CYP82Y1 Is N-Methylcanadine 1-Hydroxylase, a Key Noscapine Biosynthetic Enzyme in Opium Poppy*

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Background: The biochemistry of noscapine metabolism in opium poppy has not been established.
Results: CYP82Y1 catalyzes the efficient 1-hydroxylation of N-methylcanadine. Silencing CYP82Y1 reduced noscapine accumulation in opium poppy plants.
Conclusion: CYP82Y1 is N-methylcanadine 1-hydroxylase and functions in noscapine biosynthesis.
Significance: The enzyme catalyzing the first step in the conversion of N-methylcanadine to noscapine has been identified and characterized.

Noscapine is a phthalideisoquinoline alkaloid investigated for its potent pharmacological properties. Although structurally elucidated more than a century ago, the biosynthesis of noscapine has not been established. Radiotracer studies have shown that noscapine is derived from the protoberberine alkaloid (S)-scoulerine and has been proposed to proceed through (S)-N-methylcanadine. However, pathway intermediates involved in the conversion of N-methylcanadine to noscapine have not been identified. We report the isolation and characterization of the cytochrome P-450 CYP82Y1, which catalyzes the 1-hydroxylation of N-methylcanadine to 1-hydroxy-N-methylcanadine. Comparison of transcript and metabolite profiles of eight opium poppy chemotypes revealed four cytochrome P-450s, three from the CYP82 and one from the CYP719 families, that were tightly correlated with noscapine accumulation. Recombinant CYP82Y1 was the only enzyme that accepted (R,S)-N-methylcanadine as a substrate with strict specificity and high affinity. As expected, CYP82Y1 was abundantly expressed in opium poppy stems where noscapine accumulation is highest among plant organs. Suppression of CYP82Y1 using virus-induced gene silencing caused a significant reduction in the levels of noscapine, narcotine, and a putative downstream secoberbine intermediate and also resulted in increased accumulation of the upstream pathway intermediates scoulerine, tetrahydrocolumbamine, canadine, and N-methylcanadine. The combined biochemical and physiological data support the 1-hydroxylation of (S)-N-methylcanadine catalyzed by CYP82Y1 as the first committed step in the formation of noscapine in opium poppy.

Noscapine, also known as narcotine, anarcastine, or gnoscapine, is a benzylisoquinoline alkaloid (BIA)3 belonging to the phthalideisoquinoline alkaloid subgroup and displaying potent pharmacological properties. Noscapine was first detected in 1803 by the French industrial chemist Charles Derosne and referred to as “sel narcotique de Derosne” (Derosne’s narcotic salt) (1). Although more than two centuries have passed since the discovery of noscapine, the chemistry and molecular biochemistry of phthalideisoquinoline alkaloid biosynthesis have not been established. Extensive radiotracer experiments in the 1960s showed that the structural scaffold of noscapine shares the same biosynthetic origin as other BIAs. Labeled noscapine was isolated when opium poppy (Papaver somniferum) plants were fed radioactive tyrosine, norlaudanosoline, or laudanosoline (2, 3), suggesting that the noscapine carbon skeleton, apart from the lactonic carbonyl group, is formed by the condensation of two tyrosine derivatives. Later studies with labeled laudanosoline and reticuline showed that the lactone carbonyl carbon of phthalideisoquinoline alkaloids originates from the N-methyl moiety of the same pathway intermediate yielding the C8 position of protoberberine and protoine alkaloids. The biosynthetic relationship among these alkaloid subgroups was further demonstrated by the efficient incorporation of labeled scoulerine and canadine into noscapine (2, 4). Recently, the role of scoulerine and canadine as intermediates in noscapine biosynthesis has been corroborated by the isolation and characterization of the genes encoding SOMT1 (scoulerine 9-O-methyltransferase) and the cytochrome P-450 canadine synthase (CYP719A21) from opium poppy (see Fig. 1) (5, 6). A concomitant decrease in the levels of noscapine, tetrahydrocolumbamine, and canadine in plants with suppressed SOMT and CYP719A21 transcript abundance was demonstrated (5, 6). The detection of several naturally occurring phthalideisoquinoline alkaloids such as macrantaldehyde, papaveroxynoline, papaveroxine, and narcotinohemiacetal in certain Papaver species (7) suggested a biosynthetic process involving the

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The abbreviations used are: BIA, benzylisoquinoline alkaloid; CID, collision-induced dissociation; CPR, cytochrome P-450 reductase; CYP, cytochrome P-450; ESI, electrospray ionization; NOS, noscapine synthase; TRV, tobacco rattle virus; VIGS, virus-induced gene silencing; qRT, quantitative real time.
N-methylation of a protoberberine alkaloid, most likely canadine, yielding N-methylcanadine (8), which is hydroxylated at C1 and subsequently methylated to form 1-methoxy-N-methylcanadine (9). Hypothetical C8 oxidation of 1-methoxy-N-methylcanadine yields macranaldehyde. Hydroxylation of macranaldehyde at C13 would produce narcotinehemiacetal, which could be dehydrogenated to noscapine.

The recent discovery of a 10-gene cluster in the opium poppy genome putatively encoding all noscapine biosynthetic enzymes, except TNMT (tetrahydroprotoberberine N-methyltransferase), operating downstream of (S)-scoulerine prompted a re-evaluation of the pathway model (10). Accordingly, N-methylcanadine was proposed to undergo ring opening via C8 oxidation catalyzed by a cytochrome P-450 (CYP) yielding a 3-hydroxysecoberbine intermediate potentially prior to C4 hydroxylation, corresponding to C1 on N-methylcanadine (see Fig. 1). Subsequent hydroxylation followed by O-methylation at C4’ and O-acetylation of the 3-hydroxy group on the secoberbine intermediate was suggested to yield 3-O-acetyl-papaveroxine, which would be O-deacetylated by a carboxylesterase to form narcotinehemiacetal. The hemiacetal moiety of narcotinehemiacetal ultimately yields the lactone ring in noscapine. The final step was validated by the recent isolation and characterization of noscapine synthase (NOS), a short chain dehydrogenase/reductase that converts narcotinehemiacetal to noscapine in opium poppy (see Fig. 1) (11).

