

Supporting information

Electrocatalytic Selenocyclization for the Synthesis of Selenium-Containing Oxazolidinones

Yiyi Chen,^a Yi Xu,^a Zihao Ma,^a Hanlin Wang,^a Qisheng Chen,^a Shuangquan Zhang,^a Kun Feng,^a Xiaohui Chen,^{b*} Xianqiang Kong,^{a*}

^[a] School of Chemical Engineering and Materials, Changzhou Institute of Technology, No. 666 Liaohe Road, Changzhou 213032, China

^[b] School of Energy and Materials, Shanghai Polytechnic University, No. 2360 Jinhai Road, Shanghai 201209, China

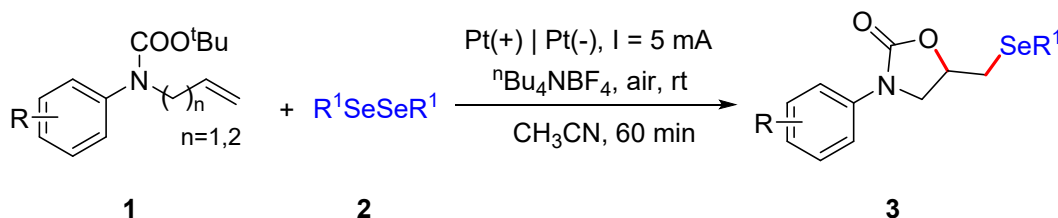
E-mail: kongxq@czu.cn; xhchen@sspu.edu.cn

Table of contents	Page
1. General information	S3
2. General procedure	S3
3. Procedure for gram-scale experiment	S3
4. Mechanistic studies	S16
5. Copies of NMR spectra	S17

1. General information

Commercial reagents and solvents were obtained from the commercial providers and used without further purification. The products were purified using a commercial flash chromatography system or a regular glass column. TLC was developed on silica gel 60 F254 glassplates. ^1H NMR (500 MHz), ^{13}C NMR (151 MHz) and ^{19}F NMR (376 MHz) spectra were recorded on a Bruker NMR apparatus. The chemical shifts are reported in δ (ppm) values (^1H and ^{13}C NMR relative to CDCl_3 , δ 7.26 ppm for ^1H NMR and δ 77.0 ppm for ^{13}C NMR). Or alternatively, ^1H NMR chemical shifts were referenced to tetramethylsilane signal (0 ppm). Multiplicities are recorded by s (singlet), d (doublet), t (triplet), q (quartet), p (pentet), h (hextet), m (multiplet), dd (doublet of doublets), dt (doublet of triplets), td (triplet of doublets), tq (triplet of quartets), ddd (doublet of doublet of doublets), and br (broad). Coupling constants (J), are reported in Hertz (Hz).

2. General Procedure for Experiments

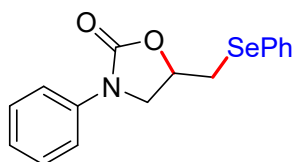


An undivided cell was equipped with Pt anode (1.0 cm $\times$ 1.0 cm $\times$ 0.1 cm) and Pt cathode (1.0 cm $\times$ 1.0 cm $\times$ 0.1 cm). To the cell was added **1** (0.2 mmol), **2** (0.24 mmol), $n\text{Bu}_4\text{NBF}_4$ (0.1 mmol), CH_3CN (4.0 mL). The mixture was electrolyzed using constant current conditions ($I = 5 \text{ mA}$) at room temperature under magnetic stirring for 1 h, the solvent was removed under reduced pressure. The residue was poured into a saturated aqueous solution of NaCl and the product was then extracted with CH_2Cl_2 (3 $\times$ 20 mL), dried over Na_2SO_4 , and concentrated in vacuo. The residue was purified by column chromatography on silica gel using a mixture of petroleum ether/EtOA eluent to afford the desired pure product **3**.

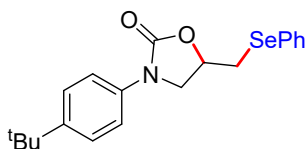
3. General Procedure for Gram-Scale Experiments

An undivided cell was equipped with Pt anode (1.0 cm $\times$ 1.0 cm $\times$ 0.1 cm) and Pt cathode (1.0 cm $\times$ 1.0 cm $\times$ 0.1 cm). To the cell was added **1a** (1.0 mmol, 0.233 g), **2a** (1.2 mmol, 0.374 g), $n\text{Bu}_4\text{NBF}_4$ (0.5 mmol, 0.165 g), CH_3CN (20.0 mL). The mixture was electrolyzed using constant current conditions ($I =$

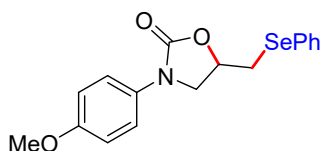
5 mA) at room temperature under magnetic stirring for 5 h. Then, the solvent was removed under reduced pressure. The residue was poured into a saturated aqueous solution of NaCl and the product was then extracted with CH₂Cl₂ (3×20 mL), dried over Na₂SO₄, and concentrated in vacuo. The residue was purified by column chromatography on silica gel using a mixture of petroleum ether/EtOA eluent to afford the desired pure product **3a** (0.3 g, 90%).



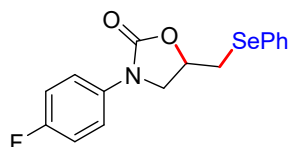
3-phenyl-5-((phenylselanyl)methyl)oxazolidin-2-one (3a), 59.9 mg, 90% yield. ¹H NMR (500 MHz, CDCl₃) δ 7.56 (dd, *J* = 7.5, 2.1 Hz, 2H), 7.48 (d, *J* = 7.9 Hz, 2H), 7.41 - 7.26 (m, 5H), 7.12 (t, *J* = 7.4 Hz, 1H), 4.71 (tdd, *J* = 8.8, 6.3, 4.3 Hz, 1H), 4.09 (t, *J* = 8.8 Hz, 1H), 3.77 (dd, *J* = 9.1, 6.4 Hz, 1H), 3.36 (dd, *J* = 12.9, 4.3 Hz, 1H), 3.05 (dd, *J* = 12.9, 9.1 Hz, 1H). ¹³C NMR (151 MHz, CDCl₃) δ 154.4, 138.2, 133.7, 129.6, 129.1, 128.1, 128.0, 124.1, 118.2, 71.8, 50.2, 31.1. HRMS (ESI) *m/z*: [M + H]⁺ Calcd for C₁₆H₁₆NO₂Se 334.0346; Found 334.0357. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



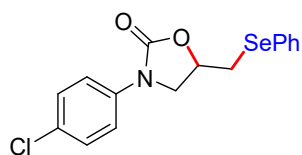
3-(4-(tert-butyl)phenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3b), 70.8 mg, 91% yield. ¹H NMR (500 MHz, CDCl₃) δ 7.56 (d, *J* = 7.8 Hz, 2H), 7.38 (q, *J* = 8.9 Hz, 4H), 7.33 - 7.26 (m, 3H), 4.70 (dtd, *J* = 13.0, 8.9, 7.6, 4.4 Hz, 1H), 4.08 (t, *J* = 8.8 Hz, 1H), 3.75 (dd, *J* = 9.1, 6.3 Hz, 1H), 3.35 (dd, *J* = 12.9, 4.4 Hz, 1H), 3.04 (dd, *J* = 12.9, 9.0 Hz, 1H), 1.30 (s, 9H). ¹³C NMR (151 MHz, CDCl₃) δ 154.5, 147.2, 135.6, 133.7, 129.6, 128.1, 128.0, 126.0, 118.1, 71.8, 50.4, 34.4, 31.4, 31.1. HRMS (ESI) *m/z*: [M + H]⁺ Calcd for C₂₀H₂₄NO₂Se 390.0972; Found 390.0973. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



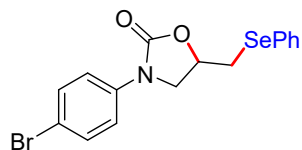
3-(4-methoxyphenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3c), 63.2 mg, 87% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.4, 2.2$ Hz, 2H), 7.37 (d, $J = 9.1$ Hz, 2H), 7.34 - 7.26 (m, 3H), 6.88 (d, $J = 9.1$ Hz, 2H), 4.70 (tdd, $J = 8.7, 6.4, 4.3$ Hz, 1H), 4.05 (t, $J = 8.8$ Hz, 1H), 3.78 (s, 3H), 3.73 (dd, $J = 9.0, 6.4$ Hz, 1H), 3.35 (dd, $J = 12.9, 4.4$ Hz, 1H), 3.05 (dd, $J = 12.8, 9.1$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 156.5, 154.8, 133.7, 131.4, 129.6, 128.2, 128.0, 120.4, 114.4, 71.8, 55.6, 50.9, 31.2. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{18}\text{NO}_3\text{Se}$ 364.0452; Found 364.0454. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



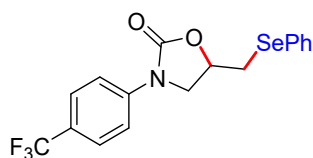
3-(4-fluorophenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3d), 58.3 mg, 83% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.5, 2.0$ Hz, 2H), 7.48 - 7.40 (m, 2H), 7.35 - 7.26 (m, 3H), 7.04 (dd, $J = 9.2, 8.1$ Hz, 2H), 4.72 (tdd, $J = 8.7, 6.4, 4.3$ Hz, 1H), 4.07 (t, $J = 8.7$ Hz, 1H), 3.74 (dd, $J = 9.0, 6.4$ Hz, 1H), 3.36 (dd, $J = 12.9, 4.3$ Hz, 1H), 3.05 (dd, $J = 12.9, 9.0$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 159.3 (d, $J = 244.1$ Hz), 154.5, 134.3 (d, $J = 2.8$ Hz), 133.7, 129.6, 128.1, 127.9, 120.0 (d, $J = 7.9$ Hz), 115.8 (d, $J = 22.5$ Hz), 71.6, 50.5, 31.0. ^{19}F NMR (376 MHz, CDCl_3) δ -118.37. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{16}\text{H}_{15}\text{FNO}_2\text{Se}$ 352.0252; Found 352.0250. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



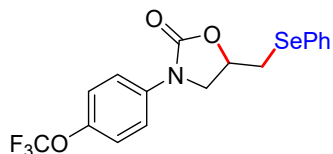
3-(4-chlorophenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3e), 65.3 mg, 89% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.5, 2.0$ Hz, 2H), 7.44 (d, $J = 9.0$ Hz, 2H), 7.39 - 7.26 (m, 5H), 4.73 (tdd, $J = 8.8, 6.3, 4.2$ Hz, 1H), 4.08 (t, $J = 8.8$ Hz, 1H), 3.75 (dd, $J = 9.1, 6.4$ Hz, 1H), 3.37 (dd, $J = 12.9, 4.3$ Hz, 1H), 3.06 (dd, $J = 12.9, 9.1$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.3, 136.8, 133.8, 129.7, 129.4, 129.2, 128.3, 127.9, 119.4, 71.9, 50.3, 31.1. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{16}\text{H}_{15}\text{ClNO}_2\text{Se}$ 367.9957; Found 367.9958. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



(4-bromophenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3f), 71.5 mg, 87% yield. $^1\text{H NMR}$ (500 MHz, CDCl_3) δ 7.55 (d, J = 6.5 Hz, 2H), 7.50 - 7.41 (m, 2H), 7.41 - 7.34 (m, 2H), 7.29 (d, J = 6.6 Hz, 3H), 4.71 (dtt, J = 8.7, 4.5, 2.2 Hz, 1H), 4.03 (dd, J = 8.8, 4.6 Hz, 1H), 3.71 (q, J = 6.1 Hz, 1H), 3.34 (dd, J = 12.9, 3.9 Hz, 1H), 3.04 (dd, J = 12.9, 9.0 Hz, 1H). $^{13}\text{C NMR}$ (151 MHz, CDCl_3) δ 154.2, 137.3, 133.7, 132.0, 129.6, 128.2, 127.8, 119.7, 116.9, 71.8, 50.1, 31.0. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{16}\text{H}_{15}\text{BrNO}_2\text{Se}$ 411.9451; Found 411.9450. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

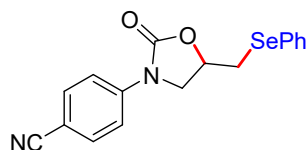


5-((phenylselanyl)methyl)-3-(4-(trifluoromethyl)phenyl)oxazolidin-2-one (3g), 70.6 mg, 88% yield. $^1\text{H NMR}$ (500 MHz, CDCl_3) δ 7.61 (s, 4H), 7.57 (dd, J = 7.8, 1.9 Hz, 2H), 7.38 - 7.27 (m, 3H), 4.77 (tdd, J = 8.8, 6.3, 4.2 Hz, 1H), 4.13 (t, J = 8.8 Hz, 1H), 3.79 (dd, J = 9.1, 6.3 Hz, 1H), 3.39 (dd, J = 13.0, 4.2 Hz, 1H), 3.08 (dd, J = 13.0, 9.1 Hz, 1H). $^{13}\text{C NMR}$ (151 MHz, CDCl_3) δ 154.2, 141.2, 133.9, 129.7, 128.4, 127.8, 126.4 (q, J = 2.0 Hz), 117.8, 72.0, 50.1, 31.1. $^{19}\text{F NMR}$ (376 MHz, CDCl_3) δ -62.16. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{15}\text{F}_3\text{NO}_2\text{Se}$ 402.0220; Found 402.0222. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

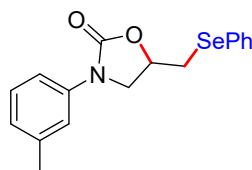


5-((phenylselanyl)methyl)-3-(4-(trifluoromethoxy)phenyl)oxazolidin-2-one (3h), 68.4 mg, 82% yield. $^1\text{H NMR}$ (500 MHz, CDCl_3) δ 7.56 (dd, J = 7.6, 1.9 Hz, 2H), 7.53 - 7.46 (m, 2H), 7.37 - 7.26 (m, 3H), 7.20 (d, J = 8.7 Hz, 2H), 4.73 (tdd, J = 8.7, 6.3, 4.3 Hz, 1H), 4.08 (t, J = 8.7 Hz, 1H), 3.75 (dd, J = 9.0, 6.4 Hz, 1H), 3.36 (dd, J = 12.9, 4.3 Hz, 1H), 3.06 (dd, J = 13.0, 9.0 Hz, 1H). $^{13}\text{C NMR}$ (151 MHz, CDCl_3) δ 154.4, 145.3, 136.9, 133.8, 133.8, 129.7, 128.3, 127.9, 121.9, 119.3, 71.9, 50.3, 31.0. $^{19}\text{F NMR}$ (376 MHz, CDCl_3) δ -58.13. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{15}\text{NF}_3\text{O}_3\text{Se}$ 418.0169; Found 418.0168.

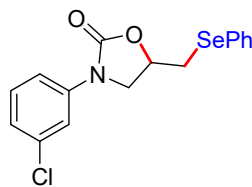
The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



4-(2-oxo-5-((phenylselanyl)methyl)oxazolidin-3-yl)benzonitrile (3i), 64.4 mg, 90% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.62 (d, $J = 2.3$ Hz, 4H), 7.56 (dd, $J = 7.8, 1.8$ Hz, 2H), 7.38 - 7.27 (m, 3H), 4.78 (tdd, $J = 8.7, 6.3, 4.2$ Hz, 1H), 4.12 (t, $J = 8.8$ Hz, 1H), 3.78 (dd, $J = 9.2, 6.4$ Hz, 1H), 3.38 (dd, $J = 13.0, 4.2$ Hz, 1H), 3.08 (dd, $J = 13.0, 9.0$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 153.8, 142.0, 133.8, 133.7, 133.7, 133.2, 129.6, 128.3, 127.7, 118.7, 117.9, 106.9, 72.0, 49.8, 30.9. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{15}\text{N}_2\text{O}_2\text{Se}$ 359.0299; Found 359.0298. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

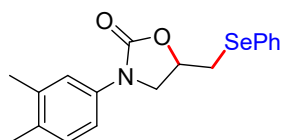


5-((phenylselanyl)methyl)-3-(m-tolyl)oxazolidin-2-one (3j), 59.0 mg, 85% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.4, 2.2$ Hz, 2H), 7.36 - 7.22 (m, 6H), 6.94 (d, $J = 6.8$ Hz, 1H), 4.70 (tdd, $J = 8.7, 6.3, 4.3$ Hz, 1H), 4.08 (t, $J = 8.8$ Hz, 1H), 3.76 (dd, $J = 9.1, 6.4$ Hz, 1H), 3.35 (dd, $J = 12.9, 4.3$ Hz, 1H), 3.04 (dd, $J = 12.8, 9.1$ Hz, 1H), 2.35 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.4, 139.1, 138.1, 133.7, 133.7, 133.7, 129.6, 128.9, 128.1, 128.0, 125.0, 119.0, 115.4, 71.7, 50.4, 31.1, 21.7. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{18}\text{NO}_2\text{Se}$ 348.0503; Found 348.0505. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

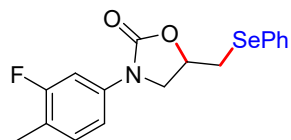


3-(3-chlorophenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3k), 61.7 mg, 84% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.61 - 7.53 (m, 2H), 7.51 (t, $J = 2.1$ Hz, 1H), 7.42 - 7.36 (m, 1H), 7.35 - 7.25 (m,

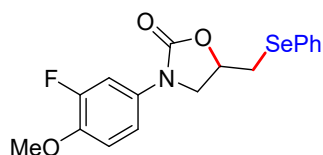
4H), 7.14 – 7.06 (m, 1H), 4.73 (tdd, $J = 8.8, 6.3, 4.2$ Hz, 1H), 4.06 (t, $J = 8.8$ Hz, 1H), 3.73 (dd, $J = 9.1, 6.3$ Hz, 1H), 3.36 (dd, $J = 13.0, 4.3$ Hz, 1H), 3.05 (dd, $J = 12.9, 9.0$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.1, 139.3, 134.9, 133.7, 130.1, 129.6, 128.2, 127.8, 124.1, 118.2, 116.1, 71.9, 50.1, 31.0. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{16}\text{H}_{15}\text{ClNO}_2\text{Se}$ 367. 9957; Found 367. 9955. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



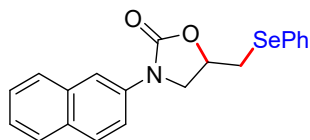
3-(3,4-dimethylphenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3l), 61.4 mg, 85% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.3, 2.2$ Hz, 2H), 7.36 - 7.26 (m, 4H), 7.17 (dd, $J = 8.2, 2.4$ Hz, 1H), 7.09 (d, $J = 8.2$ Hz, 1H), 4.68 (tdd, $J = 8.8, 6.3, 4.3$ Hz, 1H), 4.05 (t, $J = 8.8$ Hz, 1H), 3.73 (dd, $J = 9.1, 6.3$ Hz, 1H), 3.34 (dd, $J = 12.8, 4.3$ Hz, 1H), 3.03 (dd, $J = 12.9, 9.1$ Hz, 1H), 2.25 (s, 3H), 2.22 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.5, 137.4, 135.9, 133.6, 132.6, 130.1, 129.6, 128.1, 128.0, 119.8, 115.9, 71.7, 50.5, 31.1, 20.1, 19.2. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{18}\text{H}_{20}\text{NO}_2\text{Se}$ 362.0659; Found 362.0657. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



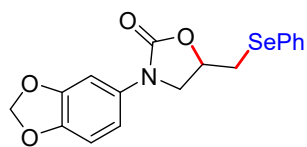
3-(3-fluoro-4-methylphenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3m), 62.8 mg, 86% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (d, $J = 7.2$ Hz, 2H), 7.30 (dd, $J = 7.4, 4.3$ Hz, 4H), 7.18 - 7.01 (m, 2H), 4.72 (ddt, $J = 8.9, 5.9, 3.0$ Hz, 1H), 4.05 (dd, $J = 8.8, 6.0$ Hz, 1H), 3.78 - 3.67 (m, 1H), 3.35 (dd, $J = 9.3, 3.7$ Hz, 1H), 3.04 (ddd, $J = 12.2, 9.0, 2.9$ Hz, 1H), 2.23 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 161.3 (d, $J = 244.4$ Hz), 154.2, 137.3, 137.3, 133.7, 131.6 (d, $J = 6.4$ Hz), 129.6, 128.2, 127.9, 120.4, 120.3, 113.2 (d, $J = 3.4$ Hz), 105.7 (d, $J = 27.9$ Hz), 71.8, 50.3, 31.1, 14.1. ^{19}F NMR (376 MHz, CDCl_3) δ -114.93. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{17}\text{FNO}_2\text{Se}$ 366.0409; Found 366.0409. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



3-(3-fluoro-4-methoxyphenyl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3n), 64.0 mg, 84% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.5, 2.1$ Hz, 2H), 7.39 (dd, $J = 13.1, 2.7$ Hz, 1H), 7.35 - 7.26 (m, 3H), 7.07 (ddd, $J = 8.9, 2.8, 1.5$ Hz, 1H), 6.92 (t, $J = 9.1$ Hz, 1H), 4.71 (tdd, $J = 8.7, 6.3, 4.3$ Hz, 1H), 4.04 (t, $J = 8.7$ Hz, 1H), 3.86 (s, 3H), 3.70 (dd, $J = 9.0, 6.4$ Hz, 1H), 3.35 (dd, $J = 13.0, 4.3$ Hz, 1H), 3.05 (dd, $J = 12.9, 9.0$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.3, 152.2 (d, $J = 245.7$ Hz), 144.3 (d, $J = 10.7$ Hz), 133.7, 131.7 (d, $J = 9.2$ Hz), 129.6, 128.1, 127.9, 113.8 (d, $J = 3.6$ Hz), 113.7 (d, $J = 2.9$ Hz), 107.6 (d, $J = 23.5$ Hz), 71.7, 56.6, 50.4, 31.1. ^{19}F NMR (376 MHz, CDCl_3) δ -132.35. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{17}\text{FNO}_3\text{Se}$ 382.0358; Found 382.0359. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

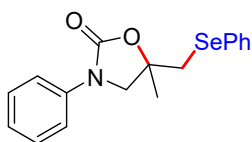


3-(naphthalen-2-yl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3o), 61.3 mg, 80% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.93 - 7.80 (m, 3H), 7.58 (dd, $J = 6.5, 2.9$ Hz, 2H), 7.52 (dtd, $J = 14.1, 6.9, 1.6$ Hz, 2H), 7.49 - 7.40 (m, 2H), 7.37 - 7.25 (m, 3H), 4.91 (tdd, $J = 8.4, 6.4, 4.3$ Hz, 1H), 4.14 (t, $J = 8.8$ Hz, 1H), 3.84 (dd, $J = 9.2, 6.5$ Hz, 1H), 3.44 (dd, $J = 13.0, 4.3$ Hz, 1H), 3.24 (dd, $J = 12.9, 8.6$ Hz, 1H). ^{13}C NMR (151 MHz, Chloroform- d) δ 156.6, 134.7, 133.8, 133.6, 129.9, 129.6, 128.7, 128.7, 128.3, 128.0, 127.1, 126.6, 125.7, 124.8, 122.4, 73.0, 54.0, 31.3. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{20}\text{H}_{28}\text{NO}_2\text{Se}$ 384.0503; Found 384.0505. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

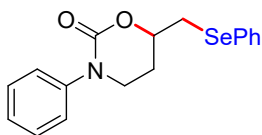


3-(benzo[d][1,3]dioxol-5-yl)-5-((phenylselanyl)methyl)oxazolidin-2-one (3p), 64.1 mg, 85% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.3, 2.2$ Hz, 2H), 7.36 - 7.26 (m, 3H), 7.24 (d, $J = 2.3$ Hz, 1H),

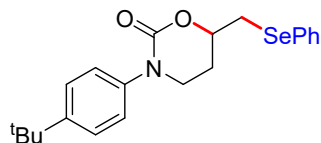
6.76 (d, $J = 8.4$ Hz, 1H), 6.69 (dd, $J = 8.5, 2.3$ Hz, 1H), 5.95 (s, 2H), 4.69 (tdd, $J = 8.8, 6.4, 4.3$ Hz, 1H), 4.04 (t, $J = 8.7$ Hz, 1H), 3.71 (dd, $J = 9.1, 6.4$ Hz, 1H), 3.35 (dd, $J = 12.9, 4.3$ Hz, 1H), 3.05 (dd, $J = 12.9, 9.1$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.6, 148.2, 144.4, 133.6, 132.6, 129.6, 128.1, 128.0, 111.5, 108.0, 101.5, 101.5, 71.7, 51.0, 31.1. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{16}\text{NO}_4\text{Se}$ 378.0245; Found 378.0247. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



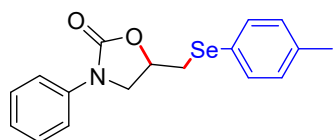
5-methyl-3-phenyl-5-((phenylselanyl)methyl)oxazolidin-2-one (3q), 61.1 mg, 88% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.53 (dd, $J = 7.5, 2.0$ Hz, 2H), 7.46 (d, $J = 7.6$ Hz, 2H), 7.38 - 7.31 (m, 2H), 7.28 - 7.22 (m, 3H), 7.12 (t, $J = 7.4$ Hz, 1H), 3.92 (d, $J = 9.0$ Hz, 1H), 3.69 (d, $J = 9.0$ Hz, 1H), 3.27 (q, $J = 13.0$ Hz, 2H), 1.60 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.0, 138.3, 133.5, 133.4, 133.4, 129.6, 129.5, 129.1, 127.8, 124.1, 118.3, 78.8, 55.2, 38.1, 26.2. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{18}\text{NO}_2\text{Se}$ 348.0503; Found 348.0506. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



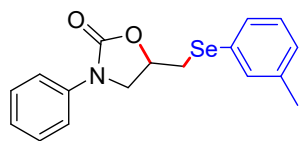
3-phenyl-6-((phenylselanyl)methyl)-1,3-oxazinan-2-one (3r), 59.7 mg, 86% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.55 (dd, $J = 7.3, 2.3$ Hz, 2H), 7.38 (t, $J = 7.8$ Hz, 2H), 7.34 - 7.24 (m, 6H), 4.52 (dddd, $J = 10.3, 9.0, 4.4, 2.6$ Hz, 1H), 3.73 (td, $J = 11.5, 4.7$ Hz, 1H), 3.63 (ddd, $J = 11.8, 5.6, 3.1$ Hz, 1H), 3.39 (dd, $J = 13.0, 4.4$ Hz, 1H), 3.05 (dd, $J = 13.0, 9.0$ Hz, 1H), 2.41 (ddt, $J = 13.8, 5.1, 2.9$ Hz, 1H), 2.02 (dtd, $J = 13.8, 10.8, 5.6$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 152.3, 142.6, 132.7, 129.4, 129.2, 129.1, 127.5, 126.9, 125.8, 76.9, 47.8, 31.0, 26.9. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{18}\text{NO}_2\text{Se}$ 348.0503; Found 348.0503. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



3-(4-(tert-butyl)phenyl)-6-((phenylselanyl)methyl)-1,3-oxazinan-2-one (3s), 70.9 mg, 88% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.55 (dd, $J = 5.0, 2.1$ Hz, 2H), 7.39 (d, $J = 8.7$ Hz, 2H), 7.30 (dd, $J = 5.2, 2.0$ Hz, 3H), 7.21 (d, $J = 8.7$ Hz, 2H), 4.51 (ttd, $J = 10.5, 9.2, 4.1, 2.5$ Hz, 1H), 3.71 (td, $J = 11.6, 4.7$ Hz, 1H), 3.62 (ddd, $J = 11.9, 5.6, 3.0$ Hz, 1H), 3.40 (dd, $J = 13.0, 4.4$ Hz, 1H), 3.04 (dd, $J = 13.0, 9.1$ Hz, 1H), 2.45 - 2.37 (m, 1H), 2.02 (dtd, $J = 14.0, 10.8, 5.5$ Hz, 1H), 1.30 (s, 9H). ^{13}C NMR (151 MHz, CDCl_3) δ 152.5, 149.8, 140.0, 132.8, 129.5, 129.1, 127.6, 126.3, 125.3, 76.9, 48.0, 34.6, 31.3, 31.0, 27.0. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{21}\text{H}_{26}\text{NO}_2\text{Se}$ 404.1130; Found 404.1130. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

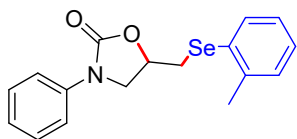


3-phenyl-5-((p-tolylselanyl)methyl)oxazolidin-2-one (3t), 59.7 mg, 86% yield. ^1H NMR (500 MHz, $\text{CDCl}_3/\text{CDCl}_3$) δ 7.47 (dd, $J = 15.4, 7.9$ Hz, 4H), 7.35 (t, $J = 7.8$ Hz, 2H), 7.11 (t, $J = 6.7$ Hz, 3H), 4.68 (ddt, $J = 13.4, 8.9, 3.6$ Hz, 1H), 4.08 (t, $J = 8.8$ Hz, 1H), 3.76 (dd, $J = 9.1, 6.4$ Hz, 1H), 3.31 (dd, $J = 12.8, 4.3$ Hz, 1H), 2.99 (dd, $J = 12.8, 9.2$ Hz, 1H), 2.33 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.5, 138.4, 138.2, 134.2, 130.4, 129.1, 124.1, 124.06, 118.3, 71.8, 50.3, 31.3, 21.2. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{18}\text{NO}_2\text{Se}$ 348.0503; Found 348.0503. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

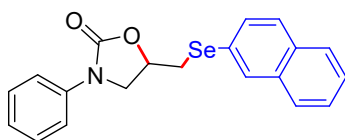


3-phenyl-5-((m-tolylselanyl)methyl)oxazolidin-2-one (3u), 59.0 mg, 85% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.49 (d, $J = 7.6$ Hz, 2H), 7.41 - 7.31 (m, 4H), 7.18 (t, $J = 7.6$ Hz, 1H), 7.16 - 7.09 (m, 2H), 4.71 (tdd, $J = 8.9, 6.3, 4.2$ Hz, 1H), 4.10 (t, $J = 8.8$ Hz, 1H), 3.78 (dd, $J = 9.2, 6.3$ Hz, 1H), 3.36 (dd, $J = 12.9, 4.2$ Hz, 1H), 3.03 (dd, $J = 12.8, 9.2$ Hz, 1H), 2.32 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.5, 139.5, 138.2, 134.3, 130.6, 129.4, 129.1, 129.0, 127.7, 124.2, 118.3, 71.8, 50.3, 31.0, 21.4. HRMS (ESI) m/z : $[\text{M}$

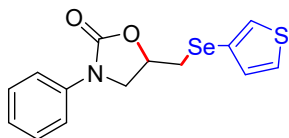
+ H]⁺ Calcd for C₁₇H₁₈NO₂Se 348.0503; Found 348.0503. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



3-phenyl-5-((o-tolylselanyl)methyl)oxazolidin-2-one (3v), 55.5 mg, 80% yield. ¹H NMR (500 MHz, CDCl₃) δ 7.51 (dd, *J* = 12.5, 7.9 Hz, 3H), 7.40 - 7.32 (m, 2H), 7.22 (d, *J* = 6.6 Hz, 2H), 7.13 (t, *J* = 7.3 Hz, 2H), 4.71 (tdd, *J* = 8.8, 6.3, 4.3 Hz, 1H), 4.10 (t, *J* = 8.8 Hz, 1H), 3.80 (dd, *J* = 9.1, 6.3 Hz, 1H), 3.34 (dd, *J* = 12.8, 4.3 Hz, 1H), 3.03 (dd, *J* = 12.8, 9.3 Hz, 1H), 2.45 (s, 3H). ¹³C NMR (151 MHz, CDCl₃) δ 154.5, 140.4, 138.2, 133.3, 130.6, 129.2, 128.9, 128.2, 127.0, 124.2, 118.3, 71.7, 50.4, 30.0, 22.7. HRMS (ESI) *m/z*: [M + H]⁺ Calcd for C₁₇H₁₈NO₂Se 348.0503; Found 348.0503. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

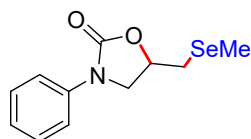


5-((naphthalen-2-ylselanyl)methyl)-3-phenyloxazolidin-2-one (3w), 62.8 mg, 82% yield. ¹H NMR (500 MHz, CDCl₃) δ 8.40 (d, *J* = 8.4 Hz, 1H), 7.88 (dd, *J* = 13.8, 7.6 Hz, 3H), 7.56 (dt, *J* = 24.3, 6.9 Hz, 2H), 7.43 (d, *J* = 7.5 Hz, 2H), 7.41 - 7.30 (m, 3H), 7.11 (t, *J* = 7.4 Hz, 1H), 4.63 (tdd, *J* = 8.7, 6.4, 4.4 Hz, 1H), 4.03 (t, *J* = 8.8 Hz, 1H), 3.73 (dd, *J* = 9.1, 6.3 Hz, 1H), 3.37 (dd, *J* = 12.7, 4.5 Hz, 1H), 3.08 (dd, *J* = 12.7, 9.0 Hz, 1H). ¹³C NMR (151 MHz, CDCl₃) δ 154.4, 138.2, 134.5, 134.4, 134.3, 129.8, 129.2, 129.0, 127.6, 127.4, 127.2, 126.7, 126.0, 124.2, 118.3, 71.9, 50.3, 31.3. HRMS (ESI) *m/z*: [M + H]⁺ Calcd for C₂₀H₁₈NO₂Se 384.0503; Found 384.0504. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

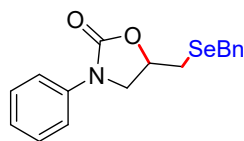


3-phenyl-5-((thiophen-3-ylselanyl)methyl)oxazolidin-2-one (3x), 52.9 mg, 78% yield. ¹H NMR (500 MHz, CDCl₃) δ 7.50 (d, *J* = 7.6 Hz, 2H), 7.43 (dd, *J* = 3.0, 1.2 Hz, 1H), 7.41 - 7.30 (m, 3H), 7.21 - 7.09

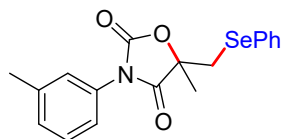
(m, 2H), 4.72 (tdd, $J = 8.7, 6.4, 4.4$ Hz, 1H), 4.12 (t, $J = 8.8$ Hz, 1H), 3.77 (dd, $J = 9.1, 6.3$ Hz, 1H), 3.27 (dd, $J = 12.9, 4.4$ Hz, 1H), 2.98 (dd, $J = 12.8, 8.9$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.5, 138.2, 132.6, 129.4, 129.2, 127.4, 124.3, 120.7, 118.3, 71.9, 50.3, 31.8. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{14}\text{H}_{24}\text{NO}_2\text{SSe}$ 339.9910; Found 339.9912. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



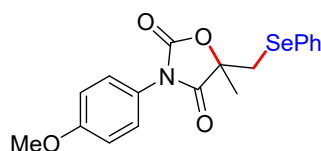
5-((methylselanyl)methyl)-3-phenyloxazolidin-2-one (3y), 50.4 mg, 93% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.54 (d, $J = 7.4$ Hz, 2H), 7.42 - 7.35 (m, 2H), 7.14 (t, $J = 7.4$ Hz, 1H), 4.83 (tdd, $J = 8.2, 6.5, 4.5$ Hz, 1H), 4.16 (t, $J = 8.7$ Hz, 1H), 3.81 (dd, $J = 9.0, 6.4$ Hz, 1H), 2.95 (dd, $J = 13.1, 4.5$ Hz, 1H), 2.82 (dd, $J = 13.1, 7.9$ Hz, 1H), 2.12 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.5, 138.2, 129.1, 124.1, 118.3, 72.2, 50.4, 28.6, 5.4. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{11}\text{H}_{14}\text{NO}_2\text{Se}$ 272.0190; Found 272.0189. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



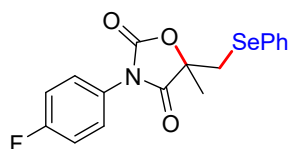
5-((benzylselanyl)methyl)-3-phenyloxazolidin-2-one (3z), 40.3 mg, 58% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.48 (d, $J = 8.1$ Hz, 2H), 7.36 (t, $J = 8.0$ Hz, 2H), 7.30 (d, $J = 4.4$ Hz, 4H), 7.22 (dq, $J = 8.6, 4.3, 3.8$ Hz, 1H), 7.13 (t, $J = 7.4$ Hz, 1H), 4.63 (tdd, $J = 8.2, 6.6, 4.5$ Hz, 1H), 4.00 (t, $J = 8.7$ Hz, 1H), 3.89 (s, 2H), 3.62 (dd, $J = 9.0, 6.5$ Hz, 1H), 2.86 (dd, $J = 13.2, 4.5$ Hz, 1H), 2.76 (dd, $J = 13.2, 7.8$ Hz, 1H). ^{13}C NMR (151 MHz, CDCl_3) δ 154.5, 138.6, 138.2, 129.1, 129.1, 128.8, 127.3, 124.2, 118.4, 72.1, 50.4, 28.1, 26.9. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{17}\text{H}_{18}\text{NO}_2\text{Se}$ 348.0503; Found 348.0503. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



5-methyl-5-((phenylselanyl)methyl)-3-(m-tolyl)oxazolidine-2,4-dione (5a), 56.3 mg, 75% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.57 (dd, $J = 7.5, 2.1$ Hz, 2H), 7.36 (t, $J = 7.7$ Hz, 1H), 7.32 - 7.25 (m, 3H), 7.25 - 7.14 (m, 3H), 3.43 (q, $J = 13.8$ Hz, 2H), 2.39 (s, 3H), 1.75 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 173.7, 153.4, 139.6, 133.7, 133.7, 130.9, 130.0, 129.5, 129.3, 129.0, 128.1, 126.4, 123.0, 85.3, 34.5, 23.0, 21.5. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{18}\text{H}_{18}\text{NO}_3\text{Se}$ 376.0452; Found 376.0455. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

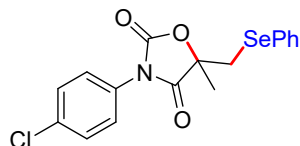


3-(4-methoxyphenyl)-5-methyl-5-((phenylselanyl)methyl)oxazolidine-2,4-dione (5b), 61.0 mg, 78% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.56 (dd, $J = 7.5, 2.1$ Hz, 2H), 7.32 (d, $J = 9.0$ Hz, 2H), 7.29 - 7.20 (m, 3H), 6.98 (d, $J = 8.9$ Hz, 2H), 3.82 (s, 3H), 3.47 - 3.36 (m, 2H), 1.74 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 173.8, 159.9, 153.6, 133.7, 129.4, 129.0, 128.1, 127.3, 123.6, 114.7, 85.4, 55.6, 34.6, 22.9. HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ Calcd for $\text{C}_{18}\text{H}_{18}\text{NO}_4\text{Se}$ 392.0401; Found 392.0400. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

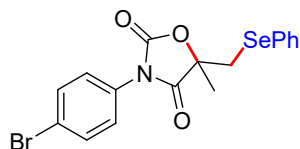


3-(4-fluorophenyl)-5-methyl-5-((phenylselanyl)methyl)oxazolidine-2,4-dione (5c), 53.1 mg, 70% yield. ^1H NMR (500 MHz, CDCl_3) δ 7.55 (d, $J = 7.9$ Hz, 2H), 7.41 (dd, $J = 9.0, 4.7$ Hz, 2H), 7.32 - 7.23 (m, 3H), 7.18 (t, $J = 8.6$ Hz, 2H), 3.49 - 3.39 (m, 2H), 1.77 (s, 3H). ^{13}C NMR (151 MHz, CDCl_3) δ 173.5, 162.5 (d, $J = 249.8$ Hz), 153.2, 133.7, 129.5, 128.9, 128.2, 127.8 (d, $J = 8.9$ Hz), 127.0 (d, $J = 3.1$ Hz), 116.5 (d, $J = 23.1$ Hz), 85.6, 34.6, 23.0. ^{19}F NMR (376 MHz, CDCl_3) δ -111.50. HRMS (ESI) m/z : $[\text{M} +$

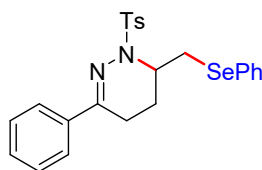
$[H]^+$ Calcd for $C_{20}H_{24}NO_2Se$ 390.0972; Found 390.0973. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



3-(4-chlorophenyl)-5-methyl-5-((phenylselanyl)methyl)oxazolidine-2,4-dione (5d), 57.7 mg, 73% yield. 1H NMR (500 MHz, $CDCl_3$) δ 7.54 (dd, J = 7.7, 1.8 Hz, 2H), 7.49 - 7.42 (m, 2H), 7.42 - 7.34 (m, 2H), 7.32 - 7.20 (m, 3H), 3.48 - 3.37 (m, 2H), 1.76 (s, 3H). ^{13}C NMR (151 MHz, $CDCl_3$) δ 173.3, 152.9, 134.8, 133.6, 129.6, 129.5, 129.5, 128.8, 128.2, 126.9, 85.5, 34.5, 22.9. HRMS (ESI) m/z : $[M + H]^+$ Calcd for $C_{17}H_{15}NO_3Se$ 380.0201; Found 380.0203. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).

