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Recent progress in the metabolic engineering of alkaloids in plant systems

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Plant alkaloids have a rich chemical ecology that has been exploited for medicinal purposes for thousands of years. Despite being highly represented within today's pharmacopoeia, relatively little is known about the biosynthesis, regulation and transport of these molecules. Understanding how nature synthesizes plant alkaloids will enhance our ability to overproduce—that is, to metabolically engineer—these medicinally useful compounds as well as new-to-nature compounds (with potentially improved bioactivity) derived from these natural scaffolds. Recent progress in the metabolic engineering of nitrogen-containing plant natural products—specifically the monoterpene indole alkaloids, the benzyloquinoline alkaloids and the glucosinolates—was made possible through the characterization of various components in both native and engineered enzymatic pathways. The subsequent reconfiguration and tuning of these biological 'parts' has enabled the production of selected products at increasingly higher titers.

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Introduction

Plant natural products have been exploited for thousand of years. The alkaloids, particularly, have a long and storied narrative. This history is highlighted in the life of Cleopatra, who used alkaloid-containing extracts from *belladonna* (Italian for 'beautiful woman') to dilate her pupils so as to increase her beauty and thereby disarm her enemies [1]. Far from being confined to ancient chronicles, the alkaloids retain a palpable presence in

today's clinics. For example, optometrists still apply eye drops containing the alkaloid atropine, an active component of *belladonna*, to dilate the pupil during routine eye exams [1]. It is unsurprising that most alkaloids are bioactive given that the evolutionary process selects for products that confer an advantage to the producing organism. Despite the rich ethnopharmacological tradition and high usage of alkaloids in the modern era, relatively little is known about the biosynthesis, regulation and transport of these molecules. This lack of knowledge fundamentally impedes our ability to co-opt nature's machinery in order to overproduce—that is, to metabolically engineer—these valuable compounds.

Notably, many drug screening efforts exclude plant natural products because of their high production costs and instead screen larger numbers of simpler molecules, which can be produced inexpensively and in fewer chemical steps [2]. Given the challenges of getting these molecules through the drug pipeline, we contend that natural products—including plant alkaloids—should be included in drug screens. Plant natural products have a high success rate as candidates and leads [3,4]. While the chemical syntheses of plant natural products, particularly the alkaloids, are dramatically improving [5], many syntheses are still too lengthy for commercial production or require industrially impractical separation steps. Therefore, alternative production platforms must be developed, evaluated and instituted. An increasing body of work enlists microbes as well as cell and tissue cultures to produce these valuable plant-derived products [4]. Biological systems have the potential to be scalable and selective, while simultaneously being more environmentally friendly and—importantly—less expensive than synthetic reactions [4].

In this Opinion, we highlight recent metabolic engineering efforts designed to improve production of selected plant-derived alkaloids. We focus on the monoterpene indole alkaloids (MIAs), the benzyloquinoline alkaloids (BIAs) and the glucosinolates. Though not classically classified as alkaloids, the glucosinolates are nitrogen-containing compounds that have been the subject of a compelling body of research that will inform the forward engineering of all plant natural products. These three classes of plant-derived nitrogen-containing natural products have been the subject of recent research efforts aimed at discovering and manipulating cellular activities, which include enzymatic function, metabolite transport and regulatory control. Ultimately this work may lead to biotechnologically useful enzymes and new

drug candidates. Throughout this opinion, we also highlight the challenges that arise in attempting to chart the underexplored landscape of plant biosynthesis and discuss the grand challenges that remain in the metabolic engineering of plant alkaloid products.

