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Incorporation of copper ions into crystals of T2 copper-depleted laccase from *Botrytis aclada*

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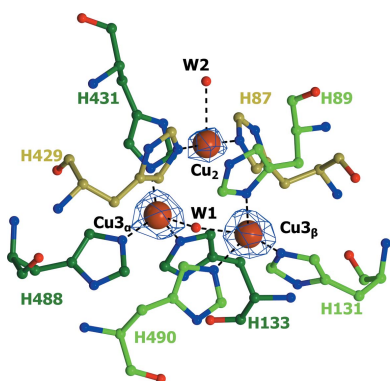
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Laccases belong to the class of multicopper oxidases catalyzing the oxidation of phenols accompanied by the reduction of molecular oxygen to water without the formation of hydrogen peroxide. The activity of laccases depends on the number of Cu atoms per enzyme molecule. The structure of type 2 copper-depleted laccase from *Botrytis aclada* has been solved previously. With the aim of obtaining the structure of the native form of the enzyme, crystals of the depleted laccase were soaked in Cu⁺- and Cu²⁺-containing solutions. Copper ions were found to be incorporated into the active site only when Cu⁺ was used. A comparative analysis of the native and depleted forms of the enzymes was performed.

1. Introduction

Laccases (EC 1.10.3.2; benzenediol:oxygen oxidoreductases) belong to the class of multicopper oxidases catalyzing the oxidation of phenols accompanied by the reduction of molecular oxygen to water. Laccases are composed of two or three structurally similar cupredoxin-like domains. These enzymes have been well studied both by biochemical (Baldrian, 2006) and structural (Hakulinen & Rouvinen, 2015) methods. Owing to their broad substrate specificity and the formation of non-toxic products, laccases are of interest in biotechnology (Xu, 2002).

The active site of laccases comprises four Cu atoms classified into three types depending on their spectroscopic characteristics (Solomon *et al.*, 1996). The type 1 copper ion can be detected in the ESR spectrum and gives a characteristic absorption at around 600 nm. This ion has a trigonal planar coordination through the N^ε atoms of two histidines and the S atom of one cysteine. Type 2 and type 3 copper ions form a T2/T3 cluster responsible for the reduction of molecular oxygen to water. The type 2 copper ion is detected in the ESR spectrum and does not show absorption in the optical region. Two type 3 copper ions are antiferromagnetically coupled through an oxygen-containing ligand and consequently do not exhibit an ESR signal, but show an absorption maximum at around 330 nm. The type 2 copper ion adopts a square-planar coordination geometry with two histidines and one or two oxygen-containing ligands. Each type 3 copper ion is coordinated in a tetrahedral geometry by three histidines and one oxygen-containing ligand, through which it is linked to another type 3 copper ion.



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Table 1
Macromolecule-production information.

Source organism	<i>B. aclada</i>
Forward primer 5BAPm1	5'-ATACACGTGCAAGATGAAGTATTTACAGTC-TTACTGC-3'
Reverse primer 3BAXba1	5'-ATATCTAGACTTAAATCCAGAATCGTCTCC-3'
Expression vector	pPICZB and pPICZ α A
Expression host	<i>P. pastoris</i>
Complete amino-acid sequence of the construct produced	<p> MKYFTVFTALTALFAQASASAIPAVRSTLTPRQN- TTASCANSATSRSCWGEYSIDTNWYDVTPTGV- TREYWLSVENSTITPDGYTRSAMTFNGTVPGP- AIIADWGDNLIHVTNNLEHNGTSHWHGIRQ- LGSLEYDGVPGVTQCPIAPGDTLTYKFQVTQY- GTTWYHSHFSLQYGDGLFGPLIINGPATADYD- EDVGVIFLQDWAHESVFEIWDARLGAPPALE- NTLMNGTNTFDCSASTDPNCVGGGKFKELTFV- EGTKYRLRLINVGIDSHFEFAIDNHTLTVIAN- DLVPIVPTYTDTLLIGIGQRVDVIVEANAAD- NYWIRGNWGTTCSTNNEANATGILRYDSSSI- ANPTSVGTTPRGTCEDEPVASLVPHLALDVGG- YSLVDEQVSSAFTNYFTWTINSSLLLDWSSP- TTLKIFNNETIFPTEYVVALEQTNANEWV- VYIEDLTGFGIWHPIHLHGDFIVAQETDVF- NSDEPAKFNLVNPPRRDVAALPGNGYLATAF- KLDNPGSWLLHCHIAWHASEGLAMQFVESQSS- IAVKMTDTAIFEDTCANWNAYTPTQLFAEDDS- GI </p>

