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Supporting Information

Temperature-dependent synthesis of vinyl sulfones and β-hydroxy sulfones from *t*-butylsulfinamide and alkenes under aerobic conditions

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1. General information

All the chemicals were obtained from Tianjin Kermel Chemical Reagent Co., Ltd. and used as received. All reactions were monitored by TLC and Silica gel was purchased from Qing Dao Hai Yang Chemical Industry Co. ¹H, ¹³C NMR spectra were recorded on a Bruker Avance 400 MHz spectrometer operating at 400.13, 100.61 MHz, respectively. NMR spectra were recorded in CDCl₃ at room temperature (20 ± 2 °C). High resolution mass spectra (HRMS) of the products were obtained on a Bruker Daltonics micro TOF-QII spectrometer.

2. Experimental procedures for the synthesis of (E)-(2-(tert-butylsulfonyl)vinyl)benzene



Styrene 0.5 mmol (0.051 g), tert-butyl sulfinamide 1.0 mmol (0.121 g), phosphorous acid 1.0 mmol (0.082 g), copper sulfate pentahydrate 0.1 mmol (0.25 g) and trifluoroacetic acid 0.5 mmol (0.057 g) were mixed in a 25 mL round-bottomed flask and refluxed at 120 °C for 5 h without solvent. The mixture was quenched with water (10 mL), extracted with CH₂Cl₂ (3×15 mL). The combined organic layers were washed with brine (15 mL) and dried over anhydrous Na₂SO₄. After filtration, the solvent was evaporated in vacuo. The crude product was purified by silica gel chromatography (petroleum ether : ethyl acetate = 20 : 1, v : v) to give the desired products.

3. Experimental procedures for the synthesis of 2-(tert-butylsulfonyl)-1-phenylethan-1-ol

Styrene 0.5 mmol (0.051 g), tert-butyl sulfinamide 1.0 mmol (0.121 g), phosphorous acid 1.0 mmol (0.082 g), copper sulfate pentahydrate 0.1 mmol (0.25 g) and trifluoroacetic acid 0.5 mmol (0.057 g) were mixed in a 25 mL round-bottomed flask and stirred at 40 °C for 12 h without solvent. The mixture was diluted with water (10 mL), extracted with CH_2Cl_2 (3 × 15 mL). The combined organic layers were washed with brine (15 mL) and dried over anhydrous Na₂SO₄. After filtration, the solvent was evaporated in vacuo. The crude product was purified by silica gel chromatography (petroleum ether : ethyl acetate = 10 : 1, v : v) to give the desired products.

4. ¹⁸O labeling experiment



Chemical Formula: C₁₂H₁₆O¹⁸OS Exact Mass: 227.0986 [M+H]⁺



5. Characterization data for products (3a-3r)

(E)-(2-(tert-butylsulfonyl)vinyl)benzene (3a).



Yield 76%. White solid. M.p. 98-100 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm) 7.59 (d, J = 15.5 Hz, 1H, 5-H), 7.54-7.52 (m, 2H, 2-H), 7.44-7.42 (m, 3H, 3-H, 4-H), 6.84 (d, J = 15.5 Hz, 1H, 6-H), 1.42 (s, 9H, 8-H). ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 146.4 (5-C), 132.5 (1-C), 131.3 (4-C), 129.1 (3-C), 128.6 (2-C), 120.7 (6-C), 58.9 (7-C), 23.4 (8-C). HRMS: C₁₂H₁₆O₂S [M + H]⁺ 225.0949, found 225.0944.

(E)-1-bromo-4-(2-(tert-butylsulfonyl)vinyl)benzene (3b).



