

Description of a Non-Canonical Mn(II)-Oxidation Site in Peroxidases

Elena Fernández-Fueyo,^{§,‡} Irene Davó-Siguero,[§] David Almendral,[§] Dolores Linde,[§] Maria Camilla Baratto,^{#,⊥} Rebecca Pogni,^{#,⊥} Antonio Romero,[§] Victor Guallar,^{¶,†} and Angel T. Martínez*,§

[§]Centro de Investigaciones Biológicas, Consejo Superior de Investigaciones Científicas (CSIC), E-28006 Madrid, Spain

[#]Department of Biotechnology, Chemistry and Pharmacy, University of Siena, I-53100 Siena, Italy

¹Consorzio per lo Sviluppo dei Sistemi a Grande Interfase (CSGI), 50019 Florence, Italy

[¶]Barcelona Supercomputing Center, E-08034 Barcelona, Spain

[†]Institució Catalana de Recerca i Estudis Avancats (ICREA), E-08010 Barcelona, Spain

Supporting Information

ABSTRACT: A dye-decolorizing peroxidase (DyP) from Pleurotus ostreatus (PosDyP4) catalyzes the oxidation of Mn^{2+} to Mn^{3+} , in the presence of H₂O₂, with an efficiency similar to the well-known manganese peroxidases and versatile peroxidases from this and other white-rot fungi. PosDyP4 has been overexpressed in Escherichia coli as an active enzyme, and its crystal structure has been solved at 1.56 Å resolution. A combination of substrate diffusion simulations on the solved structure using the PELE software, electron paramagnetic resonance, and site-directed mutagenesis led to identification of the residues involved in Mn²⁺ oxidation. The oxidation site in PosDyP4 is different than the conserved site in the other Mn-oxidizing peroxidases mentioned above, and it includes four acidic residues (three aspartates and one



glutamate) located at the surface of the protein. Moreover, since the Mn²⁺ ion is not in direct contact with the heme propionates, a tyrosine residue participates in the electron transfer to the cofactor, being the only essential individual residue for PosDyP4 oxidation of the metal ion. The four acidic residues contribute to Mn²⁺ binding in different extents, with the glutamate also involved in the initial electron transfer to the key tyrosine, as confirmed by the >50-fold decreased k_{cat} after removing its side-chain carboxylic group. A second electron transfer pathway operates in PosDyP4 for the oxidation of aromatics and dyes starting at a surface tryptophan, as reported in other fungal and prokaryotic DyPs, and connecting with the final part of the Mn^{2+} oxidation route. Both tryptophanyl and tyrosyl radicals, potentially involved in catalysis, were detected by electron paramagnetic resonance of the native enzyme and its tryptophan-less variant, respectively.

KEYWORDS: Mn-oxidation site, dye-decolorizing peroxidase, crystal structure, ligand diffusion, PELE, electron transfer, QM/MM calculations, EPR spectroscopy

INTRODUCTION

Peroxidases (EC1.11.1) can oxidize a variety of substrates using peroxides as final electron acceptors. They were traditionally classified into animal and nonanimal peroxidases, the second group being subdivided into three classes according to their origin, with fungal peroxidases forming class II. However, the recent discovery of new types of secreted fungal peroxidases has led to the establishment of two novel (super)families: the heme-thiolate peroxidases (HTP), in which classical chloroperoxidase and novel unspecific peroxygenases are included; and the dye-decolorizing peroxidases (DyP),²⁻⁵ classified together with chlorite dismutases and Escherichia coli EfeB proteins in the CDE superfamily.⁶

The first DyP was discovered in a Bierkandera adusta culture described as Geotrichum candidum.⁷ Its crystal structure was available in 2011,⁸ followed by that of Auricularia auriculajudae DyP.9 Their tertiary structure revealed a very different topology from the other (super)families mentioned above. Nevertheless, the DyP active site presents some similarities with class II peroxidases, such as a histidine residue as heme iron ligand,¹⁰ while a cysteine occupies this position in HTPs.² However, the second histidine at the opposite side of heme in classical peroxidases is not conserved in DyPs, which generally present an aspartic acid at this position.^{5,11} Moreover, contrarily to class-II peroxidases, DyPs are not exclusive of fungi, with types A-C corresponding to prokaryotic enzymes and fungal DyPs preferentially included in type D.¹²

Received: June 13, 2018 Revised: July 25, 2018 Published: July 31, 2018

Since the discovery of the first DyP, a number of other family members has been studied, as wild type or heterologously expressed enzymes.⁵ After in vitro activation of fungal DyPs expressed as inclusion bodies,¹³ *Pleurotus ostreatus* DyP4 (*Pos*DyP4) and a few other fungal DyPs have been recently expressed in *E. coli* as active proteins.^{14,15} Prokaryotic DyPs are easier to be expressed in *E. coli* facilitating different mechanistics and related studies.^{16–19}

DyPs present a broad substrate specificity, being able to oxidize substituted phenols, azo dyes, and even lignin model compounds, among other substrates, although nonphenolic aromatics with very low activity (see Linde et al.⁵ and Loncar et al.²⁰ and references herein). Nevertheless, anthraquinonoid dyes, used in first DyP descriptions,⁷ appear as their best substrates. Additionally, the ability to oxidize Mn²⁺ to Mn³⁺ has been reported in DyPs from *Amycolatopsis* sp. 7Siv2 (*AspDyP2*),²¹ *P. ostreatus* (*PosDyP4*),¹⁵ and very recently from *Irpex lacteus*,²² as well as, with much lower activity levels, in a few other bacterial^{21,23-26} and one more fungal²⁷ DyP. Interestingly, *PosDyP4* oxidizes Mn²⁺ with similar efficiency to the fungal ligninolytic manganese peroxidases (MnP) and versatile peroxidases (VP).¹⁵

Despite the variety of substrates reported, the ecophysiological role of DyPs still remains unclear. Their ability to oxidize lignin related compounds, together with their higher abundance in white-rot (ligninolytic) fungi compared with other basidiomycetes,⁵ and their widespread presence in forest soil fungal transcriptomes,²⁸ support a contribution to degradation of lignin and lignin-derived compounds. Furthermore, *PosDyP4* is one of the most abundant enzymes when *P. ostreatus* grows on lignocellulosic biomass.¹⁵ In the above context, DyPs have been described as bacterial "lignin peroxidases",²⁹ although they preferentially act on phenolic technical lignins and lignin-derived substrates.^{5,30–32}

In the present study, we identified and characterize the DyP Mn^{2+} -oxidation site thanks to a combination of different techniques (such as X-ray diffraction, computational calculations, electron paramagnetic resonance, and directed mutagenesis). Interestingly, the site was determined to be different than the canonical Mn^{2+} -oxidation one found in class-II peroxidases because of its superficial location and involvement of a long-range electron pathway to heme.

