Rapid Identification of Enzyme Variants for Reengineered Alkaloid Biosynthesis in Periwinkle

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SUMMARY

Monoterpene indole alkaloids from *Catharanthus roseus* (Madagascar periwinkle), such as the anticancer agents vinblastine and vincristine, have important pharmacological activities. Metabolic engineering of alkaloid biosynthesis can provide an efficient and environmentally friendly route to analogs of these synthetically challenging and pharmaceutically valuable natural products. However, the narrow substrate scope of strictosidine synthase, the enzyme at the entry point of the pathway, limits a pathway engineering approach. We demonstrate that with a different expression system and screening method it is possible to rapidly identify strictosidine synthase variants that accept tryptamine analogs not turned over by the wild-type enzyme. The variants are used in stereoselective synthesis of β-carboline analogs and are assessed for biosynthetic competence within the terpene indole alkaloid pathway. These results present an opportunity to explore metabolic engineering of "unnatural" product production in the plant periwinkle.

INTRODUCTION

Monoterpene indole alkaloids (MIAs) from *Catharanthus roseus* (Madagascar periwinkle) are structurally complex natural products with well-established pharmacognosy [1, 2]. The MIA biosynthetic pathway produces the antihypertensive compound ajmalicine, **1**; the topoisomerase II inhibitor serpentine, **2**; and the anticancer agents vinblastine, **3**, and vincristine, **4** (Figure 1) [1, 3, 4]. Interestingly, derivatives of vinblastine such as Vinflunine (4′-deoxy-20′,20′-difluoro-C′-norvincaleukoblastine), **5**, show excellent anticancer activity in the clinic, indicating that small changes to the alkaloid scaffold can improve or modulate bioactivity (Figure 1) [5–7]. However, the current supply of MIA analogs is primarily dependent on semisynthesis from isolated pathway intermediates that are produced in low yields.

Metabolic engineering of a biosynthetic pathway can provide an efficient and environmentally friendly route to novel natural products or natural product derivatives. We previously demonstrated that the MIA biosynthetic pathway in *C. roseus* root cultures and seedlings can utilize nonnatural substrates to synthesize an array of unnatural MIA analogs [8]. However, the low substrate promiscuity of the enzyme at the entry point in the pathway, strictosidine synthase (STS), limits the scope of precursor-directed biosynthesis efforts. STS catalyzes the stereoselective Pictet-Spengler reaction between tryptamine, **6a**, and secologanin, **7**, to yield the tetrahydro-β-carboline (3*S*)-strictosidine, **8a**, as a single diastereomer (Figure 2) [9]. Deglucosylation of **8a** by the second enzyme of the pathway, strictosidine glucosidase (SG), produces the aglucone, **9**, that is, in turn, enzymatically converted into the structurally diverse group of MIAs produced in *C. roseus* (Figure 2) [1, 2].

Reengineering STS substrate specificity is key for biosynthetic production of many MIA analogs. We recently reported that a secologanin analog already accepted by wild-type STS is selectively turned over by an STS mutant (Asp177Ala) [10]. The observed switch in selectivity could be rationalized based on the crystal structure of STS from *Rauvolfia serpentina* in complex with secologanin (PDB ID: 2FPC) [11]. The efficiency of the rational redesign of enzyme active sites can often be improved by using saturation mutagenesis protocols. However, the ineffectiveness of current methods to assess STS activity limits this "random" rational approach. Therefore, to rapidly obtain STS mutants with expanded tryptamine specificity, we developed a streamlined expression system and colorimetric assay for the Pictet-Spengler reaction. Herein, we describe the discovery of two STS mutants using this discovery strategy that have broadened substrate specificity. We highlight the potential applications of these mutants in biosynthesis of unnatural alkaloid analogs.

RESULTS AND DISCUSSION

Development of a Colorimetric Assay for STS Activity

The time-consuming high-performance liquid chromatography (HPLC) assay used to measure catalytic activity limits rapid reengineering of STS substrate specificity. There are no reports of a high-throughput assay for Pictet-Spengler activity. However, previous literature reports noted that a yellow pigment forms after deglycosylation of **8a** by SG [12, 13]. Inspired by these results, we developed
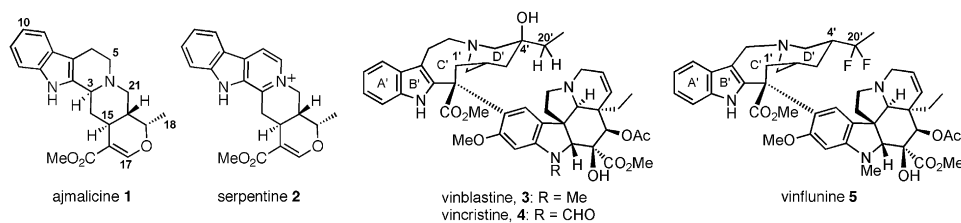
Figure 1. Structures of Selected Medicinal MIAs Produced or Derived from Madagascar Periwinkle, *C. roseus*

Ajmalicine, **1**, acts as an antihypertensive agent; serpentine, **2**, is a topoisomerase II inhibitor; vinblastine, **3**, and vincristine, **4**, are anticancer agents; and Vinflunine, **5**, is a modified vinblastine analog with promising bioactivity.

an assay for visual detection of STS activity that could be adapted to medium- or high-throughput screening.

In vitro deglycosylation by SG yields an equilibrium mixture of isomers of 4,21-dehydrogeissoschizine, **10**, and cathenamine, **11** (Figure 2) [14]. Previous literature suggested that the yellow chromophore was a low-molecular weight degradation product of **10** (dihydroflavopereirine) [15]. However, liquid chromatography coupled to high-resolution mass spectrometry (LC-HRMS) of yellow-colored assay mixtures produced by us indicated the formation of compounds with *m/z* 493, 491, and 489 from the initial deglycosylation products **10** and **11** (*m/z* 351). The compound corresponding to *m/z* 493 disappeared after overnight incubation to form compounds with *m/z* 491 and 489 exclusively.

It has been reported that deglycosylated **8a** can form amine adducts in the presence of excess ammonium sulfate, and that these adducts can further rearrange to form a dihydropyridine ring, which can easily air oxidize to an aromatic pyridine ring [16]. HRMS suggested that the compounds in our assay corresponding to *m/z* 493, 491, and 489 have molecular formulae consistent with tryptamine adducts of **10** in different oxidation states (see the
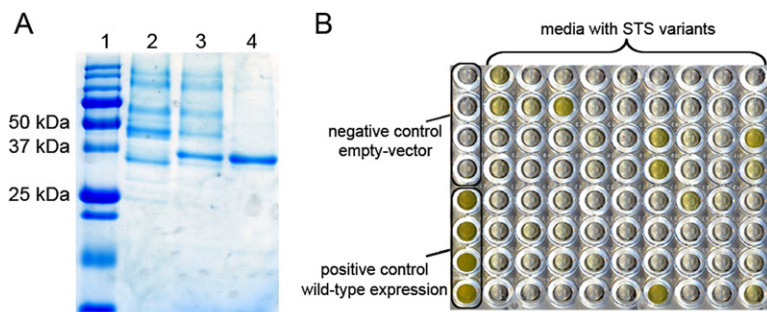
Supplemental Data available with this article online). Purification of the compounds corresponding to *m/z* 491 and 489 and analysis by 1D- and 2D-NMR suggested that the tryptamine adducts formed in the assay undergo rearrangement and air oxidation, similar to what is reported by Heinstein et al. [16] (Figure 2) (Supplemental Data). The UV-VIS spectrum of the compound corresponding to *m/z* 489 showed absorption maxima at 314 nm and 427 nm; *m/z* 491 had one unique absorption maximum at 470 nm (data not shown).

