

SUPPORTING INFORMATION

**Gold-catalysed facile access to indene scaffold via sequential
C-H functionalization and 5-*endo-dig* carbocyclization**

Ben Ma, Ziang Wu, Ben Huang, Lu Liu and Junliang Zhang**

*Shanghai Key Laboratory of Green Chemistry and Chemical Processes, School of Chemistry and
Molecular Engineering, East China Normal University, 3663 North Zhongshan Road, Shanghai
200062, P. R. China.*

Fax: (+86)-021-6223-3213; E-mail : lliu@chem.ecnu.edu.cn, jlzhang@chem.ecnu.edu.cn

Content:

1. General Information.....	S2
2. Table S1.....	S3
3. General procedure for the cascade reactions.....	S4
4. Transformation of products.....	S14
5. References.....	S15
6. Crystal structures of 3aa, 8 and 10.....	S16
7. NMR Spectra of new compounds.....	S16

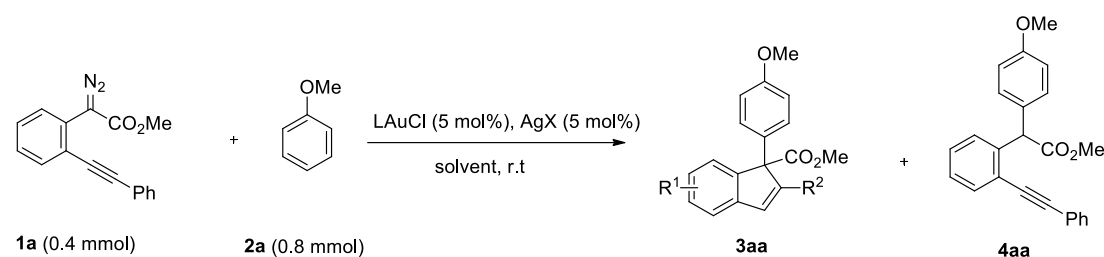
1. General Information:

Unless otherwise noted, all reactions were carried out in standard Schlenk techniques with magnetic stirring bar under air. Materials obtained from commercial suppliers were used directly without further purification. ^1H NMR spectra were recorded on a BRUKER 400 (400 MHz) spectrometer or a Bruker 300 MHz spectrometer in CDCl_3 . Chemical shifts are reported in ppm with tetramethylsilane (TMS: 0 ppm) with the solvent resonance as the internal standard. Data are reported as follows: chemical shift, multiplicity (s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet), coupling constants (Hz), and integration. ^{13}C NMR spectra were recorded on a BRUKER 400 (100 MHz) spectrometer and a Bruker 300 (75.0 MHz) spectrometer in CDCl_3 with complete proton decoupling. Chemical shifts are reported in ppm with the deuterium solvent as the internal standard (e.g. CDCl_3 : 77.0 ppm).

Anhydrous tetrahydrofuran (THF), toluene, 1,2-Dimethoxyethane (DME) and diethyl ether (Et_2O) were distilled from sodium and benzophenone to use; AgOTf , AgSbF_6 , AgNTf_2 , AgBF_4 , and AgPF_6 were purchased from Alfa-Aesar Company and used directly.

Reactions were monitored by thin layer chromatography (TLC) using silicycle pre-coated silica gel plates. Flash column chromatography was performed on silica gel 60 (particle size 200-400 mesh ASTM, purchased from Yantai, China) and eluted with hexane/ethyl acetate or hexane/ CH_2Cl_2 .

2. Table S1. Optimization of Reaction Conditions. ^a



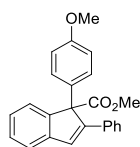
Entry	AuLCl	AgX	Solvent	3 : 4^b	Yield (%) ^c (3aa : 4aa)
1	(2,4- ^t Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgSbF ₆	DCM	5 : 1	(75/15)
2	(2,4- ^t Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgSbF ₆	DCE	6 : 1	77/13
3	(2,4- ^t Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgSbF ₆	CHCl ₃	5 : 1	75/15
4	(2,4- ^t Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgSbF ₆	THF	6 : 1	60/10
5	(2,4- ^t Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgSbF ₆	Et ₂ O	8 : 1	70/9
6	(2,4- ^t Bu ₂ C ₆ H ₃ O) ₃ PAuCl	AgSbF ₆	Toluene	>20 : 1	(80/3)
7	AuCl ₃	AgSbF ₆	DCM	-	-
8	IPrAuCl	AgSbF ₆	DCM	4 : 1	75/17
9	PicAuCl	AgSbF ₆	DCM	2 : 1	50/25
10	Ph ₃ PAuCl	AgSbF ₆	DCM	>20 : 1	(75/3)
11	Ph₃PAuCl	AgOTf	DCM	>20 : 1	(87/3)
12	Ph ₃ PAuCl	AgNTf ₂	DCM	>20 : 1	83/4
13	Ph ₃ PAuCl	AgBF ₄	DCM	15 : 1	80/6
14	Ph ₃ PAuCl	AgOMs	DCM	10 : 1	75/7
16	Ph ₃ PAuCl	AgOTf	DCE	>20 : 1	(85/4)
17	Ph ₃ PAuCl	AgOTf	CHCl ₃	>20 : 1	85/3
18	Ph ₃ PAuCl	AgOTf	THF	5 : 1	65/13
19	Ph ₃ PAuCl	AgOTf	Toluene	>20 : 1	(87/4)
20	-	AgOTf	DCM	<1 : 20	(0/60)
21	Ph ₃ PAuCl	-	DCM	-	-

^a Reaction conditions: **1a** (0.2 mmol), **2a** (0.4 mmol), Ph₃PAuCl (5.5 mol%), AgOTf (5 mol%) in dry solvent (2 mL) at room temperature under Ar. ^b Determined by NMR of crude products. ^c NMR yield. the numbers in parenthesis are isolate yields.

3. General procedure for the cascade reactions

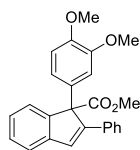
In a dried glass tube, a solution of Ph_3PAuCl (10.97 mg, 0.022 mmol, 5.5 mol%), AgOTf (5.13 mg, 5 mol%) in DCM (2 mL) was stirred at room temperature for 15 mins. After AgCl was filtered off, anisole (86.4 mg, 0.8 mmol) was added to the reaction mixture at room temperature. Then a solution of methyl 2-diazo-2-(2-(phenylethynyl)phenyl)acetate **1a** (110 mg, 0.4 mmol) in 1 mL DCM was introduced into the reaction mixture by a syringe in 5 mins. The resulting mixture was continually stirred at room temperature for 10 min and **1a** was consumed completely determined by TLC analysis. After being filtrated through celite and concentrated, the residue was purified by column chromatography on silica gel to afford the desired product.

1) Methyl 1-(4-methoxyphenyl)-2-phenyl-1H-indene-1-carboxylate (**3aa**)



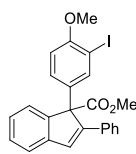
The general procedure was followed using **1a** (0.4 mmol) and **2a** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **3aa** (123.8 mg, 87%) was obtained as a white solid, m.p. 139-140 °C. $^1\text{H NMR}$ (400 MHz, CDCl_3) δ 7.43-7.35 (m, 5H), 7.30-7.16 (m, 7H), 6.75 (d, J = 8.8 Hz, 2H), 3.74 (s, 3H), 3.60 (s, 3H); $^{13}\text{C NMR}$ (100 MHz, CDCl_3) δ 172.5, 158.5, 150.3, 149.2, 142.6, 134.3, 130.3, 129.4, 128.7, 128.2, 127.9, 127.6, 127.1, 126.3, 123.8, 121.5, 113.7, 68.1, 55.1, 52.7; **MS** (EI): m/z (%): 356 (M^+ , 50.66), 297 (100); **HRMS** (EI) calculated for $[\text{C}_{24}\text{H}_{20}\text{O}_3]$: 356.1412, found: 356.1415.

2) Methyl 1-(3, 4-dimethoxyphenyl)-2-phenyl-1H-indene-1-carboxylate (**3ab**)



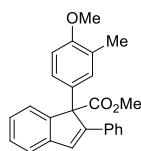
The general procedure was followed using **1a** (0.4 mmol) and **2b** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3ab** (134.4 mg, 87%) was obtained as a white solid, m.p. 149-150 °C. $^1\text{H NMR}$ (400 MHz, CDCl_3) δ 7.42-7.16 (m, 10 H), 6.95-6.91 (m, 2H), 6.70 (d, J = 8.0 Hz, 1H), 3.80 (s, 3H), 3.70 (s, 3H), 3.60 (s, 3H); $^{13}\text{C NMR}$ (100 MHz, CDCl_3) δ 172.5, 150.4, 148.9, 148.5, 148.2, 142.6, 134.5, 130.6, 129.5, 128.2, 128.0, 127.6, 127.2, 126.2, 123.8, 121.6, 119.9, 111.4, 110.9, 68.3, 55.8, 55.7, 52.7; **MS** (EI): m/z (%): 386 (M^+ , 62.46), 327 (100); **HRMS** (EI) calculated for $[\text{C}_{25}\text{H}_{22}\text{O}_4]$: 386.1518, found: 386.1520.

3) Methyl 1-(3-iodo-4-methoxyphenyl)-2-phenyl-1H-indene-1-carboxylate (**3ac**)



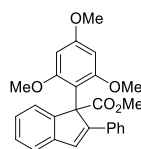
The general procedure was followed using **1a** (0.4 mmol) and **2c** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **3ac** (150.3 mg, 78%) was obtained as a white solid, m.p. 131-132 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.85 (d, *J* = 2.4 Hz, 1H), 7.45-7.10 (m, 11H), 6.61 (d, *J* = 8.8 Hz, 1H), 3.77 (s, 3H), 3.58 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.1, 157.2, 149.9, 148.7, 142.6, 138.5, 134.0, 132.4, 129.6, 128.8, 128.3, 128.1, 127.7, 127.1, 126.4, 123.6, 121.7, 110.5, 85.7, 67.3, 56.2, 52.7; **MS** (EI): *m/z* (%): 482 (M⁺, 83.65), 423 (100); **HRMS** (EI) calculated for [C₂₄H₁₉O₃I]: 482.0379, found: 482.0375.

4) Methyl 1-(4-methoxy-3-methylphenyl)-2-phenyl-1H-indene-1-carboxylate (**3ad**)



The general procedure was followed using **1a** (0.4 mmol) and **2d** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **3ad** (111.1mg, 75%) was obtained as a white solid, m.p. 151-152 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.43-7.35 (m, 5H), 7.28-7.11 (m, 7H), 6.63 (d, *J* = 8.4 Hz, 1H), 3.72 (s, 3H), 3.58 (s, 3H), 2.09 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.6, 156.7, 150.2, 149.3, 142.5, 134.3, 129.71, 129.68, 129.3, 128.1, 127.8, 127.5, 127.1, 126.3, 126.2, 125.9, 123.8, 121.4, 109.5, 68.1, 55.1, 52.6, 16.5; **MS** (EI): *m/z* (%): 370 (M⁺, 42.94), 311 (100); **HRMS** (EI) calculated for [C₂₅H₂₂O₃]: 370.1569, found: 370.1573.

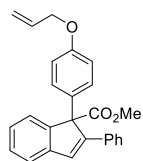
5) Methyl 2-phenyl-1-(2, 4, 6-trimethoxyphenyl)-1H-indene-1-carboxylate (**3ae**).



The general procedure was followed using **1a** (0.4 mmol) and **2e** (0.8 mmol). After purification by column chromatography (PE/EtOAc 3:1), **3ae** (161.4 mg, 97%) was obtained as a white solid, m.p. 189-190 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.52-7.40 (m, 3H), 7.33 (d, *J* = 7.2 Hz, 1H), 7.26-7.04 (m, 6H), 6.21 (s, 1H), 5.98 (s, 1H), 3.78 (s, 3H), 3.73 (s, 3H), 3.53 (s, 3H), 3.11 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 160.2, 159.5, 159.1, 150.2, 148.1, 130.0, 129.7, 128.7, 127.3, 125.1, 124.3, 123.8, 120.6, 120.5, 119.6, 116.7, 109.0, 92.4, 91.9, 55.8, 55.5, 55.4, 55.2; **MS** (EI): *m/z* (%): 416 (M⁺, 40.87), 417 (M⁺, 10.90), 384 (100); **HRMS** (EI) calculated for

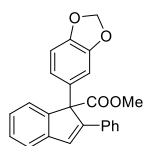
[C₂₆H₂₄O₅]: 416.1624, found: 416.1625.

6) Methyl 1-(4-(allyloxy)phenyl)-2-phenyl-1H-indene-1-carboxylate (3af)



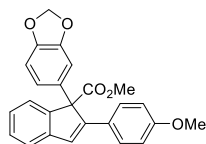
The general procedure was followed using **1a** (0.4 mmol) and **2f** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **3af** (91.6 mg, 60%) was obtained as a white solid, m.p. 109-110 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.46-7.32 (m, 5H), 7.31-7.12 (m, 7H), 6.75 (dd, *J* = 6.4, 2.4 Hz, 2H), 6.05-5.96 (m, 1H), 5.36 (dd, *J* = 17.2, 1.6 Hz, 1H), 5.24 (dd, *J* = 10.8, 1.6 Hz, 1H), 4.45 (dt, *J* = 6.4, 1.6 Hz, 2H), 3.59 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.5, 157.6, 150.3, 149.2, 142.6, 134.2, 133.2, 130.5, 129.4, 128.7, 128.2, 127.9, 127.6, 127.1, 126.3, 123.8, 121.5, 117.7, 114.4, 68.7, 68.1, 52.7; **MS** (EI): *m/z* (%): 382 (M⁺, 100); **HRMS** (EI) calculated for [C₂₆H₂₂O₃]: 382.1569, found: 382.1571.

7) Methyl 1-(benzo[d][1,3]dioxol-5-yl)-2-phenyl-1H-indene-1-carboxylate (3ag)



The general procedure was followed using **1a** (0.4 mmol) and **2g** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **3ag** (121.3 mg, 82%) was obtained as a white solid, m.p. 189-190 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.46-7.34 (m, 5H), 7.30-7.14 (m, 5H), 6.90 (dd, *J* = 8.0, 2.0 Hz, 1H), 6.84 (d, *J* = 2.0 Hz, 1H), 6.64 (d, *J* = 8.0 Hz, 1H), 5.86 (s, 2H), 3.59 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.3, 150.3, 149.1, 147.5, 146.6, 142.5, 134.2, 132.0, 129.5, 128.2, 128.0, 127.6, 127.2, 126.3, 123.8, 121.6, 120.9, 108.2, 108.0, 101.0, 68.4, 52.7; **MS** (EI): *m/z* (%): 370 (M⁺, 54.41), 311 (100); **HRMS** (EI) calculated for [C₂₄H₁₈O₄]: 370.1205, found: 370.1208.

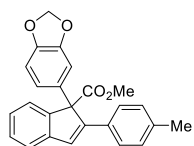
8) Methyl 1-(benzo[d][1,3]dioxol-5-yl)-2-(4-methoxyphenyl)-1H-indene-1-carboxylate (3bg)



The general procedure was followed using **1b** (0.4 mmol) and **2g** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3bg** (139.2 mg, 87%) was obtained as a white solid, m.p. 158-159 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.39-7.31 (m, 6H), 7.26-7.21 (m, 1H), 6.92-6.90 (m, 1H), 6.85-6.80 (m, 3H), 6.65 (d, *J* = 8.0 Hz, 1H), 5.87 (dd, *J* =

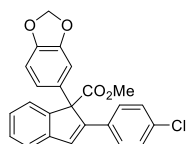
2.4, 1.6 Hz, 2H), 3.78 (s, 3H), 3.59 (s, 3H) ; ^{13}C NMR (100 MHz, CDCl_3) δ 172.4, 159.2, 149.9, 148.9, 147.5, 146.6, 142.8, 132.2, 128.5, 127.9, 127.7, 126.8, 125.9, 123.7, 121.2, 120.9, 113.7, 108.3, 108.0, 101.0, 68.3, 55.2, 52.7; **MS** (EI): m/z (%): 400 (M^+ , 57.60), 341 (100); **HRMS** (EI) calculated for $[\text{C}_{24}\text{H}_{18}\text{O}_4]$: 400.1311, found: 400.1315.

9) Methyl 1-(benzo[d][1,3]dioxol-5-yl)-2-(p-tolyl)-1H-indene-1-carboxylate (3cg)



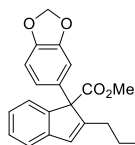
The general procedure was followed using **1c** (0.4 mmol) and **2g** (0.8 mmol). After purification by column chromatography (PE/EtOAc 8:1), **3cg** (130.5 mg, 85%) was obtained as a white solid, m.p. 186-187 °C. ^1H NMR (400 MHz, CDCl_3) δ 7.37-7.24 (m, 6H), 7.16-7.10 (m, 1H), 7.07 (d, J = 8.0 Hz, 2H), 6.91 (dd, J = 8.4, 2.0 Hz, 1H), 6.86 (d, J = 2.0 Hz, 1H), 6.64 (d, J = 8.4 Hz, 1H), 5.85 (s, 2H), 3.58 (s, 3H), 2.30 (s, 3H) ; ^{13}C NMR (100 MHz, CDCl_3) δ 172.3, 150.2, 149.0, 147.5, 146.6, 142.7, 137.6, 132.1, 131.3, 129.0, 128.6, 127.9, 127.0, 126.1, 123.7, 121.4, 120.9, 108.3, 108.0, 100.9, 68.2, 52.6, 21.2; **MS** (EI): m/z (%): 384 (M^+ , 4.65), 152 (100); **HRMS** (EI) calculated for $[\text{C}_{25}\text{H}_{20}\text{O}_4]$: 384.1362, found: 384.1360.

10) Methyl 1-(benzo[d][1,3]dioxol-5-yl)-2-(4-chlorophenyl)-1H-indene-1-carboxylate (3dg)



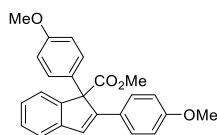
The general procedure was followed using **1d** (0.4 mmol) and **2g** (0.8 mmol). After purification by column chromatography (PE/EtOAc 8:1), **3dg** (121.2 mg, 75%) was obtained as a red solid, m.p. 152-153 °C. ^1H NMR (400 MHz, CDCl_3) δ 7.42-7.15 (m, 9H), 6.86 (dd, J = 8.4, 2.0 Hz, 1H), 6.77 (d, J = 2.0 Hz, 1H), 6.66 (d, J = 8.4 Hz, 1H), 5.89 (dd, J = 2.4, 1.6 Hz, 2H), 3.60 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 172.0, 149.0, 147.7, 146.8, 142.2, 133.4, 132.7, 131.7, 130.0, 128.4, 128.1, 126.6, 124.0, 121.7, 120.7, 108.1, 108.0, 101.1, 68.5, 52.8; **MS** (EI): m/z (%): 404 (M^+ , 43.72), 406 ($[\text{M}+2]^+$, 11.97), 345 (100); **HRMS** (EI) calculated for $[\text{C}_{24}\text{H}_{17}\text{O}_4\text{Cl}]$: 404.0815, found: 404.0811.

11) Methyl 1-(benzo[d][1,3]dioxol-5-yl)-2-propyl-1H-indene-1-carboxylate (3eg)



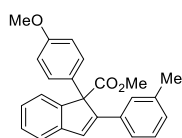
The general procedure was followed using **1e** (0.4 mmol) and **2g** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **3eg** (100.8 mg, 75%) was obtained as a yellow oil. **¹H NMR** (400 MHz, CDCl₃) δ 7.40-7.36 (m, 1H), 7.29-7.23 (m, 2H), 7.15-7.09 (m, 1H), 6.70 (d, *J* = 8.4 Hz, 1H), 6.56-6.50 (m, 2H), 6.43 (d, *J* = 1.6 Hz, 1H), 5.90 (dd, *J* = 2.8, 1.6 Hz, 2H), 3.72 (s, 3H), 2.44-2.34 (m, 1H), 2.10-1.98 (m, 1H), 1.70-1.58 (m, 1H), 1.56-1.47 (m, 1H), 0.93 (t, *J* = 7.2 Hz, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.4, 154.5, 147.6, 146.7, 146.6, 144.3, 133.0, 128.0, 126.4, 125.1, 124.8, 120.6, 120.4, 108.1, 107.8, 101.0, 70.4, 52.6, 30.3, 20.9, 14.1; **MS** (EI): *m/z* (%): 336 (*M*⁺, 100); **HRMS** (EI) calculated for [C₂₁H₂₀O₄]: 336.1362, found: 336.1363.

12) Methyl 1, 2-bis (4-methoxyphenyl)-1H-indene-1-carboxylate (**3ba**)



The general procedure was followed using **1b** (0.4 mmol) and **2a** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3ba** (134.2 mg, 87%) was obtained as a white solid, m.p. 157-158 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.40-7.24 (m, 8H), 7.14-7.01 (m, 1H), 6.81-6.72 (m, 4H), 3.77 (s, 3H), 3.73 (s, 3H), 3.59 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.7, 159.1, 158.5, 150.0, 149.1, 142.9, 130.6, 128.7, 128.5, 127.8, 127.6, 127.0, 125.8, 123.6, 121.1, 113.7, 113.6, 68.1, 55.2, 55.1, 52.6; **MS** (EI): *m/z* (%): 386 (*M*⁺, 55.93), 327 (100); **HRMS** (EI) calculated for [C₂₅H₂₂O₄]: 386.1518, found: 386.1522.

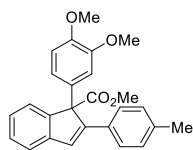
13) Methyl 1-(4-methoxyphenyl)-2-(*m*-tolyl)-1H-indene-1-carboxylate (**3fa**)



The general procedure was followed using **1f** (0.4 mmol) and **2g** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **3fa** (119.8 mg, 81%) was obtained as a white solid, m.p. 142-143 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.39-7.24 (m, 7H), 7.16-7.01 (m, 4H), 6.75-6.71 (m, 2H), 3.72 (s, 3H), 3.59 (s, 3H), 2.29 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.6, 158.5, 150.6, 149.2, 142.7, 137.6, 134.2, 130.4, 129.2, 128.8, 128.4, 128.1, 127.9, 127.7, 126.2, 124.4, 123.7, 121.4, 113.6, 68.1, 55.1, 52.6, 21.5; **MS** (EI): *m/z* (%): 370 (*M*⁺,

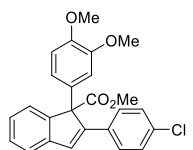
53.30), 311 (100); **HRMS** (EI) calculated for [C₂₅H₂₂O₃]: 370.1569, found: 370.1565.

14) Methyl 1-(3,4-dimethoxyphenyl)-2-(p-tolyl)-1H-indene-1-carboxylate (3cb)



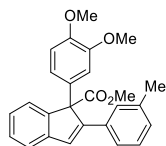
The general procedure was followed using **1c** (0.4 mmol) and **2b** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3cb** (120 mg, 75%) was obtained as a white solid, m.p. 132-133 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.45-7.26 (m, 6H), 7.17-7.13 (m, 1H), 7.07 (d, *J* = 2.0 Hz, 2H), 6.98 (d, *J* = 2.0 Hz, 1H), 6.93 (dd, *J* = 8.4, 2.0 Hz, 1H), 6.69 (d, *J* = 8.4 Hz, 1H), 3.80 (s, 3H), 3.72 (s, 3H), 3.59 (s, 3H), 2.31 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.6, 150.2, 148.8, 148.3, 147.9, 142.7, 137.5, 131.4, 130.7, 128.9, 128.6, 127.9, 127.0, 126.0, 123.6, 121.4, 119.7, 111.1, 110.6, 68.0, 55.7, 55.6, 52.7, 21.2; **MS** (EI): *m/z* (%): 400 (M⁺, 5.02), 311 (100); **HRMS** (EI) calculated for [C₂₆H₂₄O₄]: 400.1675, found: 400.1673.

15) Methyl 2-(4-chlorophenyl)-1-(3,4-dimethoxyphenyl)-1H-indene-1-carboxylate (3db)



The general procedure was followed using **1d** (0.4 mmol) and **2b** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3db** (120.9 mg, 72%) was obtained as a white solid, m.p. 156-157 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.43-7.15 (m, 9H), 6.91-6.86 (m, 2H), 6.71 (d, *J* = 9.2 Hz, 1H), 3.81 (s, 3H), 3.72 (s, 3H), 3.61 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.2, 148.9, 148.8, 148.5, 148.1, 142.2, 133.4, 132.8, 130.2, 130.0, 128.4, 128.3, 128.0, 126.5, 123.8, 121.7, 119.6, 110.8, 68.3, 55.7, 55.6, 52.8; **MS** (EI): *m/z* (%): 420 (M⁺, 7.42), 422 ([M+2]⁺, 2.89), 311 (100); **HRMS** (EI) calculated for [C₂₅H₂₁O₄Cl]: 420.1128, found: 420.1125.

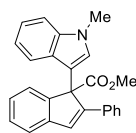
16) Methyl 1-(3,4-dimethoxyphenyl)-2-(m-tolyl)-1H-indene-1-carboxylate (3fb)



The general procedure was followed using **1f** (0.4 mmol) and **2b** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3fb** (115.2 mg, 72%) was obtained as a white solid, m.p. 158-159 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.40-7.26 (m, 5H), 7.16-7.13 (m, 3H), 7.04-7.03 (m, 1H), 6.98 (d, *J* = 2.0 Hz, 1H), 6.90 (dd, *J* = 8.4, 2.0 Hz, 1H), 6.68 (d, *J* = 8.4

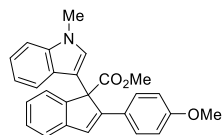
Hz, 1H), 3.79 (s, 3H), 3.71 (s, 3H), 3.60 (s, 3H), 2.30 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 172.5, 150.5, 148.8, 148.3, 148.0, 142.6, 137.6, 134.2, 130.6, 129.2, 128.5, 128.0, 127.9, 127.7, 126.1, 124.4, 123.6, 121.5, 119.8, 111.2, 110.6, 68.1, 55.7, 55.6, 52.7, 21.5; **MS** (EI): m/z (%): 400 (M^+ , 65.2), 341 (100); **HRMS** (EI) calculated for $[\text{C}_{26}\text{H}_{24}\text{O}_4]$: 400.1675, found: 400.1672.

17) Methyl 1-(1-methyl-1H-indol-3-yl)-2-phenyl-1H-indene-1-carboxylate (3ah)



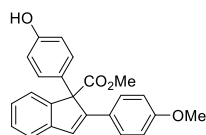
The general procedure was followed using **1a** (0.4 mmol) and **2h** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3ah** (125.8 mg, 83%) was obtained as a white solid, m.p. 89-90 °C. ^1H NMR (400 MHz, CDCl_3) δ 7.69 (s, 1H), 7.50-7.44 (m, 4H), 7.34-7.00 (m, 8H), 6.92 (d, J = 8.0 Hz, 1H), 6.78-6.76 (m, 1H), 3.74 (s, 3H), 3.59 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 172.9, 149.8, 148.4, 142.8, 136.8, 134.5, 129.2, 128.9, 128.2, 127.9, 127.4, 126.6, 126.2, 125.8, 123.5, 121.4, 121.1, 120.2, 118.8, 109.8, 109.0, 63.8, 52.7, 32.9; **MS** (EI): m/z (%): 379 (M^+ , 42.72), 320 (100); **HRMS** (EI) calculated for $[\text{C}_{26}\text{H}_{21}\text{NO}_2]$: 379.1572, found: 379.1576.

18) Methyl 2-(4-methoxyphenyl)-1-(1-methyl-1H-indol-3-yl)-1H-indene-1-carboxylate (3bh)



The general procedure was followed using **1b** (0.4 mmol) and **2h** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **3bh** (139.1 mg, 85%) was obtained as a yellow solid, m.p. 167-168 °C. ^1H NMR (400 MHz, CDCl_3) δ 7.66 (s, 1H), 7.54-7.40 (m, 4H), 7.34-7.20 (m, 2H), 7.16 (d, J = 8.0 Hz, 1H), 7.06-6.95 (m, 3H), 6.79-6.70 (m, 3H), 3.73 (s, 3H), 3.71 (s, 3H), 3.58 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 173.0, 159.0, 149.6, 148.3, 143.1, 136.9, 129.1, 127.9, 127.8, 127.4, 127.1, 125.9, 125.7, 123.4, 121.2, 121.0, 120.3, 118.9, 113.7, 110.3, 108.9, 63.9, 55.1, 52.6, 32.8; **MS** (EI): m/z (%): 409 (M^+ , 52.13), 350 (100); **HRMS** (EI) calculated for $[\text{C}_{27}\text{H}_{23}\text{NO}_3]$: 409.1678, found: 409.1672.

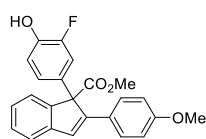
19) Methyl 1-(4-hydroxyphenyl)-2-(4-methoxyphenyl)-1H-indene-1-carboxylate (6a)



The general procedure was followed using **1b** (0.4 mmol) and **5a** (0.8 mmol).

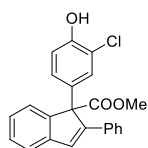
After purification by column chromatography (PE/EtOAc 3:1), **6ba** (108.6 mg, 73%) was obtained as a white solid, m.p. 90-91 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.36-7.31 (m, 4H), 7.27-7.21 (m, 4H), 7.13-7.09 (m, 1H), 6.68 (dd, *J* = 6.8, 2.0 Hz, 2H), 6.64 (dd, *J* = 6.8, 2.0 Hz, 2H), 5.18 (s, 1H), 3.76 (s, 3H), 3.58 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.8, 159.1, 154.7, 149.9, 149.0, 142.9, 130.6, 128.9, 128.5, 127.8, 127.6, 127.0, 125.8, 123.6, 121.2, 115.2, 113.7, 68.1, 55.2, 52.7; **MS** (EI): *m/z* (%): 372 (M⁺, 55.42), 313 (100); **HRMS** (EI) calculated for [C₂₄H₂₀O₄]: 372.1362, found: 372.1368.

20) Methyl 1-(3-fluoro-4-hydroxyphenyl)-2-(4-methoxyphenyl)-1H-indene-1-carboxylate (**6b**)



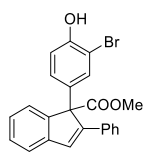
The general procedure was followed using **1b** (0.4 mmol) and **5b** (0.8 mmol). After purification by column chromatography (PE/EtOAc 3:1), **6bb** (117.0 mg, 75%) was obtained as a red oil. **¹H NMR** (400 MHz, CDCl₃) δ 7.37-7.19 (m, 7H), 7.15-7.11 (m, 1H), 7.07-7.04 (m, 1H), 6.83-6.79 (m, 3H), 5.13 (d, *J* = 2.4 Hz, 1H), 3.78 (s, 3H), 3.58 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.3, 159.3, 149.5 (d, *J* = 18.1 Hz), 148.6, 142.9, 142.5 (d, *J* = 14.3 Hz), 131.5 (d, *J* = 6.0 Hz), 128.4, 128.1, 127.7, 126.6, 125.9, 124.0, 123.5, 121.3, 116.9, 115.1 (d, *J* = 20.2 Hz), 113.8, 67.6, 55.2, 52.7; **¹⁹F NMR** (282 MHz, CDCl₃) δ -139.977; **MS** (EI): *m/z* (%): 390 (M⁺, 56.80), 84 (100); **HRMS** (EI) calculated for [C₂₄H₁₉O₄F]: 390.1267, found: 390.1264.

21) Methyl 1-(3-chloro-4-hydroxyphenyl)-2-phenyl-1H-indene-1-carboxylate (**6c**)



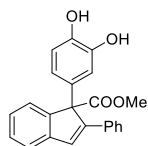
The general procedure was followed using **1a** (0.4 mmol) and **5c** (0.8 mmol). After purification by column chromatography (PE/EtOAc 3:1), **6ac** (108.2 mg, 72%) was obtained as a white solid, m.p. 136-137 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.45 (d, *J* = 2.4 Hz, 1H), 7.41-7.38 (m, 4H), 7.34-7.22 (m, 5H), 7.19-7.14 (m, 2H), 6.82 (d, *J* = 8.8 Hz, 1H), 5.48 (s, 1H), 3.59 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.1, 150.4, 149.9, 148.7, 142.6, 133.9, 131.6, 129.6, 128.3, 128.2, 128.1, 127.8, 127.7, 127.0, 126.4, 123.6, 121.7, 119.6, 116.0, 67.5, 52.8; **MS** (EI): *m/z* (%): 376 (M⁺, 49.84), 378 ([M+2]⁺, 15.68), 317 (100); **HRMS** (EI) calculated for [C₂₃H₁₇O₃Cl]: 376.0866, found: 376.0869.

