

Supporting Information

**Enantioselective Synthesis of Triarylmethanes via Organocatalytic Transfer
Hydrogenation of *para*-Quinone Methides**

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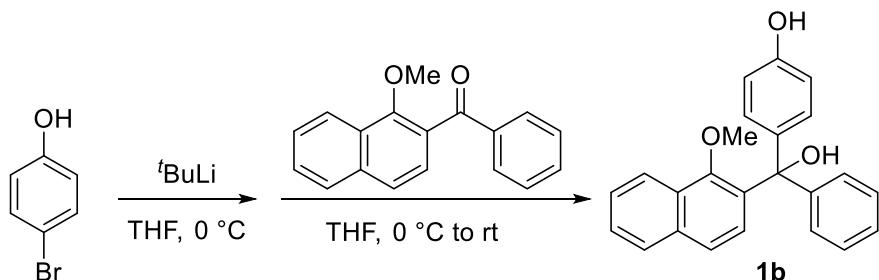
NMR and HPLC Traces

I. General Information

Flash column chromatography was performed over silica gel (200-300 mesh) purchased from Qindao Puke Co., China. All air or moisture sensitive reactions were conducted in oven-dried glassware under nitrogen atmosphere using anhydrous solvents. Anhydrous dichloromethane and tetrahydrofuran were purified by the Innovative® solvent purification system. Anhydrous solvents were purchased from Sigma-Aldrich® and J&K® and used as received. ^1H , ^{13}C , ^{19}F and ^{31}P NMR spectra were collected on a Bruker AV 500 MHz or 400 MHz or 300 MHz NMR spectrometer using residue solvent peaks as an internal standard (^1H NMR: CDCl_3 at 7.26 ppm, acetone- d_6 at 2.05 ppm, ^{13}C NMR: CDCl_3 at 77.0 ppm, acetone- d_6 at 205.9 ppm). Mass spectra were collected on an Agilent GC/MS 5975C system or an API QSTAR XL System. Optical rotations were measured on a Shanghai Shenguang® polarimeter with $[\alpha]_D$ values reported in degrees; concentration (c) is in 10 mg/mL. The enantiomeric excesses were determined by chiral HPLC using an Agilent 1200 LC instrument with a Daicel Chiralcel OJ-H column or a Daicel Chiralpak AD-H, IC, AS-H column.

II. Synthesis of Tertiary Alcohol Substrates

All compounds were synthesized according to literature,¹ and the data of **1a**, **1c – 1p**, **1a'**, **1b'** were consistent with those reported in the literature.



4-(Hydroxy(1-methoxynaphthalen-2-yl)(phenyl)methyl)phenol (1b). At 0 °C, *tert*-butyllithium (1.6 M in pentane, 15.5 mL, 24.8 mmol) was slowly added to a stirred solution of 4-bromophenol (1.30 g, 7.5 mmol) in THF (50 mL). The resulting mixture was stirred at 0 °C for 2 hours. Then a solution of (1-methoxynaphthalen-2-yl)(phenyl)methanone (3.0 mmol, 684 mg) in THF (10 mL) was added and the reaction mixture was kept stirring overnight at room temperature. A saturated aqueous solution of NH₄Cl was added to quench the reaction. The reaction mixture was extracted with EtOAc (3 × 30 mL). The combined organic layers were dried over anhydrous Na₂SO₄, filtered, and concentrated. The residue was purified by silica gel flash chromatography (eluent: hexanes/EtOAc = 15:1 → 10:1) to afford the pure tertiary alcohol **1b** as pale yellow solid in 58% yield (621 mg).

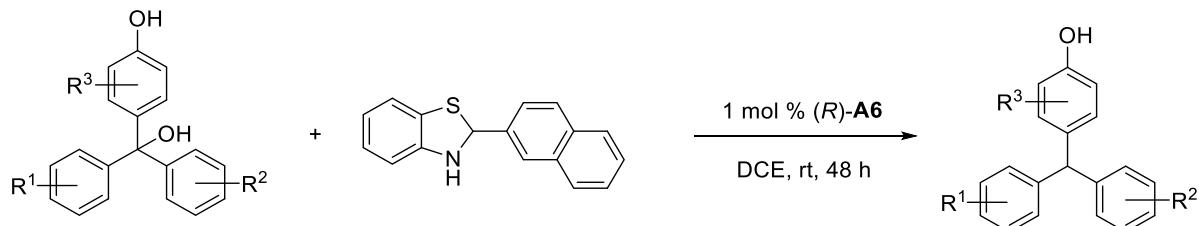
¹H NMR (500 MHz, acetone-*d*₆) δ 8.05 (d, *J* = 8.2 Hz, 1H), 7.96 – 7.86 (m, 1H), 7.61 – 7.47 (m, 3H), 7.41 – 7.23 (m, 5H), 7.19 – 7.10 (m, 2H), 6.89 – 6.75 (m, 3H), 5.93 (s, 1H), 3.34 (s, 3H), 2.88 (s, 1H).

¹³C NMR (126 MHz, acetone-*d*₆) δ 157.1, 154.3, 148.4, 138.71, 137.9, 135.4, 129.9, 128.8, 128.6, 128.5, 128.2, 127.8, 127.6, 126.9, 126.8, 123.5, 122.7, 115.0, 82.4, 61.9.

HRMS (ES+) Calcd for C₂₄H₂₀O₃Na [M + Na]⁺: 379.1310, Found: 379.1306.

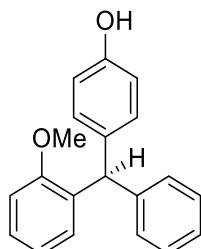
III. Catalytic Asymmetric Synthesis of Triarylmethanes

General Procedure.



At 0 °C or room temperature, to an oven-dried 10-mL vial charged with a solution of the tertiary alcohol **1** (0.3 mmol) and 2-naphthylbenzothiazoline (0.6 mmol) in DCE (2.5 mL) was slowly added a solution of the catalyst (R)-**A6** (3.0 mg, 3.0 µmol, 1.0 mol%) in DCE (0.5 mL). The reaction mixture was stirred at room temperature for 48 h (or otherwise noted temperature and time). Next, Na₂CO₃ (212 mg, 2.0 mmol) was added. The mixture was stirred for 10 min and concentrated under reduced pressure. The residue was purified by silica gel flash chromatography to afford the desired product.

Unless noted otherwise, all the racemic products (used for HPLC reference in determining the ee value) were prepared from the same reaction using a simple achiral phosphoric acid as the catalyst.



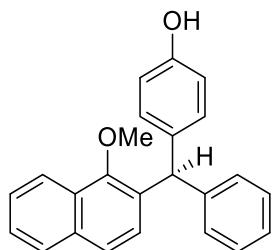
(R)-4-((2-Methoxyphenyl)(phenyl)methyl)phenol (2a) was prepared as a pale yellow foam from 4-(hydroxy(2-methoxyphenyl)(phenyl)methyl)phenol (**1a**) (91.9 mg, 0.3 mmol) and 2-naphthylbenzothiazoline (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 91% yield (79.0 mg) and 95% ee. ¹H NMR (400 MHz, CDCl₃) δ 7.32 – 7.12 (m, 4H), 7.11 – 7.02 (m, 2H), 6.98 – 6.90 (m,

2H), 6.90 – 6.79 (m, 3H), 6.75 – 6.64 (m, 2H), 5.85 (s, 1H), 4.06 (brs, 1H), 3.70 (s, 3H).

¹³C NMR (100 MHz, CDCl₃) δ 157.0, 153.8, 144.2, 136.0, 132.9, 130.5, 130.2, 129.3, 128.0, 127.4, 125.9, 120.2, 114.9, 110.7, 55.6, 48.7.

[α]_D²⁵: -2.5 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 6.8 min (minor), 7.2 min (major).

HRMS (CI+) Calcd for C₂₀H₁₈O₂ [M]⁺: 290.1307, Found: 290.1297.



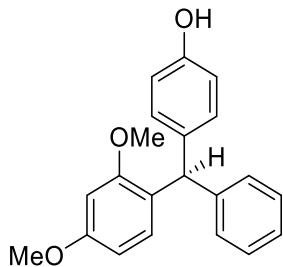
(R)-4-((1-Methoxynaphthalen-2-yl)(phenyl)methyl)phenol (2b) was prepared as pale yellow oil from 4-(hydroxy(1-methoxynaphthalen-2-yl)(phenyl)methyl)phenol (**1b**) (106.9 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 97% yield (99.2 mg) and 97% ee .

¹H NMR (300 MHz, acetone-*d*₆) δ 8.21 (s, 1H), 8.05 – 7.95 (m, 1H), 7.88 (d, *J* = 9.0 Hz, 1H), 7.86 – 7.78 (m, 1H), 7.43 (d, *J* = 9.0 Hz, 1H), 7.32 – 7.25 (m, 2H), 7.25 – 7.06 (m, 7H), 6.82 – 6.73 (m, 2H), 6.53 (s, 1H), 3.66 (s, 3H).

¹³C NMR (75 MHz, acetone-*d*₆) δ 155.6, 155.5, 144.5, 133.8, 133.3, 130.2, 130.1, 129.2, 128.9, 128.7, 127.9, 125.9, 125.8, 125.5, 125.0, 123.2, 115.1, 114.9, 56.1, 46.7.

[α]_D²⁵: +5.6 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralpak IC column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 5.3 min (minor), 8.0 min (major).

HRMS (ES-) Calcd for C₂₄H₁₉O₂ [M - H]⁺: 339.1385, Found: 339.1387.



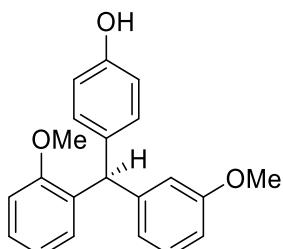
(R)-4-((2,4-dimethoxyphenyl)(phenyl)methyl)phenol (2c) was prepared as a yellow oil from 4-((2,4-dimethoxyphenyl)(hydroxy)(phenyl)methyl)phenol (**1c**) (100.9 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 99% yield (95.2 mg) and 95% ee.

¹H NMR (300 MHz, acetone-*d*₆) δ 8.19 (s, 1H), 7.34 – 7.21 (m, 2H), 7.21 – 7.11 (m, 1H), 7.11 – 7.01 (m, 2H), 6.95 – 6.84 (m, 2H), 6.74 (ddd, *J* = 8.2, 5.4, 2.9 Hz, 3H), 6.54 (d, *J* = 2.4 Hz, 1H), 6.43 (dd, *J* = 8.4, 2.5 Hz, 1H), 5.73 (s, 1H), 3.76 (s, 3H), 3.70 (s, 3H).

¹³C NMR (75 MHz, acetone-*d*₆) δ 160.1, 158.4, 156.1, 145.5, 135.5, 130.9, 130.7, 129.6, 128.5, 126.2, 125.7, 115.3, 104.5, 98.8, 55.5, 55.1, 48.7.

[α]_D²⁵: -11.4 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiraldak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 10.2 min (major), 10.8 min (minor).

HRMS (Cl⁺) Calcd for C₂₁H₂₀O₃ [M]⁺: 320.1412, Found: 320.1416.



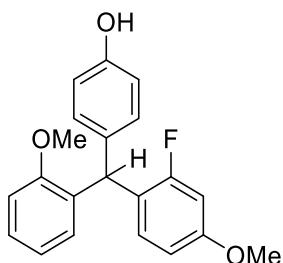
(S)-4-((2-methoxyphenyl)(3-methoxyphenyl)methyl)phenol (2d) was prepared as a white foam from 4-(hydroxy(2-methoxyphenyl)(3-methoxyphenyl)methyl)phenol (**1d**) (100.9 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 98% yield (94.5 mg) and 94% ee.

¹H NMR (300 MHz, acetone-*d*₆) δ 8.22 (s, 1H), 7.27 – 7.10 (m, 2H), 6.96 (d, *J* = 8.2 Hz, 1H), 6.94 – 6.82 (m, 4H), 6.74 (dd, *J* = 6.7, 4.7 Hz, 3H), 6.68 – 6.57 (m, 2H), 5.80 (s, 1H), 3.72 (s, 3H), 3.70 (s, 3H).

¹³C NMR (75 MHz, acetone-*d*₆) δ 160.2, 157.6, 156.2, 146.7, 135.0, 133.3, 130.7, 130.4, 129.4, 128.0, 122.1, 120.5, 115.9, 115.4, 111.2, 55.5, 54.9, 49.1.

[α]_D²⁵: -4.2 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 9.4 min (minor), 10.2 min (major).

HRMS (ES-) Calcd for C₂₁H₁₉O₃ [M - H]⁺: 319.1340, Found: 319.1420.



4-((2-Fluoro-4-methoxyphenyl)(2-methoxyphenyl)methyl)phenol (2e) was prepared as a yellow foam from 4-((2-fluoro-4-methoxyphenyl)(hydroxy)(2-methoxyphenyl)methyl)phenol (**1e**) (106.3 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure with 10 mol% of catalyst at 0 °C for 72 h (eluent: hexanes/EtOAc = 15:1 → 10:1) in 90% yield (90.4 mg) and 91% ee.

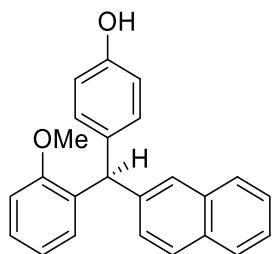
¹H NMR (400 MHz, CDCl₃) δ 7.25 – 7.21 (m, 1H), 6.95 (d, *J* = 8.5 Hz, 2H), 6.89 – 6.82 (m, 3H), 6.79 – 6.72 (m, 3H), 6.63 – 6.57 (m, 2H), 6.01 (s, 1H), 4.91 (s, 1H), 3.78 (s, 3H), 3.72 (s, 3H).

¹³C NMR (100 MHz, CDCl₃) δ 161.1 (d, *J* = 244.9 Hz), 159.2 (d, *J* = 10.9 Hz), 156.9, 153.8, 135.0, 131.9, 130.8 (d, *J* = 6.1 Hz), 130.2, 129.8, 127.6, 123.2 (d, *J* = 15.1 Hz), 120.2, 115.0, 110.7, 109.1 (d, *J* = 2.9 Hz), 101.5 (d, *J* = 27.8 Hz), 55.6, 55.4, 41.3 (d, *J* = 2.6 Hz).

¹⁹F NMR (376 MHz, CDCl₃) δ -114.19.

$[\alpha]_D^{25}$: -12.8 ($c = 1.0$, CHCl_3). HPLC analysis of the product: Daicel Chiraldak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 9.4 min (minor), 10.2 min (major).

HRMS (Cl⁺) Calcd for $\text{C}_{21}\text{H}_{19}\text{FO}_3$ [M]⁺: 338.1318, Found: 338.1313.

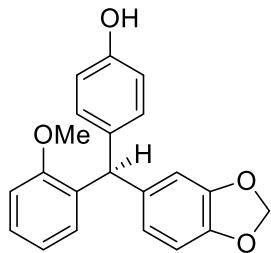


(S)-4-((2-Methoxyphenyl)(naphthalen-2-yl)methyl)phenol (2f) was prepared as a pale yellow oil from 4-(hydroxy(2-methoxyphenyl)(naphthalen-2-yl)methyl)phenol (**1f**) (106.9 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 92% yield (93.5 mg) and 94% ee.

¹H NMR (300 M Hz, acetone-*d*₆) δ 8.25 (s, 1H), 7.90 – 7.67 (m, 3H), 7.53 – 7.36 (m, 3H), 7.36 – 7.15 (m, 2H), 7.03 – 6.95 (m, 3H), 6.91 (ddd, *J* = 8.1, 7.4, 1.4 Hz, 2H), 6.84 – 6.76 (m, 2H), 6.03 (s, 1H), 3.70 (s, 3H).

¹³C NMR (75 M Hz, acetone-*d*₆) δ 157.4, 156.1, 142.7, 134.7, 133.8, 133.0, 132.5, 130.8, 130.4, 128.6, 127.9, 127.9, 127.8, 127.8, 127.4, 126.2, 125.7, 120.4, 115.3, 111.1, 55.3, 49.2. $[\alpha]_D^{25}$: +8.0 ($c = 1.0$, CHCl_3). HPLC analysis of the product: Daicel Chiraldak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 10.8 min (minor), 12.1 min (major).

HRMS (Cl⁺) Calcd for $\text{C}_{24}\text{H}_{20}\text{O}_2$ [M]⁺: 340.1463, Found: 340.1464.



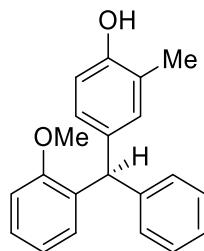
(S)-4-(Benzo[d][1,3]dioxol-5-yl(2-methoxyphenyl)methyl)phenol (2g) was prepared as a yellow oil from 4-(hydroxy(2-methoxyphenyl)(naphthalen-2-yl)methyl)phenol (**1g**) (106.9 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[d]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 99% yield (99.1 mg) and 94% ee.

¹H NMR (300 MHz, acetone-*d*₆) δ 8.20 (s, 1H), 7.20 (ddd, *J* = 8.3, 6.3, 2.9 Hz, 1H), 7.02 – 6.81 (m, 5H), 6.81 – 6.68 (m, 3H), 6.57 (d, *J* = 1.7 Hz, 1H), 6.53 (ddd, *J* = 8.0, 1.7, 0.5 Hz, 1H), 5.92 (s, 2H), 5.77 (s, 1H), 3.71 (s, 3H).

¹³C NMR (75 MHz, acetone-*d*₆) δ 157.3, 155.9, 147.8, 146.1, 138.9, 135.1, 133.2, 130.4, 130.1, 127.8, 122.4, 120.3, 115.2, 111.0, 109.9, 107.9, 101.2, 55.3, 48.6.

[α]_D²⁵: -6.8 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiraldpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 12.6 min (minor), 15.2 min (major).

HRMS (CI+) Calcd for C₂₁H₁₈O₄[M]⁺: 334.1205, Found: 334.1211.



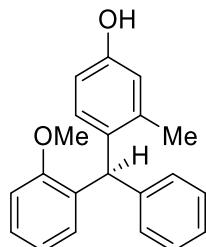
(R)-4-((2-Methoxyphenyl)(phenyl)methyl)-2-methylphenol (2h) was prepared as a colorless oil from 4-(hydroxy(2-methoxyphenyl)(phenyl)methyl)-2-methylphenol (**1h**) (96.1 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[d]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 81% yield (73.9 mg) and 72% ee.

¹H NMR (300 MHz, CDCl₃) δ 7.33 – 7.12 (m, 4H), 7.13 – 7.02 (m, 2H), 6.93 – 6.80 (m, 4H), 6.74 (dd, *J* = 8.2, 2.1 Hz, 1H), 6.62 (d, *J* = 8.2 Hz, 1H), 5.82 (s, 1H), 4.82 (s, 1H), 3.69 (s, 3H), 2.15 (s, 3H).

¹³C NMR (75 MHz, CDCl₃) δ 157.0, 152.0, 144.3, 135.9, 132.9, 132.0, 130.3, 129.3, 128.0, 127.9, 127.4, 125.8, 123.3, 120.2, 114.5, 110.7, 55.6, 48.7, 15.8.

[α]_D²⁵: -12.1 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralpak AS-H column; 5% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 15.9 min (minor), 17.1 min (major).

HRMS (ES-) Calcd for C₂₁H₁₉O₂ [M - H]⁺: 303.1385, Found: 303.1380.



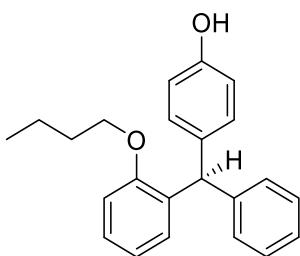
(R)-4-((2-Methoxyphenyl)(phenyl)methyl)-3-methylphenol (2i) was prepared as a colorless oil from 4-(hydroxy(2-methoxyphenyl)(phenyl)methyl)-3-methylphenol (**1i**) (96.1 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 99% yield (91.0 mg) and 73% ee.

¹H NMR (400 MHz, CDCl₃) δ 7.30 – 7.14 (m, 4H), 7.09 – 7.00 (m, 2H), 6.85 (t, *J* = 8.2 Hz, 2H), 6.77 (dd, *J* = 7.5, 1.7 Hz, 1H), 6.64 (d, *J* = 8.0 Hz, 2H), 6.53 (dd, *J* = 8.4, 2.7 Hz, 1H), 5.92 (s, 1H), 4.80 (s, 1H), 3.71 (s, 3H), 2.14 (s, 3H).

¹³C NMR (100 MHz, CDCl₃) δ 157.1, 153.6, 143.6, 138.3, 134.7, 132.5, 130.2, 130.2, 129.5, 128.1, 127.4, 125.9, 120.2, 117.2, 112.0, 110.6, 55.7, 45.6, 19.7.

[α]_D²⁵: -12.3 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralcel OJ-H column; 3% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 33.9 min (minor), 38.3 min (major).

HRMS (ES-) Calcd for C₂₁H₁₉O₂ [M - H]⁺: 303.1385, Found: 303.1380.



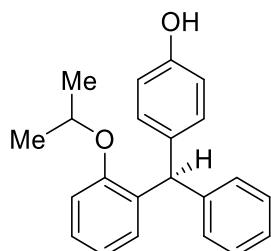
(R)-4-((2-Butoxyphenyl)(phenyl)methyl)phenol (2j) was prepared as a colorless oil from 4-((2-butoxyphenyl)(hydroxy)(phenyl)methyl)phenol (**1j**) (104.5 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure with 10 mol% of catalyst at 0 °C for 72 h (eluent: hexanes/EtOAc = 15:1 → 10:1) in 95% yield (94.4 mg) and 94% ee.

¹H NMR (300 MHz, CDCl₃) δ 7.28 – 7.11 (m, 4H), 7.11 – 7.02 (m, 2H), 6.99 – 6.89 (m, 2H), 6.86 – 6.77 (m, 3H), 6.72 – 6.62 (m, 2H), 5.82 (s, 1H), 5.19 (brs, 1H), 3.82 (t, *J* = 6.2 Hz, 2H), 1.64 – 1.44 (m, 2H), 1.36 – 1.11 (m, 2H), 0.82 (t, *J* = 7.4 Hz, 3H).

¹³C NMR (75 MHz, CDCl₃) δ 156.5, 153.6, 144.2, 136.1, 133.2, 130.5, 130.0, 129.3, 128.0, 127.3, 125.8, 119.9, 114.9, 111.4, 67.6, 49.2, 31.2, 19.0, 13.7.

[α]_D²⁵: -24.5 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiraldapak AS-H column; 5% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 10.5 min (minor), 11.2 min (major).

HRMS (ES-) Calcd for C₂₃H₂₃O₂[M - H]⁺: 331.1698, Found: 331.1699.



(R)-4-((2-Isopropoxypyhenyl)(phenyl)methyl)phenol (2k) was prepared as a colorless oil from 4-(hydroxy(2-isopropoxypyhenyl)(phenyl)methyl)phenol (**1k**) (100.3 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol)

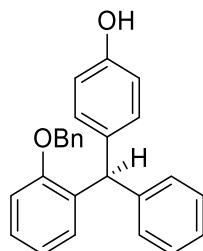
according to the General Procedure with 10 mol% of catalyst at -20 °C for 7 days (eluent: hexanes/EtOAc = 15:1 → 10:1) in 90% yield (86.0 mg) and 92% ee.

¹H NMR (300 MHz, CDCl₃) δ 7.30 – 7.20 (m, 2H), 7.20 – 7.12 (m, 2H), 7.12 – 7.03 (m, 2H), 7.01 – 6.90 (m, 2H), 6.88 – 6.77 (m, 3H), 6.75 – 6.65 (m, 2H), 5.79 (s, 1H), 5.01 (s, 1H), 4.44 (dt, *J* = 12.1, 6.0 Hz, 1H), 1.07 (dd, *J* = 8.4, 6.0 Hz, 6H).

¹³C NMR (75 MHz, CDCl₃) δ 155.3, 153.6, 144.2, 136.1, 134.2, 130.6, 130.2, 129.4, 128.0, 127.2, 125.8, 119.9, 114.9, 113.0, 69.9, 49.3, 21.9, 21.8.

[α]_D²⁵: -17.3 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralcel OJ-H column; 5% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 17.9 min (major), 20.6 min (minor).

HRMS (ES-) Calcd for C₂₂H₂₁O₂[M - H]⁺: 317.1542, Found: 317.1541.



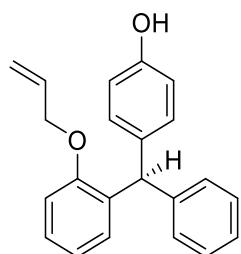
(R)-4-((2-(Benzylxy)phenyl)(phenyl)methyl)phenol (2l) was prepared as a colorless oil from 4-((2-(benzylxy)phenyl)(hydroxy)(phenyl)methyl)phenol (**1l**) (114.7 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure with 10 mol% of catalyst at 0 °C for 72 h (eluent: hexanes/EtOAc = 15:1 → 10:1) in 86% yield (94.0 mg) and 94% ee.

¹H NMR (300 MHz, acetone-*d*₆) δ 8.31 (s, 1H), 7.35 – 7.11 (m, 9H), 7.12 – 7.00 (m, 3H), 6.95 – 6.81 (m, 4H), 6.80 – 6.72 (m, 2H), 5.88 (s, 1H), 5.04 (s, 2H).

¹³C NMR (75 MHz, acetone-*d*₆) δ 156.5, 156.3, 145.0, 138.0, 135.0, 133.8, 130.9, 130.5, 129.8, 128.7, 128.6, 128.0, 127.9, 127.6, 126.4, 120.7, 115.5, 112.5, 69.9, 49.8.

[α]_D²⁵: -18.9 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiraldak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 9.6 min (major), 12.4 min (minor).

HRMS (Cl⁺) Calcd for C₂₆H₂₂O₂[M]⁺: 366.1620, Found: 366.1629.



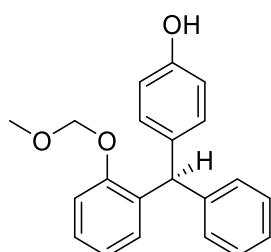
(R)-4-((2-(Allyloxy)phenyl)(phenyl)methyl)phenol (2m) was prepared as a colorless oil from 4-((2-(allyloxy)phenyl)(hydroxy)(phenyl)methyl)phenol (**1m**) (99.7 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[d]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 98% yield (93.1 mg) and 93% ee.

¹H NMR (300 MHz, acetone-d₆) δ 8.20 (s, 1H), 7.29 – 7.23 (m, 2H), 7.23 – 7.13 (m, 2H), 7.13 – 7.05 (m, 2H), 7.01 – 6.82 (m, 5H), 6.81 – 6.71 (m, 2H), 5.98 – 5.79 (m, 2H), 5.26 (dq, *J* = 17.3, 1.8 Hz, 1H), 5.11 (dq, *J* = 10.6, 1.6 Hz, 1H), 4.48 (dt, *J* = 4.6, 1.6 Hz, 2H).

¹³C NMR (75 MHz, acetone-d₆) δ 156.6, 156.2, 145.1, 135.1, 134.2, 133.9, 130.8, 130.5, 129.8, 128.6, 127.9, 126.4, 120.8, 116.3, 115.4, 112.6, 68.9, 49.6.

[α]_D²⁵: -17.7 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiraldapak AD-H column; 2% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 38.1 min (major), 41.8 min (minor).

HRMS (ES-) Calcd for C₂₂H₁₉O₂[M - H]⁺: 315.1385, Found: 315.1382.



(R)-4-((2-(Methoxymethoxy)phenyl)(phenyl)methyl)phenol (2n) was prepared as a colorless oil from 4-(hydroxy(2-(methoxymethoxy)phenyl)(phenyl)methyl)phenol (**1n**) (100.9 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[d]thiazole (158.0 mg,

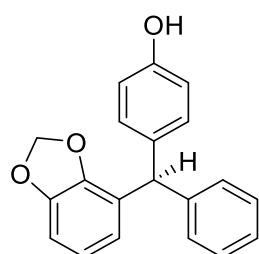
0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 99% yield (95.6 mg) and 92% ee.

¹H NMR (400 MHz, acetone-*d*₆) δ 8.22 (s, 1H), 7.33 – 7.23 (m, 2H), 7.22 – 7.14 (m, 2H), 7.13 – 7.05 (m, 3H), 6.95 – 6.84 (m, 4H), 6.80 – 6.74 (m, 2H), 5.86 (s, 1H), 5.09 (s, 2H), 3.15 (s, 3H).

¹³C NMR (75 MHz, acetone-*d*₆) δ 156.2, 155.0, 145.0, 135.0, 134.3, 130.8, 130.5, 129.7, 128.5, 127.8, 126.4, 121.6, 115.4, 114.5, 94.3, 55.5, 49.6.

[α]_D²⁵: +3.8 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiraldpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 7.2 min (major), 7.7 min (minor).

HRMS (ES-) Calcd for C₂₁H₁₉O₃[M - H]⁺: 319.1334, Found: 319.1334.



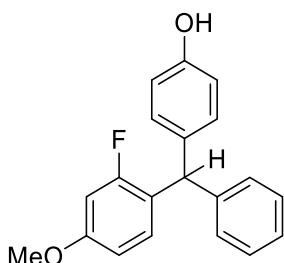
(R)-4-(Benzo[*d*][1,3]dioxol-4-yl(phenyl)methyl)phenol (2o) was prepared as a colorless oil from 4-(benzo[*d*][1,3]dioxol-4-yl(hydroxy)(phenyl)methyl)phenol (**1o**) (96.1 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 90% yield (82.1 mg) and 72% ee.

¹H NMR (300 MHz, CDCl₃) δ 7.33 – 7.17 (m, 3H), 7.17 – 7.08 (m, 2H), 7.05 – 6.93 (m, 2H), 6.82 – 6.68 (m, 4H), 6.53 – 6.39 (m, 1H), 5.87 (s, 2H), 5.58 (s, 1H), 4.89 (s, 1H).

¹³C NMR (75 MHz, CDCl₃) δ 154.0, 147.1, 145.3, 142.9, 134.9, 130.3, 129.1, 128.3, 126.4, 125.8, 122.6, 121.3, 115.1, 106.9, 100.7, 49.6.

[α]_D²⁵: -5.2 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiraldpak AD-H column; 3% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 48.9 min (minor), 53.1 min (major).

HRMS (ES-) Calcd for C₂₀H₁₅O₃ [M - H]⁺: 303.1006, Found: 303.1021.



4-((2-Fluoro-4-methoxyphenyl)(phenyl)methyl)phenol (2p) was prepared as a colorless oil from 4-((2-fluoro-4-methoxyphenyl)(hydroxy)(phenyl)methyl)phenol (**1p**) (97.3 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[d]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 80% yield (75.0 mg) (-20 °C, 10 mol % CPA) and 80% ee.

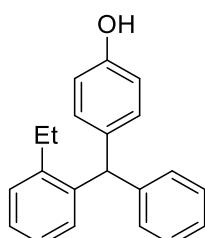
¹H NMR (300 MHz, acetone-*d*₆) δ 8.28 (s, 1H), 7.35 – 7.24 (m, 2H), 7.24 – 7.16 (m, 1H), 7.12 (d, *J* = 7.3 Hz, 2H), 6.95 (d, *J* = 8.5 Hz, 2H), 6.87 (dd, *J* = 10.2, 7.6 Hz, 1H), 6.80 (d, *J* = 8.0 Hz, 2H), 6.74 – 6.63 (m, 2H), 5.69 (s, 1H), 3.78 (d, *J* = 1.0 Hz, 3H).

¹³C NMR (75 MHz, acetone-*d*₆) δ 161.4 (d, *J* = 244.9 Hz), 160.1 (d, *J* = 11.1 Hz), 156.30, 144.1, 134.2, 131.5 (d, *J* = 6.0 Hz), 130.5, 129.4, 128.6, 126.6, 123.7 (d, *J* = 15.1 Hz), 115.5, 109.9 (d, *J* = 3.0 Hz), 101.7 (d, *J* = 26.1 Hz), 55.4, 48.5 (d, *J* = 2.4 Hz).

¹⁹F NMR (282 MHz, CDCl₃) δ -114.29.

[α]_D²⁵: -14.2 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralpak IC column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 5.1 min (minor), 5.8 min (major).

HRMS (ES-) Calcd for C₂₀H₁₆FO₂ [M - H]⁺: 307.1134, Found: 307.1136.



4-((2-Ethylphenyl)(phenyl)methyl)phenol (3a) was prepared as a colorless oil from 4-((2-ethylphenyl)(hydroxy)(phenyl)methyl)phenol (**1a'**) (91.3 mg, 0.3 mmol) and 2-

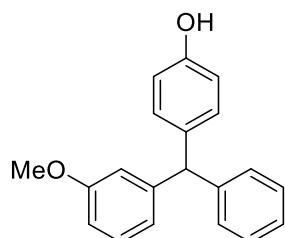
(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 96% yield (83.0 mg) and 55% ee.

¹H NMR (400 MHz, CDCl₃) δ 7.27 – 7.15 (m, 5H), 7.10 – 7.03 (m, 3H), 6.90 (d, *J* = 8.5 Hz, 2H), 6.83 (d, *J* = 7.6 Hz, 1H), 6.73 – 6.71 (m, 2H), 5.70 (s, 1H), 4.87 (brs, 1H), 2.59 (q, *J* = 7.5, 2H), 1.12 (t, *J* = 7.5, 3H).

¹³C NMR (100 MHz, CDCl₃) δ 153.8, 144.1, 142.2, 141.7, 136.0, 130.7, 129.7, 129.5, 128.5, 128.2, 126.5, 126.1, 125.6, 115.1, 51.7, 25.5, 15.0.

[α]_D²⁵: +2.6 (*c* = 1.0, CHCl₃). HPLC analysis of the product: Daicel Chiralpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 5.2 min (minor), 5.6 min (major).

HRMS (ES-) Calcd for C₂₁H₁₉O [M - H]⁺: 287.1441, Found: 287.1434.



4-((3-Methoxyphenyl)(phenyl)methyl)phenol (3b) was prepared as a colorless oil from 4-(hydroxy(3-methoxyphenyl)(phenyl)methyl)phenol (**1b'**) (91.9 mg, 0.3 mmol) and 2-(naphthalen-2-yl)-2,3-dihydrobenzo[*d*]thiazole (158.0 mg, 0.6 mmol) according to the General Procedure (eluent: hexanes/EtOAc = 15:1 → 10:1) in 91% yield (79.1 mg) and 21% ee.

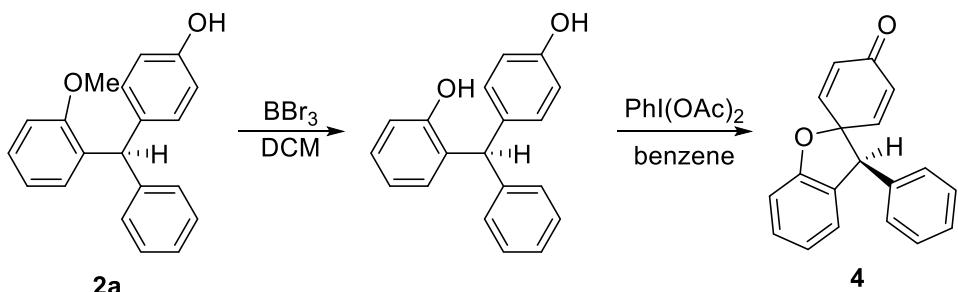
¹H NMR (400 MHz, CDCl₃) δ 7.32 – 7.14 (m, 4H), 7.14 – 7.05 (m, 2H), 6.95 (d, *J* = 8.4 Hz, 2H), 6.84 – 6.59 (m, 5H), 5.44 (s, 1H), 5.14 (brs, 1H), 3.72 (s, 3H).

¹³C NMR (101 MHz, CDCl₃) δ 159.5, 154.0, 145.9, 144.0, 136.0, 130.5, 129.3, 129.2, 128.3, 126.2, 122.0, 115.5, 115.1, 111.2, 55.9, 55.1.

$[\alpha]_D^{25}$: -4.2 ($c = 1.0$, CHCl_3). HPLC analysis of the product: Daicel Chiralpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 11.6 min (major), 12.4 min (minor).

HRMS (CI-) Calcd for $\text{C}_{20}\text{H}_{17}\text{O}_2 [\text{M} - \text{H}]^+$: 289.1229, Found: 289.1231.

IV. Product Derivatizations



(S)-3-Phenyl-3*H*-spiro[benzofuran-2,1'-cyclohexane]-2',5'-dien-4'-one (4). At 0 °C under N₂ atmosphere, to a 10-mL flask equipped with a stir bar were added a solution of **2a** (145.0 mg, 0.5 mmol, 1.0 equiv) in anhydrous CH₂Cl₂ (5 mL) and BBr₃ (1.0 M in CH₂Cl₂, 2.5 mL, 2.5 mmol, 5.0 equiv) dropwise. Then the reaction mixture was stirred at room temperature for 1 h before it was quenched with saturated aqueous NaHCO₃ solution. The aqueous layer was separated and extracted with CH₂Cl₂ (3 × 10 mL), then the combined organic layers were concentrated under reduced pressure. The residue was purified by flash column chromatography (eluent: hexanes/EtOAc = 3:1) to give the diol.

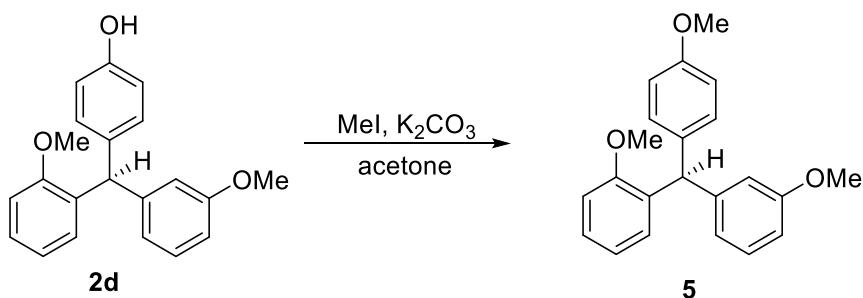
To a 10-mL flask equipped with a stir bar were added the above diol in benzene (5 mL) and (diacetoxyiodo)benzene (161.5 mg, 0.5 mmol, 1.0 equiv) in one portion at room temperature. The reaction mixture was stirred for 30 min and then concentrated under reduced pressure. The residue was purified by flash column chromatography (eluent: hexanes/EtOAc = 10:1) to give dienone **4** as a colorless oil in 52% yield (72.1 mg) and 95% ee over two steps.

¹H NMR (400 MHz, CDCl₃) δ 7.34 – 7.22 (m, 4H), 7.18 – 7.06 (m, 2H), 7.06 – 6.92 (m, 4H), 6.51 (dd, *J* = 10.3, 3.1 Hz, 1H), 6.28 (dd, *J* = 10.1, 2.0 Hz, 1H), 5.88 (dd, *J* = 10.3, 1.9 Hz, 1H), 4.78 (s, 1H).

¹³C NMR (101 MHz, CDCl₃) δ 185.1, 158.4, 147.1, 145.4, 136.6, 129.5, 128.7, 128.6, 128.5, 128.4, 128.4, 128.0, 126.1, 121.8, 110.9, 85.0, 57.8.

$[\alpha]_D^{25}$: -2.2 ($c = 1.0$, CHCl_3). 95% ee, HPLC analysis of the product: Daicel Chiraldpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 6.5 min (minor), 6.9 min (major).

HRMS (Cl⁺) Calcd for $\text{C}_{19}\text{H}_{15}\text{O}_2$ [M + H]⁺: 275.1072, Found: 275.1064.



(S)-2-((3-Hydroxyphenyl)(4-hydroxyphenyl)methyl)phenol (5). To a 10-mL flask equipped with a stir bar was added a solution of **2d** (96.7 mg, 0.3 mmol, 1.0 equiv) in acetone (3 mL). Then potassium carbonate (103.7 mg, 0.75 mmol, 2.5 equiv) and iodomethane (85.1 mg, 0.5 mmol, 2.0 equiv) were sequentially added. The reaction mixture was stirred under reflux for 10 h and cooled to room temperature. The suspension was filtered through celite and poured into a mixture of EtOAc (10 mL) and brine (10 mL), and washed with EtOAc (2 × 10 mL). The combined organic layers were concentrated under reduced pressure, and the residue was purified by flash column chromatography (eluent: hexanes/EtOAc = 15:1) to afford **5** as a colorless oil in 89% yield (89.2 mg) and 95% ee.

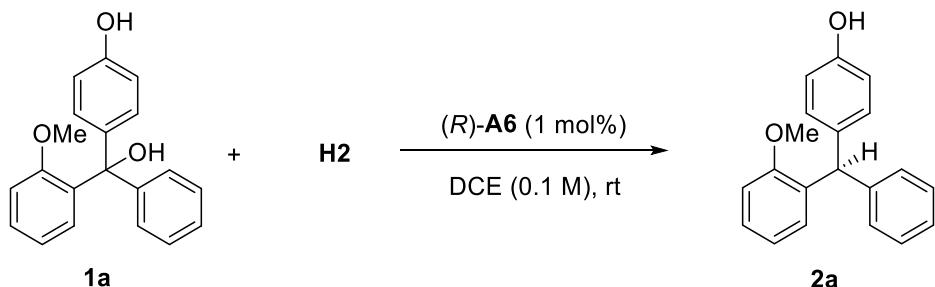
¹H NMR (300 MHz, CDCl_3) δ 7.26 – 7.16 (m, 2H), 7.09 – 6.98 (m, 2H), 6.95 – 6.87 (m, 3H), 6.87 – 6.79 (m, 2H), 6.76 (dd, $J = 8.2, 2.5$ Hz, 1H), 6.70 (dd, $J = 10.5, 4.8$ Hz, 2H), 5.88 (s, 1H), 3.80 (s, 3H), 3.75 (s, 3H), 3.74 (s, 3H).

$[\alpha]_D^{25}$: +4.2 ($c = 1.0$, CHCl_3). HPLC analysis of the product: Daicel Chiraldpak AD-H column; 10% *i*-PrOH in hexanes; 1.0 mL/min; retention times: 7.0 min (minor), 7.3 min (major).

This is a known compound. The characterization data are consistent with the literature report.²

V. Mechanistic Experiments

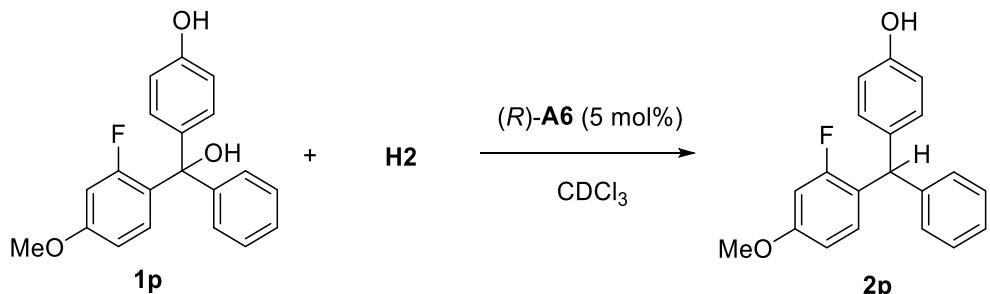
(1) Monitoring the ee values during the reaction process.



Time (h)	Ee of 1a (%)	Ee of 2a (%)
0.1	0	97
0.5	0	97
2.0	0	96
5.0	0	96
8.0	— ^a	96
24.0	— ^a	96

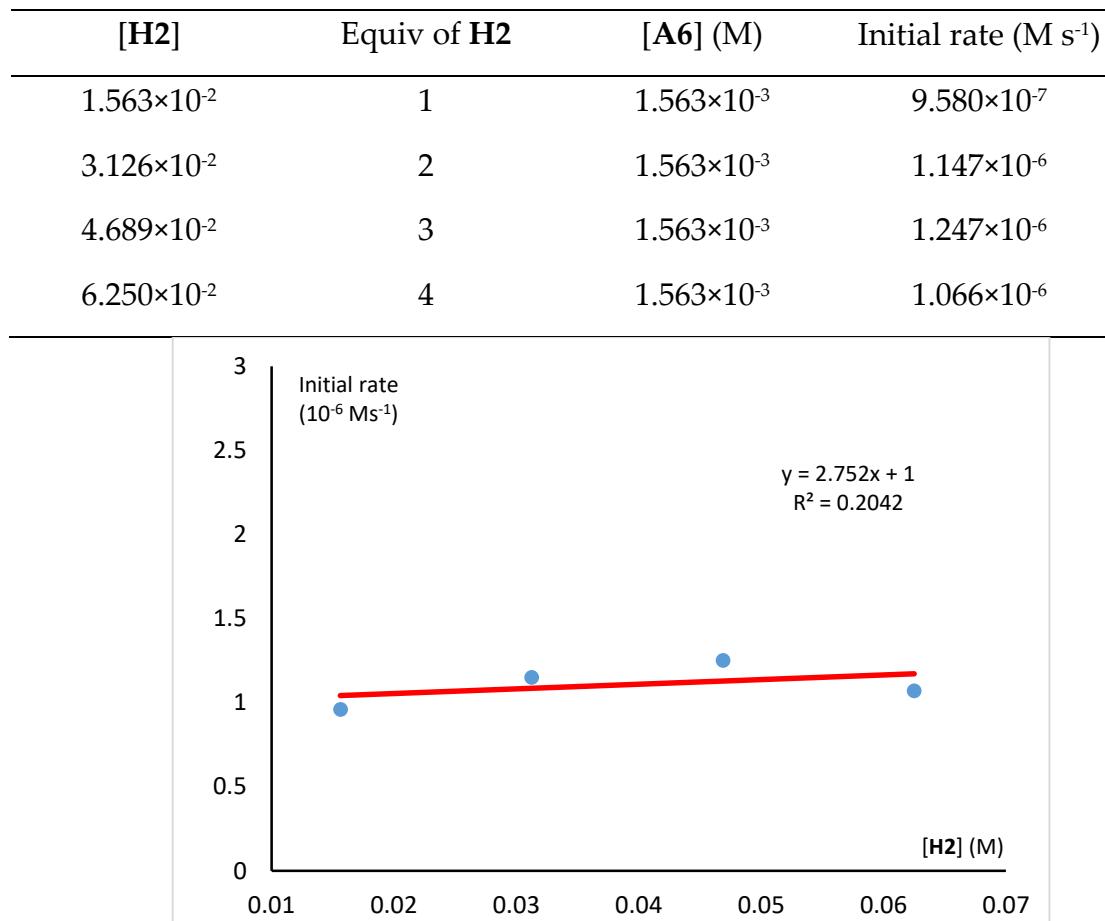
^a **1a** was consumed at this time point.

(2) Kinetic study: Reaction order in hydride H2.