If this assay is to be used for STS substrate reengineering, SG must accept analogs of **8a**, and the resulting deglycosylated adducts must form the pigment. To determine the scope of the assay, we synthesized a variety of strictosidine analogs (maleic acid buffer, 10 mM [pH 2]) to obtain a diastereomeric mixture of strictosidine analogs that were then subjected to the assay conditions. Pigment formation in the presence of excess tryptamine, **8a**, or tryptamine analogs yielded strictosidine, **8a**, or strictosidine analogs with substituents on the indole ring or on the side chain (data not shown). Careful analysis showed that the purified strictosidine analog **8m** (*m/z* 561, 531+30) formed compounds corresponding to the
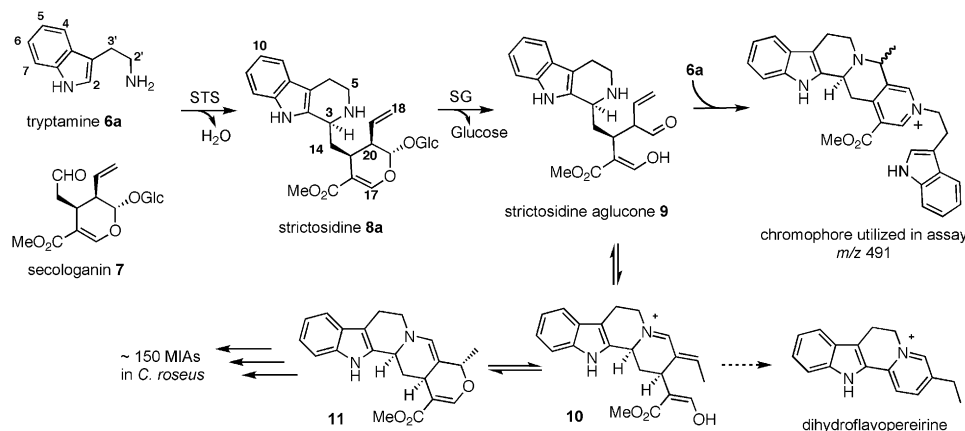
Figure 2. STS-Catalyzed Formation of 8a via a Pictet-Spengler Reaction between 6a and 7

The reaction proceeds via an iminium intermediate, which undergoes electrophilic aromatic substitution and cyclization to give **8a**. Deglucosylation by a dedicated β-glucosidase forms an aglucone, **9**, that rearranges into dehydrogeissochizine, **10**, and cathenamine, **11**. The deglucosylated products are then enzymatically converted into the MIA structures observed in periwinkle. However, aglucone, **9**, can be intercepted by amine nucleophiles to form colored adducts.

expected aglucone analogs **9m** and **10m** after deglycosylation by SG (*m/z* 381, 351+30). Only after addition of excess tryptamine analog **6m** to the deglycosylated products did the color form and the expected masses of the analogous tryptamine adducts appear (*m/z* 553, 551, and 549, M+60). Control assays lacking tryptamine or a tryptamine analog did not form the putative adducts and were not colored (Supplemental Data). Because of the slow rate of chromophore formation relative to strictosidine formation, this assay cannot be used as a quantitative coupled assay of STS activity. Nevertheless, this assay appeared to be a robust method for qualitative detection of STS activity.

Expression System for Extracellular STS Activity

To facilitate screening for STS activity, we constructed a plasmid that allows for export of STS to the media upon expression in *Saccharomyces cerevisiae*. Although the *sts* gene from *C. roseus* contains a putative native signal peptide (residues 1–26), expression in yeast did not result in significant protein export in *S. cerevisiae*. Replacement of the nucleotides encoding the first 26 residues with an N-terminal yeast mating-factor α signal sequence [17] efficiently directed the export of STS to the culture media (Figure 3A, lane 2). A C-terminal FLAG-peptide was also introduced to allow for purification by affinity chromatography (Figure 3A, lanes 3 and 4) [18].

Characterization of purified wild-type STS expressed in yeast by a quantitative HPLC assay afforded kinetic constants (V_{max} = 21 U mg^{-1}, K_M = 1.4 μM; turnover number assuming all enzyme is active: k_{cat} = 78 min^{-1}) (Table 1). These values are consistent with constants reported for STS purified from *C. roseus* (average of several isoforms: V_{max} = 14 U mg^{-1}, K_M = 9 μM) [19] and from heterologously expressed STS from the medicinal plant *Rauvolfia serpentina* (k_{cat} = 78 min^{-1}, K_M = 6 μM, 88% sequence similarity to *C. roseus* STS) [11]. Mutagenesis studies of *R. serpentina* STS suggested that Glu309 is important for catalysis (Glu309 corresponds to Glu315 in the *C. roseus* protein sequence) (Figure 4A) [11]. The >1000-fold reduced turnover number of the analogous mutant of *C. roseus* STS (Glu315Ala, V_{max} = 0.012 U mg^{-1}, K_M = 0.44 mM) confirms this conclusion. When subjected to the colorimetric assay conditions, Glu315Ala produced

a faint yellow pigment (data not shown), suggesting that this screen is sufficiently sensitive to detect weakly active mutants.

Validation of Mutagenesis and Screening Strategy

The crystal structure of *R. serpentina* STS in complex with tryptamine (PDB ID: 2FPB) was used to evaluate residues involved in substrate binding and catalysis [11]. The relatively high B factor of **6a** in the crystal structure (3-fold higher than the remainder of the protein) suggests that **6a** may adopt a relatively flexible orientation in the active site [11]. Moreover, the known substrate specificity of STS cannot always be rationalized from the structural data; for example, it is not clear why 5-hydroxytryptamine, **6f**, is weakly turned over while 5-methyltryptamine, **6l**, is not. To expand our rational redesign approach, we used saturation mutagenesis in combination with our screen to probe all 20 naturally occurring amino acids at selected residues in the tryptamine-binding pocket.

Codons were randomized by using primers containing NNK/MNN degenerate codons (32 sequence possibilities). To validate the mutagenesis strategy, expression system, and screening method, a saturation mutagenesis library of a tyrosine residue distant from the active site (Tyr167Ala greatly reduces STS catalytic activity, unpublished data) was constructed. The mutated gene and the linearized expression vector were transformed into the yeast expression host (*S. cerevisiae* BJ5465), and homologous recombination resulted in the production of the desired STS variant library. Colonies were inoculated (88 per mutated site, an oversampling that gives 94% certainty that the library is complete [20]) into expression media in deep-well, 96-well plates. The culture media were then transferred to 96-well microtiter plates containing the assay components **6a**, **7**, and SG. A positive control from a yeast strain expressing wild-type and a negative (empty-vector) control were included in the screen (Figure 3B, column 1). Five clones that led to the formation of a strong, yellow pigment were sequenced, and each contained the wild-type gene. Cultures lacking pigment yielded genes with a variety of mutations at tyrosine 167. This test assay with the natural substrate **6a** indicated that the screen was robust enough to use to search for STS variants with altered substrate specificity.