22) Methyl 1-(3-bromo-4-hydroxyphenyl)-2-phenyl-1H-indene-1-carboxylate (6d)



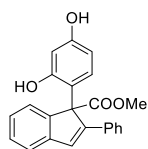
The general procedure was followed using **1a** (0.4 mmol) and **5d** (0.8 mmol). After purification by column chromatography (PE/EtOAc 3:1), **6ad** (132.7 mg, 79%) was obtained as a white solid, m.p. 189-190 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.60 (d, *J* = 2.4 Hz, 1H), 7.42-7.16 (m, 11H), 6.83 (d, *J* = 8.4 Hz, 1H), 5.42 (s, 1H), 3.59 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.1, 151.3, 149.9, 148.7, 142.6, 133.9, 132.0, 131.0, 129.6, 128.6, 128.4, 128.2, 127.8, 127.0, 126.5, 123.6, 121.7, 115.8, 110.0, 67.4, 52.8; **MS** (EI): *m/z* (%): 420 (M⁺, 52.16), 422 ([M+2]⁺, 52.41), 361 (100); **HRMS** (EI) calculated for [C₂₃H₁₇O₃Br]: 420.0361, found: 420.0368.

23) Methyl 1-(3, 4-dihydroxyphenyl)-2-phenyl-1H-indene-1-carboxylate (6e)



The general procedure was followed using **1a** (0.4 mmol) and **6e** (0.8 mmol). After purification by column chromatography (PE/EtOAc 2:1), **6ae** (97.4 mg, 68%) was obtained as a white solid, m.p. 100-101 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.47-7.15 (m, 10H), 6.93 (d, *J* = 2.0 Hz, 1H), 6.86 (dd, *J* = 8.4, 2.0 Hz, 1H), 6.70 (d, *J* = 8.4 Hz, 1H), 5.48 (br, 1H), 5.42 (br, 1H), 3.60 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.5, 150.3, 149.1, 143.1, 142.9, 142.5, 134.3, 131.0, 129.4, 128.2, 128.0, 127.6, 127.2, 126.3, 123.8, 121.5, 120.5, 115.2, 114.9, 68.1, 52.7; **MS** (EI): *m/z* (%): 358 (M⁺, 63.92), 299 (100); **HRMS** (EI) calculated for [C₂₃H₁₈O₄]: 358.1205, found: 358.1201.

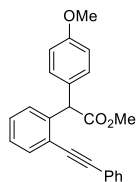
24) Methyl 1-(2, 4-dihydroxyphenyl)-2-phenyl-1H-indene-1-carboxylate (6f)



The general procedure was followed using **1a** (0.4 mmol) and **5f** (0.8 mmol). After purification by column chromatography (PE/EtOAc 5:1), **6af** (90.2 mg, 63%) was obtained as a white solid, m.p. 128-129 °C. **¹H NMR** (400 MHz, DMSO-*d*₆) δ 9.78 (s, 1H), 9.22 (s, 1H), 7.69 (t, *J* = 7.2 Hz, 3H), 7.51 (s, 1H), 7.38-7.13 (m, 6H), 6.42-6.36 (m, 2H), 5.98 (dd, *J* = 8.4, 2.0 Hz, 1H), 3.45 (s, 3H), 3.38 (s, 3H); **¹³C NMR** (100 MHz, DMSO-*d*₆) δ 171.4, 157.5, 156.3, 150.4, 148.0, 141.8, 134.1, 130.7, 128.0, 127.8, 127.5, 127.3, 127.2, 125.9, 124.2, 121.2, 116.4, 105.9, 103.4,

66.9, 51.9 ; **MS** (EI): m/z (%): 358 (M^+ , 9.51), 59 (100); **HRMS** (EI) calculated for $[C_{23}H_{18}O_4]$: 358.1205, found: 358.1209.

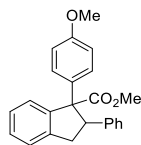
25) Methyl 2-(4-methoxyphenyl)-2-(2-(phenylethynyl)phenyl)acetate (4aa)



The general procedure was followed using **1f** (0.4 mmol) and **2b** (0.8 mmol). After purification by column chromatography (PE/EtOAc 10:1), **4aa** (4.7 mg, 3%) was obtained as a yellow oil. **¹H NMR** (400 MHz, $CDCl_3$) δ 7.57-7.50 (m, 3H), 7.38-7.22 (m, 8H), 6.87 (dd, J = 6.8, 2.4 Hz, 2H), 5.60 (s, 1H), 3.76 (s, 3H), 3.70 (s, 3H); **¹³C NMR** (100 MHz, $CDCl_3$) δ 173.0, 158.8, 140.9, 132.2, 131.5, 129.9, 129.7, 128.5, 128.4, 128.3, 128.1, 127.0, 123.0, 114.0, 94.5, 87.5, 55.2, 54.2, 52.3; **MS** (EI): m/z (%): 356 (M^+ , 6.94), 297 (100); **HRMS** (EI) calculated for $[C_{23}H_{18}O_4]$: 356.1412, found: 356.1414.

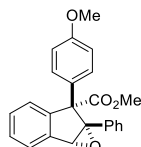
4. Transformation of products

1) Methyl 1-(4-methoxyphenyl)-2-phenyl-2,3 dihydro-1H-indene-1-carboxylate (7)



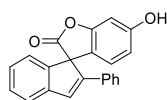
The solution of compound **3aa** (0.2 mmol) and Pd/C (contain 10% Pd) (5 mol%) in CH₃OH (2 mL) under 1atm pressure of hydrogen at room temperature for 5 hours and the reaction purified by silica chromatography, **7** (68.7 mg, 96%) was obtained as a colorless oil. **¹H NMR** (400 MHz, CDCl₃) δ 7.52-7.50 (m, 1H), 7.33-7.29 (m, 3H), 7.22-7.16 (m, 5H), 7.05-7.03 (m, 2H), 6.85-6.81 (m, 2H), 3.95 (t, *J* = 6.8 Hz, 1H), 3.79 (s, 3H), 3.42 (dd, *J* = 15.2, 6.4 Hz, 1H), 3.32 (s, 3H), 3.27 (dd, *J* = 15.2, 7.2 Hz, 1H); **¹³C NMR** (100 MHz, CDCl₃) δ 172.7, 158.5, 143.5, 141.2, 134.4, 129.1, 128.6, 127.8, 127.7, 127.5, 126.7, 126.6, 124.7, 113.4, 68.4, 58.7, 55.2, 51.4, 37.4; **MS** (EI): *m/z* (%): 358 (M⁺, 12.07), 299 (100); **HRMS** (EI) calculated for [C₂₄H₂₂O₃]: 358.1569, found: 358.1573.

2) Methyl 1-(4-methoxyphenyl)-2-phenyl-2,3 dihydro-1H-indene-1-carboxylate (8)



Added *m*-CPBA (0.4 mmol, 2 eq) to the solution of compound **3aa** (0.2 mmol) in CH₂Cl₂ (2 mL), and then reflux for 5 hours **3aa** was consumed completely determined by TLC analysis. Purified by silica chromatography, **8** (dr: 1.4 : 1, 68.7 mg, 75%) was obtained as a white solid (only one isomer can isolated 23.8 mg, 32%), m.p. 185-186 °C. **¹H NMR** (400 MHz, CDCl₃) δ 7.59 (dd, *J* = 10.0, 7.2 Hz, 2H), 7.41-7.35 (m, 2H), 7.18-7.10 (m, 5H), 6.64-6.56 (m, 4H), 4.27 (s, 1H), 3.83 (s, 3H), 3.72 (s, 3H); **¹³C NMR** (100 MHz, CDCl₃) δ 171.7, 158.8, 145.6, 140.9, 134.6, 131.6, 129.4, 129.3, 129.2, 128.9, 127.7, 127.6, 127.1, 124.6, 113.4, 73.5, 68.2, 64.4, 55.2, 52.5; **MS** (EI): *m/z* (%): 372 (M⁺, 6.37), 313 (100); **HRMS** (EI) calculated for [C₂₄H₂₀O₄]: 372.1362, found: 372.1359.

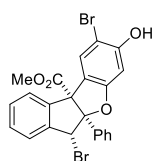
3) 6-hydroxy-2'-phenyl-2H-spiro[benzofuran-3,1'-inden]-2-one (9)²



The solution of compound **5a** (0.2 mmol) and TFA (0.21 mmol) in toluene (6 mL) was heated to 90 °C for 3 hours and the reaction purified by silica chromatography, **9** (60.6 mg, 93%) was obtained as a white solid, m.p. 167-168 °C. **¹H NMR** (400 MHz, DMSO-*d*₆) δ 10.1 (s,

1H), 7.81 (s, 1H), 7.55 (d, $J = 7.6$ Hz, 1H), 7.38-7.18 (m, 7H), 6.99 (d, $J = 7.6$ Hz, 1H), 6.92 (d, $J = 2.0$ Hz, 1H), 6.61 (d, $J = 8.4$ Hz, 1H), 6.62 (dd, $J = 8.4, 2.4$ Hz, 1H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 174.9, 159.0, 154.4, 146.4, 145.5, 143.1, 138.8, 132.1, 129.0, 128.8, 128.2, 126.8, 125.4, 124.1, 122.1, 121.9, 117.0, 112.3, 99.3, 62.8; **MS** (EI): m/z (%): 326 (M^+ , 4.14), 197 (100); **HRMS** (EI) calculated for $[\text{C}_{22}\text{H}_{14}\text{O}_3]$: 326.0943, found: 326.0948.

4) Methyl 2,6-dibromo-3-hydroxy-5a-phenyl-5a,6-dihydro-10bH-indeno[2,1-b]benzofuran-10b-carboxylate-6-hydroxy-2'-phenyl-2H-spiro[benzofuran-3,1'-inden]-2-one (10)

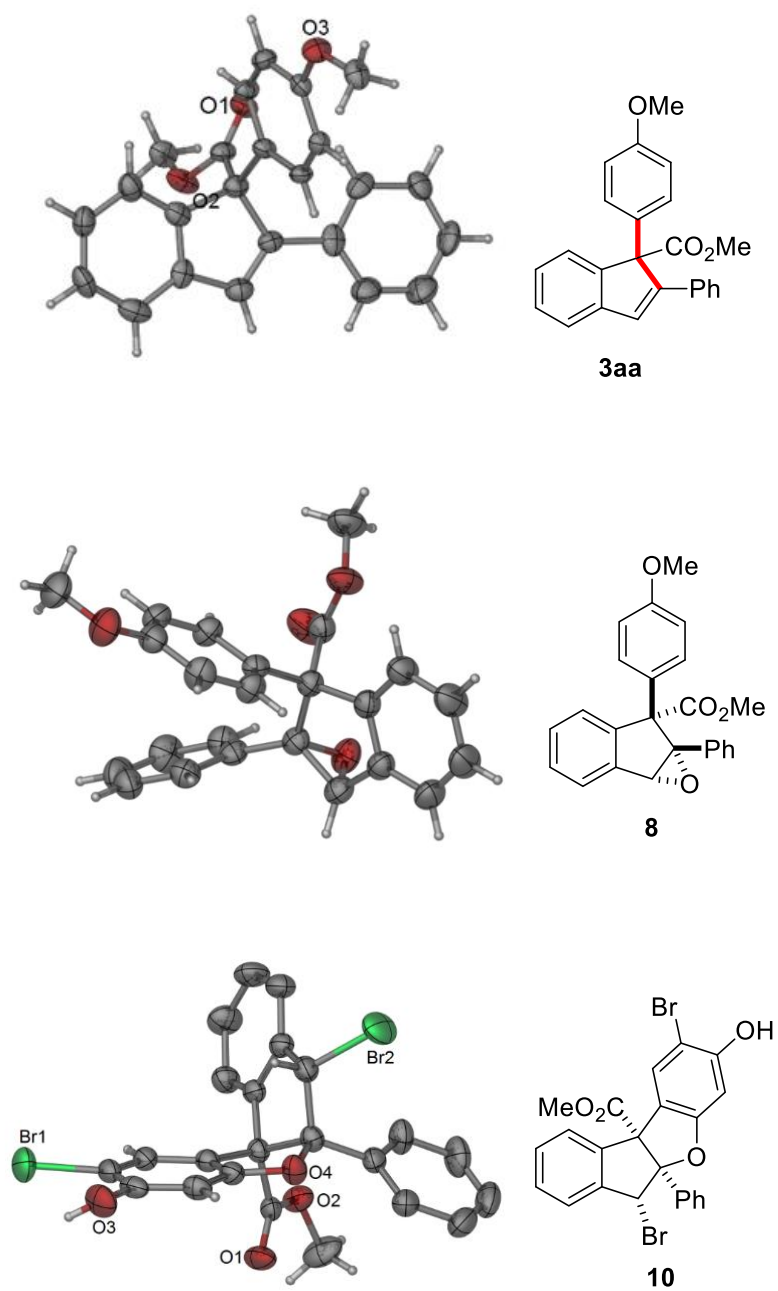


To the solution of compound **5a** (0.2 mmol) in CH_2Cl_2 (1 mL) added Br_2 (dissolved in 1 mL CH_2Cl_2) slowly, then the mixture stirred for 30 mins monitored by TLC and quenched by saturated solution of $\text{Na}_2\text{S}_2\text{O}_3$ at room temperature. The extracts with ethyl acetate were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude product was purified by column chromatography to give white solid **10** (60.6 mg, 93%), m.p. 177-178 °C. ^1H NMR (400 MHz, CDCl_3) δ 7.76 (d, $J = 7.6$ Hz, 1H), 7.75-7.44 (m, 3H), 7.28-7.24 (m, 5H), 7.16 (s, 1H), 6.80 (s, 1H), 5.78 (s, 1H), 5.62 (s, 1H), 3.11 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 169.8, 159.0, 153.7, 140.3, 139.1, 137.2, 129.8, 129.0, 128.5, 127.6, 126.4, 125.8, 125.5, 125.0, 122.2, 106.9, 102.4, 99.9, 69.6, 61.2, 52.2; **MS** (EI): m/z (%): 514 (M^+ , 23.23), 516 ($[\text{M}+2]^+$, 59.66), 518 ($[\text{M}+4]^+$, 30.47), 377 (100); **HRMS** (EI) calculated for $[\text{C}_{23}\text{H}_{16}\text{Br}_2\text{O}_4]$: 516.1543, found: 516.1548.

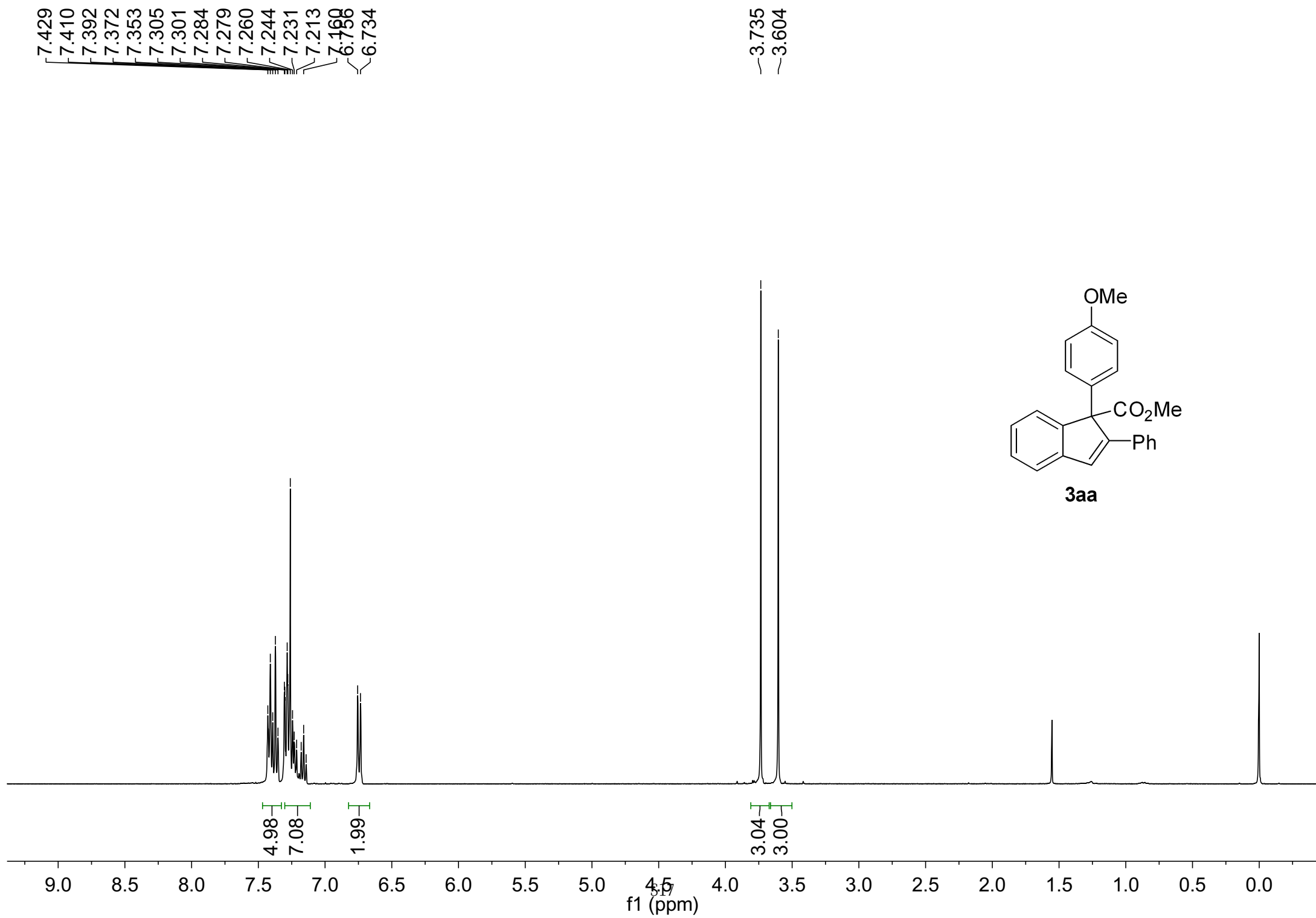
5 References

- [1] P. Cheng, J.-J. Cheng and J.-B. Wang, *Adv. Synth. Catal.*, 2008, **350**, 2359.
- [2] X. Li, S.-S. Hu, Z.-G. Xi, L. Zhang, S.-Z. Luo and J.-P. Cheng. *J. Org. Chem.* 2010, **75**, 8697.

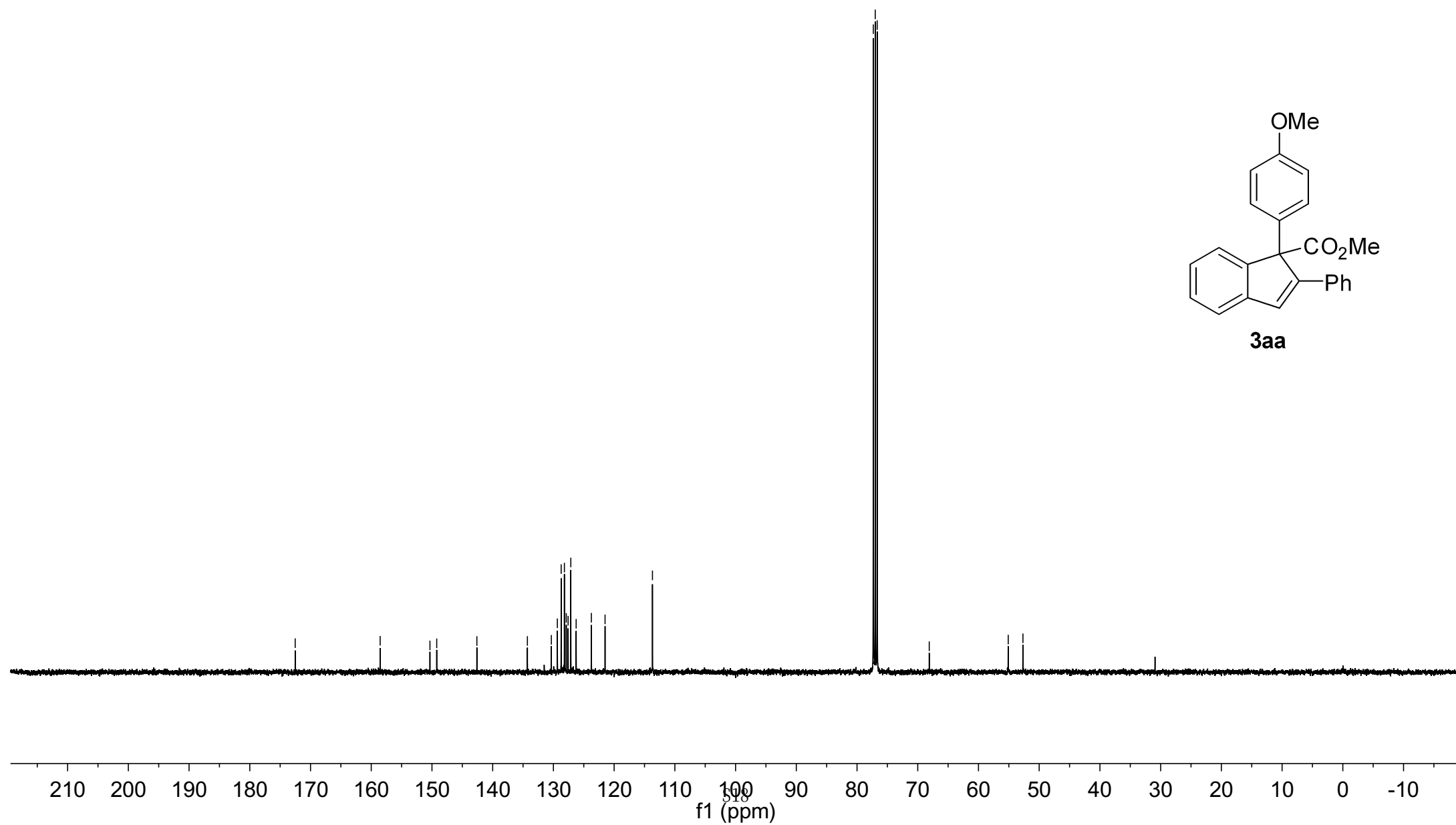
6 Figure S1. X-ray structure of 3aa, 8 and 10.

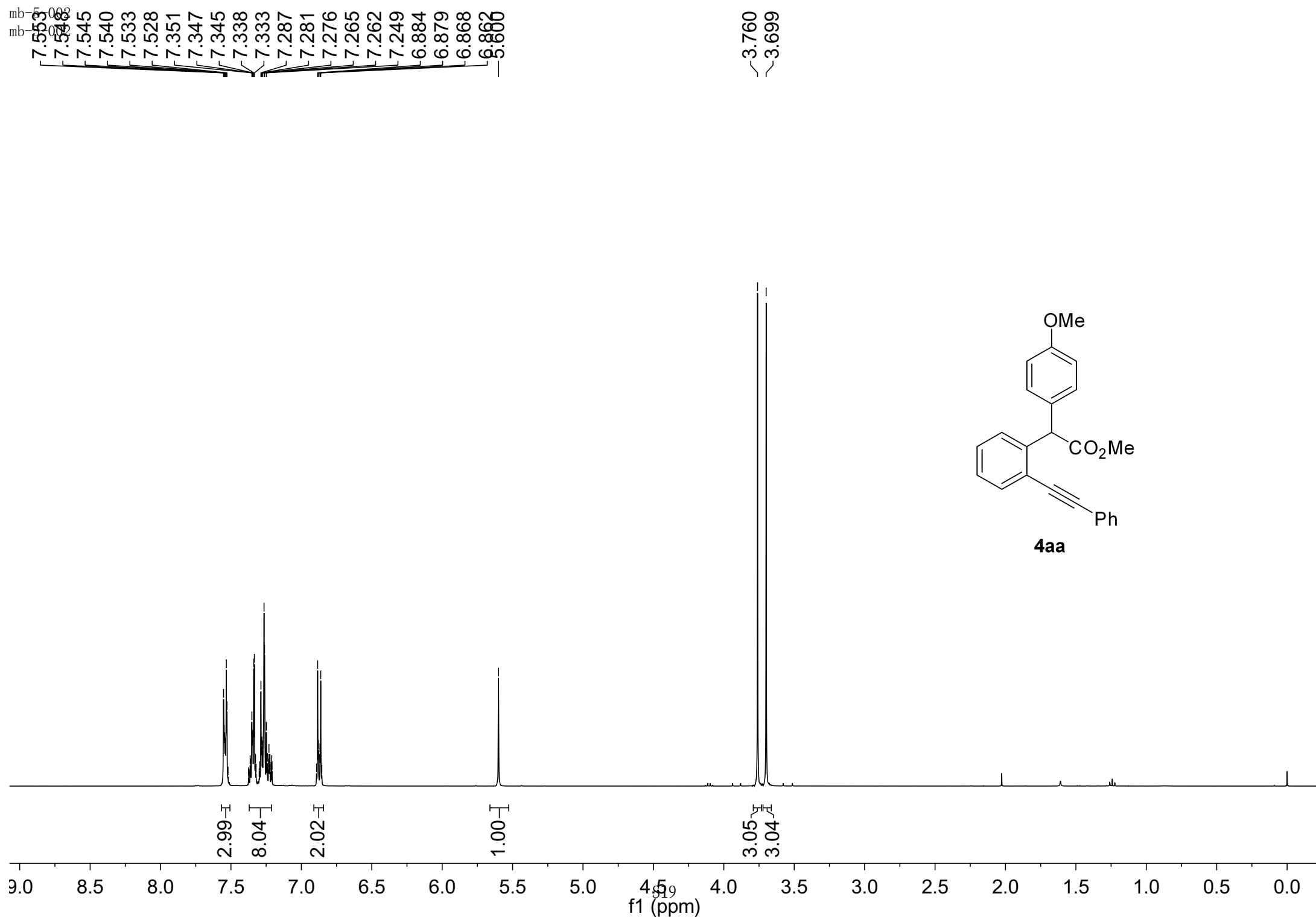


7 NMR spectra of new compounds

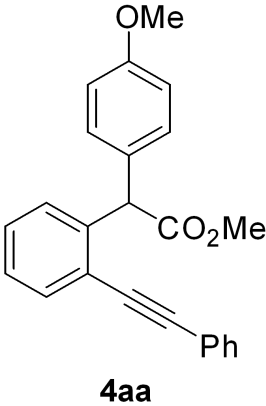
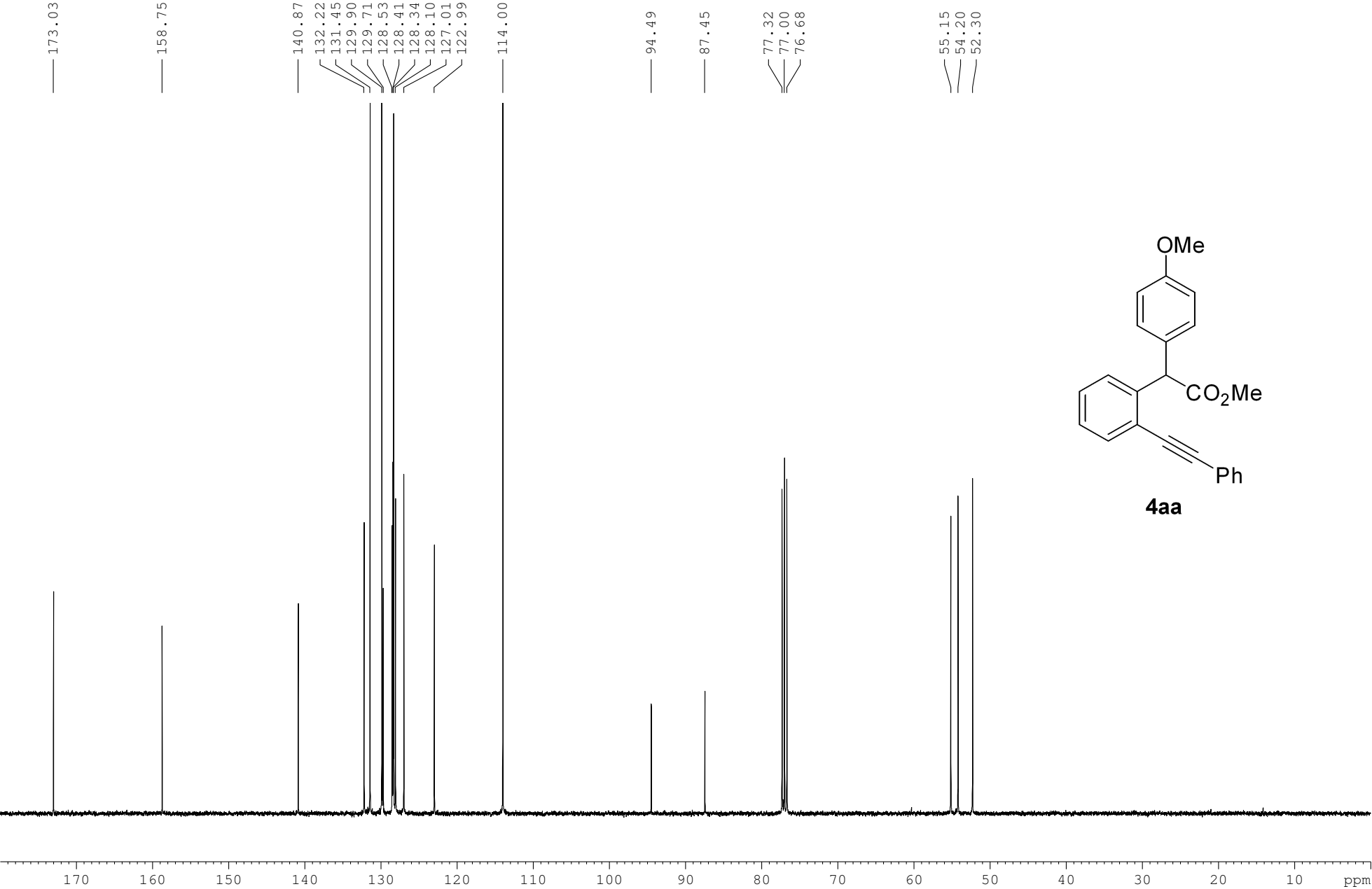


