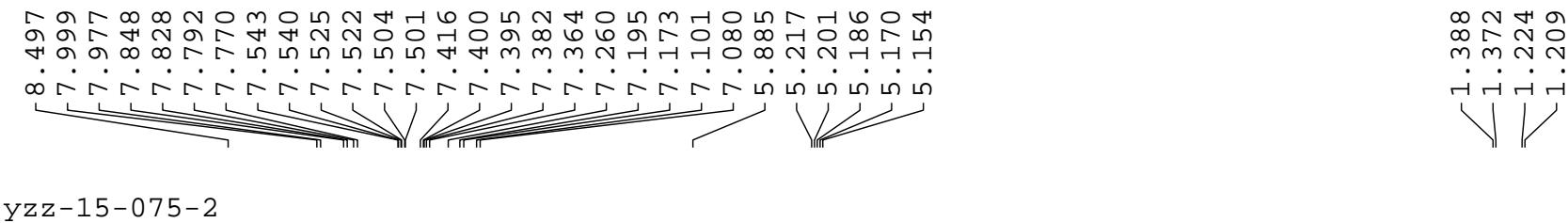
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7.818
7.798
7.761
7.739
7.514
7.512
7.494
7.476
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7.371
7.353
7.334
7.227
7.216
7.237
7.232
7.144
7.129
7.108
5.880
5.189
5.174
5.158
5.142
5.127

1.361
1.345
1.196
1.181



yzz-15-064-2-c





yzz-15-075-2-c

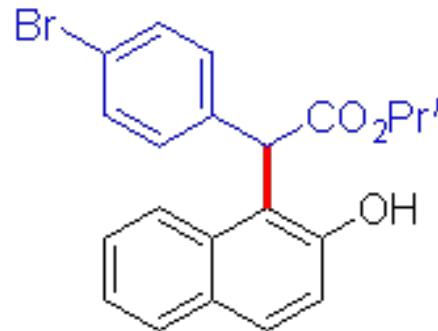
— 175.31

153.92
135.92
133.17
131.62
130.48
129.41
129.32
128.96
127.19
123.29
121.59
121.23
120.43
114.42

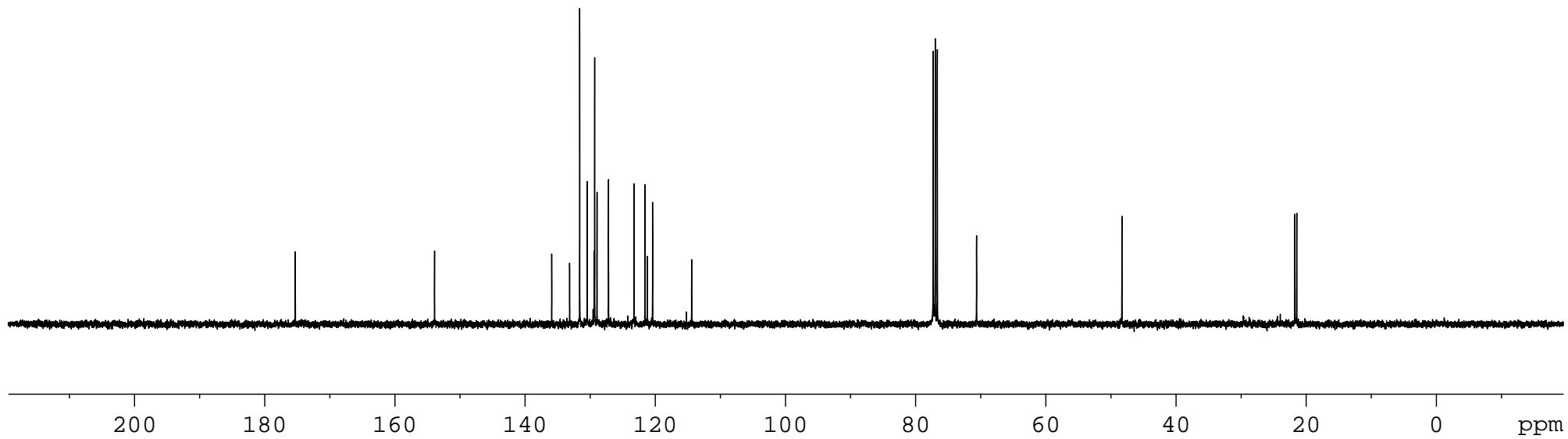
77.32
77.00
76.68
70.65

— 48.32

21.75
21.46



7e



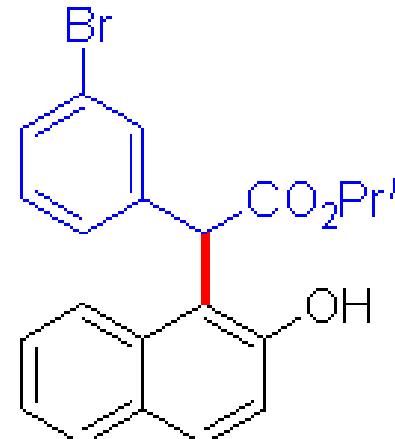
yzz-15-065-1

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7.974
7.852
7.832

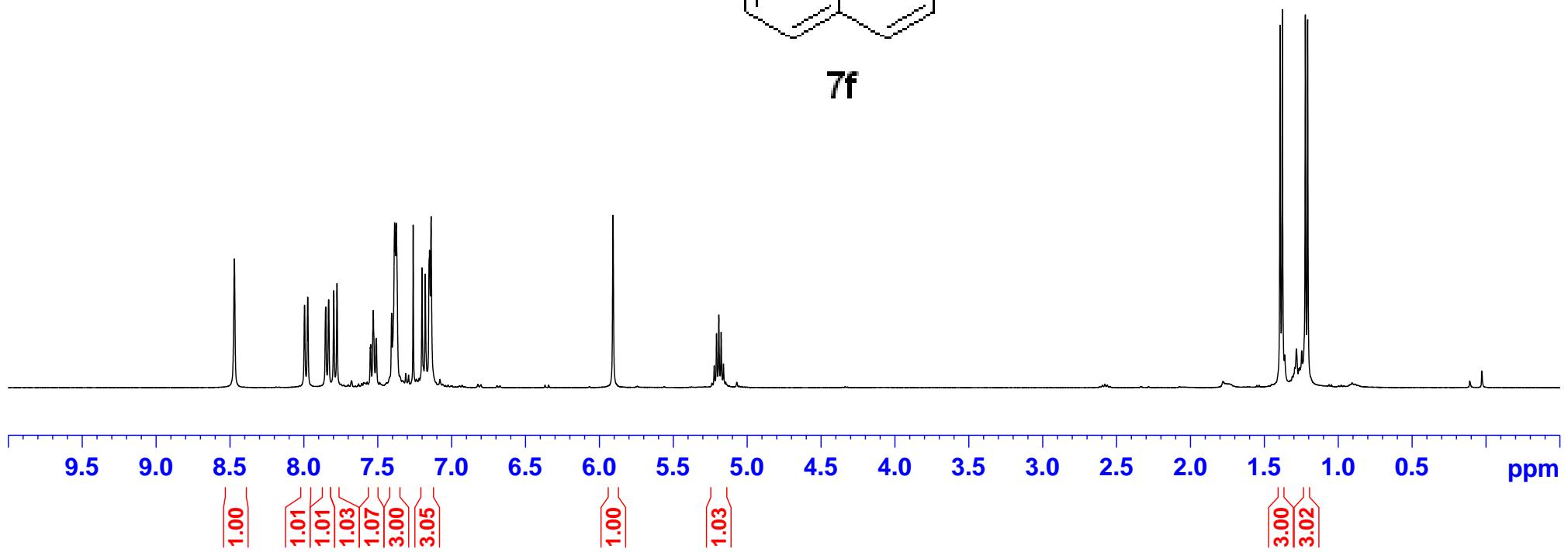
7.798
7.776
7.755
7.732
7.550
7.548
7.530

7.512
7.509
7.406
7.386
7.381
7.375
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7.139
5.908
5.224
5.208
5.192
5.177
5.161

