Supporting Information

Stereodivergent Access to Enantioenriched Epoxy Alcohols with Three Stereogenic Centers via Ruthenium-Catalyzed Transfer Hydrogenation

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I. General Information

Commercially available materials purchased from Meryer, Aladdin, and Adamas were used as received; unless otherwise noted, all reactions and manipulations involving air- and moisture-sensitive compounds were performed using standard Schlenk techniques. Catalyst **C** was synthesized using the same procedures as shown in the literature (*J. Am. Chem. Soc.* **2011**, *133*, 14960–14963). All solvents were purified and dried using normal procedures. Proton nuclear magnetic resonance (1 H NMR) spectra were recorded on a Bruker AVANCE III HD400 (400 MHz) spectrometer. Chemical shifts were recorded in parts per million (ppm, δ) relative to tetramethylsilane (δ = 0.00 ppm) or chloroform (δ = 7.26 ppm). 1 H NMR splitting

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patterns are designated as singlet (s), doublet (d), triplet (t), quartet (q), dd (doublet of doublets), m (multiplet), and etc. All first-order splitting patterns were assigned on the basis of the appearance of the multiplet. Splitting patterns that could not be easily interpreted are designated as multiplet (m) or broad (br). Carbon nuclear magnetic resonance (¹³C NMR) spectra were recorded on a Bruker AVANCE III HD400 (100 MHz) spectrometer. High resolution mass spectral analysis (HRMS) was performed on Thermo Fisher Scientific LTQ FT Ultra mass spectrometer. The determination of ee was performed via chiral HPLC analysis using Shimadzu LC-20AD HPLC workstation. X-ray crystallography analysis was performed on Agilent SuperNova X-ray diffractionmeter. Optical rotations were measured using a 1 mL cell with a 5 dm path length on an INESA SGW-1 polarimeter and are reported as follows: [α]^{rt}_D (cin g per 100 mL solvent). Analytical thin-layer chromatography (TLC) was carried out on WFH-203 F254 pre-coated silica gel plate (0.2 mm thickness). Visualization was performed using a UV lamp or 2,4-dinitrophenylhydrazine or potassium permanganate stain.

II. X-ray crystallographic analysis

X-ray crystallographic analysis of 3i

Method for single crystal cultivation: a solid sample (10–20 mg) was dissolved in CHCl₃ (100–200 μ L) in a vial at room temperature, and MeOH (300–600 μ L) was added into the above solution slowly while keeping the sample all dissolved. Then vial was sealed with a piece of parafilm and stayed quietly for several days to allow the slow evaporation of the solution until a single crystal was obtained.

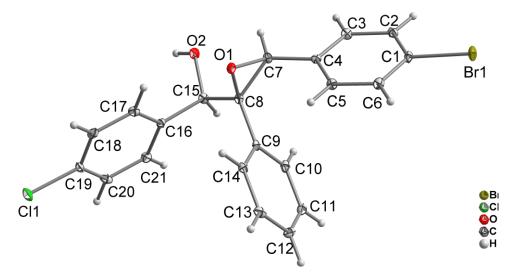


Table S1. Crystal data and structure refinement for data of compound 3i:

Identification code 3i

Empirical formula $C_{21}H_{16}BrClO_2$

Formula weight 415.70

Temperature/K 100.(2)

Crystal system monoclinic

Space group P2₁

a/Å 8.3009(2)

b/Å 10.9773(3)

c/Å 9.7497(3)

 α /° 90

 β /° 91.1200(10)

γ/° 90

Volume/ $Å^3$ 888.24(4)

Z 2

 $\rho_{calc}g/cm^3 1.554$

 μ/mm^{-1} 2.476

F(000) 420.0

Crystal size/mm³ $0.200 \times 0.200 \times 0.200$

Radiation ? ($\lambda = 0.71073$)

2Θ range for data collection/° 6.16 to 55.02

Index ranges $-10 \le h \le 10, -14 \le k \le 14, -12 \le l \le 12$

Reflections collected 18512

Independent reflections 3988 [$R_{int} = 0.0339$, $R_{sigma} = 0.0392$]

Data/restraints/parameters 3988/1/228

Goodness-of-fit on F² 1.052

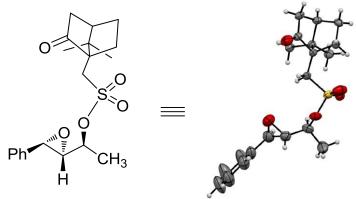
Final R indexes [I>= 2σ (I)] $R_1 = 0.0196$, $wR_2 = 0.0473$

Final R indexes [all data] $R_1 = 0.0201$, $wR_2 = 0.0474$

Largest diff. peak/hole / e Å⁻³ 0.48/-0.34

Flack parameter 0.046(3)

X-ray crystallographic analysis of 3m derivative (CCDC 2002835):



derivative of **3m** (95% ee after recrystallization)

Table S2. Crystal data and structure refinement for data of 3m derivative:

Identification code 3m derivative Empirical formula $C_{20}H_{26}NO_5S$ Formula weight 380.48 Temperature/K 230(2) Crystal system monoclinic Space group $P2_1$ a/Å 6.7030(5) h/Å 8.4049(6) c/Å 17.5344(14) $\alpha/^{\circ}$ 90 β/° 98.188(3) $\gamma/^{\circ}$ 90 Volume/Å³ 977.78(13) \mathbf{Z} 2 $\rho_{calc}g/cm^3$ 1.292

 $\rho_{calc}g/cm^{-1}$ 1.292 μ/mm^{-1} 0.193 F(000) 408.0

Crystal size/mm³ $0.1 \times 0.1 \times 0.1$ Radiation $MoK\alpha (\lambda = 0.71073)$

2Θ range for data collection/° 6.14 to 57.43

Index ranges $-9 \le h \le 9$, $-11 \le k \le 11$, $-23 \le l \le 23$

Reflections collected 27886

Independent reflections 5054 [$R_{int} = 0.0333$, $R_{sigma} = 0.0248$]

Data/restraints/parameters 5054/1/238Goodness-of-fit on F² 1.002

Final R indexes [I>= 2σ (I)] $R_1 = 0.0410$, $wR_2 = 0.1159$ Final R indexes [all data] $R_1 = 0.0466$, $wR_2 = 0.1218$

Largest diff. peak/hole / e Å⁻³ 0.47/-0.20 Flack parameter 0.05(2)

X-ray crystallographic analysis of 2n (CCDC 2002834):

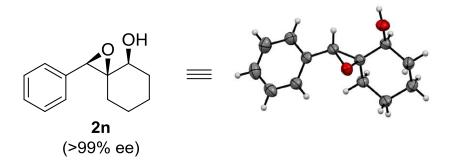


Table S3. Crystal data and structure refinement for data of 2n

Identification code 2n

Crystal system orthorhombic

 Space group
 P212121

 a/Å
 5.5579(2)

 b/Å
 7.7831(2)

 c/Å
 25.2947(7)

α/° 90 β/° 90 γ/° 90

Volume/ $Å^3$ 1094.19(6)

 $\begin{array}{ccc} Z & & 4 \\ & \rho_{calc} g/cm^3 & & 1.240 \\ & \mu/mm^{-1} & & 0.653 \\ F(000) & & 440.0 \end{array}$

Crystal size/mm³ $0.15 \times 0.1 \times 0.1$ Radiation $CuK\alpha (\lambda = 1.54184)$ 2Θ range for data collection/° 11.896 to 144.114

Index ranges $-2 \le h \le 6, -9 \le k \le 8, -30 \le l \le 31$

Reflections collected 3096

Independent reflections 1887 [$R_{int} = 0.0128$, $R_{sigma} = 0.0176$]

Data/restraints/parameters 1887/0/139

Goodness-of-fit on F^2 1.072

Final R indexes [I>=2 σ (I)] $R_1 = 0.0312$, $wR_2 = 0.0801$ Final R indexes [all data] $R_1 = 0.0322$, $wR_2 = 0.0810$

Largest diff. peak/hole / e Å⁻³ 0.18/-0.12 Flack parameter -0.04(11)

III. General procedures for the preparation of substrates

1. Typical procedure for the synthesis of substrates 1a–1h and 1l:

A solution of 1,2-diphenylethan-1-one **S1** (3.92 g, 20.0 mmol), 4-bromobenzaldehyde **S2** (3.68 g, 20.0 mmol), AcOH (1.0 mL) and piperidine (0.4 mL) in 50 mL toluene was refluxed overnight using a Dean-stark apparatus. After cooling to room temperature, the reaction was quenched by the addition of saturated aqueous NaHCO₃ solution (10 mL). The aqueous layer was extracted with ethyl acetate (3 x 100 mL) and the combined organic layer was dried over anhydrous Na₂SO₄, filtered and concentrated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 50:1) to afford product **S3** (2.61 g, 36% yield).

A solution of **S3** (0.51 g, 1.41 mmol) in 30 mL MeOH was added H_2O_2 (0.8 mL, 30% in water) dropwised at 0 °C. Then under this temperature, 0.8 mL aqueous NaOH solution (6 N) was slowly added into reaction mixture. The reaction suspension was monitored by TLC until no **S3** remained. Then the reaction was quenched with aqueous $Na_2S_2O_3$ solution, and then extracted with ethyl acetate (3 x 30 mL). The combined organic layer was dried over anhydrous Na_2SO_4 , filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 30:1) to afford product **1a** (0.45 g, 84% yield).

2. Typical procedure for the synthesis of substrate 1i:

A solution of methyl **S4** (0.40 g, 1.0 mmol) and 3-chloroperoxybenzoic acid (0.52 g, 3.0 mmol) in 10 mL CH_2Cl_2 were stirred for 2 h at room temperature. Then the reaction was quenched with aqueous $Na_2S_2O_3$ solution and extracted with ethyl acetate (3 x 20 mL). The combined organic layer was dried over anhydrous Na_2SO_4 , filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 50:1) to afford product **1i** (0.28 g, 68% yield).

3. Typical procedure for the synthesis of substrate 1k:

1,2-Diphenylethan-1-one **S5** (2.0 g, 10.0 mmol) and K_2CO_3 (1.48 g, 10.7 mmol) were added in 50 mL EtOH:acetone (v:v = 1:1). Then formaldehyde (1.5 mL, 18.0 mmol) was added at room temperature. After 4 h, the aqueous layer was extracted with ethyl acetate (3 x 50 mL) and the combined organic layer was dried over anhydrous Na_2SO_4 , filtered and concentrated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 50:1) to afford product **S6** (0.61 g, 29% yield).

A solution of **S6** (0.6 g, 2.9 mmol) in 30 mL MeOH was added H_2O_2 (3.0 mL, 30% in water) dropwised at 0 °C. Then under this temperature, 3.0 mL aqueous NaOH solution (6 N) was slowly added into reaction mixture. The reaction suspension was monitored by TLC until no **S6** remained. Then the reaction was quenched with aqueous $Na_2S_3O_2$ solution, and then extracted with ethyl acetate (3 x 30 mL). The combined organic layer was dried over anhydrous Na_2SO_4 , filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 30:1) to afford product **1k** (0.48 g, 74% yield).

4. Typical procedure for the synthesis of substrate 1m:

A solution of **S7** (1.88 g, 12.9 mmol) in 20 mL MeOH was added H_2O_2 (4.0 mL, 30% in water) dropwised at 0 °C. Then under this temperature, 3.8 mL aqueous NaOH solution (2 N) was slowly added into reaction mixture. The reaction suspension was slowly warmed to room temperature and monitored by TLC until no **S7** remained. Then the reaction was quenched with aqueous $Na_2S_3O_2$ solution, and then extracted with ethyl acetate (3 x 60 mL). The combined organic layer was dried over anhydrous Na_2SO_4 , filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 40:1) to afford product **1m** (1.42 g, 68% yield).

5. Typical procedure for the synthesis of substrate 1n:

A solution of cyclohexanone **S8** (7.36 g, 75.0 mmol) and benzaldehyde **S9** (5.3 g, 50.0 mmol) in 50 mL aqueous KOH solution (1 N) was heated to reflux for 3 h. After cooling to room temperature, the aqueous layer was extracted with ethyl acetate (3 \times 120 mL) and the combined organic layer was dried over anhydrous Na₂SO₄, filtered and concentrated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 30:1) to afford product **S10** (5.58 g, 60% yield).

A solution of **S10** (2.49 g, 13.4 mmol) in 35 mL MeOH was added H_2O_2 (5.0 mL, 30% in water) dropwised at room temperature. Then under this temperature, 2.5 mL aqueous NaOH solution (6 N) was slowly added into reaction mixture. The reaction suspension was monitored by TLC until no **S10** remained. Then the reaction was quenched with aqueous $Na_2S_3O_2$ solution, and then extracted with ethyl acetate (3 x 100 mL). The combined organic layer was dried over anhydrous Na_2SO_4 , filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 50:1) to afford product **1n** (1.30 g, 48% yield).

6. Typical procedure for the synthesis of substrate 10:

A solution of propiophenone **S11** (4.02 g, 30.0 mmol), benzaldehyde **S9** (6.36 g, 60.0 mmol), 60 mL aqueous KOH solution (0.1 N) and EtOH (60 mL) was stirred at room temperature for 3 d. After cooling to room temperature, the aqueous layer was extracted with ethyl acetate (3 x 150 mL) and the combined organic layer was dried over anhydrous Na_2SO_4 , filtered and concentrated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 30:1) to afford product **S12** (4.19 g, 63% yield).

A solution of **S12** (3.00 g, 13.5 mmol) in 60 mL MeOH was added H_2O_2 (6.0 mL, 30% in water) dropwised at room temperature. Then under this temperature, 3.0 mL aqueous NaOH solution (6 N) was slowly added into reaction mixture. The reaction suspension was monitored by TLC until no **S12** remained. Then the reaction was quenched with aqueous $Na_2S_3O_2$ solution, and then extracted with ethyl acetate (3 x 100 mL). The combined organic layer was dried over anhydrous Na_2SO_4 , filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 50:1) to afford product **10** (1.90 g, 59% yield).

IV. Typical procedure for the transfer hydrogenation:

A mixture of epoxy ketone **1a** (75.6 mg, 0.2 mmol) and the Ru catalyst (2.5 mg, 2 mol%) in 1.0 mL formic acid/triethylamine (v:v = 1:3) mixture was stirred at 40 $^{\circ}$ C under argon atmosphere. After completion of the reaction as indicated by TLC, it was extracted with ethyl acetate (3 x 10 mL) and the organic extract was dried over anhydrous Na₂SO₄ followed by the removal of solvent in a rotary evaporator. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 20:1) to afford product **2a** (35.1 mg, 46% yield) and **3a** (34.2 mg, 45% yield).

V. Procedures for derivatizations of products

1. Synthesis of compound 4a:

A solution of 3c (17.0 mg, 0.05 mmol) and Dess-Martin periodine (61.0 mg, 0.09 mmol) in 5.0 mL dry CH_2Cl_2 was stirred at room temperature under argon atmosphere. The reaction suspension was monitored by TLC until no 3c remained. Then the reaction mixture was extracted with ethyl acetate (3 x 10 mL). The combined organic layer was dried over anhydrous Na_2SO_4 , filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 5:1) to afford product 4a (15.3 mg, 96% yield).

2. Synthesis of compound 4b:

To a round-bottomed flask equiped with a dropping funnel, a condenser, and a magnetic stirrer was added 3c (10.0 mg, 0.03 mmol), NaN₃ (4.0 mg, 0.06 mmol), NH₄Cl (3.3 mg, 0.06 mmol) and 0.4 mL DMF. The reaction mixture was heated to 100 $\,^{\circ}$ C and stirred continuously for 8 h in an over-dried pressure-tight reaction tube. Then the mixture was extracted with CH₂Cl₂ (3 x 10 mL). The combined organic layer was dried over anhydrous Na₂SO₄, filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 10:1) to afford product 4b (10.7 mg, 98% yield).

3. Synthesis of compound 4c:

To a round-bottomed flask equipped with a dropping funnel, a condenser, and a magnetic stirrer was added 3c (17.0 mg, 0.05 mmol), NaH (4.0 mg, 0.13 mmol) and 5.0 mL THF at 0 $\,^{\circ}$ C under argon atmophere. After stirring for 40 min, BnBr (25.9 mg, 0.15 mmol) and n-Bu₄NI (9.2 mg, 0.005 mmol) were added into the reaction mixture at 0 $\,^{\circ}$ C. The the reaction resulting solution was slowly warm to room temperature and then stirred for additional 3 h. Then the reaction was quenched with aqueous saturated NH₄Cl solution, and then extracted with ethyl acetate (3 x 10 mL). The combined organic layer was dried over anhydrous Na₂SO₄, filtered and evaporated under reduced pressure. The residue was purified by chromatography (petroleum ether/ethyl acetate, v:v = 20:1) to afford product 4c (18.0 mg, 88% yield).

VI. Characterizations of new compounds

(3-(4-bromophenyl)-2-phenyloxiran-2-yl)(phenyl)methanone (1a): White solid, mp 149–150 °C, 0.45 g, 84% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.14–8.11 (m, 2H), 7.57–7.53 (m, 1H), 7.46–7.43 (m, 4H), 7.32–7.28 (m, 2H), 7.23–7.16 (m, 3H), 7.08–7.05 (m, 2H), 4.47 (s, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 195.0, 133.99, 133.95, 132.5, 131.7, 131.2, 130.0, 128.7, 128.6, 128.5, 128.4, 127.5, 122.3, 71.3, 62.6; HRMS (ESI, m/z): calcd. for C₂₁H₁₅O₂BrH⁺ 379.0328, found 379.0322; IR (KBr thin film, cm⁻¹): v 3007, 2989, 1689, 1592, 1448, 1260, 1183, 1007, 1010, 780, 760, 641.

(3-(4-chlorophenyl)-2-phenyloxiran-2-yl)(phenyl)methanone (1b): White solid, mp 100–101 °C, 0.28 g, 60% yield. ¹H NMR (400 MHz, CDCl₃) δ 8.10–8.07 (m, 2H), 7.52–7.47 (m, 1H), 7.41–7.37 (m, 4H), 7.20–7.05 (m, 7H), 4.44 (s, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 194.9, 134.0, 133.9, 132.0, 131.7, 129.9, 128.6, 128.5, 128.3, 128.2, 128.1, 127.5, 71.3, 62.5;

HRMS (ESI, m/z): calcd. for $C_{21}H_{15}O_2ClH^+$ 335.0833, found 335.0837; IR (KBr thin film, cm⁻¹): v 3060, 1689, 1609, 1578, 1501, 1495, 1181, 1087, 1013, 817, 746, 698.

(3-(4-fluorophenyl)-2-phenyloxiran-2-yl)(phenyl)methanone (1c): White solid, mp 93–94 $^{\circ}$ C, 0.34 g, 76% yield. 1 H NMR (600 MHz, CDCl₃) δ 8.15–8.12 (m, 2H), 7.57–7.53 (m, 1H), 7.46–7.43 (m, 4H), 7.23–7.14 (m, 5H), 6.88–6.84 (m, 2H), 4.50 (s, 1H); 13 C NMR (150 MHz, CDCl₃) δ 195.2, 162.6 (d, J = 245.4 Hz), 134.1, 133.9, 131.8, 130.0, 129.2 (d, J = 2.5 Hz), 128.7, 128.54, 128.48, 128.3, 127.6, 115.1 (d, J = 21.7 Hz), 71.3, 62.6; HRMS (ESI, m/z): calcd. for C₂₁H₁₅O₂FH⁺ 319.1129, found 319.1131; IR (KBr thin film, cm⁻¹): v 3037, 1687, 1601, 1582, 1510, 1446, 1275, 1187, 1031, 838, 775, 698.

(3-(naphthalen-1-yl)-2-phenyloxiran-2-yl)(phenyl)methanone (1d): Yellow solid, mp $109-110^{\circ}\text{C}$, 0.25 g, 51% yield. ^{1}H NMR (600 MHz, CDCl₃) δ 8.38–8.36 (m, 1H), 8.20–8.16 (m, 2H), 7.75 (d, J=8.2 Hz, 1H), 7.65 (d, J=8.2 Hz, 1H), 7.61–7.54 (m, 2H), 7.50–7.41 (m, 6H), 7.34–7.30 (m, 1H), 7.05–6.98 (m, 3H), 5.05 (s, 1H); ^{13}C NMR (150 MHz, CDCl₃) δ 195.4, 134.1, 134.0, 133.0, 132.1, 130.9, 130.2, 129.3, 128.7, 128.5, 128.3, 128.2, 128.0, 126.7, 126.5, 126.0, 124.9, 124.6, 123.6, 70.7, 63.2; HRMS (ESI, m/z): calcd. for $\text{C}_{25}\text{H}_{18}\text{O}_2\text{H}^+$ 351.1380, found 351.1381; IR (KBr thin film, cm⁻¹): v 2983, 1681, 1599, 1517, 1457, 1392, 1275, 1266, 1181, 1019, 855, 698.

(4-chlorophenyl)(2,3-diphenyloxiran-2-yl)methanone (**1e**): White solid, mp 114–115 °C, 0.38 g, 80% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.10–8.07 (m, 2H), 7.42–7.40 (m, 4H), 7.21–7.15 (m, 8H), 4.51 (s, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 194.2, 140.4, 133.2, 132.5,

131.7, 131.4, 129.0, 128.5, 128.3, 128.2, 128.0, 127.5, 126.8, 71.2, 63.2; HRMS (ESI, m/z): calcd. for $C_{21}H_{15}ClO_2H^+$ 335.0833, found 335.0834; IR (KBr thin film, cm⁻¹): v 3063, 3001, 1683, 1583, 1450, 1280, 1178, 1019, 845, 766, 752, 701.

(3-chlorophenyl)(2,3-diphenyloxiran-2-yl)methanone (1f): Yellow solid, mp 68–69 $^{\circ}$ C, 0.33 g, 70% yield. 1 H NMR (600 MHz, CDCl₃) δ 8.14–8.11 (m, 2H), 7.59–7.55 (m, 1H), 7.48–7.45 (m, 3H), 7.34–7.32 (m, 1H), 7.22–7.17 (m, 5H), 7.14–7.10 (m, 2H), 4.52 (s, 1H); 13 C NMR (150 MHz, CDCl₃) δ 194.8, 134.22, 134.18, 134.1, 133.9, 132.9, 130.0, 129.5, 128.73, 128.65, 128.3, 128.1, 127.7, 126.7, 125.8, 70.7, 63.3; HRMS (ESI, m/z): calcd. for $C_{21}H_{15}O_2H^+$ 335.0833, found 335.0835; IR (KBr thin film, cm⁻¹): v 3074, 1680, 1598, 1478, 1283, 1178, 1084, 1022, 857, 769, 755, 695.

(3-(4-bromophenyl)-2-(4-chlorophenyl)oxiran-2-yl)(phenyl)methanone (1g): White solid, mp 91–92 °C, 0.30 g, 51% yield. 1 H NMR (600 MHz, CDCl₃) δ 8.11–8.08 (m, 2H),7.58–7.55 (m, 1H), 7.47–7.43 (m, 2H), 7.39–7.36 (m, 2H), 7.34–7.31 (m, 2H), 7.20–7.17 (m, 2H), 7.06–7.02 (m, 2H), 4.46 (s, 1H); 13 C NMR (150 MHz, CDCl₃) δ 194.7, 134.7, 134.1, 133.8, 132.1, 131.4, 130.3, 129.9, 128.9, 128.73, 128.71, 128.3, 122.5, 70.8, 62.7; HRMS (ESI, m/z): calcd. for $C_{21}H_{14}O_{2}ClBrH^{+}$ 412.9938, found 412.9942; IR (KBr thin film, cm $^{-1}$): v 2938, 1677, 1604, 1498, 1450, 1319, 1280, 1175, 1019, 772, 749, 695.

(3-(4-bromophenyl)-2-(p-tolyl)oxiran-2-yl)(phenyl)methanone (1h): Colourless liquid, 0.49 g, 89% yield. 1 H NMR (600 MHz, CDCl₃) δ 8.15–8.12 (m, 2H), 7.55–7.51 (m, 1H),

7.45–7.41 (m, 2H), 7.35–7.30 (m, 4H), 7.10–7.07 (m, 2H), 7.03–7.01 (m, 2H), 4.47 (s, 1H), 2.22 (s, 3H); 13 C NMR (150 MHz, CDCl₃) δ 195.1, 138.4, 134.1, 133.9, 132.7, 131.2, 130.0, 129.2, 128.64, 128.60, 128.5, 127.4, 122.2, 71.4, 62.6, 21.3; HRMS (ESI, m/z): calcd. for $C_{22}H_{17}O_2BrH^+$ 393.0485, found 393.0488; IR (KBr thin film, cm⁻¹): v 2989, 1686, 1601, 1541, 1493, 1444, 1277, 1186, 1076, 888, 772, 698.

