

Supporting Information

Nickel-Catalyzed Direct Arylation of Azoles with Aryl Bromides

Hitoshi Hachiya, Koji Hirano, Tetsuya Satoh, and Masahiro Miura*

*Department of Applied Chemistry, Faculty of Engineering, Osaka University, Suita, Osaka 565-0871,
Japan*

Contents

Instrumentation and Chemicals	S1
Experimental Procedure for the Nickel-Catalyzed Direct Arylation of Azoles with Aryl Bromides	S2
Characterization Data for Products	S3–S26

Instrumentation and Chemicals

^1H and ^{13}C NMR spectra were recorded at 400 MHz and 100 MHz, respectively, for CDCl_3 solutions. MS data were obtained by EI. GC analysis was carried out using a silicon OV-17 column (i. d. 2.6 mm x 1.5 m) or a CBP-1 capillary column (i. d. 0.5 mm x 25 m). TLC analyses were performed on commercial glass plates bearing 0.25-mm layer of Merck Silica gel 60F₂₅₄. Silica gel (Wakogel 200 mesh) was used for column chromatography.

Unless otherwise noted, materials obtained from commercial suppliers were used without further purification. Diglyme and *o*-xylene were freshly distilled from CaH_2 prior to use. NiBr_2 and 2,9-dimethyl-1,10-phenanthroline hydrate were purchased from Wako Pure Chemical Co. $\text{NiBr}_2 \cdot \text{diglyme}$ and 1,10-phenanthroline were obtained from Aldrich. Zn powder was available from Kanto Chemical Co. 5-Aryloxazoles **6a** and **6b** were prepared by the van Lausen reaction with TosMIC and the corresponding aldehydes.¹ All reactions were carried out under nitrogen atmosphere.

¹ van Leusen, A. M.; Hoogenboom, B. E.; Sinderius, H. *Tetrahedron Lett.* **1972**, 13, 2369.

Typical Procedure for the Nickel-Catalyzed Direct Arylation of Azoles with Aryl Bromides

Synthesis of 2a (Table 2, entry 1): NiBr₂ (11 mg, 0.050 mmol), 1,10-phenanthroline (11 mg, 0.060 mmol), LiOt-Bu (160 mg, 2.0 mmol), benzothiazole (**1**, 68 mg, 0.50 mmol), 4-bromotoluene (**2a**, 103 mg, 0.60 mmol), diglyme (3.0 mL), and dibenzyl (ca. 50 mg, internal standard) were placed in a 20 mL two-necked reaction flask equipped with a reflux condenser. After being stirred for 10 min at room temperature, the solution was heated at 150 °C for 4 h. The consumption of **1** was confirmed by GC analysis, and the resulting mixture was then quenched with water. The mixture was extracted with diethyl ether, and the combined organic layer was dried over sodium sulfate. Concentration in vacuo followed by silica gel column purification gave 2-(4-methylphenyl)benzothiazole (**3a**, 74 mg, 0.33 mmol) in 67% yield.

Synthesis of 5c (Table 4, entry 1): NiBr₂•diglyme (18 mg, 0.050 mmol), 2,9-dimethyl-1,10-phenanthroline hydrate (13 mg, 0.060 mmol), LiOt-Bu (160 mg, 2.0 mmol), benzoxazole (**4**, 119 mg, 1.0 mmol), 2-bromotoluene (**2c**, 86 mg, 0.50 mmol), Zn powder (16 mg, 0.25 mmol), *o*-xylene (3.0 mL), and 1-methylnaphthalene (ca. 50 mg, internal standard) were placed in a 20 mL two-necked reaction flask equipped with a reflux condenser. The solution was stirred for 10 min at room temperature and then heated at 150 °C. After the consumption of **2c** was checked by GC analysis, the resulting mixture was allowed to cool to room temperature and quenched with water. Extraction with diethyl ether, evaporation, and purification by column chromatography on silica gel afforded 2-(2-methylphenyl)benzoxazole (**5c**, 55 mg, 0.27 mmol) in 53% yield.

Characterization Data for Compounds

¹H and ¹³C NMR spectra for all compounds are attached in the last part. Compounds **3a**,² **3b**,³ **3c**,⁴ **3e**,⁵ **3f**,⁶ **3g**,⁷ **3h**,⁶ **3i**,⁸ **3j**,⁹ **3a'**,¹⁰ **5c**,¹¹ and **5e**¹² are found in the literature.

2-(4-Methylphenyl)benzothiazole (3a) m.p. 80-82 °C; ¹H NMR (400 MHz, CDCl₃) δ 2.44 (s, 3H), 7.31 (d, *J* = 8.1 Hz, 2H), 7.36-7.40 (m, 1H), 7.47-7.52 (m, 1H), 7.90 (d, *J* = 8.1 Hz, 1H), 8.00 (dt, *J* = 8.1 Hz, 1.8 Hz, 2H), 8.07 (d, *J* = 8.1 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 21.71, 121.77, 123.27, 125.20, 126.44, 127.70, 129.92, 131.20, 135.18, 141.62, 154.41, 168.44; HRMS *m/z* (M⁺) calcd for C₁₄H₁₁NS: 225.0612, found: 225.0607.

2-(3-Methylphenyl)benzothiazole (3b) m.p. 63-66 °C; ¹H NMR (400 MHz, CDCl₃) δ 2.47 (s, 3H), 7.32 (d, *J* = 7.7 Hz, 1H), 7.39 (t, *J* = 7.7 Hz, 2H), 7.48-7.53 (m, 1H), 7.87-7.92 (m, 2H), 7.96 (s, 1H), 8.09 (d, *J* = 8.1 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 21.55, 121.80, 123.40, 125.08, 125.32, 126.48, 128.22, 129.12, 132.00, 133.78, 135.27, 139.07, 154.38, 168.52; HRMS *m/z* (M⁺) calcd for C₁₄H₁₁NS: 225.0612, found: 225.0605.

2-(2-Methylphenyl)benzothiazole (3c) m.p. 51-53 °C; ¹H NMR (400 MHz, CDCl₃) δ 2.69 (s, 3H), 7.31-7.45 (m, 4H), 7.51-7.55 (m, 1H), 7.78 (d, *J* = 7.7 Hz, 1H), 7.95 (d, *J* = 7.3 Hz, 1H), 8.13 (d, *J* = 8.1 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 21.53, 121.55, 123.60, 125.28, 126.30, 126.32, 130.19, 130.74, 131.74, 133.33, 135.83, 137.48, 154.04, 168.18; HRMS *m/z* (M⁺) calcd for C₁₄H₁₁NS: 225.0612, found: 225.0603.

2-(2,5-Dimethylphenyl)benzothiazole (3d) oil; ¹H NMR (400 MHz, CDCl₃) δ 2.38 (s, 3H), 2.61 (s, 3H), 7.17-7.24 (m, 2H), 7.37-7.41 (m, 1H), 7.48-7.52 (m, 1H), 7.59 (s, 1H), 7.92 (d, *J* = 8.4 Hz, 1H), 8.10 (d, *J* = 8.1 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 21.07 (Two signals merged.), 121.53, 123.55, 125.21, 126.28, 131.02, 131.22, 131.69, 133.11, 134.29, 135.84, 135.87, 153.98, 168.36; HRMS *m/z*

² Egi, M.; Liebeskind, L. S. *Org. Lett.* **2003**, 5, 2369.

³ Fukuhara, T.; Hasegawa, C.; Hara, S. *Synthesis* **2007**, 1528.

⁴ Kodomari, M.; Tamaru, Y.; Aoyama, T. *Synth. Commun.* **2004**, 34, 3029.

⁵ Chakraborti, A. K.; Selvam, C.; Kaur, G.; Bhagat, S. *Synlett* **2004**, 851.

⁶ Matsushita, H.; Lee, S.-H.; Joung, M.; Clapham, B.; Janda, K. D. *Tetrahedron Lett.* **2004**, 45, 313.

⁷ Lianqing, C.; Yang, C.; Li, S.; Qin, J. *Spectrochim. Acta. Part A* **2007**, 68, 317.

⁸ Inamoto, K.; Hasegawa, C.; Hiroya, K.; Doi, T. *Org. Lett.* **2008**, 10, 5147.

⁹ Itoh, T.; Mase, T. *Org. Lett.* **2007**, 9, 3687.

¹⁰ Jensen, J.; Skjaerbaek, N.; Vedsø, P. *Synthesis* **2001**, 128.

¹¹ Derridj, F.; Djebbar, S.; Benali-Baitich, O.; Doucet, H. *J. Organomet. Chem.* **2008**, 693, 135.

¹² Anderson, B. A.; Haru, N. K. *Synthesis* **1996**, 583.

(M⁺) calcd for C₁₅H₁₃NS: 239.0769, found: 239.0772.

2-(1-Naphthyl)benzothiazole (3e) m.p. 78-80 °C; ¹H NMR (400 MHz, CDCl₃) δ 7.45-7.49 (m, 1H), 7.55-7.67 (m, 4H), 7.94-8.02 (m, 4H), 8.22 (d, *J* = 8.1 Hz, 1H), 8.96 (d, *J* = 8.8 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 121.61, 123.80, 125.20, 125.51, 126.13, 126.48, 126.72, 127.84, 128.62, 129.59, 130.91, 131.09, 131.28, 134.26, 135.70, 154.40, 167.80; HRMS *m/z* (M⁺) calcd for C₁₇H₁₁NS: 261.0612, found: 225.0619.

2-(4-Methoxyphenyl)benzothiazole (3f) m.p. 115-116 °C; ¹H NMR (400 MHz, CDCl₃) δ 3.88 (s, 3H), 7.01 (d, *J* = 8.8 Hz, 2H), 7.36 (t, *J* = 7.7 Hz, 1H), 7.48 (t, *J* = 8.1 Hz, 1H), 7.88 (d, *J* = 8.4 Hz, 1H), 8.05 (d, *J* = 8.8 Hz, 2H), 8.03-8.06 (m, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 55.65, 114.58, 121.69, 123.05, 124.98, 126.38, 126.70, 129.32, 135.10, 154.48, 162.15, 168.03; HRMS *m/z* (M⁺) calcd for C₁₄H₁₁NOS: 241.0561, found: 241.0549.

2-(4-Fluorophenyl)benzothiazole (3g) m.p. 100-102 °C; ¹H NMR (400 MHz, CDCl₃) δ 7.18 (dd, *J* = 8.6 Hz, 8.4 Hz, 2H), 7.37-7.41 (m, 1H), 7.47-7.52 (m, 1H), 7.90 (d, *J* = 8.1 Hz, 1H), 8.05-8.11 (m, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 116.38 (d, *J* = 22.2 Hz), 121.84, 123.44, 125.48, 126.64, 129.77 (d, *J* = 9.1 Hz), 130.24 (d, *J* = 3.1 Hz), 135.31, 154.37, 164.71 (d, *J* = 251.8 Hz), 166.96; HRMS *m/z* (M⁺) calcd for C₁₃H₈FNS: 229.0361, found: 229.0359.

2-(4-Trifluoromethylphenyl)benzothiazole (3h) m.p. 161-162 °C; ¹H NMR (400 MHz, CDCl₃) δ 7.41-7.45 (m, 1H), 7.51-7.55 (m, 1H), 7.75 (d, *J* = 8.1 Hz, 2H), 7.93 (d, *J* = 7.7 Hz, 1H), 8.11 (d, *J* = 7.7 Hz, 1H), 8.21 (d, *J* = 8.1 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ 121.74, 123.65, 123.83 (q, *J* = 272.4 Hz), 125.79, 126.01 (q, *J* = 3.8 Hz), 126.66, 127.78, 132.48 (q, *J* = 32.8 Hz), 135.23, 136.79, 154.07, 166.04; HRMS *m/z* (M⁺) calcd for C₁₄H₈F₃NS: 279.0330, found: 279.0331.

4-(2-Benzothiazolyl)benzonitrile (3i) m.p. 169-170 °C; ¹H NMR (400 MHz, CDCl₃) δ 7.43-7.48 (m, 1H), 7.53-7.57 (m, 1H), 7.78 (dt, *J* = 8.4 Hz, 1.5 Hz, 2H), 7.95 (d, *J* = 8.1 Hz, 1H), 8.11 (d, *J* = 7.7 Hz, 1H), 8.20 (dt, *J* = 8.4 Hz, 1.8 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ 114.36, 118.46, 122.00, 124.03, 126.28, 127.02, 128.13, 132.96, 135.53, 137.70, 154.26, 165.52; HRMS *m/z* (M⁺) calcd for C₁₄H₁₀N₂S: 236.0408, found: 236.0405.

1-{4-(2-Benzothiazolyl)phenyl}ethanone (3j) m.p. 187-188 °C; ¹H NMR (400 MHz, CDCl₃) δ 2.66 (s, 3H), 7.41-7.45 (m, 1H), 7.51-7.55 (m, 1H), 7.94 (d, *J* = 7.7 Hz, 1H), 8.07 (dt, *J* = 8.4 Hz, 1.8 Hz, 2H), 8.11 (d, *J* = 8.1 Hz, 1H), 8.19 (dt, *J* = 8.4 Hz, 1.8 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ 26.93, 121.94, 123.85, 125.95, 126.83, 127.87, 129.17, 135.53, 137.74, 138.81, 154.37, 166.63, 197.46; HRMS *m/z* (M⁺) calcd for C₁₅H₁₁NOS: 253.0561, found: 253.0560.

