Highly stereoselective synthesis of spirocyclopropylthiooxindoles and biological evaluation

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1. General information

Reactions were monitored by thin layer chromatography using UV light or KMnO₄ to visualize the course of reaction. Purification of reaction products was carried out by flash chromatography on silica gel. Chemical yields refer to pure isolated substances. The $[\alpha]_D$ was recorded using Autopol VI High Accuracy Polarimeter. The infrared (IR) spectra were obtained using a Shimadzu Irtracer-100 infrared spectrometer. Chiral HPLC analysis was performed on a Shimadzu LC-20AD instrument using Daicel Chiracel columns at 25 °C and a mixture of HPLC-grade hexane and isopropanol as eluent. ¹H, ¹³C, ¹⁹F NMR spectra were obtained using a Bruker DPX-400 MHz spectrometer. Chemical shifts were reported in ppm from CDCl₃ with the solvent resonance or TMS as the internal standard. The following abbreviations were used to designate chemical shift multiplicities: s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet.

Anhydrous CH₂Cl₂ was prepared by distillation over P₂O₅ and then from CaH₂. Anhydrous toluene, *n*-hexane, Et₂O, THF and MTBE were prepared by distillation over sodium-benzophenone ketyl prior to use. The 3-diazothiooxindole 1^1 and alkenes 2^2 were prepared according to the corresponding literature reports. Chiral Rh₂(*R*-BTPCP)₄ and Rh₂(*R*-DOSP)₄ were purched from Strem. Chiral Rh₂(*S*-PTPA)₄, Rh₂(*S*-PTTL)₄, Rh₂(*S*-TFPTTL)₄ and Rh₂(*S*-TCPTTL)₄ were purched from TCI.

Entry	Chemical name	Abbreviation
1	Petroleum ether	PE
2	Tetrahydrofuran	THF
3	Ethyl acetate	EtOAc
4	Dichloromethane	CH ₂ Cl ₂
5	Methyl tert-butyl ether	MTBE

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2. General procedure for enantioselective cyclopropanation and experimental data.



To a 10 mL Schlenk tube were charged with $Rh_2(S-TCPTTL)_4$ (3.0 mg, 0.0015 mmol, 0.5 mol %), Et₂O (6.0 mL) was added followed by the alkenes **2** (3.0 mol, 10.0 equivs), after the reaction was cooled to 0 °C and stirred for 0.5 hour, 3-diazothiooxindole **1** (0.3 mmol) was added directly. The resulting mixture was stirred at 0 °C till full conversion of **1** by TLC analysis. Then, the reaction mixture was rapidly passed through a short pad of silica gel, and washed with Et₂O. The resulting organic solution was concentrated in vacuo to give the crude product. To determine the diastereoselectivity, the residue was first dissolved in CDCl₃ or acetone-d₆, and took some samples for ¹H NMR analysis of crude reaction mixture. Then the sample for analysis and rest crude product were recombined for column chromatography purification to afford products **3** or **4**, using PE/EtOAc (60/1, v/v) as the eluent.

Product **3a** was obtained in 92% yield, with a dr ratio of 10:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 119-121 °C); HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 9.34 min, t_r (major) = 7.07 min) gave the isomeric composition of the major isomer: 97% ee; $[\alpha]_D^{25} = 13.3$ (c = 0.18, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.29-7.21 (m, 4H), 7.15-7.11 (m, 2H), 6.95-6.92 (m, 1H), 5.71-5.70 (m, 1H), 3.42 (t, J = 8.8 Hz, 1H), 2.38 (dd, J =9.3, 4.6 Hz, 1H), 2.03 (dd, J = 8.4, 4.6 Hz, 1H), 1.96 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 203.78, 134.98, 134.95, 134.22, 131.32, 130.46, 128.47, 127.82, 127.79, 122.83, 122.45, 43.17, 42.09, 26.79, 21.15; IR (ATR): 3024, 1684, 1572, 1497, 1196, 986, 806 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₄NaOS [M+Na]⁺: 289.0658, Found: 289.0653. Product **3b** was obtained in 92% yield, with a dr ratio of 7:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 137-139 °C); HPLC analysis (Chiralcel OJ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 12.04 min, t_r (major) = 10.24 min) gave the isomeric

composition of the major isomer: 96% ee; $[\alpha]_D^{25} = 13.3$ (c = 0.24, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.25-7.23 (m, 1H), 7.11-7.09 (m, 2H), 7.04-7.02 (m, 2H), 6.97-6.94 (m, 1H), 5.79-5.78 (m, 1H), 3.41 (t, J = 8.8 Hz, 1H), 2.39 (dd, J = 9.3, 4.6 Hz, 1H), 2.33 (s, 3H), 2.05-2.02 (m, 1H), 2.01 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 203.78, 137.49, 135.08, 134.92, 131.31, 131.08, 130.27, 129.12, 127.75, 122.84, 122.40, 43.32, 42.14, 26.86, 21.23, 21.20; IR (ATR): 3024, 2920, 1684, 1516, 1190, 988, 770 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₆NaOS [M+Na]⁺: 303.0814, Found: 303.0805.

Product **3c** was obtained in 88% yield, with a dr ratio of 6:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 116-118 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 7.06 min, t_r (major) = 10.08 min) gave the isomeric

composition of the major isomer: 90% ee; $[\alpha]_D^{25} = -12.5$ (c = 0.16, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.33-7.30 (m, 2H), 7.25-7.23 (m, 1H), 7.09-7.06 (m, 2H), 6.96-6.94 (m, 1H), 5.65-5.64 (m, 1H), 3.44 (t, J = 8.8 Hz, 1H), 2.40 (dd, J = 9.3, 4.5 Hz, 1H), 2.04 (dd, J = 8.3, 4.6 Hz, 1H), 1.96 (s, 3H), 1.31 (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ 203.85, 150.96, 135.14, 134.84, 131.32, 131.29, 130.28, 127.68, 125.31, 122.90, 122.35, 43.15, 41.83, 34.68, 31.40, 27.12, 21.05; IR (ATR): 2963, 2918, 1694, 1364, 1194, 989, 772 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₁H₂₂NaOS [M+Na]⁺: 345.1284, Found: 345.1283.



Product **3d** was obtained in 94% yield, with a dr ratio of 13:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 133-135 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 15/85, 1.0 mL/min, 230 nm; t_r (minor) = 6.32 min, t_r (major) = 10.06 min) gave

the isomeric composition of the major isomer: 95% ee; $[\alpha]_D^{25} = 41.8$ (c = 0.24, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.32-7.30 (m, 2H), 7.25-7.23 (m, 1H), 7.14-7.12 (m, 2H), 6.97-6.94 (m, 1H), 5.69-5.68 (m, 1H), 4.55 (s, 2H), 3.41 (t, J = 8.8 Hz, 1H), 2.40 (dd, J = 9.3, 4.7 Hz, 1H), 2.03 (dd, J = 3.4

8.4, 4.7 Hz, 1H), 1.99 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 203.66, 137.20, 135.13, 134.70, 134.67, 131.34, 130.90, 128.76, 127.97, 122.81, 122.53, 45.91, 43.10, 41.53, 26.66, 21.16; IR (ATR): 2916, 2849, 1682, 1464, 1368, 1196, 993, 814 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₅ClNaOS [M+Na]⁺: 337.0424, Found: 337.0424.

Product **3e** was obtained in 90% yield, with a dr ratio of 7:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 106-108 °C); HPLC analysis (Chiralcel OZ-H, ¹PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 9.73 min, t_r (major) = 13.04 min) gave the isomeric composition of the major isomer: 96% ee; $[\alpha]p^{25} = 25.0$ (c = 0.24, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.25-7.23 (m, 1H), 7.12-7.08 (m, 2H), 6.99-6.95 (m, 3H), 5.71-5.70 (m, 1H), 3.35 (t, J =8.8 Hz, 1H), 2.39 (dd, J = 9.4, 4.7 Hz, 1H), 2.01 (s, 3H), 1.98 (dd, J = 8.3, 4.7 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 203.59, 162.29 (d, J = 246.9 Hz), 135.12, 134.69, 132.11 (d, J = 8.2 Hz), 131.43, 130.15 (d, J = 3.3 Hz), 128.00, 122.79, 122.60, 115.55 (d, J = 21.6 Hz), 43.13, 41.15, 26.82, 21.22; ¹⁹F NMR (376 MHz, CDCl₃): δ -113.94; IR (ATR): 2918, 1690, 1605, 1512, 1464, 1231, 839, 772 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₃FNaOS [M+Na]⁺: 307.0563, Found: 307.0559.



Product **3f** was obtained in 85% yield, with a dr ratio of 9:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 139-141 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 3/97, 1.0 mL/min, 254 nm; t_r (minor) = 7.07 min, t_r (major) = 9.76 min) gave the isomeric

composition of the major isomer: 97% ee; $[\alpha]_D^{25} = 57.6$ (c = 0.25, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.26-7.23 (m, 3H), 7.08-7.06 (m, 2H), 6.98-6.96 (m, 1H), 5.73-5.72 (m, 1H), 3.34 (t, J = 8.8 Hz, 1H), 2.39 (dd, J = 9.3, 4.7 Hz, 1H), 2.02 (s, 3H), 1.98 (dd, J = 8.4, 4.7 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 203.54, 135.21, 134.55, 133.73, 132.88, 131.79, 131.43, 128.69, 128.10, 122.76, 122.66, 43.12, 41.15, 26.53, 21.27; IR (ATR): 2916, 2849, 1692, 1495, 1190, 835, 770 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₃ClNaOS [M+Na]⁺: 323.0268, Found: 323.0265.



Product **3g** was obtained in 88% yield, with a dr ratio of 13:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 146-148 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0

mL/min, 254 nm; t_r (minor) = 11.59 min, t_r (major) = 17.73 min) gave the isomeric composition of the major isomer: 97% ee; $[\alpha]_D^{25} = 22.9$ (c = 0.14, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.42-7.39 (m, 2H), 7.26-7.23 (m, 1H), 7.02-6.96 (m, 3H), 5.73-5.72 (m, 1H), 3.32 (t, J = 8.8 Hz, 1H), 2.39 (dd, J = 9.3, 4.7 Hz, 1H), 2.02 (s, 3H), 1.98 (dd, J = 8.4, 4.7 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 203.50, 135.21, 134.50, 133.39, 132.12, 131.64, 131.42, 128.11, 122.74, 122.66, 121.83, 43.07, 41.18, 26.44, 21.27; IR (ATR): 2916, 2849, 1690, 1489, 1192, 991, 808 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₃BrNaOS [M+Na]⁺: 366.9763, Found: 366.9753.

Product **3h** was obtained in 83% yield, with a dr ratio of 14:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 91-93 °C); HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 10.03 min, t_r (major) = 8.88 min) gave the isomeric composition of the major isomer: 98% ee; $[\alpha]p^{25} = 19.1$ (c = 0.23, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.40-7.38 (m, 1H), 7.34-7.33 (m, 1H), 7.26-7.24 (m, 1H), 7.15-7.11 (m, 1H), 7.04-6.96 (m, 2H), 5.77-5.76 (m, 1H), 3.35 (t, J = 8.8 Hz, 1H), 2.37 (dd, J = 9.3, 4.8 Hz, 1H), 2.03 (s, 3H), 1.99 (dd, J =8.3, 4.8 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 203.47, 136.66, 135.21, 134.39, 133.25, 131.41, 130.95, 129.98, 129.28, 128.16, 122.82, 122.68, 122.47, 43.07, 41.01, 26.32, 21.24; IR (ATR): 2918, 2849, 1690, 1468, 1194, 993, 768 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₃BrNaOS [M+Na]⁺: 366.9763, Found: 366.9758.



Product **3i** was obtained in 78% yield, with a dr ratio of 6:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 151-153 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 11.89 min, t_r (major) = 13.34 min) gave the isomeric

composition of the major isomer: 97% ee; $[\alpha]_D^{25} = -49.5$ (c = 0.21, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.46-7.35 (m, 3H), 7.24-7.22 (m, 1H), 7.18-7.14 (m, 1H), 6.98-6.95 (m, 1H), 5.67-5.66 (m, 1H), 3.24 (t, J = 8.8 Hz, 1H), 2.45 (dd, J = 9.1, 4.9 Hz, 1H), 2.05 (dd, J = 8.4, 4.9 Hz, 1H), 1.97 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 203.79, 134.92, 134.55, 134.54, 132.83, 131.95, 131.18, 129.44, 128.02, 127.18, 122.53, 121.60, 43.20, 43.19, 26.30, 21.20; IR (ATR): 2920, 2855, 1680, 1466, 1192, 991, 752 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₃BrNaOS [M+Na]⁺: 366.9763, Found: 366.9753.

S S S Product **3j** was obtained in 98% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 160-162 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 3/97, 1.0 mL/min, 254 nm; tr (minor) = 7.47 min, tr (major) = 9.63 min) gave the

isomeric composition of the major isomer: 93% ee; $[\alpha]_D^{25} = 48.4$ (c = 0.19, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.82-7.79 (m, 2H), 7.73-7.70 (m, 2H), 7.52-7.47 (m, 2H), 7.25-7.23 (m, 1H), 7.16-7.13 (m, 1H), 6.92-6.90 (m, 1H), 5.79-5.78 (m, 1H), 3.58 (t, J = 8.8 Hz, 1H), 2.49 (dd, J = 9.3, 4.6 Hz, 1H), 2.19 (dd, J = 8.4, 4.6 Hz, 1H), 1.81 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 203.78, 135.08, 134.86, 133.21, 132.87, 131.83, 131.36, 129.13, 128.27, 128.21, 127.89, 127.83, 127.80, 126.41, 126.29, 122.71, 122.53, 43.41, 42.47, 26.81, 21.12; IR (ATR): 2918, 2849, 1690, 1603, 1194, 773, 748 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₁H₁₆NaOS [M+Na]⁺: 339.0814, Found: 339.0817.

Product **3k** was obtained in 86% yield, with a dr ratio of 5:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a yellowish solid (m.p. 115-117 °C); HPLC analysis (Chiralcel OD-H, ¹PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 9.59 min, t_r (major) = 7.47 min) gave the isomeric composition of the major isomer: 90% ee; $[\alpha]p^{25} = -59.1$ (c = 0.21, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.27-7.25 (m, 1H), 7.21-7.20 (m, 1H), 7.02-7.00 (m, 1H), 6.97-6.94 (m, 1H), 6.90-6.89 (m, 1H), 6.03-6.02 (m, 1H), 3.36 (t, J = 8.6 Hz, 1H), 2.46 (dd, J = 9.2, 4.7 Hz, 1H), 2.09-2.06 (m, 1H), 2.08 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 203.18, 137.86, 135.27, 134.40, 131.35, 128.62, 128.20, 126.83, 126.10, 122.57, 122.30, 43.94, 36.14, 27.70, 21.29; IR (ATR): 3103, 2922, 2855, 1684, 1572, 1470, 1194, 978, 698 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₅H₁₂NaOS₂ [M+Na]⁺: 295.0222, Found: 295.0214.



Product **31** was obtained in 99% yield, with a dr ratio of 16:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 89-91 °C); HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 8.78 min, t_r (major) = 6.64 min) gave the isomeric composition

of the major isomer: 96% ee; $[\alpha]_D^{25} = 55.7$ (c = 0.23, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.29-7.21 (m, 3H), 7.16-7.11 (m, 2H), 6.78-6.77 (m, 1H), 5.56-5.55 (m, 1H), 3.42 (t, J = 8.9 Hz, 1H),

2.38 (dd, J = 9.4, 4.5 Hz, 1H), 2.28 (s, 3H), 2.01 (dd, J = 8.4, 4.6 Hz, 1H), 1.94 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): § 203.86, 134.89, 134.35, 131.58, 131.16, 130.48, 128.91, 128.44, 127.74, 120.18, 43.69, 42.13, 26.93, 21.05, 20.61; IR (ATR): 2911, 2857, 1678, 1584, 1375, 1103, 743 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₆NaOS [M+Na]⁺: 303.0814, Found: 303.0818.

Product **3m** was obtained in 96% yield, with a dr ratio of 16:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a colorless oil; HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 14.89 min, t_r (major) = 11.96 min) gave the isomeric composition of the major isomer: 82% ee; $[\alpha]_D^{25} = 65.2$ (c = 0.27, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.29-7.24 (m, 3H), 7.14-7.12 (m, 2H), 6.93-6.92 (m, 1H), 6.36-6.34 (m, 1H), 5.83-5.81 (m, 1H), 3.71 (s, 3H), 3.37 (t, J = 8.9 Hz, 1H), 2.35 (dd, J = 9.4, 4.7 Hz, 1H), 1.98 (dd, J = 8.3, 4.7 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 203.65, 158.90, 135.80, 134.36, 130.41, 128.57, 127.80, 126.90, 122.57, 111.49, 108.70, 55.50, 42.76, 41.43, 26.38; IR (ATR): 2999, 2835, 1686, 1601, 1489, 1032, 980, 696 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₄NaO₂S [M+Na]⁺: 305.0607, Found: 305.0599.



Product **3n** was obtained in 93% yield, with a dr ratio of 15:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 92-94 °C); HPLC analysis (Chiralcel OD-H, ⁱPrOH/hexane = 1/99, 1.0 mL/min, 230

nm; t_r (minor) = 11.51 min, t_r (major) = 7.17 min) gave the isomeric composition of the major isomer: 84% ee; $[\alpha]_D^{25} = 95.5$ (c = 0.31, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 8.27-8.24 (m, 3H), 8.14-8.12 (m, 2H), 7.96-7.94 (m, 1H), 7.74-7.70 (m, 1H), 6.78-6.76 (m, 1H), 4.44 (t, J = 8.9 Hz, 1H), 3.40 (dd, J = 9.3, 4.6 Hz, 1H), 3.32 (s, 3H), 3.04 (dd, J = 8.4, 4.6 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 203.40, 134.98, 134.70, 134.25, 132.00, 130.43, 128.55, 127.99, 127.84, 125.17, 119.19, 43.82, 42.29, 27.02, 20.76; IR (ATR): 2918, 1690, 1587, 1368, 1103, 986, 741 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₇H₁₄NaOS [M+Na]⁺: 289.0658, Found: 289.0648.

Product **30** was obtained in 93% yield, with a dr ratio of 12:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 151-153 °C); HPLC analysis (Chiralcel OD-H, ⁱPrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 12.57 min, t_r (major) = 8.85 min) gave the isomeric composition of the

major isomer: 88% ee; $[\alpha]_D^{25} = -30.0$ (c = 0.61, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.36-7.26 (m, 4H), 7.15-7.10 (m, 3H), 5.89-5.88 (m, 1H), 3.49 (t, J = 9.0 Hz, 1H), 2.45 (dd, J = 9.4, 4.8 Hz, 1H), 2.09 (dd, J = 8.5, 4.8 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 202.38, 136.86, 133.49, 133.19, 131.37, 130.33, 128.80, 128.30, 127.20, 123.68, 122.20, 43.38, 42.80, 27.27; IR (ATR): 2924, 1697, 1560, 1500, 1269, 986, 814 cm⁻¹; HRMS (EI): Exact mass calcd for C₁₆H₁₁OSCl [M]⁺: 286.0219, Found: 286.0218.

