

Electronic Supporting Information

Catalytic enantioselective synthesis of α -nitro epoxides via aminolytic kinetic resolution

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General Methods

All reactions requiring dry or inert conditions were conducted in flame dried glassware under a positive pressure of nitrogen. THF and DCM were freshly distilled prior to use respectively over LiAlH₄ and calcium hydride and stored under nitrogen, all other solvents were dried over molecular sieves. Molecular sieves (Aldrich Molecular Sieves, 3 Å, 1.6 mm pellets) were activated under vacuum at 200°C overnight. Reactions were monitored by thin layer chromatography (TLC) on Merck silica gel plates (0.25 mm) and visualised by UV light or by phosphomolybdic acid/ethanol spray test. Flash chromatography was performed on Merck silica gel (60, particle size: 0.040–0.063 mm). ¹H NMR and ¹³C NMR spectra were recorded on Bruker Avance-400 spectrometer, Bruker Avance-300 spectrometer and Bruker Avance-250 spectrometer at room temperature in CDCl₃ as solvent. Chemical shifts for protons are reported using residual CHCl₃ as internal reference (δ = 7.26 ppm). Carbon spectra were referenced to the shift of the ¹³C signal of CDCl₃ (δ = 77.0 ppm). Optical rotation of compounds was performed on a Jasco Dip-1000 digital polarimeter using the Na lamp (582 nm). FTIR spectra were recorded as thin films on KBr plates using Bruker Tensor 27 spectrometer and absorption maxima are reported in wavenumber (cm⁻¹). ESI-MS was performed using a Bio-Q triple quadrupole mass spectrometer (Micromass, Manchester, UK) equipped with an electrospray ion source. Elemental analyses were carried out by using Flash EA 1112 (Thermo Electron Corporation) analyzer. Melting points were measured with a Stuart Model SMP 30 melting point apparatus. Petrol ether (PE) refers to light petroleum ether (boiling point 40-60°C). Anhydrous toluene, dry methanol and all starting materials (unless otherwise noted) were purchased from Aldrich and used as received.

Catalysts **I**, **III** were purchased from Aldrich and compound **IV** from Strem Chemicals and used as received. Catalysts **II**,¹ **V**,² **VI**,² **VII**² and **VIII**³ were prepared according to the literature. Enantiomeric excesses of α -nitroepoxides **1a**, **1e-m** and compound **4a** were determined by HPLC (Waters-Breeze 2487, UV dual λ absorbance detector and 1525 Binary HPLC Pump) using Daicel chiral columns.

Experimental Procedures and Compounds Characterization Data

General procedure for the epoxidation of nitroalkenes

The (*E*)- α,β -disubstituted nitroalkenes were synthesized according to the literature.⁴

α -Nitroepoxides were prepared according to published procedures.⁵ To a stirred ice-bath cold suspension of (*E*)- α,β -disubstituted nitroalkene (6 mmol) in methanol (12 mL) and hydrogen peroxide 50% aqueous solution (450 μ L, 7.8 mmol), aqueous NaOH 2M (1.5 mL, 3 mmol) was added rapidly (ca. 5 minutes) with stirring. The reaction was stirred at 0°C for 1 h (5 h for compound **1h**). Then, water was added (30 mL), extracted with diethyl ether (3 x 30 mL) and the combined ethereal extracts were washed with brine (40 mL), dried with Na₂SO₄ and concentrated under vacuum. The residue was purified by flash chromatography (PE/ diethyl ether 100:5) to give compounds **1a**, **1e-m** (13-72% yields).

General procedure for the aminolytic kinetic resolution (AKR) of racemic α -nitro epoxides

A sample vial was charged with nitroepoxide **1** (0.20 mmol) and catalyst **VI** (22.5 mg, 0.04 mmol) in anhydrous toluene (2 mL). Aniline (55 μ L, 0.60 mmol) was added and the reaction stirred at room temperature for 84-115 h, monitored by TLC (eluent PE/ diethyl ether 95:5 or 90:10 and PE/ ethyl acetate 90:10 only for compounds **1,3g** and **1,3k**). The enantioenriched α -nitroepoxides **1a**, **1e-m** and products **3a-3e-m** were isolated by flash chromatography (eluting from PE/ diethyl ether 100:2 to 100:5 and to 80:20 only for compound **3m**). In particular, aniline and α -amino ketone **3m** have the same polarity. To remove aniline from the mixture of the two products, recovered after silica gel chromatography, the mixture was diluted with Et₂O and washed with water. As a general note, α -amino ketone **3** showed to be relatively unstable compounds and they have to be stored under nitrogen at low temperature.

General procedure for the kinetic resolution of **1a**-($\pm$) with thiol **5**

A sample vial was charged with nitroepoxide **1a**-($\pm$) (17.9 mg, 0.10 mmol), K₂CO₃ (2.8 mg, 0.02 mmol) and catalyst **VI** (8.5 mg, 0.015 mmol) in anhydrous toluene (1 mL). Then, 2-naphthalenethiol (19.2 mg, 0.12 mmol) was added and the reaction stirred at room temperature for 70 h, monitored by TLC (eluent PE/ diethyl ether 95:5). The enantioenriched α -nitroepoxide **1a** (4.5 mg, 25% yield) and product **6** (21.3 mg, 73% yield) were isolated by flash chromatography (eluting from PE/ diethyl ether 100:1 to 80:20).

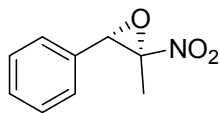
General procedure for the kinetic resolution of **1a**-(±) with diamine **7**

A sample vial was charged with nitroepoxide **1a**-(±) (26.9 mg, 0.15 mmol) and catalyst **VI** (12.7 mg, 0.022 mmol) in anhydrous toluene (1.5 mL). Then *o*-phenylenediamine (19.5 mg, 0.18 mmol) was added and the reaction was stirred at room temperature for 47 h, monitored by TLC (eluent PE/ diethyl ether 95:5). The enantioenriched α -nitroepoxide **1a** (8.6 mg, 32% yield) and product **8** (18.5 mg, 56% yield) were isolated by flash chromatography (eluting from PE/ diethyl ether 100:2 to 90:10).

General procedure for one-pot stereoselective ring-opening/reduction sequence to amino alcohol **4a**

To a solution of nitroepoxide **1a**-(±) (22.6 mg, 0.126 mmol) in dry CHCl₃ (315 μ L), pyrrolidine (21 μ L, 0.252 mmol) was added at 0 °C. The resulting mixture was stirred for 6 hours at 0 °C (TLC eluent PE/ diethyl ether 95:5). After completion, dry MeOH (105 μ L) was added at 0 °C, followed by dry CeCl₃ (31.1 mg, 0.126 mmol, finely ground CeCl₃•7H₂O dried at 100 °C under vacuum overnight). The reaction mixture was stirred for 10 min at the same temperature and then NaBH₄ (4.8 mg, 0.126 mmol) was added. After stirring at 0 °C for 2.5 h, a second portion of NaBH₄ (4.8 mg, 0.126 mmol) and CeCl₃ (31.1 mg, 0.126 mmol) was added, and stirring continued for additional 3 h. After complete conversion of the ketone to the alcohol (TLC eluent ethyl acetate/ MeOH 10 mL : 1 mL with 6 drops of NH₄OH, stained with permanganate) the reaction was quenched by diluting with brine (20 mL) and extracting with ethyl acetate (3x20 mL). The combined organic layers were washed with brine, dried over anhydrous Na₂SO₄ and concentrated under reduced pressure. Purification by flash chromatography eluting with CH₂Cl₂ and then with diethyl ether afforded *anti*-1,2-amino alcohol **4a** in 78% yield (20.2 mg). Absolute configuration of (1*R*,2*S*)-1-phenyl-1-(pyrrolidin-1-yl)propan-2-ol **4a** was determined by comparison of optical rotation with the literature.⁶ The absolute configuration of α -nitroepoxide **1a** was assigned to be (2*R*,3*S*) and the absolute configuration of α -nitroepoxides **1** was assigned to be (2*R*,3*S*) by analogy.

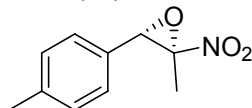
(2*R*, 3*S*)-2-Methyl-2-nitro-3-phenyloxirane (1a)



Data for this compound were consistent with those reported in the literature.^{5c}

Yellow oil, 10 mg, 28 % yield. $[\alpha]_D^{17} = -36.7$ (*c* 0.48, CHCl₃), *ee* 72%. **FTIR** ν_{max} (KBr)/cm⁻¹ 3066, 3035, 2947, 1558, 1451, 1357, 1159, 1107, 901, 770, 754, 703, 602. **¹H NMR** (CDCl₃, 400 MHz): δ 7.46-7.40 (m, 3H), 7.32-7.29 (m, 2H), 4.54 (s, 1H), 1.80 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 130.9, 129.4, 128.7, 126.4, 88.9, 62.6, 12.4. HPLC analysis with Chiralpak AS-H column, 95:5 *n*-hexane:2-propanol, 1 mL/min, 254 nm; minor enantiomer $t_R = 6.1$ min, major enantiomer $t_R = 6.8$ min.

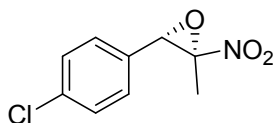
(2*R*, 3*S*)-2-Methyl-2-nitro-3-p-tolyloxirane (1e)



Data for this compound were consistent with those reported in the literature.^{5c}

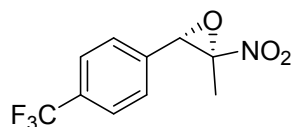
Yellow oil, 6.6 mg, 17% yield. $[\alpha]_D^{24} = -33.5$ (*c* 0.50, CHCl₃), *ee* 84%. **FTIR** ν_{max} (KBr)/cm⁻¹ 3028, 2925, 1559, 1452, 1346, 1159, 1106, 903, 848, 817, 772. **¹H NMR** (CDCl₃, 300 MHz): δ 7.25-7.16 (m, 4H), 4.50 (s, 1H), 2.38 (s, 3H), 1.80 (d, 3H, *J* = 1.3 Hz). **¹³C NMR** (CDCl₃, 100 MHz): δ 139.5, 129.4, 127.9, 126.3, 89.0, 62.8, 21.3, 12.4. **MS** (ESI *m/z*) 216.5 [MNa⁺, 12%]. HPLC analysis with Chiralpak IE-3 column, 98:2 *n*-hexane:2-propanol, 0.7 mL/min, 220 nm; minor enantiomer $t_R = 16.2$ min, major enantiomer $t_R = 14.2$ min.

(2*R*, 3*S*)-3-(4-Chlorophenyl)-2-methyl-2-nitrooxirane (1f)



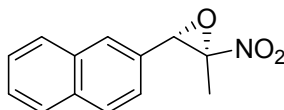
White solid, 12.4 mg, 29% yield. **Mp** 55.9-57.9 °C. $[\alpha]_D^{22} = -53.8$ (*c* 0.65, CHCl₃), *ee* 86%. **FTIR** ν_{max} (KBr)/cm⁻¹ 1561, 1495, 1436, 1352, 1159, 1091, 1015, 900, 763. **¹H NMR** (CDCl₃, 400 MHz): δ 7.41 (d, 2H, *J* = 8.2 Hz), 7.25 (d, 2H, *J* = 8.2 Hz), 4.52 (s, 1H), 1.79 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 135.6, 129.4, 129.1, 127.8, 88.6, 62.0, 12.4. Elemental analysis calcd (%) for C₉H₈ClNO₃: C, 50.60; H, 3.77; N, 6.56; found C, 50.89; H, 3.97; N, 6.60. HPLC analysis with Chiralpak AS-H column, 90:10 *n*-hexane:2-propanol, 1 mL/min, 220 nm; minor enantiomer $t_R = 6.9$ min, major enantiomer $t_R = 7.5$ min.

(2*R*, 3*S*)-2-Methyl-2-nitro-3-(4-(trifluoromethyl)phenyl)oxirane (1g)



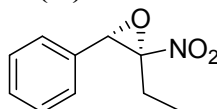
Yellow solid, 18 mg, 36% yield. **Mp** 58.2-62.6 °C. $[\alpha]_D^{23} = -30.9$ (*c* 0.53, CHCl₃), *ee* 77%. **FTIR** ν_{max} (KBr)/cm⁻¹ 1561, 1325, 1167, 1127, 1067, 1019, 857, 832. **¹H NMR** (CDCl₃, 250 MHz): δ 7.69 (d, 2H, *J* = 8.1 Hz), 7.45 (d, 2H, *J* = 8.1 Hz), 4.61 (s, 1H), 1.78 (s, 3H). **¹³C NMR** (CDCl₃, 75 MHz): δ 134.9, 131.7 (q, ²*J*_(C-C-F) = 32.8 Hz), 126.9, 125.9, 125.8, 123.6 (q, ¹*J*_(C-F) = 271.0 Hz), 88.5, 61.8, 12.4. **MS** (ESI *m/z*) 285.6 [MK⁺, 20%]. Elemental analysis calcd (%) for C₁₀H₈F₃NO₃: C, 48.59; H, 3.26; N, 5.67; found C, 48.94; H, 3.48; N, 5.40. HPLC analysis with Chiralpak IE-3 column, 98:2 *n*-hexane:2-propanol, 0.7 mL/min, 220 nm; minor enantiomer *t_R* = 18.4 min, major enantiomer *t_R* = 13.2 min.

(2*R*, 3*S*)-2-Methyl-3-(naphthalen-2-yl)-2-nitrooxirane (1h)



Pale yellow solid, 11.9 mg, 26% yield. **Mp** 63.8-65.7 °C. $[\alpha]_D^{20} = -72.2$ (*c* 0.67, CHCl₃), *ee* 95%. **FTIR** ν_{max} (KBr)/cm⁻¹ 3057, 2925, 1559, 1343, 1105, 893, 861, 821, 756. **¹H NMR** (CDCl₃, 300 MHz): δ 7.92-7.85 (m, 3H), 7.79 (s, 1H), 7.58-7.53 (m, 2H), 7.39 (dd, 1H, *J*₁ = 8.4 Hz, *J*₂ = 1.5 Hz), 4.71 (s, 1H), 1.83 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 133.6, 132.8, 128.8, 128.3, 128.0, 127.9, 126.9, 126.1, 123.2, 80.0, 62.9, 12.5. **MS** (ESI *m/z*) 230.0 [MH⁺, 6%], 248.6 [M+H₃O⁺, 38%]. Elemental analysis calcd (%) for C₁₃H₁₁NO₃: C, 68.11; H, 4.84; N, 6.11; found C, 68.44; H, 5.10; N, 6.24. HPLC analysis with Chiralpak IE-3 column, 98:2 *n*-hexane:2-propanol, 0.7 mL/min, 254 nm; minor enantiomer *t_R* = 24.7 min, major enantiomer *t_R* = 20.3 min.

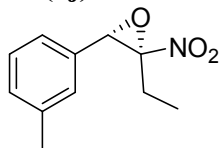
(2*R*, 3*S*)-2-Ethyl-2-nitro-3-phenyloxirane (1i)



Yellow oil, 8 mg, 21% yield. $[\alpha]_D^{22} = -25.1$ (*c* 0.67, CHCl₃), *ee* 92%. **FTIR** ν_{max} (KBr)/cm⁻¹ 2918, 1557, 1458, 1435, 1351, 938, 813, 768. **¹H NMR** (CDCl₃, 300 MHz): δ 7.44-7.38 (m, 3H), 7.33-7.29 (m, 2H), 4.52 (s, 1H), 2.55-2.41 (m, 1H), 1.69 (dq, 1H, *J*₁ = 15.1 Hz, *J*₂ = 7.3 Hz), 1.07 (t, 3H, *J*₁ = 7.4 Hz). **¹³C NMR** (CDCl₃, 100 MHz): δ 131.1, 129.4, 128.7, 126.3, 92.5, 63.2, 19.5, 7.6. **MS** (ESI *m/z*) 193.8 [MH⁺, 7%], 216.5 [MNa⁺, 15%]. Elemental analysis calcd (%) for C₁₀H₁₁NO₃: C, 62.17; H, 5.74; N, 7.25; found C, 62.46; H, 5.50; N, 7.48. HPLC analysis with Chiralpak IE-3

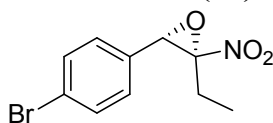
column, 98:2 *n*-hexane:2-propanol, 0.7 mL/min, 220 nm; minor enantiomer t_R = 13.7 min, major enantiomer t_R = 11.9 min.

(2*R*, 3*S*)-2-Ethyl-2-nitro-3-*m*-tolylloxirane (1j)



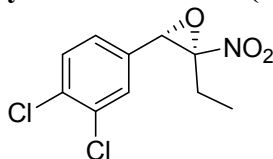
Yellow oil, 12.8 mg, 31% yield. $[\alpha]_D^{23}$ = -30.6 (*c* 0.57, CHCl₃), *ee* 92%. **FTIR** ν_{max} (KBr)/cm⁻¹ 2984, 2945, 1559, 1462, 1435, 1351, 1152, 1083, 969, 815, 790. **¹H NMR** (CDCl₃, 400 MHz): δ 7.30 (t, 1H, *J* = 7.5 Hz), 7.23-7.18 (m, 1H), 7.12-7.07 (m, 2H), 4.47 (s, 1H), 2.48 (dq, 1H, *J*₁ = 15.1 Hz, *J*₂ = 7.4 Hz), 2.38 (s, 3H), 1.69 (dq, 1H, *J*₁ = 15.1 Hz, *J*₂ = 7.4 Hz), 1.08 (t, 3H, *J* = 7.4 Hz). **¹³C NMR** (CDCl₃, 75 MHz): δ 138.6, 131.0, 130.1, 128.6, 126.9, 123.3, 92.6, 63.3, 21.4, 19.5, 7.6. **MS** (ESI *m/z*) 207.9 [MH⁺, 8%], 230.5 [MNa⁺, 10%]. Elemental analysis calcd (%) for C₁₁H₁₃NO₃: C, 63.76; H, 6.32; N, 6.76; found C, 64.07; H, 6.53; N, 6.63. HPLC analysis with Chiralpak IE-3 column, 98:2 *n*-hexane:2-propanol, 0.7 mL/min, 254 nm; minor enantiomer t_R = 13.4 min, major enantiomer t_R = 11.6 min.

(2*R*, 3*S*)-3-(4-Bromophenyl)-2-ethyl-2-nitrooxirane (1k)



White solid, 18.2 mg, 33% yield. **Mp** 59.6-60.3 °C. $[\alpha]_D^{21}$ = -29.0 (*c* 0.53, CHCl₃), *ee* 90%. **FTIR** ν_{max} (KBr)/cm⁻¹ 2984, 1557, 1489, 1463, 1434, 1346, 1071, 1011, 936, 808. **¹H NMR** (CDCl₃, 300 MHz): δ 7.56 (d, 2H, *J* = 8.4 Hz), 7.19 (d, 2H, *J* = 8.4 Hz), 4.48 (s, 1H), 2.46 (dq, 1H, *J*₁ = 15.1 Hz, *J*₂ = 7.4 Hz), 1.66 (dq, 1H, *J*₁ = 15.1 Hz, *J*₂ = 7.4 Hz), 1.07 (t, 3H, *J* = 7.4 Hz). **¹³C NMR** (CDCl₃, 100 MHz): δ 132.0, 130.1, 127.9, 123.7, 92.2, 62.6, 19.5, 7.6. **MS** (ESI *m/z*) 225.8 [M⁺-NO₂, 5%], 261.7 [M⁺-NO₂+2H₂O, 20%]. Elemental analysis calcd (%) for C₁₀H₁₀BrNO₃: C, 44.14; H, 3.70; N, 5.15; found C, 43.87; H, 3.88; N, 5.37. HPLC analysis with Chiralpak IE-3 column, 98:2 *n*-hexane:2-propanol, 0.7 mL/min, 220 nm; minor enantiomer t_R = 17.4 min, major enantiomer t_R = 14.6 min.

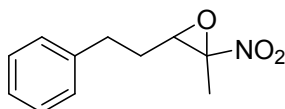
(2*R*, 3*S*)-3-(3,4-Dichlorophenyl)-2-ethyl-2-nitrooxirane (1l)



Yellow oil, 18.1 mg, 35% yield. $[\alpha]_D^{25}$ = -18.3 (*c* 0.76, CHCl₃), *ee* 61%. **FTIR** ν_{max} (KBr)/cm⁻¹ 2984, 2944, 1561, 1474, 1435, 1350, 1133, 1033, 944, 811. **¹H NMR** (CDCl₃, 400 MHz): δ 7.51 (d, 1H, *J* = 8.3 Hz), 7.40 (d, 1H, *J* = 1.9 Hz), 7.16 (dd, 1H, *J*₁ = 8.3 Hz, *J*₂ = 1.9 Hz), 4.48 (s, 1H), 2.46

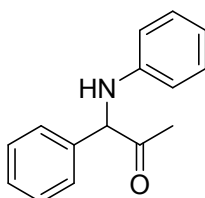
(dq, 1H, $J_1 = 15.1$ Hz, $J_2 = 7.4$ Hz), 1.65 (dq, 1H, $J_1 = 15.1$ Hz, $J_2 = 7.4$ Hz), 1.09 (t, 3H, $J = 7.4$ Hz). **^{13}C NMR** (CDCl_3 , 75 MHz): δ 133.8, 133.3, 131.3, 130.9, 128.2, 125.6, 92.0, 61.8, 19.5, 7.6. **MS** (ESI m/z) 251.6 [$\text{M}^+ - \text{NO}_2 + 2\text{H}_2\text{O}$, 100%]. Elemental analysis calcd (%) for $\text{C}_{10}\text{H}_9\text{Cl}_2\text{NO}_3$: C, 45.83; H, 3.46; N, 5.34; found C, 46.16; H, 3.24; N, 5.55. HPLC analysis with Chiralpak IE-3 column, 98:2 *n*-hexane:2-propanol, 0.7 mL/min, 254 nm; minor enantiomer $t_R = 17.3$ min, major enantiomer $t_R = 13.5$ min.

2-Methyl-2-nitro-3-phenethyloxirane (1m)



Yellow oil, 14.5 mg, 35% yield. $[\alpha]_D^{25} = -4.6$ (c 0.88, CHCl_3), *ee* 16%. **FTIR** ν_{max} (KBr)/ cm^{-1} 3027, 2924, 1557, 1496, 1455, 1388, 1357, 1112, 1083, 751, 699. **^1H NMR** (CDCl_3 , 400 MHz): δ 7.35-7.24 (m, 3H), 7.21-7.16 (m, 2H), 3.48 (t, 1H, $J = 6.3$ Hz), 2.95-2.87 (m, 1H), 2.83-2.74 (m, 1H), 2.06-1.95 (m, 1H), 1.93-1.83 (m, 1H), 1.74 (s, 3H). **^{13}C NMR** (CDCl_3 , 100 MHz): δ 139.6, 128.8, 128.3, 126.7, 62.4, 31.9, 29.8, 13.5. Elemental analysis calcd (%) for $\text{C}_{11}\text{H}_{13}\text{NO}_3$: C, 63.76; H, 6.32; N, 6.76; found C, 64.11; H, 6.58; N, 6.65. HPLC analysis with Chiralpak AS-H column, 99:1 *n*-hexane:2-propanol, 0.8 mL/min, 220 nm; minor enantiomer $t_R = 12.7$ min, major enantiomer $t_R = 11.5$ min.

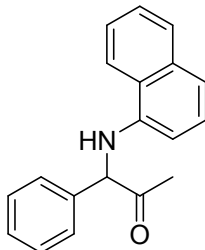
1-Phenyl-1-(phenylamino)propan-2-one (3a)



Data for this compound were consistent with those reported in the literature.⁷

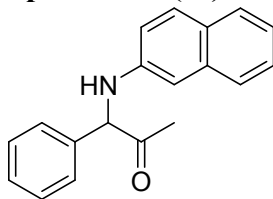
Wax, 29.3 mg, 65% yield. $[\alpha]_D^{27} = +27.9$ (c 0.50, CHCl_3), *ee* 16%. **FTIR** ν_{max} (KBr)/ cm^{-1} 3393, 2863, 1714, 1600, 1505, 1312, 1230, 1165, 745, 692. **^1H NMR** (CDCl_3 , 400 MHz): δ 7.55-7.32 (m, 5H), 7.13-7.06 (m, 2H), 6.73-6.66 (m, 1H), 6.63-6.57 (m, 2H), 5.01 (brd, 1H), 2.14 (s, 3H). **^{13}C NMR** (CDCl_3 , 75 MHz): δ 203.8, 145.4, 137.6, 129.7, 129.2, 129.1, 128.8, 127.8, 118.0, 113.7, 68.4, 26.7. **MS** (ESI m/z) 248.5 [MNa^+ , 10%]. HPLC analysis with Chiralpak IE-3 column, 95:5 *n*-hexane:2-propanol, 0.7 mL/min, 254 nm; minor enantiomer $t_R = 18.8$ min, major enantiomer $t_R = 15.6$ min.

1-(naphthalen-1-ylamino)-1-phenylpropan-2-one (3b)



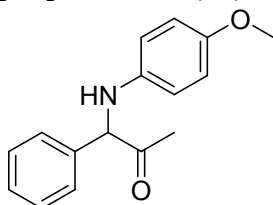
Red solid, 12.4 mg, 30% yield. **Mp** 81 °C (Decomp.). **FTIR** ν_{max} (KBr)/cm⁻¹ 3415, 3064, 1715, 1625, 1582, 1526, 1476, 1165, 769, 701. **¹H NMR** (CDCl₃, 400 MHz): δ 8.09 (d, 1H, J = 8.2 Hz), 7.76 (d, 1H, J = 8.0 Hz), 7.58-7.44 (m, 3H), 7.41-7.29 (m, 3H), 7.20-7.13 (m, 2H), 6.36-6.30 (m, 2H), 5.15 (d, 1H, J = 2.7 Hz), 2.20 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 204.0, 140.9, 137.8, 134.3, 129.3, 128.6, 128.5, 127.8, 126.3, 125.8, 124.9, 123.4, 120.2, 117.6, 105.3, 68.2, 26.6. **MS** (ESI m/z) 276.7 [MH⁺, 28%], 298.6 [MNa⁺, 100%]. Elemental analysis calcd (%) for C₁₉H₁₇NO: C, 82.88; H, 6.22; N, 5.09; found C, 83.25; H, 6.47; N, 4.94.

1-(naphthalen-2-ylamino)-1-phenylpropan-2-one (3c)



Red solid, 12.8 mg, 31% yield. **Mp** 83 °C (Decomp.). **FTIR** ν_{max} (KBr)/cm⁻¹ 3398, 3057, 1714, 1629, 1519, 1482, 1358, 1190, 827, 748, 701. **¹H NMR** (CDCl₃, 400 MHz): δ 7.64-7.58 (m, 2H), 7.54-7.45 (m, 3H), 7.42-7.37 (m, 2H), 7.34-7.28 (m, 1H), 7.18-7.13 (m, 1H), 6.96 (dd, 1H, J_1 = 8.8 Hz, J_2 = 2.5 Hz), 6.63-6.60 (m, 2H), 5.61 (brs, 1H), 5.13 (d, 1H, J = 4.3 Hz), 2.18 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 203.9, 143.5, 137.8, 134.8, 129.3, 129.0, 128.5, 127.8, 127.6, 126.2, 125.9, 122.1, 118.1, 105.4, 68.1, 26.7. **MS** (ESI m/z) 276.7 [MH⁺, 26%], 298.6 [MNa⁺, 100%]. Elemental analysis calcd (%) for C₁₉H₁₇NO: C, 82.88; H, 6.22; N, 5.09; found C, 83.22; H, 6.50; N, 4.84.

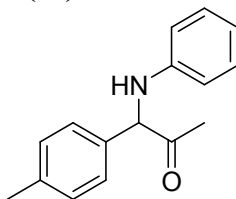
1-(4-methoxyphenylamino)-1-phenylpropan-2-one (3d)



Ochre wax, 14.6 mg, 38% yield. **FTIR** ν_{max} (KBr)/cm⁻¹ 3404, 1719, 1654, 1513, 1239, 1178, 1035, 820, 762, 701. **¹H NMR** (CDCl₃, 300 MHz): δ 7.48-7.30 (m, 5H), 6.69 (d, 2H, J = 8.8 Hz), 6.50 (d, 2H, J = 8.8 Hz), 5.15 (brs, 1H), 4.93 (s, 1H), 3.68 (s, 3H), 2.12 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 204.4, 152.1, 140.3, 138.3, 129.2, 128.3, 127.8, 114.8, 114.5, 69.1, 55.7, 26.7. **MS** (ESI

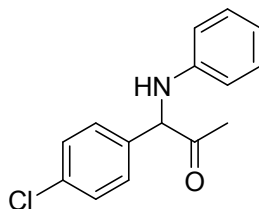
m/z) 255.7 [MH^+ , 13%], 278.0 [MNa^+ , 16%]. Elemental analysis calcd (%) for $\text{C}_{16}\text{H}_{17}\text{NO}_2$: C, 75.27; H, 6.71; N, 5.49; found C, 75.58; H, 7.00; N, 5.72.

1-(phenylamino)-1-p-tolylpropan-2-one (3e)



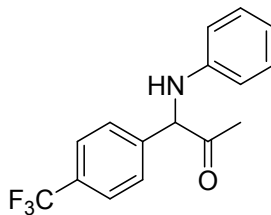
Red wax, 37.8 mg, 79% yield. **FTIR** ν_{max} (KBr)/ cm^{-1} 3422, 2921, 1707, 1668, 1601, 1496, 1444, 1314, 1159, 1113, 1030, 901, 823, 744. **^1H NMR** (CDCl_3 , 400 MHz): δ 7.32 (d, 2H, $J=7.7$ Hz), 7.17 (d, 2H, $J=7.7$ Hz), 7.13-7.06 (m, 2H), 6.72-6.66 (m, 1H), 6.64-6.58 (m, 2H), 4.99 (brs, 1H), 2.33 (s, 3H), 2.13 (s, 3H). **^{13}C NMR** (CDCl_3 , 75 MHz): δ 204.2, 145.8, 138.2, 134.8, 130.4, 129.9, 129.5, 129.1, 127.6, 117.6, 113.3, 67.8, 26.6, 21.1. **MS** (ESI m/z) 240.8 [MH^+ , 10%], 262.8 [MNa^+ , 14%]. Elemental analysis calcd (%) for $\text{C}_{16}\text{H}_{17}\text{NO}$: C, 80.30; H, 7.16; N, 5.85; found C, 79.98; H, 7.41; N, 6.06.

1-(4-chlorophenyl)-1-(phenylamino)propan-2-one (3f)



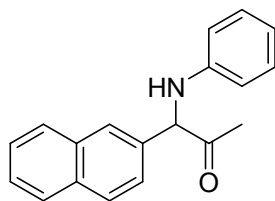
Red wax, 34.9 mg, 67% yield. **FTIR** ν_{max} (KBr)/ cm^{-1} 3449, 1715, 1670, 1601, 1542, 1500, 1489, 1441, 1315, 1162, 1091, 836, 752, 693. **^1H NMR** (CDCl_3 , 400 MHz): δ 7.46-7.33 (m, 4H), 7.14-7.07 (m, 2H), 6.70-6.66 (m, 1H), 6.55-6.50 (m, 2H), 4.98 (brs, 1H), 2.14 (s, 3H). **^{13}C NMR** (CDCl_3 , 100 MHz): δ 203.2, 145.5, 136.6, 131.7, 129.4, 129.2, 129.1, 117.9, 113.3, 67.4, 26.6. **MS** (ESI m/z) 260.5 [MH^+ , 37%], 282.5 [MNa^+ , 10%]. Elemental analysis calcd (%) for $\text{C}_{15}\text{H}_{14}\text{ClNO}$: C, 69.36; H, 5.43; N, 5.39; found C, 69.70; H, 5.60; N, 5.21.

1-(phenylamino)-1-(4-(trifluoromethyl)phenyl)propan-2-one (3g)



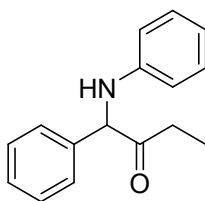
Yellow wax, 28.1 mg, 48% yield. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 3392, 3052, 2926, 1721, 1603, 1505, 1417, 1325, 1279, 1166, 1125, 1066, 1017, 845, 751, 692. **¹H NMR** (CDCl₃, 400 MHz): δ 7.67-7.58 (m, 4H), 7.12-7.07 (m, 2H), 6.70-6.66 (m, 1H), 6.53-6.48 (m, 2H), 5.47 (brs, 1H), 5.06 (brs, 1H), 2.16 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 202.7, 145.5, 142.3, 130.7 (q, $^2J_{\text{C-C-F}} = 32.3$ Hz), 129.3, 128.2, 126.2 (q, $^3J_{\text{C-C-C-F}} = 3.6$ Hz), 123.9 (q, $^1J_{\text{C-F}} = 270.5$ Hz), 118.1, 113.3, 67.8, 26.8. **MS** (ESI m/z) 293.6 [MH⁺, 17%], 316.7 [MNa⁺, 21%]. Elemental analysis calcd (%) for C₁₆H₁₄F₃NO: C, 65.52; H, 4.81; N, 4.78; found C, 65.79; H, 4.97; N, 4.63.

