X-ray Structure of a Hydroxylase–Regulatory Protein Complex from a Hydrocarbon-Oxidizing Multicomponent Monooxygenase, *Pseudomonas* sp. OX1 Phenol Hydroxylase,[†,‡]

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Abstract

Phenol hydroxylase (PH) belongs to a family of bacterial multicomponent monooxygenases (BMMs) with carboxylate-bridged diiron active sites. Included are toluene/*o*-xylene (ToMO) and soluble methane (sMMO) monooxygenase. PH hydroxylates aromatic compounds, but unlike sMMO, it cannot oxidize alkanes despite having a similar dinuclear iron active site. Important for activity is formation of a complex between the hydroxylase and a regulatory protein component. To address how structural features of BMM hydroxylases and their component complexes may facilitate the catalytic mechanism and choice of substrate, we determined X-ray structures of native and SeMet forms of the PH hydroxylase (PHH) in complex with its regulatory protein (PHM) to 2.3 Å resolution. PHM binds in a canyon on one side of the $(\alpha\beta\gamma)_2$ PHH dimer, contacting α-subunit helices A, E, and F ~12 Å above the diiron core. The structure of the dinuclear iron center in PHH resembles that of mixed-valent MMOH, suggesting an Fe(II)Fe(III) oxidation state. Helix E, which comprises part of the iron-coordinating four-helix bundle, has more π-helical character than analogous E helices in MMOH and ToMOH lacking a bound regulatory protein. Consequently, conserved active site Thr and Asn residues translocate to the protein surface, and an ~6 Å pore opens through the four-helix bundle. Of likely functional significance is a specific hydrogen bond formed between this Asn residue and a conserved Ser side chain on PHM. The PHM protein covers a putative docking site on PHH for the PH reductase, which transfers electrons to the PHH diiron center prior to O_2 activation,

suggesting that the regulatory component may function to block undesired reduction of oxygenated intermediates during the catalytic cycle. A series of hydrophobic cavities through the PHH α-subunit, analogous to those in MMOH, may facilitate movement of the substrate to and/or product from the active site pocket. Comparisons between the ToMOH and PHH structures provide insights into their substrate regiospecificities.

Bacterial multicomponent monooxygenases (BMMs)[1] are a diverse family of enzymes that hydroxylate and epoxidize an array of hydrocarbon substrates, including alkanes, alkenes, and aromatics in a regio- and enantioselective fashion (1-3). This chemistry takes place at a carboxylate-bridged diiron center similar to the ones found in the R2 subunit of type I ribonucleotide reductase (RNR-R2), stearoylacyl carrier protein (ACP) Δ^9-desaturase, and ferritin (4,5). Bacteria harboring different BMM family members, which include soluble methane monooxygenases (sMMOs), toluene monooxygenases (TMOs), phenol hydroxylase (PH), alkene monooxygenase (AMO), tetrahydrofuran monooxygenase (THFMO), and hyperthermophilic toluene monooxygenase (SsoMO), use small organic compounds as their sole source of carbon and energy. The substrate transformations performed by this family of enzymes are responsible, in part, for removing atmospheric methane, a greenhouse gas, and carcinogenic halogenated solvents like TCE from the environment (6,7). Because of the broad range of substrates oxidized by these enzymes, they have generated much interest for bioremediation and synthetic industrial applications.

BMMs like sMMO and PH consist of a 200–255 kDa dimeric hydroxylase of the form $(\alpha\beta\gamma)_2$, a cofactorless 10–16 kDa regulatory protein that enhances catalytic turnover by 30–150-fold and a FAD- and [2Fe-2S]-containing 38–40 kDa reductase that supplies the hydroxylase with electrons by consuming NADH (1-3). TMOs and SsoMOs differ slightly since they utilize an additional, Rieske protein to assist with electron transfer between the reductase and hydroxylase components (2,3). The active sites in the resting states of the methane (MMOH) and toluene/o-xylene (ToMOH) monooxygenase hydroxylases contain a diiron(III) center coordinated by four glutamate and two histidine ligands from a four-helix bundle composed of helices B, C, E, and F of the protein α-subunit (8,9). Solvent-derived water and hydroxide ions complete the octahedral coordination spheres. Sequence, spectroscopic, and structural analyses suggest that all BMM hydroxylases have nearly identical dinuclear iron centers (1,10-12). Because AMO, TMO, SsoMO, and PH are incapable of hydroxylating alkanes, whereas sMMO and THFMO are adept at this transformation, other features of the system must control substrate reactivity at their dioxygen-activated metal centers. For sMMO, methane hydroxylation is achieved exclusively by a di-μ-oxo diiron(IV) intermediate, Q, whereas other reactions can occur at a diiron(III) peroxo precursor of Q, Hperoxo (1,13, 14). A diiron(IV) intermediate has not been detected in any other BMM hydroxylase, but recent work has identified oxygenated diiron(III) species capable of oxidizing aromatic substrates (15).[2],[3]

Comparisons between the MMOH and ToMOH structures have provided insight into how the protein scaffold may tune the reactivity and substrate specificity of the diiron center by controlling the access of small molecules to the active site pocket. In particular, there is a large access channel in ToMOH linking the diiron center to the exterior environment that is absent in MMOH (8,9). The regulatory proteins in these systems may also influence reactivity by inducing different structural changes at or near the diiron active site. In addition to enhancing the catalytic rate, the regulatory protein alters the substrate regiospecificity (16-18), influences the spectroscopic and redox properties of the diiron center (1), governs movement of the substrate and product to and from the active site pocket (5), and couples NADH consumption by the reductase with substrate hydroxylation at the diiron center (18,19). Knowledge of the docking site on the hydroxylase and of structural changes at the diiron center imposed by the regulatory protein is therefore essential for improving our understanding of O_2 and C–H bond activation in BMMs.

To address these questions and improve our structural knowledge about the BMM family members, we have determined in this study the X-ray crystal structure of a complex between the hydroxylase component of phenol hydroxylase, PHH, and its regulatory protein, PHM, from *Pseudomonas* sp. OX1 to 2.3 Å resolution. The information afforded provides novel insights into the functions of the regulatory protein and the tuning of the diiron active site chemistry in BMM enzyme systems.

MATERIALS AND METHODS

Purification of Phenol Hydroxylase

The hydroxylase (PHH), regulatory protein (PHM), and reductase (PHP) from *Pseudomonas* sp. OX1 were expressed simultaneously from the pJSX148 plasmid in *Escherichia coli* JM109 cells essentially as described by Cafaro et al. (20). The procedure was modified such that 100 μM $Fe(NH_4)_2SO_4 \cdot 6H_2O$ was added to each liter of medium at induction and every hour thereafter until the cells were harvested.

The frozen cells were resuspended in a 25 mM MOPS (pH 7.0), 10% glycerol, 50 mM NaCl buffer (buffer A) supplemented with 5 mM $MgCl_2$, Pefabloc, 20 units of DNase, and PMSF and then lysed by sonication using 6×1 min pulses. Particulate debris was removed by centrifugation at 163000g for 45 min, after which the supernatant was filtered through a 0.2 μm membrane and loaded onto a DEAE column conditioned with buffer A. A salt gradient was run from 50 to 400 mM NaCl, and the hydroxylase-containing fractions were pooled and concentrated to <5 mL using an Amicon concentrator fitted with a YM100 membrane. The concentrated protein was loaded onto an S300 column preequilibrated with 25 mM MOPS (pH 8.0), 10% glycerol, and 300 mM NaCl (buffer B) and run in that same buffer. Fractions containing PHH were pooled, loaded onto a phenyl Sepharose column conditioned with 10 mM MOPS (pH 7.0), 10% glycerol, and 100 mM NaCl, and eluted by using a 100 to 0 mM NaCl gradient. The purified PHH, which contained noticeable amounts of PHM, was concentrated, frozen in liquid N_2, and stored at −80 °C.

A selenomethionine (SeMet) derivative of PHH was generated by growing BL21(DE3) Gold *E. coli* cells transformed with the pJSX148 plasmid on LeMaster's medium supplemented with 90 mg/L Se-Met and ~5 mg/L folate, vitamin B_{12}, nicotinamide, pantothenate, thiamine, riboflavin, and *p*-aminobenzoic acid (21). The protein was expressed and purified as described above.