2-(4-Methylphenyl)thiazole (3a') oil; ¹H NMR (400 MHz, CDCl₃) δ 2.40 (s, 3H), 7.26 (d, *J* =

8.4 Hz, 2H), 7.30 (d, $J = 3.3$ Hz, 1H), 7.84-7.88 (m, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 21.59, 118.50, 126.73, 129.85, 131.24, 140.43, 143.74, 168.83; HRMS m/z (M^+) calcd for $\text{C}_{10}\text{H}_9\text{NS}$: 175.0456, found: 175.0454.

2-(2-Methylphenyl)benzoxazole (5c) m.p. 64-66 °C; ^1H NMR (400 MHz, CDCl_3) δ 2.84 (s, 3H), 7.35-7.45 (m, 5H), 7.59-7.63 (m, 1H), 7.80-7.85 (m, 1H), 8.18-8.21 (m, 1H); ^{13}C NMR (100 MHz, CDCl_3) δ 22.38, 110.68, 120.37, 124.57, 125.20, 126.26, 126.49, 130.16, 131.09, 131.99, 139.08, 142.39, 150.54, 163.63; HRMS m/z (M^+) calcd for $\text{C}_{14}\text{H}_{11}\text{NO}$: 209.0841, found: 209.0831.

2-(2,5-Dimethylphenyl)benzoxazole (5d) m.p. 40-42 °C; ^1H NMR (400 MHz, CDCl_3) δ 2.43 (s, 3H), 2.78 (s, 3H), 7.22-7.27 (m, 2H), 7.35-7.39 (m, 2H), 7.58-7.62 (m, 1H), 7.80-7.84 (m, 1H), 8.02 (s, 1H); ^{13}C NMR (100 MHz, CDCl_3) δ 21.04, 21.86, 110.65, 120.30, 124.54, 125.12, 126.21, 130.62, 131.95, 131.97, 135.82, 135.95, 142.36, 150.58, 163.92; HRMS m/z (M^+) calcd for $\text{C}_{15}\text{H}_{13}\text{NO}$: 223.0997, found: 223.0999.

2-(1-Naphthyl)benzoxazole (5e) m.p. 101-104 °C; ^1H NMR (400 MHz, CDCl_3) δ 7.40-7.44 (m, 2H), 7.59-7.69 (m, 3H), 7.71-7.75 (m, 1H), 7.89-7.93 (m, 1H), 7.96 (d, $J = 8.4$ Hz, 1H), 8.05 (d, $J = 8.1$ Hz, 1H), 8.45 (dd, $J = 7.3$ Hz, 1.1 Hz, 1H), 9.50 (d, $J = 8.4$ Hz, 1H); ^{13}C NMR (100 MHz, CDCl_3) δ 110.72, 120.53, 123.91, 124.71, 125.15, 125.49, 126.55, 126.68, 128.13, 128.88, 129.54, 130.97, 132.51, 134.23, 142.59, 150.45, 163.06; HRMS m/z (M^+) calcd for $\text{C}_{17}\text{H}_{11}\text{NO}$: 245.0841, found: 245.0838.

2-(4-Methoxy-2-methylphenyl)benzoxazole (5k) m.p. 73-75 °C; ^1H NMR (400 MHz, CDCl_3) δ 2.82 (s, 3H), 3.88 (s, 3H), 6.87-6.89 (m, 2H), 7.31-7.37 (m, 2H), 7.55-7.59 (m, 1H), 7.77-7.79 (m, 1H), 8.15 (dd, $J = 7.0$ Hz, 2.2 Hz, 1H); ^{13}C NMR (100 MHz, CDCl_3) δ 22.73, 55.50, 110.45, 111.78, 117.25, 119.14, 120.00, 124.39, 124.73, 131.90, 141.20, 142.55, 150.40, 161.70, 163.74; HRMS m/z (M^+) calcd for $\text{C}_{15}\text{H}_{13}\text{NO}_2$: 239.0946, found: 239.0951.

2-(9-Phenanthrenyl)benzoxazole (5l) m.p. 169-170 °C; ^1H NMR (400 MHz, CDCl_3) δ 7.41-7.46 (m, 2H), 7.65-7.72 (m, 2H), 7.75-7.82 (m, 3H), 7.91-7.95 (m, 1H), 8.04 (d, $J = 7.7$ Hz, 1H), 8.73 (d, $J = 8.1$ Hz, 2H), 8.80 (d, $J = 8.6$ Hz, 1H), 9.55-9.57 (m, 1H); ^{13}C NMR (100 MHz, CDCl_3) δ 110.71, 110.72, 120.56, 122.78, 122.92, 123.15, 124.76, 125.60, 127.34, 127.36, 127.83, 128.88, 128.95, 130.01, 130.77, 131.03, 131.82, 131.97, 142.54, 150.43, 162.94; HRMS m/z (M^+) calcd for $\text{C}_{21}\text{H}_{13}\text{NO}$: 295.0997, found: 295.1007.

2-(2-Biphenyl)benzoxazole (5m) oil; ^1H NMR (400 MHz, CDCl_3) δ 7.23-7.34 (m, 8H), 7.47-7.53 (m, 2H), 7.56-7.61 (m, 1H), 7.69-7.73 (m, 1H), 8.11 (d, $J = 7.5$ Hz, 1H); ^{13}C NMR (100 MHz, CDCl_3) δ 110.72, 120.30, 124.50, 125.10, 126.55, 127.47, 127.73, 128.31, 129.02, 131.17, 131.18,

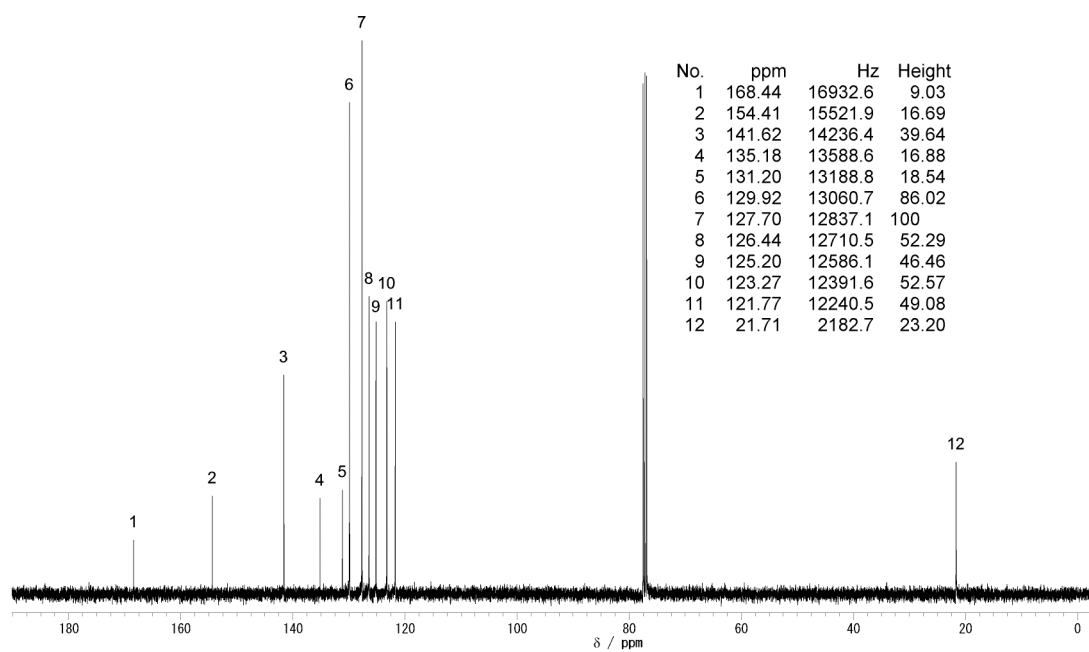
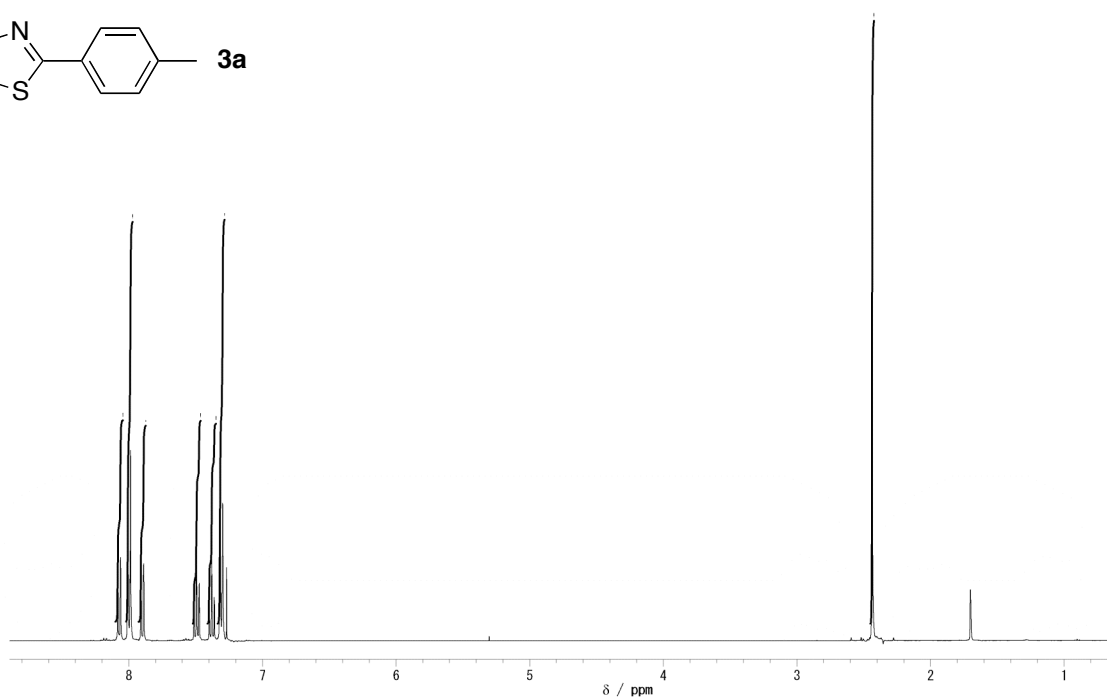
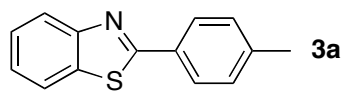
131.35, 141.22, 141.95, 142.70, 150.97, 164.08; HRMS m/z (M^+) calcd for $C_{19}H_{13}NO$: 271.0997, found: 271.0990.

2-(9-Anthracenyl)benzoxazole (5n) m.p. 164-166 °C; 1H NMR (400 MHz, $CDCl_3$) δ 7.48-7.56 (m, 6H), 7.71-7.75 (m, 1H), 7.99-8.03 (m, 1H), 8.07-8.12 (m, 4H), 8.68 (s, 1H); ^{13}C NMR (100 MHz, $CDCl_3$) δ 111.14, 120.79, 121.58, 124.88, 125.68, 125.75, 125.79, 127.58, 128.90, 131.02, 131.34, 131.61, 142.12, 151.40, 162.29; HRMS m/z (M^+) calcd for $C_{21}H_{13}NO$: 295.0997, found: 295.1007.