Members of the CYP superfamily catalyze several key conversions in BIA metabolism. Three CYP families, CYP80, CYP82, and CYP719, have been implicated in BIA biosynthesis (12). Enzymes in the CYP80 family catalyze the aromatic ring 3’-hydroxylation of N-methylcoclaurine (13), C-O coupling between 1-benzylisoquinoline monomers to form the bisbenzylisoquinoline alkaloid berbamunine (14), and intramolecular C-C phenol coupling of (S)-reticuline yielding the aporphine alkaloid (S)-corytuberine (15). Members of the CYP719 family are involved in the formation of methylenedioxy bridges on uncharacterized reactions. $R_1$ and $R_2 = H$ or OH. Partial carbon numbering is shown for N-methylcanadine and the putative secoberbine intermediate. [O], uncharacterized oxidative conversions.
protoberbine alkaloids (16–18), such as tetrahydrocolumbamine to yield canadine (Fig. 1) (6, 17), and the C-C phenol coupling of (R)-reticuline to form the promorphinan alkaloid salutaridine (19). Recently, enzymes of the CYP82 family have been shown to catalyze regiospecific aliphatic hydroxylations of (i) quaternary protoberbine alkaloids including N-methylcanadine, which causes tautomerization to the protopine backbone (20), and (ii) protopine alkaloids resulting in spontaneous intramolecular rearrangement to the benzo[c]phenanthridine scaffold (21). Although there are three CYP82s and one CYP719 in the proposed noscapine biosynthetic gene cluster (10), only CYP719A21 has been functionally characterized as canadine synthase (6). In this paper, we report the isolation and characterization of CYP82Y1 gene transcripts with opium poppy chemo-

types accumulating noscapine, compared with noscapine-free chemo-
types. CYP82Y1 catalyzed the regiospecific 1-hydrox-
ylation of the quaternary protoberbine alkaloids (R,S)-N-
methylcanadine and (S)-N-methylstylopine. The physiological role of CYP82Y1 in opium poppy was partially supported using VIGS. Formation of 1-hydroxy-

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**EXPERIMENTAL PROCEDURES**

*Plant Material*—The opium poppy chemotypes Bea’s Choice and Veronica (22) were cultivated at 20/18 °C (light/dark) in a growth chamber (Conviron, Winnipeg, Canada) with a photoperiod of 16 h and a combination of Cool White fluorescent (Sylvania, Mississauga, Canada) and incandescent lighting (5). Plant samples were collected as described previously (6).

*Chemicals*—(R,S)-Canadine was purchased from Latoxan (Valence, France). (R,S)-N-Methylcanadine was produced from (R,S)-canadine using purified recombinant *Eschscholzia cali-

fornica* TNMT (8). Enzymatic conversion was performed for 4 h with 100 μM (R,S)-canadine, 200 μM S-adenosyl-L-methio-
nine, and 75 μg of purified TNMT. Reactions were terminated by the addition of an equal volume of methanol, and precipitated protein was removed by centrifugation. The supernatant was concentrated under reduced pressure, resuspended in 0.1 N NaOH, and loaded on a Strata™-X-CW 33-μm polymeric weak cation, 30 mg ml⁻¹ column (Phenomenex, Torrance, CA). (S)-N-Methylcanadine was selectively eluted from the column using formic acid:methanol (5:95). The purity and identity of (S)-N-methylcanadine was confirmed by LC-MS/MS. (S)-N-

Methylstylopine was similarly prepared from (S)-stylopine (8). Other alkaloids were obtained as described previously (5, 8, 23). S-Adenosyl-L-methionine was purchased from Sigma-Aldrich. All other chemicals were purchased from Bioshop (Ontario, Canada).

*Phylogenetic Analysis*—Amino acid alignments were performed using ClustalW (24), and a phylogenetic tree was built from bootstrap values generated using the Geneious (Biomat-
ters, Newark, NJ) software package. GenBank™ accession numbers for sequences used to construct the phylogenetic tree are as follows: PsCYP82N4 (N-methylstylopine 14-hydroxylase), *P. somniferum* N-methylstylopine 14-hydroxylase (KC154003); PsCYP82Y1, *P. somniferum* CYP82Y1 (JQ659005); PsCYP82X1, *P. somniferum* CYP82X2 (JQ659004); CjCYP82R1, *Coptis japonica* CYP82R1 (BAF98472), PsCYP82N3 (protopine 6-hydroxylase), *P. somniferum* protopine 6-hydroxylase (KC154002); EcCYP82N2v2 (protopine 6-hydroxylase), *E. californica* protopine 6-hydroxylase (AB598834); PsCYP719A21 (canadine synthase (CAS)), *P. somniferum* canadine synthase (JQ659003); PsCYP719B1 (SalSyn), *P. somniferum* salutaridine synthase (ABR14720); PsCYP719A25 (schizanthine synthase), *P. somniferum* schizanthine synthase (GU325749); PsCYP80B3 (N-methylcoclaurine 3’-hy-

droxylase), *P. somniferum* N-methylcoclaurine 3’-hy-

droxylase (AF191722); EcCYP80B1 (N-methylcoclaurine 3’-

-hydroxylase), *E. californica* N-methylcoclaurine 3’-

-hydroxylase (AF104801); CjCYP80G2 (corytuberine synthase), *C. japonica* corytuberine synthase (AB288053); AtCYP82C2, *Arabidopsis thaliana* CYP82C2 (O49394); NcCYP82E4v1, *Nicotiana tabacum* CYP82E4v1 (DQ131886); AtCYP82G1, *A. thaliana* CYP82G1; VvCYP82C4, *Vitis vinifera* CYP82C4 (XP_002284810); and GmCYP82G4, *Glycine max* CYP82G4 (XP_003523888).

*Isolation of CYP82 Candidates*—Total RNA and alkaloid extracts from the latex of eight opium poppy chemo-
types were subjected to transcript and metabolite profiling, respectively, as described previously (5, 22). CYP82 candidates were identified among genes differentially expressed in noscapine-free (Debo-

rah, Przemko, 40, and T) and noscapine-producing (Natalia, Marianne, Roxanne, and Veronica) chemo-
types of opium poppy. The full-length coding regions of CYP82X1, CYP82X2, and CYP82Y1 were assembled in silico by searching each data-
base using the tBLASTn algorithm. Relative transcript abun-
dance was determined as the number of reads corresponding to each selected candidate compared with the total number of reads in each database (5).

*Gene Expression and Noscapine Accumulation in Different Plant Organs*—Total RNA from different opium poppy Veron-

ica organs were collected by flash freezing tissues in liquid N₂ prior to excision from the plant to preserve the latex content. Total RNA was extracted from each organ, and cDNA was syn-
thesized as described previously (5). The relative abundance of CYP82Y1 transcripts in each organ was determined by quanti-
tative real time PCR (qRT-PCR) analysis using the primer set: 5’-TGAACCTTTCGCGTGTATTA-3’ and 5’-CTATGAC-TACCGAATCTAAACAG-3’. Plant tissues were lyophilized, and alkaloids were extracted to measure noscapine content.

