

Supporting Information

The *N*-silylation of sulfoximines

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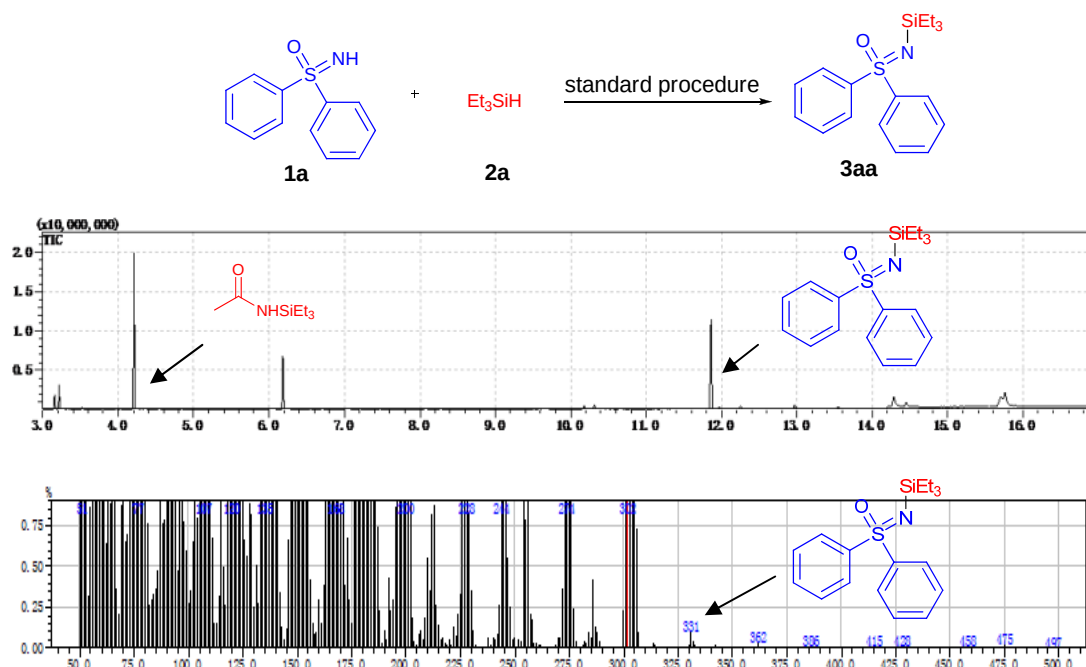
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1 General Experimental Details

Dry CH₃CN was distilled from CaH₂. Chemicals were used as received without special purification unless stated otherwise. ¹H and ¹³C NMR spectra were recorded at ambient temperature on a 400 MHz NMR spectrometer (100 MHz for ¹³C NMR). NMR results were reported in δ units, parts per million (ppm), and were referenced to CDCl₃ (δ 7.26 or 77.0 ppm) as the internal standard. The coupling constants *J* are given in Hz. IR spectra were recorded on a spectrometer using KBr discs.

General Procedure:

A sealed tube was charged with sulfoximine (0.2 mmol), triethylsilane (0.8 mmol, 4.0 equiv.), DTBP (0.6 mmol), CuI (0.02 mmol), H₂O (1.0 equiv.), CH₃CN (dry, 1.0 mL). The mixture was purged with nitrogen and kept stirring under nitrogen at 110 °C for 12 h. The reaction mixture was analyzed using GC-MS spectrometer, as shown in Figure S1. Then the mixture was concentrated in vacuum and the residue was purified by preparative TLC on GF254 (petroleum/ethyl acetate) to afford the desired product.



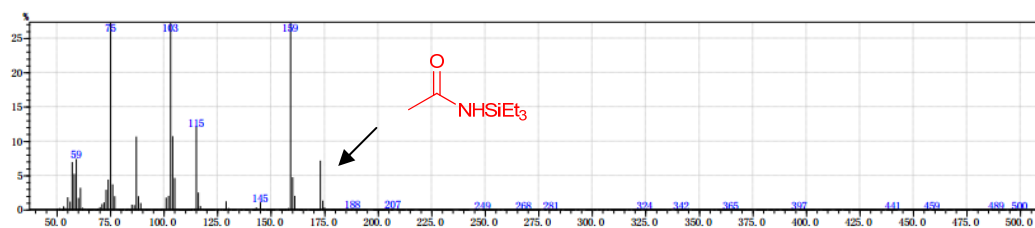


Figure S1 GC-MS spectra of **3aa**.

2 Mechanism Study Experiments

2.1 The Radical Capture Experiment

Under standard procedure, a sealed tube was charged with **1a** (0.2 mmol), **2a** (0.8 mmol), DTBP (0.6 mmol), CuI (0.02 mmol), galvinoxyl free radical (0.6 mmol), H₂O (1.0 equiv.), CH₃CN (dry, 1.0 mL). The reaction was conducted under nitrogen at 110 °C for 12 h. The reaction mixture was analyzed using GC spectrometer, as shown in Figure S2.