MATERIAL AND METHODS

DyP Production in E. coli. The PosDyP4 coding sequence (ref. no. 1069077 from the P. ostreatus genome at JGI, https:// genome.jgi.doe.gov/PleosPC15 2) was cloned in the pET15b vector carrying a N-terminal His-tag sequence (Novagen, Darmstadt, Germany), and the resulting plasmid (pET15b-1069077) was used for expression in E. coli BL21(DE3)pLysS (Novagen).¹⁵ Cultures were grown for 60 h at 16 °C in ZYM-5052 autoinducing medium³³ in the presence of 100 μ M hemin. Cells were harvested by centrifugation at 8000 rpm for 10 min at 4 °C and suspended in 100 mL of lysis buffer (10 mM Tris-HCl, pH 7.5, containing 10 mM imidazole, and 300 mM NaCl) supplemented with 2 mg \cdot mL⁻¹ lysozyme (Sigma-Aldrich, Steinheim, Germany) and DNase I (Roche Diagnostics, Mannheim, Germany). After 1 h, the suspension was sonicated, debris was removed by centrifugation at 20 000 rpm for 4 h, and the supernatant was concentrated (Amicon 10-kDa-cutoff; Millipore, Darmstadt, Germany).

The wild-type recombinant (hereinafter native) DyP and the mutated variants were purified in one single step using a 5 mL

HisTag column, at a flow rate of 5 mL·min⁻¹ of the above buffer in an ÄKTA HPLC system (GE Healthcare Bio Sciences AB, Uppsala, Sweden). After 15 mL, the retained proteins were eluted with a 0–50% gradient of 1 M imidazole in 50 mL, followed by 50–100% gradient in 5 mL and 100% NaCl in 15 mL. Peroxidase activity was followed by 2,2-azinobis(3ethylbenzothiazoline-6-sulfonic acid) (ABTS) oxidation, as described below. The appropriate fractions were pooled, concentrated, and dialyzed against 10 mM sodium tartrate (pH 5).

Purification was confirmed by sodium dodecyl sulfatepolyacrylamide gel electrophoresis (SDS-PAGE) in 12% gels stained with Coomassie brilliant blue R-250 (Sigma-Aldrich) (Figure S1), although some minor contaminating proteins were observed for some of the variants (due to the single-step purification procedure). The UV-visible spectra of native *PosDyP4* and its Mn²⁺-oxidation site variants (Figure S2) confirmed the incorporation of heme, with a characteristic Soret band at 406–407 nm. The above agrees with (i) Reinheitszahl (Rz, $A_{\text{Soret-band}}/A_{280 \text{ nm}}$) ratios of 1.4–1.8 (except for the E345A/D352A/D354A variant with a 1.1 ratio); and (ii) molar heme content, estimated by the pyridine ferrohemochrome method,¹⁷ of 0.9 for *PosDyP4* and slightly lower (0.7–0.9) for the variants, due to minor contaminating proteins.

Crystal Structures. Crystallization trials were carried out by the sitting drop vapor diffusion technique, in 96-well MRC2 plates (Swissci MRC, Molecular Dimensions, Suffolk, England) using a Cartesian Honeybee System (Genomic Solutions, Irvine, U.S.A.) and the commercial screenings from Emerald (Wizard Screens I, II and III) and Jena Biosciences (JBScreen Classic Kits 1–10). Different MnCl₂ concentrations were included in the crystallization trials. Drops consisted of 0.2 μ L of protein solution (10 mg/mL in 10 mM sodium tartrate, pH 5.0) and 0.2 μ L of precipitant and were equilibrated against a 50 μ L reservoir solution, at 295 K. Crystals suitable for X-ray data collection were obtained in 0.2 M 2-(*N*-morpholino)ethanesulfonic acid (MES), pH 5.75, containing 25% polyethylene glycol (PEG) MME 2000. Furthermore, crystal soaking in MnCl₂ solutions was attempted to solve enzyme:Mn²⁺ structures.

Crystals were mounted in nylon loops and flash-frozen in liquid nitrogen in the mother liquor using Paratone-N (Hampton Research, Aliso Viejo, U.S.A.) as cryoprotectant. All diffraction data were collected at 100 K on beamline ID30B at the ESRF (Grenoble, France) using a wavelength of 0.9762 Å and a Pilatus 6 M detector. Diffraction data were indexed and integrated using XDS,³⁴ and scaled and merged using Aimless from CCP4 program suite.³⁵

The structure was solved by molecular replacement using the crystal structure of *B. adusta* DyP (PDB: 3afv) as the search model and the program MOLREP in CCP4 suite.³⁵ The final model was obtained by consecutive rounds of refinement with RefMac, in CCP4 suite,³⁵ and alternating manual model building with Coot³⁶ using σ_A weighted $2F_o-F_c$ and F_o-F_c electron density maps. Solvent molecules and MES were added automatically and visually inspected for chemically plausible positions. A total of 5% of reflections was used to calculate the $R_{\rm free}$ value throughout the refinement process. The structures were validated with MolProbity.³⁷

X-ray data collection and refinement statistics are summarized in Table S1. The refined model consists of a partial polypeptide chain of 484 amino acids (residues 6–489),



Figure 1. *Pos*DyP4 crystal structure. (A) Ribbon representation of the overall fold with α -helices in magenta (H1 to H28), β -strands in blue forming two β -sheets (B1–B6 and B7, B8 and B11), random coil in green, C and N termini of the solved structure (Ct and Nt) included, and the heme cofactor as CPK-colored sticks (PDB: 6fsk). (B) Detail of the heme pocket with His334, Asp196, Arg360, and water134 (at the sixth coordination position of heme iron), and Trp405 and Tyr339 at 7.2 and 3.2 Å from the heme cofactor, respectively, as CPK-colored sticks.

in which the first five and the last 15 residues were not visible in the electron density maps, one heme group, one MES, and 1119 water molecules. Figures were produced with PyMOL (Schrödinger, Portland, OR, U.S.A.). The coordinates and structure factors have been deposited at the Protein Data Bank with accession code 6fsk.

Molecular Simulations. The solved crystal structure (6fsk, chain A) was used for *Pos*DyP4 molecular simulations. The system was prepared at pH 4.5 (optimal activity) with the protein preparation wizard from Schrödinger, resulting in aspartic acids 32 and 241 being protonated; histidines 49, 138, 148, 168, 191, and 220 being protonated in their N ε ; and histidines 380 and 299 being double protonated (positively charged). Crystallographic water molecules were retained because of their (potential) importance in Mn²⁺ coordination.

