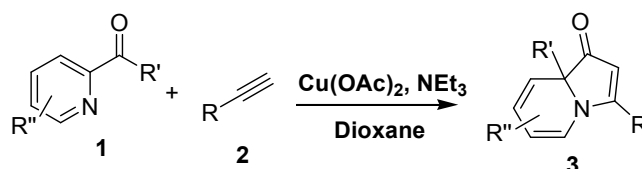


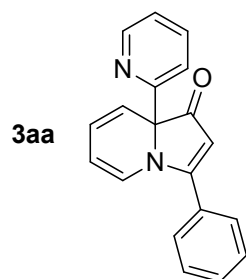
General Methods. Unless otherwise noted, materials obtained from commercial suppliers were used without further purification. ^1H spectra were recorded on 300 MHz or 400 MHz NMR spectrometers. Chemical shifts (in ppm) were referenced to tetramethylsilane ($\delta = 0$ ppm) in CDCl_3 as an internal standard. ^{13}C -NMR spectra were obtained by using the same NMR spectrometers and were calibrated with CDCl_3 ($\delta = 77.00$ ppm). Single crystal X-ray diffraction data was collected on Bruker SMART diffractometer at 273(2) K with graphite-monochromated Mo-K α radiation ($\lambda = 0.71073$ Å).

Experimental procedures and characterization datas of compounds

Typical procedure for the synthesis of indolizinone 3aa-3da.

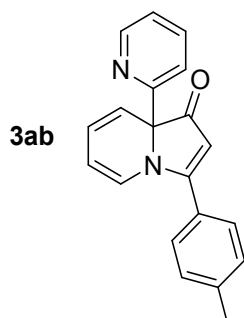


To a screw-cap vial containing a stir bar, **1** (0.3 mmol), **2** (0.6 mmol), $\text{Cu}(\text{OAc})_2$ (0.03 mmol) and NEt_3 (0.6 mmol) were added in 1,4-dioxane (2 mL). The reaction vial was fitted with a cap. The reaction vial was heated with stirring at 110 °C or 130 °C for several hours. After cooling down to room temperature and concentrated in vacuum, the residue was purified by flash chromatography on a short silica gel to provide the terminal product **3**.

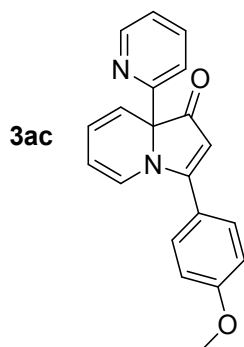


3-phenyl-8a-(pyridin-2-yl)-indolizinone (3aa). **1a** (55.2 mg, 0.3 mmol), **2a** (61.2 mg, 0.6 mmol), $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol) and NEt_3 (60.6 mg, 0.6 mmol) were added in 1,4-dioxane (2 mL). The reaction vial was heated with stirring at 110 °C for 4h. **3aa** (82 mg, 95%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 173-174 °C; IR (KBr) ν 3036, 1566, 1487, 1389, 1302 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.60 (d, $J = 4.3$ Hz,

1H), 7.68 (t, $J = 4.8$ Hz, 3H), 7.58 – 7.46 (m, 4H), 7.19 (dd, $J = 7.0, 5.1$ Hz, 1H), 6.73 (d, $J = 7.2$ Hz, 1H), 6.45 (d, $J = 9.2$ Hz, 1H), 6.12 (dd, $J = 9.2, 5.5$ Hz, 1H), 5.39 – 5.29 (m, 1H), 5.21 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.2, 175.2, 158.7, 149.5, 136.8, 131.2, 129.5, 129.1, 128.2, 124.6, 123.5, 122.7, 122.0, 120.2, 108.8, 99.5, 73.8; HRMS (ESI): Exact mass calcd for $\text{C}_{19}\text{H}_{14}\text{N}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 309.0996, found 309.0998.

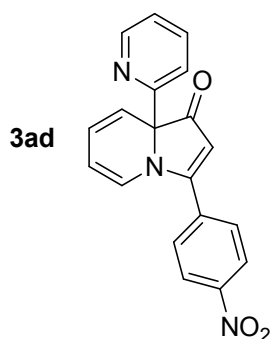


3-(p-tolyl)-8a-(pyridin-2-yl)-indolizinone (3ab). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2b** (69.6 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 4 h. **3ab** (90 mg, 100%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 185-186 °C; IR (KBr) ν 3041, 1675, 1428, 1385, 1300 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.59 (d, $J = 4.7$ Hz, 1H), 7.66 (td, $J = 7.9, 1.7$ Hz, 1H), 7.55 (d, $J = 8.1$ Hz, 2H), 7.49 (d, $J = 8.0$ Hz, 1H), 7.34 (d, $J = 8.0$ Hz, 2H), 7.20 – 7.13 (m, 1H), 6.74 (d, $J = 7.3$ Hz, 1H), 6.44 (d, $J = 9.2$ Hz, 1H), 6.10 (dd, $J = 9.2, 5.5$ Hz, 1H), 5.37 – 5.29 (m, 1H), 5.18 (s, 1H), 2.44 (s, 3H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.1, 175.4, 158.8, 149.5, 141.7, 136.7, 129.7, 128.1, 126.6, 124.7, 123.4, 122.7, 122.1, 120.1, 108.7, 99.2, 73.8, 21.7; HRMS (ESI): Exact mass calcd for $\text{C}_{20}\text{H}_{16}\text{N}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 323.1152, found 323.1155.

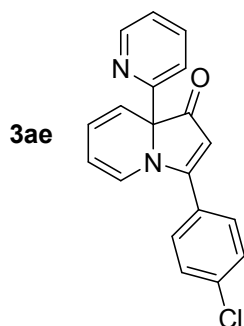


3-(4-methoxyphenyl)-8a-(pyridin-2-yl)-indolizinone (3ac). The reaction of $\text{Cu}(\text{OAc})_2$ (5.4 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol),

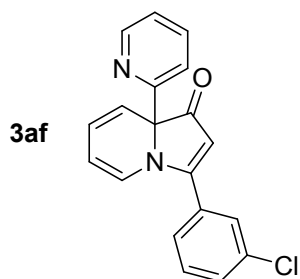
2c (79.2 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 24 h. **3ac** (81 mg, 85%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 175-176 °C; IR (KBr) ν 3023, 1675, 1496, 1388, 1256 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.58 (d, J = 4.1 Hz, 1H), 7.70 – 7.54 (m, 3H), 7.48 (d, J = 8.0 Hz, 1H), 7.17 (dd, J = 6.7, 5.0 Hz, 1H), 7.04 (d, J = 8.8 Hz, 2H), 6.77 (d, J = 7.2 Hz, 1H), 6.44 (d, J = 9.2 Hz, 1H), 6.10 (dd, J = 9.2, 5.4 Hz, 1H), 5.39 – 5.27 (m, 1H), 5.16 (s, 1H), 3.88 (s, 3H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.0, 175.1, 161.93, 158.9, 149.5, 136.7, 129.9, 124.8, 123.4, 122.6, 122.2, 121.7, 120.2, 114.5, 108.7, 98.9, 73.8, 55.5.; HRMS (ESI): Exact mass calcd for $\text{C}_{20}\text{H}_{16}\text{N}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 339.1101, found 339.1104.



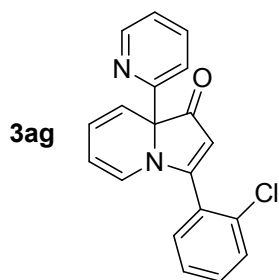
3-(4-nitrophenyl)-8a-(pyridin-2-yl)-indolizinone (3ad). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2d** (88.2 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 24 h. **3ad** (77.5 mg, 78%, n-hexane/ethyl acetate = 3:1): red solid; mp 206-207 °C; IR (KBr) ν 3104, 1683, 1521, 1349, 1289 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.57 (d, J = 4.3 Hz, 1H), 8.47 – 8.35 (m, 2H), 7.89 (d, J = 8.7 Hz, 2H), 7.69 (dd, J = 7.7, 1.4 Hz, 1H), 7.53 (d, J = 8.0 Hz, 1H), 7.26 (s, 1H), 7.22 (dd, J = 7.0, 5.3 Hz, 1H), 6.59 (d, J = 7.2 Hz, 1H), 6.42 (d, J = 9.2 Hz, 1H), 6.15 (dd, J = 9.2, 5.5 Hz, 1H), 5.44 – 5.33 (m, 1H), 5.27 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.1, 172.3, 158.3, 149.3, 137.0, 135.7, 129.4, 124.3, 123.9, 123.8, 123.0, 121.7, 120.9, 109.4, 100.6, 73.9.; HRMS (ESI): Exact mass calcd for $\text{C}_{19}\text{H}_{13}\text{N}_3\text{O}_3^+ [\text{M}+\text{Na}]^+$ 354.0842, found 354.0849.



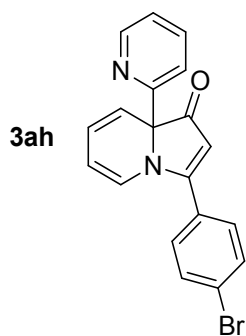
3-(4-chlorophenyl)-8a-(pyridin-2-yl)-indolizinone (3ae). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2e** (81.9 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 24 h. **3ae** (89 mg, 93%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 179-180 °C; IR (KBr) ν 3051, 1682, 1483, 1380, 1297 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.58 (d, J = 4.1 Hz, 1H), 7.74 – 7.56 (m, 3H), 7.56 – 7.44 (m, 3H), 7.19 (dd, J = 6.6, 5.0 Hz, 1H), 6.66 (d, J = 7.2 Hz, 1H), 6.42 (d, J = 9.2 Hz, 1H), 6.12 (dd, J = 9.2, 5.5 Hz, 1H), 5.41 – 5.30 (m, 1H), 5.19 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.1, 173.9, 158.5, 149.4, 137.4, 136.9, 129.6, 129.4, 127.9, 124.3, 123.6, 122.8, 121.9, 120.5, 109.0, 99.6, 73.8.; HRMS (ESI): Exact mass calcd for $\text{C}_{19}\text{H}_{13}\text{ClN}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 343.0605, found 343.0609.



3-(3-chlorophenyl)-8a-(pyridin-2-yl)-indolizinone (3af). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2f** (81.9 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 4 h. **3af** (89 mg, 93%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 152-153 °C; IR (KBr) ν 3049, 1691, 1539, 1430, 1375 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.59 (d, J = 4.2 Hz, 1H), 7.74 – 7.62 (m, 2H), 7.61 – 7.42 (m, 4H), 7.20 (dd, J = 6.8, 5.1 Hz, 1H), 6.67 (d, J = 7.2 Hz, 1H), 6.43 (d, J = 9.2 Hz, 1H), 6.13 (dd, J = 9.2, 5.4 Hz, 1H), 5.43 – 5.31 (m, 1H), 5.21 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 190.2, 173.5, 158.5, 149.5, 136.9, 135.2, 131.2, 130.4, 128.2, 126.4, 124.3, 123.6, 122.8, 121.9, 120.5, 109.1, 88.8, 73.8.; HRMS (ESI): Exact mass calcd for $\text{C}_{19}\text{H}_{13}\text{ClN}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 343.0605, found 343.0609.

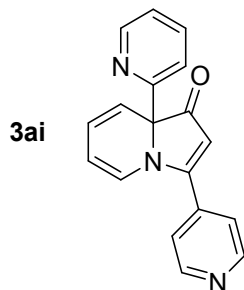


3-(2-chlorophenyl)-8a-(pyridin-2-yl)-indolizinone (3ag). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2g** (81.9 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 4 h. **3ag** (92 mg, 96%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 163-164 °C; IR (KBr) ν 3053, 1683, 1566, 1391, 1301 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.63 (d, J = 4.4 Hz, 1H), 7.67 (td, J = 7.7, 1.7 Hz, 1H), 7.60 – 7.51 (m, 3H), 7.45 (dtd, J = 14.4, 7.3, 1.7 Hz, 2H), 7.19 (ddd, J = 7.2, 4.8, 1.0 Hz, 1H), 6.48 (d, J = 9.2 Hz, 1H), 6.30 (d, J = 7.2 Hz, 1H), 6.11 (dd, J = 9.2, 5.5 Hz, 1H), 5.35 (t, J = 6.1 Hz, 1H), 5.19 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.3, 171.9, 158.3, 149.7, 136.7, 132.5, 131.9, 130.7, 130.4, 128.8, 127.3, 124.2, 123.4, 122.7, 122.5, 119.9, 109.4, 100.5, 73.4.; HRMS (ESI): Exact mass calcd for $\text{C}_{19}\text{H}_{13}\text{ClN}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 343.0604, found 343.0609.

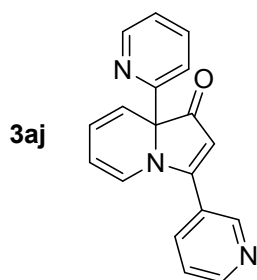


3-(4-bromophenyl)-8a-(pyridin-2-yl)-indolizinone (3ah). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2h** (108.6 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 24 h. **3ah** (100 mg, 91%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 205-206 °C; IR (KBr) ν 3086, 1680, 1480, 1430, 1291 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.57 (d, J = 4.2 Hz, 1H), 7.67 (dd, J = 10.9, 5.0 Hz, 3H), 7.52 (dd, J = 15.4, 8.2 Hz, 3H), 7.19 (dd, J = 6.9, 5.4 Hz, 1H), 6.65 (d, J = 7.2 Hz, 1H), 6.42 (d, J = 9.2 Hz, 1H), 6.12 (dd, J = 9.2, 5.5 Hz, 1H), 5.40 – 5.29 (m, 1H), 5.19 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.1, 173.9,

158.5, 149.4, 136.9, 132.4, 129.7, 128.4, 125.7, 124.3, 123.6, 122.8, 121.9, 120.5, 109.0, 99.6, 73.8.; HRMS (ESI): Exact mass calcd for $C_{19}H_{13}BrN_2O^+$ $[M+Na]^+$ 387.0097, found 387.0104.

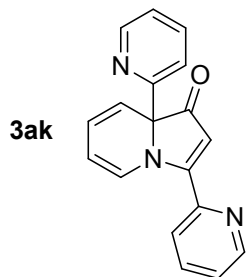


3-(pyridin-4-yl)-8a-(pyridin-2-yl)-indolizinone (3ai). The reaction of $Cu(OAc)_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2i** (61.8 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 130 °C for 24 h. **3ai** (56 mg, 64%, n-hexane/ethyl acetate = 1:2): yellow solid; mp 230-231 °C; IR (KBr) ν 3050, 1686, 1487, 1390, 1303 cm^{-1} ; 1H NMR (400 MHz, $CDCl_3$) δ 8.83 (d, J = 5.7 Hz, 2H), 8.57 (d, J = 4.1 Hz, 1H), 7.69 (td, J = 7.9, 1.6 Hz, 1H), 7.60 – 7.54 (m, 2H), 7.51 (d, J = 8.0 Hz, 1H), 7.21 (dd, J = 6.9, 5.3 Hz, 1H), 6.61 (d, J = 7.2 Hz, 1H), 6.42 (d, J = 9.2 Hz, 1H), 6.13 (dd, J = 9.2, 5.5 Hz, 1H), 5.39 (t, J = 6.3 Hz, 1H), 5.26 (s, 1H); ^{13}C NMR (75 MHz, $CDCl_3$) δ 199.2, 172.0, 158.2, 150.8, 149.4, 137.2, 137.0, 123.9, 123.7, 122.9, 122.26, 121.8, 120.7, 109.4, 100.2, 73.8.; HRMS (ESI): Exact mass calcd for $C_{18}H_{13}N_3O^+$ $[M+Na]^+$ 310.0949, found 310.0951.

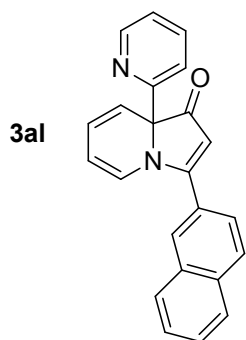


3-(pyridin-3-yl)-8a-(pyridin-2-yl)-indolizinone (3aj). The reaction of $Cu(OAc)_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2j** (61.8 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 72 h. **3aj** (68 mg, 79%, n-hexane/ethyl acetate = 1:2): yellow solid; mp 160-161 °C; IR (KBr) ν 3047, 1681, 1570, 1426, 1301 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$) δ 8.93 (s, 1H), 8.78 (d, J = 3.7 Hz, 1H), 8.56 (d, J = 4.7 Hz, 1H), 8.06 – 7.93 (m, 1H), 7.67 (td, J = 7.8, 1.7 Hz, 1H),

7.48 (dd, $J = 7.9, 4.1$ Hz, 2H), 7.23 – 7.14 (m, 1H), 6.63 (t, $J = 5.7$ Hz, 1H), 6.42 (d, $J = 9.2$ Hz, 1H), 6.13 (dd, $J = 9.2, 5.3$ Hz, 1H), 5.45 – 5.32 (m, 1H), 5.25 (d, $J = 4.5$ Hz, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.1, 171.6, 158.3, 152.1, 149.4, 148.8, 136.9, 135.7, 125.8, 124.0, 123.7, 122.9, 121.9, 120.7, 109.4, 100.2, 73.8.; HRMS (ESI): Exact mass calcd for $\text{C}_{18}\text{H}_{13}\text{N}_3\text{O}^+ [\text{M}+\text{Na}]^+$ 310.0948, found 310.0951.

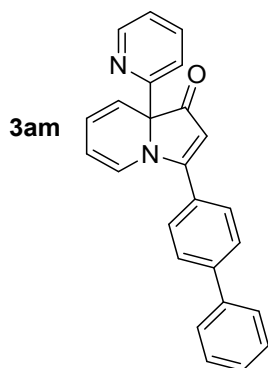


3-(pyridin-2-yl)-8a-(pyridin-2-yl)-indolizinone (3ak). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2k** (61.8 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 130 °C for 18 h. **3ak** (53 mg, 61%, n-hexane/ethyl acetate = 1:2): yellow solid; mp 155-156 °C; IR (KBr) ν 3095, 1682, 1537, 1392, 1285 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.83 (d, $J = 4.6$ Hz, 1H), 8.62 (d, $J = 4.2$ Hz, 1H), 7.88 (td, $J = 7.8, 1.6$ Hz, 1H), 7.77 (d, $J = 7.8$ Hz, 1H), 7.65 (td, $J = 7.8, 1.7$ Hz, 1H), 7.55 (t, $J = 8.3$ Hz, 2H), 7.45 (dd, $J = 6.5, 5.0$ Hz, 1H), 7.17 (dd, $J = 6.5, 5.0$ Hz, 1H), 6.51 (d, $J = 9.2$ Hz, 1H), 6.11 (dd, $J = 9.2, 5.4$ Hz, 1H), 5.44 (s, 1H), 5.42 – 5.34 (m, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.4, 171.5, 158.5, 149.8, 149.7, 149.1, 137.3, 136.8, 126.1, 125.4, 125.2, 123.5, 122.7, 122.5, 119.9, 109.2, 99.5, 74.2.; HRMS (ESI): Exact mass calcd for $\text{C}_{18}\text{H}_{13}\text{N}_3\text{O}^+ [\text{M}+\text{Na}]^+$ 310.0947, found 310.0951.

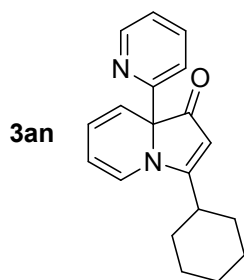


3-(naphthalen-2-yl)-8a-(pyridin-2-yl)-indolizinone (3al). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2l** (91.2 mg,

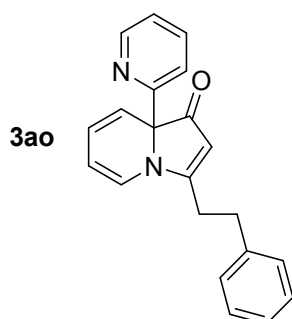
0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 130 °C for 5 h. **3al** (100 mg, 100%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 186-187 °C; IR (KBr) ν 3050, 1675, 1531, 1391, 1296 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 8.64 (d, J = 4.0 Hz, 1H), 8.19 (s, 1H), 8.00 (d, J = 8.5 Hz, 1H), 7.98 – 7.89 (m, 2H), 7.76 – 7.65 (m, 2H), 7.65 – 7.53 (m, 3H), 7.21 (dd, J = 6.7, 5.3 Hz, 1H), 6.81 (d, J = 7.2 Hz, 1H), 6.47 (d, J = 9.2 Hz, 1H), 6.14 (dd, J = 9.2, 5.5 Hz, 1H), 5.36 (t, J = 6.3 Hz, 1H), 5.32 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.3, 175.4, 158.8, 149.5, 136.8, 134.4, 132.9, 128.9, 128.66, 128.5, 127.9, 127.9, 127.2, 126.8, 124.8, 124.7, 123.5, 122.7, 122.1, 120.3, 108.8, 99.8, 73.9.; HRMS (ESI): Exact mass calcd for $\text{C}_{23}\text{H}_{16}\text{N}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 359.1150, found 359.1155.