(3-(4-bromophenyl)-2-phenyloxiran-2-yl)(4-chlorophenyl)methanone (1i): White solid, mp 154–155 °C, 0.28 g, 68% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.08–8.05 (m, 2H), 7.42–7.40 (m, 4H), 7.31–7.28 (m, 2H), 7.24–7.17 (m, 3H), 7.06–7.03 (m, 2H), 4.46 (s, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 189.9, 136.6, 128.3, 127.4, 127.3, 125.1, 124.8, 124.53, 124.47, 123.5, 118.4, 67.3, 58.7; HRMS (ESI, m/z): calcd. for C₂₁H₁₄O₂ClBrH⁺ 412.9938, found 412.9941; IR (KBr thin film, cm⁻¹): v 3011, 2989, 1691, 1592, 1487, 1277, 1175, 1101, 1007, 766, 749, 695.

(3-(4-bromophenyl)-2-(m-tolyl)oxiran-2-yl)(phenyl)methanone (1j): Yellow solid, mp 77–78 °C, 0.41 g, 74% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.13–8.10 (m, 2H), 7.55–7.52 (m, 1H), 7.45–7.42 (m, 2H), 7.32–7.30 (m, 4H), 7.08–7.06 (m, 2H), 7.02–7.00 (m, 2H), 4.44 (s, 1H), 2.22 (s, 3H); ¹³C NMR (150 MHz, CDCl₃) δ 195.0, 138.1, 134.0, 133.9, 132.6, 131.5, 131.1, 130.0, 129.4, 128.7, 128.5, 128.3, 128.0, 124.5, 122.2, 71.4, 62.7, 21.5; HRMS (ESI, m/z): calcd. for C₂₂H₁₇O₂BrH⁺ 393.0486, found 393.0488; IR (KBr thin film, cm⁻¹): v 2955, 2856, 1683, 1541, 1450, 1374, 1269, 1175, 1022, 894, 763, 761, 707.

phenyl(2-phenyloxiran-2-yl)methanone (**1k**): Yellow solid, mp 68–69 °C, 0.48 g, 74% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.04–8.02 (m, 2H), 7.54–7.50 (m, 1H), 7.47–7.43 (m, 2H), 7.42–7.29 (m, 5H), 3.39 (d, J = 5.3 Hz, 1H), 3.07 (d, J = 5.3 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 194.8, 135.7, 134.4, 133.9, 130.1, 128.9, 128.7, 128.6, 125.5, 63.3, 55.2; HRMS (ESI, m/z): calcd. for C₁₅H₁₂O₂H⁺ 225.0910, found 225.0909; IR (KBr thin film, cm⁻¹): v 2986, 1683, 1606, 1581, 1487, 1453, 1277, 1172, 1010, 854, 769, 704.

(3-methyl-2-phenyloxiran-2-yl)(phenyl)methanone (1l): Colourless liquid, 0.24 g, 71% yield. 1 H NMR (600 MHz, CDCl₃) δ 8.05–8.02 (m, 2H), 7.57–7.55 (m, 2H), 7.51–7.47 (m, 1H), 7.40–7.35 (m, 4H), 7.32–7.29 (m, 1H), 3.58 (q, J = 5.4 Hz, 1H), 1.14 (d, J = 5.4 Hz, 3H); 13 C NMR (150 MHz, CDCl₃) δ 195.7, 134.3, 133.7, 133.3, 130.0, 128.6, 128.5, 128.4, 126.9, 68.7, 59.3, 13.5; HRMS (ESI, m/z): calcd. for C₁₆H₁₄O₂H⁺ 239.1067, found 239.1068; IR (KBr thin film, cm⁻¹): v 2932, 1729, 1680, 1584, 1502, 1380, 1277, 1260, 1184, 1056, 851, 701.

1-(3-phenyloxiran-2-yl)ethan-1-one (**1m**): White solid, mp 38–39 °C, 1.42 g, 68% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.37–7.32 (m, 3H), 7.27–7.24 (m, 2H), 3.99 (d, J = 1.8 Hz, 1H), 3.48–3.47 (m, 1H), 2.17 (s, 3H); ¹³C NMR (150 MHz, CDCl₃) δ 204.3, 135.1, 129.1, 128.8, 125.8, 63.6, 57.9, 24.9; HRMS (ESI, m/z): calcd. for C₁₀H₁₀O₂H⁺ 163.0754, found 163.0753; IR (KBr thin film, cm⁻¹): v 2997, 1721, 1489, 1460, 1366, 1281, 1250, 1187, 1090, 886, 752, 698.

2-phenyl-1-oxaspiro[**2.5**]**octan-4-one** (**1n**): White solid, mp 110–111 °C, 1.30 g, 48% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.38–7.31 (m, 3H), 7.30–7.28 (m, 2H), 4.04 (s, 1H), 2.73–2.68 (m, 1H), 2.45–2.38 (m, 1H), 2.03–1.99 (m, 1H), 1.93 (td, J = 13.2, 4.2 Hz, 1H), 1.82–1.73 (m,

2H), 1.63–1.58 (m, 1H), 1.47–1.39 (m, 1H); 13 C NMR (150 MHz, CDCl₃) δ 206.7, 133.6, 128.4, 126.8, 126.7, 66.1, 64.2, 42.0, 27.5, 24.8, 23.5; HRMS (ESI, m/z): calcd. for $C_{13}H_{14}O_{2}H^{+}$ 203.1067, found 203.1066; IR (KBr thin film, cm⁻¹): v 2861, 1713, 1647, 1457, 1423, 1321, 1278, 1156, 1065, 886, 758, 704.

(2-methyl-3-phenyloxiran-2-yl)(phenyl)methanone (1o): Colourless liquid, 1.90 g, 59% yield. 1 H NMR (600 MHz, CDCl₃) δ 8.06–8.04 (m, 2H), 7.61–7.57 (m, 1H), 7.51–7.47 (m, 2H), 7.43–7.34 (m, 5H), 4.16 (s, 1H), 1.43 (s, 3H); 13 C NMR (150 MHz, CDCl₃) δ 198.3, 134.4, 133.9, 133.6, 129.4, 128.7, 128.5, 128.4, 126.7, 66.6, 61.7, 14.5; HRMS (ESI, m/z): calcd. for $C_{16}H_{14}O_{2}H^{+}$ 239.1067, found 239.1069; IR (KBr thin film, cm $^{-1}$): v 3003, 2929, 1687, 1599, 1585, 1500, 1451, 1281, 1184, 977, 755, 698.

(*R*)-((2*R*,3*S*)-3-(4-bromophenyl)-2-phenyloxiran-2-yl)(phenyl)methanol (2a): Colourless liquid, 35.1 mg, 46% yield. 1 H NMR (600 MHz, CDCl₃) δ 7.29–7.27 (m, 3H), 7.22–7.18 (m, 4H), 7.15–7.11 (m, 1H), 7.06–7.03 (m, 2H), 6.82–6.79 (m, 2H), 6.75–6.72 (m, 2H), 4.98 (s, 1H), 4.61 (s, 1H), 2.81 (d, *J* = 1.7 Hz, 1H); 13 C NMR (150 MHz, CDCl₃) δ 138.5, 133.9, 133.1, 130.9, 129.2, 128.5, 128.3, 128.0, 127.7, 127.6, 121.7, 76.3, 71.8, 60.0; HRMS (ESI, m/z): calcd. for $C_{21}H_{17}O_{2}BrNa^{+}$ 403.0304, found 403.0308; $[\alpha]_{D}^{25}$: 15.9 (c 0.7, CHCl₃); HPLC analysis: 90% *ee* (Chiralcel AD-H, 3:97 i PrOH/Hexane, 1 mL/min, 254 nm), R_{t} (major) = 18.4 min, R_{t} (minor) = 24.8 min; IR (KBr thin film, cm $^{-1}$): v 3062, 3031, 2852, 1595, 1489, 1454, 1391, 1193, 1068, 1010, 757, 699.

(*R*)-((2*S*,3*R*)-3-(4-bromophenyl)-2-phenyloxiran-2-yl)(phenyl)methanol (3a): White solid, mp 131–133 °C, 34.2 mg, 45% yield. 1 H NMR (600 MHz, CDCl₃) δ 7.24–7.17 (m, 7H),

7.08–7.00 (m, 3H), 6.87–6.81 (m, 4H), 5.04 (d, J = 6.8 Hz, 1H), 4.62 (s, 1H), 2.38 (d, J = 6.8 Hz, 1H); 13 C NMR (150 MHz, CDCl₃) δ 139.6, 134.1, 133.3, 130.9, 128.7, 128.4, 128.3, 128.2, 127.7, 127.6, 127.0, 121.6, 76.3, 71.7, 61.0; HRMS (ESI, m/z): calcd. for $C_{21}H_{17}O_2BrNa^+$ 403.0304, found 403.0303; $[\alpha]_D^{25}$: +42.6 (c 0.8, CHCl₃); HPLC analysis: 98% *ee* (Chiralcel AD-H, 3:97 i PrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 39.2 min, R_t (minor) = 22.1 min; IR (KBr thin film, cm⁻¹): v 3065, 2928, 2849, 1599, 1492, 1448, 1397, 1189, 1065, 1008, 765, 695.

(*R*)-((2*R*,3*S*)-3-(4-chlorophenyl)-2-phenyloxiran-2-yl)(phenyl)methanol (2b): Colourless liquid, 29.0 mg, 43% yield. 1 H NMR (400 MHz, CDCl₃) δ 7.31–7.27 (m, 3H), 7.23–7.19 (m, 2H), 7.16–7.11 (m, 1H), 7.08–7.03 (m, 4H), 6.90–6.86 (m, 2H), 6.77–6.74 (m, 2H), 4.99 (d, *J* = 0.9 Hz, 1H), 4.63 (s, 1H), 2.86 (d, *J* = 1.7 Hz, 1H); 13 C NMR (150 MHz, CDCl₃) δ 138.6, 133.5, 133.4, 133.2, 129.2, 128.7, 128.5, 128.3, 128.02, 127.96, 127.6, 127.0, 76.3, 71.8, 60.0; HRMS (ESI, m/z): calcd. for $C_{21}H_{17}O_2ClNa^+$ 359.0809, found 359.0811; $[\alpha]_D^{25}$: –7.6 (c 0.8, CHCl₃); HPLC analysis: 90% *ee* (Chiralcel AD-H, 3:97 i PrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 11.4 min, R_t (minor) = 14.7 min; IR (KBr thin film, cm⁻¹): v 3059, 2929, 2857, 1600, 1491, 1451, 1265, 1195, 1086, 1013, 766, 692.

(*R*)-((2*S*,3*R*)-3-(4-chlorophenyl)-2-phenyloxiran-2-yl)(phenyl)methanol (3*b*): White solid, mp 128–130 °C, 26.9 mg, 40% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.26–7.19 (m, 5H), 7.07–7.01 (m, 5H), 6.93–6.90 (m, 2H), 6.86–6.83 (m, 2H), 5.04 (s, 1H), 4.64 (s, 1H), 2.55 (s, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 139.7, 133.6, 133.41, 133.37, 128.7, 128.4, 128.2, 127.99, 127.96, 127.7, 127.6, 127.0, 76.4, 71.8, 60.9; HRMS (ESI, m/z): calcd. for C₂₁H₁₇O₂ClNa⁺ 359.0809, found 359.0810; [α]_D²⁵: +26.5 (c 0.7, CHCl₃); HPLC analysis: 91% *ee* (Chiralcel AD-H, 3:97 ^{*i*}PrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 21.7 min, R_t

(minor) = 12.8 min; IR (KBr thin film, cm⁻¹): v 3063, 2925, 2854, 1601, 1493, 1454, 1261, 1193, 1091, 1016, 763, 697.

(*R*)-((2*R*,3*S*)-3-(4-fluorophenyl)-2-phenyloxiran-2-yl)(phenyl)methanol (2c): Colourless liquid, 28.5 mg, 44% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.30–7.27 (m, 3H), 7.22–7.19 (m, 2H), 7.15–7.11 (m, 1H), 7.06–7.03 (m, 2H), 6.92–6.89 (m, 2H), 6.79–6.74 (m, 4H), 4.99 (s, 1H), 4.65 (s, 1H), 2.97 (d, *J* = 9.5 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 162.3 (d, *J* = 245.6 Hz), 138.7, 133.3, 130.6, 129.3, 128.5, 128.3, 128.0, 127.7, 127.6, 114.8 (d, *J* = 21.5 Hz), 76.3 (t, *J* = 6.9 Hz), 71.7, 60.1; HRMS (ESI, m/z): calcd. for C₂₁H₁₇O₂FNa⁺ 343.1105, found 343.1107; [α]_D²⁵: –8.2 (c 0.5, CHCl₃); HPLC analysis: 89% *ee* (Chiralcel AD-H, 3:97 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 14.4 min, R_t (minor) = 17.8 min; IR (KBr thin film, cm⁻¹): v 3055, 2925, 2869, 1605, 1497, 1452, 1268, 1199, 1085, 1007, 768, 687.

(*R*)-((2*S*,3*R*)-3-(4-fluorophenyl)-2-phenyloxiran-2-yl)(phenyl)methanol (3*c*): White solid, mp 85–87 °C, 28.8 mg, 45% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.28–7.20 (m, 5H), 7.07–7.00 (m, 3H), 6.97–6.93 (m, 2H), 6.86–6.84 (m, 2H), 6.79–6.75 (m, 2H), 5.05 (d, *J* = 6.3 Hz, 1H), 4.65 (s, 1H), 2.43 (d, *J* = 6.7 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 162.4 (d, *J* = 244.5 Hz), 139.9, 130.8 (d, *J* = 3.0 Hz), 129.4, 128.9, 128.5, 128.4, 128.3 (d, *J* = 2.4 Hz), 127.74, 127.66, 127.1, 114.9 (d, *J* = 21.6 Hz), 76.6, 71.7, 61.1; HRMS (ESI, m/z): calcd. for C₂₁H₁₇O₂FNa⁺ 343.1105, found 343.1107; [α]_D²⁵: +27.7 (c 0.3, CHCl₃); HPLC analysis: 91% *ee* (Chiralcel AD-H, 10:90 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 10.1 min, R_t (minor) = 7.2 min; IR (KBr thin film, cm⁻¹): v 3057, 2929, 2861, 1602, 1499, 1456, 1269, 1194, 1088, 1011, 771, 684.

(*R*)-((2*R*,3*S*)-3-(naphthalen-1-yl)-2-phenyloxiran-2-yl)(phenyl)methanol (2d): Yellow liquid, 38.9 mg, 55% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.00 (d, J = 8.5 Hz, 1H), 7.79–7.76 (m, 1H), 7.58 (d, J = 7.8 Hz, 1H), 7.52–7.44 (m, 2H), 7.38–7.32 (m, 5H), 7.14 (t, J = 7.6 Hz, 1H), 7.10–7.07 (m, 1H), 7.03–6.99 (m, 1H), 6.92 (t, J = 7.8 Hz, 2H), 6.81–6.79 (m, 2H), 5.20 (d, J = 2.0 Hz, 1H), 5.06 (s, 1H), 3.07 (d, J = 2.0 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 138.8, 133.8, 133.0, 130.9, 130.7, 128.8, 128.7, 128.5, 128.3, 127.8, 127.4, 126.2, 125.7, 125.2, 125.1, 124.0, 123.9, 122.8, 76.4, 71.9, 59.4; HRMS (ESI, m/z): calcd. for $C_{25}H_{20}O_2Na^+$ 375.1356, found 375.1354; $[\alpha]_D^{25}$: 79.1 (c 1.9, CHCl₃); HPLC analysis: 74% *ee* (Chiralcel AD-H, 10:90 [†]PrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 8.5 min, R_t (minor) = 21.0 min; IR (KBr thin film, cm⁻¹): v 2986, 1571, 1457, 1332, 1278, 1258, 1224, 1071, 937, 852, 775, 667.

(*R*)-((2*S*,3*R*)-3-(naphthalen-1-yl)-2-phenyloxiran-2-yl)(phenyl)methanol (3d): White solid, mp 86–87 °C, 25.9 mg, 37% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.15–8.13 (m, 1H), 7.78–7.76 (m, 1H), 7.58–7.54 (m, 2H), 7.49–7.45 (m, 1H), 7.34–7.25 (m, 5H), 7.17–7.13 (m, 2H), 6.92–6.84 (m, 5H), 5.31 (d, J = 6.8 Hz, 1H), 5.20 (s, 1H), 2.55 (d, J = 6.8 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 139.9, 134.1, 133.0, 130.9, 130.8, 128.8, 128.4, 128.3, 127.9, 127.6, 127.33, 127.26, 126.2, 125.7, 125.1, 123.9, 122.9, 76.2, 71.4, 60.1; HRMS (ESI, m/z): calcd. for C₂₅H₂₀O₂Na⁺ 375.1356, found 375.1357; [α]_D²⁵: +158.1 (c 0.6, CHCl₃); HPLC analysis: 65% *ee* (Chiralcel AD-H, 10:90 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 15.8 min, R_t (minor) = 10.0 min; IR (KBr thin film, cm⁻¹): v 3040, 1539, 1525, 1508, 1460, 1338, 1275, 1090, 900, 838, 775, 707.

(*R*)-(4-chlorophenyl)((2*R*,3*S*)-2,3-diphenyloxiran-2-yl)methanol (2e): Colourless liquid, 28.3 mg, 42% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.26–7.24 (m, 2H), 7.15–7.03 (m, 8H), 6.91 (dd, J = 7.5, 1.5 Hz, 2H), 6.76–6.74 (m, 2H), 4.97 (d, J = 1.4 Hz, 1H), 4.62 (s, 1H), 2.91 (d, J = 1.7 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 137.3, 134.5, 134.1, 133.1, 129.3, 128.9, 128.5, 128.0, 127.8, 127.6, 126.7, 75.8, 71.5, 60.4; HRMS (ESI, m/z): calcd. for C₂₁H₁₇O₂ClNa⁺ 359.0809, found 359.0812; $[\alpha]_D^{25}$: –8.8 (c 0.5, CHCl₃); HPLC analysis: 83% *ee* (Chiralcel AD-H, 3:97 ⁱPrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 19.1 min, R_t (minor) = 25.3 min; IR (KBr thin film, cm⁻¹): v 3061, 2931, 2861, 1599, 1492, 1458, 1264, 1195, 1092, 1013, 765, 682.

(*R*)-(4-chlorophenyl)((2*S*,3*R*)-2,3-diphenyloxiran-2-yl)methanol (3e): White semi-solid, 25.7 mg, 38% yield. 1 H NMR (600 MHz, CDCl₃) δ 7.23–7.20 (m, 2H), 7.16–7.13 (m, 2H), 7.09–7.01 (m, 6H), 6.98–6.96 (m, 2H), 6.86–6.83 (m, 2H), 5.03 (d, J = 6.8 Hz, 1H), 4.64 (s, 1H), 2.50 (d, J = 6.8 Hz, 1H); 13 C NMR (150 MHz, CDCl₃) δ 138.3, 134.6, 133.9, 133.3, 128.7, 128.5, 128.4, 127.8, 127.70, 127.66, 127.6, 126.6, 75.9, 71.4, 61.7; HRMS (ESI, m/z): calcd. for $C_{21}H_{17}O_2ClNa^+$ 359.0809, found 359.0810; $[\alpha]_D^{25}$: +26.2 (c 0.5, CHCl₃); HPLC analysis: 98% *ee* (Chiralcel AD-H, 3:97 i PrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 37.5 min, R_t (minor) = 21.5 min; IR (KBr thin film, cm $^{-1}$): v 3057, 2932, 2866, 1597, 1493, 1459, 1261, 1199, 1087, 1015, 759, 681.

(*R*)-(3-chlorophenyl)((2*R*,3*S*)-2,3-diphenyloxiran-2-yl)methanol (2*f*): Colourless liquid, 29.1 mg, 43% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.32–7.30 (m, 3H), 7.26–7.21 (m, 2H), 7.13–7.07 (m, 4H), 6.97–6.88 (m, 4H), 6.49 (d, J = 7.6 Hz, 1H), 4.96 (s, 1H), 4.68 (s, 1H), 2.87 (d, J = 6.2 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 138.3, 135.7, 134.2, 133.4, 129.2, 128.62, 128.59, 128.4, 128.0, 127.9, 127.8, 127.6, 127.4, 126.5, 76.2, 71.0, 60.6; HRMS (ESI, m/z): calcd. for C₂₁H₁₇O₂ClNa⁺ 359.0809, found 359.0811; $[\alpha]_D^{25}$: –19.4 (c 2.1, CHCl₃); HPLC analysis: 80% *ee* (Chiralcel AD-H, 1.5:98.5 ⁱPrOH/Hexane, 1 mL/min, 220

nm), R_t (major) = 25.3 min, R_t (minor) = 34.7 min; IR (KBr thin film, cm⁻¹): v 3058, 3027, 2869, 1598, 1501, 1451, 1267, 1199, 1095, 1005, 765, 689.

(*R*)-(3-chlorophenyl)((2*S*,3*R*)-2,3-diphenyloxiran-2-yl)methanol (3*f*): White semi-solid, 25.6 mg, 38% yield. 1 H NMR (400 MHz, CDCl₃) δ 7.29–7.22 (m, 5H), 7.13–7.10 (m, 3H), 7.04–6.98 (m, 3H), 6.94–6.90 (m, 2H), 6.88 (d, J = 7.6 Hz, 1H), 5.04 (d, J = 6.5 Hz, 1H), 4.68 (s, 1H), 2.51 (d, J = 6.6 Hz, 1H); 13 C NMR (100 MHz, CDCl₃) δ 139.3, 135.7, 134.3, 133.3, 128.9, 128.6, 128.4, 128.3, 127.84, 127.75, 127.7, 126.9, 126.8, 126.4, 76.4, 71.1, 61.7; HRMS (ESI, m/z): calcd. for $C_{21}H_{17}O_{2}CINa^{+}$ 359.0809, found 359.0808; $[\alpha]_{D}^{25}$: +19.0 (c 0.5, CHCl₃); HPLC analysis: 90% *ee* (Chiralcel AD-H, 1.5:98.5 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_{t} (major) = 51.0 min, R_{t} (minor) = 38.5 min; IR (KBr thin film, cm⁻¹): v 3059, 2931, 2868, 1598, 1500, 1455, 1268, 1195, 1097, 1007, 766, 692.

(*R*)-((2*R*,3*S*)-3-(4-bromophenyl)-2-(4-chlorophenyl)oxiran-2-yl)(phenyl)methanol (2g): Colourless liquid, 37.5 mg, 45% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.32–7.29 (m, 3H), 7.26–7.22 (m, 2H), 7.21–7.18 (m, 2H), 7.03 (d, J = 8.6 Hz, 2H), 6.83–6.80 (m, 2H), 6.69–6.66 (m, 2H), 4.94 (s, 1H), 4.63 (s, 1H), 2.74 (s, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 138.1, 134.1, 134.0, 133.3, 131.1, 128.70, 128.65, 128.6, 128.4, 128.0, 127.9, 121.9, 75.8, 71.6, 61.1; HRMS (ESI, m/z): calcd. for C₂₁H₁₆O₂ClBrNa⁺ 436.9914, found 436.9916; $[\alpha]_D^{25}$: –6.1 (c 0.5, CHCl₃); HPLC analysis: 87% *ee* (Chiralcel AD-H, 4:96 ⁱPrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 14.4 min, R_t (minor) = 23.0 min; IR (KBr thin film, cm⁻¹): v 3063, 2935, 2866, 1601, 1493, 1449, 1270, 1197, 1089, 1009, 759, 681.

(*R*)-((2*S*,3*R*)-3-(4-bromophenyl)-2-(4-chlorophenyl)oxiran-2-yl)(phenyl)methanol (3g): White solid, 123–126 °C, 24.9 mg, 30% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.29–7.19 (m, 7H), 7.10–7.01 (m, 3H), 6.88–6.84 (m, 3H), 5.05 (d, *J* = 6.6 Hz, 1H), 4.63 (s, 1H), 2.40 (d, *J* = 6.8 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 139.6, 134.1, 133.3, 130.9, 128.6, 128.3, 128.22, 128.16, 127.64, 127.56, 127.0, 121.6, 76.3, 71.6, 60.9; HRMS (ESI, m/z): calcd. for C₂₁H₁₆O₂ClBrNa⁺ 436.9914, found 436.9915; $[\alpha]_D^{25}$: +28.3 (c 0.6, CHCl₃); HPLC analysis: 93% *ee* (Chiralcel AD-H, 4:96 ⁱPrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 27.9 min, R_t (minor) = 17.6 min; IR (KBr thin film, cm⁻¹): v 3066, 2929, 2863, 1603, 1495, 1451, 1268, 1199, 1093, 1015, 761, 683.