The monoterpene indole alkaloids

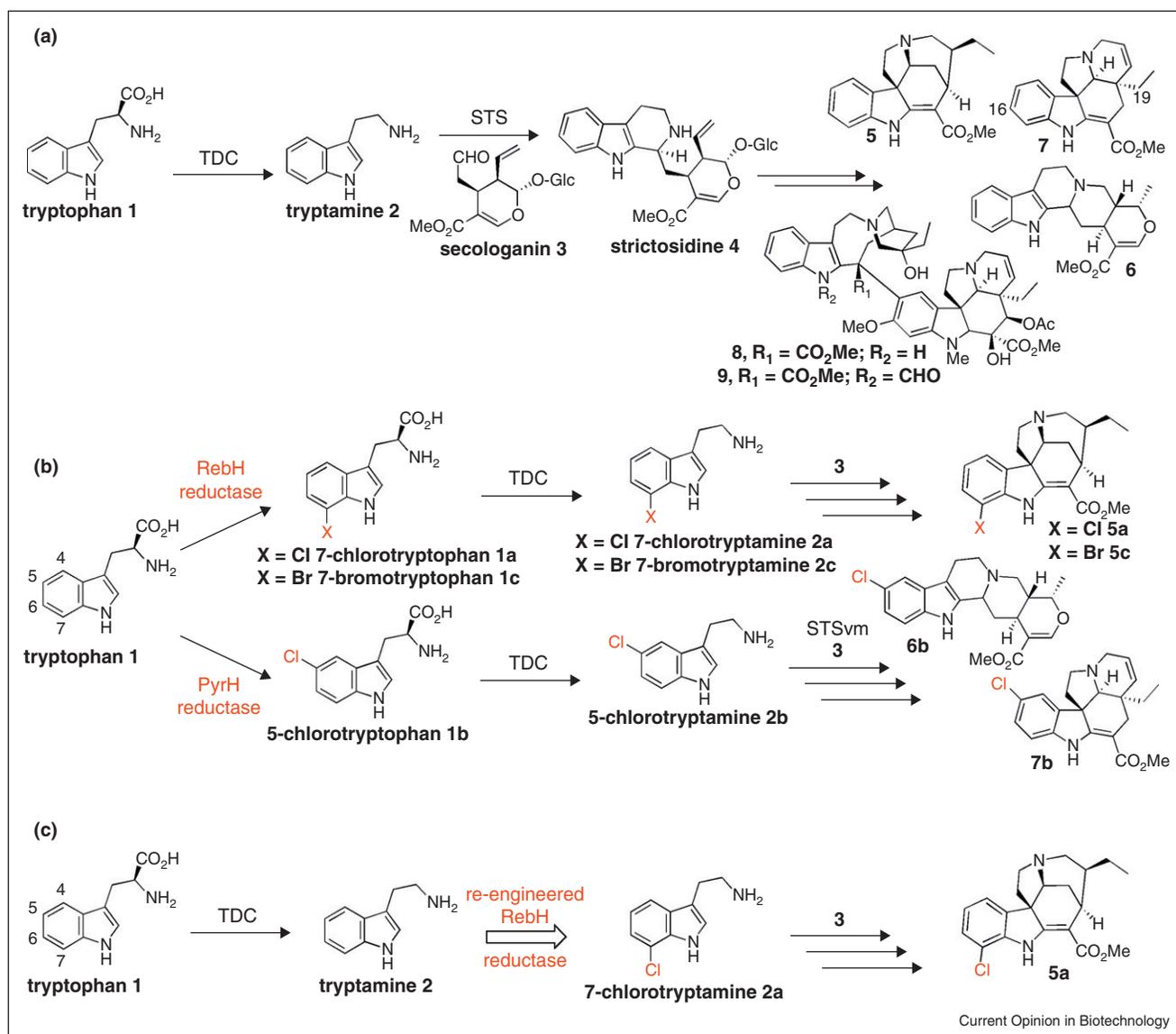
Introduction

The monoterpene indole alkaloids (MIAs) have garnered interest over the past few decades largely because of vinblastine **8** and vincristine **9**, two potent and widely prescribed anti-cancer agents that are currently produced solely through harvest from the leaves of mature periwinkle plants (*Catharanthus roseus*) [6]. The

concentrations of vinblastine and vincristine per gram of dry leaf material are approximately 0.01% and 0.003%, respectively, and are greatly dependent upon plant growth conditions [7]. Their low yields and lengthy production timeline have elicited intense efforts to engineer higher titers of these medicinally important MIAs.

The MIAs are encountered most commonly in the Apocynaceae, Loganiaceae and Rubiaceae families [8]. Most MIAs are built from the secoiridoid secologanin **3** and the indole-containing molecule tryptamine **2** (Figure 1a) [9]. Strictosidine synthase (STS) condenses these two molecules via a Pictet-Spengler condensation that forms

Figure 1



(a) The monoterpene indole alkaloid (MIA) pathway. TDC, tryptophan decarboxylase; STS, strictosidine synthase; Glc, glucose. (b) Introduction of halogenation into the MIA pathway. RebH and PyrH are both flavin-dependent halogenases from actinomycetes species; STS_{vm}, strictosidine synthase Val214Met mutant. (c) Reengineering of halogenase to preferentially chlorinate tryptamine **2** over the natural substrate tryptophan **1**.

