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Supporting Information

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1. General and Materials

Commercially available reagents were used without further purification. Solvents were treated prior to use according to the standard methods. ¹H NMR, ¹³C NMR and ¹⁹F NMR spectra were recorded at room temperature in CDCl₃ on 400 MHz instrument with TMS (tetramethylsilane) as internal standard. Enantiomeric excess was determined by HPLC analysis, using chiral column described below in detail. Optical rotations were measured by polarimeter. Flash column chromatography was performed on silica gel (200-300 mesh). All reactions were monitored by TLC analysis or NMR analysis.

Materials: Commercially available reagents were used throughout without further purification. The anhydrous solvents were also purchased without the further purification.

2. General Procedure for Synthesis of [2.2]Paracyclophane Aldimines

The [2.2]paracyclophane aldimines **1** could be synthesized from formyl[2.2]paracyclophanes **5** and *p*-toulenesulfonamide in the presence of tetraethyl orthosilicate according to the known literature procedure.^[1] The intermediate formyl[2.2]paracyclophanes **5a** and **5b** were synthesized from the [2.2]paracyclophanes **S1** according to the known literature procedures.^[2] Starting material 4-methyl[2.2]paracyclophane **S1b** was prepared from 4-formyl[2.2]paracyclophane **5a** according to the known literature procedures.^[3]



[2.2]Paracyclophanes **S1** (14.7 mmol) were dissolved in dichloromethane (50 mL) and cooled to 0 °C. Titanium tetrachloride (3.22 mL, 29.4 mmol) and dichloromethoxymethane (1.39 mL, 15.4 mmol) were added subsequently. The mixture was stirred at room temperature overnight, poured into water (30 mL) and stirred at room temperature for another 2 h. The two phases were separated, and the aqueous phase was extracted with dichloromethane (20 mL×3). The combined organic phases were dried by anhydrous sodium sulfate and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel using hexanes, ethyl acetate and dichloromethane as eluent to afford the aldehyde **5a** or **5b**.

4-Formyl-7-methyl[2.2]paracyclophane 5b: The reaction was conducted by using 4-methyl-[2.2]paracyclophane **S1b** (3.528 g, 14.7 mmol), affording **5b** 2.293 g, 62% yield, white solid, mp



= 173-175 °C, the known compound,^[4] $R_f = 0.50$ (hexanes/ethyl acetate 10/1). ¹H NMR (400 MHz, CDCl₃) δ 9.93 (s, 1H), 6.91 (s, 1H), 6.81-6.71 (m, 1H), 6.51-6.39 (m, 3H), 6.22 (s, 1H), 4.12-3.99 (m, 1H), 3.41-3.29 (m, 1H), 3.29-3.14 (m, 2H), 3.07-2.94 (m, 2H), 2.89-2.71 (m, 2H), 2.16 (s, 3H). ¹³C NMR (100

MHz, CDCl₃) δ 191.7, 144.3, 143.3, 139.4, 139.4, 139.4, 138.2, 137.6, 134.8, 132.6, 132.5, 132.1, 128.2, 35.2, 33.5, 33.3, 33.3, 20.4. HRMS: Calculated for C₁₈H₁₉O [M+H]⁺ 251.1430, found: 251.1434.

The above aldehyde **5a** or **5b** (6.1 mmol), *p*-toluenesulfonamide (1.044 g, 6.1 mmol) and tetraethyl orthosilicate (1.271 g, 1.36 mL, 6.1 mmol) were combined in a schlenk flask and heated at 160 $^{\circ}$ C under nitrogen for 12 h. After cooling to room temperature, the reaction mixture was

purified by column chromatography on silica gel using hexanes and dichloromethane as eluent to afford the corresponding aldimine **1a** or **1b**.

(±)-*N*-Tosyl[2.2]paracyclophane-4-methanimine 1a: The reaction was conducted by using 4-formyl[2.2]paracyclophane 5a (2.363 g, 10.0 mmol), affording 1a 4.174 g, 91% yield, yellow



solid, mp = 172-174 °C, new compound, $R_f = 0.50$ (hexanes/ethyl acetate 10/1). ¹H NMR (400 MHz, CDCl₃) δ 8.98 (s, 1H), 7.98-7.90 (m, 2H), 7.41-7.33 (m, 2H), 7.17 (d, J = 1.9 Hz, 1H), 6.73 (dd, J = 7.8, 1.9 Hz, 1H), 6.59 (d, J = 7.8 Hz, 1H), 6.56-6.45 (m, 2H), 6.35 (dd, J = 7.9, 1.8 Hz, 1H), 6.13 (dd, J = 7.9,

1.8 Hz, 1H), 3.95-3.83 (m, 1H), 3.27-2.92 (m, 6H), 2.88-2.79 (m, 1H), 2.43 (s, 3H). 13 C NMR (100 MHz, CDCl₃) δ 168.6, 145.5, 144.5, 141.1, 139.6, 139.1, 139.0, 136.1, 135.8, 135.8, 133.3, 133.0, 132.5, 132.2, 132.0, 129.9, 127.9, 35.5, 35.3, 34.9, 34.2, 21.8. HRMS: Calculated for C₂₄H₂₄NO₂S [M+H]⁺ 390.1522, found: 390.1525.

(±)-*N*-Tosyl-7-methyl[2.2]paracyclophane-4-methanimine 1b: The reaction was conducted by using 5b (1.529 g, 6.1 mmol), affording 1b 1.160 g, 47% yield, yellow solid, mp = 216-218 °C,



new compound, $R_f = 0.50$ (hexanes/ethyl acetate 5/1). ¹H NMR (400 MHz, CDCl₃) δ 8.97 (s, 1H), 8.05-7.82 (m, 2H), 7.42-7.31 (m, 2H), 7.07 (s, 1H), 6.75 (dd, J = 7.9, 1.9 Hz, 1H), 6.43 (dd, J = 7.9, 1.8 Hz, 1H), 6.36 (dd, J = 7.9, 1.8 Hz, 1H), 6.27-6.15 (m, 2H), 3.96-3.79 (m, 1H), 3.35-3.09 (m, 3H),

3.04-2.91 (m, 1H), 2.88-2.71 (m, 3H), 2.43 (s, 3H), 2.15 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 168.6, 145.7, 145.6, 144.4, 139.9, 139.5, 139.0, 138.2, 137.2, 136.2, 132.4, 132.3, 132.1, 130.4, 129.9, 128.3, 127.9, 35.5, 33.9, 33.6, 33.1, 21.8, 20.6. HRMS: Calculated for C₂₅H₂₆NO₂S [M+H]⁺ 404.1679, found: 404.1676.

4-Methoxy-5-formyl[2.2]paracyclophane **5c** was synthesized from the [2.2]paracyclophane **S1a** according to the known literature procedure.^[5]



The aldehyde **5c** (0.173 g, 0.7 mmol), *p*-toluenesulfonamide (0.123 g, 0.7 mmol) and tetraethyl orthosilicate (0.150 g, 0.16 mL, 0.7 mmol) were combined in a schlenk flask and heated at 160 $^{\circ}$ C under nitrogen for 12 h. After cooling to room temperature, the reaction mixture was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the corresponding aldimine **1c**.

(±)-*N*-Tosyl-4-methoxy[2.2]paracyclophane-5-methanimine 1c: 0.195 g, 72% yield, yellow solid, mp = 61-63 $^{\circ}$ C, new compound, R_f = 0.35 (hexanes/ethyl acetate 10/1). ¹H NMR (400 MHz,



CDCl₃) δ 9.09 (s, 1H), 8.02-7.90 (m, 2H), 7.41-7.33 (m, 2H), 6.73-6.64 (m, 2H), 6.54-6.48 (m, 1H), 6.45-6.36 (m, 2H), 6.05-5.96 (m, 1H), 4.23-4.06 (m, 1H), 3.76 (s, 3H), 3.36-3.23 (m, 1H), 3.17-2.94 (m, 3H), 2.80-2.59 (m, 3H), 2.44 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 167.7, 163.5, 145.9, 144.3, 141.7, 139.3, 139.2, 135.9, 133.5, 132.4, 131.9, 131.6, 131.1, 129.8, 129.3,

127.9, 126.1, 62.5, 35.6, 34.1, 33.4, 31.2, 21.7. HRMS: Calculated for $C_{25}H_{26}NO_3S$ [M+H]⁺ 420.1628, found: 420.1634.

3. General Procedure for Kinetic Resolution



A schlenk tube (25 mL) was charged with $Pd(OCOCF_3)_2$ (3.3 mg, 0.01 mmol, 5 mol%) and (R_p ,S)-L2 (5.0 mg, 0.01 mmol, 5 mol%) under nitrogen, and degassed anhydrous acetone (1.5 mL) was added. The mixture was stirred at room temperature for 1 h. The solvent was removed under vacuum to give the catalyst. Then substrate *rac*-1 (77.9 mg, 0.20 mmol), arylboronic acid (0.20 mmol) and 2,2,2-trifluoroethanol (4.0 mL) were added into the tube under nitrogen. The mixture was heated to 60 °C. After stirring at 60 °C for 15 h, the reaction mixture was cooled to room temperature, and the solvent was removed by rotary evaporation. The resulting mixture was dried under vacuum and the conversion of *rac*-1 was confirmed by ¹H NMR analysis with benzyl ether as internal standard. The solvent was removed in *vacuo*, the recovered material (-)-1 and addition product **3** were isolated by column chromatography on silica gel using hexanes and ethyl acetate as eluent.

To a solution of the addition product **3** in dry *N*,*N*-dimethylformamide (DMF, 1.0 mL) was added sodium hydride (15 mg, 0.38 mmol, 60% wt.) at 0 °C, and then allyl bromide (36 mg, 26.0 μ L, 0.30 mmol) was added dropwise. The reaction mixture was warmed to room temperature and stirred at room temperature for 2 h. Water (5.0 mL) was added, and extracted with ethyl acetate (10 mL×3). The combined organic layer was washed with brine, dried by anhydrous sodium sulfate and filtered, concentrated in *vacuo* and analyzed by crude ¹H NMR to determine diastereomeric ratio. The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the product (+)-2.

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(phenyl)methyl}-4-methylbenzenesulfonamide (2a): 48.9 mg, 48% yield, 12:1 dr, white solid, mp = 144-146 $^{\circ}$ C, new compound, R_f = 0.50 (hexanes/



ethyl acetate 10/1), 98.8% ee, $[\alpha]^{20}_{D} = 114.70$ (*c* 0.24, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 8.01-7.89 (m, 2H), 7.43-7.35 (m, 2H), 7.24-7.18 (m, 1H), 7.17-7.07 (m, 3H), 6.92-6.84 (m, 2H), 6.76-6.71 (m, 1H), 6.64-6.46 (m, 4H), 6.34 (d, *J* = 7.6 Hz, 1H), 6.20 (s, 1H), 4.82-4.69 (m, 1H), 4.66-4.50 (m, 2H), 4.05-3.90 (m, 1H), 3.46 (dd, *J* = 15.6, 3.6 Hz, 1H), 3.31-3.15 (m, 2H),

3.08-2.87 (m, 4H), 2.55-2.38 (m, 2H), 2.50 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.5, 140.9, 140.0, 139.5, 139.2, 138.9, 138.6, 136.9, 136.0, 134.0, 133.7, 132.6, 132.3, 132.2, 131.4, 130.5, 129.7, 129.4, 128.4, 128.2, 128.0, 117.2, 64.6, 48.5, 35.5, 35.4, 34.6, 34.5, 21.7. HPLC: Chiracel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 6.9 min (major) and 7.5 min. HRMS: Calculated for C₃₃H₃₃KNO₂S [M+K]⁺ 546.1864, found: 546.1861.

N-Allyl-*N*-{[2.2]paracyclophan-4-yl(phenyl)methyl}-4-methylbenzenesulfonamide (2a'):

white solid, mp = 176-178 °C, new compound, $R_f = 0.35$ (hexanes/ethyl acetate 10/1). ¹H NMR



(400 MHz, CDCl₃) δ 7.34-7.27 (m, 2H), 7.26-7.23 (m, 1H), 7.19-7.12 (m, 4H), 6.93 (d, J = 8.1 Hz, 2H), 6.87 (dd, J = 7.9, 1.7 Hz, 1H), 6.62-6.43 (m, 4H), 6.29 (s, 1H), 6.04 (dd, J = 7.9, 1.7 Hz, 1H), 5.68-5.54 (m, 2H), 4.85-4.76 (m, 1H), 4.72-4.59 (m, 1H), 4.00-3.83 (m, 1H), 3.57-3.35 (m, 3H), 3.30-3.20 (m, 1H), 3.05-2.82 (m, 4H), 2.74-2.59 (m, 1H), 2.30 (s, 3H). ¹³C NMR (100 MHz, 100 MHz, 100

CDCl₃) δ 142.5, 140.3, 140.0, 139.7, 139.3, 138.8, 137.2, 136.2, 136.1, 134.1, 133.3, 133.2, 133.0, 132.2, 129.6, 128.8, 128.7, 128.7, 127.6, 127.6, 116.0, 66.0, 49.9, 35.1, 35.0, 34.7, 33.8, 21.5. HRMS: Calculated for C₃₃H₃₃KNO₂S [M+K]⁺ 546.1864, found: 546.1827.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with phenylboronic acid, 30.7 mg, 39% yield, 93.7% ee, $[\alpha]^{20}_{D} =$ -356.96 (*c* 0.63, CHCl₃). HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.6 min and 18.9 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(*o*-tolyl)methyl}-4-methylbenzenesulfonamide (2b): 51.9 mg, 50% yield, 7:1 dr, white solid, mp = 58-60 $^{\circ}$ C, new compound, R_f = 0.60 (hexanes/ ethyl



acetate 10/1), 98.7% ee, $[\alpha]^{20}{}_{D} = 45.71$ (*c* 0.28, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.96-7.88 (m, 2H), 7.37-7.30 (m, 2H), 7.26-7.23 (m, 1H), 7.19-7.08 (m, 2H), 6.85 (t, *J* = 7.4 Hz, 1H), 6.77-6.68 (m, 3H), 6.64-6.59 (m, 1H), 6.56-6.46 (m, 3H), 6.32 (d, *J* = 7.5 Hz, 1H), 4.34-4.23 (m, 3H), 3.96-3.85 (m, 1H), 3.60-3.50 (m, 1H), 3.35-2.90 (m, 7H), 2.69 (s, 3H), 2.54-2.41 (m, 2H),

