Mode-of-action and evolution of methylenedioxy bridge forming P450s in plant specialized metabolism

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Abstract (+)-Sesamin is a major furofran-class lignan in sesame seeds and harbors characteristic two methylenedioxy bridges (MDB) that are sequentially formed from (+)-pinoresinol via (+)-piperitol by a Sesamum indicum P450, CYP81Q1. However, the molecular basis for this unique catalytic activity of CYP81Q1 has been poorly understood. To elucidate MDB formation, we tested various natural and non-natural metabolites as substrates for CYP81Q1. A synthetic (+)-SC1mr and a naturally occurring (+)-kobusin showed inhibitory effect on the production of (+)-sesamin by CYP81Q1 unlike (+)-epipinoresinol and (+)-pinoresinol, indicating the strict diastereomer and enantiomer selectivity. Homology modeling followed by site-directed mutagenesis of CYP81Q1 showed that an amino acid residue crucial for MDB formation is a unique Ala residue (A308), located in I-helix proximal to the substrate pocket, responsible to the conserved distal-Thr residue. MDB by CYP81Q1 is produced possibly through the formation of a substrate-participated hydrogen-bonding network, since single replacement of the Ala by Thr severely and specifically lowered the MDB forming activity. This hypothesis is supported by a newly identified MDB-forming enzyme CYP81Q38 from Phryma leptostachya harboring an Ala responsible to Ala308 in CYP81Q1. An evolutional perspective of CYP81Q1 is discussed in relation to another MDB-forming CYP719As functionally conserved in Ranunculales.

Key words: P450, lignan, methylenedioxy bridge, lamiales, convergent evolution.

![Diagram of naturally occurring specialized metabolites with MDB structure](image)

**Figure 1.** Naturally occurring specialized metabolites with MDB structure. Stylopine in California poppy (*Eschscholzia californica*), noscapine in opium poppy (*Papaver somniferum*), Berberine in *Coptis japonica*, lycorine in *Lycoris radiata*, piperine in peppet (*Piper nigrum*), pisatin in pea (*Pisum sativum*), pseudobaptigenin in chickpea (*Cicer arietinum*), podophyllotoxin in *Podophyllum hexandrum*, yatein in *Anthriscus sylvestris*, dihydrocubebin in *Piper cubeba*, hinokinin in *Linum corymbulosum*, safrole in *Sassafras officinale*, apiole in parsley (*Petroselinum crispum*), aristolochic acid in *Asarum sieboldii*, phrymarolin I in *Phryma leptostachya*, paulownin in *Paulownia tomentosa*.

Oxidative formation of MDB moiety is catalyzed by two discrete classes of P450 monoxygenases (P450s). Previously we identified CYP81Q1 from *S. indicum* as a piperitol/sesamin synthase (PSS) that catalyzes two sequential MDB formations on (+)-pinoresinol, yielding (+)-sesamin via (+)-piperitol (Figure 2) (Ono et al. 2006). Moreover CYP81Q2, CYP81Q3 and CYP81Q4 were isolated from *S. radiatum*, *S. alatum*, and *S. schinzianum*, respectively, showing that CYP81Q genes are structurally conserved among *Sesamum* genus.

CYP719A enzymes are another class of MDB-forming P450 for isoquinoline alkaloids (Ikezawa et al. 2003, 2007, 2009; Takemura et al. 2013; Winzer et al. 2012). More recently, CYP719A23 and CYP719A24 from *Podophyllum* spp. were shown to catalyze MDB formation from (−)-matairesinol to (−)-pluviatolide (Marques et al. 2013). Important to note, MDB formation is a unique type of oxygenation reaction because of the lack of oxygen derived from P450 in the products. In spite of general occurrence of MDB as a chemical structure observed in an array of specialized metabolites, molecular mechanism of MDB formation by P450 has been poorly understood.

In this report, we characterized MDB formation mechanism, 1) with selected lignans which inhibit MDB
formation on (+)-piperitol by CYP81Q1, and 2) by site-directed mutagenesis upon CYP81Q1 based on the homology models. Through a series of investigations, we propose a possible scheme for MDB formation. We also discuss an evolitional perspective of MDB forming enzymes in lineage-specific specialized metabolisms with identification of a novel PSS member, CYP81Q38 from *Phryma leptostachya*, in comparison to CYP719As found in Ranunculales.