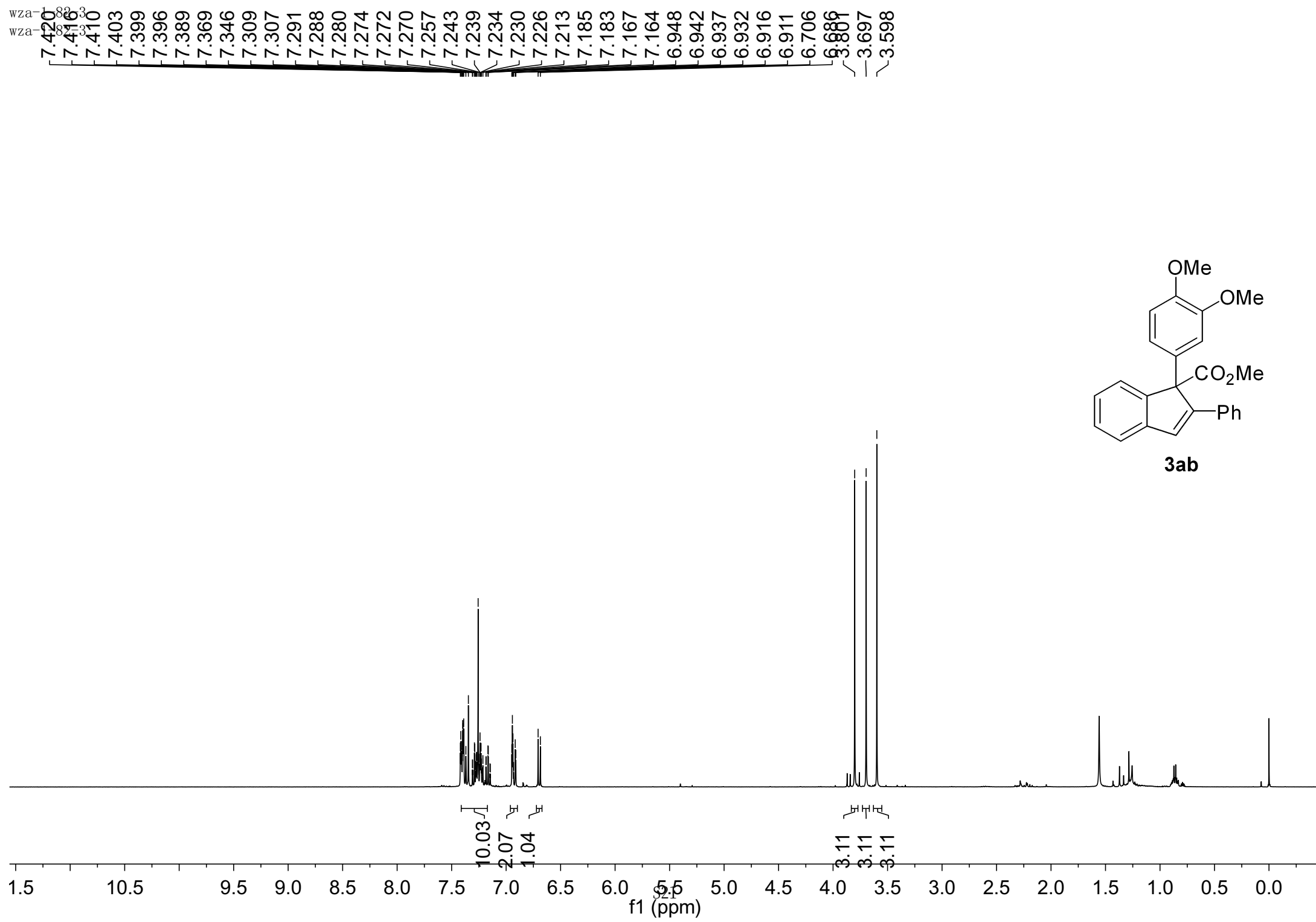
mb-3-5
mb-3-5 C

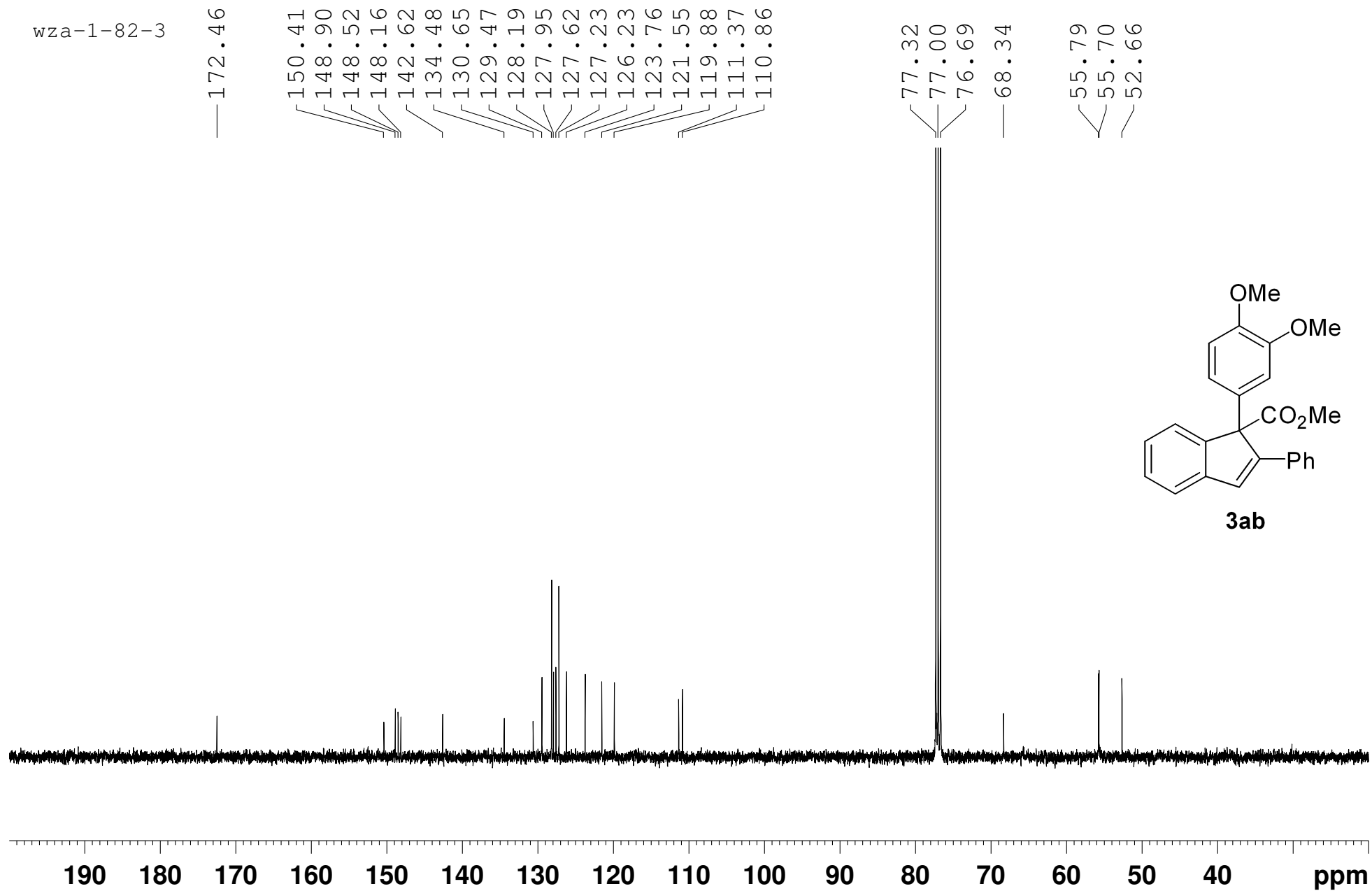




mb-5-002





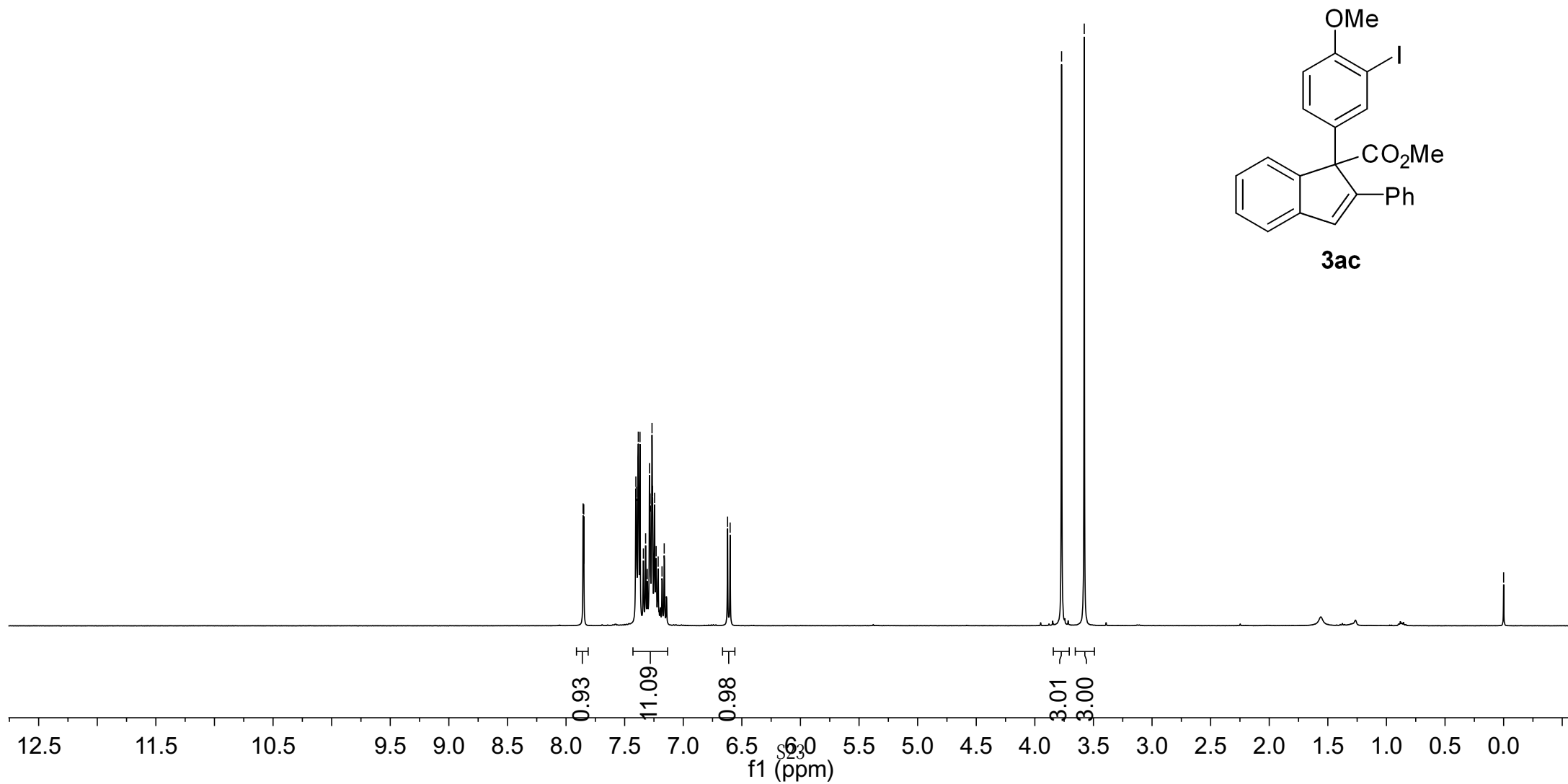


mbw-1-92
mbw-1-92

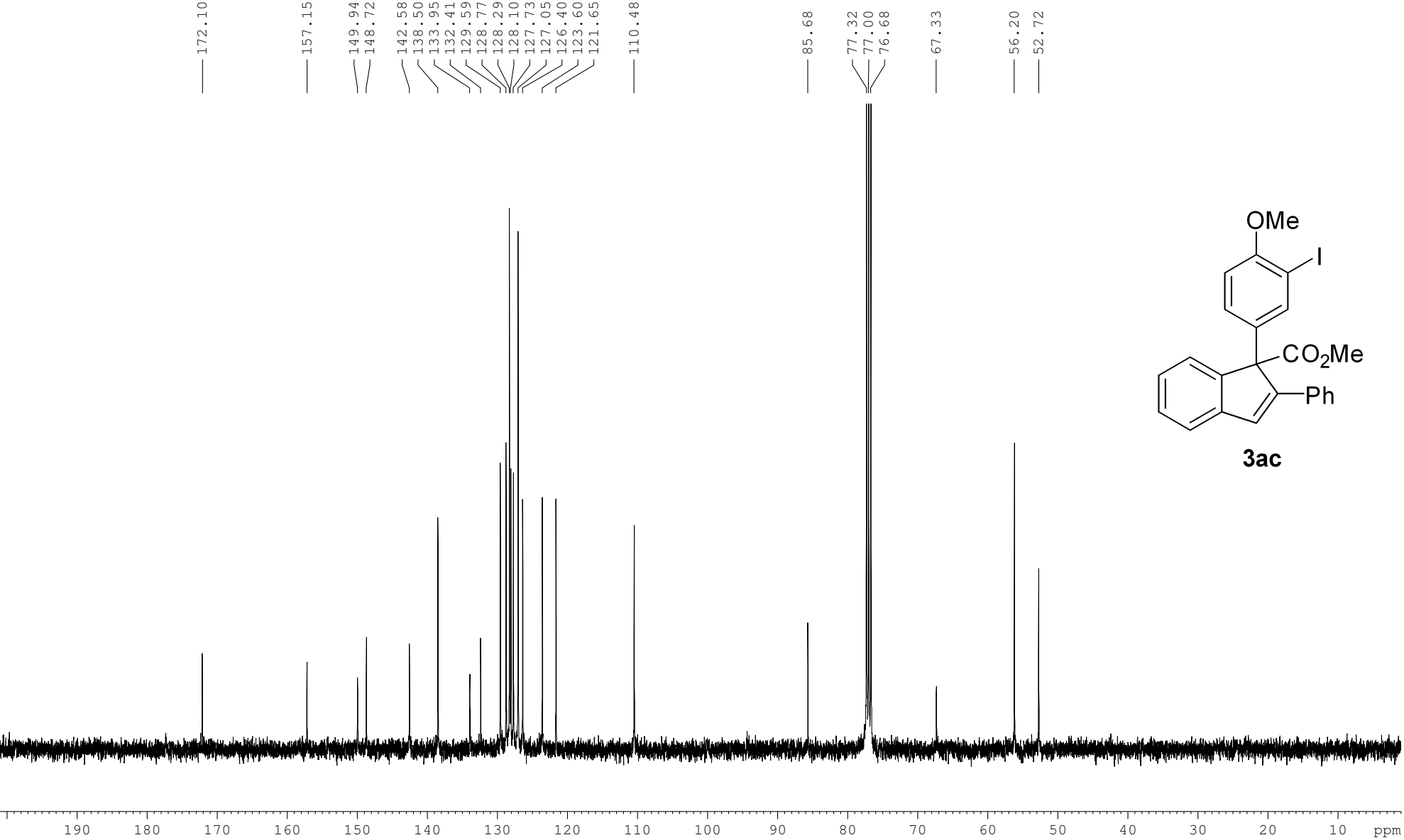
7.855
7.849
7.404
7.401
7.383
7.369
7.340
7.321
7.307
7.288
7.283
7.266
7.262
7.244
7.231
7.214
7.181
7.162
6.622
6.600

3.772
3.578

0.000



mbw-1-92-1

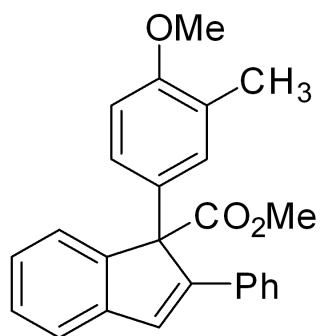


mb-5-138-1
mb-5-138-1

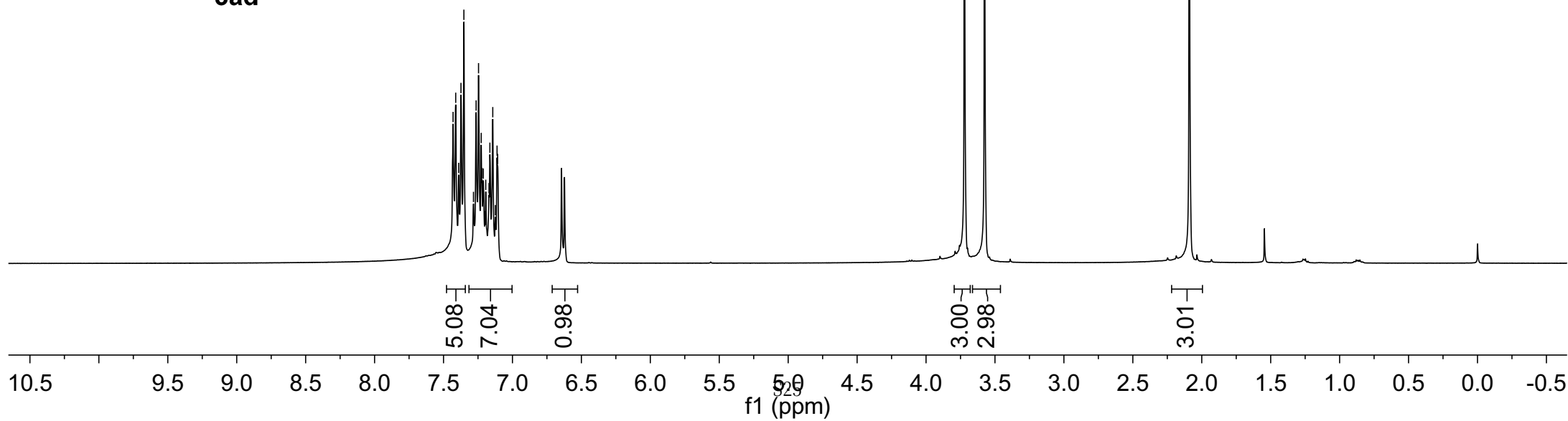
7.431
7.412
7.390
7.373
7.352
7.283
7.264
7.246
7.227
7.221
7.211
7.200
7.194
7.170
7.164
7.143
7.124
7.112
7.107

~3.722
~3.576

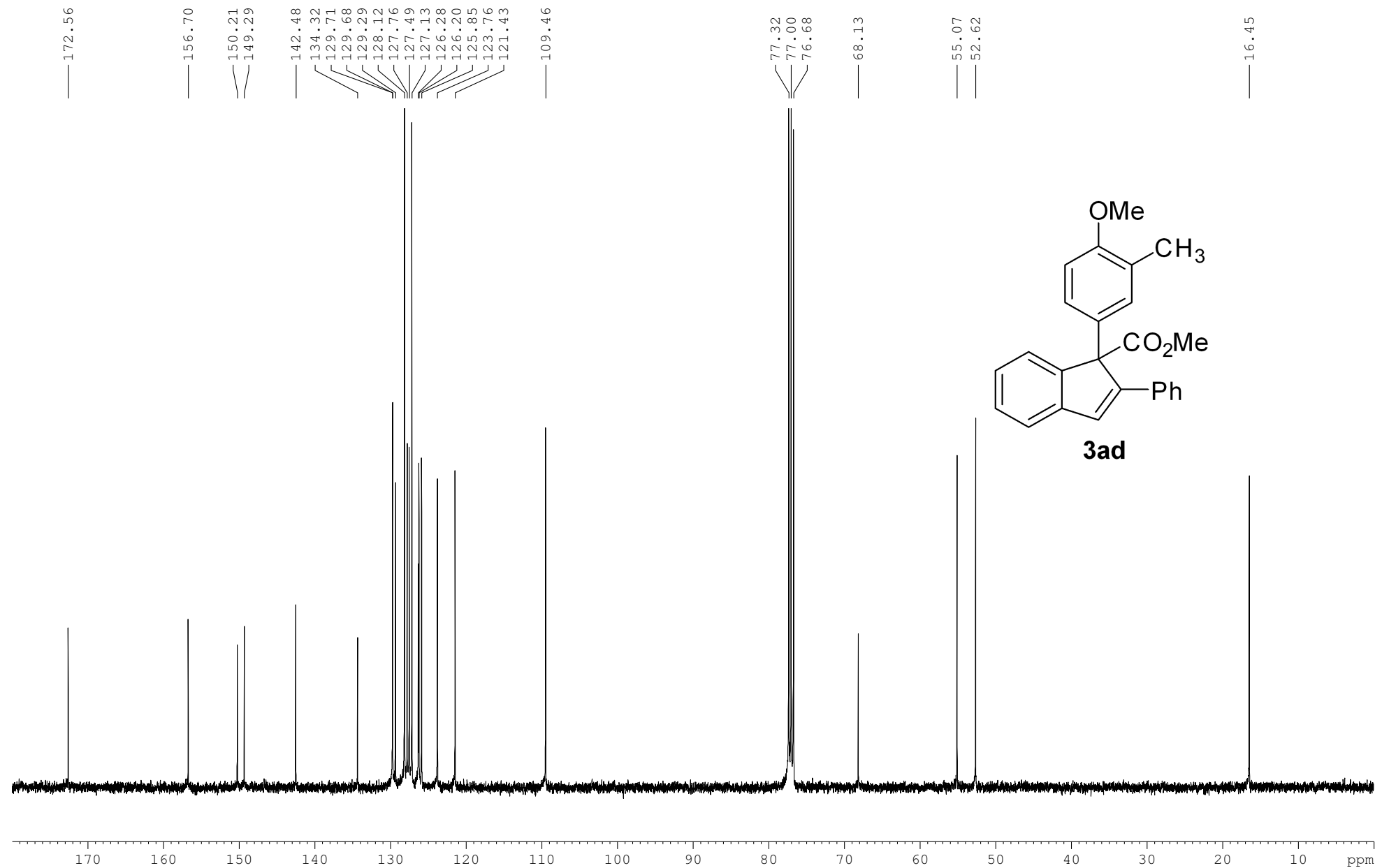
—2.090



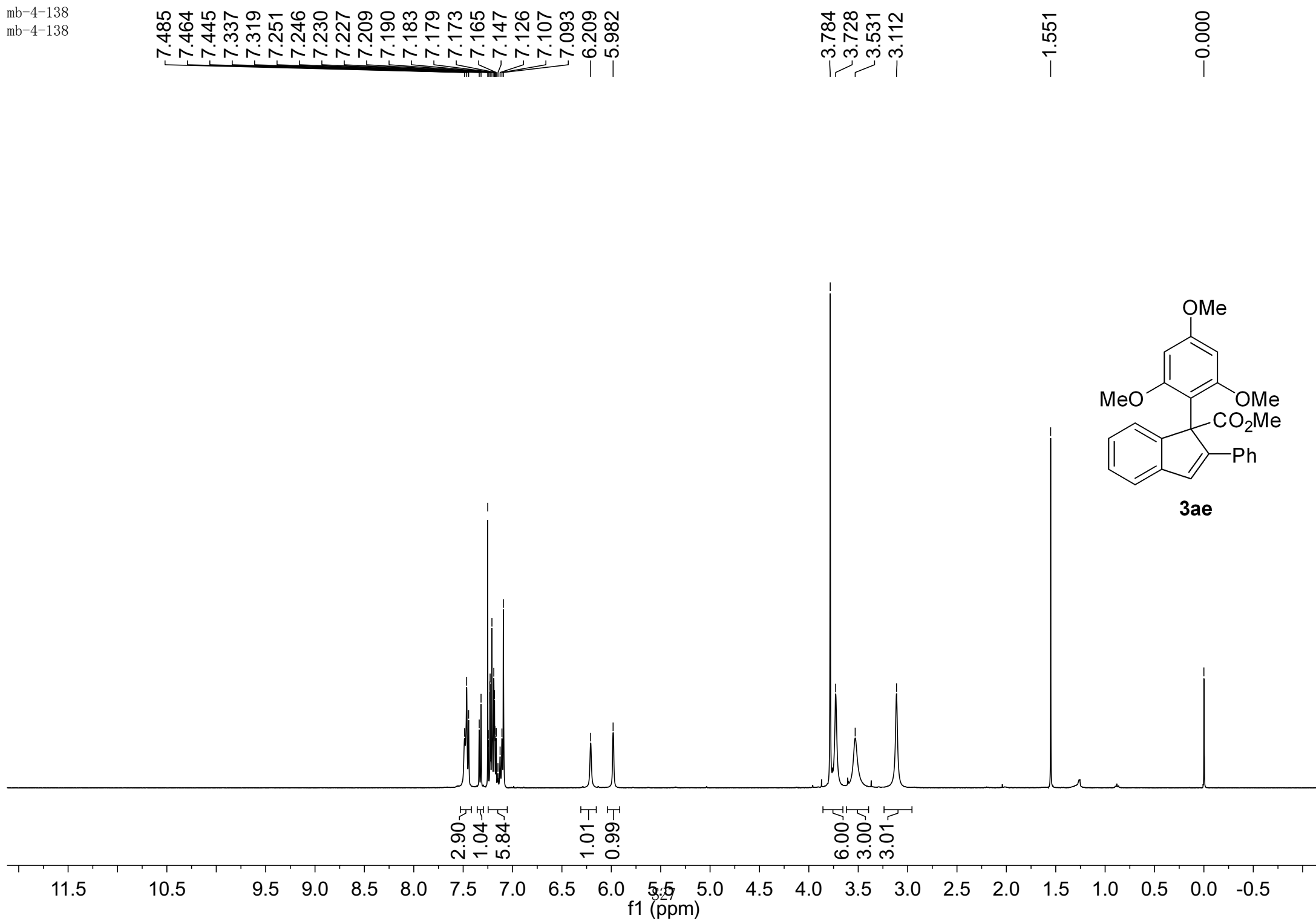
3ad



mb-5-138-1

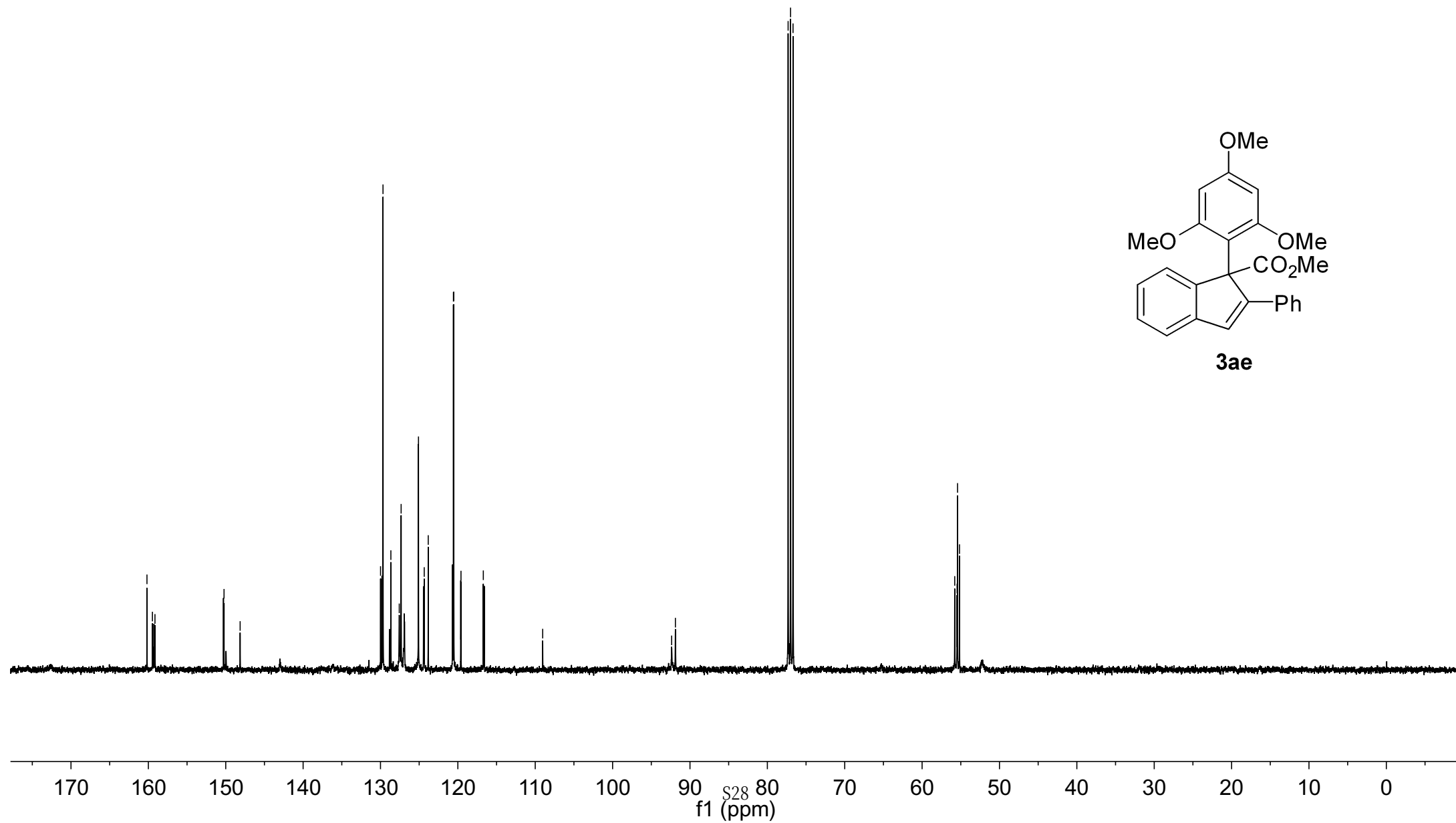


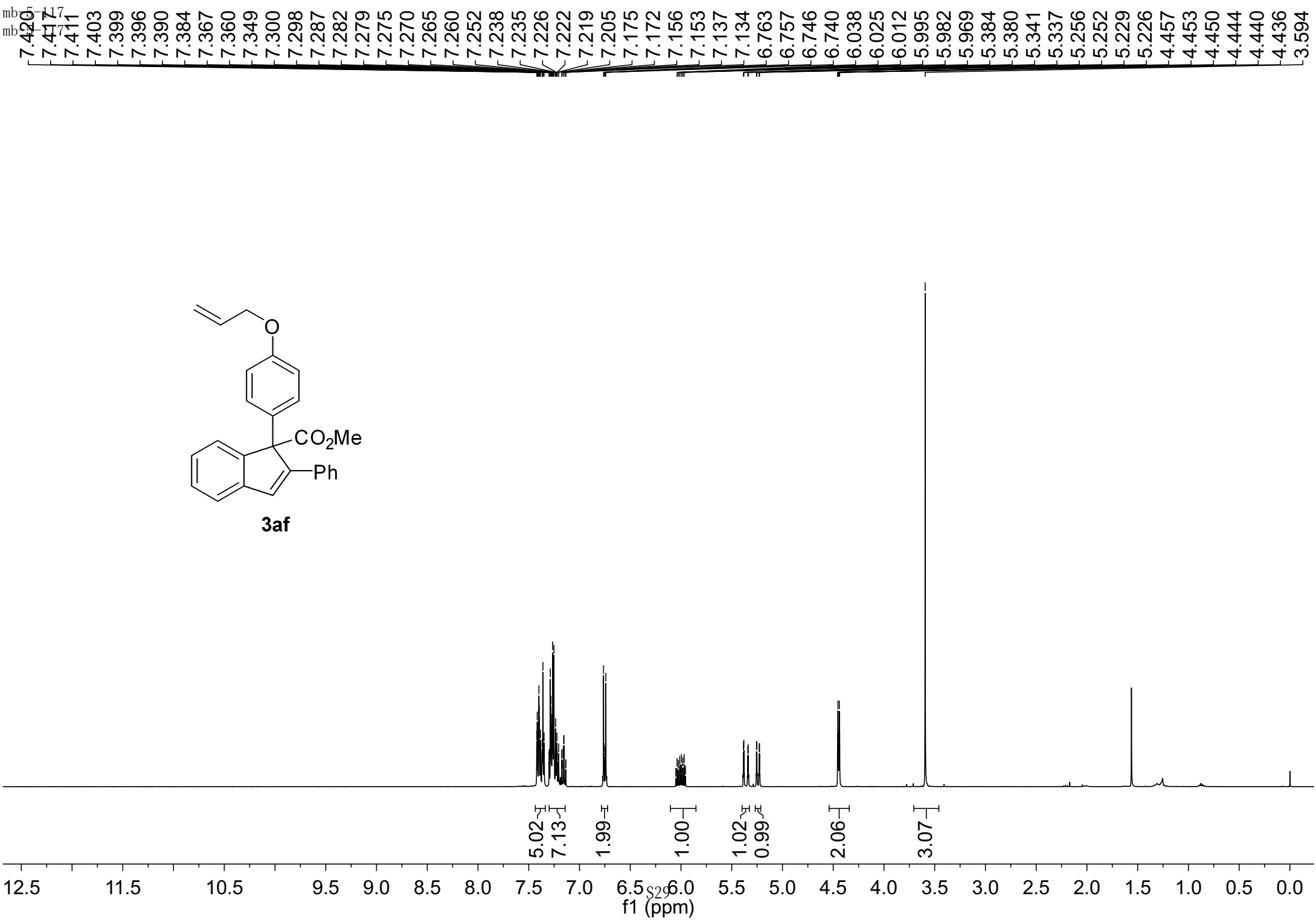
mb-4-138
mb-4-138



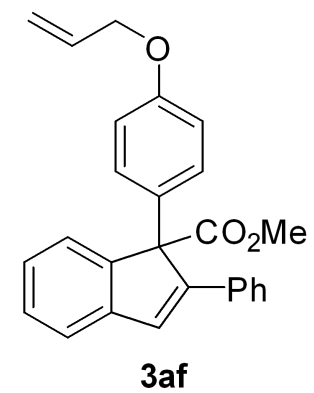
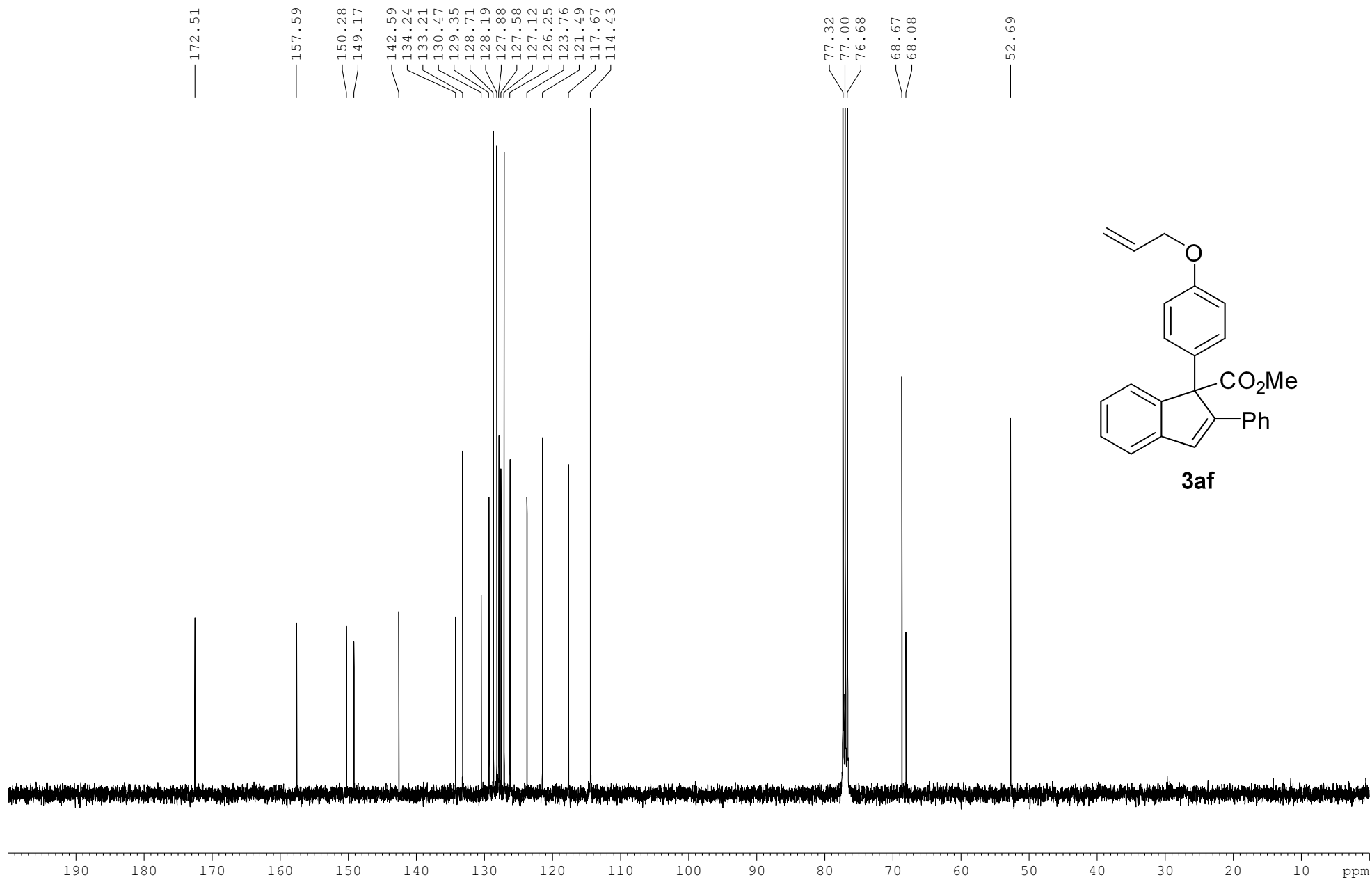
mb-4-136
mb-4-136