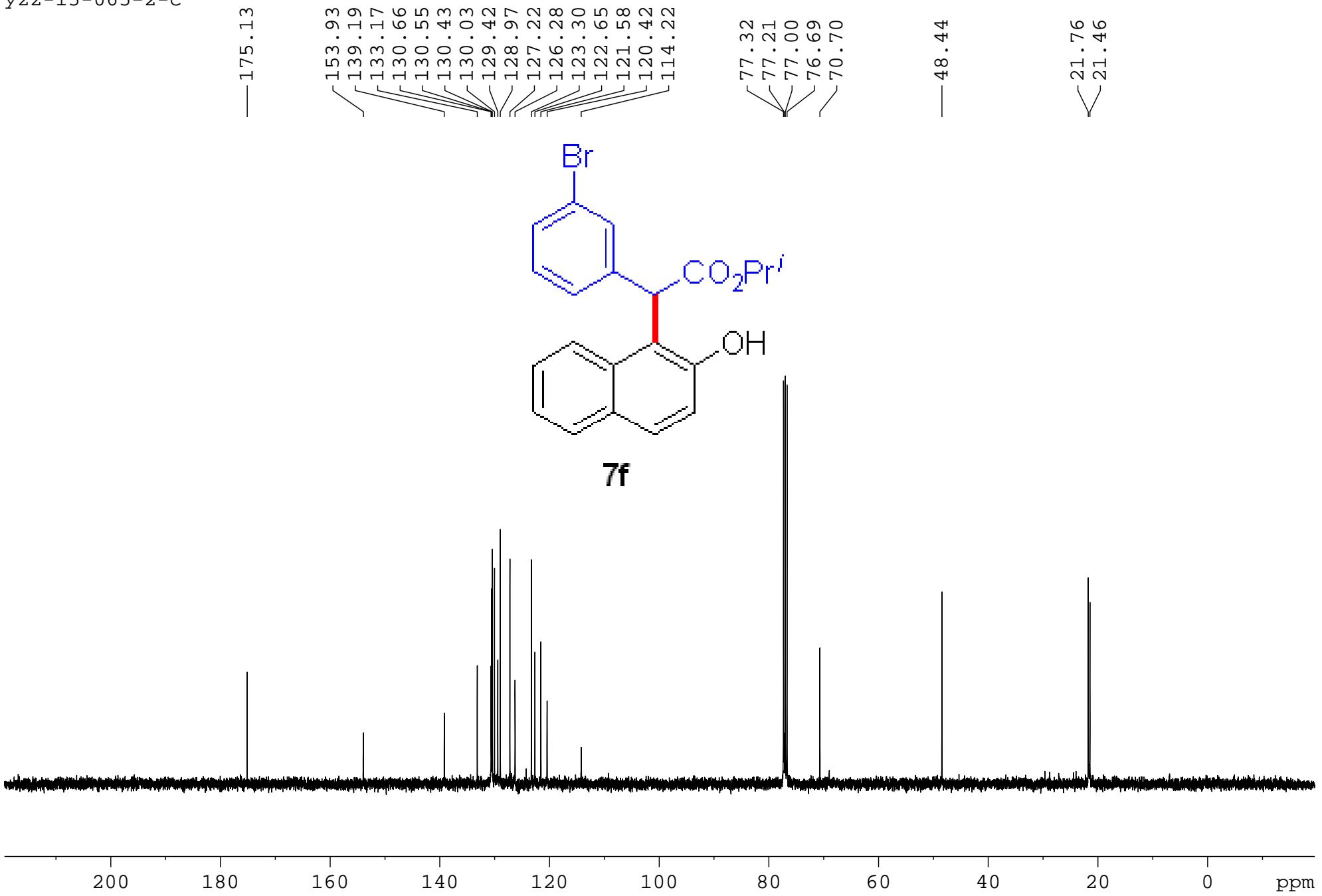
1.394
1.379
1.223
1.207

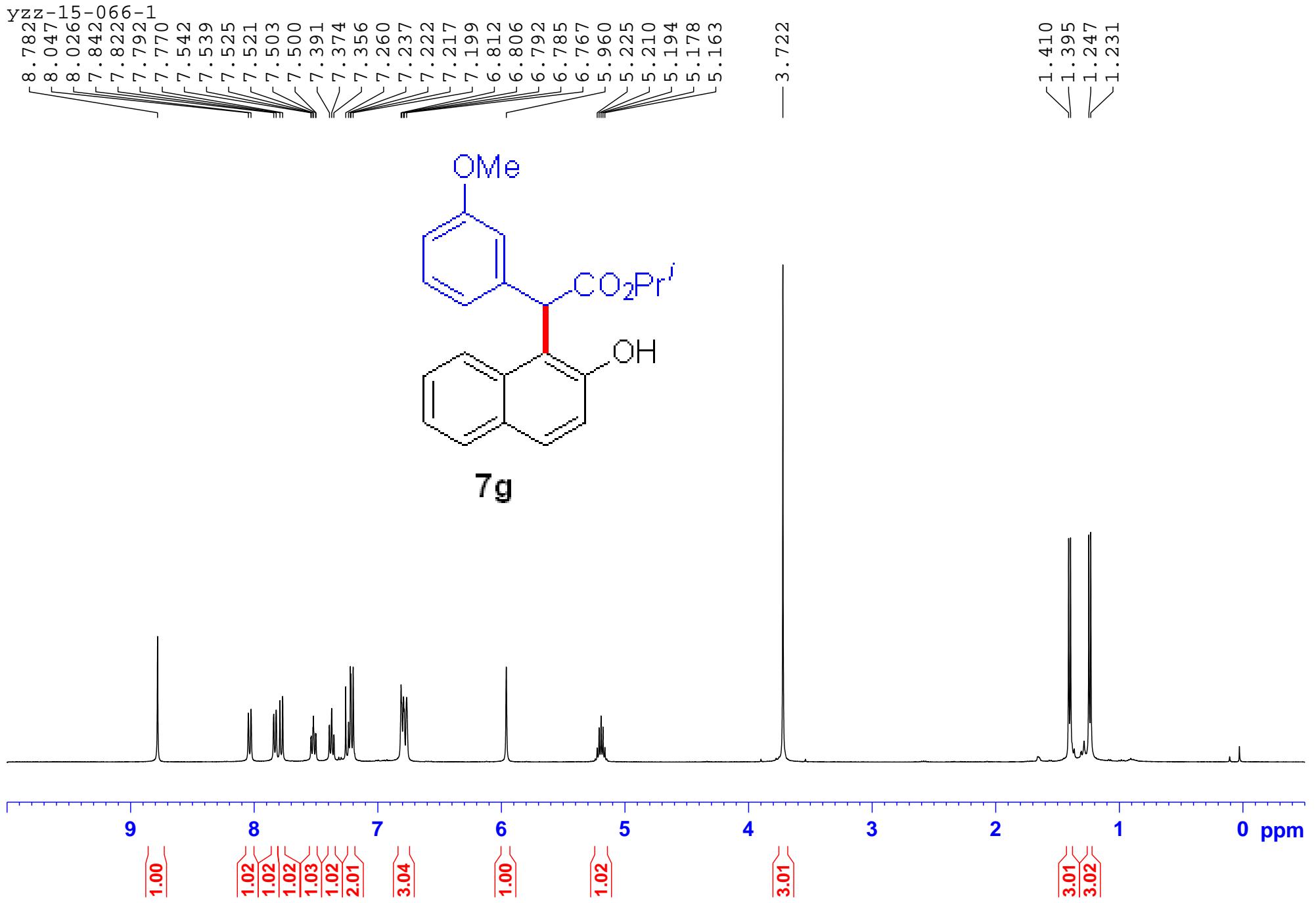


7f

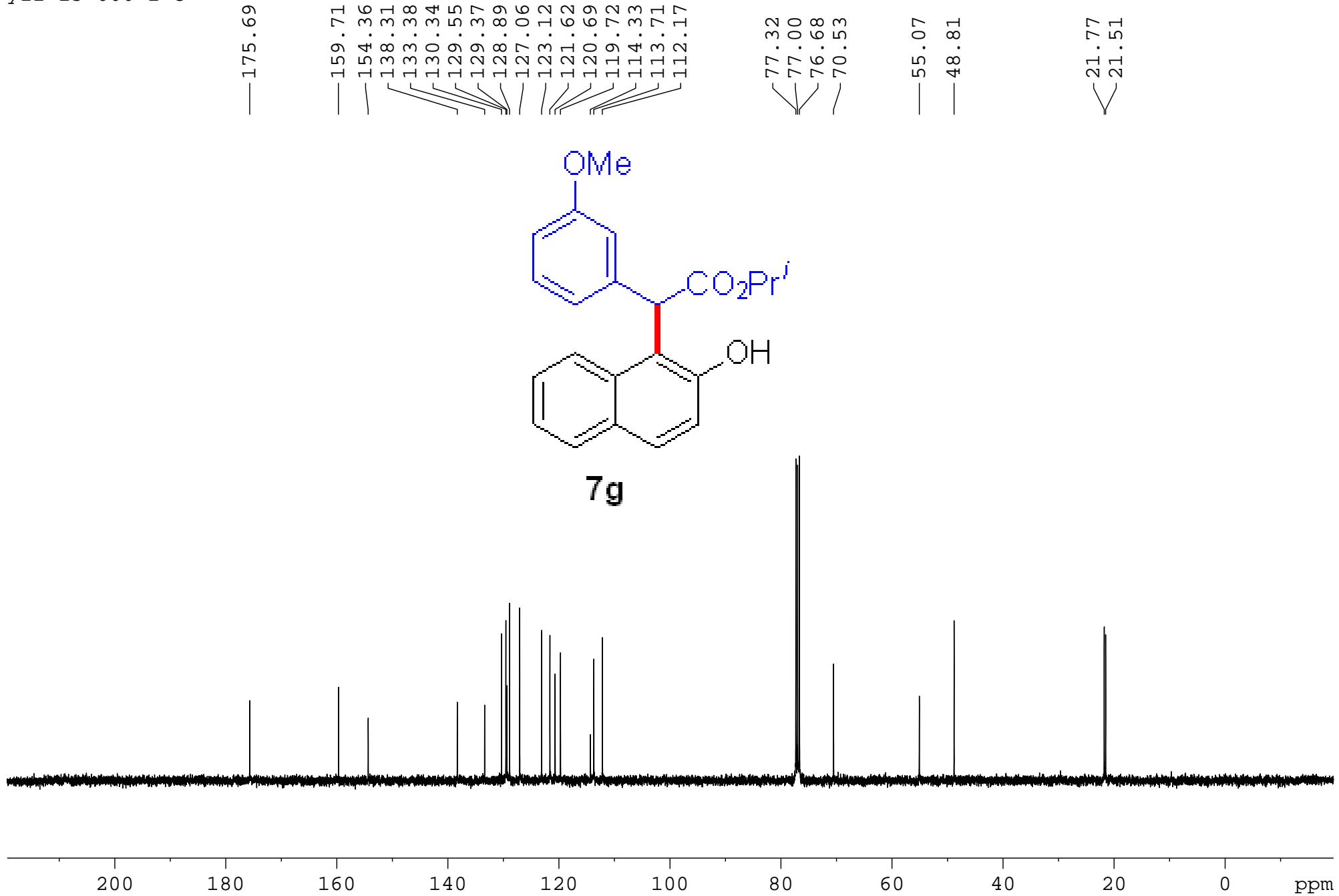


yzz-15-065-2-c





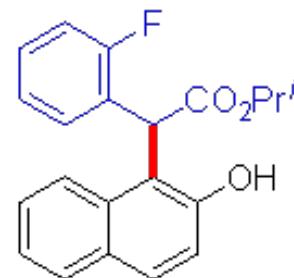
yzz-15-066-1-c



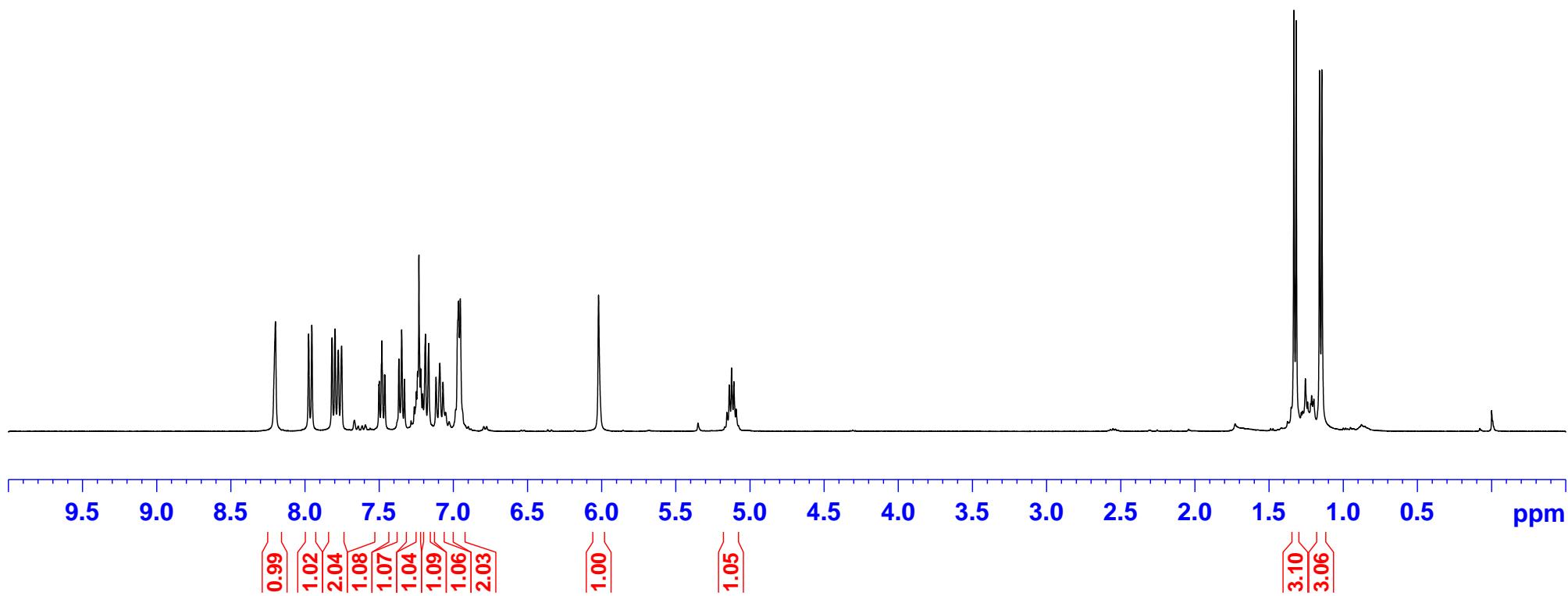
yzz-15-067-1

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7.817
7.797
7.753
7.502
7.499
7.482
7.463
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7.366
7.348
7.329
7.249
7.241
7.232
7.220
7.207
7.188
7.166
7.116
7.091
7.070
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6.021
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5.125
5.109
5.094

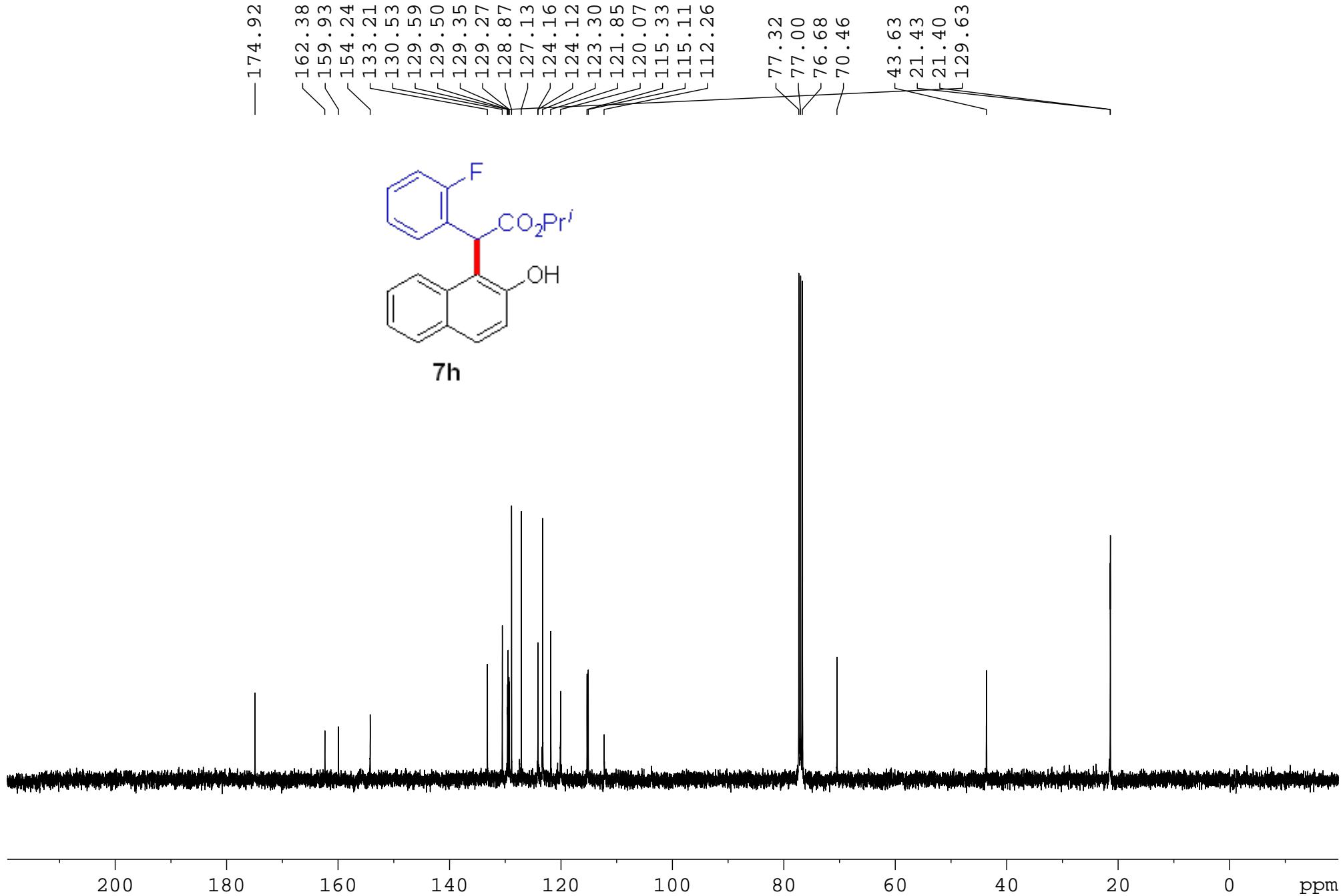
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1.317
1.161
1.145



