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# X-ray-induced catalytic active-site reduction of a multicopper oxidase: structural insights into the proton-relay mechanism and O<sub>2</sub>-reduction states

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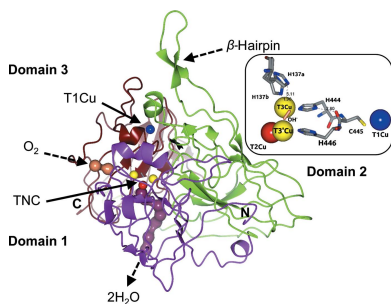
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During X-ray data collection from a multicopper oxidase (MCO) crystal, electrons and protons are mainly released into the system by the radiolysis of water molecules, leading to the X-ray-induced reduction of O<sub>2</sub> to 2H<sub>2</sub>O at the trinuclear copper cluster (TNC) of the enzyme. In this work, 12 crystallographic structures of *Thermus thermophilus* HB27 multicopper oxidase (*Tth*-MCO) in holo, apo and Hg-bound forms and with different X-ray absorbed doses have been determined. In holo *Tth*-MCO structures with four Cu atoms, the proton-donor residue Glu451 involved in O<sub>2</sub> reduction was found in a double conformation: Glu451a (~7 Å from the TNC) and Glu451b (~4.5 Å from the TNC). A positive peak of electron density above 3.5σ in an F<sub>o</sub> - F<sub>c</sub> map for Glu451a O<sup>ε2</sup> indicates the presence of a carboxyl functional group at the side chain, while its significant absence in Glu451b strongly suggests a carboxylate functional group. In contrast, for apo *Tth*-MCO and in Hg-bound structures neither the positive peak nor double conformations were observed. Together, these observations provide the first structural evidence for a proton-relay mechanism in the MCO family and also support previous studies indicating that Asp106 does not provide protons for this mechanism. In addition, eight composite structures (*Tth*-MCO-C1-8) with different X-ray-absorbed doses allowed the observation of different O<sub>2</sub>-reduction states, and a total depletion of T2Cu at doses higher than 0.2 MGy showed the high susceptibility of this Cu atom to radiation damage, highlighting the importance of taking radiation effects into account in biochemical interpretations of an MCO structure.

## 1. Introduction

Multicopper oxidases (MCOs) are a diverse family of metallo-enzymes that are widely distributed in all kingdoms, with bacterial and fungal laccases being the most numerous members (Hoegger *et al.*, 2006; Sharma *et al.*, 2007). MCOs couple four one-electron oxidations of several substrates to the four-electron reduction of O<sub>2</sub> to 2H<sub>2</sub>O using at least four Cu atoms distributed in two active sites: one type 1 copper (T1Cu) site, where an organic and/or inorganic substrate is oxidized, and a trinuclear copper cluster (TNC), where O<sub>2</sub> is bound, activated and reduced to 2H<sub>2</sub>O (Solomon *et al.*, 2001). The TNC consists of one type 2 copper (T2Cu) site and a type 3 binuclear T3Cu-T3'Cu cluster (Fig. 1; Solomon *et al.*, 1996). T1Cu has a typical electron paramagnetic resonance (EPR) signal and also forms a covalent S-Cu bond with an intense SCys→Cu<sup>II</sup> charge-transfer band at around 610 nm, which provides the distinctive blue colour of MCOs. T2Cu also has a distinctive EPR signal but it does not have any noticeable

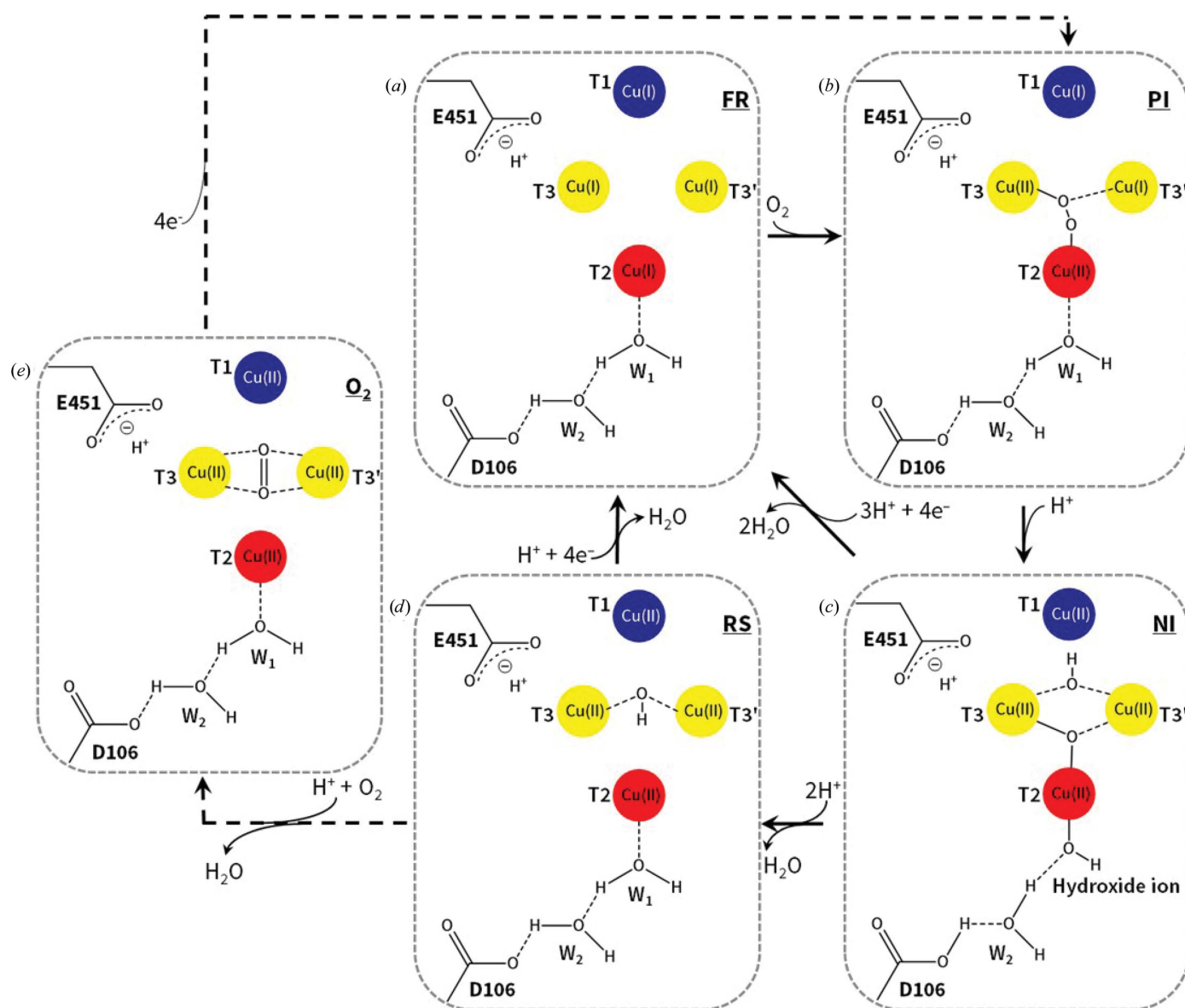


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features in the UV–visible region, while the binuclear T3Cu–T3′Cu cluster does not have an EPR signal but has an absorption band at around 330 nm (Solomon *et al.*, 2008). The binuclear T3Cu–T3′Cu cluster is coordinated by three His residues on each of the two Cu atoms, T2Cu is coordinated by two His residues and T1Cu is coordinated by two His residues and one Cys residue (Sharma *et al.*, 2007; Zhukhlistova *et al.*, 2008).

Four protons and four electrons generated by T1Cu from four substrate molecules are required for the reduction of O<sub>2</sub> at the TNC (Solomon *et al.*, 2008). Particularly, after substrate oxidation occurs at the T1Cu site, the electrons are shuttled over a distance of ~13 Å through a T1Cu<sup>II</sup>–Cys–His–T3Cu<sup>II</sup> electron-transfer pathway to the TNC (Augustine *et al.*, 2010). However, O<sub>2</sub> reduction at the TNC could also be observed in

the crystalline state without substrate oxidation, since electrons and protons are released into the crystal by the radiolysis of water molecules during X-ray data collection (Hakulinen *et al.*, 2006; Macedo *et al.*, 2009; Garman, 2010). Thus, the final crystallographic structure of an MCO is an average model of different Cu oxidation states and X-ray-induced O<sub>2</sub>-reduction states and intermediates (Hakulinen *et al.*, 2006; Ferraroni *et al.*, 2012). Accordingly, the structure of *Melanocarpus albo-myces* laccase at high absorbed X-ray dose was found to be in the resting state (RS; Fig. 1d) with the binuclear T3Cu–T3′Cu cluster coupled through a bridging hydroxide (OH<sup>−</sup>) ligand, while at low absorbed X-ray dose the structure was found with an almost symmetrically coordinated O<sub>2</sub> amidst the binuclear T3Cu–T3′Cu cluster (Fig. 1e; Hakulinen *et al.*, 2006). Since an MCO crystal is usually exposed from low to high absorbed



**Figure 1**

Schematic representation of the catalytic mechanism of O<sub>2</sub> reduction to 2H<sub>2</sub>O by MCOs. (a) FR state. (b) PI. (c) NI. (d) RS. (e) O<sub>2</sub> state. Note that the Glu451/Asp106 residues belong to *Th*-MCO. W<sub>1</sub> and W<sub>2</sub> are water molecules. The same colour code for the four Cu atoms is used in all other figures. Adapted from Bento *et al.* (2005), Quintanar *et al.* (2005) and Augustine *et al.* (2010).

radiation dose during a classic X-ray diffraction data acquisition, the latter observations revealed the reduction of O<sub>2</sub> in the crystalline state. Remarkably, in order to systematically study such catalytic reactions of metalloenzymes in the course of X-ray measurements, a multicrystal data-collection strategy based on a systematic diffusion of absorbed dose over a set of crystals has been designed (Berglund *et al.*, 2002). Using this technique of data combination from different crystals and the generation of composite structures with different absorbed X-ray doses, the catalytic mechanism of X-ray-induced reduction of O<sub>2</sub> in horseradish peroxidase has been described (Berglund *et al.*, 2002).

The reaction mechanism for the reduction of O<sub>2</sub> to 2H<sub>2</sub>O at the TNC has been extensively studied and involves two two-electron transfers (Fig. 1) starting from the fully Cu reduced (FR) enzyme with four catalytic Cu ions in the cuprous state (Fig. 1*a*; Augustine *et al.*, 2010). The first two-electron reduction from O<sub>2</sub> to the peroxide intermediate (PI) is rate-limiting (Figs. 1*a* and 1*b*), while the conversion of the PI to the native intermediate (NI) is extremely fast in the holoenzyme (Figs. 1*b* and 1*c*; Lee, George *et al.*, 2002; Palmer *et al.*, 2002; Yoon *et al.*, 2007; Yoon & Solomon, 2007). It has previously been demonstrated that protons are not involved in all steps of the reduction of O<sub>2</sub> at the TNC (Augustine *et al.*, 2007). For instance, no protons are involved in the reductive cleavage of the molecular oxygen O=O bond from O<sub>2</sub> to give the PI (Figs. 1*a* and 1*b*; Augustine *et al.*, 2007). However, there is one proton involved in the conversion of the PI to the NI, and therefore in the reductive cleavage of the peroxide <sup>-</sup>O—O<sup>-</sup> bond (Figs. 1*b* and 1*c*; Palmer *et al.*, 2002). Structurally, one O atom of the PI is coordinated to T3Cu and another O atom is coordinated to T2Cu, while in the NI one O atom is in the middle of the TNC and another O atom (OH<sup>-</sup>) is symmetrically located between the binuclear T3Cu–T3'Cu cluster (Augustine *et al.*, 2010). Thus, after the two two-electron transfers have taken place, the NI displays four catalytic Cu ions in the cupric state and forms the RS by releasing one H<sub>2</sub>O molecule (Fig. 1*d*; Yoon *et al.*, 2007). As mentioned above, the RS presents an OH<sup>-</sup> ligand bridging the binuclear T3Cu–T3'Cu cluster as well as four catalytic Cu ions in the cupric state (Quintanar *et al.*, 2005). However, the decay of the NI to the RS is slower than the turnover rate of the enzyme and therefore is not catalytically relevant (Yoon *et al.*, 2007). In fact, under catalytic conditions the cycle is completed upon reduction of the NI by a total of four electrons and the release of 2H<sub>2</sub>O molecules by the addition of three protons, regenerating the FR enzyme for the next enzyme cycle without decaying to the RS. Nevertheless, because of the lack of sufficient electrons to form the FR enzyme, the NI necessarily decays to the RS (Augustine *et al.*, 2010). Indeed, the RS is rather common in a large number of crystallographic structures of MCOs deposited in the PDB (~15 MCO structures), since the catalytic cycle for O<sub>2</sub> reduction in the crystalline state using the protons and electrons released by X-rays is rather inefficient (Hakulinen *et al.*, 2006; Kjaergaard *et al.*, 2012). Thus, since the FR enzyme is the first state of the catalytic cycle for O<sub>2</sub> reduction of MCOs (Augustine *et al.*, 2010), and

reacts immediately with O<sub>2</sub> to form the PI, the structural stabilization of the O<sub>2</sub> state (Fig. 1*e*) observed in the structure of *M. albomyces* laccase at low absorbed X-ray dose (Hakulinen *et al.*, 2006) seems to be a process that is exclusive to the crystalline state as a result of the deficiency of electrons to generate the FR enzyme at low doses.

