JBC Papers in Press. Published on December 30, 2019 as Manuscript RA119.011547 The latest version is at http://www.jbc.org/cgi/doi/10.1074/jbc.RA119.011547 *O*-Methyltransferases from sacred lotus

Isolation and characterization of two *O*-methyltransferases involved in benzylisoquinoline alkaloids biosynthesis in sacred lotus (*Nelumbo nucifera*)

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ABSTRACT

Benzylisoquinoline alkaloids (BIAs) are a major class of plant metabolites with many pharmacological benefits. Sacred lotus (Nelumbo nucifera) is an ancient aquatic plant of medicinal value owing to antiviral and immunomodulatory activities linked to its constituent BIAs. Although more than 30 BIAs 1-benzylisoquinoline, belonging to the aporphine, and bisbenzylisoquinoline structural subclasses and displaying a predominant Renantiomeric conformation have been isolated from N. nucifera, its BIA biosynthetic genes and enzymes remain unknown. Herein, we report the isolation and biochemical characterization of two O-methyltransferases (OMTs) involved in BIA biosynthesis in sacred lotus. Five homologous genes, designated NnOMT1-5 and encoding polypeptides sharing > 40% amino acid sequence identity, were expressed in Escherichia coli. Functional characterization of the purified recombinant proteins revealed that NnOMT1 is a regiospecific 1-benzylisoquinoline 6-O-methyltransferase (6OMT) accepting both R and S substrates, whereas NnOMT5 is mainly 7-O-methyltransferase (7**O**MT), a with relatively minor 60MT activity and a strong stereospecific preference for S enantiomers. Available aporphines were not accepted as substrates by either enzyme, suggesting that O-

methylation precedes BIA formation from 1benzylisoquinoline intermediates. K_m values for NnOMT1 and NnOMT5 were 20 µM and 13 uM for (R,S)-norcoclaurine and (S)-Nmethylcoclaurine, respectively, similar to those for OMTs from other BIA-producing plants. Organ-based correlations of alkaloid content, OMT activity in crude extracts, and OMT gene expression supported physiological roles for NnOMT1 and NnOMT5 in BIA metabolism, occurring primarily in young leaves and embryos of sacred lotus. In summary, our work identifies two OMTs involved in BIA metabolism in the medicinal plant N. nucifera.

INTRODUCTION

Benzylisoquinoline alkaloids (BIAs) are a major class of plant specialized metabolites, many of which possess significant pharmacological value. The 1-benzylisoquinoline (S)-norcoclaurine has been recognized as the common intermediate to all biosynthetically characterized BIAs (Figure 1), enantioselective and is formed via the condensation of the tyrosine derivatives, dopamine and 4-hydroxyphenylacetaldehyde, catalyzed by norcoclaurine synthase (NCS) (1). Functional group modifications, especially O- and Nmethylations, of the 1-benzylisoquinoline scaffold combined with various C-C and C-O coupling reactions yield additional structural subclasses including aporphines (e.g. magnoflorine), bisbenzylisoquinolines (e.g. dauricine), pavine

(e.g. eschscholtzidine), protoberberines (e.g. berberine), morphinans (e.g. morphine), phthalideisoquinolines (e.g. noscapine), and benzo[c]phenanthridines (e.g. sanguinarine) (2).

The natural occurrence of BIAs is restricted to a relatively small number of plant families primarily in the order Ranunculales, which includes the opium poppy (Papaver somniferum), an extensively investigated model species that remains the sole commercial source for the narcotic analgesic morphine and the antitussive and potential anticancer drug noscapine (3). BIAs have also been detected in other plant including the Cornales, Laurales. orders Magnoliales, Piperales, Sapindales, and Proteales, although BIA metabolism in members of these taxa has not yet been empirically determined (4). Sacred lotus (Nelumbo nucifera Gaertn.; Figure 2a), is an aquatic basal eudicot in the order Proteales that has been cultivated as an edible crop for nearly seven millennia, and is highly valued in both Chinese and Ayurvedic traditional medicine for its healing properties (5). The pharmacological properties of sacred lotus are largely the result of the BIA constituents in the plant. For instance, 1benzylisoquinolines such as norcoclaurine and its 6-*O*-methylated derivative coclaurine have exhibited anti-HIV activity (6), whereas the 6.7-*O*,*O*- and *N*-methylated derivative armepavine has displayed promising immunomodulatory effects for the treatment of autoimmune diseases such as systemic lupus erythematosus (7). In addition, major aporphines (e.g. nuciferine) and bisbenzylisoquinolines (e.g. neferine) have shown potential for the treatment of lung, liver, ovarian and breast cancers (8-12). However, none of the biosynthetic genes and enzymes responsible for the BIA biosynthesis in sacred lotus have been characterized (13).

Based on the reported alkaloid profile in sacred lotus and well characterized pathways in opium poppy, BIA metabolism is predicted to begin with the formation of norcoclaurine and proceed via a limited number of enzyme types including one or more O-methyltransferases (OMTs), N-methyltransferases, and cytochrome P450 monooxygenases potentially belonging to and **CYP719** subfamilies CYP80 (13).Specifically, the formation of known 1benzylisoquinoline alkaloids in sacred lotus would involve only O-methylations and N-methylations

of norcoclaurine (**Figure 1**). Since *O*-methylation contributes substantially to the diversity of BIAs in sacred lotus, we targeted the isolation and functional characterization of OMTs.

Methylations play a central role in the functionalization of specialized metabolites (14). In particular, OMTs involved in BIAs metabolism catalyze methyl transfer to the hydroxyl group of an alkaloid substrate, using S-adenosyl-Lmethionine (SAM) as the methyl donor group and proceeding via a S_N2-like nucleophilic attack vielding a methylated alkaloid product and Sadenosyl-L-homocysteine (SAH) (15). OMTs are typically homodimeric or, as recently discovered, heterodimeric enzymes (16,17). Within each monomer, the C-terminal domain adopts a canonical Rossmann fold (i.e. a β -strand sandwiched by two α -helices) and plays a fundamental role in substrate binding and catalysis, whereas the N-terminal domain is of central importance in dimerization (15,18-20).

In opium poppy, the methylation of (S)norcoclaurine to (S)-coclaurine by (R,S)norcoclaurine 6-O-methyltransferase (Ps6OMT) represents the first tailoring reaction in BIA metabolism (21,22). Recent crystallographic studies on 6OMT from meadow rue (Thalictrum flavum, Tf6OMT) have revealed key determinants for substrate recognition and turnover (18). In opium poppy, at least six other OMTs involved in BIA biosynthesis have been functionally characterized. including (R,S)-3'-hydroxy-Nmethylcoclaurine 4'-O-methyltransferase 2 (*R*,*S*)-reticuline (Ps4'OMT2), 7-0methyltransferase (Ps7OMT) and norreticuline 7-O-methyltransferase (PsN7OMT), all involved in 1-benzylisoquinoline biosynthesis (i.e. reticuline, laudanine and norlaudanine), as well as 4'-Odesmethyl-3-O-acetylpapaveroxine 4'-0methyltransferase (PsOMT2:PsOMT3 and PsOMT2:Ps6OMT heterodimers) and scoulerine 9-*O*-methyltransferase (PsSOMT1), which function in phthalideisoquinoline (e.g. noscapine) metabolism (16,17,19,21-26). Although the isolation of a specific 3'OMT has been elusive, low 3'-O-methyltransferase activity has been detected for PsSOMT1, as well as for SOMTs from related species such as yellow horned poppy (Glaucium flavum, GfOMT6) and California poppy (Eschscholzia californica, EcSOMT) (26-28). Finally, a columbamine *O*-methyltransferase

has been isolated from Japanese goldthread (*Coptis japonica*; CjCoOMT) (29). Although several functionally related OMTs have been investigated, these enzymes have been isolated from fewer than ten species all from the Ranunculales (20).