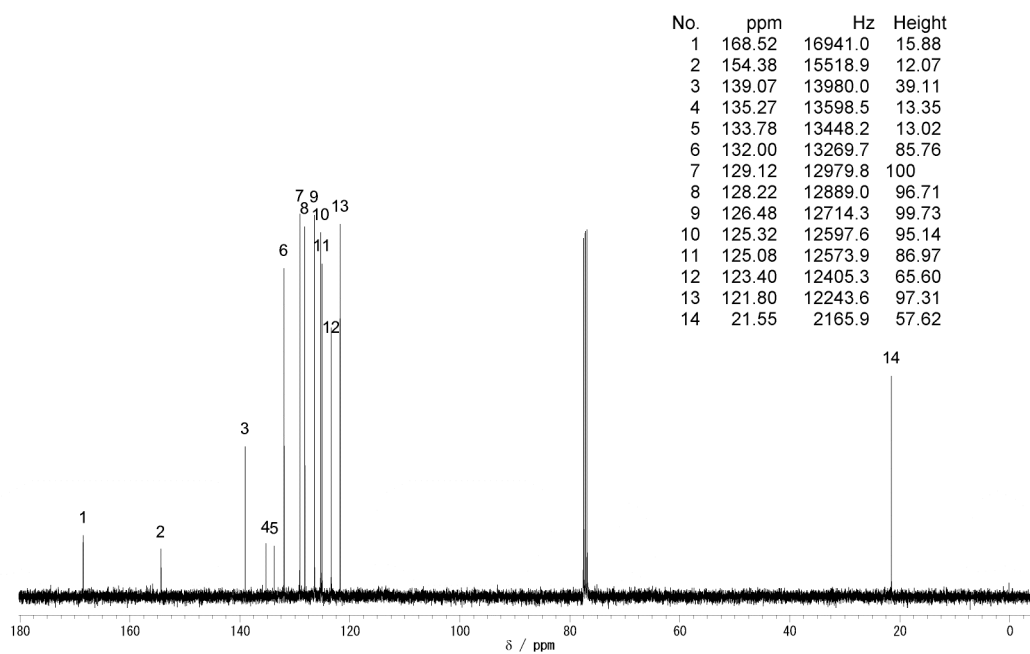
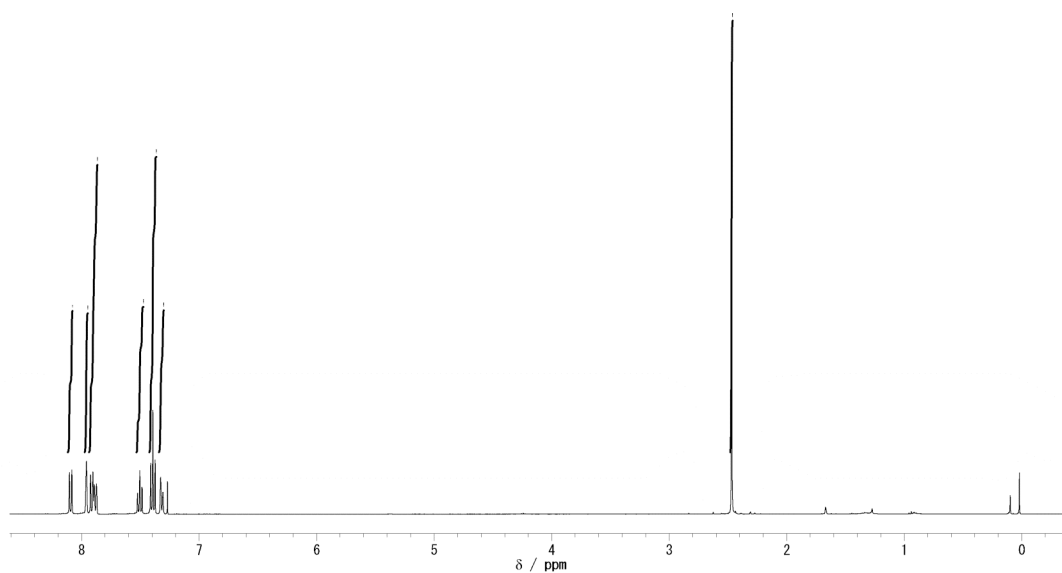
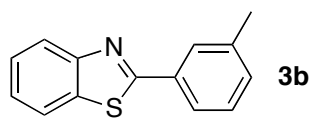
2-(9-Phenanthrenyl)-5-phenyloxazole (7al) m.p. 180-182 °C; 1H NMR (400 MHz, $CDCl_3$) δ 7.40 (t, $J = 7.3$ Hz, 1H), 7.51 (t, $J = 7.3$ Hz, 2H), 7.63 (s, 1H), 7.64-7.68 (m, 1H), 7.72-7.79 (m, 3H), 7.82-7.85 (m, 2H), 8.01-8.03 (dd, $J = 7.2$ Hz, 1.1 Hz, 1H), 8.57 (s, 1H), 8.71 (d, $J = 8.4$ Hz, 1H), 8.77-8.80 (m, 1H), 9.41-9.45 (m, 1H); ^{13}C NMR (100 MHz, $CDCl_3$) δ 122.87, 123.06, 123.11, 123.65, 124.63, 127.19, 127.23, 127.25, 127.65, 128.26, 128.35, 128.78, 128.80, 129.22, 129.73, 129.86, 130.95, 131.02, 131.51, 151.32, 161.16; HRMS m/z (M^+) calcd for $C_{23}H_{15}NO$: 321.1154, found: 321.1151.

2-(9-Phenanthrenyl)-5-(4-trifluoromethylphenyl)oxazole (7bl) m.p. 165-167 °C; 1H NMR (400 MHz, $CDCl_3$) δ 7.64-7.78 (m, 7H), 7.87 (d, $J = 8.4$ Hz, 2H), 8.01 (d, $J = 7.7$ Hz, 1H), 8.54 (s, 1H), 8.70 (d, $J = 8.1$ Hz, 1H), 8.75-8.78 (m, 1H), 9.37-9.41 (m, 1H); ^{13}C NMR (100 MHz, $CDCl_3$) δ 122.65, 122.90, 123.16, 124.19 (q, $J = 271.6$ Hz), 124.60, 125.35, 126.23 (q, $J = 3.8$ Hz), 127.10, 127.31, 127.34, 127.72, 128.60, 128.66, 129.78, 130.20, 130.40 (q, $J = 32.8$ Hz), 130.81, 131.02, 131.39, 131.64, 149.82, 161.96; HRMS m/z (M^+) calcd for $C_{24}H_{14}F_3NO$: 389.1027, found: 389.1022.

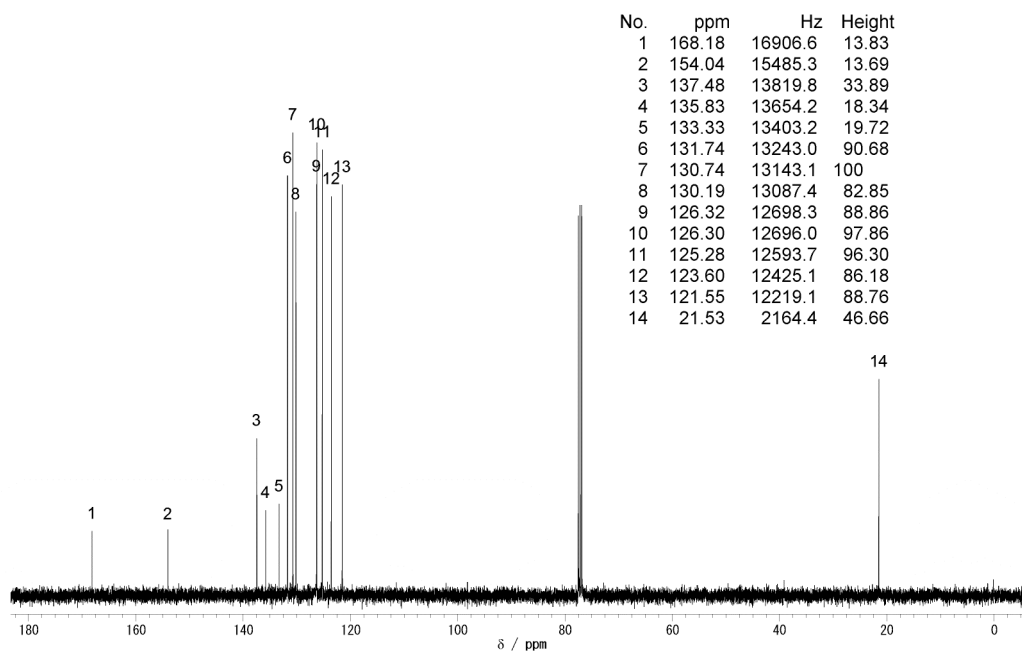
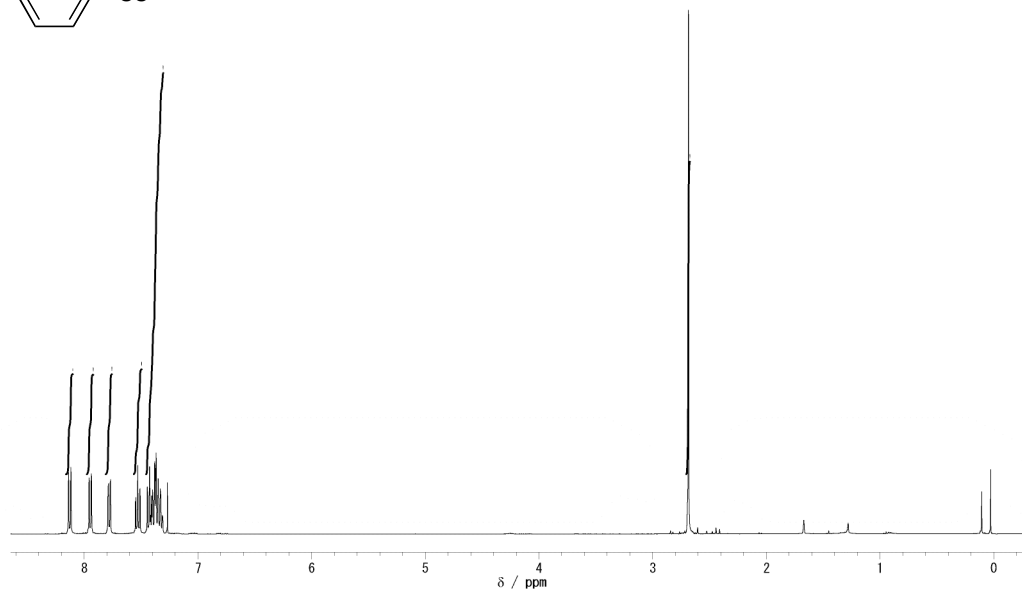
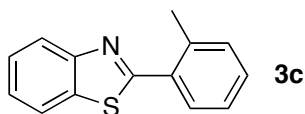
[¹H and ¹³C NMR Spectra of **3a**]



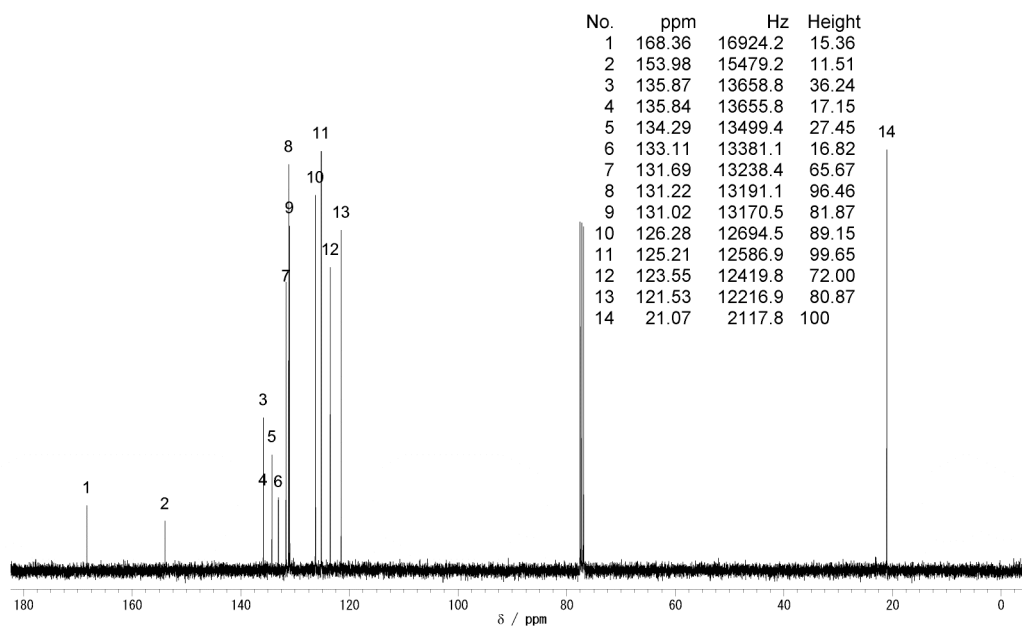
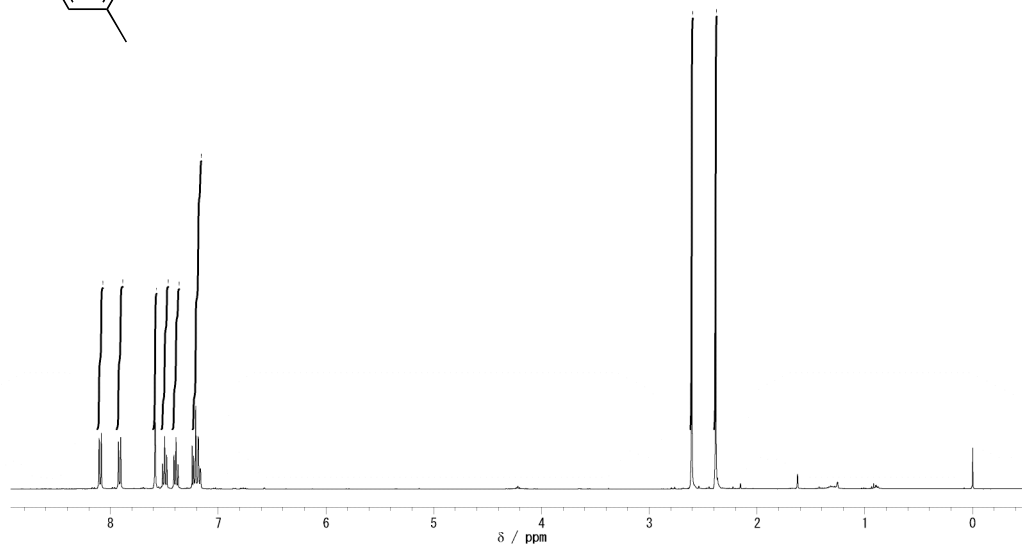
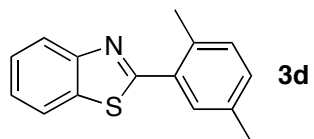
[¹H and ¹³C NMR Spectra of **3b**]