MCOs have a pair of highly conserved acidic residues in the second coordination sphere of the TNC which are involved in the proton-relay mechanism for O<sub>2</sub> reduction mentioned above (Quintanar *et al.*, 2005; Augustine *et al.*, 2007; Kataoka *et al.*, 2009; Bento *et al.*, 2010; Chen, Durão *et al.*, 2010); specifically, a Glu residue next to T3Cu or in the middle of the TNC in the entry channel for O<sub>2</sub> and an Asp residue close to T2Cu in the exit channel for the water molecules produced. The pair of Glu/Asp residues in CotA from *Bacillus subtilis* (Bento *et al.*, 2010; Chen, Durão *et al.*, 2010), CueO from *Escherichia coli* (Kataoka *et al.*, 2009) and Fet3p from *Saccharomyces cerevisiae* (Quintanar *et al.*, 2005; Augustine *et al.*, 2007) are Glu498/Asp116, Glu506/Asp112 and Glu487/Asp94, respectively. As expected, *Thermus thermophilus* HB27 multicopper oxidase (*Tth*-MCO; Miyazaki, 2005; Serrano-Posada *et al.*, 2011) also possesses these two conserved acidic residues: Glu451/Asp106 (Fig. 1). It has been strongly suggested that Glu498 in CotA, Glu506 in CueO and Glu487 in Fet3p is the only proton-donor residue that plays a crucial role in the protonation events at the TNC, channelling the four protons required for O<sub>2</sub> reduction. In contrast, Asp116 in CotA, Asp112 in CueO and Asp94 in Fet3p is not a proton-donor residue but is important in the decay of the PI to the NI by the stabilization of both an H<sub>2</sub>O molecule (W<sub>1</sub>; Fig. 1) and an OH<sup>-</sup> ion (Fig. 1*c*), which are coordinated to T2Cu in the PI and the NI, respectively (Quintanar *et al.*, 2005; Augustine *et al.*, 2007; Kataoka *et al.*, 2009; Bento *et al.*, 2010; Chen, Durão *et al.*, 2010; Silva, Damas *et al.*, 2012). In fact, both H<sub>2</sub>O/OH<sup>-</sup> species interact in each intermediate with the side chain of the Asp residue through a structural water molecule (W<sub>2</sub>; Fig. 1). It has been shown that the absence of these two Glu/Asp residues, as in several mutants of CotA (Chen, Durão *et al.*, 2010; Silva, Damas *et al.*, 2012), CueO (Kataoka *et al.*, 2009) and Fet3p (Quintanar *et al.*, 2005; Augustine *et al.*, 2007), results in a severe catalytic impairment of O<sub>2</sub> reduction, reinforcing their importance in the proton-relay mechanism. Despite extensive study of the reduction of O<sub>2</sub> at the TNC in MCOs, the structural details of the proton-relay mechanism involved in this catalytic reaction have not currently been described.

*Tth*-MCO is a thermophilic metalloenzyme which exhibits oxidase activity towards laccase substrates such as guaiacol and ABTS, with an optimal temperature of 365 K for ABTS oxidation and a half-time of thermal inactivation of over 14 h at 353 K (Miyazaki, 2005). In this work, 12 different crystallographic structures of *Tth*-MCO have been determined in order to study the X-ray-induced catalytic active-site reduction of the enzyme and the role of the Glu451/Asp106 residues in the proton-relay mechanism. Firstly, the *Tth*-MCO structure at 1.5 Å resolution and three inactive forms of the enzyme at 1.7 Å resolution, apo *Tth*-MCO, Hg-*Tth*-MCO and

Hg-*Tth*-MCO-2h, were determined using classic data acquisition from a single crystal. Secondly, eight composite structures (*Tth*-MCO-C1–8) at 1.8 Å resolution with different absorbed X-ray doses were determined using a multicrystal data-collection strategy. In the latter case, different states of the X-ray-induced reduction of O<sub>2</sub> were trapped. The noticeably different behaviour of the Glu451/Asp106 residues in active and inactive forms of *Tth*-MCO provided structural details of the proton-relay mechanism and indicated that they act together, providing part of the necessary driving force to reduce O<sub>2</sub>. Moreover, the four catalytic Cu atoms of *Tth*-MCO revealed uneven radiation damage by absorbed X-rays, where T2Cu is completely depleted at doses of >0.2 MGy.

## 2. Experimental procedures

All chemical supplies were analytical grade and were purchased from Sigma–Aldrich unless stated otherwise.

### 2.1. Gene cloning, protein expression, purification and crystallization

Gene cloning, protein expression, protein purification and crystallization of *Tth*-MCO and apo *Tth*-MCO from *T. thermophilus* HB27 were performed as described previously (Serrano-Posada *et al.*, 2011). The purified recombinant *Tth*-MCO was the mature form of the enzyme, a 439-residue protein without the first 22 residues (Met1–Ala22), which are part of the signal peptide (Miyazaki, 2005), and exhibited the typical blue colour of MCOs. Additionally, two inactive Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h forms of *Tth*-MCO with Hg instead of Cu were obtained at 278 K by soaking two apo *Tth*-MCO crystals with 5 mM mercury(II) chloride for 5 min and 2 h, respectively. For the multicrystal data-collection strategy, eight different *Tth*-MCO crystals of approximate dimensions  $\sim 0.025 \times 0.05 \times 0.1$  mm were grown at 278 K by the hanging-drop vapour-diffusion method using the microseeding technique (Serrano-Posada *et al.*, 2011).

### 2.2. Copper content

The copper content of *Tth*-MCO and apo *Tth*-MCO was determined spectrophotometrically by the 2,2'-biquinoline method as described elsewhere (Felsenfeld, 1960). A solution of 2,2'-biquinoline at 0.5 mg ml<sup>-1</sup> in glacial acetic acid was prepared and standardized against five submicromolar solutions of copper(II) sulfate incubated with an excess of ascorbic acid (8 mM). Protein solutions (100 µl at 25 µM) were incubated for 20 min at 383 K for protein denaturation and mixed with 8 mM ascorbic acid for 10 min to reduce native enzyme cupric ions to copper(I). The latter solution was then mixed with 100 µl of the 2,2'-biquinoline solution and the Cu content was determined at  $\lambda = 546$  nm. Experiments were performed in triplicate and solutions were kept anaerobic by bubbling gaseous N<sub>2</sub> through them. All UV–visible spectra were recorded using an Evolution 1000 spectrophotometer (Thermo Scientific) using quartz cells with 1 cm path length.

### 2.3. X-ray data collection

Data collection was performed on beamline X6A of the National Synchrotron Light Source (NSLS), Brookhaven National Laboratory (BNL), New York, USA using an ADSC Quantum 270 detector. Diffraction data collection for *Tth*-MCO (12.650 keV,  $\lambda = 0.9795$  Å and 9.003 keV,  $\lambda = 1.3767$  Å), apo *Tth*-MCO (9.019 keV,  $\lambda = 1.3747$  Å) and Hg-*Tth*-MCO (14.900 keV,  $\lambda = 0.8321$  Å) has been described previously (Serrano-Posada *et al.*, 2011). A data set (12.650 keV,  $\lambda = 0.9795$  Å) was collected to a resolution of 1.7 Å for an Hg-*Tth*-MCO-2h crystal; its unit-cell parameters were  $a = 93.3$  Å,  $b = 110.1$  Å,  $c = 96.2$  Å,  $\alpha = \beta = \gamma = 90^\circ$ . X-ray absorption spectra in fluorescence mode were collected on beamline X6A using a Si(111) channel-cut monochromator (band pass  $1.9 \times 10^{-1}$  eV) and a custom-made fluorescence detector vertically positioned at an angle of 90° relative to the beam axis. Fluorescence spectra were subjected to background subtraction and normalized using the *Athena* software package (Ravel & Newville, 2005).

For the multicrystal data-collection strategy, data were collected from eight different *Tth*-MCO crystals obtained using the microseeding technique under the same conditions: 12.650 keV ( $\lambda = 0.9795$  Å), crystal-to-detector distance 200 mm,  $\Delta\varphi = 1.0^\circ$ , exposure time per image 30 s, slit size  $100 \times 100$  µm. Complete diffraction patterns of 96 images each were collected from eight *Tth*-MCO crystals at different known starting  $\varphi$  angles.

### 2.4. Data processing and model refinement

Diffraction images of *Tth*-MCO, apo *Tth*-MCO, Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h were integrated using *XDS* (Kabsch, 2010) and scaling was performed with *SCALA* from the *CCP4* suite (Winn *et al.*, 2011). For the multicrystal data-collection strategy, each of eight individual data sets was divided into sequential blocks of 12 images. These blocks from eight different crystals but having the absorbed dose in common were separately integrated using *XDS* (Kabsch, 2010) and then combined using *SORTMTZ* from the *CCP4* suite (Winn *et al.*, 2011) in order to obtain eight full composite data sets (*Tth*-MCO-C1–8) corresponding to increasing absorbed doses. Scaling was then performed as described above.

The *Tth*-MCO structure at 1.5 Å resolution was determined by a combination of the molecular-replacement (MR) and single-wavelength anomalous dispersion (SAD) techniques (Serrano-Posada *et al.*, 2011) using *Phaser* in MR-SAD mode (McCoy *et al.*, 2007). Initial model building was performed automatically using *ARP/wARP* (Langer *et al.*, 2008). The structures of the inactive forms of *Tth*-MCO at 1.7 Å resolution, apo *Tth*-MCO, Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h, as well as the active composite structures *Tth*-MCO-C1–8 at 1.8 Å resolution, were determined by MR using *Phaser* (McCoy *et al.*, 2007). The atomic coordinates of *Tth*-MCO (PDB entry 2xu9), from which all of the Cu ions, MPD molecules and water molecules had been removed, were used as a structural model for MR trials. All crystallographic



**Table 1**  
X-ray data-collection and refinement statistics.

Values in parentheses are for the last resolution shell.