1-(naphthalen-2-yl)-1-(phenylamino)propan-2-one (3h)



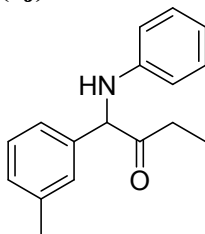
Red wax, 39.6 mg, 72% yield. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 3056, 1711, 1667, 1598, 1544, 1496, 1444, 1313, 1232, 1155, 1028, 821, 753, 693. **¹H NMR** (CDCl₃, 300 MHz): δ 7.99 (s, 1H), 7.93-7.80 (m, 2H), 7.58-7.35 (m, 5H), 7.17-7.05 (m, 2H), 6.70-6.58 (m, 2H), 5.17 (s, 1H), 2.16 (s, 3H). **¹³C NMR** (CDCl₃, 75 MHz): δ 203.9, 145.9, 135.5, 133.4, 133.2, 129.2, 129.1, 128.3, 127.8, 127.7, 127.3, 126.5, 126.3, 124.9, 117.7, 113.3, 68.3, 26.8. **MS** (ESI m/z) 276.6 [MH⁺, 39%], 298.6 [MNa⁺, 77%]. Elemental analysis calcd (%) for C₁₉H₁₇NO: C, 82.88; H, 6.22; N, 5.09; found C, 82.58; H, 6.38; N, 5.37.

1-phenyl-1-(phenylamino)butan-2-one (3i)



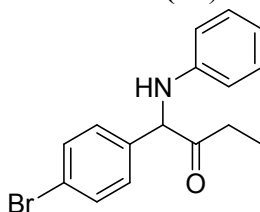
Ochre solid, 28.2 mg, 59% yield. **Mp** 84.2-86.7 °C. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 3395, 2923, 1716, 1603, 1504, 1454, 1428, 1317, 1260, 1111, 1034, 749. **¹H NMR** (CDCl₃, 400 MHz): δ 7.47-7.42 (m, 2H), 7.39-7.34 (m, 3H), 7.30 (d, 1H, $J = 7.2$ Hz), 7.12-7.05 (m, 2H), 6.65 (t, 1H, $J = 7.3$ Hz), 6.57-6.53 (m, 2H), 5.50 (brs, 1H), 5.00 (s, 1H), 2.53-2.40 (m, 2H), 0.99 (t, 3H, $J = 7.3$ Hz). **¹³C NMR** (CDCl₃, 100 MHz): δ 206.9, 146.0, 138.3, 129.14, 129.12, 128.3, 127.8, 117.6, 113.3, 67.4, 32.5, 7.86. **MS** (ESI m/z) 240.7 [MH⁺, 18%], 262.6 [MNa⁺, 43%]. Elemental analysis calcd (%) for C₁₆H₁₇NO: C, 80.30; H, 7.16; N, 5.85; found C, 80.57; H, 7.41; N, 6.12.

1-(phenylamino)-1-m-tolylbutan-2-one (3j)



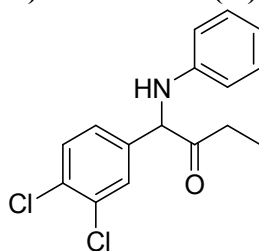
Orange wax, 33.4 mg, 66% yield. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 3386, 1716, 1603, 1505, 1428, 1321, 1111, 1035, 749, 692. **¹H NMR** (CDCl₃, 300 MHz): δ 7.31-7.23 (m, 3H), 7.16-7.08 (m, 3H), 6.68 (t, 1H, J = 7.3 Hz), 6.61-6.56 (m, 2H), 5.48 (brs, 1H), 4.98 (s, 1H), 2.48 (q, 2H, J = 7.3 Hz), 2.36 (s, 3H), 1.01 (t, 3H, J = 7.3 Hz). **¹³C NMR** (CDCl₃, 75 MHz): δ 207.1, 146.2, 138.9, 138.3, 129.13, 129.09, 128.95, 128.21, 125.0, 117.5, 113.3, 67.4, 32.4, 21.4, 7.9. **MS** (ESI m/z) 276.5 [MNa⁺, 100%]. Elemental analysis calcd (%) for C₁₇H₁₉NO: C, 80.60; H, 7.56; N, 5.53; found C, 80.28; H, 7.81; N, 5.80.

1-(4-Bromophenyl)-1-(phenylamino)butan-2-one (3k)



Red wax, 40.7 mg, 64% yield. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 3066, 1719, 1602, 1505, 1428, 1404, 1316, 1140, 1111, 1071, 1010, 833, 750, 692. **¹H NMR** (CDCl₃, 300 MHz): δ 7.50 (d, 2H, J = 8.3 Hz), 7.34 (d, 2H, J = 8.3 Hz), 7.13-7.05 (m, 2H), 6.67 (t, 1H, J = 7.2 Hz), 6.54-6.49 (m, 2H), 4.96 (s, 1H), 2.50-2.40 (m, 2H), 1.00 (t, 3H, J = 7.3 Hz). **¹³C NMR** (CDCl₃, 100 MHz): δ 206.1, 145.6, 137.4, 132.3, 129.3, 129.1, 122.2, 117.8, 113.3, 66.7, 32.4, 7.8. **MS** (ESI m/z) 318.5 [MH⁺, 45%]. Elemental analysis calcd (%) for C₁₆H₁₆BrNO: C, 60.39; H, 5.07; N, 4.40; found C, 60.13; H, 5.28; N, 4.63.

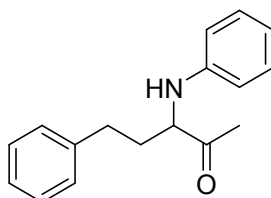
1-(3,4-Dichlorophenyl)-1-(phenylamino)butan-2-one (3l)



Red wax, 29 mg, 47% yield. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 3390, 1720, 1604, 1505, 1467, 1429, 1394, 1318, 1135, 1031, 752, 740, 692. **¹H NMR** (CDCl₃, 300 MHz): δ 7.56 (s, 1H), 7.48-7.42 (m, 1H), 7.35-7.28 (m, 1H), 7.16-7.07 (m, 2H), 6.70 (t, 1H, J = 7.3 Hz), 6.55-6.48 (m, 2H), 5.55 (brs, 1H),

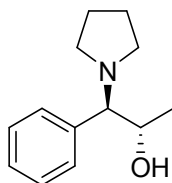
4.95 (s, 1H), 2.53-2.41 (m, 2H), 1.02 (t, 3H, $J = 7.3$ Hz). ^{13}C NMR (CDCl_3 , 75 MHz): δ 205.5, 145.4, 138.9, 133.4, 132.5, 131.1, 129.5, 129.2, 127.0, 118.1, 113.3, 66.4, 32.6, 7.7. MS (ESI m/z) 307.7 [MH^+ , 19%], 331.4 [MNa^+ , 12%]. Elemental analysis calcd (%) for $\text{C}_{16}\text{H}_{15}\text{Cl}_2\text{NO}$: C, 62.35; H, 4.91; N, 4.54; found C, 62.57; H, 5.07; N, 4.37.

5-Phenyl-3-(phenylamino)pentan-2-one (3m)



Yellow oil, 14.2 mg, 28% yield. Aniline and α -amino ketone **3m** have the same polarity. To remove aniline from the mixture of the two products, recovered after silica gel chromatography, the mixture was diluted with Et_2O and washed with water. FTIR ν_{max} (KBr)/ cm^{-1} 3441, 2918, 2850, 1705, 1635, 1602, 1506, 1454, 1260, 1093, 799, 748, 693. ^1H NMR (CDCl_3 , 400 MHz): δ 7.33-7.15 (m, 7H), 6.77-6.72 (m, 1H), 6.59-6.52 (m, 2H), 4.32 (brs, 1H), 4.07-4.02 (m, 1H), 2.94-2.84 (m, 1H), 2.80-2.65 (m, 2H), 2.19 (s, 3H), 2.03-1.93 (m, 1H). ^{13}C NMR (CDCl_3 , 100 MHz): δ 210.0, 146.8, 140.8, 129.4, 128.5, 128.4, 126.2, 118.1, 113.2, 62.8, 33.4, 31.5, 26.4. Elemental analysis calcd (%) for $\text{C}_{17}\text{H}_{19}\text{NO}$: C, 80.60; H, 7.56; N, 5.53; found C, 82.32; H, 7.72; N, 5.70.

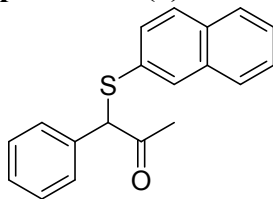
(1R,2S)-1-phenyl-1-(pyrrolidin-1-yl)propan-2-ol (4a)



Data for this compound were consistent with those reported in the literature.⁶

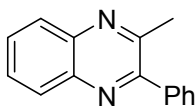
White solid, 20.2 mg, 78% yield. $[\alpha]_{\text{D}}^{24} = -53.0$ (c 0.82, CH_2Cl_2), ee 67%. ^1H NMR (CDCl_3 , 400 MHz): δ 7.32-7.27 (m, 5H), 4.27-4.21 (m, 1H), 2.99 (d, 1H, $J = 3.3$ Hz), 2.69-2.60 (m, 2H), 2.47-2.40 (m, 2H), 1.79-1.74 (m, 4H), 0.83 (d, 3H, $J = 6.4$ Hz). ^{13}C NMR (CDCl_3 , 100 MHz): δ 138.4, 129.4, 127.8, 127.4, 75.7, 66.9, 52.8, 23.3, 19.1. HPLC analysis with Chiralcel OD-H column, 99:1 n -hexane: ethanol, 1 mL/min, 220 nm; minor enantiomer $t_R = 7.5$ min, major enantiomer $t_R = 8.4$ min, ee 67 %.

1-(Naphthalen-2-ylthio)-1-phenylpropan-2-one (6)



Pink solid, 21.3 mg, 73% yield. **Mp** 70.8-74.3 °C. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 2923, 1712, 1585, 1493, 1454, 1355, 1067, 815, 744, 699, 569. **¹H NMR** (CDCl₃, 300 MHz): δ 7.81-7.68 (m, 4H), 7.48-7.44 (m, 2H), 7.41-7.31 (m, 6H), 5.10 (s, 1H), 2.21 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 203.1, 135.4, 133.5, 132.5, 131.3, 131.0, 129.3, 128.9, 128.6, 128.5, 128.3, 127.6, 127.5, 126.5, 126.4, 64.4, 27.3. **MS** (ESI m/z) 293.4 [MH⁺, 14%]. Elemental analysis calcd (%) for C₁₉H₁₆OS: C, 78.05; H, 5.52; S, 10.97; found C, 78.27; H, 5.34; S, 11.12.

2-methyl-3-phenylquinoxaline (8)



Data for this compound were consistent with those reported in the literature.⁸

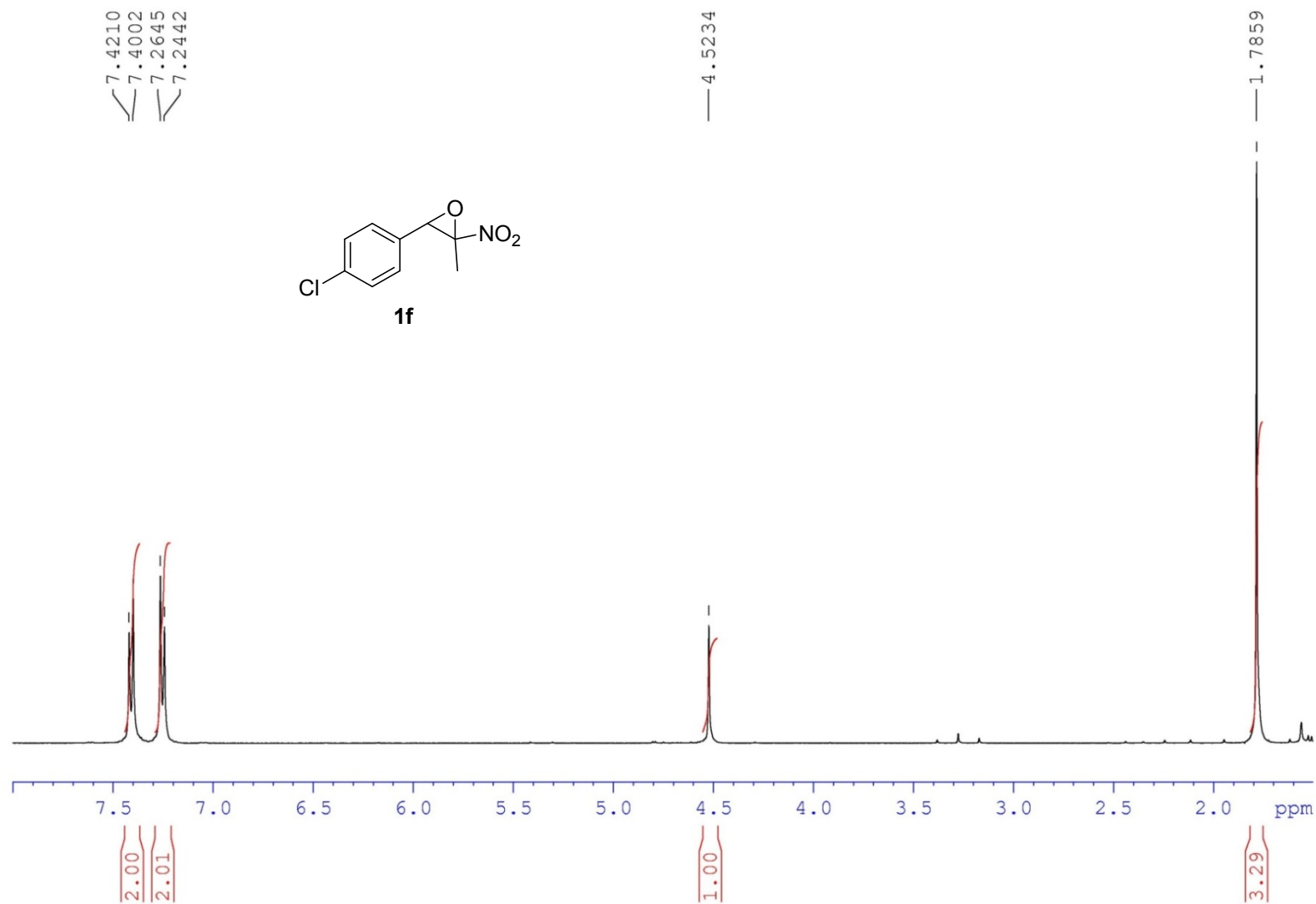
White wax, 18.5 mg, 56% yield. **FTIR** $\nu_{\max}$ (KBr)/cm⁻¹ 3060, 2924, 1562, 1483, 1445, 1396, 1374, 1343, 1191, 1132, 1006, 996, 766, 699. **¹H NMR** (CDCl₃, 400 MHz): δ 8.13-8.10 (m, 1H), 8.07-8.05 (m, 1H), 7.76-7.68 (m, 2H), 7.68-7.63 (m, 2H), 7.56-7.47 (m, 3H), 2.77 (s, 3H). **¹³C NMR** (CDCl₃, 100 MHz): δ 154.9, 152.5, 141.1, 140.9, 138.9, 129.7, 129.21, 129.17, 129.0, 128.9, 128.5, 128.2, 24.3.

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NMR Spectra

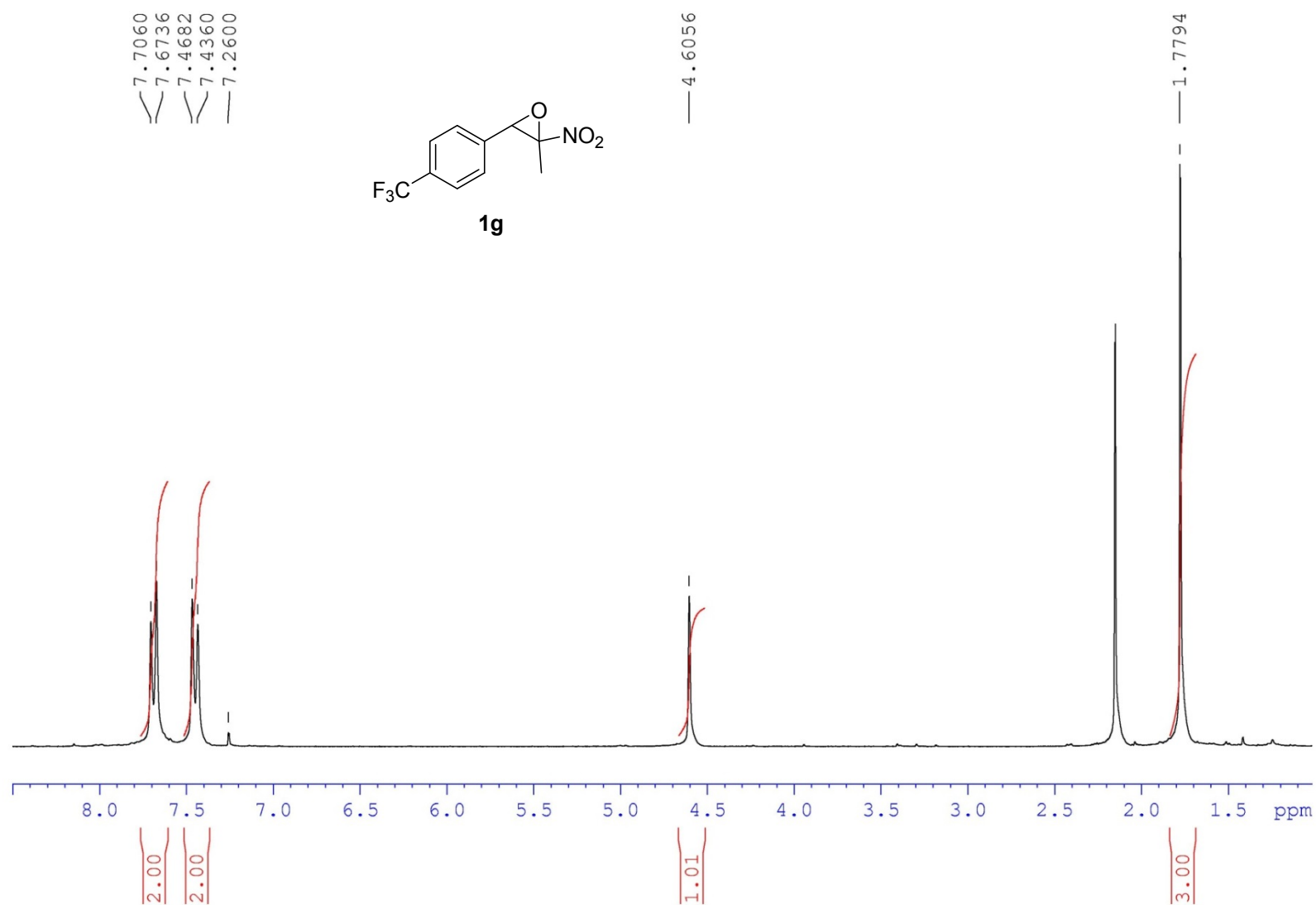
^1H NMR in CDCl_3 (400 MHz)



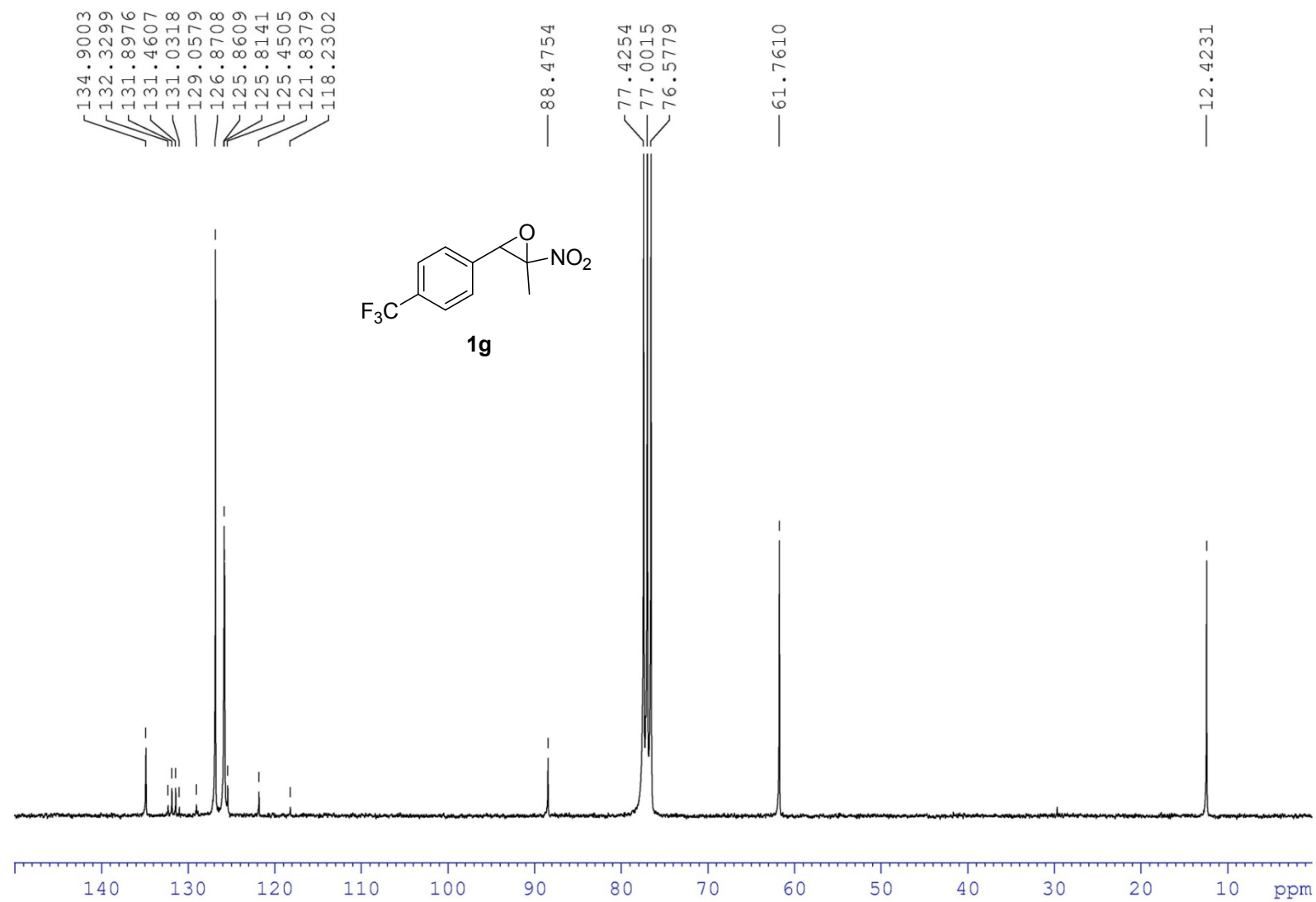
^{13}C NMR in CDCl_3 (100 MHz)



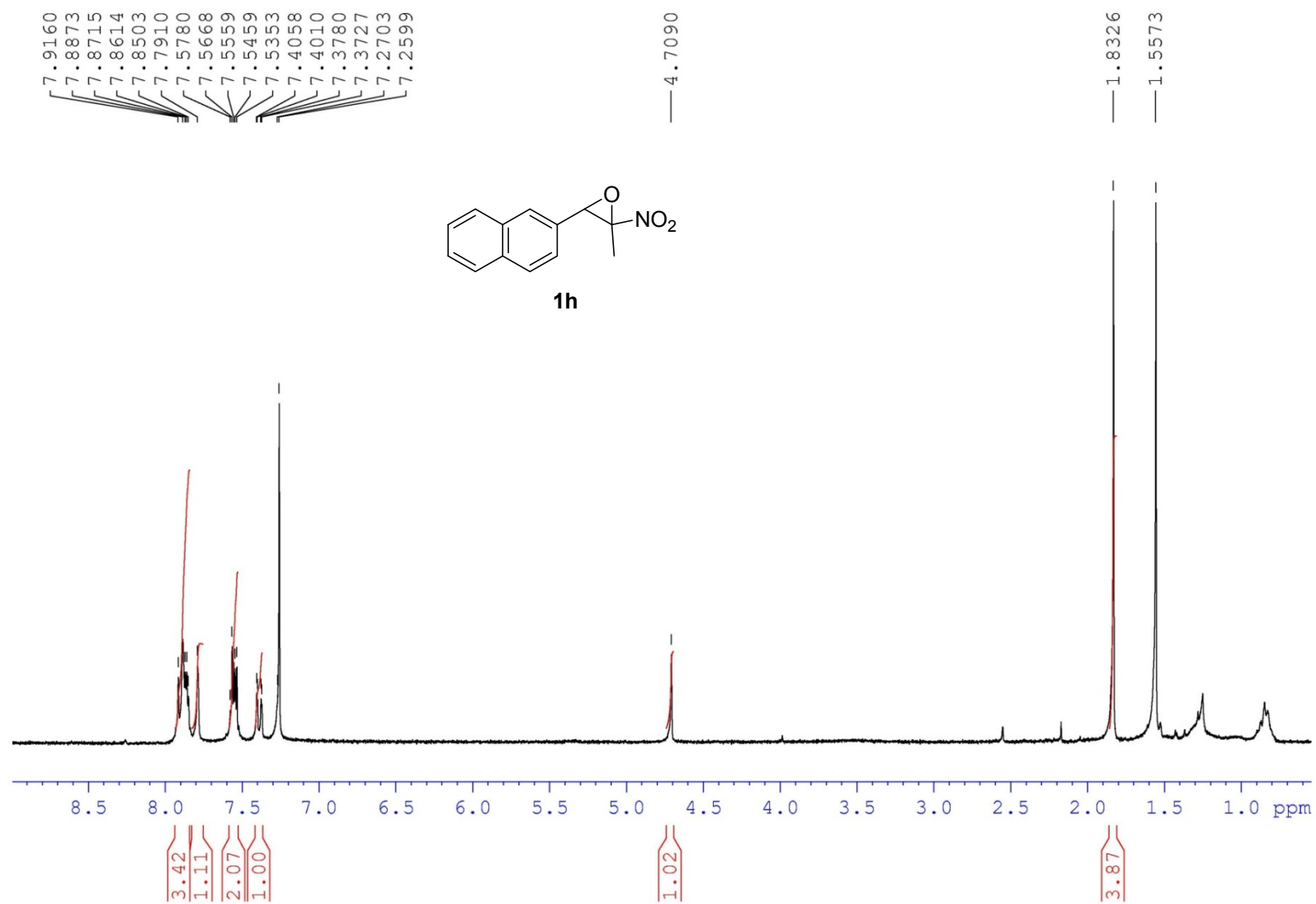
^1H NMR in CDCl_3 (250 MHz)



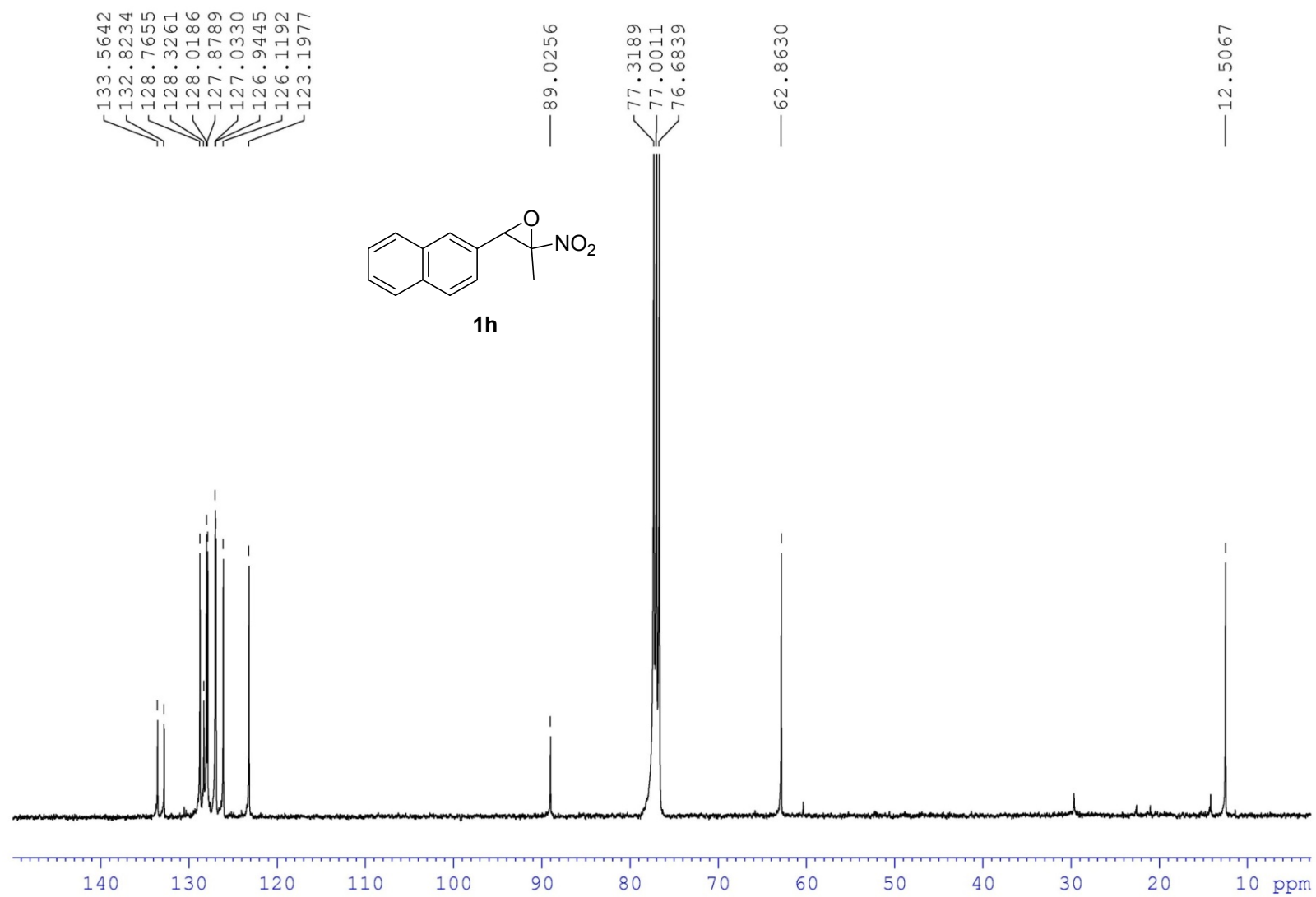
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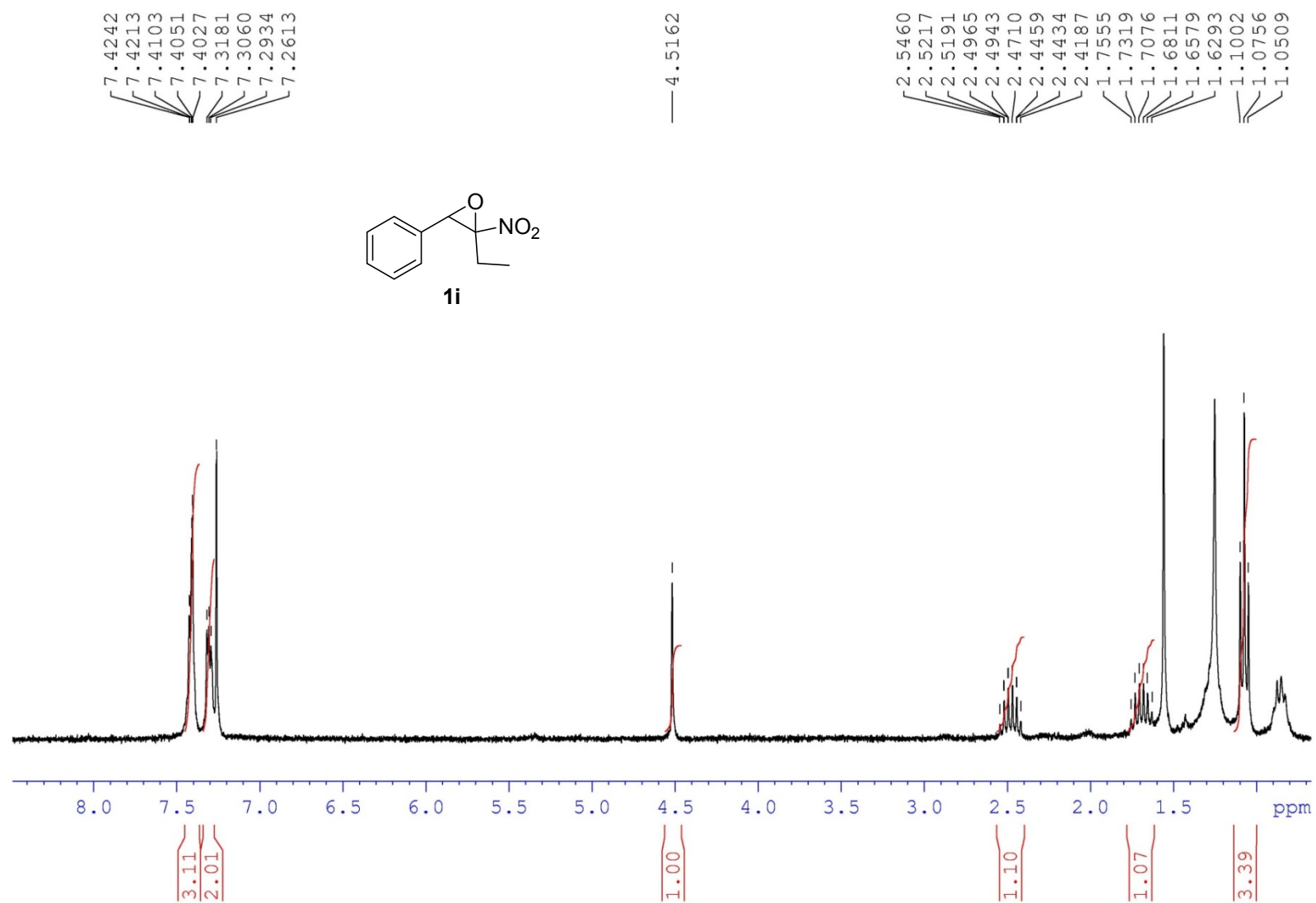
^1H NMR in CDCl_3 (300 MHz)