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Product **3p** was obtained in 82% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 149-151 °C); HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 10.31 min, t_r (major) = 8.86 min) gave the isomeric

composition of the major isomer: 85% ee; $[\alpha]_D^{25} = 77.2$ (c = 0.29, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.35-7.30 (m, 3H), 7.26-7.20 (m, 2H), 6.97-6.94 (m, 1H), 5.47-5.46 (m, 1H), 3.64 (d, J = 6.8 Hz, 1H), 3.57 (dd, J = 18.8, 6.5 Hz, 1H), 3.25 (d, J = 18.9 Hz, 1H), 3.00 (t, J = 6.7 Hz, 1H), 1.95 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.70, 145.76, 138.64, 135.24, 133.80, 132.36, 128.10, 127.62, 127.44, 126.83, 124.01, 123.14, 122.83, 47.55, 45.29, 40.91, 32.56, 21.42; IR (ATR): 2916, 2849, 1684, 1460, 1088, 808, 760 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₄NaOS [M+Na]⁺: 301.0658, Found: 301.0659.

Product **4a** was obtained in 45% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a colorless oil; HPLC analysis (Chiralpak AD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; tr (minor) = 14.94 min, tr (major) = 10.72 min) gave the isomeric composition of the major isomer: 99% ee; $[\alpha]_D^{25} = -21.7$ (c = 0.24, Acetone); ¹H NMR (400 MHz, Acetone- d_6): δ 7.44-7.42 (m, 1H), 7.29-7.26 (m, 4H), 7.22-7.17 (m, 3H), 2.55 (d, J = 5.3 Hz, 1H), 2.39 (s, 3H), 2.17 (d, J = 5.3 Hz, 1H), 1.58 (s, 3H); ¹³C NMR (100 MHz, Acetone- d_6): δ 199.04, 142.79, 136.01, 135.91, 132.91, 129.59, 128.95, 128.88, 127.68, 125.14, 123.79, 49.79, 47.46, 28.40, 22.97, 21.39; IR (ATR): 2916, 2849, 1462, 1445, 1152, 959, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₆NaOS [M+Na]⁺: 303.0814, Found: 303.0813. Product 4b was obtained in 58% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid
(m.p. 100-102 °C); HPLC analysis (Chiralcel OZ-H, ⁱPrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 12.58 min, t_r (major) = 16.48 min) gave the isomeric

composition of the major isomer: 98% ee; $[\alpha]_D^{25} = -331.4$ (c = 0.21, Acetone); ¹H NMR (400 MHz, Acetone- d_6): δ 7.43-7.41 (m, 1H), 7.21-7.13 (m, 4H), 7.08-7.06 (m, 2H), 2.53 (d, J = 5.3 Hz, 1H), 2.39 (s, 3H), 2.28 (s, 3H), 2.15 (d, J = 5.3 Hz, 1H), 1.56 (s, 3H); ¹³C NMR (100 MHz, Acetone- d_6): δ 199.08, 139.88, 137.28, 136.31, 136.06, 133.13, 129.76, 129.67, 128.99, 125.30, 123.96, 50.13, 47.50, 28.58, 23.19, 21.56, 21.26; IR (ATR): 2924, 1694, 1516, 1458, 1152, 1043, 959, 818 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₉H₁₈NaOS [M+Na]⁺: 317.0971, Found: 317.0973.

Product **4c** was obtained in 46% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 126-128 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 10.76 min, t_r (major) = 16.94 min) gave the

isomeric composition of the major isomer: 98% ee; $[\alpha]_D^{25} = -358.0$ (c = 0.20, Acetone); ¹H NMR (400 MHz, Acetone- d_6): δ 7.44-7.42 (m, 1H), 7.31-7.27 (m, 4H), 7.21-7.19 (m, 2H), 2.54 (d, J = 5.5 Hz, 1H), 2.39 (s, 3H), 2.19 (d, J = 5.5 Hz, 1H), 1.59 (s, 3H); ¹³C NMR (100 MHz, Acetone- d_6): δ 199.49, 142.01, 136.24, 135.97, 133.10, 133.09, 131.59, 129.22, 129.18, 125.40, 124.04, 50.06, 46.69, 28.65, 22.82, 21.56; IR (ATR): 2924, 2862, 1692, 1491, 1344, 1013, 899, 777 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₅ClNaOS [M+Na]⁺: 337.0424, Found: 337.0422.



Product 4d was obtained in 46% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a colorless oil; HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; tr

(minor) = 14.65 min, tr (major) = 12.44 min) gave the isomeric composition of the major isomer: 98% ee; $[\alpha]_D^{25}$ = -46.7 (c = 0.24, Acetone); ¹H NMR (400 MHz, Acetone- d_6): δ 7.30-7.17 (m, 5H), 7.04-7.01 (m, 2H), 2.55 (d, J = 5.3 Hz, 1H), 2.36 (s, 3H), 2.30 (s, 3H), 2.13 (d, J = 5.3 Hz, 1H), 1.57 (s, 3H); ¹³C NMR (100 MHz, Acetone- d_6): δ 199.12, 143.08, 136.10, 135.97, 132.80, 132.78, 130.09, 129.80, 129.14, 127.86, 122.75, 50.52, 47.70, 28.63, 23.22, 21.49, 20.70; IR

(ATR): 2926, 1699, 1497, 1144, 1045, 856, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₉H₁₈NaOS [M+Na]⁺: 317.0971, Found: 317.0965.

Product **4e** was obtained in 98% yield, with a dr ratio of 7:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a colorless oil; HPLC analysis (Chiralcel OX-H, ¹PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; tr (minor) = 11.48 min, tr (major) = 10.14 min) gave the isomeric composition of the major isomer: 94% ee; $[\alpha]_D^{25} = -24.1$ (c = 0.58, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.31-7.20 (m, 5H), 6.96-6.93 (m, 2H), 5.40-5.39 (m, 1H), 4.29 (d, J = 10.2 Hz, 1H), 3.94 (dd, J = 10.2, 1.2 Hz, 1H), 2.51 (d, J = 5.5 Hz, 1H), 2.27 (dd, J = 5.5, 1.2 Hz, 1H), 1.90 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.41, 136.42, 134.75, 134.50, 131.30, 130.84, 128.31, 128.25, 128.21, 124.43, 122.29, 50.60, 49.42, 37.65, 32.22, 21.05; IR (ATR): 3026, 2920, 1603, 1495, 1352, 1167, 934, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C18H15BrNaOS [M+Na]⁺: 380.9919, Found: 380.9912.

Product **4f** was obtained in 95% yield, with a dr ratio of 6:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a colorless oil; HPLC analysis (Chiralcel OX-H, ¹PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; tr (minor) = 13.54 min, tr (major) = 12.05 min) gave the isomeric composition of the major isomer: 94% ee; $[\alpha]_D^{25} = -12.5$ (c = 0.45, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.26-6.99 (m, 4H), 6.96-6.93 (m, 2H), 5.43-5.42 (m, 1H), 4.28 (d, J = 10.1 Hz, 1H), 3.92 (dd, J = 10.2, 1.2 Hz, 1H), 2.49 (d, J = 5.5 Hz, 1H), 2.33 (s, 3H), 2.24 (dd, J = 5.5, 1.2 Hz, 1H), 1.91 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.45, 138.06, 134.66, 134.62, 133.36, 131.14, 130.83, 128.95, 128.13, 124.47, 122.22, 50.51, 49.56, 37.88, 32.34, 21.25, 21.07; IR (ATR): 2920, 2851, 1692, 1260, 1092, 935, 802 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₉H₁₇BrNaOS [M+Na]⁺: 395.0076, Found: 395.0069.



Product **4g** was obtained in 86% yield, with a dr ratio of 6:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a colorless oil; HPLC analysis (Chiralcel OX-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 13.67 min, t_r (major) = 12.87 min) gave the isomeric composition of the

major isomer: 95% ee; $[\alpha]_D^{25} = -8.3$ (c = 0.29, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.25-7.21 (m, 5H), 6.98-6.95 (m, 1H), 5.42-5.41 (m, 1H), 4.21 (d, J = 10.3 Hz, 1H), 3.90 (d, J = 10.3 Hz, 1H), 2.50

(d, J = 5.6 Hz, 1H), 2.21 (d, J = 6.0 Hz, 1H), 1.95 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.05, 135.02, 134.93, 134.26, 134.08, 132.64, 130.97, 128.53, 128.48, 124.32, 122.50, 49.58, 49.20, 37.23, 31.89, 21.14; IR (ATR): 3044, 2920, 1686, 1570, 1493, 1236, 932, 737 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₄BrClNaOS [M+Na]⁺: 414.9529, Found: 414.9528.



Product **4h** was obtained in 92% yield, with a dr ratio of 8:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a yellowish oil; HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 10.31 min, t_r (major) = 7.84 min) gave the isomeric composition of the

major isomer: 92% ee; $[\alpha]_D^{25} = -9.0$ (c = 0.40, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.27-7.13 (m, 5H), 6.70-6.69 (m, 1H), 5.16-5.15 (m, 1H), 4.20 (d, J = 10.2 Hz, 1H), 3.86 (dd, J = 10.1, 1.1 Hz, 1H), 2.42 (d, J = 5.5 Hz, 1H), 2.21 (s, 3H), 2.16 (dd, J = 5.4, 1.1 Hz, 1H), 1.78 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.48, 136.53, 134.61, 134.43, 131.37, 131.22, 130.64, 129.27, 128.25, 128.19, 121.82, 50.60, 49.90, 37.77, 32.35, 20.94, 20.54; IR (ATR): 2857, 1682, 1495, 1233, 1036, 935, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₉H₁₇BrNaOS [M+Na]⁺: 395.0076, Found: 395.0067.

Product **4i** was obtained in 87% yield, with a dr ratio of 3:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 123-125 °C); HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 10.58 min, t_r (major) = 8.41 min) gave the isomeric composition of the major isomer: 95% ee; $[\alpha]_D^{25} = -9.5$ (c = 0.55, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.29-7.20 (m, 6H), 6.95-6.92 (m, 1H), 5.43-5.42 (m, 1H), 5.07 (dd, J = 46.8, 9.7 Hz, 1H), 4.85 (dd, J = 47.4, 9.7 Hz, 1H), 2.46 (dd, J = 5.4, 1.7 Hz, 1H), 2.17 (t, J = 5.4 Hz, 1H), 1.89 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.40, 136.15, 134.67, 134.44, 131.16, 128.48, 128.18, 128.13, 124.29, 122.30, 83.71 (d, J = 172.0 Hz), 48.94 (d, J = 21.4 Hz), 45.29 (d, J = 4.4 Hz), 28.86 (d, J = 7.7 Hz), 21.06; ¹⁹F NMR (376 MHz, CDCl₃): δ -215.54; IR (ATR): 2922, 2855, 1697, 1497, 1462, 993, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C1₈H₁₅FNaOS [M+Na]⁺: 321.0720, Found: 321.0711. The minor isomer was isolated a white solid (m.p. 143-145 °C); HPLC analysis (Chiralpak AD-H, 'PrOH/hexane = 2/98, 1.0 mL/min, 230 nm; t_r (minor) = 11.32 min, t_r (major) = 9.99 min) gave the isomeric composition of the product: 98% ee; $[\alpha]_D^{25} = -280.0$ (c = 0.16, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.37-7.29 (m, 6H), 7.18-7.16 (m, 1H), 6.96-6.95 (m, 1H), 4.78 (dd, J = 47.6, 10.4 Hz, 1H), 4.75 (dd, J = 47.6, 10.4 Hz, 1H), 2.60 (dd, J = 5.8, 4.6 Hz, 1H), 2.42 (s, 3H), 2.20 (d, J = 5.8 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 199.28(d, J = 1.5 Hz), 137.14, 135.63, 133.88, 132.53, 129.79, 128.84, 128.60, 128.11, 123.46, 123.38 (d, J = 2.7 Hz), 84.74 (d, J = 175.8 Hz), 48.43 (d, J = 21.6 Hz), 47.75 (d, J = 4.4 Hz), 24.19 (d, J = 8.0 Hz), 21.63; ¹⁹F NMR (376 MHz, CDCl₃): δ -214.53; IR (ATR): 2922, 2853, 1699, 1497, 1150, 812, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₅FNaOS [M+Na]⁺: 321.0720, Found: 321.0718.



Product **4j** was obtained in 89% yield, with a dr ratio of 5:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a yellowish solid (m.p. 76-78 °C); HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 7.83 min, t_r (major) = 7.30 min) gave the isomeric

composition of the major isomer: 96% ee; $[\alpha]_D^{25} = 15.4$ (c = 0.26, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.30-6.92 (m, 6H), 5.46-5.45 (m, 1H), 5.05 (dd, J = 46.8, 9.7 Hz, 1H), 4.83 (dd, J = 47.5, 9.7 Hz, 1H), 2.44 (dd, J = 5.4, 1.6 Hz, 1H), 2.31 (s, 3H), 2.15 (t, J = 5.3 Hz, 1H), 1.90 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.43, 137.96, 134.58, 134.54, 133.06, 131.12, 129.09, 128.04, 124.32, 122.22, 83.81 (d, J = 171.6 Hz), 48.79 (d, J = 21.5 Hz), 45.39 (d, J = 4.4 Hz), 28.94 (d, J = 7.7 Hz), 21.21, 21.07; ¹⁹F NMR (376 MHz, CDCl₃): δ -215.10; IR (ATR): 2922, 1692, 1516, 1458, 1157, 980, 808 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₉H₁₇FNaOS [M+Na]⁺: 335.0876, Found: 335.0875.



Product **4k** was obtained in 78% yield, with a dr ratio of 5:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 96-98 °C); HPLC analysis (Chiralcel OX-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 14.04 min, t_r (major) = 12.72 min) gave the isomeric

composition of the major isomer: 97% ee; $[\alpha]p^{25} = 28.2$ (c = 0.27, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.26-7.21 (m, 5H), 6.97-6.95 (m, 1H), 5.47-5.46 (m, 1H), 5.04 (dd, J = 46.9, 9.8 Hz, 1H), 4.80 (dd, J = 47.5, 9.8 Hz, 1H), 2.45 (dd, J = 5.5, 1.7 Hz, 1H), 2.12 (t, J = 5.5 Hz, 1H), 1.95 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.03, 134.87, 134.73, 134.21, 133.99, 131.26, 128.67, 128.39, 124.15, 122.51, 83.42 (d, J = 172.3 Hz), 47.98 (d, J = 21.4 Hz), 45.20 (d, J = 4.1 Hz), 28.39 (d, J = 7.6 Hz), 21.15; ¹⁹F NMR (376 MHz, CDCl₃): δ -215.09; IR (ATR): 2922, 1690, 1491, 1157, 1092, 808, 725 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₈H₁₄ClFNaOS [M+Na]⁺: 355.0330, Found: 355.0331.

CH₂F

Product **41** was obtained in 85% yield, with a dr ratio of 9:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a yellowish solid (m.p. 129-131 °C); HPLC analysis (Chiralcel OX-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 23.79 min, t_r (major) = 22.03 min) gave the

isomeric composition of the major isomer: 93% ee; $[\alpha]_D^{25} = 125.6$ (c = 0.43, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.83-7.21 (m, 8H), 6.90-6.88 (m, 1H), 5.44 (s, 1H), 5.16 (dd, J = 46.7, 9.8 Hz, 1H), 4.94 (dd, J = 47.5, 9.8 Hz, 1H), 2.56 (dd, J = 5.5, 1.7 Hz, 1H), 2.32 (t, J = 5.5 Hz, 1H), 1.65 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.35, 134.70, 134.24, 133.11, 132.97, 131.14, 128.14, 127.95, 127.71, 126.50, 126.36, 124.14, 122.33, 83.85 (d, J = 172.1 Hz), 49.02 (d, J = 21.3 Hz), 45.37, 28.90 (d, J = 7.7 Hz), 20.89; ¹⁹F NMR (376 MHz, CDCl₃): δ -214.75; IR (ATR): 2855, 1686, 1504, 1452, 1153, 978, 737 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₂H₁₇FNaOS [M+Na]⁺: 371.0876, Found: 371.0872.

Product **4m** was obtained in 73% yield, with a dr ratio of 18:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a yellowish solid (m.p. 102-104 °C); HPLC analysis (Chiralcel OD-H, ^{*i*}PrOH/hexane = 2/98, 1.0 mL/min, 230 nm; t_r (minor) = 13.50 min, t_r (major) = 9.28 min) gave the isomeric composition of the major isomer: 93% ee; $[\alpha]_D^{25} = -21.8$ (c = 0.22, Acetone); ¹H NMR (400 MHz, Acetone- d_6): δ 7.38-7.37 (m, 1H), 7.35-7.33 (m, 1H), 7.08-7.07 (m, 1H), 7.06-7.03 (m, 1H), 7.01-6.99 (m, 1H), 6.05-6.04 (m, 1H), 5.06 (dd, J = 46.8, 9.9 Hz, 1H), 4.84 (dd, J = 47.9, 9.9 Hz, 1H), 2.57 (t, J = 5.9 Hz, 1H), 2.52 (dd, J = 5.9, 1.8 Hz, 1H), 2.00 (s, 3H); ¹³C NMR (100 MHz, Acetone- d_6): δ 201.70, 139.51, 135.73, 134.89, 131.39, 130.49, 129.22, 127.68, 126.97, 124.07, 123.14, 83.97 (d, J = 170.1 Hz), 47.07 (d, J = 4.1 Hz), 44.63 (d, J = 21.6 Hz), 29.06 (d, J = 8.0 Hz), 21.08; ¹⁹F NMR (376 MHz, Acetone- d_6): δ -213.46; IR (ATR): 2924, 1705, 1653, 1470, 1236, 1113, 1034, 704 cm⁻¹; HRMS (EI): Exact mass calcd for C₁₆H₁₃FOS₂ [M]⁺: 304.0392, Found: 304.0395.



Product **4n** was obtained in 96% yield, with a dr ratio of 5:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a reddish solid (m.p. 107-109 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 10.75 min, t_r (major) = 8.75 min) gave the isomeric

composition of the major isomer: 95% ee; $[\alpha]_D^{25} = 11.6$ (c = 0.38, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.26-7.24 (m, 5H), 6.77-6.76 (m, 1H), 5.27-5.26 (m, 1H), 5.07 (dd, J = 46.8, 9.7 Hz, 1H), 4.85 (dd, J = 47.5, 9.7 Hz, 1H), 2.44 (dd, J = 5.4, 1.7 Hz, 1H), 2.28 (s, 3H), 2.14 (t, J = 5.4 Hz, 1H), 1.86 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 202.44, 136.25, 134.52, 134.34, 131.35, 130.93, 129.17, 128.40, 128.10, 121.64, 83.75 (d, J = 171.8 Hz), 48.93 (d, J = 21.3 Hz), 45.77 (d, J = 4.3 Hz), 28.91 (d, J = 7.7 Hz), 20.94, 20.51; ¹⁹F NMR (376 MHz, CDCl₃): δ -215.01; IR (ATR): 3059, 1684, 1495, 1458, 1265, 854, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₉H₁₇FNaOS [M+Na]⁺: 335.0876, Found: 335.0871.