	<i>Tth</i> -MCO	Apo <i>Tth</i> -MCO	Hg- <i>Tth</i> -MCO	Hg- <i>Tth</i> -MCO-2h
Data-collection statistics				
Space group	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>
Unit-cell parameters				
<i>a</i> (Å)	93.6	93.0	93.5	93.3
<i>b</i> (Å)	110.3	110.1	110.2	110.1
<i>c</i> (Å)	96.3	96.3	96.3	96.2
$\alpha = \beta = \gamma$ (°)	90.0	90.0	90.0	90.0
Resolution range (Å)	23.0–1.50 (1.60–1.50)	26.0–1.70 (1.80–1.70)	20.0–1.70 (1.80–1.70)	28.0–1.70 (1.80–1.70)
No. of reflections	899414	435926	542926	245055
No. of unique reflections	78896 (10654)	53818 (7154)	54841 (8585)	53273 (7694)
Completeness (%)	94.7 (83.9)	98.9 (82.1)	94.6 (91.4)	97.9 (97.5)
<i>R</i> <sub>merge</sub> (%)	8.6 (39.5)	9.0 (45.0)	13.3 (35.3)	5.5 (27.9)
CC <sub>1/2</sub> (%)	96.7 (96.3)	96.5 (96.0)	96.3 (95.8)	96.2 (95.8)
<i>I</i> / $\sigma$ ( <i>I</i> )	19.4 (4.1)	18.5 (3.8)	9.1 (3.7)	18.4 (5.1)
Multiplicity	11.4 (9.9)	8.1 (6.6)	9.9 (10.3)	4.6 (4.7)
Monomers per asymmetric unit	1	1	1	1
Refinement statistics				
Resolution range (Å)	23.0–1.50	26.0–1.70	20.0–1.70	28.0–1.70
<i>R</i> <sub>work</sub> / <i>R</i> <sub>free</sub> (%)	15.5/17.8	15.1/18.5	15.3/17.8	15.2/17.0
No. of atoms				
Protein	4259	4045	3837	3799
Ion/ligand	134	128	141	133
Water	549	525	455	504
Mean <i>B</i> values (Å <sup>2</sup> )				
Protein	13.7	16.5	17.2	15.8
Ion/ligand	33.1	37.8	38.3	34.8
Water	31.5	33.2	31.5	31.7
All atoms	16.2	18.9	19.0	18.2
Wilson plot	12.5	16.3	15.8	14.1
R.m.s.d. from ideal stereochemistry				
Bond lengths (Å)	0.01	0.02	0.02	0.01
Bond angles (°)	1.51	1.99	1.98	2.32
Coordinate error (maximum-likelihood-based)	0.19	0.18	0.17	0.17
Ramachandran plot (%)				
Most favoured regions	97.3	97.5	97.7	97.8
Additional allowed regions	2.4	2.2	1.9	2.2
Disallowed regions	0.3	0.3	0.4	0.0
PDB code	2xu9	2xuw	2xvb	4ai7

structures determined in this work belonged to the *C*-centred orthorhombic space group C222<sub>1</sub> as suggested by *POINTLESS* (Evans, 2006). Matthews coefficient calculations suggested that there was one molecule per asymmetric unit in all structures (~50% solvent content in all structures).

In all cases, refinement was performed using *PHENIX* (Adams *et al.*, 2010) until the *R*<sub>work</sub> and *R*<sub>free</sub> values were lower than 0.19 with satisfactory r.m.s. deviations from ideal bond lengths and bond angles. Refinement was alternated with manual building/refinement in *Coot* (Emsley *et al.*, 2010). 5% of the data were randomly chosen and reserved to determine *R*<sub>free</sub>. Water molecules were first automatically located using *PHENIX* and then validated in *Coot*. Meanwhile, several MPD molecules and metal ions were positioned manually in *Coot* and then refined using *PHENIX*. The occupancies of the metal atoms in all structures were adjusted so that their isotropic thermal vibration parameters were refined approximately to the values of the neighbouring atoms in the structure. For the *Tth*-MCO-C1 composite structure with the lowest absorbed X-ray dose (0.2 MGy), a symmetrical and elongated

electron density was found between the binuclear T3Cu–T3'Cu cluster and was assumed to be O<sub>2</sub>. Refinement then proceeded constraining the O=O distance to the target value of 1.21 Å. TLS refinement for the His95 residue of the composite structures *Tth*-MCO-C2–8 was performed using *PHENIX*. Changes in the electron density for the *Tth*-MCO-C1–8 structures, structural damage at each Cu atom and the decrease of the occupancy values induced by X-ray radiation were observed and analyzed as a function of the absorbed dose.

Model validation was performed using *MolProbity* (Chen, Arendall *et al.*, 2010). The graphical representations were made using *CCP4mg* v.2.7.3 (McNicholas *et al.*, 2011) and *PyMOL* (DeLano, 2002). Data-collection and refinement statistics for the *Tth*-MCO, apo *Tth*-MCO, Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h structures determined using classic single-crystal data collection are shown in Table 1. Data-collection and refinement statistics for the composite structures *Tth*-MCO-C1–8 determined using multicrystal data collection are shown in Table 2.

### 2.5. Dose calculation

Calculations of the absorbed dose, in megagrays (MGy), were performed using *RADDOSE* v.2 (Murray *et al.*, 2004; Paithankar *et al.*, 2009). The values of the beam parameters (including the energy, beam profile, size/area and flux) and the crystal properties (unit cell, space group, number of molecules per asymmetric unit, composition, size and thickness) were used in order to calculate the absorbed X-ray dose. An estimate of the X6A beam flux in photons s<sup>-1</sup> was obtained using a silicon pin diode to calibrate the beam intensity as described previously (Owen *et al.*, 2009). The sizes of the eight *Tth*-MCO crystals for the composite structures *Tth*-MCO-C1–8 were chosen in order to match the beam size, ensuring a uniform dose for all crystal regions.

## 3. Results and discussion

### 3.1. Crystallographic structures

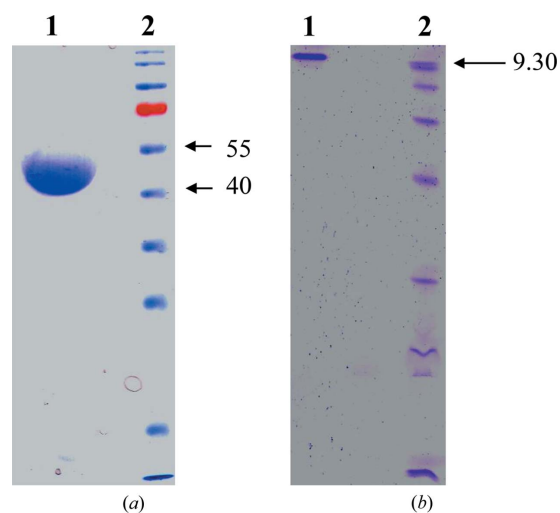
The classic single-crystal data-acquisition and multicrystal data-collection techniques were employed to determine 12 crystallographic structures of *Tth*-MCO in holo, apo and Hg-bound states and with different absorbed X-ray doses. The

**Table 2**  
X-ray data-collection and refinement statistics.

Values in parentheses are for the last resolution shell.

	<i>Tth</i> -MCO-C1	<i>Tth</i> -MCO-C2	<i>Tth</i> -MCO-C3	<i>Tth</i> -MCO-C4	<i>Tth</i> -MCO-C5	<i>Tth</i> -MCO-C6	<i>Tth</i> -MCO-C7	<i>Tth</i> -MCO-C8
Data-collection statistics								
Space group	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>	C222 <sub>1</sub>
Unit-cell parameters								
<i>a</i> (Å)	93.6	93.7	93.6	93.5	93.5	93.5	93.7	93.7
<i>b</i> (Å)	110.3	110.3	110.3	110.2	110.2	110.2	110.4	110.4
<i>c</i> (Å)	96.3	96.3	96.3	96.3	96.3	96.2	96.4	96.5
$\alpha = \beta = \gamma$ (°)	90.0	90.0	90.0	90.0	90.0	90.0	90.0	90.0
Resolution range (Å)	30.0–1.80 (1.90–1.80)	29.0–1.80 (1.90–1.80)	29.0–1.80 (1.90–1.80)	28.0–1.80 (1.90–1.80)	30.0–1.80 (1.90–1.80)	29.0–1.80 (1.90–1.80)	29.0–1.80 (1.90–1.80)	29.0–1.80 (1.90–1.80)
No. of reflections	174732	176253	151457	175957	153500	173277	175181	172277
No. of unique reflections	44803 (6341)	45193 (6600)	44546 (6378)	45117 (6463)	43857 (6385)	44430 (6520)	44918 (6345)	45336 (6589)
Completeness (%)	96.5 (95.2)	97.7 (98.7)	96.9 (95.5)	97.6 (96.6)	95.4 (95.5)	96.7 (97.8)	97.2 (95.8)	97.9 (98.4)
<i>R</i> <sub>merge</sub> (%)	13.0 (28.0)	27.6 (40.1)	12.4 (29.6)	12.6 (30.0)	13.6 (32.6)	13.4 (32.9)	13.1 (32.4)	13.5 (31.2)
CC <sub>1/2</sub> (%)	95.8 (95.4)	95.1 (94.9)	95.5 (95.1)	95.8 (95.4)	95.5 (95.0)	95.7 (95.2)	96.0 (95.4)	95.4 (94.9)
<i>I</i> / $\sigma$ ( <i>I</i> )	7.6 (4.2)	6.8 (3.5)	7.8 (4.2)	8.1 (4.2)	7.4 (4.0)	8.0 (4.1)	8.1 (4.1)	7.9 (4.0)
Multiplicity	3.9 (4.0)	3.9 (3.8)	3.4 (3.4)	3.9 (3.9)	3.5 (3.5)	3.9 (3.9)	3.9 (4.0)	3.8 (3.8)
Monomers per asymmetric unit	1	1	1	1	1	1	1	1
Refinement statistics								
Resolution range (Å)	30.0–1.80	29.0–1.80	29.0–1.80	28.0–1.80	30.0–1.80	29.0–1.80	29.0–1.80	29.0–1.80
<i>R</i> <sub>work</sub> / <i>R</i> <sub>free</sub> (%)	14.7/17.4	15.4/18.9	14.8/17.9	14.7/17.2	14.8/18.2	14.9/17.9	14.6/16.8	15.0/18.3
No. of atoms								
Protein	3798	3786	3770	3763	3746	3790	3786	3762
Ion/ligand	134	125	133	133	133	131	133	117
Water	500	516	509	498	543	512	508	501
Mean <i>B</i> values (Å <sup>2</sup> )								
Protein	12.1	12.0	12.0	11.9	11.9	12.5	13.0	12.4
Ion/ligand	31.8	30.6	31.3	31.3	31.3	32.0	32.5	31.1
Water	28.3	28.6	28.8	28.2	29.7	30.0	31.0	29.4
All atoms	14.6	14.5	14.6	14.3	14.7	15.1	15.7	14.8
Wilson plot	11.2	11.3	11.2	11.2	11.1	12.2	12.9	11.9
R.m.s.d. from ideal stereochemistry								
Bond lengths (Å)	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Bond angles (°)	1.57	1.43	1.39	1.40	1.42	1.42	1.41	1.43
Coordinate error	0.17	0.18	0.15	0.15	0.17	0.18	0.17	0.19
(maximum-likelihood-based)								
Ramachandran plot (%)								
Most favoured regions	97.6	96.9	97.8	97.8	97.8	97.5	97.6	97.4
Additional allowed regions	2.2	3.1	2.2	2.2	1.8	2.1	2.0	2.2
Disallowed regions	0.2	0.0	0.0	0.0	0.4	0.4	0.4	0.4
PDB code	2yae	2yaf	2yah	2yam	2yao	2yap	2yaq	2yar

