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Auteurs: Sonali Dhindwal, Dipak N. Patil, Mahmood Mohammadi, Michel Sylvestre, Shailly Tomar, & Pravindra Kumar
Authors:

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Biochemical Studies and Ligand-bound Structures of Biphenyl Dehydrogenase from *Pandoraea pnomenusa* Strain B-356 Reveal a Basis for Broad Specificity of the Enzyme*

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Sonali Dhindwal[‡], Dipak N. Patil[‡], Mahmood Mohammadi[§], Michel Sylvestre[§], Shailly Tomar[‡], and Pravindra Kumar^{‡1}

From the [‡]Department of Biotechnology, Indian Institute of Technology Roorkee, Roorkee, Uttarakhand 247667, India and the

[§]Institut National de Recherche Scientifique (INRS-Institut Armand-Frappier), Laval, Québec H7V 1B7, Canada

Background: BphB_{B-356} catalyzes the second step of the PCB catabolic pathway.

Result: Apo, binary, intermediate, and ternary structures were obtained.

Conclusion: Conformational changes in the substrate binding loop lead to the formation of a structurally defined pocket to catalyze a wide range of substrates.

Significance: Recognition of conformational changes in the substrate binding loop and insight into the substrate specificity.

Biphenyl dehydrogenase, a member of short-chain dehydrogenase/reductase enzymes, catalyzes the second step of the biphenyl/polychlorinated biphenyls catabolic pathway in bacteria. To understand the molecular basis for the broad substrate specificity of *Pandoraea pnomenusa* strain B-356 biphenyl dehydrogenase (BphB_{B-356}), the crystal structures of the apo-enzyme, the binary complex with NAD⁺, and the ternary complexes with NAD⁺-2,3-dihydroxybiphenyl and NAD⁺-4,4'-dihydroxybiphenyl were determined at 2.2-, 2.5-, 2.4-, and 2.1-Å resolutions, respectively. A crystal structure representing an intermediate state of the enzyme was also obtained in which the substrate binding loop was ordered as compared with the apo and binary forms but it was displaced significantly with respect to the ternary structures. These five structures reveal that the substrate binding loop is highly mobile and that its conformation changes during ligand binding, starting from a disorganized loop in the apo state to a well organized loop structure in the ligand-bound form. Conformational changes are induced during ligand binding; forming a well defined cavity to accommodate a wide variety of substrates. This explains the biochemical data that shows BphB_{B-356} converts the dihydrodiol metabolites of 3,3'-dichlorobiphenyl, 2,4,4'-trichlorobiphenyl, and 2,6-dichlorobiphenyl to their respective dihydroxy metabolites. For the first time, a combination of structural, biochemical, and molecular docking studies of BphB_{B-356} elucidate the unique ability of the enzyme to transform the *cis*-dihydrodiols of double *meta*-, *para*-, and *ortho*-substituted chlorobiphenyls.

Aerobic biodegradation of polychlorinated biphenyls (PCBs)² by bacteria occurs through an oxidative process (1–5).

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¹ To whom correspondence should be addressed. Tel.: 91-01332-285072; E-mail: pravshai@gmail.com.

² The abbreviations used are: PCB, polychlorinated biphenyl; 2,3-DB, 2,3-dihydroxybiphenyl; 4,4'-DB, 4,4'-dihydroxybiphenyl; SDR, short-chain dehydrogenase/oxidoreductase; PDB, Protein Data Bank; r.m.s., root mean square;

The biphenyl degrading pathway, encoded by the *bph* operon, metabolizes several PCB congeners to the corresponding chlorobenzoates (6). This pathway has been found in many bacteria among which *Pandoraea pnomenusa* strain B-356, *Burkholderia xenovorans* strain LB400, *Pseudomonas pseudoalcaligenes* strain KF707, and *Pseudomonas* sp. strain KKS102 have been thoroughly investigated (7–11). The *bph* pathway comprises four enzymes that in *P. pnomenusa* B-356 are encoded by *bphA-EFGBCD*. The biphenyl dioxygenase system (BphAEFG) catalyzes the first reaction, the insertion of two oxygen atoms into vicinal *ortho-meta* carbons of biphenyl to generate *cis*-2,3-dihydro-2,3-dihydroxybiphenyl. *cis*-2,3-Dihydro-2,3-dihydroxybiphenyl dehydrogenase (BphB), the second enzyme of this pathway, catalyzes a dehydrogenation reaction to produce 2,3-dihydroxybiphenyl. 2,3-Dihydroxybiphenyl 1,2-dioxygenase (BphC) and 2-hydroxy-6-oxo-6-phenylhexa-2,4-dienoate hydrolase (BphD) then sequentially metabolize 2,3-dihydroxybiphenyl to chlorobenzoate and 2-oxo-4-pentadienoate as shown in Fig. 1 (3, 5).

Many investigations have shown that the PCB-degrading ability differs among the different PCB-degrading bacteria (2, 6, 10). The biphenyl dioxygenase system critically determines which of the 209 PCB congeners can be metabolized by each strain. For example, a recent study showed that the biphenyl dioxygenase of strain B-356 metabolizes 2,6-dichlorobiphenyl, 4,4'-dichlorobiphenyl, and 2,4,4'-trichlorobiphenyl, whereas the biphenyl dioxygenase of strain LB400 metabolizes these congeners poorly. On the other hand, strain BPDO_{B-356} is unable to catalyze the *meta-para* oxygenation of 2,2',5,5'-tetrachlorobiphenyl, whereas BPDO_{LB400} catalyzes it efficiently. The versatility of BphB toward BPDO metabolites of PCB congeners has not been thoroughly investigated (12). A previous report, however, showed that both BphB_{LB400} and BphB_{B-356} are able to oxidize 3,4-dihydro-3,4-dihydroxy-2,2',5,5'-tetra-

BphB, *cis*-2,3-dihydro-2,3-dihydroxybiphenyl dehydrogenase; BphC, 2,3-dihydroxybiphenyl 1,2-dioxygenase; BphD, 2-hydroxy-6-oxo-6-phenylhexa-2,4-dienoate hydrolase; BPDD, *cis*-(2R,3S)-dihydroxy-1-phenyl-cyclohexa-4,6-diene; BPDO, biphenyl dioxygenase.

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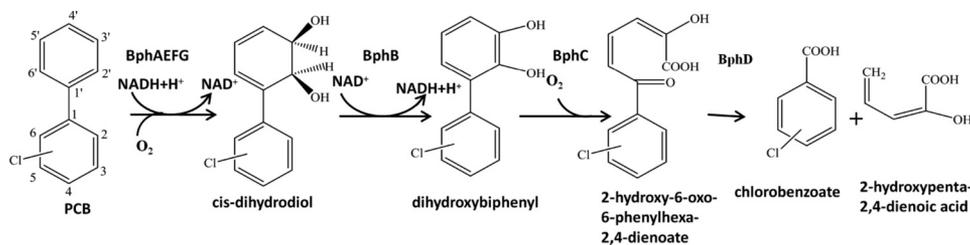


FIGURE 1. Schematic representation of aerobic degradation of biphenyl/polychlorinated degradation catabolic pathway.

chlorobiphenyl, the product resulting from the metabolism of 2,2',5,5'-tetrachlorobiphenyl by BPDO_{LB400} (13). To the best of our knowledge, the ability of BphB_{B-356} to metabolize the product generated from 2,6-dichlorobiphenyl or 2,4,4'-trichlorobiphenyl has never been examined. But recently it has been shown that the dihydrodiol dehydrogenase from *Spingomonas* sp. strain CHY-1 can oxidize a wide range of polyaromatic hydrocarbon dihydrodiols (14).

BphB occurs as a homotetramer comprised of 29.4-kDa subunits. It is an NAD⁺-dependent oxidoreductase, related to other *cis*-dihydrodiol dehydrogenases involved in aromatic degradation pathways and belongs to a very large family of short-chain dehydrogenase (the SDR family) (15, 16). Although, the crystal structure of the NAD⁺-bound form of BphB_{LB400} is known (17), there is no structure available with bound substrate, product, or any product analogs. Furthermore, the substrate binding loop in the published structure of BphB_{LB400} is disordered (17). Therefore, the structural features involved in substrate binding of the *cis*-dihydrodiol dehydrogenases from the aromatic degradation pathways are still undetermined.

To get more insight into the binding mode of the ligand with BphB, we have compared the crystal structure of the apo form of BphB_{B-356} with that of its NAD⁺-bound form (binary state) or of its ternary complex with its coenzyme-NAD⁺ and its product (2,3-dihydroxybiphenyl) or a product analog (4,4'-dihydroxybiphenyl). In addition, we also describe a structure showing an intermediate state of the substrate binding loop. These three-dimensional structures provide insight to the binding mode of ligand with the enzyme. We were able to identify the series of conformational changes in the substrate binding loop that occur during ligand binding. Additionally, our docking studies are consistent with the biochemical experiments that examine the ability of BphB_{B-356} to metabolize and accommodate a large range of chlorinated substrates.

EXPERIMENTAL PROCEDURES

Preparation of BphB Crystals—Purification, crystallization, and preliminary x-ray diffraction properties of BphB_{B-356} have been reported earlier (18). In brief, crystallization conditions for BphB_{B-356} were screened initially using commercial kits by Hampton Research (Hampton Research Inc., Aliso Viejo, CA). We were able to obtain good diffracting crystals in 0.2 M sodium malonate (pH 6.5) buffer containing 10–36% (w/v) PEG-3350.

