Construction and Application of a Functional Library of Cytochrome P450 Monoxygenases from the Filamentous Fungus *Aspergillus oryzae*  
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A functional library of cytochrome P450 monoxygenases from *Aspergillus oryzae* (AoCYPs) was constructed in which 121 isoforms were coexpressed with yeast NADPH-cytochrome P450 40 oxidoreductase in *Saccharomyces cerevisiae*. Using this functional library, novel catalytic functions of AoCYPs, such as catalytic potentials of CYP57B3 against genistein, were elucidated for the first time. Comprehensive functional screening promises rapid characterization of catalytic potentials and utility of AoCYPs.

Cytochrome P450 (CYP) enzymes constitute a large superfamily of heme-containing monooxygenases that are distributed among a wide variety of organisms. An evolutionary history of extensive divergence from a common ancestor is implied (14, 20, 28). The majority of CYPs are highly specialized and play crucial roles in meeting the metabolic requirements of particular organisms. They are involved in secondary metabolic pathways, such as detoxification of xenobiotics and synthesis of secondary metabolites (9, 30). Over the last few years, the sequence database of CYPs has greatly enlarged and continues to increase (27, 31). A compilation of CYP sequences could facilitate understanding of metabolic diversity and evolutionary history among living organisms. In addition to their biological importance, the catalytic functions of CYPs are of great interest to those involved in biotechnology (4, 5, 8, 13, 16, 42). Biocatalytic regiospecific and stereospecific oxidations by CYPs attract much attention in industry because (i) use of biocatalysts allows reduction in the number of tedious steps, such as blocking and deblocking, that are common in conventional chemical synthesis, (ii) a single product without by-product formation reduces the cost of downstream purification steps, and (iii) enantiomeric mixtures (rather than racemic mixtures) are strictly required in some fields, such as pharmaceutical industry (12, 33, 39, 42). Therefore, a rational comprehensive approach toward elucidating catalytic potentials and utilities of CYPs is required to develop bioindustrial applications (8, 19, 34).

A wide variety of filamentous fungi are used to produce economically valuable consumer items. The filamentous fungus *Aspergillus oryzae* is one of the most widely utilized microorganisms. It has been used for more than 1,000 years in Japanese fermentation industries to produce indigenous products, such as sake (rice wine), miso (soybean paste), and shoyu (soy sauce). Besides traditional fermentation technologies, much recent research into *A. oryzae* has focused on production of recombinant enzymes and primary and secondary metabolites (1, 40). The whole genome of *A. oryzae* has been sequenced and is available to the public (21). Genomic data have the potential to provide a more thorough understanding of metabolic diversity and capability of *A. oryzae*. Recently, we identified 155 genes, including 13 putative pseudogenes of *A. oryzae* CYPs (AoCYPs), and cloned 121 full-length cDNAs encoding an open reading frame (25). Molecular and functional diversity of AoCYPs is presumably important to the metabolic capability of this fungus; however, biological functions and catalytic potentials remain obscure. A rational approach to find and exploit catalytic potentials of various CYP enzymes could facilitate a myriad of biocatalytic processes. In this study, we have aimed to develop a functional screening system for AoCYPs and to increase understanding of their functional information.

**Heterologous expression of AoCYPs.** Using the isolated cDNAs, we attempted to express recombinant enzymes for a functional screening system of AoCYPs. For catalytic activity, CYPs must be associated with electron transfer systems providing reducing equivalents. Thus, systematic production of both CYPs and their redox partner(s) are important to facilitate comprehensive functional screening (19, 34). In this study, we employed *S. cerevisiae* as a host strain for heterologous expression because (i) a yeast expression system seemed appropriate to obtain recombinant eukaryotic CYP without genetic engineering of cDNA, (ii) *S. cerevisiae* has only three CYP genes, and their catalytic functions are well characterized (26), and (iii) *S. cerevisiae* is potentially useful for industrial applications as well as basic research. In addition, cytochrome P450 oxidoreductase (CPR) from *S. cerevisiae* is capable of donating reducing equivalents for various CYPs. It has been shown that a wide variety of eukaryotic CYPs, including *Aspergillus* spp., could be functional with CPR from *S. cerevisiae* (3, 23, 24, 32). Presumably, eukaryotic CYPs can exhibit catalytic activities with a heterologous CPR. Therefore, we constructed...
a yeast coexpression system of AoCYP and yeast CPR using pGYR vector (24, 35, 36) (see Material and Methods in the supplemental material). To elucidate heterologous expression of AoCYPs, we analyzed the CO difference spectrum of \textit{S. cerevisiae} cells harboring the expression plasmid (29). Figure 1 depicts a CO difference spectrum of CYP628C1 expressed in \textit{S. cerevisiae}. No peak was observed around 450 nm from a transformant harboring a plasmid without an AoCYP, indicating that the recombinant AoCYP enzyme was successfully expressed in \textit{S. cerevisiae} in an active form. These results also implied that endogenous CYPs of \textit{S. cerevisiae} did not interfere with spectroscopic analysis of recombinant AoCYPs due to their low levels of expression in this host strain (2). We have confirmed significant expression of 84 AoCYPs based upon CO difference spectra.