For substrate diffusion molecular mechanics (MM) simulations with PELE, heme charges were obtained from a quantum mechanics/molecular mechanics (QM/MM) minimization using QSite,³⁸ at the DFT M06-L(lacvp*)/OPLS level of theory. Based on previous experience in manganese heme-bound systems,³⁹ the charge of the Mn²⁺ ion was set to 1.2. The ion was then placed on five different initial positions, all of them in the bulk solvent at distances >25 Å from the heme iron. The new adaptive-PELE software⁴⁰ was used to study Mn²⁺ diffusion and binding. The adaptive protocol improves sampling in PELE by running multiple short simulations (epochs) where the initial conditions in each of them are selected through a reward function aiming at sampling nonvisited areas. In particular, three simulations of 256 processors with 20 epochs, of 12 PELE steps each, were used. We should remark that along all simulations, the ligand

moves freely, without any predetermined bias. Interaction energies, in kcal/mol, were derived as Eab – (Ea + Eb) where Eab is the total energy of the complex, Ea the energy of the "receptor" (protein + crystallographic waters), and Eb the energy of the Mn²⁺ ion; all of them obtained at the OPLS2005 level of theory with a surface GB implicit solvent model.

Oxidation QM/MM simulations were performed also with Qsite.³⁸ The QM region included: the heme (compound I, i.e. Fe⁴⁺=O and porphyrin cation radical complex), Trp405, Mn²⁺ ion, residues 215, 334 to 345, and 352 to 357, and two water molecules coordinated to the metal ion, for a total of 502 QM atoms. They were modeled at the DFT M06-L(lacvp*) level of theory, the reminder of the system being described with the OPLS-AA force field. To facilitate convergence of compound-I's electronic wave function, the triplet spin state was selected.⁴¹ For the QM/MM e-pathway calculations,⁴² monitoring the electron transfer route, the heme group, Mn²⁺ ion and its directly coordinated residues were subtracted from the QM region. Three iterations of the procedure were performed, where Trp405 and both Trp405 and Tyr339 were subtracted from the QM region in the second and third iterations, respectively.

Electron Paramagnetic Resonance (EPR) Measurements. Continuous-wave EPR X-band (9.38/9.39 GHz) measurements were performed with a Bruker E500 Elexsys Series using the Bruker ER 4122SHQE cavity, equipped with an Oxford Instruments helium continuous flow cryostat (ESR900). Enzyme activation was carried out by adding H_2O_2 in a *Pos*DyP4/peroxide molar ratio of 1:8, with a final concentration of 0.25 mM enzyme in tartrate buffer, pH 3. H_2O_2 was added directly in a 3.5 mm EPR tube, and the reaction time before freezing was ~ 10 s. The *g*-values were calculated with a strong pitch sample as reference, and an error of ± 0.0002 was estimated.

Site-Directed Mutagenesis. The D196A, D215V, Y339A, E345A, D352A, D354A, and W405S simple mutations, and the D352A/D354A double and E345A/D352A/D354A triple mutations were introduced in the *Pos*DyP4 sequence by PCR using the expression plasmid pET15b-1069077 (see above) as template, and the QuikChange kit from Stratagene (La Jolla, CA, U.S.A.). The direct and reverse primers for PCR (Table S2) were synthesized. To obtain the triple variant, the plasmid harboring the D352A/D354A mutations was used as a template and combined with the primer E345A.

The PCR reaction (50 μ L volume) was carried out in an Eppendorf (Hamburg, Germany) Mastercycler pro-S thermal cycler using 20 ng of template DNA, 500 μ M each dNTP, 125 ng of direct and reverse primers, 2.5 units of *Pfu*Turbo polymerase (Stratagene), and the manufacturer's buffer. Reaction conditions included: (i) a start cycle of 1 min at 95 °C; (ii) 18 cycles of 50 s at 95 °C, 50 s at 55 °C, and 10 min at 68 °C; and (iii) a final cycle of 10 min at 68 °C. The mutated genes were expressed in *E. coli*, and the variants were purified as described above for the wild gene and native DyP.

Enzyme Kinetics. The kinetic constants of *Pos*DyP4 and its variants were estimated from absorbance changes during substrate oxidation in 0.1 M tartrate (optimal pH values) at 25 °C in a Biomate5 spectrophotometer (Thermo Scientific). The reactions were initiated by addition of H_2O_2 to a concentration of 0.25 mM.

Oxidation of Mn^{2+} was followed at pH 4.5 by monitoring formation of Mn^{3+} -tartrate complex (ε_{238} 6.5 mM⁻¹·cm⁻¹). Reactive Blue 19 (RB19) and ABTS oxidation were assayed at pH 3.5 and monitored by RB19 disappearance (ε_{595} 10 mM⁻¹· cm⁻¹) and formation of ABTS cation radical (ε_{436} 29.3 mM⁻¹· cm⁻¹), respectively. Reactive Black 5 (RB5), 2,6-dimethoxyphenol (DMP) and veratryl alcohol (VA) oxidation were assayed at pH 3, and monitored by RB5 disappearance (ε_{598} 30 mM⁻¹·cm⁻¹), dimeric coerulignone (ε_{469} 55 mM⁻¹·cm⁻¹) formation, and veratraldehyde (ε_{310} 9.3 mM⁻¹·cm⁻¹) formation, respectively.

Means and standard errors for Michaelis constant (K_m) and turnover (k_{cat}) values were obtained by nonlinear least-squares fitting to the Michaelis–Menten model. Fitting of these constants to the normalized equation $v = (k_{cat}/K_m)[S]/(1 + [S]/K_m)$ yielded the catalytic efficiencies (k_{cat}/K_m) with their standard errors. Biphasic kinetics was observed for Mn^{2+} oxidation by some *Pos*DyP4 variants enabling calculation of two sets of kinetic constants, as found also for other DyPs.^{13,43}

RESULTS AND DISCUSSION

Solving the Crystal Structure of PosDyP4. We have solved the crystal structure of PosDyP4 at 1.56 Å resolution, after the heterologous expression in *E. coli*, as an active enzyme, of the sequence obtained from the *P. ostreatus* genome available at the DOE JGI.¹⁵ All attempts to solve the structure of native PosDyP4 in complex with Mn²⁺ were unsuccessful (an electron-density map, corresponding to the Asp215, Tyr339, Glu345, Asp352, Asp354, and water31 region discussed below, is shown in Figure S3). PosDyP4 shows the overall topology (Figure 1A) characteristic of the DyP family and other members of the CDE superfamily.^{5,6} Each of the two ferredoxin-like domains is formed by a four-stranded antiparallel β -sheet (strands B1–B6 and B7–B11 in the C- terminal and N-terminal domains, respectively) and several helices (up to a total of 28, when short helical regions are included). Two additional β -strands (B9 and B10) are located at the bottom of the N-terminal domain, which also harbors the heme cofactor.

A structural search with the *Pos*DyP4 molecular structure at the DALI server⁴⁴ shows the highest fitting (Z value) and identity (id) with the reported crystal structures of: (i) the only two fungal DyPs whose structure had been reported, from *B. adusta* and *Auricularia auricula-judae* (Z: > 51, id: >43%); (ii) two structurally related DyPs from *Amycolatopsis* sp. and *Anabaena* sp. (Z: 39, id: 28–29%); and (iii–iv) two other groups of prokaryotic DyPs and CDE- proteins with 11 (Z: 26–28, id: 15–20%) and two (Z: 12, id: 7–9%) members (see Table 1 with PDB entries and additional details).