Diffraction data for crystals of substrate-free and NAD⁺-bound forms of the enzyme were collected on an in-house x-ray setup. Before data collection, the protein crystals were cryoprotected by direct transfer to the mother liquor drop containing

10% (v/v) ethylene glycol as a cryoprotectant. Crystals of the BphB_{B-356}-NAD⁺ complex were prepared by soaking protein crystals for 15 min at room temperature in a solution comprised of 22% (w/v) PEG-3350, 0.2 M sodium malonate, and 10 mM NAD⁺. Diffraction data were collected under cryogenic conditions (~100 K).

Protein crystals were also soaked (for 50 s to 15 min) with 2,3-dihydroxybiphenyl (23-DB), the reaction product in the same solution. Similar procedures were used to soak the 4,4'-dihydroxybiphenyl (44-DB), a product analog. 2,3-Dihydroxybiphenyl and 4,4'-dihydroxybiphenyl were from Sigma and Himedia Laboratories Pvt. Ltd., respectively. Data for these crystals were collected at a synchrotron radiation source, ESRF Grenoble, at beamline 14. All the crystals that were used for soaking experiments were grown under similar conditions as that of the apocrystals.

For all crystal structures, the diffraction data were indexed, integrated, and scaled using the HKL2000 program suite (19). Initial phases for BphB_{B-356} were obtained by molecular replacement using MOLREP from the CCP4 version 6.0 software suite (20). The crystal structure of BphB_{LB400} was used as a search model (Protein Data Bank (PDB) code 1BDB) (21). Rigid body refinement was followed by iterative cycles of restrained atomic parameter refinement using the programs CNS (22) and REFMAC_5.2 (23). The program COOT was used for analysis of electron density maps and model building (24). The difference Fourier map for the NAD⁺-bound form of BphB_{B-356} clearly showed density for the NAD⁺ at above the 3 σ level, allowing NAD⁺ to be modeled into the density. Solvent molecules were added where the $F_o - F_c$ map had values above 3 σ and the $2F_o - 2F_c$ map showed a density at the 1 σ level.

Several cycles of rigid-body refinement and then restrained refinement were used to achieve acceptable R_{cryst} and R_{free} . Stereochemical properties of the models were evaluated using the program PROCHECK (25). Figures were prepared using the program PyMOL (26).

Docking Study—Molecular docking of all the substrates was carried out using the Schrödinger, Maestro suite, version 9.1 (Glide, version 2.6, Schrodinger, Inc., New York). The protein was prepared using the Maestro protein preparation wizard by addition of hydrogens, assigning the bond orders, and minimization of the protein using the OPLS2001 force-field until the r.m.s. deviation between the minimized structure and the starting structure reached 0.3 Å. All substrates were prepared using the Maestro Ligprep module (Ligprep, Schrodinger, Inc.). A receptor grid for docking was generated by the centroids of the selected residues, Ser-142, Tyr-155, Lys-159, Gly-150, Asn-149,

TABLE 1

Data collection and refinement statistics for BphB_{B-356} in apo form, in complex with NAD⁺, in complex with NAD⁺-2,3-dihydroxybiphenyl, and in complex with NAD⁺-4,4'-dihydroxybiphenyl

	BphB _{B-356} ⁻ apo	BphB _{B-356} ⁻ binary	BphB _{B-356} ⁻ intermediate	BphB _{B-356} ⁻ -ternary (2,3-dihydroxy biphenyl)	BphB _{B-356} ⁻ -ternary (4,4'-dihydroxy biphenyl)
Crystallographic data					
Space group	<i>P</i> ₄ -2 ₁ 2	<i>P</i> ₄ -2 ₁ 2	<i>P</i> ₄ -2 ₁ 2	<i>P</i> ₄ -2 ₁ 2	<i>P</i> ₄ -2 ₁ 2
Resolution	2.2	2.1	2.9	2.4	2.1
Cell dimensions					
<i>a</i> (Å)	75.7	76.1	75.7	75.4	76.2
<i>b</i> (Å)	75.7	76.1	75.7	75.4	76.2
<i>c</i> (Å)	176.9	178.0	179.0	173.6	180.6
Unique reflections	25,874	18,429	11,486	20,162	30,090
Completeness (%) (last shell)	97.0 (40.6)	97.1 (99.8)	99.6 (99.9)	95.0 (99.4)	99.9 (100.0)
<i>R</i> _{sym} (%) ^a (last shell)	10.6 (38.7)	6.5 (27.9)	13.2 (50.7)	10.4 (61.2)	12.0 (63.3)
<i>I</i> / <i>σ</i> (last shell)	13.6 (2.4)	13.8 (2.2)	8.3 (3.7)	19.3 (2.3)	19.4 (2.5)
Multiplicity (last shell)	5.5 (2.9)	2.7 (2.2)	5.8 (3.5)	12.1 (7.7)	8.9 (8.7)
Refinement					
No. of Residues	536	534	534	543	543
Water molecules	231	264	39	132	182
Resolution range (Å)	70.0–2.2	10.0–2.5	70.0–2.9	70.0–2.4	70.0–2.1
<i>R</i> _{cryst} (%)	18.1	21.7	26.8	20.8	18.7
<i>R</i> _{free} (%)	21.9	28.6	29.5	26.6	22.6
Average <i>B</i> -factors (Å ²)	A 38.5 B 38.9	A 22.5 B 20.1	A 59.9 B 59.9	A 41.7 B 41.4	A 38.3 B 38.5
Water atoms	42.7	23.8	30.6	47.2	41.4
All atoms	38.9	21.9	59.6	41.7	38.7
R.m.s. deviations on bond lengths (Å)	0.008	0.006	0.007	0.016	0.014
R.m.s. deviations bond angles (Å)	1.2	0.9	1.0	1.8	1.5
Ramachandran plot (%)					
Preferred	90.4	89.2	87.6	88.7	90.5
Allowed	9.5	10.9	12.4	11.3	9.0
Outliers	0.0	0.0	0.0	0.0	0.0

$$^a R_{\text{sym}} = \frac{\sum_{hkl} \sum_{i=1}^n |I_{hkl,i} - \overline{I_{hkl}}|}{\sum_{hkl} \sum_{i=1}^n I_{hkl,i}}$$

and the NAD⁺ molecule. These residues and the NAD⁺ form the active site of the protein (27). Glide was then used for docking the substrates using the Extra Precision (XP) mode. The best conformation was selected on the basis of Glide score and by visually inspecting the molecule (28).

Whole Cell Assays to Identify Metabolites of Chlorobiphenyls—Metabolites from 2,6-dichlorobiphenyl, 2,4,4'-trichlorobiphenyl, and 3,3'-dichlorobiphenyl (all 99% pure, from Ultra Scientific, Kingstown, RI) were analyzed according to a previously published protocol from suspensions of isopropyl β-D-1-thiogalactopyranoside-induced *Escherichia coli* DH11S pDB31[LB400-*bphFG*] harboring pQE31[B-356-*bphAE*] or a mixture containing equal amounts of isopropyl β-D-1-thiogalactopyranoside-induced *E. coli* DH11S pDB31[LB400-*bphFG*] harboring pQE31[B-356-*bphAE*] plus isopropyl β-D-1-thiogalactopyranoside-induced *E. coli* DH11S pQE31[B-356-*bphB*] (29). In the case of 2,6-dichlorobiphenyl, pQE31[B-356-*bphAE*] was replaced by pQE31[*p4-bphAE*]. The level of expression was assessed by inspection of SDS-PAGE gels. Metabolites were identified by gas chromatography-mass spectrometry (GC-MS) analysis of their butylboronate derivatives (29).

Protein Data Bank Accession Numbers—The accession codes used are 2Y93 (BphB_{B-356} apo-structure), 2Y99 (BphB_{B-356}-NAD⁺ complex), 3ZV6 (BphB_{B-356}-NAD⁺-4,4'-dihydroxybiphenyl complex), 3ZV5 (BphB_{B-356}-NAD⁺-2,3-dihydroxybiphenyl complex), and 3ZV3 (BphB_{B-356} intermediate structure).

RESULTS

Quality of the Structure and Structure Determination—The structures of the ligand-free form of BphB_{B-356} (apo), the binary

complex with NAD⁺, and the ternary complex with NAD⁺ and product (23-DB) or product analog (44-DB) were determined. The crystallographic data collection and refinement statistics for all of the structures are summarized in Table 1.

The refined model of the ligand-free BphB_{B-356} includes residues from Met¹ to Gly¹⁹⁸ and Ser²⁰⁶ to Ile²⁷⁵ in both subunits. This model lacks seven residues from 199 to 205, which form a loop at the mouth of the active site (the substrate binding loop) and six residues from the C-terminal due to insufficient interpretable electron density.

The structure of the BphB_{B-356}-NAD⁺ complex was determined at 2.5 Å resolution. An NAD⁺ moiety was clearly visible in the initial difference Fourier maps in each subunit. Binding of NAD⁺ at the coenzyme binding site and near the substrate binding site required only local structural adjustment and thus, did not alter the overall conformation of BphB_{B-356}. The loop region from Leu¹⁹⁹ to Ser²⁰⁵ is disordered in this binary structure also.