2.45 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.6, 140.0, 139.7, 139.4, 139.3, 138.5, 138.2, 137.7, 137.1, 136.0, 133.6, 133.4, 132.6, 132.1, 132.0, 131.1, 130.9, 130.4, 130.3, 129.4, 128.6, 127.9, 125.6, 115.7, 61.1, 47.4, 35.5, 35.4, 34.8, 34.1, 21.6, 20.0. HPLC: Chiracel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 5.3 min (major) and 7.0 min. HRMS: Calculated for C₃₄H₃₅NaNO₂S [M+Na]⁺ 544.2281, found: 544.2282.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 2-methylphenylboronic acid, 29.6 mg, 38% yield, 80.1% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.7 min and 18.9 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(*m*-tolyl)methyl}-4-methylbenzenesulfonamide (2c): 50.7 mg, 49% yield, 12:1 dr, white solid, mp = 63-65 $^{\circ}$ C, new compound, R_f = 0.60 (hexanes/ethyl



acetate 10/1), 98.1% ee, $[\alpha]_{D}^{20} = 135.93$ (*c* 0.32, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.98-7.88 (m, 2H), 7.44-7.35 (m, 2H), 7.13-7.07 (m, 1H), 7.03-6.95 (m, 2H), 6.73 (s, 1H), 6.67-6.57 (m, 3H), 6.55-6.45 (m, 3H), 6.32 (d, *J* = 7.6 Hz, 1H), 6.05 (s, 1H), 4.84-4.73 (m, 1H), 4.68-4.51 (m, 2H), 3.95 (dd, *J* =

Ar = $3 - MeC_6H_4$ 16.4, 7.8 Hz, 1H), 3.45 (dd, J = 16.4, 4.3 Hz, 1H), 3.28-3.13 (m, 2H), 3.09-2.86 (m, 4H), 2.51 (s, 3H), 2.48-2.38 (m, 2H), 2.12 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.4, 140.7, 140.0, 139.6, 139.2, 139.0, 138.7, 137.9, 137.0, 136.0, 134.3, 133.8, 132.7, 132.3, 132.2, 131.5, 130.5, 130.2, 129.7, 128.7, 128.3, 128.2, 126.3, 117.3, 64.6, 48.6, 35.5, 35.4, 34.6, 34.6, 21.7, 21.4. HPLC: Chiracel IC column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 15.8 min and 21.7 min (major). HRMS: Calculated for C₃₄H₃₅NaNO₂S [M+Na]⁺ 544.2281, found: 544.2282.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 3-methylphenylboronic acid, 21.7 mg, 28% yield, 91.2% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.6 min and 18.9 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(*p*-tolyl)methyl}-4-methylbenzenesulfonamide (2d): 55.2 mg, 53% yield, 13:1 dr, white solid, mp = 56-58 °C, new compound, $R_f = 0.55$ (hexanes/



ethyl acetate 10/1), 98.3% ee, $[\alpha]^{20}{}_{D} = 134.91$ (*c* 0.61, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 8.03-7.87 (m, 2H), 7.47-7.34 (m, 2H), 7.09 (d, *J* = 7.7 Hz, 1H), 6.93 (d, *J* = 7.8 Hz, 2H), 6.79-6.69 (m, 3H), 6.59 (t, *J* = 8.0 Hz, 2H), 6.55-6.45 (m, 2H), 6.32 (d, *J* = 7.6 Hz, 1H), 6.15 (s, 1H), 4.82-4.70 (m, 1H), 4.68-4.53 (m, 2H), 4.01-3.87 (m, 1H), 3.51-3.39 (m, 1H), 3.28-3.14 (m, 2H),

3.07-2.86 (m, 4H), 2.56-2.41 (m, 2H), 2.50 (s, 3H) 2.28 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.4, 140.0, 139.5, 139.3, 139.0, 138.7, 137.9, 137.7, 137.2, 136.0, 134.2, 133.7, 132.6, 132.3, 132.2, 131.5, 130.5, 129.7, 129.3, 129.0, 128.2, 117.2, 64.4, 48.5, 35.5, 35.4, 34.6, 34.6, 21.7, 21.2. HPLC: Chiracel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 7.1 min and 8.5 min (major). HRMS: Calculated for C₃₄H₃₅NaNO₂S [M+Na]⁺ 544.2281, found: 544.2284.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 4-methylphenylboronic acid, 27.8 mg, 36% yield, 88.0% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.5 min and 18.8 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(4-fluorophenyl)methyl}-4-methylbenzenesulfonam ide (2e): 51.9 mg, 49% yield, 16:1 dr, white solid, mp = 164-166 $^{\circ}$ C, new compound, R_f = 0.45



(hexanes/ethyl acetate 10/1), 96.3% ee, $[\alpha]^{20}{}_{D} = 105.59$ (*c* 0.25, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.96-7.88 (m, 2H), 7.43-7.36 (m, 2H), 7.04 (dd, *J* = 7.8, 1.8 Hz, 1H), 6.89-6.78 (m, 4H), 6.71-6.66 (m, 1H), 6.62-6.57 (m, 1H), 6.55-6.45 (m, 3H), 6.37-6.30 (m, 1H), 6.18 (s, 1H), 4.77-4.58 (m, 3H), 4.05-3.90 (m, 1H), 3.50-3.36 (m, 1H), 3.24-3.13 (m, 2H), 3.04-2.87 (m, 4H),

2.56-2.35 (m, 2H), 2.50 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 162.4 (d, $J_{C-F} = 247.7$ Hz), 143.7, 140.2, 139.6, 139.2, 138.9, 138.5, 137.0 (d, $J_{C-F} = 3.3$ Hz), 136.8, 136.1, 133.9, 133.7, 132.7, 132.5, 132.3, 131.4, 131.1 (d, $J_{C-F} = 8.1$ Hz), 130.4, 129.8, 128.1, 117.4, 115.3 (d, $J_{C-F} = 21.3$ Hz), 63.8, 48.5, 35.5, 35.4, 34.6, 34.5, 21.7. ¹⁹F NMR (376 MHz, CDCl₃) δ -113.8. HPLC: Chiracel IB column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 1.0 mL/min, retention time 5.1 min (major) and 5.5 min. HRMS: Calculated for C₃₃H₃₂FNaNO₂S [M+Na]⁺ 548.2030, found: 548.2026.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 4-fluorophenylboronic acid, 33.8 mg, 43% yield, 99.2% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.4 min and 18.6 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(4-(trifluoromethyl)phenyl)methyl}-4-methylbenze nesulfonamide (2f): 58.2 mg, 51% yield, >20:1 dr, white solid, mp = 108-110 °C, new compound,



 $R_f = 0.43$ (hexanes/ethyl acetate 10/1), 98.7% ee, $[α]^{20}_D = 106.49$ (*c* 0.20, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.96-7.87 (m, 2H), 7.45-7.36 (m, 4H), 7.08-7.00 (m, 3H), 6.72-6.67 (m, 1H), 6.64-6.58 (m, 1H), 6.56-6.47 (m, 3H), 6.35 (d, *J* = 7.6 Hz, 1H), 6.27 (s, 1H), 4.74-4.57 (m, 3H), 4.02-3.92 (m, 1H), 3.50-3.38 (m, 1H), 3.26-3.14 (m, 2H), 3.07-2.89 (m, 4H), 2.57-2.43 (m, 1H),

2.51 (s, 3H), 2.39-2.27 (m, 1H). ¹³C NMR (100 MHz, CDCl₃) δ 145.1, 143.8, 140.3, 139.6, 139.1, 138.7, 138.3, 136.2, 135.9, 133.6, 133.5, 132.7, 132.7, 132.4, 131.4, 130.4, 130.2 (q, $J_{C-F} = 32.4$ Hz), 129.8, 129.8, 128.1, 125.3 (q, $J_{C-F} = 3.7$ Hz), 124.0 (q, $J_{C-F} = 272.2$ Hz), 117.5, 64.0, 48.6, 35.4, 35.4, 34.6, 34.5, 21.7. ¹⁹F NMR (376 MHz, CDCl₃) δ -62.5. HPLC: Chiracel IB column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 5.9 min (major) and 6.6 min. HRMS: Calculated for C₃₄H₃₆F₃N₂O₂S [M+NH₄]⁺ 593.2444, found: 593.2462.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 4-trifluoromethylphenylboronic acid, 28.8 mg, 37% yield, 98.5% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.1 min and 18.3 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(4-chlorophenyl)methyl}-4-methylbenzenesulfona mide (2g): 54.1 mg, 50% yield, >20:1 dr, white solid, mp = 139-141 °C, new compound, $R_f = 0.43$



(hexanes/ethyl acetate 10/1), 98.9% ee, $[\alpha]^{20}{}_{D} = 134.50$ (*c* 0.82, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.96-7.88 (m, 2H), 7.43-7.36 (m, 2H), 7.15-7.08 (m, 2H), 7.05 (dd, *J* = 7.8, 1.4 Hz, 1H), 6.86-6.79 (m, 2H), 6.72-6.67 (m, 1H), 6.64-6.57 (m, 1H), 6.56-6.46 (m, 3H), 6.34 (d, *J* = 7.6 Hz, 1H), 6.17 (s, 1H), 6.86-6.57 (m, 2H), 6.25 (m, 2H), 6.25

Ar = 4-CIC₆H₄ 4.80-4.60 (m, 3H), 4.03-3.88 (m, 1H), 3.49-3.35 (m, 1H), 3.25-3.14 (m, 2H), 3.06-2.89 (m, 4H), 2.55-2.45 (m, 1H), 2.50 (s, 3H), 2.44-2.33 (m, 1H). ¹³C NMR (100 MHz, CDCl₃) δ 143.7, 140.2, 139.6, 139.5, 139.1, 138.8, 138.4, 136.4, 136.1, 133.9, 133.8, 133.6, 132.7, 132.6, 132.3, 131.4, 130.7, 130.4, 129.8, 128.6, 128.1, 117.5, 63.8, 48.5, 35.4, 35.4, 34.6, 34.5, 21.7. HPLC: Chiracel IB column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 6.4 min (major) and 7.0 min. HRMS: Calculated for C₃₃H₃₆ClN₂O₂S [M+NH₄]⁺ 559.2181, found: 559.2175 (³⁵Cl), 561.2160 (³⁷Cl).

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 4-chlorophenylboronic acid, 31.2 mg, 40% yield, 99.6% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.4 min and 18.6 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(3-fluorophenyl)methyl}-4-methylbenzenesulfonam ide (2h): 52.2 mg, 50% yield, >20:1 dr, white solid, mp = 134-136 °C, new compound, $R_f = 0.43$



(hexanes/ethyl acetate 10/1), 98.9% ee, $[\alpha]^{20}{}_{D} = 123.93$ (*c* 0.66, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.99-7.90 (m, 2H), 7.45-7.36 (m, 2H), 7.16-7.02 (m, 2H), 6.95-6.84 (m, 1H), 6.75-6.67 (m, 2H), 6.64-6.58 (m, 1H), 6.57-6.46 (m, 4H), 6.34 (d, *J* = 7.6 Hz, 1H), 6.16 (s, 1H), 4.78-4.59 (m, 3H), 4.04-3.93 (m, 1H), 3.49-3.38 (m, 1H), 3.26-3.14 (m, 2H), 3.06-2.89 (m, 4H), 2.57-2.46 (m,

1H), 2.51 (s, 3H), 2.46-2.37 (m, 1H). ¹³C NMR (100 MHz, CDCl₃) δ 162.7 (d, $J_{C-F} = 246.7$ Hz), 143.8, 143.5 (d, $J_{C-F} = 6.6$ Hz), 140.2, 139.6, 139.2, 138.7, 138.5, 136.2, 136.1, 133.8, 133.7, 132.7, 132.6, 132.3, 131.4, 130.4, 129.8 (d, $J_{C-F} = 8.0$ Hz), 129.8, 128.1, 125.1 (d, $J_{C-F} = 2.8$ Hz),

117.4, 116.4 (d, $J_{C-F} = 22.1$ Hz), 115.0 (d, $J_{C-F} = 21.1$ Hz), 64.0 (d, $J_{C-F} = 1.6$ Hz), 48.5, 35.4, 35.4, 34.6, 34.5, 21.7. ¹⁹F NMR (376 MHz, CDCl₃) δ -112.6. HPLC: Chiracel IA column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 6.4 min (major) and 7.2 min. HRMS: Calculated for C₃₃H₃₂FNaNO₂S [M+Na]⁺ 548.2030, found: 548.2061.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 3-fluorophenylboronic acid, 31.9 mg, 41% yield, 97.4% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.2 min and 18.3 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(3-chlorophenyl)methyl}-4-methylbenzenesulfona mide (2i): 54.4 mg, 50% yield, >20:1 dr, white solid, mp = 56-58 °C, new compound, $R_f = 0.41$



(hexanes/ethyl acetate 10/1), 99.4% ee, $[\alpha]^{20}_{D} = 114.30$ (*c* 1.10, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.98-7.89 (m, 2H), 7.47-7.38 (m, 2H), 7.21-7.15 (m, 1H), 7.10-7.01 (m, 2H), 6.82-6.75 (m, 1H), 6.73-6.68 (m, 1H), 6.63-6.58 (m, 2H), 6.58-6.47 (m, 3H), 6.34 (d, *J* = 7.6 Hz, 1H), 6.06 (s, 1H), 4.86-4.58 (m, 3H), 4.07-3.92 (m, 1H), 3.52-3.37 (m, 1H), 3.29-3.13 (m, 2H), 3.08-2.85 (m, 3H), 4.07-3.92 (m, 2H), 3.08-2.85 (m, 2H), 3.29-3.13 (m, 2H), 3.08-2.85 (m, 2H), 3.08-

4H), 2.58-2.31 (m, 2H), 2.52 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.9, 142.9, 140.2, 139.6, 139.1, 138.6, 138.4, 136.2, 136.1, 134.3, 133.8, 133.7, 132.7, 132.6, 132.3, 131.4, 130.4, 129.9, 129.6, 129.5, 128.2, 128.1, 127.4, 117.6, 64.0, 48.6, 35.4, 35.4, 34.6, 34.5, 21.7. HPLC: Chiracel IC column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 11.2 min and 13.2 (major) min. HRMS: Calculated for $C_{33}H_{36}CIN_2O_2S$ [M+NH₄]⁺ 559.2181, found: 559.2175 (³⁵Cl), 561.2155 (³⁷Cl).

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 3-chlorophenylboronic acid, 32.7 mg, 42% yield, 98.4% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.3 min and 18.5 min (major).

(+)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(4-methoxyphenyl)methyl}-4-methylbenzenesulfon amide (2j): 51.6 mg, 48% yield, 13:1 dr, white solid, mp = 65-67 $^{\circ}$ C, new compound, R_f = 0.25



(hexanes/ethyl acetate 10/1), 96.5% ee, $[\alpha]_{D}^{20} = 135.32$ (*c* 0.30, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.97-7.88 (m, 2H), 7.42-7.35 (m, 2H), 7.07 (dd, *J* = 7.8, 1.5 Hz, 1H), 6.80-6.74 (m, 2H), 6.73-6.69 (m, 1H), 6.68-6.62 (m, 2H), 6.61-6.44 (m, 4H), 6.32 (d, *J* = 7.6 Hz, 1H), 6.13 (s, 1H), 4.81-4.72 (m, 1H), 4.70-4.59 (m, 2H), 3.98-3.89 (m, 1H), 3.75 (s, 3H), 3.47-3.38 (m, 1H),

3.25-3.14 (m, 2H), 3.05-2.88 (m, 4H), 2.56-2.43 (m, 2H), 2.49 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 159.2, 143.4, 140.0, 139.5, 139.3, 139.0, 138.7, 137.4, 136.0, 134.3, 133.7, 133.1, 132.7, 132.3, 132.2, 131.5, 130.6, 130.5, 129.7, 128.2, 117.2, 113.6, 64.1, 55.3, 48.5, 35.5, 35.4, 34.6, 34.6, 21.7. HPLC: Chiracel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 1.0 mL/ min, retention time 6.5 min and 7.2 min (major). HRMS: Calculated for C₃₄H₃₅KNO₃S [M+K]⁺ 576.1969, found: 576.1959.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 4-methoxyphenylboronic acid, 35.1 mg, 45% yield, 83.9% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.3 min and 18.5 min (major).