**Materials and methods**

**Chemicals**

Two sesamin metabolites, (1R,2S,5R,6S)-6-(3,4-dihydroxyphenyl)-2-(3,4-methylenedioxyphenyl)-3,7-dioxabicyclo[3,3,0]-octane ((+)-SC1) and (1R,2S,5R,6S)-2,6-bis(3,4-dihydroxyphenyl)-3,7-dioxabicyclo[3,3,0]-Octane ((+)-SC2) (Nakai...
et al. 2003) were prepared by the previously reported procedure (Urata et al. 2008). Piperitol ((+)-SC1m) and its positional isomer, (+)-SC1mr, and (+)-pinoresinol, and its positional isomers, (+)-pinoresinol mr1 (Pmr1) and (+)-pinoresinol mr2 (Pmr2) were obtained by the reaction of (+)-sesamin with diisobutylalminum hydride in toluene at reflux. (+)- and (−)-syringaresinol or (−)- and (−)-pinoresinol were prepared by the separation of synthesized dl-syringaresinol or dl-pinoresinol (Vermes et al. 1991) in the high performance liquid chromatography (HPLC) using the chiral column. (+)-Pinoresinol monomethyl ether was prepared by hydrolysis of (+)-pinoresinol monomethyl ether monoglucoside isolated from leaves of Forsythia. (+)-Kobusin was synthesized by enzymatic reaction of recombinant CYP81Q1 proteins with (+)-pinoresinol monomethyl ether prepared from Forsythia plants. (+)-Lariciresinol and (−)-matairesinol were purchased from Arbo Nova (Finland). The other compounds were described in previously works (Noguchi et al. 2008; Ono et al. 2006).

Preparation of recombinant CYP81Q1 proteins in Baculovirus/Sf-9 system

Gateway system was used to construct a plasmid for heterologous expression in Sf9/Baculovirus system kit (Life technology, CA). The cDNA of CYP81Q1 was amplified by PCR using the following two primers: 5′-GGG GAC AAG TTT GTA CAA AAA AAG ACC AGG CTA TG GAA GCT GAA ATG CTA TAT TCA T3′ (an attB1 site is underlined) and 5′-GGG GAC CAC TTT GTA CAA GAA AGC TGG GTT CA AAG GTT GGA AAC CTC AGC AAG AA-3′ (an attB2 site is underlined), and then subcloned into pENTR-D-TOPO (entry vector). The subcloned fragment was sequenced to confirm the absence of PCR errors. The cDNA subcloned in the entry vector was transferred in attR-containing destination vector, pDEST8 by LR recombination reaction.

In vitro mutagenesis of the CYP81Q1 gene was performed by recombinant PCR (Higuchi 1990) to obtain the following site-directed mutants: CYP81Q1-A308S and CYP81Q1-A308T, which have amino acid substitutions at Ala-308 replaced with serine and threonine, respectively. The primers used for PCR were as follows: CYP81Q1_full_Fw (5′-ATG GAA GCT GAA ATG CTA TAT TCA GCT CT) and CYP81Q1_full_Rv (5′-TCA AAC GTT GGA AAC CTC AGC AAG AA) for amplification of full length of CYP81Q1 and its mutants. CYP81Q1_A308S_Fw (5′-ATT GCG GAG CAT TCA TGG GTA ACT) and CYP81Q1_A308S_Rv (5′-AGT TAC AAG CGT GAA ATCC GTT CCG ACA AT) for A308S mutagenesis. CYP81Q1_A308T_Fw (5′-ATT GCG GAG CAT GAA ATCC GTT GAA ACT) and CYP81Q1_A308T_Rv (5′-AGT TAC AAG CGT GAA ATCC GTT CCG ACA AT) for A308T mutagenesis. The sequence of the Rv primer for PCR-mutagenesis was complementary to that of the corresponding Fw primer. The PCR was performed with the plasmid pENTR-D-TOPO containing cDNA of CYP81Q1 (see above) as a template using full_Fw primer and A308S-Rv, A308S-Fw and full_Rv primer, full_Fw primer and A308T-Rv, or A308T-Fw and full_Rv primer as primers, yielding A308S-a, A308S-b, A308T-a or A308T-b, respectively. The second PCR was performed with mixture of A308S-a and A308S-b or A308T-a and A308T-b as a template using full_Fw primer and full_Rv primer as primers. The amplified fragments were subcloned in pENTR-D-TOPO as described above and sequenced to confirm the absence of PCR errors. The resultant CYP81Q1-A308S and CYP81Q1-A308T in entry vector were transferred to pDEST8 as described above. Three kinds of destination vectors (CYP81Q1-WT, A308S, A308T) were transformed to E. coli (strain DH10Bac, Life technology, CA). Preparation of the recombinant CYP81Q1-baculovirus and transfection to Sf9 cells (Spodoptera frugiperda 9) were performed according to the instructions of the manufacturer (Life technology, CA). Heterologous expression of CYP81Q proteins in Sf9 cells and spectrophotometric analysis were performed as described (Saito et al. 2004).