(*R*)-((2*R*,3*S*)-3-(4-bromophenyl)-2-(*p*-tolyl)oxiran-2-yl)(phenyl)methanol (2h): Colourless liquid, 32.5 mg, 41% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.30–7.28 (m, 3H), 7.23–7.19 (m, 4H), 6.88–6.81 (m, 4H), 6.63 (d, J = 8.0 Hz, 2H), 4.97 (d, J = 1.1 Hz, 1H), 4.58 (s, 1H), 2.80 (d, J = 1.7 Hz, 1H), 2.23 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 138.6, 137.7, 134.0, 131.0, 130.9, 130.0, 129.0 128.37, 128.35, 128.2, 127.6, 121.6, 76.3, 71.6, 59.9, 21.2; HRMS (ESI, m/z): calcd. for C₂₂H₁₉O₂BrNa⁺ 417.0461, found 417.0463; [α]_D²⁵: –14.3 (c 0.5, CHCl₃); HPLC analysis: 88% *ee* (Chiralcel AD-H, 3:97 ⁱPrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 19.1 min, R_t (minor) = 23.8 min; IR (KBr thin film, cm⁻¹): v 3061, 2936, 2865, 1599, 1497, 1459, 1264, 1201, 1097, 1011, 761, 688.

(*R*)-((2*S*,3*R*)-3-(4-bromophenyl)-2-(*p*-tolyl)oxiran-2-yl)(phenyl)methanol (3h): White solid, mp 114–116 °C, 31.7 mg, 40% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.26–7.20 (m, 7H), 6.88–6.82 (m, 4H), 6.73 (d, *J* = 8.0 Hz, 2H), 5.02 (d, *J* = 6.7 Hz, 1H), 4.59 (s, 1H), 2.36 (d, *J* = 6.8 Hz, 1H), 2.19 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 139.7, 137.3, 134.2, 130.8, 130.1, 128.4, 128.29, 128.28, 128.25, 128.1, 127.0, 121.5, 76.4, 71.5, 60.9, 21.2; HRMS (ESI, m/z): calcd. for C₂₂H₁₉O₂BrNa⁺ 417.0461, found 417.0463; [α]_D²⁵: +28.6 (c 0.9, CHCl₃); HPLC

analysis: 97% *ee* (Chiralcel AD-H, 3:97 i PrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 32.6 min, R_t (minor) = 20.8 min; IR (KBr thin film, cm⁻¹): v 3060, 2938, 2862, 1601, 1499, 1463, 1261, 1200, 1095, 1013, 763, 671.

(*R*)-((2*R*,3*S*)-3-(4-bromophenyl)-2-(*m*-tolyl)oxiran-2-yl)(phenyl)methanol (2i): Colourless liquid, 30.7 mg, 39% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.29–7.20 (m, 7H), 6.96–6.88 (m, 2H), 6.82 (d, J = 7.5 Hz, 2H), 6.66 (s, 1H), 6.42 (d, J = 7.2 Hz, 1H), 4.98 (s, 1H), 4.57 (s, 1H), 2.79 (s, 1H), 2.15 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 138.5, 137.2, 133.9, 132.9, 130.8, 129.7, 128.7, 128.4, 128.3, 128.1, 127.6, 127.3, 126.2, 121.6, 76.2, 71.7, 59.9, 21.2; HRMS (ESI, m/z): calcd. for C₂₂H₁₉O₂BrNa⁺ 417.0461, found 417.0464; [α]_D²⁵: –31.0 (c 0.5, CHCl₃); HPLC analysis: 92% *ee* (Chiralcel AD-H, 1:99 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 41.7 min, R_t (minor) = 49.0 min; IR (KBr thin film, cm⁻¹): v 3059, 2931, 2861, 1600, 1496, 1460, 1263, 1201, 1098, 1009, 799, 705.

(*R*)-((2*S*,3*R*)-3-(4-bromophenyl)-2-(*m*-tolyl)oxiran-2-yl)(phenyl)methanol (3i): White solid, mp 110–112 °C, 29.9 mg, 38% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.27–7.20 (m, 7H), 6.93–6.85 (m, 4H), 6.69 (s, 1H), 6.61 (d, *J* = 6.7 Hz, 1H), 5.03 (d, *J* = 6.3 Hz, 1H), 4.58 (s, 1H), 2.40 (d, *J* = 6.6 Hz, 1H), 2.13 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 139.6, 137.1, 134.1, 133.1, 130.8, 129.2, 128.4, 128.24, 128.17, 128.1, 127.3, 126.9, 125.6, 121.5, 76.4, 71.6, 60.9, 21.2; HRMS (ESI, m/z): calcd. for C₂₂H₁₉O₂BrNa⁺ 417.0460, found 417.0463; [α]_D²⁵: +35.9 (c 0.6, CHCl₃); HPLC analysis: 90% *ee* (Chiralcel AD-H, 3:97 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 26.9 min, R_t (minor) = 18.0 min; IR (KBr thin film, cm⁻¹): v 3057, 2927, 2866, 1602, 1498, 1466, 1260, 1205, 1099, 1007, 793, 700.

(*R*)-((2*R*,3*S*)-3-(4-bromophenyl)-2-phenyloxiran-2-yl)(4-chlorophenyl)methanol (2j): Colourless liquid, 46.0 mg, 55% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.27–7.23 (m, 2H), 7.22–7.19 (m, 2H), 7.17–7.05 (m, 5H), 6.80–6.72 (m, 4H), 4.96 (d, J = 1.6 Hz, 1H), 4.55 (s, 1H), 2.82 (d, J = 1.9 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 137.1, 134.2, 133.7, 132.7, 131.0, 129.2, 128.9, 128.5, 128.3, 128.2, 127.8, 121.9, 75.7, 71.6, 59.9; HRMS (ESI, m/z): calcd. for C₂₁H₁₆O₂ClBrNa⁺ 436.9914, found 436.9916; [α]_D²⁵: –10.5 (c 0.5, CHCl₃); HPLC analysis: 54% *ee* (Chiralcel AD-H, 3:97 ⁱPrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 22.5 min, R_t (minor) = 27.2 min; IR (KBr thin film, cm⁻¹): v 3064, 2928, 2859, 1602, 1488, 1451, 1268, 1198, 1087, 1011, 767, 687.

(*R*)-((2*S*,3*R*)-3-(4-bromophenyl)-2-phenyloxiran-2-yl)(4-chlorophenyl)methanol (3j): White solid, mp 122–124 °C, 30.0 mg, 36% yield. ¹H NMR (400 MHz, CDCl₃) δ 7.27–7.20 (m, 4H), 7.14 (d, J = 8.4 Hz, 2H), 7.10–7.03 (m, 3H), 6.88–6.84 (m, 4H), 5.04 (d, J = 6.8 Hz, 1H), 4.61 (s, 1H), 2.42 (d, J = 6.8 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 138.1, 133.9, 133.8, 133.1, 130.9, 128.50, 128.45, 128.4, 128.2, 127.8, 127.7, 121.7, 75.6, 71.4, 60.9; HRMS (ESI, m/z): calcd. for C₂₁H₁₆O₂ClBrNa⁺ 436.9914, found 436.9918; [α]_D²⁵: +36.4 (c 0.6, CHCl₃); HPLC analysis: 98% *ee* (Chiralcel AD-H, 3:97 ⁱPrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 41.8 min, R_t (minor) = 21.9 min; IR (KBr thin film, cm⁻¹): v 3061, 2926, 2857, 1601, 1489, 1455, 1263, 1197, 1086, 1009, 766, 689.

(*R*)-phenyl((*S*)-2-phenyloxiran-2-yl)methanol (2k): Colourless liquid, 19.9 mg, 44% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.28–7.21 (m, 6H), 7.19–7.16 (m, 2H), 7.14–7.11 (m, 2H),

5.04 (d, J = 0.9 Hz, 1H), 3.39 (dd, J = 5.1, 0.5 Hz, 1H), 2.94 (dd, J = 5.1, 0.5 Hz, 1H), 2.74 (d, 1.7 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 138.9, 136.8, 128.33, 128.27, 128.14, 128.08, 127.5, 74.5, 64.3, 50.3; HRMS (ESI, m/z): calcd. for $C_{15}H_{14}O_2Na^+$ 249.0886, found 249.0888; $[\alpha]_D^{25}$: +71.5 (c 0.6, CHCl₃); HPLC analysis: 85% *ee* (Chiralcel AD-H, 5:95 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 15.5 min, R_t (minor) = 11.0 min; IR (KBr thin film, cm⁻¹): v 2983, 1559, 1548, 1479, 1465, 1275, 1267, 1048, 1028, 906, 778, 673.

(*R*)-phenyl((*R*)-2-phenyloxiran-2-yl)methanol (3k): Yellow liquid, 20.8 mg, 46% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.28–7.22 (m, 5H), 7.22–7.15 (m, 5H), 5.06 (d, *J* = 7.1 Hz, 1H), 3.41 (d, *J* = 5.5 Hz, 1H), 2.80 (d, *J* = 5.5 Hz, 1H), 2.34 (d, *J* = 7.1 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 139.9, 137.0, 128.4, 128.1, 128.0, 127.8, 127.3, 127.2, 75.0, 63.5, 52.4; HRMS (ESI, m/z): calcd. for C₁₅H₁₄O₂Na⁺ 249.0087, found 249.0088; [α]_D²⁵: +18.0 (c 1.5, CHCl₃); HPLC analysis: 85% *ee* (Chiralcel AD-H, 1:99 ^{*i*}PrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 27.3 min, R_t (minor) = 37.0 min; IR (KBr thin film, cm⁻¹): v 3037, 1687, 1601, 1582, 1510, 1446, 1275, 1187, 1031, 838, 775, 698.

(*R*)-((2*R*,3*S*)-3-methyl-2-phenyloxiran-2-yl)(phenyl)methanol (2l): White solid, mp 84–85 $^{\circ}$ C, 17.3 mg, 36% yield. 1 H NMR (600 MHz, CDCl₃) δ 7.30–7.25 (m, 3H), 7.25–7.19 (m, 3H), 7.18–7.15 (m, 2H), 6.98–6.96 (m, 2H), 4.86 (s, 1H), 3.63 (q, *J* = 5.5 Hz, 1H), 2.71 (d, *J* = 1.1 Hz, 1H), 1.03 (d, *J* = 5.5 Hz, 3H); 13 C NMR (150 MHz, CDCl₃) δ 139.1, 135.1, 128.4, 128.3, 128.2, 127.91, 127.88, 127.8, 76.2, 68.7, 56.4, 15.0; HRMS (ESI, m/z): calcd. for $C_{16}H_{16}O_2Na^+$ 263.1043, found 263.1040; $[\alpha]_D^{25}$: +27.8 (c 0.3, CHCl₃); HPLC analysis: 80% *ee* (Chiralcel AD-H, 5:95 i PrOH/Hexane, 1 mL/min, 254 nm), R_t (major) = 7.4 min, R_t (minor) = 8.7 min; IR (KBr thin film, cm⁻¹): v 2991, 1539, 1454, 1414, 1315, 1283, 1264, 1133, 920, 897, 767, 676.

(*R*)-((2*S*,3*R*)-3-methyl-2-phenyloxiran-2-yl)(phenyl)methanol (3l): White solid, mp 141–142 °C, 17.3 mg, 36% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.24–7.15 (m, 8H), 7.04–7.01 (m, 2H), 4.94 (d, J = 6.9 Hz, 1H), 3.68 (q, J = 5.5 Hz, 1H), 2.37 (d, J = 6.9 Hz, 1H), 1.00 (d, J = 5.5 Hz, 3H); ¹³C NMR (150 MHz, CDCl₃) δ 140.2, 135.1, 128.2, 128.0, 127.9, 127.8, 127.6, 127.0, 76.1, 68.6, 57.3, 14.7; HRMS (ESI, m/z): calcd. for C₁₆H₁₆O₂Na⁺ 263.1043, found 263.1044; [α]_D²⁵: 12.4 (c 0.6, CHCl₃); HPLC analysis: 86% *ee* (Chiralcel AD-H, 0.5:99.5 PrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 38.4 min, R_t (minor) = 28.9 min; IR (KBr thin film, cm⁻¹): v 2988, 1556, 1454, 1414, 1280, 1263, 1230, 1130, 1016, 894, 769, 678.

(*R*)-1-((2*R*,3*S*)-3-phenyloxiran-2-yl)ethan-1-ol (2m): Colourless liquid, 14.1 mg, 43% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.36–7.26 (m, 5H), 4.12–4.08 (m, 1H), 3.94 (d, *J* = 2.2 Hz, 1H), 3.08 (dd, *J* = 2.8, 2.3 Hz, 1H), 2.17 (s, 1H), 1,31 (d, *J* = 6.5 Hz, 3H); ¹³C NMR (150 MHz, CDCl₃) δ 137.0, 128.6, 128.4, 125.8, 65.7, 64.9, 54.8, 18.8; HRMS (ESI, m/z): calcd. for C₁₀H₁₂O₂Na⁺ 187.0730, found 187.0729; $\left[\alpha\right]_D^{25}$: 7.4 (c 1.8, CHCl₃); HPLC analysis: 89% *ee* (Chiralcel AD-H, 10:90 ^{*i*}PrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 5.1 min, R_t (minor) = 4.2 min; IR (KBr thin film, cm⁻¹): v 2988, 1571, 1542, 1466, 1335, 1281, 1261, 1150, 1031, 951, 778, 681.

(*R*)-1-((2*S*,3*R*)-3-phenyloxiran-2-yl)ethan-1-ol (3m): Colourless liquid, 17.4 mg, 53% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.36–7.33 (m, 2H), 7.32–7.29 (m, 1H), 7.29–7.25 (m, 2H), 3.86 (d, J = 2.1 Hz, 1H), 3.04 (dd, J = 4.6, 2.2 Hz, 1H), 1.96 (d, J = 6.0 Hz, 1H), 1.35 (d, J = 6.5 Hz, 3H); ¹³C NMR (150 MHz, CDCl₃) δ 136.8, 128.6, 128.4, 125.8, 67.3, 66.4, 56.7, 20.2; HRMS (ESI, m/z): calcd. for C₁₀H₁₂O₂Na⁺ 187.0730, found 187.0729; [α]_D²⁵: +10.8 (c 0.3, CHCl₃); HPLC analysis: 56% *ee* (Chiralcel AD-H, 5:95 ⁱPrOH/Hexane, 1 mL/min, 220

nm), R_t (major) = 11.9 min, R_t (minor) = 7.1 min; IR (KBr thin film, cm⁻¹): v 2994, 1542, 1463, 1375, 1340, 1284, 1218, 1090, 1014, 900, 772, 701.

(2S,3R,4R)-2-phenyl-1-oxaspiro[2.5]octan-4-ol (2n): Yellow solid, mp 102–103 °C, 18.8 mg, 46% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.36–7.25 (m, 5H), 4.34 (s, 1H), 3.88 (ddd, J = 10.9, 4.3, 1.6 Hz, 1H), 2.24–2.14 (m, 2H), 1.77–1.73 (m, 1H), 1.64–1.62 (m, 1H), 1.54–1.46 (m, 2H), 1.41–1.33 (m, 2H), 1.05–0.96 (m, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 135.8, 128.1, 127.4, 126.3, 70.6, 68.5, 58.8, 33.3, 27.1, 24.6, 23.9; HRMS (ESI, m/z): calcd. for $C_{13}H_{16}O_2Na^+$ 227.1043, found 227.1042; $[\alpha]_D^{25}$: +12.5 (c 0.8, CHCl₃); HPLC analysis: >99% *ee* (Chiralcel AD-H, 0.5:99.5 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 13.9 min, R_t (minor) = 17.4 min; IR (KBr thin film, cm⁻¹): v 2983, 1539, 1463, 1420, 1278, 1264, 1221, 1019, 931, 897, 772, 693.

(2*R*,3*S*,4*R*)-2-phenyl-1-oxaspiro[2.5]octan-4-ol (3n): White solid, mp 64–65 °C, 20.7 mg, 51% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.35–7.32 (m, 2H), 7.29–7.26 (m, 3H), 4.30 (s, 1H), 3.73 (td, J = 9.1, 4.2 Hz, 1H), 2.01–1.95 (m, 1H), 1.78 (d, J = 8.9 Hz, 1H), 1.79–1.75 (m, 1H), 1.64–1.59 (m, 1H), 1.56–1.52 (m, 1H), 1.52–1.43 (m, 2H), 1.42–1.34 (m, 1H), 1.25–1.19 (m, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 135.7, 128.1, 127.5, 126.4, 70.3, 67.2, 60.7, 33.9, 26.1, 23.6, 23.0; HRMS (ESI, m/z): calcd. for C₁₃H₁₆O₂Na⁺ 227.1043, found 227.1044; [α]_D²⁵: +8.2 (c 1.1, CHCl₃); HPLC analysis: 96% *ee* (Chiralcel AD-H, 0.5:99.5 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 29.5 min, R_t (minor) = 12.0 min; IR (KBr thin film, cm⁻¹): v 2929, 1539, 1463, 1420, 1278, 1264, 1221, 1019, 931, 897, 772, 693.

(*R*)-((2*R*,3*S*)-2-methyl-3-phenyloxiran-2-yl)(phenyl)methanol (2o): Colourless liquid, 17.3 mg, 36% yield. 1 H NMR (600 MHz, CDCl₃) δ 7.47–7.45 (m, 2H), 7.41–7.38 (m, 2H), 7.35–

7.31 (m, 3H), 7.28–7.24 (m, 3H), 4.83 (s, 1H), 4.51 (s, 1H), 2.74 (s, 1H), 0.98 (s, 3H); 13 C NMR (150 MHz, CDCl₃) δ 139.6, 135.4, 128.7, 128.5, 128.2, 127.7, 127.4, 126.4, 75.4, 66.1, 59.7, 13.7; HRMS (ESI, m/z): calcd. for $C_{16}H_{16}O_2Na^+$ 263.1043, found 263.1045; $[\alpha]_D^{25}$: 2.6 (c 1.2, CHCl₃); HPLC analysis: 58% *ee* (Chiralcel AD-H, 10:90 ⁱPrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 6.5 min, R_t (minor) = 4.8 min; IR (KBr thin film, cm⁻¹): v 2934, 1553, 1454, 1412, 1340, 1280, 1261, 1093, 980, 852, 775, 687.

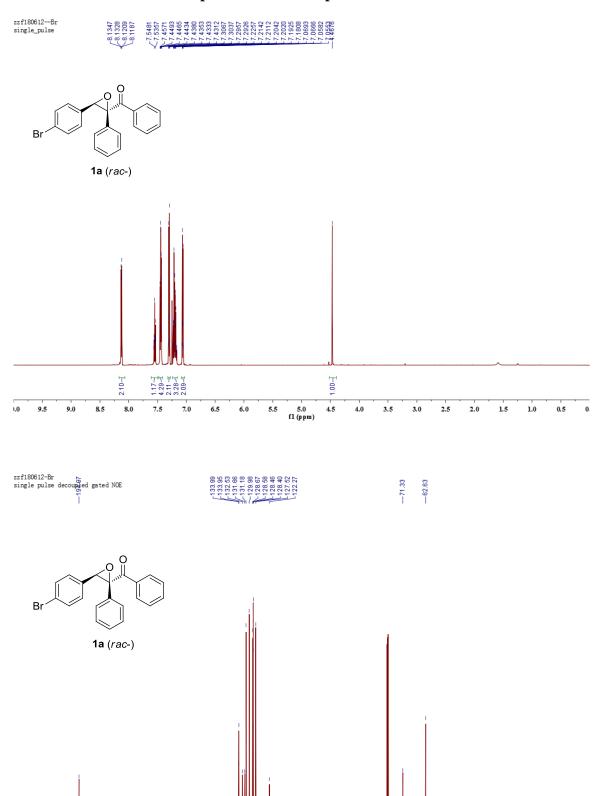
(*R*)-((2*S*,3*R*)-2-methyl-3-phenyloxiran-2-yl)(phenyl)methanol (3o): White solid, mp 97–98 $^{\circ}$ C, 25.0 mg, 52% yield. 1 H NMR (600 MHz, CDCl₃) δ 7.48–7.43 (m, 2H), 7.41–7.26 (m, 8H), 4.66 (s, 1H), 4.36 (s, 1H), 2.54 (s, 1H), 0.92 (s, 3H); 13 C NMR (150 MHz, CDCl₃) δ 140.3, 135.4, 128.7, 128.3, 128.1, 127.8, 126.6, 126.5, 77.3, 66.5, 61.7, 11.9; HRMS (ESI, m/z): calcd. for C₁₆H₁₆O₂Na⁺ 263.1043, found 263.1042; $[\alpha]_D^{25}$: +23.5 (c 0.8, CHCl₃); HPLC analysis: 34% *ee* (Chiralcel AD-H, 10:90 i PrOH/Hexane, 1 mL/min, 220 nm), R_t (major) = 6.4 min, R_t (minor) = 5.4 min; IR (KBr thin film, cm⁻¹): v 2858, 1562, 1451, 1402, 1340, 1287, 1224, 1059, 928, 852, 772, 693.

((2*R*,3*R*)-3-(4-fluorophenyl)-2-phenyloxiran-2-yl)(phenyl)methanone (4a): White solid, mp 93–94 °C, 15.3 mg, 96% yield. ¹H NMR (600 MHz, CDCl₃) δ 8.14–8.12 (m, 2H), 7.57–7.53 (m, 1H), 7.46–7.42 (m, 4H), 7.23–7.13 (m, 5H), 6.88–6.84 (m, 2H), 4.49 (s, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 195.2, 162.6 (d, J = 245.4 Hz), 134.1, 133.9, 131.8, 130.0, 129.2 (d, J = 2.5 Hz), 128.7, 128.54, 128.48, 128.3, 127.6, 115.1 (d, J = 21.7 Hz), 71.3, 62.6; HRMS (ESI, m/z): calcd. for C₂₁H₁₅O₂FH⁺ 319.1129, found 319.1132; [α]_D²⁵: 25.0 (c 0.9, CHCl₃); HPLC analysis: 91% *ee* (Chiralcel AD-H, 5:95 ⁱPrOH/Hexane, 1 mL/min), R_t (major) = 5.3 min, R_t (minor) = 4.8 min; IR (KBr thin film, cm⁻¹): v 3037, 1687, 1601, 1582, 1510, 1446, 1275, 1187, 1031, 838, 775, 698.

(1*R*,2*R*,3*R*)-3-azido-3-(4-fluorophenyl)-1,2-diphenylpropane-1,2-diol (4b): White solid, mp 78–79 °C, 10.7 mg, 98% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.16–7.09 (m, 10H), 6.91–6.88 (m, 2H), 6.85–6.82 (m, 2H), 5.53 (d, J = 3.2 Hz, 1H), 5.28 (s, 1H), 2.70 (s, 1H), 2.64 (d, J = 3.3 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) δ 162.6 (d, J = 245.3 Hz), 139.1, 138.2, 131.3 (d, J = 2.4 Hz), 130.9, 130.8, 128.14, 128.07, 128.0, 127.4, 126.8 (d, J = 7.6 Hz), 114.7 (d, J = 21.3 Hz), 80.1, 76.0, 69.7; HRMS (ESI, m/z): calcd. for C₂₁H₁₈O₂N₃Na⁺ 386.1277, found 386.1279; [α]_D²⁵: 27.4 (c 1.1, CHCl₃); HPLC analysis: 91% *ee* (Chiralcel AD-H, 15:85 ⁱPrOH/Hexane, 1 mL/min), R_t (major) = 6.6 min, R_t (minor) = 4.8 min; IR (KBr thin film, cm⁻¹): v 2108, 1556, 1508, 1409, 1340, 1278, 1258, 1221, 1090, 772, 704, 678.