(+)-N-Allyl-N-{[2.2]paracyclophan-4-yl(2-naphthyl)methyl}-4-methylbenzenesulfonamide

(2k): 54.0 mg, 48% yield, 12:1 dr, white solid, mp = 66-68 °C, new compound, $R_f = 0.38$



(hexanes/ethyl acetate 10/1), 98.4% ee, $[\alpha]^{20}{}_{D} = 155.39$ (*c* 0.87, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 8.02-7.94 (m, 2H), 7.80-7.74 (m, 1H), 7.63 (d, *J* = 8.5 Hz, 1H), 7.55-7.50 (m, 1H), 7.49-7.38 (m, 4H), 7.21-7.18 (m, 1H), 7.18-7.12 (m, 1H), 7.03 (dd, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6.63 (d, *J* = 8.5, 1.5 Hz, 1H), 6.84-6.79 (m, 1H), 6

Ar = 2-Naphthyl = 8.0 Hz, 2H), 6.53 (d, J = 7.1 Hz, 2H), 6.39-6.29 (m, 2H), 4.76-4.67 (m, 1H), 4.64-4.49 (m, 2H), 4.04-3.92 (m, 1H), 3.63-3.50 (m, 1H), 3.31-3.20 (m, 2H), 3.12-2.96 (m, 3H), 2.95-2.85 (m, 1H), 2.54 (s, 3H), 2.48-2.39 (m, 2H). ¹³C NMR (100 MHz, CDCl₃) δ 143.5, 140.1, 139.6, 139.2, 138.9, 138.7, 138.2, 136.9, 136.1, 134.1, 133.8, 133.0, 132.9, 132.7, 132.5, 132.3, 131.6, 130.5, 129.8, 128.5, 128.3, 128.1, 128.1, 127.7, 127.3, 126.4, 126.3, 117.4, 64.7, 48.7, 35.5, 35.4, 34.6, 21.7. HPLC: Chiracel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 9.0 min and 13.8 min (major). HRMS: Calculated for C₃₇H₃₅Na-NO₂S [M+Na]⁺ 580.2281, found: 580.2279.

(-)-*N*-Tosyl[2.2]paracyclophane-4-methanimine (1a): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1a with 2-naphthaleneboronic acid, 31.9 mg, 41% yield, 82.3% ee. HPLC: Chiralcel AD-3 column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.3 min and 18.5 min (major).

(+)-*N*-Allyl-*N*-{7-methyl[2.2]paracyclophan-4-yl(phenyl)methyl}-4-methylbenzenesulfona mide (21): 52.1 mg, 50% yield, 8:1 dr, white solid, mp = 183-185 °C, new compound, $R_f = 0.50$



(hexanes/ethyl acetate 10/1), 97.9% ee, $[\alpha]^{20}_{D} = 158.03$ (*c* 0.52, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 8.00-7.88 (m, 2H), 7.44-7.32 (m, 2H), 7.24-7.08 (m, 4H), 6.96-6.82 (m, 3H), 6.61 (s, 1H), 6.57-6.50 (m, 1H), 6.46-6.35 (m, 1H), 6.22 (s, 1H), 5.96 (s, 1H), 4.80-4.65 (m, 1H),

4.65-4.49 (m, 2H), 4.03-3.90 (m, 1H), 3.51-3.40 (m, 1H), 3.40-3.29 (m, 1H), 3.19 (t, J = 11.8 Hz, 1H), 3.04-2.74 (m, 4H), 2.50 (s, 3H), 2.41-2.29 (m, 2H), 2.13 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.4, 141.1, 139.2, 139.0, 138.9, 138.5, 138.2, 138.1, 136.6, 134.7, 134.2, 133.9, 132.6, 131.3, 130.8, 129.7, 129.4, 128.4, 128.2, 128.0, 127.8, 117.2, 64.5, 48.4, 34.3, 34.0, 33.6, 33.4, 21.7, 19.8. HPLC: Chiracel IB column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 6.2 min (major) and 7.1 min. HRMS: Calculated for C₃₄H₃₅NaNO₂S [M+Na]⁺ 544.2281, found: 544.2262.

(-)-*N*-Tosyl-7-methyl[2.2]paracyclophane-4-methanimine (1b): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1b with phenylboronic acid, 32.2 mg, 40% yield, 99.4% ee, $[\alpha]^{20}_{D} = -400.28$ (*c* 0.65, CHCl₃). HPLC: Chiralcel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.7 min and 20.5 min (major).

(+)-*N*-Allyl-*N*-{7-methyl[2.2]paracyclophan-4-yl(4-chlorophenyl)methyl}-4-methylbenzene sulfonamide (2m): 52.3 mg, 47% yield, 17:1 dr, white solid, mp = 55-57 °C, new compound, $R_f = 1000$ ms solid methyl so



0.50 (hexanes/ethyl acetate 10/1), 97.9% ee, $[\alpha]_{D}^{20}$ = 150.36 (*c* 0.54, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.98-7.86 (m, 2H), 7.46-7.34 (m, 2H), 7.15-7.04 (m, 3H), 6.90-6.78 (m, 3H), 6.56 (s, 1H), 6.49 (dd, J = 7.8, 1.4 Hz, 1H), 6.39 (dd, J = 7.8, 1.6 Hz, 1H), 6.18 (s, 1H), 5.96 (s, 1H), 4.81-4.57 (m, 3H), 4.03-3.90 (m, 1H), 3.48-3.27 (m, 2H), 3.23-3.10 (m

1H), 3.06-2.85 (m, 3H), 2.84-2.70 (m, 1H), 2.50 (s, 3H), 2.41-2.27 (m, 2H), 2.12 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.7, 139.7, 139.2, 138.9, 138.8, 138.6, 138.2, 138.0, 136.9, 134.1,

133.9, 133.8, 133.8, 132.6, 131.4, 130.7, 130.7, 129.8, 128.6, 128.1, 127.9, 117.5, 63.7, 48.5, 34.3, 34.1, 33.6, 33.4, 21.7, 19.8. HPLC: Chiracel IB column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 6.4 min (major) and 7.5 min. HRMS: Calculated for $C_{34}H_{38}CIN_2O_2S$ [M+NH₄]⁺ 573.2337, found: 573.2376 (³⁵Cl), 575.2360 (³⁷Cl).

(-)-*N*-Tosyl-7-methyl[2.2]paracyclophane-4-methanimine (1b): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1b with 4-chlorophenylboronic acid, 33.8 mg, 42% yield, 99.9% ee. HPLC: Chiralcel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 16.7 min and 20.5 min (major).

(+)-*N*-Allyl-*N*-{4-methoxy[2.2]paracyclophan-5-yl(4-chlorophenyl)methyl}-4-methylbenze nesulfonamide (2n): 14.0 mg, 12% yield, >20:1 dr, white solid, mp = 45-47 °C, new compound,



$$\begin{split} R_{\rm f} &= 0.45 \text{ (hexanes/ethyl acetate 10/1), } 99.4\% \text{ ee, } \left[\alpha\right]^{20}{}_{\rm D} = 6.07 \text{ (c 0.28, CHCl_3).} ^1\text{H NMR (400 MHz, CDCl_3) } \delta 7.33-7.28 \text{ (m, 2H), } 7.22-7.12 \text{ (m, 4H), } 7.11-6.99 \text{ (m, 2H), } 6.70-6.59 \text{ (m, 2H), } 6.55-6.49 \text{ (m, 1H), } 6.40-6.32 \text{ (m, 2H), } 6.28-6.22 \text{ (m, 1H), } 6.19-6.13 \text{ (m, 1H), } 5.13-4.97 \text{ (m, 1H), } 4.79-4.66 \text{ (m, 2H), } 6.40-6.32 \text{ (m, 2H), } 6.28-6.22 \text{ (m, 2H), } 6.19-6.13 \text{ (m, 2H), } 5.13-4.97 \text{ (m, 2H), } 6.40-6.32 \text{ (m, 2H), } 6.28-6.22 \text{ (m, 2H), } 6.19-6.13 \text{ (m, 2H), } 5.13-4.97 \text{ (m, 2H), } 6.40-6.32 \text{ (m, 2H), } 6.28-6.22 \text{ (m, 2H), } 6.19-6.13 \text{ (m, 2H), } 5.13-4.97 \text{ (m, 2H), } 6.40-6.32 \text{ (m, 2H), } 6.28-6.22 \text{ (m, 2H), } 6.19-6.13 \text{ (m, 2H), } 5.13-4.97 \text{ (m, 2H), } 6.40-6.32 \text{ (m, 2H), } 6.19-6.13 \text{ (m, 2H),$$

Ar = 4-CIC₆H₄ 2H), 3.99-3.80 (m, 2H), 3.65-3.53 (m, 1H), 3.15-2.72 (m, 7H), 2.91 (s, 3H), 2.36 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 158.0, 142.8, 141.5, 139.3, 139.2, 139.0, 137.8, 135.9, 135.1, 133.2, 132.4, 132.0, 131.5, 131.2, 130.9, 129.3, 129.0, 128.7, 128.1, 127.7, 116.6, 61.9, 60.9, 49.0, 34.8, 34.7, 33.2, 31.0, 21.5. HPLC: Chiracel IA column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 9.9 min (major) and 10.9 min. HRMS: Calculated for C₃₄H₃₄ClNNaO₃S [M+Na]⁺ 594.1840, found: 594.1851 (³⁵Cl), 596.1824 (³⁷Cl).

(-)-*N*-Tosyl-4-methoxy[2.2]paracyclophane-5-methanimine (1c): Kinetic resolution from the addition of [2.2]paracyclophane aldimine 1c with 4-chlorophenylboronic acid, 71.0 mg, 85% yield, 13.2% ee. $[\alpha]^{20}_{D} = -14.09$ (*c* 0.44, CHCl₃). HPLC: Chiralcel IA column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 80/20, flow = 0.8 mL/min, retention time 10.8 min (major) and 16.4 min.

4. Elaborations of Recovered Material and Product

Derivatizations of the Recovered Material (R_p) -(-)-1a Ts_N $\xrightarrow{\text{NaBH}_4, \text{ MeOH}, \text{ rt}}$ Ts_N $\stackrel{\text{Ts}_N}{\xrightarrow{\text{H}}}$

(*R_p*)-(-)-**1a:** 99% ee (*R_p*)-(-)-**4**: 99% ee, 94% yield

Sodium tetrahydroborate (38 mg, 1.0 mmol) was added to a solution of aldimine (-)-1a (78 mg, 0.20 mmol, 99% ee) in methanol (5.0 mL). The reaction was performed at room temperature overnight. The reaction mixture was quenched by addition of saturated aqueous ammonium chloride solution (15 mL). After being extracted with ethyl acetate (15 mL×3), the combined organic layer was dried by anhydrous sodium sulfate, concentrated in *vacuo*, then purification by silica gel chromatography using hexanes and ethyl acetate as eluent to gave product (-)-4.

(-)-*N*-([2.2]Paracyclophan-4-ylmethyl)-4-methylbenzenesulfonamide (4): 73 mg, 94% yield, white solid, mp = 49-51 °C, new compound, $R_f = 0.50$ (hexanes/ethyl acetate 3/1), 99% ee, $[\alpha]^{20}_{D} = -16.48$ (*c* 1.42, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.85-7.75 (m, 2H), 7.40-7.30 (m, 2H), 6.56-6.37 (m, 5H), 6.32-6.24 (m, 1H), 6.16 (s, 1H), 4.50 (t, *J* = 5.7 Hz, 1H), 4.07 (dd, *J* = 13.4, 6.1 Hz, 1H), 3.72 (dd, *J* = 13.4, 5.7 Hz, 1H), 3.25-3.14 (m, 1H), 3.13-2.96 (m, 4H), 2.95-2.83 (m, 2H), 2.81-2.70 (m, 1H), 2.46 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.6, 140.4, 139.5, 139.2, 138.1, 136.7, 135.2, 135.2, 133.4, 133.4, 133.3, 132.4, 132.2, 129.9, 128.7, 127.3, 46.3, 35.3, 34.9, 34.3, 32.9, 21.7. HPLC: Chiracel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 60/40, flow = 0.7 mL/min, retention time 15.9 min and 26.4 min (major). HRMS: Calculated for C₂₄H₂₅NaNO₂S [M+Na]⁺ 414.1498, found: 414.1502.



To a stirred mixture of aldimine (-)-1a (78 mg, 0.20 mmol, 99% ee) in methanol (10 mL) was added aqueous HCl (10 mL, 2 M in water). The resulting mixture was stirred at 60 °C for 8 h. The reaction was quenched by addition of saturated aqueous sodium bicarbonate solution (30 mL), and then extracted with dichloromethane (20 mL×3). The combined organic layer was washed with brine, dried by anhydrous sodium sulfate and filtered. The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the corresponding product (-)-**5a**.

(*R_p*)-(-)-4-Formyl[2.2]paracyclophane (5a): 45 mg, 95% yield, white solid, known compound,^[6] R_f = 0.30 (hexanes/ethyl acetate 80/1), 99% ee, $[\alpha]^{20}{}_{\rm D}$ = -181.65 (*c* 0.96, CHCl₃), [lit.^[6]: $[\alpha]^{20}{}_{\rm D}$ = 184 (*c* 0.41, CHCl₃) for 98.7% ee (*S_p*)]. ¹H NMR (400 MHz, CDCl₃) δ 9.95 (s, 1H), 7.02 (d, *J* = 1.9 Hz, 1H), 6.76-6.69 (m, 1H), 6.62-6.54 (m, 2H), 6.53-6.48 (m, 1H), 6.46-6.36 (m, 2H), 4.18-4.05 (m, 1H), 3.31-2.90 (m, 7H). ¹³C NMR (100 MHz, CDCl₃) δ 192.0, 143.3, 140.8, 139.6, 139.5, 138.2, 136.7, 136.4, 136.2, 133.4, 133.0, 132.5, 132.3, 35.4, 35.2, 35.1, 33.7. HPLC: Chiracel IC column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 0.7 mL/min, retention time 12.6 min (major) and 15.4 min.