3-([1,1'-biphenyl]-4-yl)-8a-(pyridin-2-yl)-indolizinone (3am). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2mg, 0.3 mmol), **2m** (106.8 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 130 °C for 1 h. **3am** (100 mg, 92%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 177-178 °C; IR (KBr) ν 3049, 1677, 1568, 1427, 1298 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.61 (d, J = 4.0 Hz, 1H), 7.76 (d, J = 2.8 Hz, 4H), 7.72 – 7.61 (m, 3H), 7.55 – 7.38 (m, 4H), 7.23 – 7.15 (m, 1H), 6.81 (d, J = 7.3 Hz, 1H), 6.47 (d, J = 9.2 Hz, 1H), 6.13 (dd, J = 9.2, 5.4 Hz, 1H), 5.38 (dd, J = 9.3, 3.4 Hz, 1H), 5.26 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.16, 174.94, 158.72, 149.49, 144.16, 139.86, 136.83, 129.05, 128.73, 128.26, 128.21, 127.71, 127.20, 124.70, 123.50, 122.73, 122.08, 120.29, 108.84, 99.48, 73.87.; HRMS (ESI): Exact mass calcd for $\text{C}_{25}\text{H}_{18}\text{N}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 385.1306, found 385.1311.

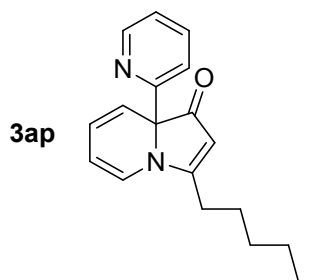


3-cyclohexyl-8a-(pyridin-2-yl)-indolizinone (3an). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **1a** (55.2 mg, 0.3 mmol), **2n** (64.8 mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 1 h. **3an** (79 mg, 90%, n-hexane/ethyl acetate = 2:1): yellow solid; mp 134-135 °C; IR (KBr) ν 3045, 1671, 1521, 1442, 1317 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.54 (d, J = 4.6 Hz, 1H), 7.61 (t, J = 7.7 Hz, 1H), 7.34 (d, J = 7.9 Hz, 1H), 7.21 – 7.09 (m, 1H), 6.67 (d, J = 7.1 Hz, 1H), 6.41 (d, J = 9.1 Hz, 1H), 6.07 (dd, J = 9.2, 5.2 Hz, 1H), 5.50 – 5.31 (m, 1H), 4.93 (s, 1H), 2.67-2.63 (m, 1H), 2.31 – 1.15 (m, 10H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.3, 182.6, 158.5, 149.3, 136.6, 123.4, 123.0, 122.5, 119.9, 109.3, 95.4, 72.9, 36.4, 31.5, 30.7, 26.1, 25.9, 25.8.; HRMS (ESI): Exact mass calcd for $\text{C}_{19}\text{H}_{20}\text{N}_2\text{O}^+ [\text{M}+\text{Na}]^+$ 315.1464, found 315.1468.

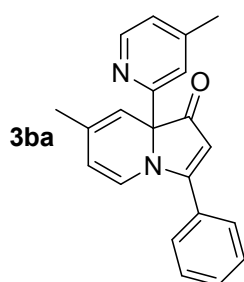


3-phenethyl-8a-(pyridin-2-yl)-indolizinone (3ao). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5mg, 0.03 mmol), NEt_3 (60.6mg, 0.6 mmol), **1a** (55.2mg, 0.3 mmol), **2o** (78mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 1 h. **3ao** (88 mg, 94%, n-hexane/ethyl acetate = 2:1): yellow solid; mp 157-158 °C; IR (KBr) ν 3043, 1683, 1565, 1443, 1312 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.55 (d, J = 4.1 Hz, 1H), 7.62 (t, J = 7.1 Hz, 1H), 7.31 (dd, J = 9.0, 5.6 Hz, 3H), 7.23 (dd, J = 8.3, 4.2 Hz, 2H), 7.19 – 7.12 (m, 1H), 6.60 (d, J = 7.1 Hz, 1H), 6.39 (d, J = 9.2 Hz, 1H), 6.09 (dd, J = 9.2, 5.4 Hz, 1H), 5.43 – 5.36 (m, 1H), 4.98 (s, 1H), 3.13 – 2.83 (m, 4H); ^{13}C NMR (75 MHz, CDCl_3) δ 199.3, 177.0, 158.4, 149.3, 139.7, 136.7, 128.7, 128.4, 126.7, 123.6, 123.0, 122.8,

122.6, 120.4, 109.6, 97.6, 73.1, 32.8, 29.2.; HRMS (ESI): Exact mass calcd for $C_{21}H_{18}N_2O^+ [M+Na]^+$ 337.1307, found 337.1311.

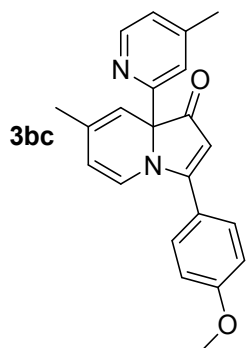


3-hexyl-8a-(pyridin-2-yl)-indolizinone (3ap). The reaction of $Cu(OAc)_2$ (5.5mg, 0.03 mmol), NEt_3 (60.6mg, 0.6 mmol), **1a** (55.2mg, 0.3 mmol), **2p** (66.0mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 1 h. **3ap** (70 mg, 80%, n-hexane/ethyl acetate = 2:1): yellow solid; mp 93-94 °C; IR (KBr) ν 3045, 1676, 1562, 1445, 1292 cm^{-1} ; 1H NMR (400 MHz, $CDCl_3$) δ 8.58 (s, 1H), 7.70 (s, 1H), 7.45 (s, 1H), 7.22 (s, 1H), 6.71 (s, 1H), 6.38 (d, J = 9.1 Hz, 1H), 6.11 (dd, J = 9.0, 5.5 Hz, 1H), 5.44 (t, J = 6.1 Hz, 1H), 4.97 (s, 1H), 2.81 – 2.53 (m, 2H), 1.89 – 0.75 (m, 13H).; ^{13}C NMR (75 MHz, $CDCl_3$) δ 199.3, 178.2, 158.5, 149.3, 136.6, 123.5, 123.1, 122.8, 122.6, 120.3, 109.3, 97.5, 73.1, 31.45, 29.0, 27.4, 26.7, 22.5, 14.0.; HRMS (ESI): Exact mass calcd for $C_{19}H_{22}N_2O^+ [M+Na]^+$ 317.1621, found 317.1624.

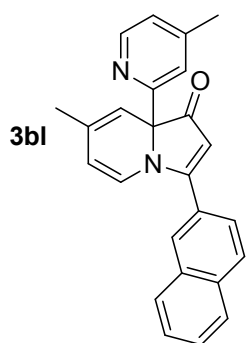


3-methyl-8a-(4-methylpyridin-2-yl)-2-phenyl-indolizinone (3ba). The reaction of $Cu(OAc)_2$ (3.7mg, 0.02 mmol), NEt_3 (40.4mg, 0.4 mmol), **1b**¹ (42.4mg, 0.2 mmol), **2a** (40.8mg, 0.4 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 1 h. **3ba** (48 mg, 77%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 181-182 °C; IR (KBr) ν 3054, 1680, 1540, 1376, 1287 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$) δ 8.43 (d, J = 4.9 Hz, 1H), 7.73 – 7.62 (m, 2H), 7.59 – 7.49 (m, 3H), 7.31 (s, 1H), 6.99 (d, J = 4.6 Hz, 1H), 6.70 (d, J = 7.2 Hz, 1H), 6.12 (d, J = 1.1 Hz, 1H), 5.21 (s, 1H), 5.20 (s, 1H), 2.35 (s, 3H), 1.84 (s, 3H); ^{13}C NMR (75 MHz, $CDCl_3$) δ 199.9, 175.0, 158.9, 149.0, 147.9,

132.1, 131.1, 129.7, 129.0, 128.2, 124.1, 123.7, 121.3, 117.1, 112.4, 99.9, 74.1, 21.3, 20.7.; HRMS (ESI): Exact mass calcd for $C_{21}H_{19}N_2O^+$ $[M+H]^+$ 315.1489, found 315.1492.

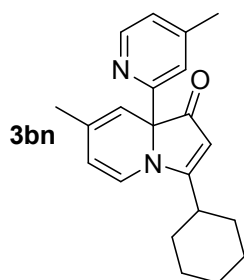


3-(4-methoxyphenyl)-8a-methyl-5-(4-methylpyridin-2-yl)-indolizinone (3bc). The reaction of $Cu(OAc)_2$ (2.7mg, 0.015 mmol), NEt_3 (30.3mg, 0.3 mmol), **1b**¹ (31.8mg, 0.15 mmol), **2c** (39.6mg, 0.3 mmol), in 1,4-dioxane (1.5 mL) were stirred at 110 °C for 24 h. **3bc** (35 mg, 68%, n-hexane/ethyl acetate = 2:1): yellow solid; mp 215-216 °C; IR (KBr) ν 3050, 1679, 1608, 1380, 1256 cm^{-1} ; 1H NMR (400 MHz, $CDCl_3$) δ 8.42 (d, J = 4.8 Hz, 1H), 7.63 (d, J = 8.4 Hz, 2H), 7.30 (s, 1H), 7.03 (d, J = 8.7 Hz, 2H), 6.98 (d, J = 4.4 Hz, 1H), 6.74 (d, J = 7.2 Hz, 1H), 6.11 (s, 1H), 5.21 (d, J = 7.3 Hz, 1H), 5.17 (s, 1H), 3.88 (s, 3H), 2.34 (s, 3H), 1.83 (s, 3H); ^{13}C NMR (75 MHz, $CDCl_3$) δ 199.8, 175.0, 161.9, 159.2, 149.1, 131.9, 129.9, 124.3, 123.6, 121.8, 121.3, 117.3, 114.4, 112.3, 99.2, 74.2, 55.5, 21.3, 20.7.; HRMS (ESI): Exact mass calcd for $C_{22}H_{21}N_2O_2^+$ $[M+H]^+$ 345.1593, found 345.1598.

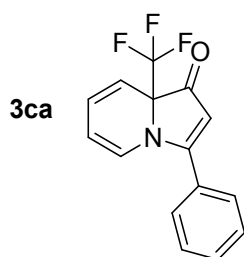


6-methyl-8a-(4-methylpyridin-2-yl)-3-(naphthalen-2-yl)-indolizinone (3bl). The reaction of $Cu(OAc)_2$ (2.7mg, 0.015 mmol), NEt_3 (30.3mg, 0.3 mmol), **1b**¹ (31.8mg, 0.15 mmol), **2l** (45.6mg, 0.3 mmol), in 1,4-dioxane (1.5 mL) were stirred at 110 °C for 24 h. **3bl** (45 mg, 82%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 217-218

°C; IR (KBr) ν 3054, 1679, 1536, 1392, 1340 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 8.64 (d, J = 4.0 Hz, 1H), 8.19 (s, 1H), 8.00 (d, J = 8.5 Hz, 1H), 7.98 – 7.89 (m, 2H), 7.76 – 7.65 (m, 2H), 7.65 – 7.53 (m, 3H), 7.21 (dd, J = 6.7, 5.3 Hz, 1H), 6.81 (d, J = 7.2 Hz, 1H), 6.47 (d, J = 9.2 Hz, 1H), 6.14 (dd, J = 9.2, 5.5 Hz, 1H), 5.36 (t, J = 6.3 Hz, 1H), 5.32 (s, 1H).; ^{13}C NMR (75 MHz, CDCl_3) δ 200.0, 175.2, 159.1, 149.1, 148.3, 134.3, 132.9, 132.1, 128.9, 128.7, 128.5, 127.9, 127.8, 127.1, 127.0, 124.8, 124.2, 123.7, 121.4, 117.2, 112.4, 100.2, 74.3, 21.3, 20.7.; HRMS (ESI): Exact mass calcd for $\text{C}_{25}\text{H}_{21}\text{N}_2\text{O}^+ [\text{M}+\text{H}]^+$ 365.1645, found 365.1648.

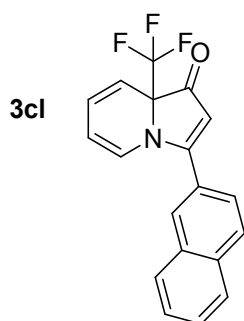


3-cyclohexyl-6-methyl-8a-(4-methylpyridin-2-yl)-indolizinone (3bn). The reaction of $\text{Cu}(\text{OAc})_2$ (2.7mg, 0.015 mmol), NEt_3 (30.3mg, 0.3 mmol), **1b**¹ (31.8mg, 0.15 mmol), **2n** (32.4mg, 0.3 mmol), in 1,4-dioxane (1.5 mL) were stirred at 110 °C for 1 h. **3bn** (51 mg, 80%, n-hexane/ethyl acetate = 2:1): yellow solid; mp 207-208 °C; IR (KBr) ν 3043, 1676, 1525, 1442, 1358 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 8.36 (d, J = 4.9 Hz, 1H), 7.17 (s, 1H), 6.94 (d, J = 4.0 Hz, 1H), 6.64 (d, J = 7.0 Hz, 1H), 6.09 (s, 1H), 5.28 (d, J = 7.2 Hz, 1H), 4.93 (s, 1H), 2.65 – 2.61 (m, 1H), 2.31 (s, 3H), 2.20 – 1.27 (m, 10H), 1.81 (s, 3H); ^{13}C NMR (75 MHz, CDCl_3) δ 200.1, 182.3, 158.8, 148.9, 147.6, 131.9, 123.4, 122.5, 121.0, 118.2, 112.9, 95.7, 73.4, 36.4, 31.5, 30.8, 26.1, 25.9, 25.8, 21.3, 20.6.; HRMS (ESI): Exact mass calcd for $\text{C}_{21}\text{H}_{25}\text{N}_2\text{O}^+ [\text{M}+\text{H}]^+$ 321.1958, found 321.1961.

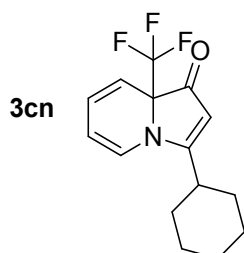


3-phenyl-8a-(trifluoromethyl)-indolizinone (3ca). The reaction of $\text{Cu}(\text{OAc})_2$ (5.5mg,

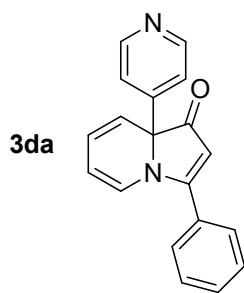
0.03 mmol), NEt₃ (60.6mg, 0.6 mmol), **1c** (52.5mg, 0.3 mmol), **2a** (61.2mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 20 h. **3ca** (58 mg, 70%, n-hexane/ethyl acetate = 12:1): yellow solid; mp 131-132 °C; IR (KBr) ν 3120, 1708, 1555, 1374, 1240 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 7.58 – 7.53(m, 5H), 6.57 (dd, J = 7.4, 0.7 Hz, 1H), 6.22 (dd, J = 9.4, 5.7 Hz, 1H), 5.80 (d, J = 9.4 Hz, 1H), 5.37 (dd, J = 7.2, 1.0 Hz, 1H), 5.35 (s, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 194.6, 178.5, 131.8, 129.3, 128.6, 127.9, 127.4, 126.1, 122.7 (q, J_{C-F} = 288.0 Hz, 1C), 113.3, 108.3, 101.8, 101.8, 69.6.; HRMS (ESI): Exact mass calcd for C₁₅H₁₁F₃NO⁺ [M+H]⁺ 278.0786, found 278.0787.



3-(naphthalen-2-yl)-8a-(trifluoromethyl)-indolizinone (3cl). The reaction of Cu(OAc)₂ (5.5mg, 0.03 mmol), NEt₃ (60.6mg, 0.6 mmol), **1c** (52.5mg, 0.3 mmol), **2l** (91.2mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 36 h. **3cl** (56 mg, 57%, n-hexane/ethyl acetate = 12:1): yellow solid; mp 144-145 °C; IR (KBr) ν 3083, 1704, 1550, 1380, 1243 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 8.09 (s, 1H), 8.04 – 7.86 (m, 3H), 7.66 – 7.58 (m, 3H), 6.65 (d, J = 7.5 Hz, 1H), 6.24 (dd, J = 9.4, 5.7 Hz, 1H), 5.83 (d, J = 9.4 Hz, 1H), 5.47 (s, 1H), 5.38 (t, J = 6.6 Hz, 1H); ¹³C NMR (75 MHz, CDCl₃) δ 194.6, 178.6, 134.6, 132.8, 129.3, 128.7, 128.5, 128.3, 128.0, 127.44, 127.40, 126.3, 125.9, 124.0, 122.8 (q, J_{C-F} = 288.0 Hz, 1C), 113.4, 108.3, 102.0, 69.6 (q, J_{C-F} = 28.5 Hz, 1C).; HRMS (ESI): Exact mass calcd for C₁₉H₁₃F₃NO⁺ [M+H]⁺ 328.0942, found 328.0944.



3-cyclohexyl-8a-(trifluoromethyl)-indolizinone (3cn). The reaction of Cu(OAc)₂ (5.5mg, 0.03 mmol), NEt₃ (60.6mg, 0.6 mmol), **1c** (52.5mg, 0.3 mmol), **2n** (64.8mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 96 h. **3cn** (45 mg, 53%, n-hexane/ethyl acetate = 20:1): yellow solid; mp 98-99 °C; IR (KBr) ν 3054, 1693, 1539, 1308, 1127 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 6.54 (d, *J* = 7.3 Hz, 1H), 6.22 (dd, *J* = 9.3, 5.6 Hz, 1H), 5.75 (d, *J* = 9.3 Hz, 1H), 5.55 – 5.43 (m, 1H), 5.05 (s, 1H), 2.57 – 2.39 (m, 1H), 2.14 – 1.16 (m, 10H); ¹³C NMR (75 MHz, CDCl₃) δ 195.0, 185.5, 127.5, 124.2, 122.7 (q, *J*_{C-F} = 288.0 Hz, 1C), 117.0, 114.1, 109.1, 97.6, 68.3 (q, *J*_{C-F} = 28.5 Hz, 1C), 36.3, 31.4, 30.3, 26.0, 25.70, 25.67.; HRMS (ESI): Exact mass calcd for C₁₅H₁₇F₃NO⁺ [M+H]⁺ 284.1257, found 284.1257.

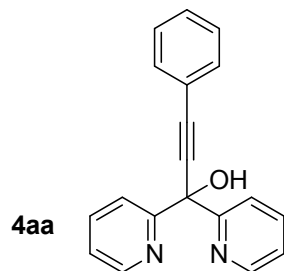


3-phenyl-8a-(pyridin-4-yl)-indolizinone (3da). The reaction of Cu(OAc)₂ (5.5mg, 0.03 mmol), NEt₃ (60.6mg, 0.6 mmol), **1d** (55.2mg, 0.3 mmol), **2a** (61.2mg, 0.6 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 63 h. **3da** (28 mg, 33%, n-hexane/ethyl acetate = 3:1): yellow solid; mp 234-235 °C; IR (KBr) ν 3093, 1676, 1536, 1484, 1390 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 8.63 (s, 2H), 7.68 – 7.52 (m, 5H), 7.46 (s, 2H), 6.74 (d, *J* = 7.3 Hz, 1H), 6.31 (d, *J* = 9.2 Hz, 1H), 6.08 (dd, *J* = 9.2, 5.5 Hz, 1H), 5.46 – 5.33 (m, 1H), 5.22 (s, 1H); ¹³C NMR (75 MHz, CDCl₃) δ 198.8, 175.1, 149.8, 149.2, 131.6, 129.3, 129.0, 128.1, 124.4, 123.8, 121.6, 109.6, 99.7, 71.3; HRMS (ESI): Exact mass calcd for C₁₉H₁₅N₂O⁺ [M+H]⁺ 287.1181, found 287.1179.

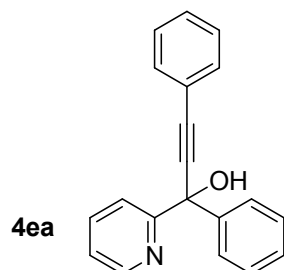
Preparation of tertiary propargylic alcohols 4aa and 4ea.

To a stirred solution of terminal alkyne (1.2 equiv) in THF was added n-BuLi (1.1 equiv, 1.6 M solution in hexanes) at -78 °C. After 5 min, a solution of pyridyl ketone (1.0 equiv) in THF was slowly added to this mixture at -78 °C. After 15 min at -78 °C,

the reaction mixture was quenched with saturated NH_4Cl . The reaction mixture was diluted with ethyl acetate and washed with aqueous NH_4Cl . The organic layer was dried over MgSO_4 and concentrated in vacuo to give a crude mixture, which was purified by silica gel column chromatography (hexane/ethyl acetate/dichloromethane) to afford the propargylic alcohol.²



3-phenyl-1,1-di(pyridin-2-yl)prop-2-yn-1-ol (4aa).² 132 mg (46%). White solid. mp 90-91 °C; IR (KBr) ν 3082, 1652, 1538, 1454, 1296 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.55 (d, J = 4.2 Hz, 2H), 7.95 (d, J = 8.0 Hz, 2H), 7.73 (td, J = 7.8, 1.7 Hz, 2H), 7.56 – 7.43 (m, 2H), 7.32 – 7.26 (m, 3H), 7.22 (dd, J = 6.7, 5.0 Hz, 2H), 6.79 (s, 1H).; ^{13}C NMR (75 MHz, CDCl_3) δ 161.0, 148.0, 137.2, 132.0, 128.5, 128.2, 122.9, 122.5, 121.4, 91.1, 85.5, 74.2.