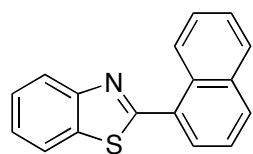
[¹H and ¹³C NMR Spectra of **3c**]



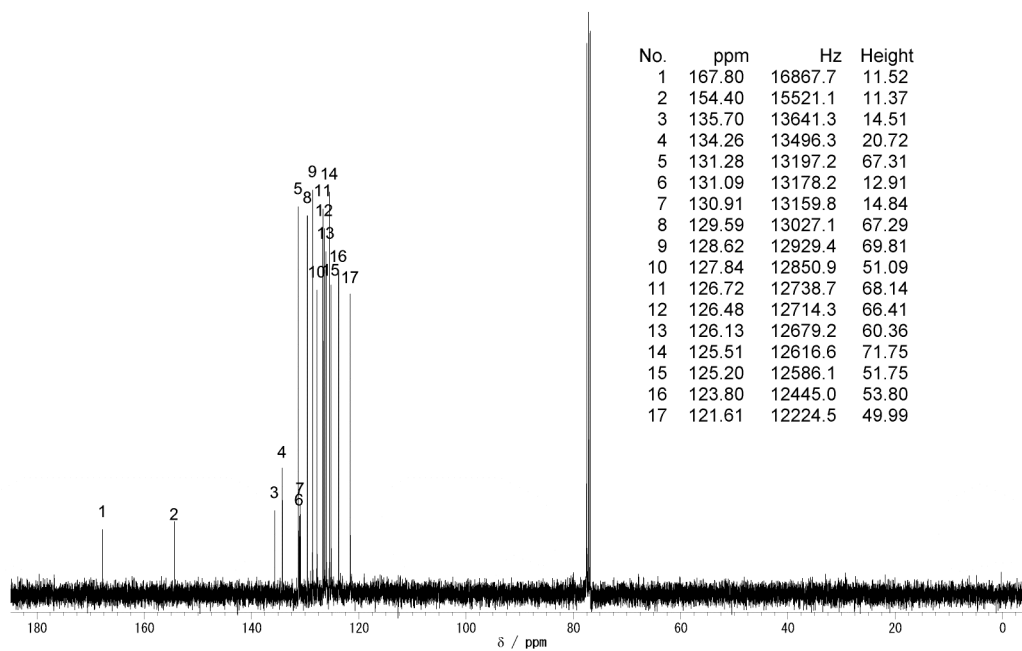
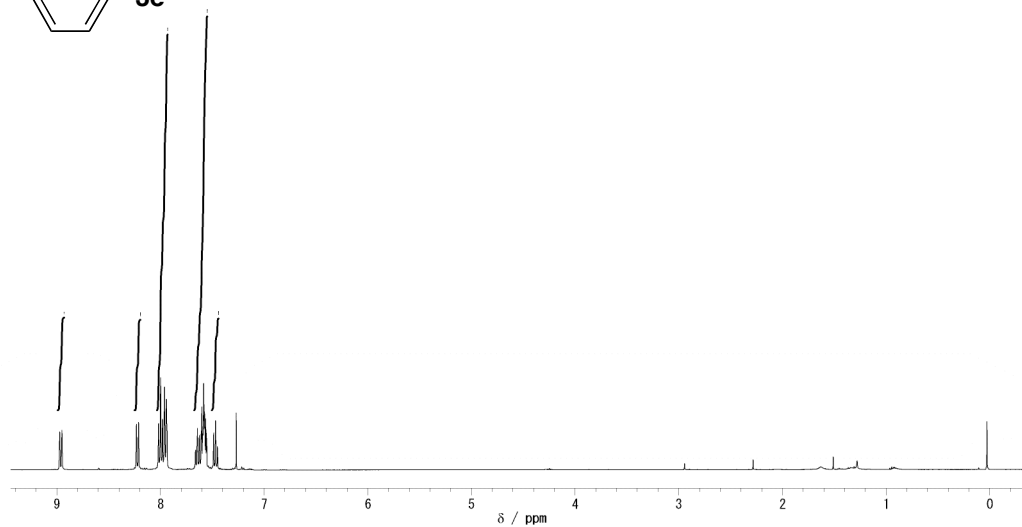
[¹H and ¹³C NMR Spectra of **3d**]



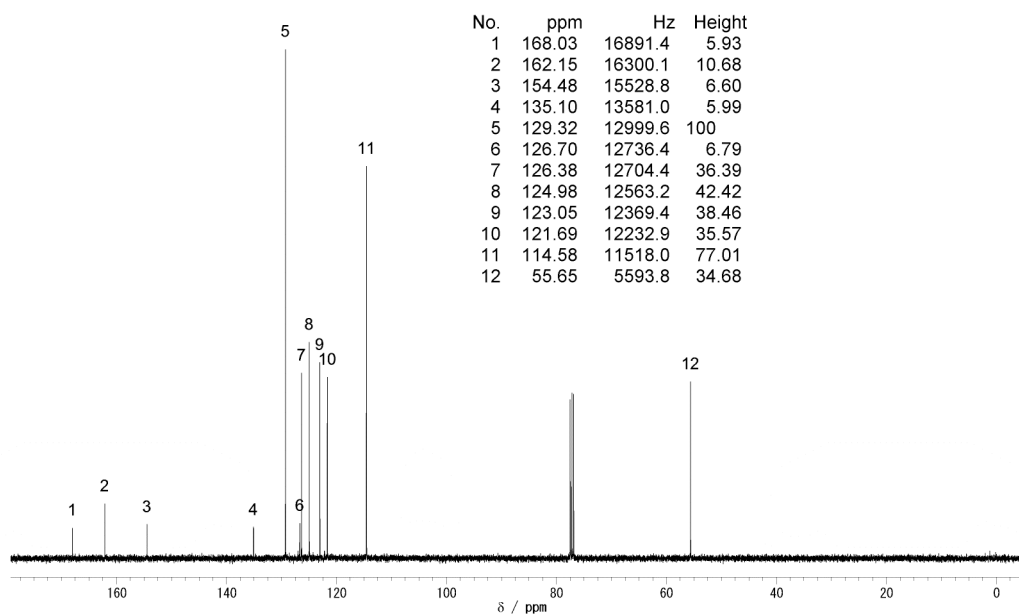
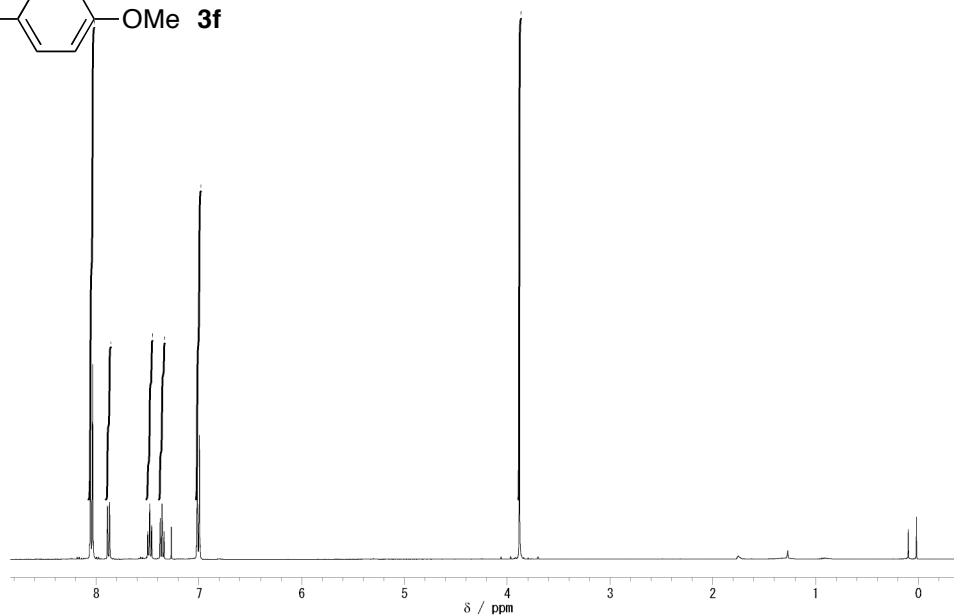
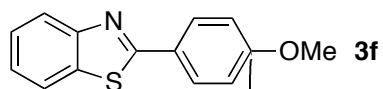
[¹H and ¹³C NMR Spectra of **3e**]



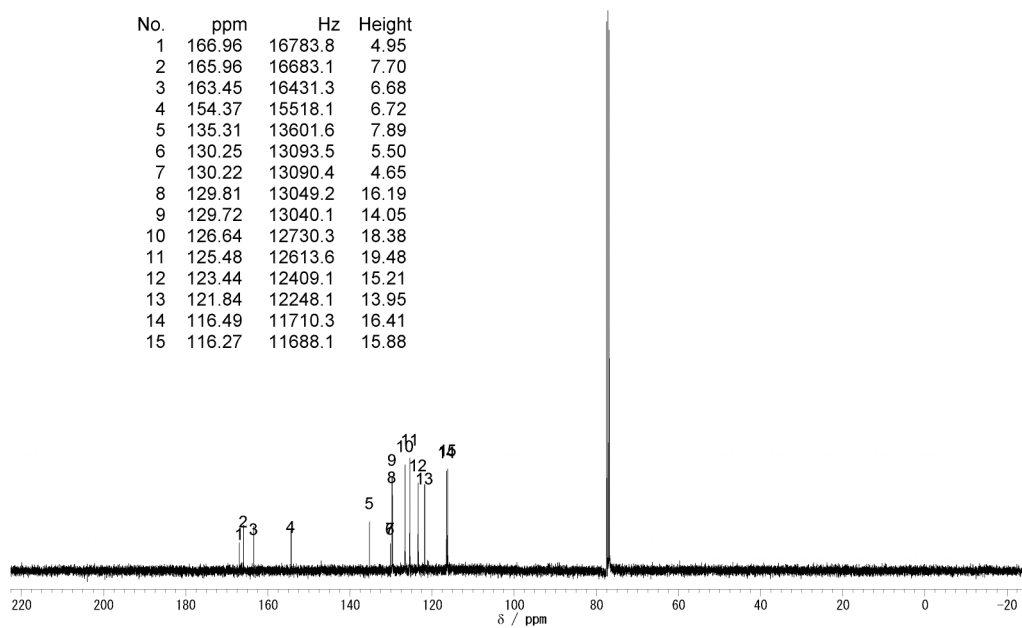
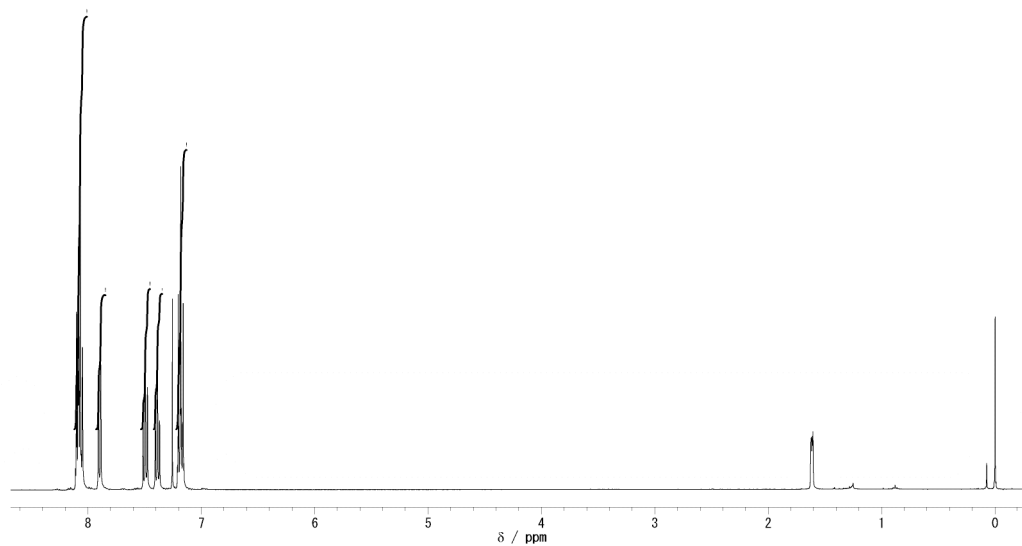
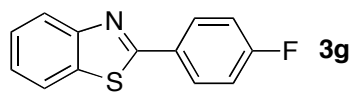
3e