Sacred lotus accumulates 1benzylisoquinolines, and derived aporphines (via C8–C2' intramolecular coupling) and bisbenzylisoquinolines (via C8-C3'/5' and C7-O-C3'/5' intermolecular coupling) (5), most of which are O-methylated at C6, C7, and/or C4' (13). The availability of a sacred lotus draft genome (30,31) greatly facilitates gene mining for candidate OMTs based on amino acid sequence similarity with respect to characterized O-methyltransferases (32-34). However, OMT candidates have so far been investigated only in terms of gene expression, with no functional characterization of the encoded proteins. In addition, enzyme stereospecificity is an intriguing feature of BIA metabolism in sacred lotus, since norcoclaurine has been isolated from the plant as both R and S enantiomers, and other BIAs, including coclaurine, norarmepavine and Nmethylcoclaurine have been exclusively reported as the R conformer (13), in contrasts with the predominantly S stereochemistry of most BIAs in opium poppy and related plants (3). Herein we report the identification and characterization of Omethyltransferases involved in BIA metabolism in sacred lotus.

RESULTS

Sacred lotus BIA profile

Alkaloids corresponding to the leaf (folded and unfolded developmental stages), rhizome, root and embryo (lotus plumule) of two Nelumbo nucifera cultivars, referred as Pink and White owing to their distinct floral coloration, were extracted and analyzed by mass spectrometry (MS) (Figure 2b,c, Table S1). Fifteen alkaloids (i.e. six 1benzylisoquinolines, four aporphines, one proaporphine and four bisbenzylisoquinolines) were identified based on their retention times and dissociation collision-induced (CID) spectra compared with available authentic standards or previously published data. 1-Benzylisoquinoline and aporphine alkaloids were prevalent in the leaves, whereas bisbenzylisoquinolines were most

abundant in the embryos. In general, the rhizome and roots contained only low BIA levels, although no bisbenzylisoquinolines were detected. Notably, the structural isomers N-methylisococlaurine and norarmepavine were exclusively detected in the Pink and White varieties, respectively. The bisbenzylisoquinolines liensinine and neferine were found only in the White variety, whereas nelumboferine and isoliensinine were the only bisbenzylisoquinolines identified in Pink and accumulated at 20- (in folded leaves) and 4-fold (in embryos) higher levels compared with the White variety. In addition, alkaloids were detected in the leaves (norarmepavine, nelumboferine and isoliensinine), rhizome (norcoclaurine, coclaurine, *N*-methylcoclaurine, norarmepavine, Nmethylisococlaurine, anonaine, roemerine, Nnornuciferine, O-nornuciferine, nuciferine and pronuciferine), roots (coclaurine. Nmethylcoclaurine, norarmepavine, Nmethylisococlaurine, armepavine, anonaine. roemerine, N-nornuciferine and O-nornuciferine) and embryo (norarmepavine and O-nornuciferine) that had not been previously reported (5.13). Overall, the alkaloid profile found for both varieties supports the occurrence of 6-O-, 7-O- and 4'-O-methyltransferases in sacred lotus.

NnOMT identification and phylogeny

Sacred lotus transcripts encoding five putative OMTs (NnOMT1-NnOMT5) were identified based on amino acid sequence identity with opium OMTs (Table S2). The predicted poppy NnOMT1-NnOMT5 translation products displayed predicted molecular weights of 38.4, 38.7, 38.8, 39.2 and 38.1 kDa and expected isoelectric points of 6.5, 7.8, 6.1, 5.7 and 5.6, respectively. Four sacred lotus OMTs shared ~80% amino acid sequence identity, whereas NnOMT5 exhibited <50% identity with respect to the others. NnOMT1 showed ~70% amino acid sequence identity compared with functionally characterized 60MTs, whereas all candidates shared ~40-60% sequence identity with previously reported 70MTs and 4'OMTs (Table S3). Phylogenetic relationships among NnOMTs and functionally characterized BIA O-methyltransferases placed NnOMT1-NnOMT4 in a single clade related to several characterized 6OMTs, whereas NnOMT5 emerged on a distal clade that also contained CjCoOMT (Figure 3).

Amino acid sequence alignment of the functionally characterized Tf6OMT and sacred lotus OMT candidates showed a high conservation of key amino acid residues, including the catalytic determinants H²⁵⁶, D²⁵⁷ and E³¹⁵ (18), except for NnOMT4, in which case D²⁵⁷ was substituted with a Y residue (Figure 4). Amino acids implicated in BIA substrate binding (G¹⁶⁵, D¹⁶⁹, C²⁵³ and D³⁰⁶) were also conserved among sacred lotus OMT candidates except for G165V and C253W substitutions in NnOMT2, G165A in NnOMT3, G165S in NnOMT4, and G165A, D169H and C253S in NnOMT5. The main residues involved in SAM binding including T^{170} , G^{195} (motif I), D^{218} (motif II), D^{238} (motif III), and K^{252} (motif IV) were conserved across sacred lotus OMT candidates, excluding D²¹⁸ (substituted by G in NnOMT4) and D^{238} (substituted by N in NnOMT3). Two methionine residues M¹⁶⁶ and M³⁰⁷ assisting in the isoquinoline placing through sulfur-aromatic interactions were also conserved in lotus OMT candidates. The aromatic residues W¹⁴⁹ and F¹⁶², which interact with both SAM and BIA substrates, were mutated, W¹⁴⁹ to C, S and I in NnOMT3, NnOMT4 and NnOMT5, respectively, and F¹⁶² to L in NnOMT5. The conserved GxGxGx sequence in motif I. a hallmark of SAMdependent methyltransferases (14), was fully maintained in all the candidates except NnOMT5, in which the middle G residue was mutated to C.

NnOMT purification and characterization

Sacred lotus recombinant His₆-tagged OMTs produced in Escherichia coli were purified using cobalt-affinity chromatography and detected on an immunoblot using an anti-His₆ antibody (Figure S1). The molecular weights of the purified proteins and the predicted translation products were similar. Recombinant proteins were initially screened for enzymatic activity using 15 potential substrates representing BIA the 1benzylisoquinoline, aporphine, protoberberine and morphinan structural subclasses. The potential 1benzylisoquinoline substrates differentially contained free hydroxyl groups at C6, C7, C3' and/or C4', and both R and S enantiomers were in some cases available (Table S4). Reaction products showed an increase of 14 Da with respect

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to the corresponding substrate and were detected in reactions containing NnOMT1 or NnOMT5 with various 1-benzylisoquinoline incubated substrates (Figure 5, Figure S2). (R,S)-Norcoclaurine and (S)-norlaudanosoline were Omethylated by both NnOMT1 and NnOMT5, whereas (R)-norlaudanosoline was only accepted by NnOMT1. In addition, NnOMT5 catalyzed the *O*-methylation of (S)-coclaurine. (S)-Nmethylcoclaurine and (S)-reticuline with similar efficiencies, but (R)-reticuline was not accepted. Single peaks were detected for all the reactions, indicating the formation of only one product.

Reaction products corresponding to the 6-*O*-methylation norcoclaurine of and norlaudanosoline (coclaurine [m/z 286] and 6-Omethylnorlaudanosoline [m/z 302], respectively) and the 7-O-methylation of N-methylcoclaurine (armepavine [m/z 314]), were unambiguously identified by comparison with available authentic standards. Enzyme assays with coclaurine and reticuline were subjected to positive-mode electrospray ionization (ESI[+]) high performance liquid chromatography (HPLC)-tandem mass spectrometry (MS/MS) for the accurate characterization of reaction products by CID fragmentation analysis to determine whether Omethylation was associated with the isoquinoline (C7) or benzyl moiety of coclaurine (C4') and reticuline (C3') (Figure S3). ESI[+]-CID of 1benzylisoquinolines at low ionization energy yielded isoquinoline and/or benzyl moieties as major ion fragments; thus, the addition of 14 Da to either fragment with respect to the parent ion revealed the regiospecificity of the reaction. In this regard, coclaurine showed two main fragments at m/z 178 and m/z 107, corresponding to the isoquinoline and benzyl moieties, respectively. After incubation with NnOMT5, the m/z 107 fragment was retained in the reaction product, indicating that the methyl group was added to the isoquinoline moietv (i.e. 7-O-methylation) resulting in the formation of norarmepavine, an alkaloid found in sacred lotus (the other possible reaction product, 4'-O-methylcoclaurine, has not been reported in this plant). Similarly, the reaction product of NnOMT5 incubated with (S)-reticuline showed a mass increase of the isoquinoline moiety from m/z 192 to m/z 206, corresponding to the 7-O-methylated product laudanine, whereas the mass of the benzyl moiety (m/z 137) remained

unchanged. It is notable that neither reticuline nor laudanine have been reported in sacred lotus. 4'OMT activity was not detected with any combination of tested substrates and OMT candidates.