Table 1. Crystal Structures Related to *Pos*DyP4 from Search at the DALI Server⁴⁴ Including 2 Fungal Followed by 15 Prokaryotic Proteins^{*a*}

		PDB	Å	Z	%
	Bjerkandera adusta	3afv	1.4	54	48
	Auricularia auricula-judae	4au9	2.1	51	43
	Amycolaptosis sp.	4g2c	2.3	39	28
	Anabaena sp.	5c2i	1.9	39	29
	Thermobifida fusca	5fw4	1.8	28	20
	Shewanella oneidensis	2iiz	2.3	28	17
	Escherichia coli	3072	2.0	28	20
	Thermomonospora curvata	5jxu	1.8	28	18
	Streptomyces lividans	5mjh	1.5	27	19
	Streptomyces coelicolor	4gt2	1.8	27	19
	Bacteroides thetaiotaomicron	2gvk	1.6	27	16
	Enterobacter ligninolyticus	5vj0	1.9	26	28
	Rhodococus jostii	3veg	2.4	27	15
	Thermobifida cellulosilytica	4gs1	1.7	26	19
	Vibrio cholera	5de0	2.2	26	19
	Candidatus Nitrospira difluvii	3nn2	1.9	12	9
	Thermus thermophilus	1vdh	2.0	12	7
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"See Figure S4 for 3D models. PDB entries, resolution (A), Z value and identity with *Pos*DyP4 (%) are provided.

Although the sequence identity is low, around 50% among fungal members and below 30% between the fungal and prokaryotic members, there are noteworthy similarities in the tertiary structure of the above DyPs and related CDE proteins (Figure S4). A peculiarity of the *P. ostreatus, Amycolatopsis* sp. and *Anabaena* sp. DyPs is a prominent helix-8 (red asterisks in Figure S4).^{12,21} Interestingly, it has been proposed to include these two prokaryotic proteins in a new DyP type together with the fungal DyPs.⁴⁵ AspDyP2 and *Pos*DyP4 also share its ability to efficiently oxidize Mn²⁺, as described below.

In contrast with class-II peroxidases, which have a main heme access channel (in front of the heme δ -meso carbon) and eventually a small second channel in front of one of the heme propionates (Mn²⁺ channel),¹⁰ *Pos*DyP4 only has a narrow access channel connecting the solvent with the upper side of the heme pocket (Figure S5) and lacks a channel at the vicinity of the heme propionates. This led to the exposed Mn²⁺ oxidation site described here.

The heme pocket region in *Pos*DyP4 (Figure 1B) shows several conserved residues including His334, which appears as a ligand of the heme iron, and Asp196 occupying a mirror position at the opposite side of the cofactor together with neighboring Arg360, and Trp405 and Tyr339 mentioned below at ~7 and ~3 Å from the heme, respectively. Asp196 and Trp405 are homologous to residues involved in catalysis by other fungal DyPs.⁵ To confirm their role in *Pos*DyP4, they were substituted by alanine and serine, respectively, and the activities (on ABTS, DMP, Mn^{2+} , RB5, RB19 and VA) were analyzed on different substrates (data not shown in tables). Asp196, located over the heme plane, is essential in *Pos*DyP4, since no activity was detected in the D196A variant. This is in agreement with a role in the heterolytic cleavage of H₂O₂ to form compound I, as reported in other fungal^{5,11} and some bacterial DyPs.²⁷

Mutation of the solvent-exposed Trp405 (W405S variant) suppressed VA oxidation and decreased the activity toward all other substrates with the only exception of Mn^{2+} , whose oxidation was unaffected. This tryptophan would be homologous to *AauDyP* Trp377 involved in oxidation of high redox-potential substrates, such as VA, by formation of a solvent-exposed aromatic radical, followed by long-range electron transfer from the substrate to the heme.⁴³ Formation of a similar catalytically active tryptophanyl radical has been recently demonstrated also for *Thermomonospora curvata* DyP,¹⁸ while reactive tyrosyl radicals have been reported in other prokaryotic DyPs.⁴⁶ Interestingly, the Trp405 radical was detected by EPR (see below) after peroxide activation of *Pos*DyP4, and the W405S variant was used in EPR experiments, as described below.

Diffusion Simulations Revealing an Unusual Mn^{2+} Binding Site. Attempts to obtain the crystal structure of a *Pos*DyP4:Mn²⁺ complex failed. Therefore, to identify the Mn²⁺-oxidation site, a free (nonbiased) exploration of Mn²⁺ diffusion with adaptive PELE⁴⁰ was performed by mapping the entire enzyme surface. Figure 2 shows the protein–ligand interaction energy profile against the distance from the Mn²⁺ ion to the Asp354 mentioned below.

While the ion finds multiple small cavities (local minima) in various sites of the surface, all simulations indicated a highly acidic area (Figure 3) where Mn^{2+} binds preferentially coordinated by the Glu345, Asp352 and Asp354 carboxyl groups at ~2 Å distance (Figure 4). These three acidic



Figure 2. Results of Mn^{2+} diffusion with adaptive-PELE. The protein–ligand interaction energy is shown against the distance to the side-chain carboxyl CG atom in Asp354. Each color (symbol) corresponds to a different trajectory. The best energy positions were located near Asp354 (red box), and one of them is shown in Figures 3 and 4



Figure 3. Solvent access surface of *Pos*DyP4, colored by electrostatic potential (blue, positive; and red, negative) showing the highly acidic region (with Asp215, Glu345, Asp352, and Asp354) surrounding the Mn^{2+} ion (green sphere) after its diffusion by PELE (on PDB: 6fsk).



Figure 4. Four acidic residues and two water molecules (red spheres) involved in the coordination of Mn^{2+} (purple sphere) by *Pos*DyP4 and Tyr339 located between the Mn^{2+} -binding site and the heme cofactor (residues and heme as CPK-colored sticks), in one snapshot from the best ligand-binding region in PELE simulations (red square in Figure 3). Some distances are indicated in Å.

residues, along with Asp215 (at a longer distance from the Mn^{2+} ion) and two water molecules, form an octahedral coordination environment. Tyr339, another residue involved in *PosDyP4* oxidation of Mn^{2+} , as shown by site-directed mutagenesis, is far from the metal ion (4.9 Å) but at 2.6 Å from Glu345 (Figure 4) bridging the Mn^{2+} -binding site with the heme cofactor. Comparison between the lowest interaction energy structure and the (apo) crystal structure without Mn^{2+} reveals some side-chain rearrangement in several amino acids during binding of the metal ion (Figure S6).