Activity and Iron Content

The enzyme activity, measured by using a catechol 2,3-dioxygenase (C2,3O) assay as described previously (20), was 470 ± 40 milliunits/mg. The C2,3O protein required for the assay was expressed and purified as described previously (20). The iron content of PHH was determined to be 3.6 Fe atoms/dimer by a colorimetric ferrozine assay (22).

Crystallization and Collection of X-ray Diffraction Data

Crystals were grown at 20 °C by the hanging drop vapor diffusion method. A protein solution containing 35 μM PHH in 10 mM MES (pH 7.1) and 10% glycerol was combined with an equal volume of crystallization buffer containing 100 mM Tris (pH 7.0), 150 mM Na_2MoO_4, 5% glycerol, and 17–20% PEG 8000 (w/w). This condition was initially identified by using the Hauptman-Woodward Institute high-throughput screen. The addition of varying amounts of PHM to the crystallization buffer did not improve the occupancy of the regulatory protein in the crystals (vide infra). Before data collection, PHH crystals were frozen in a cryo solution containing the precipitant and 20% glycerol. Data at the Se peak wavelength were collected on beamline 9-1 at SSRL, and the native data set was collected at APS on the NECAT 8BM beamline. The data were indexed and scaled with HKL2000 (23). The space group is $P2_12_12_1$, and the unit cell dimensions are 87.8 Å × 146.3 Å × 190.0 Å. Further experimental details about data collection and refinement are given in Table 1.

Structure Determination

Molecular replacement was attempted initially by using the most structurally conserved regions of the MMOH and ToMOH α- and β-subunits. None of the starting α, $α_2$, β, $β_2$, αβ, and $α_2β_2$ models used for molecular replacement afforded good starting phases, even though the PHH α- and β-subunits fold in a manner similar to the folding of these subunits in MMOH and ToMOH (see below).

Single-wavelength anomalous dispersion data for the selenomethionine derivative were then used to determine a set of starting phases. Attempts to determine the heavy atom substructure by using CNS (24), Solve (25), and SHELX (26) failed. SnB (27) was the only program able to locate 71 of 77 Se atoms in the asymmetric unit by using the peak wavelength. Native data were included in the phasing by refinement of both real and anomalous occupancies with MLPHARE from the CCP4 suite of programs (28). Initial phases were calculated with MLPHARE, improved, and extended to the limit of the native data by density modification with DM (28). Resolve (25) was used to autobuild ~50% of PHH, after which the remainder of the molecule was manually built in XtalView (29) and O (30) and refined with REFMAC5 (28) and CNS (24). Because of structural and occupancy differences, models for both the native and SeMet data sets were generated. The relative occupancy of PHM was estimated by refining the PHM occupancies after its *B*-factors were fixed to reflect the average value of the entire PHH molecule. A Ramachandran plot calculation using PROCHECK (31) indicated that 99.8% of the residues occupied allowed regions.

RESULTS AND DISCUSSION

PHH Global Fold

The PH hydroxylase component has an $(αβγ)_2$ homodimeric composition with a noncrystallographic 2-fold symmetry axis similar to those in the ToMOH and MMOH structures (Figure 1A and Figure S1A–C of the Supporting Information) (8, 9). The folds of the PHH α- and β-subunits are nearly identical to those of the respective subunits in ToMOH and MMOH. The rms deviations between the α-carbon atoms are 1.5–1.6 Å for all possible α-to-α and β-to-β combinations. The largest topological differences between these evolutionarily related hydroxylase proteins occur for the γ-subunit, both in its fold and in its location. For PHH and

ToMOH, the γ-subunits are positioned in the top left and right quadrants (11 and 1 o'clock, respectively) of the protein with similar ferredoxin-like folds despite having sequences that are only 13% identical (Figure S2) (9). Unlike the ToMOH γ-subunit, however, the PHH γ-subunit has an unstructured 22-amino acid N-terminus that forms extensive contacts with the α-subunit. For MMOH, the mostly α-helical γ-subunits localize to the far left- and right-hand sides of the protein (9 and 3 o'clock, respectively) and contact both the α- and β-subunits (Figure S1). Such structural differences are not surprising given the poor sequence conservation for this subunit among the different BMM family members. The function of this subunit, which is absent in some BMMs like alkene monooxygenases, is unclear (2, 3, 11).

PHM Fold and Binding Site on the Hydroxylase

The interface between the αβγ protomers of PHH forms a canyon, which we previously suggested would be the binding site for the BMM regulatory protein and reductase components on their hydroxylases (5,8,32,33). On one face of the PHH protein, a single molecule of PHM binds in the canyon region across α-subunit helices A, E, F, and H, just 12 Å above the diiron center at its closest distance. On the opposite face of the hydroxylase, both a γ-subunit from a different, symmetry-related hydroxylase molecule in the unit cell and the N-terminus of the β-subunit occupy the space defined by the PHM binding site (Figure S3). The occupancy of the PHM component was estimated to be 40 and 50% in the native and selenomethionine (SeMet) structures, respectively. These different occupancies presumably reflect differing amounts of PHM that copurified with the hydroxylase component. We therefore generated models of the PHHPHM complex from both the native and SeMet data.

The PHM global fold comprises three α-helices flanked by two nearly perpendicular β-sheets (Figure 1B). The fold is nearly identical to those previously reported for the NMR and crystal structures of the methane monooxygenase (MMOB; PDB entries 1CKV and 2MOB) and toluene 4-monooxygenase (T4MOD; PDB entries 2BF5 and 1G10) regulatory proteins (34-37). The rms deviations between PHM and the T4MOD and MMOB proteins are between 1.5 and 1.6 Å for all backbone atoms (Figure S4). Comparisons between hydroxylase-bound PHM, T4MOD, and MMOB suggest there may be no significant structural rearrangements of the peptide backbone when these regulatory proteins form a complex with their respective hydroxylases.

Primarily in contact with the α-subunit of one protomer, PHM interacts with PHH helices A, E, F, and H via β-strands 1, 5, and 6, α-helix 3, the N-terminal region of α-helix 1, the loop between β-strands 2 and 3, and the peptide chain termini (Figures 1B and S5). The PHM binding site on the surface of the PHH α-subunit is formed by portions of the helices including residues 64–68 (helix A), 200–207 (helix E), and 226–245 (helix F), which collectively create a concave surface that complements the protrusive binding face of the regulatory protein. The PHH-PHM binding interaction is primarily hydrophobic, with some flanking polar and charged residues (Figure S1D). The N- and C-termini of the PHH regulatory protein are oriented away from the canyon region toward solvent and form some contacts with α-subunit helix H (Figure 1B).

Sequence alignments of the regulatory proteins from representative BMM family members indicate that very few of the amino acids are strictly conserved (Figure S5). Residues at the binding interface appear to be conserved only within a subfamily, suggesting why regulatory protein cross-reactivity with hydroxylases from different BMM subfamilies is rarely observed. Nevertheless, we generated electrostatic surface maps of what might be the homologous hydroxylase binding faces of T4MOD and MMOB from the information in Figure S5. As depicted in panels E and F of Figure S1, these protein surfaces, like that in PHM, have significant hydrophobic character, but with a somewhat greater number of charged residues. NMR line broadening, mutagenesis, cross-linking, and spin-labeling studies probing interactions of MMOB with MMOH are highly consistent with a similar surface being used

by MMOB to bind its hydoxylase (Figure S5) (34,38-40). Previously published spin-labeling experiments by our laboratory identified a different and probably incorrect MMOB binding face (41). A recent discussion of component interactions in the sMMO system is available (40).

The PHH–PHM crystal structure presented here thus serves as an excellent template for modeling interactions of the folded regulatory protein core with its cognate hydroxylase in related BMM systems. Although the unstructured Nand C-termini of some regulatory proteins (Figure S5), such as those on MMOB that affect steps in the catalytic cycle (42-45), cannot be placed on the hydroxylase surface with any degree of certainty, the specific interactions identified here (vide infra) should be quite valuable in guiding future mutagenesis, structural, and mechanistic studies of all the BMM enzymes.