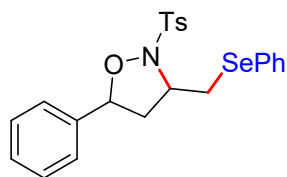


3-(4-bromophenyl)-5-methyl-5-((phenylselanyl)methyl)oxazolidine-2,4-dione (5e), 65.8 mg, 75% yield. 1H NMR (500 MHz, $CDCl_3$) δ 7.61 (d, J = 8.7 Hz, 2H), 7.54 (dd, J = 7.8, 1.8 Hz, 2H), 7.33 (d, J = 8.7 Hz, 2H), 7.29 - 7.23 (m, 3H), 3.48 - 3.38 (m, 2H), 1.76 (s, 3H). ^{13}C NMR (151 MHz, $CDCl_3$) δ 173.3, 152.9, 133.7, 132.7, 130.1, 129.5, 128.8, 128.2, 127.2, 122.9, 85.6, 34.5, 23.0. HRMS (ESI) m/z : $[M + H]^+$ Calcd for $C_{17}H_{15}BrNO_3Se$ 439.9401; Found 439.9402. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 10 ~ 20%, v/v).



3-phenyl-6-((phenylselanyl)methyl)-1-tosyl-1,4,5,6-tetrahydropyridazine (7a), 92.0 mg, 95% yield. 1H NMR (400 MHz, $CDCl_3$) δ 7.80 (d, J = 8.1 Hz, 2H), 7.67 (dd, J = 7.5, 2.3 Hz, 2H), 7.58 (dd, J = 7.7, 1.8 Hz, 2H), 7.44 - 7.28 (m, 6H), 7.26 (d, J = 8.2 Hz, 2H), 4.50 (d, J = 11.4 Hz, 1H), 3.35 (dd, J = 12.9, 2.6 Hz, 1H), 2.79 (t, J = 12.3 Hz, 1H), 2.60 - 2.46 (m, 2H), 2.38 (s, 3H), 2.28 - 2.10 (m, 1H), 1.57 (dq, J = 13.0, 6.6, 5.1 Hz, 1H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 148.0, 143.9, 136.7, 135.9, 132.6, 129.6, 129.5, S15

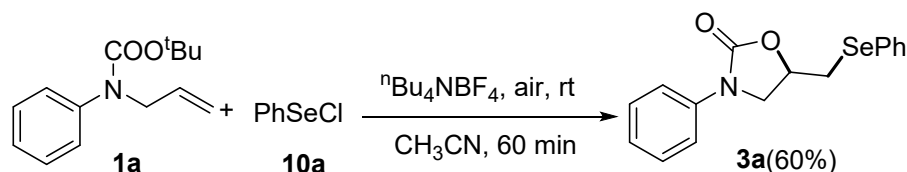
129.4, 128.9, 128.5, 128.0, 127.5, 125.3, 51.4, 28.1, 21.7, 19.7, 17.8. HRMS (ESI) m/z : $[M + H]^+$ Calcd for $C_{24}H_{25}N_2O_2Se$ 485.0802; Found 485.0804. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 2 ~ 5%, v/v).



5-phenyl-3-((phenylselanyl)methyl)-2-tosyloxazolidine (9a), 88.0 mg, 93% yield. **dr** = **1.2:1**. 1H NMR (400 MHz, $CDCl_3$) δ 7.75 (d, J = 8.0 Hz, 2H), 7.67 (d, J = 8.0 Hz, 1.66H), 7.58 (qd, J = 4.3, 2.9, 1.8 Hz, 3.71H), 7.36 - 7.20 (m, 18.4H), 5.16 (dd, J = 11.0, 5.7 Hz, 0.83H), 4.94 (dd, J = 10.5, 5.6 Hz, 1H), 4.43 (dt, J = 12.6, 6.3 Hz, 1H), 4.31 (td, J = 9.2, 5.1 Hz, 0.83H), 3.57 (dd, J = 12.6, 4.5 Hz, 1H), 3.46 (dd, J = 12.8, 5.0 Hz, 0.83H), 3.11 (dd, J = 12.6, 9.4 Hz, 1H), 3.07 - 3.00 (m, 0.83H), 2.83 (dt, J = 12.7, 6.5 Hz, 1H), 2.54 (dd, J = 12.9, 5.8 Hz, 0.84H), 2.42 (s, 5.5H), 2.22 - 2.14 (m, 1H), 2.14 - 2.07 (m, 0.83H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 145.1, 145.1, 136.6, 136.2, 133.0, 132.7, 132.4, 131.2, 129.7, 129.6, 129.5, 129.4, 129.3, 129.3, 128.9, 128.8, 128.7, 128.6, 128.5, 127.5, 127.4, 127.1, 126.9, 83.3, 82.4, 61.9, 61.1, 43.3, 40.8, 32.5, 31.2, 21.7, 21.7. HRMS (ESI) m/z : $[M + H]^+$ Calcd for $C_{23}H_{24}NO_3SSe$ 474.0642; Found 474.0641. The isolated yield was obtained by chromatography column on silica gel (gradient eluent of EtOAc in Petroleum ether: 2 ~ 10%, v/v).

4. Mechanistic studies

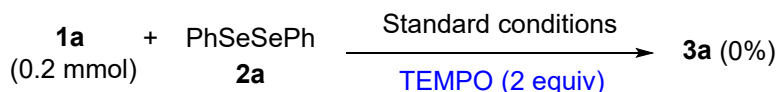
a) Synthesis of Selenium-Containing Oxazolidinones with Phenylselenium Chloride as the Selenium Source



To the cell was added **1a** (0.2 mmol, 46.6 mg), **10a** (0.24 mmol, 45.96 mg), nBu_4NBF_4 (0.1 mmol, 32.93 mg), CH_3CN (4.0 mL). The mixture was stirred at room temperature for 60 min. Then, the solvent was removed under reduced pressure. The residue was poured into a saturated aqueous solution of NaCl and

the product was then extracted with CH₂Cl₂ (3×20 mL), dried over Na₂SO₄, and concentrated in vacuo. The residue was purified by column chromatography on silica gel using a mixture of petroleum ether/EtOA eluent to afford the desired pure product **3a** (40 mg, 60%).

b) Radical trapping experiments

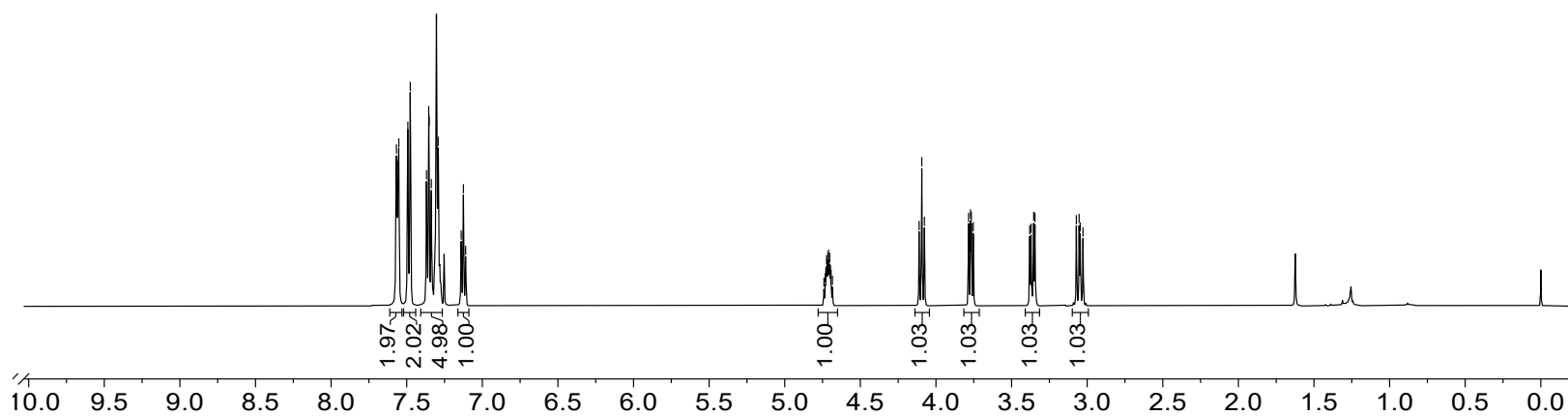
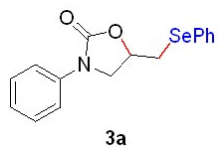
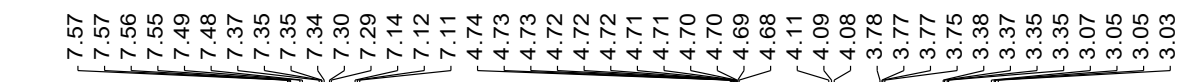


An undivided cell was equipped with Pt anode (1.0 cm × 1.0 cm × 0.1 cm) and Pt cathode (1.0 cm × 1.0 cm × 0.1 cm). To the cell was added **1a** (0.2 mmol, 46.6 mg), **2a** (0.24 mmol, 74.88 mg), *n*Bu₄NBF (0.1 mol, 32.93 mg), TEMPO (0.4 mmol, 62.5 mg), CH₃CN (4.0 mL). The mixture was electrolyzed using constant current conditions (*I* = 5 mA) at room temperature under magnetic stirring for 60 min. After the reaction was completed, analysis by GC-MS did not detect the target compound **3a**.

6. Copies of NMR spectra

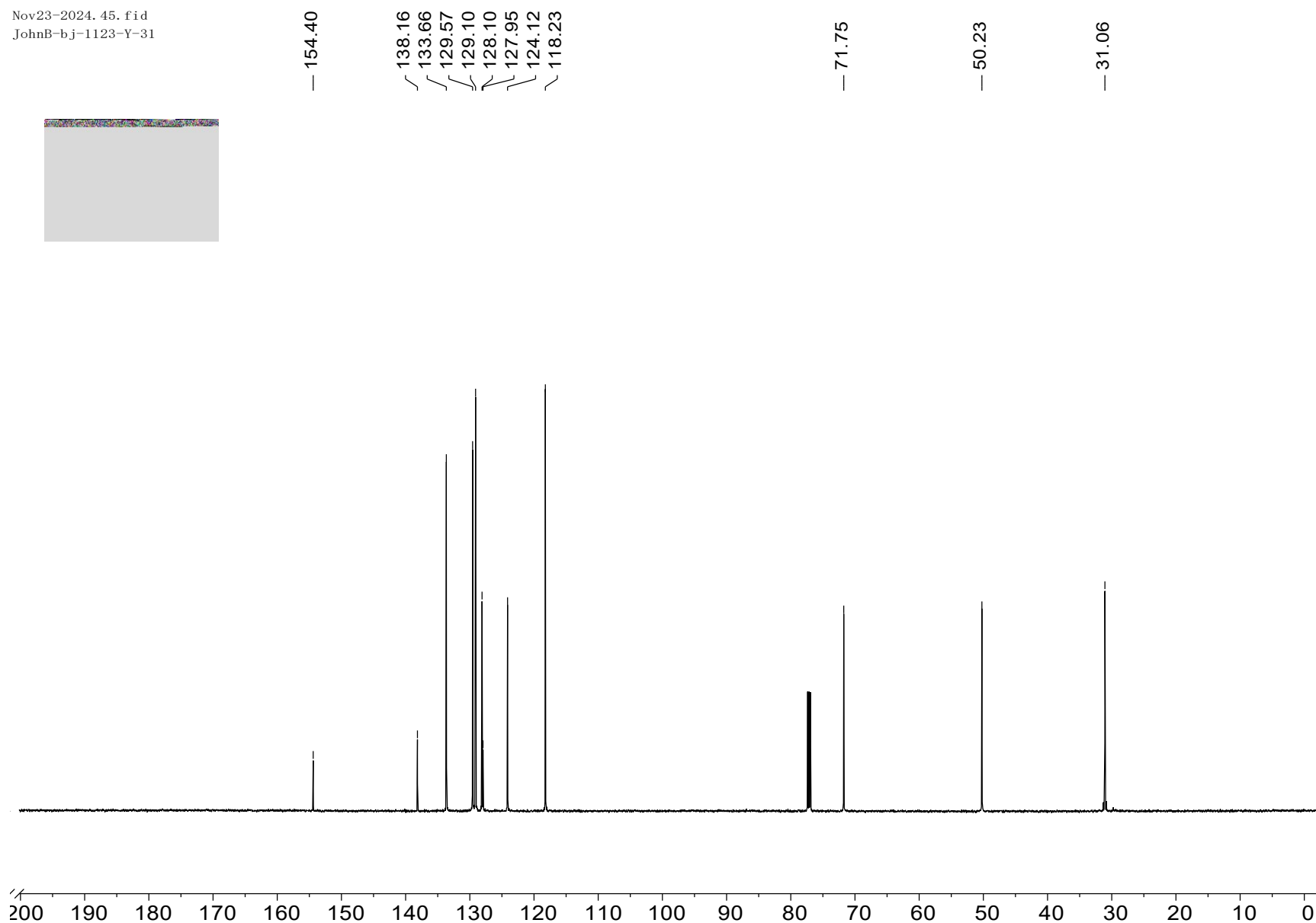
^1H NMR (500 MHz, CDCl_3) spectrum of **3a**.

31.1.fid



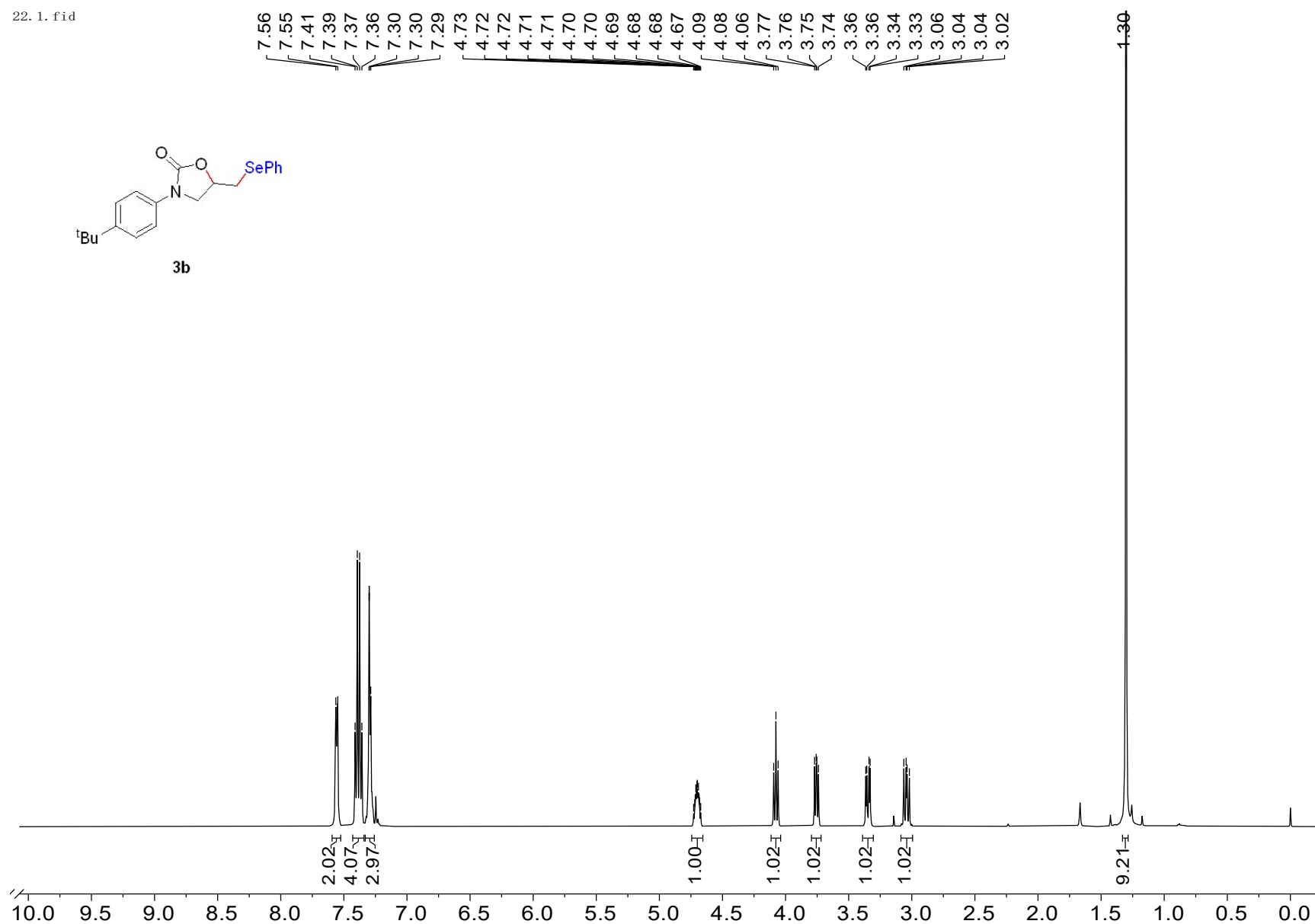
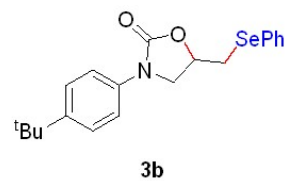
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3a**.

Nov23-2024. 45. fid
JohnB-bj-1123-Y-31



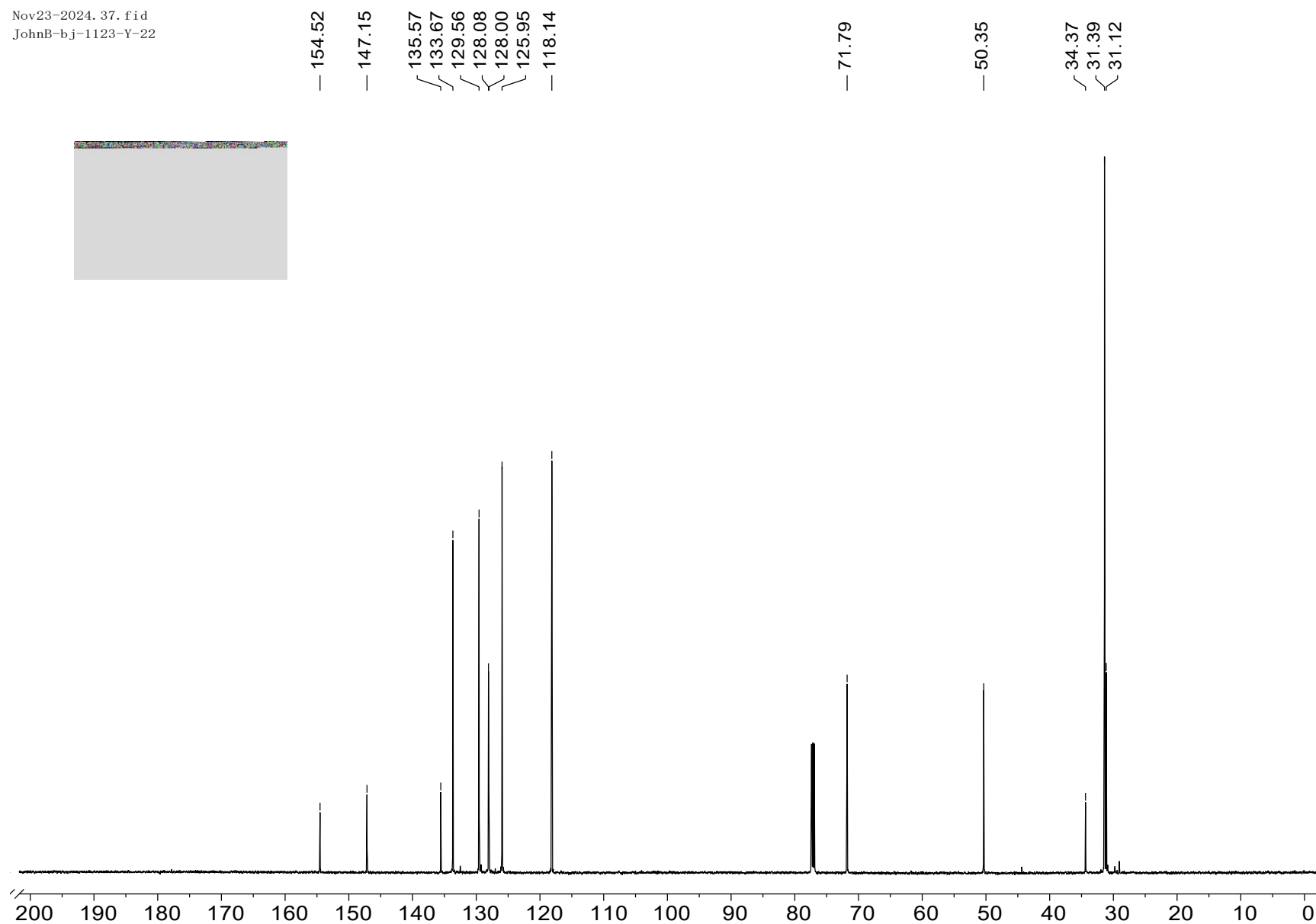
^1H NMR (500 MHz, CDCl_3) spectrum of **3b**.

22.1.fid



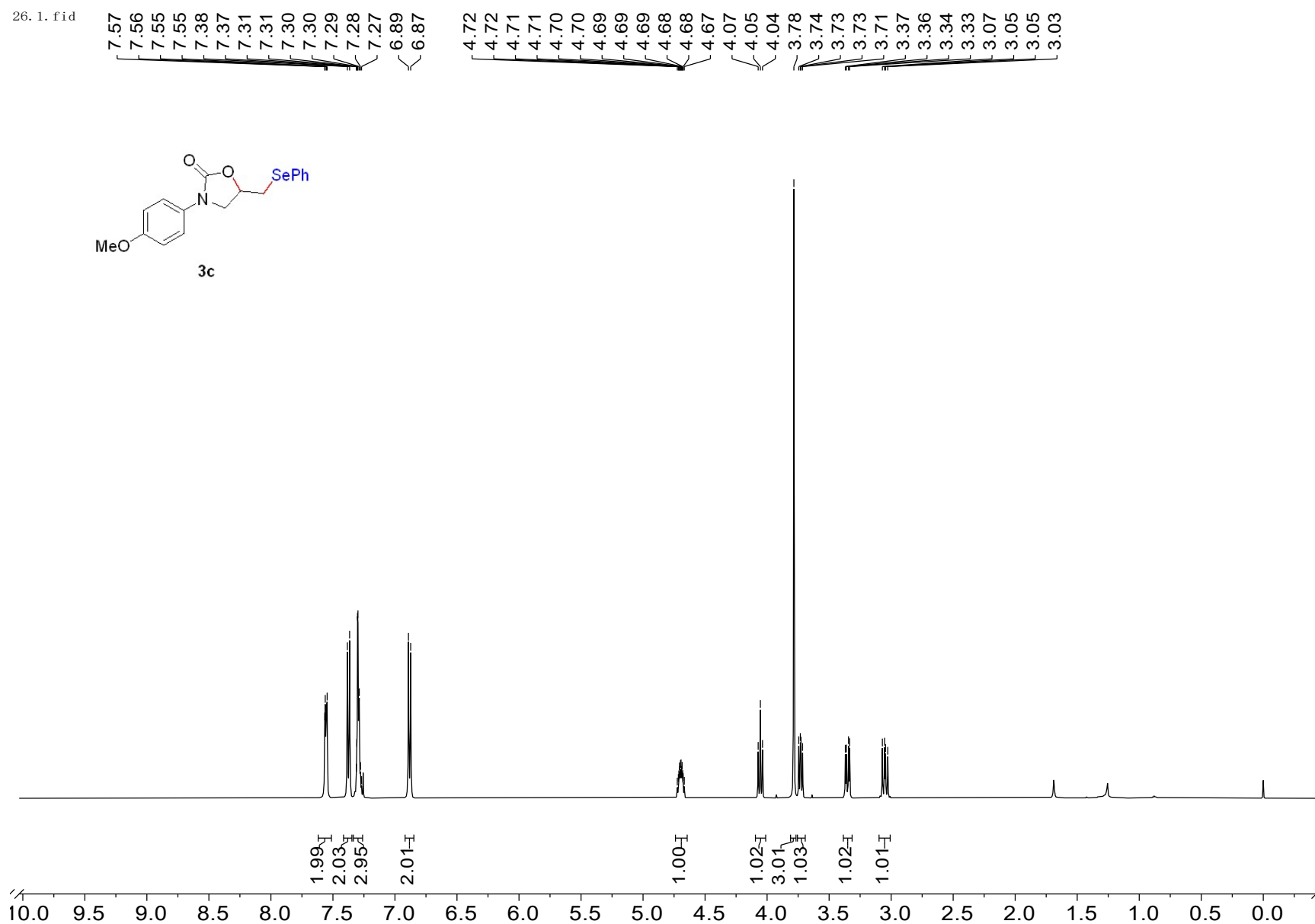
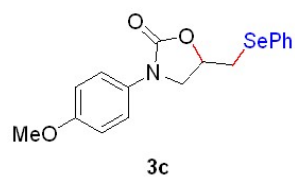
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3b**.

Nov23-2024. 37. fid
JohnB-bj-1123-Y-22



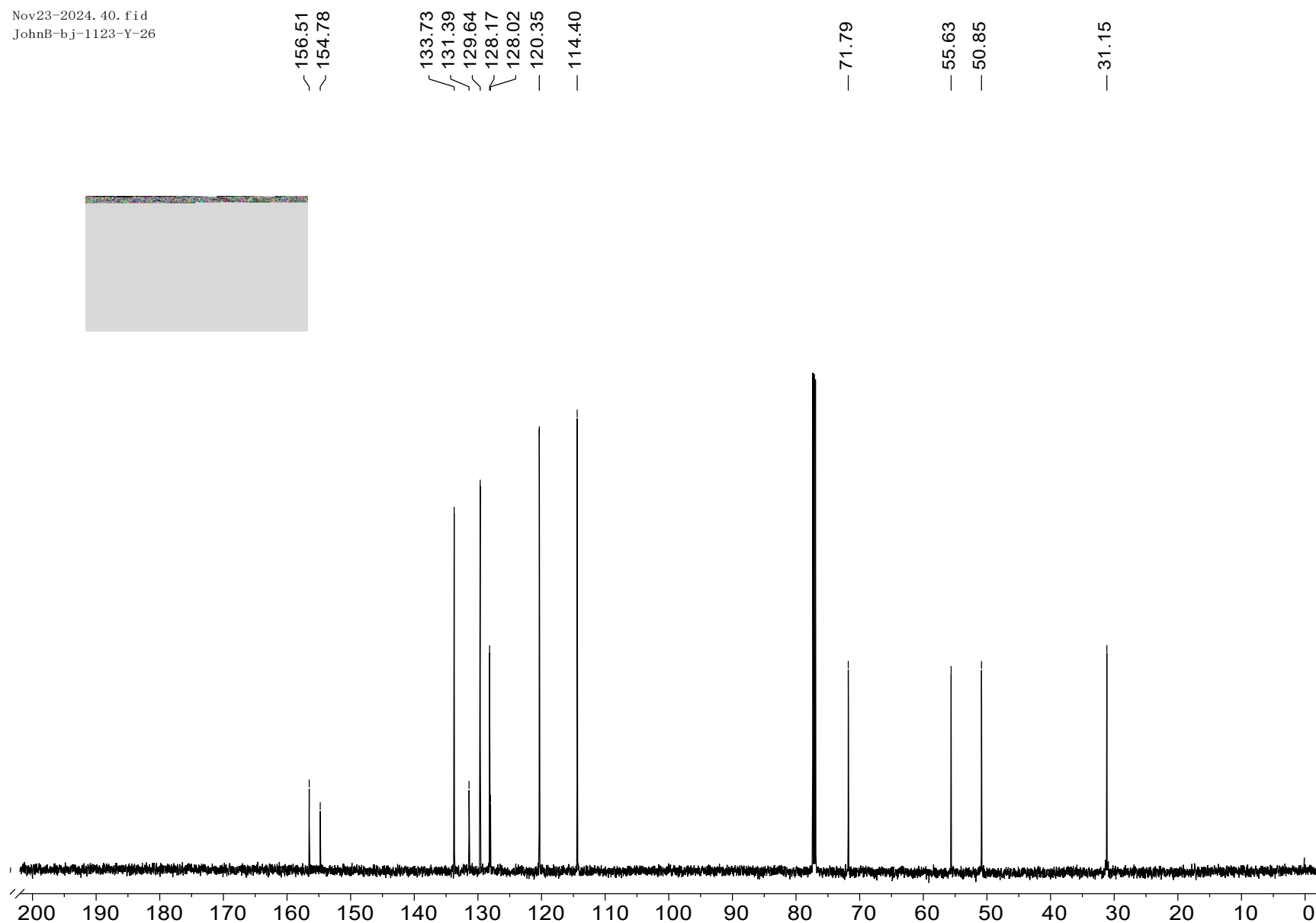
^1H NMR (500 MHz, CDCl_3) spectrum of **3c**.

26.1.1.fid

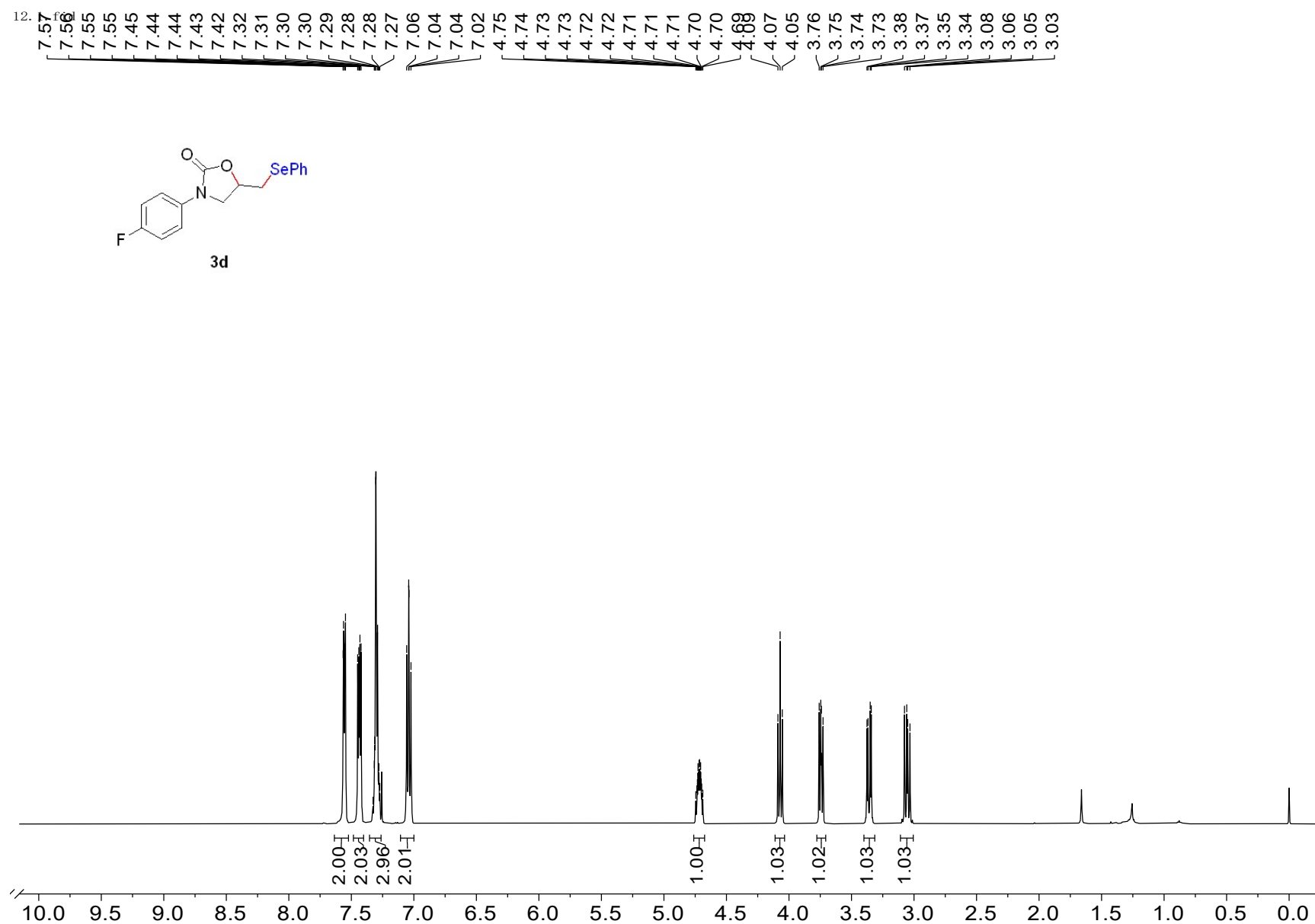


^{13}C NMR (151 MHz, CDCl_3) spectrum of **3c**.

Nov23-2024. 40. fid
JohnB-bj-1123-Y-26

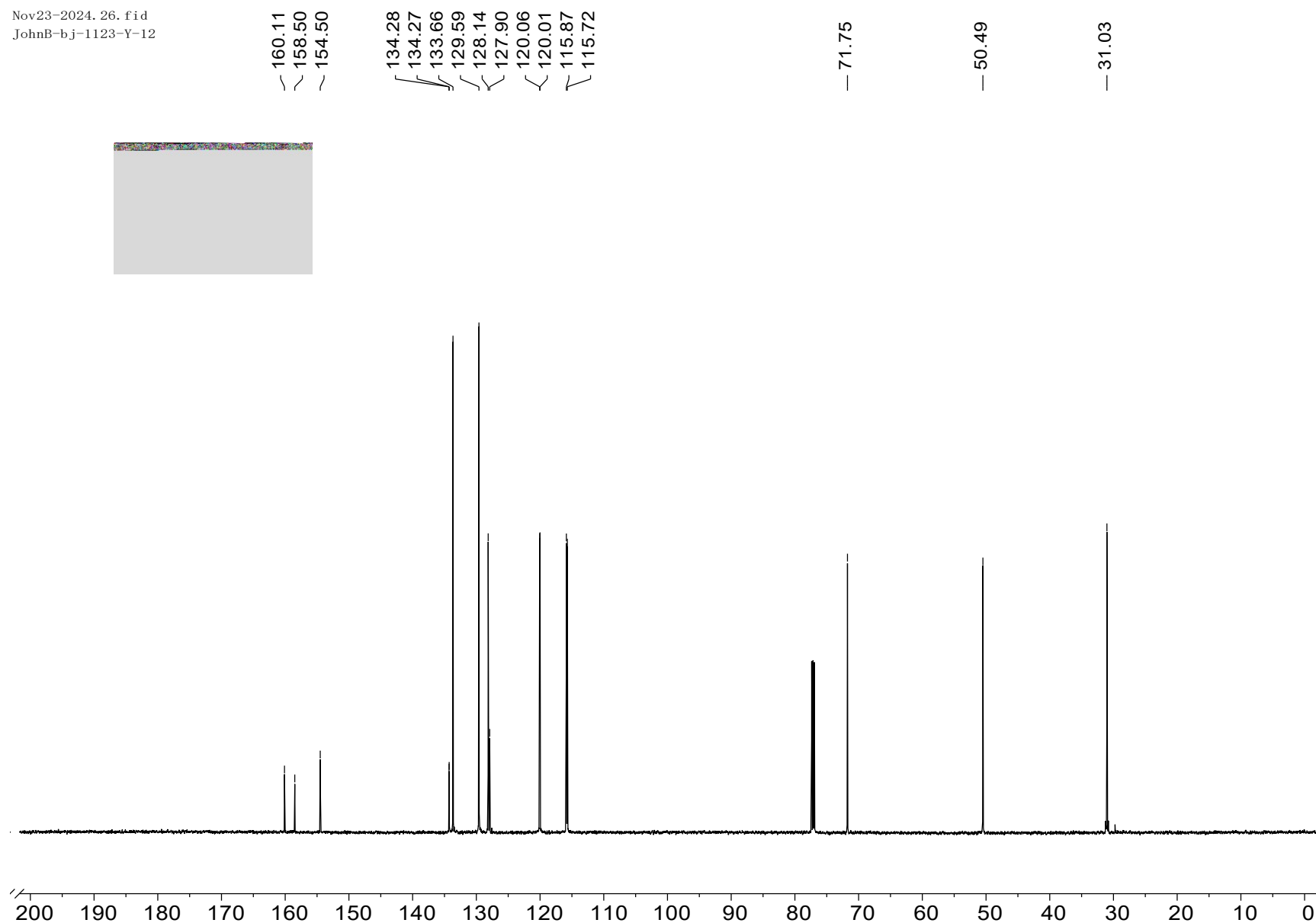


^1H NMR (500 MHz, CDCl_3) spectrum of **3d**.