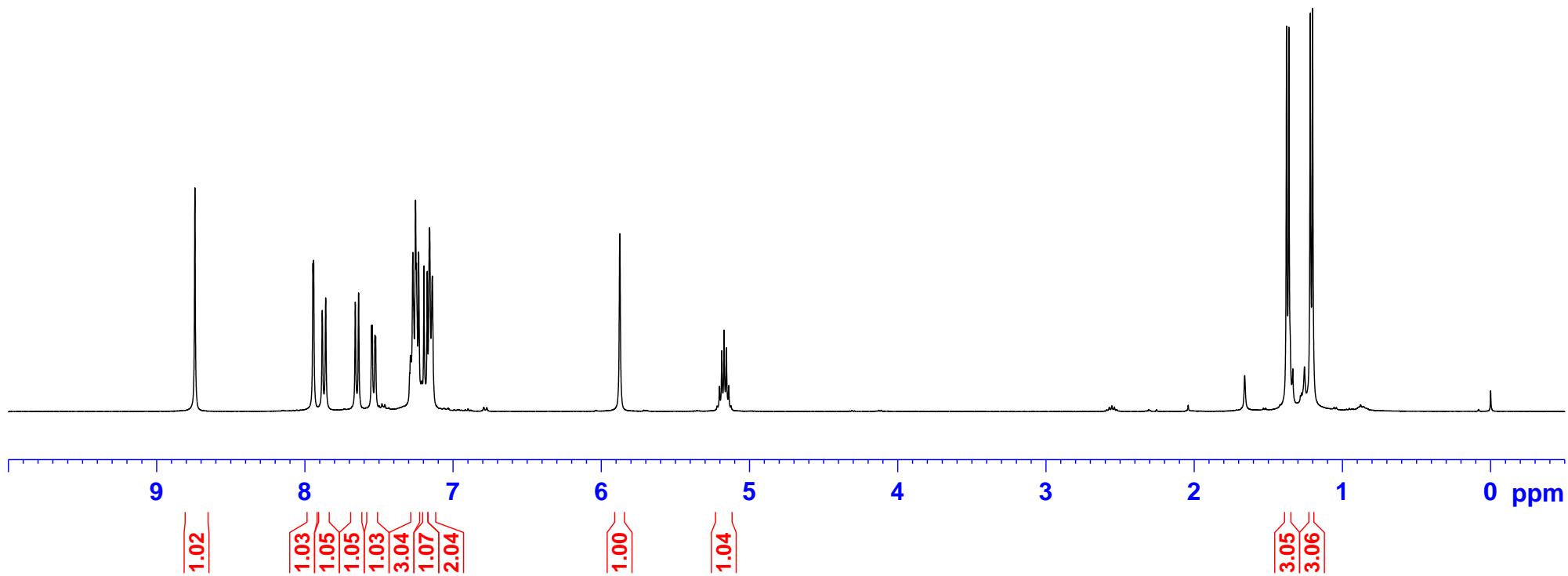
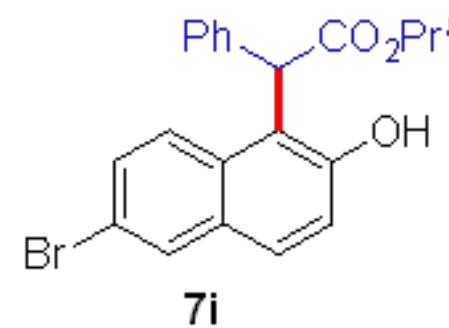
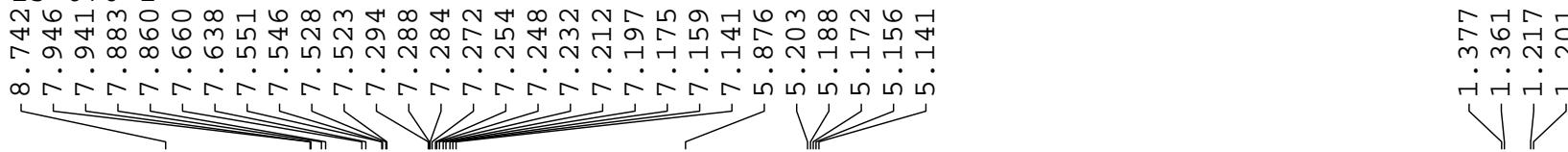
7h



yzz-15-067-1-c



yzz-15-070-1



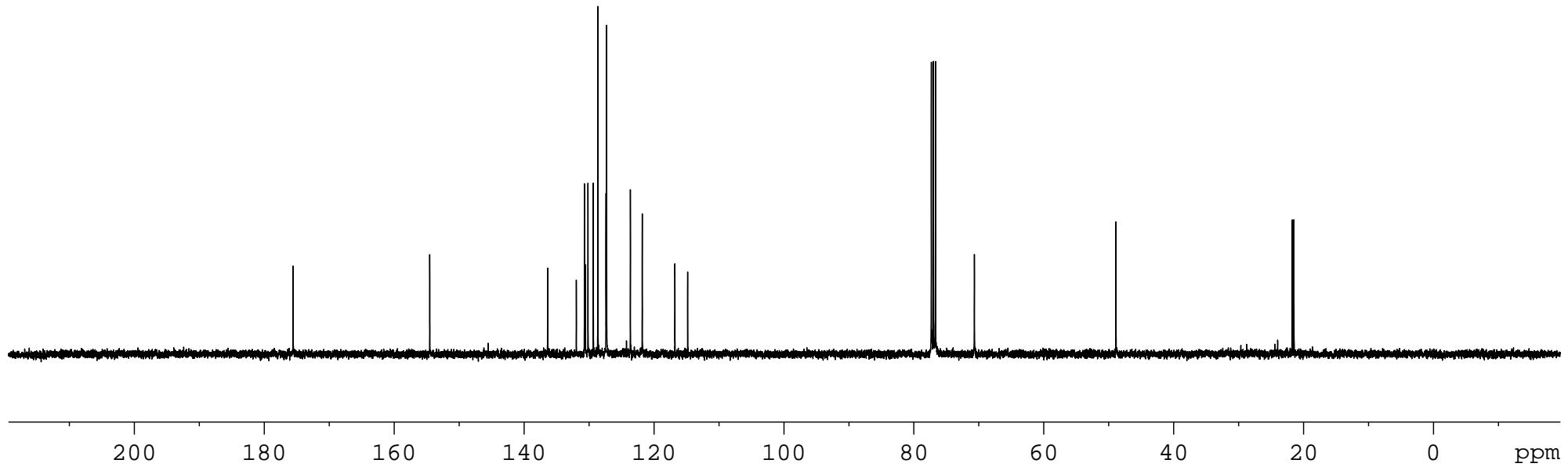
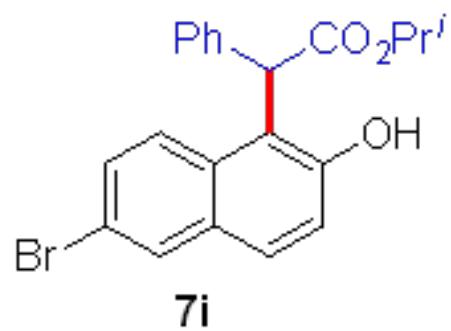
yzz-15-070-1-c

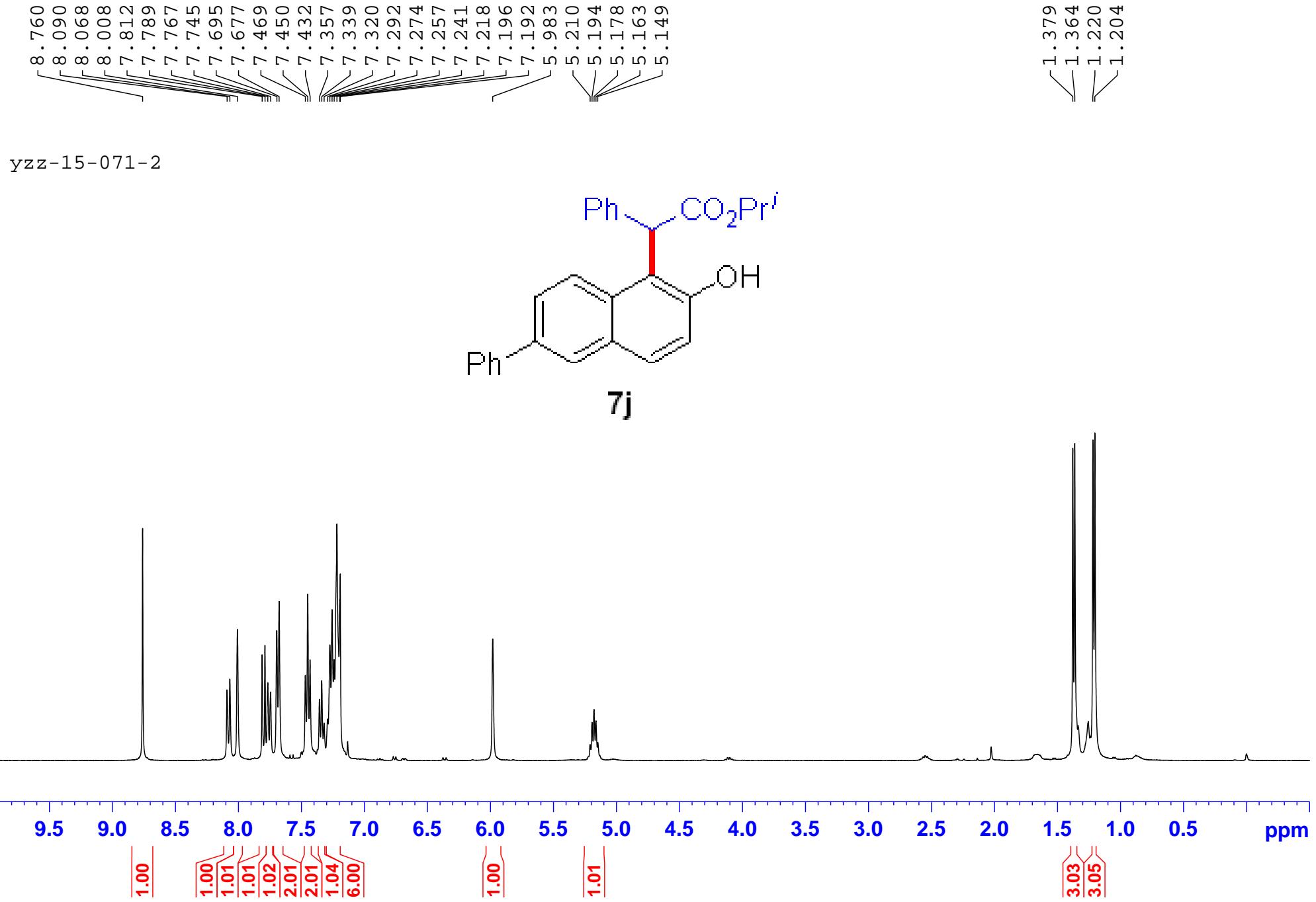
— 175.56

154.53
136.39
131.97
130.72
130.58
130.17
129.36
128.66
127.40
127.31
123.65
121.82
116.79
114.83