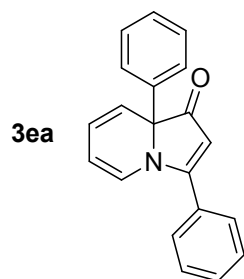


1,3-diphenyl-1-(pyridin-2-yl)prop-2-yn-1-ol (4ea).³ 180 mg (63%). White solid. mp 69-70 °C; IR (KBr) ν 3067, 1679, 1592, 1434, 1268 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 8.55 (d, J = 4.7 Hz, 1H), 7.77 – 7.60 (m, 3H), 7.51 (dd, J = 12.6, 4.9 Hz, 3H), 7.40 – 7.19 (m, 7H), 6.62 (s, 1H).; ^{13}C NMR (101 MHz, CDCl_3) δ 161.2, 146.9, 144.2, 137.6, 131.9, 128.6, 128.3, 128.3, 127.9, 126.5, 122.9, 122.5, 121.8, 91.0, 86.6, 73.6.

$\text{Cu}(\text{OAc})_2$ -Catalyzed Transformation from 4 to 3.

The mixture of $\text{Cu}(\text{OAc})_2$ (5.5 mg, 0.03 mmol), NEt_3 (60.6 mg, 0.6 mmol), **4aa** (85.8 mg, 0.3 mmol), in 1,4-dioxane (2 mL) were stirred at 110 °C for 5 h. Column chromatography on silica gel (n-hexane/ethyl acetate = 3:1) to afford 86 mg (100%)

of **3aa**.



3,8a-diphenyl-lindolizinone (3ea). The reaction of $\text{Cu}(\text{OAc})_2$ (2.5 mg, 0.0138 mmol), NEt_3 (27.8 mg, 0.276 mmol), **4ea** (39.3 mg, 0.138 mmol), in 1,4-dioxane (1 mL) were stirred at 110 °C for 4 h. **3ea** (34 mg, 88%, n-hexane/ethyl acetate = 3:1): yellow solid. mp 125-126 °C; IR (KBr) ν 3056, 1683, 1554, 1434, 1291 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 7.63 (dd, J = 6.8, 3.0 Hz, 2H), 7.58 – 7.49 (m, 5H), 7.39 – 7.28 (m, 3H), 6.69 (dd, J = 7.2, 0.7 Hz, 1H), 6.41 – 6.31 (m, 1H), 6.05 (dd, J = 9.3, 5.4 Hz, 1H), 5.42 – 5.32 (m, 1H), 5.20 (s, 1H); ^{13}C NMR (75 MHz, CDCl_3) δ 200.8, 174.6, 140.4, 131.2, 129.5, 129.2, 128.7, 128.1, 127.9, 124.5, 124.2, 122.9, 109.5, 99.5, 72.1.

Reference:

- 1) Collman, J. P.; Zhong, M.; Wang, Z.; Rapta, M. *Org. Lett.* **1999**, *13*, 2122.
- 2) Cho, H.; Kim, I. *Tetrahedron*. **2012**, *68*, 5464.
- 3) Yan, B.; Zhou Y. B.; Liu Y. H. *J. Org. Chem.* **2007**, *72*, 7783.

Fig.S1. GC-MS analysis of the mixture after the completion of the reaction between **1a** and **2a**.

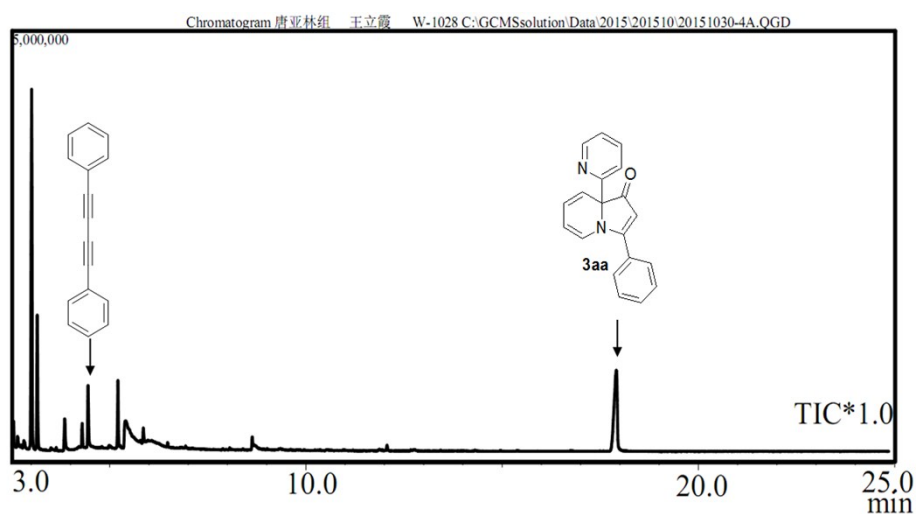


Fig.S2. X-ray crystal structure of **3af**

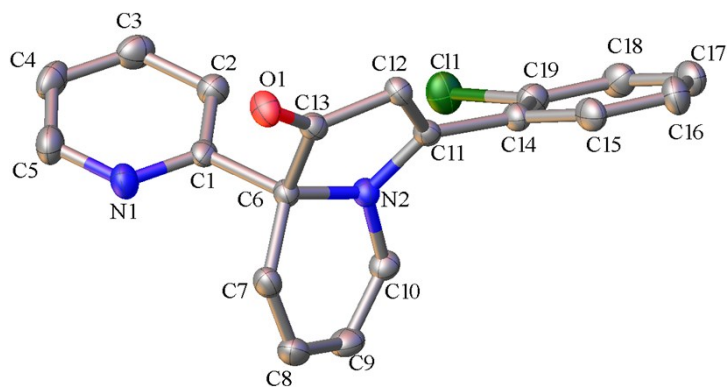
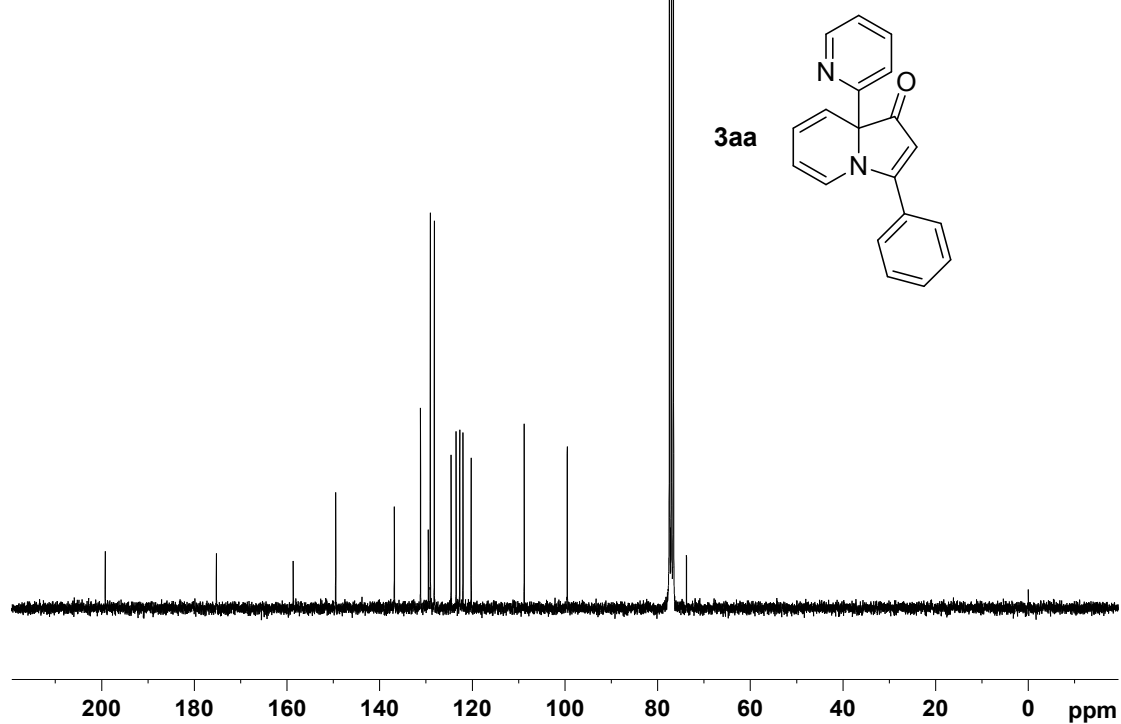
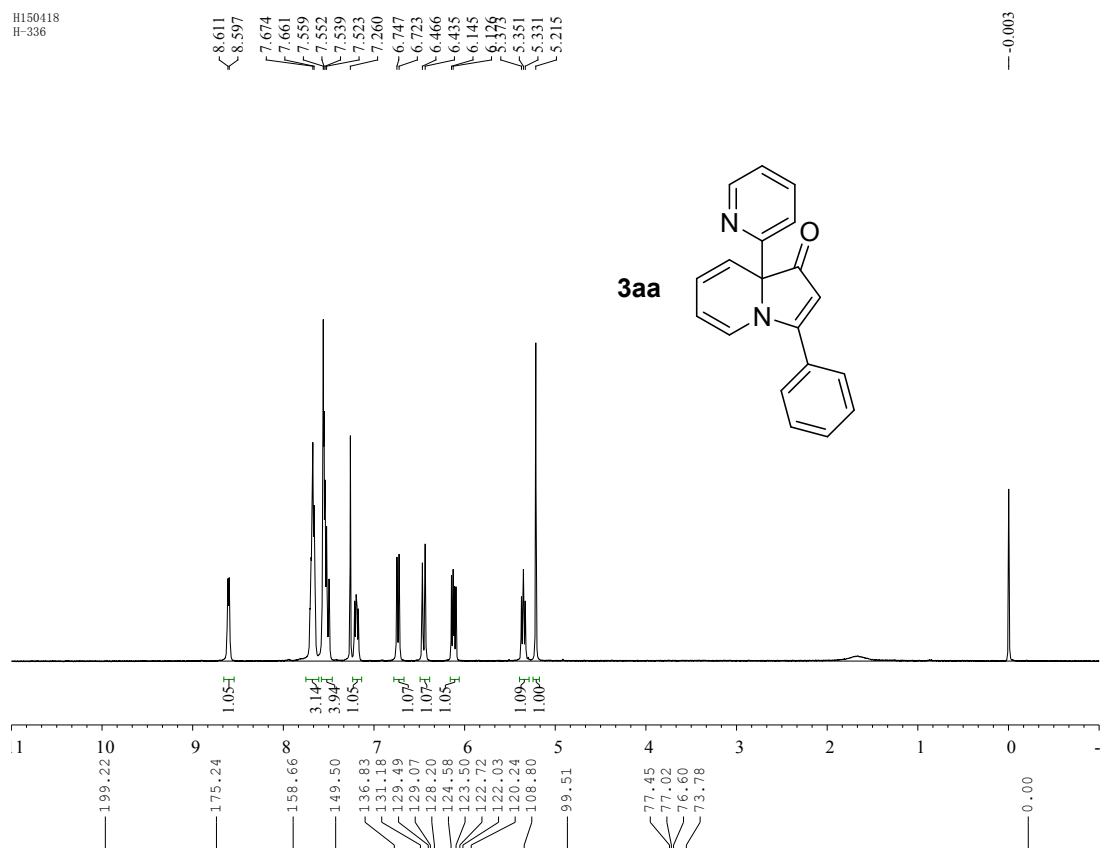


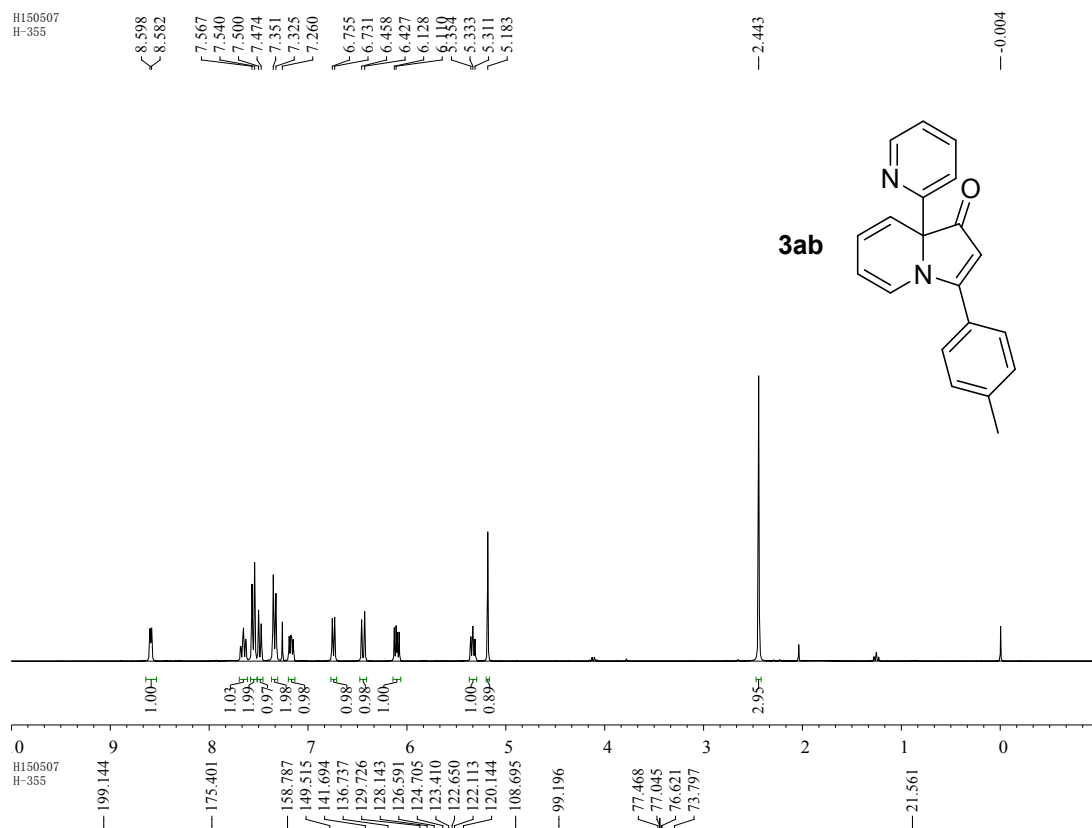
Table.S1. Crystal data and structural refinement for 3af and gem-diol complex.

compound	3af	gem-diol complex
empirical formula	C ₁₉ H ₁₃ ClN ₂ O	C ₃₄ H ₄₆ CuN ₄ O ₁₄
<i>M_r</i>	320.76	798.29
crystal size [mm ³]	0.31 x 0.29 x 0.28	0.20 x 0.14 x 0.10
crystal system	triclinic	monoclinic
space group	P-1	C12/c1
a [Å]	7.400(3)	21.170(8)
b [Å]	10.351(3)	7.831(3)
c [Å]	11.137(4)	24.511(9)
α [deg]	63.632(9)	90
β [deg]	84.599(15)	114.363(4)
γ [deg]	80.315(16)	90
V [Å ³]	753.3(4)	3702(2)
ρ [g/cm ³]	1.414	1.432
Z	2	4
T [K]	173(2)	173(2)
R1,wR2 [I>2σ(I)]	0.0613, 0.2141	0.0795, 0.1832
R1,wR2 (all data)	0.0633, 0.2158	0.0873, 0.1876
quality of fit	1.788	1.213

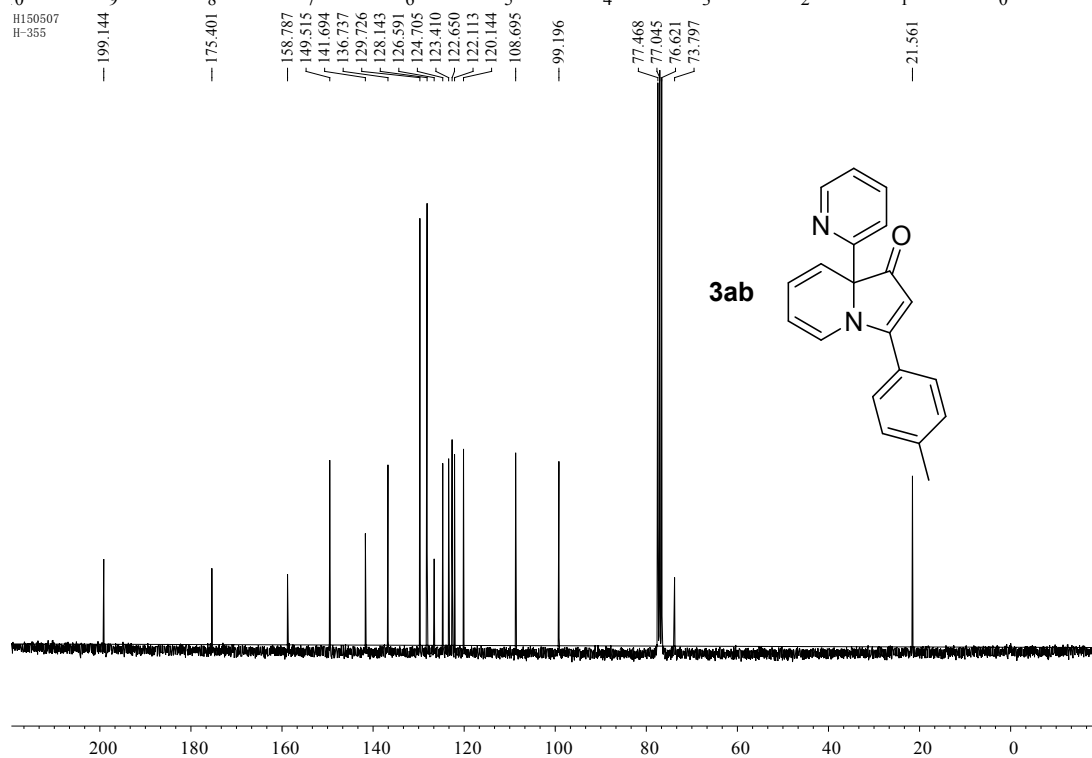
H150418
H-336



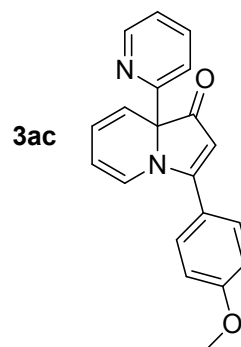
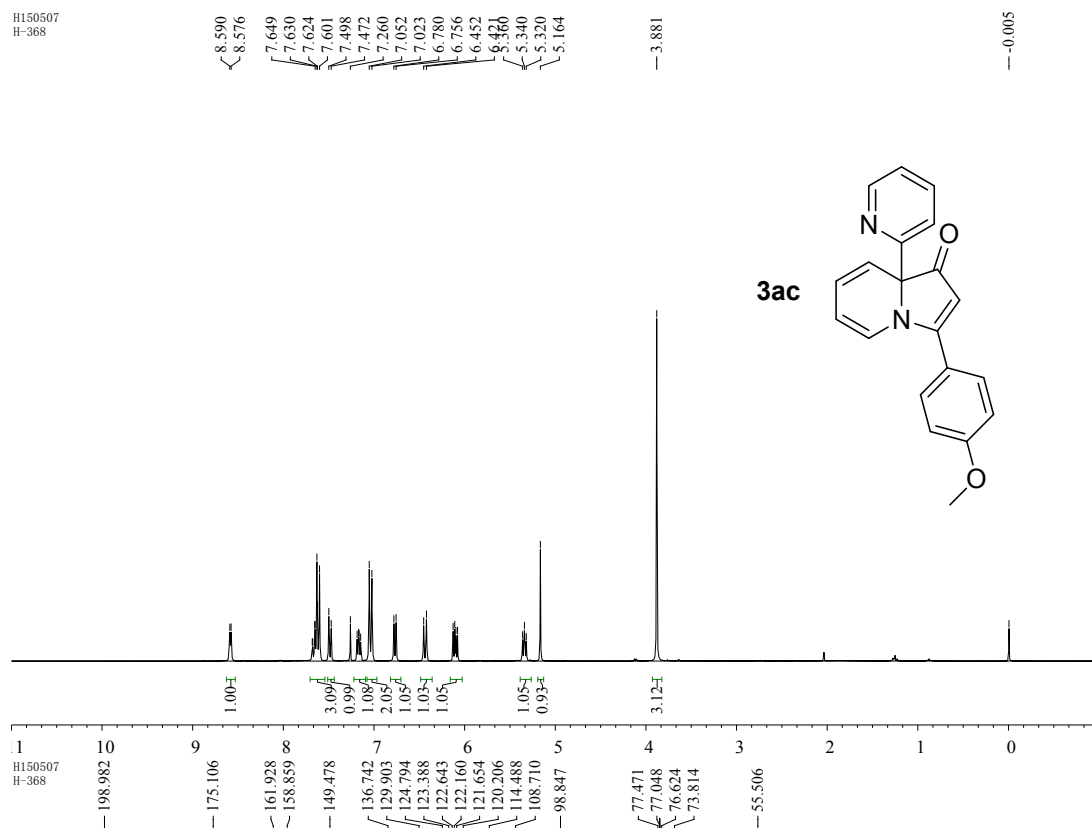
H150507
H-355