NnOMT1 displayed strict regiospecificity with respect to the C6 hydroxyl and showed similar turnover using racemic norcoclaurine and individual enantiomers of norlaudanosoline, the 3'hydroxylated derivative of norcoclaurine (**Figure 5**). NnOMT5 non-preferentially methylated the C7 hydroxyl of *N*-methylated (*N*-methylcoclaurine and reticuline) or *N*-demethylated (coclaurine) substrates, and also acted on the C6 hydroxyl of (*R*,*S*)-norcoclaurine and (*S*)-norlaudanosoline. The 6-*O*-methylation activity of NnOMT5 was not accompanied by secondary methylations (an increase of 28 Da). Contrary to NnOMT1, NnOMT5 showed a strict stereospecificity for the substrates in the *S* conformation.

Both enzymes showed a pH optimum of 8.0 using (R,S)-norcoclaurine (NnOMT1) or (S)-Nmethylcoclaurine (NnOMT5) as substrates and maximum catalytic activity at 30 °C and 37 °C, respectively (Figure S4). NnOMT1 and NnOMT5 displayed $K_{\rm m}$ values of 20 μ M and 152 μ M for (R,S)-norcoclaurine, respectively, whereas NnOMT5 exhibited a K_m value of 13 μ M and a K_i of 870 µM for (S)-N-methylcoclaurine (Table 1, Figure S5). Catalytic efficiencies (k_{cat}/K_m) were similar for the 6OMT activity of NnOMT1 and the 70MT activity of NnOMT5. The 70MT activity of NnOMT5 was 50-fold higher than the corresponding 6OMT activity using (R,S)norcoclaurine as the substrate.

NnOMT and BIA correlations in the plant

Alkaloid content, *O*-methyltransferase activity and *OMT* gene expression were measured in leaves, rhizome, roots and embryos for two sacred lotus varieties. Alkaloid content was determined for norcoclaurine, coclaurine, *N*-methylcoclaurine and armepavine, for all of which authentic standards were available (**Figure S6**). In both the Pink and White varieties, 1-benzylisoquinoline intermediates were predominantly detected in the leaves (**Figure 6a, Table S5**). In the White variety, norcoclaurine was the most abundant alkaloid, with folded leaves accumulating 7-, 70- and 1400-fold higher levels compared with the

embryo, rhizome and root, respectively. The Pink cultivar was richer in *N*-methylcoclaurine, with the folded leaves containing 3-, 40- and 90- fold higher levels than the rhizome, root and embryo, respectively.

Crude plant protein extracts incubated with (R,S)-norcoclaurine, (S)-N-methylcoclaurine or (R)-armepavine revealed similarities between the Pink and White varieties (**Figure 6b, Table S6**). 6OMT activity was most abundant in folded leaves and embryos, whereas 7OMT activity was predominant in the unfolded leaves. 6OMT activity was 5- and 4-fold higher than 7OMT activity in folded leaves and embryos, respectively. 7OMT activity was 8-fold higher than 6OMT in the White variety, whereas 6OMT activity was not detected in unfolded leaves of the Pink variety. Rhizomes and roots showed only low levels of 6OMT and 7OMT activities. 4'OMT activity was not detected.

Gene expression profiles were similar for both the Pink and White varieties (**Figure 6c**, **Table S7**). *NnOMT1* and *NnOMT5* transcripts were most abundant in leaves and were 3- to 30fold higher compared with other *OMT* genes. The levels of all five *NnOMT* transcripts were similar in the embryos of both varieties.

DISCUSSION

We have shown that two cDNAs from sacred lotus efficiently catalyze the regiospecific 6-Omethylation of norcoclaurine to coclaurine (NnOMT1) and 7-O-methylation of coclaurine and N-methylcoclaurine to norarmepavine and armepavine, respectively (NnOMT5). NnOMT1 was not stereospecific whereas NnOMT5 accepted only S enantiomers. Correlation between transcript levels, enzyme activity and alkaloid content supported a physiological role for NnOMT1 and NnOMT5 in BIA metabolism.

Despite sharing an apparently common biosynthetic pathway to *N*-methylcoclaurine (**Figure 1**), sacred lotus and opium poppy exhibit substantial differences in their alkaloid profiles. In this regard, whereas reticuline constitutes a key branch-point intermediate in the biosynthesis of most BIAs in opium poppy (3), sacred lotus does not accumulate reticuline and other 1benzylisoquinolines containing a C3' functional group, or reticuline-derived alkaloids including phthalideisoquinolines, benzo[c]phenanthridines, protoberberines and morphinans, which occur in members of the Ranunculales (13). In contrast, aporphines and bisbenzylisoquinolines are the major BIAs in sacred lotus, although alkaloid profiles vary among nearly 600 known varieties, across different plant organs and and developmental stages (32-37). In addition, sacred lotus aporphine alkaloids do not display substitutions in the benzyl moiety (13), in contrast with the reticuline-derived aporphines (e.g. magnoflorine, corytuberine and glaucine) found in members of the Ranunculales (27,38,39).

The sacred lotus BIA profile showed a preferential accumulation of 1-benzylisoquinolines aporphines in leaves. and and bisbenzylisoquinolines in embryos (Figure 2b), in agreement with previous investigations (36.37.40). Alkaloids containing 6-O-, 7-O- and/or 4'-Omethylations were detected in Pink and White varieties, supporting the search for specific Omethyltransferase activities in the plant. OMT gene candidates have been predicted in sacred lotus based on sequence similarity to functionally characterized OMTs from BIA-accumulating species (32-34), but the purported catalytic activities has not been assessed. We show that two of these candidates. NnOMT1 and NnOMT5. catalvze the *O*-methylation of 1benzylisoquinolines.

Fifteen BIA substrates possessing diverse structural backbones were screened, including 1benzylisoquinolines with a free hydroxyl at C6, C7 and C4'. As described for several 6OMTs from the Ranunculales (41,42), NnOMT1 displayed strict C6 regiospecificity and accepted both R and S 1-(Figure benzylisoquinoline conformers 5). NnOMT1 contains all conserved catalytic and substrate-binding residues occurring in Tf6OMT (Figure 4). In Tf6OMT, the O-methyltransfer H^{256} base-assisted reaction occurs via deprotonation of the substrate C6 hydroxyl, followed by a nucleophilic attack on the SAM methyl group. Interactions of H^{256} and D^{257} with the C6 hydroxyl group provide proper orientation and E³¹⁵ contributes to the strong basicity in the H²⁵⁶ NE2 required for catalysis (18). Since NnOMT1 and Tf6OMT possess equivalent key catalytic residues (H $^{256},$ \dot{D}^{257} and $E^{315})$ and share ~70% amino acid sequence identity, a similar catalytic mechanism is expected for NnOMT1.

unique sacred lotus OMT, catalyzed the methyltransfer to the C6 hydroxyl of (R,S)norcoclaurine and (S)-norlaudanosoline, but more efficiently to the C7 hydroxyl of (S)-coclaurine. (S)-N-methylcoclaurine and (S)-reticuline (Table 1). Despite the closest phylogenetic relationship between NnOMT5 and CiCoOMT, their substrate ranges are different. NnOMT5 did not accept the tested protoberberines (Table S4), whereas CjCoOMT preferred columbamine and other protoberberines, but did not accept the 1benzylisoquinolines norlaudanosoline, coclaurine and reticuline (29). NnOMT5 also showed a close relationship with a sister clade containing PsOMT2, which together with PsOMT3 or Ps6OMT forms the heterodimeric enzyme capable of O-methylate a seco-berbine intermediate in noscapine biosynthesis (16,17). Notably, this sister clade also included Coptis chinensis 60MT1 (Cc6OMT1), which was reported to catalyze either 6-O- or 7-O-methylation of (S)-norcoclaurine (43). Dual 6-O- and 7-O-methyltransferase activity has also been reported for Cc6OMT2, along with enzymes from Coptis teeta (Ct7OMT), P. somniferum (Ps6OMT and Ps7OMT) and G. flavum (GfOMT1) (22,27,43).