Table 1. Kinetic Constants for STS Variants Varying Tryptamine 6a or an Analog of 6a

Variant, Substrate	V_{max} (U mg^{-1})[a]	K_M (mM)	V_{max}/K_M (U mg^{-1} mM^{-1})[a]
Wild-type, **6a**	21	0.0014	15,000
Glu315Ala, **6a**	0.012	0.44	0.027
Val214Met, **6a**	3.1	0.42	7.5
Val214Met, **6l**[b]	0.051	0.37	0.14
Phe232Leu, **6a**	0.24	0.62	0.39
Phe232Leu, **6m**[b]	0.23	0.94	0.24

Kinetic assays were performed in 100 mM sodium phosphate buffer (pH 7.0) using an internal standard (naphthyl acetic acid), and samples were analyzed by HPLC for formation of product. The error limit for kinetic assays is estimated at 20%. V and V/K parameters are obtained by nonlinear regression analysis by using the Northrop equation, and K_M is obtained from the Michaelis-Menten equation.
[a] 1 U = formation of 1 μmol product (**8a** or analog) per minute.
[b] Compounds **6l** and **6m** are not accepted by wild-type STS or Glu315Ala STS.

Selection of Substrates for Screening and Residues for Mutagenesis

With the exception of Glu315, which is crucial for the catalysis of the Pictet-Spengler reaction, the tryptamine-binding site consists primarily of hydrophobic and aromatic residues, including Trp153, Tyr155, Val171, Val182, Val214, and Phe232 (Figure 4A). Therefore, the pocket may simply serve to foster tryptamine binding to the enzyme in a productive orientation that promotes iminium formation with the aldehyde substrate and cyclization to form the tetrahydro-β-carboline. We hypothesized that mutagenesis of residues in the tryptamine-binding pocket will allow for productive binding and catalysis of substrates not accepted by the wild-type enzyme.

We selected substrates for screening based on previous work on STS substrate specificity (Figure 4B). Tryptamine analogs with a methyl substituent at the 4 or 7 position (**6c**, **6j**) are accepted by the wild-type enzyme (11- and 27-fold increased K_M and 5- and 3-fold decreased k_{cat} with **6c** and **6j**, respectively) [21]. Substitutions at the 5 and 6 positions are less tolerated: although 5-hydroxytryptamine, **6f**, and 6-methoxytryptamine, **6h**, are weakly turned over by the wild-type enzyme, 5-methyltryptamine, **6l**, is not a substrate. We selected **6l** as a model compound to assay mutants with increased tolerance to 5 indole substitutions. The 2′ position of the tryptamine side chain is also highly sensitive toward substitution. A methyl group (racemic starting material) is only weakly tolerated at the 2′ position [21]. We determined that (2′R)-tryptophanol, **6m**; (2′S)-tryptophanol, **6p**; (2′R)-tryptophan, **6q**; (2′S)-tryptophan, **6r**; and (2′R)-tryptophan methyl ester, **6s**, are not accepted by the wild-type enzyme. Compounds **6m** and **6p** were selected for screening.

The intriguing limitation at the 5 position may be due to the proximity of Val171 or Val214 (closest distances: 4.4 Å

and 3.2 Å, respectively). The 2′ position is close to the reactive amine, and large substituents at this position may interfere with the catalytic machinery. In addition, Phe232 could prevent reaction with tryptamine analogs that have substitutions at the 2′-proR hydrogen atom (closest distance: 3.8 Å), while Glu315 could prevent the proS substitutions (3.3 Å).

We anticipated that saturation mutagenesis of residues Val171 or Val214 could help turnover of **6l**, and mutagenesis of Phe232 or Glu315 would allow for reaction with **6m** and **6p** (Figure 5). These four residues formed the basis for our first efforts to alter the tryptamine substrate specificity for STS.

Discovery of Active Mutants

The Val171, Val214, Phe232, or Glu315 libraries were expressed in 96-well plates. After protein expression, the media containing the enzyme were transferred to microtiter plates for screening by using an assay solution containing the appropriate analogs of **6a**, as well as **7** and SG. Color formation was detected visually after 24 hr of incubation (Supplemental Data). The Val171 and Glu315 libraries did not produce positive hits toward **6l**, **6m**, or **6p** (Figure 5). However, several members of the Val214 library showed activity toward **6l**, as evidenced by the colorimetric assay. Interestingly, sequencing of three active mutants revealed a Val214Met mutation. The methionine substitution eliminates the side chain β branching that could hinder reaction with **6l** in wild-type STS. Since no mutants containing smaller amino acids at Val214 were isolated from the screen, we speculate that a longer side chain is key for orienting the tryptamine analog optimally for catalysis. (A different *R. serpentina* STS variant that accepts **6l** was also found by J. Stöckigt and coworkers, personal communication.) Further evaluation of this mutant by LC-MS and HPLC showed that the Val214Met mutant also accepts 5-chlorotryptamine, **6n**, and 5-bromotryptamine, **6o**. Halogenated precursors are useful because the distinctive isotopic signature allows for facile detection of metabolites. Furthermore, aryl chlorides and bromides can be used to derivatize pathway products via aromatic crosscoupling reactions [22]. There is also some evidence that bromination at this position in synthetic vinblastine analogs has resulted in compounds displaying promising antitumor activity [23].

The colorimetric assay also indicated that **6m** was accepted by members of the Phe232 library (Figure 5). Sequencing of six active clones showed a substitution at residue 232 to leucine (four of the six sequenced cultures) or methionine (two of the six sequenced cultures). The 2′-proR hydrogen atom projects toward Phe232 (closest distance: 3.4 Å), and mutagenesis of Phe232 to the smaller leucine or methionine residue may help accommodate the hydroxymethylene substituent. Initial experiments with HPLC indicated that Phe232Leu was slightly more active under the assay conditions; thus, the Phe232Leu mutant was chosen for further characterization. Notably, **6p**, **6r**, and other tryptamine analogs with a substitution at the 2′-proS hydrogen are directed toward the catalytic
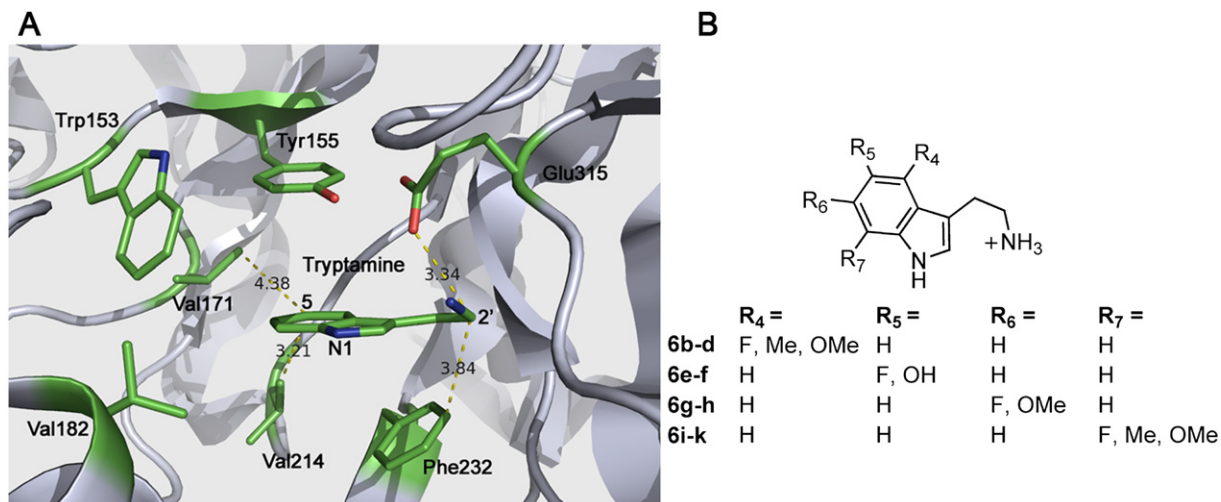
**Figure 4. Tryptamine-Binding Site and Tryptamine Substrate Scope of STS**