The structures of the ternary complexes of BphB_{B-356} with NAD⁺ and 23-DB and BphB_{B-356} with NAD⁺ and 44-DB were obtained at 2.4 and 2.1 Å, respectively. Difference Fourier maps distinctly showed the presence of NAD⁺ in both structures. Additionally, in both structures extra electron density was present at the active site near the nicotinamide ring of NAD⁺. This density was due to the presence of 23-DB and 44-DB, respectively.

Surprisingly, for both ternary structures the electron density was clearly visible for the entire substrate binding loop from Thr¹⁸⁷ to Val²¹⁶. Thus, the disordered region (Leu¹⁹⁹ to Ser²⁰⁵) in the ligand-free form and binary complex was ordered in the

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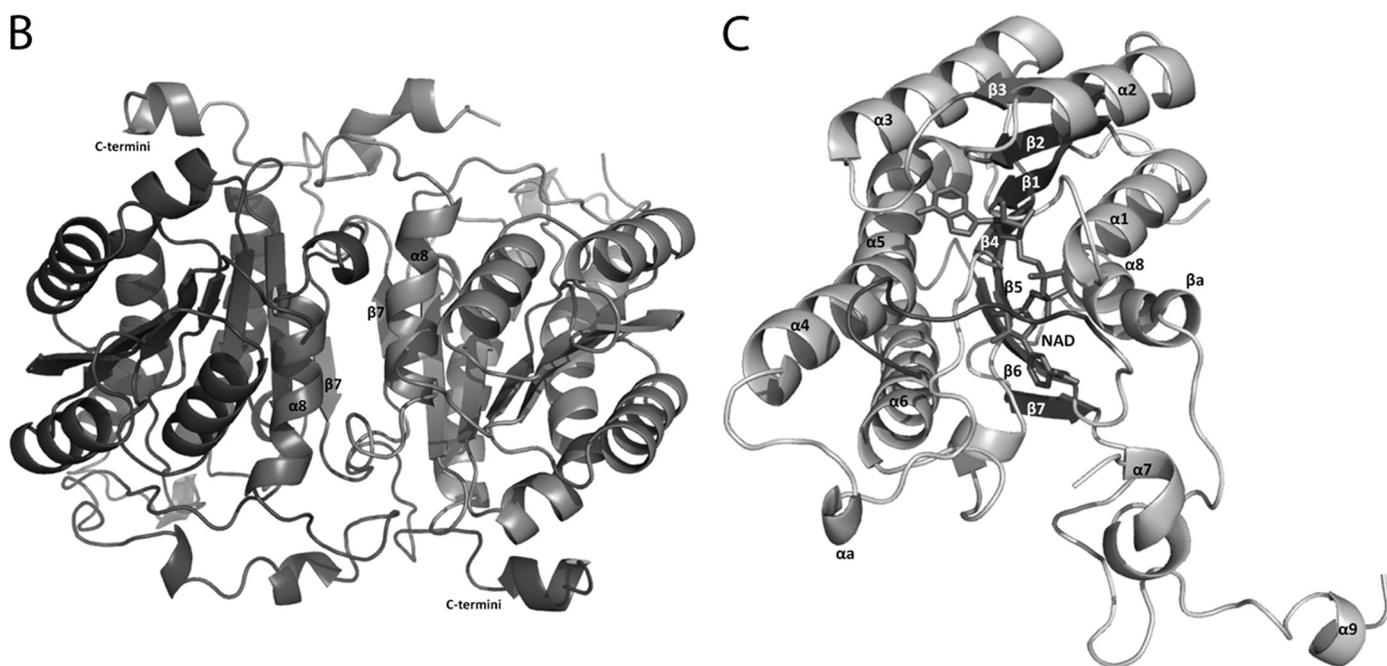
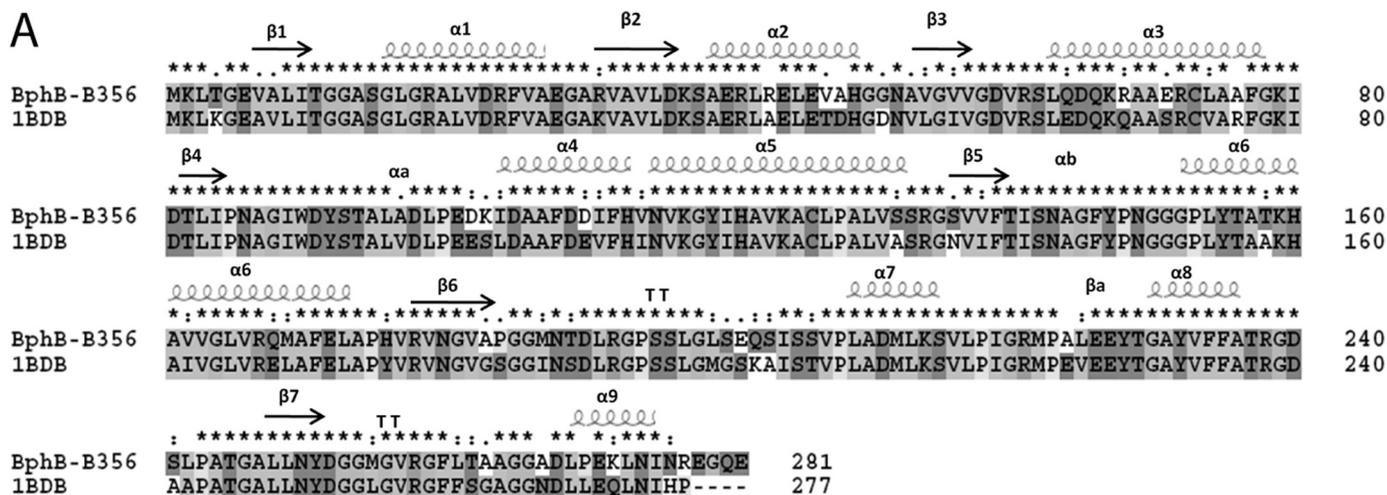


FIGURE 2. A, sequence alignment and secondary structure assignment of BphB_{B-356} and BphB_{LB400} (PDB code 1BDB). This figure was made using ClustalX (43). B, schematic diagram of BphB_{B-356} in dimeric form. The interface of the two monomers comprised of helix $\alpha 8$ and sheet $\beta 7$ from both subunits A and B is indicated. C, the overall structure of BphB_{B-356} as a monomer. Bound NAD⁺ is shown in stick form. The substrate binding loop (residues 189–197) near the active site is highlighted by the black color loop.

ternary complexes. The refined models contain two molecules per asymmetric unit for both ternary complexes.

Overall Structure—BphB_{B-356} exists as a tetramer made up of two asymmetric units. The tetrameric state of BphB_{B-356} was also confirmed by gel filtration chromatography. The two monomers in an asymmetric unit are related to each other by a noncrystallographic 2-fold axis.

As per the conventional nomenclature of SDR enzymes, the subunit contacts are named based on the three perpendicular axes *P*, *Q*, and *R* along the tetrameric arrangement of monomers (30). In BphB_{B-356}, the monomers in an asymmetric unit coincide by the *P* axis. The two monomers along this axis interact through their respective $\alpha 8$ helix and $\beta 7$ sheet as shown in Fig. 2B. The contacts between the two vicinal helices occur principally through hydrophobic interactions involving Phe²³⁴ and Phe²³⁵ from each subunit. Between $\beta 7$ sheets, no direct

interaction is seen and most of the interactions are mediated through water molecules.

The overall structure of the BphB_{B-356} monomer is similar to the previously reported structure of BphB_{LB400} (17). It can be divided into three parts: the main body (Met¹-Gly¹⁸⁶ and Leu²¹⁷-Tyr²⁵¹) arranged in a α/β folding pattern, consisting of a dinucleotide binding motif called the Rossmann-fold, the substrate binding loop (Met¹⁸⁷-Val²¹⁶), and the C-terminal region (Asp²⁵²-Ile²⁷⁵).

BphB_{B-356} monomer displays a highly similar α/β folding pattern, consisting of a dinucleotide binding motif comprised of a central β -sheet flanked by α -helices. The central unit is made up of seven core parallel β -sheets (residues 7–11($\beta 1$), 31–36($\beta 2$), 53–57($\beta 3$), 82–84($\beta 4$), 136–140($\beta 5$), 178–184($\beta 6$), and 248–251($\beta 7$)) buried between six α -helices, three on each side $\alpha 8$ (230–236), $\alpha 1$ (15–28), $\alpha 2$ (39–48) and $\alpha 3$ (63–