Product **40** was obtained in 68% yield, with a dr ratio of 18:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a reddish solid mL/min, 230 nm; tr (minor) = 11.99 min, tr (major) = 10.71 min) gave the isomeric composition of the major isomer: 96% ee; $[\alpha]_D^{25} = 58.3$ (c = 0.46, Acetone); ¹H NMR (400 MHz, Acetone- d_6): δ 7.52-7.50 (m, 1H), 7.37-7.35 (m, 1H), 7.06-7.04 (m, 2H), 6.72-6.60 (m, 2H), 6.44 (t, J = 55.8 Hz, 1H), 5.60-5.59 (m, 1H), 3.79 (s, 3H), 2.68 (dt, J = 6.1, 1.8 Hz, 1H), 2.49 (dd, J = 7.3, 6.1 Hz, 1H), 1.92 (s, 3H); ¹³C NMR (100 MHz, Acetone- d_6): δ 203.26, 161.23, 135.92, 135.08, 134.46, 134.31, 131.61, 129.57, 125.90, 123.39, 122.99 (d, J = 1.7 Hz), 115.79 (dd, J = 237.9, 236.4 Hz), 114.52 (d, J = 12.0 Hz), 55.86, 48.75 (dd, J = 29.9, 25.3 Hz), 45.90 (d, J = 4.3 Hz), 28.64 (d, J = 8.0Hz), 21.22; ¹⁹F NMR (376 MHz, Acetone- d_6): δ -114.62 (d, J = 289.5 Hz, 1F), -121.26 (d, J = 289.1Hz, 1F); IR (ATR): 2837, 1688, 1514, 1325, 1157, 930, 760 cm⁻¹; HRMS (ESI): Exact mass calcd for C₁₉H₁₆F₂NaO₂S [M+Na]⁺: 369.0731, Found: 369.0724.



Product **4p** was obtained in 62% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a reddish solid (m.p. 143-145 °C); HPLC analysis (Chiralcel OZ-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 9.98 min, t_r (major) = 8.71 min) gave the isomeric composition of the major isomer: 97% ee; $[\alpha]_D^{25} = 100.7$ (*c* = 0.27, Acetone); ¹H

NMR (400 MHz, Acetone-*d*₆): δ 7.51-7.49 (m, 1H), 7.04-7.02 (m, 1H), 6.90-6.89 (m, 1H), 6.71-6.59 (m, 2H), 6.43 (t, *J* = 55.8 Hz, 1H), 5.44-5.43 (m, 1H), 3.79 (s, 3H), 2.67 (dt, *J* = 6.1, 1.8 Hz, 1H), 2.47 (dd, *J* = 7.3, 6.0 Hz, 1H), 2.27 (s, 3H), 1.89 (s, 3H); ¹³C NMR (100 MHz, Acetone-*d*₆): δ 202.92,

161.03, 135.68, 134.92, 134.21, 134.11, 132.12, 131.11, 130.35, 123.15, 122.91 (d, J = 1.7 Hz), 115.64 (dd, J = 237.9, 235.0 Hz), 114.31 (d, J = 4.2 Hz), 55.68, 48.60 (dd, J = 29.7, 25.3 Hz), 46.22 (d, J = 4.0 Hz), 28.57 (d, J = 8.0 Hz), 20.97, 20.29; ¹⁹F NMR (376 MHz, Acetone- d_6): δ -114.62 (d, J = 289.5 Hz, 1F), -121.31 (d, J = 289.2 Hz, 1F); IR (ATR): 2963, 1682, 1516, 1252, 1153, 937, 766 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₀H₁₈F₂NaO₂S [M+Na]⁺: 383.0888, Found: 383.0878.

Product **4q** was obtained in 92% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 141-143 °C); HPLC analysis (Chiralcel OX-H, ⁷PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 6.48 min, t_r (major) = 8.09 min) gave the isomeric composition of the major isomer: 94% ee; $[\alpha]_{D}^{25}$ = -101.5 (*c* = 0.26, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.34-7.32 (m, 1H), 7.26-7.18 (m, 5H), 7.12-7.08 (m, 1H), 6.75-6.71 (m, 1H), 5.65-5.62 (m, 1H), 2.52 (d, *J* = 3.9 Hz, 1H), 2.09 (d, *J* = 3.9 Hz, 1H), 1.19 (s, 6H), 1.18 (s, 6H); ¹³C NMR (100 MHz, CDCl₃): δ 204.20, 136.54, 135.22, 135.13, 130.89, 128.35, 127.18, 127.01, 125.00, 122.72, 122.53, 84.30, 46.51, 31.40, 24.90, 24.56 (Note: the carbon attached to boron was not observed due to quadrupole broadening caused by the ¹¹B nucleus); IR (ATR): 2978, 1688, 1447, 1371, 1250, 1186, 849, 743 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₂H₂₃BNaO₃S [M+Na]⁺: 401.1357, Found: 401.1357.

Product **4r** was obtained in 91% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a yellowish solid (m.p. 121-123 °C); HPLC analysis (Chiralcel OX-H, ⁱPrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 6.40 min, t_r (major) = 7.75 min) gave the isomeric composition of the major isomer: 99% ee; $[\alpha]p^{25} = -192.0$ (c = 0.15, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.26-7.17 (m, 6H), 6.93-6.91 (m, 1H), 5.40-5.39 (m, 1H), 2.50 (d, J = 3.9 Hz, 1H), 2.07 (d, J = 3.9 Hz, 1H), 1.91 (s, 3H), 1.20 (s, 6H), 1.19 (s, 6H); ¹³C NMR (100 MHz, CDCl₃): δ 204.68, 136.71, 135.11, 134.70, 131.54, 130.99, 128.29, 128.23, 127.73, 127.05, 123.57, 122.31, 84.28, 46.41, 31.31, 24.91, 24.57, 21.13 (Note: the carbon attached to boron was not observed due to quadrupole broadening caused by the ¹¹B nucleus); IR (ATR): 2976, 1690, 1468, 1369, 1256, 1186, 847, 702 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₃H₂₅BNaO₃S [M+Na]⁺: 415.1514, Found: 415.1514. Product **4s** was obtained in 95% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 163-165 °C); HPLC analysis (Chiralcel OX-H, ¹PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; t_r (minor) = 10.77 min, t_r (major) = 13.59 min) gave the isomeric composition of the major isomer: 98% ee; $[\alpha]_D^{25} = -85.3$ (c = 0.15, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.26-7.16 (m, 5H), 6.90-6.89 (m, 1H), 6.30-6.27 (m, 1H), 5.53-5.51 (m, 1H), 3.71 (s, 3H), 2.46 (d, J =3.9 Hz, 1H), 2.01 (d, J = 4.0 Hz, 1H), 1.19 (s, 6H), 1.18 (s, 6H); ¹³C NMR (100 MHz, CDCl₃): δ 204.51, 158.83, 136.73, 136.02, 130.90, 128.32, 127.12, 127.08, 123.25, 111.21, 108.54, 84.25, 55.47, 46.01, 30.89, 24.89, 24.56 (Note: the carbon attached to boron was not observed due to quadrupole broadening caused by the ¹¹B nucleus); IR (ATR): 2974, 1694, 1489, 1371, 1287, 1186, 847, 702 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₃H₂₅BNaO4S [M+Na]⁺: 431.1463, Found: 431.1467.

Product **4t** was obtained in 87% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a yellowish solid (m.p. 117-119 °C); HPLC analysis (Chiralcel OX-H, ^{*i*}PrOH/hexane = 1/99, 1.0 mL/min, 254 nm; t_r (minor) = 7.97 min, t_r (major) = 9.14 min) gave the isomeric composition of the major isomer: 98% ee; $[\alpha]p^{25} = -186.0$ (c = 0.20, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.30-7.22 (m, 5H), 7.19-7.17 (m, 1H), 7.09-7.07 (m, 1H), 5.56-5.55 (m, 1H), 2.55 (d, J =4.0 Hz, 1H), 2.11 (d, J = 4.1 Hz, 1H), 1.19 (s, 6H), 1.18 (s, 6H); ¹³C NMR (100 MHz, CDCl₃): δ 203.29, 137.04, 135.96, 133.37, 131.11, 130.82, 128.57, 127.57, 127.08, 123.49, 122.85, 84.46, 46.51, 31.75, 24.91, 24.57 (Note: the carbon attached to boron was not observed due to quadrupole broadening caused by the ¹¹B nucleus); IR (ATR): 2978, 1458, 1371, 1250, 1188, 847, 702 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₂H₂₂BClNaO₃S [M+Na]⁺: 435.0967, Found: 435.0972.



Product **4u** was obtained in 96% yield, with a dr ratio of 13:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 79-81 °C); HPLC analysis (Chiralcel OX-H, ^{*i*}PrOH/hexane = 0.1/99.9, 1.0

mL/min, 230 nm; t_r (minor) = 13.88 min, t_r (major) = 15.51 min) gave the isomeric composition of the major isomer: 98% ee; $[\alpha]_D^{25}$ = -153.3 (*c* = 0.24, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.37-7.33

(m, 1H), 7.25-7.22 (m, 1H), 7.19-7.17 (m, 1H), 7.15-7.12 (m, 1H), 7.01-6.97 (m, 1H), 6.91-6.88 (m, 1H), 6.43-6.40 (m, 1H), 5.32-5.31 (m, 1H), 2.51 (d, J = 4.1 Hz, 1H), 2.17 (d, J = 4.2 Hz, 1H), 1.87 (s, 3H), 0.04 (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ 204.85, 141.14, 135.57, 134.35, 131.10, 131.04, 129.34, 128.47, 127.98, 127.47, 126.21, 124.32, 122.11, 47.23, 46.12, 32.22, 21.07, 0.03; IR (ATR): 2361, 1695, 1470, 1248, 1070, 841, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₀H₂₂NaOSSi [M+Na]⁺: 361.1053, Found: 361.1046.

Product 4v was obtained in 97% yield, with a dr ratio of >20:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 136-138 °C); HPLC analysis (Chiralcel OX-H, ¹PrOH/hexane = 1/99, 1.0 mL/min, 230 nm; tr (minor) = 12.79 min, tr (major) = 13.45 min) gave the isomeric composition of the major isomer: 90% ee; $[\alpha]_D^{25} = -22.5$ (c = 0.16, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.36-7.31 (m, 1H), 7.22-7.19 (m, 1H), 7.16-7.12 (m, 1H), 7.03-6.99 (m, 1H), 6.89-6.88 (m, 1H), 6.47-6.44 (m, 1H), 6.24-6.21 (m, 1H), 5.45-5.42 (m, 1H), 3.72 (s, 3H), 2.47 (d, J = 4.2 Hz, 1H), 2.11 (d, J = 4.2 Hz, 1H), 0.02 (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ 204.62, 158.64, 141.13, 135.46, 131.08, 129.23, 128.54, 128.14, 127.64, 126.25, 123.96, 110.94, 108.22, 55.49, 46.80, 45.24, 31.88, 0.03; IR (ATR): 2955, 1697, 1489, 1441, 1121, 845, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₀H₂₂NaO₂SSi [M+Na]⁺: 377.1002, Found: 377.1008.



Product **4w** was obtained in 90% yield, with a dr ratio of 11:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a reddish solid (m.p. 60-62 °C); HPLC analysis (Chiralcel OX-H, ^{*i*}PrOH/hexane = 0.5/99.5, 1.0 mL/min, 254 nm; t_r (minor) = 7.91 min, t_r (major) = 6.90 min) gave the isomeric

composition of the major isomer: 98% ee; $[\alpha]_D^{25} = -113.3$ (c = 0.12, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.36-7.32 (m, 1H), 7.24-7.21 (m, 1H), 7.15-7.11 (m, 1H), 7.02-6.97 (m, 1H), 6.75-6.74 (m, 1H), 6.45-6.42 (m, 1H), 5.17-5.16 (m, 1H), 2.50 (d, J = 4.0 Hz, 1H), 2.28 (s, 3H), 2.15 (d, J = 4.1 Hz, 1H), 1.85 (s, 3H), 0.04 (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ 204.96, 141.24, 135.53, 134.22, 131.14, 131.09, 130.85, 129.34, 128.57, 128.41, 127.94, 126.16, 121.71, 47.72, 46.15, 32.41, 20.97, 20.53, 0.05; IR (ATR): 2949, 1699, 1487, 1248, 1148, 849, 700 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₁H₂₄NaOSSi [M+Na]⁺: 375.1209, Found: 375.1208.

Product **4x** was obtained in 74% yield, with a dr ratio of 10:1 indicated by ¹H NMR analysis of crude reaction mixture; The major isomer was isolated as a white solid (m.p. 142-144 °C); HPLC analysis (Chiralcel OX-H, ¹PrOH/hexane = 0.1/99.9, 1.0 mL/min, 254 nm; tr (minor) = 9.73 min, tr (major) = 12.67 min) gave the isomeric composition of the major isomer: 92% ee; $[\alpha]_{D}^{25}$ = -123.2 (*c* = 0.38, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 7.38-7.34 (m, 1H), 7.26-7.22 (m, 2H), 7.14-7.10 (m, 2H), 7.00-6.96 (m, 1H), 6.43-6.40 (m, 1H), 5.63-5.62 (m, 1H), 2.52 (d, *J* = 4.1 Hz, 1H), 2.18 (d, *J* = 4.1 Hz, 1H), 0.88 (s, 9H), 0.02 (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ 205.02, 147.83, 141.13, 134.99, 131.28, 131.06, 129.19, 128.73, 128.26, 126.36, 124.03, 122.02, 120.80, 47.31, 46.00, 34.35, 32.12, 31.08, 0.01; IR (ATR): 2968, 1701, 1246, 1128, 990, 852, 745 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₃H₂₈NaOSSi [M+Na]⁺: 403.1522, Found: 403.1513.

3. Transformations of product 3a to 6



To a Schlenk tube was sequentially added **3a** (53.2 mg, 0.2 mmol), nitrone **5** (40.5 mg, 0.3 mmol, 1.5 equivs) and Sc(OTf)₃ (9.8 mg, 0.02 mmol, 10 mol %), followed by the addition of anhydrous DCE (2.0 mL). The reaction mixture was stirred vigorously at 50 °C until almost full consumption of **3a** by TLC analysis (about 7 h). The reaction mixture was rapidly passed through a short pad of silica gel, and washed with Et₂O. The obtained organic solution was concentrated in vacuo to give the crude product. To determine the diastereoselectivity of the product, the residue was first dissolved in CDCl₃, and took some samples for ¹H NMR analysis. Then the sample for analysis and rest crude product were recombined for column chromatography purification to afford product **6** (73.4 mg, 92%), using PE/EtOAc (50/1, v/v) as the eluent.



Product **6** was obtained in 92% yield, with a dr ratio of 3:1 indicated by ¹H NMR analysis of crude reaction mixture; The major distereoisomer was isolated as a white solid (m.p. 144-146 °C); HPLC analysis (Chiralcel OX-H, ^{*i*}PrOH/hexane = 0.5/99.5,

1.0 mL/min, 230 nm; tr (minor) = 11.70 min, tr (major) = 14.02 min) gave the isomeric composition of

the product: 98% ee; $[\alpha]_D^{25} = -143.3$ (c = 0.12, Acetone); ¹H NMR (400 MHz, CDCl₃): δ 8.42-8.41 (m, 1H), 7.47-7.44 (m, 2H), 7.40-7.36 (m, 2H), 7.34-7.30 (m, 1H), 7.15-6.85 (m, 6H), 6.69-6.66 (m, 1H), 5.69 (dd, J = 12.0, 2.5 Hz, 1H), 4.21 (s, 1H), 2.56 (s, 3H), 2.54 (s, 3H), 2.49 (dd, J = 13.3, 12.0 Hz, 1H), 2.04 (dd, J = 13.3, 2.5 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 207.62, 140.10, 139.07, 135.55, 135.47, 131.22, 130.46, 129.21, 128.96, 128.65, 128.32, 127.95, 127.74, 126.56, 122.57, 76.71, 75.46, 63.95, 44.91, 42.54, 21.86; IR (ATR): 2918, 1701, 1493, 1454, 1047, 972, 698 cm⁻¹; HRMS (ESI): Exact mass calcd for C₂₅H₂₃NNaO₂S [M+Na]⁺: 424.1342, Found: 424.1340.

4. X-ray crystallographic data of 3a, 4l and 6

Data intensity of $3a^3$ was collected using a 'XtaLAB AFC12 (RINC)' diffractometer at 99.9(9) K. Data collection and reduction were done by using Olex2 and the structure was solved with the ShelXS structure solution program using Intrinsic Phasing and refined with the ShelXL refinement package using Least Squares minimization. Crystal data for 3a: C₁₇H₁₄OS, T = 99.9(9) K, orthorhombic, P2₁₂₁₂₁, a = 7.54810(10) Å, b = 11.5576(2) Å, c = 15.3921(3) Å, $a = 90^{\circ}$, $\beta = 90^{\circ}$, $\gamma = 90^{\circ}$, V = 1342.77(4) Å³. Z = 4, $\rho_{calc} = 1.317$ g/cm³. 25631 reflections collected, 2375 [R_{int} = 0.0506, R_{sigma} = 0.0203] independent reflections, R₁ = 0.0252, wR₂ = 0.06080 ($I > 2\sigma(I)$, final), R₁ = 0.0264, wR₂ = 0.0613 (all data), GOF = 1.038, and 173 parameters.



Table S1. Crystal data and structure refinement for 3a.

Identification code	3 a
Empirical formula	$C_{17}H_{14}OS$
Formula weight	266.34
Temperature/K	99.9(9)
Crystal system	orthorhombic
Space group	P212121
a/Å	7.54810(10)
b/Å	11.5576(2)

³ Supplementary crystallographic data have been deposited at Cambridge Crystallographic Data Center (CCDC number: 2053474).

c/Å	15.3921(3)
$\alpha /^{\circ}$	90
β/°	90
$\gamma^{\prime \circ}$	90
Volume/Å ³	1342.77(4)
Ζ	4
$\rho_{calc}g/cm^3$	1.317
μ/mm^{-1}	2.029
F(000)	560.0
Crystal size/mm ³	$0.36 \times 0.22 \times 0.18$
Radiation	$CuK\alpha \ (\lambda = 1.54184)$
2Θ range for data collection/°	9.57 to 134.156
Index ranges	$\textbf{-9} \le h \le 8, \textbf{-13} \le k \le 13, \textbf{-18} \le l \le 18$
Reflections collected	25631
Independent reflections	2375 [$R_{int} = 0.0506, R_{sigma} = 0.0203$]
Data/restraints/parameters	2375/0/173
Goodness-of-fit on F ²	1.038
Final R indexes $[I \ge 2\sigma(I)]$	$R_1 = 0.0252, wR_2 = 0.0608$
Final R indexes [all data]	$R_1 = 0.0264, wR_2 = 0.0613$
Largest diff. peak/hole / e Å ⁻³	0.14/-0.15
Flack parameter	-0.012(7)

Table S2. Fractional Atomic Coordinates (×10⁴) and Equivalent Isotropic Displacement Parameters (Å²×10³) for **3a**. U_{eq} is defined as 1/3 of of the trace of the orthogonalised U_{IJ} tensor.

Atom	x	у	Z.	U(eq)
S1	-132.3(7)	5120.1(4)	2904.6(3)	25.99(14)
01	310(2)	6692.8(13)	1666.9(9)	31.0(4)
C10	3302(3)	5368.6(16)	3210.0(12)	19.7(4)
C14	1913(3)	3945.8(17)	4147.0(13)	24.0(4)
C15	1810(3)	4762.5(17)	3485.3(12)	22.1(4)
C12	5051(3)	4359.5(15)	4287.9(11)	21.2(4)
C6	5019(3)	7814.7(15)	3088.7(12)	22.6(4)
C9	2917(3)	6180.0(18)	2487.9(12)	21.7(4)
C16	1032(3)	6132.0(18)	2232.5(13)	24.8(5)
C4	7964(3)	8155.5(19)	3636.1(15)	31.5(5)
C11	4919(3)	5162.4(15)	3608.8(11)	20.4(4)
C7	3773(3)	7402.3(18)	2400.5(13)	24.6(5)
C5	6839(3)	7795.9(18)	2974.2(14)	27.1(5)

C13	3536(3)	3761.5(18)	4541.4(13)	23.2(5)
C8	4252(3)	6474.2(19)	1775.6(13)	25.7(5)
C17	6791(3)	4157.3(19)	4729.2(14)	26.9(5)
C2	5458(3)	8565.8(19)	4537.2(15)	34.8(6)
C1	4337(3)	8208.9(18)	3878.7(14)	29.7(5)
C3	7273(3)	8540.8(19)	4418.2(16)	33.1(5)

Table S3. Anisotropic Displacement Parameters ($Å^2 \times 10^3$) for **3a**. The Anisotropic displacement factor exponent takes the form: $-2\pi^2[h^2a^{*2}U_{11}+2hka^*b^*U_{12}+...]$.