Yield 64%. White solid. M.p. 86-89 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.56 (d, *J* = 8.4 Hz, 2H, 3-H), 7.52

 $(d, J = 15.6 \text{ Hz}, 1\text{H}, 5\text{-H}), 7.40 (d, J = 8.4 \text{ Hz}, 2\text{H}, 2\text{-H}), 6.84 (d, J = 15.4 \text{ Hz}, 1\text{H}, 6\text{-H}), 1.42 (s, 9\text{H}, 8\text{-H}), {}^{13}\text{C} \text{ NMR}$ (CDCl₃, 100 MHz): δ (ppm), 145.0 (5-C), 132.4 (3-C), 131.4 (1-C), 129.9 (2-C), 125.7 (4-C), 121.5 (6-C), 59.0 (7-C), 23.4 (8-C); HRMS: C1₂H₁₅BrO₂S [M + H]⁺ 303.0054, found 303.0051.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-4-chlorobenzene (3c).

$$5$$
 0 0 5 7 8 1 6 7 8 1 6 7 8

Yield 63%. White solid. M.p. 114-116 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.54 (d, J = 15.6 Hz, 1H, 5-H), 7.48-7.39 (m, 4H, 2-H, 3-H), 6.81 (d, J = 15.6 Hz, 1H, 6-H), 1.42 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 145.0 (5-C), 137.3 (4-C), 131.0 (1-C), 129.7 (2-C), 129.5 (3-C), 121.3 (6-C), 58.4 (7-C), 23.4 (8-C); HRMS: C₁₂H₁₅ClO₂S [M + H]⁺ 259.0560, found 259.0577.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-4-fluorobenzene (3d).



Yield 58%. White solid. M.p. 87-89 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.57-7.53 (m, 3H), 7.12 (m, 2H), 6.78 (d, J = 15.6 Hz, 1H), 1.42 (s, 9H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 164.4 (d, J = 251.4 Hz, 4-C), 145.1 (5-C), 130.6 (d, J = 8.7 Hz, 2-C), 128.8 (d, J = 3.2 Hz, 1-C), 120.4 (6-C), 116.3 (d, J = 21.9 Hz, 3-C), 58.9 (7-C), 23.4 (8-C); HRMS: C₁₂H₁₅FO₂S [M + H]⁺ 243.0855, found 243.0851.

(E)-1-bromo-3-(2-(tert-butylsulfonyl)vinyl)benzene (3e).



Yield 58%. White solid. M.p. 127-128 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.70 (m, 1H, 2-H), 7.61-7.58 (m, 1H, 4-H), 7.54 (d, *J* = 15.6 Hz, 1H, 5-H), 7.47 (d, *J* = 7.8 Hz, 1H, 10-H), 7.33 (m, 1H, 9-H), 6.86 (d, *J* = 15.6 Hz, 1H, 6-H), 1.42 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 144.7 (5-C), 134.5 (1-C), 134.1 (3-C), 131.1 (4-C), 130.7 (2-C), 127.2 (9-C), 123.2 (10-C), 122.3 (6-C), 59.0 (7-C), 23.4 (8-C); HRMS: C₁₂H₁₅BrO₂S [M + Na]⁺ 324.9874, found 324.9861.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-3-chlorobenzene (3f).



Yield 60%. White solid. M.p. 115-117 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.57-7.54 (m, 2H, 5-H, 2-H), 7.46-7.37 (m, 3H, 4-H, 9-H, 10-H), 6.87 (d, *J* = 15.6 Hz, 1H, 6-H), 1.45 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 144.8 (5-C), 135.2 (1-C), 134.2 (3-C), 131.1 (9-H), 130.4 (4-C), 128.1 (2-C), 126.8 (10-C), 122.3 (6-C), 59.0 (7-C), 23.4 (8-C); HRMS: C₁₂H₁₅ClO₂S [M + H]⁺ 259.0560, found 259.0556.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-3-fluorobenzene (3g).



