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Correction to "Oxidative Cleavage of Cellobiose by Lytic Polysaccharide Monooxygenase (LPMO)-Inspired Copper Complexes"

Andrea. C. Neira, Paulina R. Martínez-Alanis, Gabriel Aullón, Marcos Flores-Alamo, Paulino Zerón, Anna Company, Juan Chen, Johann B. Kasper, Wesley R. Browne, Ebbe Nordlander, and Ivan Castillo*

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It should be specified that in the preparative scale oxidations of cellobiose, the total volume of the reaction mixture was 2 mL at a phosphate buffer concentration of 200 mM. The column employed for HPLC-MS analysis was Poroshell 120 EC-C18 (2.7 μm), and samples were filtered through a short plug of Celite prior to analysis to eliminate the copper complexes.

We identified the peak in Figure S19 in the Supporting Information as a leaching product from the HPLC column when $\text{H}_2\text{O}_2/\text{Et}_3\text{N}$ is employed as oxidant for cellobiose degradation. Upon further analysis, the peak giving rise to $m/z = 118$ and 235 is consistent with protonated triethylamine *N*-oxide $[\text{Et}_3\text{NOH}]^+$ and $[(\text{Et}_3\text{NO})_2\text{H}]^+$, respectively.

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