NnOMT5, the most phylogenetically

Regardless of a nomenclature suggestive of regiospecific functions. OMTs involved in BIA metabolism are generally promiscuous and exhibit apparent redundancy in most plants (22,26-28,43,44). In Tf6OMT co-crystalized with (S)norlaudanosoline the C7 hydroxyl group of the BIA substrate was found in close proximity to the catalytic H²⁵⁶, suggesting that minor adjustments in substrate positioning could lead to productive 7-O-methyltransfer (18) and the observed dual 6-Oand 7-O-methyltransferase activity of some OMTs. Based on the available crystal structure for Tf6OMT (PDB 5ICE), we constructed a homology model for NnOMT5 (Figure S7). In the resulting structure, the key catalytic histidine (H²⁵¹ in NnOMT5) involved in hydrogen bonding with the target C6-OH (3.0 Å), is also near the C7-OH (3.1 Å). Analysis of residues likely involved in BIA substrate binding showed that G165A and C253S mutations did not have a significant impact (i.e. the main chain carbonyl involved in hydrogen bond formation with the BIA substrate C3' and C6 hydroxyl groups, respectively, remained in a similar position) in agreement with the observed variability of these residues in other 70MTs (i.e. G165A in Ps70MT, PsN70MT and Ct70MT, C253W in Ps70MT and Ct70MT, C253N in Ps70MT, and C253G in Ec70MT). Conversely, the D169H mutation disrupted a hydrogen bond with the C4'-OH (2.6 Å), which stabilizes the substrate in the active site. Interestingly, whereas D^{169} is a conserved residue across all 60MTs, or is replaced by E in 4'OMTs, no consensus was found in functionally characterized 70MTs (i.e. D in PsN70MT, S in Ps70MT and Ec70MT, and T in Ct70MT); thus, this residue might not play a significant role in 70MT regiospecificity.

Interestingly, when both C6 and C7 hydroxyl groups were available, NnOMT5 exclusively catalyzed 6-O-methylation (Figure S2). Similar results were reported for E. californica and C. japonica 4'OMTs, which norcoclaurine catalvze 6-O-methyltransfer, although with lower efficiency than 4'-Omethylation on already 6-O-methylated substrates (45), in support of the importance of 6-Omethylation as a purported rate-limiting step in BIAs metabolism (21). However, since NnOMT5 7-O-methylation catalyzed of (S)-Nmethylcoclaurine 50-fold more efficiently than 6-*O*-methylation of (R,S)-norcoclaurine, the physiological relevance of the latter activity is questionable compared with NnOMT1 (Table 1). It should be noted that, although coclaurine and Nmethylcoclaurine are almost certainly physiological substrates of NnOMT5 (Figure 1), other reported 1-benzylisoquinolines such as Nmethylisococlaurine and lotusine contain 7-Omethyl but C6 hydroxyl groups, which does not agree with the inability of NnOMT5 to 7-Omethylate norcoclaurine in vitro. In this regard, it is feasible that norcoclaurine N-methylation should occur first for NnOMT5 to catalyze 7-O- rather than 6-O-methylation, however the required substrate to test this possibility was not available.

NnOMT5 also preferred norcoclaurine over norlaudanosoline (**Figure 5**), whereas NnOMT1 showed no substrate preference. Nonetheless, the main catalytic activity detected for NnOMT5 was the 7-*O*-methylation of several *N*-methylated and *N*-demethylated substrates (**Figure 5**). Although our understanding of 7OMT substrate recognition and catalysis is limited, it has been suggested that the conserved residue D³⁰⁶ in PsN7OMT is able to hydrogen bond with the nitrogen of its only substrate norreticuline, whereas Ps7OMT and Ec7OMT contain smaller uncharged residues at equivalent positions presumably to ease steric hindrance with (R,S)reticuline and other *N*-methylated substrates (22,23,44), and Ct7OMT, with a D³⁰⁶ to S substitution, is able to 7-*O*-methylate (*S*)norcoclaurine (43). Notably, NnOMT5 contains a D³⁰⁶ residue (**Figure S7, Figure S8**), but catalyzes similarly efficient 7-*O*-methylation of *N*methylated and *N*-demethyl substrates (**Figure 5**).

The catalytic activity of recombinants NnOMT1 and NnOMT5 with (R,S)-norcoclaurine (S)-N-methylcoclaurine, and respectively, exhibited similar reaction efficiency (k_{cat}/K_m) , whereas NnOMT5 showed a 50-fold more efficient 7-O- versus 6-O-methylation (Table 1). The $K_{\rm m}$ of NnOMT1 with (*R*,*S*)-norcoclaurine (20 µM) is similar to that reported for Ps6OMT (10 μ M) and Tf6OMT (52 μ M) (22,41), whilst the K_m of NnOMT5 (152 µM) is substantially higher, suggesting that NnOMT1 is the main 6OMT in sacred lotus. On the other hand, the $K_{\rm m}$ of NnOMT5 with (S)-N-methylcoclaurine (13 μ M) is substantially lower compared with that for (R,S)norcoclaurine (152 µM), but similar to that reported for Ps7OMT with (S)-reticuline (16 µM) (22) and PsN7OMT with (S)-norreticuline (44 µM) (23), suggesting that NnOMT5 acts primarily as a 70MT in the plant.

The apparent lack of recombinant NnOMT (Figure 5, Figure S2) or plant crude extract activity (Figure 6b, Table S6) on substrates 3'- or 4'-hydroxyl groups partially agrees with the sacred lotus BIA profile, which lacks C3'-substituted alkaloids (13). In contrast, 4'-O-methylated BIAs were detected (Figure 2b), including isoliensinine and neferine (Figure 2c). Low 4'OMT activity has been reported for enzymes with primarily 6-O- or 7-O-methyltransferase activity, such as GfOMT2 and Ps7OMT (22,27). However, NnOMT1 and NnOMT5 did not show 4'-O-methyltransferase activity on any substrate, including (R)armepavine, although the actual substrate(s) might not have been available. The phylogenetic relationships between NnOMT5 and PsOMT2, and between NnOMT1 and PsOMT3 and Ps6OMT, suggest that 4'OMT activity in sacred lotus could also involve a heterodimeric enzyme. It is notable that in the Ranunculales, a Berberis koetineana 4'OMT (Bk4'OMT), Cj4'OMT, Ec4'OMT and

Ps4'OMT2 act only on 3'-hydroxylated substrates (22,25,27,42,45,46), suggesting that sacred lotus 4'-*O*-methyltransferase activity could involve a unique enzyme.

Assayed aporphine, protoberberine and morphinan alkaloids were not accepted as substrates (Table S4). OMTs shown to Omethylate 1-benzylisoquinolines and protoberberines occur in species accumulating both alkaloid types (26-28), although specificity for 1-benzylisoquinolines (e.g. Cj6OMT and PsN7OMT) has also been reported (23,42). Though sacred lotus does not accumulate morphinans or protoberberines (13), aporphines are major alkaloids in the leaves (34,35). The lack of NnOMT activity on aporphines suggests that the C8-C2' coupling yielding the aporphine scaffold occurs after the O-methylation of 1intermediates. benzylisoquinoline However. Dactylicapnos scandens 70MT was recently reported to *O*-methylate corvtuberine. demonstrating the plausibility of aporphines as OMTs substrates (47). It is notable that the aporphines and bisbenzylisoquinolines of sacred lotus were not available; thus, it remains possible that certain intermediates within these BIA structural types are NnOMT substrates.