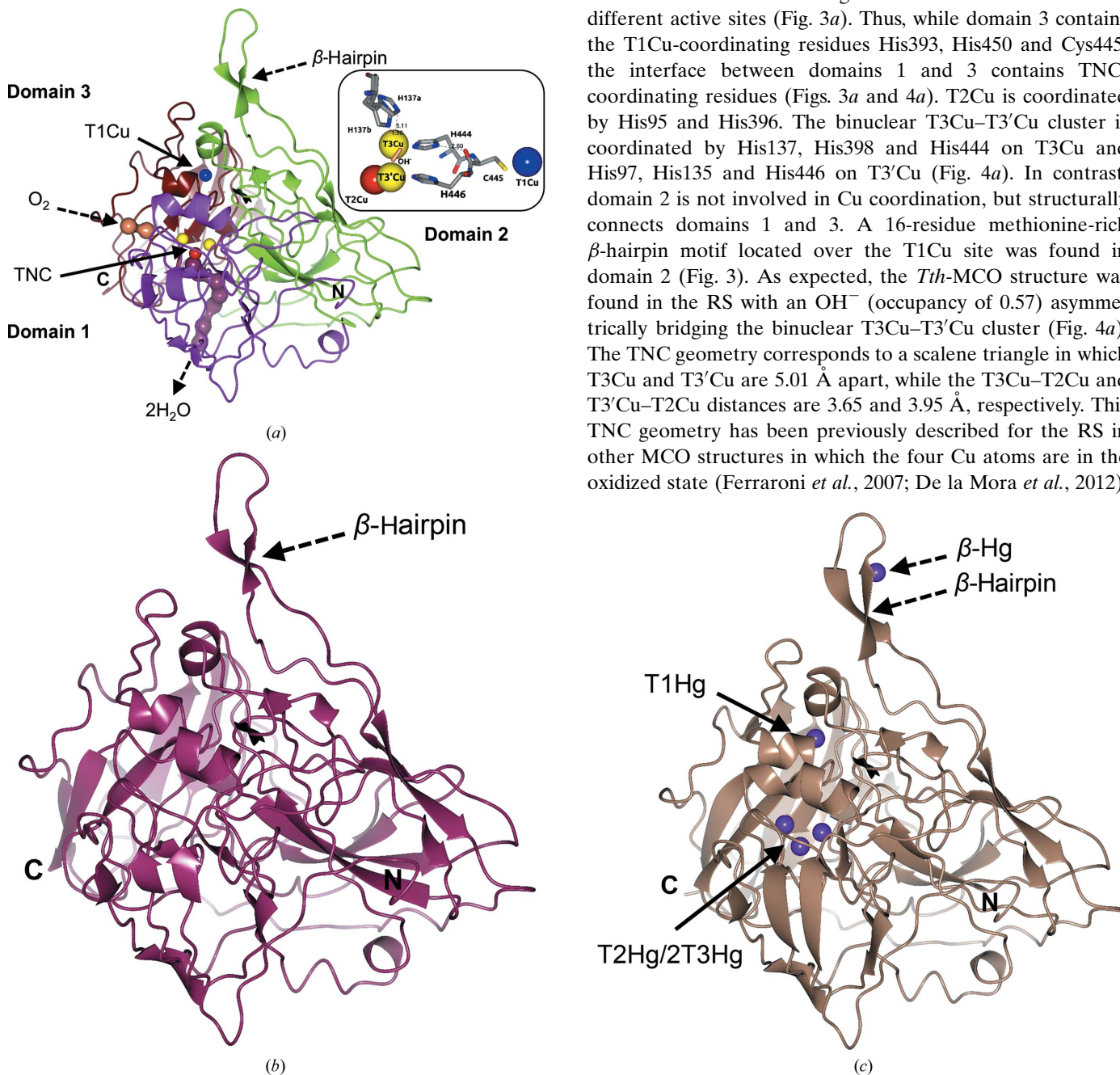
latter structures were determined with increasing doses from *Tth*-MCO-C1 (0.2 MGy, 12.5% dose) to *Tth*-MCO-C8 (1.6 MGy, 100% dose). The overall architecture of inactive forms of the enzyme (apo *Tth*-MCO, Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h) is identical to that of *Tth*-MCO; a calculated r.m.s. deviation of 0.16 Å was obtained after superimposing 439 C<sup>α</sup> atoms. The most interesting changes were observed in metal-coordinating residues and Glu451/Asp106 at the TNC, which are involved in the first and second coordination spheres, respectively. The markedly different behaviour of Glu451/Asp106 in the active and inactive forms of *Tth*-MCO provided structural details of the proton-relay mechanism of O<sub>2</sub> reduction. In addition, a methionine-rich β-hairpin motif, which seems to be involved in an extra metal-coordinating site, was also observed in all structures. Different states (the O<sub>2</sub> state, RS and FR state) of the X-ray-induced reduction of O<sub>2</sub> were trapped in the *Tth*-MCO-C1, *Tth*-MCO-C2 and *Tth*-MCO-C6 structures. The four catalytic Cu atoms exhibited a different radiation-damage behaviour from absorbed X-rays. Accordingly, while T2Cu and T3Cu showed a high susceptibility to radiation damage, both T1Cu and T3'Cu seem to be quite stable to X-rays.



**Figure 2**  
(a) Coomassie Blue-stained SDS-PAGE (12%) of purified recombinant *Tth*-MCO. Lane 1, purified *Tth*-MCO. Lane 2, molecular-weight markers (Fermentas; labelled in kDa). (b) IEF gel. Lane 1, purified *Tth*-MCO. Lane 2, isoelectric point markers (Pharmacia; pI value labelled).

**3.1.1. *Tth*-MCO structure.** *Tth*-MCO is a monomeric enzyme (Miyazaki, 2005) which showed a molecular mass of ~50 kDa on SDS-PAGE and an isoelectric point (pI) of ~9.3 on an isoelectric focusing (IEF) gel (Fig. 2). The overall *Tth*-MCO fold comprises three sequential cupredoxin globular domains: domain 1, Gly24–Ala169, domain 2, Glu170–Val343; domain 3, Val344–Gly462 (Fig. 3a). This fold was first observed in the cupredoxin family, which includes the single-domain

blue copper proteins plastocyanin and azurin (Murphy *et al.*, 1997), and was subsequently detected in two-domain (Komori *et al.*, 2009) and three-domain (Zhukhlistova *et al.*, 2008) MCOs. The cupredoxin fold basically consists of an eight-stranded Greek-key  $\beta$ -barrel comprising two  $\beta$ -sheets composed of four strands each arranged in a sandwich conformation (Murphy *et al.*, 1997). The *Tth*-MCO fold contains 35  $\beta$ -strands and five  $\alpha$ -helices distributed in the three domains. The four catalytic Cu ions are distributed among two of the three domains in two different active sites (Fig. 3a). Thus, while domain 3 contains the T1Cu-coordinating residues His393, His450 and Cys445, the interface between domains 1 and 3 contains TNC-coordinating residues (Figs. 3a and 4a). T2Cu is coordinated by His95 and His396. The binuclear T3Cu–T3'Cu cluster is coordinated by His137, His398 and His444 on T3Cu and His97, His135 and His446 on T3'Cu (Fig. 4a). In contrast, domain 2 is not involved in Cu coordination, but structurally connects domains 1 and 3. A 16-residue methionine-rich  $\beta$ -hairpin motif located over the T1Cu site was found in domain 2 (Fig. 3). As expected, the *Tth*-MCO structure was found in the RS with an OH<sup>−</sup> (occupancy of 0.57) asymmetrically bridging the binuclear T3Cu–T3'Cu cluster (Fig. 4a). The TNC geometry corresponds to a scalene triangle in which T3Cu and T3'Cu are 5.01 Å apart, while the T3Cu–T2Cu and T3'Cu–T2Cu distances are 3.65 and 3.95 Å, respectively. This TNC geometry has been previously described for the RS in other MCO structures in which the four Cu atoms are in the oxidized state (Ferraroni *et al.*, 2007; De la Mora *et al.*, 2012).



**Figure 3** Crystallographic structures of *Tth*-MCO in holo, apo and Hg-bound states. (a) High-resolution structure of *Tth*-MCO showing the cupredoxin-domain organization (domain 1, magenta; domain 2, green; domain 3, brown) with T1Cu (blue sphere) located in domain 3 and the TNC (T2Cu, red sphere; binuclear T3Cu–T3'Cu cluster, yellow spheres) placed at the interface between domains 1 and 3. The molecular oxygen-entrance channel (coral) towards T3Cu and the water-molecule exit channel (dark purple) from T2Cu are represented by water molecules found in the *Tth*-MCO structure. Right inset, close-up of four catalytic Cu atoms of *Tth*-MCO, the hydroxide ligand (coral cylinder) bridging the binuclear T3Cu–T3'Cu cluster and the electron-transfer pathway from T1Cu to the TNC. (b, c) Two inactive forms of *Tth*-MCO: (b) apo *Tth*-MCO (dark purple) and (c) Hg-*Tth*-MCO-2h (pale brown) with Hg bound (purple spheres) instead of Cu. It is notable that a fifth Hg-binding site ( $\beta$ -Hg) was observed in (c). Distances are in Å.



Furthermore, Met455 in a double conformation was found flanking T1Cu (Fig. 4*a*). This axial methionine residue has been proposed to be involved in tuning the reduction potential ( $E^\circ$ ) of T1Cu (Quintanar *et al.*, 2007) and is frequently found in enzymes with low  $E^\circ$  (Zhukhlistova *et al.*, 2008).

Interestingly, T1Cu and T3'Cu proved to be highly stable to radiation damage, with occupancies of 0.87 and 0.90, respectively. In contrast, T2Cu and T3Cu showed occupancies of 0.13 and 0.43, respectively. The low occupancies of T2Cu and T3Cu in the crystalline state indicate the high susceptibility of both Cu atoms to radiation damage, since in solution the 2,2'-biquinoline method resulted in a ratio of  $4.1 \pm 0.1$  Cu atoms per *Tth*-MCO molecule. Moreover, a solution of *Tth*-MCO before crystallization trials also showed typical absorption peaks for both the  $C445S \rightarrow T1Cu^{II}$  charge-transfer band at  $\sim 610$  nm and the shoulder of the  $T3Cu^{II} \leftarrow OH^- \rightarrow T3'Cu^{II}$  charge-transfer band at  $\sim 330$  nm in UV-visible spectra (Fig. 5*a*). This information suggests that in the crystalline state there are probably several factors that increase the susceptibility of T2Cu and T3Cu to radiation damage. For instance, partial occupancy of T3Cu could be explained by the significant increase from 1.90 to 3.11 Å of the His137 N<sup>6</sup>-T3Cu coordination-bond distance (see the inset in Fig. 3*a*). Additionally, as Augustine *et al.* (2010) have previously described for Fet3p, the electron-transfer pathway from T1Cu to TNC is conserved in *Tth*-MCO via T1Cu<sup>II</sup>-Cys445-His444-T3Cu<sup>II</sup> and involves a hydrogen bond of 2.80 Å between the carbonyl group of Cys445 and His444 N<sup>δ</sup> (see inset in Fig. 3*a*). Furthermore, unlike T3'Cu, which is flanked by Trp133 in a cation- $\pi$  interaction (Fig. 4*a*), T3Cu is located in the entry channel for the reduction of O<sub>2</sub> (Fig. 3*a*). Therefore, since T3Cu is located in both the entry channel of O<sub>2</sub> and at the end of the

electron-transfer pathway for the reduction of O<sub>2</sub>, this Cu atom is probably more affected than T3'Cu by electrons released by X-rays during data collection. On the other hand, although the location of T2Cu in the exit channel for the water molecules produced is structurally important for the catalytic