(4c): White solid, mp 94–95 °C, 18.0 mg, 88% yield. ¹H NMR (600 MHz, CDCl₃) δ 7.37–7.28 (m, 8H), 7.24–7.22 (m, 2H), 7.11–7.08 (m, 1H), 7.06–7.03 (m, 2H), 6.91–6.85 (m, 4H), 6.76–6.72 (m, 2H), 4.74 (s, 1H), 4.64 (d, J = 12.1 Hz, 1H), 4.57 (s, 1H), 4.68 (d, J = 12.1 Hz, 1H); ¹³C NMR (150 MHz, CDCl₃) 162.2 (d, J = 244.2 Hz), 138.2, 137.3, 134.2, 131.0 (d, J = 2.4 Hz), 129.1, 128.5, 128.3, 128.2, 128.1, 128.0, 127.7, 127.6, 127.5, 127.4, 114.7 (d, J = 21.5 Hz), 83.3, 71.1, 71.0, 60.7; HRMS (ESI, m/z): calcd. for C₂₈H₂₃FO₂H⁺ 411.1755, found 411.1756; $[\alpha]_D^{25}$: +11.4 (c 2.9, CHCl₃); HPLC analysis: 96% *ee* (Chiralcel AD-H, 1:99 ⁱPrOH/Hexane, 1 mL/min), R_t (major) = 17.3 min, R_t (minor) = 5.9 min; IR (KBr thin film, cm⁻¹): v 1618, 1565, 1537, 1451, 1394, 1335, 1281, 1264, 1099, 1025, 778, 690.

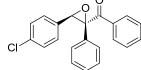
VII. ¹H NMR and ¹³C NMR spectra of new compounds

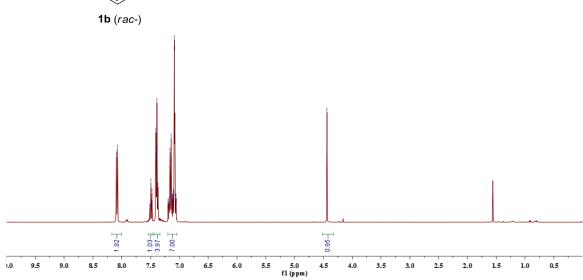


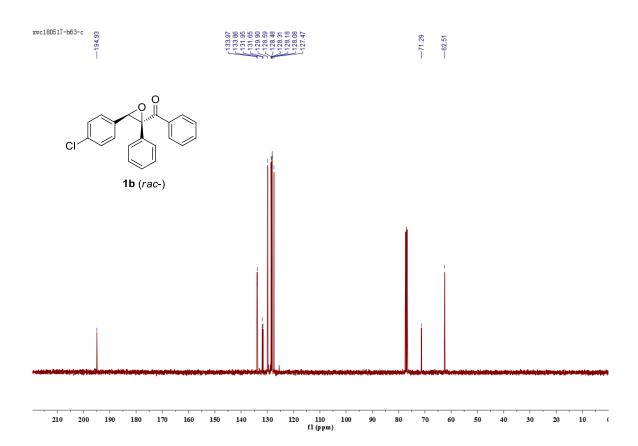
110 f1 (ppm)



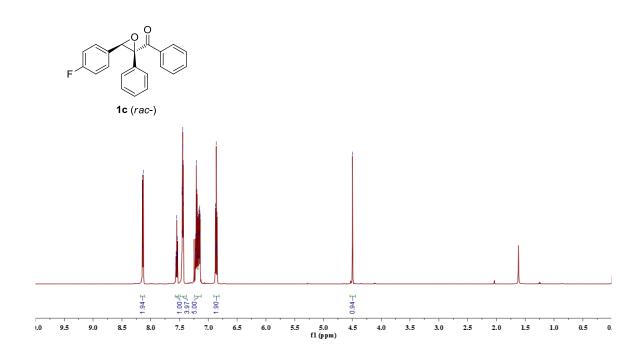


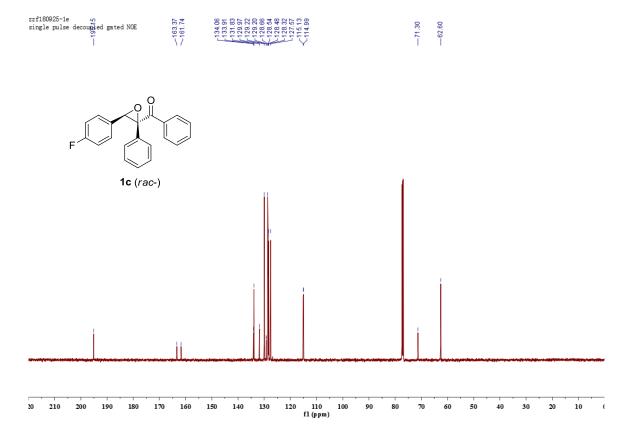




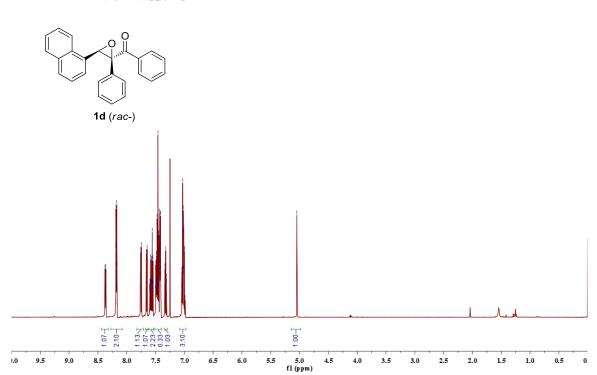


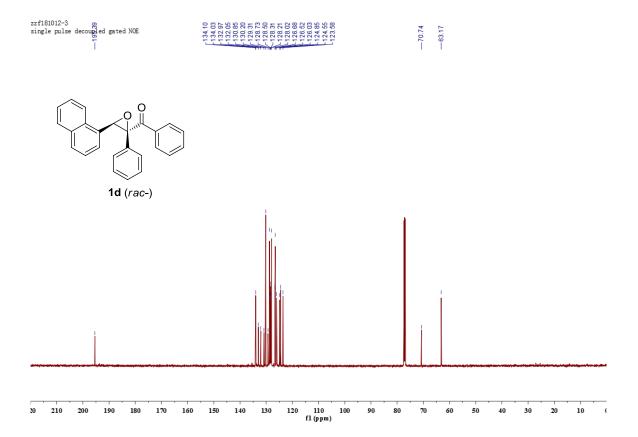


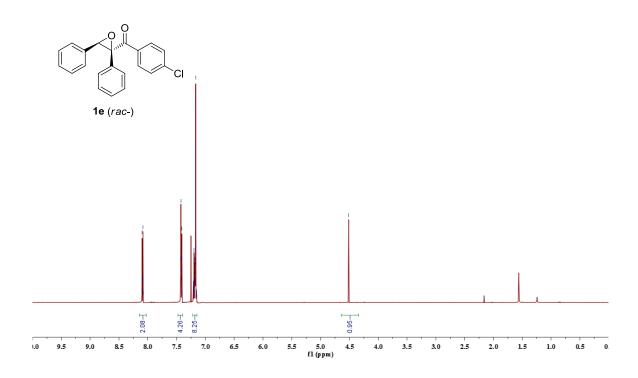


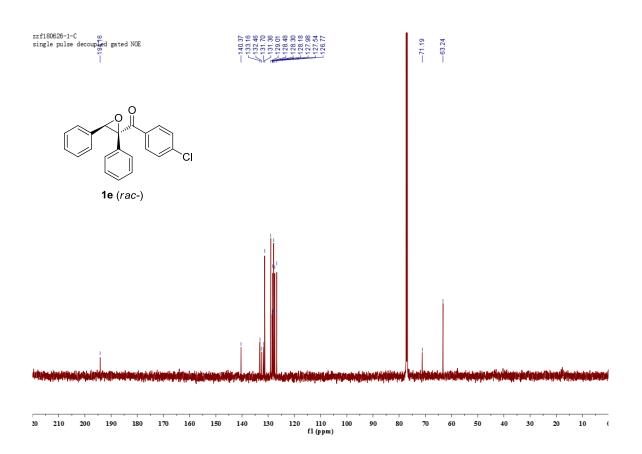


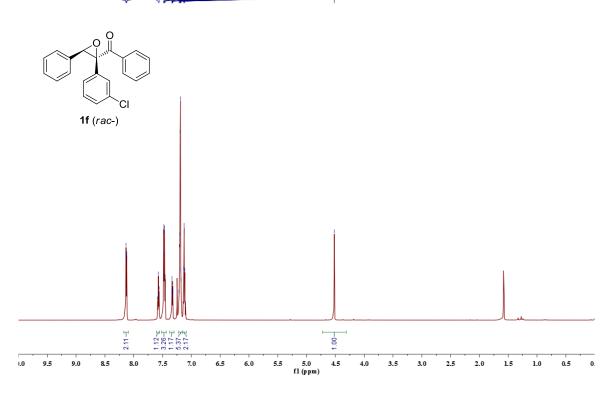


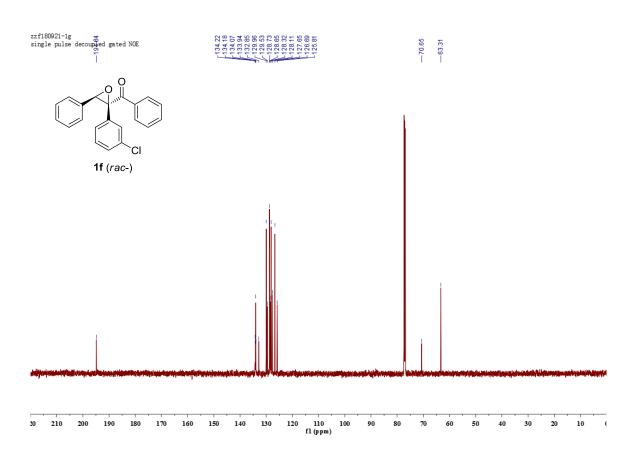












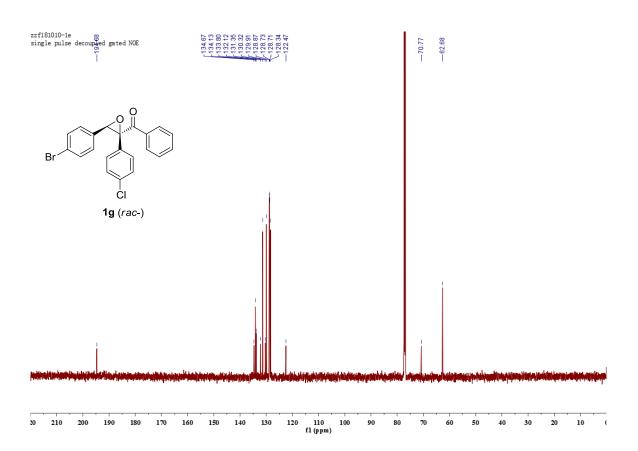
1.00-I

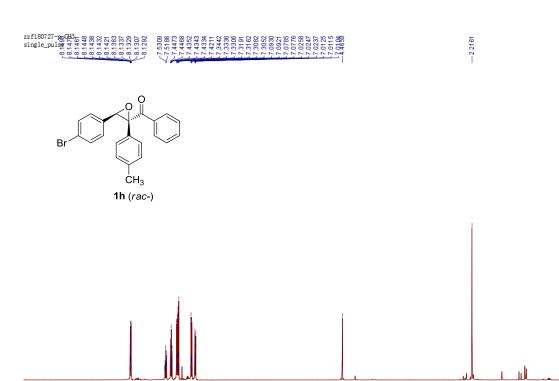
5.0 f1 (ppm)

2.04∃

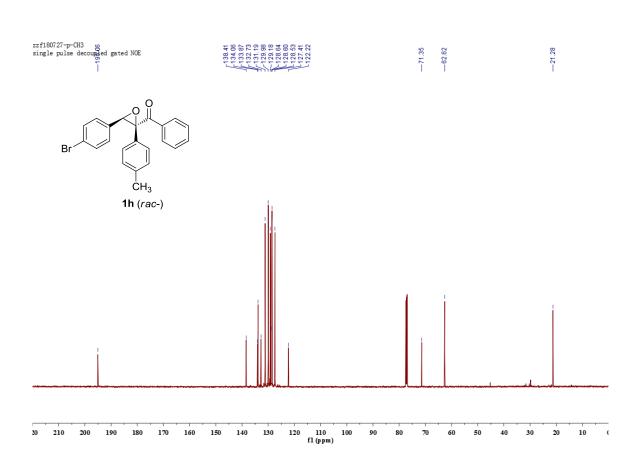
8.0

2.07 2.07 2.07 2.00 1.99 2.02

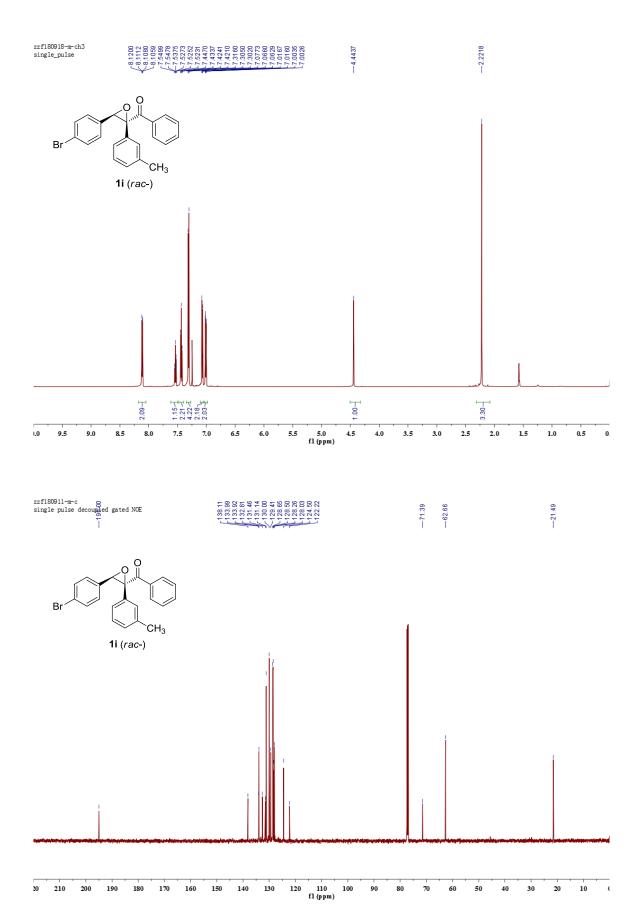


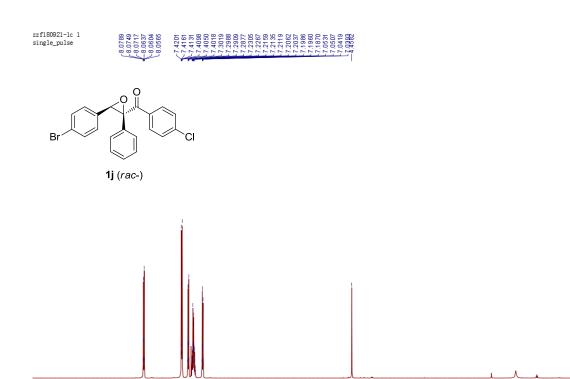


1.95 ±



5.0 f1 (ppm) 3.08±

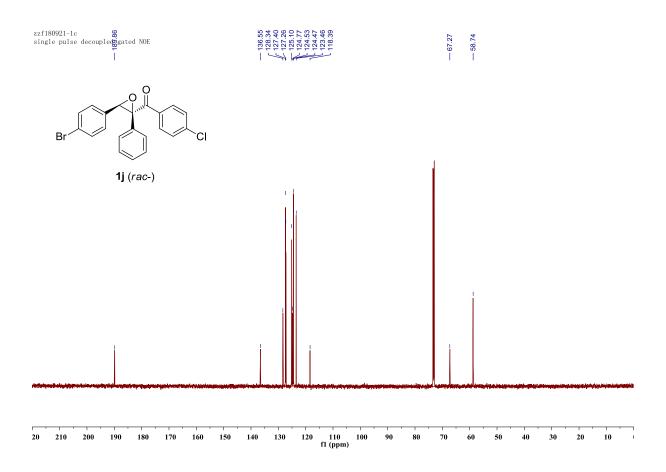




2.08-J

8.0

2.094 2.094 2.094 2.094

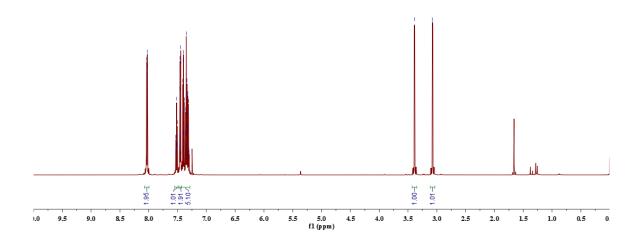


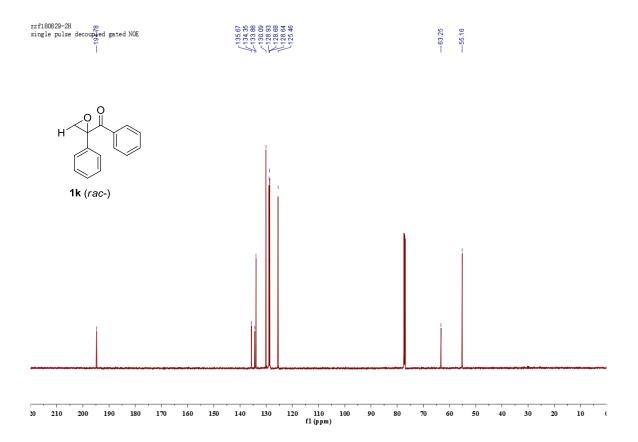
1.00-J

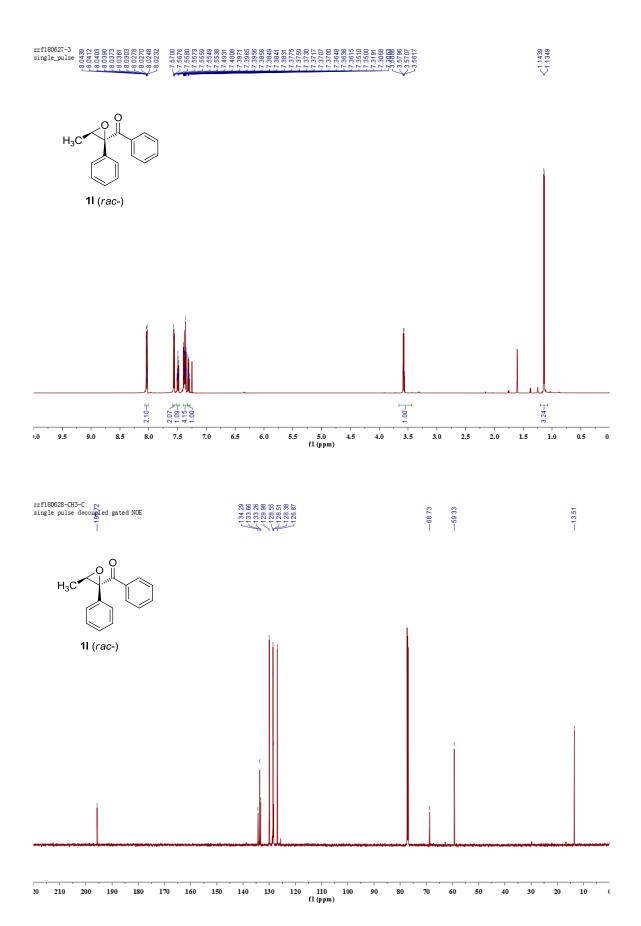
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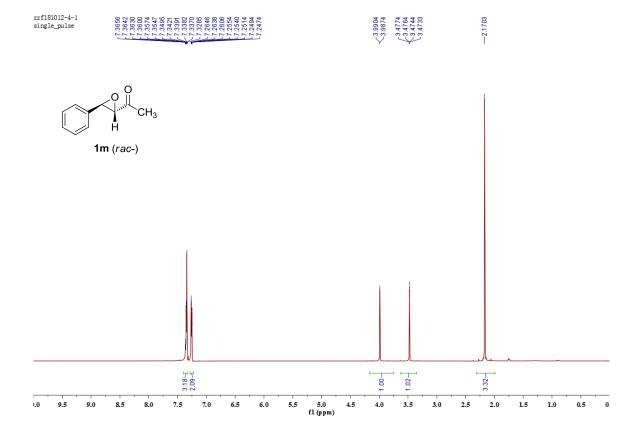


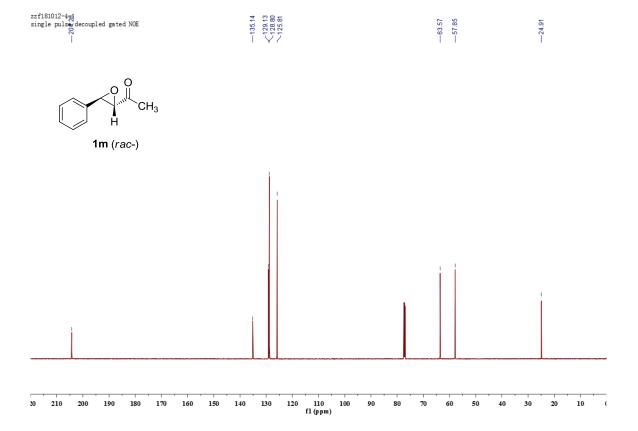
1k (rac-)