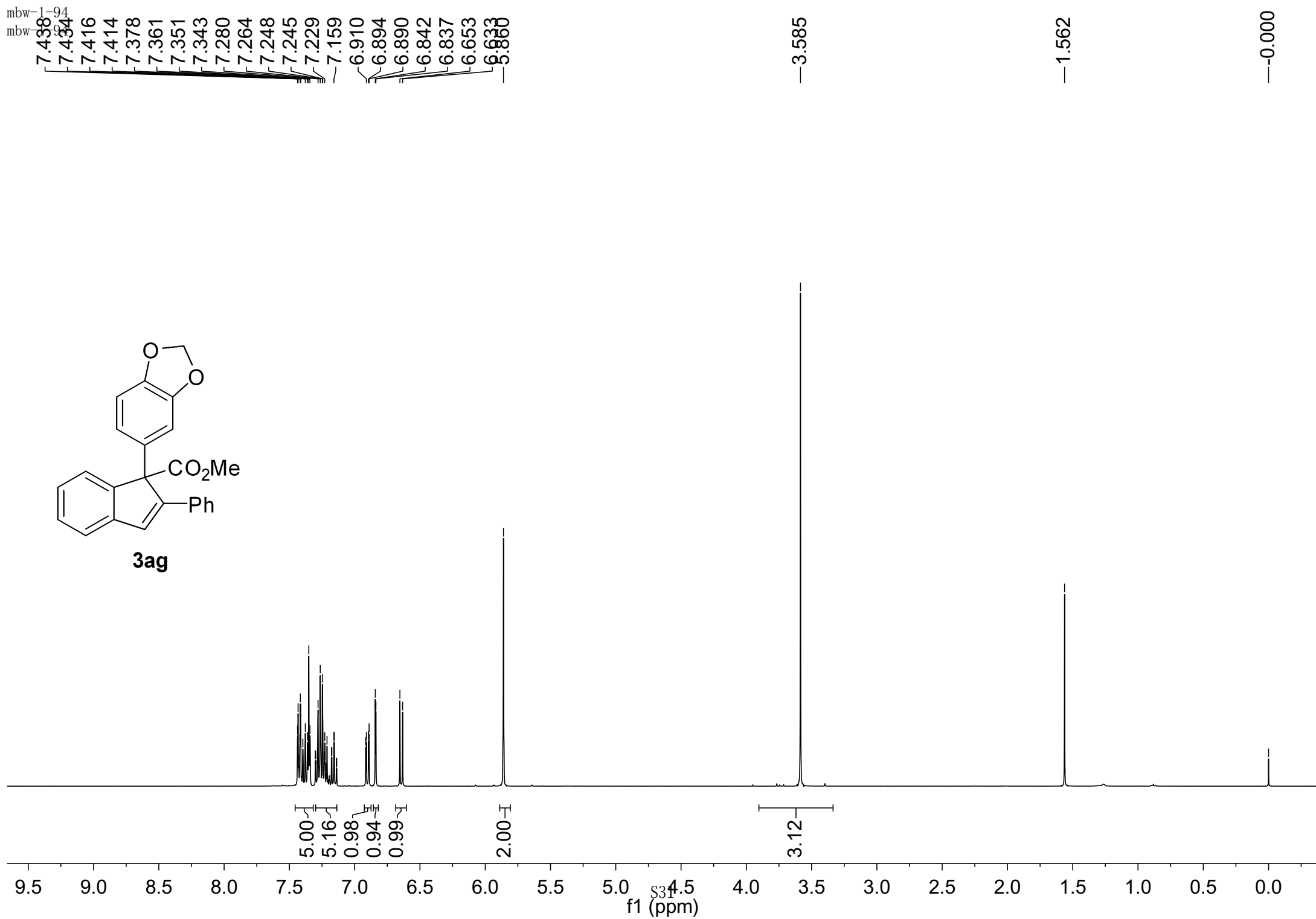
160.181
159.476
159.147
150.220
148.139
129.990
129.674
128.658
127.332
125.089
124.346
123.827
120.573
120.528
119.595
116.710
109.636
92.370
91.885
77.317
77.000
76.682
55.784
55.480
55.418
55.165



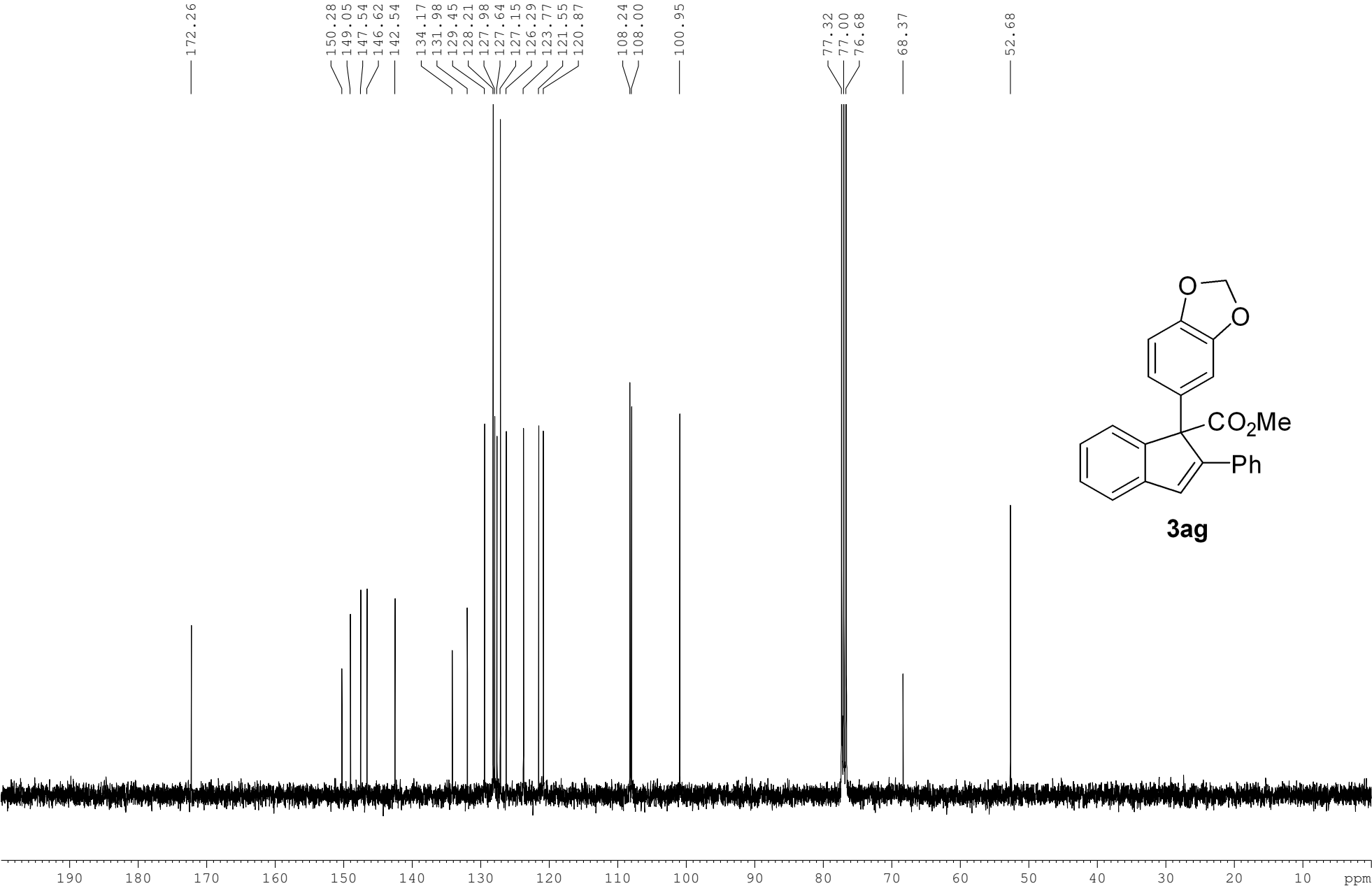


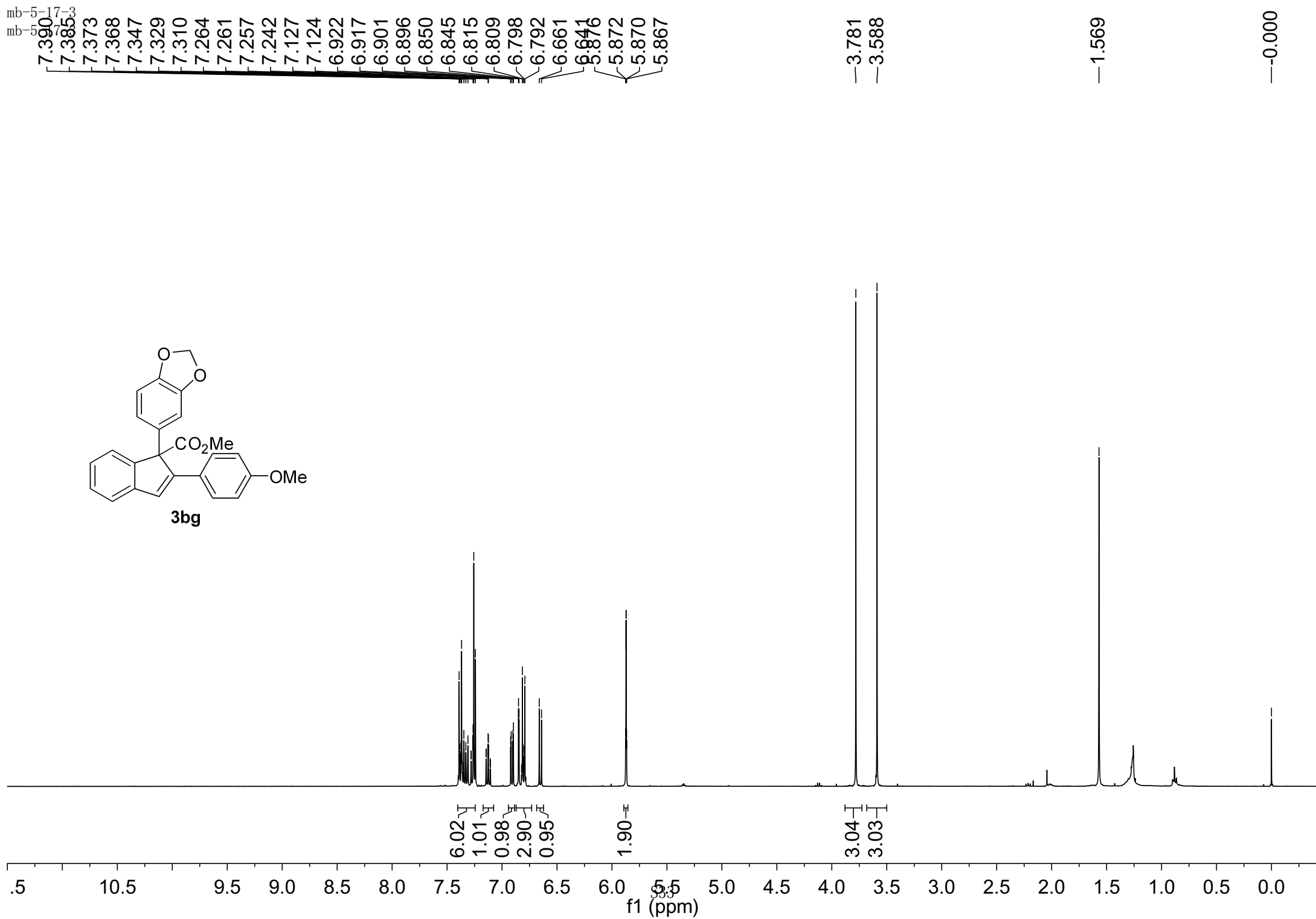
mb-5-117





mbw-1-94





mb-5-17-3
mb-5-17-3

—172.411

—159.193

149.866

148.887

147.547

146.593

142.823

132.233

128.482

127.914

127.651

126.834

125.855

123.661

121.214

120.855

113.698

108.264

108.003

100.953

77.318

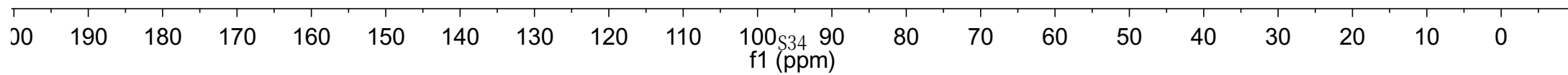
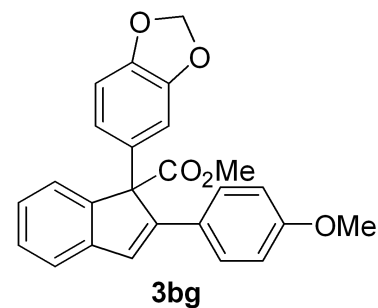
77.000

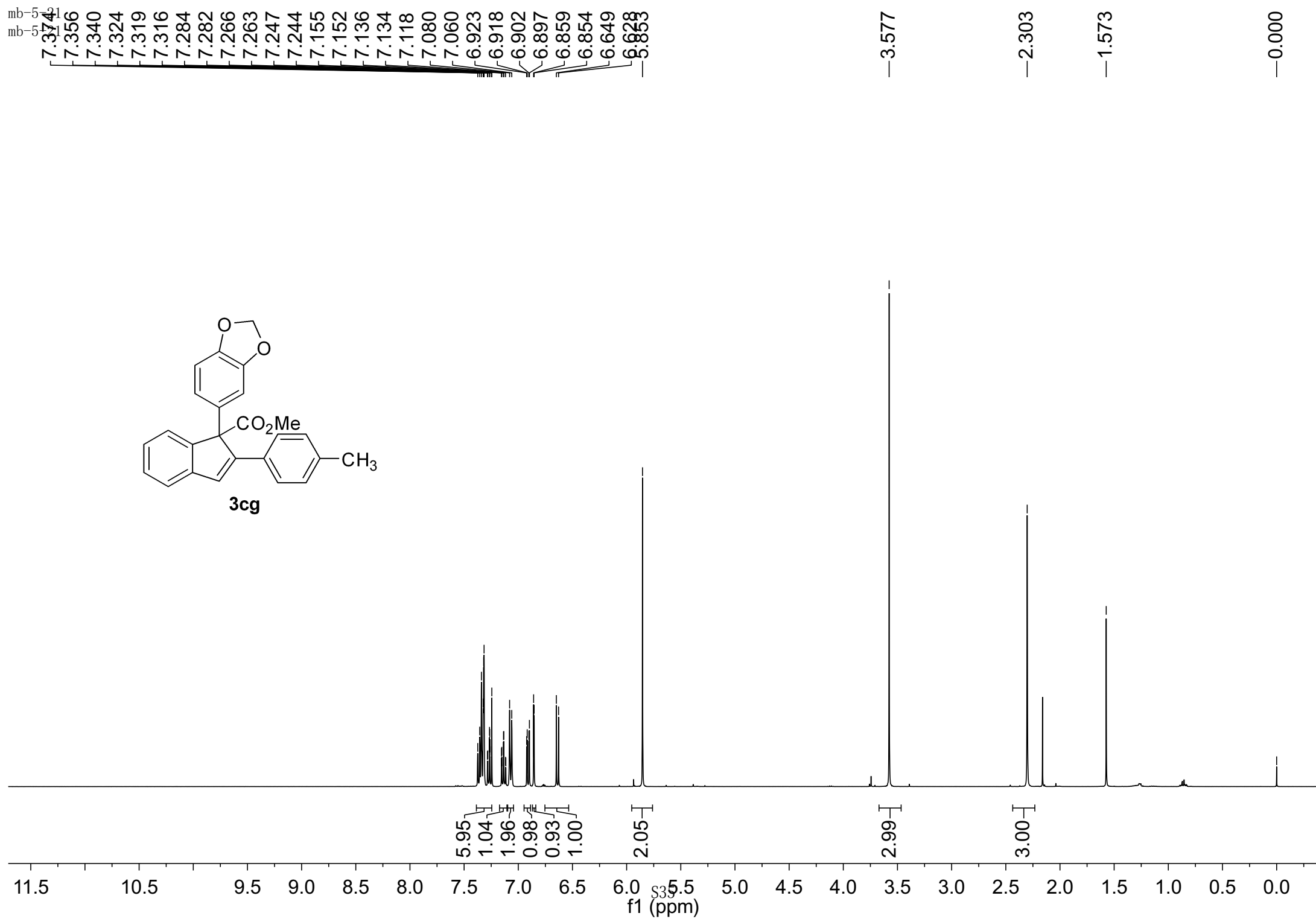
76.683

—68.337

~55.191

~52.678





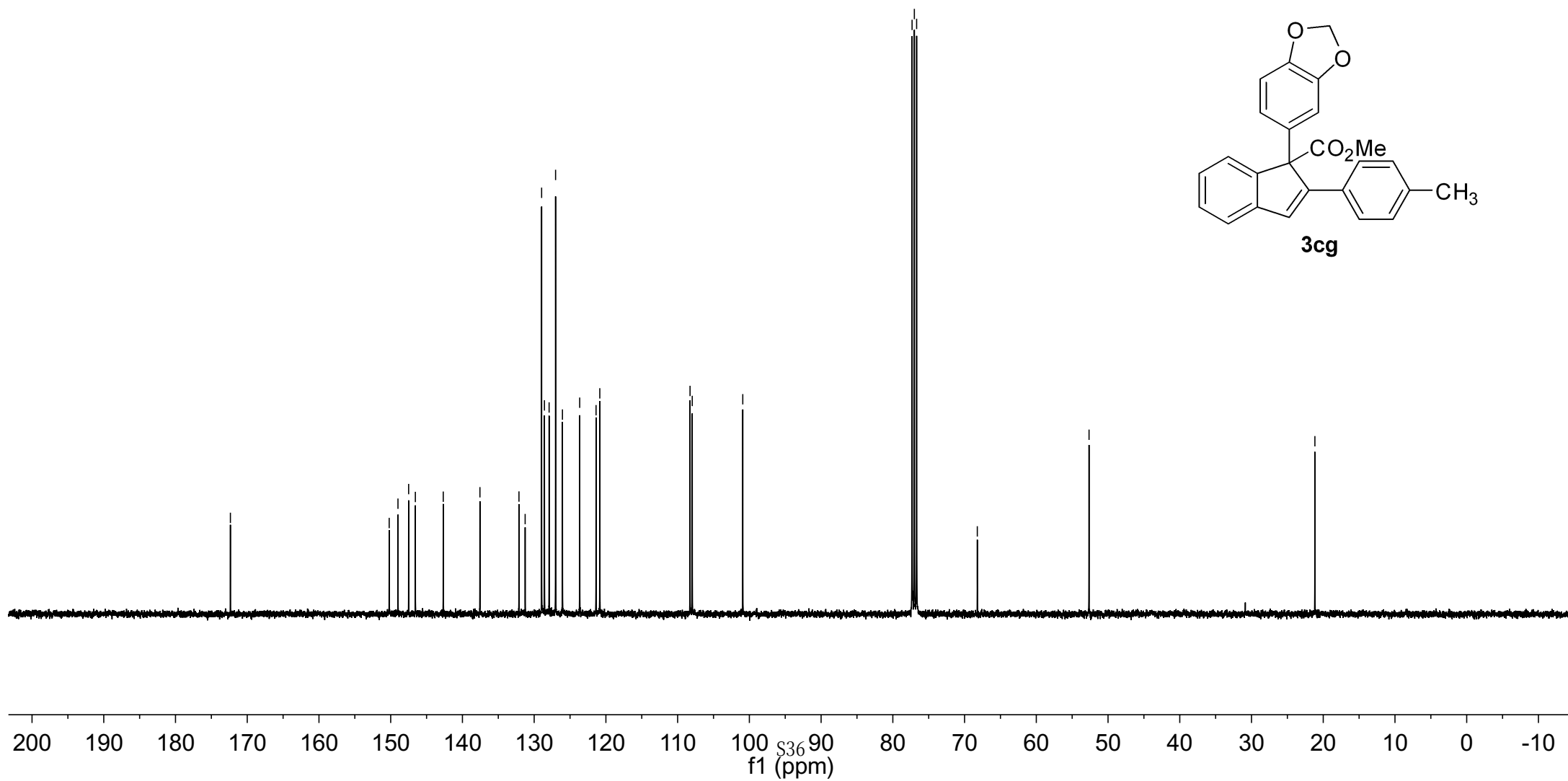
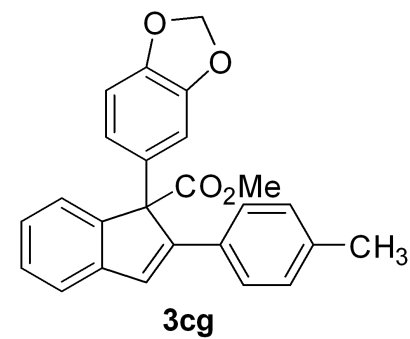
mb-5-21
mb-5-21 C

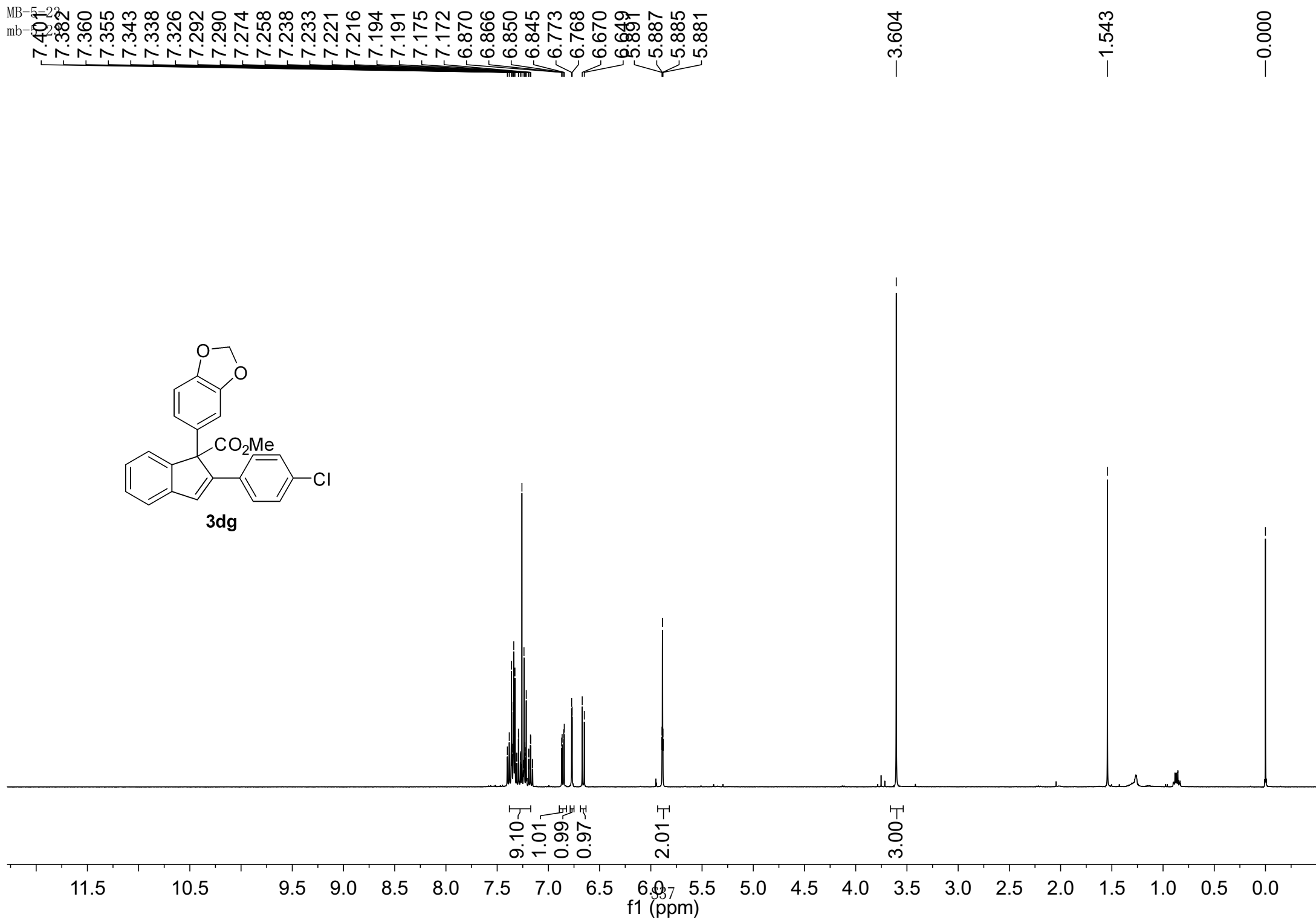
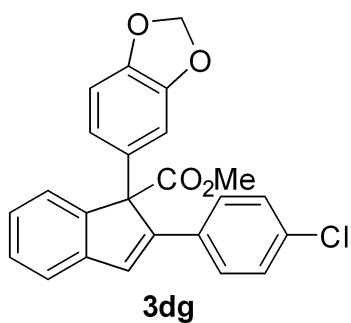
172.346
150.210
148.987
147.499
146.563
142.674
137.561
132.120
131.257
128.961
128.567
127.916
127.015
126.057
123.653
121.373
120.851
108.270
107.963
100.918