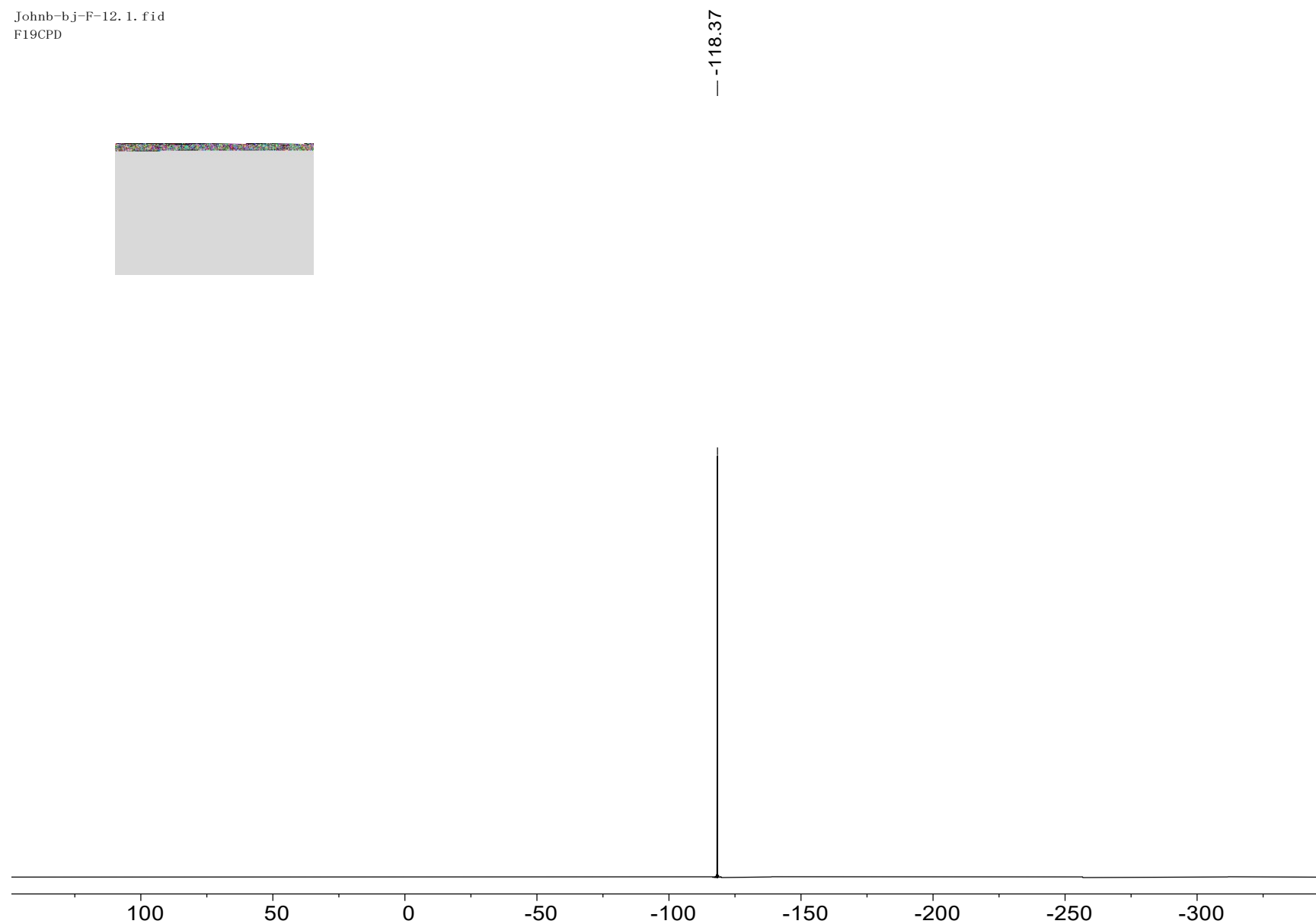
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3d**.

Nov23-2024, 26. fid
JohnB-bj-1123-Y-12



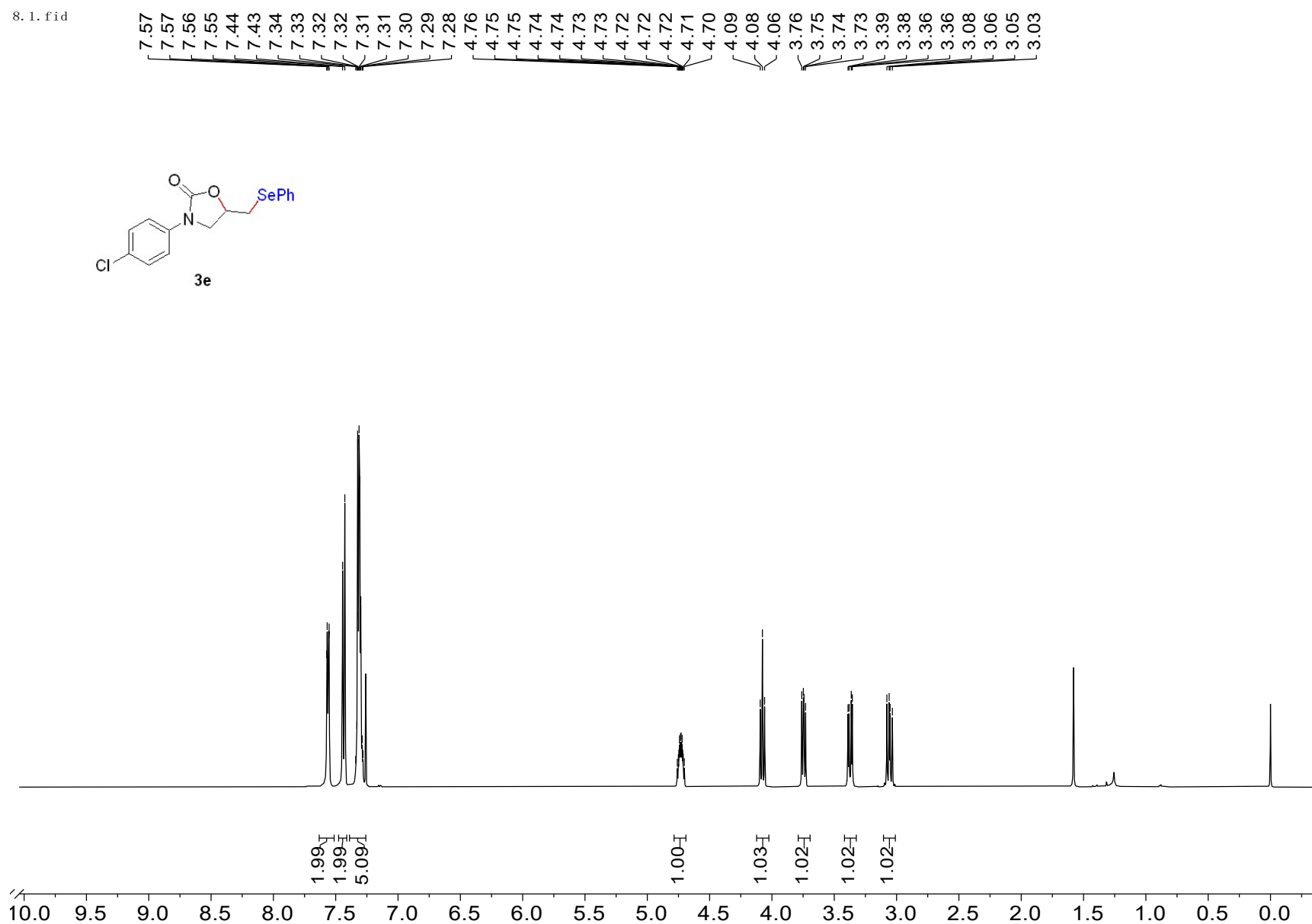
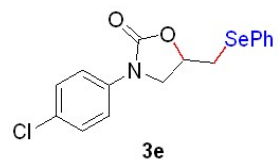
^{19}F NMR (376 MHz, CDCl_3) spectrum of **3d**.

Johnb-bj-F-12. 1. fid
F19CPD



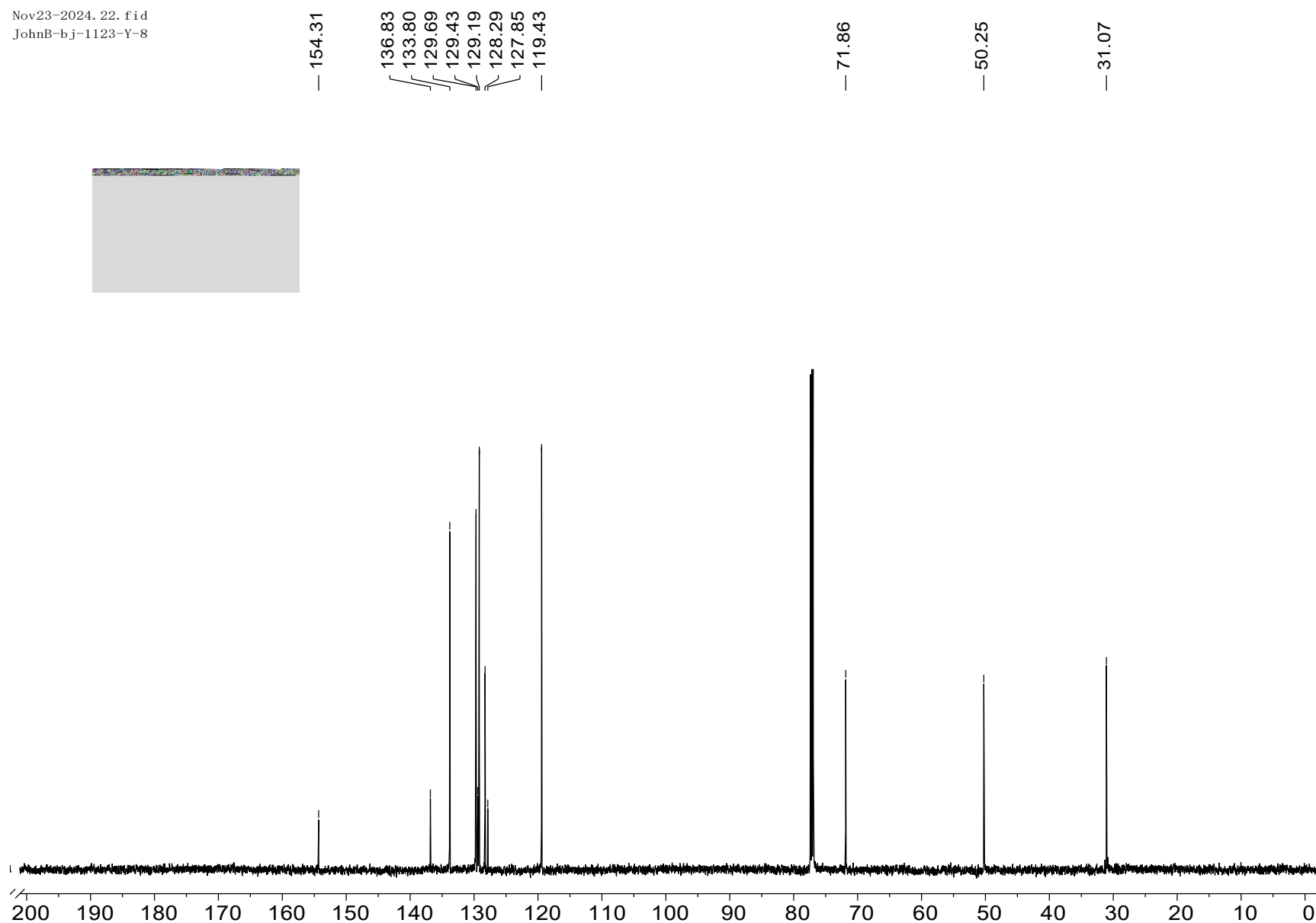
^1H NMR (500 MHz, CDCl_3) spectrum of **3e**.

8.1.fid



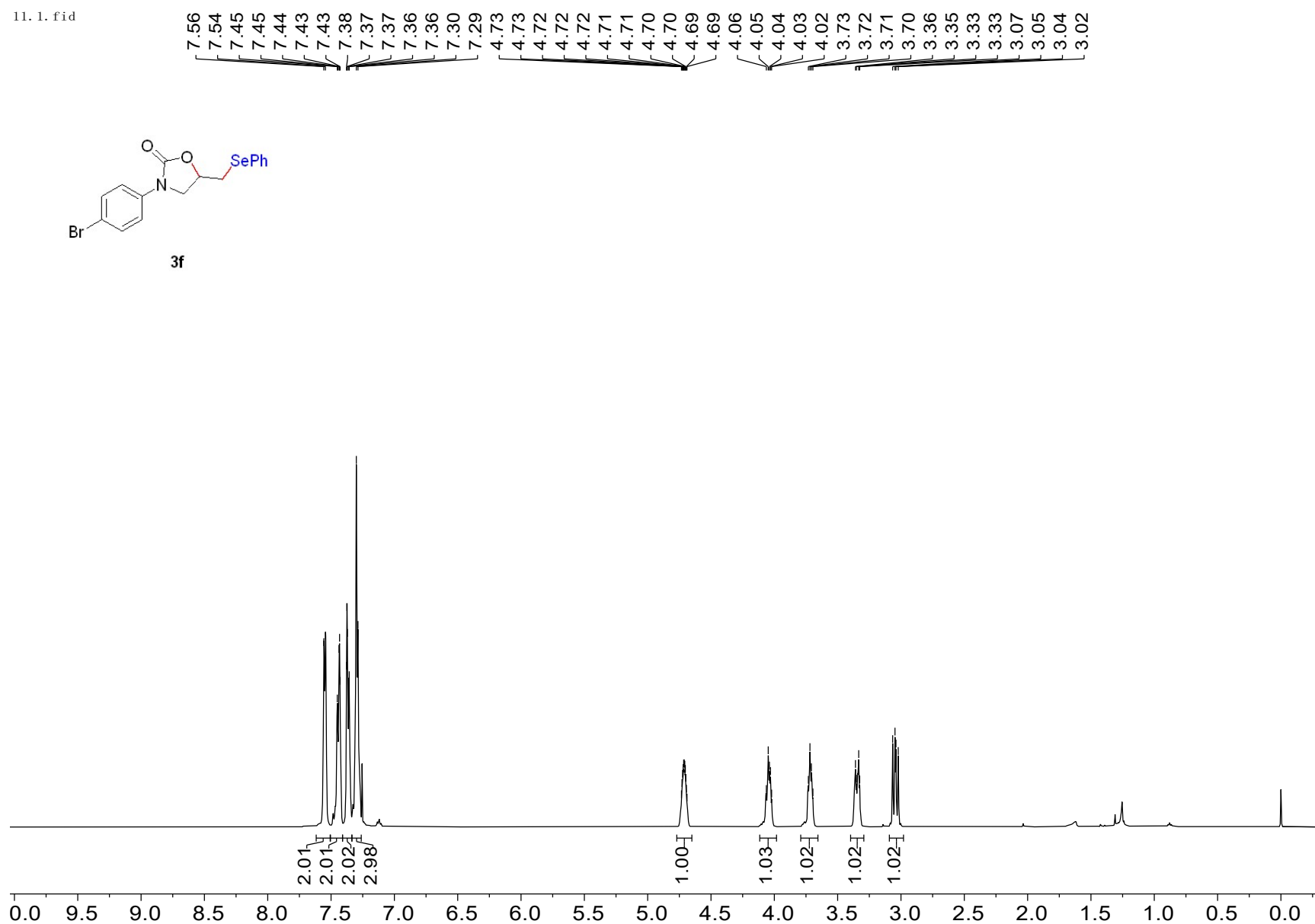
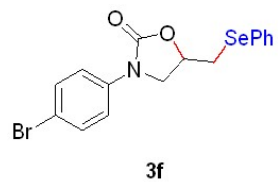
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3e**.

Nov23-2024, 22. fid
JohnB-bj-1123-Y-8



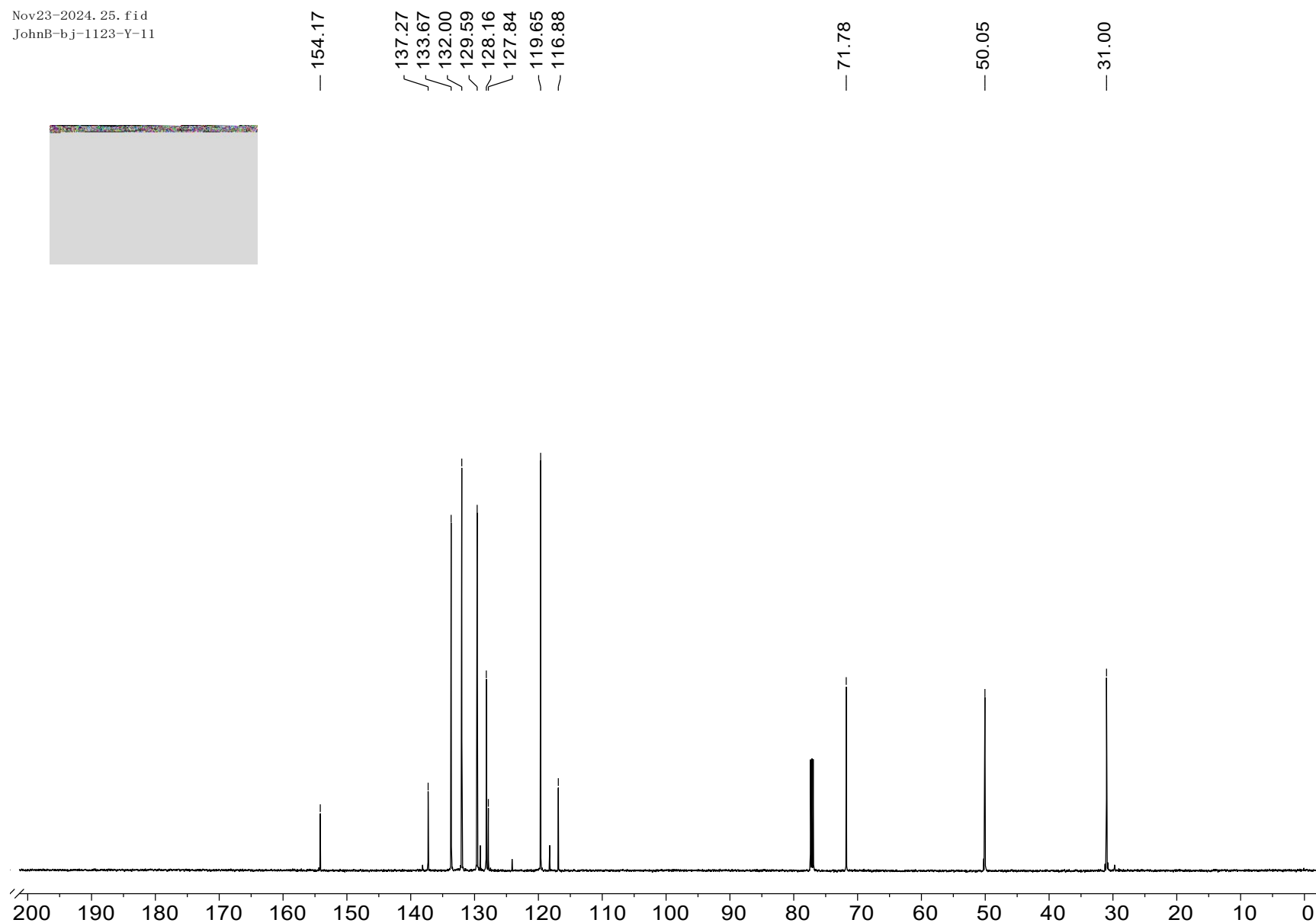
^1H NMR (500 MHz, CDCl_3) spectrum of **3f**.

11.1.fid



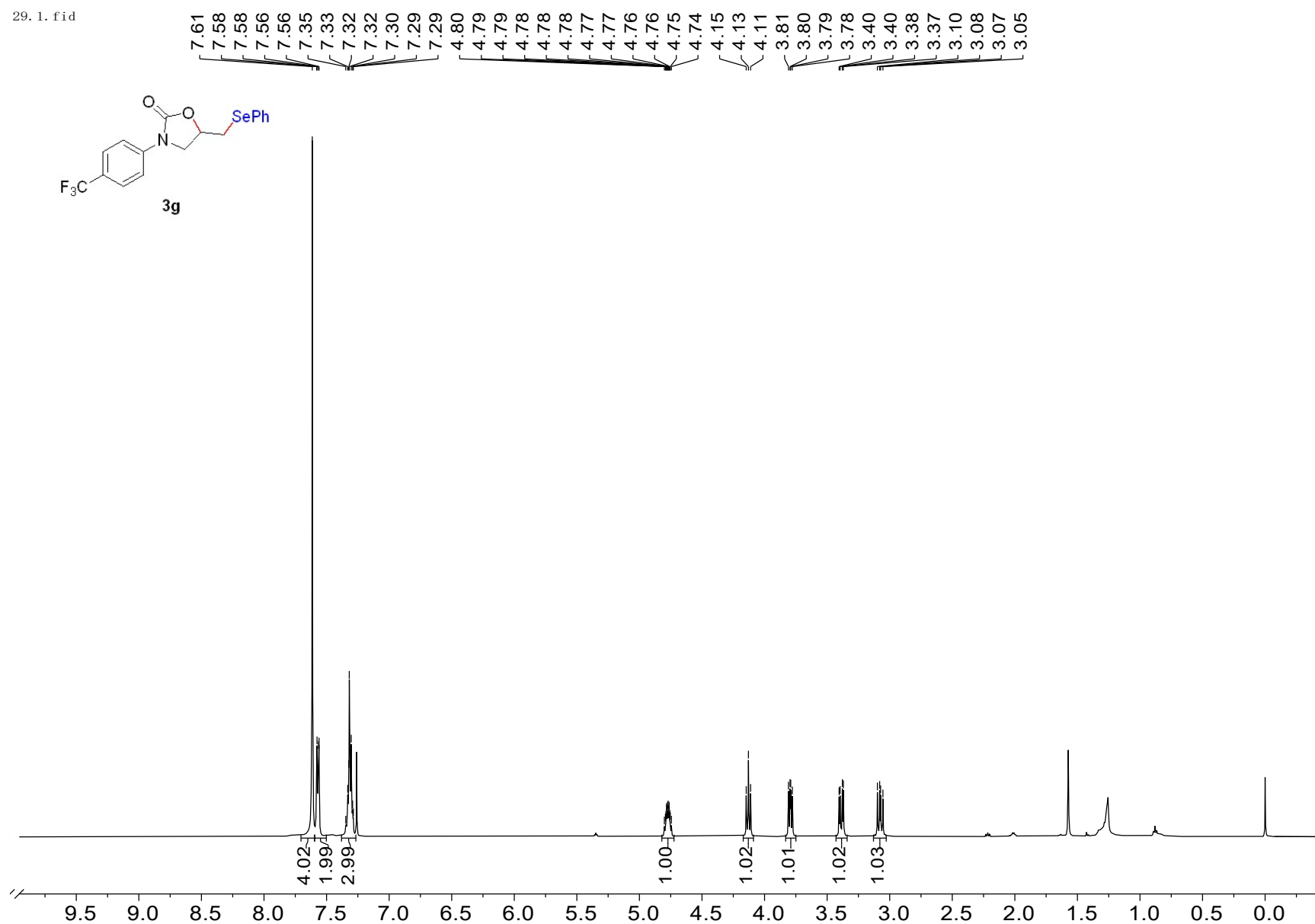
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3f**.

Nov23-2024. 25. fid
JohnB-bj-1123-Y-11



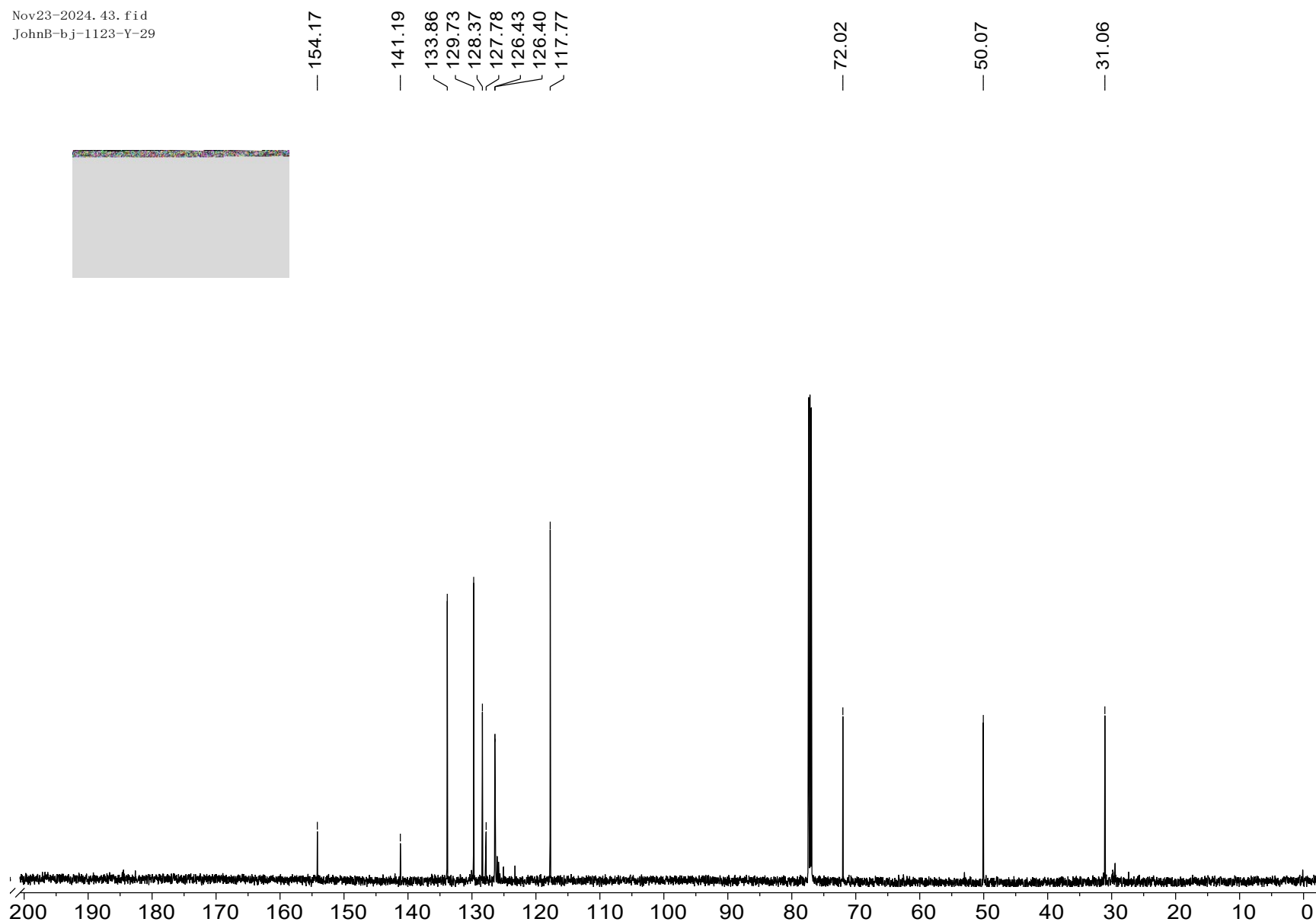
^1H NMR (500 MHz, CDCl_3) spectrum of **3g**.

29.1.fid



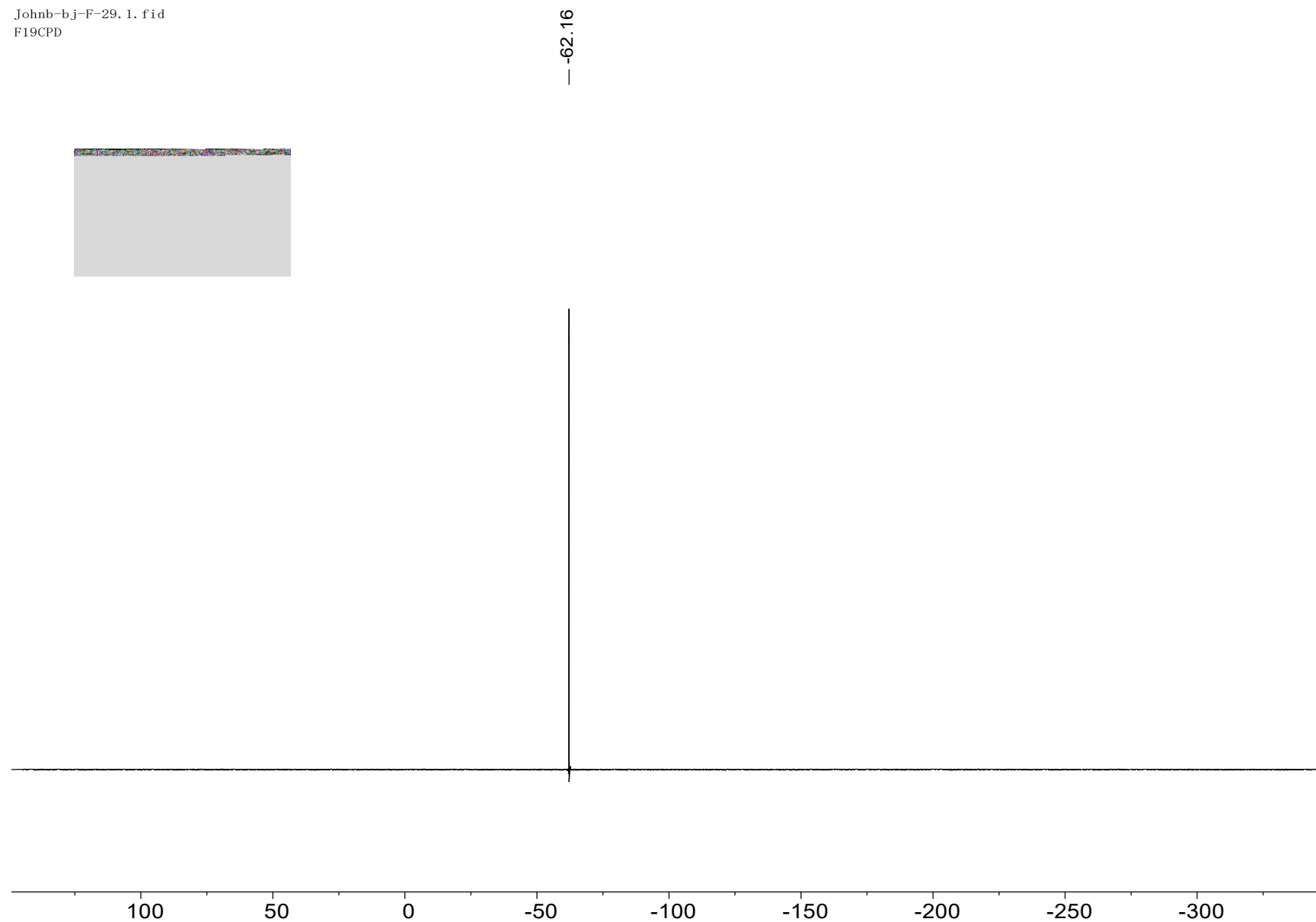
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3g**.

Nov23-2024. 43. fid
JohnB-bj-1123-Y-29

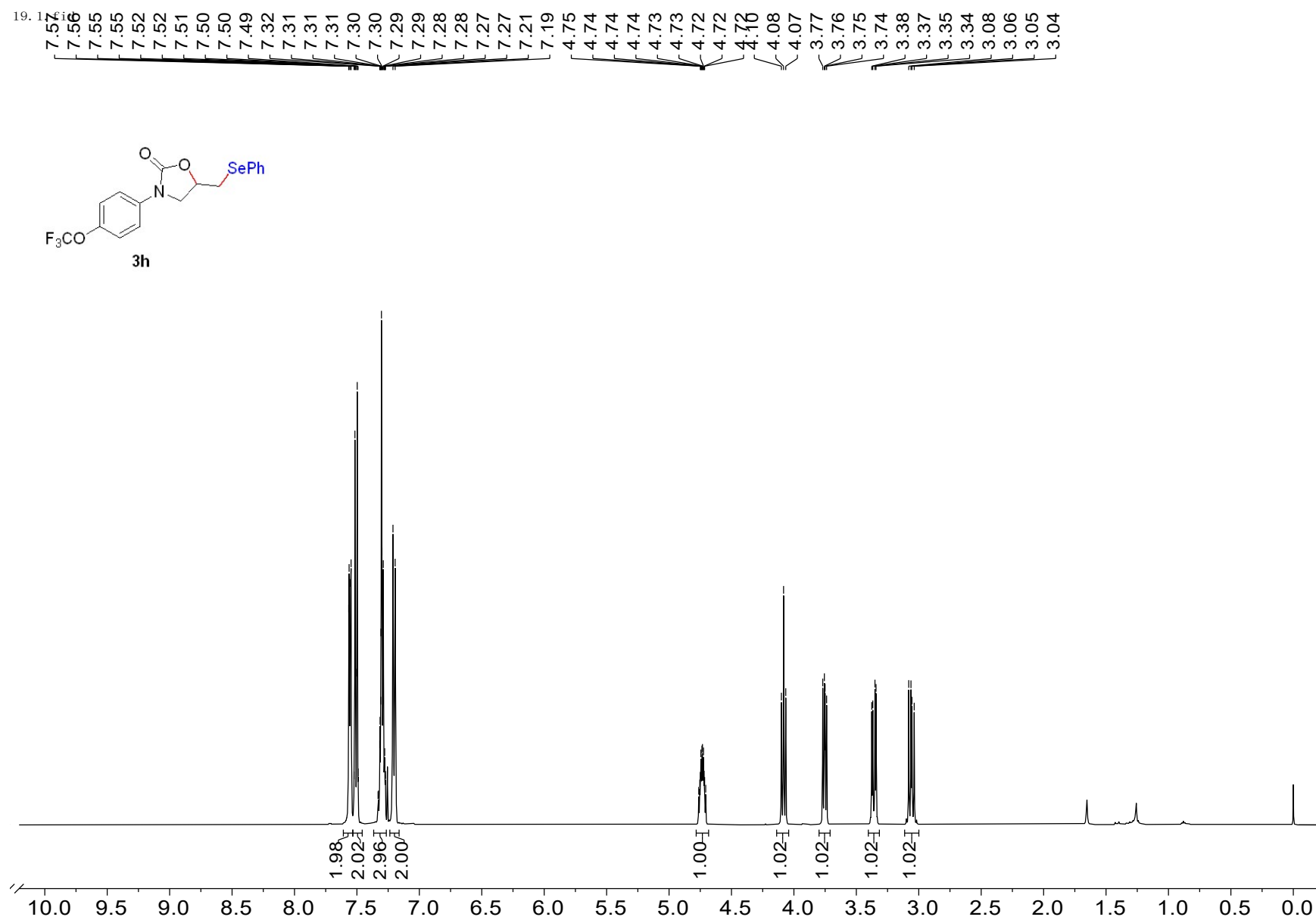


^{19}F NMR (376 MHz, CDCl_3) spectrum of **3g**.

Johnb-bj-F-29, 1. fid
F19CPD

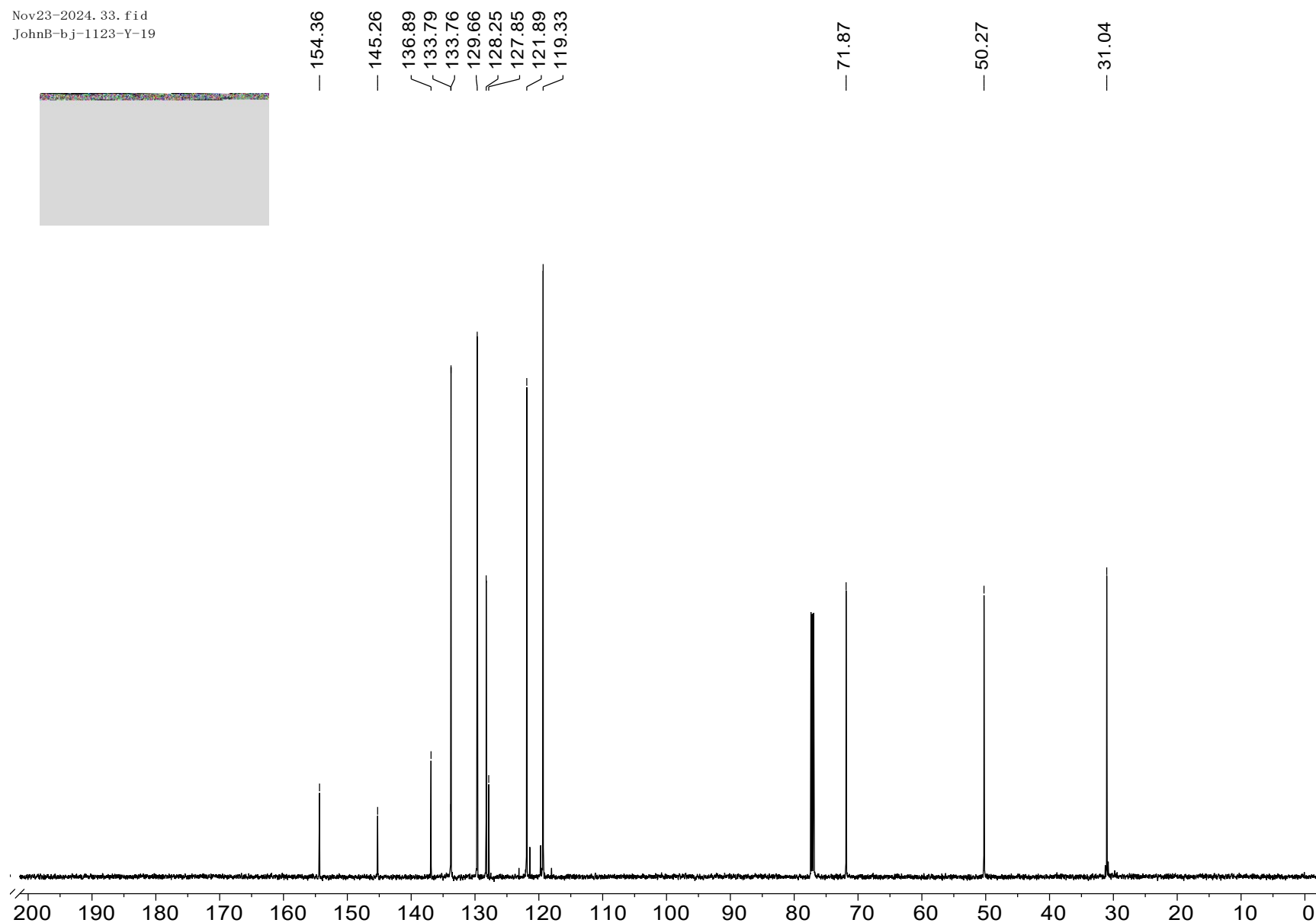


^1H NMR (500 MHz, CDCl_3) spectrum of **3h**.



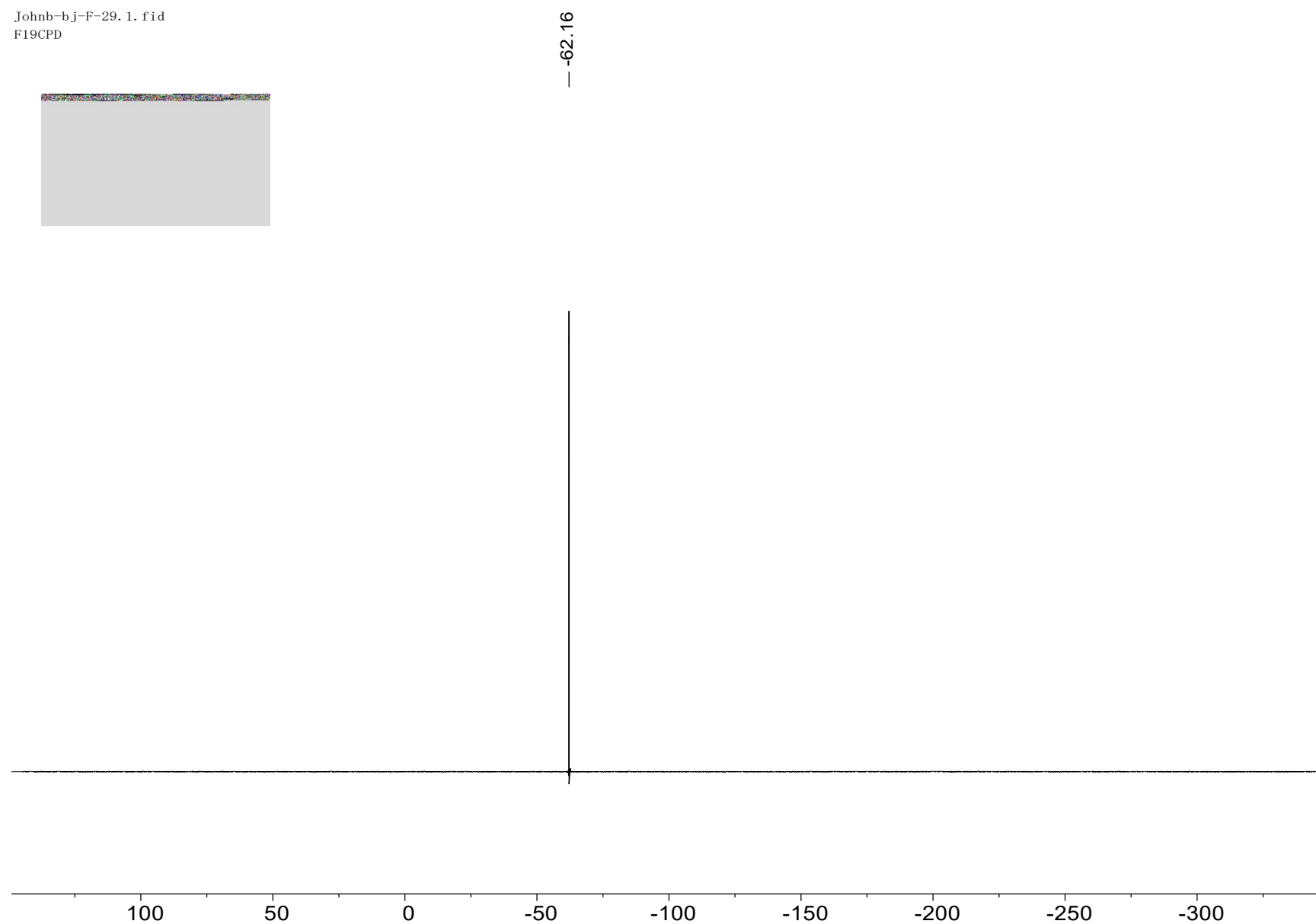
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3h**.