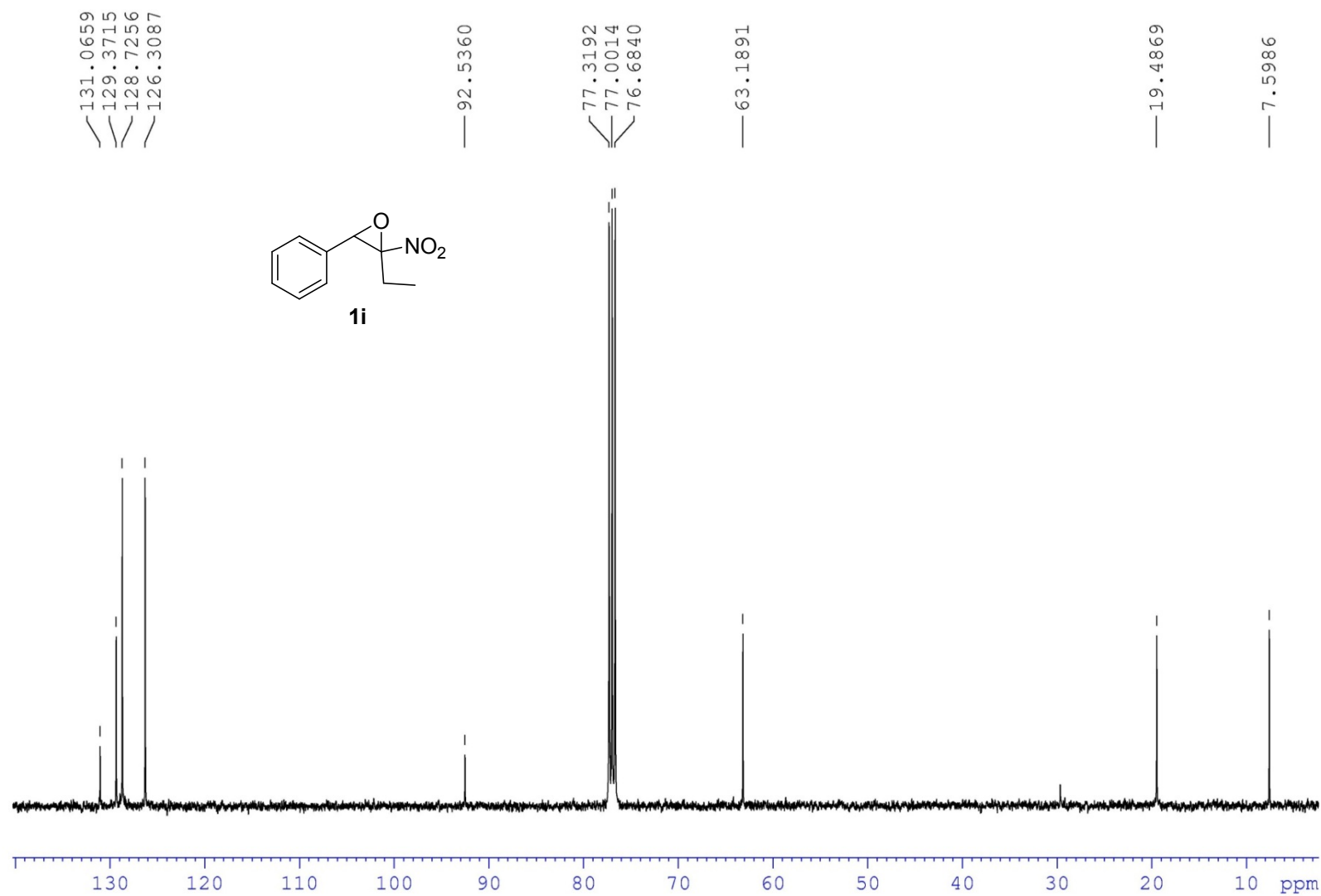
^{13}C NMR in CDCl_3 (100 MHz)



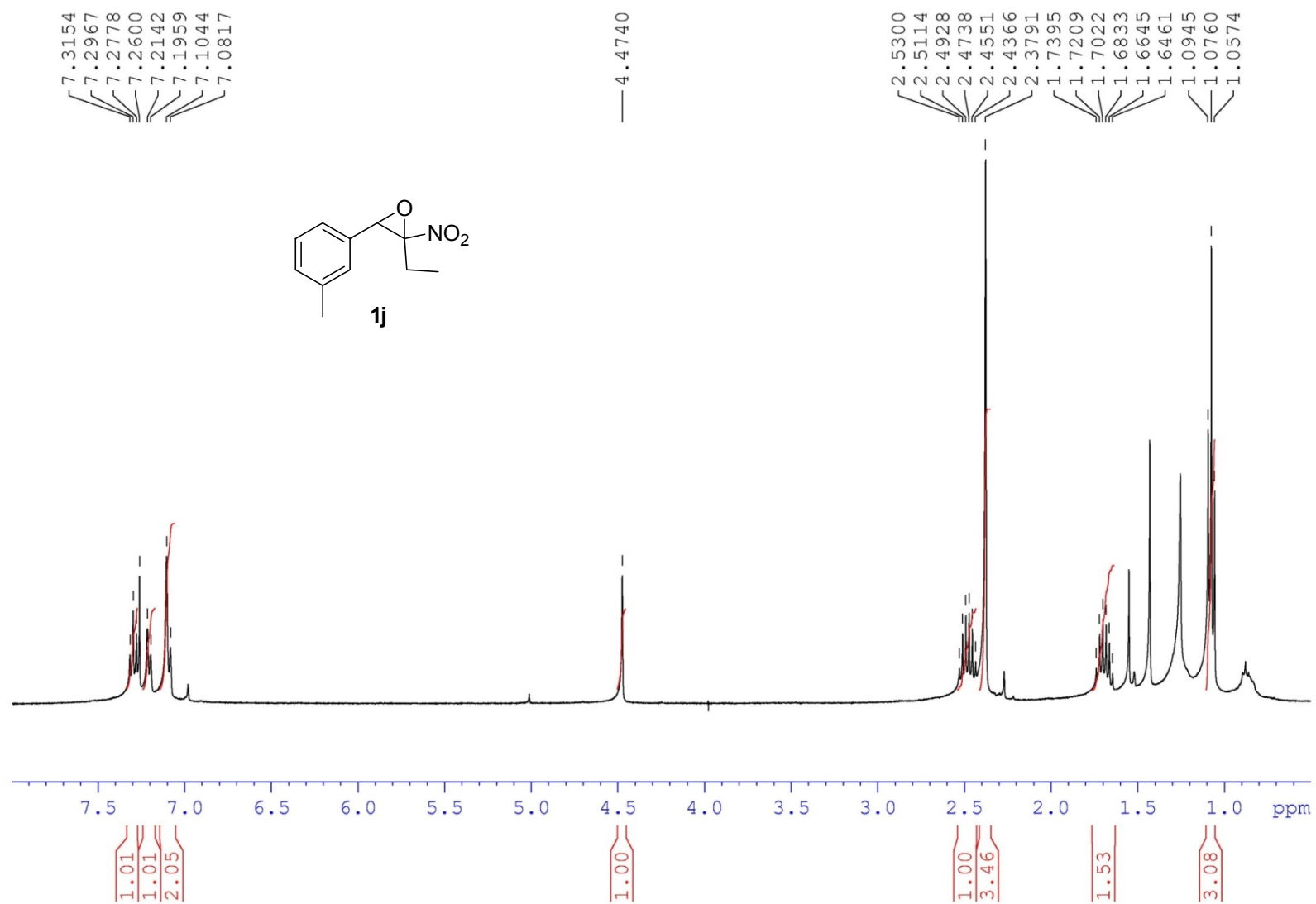
^1H NMR in CDCl_3 (300 MHz)



^{13}C NMR in CDCl_3 (100 MHz)



^1H NMR in CDCl_3 (400 MHz)



^{13}C NMR in CDCl_3 (75 MHz)

— 138.6077
— 130.9917
— 130.1374
— 128.6143
— 126.8547
— 123.3469

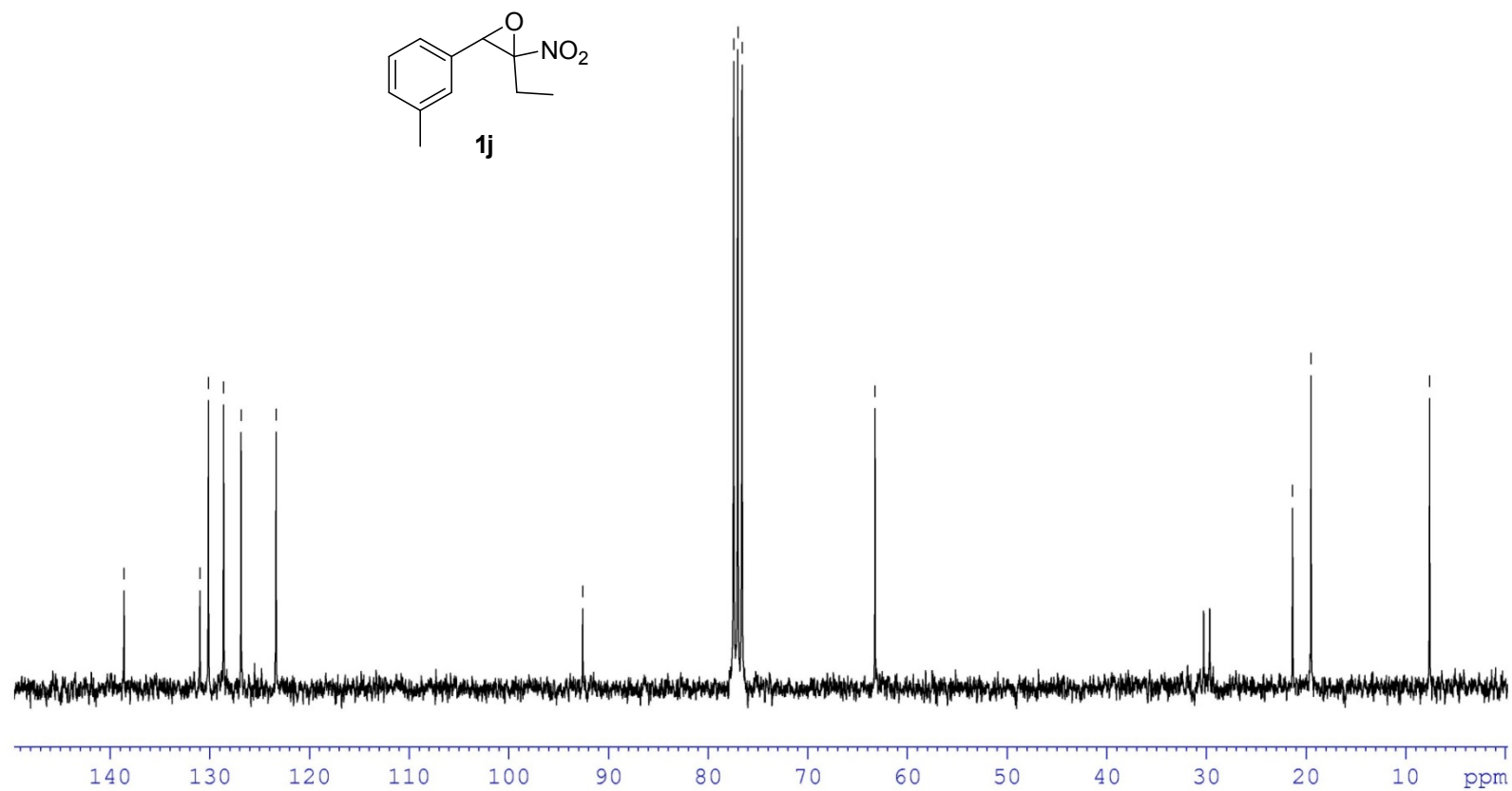
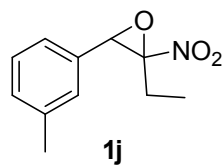
— 92.5725

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— 77.0029
— 76.5793

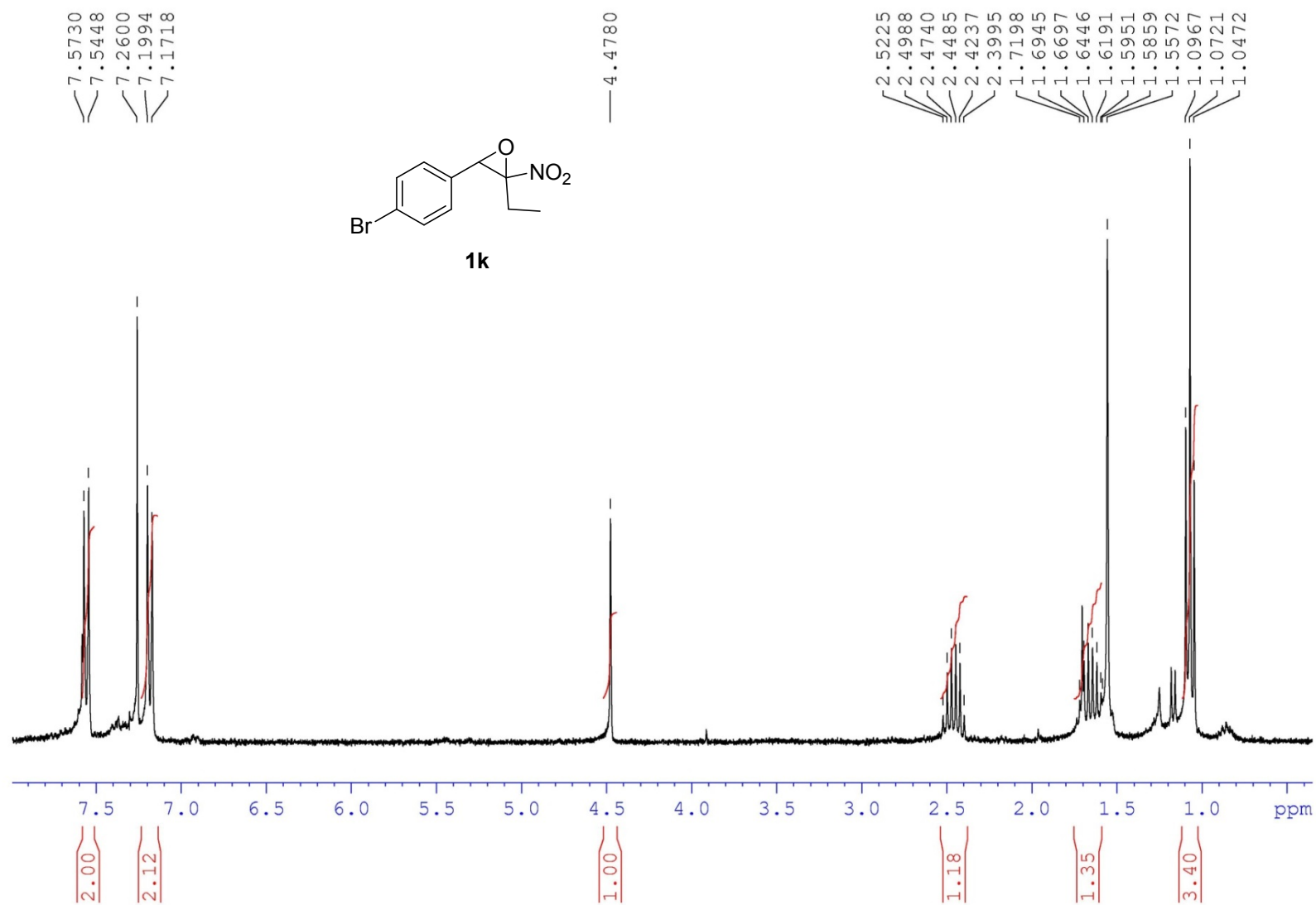
— 63.2718

— 21.3675
— 19.5095

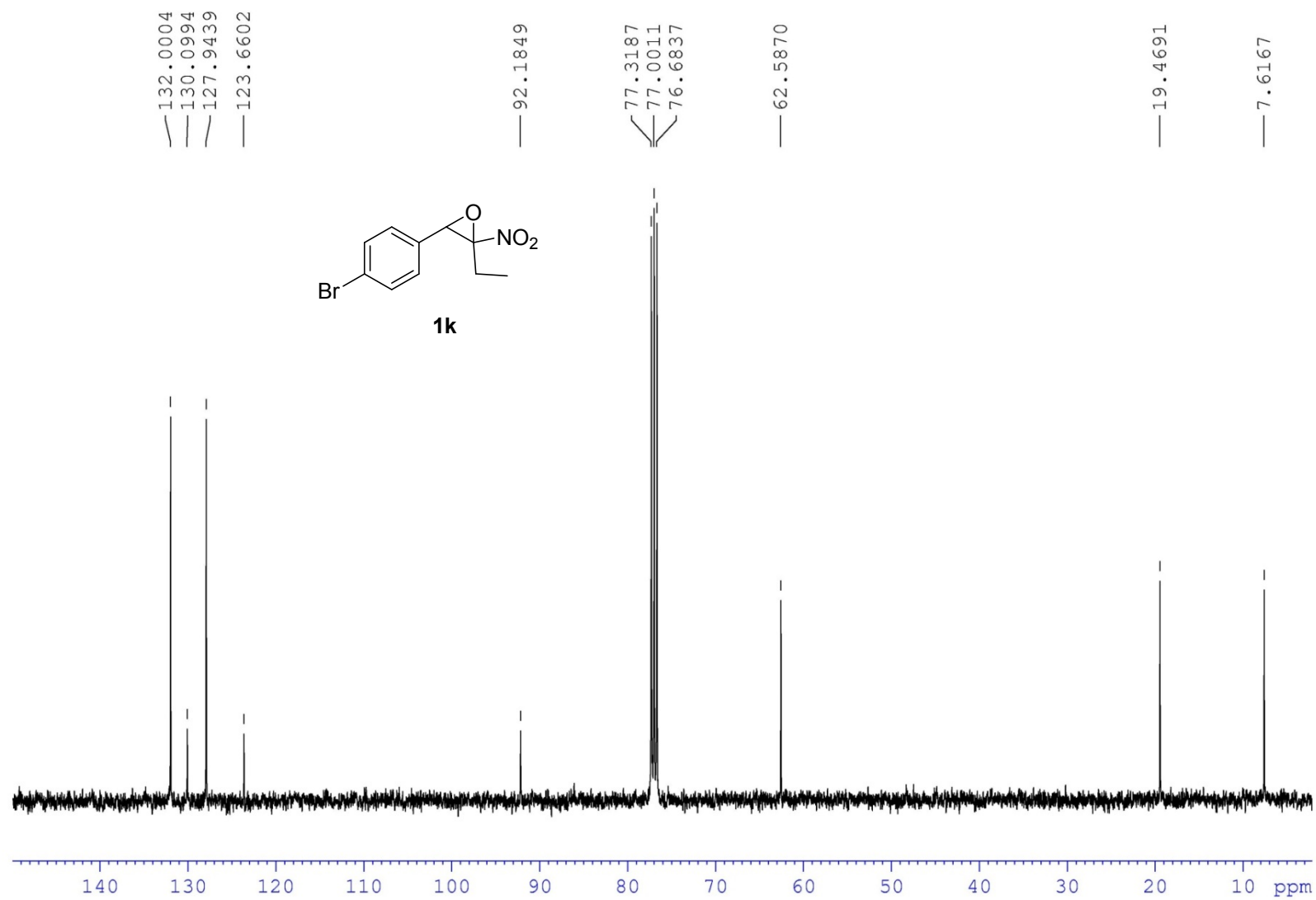
— 7.6121



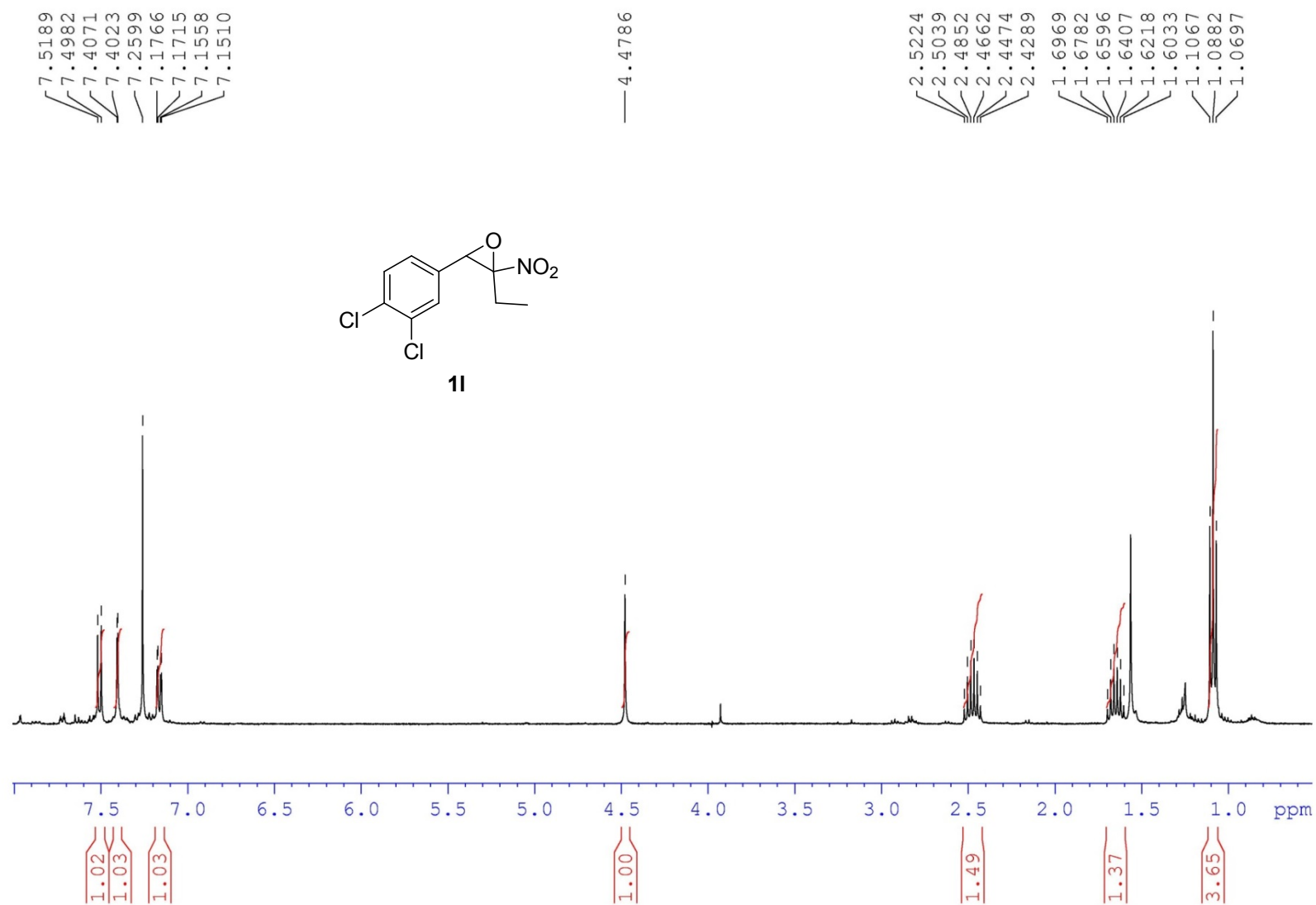
¹H NMR in CDCl₃ (300 MHz)



^{13}C NMR in CDCl_3 (100 MHz)



^1H NMR in CDCl_3 (400 MHz)



^{13}C NMR in CDCl_3 (75 MHz)

133.8413
133.3376
131.2798
130.8845
128.2404
125.5958

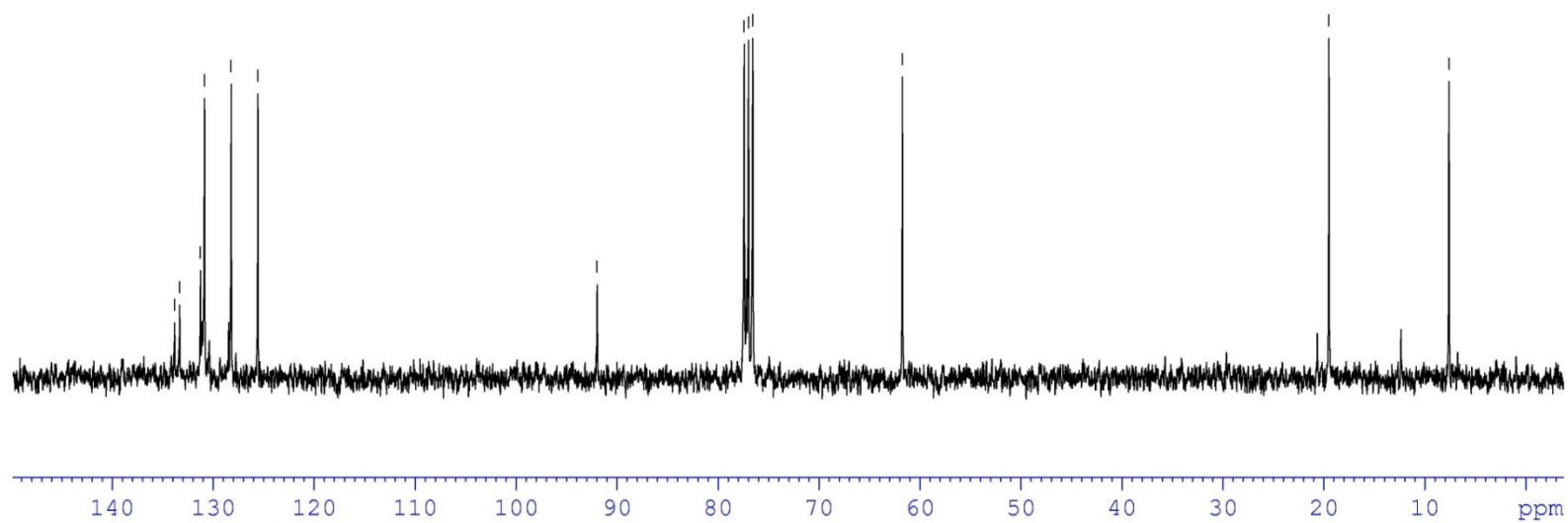
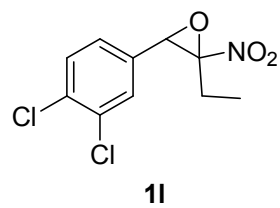
— 91.9920

