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Copper complexes as bioinspired models for Lytic Polysaccharide Monoxygenases.

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Supporting Information Placeholder

ABSTRACT: We report here two copper complexes as first functional models for Lytic Polysaccharide Monoxygenases, mononuclear copper-containing enzymes involved in recalcitrant polysaccharide breakdown. These complexes feature similar structural and spectroscopic properties to that of the enzyme. In addition, they catalyze the oxidative cleavage of the model substrate *p*-nitrophenyl- β -D-glucopyranoside. More importantly, a particularly stable Cu(II)-hydroperoxide intermediate is detected in the reaction conditions.

Much attention has been recently directed to Lytic Polysaccharide Monoxygenases (LPMOs) for their boosting activity during enzymatic conversion of biomass recalcitrant polysaccharides (cellulose, chitin etc) *via* an oxidative cleavage mechanism.¹ LPMOs are copper-containing monoxygenases catalyzing the oxidation of glycosidic bonds creating new polymer chain breaks, which facilitates the action of classical glycoside hydrolases. Dioxygen activation at the copper site in the presence electron donors (ascorbate, proteic partner, etc) leads to the hydroxylation of an inert C-H bond at the glycosidic linkage (Figure 1). This glycosidic bond is further cleaved off upon elimination of a water molecule. Hydroxylation at position C1 is preferred but LPMOs performing the oxidative cleavage of cellulose at position C4 has also been reported.^{1,2} LPMOs harbor a solvent exposed mononuclear copper active site at the center of a flat surface that interacts with the polysaccharide substrate.³ The active site copper ion is coordinated by two histidine residues, including the *N*-terminal histidine that is bound both by the side chain imidazole and the main-chain amino group ('Histidine-brace motif'). A tyrosine or a phenylalanine residue is located

in axial position (Figure 1).¹ Finally, the imidazole moiety of the *N*-terminal histidine is *N*-methylated in fungal LPMOs.

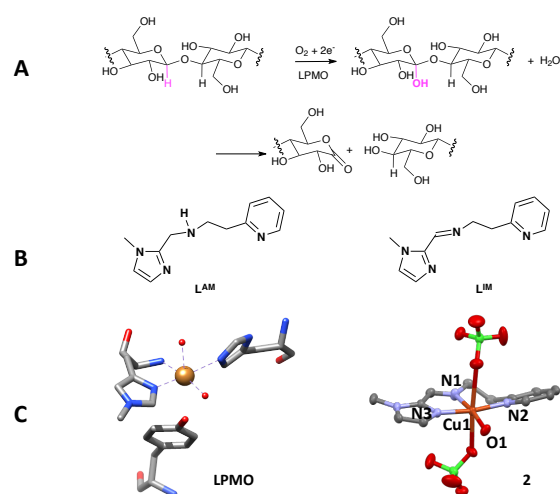
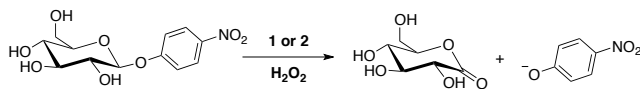


Figure 1. A) Oxidative cleavage of polysaccharides catalyzed by LPMO. B) Ligands used in the present study. C) Active site of LPMO from *Thermoascus auriantacus* (pdb 2yet) and ORTEP diagram of complex 2 (30% thermal ellipsoid plots). Hydrogen atoms were omitted for clarity. Selected distances in Å: Cu1-N1: 2.000(2), Cu1-N2: 2.001(2), Cu1-N3: 1.973(2), Cu1-O1: 1.9771(19).

LPMOs are members of the mononuclear copper containing monoxygenases in which reactive copper-oxygen species are involved in C-H bond hydroxylation.⁴ In the case of LPMOs, the nature of the copper-oxygen intermediates involved in C-H bond activation is still debated.⁵ Several copper-oxygen intermediates have been characterized using biomimetic complexes,⁶ and much can be learnt on LPMOs mechanism by using a small model approach. Recently, a Cu(II) complex based on bis(benzimidazole)amine ligand was prepared as structural model for LPMOs but no oxidative reactivity was reported.⁷ We have prepared two ligands