To a solution of (-)-1a (78 mg, 0.20 mmol, 99% ee) in dry tetrahydrofuran (7.0 mL) was added PhMgBr (0.50 mL, 2 M in THF, 1.0 mmol) at 0 °C under nitrogen. The reaction mixture was allowed to warm to room temperature and stirred at room temperature overnight. Water (5.0 mL) was added, and extracted with ethyl acetate (10 mL×3). The combined organic layer was washed with brine, dried by anhydrous sodium sulfate and filtered. The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford sulfonamide **3a**.

To a solution of the above sulfonamide **3a** in dry *N*,*N*-dimethylformamide (5.0 mL) was added sodium hydride (24 mg, 1.0 mmol, 60% wt.) at 0 °C, and then allyl bromide (121 mg, 86.0 μ L, 1.00 mmol) was added dropwise. The reaction mixture was warmed to room temperature and stirred at room temperature overnight. Water (10 mL) was added, and extracted with ethyl acetate (10 mL×3). The combined organic layer was washed with brine, dried by anhydrous sodium sulfate and filtered. The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the desirable product (-)-**2a**.

(-)-*N*-Allyl-*N*-{[2.2]paracyclophan-4-yl(phenyl)methyl}-4-methylbenzenesulfonamide (2a): 99 mg, 98% yield, white solid, new compound, $R_f = 0.50$ (hexanes/ethyl acetate 10/1), 99% ee, $[\alpha]^{20}_D = -129.07$ (*c* 1.37, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.97-7.89 (m, 2H), 7.43-7.33 (m, 2H), 7.24-7.17 (m, 1H), 7.17-7.07 (m, 3H), 6.91-6.84 (m, 2H), 6.77-6.70 (m, 1H), 6.63-6.47 (m, 4H), 6.34 (d, *J* = 7.6 Hz, 1H), 6.20 (s, 1H), 4.80-4.68 (m, 1H), 4.64-4.48 (m, 2H), 4.04-3.91 (m, 1H), 3.53-3.39 (m, 1H), 3.30-3.15 (m, 2H), 3.08-2.86 (m, 4H), 2.53-2.37 (m, 2H), 2.50 (s, 3H). ¹³C NMR (100 MHz, CDCl₃) δ 143.5, 140.9, 140.0, 139.5, 139.2, 138.9, 138.6, 136.9, 136.0, 134.0, 133.7, 132.7, 132.3, 132.2, 131.4, 130.5, 129.7, 129.4, 128.4, 128.2, 128.0, 117.2, 64.6, 48.5, 35.5, 35.4, 34.6, 34.5, 21.7. HPLC: Chiracel AD-H column, 254 nm, 30 °C, *n*-Hexane/ *i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 6.8 min and 7.4 min (major).



To a suspension of (-)-**1a** (78 mg, 0.20 mmol, 99% ee) and zinc powder (24 mg, 0.36 mmol) in dry *N*,*N*-dimethylformamide (4.0 mL) was slowly added allyl bromide (36 mg, 26.0 μ L, 0.30 mmol) under nitrogen and the solution was stirred at 80 °C for 4 h. After the addition of saturated aqueous ammonium chloride solution (15 mL), the solution was extracted with ethyl acetate (15 mL×3). The combined organic layer was washed by brine, dried by anhydrous sodium sulfate and filtered. The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford sulfonamide **S2**.

To a solution of the above sulfonamide **S2** in dry *N*,*N*-dimethylformamide (2.0 mL) was added sodium hydride (30 mg, 0.75 mmol, 60% wt.) at room temperature, and then allyl bromide (73 mg, 50.0 μ L, 0.60 mmol) was added dropwise. The reaction mixture was stirred at room temperature overnight. Water (10 mL) was added, and extracted with ethyl acetate (15 mL×3). The combined organic layer was washed with brine, dried by anhydrous sodium sulfate and filtered. The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the compound **S3**.

To a solution of crude product **S3** in dichloromethane (5.0 mL) was added Grubb's second generation catalyst (17 mg, 0.02 mmol, 10 mol%). The mixture was refluxed overnight and concentrated. The resulting crude product was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the desiable product (R_p ,R)-(+)-**6**.

(+)-2-([2.2]Paracyclophan-4-yl)-1-tosyl-1,2,3,6-tetrahydropyridine (6): 80 mg, 90% yield, white solid, mp = 149-151 °C, new compound, $R_f = 0.30$ (hexanes/ethyl acetate 10/1), 99% ee, $[\alpha]^{20}_D = 73.22$ (*c* 0.62, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.94-7.82 (m, 2H), 7.41-7.32 (m, 2H), 7.12-7.01 (m, 1H), 6.64 (s, 1H), 6.63-6.54 (m, 2H), 6.54-6.48 (m, 1H), 6.45-6.37 (m, 2H), 5.48-5.34 (m, 3H), 4.23-3.91 (m, 2H), 3.35-3.23 (m, 2H), 3.22-2.93 (m, 6H), 2.51-2.34 (m, 1H), 2.46 (s, 3H), 2.26-2.14 (m, 1H). ¹³C NMR (100 MHz, CDCl₃) δ 143.4, 140.0, 139.9, 139.5, 139.0, 137.8, 136.0, 136.0, 133.4, 132.5, 132.3, 131.1, 129.9, 129.7, 127.1, 123.2, 122.1, 52.2, 42.1, 35.3, 34.9, 33.8, 31.2, 21.6. HPLC: Chiracel AD-H column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 90/10, flow = 1.0 mL/min, retention time 13.4 min (major) and 14.2 min. HRMS: Calculated for C₂₈H₃₀NO₂S [M+H]⁺ 444.1992, found: 444.1971.

Removal of the Tosyl Group



Sodium (92 mg, 4.0 mmol, 20 equiv; washed free of oil in hexane) was added to a vigorously stirred suspension of naphthalene (513 mg, 4.00 mmol) in tetrahydrofuran (15 mL) at room temperature. The resulting green suspension was stirred for 4 h at room temperature, then was transferred to a solution of (S_p,S) -(+)-**2a** (101 mg, 0.20 mmol) in tetrahydrofuran (10 mL) at -78 °C. The dark green solution was stirred at -78 °C for 2 h. Water (20 mL) was added to the solution at -78 °C. The reaction mixture was extracted with ethyl acetate (20 mL×3). The combined organic layer was washed with brine, dried by anhydrous sodium sulfate and filtered. The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the desirable product (+)-**7**.

(+)-*N*-([2.2]Paracyclophan-4-yl(phenyl)methyl)prop-2-en-1-amine (7): 67 mg, 94% yield, white solid, mp = 99-101 °C, new compound, $R_f = 0.35$ (hexanes/ethyl acetate 10/1), 99% ee, $[\alpha]^{20}_D = 174.32$ (*c* 0.30, CHCl₃). ¹H NMR (400 MHz, CDCl₃) δ 7.25-7.19 (m, 2H), 7.18-7.12 (m, 3H), 6.88 (s, 1H), 6.67-6.60 (m, 1H), 6.60-6.49 (m, 3H), 6.48-6.42 (m, 1H), 6.35 (d, *J* = 7.6 Hz, 1H), 6.19-6.05 (m, 1H), 5.42-5.23 (m, 2H), 4.76 (s, 1H), 3.41-3.02 (m, 9H), 2.77-2.63 (m, 1H). ¹³C NMR (100 MHz, CDCl₃) δ 143.9, 142.9, 140.1, 139.8, 139.6, 137.2, 136.6, 135.7, 133.8, 133.1, 132.0, 131.5, 129.7, 129.3, 128.4, 128.1, 126.9, 116.6, 62.9, 50.4, 35.5, 35.4, 34.6, 34.1. HPLC: Chiracel IA column, 254 nm, 30 °C, *n*-Hexane/*i*-PrOH = 98/2, flow = 1.0 mL/min, retention time 5.0 min (major) and 5.7 min. HRMS: Calculated for C₂₆H₂₈N [M+H]⁺ 354.2216, found: 354.2213.

5. Scale-up Reaction



A schlenk tube (50 mL) was charged with Pd(OCOCF₃)₂ (16.6 mg, 0.05 mmol, 5 mol%) and (R_p ,S)-L2 (24.8 mg, 0.05 mmol, 5 mol%) under nitrogen, and degassed anhydrous acetone (5.0 mL) was added. The mixture was stirred at room temperature for 1 h. The solvent was removed under vacuum to give the catalyst. Then substrate *rac*-**1a** (389 mg, 1.00 mmol), phenylboronic acid (122 mg, 1.0 mmol) and 2,2,2-trifluoroethanol (20 mL) were added into the tube under nitrogen. The mixture was heated to 60 °C. After stirring at 60 °C for 15 h, the reaction mixture was cooled to room temperature, and the solvent was removed by rotary evaporation. The resulting mixture was dried under vacuum and the conversion of *rac*-**1a** (51% conv.) was confirmed by ¹H NMR analysis with benzyl ether as internal standard. The solvent was removed in *vacuo*, recovered material (-)-**1a** (165.5 mg, 43% yield with 96.0% ee) and addition product **3a** were isolated by column chromatography on silica gel using hexanes and ethyl acetate as eluent.

To a solution of the addition product **3a** in dry *N*,*N*-dimethylformamide (DMF, 8.0 mL) was added sodium hydride (48 mg, 1.20 mmol, 60% wt.) at 0 °C, and then allyl bromide (242 mg, 0.17 mL, 2.00 mmol) was added dropwise. The reaction mixture was warmed to room temperature and stirred at room temperature for 4 h. Water (10 mL) was added, and extracted with ethyl acetate (15 mL×3). The combined organic layer was washed with brine, dried by anhydrous sodium sulfate and filtered, concentrated in *vacuo* and analyzed by crude ¹H NMR to determine diastereomeric ratio (> 20:1 dr). The solvent was removed in *vacuo*, and the residue was purified by column chromatography on silica gel using hexanes and ethyl acetate as eluent to afford the product (+)-**2a** (255.0 mg, 50% yield with 98.1% ee).

6. Determination of Absolute Configuration

6.1 Determination of the Absolute Configuration of (+)-2a

To determine the absolute configuration of (+)-2a, a single crystal of the 2a was grown from its solution in dichloromethane and *n*-hexane. *n*-Hexane (3.0 mL) was slowly added into the solution of 2a in dichloromethane (3.0 mL) at room temperature, then the solvent was slowly evaporated and single crystal was obtained after 6 days. The structure in Figure S1 showed the absolute configuration of (+)-2a is (S_p ,S). The CCDC number is 1891926. These details can be obtained free of charge *via* www.ccdc.com. ac.uk/data_request/cif from the Cambridge Crystallographic Data Centre.



Figure S1. X-ray Crystallographic Analysis of (S_p,S)-2a

6.2 Determination of the Absolute Configuration of (+)-21

To determine the absolute configuration of (+)-2l, a single crystal of the 2l was grown from its solution in dichloromethane and *n*-hexane. *n*-Hexane (1.5 mL) was slowly added into the solution of 2l in dichloromethane (1.5 mL) at room temperature, then the solvent was slowly evaporated and single crystal was obtained after 3 days. The structure in Figure S2 showed the absolute configuration of (+)-2l is (S_p ,S). The CCDC number is 1910494. These details can be obtained free of charge *via* www.ccdc.com. ac.uk/data_request/cif from the Cambridge Crystallographic Data Centre.



Figure S2. X-ray Crystallographic Analysis of (S_p,S)-2l

6.3 Determination of the Absolute Configuration of (+)-6

To determine the absolute configuration of (+)-6, a single crystal of the 6 was grown from its solution in dichloromethane and *n*-hexane. *n*-Hexane (3.0 mL) was slowly added into the solution of 6 in dichloromethane (3.0 mL) at room temperature, then the solvent was slowly evaporated and single crystal was obtained after 5 days. The structure in **Figure S3** showed the absolute configuration of (+)-6 is (R_p ,R). The CCDC number is 1917426. These details can be obtained free of charge *via* www.ccdc.com. ac.uk/data_request/cif from the Cambridge Crystallographic Data Centre.



Figure S3. X-ray Crystallographic Analysis of (R_p, R) -6

7. References

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-21.75

13C NMR YZ-4-22 in CDCl3









13C NMR YZ-4-84 in CDCI3



~21.78





13C NMR YZ-6-7 in CDCl3





rac-1c ¹³C NMR (100 MHz, CDCl₃)





S25


































1H NMR YZ-4-74BTM in CDCI3







1H NMR YZ-4-74BTM in CDCI3



-62.52

19F NMR YZ-4-74BTM in CDCI3







1H NMR YZ-4-72ATM in CDCI3







1H NMR YZ-4-74ATM in CDCI3





19F NMR YZ-4-74ATM in CDCI3 Ār ∠Ts $Ar = 3-FC_6H_4$ **2h** ¹⁹F NMR (376 MHz, CDCl₃) 10 0 -10 -20 -30 -40 -50 -60 -70 -80 -90 -100 -110 -120 -130 -140 -150 -160 -170 -180 -190 -200 -210 f1 (ppm)









Ar = 3-CIC₆H₄ **2i** ¹H NMR (400 MHz, CDCl₃)





13C NMR YZ-4-77TM in CDCI3









1H NMR YZ-4-65BTM in CDCI3









1H NMR YZ-4-68TM in CDCI3







1H NMR YZ-4-85ATM in CDCI3









1H NMR YZ-4-85BTM in CDCI3







1H NMR YZ-6-8TM in CDCI3











S58











-192.02

н

5a ¹³C NMR (100 MHz, CDCl₃)







35.45 35.40 734.57 34.49



S63









1H NMR YZ-5-4 in CDCI3



7 ¹H NMR (400 MHz, CDCl₃)

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Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1003079.D Sample Name: YZ-4-58BSM(RAC)

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Acq. Instrument	:	仪器 1 Location : Vial 1
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		Inj Volume : 5.000 µl
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		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:07:23
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n
		TO





-----Area Percent Report

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Dilution:		:	1.0000	
Use Multiplier & D	ilution	Factor with	ISTDs	
Signal 1: VWD1 A,	Waveleng	th=254 nm		
Peak RetTime Type	Width	Area	Height	Area
Peak RetTime Type # [min]	Width [min]	Area [mAU*s]	Height [mAU]	Area %
Peak RetTime Type # [min] 	Width [min] 	Area [mAU*s] 	Height [mAU] 	Area *
Peak RetTime Type # [min] 1 16.115 BV	Width [min] 0.2366	Area [mAU*s] 1152.96143	Height [mAU] 76.14727	Area % 49.9519
Peak RetTime Type # [min] 1 16.115 BV 2 18.177 VB	Width [min] 0.2366 0.2685	Area [mAU*s] 1152.96143 1155.18225	Height [mAU] 76.14727 67.21445	Area * 49.9519 50.0481
Peak RetTime Type # [min] 1 16.115 BV 2 18.177 VB	Width [min] 0.2366 0.2685	Area [mAU*s] 1152.96143 1155.18225	Height [mAU] 76.14727 67.21445	Area * 49.9519 50.0481