77.4261
77.0016
76.5781

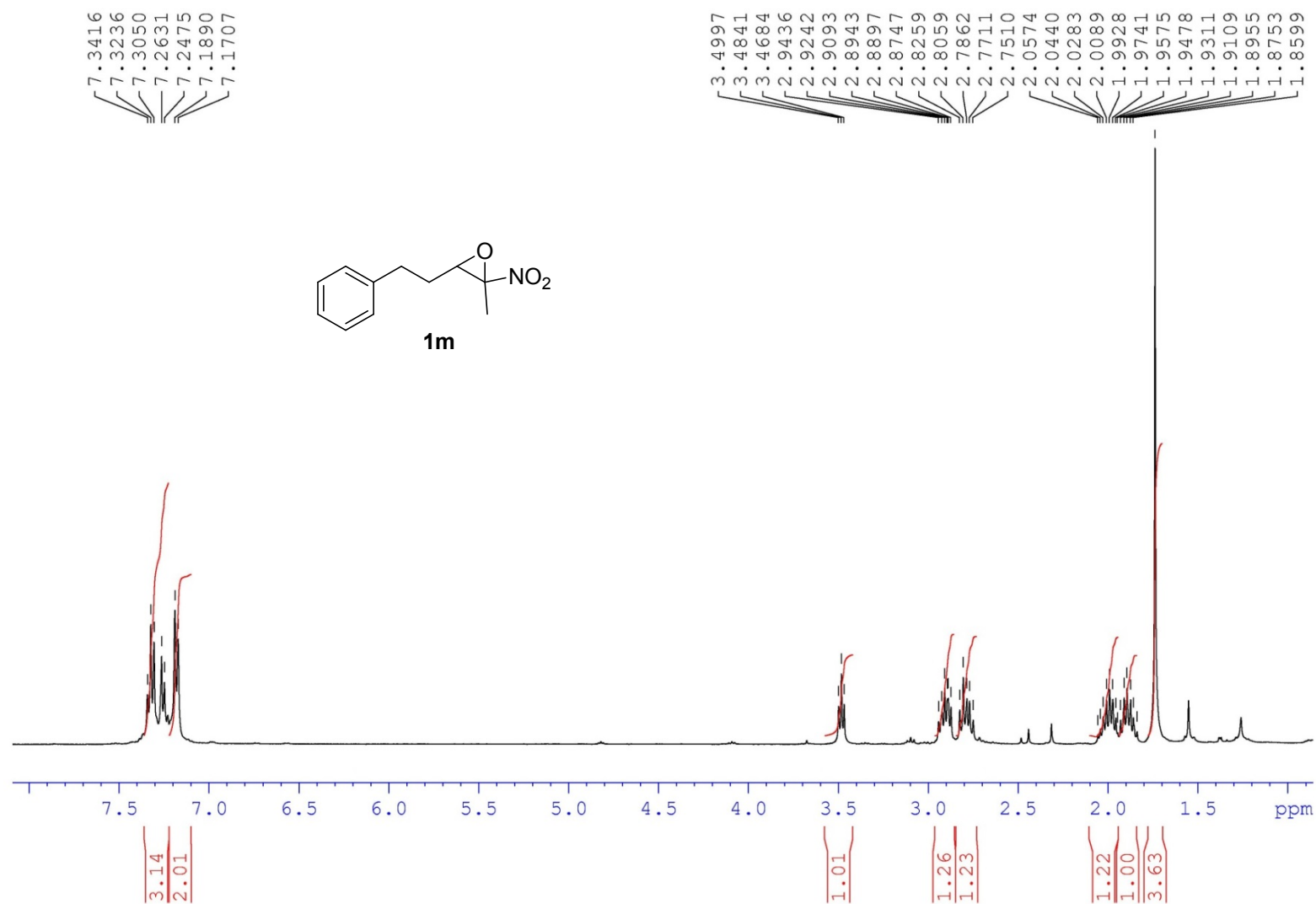
— 61.7522

— 19.5083

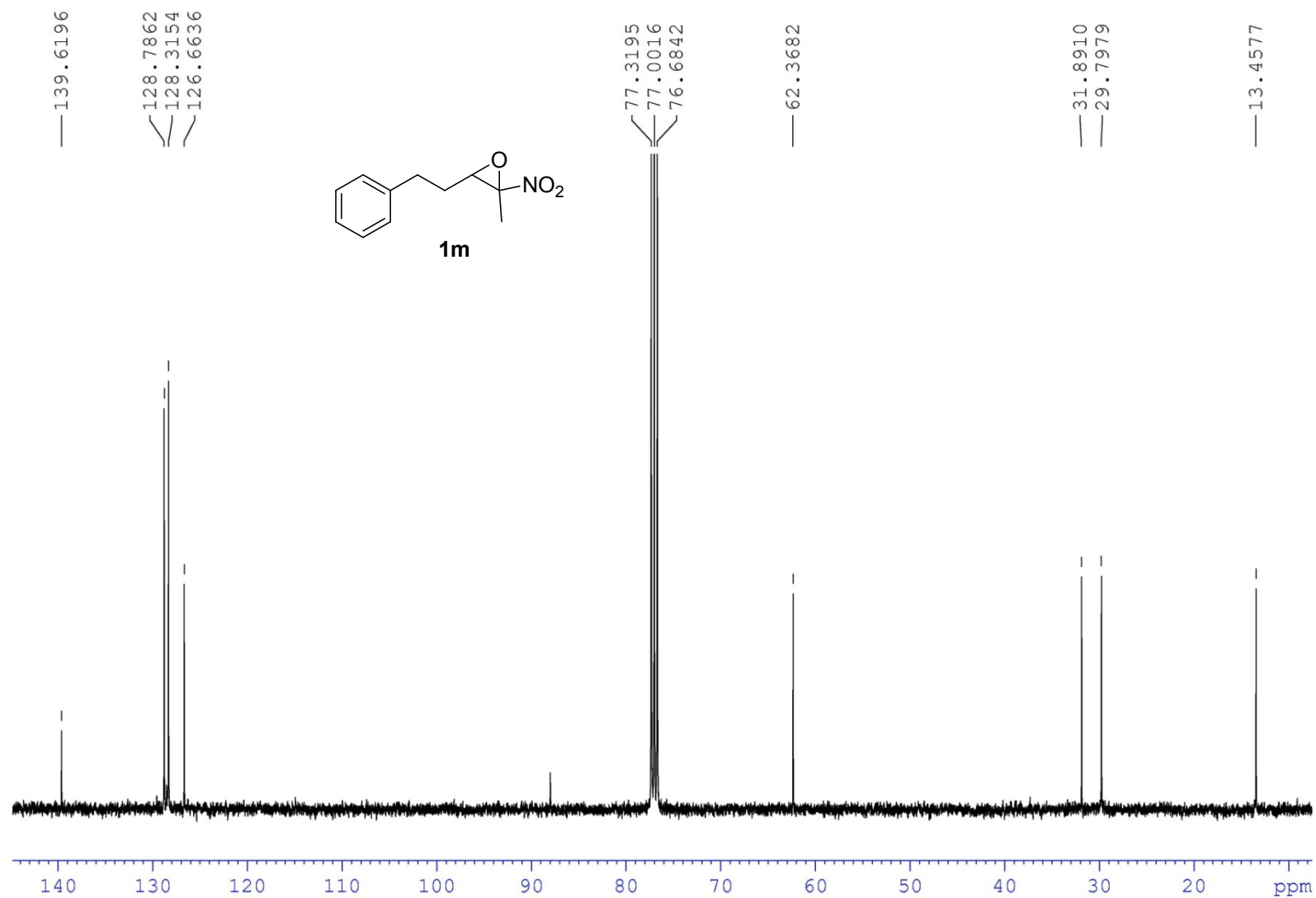
— 7.6208



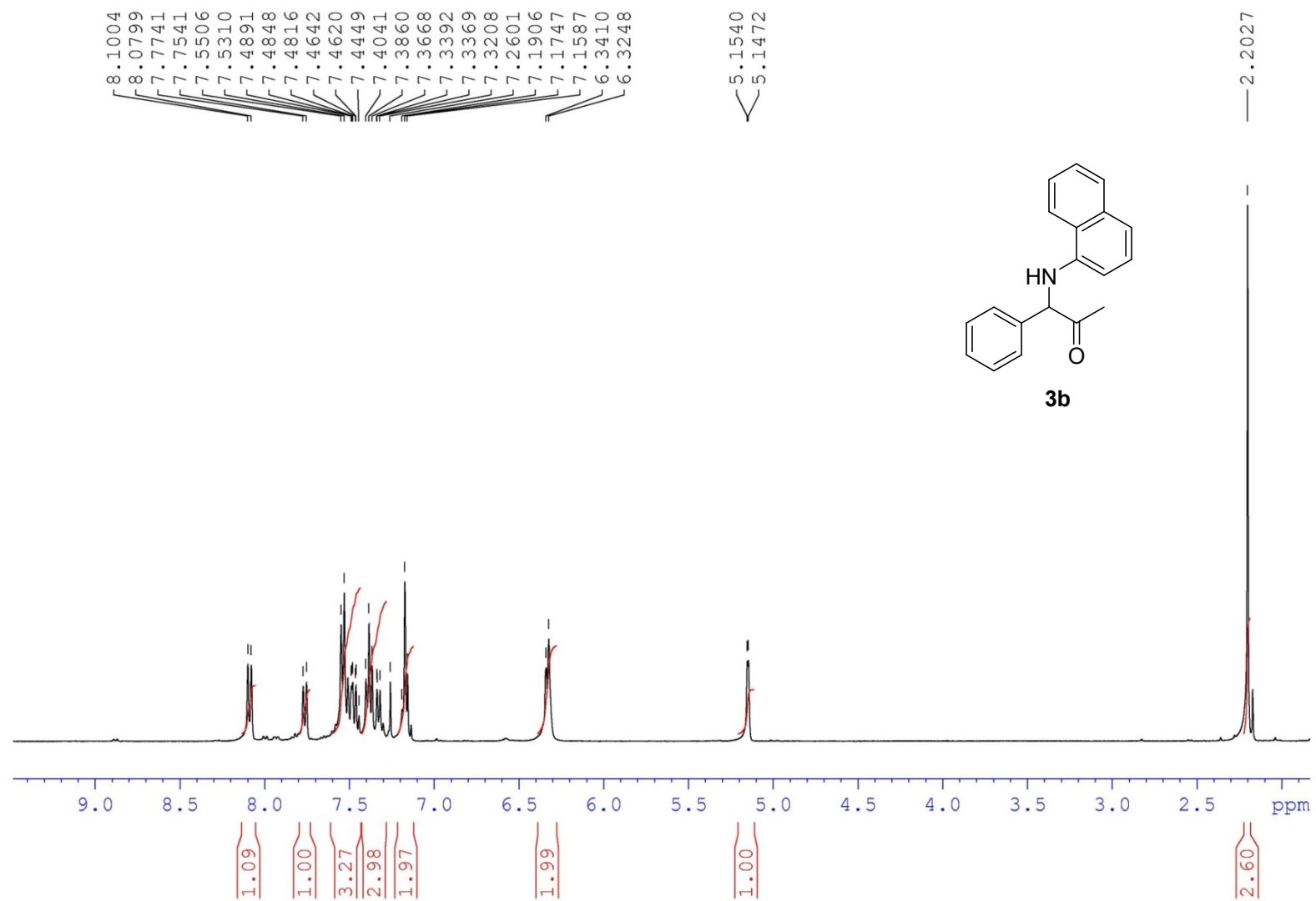
^1H NMR in CDCl_3 (400 MHz)



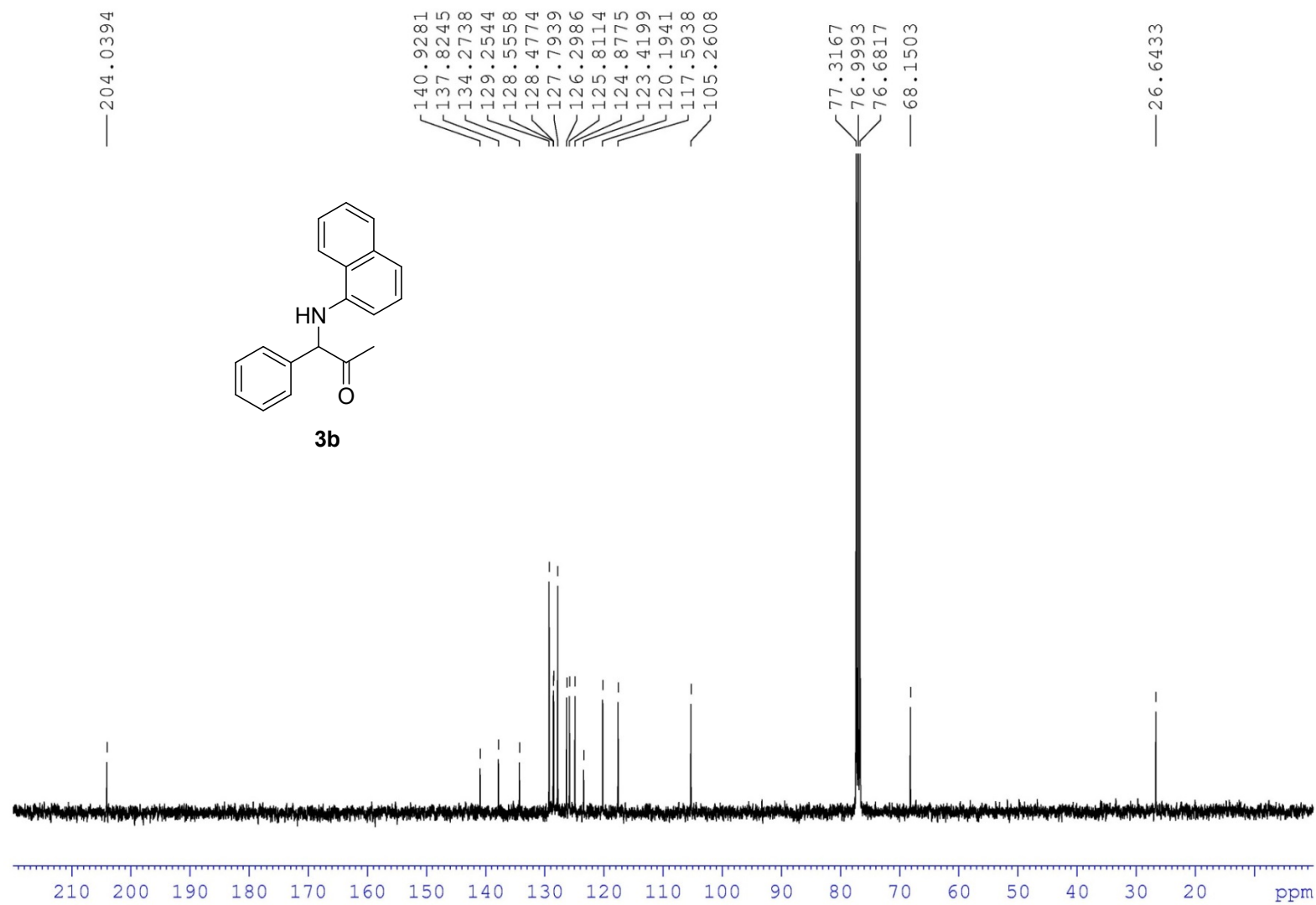
^{13}C NMR in CDCl_3 (100 MHz)



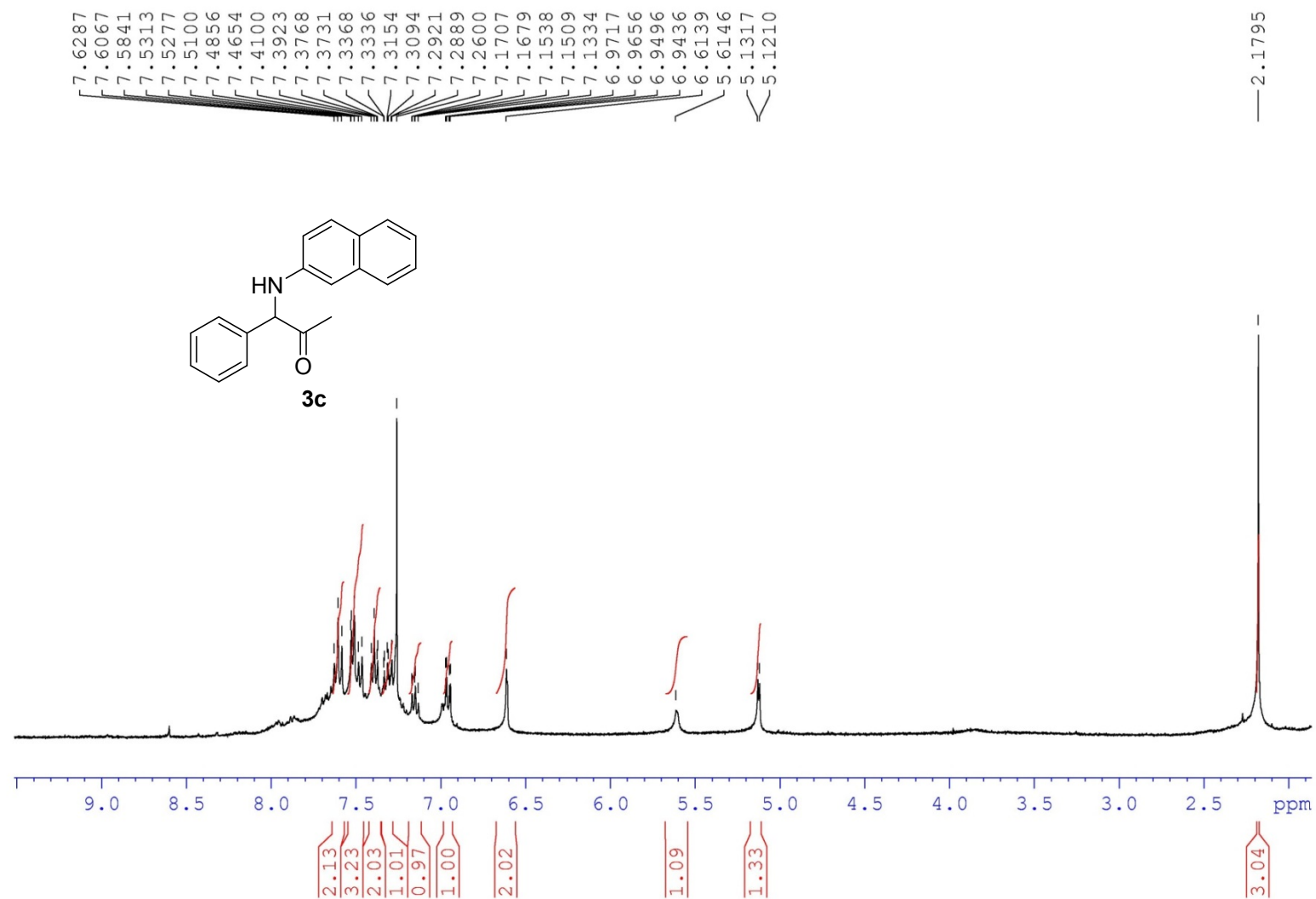
^1H NMR in CDCl_3 (400 MHz)



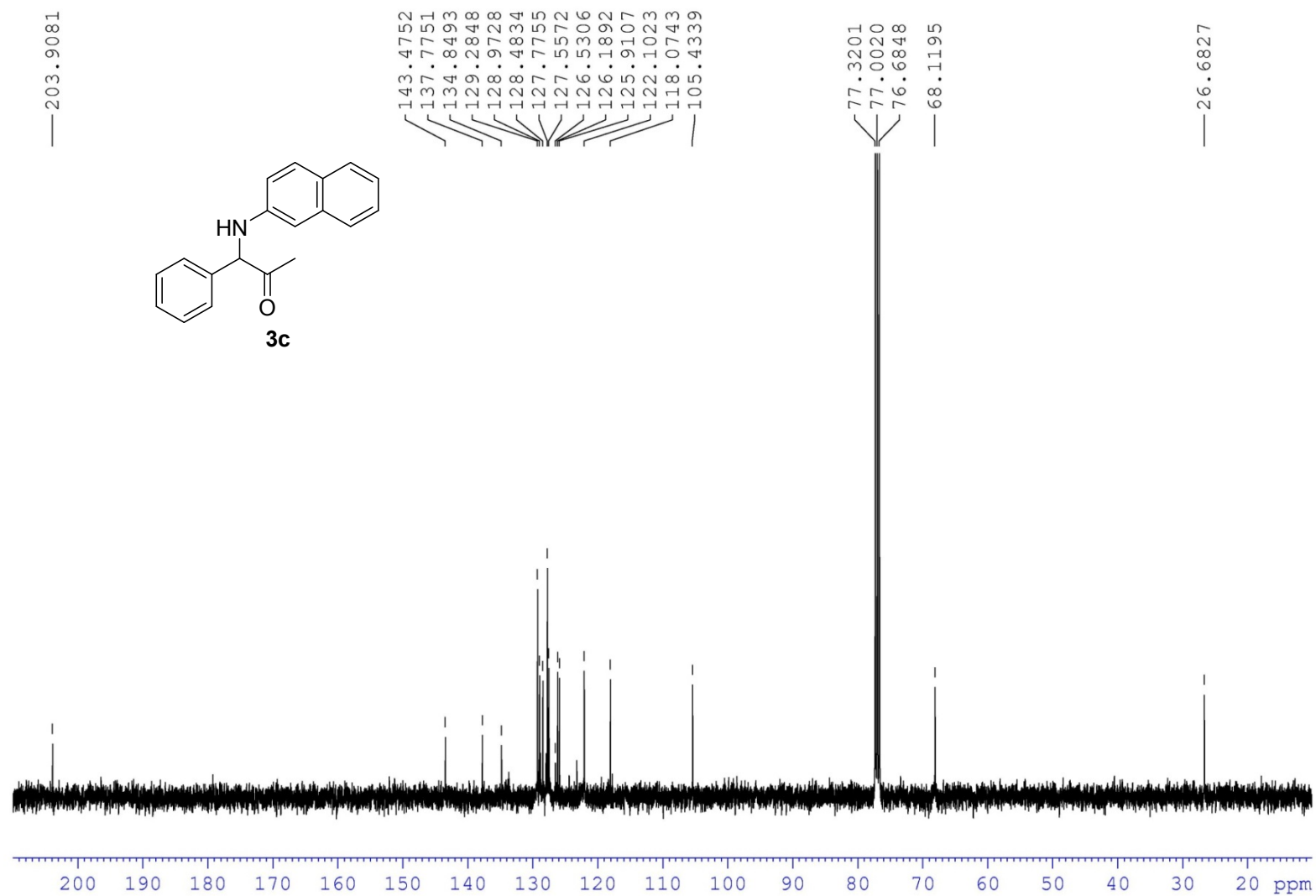
^{13}C NMR in CDCl_3 (100 MHz)



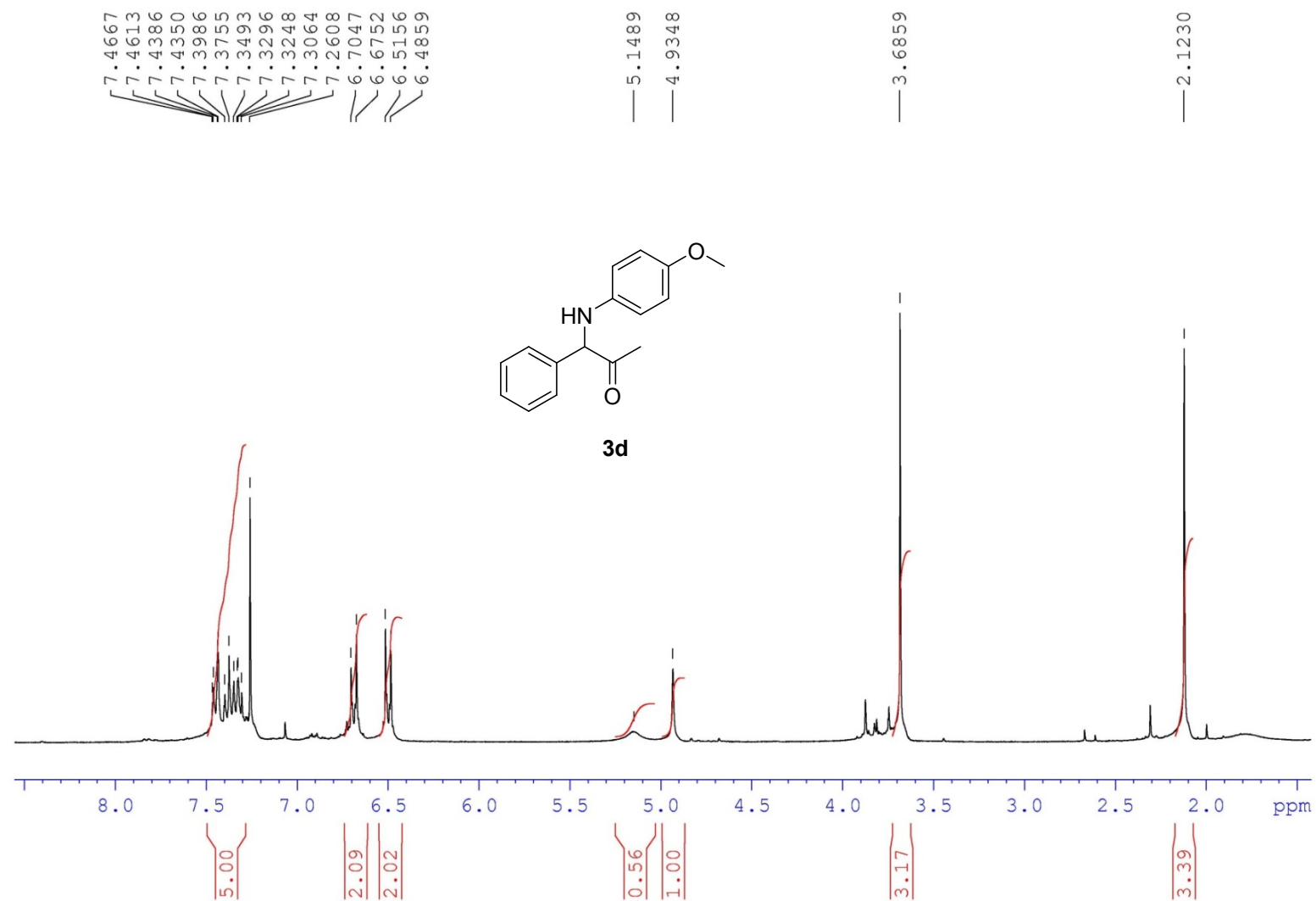
^1H NMR in CDCl_3 (400 MHz)



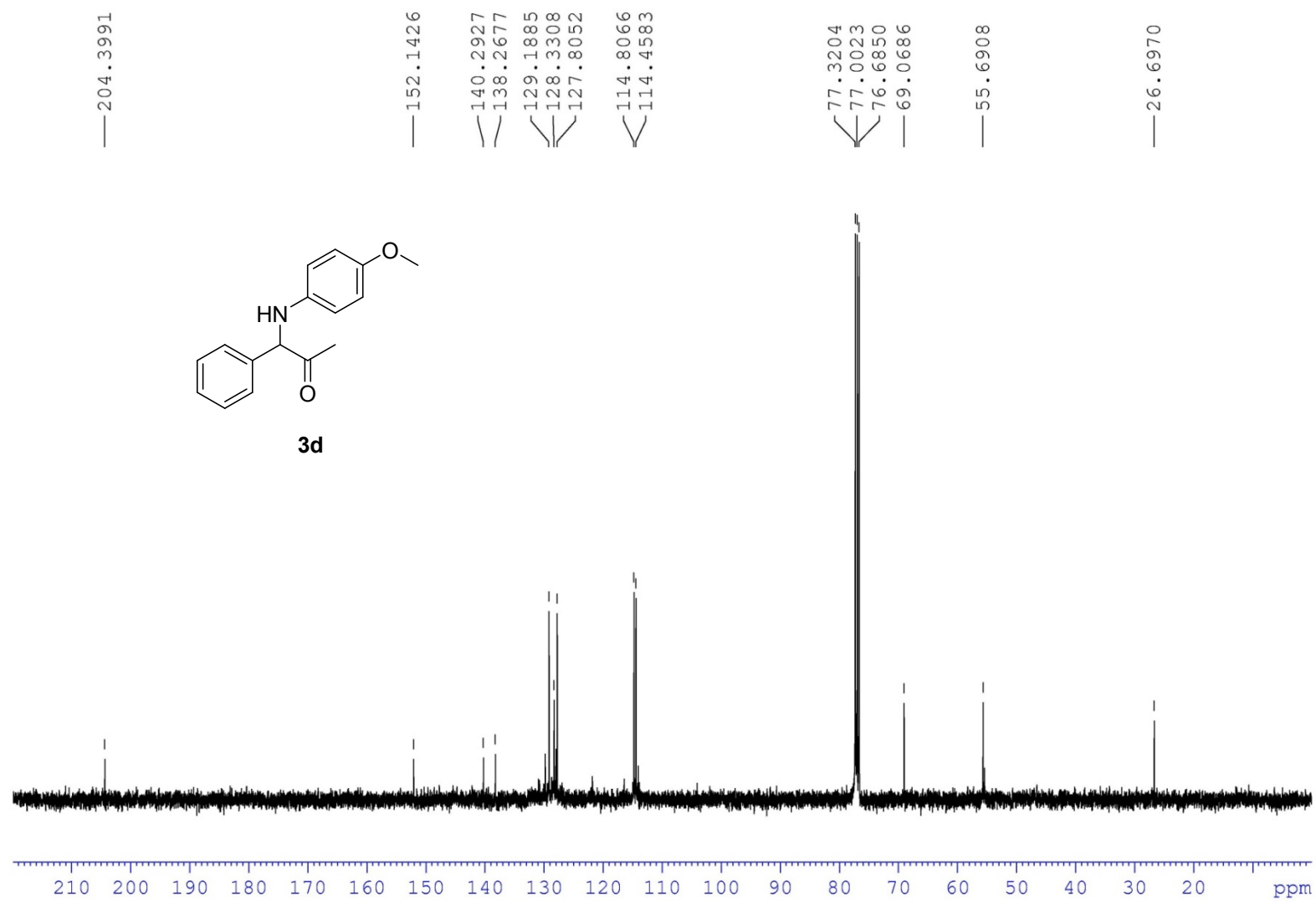
^{13}C NMR in CDCl_3 (100 MHz)



^1H NMR in CDCl_3 (300 MHz)



^{13}C NMR in CDCl_3 (100 MHz)



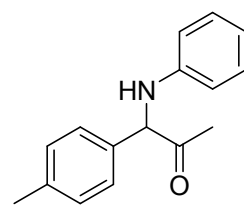
^1H NMR in CDCl_3 (400 MHz)

7.3308
7.3116
7.2599
7.1787
7.1593
7.1141
7.0931
7.0746
6.7079
6.6898
6.6720
6.6229
6.6050

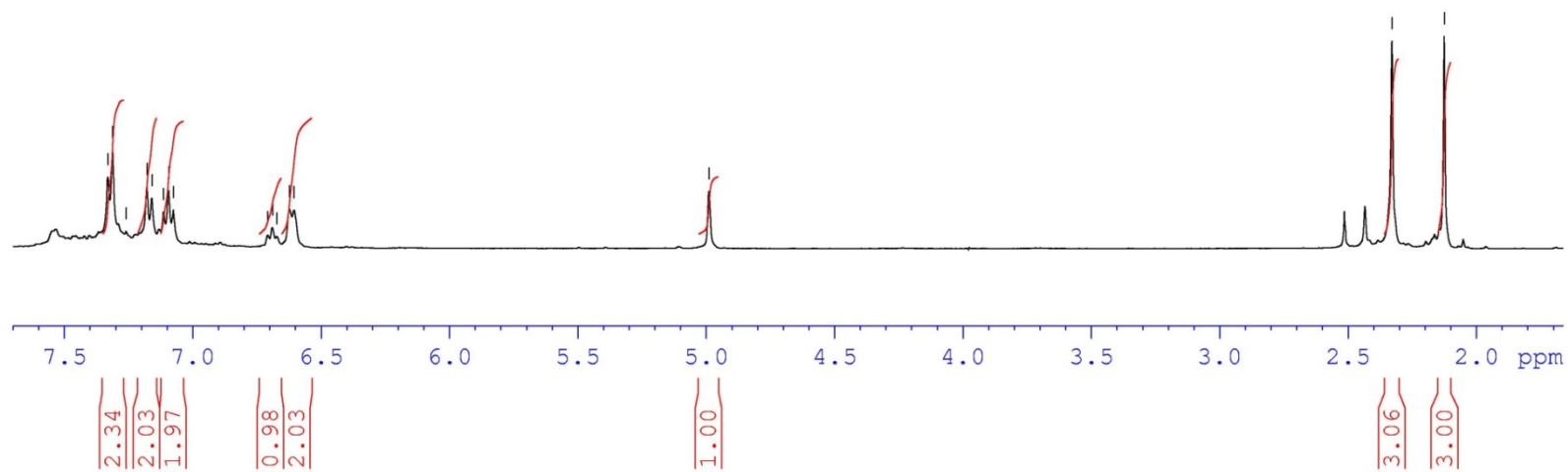
— 4.9891

— 2.3289

— 2.1254



3e



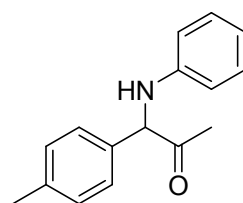
^{13}C NMR in CDCl_3 (75 MHz)

— 204.1632

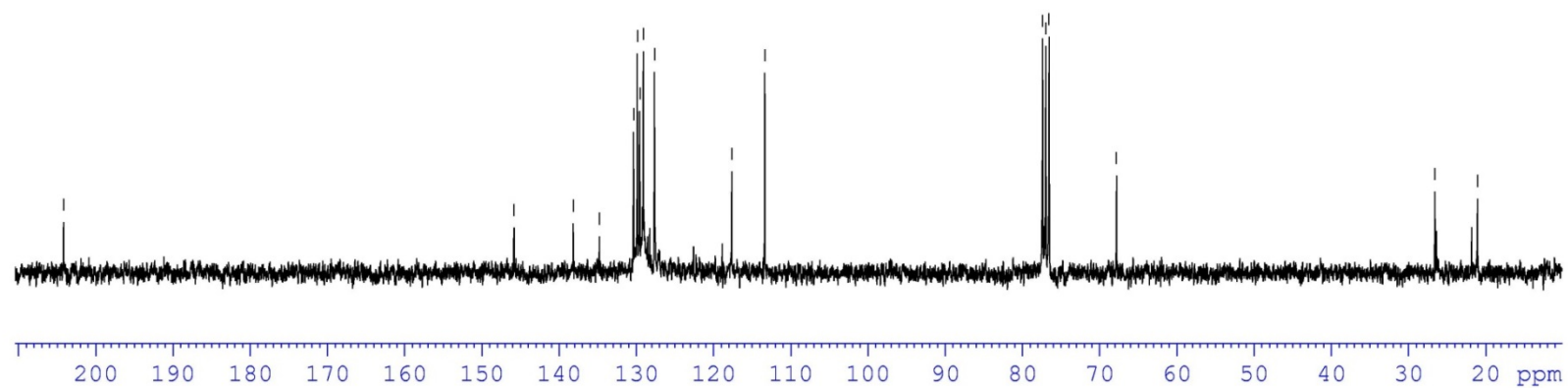
— 145.8416
 — 138.1569
 — 134.7725
 — 130.3597
 — 129.8625
 — 129.5409
 — 129.0750
 — 127.6382
 — 117.6286
 — 113.3487

77.4209
 — 76.9969
 — 76.5734
 — 67.8437

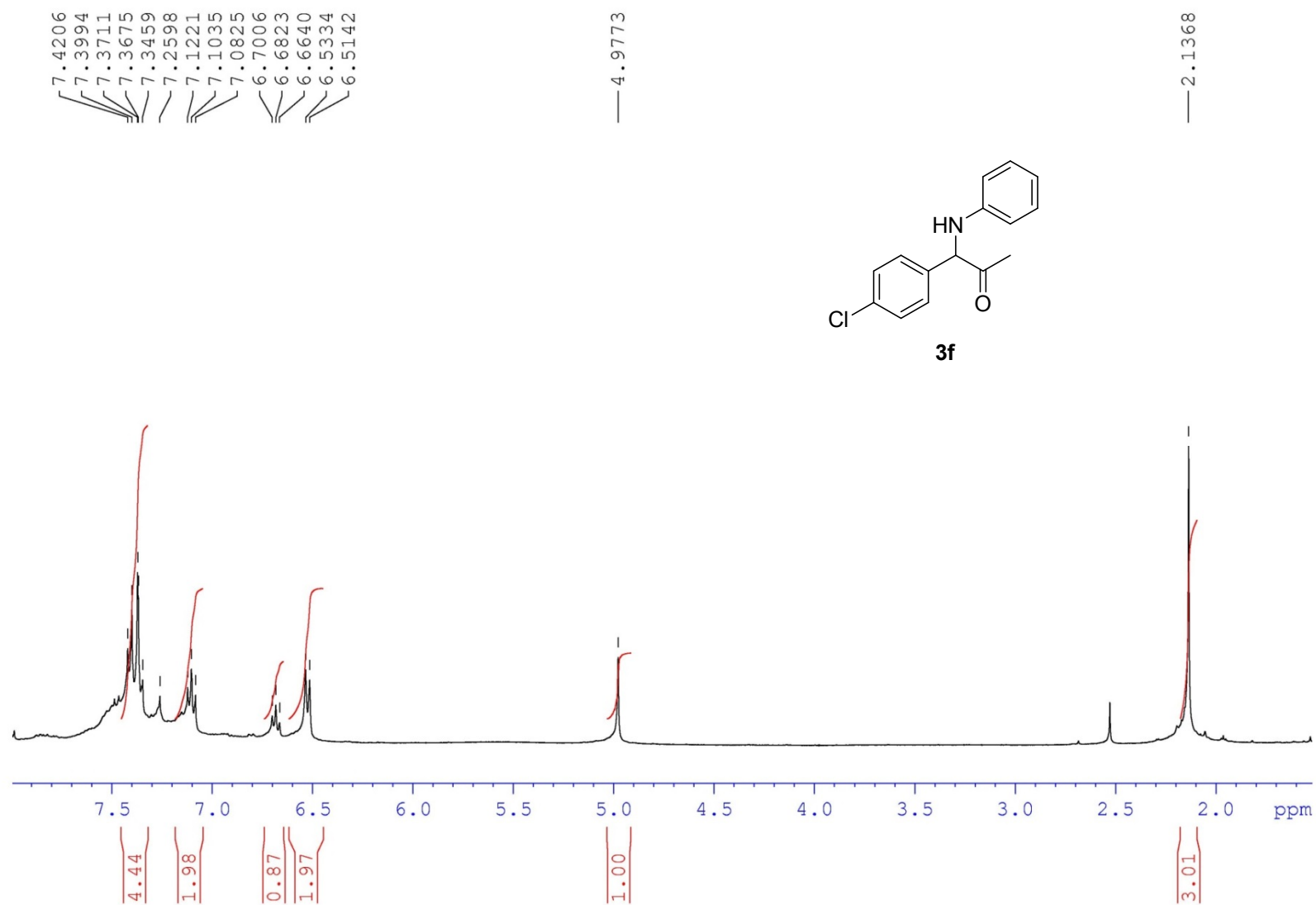
— 26.6015
 — 21.0867



3e



^1H NMR in CDCl_3 (400 MHz)