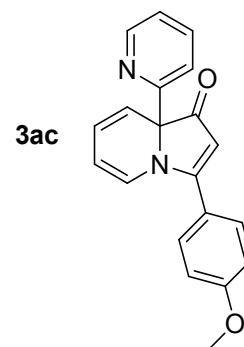
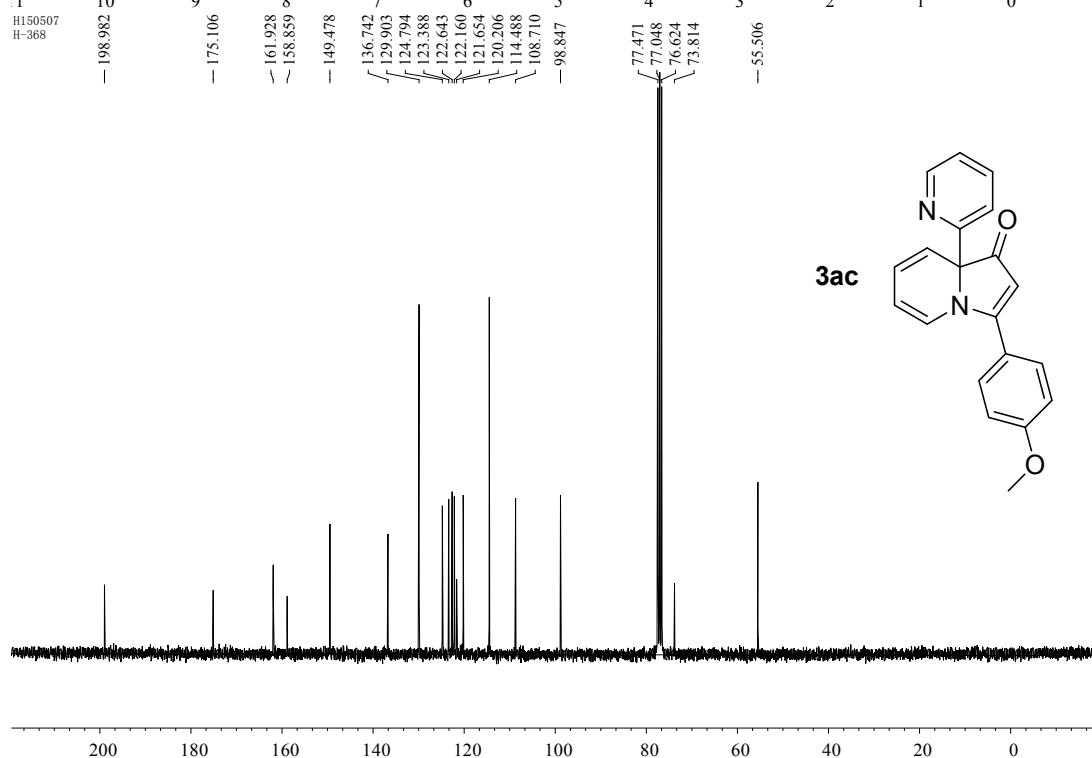
H150507
H-355



H150507
H-368



H150507
H-368

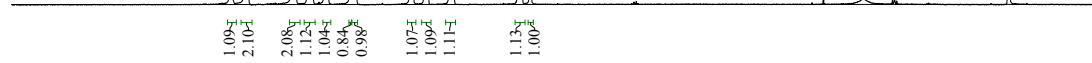
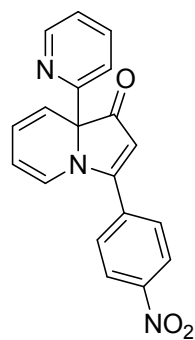


h150523
h-388

8.582
8.567
8.422
8.415
8.399
8.392
8.385
7.900
7.871
7.709
7.540
7.513
7.260
6.574
6.440
6.409
6.170
5.477
5.391
5.369
5.275

— -0.003

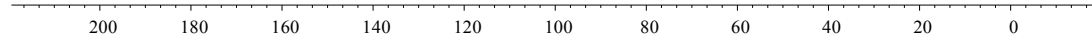
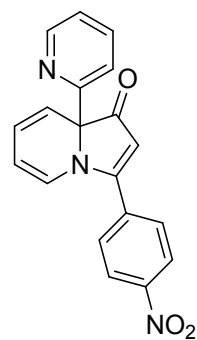
3ad



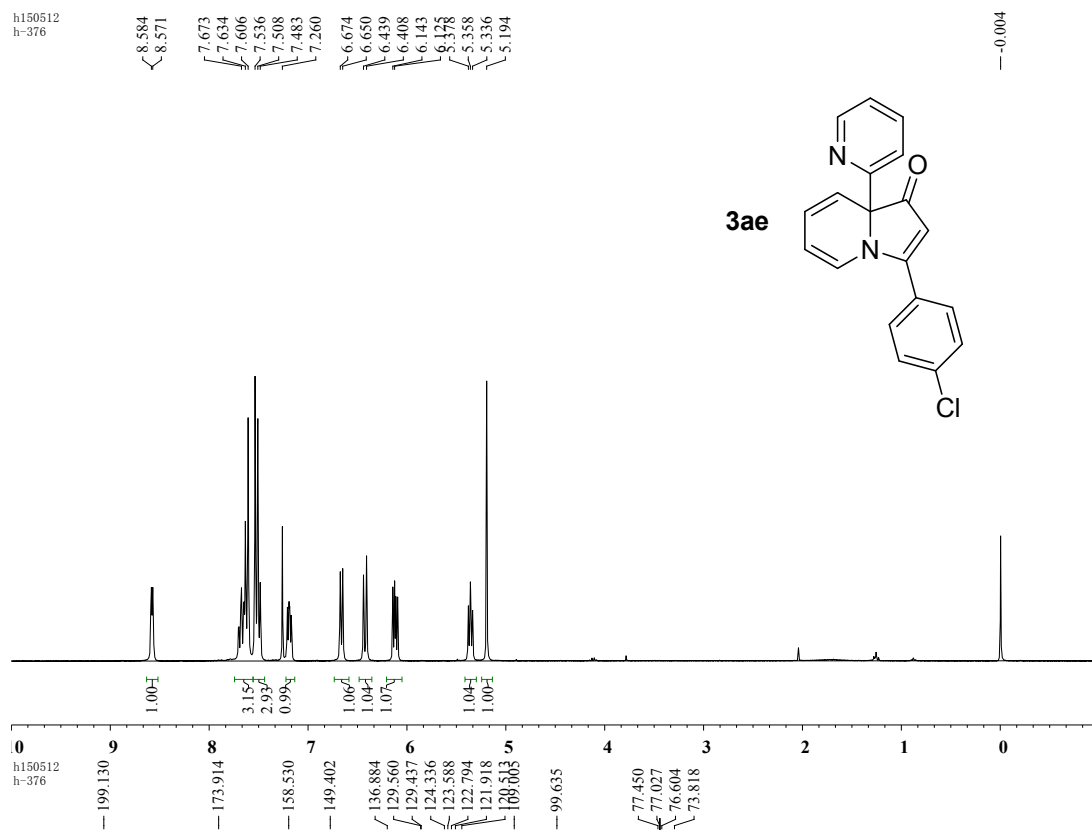
h150530
h-388

199.128
172.301
158.247
149.338
137.041
129.421
124.282
123.916
123.789
122.972
121.715
120.880
100.556
77.446
77.022
76.599
73.922

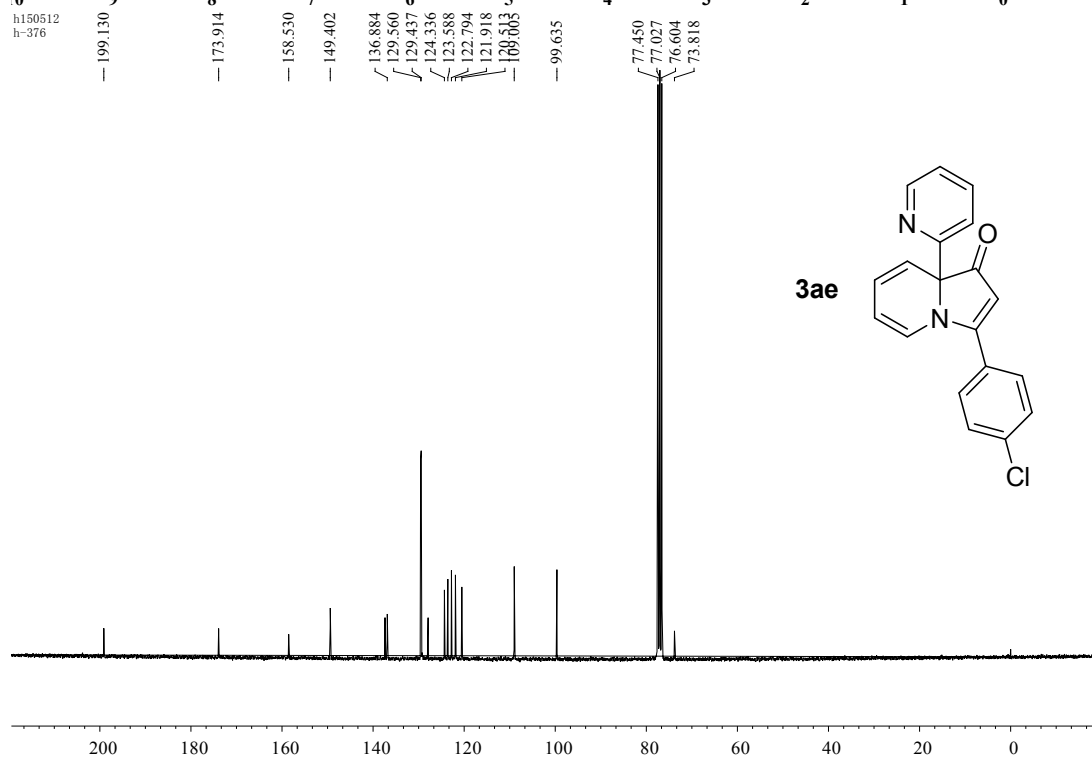
3ad



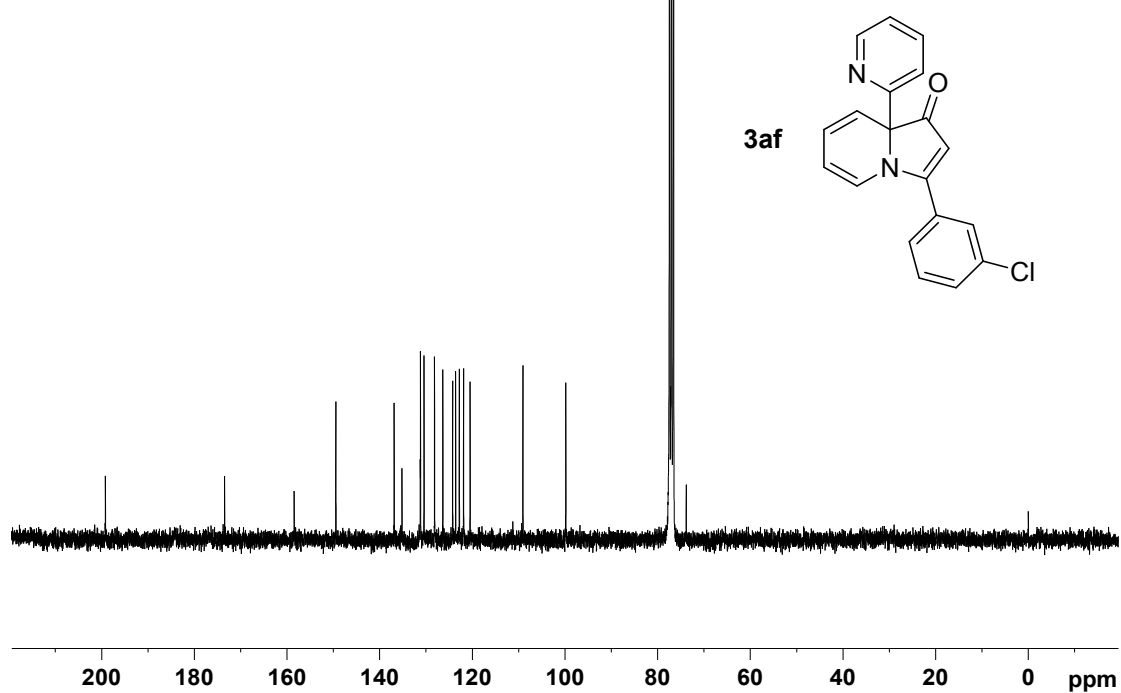
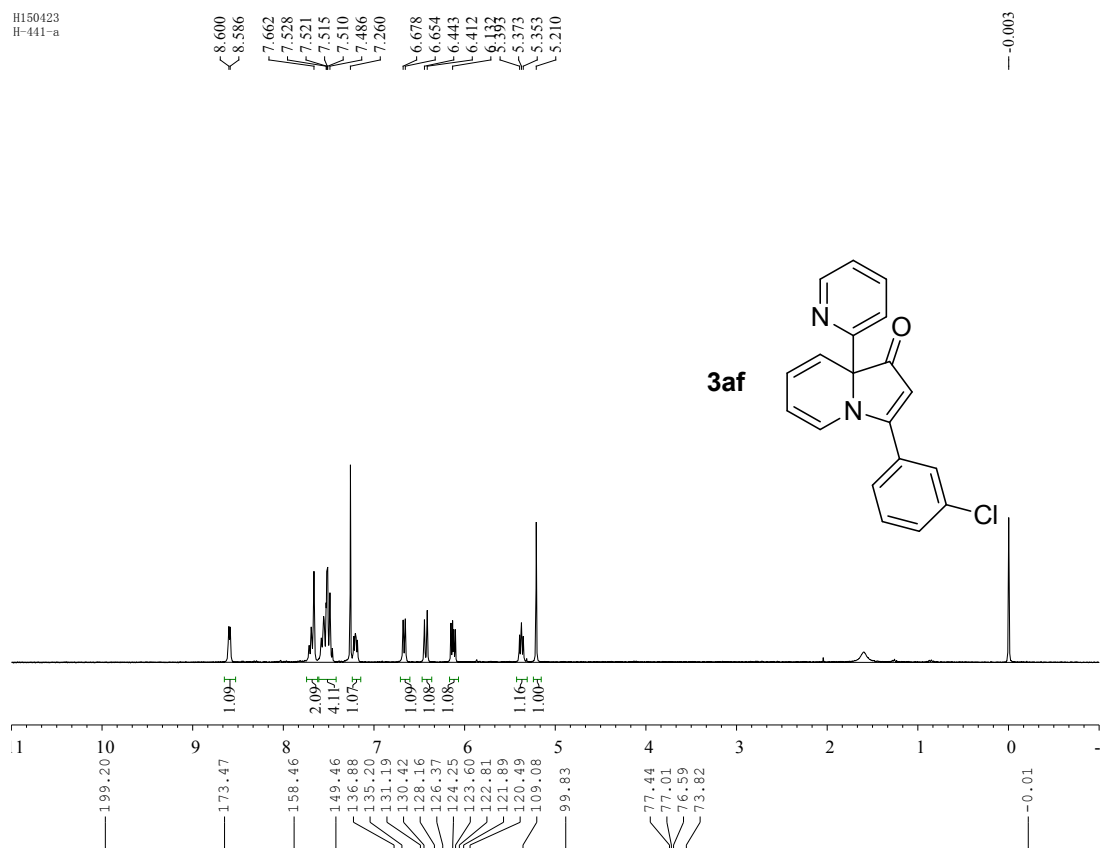
h150512
h-376



h150512
h-376



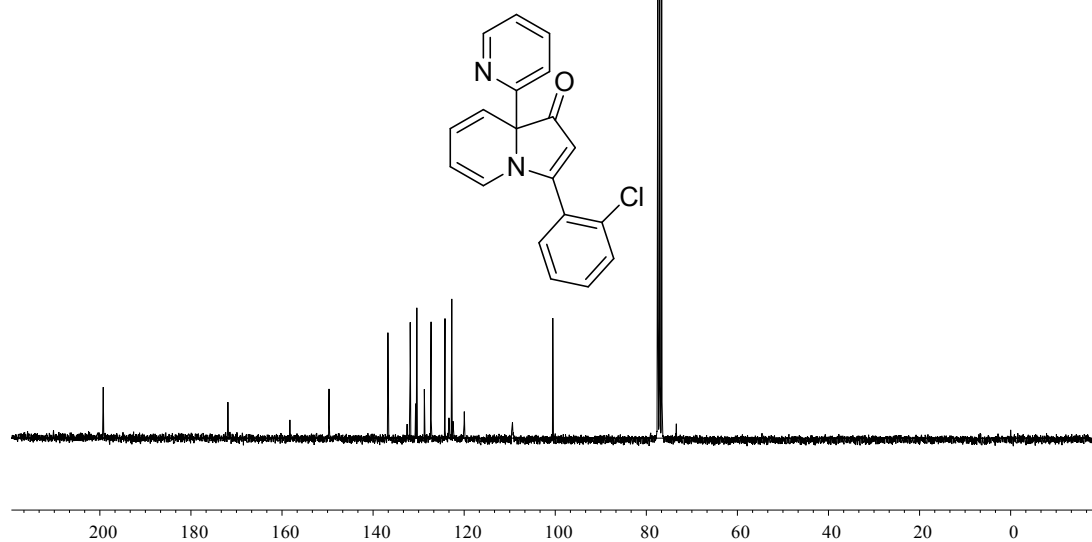
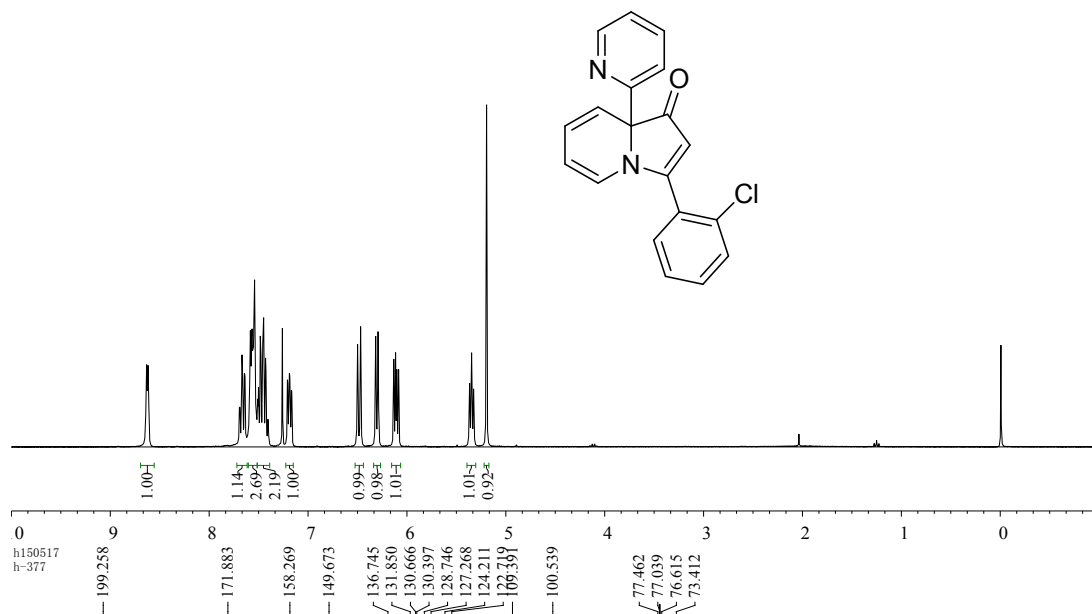
H150423
H-441-a



h150517
h-377

8.632
8.618
7.583
7.566
7.545
7.541
7.483
7.455
7.449
7.260
6.499
6.468
6.316
6.292
6.133
5.166
5.146
5.326
5.195

