## **Author Correction:** Lactonization as a general route to $\beta$ -C( $sp^3$ )–H **functionalization**

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In this Article, the instability of the β-lactone intermediates led to ring opening on storage and resulted in the formation of impurities. We have re-made volatile lactones 2a, 2c, 2d, 2e and 2ac and re-recorded their nuclear magnetic resonance (NMR) spectra immediately after the reaction and also after filtration through silica. We have also hydrolysed 2a, 2c, 2d, 2e, 2ac, 2g and 2h to their corresponding β-hydroxy acid products and recorded the spectra of the resulting compounds: 2a', 2c', 2d', 2e', 2g', 2h' and 2ac'. In addition, we have re-made compounds 3j, 3l, 3m and 3o, owing to the presence of impurities, and the <sup>1</sup>H NMR spectrum of each compound was recorded immediately after purification. In the Supplementary Information of the original Article the spectra after the reaction and then after filtration through silica for compounds 2a, 2c, 2d, 2e, 2ac, and re-made compounds 3j, 3l, 3m and 30 have been corrected. For transparency, we present the corrected and originally published spectra of compounds 2a, 2c, 2d, 2e, 2ac, 3j, 31, 3m and 3o and also the spectra of compounds 2a', 2c', 2d', 2e', 2g', **2h'** and **2ac'** in the Supplementary Information of this Amendment. The synthetic procedures have been updated in the Supplementary Information section 'General Procedure for β-C(sp³)–H Lactonization' accordingly, and the original version is in the Supplementary Information of this Amendment for comparison. We thank the readers who brought these issues to our attention. The Supplementary Information of the original paper has been corrected.

Supplementary information is available in the online version of this Amendment.