^{13}C NMR in CDCl_3 (100 MHz)

— 203.2103

— 145.5483

— 136.6251

— 131.7290

— 129.4252

— 129.1732

— 129.0631

— 117.9192

— 113.3207

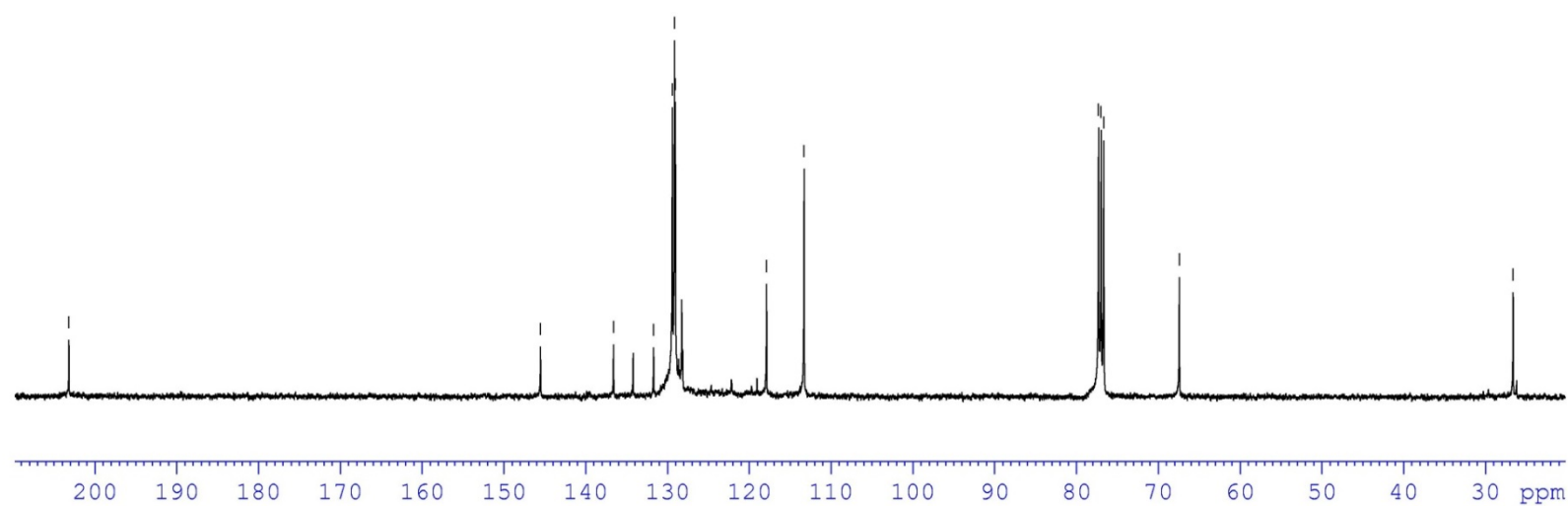
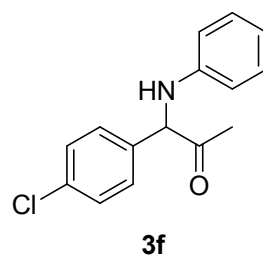
— 77.3185

— 77.0007

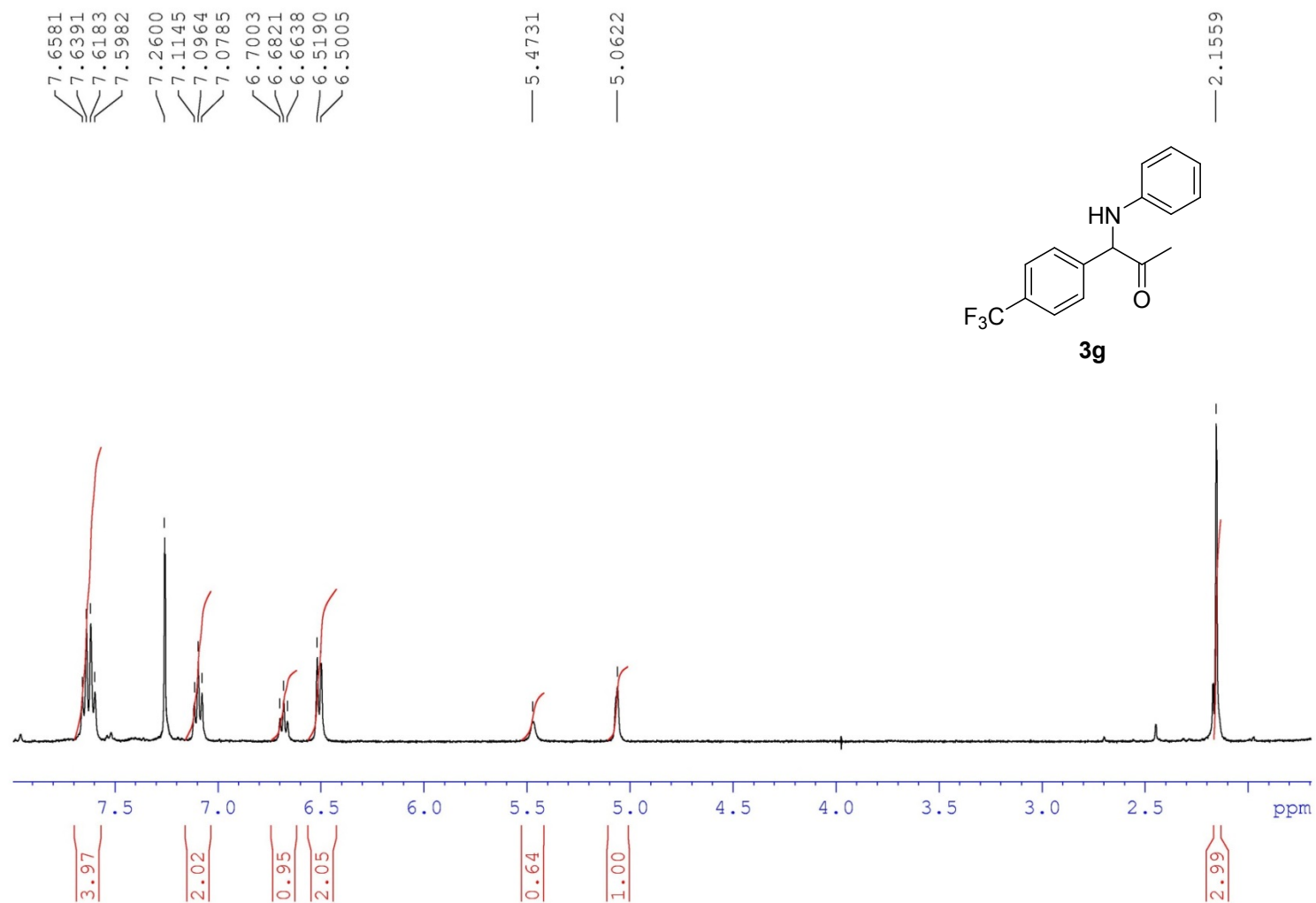
— 76.6829

— 67.4467

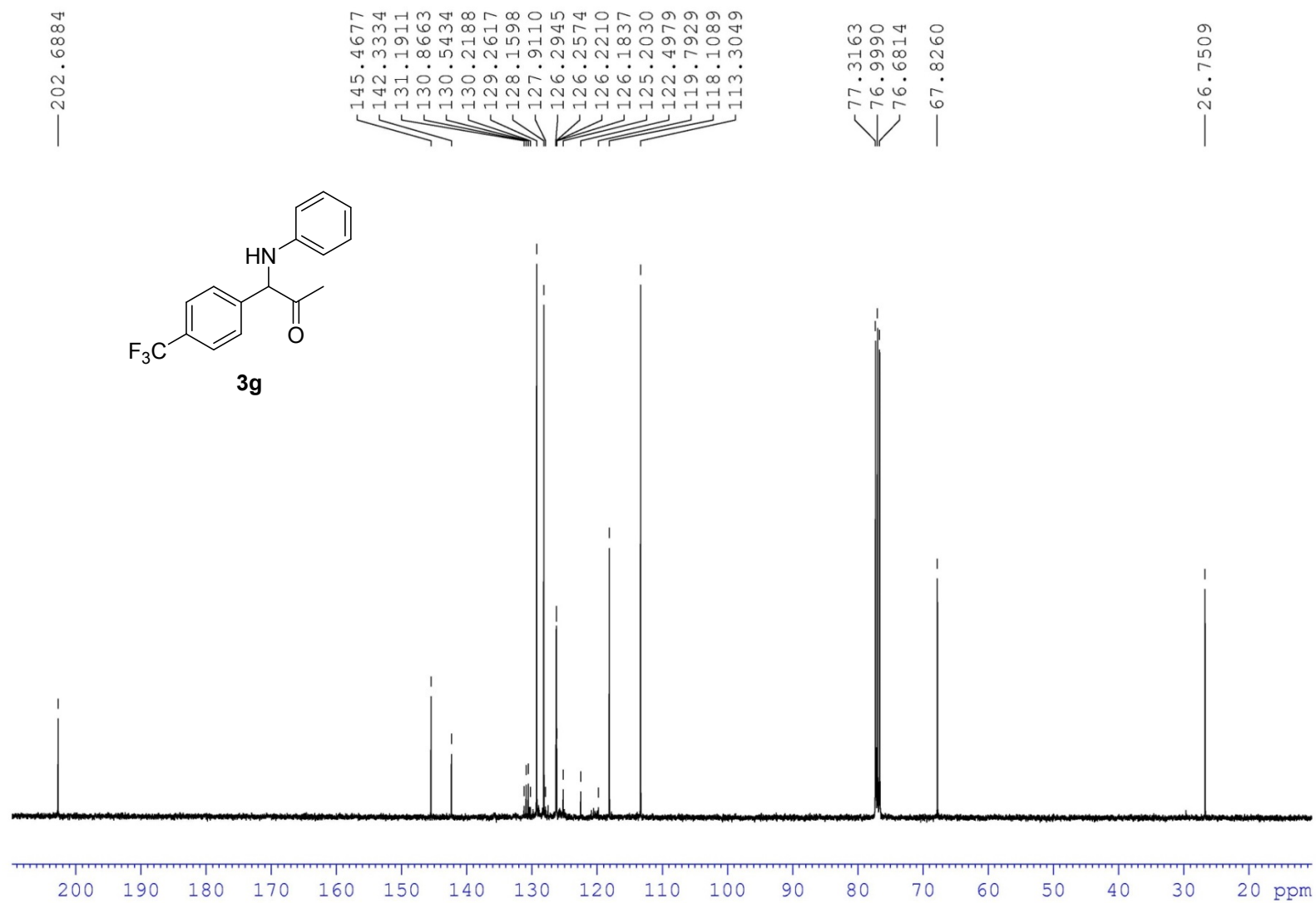
— 26.6397



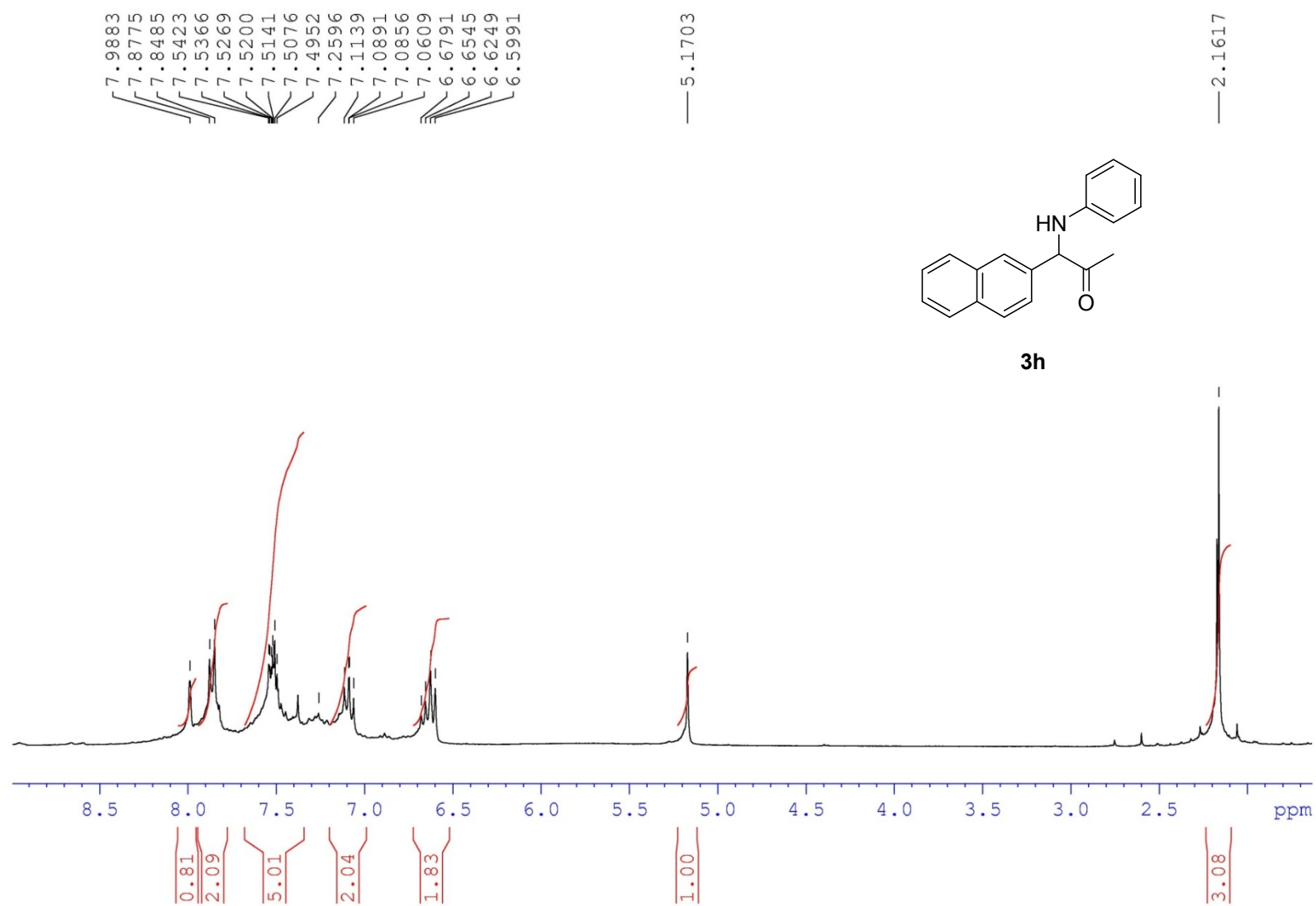
^1H NMR in CDCl_3 (400 MHz)



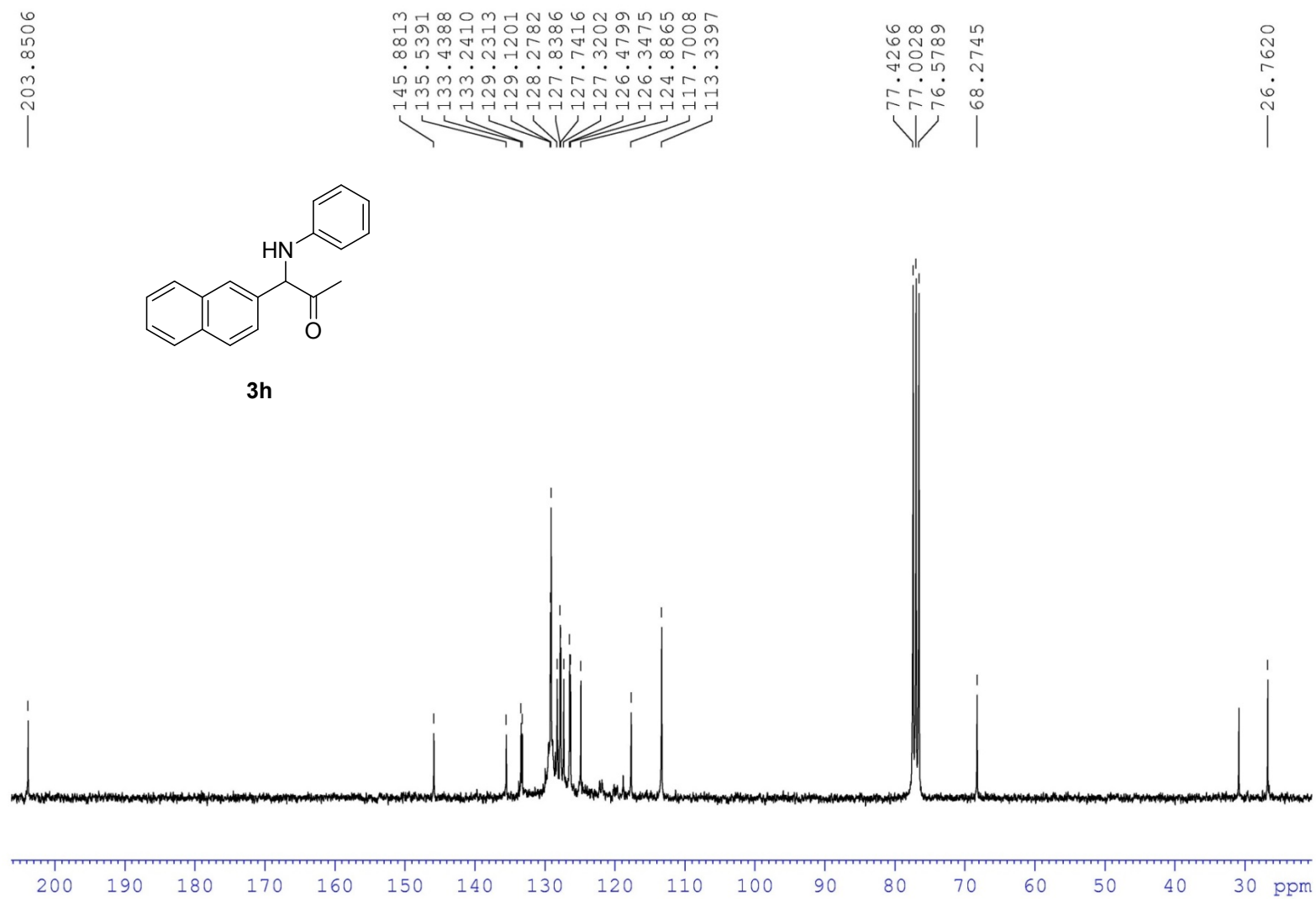
^{13}C NMR in CDCl_3 (100 MHz)



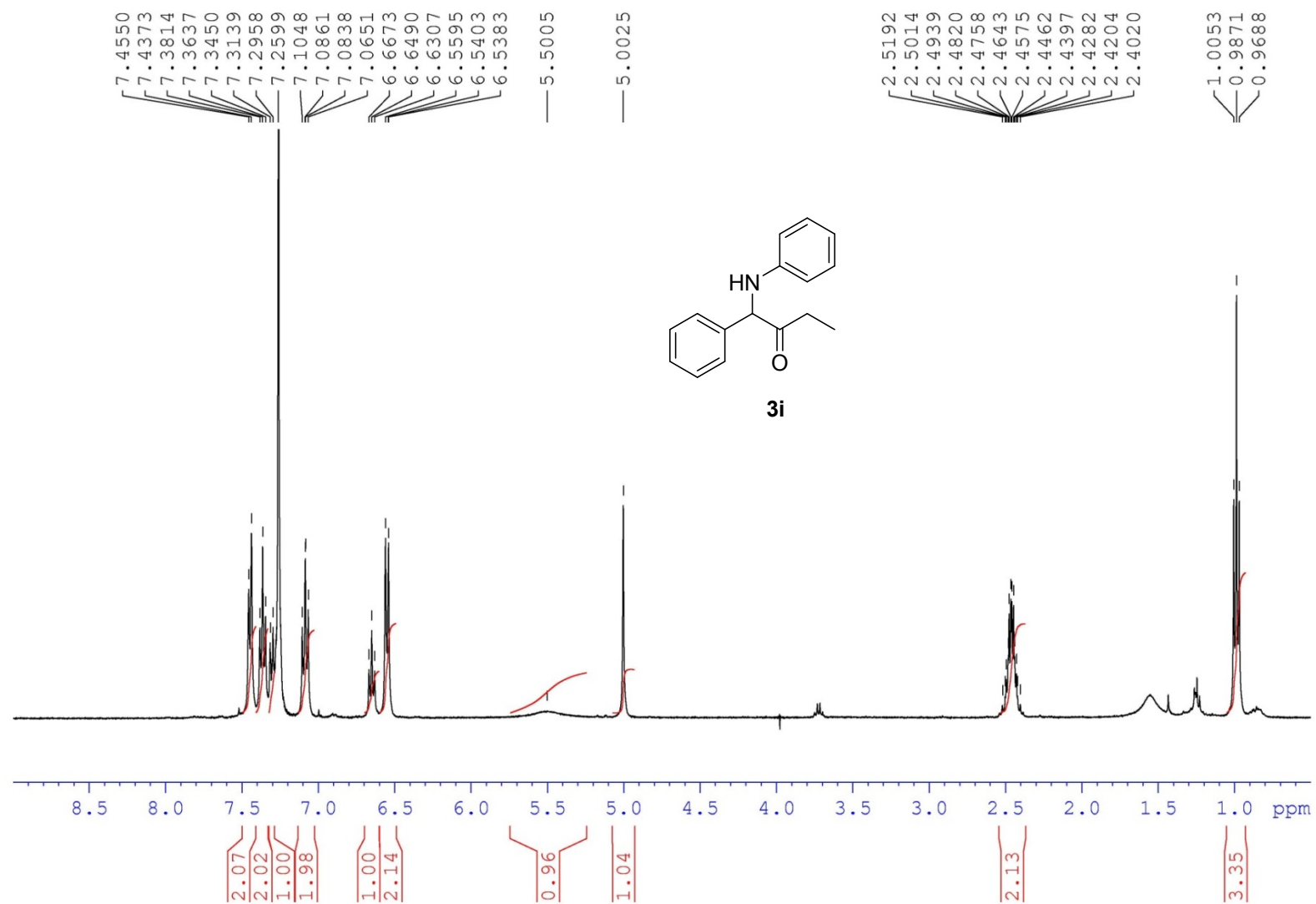
^1H NMR in CDCl_3 (300 MHz)



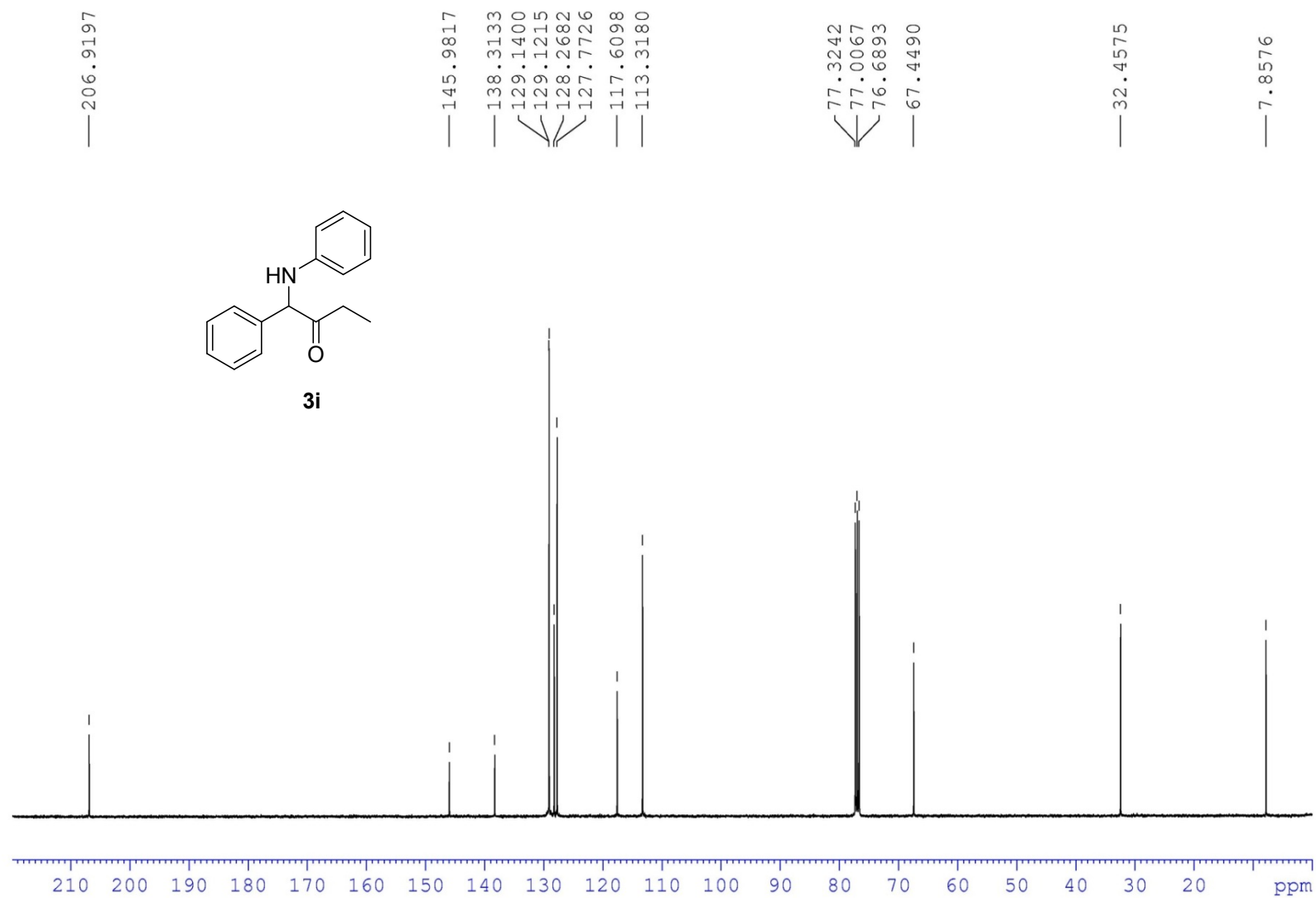
^{13}C NMR in CDCl_3 (75 MHz)



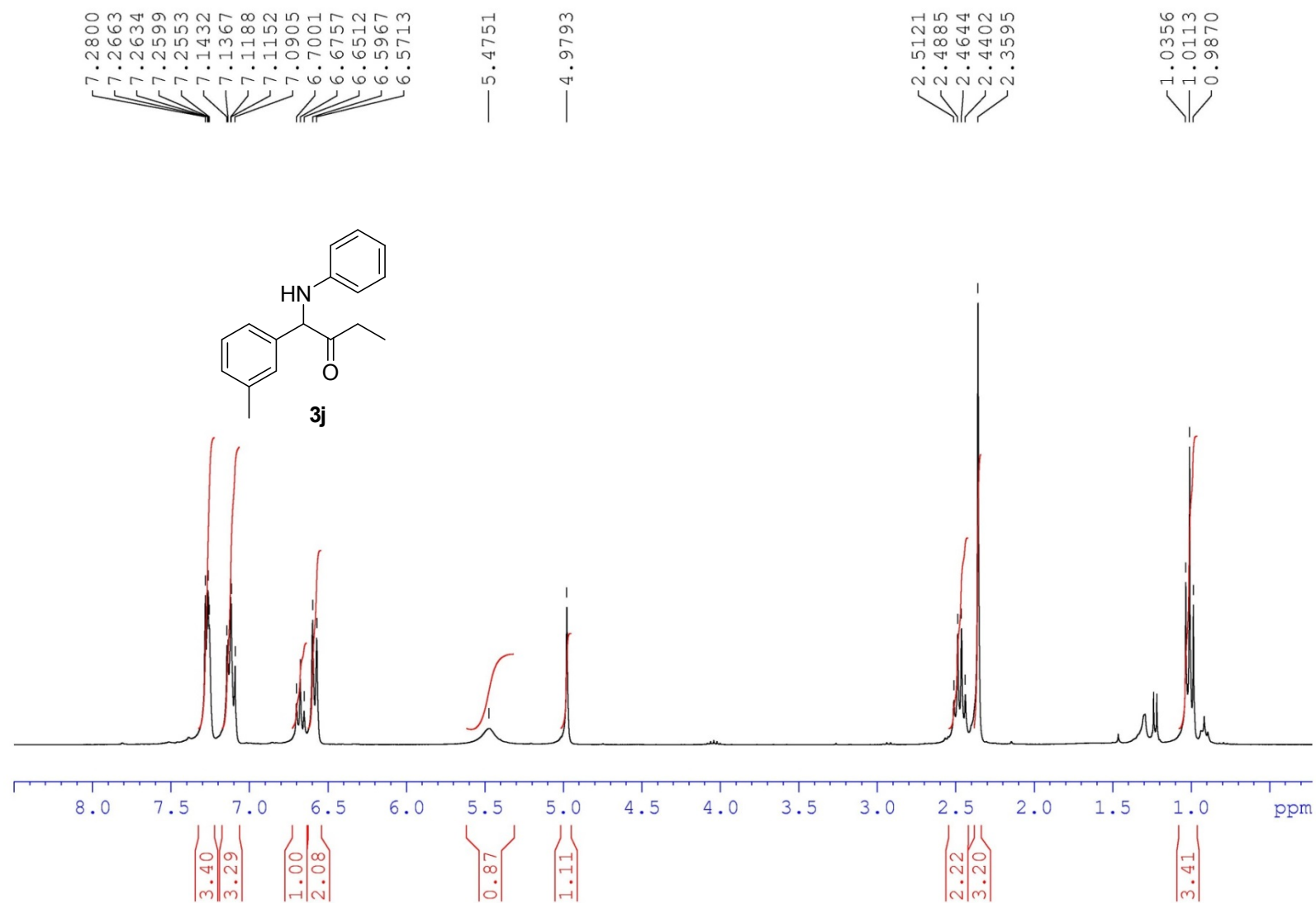
^1H NMR in CDCl_3 (400 MHz)