Enzyme assays and kinetics

Enzyme assay procedure and HPLC analysis of reaction mixtures were described in our previous works (Noguchi et al. 2008; Ono et al. 2006). For relative inhibition assay, 100 μM of (+)-piperitol and tested lignans were used as substrate and inhibitor, respectively. Chemical-structure confirmation of the product in the reaction of CYP81Q1 with (+)-pinoresinol monomethyl ether was performed by a high-performance liquid chromatography (20A series of Prominance) coupled to ion trap time-of-flight mass spectrometry (LC-IT-TOF-MS, Shimadzu, Japan) with a Develosil C30-UG-3 column (2.0 mm i.d.×150 mm, Nomura Chemical, Japan). In LC conditions, Samples were eluted through the column with a gradient of water and acetonitrile containing 0.1% formic acid (0 min, 65:35; 20 min, 0:100; 27 min, 0:100; 27.1 min, 65:35; 38 min, 65:35). IT-TOF-MS measurements with an electrospray ionization (ESI) source in positive mode were carried out under the following conditions. The voltages of interface, skimmer, and detector were 4.5 kV, 6.5 kV, and 1.68 kV. The scan range was m/z 300–1000. The CDL and heat block temperatures were 200°C. Liquid nitrogen (purity >99.999%) was used as the source of nebulizer gas and drying gas. Enzyme-reaction products were detected by the correspondent ions such as a protonated ion and a dehydrated ion. The initial velocity assays for CYP81Q1 and its mutants were carried out under steady-state conditions using the standard assay system (see above) with varying concentrations of substrates. The kinetic parameters and their standard errors were determined at nine substrate concentrations (2–100 μM) by fitting the initial velocity data to the Michaelis–Menten equation by means of a non-linear regression analysis (Leatherbarrow 1990). Inhibition by (+)-SC1mr was kinetically analyzed under steady-state conditions using the standard assay system (see above) with 20 μM of (+)-SC1mr. The data were fitted to the competitive inhibition equation (Eq. 1) using least-squares analysis.
\[
\frac{1}{v} = \frac{1}{V_{\text{max}}} + \frac{K_m [I + ([2]/K_i)]}{(V_{\text{max}}[S])} \quad \text{(Eq. 1)}
\]
where \(v\) and \(V_{\text{max}}\) are the initial velocity and the maximum velocity, respectively, \(K_m\) and \(K_i\) are the Michaelis constant and inhibition constant for \((+)-\text{SC1mr}\), respectively, and \([S]\) and \([I]\) denote concentrations of \((+)-\text{piperitol}\) and \((+)-\text{SC1mr}\), respectively.

**Homology modeling**

The three-dimensional structural model of CYP81Q1 was constructed with homology modeling module within Insight II (Accelrys, CA) using the crystal structure data of P450 BM3 (CYP102A1, pdb code: 1bu7) as a template. The crystal structure had two water molecules near the active site of the monooxygenation reaction, thus two water molecules were placed in the same position of the initially constructed model of CYP81Q1. The water molecule 2 (w2) has been known as one of important sites for the monooxygenation reaction. The water molecule 1 (w1) placed near both w2 and the oxygen atom in the iron(IV)oxo species. We predicted that w1 was important species for MDB formation because of replacement of conserved Thr residue to Ala in CYP81Q1. The substrate, \((+)-\text{pinoresinol},\) was then docked into the substrate-binding space near the catalytic center. The constructed \((+)-\text{pinoresinol-bound CYP81Q1}\) was energy-minimized with molecular mechanics, followed by optimization with molecular dynamics using Discover 3 (Accelrys, CA). To construct the model structures of \((+)-\text{molecular dynamics using Discover 3 (Accelrys, CA). To with molecular mechanics, followed by optimization with \((+)-\text{SC1mr}\), \((+)-\text{SC1mr}\), and CYP81Q1-A308T.

