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Addition/Correction

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Correction to "Asymmetric Reduction of *tert*-Butanesulfinyl Ketimines by N-Heterocyclic Carbene Boranes"

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Page 11444. We regret that there were unintentional nomenclature errors in the title of the compounds. The first "R" in the names of compound **7a−r** should be changed to "S" as follows:

(*S*)-2-Methyl-*N*-((*R*)-1-phenylethyl)propane-2-sulfinamide (7a)[15]. A colorless oil (50.6 mg, 90%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.30−7.36 (m, 5H), 4.57 (m, 1 H), 3.46 (s, 1H), 1.53 (d, J = 6.4 Hz, 3H), 1.26 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 144.1, 128.7, 127.8, 126.6, 55.5, 54.0, 22.8, 22.6.

(*S*)-2-Methyl-*N*-((*R*)-1-*p*-tolylethyl)propane-2-sulfinamide (7b). A colorless oil (52.0 mg, 87%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.26 (d, J = 8.0 Hz, 2H), 7.17 (d, J = 7.6 Hz, 2H), 4.54 (m, 1H), 3.42 (s, 1H), 2.36 (s, 3H), 1.51 (d, J = 6.8 Hz, 3H), 1.25 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 114.2, 137.5, 129.4, 126.5, 55.4, 53.7, 22.7, 22.6, 21.1. HRMS (ESI-TOF) m/z: $[M + H]^+$ calcd for $C_{13}H_{21}NOS$ 240.1417, found 240.1427.

(*S*)-*N*-((*R*)-1-(4-Methoxyphenyl)ethyl)-2-methylpropane-2-sulfinamide (7c)[15]. A colorless oil (56.1 mg, 88%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.29 (d, J = 7.6 Hz, 2H), 6.90 (d, J = 8.8 Hz, 2H), 4.53 (m, 1H), 3.82 (s, 3H), 3.43 (s, 1H), 1.51 (d, J = 6.4 Hz, 3H), 1.25 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 159.2, 136.2, 127.7, 114.1, 55.4, 55.3, 53.4, 22.7, 22.6.

(*S*)-*N*-((*R*)-1-(4-Chlorophenyl)ethyl)-2-methylpropane-2-sulfinamide (7d)[16]. A colorless oil (57.8 mg, 89%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.26−7.38 (m, 4H), 4.53 (m, 1H), 3.41 (s, 1H), 1.50 (d, J = 6.8 Hz, 3H), 1.25 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 142.5, 133.5, 128.9, 128.0, 55.6, 53.5, 22.8, 22.6. HRMS (ESI-TOF) m/z: $[M + H]^+$ calcd for $C_{12}H_{18}ClNOS$ 260.0870, found 260.0880.

(*S*)-2-Methyl-*N*-((*R*)-1-(4-(trifluoromethyl)phenyl)-ethyl)propane-2-sulfinamide (7e)[15]. A colorless oil (60.1 mg, 82%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.63 (d, J = 8.4 Hz, 2H), δ 7.49 (d, J = 8.4 Hz, 2H), 4.62 (m, 1H), 3.47 (d, J = 2.0 Hz, 1H), 1.55 (d, J = 6.4 Hz, 3H), 1.26 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 148.0, 127.0, 125.7, 125.7, 55.7, 53.9, 22.9, 22.5.

(*S*)-*N*-((*R*)-1-(4-Cyanophenyl)ethyl)-2-methylpropane-2-sulfinamide (7f)[15]. A colorless oil (59.5 mg, 95%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.62 (d, J = 8.4 Hz, 2H), 7.47 (d, J = 8.4 Hz, 2H), 4.58 (m, 1H), 3.59 (d, J = 3.6 Hz, 1H), 1.51 (d, J = 6.4 Hz, 3H), 1.22 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 149.3, 132.6, 127.4, 118.6, 111.6, 55.8, 54.1, 22.9, 22.5.

(*S*)-*N*-((*R*)-1-(3-Fluorophenyl)ethyl)-2-methylpropane-2-sulfinamide (7g). A colorless oil (52.3 mg, 86%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 6.97−7.28 (m, 4H), 4.56 (m, 1H), 3.48 (s, 1H), 1.52 (d, J = 6.8 Hz, 3H), 1.26 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 163.0 (d, J = 245.0 Hz), 146.7 (d, J = 7.0 Hz), 130.3 (d, J = 8.0 Hz), 122.3 (d, J = 3.0 Hz), 114.7 (d, J = 21.0 Hz), 113.4 (d, J = 22.0 Hz), 55.6, 53.6, 22.8, 22.6. HRMS (ESI-TOF) m/z: $[M + H]^+$ calcd for $C_{12}H_{18}FNOS$ 244.1166, found 244.1174.

