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# Neutron and Atomic Resolution X-ray Structures of a Lytic Polysaccharide Monooxygenase Reveal Copper-Mediated Dioxygen Binding and Evidence for N-Terminal Deprotonation

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## Supporting Information

**ABSTRACT:** A 1.1 Å resolution, room-temperature X-ray structure and a 2.1 Å resolution neutron structure of a chitin-degrading lytic polysaccharide monooxygenase domain from the bacterium *Jonesia denitrificans* (JdLPMO10A) show a putative dioxygen species equatorially bound to the active site copper. Both structures show an elongated density for the dioxygen, most consistent with a Cu(II)-bound peroxide. The coordination environment is consistent with Cu(II). In the neutron and X-ray structures, difference maps reveal the N-terminal amino group, involved in copper coordination, is present as a mixed ND<sub>2</sub> and ND<sup>-</sup>, suggesting a role for the copper ion in shifting the pK<sub>a</sub> of the amino terminus.

Lytic polysaccharide monooxygenases (LPMOs) are copper-dependent enzymes found in bacteria and fungi, utilizing an oxidative mechanism to cleave polysaccharides.<sup>1,2</sup> This family of enzymes has wide-ranging applications in biofuel production, as they are able to degrade a variety of polysaccharides, notably crystalline cellulose. The enzymes utilize a single copper ion to activate oxygen. While the catalytic mechanism of LPMOs remains uncertain, it is thought that catalysis involves initial formation of a superoxide by electron transfer from the reduced copper ion.<sup>3–6</sup> The copper ion is in a conserved histidine brace motif, using the side chain imidazole groups of two histidines, one of which is the N-terminal amino acid, along with the N-terminus itself, to coordinate the copper.<sup>7</sup> A number of X-ray crystallographic structures are currently available for LPMOs from fungal and bacterial species.<sup>7–13</sup> Identifying the oxidation state of the catalytic copper ion and copper-bound oxygen species, the protonation states of active site residues, and copper-bound ligands may offer potential clues about the enzyme's function.

Neutron diffraction has the ability to experimentally determine hydrogen atom positions, as the coherent scattering

length of deuterium atoms, introduced into exchangeable moieties such as amides, amines, and hydroxyls, is 6.67 fm, comparable to those of other heavier elements in macromolecules like C, N, and O. This makes neutron diffraction a useful tool for determining protonation states and estimating pK<sub>a</sub> values of critical catalytic residues, and for unambiguously orienting hydrogen bonding donor and acceptor pairs and solvent molecules.<sup>14</sup> Furthermore, radiation damage is negligible, allowing for data sets to be collected at more physiological temperatures.

To probe the protonation states of active site residues and species bound to the copper, neutron diffraction data were collected to 2.1 Å resolution on a large crystal of a chitin-degrading LPMO domain from the Gram-positive bacterium *Jonesia denitrificans* (JdLPMO10A), which uses an oxidative mechanism to cleave at the C1 position in the scissile glycosidic bond.<sup>15</sup> Labile hydrogen atoms in the crystal were exchanged using D<sub>2</sub>O-containing mother liquor, to make them visible in nuclear density maps as deuteriums. The model was refined to R<sub>free</sub> and R<sub>cryst</sub> values of 26.5 and 18.7%, respectively, using the program Phenix (Table S1). The crystal used for the experiment was not exposed to X-rays, avoiding potential structural changes due to X-ray-induced photoreduction, the occurrence of which has been well-documented by other groups that have studied copper proteins.<sup>9,10</sup> Using a separate crystal grown under identical crystallization conditions, room-temperature X-ray diffraction data to 1.1 Å were collected at beamline 4.2.2 at the Advanced Light Source, with careful efforts to minimize radiation damage and the potential for photoreduction, as described previously.<sup>15</sup> The X-ray structure of JdLPMO10A was determined and refined to R<sub>free</sub> and R<sub>cryst</sub> values of 12.8 and 11.4%, respectively (Table S1), extending the resolution of our previous 1.55 Å structure collected at 100 K.<sup>8</sup>