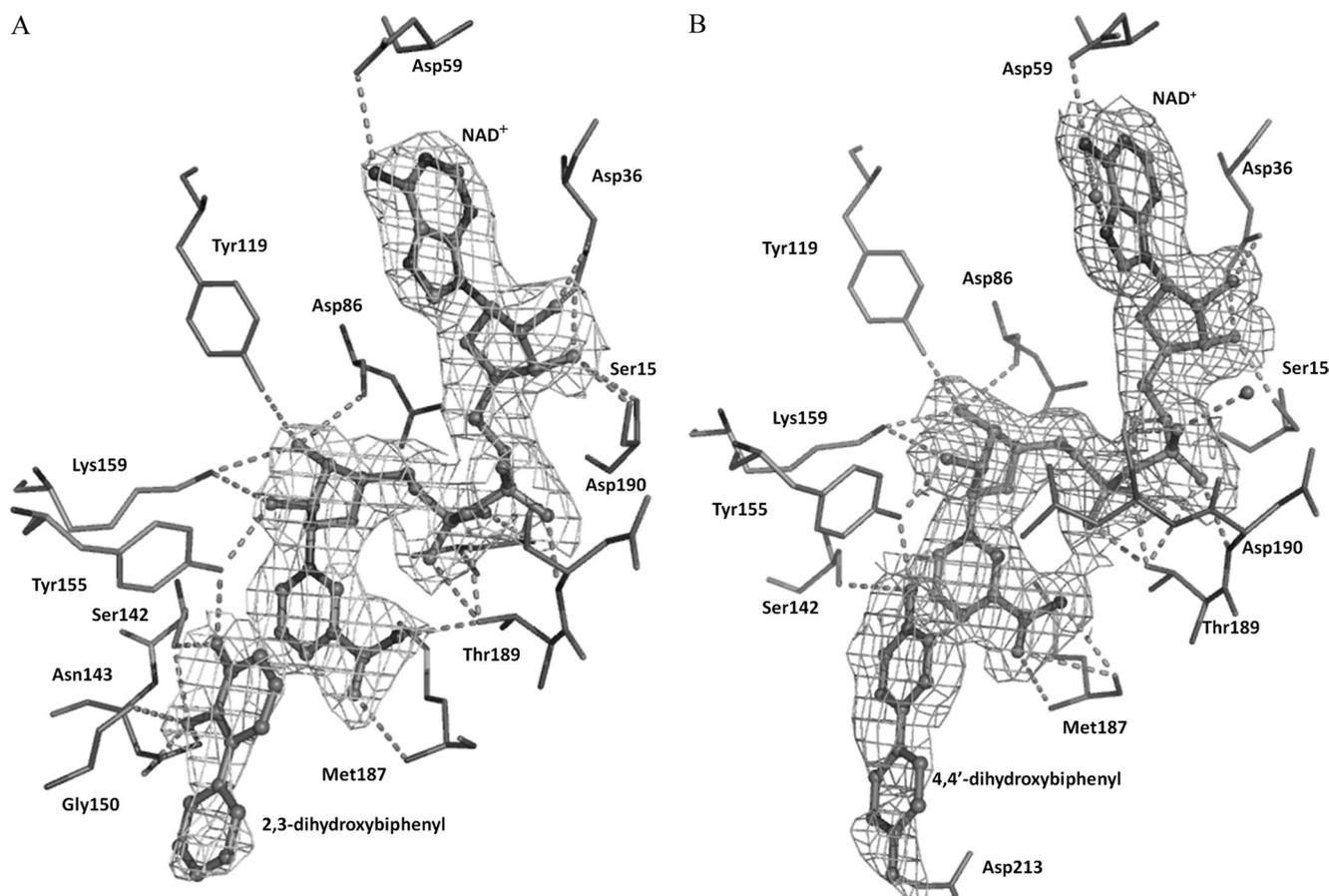


FIGURE 3. Displaying the bound NAD^+ in $\text{BphB}_{\text{B-356}}$ with (A) 2,3-dihydroxybiphenyl and (B) 4,4'-dihydroxybiphenyl (shown as ball and stick models) with $2F_{\text{obs}} - F_{\text{calc}}$ electron density contoured at 1σ and 0.8σ , respectively. Residues at hydrogen bonding distance to NAD^+ and the product/product analog are shown as stick models.

77), $\alpha 5$ (115–133), $\alpha 6$ (153–173) as shown in Fig. 2, A and C. The two helices, αa and αb , and turn βa protrude out of the main body. The region between sheet $\beta 6$ and helix $\alpha 8$ is the substrate binding region and corresponds to the disordered region (Leu¹⁹⁹–Ser²⁰⁵) in the structures of the apoenzyme and binary complex.

The C-terminal region consists of a loop and a small helix oriented above the $\alpha 5$ and $\alpha 6$ helices of the vicinal molecule in the asymmetric unit as shown in Fig. 2B. For each subunit, Gly²⁶⁵, Gly²⁶⁶, and Ala²⁶⁷ of the C-terminal region interacts with Phe¹⁷¹, Ala¹⁷⁴, and His¹⁷⁶, respectively, located on helix $\alpha 6$ of the vicinal subunit. The involvement of the C termini in the subunit-subunit interaction could be of functional significance in tetrameric assembly.

Binary Complex with Coenzyme NAD^+ —In the binary complex of the protein, both subunits in an asymmetric unit contain NAD^+ bound at the coenzyme binding site. NAD^+ is visible in the electron density map and is bound in a similar fashion as in other SDR family enzymes. It is placed with its adenosine ribosyl-OH group held by Asp³⁶ and its nicotinamide ring toward the active site of the enzyme where the nicotinamide ring lies parallel to the indole ring of Trp⁹⁰. NAD^+ interacts with the protein in a similar manner as in $\text{BphB}_{\text{LB400}}$ (17). Superposition of the binary structure and the apo form gives an r.m.s deviation of 0.3 Å (3559 to 3559 atoms). The structure shows a local conformational change in the C $^{\alpha}$ chain and side chains of some

of the residues surrounding the coenzyme binding site allowing the accommodation of NAD^+ in the cavity. These conformational changes are visible at the substrate binding loop from Met¹⁸⁷ to Arg¹⁹². As compared with the apo form, these amino acids move toward the pyrophosphate moiety and nicotinamide ring of NAD^+ .

Ternary Complex with 2,3-Dihydroxybiphenyl and 4,4'-Dihydroxybiphenyl—Crystal complexes of $\text{BphB}_{\text{B-356}}$ -44-DB and $\text{BphB}_{\text{B-356}}$ -23-DB were prepared by soaking with NAD^+ and product or product analog for about 15 min. The difference Fourier maps distinctly showed the presence of NAD^+ in both structures and also showed bulky electron density in the active site of the complexes. 23-DB and 44-DB molecules were modeled in the available electron density at the active site for their respective complexes. The refined ternary complexes of the $\text{BphB}_{\text{B-356}}$ crystal structures show that product and product analogs bind to the active site essentially in the same manner (Fig. 3, A and B). As compared with the apo and binary forms of the enzyme, the ternary structure shows conformational changes in the residues surrounding the active site and the substrate binding loop. The distal rings of 23-DB and 44-DB were not very well defined in the electron density map and have higher *B*-factor than the average *B*-factor. To confirm the product/product analog binding to the active site, the ternary structures were refined without adding the coordinates of the product or product analog and it was observed that the

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ternary structure still has conformational changes. This suggests that substrates, product, or product analogs require conformational changes in the enzyme upon binding. This has also been mentioned for other enzymes that belong to the SDR family (31–34). It has been reported earlier that conformational changes occur in the substrate binding loop to shield the active site from bulk solvent, which is essential to allow hydride transfer during the catalysis (34, 35). It could also be seen that as opposed to the BphB_{B-356} structure in the apo and binary forms, the active site in the ternary structure is devoid of any water molecule as shown in Fig. 4, A–C.

Superposition of C α atoms of the BphB_{B-356}-44-DB crystal structure with that of the apo-enzyme and binary structure of BphB_{B-356} exhibit r.m.s deviations of 0.3 Å (3530 to 3530 atoms) and 0.3 Å (3526 to 3526 atoms), respectively. In the case of the BphB_{B-356}-23-DB structure, the r.m.s. deviations compared with the apoenzyme and the binary structure were 0.4 Å (3550 to 3550 atoms) and 0.4 Å (3557 to 3557 atoms), respectively. The overall structure of each ternary complex is similar to the binary structure of BphB_{B-356}, except that the substrate binding region (from residue Gly¹⁹⁸ to Ser²⁰⁶) is complete and clearly visible in the electron density map for both ternary complexes.

In the ternary complex, the complete substrate binding loop comes close to the active site and forms a compact cavity where the product/product analog binds. The same residues interact with NAD⁺ in the ternary and binary structures. However, conformational changes of residues from the substrate binding loop have been observed. Residues from Gly¹⁸⁶ to Leu¹⁹¹ (which are also modeled in the binary structure), now make more interactions with NAD⁺. Met¹⁸⁷ forms two H-bonds from its main chain atoms to the amide group of the nicotinamide ring. This interaction was not present in the NAD⁺-bound structure (binary complex). The nicotinamide ring of the NAD⁺ makes hydrophobic interaction with Leu¹⁹¹, Ile²⁰⁴, and Leu²⁰⁹ in the ternary complexes, but these interactions are not seen in the binary complex.

Active Site Architecture—The active site of the enzyme is formed by a tetrad of residues (Ser¹⁴², Asn¹¹⁵, Tyr¹⁵⁵, and Lys¹⁴⁹) and the nicotinamide ring of NAD⁺ (13, 36, 37). The substrate binding loop in the ternary complex forms a cavity at the nicotinamide end of the NAD⁺ to accommodate 44-DB/23-DB.

The product 23-DB fits well at the active site in a deep hydrophobic cleft. The hydroxylated ring of 23-DB places itself between the NAD⁺ nicotinamide ring and the indole ring of Trp⁹⁰. Ser¹⁴², one of the catalytic residues at the loop between β 5 and α 6, holds the product by formation of two hydrogen bonds with 23-DB, one with its 2-OH group (Ser-OH—23DB-2OH = 3.5 Å) and the other with its 3-OH group (Ser-OH—23DB-3OH = 2.1 Å). The side chain of Asn¹⁴³ makes a H-bond with the 2-OH of 23-DB (Asn¹⁴³-ND2—23DB-2OH = 2.6 Å and Asn¹⁴³-OD1—23DB-2OH = 2.5 Å). Finally, the hydroxyl group of Tyr¹⁵⁵ interacts with the 3-OH of 23-DB (Tyr-OH—2OH-23DB = 3.2 Å).