Yield 59%. White solid. M.p. 85-89 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.58 (d, *J* = 15.6 Hz, 1H, 5-H), 7.46-7.40 (m, 1H, 9-H), 7.33 (d, *J* = 7.6 Hz, 1H, 10-H), 7.25 (m, 1H, 4-H), 7.18 (m, 1H, 2-H), 6.86 (d, *J* = 15.6 Hz, 1H, 6-H), 1.45 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 164.2 (d, *J* = 246.4 Hz, 3-C), 145.0 (d, *J* = 2.6 Hz, 5-C), 134.6 (d, *J* = 7.6 Hz, 1-C), 130.8 (d, *J* = 8.2 Hz, 9-C), 124.6 (d, *J* = 2.8 Hz, 10-C), 122.2 (6-C), 118.2 (d, *J* = 21.2 Hz, 4-C), 114.8 (d, *J* = 22 Hz, 2-C), 59.0 (7-C), 23.4 (8-C); HRMS: C₁₂H₁₅FO₂S [M + H]⁺ 243.0855, found 243.0851.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-4-methylbenzene (3h).



Yield 79%. White solid. M.p. 127-130 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.55 (d, J = 15.4 Hz, 1H, 5-H), 7.42 (d, J = 8.0 Hz, 2H), 7.23 (d, J = 8.0 Hz, 2H), 6.76 (d, J = 15.6 Hz, 1H), 2.39 (s, 3H), 1.42 (s, 9H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 146.5 (5-C), 141.9 (4-C), 129.8 (3-C), 129.8 (1-C), 128.6 (2-C), 119.4 (6-C), 58.9 (7-C), 23.4 (8-C), 21.5 (9-C); HRMS: Cl₃H₁₈O₂S [M + H]⁺ 239.1106, found 239.1104.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-3-methylbenzene (3i).



Yield 75%. White solid. M.p. 68-72 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.56 (d, J = 15.6 Hz, 1H, 5-H), 7.34-7.26 (m, 4H, 2-H, 4-H, 9-H, 10-H), 6.80 (d, J = 15.6 Hz, 1H, 6-H), 2.39 (s, 3H, 11-H), 1.42 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 146.6 (5-C), 138.9 (3-C), 132.5 (1-C), 132.1 (9-C), 129.1 (4-C), 129.0 (2-C), 125.8 (10-C), 120.4 (6-C), 58.9 (7-C), 23.4 (8-C), 21.3 (11-C); HRMS: C1₃H₁₈O₂S [M + H]⁺ 239.1106, found 239.1105.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-2-methylbenzene (3j).



Yield 70%. White solid. M.p. 108-109 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.86 (d, J = 15.4 Hz, 1H, 5-H), 7.51 (m, 1H, 10-H), 7.34 (m, 1H, 9-H), 7.25, (m, 2H, 4-H, 3-H), 6.73 (d, J = 15.4 Hz, 1H, 6-H), 2.44 (s, 3H, 11-H), 1.43 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 144.5 (5-C), 138.0 (1-C), 131.7 (2-C), 131.1 (4-C), 131.0 (3-C), 126.9 (10-C), 126.6 (9-C), 121.7 (6-C) 58.9 (7-C), 23.4 (8-C), 21.3 (11-C); HRMS: C₁₃H₁₈O₂S [M + H]⁺ 239.1106, found 239.1112.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-4-(chloromethyl)benzene (3k).



Yield 66%. White solid. M.p. 124-126 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.59 (d, J = 15.6 Hz, 1H,), 7.55 (d, J = 8.2 Hz, 2H, 2-H), 7.47 (d, J = 8.4 Hz, 2H), 6.85 (d, J = 15.6 Hz, 1H, 6-H), 1.42 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 145.6 (5-C), 140.6 (1-C), 129.3 (2-C), 128.9 (3-C), 121.3 (6-C), 58.5 (7-C), 23.4 (8-C), 18.4 (11-C); HRMS: C₁₃H₁₇ClO₂S [M + H]⁺ 273.0716, found 273.0711.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-4-methoxybenzene (3l).



Yield 77%. White solid. M.p. 139-141 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.55-7.48 (m, 3H, 2-H, 5-H), 6.94 (d, *J* = 8.6 Hz, 2H, 3-H), 6.68 (d, *J* = 15.4 Hz, 1H, 6-H), 3.86 (s, 3H, 9-H), 1.42 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 162.1 (4-C), 146.1 (5-C), 130.3 (2-C), 125.2 (1-C), 117.7 (6-C), 114.5 (3-C), 58.9 (7-C), 55.5 (9-C), 23.5 (8-C); HRMS: C₁₃H₁₈O₃S [M + H]⁺ 255.1055, found 255.1055.