**Molecular cloning and RT-PCR of CYP81Q38**

*Phryma leptostachya* naturally grown in Rokko Mountain in Kobe, Hyogo prefecture was collected in summer 2007. Total RNAs were extracted from each organ (roots, leaves, stem, and flowers) with RNeasy Plant Mini kit (Qiagen, CA), and then reverse-transcribed into cDNA with SuperScript III (Invitrogen, CA) as described previously (Noguchi et al. 2009). A cDNA fragment responsible for CYP81Q38 was amplified by RT-PCR with the CYP81 primers of CYP81Q-Fw (5′-ATG GAA GCT GAA ATG CTG CTA TAT TCA) and CYP81Q-Rv (5′-TCA AAC GTT GGA AAC CTG ACG AAG AA) on cDNAs prepared from roots. The PCR with ExTag DNA polymerase (Takara Bio, Japan) was run at 94°C for 3 min followed by 28 cycles at 94°C for 1 min, at 54°C for 1 min and at 72°C for 2 min. *Phryma* 5.8S rRNA (accession DQ533822) was amplified with the primers of Pl-rRNA-Fw (5′-AGG ATC ATT GTC GAA ACC TGC AA-3′) and Pl-rRNA-Rv (5′-GGG CGT CGT TGG ACA CGC GTT-3′).

**Lignan biotransformation by yeast cells expressing CYP81Q38**

cDNA of CYP81Q38 was cloned in the yeast expression vector, pYE22m as described previously (Ono et al. 2006). The resulting vector for CYP81Q38 expression, termed pYE22m-PlCYP81Q38, was used to transform a yeast strain, INVsc (Invitrogen, CA) by conventional method as described previously (Ono et al. 2006). 100 μl of saturated liquid cultures of yeast cells harboring pYE22m-PlCYP81Q38 were transferred to 3 ml of fresh medium supplemented with 100 μM of \((+)-\text{piperitol.}\) After 48 h of culture at 30°C, the cultures were mixed with equal volume of methanol (final 50% methanol), sonicated together with the medium and centrifuged at 21,000 × g for 10 min at 4°C. The supernatant was filtered and subjected to HPLC analysis using a Develosil C30-UG-5 column (Nomura Chemical, Japan) with gradient elution program starting from 15% acetonitrile with 0.1% trifluoroacetic acid (TFA) to 80% acetonitrile with 0.1% TFA in 8 min, followed by isocratic elution for 7 min at a flow rate of 1 ml/min. The peaks were detected using 2996 PDA Detector (Waters, MA) at 283 nm. The UV spectrum and the retention time (12 min) of the product peak were identical to those of \((+)-\text{sesamin standard.}\) Yeast cells transformed with the empty vector pYE22m was used as a negative control. The identity of the product was further confirmed by co-chromatography analysis by mixing \((+)-\text{sesamin standard with the extracts.}\)

**Phylogenetic analysis**

The amino acid sequences of CYP81Qs and CYP719As were aligned by taking consideration of codon position using ClustalW bundled in SEAVIEW 4.2.4 (Gouy et al. 2010; Thompson et al. 1994). All positions containing gaps and missing data were eliminated from the further analysis. The unrooted phylogenetic trees of them were reconstructed by neighbor-joining methods from translated amino acid sequences. The alignment used to generate the tree shown in Figure 5C is available as Supplemental Data Set. The neighbor-joining tree was reconstructed by SEAVIEW with the matrix of the evolutionary distances calculated by Poisson correction for the multiple substitutions. The reliability of reconstructed tree was evaluated by bootstrap test with 1000 replicates.

**Results**

**Heterologous expression of CYP81Q1**

After SDS-PAGE of microsomal preparation from Sf9 cells infected with baculovirus expressing CYP81Q1, whole proteins were stained with CBB. A major band was detected around 50 kDa, which is consistent with the estimated molecular weight of CYP81Q1. Moreover reduced CO-difference spectra of the microsomal preparation clearly showed a sharp absorbance at 450 nm (Figure 3A). Enzyme activity assays of the microsome fraction expressing CYP81Q1 using \((+)-\text{pinoresinol and NADPH as substrate and co-substrate, respectively,}
Characterization of Methyleneoxy bridge forming P450s

resulted in the production of (+)-sesamin and (+)-piperitol (Figure 3B). These data clearly indicate that recombinant CYP81Q1 protein expressed in the baculovirus/Sf-9 cells system is functionally active.

