

Correction to Direct Access to Functionalized Azepanes by Cross-Coupling with α -Halo Eneformamides

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Supporting Information

C olvent peaks and in some cases resonances corresponding to minor diastereomers, rotamers, or impurities had been removed from the ¹H NMR spectra reported for compounds 1a, 5, 6, 7, 8, 12, 14–16, 20, 25, 26, 28, 29, and 32 by one of us (TKB). 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 except for diastereomeric ratios for 8 (92:8 E/Z) and 14 (90:10 E/Z). The purities calculated on the basis of the revised spectra and corrected yields are as follows: 1a (97% purity, 85% yield), 5 (96% purity, 72% yield), 6 (99% purity, 63% yield), 7 (95% purity, 73% yield), 8 (99% purity; 92:8 diastereomeric ratio; peaks corresponding to a minor diastereomer were deleted), 12 (80% purity, 52% yield), 14 (97% purity, 82% yield of a 9:1 diastereomeric mixture), 15 (96% purity, 86% yield), 16 (95% purity, 88% yield), 20 (95% purity, 15% yield), 25 (95% purity, 83% yield), 26 (99% purity, 80% yield), 28 (91% purity, 80% yield), 29 (95% purity, 67% yield), and 32 (90% purity, 55% vield).

ASSOCIATED CONTENT

S Supporting Information

Revised file containing experimental data and spectra; FIDs for the compounds described above. This material is available free of charge via the Internet at http://pubs.acs.org.

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