The activity of laccase preparations depends on the number of Cu atoms per enzyme molecule. Thus, one way of investigating laccases is to study enzyme preparations in which the copper ions have been partially removed. The type 2 copper ion is more easily removed from the laccase molecule compared with the other copper ions (Malkin *et al.*, 1969). On the one hand, procedures have been developed for the preparation of inactive type 2 copper-depleted forms of the enzyme based on treatment with copper chelators in the presence of reducing agents (Reinhammar & Oda, 1979; Koroleva *et al.*, 2001). On the other hand, the partial loss of Cu atoms from the active site can occur spontaneously during the production and storage of laccases.

In some X-ray diffraction structures of laccases which had not been subjected to a copper-chelation procedure, the type 2 copper ion was either absent or had partial occupancy (Glazunova *et al.*, 2015; Ducros *et al.*, 1998; Osipov *et al.*, 2014). The restoration of depleted forms of laccases with a simultaneous increase in their activity is accomplished by treatment of the enzyme with copper salts. The insertion of a type 2 copper ion into fungal, plant and bacterial laccases using Cu⁺ ions has been described in the literature (Malkin *et al.*, 1969; Reinhammar & Oda, 1979; Koroleva *et al.*, 2001). The X-ray crystal structure of the fungal type 2 copper-depleted laccase from *Corioliopsis caperata* showed that a copper ion was inserted into the active site only when Cu⁺ ions were used, whereas the use of Cu²⁺ ions did not give the desired result (Glazunova *et al.*, 2015). However, it was demonstrated that the incorporation of a type 2 copper ion into the bacterial laccase from *Bacillus subtilis* occurs in the presence of either Cu⁺ or Cu²⁺ ions (Durão *et al.*, 2008).

The laccase from the ascomycete *Botrytis aclada* has previously been isolated and characterized biochemically (Kittl *et al.*, 2012). The structures of this enzyme (hereafter

Table 2
Crystallization.

Method	Vapour diffusion
Plate type	VDX
Temperature (K)	298
Protein concentration (mg ml ⁻¹)	20
Buffer composition of protein solution	25 mM sodium acetate pH 5.0
Composition of reservoir solution	1.8 M ammonium sulfate
Volume and ratio of drop	2 μ l, 1:1
Volume of reservoir	500 μ l

referred to as T2D) and its L499M mutant were determined at 1.7 Å resolution (Osipov *et al.*, 2014). Although the laccase was not subjected to a depletion procedure, the type 2 copper ion was absent in both structures and the residue His429, *i.e.* one of the two histidines that are involved in the coordination of the type 2 copper ion, points towards Cu_{3 α} .

The aim of the study presented here is to determine the structure of *B. aclada* laccase containing the complete set of copper ions. For this purpose, crystals of the type 2 copper-depleted form of *B. aclada* laccase were soaked in solutions containing Cu⁺ or Cu²⁺ ions. Restoration of the native form of the enzyme was only observed in the experiment employing Cu⁺ ions.