The occurrence of most BIAs as Rconformers is another unique feature in sacred lotus compared with members of the Ranunculales (13). The importance of the stereochemistry of 1benzylisoquinoline substrates is evident in opium poppy, in which reticuline conformation determines the formation of major end-point alkaloids, with morphine derived from (R)reticuline, and other alkaloids such as noscapine and sanguinarine produced from (S)-reticuline. BIA stereochemistry is initially established by the stereoselectivity of norcoclaurine synthase (NCS). Although all characterized NCS isoforms from the Ranunculales yield exclusively (S)-norcoclaurine (48-50), norcoclaurine in sacred lotus has been detected as both S and R enantiomers (6,51,52) and consequently, the presence of both enantiomeric forms of norcoclaurine-derived alkaloids depends on the stereospecificity of downstream enzymes. We show that NnOMT1 catalyzed the 6-Omethylation of (R)- and (S)-norlaudanosoline with similar turnover efficiency, whereas NnOMT5 accepted only (S)-norlaudanosoline and (S)reticuline (Figure 5, Table S4). Previously

characterized OMTs have either shown a lack of stereospecificity (e.g. Cj6OMT, Ps6OMT and Ps7OMT) or strict *S* stereospecificity (Bk4'OMT) (22,42,53). Our results support the formation of (R)-coclaurine and (S)-armepavine as previously reported in lotus (6,54). However, despite NnOMT5 showed strict stereospecificity toward Sneither norlaudanosoline enantiomers. nor reticuline are physiologically relevant substrates and it is possible that the enzyme catalyzes the of (*R*)-coclaurine and turnover (R)-Nmethylcoclaurine in the formation of the reported compounds (*R*)-norarmepavine and (R)armepavine, respectively (55). Unfortunately, none of the relevant R enantiomers were available as potential substrates.

Previous work on the BIA profile in sacred lotus focused on the quantification of aporphines and bisbenzylisoquinolines in leaves and embryos, respectively, but limited information has been provided for 1-benzylisquinoline intermediates (34,36). We quantified key alkaloids produced in the early steps of the pathway (i.e. norcoclaurine, coclaurine, N-methylcoclaurine and armepavine) in different sacred lotus organs (Figure 6a). Although 1-benzylisoquinolines were predominant in leaves, the detected compounds were likely intermediates leading to aporphine and bisbenzylisoquinoline end products. The low levels of 1-benzylisoquinolines in embryos could also reflect their transformation to predominant bisbenzylisoquinolines (36,37,40,56).

Assays containing crude plant extracts (R,S)-norcoclaurine, incubated with (S)-Nmethylcoclaurine and (R)-armepavine showed 6-O- and 7-O-methyltransferase activity in all organs (Figure 6b), but mostly in leaves and embryos. *OMT* transcripts were also expressed in all organs, although NnOMT1 and NnOMT5 showed higher expression levels, mainly in leaves (Figure 6c). These results are consistent with a recent report on *NnOMT* expression at different leaf developmental stages, whereby NnOMT1 and NnOMT5 transcript levels were up to 10-fold higher compared with NnOMT2-NnOMT4 (34). A lag between NnOMT expression and aporphine alkaloid accumulation was also noted in this study, which is potentially associated with the involvement of OMTs early in BIA metabolism and a requirement for the expression of additional genes to complete aporphine and bisbenzylisoquinoline biosynthesis.

It is notable that since this earlier study did not perform a functional characterization of candidate genes, NnOMT5 was reported as a 4'OMT (34). In agreement with the suggested order of *O*methylations in other plants (21,45), NnOMT1 appears to act earlier than NnOMT5 in the leaves since 6OMT activity decreased in unfolded leaves whereas 7OMT activity increased (**Figure 6b**). Similarly, *NnOMT1* transcript levels were higher in folded leaves than unfolded leaves, whereas *NnOMT5* transcript levels remained constant (**Figure 6c**), consistent with a previous report (34).

Although NnOMT1 and NnOMT5 transcript levels were 5- to 15-fold lower in embryos compared with folded leaves, specific 6and 7-O-methyltransferase activities were similar in both organs, likely owing to the prevalence of photosynthetic leaf proteins. Interestingly, the transcripts encoding CYP80G. levels of purportedly involved in aporphine biosynthesis, in leaves were 5 times higher than levels of transcripts encoding CYP80A, potentially involved in bisbenzylisoquinoline biosynthesis, in embryos (33), suggesting that lower biosynthetic gene transcript levels in embryos might be sufficient in this organ. However, neither CYP80G or CYP80A orthologs in sacred lotus were functionally characterized, so it is not certain that these transcripts are relevant to BIA metabolism. Bisbenzylisoquinolines biosynthesis has been proposed to occur in leaves, with alkaloids subsequently transported to embryos (36). NCS activity (4) and purported NnNCS transcripts (57) were detected in leaves, suggesting that BIA biosynthesis occurs in this organ. The detection of *O*-methyltransferase activity indicates that embryos are also involved in alkaloid biosynthesis.

From an evolutionary perspective, it is not known whether OMT recruitment occurred repeatedly or if an ancestrally promiscuous OMT gave rise to functionally distinct enzymes (20). The 6-O-methyltransferase activity of NnOMT1 was correctly predicted based on similarity to Tf6OMT and other 6OMTs. However, the 7-Omethyltransferase activity of NnOMT5 was unexpected owing to its relatively low similarity to characterized 70MTs from the Ranunculales (Figure 3, Table S3), suggesting a possible independent recruitment in sacred lotus. The characterization of other enzymes (e.g. norcoclaurine coclaurine Nsynthase,

methyltransferase, and aporphine and bisbenzylisoquinoline synthases) will provide additional key insights into the evolutionary history of BIA biosynthesis in sacred lotus.

EXPERIMENTAL PROCEDURES

Plant material

Seeds for two sacred lotus (*Nelumbo nucifera*) varieties, named Pink and White owing to their distinct flower colors, were germinated in water according to the instructions of the vendor (<u>www.rarexoticseeds.com</u>). After the emergence of embryogenic leaves, the seedlings were planted in pots containing heavy loam soil and submerged in an artificial pond. Plants were grown at 30 °C and under high-intensity lighting with a 16 h photoperiod. Alternatively, seeds were carefully opened with a bench press and the embryos were isolated. All harvested plant samples were immediately frozen in liquid nitrogen and stored at -80 °C until used.

Chemicals and reagents

(R,S)-Norcoclaurine, (R)-norlaudanosoline, (S)norlaudanosoline, (S)-coclaurine and (13S, 14R)-1,13-dihydroxy-N-methylcanadine were purchased from Toronto Research Chemicals (ON, Canada); (R)-reticuline was purchased from Santa Cruz Biotechnology (CA, USA); (R)-armepavine was purchased from MuseChem (NJ, USA); (S)scoulerine was purchased from ChromaDex (CA, USA); (+)-bulbocapnine and (+)-isocorvdine were purchased from Sequoia Research Products (Pangbourne, UK); and boldine and S-adenosyl-Lmethionine (SAM) were purchased from Sigma-Aldrich. (S)-Reticuline was a gift from Tasmanian Alkaloids Pty Ltd (Westbury, Australia), and morphine and codeine were gifts from Sanofi-Aventis (Paris, France). (S)-N-Methylcoclaurine was generated from (S)-coclaurine using purified recombinant NMT1 from Glaucium flavum (58). Enzymatic conversion was performed at 30 °C, with 1 mM (S)-coclaurine, 1 mM SAM, and 20 µg recombinant GfNMT1. Reactions were extracted by ethyl acetate three times and the product identity and purity confirmed by HPLC-MS/MS.

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All other chemicals were purchased from Sigma-Aldrich or Bioshop Canada (ON, Canada).