(L^{AM} and L^{IM}) providing a N₃-coordination sphere for a copper ion and featuring N-methylated imidazole moiety (Figure 1).⁸ The corresponding complexes [(L^{AM})Cu(CH₃CN)](ClO₄)₂ (**1**) and [(L^{IM})Cu(OH₂)](ClO₄)₂·H₂O (**2**) were prepared. X-ray diffraction analysis have shown that, in both cases, the Cu(II) ions are in pseudo-octahedral geometries and are coordinated in the equatorial plane by the 3 nitrogen donors from the ligands (Figure 1 and SI). The Cu-N distances are found ranging from ca. 1.97 to 2.03 Å and are within the range of distances found in similar complexes⁸ as well as in the enzymatic systems.^{1,3} The Electron Paramagnetic Resonance (EPR) spectra of the complexes were recorded in aqueous solution (SI). The EPR parameters obtained from simulation with axial set of parameters are the following $g_{\parallel} = 2.260$, $|A_{\parallel}| = 530$ MHz, $g_{\perp} = 2.059$ for **1** and $g_{\parallel} = 2.265$, $|A_{\parallel}| = 530$ MHz, $g_{\perp} = 2.060$ for **2**. These parameters are in good agreement with those obtained for LPMOs.^{1,3,9} The redox potentials for the Cu(II)/Cu(I) couples in **1** and **2** were measured by cyclic voltammetry and redox titration. These potentials are found ranging from 5-50 mV *vs.* SHE depending on the method used and on the conditions (SI). These potential are much lower than those observed for the enzymatic systems, which range from 150-370 mV.^{1,9}

Scheme 1. Oxidative cleavage of *p*-nitrophenyl-β-D-glucopyranoside



The LPMO-like reactivity of the complexes was evaluated in the presence of hydrogen peroxide and *p*-nitrophenyl-β-D-glucopyranoside was used as a model substrate (Scheme 1). The formation of *p*-nitrophenolate was monitored under various conditions by spectrophotometry ($\lambda_{\max} = 400$ nm, $\epsilon = 18500$ mol⁻¹Lcm⁻¹).¹⁰ The initial velocities were found dependent on the concentrations of complexes and are up to 6-times higher than that obtained in the presence of a copper salt (Figure 2). The reaction rates were also dependent on hydrogen peroxide or substrate concentration (SI). Consequently, after 10 minutes of reaction, a 6-fold improvement in oxidation yield is obtained in the presence of the complexes as compared to CuSO₄ (0.1 mM of catalyst, see SI). After longer reaction times (24 h), up to 500 μM of *p*-nitrophenolate are produced using 10 μM of complexes (50 TON). In absence of either hydrogen peroxide or complex, low amount of *p*-nitrophenolate is detected (30-50 μM) suggesting that oxidative processes are involved rather than hydrolysis. In addition, gluconic acid (arising from gluconolactone) was evidenced by chromatographic analysis and ESI-MS analysis

in the reaction mixtures (**1** or **2** + H₂O₂) after 24 hours, thus strongly supporting oxidative cleavage similar to that catalyzed by LPMO (SI). Stepwise addition of hydrogen peroxide (15 mM every 2h, 5 additions) increases the reaction yield and 2.5 mM of *p*-nitrophenolate are detected after 24h (using 50 μM of catalyst). Each addition is followed by the generation of ca. 300 μM of *p*-nitrophenolate (6 TON) indicating that the reactive species can be regenerated by consecutive addition of hydrogen peroxide.

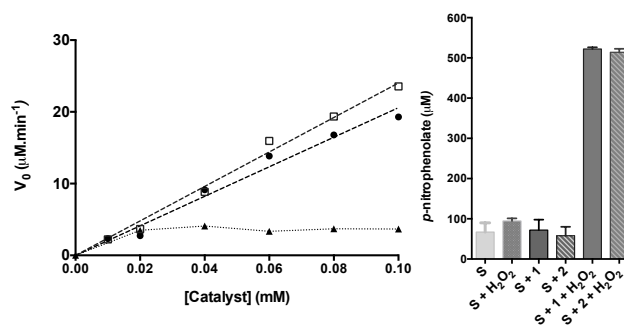


Figure 2. Reaction of *p*-nitrophenyl-β-D-glucopyranoside (S) with complexes **1** and **2** and H₂O₂ in carbonate buffer 100 mM pH 10.5 and at 30°C with [H₂O₂]=20 mM and [S] =20 mM. Left: initial velocities as a function of complex concentration for complexes **1** (●) and **2** (□) and for CuSO₄ (▲) Right: concentration of *p*-nitrophenolate obtained after 24 hours of reaction with [**1** or **2**]=0.01 mM.