strictosidine 4, which is believed to ultimately succumb to either of two chemical fates [9,10]. If the plant is not under herbivore attack, strictosidine 4 is deglycosylated and rearranged into the over 3000 MIAs found in nature [10]. Madagascar periwinkle has a subset of approximately 130 MIAs. Alternatively, if the plant is under herbivore attack, the strictosidine 4 pool (estimated to be approximately 10 mM in periwinkle leaf epidermal cells hormonally treated to mimic herbivore attack) can be directed to the nucleus for mass deglycosylation, leading to a reactive dialdehyde species capable of cross-linking proteins [10]. This mechanism has been dubbed the strictosidine nuclear ‘bomb’ in reference to the ‘mustard oil bomb’ mechanism of glucosinolate biosynthesis (see below) [10]. Importantly, many of the MIA metabolites themselves have also been implicated in plant defense strategies [11].

Obtaining the building blocks for metabolic engineering efforts—A case study on the discovery of P450s involved in MIA biosynthesis

The enzymatic pathways leading to the MIAs have not yet been fully elucidated in any organism. These uncharacterized biochemical steps may utilize novel chemistries or possess informative and interesting specificities that enable the enzymes to be employed in various synthetic metabolic pathway designs [12,13]. Notably, many plants—including MIA producers—are predicted to contain a high percentage of cytochromes P450 (P450s). Some estimates place P450s at approximately 1% of representative plant genomes [14], over 5-fold higher than the proportion of P450s found in the human genome [15]. By using molecular oxygen to tailor hydrocarbon skeletons, P450s facilitate a panel of difficult chemical transformations and are consequently utilized in many alkaloid biosynthetic pathways [13]. P450s have also been successfully engineered for biotechnological purposes [15,16]. Various technologies, such as nanodiscs [17], and N-terminus reengineering efforts [16] have improved the expression of membrane-bound P450s, making this class of enzymes accessible to a full suite of biochemical and biophysical characterization techniques.

Given the high sequence similarity of P450s, identifying a P450 that facilitates a specific biochemical reaction within a biosynthetic pathway remains a challenge. This has greatly slowed the discovery and characterization of new P450s within the plant kingdom. However, Giddings *et al.* recently used co-expression analysis to identify P450s with expression profiles similar to known MIA biosynthetic genes [12]. By functionally assaying these candidates in *Saccharomyces cerevisiae*, Giddings *et al.* discovered one P450 (CYP71BJ1) that hydroxylated the 19 position of either lochnericine or tabersonine 7, an intermediate that is positioned at a metabolic branch point [12]. Hydroxylation of tabersonine 7 at the 16 position commits the intermediate to vindoline and

vinblastine 8 biosynthesis, whereas hydroxylation at the 19 position commits the molecule to 19-O-acetyl-hörhammericine formation [12].

Engineering ‘unnatural’ natural products

Plant alkaloids often require modification to improve their pharmacological properties prior to human consumption. Halogenation, particularly fluorination and chlorination, is a pervasive modification in successful pharmaceutical candidates [18,19]. Halogens often confer the potency of a drug, alter its pharmacokinetics or function as site-specific handles for subsequent modification [18–20]. Halogens can be introduced into MIA pathways by a number of methods. One particular example employed mutasynthesis, a process whereby natural biosynthesis is first blocked by genetic silencing of the natural precursor, and then rescued by feeding with structural analogs of the precursor [21]. In this case, in conjunction with the RNAi-mediated knockdown of tryptophan decarboxylase (TDC), unnatural tryptamine 2 analogs were added to a chemically ‘silent’—non-alkaloid producing—background and fluorinated MIA analogs were observed [21]. In a separate engineering strategy, strictosidine synthase (STS)—the enzyme situated at the first committed step of MIA biosynthesis—was engineered to accept an expanded range of halogenated tryptamine 2 precursors [22]; the utility of this enzyme was demonstrated *in planta* by feeding previously unaccepted unnatural precursors to *C. roseus* hairy roots [23].

Finally, Runguphan *et al.* interfaced RebH and PyrH—two tryptophan halogenases isolated from soil-dwelling actinomycetes species—with the MIA metabolism of periwinkle to produce halogenated natural products *de novo* [24**] (Figure 1b). However, the RebH-overexpressing and RebF-overexpressing lines also displayed a brown and slow growth morphology [24**]. Runguphan *et al.* hypothesized that this morphology was the result of the accumulation of 7-chlorotryptan 1a, an analog of primary metabolite L-tryptophan 1 that is somewhat structurally similar to 4-chloroindole-3-acetic acid, an auxin known to be involved in regulating plant growth [24**].