*Cloning and Expression of CYP82 Candidates*—Full-length coding regions of CYP82X1 and CYP82Y1 were amplified from cDNA derived from total stem RNA of the Bea’s Choice chemo-
type using Takara Ex Taq DNA polymerase (Fisher Sci-

tentific, Ottawa, Canada) and the following primer sets: CYP82X1, 5’-GGGGCCGCGCCATGGAGTTATTCATGTAAGAGCCGC; CYP82Y1, 5’-ATTAGGCCGCGCCACATGCGGATTTGTTATGATGCAATG-3’ and 5’-CTATGAC-TACCGAATCTAAACAG-3’. A synthetic CYP82X2 gene based on native nucleotide sequences was constructed (Genscript, Piscataway, NJ) by replacing the first 60 N-terminal amino acids of CYP82X2 with 43 N-terminal amino acids from lettuce germacrene A oxidase to improve heterologous expression in yeast (25). For heterol-

ogous production of FLAG-tagged CYP82 cDNAs, the cognate
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cDNAs were ligated into the NotI and SpeI restriction sites of the dual plasmid pESC-leu2d::CPR (20, 26) yielding pESC-Leu2d::CYP82X1/CPR, pESC-Leu2d::CYP82X2/CPR, and pESC-Leu2d::CYP82Y1/CPR. Yeast harboring pESC-leu2d::CPR was used as the negative control.

Yeast Culture, Preparation of Microsomes, and Immunoblot Analysis—The protease-deficient Saccharomyces cerevisiae strain YPL 154c: Pep4 was transformed with pESC-Leu2d::CYP82X1/CPR, pESC-Leu2d::CYP82X2/CPR, and pESC-Leu2d::CYP82Y1/CPR. Transformed yeast were used to inoculate 2 ml of synthetic complete (SC) medium lacking leucine (SC−Leu), but containing 2% (w/v) glucose, and cultured overnight on a gyratory shaker with gentle agitation (60 rpm). Reactions in vitro by varying the alkaloid substrate concentration in the standard CYP82X1/CPR were collected immediately prior to anthesis as described previously (5). Infiltration with A. tumefaciens was confirmed by detection of the RNA corresponding to the TRV2 transgene cassette in VIGS-treated plants using TRV-MCS primer sets specific to sequences (i) flanking the multiple cloning site of pTRV2 (5′-GGTCAAGGTAGTCTAGAGAG-3′ and 5′-CGAGAATGTCATCTCAGG-3′) (29) and (ii) encoding GAPDH as a positive control (5′-CTATTTTGAAGGGTGGA- GC-3′ and 5′-GTATTTGCGGAGAGTGG-3′). Latex samples from infiltrated plants were lyophilized, resuspended in methanol at a concentration of 0.1 μg μl−1, and extracted overnight at −80°C. Transcript analysis of infiltrated plants was performed by qRT-PCR using a 7300 real time PCR system (Applied Biosystems, Burlington, Canada) for 40 cycles of template denaturation, primer annealing, and primer extension. Each 10-μl PCR contained 1 μl of cDNA, 300 nm forward and reverse primers, and a 1× KAPA SYBR FAST qPCR kit (Kapa Biosystems, Boston, MA). The opium poppy gene encoding ubiquitin was used as an endogenous reference and amplified with the primer set: 5′-CCATTGGGTGTCTCGTCTAC-3′ and 5′-CAAGCCATAGCTGAAACACC-3′. Primer sets used to perform qRT-PCR on other genes were: CYP82Y1, 5′-ACACTTTCCGCTTGTATTA-3′ and 5′-CTATGACTACCCGAGATTCTAACAG-3′; CYP82X1, 5′-GGGAAAGAAAAAGGACACAAATGC-3′ and 5′-CActGGTTTACCACTTCCCCAG-3′; CYP82X2, 5′-ACGAATGGTTGGCTGACATGAT-3′ and 5′-AAAGACCTTCTCAATGCTCT-3′; CYP719A2 (CAS), 5′-CTACCAATCCATCTGCATTACCTGCG-3′ and 5′-GGTAACTTGGTGGACAGCTGC-3′. Recombinant enzymes were detected by immunoblot analysis as described previously (6).

CYP82 Enzyme Assays—Standard enzyme assays were performed at 30°C for 30 min in 200 μl of 100 mM HEPES-NaOH, pH 7.5, containing 5 mg of total microsomal proteins, 500 μM NADPH, and 50 μM (R, S)-canadine or (R, S)-N-methylcanadine on a gyratory shaker with gentle agitation (60 rpm). Reactions were stopped by the addition of 800 μl of methanol. Microsomal protein extracted from yeast harboring pESC-leu2d::CPR was used as a negative control.

In Vitro Biochemical Characterization of CYP82Y1—Standard in vitro enzyme assays were performed as described above. A variety of alkaloids belonging to several different BIA subgroups (1-benzylisoquinoline, morphinan, protoberberine, aporphine, benzo[c]phenanthidine, protopine, phthalidisoquinoline, and bisbenzylisoquinoline) were tested to determine the substrate range of CYP82Y1 (see Table 1). Steady-state enzyme kinetics for (R, S)-N-methylcanadine were determined by varying the alkaloid substrate concentration in the standard in vitro assay between 0 and 400 μM at a fixed concentration of 500 μM NADPH. Kinetic constants for CYP82Y1 were calculated by fitting initial velocity versus substrate concentration to the Michaelis-Menten equation using the GraphPad Prism 5.0 (GraphPad Software, San Diego, CA).

Virus-induced Gene Silencing—The tobacco mosaic virus (TRV) vector system (28) was used to suppress transcript levels of CYP82Y1 in opium poppy plants. A unique sequence encompassing part of the 3′-UTR and coding region of CYP82Y1 was amplified with the primer set: 5′-TTAATCAGACTCACCCACGGACACATGATGAC-3′ and 5′-AGGTCGAGATCCACCAAAAGGAAATATTATAAG-3′. The fragment was cloned into pTRV2, and vectors were mobilized in Agrobacterium tumefaciens as described previously (6). Apical meristems of 2−3-week-old seedlings were infiltrated with a 1:1 mixture of A. tumefaciens harboring pTRV1 and either pTRV2::CYP82Y1 or pTRV2. Infiltrated plants were grown in the greenhouse for 8−10 weeks. Visual confirmation of gene silencing was monitored using the pTRV2-PDS construct encoding phytoene desaturase (29). Latex and stems of infiltrated opium poppy were collected immediately prior to anthesis as described previously (5).
vent B (acetonitrile), 6–7 min ramp to 99% solvent B, 7–9 min isocratic at 99% solvent B, and 9–13 min ramp to 0% solvent B; (ii) for VIGS, 0–6 min 60% solvent B, 6–9 min ramp to 99% solvent B, 9–14 min isocratic at 99% solvent B, and 14–18 min ramp to 0% solvent B. Samples were injected into the mass analyzer via an electrospray ionization (ESI) probe inlet. Ions were generated and focused using the following parameters: (i) for enzyme assays, capillary voltage, 4000 kV; gas flow, 9 liter min⁻¹, fragmentor voltage, 100 V; nebulizer pressure, 40 psi; gas temperature, 330 °C; (ii) for VIGS, capillary voltage, 4000 kV; gas flow, 10 liter min⁻¹; fragmentor voltage, 100 V; nebulizer pressure, 50 psi; gas temperature, 350 °C. Mass spectrometry data were acquired in positive ion mode in the range of m/z 200–700. Positive mode electrospray ionization (ESI) collision-induced dissociation (CID) spectra were recorded using collision energy of 25.0 eV applied in quadrupole 2 and an argon collision gas pressure of 1.8 × 10⁻³ torr. MS² fragments were generated by heated ESI with the following parameters: heater, 400 °C; sheath gas, 60 au; auxiliary gas, 20 au; spray voltage, 3 kV. Ion interface settings were 380 °C and 38 V (capillary) and 85 V (tube lens). MS³ experiments were performed using an LTQ-Orbitrap XL equipped with a syringe pump and an Accela HPLC system (Thermo- Fisher Scientific). Alkaloids (1 μg ml⁻¹) were introduced continuously with a syringe pump (5 μl min⁻¹) into the HPLC flow at a rate of 500 ml min⁻¹ using acetonitrile, and positive ions were generated by heated ESI with the following parameters: heater, 400 °C; sheath gas, 60 au; auxiliary gas, 20 au; spray voltage, 3 kV. Ion interface settings were 380 °C and 38 V (capillary) and 85 V (tube lens). MS³ experiments were performed by conducting CID on target ions isolated in the linear ion trap followed by high resolution (60,000 full width at half-maximum; <2 ppm error) mass analysis of the resulting fragment ions in the Orbitrap XL. Full scan data were collected in centripetal mode over mass ranges extending from the lowest permissible value up to 10 atomic mass units beyond the parent ion. Detection methods consisted of three scan events of ~1 s each with a total run time of 10 min. External and internal instrument calibration ensured an error of <2 ppm.