2. Materials and methods

2.1. Purification and crystallization of laccase

Laccase from *B. aclada* (Table 1) was recombinantly expressed in the yeast *Pichia pastoris* and purified as described by Kittl *et al.* (2012). Attempts to obtain crystals of the native form of laccase failed. Therefore, the enzyme was deglycosylated (Osipov *et al.*, 2014). The crystallization conditions have been described in detail in a previous study (Osipov *et al.*, 2014). Crystals were grown by the vapour-diffusion technique. A 1.8 M solution of ammonium sulfate in water was used as the reservoir solution. The protein solution consisted of 20 mg ml⁻¹ protein in 25 mM sodium acetate buffer pH 5.0. A 2 μ l drop composed of equal volumes of the protein and reservoir solutions was used (Table 2). Crystals appeared within 3 d and reached maximum dimensions of 0.2 \times 0.1 \times 0.05 mm in one month.

2.2. Preparation of complexes of T2D with Cu⁺ and Cu²⁺

The complexes were obtained by soaking crystals of the enzyme in reservoir solution containing 0.6 mM (saturated solution) CuCl (T2D+Cu⁺ complex) or 10 mM CuSO₄ (T2D+Cu²⁺ complex). The soaking times were 10 min and 1 d, respectively.

2.3. Data collection and processing

X-ray data sets for T2D+Cu⁺ and T2D+Cu²⁺ were collected on the K4.4e beamline at the Belok station at the Kurchatov synchrotron-radiation source at 100 K under a nitrogen flow at a wavelength of 0.98 Å using a Rayonix SX165 detector. Before X-ray data collection, the crystals were placed for 5 s in reservoir solution supplemented with 20%(v/v) glycerol.

Table 3
Data collection and processing.

Values in parentheses are for the outer shell.

Data set	T2D+Cu ⁺	T2D+Cu ²⁺
Beamline	Beamline K4.4e, Kurchatov SNC	
Wavelength (Å)	0.98	
Temperature (K)	100	
Detector	Rayonix SX165 CCD	
Rotation range per image (°)	1.0	
Total rotation range (°)	154.0	119.0
Space group	C2	
<i>a</i> , <i>b</i> , <i>c</i> (Å)	69.4, 113.1, 79.6	70.1, 113.9, 80.0
α , β , γ (°)	90, 109.0, 90	90, 108.8, 90
Mosaicity (°)	0.37	0.26
Resolution range (Å)	30–2.30 (2.44–2.30)	30–1.83 (1.94–1.83)
Total No. of reflections	81583 (12203)	124585 (16545)
No. of unique reflections	25106 (3913)	50252 (7615)
Completeness (%)	96.7 (94.6)	96.1 (90.5)
Multiplicity	3.3 (3.1)	2.5 (2.2)
$\langle I/\sigma(I) \rangle$	16.3 (2.3)	20.8 (5.1)
<i>R</i> _{meas}	0.071 (0.60)	0.039 (0.23)
Overall <i>B</i> factor from Wilson plot (Å ²)	37.7	24.4

The X-ray data sets were processed using the *XDS* package (Kabsch, 2010*a,b*). The data-collection statistics are summarized in Table 3.

2.4. Structure solution and refinement

The crystals of the complexes were isomorphous to the crystals of T2D. The structures were refined with *REFMAC5* (Murshudov *et al.*, 2011). All crystallographic calculations were carried out using the *CCP4* suite (Winn *et al.*, 2011). During the refinement of the copper ions with full occupancy, the *F*_o – *F*_c difference map contained essential peaks in the region of the copper ions and the *B* factors of the copper ions exceeded the *B* factors of the ligands. Therefore, in the final stages of refinement the occupancies of the copper ions were refined manually. Manual correction of the occupancies was followed by *B*-factor refinement. In the final model, the *B* factors of the copper ions were approximately equal to the *B* factors of the ligands, and the *F*_o – *F*_c difference map did not contain essential peaks in the region of the copper ions. Visual inspection and manual rebuilding of the models was carried out using the *Coot* interactive graphics program (Emsley *et al.*, 2010). Water molecules were manually added to the structures based on analysis of the difference electron-density maps. The quality of the structures was evaluated with *SFCHECK* (Vaguine *et al.*, 1999) and *PROCHECK* (Laskowski *et al.*, 1993). The figures were drawn with *CCP4mg* (McNicholas *et al.*, 2011). *MolProbity* (Chen *et al.*, 2010) was used for Ramachandran analysis. Structure-solution and refinement statistics are summarized in Table 4.