The distal part of the ligand binding cavity is surrounded by hydrophobic residues to provide an environment suitable to accommodate the nonpolar portion of the product. These res-

idues are Gly¹⁵⁰, Gly¹⁸⁶, Ile²⁰⁴, Leu²⁰⁹, Asp²¹³, Met²¹², and Met²⁵⁵.

With regard to the other ternary complex, 44-DB is also located at the active site of enzyme. The hydroxyl group of the ring that is directed toward NAD⁺ forms H-bonds with Tyr¹⁵⁵ and Ser¹⁴². As the other ring also has one -OH group, 44-DB is displaced compared with 23-DB to form a H-bond with Asp²¹³. The hydrophobic interactions observed in 44-DB are similar to those occurring in the 23-DB-bound form of the enzyme.

The Intermediate State of the Substrate Binding Loop—We obtained a crystal structure of BphB_{B-356} where the soaking time of the crystal with NAD⁺ and product was less than a minute. In this structure we could not find proper density for either NAD⁺ or 23-DB, but did observe electron density for the complete substrate binding loop from residues Gly¹⁹⁸ to Ser²⁰⁶.

A comparison of this structure and ternary structures bound with 23-DB or 44-DB shows that the substrate binding loop occupies an entirely different space within the crystal structure. The distance between main chain carbons within the substrate binding loop of this short-soak structure and the ternary structures of 12.5 Å at Glu²⁰¹ to 15.9 Å at Ser²⁰⁵ are shown in Fig. 5. The position of the substrate binding loop in these two structures is similar to the opposite extremes of a arc of the swing.

As the soaking time was very short, this state could be said to have captured the structure of the substrate binding loop as an intermediate step between that of the apoenzyme and the ternary structure. The substrate binding loop was disordered in the apoenzyme and binary structure, whereas in this structure the substrate binding loop is well organized, perhaps having caught the enzyme in a process of ordering its substrate binding loop in the presence of product prior to its movement toward the active site to form a proper cavity as found in the ternary structure. The snapshots of the ligand binding loop in three different states are clearly shown in Fig. 5. A surface view of apo, binary, ternary, and intermediate states is shown in Fig. 4, exhibiting the differences in structure of the different enzyme forms.

Comparison of BphB_{B-356} with Other Enzymes in SDR Family—Structural and sequence alignments confirm that BphB_{B-356} is very similar to BphB_{LB400}, with some variations in the active site region. Comparison of the binary structure with BphB_{LB400} (PDB code 1BDB) shows that most of the structural differences are observed at the N- and C-terminal regions of the protein and in the substrate binding loop region. The BphB_{LB400} structure is available in one form only, the binary structure complex with bound coenzyme. Its superposition with the binary structure of BphB_{B-356} gives an r.m.s. deviation of 0.5 Å (1666 to 1666 atoms). The relative positions of the residues that form the active site, Ser¹⁴², Tyr¹⁵⁵, Lys¹⁵⁹, Gly¹⁵⁰, and Asn¹⁴³, are similar in both molecules. The orientation of the NAD⁺ molecule in both BphB_{LB400} and binary BphB_{B-356} is almost the same.

A Blastp and Dali search for structurally related proteins against BphB_{B-356} shows that the closest homologous proteins belong to the SDR family (38). Despite their low sequence identity, these proteins share very well conserved overall topology. The major difference lies in the structure of the substrate binding loop as shown in Fig. 6. Structural alignment of these sequences shows that the substrate binding region in other pro-

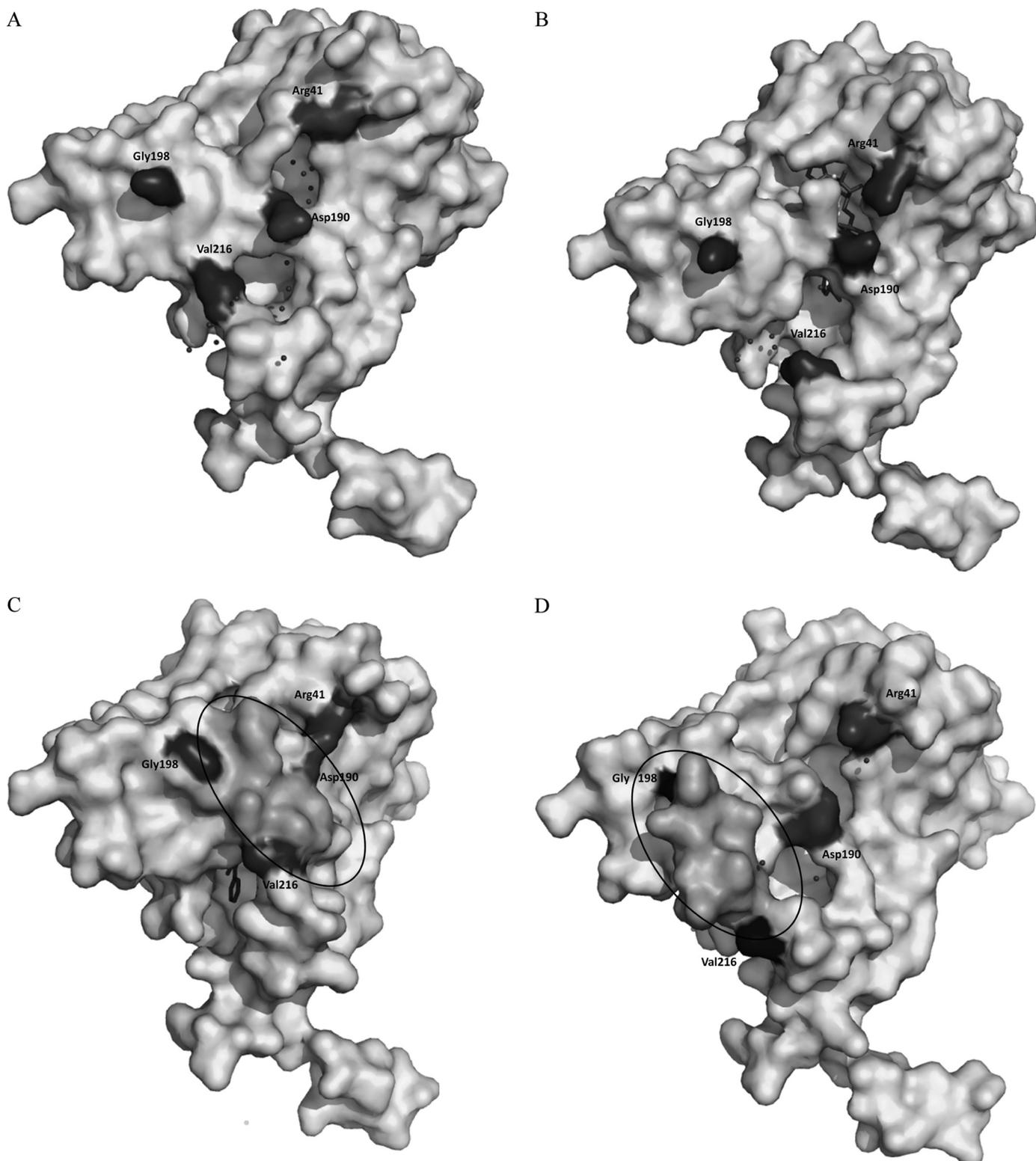


FIGURE 4. The surface representation of BphB_{B-356} in (A) apo form, (B) binary form, and (C) ternary structures in the presence of 23-DB and the (D) intermediate state. Residues Asp¹⁹⁰ and Arg⁴¹ are highlighted; in apo and binary structures they appear as separate residues, whereas in the ternary structure they appear to be joined due to the interaction between them. The circled region is part of substrate binding loop between Gly¹⁹⁸ and Val²¹⁶, which is not visible in apo and binary forms but is visible in the ternary and intermediate structures. Shift in the substrate binding loop in the ternary and intermediate regions could be seen. The spheres surrounding the coenzyme binding site and the active site show water molecules.

teins is made up of two helices joined by a small loop, unlike the elongated loop-like structure found in BphB_{B-356}. The one exception is the alcohol dehydrogenase from *Drosophila leba-*

nonensis (PDB code 1B15), which was not found in the Blastp or Dali search. It has been mentioned earlier that although the overall structure of SDR enzymes remains similar, their ability

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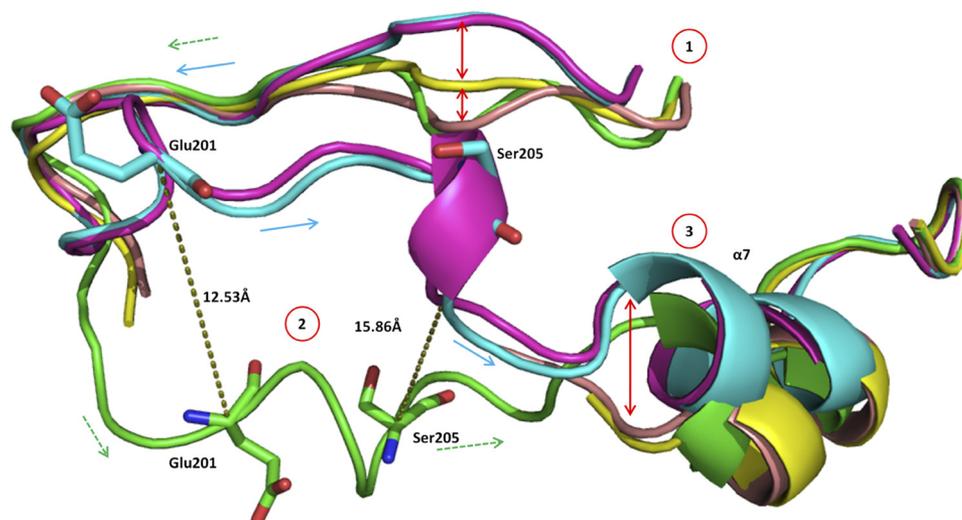


FIGURE 5. The aligned schematic view of the substrate binding loop of BphB_{B-356} in apo (peach), binary (yellow), ternary (with 23-DB (cyan) and 44-DB (magenta)), and intermediate states (green). Numerals in circles show the conformational changes in different structural forms at three positions. Dotted lines in dark olive green show the distance between Glu²⁰¹ and Ser²⁰⁵ in the ternary and intermediate forms. Smooth arrows (cyan) and dotted arrows (green colors) differentiate the direction of movement of ternary and intermediate states, respectively.