**NMR Analysis**—1-Hydroxy-N-methylcanadine was produced from N-methylcanadine in scaled-up CYP82Y1 reactions terminated with an equal volume of methanol. Precipitated protein was removed by centrifugation, the supernatant was concentrated under reduced pressure, resuspended in 10 mM ammonium acetate:acetonitrile (95:5), and loaded on a Strata™-X-33μ polymeric reversed phase, 200 mg ml⁻¹ column (Phenomenex, Torrance, CA) to remove non-alkaloid components. N-Methylcanadine and 1-hydroxy-N-methylcanadine were eluted from the column using 10 mM ammonium acetate:acetonitrile (70:30), and the sample was concentrated under reduced pressure. Compounds were separated on a 250-μm silica-coated glass TLC plate using a mobile phase of isopropanol:ammonia hydroxide:water 9:2:2 (Moravik Biochemical, Brea, CA). Bands corresponding to N-methylcanadine and 1-hydroxyl-N-methylcanadine were excised, eluted with 50% (v/v) ethanol, dried under reduced pressure, and lyophilized with water. One milligram of each compound was dissolved in CD₃OD, and ¹H COSY NMR spectra were recorded in 5-mm Shigemi NMR tubes (Shigemi, Allison Park, PA) on a Varian VNMRS three-channel 600 MHz spectrometer. Data were interpreted based on ¹H NMR spectra for (−)-cis-N-methylcanadine reported previously (31, 32).

**RESULTS**

Identification of CYP Candidates—Available opium poppy chemotypes display two distinct phenotypes with respect to the accumulation of noscapine (Fig. 2A) (5). Whereas the chemotypes Marianne, Natasha, Roxanne, and Veronica accumulate high levels of noscapine, the chemotypes Deborah, 40, T, and Przemko display only trace to undetectable noscapine content. Marianne contained the highest levels of noscapine, and Roxanne and Veronica accumulated papaverine in addition to noscapine. Compared with noscapine-free plants, the high noscapine chemotypes accumulated substantially less morphinan alkaloids.

Comparing the Marianne and Deborah transcriptome databases for differentially expressed genes revealed a variety of CYPs, O-methyltransferases (5), a short chain dehydrogenase/reductase (11), an acetyltransferase, and a carboxyl esterase for...
which transcripts were detected only in noscapine-producing chemotypes. Focusing on CYPs that displayed (i) sequence similarity with previously reported BIA biosynthetic enzymes (i.e., members of the CYP80, CYP82, and CYP719 families) and (ii) gene expression profiles that strictly correlated with the accumulation of noscapine in different opium poppy chemotypes yielded four candidate genes. One candidate belongs to CYP719 family, whereas three others are members of the CYP82 family (Fig. 2B). Other characterized CYPs involved in BIA biosynthesis did not exhibit a transcript accumulation pattern that correlated with the accumulation of noscapine. Except for CYP719A21, which has been characterized as canadine synthase (6), biochemical functions have not been assigned for CYP82X1, CYP82X2, and CYP82Y1.

The CYP82X1, CYP82X2, and CYP82Y1 cDNAs contained open reading frames encoding 540, 554, and 556 amino acids, respectively. Phylogenetic analysis based on an alignment of CYP protein sequences involved in BIA biosynthesis highlighted the distinction between CYP82 members, and the CYP719 and CYP80 families (Fig. 3). CYP82Y1 shares the highest amino acid sequence identity with other opium poppy CYPs shown or purported to play a role in BIA metabolism: CYP82N4 (53%), CYP82X1 (50%), CYP82N3 (47%), putatively protopine 6-hydroxylase, and CYP82X2 (43%). Among functionally characterized CYPs from other plant species, CYP82Y1 displayed the highest amino acid sequence identity with EcCYP82N2v2 (44%), which is protopine 6-hydroxylase (21). Based on substantial amino acid sequence similarity with CYP82N4, which was functionally characterized as N-methylstilopine 14-hydroxylase and also acted on N-methylcanadine (20), CYP82Y1 might also accept quaternary protoberberine alkaloids as substrates.

Expression and Characterization of CYP82Y1—A galactose-inducible yeast dual expression vector pESC-Leu2d was used to simultaneously express opium poppy CYP82 and CPR cDNAs in S. cerevisiae. The CYP82 coding regions were inserted into the multiple cloning site of pESC-Leu2d under the control of the Gal-10 promoter and in-frame with the FLAG epitope. In the same pESC-Leu2d vector, CPR was inserted under the Gal-1 promoter and in-frame with the c-Myc epitope. Co-expression of CYP82 and CPR cDNAs was confirmed by immunoblot analysis of microsomal fractions prepared from S. cerevisiae cultures harboring the pESC-leu2d:CYP82X1/CPR, pESC-leu2d:CYP82X2/CPR, and pESC-leu2d:CYP82Y1/CPR vectors using FLAG and c-Myc antibodies to detect epitope-tagged recombinant proteins (Fig. 4A).
NR

was accepted as a substrate, although with considerably lower efficiency (38% of the substrate turnover rate compared with (R,S)-N-methylcanadine). Microsomal fractions of yeast harboring pESC-leu2d::CPR (i.e., lacking a CYP82 candidate) did not catalyze the formation of any products with either (R,S)-N-methylcanadine or (S)-N-methylstylopine. CYP82Y1 showed an absolute requirement for the quaternary N-methyl moiety because neither (R,S)-canadine nor (S)-stylopine were accepted as substrates. No compounds belonging to other BIA subgroups were accepted as substrates. Notably, no conversions to compounds with m/z reduced by 2 atomic mass units compared with the potential substrate were detected, ruling out the possible occurrence of C-C bond formation. The increased m/z by 16 atomic mass units of the reaction products compared with the m/z of N-methylcanadine and N-methylstylopine suggested the hydroxylation of these substrates.