Atom	U ₁₁	U ₂₂	U ₃₃	U ₂₃	U ₁₃	U_{12}
S 1	17.8(3)	33.1(3)	27.0(2)	-0.6(2)	-1.8(2)	-1.0(2)
01	28.3(9)	41.4(9)	23.4(7)	1.8(6)	-5.8(7)	6.3(7)
C10	21.1(11)	20.5(9)	17.5(9)	-3.1(8)	1.9(8)	0.9(8)
C14	22.3(12)	24.0(10)	25.6(10)	-0.4(8)	4.6(9)	-3.4(8)
C15	20.2(11)	24.3(10)	22.0(9)	-4.4(8)	1.1(8)	0.9(9)
C12	23.6(11)	20.0(9)	19.9(9)	-0.9(7)	2.3(9)	0.5(9)
C6	23.5(11)	19.0(9)	25.3(10)	4.5(7)	-0.7(9)	0.0(9)
С9	23.4(12)	23.5(10)	18.2(9)	0.4(8)	-0.1(8)	1.6(9)
C16	24.8(12)	29.1(10)	20.5(10)	-5.0(9)	0.0(9)	2.7(9)
C4	24.2(12)	28.6(11)	41.9(13)	5.7(10)	-2.9(10)	-3.0(10)
C11	19.2(10)	21.7(9)	20.4(9)	-0.5(7)	3.1(8)	-0.6(9)
C7	25.9(13)	24.5(10)	23.4(10)	5.6(8)	0.9(9)	0.4(9)
C5	28.6(13)	25.0(10)	27.7(11)	4.4(9)	3.3(10)	-2.7(9)
C13	27.1(12)	21.4(10)	21.1(10)	1.0(8)	2.7(8)	0.9(8)
C8	22.7(11)	36.4(12)	18.1(9)	2.8(9)	1.0(8)	-0.7(9)
C17	25.1(13)	28.4(11)	27.3(11)	6.0(9)	-1.3(9)	0.8(9)
C2	37.7(15)	32.6(12)	33.9(12)	-9.3(10)	-2.2(10)	2.8(10)
C1	26.4(13)	29.8(11)	32.9(12)	-3.8(9)	0.6(9)	3.0(9)
C3	35.1(14)	27.0(11)	37.3(13)	-2.0(9)	-9.2(11)	-2.8(10)

Table S4. Bond Lengths for 3a.

Atom	Atom	Length/Å	Atom	Atom	Length/Å
S 1	C15	1.766(2)	C6	C7	1.494(3)
S 1	C16	1.792(2)	C6	C5	1.385(3)
01	C16	1.215(3)	C6	C1	1.397(3)
C10	C15	1.392(3)	C9	C16	1.477(3)
C10	C9	1.483(3)	C9	C7	1.559(3)

C10	C11	1.386(3)	С9	C8	1.527(3)
C14	C15	1.391(3)	C4	C5	1.390(3)
C14	C13	1.384(3)	C4	C3	1.385(3)
C12	C11	1.401(3)	C7	C8	1.485(3)
C12	C13	1.392(3)	C2	C1	1.383(3)
C12	C17	1.497(3)	C2	C3	1.383(4)

 Table S5. Bond Angles for 3a.

Atom	Atom	Atom	Angle/°	Atom	Atom	Atom	Angle/°
C15	S 1	C16	92.16(10)	C16	C9	C7	114.22(18)
C15	C10	C9	112.80(18)	C16	C9	C8	116.92(17)
C11	C10	C15	119.39(18)	C8	C9	C7	57.52(14)
C11	C10	C9	127.81(19)	01	C16	S 1	122.83(17)
C13	C14	C15	118.38(19)	01	C16	С9	127.1(2)
C10	C15	S 1	113.57(15)	C9	C16	S 1	110.09(15)
C14	C15	S 1	125.16(16)	C3	C4	C5	120.2(2)
C14	C15	C10	121.27(19)	C10	C11	C12	120.46(19)
C11	C12	C17	120.30(19)	C6	C7	C9	119.24(17)
C13	C12	C11	118.6(2)	C8	C7	C6	122.4(2)
C13	C12	C17	121.06(17)	C8	C7	С9	60.15(13)
C5	C6	C7	121.92(19)	C6	C5	C4	120.5(2)
C5	C6	C1	118.8(2)	C14	C13	C12	121.86(19)
C1	C6	C7	119.3(2)	C7	C8	С9	62.32(14)
C10	C9	C7	123.80(18)	C1	C2	C3	120.2(2)
C10	C9	C8	123.33(19)	C2	C1	C6	120.7(2)
C16	C9	C10	111.37(18)	C2	C3	C4	119.6(2)

Table S6. Hydrogen Atom Coordinates (Å×104) and Isotropic Displacement Parameters (Å²×10³) for **3a**.

Atom	x	У	Ζ	U(eq)
H14	915.53	3533.69	4320.13	29
H4	9184.3	8137.47	3553.59	38
H11	5921.97	5559.71	3424.37	25
H7	2984.99	7999.51	2166.39	30
H5	7311.51	7540.69	2450.19	33
H13	3617.12	3222.86	4988.51	28
H8A	5427.2	6140.36	1821.75	31

H8B	3808.83	6543.3	1186.34	31
H17A	7508.06	3655.86	4377.71	40
H17B	6591.36	3803.35	5284.89	40
H17C	7388.72	4882.81	4807.97	40
H2	4988.84	8823.27	5061.43	42
H1	3117.34	8231.59	3962.78	36
H3	8026.87	8781.28	4860.84	40

Data intensity of **4l**⁴ was collected using a 'XtaLAB AFC12 (RINC)' diffractometer at 99.99(10) K. Data collection and reduction were done by using Olex2 and the structure was solved with the ShelXS structure solution program using Intrinsic Phasing and refined with the ShelXL refinement package using Least Squares minimization. Crystal data for **4l**: C₂₂H₁₇FOS, T = 99.99(10) K, monoclinic, P2₁, a = 6.6246(2) Å, b = 12.6387(3) Å, c = 10.4032(2) Å, $a = 90^{\circ}$, $\beta = 90.958(2)^{\circ}$, $\gamma = 90^{\circ}$, V = 870.90(4) Å³. Z = 2, $\rho_{calc} = 1.329$ g/cm³. 16182 reflections collected, 3057 [R_{int} = 0.0661, R_{sigma} = 0.0370] independent reflections, R₁ = 0.0705, wR₂ = 0.1874 ($I > 2\sigma(I)$, final), R₁ = 0.0743, wR₂ = 0.1912 (all data), GOF = 1.038, and 227 parameters.



 Table S7. Crystal data and structure refinement for 41.

41
$C_{22}H_{17}FOS$
348.41
99.99(10)
monoclinic
P21
6.6246(2)
12.6387(3)
10.4032(2)
90

⁴ Supplementary crystallographic data have been deposited at Cambridge Crystallographic Data Center (CCDC number: 2053475).

β/°	90.958(2)
$\gamma^{/\circ}$	90
Volume/Å ³	870.90(4)
Ζ	2
$\rho_{calc}g/cm^3$	1.329
μ/mm^{-1}	1.782
F(000)	364.0
Crystal size/mm ³	$0.36 \times 0.28 \times 0.12$
Radiation	$CuK\alpha (\lambda = 1.54184)$
2Θ range for data collection/°	8.5 to 134.03
Index ranges	$-7 \le h \le 7, -15 \le k \le 15, -12 \le l \le 12$
Reflections collected	16182
Independent reflections	$3057 [R_{int} = 0.0661, R_{sigma} = 0.0370]$
Data/restraints/parameters	3057/1/227
Goodness-of-fit on F ²	1.038
Final R indexes [I>= 2σ (I)]	$R_1 = 0.0705, wR_2 = 0.1874$
Final R indexes [all data]	$R_1 = 0.0743, wR_2 = 0.1912$
Largest diff. peak/hole / e Å ⁻³	0.74/-0.40
Flack parameter	0.017(14)

Table S8. Fractional Atomic Coordinates (×10⁴) and Equivalent Isotropic Displacement Parameters (Å²×10³) for **41**. U_{eq} is defined as 1/3 of of the trace of the orthogonalised U_{IJ} tensor.

Atom	x	У	\boldsymbol{z}	U(eq)
S 1	721(2)	5426.5(15)	9719.0(15)	47.0(5)
F1	-511(7)	3154(4)	6863(6)	69.0(15)
01	775(8)	3333(5)	9860(5)	57.1(15)
C6	4019(9)	5230(5)	8350(5)	34.9(14)
C5	5781(8)	5554(5)	7752(5)	33.3(13)
C4	6291(10)	6629(5)	7741(6)	34.4(14)
C7	3268(9)	4130(5)	8471(6)	35.6(14)
C10	3018(9)	3410(5)	7270(6)	36.1(14)
C19	4278(11)	4917(5)	3692(7)	42.6(16)
C1	2802(10)	5975(6)	8942(6)	39.6(15)
C3	5054(11)	7360(6)	8333(7)	41.8(16)
С9	4584(10)	3169(6)	8292(6)	38.4(14)
C14	5703(10)	4247(5)	4176(7)	43.5(16)
C12	3536(10)	3901(6)	6008(6)	40.8(15)
C17	6456(14)	5192(6)	1849(7)	55(2)

C22	8195(10)	6987(6)	7069(7)	41.8(16)
C2	3311(12)	7051(6)	8929(6)	42.2(16)
C15	7555(11)	4046(6)	3583(7)	46.2(16)
C16	7940(13)	4521(6)	2432(7)	51.0(17)
C8	1551(9)	4105(6)	9395(6)	44.7(18)
C18	4724(11)	5406(6)	2458(6)	47.1(15)
C11	1283(10)	2632(6)	7185(8)	47.0(17)
C13	5269(11)	3722(6)	5400(6)	44.2(16)
C21	2147(13)	4658(6)	5458(7)	49.3(18)
C20	2459(11)	5133(6)	4314(7)	48.1(17)

Table S9. Anisotropic Displacement Parameters ($Å^2 \times 10^3$) for **41**. The Anisotropic displacement factor exponent takes the form: $-2\pi^2[h^2a^{*2}U_{11}+2hka^*b^*U_{12}+...]$.

Atom	U11	U22	U33	U23	U13	U12
S1	32.9(8)	68.2(11)	40.2(8)	10.3(8)	6.1(6)	8.3(8)
F1	35(2)	67(3)	105(4)	37(3)	-22(2)	-13(2)
01	46(3)	72(4)	53(3)	26(3)	12(2)	0(3)
C6	31(3)	48(4)	25(3)	7(3)	-6(2)	2(3)
C5	26(3)	45(3)	29(3)	10(3)	-3(2)	-1(3)
C4	31(3)	45(3)	27(3)	3(2)	-10(2)	2(3)
C7	29(3)	49(4)	29(3)	14(3)	0(2)	-4(3)
C10	32(3)	46(4)	31(3)	10(3)	-7(2)	-6(3)
C19	40(4)	33(3)	54(4)	-9(3)	-16(3)	4(3)
C1	30(3)	63(4)	25(3)	4(3)	-6(2)	3(3)
C3	41(4)	47(4)	37(3)	-2(3)	-8(3)	-1(3)
C9	31(3)	48(4)	36(3)	12(3)	-2(3)	0(3)
C14	29(3)	39(4)	62(4)	-17(3)	-3(3)	0(3)
C12	40(4)	50(4)	32(3)	3(3)	-7(3)	-16(3)
C17	78(6)	47(4)	41(3)	-3(3)	3(3)	-18(4)
C22	36(4)	45(4)	44(4)	10(3)	-4(3)	-5(3)
C2	46(4)	49(4)	31(3)	-3(3)	-2(3)	13(3)
C15	42(4)	49(4)	47(4)	-3(3)	-1(3)	5(3)
C16	53(4)	53(4)	47(4)	-2(3)	5(3)	-2(4)
C8	21(3)	75(5)	38(3)	17(3)	2(2)	2(3)
C18	47(4)	45(4)	49(3)	-2(3)	-6(3)	0(4)
C11	31(4)	54(4)	56(4)	16(3)	-6(3)	-6(3)
C13	37(4)	61(4)	35(3)	-1(3)	-5(3)	-13(3)
C21	64(5)	44(4)	40(4)	0(3)	-9(3)	2(3)

C20	46(4)	47(4)	50(4)	2(3)	-5(3)	8(3)
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Table S10. Bond Lengths for 4l.

Atom	Atom	Length/Å	Atom	Atom	Length/Å
S1	C1	1.752(7)	C10	C11	1.514(9)
S1	C8	1.792(8)	C19	C14	1.359(10)
F1	C11	1.395(8)	C19	C18	1.460(11)
01	C8	1.208(9)	C19	C20	1.404(11)
C6	C5	1.394(8)	C1	C2	1.402(11)
C6	C7	1.483(9)	C3	C2	1.376(11)
C6	C1	1.389(9)	C14	C15	1.406(10)
C5	C4	1.400(10)	C14	C13	1.469(10)
C4	C3	1.386(10)	C12	C13	1.339(10)
C4	C22	1.521(10)	C12	C21	1.439(10)
C7	C10	1.552(9)	C17	C16	1.425(12)
C7	С9	1.509(10)	C17	C18	1.347(11)
C7	C8	1.502(8)	C15	C16	1.367(11)
C10	С9	1.504(9)	C21	C20	1.352(10)
C10	C12	1.498(8)			

Table S11. Bond Angles for 4l.

Atom	Atom	Atom	Angle/°	Atom	Atom	Atom	Angle/°
C1	S1	C8	92.0(3)	C6	C1	S 1	113.8(6)
C5	C6	C7	126.7(6)	C6	C1	C2	120.7(6)
C1	C6	C5	119.7(6)	C2	C1	S 1	125.4(5)
C1	C6	C7	113.5(6)	C2	C3	C4	121.2(7)
C6	C5	C4	119.5(6)	C10	C9	C7	62.0(4)
C5	C4	C22	119.6(6)	C19	C14	C15	123.8(7)
C3	C4	C5	119.9(6)	C19	C14	C13	117.3(7)
C3	C4	C22	120.5(6)	C15	C14	C13	119.0(6)
C6	C7	C10	120.9(5)	C13	C12	C10	123.7(6)
C6	C7	С9	123.3(5)	C13	C12	C21	118.2(6)
C6	C7	C8	109.4(6)	C21	C12	C10	118.0(6)
С9	C7	C10	58.8(4)	C18	C17	C16	120.4(7)
C8	C7	C10	115.5(6)	C3	C2	C1	118.9(6)
C8	C7	С9	120.5(6)	C16	C15	C14	118.8(7)
С9	C10	C7	59.1(4)	C15	C16	C17	119.8(7)

C9	C10	C11	115.1(5)	01	C8	S1	123.0(5)
C12	C10	C7	116.1(5)	01	C8	C7	127.2(7)
C12	C10	C9	122.6(6)	C7	C8	S1	109.8(5)
C12	C10	C11	113.7(5)	C17	C18	C19	121.0(7)
C11	C10	C7	119.9(6)	F1	C11	C10	110.5(6)
C14	C19	C18	116.1(7)	C12	C13	C14	121.0(7)
C14	C19	C20	123.1(7)	C20	C21	C12	122.6(7)
C20	C19	C18	120.7(6)	C21	C20	C19	117.7(7)

Table S12. Hydrogen Atom Coordinates ($Å \times 10^4$) and Isotropic Displacement Parameters ($Å^2 \times 10^3$) for **41**.

x	У	Z	U(eq)
6612.46	5060	7362.47	40
5408.78	8072.19	8328.2	50
5987.7	3286.36	8082.76	46
4349.51	2564.59	8844.57	46
6685.66	5483.42	1043.72	67
9185.66	7203.49	7699.86	63
7881.06	7570.65	6510.43	63
8720.92	6411.84	6573.2	63
2484.69	7548.07	9317.14	51
8502.24	3598.01	3965.86	55
9164.98	4407.68	2031.03	61
3798.68	5872.68	2085.81	57
1565.23	2101.09	6538.72	56
1139.68	2275.96	8004.06	56
6211.99	3258.35	5759.11	53
988.92	4826.55	5907.53	59
1501.96	5587.31	3952.44	58
	x 6612.46 5408.78 5987.7 4349.51 6685.66 9185.66 7881.06 8720.92 2484.69 8502.24 9164.98 3798.68 1565.23 1139.68 6211.99 988.92 1501.96	xy6612.4650605408.788072.195987.73286.364349.512564.596685.665483.429185.667203.497881.067570.658720.926411.842484.697548.078502.243598.019164.984407.683798.685872.681565.232101.091139.682275.966211.993258.35988.924826.551501.965587.31	xyz6612.4650607362.475408.788072.198328.25987.73286.368082.764349.512564.598844.576685.665483.421043.729185.667203.497699.867881.067570.656510.438720.926411.846573.22484.697548.079317.148502.243598.013965.869164.984407.682031.033798.685872.682085.811565.232101.096538.721139.682275.968004.066211.993258.355759.11988.924826.555907.531501.965587.313952.44

Data intensity of 6^5 was collected using a 'XtaLAB AFC12 (RINC)' diffractometer at 100.00(10) K. Data collection and reduction were done by using Olex2 and the structure was solved with the ShelXS structure solution program using direct methods and refined by full-matrix least-squares on F^2 with anisotropic displacement parameters for non-H atoms using SHELX-97. Hydrogen atoms were added at their geometrically idea positions and refined isotropically. Crystal data for **6**:

⁵ Supplementary crystallographic data have been deposited at Cambridge Crystallographic Data Center (CCDC number: 2053476).

C₂₅H₂₃NO₂S, *T* = 100.00(10) K, orthorhombic, P2₁2₁2₁, *a* = 8.40850(10) Å, *b* = 9.74720(10) Å, *c* = 25.2684(3) Å, $\alpha = 90^{\circ}$, $\beta = 90^{\circ}$, $\gamma = 90^{\circ}$, *V* = 2070.98(4) Å³. Z = 4, $\rho_{calc} = 1.288 \text{ g/cm}^3$. 48728 reflections collected, 4211 [R_{int} = 0.0860, R_{sigma} = 0.0316] independent reflections, R₁ = 0.0362, wR₂ = 0.0892 (*I* > 2 σ (*I*), final), R₁ = 0.0385, wR₂ = 0.0913 (all data), GOF = 1.107, and 264 parameters.



Table S13. Crystal data and structure refinement for 6.