Page 1 of 2







-----Area Percent Report

		rea reroem	C Report		
Sorted By Multiplier: Dilution: Use Multiplier & S	: Dilution	Signal : : Factor wit	1.0000 1.0000 n ISTDs		N ^{-Ts}
Signal 1: VWD1 A,	Waveleng	th=254 nm			
Peak RetTime Type	Width	Area	Height	Area	(-)-1a
# [min]	[min]	[mAU*s]	[mAU]	*	Kinetic resolution
					Triffette resolution
1 16.616 VB	0.2488	61.41887	3.79507	3.1303	from PhB(OH) ₂
2 18.867 BB	0.2875	1900.67249	102.88990	96.8697	()2
	_				Dave 1 of 2

仪器 1 5/30/19 8:45:32

Page 1 of 2

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002465.D Sample Name: YZ-4-60A

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		Inj Volume : 5.000 µl					
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M					
Last changed	:	1/15/19 13:26:04					
		(modified after loading)					
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M					
Last changed	:	5/30/19 10:10:26					
		(modified after loading)					
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n					
		The second se					



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Dilution:		:	1.0000	
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Signal 1: VWD1 A,	Wavelen(gth=254 nm		
Peak Retlime Type	Width	Area	Height	Area
# [min]	[min]	[mAU*s]	[mAU]	*
1 16.672 BB	0.2563	293.14850	17.42010	9.9678
2 18.944 BB	0.2912	2647.81104	140.90161	90.0322
仪器 1 5/30/19 10:10:3	2			



Kinetic resolution from 2-MeC₆H₄B(OH)₂

Page 1 of 2

Data File C:\CHEM32\l Sample Name: ¥Z-4-60B	\DATA\2H0U2019\\$IG1002469.D
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Acq. Instrument :	仪器 1 Location : Vial 91
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	Inj Volume : 5.000 μl
Acq. Method :	C:\CHEM32\1\METHODS\DEF_LC11.M
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	(modified after loading)
Analysis Method :	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed :	5/30/19 9:51:56
	(modified after loading)
Sample Info :	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n



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Area Percent Report Sorted By Signal : : 1.0000 : 1.0000 Multiplier: Dilution: Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area

#	[min]	[min]	[mAU*s]	[mAU]	*
1	16.607 VB	0.2640	56.11112	3.20973	4.4128
2	18.868 BV	0.2869	1215.42883	65.99891	95.5872

仪器 1 5/30/19 9:52:11

from 3-MeC₆H₄B(OH)₂ Page 1 of 2

(-)-1a

Kinetic resolution

≪_N∠Ts

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002527.D Sample Name: YZ-4-62

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		Inj Volume : 10.000 µl						
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M						
Last changed	:	1/22/19 16:31:20						
		(modified after loading)						
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M						
Last changed	:	5/30/19 10:18:44						
		(modified after loading)						
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n						
		1						





-----Area Percent Report

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Multiplier:		:	1.0000	
Dilution:		:	1.0000	
Use Multiplier & D:	ilution	Factor with	n ISTDs	
Signal 1: VWD1 A, U	Javelenç	gth=254 nm		
Peak RetTime Type	Width	Area	Height	Area
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-				
1 16.536 BB	0.2701	334.76022	18.94763	5.9673
2 18.779 BB	0.3074	5275.15674	261.39069	94.0327
器 1 5/30/19 10:18:53	3			



Kinetic resolution from 4-MeC₆H₄B(OH)₂

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Page 1 of 2

Data File C:\CHEM32 Sample Name: YZ-4-6	\1 5A	\DATA\ZHOU2019\SIG1002562.D
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Acq. Instrument	:	仪器 1 Location : Vial 1
Injection Date	:	2/23/19 14:27:02
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M
Last changed	:	2/23/19 14:06:41
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:21:45
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n
		n





Area Percent Report Sorted By Signal . Multiplier: : 1.0000 Dilution: : 1.0000 Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm

Peak RetTime Type Width Height Area Area # [min] [min] [mAU*s] [mAU] * 1 16.398 BB 0.2462 12.59610 7.72629e-1 0.3868 2 18.586 BB 0.2806 3243.52002 181.41284 99.6132

仪器 1 5/30/19 10:21:50

(-)-1a

≷_N∕Ts

Kinetic resolution from 4-FC₆H₄B(OH)₂

Page 1 of 2

Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002673.D Sample Name: YZ-4-74B

	==	
Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location : Vial 1
Injection Date	:	3/11/19 21:25:13
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M
Last changed	:	3/11/19 21:22:09
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:53:00
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHOU2019\SIG1002673.D)



Area Percent Report

Sorted By	:	Signal		
Multiplier:		:	1.0000	
Dilution:		:	1.0000	
Use Multiplier	& Dilution	Factor wit	h ISTDs	

Signal 1: VWD1 A, Wavelength=254 nm

I	e ak	RetTime	Type	Width	Area	Height	Area	
	#	[min]		[min]	[mAU*s]	[mAU]	*	
-								
	1	16.132	ΒV	0.2730	24.26760	1.32900	0.7670	
	2	18.275	VV	0.2746	3139.73730	177.32698	99.2330	



Page 1 of 2

(-)-1a

Kinetic resolution

from 4-CF₃C₆H₄B(OH)₂

≿_N∠Ts



Acq. Operator	:				
Acq. Instrument	:	仪器 1 Location: Vial 1			
Injection Date	:	3/7/19 14:52:57			
		Inj Volume : 5.000 µl			
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	3/7/19 13:49:58			
		(modified after loading)			
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	5/30/19 10:37:16			
		(modified after loading)			
Sample Info	:	AD-3, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm			

Additional Info : Peak(s) manually integrated VWD1A Wavelength=254nm (ZHOU2019%)G1002634.D)



Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier & Signal 1: VWD1 A	: Sign : Dilution Factor : ., Wavelength=254 p	al 1.0000 1.0000 with ISTDs	
Peak RetTime Typ	e Width Area	Height Area	
# [min]	[min] [mAU*s] [mAU] %	Kinot
	-		Killer
1 16.402 BB	0.2663 4.59	826 2.65129e-1 0.2218	from 4-
2 18.608 BB	0.2771 2068.83	301 115.40721 99.7782	irein 1



(-)-**1a** Kinetic resolution from 4-CIC₆H₄B(OH)₂

仪器 1 5/30/19 10:37:28

Page 1 of 2
Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002672.D Sample Name: YZ-4-74A

Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location : Vial 1
Injection Date	:	3/11/19 20:48:19
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M
Last changed	:	3/11/19 20:25:14
		(modified after loading)
Analysis Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:44:02
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHOU2019\SIG1002672.D)



Area Percent Report

Sorted By	:	Sig	nal
Multiplier:		:	1.0000
Dilution:		:	1.0000
Use Multiplier	& Dilution	Factor	with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	*
1	16.207	VV	0.6405	16.37306	3.20245e-1	1.3236
2	18.307	ΒV	0.2758	1220.64233	68.52013	98.6764



Page 1 of 2

(-)-**1a** Kinetic resolution

from 3-FC₆H₄B(OH)₂

≈_N∠Ts

Data File C:\CHEM32 Sample Name: YZ-4-7	\1 7	\DATA\ZHOU2019\SIG100270	2.D
Acq. Operator	:		
Acq. Instrument	:	仪器 1	Location : Vial 1
Injection Date	:	3/16/19 16:34:44	
			Inj Volume : 5.000 µl
Acc. Method		C:\CHEM32\l\METHODS\DEE	LCII.M

nog. neenou	•	C. JOILINGS JI JILLINGS JELL_BOIL.IN
Last changed	:	3/16/19 16:28:23
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 11:00:32
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm





Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier « Signal 1: VWD1 A,	: Dilution , Waveleng	Signal : Factor with th=254 nm	1.0000 1.0000 ISTDs		N-Ts
Peak RetTime Type # [min] 1 16.331 BB 2 18.511 BB	<pre>Width [min] - 0.2472 0.2764</pre>	Area [mAU*s] 17.21958 2096.09521	Height [mAU] 1.09678 117.34330	Area % 0.8148 99.1852	(-)- 1a Kinetic resolution from 3-ClC ₆ H ₄ B(OH) ₂

仪器 1 5/30/19 11:00:40

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002563.D Sample Name: YZ-4-65B

Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location: Vial 1
Injection Date	:	2/23/19 14:49:53
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	2/23/19 14:48:56
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:27:42
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n
		m





-----Area Percent Report

Sorted By	:	Signal		
Multiplier:		:	1.0000	
Dilution:		:	1.0000	
Use Multiplier & D	ilution	Factor with	n ISTDs	
Signal 1: WWD1 A,	Wavelen	gth=254 nm		
Peak RetTime Type	Width	Area	Height	Area
# [min]	[min]	[mAU*s]	[mAU]	*
1 16.335 BB	0.2425	347.60333	22.21527	8.0739
2 18.525 VB	0.2768	3957.66699	221.08186	91.9261
器 1 5/30/19 10:27:4	7			



Kinetic resolution from 4-MeOC₆H₄B(OH)₂



Page 1 of 2

Data File C:\CHEM32\ Sample Name: YZ-4-68	,1\DATA\ZHOU2019\SIG1002615.D 3	
Acq. Operator	:	
Acc. Instrument	: 仪器 1	Location : Vial 1

Acq. Instrument	:	仪器 l Location : Vial l
Injection Date	:	3/2/19 19:08:27
		Inj Volume : 5.000 μl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	3/2/19 18:34:41
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:34:34
		(modified after loading)







Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier « : Signal 1: VWD1 A,	: Dilution Waveleng	Signal : Factor with th=254 nm	1.0000 1.0000 ISTDs		N-Ts
Peak RetTime Type # [min] 	Width [min] 0.2409 0.2793	Area [mAU*s] 329.95752 3406 98022	Height [mAU] 21.27563	Area % 8.8296	(-)- 1a Kinetic resolution from 2-naphthylB(OH) ₂

仪器 1 5/30/19 10:34:39

Data File C:\CHEM32\1\DATA\ZHOU2018\SIG1002068.D Sample Name: YZ-4-34A(RAC)

Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location: Vial 91
Injection Date	:	12/8/18 10:42:08
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	12/8/18 10:40:33
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 9:41:08
		(modified after loading)
Sample Info	:	AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n
		70





Area Percent Report

Sorted By		Simal		
Multiplier:	•	:	1.0000	
Dilution:			1.0000	
Use Multiplier & D	ilution	Factor with	i ISTDs	
Signal 1: VWDl A,	Wavelenç	th=254 nm		
Peak RetTime Type	Width	Area	Height	Area
# [min]	[min]	[mAU*s]	[mAU]	*
1 6.803 VV	0.1468	616.60956	64.94445	49.8919
2 7 446 VB	0.1690	619.28156	57.78165	50.1081
5 71440 VD				

Ph N^{-Ts}

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Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002454.D



Area Percent Report



仪器 1 5/30/19 9:44:53

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Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002511.D Sample Name: YZ-4-60ATM-rac

Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location: Vial 91
Injection Date	:	1/20/19 13:43:52
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	1/20/19 13:36:11
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:13:31
		(modified after loading)
Sample Info	:	AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n
		70





Area Percent Report

Sorted By	:	Signal		
Multiplier:		:	1.0000	
Dilution:		:	1.0000	
Use Multiplier « D	ilution	Factor with	n ISTDs	
Signal 1: VWD1 A,	Wavelenç	gth=254 nm		
Peak RetTime Type	Width	Area	Height	Area
# [min]	[min]	[mAU*s]	[mAU]	*
1 5.288 VB	0.1161	1773.82800	234.77663	49.8177
2 7.045 VB	0.1554	1786.80872	174.56734	50.1823
仪器 1 5/30/19 10:13:3	7			

 $Ar = 2-MeC_6H_4$ rac-2b



Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location : Vial 91
Injection Date	:	1/20/19 14:00:58
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	1/20/19 13:53:05
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:15:57
		(modified after loading)
Sample Info	:	AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 m
		-





Area Percent Report



Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002518.D Sample Name: YZ-4-60BTM-rac

Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location : Vial 91
Injection Date	:	1/20/19 16:14:19
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	1/20/19 16:08:14
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 9:56:17
		(modified after loading)
Sample Info	:	IC, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 nm

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHOU2019\SIG1002518.D)



Area Percent Report

Sorted By		Sim	nal
Multiplier:		:	1.0000
Dilution:		:	1.0000
Use Multiplier &	Dilution	Factor	with ISTDs
Signal 1: VWD1 A	, Wavelenç	gth=254	nm

仪器 1 5/30/19 9:56:37

Peak.	Retlime	Type	Width	Area	Height	Area	
#	[min]		[min]	[mAU*s]	[mAU]	*	
1	15.877	BB	0.4098	2122.69409	80.99484	49.8428	
2	21.863	BB	0.5953	2136.08423	55.22515	50.1572	



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Sample Info : IC, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 nm





Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier &	: Dilution	Signal : : Factor with	1.0000 1.0000 h ISTDs		Âr
Signal 1: VWD1 A,	, Waveleng	th=254 nm			
Peak RetTime Type # [min]	e Width [min]	Area [mAU*s] 	Height [mAU]	Area %	Ar = 3-MeC ₆ H∠ (+)- 2c
1 15.810 BB 2 21.749 VB	0.4197	45.00853 4762.27295	1.66333 122.14498	0.9363 99.0637	(1) 20

仪器 1 5/30/19 10:00:42

Page 1 of 2

∠Ts

Data File C:\CHEM32\1\DATA\ZHOU-19\YZN011706.D Sample Name: YZ-4-62TMrac

Acq. Operator	:					
Acq. Instrument	:	Instrument 1	Location	:	-	
Injection Date	:	1/24/2019 8:46:16 PM				
Acq. Method	:	C:\CHEM32\1\METHODS\DEF LC11.M				
Last changed	:	1/24/2019 8:28:38 PM				
		(modified after loading)				
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M				
Last changed	:	5/30/2019 11:42:24 AM				
		(modified after loading)				
Sample Info	:	AD-H, Hexane/i-PrOH = 90/10, 1	.0 mL/min.	, 30	oC,	254nm



Sorted By : Signal : 1.0000 : 1.0000 Multiplier: Dilution: Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area $Ar = 4 - MeC_6H_4$ 2 8.517 BB 0.1819 825.60114 70.45312 49.9218 rac-2d 1653.78821 155.63210 Totals : -----

*** End of Report ***

Acd. Obergrot						
Acq. Instrument	:	Instrument 1	Location	:	-	
Injection Date	:	1/24/2019 9:20:35 PM				
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M				
Last changed	:	1/24/2019 9:08:41 PM				
		(modified after loading)				
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M				
Last changed	:	5/30/2019 2:12:38 PM				
		(modified after loading)				
Sample Info	:	AD-H, Hexane/i-PrOH = 90/10, 1.	.0 mL/min,	, 30	oC,	254nm







Instrument 1 5/30/2019 11:42:28 AM

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∕Ts

Instrument 1 5/30/2019 2:12:44 PM

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002572.D Sample Name: YZ-4-65ATM(RAC)