**Structural Elucidation of the CYP82Y1 Reaction Product**—High resolution MS³ analysis suggested aromatic ring hydroxylation at positions C1 or C4 on the isoquinoline moiety of N-methylcanadine. The ion trap portion of a hybrid LTQ-Orbitrap-XL instrument isolated and fragmented parent ions and subsequent daughter ions, and the Orbitrap analyzer acquired high resolution mass data for all MS² and MS³ fragment ions. Because authentic standards were not available, mass fragmentation data for the enzymatic reaction product (1-hydroxy-N-methylcanadine) and the substrate (N-methylcanadine) were compared with authentic standards of the phthalideisoquinoline alkaloids narcotoline and hydastine. Narcotoline and 1-hydroxyl-N-methylcanadine both possess a 1-hydroxyl group, whereas hydastine and N-methylcanadine are not hydroxylated at the C1 position. CID of both phthalideisoquinoline and protoberberine alkaloids generally yielded a principal isoquinoline fragment (33), which allowed MS³ fragmentation analysis of equivalent MS² daughter ions. Narcotoline and the N-methylcanadine reaction product yielded isoquinoline fragments at 206.08124 and 206.08123 m/z, respectively, corresponding to an elemental formula of C₁₁H₁₂O₂N and indicating that either C1 or C4 hydroxylation of N-methylcanadine had occurred (Fig. 5, A and B). In contrast, MS² of hydastine and the N-methylcanadine substrate both yielded fragments at 190.08425 m/z corresponding to an elemental formula of C₁₁H₁₂O₂N, which is consistent with an isoquinoline moiety lacking a 1-hydroxyl moiety (Fig. 5, C and D). To support the 1-hydroxylation of N-methylcanadine, CID analysis of all isoquinoline ions was performed. MS³ revealed nearly identical spectra for both 1-hydroxylated isoquinoline moieties of 1-hydroxyl-N-methylcanadine and narcotoline (Fig. 5, E and F) and similarly identical spectra for both nonhydroxylated isoquinoline ions (Fig. 5, G and H) of N-methylcanadine and hydastine. Because the product derived from N-methylstylopine produced a similar MS/MS fragmentation pattern, we conclude that a regiospecifically identical hydroxylation event occurred with N-methylstylopine (m/z 338.2) yielding 1-hydroxy-N-methylstylopine.

The CYP82Y1 reaction product derived from N-methylcanadine was confirmed as 1-hydroxyl-N-methylcanadine by NMR spectroscopy. The 1H COSY NMR (600 MHz, CD₃OD, δH) spectrum of N-methylcanadine showed 16 proton signals con-

### TABLE 1

Benzylisoquinoline alkaloids tested as potential CYP82Y1 substrates

| Alloisocryptopine | Cypretopine | Papaverine |
|-------------------|-------------|------------|
| Berbine           | Dehydroreticuline | Pavine     |
| Berberine         | (-)-Glucine  | Protopine  |
| (-)-Bolinde       | Hydrostine   | Reticuline |
| (-)-Bulbocapnine  | (-)-Isobehen | Salutaridine |
| Bicuculline       | (S)-N-Methylcanadine | Sanguinarine |
| Canadine          | Narcotheimiecetal | Stylopine |
| Chelerythine      | Narcotheine | (R,S)-Tetrahydrocolumbamine |
| (-)-Coclaurine    | (R,S)-Norlaudanosoline | (R,S)-Tetrahydropalmatine |
| Cortexine         | Nocaspine   | Thebaine   |

(R,S)-Canadine and (R,S)-N-methylcanadine were tested as potential substrates for the three CYP82 candidates. In the presence of NADPH, only N-methylcanadine (m/z 354) was accepted by CYP82Y1, resulting in the formation of a reaction product with m/z 370 (Fig. 4B). CYP82X1 and CYP82X2 did not show activity with (R,S)-canadine or (R,S)-N-methylcanadine. The substrate specificity of CYP82Y1 was determined using 36 BAs representing several different structural subgroups including 1-benzylisoquinoline, protoberberine, benzopenanthridine, aporphine, protopine, morphinan, pavine, and phthalideisoquinoline alkaloids (Table 1). In addition to (R,S)-N-methylcanadine (m/z 354), only (S)-N-methylstylopine...
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Physiological Role of CYP82Y1 in Noscapine Biosynthesis—A unique region of the coding region and 3′-UTR was used to construct pTRV2–82Y1 to suppress CYP82Y1 transcript levels in opium poppy plants using VIGS (Fig. 7A). The Bea’s Choice chemoype accumulates noscapine in addition to papaverine, morphine, and other BIAs. TRV infection of infiltrated plants was confirmed by reverse transcription-PCR amplification of TRV2 RNA (Fig. 7B). CYP82Y1 transcript levels were significantly reduced in plants infiltrated with A. tumefaciens harboring the pTRV2–82Y1 construct compared with the empty pTRV2 vector control (Fig. 7C). Total alkaloid content and the levels of major alkaloids including morphine, codeine, reticuline, thebaine, and papaverine were not altered in CYP82Y1-silenced plants compared with controls (Fig. 7, D and E). However, the suppression of CYP82Y1 transcript levels significantly reduced the accumulation of noscapine (Fig. 7D). CYP82Y1-silenced plants also accumulated significantly higher levels of scoulerine, canadine, N-methylcanadine, and tetrahydropalmatine. Suppression of CYP82Y1 transcription abundance also caused a significant reduction in the levels of narcotoline and an alkaloid displaying a secoberville-type CID spectrum (Fig. 7E). Notably, CYP82Y1 silencing also significantly reduced the relative abundance of SOMT1, SOMT2, CAS, CYP82X1, and CYP82X2 transcripts, but did not affect transcript levels of other tested genes (Fig. 8).

Expression of CYP82Y1 in Opium Poppy—Noscapine levels were highest in opium poppy stems, but the alkaloid was detected in all organs (Fig. 9A). Expression analysis by qRT-PCR also revealed the occurrence of CYP82Y1 transcripts in all plant organs, with the highest levels detected in stems (Fig. 9B).