Substrate specificity of CYP81Q1

We previously showed that CYP81Q1 specifically utilized (+)-piroresinol and (+)-piperitol as substrates but did not accept (+)-phyllicigenin, (+)-epipinoresinol, (+)-piroresinol 4'-O-glucoside or (+)-sesamolinol (Ono et al. 2006). To further investigate substrate specificity of CYP81Q1 in detail, seven naturally occurring lignans, (+)-piroresinol monomethylether, (+)-kobusin, (+)-syringaresinol, (-)-syringaresinol, (-)-lariciresinol, (-)-piroresinol, (-)-matairesinol and an artificial synthesized lignan, (+)-SC1mr were prepared and tested. In the reaction with (+)-piroresinol monomethylether, CYP81Q1 produced a new product at retention time 14.9 min (Figure 3C), which was determined to be (+)-kobusin with \([M+H]^+ = 371\) and \([M+H−H_2O]^+ = 353\) by LC-IT-TOF-MS. In contrast, none of the other lignans were served as substrates for CYP81Q1 in the presence of NADPH, confirming its specific substrate recognition for the same configuration to (+)-sesamin.

To gain insights of molecular mechanism in substrate recognition by CYP81Q1, some of these lignans were further tested as inhibitors for MDB formation from (+)-piperitol to (+)-sesamin by CYP81Q1 that was set as 100% activity in the absence of inhibitory chemicals (Figure 2). Neither (+)-epipinoresinol nor (-)-piroresinol showed apparent inhibitory effects on MDB formation with the relative activity of 97% and 94%, respectively, whereas (+)-kobusin and (+)-SC1mr showed significant inhibitory effects (relative activity of 69% and 56%, respectively) (Table 1). The kinetic parameter of (+)-SC1mr with the highest inhibition (relative activity of 56%) was determined to be \(K_i = 19.8 \mu M\). These results suggest that (+)-kobusin and (+)-SC1mr are accessible to the substrate pocket of CYP81Q1, by which MDB formation for the bona fide substrate, (+)-piperitol is inhibited (block out), whereas (+)-epipinoresinol and (-)-piroresinol are not structurally accessible to the substrate pocket of CYP81Q1 due to their differential configuration.

To confirm this hypothesis, two additional synthetic lignans, (+)-piroresinol-mr1 ((+)-Prmr1) and (+)-piroresinol-mr2 ((+)-Prmr2) were further tested for the inhibition assay. As expected, (+)-Prmr1 and (+)-Prmr2 showed relative high inhibition activity (relative activity of 69% and 79%, respectively) (Table 1). Collectively these results show that CYP81Q1 strictly discriminates substrates; (+) epi-mer and (-)-enantiotmer. In addition, the results using (+)-Prmr1 and (+)-Prmr2 also demonstrate that CYP81Q1 discriminates the position of the two functional groups

Table 1. Relative Inhibitory activity for MDB formation by CYP81Q1. MDB formation activity for (+)-piperitol in the absence of other lignans was set as 100%.

<table>
<thead>
<tr>
<th>CYP81Q1(WT_A308)</th>
<th>Relative activity (%)</th>
<th>(K_i (\mu M))</th>
</tr>
</thead>
<tbody>
<tr>
<td>(+)-Piperitol</td>
<td>100</td>
<td></td>
</tr>
<tr>
<td>(+)-Epipinoresinol</td>
<td>97</td>
<td></td>
</tr>
<tr>
<td>(-)-Pirosinol</td>
<td>94</td>
<td></td>
</tr>
<tr>
<td>(+)-Pirosinol mr2(Pmr2)</td>
<td>79</td>
<td></td>
</tr>
<tr>
<td>(+)-Pirosinol mr1(Pmr1)</td>
<td>69</td>
<td></td>
</tr>
<tr>
<td>(+)-Kobusin</td>
<td>69</td>
<td></td>
</tr>
<tr>
<td>(+)-SC1mr</td>
<td>56</td>
<td>19.8±3.5</td>
</tr>
</tbody>
</table>
(a hydroxy group and a methoxy group) on the aromatic ring of substrates, further establishing the substrate selectivity of CYP81Q1.

**Homology modeling of CYP81Q1**

As previously indicated, MDB forming P450s do not possess conserved Thr residue in the distal I helix that is considered to be crucial for O–O bond scission in the P450 reaction cycle (Imai et al. 1989; Ono et al. 2006). However, all CYP81Qs and CYP719As that have been identified so far have Ala and Ser residues, respectively, at the corresponding position in the distal I helix, suggesting that the Thr residue conserved among other P450s is not indispensable for MDB formation (Figure 4). In turn, Ala or Ser residue observed in MDB forming P450s are likely to be involved in the catalyses for MDB formation.