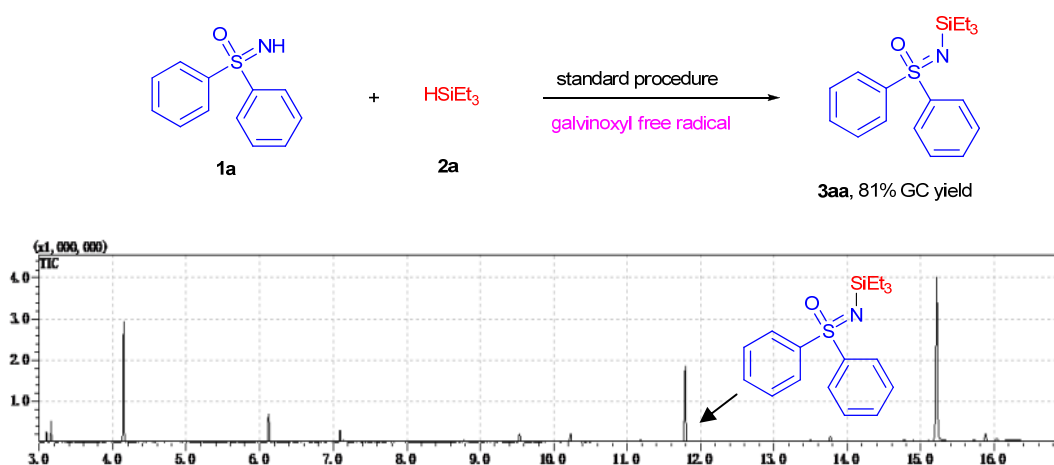


Figure S2 GC spectra of free radical capture result.

2.2 Determination of Reaction Intermediates.

Under standard procedure, a sealed tube was charged with **2e** (0.8 mmol), DTBP (0.6 mmol), CuI (0.2 mmol), CH₃CN (dry, 1.0 mL). The reaction was conducted under nitrogen at 110 °C for 12 h. Then, the reaction mixture was analyzed using GC-MS spectrometer, as shown in Figure S3.

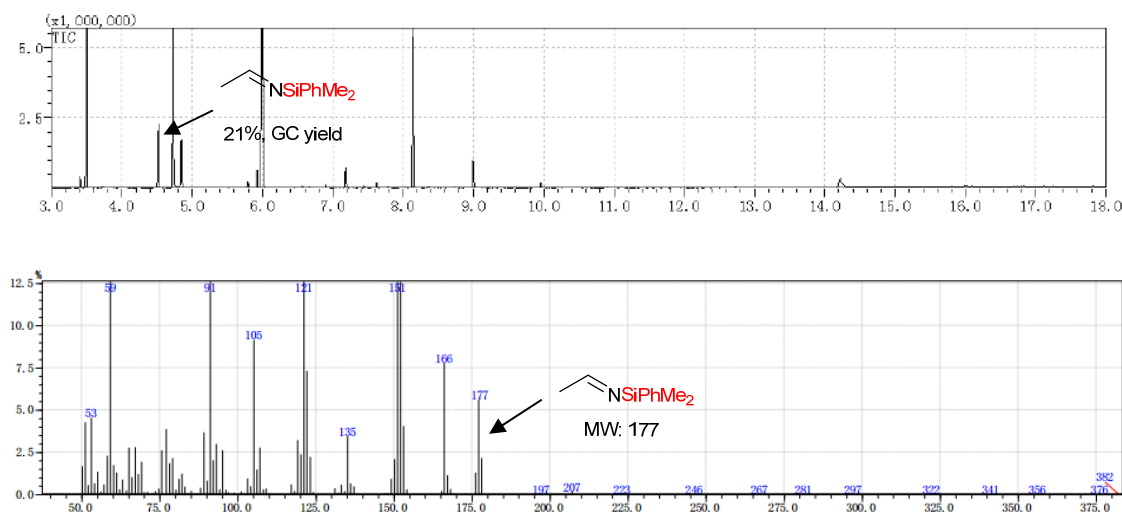


Figure S3

Under standard procedure, a sealed tube was charged with **1a** (0.2 mmol), **2a** (0.8 mmol), DTBP (0.6 mmol), CuI (0.2 mmol), acetamide (0.8 mmol), ethyl acetate (dry, 1.0 mL), H₂O (1.0 equiv). The reaction was conducted under nitrogen at 110 °C for 12 h. Then, the reaction mixture was analyzed using GC spectrometer, as shown in Figure S4.

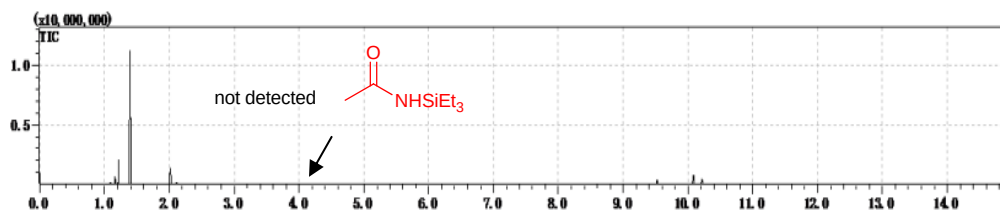


Figure S4

Under standard procedure, a sealed tube was charged with **1a** (0.2 mmol), DTBP (0.6 mmol), *N*-(trimethylsilyl)acetamide (0.8 mmol), H₂O (1.0 equiv.), CH₃CN (dry, 1.0 mL). The reaction was conducted under nitrogen at 110 °C for 12 h. Then, the reaction mixture was analyzed using GC-MS spectrometer, as shown in Figure S5.