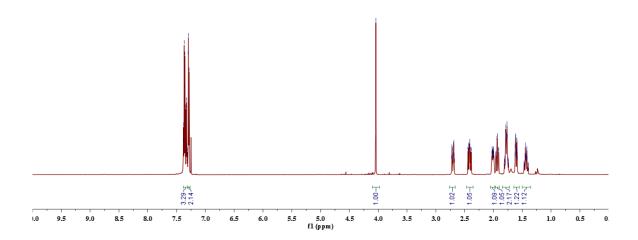


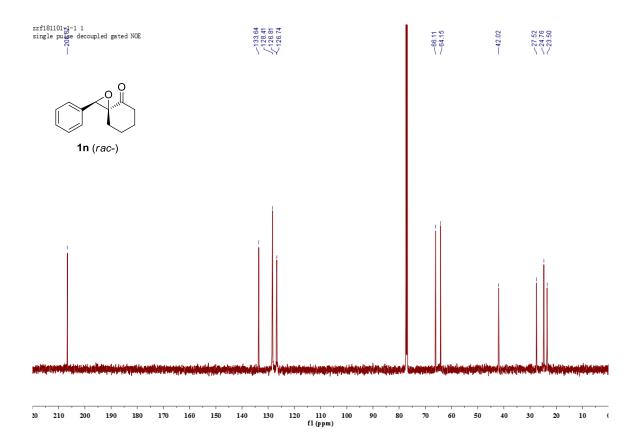


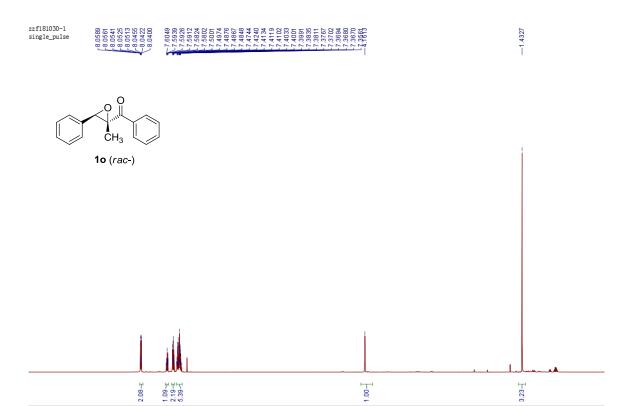




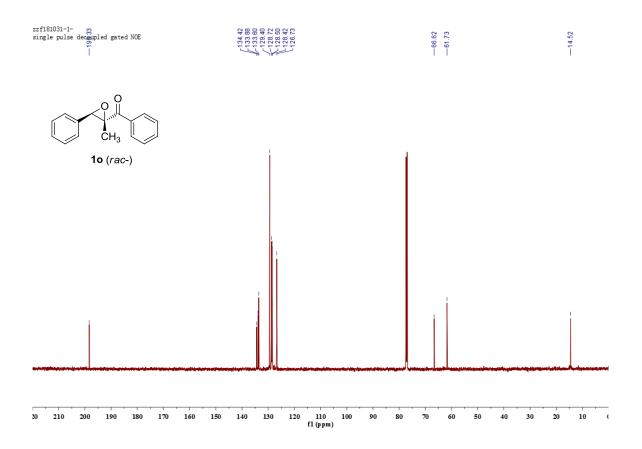


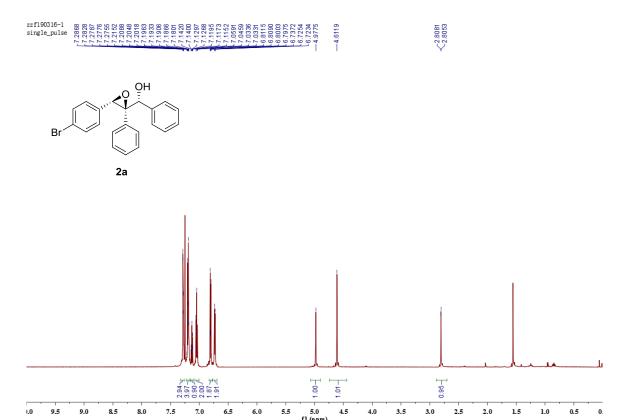


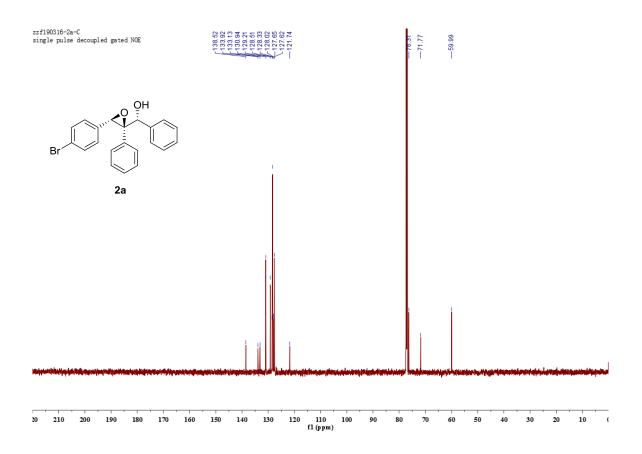


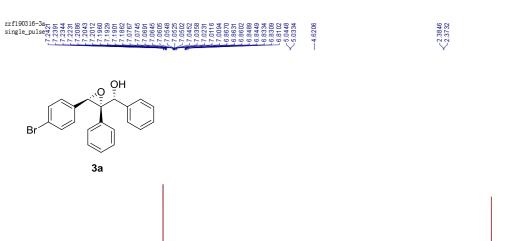


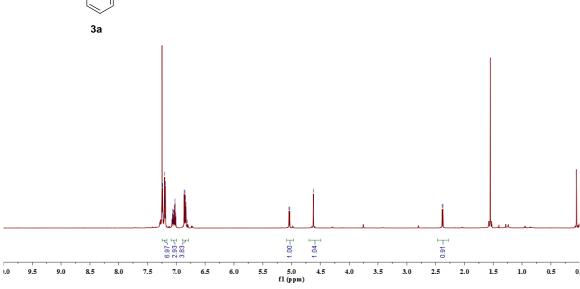
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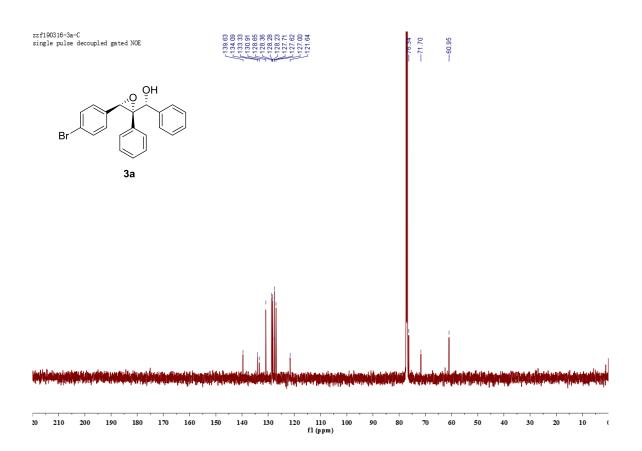