Acq. Operator :	
Acq. Instrument :	仪器 1 Location: Vial 1
Injection Date :	2/26/19 9:40:56
	Inj Volume : 5.000 μl
Acq. Method :	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed :	2/26/19 9:37:52
	(modified after loading)
Analysis Method :	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed :	5/30/19 10:23:50
	(modified after loading)
Sample Info :	IB, n-hexane/i-PrOH =80/20, 1.0 mL/min, 30 oC, 254 nm





Area Percent Report

Sorted By		Sim	nal
Multiplier:	-	:	1.0000
Dilution:		:	1.0000
Use Multiplier	& Dilution	Factor	with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

仪器 1 5/30/19 10:24:02

Peak	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	*
1	5.088	VV	0.1074	1295.38733	190.37517	49.6614
2	5.427	VV	0.1076	1313.04907	183.26695	50.3386







Acq. Operator	:							
Acq. Instrument	:	仪器 1		Location	:	Vial	1	
Injection Date	:	2/26/19 10:10:55						
			In	j Volume	:	5.00)O μ)	1
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11	. M					
Last changed	:	2/26/19 9:55:52						
		(modified after loading)						
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11	. M					
Last changed	:	5/30/19 10:26:07						
		(modified after loading)						
Sample Info	:	IB, n-hexane/i-PrOH =80/20,	1.0	mL/min,	30	οC,	254	nn





Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier & Signal 1: VWD1 A	: Dilution , Waveleng	Signal : Factor with th=254 nm	1.0000 1.0000 A ISTDs		Ar
Peak RetTime Typ # [min]	e Width [min]	Area [mAU*s]	Height [mAU]	Area %	$Ar = 4 - FC_6 I$
1 5.111 VV 2 5.450 VV	0.1058	976.24976 19.14368	146.41737 2.04433	98.0768 1.9232	(+)-26



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仪器 1 5/30/19 10:26:12

Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002689.D Sample Name: YZ-4-74BTM(RAC)

Acq. Operator :	
Acq. Instrument :	: 仪器 1 Location : Vial 1
Injection Date :	3/13/19 15:35:31
	Inj Volume : 5.000 µl
Acq. Method :	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed :	3/13/19 15:25:54
	(modified after loading)
Analysis Method :	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed :	5/30/19 10:54:57
	(modified after loading)
Sample Info :	IB, n-hexane/i-PrOH =90/10, 1.0 mL/min, 30 oC, 254 nm

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHO U2019\S1G1002689.D)



Area Percent Report

Sorted By	:	Sign	nal
Multiplier:		:	1.0000
Dilution:		:	1.0000
Use Multiplier	& Dilution	Factor	with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak R	etTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	25
-						
1	6.035	VV	0.1249	1574.70276	197.71895	50.1111
2	6.689	VB	0.1333	1567.71960	180.62619	49.8889



Sample Info : IB, n-hexane/i-PrOH =90/10, 1.0 mL/min, 30 oC, 254 nm

Additional Info : Peak(s) manually integrated VWD1A Wavelength=254nm (ZHOU2019%)G1002690.D)



Area Percent Report

Sorted By	:	Signal	
Multiplier:		: 1.0	0000
Dilution:		: 1.0	0000
Jse Multiplier	& Dilution	Factor with IS	STDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak RetTime Type Width Area	Height Area
# [min] [min] [mAU*s]	[mAU] %
-	
1 5.938 VV 0.1197 3623.50244	461.16406 99.3273
2 6.598 VV 0.1821 24.54160	1.90772 0.6727



(+)-**2f**

仪器 1 5/30/19 10:55:10

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仪器 1 5/30/19 10:58:34

Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002649.D Sample Name: YZ-4-72ATM RAC

	= :	
Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location: Vial 1
Injection Date	:	3/9/19 9:45:10
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	3/9/19 9:38:16
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:39:35
		(modified after loading)
Sample Info	:	IB, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHOU2019\SIG1002649.D)



Area Percent Report

Sorted By	:	Sign	nal
fultiplier:		:	1.0000
ilution:		:	1.0000
Jse Multiplier	& Dilution	Factor	with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak	RetTime	Type	Width	Area	Height	Area	
#	[min]		[min]	[mAU*s]	[mAU]	*	
1	6.386	VV	0.1217	1653.46570	214.99780	49.9937	
2	6.950	VV	0.1280	1653.88330	200.99446	50.0063	



Ar

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002654.D Sample Name: YZ-4-72ATM

Acq. Operator	:				
Acq. Instrument	:	仪器 1 Location: Vial 1			
Injection Date	:	3/9/19 14:20:02			
		Inj Volume : 5.000 µl			
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	3/9/19 14:17:28			
		(modified after loading)			
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	5/30/19 10:41:45			
		(modified after loading)			
Sample Info	:	IB, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm			





Area Percent Report

Sorted By Multiplier: Dilution: Jse Multiplier &	: Dilution)	Signal : : Factor wit	1.0000 1.0000 n ISTDs] ₁
3ignal l: VWDl A	, Waveleng	th=254 nm			L	
Peak RetTime Typ	e Width	Area	Height	Area	Ar	=
# [min]	[min]	[mAU*s]	[mAU]	*	7.0	
	-					(
1 6.382 VV	0.1170 2	2098.80957	275.09128	99.4442		
2 6.951 VV	0.1537	11.73071	1.08846	0.5558		



仪器 1 5/30/19 10:39:44

Page 1 of 2

仪器 1 5/30/19 10:41:48

Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002685.D Sample Name: YZ-4-74ATM(RAC)

Acq. Operator :					
Acq. Instrument :	仪器 1 Location : Vial 1				
Injection Date :	3/13/19 11:35:41				
	Inj Volume : 5.000 µl				
Acq. Method :	C:\CHEN32\1\METHODS\DEF_LC11.M				
Last changed :	3/13/19 11:33:58				
	(modified after loading)				
Analysis Method :	C:\CHEN32\1\METHODS\DEF_LC11.M				
Last changed :	5/30/19 10:46:04				
	(modified after loading)				
Sample Info :	IA, n-hexane/i-PrOH =90/10, 1.0 mL/min, 30 oC, 254 nm				

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHO U2019\S1G1002685.D)



Area Percent Report

:	Signal					
	:	1.0000				
	:	1.0000				
& Dilution	Factor wit	h ISTDs				
	: « Dilution	: Signal : : & Dilution Factor wit	: Signal : 1.0000 : 1.0000 & Dilution Factor with ISTDs	: Signal : 1.0000 : 1.0000 & Dilution Factor with ISTDs	: Sigmal : 1.0000 : 1.0000 & Dilution Factor with ISTDs	: Sigmal : 1.0000 : 1.0000 & Dilution Factor with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak	RetTime	Type	Width	Area	Height	Area	
#	[min]		[min]	[mAU*s]	[mAU]	*	
1	6.444	VV	0.1419	1170.60938	128.92268	49.6684	
2	7.340	VB	0.1576	1186.24072	113.80168	50.3316	





Sample Info : IA, n-hexane/i-PrOH =90/10, 1.0 mL/min, 30 oC, 254 nm





Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier &	: Dilution	Signal : : Factor with	1.0000 1.0000 n ISTDs			ĺ
Signal 1: VWD1 A	., Waveleng	th=254 nm				Í
Peak RetTime Typ # [min]	e Width [min]	Area [mAll*s]	Height [mAII]	Area %	Ar =	=
	-					1
1 6.397 VV 2 7.238 VV	0.1351 0.1987	3452.54590 19.35161	390.52866 1.38728	99.4426 0.5574		`



仪器 1 5/30/19 10:46:10

Page 1 of 2

仪器 1 5/30/19 10:48:51

Data File C:\CHEM32\1\DATA\ZH0U-19\YZN012032.D Sample Name: YZ-4-77TM (RAC)

Acq. Operator	:					
Acq. Instrument	:	Instrument l Location : -				
Injection Date	:	3/18/2019 9:39:09 PM				
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M				
Last changed	:	3/18/2019 9:07:39 PM				
		(modified after loading)				
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M				
Last changed	:	5/30/2019 3:06:15 PM				
		(modified after loading)				
Sample Info	:	IC, Hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 nm				



Area Percent Report · Sorted By : Signal : 1.0000 Multiplier: ∠Ts Dilution: Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area # [min] [min] mAU *s [mAU] * $Ar = 3-CIC_6H_4$ 1 11.219 BB 0.2505 1088.86633 67.61742 49.9980 2 13.247 BB 0.3161 1088.95276 53.49617 50.0020 rac-**2i** Totals : 2177.81909 121.11358 -----

*** End of Report ***

Instrument 1 5/30/2019 3:06:24 PM

Page 1 of 1

Data File C:\CHEM32\1\DATA\ZH0U-19\YZN012033.D Sample Name: YZ-4-77TM

Acq. Operator	:					
Acq. Instrument	:	Instrument 1	Location	:	-	
Injection Date	:	3/18/2019 9:58:13 PM				
Acq. Method	:	C:\CHEM32\1\METHODS\DEF LC11.M				
Last changed	:	3/18/2019 9:53:52 PM				
		(modified after loading)				
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M				
Last changed	:	5/30/2019 3:12:11 PM				
		(modified after loading)				
Sample Info	:	IC, Hexane/i-PrOH = 80/20, 0.8	mL/min, 3	30 ol	C, 254 n	m







Instrument 1 5/30/2019 3:12:14 PM

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002579.D Sample Name: YZ-4-65BTMRAC

Acq. Operator	:				
Acq. Instrument	:	仪器 1 Location: Vial 1			
Injection Date	:	2/26/19 19:28:36			
		Inj Volume : 5.000 µl			
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M			
Last changed	:	2/26/19 19:26:35			
		(modified after loading)			
Analysis Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M			
Last changed	:	5/30/19 10:30:25			
		(modified after loading)			
Sample Info	:	AD-H, n-hexane/i-PrOH =80/20, 1.0 mL/min, 30 oC, 254 nm			

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHOU2019\SIG1002579.D)



Area Percent Report

Sorted By	:	Sign	nal
Multiplier:		:	1.0000
Dilution:		:	1.0000
Use Multiplier	& Dilution	Factor	with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak H	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	*
1	6.476	VV	0.1500	1364.39746	139.67671	49.8383
2	7.198	VB	0.1669	1373.24829	126.26674	50.1617



Ar



Acq. Operator								
Acq. Instrument	仪器 1 Location: Vial 1							
Injection Date	2/26/19 19:42:14							
	Inj Volume : 5.000 µl							
Acq. Method	C:\CHEN32\1\METHODS\DEF_LC11.M							
Last changed	2/26/19 19:37:55							
	(modified after loading)							
Analysis Method	C:\CHEN32\1\METHODS\DEF_LC11.M							
Last changed	5/30/19 10:32:27							
	(modified after loading)							
Sample Info	AD-H, n-hexane/i-PrOH =80/20, 1.0 mL/min, 30 oC, 254 nm							

Additional Info : Peak(s) manually integrated VWD1A Wavelength=254nm (ZHOU2019%)G1002580.D)



Area Percent Report _____

Sorted By Multiplier:	:	Signal :	1.0000		
Dilution:		:	1.0000		
Use Multiplier Signal 1: VWD1	& Dilution A, Wavelen(Factor with oth=254 nm	ı ISTDs		
Peak RetTime T	ype Width	Area	Height	Area	Ár = 4-Me
# [min]	[min]	[mAU*s]	[mAU]	*	
-					(+)-
1 6.516 V	B 0.1543	52.88145	5.21458	1.7472	()
2 7.235 B	B 0.1635	2973.78955	280.84409	98.2528	



-2j

仪器 1 5/30/19 10:30:28

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仪器 1 5/30/19 10:32:35

Data File C:\CHEM32\1\DATA\ZH0U-19\YZN011868.D Sample Name: YZ-4-68TMRAC

Acq. Operator	:							
Acq. Instrument	:	Instrument 1	Location	:	-			
Injection Date	:	3/4/2019 2:21:07 PM						
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M						
Last changed	:	3/4/2019 2:12:49 PM						
		(modified after loading)						
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M						
Last changed	:	5/30/2019 3:01:05 PM						
		(modified after loading)						
Sample Info	:	AD-3, Hexane/i-PrOH = 90/10, 1	.0 mL/min	, 30	oC,	254	nm	





-----*** End of Report ***

Instrument 1 5/30/2019 3:01:09 PM

Page 1 of 1

Data File C:\CHEM32\1\DATA\ZH0U-19\YZN011869.D Sample Name: YZ-4-68TM

Acq. Operator	:						
Acq. Instrument	:	Instrument 1	Location	:	-		
Injection Date	:	3/4/2019 2:42:07 PM					
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M					
Last changed	:	3/4/2019 2:37:25 PM					
		(modified after loading)					
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M					
Last changed	:	5/30/2019 3:03:12 PM					
		(modified after loading)					
Sample Info	:	AD-3, Hexane/i-PrOH = 90/10, 1	.0 mL/min,	30	oC,	254	nm







Instrument 1 5/30/2019 3:03:18 PM

Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1002783.D Sample Name: YZ-4-85SM(RAC)

Acq. Operator	:						
Acq. Instrument	:	仪器 1 Location: Vial 1					
Injection Date	:	3/28/19 15:10:36					
		Inj Volume : 5.000 µl					
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M					
Last changed	:	3/28/19 15:07:42					
		(modified after loading)					
Analysis Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M					
Last changed	:	5/30/19 11:10:50					
		(modified after loading)					
Sample Info	:	AD-H, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm					

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHOU2019\SIG1002783.D)



Area Percent Report

Sorted By	:	Signal	
Multiplier:		:	1.0000
Dilution:		:	1.0000
Use Multiplier	& Dilution	Factor wi	th ISTDs
····			

Signal 1: VWD1 A, Wavelength=254 nm

Peak	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	*
1	16.750	VB	0.3398	1703.34778	77.67725	50.0414
2	20.564	BB	0.4090	1700.52625	64.20832	49.9586

仪器 1 5/30/19 11:11:04

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rac-**1b**

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Ts

H₃C

Data File C:\CHEM32 Sample Name: YZ-4-8	\1 5A	\DATA\ZHOU2019\SIG100278 SM	5.D			
	==				==	
Acq. Operator	:					
Acq. Instrument	:	仪器 1	L	ocation	:	Vial 1
Injection Date	:	3/28/19 16:11:29				
-			Inj	Volume	:	5.000 µl
Acg Method		C • \ CHEM3 2 \ 1 \ METHODS \ DEE	LCII M			

Last changed	:	3/28/19 16:03:42
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 11:13:02
		(modified after loading)
Sample Info	:	AD-H, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm





Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier & I Signal 1: VWD1 A,	: Dilution Waveleng	Signal : Factor wit 9th=254 nm	1.0000 1.0000 h ISTDs		H ₃ C. N _{Ts}
Peak RetTime Type # [min]	Width [min]	Area [mAU*s]	Height [mAU]	Area %	(-)- 1b
 1 16.678 VB 2 20.478 VB	0.3780	5.37114 1792.24390	2.19076e-1 67.75414	 0.2988 99.7012	Kinetic resolution from PhB(OH) ₂