Finally, the addition of H₂O₂ (in buffers) or H₂O₂/Et₃N (in water) on solutions of **1** or **2** is followed by changes in the UV-visible spectra and intense absorption bands centered at 305 nm with shoulders around 375 nm appear for both complexes (Figure 3). These species are stable from minutes to several hours depending on the conditions (concentrations, pH) and then slowly decay. The intermediates still display d-d transitions around 650 nm which suggests the maintenance of Cu(II) redox state. In the literature, addition of H₂O₂ in the presence of Et₃N on different copper complexes in organic solvents and at low temperature has led to the identification of several CuOOH intermediates displaying charge-transfer transitions in the 350-400 nm region.⁵ Yet, there are only few data on copper-oxygen intermediates formed in aqueous solutions and to the best of our knowledge only few bridged-peroxo intermediates have been observed.¹¹ In the present case, no significant decrease of EPR intensities is observed in different conditions and the intermediates display similar EPR spectra than the parent complexes (SI). It is therefore possible that the copper coordination sites of **1** or **2** are substituted with hydroperoxide ligand without altering significantly the copper coordination geometry, as already observed for some CuOOH species.¹² Correlation between formation/decay of the intermediates and production of *p*-nitrophenolate was unfortunately difficult due

to spectrophotometric overlap between the substrate and the intermediate (SI).

Density Functional Theory (DFT) calculations were undertaken to get deeper insight into the structure of the intermediate. The starting complex was subjected to geometry optimization and different structures were considered including replacement of acetonitrile by a water ligand (SI) and provided pentacoordinated copper centers (one perchlorate was decoordinated). The structure of 1^* ($[L^{\text{AM}}\text{Cu}(\text{OH}_2)(\text{ClO}_4)]^+$) was used as starting point to compute structures of the intermediate. Different mononuclear intermediates displaying peroxo and hydroperoxo ligands were considered (SI) and TDDFT calculations were carried out on their optimized structures to get insight into their optical properties. Comparison between the predicted and experimental spectra indicated that the best model consists in one hydroperoxo and one water bound to the copper, namely 1^*-OOH (SI). The calculations reproduce the experimental spectra of the initial complex and of the intermediate and in particular a ligand-to-metal CT in 1^* and a peroxo-to-metal CT in 1^*-OOH calculated at 278 and 331 nm respectively, that are found at 265 and 305 nm experimentally (the shift is also well reproduced). Finally, computed electronic structure and magnetic properties show that 1^* and 1^*-OOH display singly occupied molecular orbitals (SOMOs) with dominant Cu $3d_{x^2-y^2}$ character, in agreement with the computed and experimental EPR parameters (SI).

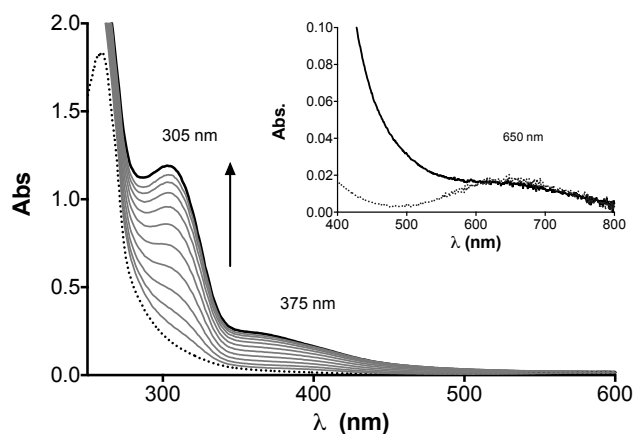


Figure 3. Absorption spectra of **1** at 0.25 mM in aqueous solution (----) upon addition of 25 mM of hydrogen peroxide at room temperature. Formation of an intermediate (—) over 15 minutes reaction time. Inset: zoom around the d-d transitions region.

Some CuOOH intermediates were shown to display oxidative activities.¹³ However, it was also suggested that high-valent copper-oxo species or radicals derived from hydroperoxo intermediates could be responsible for the observed reactivities.¹⁴ Therefore the role of CuOOH species in the reactiv-

ity of complexes **1** and **2** remains to be elucidated.⁸ Interestingly, copper complexes were recently shown to efficiently perform alkane oxidation in the presence of H₂O₂ through Fenton-like mechanisms.¹⁵ Although plausible with model complexes, Fenton mechanism would not be expected for the selective oxidations (C1 or C4) performed by LPMO enzymes. This work however sheds light on species that should be considered in discussions of mechanism of mononuclear copper-containing systems.

ASSOCIATED CONTENT

Supporting Information containing experimental, crystallographic, EPR, redox and calculation data is available free of charge on the ACS Publications website.

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Notes

[§]Dimethylpyrroline N-oxide (DMPO) was used as radical trapping agent. Similar quantities of DMPO-OH radicals were detected by EPR in the reaction conditions, in presence or in absence of complexes, rendering any conclusion difficult here.

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Two copper(II) complexes are described as models for the Lytic Polysaccharide Monooxygenases, copper-containing enzymes involved in oxidative cleavage of recalcitrant polysaccharides. These complexes are able to catalyze the oxidative cleavage of a model substrate, *p*-nitrophenyl- β -D-glucopyranoside, in the presence of hydrogen peroxide. An possible CuOOH intermediate is detected in the reaction conditions.