Identification code	6
Empirical formula	C25H23NO2S
Formula weight	401.50
Temperature/K	100.00(10)
Crystal system	orthorhombic
Space group	P212121
a/Å	8.40850(10)
b/Å	9.74720(10)
c/Å	25.2684(3)
α /°	90
β/°	90
$\gamma^{\prime \circ}$	90
Volume/Å ³	2070.98(4)
Ζ	4
$\rho_{calc}g/cm^3$	1.288
μ/mm^{-1}	1.548
F(000)	848.0
Crystal size/mm ³	$0.38 \times 0.12 \times 0.08$
Radiation	$CuK\alpha$ ($\lambda = 1.54184$)
2Θ range for data collection/°	6.996 to 149.406
Index ranges	$\text{-10} \le h \le 10, \text{-12} \le k \le 12, \text{-31} \le l \le 31$
Reflections collected	48728
Independent reflections	4211 [$R_{int} = 0.0860, R_{sigma} = 0.0316$]
Data/restraints/parameters	4211/0/264
Goodness-of-fit on F ²	1.107

Final R indexes $[I \ge 2\sigma(I)]$	$R_1 = 0.0362, wR_2 = 0.0892$
Final R indexes [all data]	$R_1 = 0.0385, wR_2 = 0.0913$
Largest diff. peak/hole / e Å ⁻³	0.19/-0.31
Flack parameter	0.012(8)

Table S14. Fractional Atomic Coordinates ($\times 10^4$) and Equivalent Isotropic Displacement Parameters ($\mathring{A}^2 \times 10^3$) for **6**. U_{eq} is defined as 1/3 of of the trace of the orthogonalised U_{IJ} tensor.

Atom	x	У	Z	U(eq)
C1	7691(3)	1430(3)	3547.6(11)	24.8(5)
C2	7482(4)	222(3)	3827.7(12)	30.9(7)
C3	7140(4)	292(3)	4362.7(12)	32.5(7)
C4	6996(3)	1546(3)	4622.7(10)	27.7(6)
C5	7225(3)	2751(3)	4330.2(10)	22.7(6)
C6	7581(3)	2717(2)	3793.4(10)	20.4(5)
C7	7830(3)	3923(2)	3424.6(9)	19.1(5)
C8	8240(3)	3313(3)	2876.3(10)	22.9(5)
C9	6301(3)	4804(2)	3335.2(9)	18.8(5)
C10	4887(3)	3939(2)	3177.6(10)	19.6(5)
C11	4471(4)	3864(3)	2642.2(10)	24.2(6)
C12	3224(4)	3033(3)	2477.8(11)	27.8(6)
C13	2372(4)	2275(3)	2845.7(12)	29.5(6)
C14	2744(3)	2371(3)	3376.9(11)	27.7(6)
C15	4002(3)	3190(3)	3542.8(11)	24.5(6)
C16	9213(3)	4892(2)	3581.0(10)	19.8(5)
C17	8734(3)	5909(2)	4008.7(9)	18.8(5)
C18	4591(3)	6478(3)	3774.1(11)	26.0(6)
C19	9911(3)	7044(2)	4113.2(10)	18.6(5)
C20	10206(3)	7446(3)	4632.2(10)	24.3(5)
C21	11212(4)	8538(3)	4743.8(10)	29.1(6)
C22	11951(4)	9240(3)	4333.6(11)	28.3(6)
C23	11661(3)	8850(2)	3813.2(10)	24.8(6)
C24	10642(3)	7768(2)	3703.3(10)	21.6(5)
C25	6591(5)	1613(3)	5201.9(11)	41.3(8)
N1	5997(3)	5597(2)	3823.3(8)	19.5(5)
01	8597(2)	3986.2(19)	2494.8(7)	27.7(4)
O2	7297(2)	6606.0(17)	3851.0(7)	19.8(4)
S 1	8101.3(9)	1504.3(7)	2864.6(3)	28.08(17)

Table S15. Anisotropic Displacement Parameters ($Å^2 \times 10^3$) for **6**. The Anisotropic displacement factor exponent takes the form: $-2\pi^2[h^2a^{*2}U_{11}+2hka^*b^*U_{12}+...]$.

Atom	U11	U ₂₂	U ₃₃	U ₂₃	U ₁₃	U ₁₂
C1	21.7(14)	17.2(11)	35.5(13)	-2.1(11)	2.9(11)	-0.5(11)
C2	31.5(17)	15.7(12)	45.7(17)	1.1(11)	3.6(14)	1.1(11)
C3	33.5(18)	19.2(12)	44.9(16)	10.9(12)	-2.3(14)	0.2(12)
C4	27.6(15)	26.6(13)	28.9(13)	8.9(11)	-3.2(11)	-3.5(13)
C5	25.3(15)	19.7(12)	23.2(12)	2.5(9)	-2.6(10)	-2.0(10)
C6	19.7(13)	15.7(12)	25.9(12)	2.0(9)	-2.6(10)	0.6(10)
C7	21.6(14)	16.2(11)	19.4(11)	-1.3(9)	2.1(10)	0.9(9)
C8	21.6(14)	22.5(12)	24.6(12)	-3.7(10)	0.1(11)	0.6(11)
C9	22.6(14)	15.5(11)	18.2(11)	-0.9(9)	0.4(10)	0.9(10)
C10	20.5(13)	13.9(10)	24.3(12)	-3.1(9)	-1.4(10)	1.7(10)
C11	28.2(15)	19.6(12)	24.7(12)	-0.5(10)	-2.1(10)	2.2(11)
C12	27.7(16)	25.6(13)	30.2(13)	-6.8(10)	-7.0(12)	3.4(11)
C13	26.2(15)	18.7(12)	43.5(15)	-7.2(11)	-5.5(13)	0.9(11)
C14	25.7(15)	18.2(12)	39.2(15)	-0.4(11)	3.3(12)	-0.7(11)
C15	25.7(15)	22.0(13)	25.8(12)	0.8(10)	0.2(11)	-0.4(10)
C16	22.2(14)	15.6(11)	21.4(11)	-0.1(9)	1.7(10)	0.4(10)
C17	20.8(13)	17.2(11)	18.4(11)	0.4(9)	0.1(9)	2.5(10)
C18	25.4(15)	22.7(12)	29.9(13)	-4.4(11)	-0.6(11)	2.3(12)
C19	18.9(14)	15.3(10)	21.5(12)	0.7(9)	-1.6(10)	2.2(9)
C20	30.0(15)	22.8(12)	19.9(12)	1.6(10)	-1.4(10)	-2.0(12)
C21	35.7(17)	28.9(13)	22.7(12)	-4.0(11)	-6.2(11)	-7.2(13)
C22	30.4(16)	19.8(12)	34.9(14)	-1.7(10)	-5.2(13)	-5.2(12)
C23	29.1(16)	18.6(12)	26.8(12)	1.8(9)	2.7(12)	-1.0(11)
C24	26.8(15)	16.3(12)	21.6(12)	-0.2(9)	-0.3(10)	0.4(11)
C25	56(2)	39.8(17)	28.5(14)	11.8(13)	-1.9(14)	-10.4(17)
N1	18.4(12)	18.2(10)	22.0(10)	-4.0(8)	0.5(9)	-1.7(8)
01	35.6(12)	27.1(9)	20.5(8)	-2.4(7)	5.7(8)	-1.0(8)
O2	19.4(9)	14.8(8)	25.2(8)	-1.7(7)	-1.7(7)	-0.3(7)
S 1	32.8(4)	17.6(3)	33.9(3)	-7.9(3)	9.2(3)	-2.9(3)

Table S16. Bond Lengths for 6.

Atom	Atom	Length/Å	Atom	Atom	Length/Å
C1	C2	1.385(4)	C10	C15	1.392(4)
C1	C6	1.403(3)	C11	C12	1.389(4)

C1	S 1	1.761(3)	C12	C13	1.387(4)
C2	C3	1.384(4)	C13	C14	1.381(4)
C3	C4	1.393(4)	C14	C15	1.390(4)
C4	C5	1.401(3)	C16	C17	1.521(3)
C4	C25	1.504(4)	C17	C19	1.508(4)
C5	C6	1.390(4)	C17	O2	1.442(3)
C6	C7	1.515(3)	C18	N1	1.467(3)
C7	C8	1.547(3)	C19	C20	1.391(4)
C7	С9	1.562(4)	C19	C24	1.396(4)
C7	C16	1.549(4)	C20	C21	1.388(4)
C8	O1	1.204(3)	C21	C22	1.389(4)
C8	S 1	1.767(3)	C22	C23	1.390(4)
C9	C10	1.510(4)	C23	C24	1.387(4)
C9	N1	1.478(3)	N1	O2	1.472(3)
C10	C11	1.399(4)			

Table S17. Bond Angles for 6.

Atom	Atom	Atom	Angle/°	Atom	Atom	Atom	Angle/°
C2	C1	C6	121.7(2)	C15	C10	C9	122.6(2)
C2	C1	S 1	124.1(2)	C15	C10	C11	118.7(2)
C6	C1	S 1	114.2(2)	C12	C11	C10	120.6(3)
C3	C2	C1	119.0(3)	C13	C12	C11	120.0(3)
C2	C3	C4	121.4(3)	C14	C13	C12	119.9(3)
C3	C4	C5	118.4(2)	C13	C14	C15	120.3(3)
C3	C4	C25	121.1(2)	C14	C15	C10	120.5(3)
C5	C4	C25	120.5(3)	C17	C16	C7	112.3(2)
C6	C5	C4	121.6(2)	C19	C17	C16	115.4(2)
C1	C6	C7	114.4(2)	O2	C17	C16	109.4(2)
C5	C6	C1	117.9(2)	O2	C17	C19	104.64(19)
C5	C6	C7	127.7(2)	C20	C19	C17	119.2(2)
C6	C7	C8	106.46(19)	C20	C19	C24	118.6(2)
C6	C7	C9	113.7(2)	C24	C19	C17	122.0(2)
C6	C7	C16	114.8(2)	C21	C20	C19	121.1(2)
C8	C7	C9	105.40(19)	C20	C21	C22	119.9(2)
C8	C7	C16	107.2(2)	C21	C22	C23	119.5(3)
C16	C7	С9	108.65(19)	C24	C23	C22	120.4(2)
C7	C8	S 1	112.58(17)	C23	C24	C19	120.5(2)

01	C8	C7	124.3(2)	C18	N1	C9	112.0(2)
01	C8	S1	123.1(2)	C18	N1	O2	102.21(18)
C10	C9	C7	112.28(19)	O2	N1	C9	105.14(18)
N1	C9	C7	107.99(19)	C17	O2	N1	108.69(16)
N1	C9	C10	112.1(2)	C1	S 1	C8	92.14(12)
C11	C10	С9	118.8(2)				

Table S18. Hydrogen Atom Coordinates ($Å \times 10^4$) and Isotropic Displacement Parameters ($Å^2 \times 10^3$) for 6.

Atom	x	у	Z	U(eq)
H2	7570.32	-621.95	3658.87	37
H3	7004.24	-516.78	4552.92	39
H5	7135.97	3594.59	4499.61	27
H9	6515.24	5454.51	3048.28	23
H11	5034.09	4375.56	2394.38	29
H12	2961.04	2983.78	2120.88	33
H13	1551.84	1702.72	2735.07	35
H14	2150.45	1886.35	3624.61	33
H15	4255.29	3237.84	3900.66	29
H16A	10100.68	4346.93	3706.81	24
H16B	9562.9	5392.13	3270.3	24
H17	8535.84	5410.34	4338.98	23
H18A	4648.25	6985.05	3448.94	39
H18B	4553.64	7104.84	4066.82	39
H18C	3649.53	5920.39	3774.19	39
H20	9722.47	6975.32	4908.89	29
H21	11389.73	8798.75	5092.8	35
H22	12635.26	9965.79	4406.53	34
H23	12153.57	9317.33	3537.2	30
H24	10443.3	7523.47	3353.74	26
H25A	5498.25	1878.11	5242.87	62
H25B	7262.12	2276.28	5372.48	62
H25C	6754.36	729.08	5360.04	62

5. Biological activity evaluation

Cells culture

Human cancer cell lines HCT116, SJSA-1, MCF-7 were purchased from the American Tissue Culture Collection (ATCC), and normal cell lines HUVEC were obtained from ScienCell Research

Laboratories. Cells were cultured aseptically at 37°C with 5% CO₂ using McCoy's 5A, RPMI-1640, MEM and ECM (Gibco) with 10% (v/v) FBS and 1% (v/v) penicillin-streptomycin (Sigma).

CCK-8 assay

The *in vitro* inhibitory effect on cell proliferation was measured using CCK-8 assay. Different kinds of cells were seeded in 96-well plates at a concentration of 3,000 cells/well. After 24 h incubation, compounds were added into each well at 50 µM. Seventy-two hours later, the old medium was replaced with fresh medium containing 10% CCK-8, and the cells were incubated for additional 4 h. The optical density was measured at 450 nm and 620 nm (reference wavelength) using a microplate reader (Berthold). If the inhibition rate of compounds on cells is higher than 70%, the IC₅₀ value was determined by testing the inhibitory effects of the compound with 8 gradient-dilution concentrations. The IC₅₀ value was calculated using GraphPad software.

6. NMR spectra
































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															f1	(ppn	n)																

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T ' T ____ 50 -60 -70 -80 -90 -100 -110 -120 -130 -140 -150 -160 -170 -180 -190 -200 -210 -220 -230 -240 -250 -260 -270 -280 -290 -300 -310 -320 -330 -340 -35 f1 (ppm)

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50 -60 -70 -80 -90 -100 -110 -120 -130 -140 -150 -160 -170 -180 -190 -200 -210 -220 -230 -240 -250 -260 -270 -280 -290 -300 -310 -320 -330 -340 -35 fl (ppm)







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T . L ____ 60 -70 -80 -90 -100 -110 -120 -130 -140 -150 -160 -170 -180 -190 -200 -210 -220 -230 -240 -250 -260 -270 -280 -290 -300 -310 -320 -330 -340 -350 -36 f1 (ppm)

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60 -70 -80 -90 -100 -110 -120 -130 -140 -150 -160 -170 -180 -190 -200 -210 -220 -230 -240 -250 -260 -270 -280 -290 -300 -310 -320 -330 -340 -350 -36 fl (ppm)



















10 0 -10 -20 -30 -40 -50 -60 -70 -80 -90 -100 -110 -120 -130 -140 -150 -160 -170 -180 -190 -200 -210 fl (ppm)








10 0 -10 -20 -30 -40 -50 -60 -70 -80 -90 -100 -110 -120 -130 -140 -150 -160 -170 -180 -190 -200 -210 fl (ppm)















































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Analysis Report

<Sample Information>

Sample Name Sample ID Data Filename Method Filename	: pbw-pc-79-asy-odh-99-1-1.0ml : : pbw-pc-79-asy-odh-99-1-1.0ml1.lcd : 1.0ml-254-230.lcm		
Vial #	: : 1-1 : 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	2020/8/28 10:09:19 2020/8/28 10:24:32	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Analysis Report

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Sample Name	: pbw-pb-142-2-rac-odh-99-1-	1.0ml	
Data Filename Method Filename	: pbw-pb-142-2-rac-odh-99-1-	1.0ml1.lcd	
Batch Filename	:		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/9/10 20:54:51	Acquired by	: System Administrato
Date Processed	: 2020/9/10 21:07:42	Processed by	: System Administrato

<Chromatogram>



Detecto	or A Chann	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	6.967	4294000	461567	50.016
2	9.372	4291234	348494	49.984
Total		8585234	810061	

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D:\Data\PBW\pbw-pb-142-2-rac-odh-99-1-1.0ml1.lcd

2020/9/10 21:22:51 Page 1 / 1

Analysis Report

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: pbw-pc-56-1-asy-ojh-99-1-1.0ml ; ; pbw-pc-56-1-asy-ojh-99-1-1.0ml1.lcd ; 1.0ml-254-230.lcm				
1-1	Sample Type	: Unknown		
: 200L : 2020/8/31 14:22:01 : 2020/8/31 14:48:13	Acquired by Processed by	: System Administrator : System Administrator		
	: pbw-pc-56-1-asy-ojh-99-1-1.0ml pbw-pc-56-1-asy-ojh-99-1-1.0ml1.lcc 1.0ml-254-230.lcm 1-1 20 uL 2020/8/31 14:22:01 2020/8/31 14:48:13	pbw-pc-56-1-asy-ojh-99-1-1.0ml pbw-pc-56-1-asy-ojh-99-1-1.0ml1.lcd 1.0ml-254-230.lcm 1-1 Sample Type 20 uL 2020/8/31 14:22:01 Acquired by 2020/8/31 14:48:13 Processed by		

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area 1 10.236 6577362 2 12.039 141437 Total 6718799 Height 292807 6237 299045 Conc. 97.895 2.105

Analysis Report

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Sample Name	: pbw-pb-148-1-rac-ojh-99-1-1	.0ml	
Data Filename	: pbw-pb-148-1-rac-ojh-99-1-1	.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Vial #	1-1	Sample Type	: Unknown
Injection Volume	: 20 uL	. ,	
Date Acquired	: 2020/8/19 14:36:06	Acquired by	: System Administrator
Date Processed	: 2020/8/31 14:58:12	Processed by	: System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	10.050	6973961	290110	50.089
2	11.872	6949276	227272	49.911
Total		13923237	517382	

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D:\Data\PBW\catalyst2\pbw-pb-148-1-rac-ojh-99-1-1.0ml1.lcd

2020/8/31 10:22:40 Page 1 / 1

Analysis Report

<Sample Information>

Sample Name	: pbw-pc-58-1-asy-ozh-99-1-1	.0ml		
Data Filename	: pbw-pc-58-1-asy-ozh-99-1-1.0ml1.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Batch Filename				
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	· 20 ul	- Pression of the		
Date Acquired	2020/8/31 0:37:48	Acquired by	· System Administrator	
Date Processed	2020/9/31 10:11:56	Processed by	: System Administrator	
Date Flocesseu	. 2020/0/31 10.11.00	Frocessed by	. System Auministrator	

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area 1 7.057 305185 2 10.082 5639981 Total 5945166 Height 33360 365414 398774 Conc. 5.133 94.867

Analysis Report

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Sample Name	: pbw-pc-7-7-rac-ozh-99-1-1.0ml		
Data Filename	: pbw-pc-7-7-rac-ozh-99-1-1.0ml1.lcd		
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/8/20 20:48:35	Acquired by	: System Administrator
Date Processed	: 2020/8/31 10:21:42	Processed by	: System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	7.067	5265521	513327	49.903
2	10.190	5285901	340909	50.097
Total		10551422	854236	

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2020/8/31 21:13:57 Page 1 / 1

Analysis Report

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Sample Name	: pbw-pc-56-5-asy-ozh-85-15-1 0ml			
Sample ID				
Sample ID				
Data Filename	: pbw-pc-56-5-asy-ozh-85-15-1.0ml1.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Batch Filename	•			
Vial #	1.1	Sample Type	Linknown	
via #	00.1	Sample Type	. OHKHOWH	
Injection volume	: 20 UL			
Date Acquired	: 2020/8/31 20:52:17	Acquired by	: System Administrator	
Date Processed	· 2020/8/31 21:09:55	Processed by	 System Administrator 	
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<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	6.321	157584	19656	2.349
2	10.059	6551991	388154	97.651
Total		6709575	407809	

Analysis Report

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Sample Name	: pbw-pc-8-4-rac-ozh-85-15-1	.0ml	
Sample ID			
Data Filename	: pbw-pc-8-4-rac-ozh-85-15-1	.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	:		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/8/21 20:24:34	Acquired by	: System Administrator
Date Processed	: 2020/8/31 21:13:01	Processed by	: System Administrator

<Chromatogram>



Detecto	or A Channe	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	6.309	10920056	1205824	50.023
2	9.971	10910206	659827	49.977
Total		21830262	1865651	