Diiron Center

The diiron center of PHH is located in the α-subunit four-helix bundle 12 Å below the surface. Both protomers in the native and SeMet structures have nearly similar diiron active sites despite the binding of PHM to only one side of the hydroxylase. The iron atoms are coordinated by a bridging or semibridging glutamate, Glu-138, the δ-nitrogen atoms of His-141 and His-236, which lie distal to the active site pocket, and three additional glutamate residues, Glu-108, Glu-199, and Glu-233 (Figures 2 and S6). Glu-108 coordinates to Fe1 and hydrogen bonds to a terminal water molecule bound to the same iron atom. This structural motif is observed in all known BMM hydroxylase structures in their various oxidation and product-bound states (9, 33, 46-50). In the native structure and the SeMet α-subunit with bound PHM, Fe2 is coordinated by Glu-233 in a bidentate chelating mode, the geometry of which is most similar to that of the mixed-valent, Fe(II)Fe(III) form of MMOH (MMOH$_{mv}$) (Figure 2C). In the other SeMet protomer without bound PHM, Glu-233 appears to coordinate to Fe2 by using only one oxygen atom, the other forming a hydrogen bond to the terminal water on Fe1. For most of the active sites, Glu-199 coordinates to Fe2 via just one oxygen atom while the other dangling oxygen atom is in hydrogen bond contact with nearby residues Tyr-115 and Gln-134 (Figures 2 and 3A). This ligand appears to adopt a bidentate chelating geometry in the SeMet α-subunit without bound PHM. The final iron ligand that we could identify is a water molecule bound exclusively to Fe1. Electron density corresponding to a bridging or semibridging water-derived ligand, like hydroxide or oxide, between the iron atoms was not observed. In the higher-resolution native structure, the Fe1–Fe2 distances are 3.6 and 3.3 Å for the PHM-bound and unbound protomers, respectively. The Fe1–Fe2 distances in the lower-resolution SeMet structure are 3.0–3.1 Å for both active sites. These shorter Fe–Fe distances may be a consequence of a fully bridging Glu-138 residue versus the semibridging one in the native structure. Overall, Fe1 has a five-coordinate, square-pyramidal geometry, whereas Fe2 adopts a more distorted structure.

The identity of the Fe2-coordinating histidine N-donor is notably different among the PHH, MMOH, and ToMOH dinuclear iron centers. In ToMOH, this histidine coordinates to iron by using its ϵ-nitrogen atom, whereas in MMOH and PHH, the histidine δ-nitrogen atom is employed. When the ToMOH structure was first determined, it was unclear whether this structural difference might influence the hydroxylation chemistry (9). Since PH and ToMO hydroxylate similar aromatic substrates at nearly equivalent rates (20), the structural difference is presumably of little or no consequence to this property of the dimetallic center.

At present, it is difficult to discern the true oxidation state of the different dinuclear iron centers in PHH enzymes. As purified, the enzyme has an optical band at 350 nm that is indicative of a (μ-oxo)diiron(III) species comprising ~68–96% of the sample (ϵ_{350} = 4800–6000 M^{-1} cm^{-1} per dinuclear iron cluster) (12). PHH from *Pseudomonas* sp. strain CF600 is 81% identical to the *P. stutzeri* OX1 enzyme and forms a complex with its regulatory component

involving the α-subunit as deduced by chemical cross-linking studies (51). Spectroscopic investigations of this enzyme indicate that the oxidized form contains both (μ-oxo)- and di-μ-(hydroxo)diiron(III) metal centers (12) and that the predominant, catalytically competent species is the oxo-bridged diiron(III) form. Preliminary EPR spectroscopic analysis of samples of *P. stutzeri* OX1 PHH in its resting and cryo-reduced states reveals that the purified protein contains a heterogeneous mixture of diiron(III) and mixed-valent Fe(II)Fe(III) species, the relative concentrations of which are at present unknown.[4] If the PHH metal centers in these crystal structures were fully oxidized, one would expect to see either a short Fe–Fe distance of 3.0 Å and a dihydroxo-bridged dimetallic cluster (9,48,49) or a longer Fe–Fe distance of 3.2–3.6 Å and an oxo-bridged cluster (4,5,52). None of the PHH diiron centers, however, has an observable bridging oxide or hydroxide ligand. Given that the geometries of the active site metal clusters are most similar to that of the mixed-valent Fe(II)Fe(III) form of MMOH (49), one possibility is that photoreduction by synchrotron radiation occurred to produce the mixed-valent oxidation state. A similar phenomenon has been reported previously for structures of Δ^9-desaturase and RNR-R2, although the oxidized forms of these proteins were reduced by two electrons (53-55). The structures of the PHH diiron centers presented here are not charge-balanced, however, and would require an additional negatively charged ligand, such as a bridging hydroxide ion, to achieve a neutral state similar to that observed in MMOH$_{mv}$ (Figure 2C). Although $|2F_o| - |F_c|$ and $|F_o| - |F_c|$ simulated-annealing omit electron density maps have not allowed us to identify such a ligand, the data quality at 2.3 and 2.7 Å resolution may not be sufficient to reveal a coordinating oxygen atom between the iron atoms (56).

A second possibility is that the metal centers have each been photoreduced by two electrons to the diiron(II) state, rendering the PHH active sites charge neutral. The structures do not, however, resemble the reduced forms of MMOH and ToMOH, which both have a 3.3–3.4 Å metal–metal distance and a glutamate residue that coordinates to Fe2 in a bidentate chelating fashion and simultaneously bridges the two iron atoms with one of its O atoms (48,49,57). Because the active sites on the PHM-bound and unbound sides of PHH are so similar, it is unclear whether PHM affects them both in a similar manner by an allosteric effect or not at all. The incomplete PHM occupancy may also be a factor. Further spectroscopic and structural investigations are required to improve our understanding of the PHH dinuclear iron center and to discern how the diiron site geometries are influenced by the oxidation state and the presence of bound PHM.

Second-Sphere Hydrogen Bonding Patterns

The second coordination sphere is critical to the formation and stability of the BMM diiron centers, providing ligand conformational constraints in the PHH, ToMOH, and MMOH structures (5). For PHH, the geometry of Glu-108, which coordinates to Fe1, is stabilized by a hydrogen bonding network involving Gln-145 and Ser-105 (Figure 3). A similar pattern exists in ToMOH involving Gln-141 and His-96 (9), but not in MMOH, which has a cysteine residue positioned 4.5 Å from the coordinating glutamate (8). Gln-145 is conserved among all PH and TMO family members, whereas Ser-105 and His-96 are not. Substituting the corresponding Gln with a Cys in toluene 4-monooxygenase (T4MO) resulted in a variant with a 10-fold diminished turnover rate that was still incapable of hydroxylating methane (58). A Gln-to-Val substitution in the same enzyme resulted in a 100-fold decrease in the rate of turnover compared to that of the wild-type protein. These results and the crystal structure presented here suggest that stabilization of the Fe1-coordinating terminal glutamate by hydrogen bonding may be critical for reactivity in the TMO and PH systems. The importance of such hydrogen bonding involving the analogous residue in MMOH, Glu-114, is less obvious because this residue has

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[4]V. Izzo, R. Davydov, M. H. Sazinsky, S. J. Lippard, and B. M. Hoffman, unpublished results.

no hydrogen bonding partners other than an active site pocket water molecule that is not always detected in every crystal structure (49).

At the other end of the diiron center, there is extensive hydrogen bonding to conserved Gln-134, which may be essential for stabilizing the geometry by which the bridging Glu-138 and terminal Glu-199 residues coordinate to Fe2 (Figure 3). Glu-199 may also form a weak hydrogen bond with Tyr-115; this residue is a phenylalanine in homologous PH family members, however. The same general arrangement of amino acids is preserved in MMOH (9). ToMOH utilizes a water molecule (or hydronium ion) to mediate interactions between the Fe-coordinating glutamates and a second-sphere glutamate, Glu-111. In the PHH structure, hydrogen bonds between two ordered water molecules link Gln-134 with conserved Arg-235, an arrangement that differs from that in the other hydroxylase proteins (see below). It is clear through these comparisons that BMMs employ a variety of schemes to stabilize the coordination spheres around the two iron atoms, but it is unknown whether and how these different architectures might contribute to the dissimilar reactivity of the enzymes. Comparisons of the native and SeMet α-subunits with and without bound PHM do not reveal any significant alterations in these hydrogen bonding patterns.