— 48.90

21.75
21.50





yzz-15-071-2_c

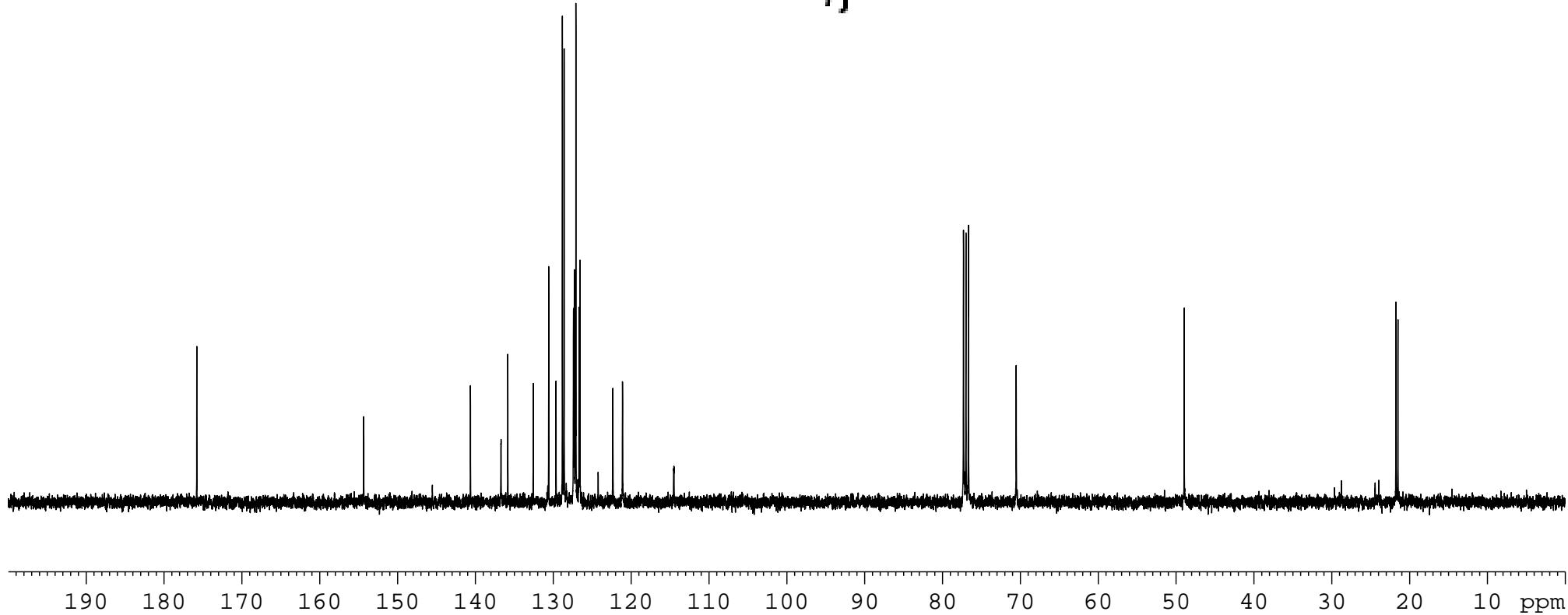
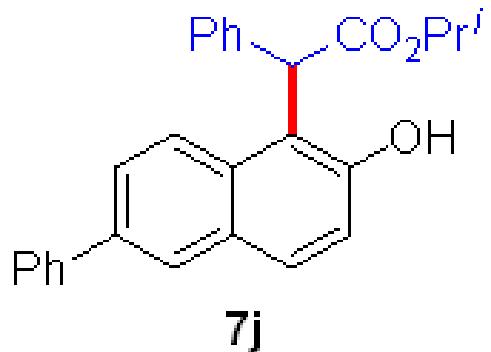
— 175.80 ppm

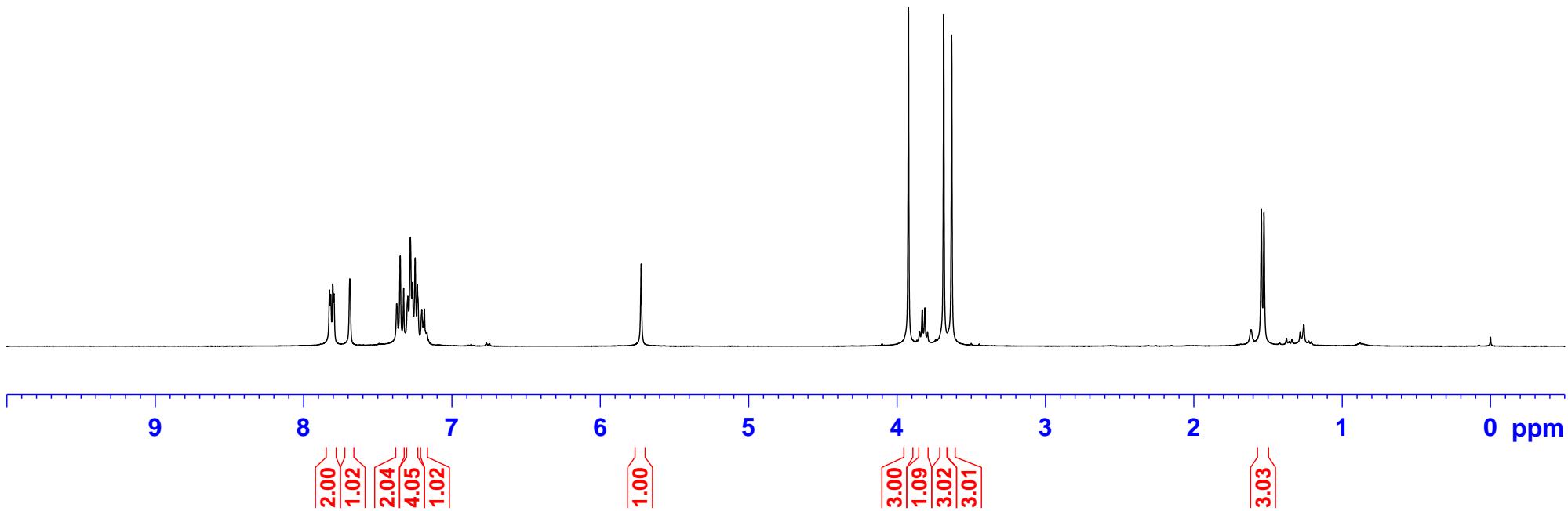
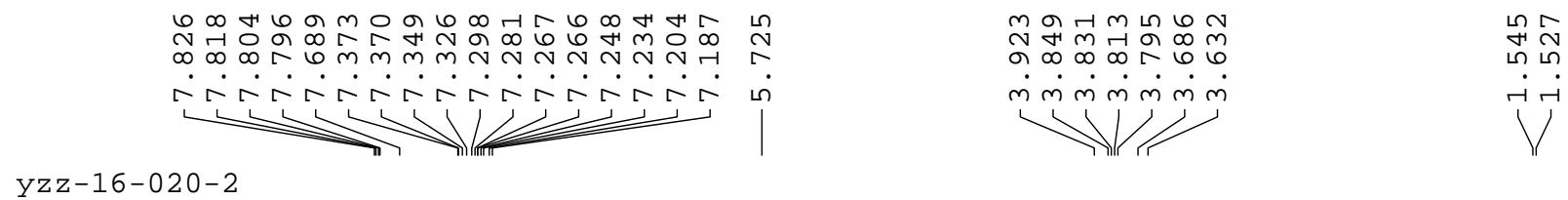
154.36
140.63
136.70
135.84
132.58
130.59
129.64
128.82
128.61
127.40
127.29
127.15
127.09
126.69
126.58
122.35
121.11
114.52

77.32
77.00
76.68
70.55

48.96

21.77
21.51





yzz-16-020-1-c

174.90
173.89

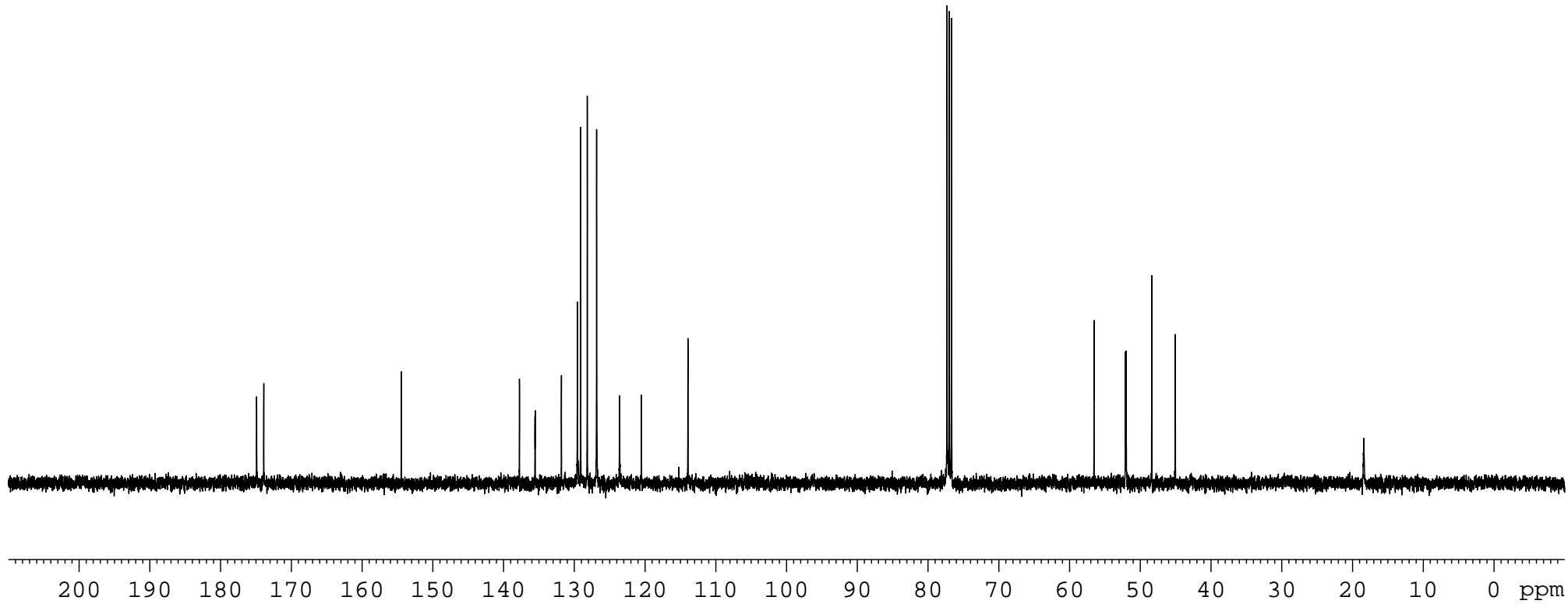
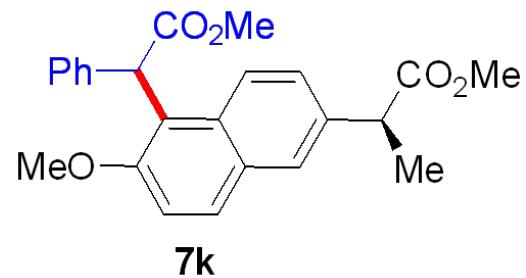
154.45

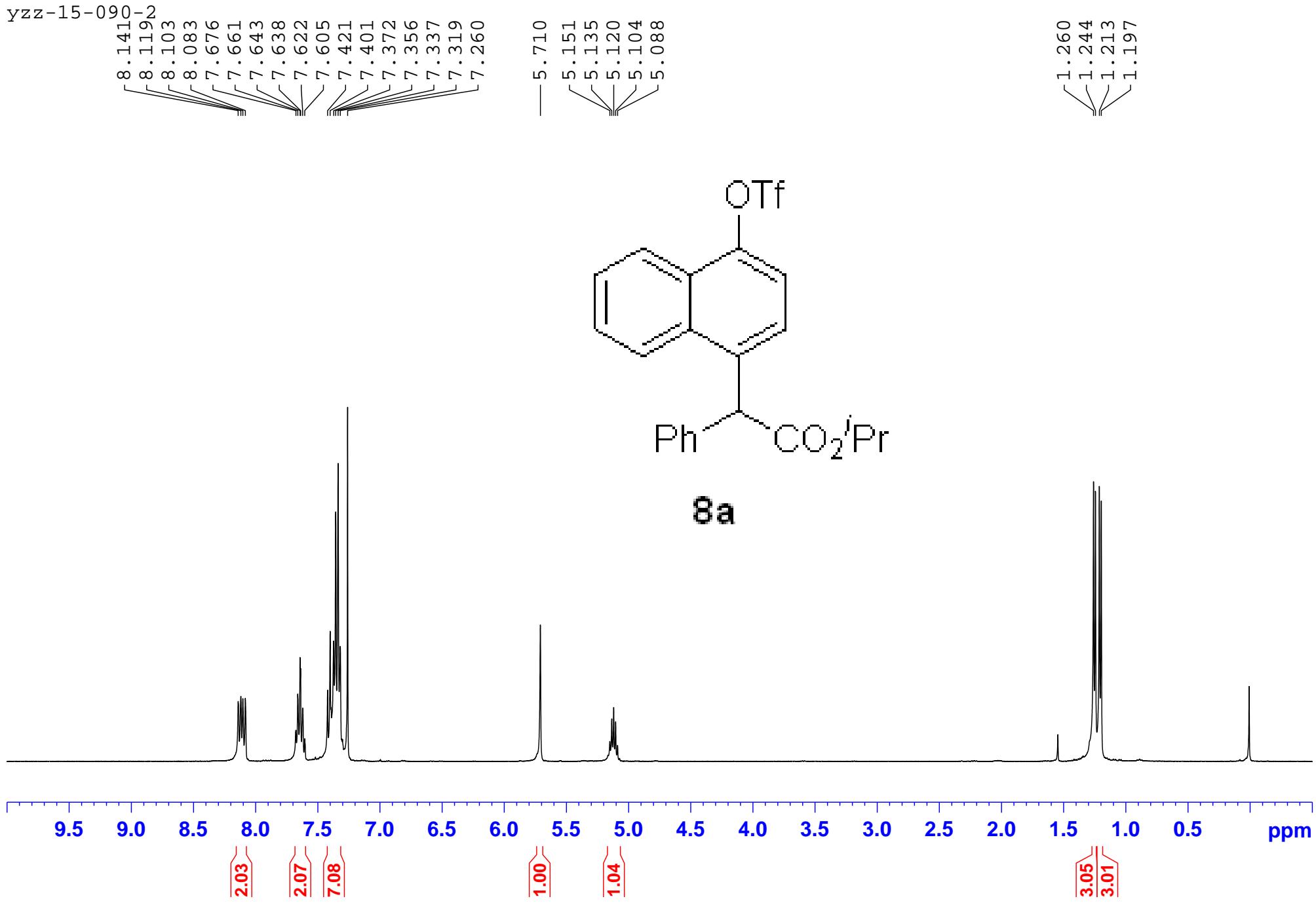
137.77
135.57
135.54
131.84
129.57
129.12
128.18
126.86
123.61
120.53
113.95