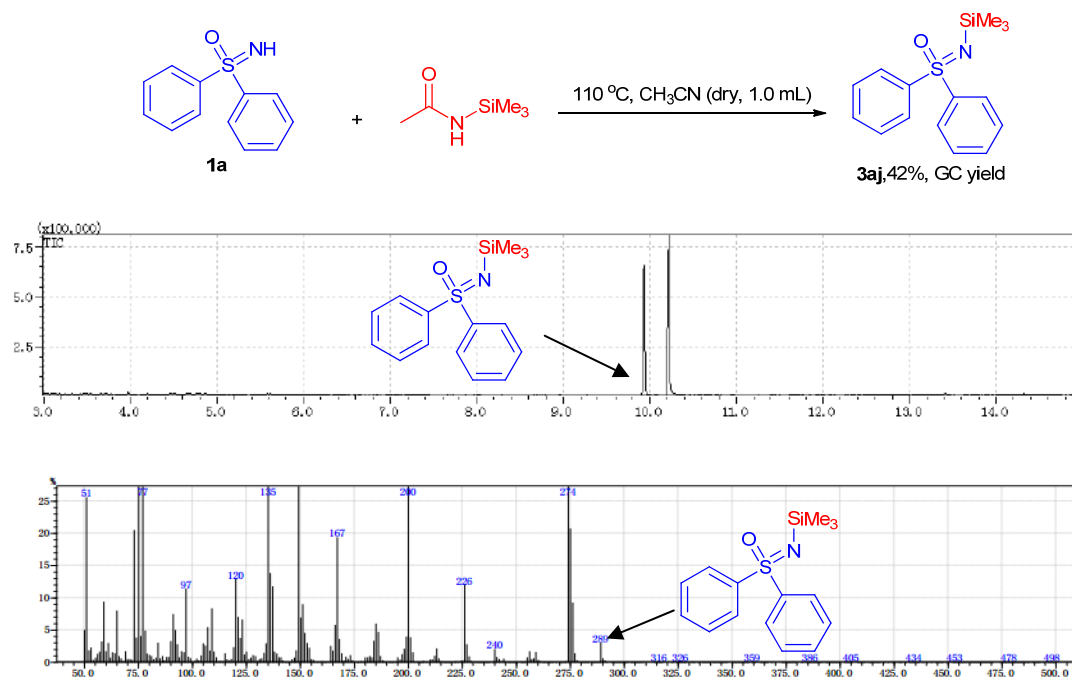
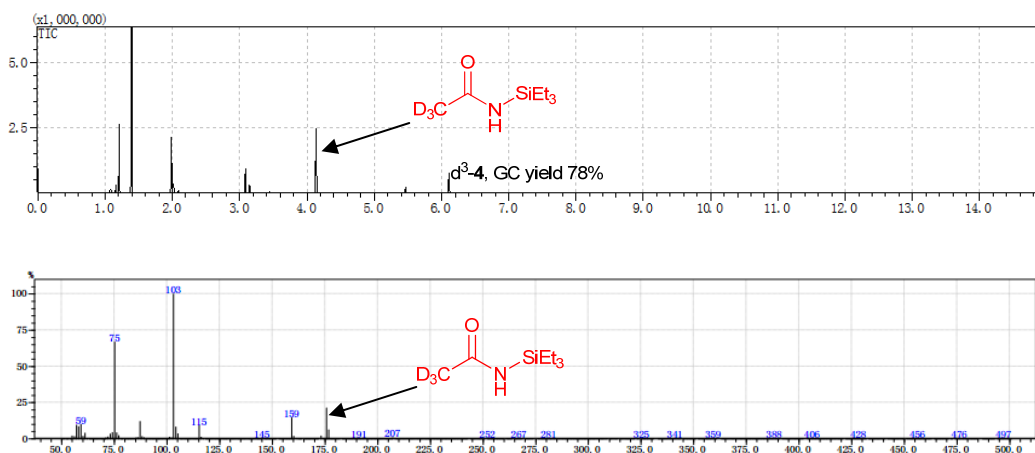
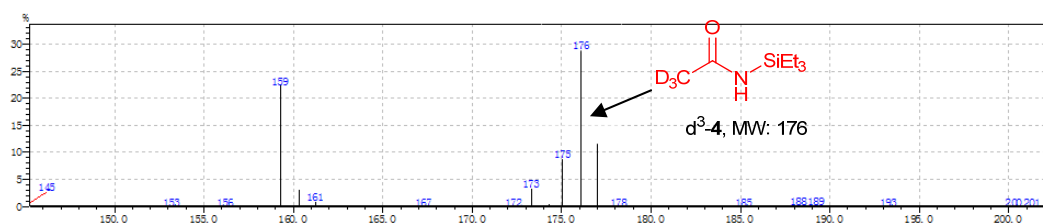


Figure S5 GC-MS spectra of **3aj**.