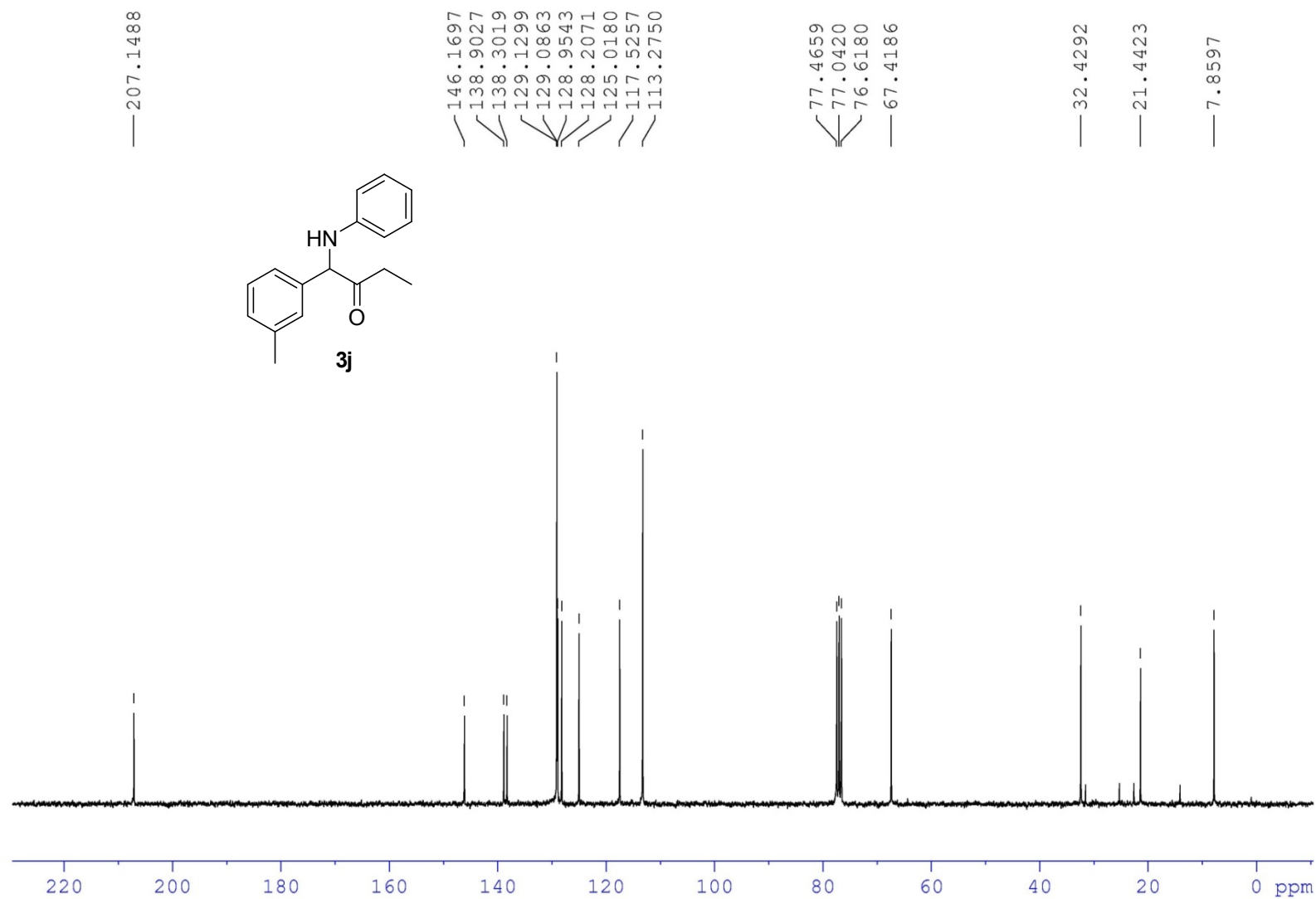
^{13}C NMR in CDCl_3 (100 MHz)



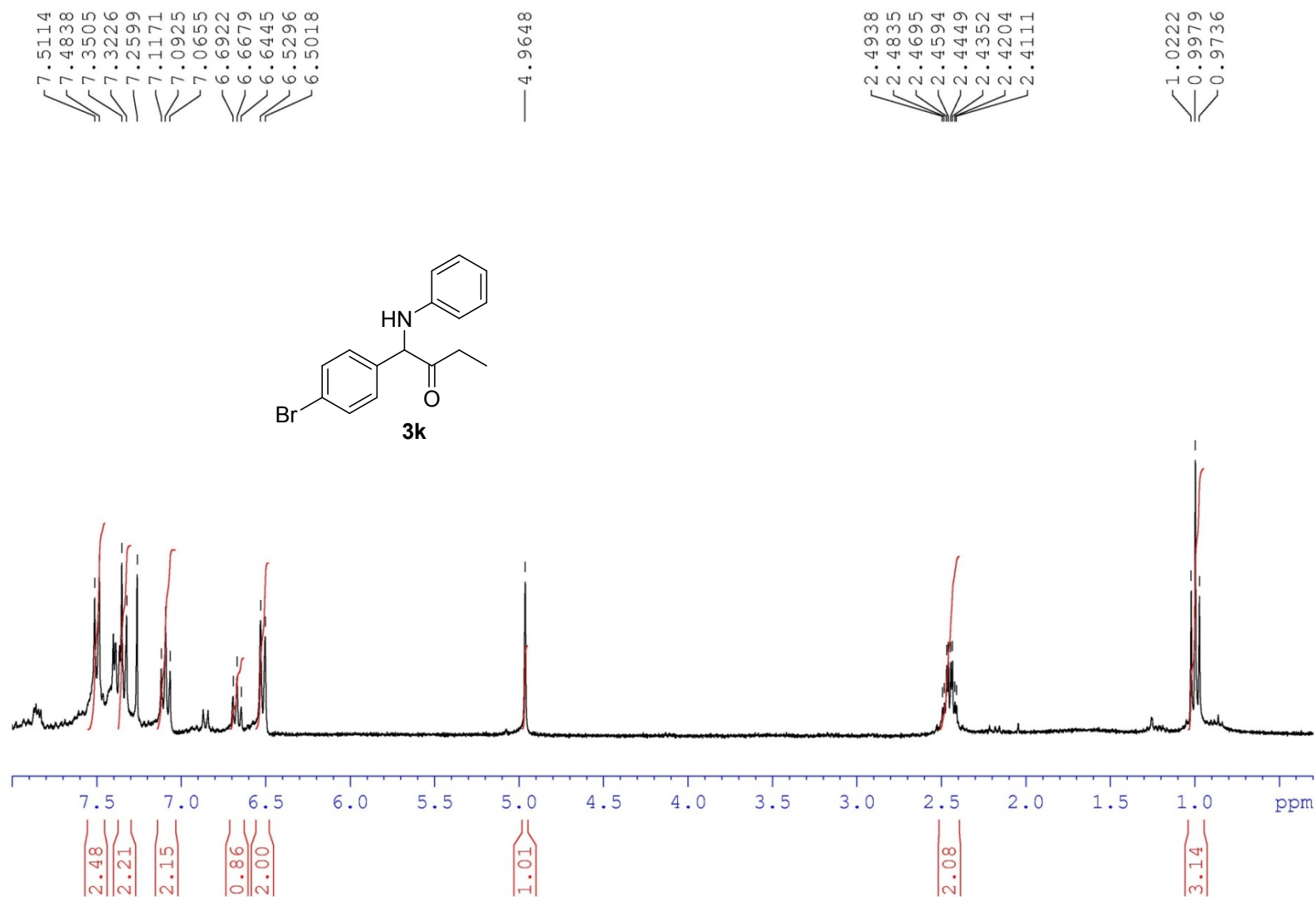
^1H NMR in CDCl_3 (300 MHz)



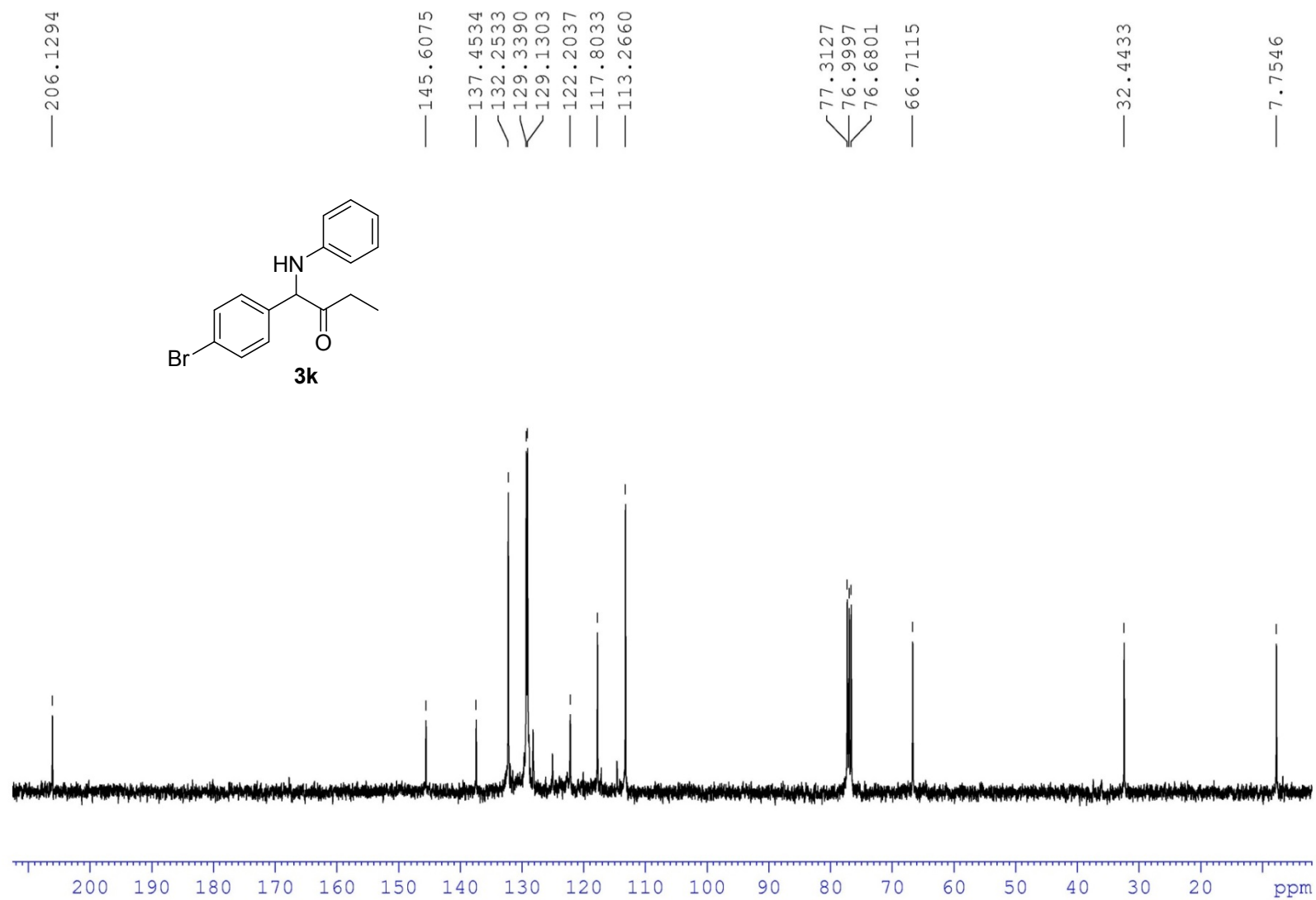
^{13}C NMR in CDCl_3 (75 MHz)



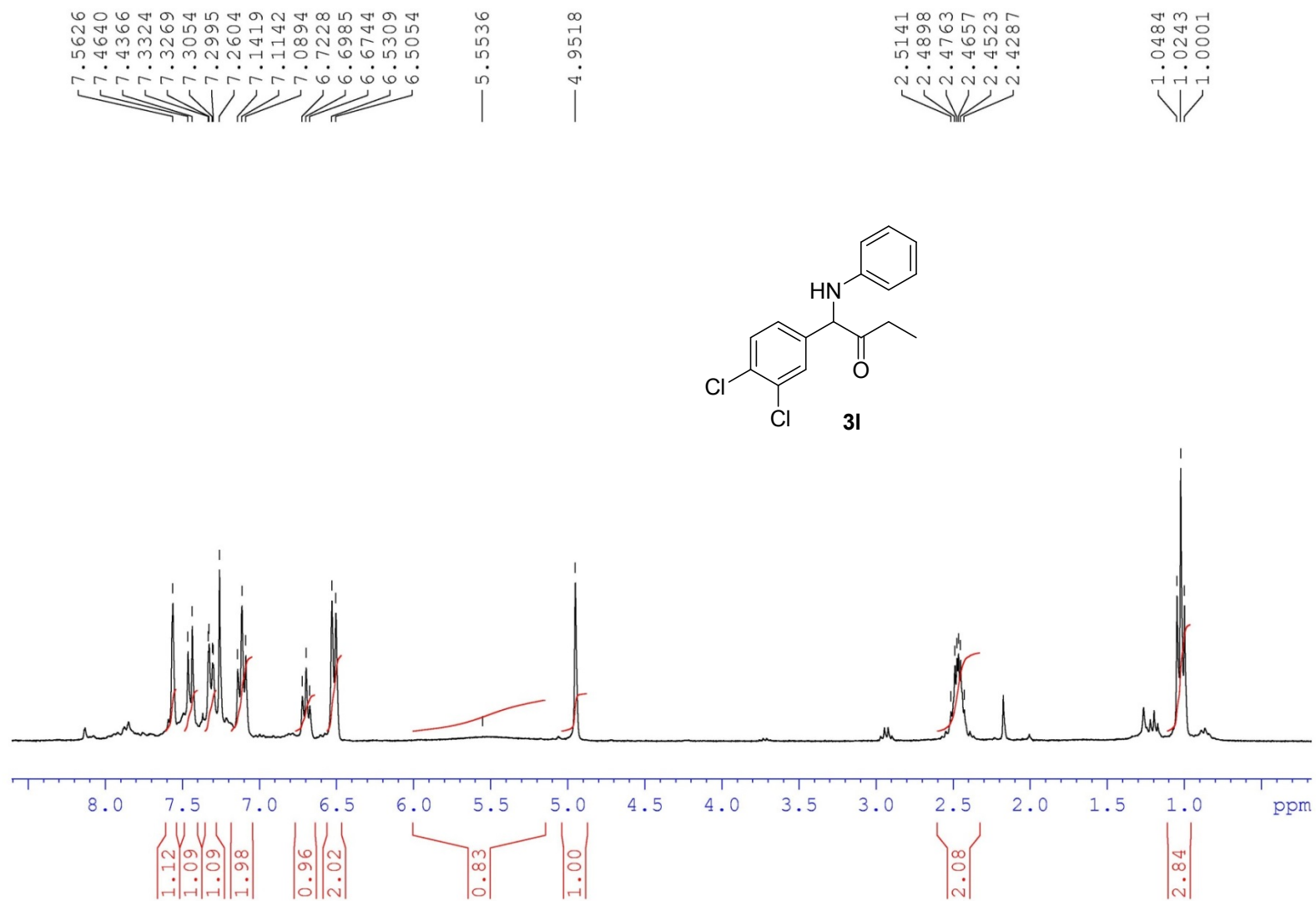
^1H NMR in CDCl_3 (300 MHz)



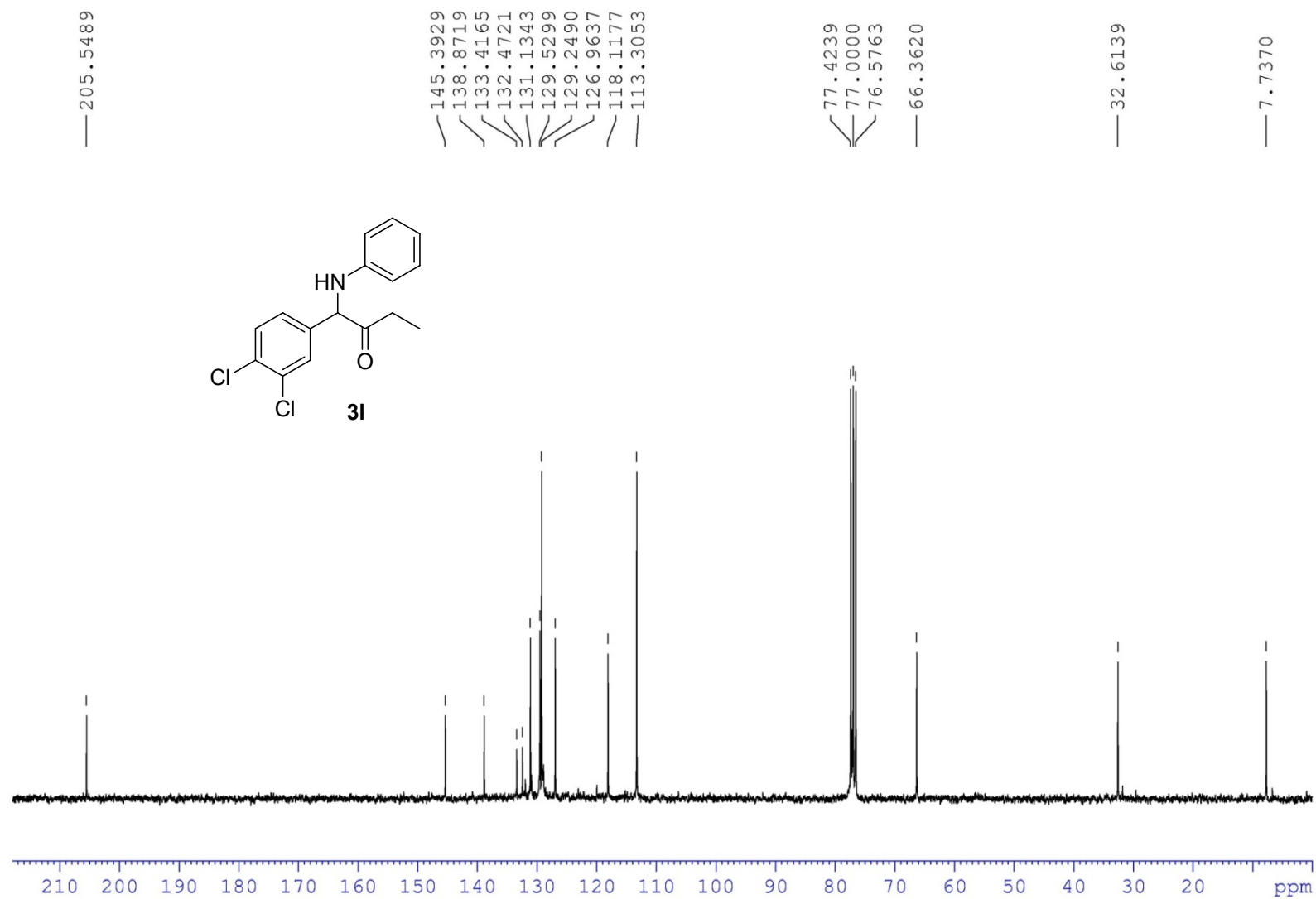
^{13}C NMR in CDCl_3 (100 MHz)



^1H NMR in CDCl_3 (300 MHz)



^{13}C NMR in CDCl_3 (75 MHz)

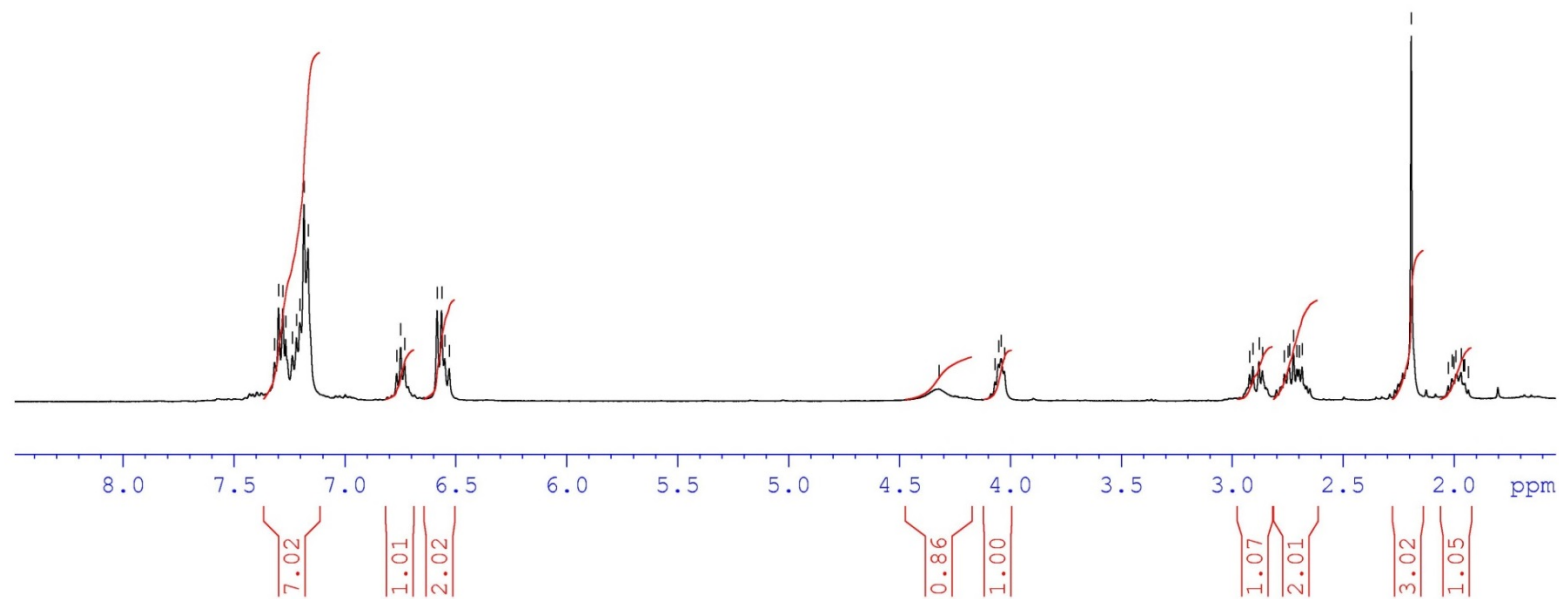
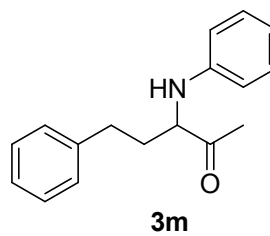


^1H NMR in CDCl_3 (400 MHz)

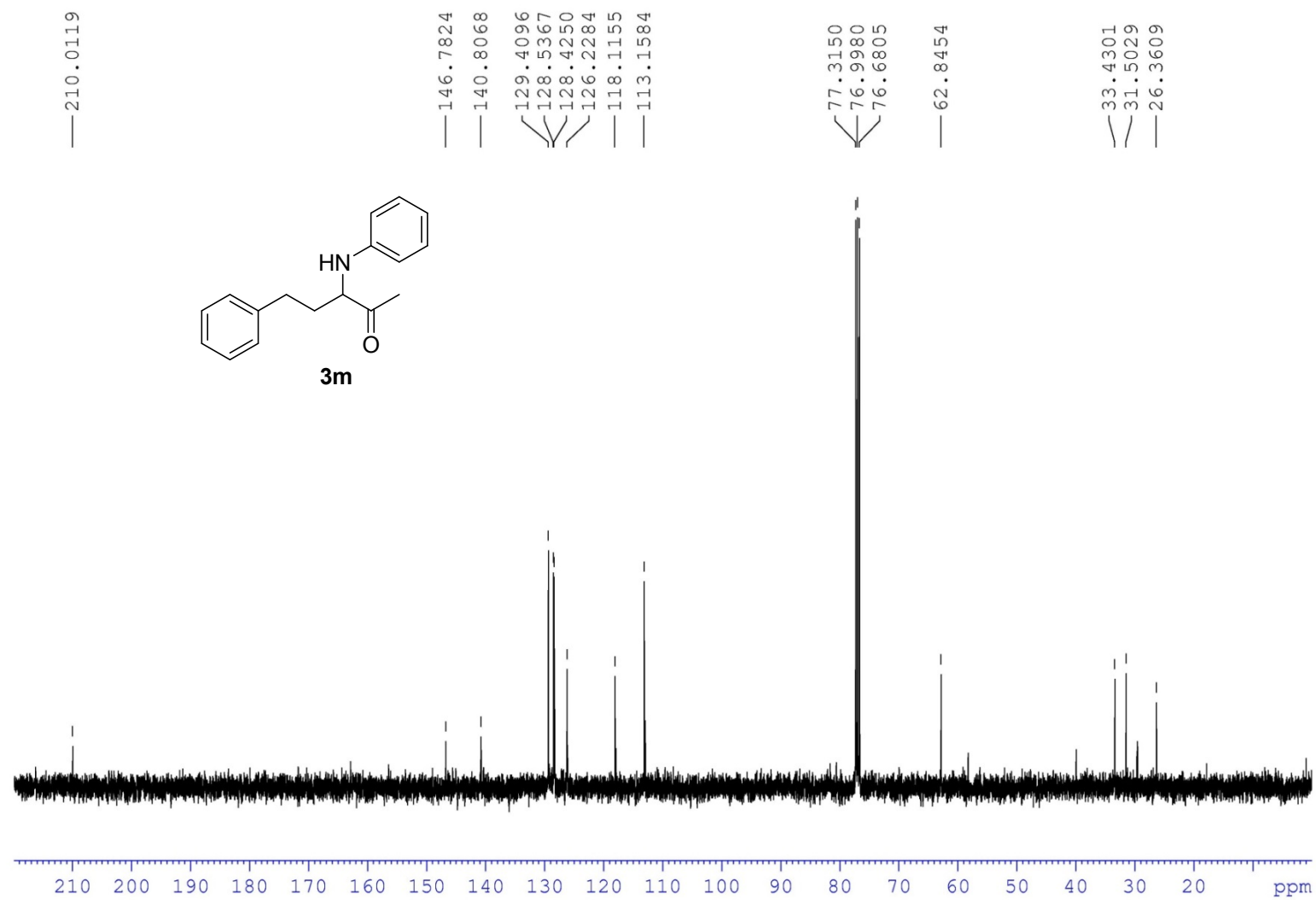
7.3167
7.2985
7.2797
7.2670
7.2361
7.2180
7.2029
7.1839
7.1655
6.7658
6.7483
6.7309
6.5839
6.5637
6.5496
6.5288

4.3219
4.0695
4.0534
4.0411
4.0270

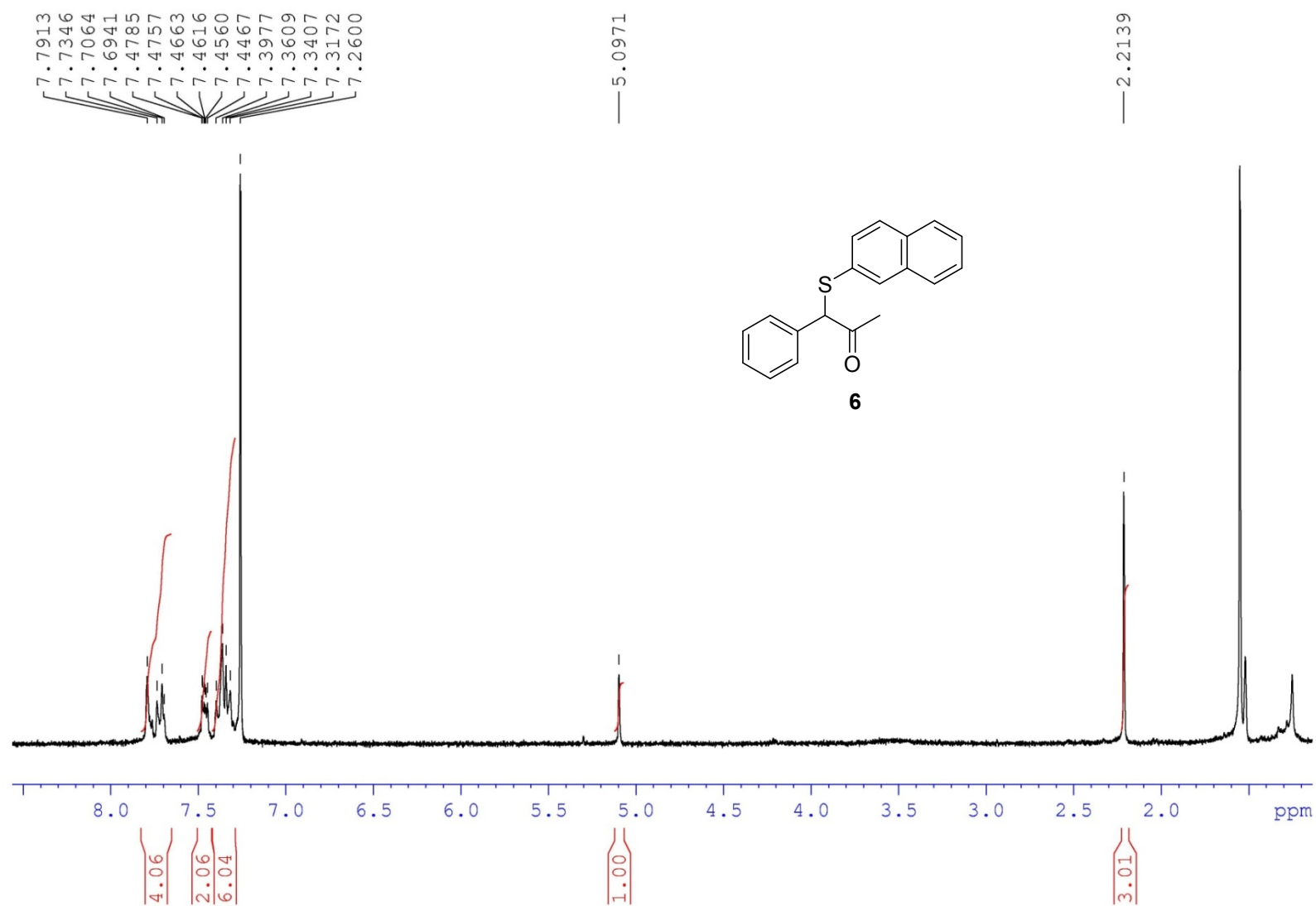
2.9207
2.9064
2.8790
2.8634
2.7646
2.7477
2.7416
2.7232
2.7080
2.6984
2.6844
2.1926
2.0257
2.0086
2.0021
1.9912



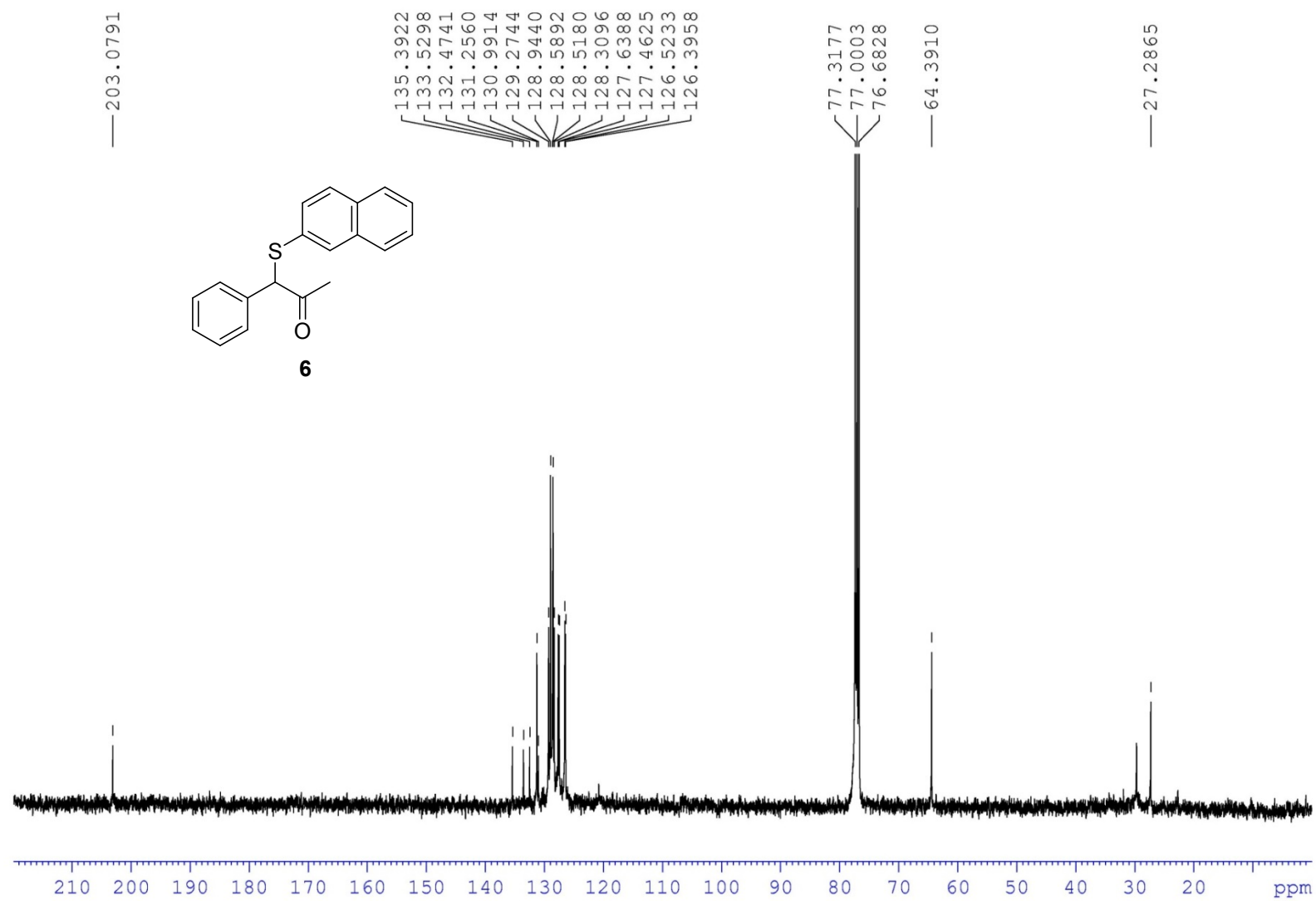
^{13}C NMR in CDCl_3 (100 MHz)