77.318
77.000
76.682
68.224

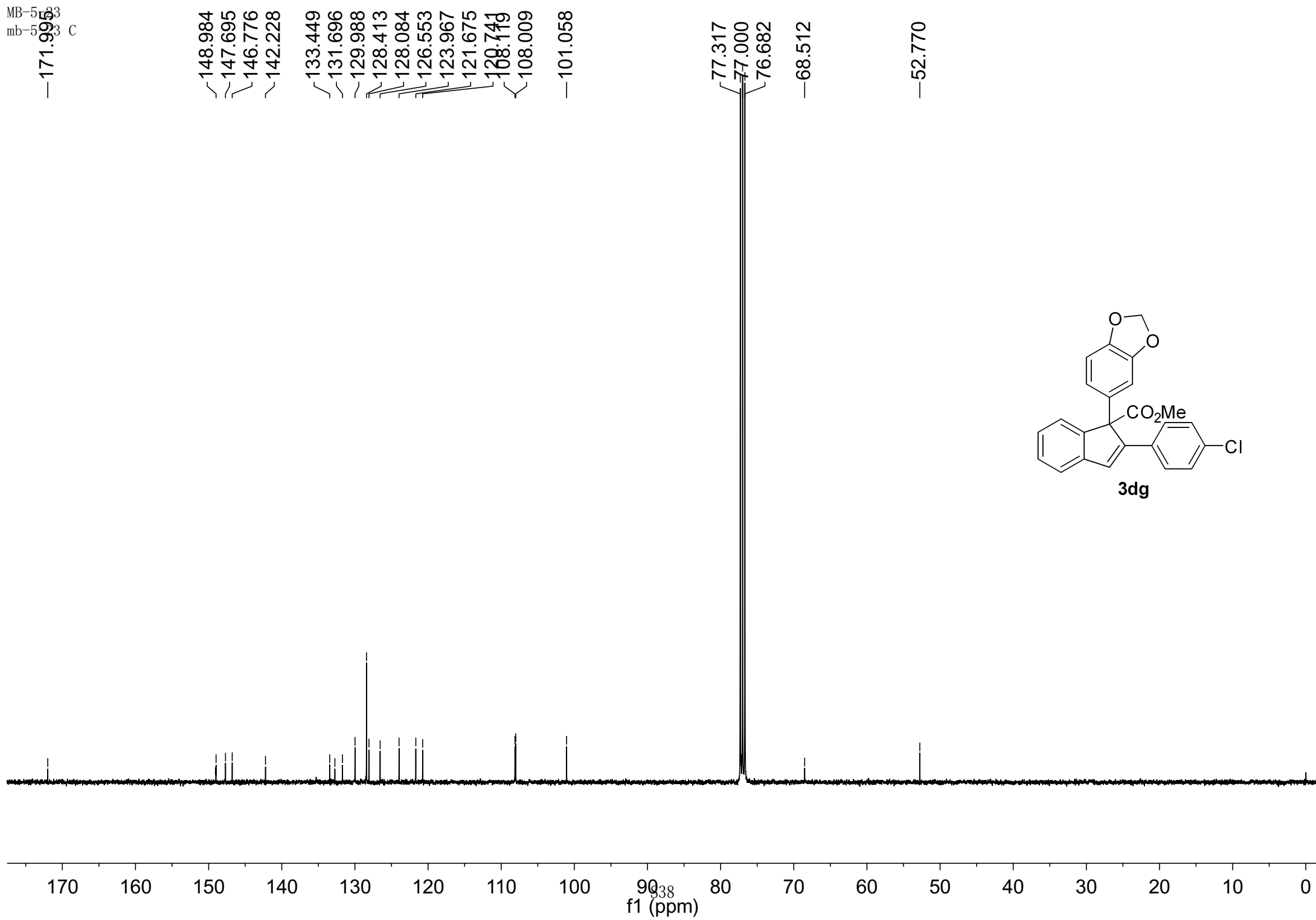
52.648

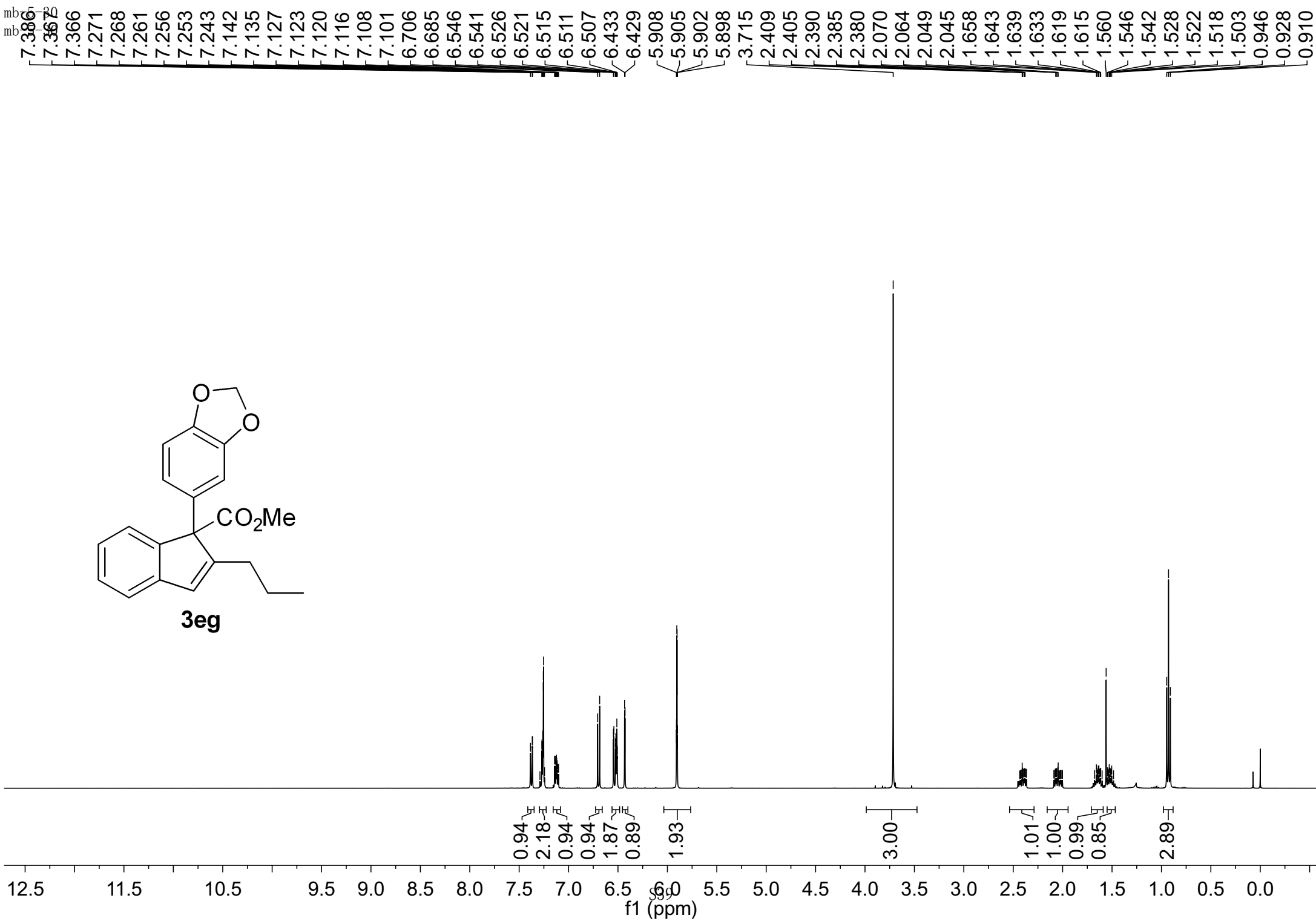
21.163



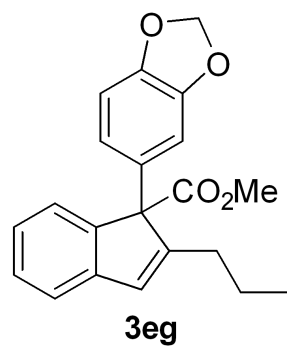


MB-5133
mb-5133 C





mb-5-30
mb-5-30



—172.435

154.513

147.611

146.659

146.582

144.301

132.981

128.011

126.387

125.094

124.804

120.567

120.441

108.107

107.785

101.032

77.317

77.000

76.682

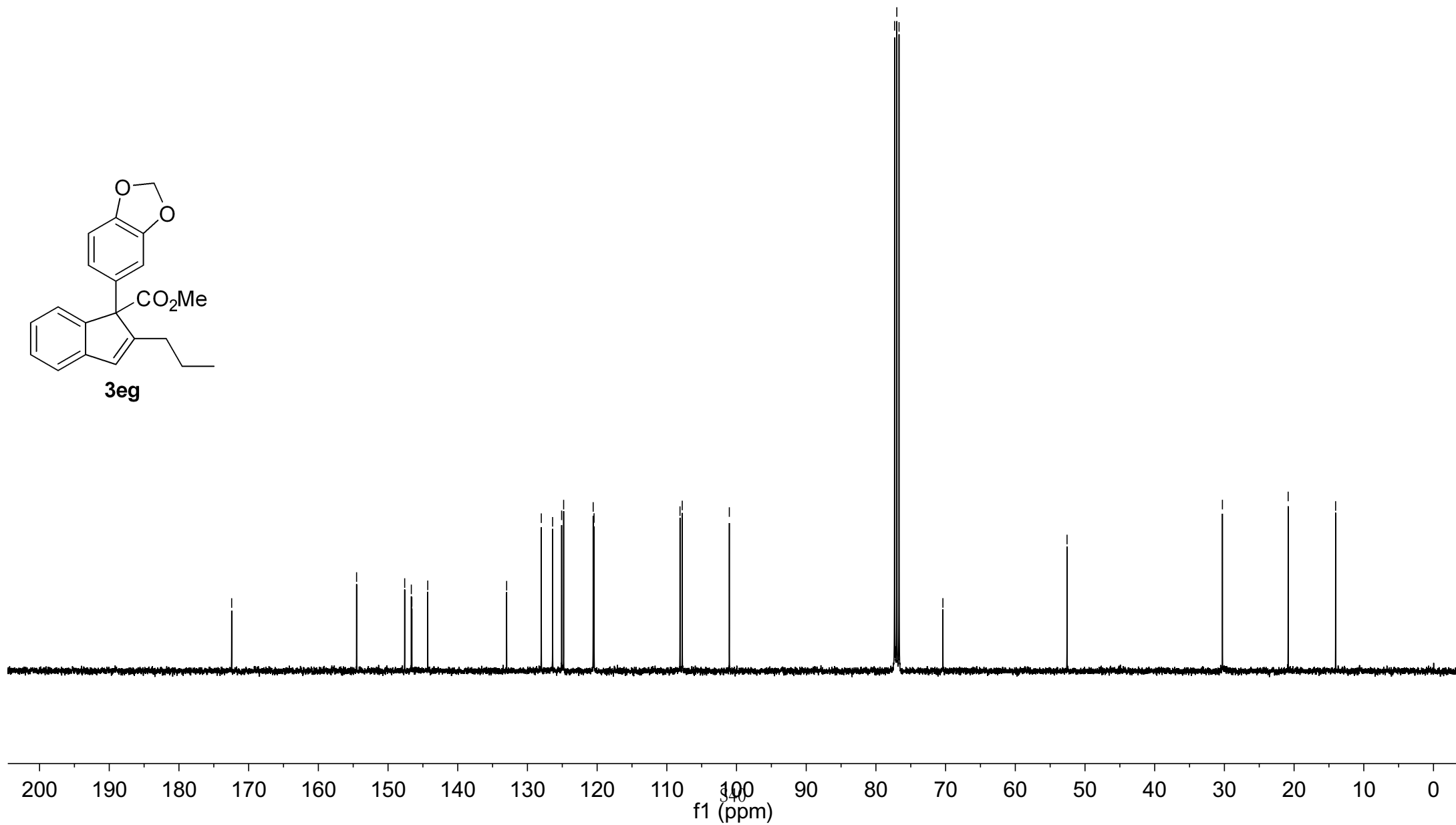
70.389

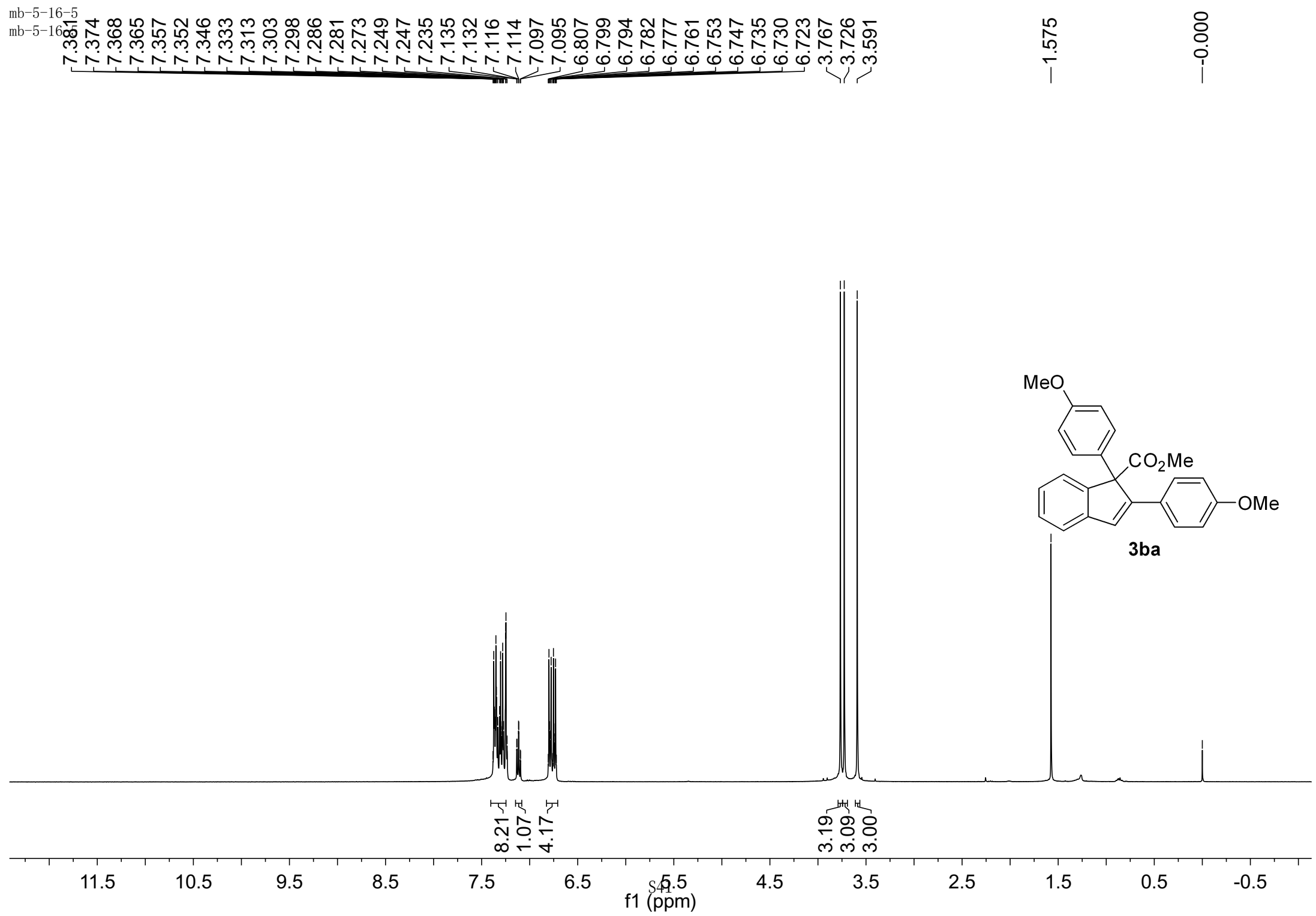
—52.576

—30.286

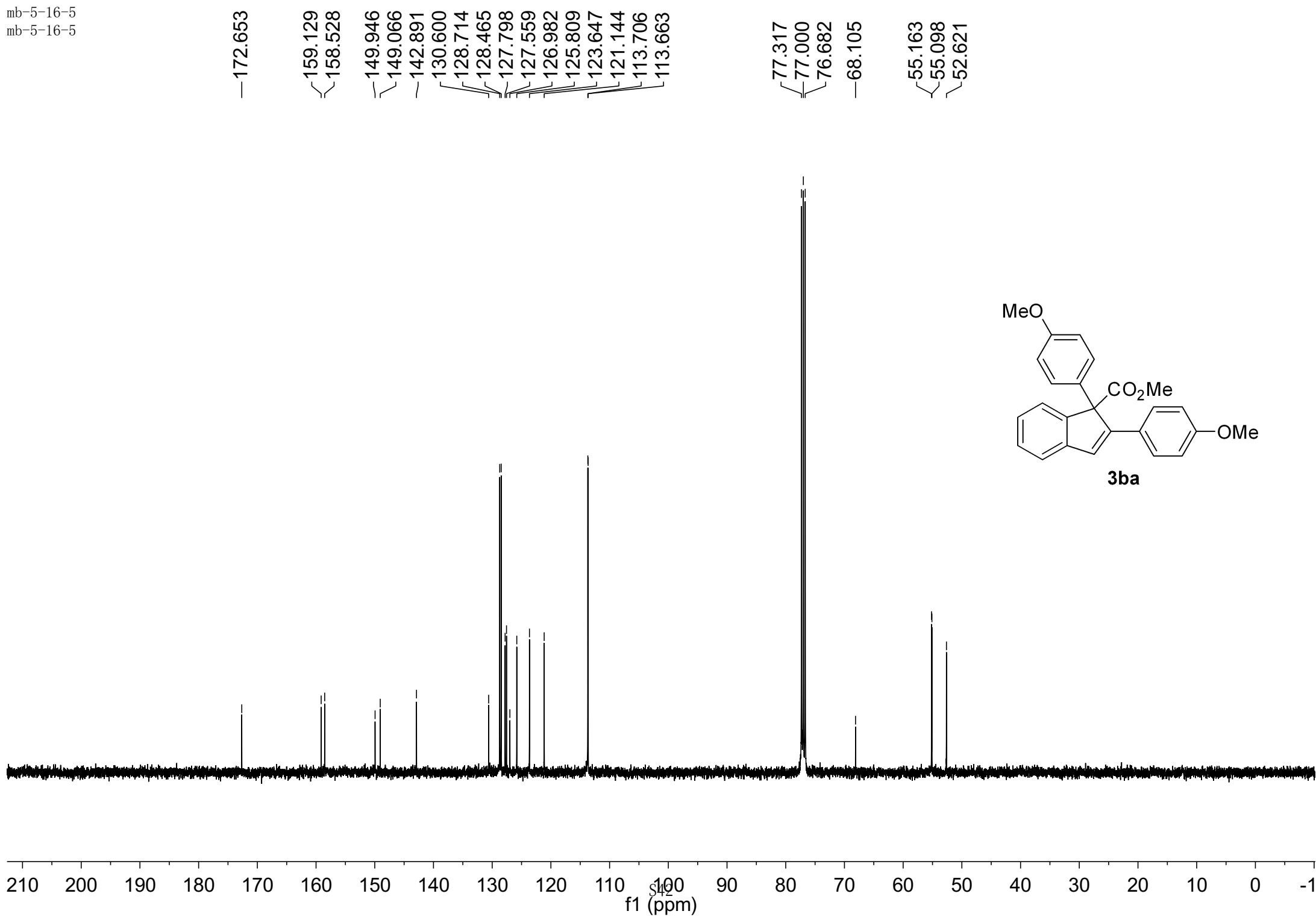
—20.851

—14.042

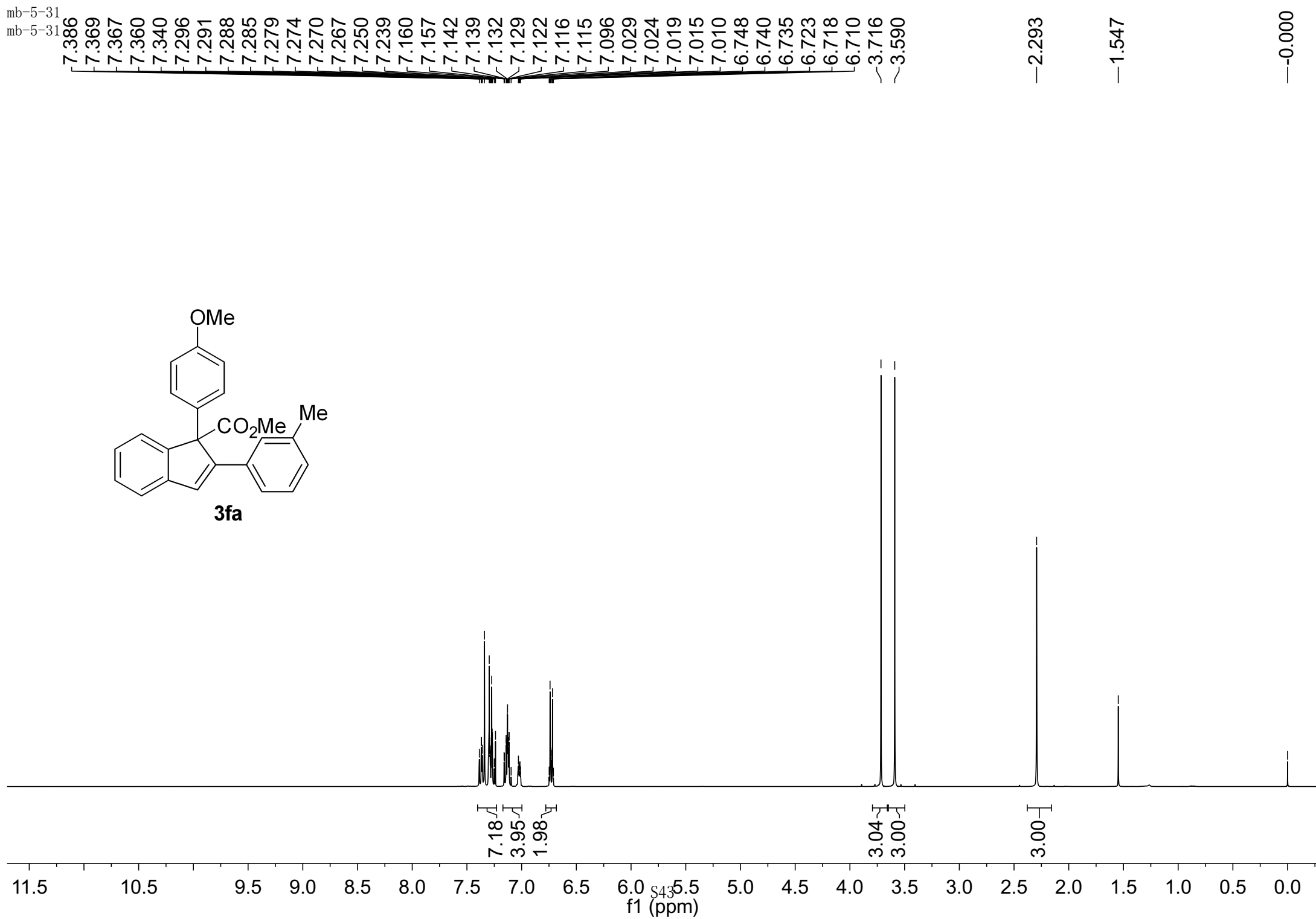
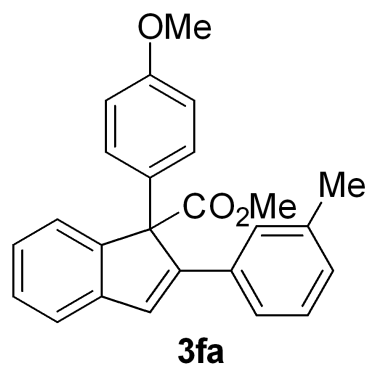




mb-5-16-5
mb-5-16-5



mb-5-31
mb-5-31



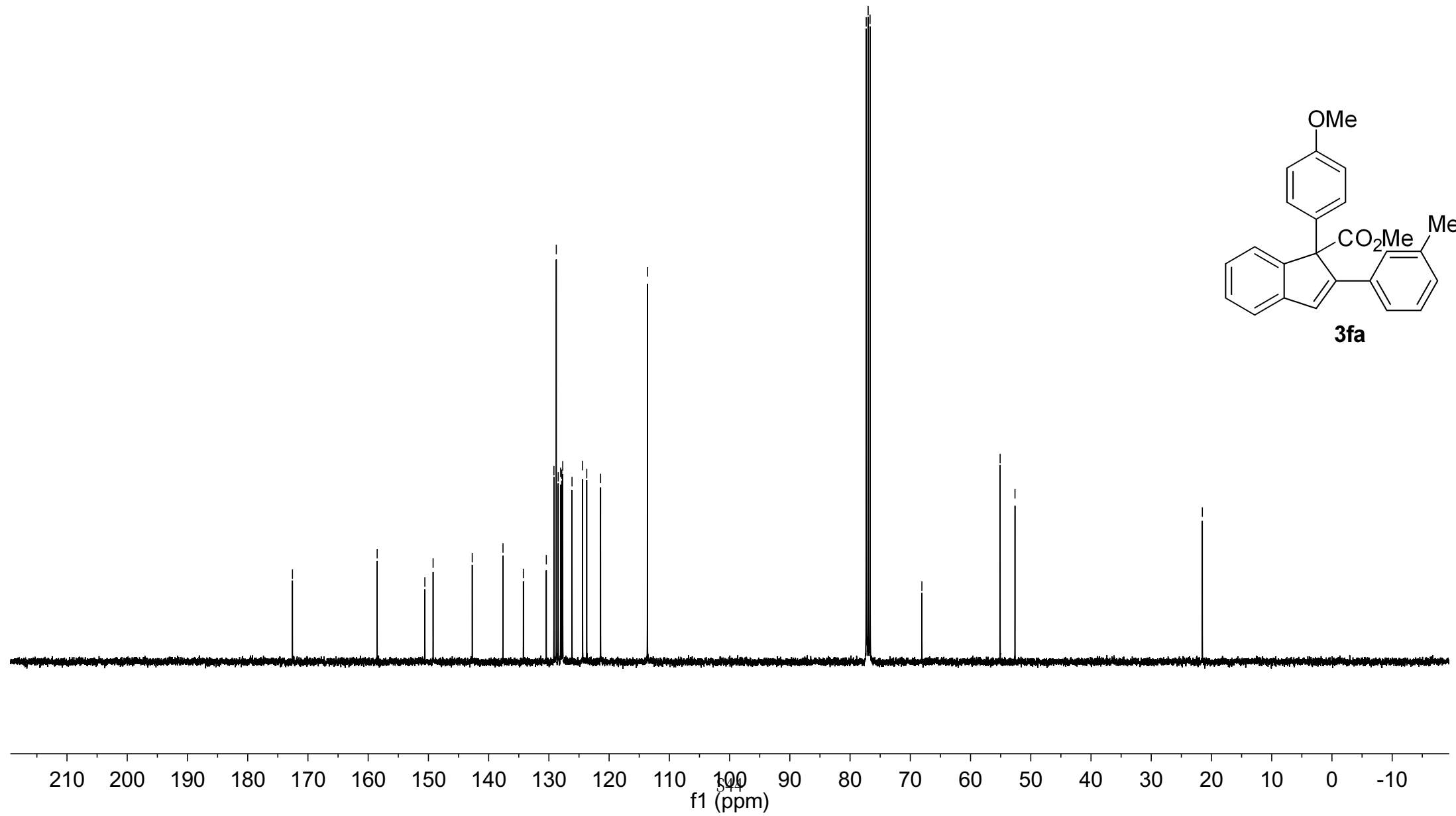
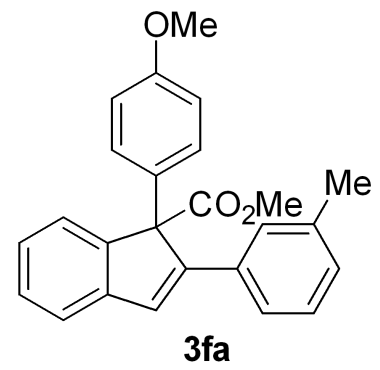
mb-5-31
mb-5-31 C

172.571
158.508
150.592
149.221
142.724
137.619
134.235
130.429
129.162
128.772
128.448
128.055
127.860
127.697
126.159
124.402
123.712
121.416
113.643

77.317
77.000
76.682
68.076

55.086
52.620

21.516

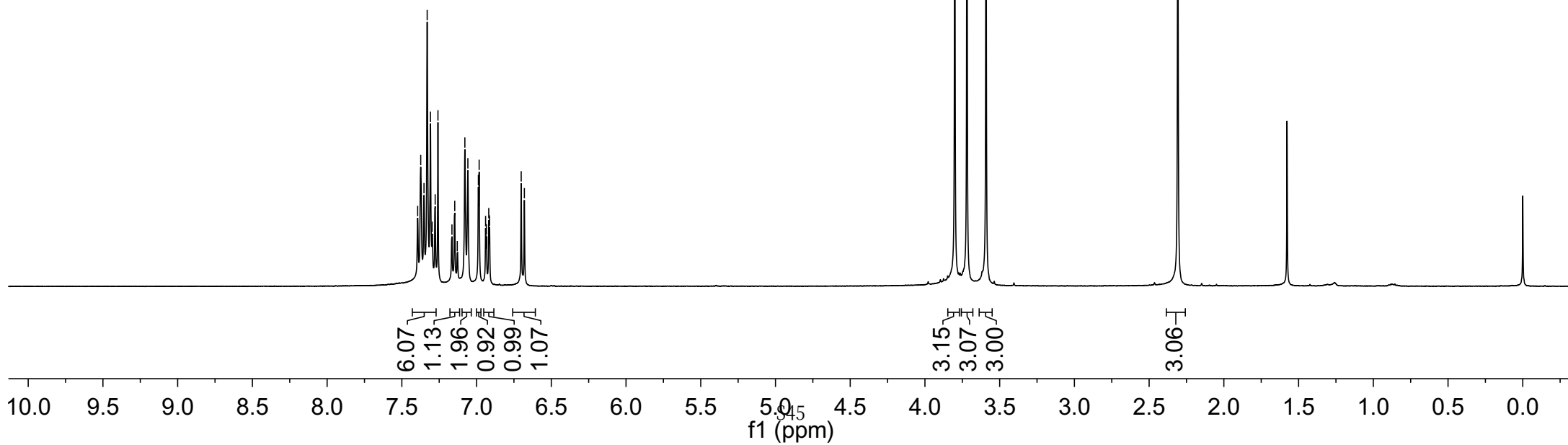
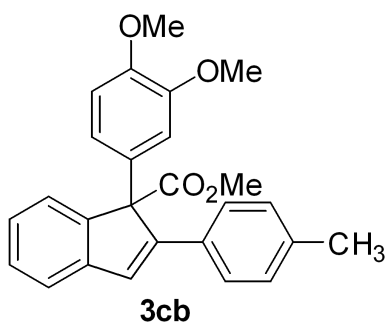


mb-5-139-1
mb-5-139-1

7.394
7.374
7.352
7.330
7.308
7.297
7.277
7.258
7.165
7.146
7.129
7.078
7.058
6.988
6.983
6.940
6.934
6.918
6.913
6.701
6.680

3.801
3.719
3.591

2.308



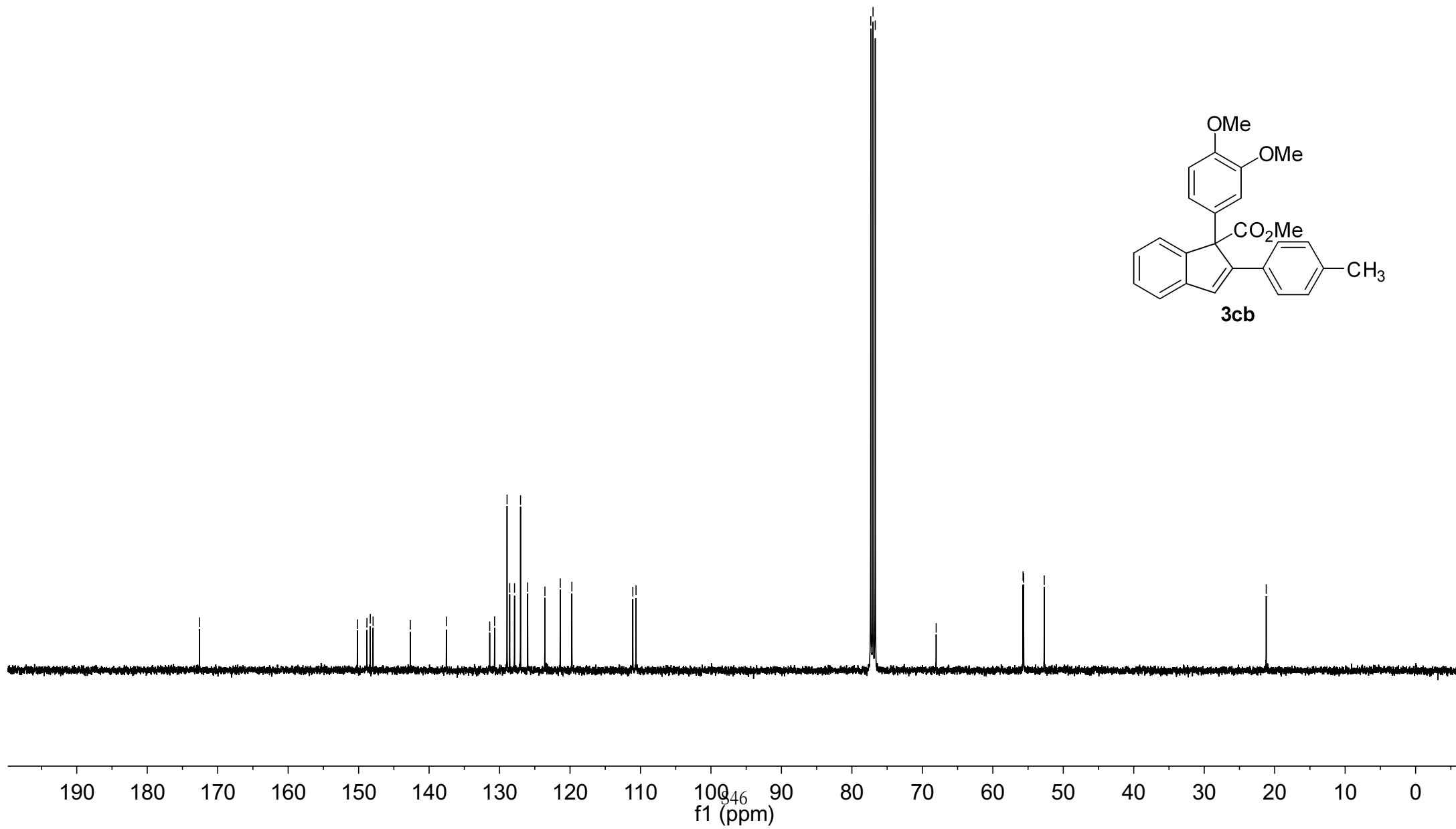
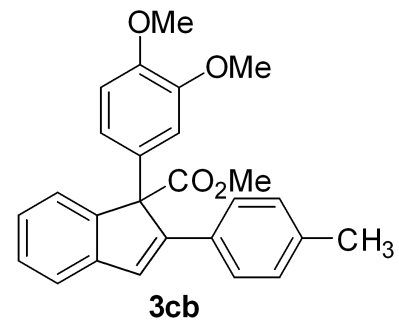
mb-5-139-1
mb-5-139-1

172.583
150.172
148.821
148.344
147.948
142.661
137.549
131.398
130.705
128.936
128.561
127.877
127.023
126.010
123.563
121.386
119.738
111.090
110.634