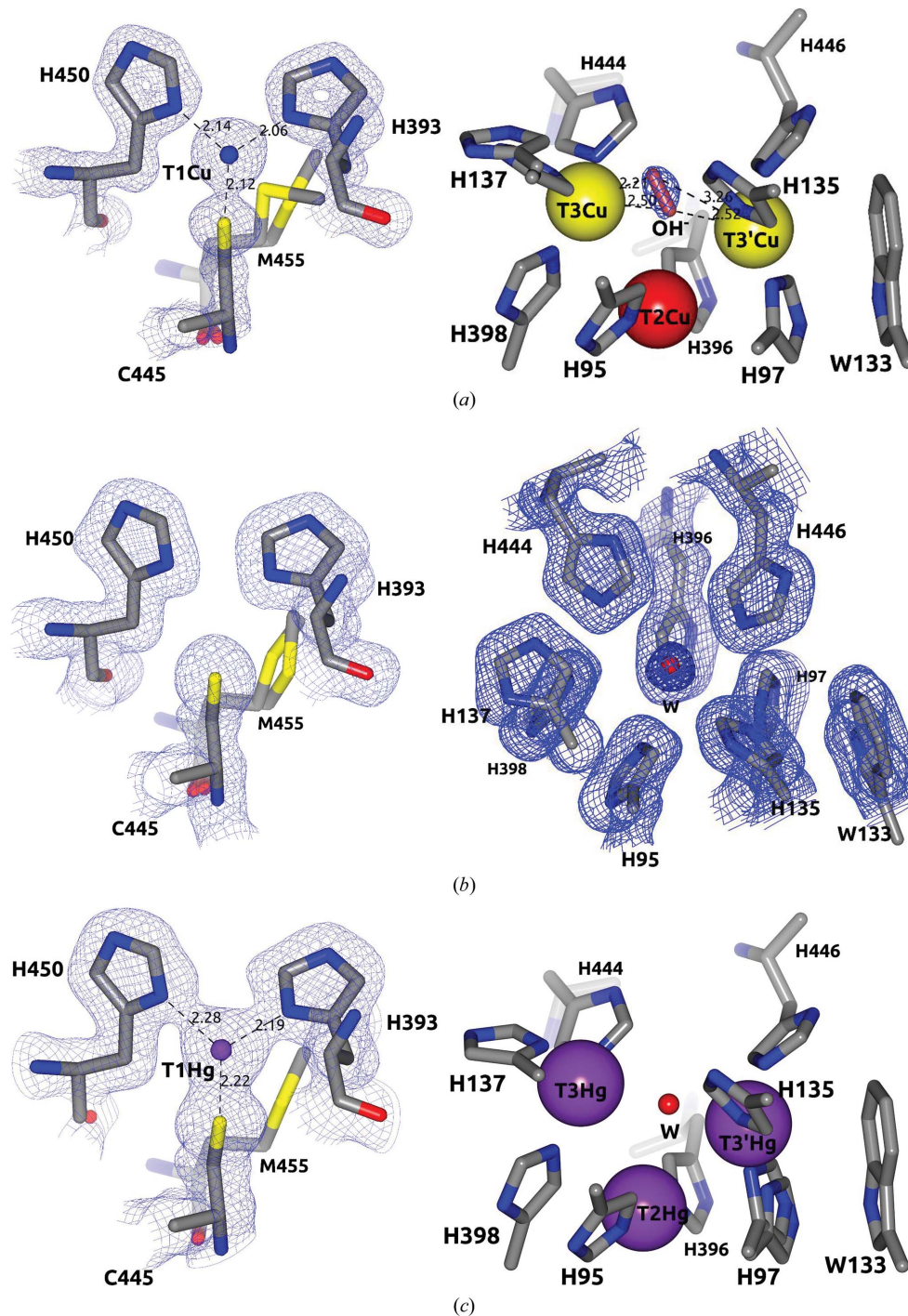


Figure 4

Geometry of the T1 site and TNC. (*a*) *Tth*-MCO. Note the hydroxide ligand (coral cylinder) in the RS asymmetrically bridging the binuclear T3Cu-T3'Cu cluster and Met455 flanking T1Cu. (*b*) Apo *Tth*-MCO structure. Note the absence of electron density for T1Cu and the TNC (superior view). A water molecule (red sphere) in the same position as OH<sup>-</sup> observed in (*a*) was found in this inactive apo form. (*c*) Hg-*Tth*-MCO-2h. Note that the view of TNC is the same that in (*a*) and a water molecule (red sphere) bridging the T3Cu-T3'Cu cluster was also found in this inactive form with Hg bound. The  $2F_o - F_c$  electron-density map contoured at the  $1\sigma$  level is drawn in blue. Metal ions are represented as spheres. Distances are in Å.



activity of the enzyme (Fig. 3*a*), the partial occupancy of T2Cu suggests that at this position specially designed for the removal of water molecules in MCOs during the reduction of O<sub>2</sub>, T2Cu could be continually exposed to hydrated electrons released by X-rays.

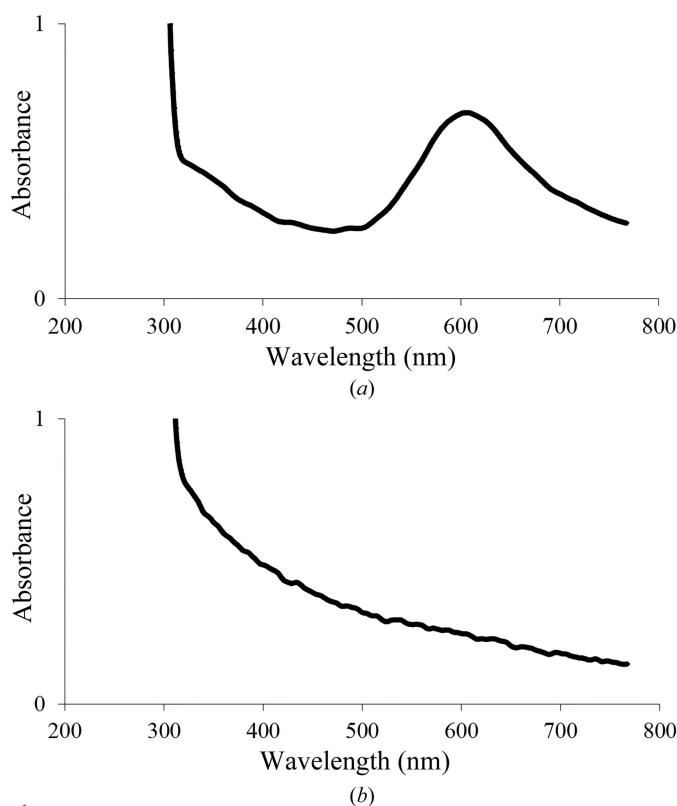
**3.1.2. Apo *Tth*-MCO, Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h.** To assess the significantly different behaviour between the active and inactive forms of *Tth*-MCO, three inactive structures of the apo and Hg-bound forms were determined. As expected, no electron density was observed for the four catalytic Cu atoms in apo *Tth*-MCO (Figs. 3*b* and 4*b*), indicating that the overexpression and purification processes did not incorporate Cu atoms into this inactive form of *Tth*-MCO. In solution, the 2,2'-biquinoline method also resulted in a ratio of  $0.4 \pm 0.1$  Cu atoms per apo *Tth*-MCO molecule and a UV-visible spectrum revealed the absence of the typical absorption peaks at both  $\sim 610$  and  $\sim 330$  nm (Fig. 5*b*).

Two inactive derivatives, Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h, were prepared with the spectroscopically silent and redox-inactive Hg (Palmer *et al.*, 2001) instead of Cu. Remarkably, the coordination of Hg atoms with partial occupancies in both the inactive Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h (Fig. 3*c*) structures, obtained with different soaking times from two different apo *Tth*-MCO crystals, occurred in the four vacant Cu sites. Together with this observation, a fifth Hg-binding ( $\beta$ -Hg) site was also found in the methionine-rich  $\beta$ -hairpin motif (Fig. 3*c*). Independently, X-ray fluorescence (XRF)

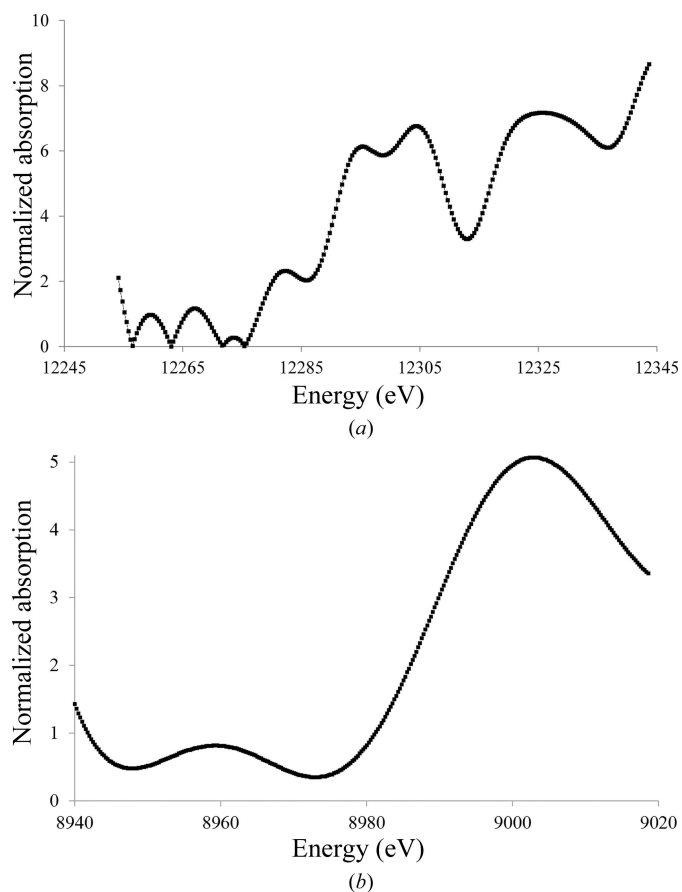
spectra were collected from an Hg-*Tth*-MCO-2h crystal before data collection. The *L*<sub>III</sub> absorption edges of Hg-*Tth*-MCO-2h were observed at 12.295, 12.304 and 12.325 keV in the fluorescence spectrum (Fig. 6*a*), correlating with the *L*<sub>III</sub> absorption edges experimentally determined by Ji *et al.* (2001). In addition, a single characteristic peak of copper(II) was always observed at 8.999 keV in the XRF spectrum from *Tth*-MCO crystals before data collection (Fig. 6*b*). A water molecule bridging the binuclear T3Cu–T3'Cu site was also found in all inactive structures (Figs. 4*b* and 4*c*), indicating the structural relevance of the O<sub>2</sub>-reduction site at the TNC, in which different states and intermediates of the catalytic cycle are located.

**3.1.3. Composite structures *Tth*-MCO-C1–8 obtained with different absorbed X-ray doses.** To identify the states and/or the intermediates in the X-ray-induced O<sub>2</sub> reduction, eight composite structures corresponding to increasing absorbed dose levels from *Tth*-MCO-C1 (0.2 MGy, 12.5% dose) to *Tth*-MCO-C8 (1.6 MGy, 100% dose) were determined using the multiscrystal data-collection technique.

Interestingly, the *Tth*-MCO-C1 structure at a low absorbed dose was found with an O<sub>2</sub> molecule (occupancy of 0.4) almost symmetrically coordinated amongst the binuclear



**Figure 5**  
Spectra of purified *Tth*-MCO in 20 mM Tris–HCl pH 8.0. (a) Holo form (450 μM). (b) Apo form (500 μM). Note that although the concentration in (b) is higher than that in (a), the typical absorption peaks at both  $\sim 330$  and  $\sim 610$  nm were not observed.



**Figure 6**  
X-ray absorption spectra of *Tth*-MCO crystals in the holo form and with Hg bound before X-ray data collection. (a) Hg-*Tth*-MCO-2h single crystal showing the *L*<sub>III</sub> absorption edges of protein-bound Hg. (b) *Tth*-MCO single crystal showing the Cu K absorption edge of the four native copper(II) atoms of the enzyme.

T3Cu–T3'Cu cluster (Fig. 7a). This state is structurally similar to the previously described O<sub>2</sub> state for low absorbed dose laccase from *M. albomyces* (PDB entry 2ih8; Hakulinen *et al.*, 2006) and CotA (PDB entry 1w6l; Bento *et al.*, 2005). The finding of the O<sub>2</sub> state in *Tth*-MCO-C1 suggests that the electron population released in the crystal by the X-ray radiolysis of water molecules is relatively deficient at doses of  $\leq 0.2$  MGy. In agreement with this observation, the stabilization of O<sub>2</sub> also suggests that four Cu atoms are oxidized (Fig. 1e) and that the FR state has not yet been formed.

Notably, the *Tth*-MCO-C2 structure at 25.0% dose was found to have an OH<sup>−</sup> ion (occupancy of 0.7) coordinated between the binuclear T3Cu–T3'Cu cluster (Fig. 7b), indicating that the population of protons and electrons in the crystal significantly increased from *Tth*-MCO-C1 to generate the RS in *Tth*-MCO-C2. This is particularly interesting with regard to the experiments on Fet3p (Augustine *et al.*, 2010), since in the path from the *Tth*-MCO-C1 to the *Tth*-MCO-C2 structures the system progresses through the following states and intermediates: O<sub>2</sub> state (*Tth*-MCO-C1)→PI→NI→RS (*Tth*-MCO-C2; Fig. 1). However, whilst it is well known that the FR state is the first state of the catalytic cycle for the reduction of O<sub>2</sub> by MCOs (Augustine *et al.*, 2010) and that it reacts immediately with O<sub>2</sub> to form the PI, the structural stabilization of the O<sub>2</sub> state in the *Tth*-MCO-C1 structure seems to be a process that is exclusive to the crystalline state as the result of a deficiency of electrons to generate the FR state at doses of  $\leq 0.2$  MGy. As also mentioned above, in the absence of sufficient electrons to form the FR state in solution, MCOs are usually observed in the RS (Augustine *et al.*, 2010). Consequently, the formation of the O<sub>2</sub> state in *Tth*-MCO-C1 remains unclear, and in fact even the nature of the RS of MCOs is a question that still remains to be answered (Bento *et al.*, 2010). Unsurprisingly, the PI and NI cannot be trapped under the conditions of these crystallographic experiments on *Tth*-MCO owing to their rate of decay and the intrinsic nature of the intermediates.