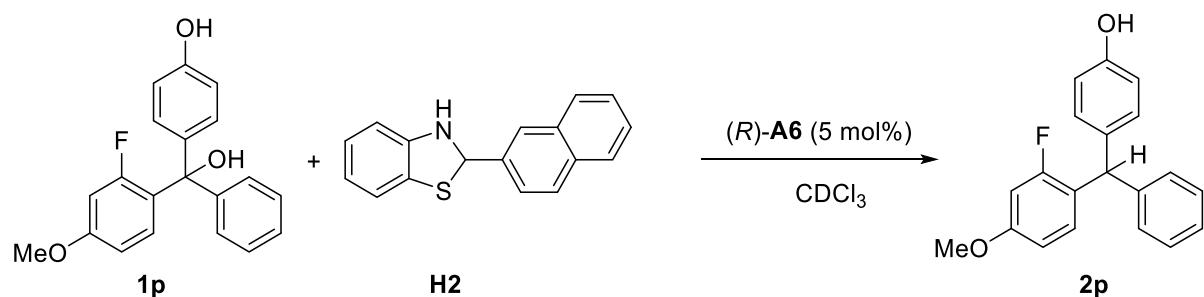


In four oven-dried NMR tubes, each was charged with a solution of substrate **1p** (8.1 mg, 0.0125 mmol, 1 equiv) in CDCl_3 (0.3 mL). In other four separate 4-mL vials, each was charged with a solution of (*R*)-**A6** (1.3 mg, 1.25 μmol) and the hydride source **H2** (6.6 mg, 1 equiv; 13.2 mg, 2 equiv; 19.8 mg, 3 equiv; 26.4 mg, 4 equiv) in CDCl_3 (1 mL). An aliquot of this solution (0.5 mL) was added to the above NMR tube containing **1p**. Then, the formation of the product was monitored by NMR. The concentration of

product **2p** was determined by ^{19}F NMR. A linear trend representing the first 10% conversion was fitted to determine the initial rates of the reaction. The data indicated zero-order kinetics in hydride **H2**.



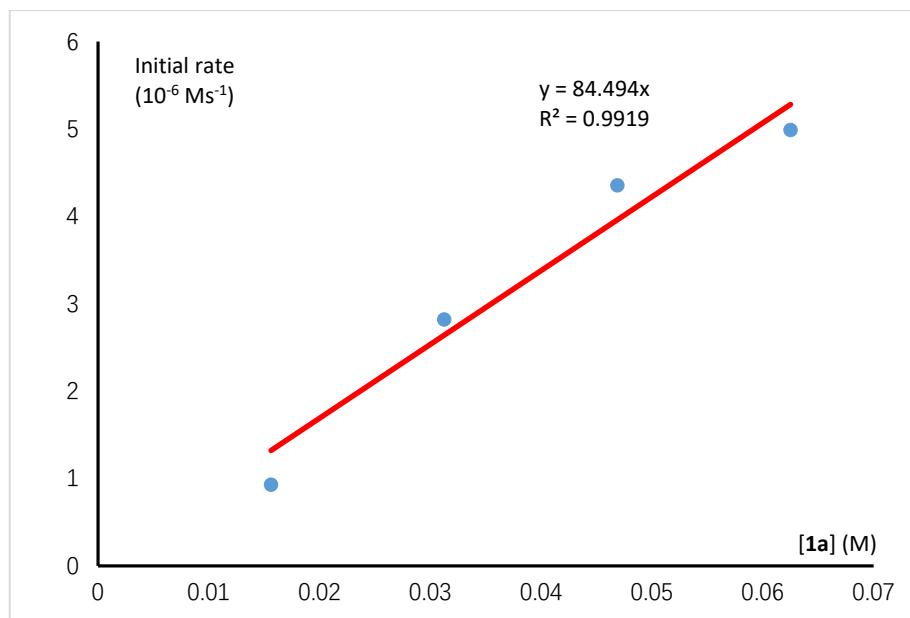
(3) Kinetic study: Reaction order in triarylmethanol.



In four oven-dried NMR tubes, each was charged with a solution of the hydride source **H2** (3.3 mg, 0.0125 mmol, 1 equiv) and catalyst **A6** (1.3 mg, 1.25 μmol) in CDCl_3 (0.3 mL). In other four separate 4-mL vials, each was charged with **1p** (16.2 mg, 1 equiv;

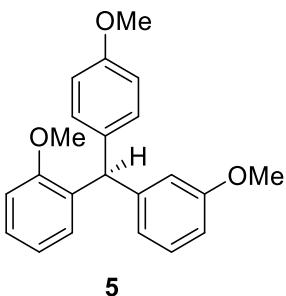
32.4 mg, 2 equiv; 48.6 mg, 3 equiv; 64.8 mg, 4 equiv) in CDCl_3 (1 mL). An aliquot of this solution (0.5 mL) was added to the above NMR tube. Then, the formation of the product was monitored by NMR. The concentration of product **2p** was determined by ^{19}F NMR. A linear trend representing the first 10% conversion was fitted to determine the initial rates of the reaction. The data indicated first-order kinetics in triarylmethanol.

[1p]	Equiv of 1p	[A6] (M)	Initial rate (M s^{-1})
1.563×10^{-2}	1	1.563×10^{-3}	9.304×10^{-7}
3.126×10^{-2}	2	1.563×10^{-3}	2.821×10^{-6}
4.689×10^{-2}	3	1.563×10^{-3}	4.357×10^{-6}
6.250×10^{-2}	4	1.563×10^{-3}	4.991×10^{-6}



VI. Determination of the Absolute Stereochemistry

The absolute stereochemistry of product derivative **5** was assigned to be *S* based on comparison of the optical rotation value with the literature value of the known compound.² The absolute configuration of other products was assumed by analogy.

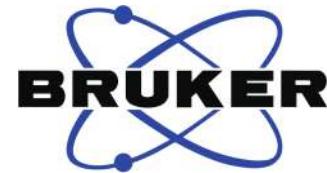
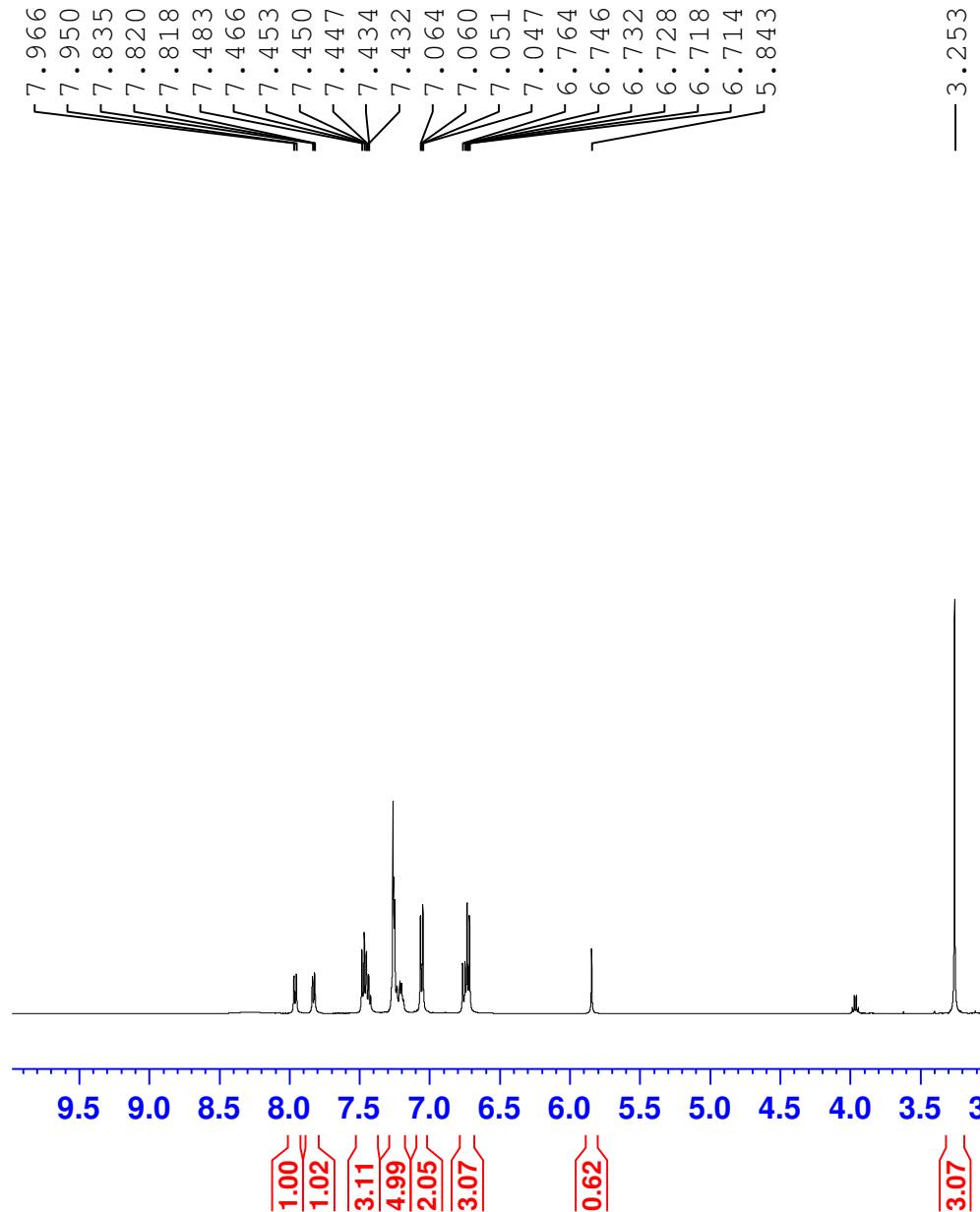


(*S*)-1-Methoxy-2-((3-methoxyphenyl)(4-methoxyphenyl)methyl)benzene. The literature optical rotation value for (*S*)-**5** in 95% ee was reported to be $[\alpha]_D^{25}$: +0.15 ($c = 1.44$, CHCl_3).² The measured value of our product in 95% ee is $[\alpha]_D^{25}$: +4.2 ($c = 1.0$, CHCl_3).

VII. References

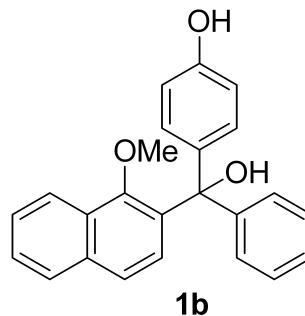
- (1) X. Li, M. Duan, Z. Deng, Q. Shao, M. Chen, G. Zhu, K. N. Houk and J. Sun, *Nat. Catal.* 2020, **3**, 1010–1019.
- (2) Y. Huang and T. Hayashi, *J. Am. Chem. Soc.* 2015, **137**, 7556–7559.

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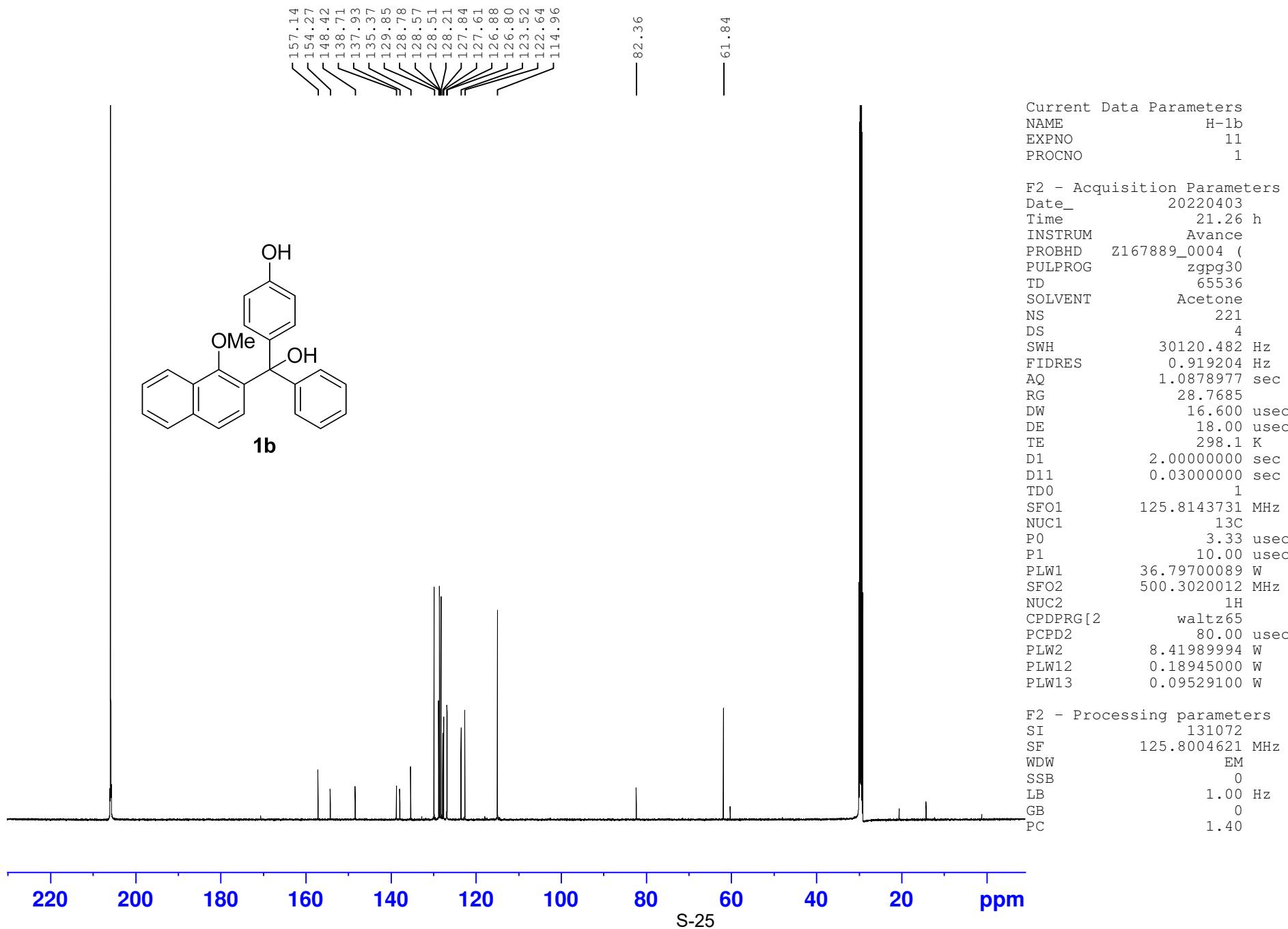
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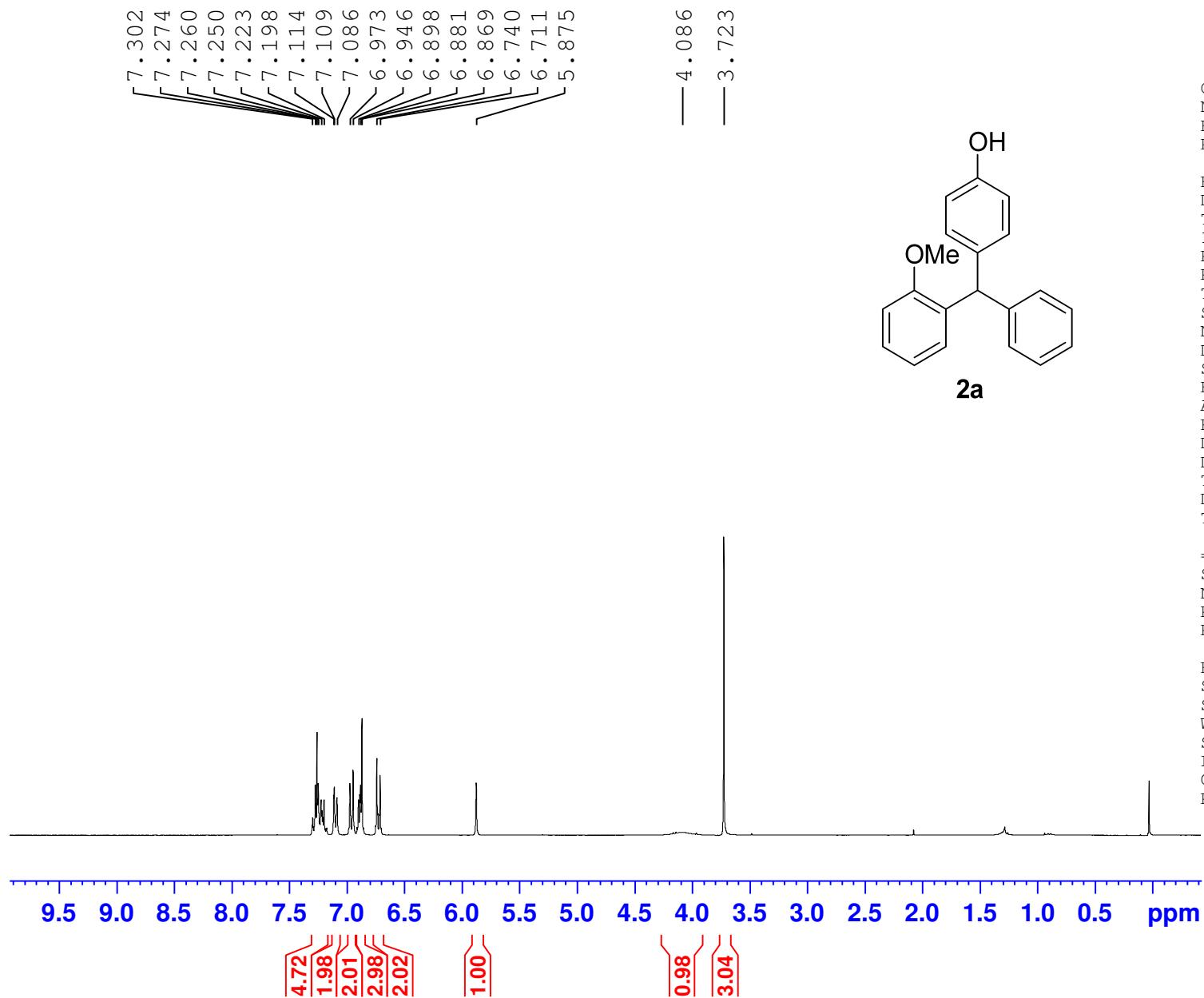


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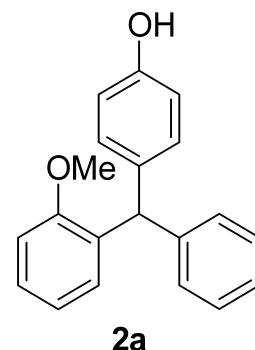
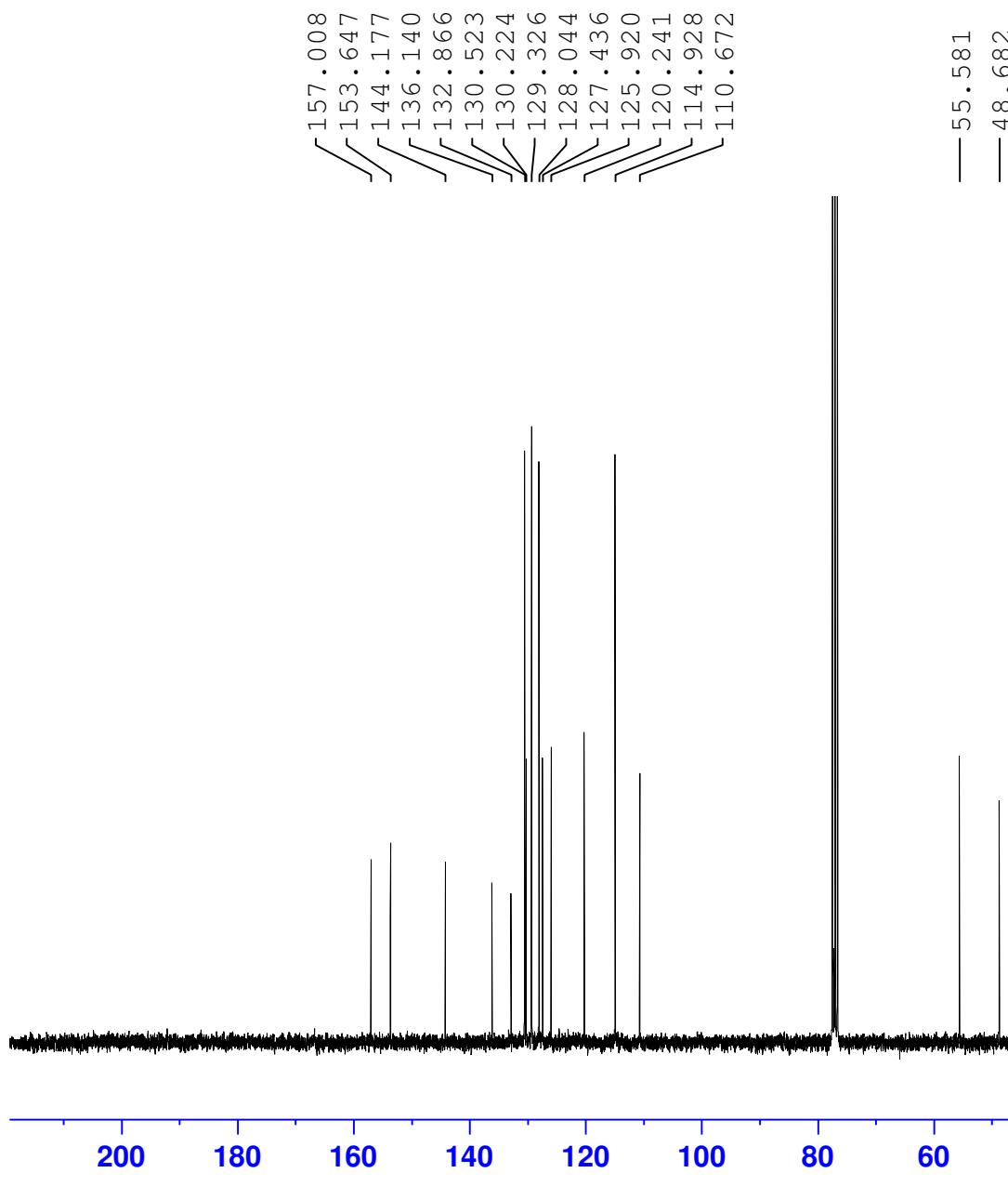
H-1b-c



3sjwei 2765 zy-3-10 1h cdcl3



3sjwei 2766 zy-3-10 13c cdcl3



Current Data Parameters
NAME ZY-3-10-c-fr
EXPNO 2766
PROCNO 1

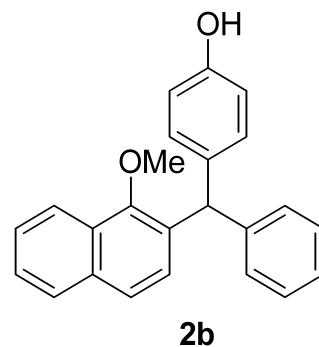
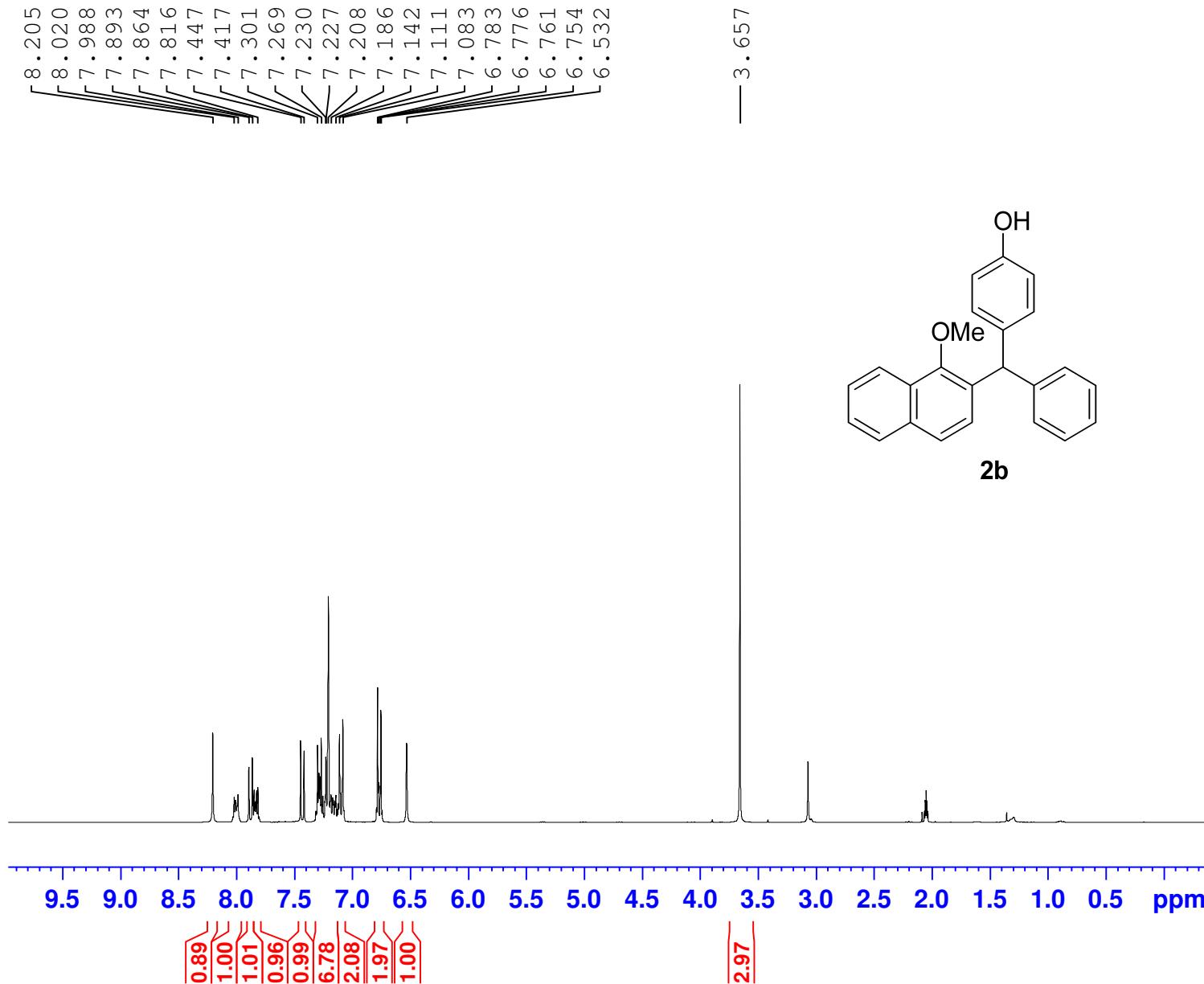
F2 - Acquisition Parameters
Date_ 20210329
Time 22.20
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 463
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 296.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 ======
SFO1 75.4752949 MHz
NUC1 13C
P1 9.50 usec
PLW1 34.20000076 W

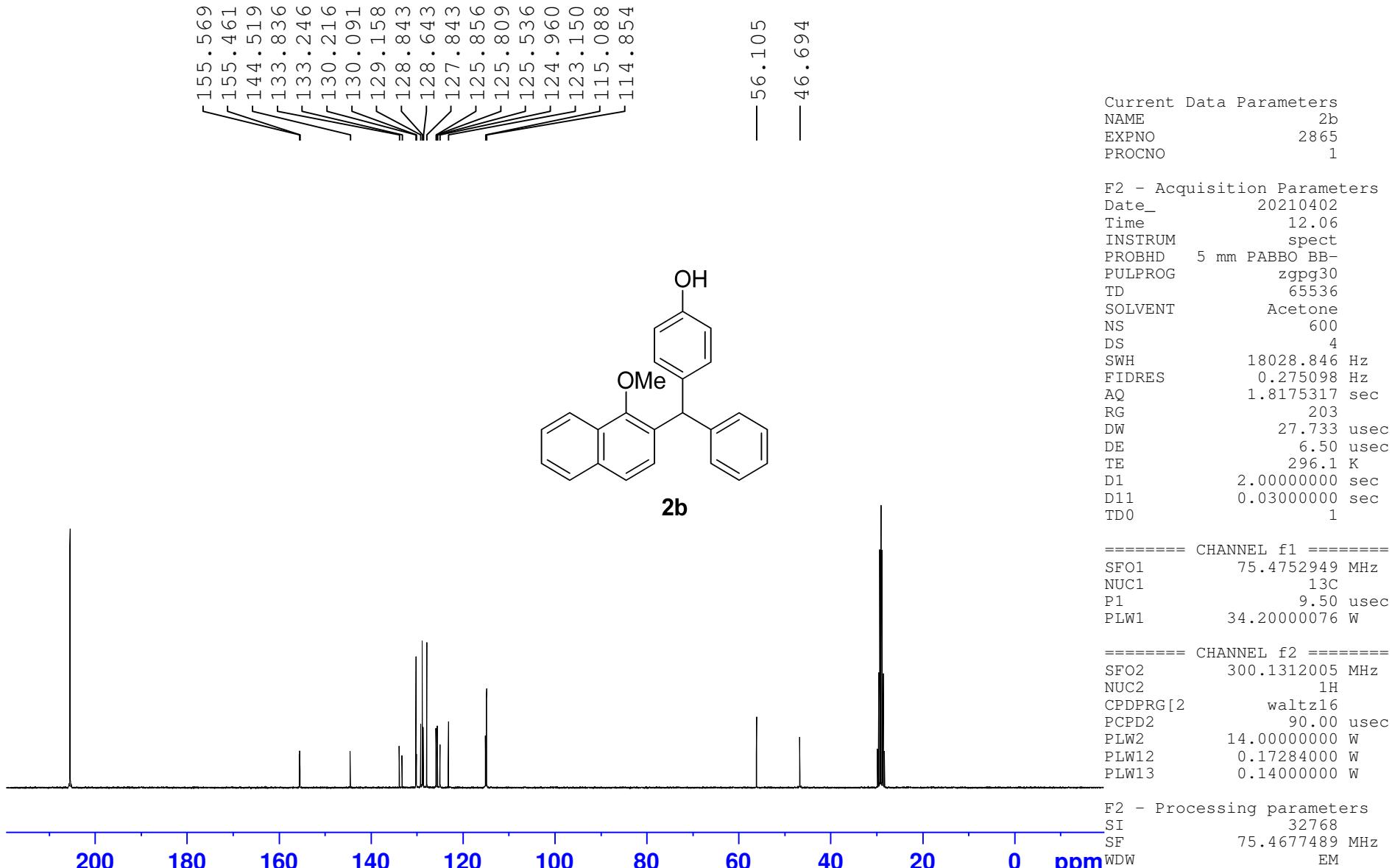
===== CHANNEL f2 ======
SFO2 300.1312005 MHz
NUC2 1H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 14.00000000 W
PLW12 0.17284000 W
PLW13 0.14000000 W

F2 - Processing parameters
SI 32768
SF 75.4677537 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

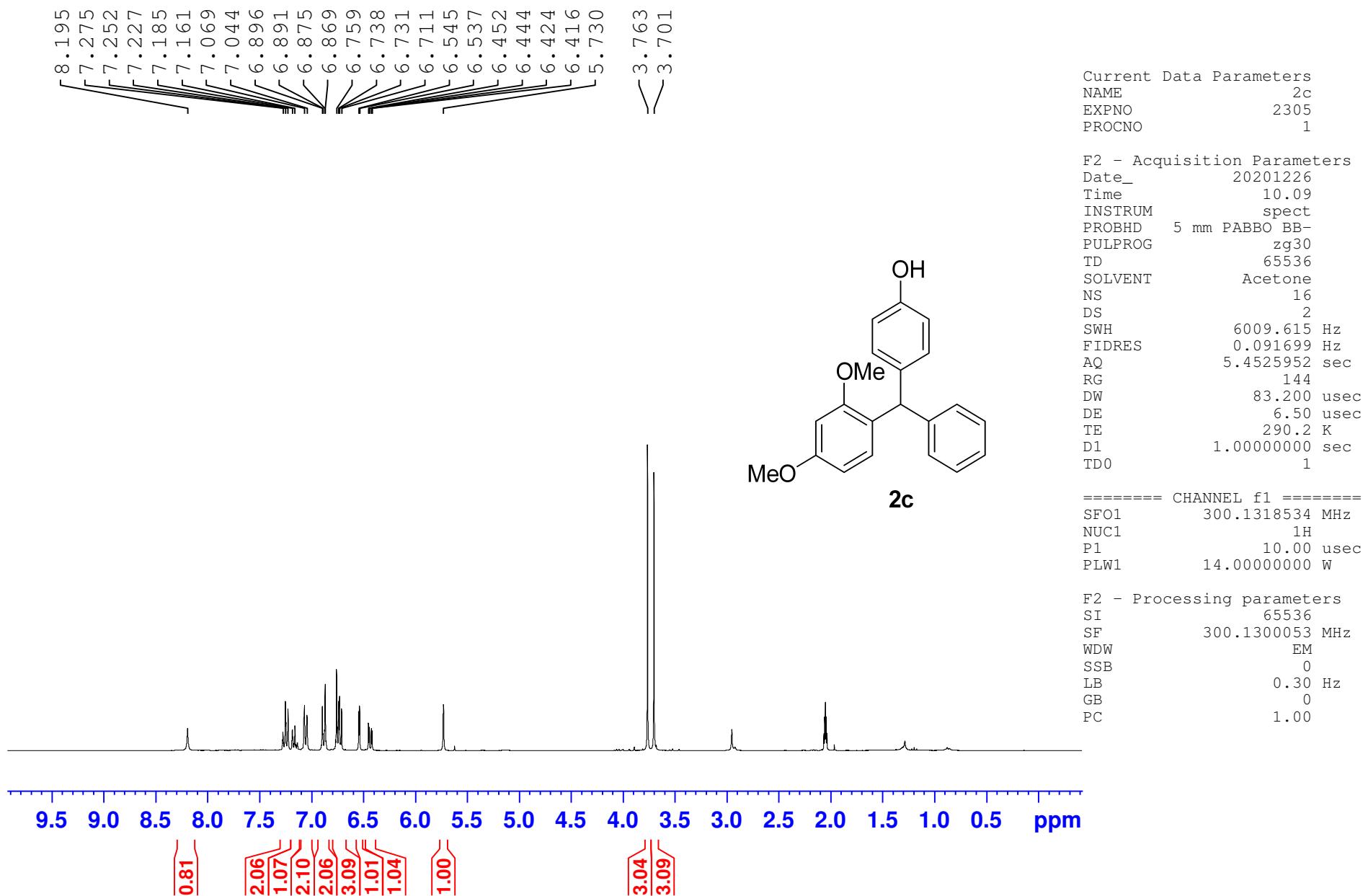
3sjwei 2851 zy-3-16 1h acetone



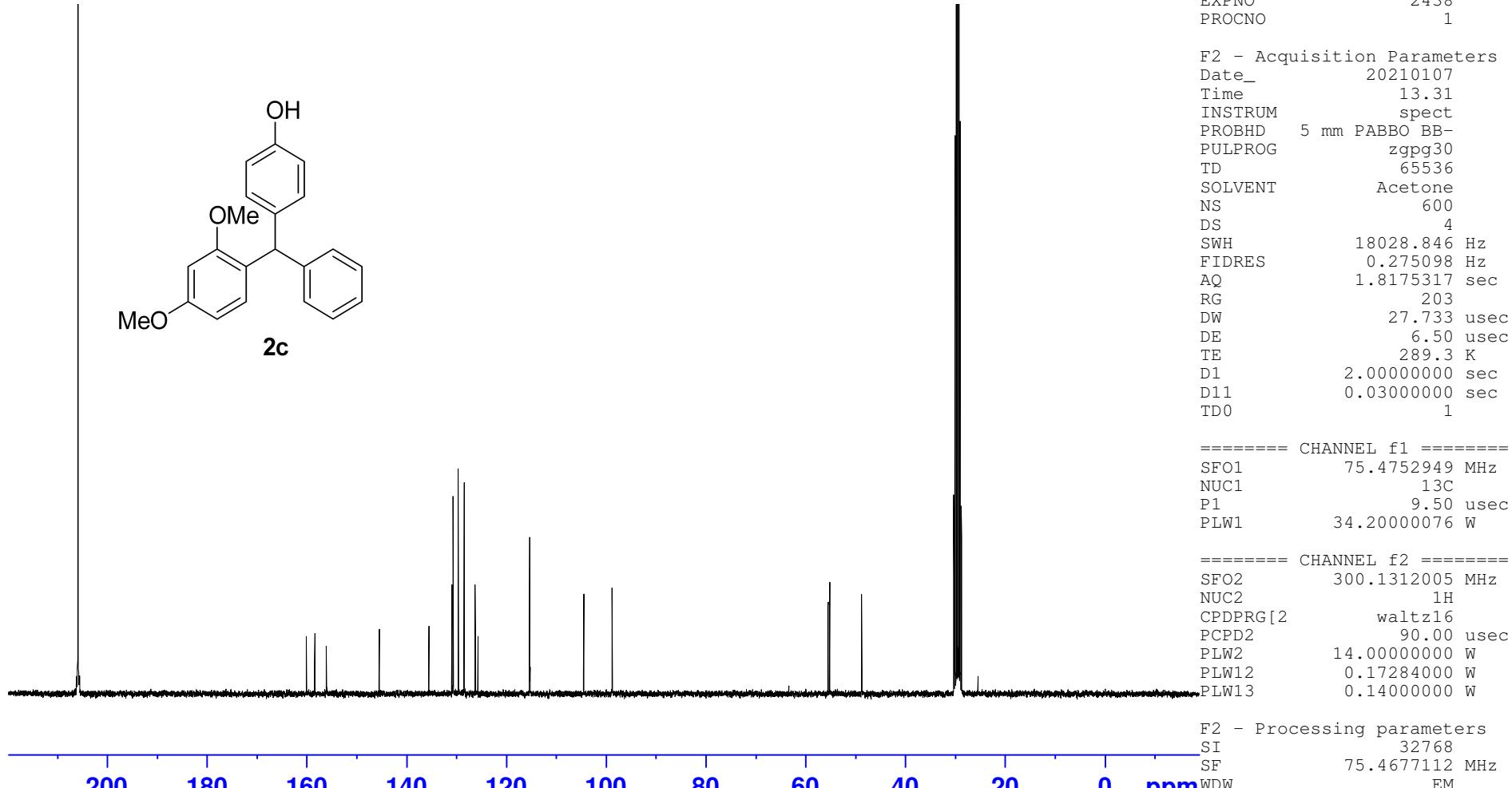
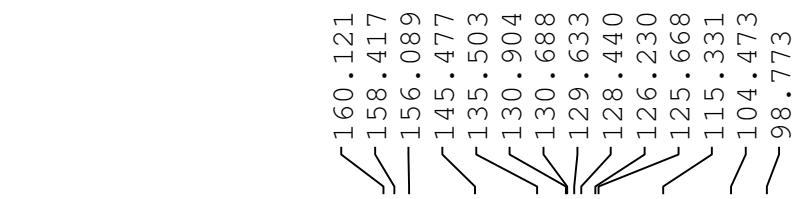
3sjwei 2865 zy-3-16 13c acetone



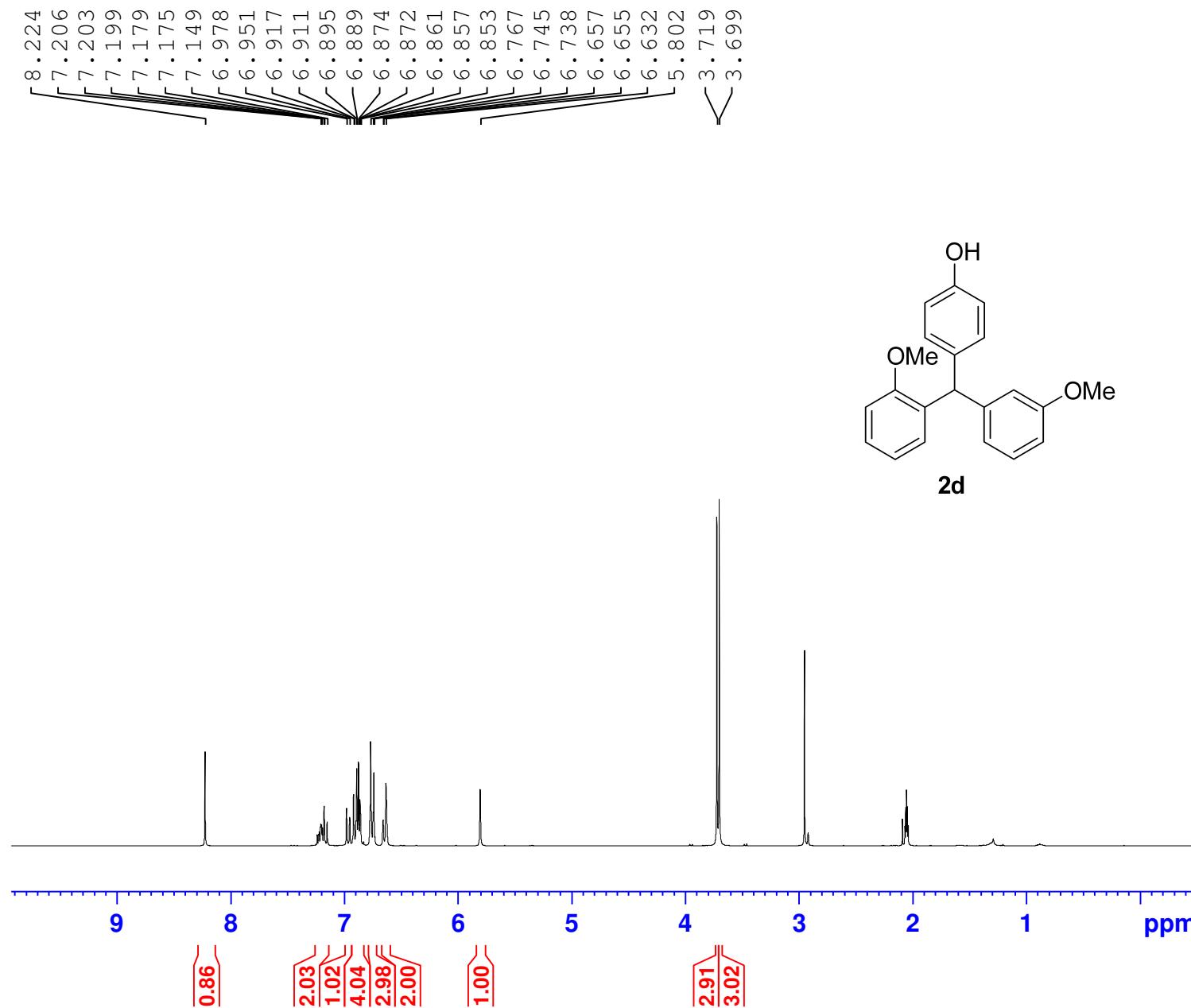
3sjwei 2305 zy-3-14f 1h acetone

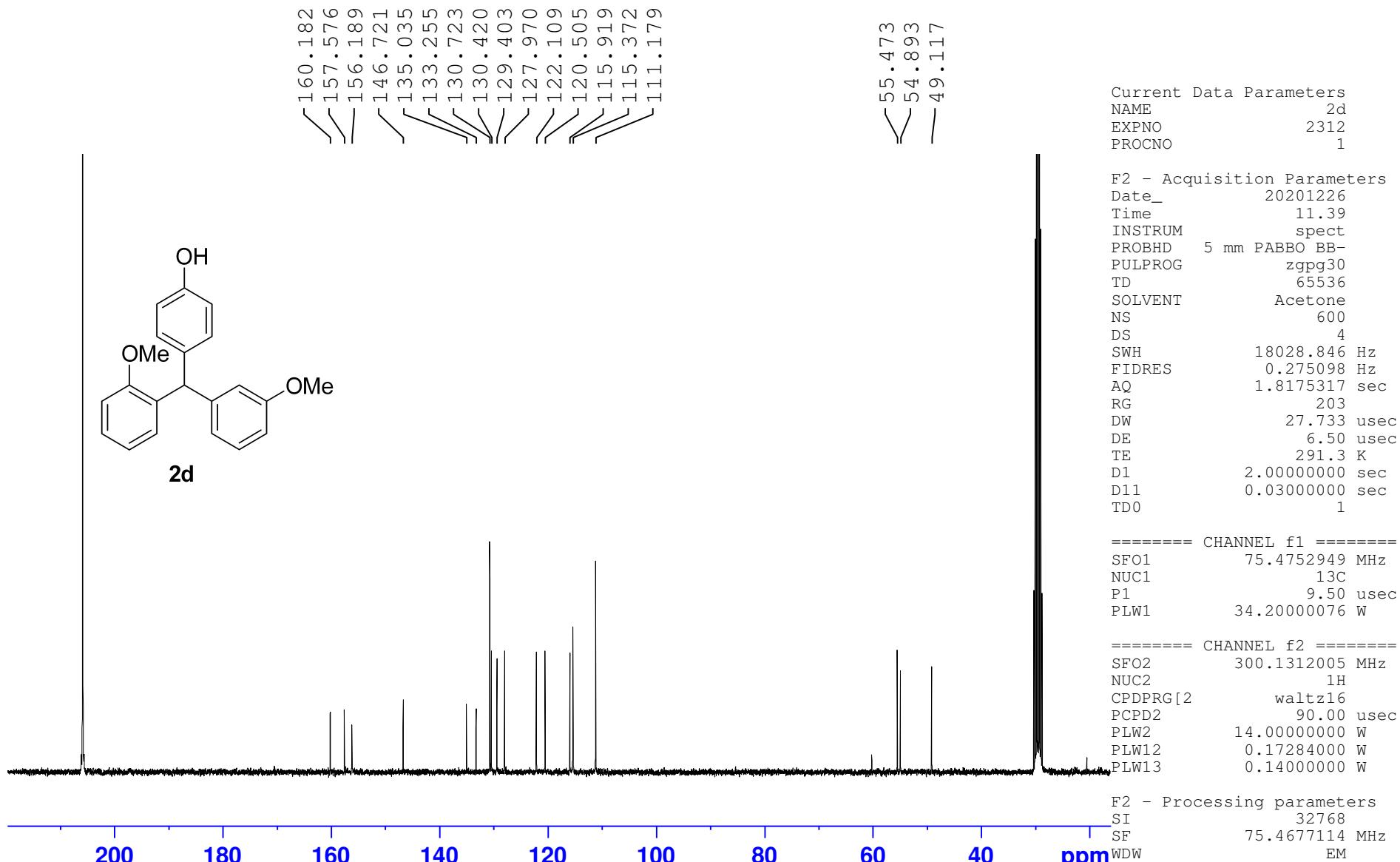


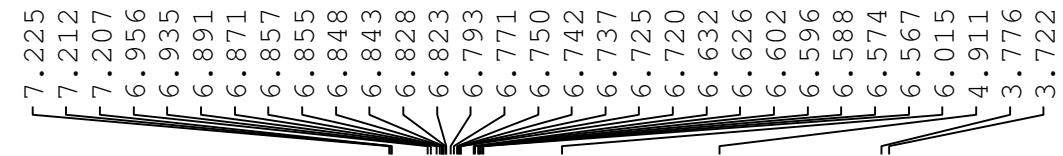
3sjwei 2438 zy-3-14f 13c acetone



3sjwei 2585 zy-3-11c 1h acetone





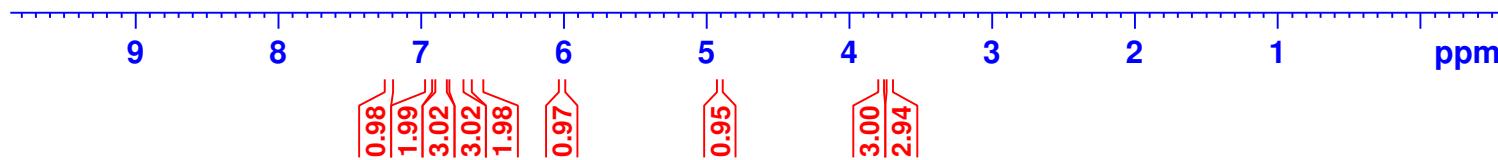
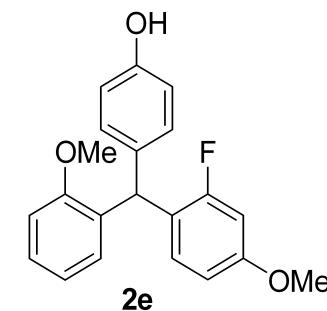