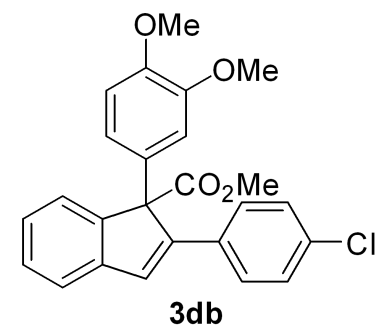
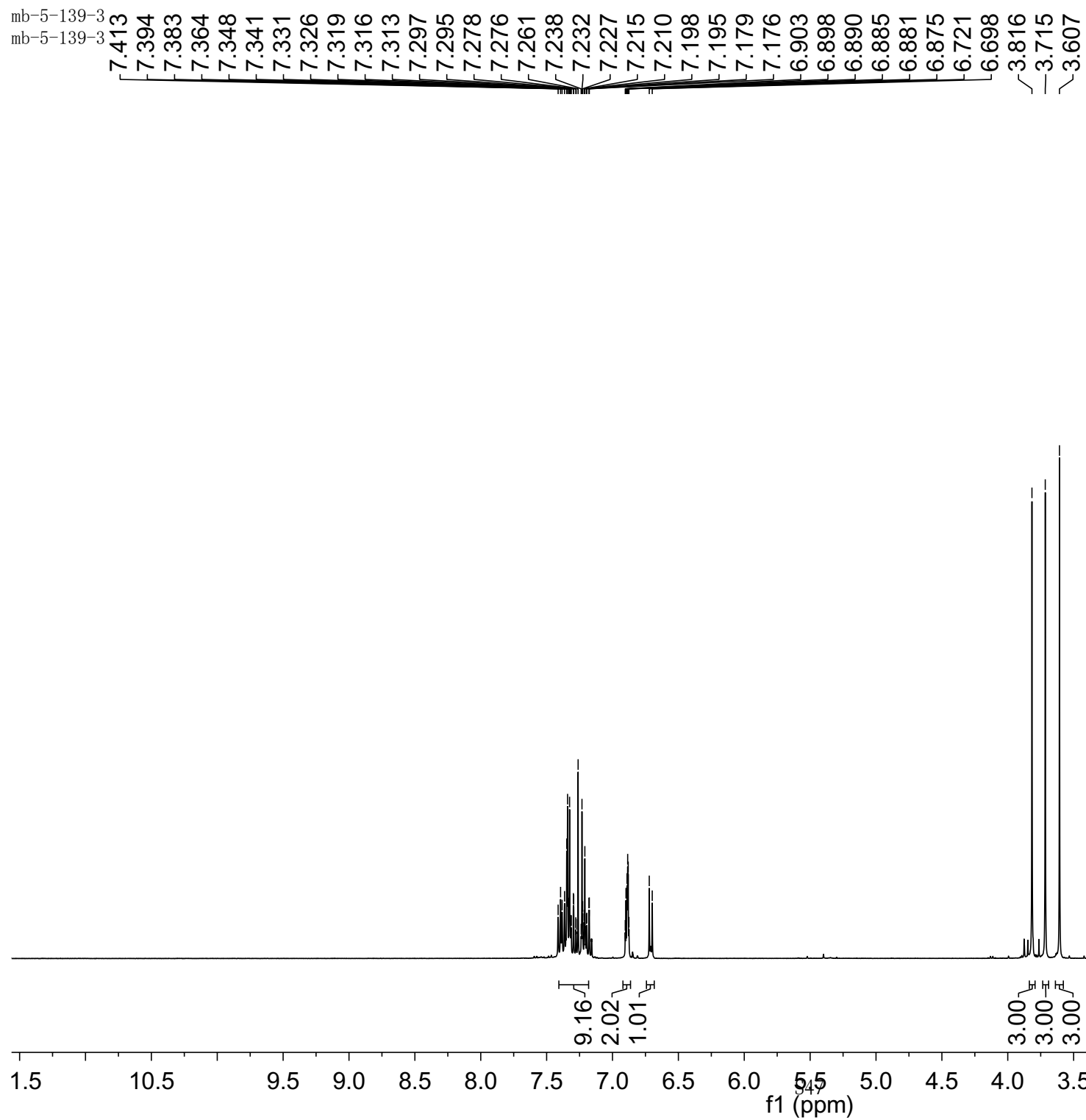
77.317
77.000
76.682
68.039

55.712
55.627
52.706

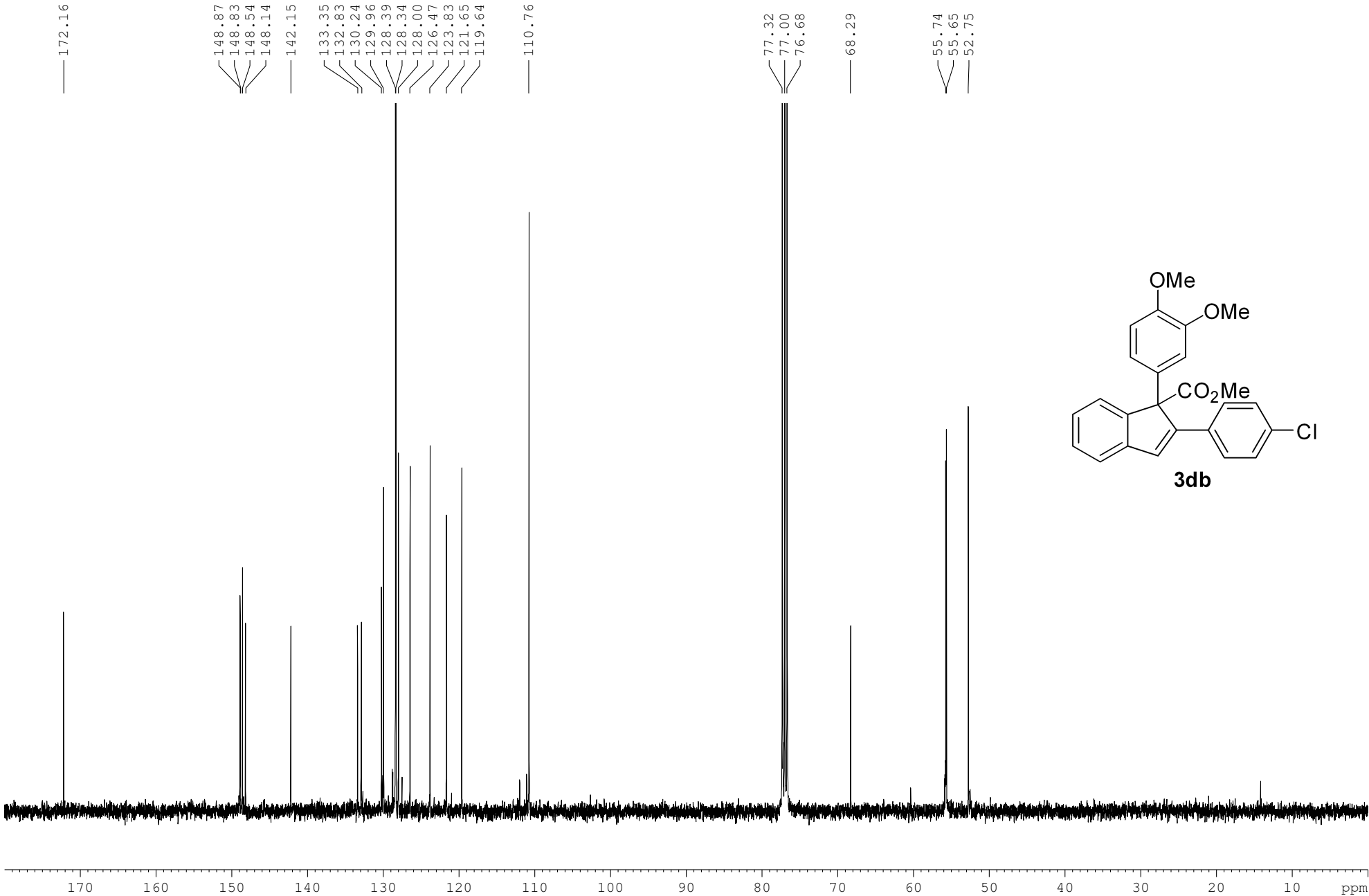
21.200

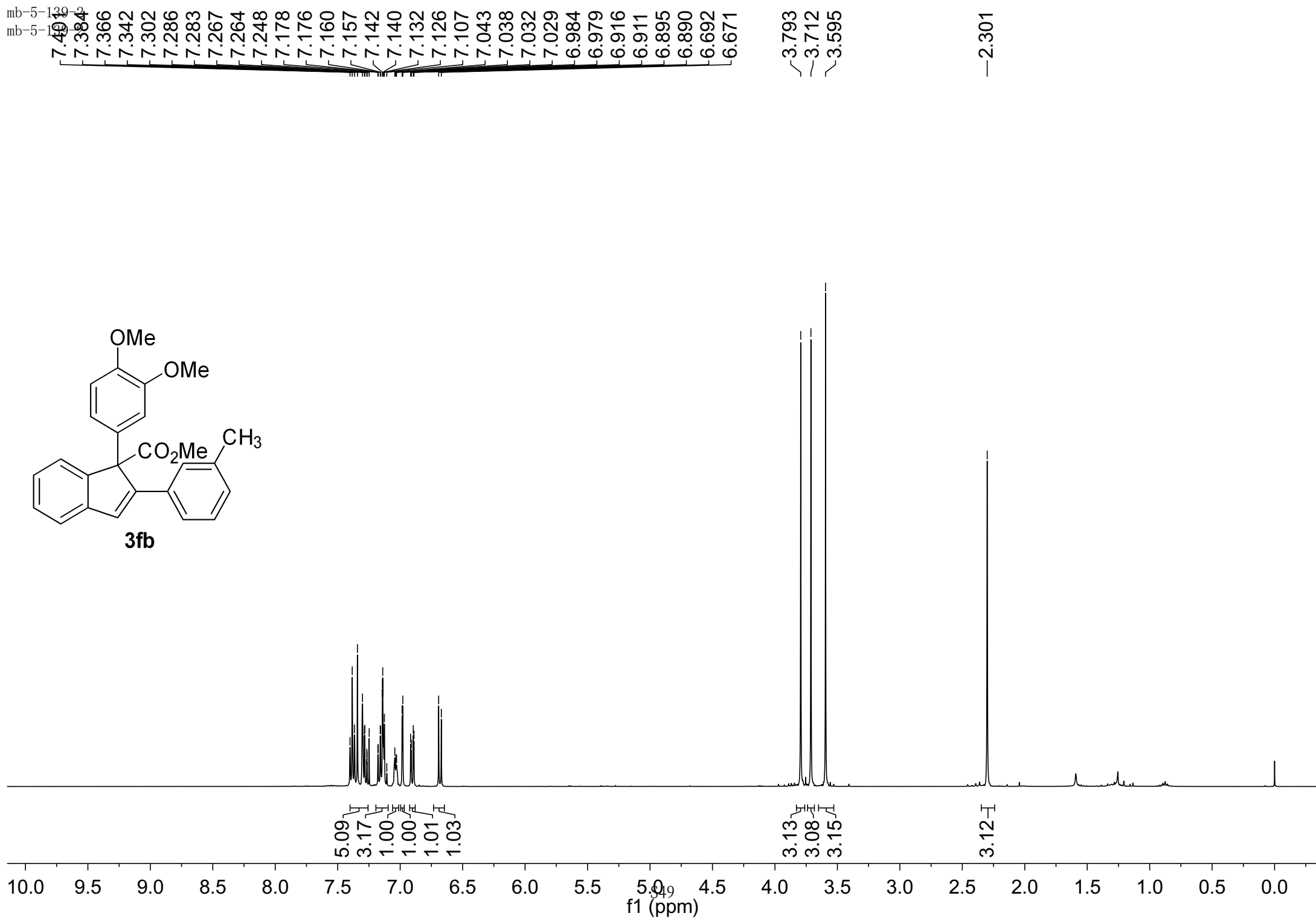


mb-5-139-3
mb-5-139-3

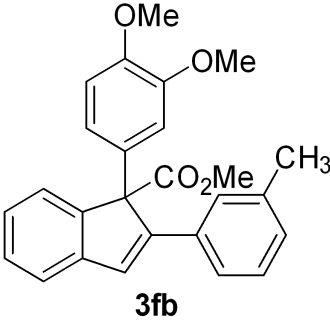
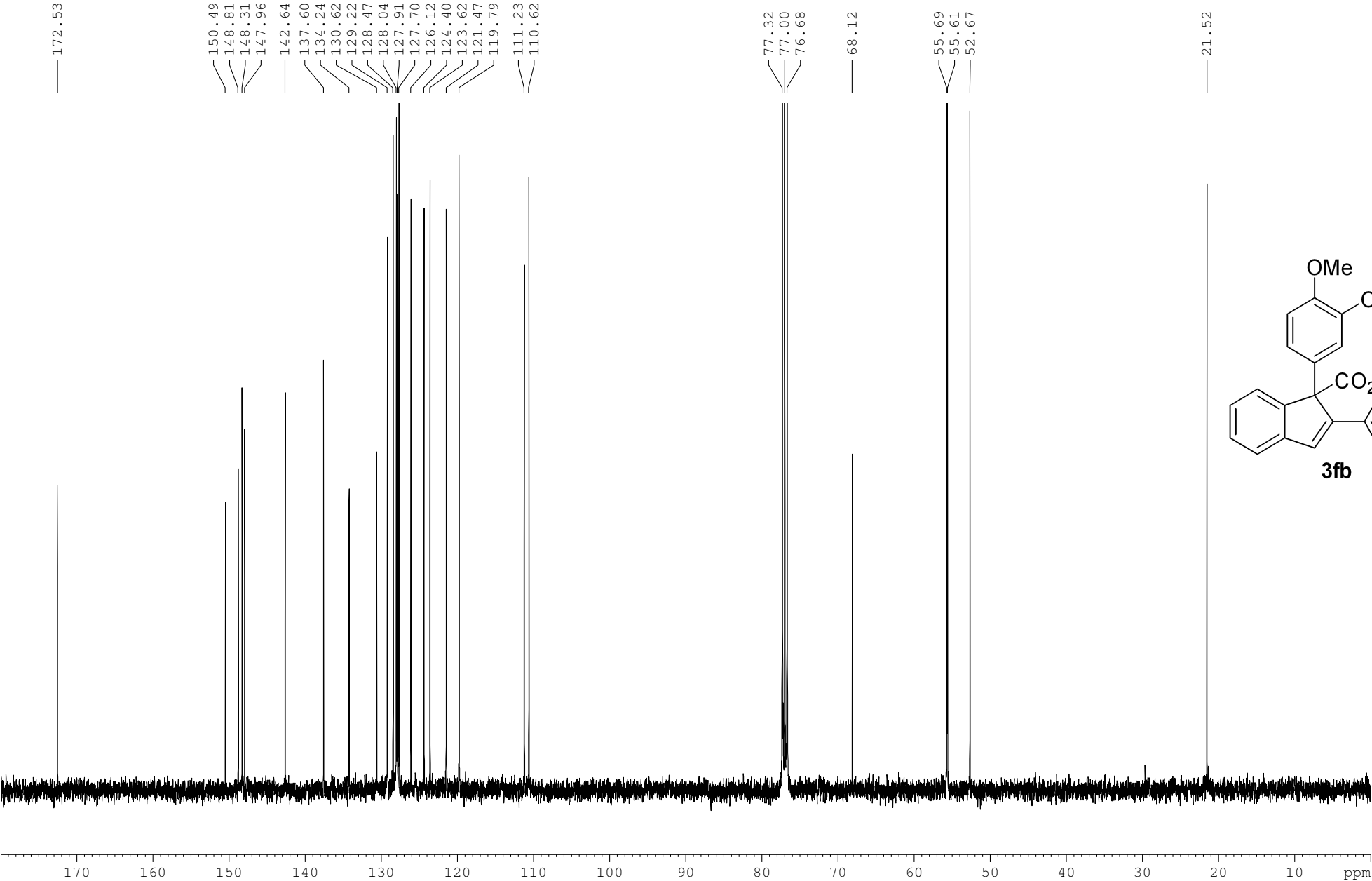


mb-5-139-3

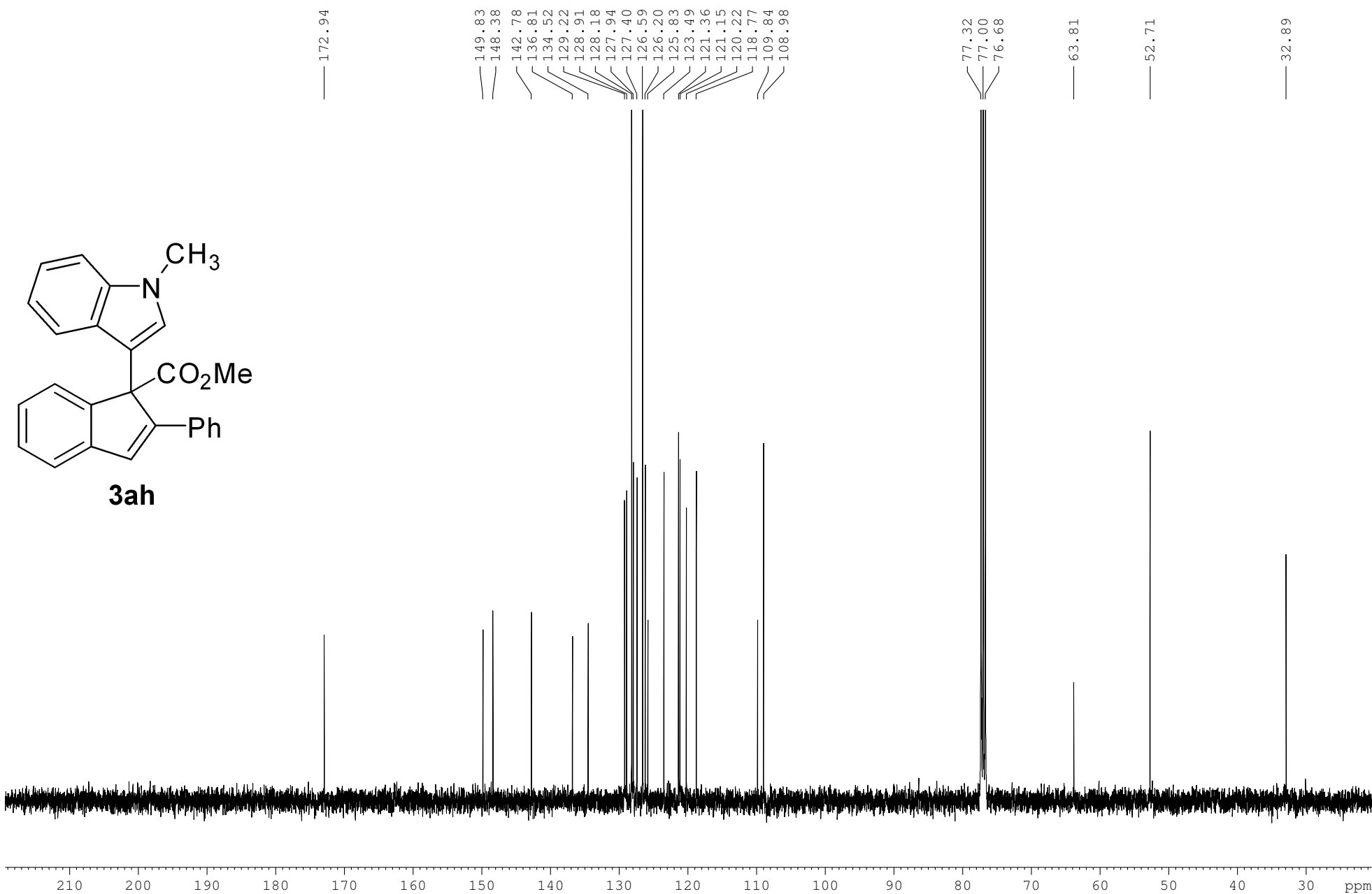


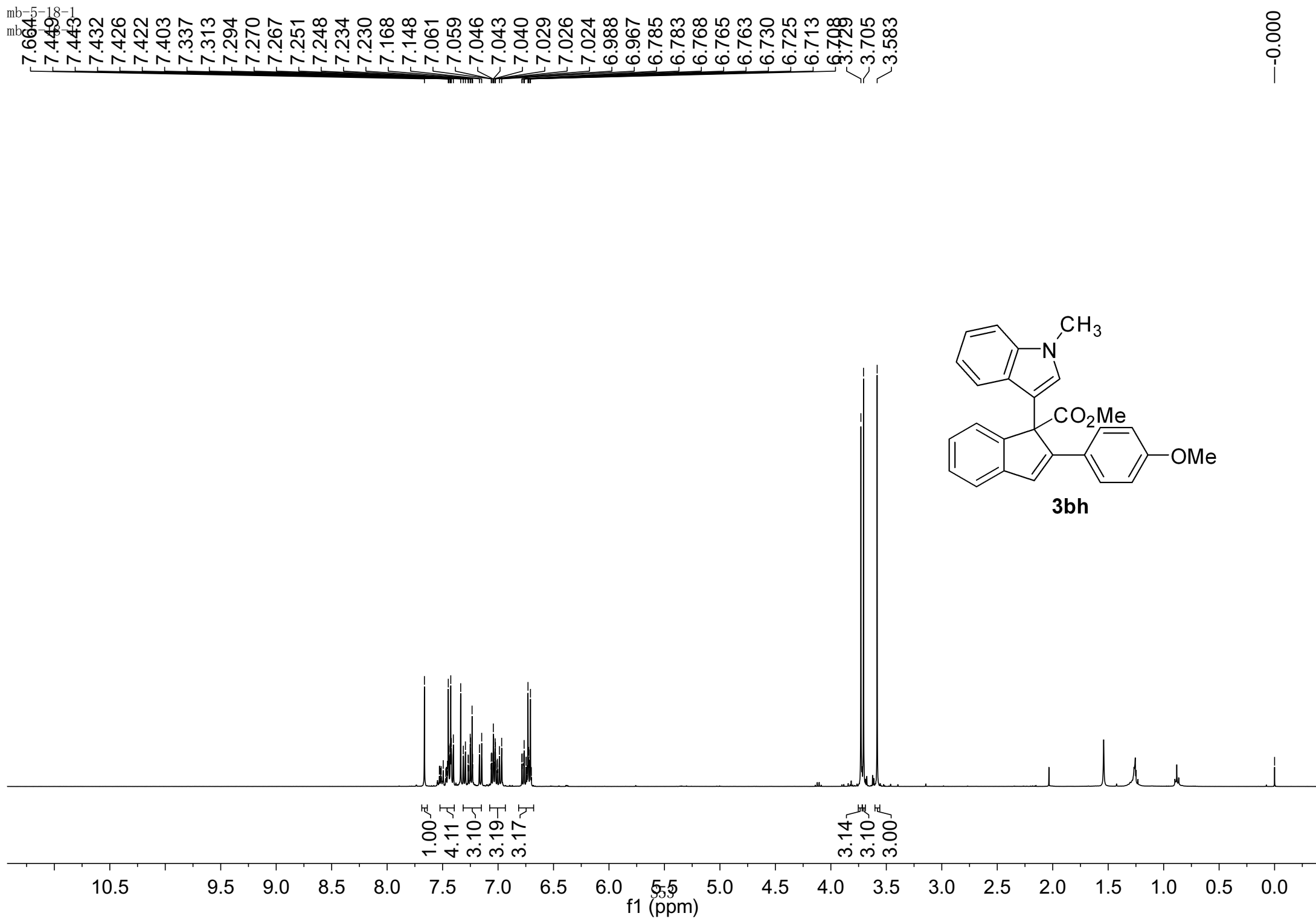


mb-5-139-2



mb-5-139-4





mb-5-18-1
mb-5-18-1

—173.020

—158.990

149.587

148.251

143.118

136.903

129.110

127.934

127.848

127.424

127.149

125.942

125.737

123.406

121.183

120.986

120.344

118.799

113.656

110.254

108.946

77.318

77.000

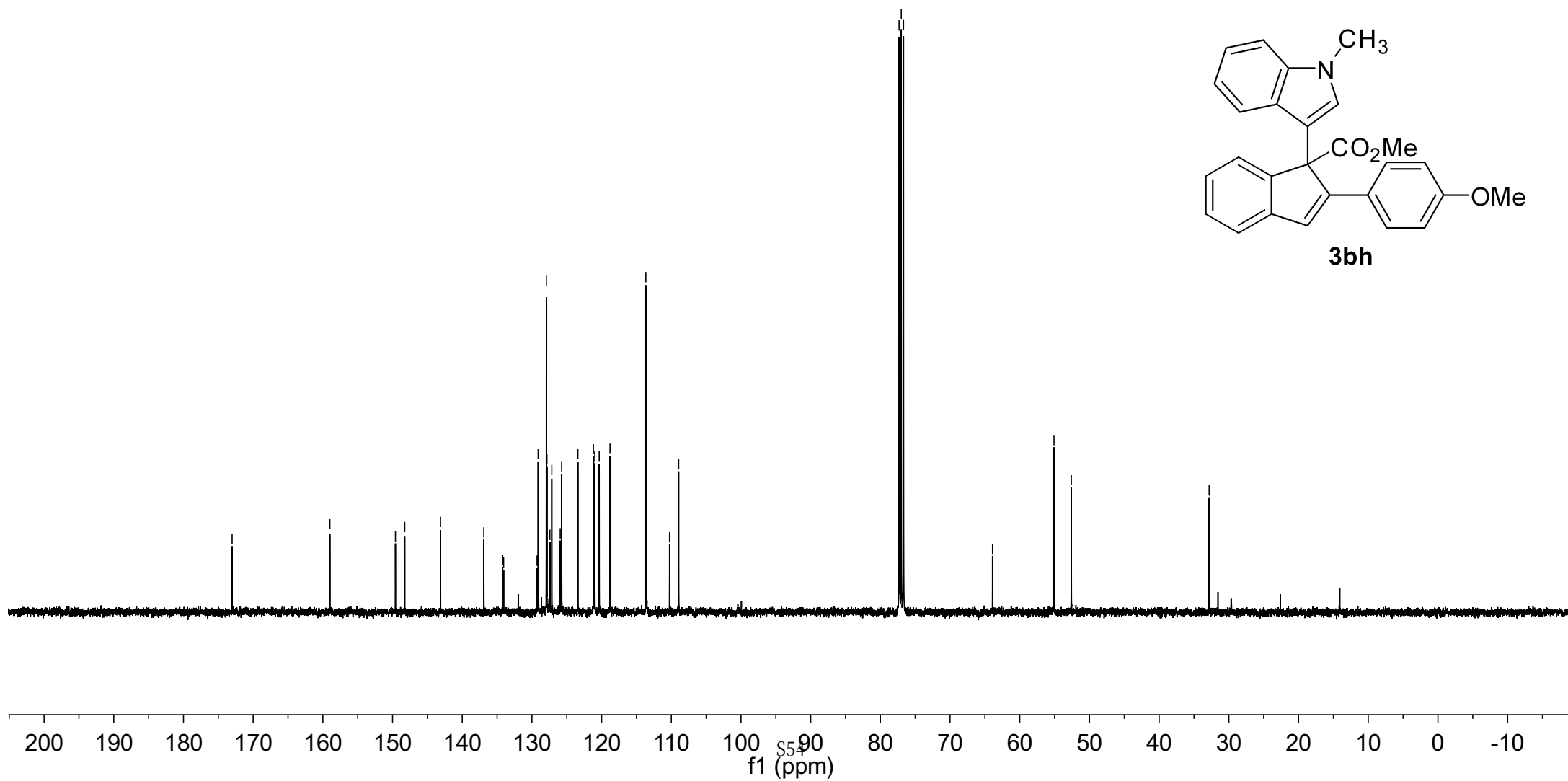
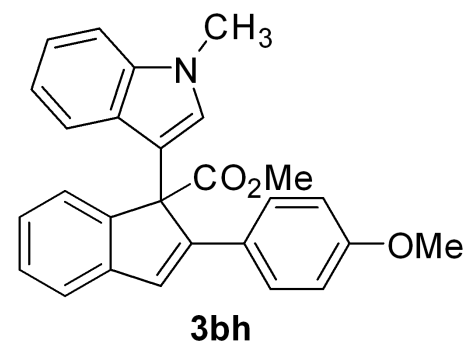
76.682