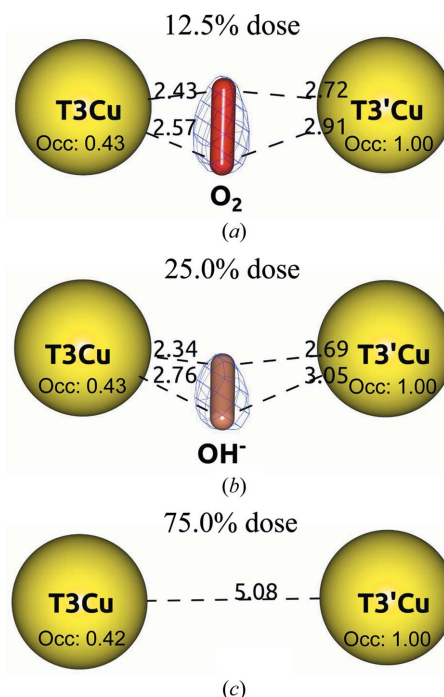
The RS was always observed in the *Tth*-MCO-C3 (37.5% dose) to *Tth*-MCO-C5 (62.5% dose) structures, indicating that the FR state is not stable enough to be trapped, regardless of the increasing levels of absorbed dose. However, when the dose reached 75.0% in the *Tth*-MCO-C6 structure, no state or intermediate was found at the TNC; a distance of 5.08 Å between the binuclear T3Cu–T3'Cu cluster was also observed (Fig. 7c). This observation indicates that the FR state seems to be trapped at 75.0% dose, because in accordance with the mechanism of O<sub>2</sub> reduction at the TNC (Fig. 1), the RS (*Tth*-MCO-C5) should generate the FR state (*Tth*-MCO-C6) for the next enzyme cycle (Quintanar *et al.*, 2005). Naturally, the structural stabilization of the O<sub>2</sub> state observed in *Tth*-MCO-C1 cannot be obtained again considering that the population of protons and electrons is significant at this dose. In addition, the FR state is the only state in the reduction of the O<sub>2</sub> in MCOs that presents a binuclear T3Cu–T3'Cu cluster distance of  $>5$  Å (Augustine *et al.*, 2010). Remarkably, the FR state in *Tth*-MCO-C6 is also structurally similar to the previously reported FR state for CotA (PDB entry 2bhf;

Bento *et al.*, 2005), which displayed a binuclear T3Cu–T3'Cu cluster distance of 5.10 Å. Finally, according to the mechanism of O<sub>2</sub> reduction at the TNC (Fig. 1), the FR state should generate the RS if the system has progressed as follows: FR (*Tth*-MCO-C6)→PI→NI→RS (*Tth*-MCO-C7). In fact, the last two composite structures *Tth*-MCO-C7 (87.5% dose) and *Tth*-MCO-C8 (1.6 MGy, 100% dose) were found in the RS.

Accordingly, the composite structures *Tth*-MCO-C1–8 support previous observations indicating that the final crystallographic structure of an MCO obtained using classic single-crystal data acquisition such as *Tth*-MCO (Fig. 3a) is an average model of different Cu oxidation states and X-ray-induced O<sub>2</sub>-reduction states and intermediates. As also mentioned above, *Tth*-MCO was found in the RS (Figs. 3a and 4a); this is the statistically relevant state in a full data set of this enzyme, as the composite structures *Tth*-MCO-C1–8 have shown.

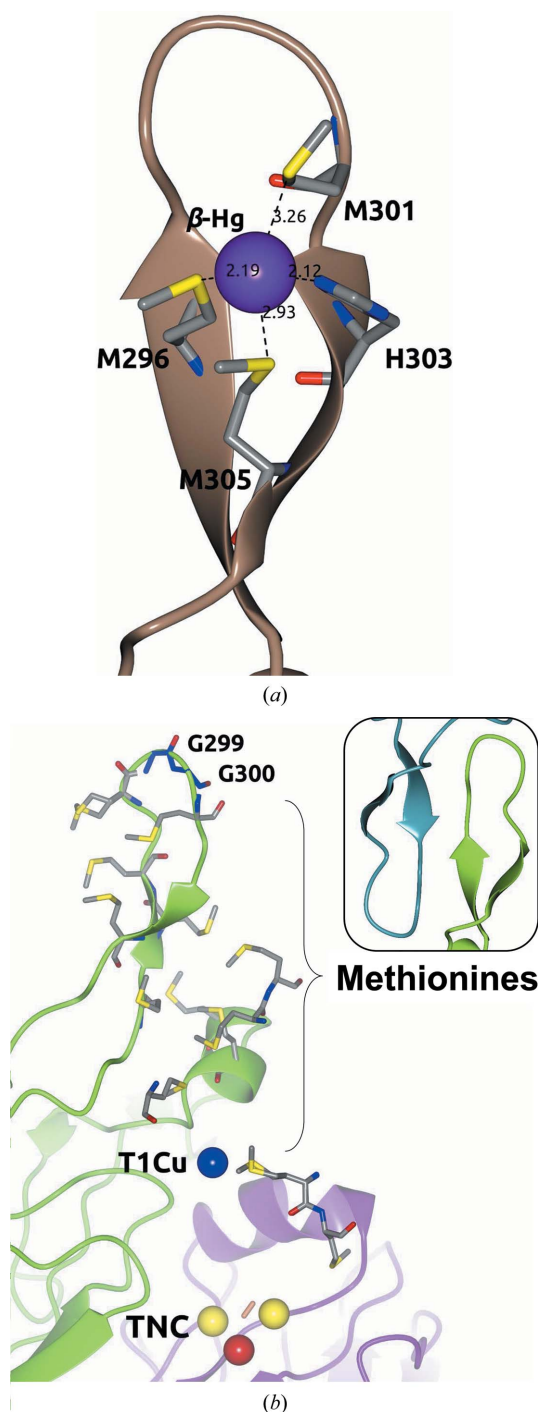
### 3.2. Methionine-rich $\beta$ -hairpin motif

The MCOs found in many bacterial copper-detoxification systems have both methionine-rich structural motifs and copper(I) oxidase activity (Cha & Cooksey, 1991; Lee, Grass *et al.*, 2002; Fernandes *et al.*, 2007). Interestingly, a 16-residue methionine-rich  $\beta$ -hairpin motif (Ala292–Gln307; Fig. 3) over the T1Cu site was observed in all of the crystallographic structures determined in this work. This structural motif contains six methionines, Met293, Met295, Met296, Met298,



**Figure 7**  
Different states of O<sub>2</sub> reduction found at the TNC of the *Tth*-MCO composite structures. (a) *Tth*-MCO-C1 in the O<sub>2</sub> state. (b) *Tth*-MCO-C2 in the RS. (c) *Tth*-MCO-C6 in the FR state. Note that the O<sub>2</sub> (red cylinder) is bridging the binuclear T3Cu–T3'Cu cluster at a dose of 0.2 MGy (12.5% dose) in (a), as well as the different occupancy values related to the radiation damage of both Cu atoms. The  $2F_o - F_c$  electron-density map contoured at the  $1\sigma$  level is drawn in blue. Metal ions are represented as spheres. Distances are in Å.

Met301 and Met305, of the 13 in total in the primary sequence of the enzyme. In addition, the  $\beta$ -Hg (Figs. 3c and 8a) observed in both proteins with bound Hg is coordinated by four residues of the  $\beta$ -hairpin motif: Met296, Met301, Met305 and His303. However, low occupancies of 0.10 ( $B$  value of 83.4  $\text{\AA}^2$ ) in Hg-*Tth*-MCO and 0.20 ( $B$  value of 58.5  $\text{\AA}^2$ ) in



**Figure 8** Methionine-rich  $\beta$ -hairpin motif over the T1Cu site. (a)  $\beta$ -Hg coordinated in the  $\beta$ -hairpin motif of Hg-*Tth*-MCO-2h. (b) Methionine distribution between the top (Gly299-Gly300; blue cylinders) of the  $\beta$ -hairpin motif and T1Cu of *Tth*-MCO. Right inset: the  $\beta$ -hairpin motif (green) of *Tth*-MCO stabilized by the same motif (dark cyan) of the crystallographic neighbour. Metal ions are represented as spheres. Distances are in  $\text{\AA}$ .

Hg-*Tth*-MCO-2h were found for this extra Hg atom. This observation corresponds to the structural dynamics of the local environment of the  $\beta$ -Hg, which displayed average  $B$  values of 77.2 and 68.2  $\text{\AA}^2$  for Hg-*Tth*-MCO and Hg-*Tth*-MCO-2h, respectively. No electron density was observed at the  $\beta$ -Hg site for *Tth*-MCO and apo *Tth*-MCO.

Similar structural motifs over the T1Cu site have been described previously in other MCOs such as McoP from *Pyrobaculum aerophilum* (PDB entry 3aw5; Sakuraba *et al.*, 2011), McoC from *Campylobacter jejuni* (PDB entry 3zx1; Silva, Durão *et al.*, 2012) and CueO (PDB entry 1kv7; Roberts *et al.*, 2002), but the methionine-rich  $\beta$ -hairpin motif is unique to *Tth*-MCO (Fig. 3). Molecular-dynamics simulations have shown that this *Tth*-MCO motif experiences conformational changes which enable exposure of the T1Cu site for substrate oxidation (Bello *et al.*, 2012), regardless of its stabilization by the same motif in a crystallographic neighbour (see the inset in Fig. 8b).

It has also previously been shown that the methionine-rich CueO motif is involved in several copper(I)-coordinating sites apart from the four catalytic Cu atoms, which increase the metal oxidase activity to oxidize copper(I) to the less toxic copper(II) (Roberts *et al.*, 2003; Singh *et al.*, 2011). Indeed, the removal of the extra copper(I)-binding sites through the mutation of copper(I)-coordinating residues leads to a catalytic impairment of copper(I) oxidation (Singh *et al.*, 2011). Moreover, deletion of the methionine-rich motif decreased the copper(I) oxidase activity and increased the organic substrate oxidase activity through increased access to T1Cu (Kataoka *et al.*, 2007). Nevertheless, the mechanism of copper(I) oxidation is still unknown, but it has been suggested that it involves an extended electron-transfer pathway from the extra copper(I)-binding sites to T1Cu and then to the TNC (Singh *et al.*, 2011).

Consequently, it appears to be very probable that the methionine-rich  $\beta$ -hairpin motif of *Tth*-MCO is involved in extra copper(I)-binding sites. Nevertheless, electron transfer from the extra copper(I)-binding sites to T1Cu of *Tth*-MCO seems to be difficult, since there is a long distance of  $\sim 23$   $\text{\AA}$  from a hypothetical copper(I) placed in the same position as  $\beta$ -Hg to T1Cu. Notably, *Tth*-MCO possesses 13 methionines in the primary sequence, which are all located between the top (Gly299-Gly300) of the  $\beta$ -hairpin motif and T1Cu (Fig. 8b), suggesting that several positions for extra copper(I)-binding sites, differing from the  $\beta$ -Hg site, may be present in the enzyme. As mentioned above, *Tth*-MCO exhibits laccase activity towards organic substrates such as guaiacol and ABTS (Miyazaki, 2005). However, this information suggests that *Tth*-MCO also displays important structural features over the T1Cu site that could be related to its oxidase activity towards inorganic substrates, although there is no experimental evidence to support this view.

### 3.3. Proton-relay mechanism for the X-ray-induced reduction of $\text{O}_2$ to $2\text{H}_2\text{O}$

To assess the role in the crystalline state of the highly conserved pair of acidic residues Glu451/Asp106 involved in



the proton-relay mechanism for the X-ray-induced reduction of O<sub>2</sub>, and in particular their behaviour in active and inactive forms of *Tth*-MCO, several crystallographic structures of the enzyme in holo, apo and Hg-bound forms and with different X-ray absorbed doses were analyzed.