To circumnavigate this problem, Glenn *et al.* employed structure-guided protein design to engineer a halogenase that preferentially chlorinated tryptamine 2, a more direct MIA precursor (Figure 1c) [25]. Microgram per gram fresh weight quantities of 12-chloro-19,20-dihydroakuamicine 5a were observed with this strategy, but neither 7-chlorotryptophan 1a nor 7-chlorotryptamine 2a accumulated in *planta*, indicating the chlorinated precursor was being effectively shuttled into MIA metabolism [25]. Engineering halogenation into MIA metabolism highlights an important need to interface specialized metabolism with primary carbon and nitrogen metabolism.

The benzyloquinoline alkaloids

Introduction

Benzyloquinoline alkaloids (BIA) are found mainly in the Papaveraceae, Ranunculaceae, Berberidaceae and Menispermaceae plant families. Approximately 2500 BIAs have been isolated to date [26]. This class of compounds has been used throughout human history and contains pharmaceuticals that are still widely used today, including the narcotic and analgesic morphine 28, the cough suppressant codeine 26, the muscle relaxant papaverine, and the anti-microbial agents sanguinarine 19 and berberine 29. All known BIAs, like the MIAs, are derived from a single intermediate, which, for this class of compounds, is norcoclaurine 13. Norcoclaurine synthase (NCS) catalyzes the condensation between dopamine 11 and 4-hydroxyphenylacetaldehyde 12 to yield the central intermediate, norcoclaurine 13. Notably, the biosynthetic pathways of several benzyloquinoline alkaloids—morphine 28, sanguinarine 19 and berberine 29—have been fully elucidated at the genetic level, which has enabled sophisticated metabolic engineering approaches. The application of metabolic engineering strategies for BIAs has focused predominantly on improving the yields of specific alkaloid compounds that exhibit medicinal value.

Enzyme discovery and engineering in BIA pathways

Several outstanding efforts in enzyme discovery have been reported for BIA biosynthetic pathways. In a recent effort, Hagel *et al.* characterized two O-demethylases that are involved in morphine biosynthesis, completing the characterization of the morphinan pathway (Figure 2b) [27]. This work also clearly highlighted how co-expression analysis can be used to discover enzymes with unprecedented catalytic function. These enzymes offer the first examples of non-heme iron(II) oxoglutarate dioxygenases capable of catalyzing O-demethylation. Codeine-O-demethylase (CODM) regioselectively demethylates codeine 26 and thebaine 22 at the 3-position, while thebaine-6-O-demethylase (T6ODM) demethylates thebaine 22 and oripavine 24 at the 6-position. Swapping amino acid regions between the two demethylases resulted in a CODM mutant that selectively demethylates codeine (Figure 2b) [28]. This mutant—which effectively sidesteps oripavine 24 production by committing thebaine 22 to just one of two possible routes—could potentially impact titers of codeine 26 and morphine 28 in subsequent metabolic engineering efforts. Collectively, these studies highlight how characterizing individual pathways steps and understanding their specificity and selectivity can both inform and enable metabolic engineering efforts.

While transcript analysis has proven to be spectacularly successful in elucidating the demethylases of morphine 28 biosynthesis, Winzer *et al.* provide a rare example of gene clustering in a BIA pathway [29]. The authors describe a 10-gene cluster in the poppy genome that putatively encodes the entire biosynthetic pathway of

the BIA noscapine 15. This is the first gene cluster discovered for an alkaloid pathway, and it is the largest plant gene cluster discovered to date. The authors further successfully silenced six of the ten proposed genes using VIGS to validate their role in noscapine 15 biosynthesis [29]. This study indicates that genomic data, in addition to expression data, can be used to decipher alkaloid pathways in plants.