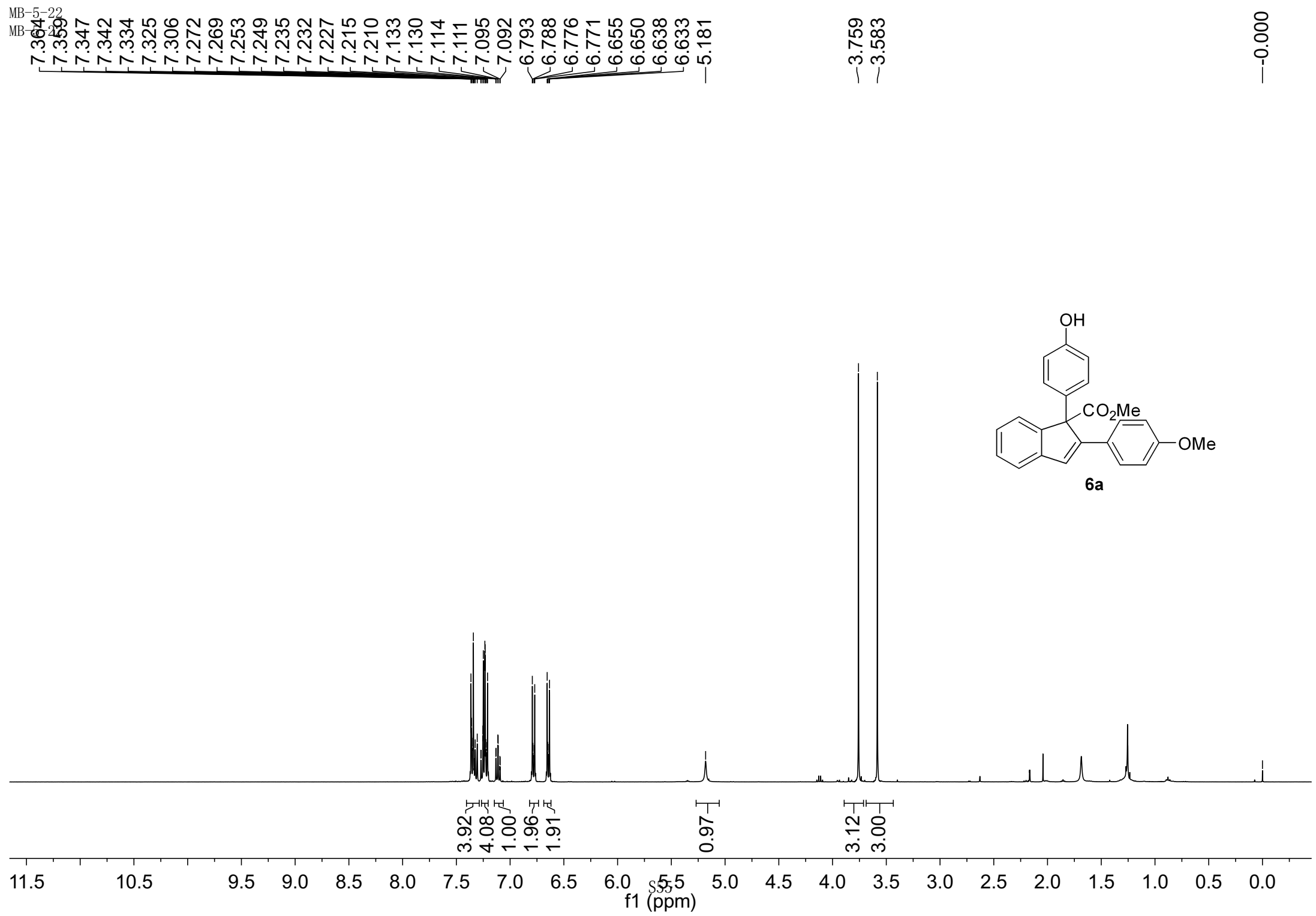
—63.910

~55.088

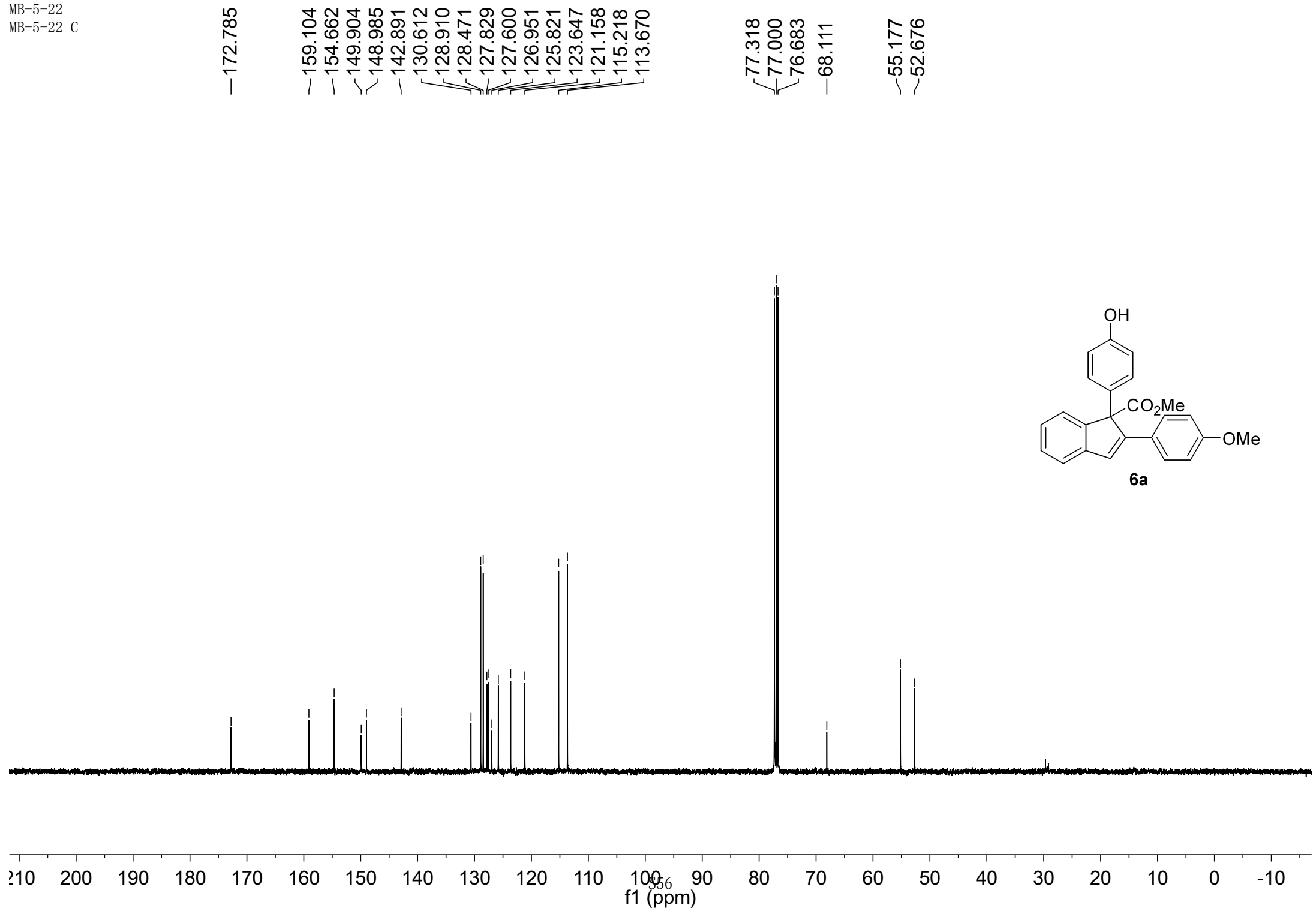
~52.602

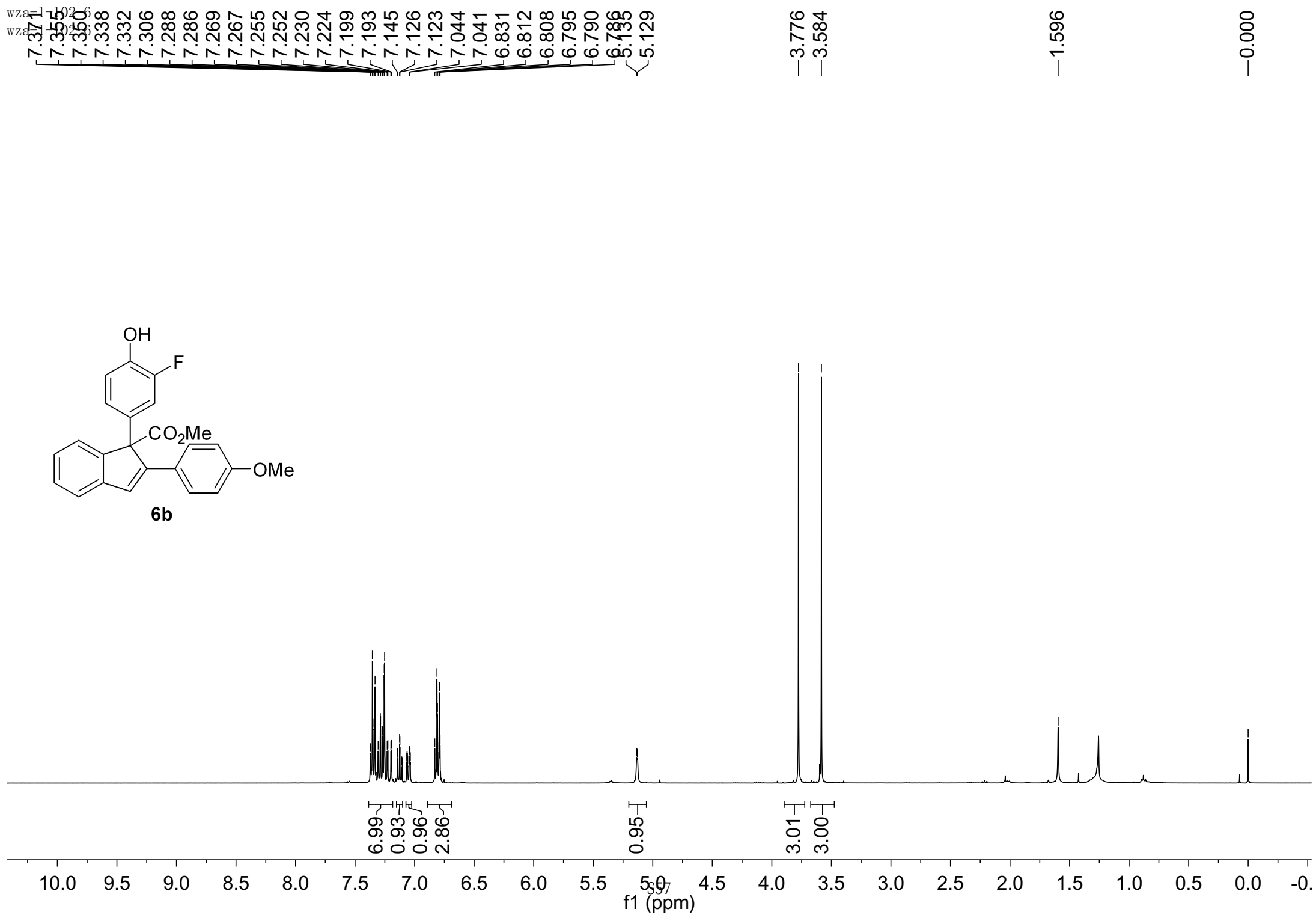
—32.828





MB-5-22
MB-5-22 C





wza-1-102-6
wza-1-102-6

—172.322

—159.273

149.598

149.417

148.567

142.876

142.583

142.440

137.482

131.422

128.356

128.064

127.716

126.595

125.945

124.021

123.989

123.458

121.317

116.915

115.251

115.049

113.816

77.318

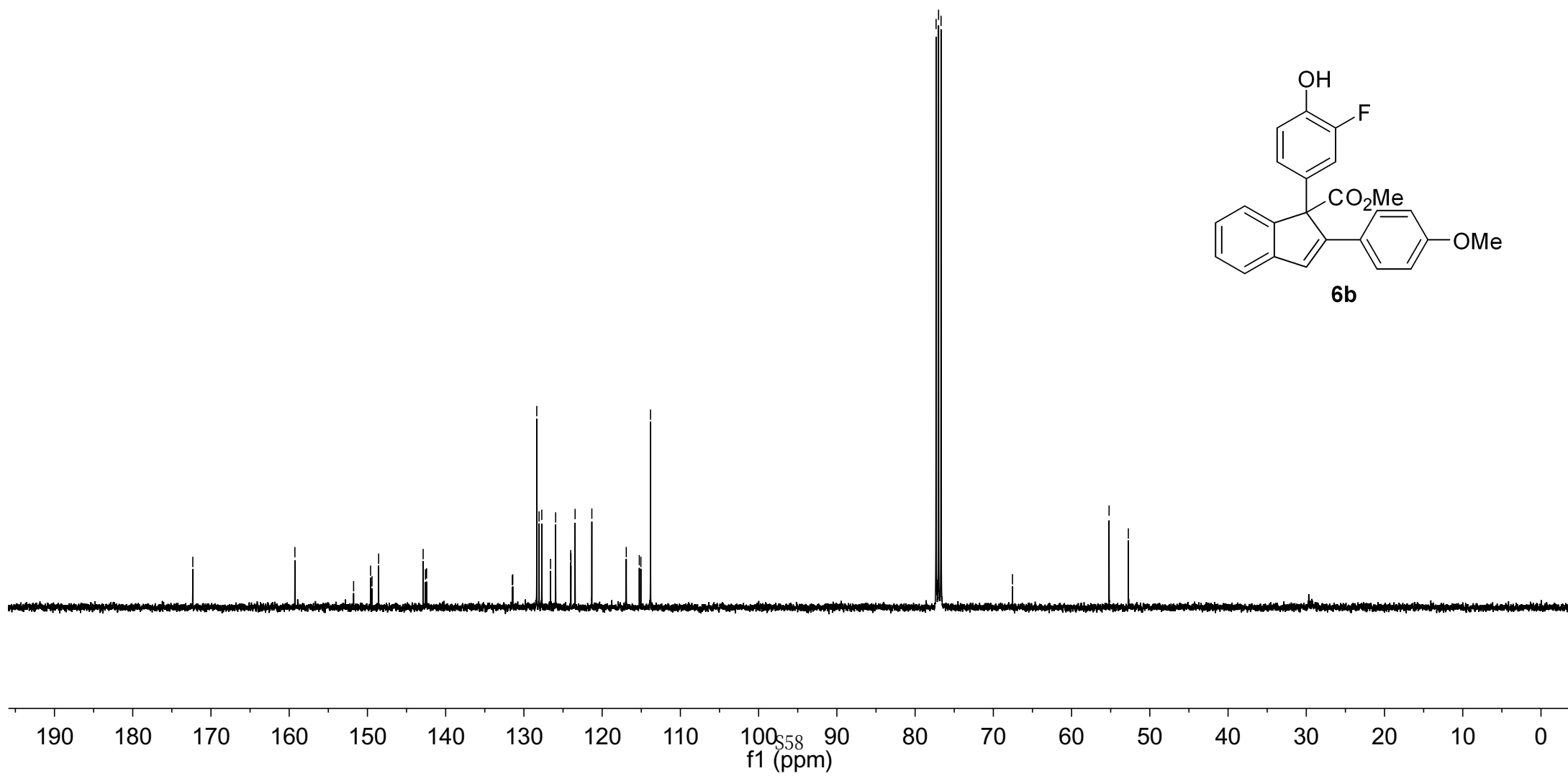
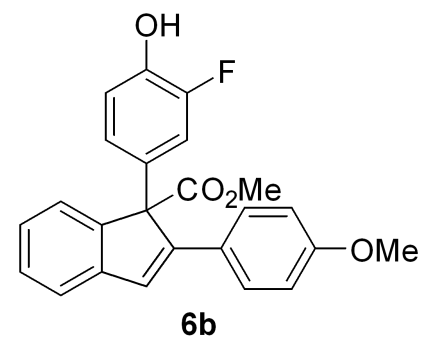
77.000

76.683

—67.554

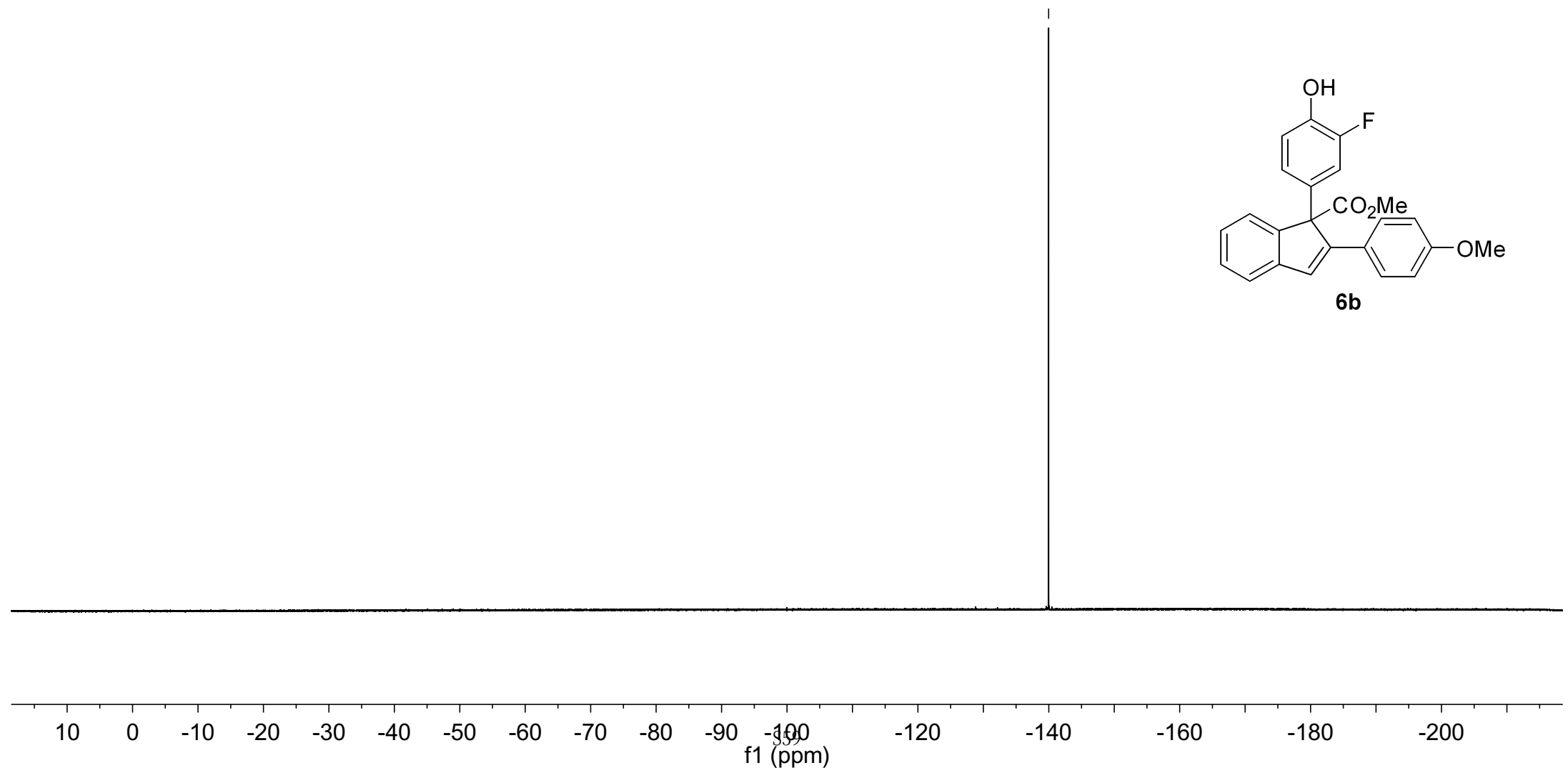
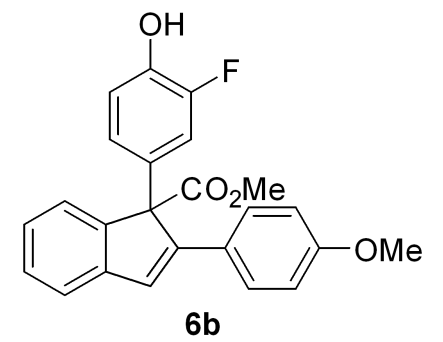
~55.204

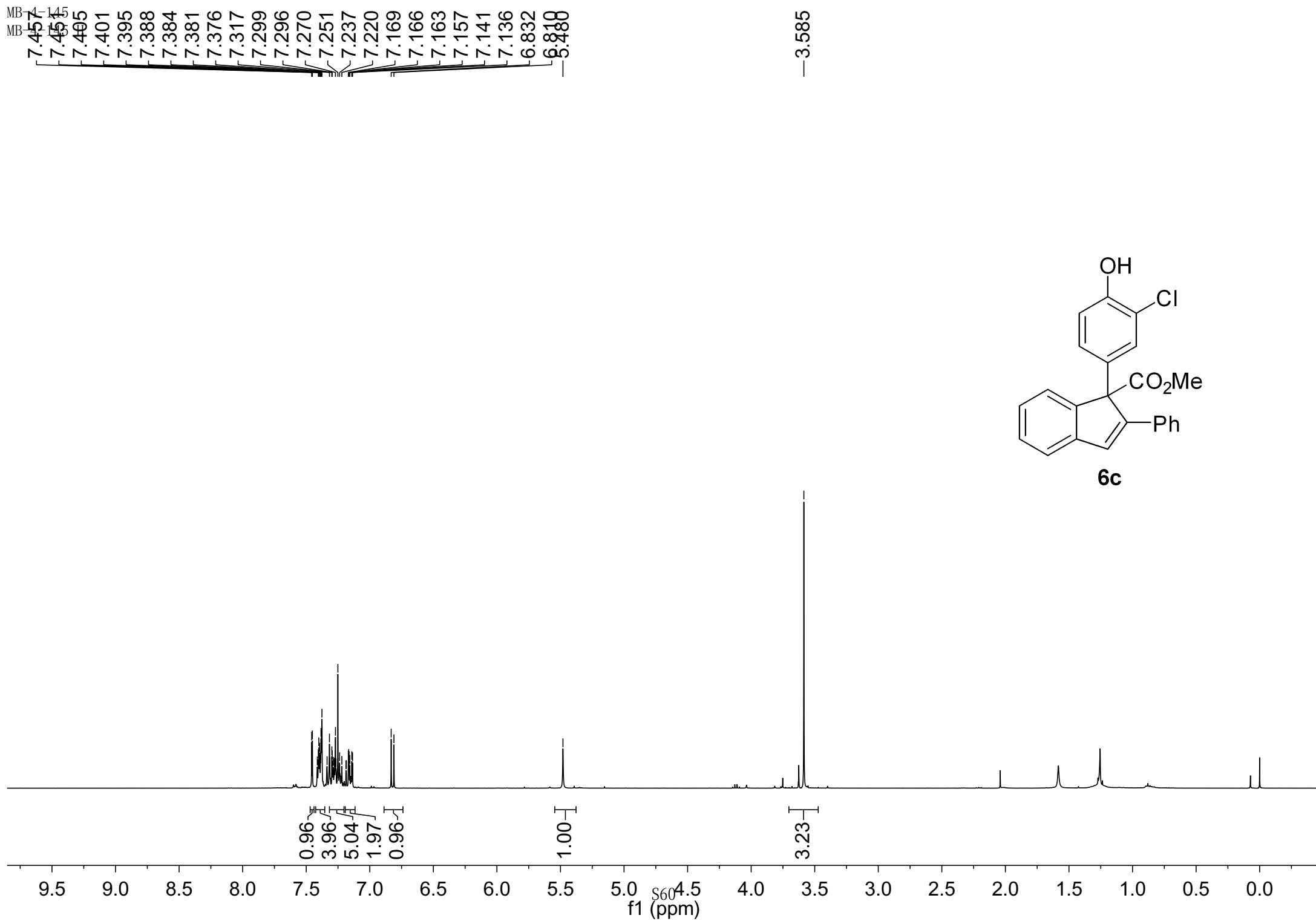
~52.744



wza-1-102
wza-1-102 F

--139.977





MB-4-145
MB-4-145 C

—172.148

150.409

149.917

148.662

142.604

133.869

131.621

129.615

128.343

128.166

128.154

127.787

127.782

127.022

126.435

123.577

121.690

119.633

116.000

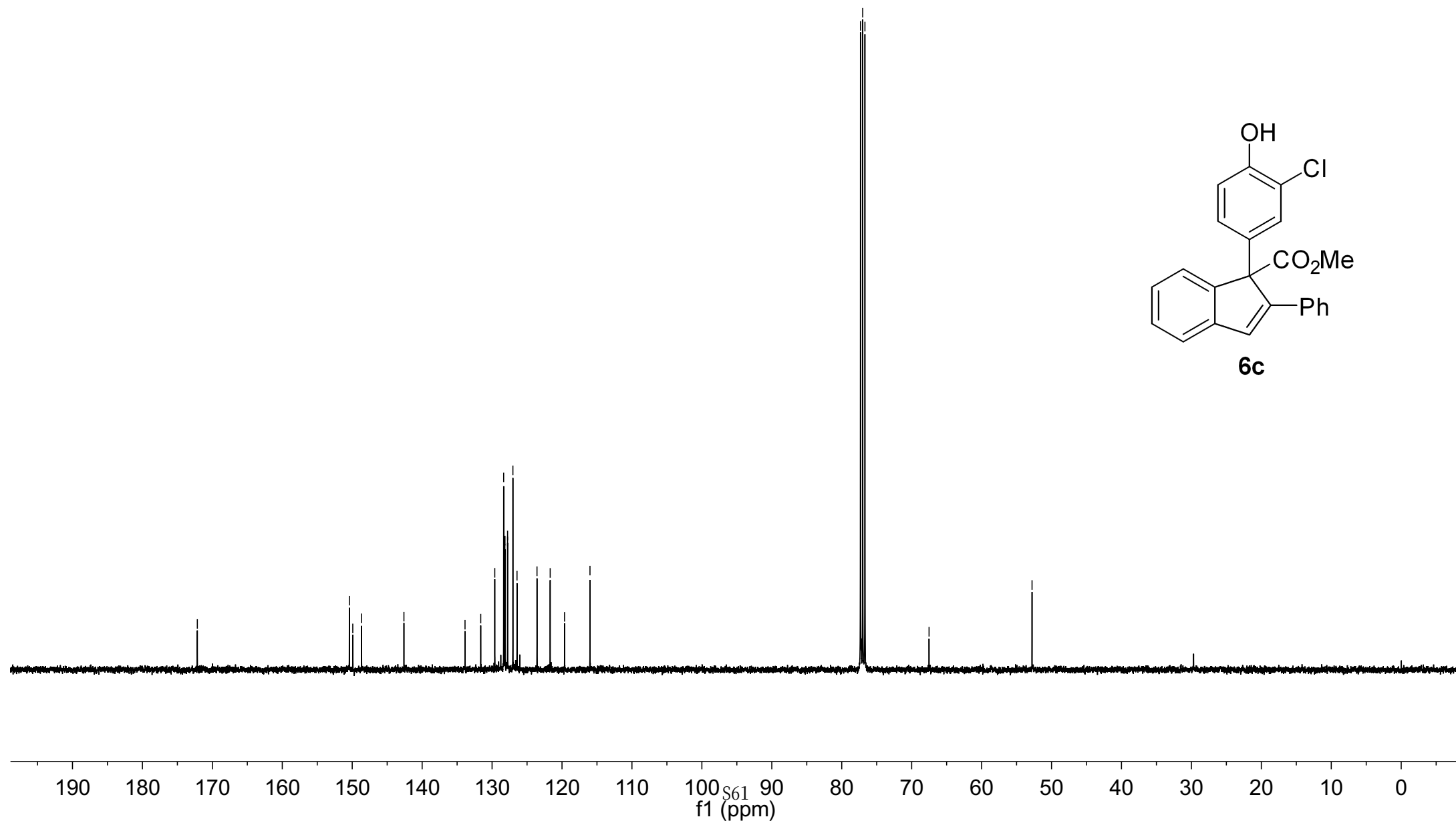
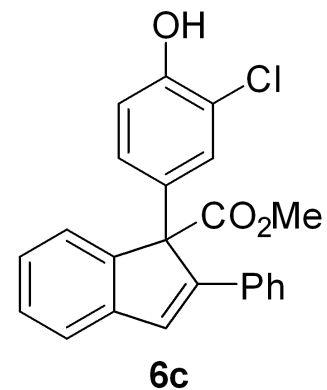
77.318

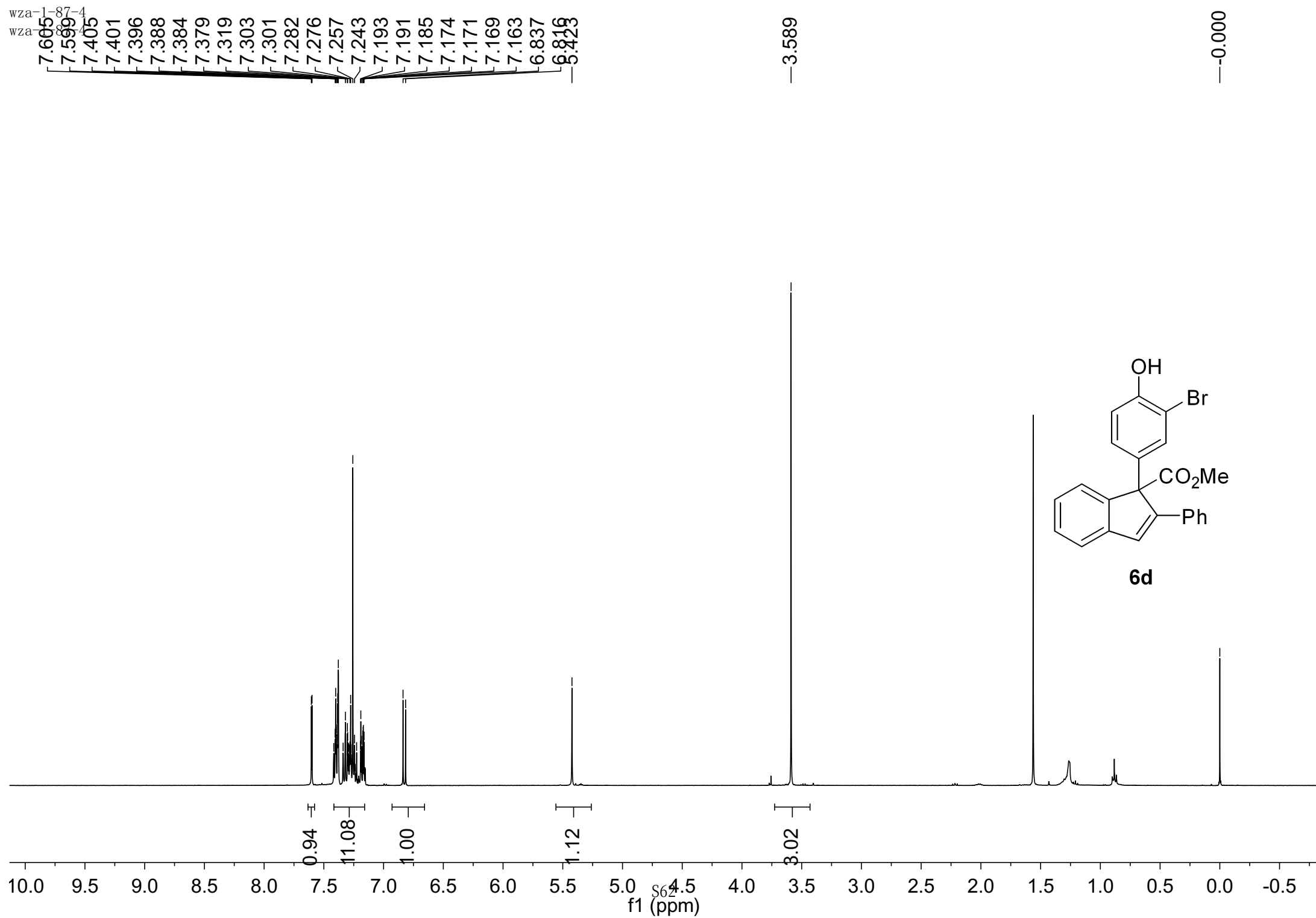
77.000

76.683

—67.514

—52.787





wza-1-87-4
wza-1-87-4

—172.147

151.331

149.918

148.658

142.609

133.870

132.011

131.048

129.635

128.571

128.355

128.178

127.804

127.036

126.452

123.589

121.703

115.812

110.014

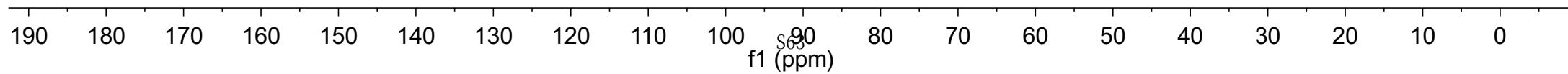
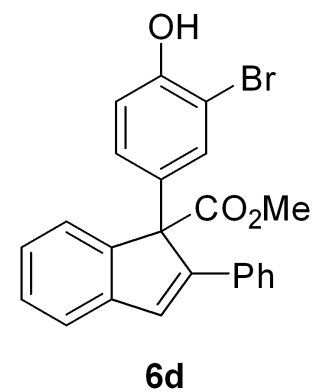
77.317