A Conserved Mononuclear Zn^{2+} Site

In the C-terminal domain of the PHH α-subunit, 24 Å from the diiron center, there exists a four-cysteine site that coordinates a single metal ion with a tetrahedral geometry and an average metal–sulfur distance of 2.3 Å (Figures 1A and 4). Sequence homology indicates that the coordinating cysteines are conserved only among members of the PH and AMO families (Figure S7). UV–vis, EPR, and Mössbauer spectroscopic studies of recombinant PHH from *P. stutzeri* OX1 and native PHH from *Pseudomonas* sp. strain CF600 do not reveal the presence of a thiolate rich mononuclear iron center, or any other mononuclear paramagnetic metal center, in any of the protein preparations (12).[3],[4] Anomalous difference electron density maps generated from data collected at the selenium edge reveal much stronger peaks at the mononuclear metal site than at the diiron center (Table 2), suggesting that the metal at this position is a stronger absorber of X-rays at 12 660 eV than iron, the K-edge of which is 7111 eV. The K-edge of zinc occurs at 9659 eV. Given the ligand content, geometry, and absence of a spectroscopic signal, Zn^{2+} is most likely bound to this position. In addition, it was observed previously that the toluene 2-monooxygenase hydroxylase, a PH family member, is purified with two zinc and four iron atoms (59). In light of this information, we conclude that zinc is the physiologically preferred metal ion bound to the four-cysteine site and suggest that it may be important for stabilizing the fold of the α-subunit C-terminal domain.

Conserved Active Site Residues and Changes in Helix E

Situated just above the diiron center on helix E in PHH are two highly conserved residues found in every BMM, Thr-203 and Asn-204 (Thr-213 and Asn-214 in MMOH, respectively). Thr-203 is positioned at the interface between helices E and F above the Fe2-coordinating Glu-233 (Figure 5A), which undergoes oxidation state-dependent carboxylate shifts in MMOH and ToMOH (48, 49, 57). Asn-204 is located on the surface of the protein, 12 Å from the diiron site. Several ordered water molecules link these Thr and Asn residues to the diiron center by hydrogen bonds. In MMOH and ToMOH, the Thr residue forms the "roof" of the active site pocket. The Asn residue lies between helices E and F just above the iron-coordinating Glu that undergoes a carboxylate shift upon reduction of the diiron center (Figure 5B) (48, 49). The rotameric conformation of this Asn also changes with the active site oxidation state; it points inward toward the diiron center in the mixed-valent and reduced forms and away from it when the iron atoms are oxidized (Figure 5B) (48, 49). Mutagenesis studies on TMOs suggest that the Thr is not essential for catalysis under steady state conditions (60), whereas the Asn is important for turnover and protein component interactions.[5]

The positions of this pair of Thr and Asn residues in both protomers from the native and SeMet PHH structures differ significantly from what has been observed for oxidized, reduced, and mixed-valent MMOH (8,46,48,49) as well as for oxidized and Mn^{2+}-substituted ToMOH (9, 57). The alternative positions that we observe here are the consequence of a different helix E peptide backbone configuration (Figure 5D). Residues 193–202 form a π-helix; residues 205–217 are α-helical, and Thr-203 and Asn-204 mark the transition from the π-to the α-helical forms. In MMOH and ToMOH, the backbone atoms of Thr-213 and Asn-214 assume an α-helical configuration (Figure 5B,F), whereas the residues preceding these two form a π-helix. The alternative configuration of helix E in PHH has been observed once before, in a 6-bromohexan-1-ol-bound structure of MMOH (Figure 5C,E) (33). In this particular MMOH crystal structure, residues 212–216 adopt a π-helical configuration to accommodate the bound product analogue by increasing the volume of the active site pocket. As a result, the orientation of the conserved Thr and Asn residues changes to adopt a configuration nearly identical to the one observed in PHH. We previously hypothesized that such a helical reconfiguration in MMOH may reflect one way by which the regulatory protein induces a change in the active site pocket to alter substrate regiospecificity and accommodate larger substrates. The PHH structure marks the second time that this alternative configuration of helix E has been observed and strongly supports our previous suggestion that the flexibility exhibited by this segment of the α-subunit may play a significant, and largely unexplored, role in the functions of BMMs. Because helix E in all of the PHH α-subunits adopts this configuration, we cannot conclude that PHM is directly responsible for this perturbation without further investigation. If both the conformational changes in helix E and the conserved Asn are relevant to the functions of BMMs, an additional challenge will be to understand when these conformational changes take place during the catalytic cycle.

Structural Changes in Helix F

The electron density for helix F residues 225–231 in the SeMet PHH α-subunit where PHM is bound at higher occupancy is different than that in the other α-subunits (Figure S8), suggesting that PHM may induce a slight reconfiguration, or disordering, of this helix when it is bound to the hydroxylase. The best interpretation of the electron density in this region, which may be a combination of the PHM-bound and unbound forms, is presented in Figure 6. In most α-subunits, F227 and Q230 comprise part of the canyon surface and typically point toward solvent. When the regulatory protein binds, PHM side chains Q10 and I71 occupy the former positions of Q230 and F227, respectively (Figure 6B), and presumably force helix F to reconfigure. The occupancy of PHM in the native structure is insufficient for observation of these differences or identification of the positions of the specific PHM side chains. The major consequences of the helix F rearrangement are the movement of Q230 toward the diiron center, such that it lies above Fe2-coordinating E233, and a shift in the position of the F207 side chain, such that it is closer to the terminal water ligand on Fe1. Residue Q230 is highly conserved among all of the different BMM subfamilies except that of sMMO, where it is a Glu in MMOH and butane monooxygenase (BMOH). It is unknown how mutations of this Gln would affect hydroxylation, electron transfer, or proton transfer chemistry. Residue F207 is conserved among the aromatic hydroxylating BMMs but is a Leu or Ile in the other BMM systems. The conformational changes in helix F do not alter any of the second-sphere hydrogen bonding networks, but they do decrease slightly the solvent accessibility of the diiron center.

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[5]E. Cadieux, L. J. Murray, M. S. McCormick, M. H. Sazinsky, and S. J. Lippard, unpublished results.

Possible Consequences of Regulatory Protein Binding on Transfer of Electrons to the Hydroxylase

Behind the diiron center is an extensive hydrogen bonding network extending from the active site to the surface of the hydroxylase canyon at α-subunit helix A (Figure 3B). Most of the residues contributing to this network are strictly conserved among the different BMM families (3), suggesting that they are critical to the function and folding of the protein. Several other diiron proteins, including bacterioferritin and rubrerythrin, have similar hydrogen bonding patterns that extend 10–14 Å to a redox active cofactor, such as a porphyrin or mononuclear iron–sulfur center (61, 62). In the R2 subunit of RNR, a tryptophan residue in an analogous hydrogen bonding network behind the diiron center facilitates generation of the catalytically important Fe(III)Fe(IV) X intermediate and tyrosyl radical (63). More recently, this pathway in Δ9-desaturase has been implicated in electron transfer (64). On the basis of these observations, we hypothesized that this hydrogen bonding network in BMMs may represent the electron transfer pathway from the reductase to the diiron center (5). Investigations of the electron transfer reactions between MMOH and its reductase, MMOR, indicate the [2Fe-2S] cluster of the latter and the MMOH diiron center may be 11–14 Å apart when this protein complex forms (65). Using this distance constraint, an analysis of the PHH surface structure places the reductase binding site very close to that of the PHM component. If our assumptions about the relative location of reductase binding site are correct, the PH regulatory and regulatory proteins must share a similar binding site on both faces of the hydroxylase surface.