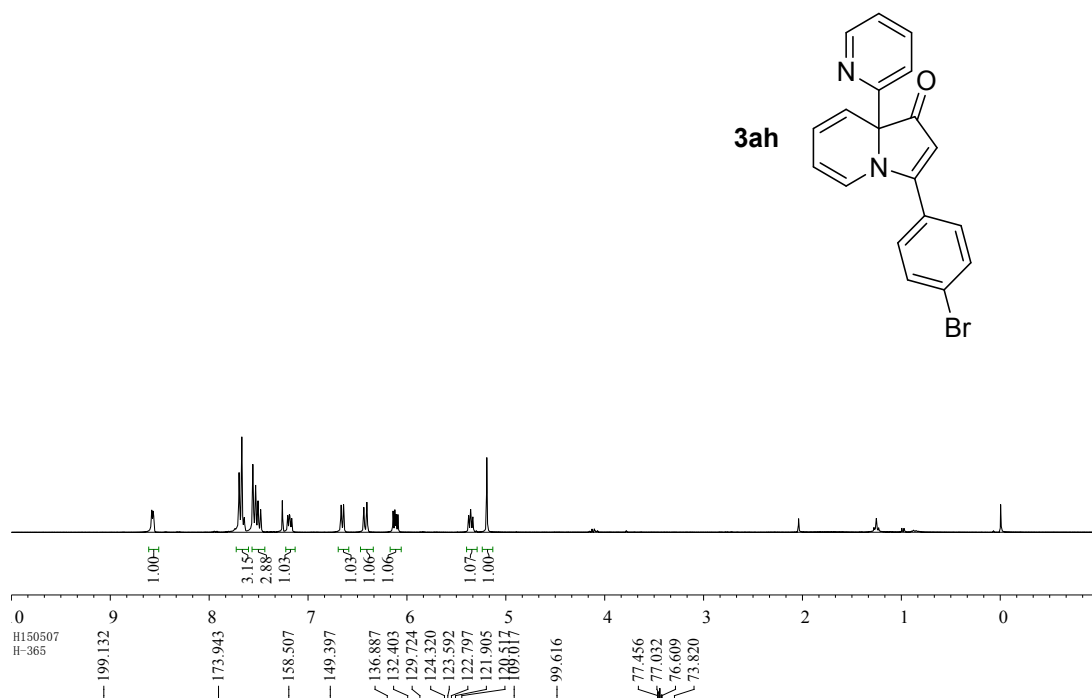
-0.007



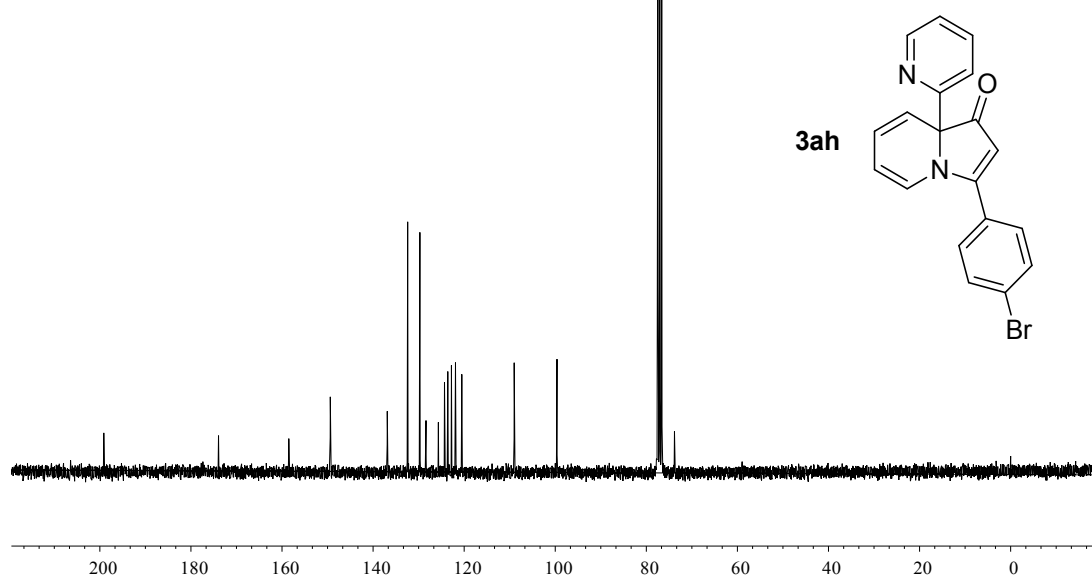
H150507
H-365

8.580
8.566
7.697
7.670
7.558
7.530
7.506
7.480
7.260
6.665
6.641
6.436
6.405
6.140
5.375
5.355
5.333
5.192

-0.004



H150507
H-365

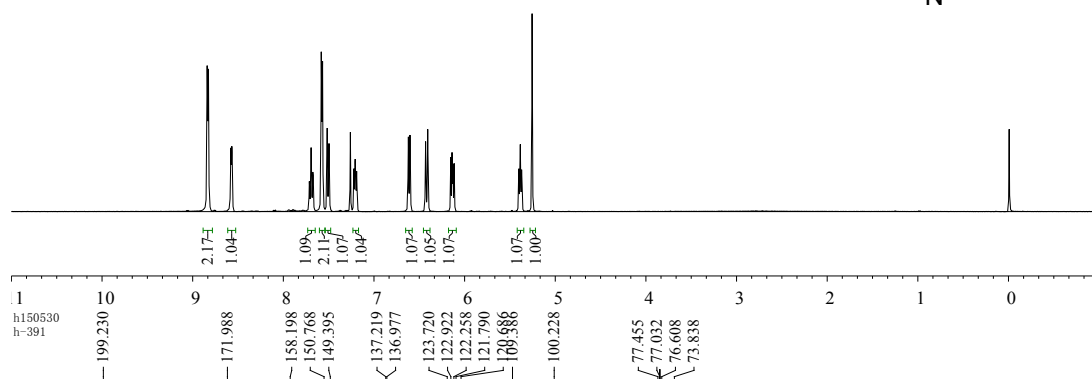
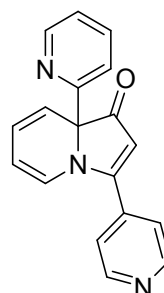


h150529
h-391

8.840
8.826
8.578
8.568
7.582
7.570
7.567
7.516
7.496
7.260
6.621
6.603
6.430
6.407
5.402
5.386
5.371
5.256

— -0.008

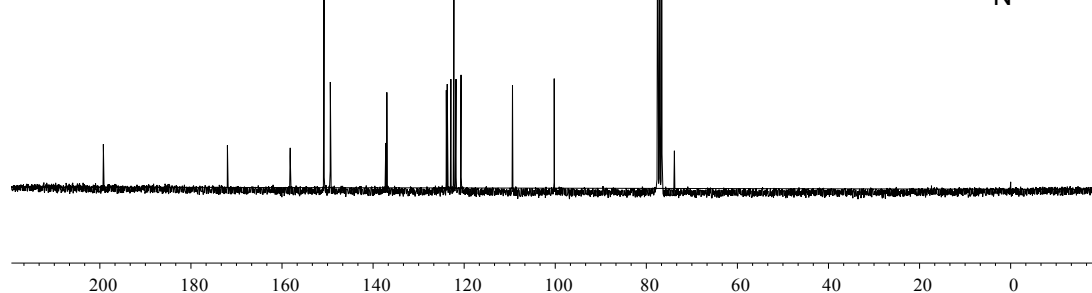
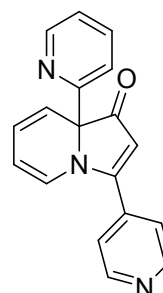
3ai



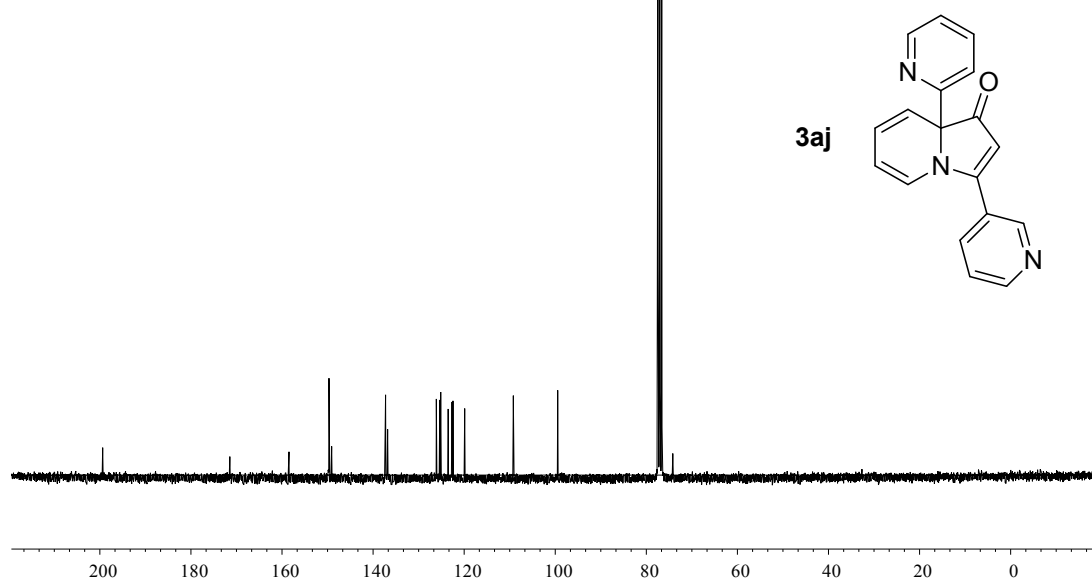
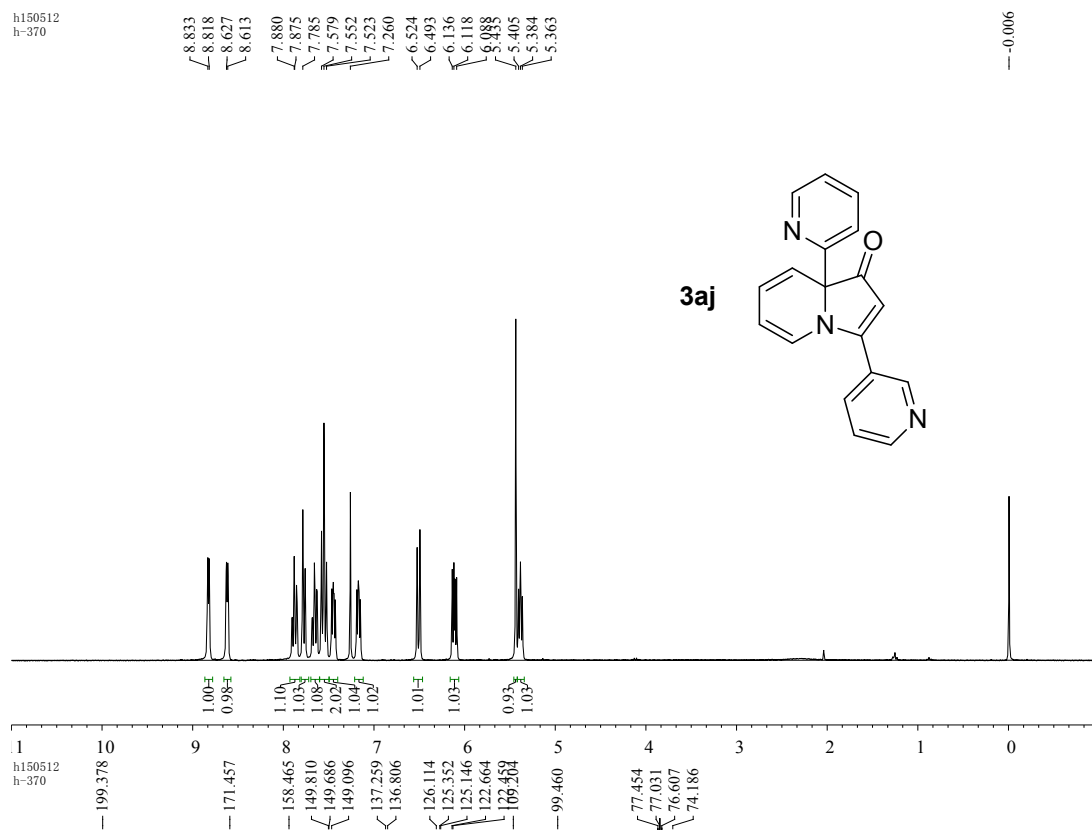
h150530
h-391

199.230
171.988
158.198
150.768
149.395
137.219
136.977
123.720
122.922
122.258
121.790
108.588
100.228
77.455
77.032
76.608
73.838

3ai



h150512
h-370

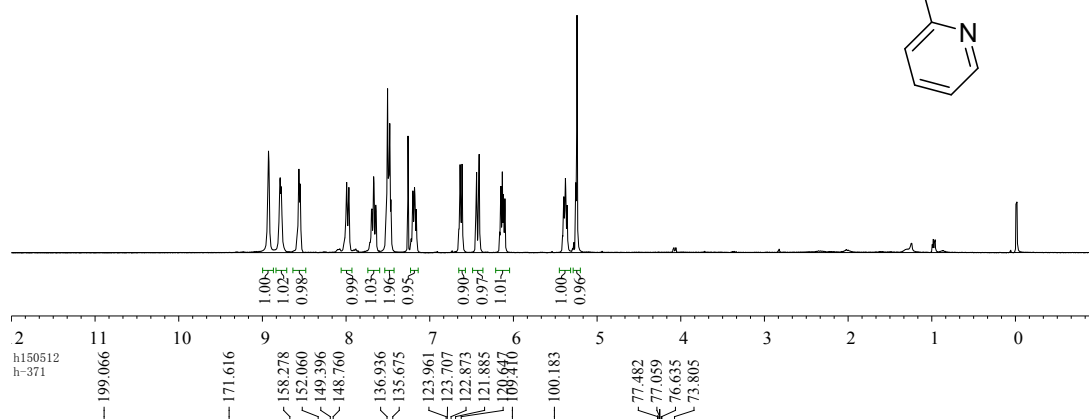
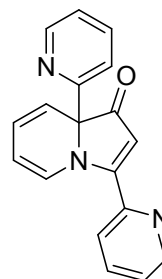


h150512
h-371

8.927
8.787
8.775
8.564
8.549
7.993
7.671
7.504
7.491
7.478
7.260
6.637
6.614
6.440
6.409
6.133
5.378
5.358
5.255
5.240

— -0.006

3ak



h150512
h-371

— 199.066

— 171.616

158.278

152.060

149.396

148.760

136.936

135.675

123.961

123.707

122.873

121.885

120.470

100.183

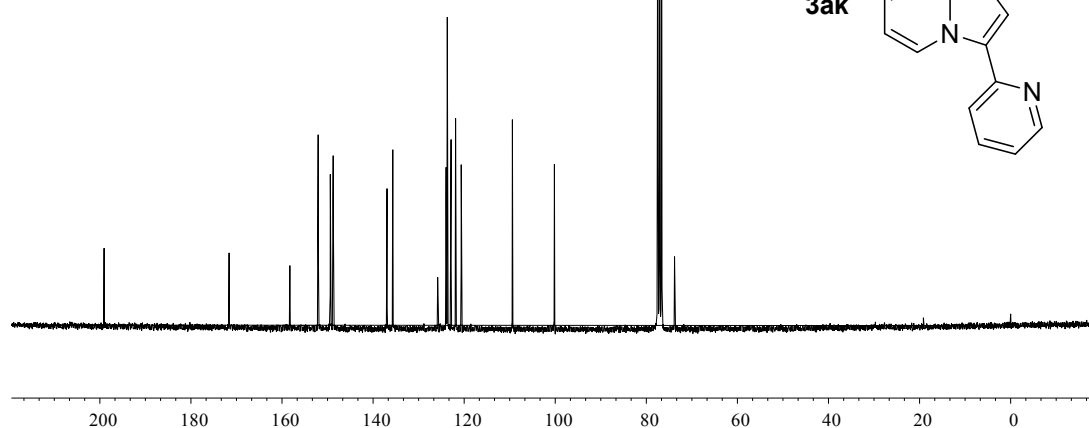
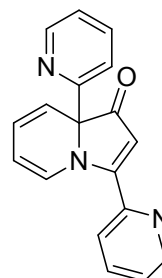
77.482

77.059

76.635

73.805

3ak

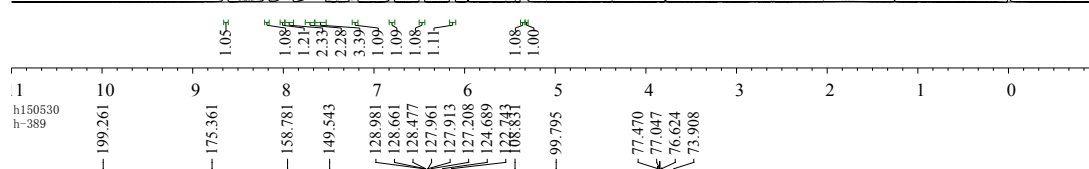
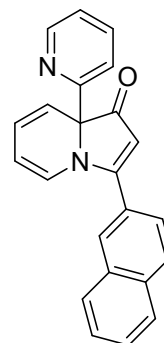


h150529
h-389-2

8.642
8.632
8.187
8.015
7.993
7.722
7.622
7.618
7.614
7.607
7.599
7.595
7.575
7.260
5.449
5.358
5.343
5.323

0.001

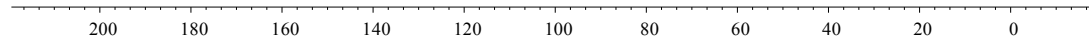
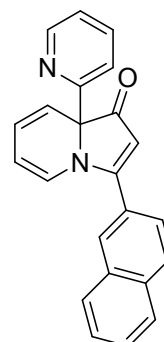
3aI



h150530
h-389

199.261
175.361
158.781
149.543
128.981
128.661
128.477
127.961
127.913
127.208
124.689
108.843
99.795
77.470
77.047
76.624
73.908

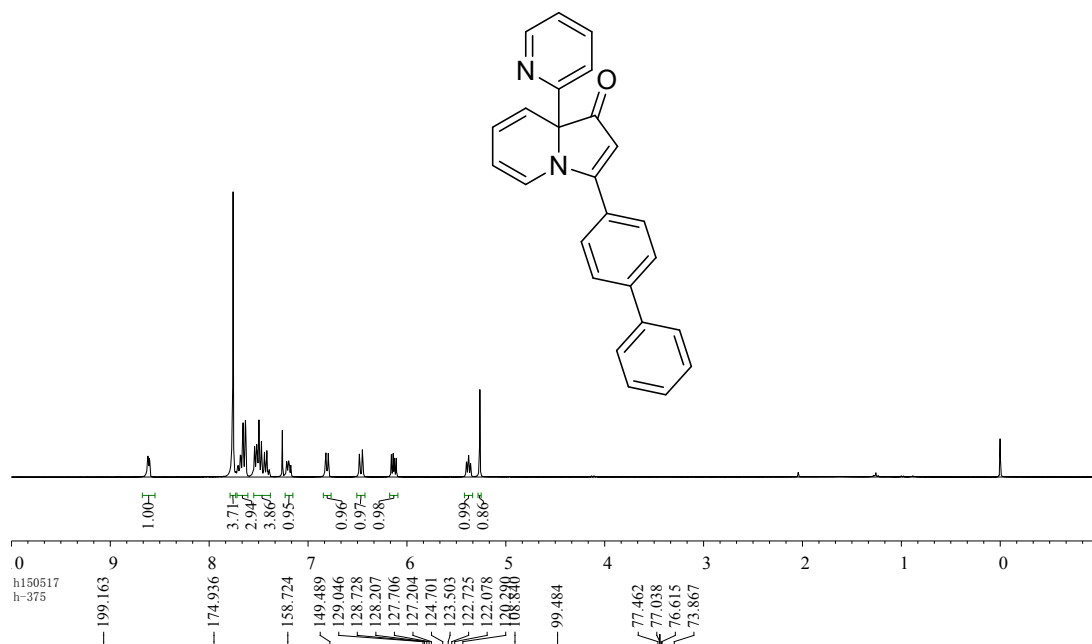
3aI



h150517
h-375

8.618
8.605
7.768
7.758
7.657
7.633
7.540
7.520
7.514
7.497
7.471
7.441
7.417
7.260
6.820
6.451
5.398
5.396
5.376
5.356
5.262