77.32
77.00
76.68

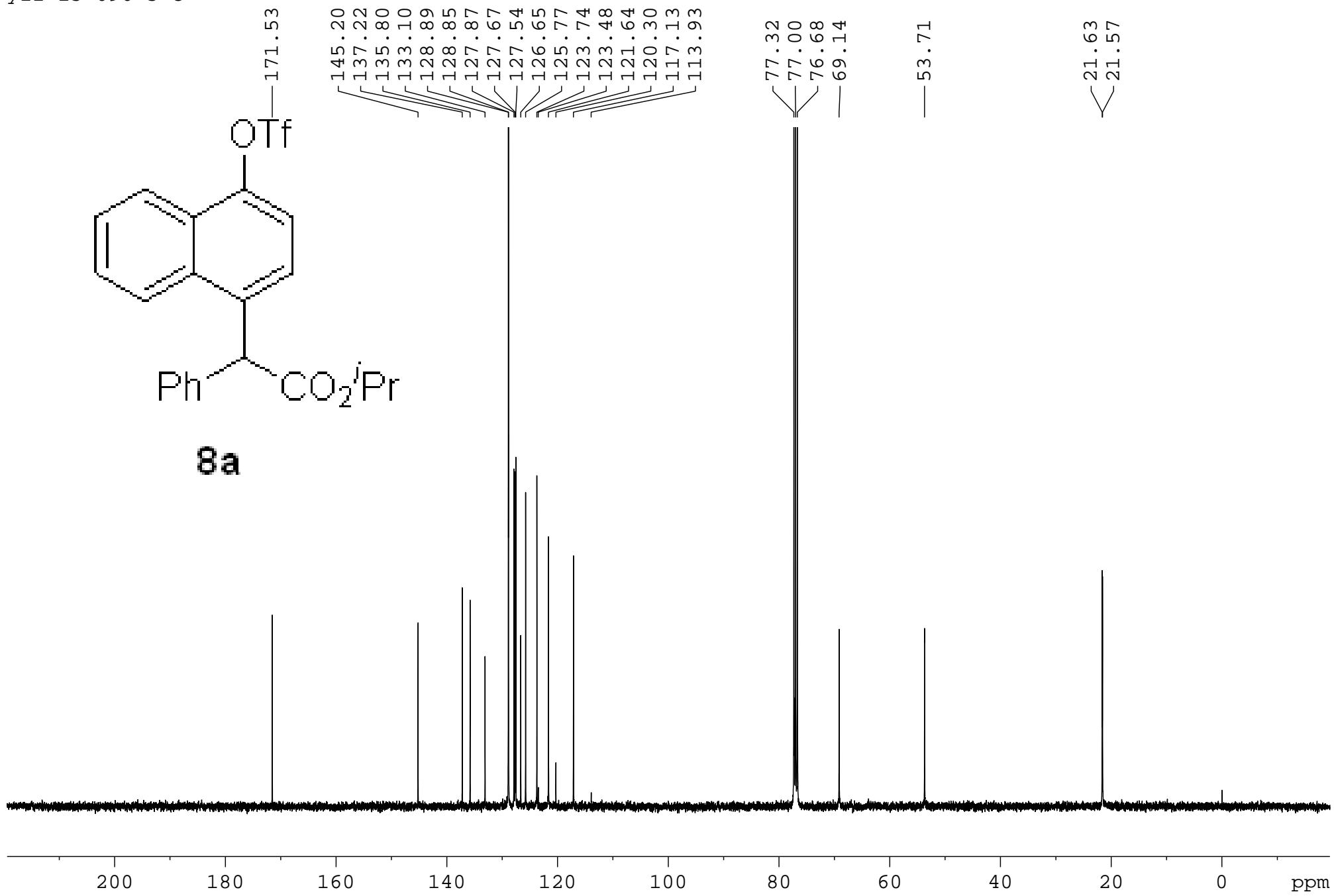
56.54
52.10
51.99
48.39
45.06

18.41





yzz-15-090-3-c

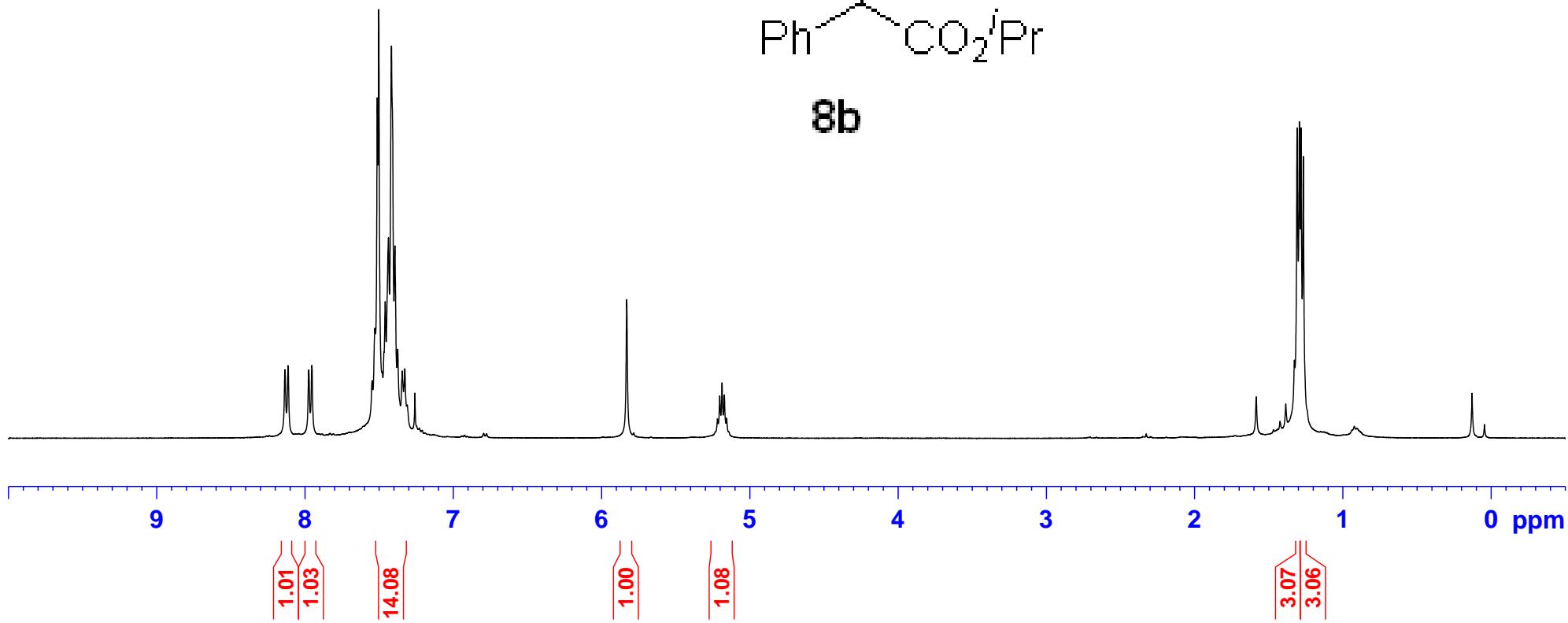
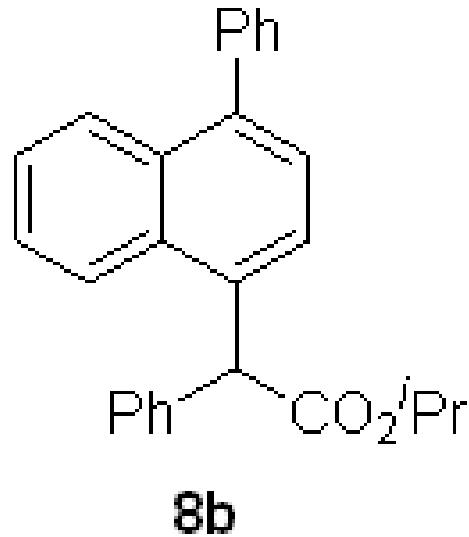


yzz-15-085-1

8.135
8.114
7.974
7.954
7.529
7.513
7.502
7.468
7.437
7.459
7.393
7.375

5.820
5.219
5.203
5.188
5.172
5.157

1.310
1.294
1.283
1.268



yzz-15-085-1

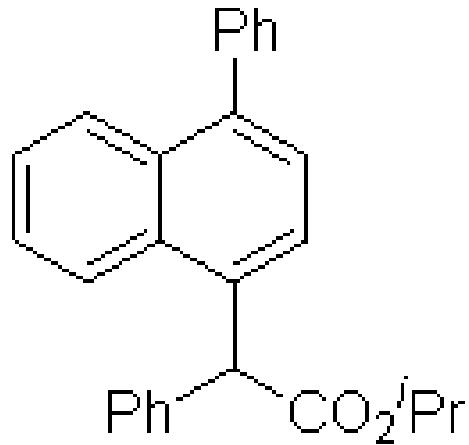
— 172.27

140.66
140.09
138.07
134.23
132.15
131.91
130.10
129.01
128.63
128.19
127.31
127.22
127.06
126.42
126.20
125.76
125.65
123.46