This finding has important implications for how the regulatory protein influences the properties of the hydroxylase diiron center. One attractive scenario is that in which regulatory protein binding prevents the reductase from binding to the hydroxylase and donating extra electrons to the dioxygen-activated diiron center, quenching the peroxo or Q intermediates before they can achieve their hydroxylation chemistry. Such uncoupling, or premature reduction of the oxygenated metal cluster, would not only consume the reactive diiron species but also wastefully deplete the NADH supply of the cell. After completion of the hydroxylation reaction, the regulatory protein may temporarily dissociate from the hydroxylase so that the reductase can bind and reinitiate the catalytic cycle. This scenario would explain why, in the sMMO system, binding of MMOB to MMOH shifts the redox potentials to disfavor reduction of the diiron center, whereas reductase binding shifts the potentials so that reduction is favored (65-68). It is also possible that BMMs function via half-sites reactivity. During the catalytic cycle, the regulatory protein may bind on one face of the hydroxylase to facilitate catalysis while the reductase binds temporarily to the other side to reinitiate the cycle. Such a model may explain why excess regulatory protein often inhibits BMM hydroxylation chemistry, whereas substoichiometric amounts lead to uncoupling (18,19). Pinning down the exact function of the regulatory protein has been a difficult task. Often, it has been termed a "coupling protein" because it couples hydroxylation with electron transfer and an "effector protein" because it effects substrate hydroxylation. The PHH–PHM structure demonstrates how it may carry out both functions.

Access of Substrate to the Diiron Center

The α-subunit scaffold is important for controlling the entrance and egress of small molecules to and from the diiron center. The pathway for substrate entrance, such as the hydrophobic cavities or channels that we previously identified in MMOH and ToMOH, may play an important role in determining the substrate specificity and reactivity of these enzymes (8,9, 33,69). Surface calculations of the PHH α-subunits using PyMOL (70) reveal the presence of three cavities leading from the active site pocket (cavity 1) to the protein surface near the γ-subunit (cavity 3) (Figure 7A). Cavities 1 and 3 have direct access to the exterior environment of the protein, whereas access to cavity 2 is gated by Leu-107 and Leu-272 at one end near the active site pocket and Leu-399 and Val-455 on the other near cavity 3. This putative route for

the access of small molecules to the active site involves mostly hydrophobic residues. It traverses the α-subunit in a manner that is analogous to the pathway defined by the cavities in MMOH and ToMOH but different from the 40 Å channel in ToMOH (9,33,57,69). Although the PHH cavities are large enough to accommodate aromatic substrates, in the absence of structures containing bound product or substrate, it is difficult to know whether these cavities are relevant to the function of PHH.

Surface calculations of the PHH α-subunits without bound PHM reveal a large pore between helices E and F that offers the most direct route for access of small molecules to the active site pocket. This pore is ~6 Å in diameter, lies above the diiron center, and is lined with the side chains of Thr-203, Asn-204, Phe-207, Glu-233, and Met-237 (Figure 7B), all of which are conserved among the different PH enzymes. The pore diameter may be large enough to accommodate phenol and alternative substrates like xylene, dimethylcatechol, and naphthalene, which have diameters between 5.0 and 7.0 Å. Several water molecules occupy the pore and delineate how a substrate may move through the four-helix bundle in the absence of PHM. The 6-bromohexanol-bound form of MMOH, which has a helix E configuration similar to that in PHH, does not have such an obvious mode of entry through helices E and F. The absence of an analogous pore in the MMOH structure may reflect the fact that the peptide backbones of helices E and F near the diiron center are closer together by approximately 2.0 Å, measuring from the C R atoms of Asn-214 and Gln-240 (Figure 5E). In some, but not all, crystal structures of reduced MMOH in which helix E has its typical configuration, a pore with a narrow 2.0 Å diameter forms as a result of Asn-214 (Asn-204 in PHH) shifting to point inward, toward the diiron center (33). This pore may become large enough to accommodate small substrates like methane, dioxygen, and protons in the form of H_3O^+.

Surface calculations of the PHH α-subunits with bound PHM reveal that direct access to the diiron center through helices E and F is restricted in this conformation of the complex (Figure 8). Binding of the regulatory protein also induces F207 and Q230 to reposition as described above, further limiting access to the diiron center. Previously, we suggested a similar barrier-like role for MMOB in the sMMO system because it significantly slowed the rate of diffusion of iron into and out of the active site during metal reconstitution studies (32). Investigations of *Methylosinus trichosporium* OB3b MMOB mutants in which residues 108–111 were changed to Ala revealed variable substrate access to the diiron center. On the basis of these results, MMOB was proposed to generate a size-selective pore on the hydroxylase surface that would serve as a substrate-specific "sieve" (44,71,72). We find no evidence for a PHM-generated substrate-specific pore in the PHH–PHM complex or in models of the MMOH–MMOB structure (not shown), suggesting that the MMOB mutations, which correspond to residues 70–73 in PHM (Figures 6B and S5), may simply decrease the diffusion barrier at the MMOH–MMOB interface to modify the access of the substrate and product to and from the diiron center. Kinetic and computational studies of the reactions of intermediate Q with a variety of substrates have revealed two mechanistic limits, depending upon whether access to the diiron center is rate-determining (73). These results suggest how the regulatory protein might be responsible for controlling the kinetics of these processes in the sMMO system.

The PHM equivalent of S110 in *M. trichosporium* MMOB is S72. The side chain of this residue forms a hydrogen bond to conserved Asn-204 in both the native and SeMet crystal structures (Figures 6B and 8) and is highly conserved among the different BMM regulatory proteins. The link between these two residues has not yet been fully explored. Given that the PH enzymes act on substrates larger than MMOH and that only a deep crevice is observed on the surface of the PHH–PHM complex (Figure 8), the aromatic substrates may enter the PH hydroxylase by an alternative route, possibly via cavity 3. If a pore through the E and F helices of the α-subunit forms at specific stages of the catalytic cycle, such as when the diiron center is reduced and Asn-204 undergoes a rotameric shift, it may allow passage of O_2, H_2O, or H_3O^+ to or from

the active site. Collectively, these findings reveal how the regulatory protein may control the access of small molecules and ions to the diiron center and possibly offers some type of protection of the dioxygen-activated metal center as it traverses the reaction cycle.

Active Site Pocket and Substrate Specificity

The arrangement of side chains in the active site pockets of BMMs is crucial for determining substrate specificity. Phenol hydroxylases, for the most part, are ortho-hydroxylating enzymes, whereas toluene monooxygenases hydroxylate in the para position (58, 74). PHH and ToMOH from *P. stutzeri* OX1, however, have relaxed regiospecificities and typically afford 70% *o*-cresol and 45% *p*-cresol, respectively (20,75,76). For alternative substrates, PH and ToMO perform highly regiospecific reactions. For example, PH and ToMO yield 96% 3-methylcatechol and 95% 4-methylcatechol, respectively, when *m*-cresol is used as the substrate (76,77). The hydrophobic residues that line the active site pockets of PHH and ToMOH are mostly conserved among their specific subfamilies (Table 3) (3). A superimposition of the PHH and ToMOH active sites indicates that the side chains occupy nearly identical positions except for those residues on helix E because of its alternative conformation (Figure 9). Single-ring aromatic substrates can be most effectively docked manually into the active site by orienting their ring plane perpendicular to the Fe–Fe vector. A similar orientation for the aromatic ring has been suggested previously for ToMOH (77). On the basis of DFT calculations and structural investigations of MMOH, substrate hydroxylation presumably occurs at a bridging position between the two iron atoms (78). By docking the oxygen atom of a pentamethylphenol at such a position, we readily see that ortho hydroxylation is favored if a ring substituent points above or below the plane of the iron atoms and bridging oxygen (carbons 2 and 6 in Figure 9), whereas para and meta hydroxylation are favored only if the ring substituent is positioned at the back of the active site pocket. Para hydroxylation may be disfavored in PH enzymes because of steric interaction between Leu-107 and the substituent on the C4 atom (Figure 9). For TMOs, the residue analogous to Leu-107 is typically a glycine or a glutamate (Glu-103) in ToMOH that points away from the active site pocket. A smaller residue at this position would provide enough space to orient a substituent on C4 toward the back of the active site pocket to favor hydroxylation of the para carbon atom. Mutants of both T4MOH and ToMOH in which this residue was changed to leucine afforded 55–60% ortho hydroxylation (58,77), suggesting that the space occupied by this residue is important for steering substrates. Conversely, mutagenesis studies on a related phenol hydroxylase family member demonstrated that changing the residue analogous to Leu-107 to an alanine produced an enzyme that yielded 50% *o*-cresol, compared to 98% for the wild-type form (74). Moreover, this variant was capable of hydroxylating naphthalene, anthracene, fluorene, and phenanthrene at increased rates. Thus, increasing the depth of the active site pocket promotes hydroxylation of larger substrates.