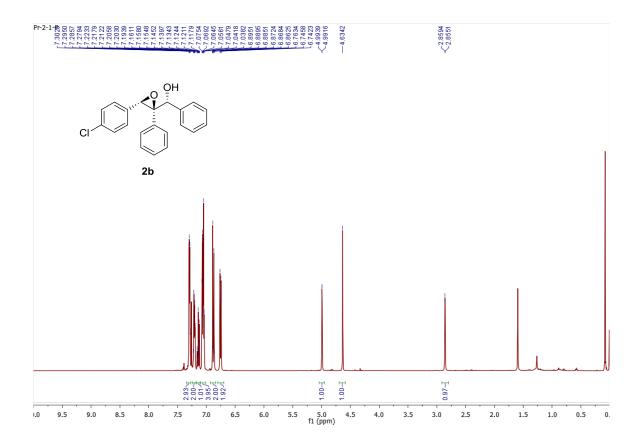


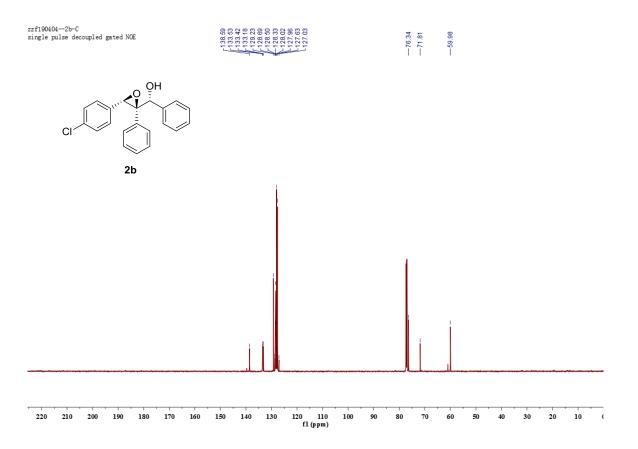


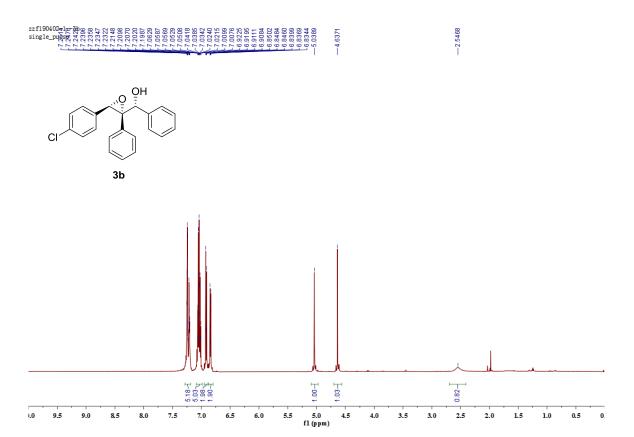


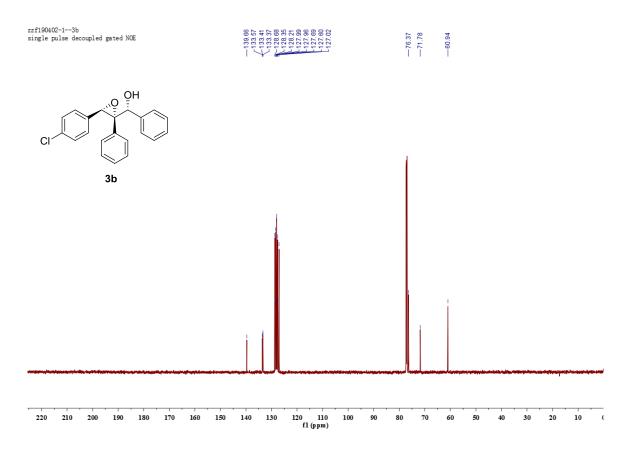


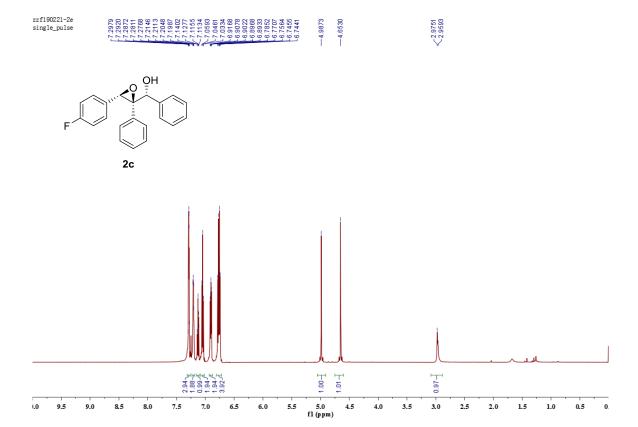


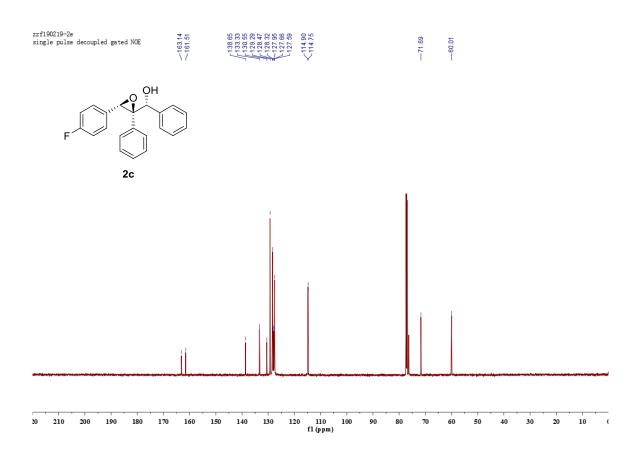


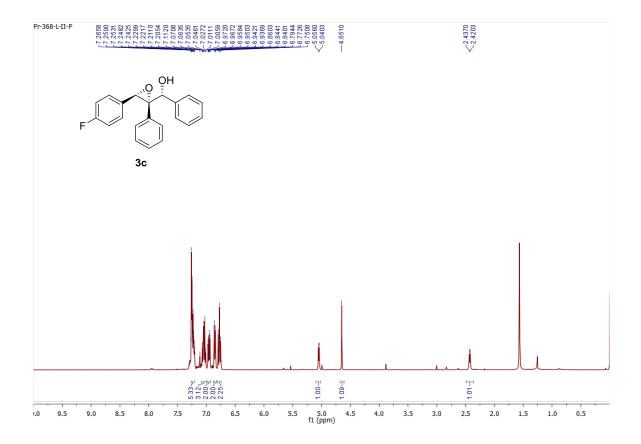


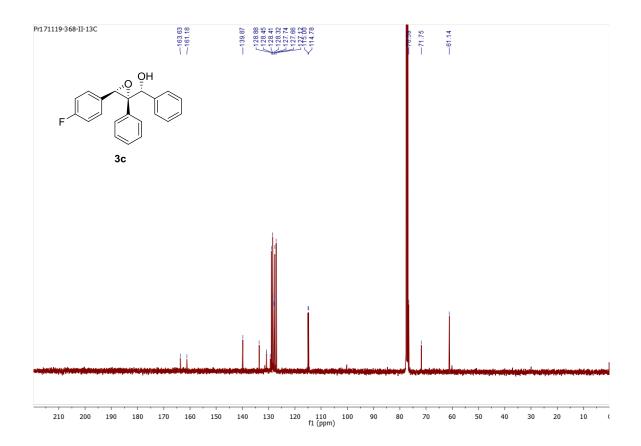


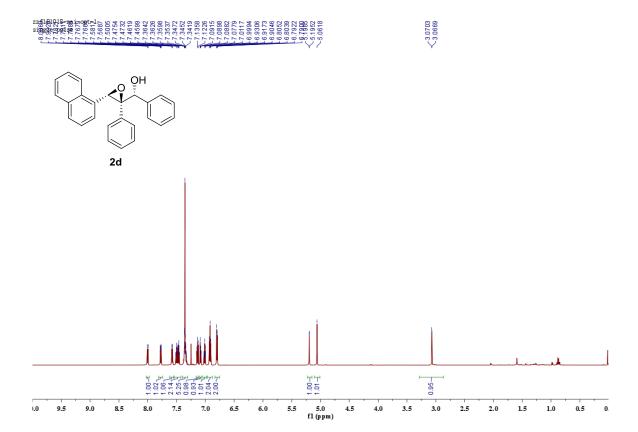


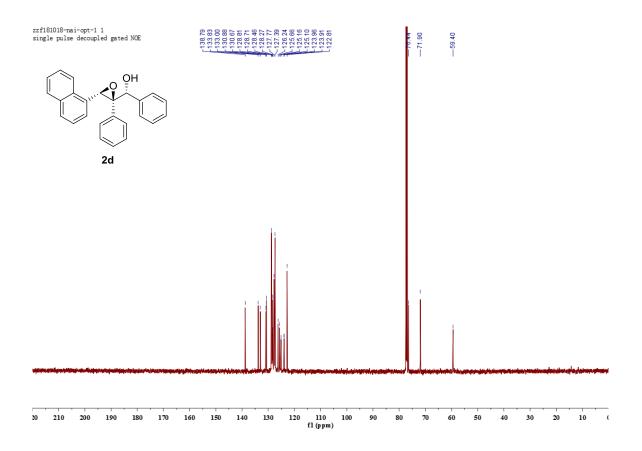


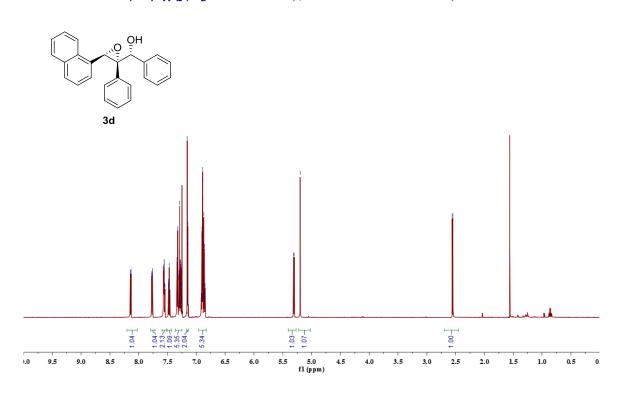


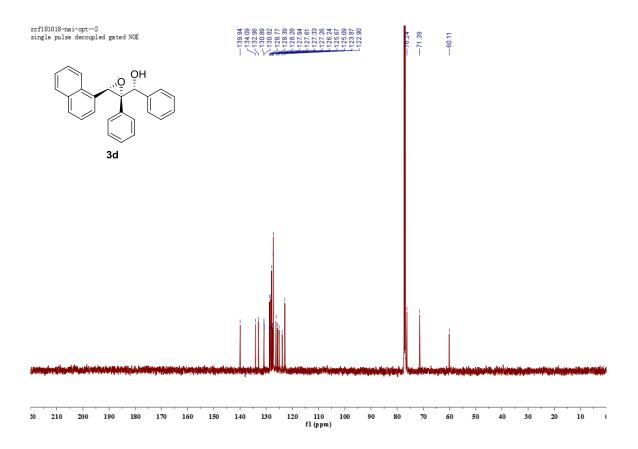


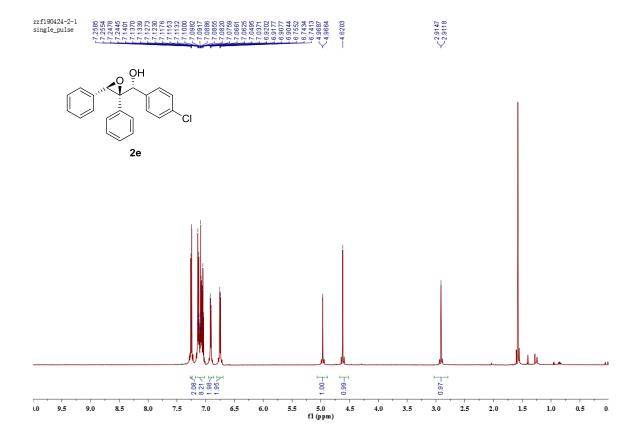


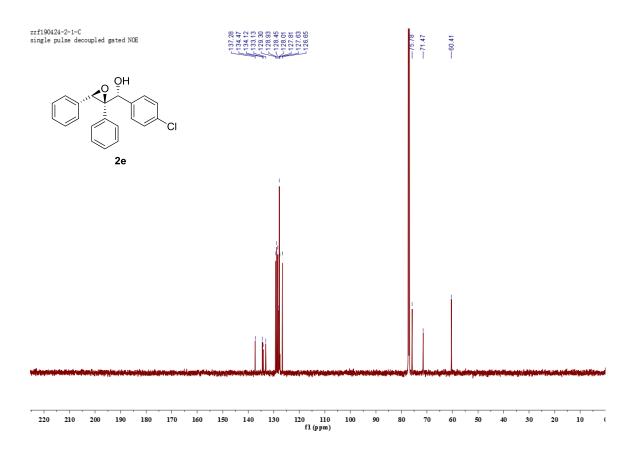


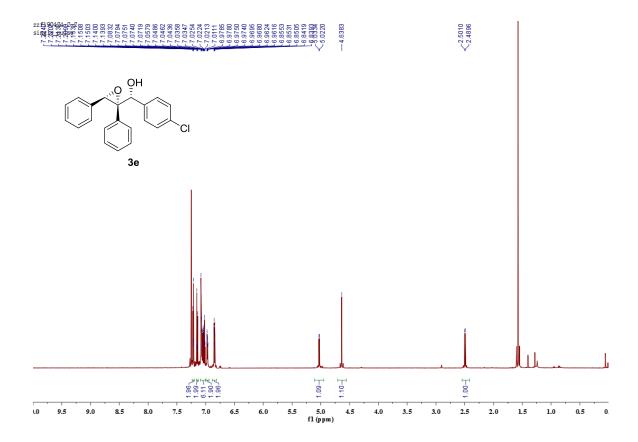


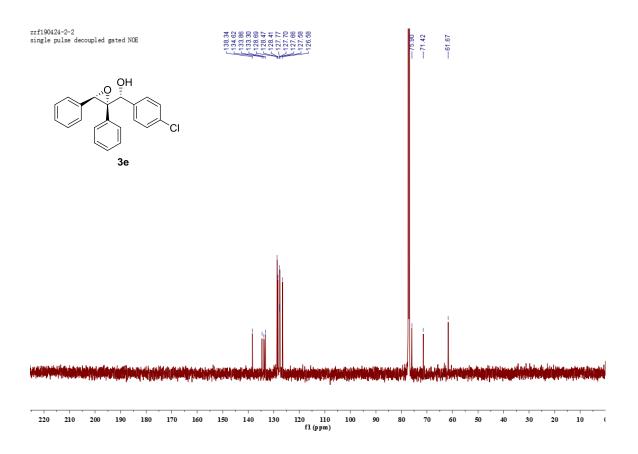


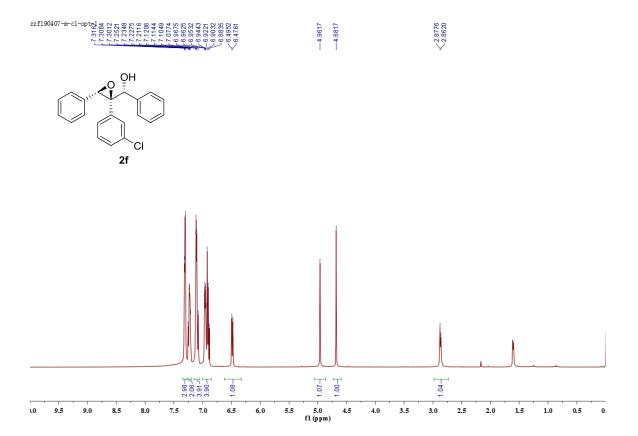


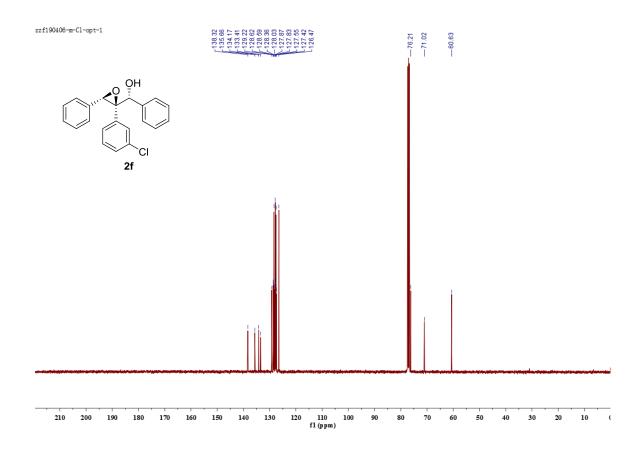


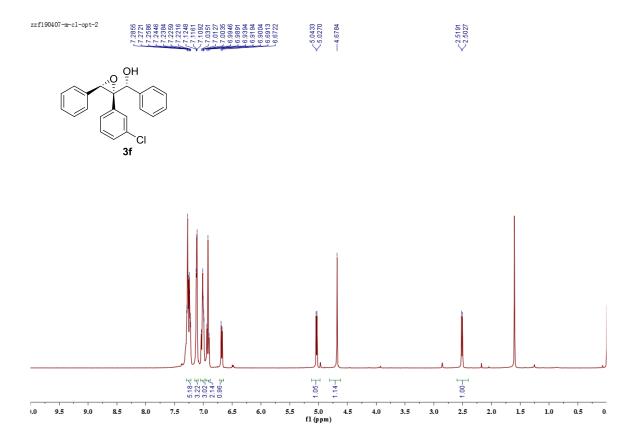


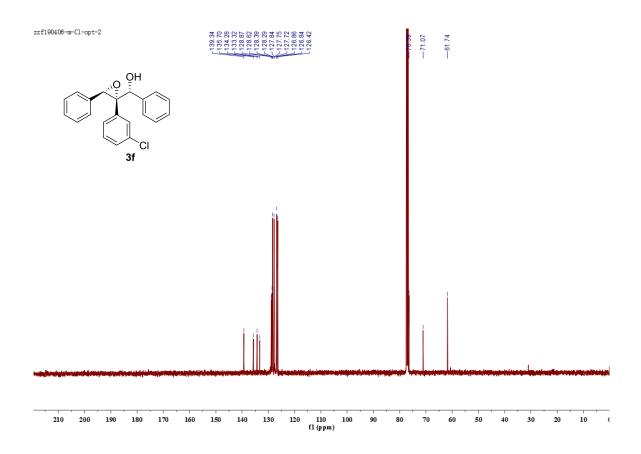


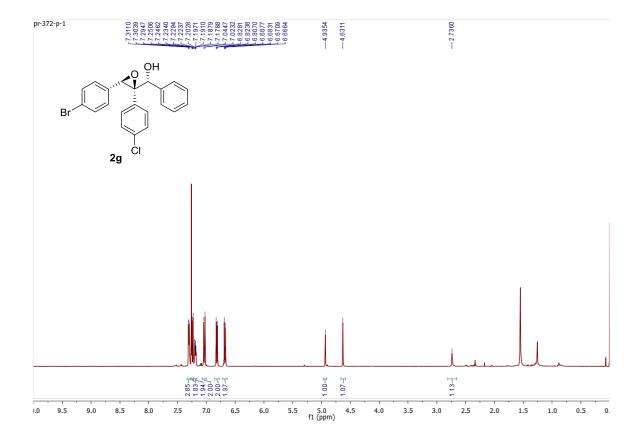


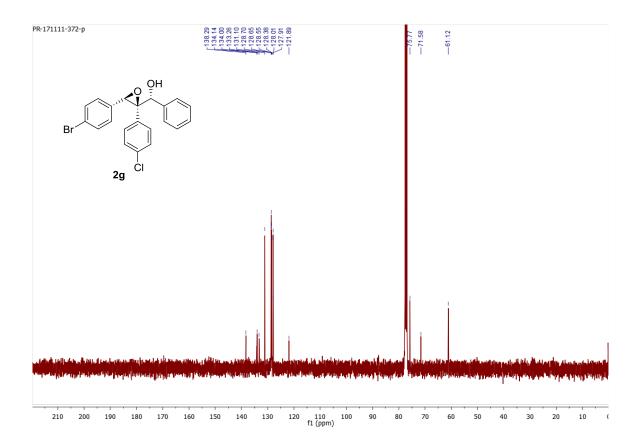


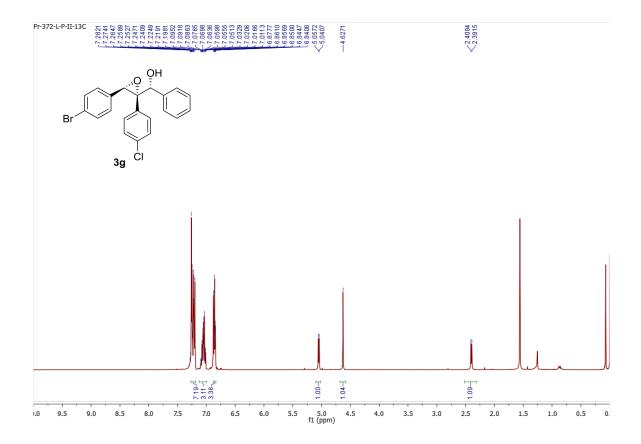


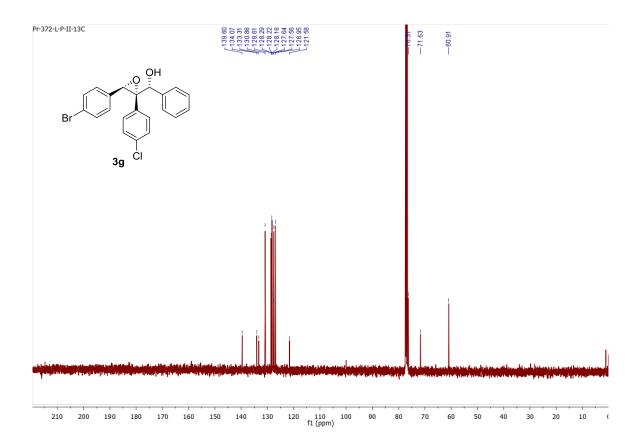


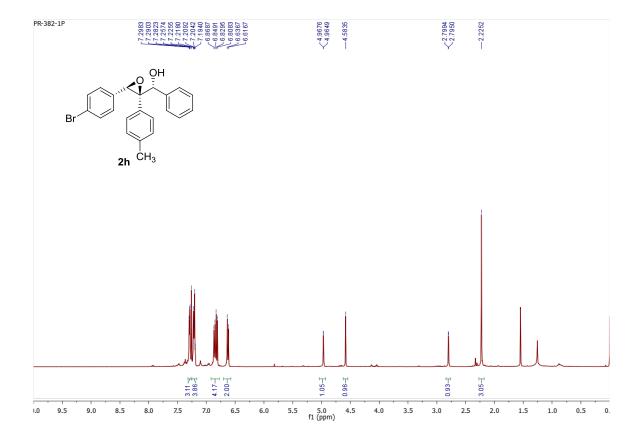


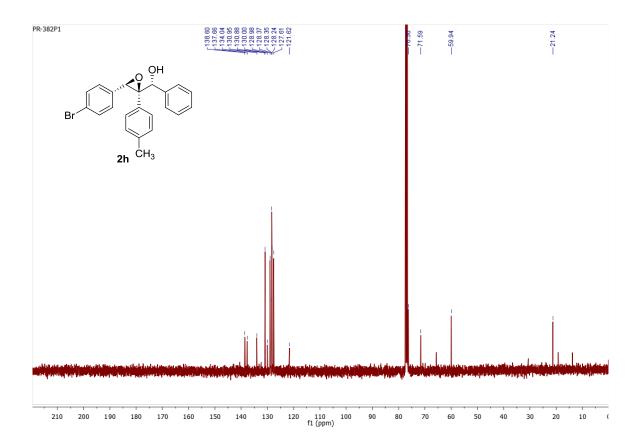


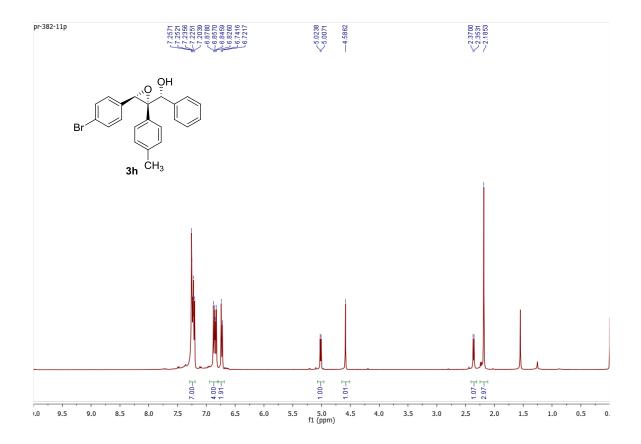


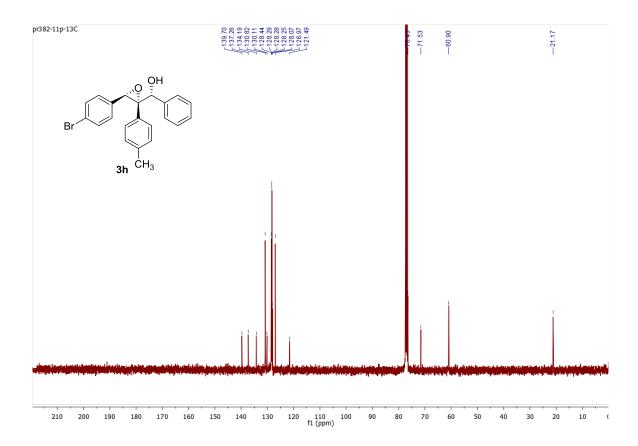


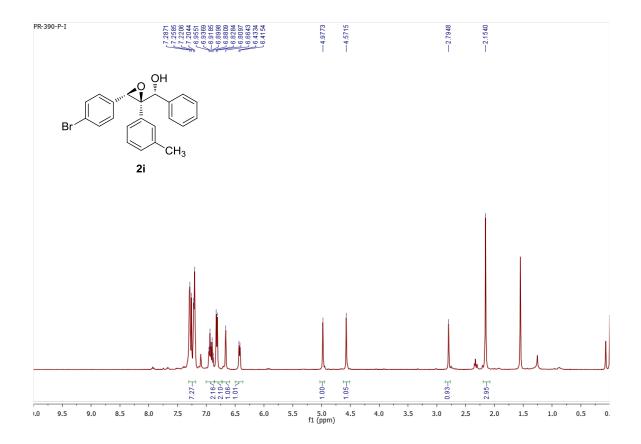


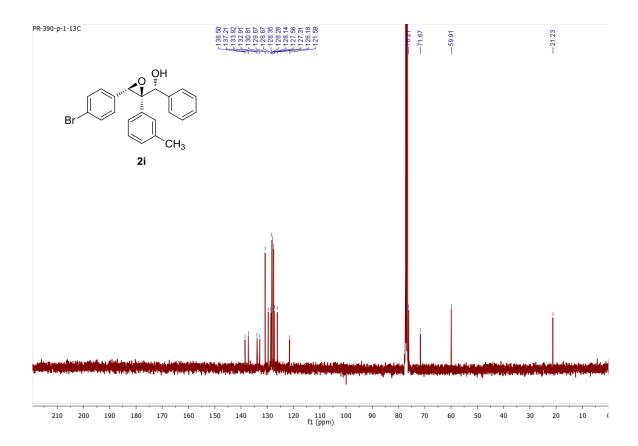


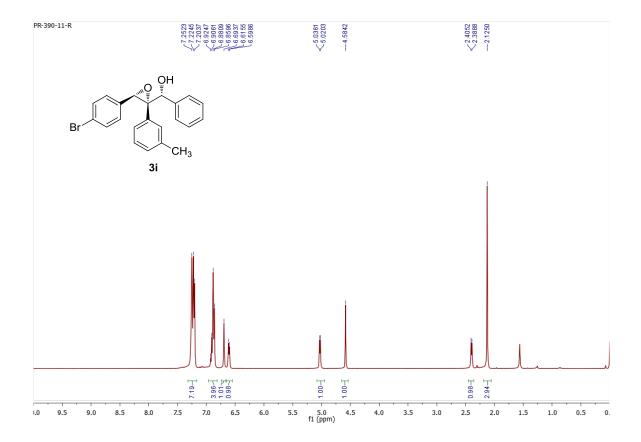


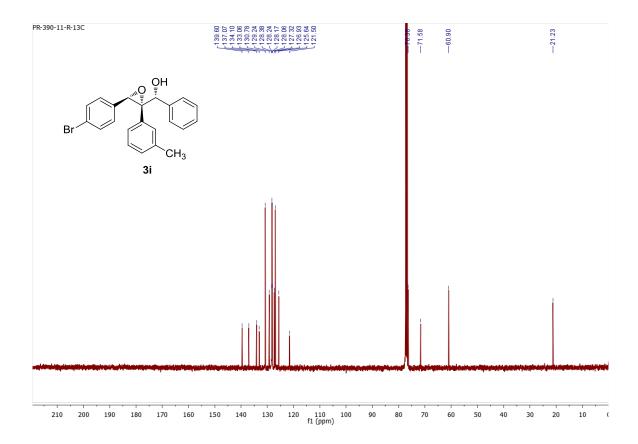


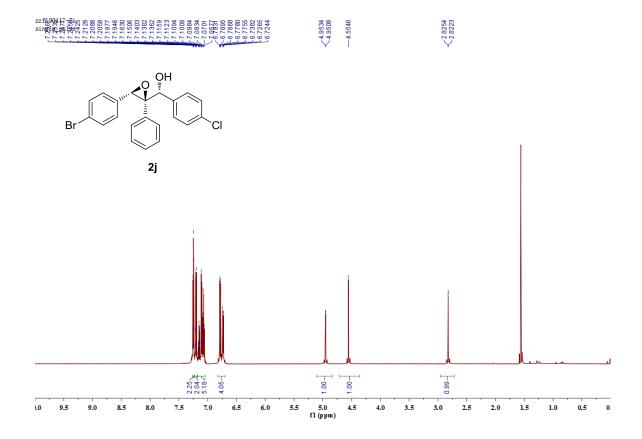


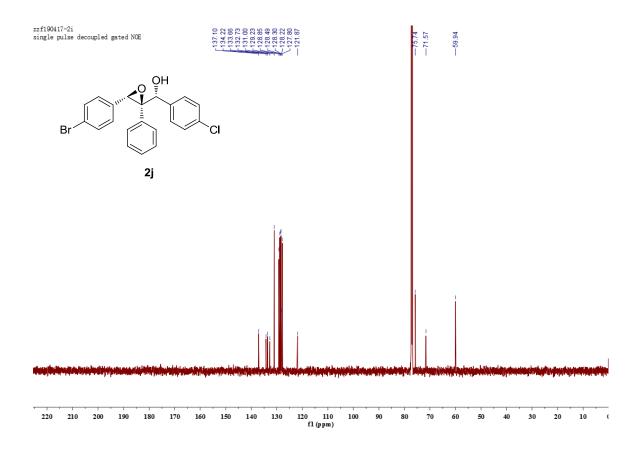


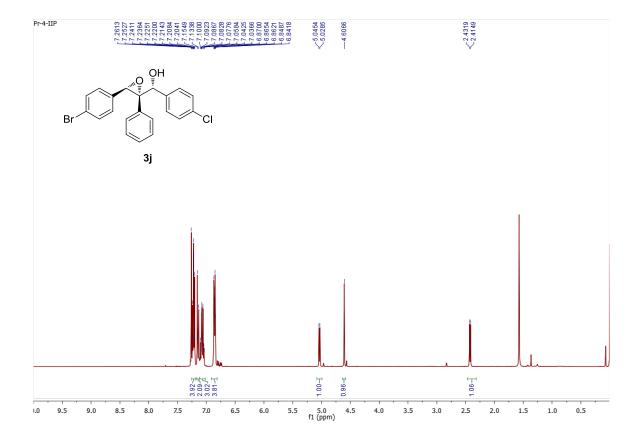


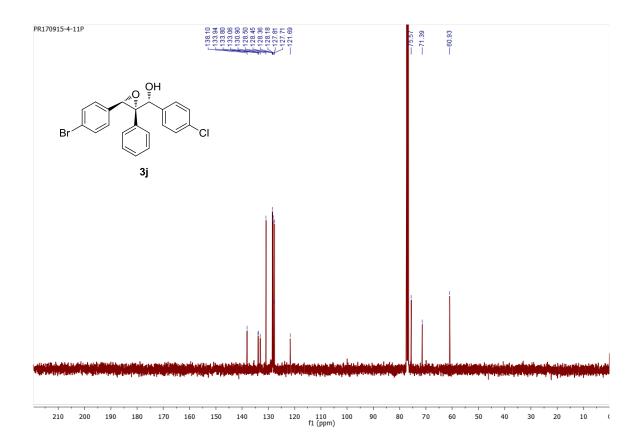


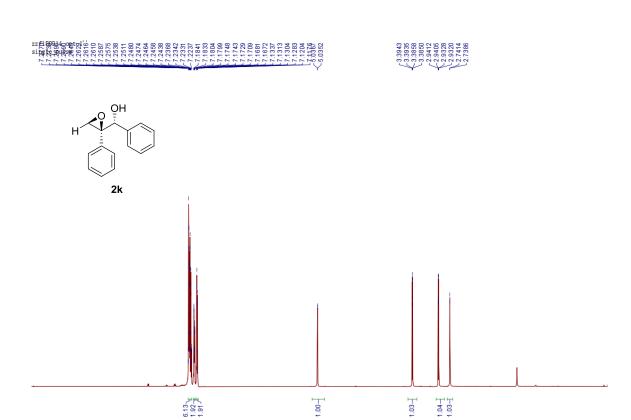




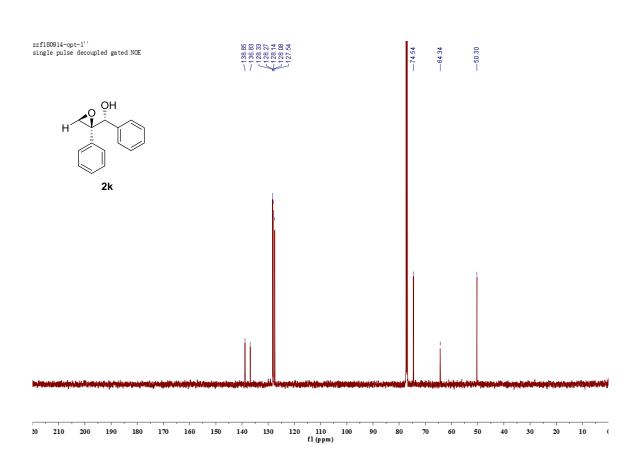








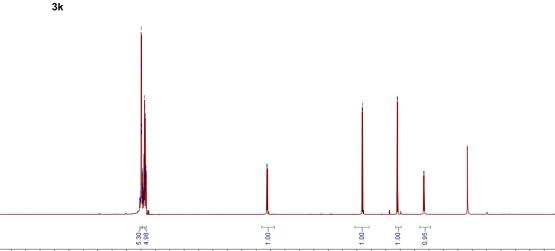
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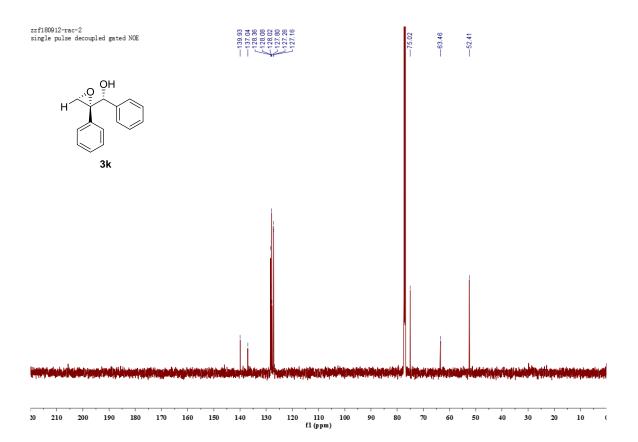






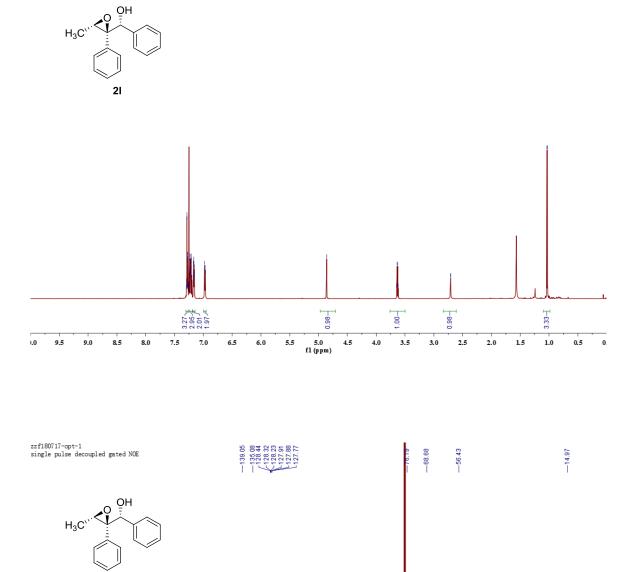


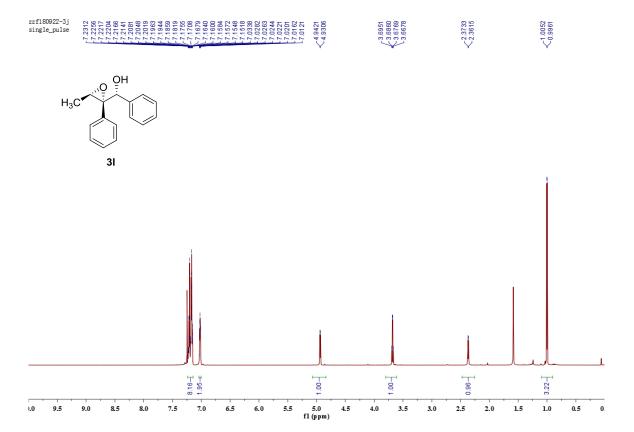


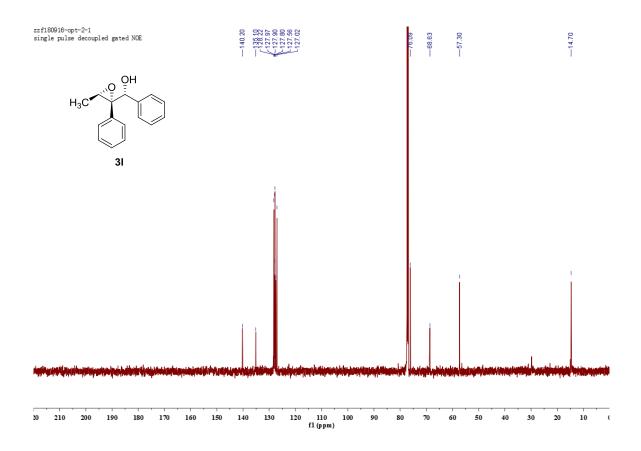


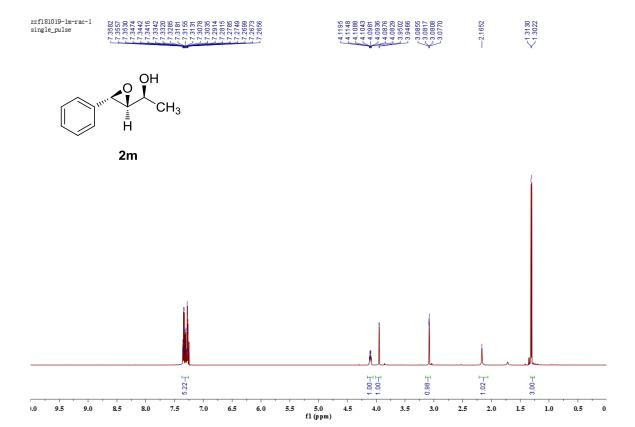
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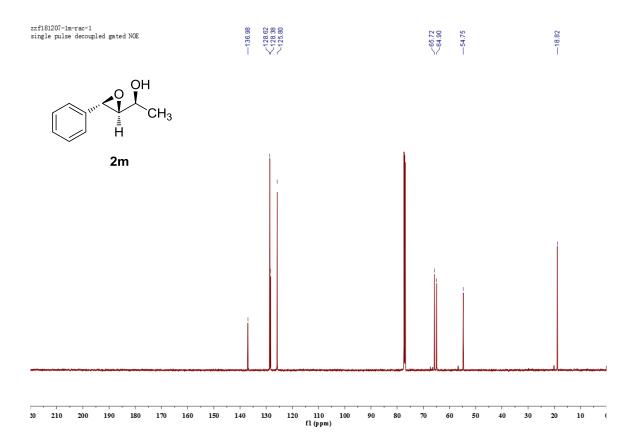
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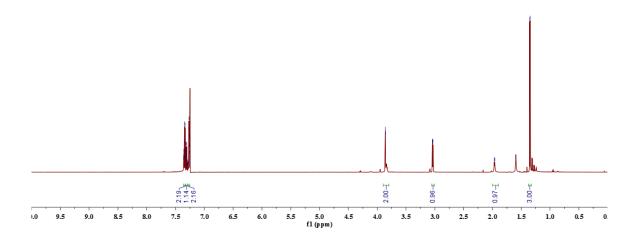


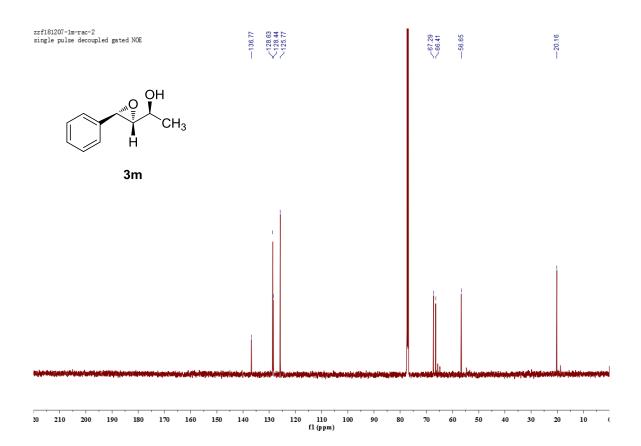


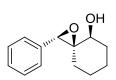




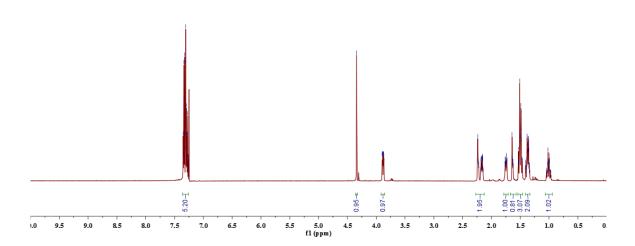
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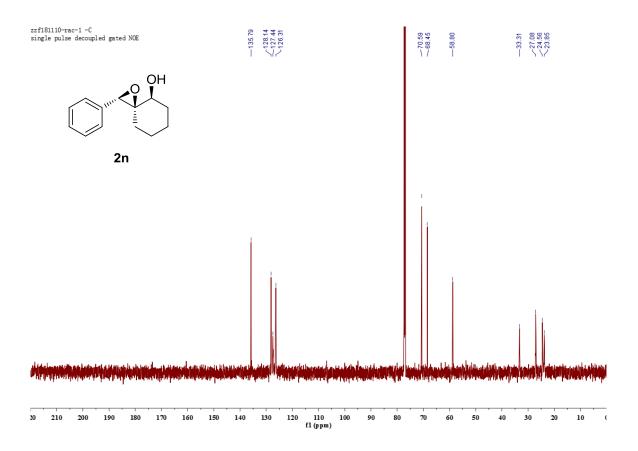