Under standard procedure, a sealed tube was charged with **2a** (0.8 mmol) DTBP (0.6 mmol), CuI (0.2 mmol), solvent (1.0 mL). The reaction was conducted under nitrogen at 110 °C for 12 h. Then, the reaction mixture was analyzed using GC-MS spectrometer, as shown in Figure S6.

CD₃CN + H₂O (1.0 equiv.)





CH_3CN (dry) + H_2^{18}O (0.1 mL)

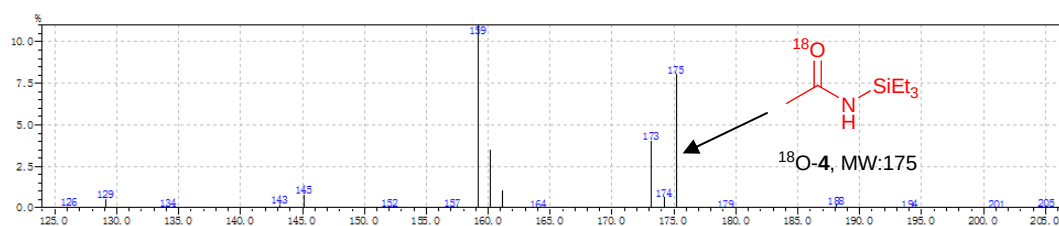
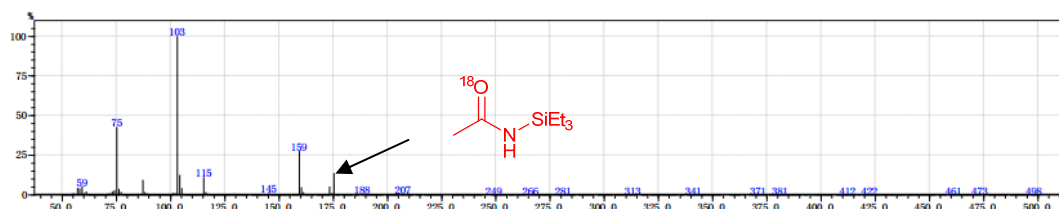
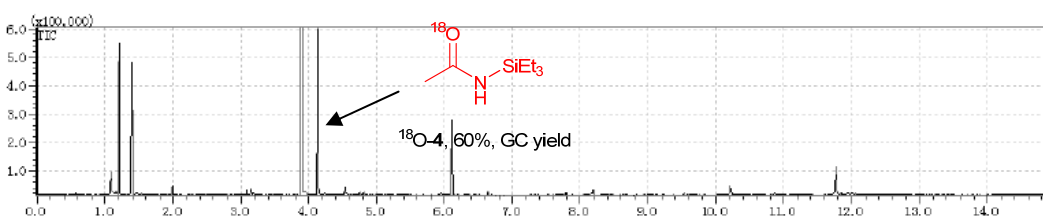
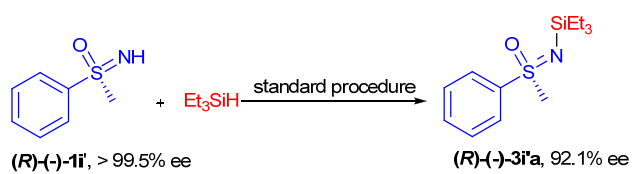