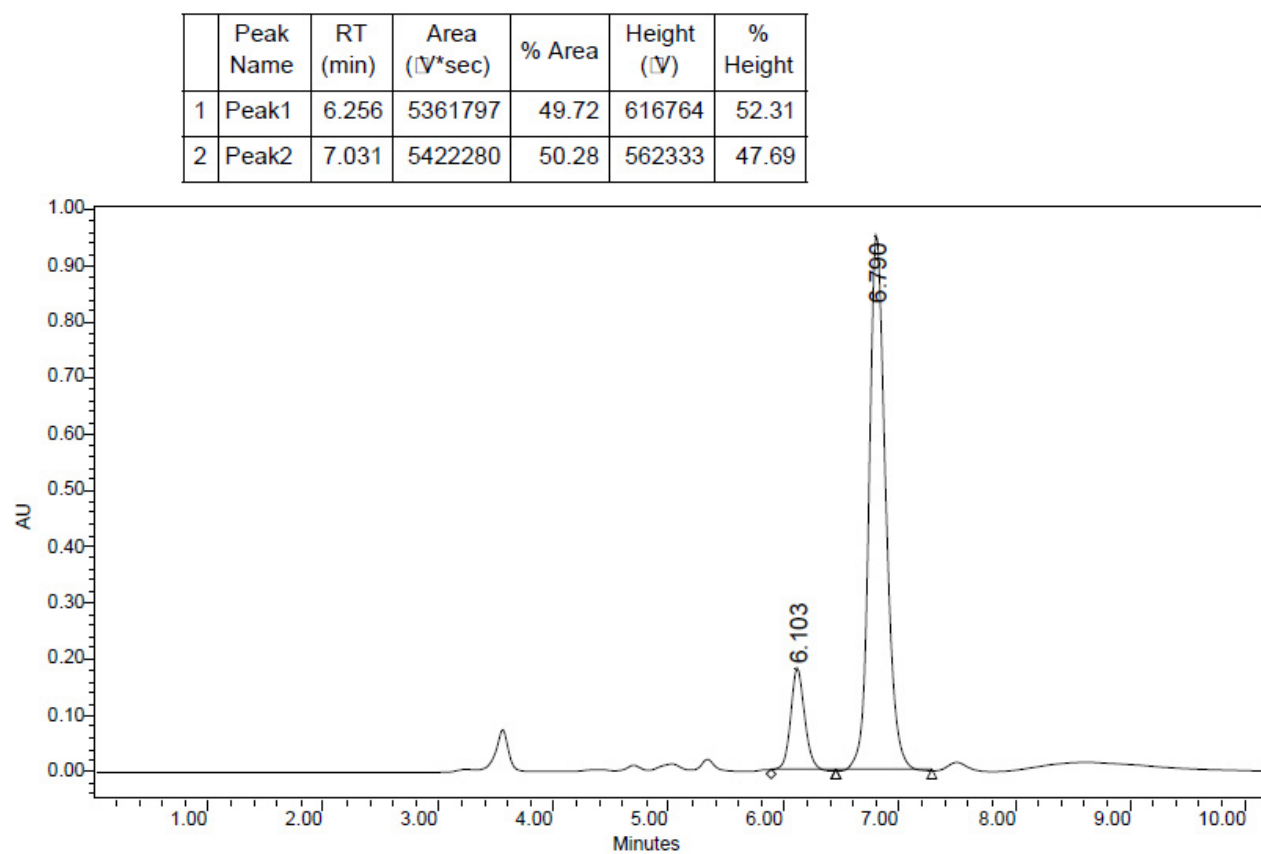
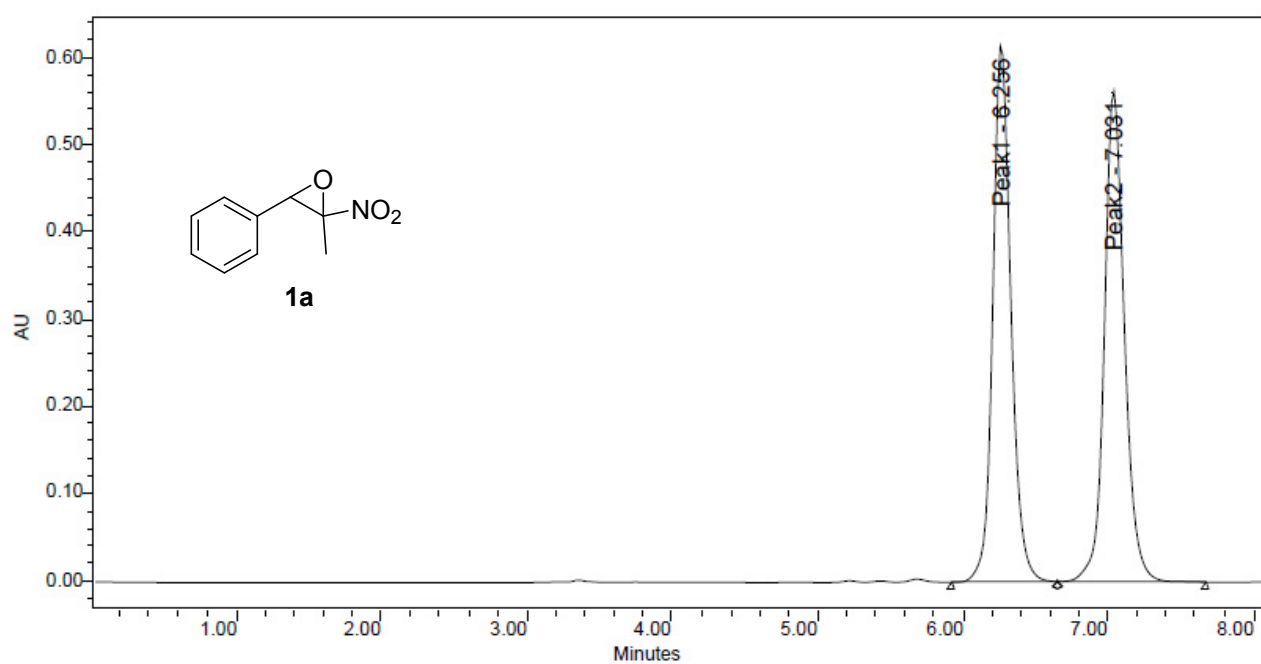
^1H NMR in CDCl_3 (300 MHz)

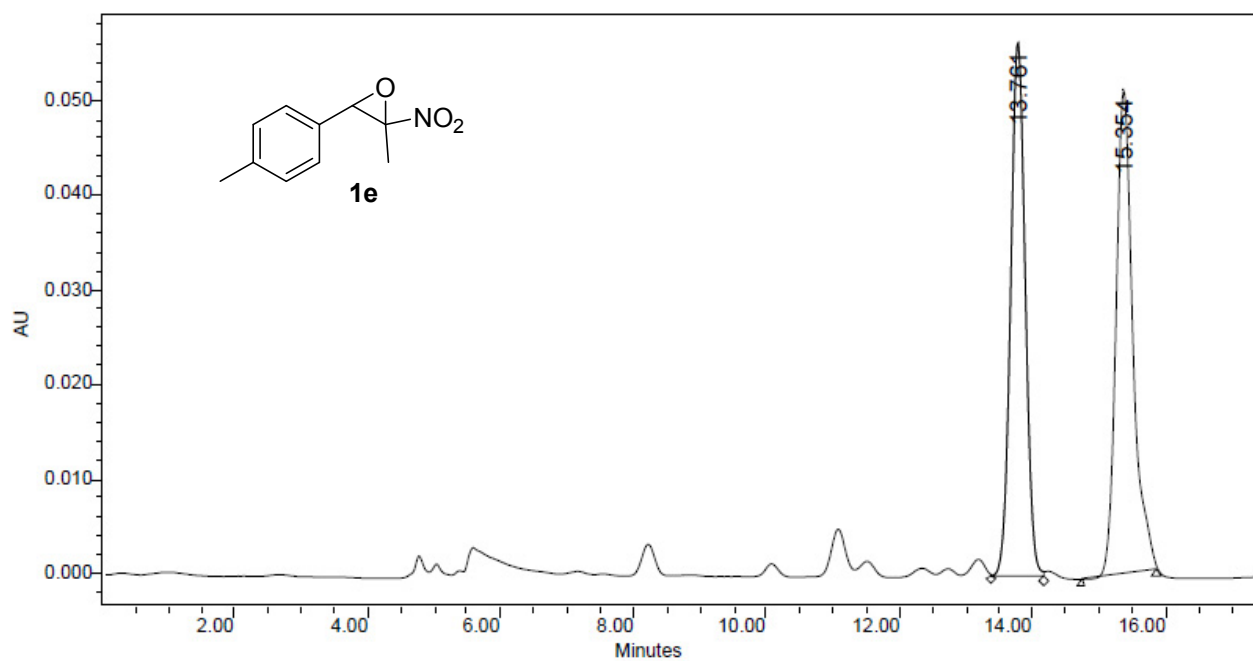


^{13}C NMR in CDCl_3 (100 MHz)

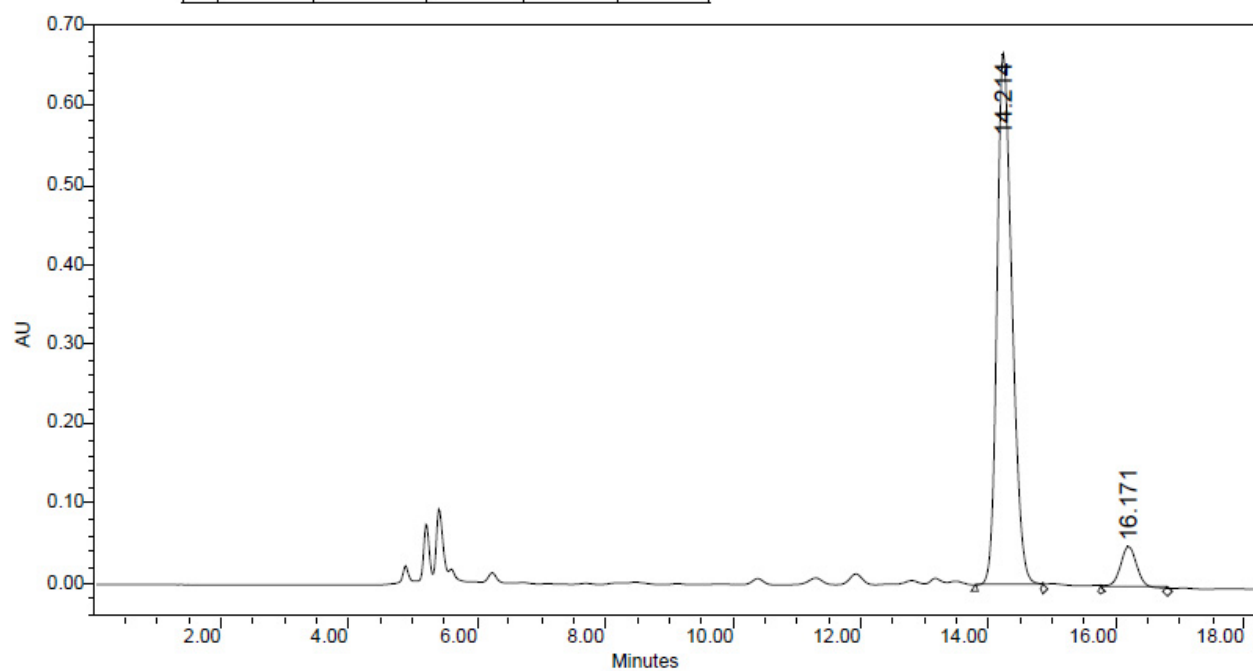


HPLC Chromatograms

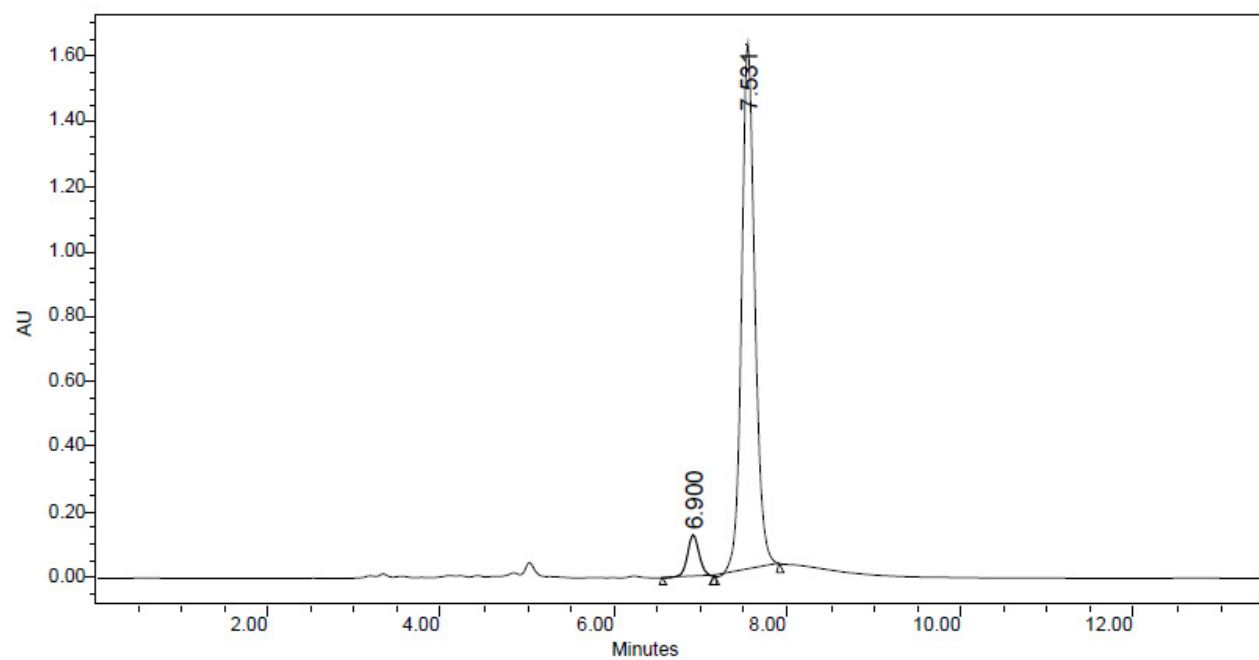
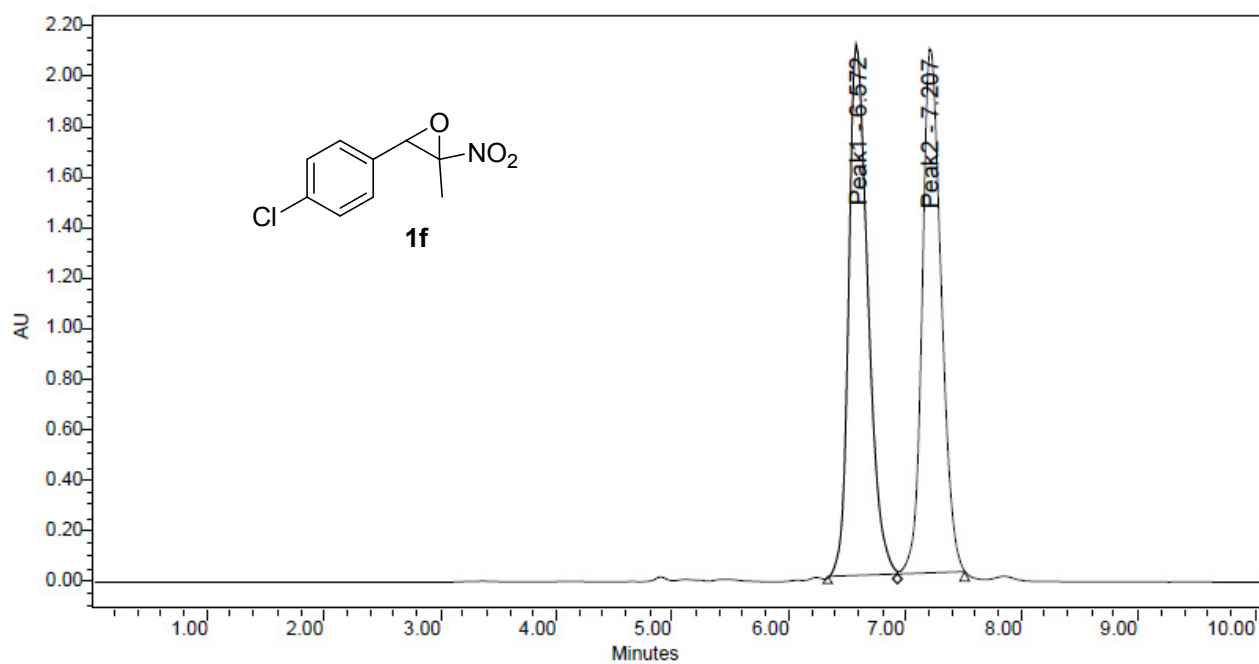


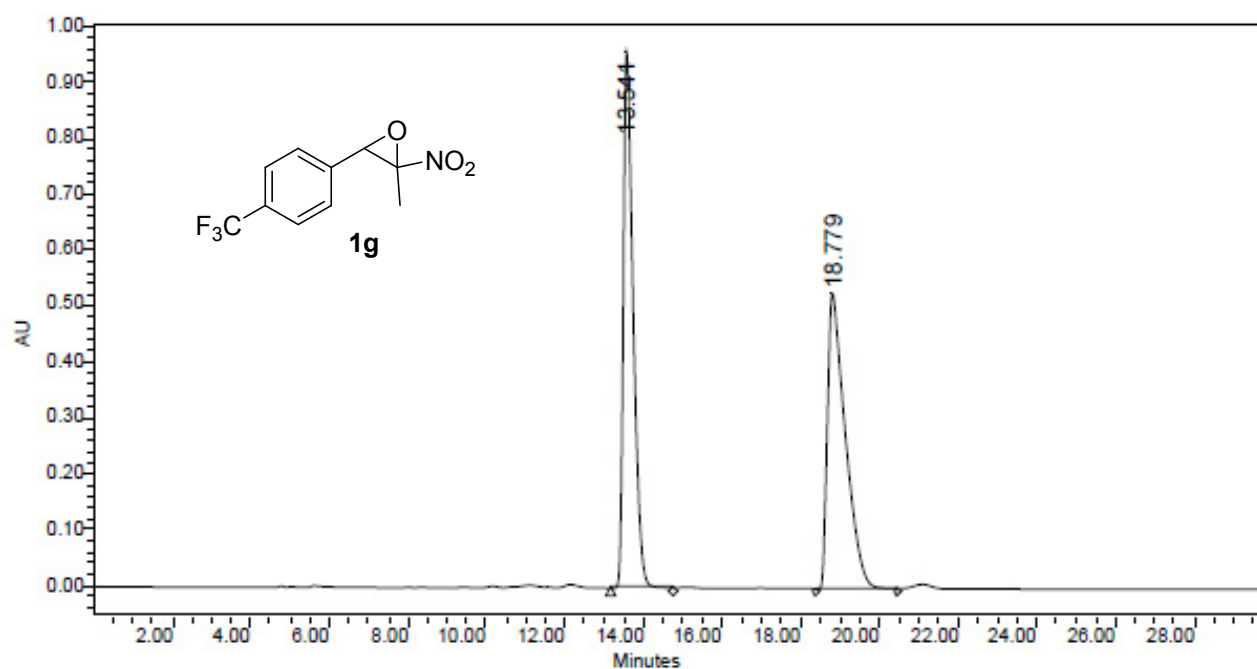


	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	13.761	864135	49.01	56472	52.53
2	15.354	899101	50.99	51032	47.47

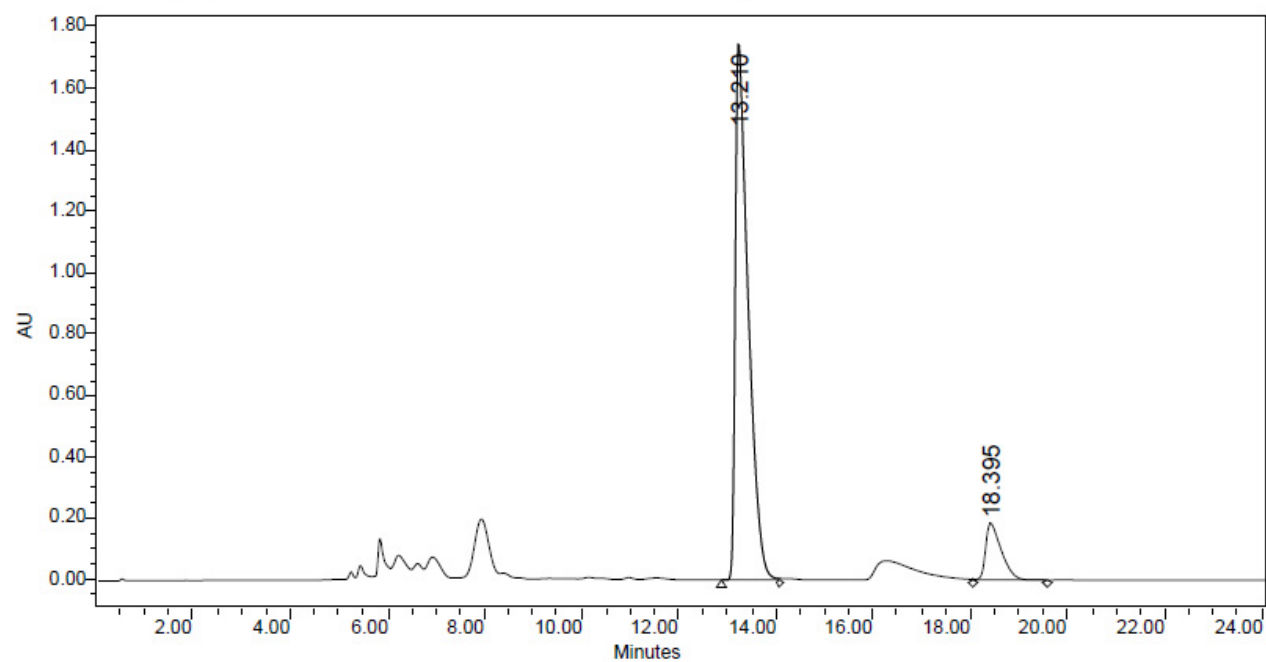


	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	14.214	10970925	92.06	667355	92.80
2	16.171	945669	7.94	51757	7.20

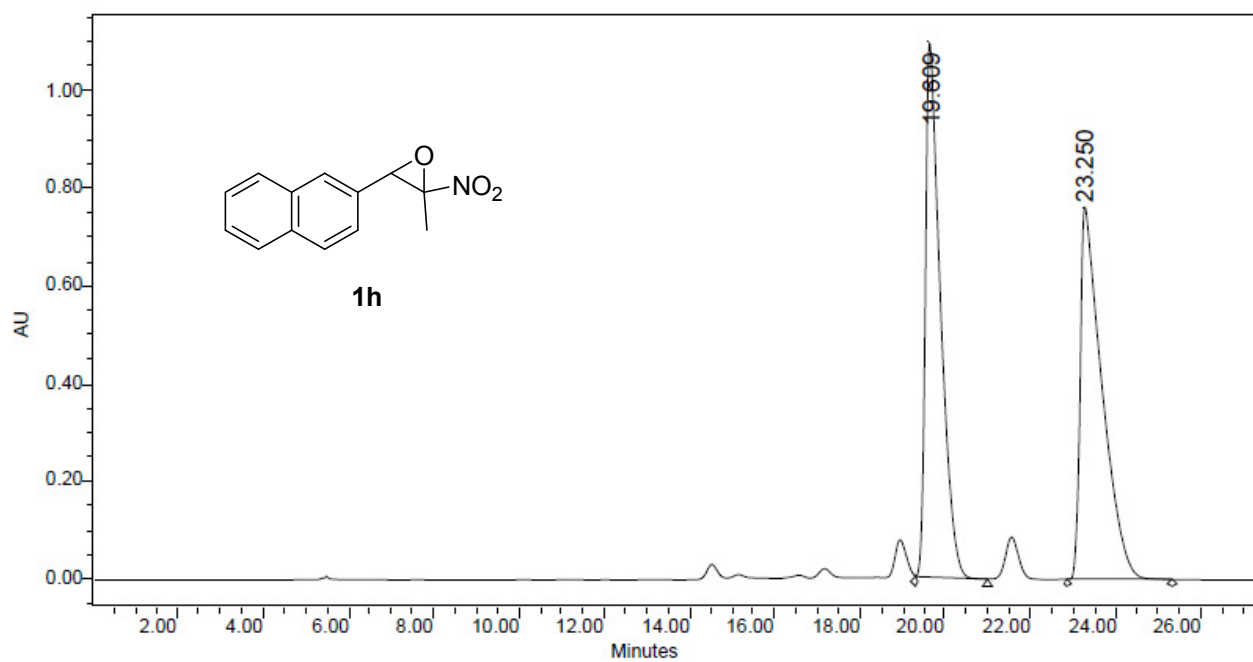




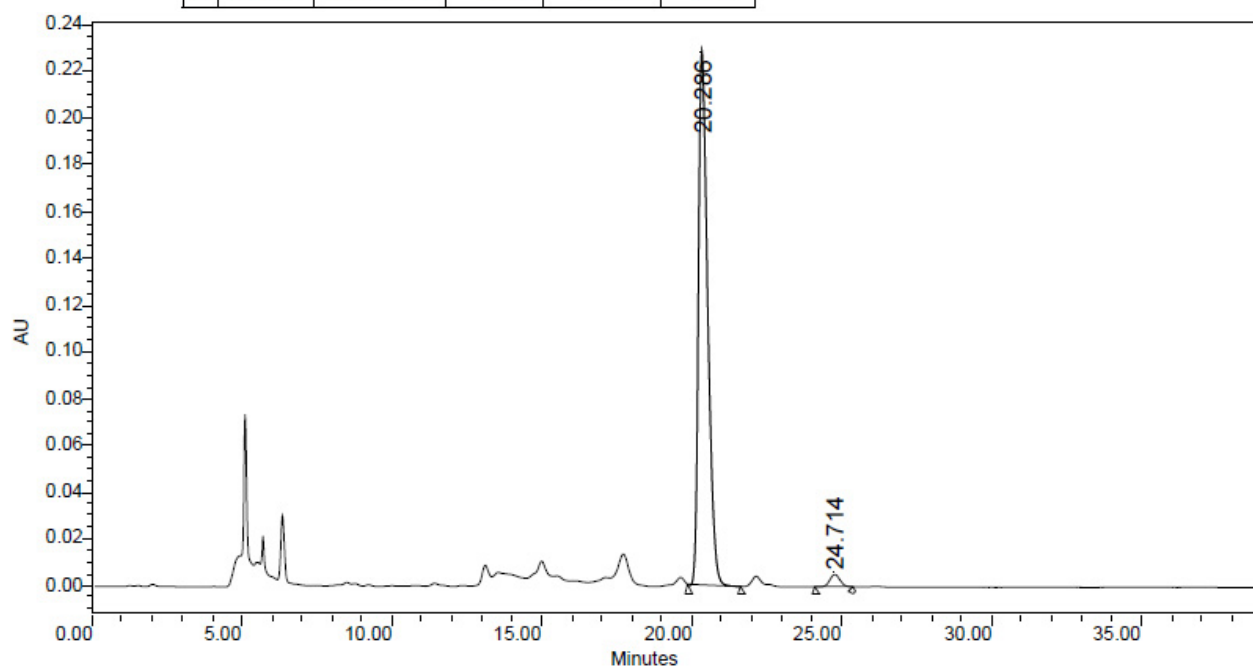
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	13.541	16667643	50.01	962702	64.58
2	18.779	16663753	49.99	528055	35.42



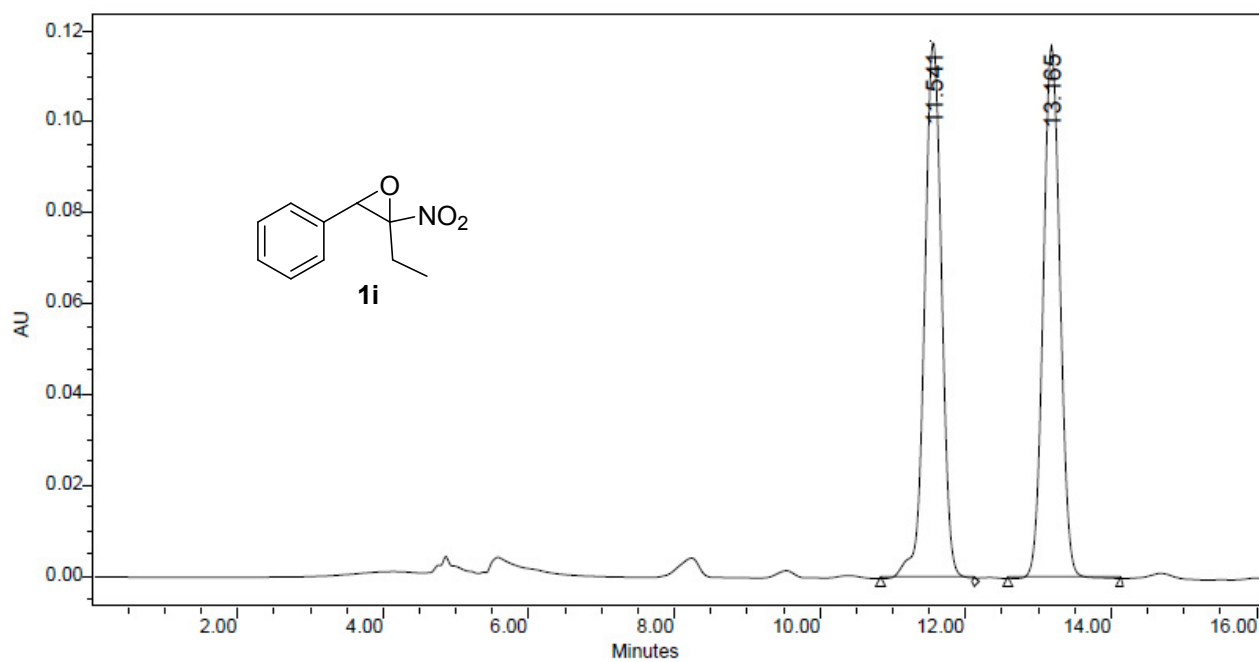
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	13.210	32134640	88.53	1756444	90.47
2	18.395	4163943	11.47	185022	9.53



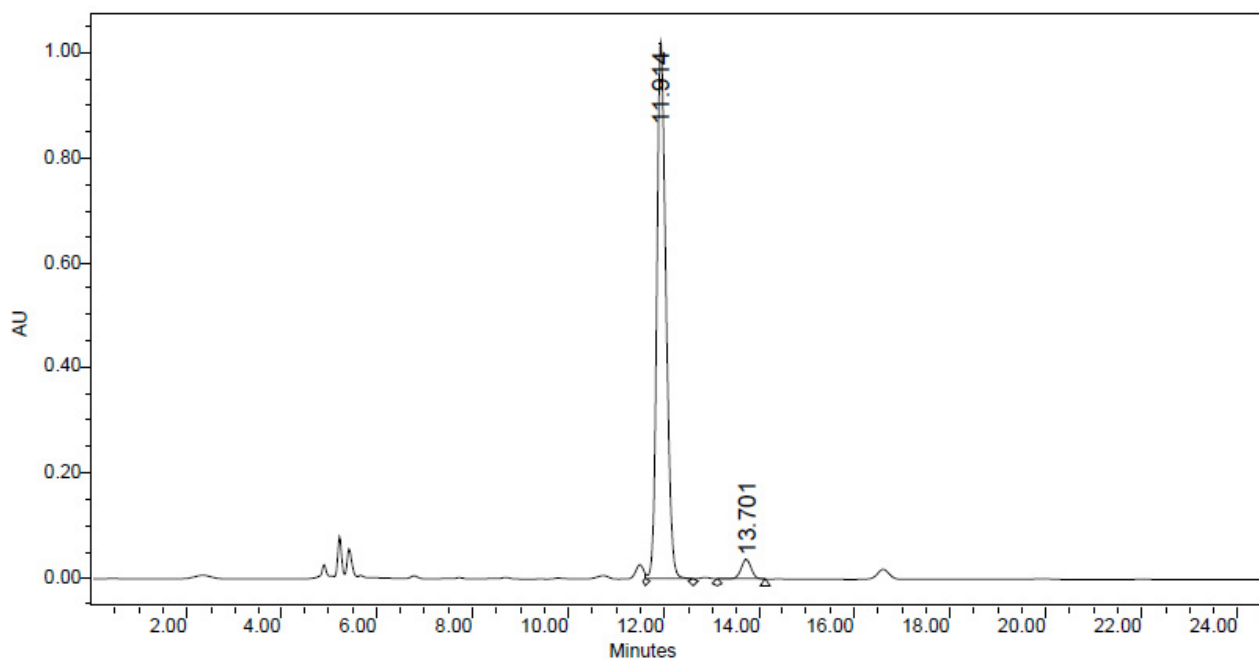
	RT (min)	Area (AU*sec)	% Area	Height (AU)	% Height
1	19.609	26936455	49.84	1099274	59.03
2	23.250	27112984	50.16	762846	40.97