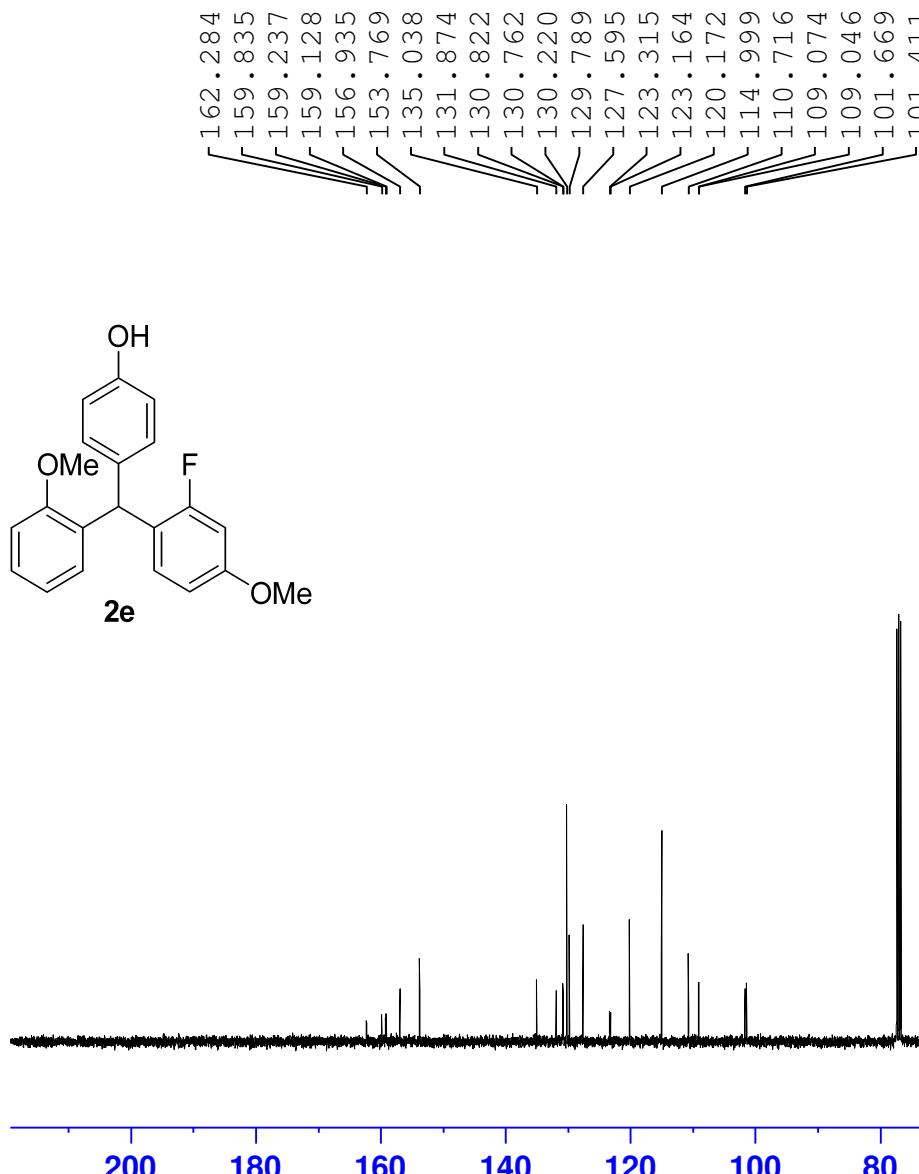
Current Data Parameters
 NAME 2eNMR
 EXPNO 1
 PROCNO 1

F2 - Acquisition Parameters
 Date_ 20201210
 Time 19.40
 INSTRUM spect
 PROBHD 5 mm PABBO BB/
 PULPROG zg30
 TD 65536
 SOLVENT CDCl3
 NS 14
 DS 0
 SWH 8012.820 Hz
 FIDRES 0.122266 Hz
 AQ 4.0894465 sec
 RG 45.67
 DW 62.400 usec
 DE 6.50 usec
 TE 296.7 K
 D1 1.0000000 sec
 TD0 1

===== CHANNEL f1 ======
 SFO1 400.1324710 MHz
 NUC1 1H
 P1 14.50 usec
 PLW1 11.99499989 W

F2 - Processing parameters
 SI 65536
 SF 400.1300101 MHz
 WDW EM
 SSB 0
 LB 0.30 Hz
 GB 0
 PC 1.00





162.284
159.835
159.237
159.128
156.935
153.769
135.038
131.874
130.822
130.762
130.220
129.789
127.595
123.315
123.164
120.172
114.999
110.716
109.074
109.046
101.669
101.411

55.608
55.448
41.362
41.336

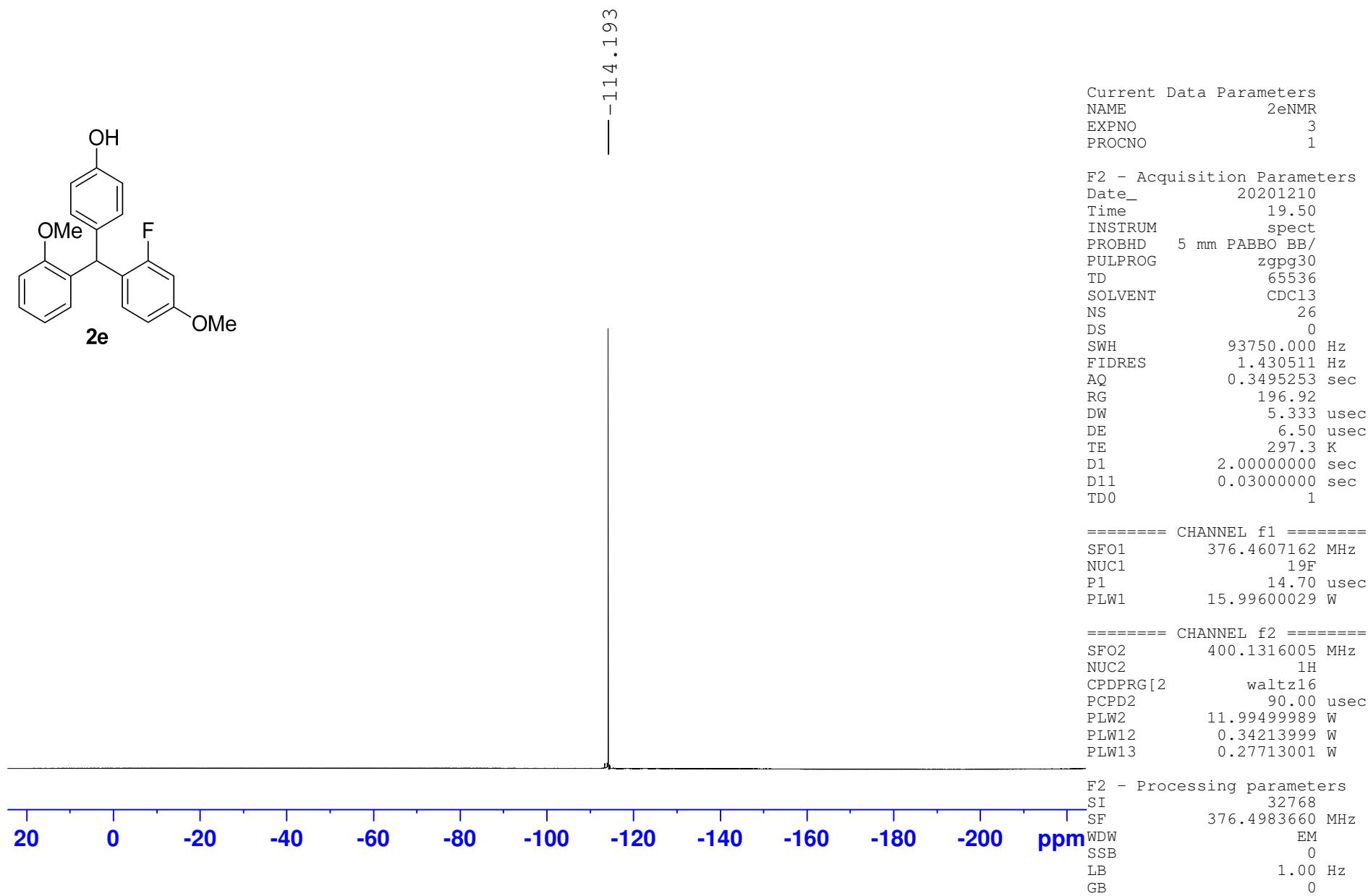
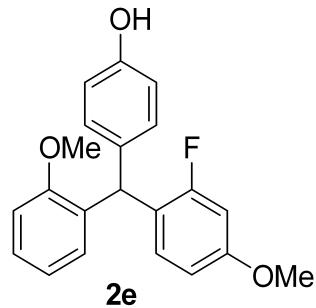
F2 - Processing parameters
SI 32768
SF 100.6127751 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

Current Data Parameters
NAME 2eNMR
EXPNO 2
PROCNO 1

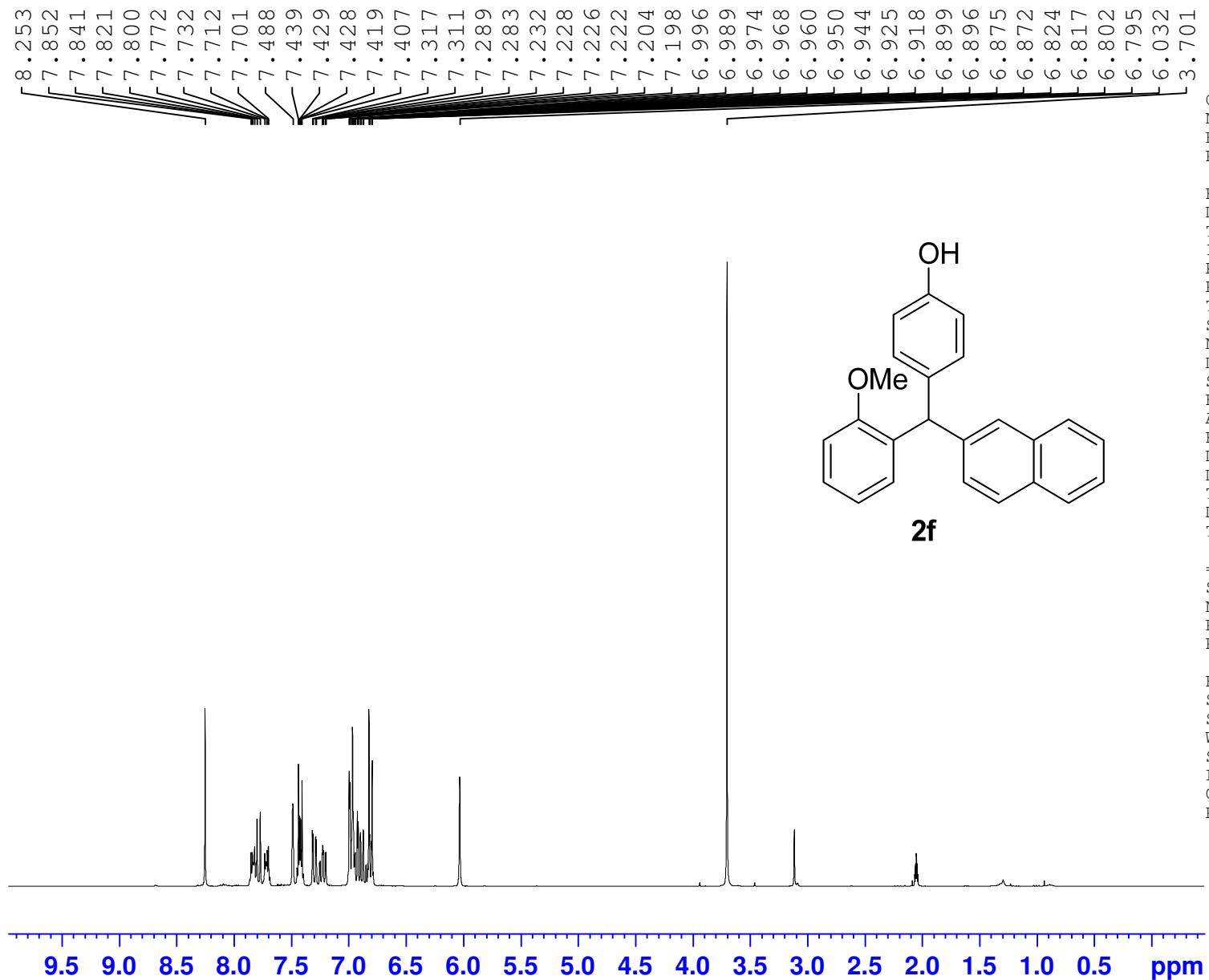
F2 - Acquisition Parameters
Date_ 20201210
Time 19.43
INSTRUM spect
PROBHD 5 mm PABBO BB/
PULPROG zgpg30
TD 65536
SOLVENT CDCl₃
NS 44
DS 0
SWH 24038.461 Hz
FIDRES 0.366798 Hz
AQ 1.3631488 sec
RG 196.92
DW 20.800 usec
DE 6.50 usec
TE 297.3 K
D1 2.0000000 sec
D11 0.0300000 sec
TD0 1

===== CHANNEL f1 ======
SFO1 100.6228298 MHz
NUC1 13C
P1 9.70 usec
PLW1 46.98899841 W

===== CHANNEL f2 ======
SFO2 400.1316005 MHz
NUC2 1H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 11.99499989 W
PLW12 0.34213999 W
PLW13 0.27713001 W



3sjwei 4273 zy-3-22 1h acetone

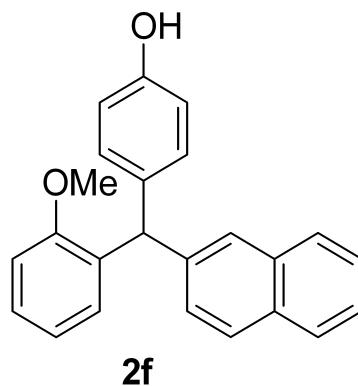


F2 - Acquisition Parameters
Date_ 20210624
Time 12.13
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT Acetone
NS 16
DS 2
SWH 6009.615 Hz
FIDRES 0.091699 Hz
AQ 5.4525952 sec
RG 64
DW 83.200 usec
DE 6.50 usec
TE 296.1 K
D1 1.00000000 sec
TD0 1

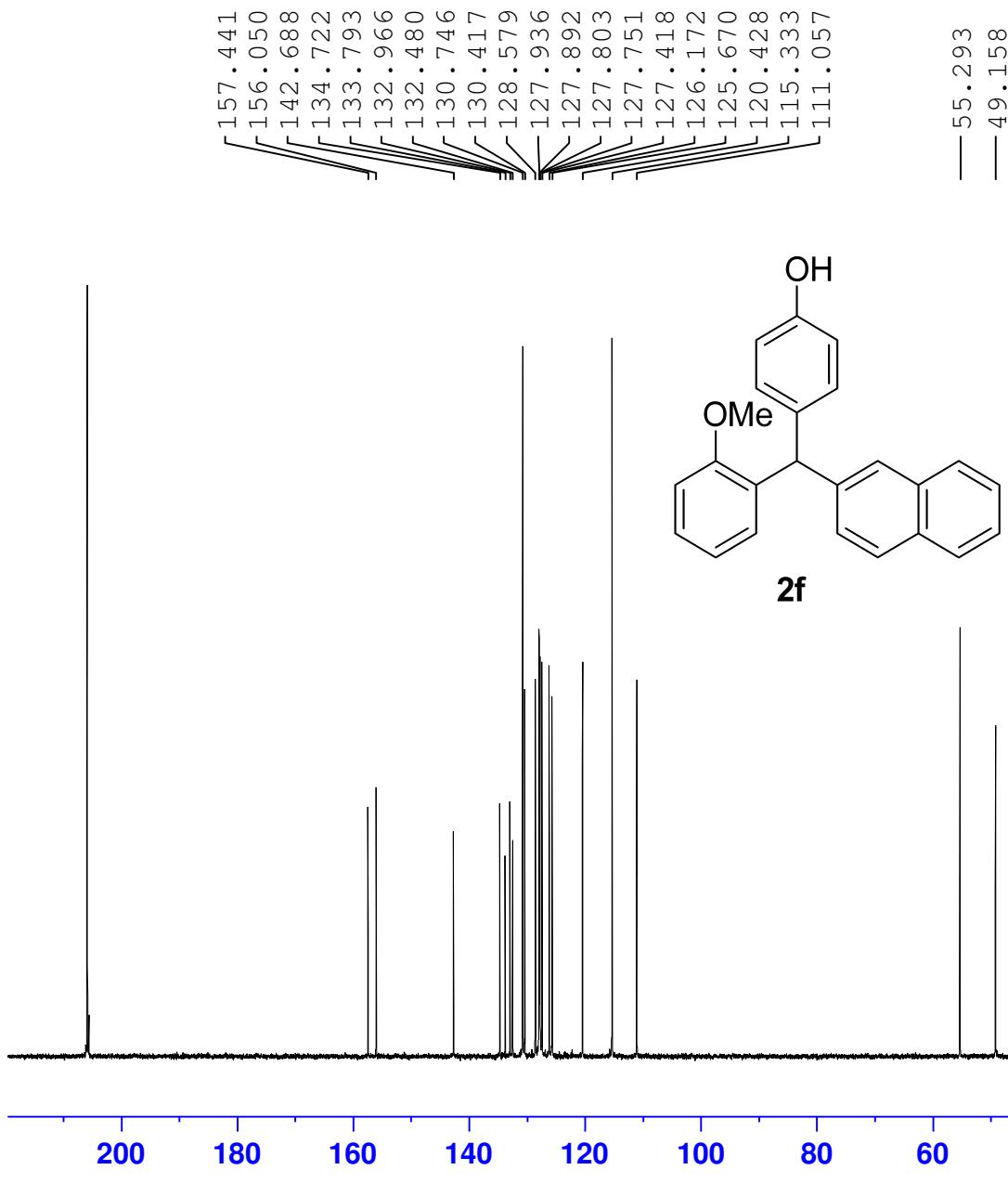
===== CHANNEL f1 ======

SFO1 300.1318534 MHz
NUC1 1H
P1 10.00 usec
PLW1 14.00000000 W

F2 - Processing parameters
SI 65536
SF 300.1300050 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



3sjwei 2814 zy-3-22 13c acetone



Current Data Parameters
NAME ZY-3-22-c-fr
EXPNO 2814
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210330
Time 22.03
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT Acetone
NS 600
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 296.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 ======

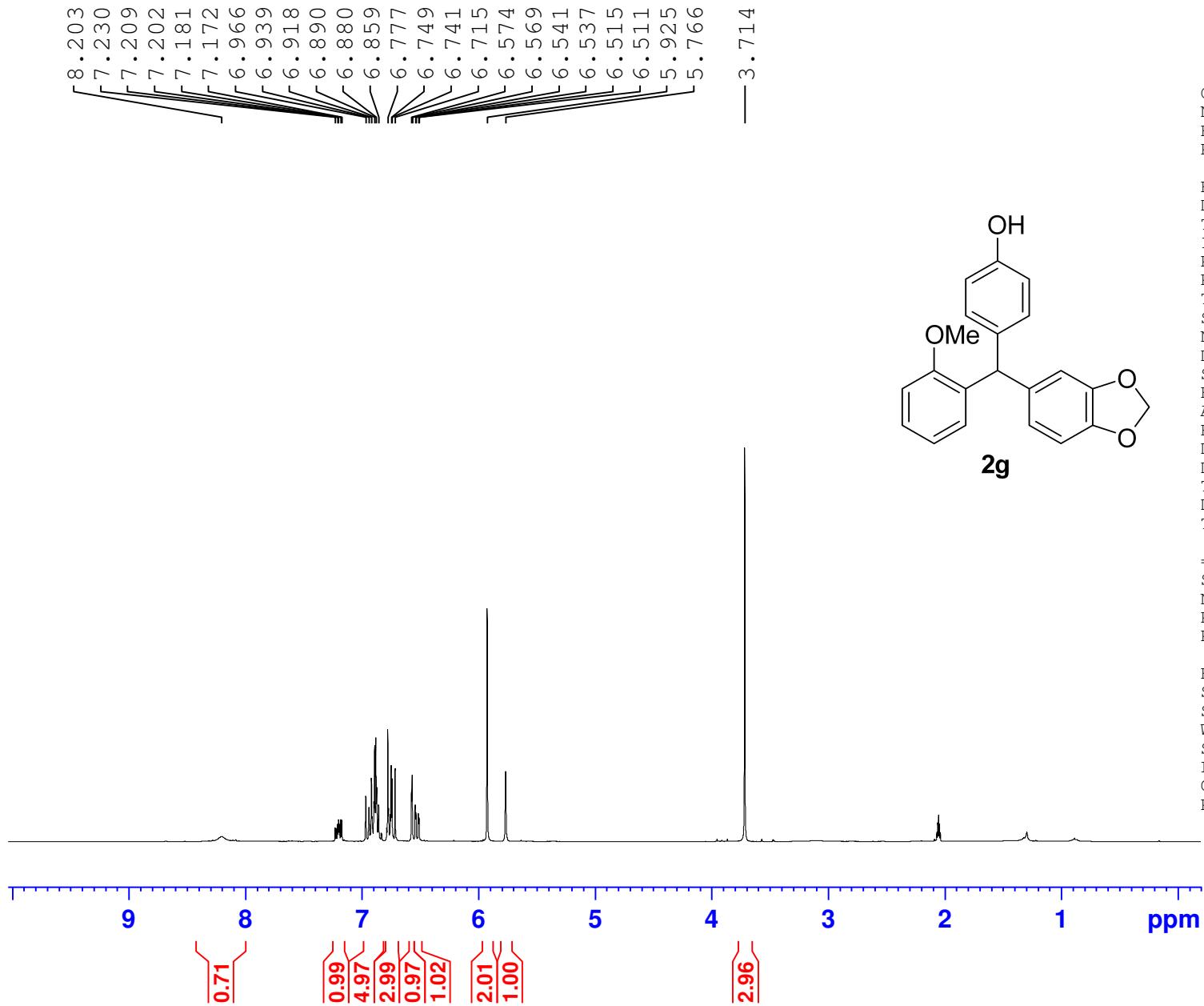
SFO1 75.4752949 MHz
NUC1 ¹³C
P1 9.50 usec
PLW1 34.20000076 W

===== CHANNEL f2 ======

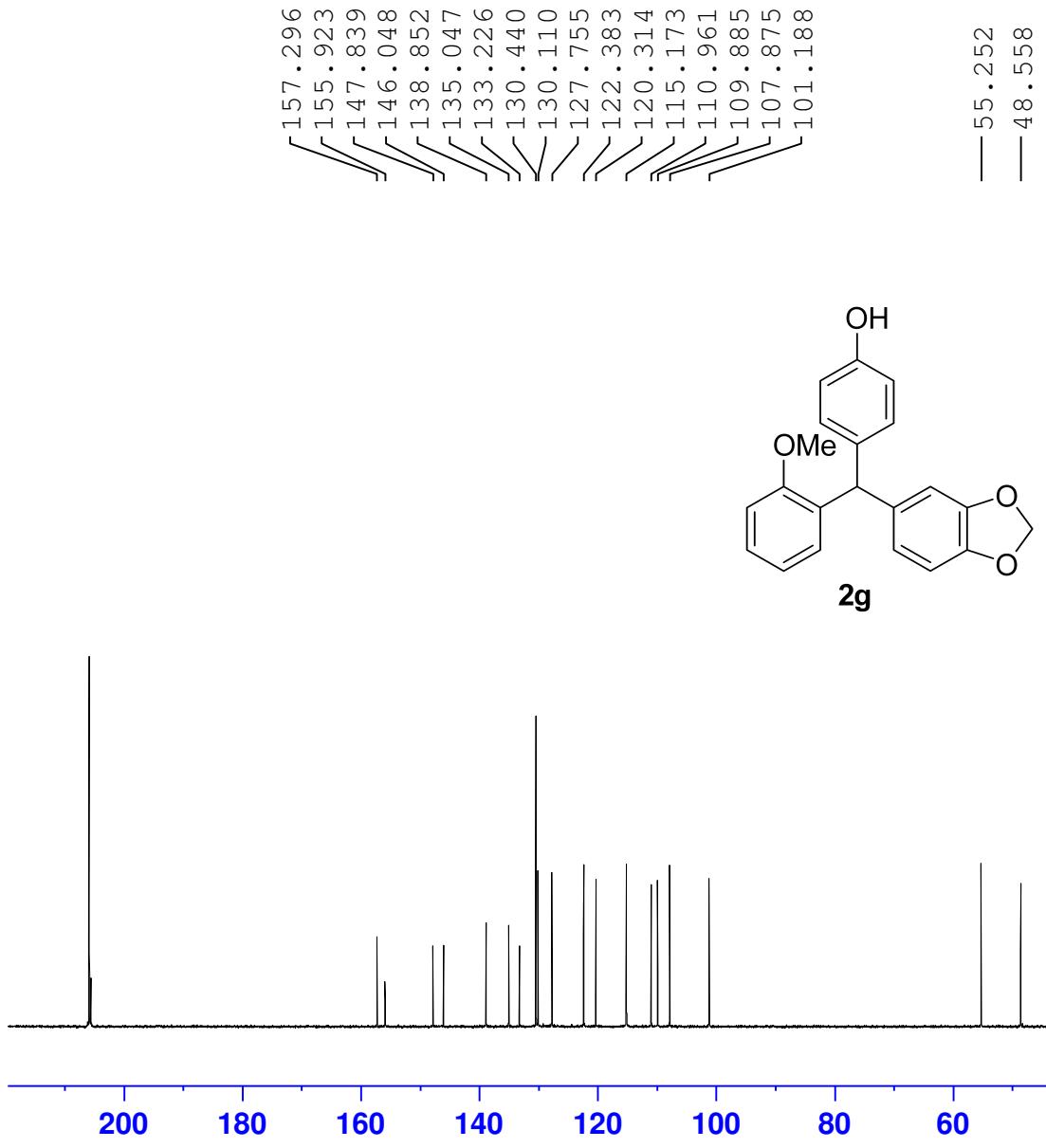
SFO2 300.1312005 MHz
NUC2 ¹H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 14.00000000 W
PLW12 0.17284000 W
PLW13 0.14000000 W

F2 - Processing parameters
SI 32768
SF 75.4677309 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

3sjwei 4272 zy-3-17a 1h acetone



3sjwei 2743 zy-3-17a 13c acetone



Current Data Parameters
NAME 2g
EXPNO 2743
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210318
Time 11.42
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT Acetone
NS 600
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 292.5 K
D1 2.0000000 sec
D11 0.0300000 sec
TD0 1

===== CHANNEL f1 ======

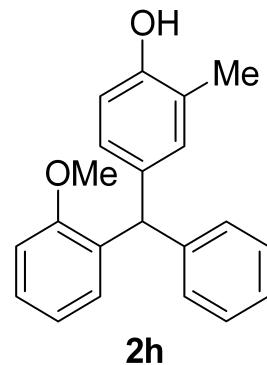
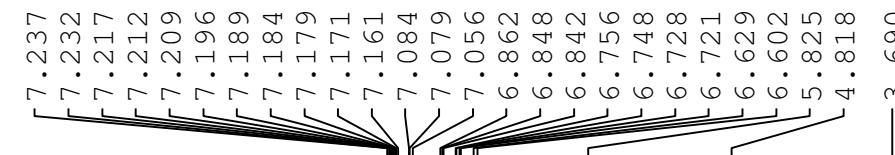
SFO1 75.4752949 MHz
NUC1 13C
P1 9.50 usec
PLW1 34.20000076 W

===== CHANNEL f2 ======

SFO2 300.1312005 MHz
NUC2 1H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 14.00000000 W
PLW12 0.17284000 W
PLW13 0.14000000 W

F2 - Processing parameters
SI 32768
SF 75.4677302 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

3sjwei 4371 zy-3-20d 1h cdcl3



2h



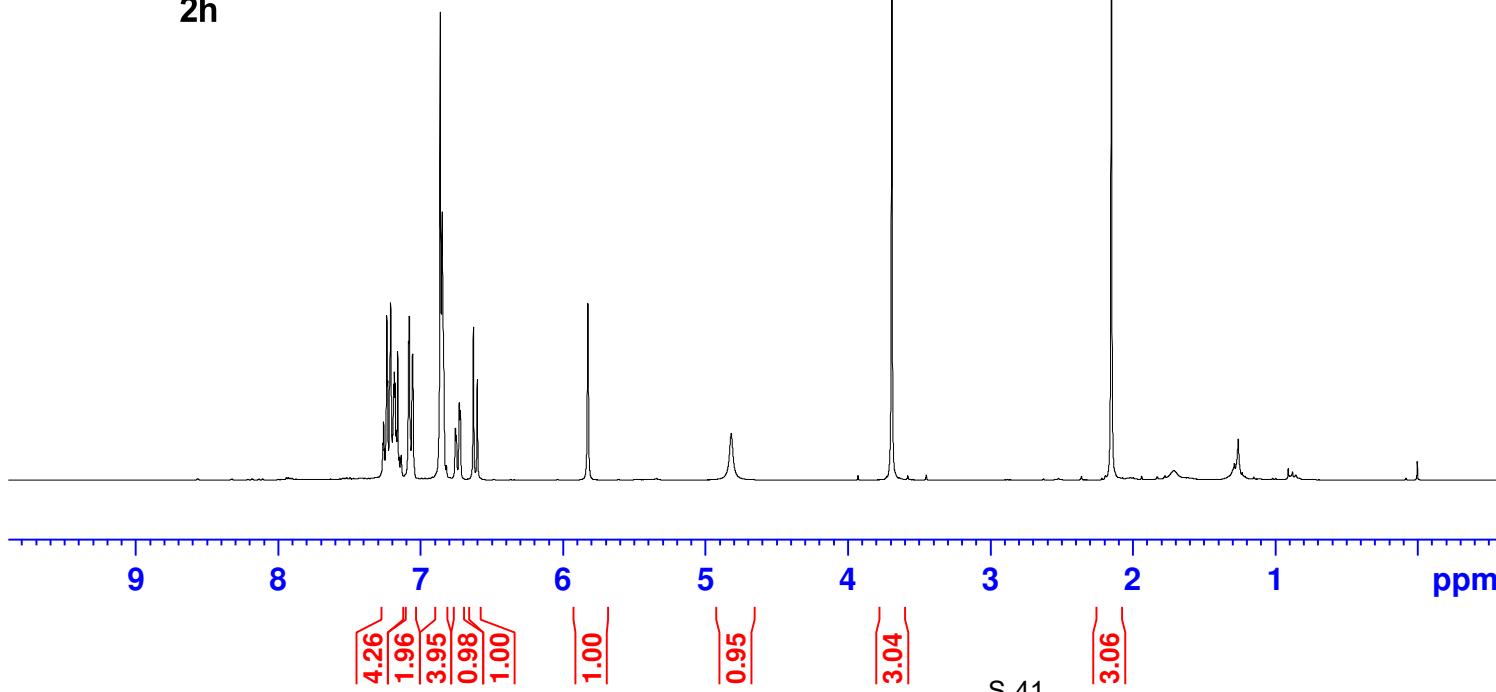
Current Data Parameters
NAME ZY-3-20D-h-fr
EXPNO 4371
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210701
Time 9.58
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 6009.615 Hz
FIDRES 0.091699 Hz
AQ 5.4525952 sec
RG 80.6
DW 83.200 usec
DE 6.50 usec
TE 296.2 K
D1 1.00000000 sec
TD0 1

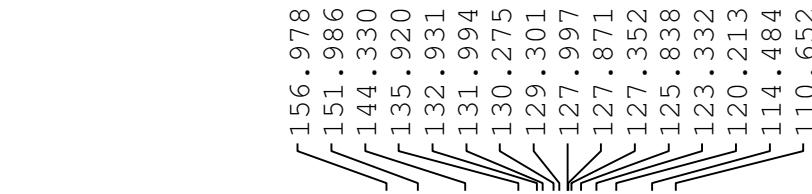
===== CHANNEL f1 ======

SFO1 300.1318534 MHz
NUC1 1H
P1 10.00 usec
PLW1 14.00000000 W

F2 - Processing parameters
SI 65536
SF 300.1300225 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



3sjwei 4394zy-3-20d 13c cdcl3



— 55.586
— 48.655

— 15.825

Current Data Parameters
NAME ZY-3-20D-c-fr
EXPNO 4394
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210702
Time 11.21
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 1024
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 296.2 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

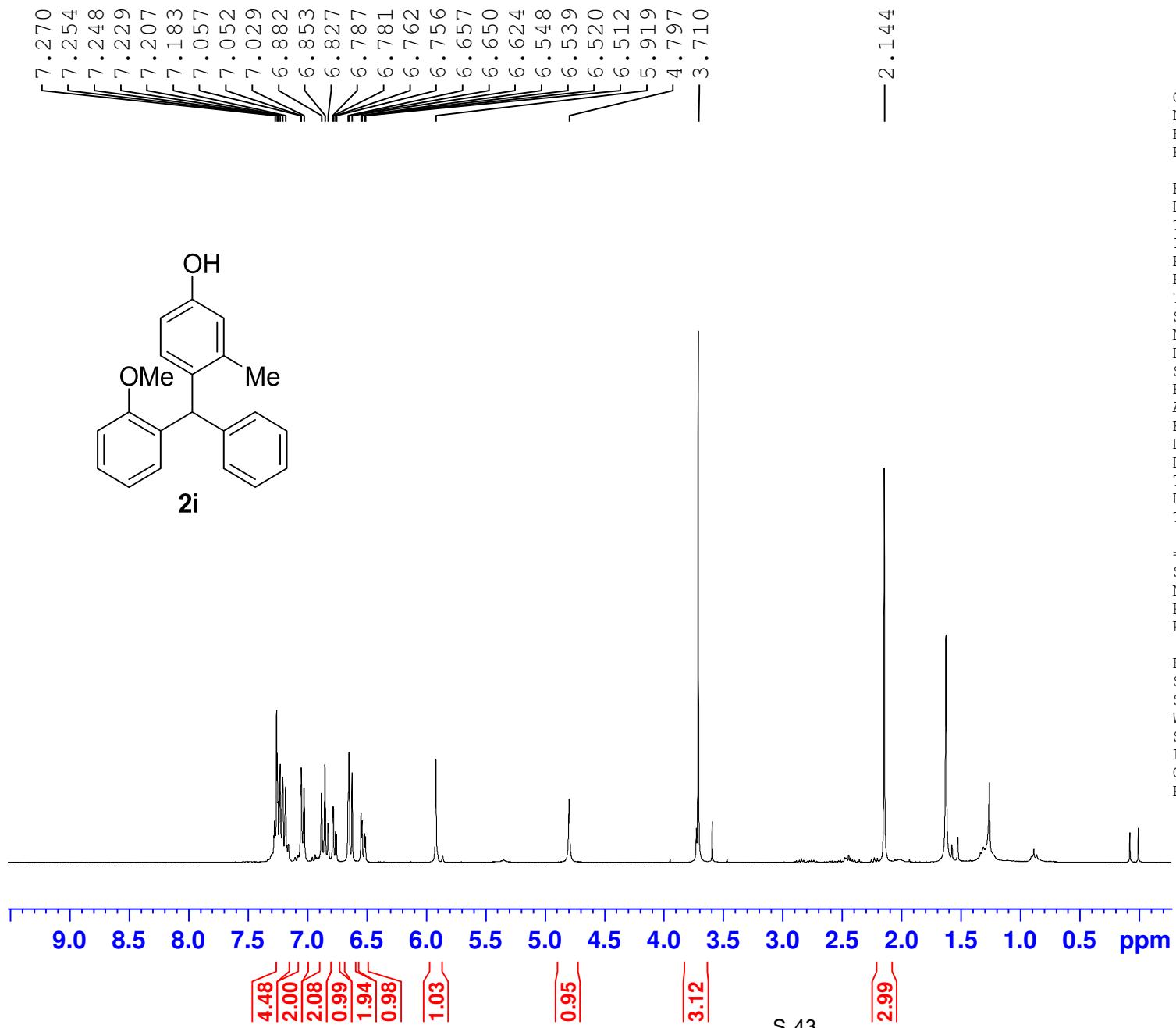
===== CHANNEL f1 ======
SFO1 75.4752949 MHz
NUC1 13C
P1 9.50 usec
PLW1 34.20000076 W

===== CHANNEL f2 ======
SFO2 300.1312005 MHz
NUC2 1H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 14.00000000 W
PLW12 0.17284000 W
PLW13 0.14000000 W

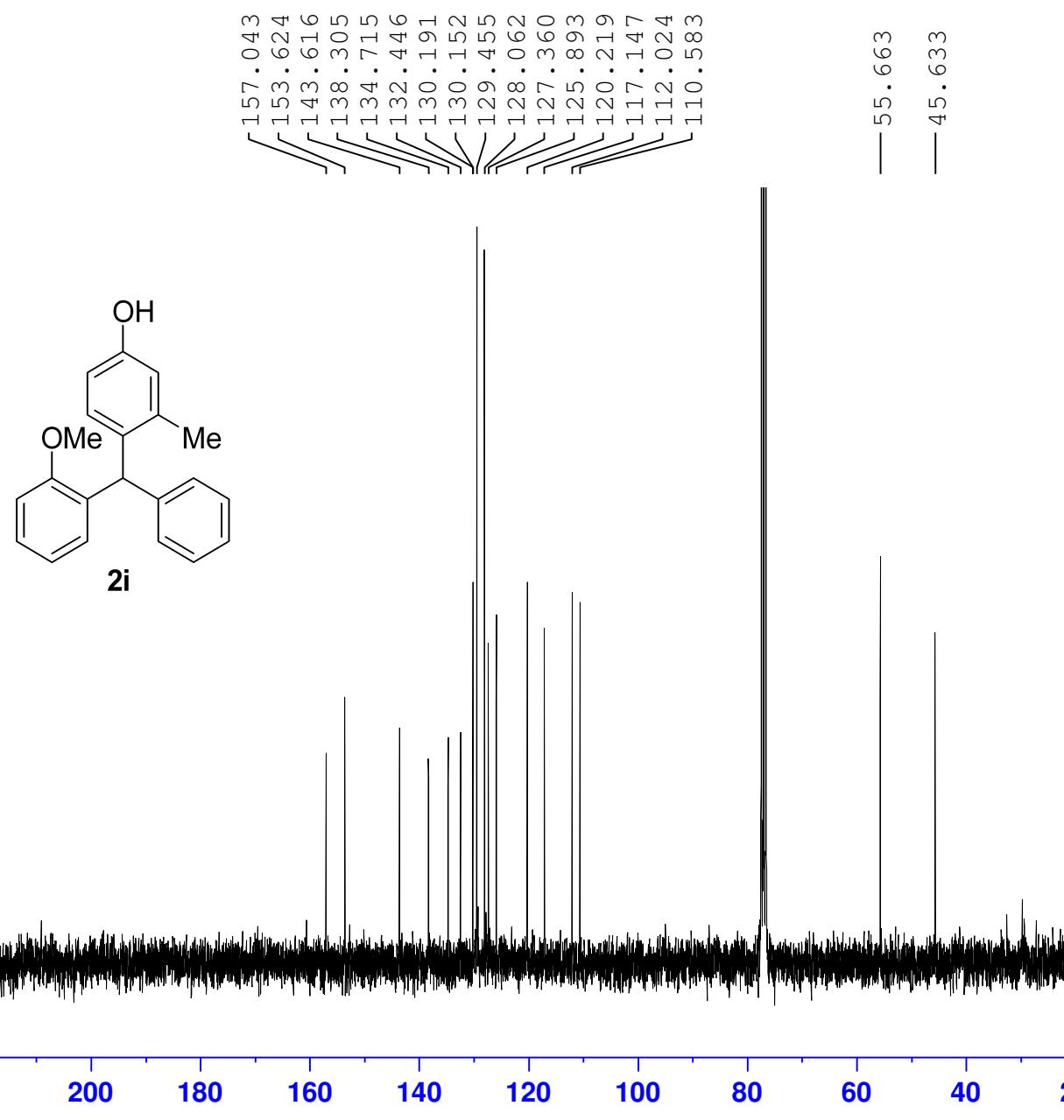
F2 - Processing parameters
SI 32768
SF 75.4677566 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

200 180 160 140 120 100 80 60 40 20 0 ppm

3sjwei 4415 zy-3-17e 1h cdcl3



3sjwei 4416 zy-3-17e 13c cdcl3



Current Data Parameters
NAME ZY-3-17E-c-fr
EXPNO 4416
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210703
Time 13.48
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 1024
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 296.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 ======

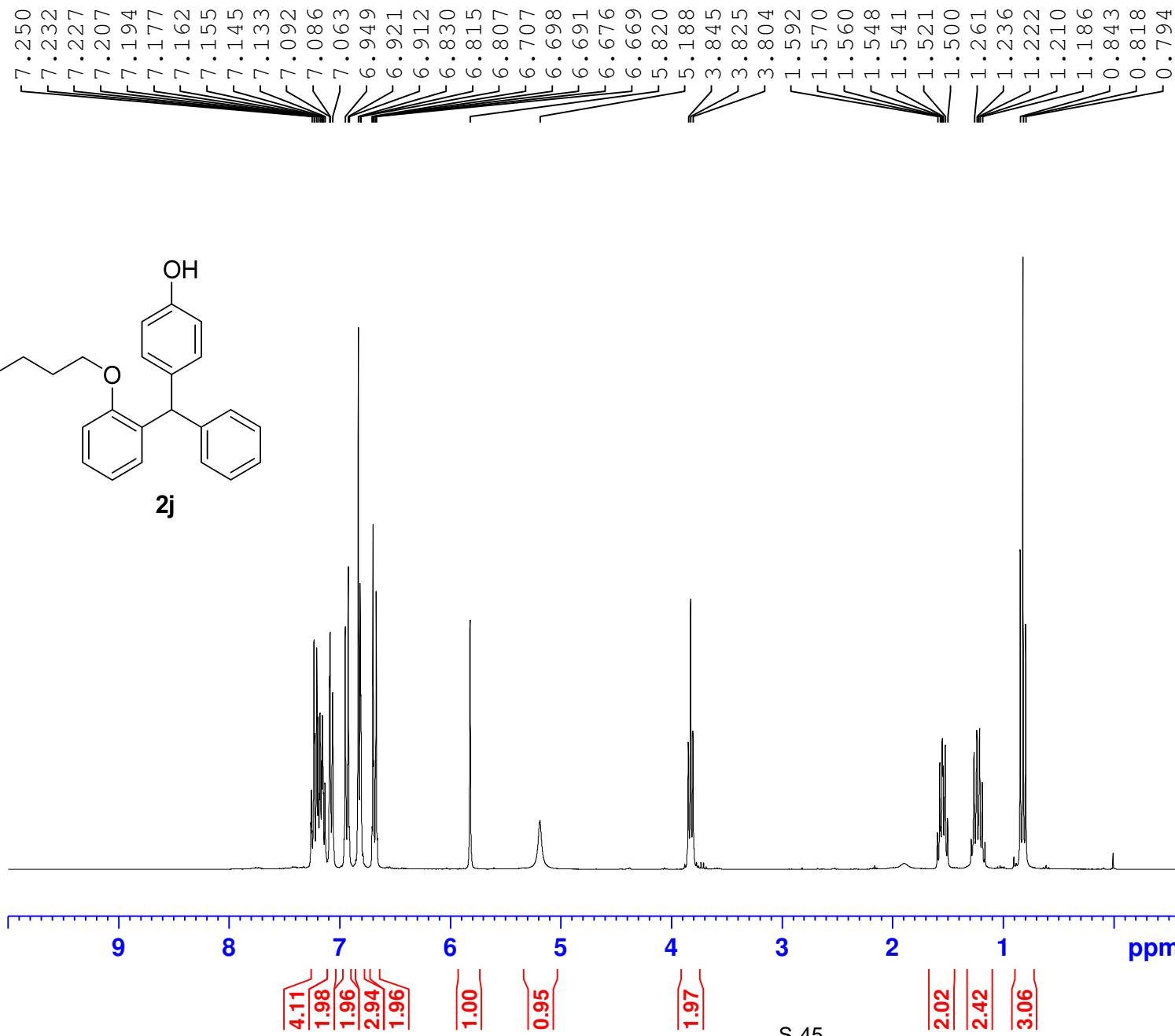
SFO1	75.4752949 MHz
NUC1	^{13}C
P1	9.50 usec
PLW1	34.20000076 W