DISCUSSION

CYP82Y1 was shown to catalyze the conversion of (R,S)-N-methylcanadine to (R,S)-1-hydroxy-N-methylcanadine and also accepted (S)-N-methyllystolipine as a substrate. Targeted metabolite and transcript profiling of eight opium poppy chemotypes that either accumulate or lack noscapine was used to identify CYP candidates potentially involved in phthalideisoquinoline alkaloid metabolism (Fig. 2). The same approach was used to isolate scoulerine 9-O-methyltransferase (SOMT1), canadine synthase (CYP719A21), and noscapine synthase (NOS) (Fig. 1), along with two other O-methyltransferases (SOMT2 and SOMT3) (5, 6, 11). CYP82Y1, CYP82X1, CYP82X2, and CYP719A21 transcripts were detected in 454 pyrosequencing databases of opium poppy chemotypes accumulating noscapine or narcotoline (Fig. 2). These CYPs were also reported as four of a ten-member genomic cluster consisting of biosynthetic genes absent in a noscapine-free opium poppy chemotype (10). The extended correlation between the occurrence of CYP transcripts corresponding to members of the ten-gene cluster and noscapine in several different opium poppy chemotypes suggests that the presence or absence of the
FIGURE 7. Virus-induced gene silencing confirms the involvement of CYP82Y1 in noscapine biosynthesis. A, region (gray line) of the CYP82Y1 cDNA inserted into the pTRV vector used to suppress CYP82Y1 transcript levels. The thick black line represents the coding region, whereas the flanking thin lines indicate noncoding 5′- and 3′-untranslated regions. Solid arrows show annealing sites of primers used to amplify cDNA fragments, whereas dotted arrows show annealing sites of primers used for qRT-PCR analysis. B, ethidium bromide-stained agarose gel showing the detection by reverse transcription-PCR of TRV coat protein transcripts in total RNA extracted from opium poppy plants infiltrated with A. tumefaciens cultures harboring pTRV1 and pTRV2 constructs. C, relative CYP82Y1 transcript abundance in control (pTRV2) and silenced (pTRV2–82Y1) plants. D, total ion chromatograms showing the major alkaloid profiles of control (pTRV2) and CYP82Y1-silenced (pTRV2–82Y1) plants. E, relative abundance of major alkaloids and other alkaloids affected by the suppression of CYP82Y1 transcript levels in control (pTRV2) and silenced (pTRV2–82Y1) plants. Asterisks represent significant differences determined using an unpaired, two-tailed Student t test (p < 0.05).
gene cluster determines the ability to produce phthalideisoquinoline alkaloids.

Three CYP families have been implicated in BIA metabolism (12). Whereas several enzymes in the CYP80 and CYP719 families have been shown to catalyze aromatic hydroxylation and inter- and intramolecular C-O and C-C coupling reactions involved in 1-benzylisoquinoline, protoberberine, aporphine, bisbenzylisoquinoline, and morphinan alkaloid biosynthesis, only two members in the CYP82 family have been functionally characterized: CYP82N4 catalyzes aliphatic hydroxylation at C14 of N-methylcanadine and N-methylstylopine, resulting in tautomerization to allocryptopine and protopine, respectively (20), whereas CYP82N2v2 from *E. californica* catalyzes aliphatic hydroxylation at C6 of protopine and allocryptopine leading to intramolecular rearrangement to the benzo[c]phenanthridine scaffold (21). CYP82Y1 displays considerable sequence identity with CYP82N4 (53%) and CYP82N2v2 (44%) and, similar to CYP82N4, also accepted only N-methylcanadine and N-methylstylopine as substrates. In contrast, CYP82Y1 and CYP82N4 catalyze regiospecific hydroxylation at C1 and C14, respectively. Also, CYP82N4 converted both N-methylstylopine and N-methylcanadine at similar rates (20), whereas PsCYP82Y1 converted N-methylcanadine with greater efficiency than N-methylstylopine. Alterations in the residues responsible for substrate recognition and alkaloid binding might be attributed to differences in substrate preference and oxidation activity among CYP82 variants. In contrast with the KPIAPXXXPH substrate recognition site motif of enzymes in the CYP719 family, CYP82Y1, CYP82X1, and CYP82X2 contain a distinct YPA(G/S)XXX(E/D)R domain and a con-
served Gly-370 residue. However, whereas the residues specific for alkaloid binding in N-methylstylopine 14-hydroxylase (CYP82N4) (20) are Ile and Leu, the corresponding residues in CYP82Y1 are Leu and Ser (6). CYP82N2v2 was relatively promiscuous with protopine alkaloid substrates, including protopine, allocryptopine, 13-oxoprotopine, and corycavine (21). Other substituted guateryn protoberberine alkaloids beyond N-methylcanadine and N-methylstylopine could also serve as substrates for CYP82Y1 and CYP82N4, although such reactions might not be physiologically relevant. A narrow substrate range is common among CYPs involved in the biosynthesis of BIAs (16–19) and other plant-specialized metabolites. For example, CYP82G1 from tobacco (N. tabacum), which catalyzes the final step in the biosynthesis of common homoterpenic volatiles, displays substrate specificity for (E,E)-geranylinalool and its C15 analog (E)-nerolidol (34), and CYP82C2 and CYP82C4 from mouse ear cress (A. thaliana) show substrate specificity for 8-methoxypсорalen (35).

A major unresolved aspect of noscapine biosynthesis concerns the transformation of the protoberberine to the secoberbine backbone as a component in the ultimate formation of the phthalideisoquinoline scaffold. On structural grounds, secobereine alkaloids, such as macrancalene and macrancaldine, were considered as possible intermediates between a protoberberine precursor, such as scoulerine or 1-methoxyscoulerine, and noscapine (36). An alternative proposal involves the stereospecific N-methylation of scoulerine followed by multi-stage oxidation to egenine, which is further oxidized to phthalideisoquinoline alkaloids, such as bicuculline (7). A protoberberine, such as scoulerine, has also been hypothesized to potentially undergo N-methylation and C8 oxidation to the secobерbine macranaldehyde, which could be reduced to macrantaline or oxidized to macrantoridine (7). Theoretically, macrantaldehyde could undergo oxidation to narcoticsinecacetal (37). As such, a hypothetical scheme for noscapine biosynthesis in opium poppy was proposed whereby N-methylcanadine is hydroxylated at C1 and possibly O-methylated to form 1-methoxy-N-methylcanadine (9). Oxidation of 1-methoxy-N-methylcanadine at C8 was seen as the entry point to the formation of a secoberbine intermediate, such as macranaldehyde. The drawback of this hypothesis is the lack of empirical evidence for the occurrence of secoberbine alkaloids in opium poppy. Most known secoberbine alkaloids have been isolated from different Papaver species such as Papaver pseudo-oriental, Papaver armeniacum, and Papaver fugax (Sariyar 2002). However, the proposed pathway extrapolated from the partial physiological characterization of a putative noscapine biosynthetic gene cluster supports the natural occurrence of secoberbine alkaloids in opium poppy (10). The individual suppression of transcript levels for six of ten clustered genes using VIGS was interpreted to suggest a biosynthetic route involving an initial ring opening (i.e., cleavage of the berberine bridge) on N-methylcanadine catalyzed by either CYP82X1 or CYP82Y1 and yielding a secoberbine intermediate. CYP82X2 was proposed to then catalyze hydroxylation at the C3 position, yielding a derivative of this secoberbine intermediate. The two schemes agree on the involvement of an enzymatic product derived from N-methylcanadine as the entry point to unidentified secoberbine intermediates in noscapine biosynthesis.