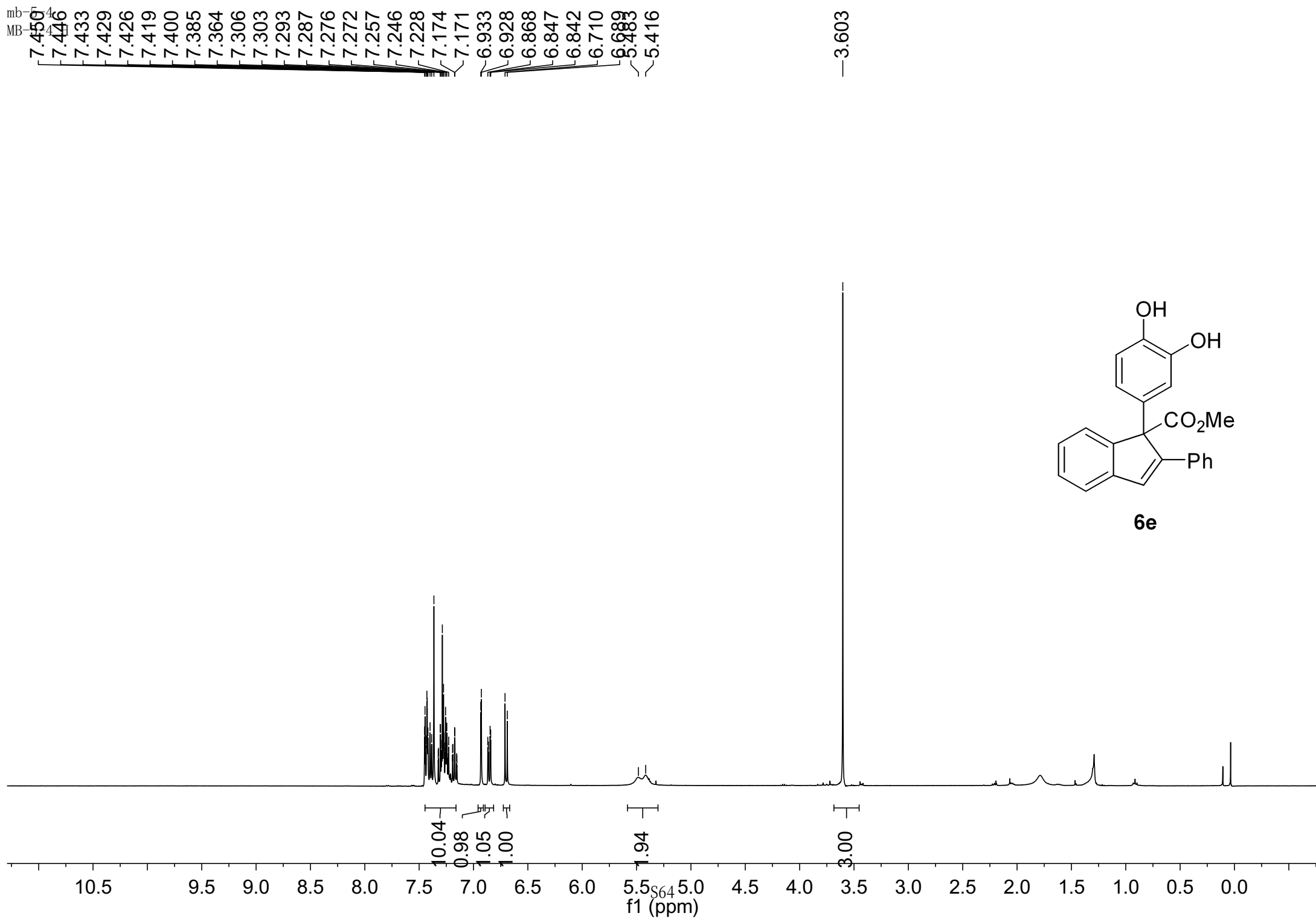
77.000

76.683

—67.432

—52.801





mb-5-4
mb-5-4

—172.495

150.266

149.054

143.105

142.853

142.542

134.273

131.039

129.414

128.192

127.951

127.624

127.161

126.280

123.824

121.506

120.451

115.217

114.857

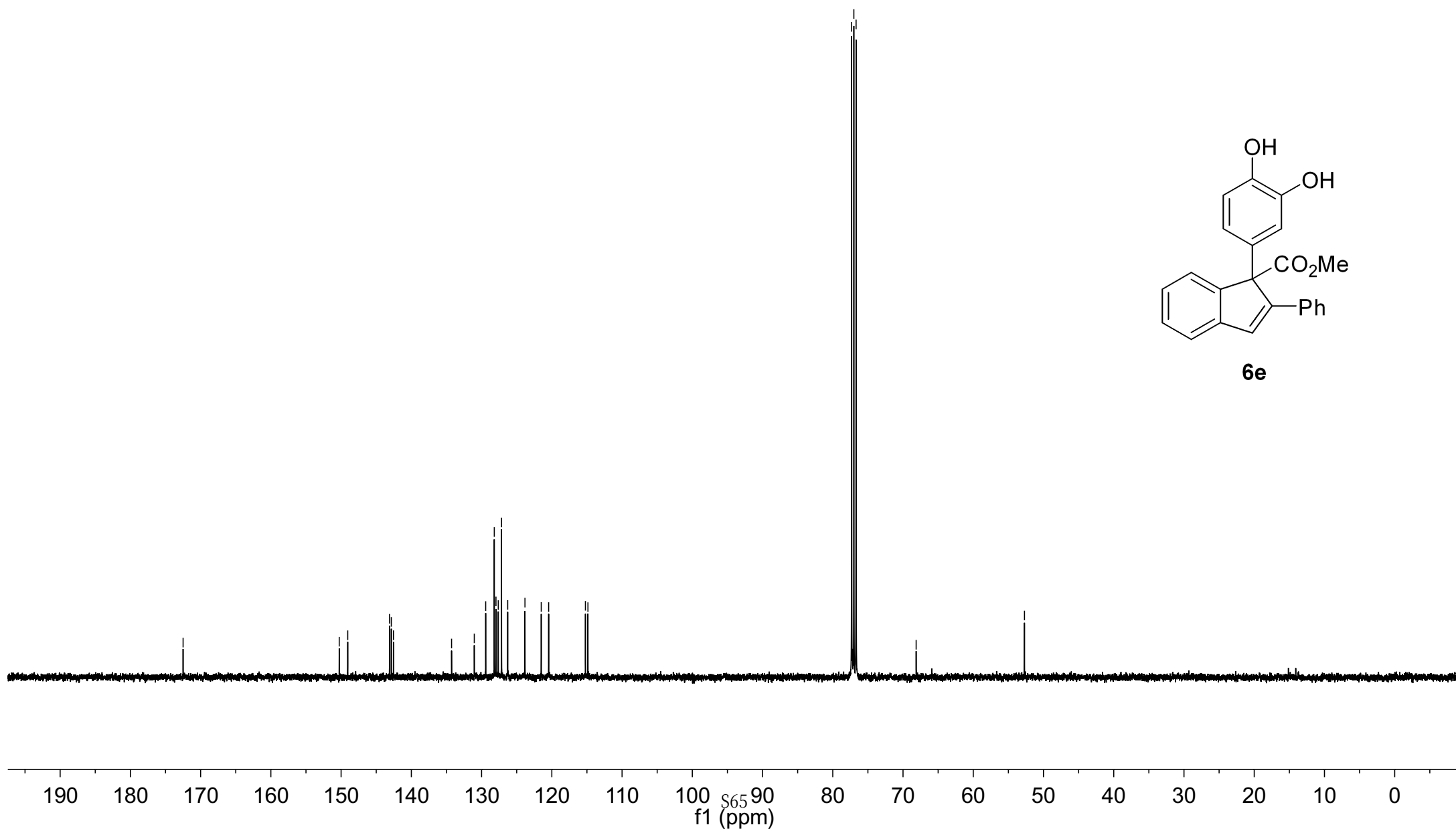
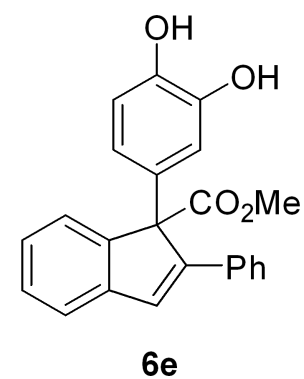
77.317

77.000

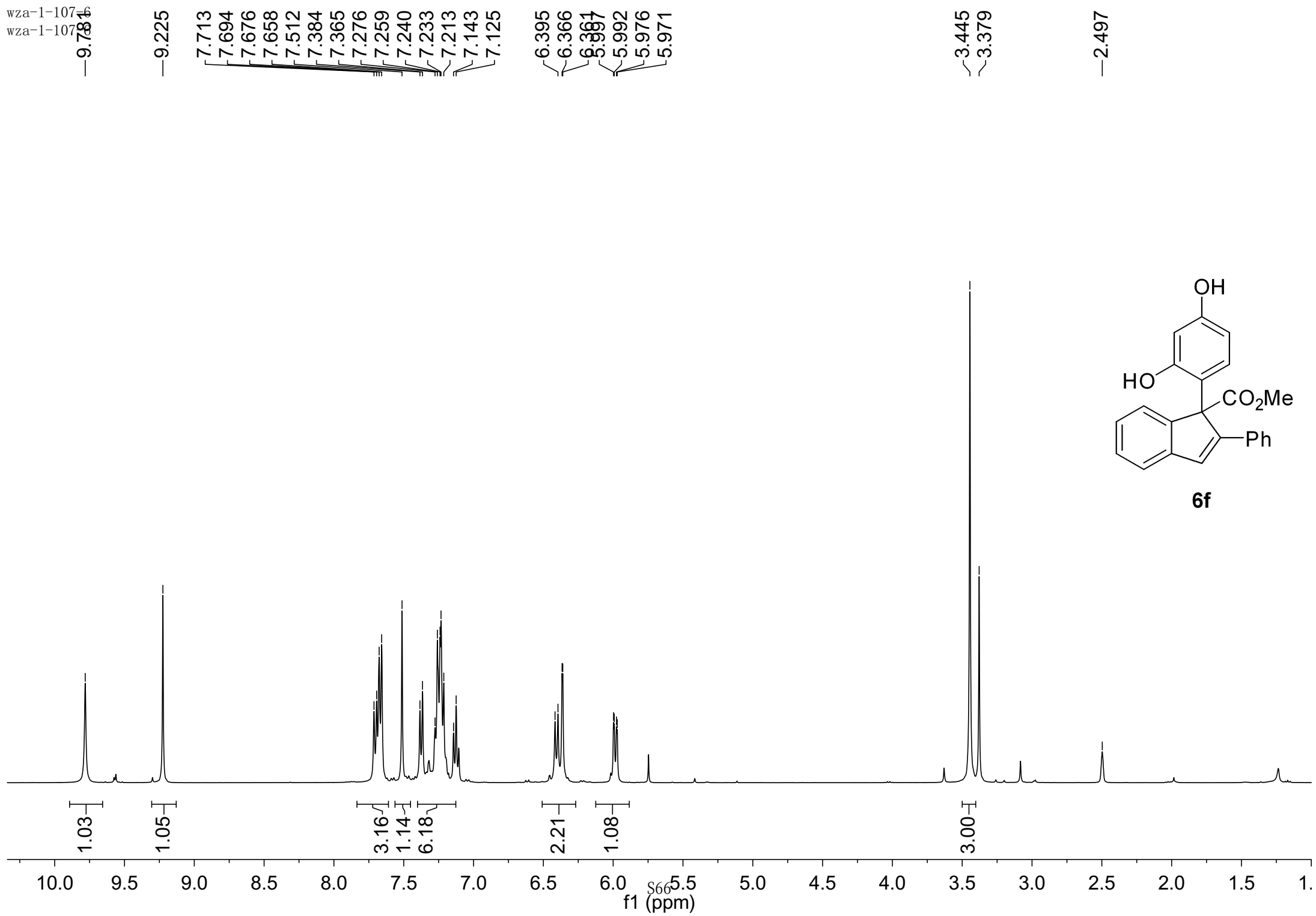
76.682

—68.131

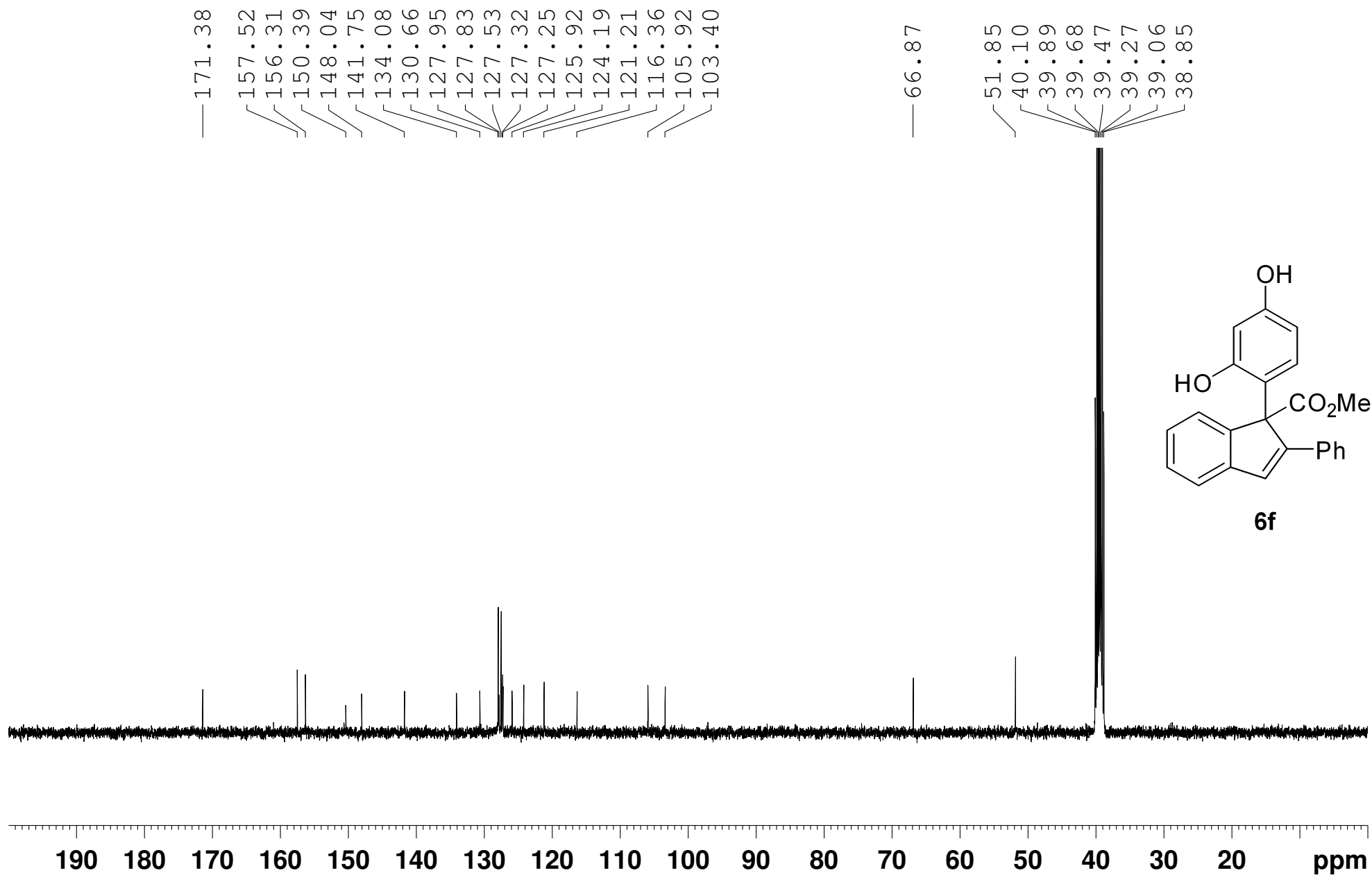
—52.709



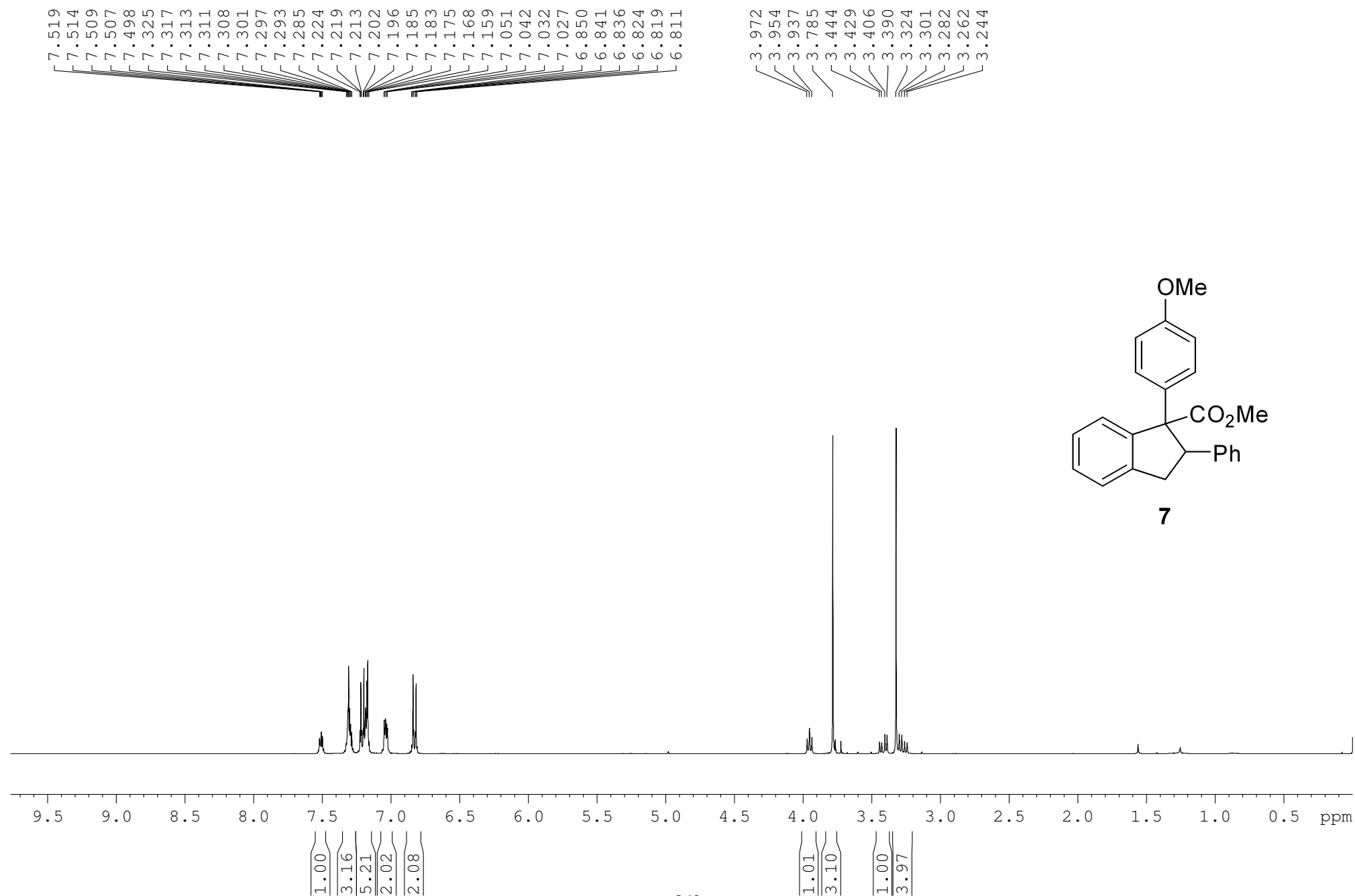
wza-1-107-6
wza-1-107



wza-1-103-1



mb-5-37-1



mb-5-37
mb-5-37

—172.732

—158.455

143.478

141.155

134.429

129.102

128.604

127.820

127.685

127.461

126.685

126.656

124.652

113.361

77.317

77.000

76.682

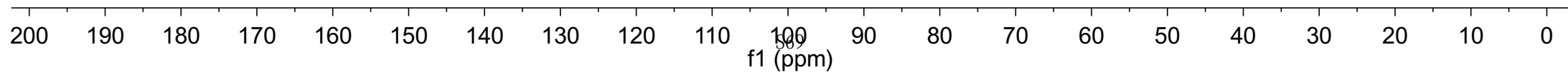
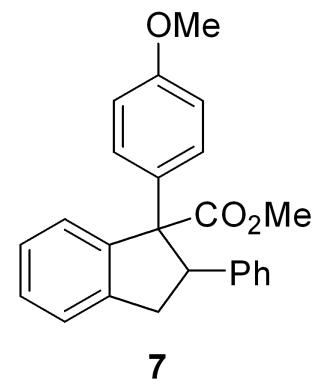
—68.429

—58.683

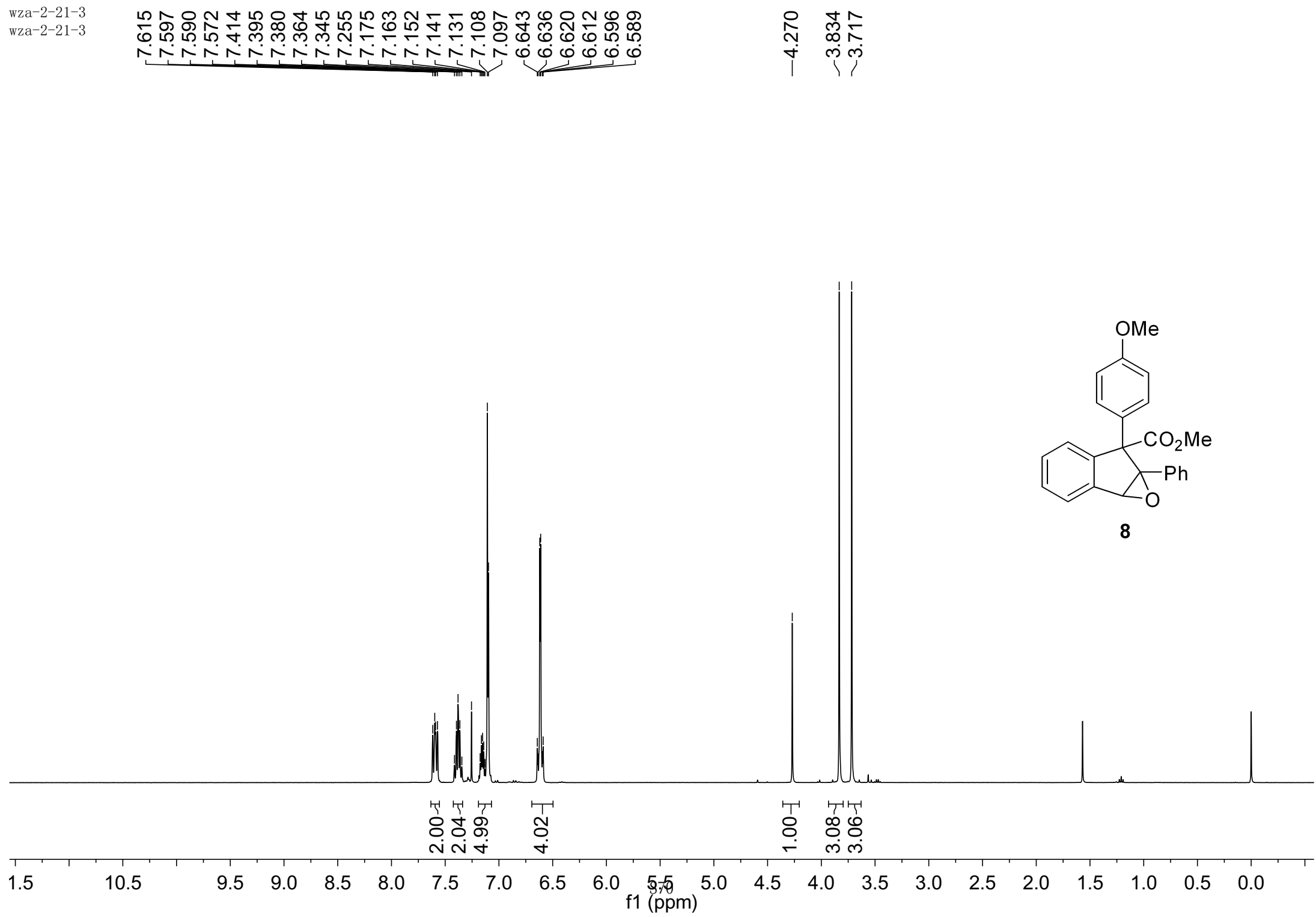
—55.165

—51.439

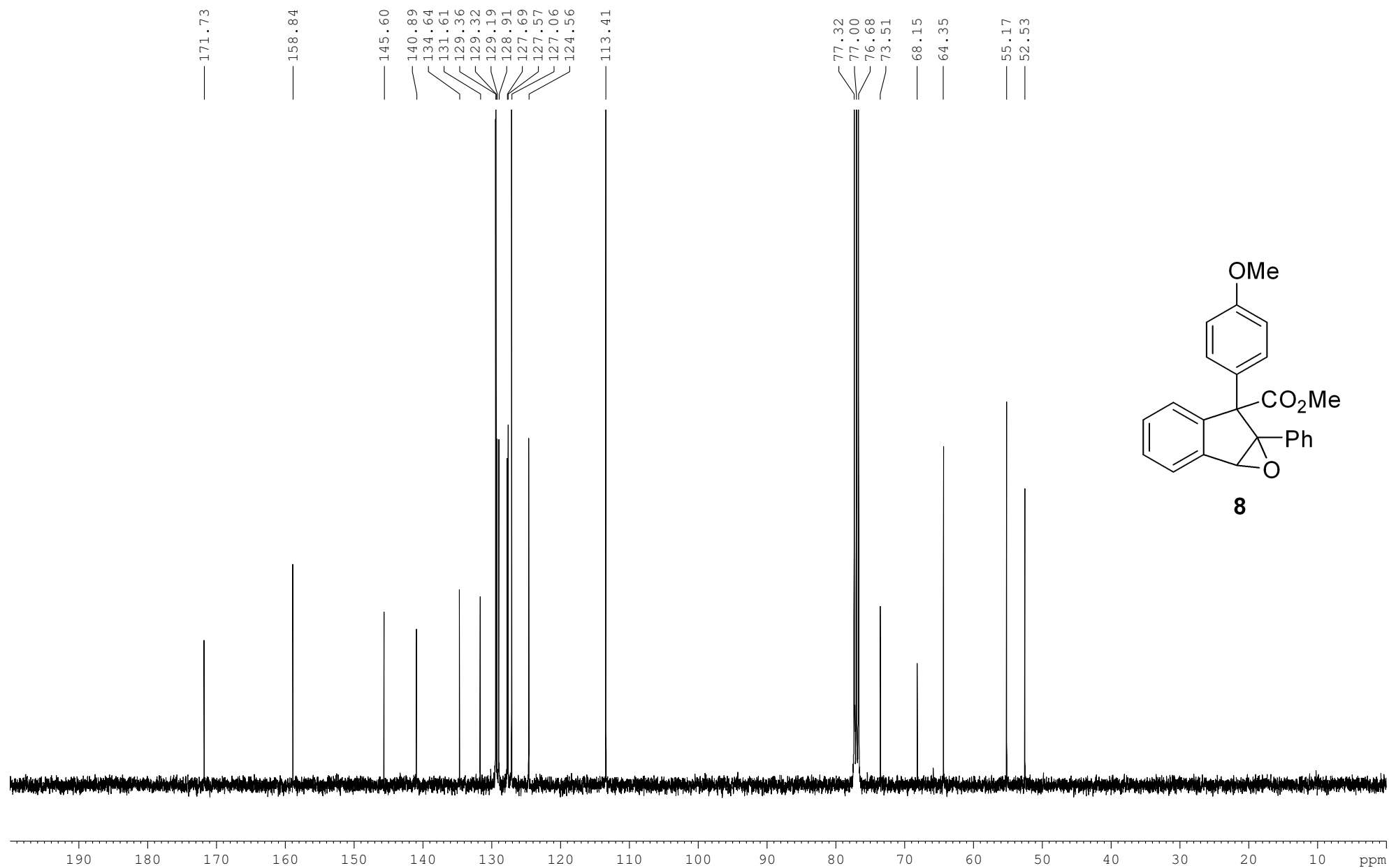
—37.448



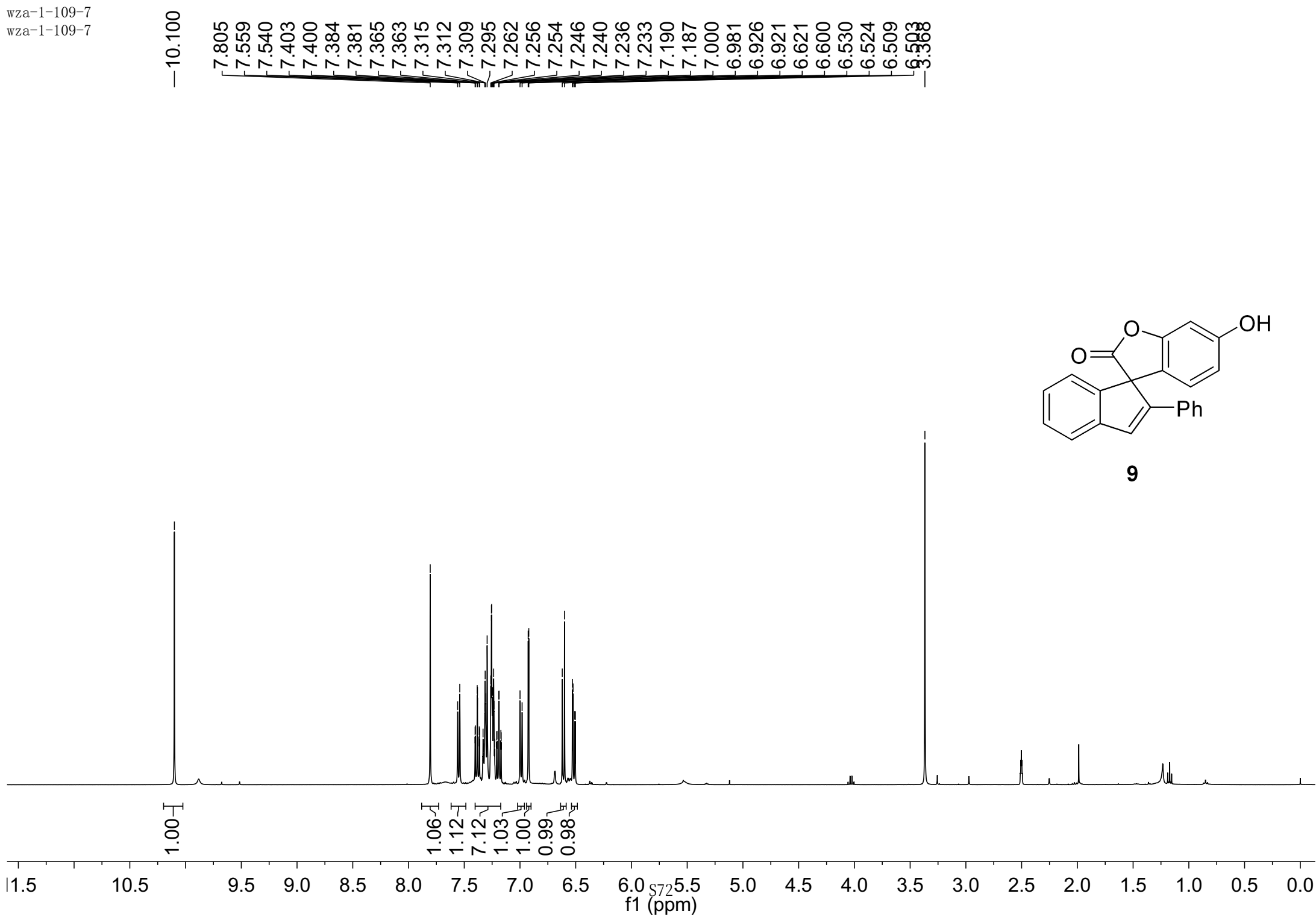
wza-2-21-3
wza-2-21-3

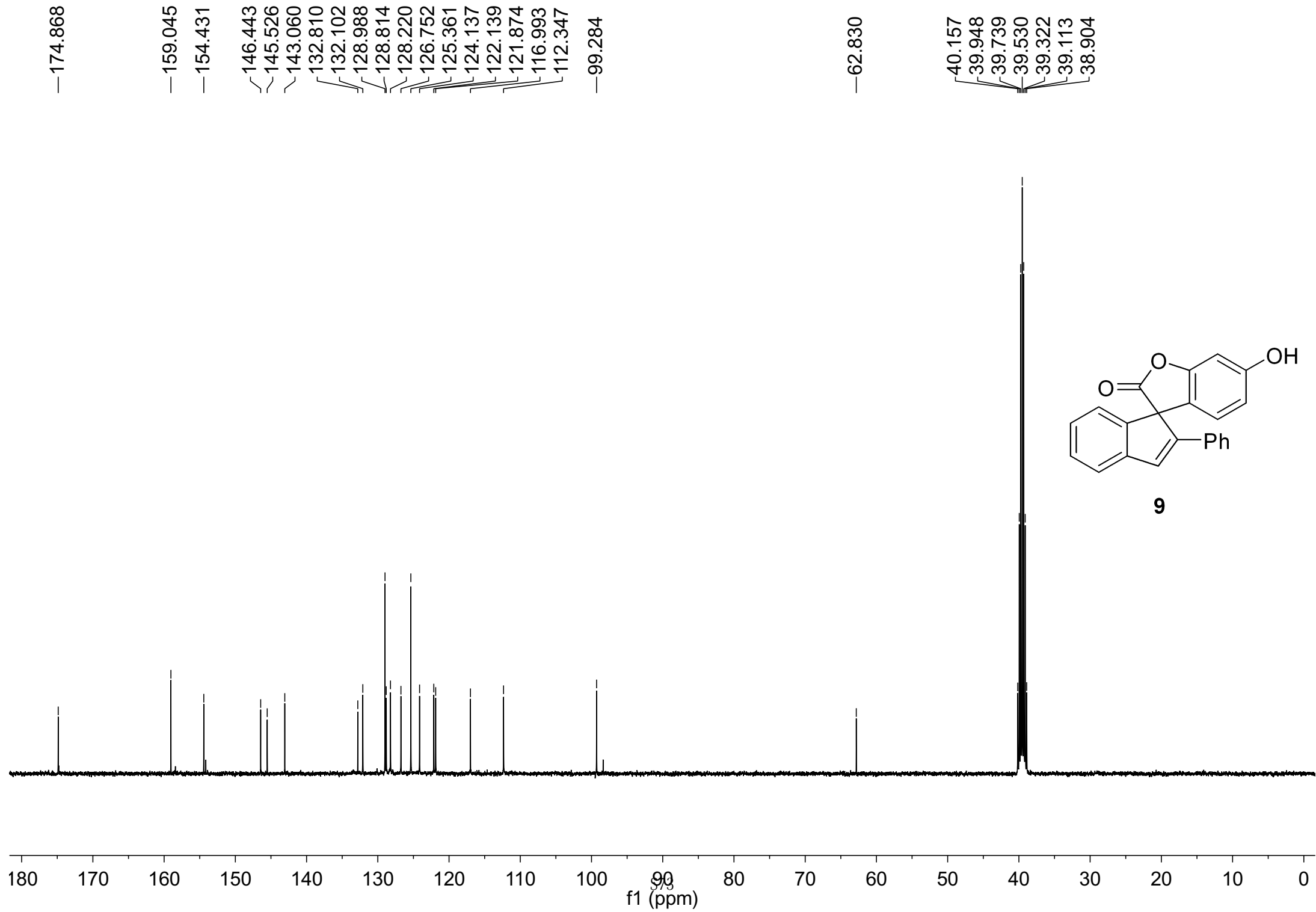


wza-2-21-3



wza-1-109-7
wza-1-109-7





wza-1-19-2-1
wza-1-19-2-1