Nov23-2024. 33. fid
JohnB-bj-1123-Y-19



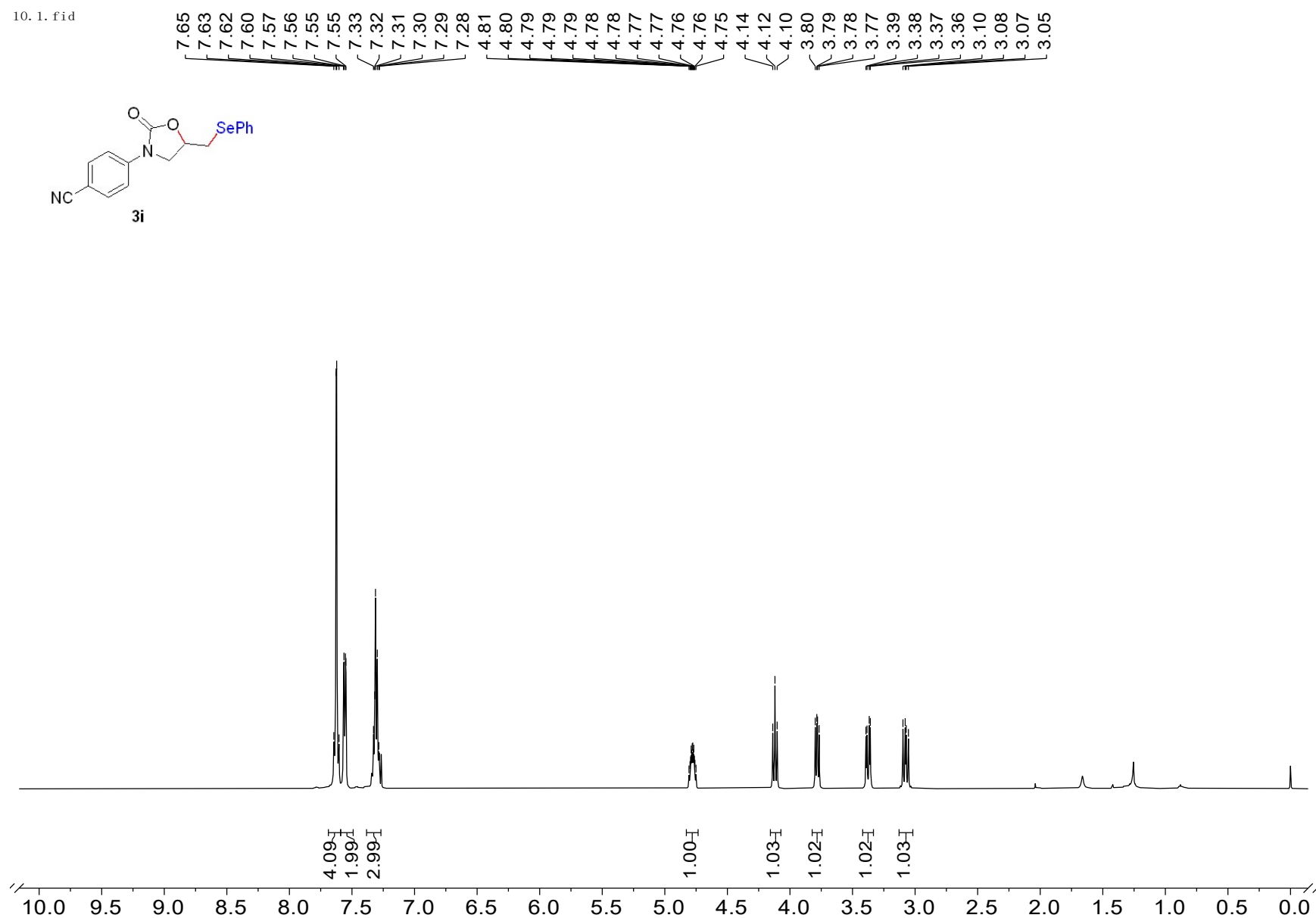
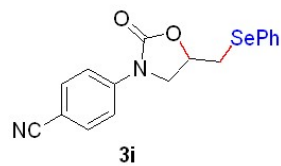
^{19}F NMR (376 MHz, CDCl_3) spectrum of **3h**.

Johnb-bj-F-29, 1. fid
F19CPD



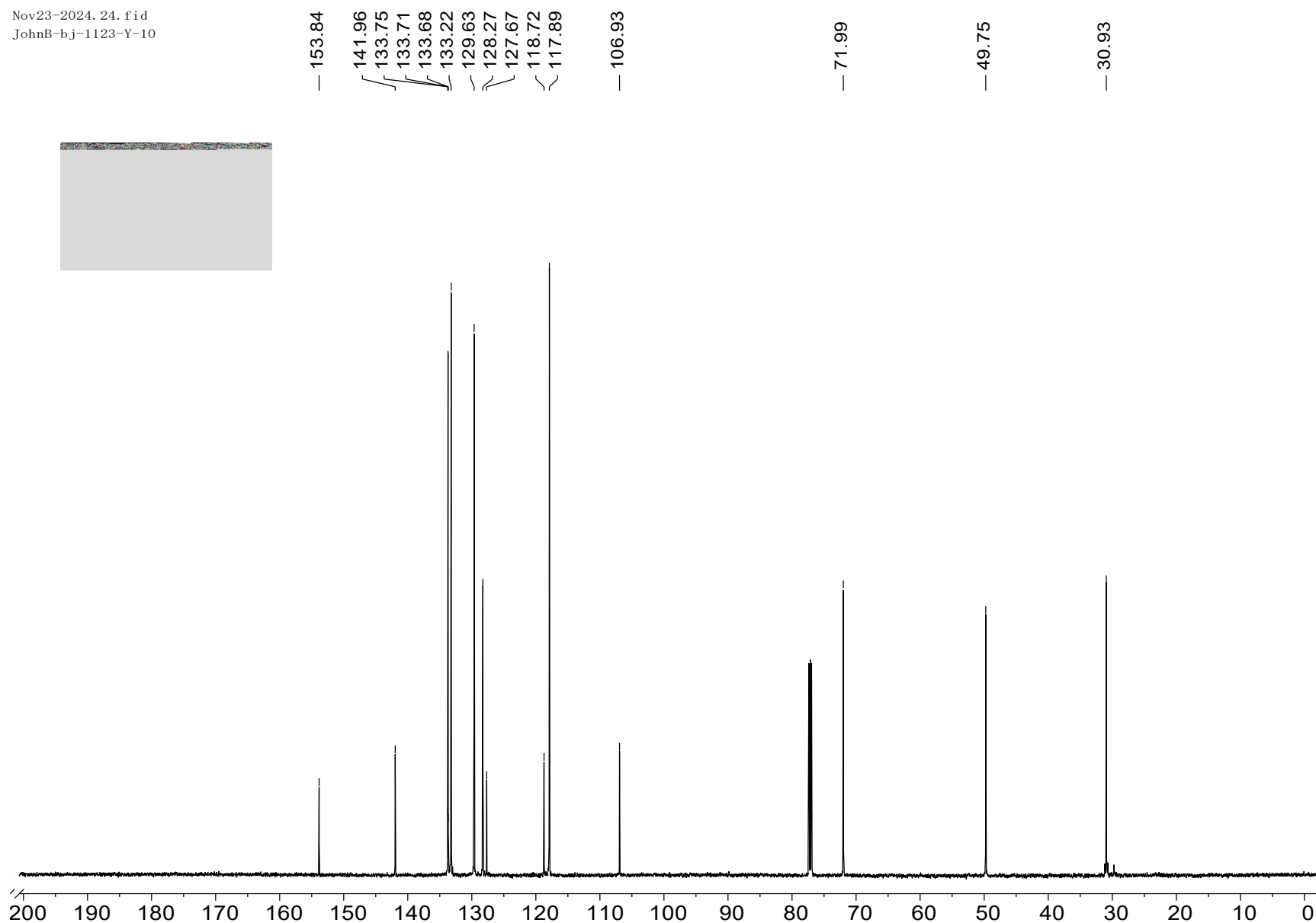
^1H NMR (500 MHz, CDCl_3) spectrum of **3i**.

10.1.fid

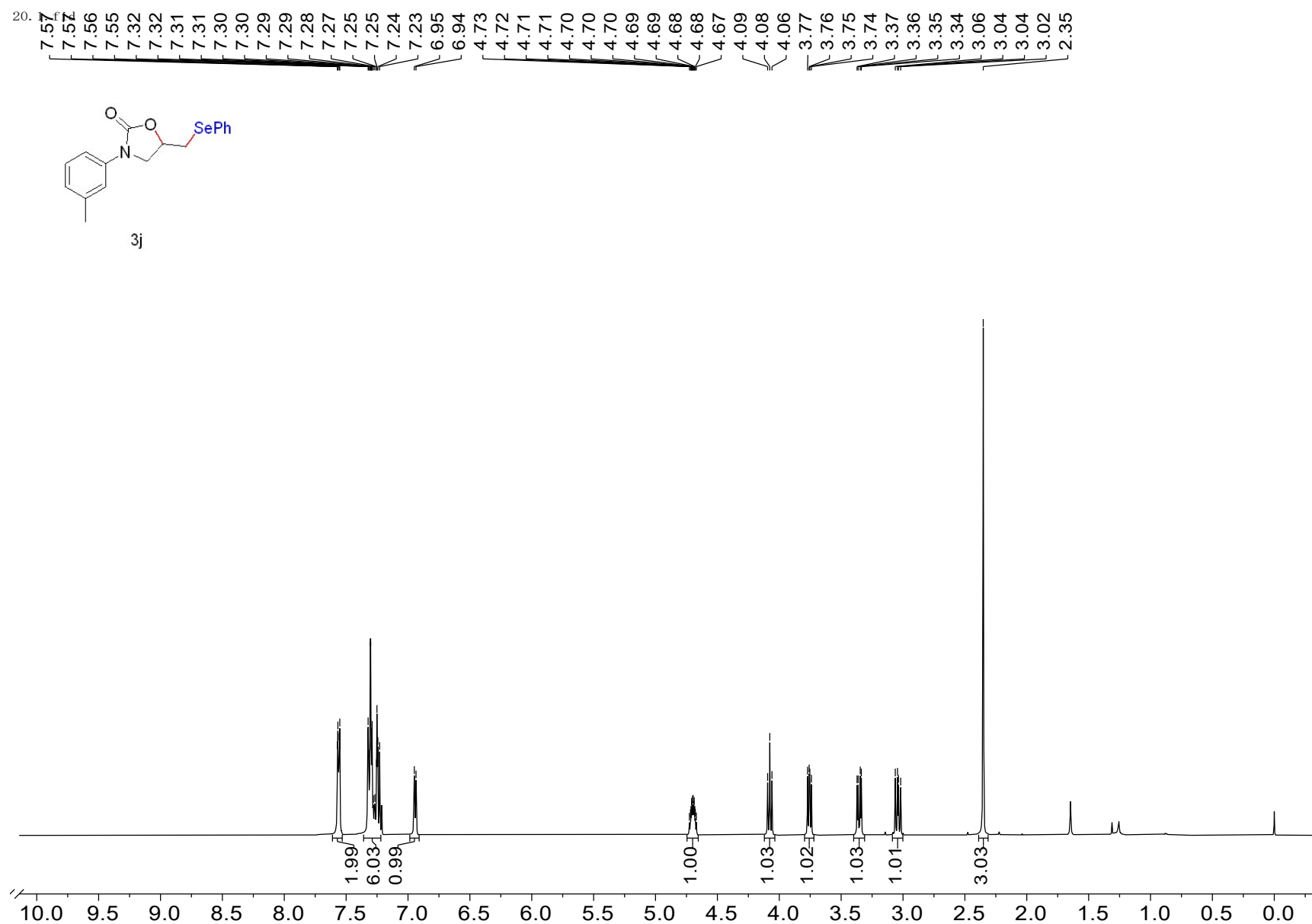


^{13}C NMR (151 MHz, CDCl_3) spectrum of **3i**.

Nov23-2024, 24. fid
JohnB-bj-1123-Y-10

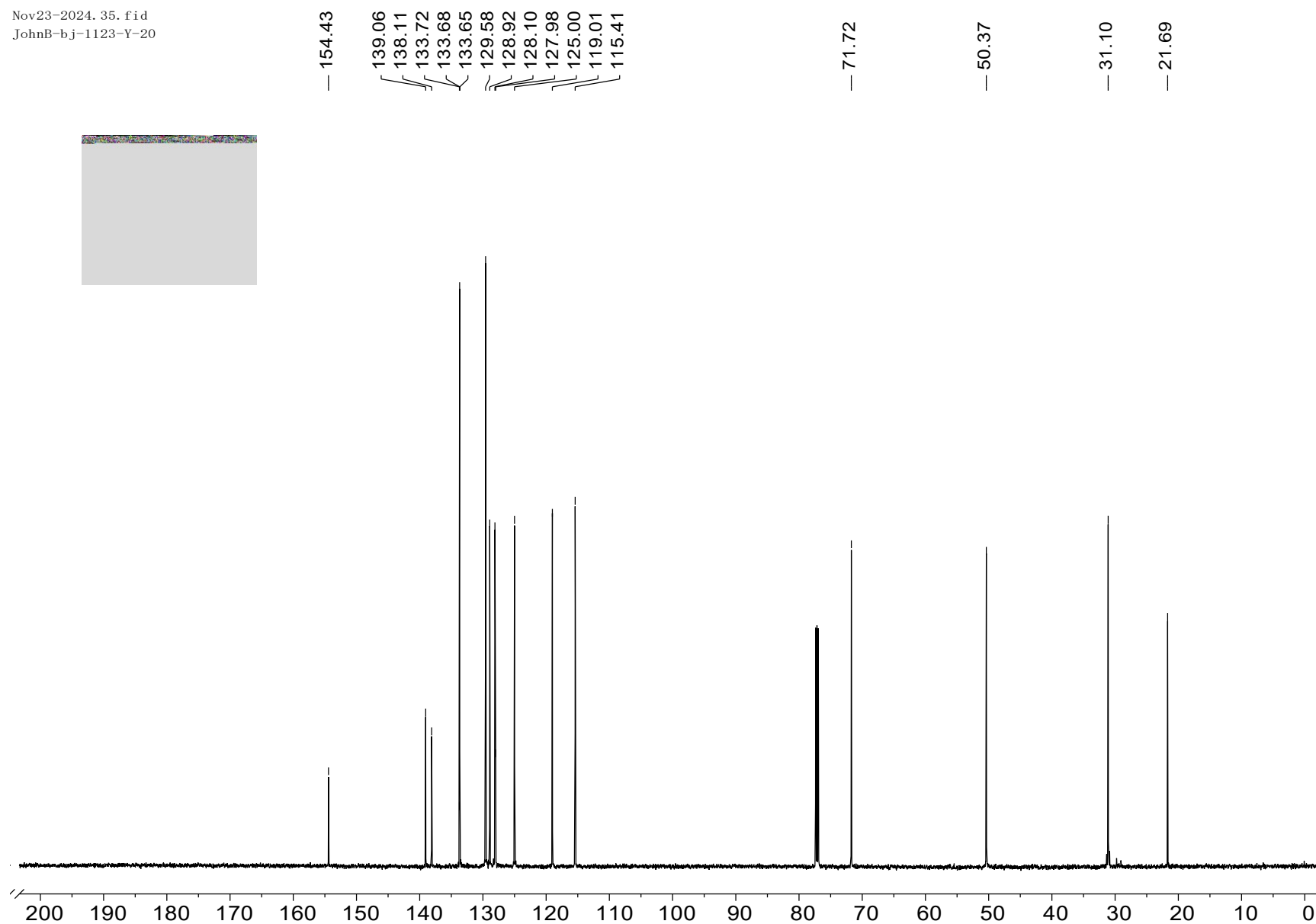


^1H NMR (500 MHz, CDCl_3) spectrum of **3j**.

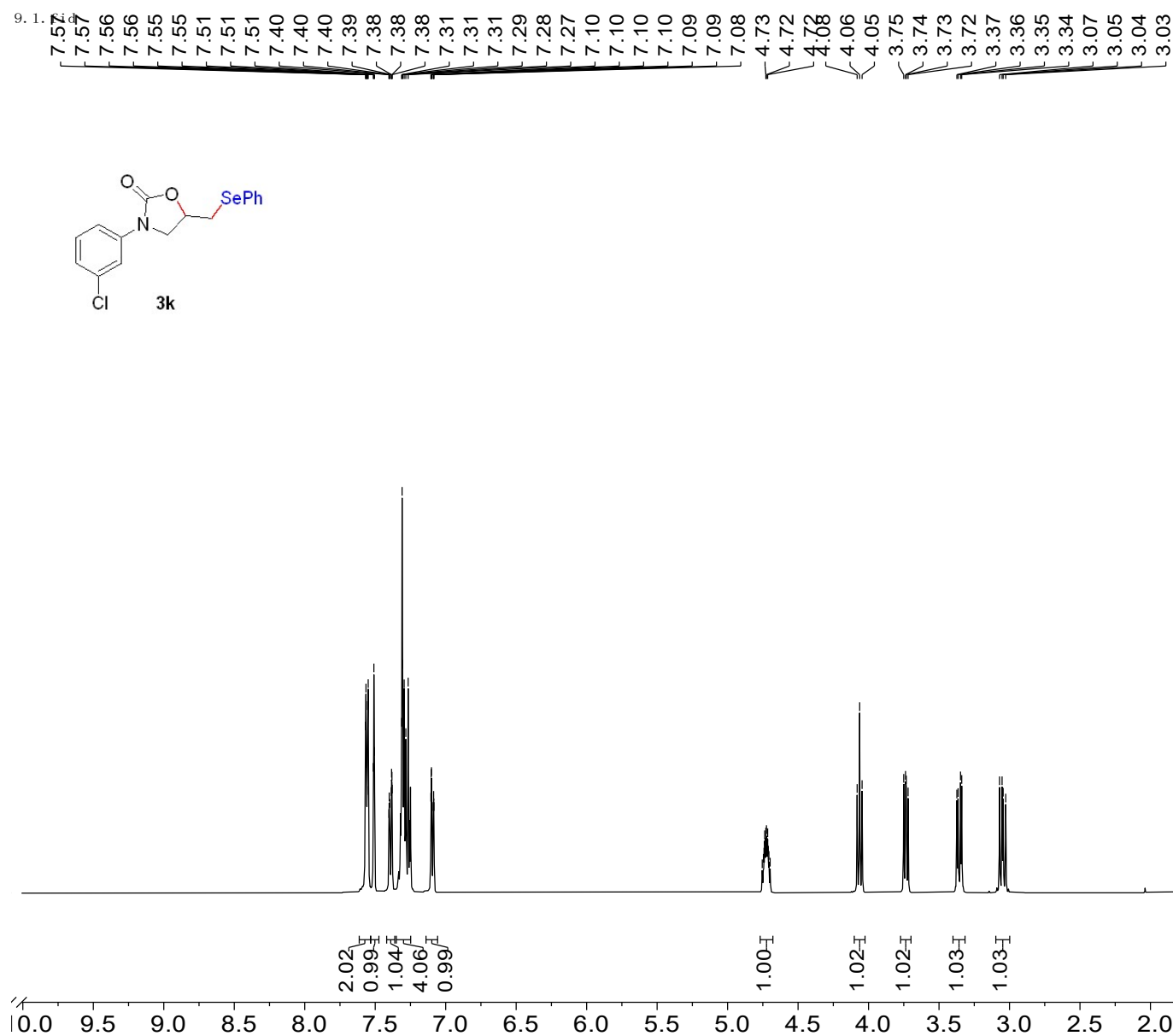


^{13}C NMR (151 MHz, CDCl_3) spectrum of **3j**.

Nov23-2024. 35. fid
JohnB-bj-1123-Y-20

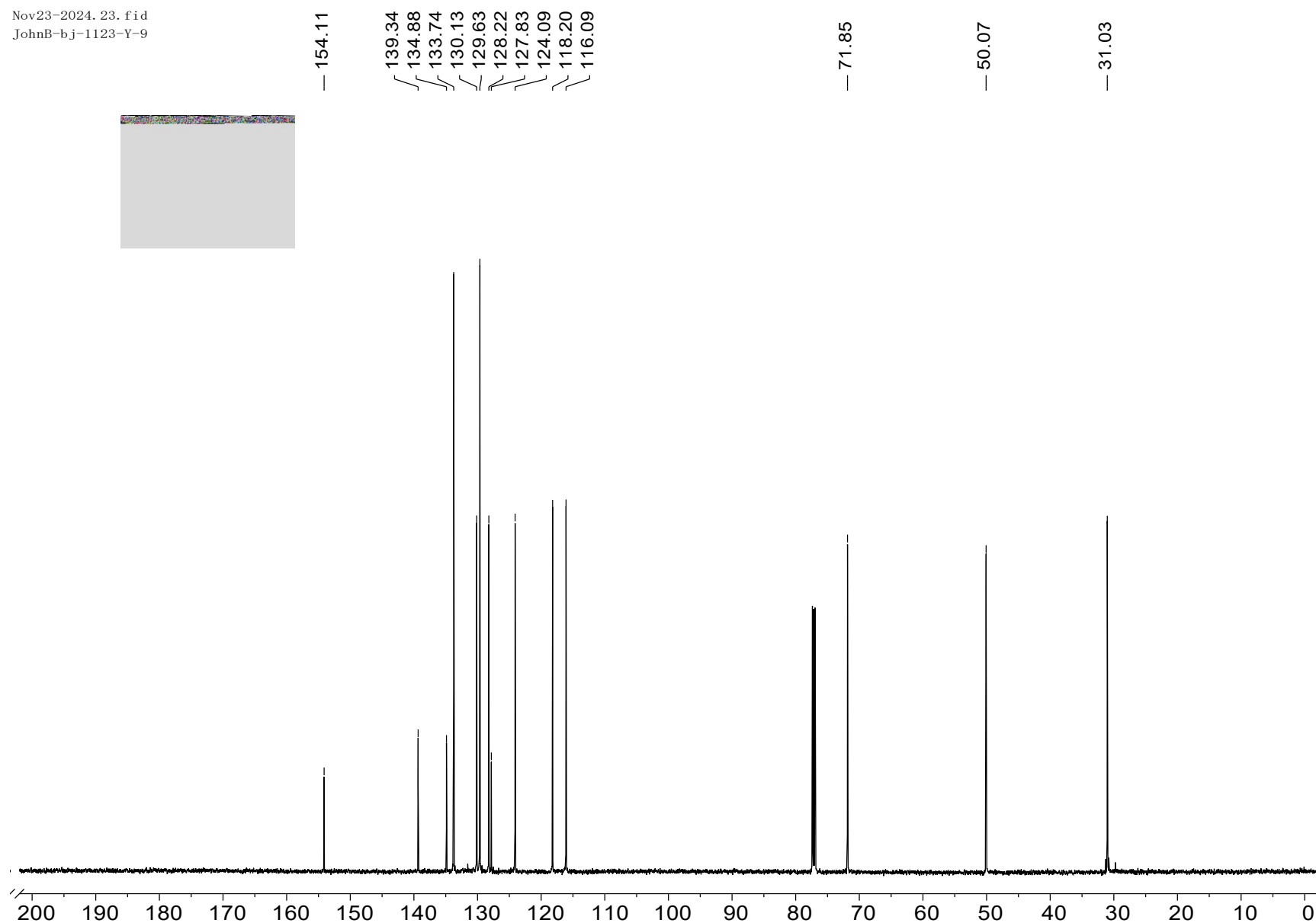


^1H NMR (500 MHz, CDCl_3) spectrum of **3k**.

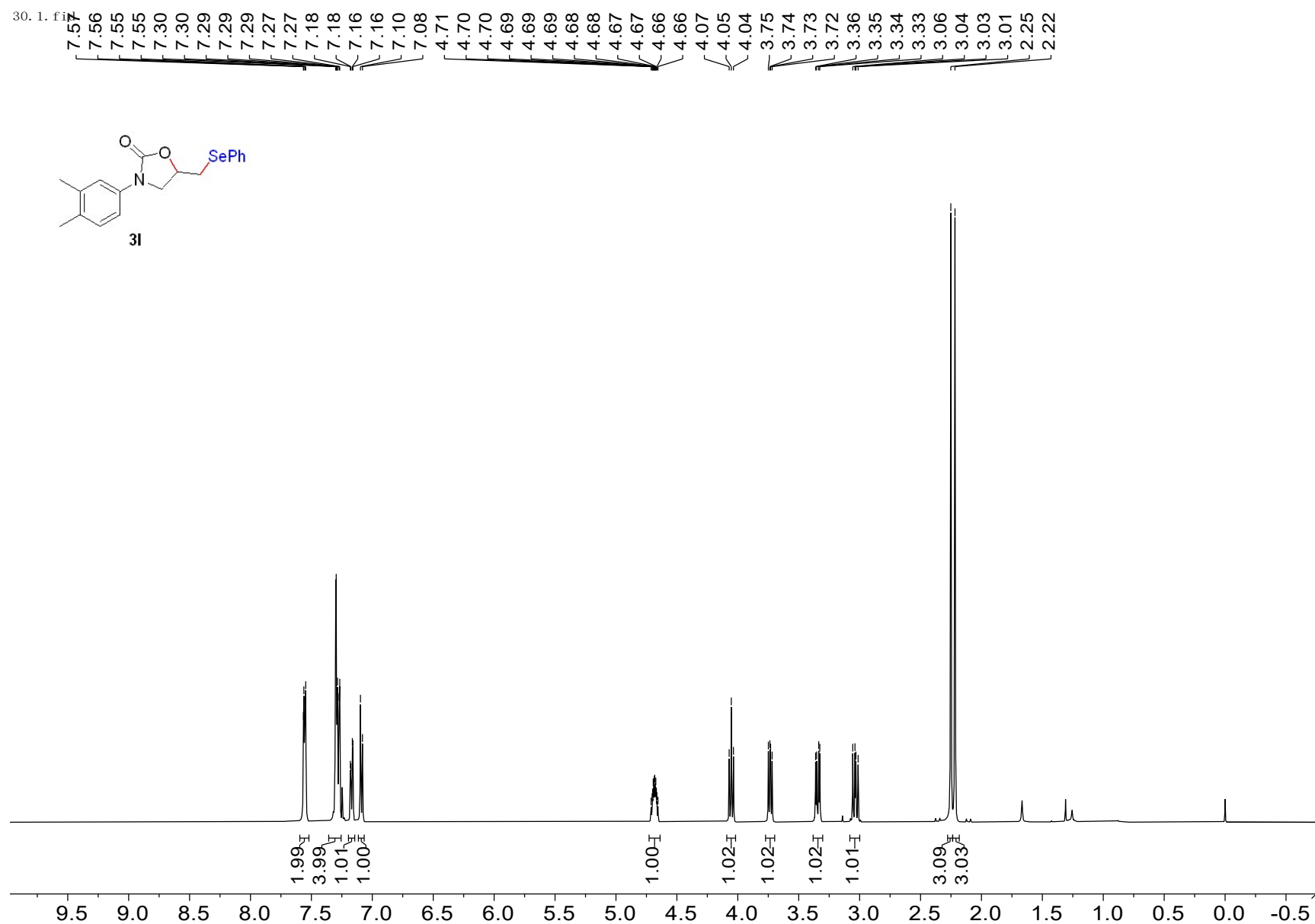


^{13}C NMR (151 MHz, CDCl_3) spectrum of **3k**.

Nov23-2024, 23. fid
JohnB-bj-1123-Y-9

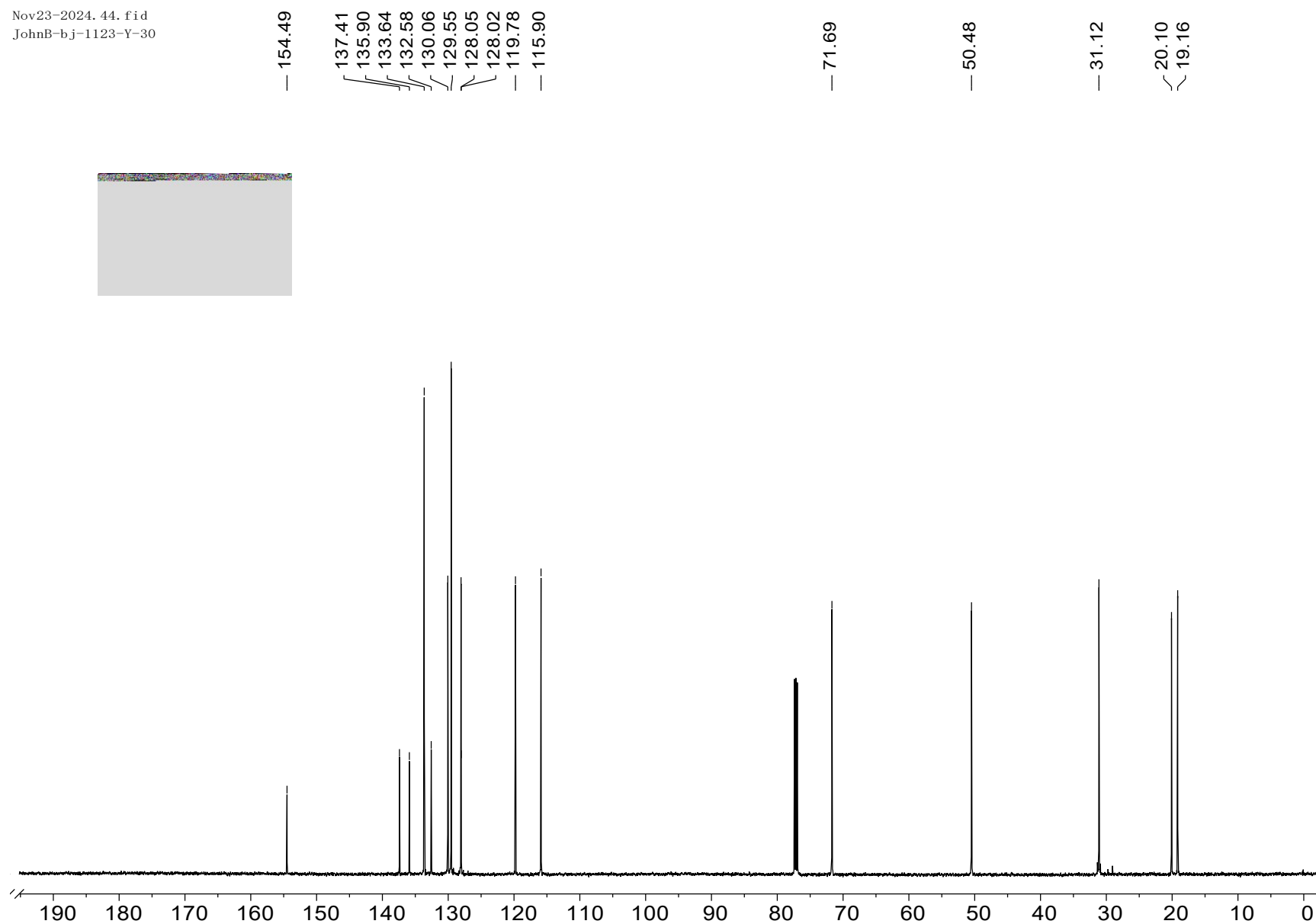


^1H NMR (500 MHz, CDCl_3) spectrum of **3l**.



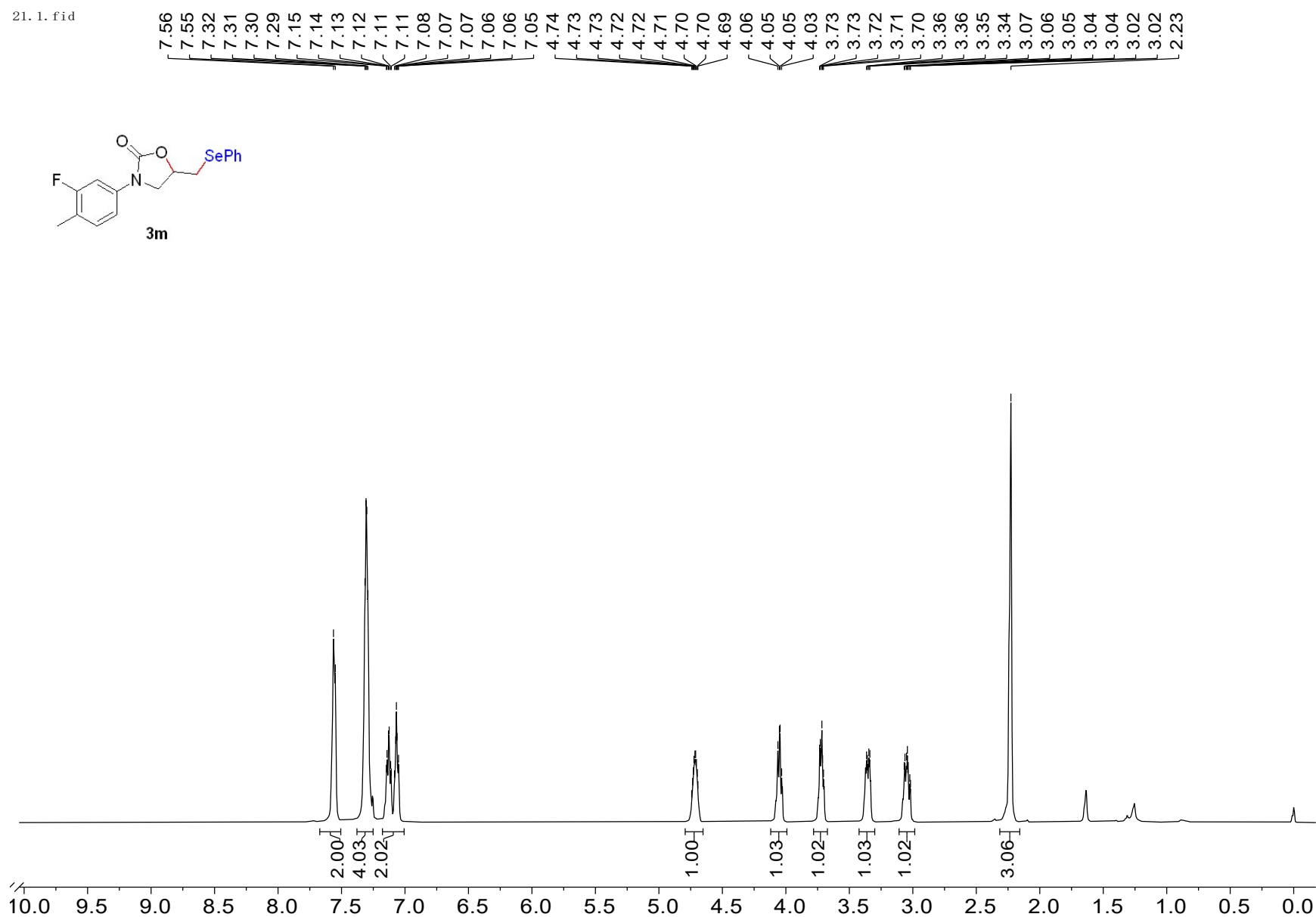
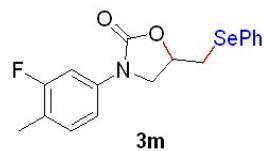
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3l**.

Nov23-2024. 44. fid
JohnB-bj-1123-Y-30



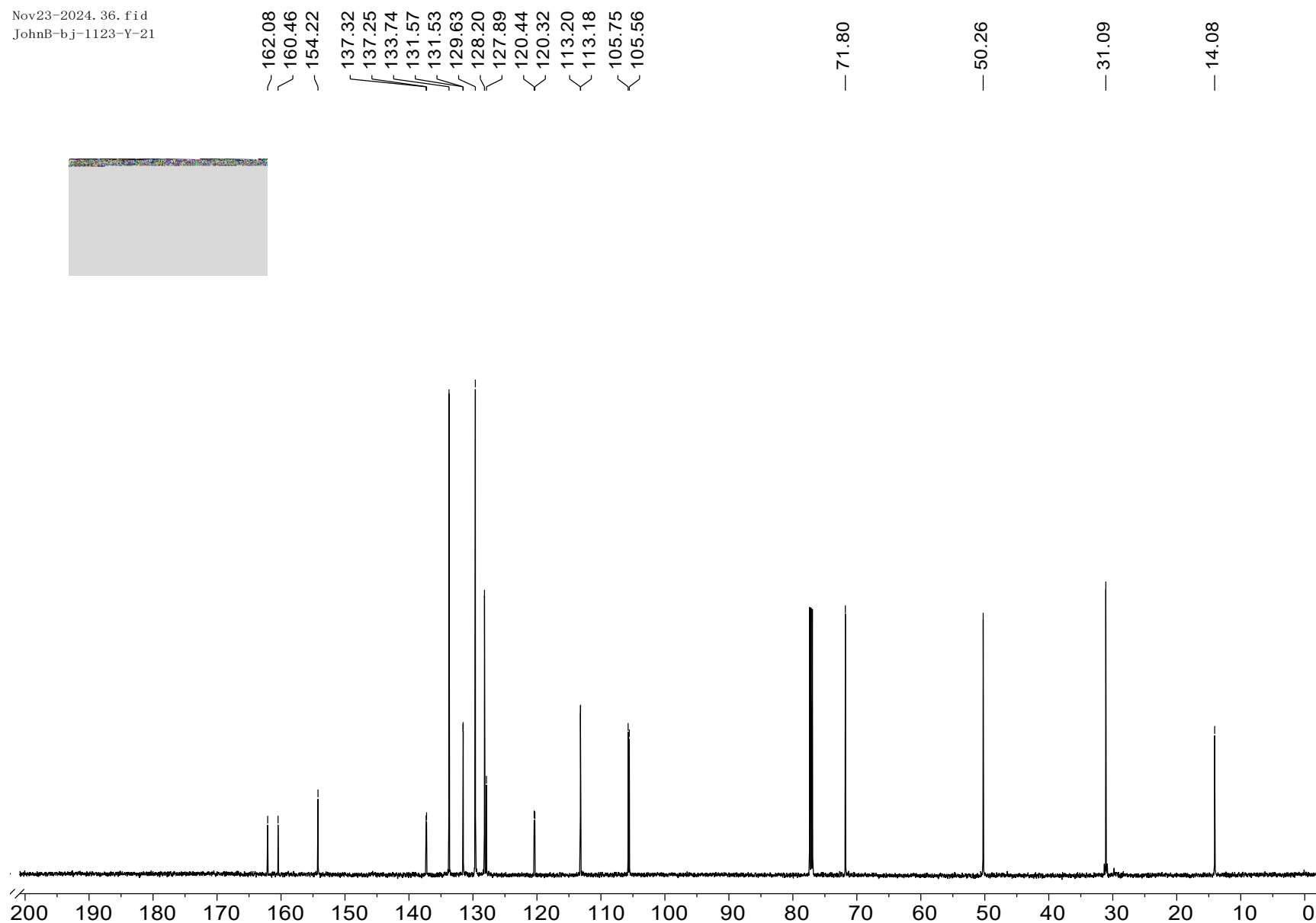
^1H NMR (500 MHz, CDCl_3) spectrum of **3m**.