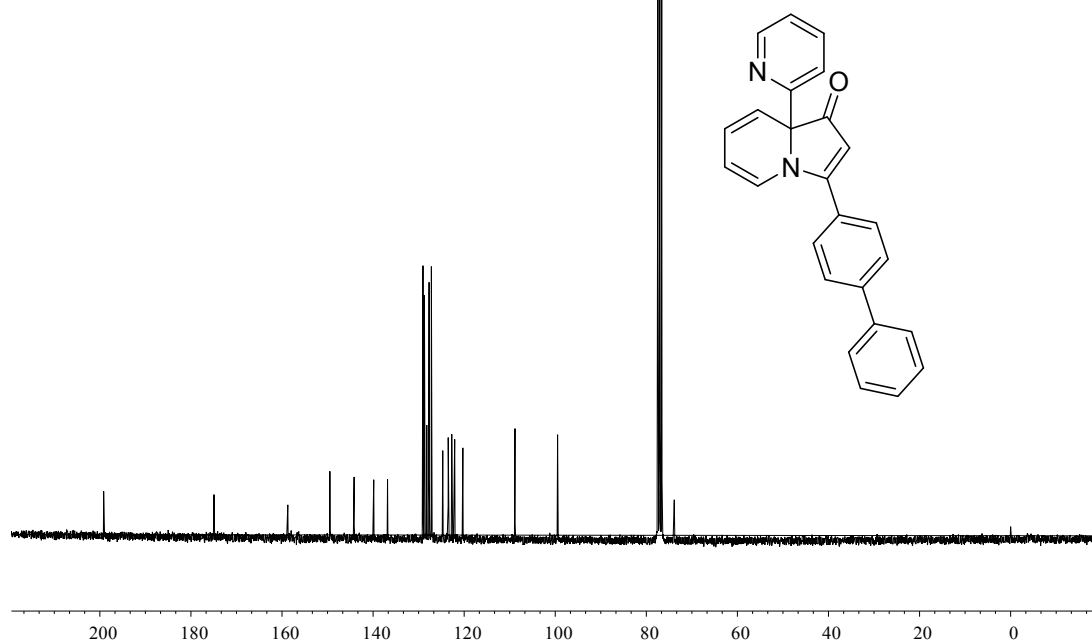
— 0.001



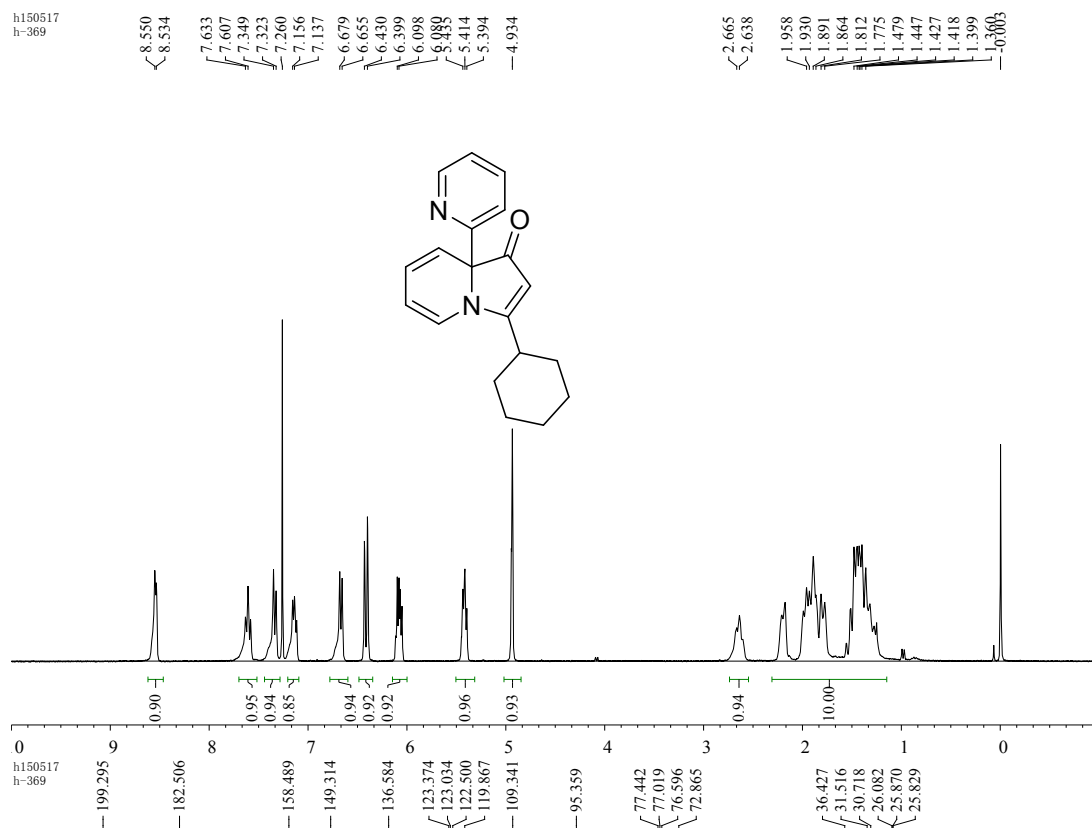
h150517
h-375

199.163
174.936
158.724
149.489
129.046
128.728
128.207
127.706
127.204
124.701
123.503
122.725
122.078
108.830
99.484

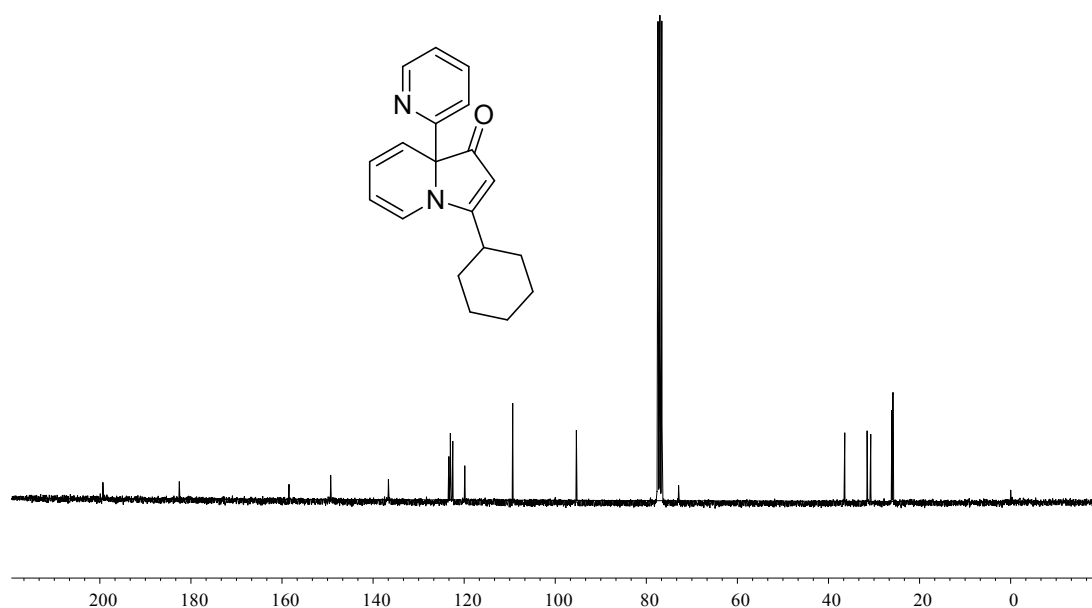
77.462
77.038
76.615
73.867



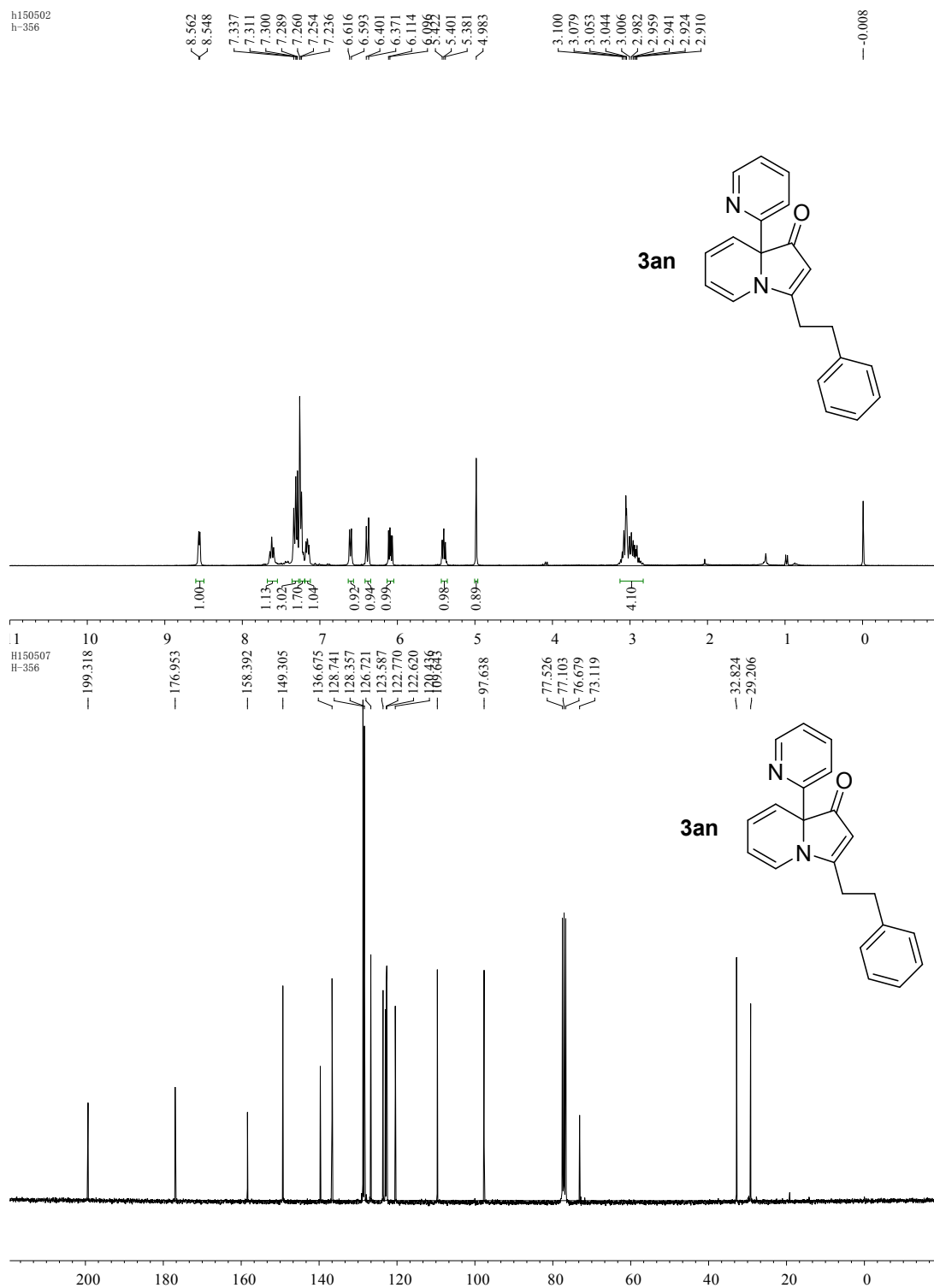
h150517
h-369



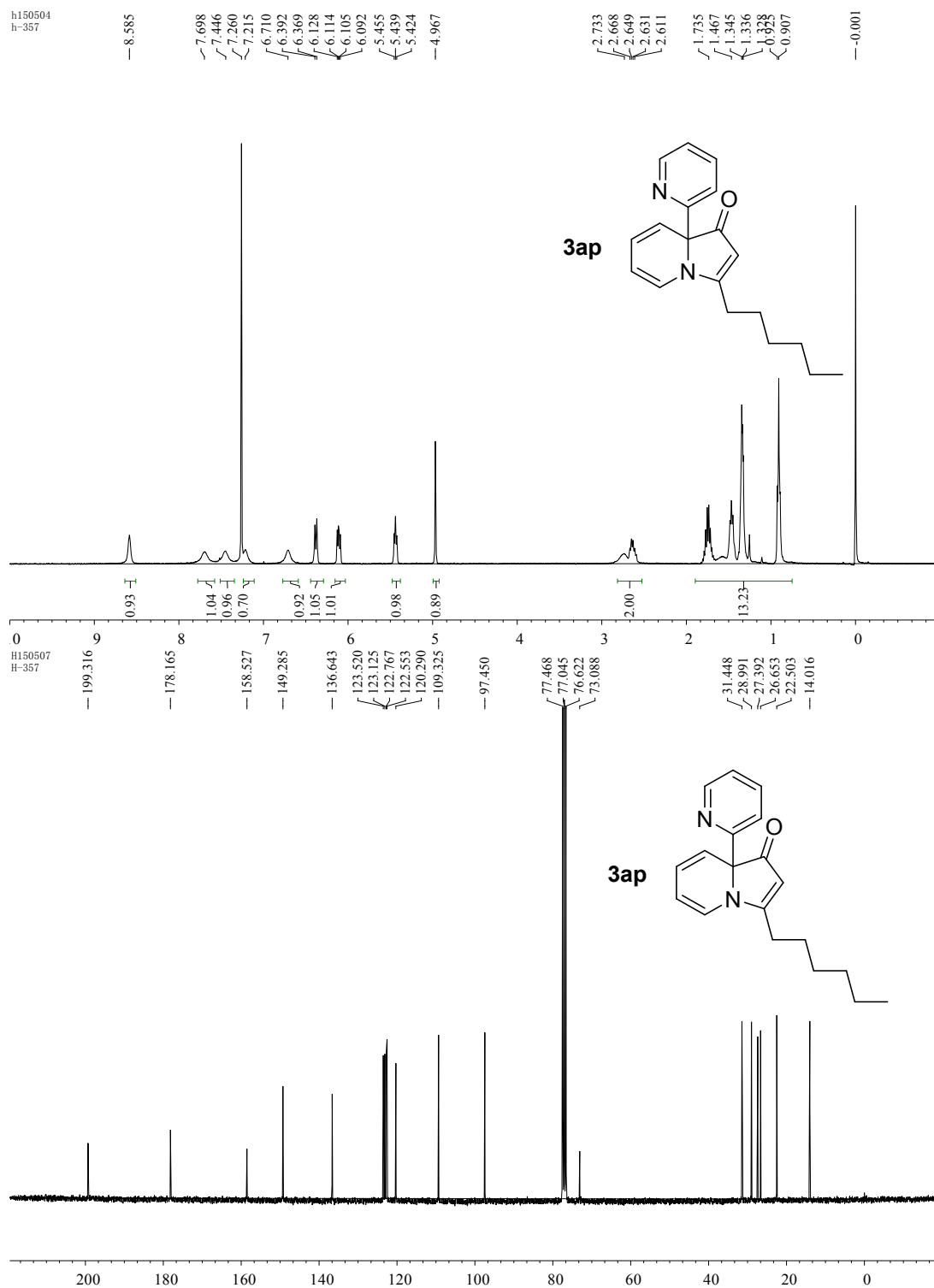
h150517
h-369



h150502
h-356



h150504
h-357

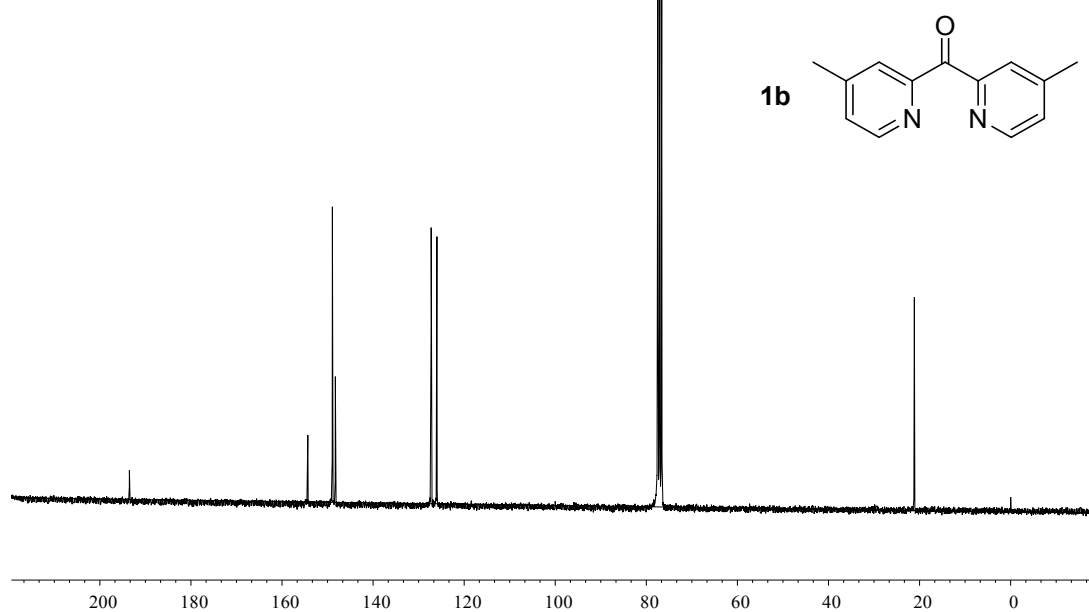
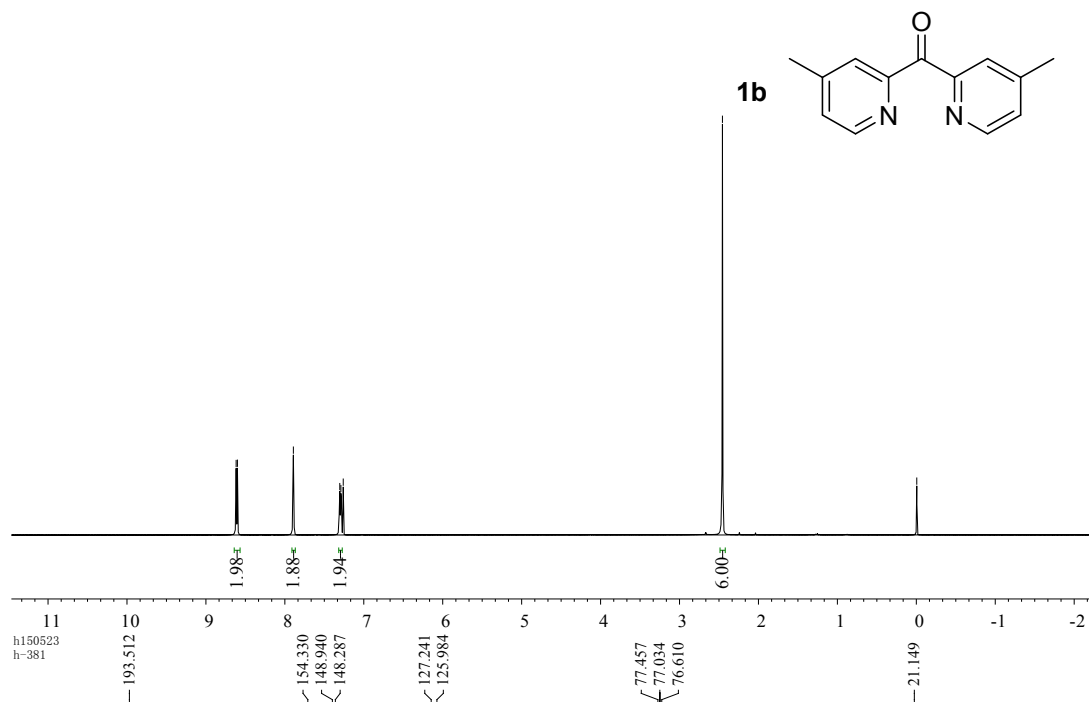


h150517
h-381

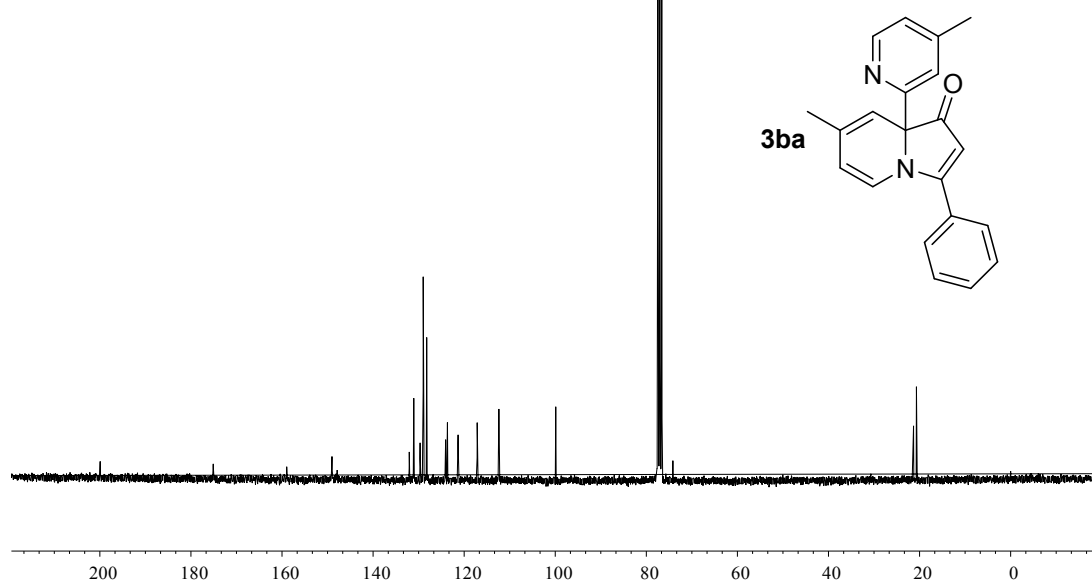
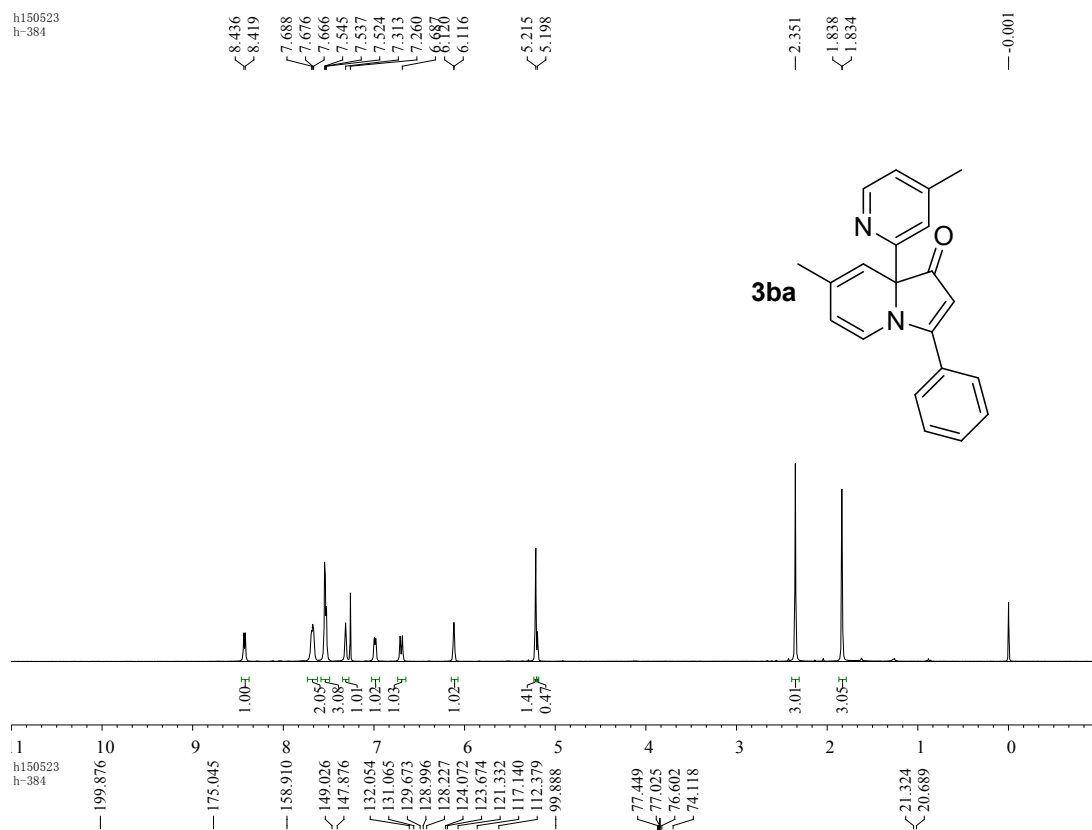
8.617
8.601
7.892
7.304
7.290
7.260