(E)-1-(tert-butyl)-4-(2-(tert-butylsulfonyl)vinyl)benzene (3m)



Yield 78%. White solid. M.p. 134-135 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.58 (d, J = 15.6 Hz, 1H, 5-H), 7.48 (m, 4H, 2-H, 3-H), 6.80 (d, J = 15.6 Hz, 1H, 6-H), 1.43 (s, 9H, 8-H),1.35 (s, 9H, 9-H) ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 155.0 (4-H), 146.4 (5-H), 129.8 (1-H), 128.4 (2-H), 126.3 (3-H), 119.6 (6-H), 58.9 (7-C), 35.0 (9-C), 31.1 (10-C), 23.4 (8-C); HRMS: C₁₆H₂₄O₂S [M + H]⁺ 281.1575, found 281.1582.

(E)-2-(2-(tert-butylsulfonyl)vinyl)naphthalene (3n).



Yield 77%. White solid. M.p. 128-129 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.99 (s, 1H, 1-H), 7.90-7.87 (m, 3H, 8-H, 5-H, 4-H), 7.77 (d, J = 17.8 Hz, 1H, 11-H), 7.65 (dd, J = 1.6 Hz, J = 8.6 Hz, 1H, 3-H), 7.60-7.54 (m, 2H, 6-H, 7-H), 6.95 (d, J = 15.4 Hz, 1H, 12-H), 1.48 (s, 9H, 14-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 146.5 (5-C), 134.6 (13-C), 133.2 (1-C), 130.9 (3-C), 130.0 (4-C), 129.0 (12-C), 128.7 (14-C), 127.9 (10-C, 11-C), 127.1 (9-C), 123.4 (2-C), 120.7 (6-C), 59.0 (7-C), 23.5 (8-C); HRMS: C₁₆H₁₈O₂S [M + H]⁺ 275.1106, found 275.1106.

(E)-(1-(tert-butylsulfonyl)prop-1-en-2-yl)benzene (30).



Yield 68%. White solid. M.p. 131-132 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.46-7.40 (m, 5H, 2-H, 3-H, 4-H), 6.38 (s, 1H, 6-H), 2.58 (s, 3H, 9-H), 1.44 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 156.8 (5-C), 141.0 (1-C), 129.8 (3-C), 128.8(4-C), 126.3 (2-C), 120.4 (6-C), 58.5 (7-C), 23.2 (8-C), 14.1 (9-C); HRMS: C₁₃H₁₈O₂S [M + H]⁺ 239.1106, found 239.1101.

(E)-1-(1-(tert-butylsulfonyl)prop-1-en-2-yl)-4-methylbenzene (3p).



Yield 74%. White solid. M.p. 129-131 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.37 (d, J = 8.2 Hz, 2H, 3-H), 7.23 (d, J = 8.2 Hz, 2H, 2-H), 6.39 (d, J = 1.2 Hz, 1H, 6-H), 2.58 (d, J = 1.2 Hz, 3H, 12-H), 2.40 (s, 3H, 11-H) 1.45 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 156.6 (5-C), 140.1 (4-C), 137.9 (1-C), 129.5 (3-C), 126.3 (2-C), 119.5 (6-C), 60.0 (7-C), 23.2 (8-C), 21.2 (11-C), 17.1 (12-C); HRMS: C₁₄H₂₀O₂S [M + H]⁺ 253.1262, found 253.1259.

(2-(tert-butylsulfonyl)ethene-1,1-diyl)dibenzene (3q).



Yield 68%. White solid. M.p. 112-114 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.43-7.28 (m, 10H, 2-C, 3-C, 4-C), 6.77 (s, 1H, 6-C), 1.45 (s, 9H, 8-C), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 157.6 (5-C), 140.3 (9-C), 135.9 (1-C), 130.2 (12-C), 129.7 (11-C), 128.9 (4-C), 128.7 (3-C), 128.3 (10-C), 127.5 (2-C), 119.6 (6-C), 59.6 (7-C), 23.6 (8-C); HRMS: C₁₈H₂₀O₂S [M + Na]⁺ 323.1082, found 323.1084.