(A) Tryptamine-binding site of STS from *R. serpentina* (PDB ID: 2FPB, residues are numbered according to the *C. roseus* peptide sequence). Side chains interacting with tryptamine are in stick representation. The protein main chain is represented in cartoon form.

(B) Tryptamine analogs with indole ring substitutions that are accepted by wild-type STS [21].

Glu315 residue. Continued redesign of mutants that accept these substituents would likely be unwarranted, as Glu315 is required for catalysis. Turnover of L-tryptophan, **6r**, by STS could impact the cellular supply of tryptophan and was likely subject to strong negative selection during the evolution of STS.
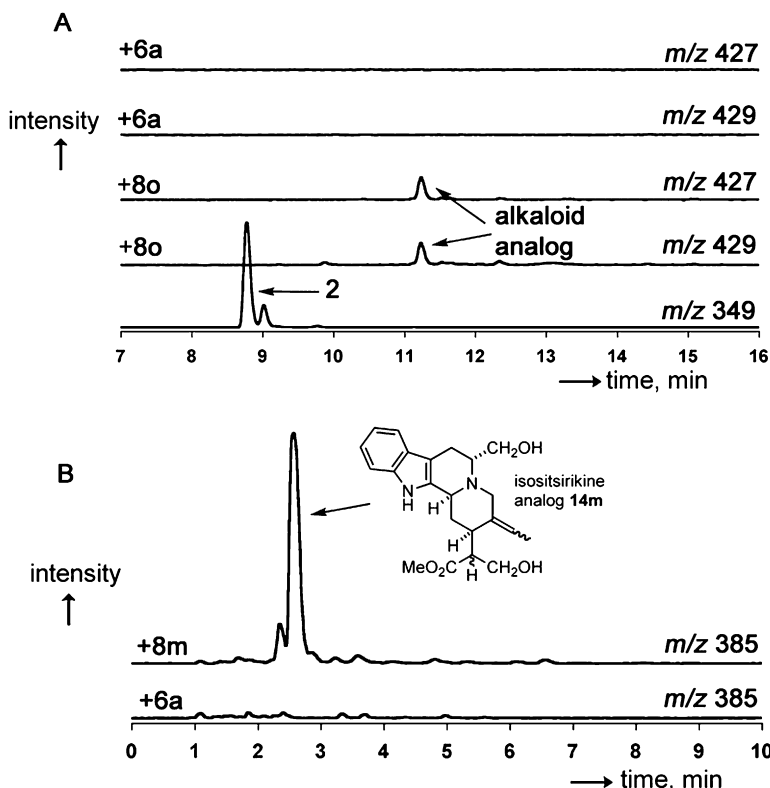
Kinetic Analysis of STS Mutants

Larger-scale (500 ml) expression of the enzyme mutants in yeast followed by purification of the concentrated media with anti-FLAG antibody resin afforded Val214Met and Phe232Leu in apparent homogeneity, as indicated by SDS-PAGE (data not shown). Steady-state kinetic analysis of Val214Met and Phe232Leu was performed with both the natural substrate **6a** and the substrate analogs **6l** and **6m** (Table 1). The mutants each have reduced

catalytic capacity compared to the wild-type enzyme ($V_{max,wt}/V_{max,mutant}$) using **6a** (7-fold and 90-fold, respectively). In addition to the 90-fold reduced V_{max}, Phe232Leu shows a >400-fold increased Michaelis constant (Table 1). This may be attributed to loss of the aromatic residue, since phenylalanine may help orient the substrate correctly for catalysis or stabilize high-energy intermediates in the reaction pathway. A comparison of the catalytic efficiency (V_{max}/K_M) of the mutants using **6a** or the respective analogs of **6a** suggested that both mutants are selective for tryptamine in the presence of the analog (50-fold and 1.5-fold for Val214Met and Phe232Leu, respectively, Table 1). Lowering the amount of **6a** in cell culture by inhibiting tryptophan decarboxylase—the enzyme directly responsible for tryptamine biosynthesis—may be necessary to suppress formation of native MIA analogs.
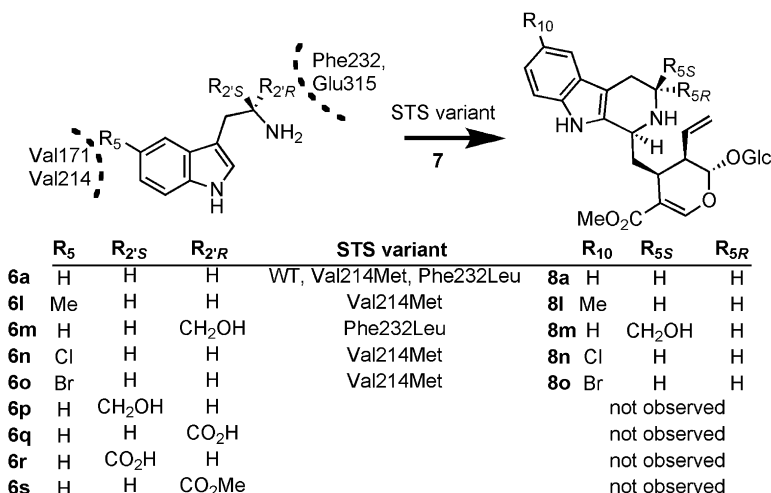
Figure 5. Expanded Substrate Scope: Tryptamine Analogs Assayed with STS Variants Used in This Study

	R_5	R_{2'S}	R_{2'R}	STS variant		R_{10}	R_{5S}	R_{5R}
6a	H	H	H	WT, Val214Met, Phe232Leu	8a	H	H	H
6l	Me	H	H	Val214Met	8l	Me	H	H
6m	H	H	CH_2OH	Phe232Leu	8m	H	CH_2OH	H
6n	Cl	H	H	Val214Met	8n	Cl	H	H
6o	Br	H	H	Val214Met	8o	Br	H	H
6p	H	CH_2OH	H		not observed			
6q	H	H	CO_2H		not observed			
6r	H	CO_2H	H		not observed			
6s	H	H	CO_2Me		not observed			

Figure 6. LC Traces of Selected Alkaloid Analogs Formed by Feeding Strictosidine Analogs to 2-Week-Old *C. roseus* Hairy Root Cultures

(A) Formation of putative serpentine or a serpentine isomer analog upon feeding of **8o** to periwinkle. The two top traces show that no compounds corresponding to *m/z* 427 and 429 exist in the culture supplemented with tryptamine, **6a**. The bottom trace shows the elution of an authentic standard of serpentine, **2**. The remaining two traces show the formation of an alkaloid analog displaying the expected isotopic pattern that could be a brominated MIA analog.