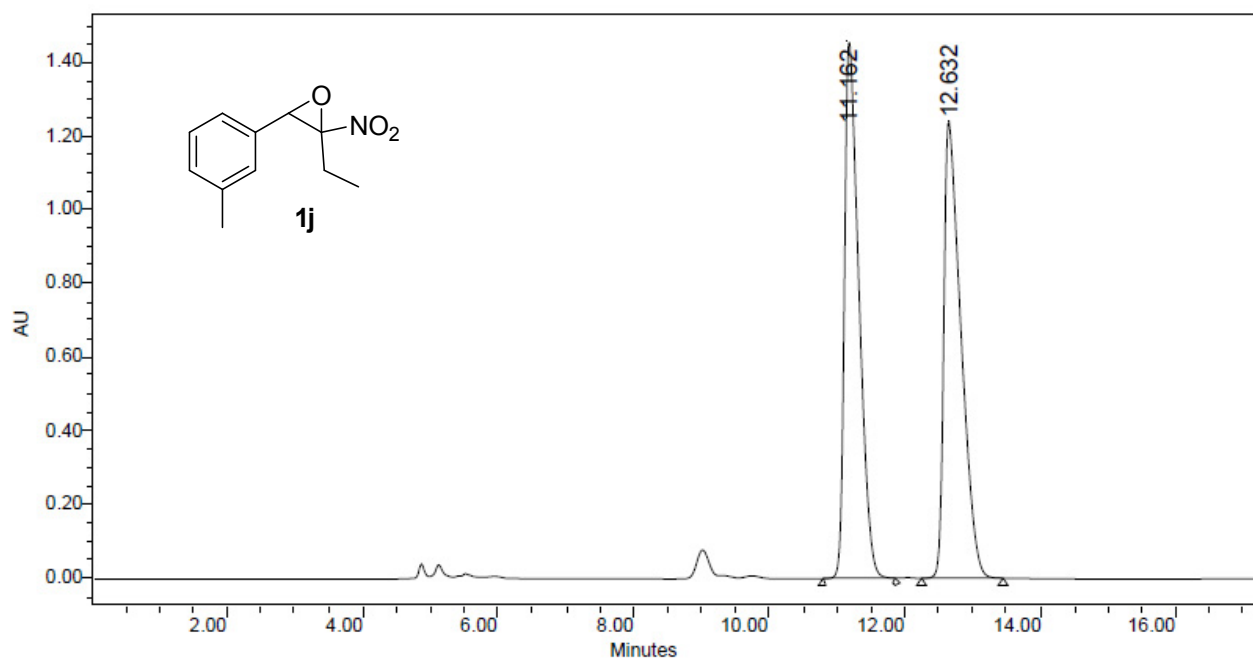
	RT (min)	Area (AU*sec)	% Area	Height (AU)	% Height
1	20.286	4915967	97.43	228886	97.69
2	24.714	129924	2.57	5403	2.31



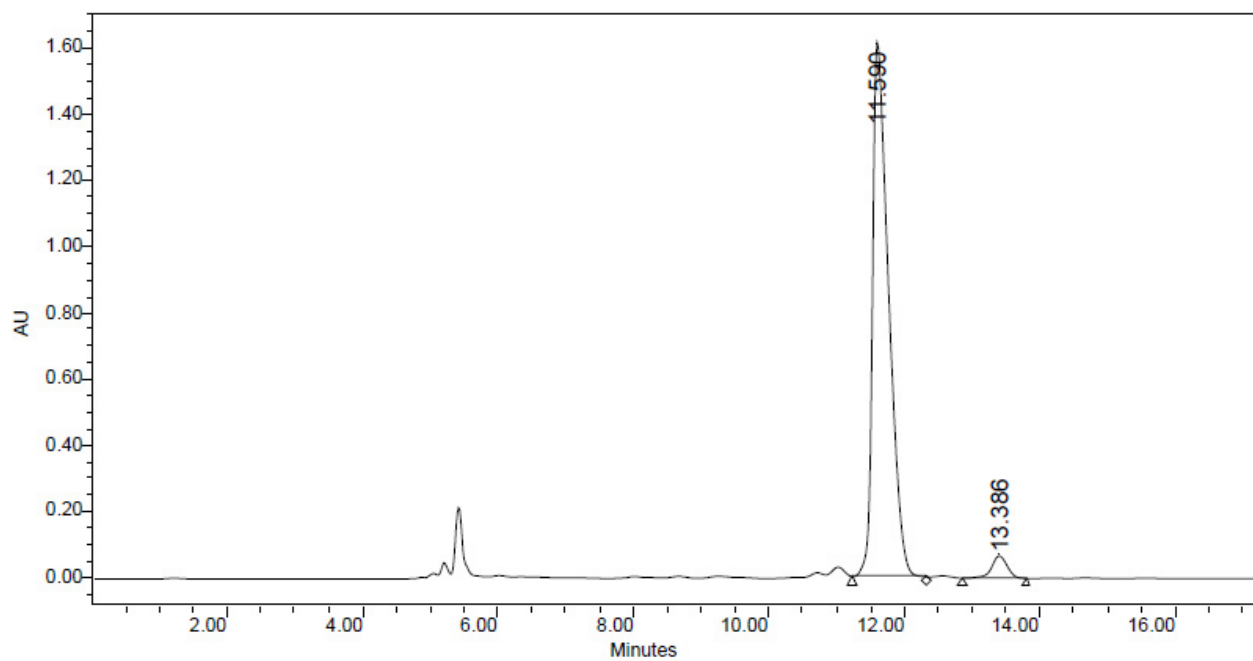
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	11.541	1923611	50.50	118170	50.21
2	13.165	1885766	49.50	117159	49.79



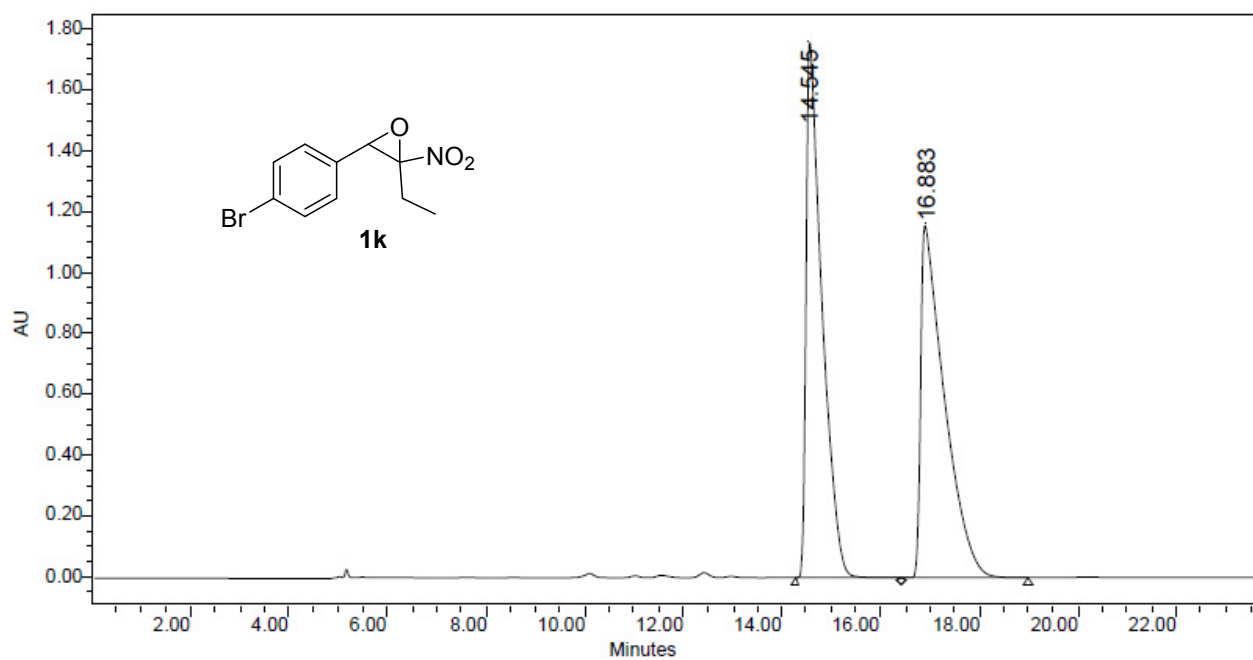
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	11.914	13675053	96.06	1023689	96.47
2	13.701	560750	3.94	37491	3.53



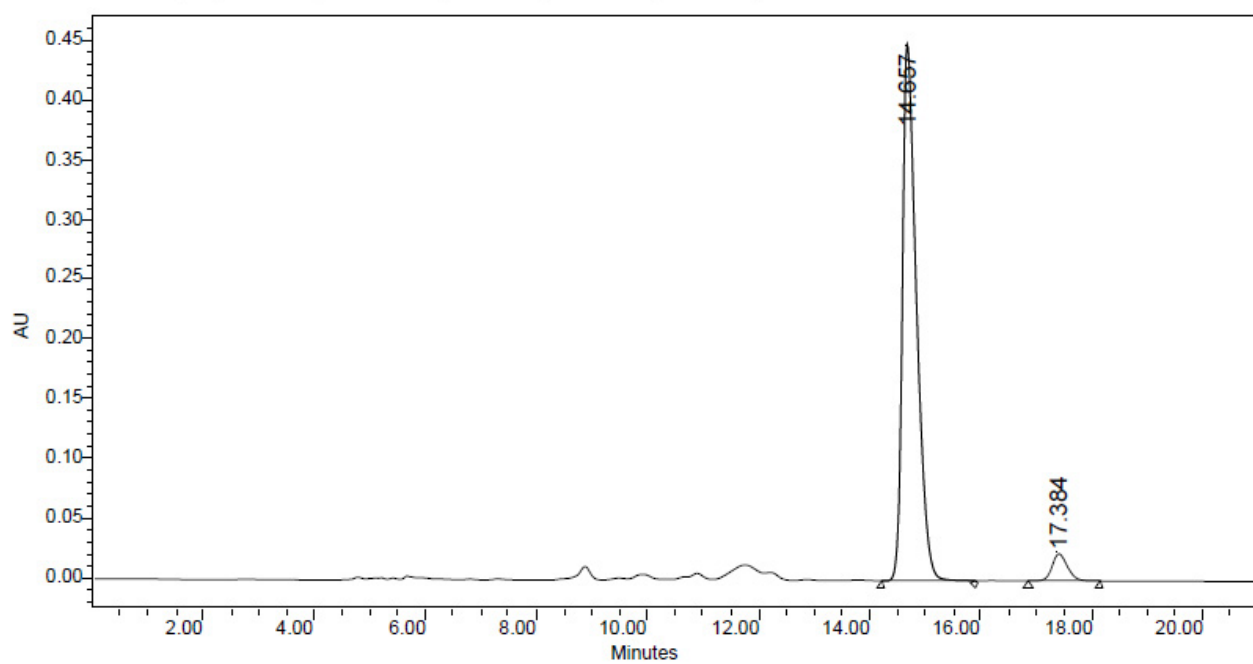
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	11.162	21624916	49.80	1457832	54.04
2	12.632	21802428	50.20	1239960	45.96



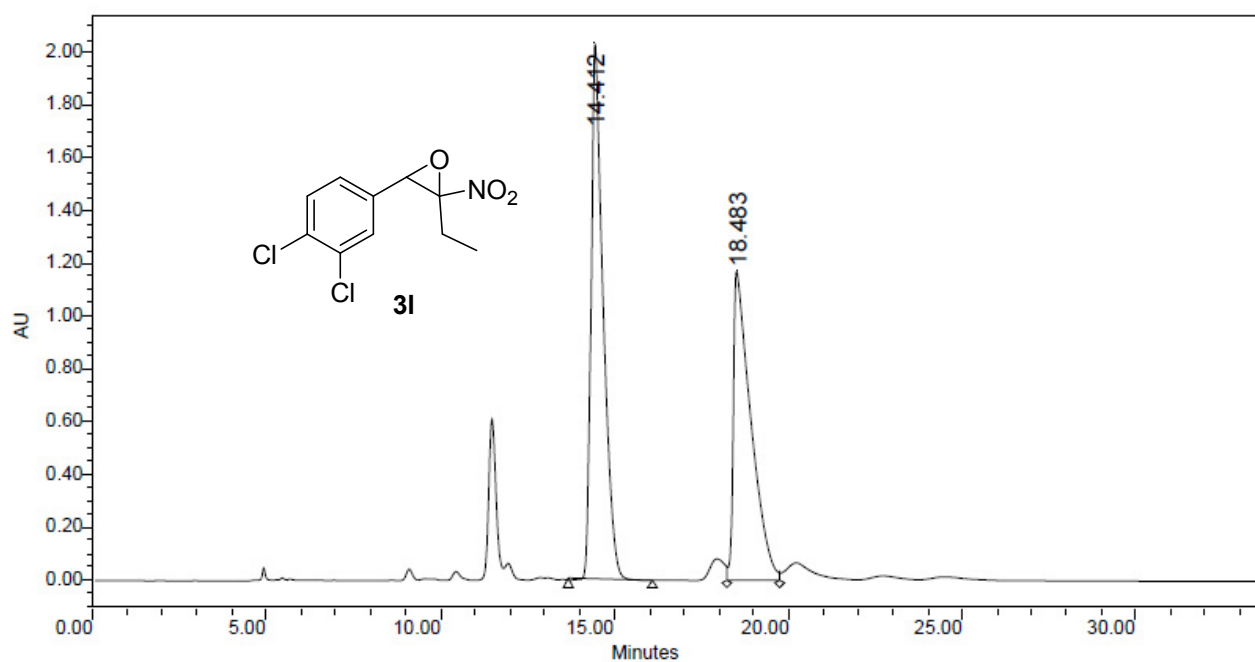
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	11.590	26703128	96.06	1621520	95.97
2	13.386	1096052	3.94	68014	4.03



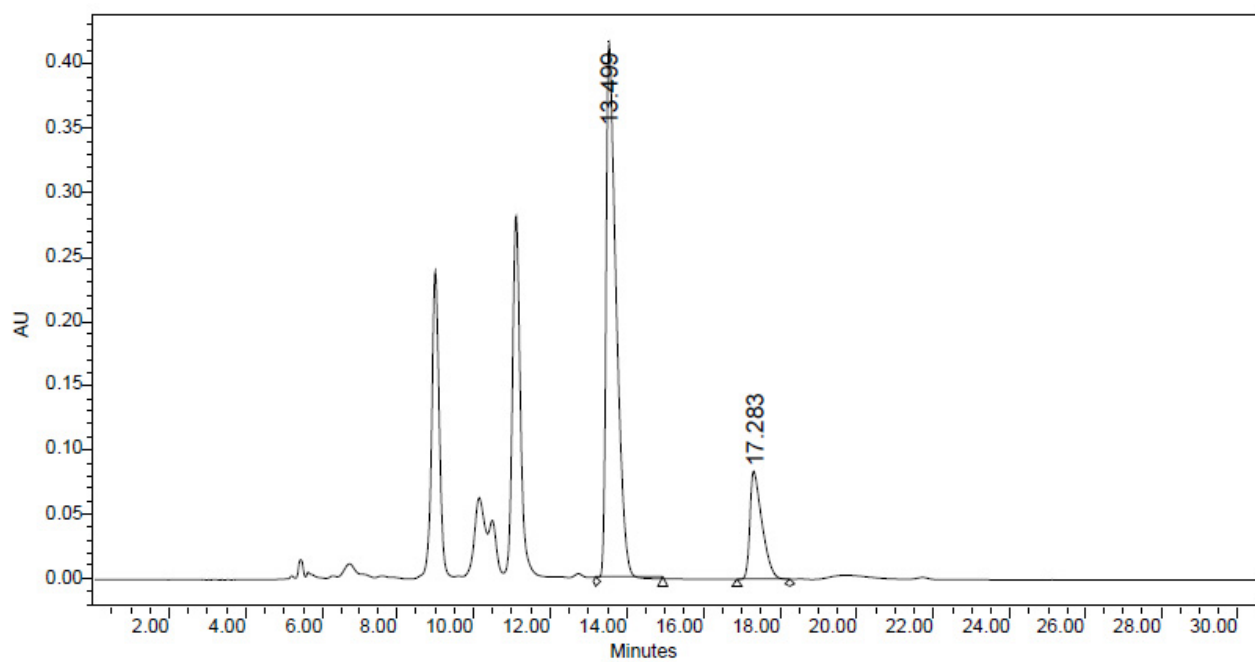
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	14.545	39250385	50.50	1766722	60.46
2	16.883	38466589	49.50	1155372	39.54



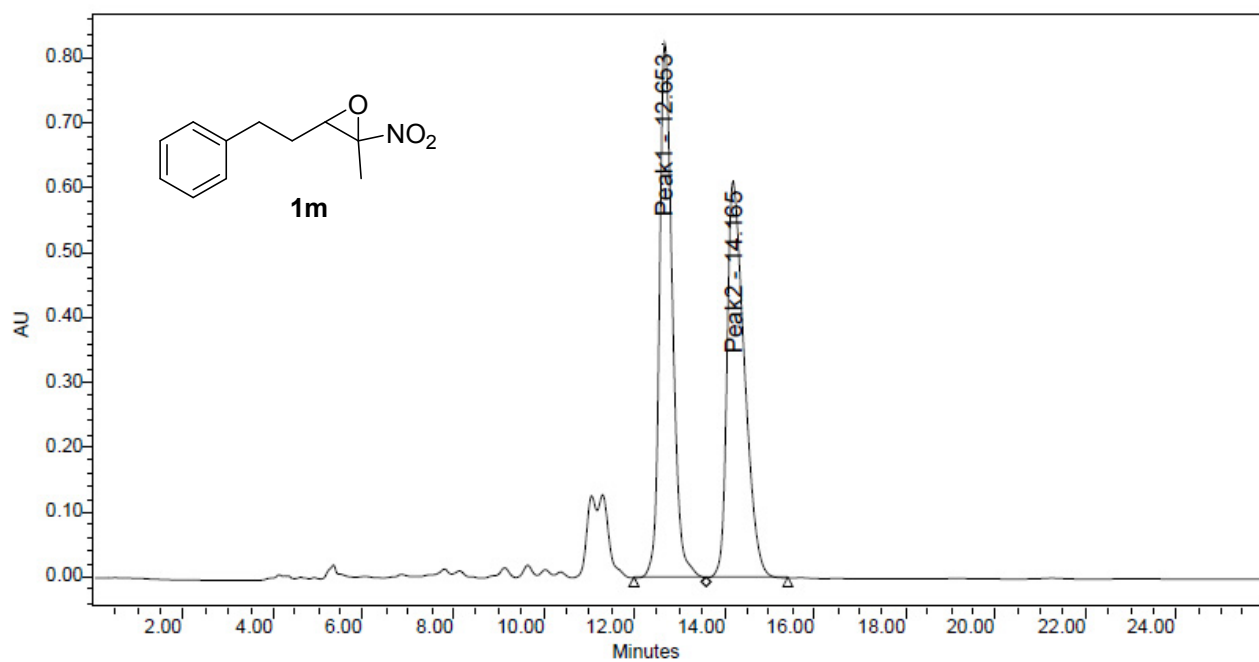
	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	14.657	7879406	94.74	449920	95.20
2	17.384	437210	5.26	22678	4.80



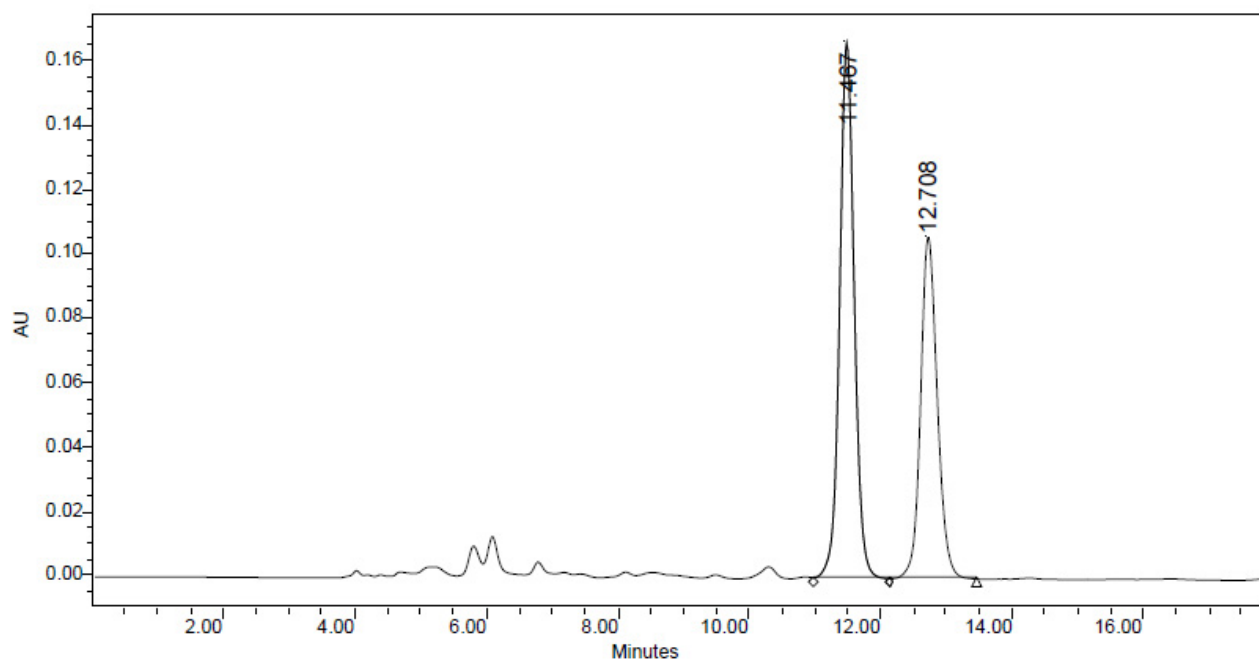
	RT (min)	Area (AU*sec)	% Area	Height (AU)	% Height
1	14.412	48990714	54.39	2036442	63.44
2	18.483	41082265	45.61	1173808	36.56



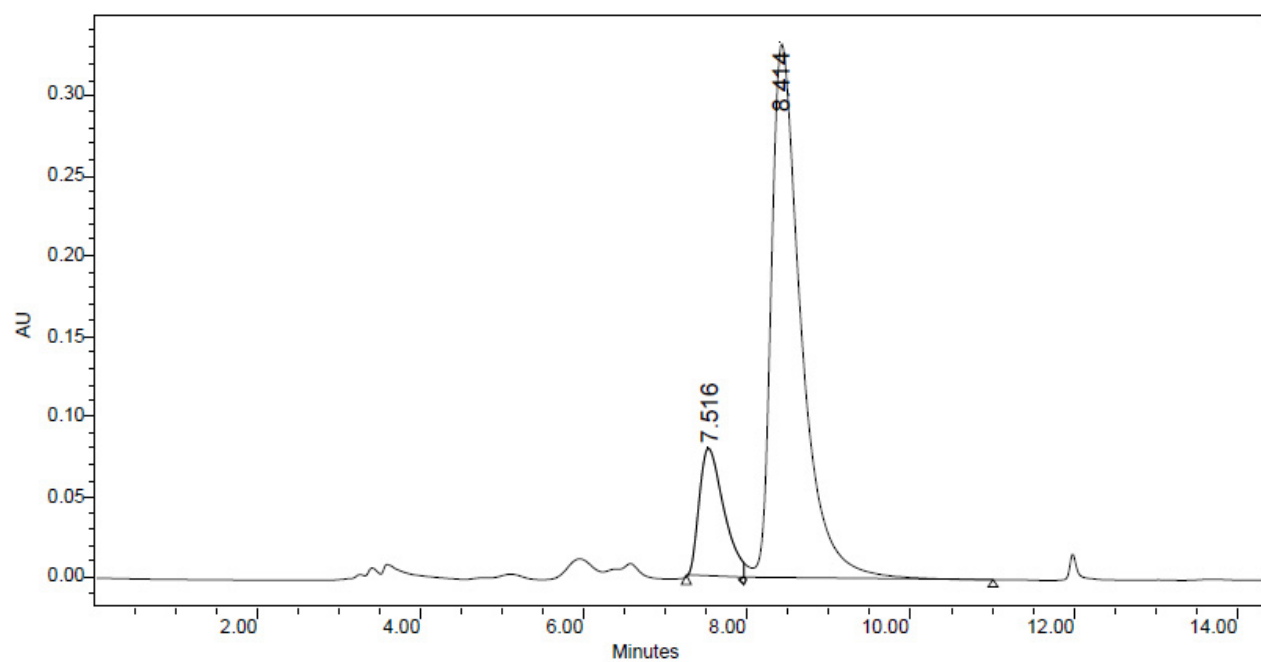
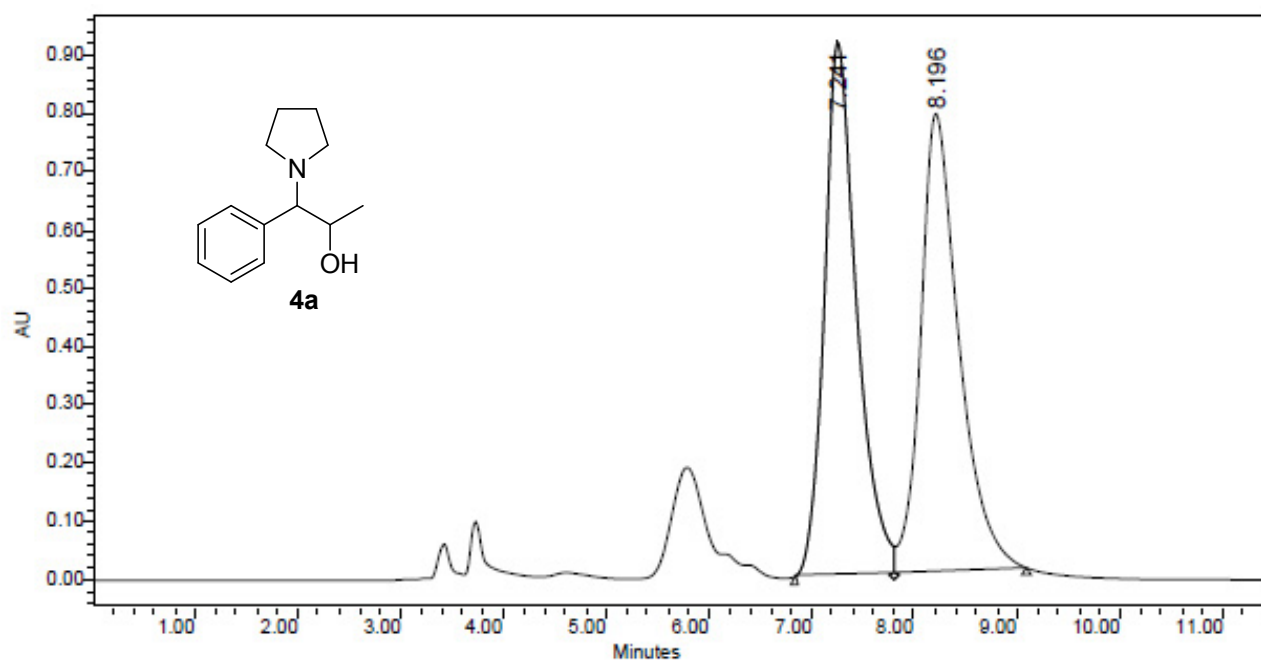
	RT (min)	Area (AU*sec)	% Area	Height (AU)	% Height
1	13.499	7654713	80.51	415746	83.16
2	17.283	1853334	19.49	84187	16.84

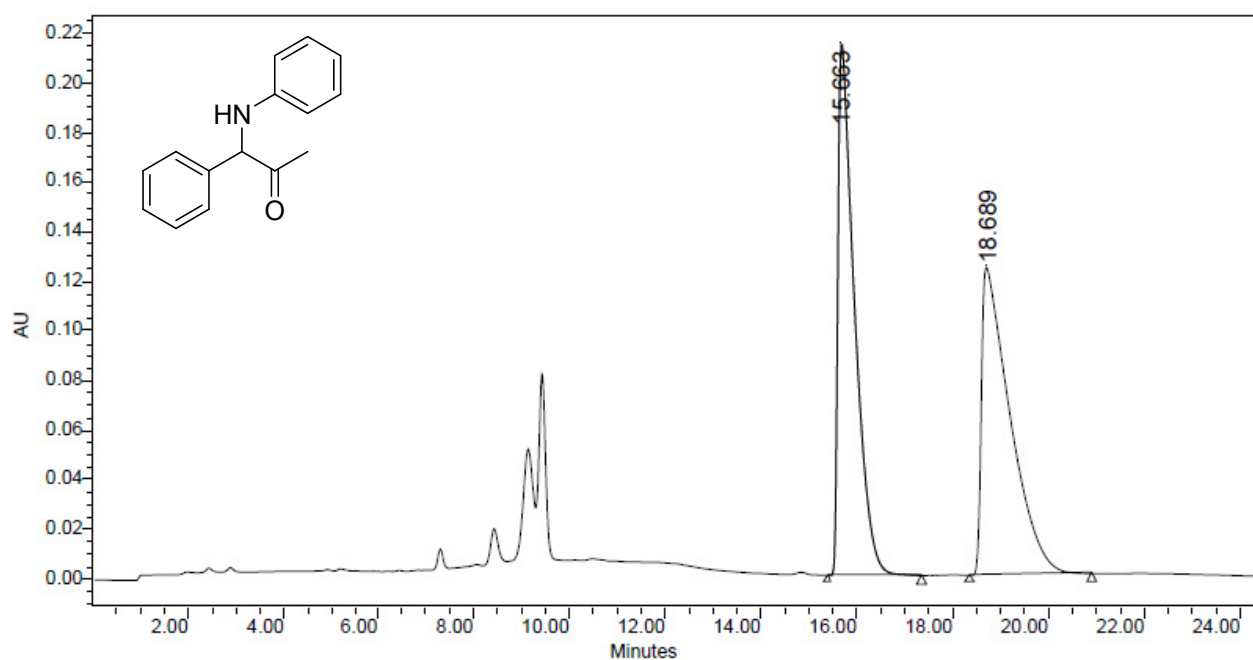


	Peak Name	RT (min)	Area (UV*sec)	% Area	Height (UV)	% Height
1	Peak1	12.653	16423500	50.02	826511	57.44
2	Peak2	14.165	16412094	49.98	612332	42.56

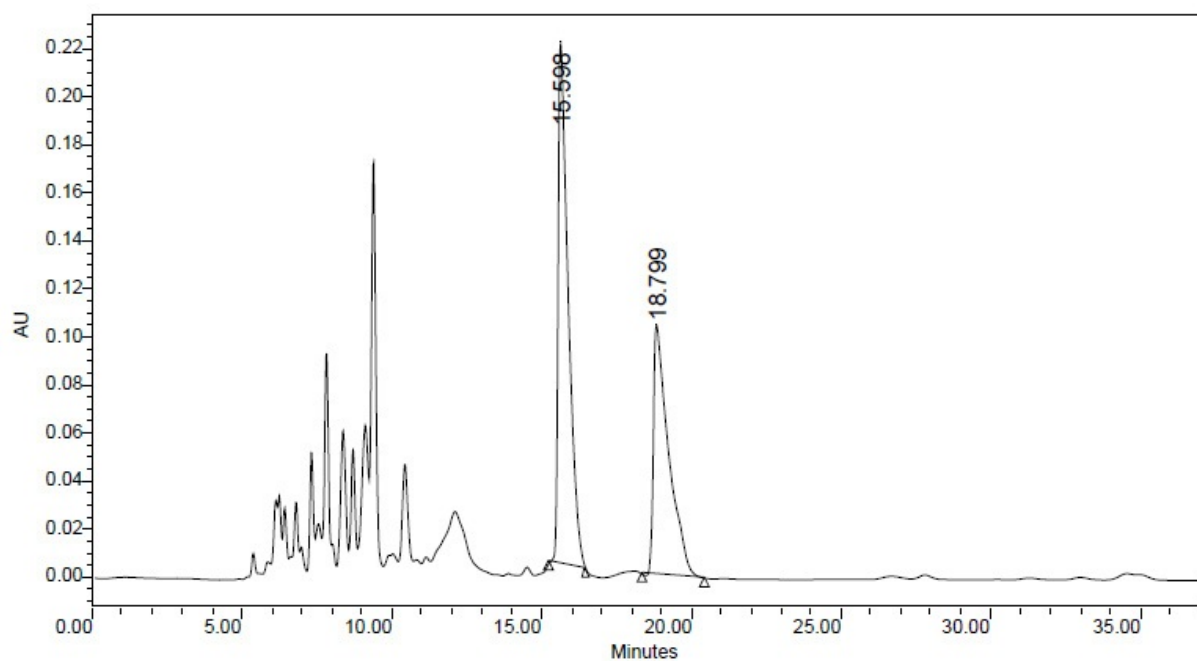


	RT (min)	Area (UV*sec)	% Area	Height (UV)	% Height
1	11.467	2574653	57.82	166210	61.02
2	12.708	1877866	42.18	106186	38.98





	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	15.663	5135581	49.95	214295	63.32
2	18.689	5144843	50.05	124154	36.68



	RT (min)	Area (V*sec)	% Area	Height (V)	% Height
1	15.598	5075409	57.82	216692	67.56
2	18.799	3702638	42.18	104049	32.44