Plant alkaloid extraction and LC-MS analysis

Nelumbo nucifera samples corresponding to folded and unfolded leaves, rhizomes, roots and embryos were ground to a fine powder under liquid nitrogen using a mortar and pestle. Alkaloid extraction was performed using a modified version of a previously reported protocol for sacred lotus (36). Briefly, 1 mL methanol was added to ~100 mg of plant material, and samples were sonicated in an ultrasonic bath for 30 min at room temperature, centrifuged at 14,000 g for 10 min, and the supernatant was filtered (0.2 µm Acrodisc syringe filter, Pall Corporation, NY, USA). The insoluble material was dried and weighed. Samples (5 µL) were fractionated on a Zorbax C18 column, 2.1 x 50 mm, 1.8 µm (Agilent, California, USA) using a Dionex UltiMate 3000 HPLC system (Thermo Scientific) at a flow rate of 500 uL/min. The mobile phase consisted of solvent A (10 mM ammonium acetate, pH 5.5, 5% (v/v) acetonitrile) and solvent B (100% acetonitrile), starting at 100% (v/v) Solvent A, and ramping from 0 to 20%(v/v) Solvent B over 5 min, 20 to 50% (v/v)Solvent B over 3 min, 50 to 100% (v/v) Solvent B over 3 min, remaining isocratic at 100% (v/v) Solvent B over 2 min, ramping from 100 to 0% (v/v) Solvent B over 0.1 min, and remaining isocratic at 100% (v/v) Solvent A over 1.9 min. The total run time was 15 min and data were collected for the first 10 min. Heated ESI source and interface conditions were operated in positive ion mode as follows: vaporizer temperature 400 °C, source voltage 3kV, sheath gas 60 au, auxiliary gas 20 au, capillary temperature 380 °C, capillary voltage 6 V, tube lens 45 V. A LTQ-Orbitrap-XL (Thermo Fisher) mass spectrometer was operated using LTQ Tune Plus v. 2.5.5 SP1 and Xcalibur software (version 2.1.0.1140), with additional analyses using the QualBrowser feature of Xcalibur. Internal and external calibration, tuning and general operations were performed as previously reported (27,59). Error was maintained at < 2 ppm. Exact masses, retention times and CID spectra of available authentic standards, or previously published data, was used for alkaloid identification (Table **S1**, Figure **S6**). Quantification of norcoclaurine, coclaurine, N-

methylcoclaurine and armepavine was performed using authentic standards.

NnOMT identification and phylogeny

A previously reported Nelumbo nucifera draft genome (30,31) was searched to identify sequences encoding potential O-methyltransferase involved in BIA biosynthesis using the tBLASTn algorithm and functionally characterized OMTs from Papaver somniferum as queries (Ps6OMT, Ps7OMT, PsN7OMT and Ps4'OMT2). Five candidates (NnOMT1-NnOMT5) sharing >40% amino acid sequence identity with query sequences were isolated. The GenBank accession codes were as follows: XM 010245752, XM 010249599, XM 010249600, XM 010273389. and XM 010277761) (Table S2). Protein molecular weight and isoelectric point predictions were made using the Geneious software package (Biomatters, NJ, USA). Amino acid sequence alignments were performed using the default parameters of the MUSCLE algorithm (60) implemented in MEGA X (61). Evolutionary history was inferred using the Maximum Likelihood method established on the JTT matrix-based model (62). The tree with the highest log likelihood was drawn, with branch lengths proportional to the number of substitutions per site, and nodes labelled according to the percentage of trees in which the associated taxa clustered together in the bootstrap test based on 1000 replicates. The amino acid percent identity matrix among Nelumbo nucifera (Nn) and other OMTs was performed using Clustal Omega (63) The GenBank accession codes were as follows: (MH165875); Cc6OMT2 Cc6OMT1 (MH165876); Cj4'OMT (D29812); Ci6OMT (D29811); CjCoOMT (AB073908); CiSOMT (D29809); Ct7OMT (MH165877); **CtSOMT** (MH165874); Ec4'OMT (AB745041); Ec7OMT (AB232153); EcSOMT (LC171865); GfOMT1 (KP176693); GfOMT2 (KP176694); GfOMT6 (KP176698); GfOMT7 (KP176699); Ps4'OMT2 (AY217334); Ps6OMT (AY217335); Ps7OMT (AY268893); PsN7OMT (FJ156103); PsSOMT1 (JN185323); PsSOMT2 (MH029292); PsSOMT3 (MH029294); Tf6OMT (AY610507); Tt6OMT1 (AF064693); Tt6OMT2 (AF064694). Abbreviations were as follows: Cc, Coptis chinensis; Cj, Coptis japonica; Ct, Coptis teeta; Ec, Eschscholzia californica; Gf, Glaucium

flavum; Nn, Nelumbo nucifera; Ps, Papaver somniferum; Tf, Thalictrum flavum; Tt, Thalictrum tuberosum; 4'OMT, 3'-hydroxy-Nmethylcoclaurine 4'-O-methyltransferase; 60MT, 6-*O*-methyltransferase: norcoclaurine 70MT. 7-O-methyltransferase; reticuline CoOMT, columbamine O-methyltransferase; N7OMT, 7-O-methyltransferase; norreticuline SOMT. scoulerine 9-O-methyltransferase. The amino acid sequences of functionally characterized OMTs involved in BIAs biosynthesis used in the present work are provided in Table S9.

NnOMT cDNA isolation and expression

Plant tissues were ground with a mortar and pestle to a fine powder under liquid nitrogen. Total RNA was extracted using the cetyltrimethylammonium bromide (CTAB) method, as previously reported for Nelumbo nucifera tissues (64). RNA quality was confirmed by A_{260/280} and A_{260/230} absorbance measurements using a NanoDrop ND-1000 (Thermo Fisher Scientific) spectrophotometer, and visualization by agarose gel electrophoresis. After genomic DNA removal (AccuRT Genomics DNA Removal kit, Applied Biological Materials, BC, Canada), first-strand cDNA synthesis was performed on 1 µg of RNA using 5X All-in-One RT MasterMix kit, according to the instructions of the manufacturer (Applied Biological Materials, BC, Canada). NnOMT open reading frames were amplified from cDNA using Q5 High Fidelity DNA polymerase (New England Biolabs, MA, USA) and sequence-specific primers (Table S8a) under the following conditions: 98 °C for 30 sec; 35 cycles of 98 °C for 10 sec, 52-65 °C (optimized for each NnOMT) for 30 sec, and 72 °C for 30 sec; 72 °C for 2 min. Amplicons were cloned in a pMiniT 2.0 vector, according to the instructions of the manufacturer (New England Biolabs, MA, USA) and used to transform E. coli TOP 10 strain for colony PCR screening. Plasmids from positive colonies were purified (GeneJet Plasmid Miniprep kit, Thermo-Fisher Scientific, MA, USA) and sequenced. PCR products were purified (GeneJet Gel Extraction Kit, Thermo-Fisher Scientific, MA, USA) and ligated into pRSET-A (Invitrogen, CA, USA) to construct the expression vectors using one-step sequence- and ligation-independent cloning (SLIC) (65). Full-length NnOMT coding regions were cloned in frame with sequences

encoding an N-terminal His₆-tag using specific primers (**Table S8a**).