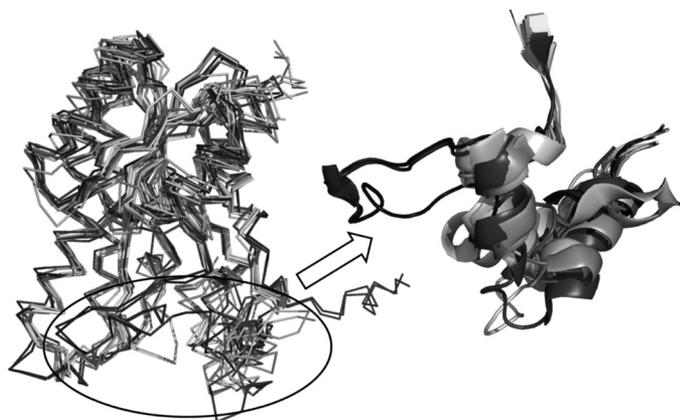


FIGURE 6. The alignment of the top 5 PDB hits obtained from Blastp and top 5 hits obtained by Dali. The circle highlights the difference in the substrate binding loop between BphB_{B-356} (in black) and other PDB structures (light color). PDB code 1BDB and BphB_{B-356} are shown in black color, and other structures are shown in light colors. These PDB codes are: 2HQ1, 3OEC, 3PKO, 2UVD, 2PNF, 2WSB, 1YDE, 2WDZ, 1HDC, and 1NFR.

to metabolize a wide range of substrate is attributed to their flexible substrate binding loop (39, 40). It could be speculated that the differences in the substrate binding loops of BphB_{B-356} and BphB_{LB400} could account for their differing specificity toward various biphenyl derivatives as substrates.

Reactivity of BphB_{B-356} with Polychlorinated Biphenyls—The reactivity of BphB_{B-356} with three different PCBs was examined. As expected from previous work with purified enzymes (12) and shown in Fig. 7, an isopropyl β-D-1-thiogalactopyranoside-induced resting cell suspension of *E. coli* harboring pDB31[LB400**bphFG**] plus pQE31[B-356 **bphAE**] produces one metabolite from each 3,3'-dichlorobiphenyl and 2,4,4'-trichlorobiphenyl, which have previously been identified as the *cis*-5,6-dihydro-5,6-dihydroxy-3,3'-dichlorobiphenyl and *cis*-2',3'-dihydro-2',3'-dihydroxy-2,4,4'-trichlorobiphenyl, respectively. Neither of these dihydrodiol metabolites are found in the co-culture of *E. coli* harboring pDB31-[LB400**bphFG**] plus pQE31[B-356 **bphAE**] plus *E. coli* harbor-

ing pQE31[B-356 **bphB**]. Instead, GC-MS analysis of the metabolites produced by the co-culture showed that 3,3'-dichlorobiphenyl was converted to a dihydroxy-dichlorobiphenyl metabolite and 2,4,4'-trichlorobiphenyl was converted to a dihydroxy-trichlorobiphenyl. In the case of 2,6-dichlorobiphenyl, BPDO_{B-356} produces principally the 2',3'-dihydro-2',3'-dihydroxy-2,6-dichlorobiphenyl and small amounts of 3',4'-dihydro-3',4'-dihydroxy-2,6-dichlorobiphenyl. Based on previous observations, to obtain larger amounts of both metabolites, we have used *E. coli* harboring pDB31[LB400**bphFG**] plus pQE31[*p4 bphAE*]. In this case the co-culture comprised of *E. coli* harboring pDB31[LB400**bphFG**] plus pQE31[*p4bphAE*] plus *E. coli* harboring pQE31[B-356 **bphB**] produced two dihydroxy-2,6-dichlorobiphenyl, which shows that both the 2,3- and 3,4-dihydrodiol metabolites produced from 2,6-dichlorobiphenyl were metabolized by BphB_{B-356}. These results are in agreement with previous data showing that both BphB_{B-356} and BphB_{LB400} are able to catalyze the hydroxylation of 3,4-dihydro-3,4-dihydroxy-2,2',5,5'-tetrachlorobiphenyl.

Docking of Substrates at the Active Site—To get the structural details of the substrate/product binding on the basis of the above biochemical data, the dihydrodiols of 2,4,4'-trichlorobiphenyl, 2,6-dichlorobiphenyl, and 3,3'-dichlorobiphenyl were docked at the active site of BphB_{B-356}. The product of BPDO, *cis*-(2*R*,3*S*)-dihydroxy-1-phenyl-cyclohexa-4,6-diene (BPDD), which is the substrate for BphB was first docked at the putative active site composed of the strictly conserved residues Ser¹⁴², Tyr¹⁵⁵, and Lys¹⁵⁹ as well as the nicotinamide end of NAD⁺. A combination of Glide scores and visual inspection of the docked conformation were taken into account to select the most probable binding mode. The substrate docked in a deep hydrophobic cleft of the cavity. The nonpolar part of the substrate forms hydrophobic interactions with residues Val²⁰⁷, Met²¹², Val²¹⁶, and Met²⁵⁵. The polar portion of the substrate places itself between the nicotinamide ring and the indole ring of Trp⁹⁰. Of the two hydroxyl groups, 2-OH of BPDD forms an H-bond with