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2020/9/10 22:12:25 Page 1 / 1

2020/8/21 18:20:50 Page 1 / 1

Analysis Report

<Sample Information>

Sample Name Sample ID Data Filename Method Filename	: pbw-pc-56-4-asy-ozh-99-1-1.0m : : pbw-pc-56-4-asy-ozh-99-1-1.0m : 1.0ml-254-230.lcm	I I1.lcd	
Vial #	: 1-1 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	: 2020/8/31 10:07:47 : 2020/8/31 10:40:56	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Detector A Channel 2 230nm

4						
I	Peak#	Ret. Time	Area	Height	Conc.	
[1	9.734	122973	10265	1.996	
Γ	2	13.035	6036547	332918	98.004	
Γ	Total		6159520	343184		

Analysis Report

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Sample Name	pbw-pc-7-8-rac-ozh-99-1-1.0ml		
Data Filename	: pbw-pc-7-8-rac-ozh-99-1-1.0ml1.lcd		
Method Filename	: 1.0ml-254-230.lcm		
Vial #	1_1	Sample Type	· Linknown
Injection Volume	20 uL	Campie Type	· Onknown
Date Acquired	: 2020/8/20 21:03:06	Acquired by	: System Administrator
Date Processed	: 2020/8/21 9:51:51	Processed by	: System Administrator

<Chromatogram>



Detect	or A Channe	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	9.778	3258068	246244	50.071
2	13.391	3248775	177952	49.929
Total		6506843	424196	

D:\Data\PBW\catalyst2\pbw-pc-56-4-asy-ozh-99-1-1.0ml1.lcd

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2020/8/31 18:06:02 Page 1 / 1

Analysis Report

<Sample Information>

Sample Name	: pbw-pc-56-3-asy-ozh-97-3-	1.0ml	
Data Filename	: pbw-pc-56-3-asy-ozh-97-3-	1.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/8/31 17:15:59	Acquired by	: System Administrator
Date Processed	: 2020/8/31 17:38:02	Processed by	: System Administrator

<Chromatogram>



Analysis Report

Sample Name Sample ID Data Filename Method Filename	pbw-pb-148-4-rac-ozh-97-3 pbw-pb-148-4-rac-ozh-97-3 1.0ml-254-230.lcm	-1.0ml -1.0ml1.lcd	
/ial #	: : 1-1 : 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	: 2020/8/31 17:45:01 : 2020/8/31 18:04:47	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>

<Sample Information>



Peak#	Ret. Time	Area	Height	Conc.
1	7.071	2349178	236673	49.937
2	9.767	2355146	169640	50.063
Total		4704324	406313	

Detect	or A Chann	el 1 254nm	11 1 1 1 1	
Peak#	Ret. Time	Area	Height	Conc.
1	7.073	58483	6838	1.458
2	9.759	3951631	284851	98.542
Total		4010114	291689	

D:\Data\PBW\pbw-pb-148-4-rac-ozh-97-3-1.0ml1.lcd

2020/9/10 21:41:33 Page 1 / 1

2020/8/31 12:02:32 Page 1 / 1

Analysis Report

<Sample Information>

•			
Sample Name Sample ID Data Filename Method Filename	: pbw-pc-56-2-asy-ozh-99-1-1.0m : pbw-pc-56-2-asy-ozh-99-1-1.0m : 1.0ml-254-230.lcm	I I1.lcd	
Vial #	1-1	Sample Type	: Unknown
Date Acquired Date Processed	: 20 uL : 2020/8/31 10:49:06 : 2020/8/31 11:17:18	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Detector A Channel 1 254nm

Delect				
Peak#	Ret. Time	Area	Height	Conc.
1	11.585	77932	5300	1.628
2	17.733	4709467	181359	98.372
Total		4787400	186659	

Analysis Report

<Sample Information>

Sample Name	: pbw-pc-148-3-rac-ozh-99-1	1-1.0ml				
Data Filename	pbw-pc-148-3-rac-ozh-99-1-1.0ml1.lcd					
Method Filename	: 1.0ml-254-230.lcm					
Batch Filename	:					
/ial #	: 1-1	Sample Type	: Unknown			
niection Volume	: 20 ul					
Date Acquired	2020/8/31 11:30:29	Acquired by	· System Administrator			
Date Processed	2020/8/31 12:01:41	Processed by	· System Administrator			
Juic I loooood	. LOLOIOIOI IL.OI.III		. Of otomin tarihinoti ator			

<Chromatogram>



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Peak#	Ret. Time	Area	Height	Conc.
1	11.622	3461924	204312	49.964
2	17.891	3466854	131210	50.036
Total		6928779	335522	

D:\Data\PBW\catalyst2\pbw-pc-56-2-asy-ozh-99-1-1.0ml1.lcd

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2020/8/21 16:28:14 Page 1 / 1

Analysis Report

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Sample Name	pbw-pc-128-asy-odh-99-1-1.0ml		
Data Filename Method Filename Batch Filename	pbw-pc-128-asy-odh-99-1-1.0ml1.lcd 1.0ml-254-230.lcm		
Vial #	: 1-1 : 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	: 2020/10/14 16:13:28 : 2020/10/14 16:31:13	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	8.876	7989373	634054	98.796
2	10.034	97387	8275	1.204
Total		8086760	642329	

Analysis Report

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Sample Name	pbw-pc-8-2-rac-odh-99-1-1.0ml		
Data Filename	: pbw-pc-8-2-rac-odh-99-1-1.0ml1.lcd		
Batch Filename	: 1.0111-234-230.1011		
Vial # Injection Volume	: 1-1 : 20 uL	Sample Type	: Unknown
Date Acquired	2020/8/15 17:03:43	Acquired by	: System Administrator

<Chromatogram>



Detect	or A Channe	1 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	8.407	3251723	289076	49.881
2	9.612	3267291	251270	50.119
Total		6519014	540345	

D:\Data\PBW\pbw-pc-8-2-rac-odh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-128-asy-odh-99-1-1.0ml1.lcd

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Sample Name Sample ID Data Filename Method Filename	: pbw-pc-60-asy-ozh-99-1-1.0ml : : pbw-pc-60-asy-ozh-99-1-1.0ml1.lcd : 1.0ml-254-230.lcm		
Batch Filename Vial #	: 1-1 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	: 2020/8/31 10:27:28 : 2020/8/31 11:35:33	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area Height 4817 249306 254124 Conc. 1.642 98.358 78175 4682397 4760571 1 2 11.890 13.343 Total

Analysis Report

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Sample Name	pbw-pc-8-1-rac-ozh-99-1-1.0ml		
Data Filename	pbw-pc-8-1-rac-ozh-99-1-1.0ml2.lcd		
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	. 1 1	Sample Tupe	Linknown
Injection Volume	: 20 uL	Sample Type	. Onknown
Date Acquired	: 2020/8/31 11:12:19	Acquired by	: System Administrator
Date Processed	: 2020/8/31 11:42:46	Processed by	: System Administrator

<Chromatogram>



Detect	Detector A Channel 2 230nm			
Peak#	Ret. Time	Area	Height	Conc.
1	11.880	6002300	359098	49.272
2	13.353	6179586	335332	50.728
Total		12181886	694430	

D:\Data\PBW\pbw-pc-8-1-rac-ozh-99-1-1.0ml2.lcd

D:\Data\PBW\catalyst2\pbw-pc-60-asy-ozh-99-1-1.0ml1.lcd

<Sample Information>

•			
Sample Name	: pbw-pc-58-2-asy-ozh-97-3-1.0ml		
Sample ID			
Data Filename	: pbw-pc-58-2-asy-ozh-97-3-1.0ml1.	lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	1-1	Sample Type	Unknown
Injection Volume	· 20 ul	earrible ()pe	
Date Acquired	: 2020/8/31 16:41:56	Acquired by	· System Administrator
Date Acquired	2020/0/31 10.41.00	Acquired by	. System Administrator
Date Processed	2020/8/31 17:09:22	Processed by	System Administrator

<Chromatogram>



Detector A Channel 1 254nm Peak#Ret. Time Area Height Conc. 1 7.467 233639 24016 3.677 2 9.626 6120202 431013 96.323 Total 6353841 455029

Analysis Report

<sample inform<="" th=""><th>nation></th><th></th><th></th></sample>	nation>		
Sample Name	pbw-pc-7-3-rac-ozh-97-3-1.0ml		
Data Filename	pbw-pc-7-3-rac-ozh-97-3-1.0ml1.lcd		
Batch Filename	: 1.0ml-254-230.lcm		
Vial #	: 1-1	Sample Type	: Unknown
Date Acquired Date Processed	: 20 0L : 2020/8/31 17:31:02 : 2020/8/31 17:50:18	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



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Detector A Channel 1 254nm				1
Peak#	Ret. Time	Area	Height	Conc.
1	7.688	5164227	469778	50.045
2	10.043	5154894	355087	49.955
Total		10319121	824865	

D:\Data\PBW\catalyst2\pbw-pc-58-2-asy-ozh-97-3-1.0ml1.lcd

D:\Data\PBW\pbw-pc-7-3-rac-ozh-97-3-1.0ml1.lcd

<Sample Information>

Sample Name	pbw-pc-58-3-asy-odh-99-	1-1.0ml	
Sample ID			
Data Filename	: pbw-pc-58-3-asy-odn-99-	1-1.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 ul	1 11	
Date Acquired	2020/9/1 9:01:37	Acquired by	 System Administrator
Date Processed	2020/0/1 0:57:00	Processed by	: System Administrator
Date Flocesseu	. 2020/9/1 9.57.09	Frocessed by	. System Authinistrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	7.466	3666182	362797	94.306
2	9.592	221375	20271	5.694
Total		3887557	383068	

Analysis Report

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Sample Name Sample ID Data Filename Method Filename	: pbw-pc-45-1-rac-odh-99-1-1.0ml : : pbw-pc-45-1-rac-odh-99-1-1.0ml4.lcc : 1.0ml-254-230.lcm	i	
Batch Filename Vial #	1-1 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	: 2020/9/1 10:34:34 : 2020/9/1 11:13:50	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Detect	Detector A Channel 1 254nm			
Peak#	Ret. Time	Area	Height	Conc.
1	7.426	3992149	407129	51.975
2	9.542	3688721	296585	48.025
Total		7680871	703713	

D:\Data\PBW\catalyst2\pbw-pc-58-3-asy-odh-99-1-1.0ml1.lcd

D:\Data\PBW\catalyst2\pbw-pc-45-1-rac-odh-99-1-1.0ml4.lcd

2020/9/9 10:41:43 Page 1 / 1

Analysis Report

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Sample Name	pbw-pc-74-asy-odh-99-1-1.0ml			
Data Filename	pbw-pc-74-asy-odh-99-1-1.0ml1.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Batch Filename				
Vial #	1-1	Sample Type	· Unknown	
Injection Volume	20 ul	oumpie type	. Onitalouni	
Date Acquired	2020/0/0 10:22:33	Acquired by	· System Administrator	
Date Record	2020/0/0 10:20:14	Proceed by	System Administrator	
Date Frocessed	. 2020/9/9 10.39.14	Frocessed by	. System Administrator	

<Chromatogram>



Analysis Report

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Sample Name	pbw-pc-69-rac-odh-99-1-1.0ml		
Data Filename	pbw-pc-69-rac-odh-99-1-1.0ml1.lcd		
Method Filename Batch Filename	: 1.0ml-254-230.lcm		
Vial #	1-1	Sample Type	: Unknown
Date Acquired	: 2020/9/9 10:08:44	Acquired by	: System Administrator
Date Processed	: 2020/9/9 10:36:05	Processed by	: System Administrator

<Chromatogram>



SFeak Table

	Detect	or A Chann	el 1 254nm		
1	Peak#	Ret. Time	Area	Height	Conc.
1	1	6.683	1870086	203090	50.449
1	2	8.804	1836831	151585	49.551
i	Total		3706917	354675	

D:\Data\PBW\pbw-pc-74-asy-odh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-69-rac-odh-99-1-1.0ml1.lcd

2020/9/15 11:21:05 Page 1 / 1

Analysis Report

<sample information=""></sample>				
Sample Name Sample ID	: pbw-pc-82-asy-odh-99-1-1.0ml			
Data Filename	: pbw-pc-82-asy-odh-99-1-1.0ml1.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Batch Filename				
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL			
Date Acquired	: 2020/9/15 10:56:17	Acquired by	: System Administrator	
Date Processed	: 2020/9/15 11:24:24	Processed by	: System Administrator	

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area 1 11.957 5371595 2 14.892 538552 Total 5910147 Height 321369 29338 350707 Conc. 90.888 9.112

Analysis Report

<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>		
Sample Name Sample ID Data Filename Method Filename	: pbw-pc-81-rac-odh-99-1-1.0ml : : pbw-pc-81-rac-odh-99-1-1.0ml1.lcd : 1.0ml-254-230.lcm		
Batch Filename Vial #	1-1	Sample Type	: Unknown
Date Acquired Date Processed	: 20 UL : 2020/9/15 10:34:50 : 2020/9/15 11:19:59	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Detect	or A Channe	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	12.124	5177085	306477	50.812
2	14.929	5011551	236906	49.188
Total		10188636	543384	

D:\Data\PBW\pbw-pc-81-rac-odh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-82-asy-odh-99-1-1.0ml1.lcd

<sample information=""></sample>					
Sample Name Sample ID Data Filename	: pbw-pc-103-asy-odh-99-1-1.0ml : : pbw-pc-103-asy-odh-99-1-1.0ml1.I	cd			
Method Filename	: 1.0ml-254-230.lcm				
Batch Filename	:				
Vial #	: 1-1	Sample Type	: Unknown		
Injection Volume	: 20 uL				
Date Acquired	: 2020/9/25 10:28:38	Acquired by	: System Administrator		
Date Processed	: 2020/9/25 10:52:32	Processed by	: System Administrator		

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	7.172	9595667	1000371	92.221
2	11.509	809462	59334	7.779
Total		10405129	1059704	

Analysis Report

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100	- 32	10.02		

Sample Name	: pbw-pc-102-rac-odh-99-1-1.0ml		
Data Filename	: pbw-pc-102-rac-odh-99-1-1.0ml1.lcd		
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	:		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/9/25 9:20:58	Acquired by	: System Administrator
Date Processed	: 2020/9/25 11:01:18	Processed by	: System Administrator

<Chromatogram>



SFeak Table

Detect	or A Channe	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	7.201	9611064	1013443	51.157
2	11.554	9176405	570698	48.843
Total		18787469	1584140	

D:\Data\PBW\pbw-pc-102-rac-odh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-103-asy-odh-99-1-1.0ml1.lcd

2020/9/9 10:33:18 Page 1 / 1

Analysis Report

<Sample Information>

Sample Name Sample ID	pbw-pc-72-asy-odh-99-1-1.0ml		
Data Filename	: pbw-pc-72-asy-odn-99-1-1.0m11.icd		
Method Filename	. 1.0mi-254-230.icm		
Batch Filename			
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/9/9 9:53:27	Acquired by	: System Administrator
Date Processed	: 2020/9/9 10:19:21	Processed by	: System Administrator

<Chromatogram>



Detector A Channel 2 230nm Peak#/ Ret. Time Area Height Conc. 1 8.849 5417426 455955 94.183 2 12.574 334625 22287 5.817 Total 5752051 478242 5.817

Analysis Report

<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>		
Sample Name	: pbw-pc-61-rac-odh-99-1-1.0ml		
Data Filename	pbw-pc-61-rac-odh-99-1-1.0ml1.lcd		
Batch Filename	1.0mi-254-230.icm		
Vial # Injection Volume	: 1-1 : 20 uL	Sample Type	: Unknown
Date Acquired	2020/9/8 10:09:26	Acquired by	: System Administrator
Date Frocessed	. 2020/9/0 15.05.40	Flocessed by	. System Auministrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	8.806	5050769	429698	50.112
2	12.327	5028171	301135	49.888
Total		10078940	730834	

D:\Data\PBW\pbw-pc-61-rac-odh-99-1-1.0ml1.lcd

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Sample Name Sample ID	pbw-pc-58-5-asy-odh-99-	1-1.0ml				
Data Filename	: pbw-pc-58-5-asy-odh-99-1-1.0ml1.lcd : 1.0ml-254-230.lcm					
Method Filename						
Batch Filename	1					
Vial #	: 1-1	Sample Type	: Unknown			
Injection Volume	: 20 uL					
Date Acquired	: 2020/9/1 8:47:11	Acquired by	: System Administrator			
Date Processed	: 2020/9/1 9:10:42	Processed by	: System Administrator			

<Chromatogram>



Detector A Channel 1 254nm Peak#Ret. Time Area Height Conc. 1 8.861 3147385 254840 92.376 2 10.305 259778 21824 7.624 Total 3407163 276664

Analysis Report

<sample information=""></sample>								
Sample Name	pbw-pc-7-1-rac-odh-99-1-1.0ml							
Data Filename	pbw-pc-7-1-rac-odh-99-1-1.0ml1.lcd							
Method Filename	: 1.0ml-254-230.lcm							
Batch Filename		Comple Ture	i Universita					
Injection Volume	20 ul	Sample Type	Unknown					
Date Acquired	2020/8/17 9:59:30	Acquired by	: System Administrator					
Date Processed	: 2020/9/1 9:21:18	Processed by	: System Administrator					

<Chromatogram>



Detector A Channel 4 054

Detect	lor A Channel 1 254nm				
Peak#	Ret. Time	Area	Height	Conc.	
1	8.409	1563634	136458	49.921	
2	9.731	1568610	119571	50.079	
Total		3132244	256029		

D:\Data\PBW\pbw-pc-7-1-rac-odh-99-1-1.0ml1.lcd

D:\Data\PBW\catalyst2\pbw-pc-58-5-asy-odh-99-1-1.0ml1.lcd
<Sample Information>

Sample Name	: pbw-pc-58-4-asy-adh-99-1-1	.0ml		
Data Filename Method Filename	: pbw-pc-58-4-asy-adh-99-1-1.0ml2.lcd : 1.0ml-254-230.lcm			
Batch Filename Vial #	: 1-1 20 ul	Sample Type	: Unknown	
Date Acquired Date Processed	: 2020/8/31 16:10:52 : 2020/8/31 16:48:06	Acquired by Processed by	: System Administrator : System Administrator	

<Chromatogram>



Detector A Channel 2 230nm Peak#Ret. Time Area Height Conc. 1 10.723 3724287 246009 99.472 2 14.941 19779 1080 0.528 Total 3744066 247089

Analysis Report

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Sample Name	: pbw-pc-7-2-2-rac-adh-99-1-1	.0ml		
Data Filename	pbw-pc-7-2-2-rac-adh-99-1-1.0ml3.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Batch Filename				
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL			
Date Acquired	: 2020/8/31 15:50:16	Acquired by	: System Administrator	
Date Processed	: 2020/8/31 16:14:45	Processed by	: System Administrator	

<Chromatogram>



Detector & Channel 2 230nm				
Ret. Time	Area	Height	Conc.	
10.759	2638454	175311	50,198	
14.848	2617591	85368	49.802	
	5256045	260679		
	or A Chann Ret. Time 10.759 14.848	or A Channel 2 230nm Ret. Time Area 10.759 2638454 14.848 2617591 5256045	A Channel 2 230nm Ret. Time Area Height 10.759 2638454 175311 14.848 2617591 85368 5256045 260679	