**3.3.1. *Tth*-MCO, apo *Tth*-MCO and Hg-*Tth*-MCO-2h.** In active holo *Tth*-MCO (Fig. 3a), which contains the four catalytic Cu atoms, a double conformation Glu451a (~7 Å to the TNC; occupancy of 0.56) and Glu451b (~4.5 Å to the TNC; occupancy of 0.44) of the proton-donor residue Glu451 located in the entry channel of O<sub>2</sub> was observed (Fig. 9a). A positive peak of electron density above 3.5σ in an  $F_o - F_c$  map for Glu451a O<sup>ε2</sup> strongly suggests a carboxyl functional group at the side chain, while its significant absence at the side chain of Glu451b suggests a carboxylate functional group. This observation indicates that in a proton-relay mechanism the Glu451a conformation seems to accept the protons for O<sub>2</sub> reduction, while the Glu451b conformation releases the protons to the TNC in a manner that appears to take place as follows: Glu451a (RCOOH) → Glu451b (RCOO<sup>-</sup> + H<sup>+</sup>) → TNC (Fig. 9a). In addition, it has previously been shown that the most likely proton-relay path from Glu451b to the TNC is through a structural water molecule (W<sub>3</sub>; Fig. 9a; Bento *et al.*, 2005, 2010; Augustine *et al.*, 2007; Kataoka *et al.*, 2009). According to the latter, structural data for *Tth*-MCO also suggest that apart from W<sub>3</sub>, which is at a distance of 2.84 Å from Glu451b O<sup>ε2</sup>, a second path with a distance of 2.90 Å between Glu451b O<sup>ε1</sup> and His137a N<sup>ε</sup> is also plausible (Fig. 9a).

A theoretical calculation has shown that Glu451 presents an unusually high pK<sub>a</sub> of 9.9 (Bello *et al.*, 2012), which is probably the result of interactions with other carboxylic groups at

the TNC (Harris & Turner, 2002; Castañeda *et al.*, 2009). Accordingly, the perturbed pK<sub>a</sub> value of Glu451 seems to be governed by its interaction with the Asp452 residue (pK<sub>a</sub> = 4.40, as calculated using the *PROPKA* server; Olsson *et al.*, 2011), which is at a hydrogen-bonding distance of 3.14 Å from Glu451a O<sup>ε2</sup> (Fig. 9a). Remarkably, the Asp452 residue is also conserved in other extensively studied bacterial MCOs such as CueO (Asp507; PDB entry 1kv7; Roberts *et al.*, 2002) and CotA (Asp499; PDB entry 1gsk; Enguita *et al.*, 2003), but in the latter cases the double conformation of the proton-donor residue was not observed. On the other hand, the structural factors that govern the unusual pK<sub>a</sub> value of the proton-donor residues of MCOs from fungi (Taylor *et al.*, 2005; De la Mora *et al.*, 2012) are not obvious from inspection of their crystal structures. For instance, the proton-donor residue Glu487 of Fet3p is in the middle of the TNC, and although there are several acidic residues around this residue none of them seems to explain the unusual pK<sub>a</sub> value of Glu487 (PDB entry 1zpu; Taylor *et al.*, 2005). However, when a double conformation of Glu487 is drawn in *Coot* (Emsley *et al.*, 2010), interactions with several neighbouring carboxylic groups occur, but in particular with the Glu134 residue, which would be at a hydrogen-bonding distance from Glu487. As expected, a theoretical calculation using the *PROPKA* server also showed that Asp106 (pK<sub>a</sub> = 1.13) is not a proton-donor residue in the proton-relay mechanism of *Tth*-MCO (Fig. 9a) but is in fact important for the decay of the PI to the NI (Figs. 1b and 1c; Quintanar *et al.*, 2005).

As mentioned above, the decay rate of both the PI and the NI restricts their structural characterization as single species, making them difficult to trap in the crystalline state. Furthermore, apart from the rest states and intermediates in

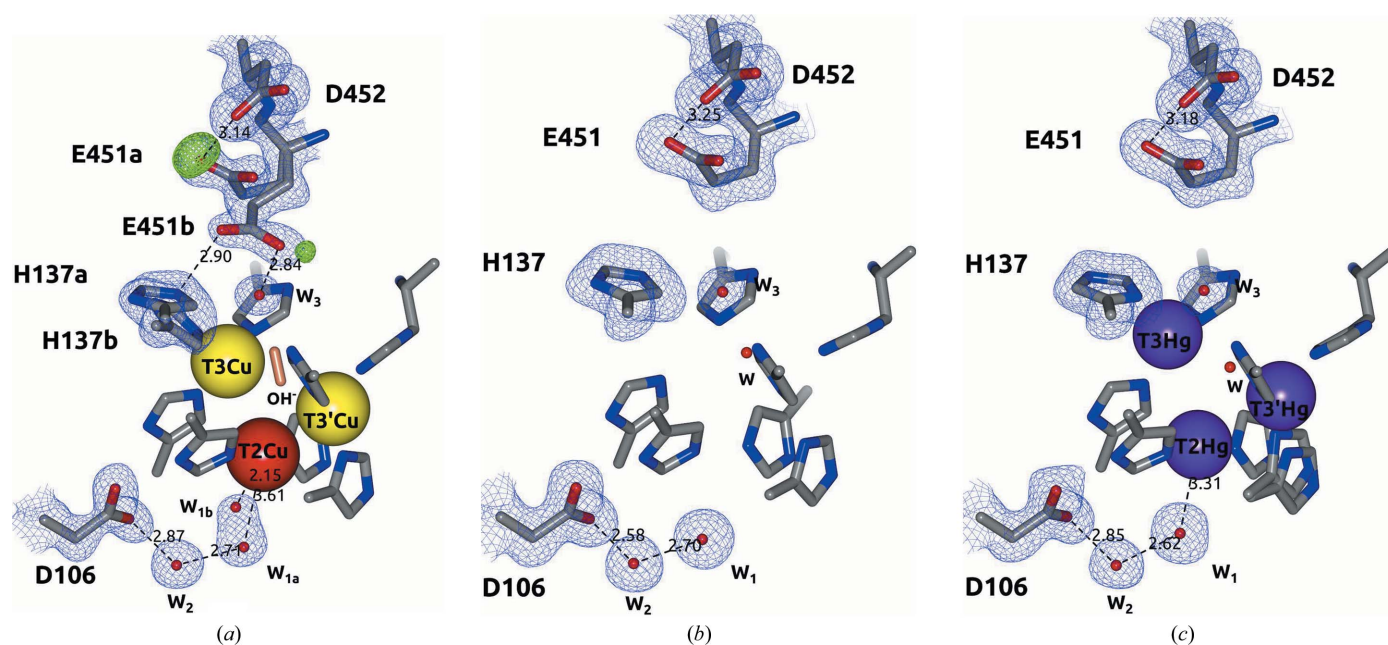


Figure 9

Proton-relay mechanism for the reduction of O<sub>2</sub> to 2H<sub>2</sub>O at the TNC of active and inactive forms of *Tth*-MCO. Note the presence of a double conformation of Glu451 in the active form of *Tth*-MCO (a) and its absence in the inactive forms apo *Tth*-MCO (b) and Hg-*Tth*-MCO-2h (c). The  $2F_o - F_c$  (blue) and  $F_o - F_c$  (green) electron-density maps are contoured at the 1σ and 3.5σ levels, respectively. Metal ions and water molecules are represented as spheres. Distances are in Å.

O<sub>2</sub> reduction, which present a H<sub>2</sub>O molecule coordinated to T2Cu (Bento *et al.*, 2005), the NI is the only species that displays an OH<sup>−</sup> coordinated to T2Cu (Fig. 1; Yoon *et al.*, 2007). As also mentioned above, it has previously been shown that these H<sub>2</sub>O/OH<sup>−</sup> species coordinated to T2Cu are always stabilized by the side chain of a highly conserved acidic residue (Asp106 in *Tth*-MCO) through a structural H<sub>2</sub>O molecule (W<sub>2</sub>; Fig. 9a; Quintanar *et al.*, 2005). Interestingly, when an H<sub>2</sub>O molecule (W<sub>1a</sub>; Fig. 9a) at a T2Cu–H<sub>2</sub>O coordination bond distance of 3.61 Å was modelled in *Tth*-MCO, a positive peak of electron density above 3.5σ in a F<sub>o</sub> – F<sub>c</sub> map was observed close to T2Cu. The latter case was explained when a second H<sub>2</sub>O molecule (W<sub>1b</sub>; Fig. 9a) with a T2Cu–OH<sup>−</sup> coordination-bond distance of 2.15 Å was modelled in a positive peak of electron density. Since an MCO structure determined by classic single-crystal data collection is an average model of different O<sub>2</sub>-reduction states and intermediates, this information suggests that populations of H<sub>2</sub>O/OH<sup>−</sup> species coordinated to T2Cu, and stabilized by Asp106 through W<sub>2</sub>, are found in the *Tth*-MCO structure (Fig. 1). Nevertheless, we deduce that the RS is the statistically relevant state for the *Tth*-MCO structure (Figs. 3a and 4a), because apart from H<sub>2</sub>O/OH<sup>−</sup> species coordinated to T2Cu, an OH<sup>−</sup> ion bridging the binuclear T3Cu–T3'Cu cluster was also observed.

To demonstrate that the proton-relay mechanism is not preserved in the inactive structures of *Tth*-MCO, three forms of the enzyme in the apo form and with Hg bound were analyzed (Figs. 3b and 3c). In fact, only one conformation of the proton-donor residue Glu451 was observed at ~7 Å to the TNC (Figs. 9b and 9c), in the equivalent position to Glu451a in active *Tth*-MCO. Notably, no electron density was found in an F<sub>o</sub> – F<sub>c</sub> map for Glu451, suggesting that Glu451 is

deprotonated and the proton-relay mechanism is not undertaken in these inactive apo and Hg-bound forms. Moreover, a single H<sub>2</sub>O molecule (W<sub>1</sub>; Figs. 9b and 9c) at the same position as W<sub>1a</sub> was observed in *Tth*-MCO (Fig. 9a), indicating that Asp106 does not play any role in O–O bond breakage in these inactive forms of *Tth*-MCO (Figs. 1b and 1c; Quintanar *et al.*, 2005).

**3.3.2. Composite structures *Tth*-MCO-C1–8 with different absorbed X-ray doses.** The proton-relay mechanism for the reduction of O<sub>2</sub> was also observed in the active composite structures *Tth*-MCO-C1–8 (Fig. 10). A positive peak of electron density in an F<sub>o</sub> – F<sub>c</sub> map at a dose of 0.2 MGy again suggests a carboxyl functional group for Glu451a O<sup>ε2</sup> of *Tth*-MCO-C1 (Fig. 10a). Furthermore, the Glu451b O<sup>ε2</sup>–H137a N<sup>ε</sup> and Glu451b O<sup>ε1</sup>–W<sub>3</sub> distances of 2.87 and 2.73 Å, respectively, do not show a preferred proton-relay path from Glu451b to TNC at this dose (Fig. 10a).

As mentioned above, protons are not involved in formation of the PI (Augustine *et al.*, 2007), and the population of electrons released by the radiolysis of water molecules in the crystals of *Tth*-MCO is relatively deficient at doses of ≤0.2 MGy. In consequence, the O<sub>2</sub> state at the TNC of the *Tth*-MCO-C1 structure was stable enough to be trapped (Figs. 7a and 10a), despite a double conformation of Glu451 being found in the proton-relay mechanism. Remarkably, the Glu451b O<sup>ε2</sup>–His137a N<sup>ε</sup> distance was reduced from 2.87 Å in *Tth*-MCO-C1 to 2.70 Å in *Tth*-MCO-C2, while the Glu451b O<sup>ε1</sup>–W<sub>3</sub> distance increased from 2.73 Å in *Tth*-MCO-C1 to 2.85 Å in *Tth*-MCO-C2 (Figs. 10a and 10b). As also mentioned above, protons and electrons are involved from the O<sub>2</sub> state (*Tth*-MCO-C1) to the RS (*Tth*-MCO-C2), so we deduce that the changes in the Glu451b O<sup>ε2</sup>–His137a N<sup>ε</sup> and Glu451b O<sup>ε1</sup>–W<sub>3</sub> distances are significantly more informative if we take into