===== CHANNEL f2 ======

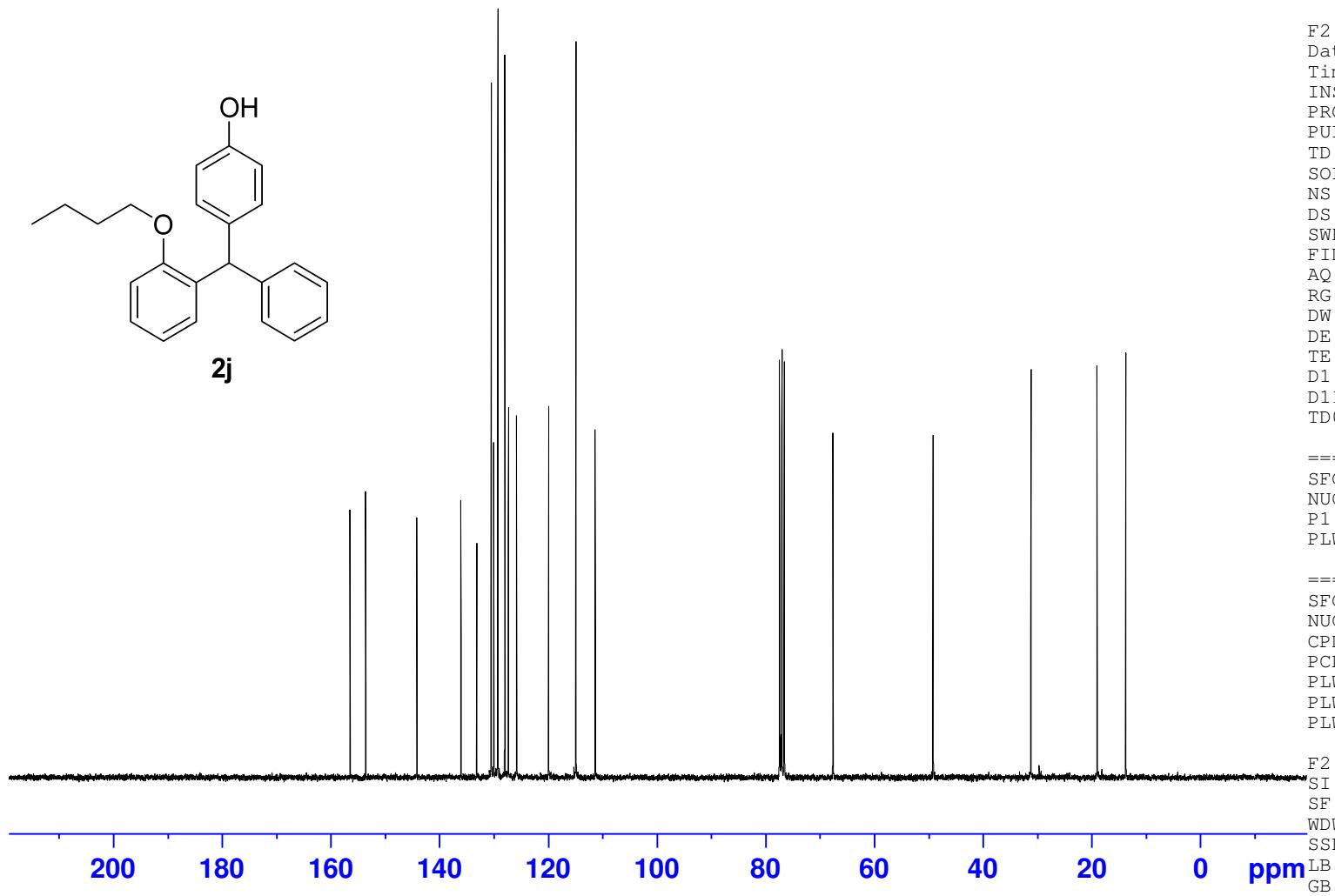
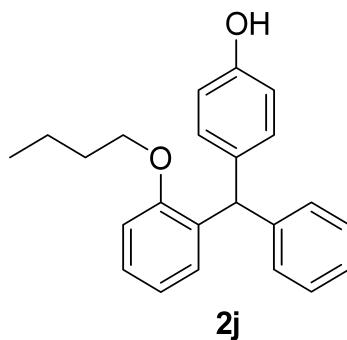
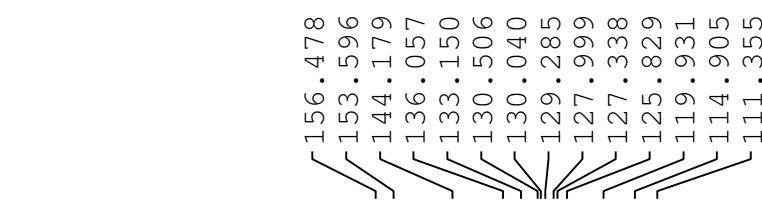
SFO2	300.1312005 MHz
NUC2	^1H
CPDPRG[2	waltz16
PCPD2	90.00 usec
PLW2	14.00000000 W
PLW12	0.17284000 W
PLW13	0.14000000 W

F2 - Processing parameters
SI 32768
SF 75.4677515 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

3sjwei 4386 zy-3-20b 1h cdcl3



3sjwei 4413 zy-3-20b 13c cdcl3



Current Data Parameters
NAME ZY-3-20B-c-fr
EXPNO 4413
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210703
Time 11.25
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 1024
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 296.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 ======

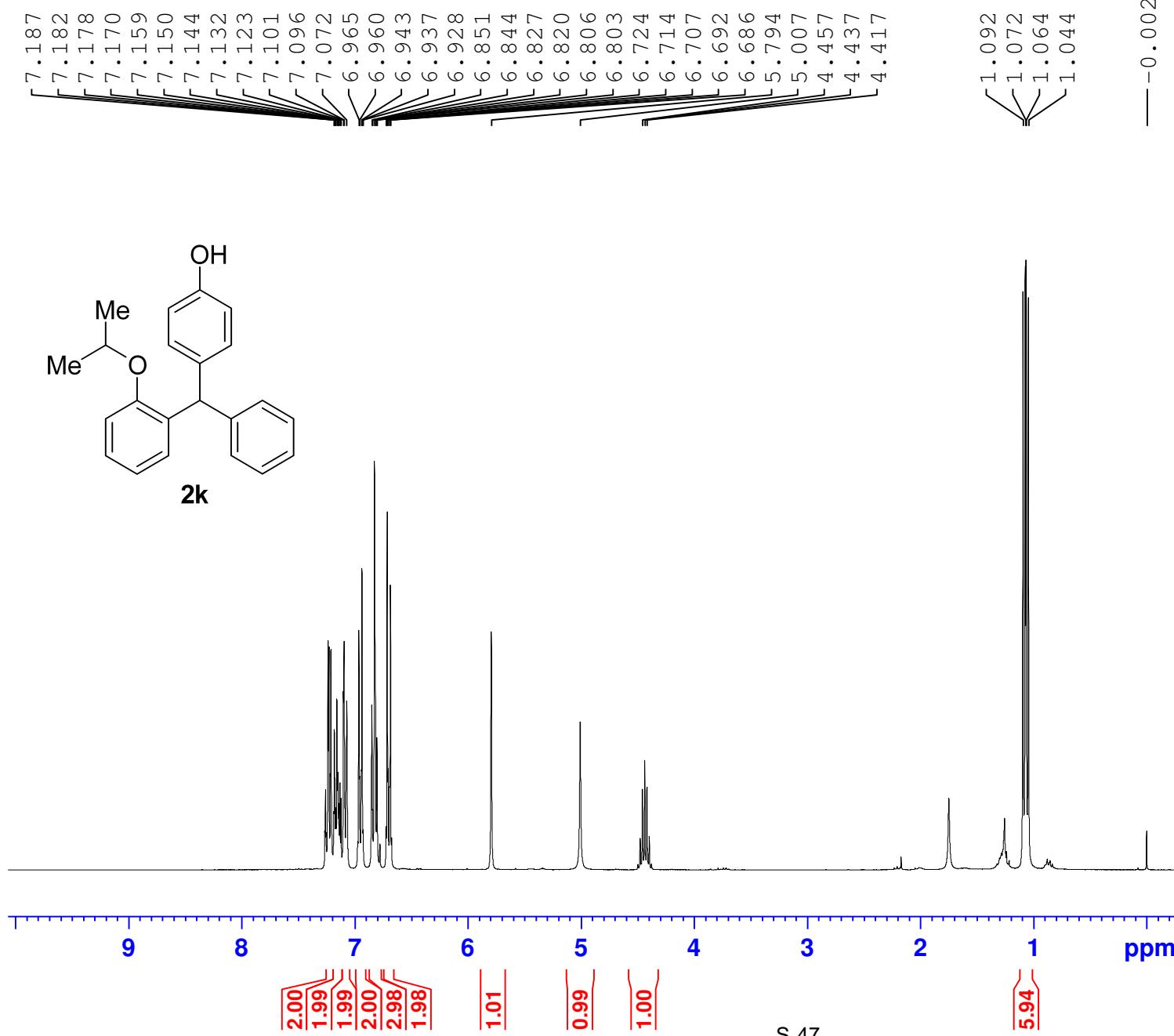
SFO1 75.4752949 MHz
NUC1 ^{13}C
P1 9.50 usec
PLW1 34.20000076 W

===== CHANNEL f2 ======

SFO2 300.1312005 MHz
NUC2 ^1H
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 14.00000000 W
PLW12 0.17284000 W
PLW13 0.14000000 W

F2 - Processing parameters
SI 32768
SF 75.4677585 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

3sjwei 4384 zy-3-17d 1h cdcl3



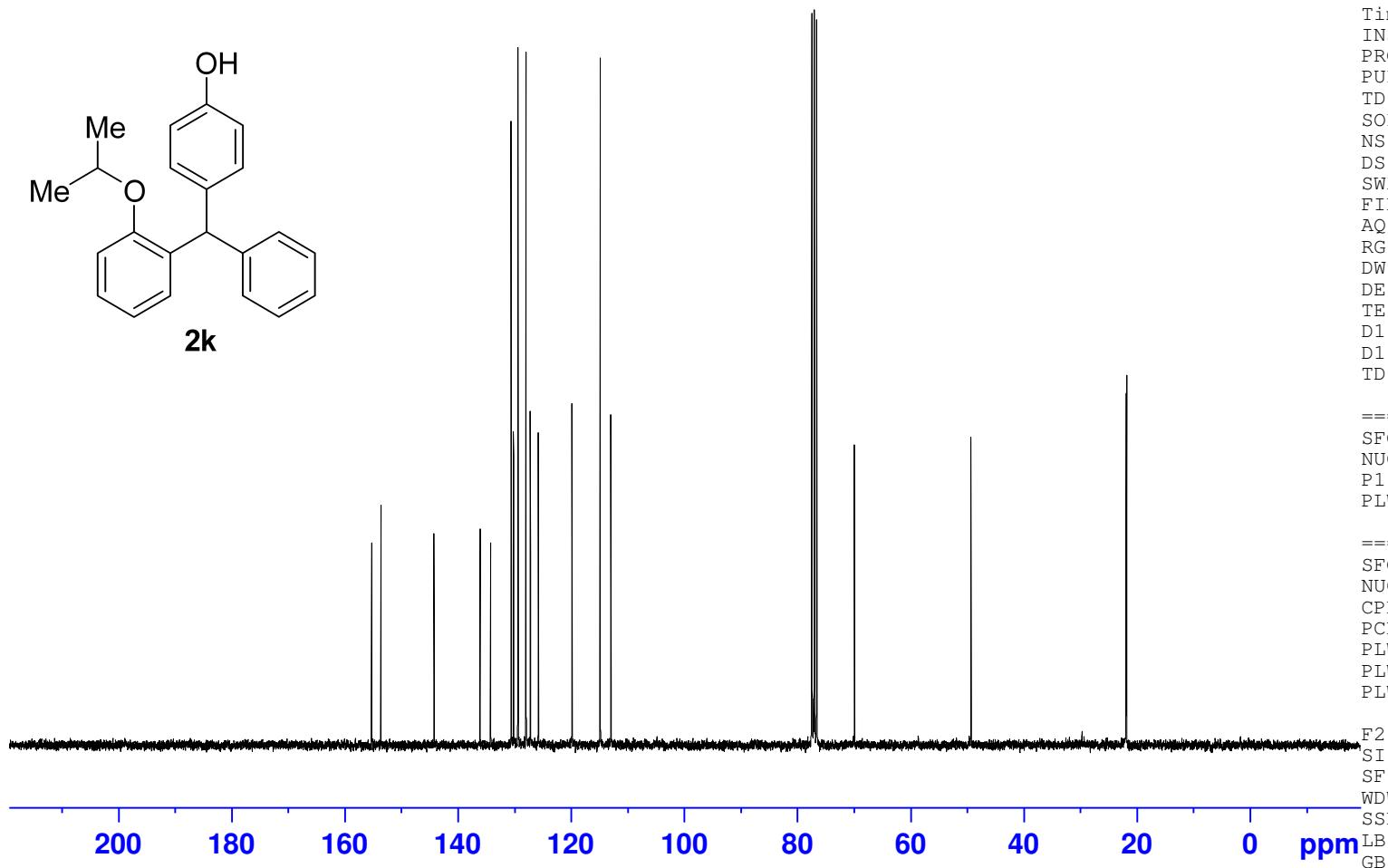
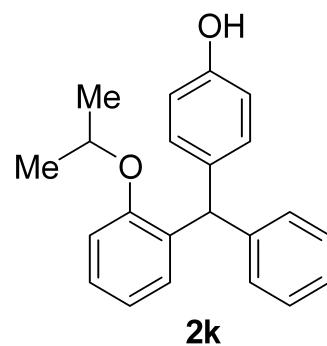
Current Data Parameters
NAME ZY-3-17D-h-fr
EXPNO 4384
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210702
Time 9.26
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 6009.615 Hz
FIDRES 0.091699 Hz
AQ 5.4525952 sec
RG 101
DW 83.200 usec
DE 6.50 usec
TE 296.1 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
SFO1 300.1318534 MHz
NUC1 1H
P1 10.00 usec
PLW1 14.00000000 W

F2 - Processing parameters
SI 65536
SF 300.1300165 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

3sjwei 4414 zy-3-17d 13c cdcl3



Current Data Parameters
NAME ZY-3-17D-c-fr
EXPNO 4414
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210703
Time 12.35
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 1024
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 50.8
DW 27.733 usec
DE 6.50 usec
TE 296.2 K
D1 2.0000000 sec
D11 0.0300000 sec
TD0 1

===== CHANNEL f1 ======

SFO1	75.4752949 MHz
NUC1	¹³ C
P1	9.50 usec
PLW1	34.20000076 W

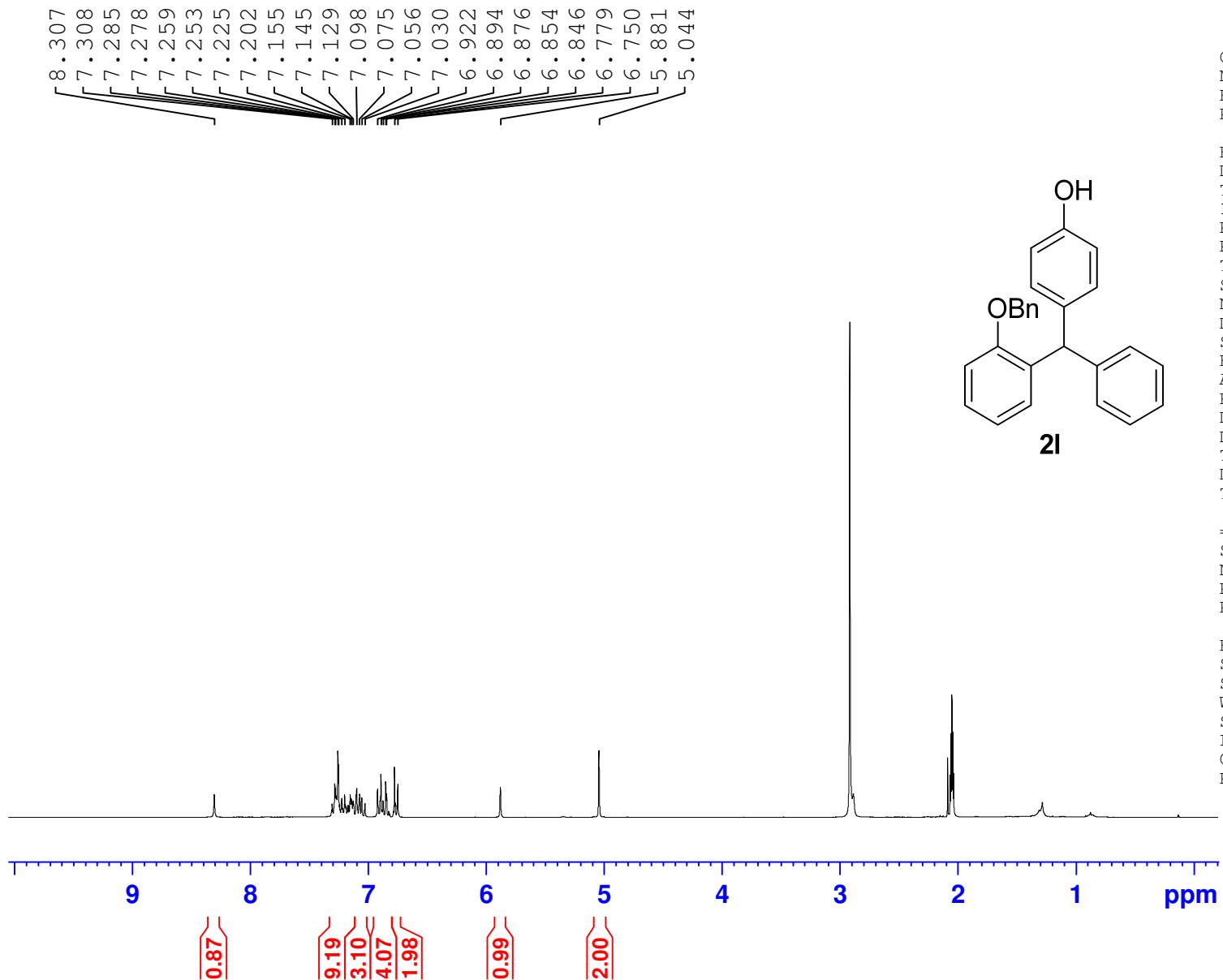
===== CHANNEL f2 ======

SFO2	300.1312005 MHz
NUC2	¹ H
CPDPRG[2]	waltz16
PCPD2	90.00 usec
PLW2	14.00000000 W
PLW12	0.17284000 W
PLW13	0.14000000 W

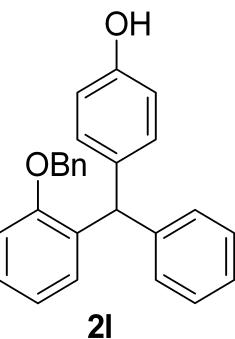
F2 - Processing parameters

SI	32768
SF	75.4677543 MHz
WDW	EM
SSB	0
LB	1.00 Hz
GB	0
PC	1.40

3sjwei 2712 zy-3-11b 1h acetone



3sjwei 2327 zy-3-11b 13c acetone

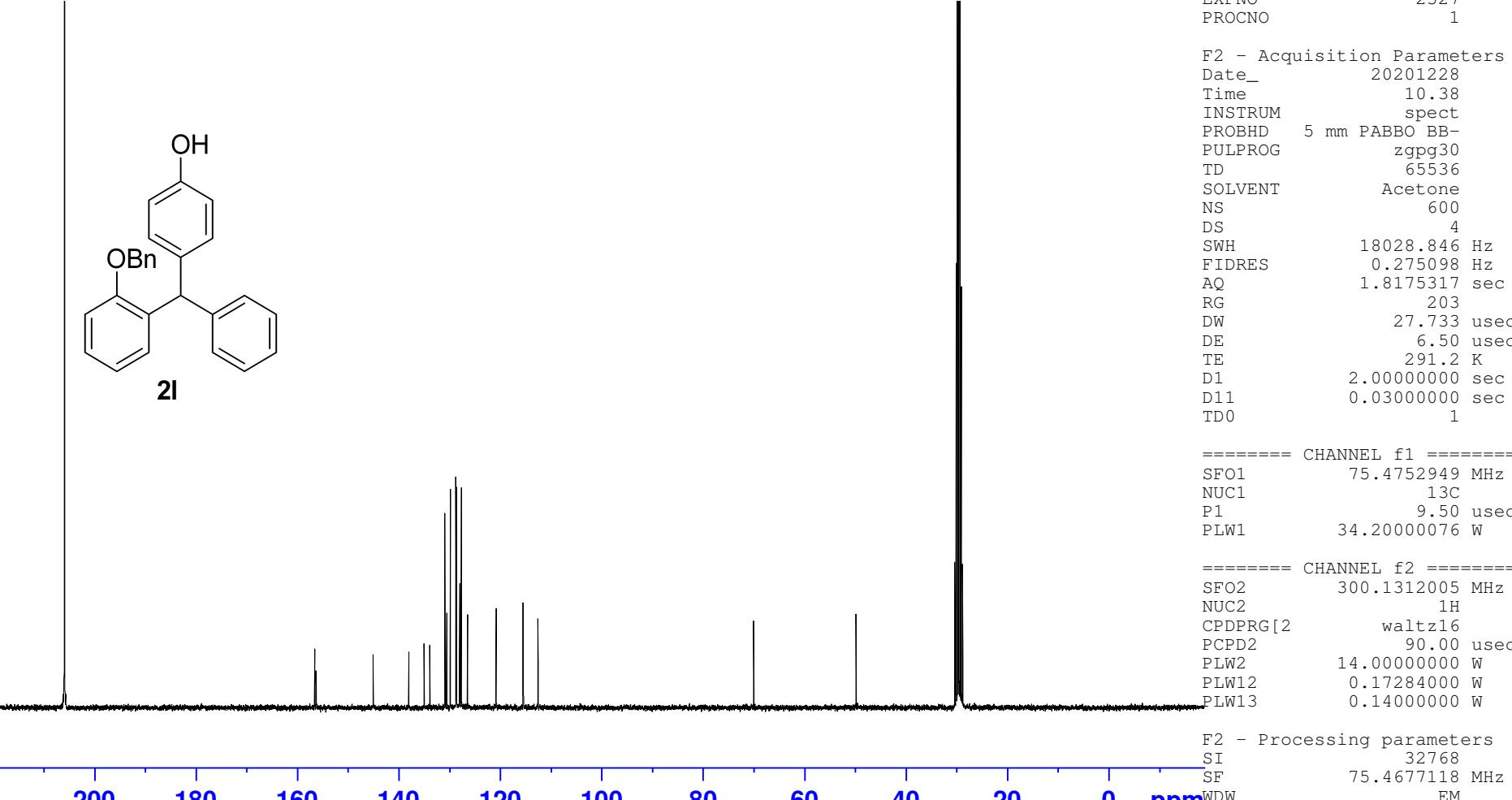


2l

156.535
156.272
145.027
137.963
134.968
133.839
130.862
130.480
129.786
128.685
128.547
127.948
127.923
127.627
126.386
120.736
115.449
112.482

— 69.941

— 49.785



Current Data Parameters
NAME ZY-3-11B-c-fr
EXPNO 2327
PROCNO 1

F2 - Acquisition Parameters
Date_ 20201228
Time 10.38
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT Acetone
NS 600
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 291.2 K
D1 2.0000000 sec
D11 0.0300000 sec
TD0 1

===== CHANNEL f1 ======

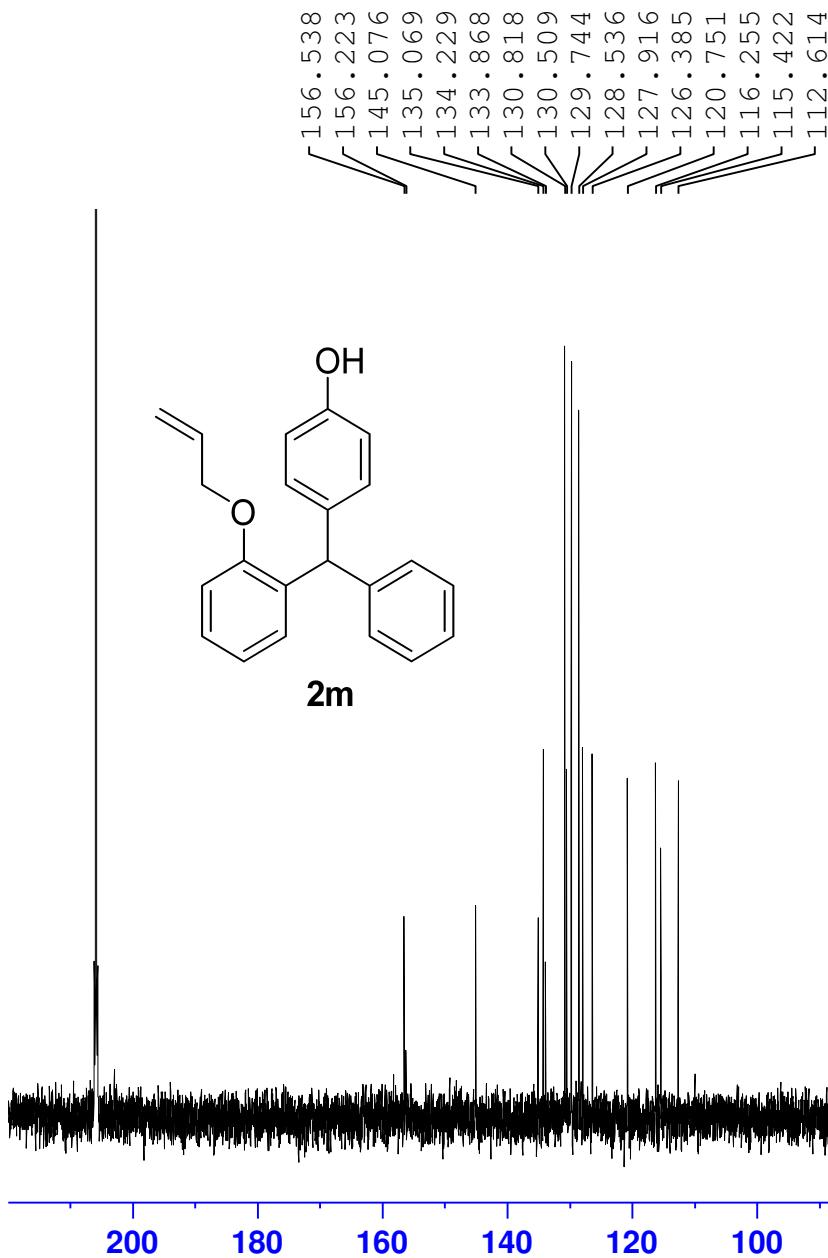
SFO1 75.4752949 MHz
NUC1 ¹³C
P1 9.50 usec
PLW1 34.20000076 W

===== CHANNEL f2 ======

SFO2 300.1312005 MHz
NUC2 ^{1H}
CPDPRG[2] waltz16
PCPD2 90.00 usec
PLW2 14.00000000 W
PLW12 0.17284000 W
PLW13 0.14000000 W

F2 - Processing parameters
SI 32768
SF 75.4677118 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

3sjwei 2439 zy-3-14c 13c acetone



— 68.920
— 49.564

Current Data Parameters
NAME ZY-3-17B-c-fr
EXPNO 2439
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210107
Time 14.12
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT Acetone
NS 600
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 289.2 K
D1 2.0000000 sec
D11 0.0300000 sec
TD0 1

===== CHANNEL f1 ======

SFO1	75.4752949 MHz
NUC1	¹³ C
P1	9.50 usec
PLW1	34.20000076 W

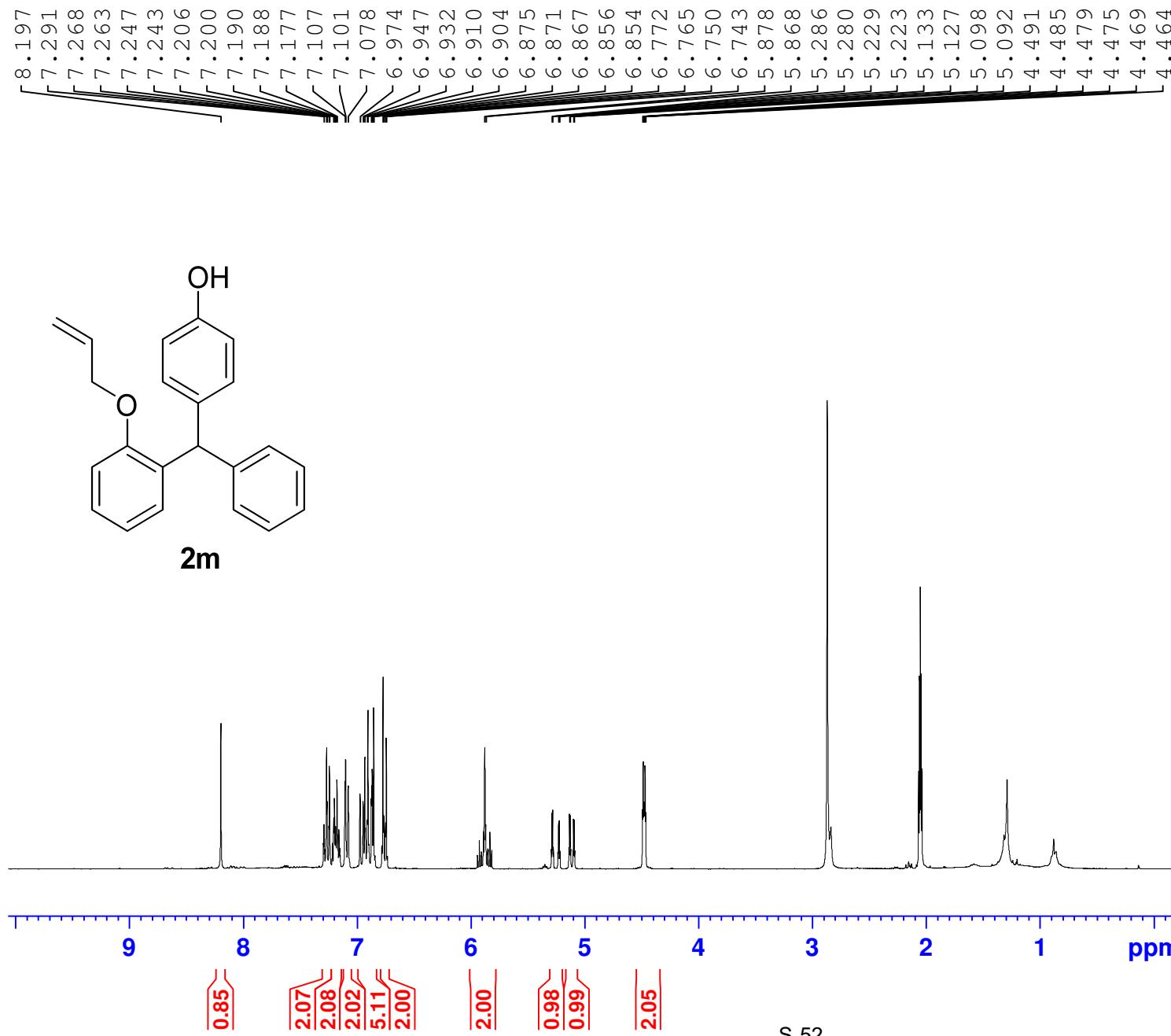
===== CHANNEL f2 ======

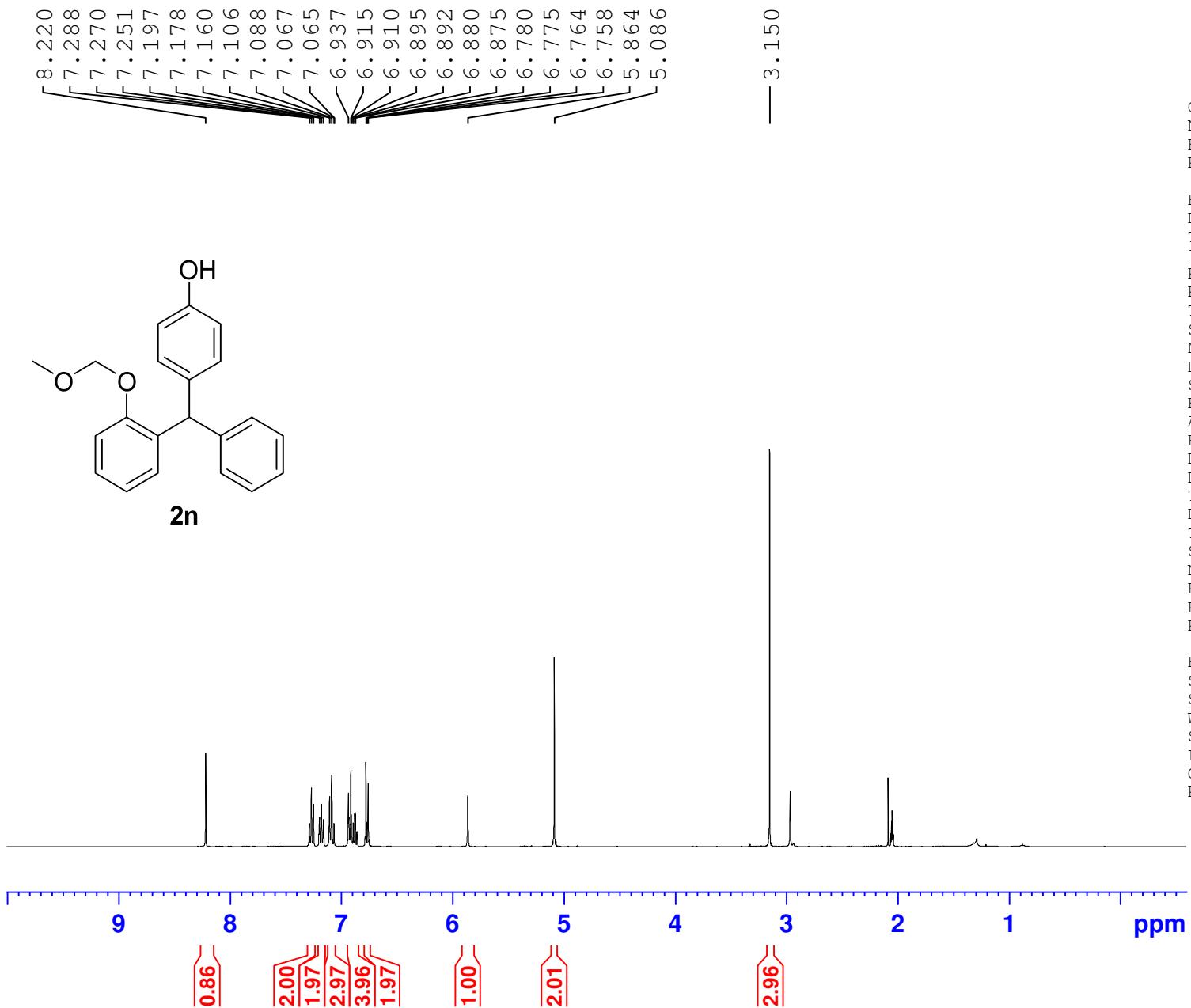
SFO2	300.1312005 MHz
NUC2	¹ H
CPDPRG[2	waltz16
PCPD2	90.00 usec
PLW2	14.00000000 W
PLW12	0.17284000 W
PLW13	0.14000000 W

F2 - Processing parameters

SI	32768
SF	75.4677083 MHz
WDW	EM
SSB	0
LB	1.00 Hz
GB	0
PC	1.40

3sjwei 4338 zy-3-14c 1h acetone

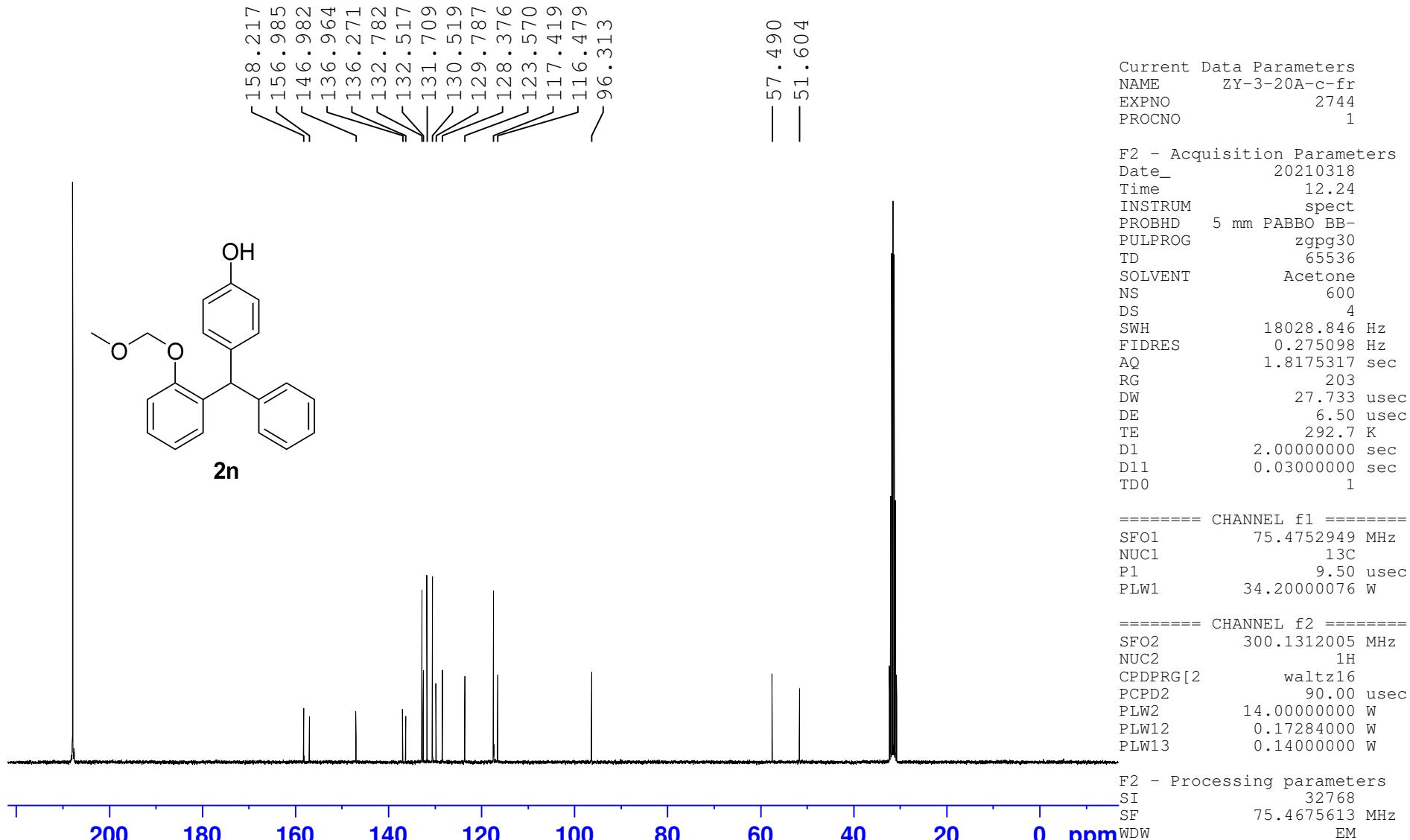




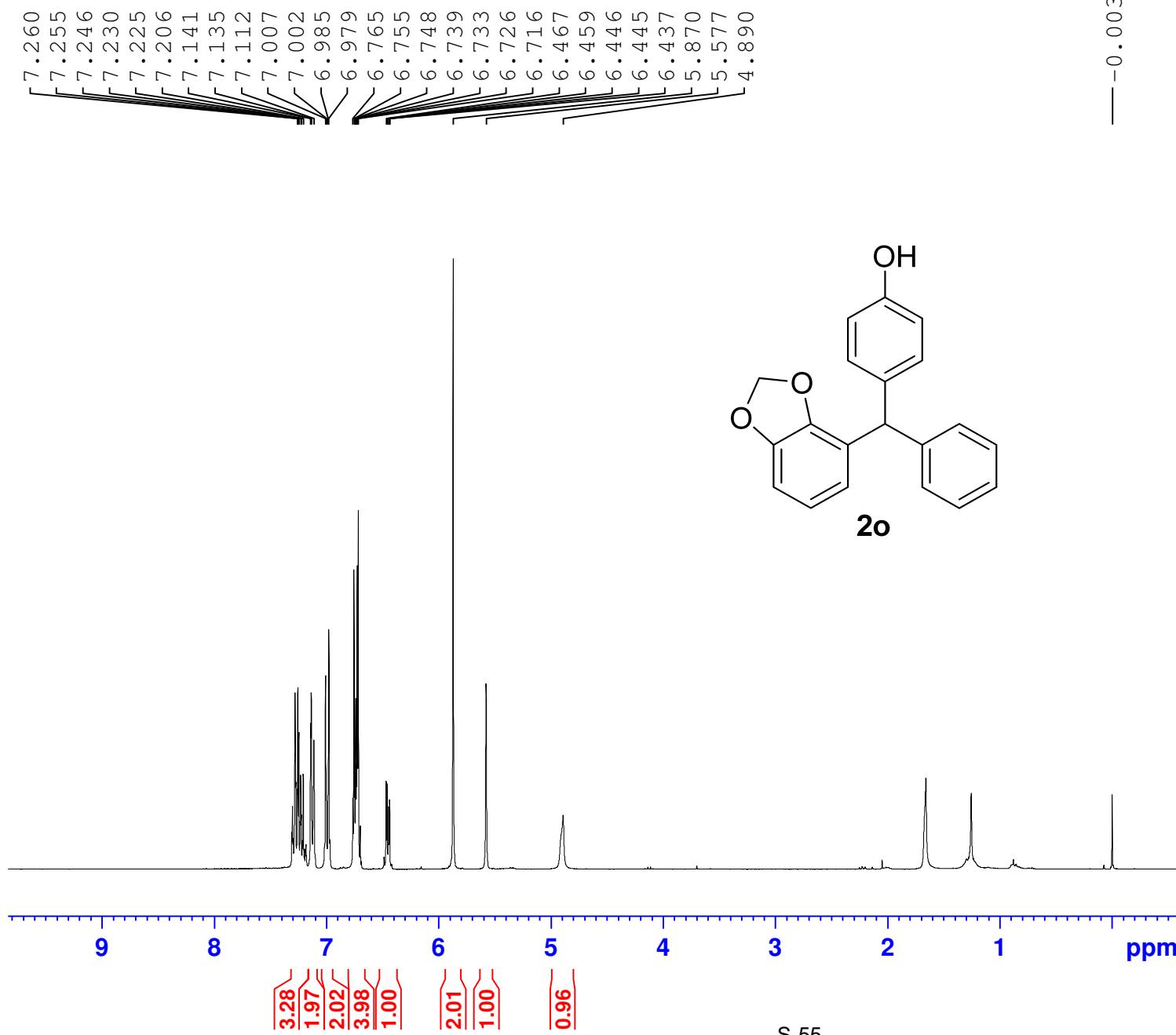
Current Data Parameters
 NAME 2n
 EXPNO 8
 PROCNO 1

F2 - Acquisition Parameters
 Date_ 20210328
 Time 11.29 h
 INSTRUM Avance
 PROBHD Z116098_0833 (zg30
 PULPROG 65536
 TD Acetone
 NS 16
 DS 2
 SWH 8196.722 Hz
 FIDRES 0.250144 Hz
 AQ 3.9976959 sec
 RG 79.7194
 DW 61.000 usec
 DE 13.54 usec
 TE 295.0 K
 D1 1.0000000 sec
 TD0 1
 SFO1 400.1324708 MHz
 NUC1 1H
 P0 3.33 usec
 P1 10.00 usec
 PLW1 20.73200035 W

F2 - Processing parameters
 SI 65536
 SF 400.1300069 MHz
 WDW EM
 SSB 0
 LB 0.30 Hz
 GB 0
 PC 1.00



3sjwei 4820 zy-4-44e 1h cdcl3



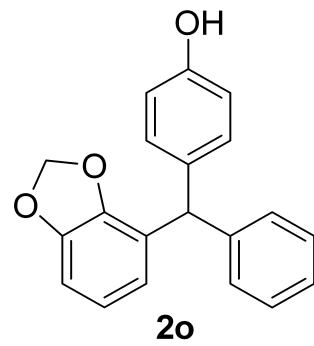
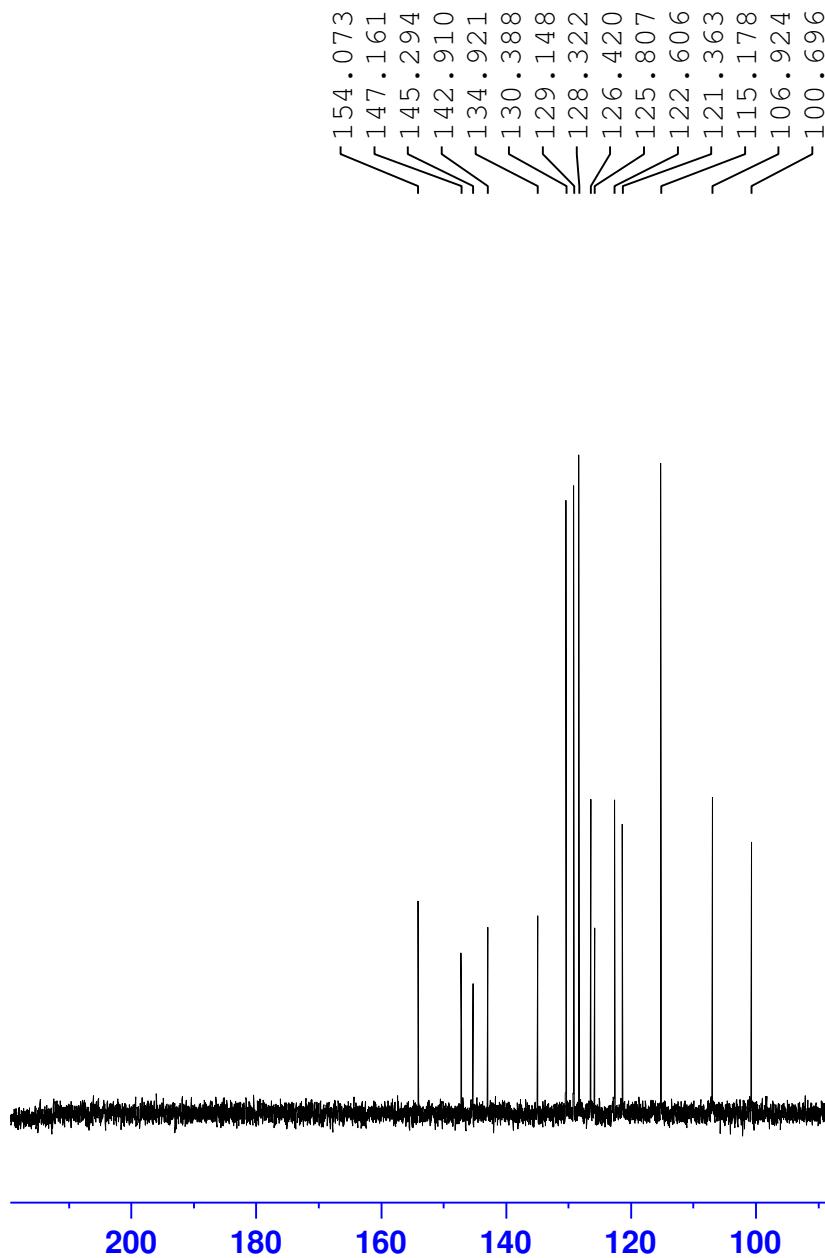
Current Data Parameters
NAME ZY-3-36-h-fr
EXPNO 4820
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210723
Time 11.29
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl3
NS 16
DS 2
SWH 6009.615 Hz
FIDRES 0.091699 Hz
AQ 5.4525952 sec
RG 181
DW 83.200 usec
DE 6.50 usec
TE 296.1 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 =====
SFO1 300.1318534 MHz
NUC1 1H
P1 10.00 usec
PLW1 14.00000000 W