21.1.fid



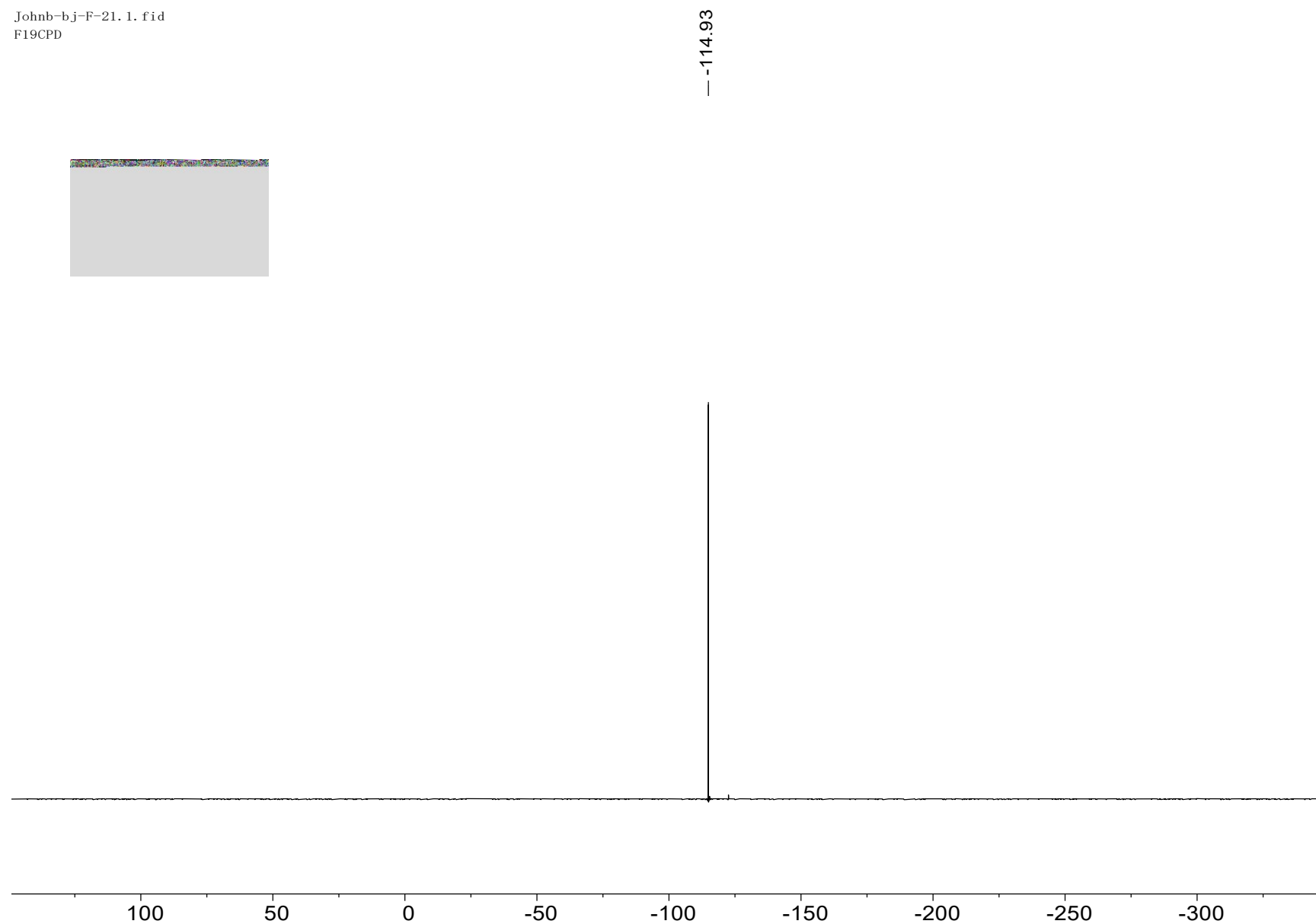
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3m**.

Nov23-2024, 36, fid
JohnB-bj-1123-Y-21

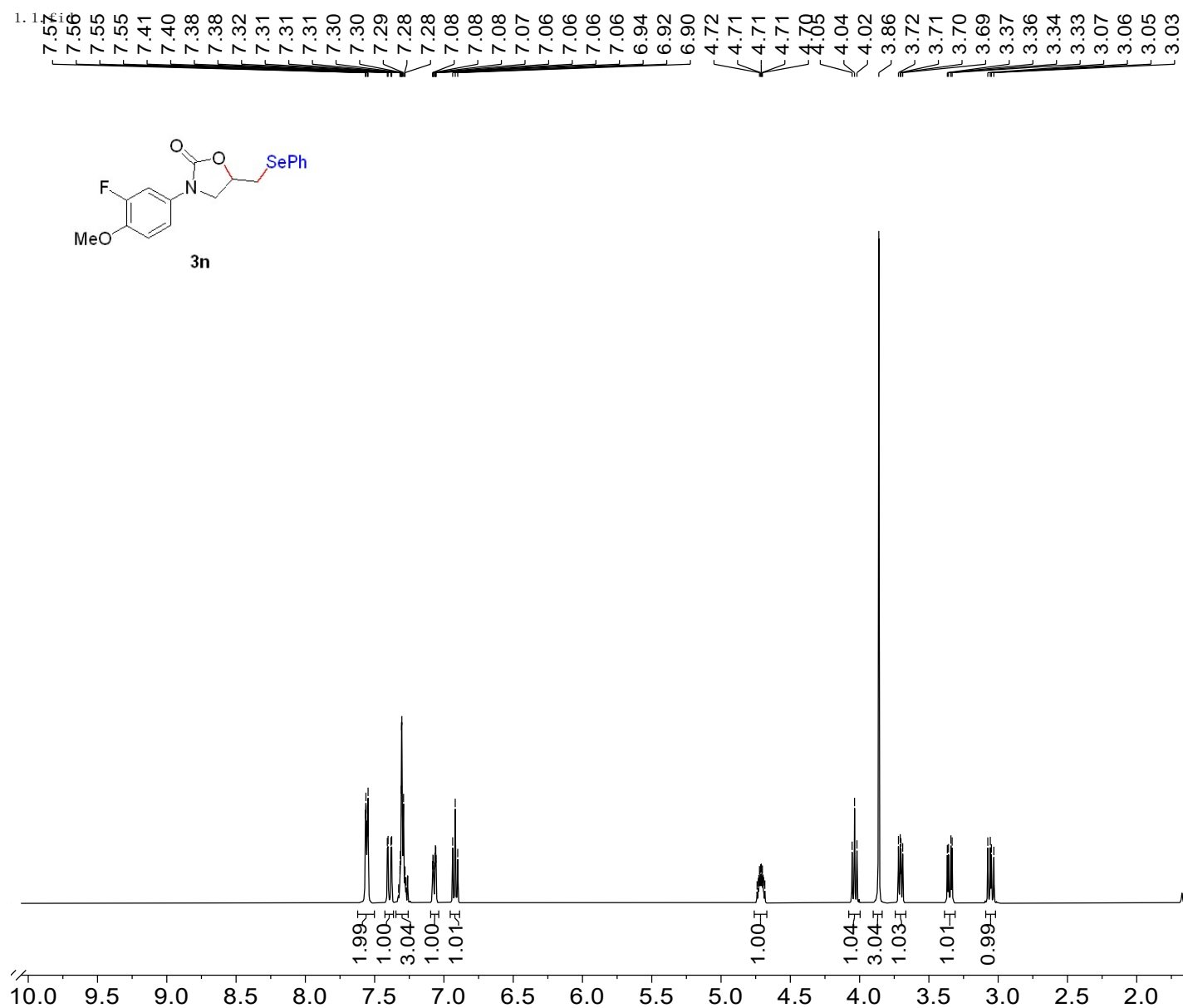


^{19}F NMR (376 MHz, CDCl_3) spectrum of **3m**.

Johnb-bj-F-21. 1. fid
F19CPD

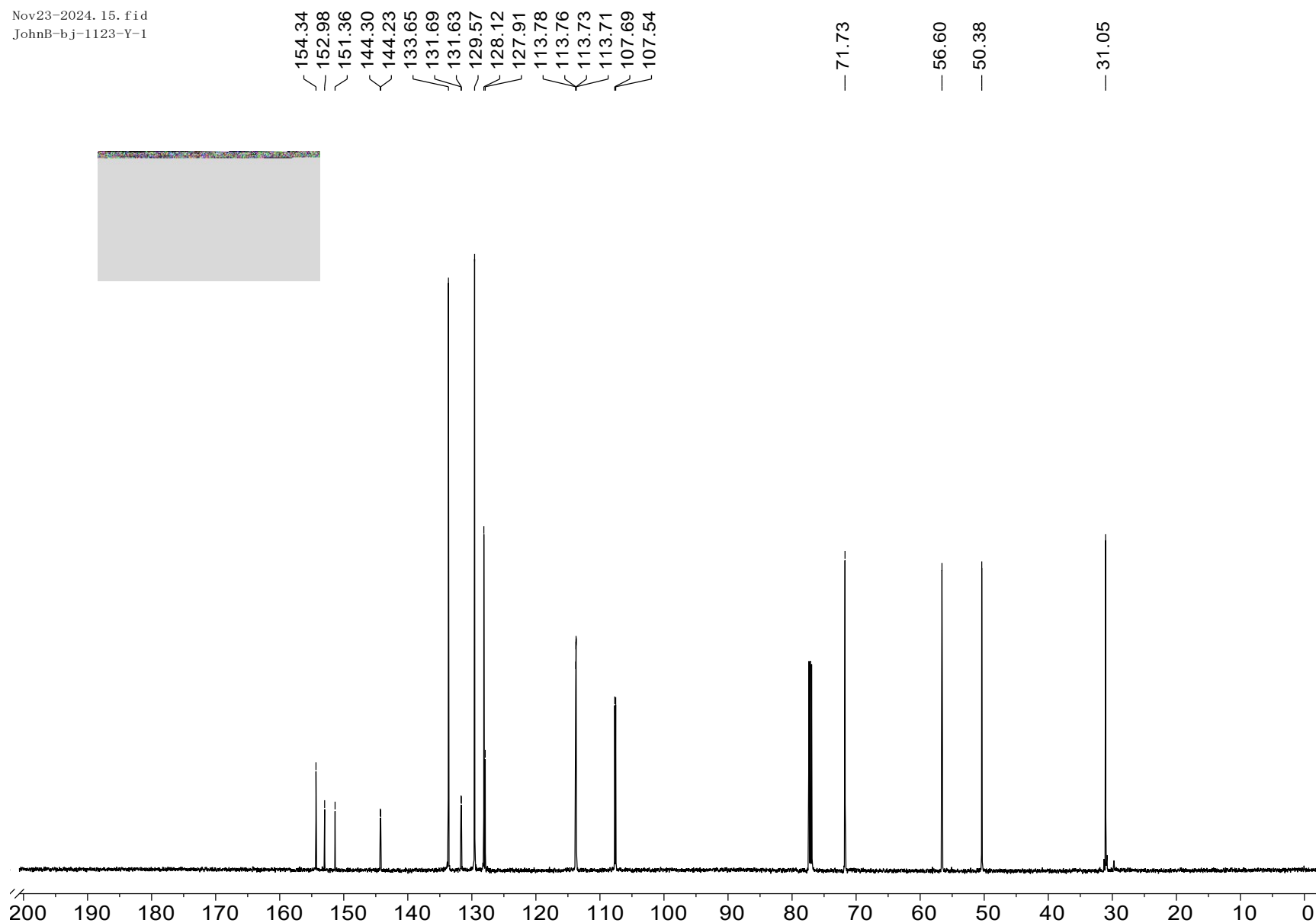


^1H NMR (500 MHz, CDCl_3) spectrum of **3n**.



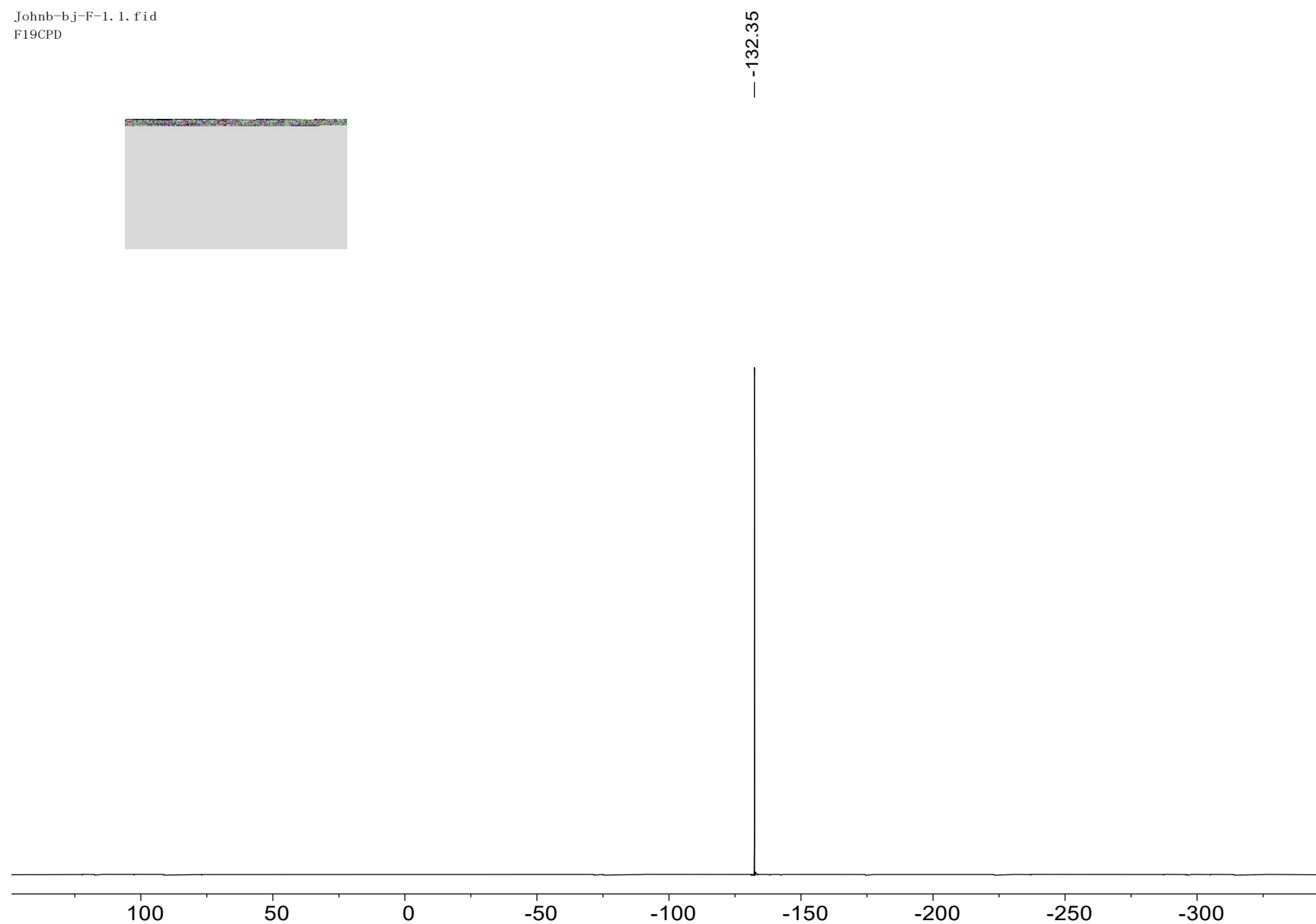
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3n**.

Nov23-2024. 15. fid
JohnB-bj-1123-Y-1

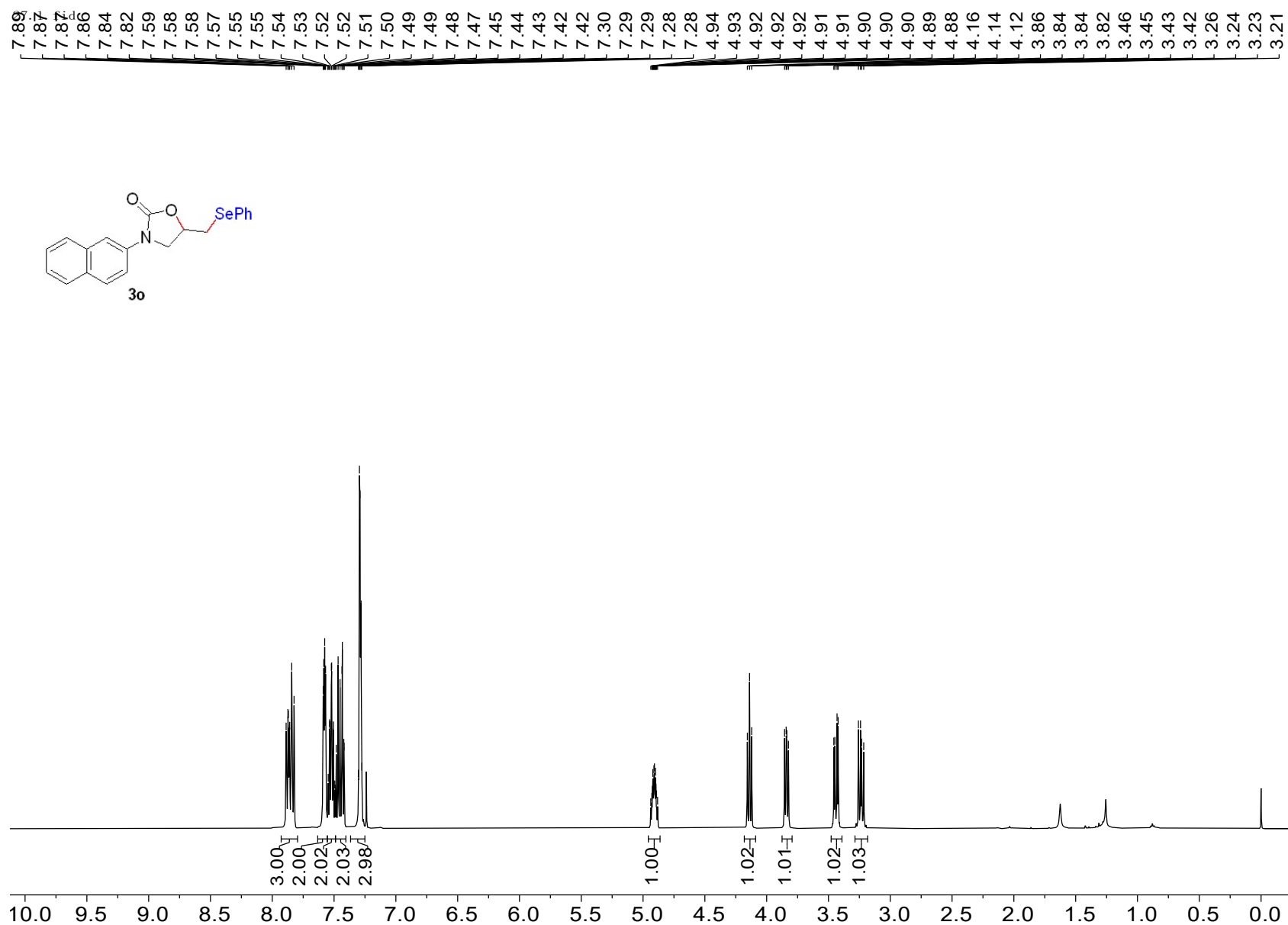


^{19}F NMR (376 MHz, CDCl_3) spectrum of **3n**.

Johnb-bj-F-1. 1. fid
F19CPD

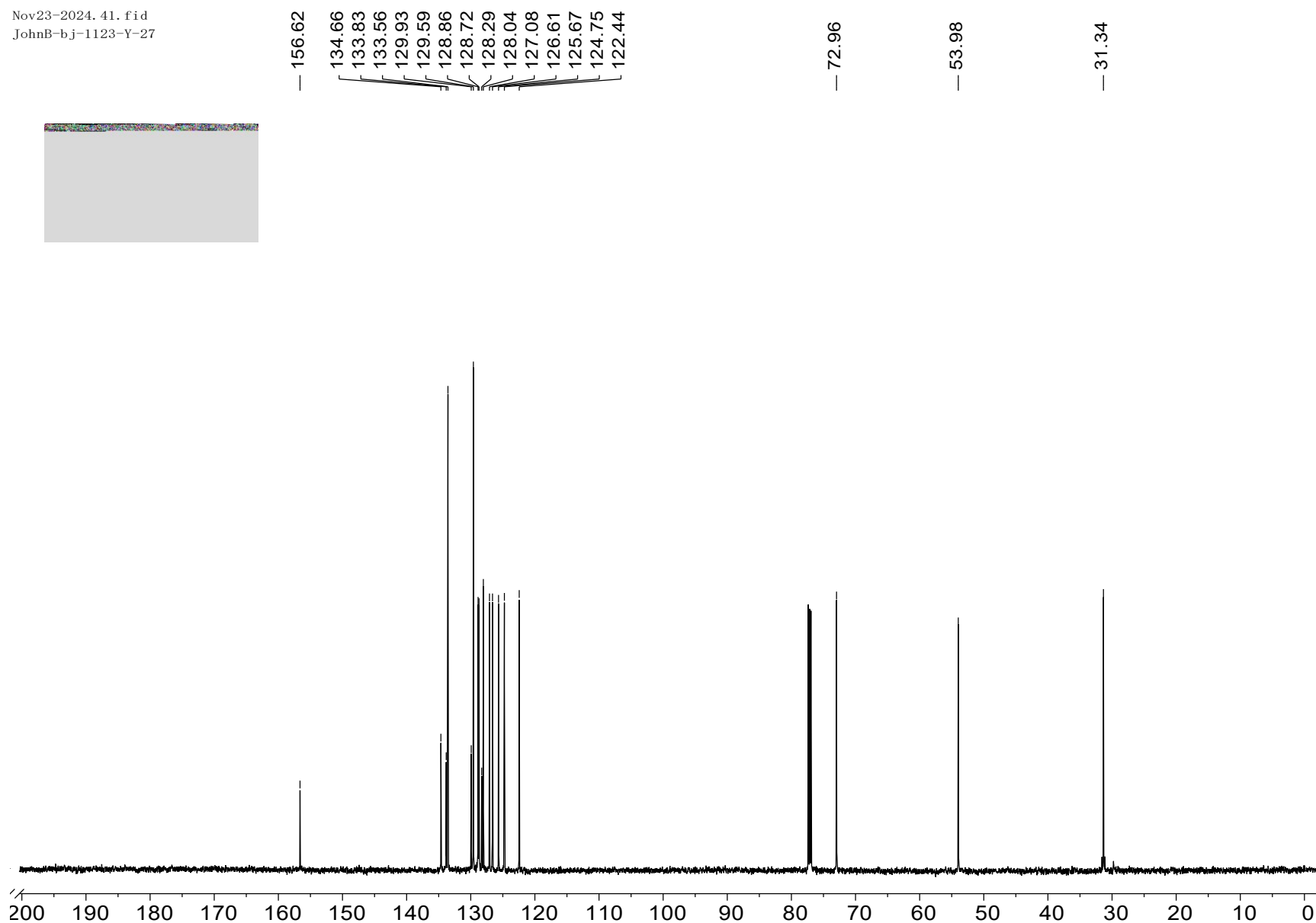


^1H NMR (500 MHz, CDCl_3) spectrum of **30**.



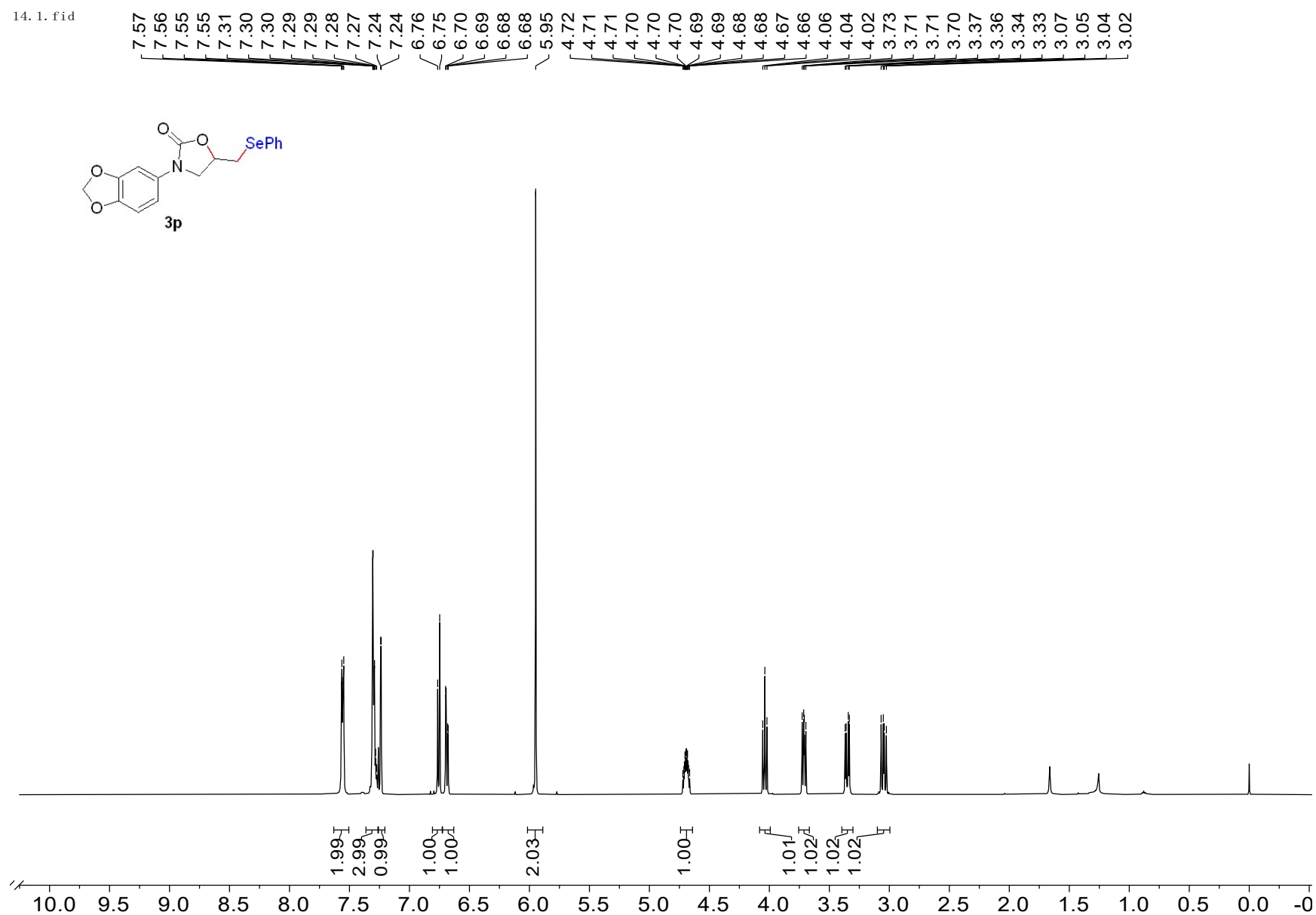
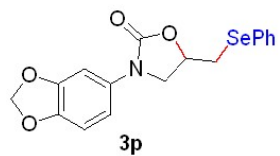
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3o**.

Nov23-2024, 41, fid
JohnB-bj-1123-Y-27



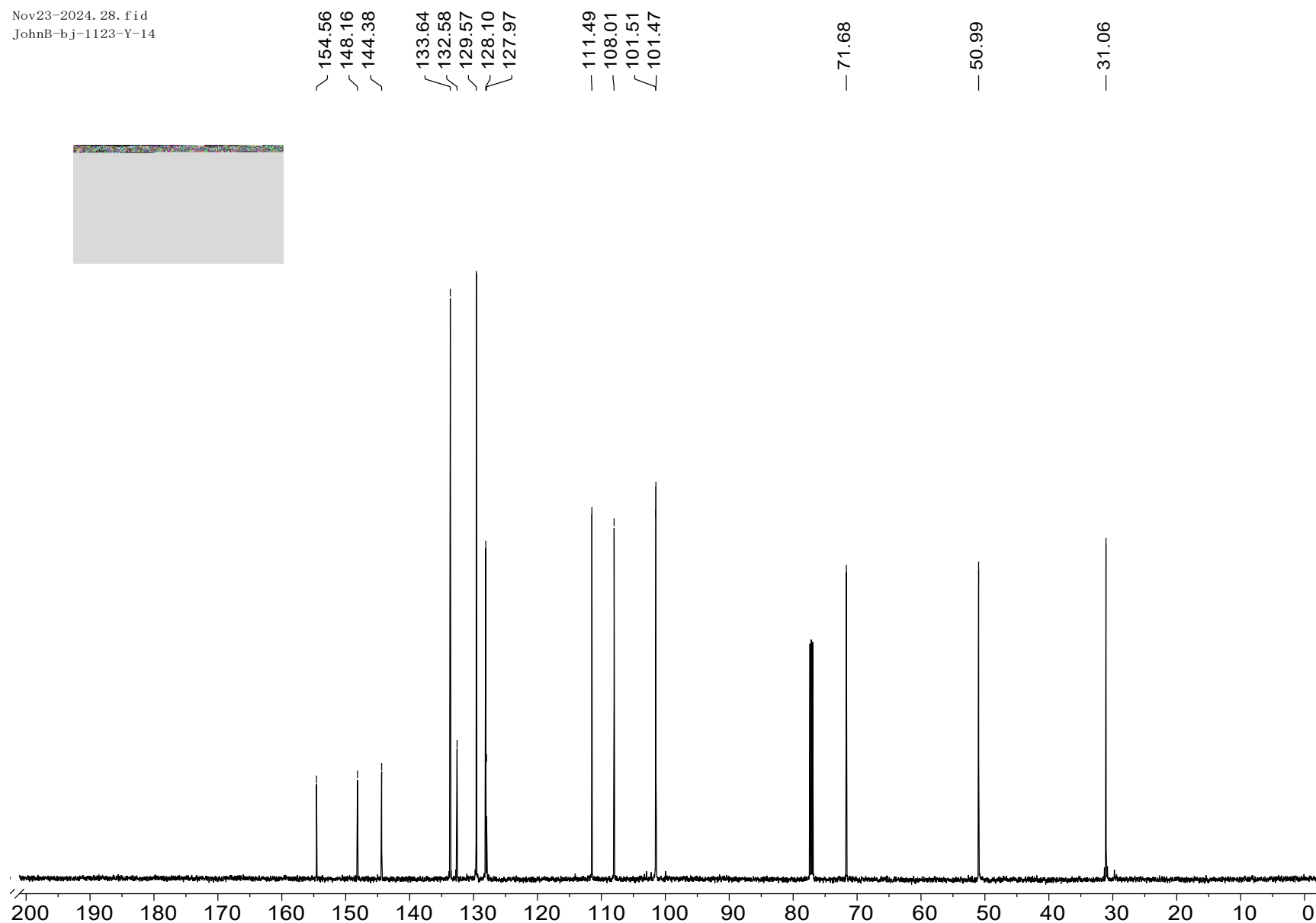
^1H NMR (500 MHz, CDCl_3) spectrum of **3p**.

14. 1. fid



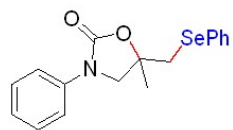
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3p**.

Nov23-2024. 28. fid
JohnB-bj-1123-Y-14

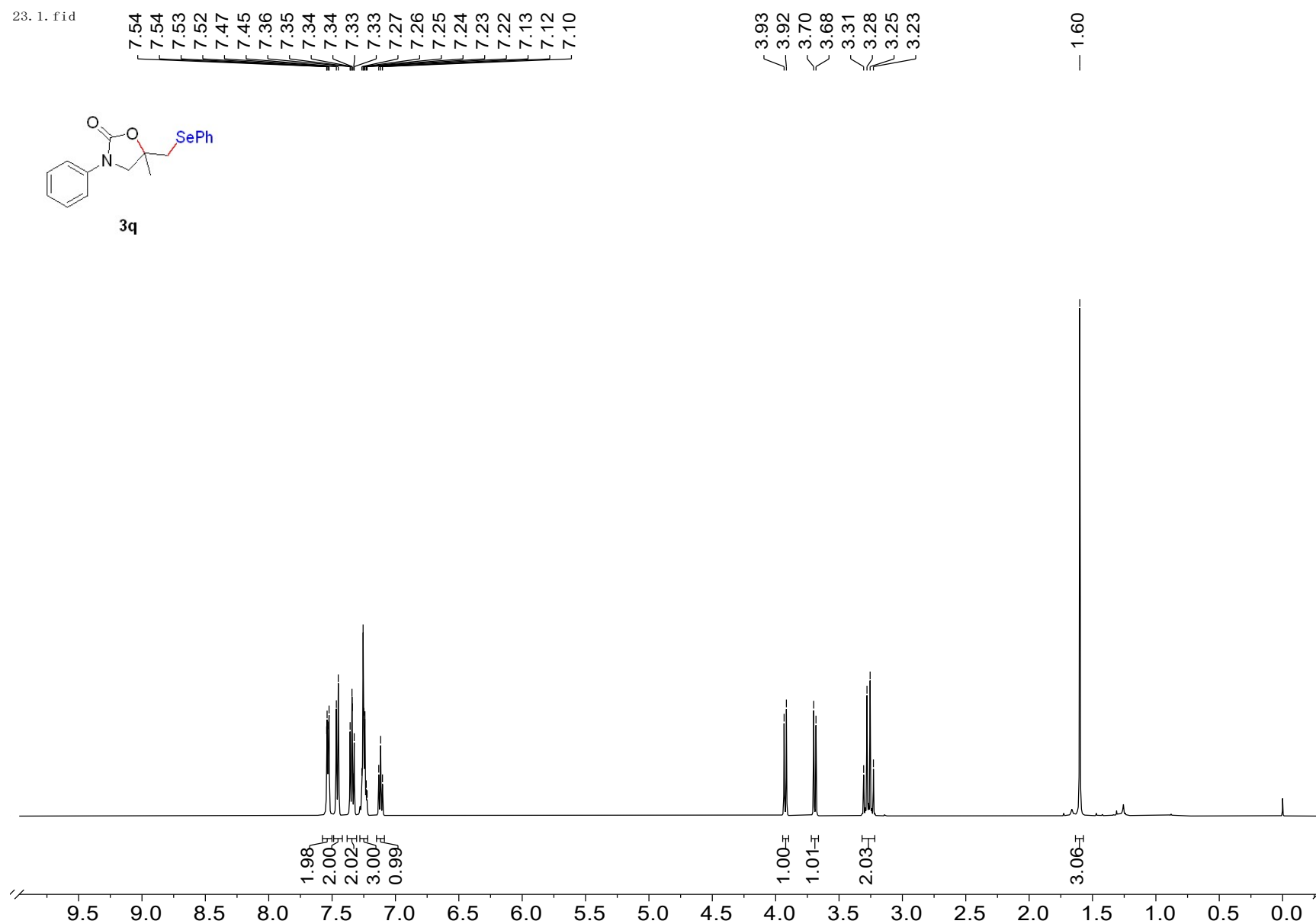


^1H NMR (500 MHz, CDCl_3) spectrum of **3q**.

23.1.fid

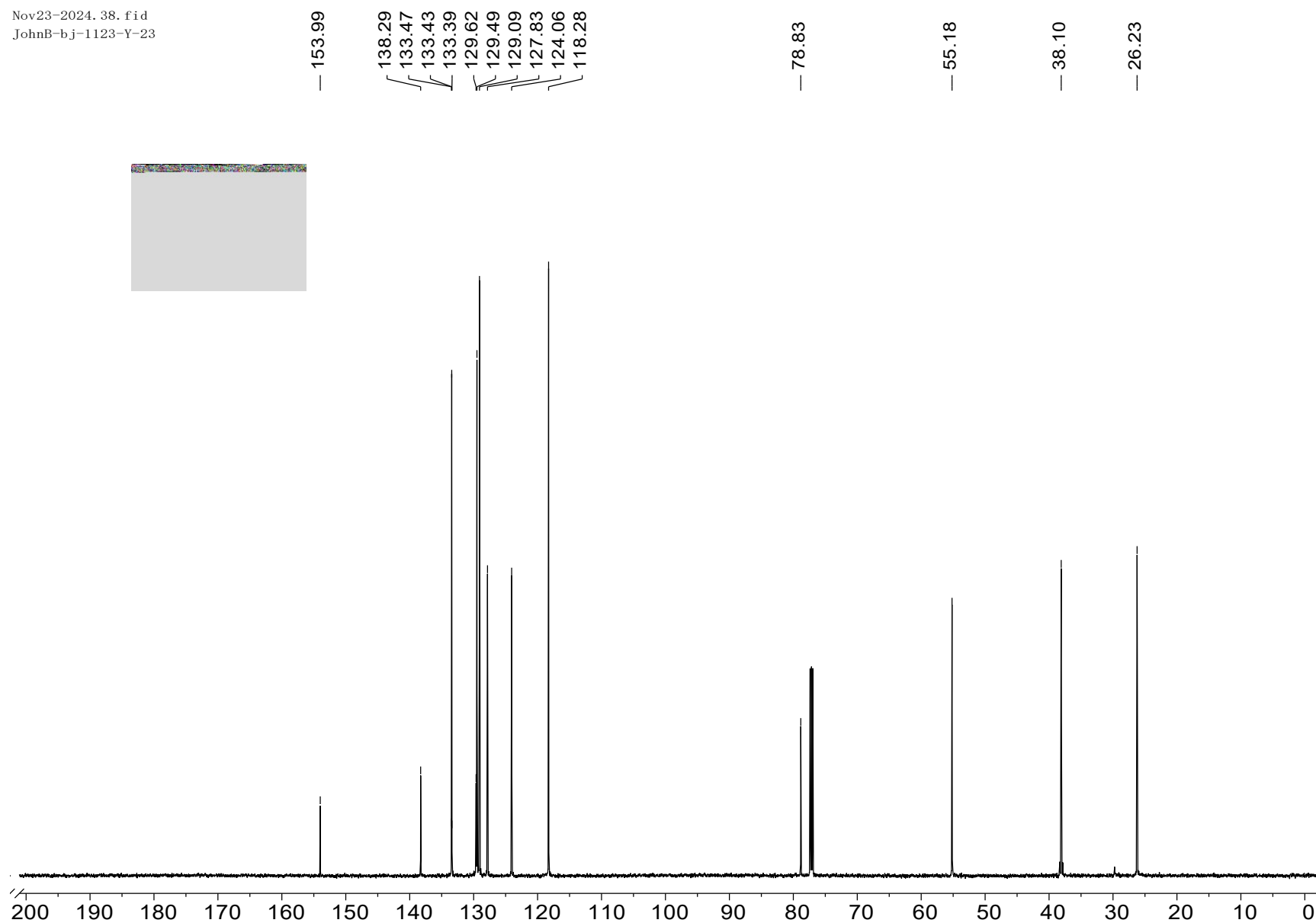


3q

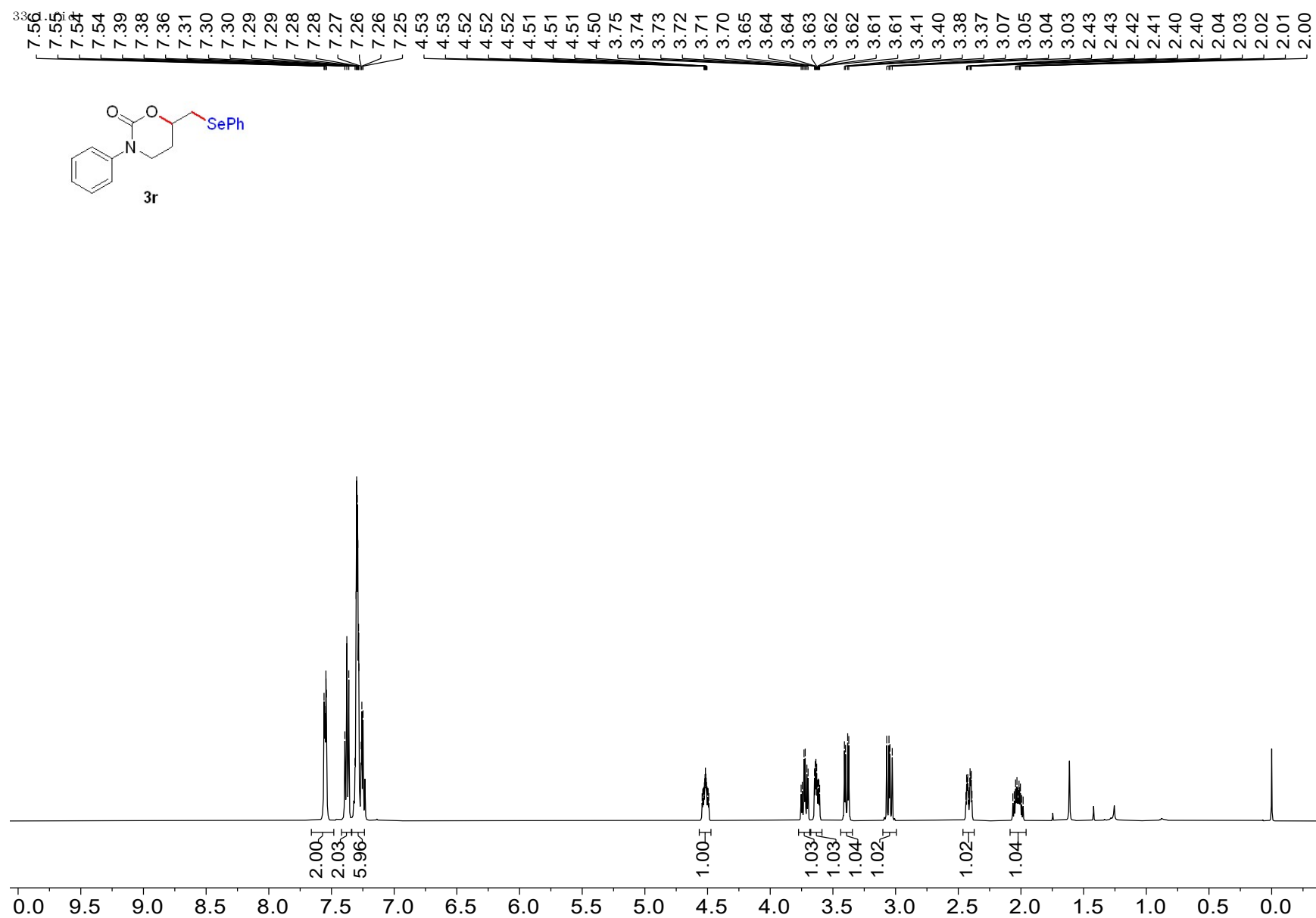


^{13}C NMR (151 MHz, CDCl_3) spectrum of **3q**.