2.456

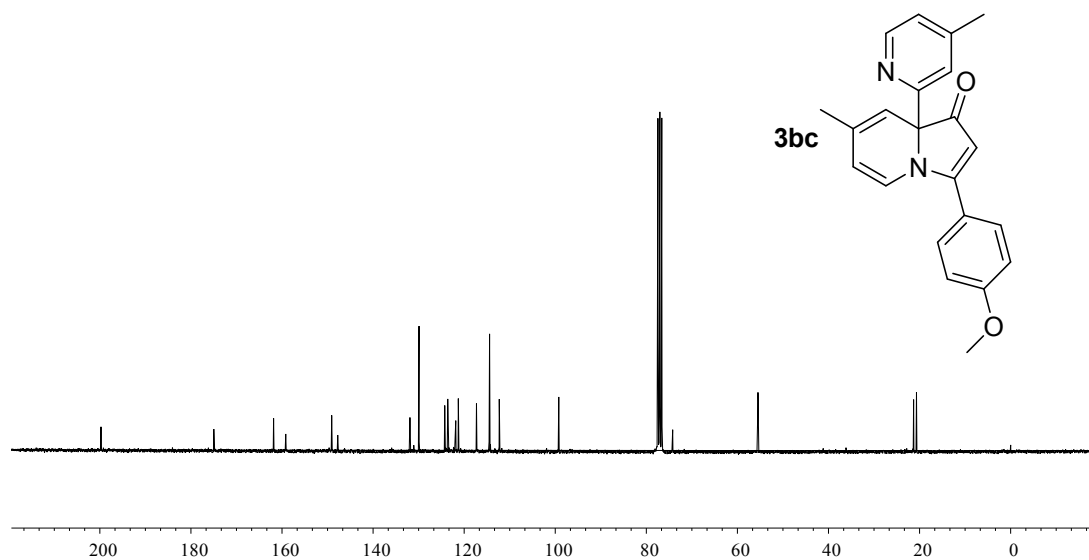
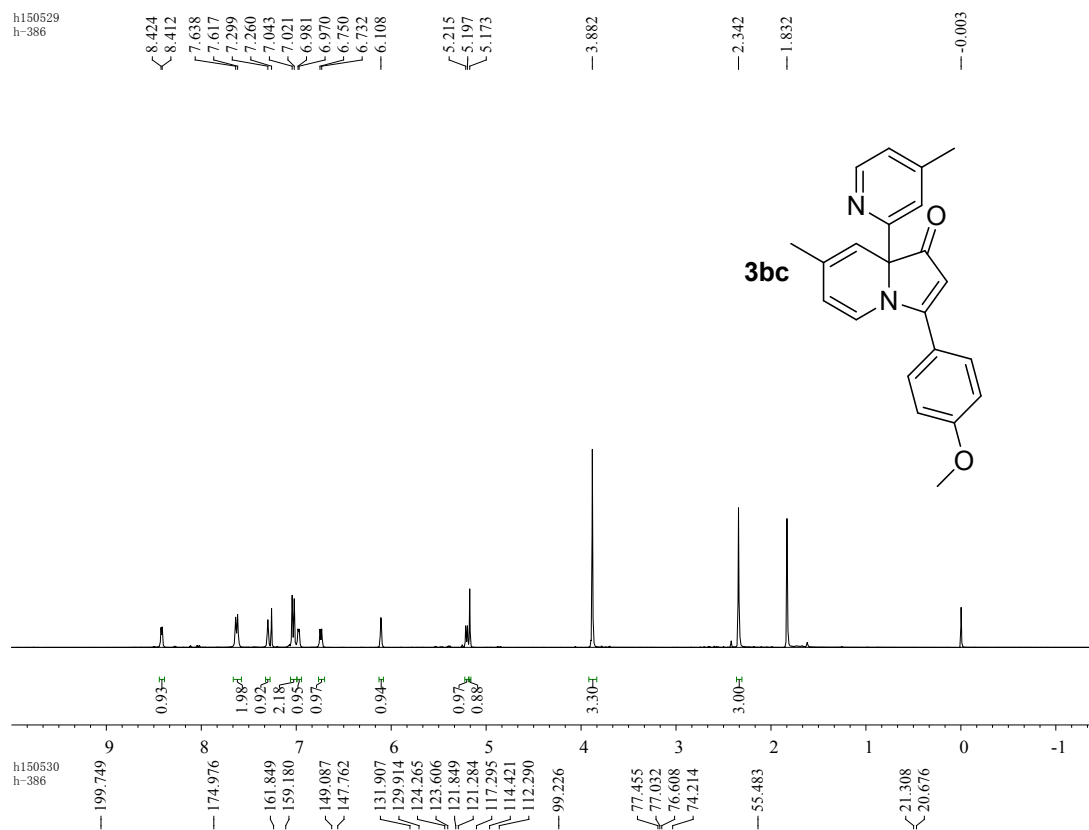
-0.006

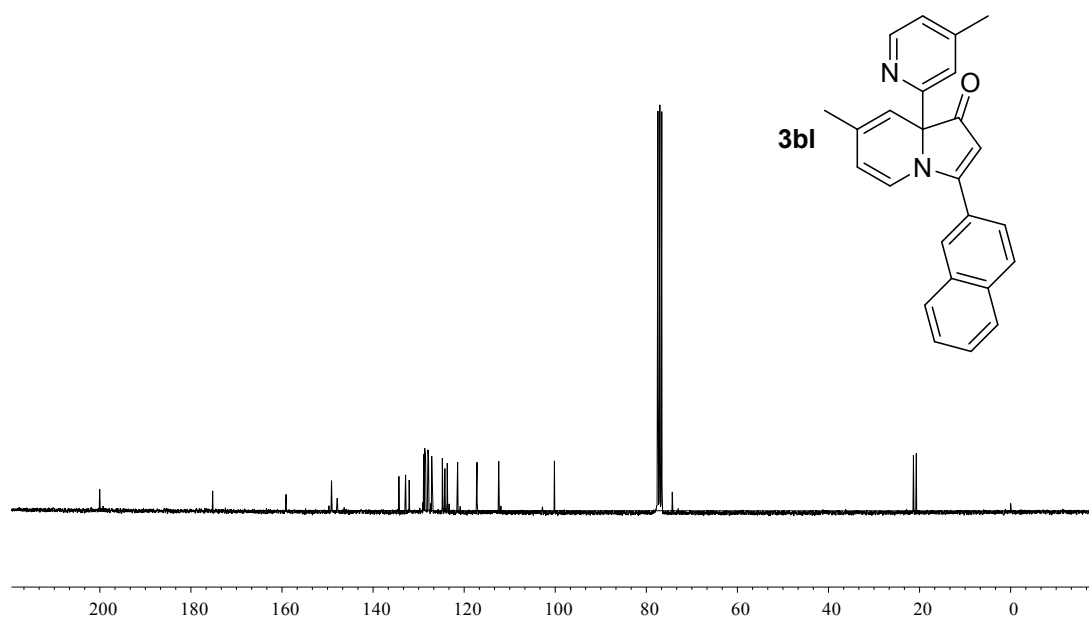
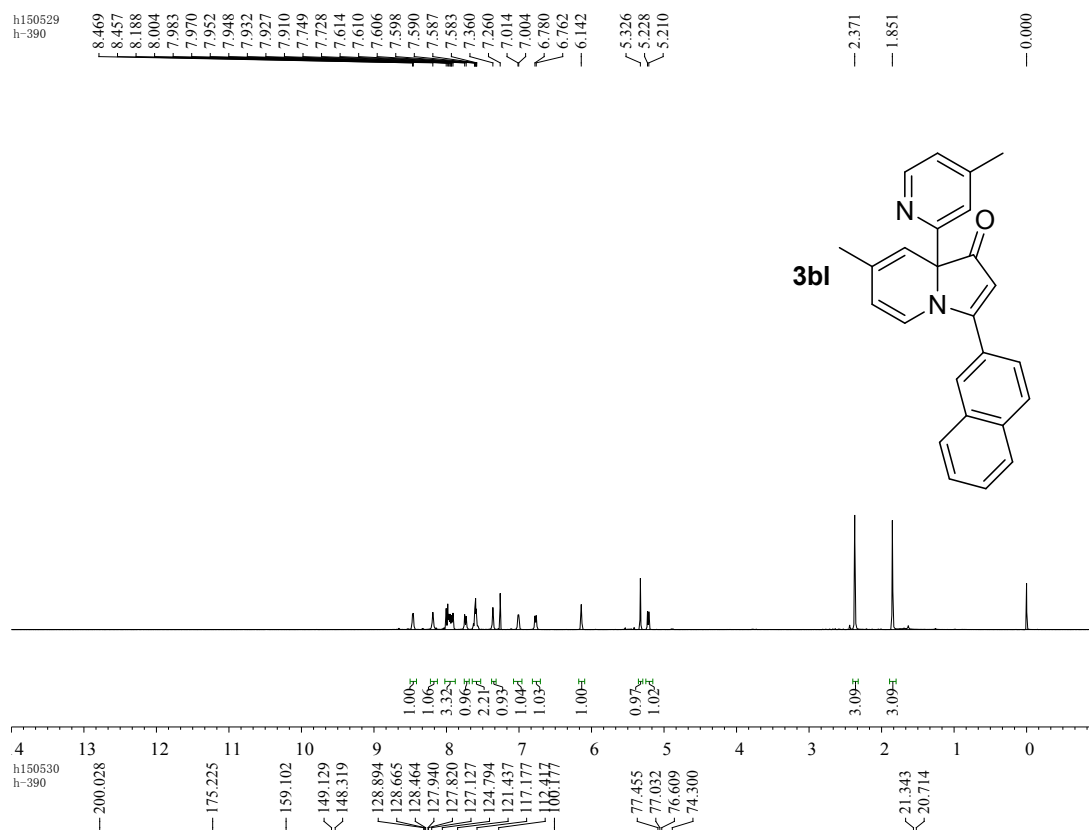


h150523
h-384

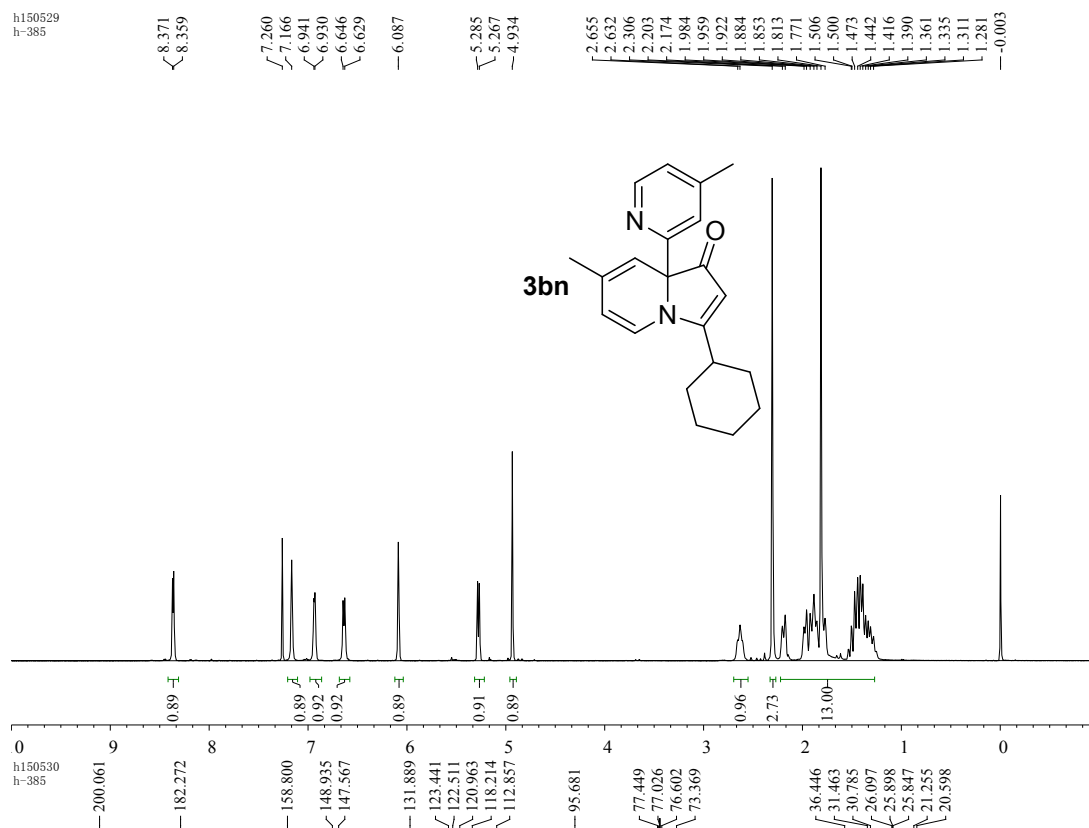


h150529
h-386

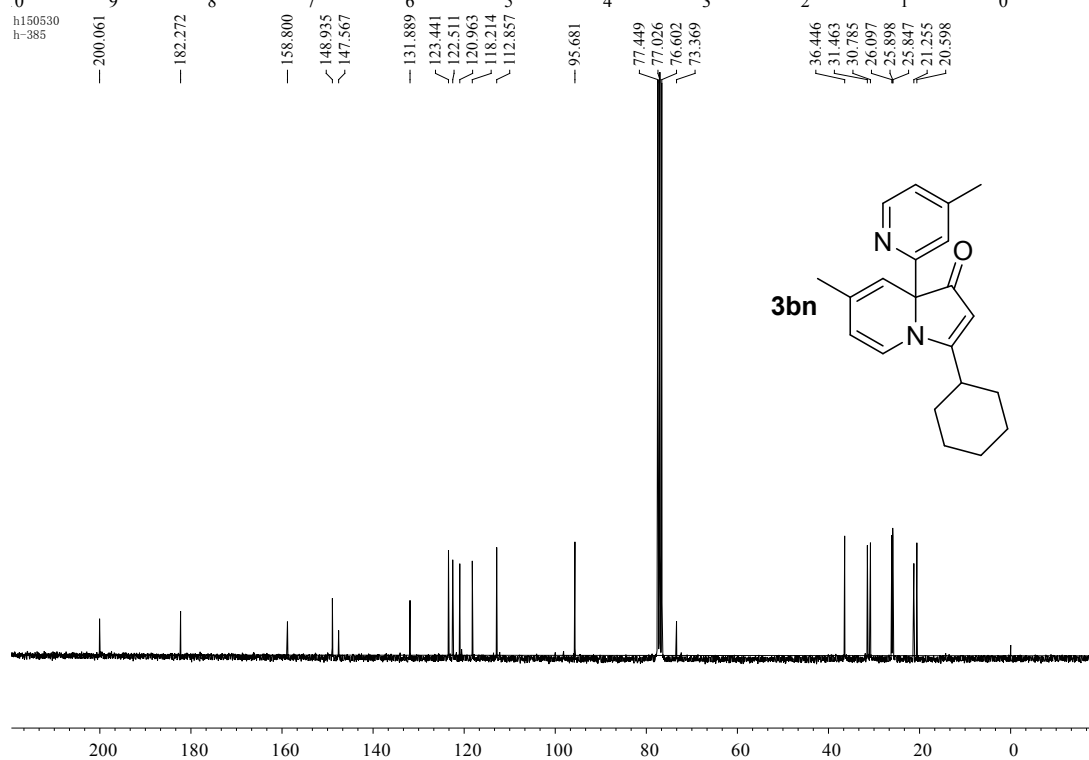




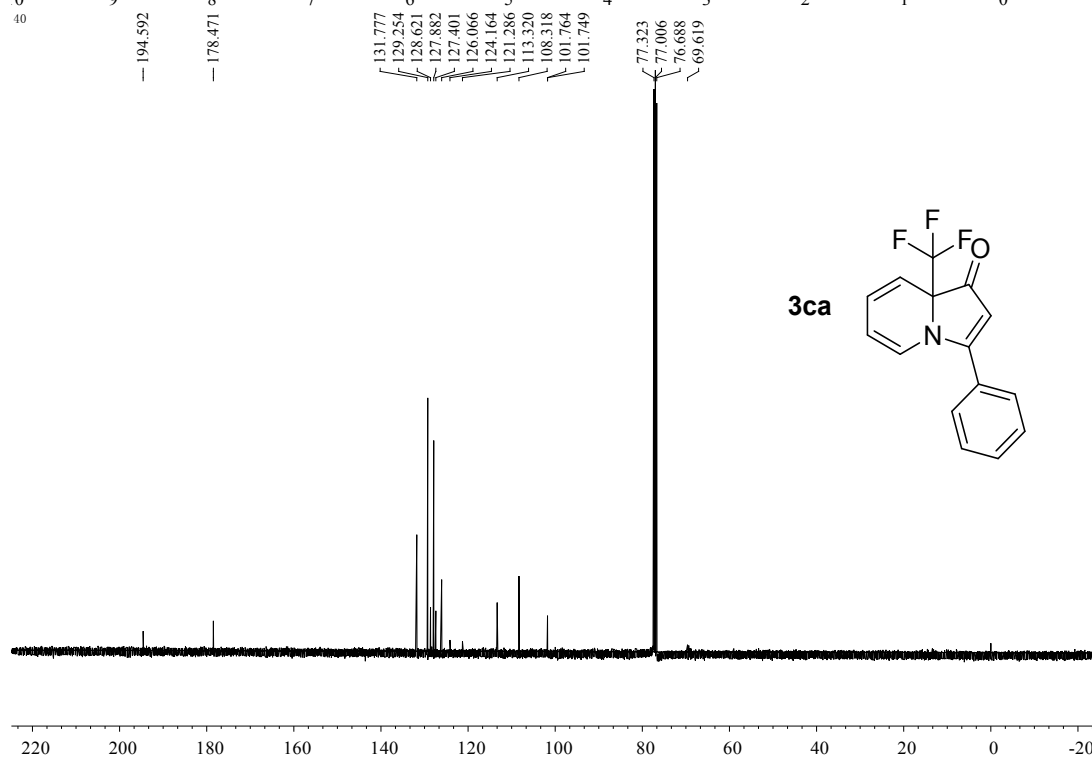
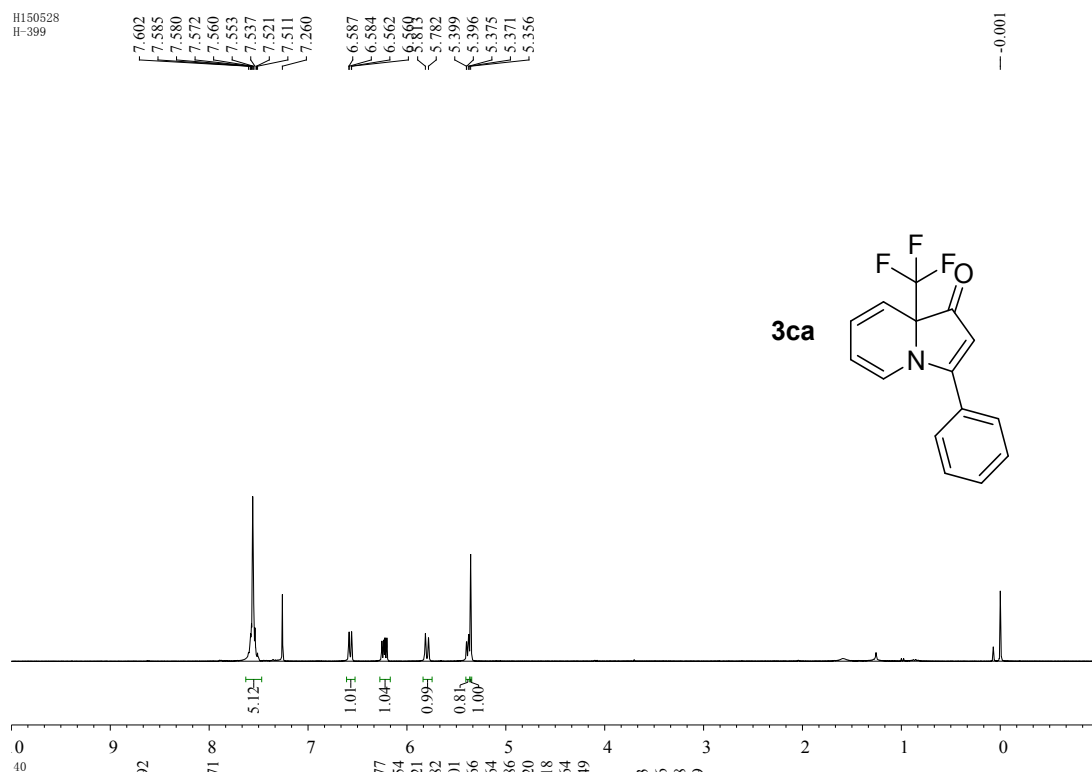
h150529
h-385