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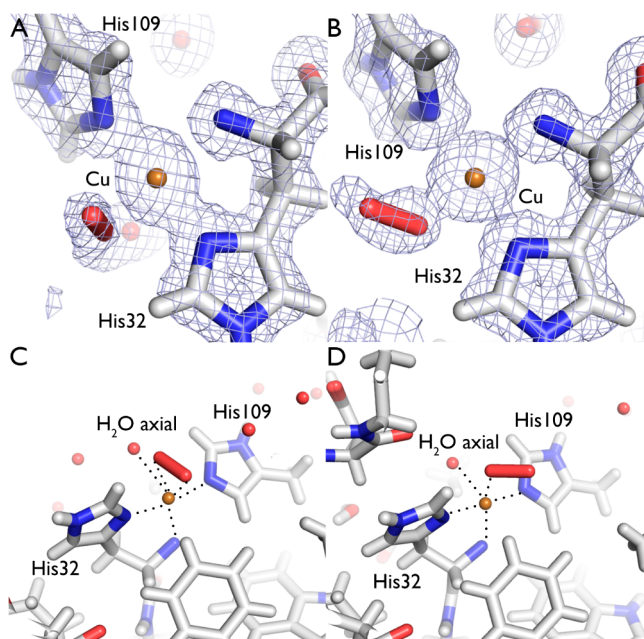
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Together, these data provide insights regarding the oxidation state of the catalytic copper ion, potential oxygen species bound to the enzyme, and the protonation state of residues near the copper site.

The protein crystallizes with two molecules occupying distinct microenvironments in the asymmetric unit. In the active site of the X-ray structure (Figure 1), the catalytic copper



**Figure 1.**  $2F_o - F_c$  electron density (lavender) for (A) molecule A and (B) molecule B of the 1.1 Å room-temperature X-ray structure, contoured at  $1.0\sigma$ , showing the active site coordination environment. The oxygen species coordinated to the copper ion has been modeled as peroxide with an O–O bond length of 1.5 Å. Active site copper coordination environment for (C) molecule A and (D) molecule B. Coordination distances are detailed in Table S2.

ion is coordinated by a so-called histidine brace found in all LPMOs that utilize the side chains of two conserved histidine residues, the N-terminal residue His32 and His109, and the N-terminal amino group of His32 (residues 1–31 comprise the signal sequence).<sup>7–9,11,13,16</sup> As shown in panels C and D of Figure 1, the copper coordination environments in the two monomers are similar, though not identical. The three nitrogen atoms form an equatorial plane. The final equatorial site in the X-ray structure is modeled as a dioxygen species, as confirmed by omit maps depicting strong  $F_o - F_c$  difference density when one of the oxygens is omitted from the dioxygen (Figure S1). The two molecules exhibit slightly different modes of oxygen binding. In molecule A, the dioxygen appears to be in a bidentate coordination environment, while in molecule B, the dioxygen is coordinated end-on (Table S2). The distal axial coordination site is occupied by water. Access to the proximal axial site is restricted by Phe164, as is commonly observed in LPMOs, which carry a Phe or a Tyr at this position.

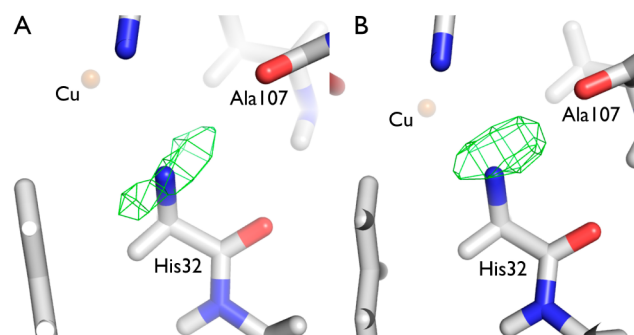
In the X-ray crystallographic structure, the refined coordination distances between the copper ion and the nitrogen ligands average to 2.0–2.2 Å (Figure 1C,D and Table S2). The square bipyramidal geometry and the presence of water/oxygen ligands suggest a Cu(II) oxidation state, consistent with other crystal structures of LPMO enzymes reported in the literature.<sup>9,10,17</sup> In an EXAFS and XANES study of a

tetrapeptide–copper complex, the Cu(II)–nitrogen distance is 1.9 Å and the Cu(III)–nitrogen distance is slightly shorter, 1.8 Å.<sup>18</sup> The coordination distances seen in the X-ray crystallographic structure do not support a Cu(III) state. The geometry of the coordination is also distinct from the T-shaped coordination seen in Cu(I) complexes.