Electron Transfer Route Mapped by QM/MM Simulations. Figure 5A shows the total spin density obtained in QM/MM modeling of the Mn^{2+} oxidation process, with over 500 atoms being included in the quantum calculations. As in previous models of cytochromes's compound I,⁴¹ we observe two unpaired electrons in the iron-oxo moiety, localized in the d_{xz}/d_{yz} and p_x/p_y orbitals, respectively. More relevant is the nature of the third unpaired electron, largely localized in the Mn^{2+} d orbitals. This indicates that compound I is capable of extracting one electron from the Mn^{2+} ion. In addition, we observe a minor component on Trp405, supporting the role of this residue as a catalytic radical, as shown by EPR and directed mutagenesis.



Figure 5. QM/MM simulations of Mn^{2+} oxidation and electron transfer to heme. (A) Spin density in the QM/MM calculations for Mn^{2+} oxidation in the triplet state. Residues in the QM region are shown in thin sticks, together with a backbone trace in a cartoon style. The heme is shown in green and a thicker stick format. (B) Electron transfer route from the QM/MM e-pathway analysis. The spin densities of the first three iterations are shown (in order): Trp405 (blue), Tyr339 (green), and Thr338 (orange). These three regions and the environment of the donor (Mn^{2+}) and acceptor (heme) are shown in sticks; the remainder of the system is in wire style. The enzyme backbone is shown in a cartoon style.

Figure 5B shows the results of applying the QM/MM epathway procedure, where we aim at mapping the electron transfer pathway. In this type of simulation, we do not include the donor and acceptor (heme and Mn^{2+}) in the QM region, focusing instead in the transfer region.⁴² The results show the first three iterations of the procedure: Trp405 in blue, Tyr339 in green, and Thr338 in orange. Thus, the analysis indicates that the Mn^{2+} oxidation and the above-mentioned Trp405 electron transfer pathways might converge in the backbone of Thr338, which is located next to the axial histidine. A close look at the carbonyl backbone of this threonine indicates a large hydrophobic environment, raising the energy levels of its π orbitals and facilitating the electron transport.

Protein Radical Formation As Shown by EPR. In the EPR spectrum of the native *Pos*DyP4 resting state, the heme Fe^{3+} is in a rhombically distorted high spin state with an almost undetectable amount of low spin state (Figure S7, black line). After the addition of H₂O₂ (Figure S7, red line), the Fe³⁺ signal decreased and the appearance of a radical species was observed, a close up of which is shown in Figure 6A. The line shape and the *g* value equal to 2.0041 are similar to that reported for the H₂O₂-activated *Aau*DyP, which was assigned to a tryptophanyl radical in combination with other minor radical species.^{43,47}



Figure 6. Twenty mT scan width X-band EPR spectra of 0.25 mM native *Pos*DyP4 (A) and W405S variant (B) in tartrate buffer, pH 3, after the addition of 8 equiv of H_2O_2 . Conditions were 5 mW microwave power, 0.2 mT modulation amplitude, and $\nu = 9.39$ (A) or 9.38 (B) GHz. The EPR spectra are normalized (see Figures S7 and S8 for entire spectrum, before and after H_2O_2 addition, and for comparison of non-normalized signals, respectively).

Table 2. Steady-State Kinetic Parameters, Including K_m (μ M), k_{cat} (s⁻¹), and K_m/k_{cat} (s⁻¹·mM⁻¹), for Oxidation of Mn²⁺ by *Pos*DyP4 and Seven Variants Mutated at the Mn²⁺-Oxidation Site (with Two Sets of Constants in Three of Them, Characterized by Higher and Lower Affinity for Mn²⁺)^a

		DyP4	Y339A	D215A	E345A	D352A	D354A	D352A/D354A	E345A/D352A/ D354A
Mn ²⁺ higher affinity	$K_{\rm m}$	290 ± 30	-	4400 ± 1000	630 ± 90	350 ± 120	260 ± 10	300 ± 40	1400 ± 160
	$k_{\rm cat}$	56 ± 2	0	28 ± 2	1 ± 0	13 ± 2	10 ± 0	25 ± 1	11 ± 0
	$k_{\rm cat}/K_{\rm m}$	200 ± 20	0	6 ± 0	2 ± 0	36 ± 0	39 ± 1	85 ± 5	8 ± 0
Mn ²⁺ lower affinity	$K_{\rm m}$	-	-	-	-	3500 ± 900	4300 ± 700	2300 ± 700	-
	$k_{\rm cat}$	0	0	0	0	40 ± 3	39 ± 2	73 ± 5	0
	$k_{\rm cat}/K_{\rm m}$	0	0	0	0	11 ± 1	9 ± 1	32 ± 4	0
^{<i>a</i>} Means and 95% co	onfidence	limits from	reactions	at 25 °C in 0.1	M tartrate,	pH 4.5.			

Radicals immobilized in proteins are mainly represented by tryptophanyl and/or tyrosyl radicals, whose identification and assignment might lead to ambiguities. In both cases, the dominant splitting in the EPR spectrum arises from distinct hyperfine couplings of the β -methylene protons β_1 and β_2 at C_{β} , attached to C3 or C1 of the tryptophan or tyrosine ring, respectively.⁴⁷ The hyperfine coupling constants of these two β protons govern the EPR spectrum and depend on the sidechain orientation measured as the dihedral angle (ϑ) for each of the β protons according to the Heller-McConnell relation.⁴⁷ When the dihedral angles are equivalent, the side chain is either in the tryptophan ring plane or perpendicular to it. In the first case, the dihedral angle for both β protons is $\approx 30^{\circ}$ and in the latter case $\approx 60^{\circ}$.

The crystallographic dihedral angles for the β protons of Trp405 (PDB: 6fsk) are $\vartheta_1 \approx |23.5^\circ|$ for H- β_1 and $\vartheta_2 \approx |35.5^\circ|$ for H- β_2 . These values correspond to almost equivalent isotropic coupling constants (2.18 mT for H- β_1 and 1.72 mT for H- β_1 , calculated from the Heller–McConnell relation) and are in agreement with the $A_{iso} = 1.92$ mT value determined from the experimental spectrum where the tryptophanyl radical represents the main contribution. For AauDyP the crystallographic dihedral angles for the β protons are $\vartheta_1 \approx |30^\circ|$ for H- β_1 and $\vartheta_2 \approx |40^\circ|$ for H- β_2 revealing approximately the same orientation for both the catalytically active Trp405 and Trp377 in *Pos*DyP4 and *Aau*DyP, respectively.^{43,47}

The EPR spectrum of the W405S variant, which was shown to be inactive on all the substrates assayed except on Mn^{2+} , also shows a high-spin Fe³⁺. After the addition of H₂O₂, a low intensity signal of a protein radical species is observed, lacking the tryptophanyl radical component observed in the native *Pos*DyP4 spectrum (Figures 6B and S8). The nonstructured signal has a line width around 1 mT, and its g value (2.0045) suggests a tyrosyl radical, similar to those detected in the H₂O₂-activated form of several peroxidases.^{43,47,48}

The *Pos*DyP4 crystal structure shows Tyr339, and several phenylalanine residues, in close proximity to Trp405. Therefore, the mixed protein radical in the EPR spectrum of activated *Pos*DyP4 includes components of both Trp405 (its main contribution being also confirmed by the spectrum of the W405S variant lacking the tryptophanyl component) and Tyr339 radicals. Then, the tyrosyl signal observed in the EPR spectrum of the W405S variant most probably corresponds to Tyr339, whose radical signal is affected by the neighboring phenylalanine residues.