Figure S6. Results of deuterium-labeling experiments

3 Determination of ee value.



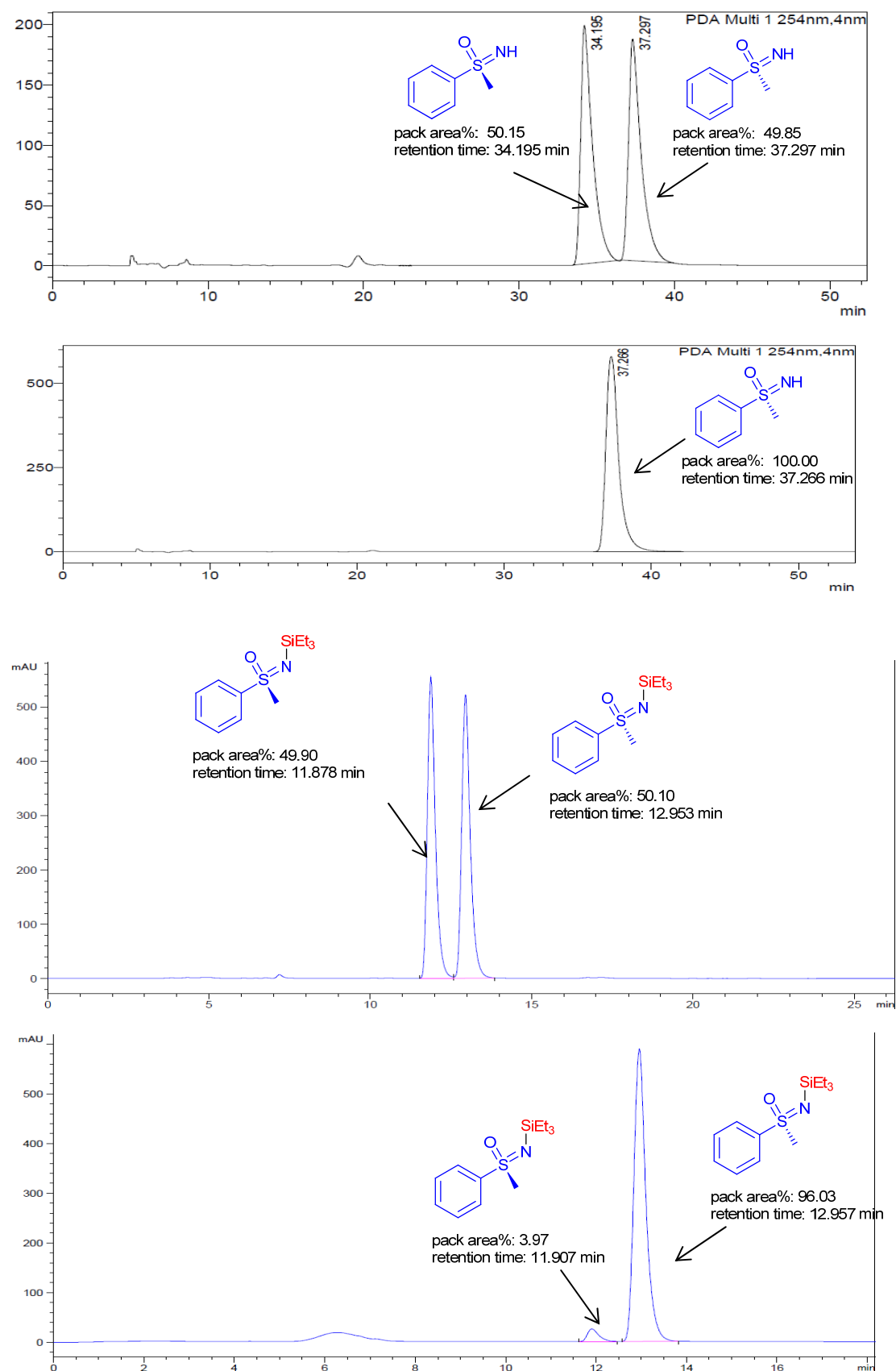
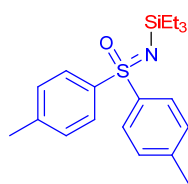
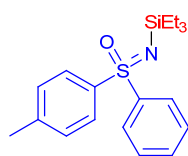


Figure S7. The ee value of (R)-(-)-3i'a detected by HPLC.

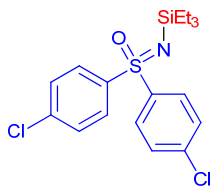
4 Characterization Data for the Products



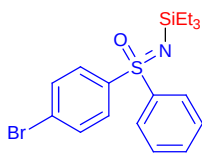
N-(Triethylsilyl)-4,4'-dimethyldiphenyl sulfoximine (3ba): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (54.3 mg, 76% yield) as a colorless liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.81-7.79 (m, 4H), 7.20 (s, 4H), 2.34 (s, 6H), 0.96-0.92 (m, 9H), 0.60-0.54 (m, 6H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 143.7, 142.0, 129.3, 127.2, 21.3, 7.8, 6.6. MS (EI): 359 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{20}\text{H}_{30}\text{NOSSi}$ ($\text{M}+\text{H}$) $^+$ 360.1812, found 360.1812. IR (KBr) ν 3059, 1597, 1490, 1457, 1412, 1159, 1096 cm^{-1} .



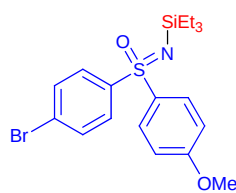
N-(Triethylsilyl)-4-methyldiphenyl sulfoximine (3ca): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (56.8 mg, 82% yield) as a yellowish liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.93-7.91 (m, 2H), 7.81 (d, J = 8.2 Hz, 2H), 7.41-7.39 (m, 3H), 7.20 (d, J = 8.1 Hz, 2H), 2.34 (s, 3H), 0.94 (t, J = 7.9 Hz, 9H), 0.58 (q, J = 7.8 Hz, 6H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 146.5, 143.3, 142.2, 131.4, 129.3, 128.6, 127.3, 127.1, 21.3, 7.2, 6.6. MS (EI): 345 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{19}\text{H}_{28}\text{NOSSi}$ ($\text{M}+\text{H}$) $^+$ 346.1655, found 346.1657. IR (KBr) ν 3062, 1598, 1457, 1318, 1294, 1095 cm^{-1} .



N-(Triethylsilyl)-4,4'-dichlorodiphenyl sulfoximine (3da): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (62.5 mg, 78% yield) as a yellowish liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.85 (d, J = 8.6 Hz, 4H), 7.39 (d, J = 8.6 Hz, 4H), 0.93 (t, J = 7.9 Hz, 9H), 0.58 (q, J = 7.8 Hz, 6H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 144.4, 138.4, 129.1, 128.7, 7.1, 6.5. MS (EI): 399 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{18}\text{H}_{24}\text{Cl}_2\text{NOSSi}$ ($\text{M}+\text{H}$) $^+$ 400.0719, found 400.0719. IR (KBr) ν 3087, 1643, 1577, 1474, 1320, 1163, 1013 cm^{-1} .