## PROTONATION STATES IN THE ACTIVE SITE

The protonation states of active site residues were examined in the nuclear density maps. Nuclear density for the copper ion is weak, as has been observed for metals in other metalloenzymes such as carbonic anhydrase, xylose isomerase, and DFPase.<sup>19–21</sup>

Of particular interest is the protonation state of the amino terminus. The amino terminus in molecule A is observed as  $\text{ND}_2$  based on a difference  $F_o - F_c$  map calculated without the deuterium atoms, which shows a planar difference peak for the two deuteriums. Notably though, the N-terminus in molecule B shows an asymmetric  $F_o - F_c$  difference peak in the nuclear density maps, suggesting the presence of a mixed  $\text{ND}_2$  and  $\text{ND}^-$  species (Figure 2). This was further confirmed using nuclear  $F_o - F_c$  omit maps (Figure S2).

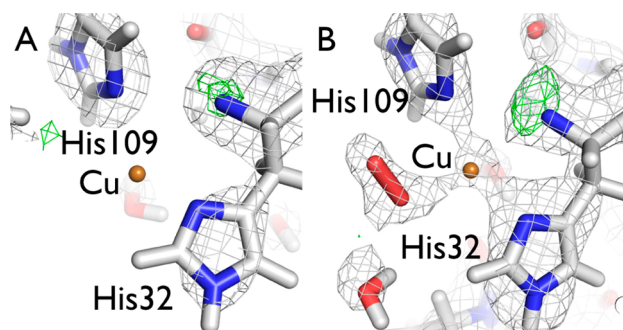


**Figure 2.** Active site of the two LPMO molecules in the asymmetric unit, showing  $F_o - F_c$  nuclear difference density (green) contoured at  $3.0\sigma$  corresponding to (A) a  $\text{ND}_2$  in molecule A and (B) a putative  $\text{ND}^-$  species in molecule B, with the deuterium pointing toward the carbonyl of Ala107.

To further examine this observation, the electron density around the amino terminus was re-examined in the room-temperature X-ray structure. The very high resolution (1.1 Å) is such that the  $F_o - F_c$  difference density for a limited number of hydrogen (or deuterium) atoms can be discerned. Although no difference density was seen in molecule A, on the basis of these difference maps, the protonation state of the N-terminal amino group in molecule B appears to be also a mixed  $\text{ND}_2$  and  $\text{ND}^-$ , similar to the neutron structure (Figure S3A). *JdLPMO10A* was crystallized at pH 7.0, and given the observation of mixed  $\text{ND}^-$  and  $\text{ND}_2$  species, the  $\text{pK}_a$  of the amino terminus is estimated to be approximately 7–7.5. Of note are electron density maps in the high resolution (1.37 Å) structure of *Neurospora crassa* LPMO9M (Protein Data Bank entry 4EIS),<sup>22</sup> crystallized at pH 8.5, which have the N-terminus modeled as an  $\text{NH}_2$ . One of the hydrogen atoms has a negative  $F_o - F_c$  peak, indicating the presence of an  $\text{NH}^-$  group (Figure S3B). It is likely that a similar protonation state for the N-terminus may be found in other members of the LPMO family, and this may be of significance to the reaction mechanism.

## OXYGEN SPECIES

In the neutron structure, one of the two molecules in the asymmetric unit shows what appears to be a dioxygen species. The density strongly suggests that the copper ion in molecule B coordinates a dioxygen species, although the resolution of the structure does not easily permit its unambiguous identification (Figure 3). The density appears to be too long for an O=O



**Figure 3.**  $2F_o - F_c$  composite omit nuclear density in the active site region (gray), contoured at  $1.5\sigma$ , for (A) molecule A and (B) molecule B.  $F_o - F_c$  difference density (green), contoured at  $3.0\sigma$ , is overlaid. No difference density is seen around the oxygen species that is present in molecule B.