F2 - Processing parameters
SI 65536
SF 300.1300112 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

3sjwei 4821 zy-4-44e 13c cdcl3



Current Data Parameters
NAME ZY-3-36-c-fr
EXPNO 4821
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210723
Time 12.16
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT CDCl3
NS 700
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 296.2 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 ======

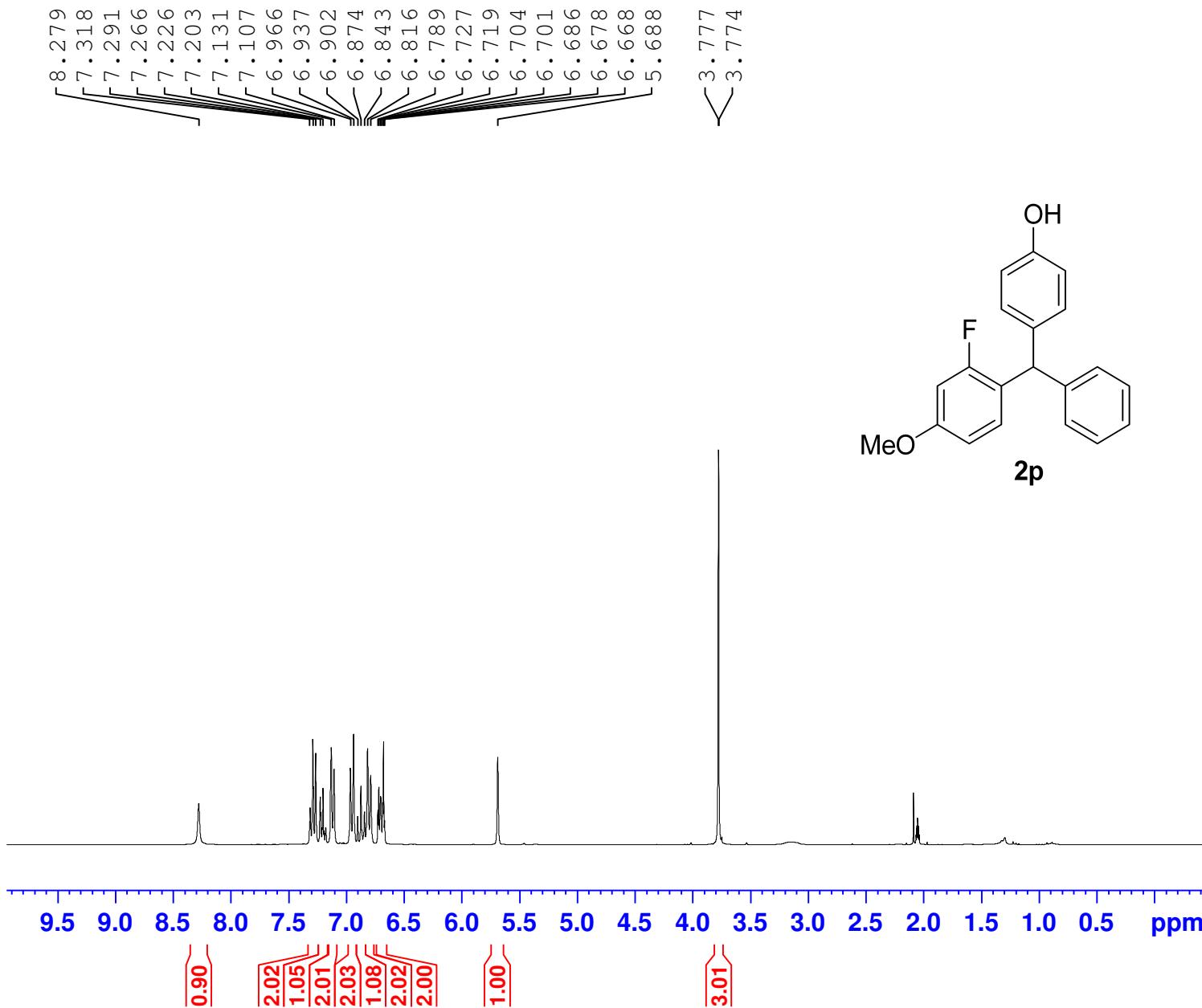
SFO1	75.4752949 MHz
NUC1	13C
P1	9.50 usec
PLW1	34.20000076 W

===== CHANNEL f2 ======

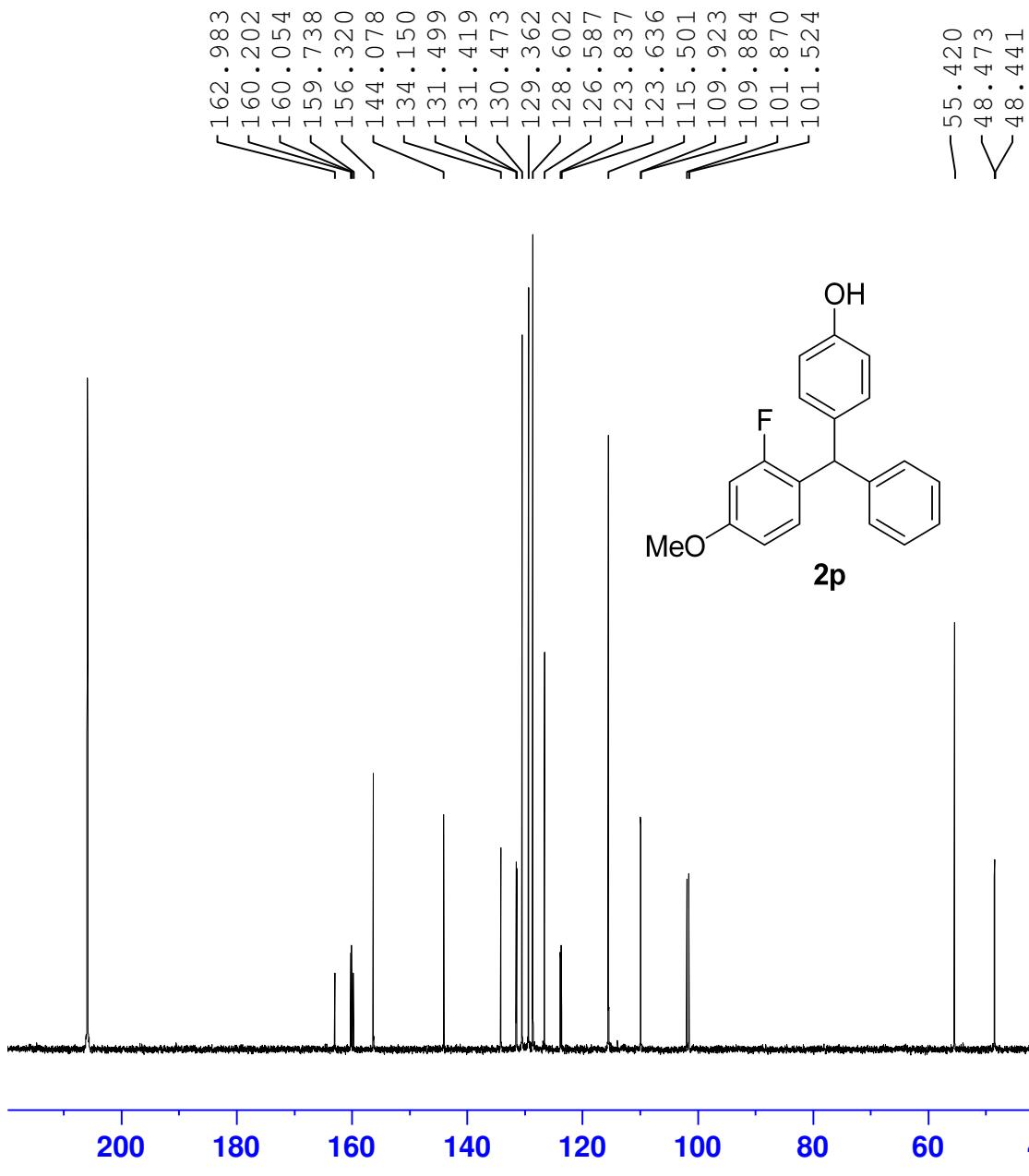
SFO2	300.1312005 MHz
NUC2	1H
CPDPRG[2	waltz16
PCPD2	90.00 usec
PLW2	14.00000000 W
PLW12	0.17284000 W
PLW13	0.14000000 W

F2 - Processing parameters
SI 32768
SF 75.4677485 MHz
WDW EM
SSB 0
LB 1.00 Hz
GB 0
PC 1.40

3sjwei 2935 zy-3-27g 1h acetone



3sjwei 2936 zy-3-27g 13c acetone



Current Data Parameters
NAME ZY-3-27G-c-fr
EXPNO 2936
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210408
Time 12.02
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgpg30
TD 65536
SOLVENT Acetone
NS 600
DS 4
SWH 18028.846 Hz
FIDRES 0.275098 Hz
AQ 1.8175317 sec
RG 203
DW 27.733 usec
DE 6.50 usec
TE 296.1 K
D1 2.00000000 sec
D11 0.03000000 sec
TD0 1

===== CHANNEL f1 ======

SFO1	75.4752949 MHz
NUC1	¹³ C
P1	9.50 usec
PLW1	34.20000076 W

===== CHANNEL f2 ======

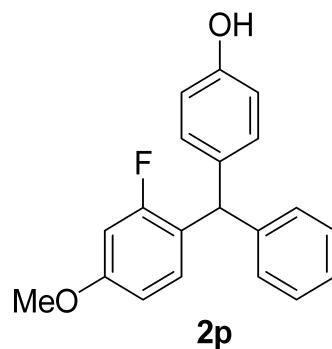
SFO2	300.1312005 MHz
NUC2	¹ H
CPDPRG[2]	waltz16
PCPD2	90.00 usec
PLW2	14.00000000 W
PLW12	0.17284000 W
PLW13	0.14000000 W

F2 - Processing parameters

SI	32768
SF	75.4677215 MHz
WDW	EM
SSB	0
LB	1.00 Hz
GB	0
PC	1.40

3sjwei 2908 zy-3-27g 19f cdcl3

-114.289



Current Data Parameters
NAME 0407sjw
EXPNO 2908
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210407
Time 10.22
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zgfhigqn.2
TD 131072
SOLVENT CDCl3
NS 16
DS 4
SWH 66964.289 Hz
FIDRES 0.510897 Hz
AQ 0.9786710 sec
RG 203
DW 7.467 usec
DE 6.50 usec
TE 296.2 K
D1 1.00000000 sec
D11 0.03000000 sec
D12 0.00002000 sec
TD0 1

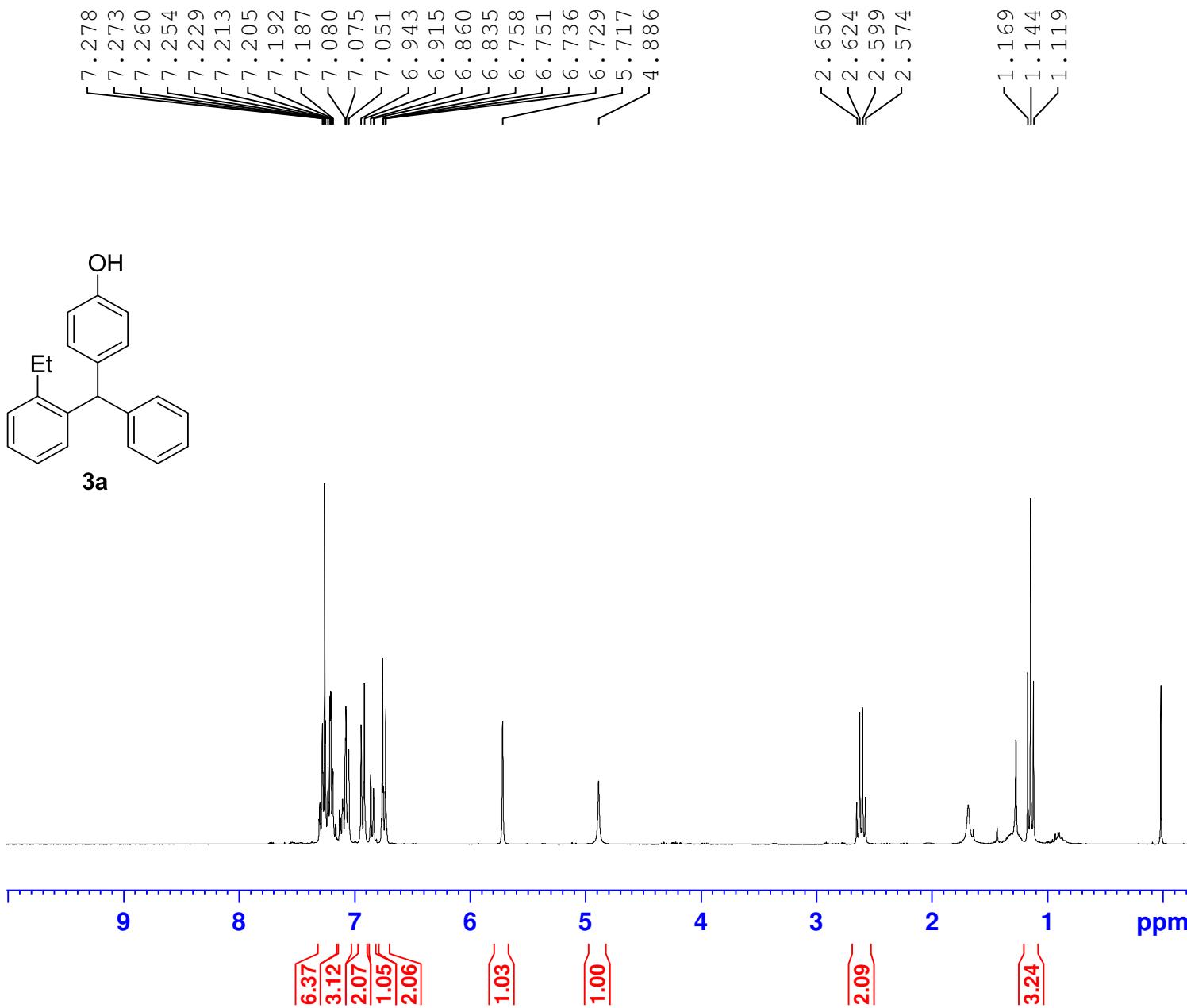
===== CHANNEL f1 ======
SFO1 282.3761148 MHz
NUC1 19F
P1 14.50 usec
PLW1 10.39999962 W

===== CHANNEL f2 ======
SFO2 300.1312005 MHz
NUC2 1H
CPDPGRG[2] waltz16
PCPD2 90.00 usec
PLW2 14.00000000 W
PLW12 0.17284000 W

F2 - Processing parameters
SI 65536
SF 282.4043552 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



3sjwei 4313 h-100-1 1h cdcl₃



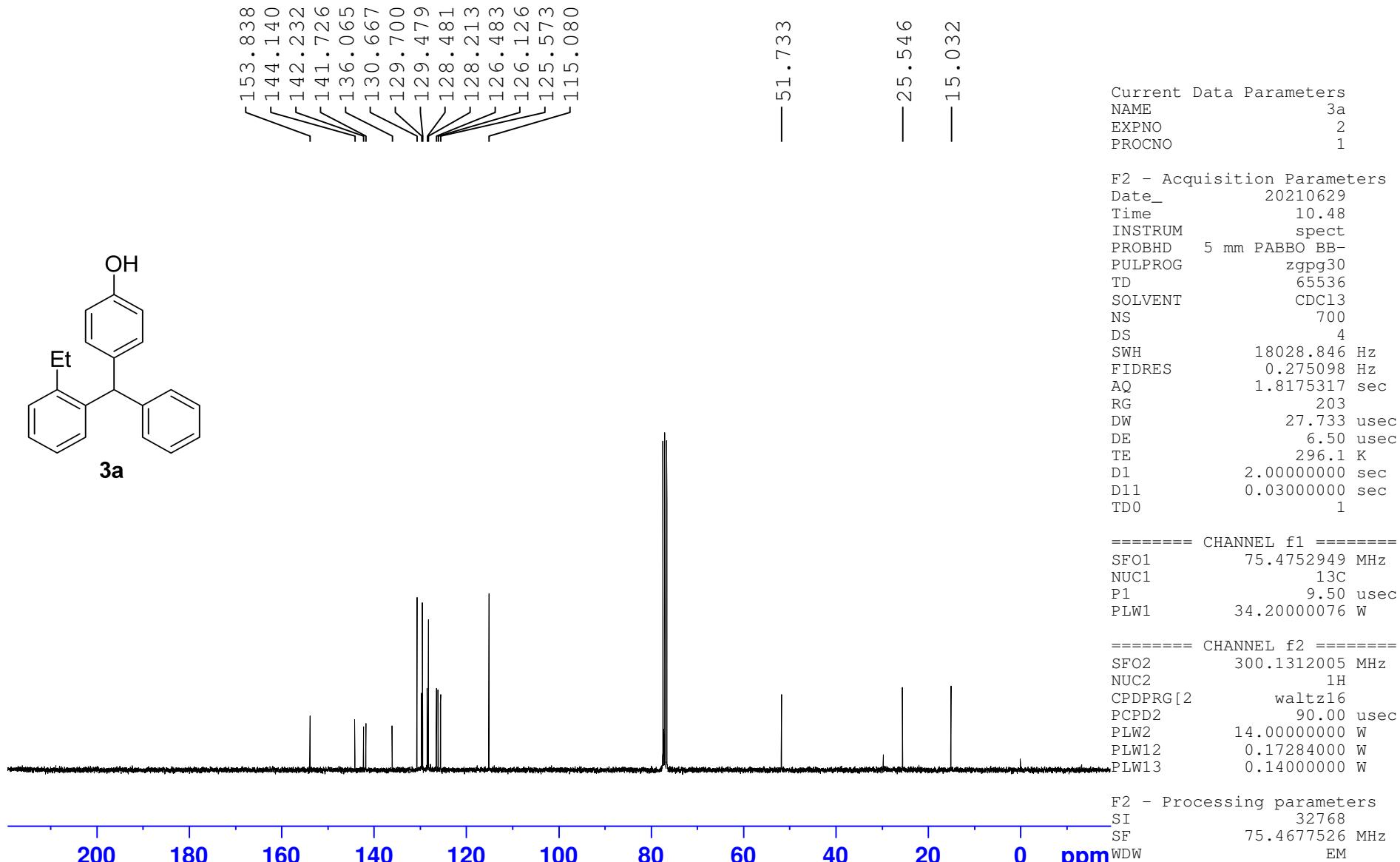
Current Data Parameters
NAME 3a
EXPNO 1
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210628
Time 9.11
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl₃
NS 16
DS 2
SWH 6009.615 Hz
FIDRES 0.091699 Hz
AQ 5.4525952 sec
RG 144
DW 83.200 usec
DE 6.50 usec
TE 296.2 K
D1 1.00000000 sec
TD0 1

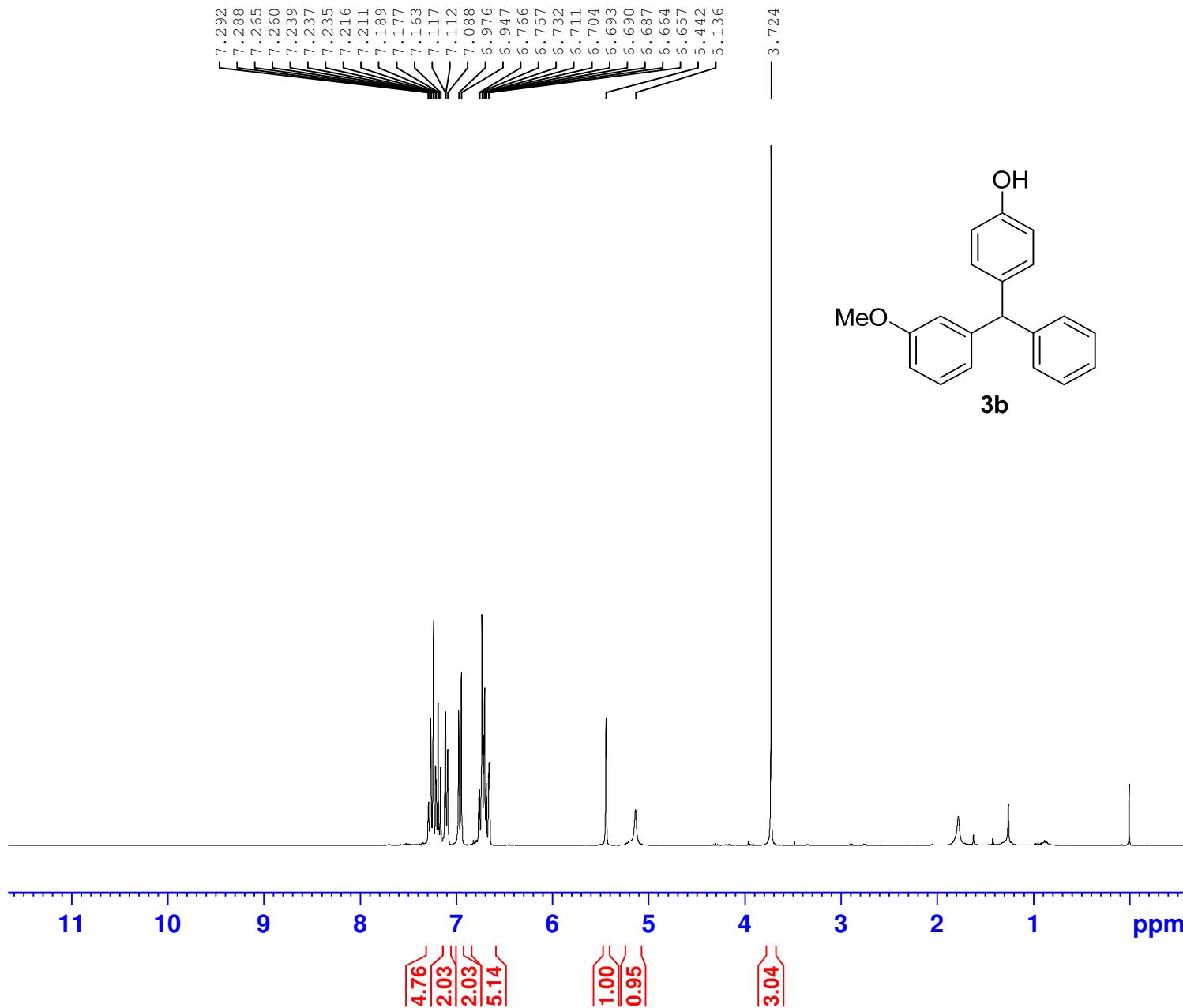
===== CHANNEL f1 ======

SFO1 300.1318534 MHz
NUC1 1H
P1 10.00 usec
PLW1 14.00000000 W

F2 - Processing parameters
SI 65536
SF 300.1300071 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



3sjwei 4314 h-100-2 1h cdcl₃

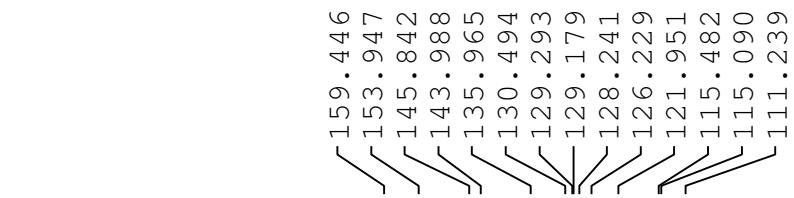


Current Data Parameters
NAME H-3b
EXPNO 1
PROCNO 1

F2 - Acquisition Parameters
Date_ 20210628
Time 9.17
INSTRUM spect
PROBHD 5 mm PABBO BB-
PULPROG zg30
TD 65536
SOLVENT CDCl₃
NS 16
DS 2
SWH 6009.615 Hz
FIDRES 0.091699 Hz
AQ 5.4525952 sec
RG 114
DW 83.200 usec
DE 6.50 usec
TE 296.1 K
D1 1.00000000 sec
TD0 1

===== CHANNEL f1 ======
SFO1 300.1318534 MHz
NUC1 1H
P1 10.00 usec
PLW1 14.00000000 W

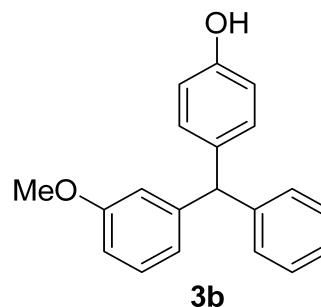
F2 - Processing parameters
SI 65536
SF 300.1300147 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00



55.923
55.110

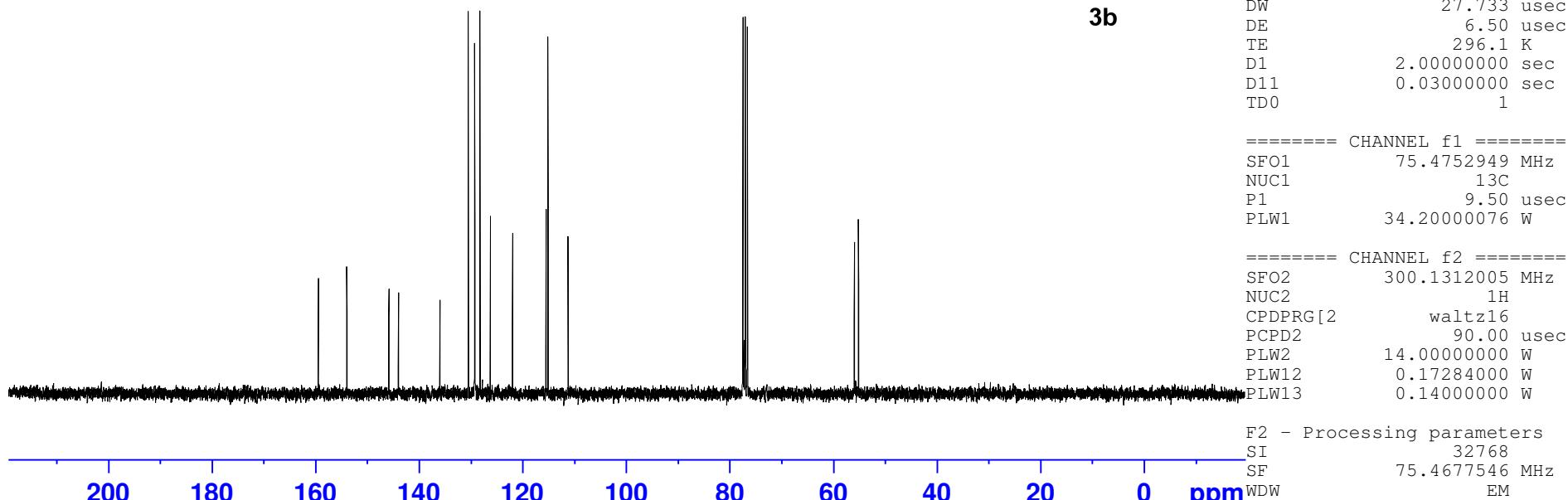
Current Data Parameters
 NAME 3b
 EXPNO 1
 PROCNO 1

F2 - Acquisition Parameters
 Date_ 20210629
 Time 11.01
 INSTRUM spect
 PROBHD 5 mm PABBO BB-
 PULPROG zgpg30
 TD 65536
 SOLVENT CDCl3
 NS 156
 DS 4
 SWH 18028.846 Hz
 FIDRES 0.275098 Hz
 AQ 1.8175317 sec
 RG 203
 DW 27.733 usec
 DE 6.50 usec
 TE 296.1 K
 D1 2.00000000 sec
 D11 0.03000000 sec
 TD0 1

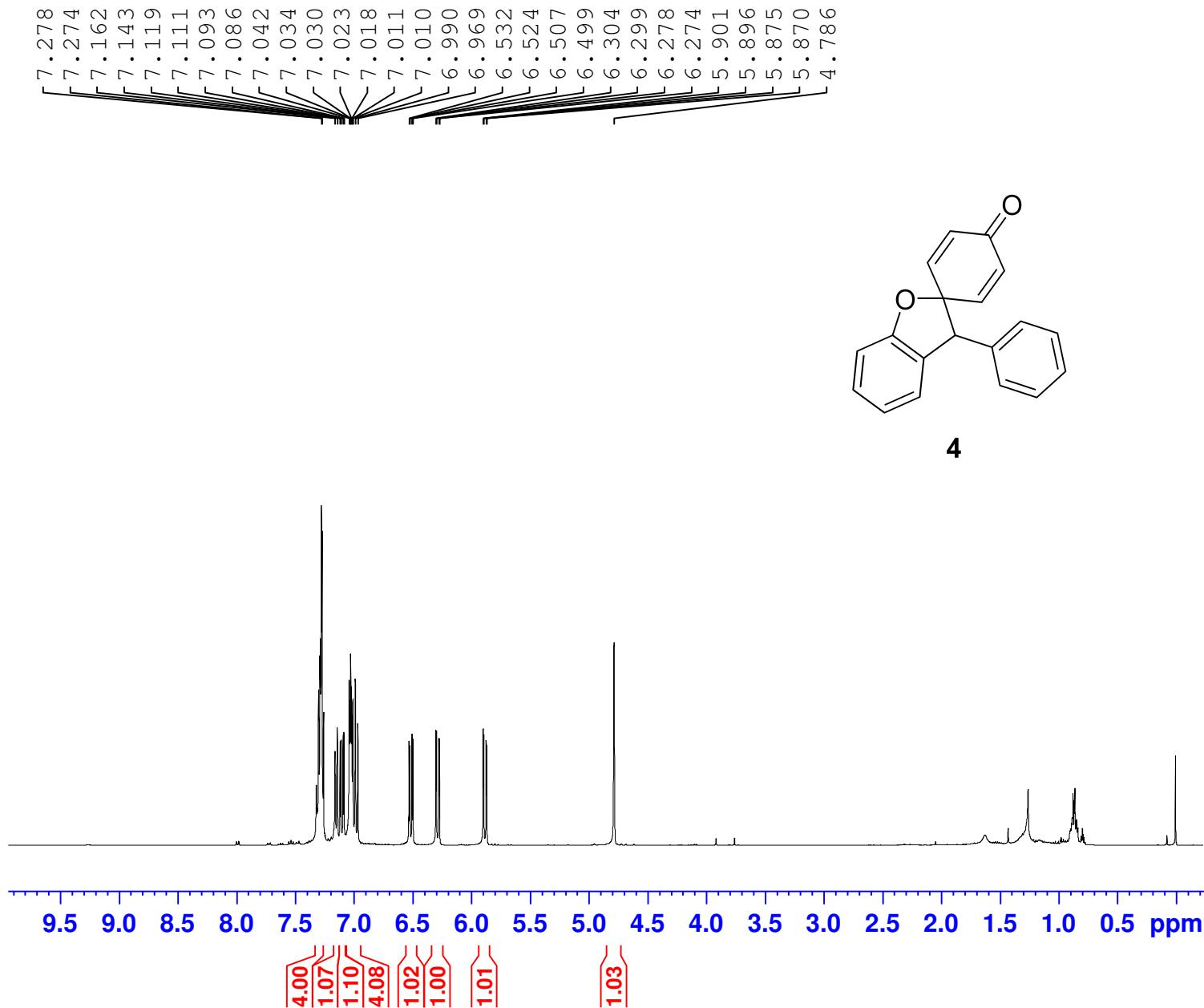


===== CHANNEL f1 ======
 SFO1 75.4752949 MHz
 NUC1 13C
 P1 9.50 usec
 PLW1 34.20000076 W

===== CHANNEL f2 ======
 SFO2 300.1312005 MHz
 NUC2 1H
 CPDPRG[2] waltz16
 PCPD2 90.00 usec
 PLW2 14.00000000 W
 PLW12 0.17284000 W
 PLW13 0.14000000 W



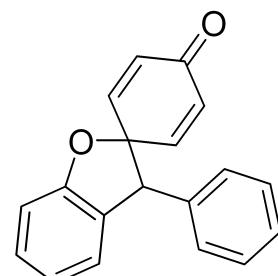
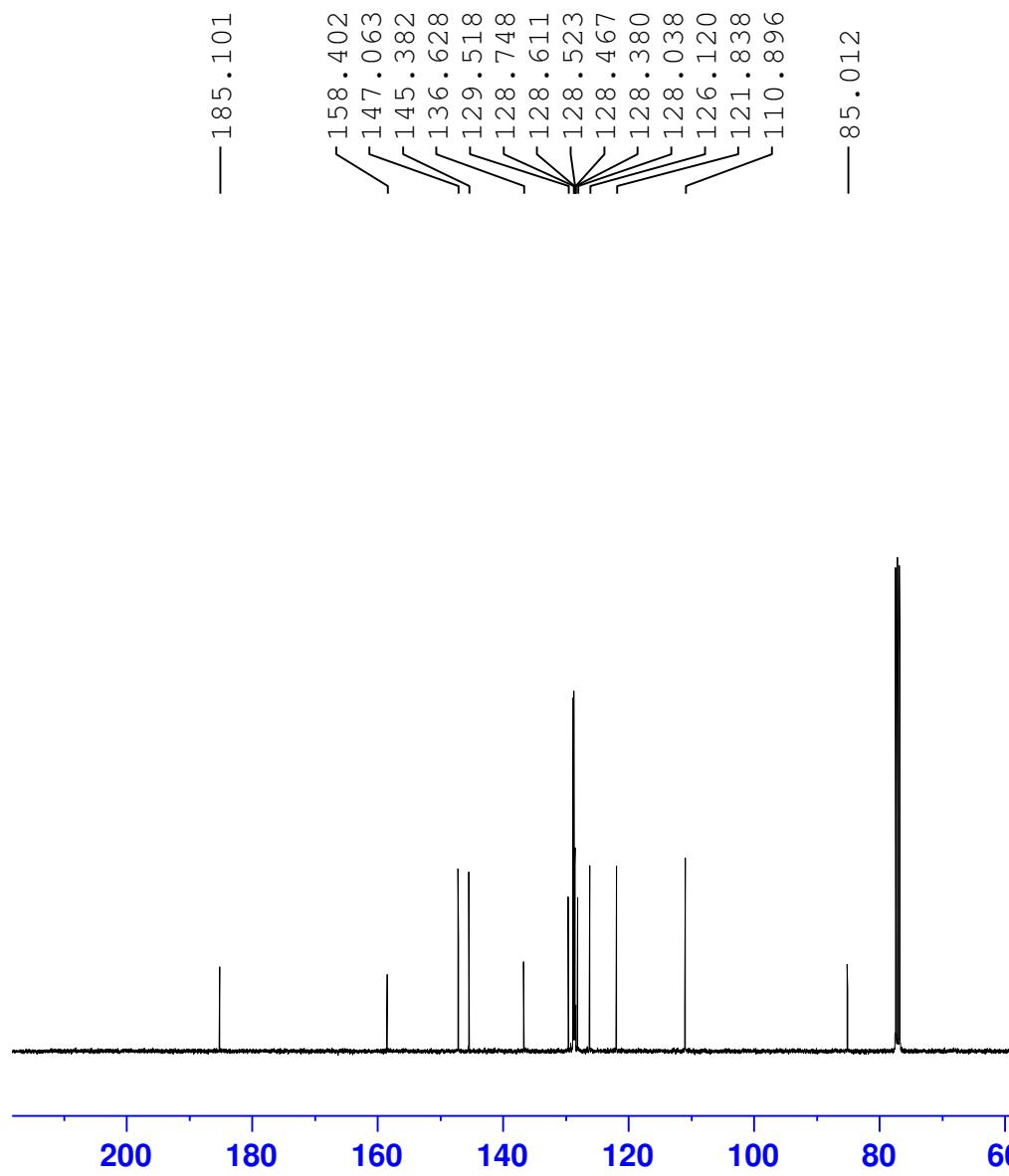
F2 - Processing parameters
 SI 32768
 SF 75.4677546 MHz
 WDW EM
 SSB 0
 LB 1.00 Hz
 GB 0
 PC 1.40



Current Data Parameters
 NAME 4
 EXPNO 1
 PROCNO 1

F2 - Acquisition Parameters
 Date_ 20220224
 Time 23.59 h
 INSTRUM Avance
 PROBHD Z116098_0833 (zg30
 PULPROG 65536
 TD 16
 SOLVENT CDCl₃
 NS 2
 DS 16
 SWH 8196.722 Hz
 FIDRES 0.250144 Hz
 AQ 3.9976959 sec
 RG 101
 DW 61.000 usec
 DE 13.54 usec
 TE 294.8 K
 D1 1.0000000 sec
 TD0 1
 SFO1 400.1324708 MHz
 NUC1 1H
 P0 3.33 usec
 P1 10.00 usec
 PLW1 20.73200035 W

F2 - Processing parameters
 SI 65536
 SF 400.1300097 MHz
 WDW EM
 SSB 0
 LB 0.30 Hz
 GB 0
 PC 1.00

**4**

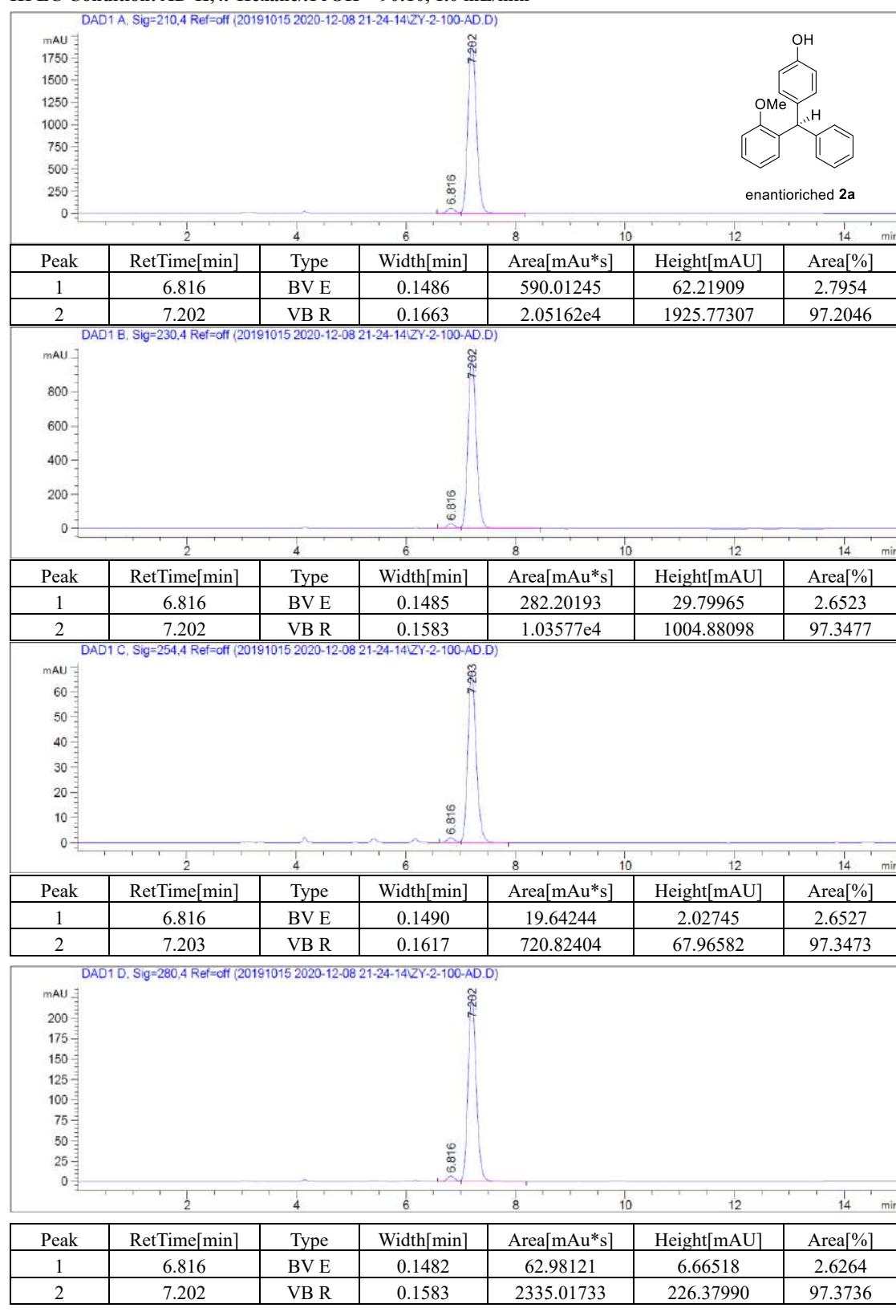
Current Data Parameters
 NAME 4
 EXPNO 2
 PROCNO 1

F2 - Acquisition Parameters
 Date_ 20220225
 Time 0.59 h
 INSTRUM Avance
 PROBHD Z116098_0833 (zgpg30
 PULPROG 65536
 TD 1024
 SOLVENT CDCl3
 NS 4
 DS 23809.523 Hz
 SWH 0.726609 Hz
 FIDRES 1.3762560 sec
 AQ 50.1934
 RG 21.000 usec
 DW 6.50 usec
 DE 295.3 K
 TE 2.00000000 sec
 D1 0.03000000 sec
 D11 1
 TD0 100.6228298 MHz
 SFO1 13C
 NUC1 3.33 usec
 P0 10.00 usec
 PLW1 87.89900208 W
 SFO2 400.1316005 MHz
 NUC2 1H
 CPDPRG[2] waltz65
 PCPD2 90.00 usec
 PLW2 20.73200035 W
 PLW12 0.25595000 W
 PLW13 0.12874000 W

F2 - Processing parameters
 SI 32768
 SF 100.6127748 MHz
 WDW EM
 SSB 0
 LB 1.00 Hz
 GB 0
 PC 1.40

Sample Name: ZY-2-100-AD

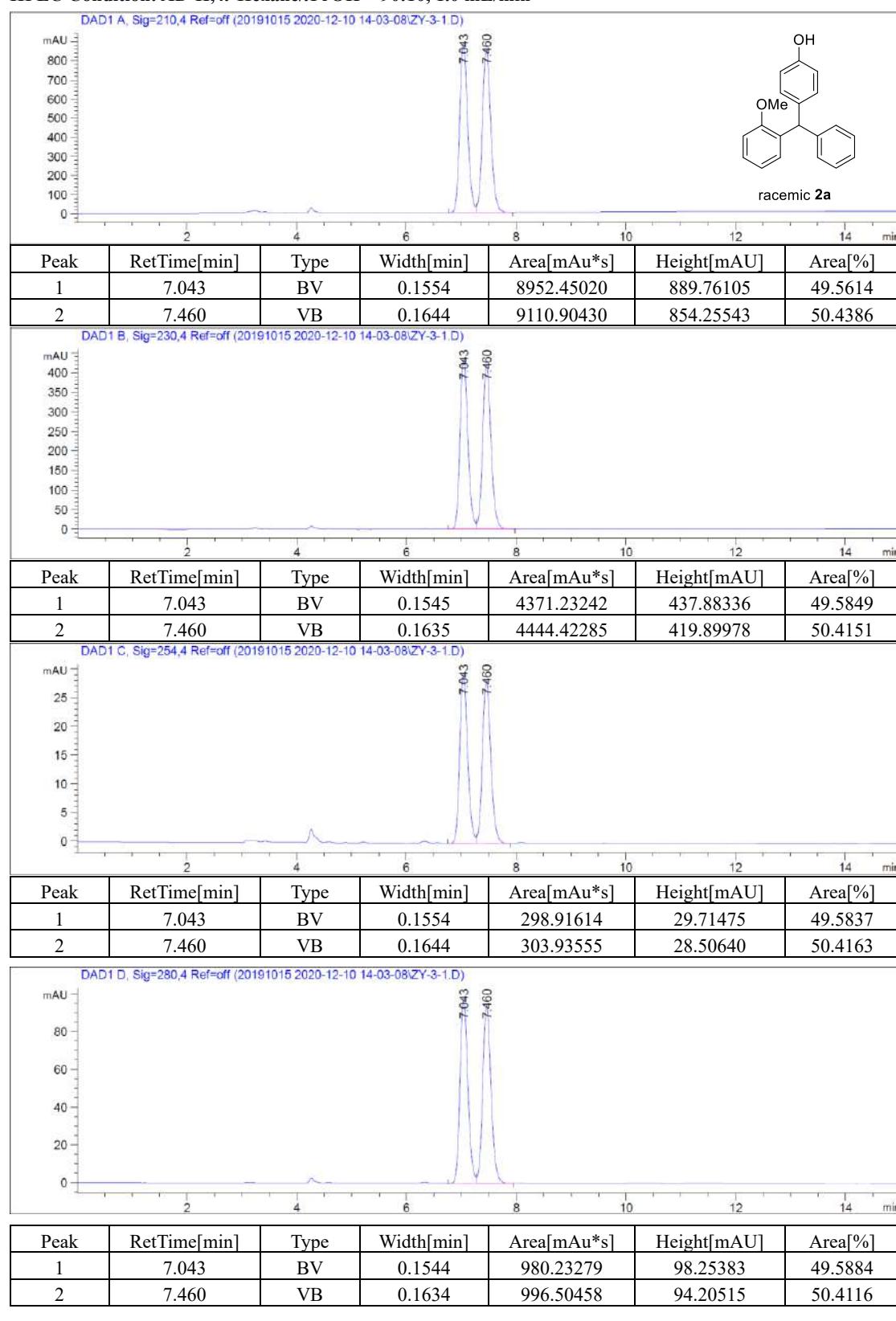
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-1