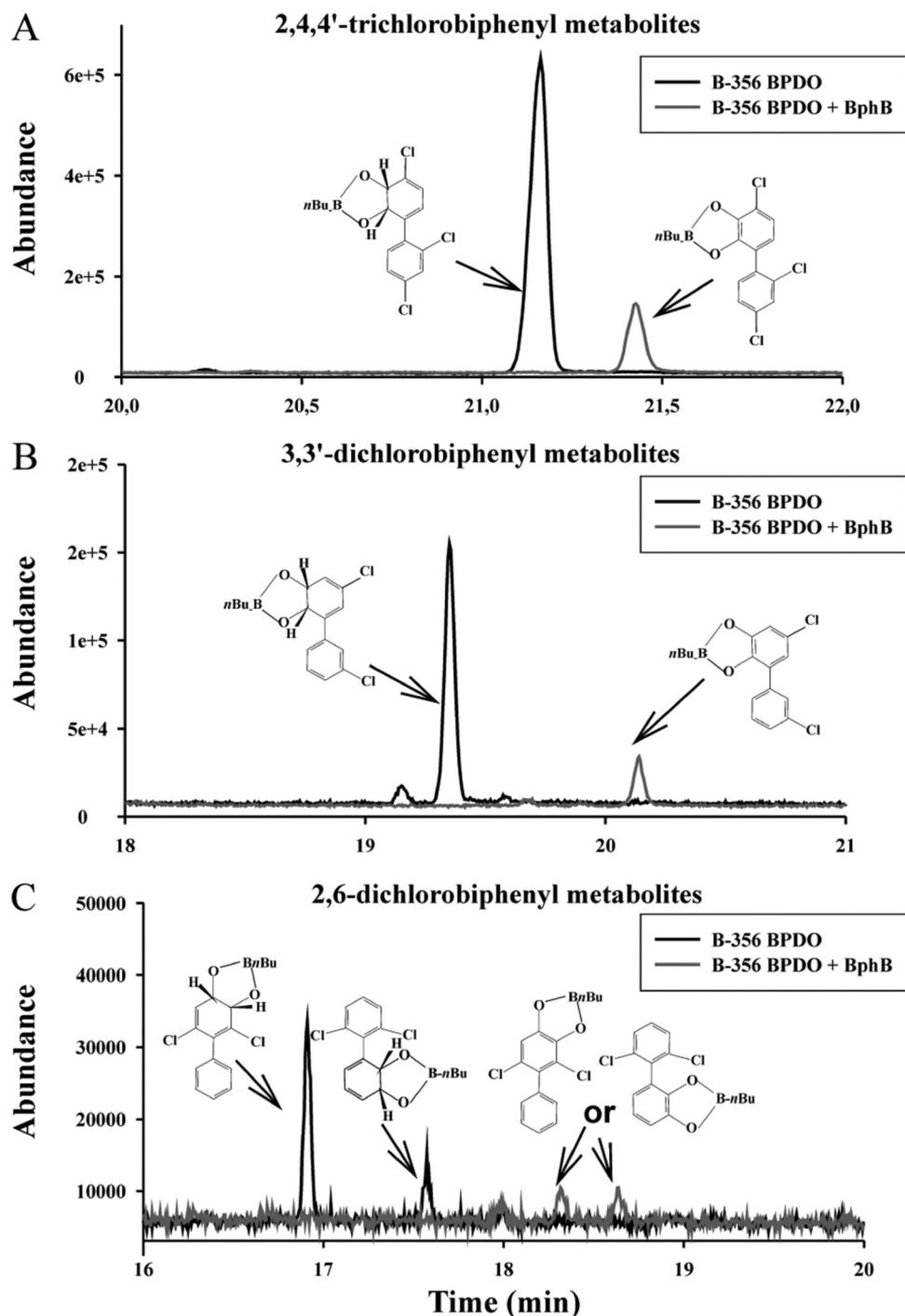


FIGURE 7. GC-MS spectra of butylboronate-derived metabolites produced from (A) 2,4,4'-trichlorobiphenyl, (B) 3,3'-dichlorobiphenyl, and (C) 2,6-dichlorobiphenyl by *E. coli* clones producing BPDO_{B-356} (black line) or BPDO_{B-356} plus BphB_{B-356} (gray line). The metabolites were extracted with ethyl acetate, derivatized with butylboronate, and injected in the GC-MS chromatograph as described under "Experimental Procedures." The figure shows the dihydrodiols produced from these congeners by *E. coli* cells producing BPDO_{B-356} only are not found in cultures producing BPDO_{B-356} plus BphB_{B-356}. Instead, the metabolites detected in those cultures (gray lines) exhibit GC-MS features showing they are catechol metabolites. This shows BphB_{B-356} oxidized the dihydrodiol metabolites produced from these three congeners, including the 3,4-dihydroxy metabolite produced from 2,6-dichlorobiphenyl.

Gly¹⁵⁰ (Gly-C = O-2-OH, 3.0 Å) and Asn¹⁴³ (Asn-ND2-2-OH, 2.9 Å). The other hydroxyl group, 3-OH, interacts with the side chains of Ser¹⁴² (OG-3-OH, 3.1 Å) and Tyr¹⁵⁵ (OH-3-OH, 3.1 Å). For this conformation a G-score of -7.7 was obtained (numbering of the hydroxyl groups is done according to Fig. 1). Docking studies of the other substrates were carried out using the same grid prepared for the docking of BPDD, such that the same docking parameters could be used for all other substrates.

The dihydro-dihydroxylated product of 2,4,4'-trichlorobiphenyl, 3,3'-dichlorobiphenyl, and both products of 2,6-dichlorobiphenyl were docked. Table 2 lists the hydroxylated products of these chlorinated biphenyls and the Glide score obtained after docking them at the active site.

All the compounds docked well at the active site, lie at H-bonding distance from the active site residues Ser¹⁴², Tyr¹⁵⁵, and Asn¹⁴³, and also interact with Gly¹⁵⁰ similar to 23-DB in the

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TABLE 2

Listing different substrates docked with their glide score

Compound name/PCB	Product as obtained after metabolism by BphA	Glide score
Biphenyl (prototype)	<i>cis</i> -(2 <i>R</i> ,3 <i>S</i>)-Dihydroxy-1-phenyl-cyclohexa-4,6-diene (BPDD)	-7.7
2,6-Dichlorobiphenyl	2,6-Dichloro-2',3'-dihydroxybiphenyl	-8.1
	2,6-Dichloro-3',4'-dihydro-3',4'-dihydroxybiphenyl	-8.0
3,3'-Dichlorobiphenyl	5,6-Dihydro-5,6-dihydroxy-3,3'-dichlorobiphenyl	-7.6
2,4,4'-Trichlorobiphenyl	2,3-Dihydro-2,3-dihydroxy-2,4,4'-trichlorobiphenyl	-7.8

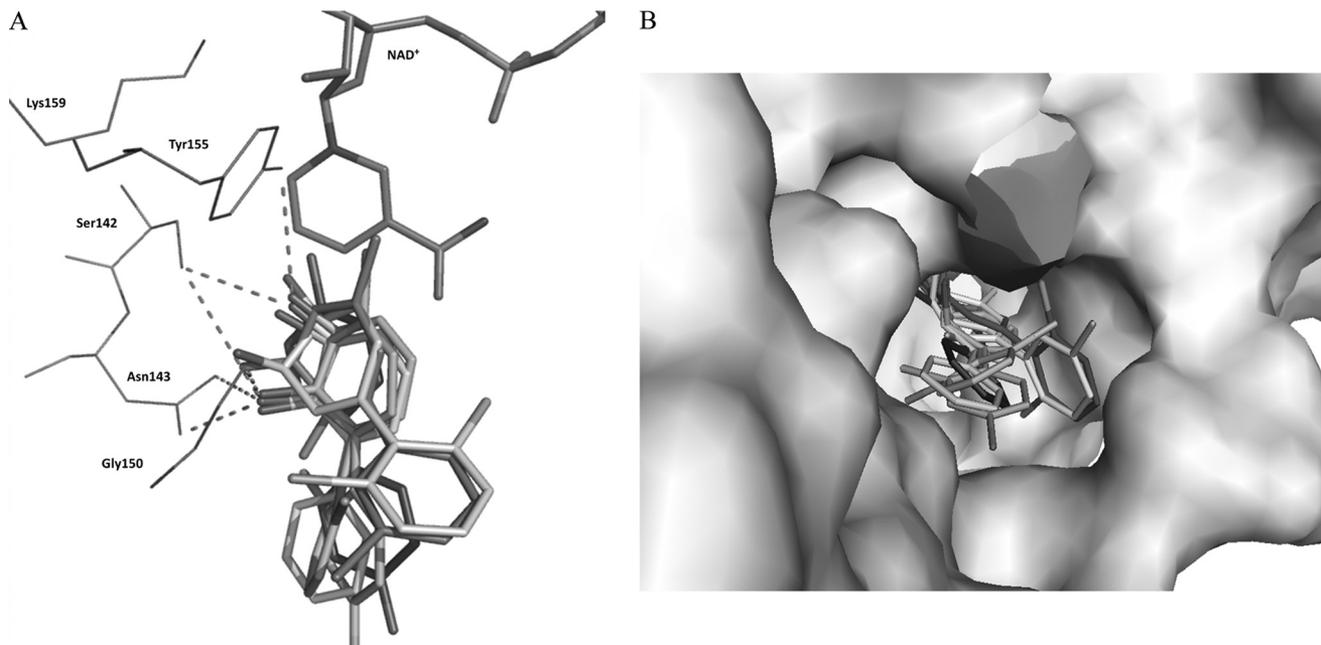


FIGURE 8. A, the superimposed docked structure of *cis*-(2*R*,3*S*)-dihydroxy-1-phenyl-cyclohexa-4,6-diene, 2,3-dihydro-2,3-dihydroxy-2,4,4'-trichlorobiphenyl, 2',3'-dihydro-2',3'-dihydroxy-2,6-dichlorobiphenyl, 3',4'-dihydro-3',4'-dihydroxy-2,6-dichlorobiphenyl, 5,6-dihydro-5,6-dihydroxy-3,3'-dichlorobiphenyl, and product 2,3-dihydroxybiphenyl with NAD⁺ in stick model. B, the aligned substrates in surface representation of protein highlighting the cavity large enough to accommodate different substrates with different orientations of their distal rings.

ternary structure as shown in Fig. 8A. The distal ring of the docked compounds makes hydrophobic interaction with Gly¹⁸⁶, Ile²⁰⁴, Val²⁰⁷, Leu²⁰⁹, Met²¹², Val²¹⁶, and Met²⁵⁵. Asn¹⁴³ is conserved in all related *cis*-dihydrodiol dehydrogenases that use biphenyl-, phenyl-, and toluene derivatives as substrates (17). It is noteworthy that in the ternary structure and all of the docked structures of BphB_{B-356}, one of the hydroxyl groups of the substrate or product binds with Asn¹⁴³, suggesting this residue could be responsible for substrate recognition.

DISCUSSION

Proposed Structural Changes during Binding of a Ligand—We have determined the crystal structures in four different states: the apo-enzyme, as a binary complex with the coenzyme NAD⁺, as a ternary complex with NAD⁺ and either product or product analog, and an intermediate state of the substrate binding loop in BphB_{B-356} for the first time.

The comparative crystal structure analysis of the four different forms suggests that major structural changes occur in BphB_{B-356} when the ligand accesses the active site in the presence of coenzyme. The most obvious changes were observed at the substrate binding region and they involved three segments of the protein as shown in Fig. 5.

First, upon substrate binding, residues Gly¹⁸⁶ to Leu¹⁹¹ move toward NAD⁺. Changes taking place on this portion of the pro-

tein result in an increased number of interactions with the NAD⁺ molecule. Compared with the apoenzyme and the binary structure, an interesting shift in the main chain of Asp¹⁹⁰ is seen in the ternary structure. In the ternary complex, Asp¹⁹⁰ is displaced by 4.1 Å from its position in apostructure as shown in Fig. 9. At its new position, it interacts with Arg⁴¹ of helix α 2. This interaction could probably help in holding the substrate binding loop at its new position.