(E)-1-(2-(tert-butylsulfonyl)vinyl)-4-methoxybenzene (3r).



Yield 36%. White solid. M.p. 126-127 °C. ¹H NMR (CDCl₃, 400 MHz): δ (ppm), 7.50 (s, 1H, 5-H), 7.44 (d, J = 8.6 Hz, 2H, 2-H), 6.96 (d, J = 8.6 Hz, 2H, 3-H), 3.86 (s, 3H, 10-H), 2.36 (d, J = 1.2 Hz, 3H, 9-H), 1.41 (s, 9H, 8-H), ¹³C NMR (CDCl₃, 100 MHz): δ (ppm), 160.5 (4-C), 142.0 (5-C), 131.7 (2-C), 114.1 (3-C), 60.6 (10-C), 55.4 (7-C), 24.0 (8-C), 16.3 (9-C); HRMS: C₁₄H₂₀O₃S [M + H]⁺ 291.1031, found 291.1031.

6. Characterization data for products (4a-4k)

2-(tert-butylsulfonyl)-1-phenylethan-1-ol (4a)



Yield 69%. White solid. M.p. 102-103 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.44 (s, 9H, 8-H), 3.12 (dd, J = 1.2 Hz, J = 13.2 Hz, 1H, 6-H), 3.33 (dd, J = 10.4 Hz, J = 13.6 Hz, 1H, 6-H), 3.92 (d, J = 1.2 Hz, 1H, -OH), 5.54 (d, J = 10.0 Hz, 1H, 5-H), 7.34 (m, 1H, 4-H), 7.37-7.45 (m, 4H, 2, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.2 (8-C), 54.1 (6-C), 59.8 (7-C), 67.6 (6-C), 125.7 (2-C), 128.3 (4-C), 128.8 (3-C), 140.0 (1-C). HRMS: C₁₂H₁₈O₃S [M + Na]⁺: m/z 265.0869, Found 265.0869.

1-(4-bromophenyl)-2-(*tert*-butylsulfonyl)ethan-1-ol (4b)



Yield 73%. White solid. M.p. 133-135 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.44 (s, 9H, 8-H), 3.07 (dd, J = 1.2 Hz, J = 9.2 Hz, 1H, 6-H), 3.26 (dd, J = 10.0 Hz, J = 13.2 Hz, 1H, 6-H), 3.97 (s, 1H, -OH), 5.50 (d, J = 10.0 Hz, 1H, 5-H), 7.31 (d, J = 8.4 Hz, 2H, 2-H), 7.53 (d, J = 8.4 Hz, 2H, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.1 (8-C), 53.8 (6-C), 60.0 (7-C), 67.0 (5-C), 122.2 (4-C), 127.5 (2-C), 131.2 (3-C), 140.1 (1-C). HRMS: C₁₂H₁₇BrO₃S [M + Na]⁺: m/z 342.9974, Found 342.9956.

2-(tert-butylsulfonyl)-1-(4-chlorophenyl)ethan-1-ol (4c)



Yield 63%. White solid. M.p. 111-114 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.43 (s, 9H, 8-H), 3.07 (dd, J = 1.6 Hz, J = 13.6 Hz, 1H, 6-H), 3.29 (dd, J = 10.4 Hz, J = 13.6 Hz, 1H, 6-H), 4.01 (d, J = 2.0 Hz, 1H, -OH), 5.48 (d, J = 10.0 Hz, 1H, 5-H), 7.33-7.38 (m, 4H, 2, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.1 (8-C), 53.9 (6-C), 59.9 (7-C), 67.0 (5-C), 127.2 (2-C), 128.9 (3-C), 134.0 (4-C), 149.7 (1-C). HRMS: C₁₂H₁₇ClO₃S [M + Na]⁺: m/z 299.0479, Found 342.9959.