In essence, the PHH active site pocket is shallower than that of ToMOH. Because the regulatory proteins in these different BMM systems alter the product distributions by influencing the structure of the active site pocket (17,18), possibly by changing the configuration of helix E, it is difficult to account fully for all of the residues and structural features that are responsible for orienting substrates in the absence of a crystal structure of the hydroxylase-regulatory protein complex at higher occupancy and in different oxidation states. Nevertheless, our structures of PHH and ToMOH, in addition to revealing the two different configurations of helix E, provide an improved understanding of the active site pockets and an initial structural basis from which one can proceed to rationally engineer these enzymes for industrial and bioremediation applications.

CONCLUSIONS AND PROSPECTS

The fundamental properties that are responsible for controlling the substrate reactivity of the BMM diiron centers lie within the protein scaffold of the hydroxylase subunit. The structure of the PHH–PHM complex represents a key step toward unraveling the principles that underlie the chemistry performed by the carboxylate-bridged diiron centers of these proteins. The structure indicates that PHM functions to block the entrance of the solvent and substrate to the diiron active site through the four-helix bundle, alter the configuration of helices E and F in a manner that could enhance the overall activity of the metal cluster, shape the active site cavity to control product distributions, and most likely affect electron transfer from the reductase by partially or wholly covering its binding site on the hydroxylase canyon walls. Sharing of the hydroxylase surface by the reductase and regulatory protein components would imply that a critical role for the regulatory proteins in all BMMs is to prevent reduction of the dioxygen-activated diiron center at inappropriate times during the catalytic cycle. The recombinant PH enzyme system is ideal for testing the various structural control features, such as hydrogen bonding networks, cavity gates, active site pores, and helix rearrangements, that we suggest may be functionally significant for substrate entrance, electron transfer, and component interactions. Moreover, the PH system will allow future testing of mechanistic theories related to the different reactivities of BMMs.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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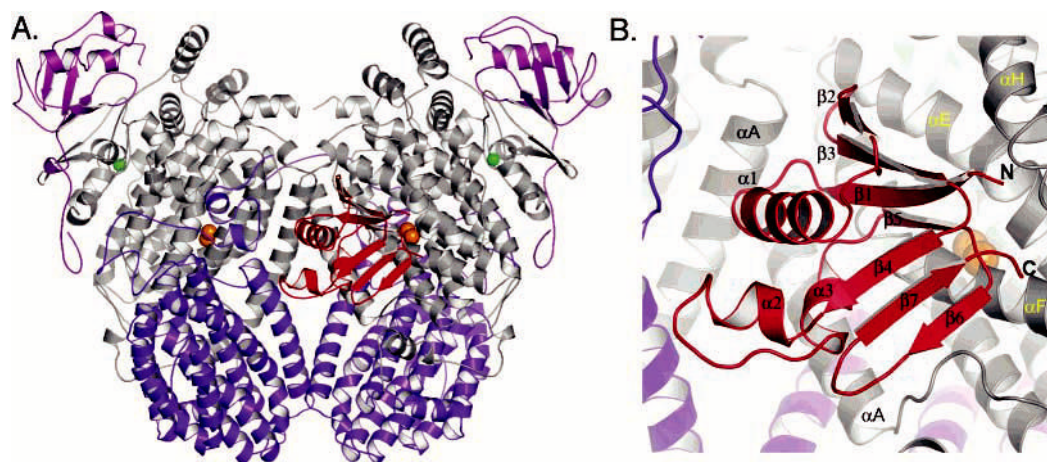
Figure 1.
(A) Global structure of the PHH–PHM complex and (B) enlarged view of PHM in the PHH canyon. The PHH α-, β-, and γ-subunits are colored gray, purple, and pink, respectively, and PHM is colored red. Iron and zinc atoms are depicted as orange and green spheres, respectively. All figures were generated by using PyMOL (70).
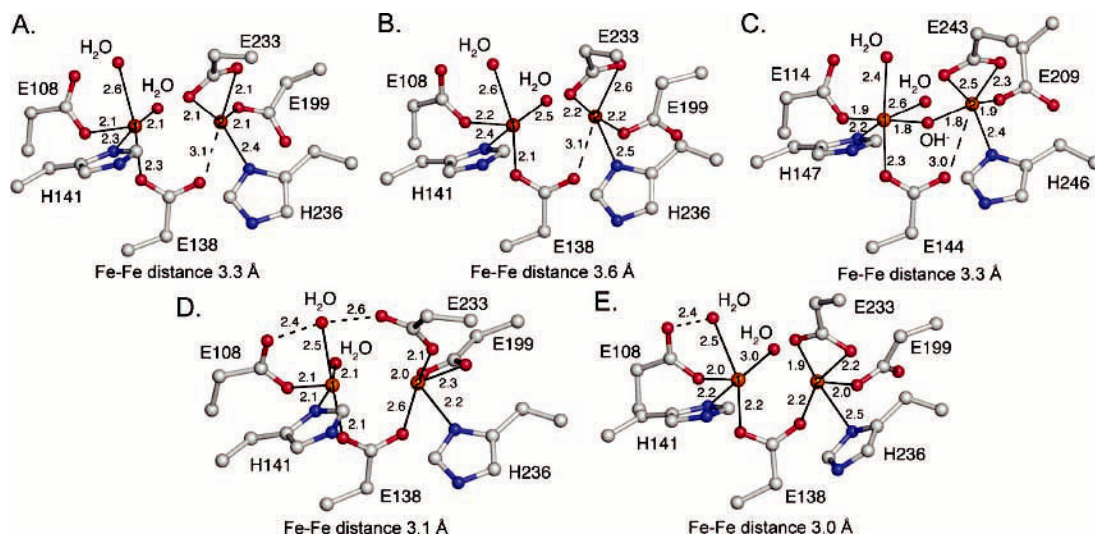
Figure 2.