These EPR results substantiate the QM/MM calculations that: (i) identified Trp405 as the residue with the highest tendency to form a radical (highest spin density) at the heme neighborhood (Figure 5A); and (ii) predicted that this

tendency would pass to Tyr339 in the W405S variant, as shown when excluding Trp405 from the QM region (Figure 5B).

Functionality of the Mn^{2+} -Oxidation Site Confirmed by Mutagenesis. According to the computational predictions, four acidic amino acids (Asp215, Glu345, Asp352, and Asp354) could be involved in the binding of Mn^{2+} , with Tyr339 also located at the cation vicinity (Figure 4). To confirm the role of the above residues, we replaced them for alanines and seven mutated variants, carrying from 1 to 3 mutations, were characterized.

As shown in Table 2, all the variants exhibited a reduced Mn^{2+} oxidation activity, while their activity on ABTS (which initially was only slightly lower than the one found on Mn^{2+}) remained basically unchanged (Table S3). This confirms the specific role of these residues in Mn^{2+} binding and oxidation. Tyr339 appears as the most important residue because no Mn^{2+} oxidation was observed for the Y339A variant. Glu345 seems to be also a key residue because its E345A variant exhibited a 50-fold decrease in the k_{cat} and a 2-fold increase in the K_m , resulting in an overall 100-fold lower catalytic efficiency (Table 2 and Figure 7). In contrast, substitution of Asp215, Asp352, or Asp354 had a lesser effect in terms of



Figure 7. Kinetics of Mn^{2+} oxidation by *Pos*DyP4 and seven simple, double and triple variants (substrate concentration in a logarithmic scale). Biphasic kinetics are evident for several variants, such as D352A/D354A, compared with native DyP4 (see Table 2). Means and 95% confidence limits.

 $Mn^{2+} k_{cat}$ which was less than 2-fold reduced, most probably due to suboptimal positioning of the substrate ion.

Nevertheless, changing the latter three residues impaired the Mn^{2+} affinity, resulting in an increase of the K_m value from 9to 15-fold. Interestingly, the Asp352 and Asp354 substitutions, individually or together, resulted in a biphasic kinetics, as shown in Figure 7 (see Figure S9 for representation using a nonlogarithmic scale). These results enabled calculation of two sets of kinetic constants, as described previously,^{13,43} characterized by their higher and lower affinities for Mn^{2+} (which would correspond to different binding sites or geometries) (Table 2). The combination of mutations (D352A/D354A or E345A/D352A/D354A) strongly impaired the enzyme activity, but no Mn^{2+} oxidation was only found for the Y339A variant.

New Mn²⁺-Oxidation Site in DyPs. The well characterized Mn²⁺ oxidation site in fungal MnPs and VPs comprises three acid residues (Glu35, Glu39, and Asp179 in *Phaner*ochaete chrysosporium MnP, PDB: 1mnp; and Glu36, Glu40 and Asp175 in *Pleurotus eryngii* VP, PDB: 2boq) and one heme propionate.^{10,49} This site includes a narrow access channel toward the heme propionate (or manganese channel), different from the wider heme access channel that provides access to H_2O_2 and some reducing substrates to the heme edge.¹⁰ In the above channel, the Mn²⁺ ion is located at ~2 Å from the heme propionate, allowing direct electron transfer. However, in *Pos*DyP4, the Mn²⁺ binding site is located at the protein surface further from the heme (~11 Å), and an electron transfer pathway is required. This pathway involves Tyr339, as predicted by the QM/MM calculations and confirmed by the mutagenesis experimental results described above.

Up to this study, two Mn²⁺-binding sites have been reported in DyPs from *Rhodococcus jostii* RHA1,²⁶ and its N246A variant with improved Mn²⁺-oxidation activity,²³ and from *Amycolatopsis* sp. 75iv2.²¹ Although crystallographic evidence was provided in these cases, including Mn-anomalous map of the R. jostii variant, no experimental confirmation on their involvement in the Mn²⁺ oxidation reaction has been reported. Even though the location of the manganese-binding site in R. jostii DyPB is reminiscent (but at 4 Å distance from heme) of the one described for class II peroxidases, metal coordnation is limited to only water molecules (PDB: 4hov) hindering its fixation and electron transfer. In contrast, the Mn²⁺⁻binding site in the Amycolatopsis sp. DyP2, which has much better Mnoxidation activity than R. jostii DyP,²¹ is analogous to the one characterized here in PosDyP4, appearing as mirror images when the heme cofactors are superimposed (Figure 8). In both cases, the manganese is located at the protein surface (Figure S10), far from the heme group, and a direct oxidation is not possible. However, this limitation is solved by transferring the electron via an intermediate tyrosine residue, as demonstrated here for PosDyP4. Interestingly, the more distant residue in the PosDyP4Mn²⁺-binding site (Asp215) is equivalent to AspDyP2 Glu273, which was reported by Brown et al.²¹ to occupy a second position in the coordination sphere of the Mn²⁺ ion. The latter authors also reported that the four amino acids involved in Mn²⁺ oxidation (one tyrosine and three acidic residues) are conserved in a few other DyP-type prokaryotic sequences, in addition to AspDyP4.²¹

CONCLUSIONS

As illustrated in the scheme of Figure 9 (upper arrow), we demonstrate for the first time (using directed mutagenesis)



Figure 8. Superposition of the heme cofactors showing the mirror position of the Mn^{2+} oxidation sites in the DyPs of *P. ostreatus* (after ligand diffusion with PELE on PDB: 6fsk) and *Amycolatopsis* sp. (as found in PDB: 4g2c). Residues are shown as CPK-colored sticks with blue and white carbons, respectively, and Mn^{2+} ions as spheres. Residue numbering in AspDyP2 is based on Brown et al.²¹ Some distances are indicated in angstroms.



Figure 9. Proposed long-range electron transfer pathways (thick arrows) to the PosDyP4 heme for the oxidation of Mn^{2+} at the new site described (dashed circle) and oxidation of aromatics at the catalytic Trp405. Displacement of several side chains during Mn^{2+} binding are shown by superposition of the molecular structures before (CPK colored) and after (cyan colored) PELE simulations.

that oxidation of Mn^{2+} by a fungal DyP, from *P. ostreatus*, and probably also by some prokaryotic DyPs, such as that from Amycolatopsis sp. 75iv2, is produced at the protein surface (>11 Å from the heme) implying a long-range electron transfer pathway. Four acidic residues (and two water molecules) coordinate the metal ion at this position. One of these residues (Glu345) would receive the electron from Mn²⁺ transferring it to a neighbor tyrosine (Tyr339), which is absolutely necessary for the oxidation reaction to take place. Among fungal ligninolytic peroxidases, the involvement of a tyrosine residue in electron transfer from reducing substrate to heme has been reported only for the lignin peroxidase of *Trametopsis cervina*.⁴⁸ In contrast, tyrosine involvement in electron transfer for Mn²⁺oxidation, as suggested for the first time by Brown et al.²¹ in AspDyP2, seems to be produced in distant DyPs of both eukaryotic and prokaryotic origins.