Recombinant protein purification

The pRSET-A-NnOMT expression plasmids were used to transform the E. coli Rosetta (DE3) pLysS (EMD Millipore, MA, USA) strain and single colonies were used to inoculate 50 mL LB medium supplemented with 100 µg/mL ampicillin and 35 ug/mL chloramphenicol. Cultures were grown at 30 °C with orbital shaking at 200 rpm for 16 h and used to inoculate 1 L LB media (100 µg/mL ampicillin and 35 µg/mL chloramphenicol) to a starting OD₆₀₀ of 0.1. Cultures were grown at 30 °C until $OD_{600} \sim 0.6$, cooled to 16 °C and the production of recombinant proteins was induced by the addition of isopropyl β -D-thiogalactoside (IPTG) to a final concentration of 0.1 mM. Cultures were kept at 16 °C with shaking at 200 rpm for 16 h and cells harvested by centrifugation at 5,000 g for 20 min at 4 °C. Cell pellets were resuspended in 40 mL protein extraction buffer (100 mM Tris-HCl pH 7.5, 300 mM NaCl, 10% [v/v] glycerol) supplemented with 1 mg/mL lysozyme and sonicated on ice for 3 min (10 sec on, 30 sec off). The crude lysate was centrifuged at 16,000 g for 20 min at 4 °C to remove cellular debris. The cleared supernatant was loaded onto 1 mL of equilibrated Talon cobalt affinity resin (Clontech, CA, USA) and incubated at 4 °C for 30 min with gentle shaking. The resin was washed with 20 mL of protein extraction buffer, followed by 5 mL of protein extraction buffer containing 20 mM imidazole. Purified protein was eluted using 5 mL of protein extraction buffer containing 200 mM imidazole, and subsequently concentrated and desalted by repeated ultrafiltration on an Amicon Ultra 30K column (EMD Millipore, MA, USA) in storage buffer (100 mM Tris-HCl pH 7.5, 10% (v/v) glycerol, 1 mM β -mercaptoethanol). Purified His₆-tagged protein concentration was determined using the Bradford reagent according to the instructions of the manufacturer (Thermo Fisher, IL. USA) and using bovine serum albumin as the standard. Protein purity was assessed by SDS-PAGE using a 10% (w/v) gel, which was subsequently transferred to a nitrocellulose membrane. Protein blots were blocked overnight with 5% (w/v) skim milk and incubated for 1 h with a 1:1 000 dilution of His₆-antibody. After washing with TBST buffer (200 mM Tris-HCl pH 7.6, 1.5 M NaCl, 0.1% (v/v) Tween 20), the blots were incubated for 1 h with a 1:10 000 dilution of goat anti-mouse IgG antibodies conjugated with horseradish peroxidize (Bio-Rad). Immunoblots were washed and developed using SuperSignal West Pico chemiluminescent substrate, according to the instructions of the manufacturer (Thermo Scientific, IL, USA) and the bands were visualized using an Amersham Imager 600 (GE Healthcare, IL, USA).

Plant protein extraction

Plant tissues were ground to a fine powder under liquid nitrogen using a mortar and pestle. Total proteins were extracted by adding 5 mL of ice-cold extraction buffer A containing 100 mM Tris-HCl pH 7.5, 10% (v/v) glycerol, 1% (w/v)polyvinylpyrrolidone 40, 5 mM dithiothreitol (DTT) and 1 mM phenylmethylsulfonyl fluoride (PMFS) to ~ 1 g of plant material, and sonicated in an ultrasonic bath for 5 min at 4 °C. Plant debris was eliminated by centrifugation at 14,000 g for 10 min, and the supernatant was desalted in a PD-10 column (GE Healthcare, IL, USA) and eluted in extraction buffer without DTT and PMFS. The solution was concentrated by repeated ultrafiltration using an Amicon Ultra 30K column (EMD Millipore, MA, USA) in storage buffer (100 mM Tris-HCl pH 7.5, 10% glycerol, 1 mM βmercaptoethanol). Crude plant protein concentration was determined using the Bradford reagent according to the instructions of the manufacturer (Thermo Fisher, IL, USA) and using bovine serum albumin as the standard.

Enzyme assays

Substrate-acceptance assays were performed in 100 mM Tris-HCl pH 7.5, using 100 μ M alkaloid, 200 μ M SAM and 5 μ g of purified recombinant protein in a reaction volume of 50 μ L. Negative control assays were performed using purified recombinant protein denatured in boiling water for 10 min. Assays were incubated at 30 °C and quenched after 4 h with 100 μ L acetonitrile, centrifuged at 17,000 g for 40 min to precipitate proteins, and the supernatant analyzed by LC-MS/MS. Product formation, determined as an increase of 14 Da with respect to the substrate, was

confirmed by comparison of retention times and CID spectra with respect to authentic standards, or data reported in the literature. Substrate conversion rates were calculated based on substrate loss. NnOMT1 and NnOMT5 pH-optimum assays were conducted in 25 mM sodium citrate (pH 5.0-6.0), 50 mM sodium phosphate (pH 6.0-7.5), and 100 mM Tris-HCl (pH 7.5-9.0) using 100 µM (R,S)norcoclaurine or 100 µM (S)-N-methylcoclaurine, respectively, 200 µM SAM and 1 µg purified recombinant protein in a reaction volume of 50 μL. Assays were incubated within the linear range for product formation at 30 °C for 5 min and stopped with 100 µL acetonitrile. Temperature optima were determined under identical conditions except that 100 mM Tris-HCl, pH 8 and incubation temperatures between 4-60 °C were used. Kinetic parameters were measured for 5 min under the same conditions except that NnOMT1 was incubated with (R,S)-norcoclaurine at 30 °C and NnOMT5 with either (R,S)-norcoclaurine or (S)-N-methylcoclaurine at 37 °C, using alkaloid concentrations from 5 to 400 µM and a fixed SAM concentration of 400 µM. Product formation for NnOMT1 and NnOMT5 was quantified by comparison with authentic standards of coclaurine and armepavine, respectively. All assays were performed in triplicates. Saturation curves and kinetic parameters were determined based on the Michaelis-Menten equation (NnOMT1 and NnOMT5 incubated with (R,S)-norcoclaurine) or a substrate-inhibition equation (NnOMT5 incubated with (S)-N-methylcoclaurine), using Prism 5 (GraphPad, CA, USA).

Plant protein assays were performed using 50 μ g of crude protein in 100 mM Tris-HCl pH 7.5, using 200 μ M alkaloid and 400 μ M SAM in a 100 μ L reaction. Negative control assays were performed using crude plant protein extracts denatured in boiling water for 10 min. Reactions were incubated overnight at 30 °C, quenched with 200 μ L acetonitrile and analyzed by LC-MS/MS. Product formation, determined as an increase of 14 Da with respect to the substrate, was confirmed by comparison of retention times and CID spectra with respect to authentic standards.

HPLC-MS/MS

HPLC-MS/MS was performed using a 1200 HPLC instrument coupled to a 6410 triple quadrupole MS

(Agilent, CA, USA). Samples (5 µL) were injected onto a Poroshell 120 SB-C18 HPLC column, 2.1 x 50 mm, 2.7-µm particle size (Agilent, CA, USA). Analytes were eluted using a mobile phase gradient of Solvent A (10 mM ammonium acetate, pH 5.5, 5% (v/v) acetonitrile) and Solvent B (100% acetonitrile) at a flow rate of 600 µL/min. The gradient started at 100% (v/v) Solvent A, ramped linearly to 60% (v/v) Solvent B by 8 min, further increased linearly to 99% (v/v) Solvent B over 2 min, remained isocratic at 99% (v/v) Solvent B from 10 to 11 min, and returned to 100% (v/v) Solvent A at 11.1 min for a 3 min reequilibration period. Analytes were applied to the mass analyzer using an electrospray ionization probe operating in positive mode with the following conditions: capillary voltage 4 000 V, fragmentor voltage 110 V, source temperature 350 °C, nebulizer pressure 50 psi, gas flow 10 L/min. For full-scan analysis, quadrupole 1 and 2 were set to radio frequency only, whereas the third quadrupole scanned from m/z 200 to m/z 700. ESI[+] and CID spectra were analyzed, the precursor m/z was selected in quadrupole 1 and collision energy of 25 eV and an argon collision gas pressure of 1.8×10^{-3} torr were applied in quadrupole 2. The resulting MS^2 fragments were resolved by quadrupole 3 scanning from 30 m/z to 5 m/z greater than the precursor ion m/z. Compounds were identified based on retention times and ESI[+]-CID spectra compared with authentic standards or published spectral data.