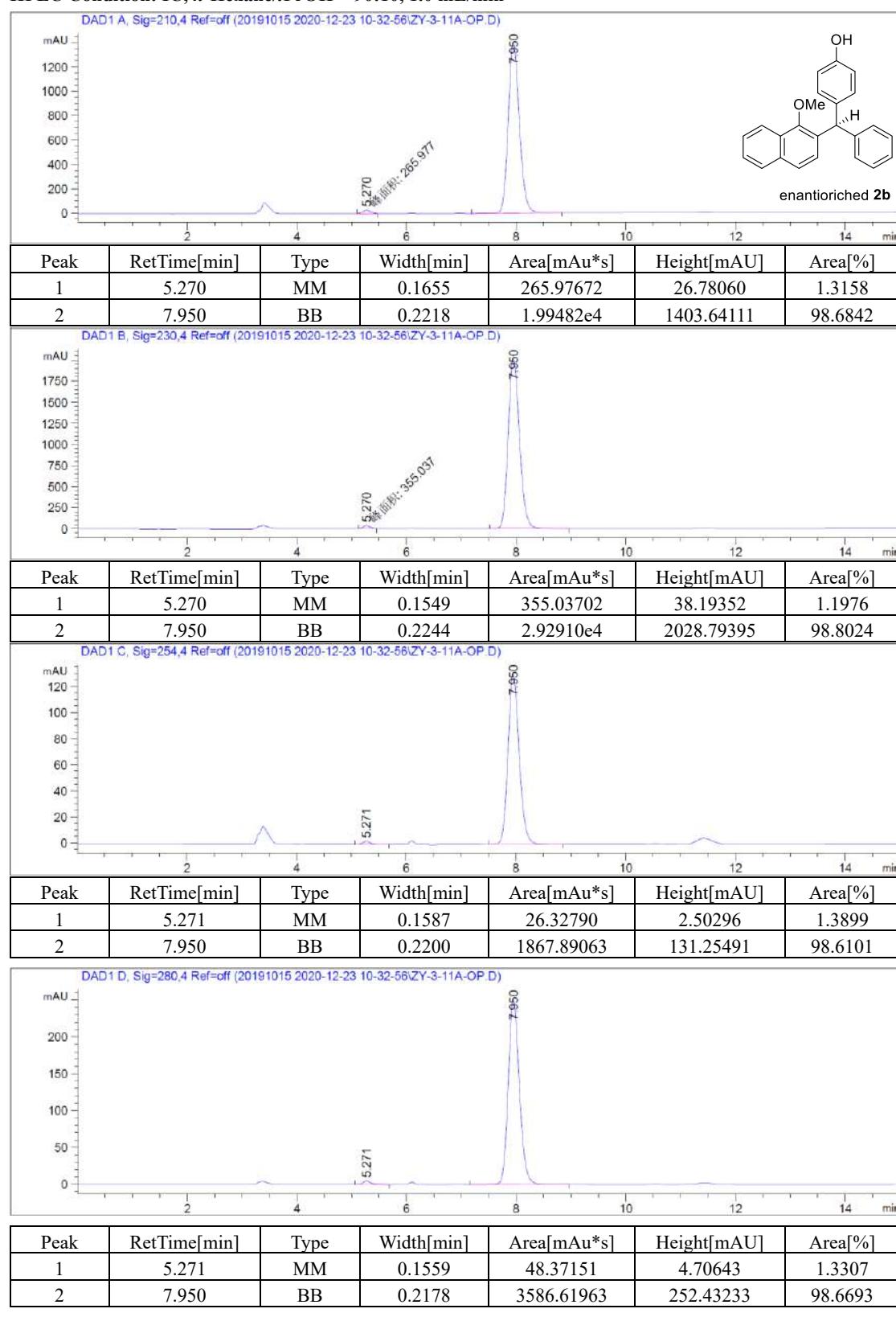
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-11-A-OP

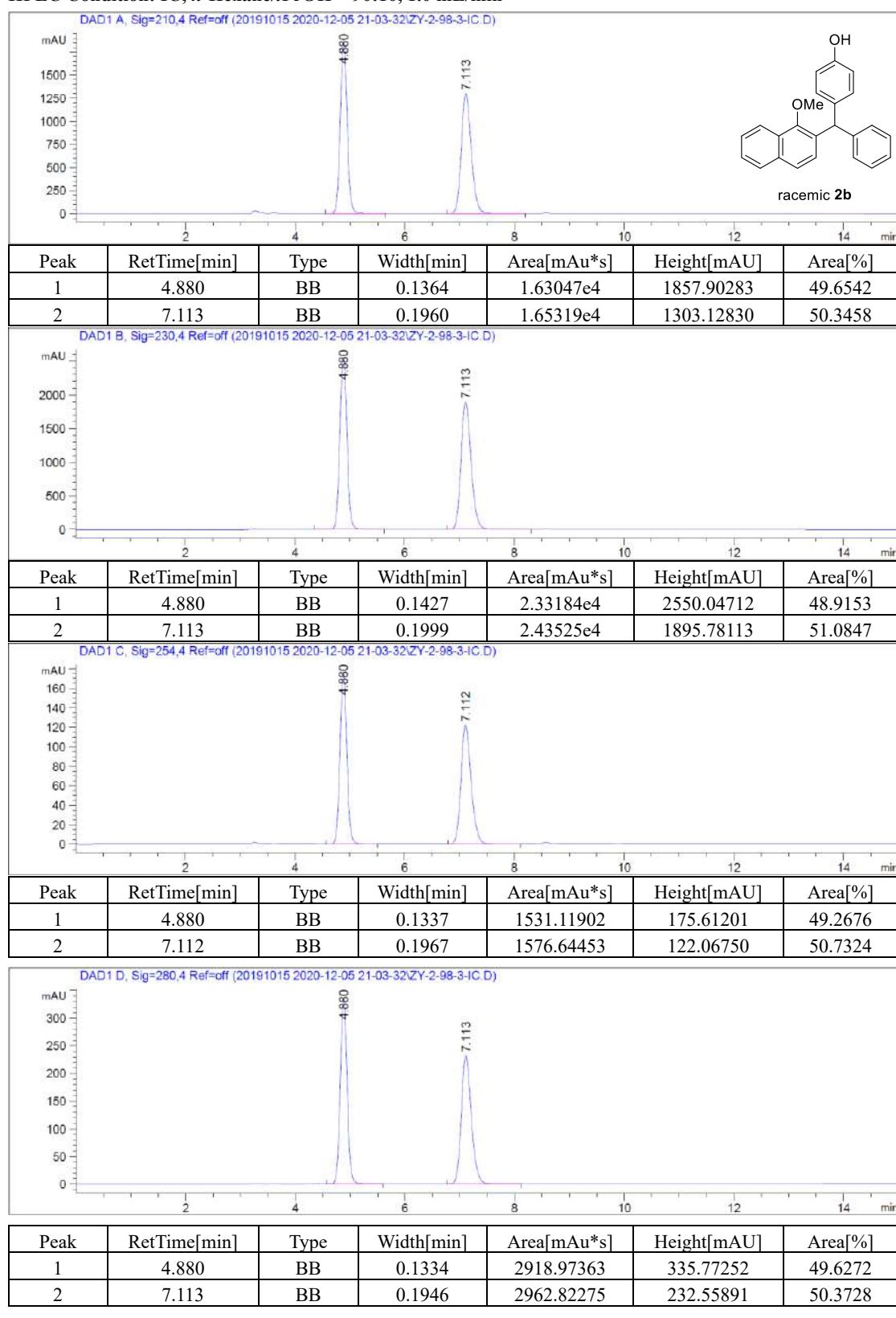
HPLC Condition: IC, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-2-98-3-IC

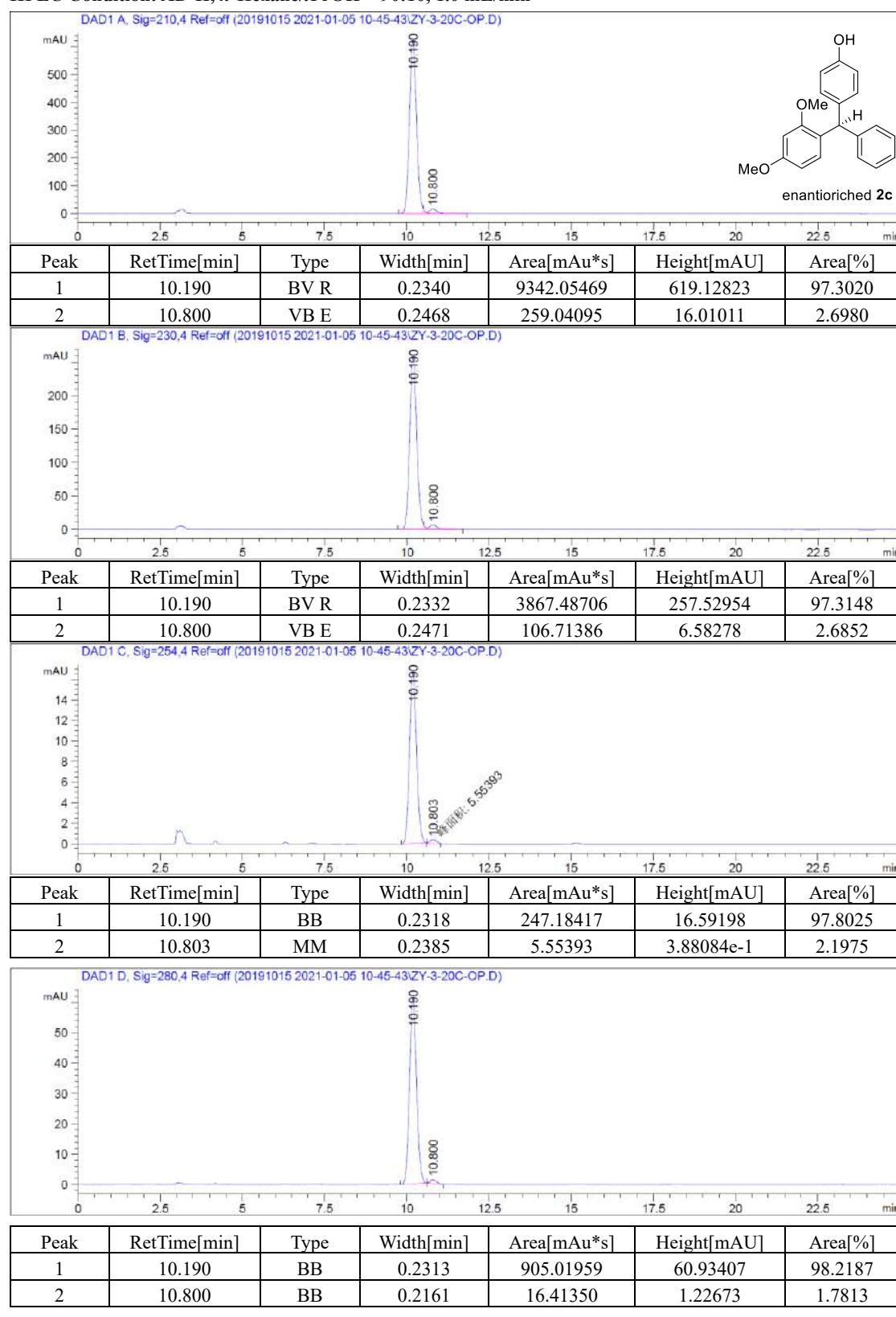
HPLC Condition: IC, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-20C-OP

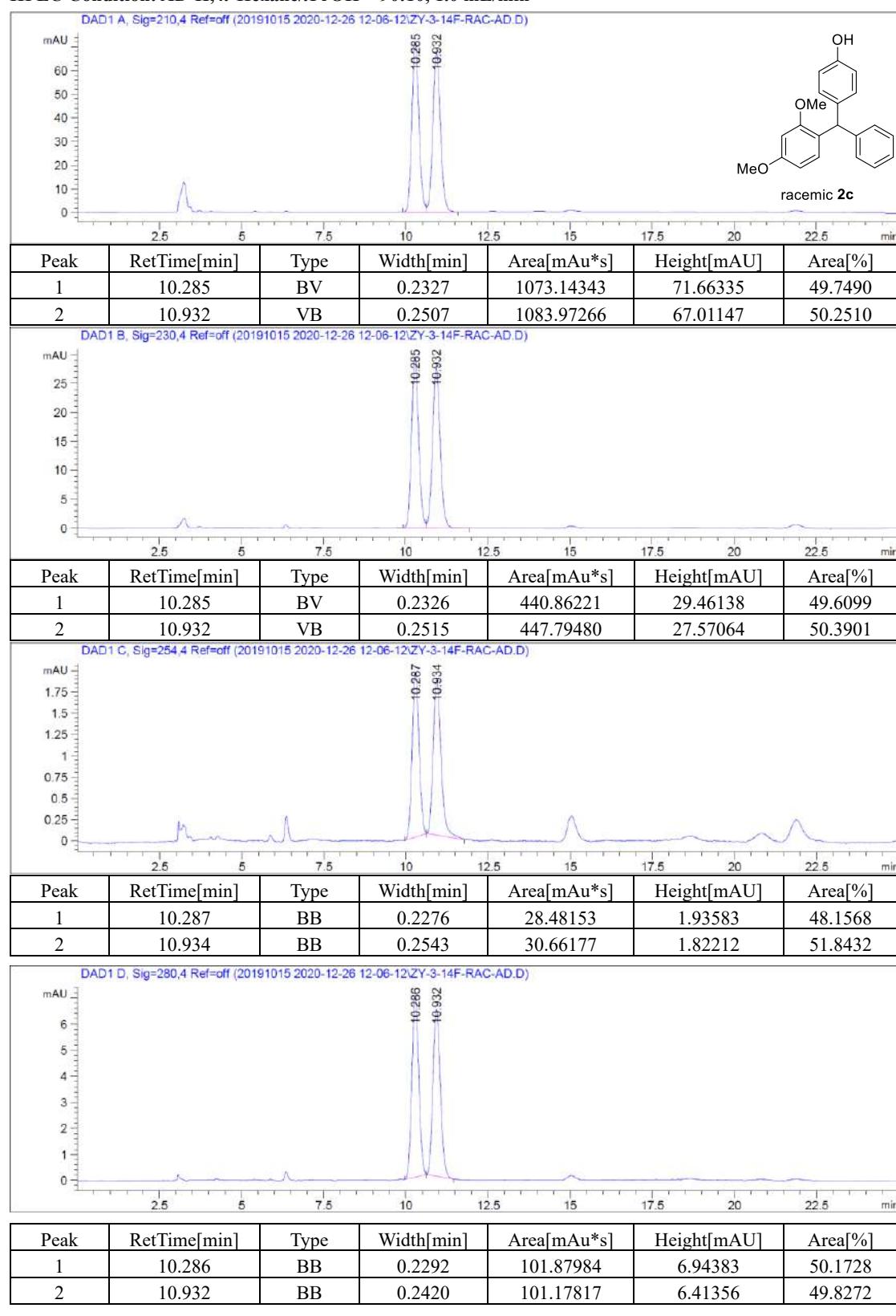
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-14F-RAC-AD

HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-11C-OP

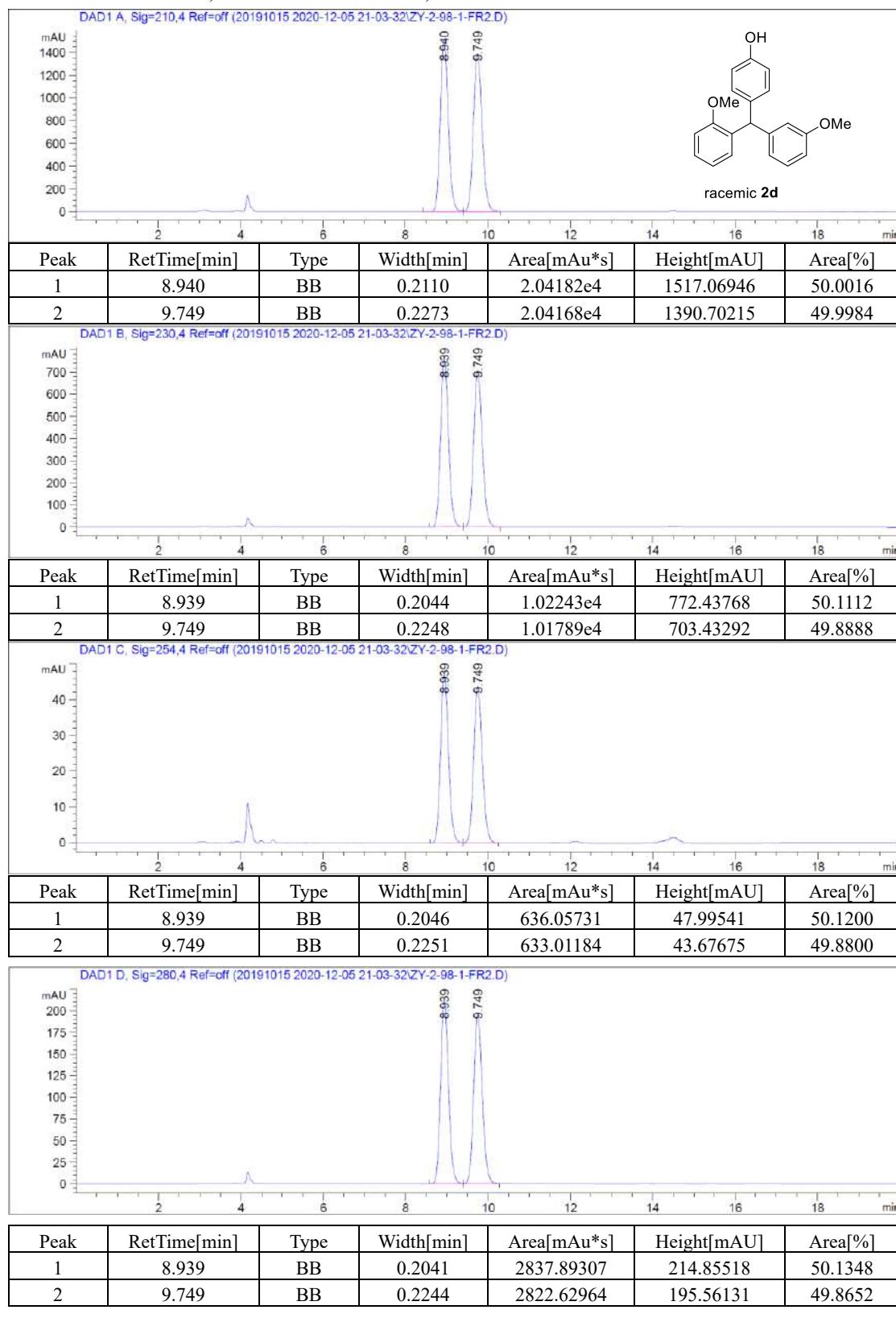
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-2-98-1-FR2

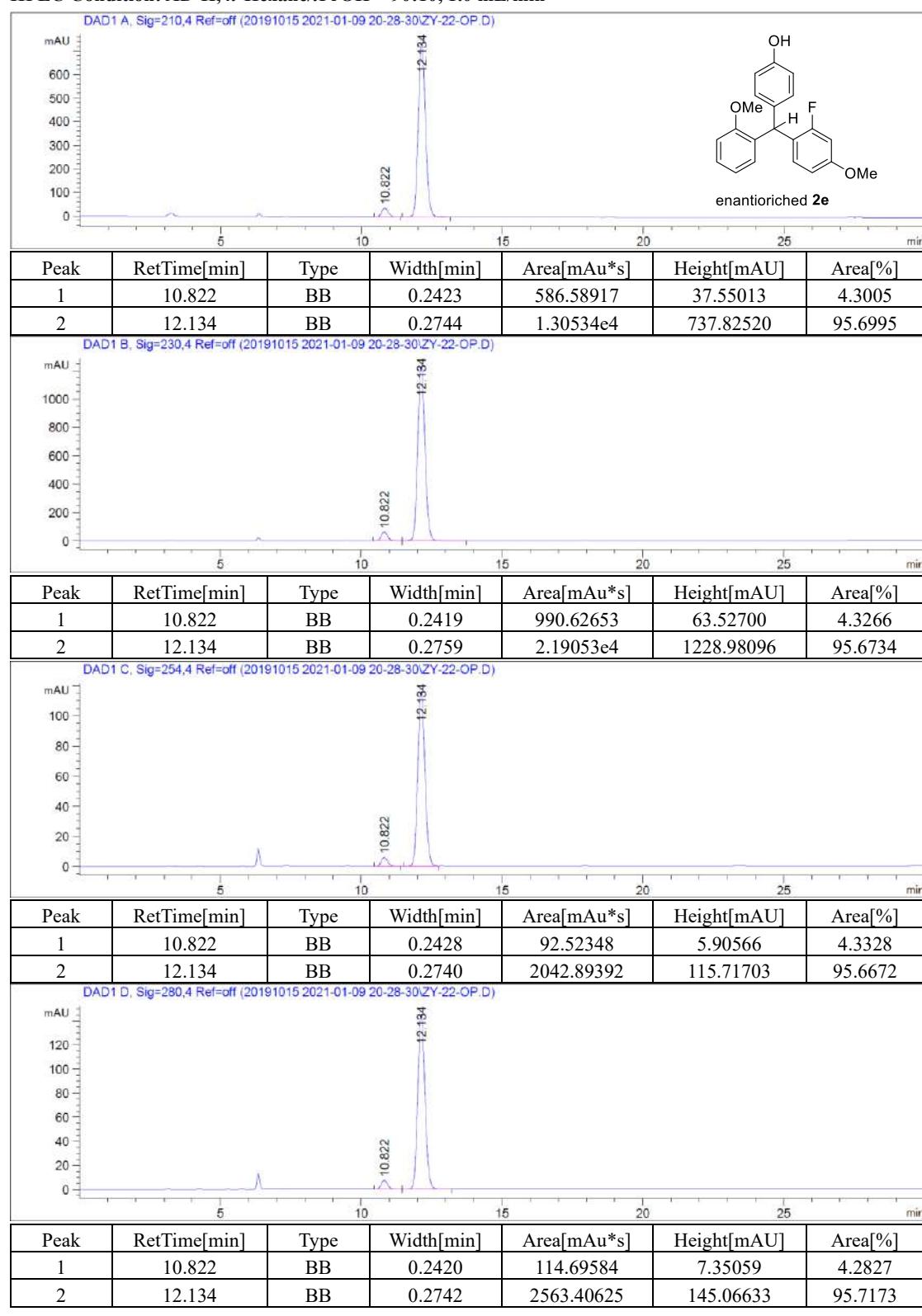
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-22-OP

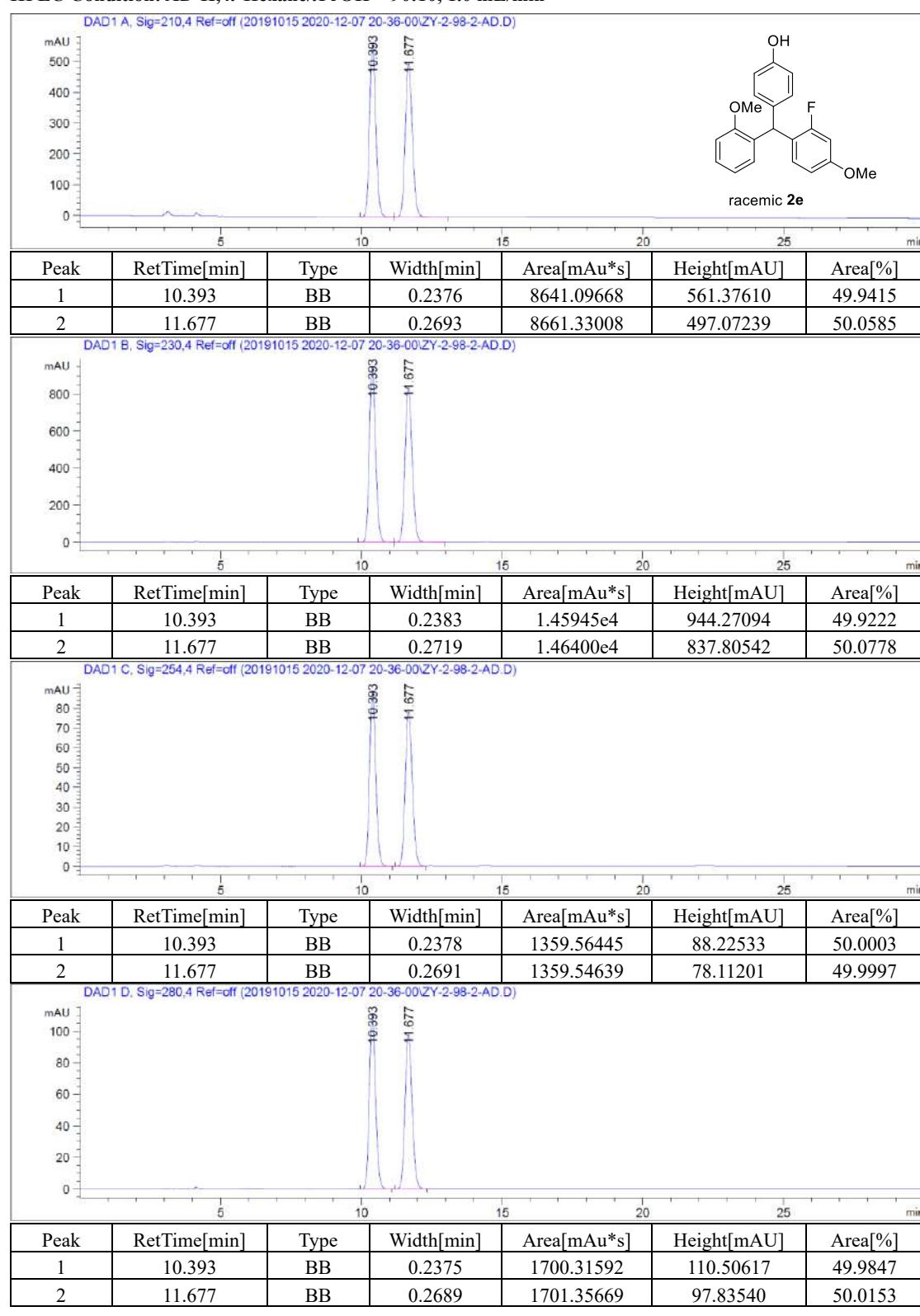
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-2-98-2-AD

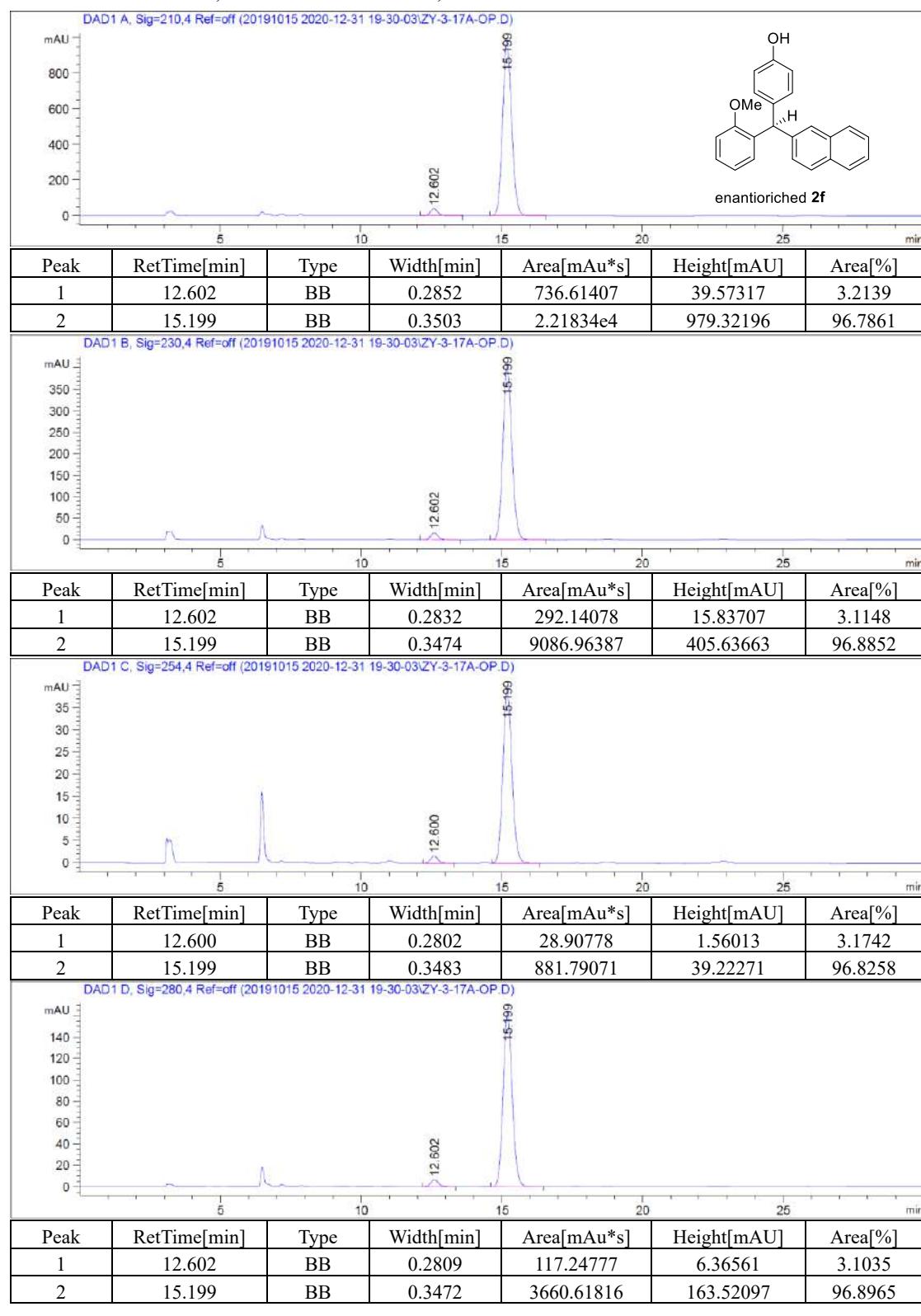
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-17A-OP

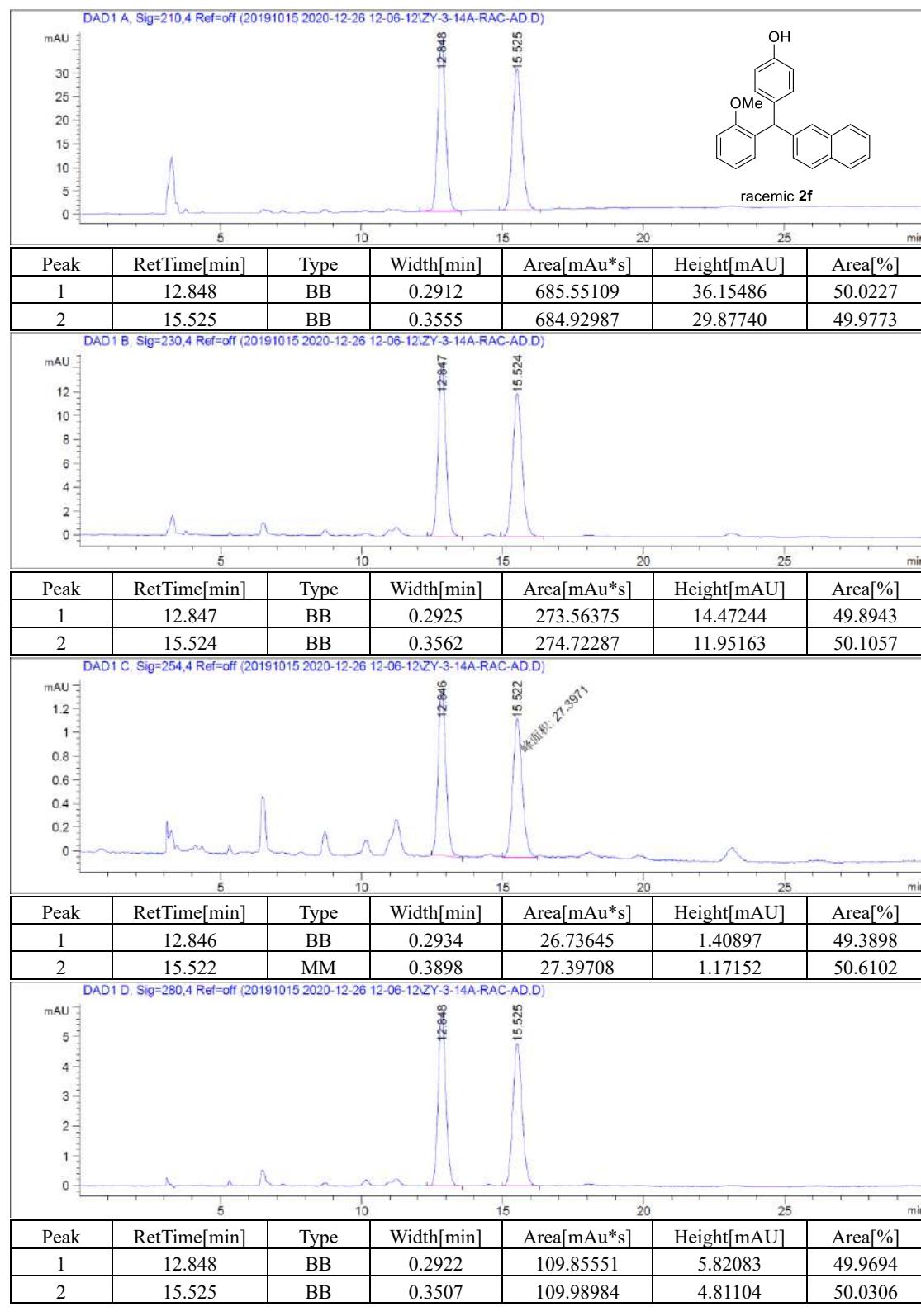
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-14A-RAC-AD

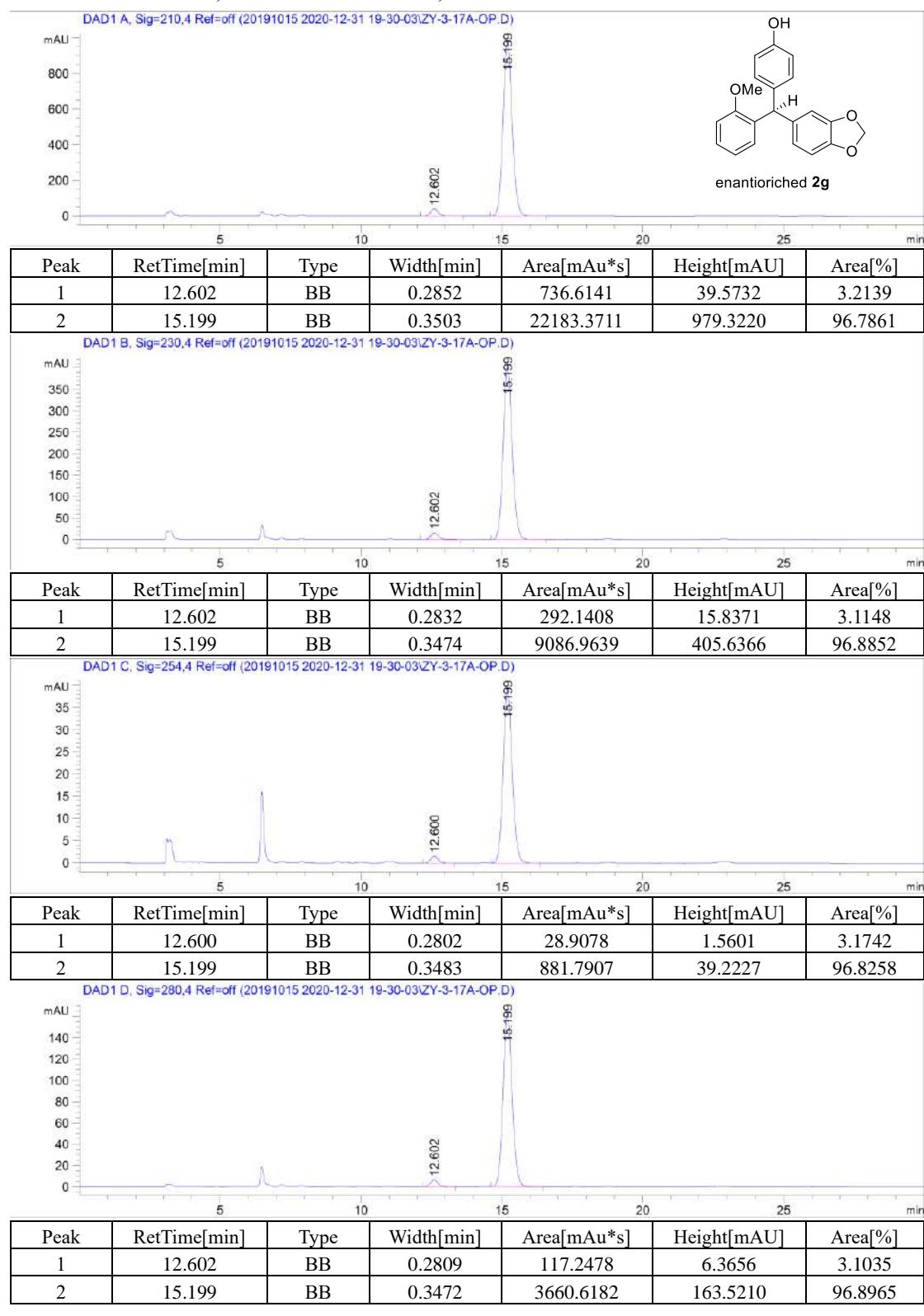
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-17A-OP

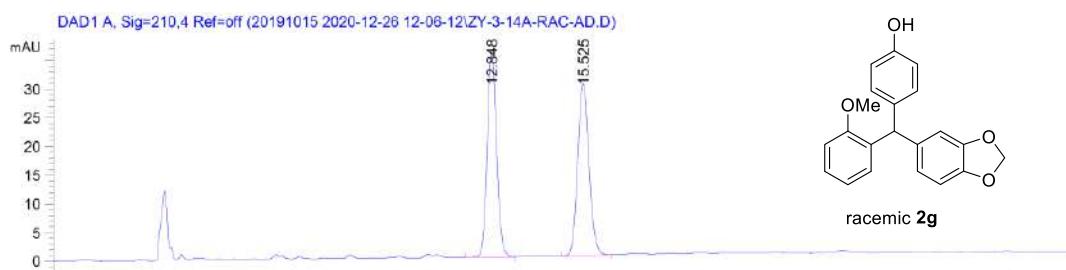
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



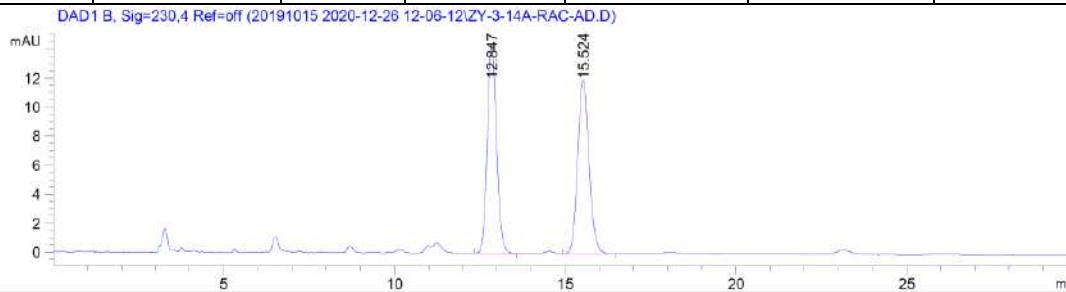
End of Report

Sample Name: ZY-3-14A-RAC-AD

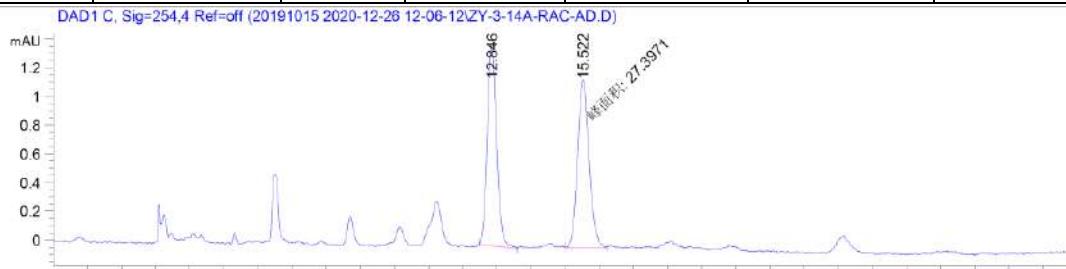
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



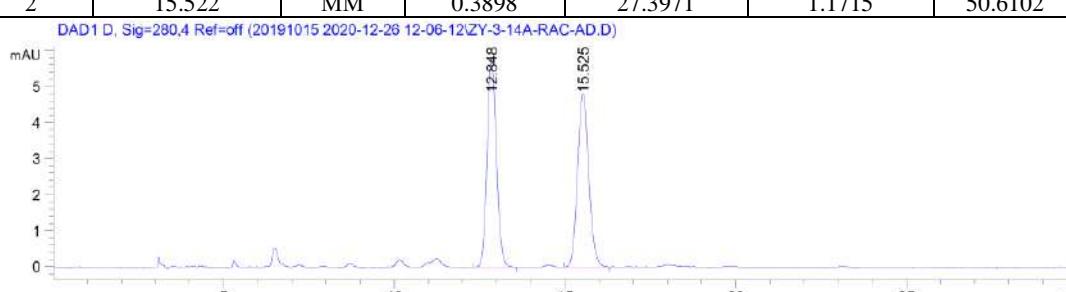
Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	12.848	BB	0.2912	685.5511	36.1549	50.0227
2	15.525	BB	0.3555	684.9299	29.8774	49.9773



Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	12.847	BB	0.2925	273.5638	14.4724	49.8943
2	15.524	BB	0.3562	274.7229	11.9516	50.1057



Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	12.846	BB	0.2934	26.7365	1.4090	49.3898
2	15.522	MM	0.3898	27.3971	1.1715	50.6102

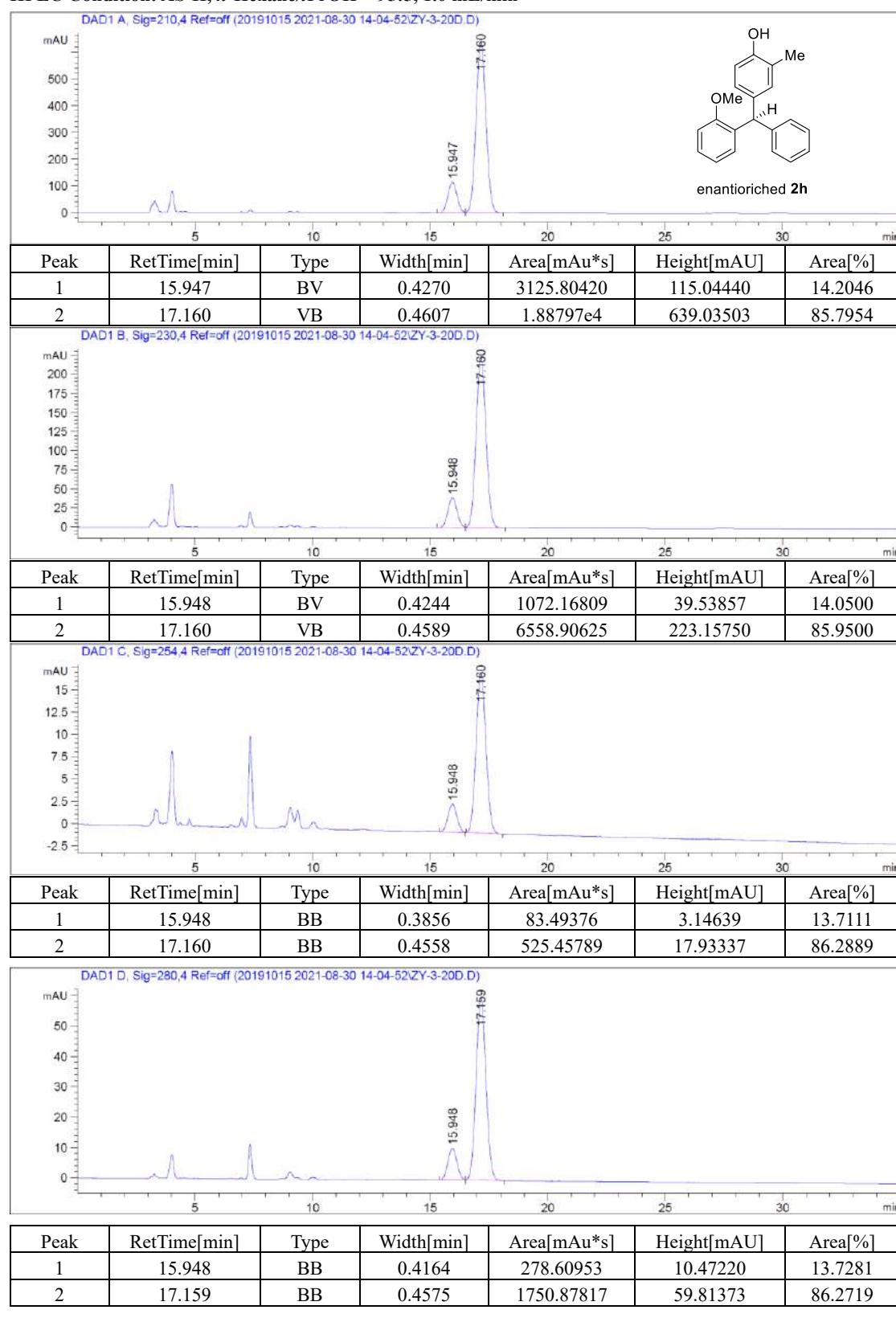


Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	12.848	BB	0.2922	109.8555	5.8208	49.9694
2	15.525	BB	0.3507	109.9898	4.8110	50.0306

End of Report

Sample Name: ZY-3-20D

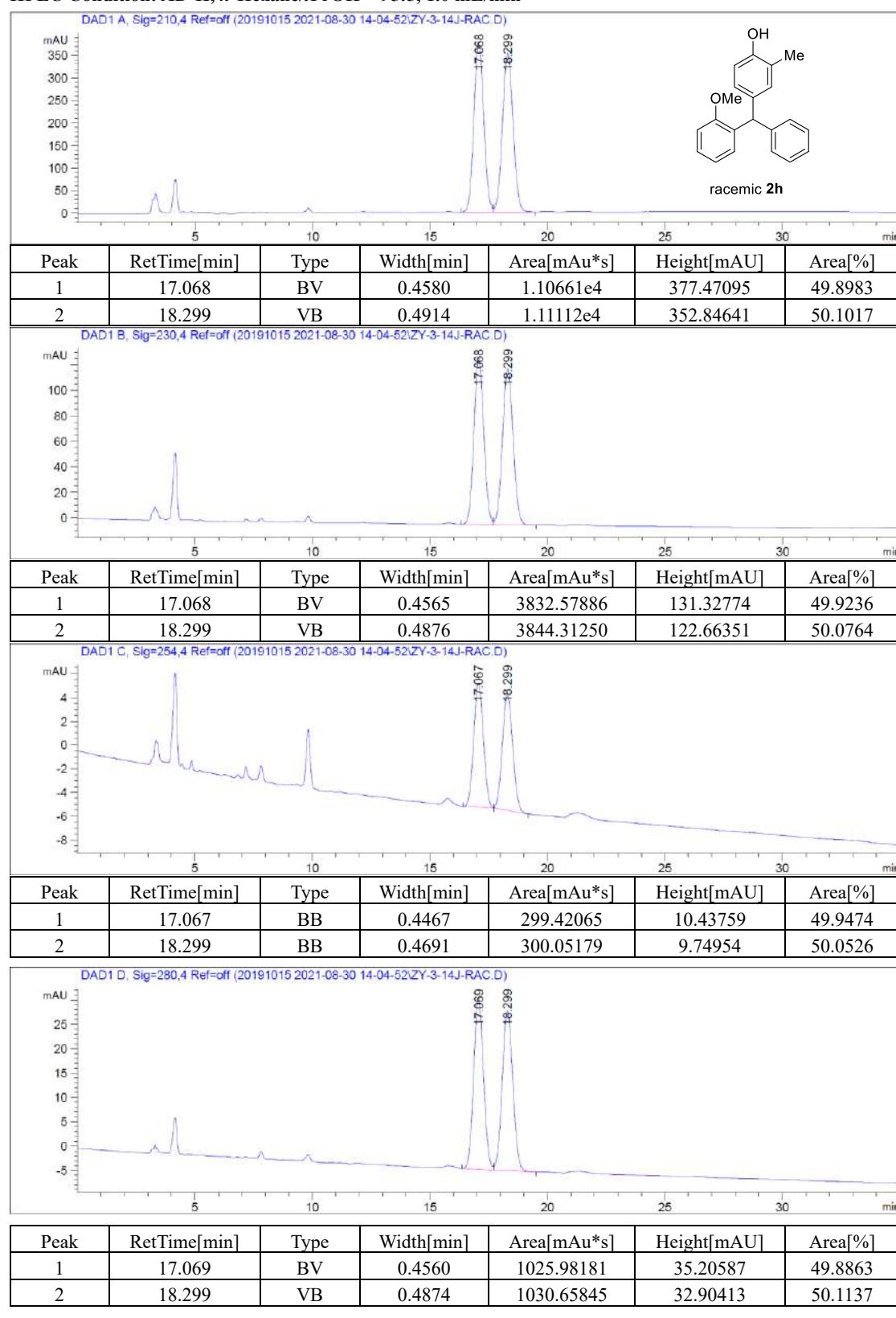
HPLC Condition: AS-H, *n*-Hexane/iPrOH = 95:5, 1.0 mL/min