Nov23-2024, 38, fid
JohnB-bj-1123-Y-23

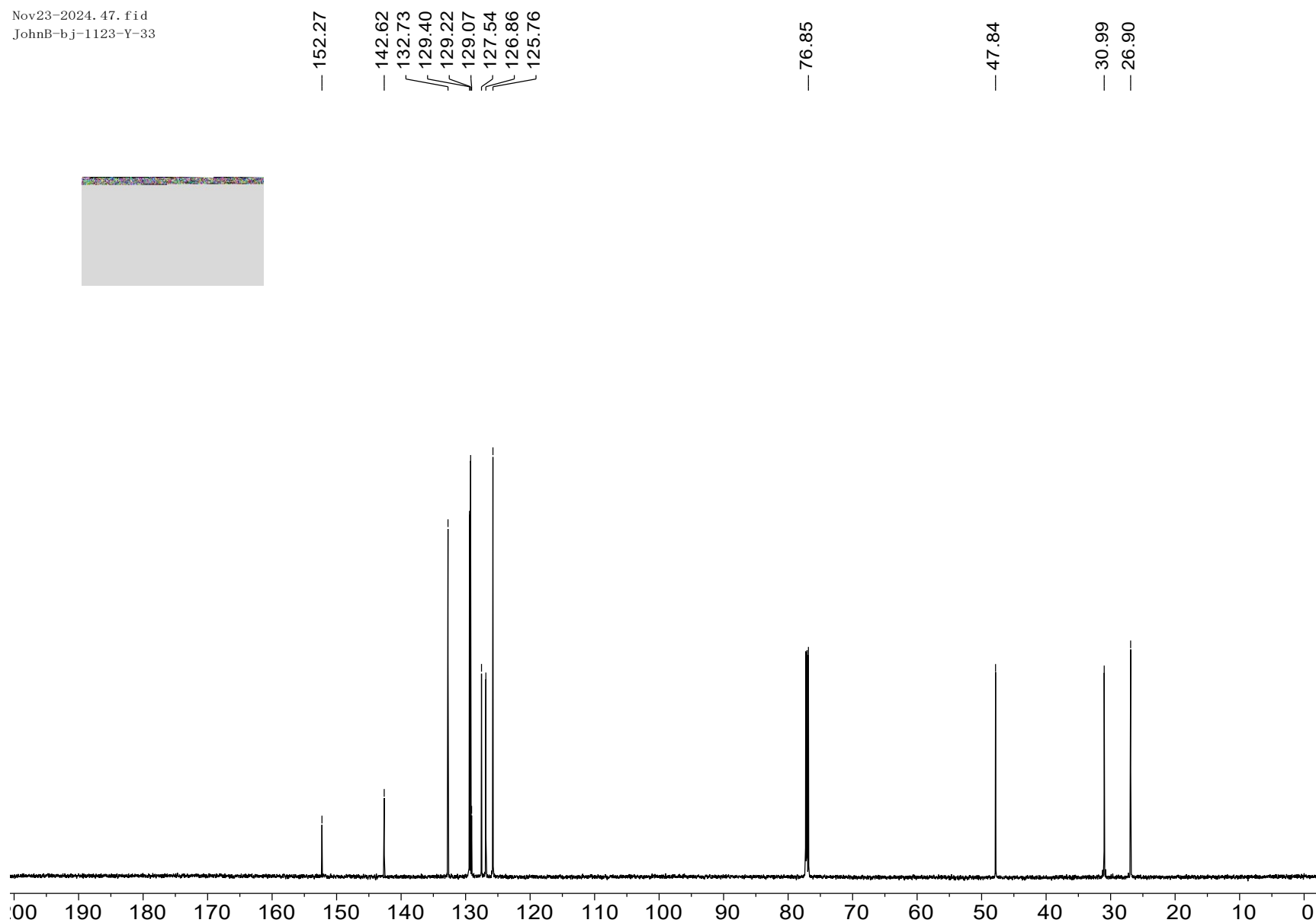


^1H NMR (500 MHz, CDCl_3) spectrum of **3r**.

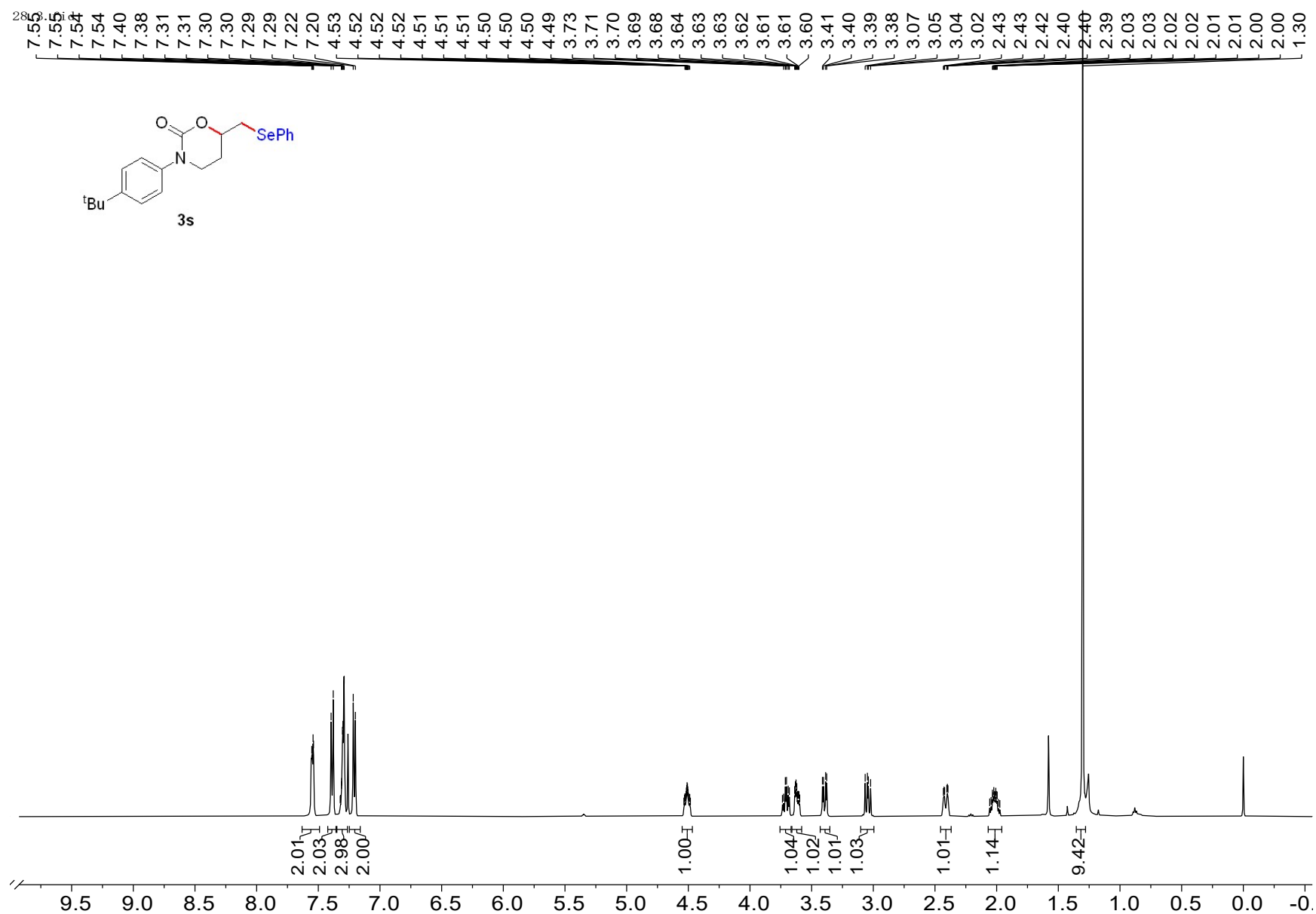


^{13}C NMR (151 MHz, CDCl_3) spectrum of **3r**.

Nov23-2024, 47. fid
JohnB-bj-1123-Y-33



^1H NMR (500 MHz, CDCl_3) spectrum of **3s**.



^{13}C NMR (151 MHz, CDCl_3) spectrum of **3s**.

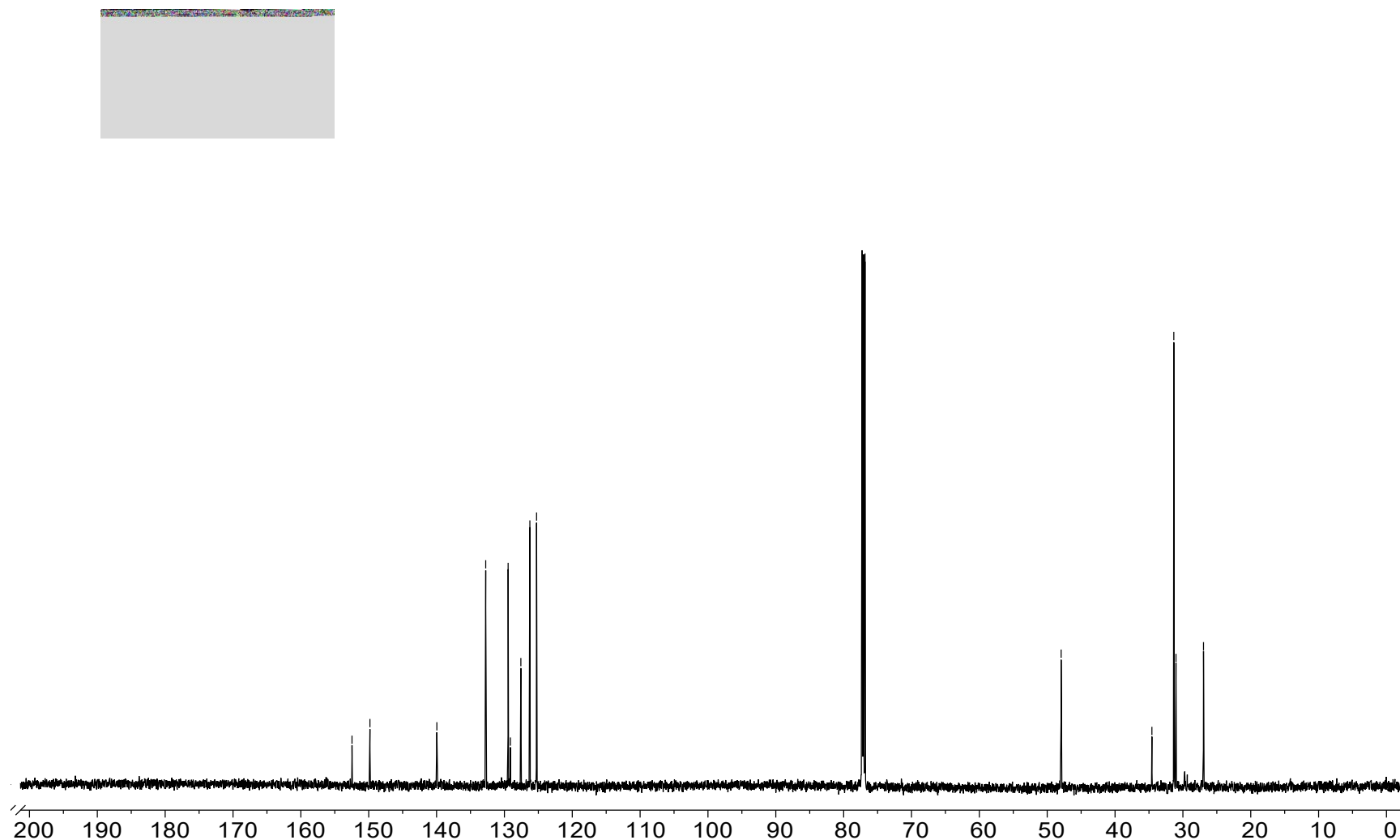
Nov23-2024, 42, fid
JohnB-bj-1123-Y-28

~ 152.46
~ 149.82
— 139.96
/ 132.75
/ 129.45
/ 129.12
~ 127.58
/ 126.25
/ 125.26

— 76.85

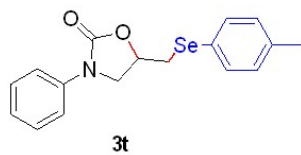
— 47.95

/ 34.57
/ 31.34
/ 31.02
~ 26.96

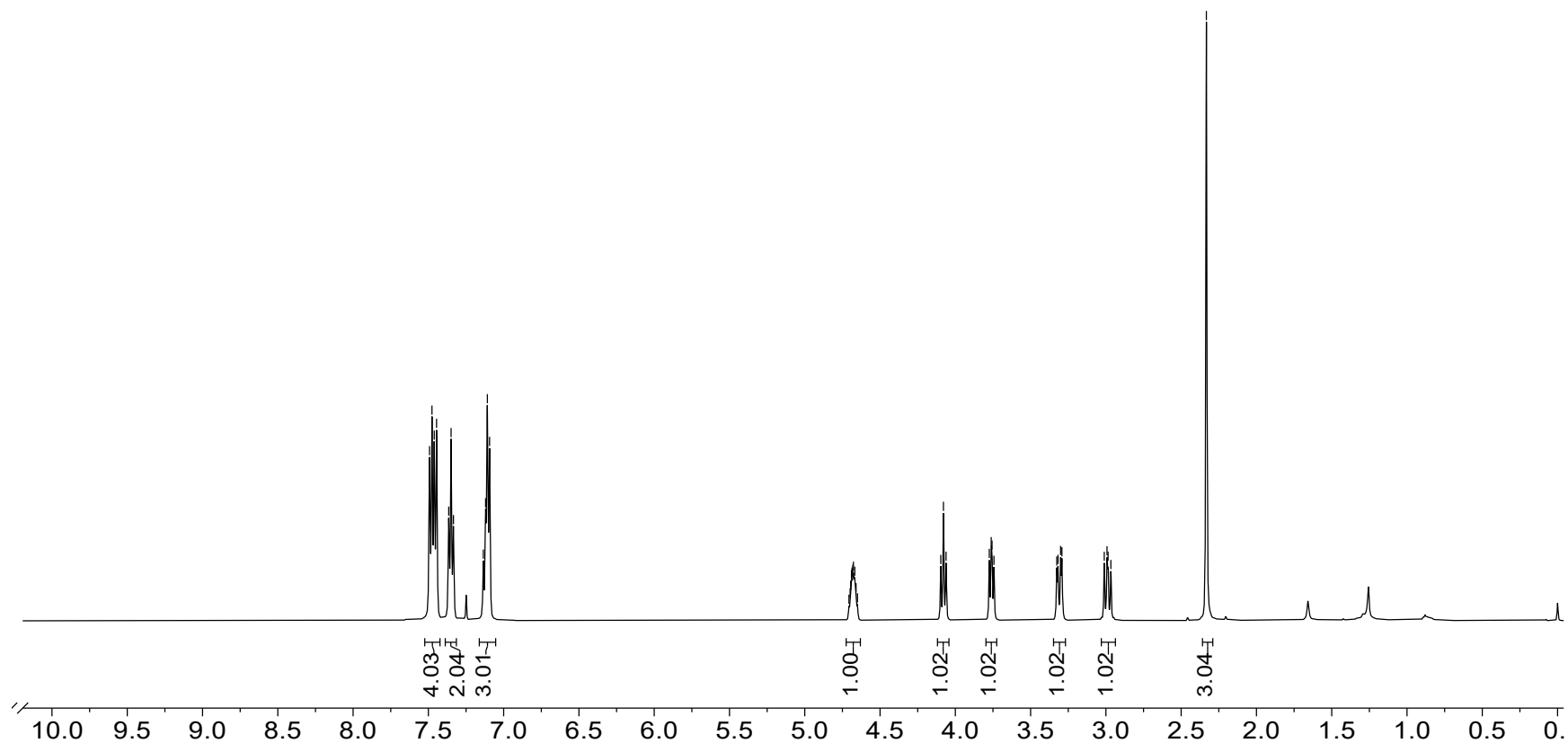


^1H NMR (500 MHz, CDCl_3) spectrum of **3t**.

17. 1. fid

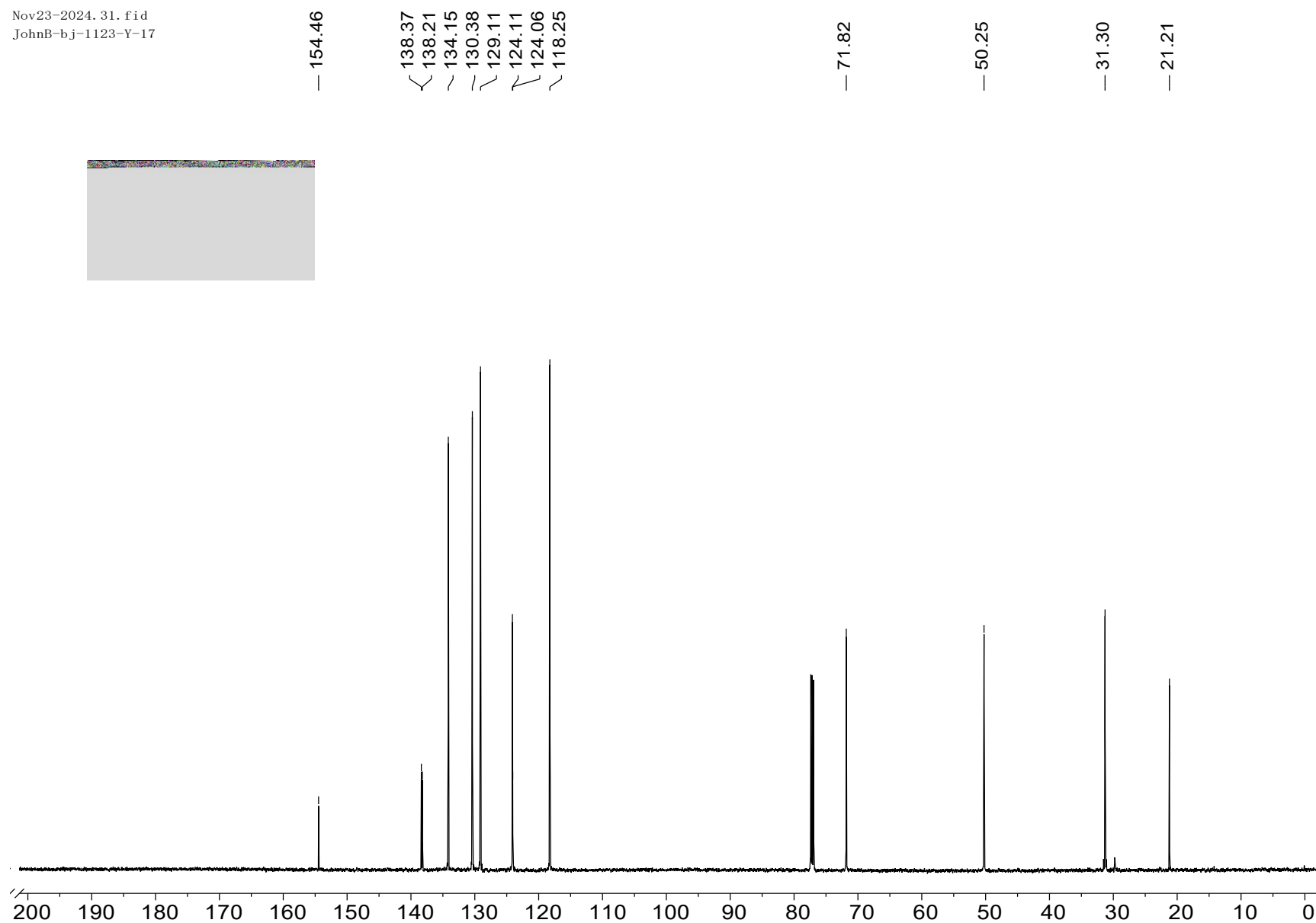


7.49 7.48 7.46 7.45 7.36 7.35 7.33 7.13 7.12 7.11 7.09
4.71 4.70 4.69 4.69 4.68 4.68 4.67 4.66 4.65 4.10 4.08 4.06 3.77 3.76 3.76 3.74 3.33 3.32 3.30 3.29 3.01 2.99 2.99 2.97 2.33



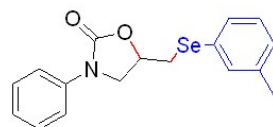
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3t**.

Nov23-2024, 31, fid
JohnB-bj-1123-Y-17

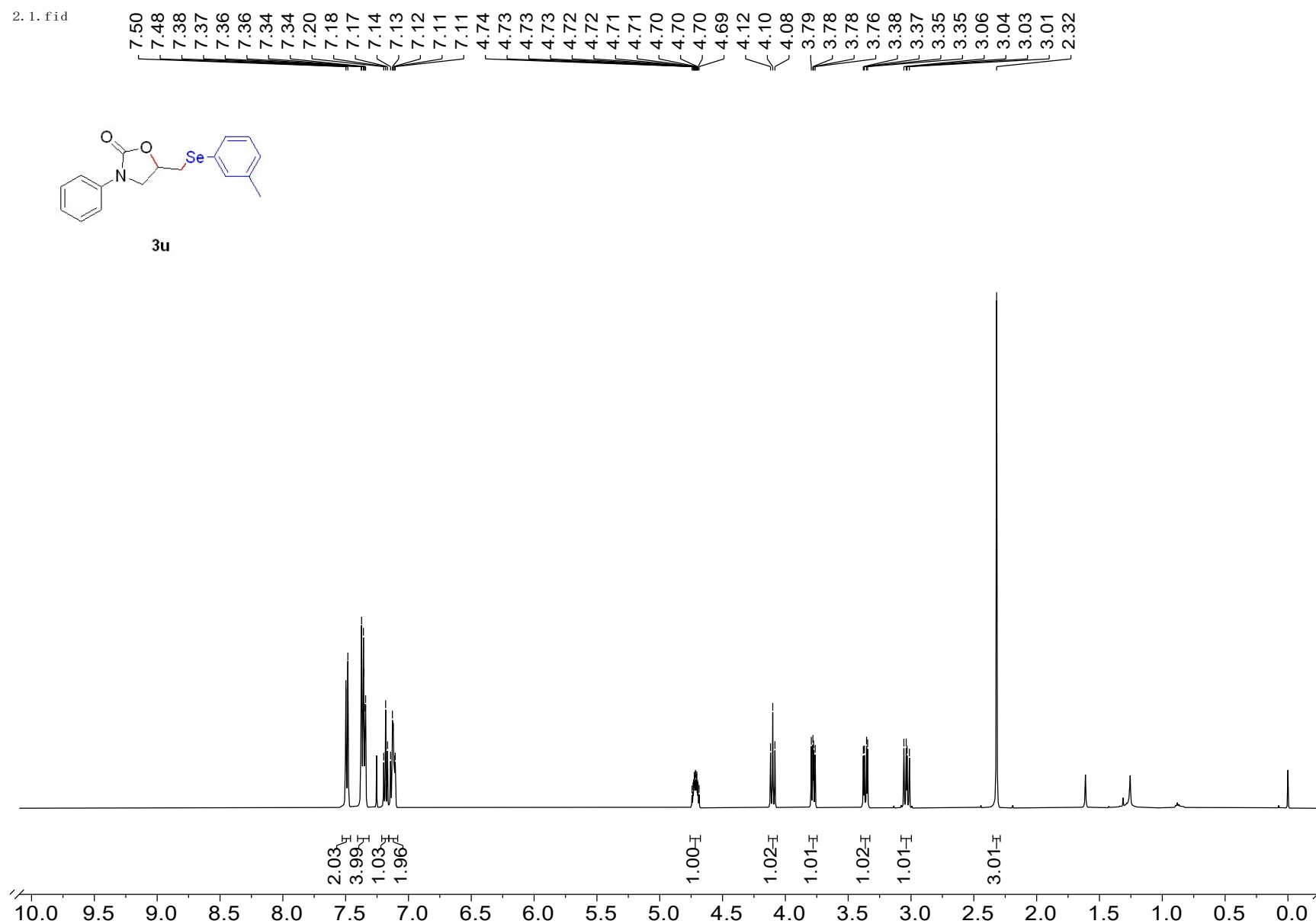


^1H NMR (500 MHz, CDCl_3) spectrum of **3u**.

2.1. fid

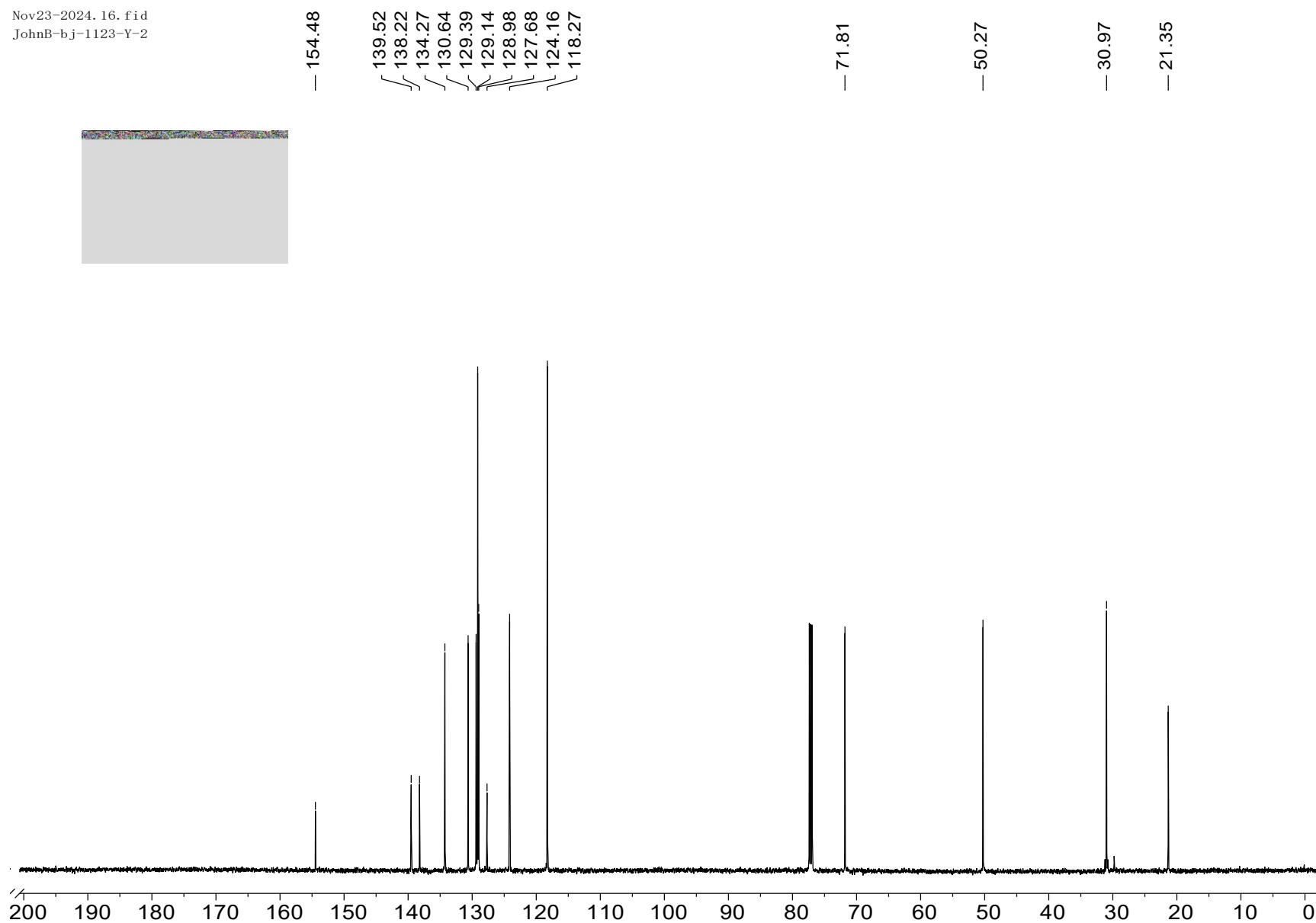


3u



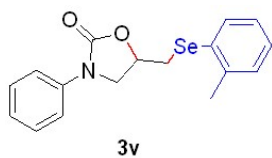
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3u**.

Nov23-2024, 16. fid
JohnB-bj-1123-Y-2

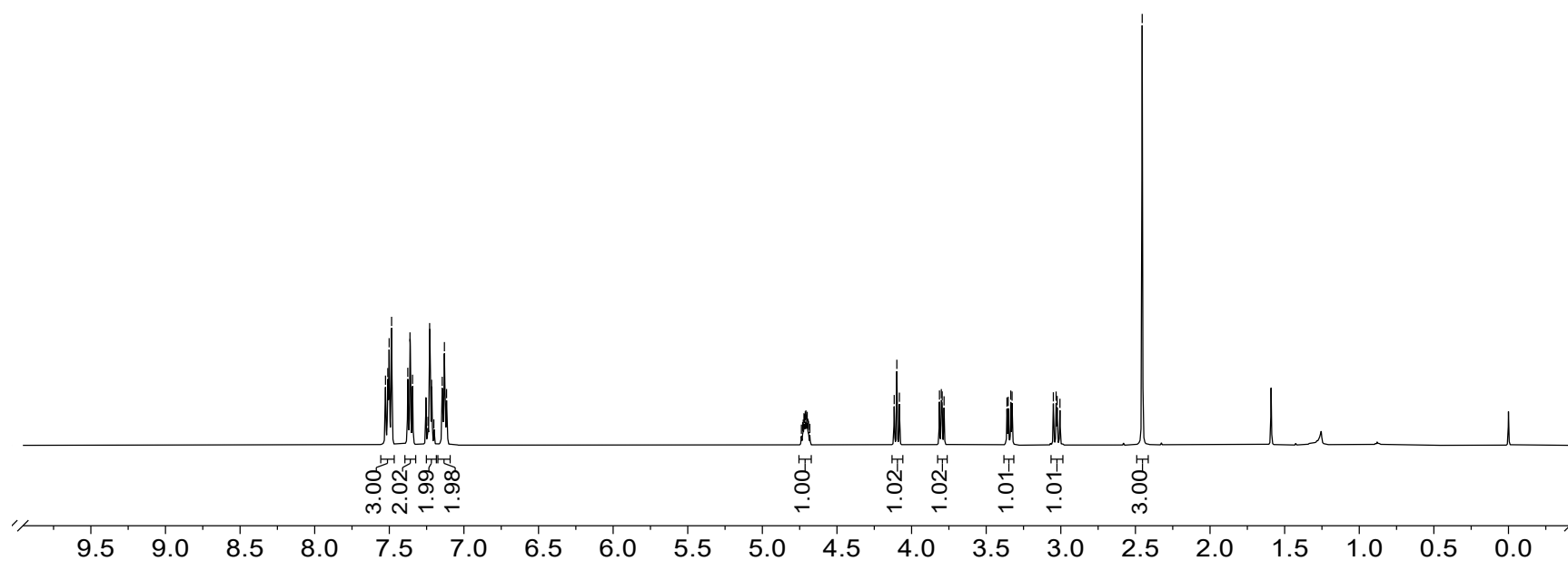


^1H NMR (500 MHz, CDCl_3) spectrum of **3v**.

5.1.fid

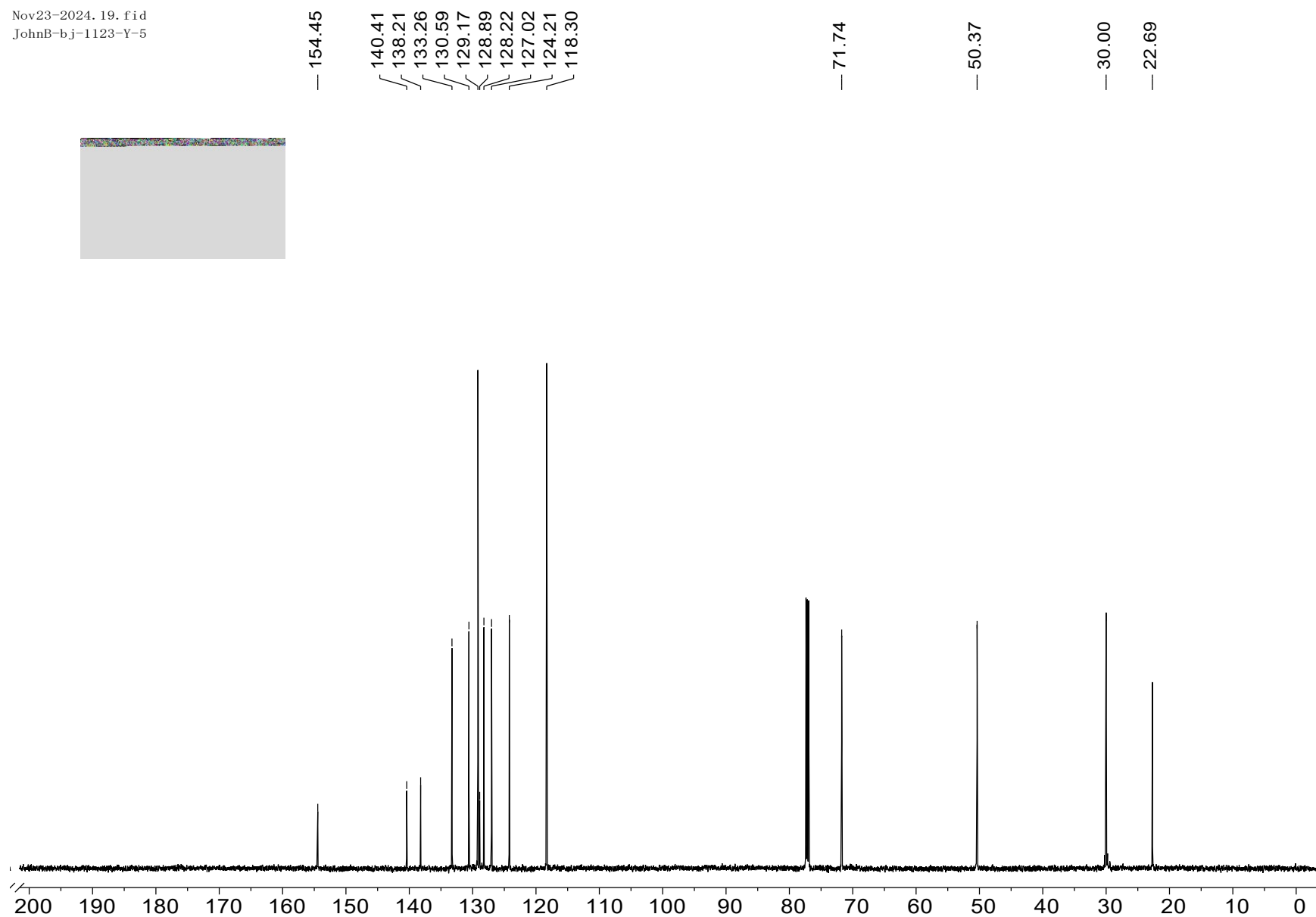


7.53
7.51
7.50
7.48
7.38
7.36
7.34
7.24
7.23
7.22
7.20
7.15
7.13
7.12
4.74
4.73
4.73
4.72
4.72
4.71
4.71
4.70
4.70
4.69
4.69
4.68
4.12
4.10
4.08
3.81
3.80
3.80
3.78
3.36
3.35
3.34
3.33
3.05
3.03
3.02
3.01
2.45



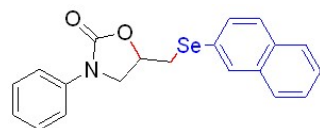
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3v**.