(B) Formation of isositsirikine, **14m**, in a feeding study of **8m**. The top trace shows the expected mass of the alkaloid analog; the bottom trace shows that no naturally occurring alkaloid of *m/z* 385 coelutes with **14m**.

Preparative Enzymatic Synthesis of 8l–8n and Characterization of Products

We used the Val214Met and Phe232Leu mutants in the stereoselective preparative (mmol-scale) synthesis of (3S)-strictosidine analogs **8l–8n**. Although several chemical strategies for Pictet-Spengler reactions exist, formation of **8a** in acidic aqueous media gave a low-yielding diastereomeric mixture of products (3R and 3S), as well as numerous side products (data not shown). Expression of the *sts* mutants in *E. coli* afforded crude cell extracts containing either Val214Met or Phe232Leu. Despite the reduced V$_{max}$ of the mutants relative to wild-type enzyme (Table 1), the extracts were conveniently used to synthesize **8l–8n** on a mmol-scale. Gratifyingly, side products were minimal, and the expected (3S)-diasteromer was formed exclusively (d.r. > 99.5%). Preparative HPLC purification was used to separate the products from the unreacted starting materials. NMR and HRMS validated the structures of the predicted strictosidine analogs (Experimental Procedures).

Biosynthetic Competence of Strictosidine Analogs

We envision that Val214Met and Phe232Leu will be useful for metabolic pathway engineering of alkaloid biosynthesis. Previous reports demonstrate that MIA pathways accept and process tryptamine and secologanin analogs to form unnatural alkaloids [8, 10]. However, it is important to validate whether the strictosidine analogs that are produced by STS mutants are in fact accepted by the remainder of the pathway.

Decades of research on *C. roseus* plants and differentiated cell cultures have yielded a wealth of structural and biochemical information for the alkaloid profile, thereby facilitating analysis of the complex alkaloid production in *C. roseus* [1]. LC-MS analysis of hairy root cultures supplemented with the natural substrate tryptamine **6a** showed, as expected, formation of the heteroyohimbine alkaloids ajmalicine, **1** (*m/z* 353); serpentine, **2** (*m/z* 349); and yohimbine, **12** (*m/z* 355), as well as the more complex alkaloid akuammacine, **13** (*m/z* 323) [8].

To assess the metabolic fate of tryptamine analogs that are not accepted by the wild-type enzyme, the corresponding strictosidine analogs have to be preformed and then fed to the hairy root cell cultures. Reports in the literature indicated that the relatively large strictosidine biosynthetic intermediate could be taken up by plant cell culture [24]. *C. roseus* hairy root cultures were supplemented with preformed **8m** (product of Phe232Leu) and **8o** (product of Val214Met) (Figure 5) at a final concentration of 0.5 mM, and the alkaloid content was extracted from the media and the roots by using established protocols [25]. Analysis of the media and the extracted cell cultures by LC-MS suggested the presence of compounds with masses corresponding to MIA analogs (with M+30 for alkaloids derived from **8m** and M+78/80 for alkaloids derived from **8o** (Figure 6 and Supplemental Data). These compounds were not observed in control cultures that contained only the natural substrate tryptamine, **6a**. Compounds derived from **8o** could be identified by the characteristic isotopic distribution expected for brominated

molecules ($^{79}Br/^{81}Br$), providing further support that these metabolites are derived from the halogenated strictosidine precursor (Figure 6). Previous directed biosynthesis experiments indicated that the MIA pathway tolerates substituents on the indole ring [8]; thus, it was not surprising to see evidence for incorporation of **8o** into products with molecular masses corresponding to analogs of the known MIAs **1**, **2** (Figure 6A), **12**, and **13** (Supplemental Data).

For this study, we isolated and characterized an alkaloid analog derived from **8m**. A metabolite derived from **8m** was chosen because while tryptamine analogs with substituents on the indole ring have been previously shown by rigorous structural characterization to be widely incorporated into the MIA pathway, the effect of a tryptamine side chain substituent has not yet been explored. Furthermore, while alkaloids derived from **8o** contain a unique isotopic signature from the bromine atom, there is no distinctive isotopic signal for alkaloids derived from **8m**. The major product derived from **8m** was therefore purified and characterized by NMR spectroscopy to validate the formation of a terpene indole alkaloid analog.

Initial analysis of cultures supplemented with **8m** by LC-MS suggested the formation of an analog of a yohimbine-type alkaloid (m/z 385, 353+30, **14m**) and an analog of ajmalicine, **1**, or an isomer of **1** (m/z 383, 353+30, **1m**, Supplemental Data). Additionally, an unknown analog with m/z 426 (396+30), **15m**, was formed (Supplemental Data). The absence of m/z 379 in the media and extracts suggests that the serpentine analog **2m** cannot form from **8m**. This could be due to the absence of a (5-pro-R)-hydrogen atom in the substrate, or the intracellular localization of the biosynthetic intermediates may be altered. A compound corresponding to the analog of the more complex alkaloid **13** was also not observed, suggesting that the (unknown) enzyme forming akuammacine, **13**, does not accept **8m** as a precursor. Isolation, cloning, and reengineering of these enzymes will be key for studies relating to the biosynthesis of these analogs. Interestingly, while the cell culture media did not appear to contain any naturally occurring MIAs, analogs **14m** and **15m** were present in the media. Although relatively little is known about export mechanisms in plant cell culture, it is intriguing to speculate that the plant cell cultures exported these products into the media [26].

Compounds **14m** and **15m**, found in both the media and in root extracts, appeared to be the major unnatural products. Preparative HPLC purification afforded **14m** (2 mg, 3% yield) and **15m** (1 mg, 1%) from 500 ml plant cell culture. Compound **15m** proved intractable to structural assignment, as it was prone to decomposition and consisted of several isomers that could not be separated. The observations of NMR signals for indole aromatic protons strongly suggested assignment as an indole alkaloid (δ 7.41 [d, 1H, J 7.7, H-9], 7.34 [d, 1H, J 8.2, H-12], 7.14 [t, 1H, J 7.6, H-10], 7.04 [t, 1H, J 7.5, H-11]), but no natural alkaloid corresponding to either the proposed parent ion m/z of 396 or the observed m/z 426 has been reported from C. roseus. However, NMR analysis revealed that

14m is an analog of the MIA isositsirikine (Figure 6) [27, 28]. This compound was shown to have an unsubstituted indole (7.5–7.1 ppm, 4H), an intact hydroxymethylene group (4.0 and 3.8 ppm, 2H; compare to 4.0 and 3.8 ppm for **8m**), and the methyl ester (s, 3.8 ppm, 3H). Also, the spectrum revealed an ethylidene group containing a trisubstituted alkene, since one olefinic hydrogen atom appears as a quartet (6.0 ppm, J = 6.8 Hz) coupled to three hydrogen atoms that appear as a doublet (1.9 ppm, J = 6.8 Hz). There are no reports on isolation of isositsirikine from either roots or hairy root cultures of C. roseus, suggesting that unusual biosynthetic precursors may form unexpected products [1].