(1.21 Å) or superoxo (1.33 Å) species and is most consistent with the bond length of a peroxide (1.49 Å). Nevertheless, the bond length of the dioxygen species may be influenced by binding to copper. The shape of the  $2F_o - F_c$  density and the lack of  $F_o - F_c$  density in the vicinity of the dioxygen suggest that the dioxygen species is not protonated, which likely excludes hydrogen peroxide (as DO-OD) or hydroperoxo (as  $^-\text{O}-\text{OD}$ ) as the bound species, as the D scatters neutrons strongly. Still, we cannot rule out flexibility in binding of the dioxygen species, which could obscure the strong scattering contribution from a bound deuterium. Interestingly, putative oxygen species have been observed previously in LPMOs at axial positions and farther from the copper.<sup>22,23</sup> Here, we show crystallographic evidence of a dioxygen species in an equatorial position that is clearly interacting with the copper. The position of this species is compatible with the recently determined X-ray structure of an LPMO in complex with a cello-oligosaccharide and the mechanistic inferences made from the structure, as well as recent crystallographic data regarding an LPMO from *N. crassa*, discussed in more detail below.<sup>11,23</sup>

## FUNCTIONAL IMPLICATIONS

The bracing N-terminal group contains mixed  $\text{ND}_2$  and  $\text{ND}^-$  as observed in both the X-ray and neutron structures. The  $\text{pK}_a$  values for copper-coordinated amines in model  $\text{Cu(III)}$  complexes have been reported to range from 8 to 10.<sup>24</sup> We have not found reports for  $\text{pK}_a$ 's of  $\text{Cu(II)}$ -coordinated amines. The  $\text{pK}_a$  values for  $\text{Cu(II)}$ -coordinated amides have been reported to be between 7.8 and 8.8, illustrating that coordination of nitrogen by copper can significantly lower the  $\text{pK}_a$ , in the case of amides, to near-physiological pH values.<sup>25</sup> Given that the protein was crystallized at pH 7.0, well within the pH range of enzymatic activity for bacterial LPMOs (e.g., refs 1, 26, and 27), the  $\text{pK}_a$  for deprotonation of the bracing amine ( $-\text{NH}_2$ ), may be in the neutral pH range. As noted by Quinlan et al. and Walton and Davies,<sup>3</sup> a deprotonated amino terminus could stabilize a potential  $\text{Cu(III)-OH}$  intermediate.

The observation of mixed  $\text{ND}_2$  and  $\text{ND}^-$  in the two molecules in the crystal asymmetric unit is plausible, given their different crystal packing environments.

A set of ultra-high resolution X-ray and neutron structures of *NcLPMO9D*, an LPMO from the fungus *N. crassa* (also known as NCU01050 and PMO-2), crystallized at pH 5.6, was recently reported.<sup>23</sup> While *NcLPMO9D* and *JdLPMO10A* share a similar  $\beta$ -sandwich inner core fold and active site environment, they exhibit low levels of sequence ( $\sim 10\%$  identical) and structural similarity and have different preferred substrates.<sup>17</sup> *JdLPMO10A* cleaves glycosidic bonds by oxidizing C1 of the  $\beta 1-4$  linkage in chitin, while *NcLPMO9D* oxidizes C4 of the  $\beta 1-4$  linkage in cellulose. *NcLPMO9D* contains an axial tyrosine that coordinates the catalytic copper, albeit with an elongated  $\text{Cu-O}$  separation, while in *JdLPMO10A*, the tyrosine position is occupied by a phenylalanine that does not coordinate the copper. From the neutron structure of *NcLPMO9D*, it was suggested that a neutral or protonated histidine residue (His157) in the vicinity of the active site, distinct from the histidine residues coordinating the catalytic copper, may be critical in the reaction mechanism by interacting with and preorganizing  $\text{O}_2$  prior to it entering the copper center. Notably, there is no analogous residue in *JdLPMO10A* or other chitin-active LPMO10s,<sup>16</sup> whereas a similar histidine occurs in some, but far from all, cellulose-active LPMO10s. Still, most importantly, the X-ray structural studies of ascorbate-treated *NcLPMO9D* crystals revealed an equatorially bound dioxygen in one of the molecules in the asymmetric unit, which was modeled as a peroxo species directly coordinating the catalytic copper ion, adopting a similar end-on binding configuration (Table S2).<sup>23</sup>

While many of the details of the LPMO mechanism remain to be resolved, the data presented here and in other recent studies<sup>11,23</sup> strongly indicate that catalysis involves equatorial binding of a reactive oxygen species, as anticipated by Kjærgaard et al.<sup>4</sup> Importantly, while LPMOs show large sequence variation and even display considerable differences in the axial copper coordination positions,<sup>9,17,28</sup> this study of a bacterial LPMO and other recent studies that include fungal LPMOs suggest a high degree of mechanistic similarity.

## ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.biochem.7b00019.

Materials and Methods, Tables S1 and S2, and Figures S1–S3 (PDF)

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The authors declare no competing financial interest.

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