Nov23-2024, 19, fid
JohnB-bj-1123-Y-5

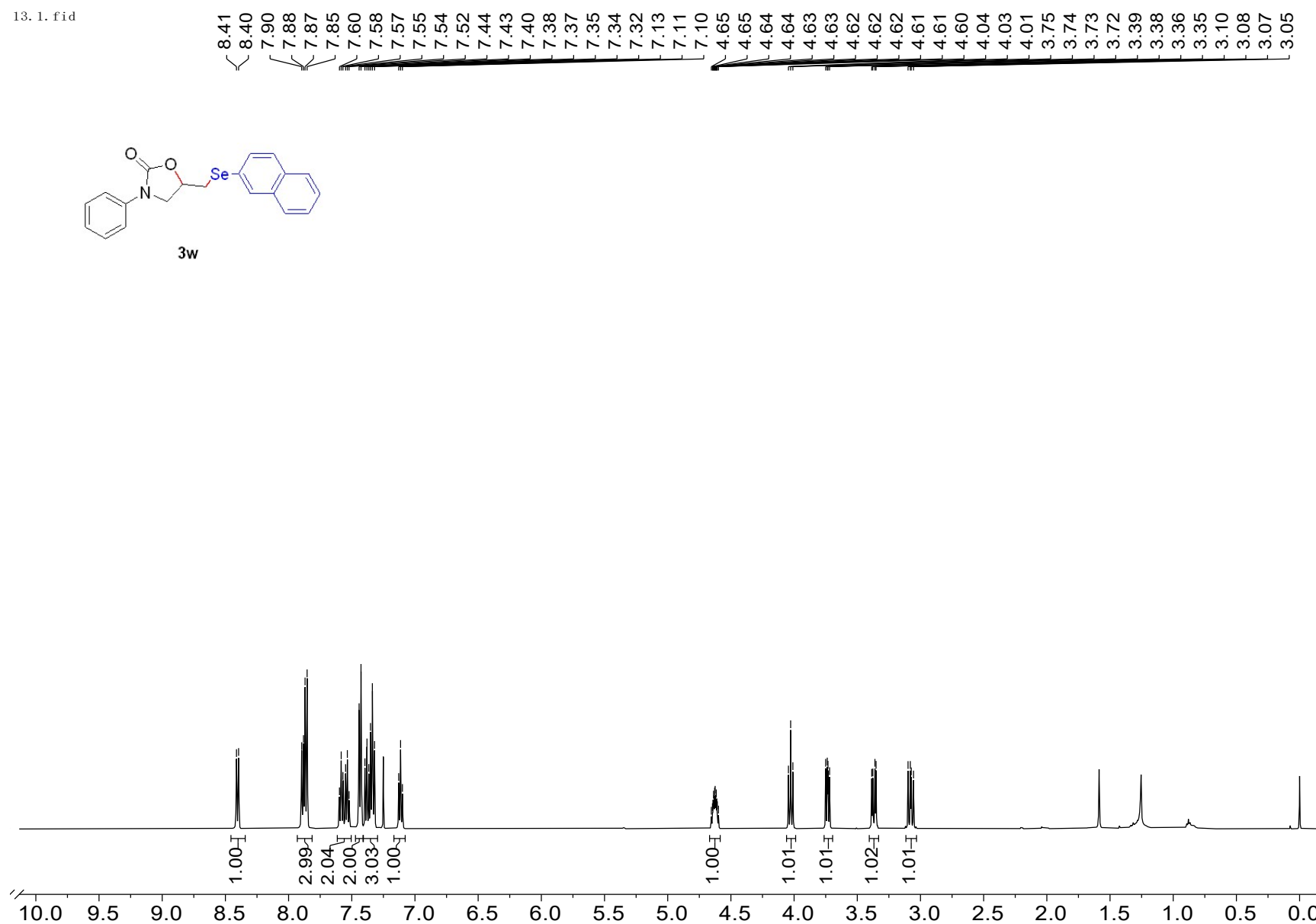


^1H NMR (500 MHz, CDCl_3) spectrum of **3w**.

13.1.1.fid

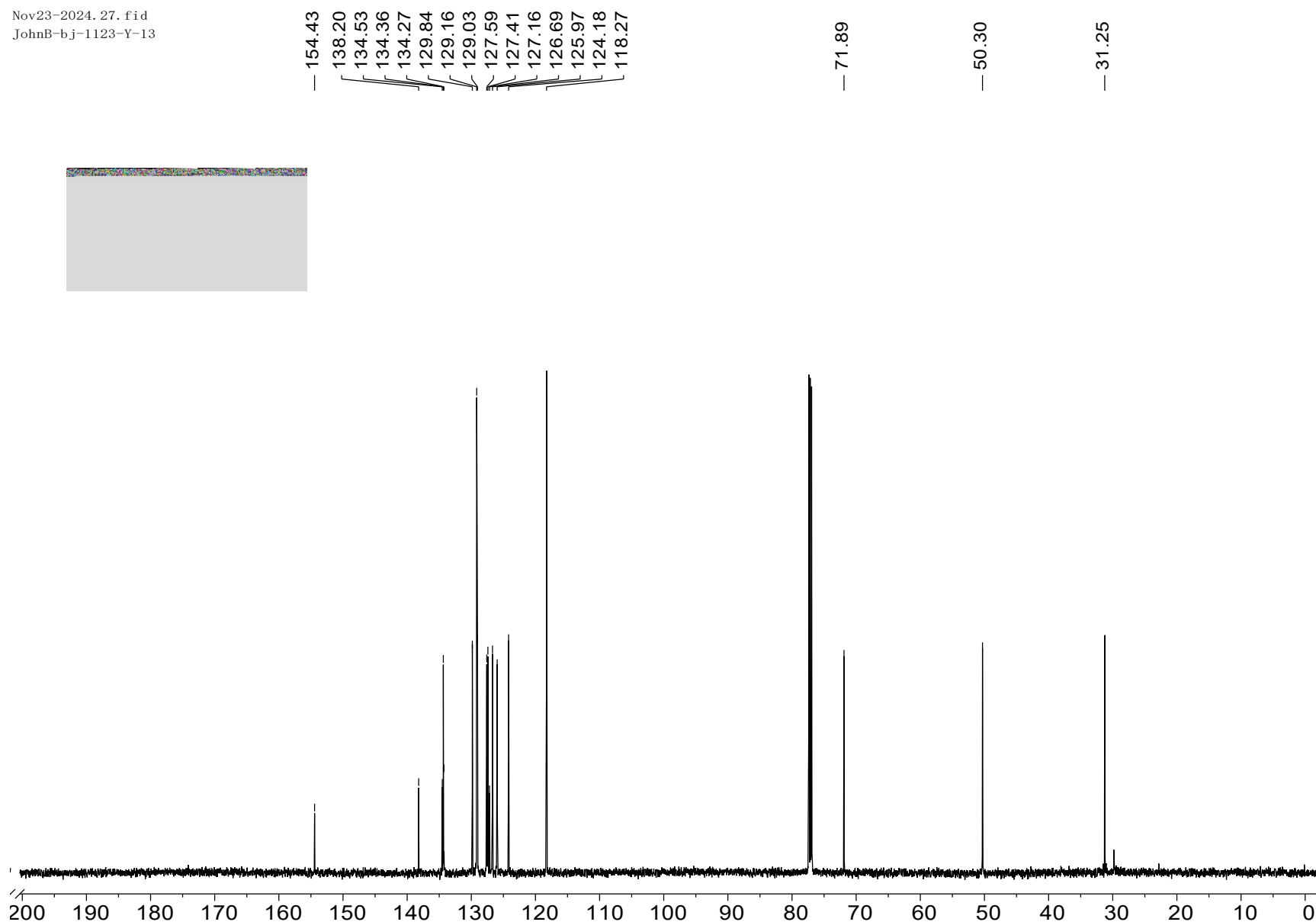


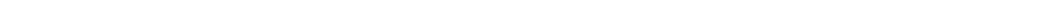
3w

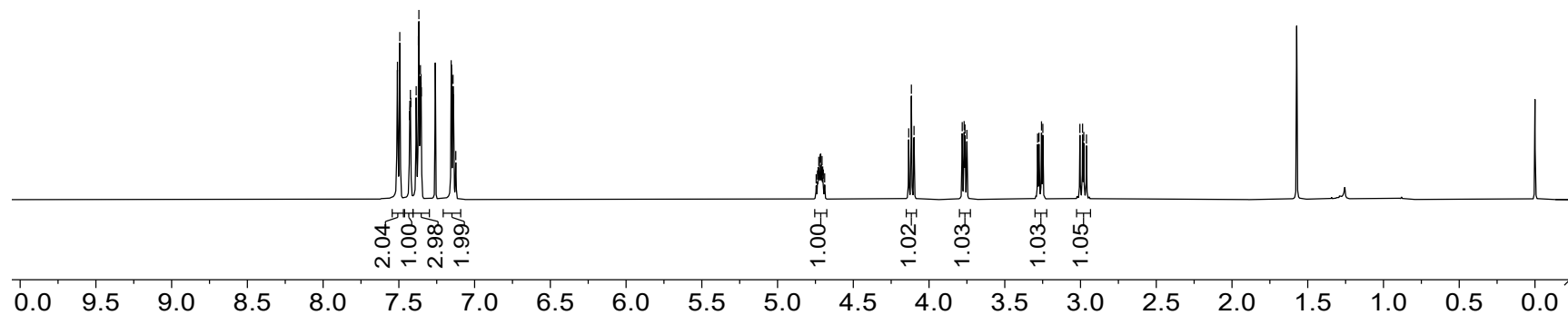
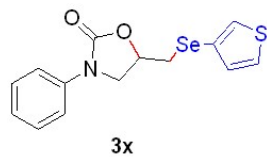


^{13}C NMR (151 MHz, CDCl_3) spectrum of **3w**.

Nov23-2024. 27. fid
JohnB-bj-1123-Y-13

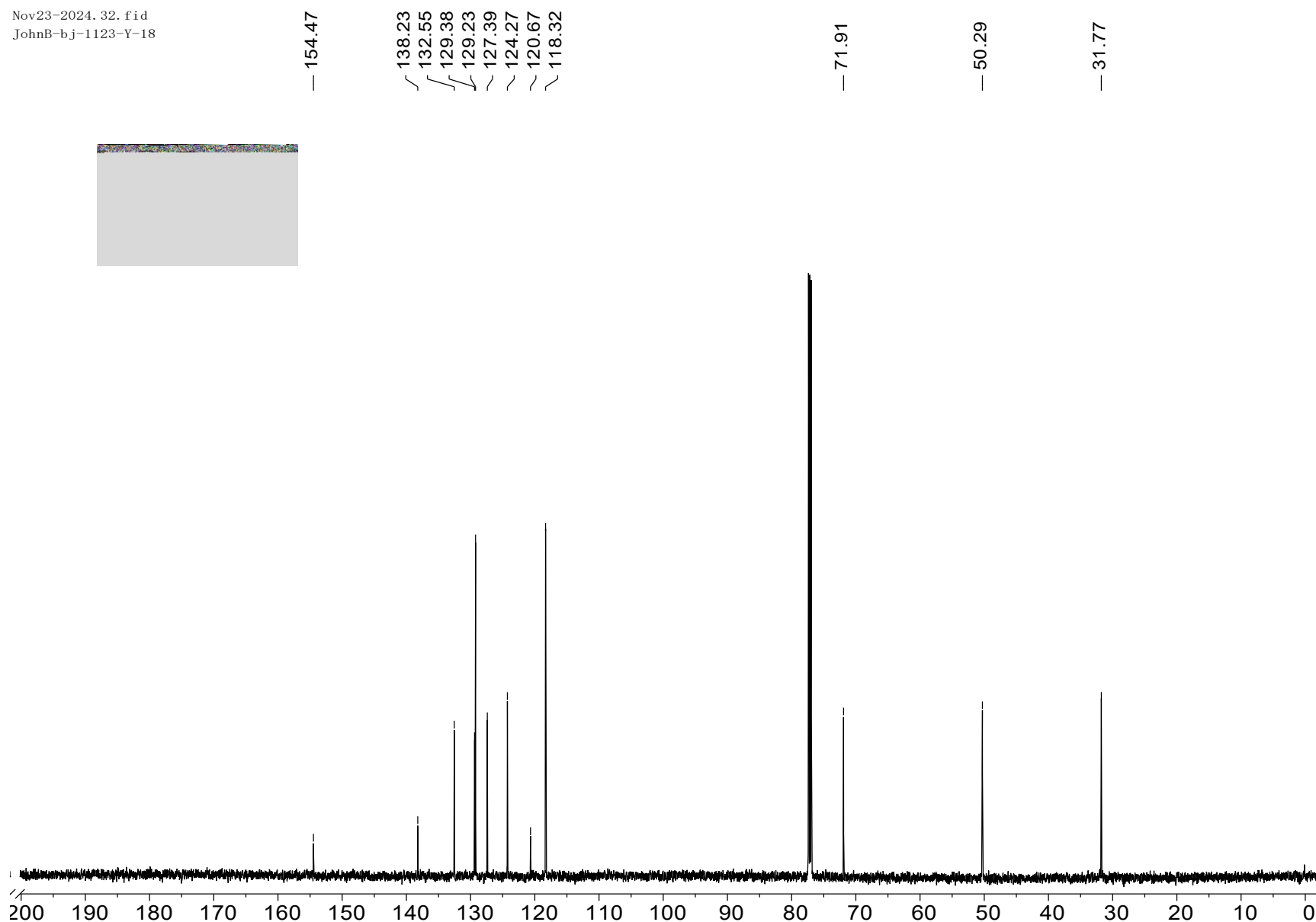


18.1. 



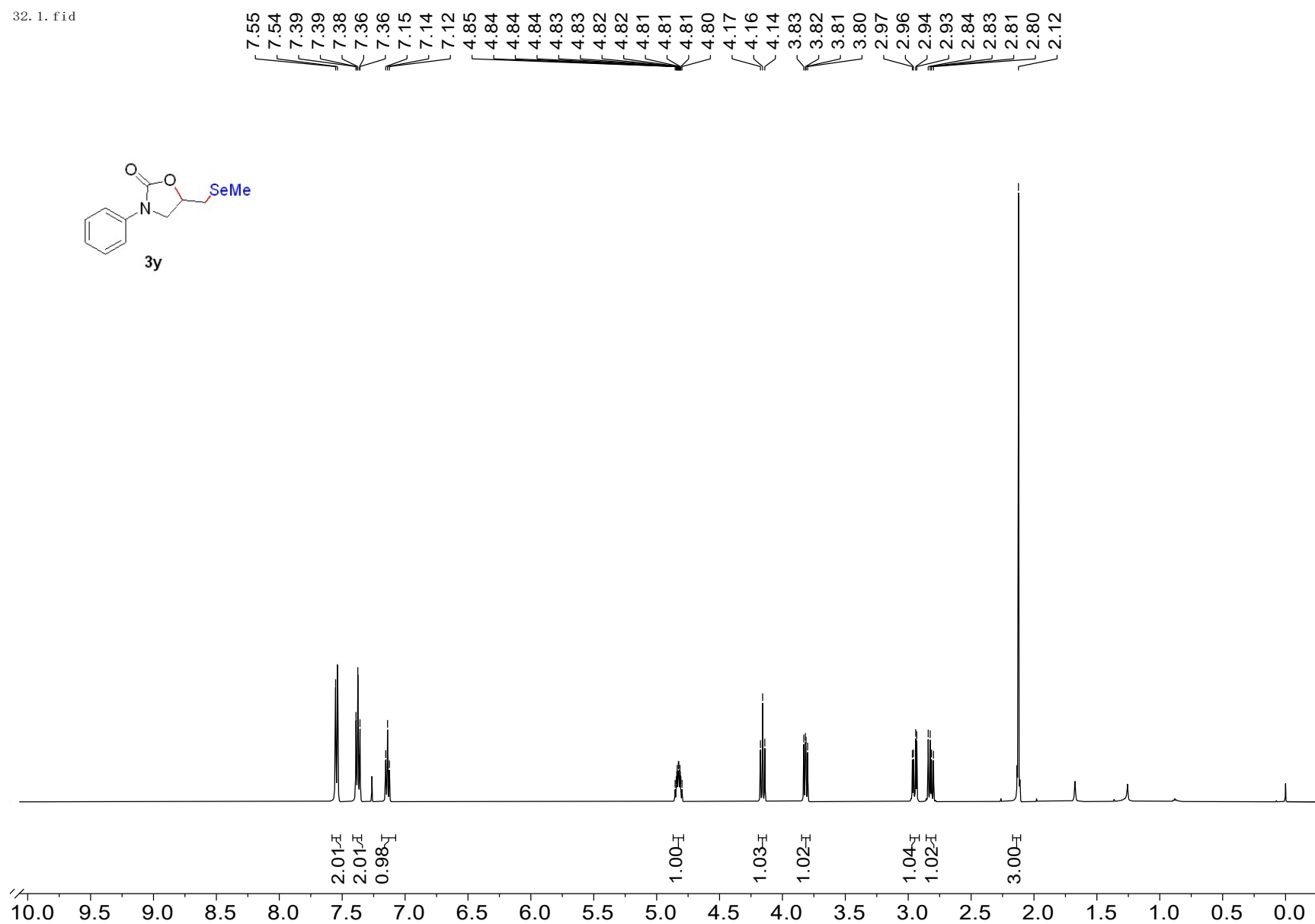
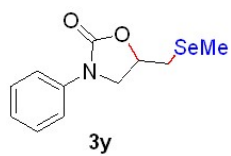
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3x**.

Nov23-2024, 32, fid
JohnB-bj-1123-Y-18



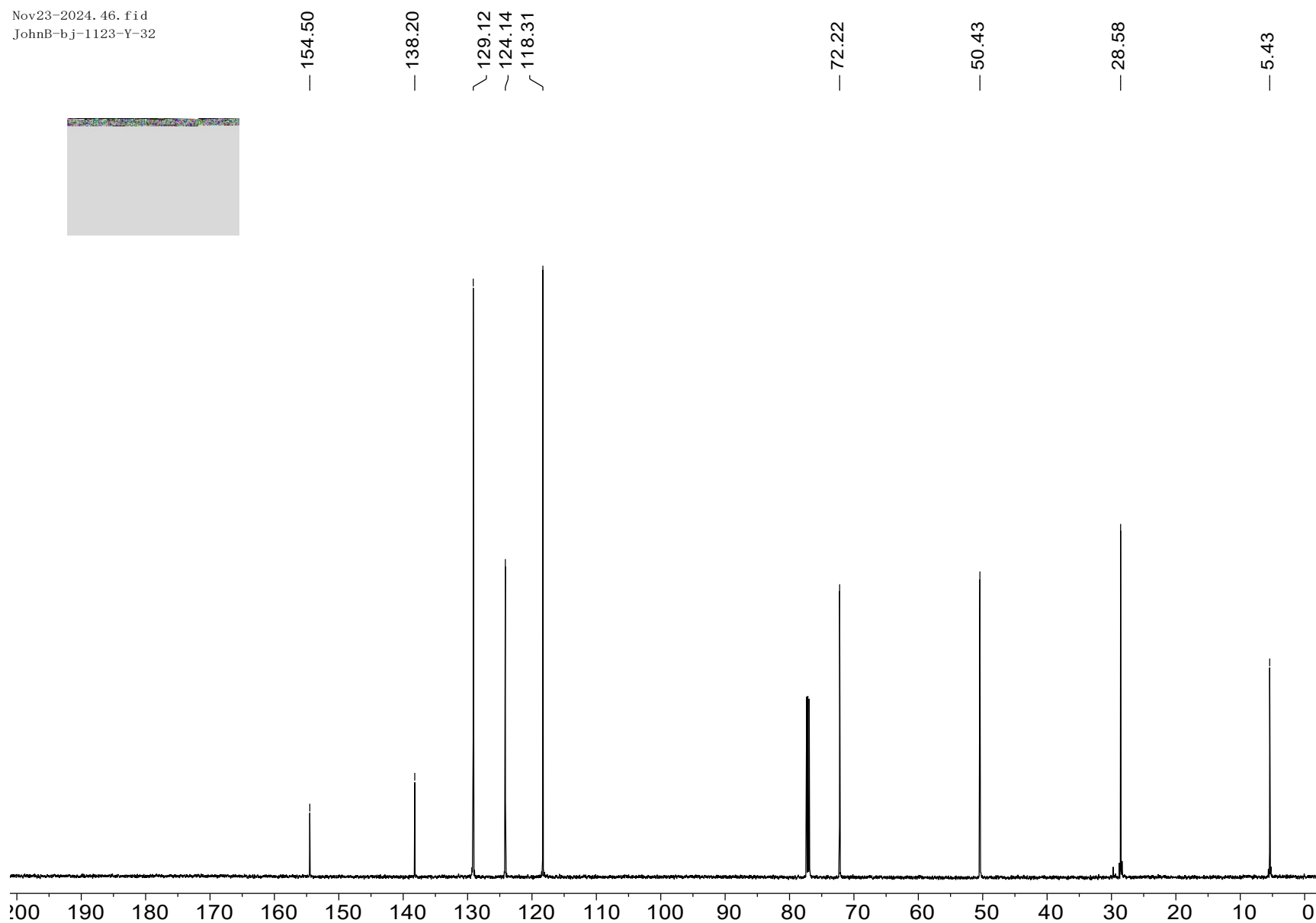
^1H NMR (500 MHz, CDCl_3) spectrum of **3y**.

32.1.fid



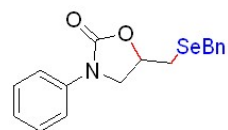
^{13}C NMR (151 MHz, CDCl_3) spectrum of **3y**.

Nov23-2024, 46, fid
JohnB-bj-1123-Y-32

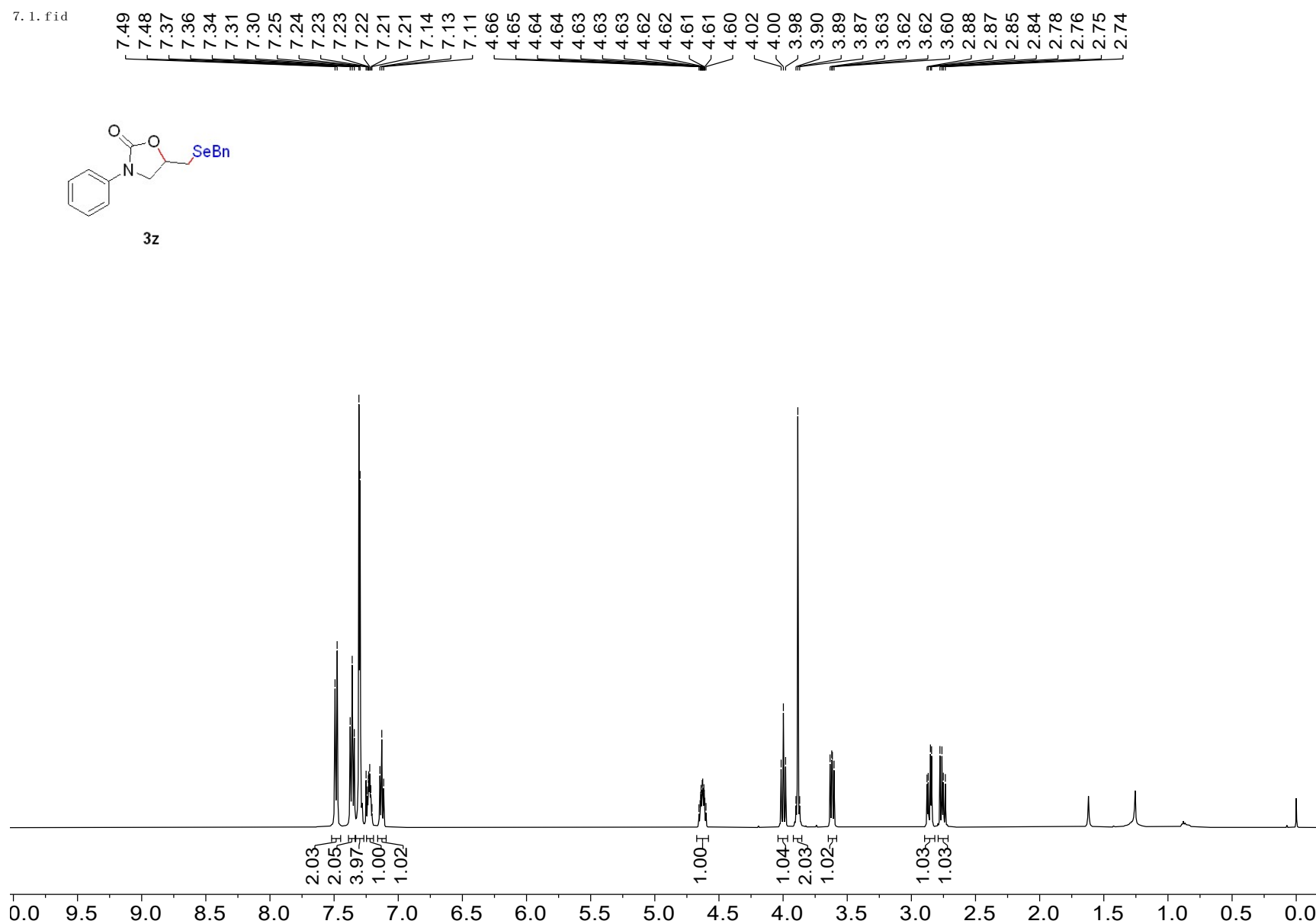


^1H NMR (500 MHz, CDCl_3) spectrum of **3z**.

7.1. fid

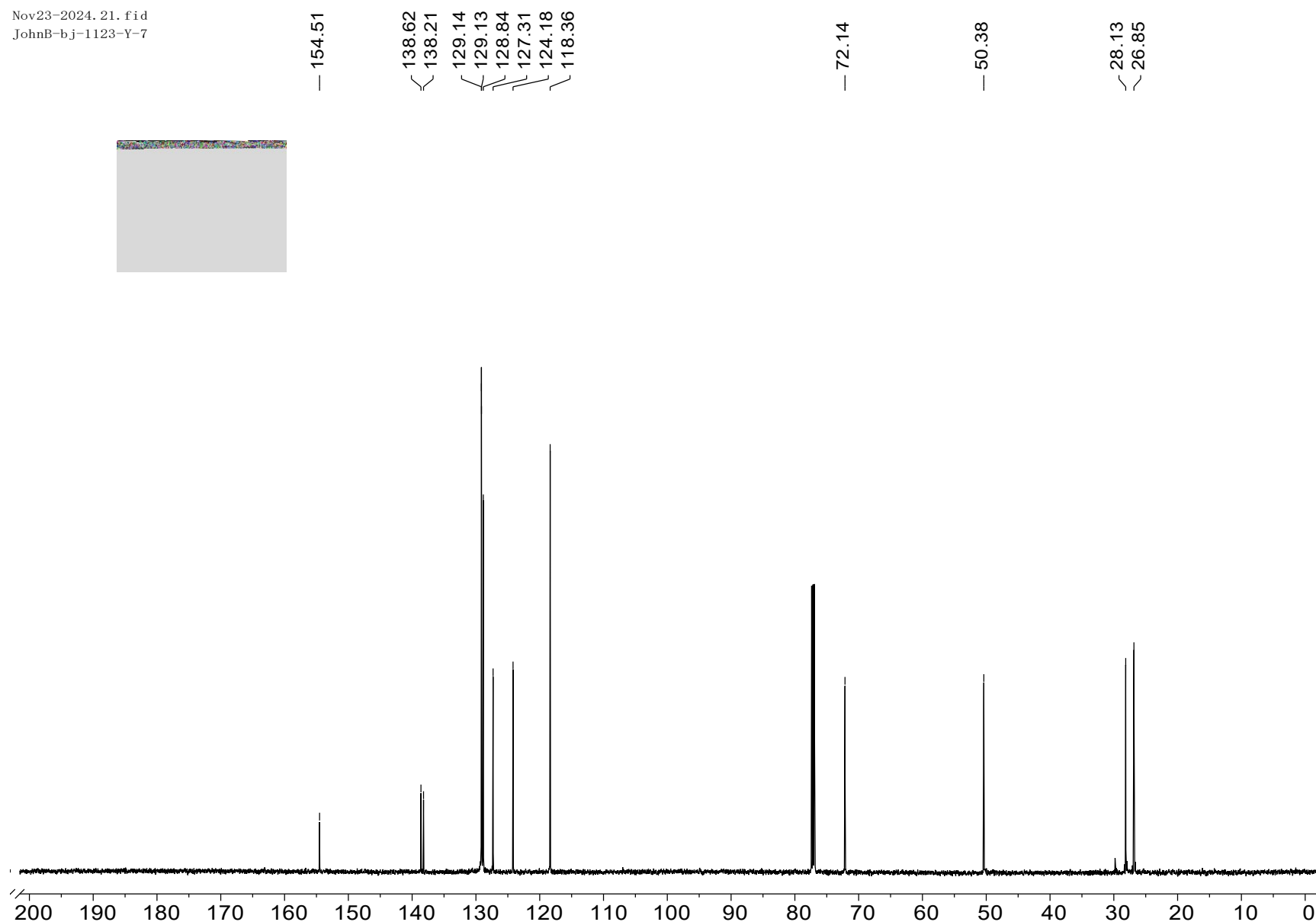


3z



^{13}C NMR (151 MHz, CDCl_3) spectrum of **3z**.

Nov23-2024, 21. fid
JohnB-bj-1123-Y-7



^1H NMR (500 MHz, CDCl_3) spectrum of **5a**.

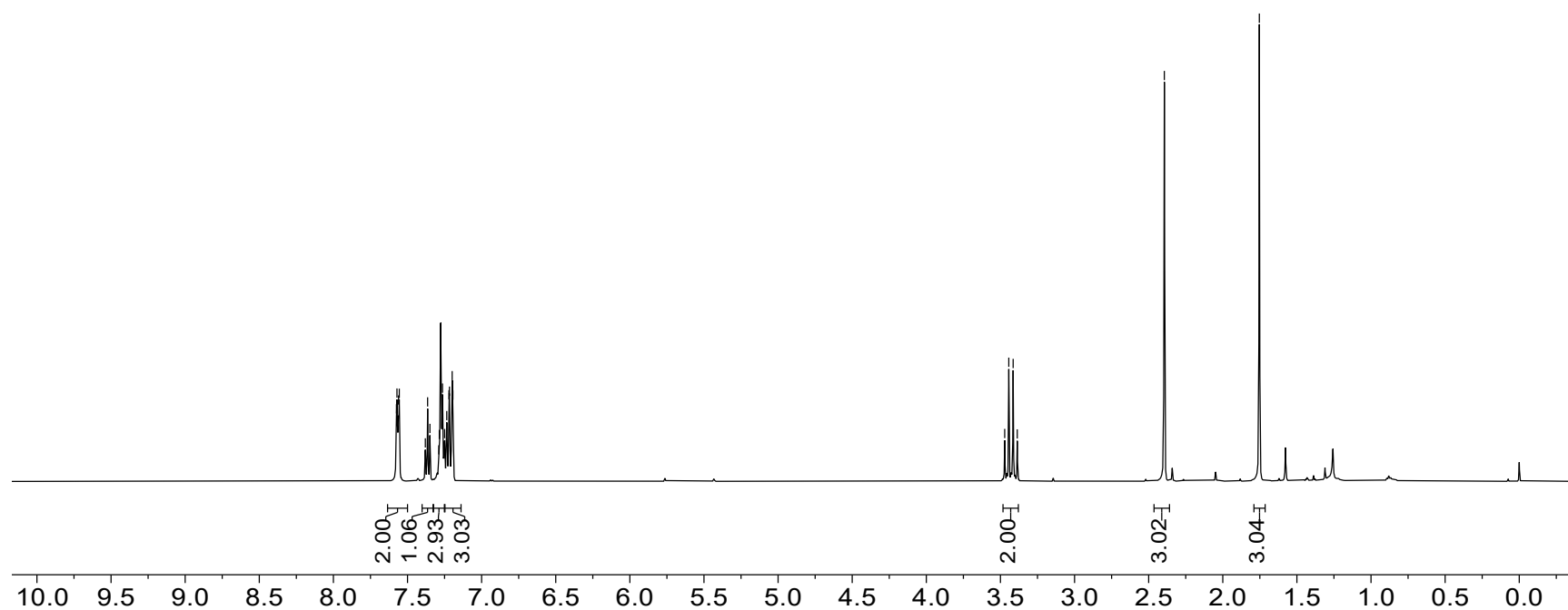
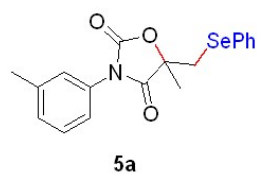
16. 1. fid

7.57
7.57
7.56
7.56
7.38
7.36
7.35
7.29
7.29
7.28
7.28
7.26
7.25
7.25
7.25
7.24
7.22
7.22
7.20
7.19

3.47
3.44
3.41
3.39

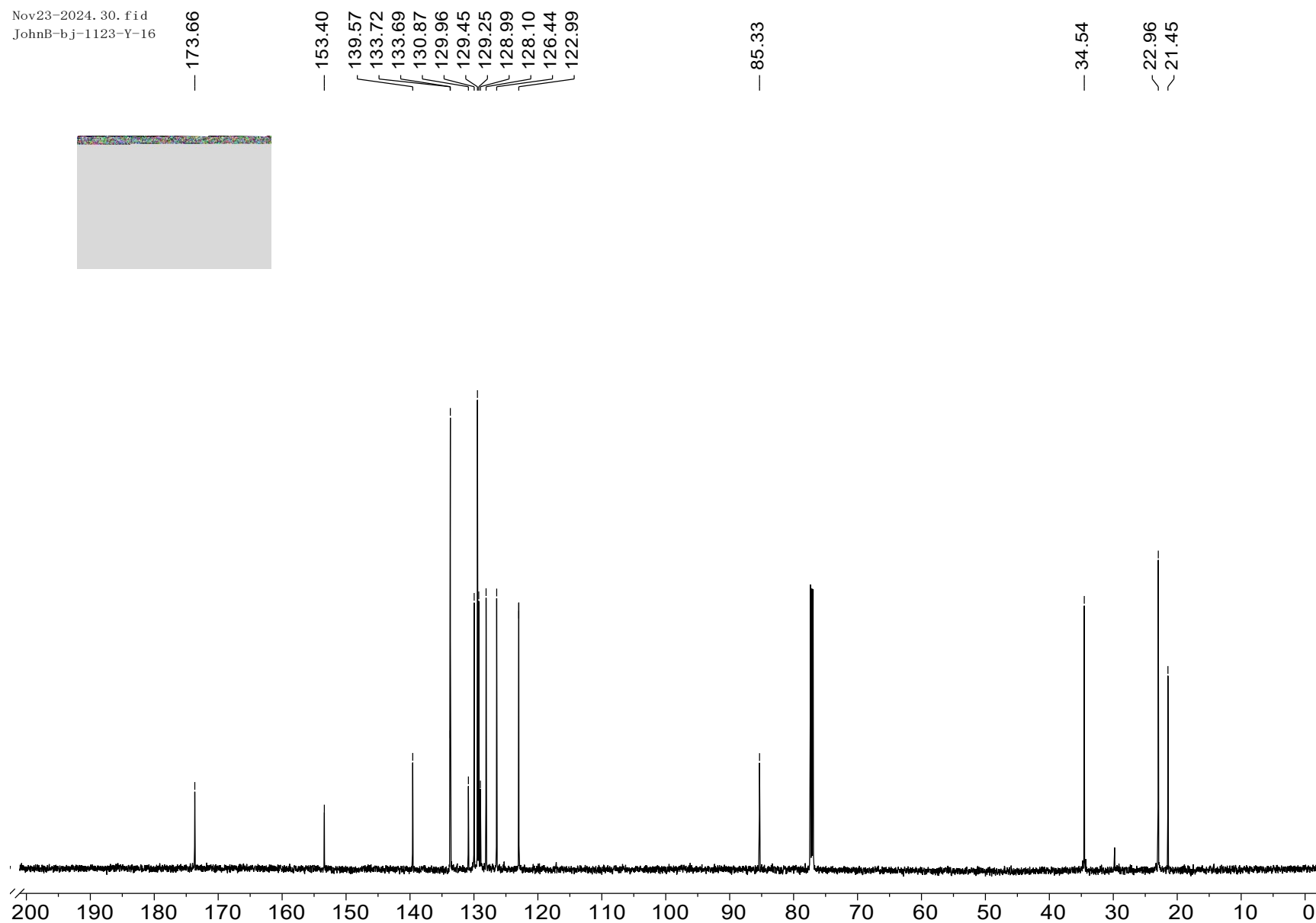
— 2.39

— 1.75



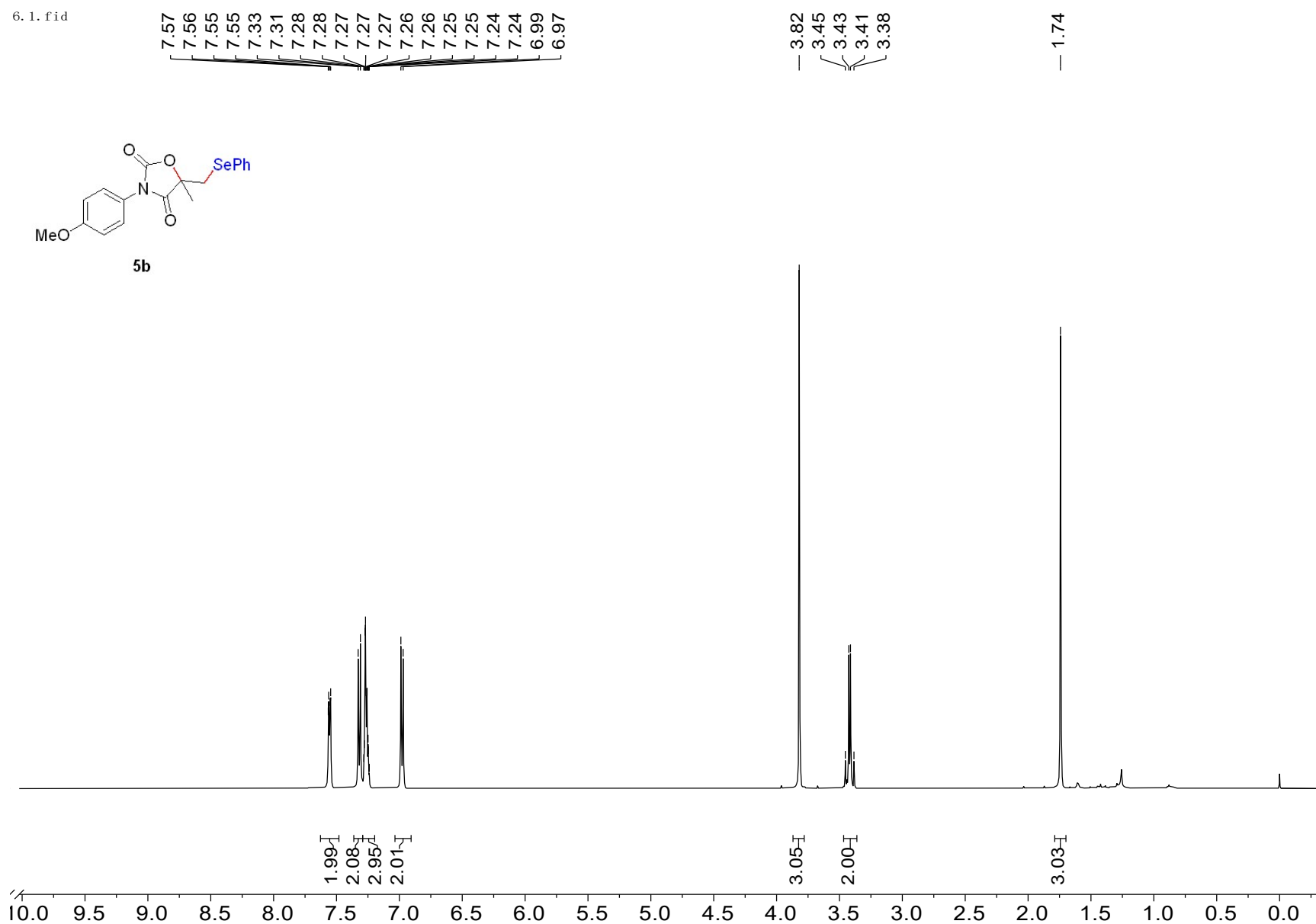
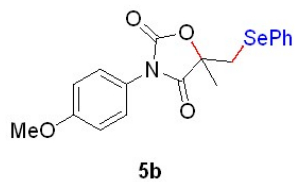
^{13}C NMR (151 MHz, CDCl_3) spectrum of **5a**.