High resolution MS² and 1H NMR analyses showed unequivocally that the CYP82Y1 reaction product derived from N-methylcanadine was 1-hydroxy-N-methylcanadine. Since none of the three CYP82 candidates accepted canadine as a substrate, and because neither CYP82X1 nor CYP82X2 acted on N-methylcanadine, the conversion of N-methylcanadine to 1-hydroxy-N-methylcanadine by CYP82Y1 clearly represents the first committed step in noscapine biosynthesis and appears crucial to further elaboration of further pathway by other enzymes. CYP82X1 and CYP82X2 are likely responsible for cleavage of the N7/C8 bridge and hydroxylation of the C13 position. 1-Hydroxylation of N-methylcanadine distinguishes the biosynthesis of noscapine and narcotine in opium poppy from the formation of other phthalideisoquinoline alkaloids, such as hydastine and bicuculline which are found mostly in plants outside of the genus Papaver and lack a C4’ hydroxyl or methoxyl moiety. Should downstream enzymes in opium poppy depend on the C1 hydroxylation of N-methylcanadine in order to catalyze subsequent ring opening and lactone bridge conversions, the biosynthesis of compounds such as hydastine and bicuculline might involve relatively unrelated genes. It will be interesting to determine whether homologs of CYP82Y1 and genomic clusters of phthalideisoquinoline alkaloid biosynthetic genes occur in plants such as goldenseal (Hydrastis canadensis) and Corydalis chaerophyllea, which accumulate hydastine and bicuculline, respectively. Interestingly, the C4’ methoxyl group contributes to the potent pharmacological properties of noscapine compared with other phthalideisoquinolines.

The VIGS-mediated suppression of CYP82Y1 transcript levels resulted in the increased accumulation of several protoberberine alkaloids including the noscapine biosynthetic intermediates scoulerine, canadine, tetrahydrocolumamine, and N-methylcanadine (Fig. 1). However, the silencing of CYP82Y1 was also associated with a significant reduction in the accumulation of transcripts encoding enzymes operating, putatively in some cases, immediately upstream (SOMT1 and CYP719A21) and downstream (CYP82X1, CYP82X2, SOMT2, and an O-acetyltransferase, AT1) of 1-hydroxy-N-methylcanadine (Fig. 8). Co-suppression of several pathway genes hinders the utility of VIGS as a functional genomics tool to validate the physiologial role of CYP82Y1 in noscapine metabolism. Nevertheless, the co-silencing of only clustered genes suggests some form of coordinated regulation, although a mechanism whereby post-transcriptional suppression of CYP82Y1 would affect the expression of other genes, many with entirely unrelated nucleotide sequences, is not known. Frequent co-expression of biosynthetic genes involved in the same metabolic pathway has been proposed as a factor in co-suppression phenomena (38). A negative correlation between intergenic distance and co-expression has also been suggested (39). The concomitant accumulation of the structurally related protoberberine tetrahydrodopalmatine, which is not a noscapine biosynthetic intermediate, supports a divergence in alkaloid pathway flux when a bottleneck is introduced at (S)-N-methylcanadine. Interestingly, the expression of biosynthetic genes outside of the gene cluster (6OMT, 4’OMT2, BBE, and TNMT)
and two clustered genes encoding enzymes operating at the end of the noscapine pathway (CXEI and NOS) were not affected by the silencing of CYP82Y1. 6OMT (norccaurine 6-O-methyltransferase) and 4’OMT2 (3’-hydroxy-N-methylccaurine 4’-O-methyltransferase) catalyze early conversions in BIA metabolism (12). In contrast, VIGS-mediated silencing of NOS did not affect the expression of other genes, which is perhaps related to the localization of CXEI and NOS in different cell type compared with other enzymes encoded by genes in the cluster (11).