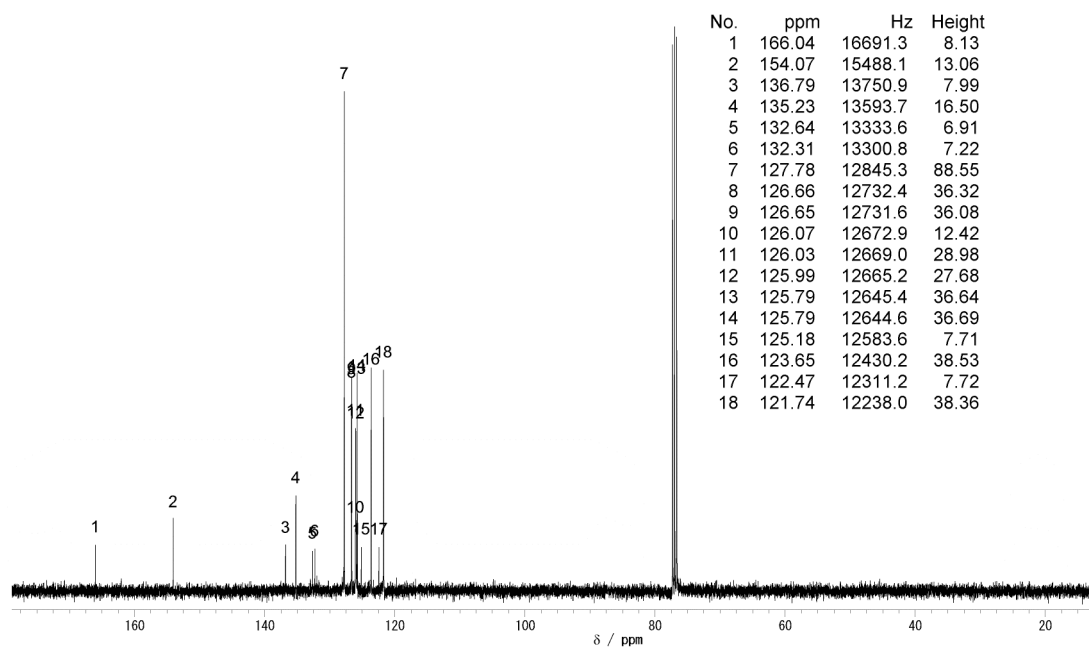
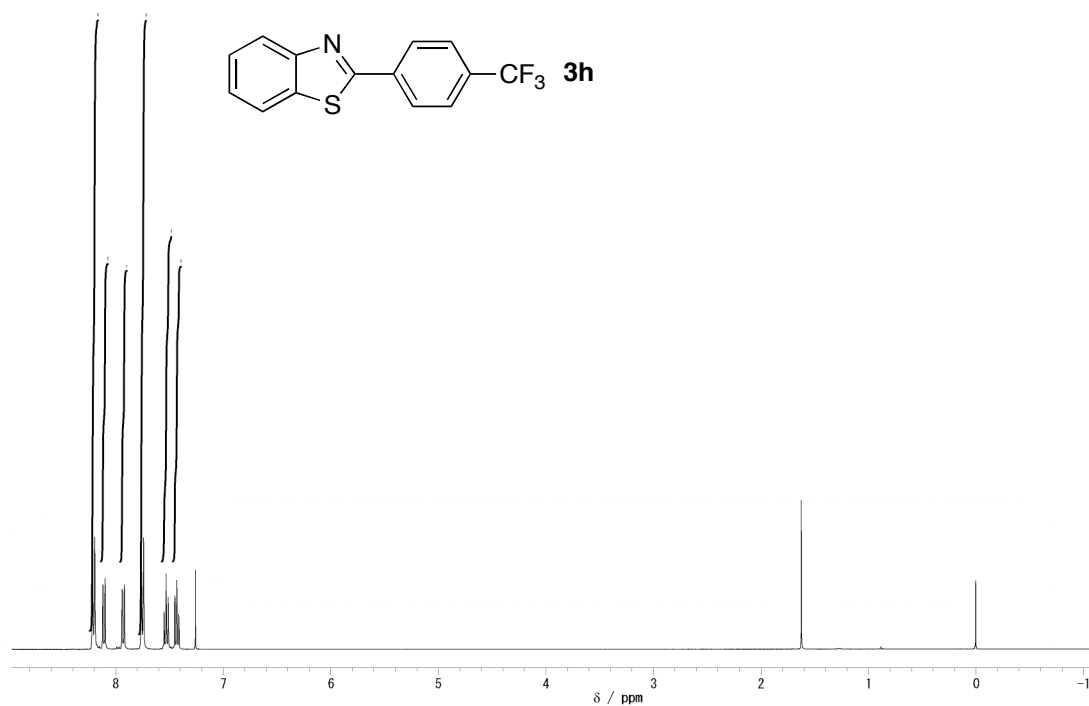
[¹H and ¹³C NMR Spectra of **3f**]



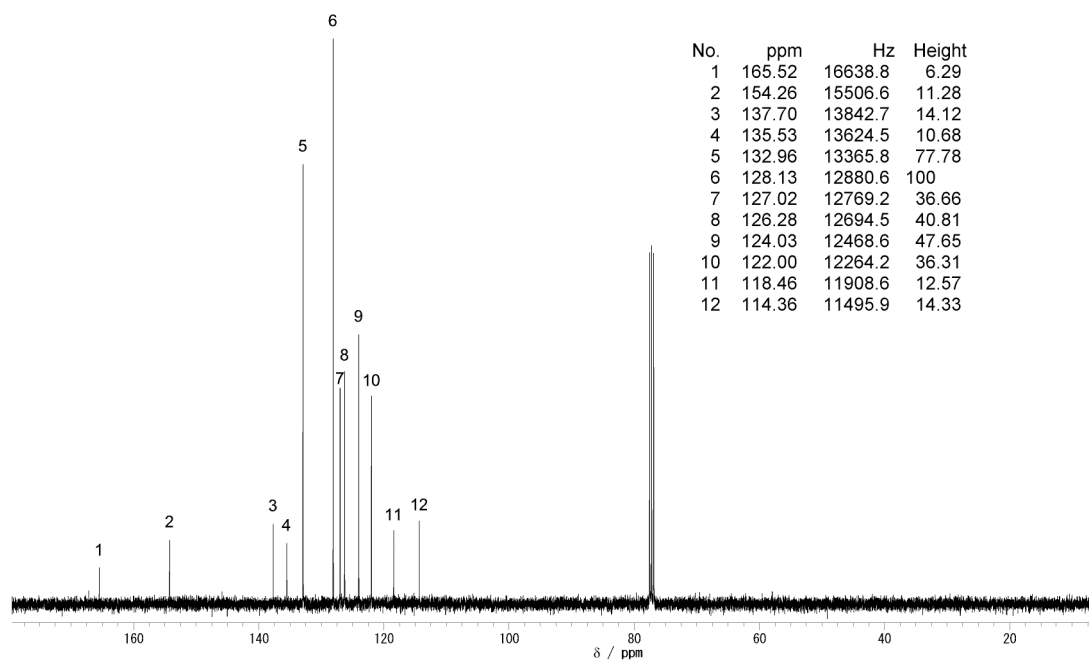
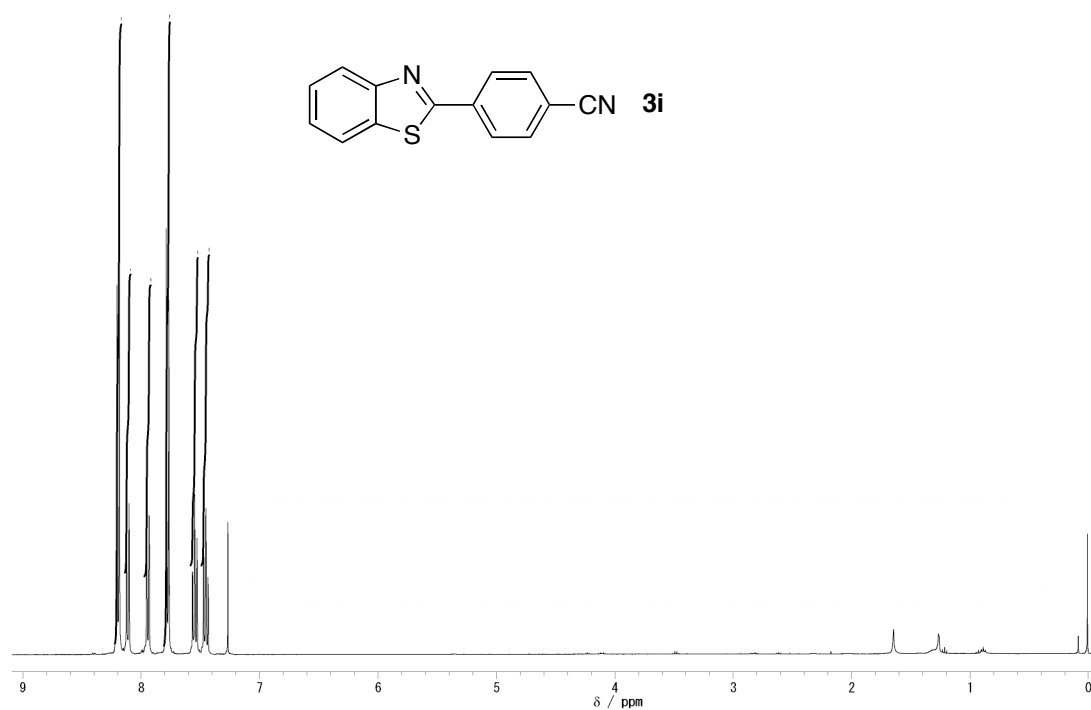
[^1H and ^{13}C NMR Spectra of **3g**]



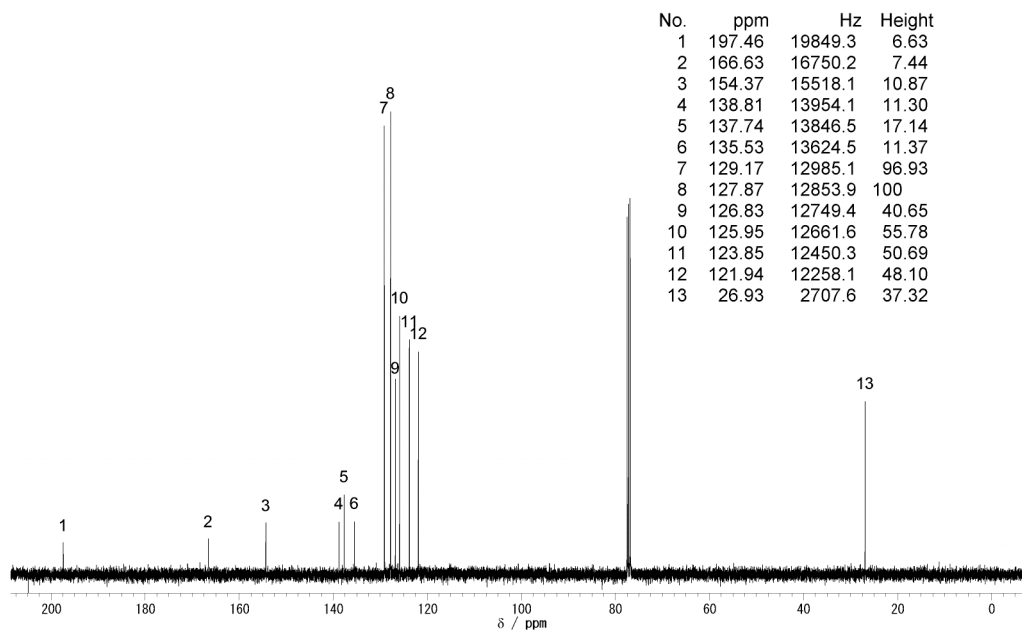
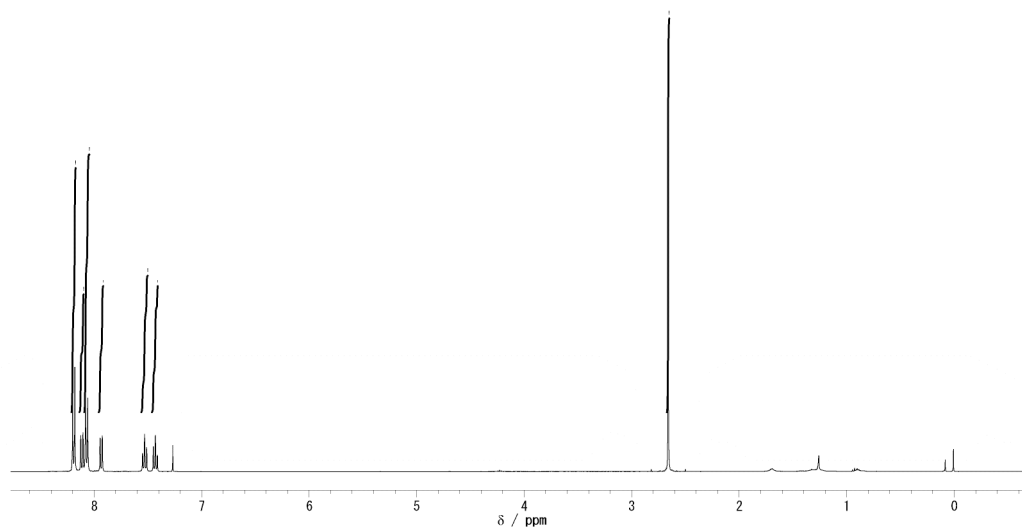
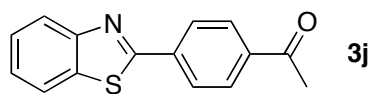
[¹H and ¹³C NMR Spectra of **3h**]