To gain structural insights into MDB formation by P450s, homology models of CYP81Q1 were constructed using the crystal structure data of P450 BM3 as a template. The putative (+)-pinoresinol-bound CYP81Q1 (WT) complex structure elucidated by homology modeling indicated that two water molecules (w1 and w2) appear to form hydrogen bonding networks to 4-hydroxy group of (+)-pinoresinol (Figure 4). The w2 is able to interact with the two carbonyl groups of the main chain of Ile303 and Ala304. These data suggest that Ala304 together with Ile303 is indispensable for the reaction.

On the other hand, the significance of Ser308 in the catalysis was examined by constructing a homology model of A308S in which Ala308 was replaced with Ser residue. The result demonstrated that Ser residue was located close to the w2 molecule but the position of w2 was analogous to that in WT. In contrast, in the model of A308T in which Ala308 was replaced with Thr residue, Thr was predicted to drastically hamper the position of the w2. These homology models support the notion that amino acid residue on the position responsible to conserved distal-Thr residue is involved in MDB formation.

**Site-directed mutagenesis of CYP81Q1**

To assess significance of the Ala308 residue for MDB formation biochemically, the two amino acid substitutions A308S and A308T were introduced into
CYP81Q1 (WT), and heterologously expressed in Sf9 cells. CBB staining and CO-difference spectra of the mutant proteins confirmed that these CYP81Q1 mutant proteins are functionally expressed (Figure 3). Using these recombinant proteins, kinetic parameters were determined. As shown in Table 2, CYP81Q1-A308S exhibited comparable substrate specificity (K_m) to those of WT, and its catalytic activity (k_cat) was moderately decreased to approximately one-third of WT.

In sharp contrast, both K_m and k_cat of CYP81Q1-A308T were significantly altered. Compared to those of WT, the K_m was increased to 6-fold and the k_cat was decreased to approximately 52-fold (Table 2). These results demonstrate that a single amino acid substitution of Ala residue by Thr residue severely causes catalytic impairment in MDB formation, unlike another substitution by Ser residue, thus support the notion that amino acid residue responsible for the conserved distal-Thr residue is involved in MDB formation. These data showed that amino acid residue on the position responsible to conserved distal-Thr residue is crucial for MDB formation.

Table 2. Kinetic analysis of CYP81Q1 and the mutant proteins.

<table>
<thead>
<tr>
<th>CYP81Q1</th>
<th>K_m (μM)</th>
<th>k_cat (S⁻¹)</th>
<th>k_cat/K_m (S⁻¹/μM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>WT_A308</td>
<td>1.98±0.30</td>
<td>0.173±0.004</td>
<td>0.087</td>
</tr>
<tr>
<td>A308S</td>
<td>1.96±0.58</td>
<td>0.051±0.002</td>
<td>0.026</td>
</tr>
<tr>
<td>A308T</td>
<td>13.33±3.90</td>
<td>0.0033±0.0003</td>
<td>0.00025</td>
</tr>
</tbody>
</table>

Identification of Phryma leptostachya CYP81Q38

Phryma leptostachya (Phrymaceae) is a lignan-producing plant, which belongs to Lamiales, and is a close relative to Sesamum plants (Pedaliaceae) (Figure 5A). Since P. leptostachya is known to accumulate phrymarolin I and other characteristic furanocoumarins with MDB in the roots (Figure 1) (Taniguchi and Oshima 1972), enzyme genes involved in the formation of MDB of