2-(tert-butylsulfonyl)-1-(4-fluorophenyl)ethan-1-ol (4d)



Yield 70%. White solid. M.p. 102-105 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.44 (s, 9H, 8-H), 3.08 (dd, J = 1.2 Hz, J = 9.2 Hz, 1H, 6-H), 3.29 (dd, J = 10.0 Hz, J = 13.2 Hz, 1H, 6-H), 3.98 (s, 1H, -OH), 5.51 (d, J = 9.6 Hz, 1H, 5-H), 7.07 (m, 2H, 2-H), 7.41 (m, 2H, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.1 (8-C), 54.0 (6-C), 59.9 (7-C), 67.0 (5-C), 115.7 (d, J = 31.7 Hz, 3-C), 127.5 (d, J = 9.4 Hz, 2-C), 136.9 (d, J = 3.7 Hz, 1-C), 161.7 (d, J = 251.8 Hz, 4-C). HRMS: C1₂H₁₇FO₃S [M + Na]⁺: m/z 283.0775, Found 283.0776.

1-(3-bromophenyl)-2-(tert-butylsulfonyl)ethan-1-ol (4e)



Yield 65%. White solid. M.p. 127-129 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.45 (s, 9H, 10-H), 3.09 (dd, J = 0.8 Hz, J = 13.6 Hz, 1H, 8-H), 3.26 (dd, J = 10.4 Hz, J = 13.6 Hz, 1H, 8-H), 4.02 (d, 1H, J = 2.0 Hz, -OH), 5.49 (d, J = 10.4 Hz, 1H, 7-H), 7.27 (d, J = 8.0 Hz, 1H, 4-H), 7.35 (d, J = 7.6 Hz, 1H, 6-H), 7.46 (d, J = 8.0 Hz, 1H, 2-H), 7.62 (s, 1H, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.1 (10-C), 53.9 (8-C), 59.9 (9-C), 66.9 (7-C), 122.9 (2-C), 124.4 (3-C), 128.9 (5-C), 130.4 (4-C), 131.4 (6-C), 143.3 (1-C). HRMS: C₁₂H₁₇BrO₃S [M + Na]⁺: m/z 342.9974, Found 342.9956.

2-(tert-butylsulfonyl)-1-(3-chlorophenyl)ethan-1-ol (4f)



Yield 63%. White solid. M.p. 103-105 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.43 (s, 9H, 10-H), 3.10 (dd, J = 1.2 Hz, J = 13.6 Hz, 1H, 8-H), 3.26 (dd, J = 10.4 Hz, J = 13.6 Hz, 1H, 8-H), 4.05 (d, 1H, J = 2.0 Hz, -OH), 5.48 (d, J = 10.4 Hz, 1H, 7-H), 7.26-7.31 (m, 3H, 2, 3, 4-H), 7.45 (s, 1H, 6-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.1 (10-C), 52.8 (8-C), 59.9 (9-C), 67.0 (7-C), 124.0 (2-C), 126.0 (3-C), 128.4 (5-C), 130.1 (4-C), 137.7 (6-C), 143.2 (1-C). HRMS: Cl₂H₁₇ClO₃S [M + Na]⁺: m/z 299.0479, Found 299.0454.

2-(tert-butylsulfonyl)-1-(3-fluorophenyl)ethan-1-ol (4g)



Yield 67%. White solid. M.p. 95-98 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.42 (s, 9H, 10-H), 3.11 (dd, J = 1.6 Hz, J = 13.6 Hz, 1H, 8-H), 3.31 (dd, J = 10.0 Hz, J = 13.2 Hz, 1H, 8-H), 4.06 (d, 1H, J = 2.0 Hz, -OH), 5.49 (d, J = 9.6 Hz, 1H, 7-H), 6.97-7.02 (m, 1H, 6-H), 7.17 (d, J = 7.6 Hz, 2H, 2, 4-H), 7.31 (t, 1H, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.1 (10-C), 53.8 (8-C), 59.9 (9-C), 67.0 (7-C), 112.8 (d, J = 22.1 Hz, 6-C), 115.1 (d, J = 20.9 Hz, 4-C), 121.4 (d, J = 9.2 Hz, 3-C), 130.4 (d, J = 8.2 Hz, 2-C), 143.8 (d, J = 7.0 Hz, 1-C), 164.1 (d, J = 24.5 Hz, 5-C). HRMS: C1₂H₁₇FO₃S [M + Na]⁺: m/z 283.0775, Found 283.0773.