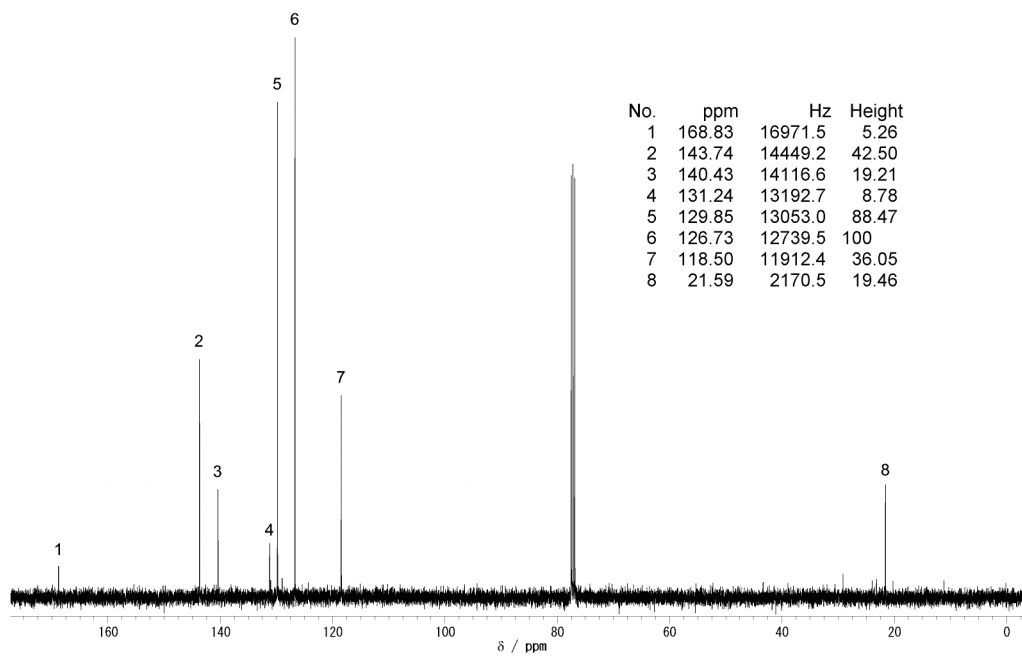
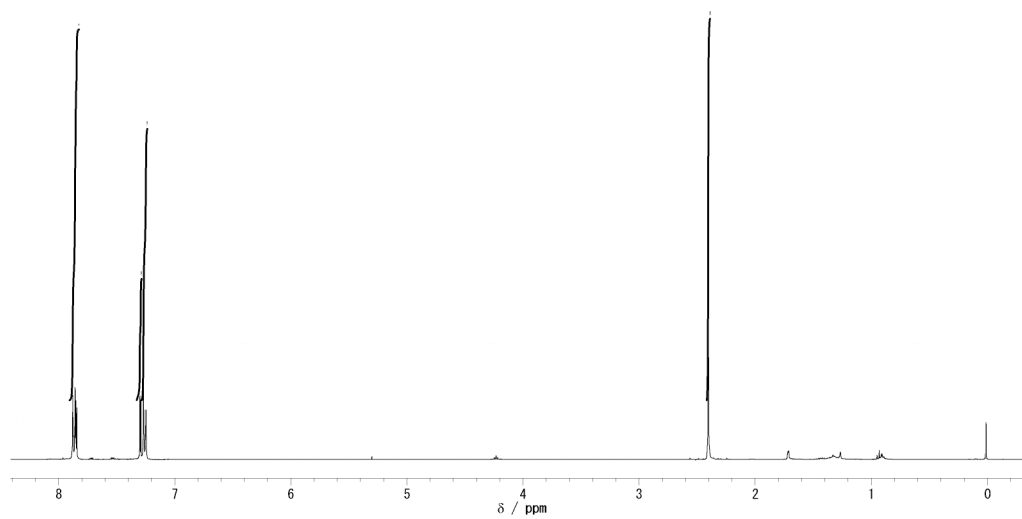
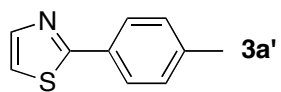
[¹H and ¹³C NMR Spectra of **3i**]



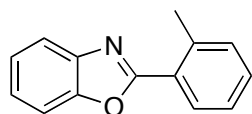
[¹H and ¹³C NMR Spectra of **3j**]



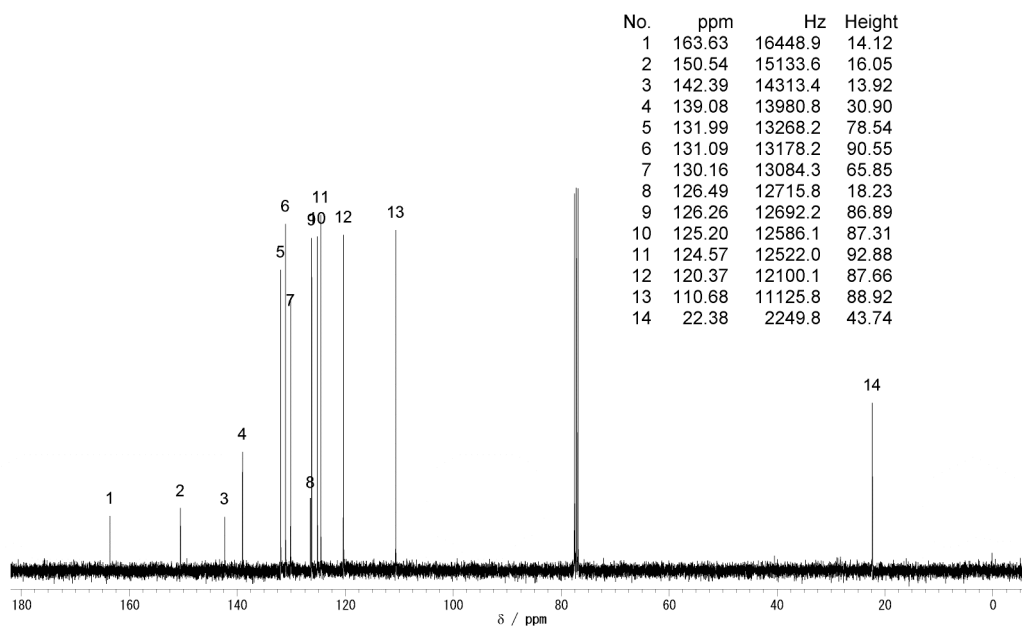
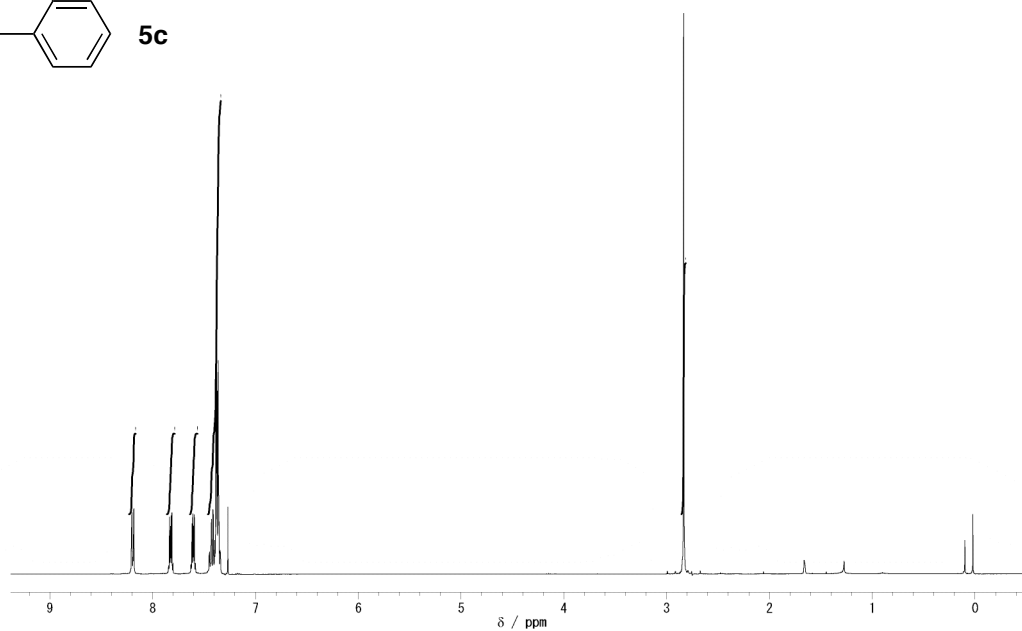
[^1H and ^{13}C NMR Spectra of **3a'**]



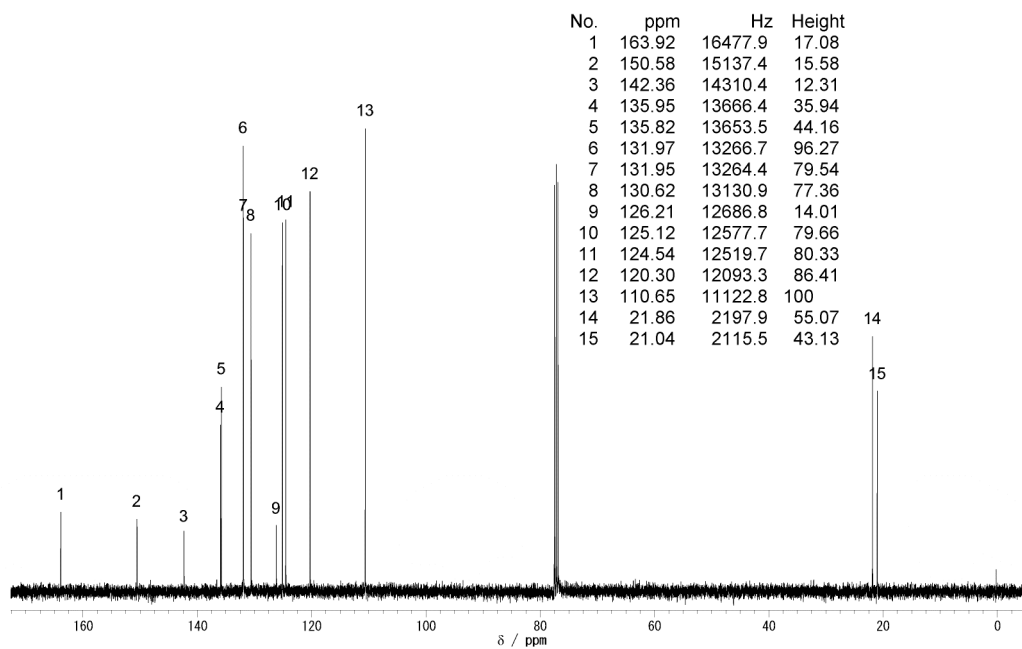
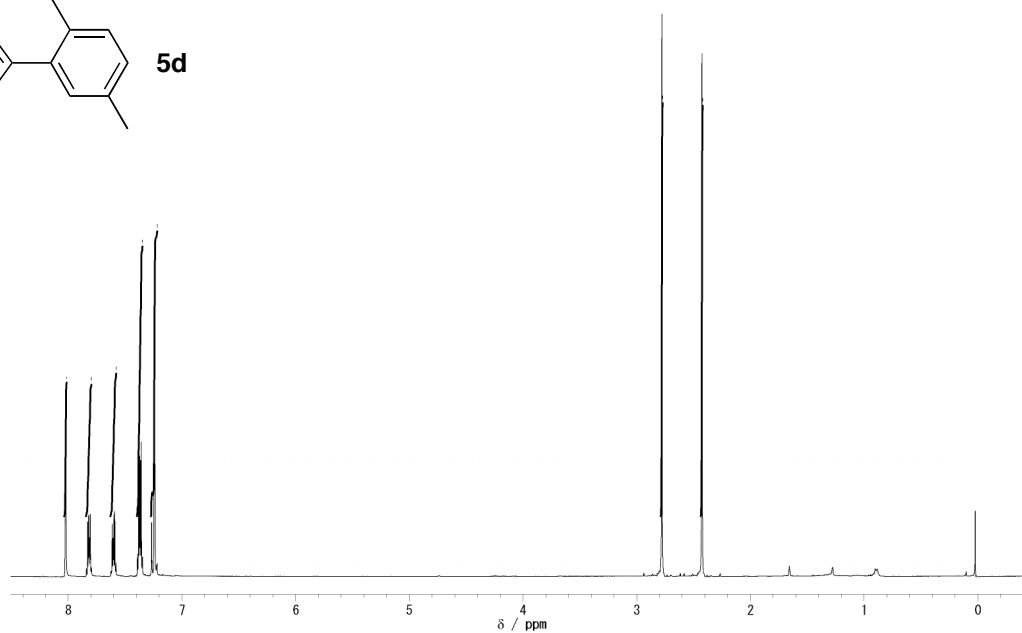
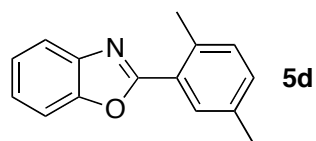
[¹H and ¹³C NMR Spectra of **5c**]