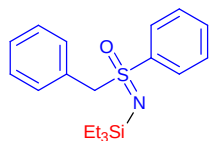


N-(Triethylsilyl)-4-bromodiphenyl sulfoximine (3ea): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (61.3 mg, 75% yield) as a colorless liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.92-7.90 (m, 2H), 7.80 (d, J = 8.5 Hz, 2H), 7.55 (d, J = 8.6 Hz, 2H), δ 7.46-7.40 (m, 2H), 0.93 (t, J = 7.9 Hz, 9H), 0.58 (q, J = 7.8 Hz, 6H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 145.7, 145.3, 131.9, 131.9, 128.9, 128.8, 127.2, 126.6, 7.2, 6.6. MS (EI): 409 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{18}\text{H}_{25}\text{BrNOSSi}$ ($\text{M}+\text{H}$) 410.0604, found 410.0603. IR (KBr) ν 3064, 1641, 1573, 1470, 1320, 1237, 1163, 1009 cm^{-1} .

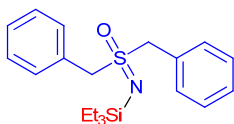


N-(Triethylsilyl)-4-bromo-4'-methoxy-diphenyl sulfoximine (3fa): TLC on GF254 (ethyl acetate: petroleum, 1: 15) give the product (59.7 mg, 68% yield) as a yellowish liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.83 (d, J = 8.9 Hz, 2H), 7.76 (d, J = 8.6 Hz, 2H), 7.53 (d, J = 8.6 Hz, 2H), 6.90 (d, J = 8.9 Hz, 2H), 3.82 (s, 3H), 0.93 (t, J = 7.9 Hz, 9H), 0.57 (q, J = 7.9 Hz, 6H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 162.4, 146.0, 137.5, 131.9, 129.4, 128.6, 126.2,

114.0, 55.5, 7.2, 6.6. MS (EI): 439 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{19}H_{27}BrNO_2SSi$ ($M+H$) $^+$ 440.0710, found 440.0710. IR (KBr) ν 3071, 1727, 1593, 1495, 1463, 1310, 1158, 1095, 1009 cm^{-1} .



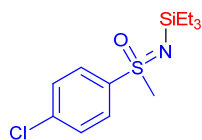
S-Benzyl-N-triethylsilyl-S-phenylsulfoximine (3ga): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (49.3 mg, 71% yield) as a colorless liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.68-7.64 (m, 2H), 7.51-7.47 (m, 1H), 7.42-7.38 (m, 2H), 7.28-7.20 (m, 3H), 7.05 (m, 2H), 4.17 (s, 2H), 0.92-0.86 (m, 1H), 0.56-0.48 (m, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 142.8, 132.2, 131.1, 129.9, 128.4, 128.1, 128.0, 67.3, 7.1, 6.6. MS (EI): 345 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{19}H_{28}NOSSi$ ($M+H$) $^+$ 346.1655, found 346.1655. IR (KBr) ν 3063, 1639, 1456, 1445, 1316, 1298, 1165 cm^{-1} .



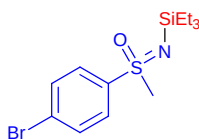
N-(Triethylsilyl) dibenzyl sulfoximine (3ha): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (33.7 mg, 47% yield) as a colorless liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.41-7.36 (m, 10H), 4.12-4.03 (m, 4H), 0.77 (t, J = 8.0 Hz, 9H), 0.34 (t, J = 7.9 Hz, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 131.1, 129.3, 128.4, 128.5, 62.7, 7.0, 6.5. MS (EI): 359 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{20}H_{30}NOSSi$ ($M+H$) $^+$ 360.1812, found 360.1811. IR (KBr) ν 3064, 1603, 1495, 1455, 1314, 1170, 1118 cm^{-1} .



S-Methyl-N-triethylsilyl-S-benzylsulfoximine (3ia): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (27.2 mg, 51% yield) as a colorless liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 8.30-7.95 (m, 2H), 7.57-7.49 (m, 3H), 3.00 (s, 3H), 0.94 (t, J = 7.9 Hz, 9H), 0.58 (q, J = 7.8 Hz, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 145.3, 132.1, 128.8, 126.9, 49.6, 7.1, 6.5. MS (EI): 269 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{13}H_{24}NOSSi$ ($M+H$) $^+$ 270.1342, found 270.1341. IR (KBr) ν 3087, 1643, 1576, 1474, 1320, 1162, 1088, 1012 cm^{-1} .

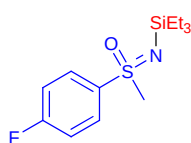


S-Methyl-N-triethylsilyl-S-4-chlorobenzyl sulfoximine (3ja): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (44.8 mg, 74% yield) as a yellow liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.83 (d, J = 8.4 Hz, 2H), 7.41 (d, J = 8.4 Hz, 2H), 2.93 (s, 3H), 0.87 (t, J = 7.9 Hz, 9H), 0.50 (q, J = 7.8 Hz, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 143.8, 138.6, 129.1, 128.5, 49.7, 7.1, 6.5. MS (EI): 303 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{13}H_{23}ClNOSSi$ ($M+H$) $^+$ 304.0953, found 304.0953. IR (KBr) ν 2957, 1635, 1581, 1434, 1312, 1170, 1001 cm^{-1} .