Figure 5. Characterization of Phryma leptostachya CYP81Q38. Inflorescence (A) and roots (B) of P. leptostachya. Size bar in photos indicates 1 cm. (C) A phylogenetic tree of MDB-forming P450s. An unrooted phylogenetic tree was constructed as described in Materials and methods. The alignment used for this analysis is available as Supplemental Data Set. The percentages of replicate trees, in which the associated taxa clustered together in the bootstrap test (1000 replicates), are shown. Bar=0.1 amino acid substitutions per site. Si, Sr, Ss, Sa, Ps, Ec, Cj, Pp, and Ph indicates Sesamum indicum, Sesamum radiatum, Sesamum schinzianum, Sesamum alatum, Phryma leptostachya, Papaver somniferum, Eschscholzia californica, Coptis japonica, Podophyllum peltatum, and P. hexandrum, respectively. (D) Gene expression analysis of CYP81Q38 in separated organs by RT-PCR. L: leaf, S: stem, F: flower, R: root (E) CYP81Q38 catalyzes the conversion of piperitol to sesamin. 3 ml cultures of yeast cells expressing CYP81Q38 were supplemented with 100μM (+)-piperitol 24h after the subculturing, and were further cultured for 48h at 30°C before extracted and subjected to HPLC analysis. pYE22m, a yeast cell line harboring an empty vector pYE22m. CYP81Q38, a yeast cell line that constitutively expresses CYP81Q38 protein. Shown is the representative data of three independent experiments.
the lignans is also expected to be expressed in the roots (Figure 5B). Based on the chemotaxonomic relationship that several lineages in Lamiales produce lignans with MDB, CYP81Q1 homologs were explored as candidates for MDB forming enzyme in *P. leptostachya*. A cDNA fragment with high structural similarity to CYP81Q1 was amplified by RT-PCR using cDNA prepared from the root as a template, and then designated as CYP81Q38 (79% amino acid sequence identity with CYP81Q1; Accession number: AB923911) by P450 nomenclature (Nelson. 2009) (Figure 5C). CYP81Q38 also have an Ala residue (Ala307), corresponding to Ala308 in CYP81Q1, at the position of distal-Thr residue (Figure 4). Semi-quantitative RT-PCR confirmed that CYP81Q38s is highly expressed in root (Figure 5D).

In order to test whether CYP81Q38 catalyzes the formation of MDB on lignans, yeast suspension cultures expressing CYP81Q38 by a constitutive GAPDH promoter were fed with possible lignans. After 48 h of incubation, no additional peaks of lignans other than tested substrates were observed in the yeast cultures harboring a control vector. In contrast, production of (+)-sesamin was detected in the CYP81Q38-expressing cultures when fed with (+)-piperitol (Figure 5E). Similarly, (+)-sesamin and (+)-piperitol were detected in the yeast when fed with (+)-pinoresinol (data not shown). However, CYP81Q38 did not produce any detectable products in the reaction with (+)-sesaminol, (+)-epipinoresinol, (-)-pinoresinol or (+)-SC1mr (data not shown).

**Discussion**

**A possible mode-of-action of MDB formation catalyzed by CYP81Q1**

CYP81Q1 is a *Sesamum indicum* PSS which specifically catalyzes (+)-sesamin formation form (+)-pinoresinol via (+)-piperitol by forming two MDBs (Figures 2, 3B). Investigation of various natural and non-natural compounds as possible substrates for MDB formation by CYP81Q1 revealed that CYP81Q1 utilizes only (+)-pinoresinol monomethylether, over all other compounds tested in the study. Formation of a MDB on (+)-pinoresinol monomethylether by CYP81Q1 resulted in (+)-kobusin (Figure 3C), and CYP81Q1 was inactive for (-)-pinoresinol, are clearly consistent with the previous report that (+)-kobusin is found in sesame seeds, and only (+)-pinoresinol is selectively metabolized into (+)-piperitol and (+)-sesamin in the presence of (±)-pinoresinols in vivo (Kato et al. 1997). Synthetic (+)-Sc1mr, (+)-Prmr1 and (+)-Prmr2 showed inhibitory effect on MDB formation, indicating that CYP81Q1 strictly discriminates configuration and position of functional groups on the aromatic ring of the substrates. This selectivity might be responsible for optical purity of (+)-sesamin and its related lignans with MDB such as (+)-sesaminol in sesame seeds. These results strongly support that CYP81Q1 is the enzyme responsible for (+)-sesamin biosynthesis in vivo.

The homology modeling approach suggests that the lack of a distal-Thr generally conserved in P450s is crucial for MDB forming P450s. The negligible activity of CYP81Q1-A308T mutant and the comparable activity of CYP81Q1-A308S mutant point out an important notion that hydrogen-bonding network is involved in their catalysis. Severe defect observed in CYP81Q1-A308T mutant demonstrated that the ordered hydrogen-bonding network in their homology models, leading to 4-hydroxy group of the substrate, is catalytically important for MDB formation. The Thr residue probably drastically affects the position of the w2, due to a characteristic methyl group of the Thr side chain. The results that hydroxyl group and methoxy group on aromatic ring of the substrates were not interchangeable also support this notion.