Large-scale synthesis of the strictosidine analogs (~600 Da) with the expensive secologanin substrate, **7**, limits the scope and rigor of these feeding studies. We anticipate that C. roseus cell lines expressing these STS mutants (using previously reported transformation methods [29]) will have the capacity to synthesize the strictosidine analogs from the more readily available tryptamine starting materials. More detailed analysis of the resulting alkaloid profiles by NMR spectroscopy would therefore be greatly facilitated. Construction of these transgenic cell lines is currently underway.

SIGNIFICANCE

Modulating the substrate specificity of strictosidine synthase (STS) is a critical step toward indole alkaloid pathway engineering to yield nonnatural alkaloids. We developed a streamlined method to expand the range of substrate analogs accepted by STS. The expression system and assay described here were used to identify two STS mutants that yield strictosidine analogs that cannot be produced by the wild-type enzyme. We anticipate that this assay can be applied to identify additional mutants with alternate substrate specificity or optimized selectivity. The assay may also facilitate the identification of functional homologs of STS and the introduction of STS activity in structurally related enzymes. The Val214Met mutant allows for formation of halogenated indole alkaloids, which may be derivatized further by crosscoupling strategies. Inspection of the isotopic pattern of crude cell lysates suggests that a brominated strictosidine analog can be incorporated into several alkaloid metabolites of C. roseus. STS mutant Phe232Leu allows for formation of a strictosidine analog with a modification at an aliphatic carbon. Feeding studies of this strictosidine analog showed that minor modifications to the side chain of tryptamine can cause major perturbations of the alkaloid profile of C. roseus. These results represent a starting point for investigations into how the STS variants can interface with the later steps of the biosynthetic machinery. Future research will seek to understand how these enzyme mutants function in the context of transgenic Madagascar periwinkle cell lines.

EXPERIMENTAL PROCEDURES

General Methods and Analytical Techniques

Secologanin was obtained by methanol extraction of *Lonicera tatarica* as described previously [21]. Tryptamine and tryptamine analogs were purchased in the highest available purity and were used without further purification. Protein was obtained as described below, and the protein concentration was determined by measuring the absorbance at 280 nm (ϵ = 50,300 M^{-1} cm^{-1}, http://www.expasy.ch/tools/protparam.html). NMR spectra were acquired on Varian Inova 500 MHz or Bruker 600 MHz spectrometers.

LC-MS analysis was performed on a Waters LC-MS system. Compounds were separated by an Acquity Ultra-Performance LC (UPLC) BEH C18 column (1.7 μm, 2.1 × 100 mm) equipped in tandem with an MS technologies Micromass LCT Premier with an electrospray ionization (ESI) source and time-of-flight (TOF) detector. The chromatography solvent system consisted of acetonitrile/water (0.1% formic acid). The capillary and sample cone voltages were 2000 V and 30 V, respectively. The desolvation and source temperature were 350°C and 100°C, respectively. The cone and desolvation gas flow rates were 20 l/hr and 700 l/hr, respectively. High-resolution mass spectra (HRMS) were obtained in accurate mass mode (W-mode) with reserpine (*m/z* 609.2812, as 1 nM solution, 40 μl/min) as reference. The HPLC-based assay was carried out on a Beckman-Coulter System Gold system equipped with a 125 solvent module, a 125 autosampler, and a 168 detector. Data were processed (integrated) by using 32Karat (v 7). A solvent system composed of acetonitrile/water (0.1% TFA) was used with a reverse-phase column (HiBar RT 250-4, RP-select B, 5 μm).

Construction of Expression Vectors

The wild-type strictosidine synthase synthetic gene (*sts*) (with codon usage optimized for expression in yeast) was amplified with PCR primers that introduce sites for XhoI and HindIII (underlined) and the FLAG tag epitope (italics): 5′-CCG<u>AAGCTT</u>T CACCAATCTTGAAGA AG-3′, 5′-CTCCTACGTTTCCTCC*GGATCTGACTACAAGGATGACGA CAACAAGTAG*<u>CTCGAG</u>CGG-3′. The primers were designed to eliminate the first 26 residues corresponding to a putative native *N*-terminal signal sequence. The PCR fragment was subcloned into the pGEM-T Easy Vector (Promega, Madison, WI) and then excised and ligated into pGAL-MF (Dualsystems Biotech AG) to obtain pSTSMF-FLAG. The hexahistidine-tagged protein expressed in *E. coli* was obtained by PCR amplification of the *sts* gene in pSTSMF-FLAG by using primers that introduce sites for NcoI and XhoI: 5′-CATG<u>CCATGGG</u>CTCACCAA TCTTGAAGAAGATC-3′, 5′-CCG<u>CTCGAG</u>GGAGGAAACGTAGGAGT TTCCC-3′; it was then ligated into pET-28a(+) to obtain pSTSMF-HIS.

Library Generation and Expression

Primers were designed to overlap with the plasmid sequence to allow for homologous recombination in vivo. Degenerate codons (NNK/ MNN) reduced the library size by 2-fold as compared to complete randomization (NNN/NNN). Two PCR reactions with the pSTSMF-FLAG as a template afforded two *sts* gene fragments. Each fragment contained a region of overlap (>20 bp) with the plasmid at one end and a region of overlap with the other gene fragment (also containing the mutagenized codon) at the opposite end. This overlap allowed for the use of homologous recombination to assemble the fragments into the *sts* mutant in the desired plasmid. Digestion of the parental DNA and agarose gel electrophoresis purification was followed by a splicing-overlap extension PCR reaction to assemble the mutated gene. Homologous recombination of the mutated gene and digested pGAL-MF vector created the saturation mutagenesis libraries. See the Supplemental Data for more information.

Yeast colonies were inoculated in 300 μl SCMM-U (*S. cerevisiae* minimal media, composition per liter: 6.7 g yeast extract without amino acids and 1.92 g yeast supplemental media without uracil) supplemented with 2% glucose and incubated overnight at 30°C in 96-well, square-bottom, deep-well plates (2 ml volume capacity per well) with shaking at 225 rpm. An aliquot of the overnight culture (100 μl) was then used to inoculate 900 μl SCMM-U supplemented with 0.5% glucose and 2% galactose and incubated for an additional 72–96 hr at 20°C. Each well contained one glass bead (5 mm) to facilitate aeration, and all plates were protected from contamination during the incubation by sterile, breathable sealing film. The plates were centrifuged (10°C, 3000 × g, 10 min), and the supernatant was used directly for screening. We verified that the supernatant could be stored with cells at 4°C for at least 4 weeks without noticeable loss of activity. 96-well plates can be recultured in SCMM-U media or supplemented with 25% glycerol and stored at −80°C for future use.