2-(tert-butylsulfonyl)-1-(m-tolyl)ethan-1-ol (4h)



Yield 65%. White solid. M.p. 89-90 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.44 (s, 9H, 10-H), 2.37 (s, 3H, 7-H), 3.11 (dd, *J* = 1.2 Hz, *J* = 13.6 Hz, 1H, 9-H), 3.31 (dd, *J* = 10.4 Hz, *J* = 13.6 Hz, 1H, 9-H), 3.90 (d, *J* = 1.2 Hz, 1H, -OH), 5.50 (d, *J* = 10.4 Hz, 1H, 8-H), 7.14 (d, *J* = 7.6 Hz, 2H, 4-H), 7.21 (d, *J* = 7.6 Hz, 2H, 2-H), 7.25-7.30 (m, 2H, 6, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 21.4 (7-C), 23.2 (11-C), 54.1 (9-C), 59.8 (10-C), 67.6 (8-C),

122.8 (2-C), 126.4 (4-C), 128.7 (3-C), 129.1 (6-C), 138.6 (5-C), 141.0 (1-C). HRMS: $C_{13}H_{20}O_{3}S [M + Na]^+$: m/z 279.1025, Found 279.1209.

2-(tert-butylsulfonyl)-1-(o-tolyl)ethan-1-ol (4i)



Yield 57%. White solid. M.p. 98-100 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.44 (s, 9H, 11-H), 2.36 (s, 3H, 7-H), 3.04 (d, J = 13.6 Hz, 1H, 9-H), 3.26 (dd, J = 10.0 Hz, J = 13.2 Hz, 1H, 9-H), 3.83 (s, 1H, -OH), 5.74 (d, J = 9.6 Hz, 1H, 8-H), 7.17 (d, J = 7.2 Hz, 1H, 4-H), 7.21-7.30 (m, 2H, 2, 3-H), 7.60 (d, J = 7.2 Hz, 1H, 5-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 18.9 (7-C), 23.2 (11-C), 52.9 (9-C), 59.7 (10-C), 64.1 (8-C), 125.4 (2-C), 126.6 (3-C), 128.1 (5-C), 130.7 (4-C), 133.9 (6-C), 139.2 (1-C). HRMS: C₁₃H₂₀O₃S [M + Na]⁺: m/z 279.1025, Found 279.1026.

1-(4-(tert-butyl)phenyl)-2-(tert-butylsulfonyl)ethan-1-ol (4j)



Yield 57%. White solid. M.p. 100-103 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.31 (s, 9H, 10-H), 1.42 (s, 9H, 8-H), 3.12 (dd, J = 1.2 Hz, J = 13.2 Hz, 1H, 6-H), 3.34 (dd, J = 10.4 Hz, J = 13.6 Hz, 1H, 6-H), 3.90 (d, J = 1.2 Hz, 1H, -OH), 5.49 (d, J = 10.4 Hz, 1H, 5-H), 7.35 (d, J = 8.4 Hz, 2H, 2-H), 7.40 (d, J = 8.4 Hz, 2H, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.2 (8-C), 31.3 (10-C), 34.6 (9-C), 54.0 (6-C), 59.8 (7-C), 67.4 (5-C), 125.4 (3-C), 125.7 (2-C), 138.1 (1-C), 151.4 (4-C). HRMS: C₁₆H₂₆O₃S [M + Na]⁺: m/z 321.1495, Found 321.1497.