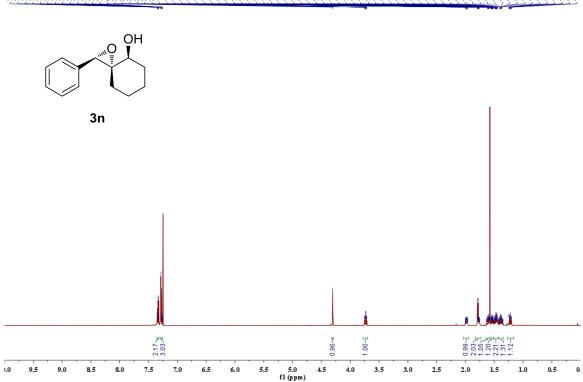


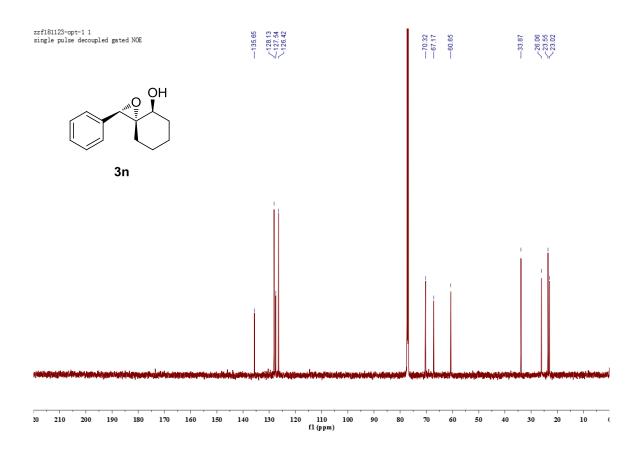


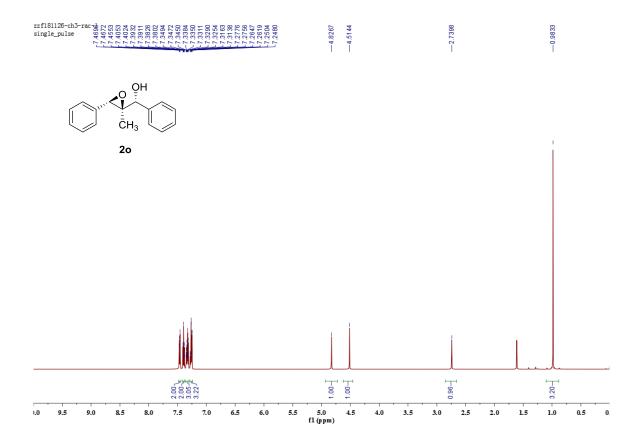
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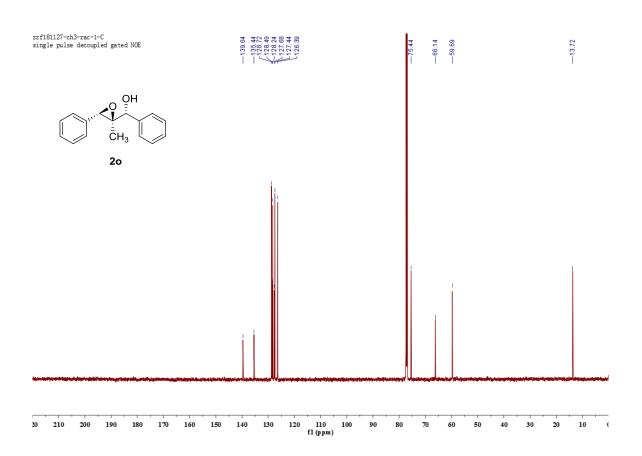


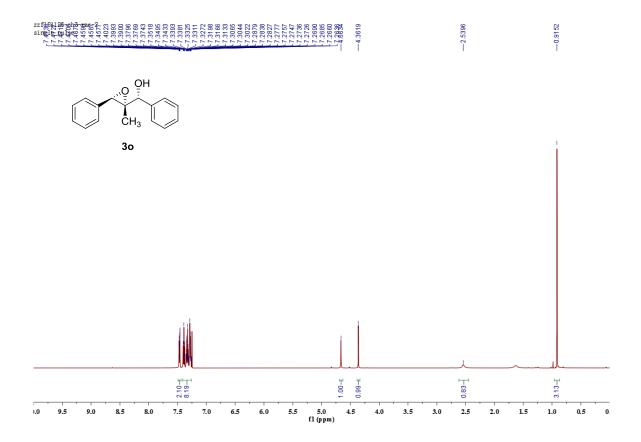


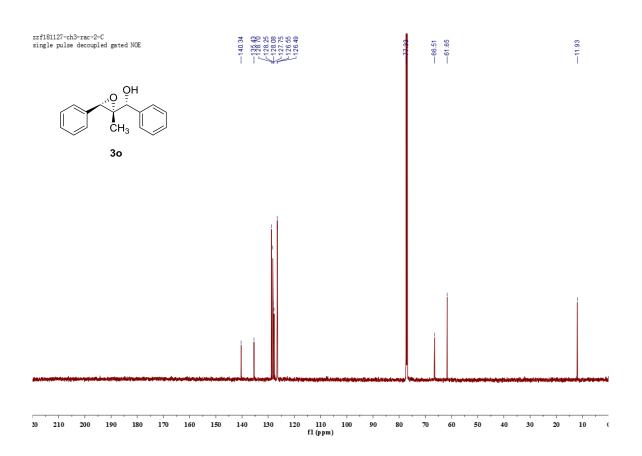




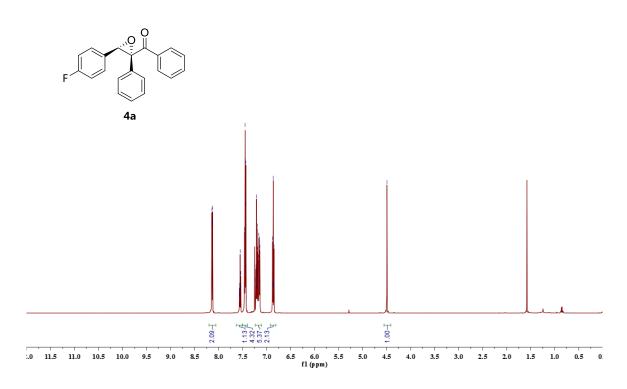


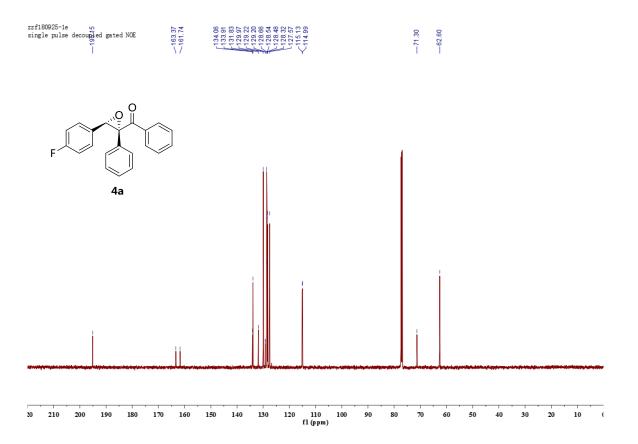


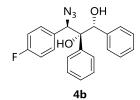


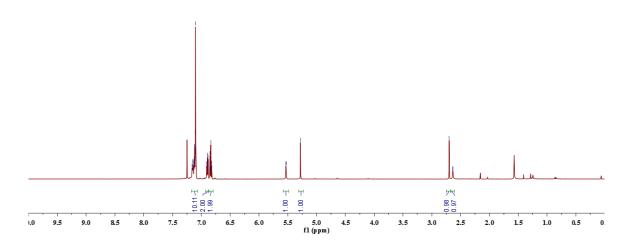


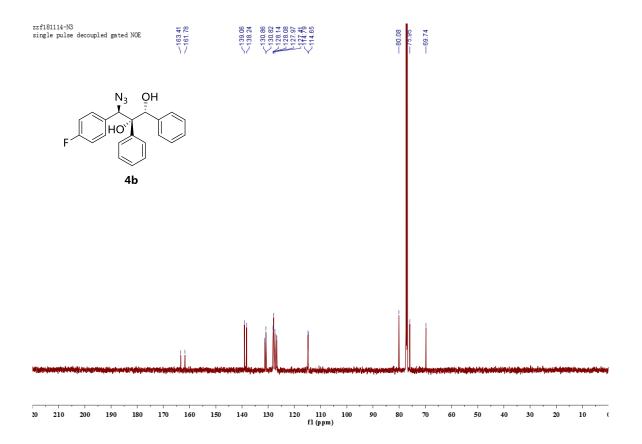




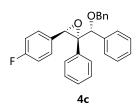


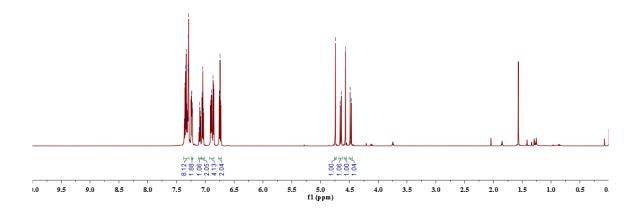


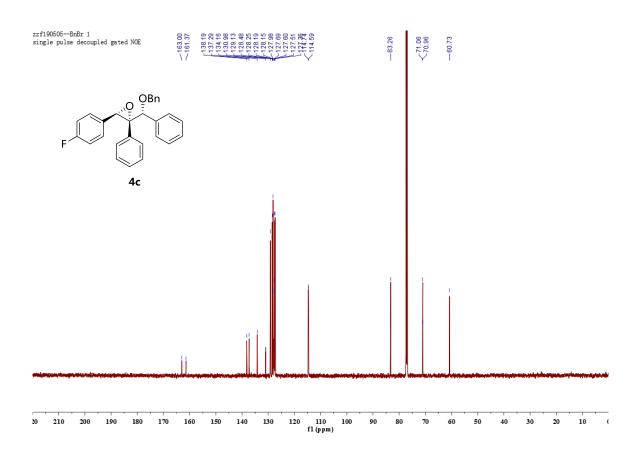




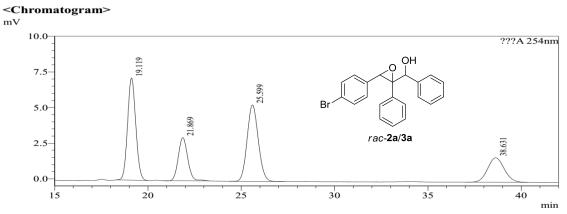






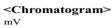


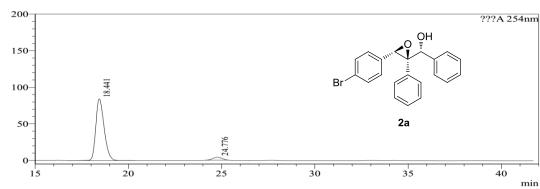
VII. HPLC spectra for ee determination



<Peak Table>

???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	19.119	219108	7174	33.458		M	
2	21.869	106002	3029	16.187		M	
3	25.599	223470	5366	34.124		M	
4	38.631	106299	1718	16.232		M	
Total		654879	17288				

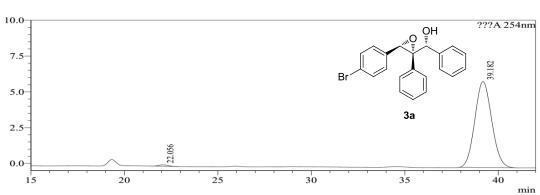




<Peak Table>

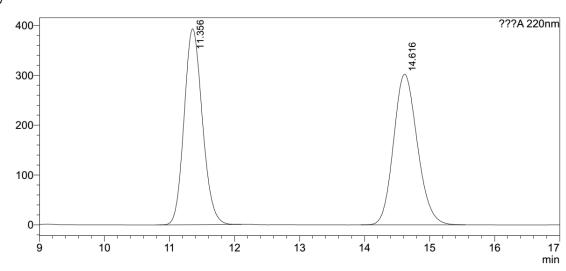
???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	18.441	2611990	84313	94.853		M	
2	24.776	141732	4017	5.147		M	
Total		2753722	88330				

<Chromatogram>



???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	22.056	3458	103	0.909		M	
2	39.182	376934	6007	99.091			
Total		380392	6110				

m\/



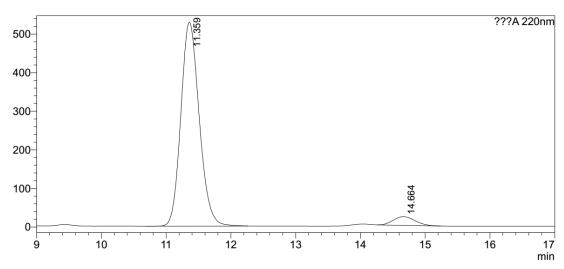
<Peak Table>

???A 220nm

11174	22011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	11.356	7661543	393042	49.873		M	
2	14.616	7700661	302395	50.127		M	
Total	l	15362205	695437				

<Chromatogram>

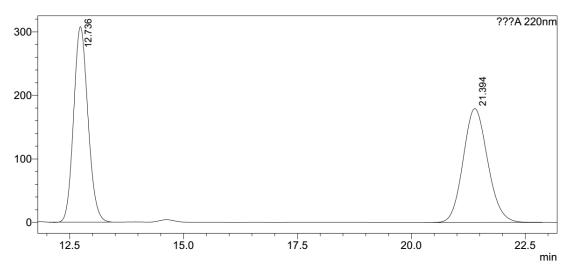
mV



<Peak Table>

11172	.2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	11.359	10413121	528467	95.189		M	
2	14.664	526292	22458	4.811		М	
Total		10939413	550925				

m٧



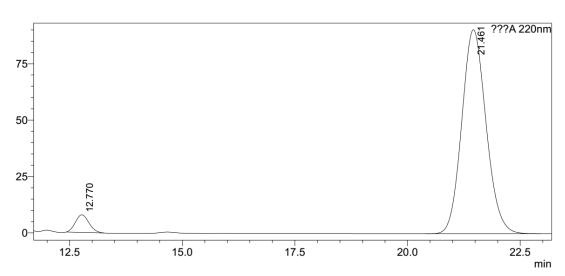
<Peak Table>

???A 220nm

:::/\	2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	12.736	6726175	307668	50.004		M	
2	21.394	6725066	179315	49.996		M	
Total		13451241	486983				

<Chromatogram>

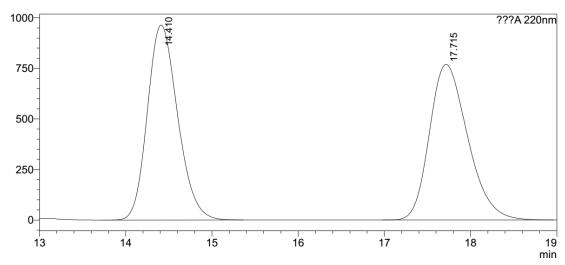
mV



<Peak Table>

/ \ 2							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	12.770	163013	7782	4.582		M	
2	21.461	3394424	90429	95.418		М	
Total		3557436	98211				

m۷



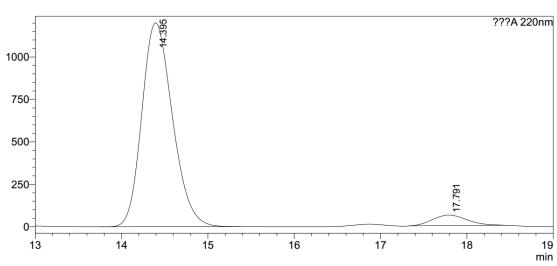
<Peak Table>

???A 220nm

	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	14.410	23536389	964336	49.793		M	
2	17.715	23731635	769385	50.207		M	
Total		47268024	1733721				

<Chromatogram>

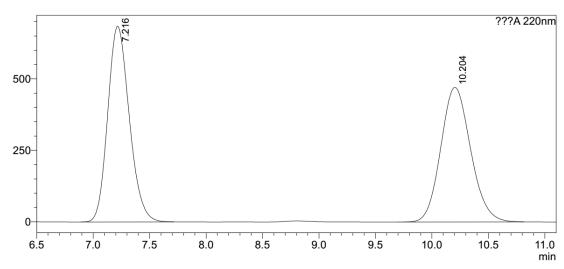
mV



<Peak Table>

Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	14.395	29603986	1200910	94.395		M	
2	17.791	1757831	62392	5.605		M	
Total		31361817	1263302				

mV



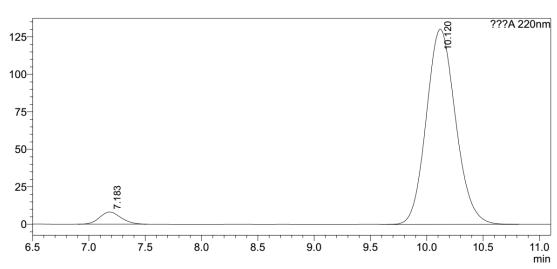
<Peak Table>

???A 220nm

/\ 2							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	7.216	9092209	684677	51.238		M	
2	10.204	8652988	470743	48.762		M	
Total		17745197	1155420				

<Chromatogram>

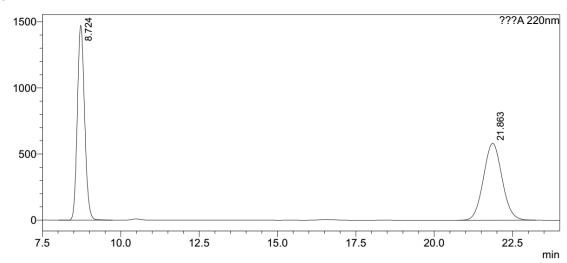
mV



<Peak Table>

!!!A	220nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	7.183	107548	8125	4.368		М	
2	10.120	2354834	130184	95.632		М	
Tota	I	2462382	138308				

mV



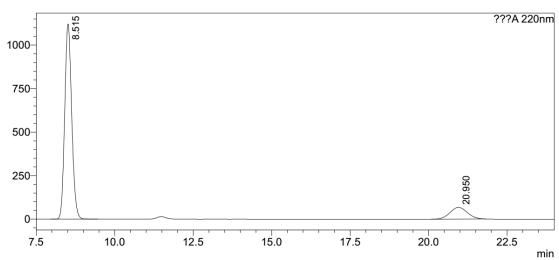
<Peak Table>

???A 220nm

:::/\2	2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	8.724	23800414	1472978	49.713		M	
2	21.863	24075627	581331	50.287		M	
Total		47876042	2054308				

<Chromatogram>

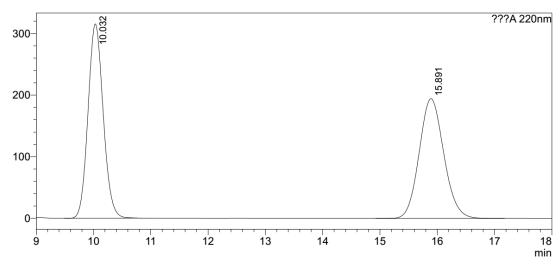
mV



<Peak Table>

$f \in \mathcal{F} \cap \mathcal{F}$	201111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	8.515	17326555	1120669	86.929		M	
2	20.950	2605372	67423	13.071		М	
Total		19931926	1188092				

mV



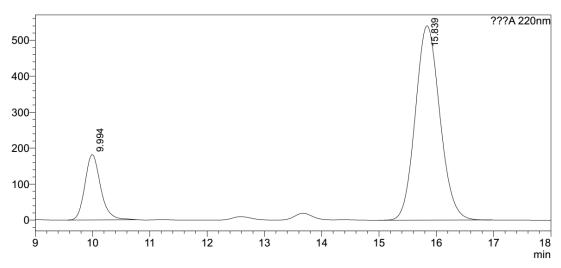
<Peak Table>

???A 220nm

	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	10.032	5822909	315519	50.092		M	
2	15.891	5801485	194483	49.908		M	
Total		11624394	510002				

<Chromatogram>

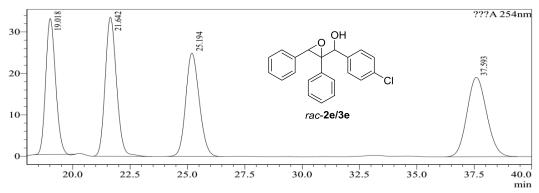
mV



<Peak Table>

Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	9.994	3462413	181667	17.711		M	
2	15.839	16086628	540070	82.289		M	
Total		19549040	721738				

mV

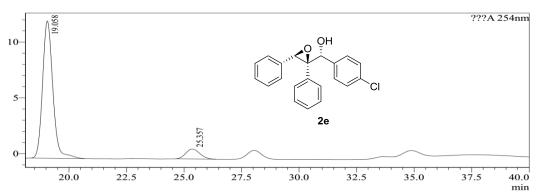


<Peak Table>

???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	19.018	981427	32804	22.809		M	
2	21.642	1155451	33556	26.854			
3	25.194	1024659	24938	23.814			
4	37.593	1141250	19141	26.524			
Total		4302787	110439				

<Chromatogram>

mV

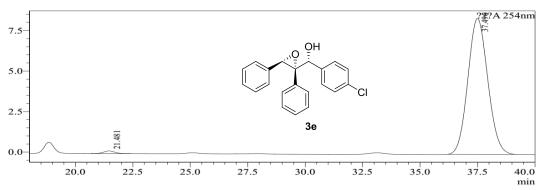


<Peak Table>

???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	19.058	382687	12312	90.959			
2	25.357	38037	899	9.041			
Total		420724	13211				

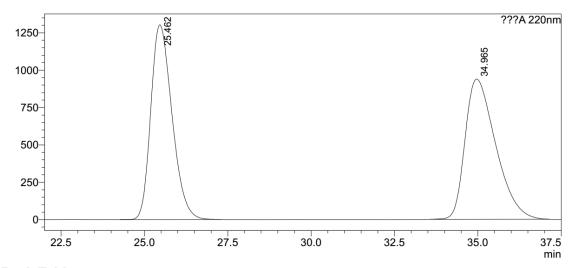
<Chromatogram>

mV



???A 2	???A 254nm											
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name					
1	21.481	5251	158	1.028		M						
2	37.496	505682	8381	98.972								
Total		510933	8539									

m۷



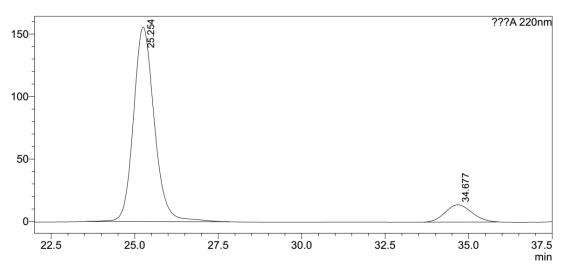
<Peak Table>

???A 220nm

:::/\ 4							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	25.462	57744367	1302569	49.036		M	
2	34.965	60014746	938444	50.964		М	
Total		117759112	2241012				

<Chromatogram>

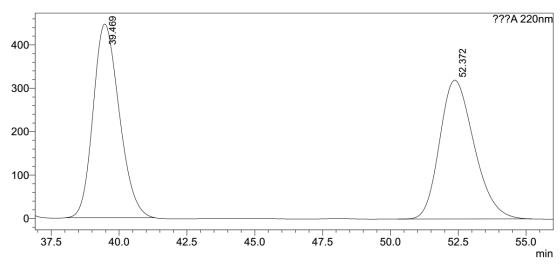
m۷



<Peak Table>

:::/\ <u> </u>	.2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	25.254	6761392	155664	89.775		M	
2	34.677	770085	13820	10.225		M	
Total		7531477	169484				

mV



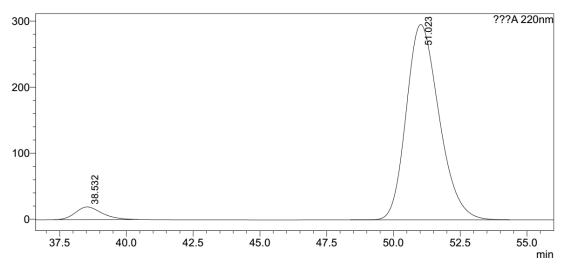
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???A 220nm

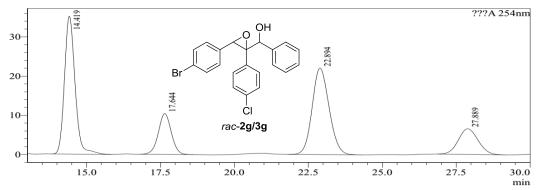
/\ 2							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	39.469	29934141	446051	51.426		M	
2	52.372	28273778	319583	48.574		М	
Total		58207918	765634				

<Chromatogram>

 mV



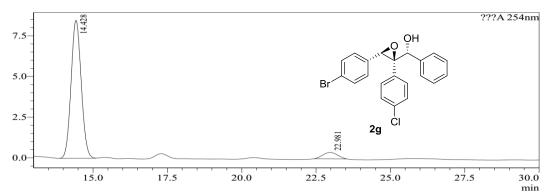
???A2	20nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	38.532	1308350	19095	4.922		M	
2	51.023	25273713	295687	95.078		М	
Total		26582063	314782				



<Peak Table>

???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	14.419	875622	35172	36.836			
2	17.644	317166	10413	13.343			
3	22.894	880474	22055	37.040			
4	27.889	303804	6533	12.781			
Total		2377066	74172				

<Chromatogram>

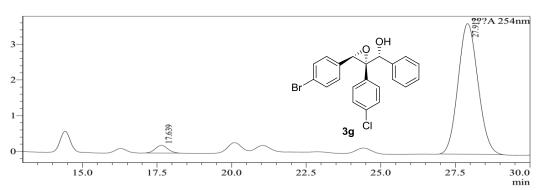


<Peak Table>

???A 2:	???A 254nm									
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name			
1	14.428	201048	8471	93.705						
2	22.981	13506	393	6.295		M				
Total		214554	8863							