Structures of the diiron center. (A) Native PH diiron center without PHM bound. (B) Native PH diiron center with PHM bound. (C) Mixed-valent MMOH diiron center. (D) SeMet PH diiron center without PHM bound. (E) SeMet PH diiron center with PHM bound. Carbon, nitrogen, oxygen, and iron atoms are colored gray, blue, red, and orange, respectively. This color scheme will be used throughout unless otherwise noted.
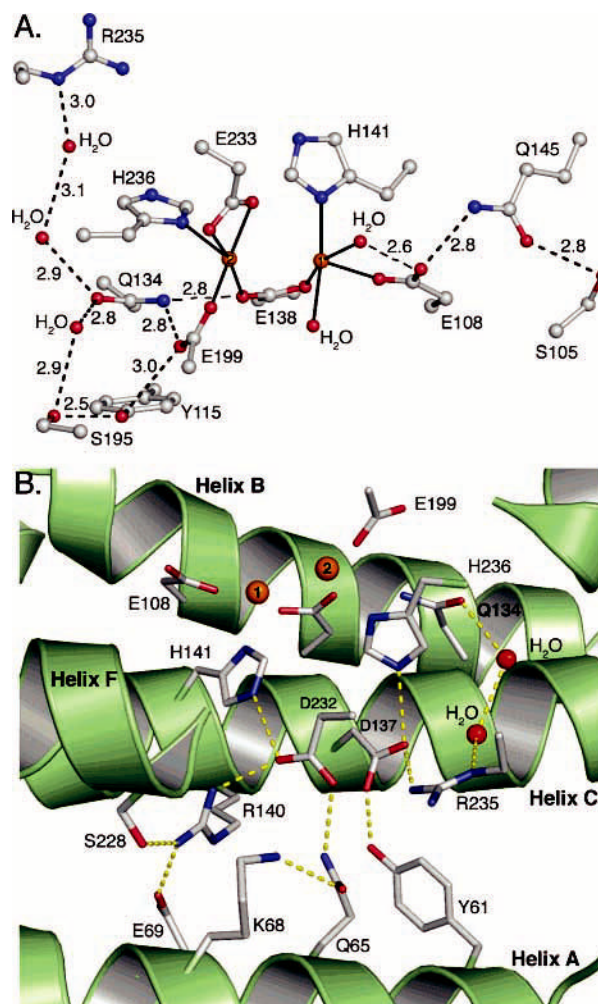
Figure 3.
Hydrogen bonding networks in the second coordination sphere. (A) Hydrogen bonding
networks flanking the diiron center. (B) Hydrogen bonding network behind the diiron center
that extends to the surface of the hydroxylase canyon region where PHM binds.
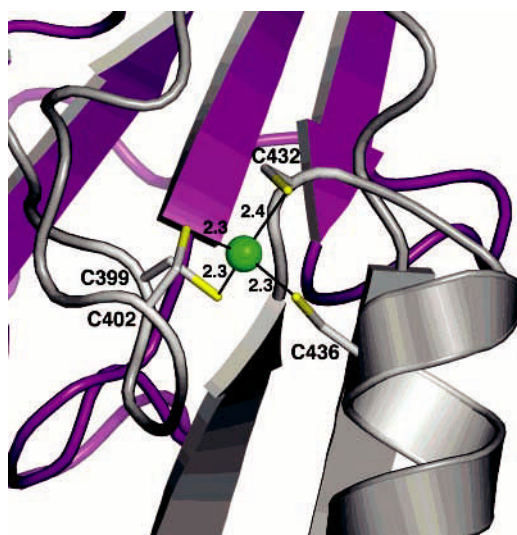
Figure 4.
Zinc binding site in the C-terminal domain of the PHH α-subunit. Zinc is depicted as a green sphere, and sulfur atoms are colored yellow. The α-subunit is depicted as a gray ribbons and the γ-subunit as purple ribbons.
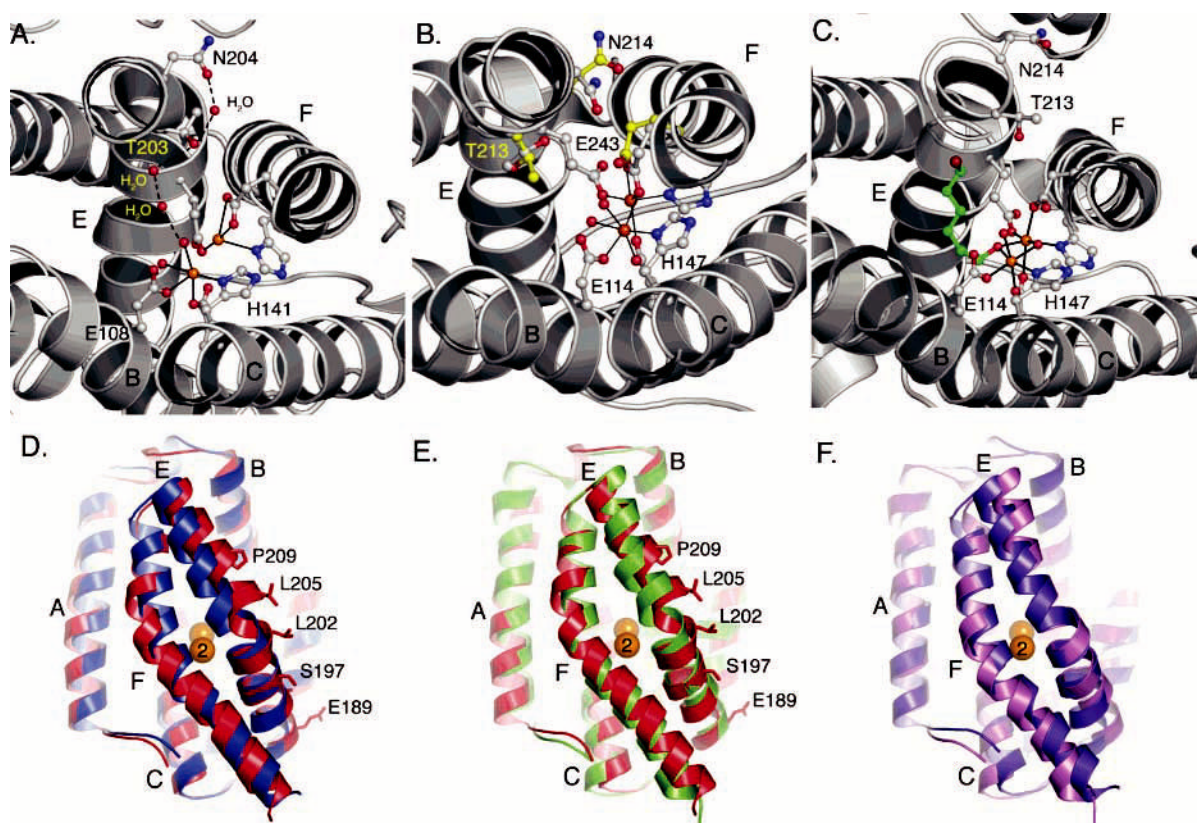
Figure 5.
Structural differences in conserved helix E residues. (A) Positions of conserved Thr-203 and Asn-204 in PHH. (B) Redox-dependent structural changes in the positions of Thr-213, Asn-214, and Glu-243 in MMOH. Carbon atoms for the reduced and oxidized forms of MMOH are colored gray and yellow, respectively. (C) Positions of conserved Thr-213 and Asn-214 in the 6-bromohexan-1-ol-bound form (green) of MMOH in which helix E adopts a more π-helical character. (D) Superimposition of the PHH (red) and ToMOH (blue) four-helix bundles. Helices A–F are labeled accordingly. (E) Superimposition of the four-helix bundle from the 6-bromohexan-1-ol-bound form of MMOH (green) with the PHH (red) four-helix bundle. (F) Superimposition of the ToMOH (violet) and MMOH (pink) four-helix bundles.
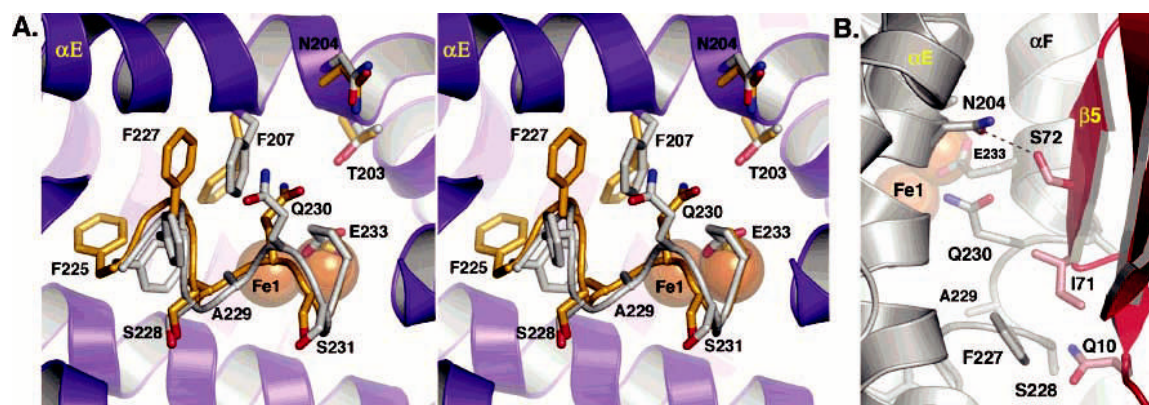
Figure 6.
Structural changes in α-subunit helix F in the SeMet structure. (A) Comparison between PHM-altered (orange) and unaltered (gray) helix F in stereo. (B) PHH (gray) and PHM (red) binding interface at α-subunit helices E and F.
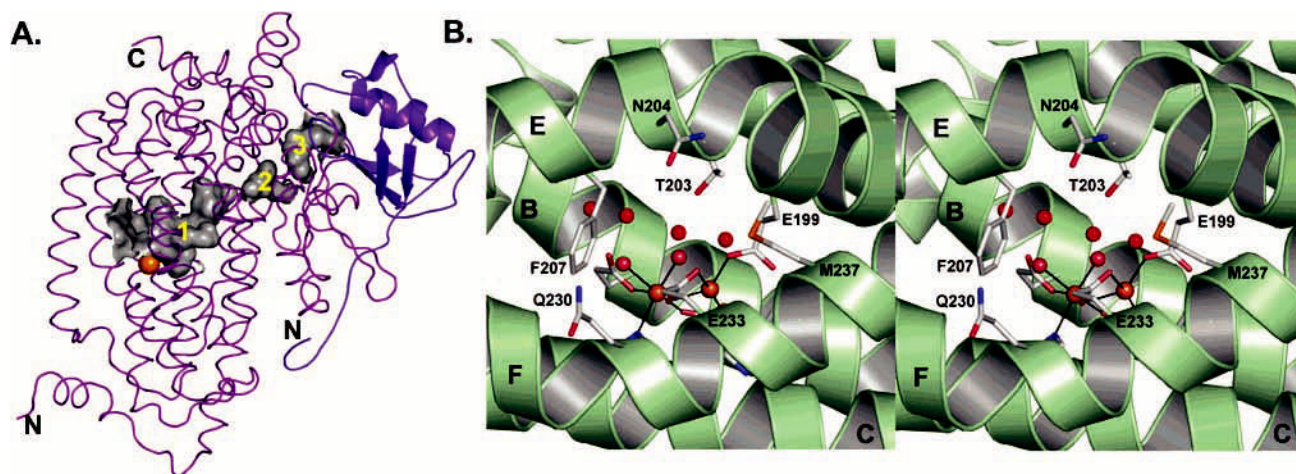
Figure 7.
Cavities and pores in the PHH α-subunit. (A) α-subunit cavities (gray) that extend from the diiron center to the γ-subunit (blue). (B) Stereoview of the conserved residues and water molecules (red spheres) contributing to the pore through the α-subunit four-helix bundle.
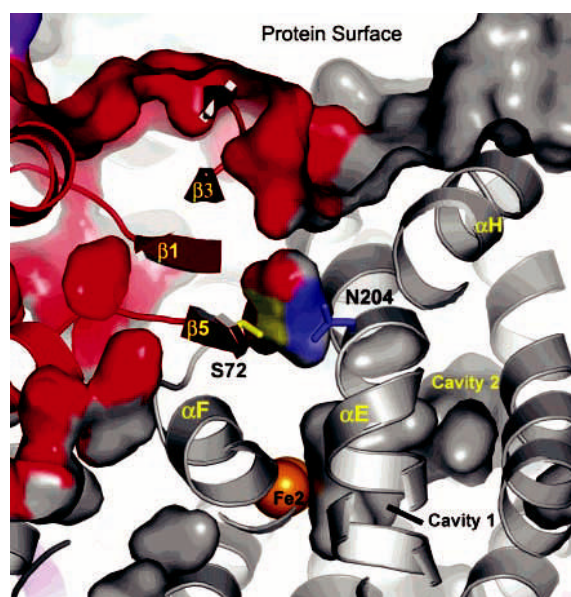
Figure 8.
Access of the substrate to the diiron center at the PHH–PHM interface. PHH (gray) and PHM (red) surfaces and cavities are depicted. Interactions between conserved N204 (blue) and S72 (yellow) and how they may gate access to the metal center are also shown.
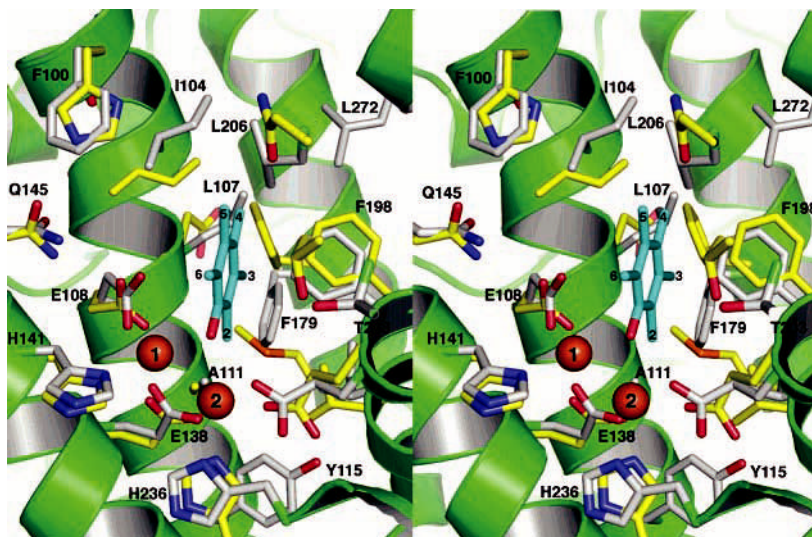
Figure 9.
Stereoview of the superimposed PHH and ToMOH substrate binding pockets. The PHH and ToMOH carbon atoms are colored gray and yellow, respectively, and helices of PHH are depicted in green. Pentamethylphenol (cyan) is docked manually into the active site pocket. The aromatic carbon atoms of the docked aromatics as well as the PHH residues contributing to the active site are labeled. The identity of the ToMOH residues can be obtained from Table 3.