On the other hand, Ser residue in the A308S mutant model does not seem to affect this hydrogen-bonding network, which is in good accordance with the result that A308 mutant exhibited the catalytic activity at the level equivalent to that of wild type CYP81Q1 (Table 2). These results substantiate the fact that distal-Thr commonly conserved among P450s is replaced with Ser in another class of MDB forming enzymes, CYP719As. Collectively, our results suggest that A308 is not a catalytic residue of CYP81Q1 but rather facilitates MDB formation by providing space for putative ‘catalytic’ water molecules. Thus, it is plausible to conclude that the establishment of substrate-participated hydrogen-bonding network triggers an intramolecular nucleophile cyclization to elaborate a MDB structure (Figure 6), and moreover, the substitution of amino acids proximal to the distal-Thr residue to Ala (probably also to Ser) is crucial in evolving MDB forming activity. It should be noted that P450cam T252A and T252S mutants produce H2O2, compared to the wild type (T252)(Imai et al. 1989). Thus, the possibility that reactive oxygen species (ROS) might be mediated in catalysis for MDB formation is not excluded.

**Molecular evolution of MDB forming P450s**

A novel cDNA encoding a MDB-forming P450, CYP81Q38, was isolated from root of *P. leptostachya* based on the sequence homology to *S. indicum* PSS, CYP81Q1. Its significant expression in roots as well as its MDB-forming activity on (+)-pinoresinol and (+)-piperitol suggests that (+)-phymarolin I and its-related lignans with MDB are biosynthesized via MDB-formation by CYP81Q38. The two PSS enzymes, CYP81Q38 from *P. leptostachya* and CYP81Q1 from *S. indicum*, are most likely derived from a common gene in their ancestral plant, considering the close phylogenetic
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relationship of *P. leptostachya* and *S. indicum* and striking overall structural similarity between CYP81Q38 and CYP81Q1, with the characteristic Ala residue at the position of distal-Thr residue, although their gene expression profiles are distinct each other (mainly in root and seed, respectively) (Figures 5C, 5D). Thus, other phylogenetically related plants producing (+)-sesamin and its structurally related metabolites with MDB might also have an orthologous gene responsible to PSS. In this context, a woody plant of Lamiales, *Paulownia*, is one candidate plant since it accumulates a large amount of (+)-sesamin and its derivative (+)-paulownin in their heartwoods often used for traditional Japanese chests (Figure 1).

It should be noted that CYP719A is also known as a class of MDB forming P450s distinct from CYP81Qs so far. CYP81Qs and CYP719As are structurally discrete as shown in Figure 5C, and are indeed derived from phylogenetically unrelated plant lineages (Lamiales and Ranunculales order): *Sesamum* (Pedaliaceae) and *Phryma* (Phrymaceae) belong to Lamiales order, whereas *Coptis japonica* (Berberidaceae), opium poppy (Ranunculaceae), and *Podophyllum hexandrum* (Papaveraceae) belong to Ranunculales order. From the viewpoint of chemotaxonomy, *Podophyllum* is a unique lineage in Ranunculales because they exceptionally produce anti-tumor lignans with MDB such as podophyllotoxin instead of isoquinoline alkaloids that are more often observed specialized metabolites in Ranunculales plants. Therefore, it is conceivable that CYP719A have differentiated to be MDB forming P450 enzymes that are originally involved in the biosynthesis of isoquinoline alkaloids in the ancestral Ranunculales plant, and later *Podophyllum* CYP719A23 and CYP719A24 were specifically recruited to lignan biosynthesis (Marques et al. 2013). Since CYP719As commonly have a Ser residue instead of Ala residue, at the position of the conserved distal-Thr residue, it is fairly feasible that the crucial substitution events at the conserved Thr residue occurred independently in the distinct MDB forming P450s (CYP81Q and CYP719A). Furthermore, this notion is also applicable to the cases that phylogenetically unrelated lineages produce the same metabolites with MDB. For instance, *Gingko* (gymnosperm) and *Magnolia* plants (primitive angiosperm) are basal lineages of seed plants and phylogenetically distant from Lamiales. However, both plants are known to produce sesamin (Umezawa 2003a, 2003b). Furthermore, KNAPSAcK database search shows that sesamin has been found in various plants more than 130 species (Afendi et al. 2012), implying that PSS have independently occurred several times during seed plant evolution. Analogously, it is also suggested that various natural products with MDB that are widespread in plants occurred as consequence of convergent evolution of MDB forming activity (Figure 1). In addition to lineage-specific local gene multiplication (Chae et al. 2014) and spatiotemporal alteration of gene expression, functional plasticity of enzyme would be a driving force to generate vast structural diversity of specialized metabolites in nature.

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