QM/MM calculations predicted the involvement of Tyr339 in the electron transfer from Mn^{2+} to the heme, as confirmed later by directed mutagenesis of *Pos*DyP4. EPR spectra are consistent with the formation of a tyrosyl radical in the H₂O₂activated W405S variant oxidizing Mn^{2+} , which in the native enzyme overlapped with a tryptophanyl radical. The latter would correspond to the solvent-exposed Trp405 radical involved in oxidation of aromatic substrates. Its formation was also predicted by QM/MM and confirmed by the EPR spectrum of the W405S variant, where only the tyrosyl radical component was observed. The abstracted electron must travel from the exposed Trp405 to the heme, the final part of the route being shared with the Mn^{2+} -oxidation pathway described above (Figure 9, lower arrow).

The new Mn^{2+} oxidation site, whose operation is demonstrated, could contribute to microbial biodegradation of lignin, acting on its phenolic moiety via Mn^{3+} chelates, as well as in the subsequent transformation of lignin-derived products. The Mn^{3+} chelates could also act on the nonphenolic lignin by promoting the formation of reactive lipid radicals, as proposed for selective degradation of lignin by some white-rot fungi.⁵⁰ DyPs could have a relevant contribution to the above reactions, especially in the case of the bacterial DyPs oxidizing Mn^{2+} , such as those from *Amycolatopsis* species, because no typical ligninolytic peroxidases have been described in prokaryotes, despite the likely bacterial contribution to degradation of lignin products and their incorporation to soil humus.⁵¹

ASSOCIATED CONTENT

S Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acscatal.8b02306.

Crystallographic data (Table S1); PCR primers (Table S2); ABTS oxidation kinetic parameters (Table S3); SDS-PAGE of PosDyP4 (Figure S1); UV-visible spectra of PosDyP4 and seven variants (Figure S2); detail of the PosDyP4 crystal electron-density map (Figure S3); overall crystal structures of PosDyP4 and 11 related proteins (Figure S4); heme access channel in PosDyP4 (Figure S5); comparison of empty Mn²⁺ binding site in the PosDyP4 crystal structure and after ion binding simulation (Figure S6); EPR spectra before and after activation by H₂O₂ (Figure S7); Intensity comparison of PosDyP4 and W405S EPR signals (Figure S8); kinetics of Mn²⁺ oxidation in nonlogarithmic scale (Figure S9); and solvent access surface of P. ostreatus and Amycolatopsis sp. DyPs showing Mn2+ binding (Figure S10) (PDF)

AUTHOR INFORMATION

Corresponding Author

*E-mail: ATMartinez@cib.csic.es.

ORCID 💿

Angel T. Martínez: 0000-0002-1584-2863

Present Address

[‡]E.F.-F.: Technical University of Delft, 2600 AA, Delft, The Netherlands

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

This work has been funded by the H2020 BBI-JU (www.bbieurope.eu) project EnzOx2 (BBI-PPP-2015-2-720297; www. enzox2.eu) and the FP7 project INDOX (KBBE-2013-7-613549; www.indoxproject.eu), together with the BIO2014-56388-R (NOESIS), BFU2016-77835-R, CTQ2016-79138-R, and (BIO2017-86559-R (GENOBIOREF) projects of the Spanish Ministry of Economy, Industry and Competitiveness, cofinanced by FEDER funds. The sequencing of *P. ostreatus* genome at JGI was supported by the DOE Office of Science, under Contract No. DE-AC02-05CH11231. We thank the staff of the ID30B beamline at ESRF (Grenoble, France) for assistance. CSGI (Florence, Italy) FFABR 2017 (Legge 232/ 2016 Comma 295, and Legge di Bilancio 2017) and MIUR (Dipartimento di Eccellenza, 2018-2022) grants are acknowledged for financial support.

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SUPPORTING INFORMATION

Description of a Non-Canonical Mn(II)-Oxidation Site in Peroxidases

Elena Fernández-Fueyo,[§] Irene Davó-Siguero,[§] David Almendral,[§] Dolores Linde,[§] Maria Camilla Baratto,[#] Rebecca Pogni,[#] Antonio Romero,[§] Victor Guallar,^{¶†} and Angel T. Martínez^{*§}

[§]Centro de Investigaciones Biológicas, CSIC, Ramiro de Maeztu 9, E-28006 Madrid, Spain; [#]Department of Biotechnology, Chemistry and Pharmacy, University of Siena, I-53100 Siena, Italy; [¶]Barcelona Supercomputing Center, Jordi Girona 29, E-08034 Barcelona, Spain; [†]ICREA, Passeig Lluís Companys 23, E-08010 Barcelona, Spain

*E-mail: ATMartinez@cib.csic.es

This Supporting Information includes crystallographic data (**Table S1**); PCR primers (**Table S2**); ABTS oxidation kinetic parameters (**Table S3**); SDS-PAGE of *Pos*DyP4 (**Figure S1**); UV-visible spectra of *Pos*DyP4 and seven variants (**Figure S2**); detail of the *Pos*DyP4 crystal electron-density map (**Figure S3**); overall crystal structures of *Pos*DyP4 and eleven related proteins (**Figure S4**); heme access channel in *Pos*DyP4 (**Figure S5**); comparison of empty Mn^{2+} binding site in the *Pos*DyP4 crystal structure and after ion binding simulation (**Figure S6**); EPR spectra before and after activation by H_2O_2 (**Figure S7**); Intensity comparison of *Pos*DyP4 and W405S EPR signals (**Figure S8**); kinetics of Mn^{2+} oxidation in non-logarithmic scale (**Figure S9**); and solvent access surface of *P. ostreatus* and *Amycolatopsis* sp DyPs showing Mn^{2+} binding (**Figure S10**) (PDF file).

Table	S1.	X-ray	data	collection	and	refinement
statisti	ics fo	r PosD	yP4 (c	lata in parei	nthesi	s correspond
to the l	ast re	esolutio	n laye	r).		