Nov23-2024, 30. fid
JohnB-bj-1123-Y-16



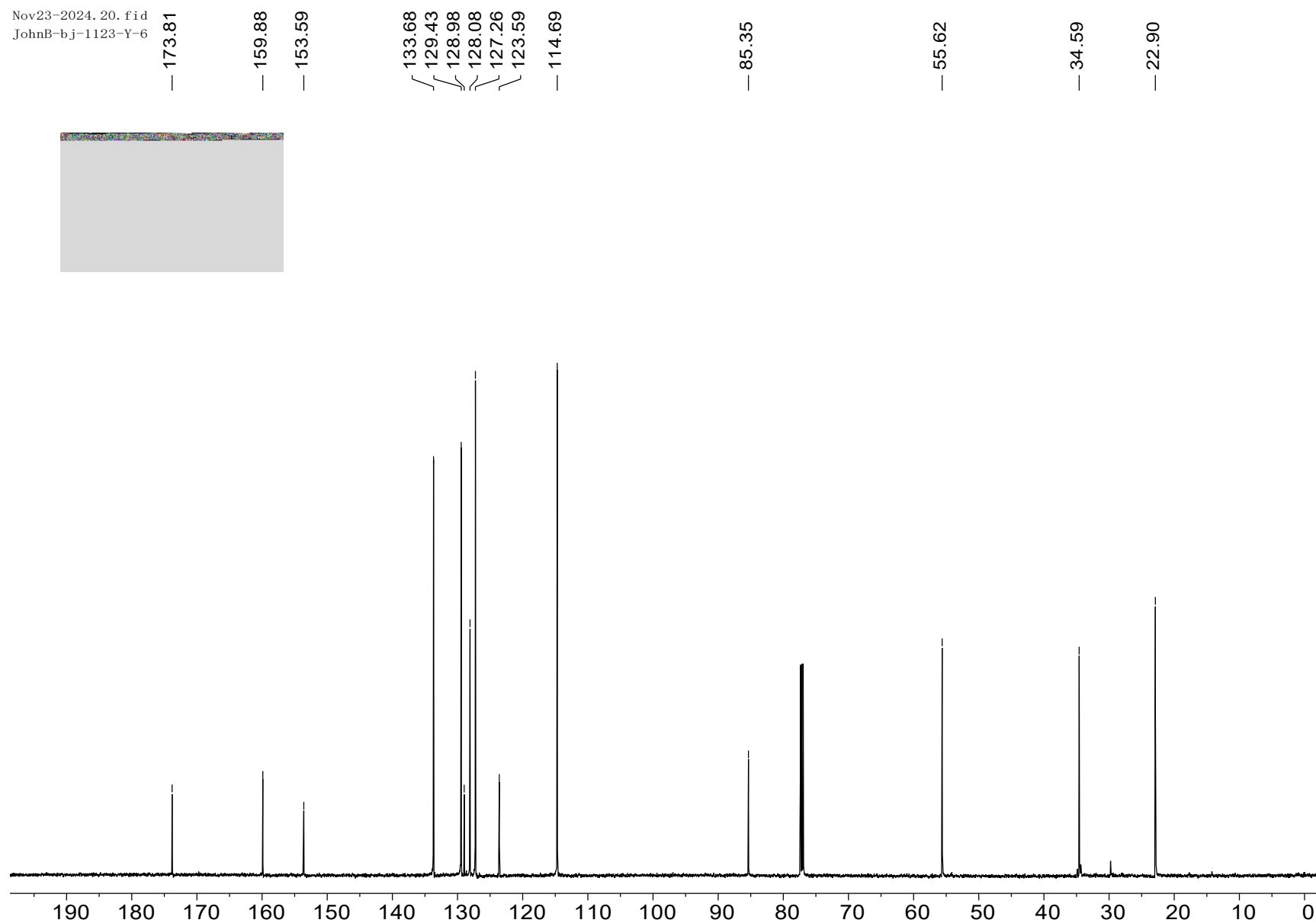
^1H NMR (500 MHz, CDCl_3) spectrum of **5b**.

6.1.fid



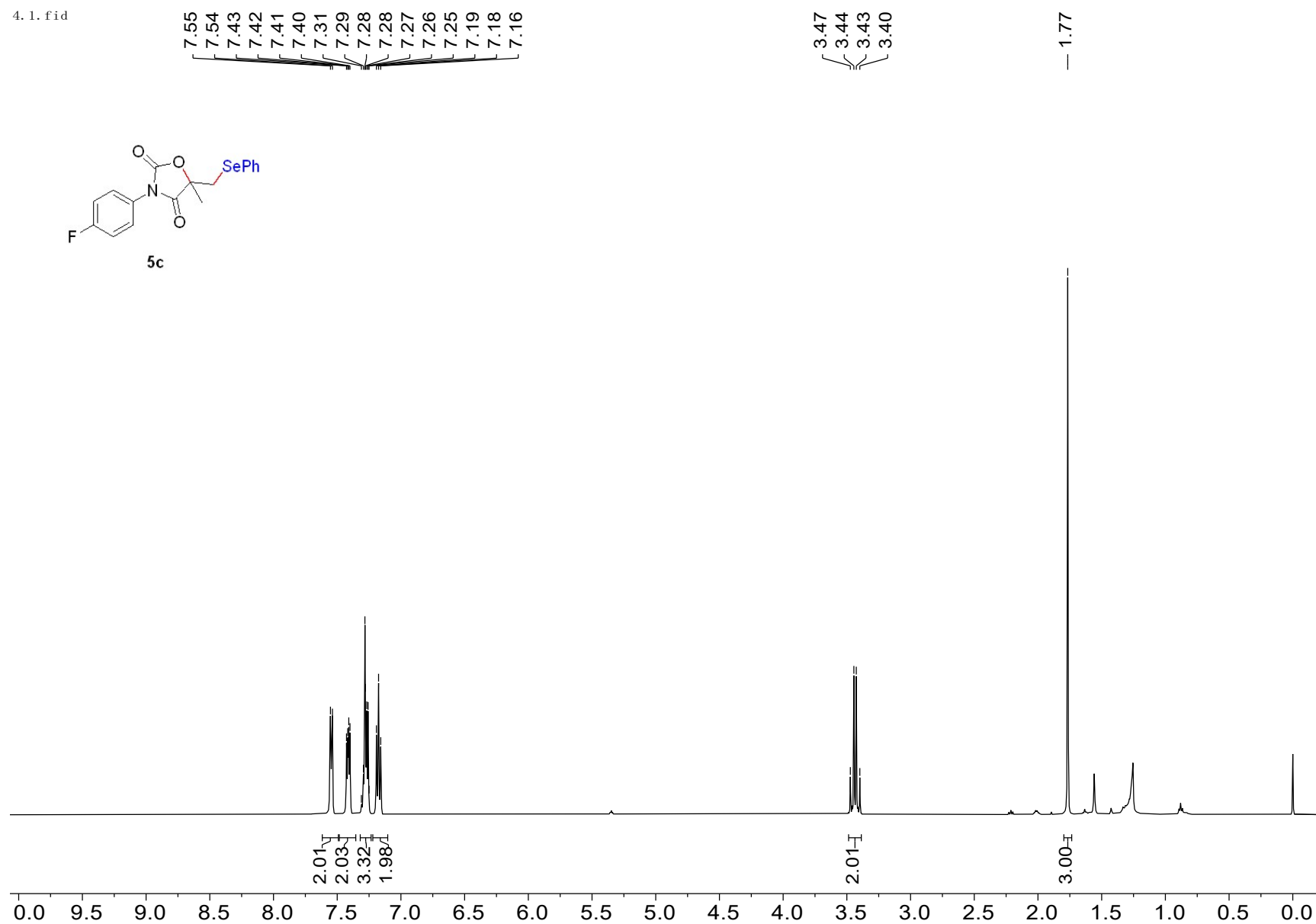
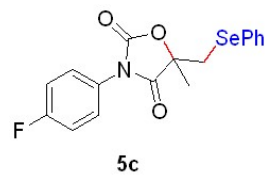
^{13}C NMR (151 MHz, CDCl_3) spectrum of **5b**.

Nov23-2024, 20. fid
JohnB-bj-1123-Y-6



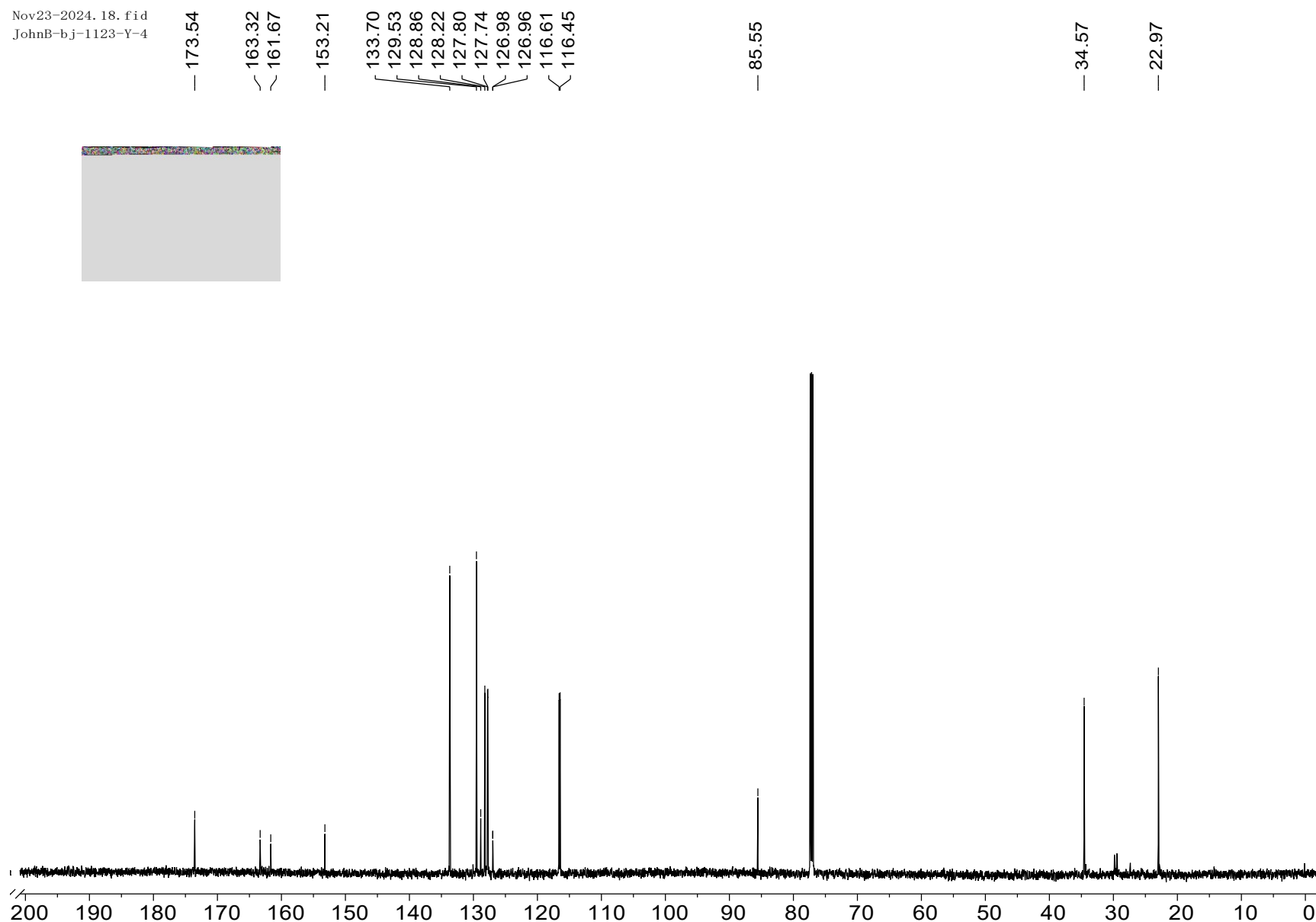
^1H NMR (500 MHz, CDCl_3) spectrum of **5c**.

4.1. fid



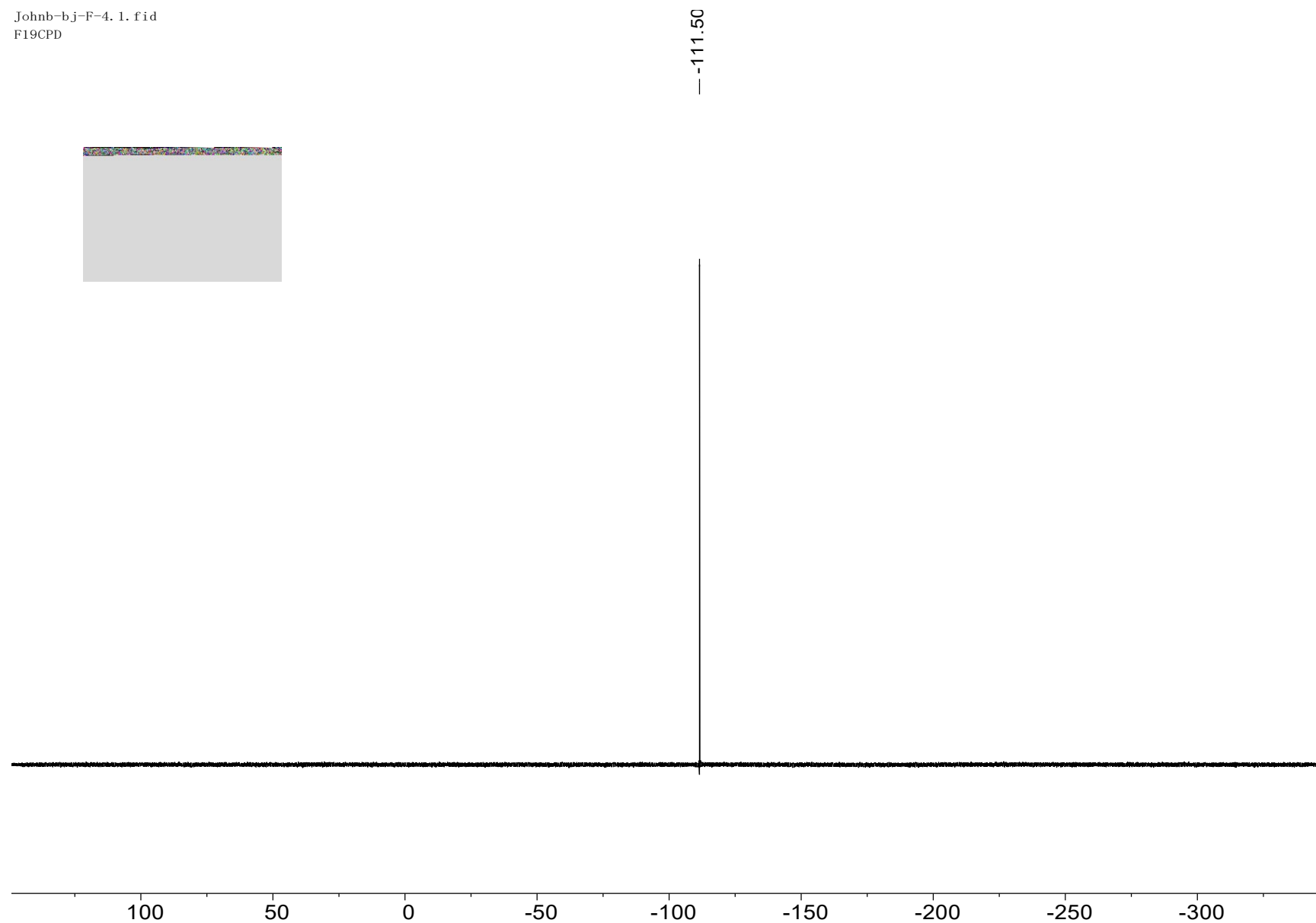
^{13}C NMR (151 MHz, CDCl_3) spectrum of **5c**.

Nov23-2024, 18. fid
JohnB-bj-1123-Y-4

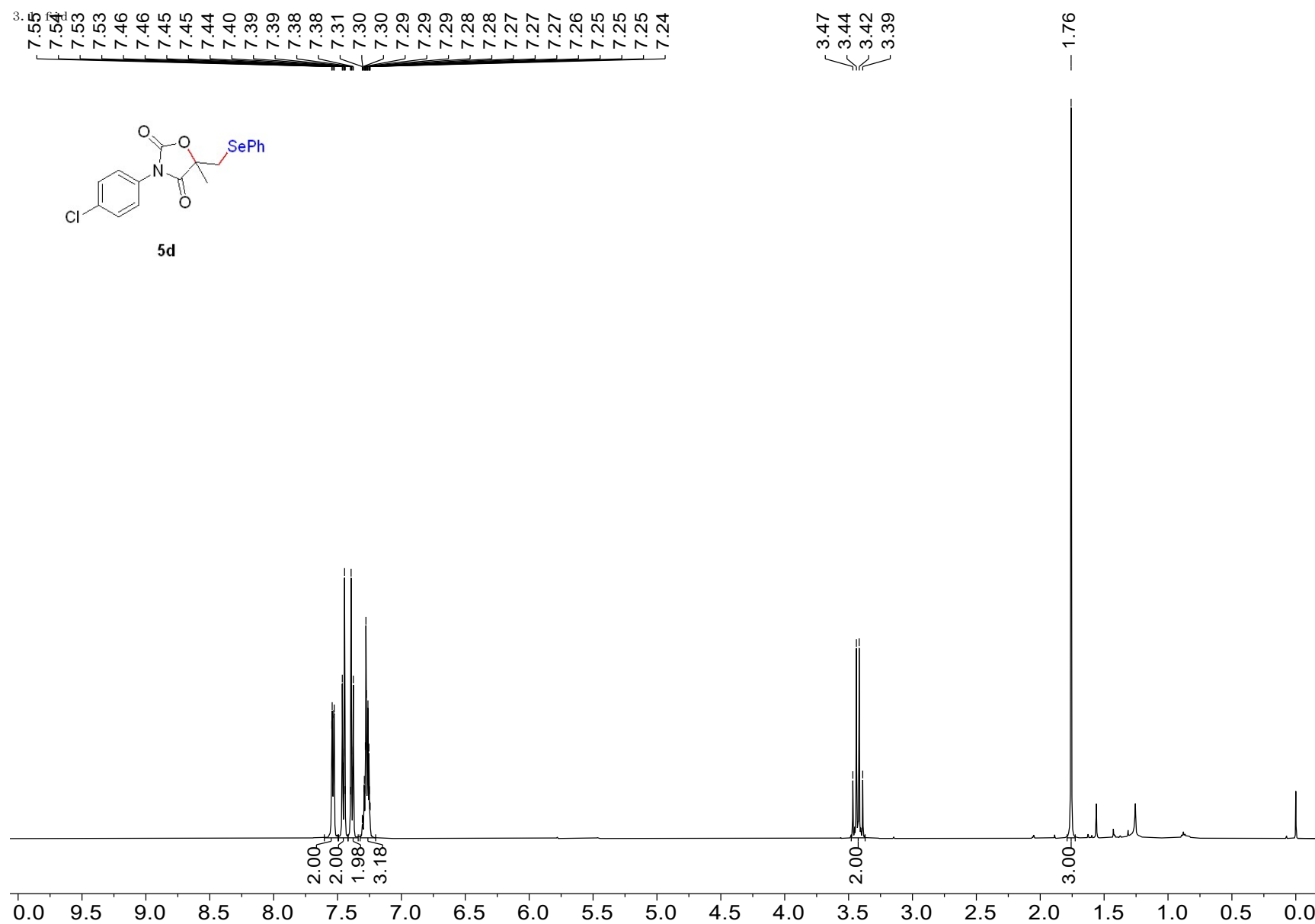


^{19}F NMR (376 MHz, CDCl_3) spectrum of **5c**.

Johnb-bj-F-4. 1. fid
F19CPD

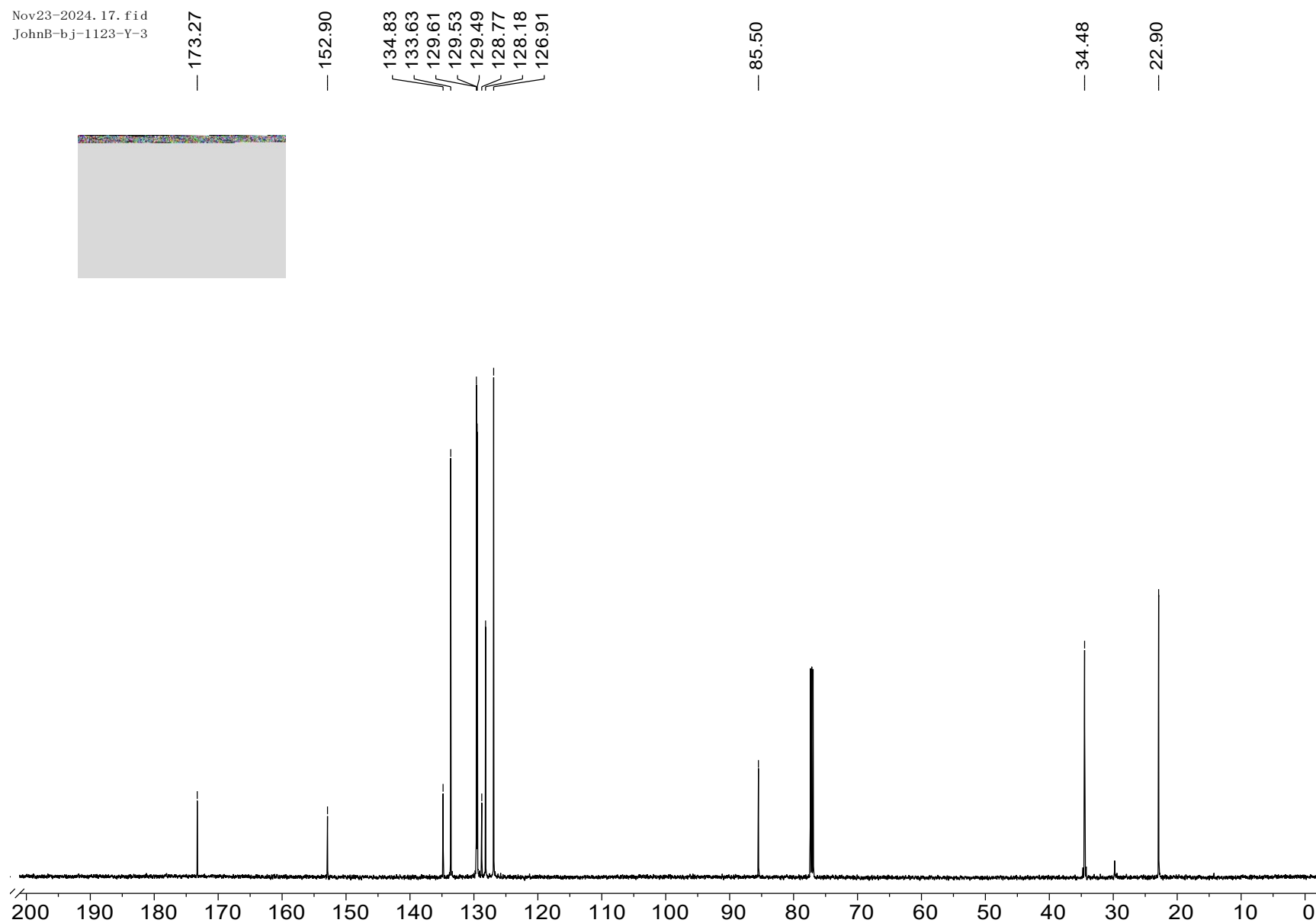


^1H NMR (500 MHz, CDCl_3) spectrum of **5d**.



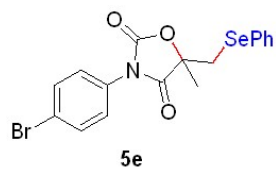
^{13}C NMR (151 MHz, CDCl_3) spectrum of **5d**.

Nov23-2024, 17. fid
JohnB-bj-1123-Y-3



^1H NMR (500 MHz, CDCl_3) spectrum of **5e**.

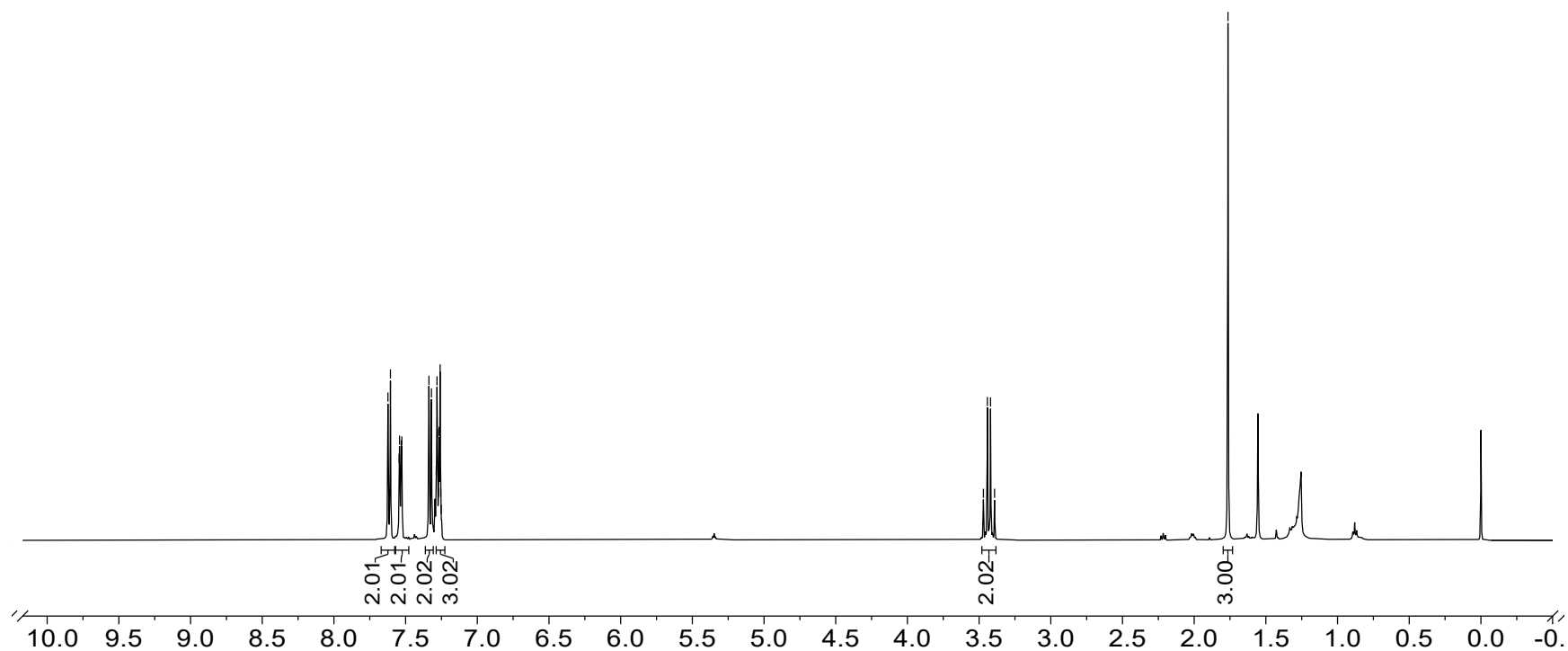
15.1.1.fid



7.62
7.61
7.55
7.54
7.53
7.53
7.34
7.32
7.32
7.29
7.28
7.28
7.27
7.27
7.26
7.26
7.25

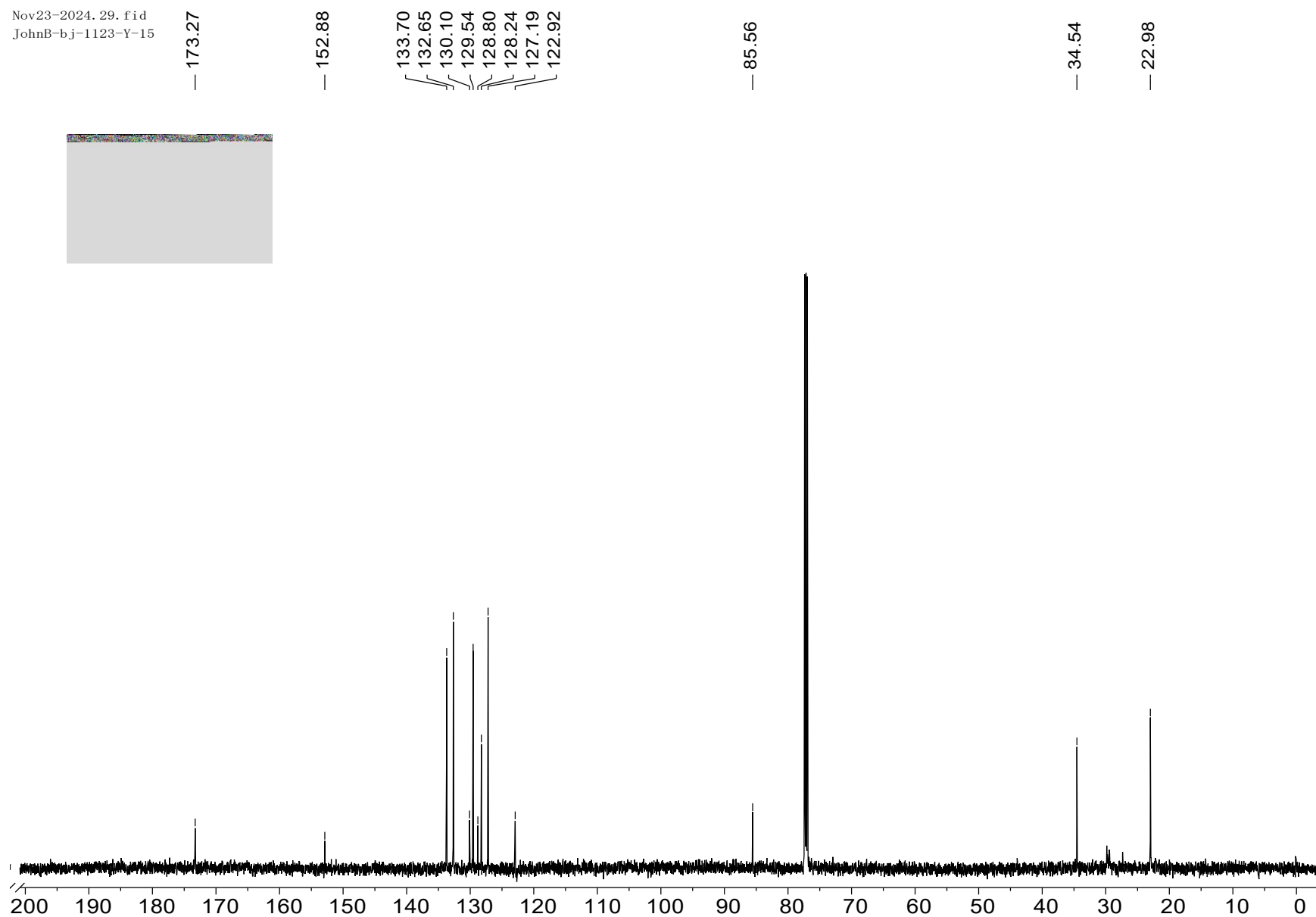
3.47
3.44
3.42
3.39

— 1.76



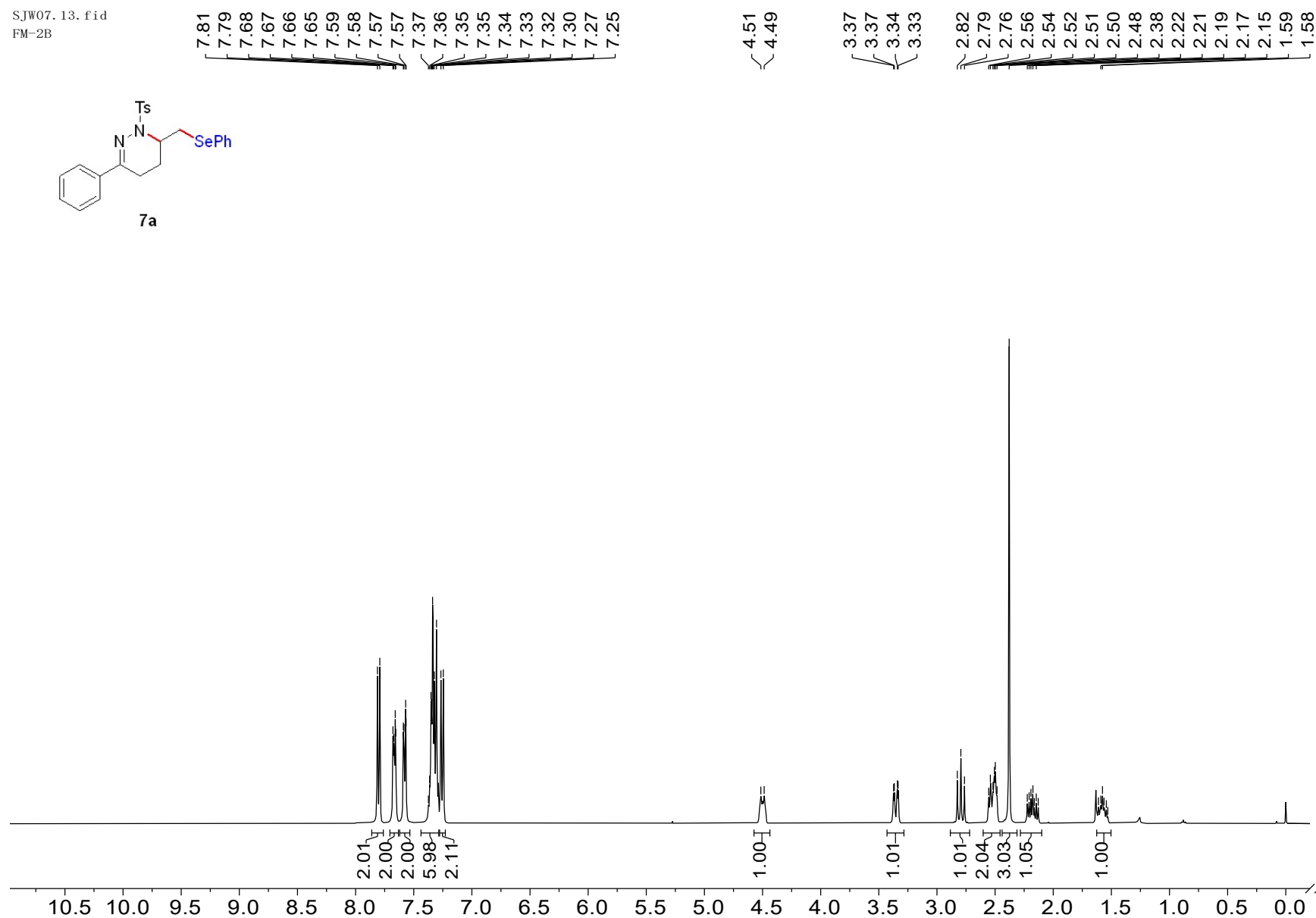
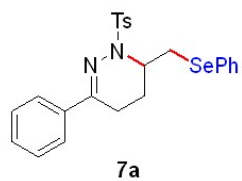
^{13}C NMR (151 MHz, CDCl_3) spectrum of **5e**.

Nov23-2024, 29, fid
JohnB-bj-1123-Y-15



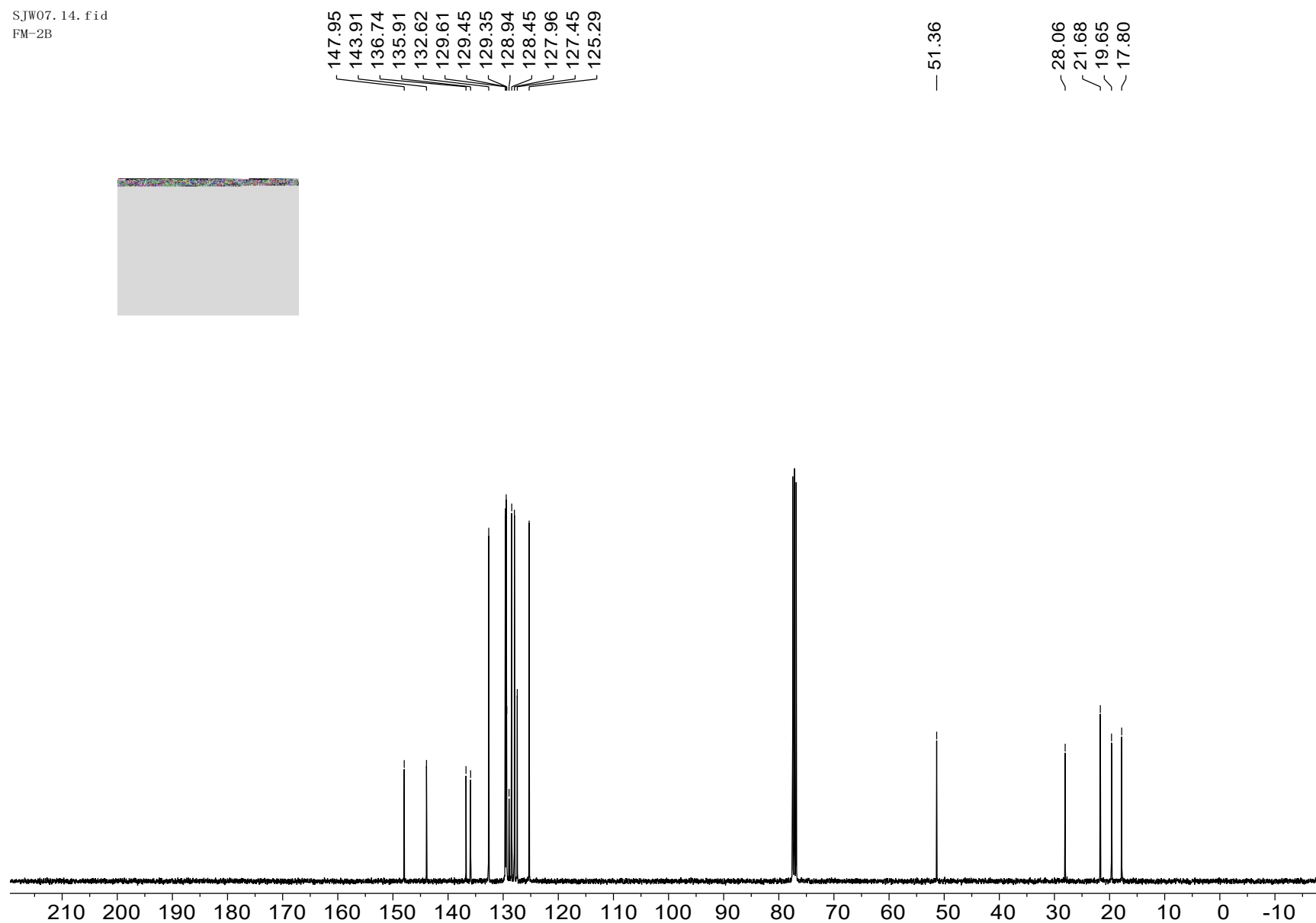
¹H NMR (500 MHz, CDCl₃) spectrum of **7a**.

SJW07.13.fid
FM-2B



^{13}C NMR (100 MHz, CDCl_3) spectrum of **7a**.

SJW07.14.fid
FM-2B



^1H NMR (500 MHz, CDCl_3) spectrum of **9a**.

SJW07.10.fid
FM-1B