仪器 1 5/30/19 11:13:08

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002783.D Sample Name: YZ-4-85SM(RAC)

Acq. Operator	:						
Acq. Instrument	:	仪器 1 Location: Vial 1					
Injection Date	:	3/28/19 15:10:36					
		Inj Volume : 5.000 μl					
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M					
Last changed	:	3/28/19 15:07:42					
		(modified after loading)					
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M					
Last changed	:	5/30/19 11:10:50					
		(modified after loading)					
Sample Info	:	AD-H, n-hexane/i-PrOH =80/20, 0.8 mL/min, 30 oC, 254 nm					

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHOU2019\SIG1002783.D)



Area Percent Report

Sorted By : Signal Multiplier: : 1.0000 Dilution: : 1.0000			
Multiplier: : 1.0000 Dilution: : 1.0000	Sorted By	:	Signal
Dilution: : 1.0000	Multiplier:		: 1.0000
and the second sec	Dilution:		: 1.0000
Use Multiplier & Dilution Factor with ISTDs	Use Multiplier	& Dilution	Factor with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

仪器 1 5/30/19 11:11:04

Peak	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	*
1	16.750	VB	0.3398	1703.34778	77.67725	50.0414
2	20.564	BB	0.4090	1700.52625	64.20832	49.9586

 H_3C Ts rac-1b

Page 1 of 2



Sample Name: YZ-4-85BSM

Acq. Instrument	:	仪器 1 Location: Vial 1
Injection Date	:	3/28/19 16:41:59
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	3/28/19 16:36:17
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 11:16:11
		(modified after loading)





Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002786.D



Area Percent Report

Sorted By Multiplier: Dilution: Use Multiplier Signal 1: VWD1	: « Dilution A, Waveleng	Signal : Factor with th=254 nm	1.0000 1.0000 n ISTDs		H ₃ C. N. Ts
Peak RetTime Ty # [min] 1 16.700 BE 2 20.486 VE	pe Width [min] 3 0.3320 3 0.4141	Area [mAU*s] 1.72371 4197.48438	Height [mAU] 8.10932e-2 157.91101	Area % 0.0410 99.9590	(-)- 1b Kinetic resolution from 4-ClC ₆ H ₄ B(OH) ₂

仪器 1 5/30/19 11:16:15

Data File C:\CHEM32\1\DATA\ZHOU-19\YZNO12208.D Sample Name: YZ-4-85ATM(RAC)

	==	
Acq. Operator	:	
Acq. Instrument	:	Instrument l Location : -
Injection Date	:	3/29/2019 6:14:54 PM
Acq. Method	:	C:\CHEM32\1\METHOD\$\DEF_LC11.M
Last changed	:	3/29/2019 5:45:37 PM
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/2019 3:14:00 PM
		(modified after loading)
Sample Info	:	IB, Hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 nm



Area Percent Report · Sorted By : Signal : 1.0000 : 1.0000 Multiplier: Dilution: Ha Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area # [min] [min] mAU *s [mAU] * rac-21 1 6.191 VB 0.1162 1327.92615 177.90462 50.1746 2 7.061 BV 0.1300 1318.68494 156.98839 49.8254 Totals : 2646.61108 334.89301 -----*** End of Report ***

Data File C:\CHEM32\1\DATA\ZHOU-19\YZN012210.D Sample Name: Y2-4-85ATM

Acq. Operator :				
Acq. Instrument :	Instrument 1	Location	: -	
Injection Date :	3/29/2019 7:21:42 PM			
Acq. Method :	C:\CHEM32\1\METHODS\DEF LC11.M			
Last changed :	3/29/2019 7:17:30 PM			
	(modified after loading)			
Analysis Method :	C:\CHEM32\1\METHODS\DEF LC11.M			
Last changed :	5/30/2019 3:16:29 PM			
	(modified after loading)			
Sample Info :	IB, Hexane/i-PrOH = 80/20, 0.8	mL/min, 3	Ю oC,	254 nm







Instrument 1 5/30/2019 3:14:14 PM

Page 1 of 1

Instrument 1 5/30/2019 3:16:35 PM

Data File C:\CHEM32\1\DATA\ZH0U-19\YZN012209.D Sample Name: YZ-4-85BTM(RAC)

	==	
Acq. Operator	:	
Acq. Instrument	:	Instrument l Location : -
Injection Date	:	3/29/2019 6:50:04 PM
Acq. Method	:	C:\CHEM32\1\METHOD\$\DEF_LC11.M
Last changed	:	3/29/2019 6:31:53 PM
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/2019 3:18:19 PM
		(modified after loading)
Sample Info	:	IB, Hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 nm



Sorted By : Signal : 1.0000 : 1.0000 Multiplier: Dilution: H_3 Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area # [min] [min] mAU *s [mAU] * $Ar = 4 - CIC_6H_4$ 1 6.393 VV 0.1201 4162.28418 533.65631 50.1410 2 7.534 VB 0.1449 4138.87744 451.23529 49.8590 rac-2m Totals : 8301.16162 984.89160 -----

*** End of Report ***

Data File C:\CHEM32\1\DATA\ZHOU-19\YZN012305.D Sample Name: YZ-4-85BTM

Acq. Operator	:		
Acq. Instrument	:	Instrument 1 Location : -	
Injection Date	:	4/4/2019 4:42:52 PM	
Acq. Method	:	C:\CHEM32\1\METHODS\DEF LC11.M	
Last changed	:	4/4/2019 4:38:27 PM	
		(modified after loading)	
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M	
Last changed	:	5/30/2019 3:21:28 PM	
		(modified after loading)	
Sample Info	:	IB, Hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 2	254nm



Area Percent Report · Sorted By : Signal : 1.0000 : 1.0000 Multiplier: Dilution: H₂(Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area
 # [min]
 [min]
 mAU
 *s
 [mAU]
 *

 1
 6.388 WV
 0.1195
 829.39233
 107.05394
 98.9261
 $Ar = 4 - CIC_6H_4$ 2 7.546 VB 0.2344 9.00354 5.42591e-1 1.0739 (+)-2m Totals : 838.39587 107.59653 -----



Instrument 1 5/30/2019 3:18:24 PM

Page 1 of 1

Instrument 1 5/30/2019 3:21:32 PM

Data File C:\CHEM32\1\DATA\ZHOU-19\YZN013592.D Sample Name: YZ-6-8SM(rac)

	==	
Acq. Operator	:	
Acq. Instrument	:	Instrument 1 Location : -
Injection Date	:	10/4/2019 7:59:13 AM
Acq. Method	:	C:\CHEM32\1\METHODS\DEF LC11.M
Last changed	:	10/4/2019 7:29:59 AM
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M
Last changed	:	10/8/2019 10:15:37 PM
		(modified after loading)
Sample Info	;	IA, Hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 nm



----|

rac-1c

Last changed : 10/4/2019 8:25:01 AM (modified after loading) Analysis Method : C:\CHEM32\1\METHODS\DEF LC11.M Last changed : 10/8/2019 10:18:11 PM (modified after loading) Sample Info : IA, Hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 nm VWD1 A, Wavelen gth=254 nm (ZHOU-19\YZN013593.D) Nam. 250 200

Data File C:\CHEM32\1\DATA\ZHOU-19\YZN013593.D

Injection Date : 10/4/2019 8:29:25 AM Acq. Method : C:\CHEM32\1\METHODS\DEF_LC11.M

Acq. Instrument : Instrument 1

Sample Name: YZ-6-8SM

Acq. Operator :



Location : -

Sorted By Signal . : 1.0000 : 1.0000 Multiplier: Dilution: Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area # [min] [min] mAU *s [mAU] ÷ - | -----| 1 10.843 VB 0.2182 2133.72656 150.12553 56.5968

(-)- 1c
Kinetic resolution
from 4-CIC ₆ H ₄ B(OH) ₂

OMe

Ts

..... *** End of Report ***

3770.04858 223.51822

2 16.373 BB 0.3431 1636.32202 73.39268 43.4032

Instrument 1 10/8/2019 10:15:39 PM

Totals :

1 10.838 VB 0.2183 1721.50732 121.04810 50.0515

2 16.360 BB 0.3434 1717.96399 76.97136 49.9485

3439.47131 198.01946

*** End of Report ***

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Page 1 of 1

Instrument 1 10/8/2019 10:18:14 PM

Totals :

Data File C:\CHEM32\1\DATA\ZH0U-19\YZN013594.D Sample Name: YZ-6-8TM(RAC)

Acq. Operator				
Acq. Instrument	: Instrument 1	Location	:	-
Injection Date	: 10/7/2019 12:17:09 AM			
Acq. Method	C:\CHEM32\1\METHODS\DEF LC11.M			
Last changed	: 10/6/2019 11:15:04 PM			
	(modified after loading)			
Analysis Method	C:\CHEM32\1\METHODS\DEF LC11.M			
Last changed	: 10/8/2019 9:25:29 PM			
	(modified after loading)			
Sample Info	: IA, Hexane/i-PrOH = 90/10, 1.0 :	mL/min, 30) oC,	254 nm



*** End of Report ***



Analysis Method		C:\CHEM32\1\METHODS\DEF LC11.M
Last changed	:	10/8/2019 11:09:42 PM
		(modified after loading)
Sample Info	:	IA, Hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 nm



				/ / / / //
Multiplier:	:	1.0000		
Dilution:	:	1.0000		
Use Multiplier & 3	Dilution Factor wit	h ISTDs		
				Ts I Ts
Signal 1: VWD1 A,	Wavelength=254 nm			Ľ ∠JY ÖMe
				\mathbf{Y}
Peak RetTime Type	Width Area	Height	Area	
# [min]	[min] mAU *s	[mAU]	*	$\boldsymbol{\nu}$
				Ar = 4-CIC ₆ H₄
1 9.925 VB	0.2147 1448.35022	104.10171	99.7152	() •
2 10.946 BB	0.3016 4.13618	1.77072e-1	0.2848	(+)-2n
Totals :	1452.48640	104.27878		

*** End of Report ***

Instrument 1 10/8/2019 9:25:32 PM

Page 1 of 1

Instrument 1 10/8/2019 11:09:49 PM

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1003079.D Sample Name: YZ-4-58BSM(RAC)

Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location: Vial 1
Injection Date	:	5/30/19 9:38:00
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 9:07:37
		(modified after loading)
Analysis Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 10:07:23
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n
		TO





Signal 1: VWD1 A, Wavelength=254 nm

Use Multiplier & Dilution Factor with ISTDs

Peak Re # [tTime Typ min]	e Width [min]	Area [mAU*s]	Height [mAU]	Area %
		-			
1 1	.6.115 BV	0.2366	1152.96143	76.14727	49.9519
2 1	8.177 VB	0.2685	1155.18225	67.21445	50.0481



rac-1a

Page 1 of 2

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002884.D Sample Name: YZ-4-89SM Acq. Operator : Acq. Instrument : 仪器 1 Location : Vial 1 Injection Date : 4/15/19 15:25:10 Inj Volume : 5.000 µl Acq. Method : C:\CHEM32\1\METHODS\DEF_LC11.M Last changed : 4/15/19 14:50:36 (modified after loading) Analysis Method : C:\CHEM32\1\METHODS\DEF LC11.M Last changed : 5/30/19 11:24:17 (modified after loading) Sample Info : AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n m

Additional Info : Peak(s) manually integrated VWD1 A, Wavelength=254nm (2H0 U20 19%161002884.D)



Area Percent Report

Sorted By . Signal ∠Ts Multiplier: : 1.0000 : 1.0000 Dilution: Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area (-)-1a # [min] [mAU*s] [mAU] * 1 16.209 BB 0.2479 4.46118 2.76951e-1 0.0620 2 18.321 BBA 0.2743 7188.56885 406.57086 99.9380

仪器 1 5/30/19 11:24:24

Data File C:\CHEM32\1\DATA\ZHOU-19\YZN012754.D Sample Name: YZ-4-90(RAC)

Acq. Operator	:						
Acq. Instrument	:	Instrument 1	Location	:	-		
Injection Date	:	5/30/2019 4:43:47 PM					
Acq. Method	:	C:\CHEM32\1\METHODS\DEF LC11.M					
Last changed	:	5/30/2019 4:22:40 PM					
		(modified after loading)					
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M					
Last changed	:	5/30/2019 6:06:15 PM					
		(modified after loading)					
Sample Info	:	AD-H, Hexane/i-PrOH = 60/40, 0.	7 mL/min,	30	oC,	254	nn



-----Area Percent Report · Sorted By : Signal : 1.0000 Multiplier: Ts Dilution: Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area # [min] [min] mAU *s [mAU] * rac-**4** 1 16.413 BB 0.3404 1280.18188 58.35740 49.9418 2 27.043 BB



*** End of Report ***

Data File C:\CHEM32\1\DATA\ZHOU-19\YZN012755.D Sample Name: YZ-4-90

Acq. Operator	:						
Acq. Instrument	:	Instrument 1	Location	:	-		
Injection Date	:	5/30/2019 5:16:44 PM					
Acq. Method	:	C:\CHEM32\1\METHODS\DEF LC11.M					
Last changed	:	5/30/2019 5:15:35 PM					
		(modified after loading)					
Analysis Method	:	C:\CHEM32\1\METHODS\DEF LC11.M					
Last changed	:	5/30/2019 6:11:10 PM					
		(modified after loading)					
Sample Info	:	AD-H, Hexane/i-PrOH = $60/40$, 0.	7 mL/min,	30	oC,	254	nm



Area Percent Report · Signal Sorted By : : 1.0000 : 1.0000 Multiplier: Ts Dilution: Use Multiplier & Dilution Factor with ISTDs Signal 1: VWD1 A, Wavelength=254 nm Peak RetTime Type Width Area Height Area
 # fmin
 fmin
 area
 neight
 keight

 # [min]
 fmin
 naW
 %
 fmAU
 %

 ---- ---- ---- ---- ---- ---- ----

 1
 15.918
 B
 0.2614
 3.77989
 1.80796e-1
 0.0887

 2
 26.383
 B
 0.5753
 4256.37598
 114.78807
 99.9113
 (-)-4 4260.15587 114.96887 Totals : -----



Instrument 1 5/30/2019 6:06:23 PM

Totals :

Page 1 of 1

Instrument 1 5/30/2019 6:11:28 PM

Data File D:\USERS\CHEMSTATION\1\DATA\ZHOU2019\YZ3000166.D Sample Name: YZ-4-99(RAC)

Acq. Operator	:	SYSTEM
Sample Operator	:	SYSTEM
Acq. Instrument	:	1260II Location: 1
Injection Date	:	5/10/2019 11:35:04 AM Inj : 1
		Inj Volume : No inj
Acq. Method	:	C:\Users\Public\Documents\ChemStation\1\Methods\def LC.M
Last changed	:	5/10/2019 9:35:07 AM by SYSTEM
		(modified after loading)
Analysis Method	:	C:\Users\Public\Documents\ChemStation\l\Methods\def LC.M
Last changed	:	5/30/2019 4:48:00 PM by SYSTEM
		(modified after loading)
Sample Info	:	IC, n-Hexane/i-PrOH = 90/10, 0.7 mL/min, 30 oC, 254 nm