Quantitative real-time PCR

RNA extraction and cDNA synthesis were performed as described above. qRT-PCR analysis was performed using PowerUp SYBR Green Master Mix and a QuantStudio-3 Real-Time PCR System (Applied Biosystems, MA, USA). Reactions (10 μ L) contained diluted cDNA (~8 ng), 5 μ L SYBR Green Master Mix (2X) (Applied Biosystems), and 500 nM of each primer (**Table S8b**). Thermal conditions were 50 °C for 2 min, 95 °C for 2 min, 40 cycles of 95 °C for 1 sec and 60 °C for 30 sec. Amplification specificity was confirmed by melt-curve analysis, generated at a ramp rate of 1.6 °C/sec to 95 °C maintained for 15 sec, 1.6 °C/sec to 60 °C maintained for 1 min, and 0.15 °C/sec to 95 °C, which was maintained for 15 sec. Primer efficiency was verified using LinRegPCR software (66) and samples with values between 1.8-2 were selected for analysis. Relative transcript abundance was calculated by the 2^{- $\Delta\Delta$ Ct} method (67), using β -actin as the endogenous reference gene (68). Transcript levels were normalized with respect to the gene showing the lowest expression level.

Homology modeling

The previously reported crystal structure from *Thalictrum flavum* norcoclaurine 6-*O*-methyltransferase (Tf6OMT) in complex with (*S*)-norlaudanosoline and *S*-adenosyl-L-homocysteine (PDB 5ICE) (18) was used as template to generate a NnOMT5 homology model using SWISS-MODEL (69). The resulting NnOMT5 model and Tf6OMT crystal structure were visualized, and figures were generated, using PyMOL Molecular Graphic System (v2.3.1, Schrödinger, LLC).

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Author contributions: Conceptualization, IMMP, PJF; Methodology, IMMP, PJF; Formal Analysis, IMMP; Investigation, IMMP; Resources, PJF; Data curation, IMMP; Writing – Original draft, IMMP; Visualization IMMP, PJF; Funding Acquisition, PJF. All authors reviewed the results and approved the final version of the manuscript.

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FOOTNOTES

Abbreviations: 4'OMT, 3'-hydroxy-*N*-methylcoclaurine 4'-*O*-methyltransferase; 6OMT, norcoclaurine 6-*O*-methyltransferase; 7OMT, reticuline 7-*O*-methyltransferase; BIA, benzylisoquinoline alkaloid; CID, collision-induced dissociation; CoOMT, columbamine *O*-methyltransferase; CYP, cytochrome P450 monooxygenase; ESI, electrospray ionization; IPTG, isopropyl β -D-thiogalactoside; N7OMT, norreticuline 7-*O*-methyltransferase; NCS, norcoclaurine synthase; NMT, *N*-methyltransferase; OD, optical density; OMT, *O*-methyltransferase; PR10, pathogenesis-related 10 protein; SOMT, scoulerine 9-*O*-methyltransferase.

Table 1: Kinetics parameters for recombinant NnOMT1 (60MT) and NnOMT5 (60MT and 7	(OMT)
determined at a fixed saturating concentration of SAM. Values represent the mean ± standard devia	tion of
three independent measurements.	

Enzyme	Substrate	Κ _m (μΜ)	V _{max} (nmol min ⁻¹ mg ⁻¹ protein)	<i>k_{cat}</i> (s ⁻¹)	k _{cat} / K _m (M ⁻¹ s ⁻¹)
NnOMT1	(R,S)-Norcoclaurine	20 ± 2	52 ± 2	0.0336	1686
NnOMT5	(R,S)-Norcoclaurine	152 ± 36	9 ± 1	0.0058	38
	(S)-N-Methylcoclaurine	13 ± 3	30 ± 2	0.0191	1465

FIGURE LEGENDS



Figure 1: Schematic representation of the proposed 1-benzylisoquinoline alkaloid biosynthetic pathway in sacred lotus (*Nelumbo nucifera*). The enzymes functionally characterized in this study (NnOMT1 and NnOMT5) are indicated, and detected alkaloids are shown in bold. Asterisk denotes the chiral center in norcoclaurine. Pink arrow, norcoclaurine synthase (NCS); green arrows, 6-O-methyltransferase (6OMT); red arrows, 7-O-methyltransferase (7OMT); blue arrow, 4'-O-methyltransferase (4'OMT); black arrows, *N*-methyltransferase (NMT).



Figure 2: (a) Diagram of sacred lotus. (b) Heatmap showing the organ-specific profile of BIAs in two varieties of sacred lotus (Pink and White). Relative abundance corresponds to the mean value of three independent replicates. Values were normalized to the sample with the highest level for each compound. (c) Structures corresponding to detected aporphine, pro-aporphine and bisbenzylisoquinoline alkaloids isolated from sacred lotus (for 1-benzylisoquinoline alkaloid structures refer to Figure 1). FL, folded leaf; UL, unfolded leaf; Rh, rhizome; Ro, roots; E, embryos.



Figure 3: Phylogenetic relationships among sacred lotus *O*-methyltransferase (NnOMT) candidates and functionally characterized OMTs from BIA-accumulating plants in the Ranunculales. The evolutionary history was inferred using the Maximum Likelihood and the JTT matrix-based model. The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (1000 replicates) is shown next to the branches. The tree with the highest log likelihood is drawn to scale, with branch lengths measured in the number of substitutions per site. Evolutionary analyses were conducted in MEGAX. Cc, *Coptis chinensis*; Cj, *Coptis japonica*; Ct, *Coptis teeta*; Ec, *Eschscholzia californica*; Gf, *Glaucium flavum*; Nn, *Nelumbo nucifera*; Ps, *Papaver somniferum*; Tf, *Thalictrum flavum*; Tt, *Thalictrum tuberosum*. 4'OMT: 3'-hydroxy-*N*-methylcoclaurine 4'-*O*-methyltransferase; 6OMT: norcoclaurine 6-*O*-methyltransferase; 7OMT: reticuline 7-*O*-methyltransferase; CoOMT: columbamine *O*-methyltransferase; N7OMT: norreticuline 7-*O*-methyltransferase; OMT: *O*-methyltransferase; SOMT: scoulerine 9-*O*-methyltransferase. Sacred lotus NnOMT1 (green box) and NnOMT5 (red box) are indicated.



Figure 4: Multiple sequence alignment among *Nelumbo nucifera O*-methyltransferase (NnOMT) candidates and *Thalictrum flavum* 60MT (Tf60MT). Residues shared by Tf60MT and sacred lotus OMTs are shaded in grey, and fully conserved residues are shaded in black. Asterisks indicate catalytic (H^{256} , D^{257} and E^{315}) and other key residues implicated in BIA (G^{165} , D^{169} , C^{253} and D^{306}) and SAM (T^{170} , G^{195} , D^{218} , D^{238} and K^{252}) binding, as reported for Tf60MT. Conserved motives I-IV are underlined.

O-Methyltransferases from sacred lotus



Figure 5: Substrate range for recombinant NnOMT1 and NnOMT5. Values represent the mean \pm standard deviation of three independent replicates. The structure of the corresponding substrates is shown; nd, not detected.



Figure 6: Organ-specific correlational analysis of 1-benzylisoquinoline alkaloids metabolism in two varieties (Pink and White) of sacred lotus. (a) Content of selected 1-benzylisoquinoline alkaloids. (b) 60MT, 70MT and 4'OMT specific activity of plant crude extracts when incubated with (R,S)-norcoclaurine, (S)-N-methylcoclaurine and (R)-armepavine, respectively. (c) Relative abundance of NnOMT1-NnOMT5 transcripts. Values represent the mean \pm standard deviation of three independent measurements. FL, folded leaf; UL, unfolded leaf; Rh, rhizome; Ro, roots; E, embryos.

Isolation and characterization of two *O*-methyltransferases involved in benzylisoquinoline alkaloids biosynthesis in sacred lotus (*Nelumbo nucifera*) Ivette M. Menéndez-Perdomo and Peter J. Facchini

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