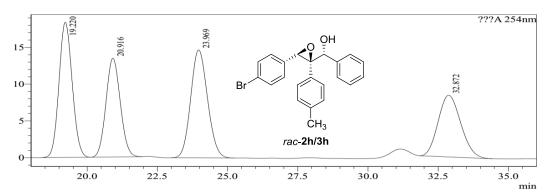
<Chromatogram>

mV



???A 254nm									
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name		
1	17.639	6123	214	3.479		M			
2	27.912	169885	3659	96.521					
Total		176007	3873						

mV

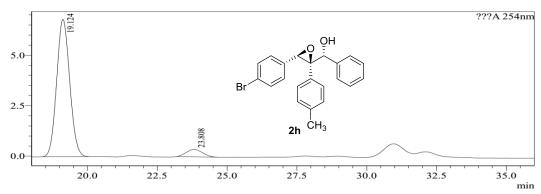


<Peak Table>

???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	19.220	603120	18385	27.886			
2	20.916	476788	13446	22.045			
3	23.969	612659	14686	28.327			
4	32.872	470223	8397	21.742			
Total		2162790	54913				

<Chromatogram>

mV

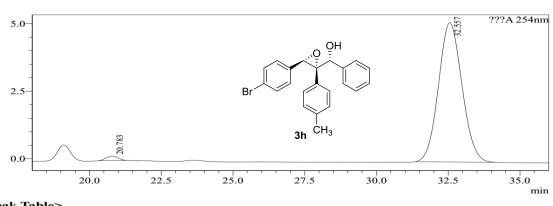


<Peak Table>

???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	19.124	221151	6808	93.788			
2	23.808	14648	364	6.212			
Total		235800	7172				

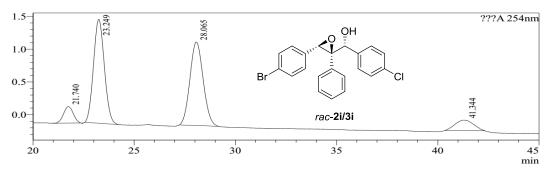
<Chromatogram>

mV



???A 25	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	20.783	4816	163	1.612			
2	32.557	293866	5162	98.388			
Total		298682	5325				

mV

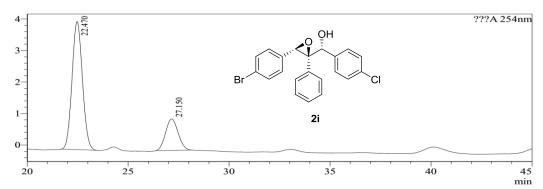


<Peak Table>

???A 2:	54nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	21.740	8387	252	6.165		M	
2	23.249	59322	1586	43.606			
3	28.065	58108	1276	42.713			
4	41.344	10226	162	7.517		M	
Total		136043	3275				

<Chromatogram>

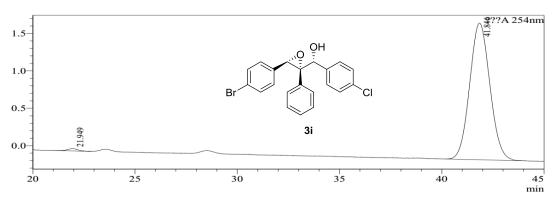
mV



<Peak Table>

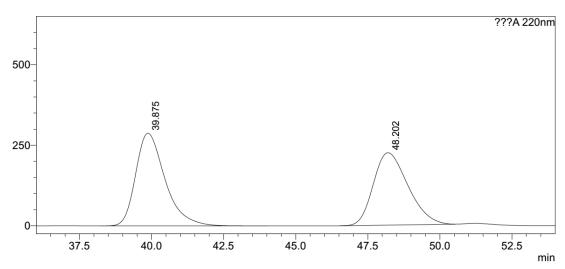
???A	254nm						
Peak	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
	22.470	145827	4068	77.236			
	27.150	42979	1002	22.764			
Tota	ıl	188806	5069				

<Chromatogram>



(((AZ)	94nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	21.949	1113	31	0.879		M	
2	41.846	125601	1827	99.121		M	
Total		126715	1858				

m۷



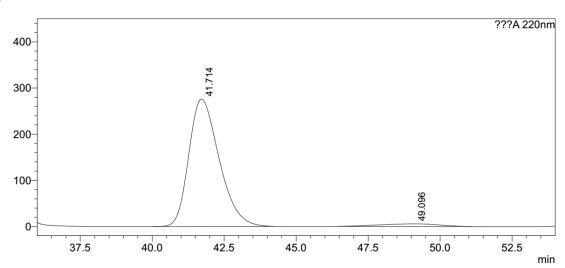
<Peak Table>

???A 220nm

11174	2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	39.875	20404430	287560	52.076			
2	48.202	18777345	224728	47.924			
Total		39181775	512287				

<Chromatogram>

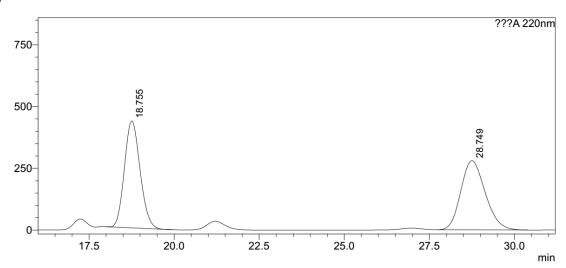
mV



<Peak Table>

Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	41.714	20269361	276178	96.054		М	
2	49.096	832691	5711	3.946		М	
Total		21102052	281889				

m۷



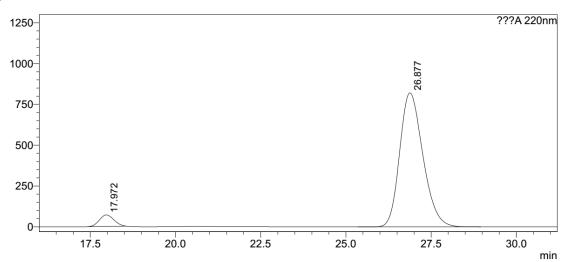
<Peak Table>

???A 220nm

:::/	ZZUIIII						
Peak	# Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	18.755	13835624	432832	49.684		M	
2	28.749	14011471	280440	50.316		М	
Tota	al	27847095	713272				

<Chromatogram>

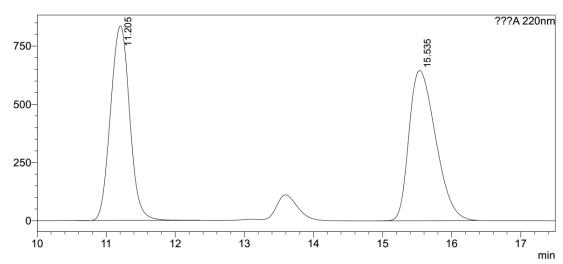
mV



<Peak Table>

:::/\2							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	17.972	2070804	70290	5.063		M	
2	26.877	38830202	820576	94.937		М	
Total		40901006	890865				

mV



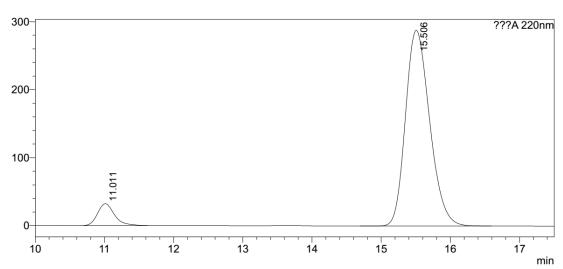
<Peak Table>

???A 220nm

111111111111111111111111111111111111111	22011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	11.205	16114237	835303	48.170		M	
2	15.535	17338726	645002	51.830		M	
Total	l	33452963	1480305				

<Chromatogram>

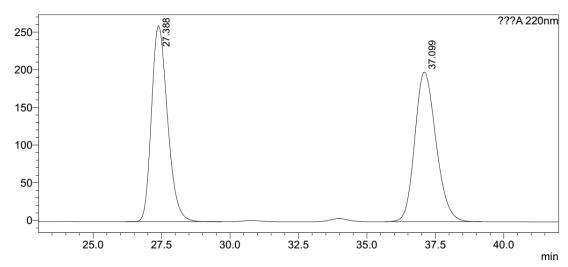
mV



<Peak Table>

:::// 4	2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	11.011	548169	32192	7.270		M	
2	15.506	6992472	287787	92.730		M	
Total		7540640	319979				

mV



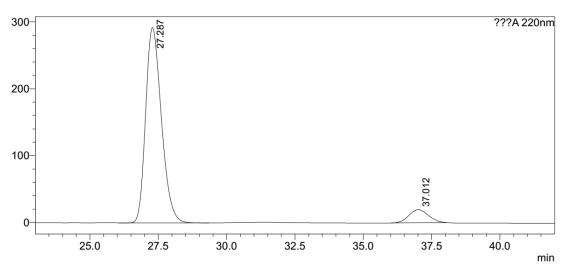
<Peak Table>

222A 220nm

(((A)	201111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	27.388	10616895	260055	49.690		M	
2	37.099	10749440	198544	50.310		M	
Total		21366335	458599				

<Chromatogram>

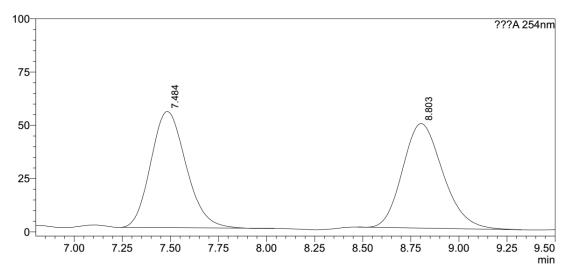
mV



<Peak Table>

-	<u>, \ </u>							
	Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
	1	27.287	11733996	292015	92.267		M	
	2	37.012	983381	19680	7.733		M	
	Total		12717377	311695				

mV



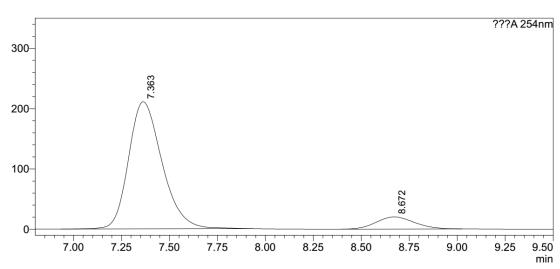
<Peak Table>

???A 254nm

<u>:::/\</u> 2	- J -1 11111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	7.484	697049	54608	49.371		M	
2	8.803	714817	49051	50.629		M	
Total		1411867	103659				

<Chromatogram>

mV

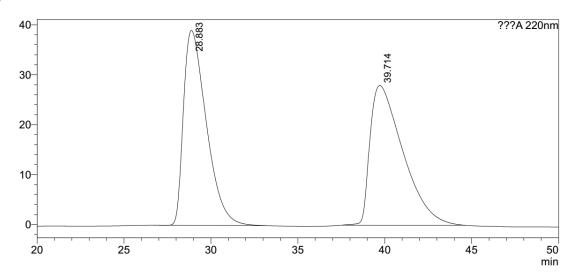


<Peak Table>

???A 254nm

Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	7.363	2570132	210869	90.231		M	
2	8.672	278272	20066	9.769		M	
Total		2848404	230936				

mV



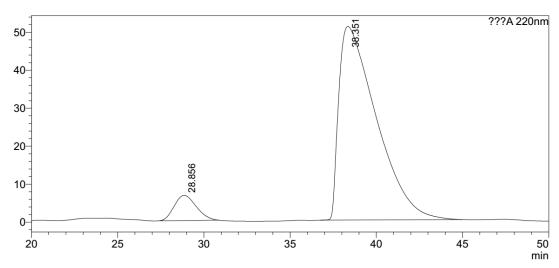
<Peak Table>

???A 220nm

::: : A Z	22011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	28.883	3573822	39014	49.220		M	
2	39.714	3687094	27959	50.780		M	
Total		7260916	66974				

<Chromatogram>

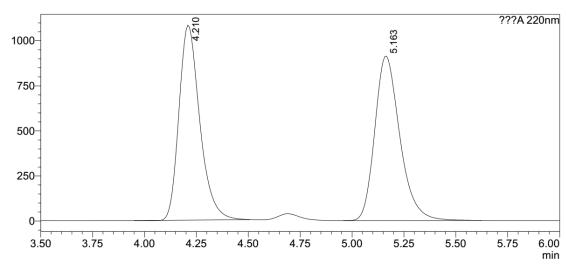
mV



<Peak Table>

:::/\∠							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	28.856	590073	6636	7.100		M	
2	38.351	7720300	50974	92.900		М	
Total		8310373	57610				

m۷



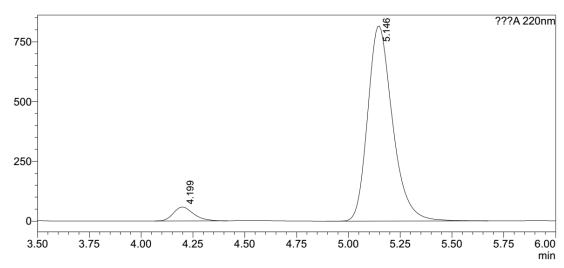
<Peak Table>

???A 220nm

<u> </u>	2201111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	4.210	7538431	1081872	49.575		M	
2	5.163	7667550	912086	50.425		M	
Total		15205981	1993958				

<Chromatogram>

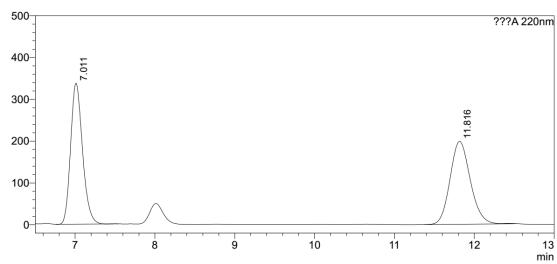
mV



<Peak Table>

:::// 2	.2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	4.199	393238	58406	5.463		M	
2	5.146	6804981	815677	94.537		М	
Total		7198219	874083				

mV



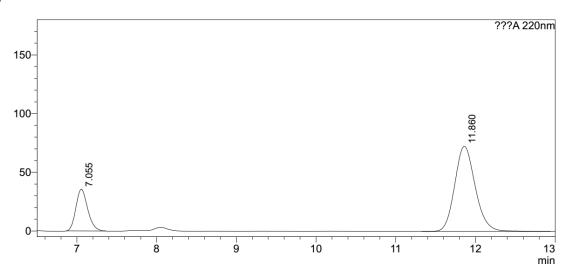
<Peak Table>

???A 220nm

<u>, \ </u>							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	7.011	3491036	337124	49.858		M	
2	11.816	3510986	198381	50.142		M	
Total		7002022	535504				

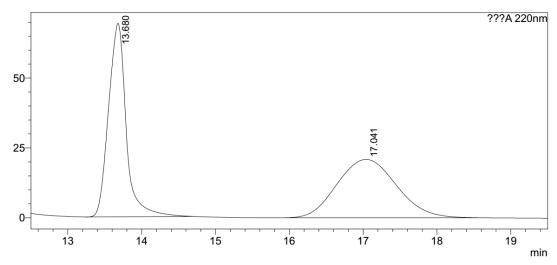
<Chromatogram>

mV



<Peak Table>

P	eak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
	1	7.055	366306	35547	21.965		М	
	2	11.860	1301407	72614	78.035			
	Total		1667713	108161				

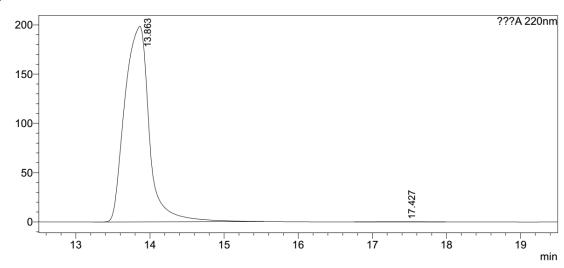


<Peak Table>

???A	220nm						
Peak	# Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	13.680	1151598	69235	49.602		M	
2	1/11/11	1170093	20895	50.398		М	
Tota	al	2321691	90129				

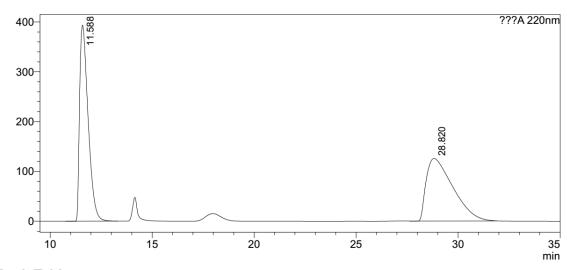
<Chromatogram>

mV



<Peak Table>

:::HZ	201111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	13.863	4561836	198555	99.867		M	
2	17.427	6059	143	0.133		M	
Total		4567894	198699				



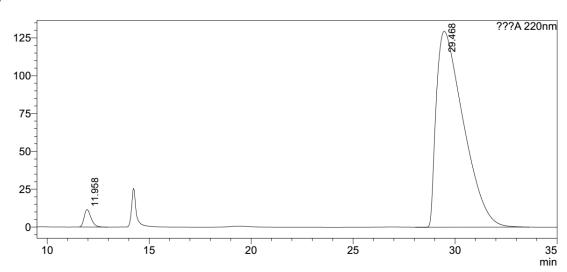
<Peak Table>

???A 220nm

Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	11.588	11464522	393238	50.065		M	
2	28.820	11434849	125601	49.935		М	
Tota	l	22899370	518839				

<Chromatogram>

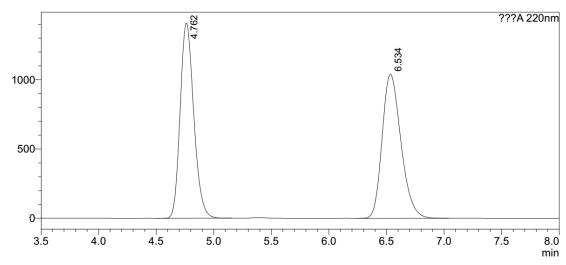
mV



<Peak Table>

	::/\ _	2011111						
F	Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
	1	11.958	273358	11547	2.234		M	
	2	29.468	11960304	129475	97.766		M	
Г	Total		12233663	141022				

mV



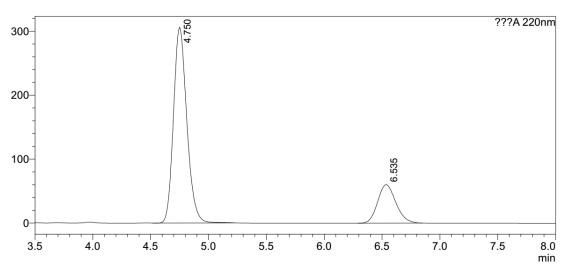
<Peak Table>

???A 220nm

:::// 2							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	4.762	11584056	1408050	49.473		M	
2	6.534	11830890	1040408	50.527		M	
Total		23414946	2448458				

<Chromatogram>

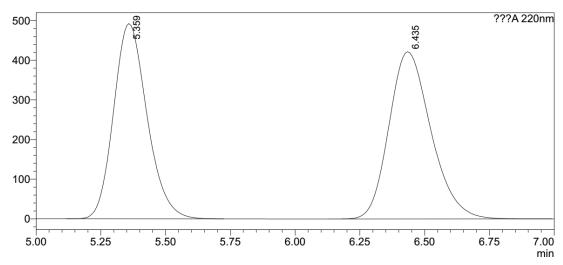
mV



<Peak Table>

((A))	. 22 0nm						
Peak	# Ret. Time	Area	Height	Conc.	Unit	Mark	Name
	1 4.750	2456863	306036	78.847		M	
	2 6.535	659114	60188	21.153		M	
Tot	al	3115977	366224				

mV



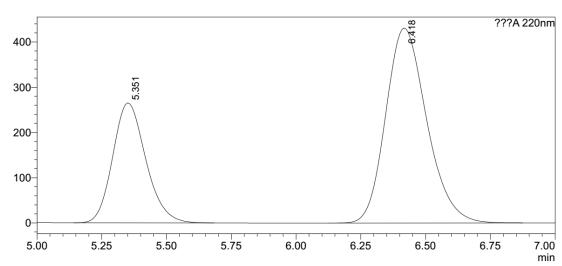
<Peak Table>

???A 220nm

<u>:::/\</u> 2	2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	5.359	4419728	491923	48.160		M	
2	6.435	4757391	421444	51.840		М	
Total		9177119	913367				

<Chromatogram>

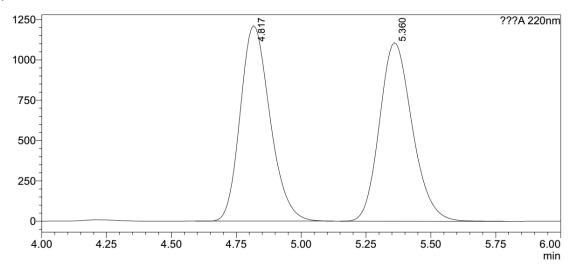
mV



<Peak Table>

???A	220nm						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	5.351	2355898	265133	33.205		M	
2	6.418	4739036	430937	66.795		M	
Tota	I	7094935	696070				

 mV



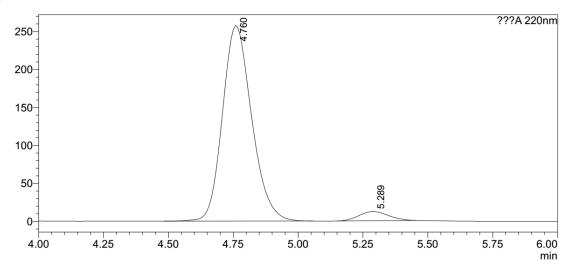
<Peak Table>

???A 220nm

<u> </u>	.201111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	4.817	9673166	1210970	49.857		М	
2	5.360	9728479	1106487	50.143		М	
Total		19401645	2317458				

<Chromatogram>

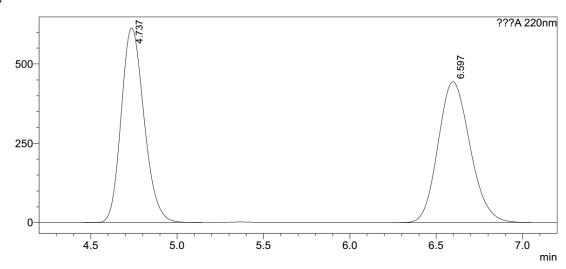
m۱



<Peak Table>

Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	4.760	2019342	257525	95.454		M	
2	5.289	96178	12095	4.546		М	
Total		2115520	269621				

mV



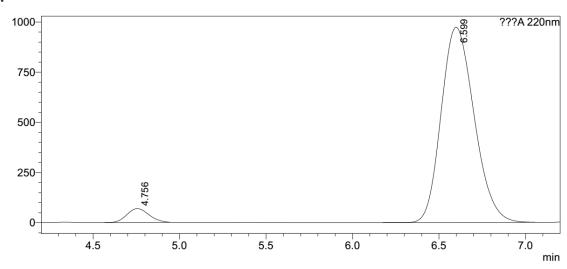
<Peak Table>

???A 220nm

:::/\							
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	4.737	5641559	613379	49.617		M	
2	6.597	5728681	443735	50.383		M	
Total		11370240	1057113				

<Chromatogram>

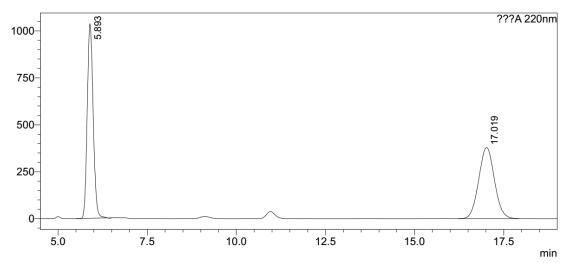
mV



<Peak Table>

:::/\ <u> </u>	.2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	4.756	616656	69211	4.558		М	
2	6.599	12912716	973132	95.442		M	
Total		13529372	1042343				

m۷



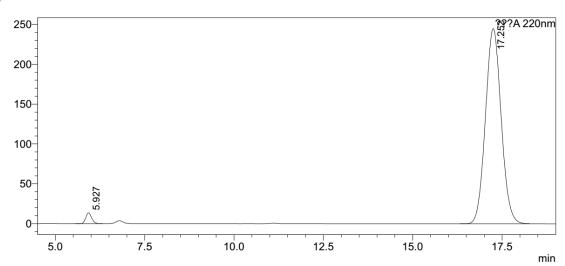
<Peak Table>

222A 220nm

:::/\ 2	2011111						
Peak#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
1	5.893	11954597	1035126	50.712		M	
2	17.019	11618916	379086	49.288		М	
Total		23573513	1414212				

<Chromatogram>

mV



<Peak Table>

: : // ZZOIIII								
Pea	ık#	Ret. Time	Area	Height	Conc.	Unit	Mark	Name
	1	5.927	154224	13520	2.014		M	
	2	17.252	7503371	245188	97.986		M	
To	otal		7657594	258707				