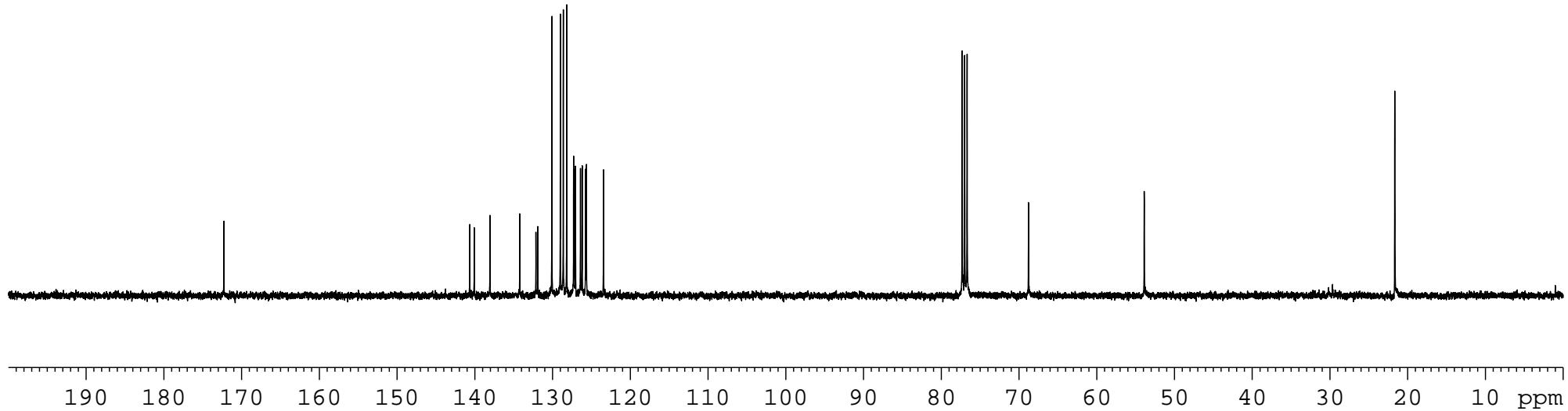
77.32
77.00
76.68
— 68.79

— 53.90

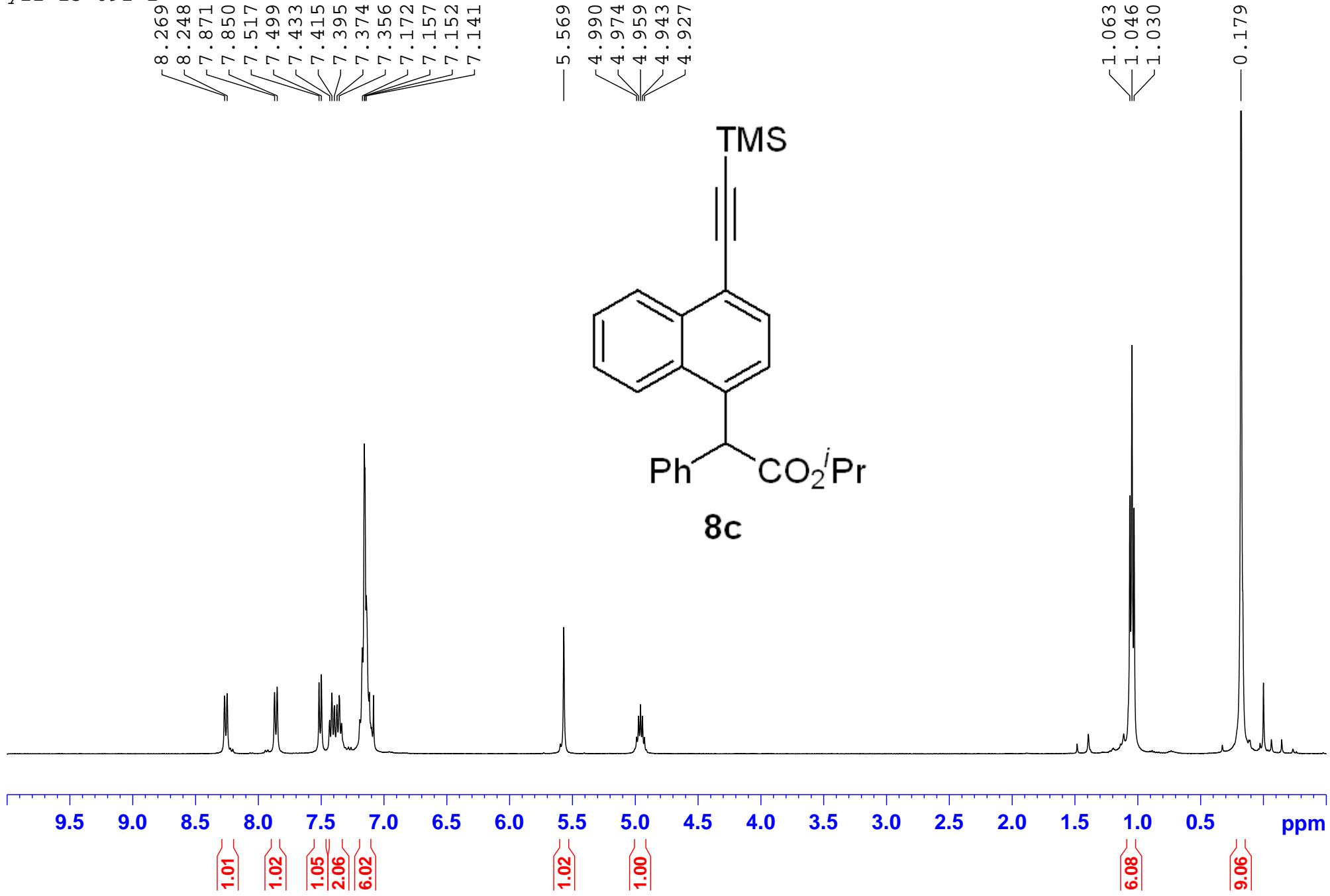
— 21.67



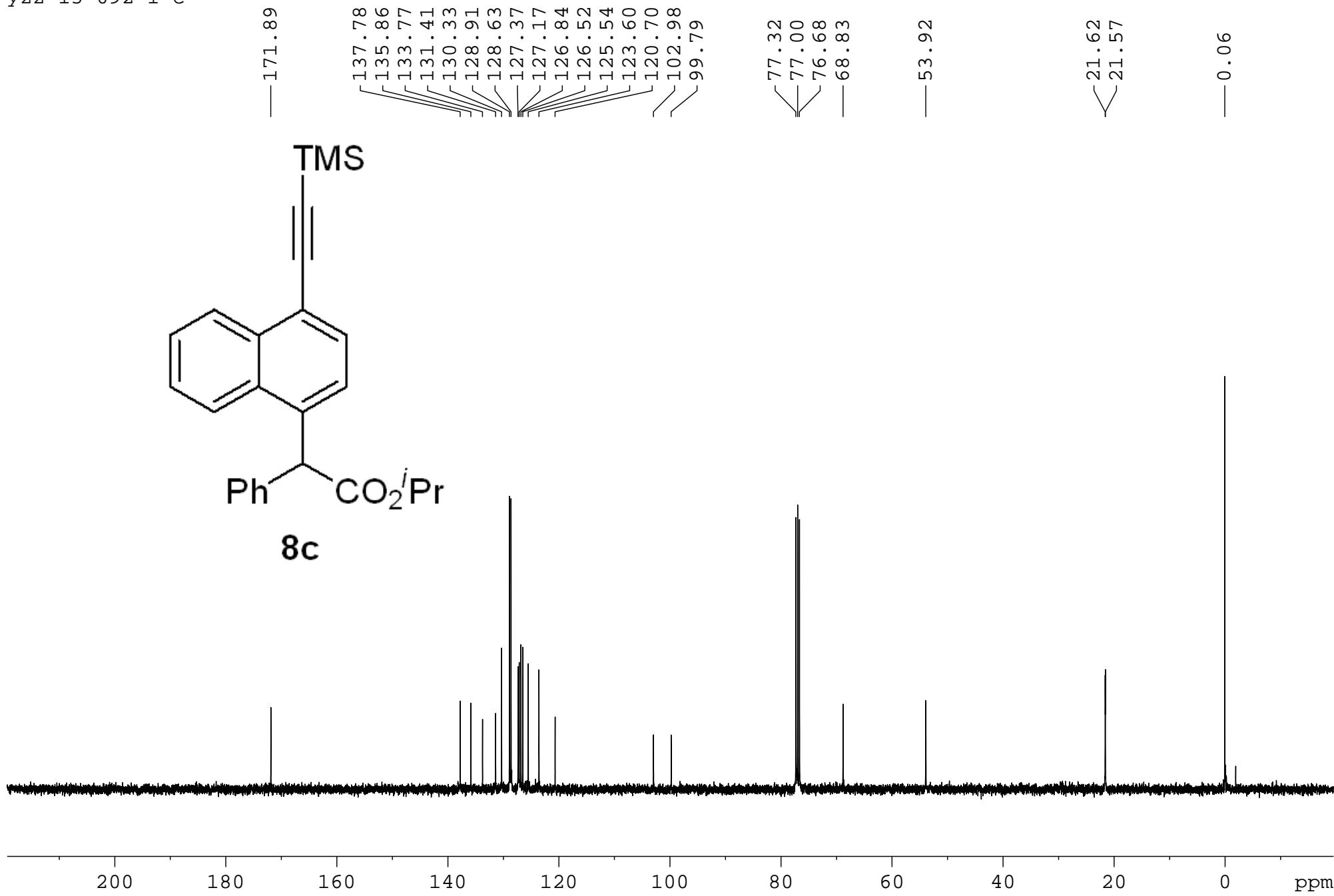
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yzz-15-092-1



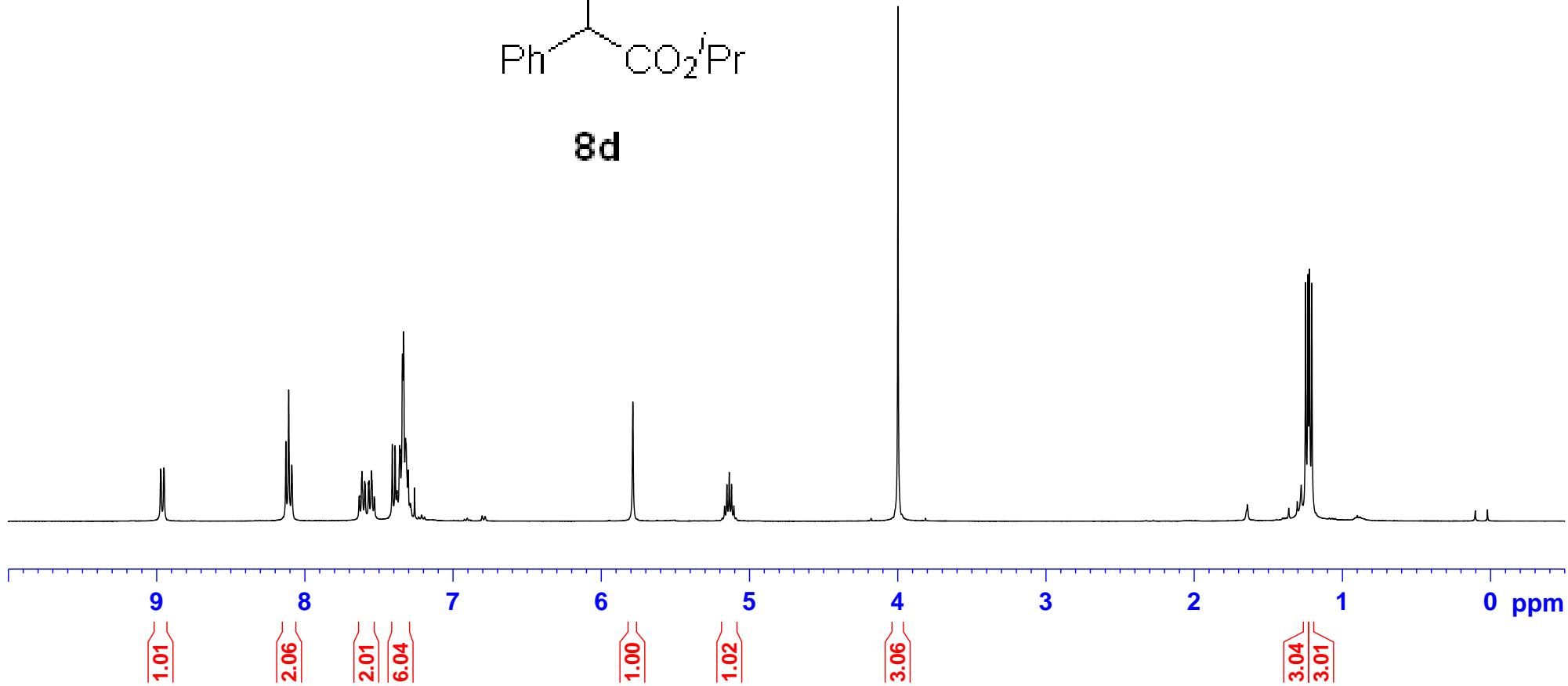
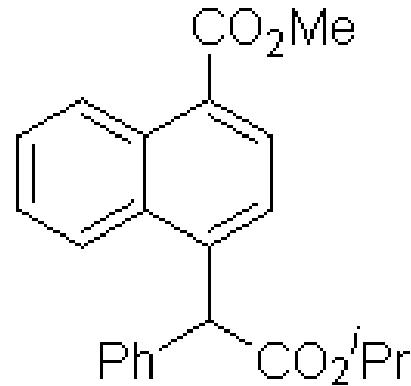
yzz-15-092-1-c



yzz-15-094-1

8.088
7.634
7.617
7.614
7.596
7.593
7.570
7.567
7.550
7.547
7.532
7.529
7.410
7.391
7.380
7.378
7.360
7.356
7.342
7.333
7.321
7.317
7.302
7.293
7.285
7.259
5.787
5.169
5.153
5.137
5.122
5.106
3.999

1.248
1.233
1.222
1.206



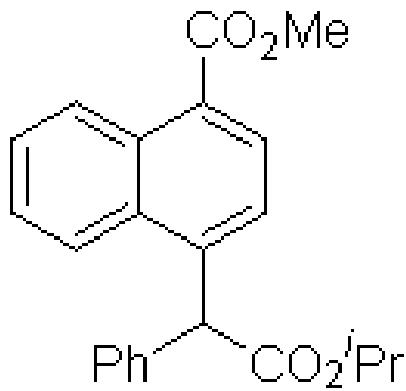
yzz-15-094-1-c

— 171.69
— 167.91

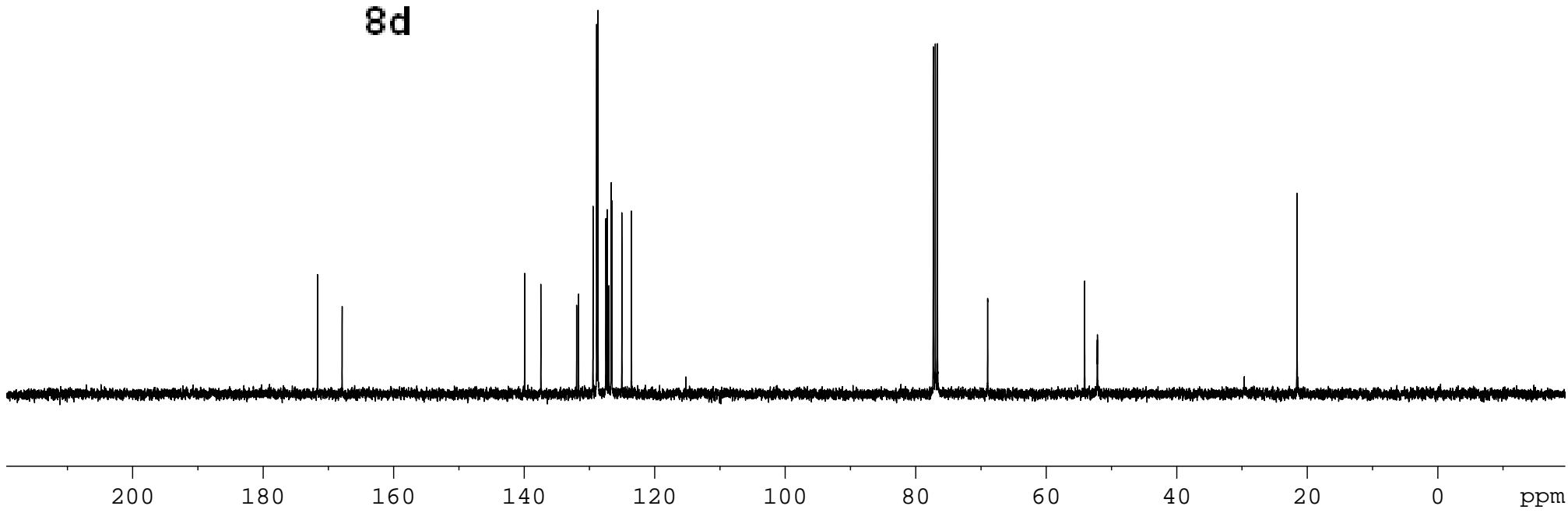
139.95
137.46
131.97
131.68
129.46
128.91
128.70
127.49
127.30
127.01
126.68
126.57
125.04
123.58

