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Additions and Corrections

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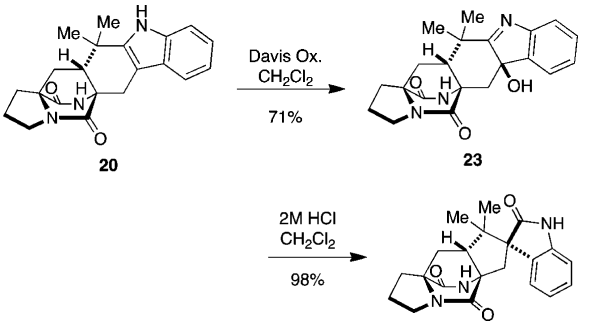
Improved Biomimetic Total Synthesis of D,L-Stephacidin A

Page 4255. The structure of product (**23**) of the reaction of **20** with Davis oxaziridine was incorrectly assigned. On the basis of the ^{13}C NMR data, the structure of **23** should be as shown below. This does not impact the bulk of the manuscript that details an improved biomimetic synthesis of stephacidin A.



To achieve the construction of the reported spiro-oxindole, compound **23** (11 mg, 0.03 mmol) was subjected to treatment with 2 M HCl (1.1 mL) in CH_2Cl_2 (1.1 mL) for 48 h at room temperature. The solution was extracted with CH_2Cl_2 (3 × 10 mL); the combined organic layer was dried over Na_2SO_4 and concentrated. The desired compound was obtained as a yellow residue (10.7 mg, 0.029 mmol) in 98% yield: ^1H NMR (400 MHz, $CDCl_3$) δ 0.80 (s, 3H), 0.86 (s, 3H), 1.71−1.65 (m, 1H), 1.83−1.78 (m, 2H), 2.09−1.98 (m, 2H), 2.20 (d, J = 14.8 Hz, 1H), 2.80−2.73 (m, 1H), 2.90 (d, J = 29.2 Hz, 1H), 3.25 (d, J = 14.8 Hz, 1H), 3.51−3.42 (m, 2H), 6.83 (d, J = 7.6 Hz, 1H), 7.02−6.98 (m, 1H), 7.21−7.17 (m, 1H), 7.40 (d, J = 7.6 Hz, 1H) 8.10 (bs, 1H); ^{13}C NMR (100 MHz, $CDCl_3$) δ 19.8, 23.3, 24.8, 29.6, 30.6, 34.4, 43.9, 46.0, 62.4, 66.7, 68.8, 77.2, 109.4, 122.2, 126.9, 128.2, 130.2, 140.9, 169.5, 174.3,

183.5; HRMS (ESI/APCI) calcd for $C_{21}H_{24}N_3O_3$ (M + H) 366.1812, found 366.1833.

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