(*S*)-2-Methyl-*N*-((*R*)-1-phenylpropyl)propane-2-sulfinamide (7h)[15]. A colorless oil (50.9 mg, 85%)> ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.30−7.36 (m, 5H), 4.31 (m, 1H), 3.41 (s, 1H), 1.80 (m, 2H), 1.25 (s, 9H), 0.81 (t, J = 14.8 Hz, 3H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 142.3, 128.6, 127.8, 127.3, 60.4, 55.7, 29.4, 22.6, 22.5.

(*S*)-*N*-((*R*)-1-(4-Methoxyphenyl)-2,2-dimethylpropyl)-2-methylpropane-2-sulfinamide (7i). A colorless oil (55.8 mg, 75%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.96 (d, J = 8.4 Hz, 2H), 6.85 (d, J = 8.4 Hz, 2H), 3.81 (d, J = 8.0 Hz, 1H), 3.60 (s, 3H), 3.11 (s, 1H), 1.24 (s, 9H), 0.95 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 159.0, 130.6, 129.2, 113.0, 66.5, 55.8, 55.1, 36.3, 26.7, 22.6. HRMS (ESI-TOF) m/z: $[M + H]^+$ calcd for $C_{16}H_{27}NO_2S$ 298.1835, found 298.1828.

(*S*)-*N*-((*R*)-(2-Hydroxyphenyl) (phenyl)methyl)-2-methylpropane-2-sulfinamide (7j)[19]. A colorless oil (39.4 mg, 52%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 6.73−7.43 (m, 9H), 5.82 (d, J = 4.8 Hz, 1H), 5.06 (d, J = 4.8 Hz, 1H), 1.32 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 155.1, 140.5, 129.1, 128.9, 128.4, 128.3, 127.5, 127.5, 119.3, 116.5, 60.2, 56.3, 22.8.

(*S*)-*N*-((*R*)-(2-Methoxyphenyl) (phenyl)methyl)-2-methylpropane-2-sulfinamide (7k)[20]. A colorless oil (38.9 mg, 49%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 6.73−7.43 (m, 9H), 5.81 (d, J = 4.4 Hz, 1H), 5.20 (d, J = 4.4 Hz, 1H), 1.32 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 156.6, 141.2, 130.8, 128.9, 128.5, 128.2, 128.0, 128.0, 127.5, 127.3, 57.4, 55.9, 55.5, 22.7.

(*S*)-2-Methyl-*N*-((*R*)-1-(pyridin-2-yl)ethyl)propane-2-sulfinamide (7l)[21]. A colorless oil (49.2 mg, 74%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 8.60 (d, J = 4.8 Hz, 1H), δ 7.68 (m, 1H), 7.30 (m, 1H), 7.19 (m, 1H), 4.85 (d, J = 4.4 Hz, 1H), 4.64 (m, 1H), 1.52 (d, J = 6.8 Hz, 3H), 1.27 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 161.9, 149.0, 136.8, 122.3, 121.0, 55.6, 55.2, 23.4, 22.7.

(*S*)-*N*-((*R*)-1-(Furan-2-yl)ethyl)-2-methylpropane-2-sulfinamide (7m)[16]. A colorless oil (48.4 mg, 90%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.38 (t, J = 3.6 Hz, 1H), 6.27−6.34 (m, 2H), 4.59 (m, 1H), 3.59 (d, J = 4.0 Hz, 1H), 1.62 (d, J = 6.8 Hz, 3H), 1.23 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 156.0, 142.0, 110.2, 106.1, 55.7, 48.6, 22.5, 20.2. HRMS (ESI-TOF) m/z: $[M + H]^+$ calcd for $C_{10}H_{17}NO_2S$ 216.1053, found 216.1059.

(*S*)-*N*-((*S*)-2-Chloro-1-(4-chlorophenyl)ethyl)-2-methylpropane-2-sulfinamide (7n)[22]. A colorless oil (66.2 mg, 90%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.28−7.38 (m, 4H), 4.68 (m, 1H), 3.87 (d, J = 5.6 Hz, 2H), 3.84 (s, 1H), 1.25 (s, 9H).