Table 1

Data Collection, Phasing, and Refinement Statistics

	Se SAD	native
data collection		
beamline	SSRL BL9-1	NE-CAT (Sector 8)
wavelength (Å)	0.979	0.979
resolution (Å)	30–2.7	30–2.3
no. of unique observations[a]	109776	108695
total no. of observations[a]	592029	773692
completeness (%)[a]	83.0 (49.8)[b]	100 (100)[b]
redundancy[a]	5.4 (4.3)[b]	7.1 (6.9)[b]
I/σ[a]	17.3 (5.0)[b]	16.5 (4.9)[b]
R_{sym}[a,c] (%)	5.9 (26.0)[b]	8.9 (44.8)[b]
Se sites used for phasing	71 of 77	
figure of merit (after density modification)	0.40 (0.75)	
refinement		
R_{work} (%)[d]	20.2	19.8
R_{free} (%)[e]	25.2	24.2
no. of molecules in ASU	1	1
no. of protein non-hydrogen atoms	16076	15307
no. of non-protein atoms	315	1042
rmsd for bond lengths (Å)	0.033	0.0065
rmsd for bond angles (deg)	2.55	1.28
average B value (Å^2)	36.0	37.7

[a] Values for the Se SAD data were calculated considering I^+ and I^- as separate reflections.

[b] Values in parentheses are for the highestresolution shell.

[c] $R_{sym} = \sum_i \sum_{hkl} |I_i(hkl) - \langle I(hkl)\rangle| / \sum_{hkl} \langle I(hkl)\rangle$, where $I_i(hkl)$ is the ith measured diffraction intensity and $\langle I(hkl)\rangle$ is the mean of the intensity for the Miller index (hkl).

[d] $R_{work} = \sum_{hkl} ||F_o(hkl)| - |F_c(hkl)|| / \sum_{hkl} |F_o(hkl)|$

[e] $R_{free} = R_{work}$ for a test set of reflections (5%).

Table 2

Heavy Atom Anomalous Peak Intensities[a]

subunit	atom	peak height	subunit	atom	peak height
α1	Fe1	11.3	α2	Fe1	11.4
	Fe2	9.62		Fe2	8.9
	Zn	20.2		Zn	22.9

[a]Peak intensities were determined by using CCP4 (28).

Table 3

Residues Comprising the BMM Active Site Substrate Binding Pocket[a]

PHH	I104	L107	A111	Q145	F179	R183[b]	I194	F198	T203	L206	F207
CF600	V105	L108	A112	Q146	F180	R184[b]	V195	F199	T204	L207	F208
TOMH	V106	L109	A113	E147	F181	A185[b]	V196	F200	T205	L208	F209
ToMOH	I100	E103	A107	Q141	F176	M180	L192	F196	T201	Q204	F205
T4MOH	I100	G103	A107	Q141	F176	I180	L192	F196	T201	Q204	F205
AMOH	L91	A94	A98	E133	G167	F172	L184	A188	T193	L196	L197
MMOH	L110	G113	A117	C151	F188	F192	L204	G208	T213	L216	I217

[a] Abbreviations: PHH, *Pseudomonas* sp. OX1 PHH; CF600, *Pseudomonas* sp. strain CF600 PHH; TOMH, *Burkholderia cepacia* G4 toluene *o*-monooxygenase hydroxylase; ToMOH, *Pseudomonas* sp. OX1 ToMOH; T4MOH, *Pseudomonas mendocina* KR1 T4MOH; AMOH, *Rhodococcus corallinus* B-276 alkene monooxygenase hydroxylase; MMOH, *Methylococcus capsulatus* (Bath).

[b] A side chain that does not contribute to the active site pocket yet has a homologous BMM family member with a side chain that does.