**Comprehensive functional screening of AoCYPs.** To develop a functional screening system, each transformant was separately inoculated into 0.5 ml culture medium and grown in 96-well DeepWell plates (see Materials and Methods in the supplemental material). Transformants accommodated in the 96-well plates were easily replicated and used for further experiments. To validate potential utility of the screening system, we initiated a functional survey using 7-ethoxycoumarin as a tentative model substrate. After 2 days of incubation, significant product formation was achieved by \textit{S. cerevisiae} harboring expression plasmids of CYP57B3, CYP62C2, CYP68Q1, CYP531E1, CYP620G1, CYP620H1, CYP620H3, CYP620H9, CYP628C1, CYP675A2, CYP5061B5, CYP5078A5, and CYP5080B2. The product was identified as 7-hydroxycoumarin by comparison of the retention time on high-performance liquid chromatography (HPLC) with an authentic standard (Fig. 2). Moreover, several AoCYPs, such as 5061B5, yielded a minor unidentified product(s), in addition to the formation of 7-hydroxycoumarin (Fig. 2). Although further information should aim to elucidate their biochemical and biophysical properties, it can be hypothesized that the natural substrate(s) of these AoCYPs might be structurally related to coumarin, such as polyketide derivatives. It is thus a good example that non-target-driven screening could provide new insight into the fascinating biology and metabolic processes of \textit{A. oryzae}. Interestingly, catalytic conversion of 7-ethoxycoumarin to 7-hydroxycoumarin was clearly demonstrated by the transformant harboring the CYP5080B2 expression plasmid; nevertheless, we could not confirm its expression by CO difference spectrum.

Such observation was also found from several AoCYPs using different substrates. Combining the results of spectroscopic analysis and bioconversion, we could conclude that at least 92 AoCYPs were functionalized in \textit{S. cerevisiae}. These results highlight the potential utility of our screening system.

Using the functional screening system, we further carried out targeted functional screening of AoCYP to produce value-added isoflavonoids. Genistein is one of the major isoflavonoids found in soybean. Because of its biological and pharmacological activities, genistein derivatives are likely to have various useful applications (6, 10, 11, 15, 17, 41). When genistein was utilized as a substrate, significant product formations were achieved by \textit{S. cerevisiae} expressing CYP57B3 (Fig. 3). Incorporation of a hydroxyl group was clearly demonstrated by LC-electrospray ionization (ESI)-mass spectrometry (MS) analysis (data not shown). Based upon \(^1\)H nuclear magnetic resonance (NMR) spectroscopic analysis, products were identified as 8-hydroxy-, 6-hydroxy-, and 3'-hydroxygenistein (Fig. 2).

![FIG. 1. CO difference spectrum of CYP628C1 expressed in \textit{S. cerevisiae}.](http://example.com/fig1.png)

![FIG. 2. Screening of AoCYPs catalyzing 7-ethoxycoumarin conversions.](http://example.com/fig2.png)
4). To the best of our knowledge, this is the first report describing catalytic activities of CYP57B3 against genistein and suggesting its possible roles in production of hydroxygenistein during soybean fermentation processes (6, 11). Based on sequence comparison, CYP57B3 seems phylogenetically close to a CYP from the plant pathogen *Nectria haematococca*; a fungus involved in biodegradation of phytoalexin pisatin (22). Although it is still unclear whether CYP57B3 exhibits catalytic activities against pisatin, it may play a biological role in the metabolism of phytoalexins with isoflavone skeletons. Biological and pharmacological potentials of the products have also been reported. 8-Hydroxygenistein shows much stronger antioxidative activity than genistein (11) and exhibits antiproliferative activity against cancer cells (15). 3'-Hydroxygenistein, known as orobol, exhibits unique pharmacological activities, such as enhancing sensitivity of human ovarian carcinoma cells against the anticancer drug paclitaxel (17) and inhibiting human immunodeficiency virus 1 integrase (41). Although hydroxylated genistein can be isolated from natural products, including fermented products, these natural compounds are limited in supply and problematic for practical utilization. Because the synthesis of isoflavones remains an important object (10, 37), it would be of great interest to utilize CYP57B3 for production of value-added rare isoflavonoids from genistein. However, further research efforts should aim to improve conversion efficiency for practical application. It has been reported that reaction efficiency can be influenced by redox partner(s) such that native combination of CYP-CPR from the same organism shows slightly better functionality than heterologous combination and several monooxygenase reactions are coupled with cytochrome *b5* (7, 18, 30, 38). Further investigation and possible reengineering of CYP57B3 and its redox partner(s) should facilitate systematic understanding for the biotechnology sector.

Through functional screening, the novel activities of AoCYPs were experimentally explored using a series of compounds, such as flavonoids, terpenoids, steroids, and pharmaceutical chemicals (Table 1). To the best of our knowledge, this is the first report experimentally demonstrating functional diversity of *Aspergillus* CYP. The functional screening system promises rapid and comprehensive characterization of the catalytic potential of AoCYPs, which could further provide novel insight into the biology and
biotechnology. In this study, we have focused on the catalytic capability of CYP57B3 to produce value-added hydroxylated products from genistein. However, the screening system should be applicable for a myriad of compounds and facilitate investigation in industrial and biological fields. A thorough understanding of AoCYP functions will open the door for advanced fungal biology and biotechnology.

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REFERENCES