2-(tert-butylsulfonyl)-1-(4-(chloromethyl)phenyl)ethan-1-ol (4k)



White solid. Yield 59%. M.p. 101-103 °C. ¹H NMR (400 MHz, CDCl₃): δ (ppm), 1.44 (s, 9H, 9-H), 3.10 (dd, J = 1.2 Hz, J = 13.2 Hz, 1H, 7-H), 3.30 (dd, J = 10.4 Hz, J = 13.6 Hz, 1H, 7-H), 3.96 (d, J = 1.6 Hz, 1H, -OH), 4.59 (s, 2H, 5-H), 5.53 (d, J = 10.4 Hz, 1H, 6-H), 7.40-7.45 (m, 4H, 2, 3-H). ¹³C NMR (100 MHz, CDCl₃): δ (ppm), 23.2 (9-C), 45.8 (5-C), 54.0 (7-C), 59.9 (8-C), 67.3 (6-C), 126.2 (2-C), 129.1 (3-C), 137.6 (4-C), 141.4 (1-C). HRMS: C₁₃H₁₉ClO₃S [M + Na]⁺: m/z 313.0636, Found 313.0636.

7. ¹H NMR, ¹³C NMR copies of products (3a-3r)



Fig.1 ¹H NMR spectrum of compound 3a



Fig.2 ¹³C NMR spectrum of compound 3a



Fig.3 ¹H NMR spectrum of compound 3b



Fig.4 ¹³C NMR spectrum of compound 3b







Fig.6 ¹³C NMR spectrum of compound 3c







Fig.8 ¹³C NMR spectrum of compound 3d



Fig.9 ¹H NMR spectrum of compound **3e**



Fig.10 ¹³C NMR spectrum of compound 3e



Fig.11 ¹H NMR spectrum of compound 3f



Fig.12 ¹³C NMR spectrum of compound 3f



Fig.13 ¹H NMR spectrum of compound 3g



Fig.14 ¹³C NMR spectrum of compound 3g







Fig.16¹³C NMR spectrum of compound 3h







Fig.18 ¹³C NMR spectrum of compound 3i





Fig.20 ¹³C NMR spectrum of compound 3j



Fig.21 ¹H NMR spectrum of compound 3k



Fig.22 ¹³C NMR spectrum of compound 3k







Fig.24 ¹³C NMR spectrum of compound 31







Fig.26 ¹³C NMR spectrum of compound 3m







Fig.28 ¹³C NMR spectrum of compound 3n







Fig.30 ¹³C NMR spectrum of compound 30







Fig.32 ¹³C NMR spectrum of compound 3p



Fig.33 ¹H NMR spectrum of compound 3q



Fig.34 ¹³C NMR spectrum of compound 3q



Fig.35 ¹H NMR spectrum of compound 3r



Fig.36 ¹³C NMR spectrum of compound 3r





Fig.37 ¹H NMR spectrum of compound 4a



Fig.38 ¹³C NMR spectrum of compound 4a



Fig.39 ¹H NMR spectrum of compound 4b



Fig.40 ¹³C NMR spectrum of compound 4b



Fig.41 ¹H NMR spectrum of compound **4**c



Fig.42 ¹³C NMR spectrum of compound 4c



Fig.43 ¹H NMR spectrum of compound 4d



Fig.44 ¹³C NMR spectrum of compound 4d



Fig.45 ¹H NMR spectrum of compound 4e



Fig.46 ¹³C NMR spectrum of compound 4e



Fig.47 ¹H NMR spectrum of compound 4f



Fig.48 ¹³C NMR spectrum of compound 4f



Fig.49 ¹H NMR spectrum of compound **4g**



Fig.50 ¹³C NMR spectrum of compound 4g



Fig.51 ¹H NMR spectrum of compound 4h



Fig.52 ¹³C NMR spectrum of compound 4h







Fig.54 ¹³C NMR spectrum of compound 4i



Fig.55 ¹H NMR spectrum of compound 4j



Fig.56 ¹³C NMR spectrum of compound 4j



Fig.57 ¹H NMR spectrum of compound 4k



Fig.58 ¹³C NMR spectrum of compound 4k