Organic LETTERS

Addition/Correction

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Correction to Total Synthesis of (−)-Histrionicotoxin

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Org. Lett. **2011**, *13*(16), 4466−4449. DOI: 10.1021/ol2018032

Supporting Information

Solvent peaks had been removed from the ^{1}H NMR spectra reported for compounds **8**, **12**, **S3**, and **S10**. Peaks of impurities had been removed from the ^{1}H NMR spectra reported for compounds **6** and **S18**. Original FIDs were located, and the spectra were reprocessed and have been replaced for the above compounds in the revised Supporting Information submitted with this correction. The spectra editing did not affect any of the conclusions of the published paper. The purities calculated on the basis of the revised spectra and corrected yields are as follows: **8** (95% purity, 91% yield), **12** (95% purity, 92% yield), **S3** (97% purity, 88% yield), and **S10** (96% purity, 86% yield). Impurities included in **6** were tri-*n*-butylphosphine and ethanol, and the purity and the yield were calculated as 58% purity and 50% yield on the basis of the revised spectrum. The sample of **S18** included some unknown impurities; thus, the exact purity could not be determined.

■ ASSOCIATED CONTENT

Supporting Information

Revised Supporting Information including reprocessed spectra. This material is available free of charge via the Internet at http://pubs.acs.org.

Published: February 3, 2014