Second, the region from Gly¹⁹⁸ to Val²⁰⁷ becomes organized. Taking into account the intermediate structure of the enzyme, it is proposed that during the substrate/ligand binding this loop moves from an open to a closed conformation as a swing attached at two hinges. In BphB_{B-356}, the hinge residues are Leu¹⁹⁷ and Leu²⁰⁹. During the swing movement of the loop, a rotation of the main chain torsion angle around hinge residues Leu¹⁹⁷ and Leu²⁰⁹ takes place and residues Gly¹⁹⁸ to Pro²⁰⁸ translate to the other side, bringing the substrate binding loop closer to the active site. The loop movement from the intermediate to the ligand bound state requires about a 15 Å displacement as shown in Fig. 5. During this movement, Leu¹⁹⁹ in the loop comes closer to Tyr⁹², which lies between sheet β 4 and helix α 4 as shown in Fig. 10. At this state, Tyr⁹² makes a conformational rotation away from Leu¹⁹⁹ and toward Glu²⁰² to allow the loop displacement and prevent short contacts with Leu¹⁹⁹. After the loop movement, the main chain carboxyl

group of Glu²⁰² at its new position holds Tyr⁹² by its side chain through hydrogen bonding, thus stabilizing the changes.

Third, a displacement in the small helix $\alpha 7$ (Leu²⁰⁹ to Val²¹⁶) was also observed. The difference was clearly visible when the apo-enzyme and binary complexes are compared with the ternary structures as shown in Fig. 5. During the movement of helix $\alpha 7$, the hinge residue Leu²⁰⁷ that causes the swing movement of the substrate binding loop also moves upwards, thus it helps in the overall displacement of the substrate binding loop toward the active site.

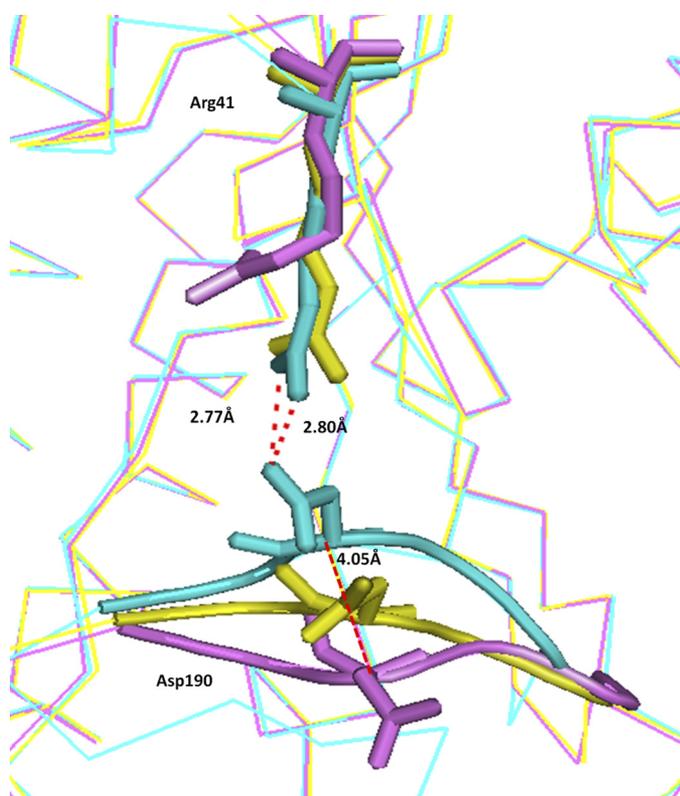


FIGURE 9. The aligned ribbon view of apo, binary, and ternary states of BphB_{B-356} are colored purple, yellow, and cyan, respectively, focusing on Asp¹⁹⁰ and Arg⁴¹. These residues in all the three structures are represented by stick model.

These observations suggest that the ligand causes major structural changes in the protein leading to the formation of a cavity for the active site. These changes could be essential for the entry and release of the substrate and product. Also it is consistent with the ordered ternary complex mechanism as proposed by Andersson *et al.* (34). In the case of BphB_{B-356}, NAD⁺ binds first without any closure of the cavity, followed by the entry of the substrate, whereas the active site is made inaccessible for solvent during hydride transfer. Generally, in the SDR family, conformational changes in the substrate binding loop occur when the substrate binds at the active site (31–34). However, this is not always the case, for example, D-3-hydroxybutyrate dehydrogenase from *Pseudomonas putida*, where coenzyme binding alone is able to induce a conformational change in this loop region that is sufficient to carry out the catalytic activity (41, 42).

In our present study Tyr⁹² and Asp¹⁹⁰ seem to play an important role during substrate binding. These residues should be targeted in future mutagenic studies to better understand their role.

Biochemical Analysis—It was reported by Gomez-Gil *et al.* (12) that BPDO_{B-356} metabolizes 2,6-dichlorobiphenyl, 2,4,4'-trichlorobiphenyl, and 3,3'-dichlorobiphenyl to corresponding dihydrodiol metabolites more efficiently than BPDO_{LB400}. It is therefore interesting to examine the capacity of BphB_{B-356} to metabolize the dihydrodiol metabolites produced from these chlorobiphenyl congeners as they represent congeners with doubly *ortho*-, *meta*-, or *para*-substitution. Furthermore, to our knowledge, the ability of a 2,3-dihydro-2,3-dihydroxybiphenyl dioxygenase to catalyze the dehydrogenation of 2',3'-dihydro-2'-3'-dihydroxy-2,4,4'-trichlorobiphenyl and 2',3'-dihydro-2'-3'-dihydroxy-2,6-dichlorobiphenyl has not yet been determined. The biochemical analysis showed that 3,3'-dichlorobiphenyl and 2,4,4'-trichlorobiphenyl were transformed to dihydroxy-dichlorobiphenyl and dihydroxy-trichlorobiphenyl metabolites, respectively. In the case of 2,6-dichlorobiphenyl, BphB_{B-356} converts both the 2',3'-dihydro-2',3'-dihydroxy-2,6-dichlorobiphenyl and 3',4'-dihydro-3',4'-dihydroxy-2,6-dichlorobiphenyl. These biochemical observation shows that BphB_{B-356} is able to transform a broad range of PCBs, specifically doubly and triply substituted *ortho*-, *meta*-, and *para*-substituted dihydrodiols.

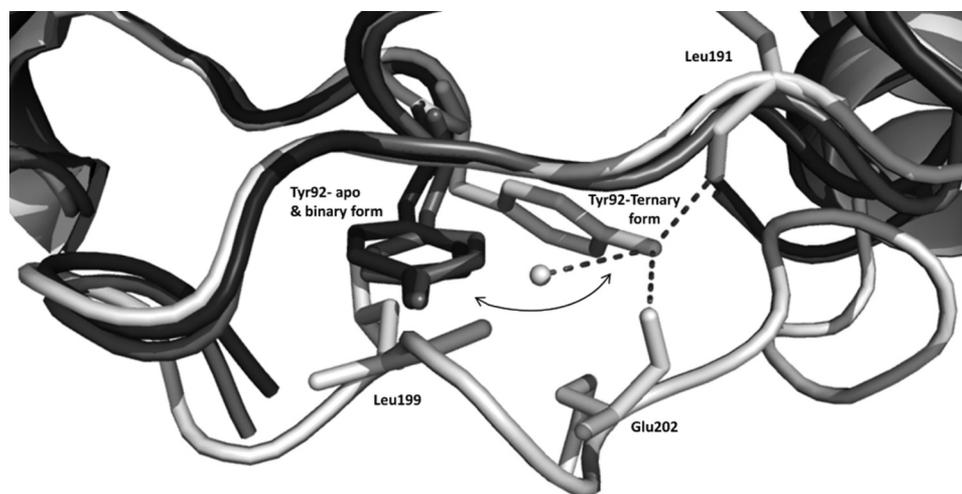


FIGURE 10. The aligned apo and binary forms of BphB_{B-356} are shown in dark colors and the ternary state is shown in white color. The figure shows the displacement of Tyr⁹² from its original positions in the apo and binary states toward the other side in the ternary state to prevent short contact from Leu¹⁹⁹ during the loop movement. Dotted lines show the H-bond that Tyr⁹² (ternary state) forms with surrounding residues.

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Docking Analysis—It was observed from the biochemistry and crystal structure complexes of product and product analogs that this enzyme is able to transform a wide range of ligands. Therefore, molecular docking was carried out to examine the binding of the doubly and triply substituted chlorinated dihydrodiols. Docking analysis of the metabolites produced by BphB_{B-356}, which were observed by the biochemical experiments revealed that all of them may have a similar mode of binding at the active site of the BphB_{B-356}. The superposition of all substrates showed that they were docked at almost the same position as in 23-DB in the ternary structure as shown in Fig. 8A. The differences in binding positions were observed at the nonhydroxylated ring of different compounds, suggesting ligands may access the active site with different orientations. The docked structures of 5,6-dihydro-5,6-dihydroxy-3,3'-dichlorobiphenyl and 3',4'-dihydro-3',4'-dihydroxy-2,6-dichlorobiphenyl were placed in such a way that the distal rings of these compounds occupy a different space than the distal rings of the other substrates.

These variations do not seem to be associated with equivalent variations in the orientation of the hydroxylated ring. Also the surface diagram, Fig. 8B, shows that the cavity size is narrow where the hydroxylated ring binds, whereas there is a much larger volume near the nonhydroxylated ring site suitable for accommodating different compounds by orienting them differently. In combination, the biochemical data and docking experiments suggests that BphB_{B-356} can accommodate both *ortho*-*meta*-hydroxylated as well as *meta*-*para*-hydroxylated substrates.

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