End of Report

Sample Name: ZY-3-14J-RAC

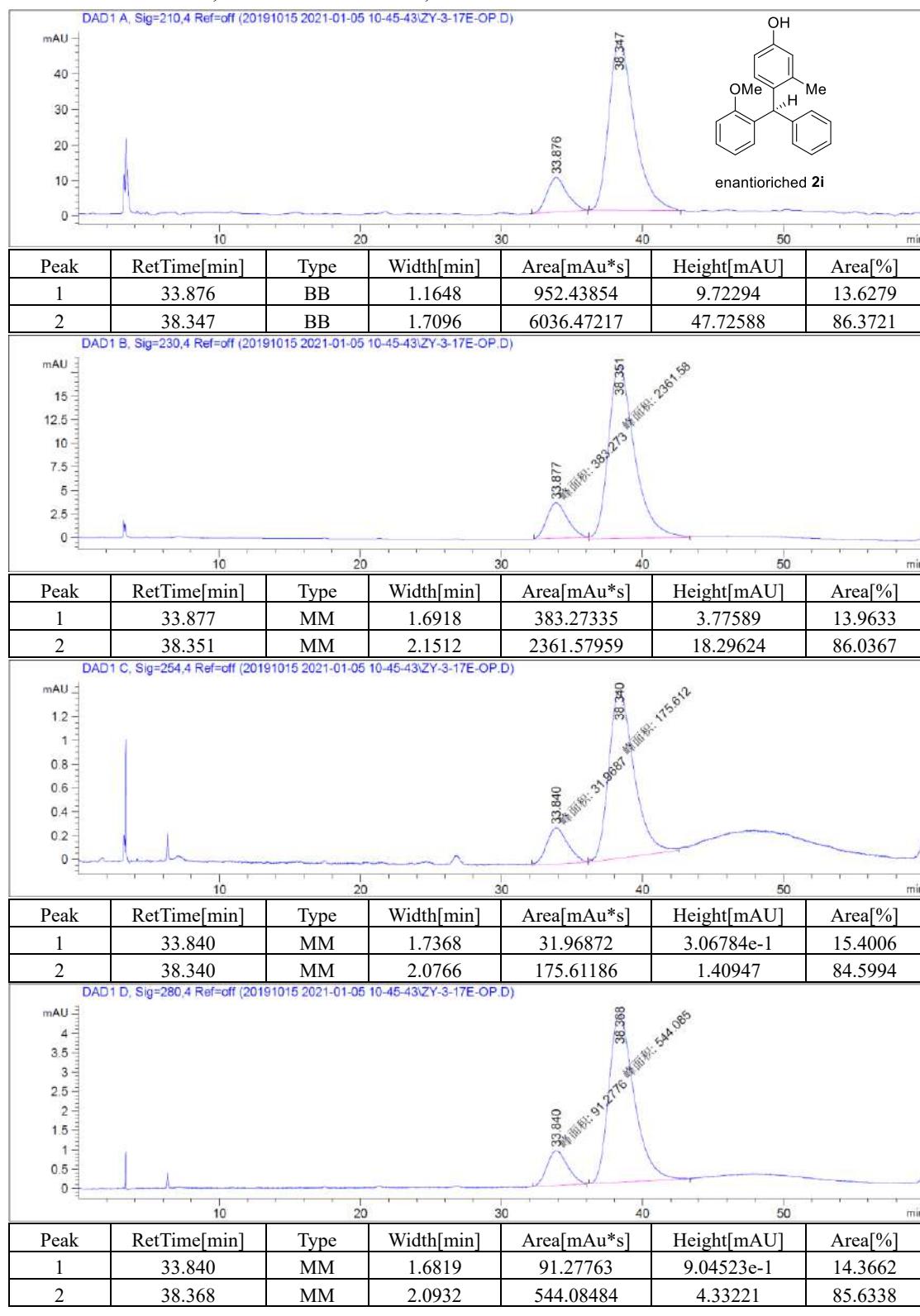
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 95:5, 1.0 mL/min



End of Report

Sample Name: ZY-3-17E-OP

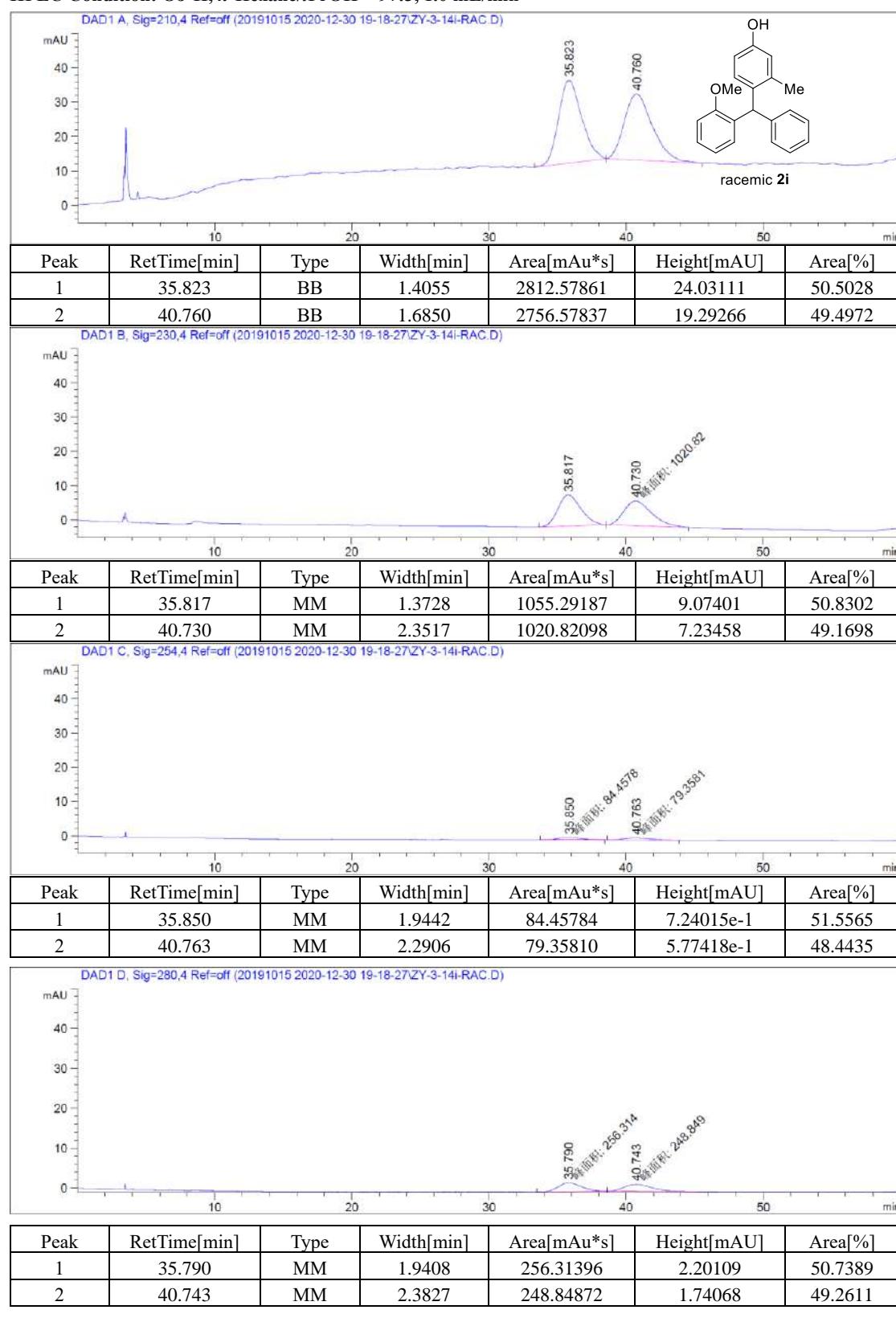
HPLC Condition: OJ-H, *n*-Hexane/iPrOH = 97:3, 1.0 mL/min



End of Report

Sample Name: ZY-3-14i-RAC

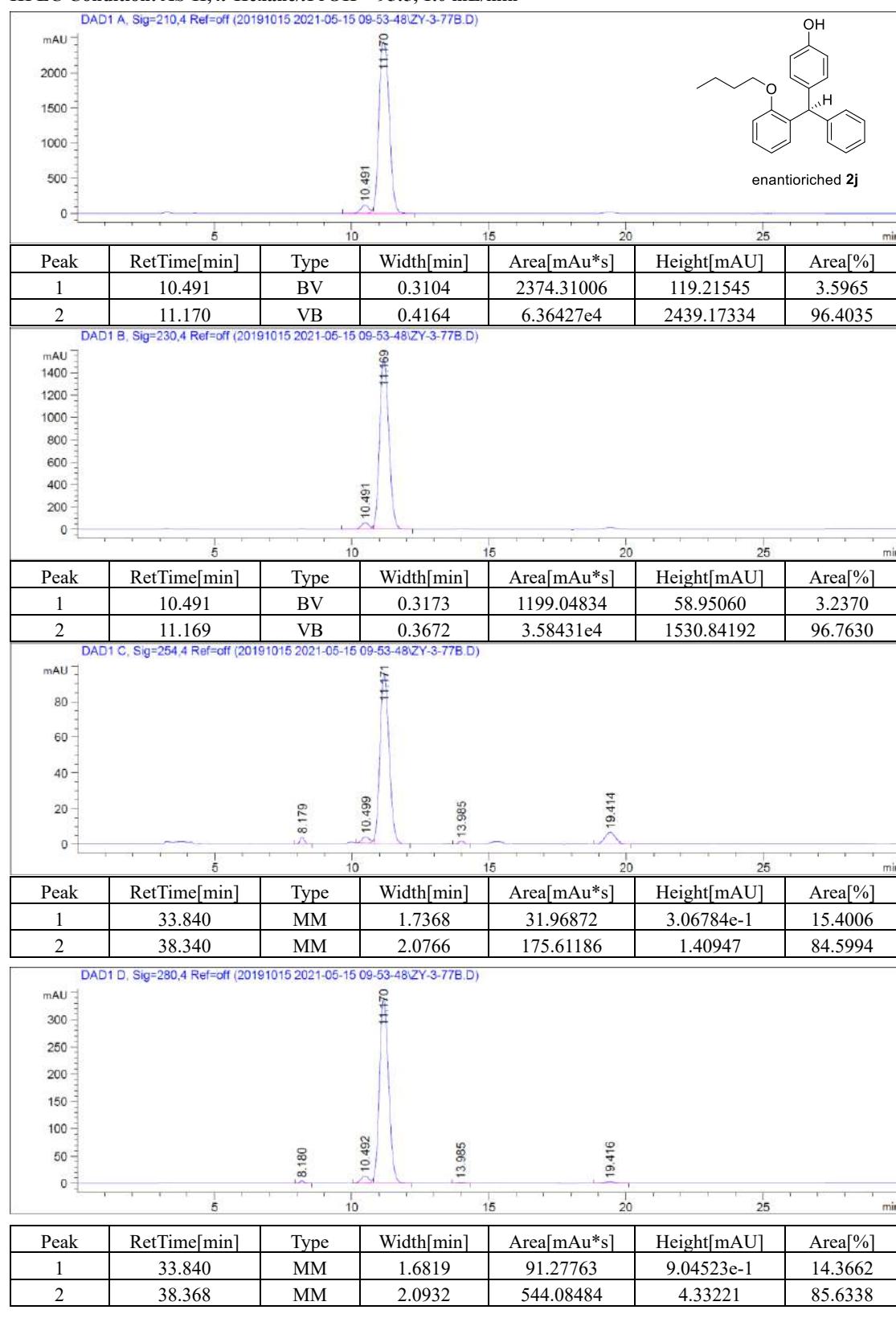
HPLC Condition: OJ-H, *n*-Hexane/iPrOH = 97:3, 1.0 mL/min



End of Report

Sample Name: ZY-3-77B

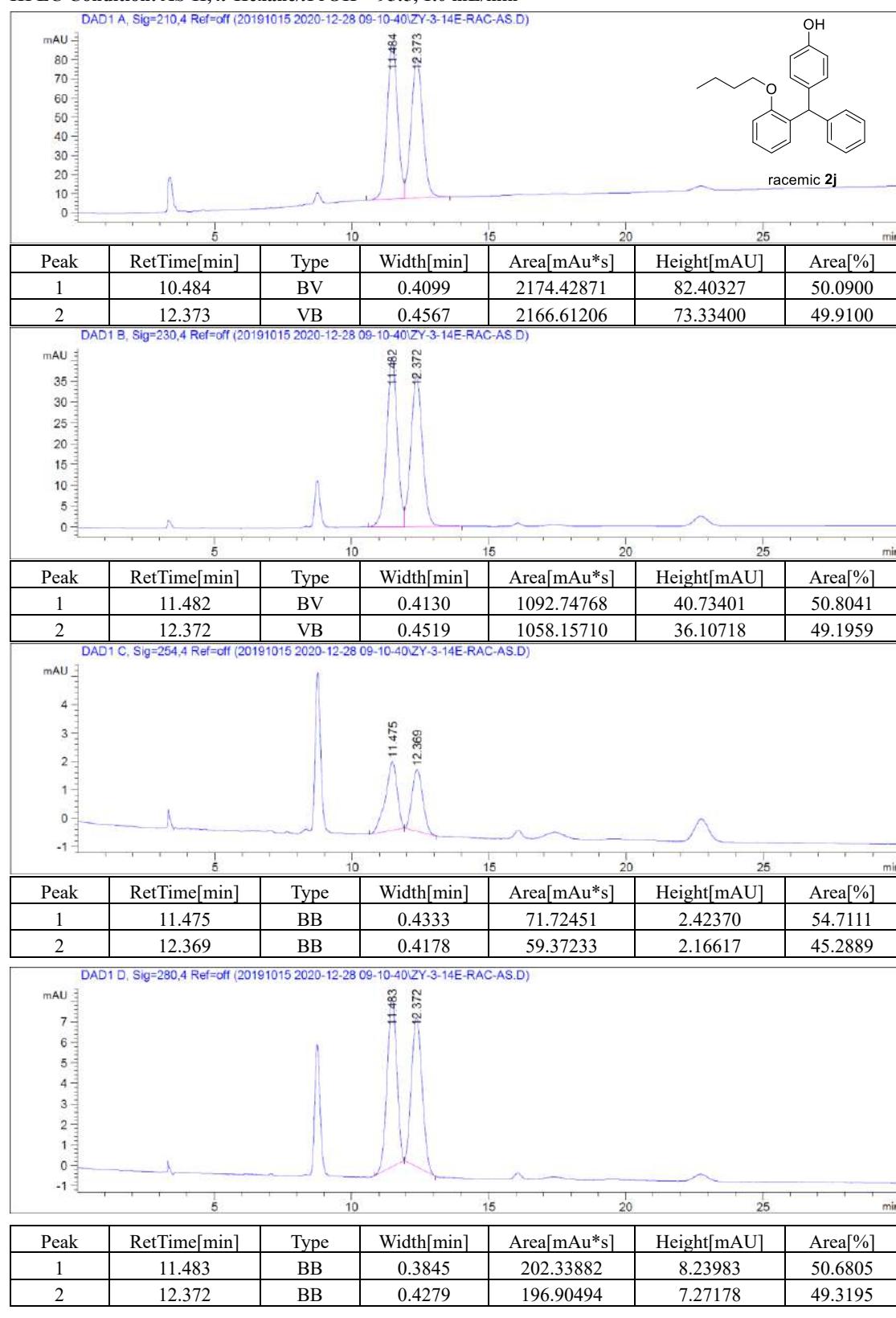
HPLC Condition: AS-H, *n*-Hexane/iPrOH = 95:5, 1.0 mL/min



End of Report

Sample Name: ZY-3-14E-RAC-AS

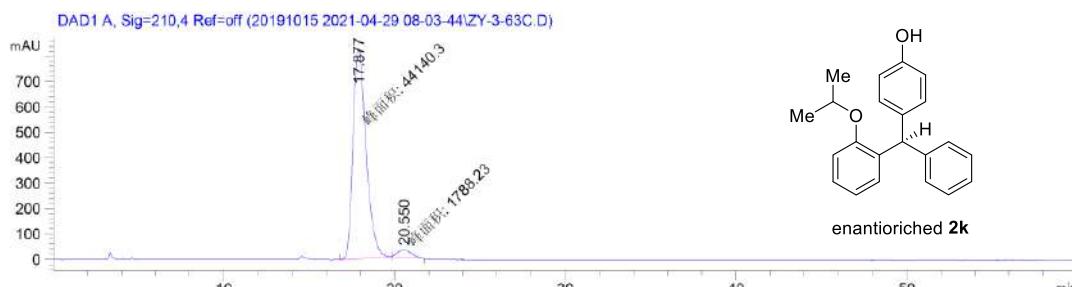
HPLC Condition: AS-H, *n*-Hexane/iPrOH = 95:5, 1.0 mL/min



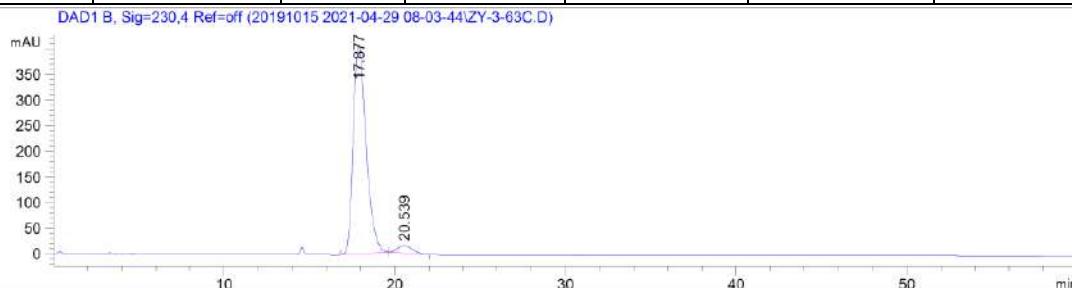
End of Report

Sample Name: ZY-3-63C

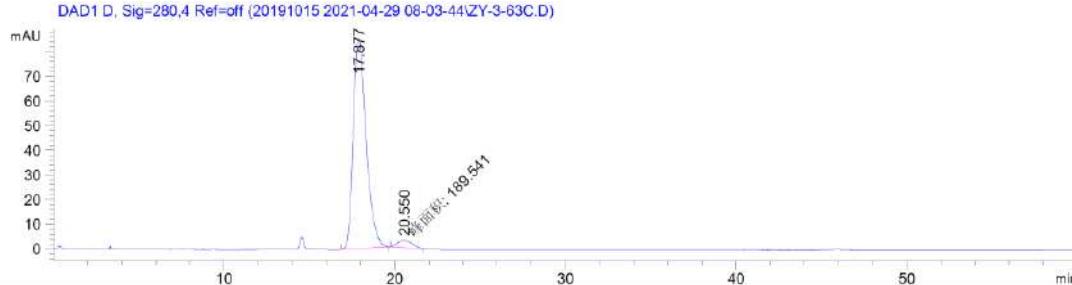
HPLC Condition: OJ-H, *n*-Hexane/iPrOH = 95:5, 1.0 mL/min



Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	17.877	BV R	0.8222	44867.2891	833.2604	94.3494
2	20.538	VB E	0.9998	2687.1086	38.3364	5.6506



Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	17.877	MM	0.8744	21467.5938	409.2059	95.9378
2	20.563	MM	0.9903	908.9843	15.2979	4.0622

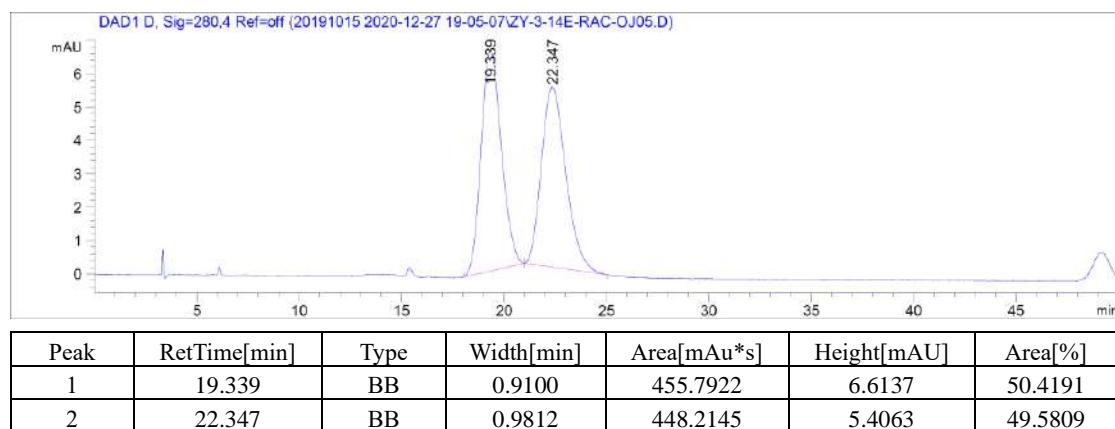
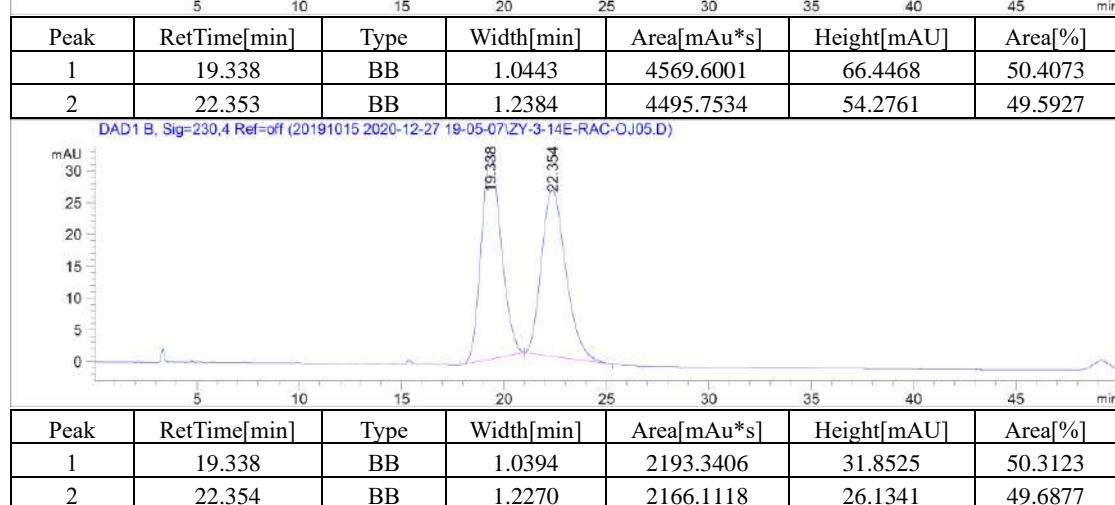
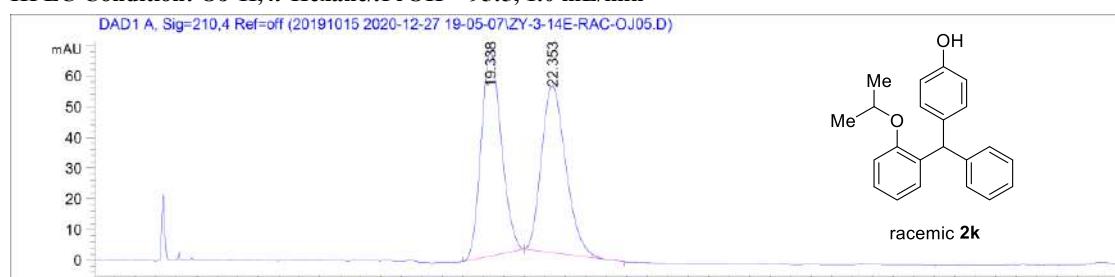


Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	17.877	MM	0.8806	4522.0562	85.5881	95.3451
2	20.550	MM	1.0660	220.7724	3.4516	4.6549

End of Report

Sample Name: ZY-3-14E-RAC-OJ05

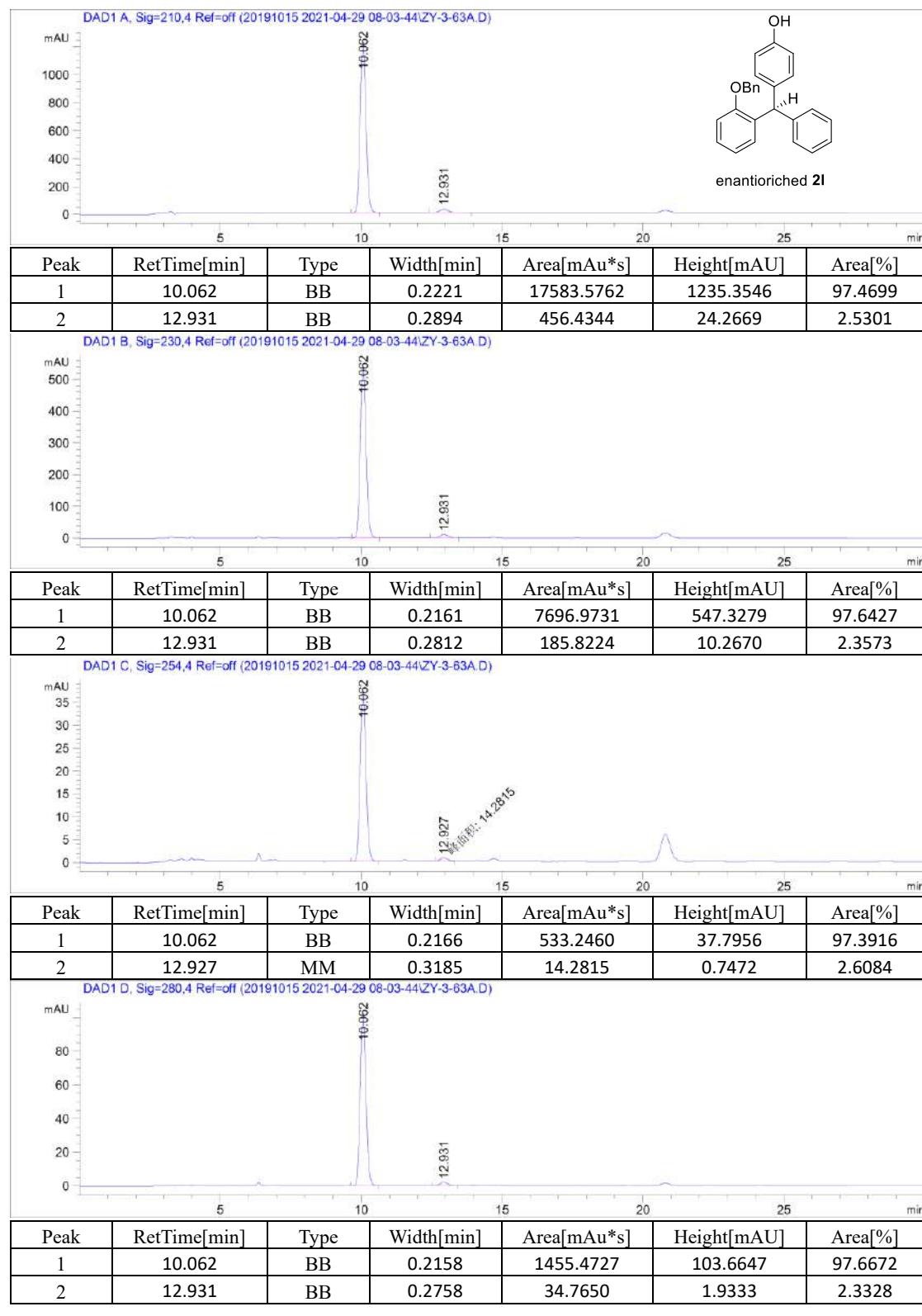
HPLC Condition: OJ-H, *n*-Hexane/iPrOH = 95:5, 1.0 mL/min



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End of Report

Sample Name: ZY-3-63A

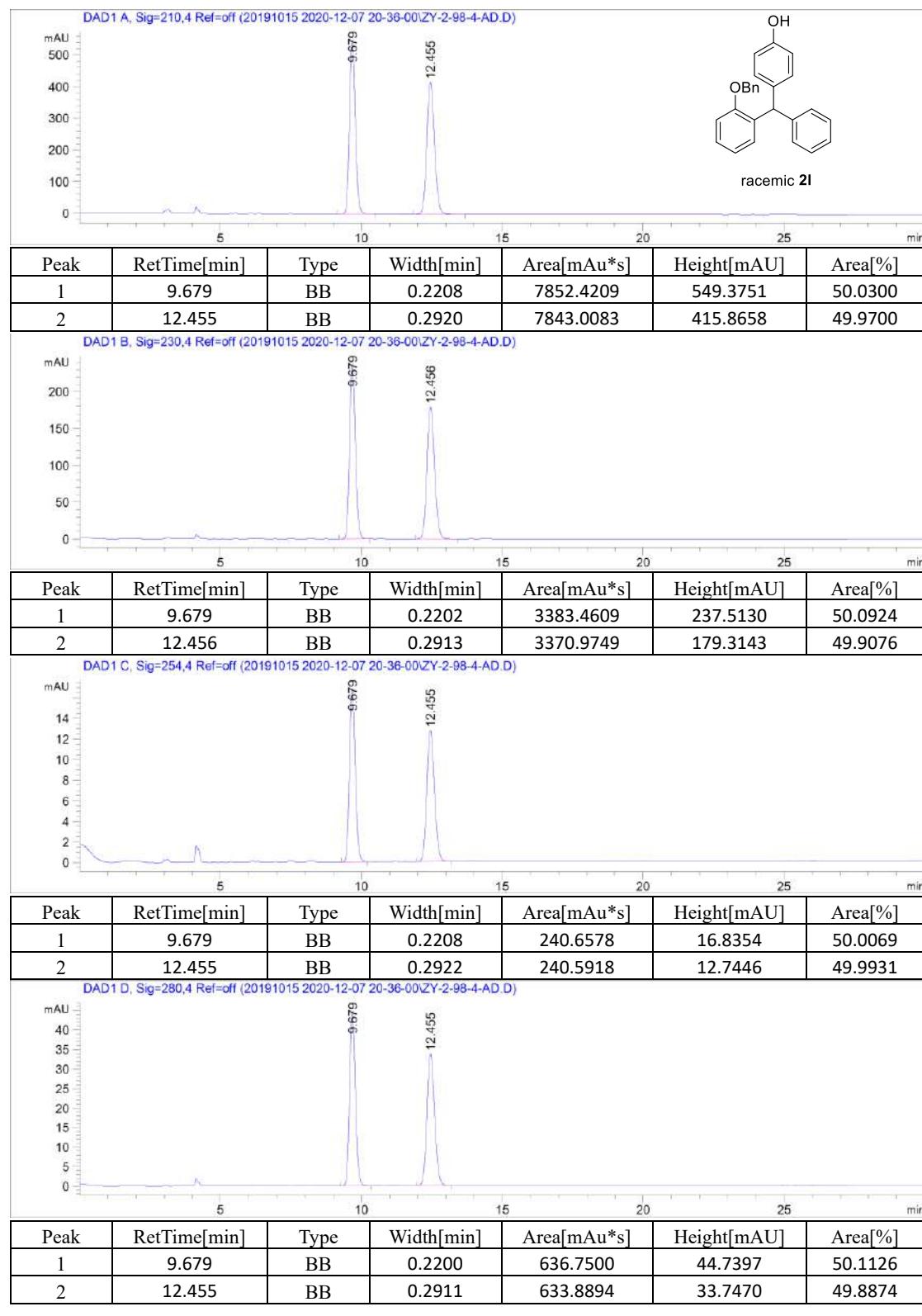
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-2-98-4-AD

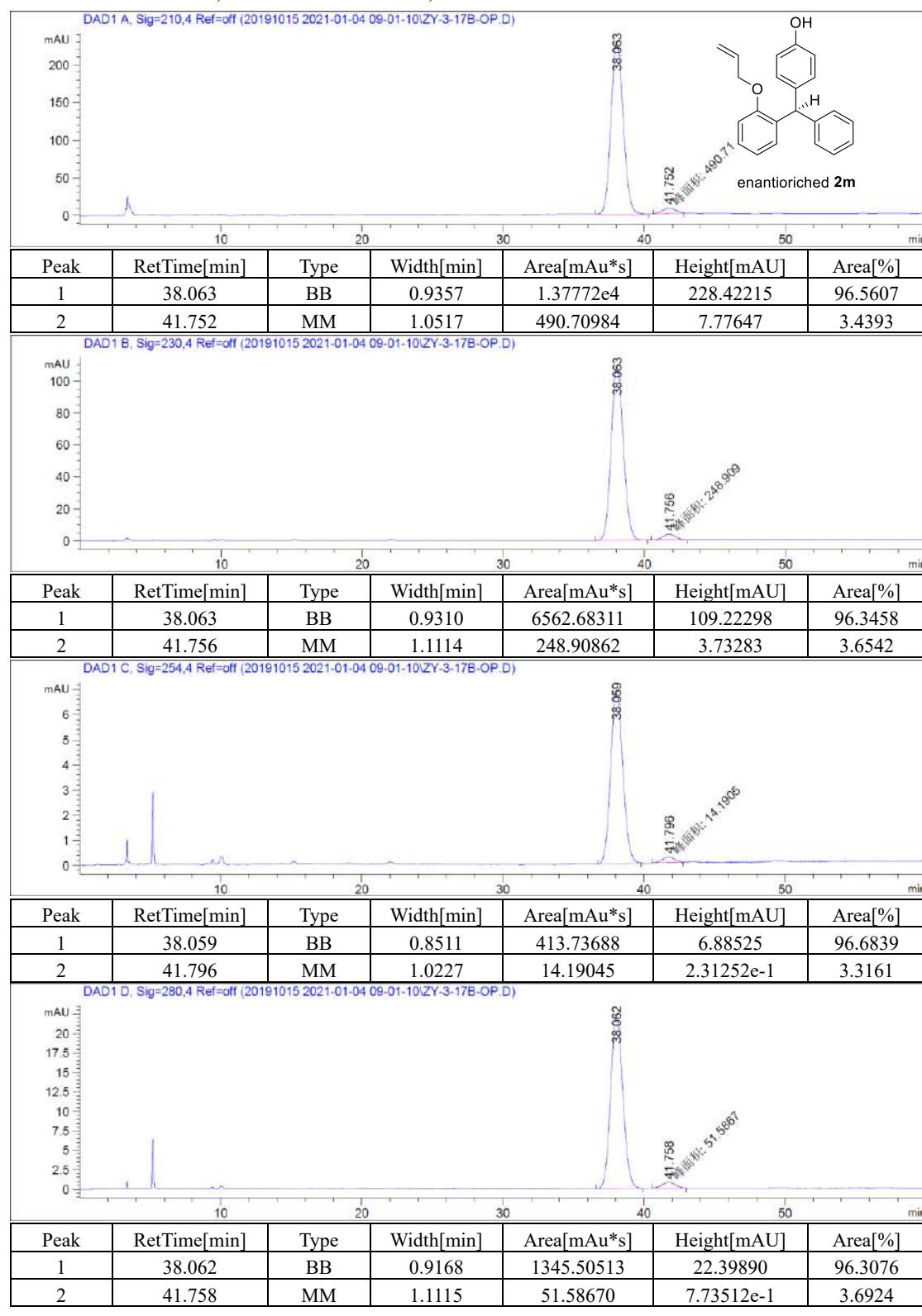
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-17B-OP

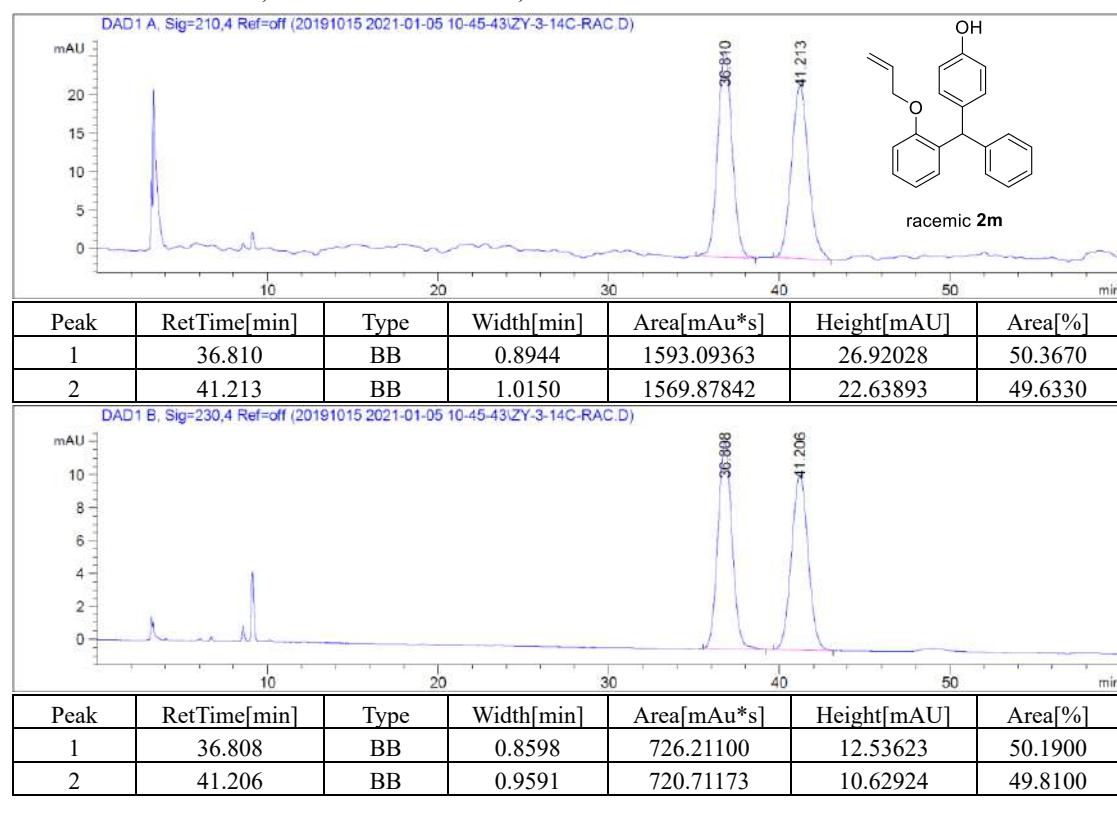
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 98:2, 1.0 mL/min



End of Report

Sample Name: ZY-3-14C-RAC

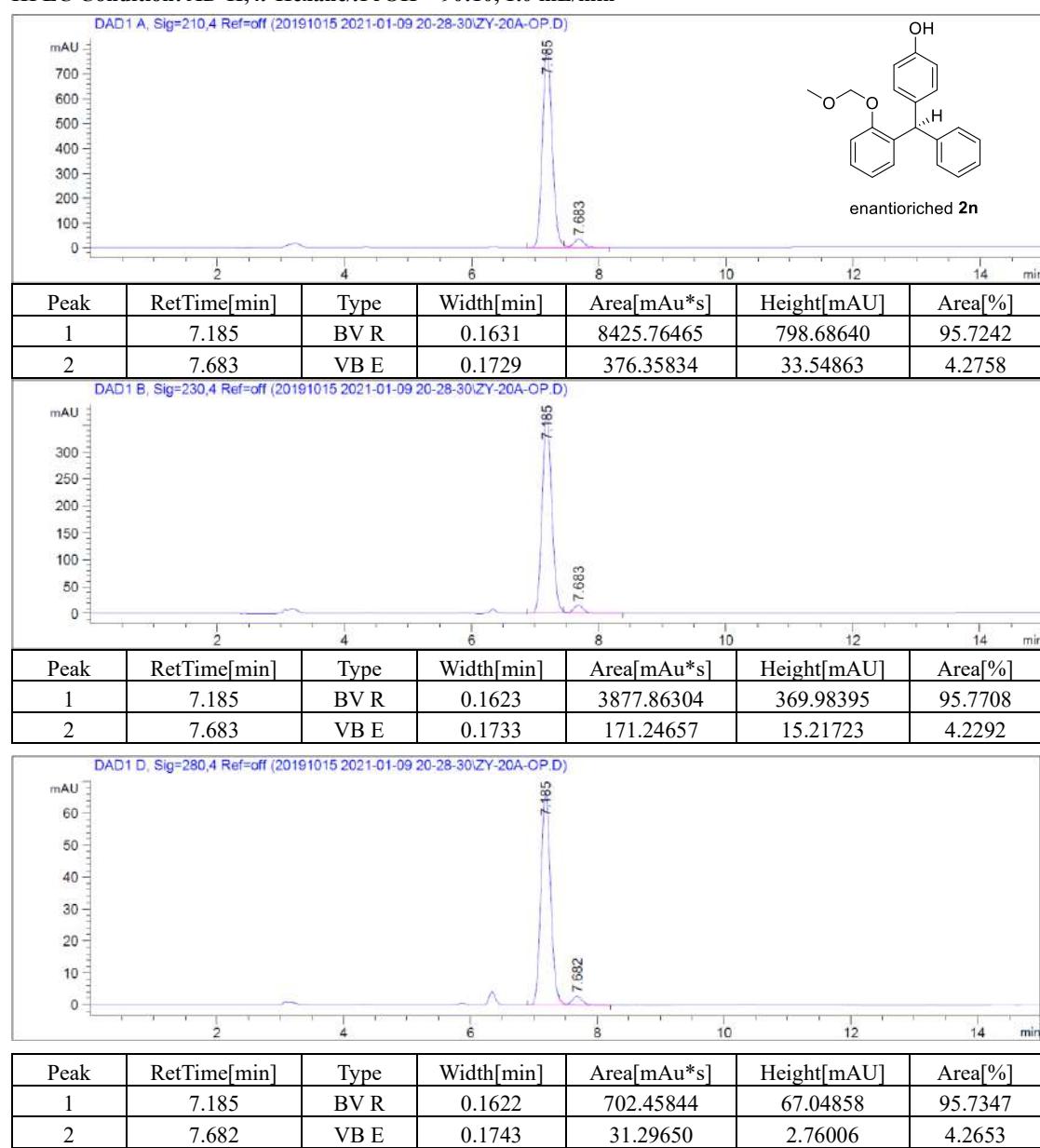
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 98:2, 1.0 mL/min



End of Report

Sample Name: ZY-20A-OP

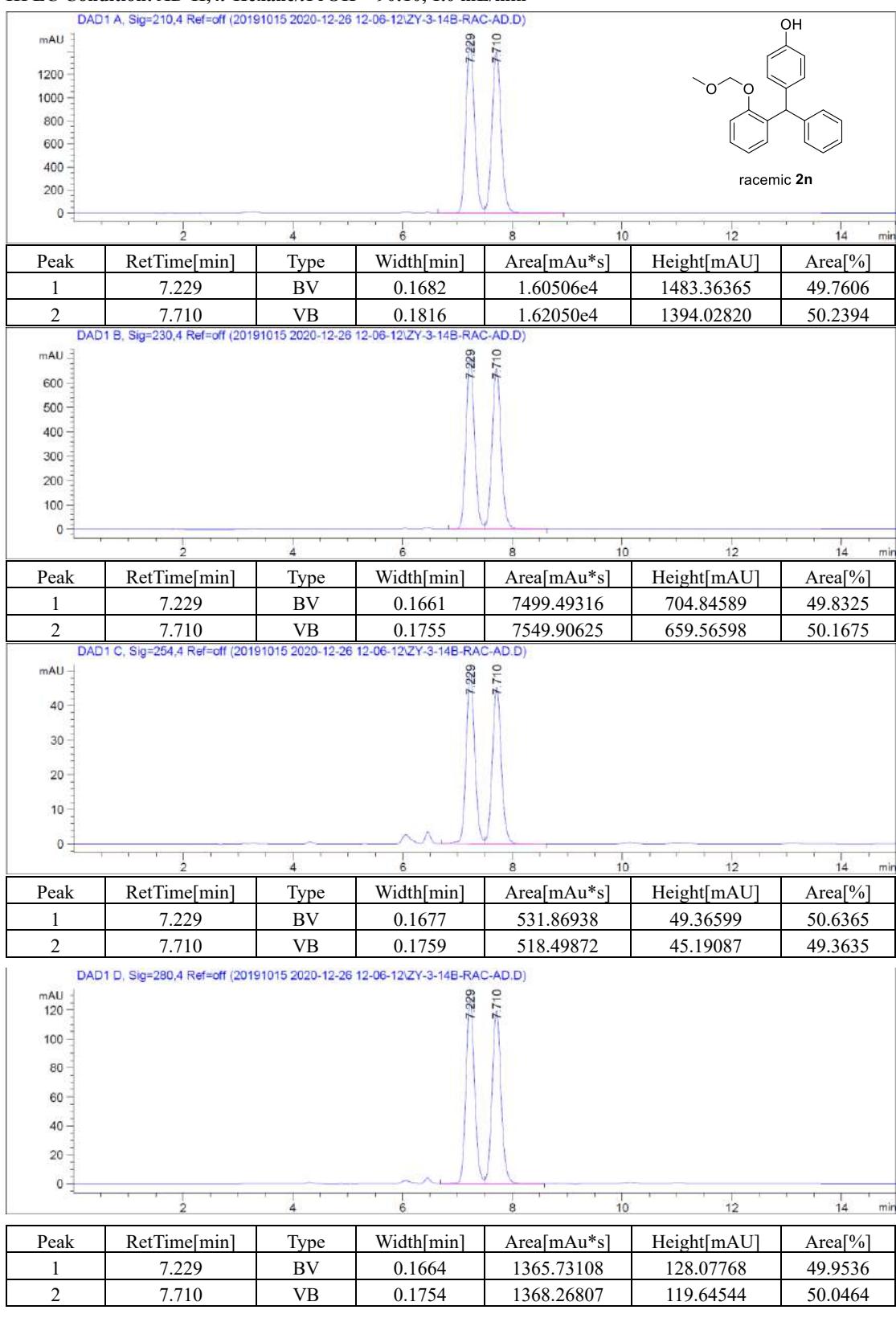
HPLC Condition: AD-H, *n*-Hexane/*i*PrOH = 90:10, 1.0 mL/min