Kinetic Analysis of STS Variants

Reaction conditions were modified to obtain less than 15% conversion for all data points used in steady-state kinetic analysis. All enzymatic reactions (0.100 ml) contained secologanin (4.0 mM) in sodium phosphate buffer (100 mM [pH 7.0]) with internal standard (0.075 mM naphthyl acetic acid) and one of several tryptamine or tryptamine analog concentrations (always added as a 10-fold dilution from stock solution). Tryptamine or tryptamine analog concentrations were at least 2-fold above and below the estimated K_M value. The HPLC detection limit did not allow for substrate concentrations lower than 1 μM. Formation of strictosidine was monitored at 280 nm by using the experimentally determined extinction coefficient of 5,140 μM^{-1}. Wild-type reactions were quenched with two equivalents of sodium hydroxide due to the fast reaction between tryptamine and secologanin. This quench resulted in the quantitative formation of strictosamide ($C_{26}H_{31}N_2O_8$ calculated: 499.2091, found: 499.2080) from strictosidine, and the extinction coefficient of the strictosamide product was experimentally determined to be 12,000 μM^{-1}. Plotted experimental data were fitted to the Michaelis-Menten and Northrop equations by nonlinear regression analysis by using Origin (v 7.0552) (OriginLab Corp., Northampton, MA) to directly afford V_{max}, V_{max}/K_M, and K_M. All kinetic assays were repeated three times to ensure reproducibility. Representative raw data for the Val214Met mutant and 5-methyltryptamine are shown in the Supplemental Data.

Feeding Study and Alkaloid Isolation

Hairy root cell cultures of *C. roseus* were inoculated in half-strength Gamborg media and were grown for 3 weeks. Enzymatic synthesis and purification afforded **8m** (ESI-MS $C_{28}H_{36}N_2O_{10}$: calculated *m/z* 561.2448 [M+H]+, found *m/z* 561.2448 [M+H]+), and chemical synthesis and purification afforded **8o** (ESI-MS $C_{27}H_{34}N_2O_9{}^{79}Br$: calculated *m/z* 609.1448 [M+H]+, found *m/z* 609.1456 [M+H]+; ESI-MS $C_{27}H_{34}N_2O_9{}^{81}Br$: calculated *m/z* 611.1427 [M+H]+, found *m/z* 611.1453 [M+H]+). Each strictosidine analog was dissolved in water to a final concentration of 0.5 mM, and the solution was filter sterilized and then added to the hairy root culture. After 1 week, the media were separated from the hairy roots, and the roots were washed with water three times. The media and roots were extracted according to standard procedures (Supplemental Data) [25, 28]. The extracted alkaloids were analyzed by LC-MS, and the two major species were isolated by using preparative HPLC and were analyzed by NMR.

NMR Characterization of Enzymatically Generated Strictosidine Analogs

(5R)-Hydroxymethylstrictosidine, 8m

^{1}H NMR (methanol-d4, 500 MHz) δ 7.80 (s, 1H, H-17), 7.46 (td, 1H, $^3J_{10}$ 7.8, $^4J_{11}$ 1.0, H-9), 7.33 (td, 1H, $^3J_{11}$ 8.2, $^4J_{10}$ 0.84, H-12), 7.15 (ddd, 1H, $^3J_{10}$ 7.1, $^3J_{12}$ 8.2, $^4J_9$ 1.1, H-11), 7.05 (ddd, 1H, $^3J_{11}$ 7.1, $^3J_9$ 8.2, $^4J_{12}$ 0.92, H-10), 5.84 (ddd, 1H $^3J_{18trans}$ 17.4, $^3J_{18cis}$ 10.6, $^3J_{20}$ 7.6, H-19), 5.80 (d, 1H, $^3J_{20}$ 8.8, H-21), 5.37 (ddd, 1H, $^3J_{19}$ 17.4, $^2J_{18cis}$ 1.3, $^4J_{20}$ 1.0, H-18*trans*), 5.28 (ddd, 1H, $^3J_{19}$ 10.6, $^2J_{18trans}$ 1.3, $^4J_{20}$ 1.0, H-18*cis*), 4.78 (d, 1H, $^3J_{2'}$ 7.9, GlcH-1′), 4.67 (dd, 1H, $^3J_{14proR}$ 10.9, $^3J_{14proS}$ 4.0, H-3), 4.03 (dd, 1H, $^3J_{23}$ 10.7, $^3J_{5a}$ 3.4, H-23), 3.94 (dd, 1H, $^2J_{6'b}$ 11.8, $^3J_{5'}$ 2.1, GlcH-6′a), 3.84–3.75 (m, 2H, H-5a, H-23), 3.78 (s, 3H, H-24), 3.62 (dd, 1H, $^2J_{6'a}$ 11.9, $^3J_{5'}$ 6.9, GlcH-6′b), 3.40 (t$_{app}$, 1H, $^3J_{2',4'}$ 9.1, GlcH-3′), 3.36–3.32 (m, 1H, GlcH-5′), 3.25–3.19

(m, 2H, GlcH-2′,4′), 3.12 (dd, 1H, $^2J_{6b}$ 16.3, $^3J_{5a}$ 4.8, H-6a), 3.04 (td, 1H, $^3J_{14proS}$ 11.2, $^3J_{20,14proR}$ 4.6, H-15), 2.89 (dd, 1H, $^2J_{6a}$ = 16.4, $^3J_{5a}$ 8.9, H-6b), 2.77 (ddd, 1H, $^3J_{15}$ 5.0, $^3J_{19}$ 7.8, $^3J_{21}$ 8.6, H-20), 2.32 (ddd, 1H, $^3J_{15}$ 4.4, $^3J_3$ 11.1, $^2J_{14proS}$ 15.2, H-14proR), 2.24 (ddd, 1H, $^3J_3$ 4.2, $^3J_{15}$ 11.4, $^2J_{14proR}$ 15.2, H-14proS); ^{13}C NMR (methanol-d_4, 125 MHz) δ 171.28, 156.93, 138.45, 135.51, 130.46, 127.50, 123.79, 120.76, 120.08, 119.24, 112.48, 108.84, 106.25, 100.56, 97.54, 78.82, 78.08, 74.78, 71.82, 63.02, 62.03, 54.10, 52.76, 51.16, 45.34, 35.38, 32.63, 21.89; ESI-MS ($C_{28}H_{37}N_2O_{10}^+$): calculated m/z 561.2448 [M+H]$^+$, found m/z 561.2448 [M+H]$^+$.

10-Methylstrictosidine, 8l
^{1}H NMR (methanol-d_4, 500 MHz) δ 7.81 (s, 1H, H-17), 7.26 (s, 1H, H-9), 7.19 (d, 1H, $^3J_{11}$ 8.3, H-12), 6.98 (dd, 1H, $^4J_9$ 1.3, $^3J_{12}$ 8.4, H-11), 5.85 (ddd, 1H, $^3J_{20}$ 7.6, $^3J_{18cis}$ 10.6, $^3J_{18trans}$ 17.5, H-19), 5.84 (d, 1H, $^3J_{20}$ 9.1, H-21), 5.35 (dd, 1H, $^2J_{18cis}$ 1.3, $^3J_{19}$ 17.4, H-18trans), 5.28 (dd, 1H, $^2J_{18trans}$ 1.1, $^3J_{19}$ 10.6, H-18cis), 4.80 (d, 1H, $^3J_{2'}$ 7.9, GlcH-1′), 4.64 (br-d, 1H, $^3J_{14proR,14proS}$ 10.6, H-3), 3.98 (dd, 1H, $^3J_{5'}$ 2.1, $^2J_{6'b}$ 11.8, GlcH-6′a), 3.80 (s, 3H, H-23), 3.73 (ddt, 1H, $^3J_{5'}$ 5.0, $^2J_{6'a}$ 12.5, GlcH-6′b), 3.49–3.42 (m, 1H, H-5a), 3.40 (t$_{app}$, 1H, $^3J_{2',4'}$ 9.1, GlcH-3′), 3.39–3.35 (m, 1H, H-5b), 3.25–3.19 (m, 2H, GlcH-2′, GlcH-4′), 3.10–2.97 (m, 3H, H-6a, H-6b, H-15), 2.75 (ddd, 1H, $^3J_{15}$ 5.1, $^3J_{19}$ 7.8, $^3J_{21}$ 8.7, H-20), 2.40 (s, 3H, H-23), 3.36–2.16 (m, 2H, H-14proS, H-14proR); ^{13}C NMR (methanol-d_4, 125 MHz) δ 171.43, 157.05, 136.67, 135.62, 130.30, 130.03, 127.82, 125.24, 119.90, 118.91, 112.10, 109.08, 106.79, 100.56, 97.43, 78.96, 78.135, 74.80, 71.88, 63.15, 53.19, 52.74, 45.52, 42.80, 34.92, 32.75, 21.71, 19.71; ESI-MS ($C_{28}H_{37}N_2O_9^+$): calculated m/z 545.2499 [M+H]$^+$, found m/z 545.2523 [M+H]$^+$.