77.32
77.00
76.68
— 68.99
— 54.13
— 52.19
— 52.15

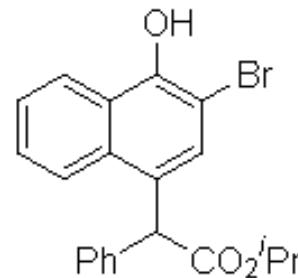
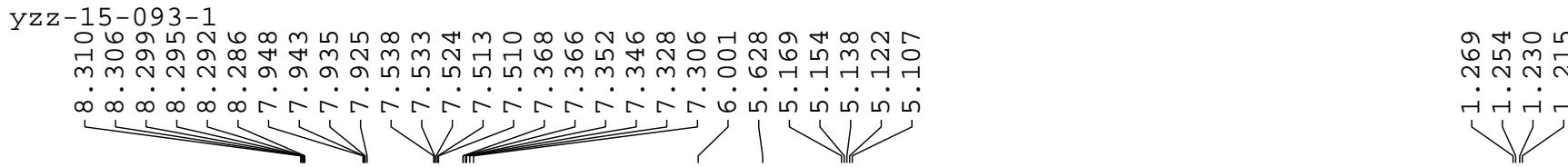
— 21.57



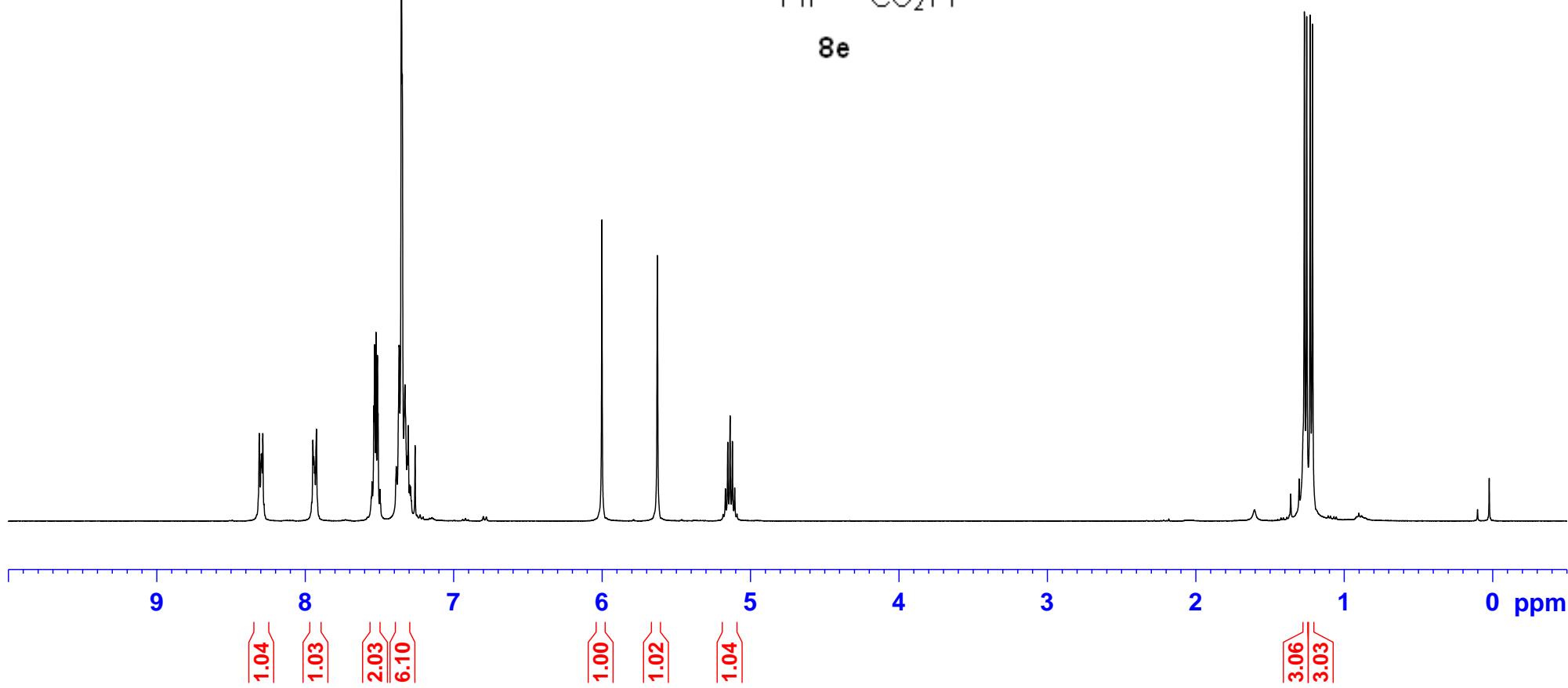
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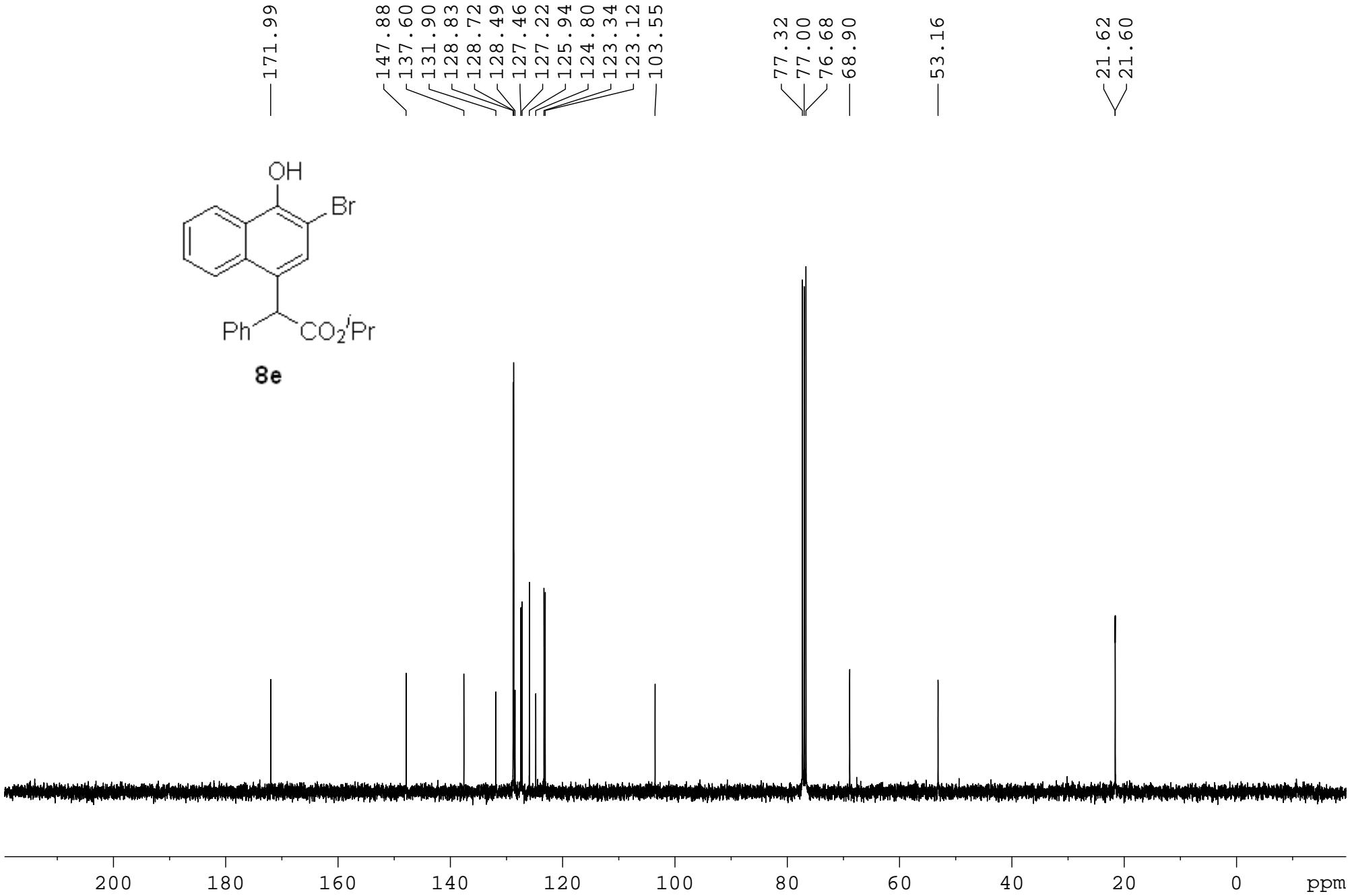
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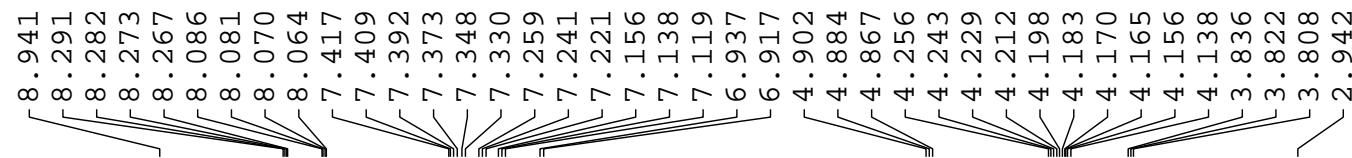


8e

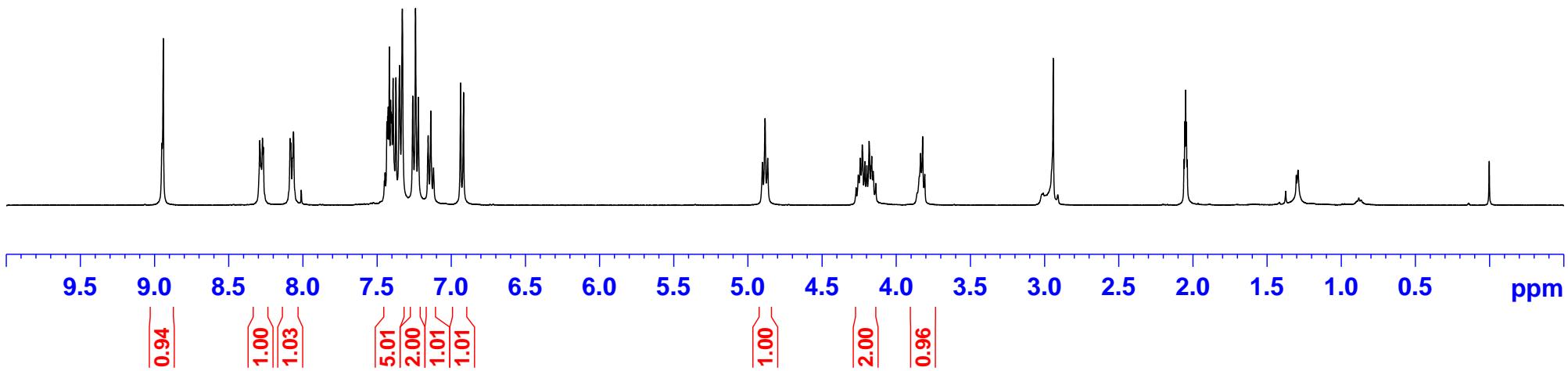
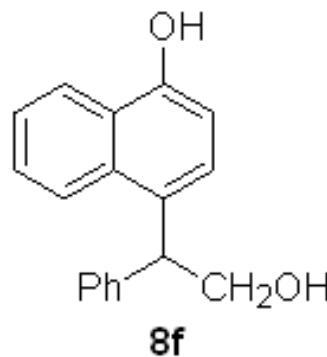