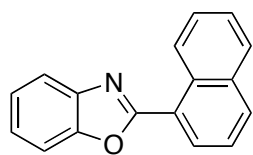
5c



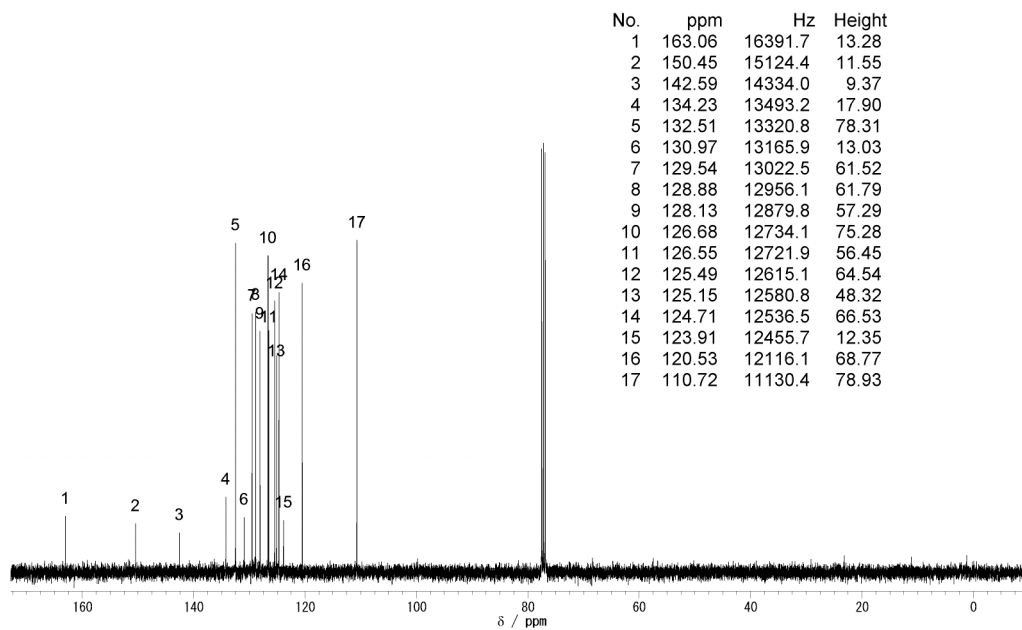
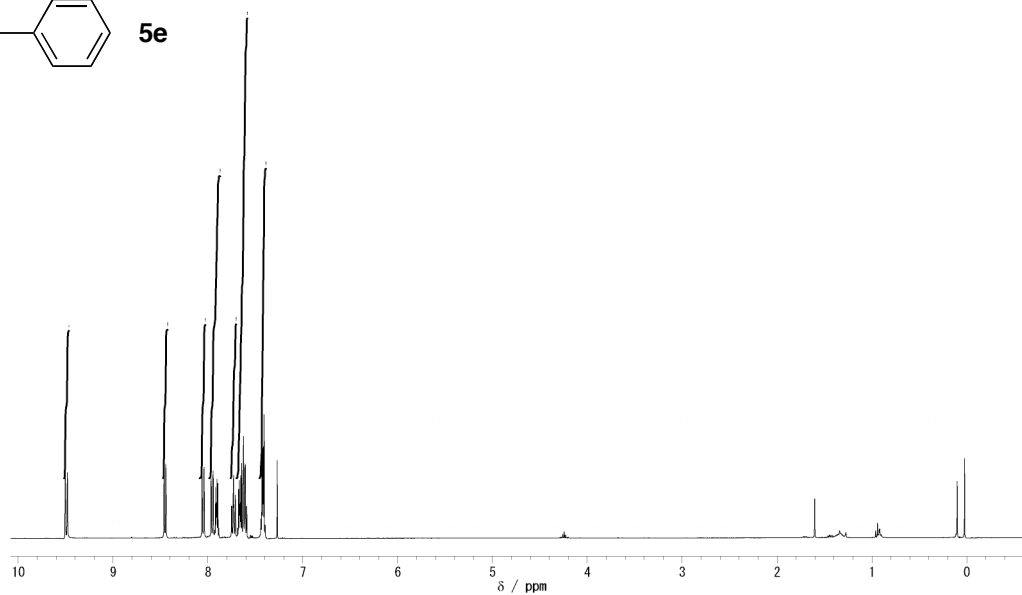
[¹H and ¹³C NMR Spectra of **5d**]



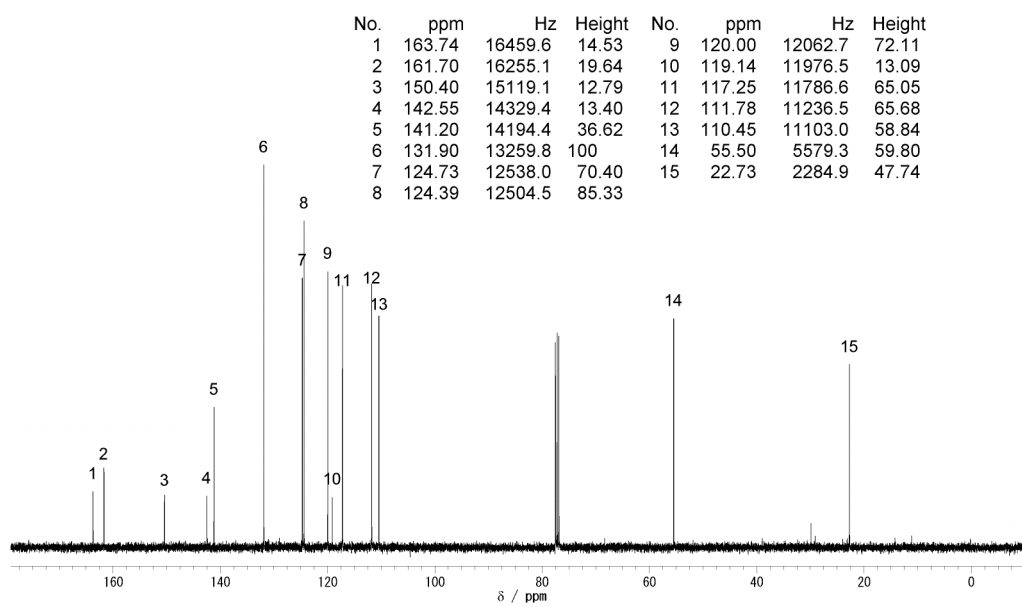
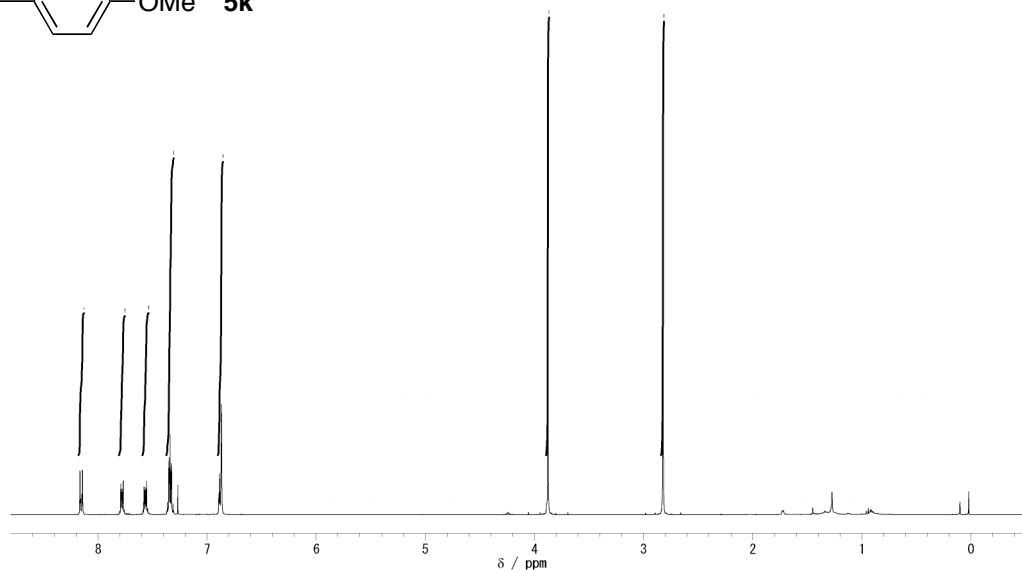
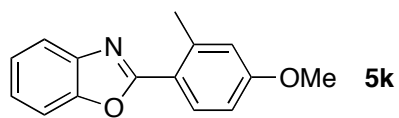
[¹H and ¹³C NMR Spectra of **5e**]



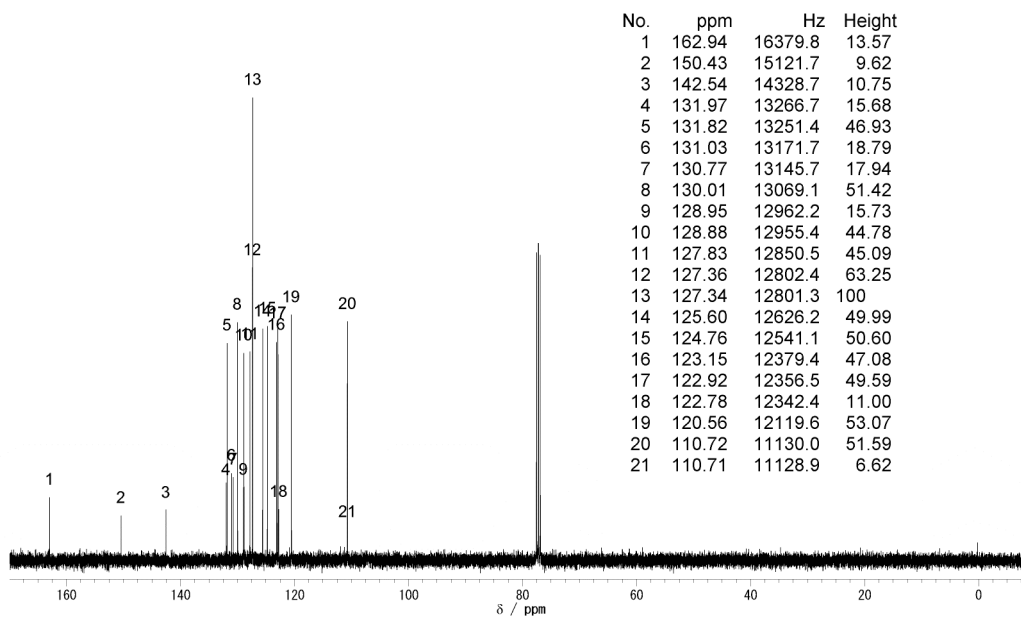
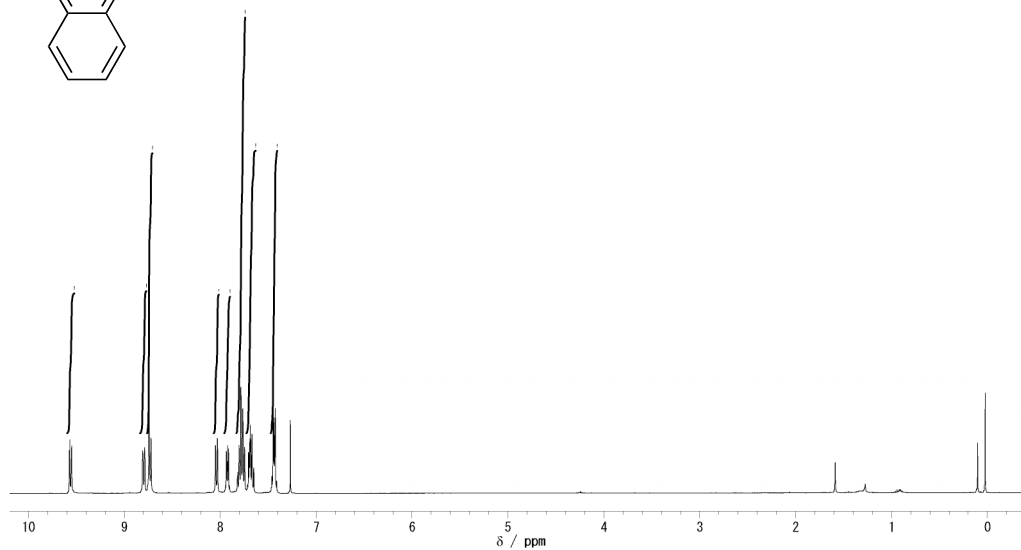
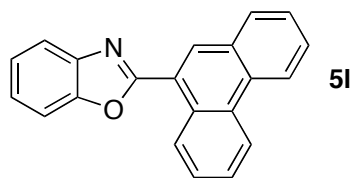
5e