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^{13}C NMR (100 MHz, $CDCl_3$) δ 137.5, 134.4, 129.0, 128.8, 67.9, 58.9, 48.3, 22.6.

(*S*)-2-Methyl-*N*-((*R*)-1,2,3,4-tetrahydronaphthalen-1-yl)propane-2-sulfinamide (7o)[15]. A white solid (50.3 mg, 80%), mp: 116 °C. ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.11−7.48 (m, 4H), 4.60 (m, 1H), 3.26 (s, 1H), 2.80 (m, 1H), 1.81−2.06 (m, 4H), 1.24 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 137.7, 137.0, 129.7, 129.2, 127.6, 126.6, 55.4, 52.8, 30.6, 29.1, 22.7, 18.2.

(*S*)-*N*-((*R*)-3,3-Dimethylbutan-2-yl)-2-methylpropane-2-sulfinamide (7p)[15]. A colorless oil (43.6 mg, 85%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 3.27 (s, 1H), 3.12 (m, 1H), 1.21 (s, 9H), 1.13 (d, J = 6.4 Hz, 3H), 0.92 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 59.0, 55.4, 34.4, 26.1, 22.6, 15.9.

(*S*)-2-Methyl-*N*-((*R*)-4-phenylbutan-2-yl)propane-2-sulfinamide (7q)[15]. A colorless oil (52.6 mg, 83%). ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.31 (d, J = 7.6 Hz, 2H), 7.21 (m, 3H), 3.43 (m, 1H), 3.16 (d, J = 4.4 Hz, 1H), 2.72 (m, 2H), 1.79−1.95 (m, 2H), 1.23 (d, J = 6.4 Hz, 3H), 1.20 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 141.5, 128.5, 128.4, 126.0, 55.3, 51.1, 39.9, 32.1, 22.5, 21.7.

(*S*)-*N*-((*S*,*E*)-1,3-Diphenylallyl)-2-methylpropane-2-sulfinamide (7r)[23]. A white solid (47.8 mg, 61%), mp: 107−109 °C. ^{1}H NMR (400 MHz, $CDCl_3$) δ 7.30−7.46 (m, 10H), 6.67 (m, 1H), 6.43 (m, 1H), 5.18 (d, J = 7.6 Hz, 1H), 3.63 (s, 1H), 1.26 (s, 9H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 140.5, 133.0, 128.7, 128.6, 128.5, 128.5, 128.4, 128.4, 128.0, 127.4, 126.4, 126.1, 40.4, 30.2, 22.7.

Page 11445. As "$[\alpha]_D$25 + 22.8 (c 0.75, CH_2Cl2)" was the optical rotation of the corresponding free amine of **8c**, it was not suitable to be placed in the data of **8c**. The data part of **8c** should be rewritten as "(*R*)-1-(4-methoxyphenyl)ethanamine hydrochloride (**8c**)[18]: a yellow solid (31.9 mg, 85%). mp: 152−155°C. ^{1}H NMR (400 MHz, $CDCl_3$) δ 8.61 (s, 2H), 7.42 (d, J = 8.4 Hz, 2H), 6.89 (d, J = 8.4 Hz, 2H), 4.33 (s, 1H), 3.81 (s, 3H), 1.65 (d, J = 6.8 Hz, 3H). ^{13}C NMR (100 MHz, $CDCl_3$) δ 159.9, 129.7, 128.3, 114.4, 55.2, 51.3, 20.5."

P11443. "$[\alpha]_D$25 + 22.8 (c 0.75, CH_2Cl_2)" should be added to the footnote of Table 3 as follows:

"[a]Unless otherwise noted, all reactions were carried out using **6** (0.25 mmol), NHC-BH_3 (0.25 mmol), and TsOH (0.125 mmol) in MeOH (1.0 mL) at −10 °C for 4 h. [b]Isolated yield. [c]Determined by ^{1}H NMR analysis of unpurified reaction mixtures. [d]CH_2Cl_2 was used as the solvent without TsOH for 8 h. [e]CH_2Cl_2 was used as the solvent for 8 h. [f]Toluene was used as the solvent. [g]The absolute configuration was determined by comparison of the optical rotation with the known compounds (corresponding free amine), $[\alpha]_D$25 + 22.8 (c 0.75, CH_2Cl_2) {literature[24] $[\alpha]_D$20 + 23.5 (c 0.9, $CHCl_3$, 95% ee)}."

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