10-Chlorostrictosidine, 8n
^{1}H NMR (methanol-d_4, 500 MHz) δ 7.81 (s, 1H), 7.47 (dd, 1H, $^5J_{12}$ 0.6, $^4J_{11}$ 2.1, H-9), 7.29 (dd, 1H, $^5J_9$ 0.6, $^3J_{12}$ 8.7, H-12), 7.11 (dd, 1H, $^4J_9$ 2.0, $^3J_{12}$ 8.7, H-11), 5.85 (ddd, 1H, $^3J_{20}$ 7.7, $^3J_{18trans}$ 10.7, $^3J_{18trans}$ 17.4, H-19), 5.84 (d, 1H, $^3J_{20}$ 9.1, H-21), 5.35 (dt$_{app}$, 1H, $^2J_{18cis}$ 1.4, $^4J_{20}$ 1.4, $^3J_{19}$ 17.4, H-18trans), 5.28 (dt$_{app}$, 1H, $^2J_{18trans}$ 1.3, $^4J_{20}$ 1.3, $^3J_{19}$ 10.7, H-18cis), 4.80 (d, 1H, $^3J_{2'}$ 7.9, H-1′), 4.68 (br-d, 1H, $^3J_{14proS,R}$ 10.4, H-3), 3.98 (dd, 1H, $^3J_{5'}$ 2.0, $^2J_{6'b}$ 11.8, H-6′a), 3.80 (s, 3H, H-23), 3.74 (td, 1H, $^3J_{6ax,6eq}$ 5.2, $^2J_{5ax}$ 12.5, H-5eq), 3.64 (dd, 1H, $^3J_5$ 7.1, $^2J_{6'a}$ 11.8, H-6′b), 3.47 (ddd, 1H, $^3J_{6eq}$ 5.6, $^3J_{6ax}$ 8.5, $^2J_{5eq}$ 12.5, H-5ax), 3.40 (t$_{app}$, 1H, $^3J_{2',4'}$ 9.1, H-3′), 3.39–3.34 (m, 1H, H-5′), 3.23 (dd, 1H, $^3J_{3'}$ 8.9, $^3J_{5'}$ 9.9, H-4′), 3.22 (dd, 1H, $^3J_{1'}$ 7.9, $^3J_{3'}$ 9.3, H-2′), 3.13–3.05 (m, 2H, H-6ax, H-15), 3.02 (td$_{app}$, 1H, $^3J_{5ax,5eq}$ 4.5, $^2J_{6ax}$ 16.3, H-6eq), 2.75 (ddd(d), 1H, $^4J_{18cis,trans}$ < 0.5, $^3J_{15}$ 5.0, $^3J_{19}$ 7.8, $^3J_{21}$ 8.8, H-20), 2.32 (ddd, 1H, $^3J_{3/15}$ 3.0, $^3J_{3/15}$ 12.1, $^2J_{14proS}$ 15.0, H-14proR), 2.23 (ddd, 1H, $^3J_{3/15}$ 3.7, $^3J_{3/15}$ 11.8, $^2J_{14proR}$ 15.2, H-14proS); ^{13}C NMR (methanol-d_4, 125 MHz) δ 171.40, 157.06, 136.70, 135.54, 132.19, 128.65, 126.51, 123.80, 119.97, 118.84, 113.67, 109.03, 107.19, 100.54, 97.41, 78.95, 78.12, 74.79, 71.87, 63.15, 53.05, 52.74, 45.50, 42.63, 34.82, 32.68, 19.52; ESI-MS ($C_{27}H_{34}N_2O_9^{35}Cl^+$): calculated m/z [M+H]$^+$ 565.1953, found m/z 565.1959; ($C_{27}H_{34}N_2O_9^{37}Cl^+$): calculated m/z [M+H]$^+$ 567.1923, found m/z 567.1945.

NMR Characterization of Isositsirikine Alkaloid Analog, 14m
^{1}H NMR (600 MHz, methanol-d_4) δ 7.51 (d, 1H, $^3J_{10}$ 7.7, H-9), 7.38 (d, 1H, $^3J_{11}$ 7.6, H-12), 7.26 (t, 1H, $^3J_{9,11}$ 7.6, H-10), 7.09 (t, 1H, $^3J_{10,12}$ 7.4, H-11), 5.96 (q, 1H, $^3J_{18}$ 6.8, H-19), 4.94–4.87 (m, 1H, H-3), 4.20 (d, 1H, $^2J_{21b}$ 14, H-21a), 4.13 (d, 1H, $^2J_{21a}$ 14, H-21b), 4.04 (dd, 1H, $^3J_{16}$ 4.0, $^2J_{17b}$ 12, H-17a), 3.82 (dd, 1H, $^3J_{16}$ 4.7, $^2J_{17a}$ 12, H-17b), 3.99–3.85 (m, 2H, H-23a, H-23b), 3.76 (s, 3H, H-22), 3.67–3.58 (m, 1H, H-5), 3.20 (dd, 1H, $^3J_5$ 7.0, $^2J_{6b}$ 14, H-6a), 3.17–3.09 (m, 1H, H-16), 2.98 (dd, 1H, $^3J_5$ 7.0, $^2J_{6a}$ 15, H-6b), 2.87–2.82 (m, 1H, H-15), 2.59–2.42 (m, 2H, H-14), 1.87 (d, 3H, $^3J_{19}$ 6.8, H-18); ESI-MS ($C_{22}H_{29}N_2O_4^+$): calculated m/z [M+H]$^+$ 385.2127, found m/z 385.2134.

Supplemental Data
Supplemental Data include a detailed protocol for *sts* gene construction and expression, representative data for kinetic analysis, a protocol for the generation of saturation mutation libraries, control experiments for assay and MS and NMR data for chromophores, a protocol for enzyme screening in 96-well plates, protocols for enzymatic synthesis of

strictosidine analogs, a detailed procedure for feeding studies, and LC-MS data of feeding experiments and are available at http://www.chembiol.com/cgi/content/full/14/8/888/DC1/.

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