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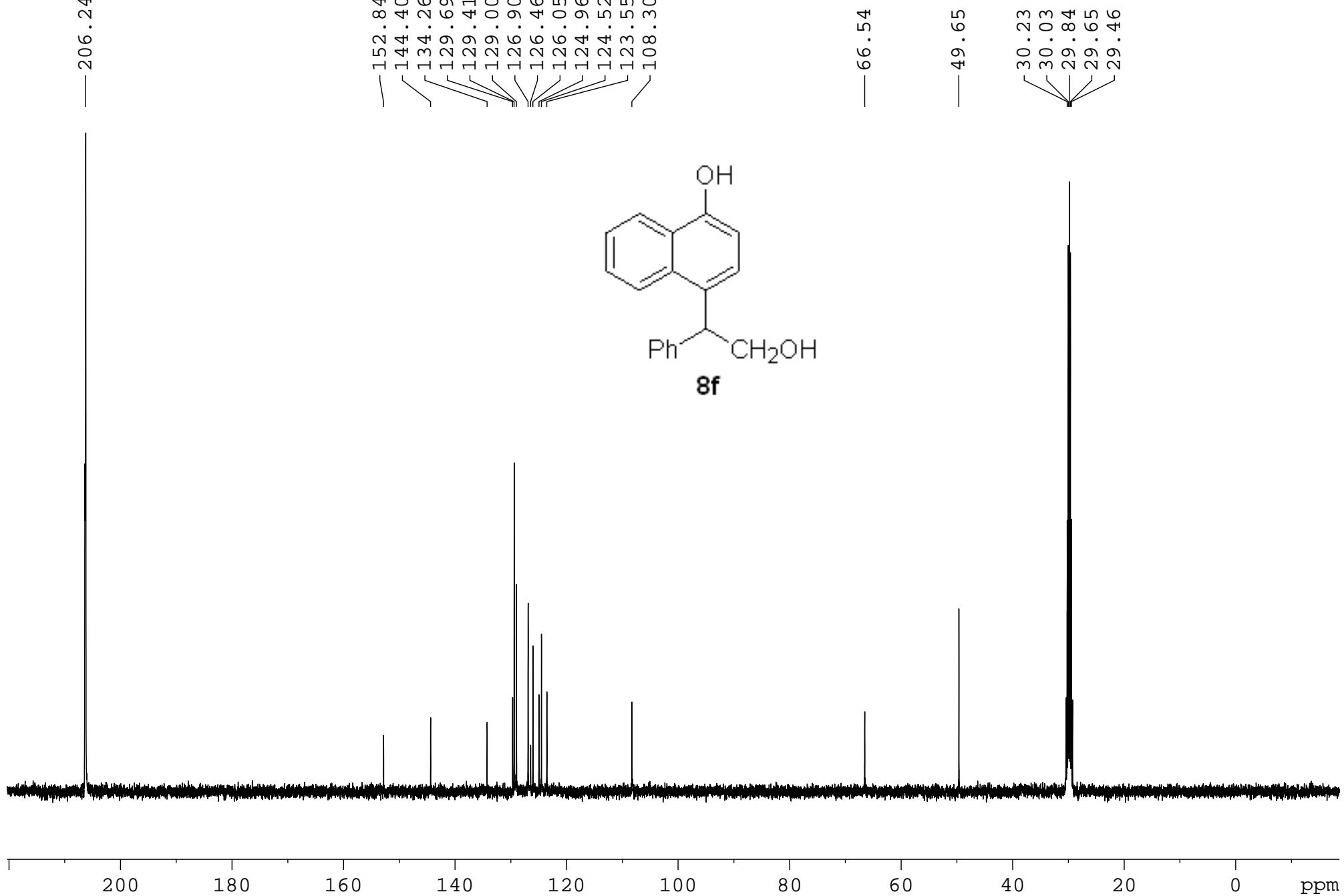


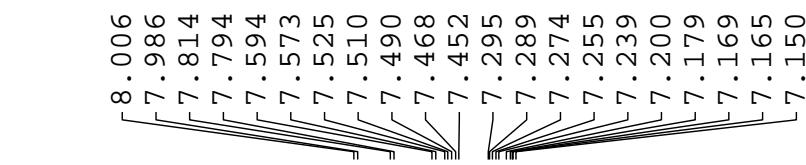


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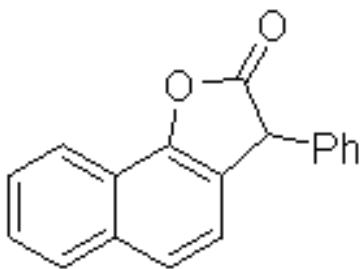


yzz-15-105-1-c

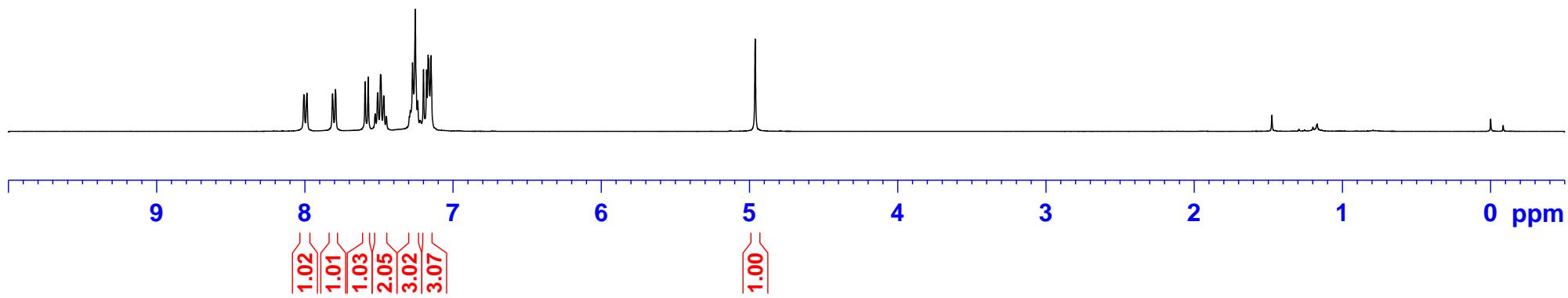




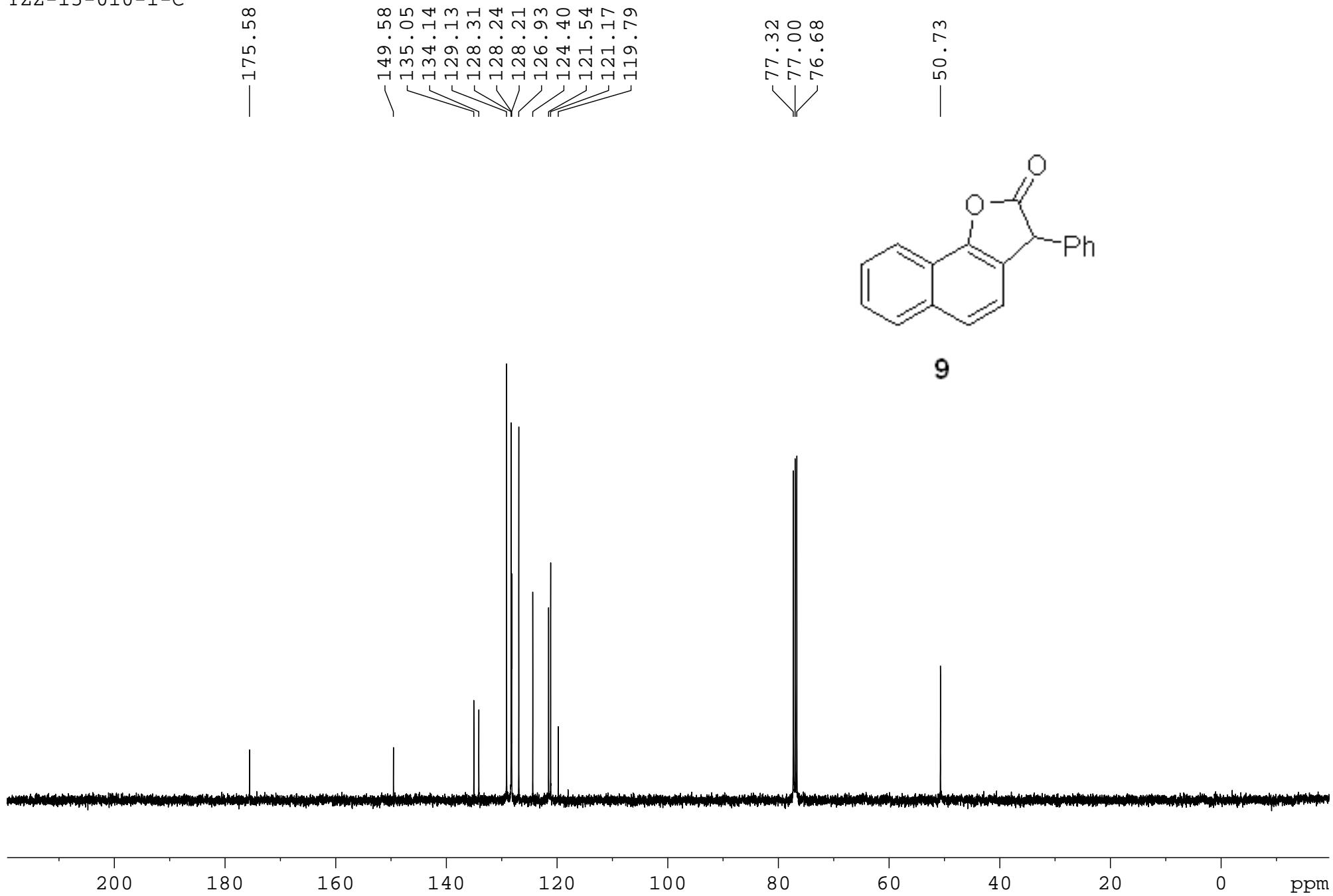
YZZ-15-010-1

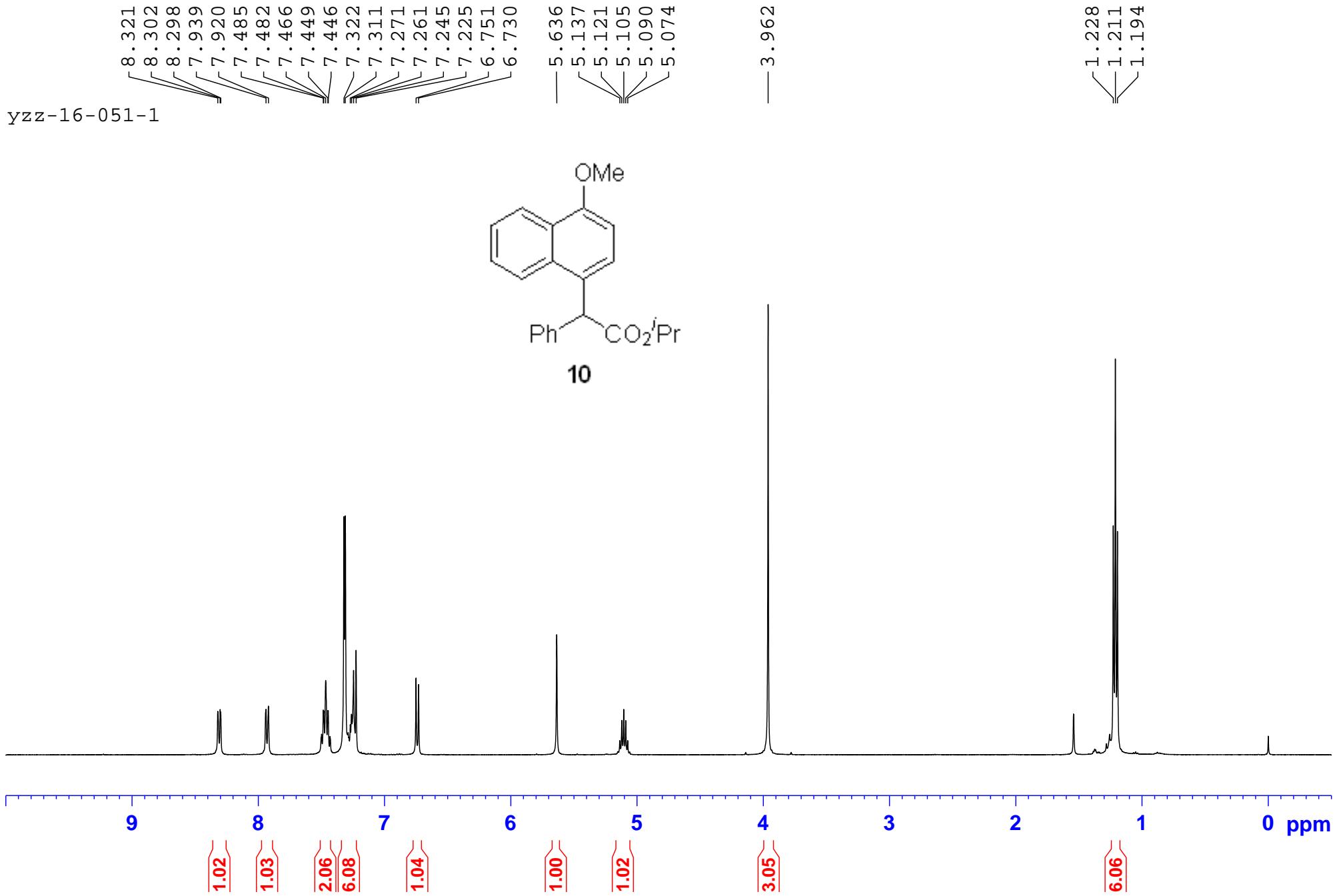


9



YZZ-15-010-1-C





yzz-16-051-1-c

— 172.52

— 154.98

138.49
132.54
128.91
128.50
127.12
126.81
126.31
126.03
124.94
123.08
122.70
— 103.15

77.32
77.00
76.68
— 68.56
55.44
55.41
53.43

— 21.66