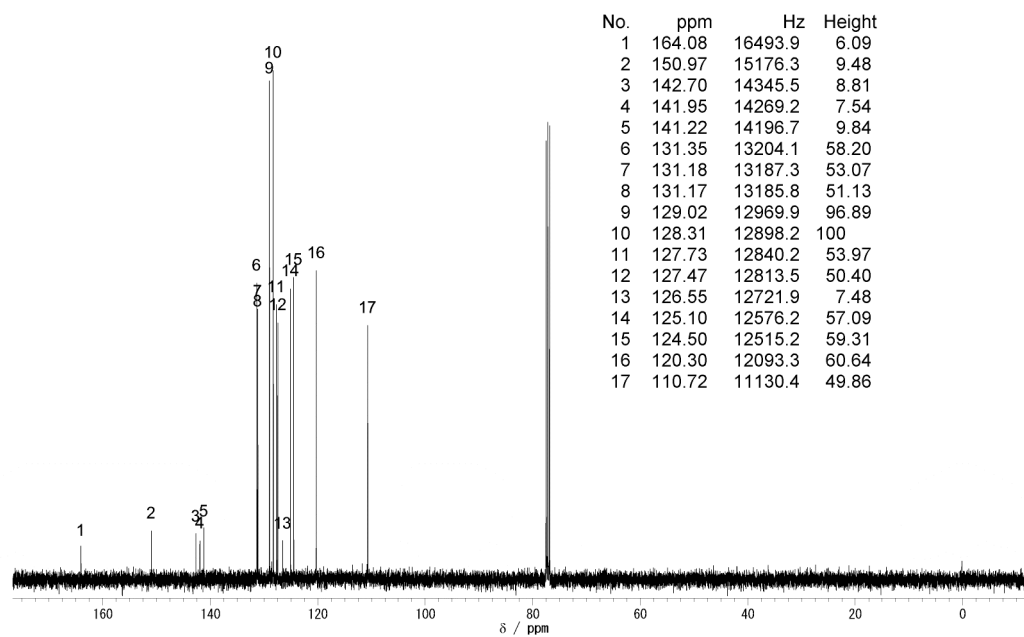
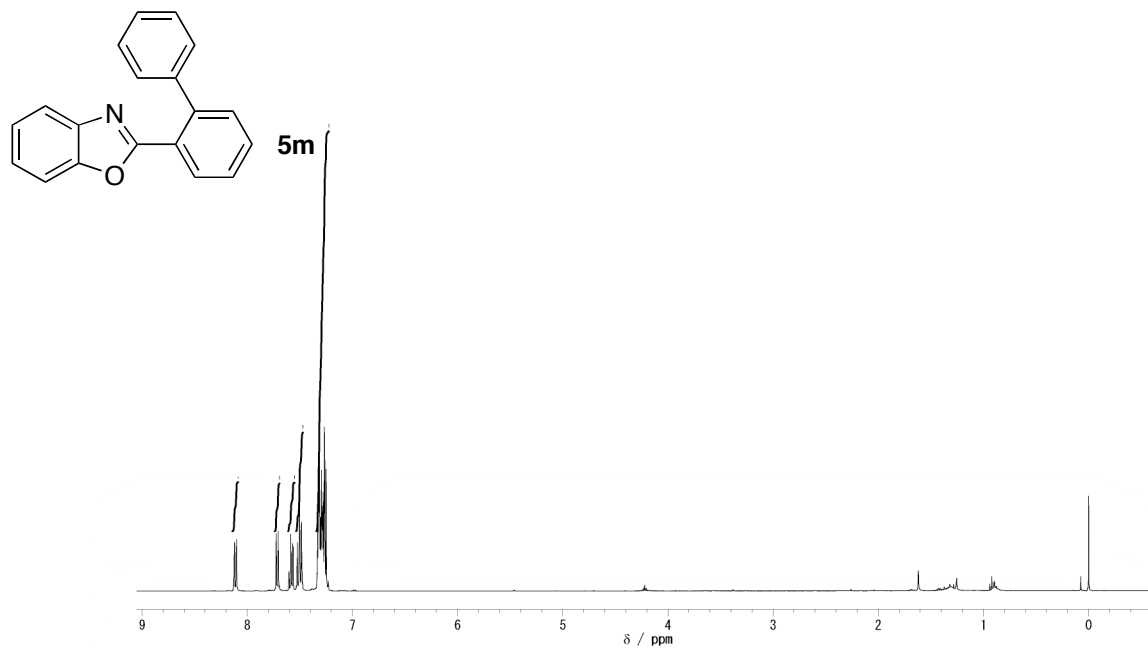
[^1H and ^{13}C NMR Spectra of **5k**]



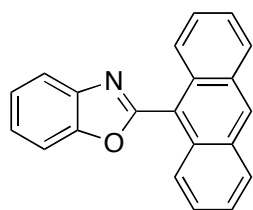
[^1H and ^{13}C NMR Spectra of **51**]



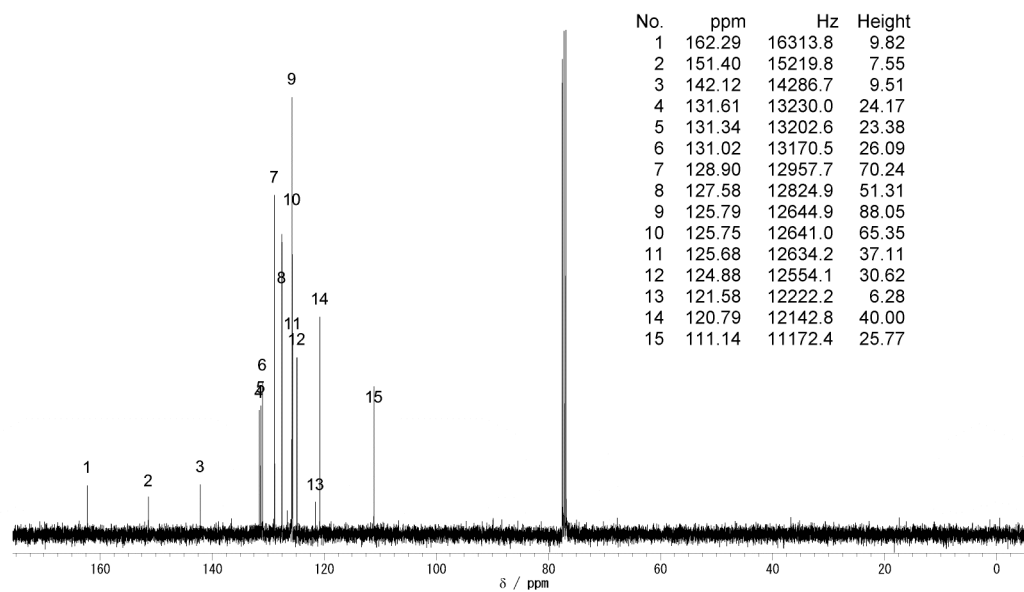
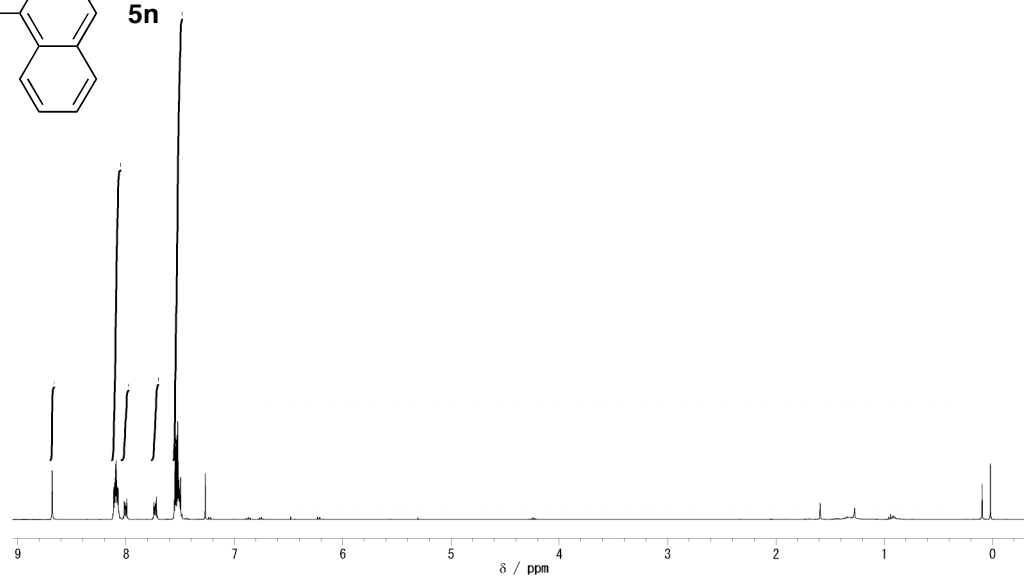
[¹H and ¹³C NMR Spectra of **5m**]



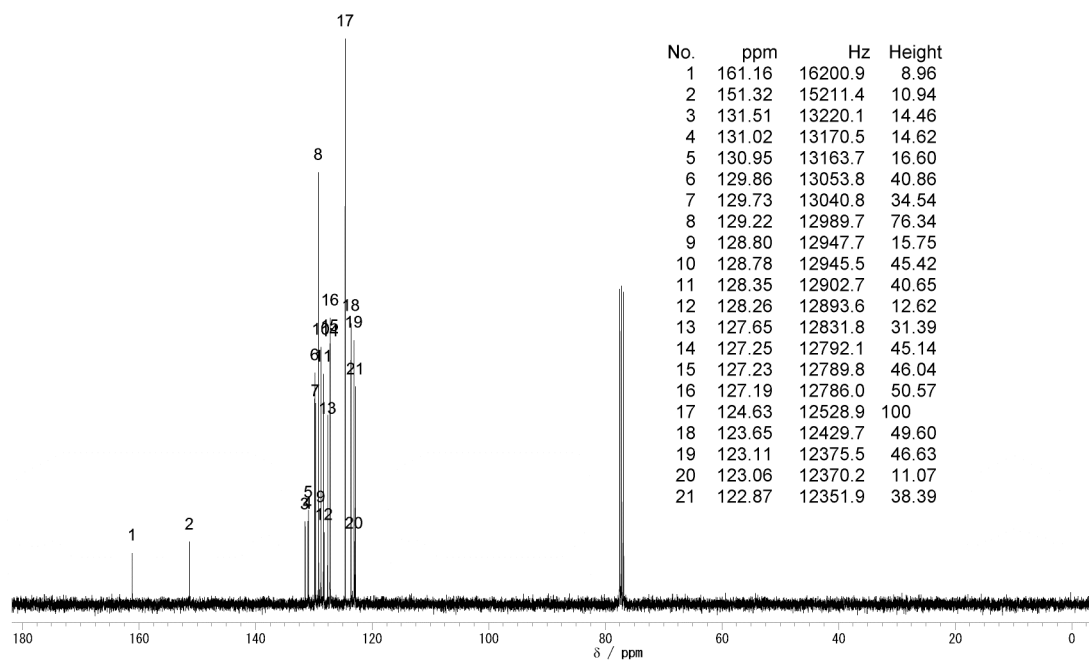
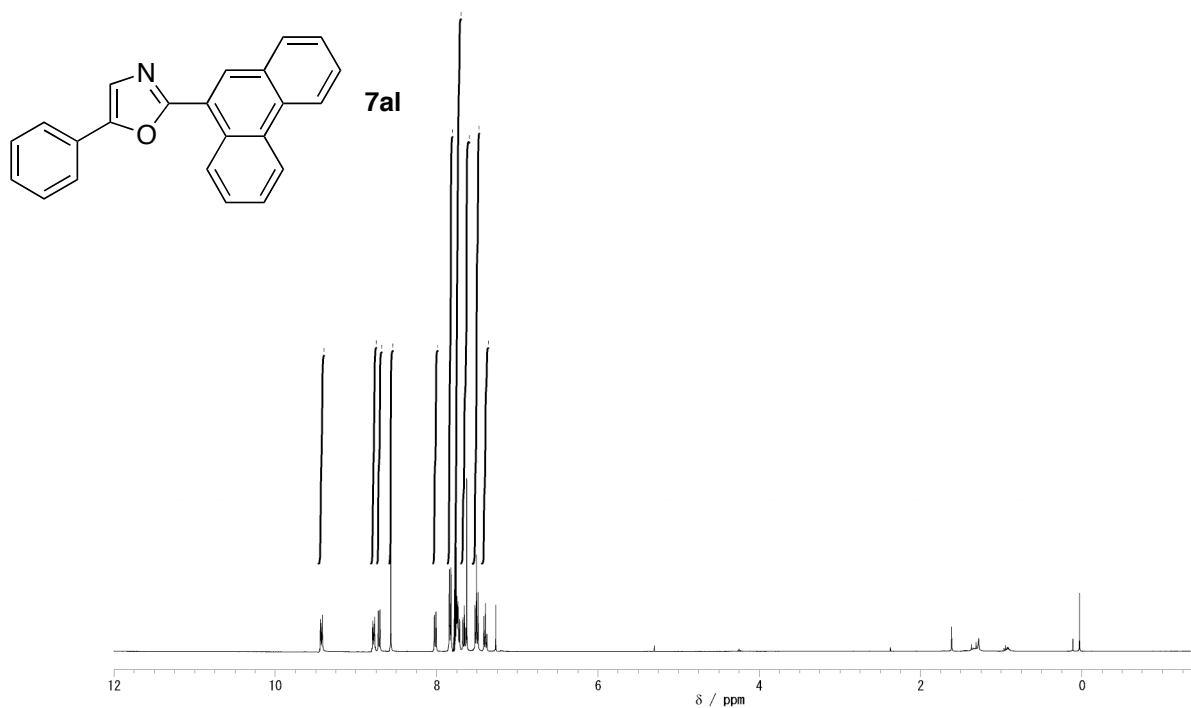
[¹H and ¹³C NMR Spectra of **5n**]



5n



[¹H and ¹³C NMR Spectra of **7al**]



[¹H and ¹³C NMR Spectra of **7bl**]