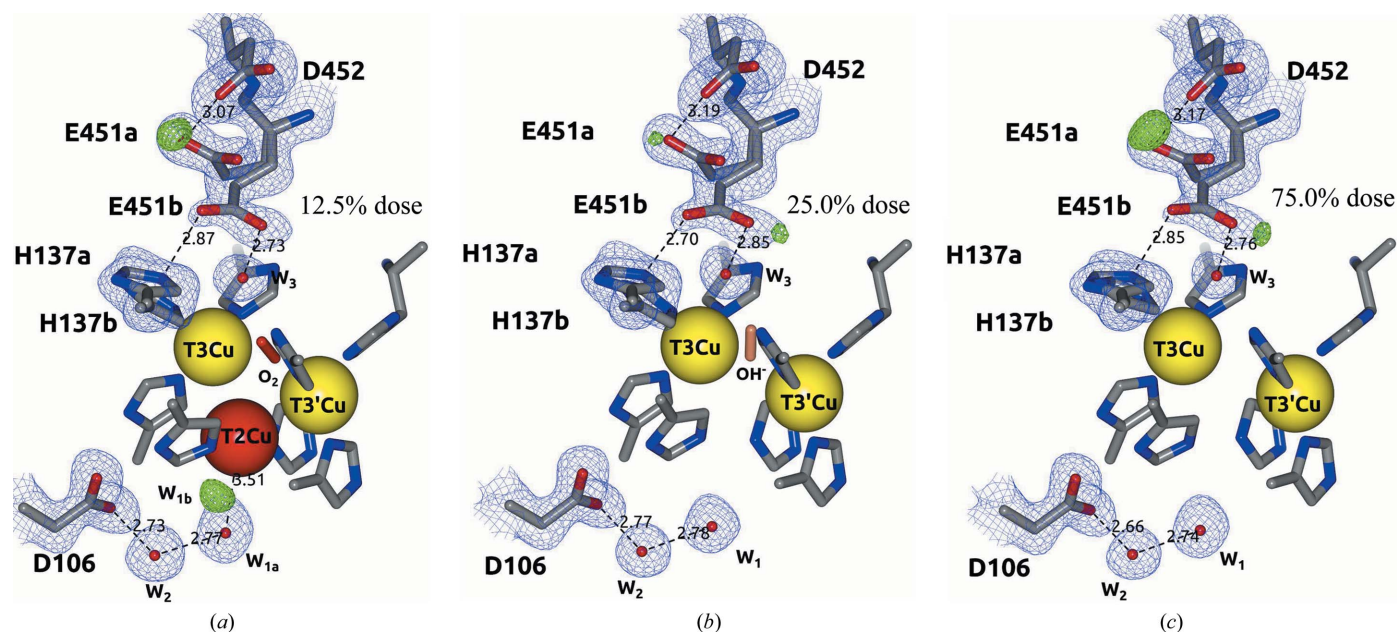


Figure 10

Proton-relay mechanism for the reduction of O<sub>2</sub> to 2H<sub>2</sub>O at the TNC of active *Tth*-MCO composite structures. (a) *Tth*-MCO-C1 (0.2 MGy; 12.5%). (b) *Tth*-MCO-C2 (25.0%). (c) *Tth*-MCO-C6 (75.0%). The 2F<sub>o</sub> – F<sub>c</sub> (blue) and F<sub>o</sub> – F<sub>c</sub> (green) electron-density maps are contoured at the 1σ and 3.5σ levels, respectively. Metal ions and water molecules are represented as spheres. Distances are in Å.

account that these two composite structures provide dynamic information for X-ray-induced reduction of O<sub>2</sub> (Figs. 10*a* and 10*b*). Additionally, the proton-relay path from Glu451b to the TNC, in which the Glu451b O<sup>ε2</sup>–His137a N<sup>ε</sup> distance is less than that between Glu451b O<sup>ε1</sup> and W<sub>3</sub>, was also observed in five of the composite structures (*Tth*-MCO-C2–5 and *Tth*-MCO-C8). This information suggests that the preferred proton-relay path in *Tth*-MCO from Glu451b to TNC seems to be as follows: Glu451b→His137a→His137b→TNC. However, the latter remains unclear, since the changes in the distances are close to the coordinate error (Table 2) and should be investigated in the future.

On the other hand, the significant absence of a positive peak of electron density in an F<sub>o</sub> – F<sub>c</sub> map for Glu451a O<sup>ε2</sup> in *Tth*-MCO-C2 (Fig. 10*b*) also suggests that a significant part of the population of protons were probably transferred to the TNC to reduce O<sub>2</sub> in *Tth*-MCO-C2 as follows: O<sub>2</sub> state (*Tth*-MCO-C1)→PI→NI→RS (*Tth*-MCO-C2). It is worth noting that except for *Tth*-MCO-C2 (Fig. 10*b*), a positive peak of electron density in an F<sub>o</sub> – F<sub>c</sub> map for Glu451a O<sup>ε2</sup> was clearly observed in all composite structures. In fact, *Tth*-MCO-C2 was the only structure determined in this work that seems to exhibit both conformations of the Glu451 residue with a carboxylate functional group.

Interestingly, a positive peak of electron density in an F<sub>o</sub> – F<sub>c</sub> map corresponding to W<sub>1b</sub> was also found close to W<sub>1a</sub> in *Tth*-MCO-C1 (Fig. 10*a*). In agreement with the information mentioned above for *Tth*-MCO (Fig. 9*a*), W<sub>1a</sub> is simultaneously coordinated to T2Cu and stabilized by Asp106 through W<sub>2</sub> in *Tth*-MCO-C1 (Fig. 10*a*). Together, this information confirms the above results around the T2Cu site for *Tth*-MCO (Fig. 9*a*), suggesting that populations of H<sub>2</sub>O/OH<sup>–</sup> species coordinated to T2Cu are also present at 0.2 MGy (Fig. 10*a*) owing to the decay of the PI to the NI in every

catalytic cycle during X-ray data collection. Naturally, only W<sub>1a</sub> was found in the *Tth*-MCO-C2–8 structures owing to the depletion of T2Cu at doses of >0.2 MGy (Fig. 10*b*). Moreover, an anisotropic displacement of His95 in the *Tth*-MCO-C2–8 structures was also observed, with a concomitant decrease in the His95 N<sup>ε</sup>–His396 N<sup>ε</sup> distance from 3.28 Å in *Tth*-MCO-C1 to 3.03 Å in *Tth*-MCO-C8.

Together, the above results revealed structural details of both the T1Cu<sup>II</sup>–Cys445–His444–T3Cu<sup>II</sup> electron-transfer pathway from T1Cu to the TNC and the proton-relay mechanism, which act together as a part of the driving force for X-ray-driven catalytic conversion of O<sub>2</sub> to 2H<sub>2</sub>O (Fig. 11).

### 3.4. Effects of absorbed radiation dose on the Cu centres of the composite structures *Tth*-MCO-C1–8

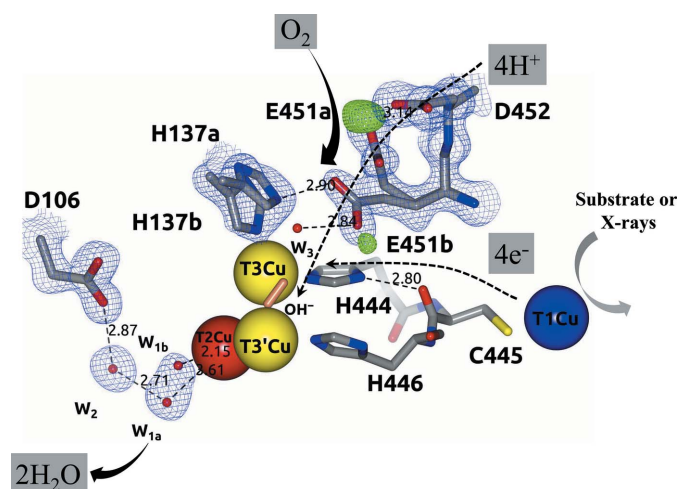
Specific structural damage to the four catalytic Cu atoms and/or the decrease in the occupancy values induced by X-ray radiation were analyzed as a function of absorbed dose in the composite structures *Tth*-MCO-C1–8. Radiation damage to T1Cu and structural changes in its local environment in the eight composite structures were negligible, with a minimal change in occupancy for this Cu atom from 1.00 (*Tth*-MCO-C1) to 0.97 (*Tth*-MCO-C8). Similarly, the T3'Cu atom of the composite structures exhibited high stability to X-rays (Fig. 7), with a final occupancy of 0.94 when the system reached 100% absorbed dose.

In contrast, the T3Cu and T2Cu atoms of the composite structures again showed a high susceptibility to radiation damage, as we observed in the *Tth*-MCO structure. In effect, the occupancy of the T3Cu atom was only 0.43 at a dose of 0.2 MGy (Fig. 7*a*). As mentioned above, this observation indicates that several factors converged once more to explain the high susceptibility of the T3Cu atom to radiation damage; for instance, the significant increase in the His137 coordination-bond distance which was observed in *Tth*-MCO (see inset in Fig. 3*a* and Fig. 9*a*) and all composite structures (Fig. 10), as well as its location in the entry channel for O<sub>2</sub> and at the end of the electron-transfer pathway from T1Cu to TNC (Fig. 11).

On the other hand, the T2Cu atom showed an occupancy of only 0.16 for *Tth*-MCO-C1 (Fig. 10*a*) and was totally depleted at a dose of >0.2 MGy (Fig. 10*b*; *Tth*-MCO-C2). In agreement with this observation, partial occupancy and depletion of the T2Cu atom has previously been reported for the structure of laccase from *Corioloropsis gallica* (Cg L; De la Mora *et al.*, 2012). In the latter case, a high susceptibility of T2Cu to radiation damage was shown and a complete depletion of T2Cu at a dose between 0.6 and 4.0 MGy was found. Interestingly, T2Cu of *Tth*-MCO displays a higher susceptibility to X-rays than T2Cu of Cg L, highlighting the importance of taking radiation effects into account in biochemical interpretations of an MCO structure.

## 4. Conclusions

We have determined 12 crystallographic structures of *Tth*-MCO in active and inactive forms and with different absorbed



**Figure 11**

Proton-relay mechanism and electron-transfer pathway from T1Cu to TNC for the X-ray-driven catalytic conversion of O<sub>2</sub> to 2H<sub>2</sub>O, taking the active *Tth*-MCO structure as a reference. The 2F<sub>o</sub> – F<sub>c</sub> (blue) and F<sub>o</sub> – F<sub>c</sub> (green) electron-density maps are contoured at the 1σ and 3.5σ levels, respectively. Metal ions and water molecules are represented as spheres. Distances are in Å.



X-ray doses at 1.5–1.8 Å resolution in order to study the X-ray-induced reduction of O<sub>2</sub> to 2H<sub>2</sub>O at the TNC of the enzyme and the proton-relay mechanism involved in this catalytic reaction. To the best of our knowledge, this is the first structural evidence in the MCO family that demonstrates the dynamic behaviour of the proton-donor residue Glu451 in the protonation events at the TNC. In the case of active *Tth*-MCO structures, the double conformation of Glu451 shows how protons are channelled to the TNC for the reduction of O<sub>2</sub> and the importance of the Asp106 residue for the decay of the PI to the NI, but not for protonation events at the TNC. Remarkably, in apo and Hg-bound structures dynamic behaviour of the proton-donor Glu451 residue was not observed, demonstrating that the proton-relay mechanism is not undertaken in these inactive forms.

X-ray-induced reduction experiments of *Tth*-MCO crystals allowed the systematic study of O<sub>2</sub> reduction at the TNC in the crystalline state. Accordingly, at a low absorbed radiation dose of 0.2 MGy the active *Tth*-MCO-C1 composite structure showed an O<sub>2</sub> molecule coordinated at the TNC, demonstrating that in the absence of sufficient electrons in the crystalline state the RS is not the only state that is sufficiently stable to be trapped in *Tth*-MCO structures and possibly found in solution, but the statistically relevant state has since been observed in six composite structures (*Tth*-MCO-C2–5, *Tth*-MCO-C7 and *Tth*-MCO-C8) as well as in the *Tth*-MCO structure determined by classical single-crystal data collection. Additionally, when the dose was increased the FR state found in the *Tth*-MCO-C6 composite structure was also stable enough to be trapped. However, we were unable to isolate composite structures of the PI and the NI coordinated at the TNC owing to their decay rates. Together, these results also demonstrate the high susceptibility of T2Cu and T3Cu to radiation damage, with the former being totally depleted at radiation doses higher than 0.2 MGy.

Finally, a 16-residue methionine-rich β-hairpin motif over the T1Cu site, which appears to have interesting biological implications and is probably involved in extra copper(I)-binding sites, as previously described for CueO (Roberts *et al.*, 2003; Kataoka *et al.*, 2007; Singh *et al.*, 2011), was observed in all structures determined in this work. Both a high concentration of methionines between the top of the β-hairpin motif and T1Cu and a fifth Hg-binding site found in the β-hairpin motif of Hg-bound structures demonstrate that extra metal-coordinating sites may be found in this methionine-rich region; however, their biological role has to be proposed in the future.

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