D:\Data\PBW\catalyst2\pbw-pc-58-4-asy-adh-99-1-1.0ml2.lcd

D:\Data\PBW\pbw-pc-7-2-2-rac-adh-99-1-1.0ml3.lcd

2020/9/23 16:05:48 Page 1 / 1

Analysis Report

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Sample Name Sample ID Data Filename Method Filename Batch Filename	pbw-pc-94-1B-asy-ozh-99 pbw-pc-94-1B-asy-ozh-99 1.0ml-254-230.lcm	9-1-1.0ml 9-1-1.0ml1.lcd	
Vial #	1-1	Sample Type	: Unknown
Injection Volume Date Acquired Date Processed	: 20 uL : 2020/9/23 14:44:50 : 2020/9/23 15:56:23	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Analysis Report

<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>			
Sample Name	pbw-pc-92-1B-rac-ozh-99-1-1	.0ml		
Data Filename	pbw-pc-92-1B-rac-ozh-99-1-1.0ml2.lcd			
Batch Filename	: 1.0111-234-230.1011			
Vial #	: 1-1	Sample Type	: Unknown	
Date Acquired	: 2020/9/23 15:27:08	Acquired by	: System Administrator	
Date Processed	: 2020/9/23 16:03:44	Processed by	: System Administrator	

<Chromatogram>



Detect	or A Channe	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	12.251	4265057	167657	50.314
2	16.253	4211741	165569	49.686
Total		8476798	333226	

D:\Data\PBW\pbw-pc-94-1B-asy-ozh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-92-1B-rac-ozh-99-1-1.0ml2.lcd

2020/9/23 16:22:57 Page 1 / 1

Analysis Report

<Sample Information>

Sample Name Sample ID Data Filename Method Filename	pbw-pc-94-2B-asy-ozh-99-1-1.0ml pbw-pc-94-2B-asy-ozh-99-1-1.0ml1.lcd		
Batch Filename Vial #	: 1-1	Sample Type	: Unknown
Injection Volume Date Acquired Date Processed	: 20 uL : 2020/9/23 15:05:24 : 2020/9/23 16:30:36	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area 1 10.763 59630 2 16.944 4902022 Conc. 1.202 98.798 Height 2821 186088 Total 4961652 188909

Analysis Report

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Sample Name Sample ID	: pbw-pc-92-2B-rac-ozh-99-1	1-1.0ml		
Method Eilename	: 1 0ml 254 220 lom	1-1.01112.1Cu		
Reteb Fileneme	. 1.0111-234-230.1011			
batch Fliename		101		
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL			
Date Acquired	· 2020/0/23 15:46:48	Acquired by	· System Administrator	
Date Acquireu	2020/0/20 10:40:40	Acquired by	. Oystern Administrator	
Date Processed	: 2020/9/23 16:21:26	Processed by	: System Administrator	

<Chromatogram>



Detect Peak#	Ret. Time	Area	Height	Conc.
1	10.644	3519845	188393	49.560
2	16.799	3582360	137070	50.440
Total		7102205	325464	

D:\Data\PBW\pbw-pc-92-2B-rac-ozh-99-1-1.0ml2.lcd

D:\Data\PBW\pbw-pc-94-2B-asy-ozh-99-1-1.0ml1.lcd

2020/10/30 17:05:04 Page 1 / 1

2020/10/7 21:10:04 Page 1 / 1

Analysis Report

<Sample Information>

-					
Sample Name	: pbw-pc-124-1-asy-ozh-99-1-1.0ml				
Data Filename	pbw-pc-124-1-asy-ozh-99-1-1 0ml1 lcd				
Method Filename	: 1.0ml-254-230.lcm				
Batch Filename	1.1	Sample Type	Linknown		
Injection Volume	: 20 uL	Sample Type	. Onknown		
Date Acquired	: 2020/10/7 17:42:16	Acquired by	: System Administrator		
Date Processed	: 2020/10/7 21:00:21	Processed by	: System Administrator		

<Chromatogram>



	Ana	lysis	Report
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<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>				
Sample Name Sample ID	: pbw-pc-118-1-rac-ozh-99-1-1	.0ml			
Data Filename	: pbw-pc-118-1-rac-ozh-99-1-1.0ml4.lcd				
Method Filename	: 1.0ml-254-230.lcm				
Batch Filename					
Vial #	: 1-1	Sample Type	: Unknown		
Injection Volume	: 20 uL				
Date Acquired	: 2020/10/7 20:37:34	Acquired by	: System Administrator		
Date Processed	: 2020/10/7 21:02:34	Processed by	: System Administrator		

<Chromatogram>



Detector A Channel 2 230nm

Deleci	OF A GHAIIII	er z z 301111		
Peak#	Ret. Time	Area	Height	Conc.
1	12.427	2322492	121166	50.50
2	14.573	2276405	78827	49.499
Total		4598898	199993	

D:\Data\PBW\catalyst2\pbw-pc-124-1-asy-ozh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-118-1-rac-ozh-99-1-1.0ml4.lcd

2020/9/23 20:04:51 Page 1 / 1

Analysis Report

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Sample Name	pbw-pc-94-5-asy-oxh-99-1	-1.0ml		
Data Filename	pbw-pc-94-5-asy-oxh-99-1-1.0ml1.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Batch Eilename				
Vial #	1_1	Sample Type	Linknown	
Injection Volume	20.01	Cample Type	. Onknown	
injection volume	. 20 UL	A second second beau	O stars Administration	
Date Acquired	2020/9/23 17:37:26	Acquired by	: System Administrator	
Date Processed	: 2020/9/23 20:02:07	Processed by	: System Administrator	

<Chromatogram>



Analysis Report

<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>				
Sample Name	pbw-pc-92-5-rac-oxh-99-1-1.0ml				
Data Filename	: pbw-pc-92-5-rac-oxh-99-1-1.0ml1.lcd				
Method Filename	: 1.0ml-254-230.lcm				
Vial #	1-1	Sample Type	Unknown		
Injection Volume	: 20 uL	campie ()pe			
Date Acquired	: 2020/9/23 16:58:47	Acquired by	: System Administrator		
Date Processed	: 2020/9/23 20:03:55	Processed by	: System Administrator		

<Chromatogram>



Detect	or A Channe	el 1 254nm		
Peak#	Ret. Time	Area	Height	Conc.
1	10.138	6003866	479733	49.989
2	11.464	6006438	418840	50.011
Total		12010304	898573	

D:\Data\PBW\pbw-pc-94-5-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-92-5-rac-oxh-99-1-1.0ml1.lcd

<Sample Information>

Sample Name	: pbw-pc-122-5-asy-oxh-99-1	1-1.0ml		
Sample ID Data Filename	nbw-nc-122-5-asy-ovh-99-1-1 0ml1 lcd			
Method Filename	1.0ml-254-230.lcm	1-1.01111.100		
Batch Filename	:			
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL			
Date Acquired	: 2020/10/7 14:55:55	Acquired by	: System Administrator	
Date Processed	: 2020/10/7 15:39:16	Processed by	: System Administrator	

<Chromatogram>



Petector A Channel 2 230nm Peak#Ret. Time Area Height Conc. 1 12.054 9597110 632769 96.839 2 13.538 313219 20417 3.161 Total 9910329 653185 5

Analysis Report

<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>				
Sample Name Sample ID	: pbw-pc-113-5-rac-oxh-99-1-1.0	ml			
Data Filename	: pbw-pc-113-5-rac-oxh-99-1-1.0ml1.lcd				
Batch Filename	: 1.0111-234-230.1011				
Vial #	: 1-1	Sample Type	: Unknown		
Injection Volume	: 20 uL		_		
Date Acquired	: 2020/10/6 16:41:14	Acquired by	: System Administrator		
Date Processed	: 2020/10/7 15:39:43	Processed by	: System Administrator		

<Chromatogram>



Detect	Detector A Channel 2 230nm					
Peak#	Ret. Time	Area	Height	Conc.		
1	11.652	12132553	841938	50.384		
2	12.980	11947650	748561	49.616		
Tota	1	24080204	1590498			

D:\Data\PBW\catalyst2\pbw-pc-122-5-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-113-5-rac-oxh-99-1-1.0ml1.lcd

2020/10/7 15:50:14 Page 1 / 1

Analysis Report

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Sample Name Sample ID Data Filename Method Filename	pbw-pc-122-6-asy-oxh-99-1-1.0ml pbw-pc-122-6-asy-oxh-99-1-1.0ml1.lcd 1.0ml-254-230.lcm			
Vial #	1-1	Sample Type	: Unknown	
Date Acquired Date Processed	: 2020/10/7 15:13:57 : 2020/10/7 15:48:02	Acquired by Processed by	: System Administrator : System Administrator	

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area Height Conc. 1 12.865 18297446 1096722 97.350 2 13.674 489050 33443 2.650 Total 18795496 1130165

Analysis Report

<sample inform<="" th=""><th>mation></th><th>•</th><th></th></sample>	mation>	•	
Sample Name	: pbw-pc-113-6-rac-oxh-99-1-1	.0ml	
Data Filename	pbw-pc-113-6-rac-oxh-99-1-1	.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	÷		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/10/6 16:58:04	Acquired by	: System Administrator
Date Processed	: 2020/10/7 15:48:43	Processed by	: System Administrator

<Chromatogram>



Delect				
Peak#	Ret. Time	Area	Height	Conc.
1	12.518	10868733	692620	50.088
2	13.242	10830755	670194	49.912
Total		21699488	1362814	

D:\Data\PBW\catalyst2\pbw-pc-122-6-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-113-6-rac-oxh-99-1-1.0ml1.lcd

<samp< th=""><th>le</th><th>Inf</th><th>orm</th><th>ati</th><th>0</th><th>n></th></samp<>	le	Inf	orm	ati	0	n>
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Sample Name Sample ID Data Filename Method Filename	pbw-pc-124-2-asy-ozh-99- pbw-pc-124-2-asy-ozh-99- 1.0ml-254-230.lcm	1-1.0ml 1-1.0ml1.lcd	
Vial #	: 1-1 : 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	: 2020/10/7 17:09:28 : 2020/10/7 17:30:02	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	7.842	8726826	732452	96,134
2	10.311	350935	24786	3.866
Total		9077761	757238	

Analysis Report

<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>		
Sample Name Sample ID Data Filename Method Filename Batch Filename	pbw-pc-118-2-rac-ozh-99-1-1 pbw-pc-118-2-rac-ozh-99-1-1 1.0ml-254-230.lcm	Oml Oml1.lcd	
Vial #	: 1-1 : 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	2020/10/6 15:13:55 2020/10/7 17:33:49	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Detect	or A Chann	el 1 254nm		
Peak#	Ret. Time	Area	Height	Conc.
1	7.618	2733158	247308	50.807
2	10.079	2646293	177690	49.193
Total		5379451	424998	

D:\Data\PBW\catalyst2\pbw-pc-124-2-asy-ozh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-118-2-rac-ozh-99-1-1.0ml1.lcd

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Analysis Report

<Sample Information>

Sample Name Sample ID	: pbw-pc-73-asy-odh-99-1-1.0ml		
Data Filename	: pbw-pc-73-asy-odh-99-1-1.0ml1.lcd		
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	1		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/9/8 11:08:11	Acquired by	: System Administrator
Date Processed	: 2020/9/8 11:34:27	Processed by	: System Administrator

<Chromatogram>



Anal	/sis	Rep	ort
	, 0.0	1.0p	

<Sample Information>

Sample Name	: pbw-pc-62-1-rac-odh-99-1-	1.0ml	
Data Filename	: pbw-pc-62-1-rac-odh-99-1-	1.0ml1.lcd	
Ratch Filename	. 1.0111-254-230.1011		
/ial #	1-1	Sample Type	: Unknown
njection Volume	: 20 uL		
Date Acquired	: 2020/9/8 10:25:26	Acquired by	: System Administrator
Jate Processed	: 2020/9/8 11:35:42	Processed by	: System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	8.425	3282611	280787	50.140
2	10.477	3264229	198067	49.860
Total		6546841	478854	

D:\Data\PBW\pbw-pc-73-asy-odh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-62-1-rac-odh-99-1-1.0ml1.lcd

2020/10/30 15:45:59 Page 1 / 1

2020/10/24 16:00:30 Page 1 / 1

Analysis Report

<Sample Information>

Sample Name	: pbw-pc-73B-asy-adh-98-2-1.0ml		
Data Filename	: pbw-pc-73B-asy-adh-98-2-1.0ml1	.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/10/24 15:48:57	Acquired by	: System Administrator
Date Processed	: 2020/10/24 16:06:40	Processed by	: System Administrator
Method Filename Batch Filename Vial # Injection Volume Date Acquired Date Processed	: 1.0ml-254-230.lcm : 1-1 : 20 uL : 2020/10/24 15:48:57 : 2020/10/24 16:06:40	Sample Type Acquired by Processed by	: Unknown : System Administrato : System Administrato

<Chromatogram>



- can	Ret. Hille	Alea	riegni	CONC.
1	9.994	19926325	1202841	98.964
2	11.316	208631	9638	1.036
Total		20134956	1212479	

Analysis Report

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<sample info<="" th=""><th>rmation></th></sample>	rmation>
Sample Name	: phw.pc.62.2 rac.adb.08.2.1.0ml

Sample ID	. pbw-pc-oz-z-rac-auri-96-z-1.0					
Data Filename	: pbw-pc-62-2-rac-adh-98-2-1.0	: pbw-pc-62-2-rac-adh-98-2-1.0ml2.lcd				
Method Filename	: 1.0ml-254-230.lcm					
Batch Filename						
Vial #	: 1-1	Sample Type	: Unknown			
Injection Volume	: 20 uL					
Date Acquired	: 2020/10/24 15:33:18	Acquired by	: System Administrator			
Date Processed	: 2020/10/24 15:58:02	Processed by	: System Administrator			

<Chromatogram>



Detec	tor A Chann Ret. Time	Area	Height	Conc.
1	10.053	3108683	202288	50.973
2	11.264	2989966	117605	49.02
Tota	d .	6098649	319893	

D:\Data\PBW\catalyst2\pbw-pc-73B-asy-adh-98-2-1.0ml1.lcd

D:\Data\PBW\pbw-pc-62-2-rac-adh-98-2-1.0ml2.lcd

2020/9/23 21:30:04 Page 1 / 1

Analysis Report

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Sample Name	pbw-pc-94-3-asy-odh-99-1-	1.0ml			
Data Filename	: hhusho-94-3-asy-odb-99-1-1 0ml1 lod				
Method Filename	: 1.0ml-254-230.lcm				
Batch Filename	:				
Vial #	: 1-1	Sample Type	: Unknown		
Injection Volume	: 20 uL	A service d bu	· Custom Administrator		
Date Acquired	2020/9/23 21:12:10	Processed by	System Administrator		
Date Trocessed	. 2020/0/20 21.01.07	i rocessed by	. Oystern Administrator		

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	7.301	11756251	1139439	98.080
2	7.826	230095	21251	1.920
Total		11986346	1160691	

Analysis Report

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Sample Name	: pbw-pc-92-3-rac-odh-99-1-1	.0ml		
Data Filename	ata Filename : pbw-pc-92-3-rac-odh-99-1-1.0ml1.lcd			
Batch Filename	: 1.0mi-254-230.icm			
Vial #	: 1-1	Sample Type	: Unknown	
Date Acquired	: 20 UL : 2020/9/23 9:06:47	Acquired by	: System Administrator	
Date Processed	: 2020/9/23 21:25:16	Processed by	: System Administrator	

<Chromatogram>



Detect	or A Channe	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	7.585	6004508	554433	50.152
2	8.071	5968122	468263	49.848
Total		11972630	1022696	

D:\Data\PBW\pbw-pc-94-3-asy-odh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-92-3-rac-odh-99-1-1.0ml1.lcd

2020/9/23 17:44:11 Page 1 / 1

Analysis Report

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Sample Name	: pbw-pc-94-4-asy-oxh-99-1-	1.0ml			
Sample ID					
Data Filename	pbw-pc-94-4-asy-oxh-99-1-1.0ml1.lcd				
Method Filename	1 0ml-254-230 lcm				
Batch Filename					
Vial #	1.1	Sample Type	: Unknown		
Vial #	20.01	Sample Type	. OTIKITOWIT		
injection volume	. 20 UL		12 17 19 19 19 19 19 19 19 19 19 19 19 19 19		
Date Acquired	: 2020/9/23 17:14:03	Acquired by	: System Administrator		
Date Processed	: 2020/9/23 17:46:09	Processed by	: System Administrator		

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area 1 12.723 13705552 2 14.036 203458 Total 13909010 Height 851833 15315 867148 Conc. 98.537 1.463

Analysis Report

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Comple Name	

Sample ID	. pbw-pc-92-4-1ac-0x11-99-1-1.0	410	
Data Filename	: pbw-pc-92-4-rac-oxh-99-1-1.0	ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	1		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/9/23 16:36:00	Acquired by	: System Administrator
Date Processed	: 2020/9/23 16:57:18	Processed by	: System Administrator

<Chromatogram>



Peak# F	Ret. Time	Area	Height	Conc.
1	12.600	9508280	604345	50.038
2	13.803	9493960	545729	49.962
Total		19002240	1150074	

D:\Data\PBW\catalyst2\pbw-pc-94-4-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-92-4-rac-oxh-99-1-1.0ml1.lcd

<Sample Information>

Sample Name	: pbw-pc-122-4-asy-oxh-99-1-1.0ml			
Sample ID	. pon pone i aoj onnoo			
Sample ID				
Data Filename	: pbw-pc-122-4-asy-oxh-99-1-1.0ml1.lcd			
Method Filename	1.0ml-254-230.lcm			
Batch Eilename				
Daton nename	·	0 I T		
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL			
Date Acquired	- 2020/10/7 14-26-51	Acquired by	· System Administrator	
Date Acquireu	. 2020/10// 14.20.01	Acquired by	. System Auministrator	
Date Processed	: 2020/10/7 15:11:34	Processed by	: System Administrator	

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	22.032	36504545	1087529	96.348
2	23.794	1383852	51214	3.652
Total		37888397	1138743	

Analysis Report

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Sample Name	: pbw-pc-113-4-rac-oxh-99-1-	1.0ml		
Data Filename	: pbw-pc-113-4-rac-oxh-99-1-1.0ml2.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Vial #	1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL	A service of hus	· Overlage Administrator	
Date Processed	: 2020/10/7 15:32:38	Processed by	: System Administrator	

<Chromatogram>



Detect	or A Chann	el 2 230nm		1
Peak#	Ret. Time	Area	Height	Conc.
1	21.555	42884768	1377457	49.803
2	23.035	43223391	1389941	50.197
Total		86108159	2767398	

D:\Data\PBW\catalyst2\pbw-pc-122-4-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-113-4-rac-oxh-99-1-1.0ml2.lcd

<Sample Information>

•				
Sample Name	pbw-pc-122-3-asy-odh-98-	2-1.0ml		
Data Filename	pbw-pc-122-3-asy-odh-98-2-1.0ml1.lcd			
Method Filename	: 1.0ml-254-230.lcm			
Batch Filename				
Vial #	1-1	Sample Type	· Unknown	
Injection Volume	20.01	Cumpic Type	. Ondional	
Date Acquired	- 2020/10/7 11-20-22	Acquired by	: System Administrator	
Date Acquired	2020/10/7 11.20.22	Acquired by	. System Administrator	
Date Processed	2020/10/7 14:34:21	Processed by	: System Administrator	

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area 1 9.283 17722233 2 13.500 608016 Total 18330249 Height 1288439 32303 1320743 Conc. 96.683 3.317