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End of Report

Sample Name: ZY-3-14B-RAC-AD

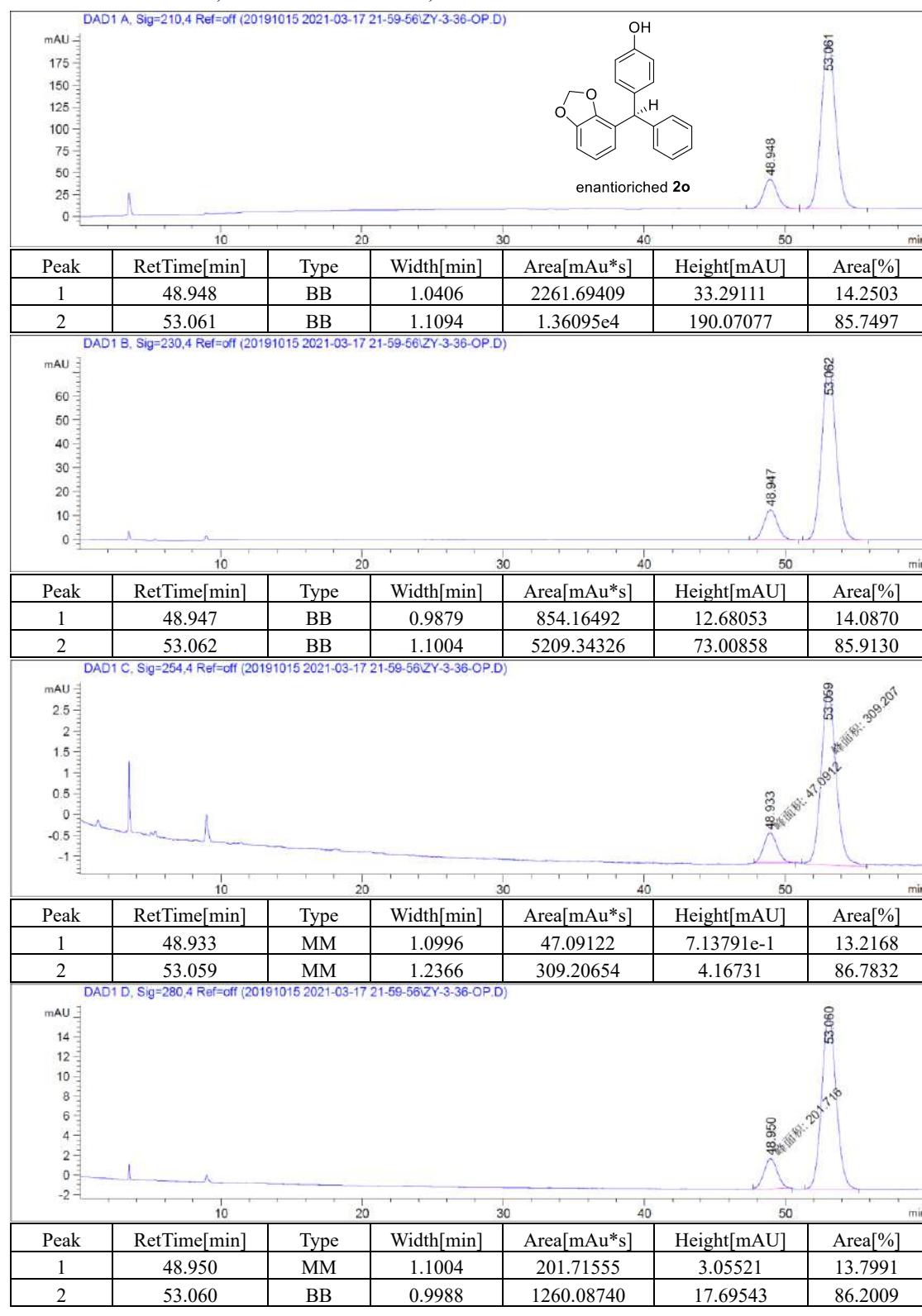
HPLC Condition: AD-H, n-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: ZY-3-36-OP

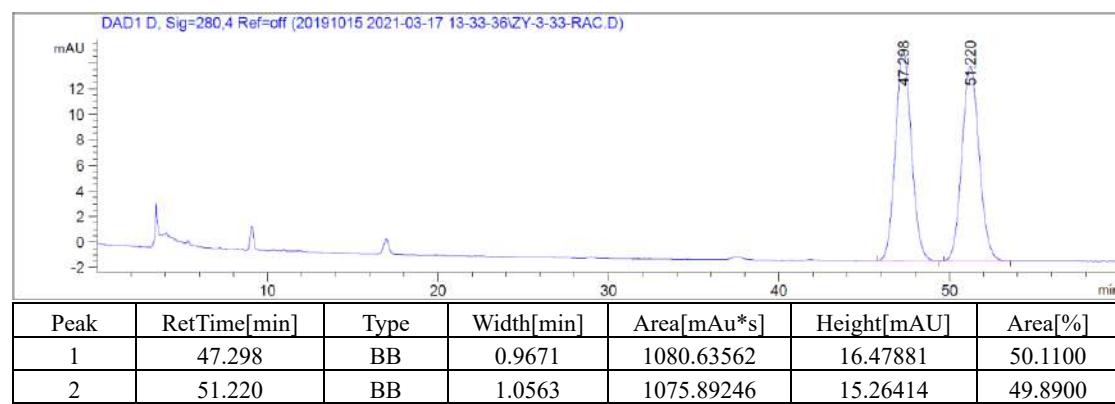
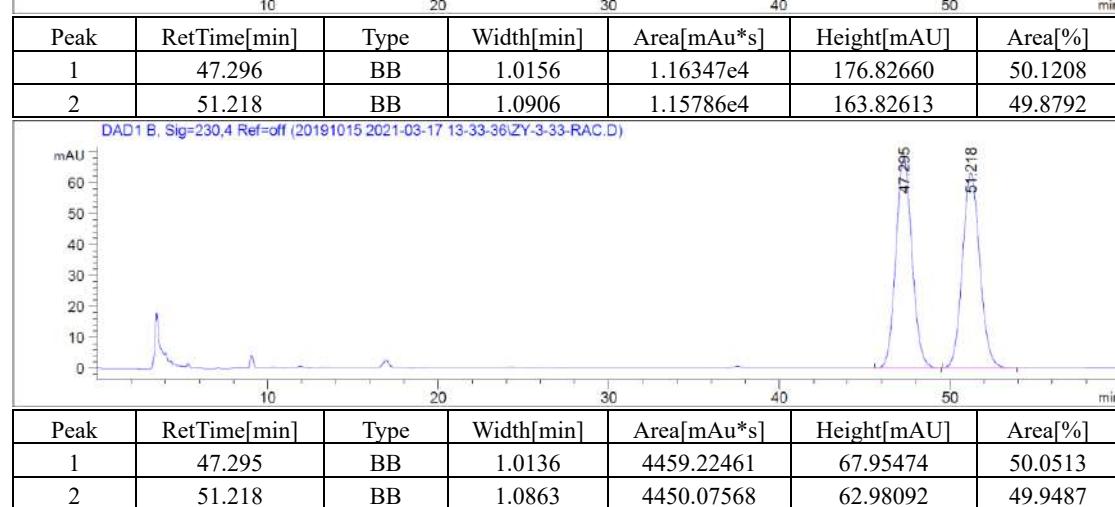
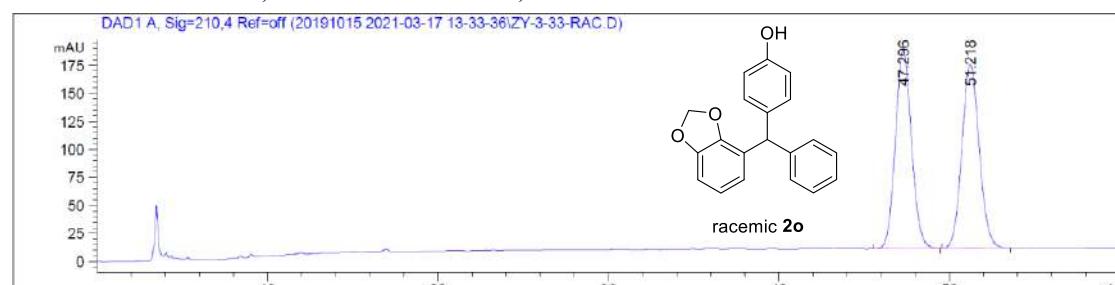
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 97:3, 1.0 mL/min



End of Report

Sample Name: ZY-3-33-RAC

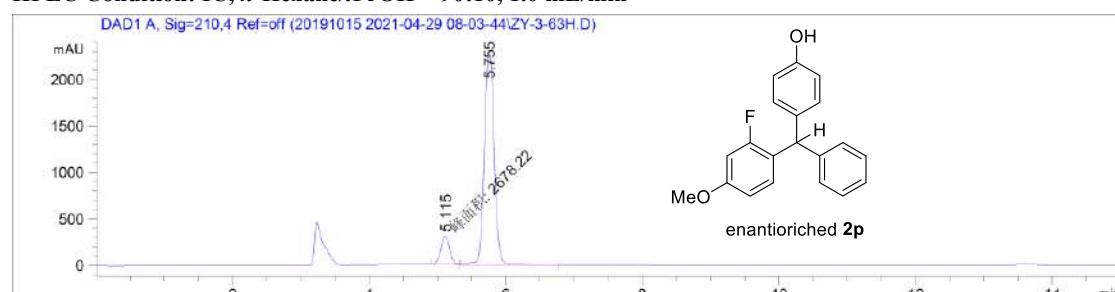
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 97:3, 1.0 mL/min



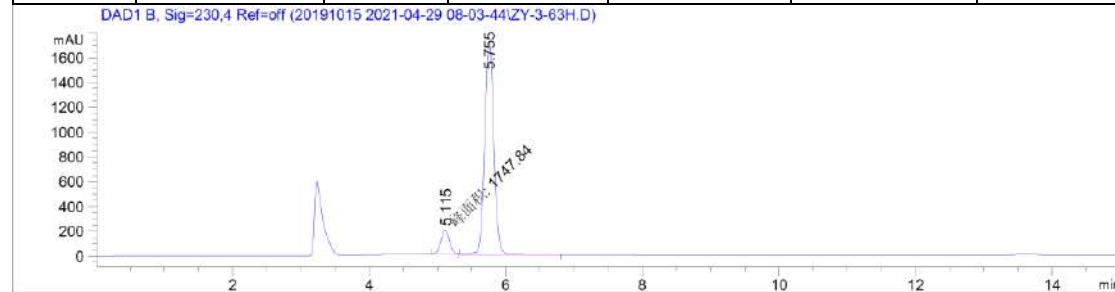
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End of Report

Sample Name: ZY-3-63H

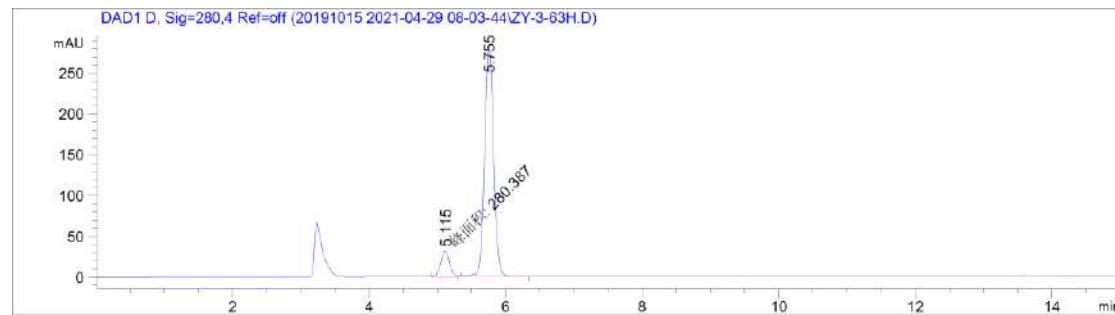
HPLC Condition: IC, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	5.115	MM	0.1464	2678.22046	304.79419	10.2911
2	5.755	VB	0.1607	2.33464e4	2294.65039	89.7089



Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	5.115	MM	0.1479	1747.8446	196.9044	9.6457
2	5.755	VB	0.1471	16372.6396	1718.7126	90.3543

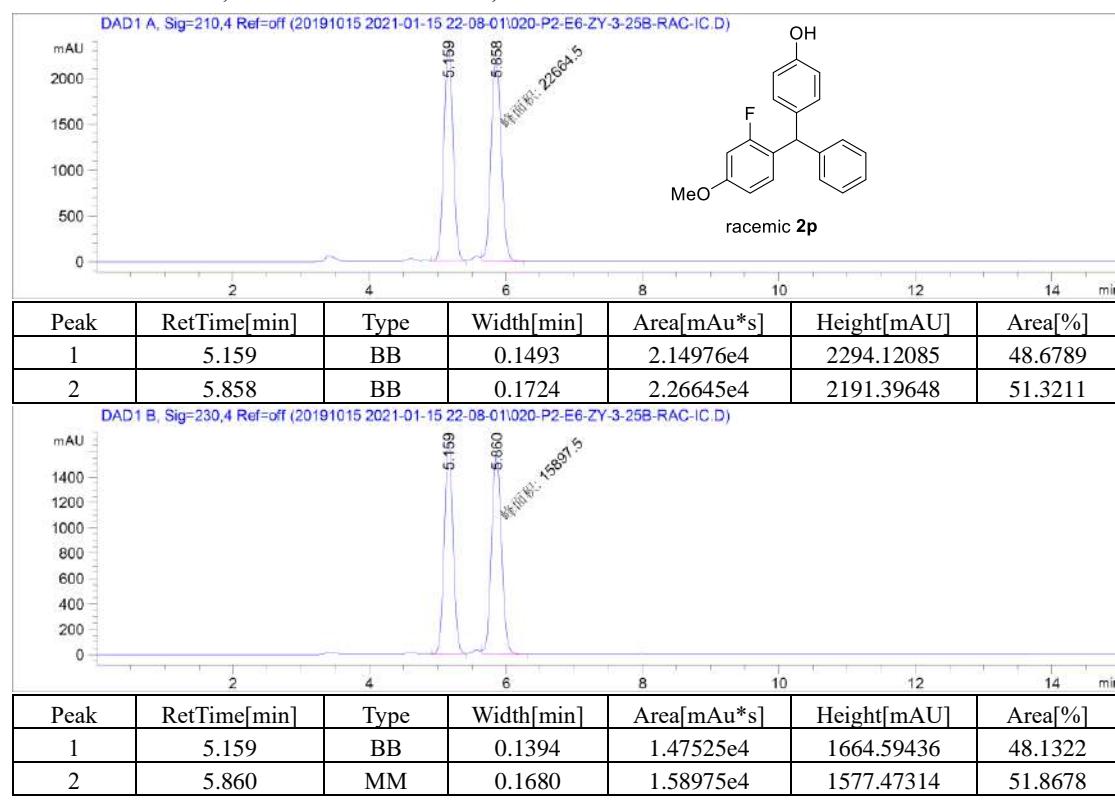


Peak	RetTime[min]	Type	Width[min]	Area[mAu*s]	Height[mAU]	Area[%]
1	5.115	MM	0.1484	280.38669	31.48812	9.7006
2	5.755	BB	0.1439	2610.01416	282.31747	90.2994

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End of Report

Sample Name: ZY-3-25B-RAC-IC

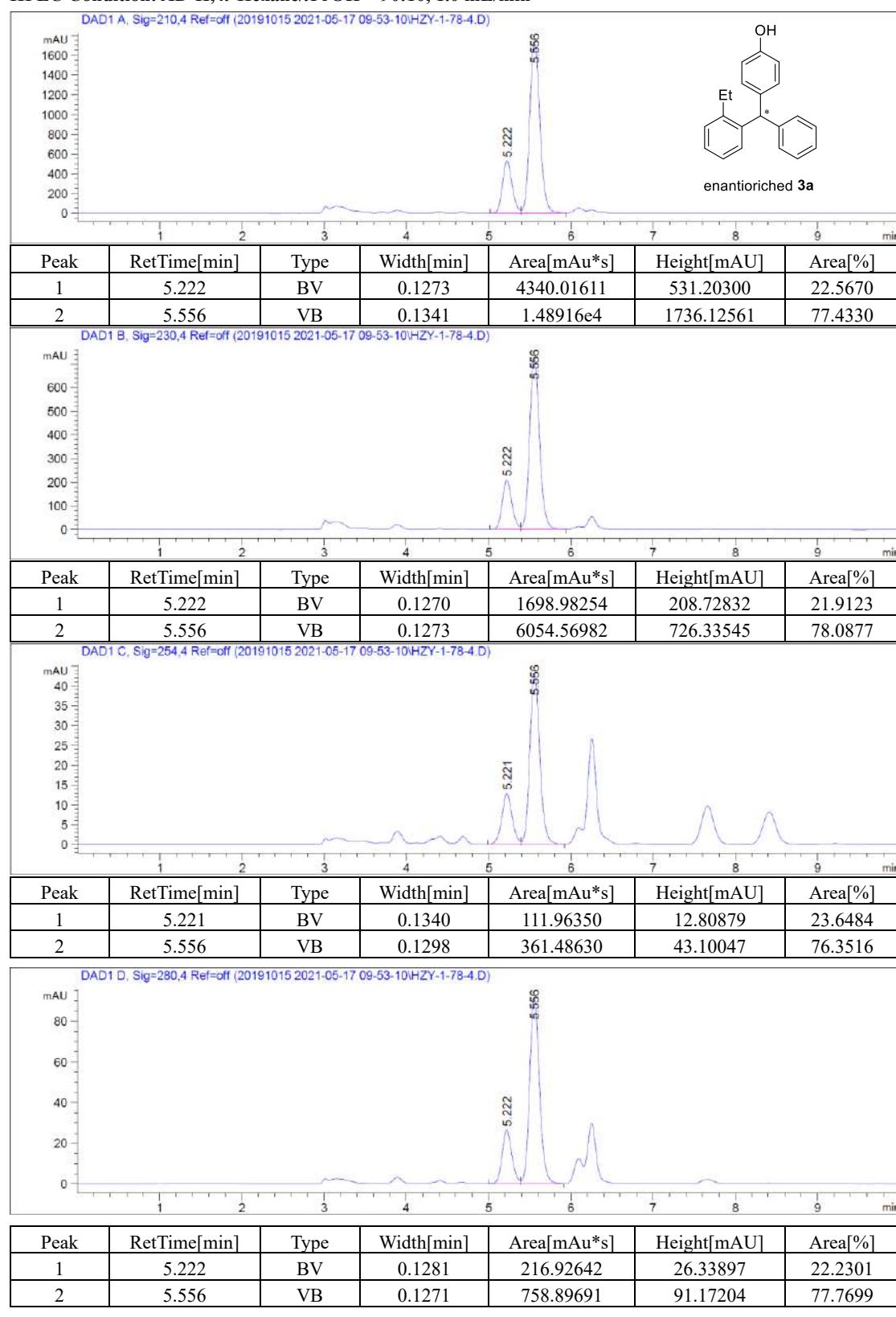
HPLC Condition: IC, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: HZY-78-4

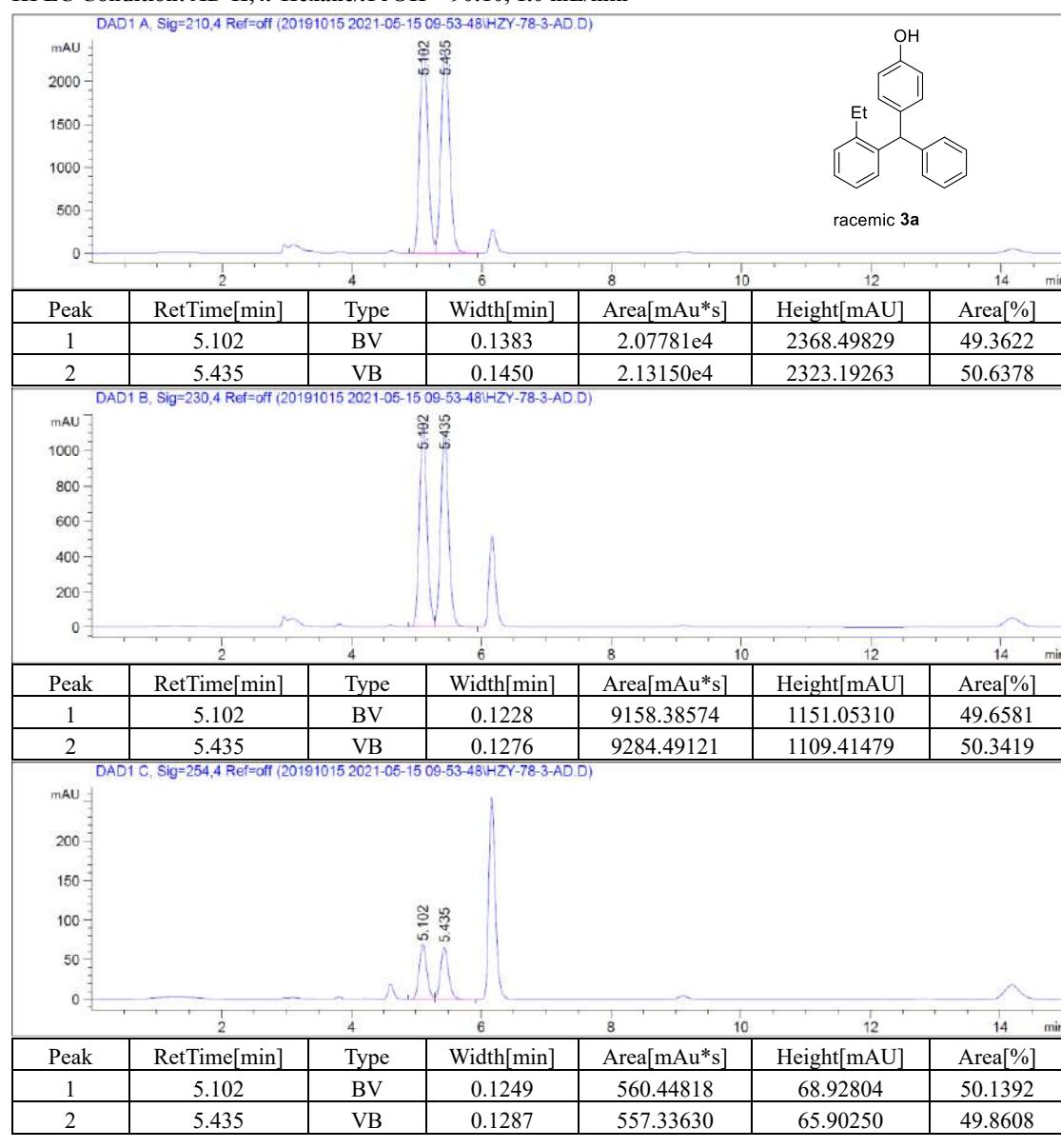
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: HZY-78-3-AD

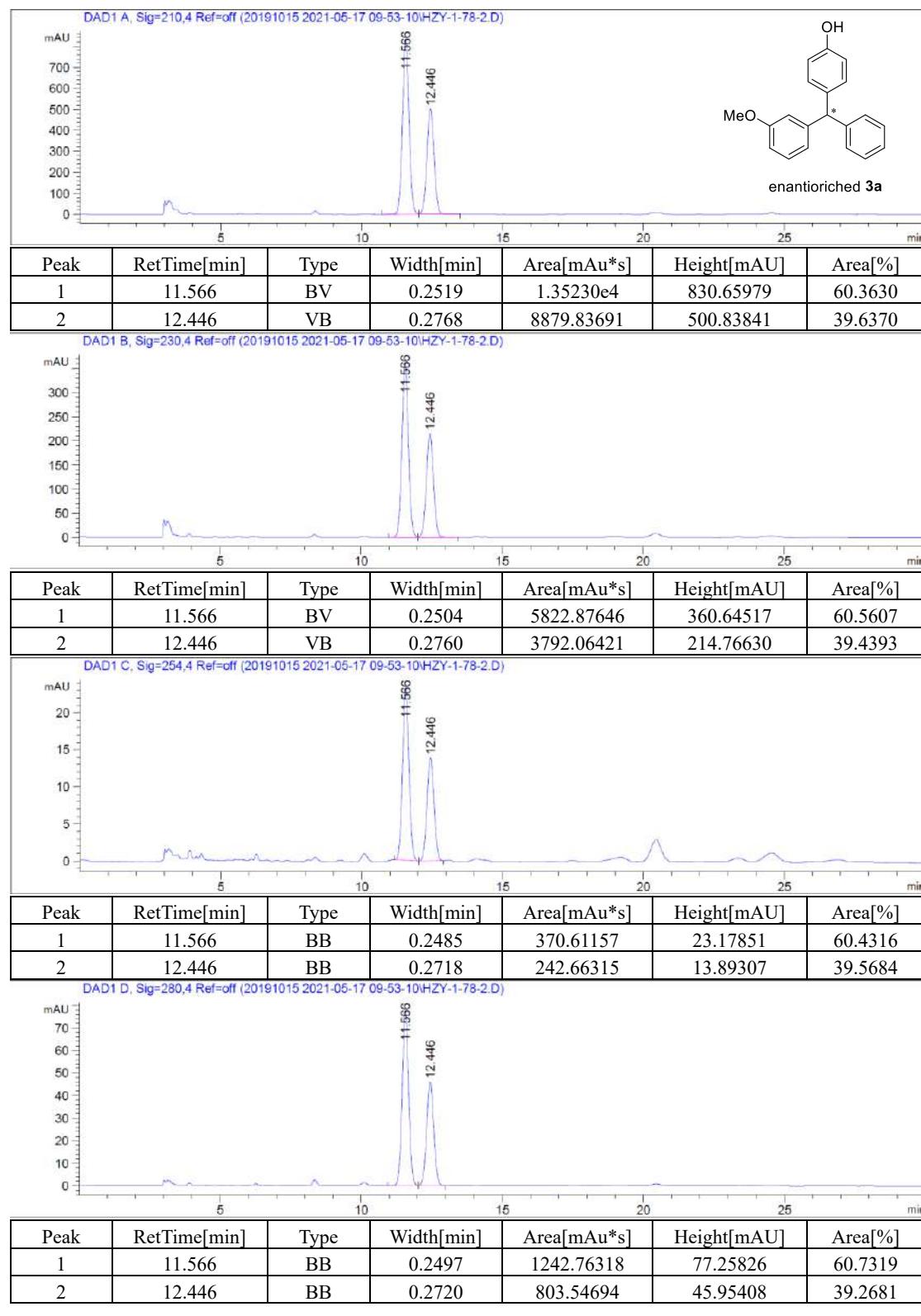
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: HZY-78-2

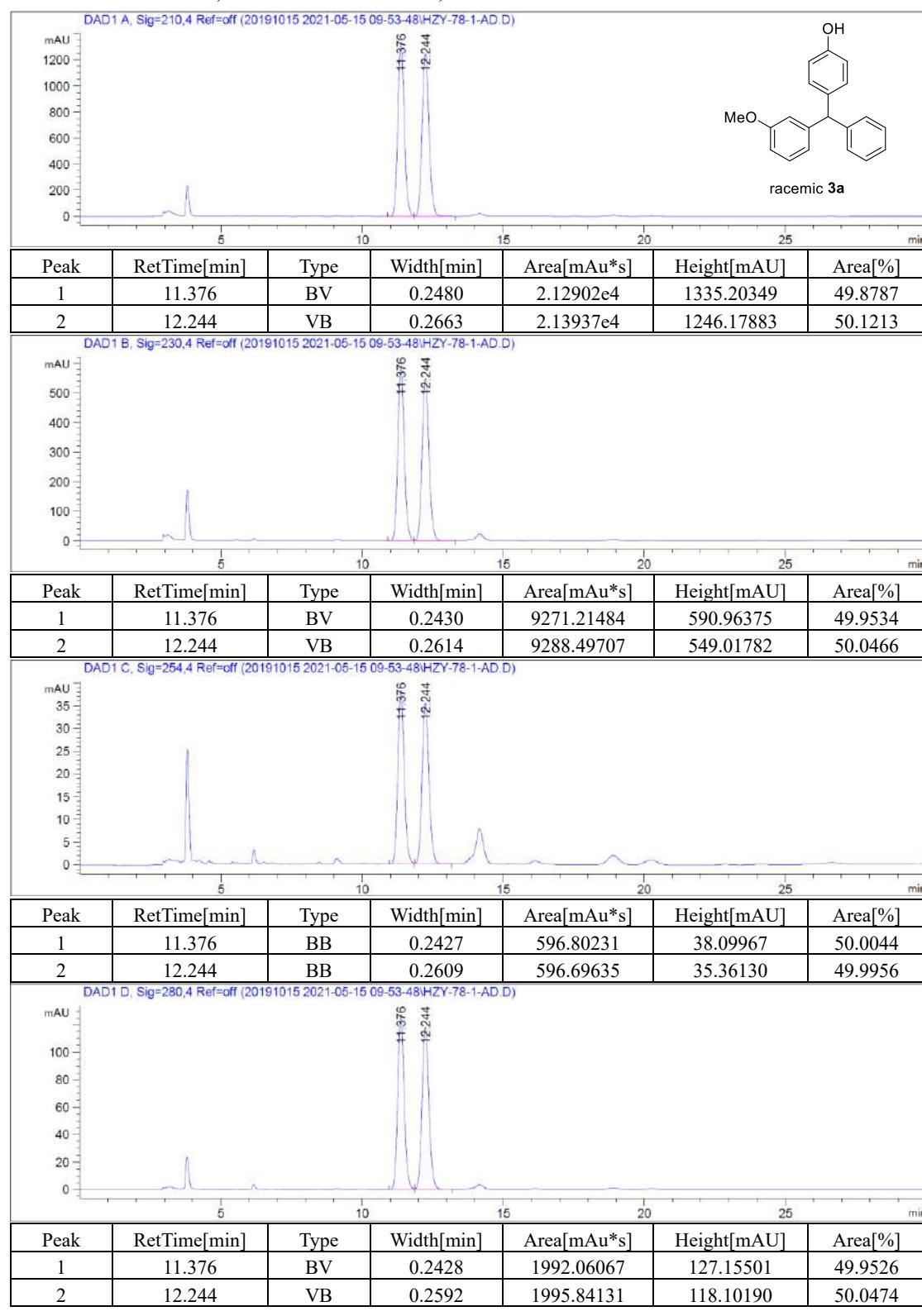
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: HZY-78-1-AD

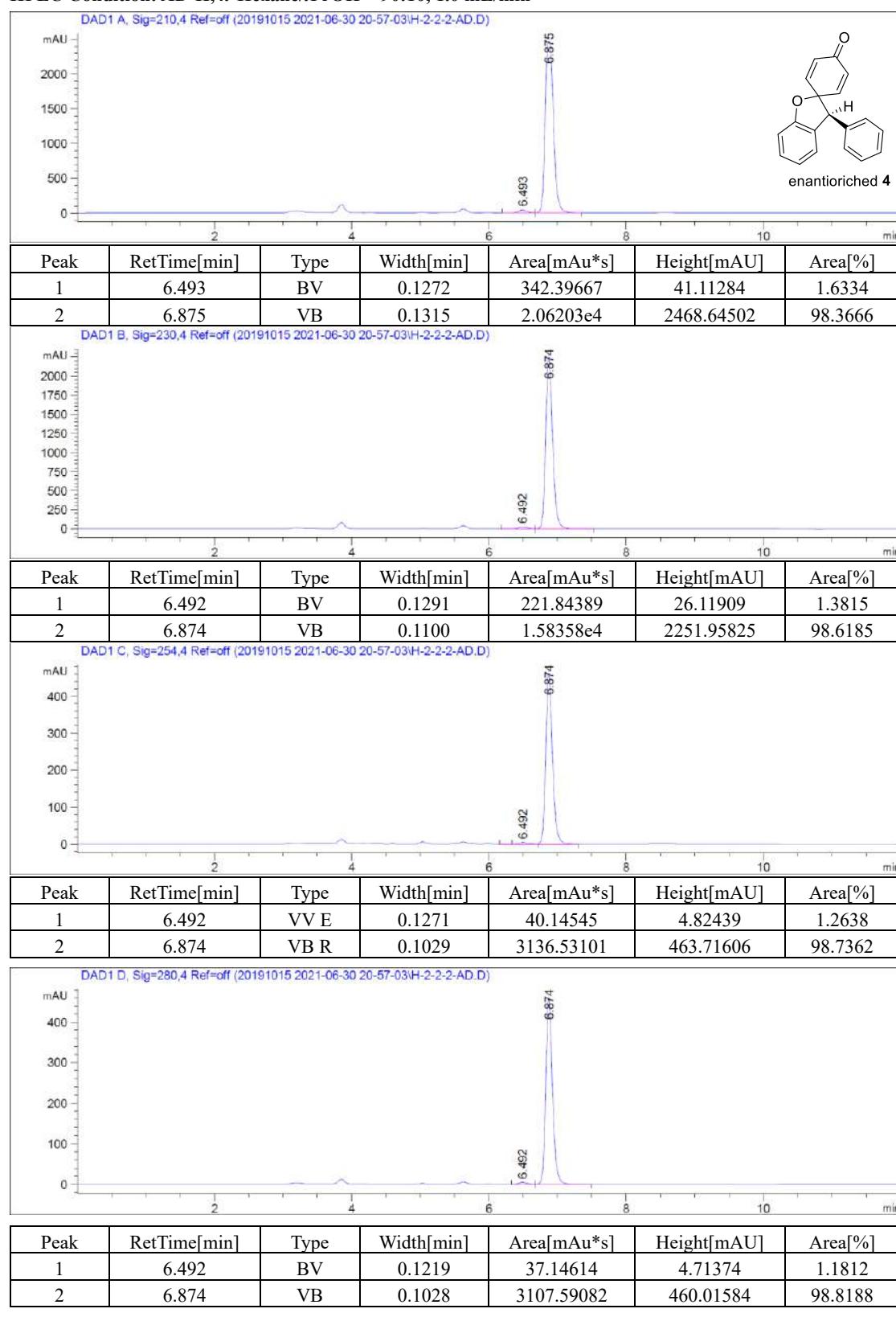
HPLC Condition: AD-H, n-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: H-2-2-2-AD

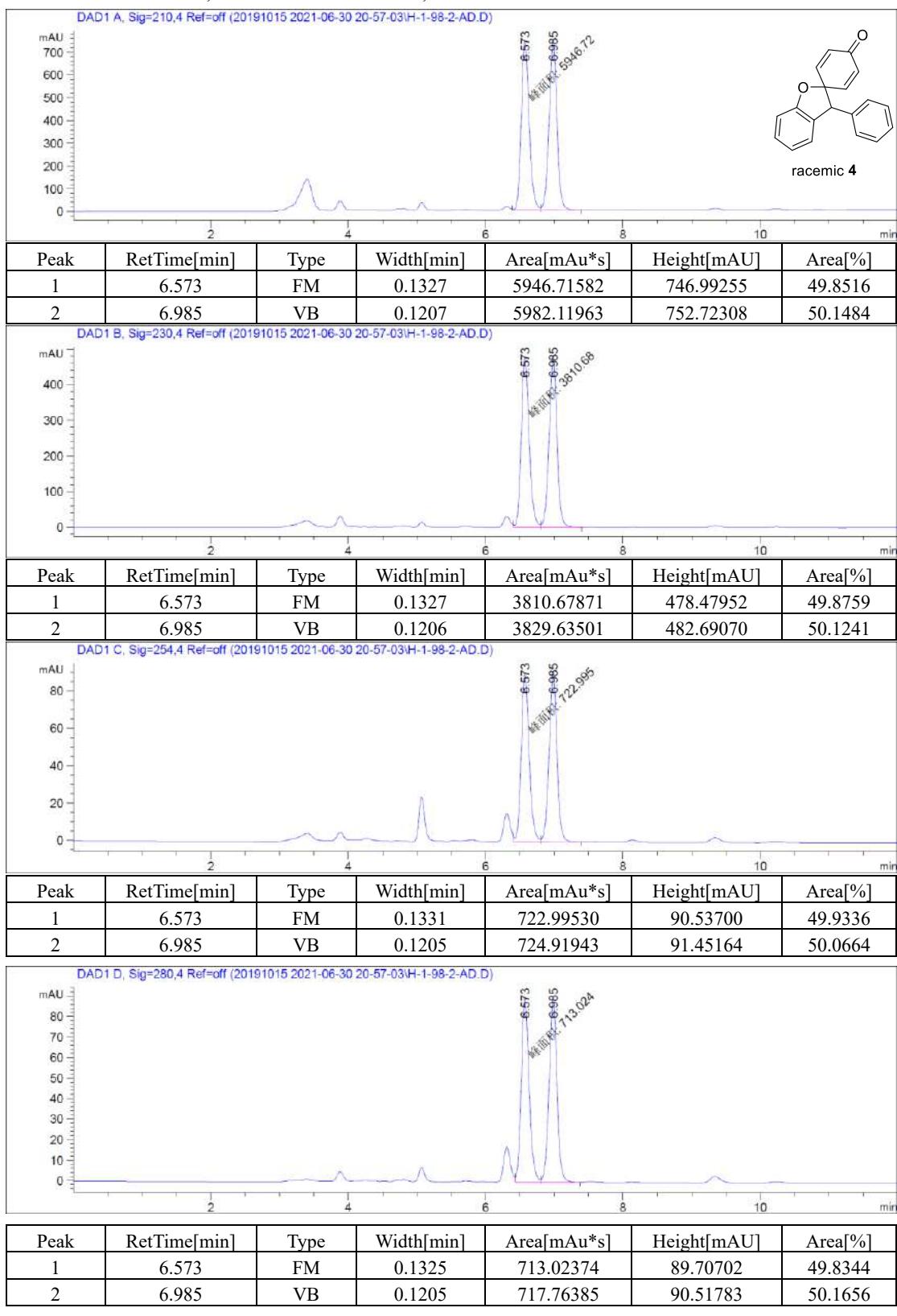
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: H-1-98-2-AD

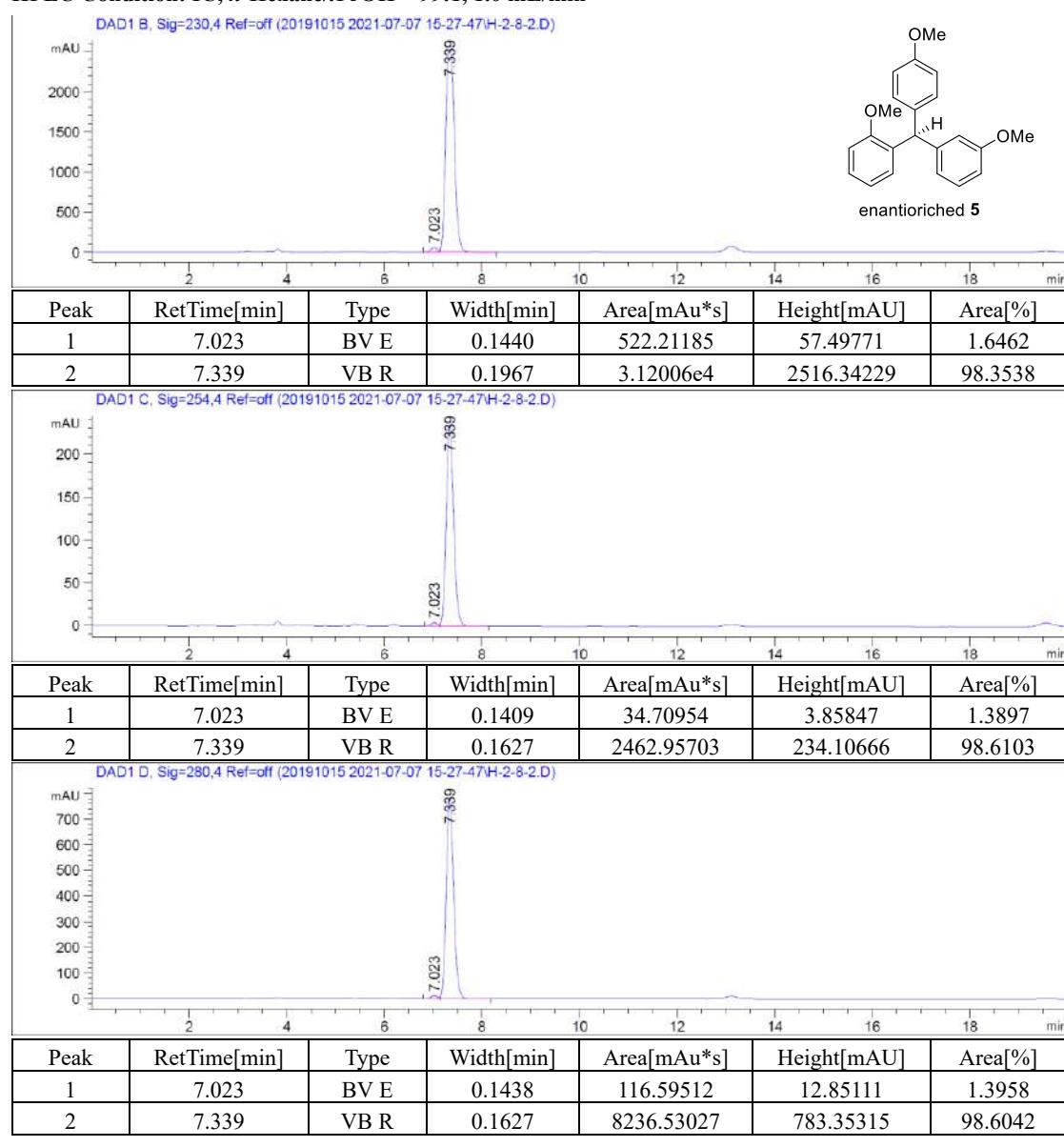
HPLC Condition: AD-H, *n*-Hexane/iPrOH = 90:10, 1.0 mL/min



End of Report

Sample Name: H-2-8-2

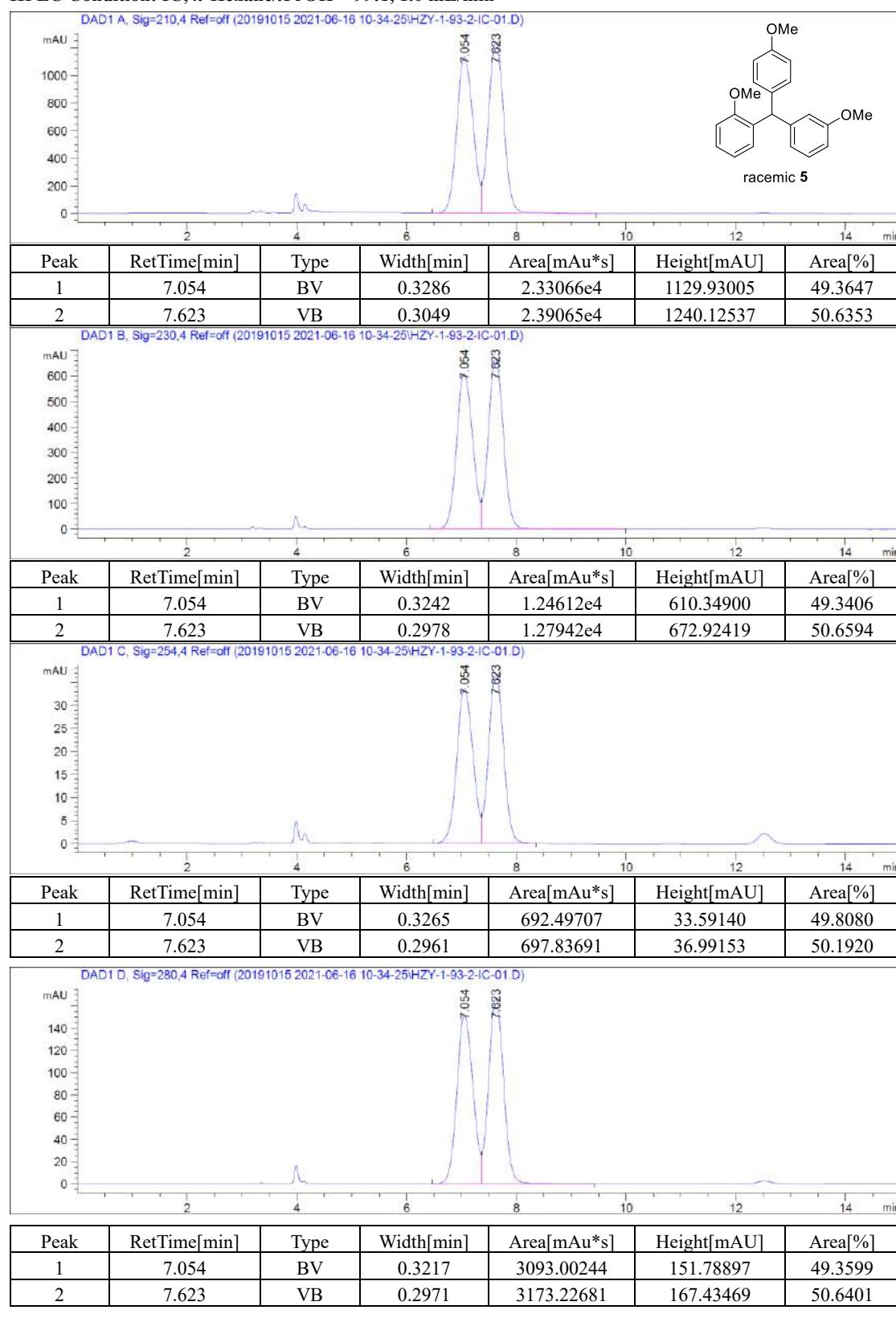
HPLC Condition: IC, *n*-Hexane/iPrOH = 99:1, 1.0 mL/min



End of Report

Sample Name: HZY-1-93-2-IC-01

HPLC Condition: IC, *n*-Hexane/iPrOH = 99:1, 1.0 mL/min



End of Report