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REFERENCES
1. Anderson, T. (1862) On the chemistry of opium. J. Chem. Soc. Perkin Trans. I 15, 446–455
2. Battersby, A. R., Hirst, M., McCalind, D. J., Southgate, R., and Staunton, J. (1968) Alkaloid biosynthesis. XII. The biosynthesis of narcotine. J. Chem. Soc. Perkin Trans. I 17, 2163–2172
3. Kleinschmidt, G., and Mothes, K. (1959) Physiology and biosynthesis of alkaloids of Papaver somniferum. Z. Naturforsch. 14B, 52–56
4. Battersby, A. R., Staunton, J., Wiltshire, H. R., Bircher, B. J., and Fuganti, C. (1975) Studies of enzyme-mediated reactions. Part V. Synthesis of (13S)-[13-3H1]scoulerine from stereospecifically labelled (S)-[α-3H1] benzyl alcohols. Stereochemistry of enzymatic reactions at saturated benzylic carbon. J. Chem. Soc. Perkin Trans. I, 1162–1171
5. Dang, T. T., and Facchini, P. J. (2012) Characterization of three O-methyltransferases involved in noscapine biosynthesis in opium poppy. Plant Physiol. 159, 618–631
6. Dang, T. T., and Facchini, P. J. (2014) Cloning and characterization of canadine synthase involved in noscapine biosynthesis in opium. FEBS Lett. 588, 198–204
7. Saziyar, G., and Shamma, M. (1986) Six alkaloids from Papaver species. Phytochemistry 25, 2403–2406
8. Liscombe, D. K., and Facchini, P. J. (2007) Molecular cloning and characterization of tetrahydroprotoberberine cis-N-methyltransferase, an enzyme involved in alkaloid biosynthesis in opium poppy. J. Biol. Chem. 282, 14741–14751
9. Facchini, P. J., Hagel, J. M., Liscombe, D. K., Loukanina, N., MacLeod, B. P., Samanani, N., and Zulak, K. G. (2007) Opium poppy. Blueprint for an alkaloid factory. Phytochem. Rev. 6, 97–124
10. Winzer, T., Gazda, V., He, Z., Kaminski, F., Kern, M., Larson, T. R., Li, Y., Meade, F., Teodor, R., Vaistij, F. E., Walker, C., Bowser, T. A., and Graham, I. A. (2012) Papaver somniferum 10-gene cluster for synthesis of the anti-cancer alkaloid noscapine. Science 336, 1704–1708
11. Chen, X., and Facchini, P. J. (2014) Short-chain dehydrogenase/reductase catalyzing the final step of noscapine biosynthesis is localized to laticifers in opium poppy. Plant J. 70.10.1111/tpj.12379
12. Hegel, J. M., and Facchini, P. J. (2013) Benzylisoquinoline alkaloid metabolism. A century of discovery and a brave new world. Plant Cell Physiol. 54, 647–672
13. Nguyen, D. T., Göpfert, J. C., Ikezawa, N., MacNevin, G., Kathiresan, M., Conrad, J., Spring, O., Ro, D. K. (2010) Biochemical conservation and characterization of CYP80G2, a cytochrome P450 that catalyzes an intramolecular C-C phenol coupling of (S)-reticuline in magnoflorine bio-
synthesis, from cultured Coptis japonica cells. J. Biol. Chem. 283, 8810–8821
14. Díaz Chávez, M. L., Rolf, M., Gesell, A., and Kutchan, T. M. (2012) Characterization of two methylenedioxy bridge-forming cytochrome P450-dependent enzymes of alkaloid formation in the Mexican prickly poppy Argemone mexicana. Arch. Biochem. Biophys. 507, 186–193
15. Ikezawa, N., Tanaka, M., Nagayoshi, M., Shinkyo, R., Sakaki, T., Inouye, K., and Sato, F. (2003) Molecular cloning and characterization of CYP719, a methylenedioxy bridge-forming enzyme that belongs to a novel P450 family, from cultured Coptis japonica cells. J. Biol. Chem. 278, 38557–38565
16. Ikezawa, N., Iwasa, K., and Sato, F. (2007) Molecular cloning and characterization of methylenedioxy bridge-forming enzymes involved in stylopine biosynthesis in Eschscholzia californica. FEBS J. 274, 1019–1035
17. Gesell, A., Rolf, M., Ziegler, J., Díaz Chávez, M. L., Huang, F.-C., and Kutchan, T. M. (2009) CYP719Bl is a salutaridine synthase, the C-C phenol-coupling enzyme of morphine biosynthesis in opium poppy. J. Biol. Chem. 284, 24432–24442
18. Beaudoin, G. A., and Facchini, P. J. (2013) Isolation and characterization of a cDNA encoding (5S)-n-methylstyrlopin 14-hydroxylase from opium poppy, a key enzyme in sanguinarine biosynthesis. Biochem. Biophys. Res. Commun. 431, 597–603
19. Takemura, T., Ikezawa, N., Iwasa, K., and Sato, F. (2013) Molecular cloning and characterization of a cytochrome P450 in sanguinarine biosynthesis from Eschscholzia californica cells. Phytochemistry 91, 100–108
20. Desgagné-Penix, I., Khan, M. F., Schrierme, D. C., Cram, D., Nowak, J., and Facchini, P. J. (2011) Integration of deep transcriptome and proteome analyses reveals the components of alkaloid metabolism in opium poppy cell cultures. BMC Plant Biol. 10, 252
21. Hagel, J. M., and Facchini, P. J. (2010) Dioxygenases catalyze the O-de-methylation steps of morphine biosynthesis in opium poppy. Nat. Chem. Biol. 6, 273–275
22. Larkin, M. A., Blackshields, G., Brown, N. P., Chenna, R., McGettigan, P. A., McWilliam, H., Valentin, F., Wallace, I. M., Wilm, A., Lopez, R., Thompson, J. D., Gibson, T. J., and Higgins, D. G. (2007) Clustal W and Clustal X version 2.0. Bioinformatics 23, 2947–2948
23. Ro, D. K., Ouellet, M., Paradiso, E. M., Burd, H., Eng, D., Paddon, C. J., Newman, J. D., and Keasling, J. D. (2008) Induction of multiple pleiotropic drug resistance genes in yeast engineered to produce an increased level of anti-malarial drug precursor, artemisinic acid. BMC Biotechnol. 8, 83
24. Pauli, H. H., and Kutchan, T. M. (1998) Molecular cloning and functional heterologous expression of two alleles encoding (S)-n-methylccaurine 3’-hydroxylase (CYP80B1), a new methyl jasmonate-inducible cytochrome P450-dependent mono-oxygenase of benzylisoquinoline alkaloid biosynthesis. Plant J. 13, 793–801
25. Pompon, D., Louerat, B., Bronine, A., and Urban, P. (1996) Yeast expression of animal and plant P450s in optimized redox environments. Methods Enzymol. 272, 51–64
26. Dinesh-Kumar, S. P., Anandarakshmi, R., Marathe, R., Schiff, M., and Liu, Y. (2003) Virus-induced gene silencing. Methods Mol. Biol. 236, 287–294
27. Hileman, L. C., Drea, S., Martino, G., Litt, A., and Irish, V. F. (2005) Virus-induced gene silencing is an effective tool for assaying gene function in the basal eukaryotic species Papaver somniferum (opium poppy). Plant J. 44, 334–341
28. Desgagné-Penix, I., Farrow, S. C., Cram, D., Nowak, J., and Facchini, P. J. (2012) Integration of deep transcript and targeted metabolite profiling of eight cultivars of opium poppy. Plant Mol. Biol. 79, 295–313
29. Binutu, O. A., and Cordell, G. A. (2000) Constituents of Zanthoxylum sprucea. Pharm. Biol. 38, 210–213
30. Lee, J. K., Cho, J. G., Song, M. C., Yoo, J. S., Lee, D. Y., Yang, H. J., Han, K. M., Kim, D. H., Oh, Y. J., Jeong, T. S., and Baek, N. I. (2009) Isolation of isoquinoline alkaloids from tuber of Corydalis tartschaninovii and their inhibition activity on low density lipoprotein oxidation. J. Korean Soc. Appl. Biol. Chem. 52, 646–654
31. Le, P. M., McCooeye, M., and Windust, A. (2013) Characterization of the

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alkaloids in goldenseal (Hydrastis canadensis) root by high resolution Orbitrap LC-MS(n). Anal. Bioanal. Chem. 405, 4487–4498

34. Lee, S., Badieyan, S., Bevan, D. R., Herde, M., Gatz, C., and Tholl, D. (2010) Herbivore-induced and floral homoterpene volatiles are biosynthesized by a single P450 enzyme (CYP82G1) in Arabidopsis. Proc. Natl. Acad. Sci. U.S.A. 107, 21205–21210

35. Kruse, T., Ho, K., Yoo, H.-D., Johnson, T., Hippely, M., Park, J.-H., Flavell, R., and Bobzin, S. (2008) In planta biocatalysis screen of P450s identifies 8-methoxypsoralen as a substrate for the CYP82C subfamily, yielding original chemical structures. Chem. Biol. 15, 149–156

36. Sariyar, G., and Phillipson, D. (1977) Macrantaline and macrantoridine, new alkaloids from a Turkish sample of Papaver pseudo-orientale. Phytochemistry 16, 2009–2013

37. Sariyar, G. (2002) Biodiversity in the alkaloids of Turkish Papaver species. Pure Appl. Chem. 74, 557–574

38. Williams, E. I., and Bowles, D. I. (2004) Coexpression of neighboring genes in the genome of Arabidopsis thaliana. Genome Res. 14, 1060–1067

39. Michalak, P. (2008) Coexpression, coregulation, and cofunctionality of neighboring genes in eukaryotic genomes. Genomics 91, 243–248