Analysis Report

<sample inform<="" th=""><th>nation></th><th></th><th></th></sample>	nation>			
Sample Name	: pbw-pc-113-3b-rac-odh-98-2	-1.0ml		
Data Filename	: pbw-pc-113-3b-rac-odh-98-2-1.0ml1.lcd			
Batch Filename	1.0ml-254-230.1cm			
Vial #	: 1-1	Sample Type	: Unknown	
Date Acquired	: 20 uL : 2020/10/7 11:50:30	Acquired by	: System Administrator	
Date Processed	: 2020/10/7 14:20:04	Processed by	: System Administrator	

<Chromatogram>



Detector A Channel 2 230nm					
Peak#	Ret. Time	Area	Height	Conc.	
1	9.222	5182608	404324	49.993	
2	13.138	5184052	246070	50.007	
Total		10366659	650394		

D:\Data\PBW\pbw-pc-113-3b-rac-odh-98-2-1.0ml1.lcd

D:\Data\PBW\catalyst2\pbw-pc-122-3-asy-odh-98-2-1.0ml1.lcd

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Sample Name	: pbw-pc-86-3-asy-ozh-99-1-1	l.0ml		
Data Filename	: pbw-pc-86-3-asy-ozh-99-1-1.0ml1.lcd			
Method Filename	1.0ml-254-230.1cm			
Batch Filename				
Viel #		Sample Tune	: Unknown	
Vidi #	. 1-1	Sample Type	. UTIKHOWH	
Injection Volume	: 20 uL			
Date Acquired	: 2020/9/17 15:50:34	Acquired by	: System Administrator	
Date Processed	2020/9/17 16:10:34	Processed by	System Administrator	
Date Trocessed	. 2020/3/11 10.10.04	Trocessed by	. Oystern Administrator	

<Chromatogram>



Analysis	Report
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<sample inform<="" th=""><th>mation></th><th></th><th></th></sample>	mation>		
Sample Name Sample ID Data Filename Method Filename Batch Filename Vial # Injection Volume Date Acquired	: pbw-pc-84-3-rac-ozh-99-1-1.0ml : pbw-pc-84-3-rac-ozh-99-1-1.0ml1.lco 1.0ml-254-230.lcm : -1 : 20 uL : 2020/9/17 15:20:53	sample Type Acquired by	: Unknown : System Administrator
Date Processed	: 2020/9/17 15:53:05	Processed by	: System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	8.805	2933417	235568	50.165
2	10.837	2914112	186057	49.835
Total		5847529	421625	

D:\Data\PBW\pbw-pc-86-3-asy-ozh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-84-3-rac-ozh-99-1-1.0ml1.lcd

<Sample Information>

Sample Name	pbw-pc-133-asy-odh-99-1-1 0ml			
Sample ID	:			
Data Filename	: pbw-pc-133-asy-odh-99-1-1.0ml1.lcd			
Method Filename	: 278-254nm.lcm			
Batch Filename				
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL			
Date Acquired	: 2020/10/16 18:39:12	Acquired by	: System Administrator	
Date Processed	: 2020/10/17 9:29:47	Processed by	: System Administrator	
Sample ID Data Filename Method Filename Batch Filename Vial # Injection Volume Date Acquired Date Processed	pbw-pc-133-asy-odh-99-1-1.0ml1.lcd 278-254m.lcm 1-1 20 uL 2020/10/16 18:39:12 2020/10/16 18:39:12	Sample Type Acquired by Processed by	: Unknown : System Administrat : System Administrat	or

<Chromatogram>



Feak Table>

Peak#	Ret. Time	Area	Height	Conc.
1	10.714	4513976	252938	97.968
2	11.991	93603	5650	2.032
Total		4607580	258587	

Analysis Report

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Sample Name	: pbw-pc-120A-rac-odh-99-1-1.	0ml-2	
Data Filename Method Filename	pbw-pc-120A-rac-odh-99-1-1.	0ml-3.lcd	
Batch Filename	:		
Vial #	: 1-1	Sample Type	: Unknown
Date Acquired	20 UL 2020/10/16 19:26:45	Acquired by	· System Administrator
Date Processed	: 2020/10/17 9:37:13	Processed by	: System Administrator

<Chromatogram>



Detect	or A Chann	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	10.593	4964864	283206	50.618
2	11.804	4843710	268133	49.382
Total		9808574	551338	

D:\Data\PBW\pbw-pc-120A-rac-odh-99-1-1.0ml-3.lcd

D:\Data\PBW\catalyst2\pbw-pc-133-asy-odh-99-1-1.0ml1.lcd

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Sample Name Sample ID	: pbw-pc-140-5-asy-ozh-99-1	-1.0ml	
Data Filename	: ppw-pc-140-5-asy-ozn-99-1	-1.0ml.ica	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	1-1	Sample Type	Unknown
Injection Volume	· 20 ul	eenthis ()he	
Date Acquired	2020/10/31 10:30:20	Acquired by	· System Administrator
Date Acquireu	. 2020/10/31 10.30.20	Acquired by	. System Auministrator
Date Processed	: 2020/10/31 10:58:08	Processed by	: System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	8.713	5165255	342067	98.487
2	9.977	79354	5896	1.513
Total		5244608	347962	

Analysis Report

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Sample Name : pbw-pc-135-5-ra	c-ozh-99-1-1.0ml	
Sample ID :	CARACTER IN DESCRIPTION OF THE	
Data Filename : pbw-pc-135-5-ra	c-ozh-99-1-1.0ml001.lcd	
Method Filename : 1.0ml-254-230.ld	m	
Batch Filename :		
Vial # : 1-1	Sample Type	: Unknown
Injection Volume : 20 uL		
Date Acquired : 2020/10/31 10:46	6:08 Acquired by	: System Administrator
Date Processed : 2020/10/31 11:03	3:59 Processed by	: System Administrator

<Chromatogram>



Detect	or A Chann	el 1 254nm		t
Peak#	Ret. Time	Area	Height	Conc.
1	8.706	3134051	213020	50.021
2	9.915	3131394	192956	49.979
Total		6265445	405976	

D:\Data\PBW\catalyst2\pbw-pc-140-5-asy-ozh-99-1-1.0ml.lcd

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Sample Name	pbw-pd-11-1-asy-oxh-99-1-	1.0ml	
Data Filename Method Filename	pbw-pd-11-1-asy-oxh-99-1- 1.0ml-254-230.lcm	1.0ml1.lcd	
Vial #	: : 1-1 : 20 ul	Sample Type	: Unknown
Date Acquired Date Processed	: 2020/11/3 10:19:50 : 2020/11/3 10:41:11	Acquired by Processed by	: System Administrator : System Administrator

<Chromatogram>



Analysis	Report
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Sample Name Sample ID	: pbw-pc-142-1-rac-oxh-99-1 :	-1.0ml	
Data Filename	: pbw-pc-142-1-rac-oxh-99-1	-1.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/11/3 9:07:30	Acquired by	: System Administrator
Date Processed	: 2020/11/3 9:24:22	Processed by	: System Administrator

<Chromatogram>



Detect	or A Chann	el 1 254nm		
Peak#	Ret. Time	Area	Height	Conc.
1	6.474	1556214	151987	50.064
2	8.589	1552258	44650	49.936
Total		3108472	196637	

D:\Data\PBW\catalyst2\pbw-pd-11-1-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-142-1-rac-oxh-99-1-1.0ml1.lcd

2020/11/17 9:30:57 Page 1 / 1

2020/10/31 16:04:16 Page 1 / 1

Analysis Report

<Sample Information>

-			
Sample Name	: pbw-pc-141-asy-oxh-99-1-1.0ml		
Data Filename	: pbw-pc-141-asy-oxh-99-1-1.0ml1.lcd	t b	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	: 1-1	Sample Type	: Unknown
injection volume	: 20 UL		0 1 1 1 1 1 1
Date Acquired	2020/10/31 15:41:12	Acquired by	System Administrator
Date Processed	2020/10/31 16:03:13	Processed by	: System Administrator

<Chromatogram>



Detector A Channel 1 254nm

Peak#	Ret. Time	Area	Height	Conc.
1	6.400	27446	2916	0.583
2	7.745	4677918	198877	99.417
Total		4705364	201793	

Analysis Report

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Sample Name Sample ID Data Filename Method Filename	: pbw-pc-136-rac-oxh-99-1-1.0ml : pbw-pc-136-rac-oxh-99-1-1.0ml2.lcd : 1.0ml-254-230.lcm			
Batch Filename				
Vial #	: 1-1	Sample Type	: Unknown	
Injection Volume	: 20 uL			
Date Acquired	2020/10/31 15:30:16	Acquired by	 System Administrator 	
Date Processed	2020/10/31 15:46:11	Processed by	· System Administrator	
Date i rocesseu	. 2020/10/01 10.40.11	i loccased by	. Oystern Administrator	

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	6.356	2468710	197262	50.23
2	7.742	2445326	95798	49.76
Total		4914036	293060	

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<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	10.774	87782	6038	1.239
2	13.591	6994752	242788	98.761
Total		7082534	248826	

Analysis Report

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Sample Name	: pbw-pc-142-3-rac-oxh-99-1-	1.0ml	
Data Filename	pbw-pc-142-3-rac-oxh-99-1-	1.0ml2.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Vial #	1-1	Sample Type	: Unknown
Injection Volume	: 20 uL	Acquired by	· Sustam Administrator
Date Processed	: 2020/11/3 11:37:33	Processed by	: System Administrator

<Chromatogram>



Detect				
Peak#	Ret. Time	Area	Height	Conc.
1	10.779	4866096	289933	49.042
2	13.694	5056172	187585	50.958
Total		9922269	477518	

D:\Data\PBW\catalyst2\pbw-pd-11-2-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-142-3-rac-oxh-99-1-1.0ml2.lcd

2020/11/10 15:20:21 Page 1 / 1

Analysis Report

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O	1 11 0		
Sample Name	: pbw-pd-11-3-asy-oxn-99-1-	1.0ml	
Sample ID	1		
Data Filename	: pbw-pd-11-3-asy-oxh-99-1-	1.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	:		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	20.01	comple type	
Date Acquired	2020/11/10 15:09:49	Acquired by	· System Administrator
Date Processed	: 2020/11/10 15:03:43	Procossed by	: System Administrator
Date Flocesseu	. 2020/11/10 15.24.49	FIDCessed by	. System Auministrator

<Chromatogram>



Analysis Report

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Sample Name	: pbw-pc-142-2-rac-oxh-99-1-1.0ml

Sample ID	: pow-pc-142-2-180-0x11-33-1-	1.0111	
Data Filename Method Filename	: pbw-pc-142-2-rac-oxh-99-1-1 : 1.0ml-254-230.lcm	1.0ml1.lcd	
Batch Filename	1		
/ial #	: 1-1	Sample Type	: Unknown
niection Volume	: 20 uL		
Date Acquired	: 2020/11/10 14:39:59	Acquired by	: System Administrator
Date Processed	: 2020/11/10 15:18:23	Processed by	: System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	7.977	3354674	252916	49.937
2	9.178	3363071	169393	50.063
Total		6717745	422309	

D:\Data\PBW\pbw-pd-11-3-asy-oxh-99-1-1.0ml1.lcd

D:\Data\PBW\pbw-pc-142-2-rac-oxh-99-1-1.0ml1.lcd

2020/11/17 9:37:00 Page 1 / 1

2020/11/7 11:02:57 Page 1 / 1

Analysis Report

<Sample Information>

Sample Name	: pbw-pd-17-4-asy-oxh-99.9-0	0.1-1.0ml	
Data Filename	: pbw-pd-17-4-asy-oxh-99.9-0	0.1-1.0ml2.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	: 1-1 : 20 ul	Sample Type	Unknown
Date Acquired	2020/11/7 10:38:49	Acquired by	System Administrator
Date Processed	: 2020/11/7 11:05:30	Processed by	: System Administrator

<Chromatogram>



Detector A Channel 2 230nm

Delect				
Peak#	Ret. Time	Area	Height	Conc.
1	13.879	67856	4000	0.952
2	15.505	7061792	280844	99.048
Total		7129649	284843	

Analysis Report

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Sample Name Sample ID Data Filename Method Filename Batch Filename	: pbw-pd-7-rac-oxh-99.9-0.1-1 pbw-pd-7-rac-oxh-99.9-0.1-1 1.0ml-254-230.lcm	.0ml .0ml2.lcd	11-1
Injection Volume	: 1-1 : 20 uL	Sample Type	: Unknown
Date Acquired	: 2020/11/7 10:18:19	Acquired by	: System Administrator
Date Processed	: 2020/11/7 10:59:43	Processed by	: System Administrator

<Chromatogram>



SI Cak Table

Detect				
Peak#	Ret. Time	Area	Height	Conc.
1	13.781	2593140	126047	50.089
2	15.584	2583885	110675	49.911
Total		5177026	236722	

D:\Data\PBW\catalyst2\pbw-pd-17-4-asy-oxh-99.9-0.1-1.0ml2.lcd

D:\Data\PBW\pbw-pd-7-rac-oxh-99.9-0.1-1.0ml2.lcd

2020/11/17 9:03:58 Page 1 / 1

Analysis Report

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: pbw-pd-17-1-asy-oxh-99-1-1	.0ml	
: pbw-pd-17-1-asy-oxh-99-1-1	.0ml2.lcd	
: 1.0ml-254-230.lcm		
: 1-1	Sample Type	: Unknown
: 20 uL		
: 2020/11/16 11:36:05	Acquired by	: System Administrator
: 2020/11/16 15:00:32	Processed by	: System Administrator
	pbw-pd-17-1-asy-oxh-99-1-1 pbw-pd-17-1-asy-oxh-99-1-1 1.0ml-254-230.lcm 1-1 20 uL 2020/11/16 11:36:05 2020/11/16 15:00:32	pbw-pd-17-1-asy-oxh-99-1-1.0ml pbw-pd-17-1-asy-oxh-99-1-1.0ml2.lcd 1.0ml-254-230.lcm 1.1 Sample Type 20 uL Sample Type 20 uL Acquired by 2020/11/16 11:36:05 Acquired by 2020/11/16 15:00:32 Processed by

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Analysis Report

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Sample Name	pbw-pd-15-1-rac-oxh-99-1-1.	0ml	
Data Filename	: pbw-pd-15-1-rac-oxh-99-1-1.	0ml4.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename		O-mails T-mail	and the loss sections.
Vial #	: 1-1	Sample Type	: Unknown
Injection volume	. 20 UL	A service of buy	· Custom Administrator
Date Acquired	2020/11/17 8.43.46	Acquired by	System Administrator
Date Processed	: 2020/11/17 9:02:31	Processed by	: System Administrator

<Chromatogram>



Detect	or A Channe	el 2 230nm		
Peak#	Ret. Time	Area	Height	Conc.
1	12.847	2562508	158531	50.067
2	13.534	2555654	136119	49.933
Total		5118162	294650	

D:\Data\PBW\pbw-pd-17-1-asy-oxh-99-1-1.0ml2.lcd

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2020/11/12 23:18:05 Page 1 / 1

Analysis Report

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Sample Name	: pbw-pd-17-3-asy-oxh-99.5-0	0.5-1.0ml	
Data Filename	pbw-pd-17-3-asy-oxh-99.5-0	0.5-1.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	•		
Vial #	1.1	Sample Type	Linknown
viai #		Sample Type	. OTKHOWIT
Injection volume	: 20 UL		
Date Acquired	: 2020/11/12 22:54:28	Acquired by	: System Administrator
Date Processed	· 2020/11/12 23·11·40	Processed by	System Administrator
Duterroocoodu	. 2020/11/12 20.11.10	Troccocca by	. Of otom / tanin lot atom

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Analysis Report

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Sample Name	: pbw-pd-15-3-rac-oxh-99.5-0.	5-1.0ml	
Data Filename Method Filename	pbw-pd-15-3-rac-oxh-99.5-0.1 1.0ml-254-230.lcm	5-1.0ml1.lcd	
Batch Filename	:		
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/11/12 22:43:01	Acquired by	: System Administrator
Date Processed	: 2020/11/12 22:56:17	Processed by	: System Administrator

<Chromatogram>



Detector A Channel 1 254nm				
Peak#	Ret. Time	Area	Height	Conc.
1	6.910	3574227	363012	50.519
2	7.877	3500751	278944	49.481
Total		7074977	641956	

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2020/11/16 11:09:18 Page 1 / 1

Analysis Report

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Sample Name Sample ID Data Filename	pbw-pd-17-6-asy-oxh-99.9-1 pbw-pd-17-6-asy-oxh-99.9-1	0.1-1.0ml 0.1-1.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename	•		
Daterrinename	1	Orana la Trans	 I. Instance account
Vial #	: 1-1	Sample Type	: Unknown
Injection Volume	· 20 ul		
Date Acquired	2020/11/16 10:27:21	Acquired by	· Sustam Administrator
Date Acquired	. 2020/11/10 10.27.21	Acquired by	. System Auministrator
Date Processed	: 2020/11/16 10:58:23	Processed by	: System Administrator
			,

<Chromatogram>



Analysis Report

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Sample Name	: pbw-pd-15-6-rac-oxh-99.9-0.1	I-1.0ml	
Data Filename	pbw-pd-15-6-rac-oxh-99.9-0.1	I-1.0ml1.lcd	
Batch Filename	: 1.0ml-254-230.1cm		
Vial #	1-1	Sample Type	: Unknown
Date Acquired	: 20 uL : 2020/11/16 10:45:45	Acquired by	System Administrator
Date Processed	: 2020/11/16 11:01:55	Processed by	: System Administrator

<Chromatogram>



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Peak#	Ret. Time	Area	Height	Conc.
1	9.672	2704895	173693	50.164
2	12.705	2687179	124075	49.836
Total		5392074	297767	

D:\Data\PBW\pbw-pd-17-6-asy-oxh-99.9-0.1-1.0ml1.lcd

D:\Data\PBW\pbw-pd-15-6-rac-oxh-99.9-0.1-1.0ml1.lcd

2020/11/3 17:11:13 Page 1 / 1

Analysis Report

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Sample Name	pbw-pd-12A-asy-oxh-99.5-	0.5-1.0ml	
Data Filename	pbw-pd-12A-asy-oxh-99.5-	0.5-1.0ml1.lcd	
Method Filename	: 1.0ml-254-230.lcm		
Batch Filename			
Vial #	1-1	Sample Type	: Unknown
Injection Volume	: 20 uL		
Date Acquired	: 2020/11/3 16:34:06	Acquired by	: System Administrator
Date Processed	: 2020/11/3 17:01:26	Processed by	: System Administrator

<Chromatogram>



Detector A Channel 2 230nm Peak# Ret. Time Area 1 11.699 76484 2 14.018 7870605 Total 7947089 Height 5782 205034 210816 Conc. 0.962 99.038

Analysis Report

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Sample Name	: pbw-pc-144-1A-rac-oxh-99.5-0.5-1.0ml

Sample Name	pbw-pc-144-1A-rac-oxh-99.5	-0.5-1.0ml	
Data Filename	pbw-pc-144-1A-rac-oxh-99.5	-0.5-1.0ml2.lcd	
Batch Filename		0	
Vial # Injection Volume	: 1-1 : 20 uL	Sample Type	: Unknown
Date Acquired	: 2020/11/3 16:52:32	Acquired by	: System Administrator
Date Processed	2020/11/3 17:09:51	Processed by	: System Administrator

<Chromatogram>



Peak#	Ret. Time	Area	Height	Conc.
1	11.276	4970596	312476	50.407
2	13.131	4890352	126288	49.593
Total		9860948	438764	

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