Area Percent Report

Sorte	d By	:	Signal			
Multi	plier	:	1.0000			
Dilut:	ion	:	1.0000			C
Do no	t use Multipl	ier & Di	ilution Fact	or with IST	Ds	
Signa.	1 1: VWD1 A,	Wavelenq	gth=254 nm			
Peak l	RetTime Type	Width	Area	Height	Area	
#	[min]	[min]	[mAU*s]	[mAU]	*	
1	12.646 BB	0.1126	1885.05750	251.01039	49.9288	
2	15.382 BB	0.2521	1890.43579	116.64406	50.0712	
Total:	з:		3775.49329	367.65445		

1260II 5/30/2019 4:48:04 PM SYSTEM



rac**-5a**

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Data File D:\USERS\CHEMSTATION\1\DATA\ZHOU2019\YZ3000164.D Sample Name: YZ-4-99







Area Percent Report

ignal
.0000
.0000
on Factor with ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	*
1	12.629	VB R	0.1026	1896.74072	281.16125	99.9318
2	15.396	VB	0.2399	1.29462	8.03803e-2	0.0682

1260II 5/30/2019 4:51:37 PM SYSTEM

Page 1 of 2

(-)-5a

Data File C:\CHEM32\1\DATA\ZHOU2018\SIG1002068.D Sample Name: YZ-4-34A(RAC)

Acq. Operator	:				
Acq. Instrument	:	仪器 1 Location: Vial 91			
Injection Date	:	12/8/18 10:42:08			
		Inj Volume : 5.000 µl			
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	12/8/18 10:40:33			
		(modified after loading)			
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	5/30/19 9:41:08			
		(modified after loading)			
Sample Info	:	AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n			
		70			





Area Percent Report

Sorted By	:	Signal		
Multiplier:		:	1.0000	
Dilution:		:	1.0000	
Use Multiplier & D	ilution	Factor with	n ISTDs	
Signal 1: VWD1 A,	Wavelenç	th=254 nm		
Peak RetTime Type	Width	Area	Height	Area
# [min]	[min]	[mAU*s]	[mAU]	÷
1 6.803 VV	0.1468	616.60956	64.94445	49.8919
2 7.446 VB	0.1690	619.28156	57.78165	50.1081
仪器 1 5/30/19 9:41:19				

Ph Ts rac-2a

Page 1 of 2







Area Percent Report

					Ph
Sorted By	:	Signal			÷
Multiplier:		:	1.0000		\land \land
Dilution:		:	1.0000		$\sim 10^{-1} \text{ M}^{-1}$
use Multiplier ۵	Dilution	Factor wit	h ISTDs		Ťs
Signal 1: VWD1 #	, Waveleng	th=254 nm			
Peak RetTime Typ	e Width	Area	Height	Area	() 22
# [min]	[min]	[mAU*s]	[mAU]	*	(- <i>)</i> -za

1	6.796 VV	0.2092	15.77220	1.06260	0.3078
2	7.383 VV	0.1618	5108.03760	489.32965	99.6922

```
仪器 1 5/30/19 11:26:33
```

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002966.D Sample Name: YZ-4-97RAC

Acq. Operator	:				
Acq. Instrument	:	仪器 1 Location : Vial 1			
Injection Date	:	5/7/19 15:17:34			
		Inj Volume : 5.000 µl			
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	5/7/19 15:10:19			
		(modified after loading)			
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M			
Last changed	:	5/30/19 11:30:52			
		(modified after loading)			
Sample Info	:	AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n			
		10			

Additional Info : Peak(s) manually integrated \WWD1 A Wavelength=254nm (ZHO U2019\S161002966.D)



-----Area Percent Report

Sorted By	:	Signal							
Multiplier:		:	1.0000						
Dilution:		:	1.0000						
Use Multiplier & D	ilution	Factor with	n ISTDs						
Signal 1: VWD1 A,	Signal 1: VWD1 A, Wavelength=254 nm								
Feak Reclime Type	widdu	ALEa	Height	Area					
# [min]	[min]	[mAU*s]	[mAU]	÷					
1 13.409 VV	0.2764	3212.07471	179.84615	49.3653					
2 14.156 VV	0.2974	3294.67456	170.52757	50.6347					
仪器 1 5/30/19 11:30:5	9								

rac-6

Page 1 of 2

Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1002967.D Sample Name: YZ-4-97 -----Acq. Operator : Acq. Instrument : 仪器 1 Location : Vial 1 Injection Date : 5/7/19 15:42:58 Inj Volume : 5.000 µl Acq. Method : C:\CHEM32\1\METHODS\DEF_LC11.M Last changed : 5/7/19 15:36:07 (modified after loading) Analysis Method : C:\CHEM32\1\METHODS\DEF LC11.M Last changed : 5/30/19 11:32:57 (modified after loading) Sample Info : AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n т





-----Area Percent Report

Sorted By	:	Signal			
Multiplier:		:	1.0000		
Dilution:		:	1.0000		
Use Multiplier «	Dilution H	Factor with	n ISTDs		Ts
Sigmal 1, VUD1 A	HOTTO LODOT				
Signal 1: VWD1 A Peak RetTime Type	, wavelengt e Width	Area	Height	Area	Ĺ
Signal 1: VWD1 A Peak RetTime Typ # [min]	, wavelengt e Width [min]	Area [mAU*s]	Height [mAU]	Area %	(+)-6
Signal 1: VWDI A Peak RetTime Typ # [min]	, wavelengt e Width [min] - -	Area [mAU*s]	Height [mAU]	Area *	(+)- 6
Signal 1: VWD1 A Peak RetTime Typ # [min] 1 13.360 VV	, wavelengt e Width [min] - - 0.2753 2	Area [mAU*s] 2854.07349	Height [mAU] 160.65536	Area * 99.2197	(+)- 6
Signal 1: VWD1 A Peak RetTime Typ # [min] 1 13.360 VV 2 14.160 VB	, wavelengt e Width [min] - - 0.2753 2 0.3117	Area [mAU*s] 2854.07349 22.44633	Height [mAU] 160.65536 1.07450	Area * 99.2197 0.7803	(+)- 6

Data File C:\CHEM32\1\DATA\ZHOU2018\SIG1002068.D Sample Name: YZ-4-34A(RAC)

Acq. Operator	:								
Acq. Instrument	:	仪器 1 Location : Vial 91							
Injection Date	:	12/8/18 10:42:08							
		Inj Volume : 5.000 µl							
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M							
Last changed	:	12/8/18 10:40:33							
		(modified after loading)							
Analysis Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M							
Last changed	:	5/30/19 9:41:08							
		(modified after loading)							
Sample Info	:	AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n							
		T0							





Area Percent Report

Sorted By	:	Signal		
Multiplier:		:	1.0000	
Dilution:		:	1.0000	
Use Multiplier & D	ilution	Factor with	n ISTDs	
Signal 1: VWD1 A, T	Wavelenç Width	gth=254 nm	Height	àrea
# [min]	[min]	[mAIIte]	[màII]	2
~ [<u></u>]		[[]	
1 6.803 VV	0.1468	616.60956	64.94445	49.8919
2 7.446 VB	0.1690	619.28156	57.78165	50.1081
仪器 1 5/30/19 9:41:19				

Ph N^{-Ts} rac-2a

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Data File C:\CHEM32\1\DATA\ZH0U2019\SIG1003031.D Sample Name: YZ-5-4SM -----Acq. Operator : Acq. Instrument : 仪器 1 Location : Vial 1 Injection Date : 5/17/19 19:45:19 Inj Volume : 5.000 µl Acq. Method : C:\CHEM32\1\METHODS\DEF_LC11.M Last changed : 5/17/19 19:05:30 (modified after loading) Analysis Method : C:\CHEM32\1\METHODS\DEF LC11.M Last changed : 5/30/19 11:37:07 (modified after loading) Sample Info : AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n m





Area Percent Report

Sorted By	:	Signal	
Multiplier:		: :	1.0000
Dilution:		: :	1.0000
Use Multiplier	& Dilution	Factor with	ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak	RetTime	Type	Width	Area	Height	Area	
#	[min]		[min]	[mAU*s]	[mAU]	*	
1	6.813	VV	0.1375	3814.08960	421.62192	99.2674	
2	7.433	VV	0.1819	28,14677	2.31449	0.7326	

```
仪器 1 5/30/19 11:37:21
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Data File D:\USERS\CHEMSTATION\1\DATA\ZH0U2019\YZ3000216.D Sample Name: YZ-5-4RAC

Acq. Operator	:	SYSTEM
Sample Operator	:	SYSTEM
Acq. Instrument	:	1260II Location : 1
Injection Date	:	5/17/2019 4:37:26 PM Inj: 1
		Inj Volume : No inj
Acq. Method	:	C:\Users\Public\Documents\ChemStation\l\Methods\def_LC.M
Last changed	:	5/17/2019 4:32:37 PM by SYSTEM
		(modified after loading)
Analysis Method	:	C:\Users\Public\Documents\ChemStation\l\Methods\def_LC.M
Last changed	:	5/30/2019 4:53:40 PM by SYSTEM
		(modified after loading)
Sample Info	:	IA, n-Hexane/i-PrOH = 98/2, 1.0 mL/min, 30 oC, 254 nm





Area Percent Report

Sorted By	:	Signal		
Multiplier	:	1.0000		
Dilution	:	1.0000		
Do not use Multiplier	6	Dilution Factor	with	ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak F	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	*
1	5.051	BB	0.1053	593.43890	85.17809	50.1176
2	5.828	BV R	0.1341	590.65308	67.34789	49.8824

1260II 5/30/2019 4:55:02 PM SYSTEM

Ph rac-7

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Data File D:\USERS\CHEMSTATION\1\DATA\ZH0U2019\YZ3000217.D Sample Name: YZ-5-4

Acq. Operator	:	SYSTEM	
Sample Operator	:	SYSTEM	
Acq. Instrument	:	1260II Location: 1	
Injection Date	:	5/17/2019 5:00:29 PM Inj: 1	
		Inj Volume : No inj	
Acq. Method	:	C:\Users\Public\Documents\ChemStation\1\Methods\def_LC.M	i
Last changed	:	5/17/2019 4:32:37 PM by SYSTEM	
		(modified after loading)	
Analysis Method	:	C:\Users\Public\Documents\ChemStation\1\Methods\def_LC.M	í
Last changed	:	5/30/2019 5:01:27 PM by SYSTEM	
		(modified after loading)	
Sample Info	:	IA, n-Hexane/i-PrOH = 98/2, 1.0 mL/min, 30 oC, 254 nm	



Area Percent Report

Sorted By	:	Sign	nal			
Multiplier	:	1.00	000			
Dilution	:	1.00	000			
Do not use Multiplier	6	Dilution	Factor	with	ISTDs	

Signal 1: VWD1 A, Wavelength=254 nm

1260II 5/30/2019 5:01:38 PM SYSTEM

Peak	RetTime	Type	Width	Area	Height	Area
#	[min]		[min]	[mAU*s]	[mAU]	25
1	5.025	VV R	0.1057	1674.16895	238.95752	99.6326
2	5.729	VB E	0.1184	6.17409	8.33798e-1	0.3674



Data File C:\CHEM32\1\DATA\ZHOU2019\SIG1003079.D Sample Name: YZ-4-58BSM(RAC)

Acq. Operator	:						
Acq. Instrument	:	仪器 1 Location: Vial 1					
Injection Date	:	5/30/19 9:38:00					
		Inj Volume : 5.000 µl					
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M					
Last changed	:	5/30/19 9:07:37					
		(modified after loading)					
Analysis Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M					
Last changed	:	5/30/19 10:07:23					
		(modified after loading)					
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n					
		TO					





Area Percent Report

Sorted By	:	Signal		
Multiplier:		:	1.0000	
Dilution:		:	1.0000	
Use Multiplier & 3	Dilution	Factor with	n ISTDs	
Signal 1: VWD1 A,	Wavelenç	gth=254 nm		
Peak RetTime Type	Width	Area	Height	Area
# [min]	[min]	[mAU*s]	[mAU]	*
1 16.115 BV	0.2366	1152.96143	76.14727	49.9519
2 18.177 VB	0.2685	1155.18225	67.21445	50.0481
仪器 1 5/30/19 10:07:	42			



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		······
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEN32\1\METHODS\DEF_LC11.M
Last changed	:	4/10/19 13:21:10
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 11:18:21
		(modified after loading)
Sample Info	:	AD-3, n-hexane/i-PrOH = 80/20, 0.8 mL/min, 30 oC, 254 n
		n





Area Percent Report



Data File C:\CHEM32\1\DATA\ZHOU2018\SIG1002068.D Sample Name: YZ-4-34A(RAC)

Acq. Operator	:	
Acq. Instrument	:	仪器 1 Location: Vial 91
Injection Date	:	12/8/18 10:42:08
		Inj Volume : 5.000 µl
Acq. Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	12/8/18 10:40:33
		(modified after loading)
Analysis Method	:	C:\CHEM32\1\METHODS\DEF_LC11.M
Last changed	:	5/30/19 9:41:08
		(modified after loading)
Sample Info	:	AD-H, n-hexane/i-PrOH = 90/10, 1.0 mL/min, 30 oC, 254 n
		70

Additional Info : Peak(s) manually integrated \WWD1 A, Wavelength=254nm (ZHOU2018%)G1002068.D)



Area Percent Report

Sorte	1 Bv	:	Signal		
Multip	plier:		:	1.0000	
Dilut:	ion:		:	1.0000	
Use Mu	ultiplier «	Dilution	Factor with	ISTDs	
Signa	1 1: VWD1 A,	Waveleng	th=254 nm		
Peak H	RetTime Type	Midth	Area	Height.	Area
Peak I #	RetTime Type [min]	Width [min]	Area [mAU*s]	Height [mAU]	Area %
Peak H # -	RetTime Type [min] 	Width [min]	Area [mAU*s] 	Height [mAU]	Area %
Peak H # - 1	RetTime Type [min] 6.803 VV	Width [min] 0.1468	Area [mAU*s] 616.60956	Height [mAU] 64.94445	Area % 49.8919

Ph N-Ts rac-2a

Page 1 of 2







Area Percent Report

Sorted By	:	Signal	
Multiplier:		:	1.0000
Dilution:		:	1.0000
Use Multiplier	& Dilution	Factor wit	h ISTDs

Signal 1: VWD1 A, Wavelength=254 nm

Peak	RetTime	Type	Width	Area	Height	Area	
#	[min]		[min]	[mAU*s]	[mAU]	25	
1	6.807	VV	0.1430	4333.33105	472.46912	99.0518	
2	7.368	VV	0.1897	41.48112	3.23173	0.9482	

仪器 1 5/30/19 11:21:10

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