Data collection:	
Wavelength (Å)	0.9762
Crystal system, space group	Orthorhombic, P2 ₁ 2 ₁ 2 ₁
Unit cell parameters (Å)	a=62.72, b=117.57, c=140.43
Resolution range (Å)*	70.22 – 1.56 (1.64 – 1.56)
N° of observations	655645 (96513)
N° of unique reflections	144926 (20996)
Redundancy	4.5 (4.6)
Completeness (%)	97.3 (97.4)
Mean Io(I)	12.7 (2.3)
R _{merge} ^a	0.065 (0.514)
R _{meas} ^b	0.083 (0.649)
CC _{1/2} ^c (%)	99.7 (74.5)
Wilson B-factor (Å ²)	17.70
Refinement:	
Resolution range (Å)	70.22 – 1.56 (1.60 – 1.56)
N° of working reflections	137552 (10100)
N° of testing reflections	7250 (553)
Completeness (%)	97.05
R _{work}	0.162 (0.307)
R _{free} #	0.199 (0.334)
Nº of non-H atoms:	
Protein	7484
Heme group	86
MES	12
Water molecules	1119
Average B factors (Ų):	
Protein	18.56
Heme group	10.72
MES	22.95
Water molecules	29.70
RMS deviations:	
Bond lengths (Å)	0.021
Bond angles (°)	2.04
Ramachandran plot statistics:	
Favoured (%)	97.8
Allowed (%)	2.0
Outliers (%)	0.2
PDB:	
ID code	6FSK

Table S2. Primers (forward and reverse) used for the generation of *Pos***DyP4 variants**The mutated codons are underlined and the mutated nucleotides are in italics.

Mutation	Туре	Sequence
D196A	fwd rev	C GCC CAT GAA CAC TTC GGC TTC CTC <u>GCC</u> GGT ATC TCC GGA GAT ACC <u>GGC</u> GAG GAA GCC GAA GTG TTC ATG GGC G
D215V	fwd rev	C CCG CTC CCA GGT CAA <u>G7T</u> CCG ATC AGA CCG GGC GCC CGG TCT GAT CGG <u>AAC</u> TTG ACC TGG GAG CGG G
Y339A	fwd rev	GCT GCT CAC ATC AGA AAG ACC \underline{GCC} CCG CGC AAC GAT TTG GAG GGC CC GG GCC CTC CAA ATC GTT GCG CGG \underline{GGC} GGT CTT TCT GAT GTG AGC AGC
E345A	fwd rev	C CCG CGC AAC GAT TTG <u>GCG</u> GGC CCG CC GG CGG GCC <u>CGC</u> CAA ATC GTT GCG CGG G
D352A	fwd rev	GGC CCG CCT TTG AAA GCT <u>GCT</u> ATT GAC AAC CG CG GTT GTC AAT <u>AGC</u> AGC TTT CAA AGG CGG GCC
D354A	fwd rev	GG CCC GCC TTTG AAA GCT GAT ATT <u>GCC</u> AAC CGT AGA ATC ATC CGC GCG GAT GAT TCT ACG GTT <u>GGC</u> AAT ATC AGC TTT CAA AGG CGG GCC
W405S	fwd rev	C CAC TTC ATT CAG GAA TCT <u>TGG</u> GCC AAT GCC C G GGC ATT GGC <u>CCA</u> AGA TTC CTG AAT GAA GTG G
D352A/D354A	fwd rev	GGC CCG CCT TTG AAA GCT <u>GCT</u> ATT GCC AAC CGT AGA ATC ATC CGC GCG GAT GAT TCT ACG GTT GGC AAT <u>AGC</u> AGC TTT CAA AGG CGG GCC

Table S3. Kinetic Constants - K_m (μ M), k_{cat} (s⁻¹) and K_m/k_{cat} (s⁻¹.mM⁻¹) - for Oxidation of ABTS by *Pos*DyP4 and Seven Variants Mutated near the Mn²⁺-Oxidation Site^a

	DyP4	Y339A	D215A	E345A	D352A	D354A	D352A/ D354A	E345A/ D352A/ D354A
K _m	790±160	590± 50	320±30	420±60	410±70	420±70	500±90	380±40
k _{cat}	280±30	240±10	70±0	170±10	160±10	170±10	230±10	200±10
k _{cat} /K _m	350±50	240±30	230±20	420±50	390±10	410±10	450±30	540±50
^a Means and 95% confidence limits from reactions at 25°C in 0.1 M tartrate, pH 3.5.								



Figure S1. SDS-PAGE of purified *Pos*DyP4 (lane **a**, arrow) and molecular-mass markers (lane **b**).



Figure S2. UV-visible spectra of *Pos*DyP4 and seven variants at the Mn^{2+} -oxidation site. The Soret band wavelength and details of the 400-700 nm region are included.



Figure S3. $2F_o$ - F_c electron density map, contoured at 1σ level, of residues surrounding several water molecules in *Pos*DyP4, and CPK-colored stick representation of these residues in PDB: 6fsk.



Figure S4. Overall structure of *Pos*DyP4 (6fsk) and selected structurally related proteins of fungal (3afv and 4au9) and prokaryotic origin (4g2c, 5c2i, 5fw4, 3o72, 5jxu, 5vj0, 3veg, 4gs1 and 5de0). See **Table 1** for species names of the different PDB entries (red asterisks indicate prominent helix-8 in three of the structures).



Figure S5. Heme access channel in *Pos*DyP4. **A.** Detail of solvent access surface with heme (green) and channel residues (CPK colors) as spheres. **B.** Semi-transparent surface showing heme and channel residues as sticks (from PDB: 6fsk).



Figure S6. Superimposition of the four acidic residues involved in Mn^{2+} binding and Tyr339 (**Figure 4**) before and after ligand diffusion with PELE (as CPK-colored sticks with white and cyan carbons, respectively). The displacement of some side chains for metal ion binding is observed.



Figure S7. X-band EPR spectra of 0.25 mM native *Pos*DyP4 in tartrate buffer, pH 3, before (black line) and after (red line) the addition of 8 eq. of H_2O_2 . Experimental conditions were 5 mW microwave power, 0.4 mT modulation amplitude, and v= 9.39 GHz.



Figure S8. Comparison of 20 mT scan width X-band EPR spectra of 0.25 mM native *Pos*DyP4 (*black line*) and W405S variant (*red line*) in tartrate buffer, pH 3, after the addition of 8 eq. of H₂O₂. Conditions were 5 mW microwave power, 0.2 mT modulation amplitude, and v= 9.39 (*Pos*DyP4) or 9.38 (W405S) GHz. The normalized spectra are shown in **Figure 6**.



Figure S9. Kinetics of Mn^{2+} oxidation by *Pos*DyP4 and seven simple, double and triple variants, with substrate concentration in arithmetic scale. See **Figure 7** for representation of the same data with substrate concentration in logarithmic scale for better showing the biphasic kinetics with some of the variants. Means and 95% confidence limits.



Figure S10. Solvent access surface of *Amycolatopsis* sp DyP2 (**A**) and *P. ostreatus* DyP4 (**B**) showing the exposed Mn^{2+} (purple sphere) oxidation site and the acidic residues involved in metal ion binding. See **Figure 8** for PDB entries.