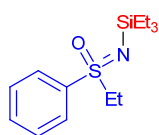


S-Methyl-N-triethylsilyl-S-4-bromophenylsulfoximine (3ka): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (48.1 mg, 69% yield) as a colorless liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.83 (d, J = 8.5 Hz, 2H), 7.65 (d, J = 8.5 Hz, 2H), 3.00 (s, 3H), 0.94 (t, J = 7.9 Hz, 9H), 0.57 (q, J = 7.8 Hz, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 144.4, 132.1, 128.7, 127.1, 49.7, 7.1, 6.5.

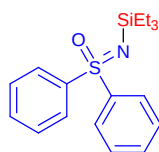
MS (EI): 347 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{13}H_{23}BrNOSSi$ ($M+H$) $^+$ 348.0448, found 348.0448. IR (KBr) ν 2953, 1640, 1573, 1464, 1322, 1157, 1009 cm^{-1} .



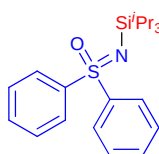
S-Methyl-N-triethylsilyl-S-4-fluorophenylsulfoximine (3la): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (36.0 mg, 63% yield) as a yellowish liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.99-7.95 (m, 2H), 7.21-7.16 (m, 2H), 3.00 (s, 1H), 0.94 (t, J = 7.9 Hz, 9H), 0.57 (q, J = 7.8 Hz, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 164.9 (d, J_{C-F} = 251.7 Hz), 141.4 (d, J_{C-F} = 5.7 Hz), 129.6 (d, J_{C-F} = 9.2 Hz), 115.9 (d, J_{C-F} = 22.3 Hz), 49.8, 7.1, 6.5. MS (EI): 287 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{13}H_{23}FNOSSi$ ($M+H$) $^+$ 288.1248, found 288.1246 IR (KBr) ν 3066, 1594, 1496, 1442, 1309, 1154, 1027 cm^{-1} .



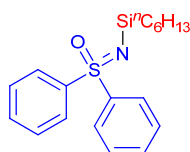
S-Ethyl-N-triethylsilyl-S-benzylsulfoximine (3ma): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (34.0 mg, 60% yield) as a colorless liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.98-7.90 (m, 2H), 7.57-7.48 (m, 3H), 3.00 (q, J = 7.2 Hz, 2H), 1.19 (t, J = 7.4 Hz, 3H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 143.2, 132.1, 128.7, 127.8, 54.7, 8.1, 7.1, 6.6. MS (EI): 283 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{14}H_{26}NOSSi$ ($M+H$) $^+$ 284.1499, found 284.1499. IR (KBr) ν 3064, 1583, 1476, 1445, 1237, 1095 cm^{-1} .



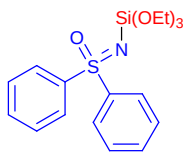
N-(Triethylsilyl) diphenyl sulfoximine (3aa): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (53.0 mg, 80% yield) as a colorless liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.95-7.92 (m, 4H), 7.43-7.38 (m, 6H), 0.94 (t, J = 7.9 Hz, 9H), 0.58 (q, J = 7.8 Hz, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 146.1, 131.6, 128.7, 127.5, 7.1, 6.5. MS (EI): 331 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{18}H_{26}NOSSi$ ($M+H$) $^+$ 332.1499, found 332.1500. IR (KBr) ν 3064, 1583, 1475, 1457, 1446, 1162, 1095 cm^{-1} .



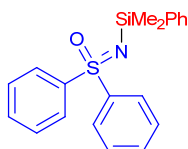
N-(Triisopropylsilyl) diphenyl sulfoximine (3ab): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (56.3 mg, 75% yield) as a yellow liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.97-7.94 (m, 4H), 7.42-7.40 (m, 6H), 1.08-1.03 (m, 21H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 146.6, 131.5, 128.7, 127.2, 18.2, 18.3, 13.2. MS (EI): 373 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{21}H_{32}NOSSi$ ($M+H$) $^+$ 374.1968, found 374.1969. IR (KBr) ν 3065, 1639, 1463, 1327, 1168, 1095 cm^{-1} .



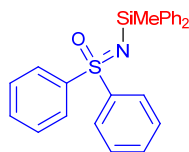
N-(Tri-n-hexylammoniumsilyl) diphenyl sulfoximine (3ac): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (84.4 mg, 85% yield) as a colorless liquid. 1H NMR ($CDCl_3$, 400 MHz): δ 7.93-7.91 (m, 4H), 7.43-7.38 (m, 6H), 1.32-1.21 (m, 24H), 0.85 (t, J = 6.6 Hz, 9H), 0.59-0.55 (m, 6H); ^{13}C NMR ($CDCl_3$, 100 MHz): δ 146.2, 131.5, 128.6, 127.3, 33.3, 31.5, 23.5, 22.6, 15.8, 14.1. MS (EI): 499 (M^+); HRMS (ESI-TOF) m/z calcd for $C_{30}H_{50}NOSSi$ ($M+H$) $^+$ 500.3377, found 500.3378. IR (KBr) ν 3064, 1584, 1466, 1446, 1319, 1296, 1164, 1095 cm^{-1} .