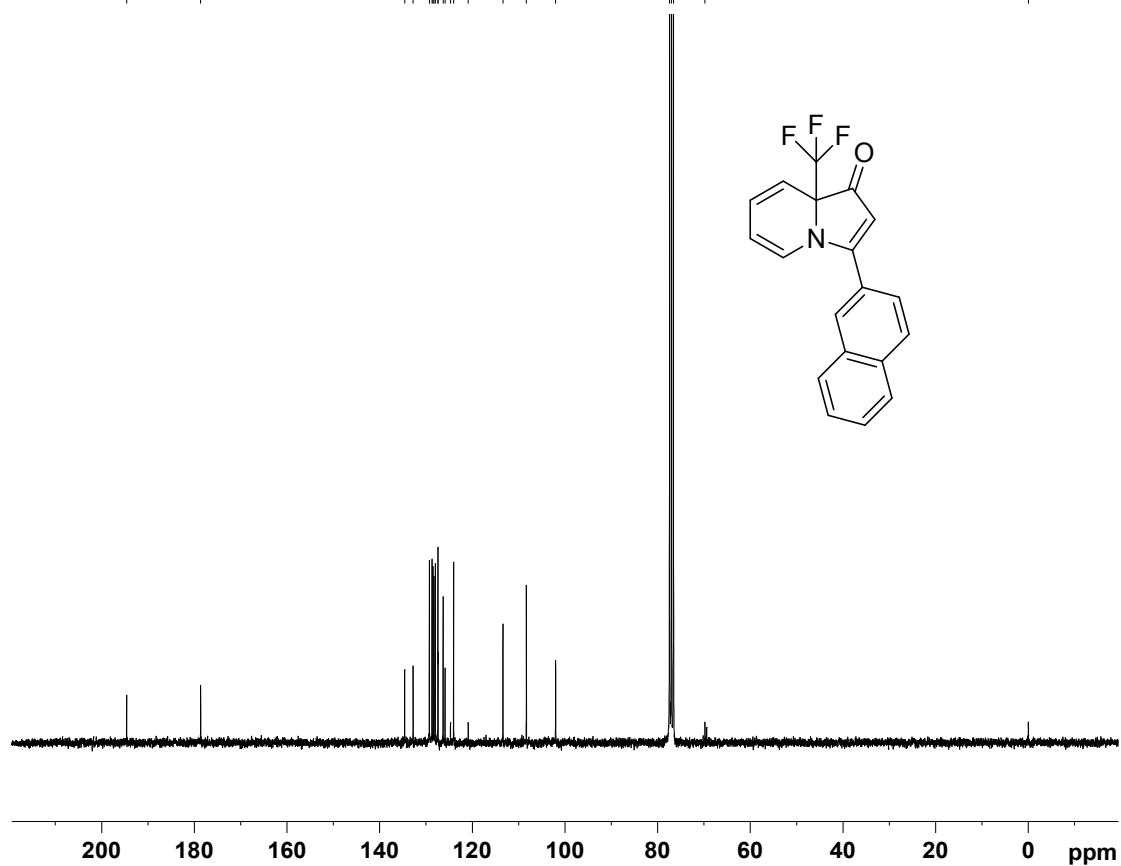
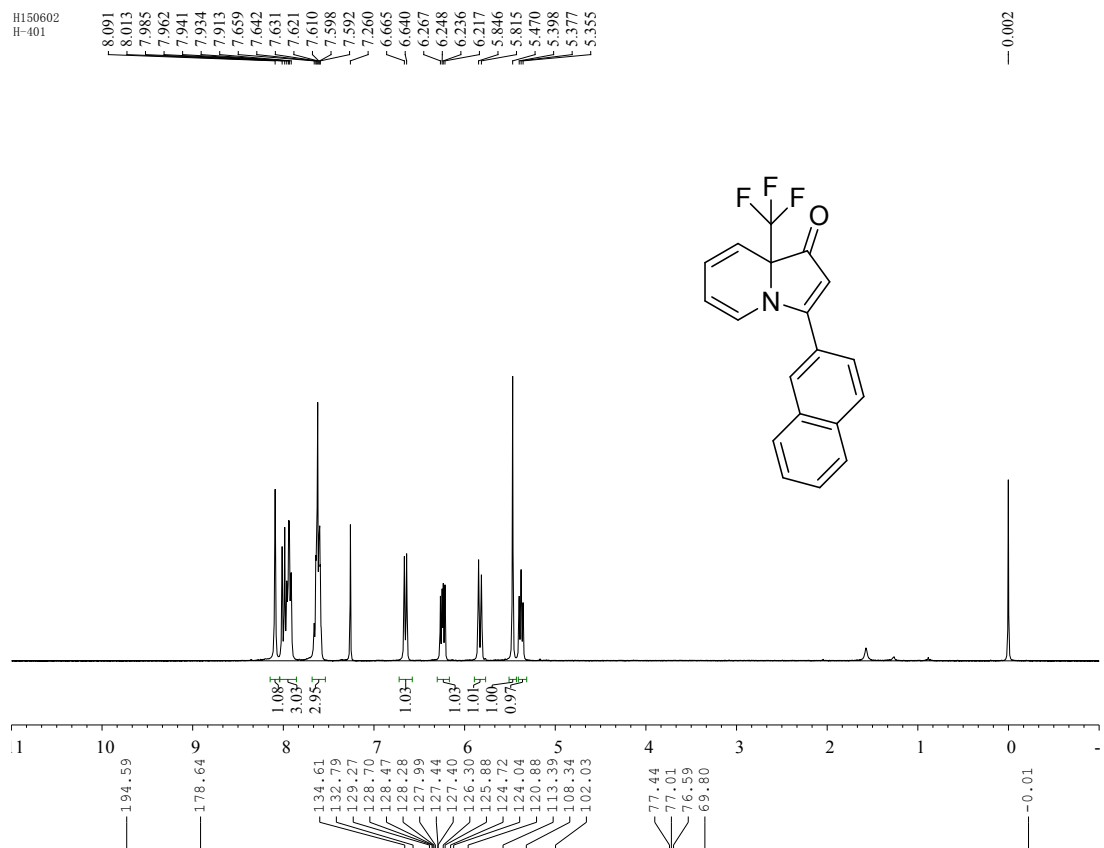
h150530
h-385



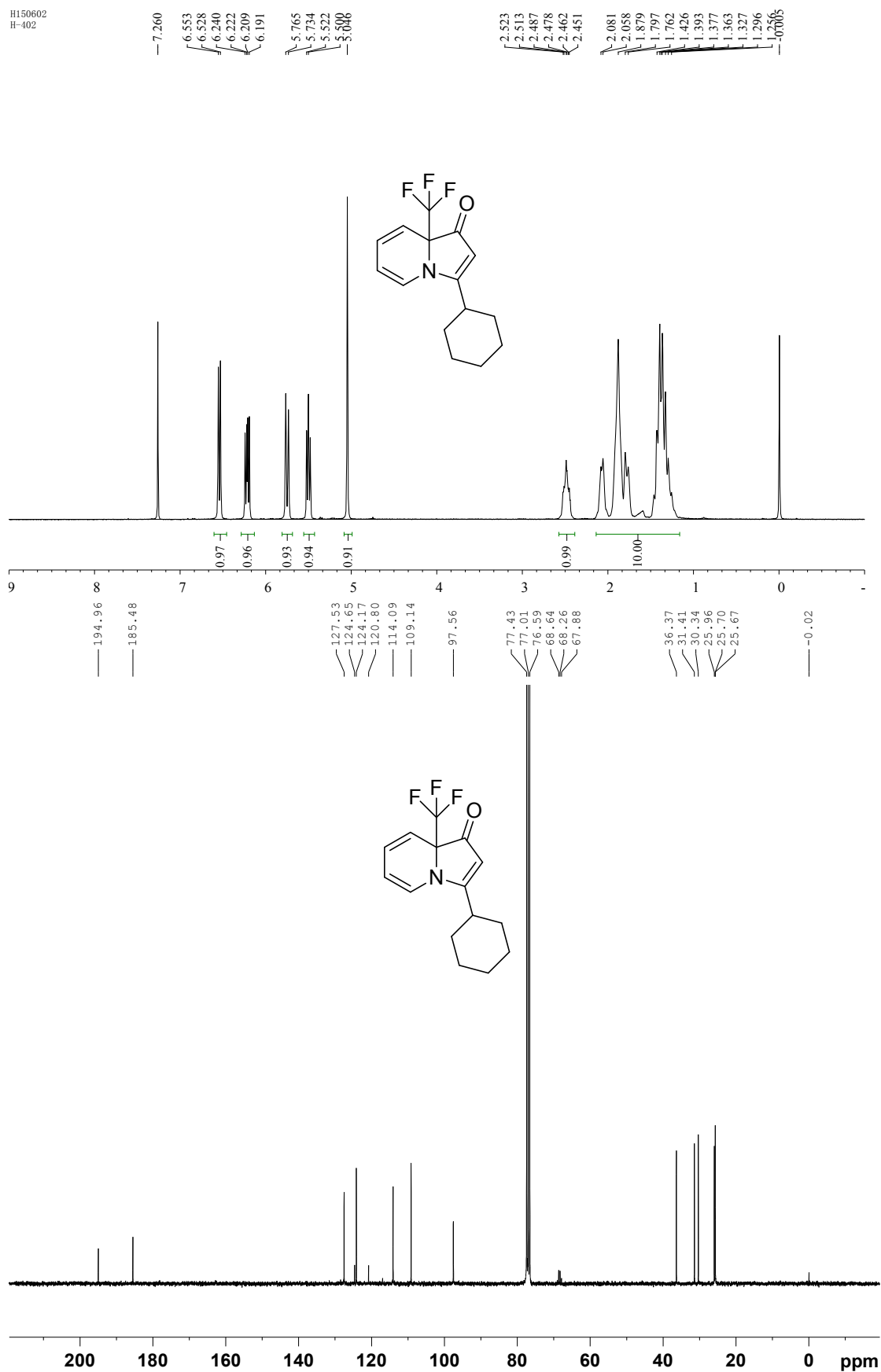
H150528
H-399



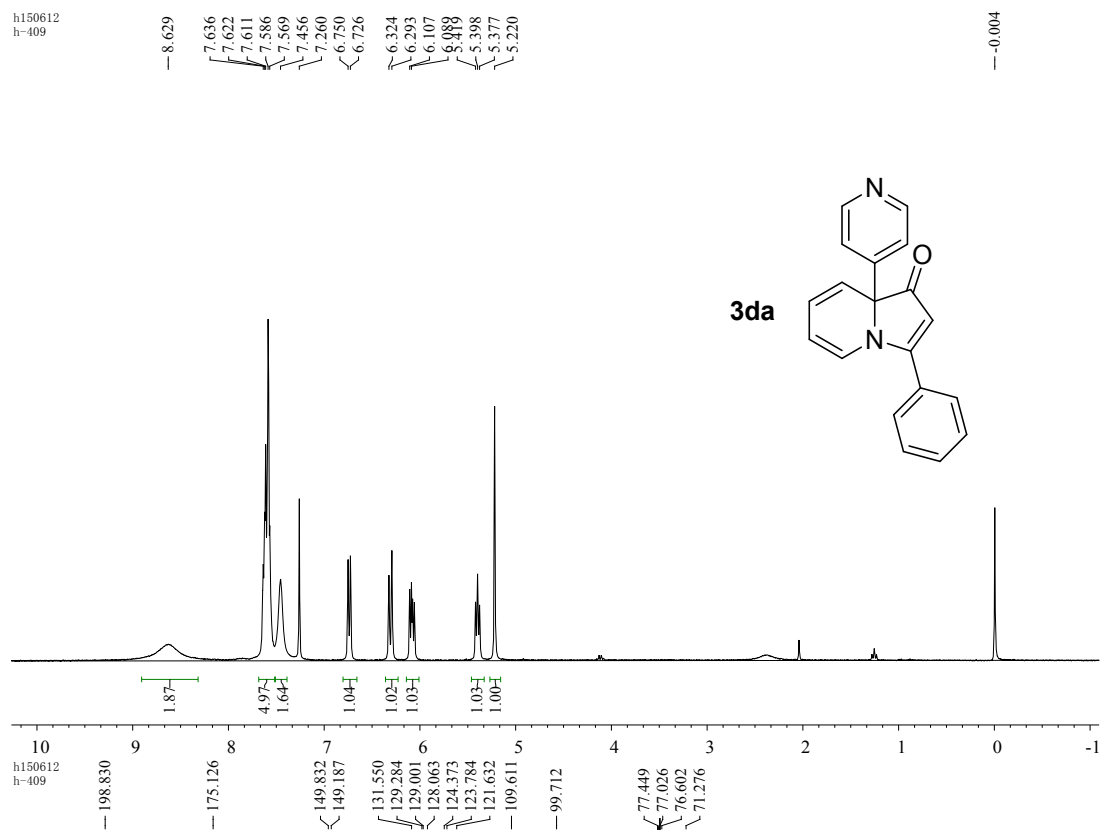
H150602
H-401



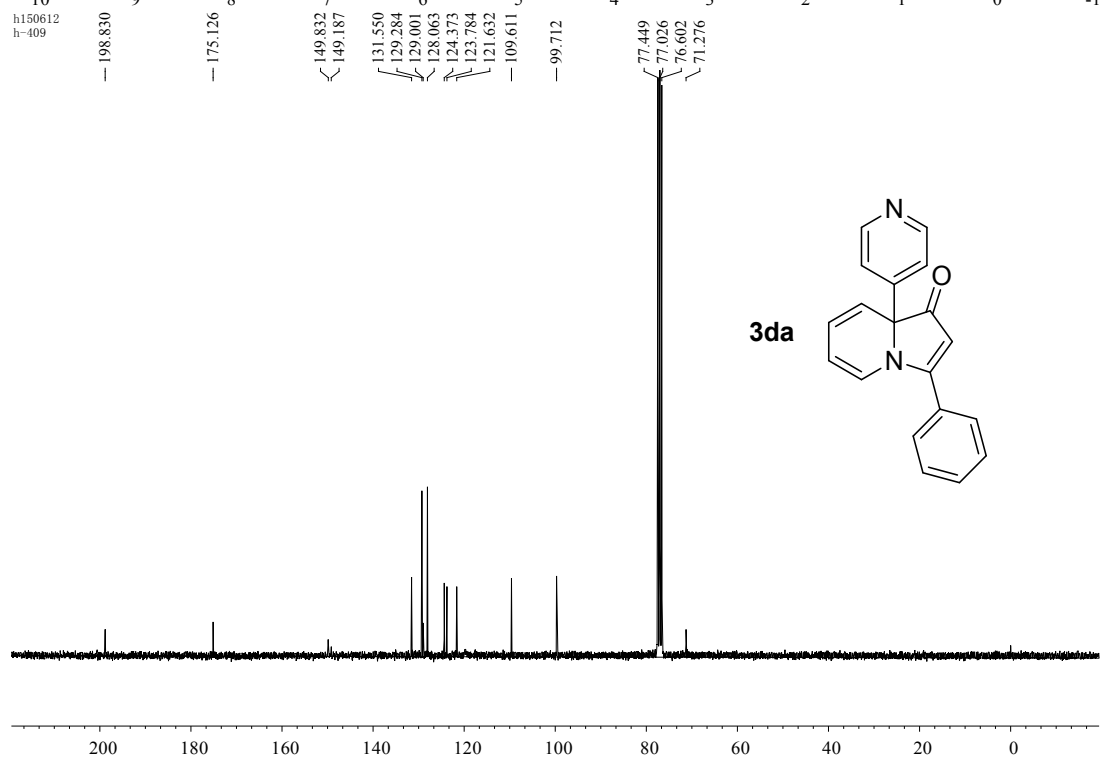
H150602
H-402



h150612
h-409



h150612
h-409

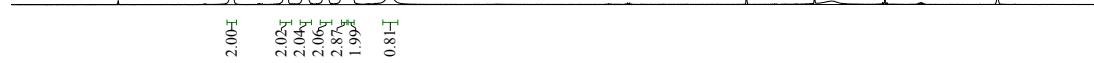
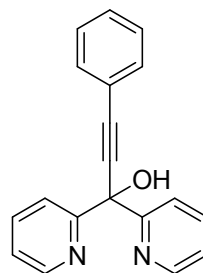


h150624
h-434

8.559
8.545
7.962
7.935
7.755
7.750
7.730
7.725
7.704
7.698
7.519
7.512
7.500
7.497
7.486
7.304
7.298
7.287
7.281
7.267
7.260
7.238
7.221
7.216
7.199
6.785

-0.000

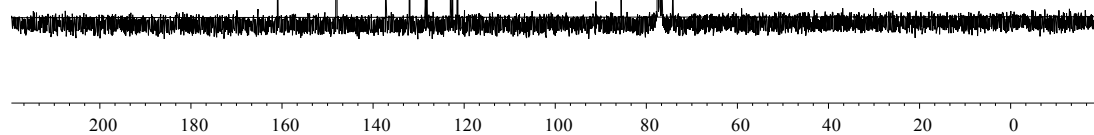
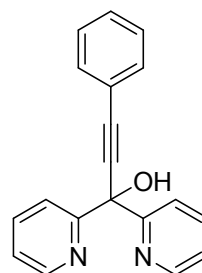
4aa



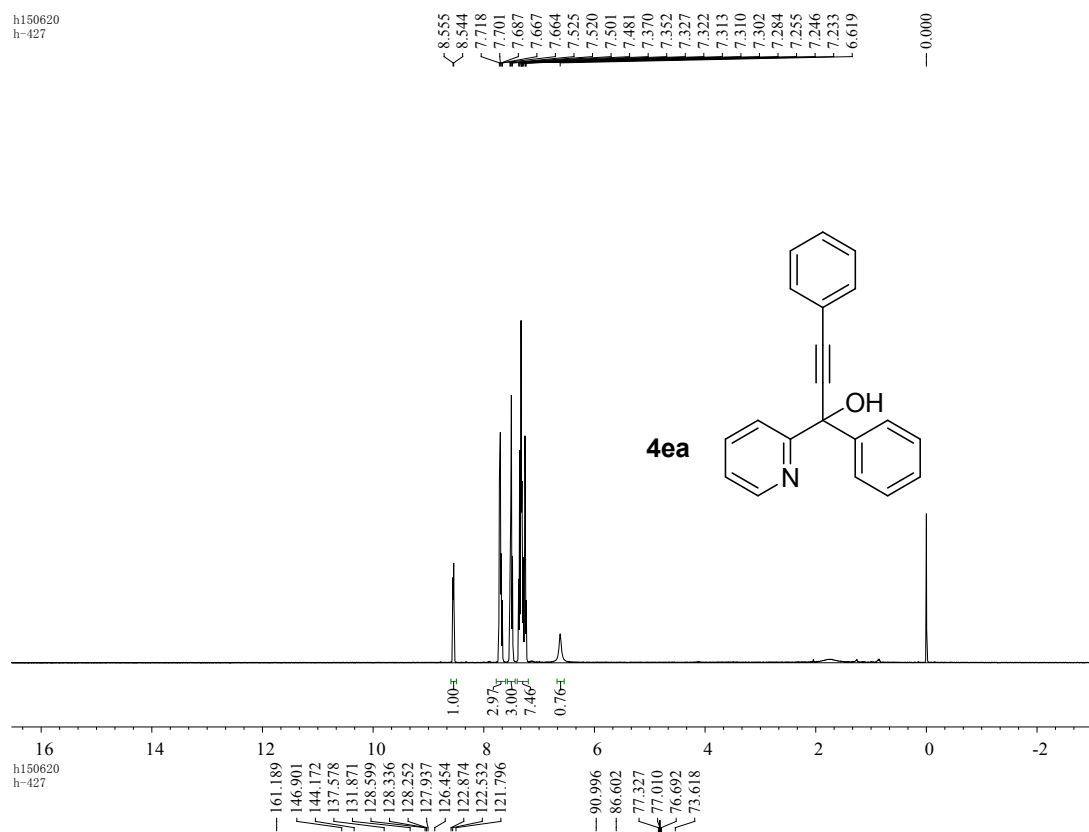
11 10 9 8 7 6 5 4 3 2 1 0 -1
h150624
h-434

160.980
148.028
137.152
132.004
128.542
128.157
122.893
122.535
121.440
91.079
85.541
77.462
77.039
76.616
74.188

4aa



h150620
h-427



h150620
h-427

