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

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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  $\mu\text{m}$ ), 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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