3. Results and discussion

The structure of T2D+Cu²⁺ was solved at 1.8 Å resolution. The crystals of T2D+Cu²⁺ have the same qualities as the initial crystals of T2D. Since the structure of T2D+Cu²⁺ is almost identical to that of T2D, it was not deposited in the PDB.

Table 4
Structure solution and refinement.

Values in parentheses are for the outer shell.

Data set	T2D+Cu ⁺	T2D+Cu ²⁺
Resolution range (Å)	30–2.30 (2.44–2.30)	30–1.83 (1.94–1.83)
Completeness (%)	97.2	96.3
No. of reflections, working set	23801 (1706)	47673 (3062)
No. of reflections, test set	1292 (111)	2579 (187)
Final <i>R</i> _{cryst}	0.181 (0.278)	0.162 (0.241)
Final <i>R</i> _{free}	0.233 (0.347)	0.204 (0.302)
No. of non-H atoms		
Protein	4087	4208
Ion	5	3
Water	155	429
Other	197	197
Total	4444	4837
R.m.s. deviations		
Bonds (Å)	0.011	0.016
Angles (°)	1.62	1.92
Average <i>B</i> factors (Å ²)		
Protein	38.6	23.3
Ion	38.3	21.2
Water	35.8	29.8
Other	58.9	37.7
Ramachandran plot		
Favoured (%)	96.2	96.8
Allowed (%)	3.6	3.2

The structures of T2D and T2D+Cu²⁺ superimposed with an r.m.s.d. of 0.09 Å using the coordinates of 539 equivalent C^α atoms. Soaking T2D crystals for 24 h in a solution containing Cu²⁺ ions at high concentrations did not lead to the insertion of a copper ion into the T2/T3 cluster.

The complexes were obtained using crystals grown in the same drop. During the preparation of the T2D+Cu⁺ crystals the quality of the crystals visually deteriorated despite the low concentration of Cu⁺ ions and the short period of soaking. The structure of T2D+Cu⁺ was solved at 2.3 Å resolution. The reduction in the diffraction limit was accompanied by an increase in the mosaicity (0.38° for the crystals of T2D+Cu⁺ versus 0.25° for the crystals of T2D). Superimposition of the T2D+Cu⁺ and T2D structures using the coordinates of 525 equivalent C^α atoms gave an r.m.s.d. of 0.16 Å. Residues 1–14 (the numbering given is according to the T2D structure) and residues 405–408 were not located in the electron-density map for T2D+Cu⁺. The latter residues are also not observed in the electron-density maps for T2D and T2D+Cu²⁺.

The main differences in the structures of T2D+Cu⁺ and T2D are related to the location of a copper ion in the type 2 site and the orientation of the side chain of His429 (Fig. 1). It should be noted that in the structure of T2D Cu2 is completely absent and the sum of occupancies of the copper ions in T2D (2.6) is almost equal to the content of copper ions per molecule for the enzyme in solution as determined by mass spectrometry (Osipov *et al.*, 2014). In the T2D+Cu⁺ structure the type 2 copper ion has a square-planar coordination formed by the His87 N^ε and His429 N^ε atoms and the O atom of a water molecule (Table 5). This situation is typical for all laccases containing type 2 copper ions. In the T2D structure His429 is coordinated to Cu_{3α} by the N^δ atom (Osipov *et al.*, 2014). Thus, the removal of type 2 copper ions could be associated

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