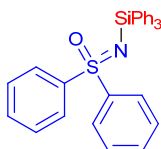
N-(Triethoxysilyl) diphenyl sulfoximine (3ad): TLC on GF254 (ethyl acetate: petroleum, 1: 15) give the product (33.4 mg, 44% yield) as a colorless liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 8.00-7.97 (m, 4H), 7.48-7.43 (m, 6H), 3.82 (q, $J = 7.0$ Hz, 6H), 1.15 (q, $J = 7.0$ Hz, 9H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 144.8, 132.1, 128.2, 127.4, 58.8, 18.0. MS (EI): 379 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{18}\text{H}_{26}\text{NO}_4\text{SSi}$ ($\text{M}+\text{H}$) $^+$ 380.1346, found 380.1346. IR (KBr) ν 3067, 1636, 1446, 1428, 1286, 1164 cm^{-1} .



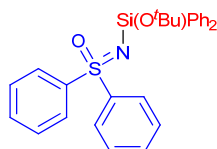
N-(Dimethylphenylsilyl) diphenyl sulfoximine (3ae): TLC on GF254 (ethyl acetate: petroleum, 1: 20) give the product (42.3 mg, 60% yield) as a colorless liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.85 (d, $J = 8.8$ Hz, 1H), 7.67 (d, $J = 6.6$ Hz, 5H), 7.36-7.30 (m, 9H), 6.88 (d, $J = 8.8$ Hz, 1H), 3.82 (s, 3H), 2.86 (s, 3H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 136.6, 135.3, 135.2, 129.8, 129.3, 129.1, 127.6, 113.9, 55.6, 49.3. MS (EI): 351 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{20}\text{H}_{22}\text{NOSSi}$ ($\text{M}+\text{H}$) $^+$ 352.1186, found 352.1185. IR (KBr) ν 3048, 1594, 1497, 1442, 1309, 1258, 1154, 1112, 1027 cm^{-1} .



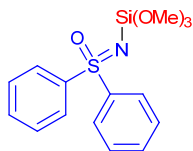
N-(Diphenylmethylsilyl) diphenyl sulfoximine (3af): TLC on GF254 (ethyl acetate: petroleum, 1: 15) give the product (61.7 mg, 75% yield) as a colorless liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.89 (d, $J = 7.2$ Hz, 4H), 7.62 (d, $J = 7.3$ Hz, 4H), 7.40-7.26 (m, 12H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 145.1, 138.4, 134.2, 129.0, 128.7, 127.5, 127.3, 0.2. MS (EI): 413 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{25}\text{H}_{24}\text{NOSSi}$ ($\text{M}+\text{H}$) $^+$ 414.1342, found 414.1344. IR (KBr) ν 3066, 1636, 1475, 1446, 1296, 1164, 1096 cm^{-1} .



N-(Triphenylsilyl) diphenyl sulfoximine (3ag): TLC on GF254 (ethyl acetate: petroleum, 1: 15) give the product (63.4 mg, 67% yield) as a colorless liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.86-7.84 (m, 4H), 7.63-7.61 (m, 6H), 7.37-7.22 (m, 15H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 144.84, 136.33, 135.3, 131.7, 129.2, 128.7, 127.5, 127.3. MS (EI): 475 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{30}\text{H}_{26}\text{NOSSi}$ ($\text{M}+\text{H}$) $^+$ 476.1499, found 476.1501. IR (KBr) ν 2953, 1640, 1573, 1464, 1321, 1157, 1010 cm^{-1} .



N-(Diphenyl-tert-butoxysilyl) diphenyl sulfoximine (3ah): TLC on GF254 (ethyl acetate: petroleum, 1: 10) give the product (43.5 mg, 46% yield) as a yellowish liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 7.87-7.85 (m, 3H), 7.67-7.65 (m, 4H), 7.40-7.21 (m, 13H), 1.26 (s, 9H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 145.0, 137.9, 134.7, 131.7, 129.0, 128.7, 127.3, 127.3, 31.9. MS (EI): 471 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{28}\text{H}_{30}\text{NO}_2\text{SSi}$ ($\text{M}+\text{H}$) $^+$ 472.1761, found 472.1761. IR (KBr) ν 3067, 1590, 1446, 1364, 1296, 1052 cm^{-1} .



N-(Trimethoxysilyl) diphenyl sulfoximine (3ai): TLC on GF254 (ethyl acetate: petroleum, 1: 15) give the product (35.2 mg, 52% yield) as a yellowish liquid. ^1H NMR (CDCl_3 , 400 MHz): δ 8.00-7.98 (m, 3H), 7.49-7.45 (m, 7H), 3.53 (s, 9H); ^{13}C NMR (CDCl_3 , 100 MHz): δ 144.6, 132.2, 128.9, 127.3, 51.0. MS (EI): 337 (M^+); HRMS (ESI-TOF) m/z calcd for $\text{C}_{15}\text{H}_{19}\text{NO}_4\text{SSi}$ ($\text{M}+\text{H}$) $^+$ 338.0877, found 338.0879. IR (KBr) ν 3061, 1644, 1446, 1301, 1232, 1184, 1094 cm^{-1} .

5 Copies of ^1H NMR and ^{13}C NMR spectra