Engineering in native hosts

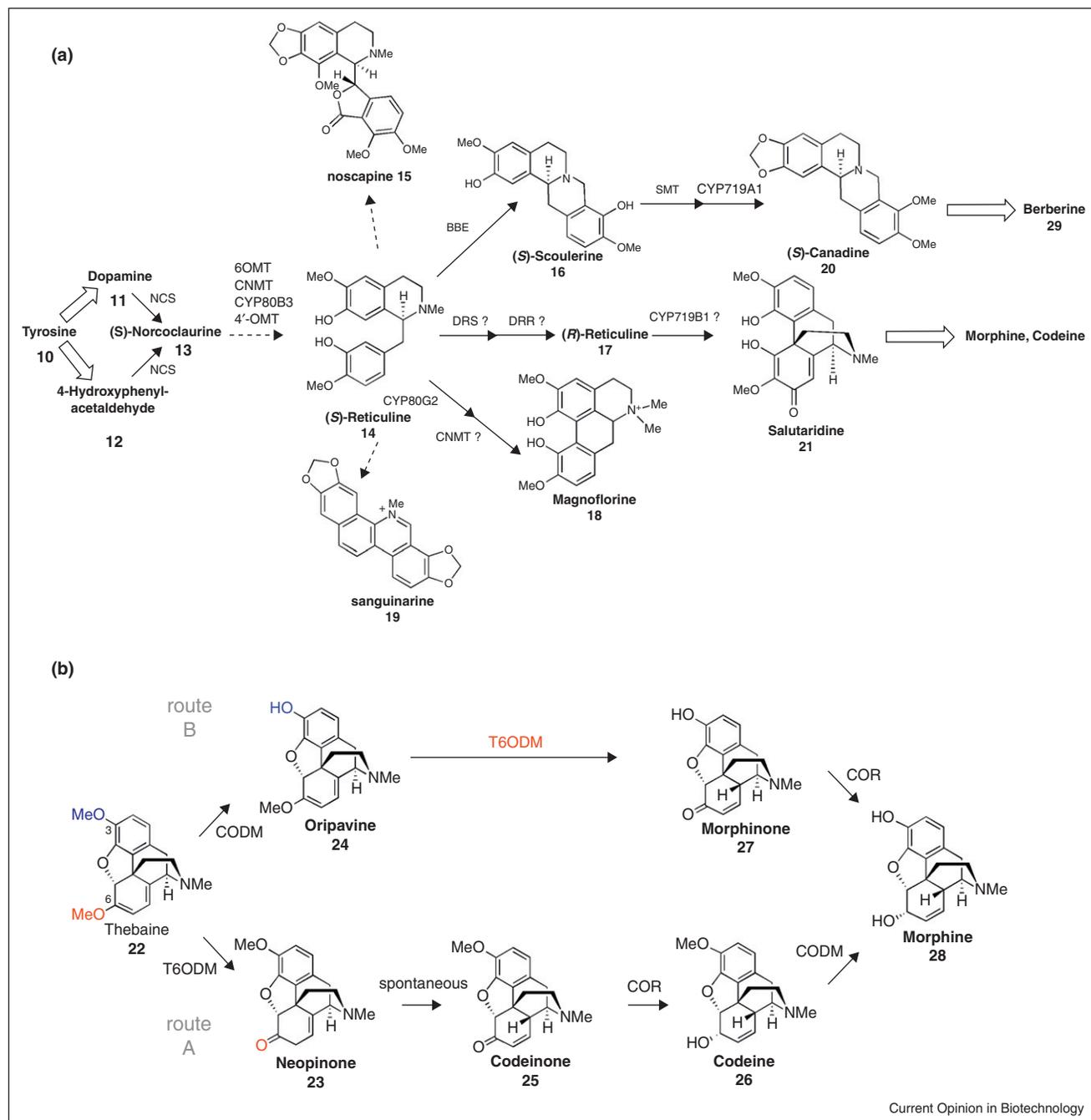
In one of the earliest attempts to engineer BIA-producing plants, RNA interference (RNAi) was used to silence the expression of codeinone reductase (COR) in the opium poppy [30]. COR, the penultimate enzyme of morphine biosynthesis, converts codeinone 25 to codeine 26 (Figure 2b). While one might anticipate that the silencing of COR would lead to elevated levels of codeinone 25, the study found instead that COR-silenced plants accumulated reticuline 14—an intermediate seven steps upstream of codeinone 25—at the expense of morphine 28, codeine 26, oripavine 24 and thebaine 22. A feedback mechanism was proposed as an explanation for the elevated levels of reticuline 14, though testing this hypothesis has yielded conflicting results [30].

Other early attempts to improve the yields of BIA alkaloids include the overexpression of berberine bridge enzyme (BBE) in *Eschscholzia californica* root cultures. This effort resulted in elevated levels of downstream alkaloids and decreased levels of amino acids, though notably levels of tyrosine 10—the amino acid employed in BIA synthesis—were unaltered [31]. Conversely, the antisense suppression of BBE expression led to the effective silencing of BIA production and increased cellular amino acid levels, though, again, tyrosine 10 levels went largely unchanged (less than two-fold higher than in control lines) [32]. Nonetheless, these two studies highlight how perturbations in alkaloid metabolism can impact primary metabolism [31,32]. More recent studies suggest, however, that the RNAi suppression of BBE in *E. californica* leads to increased accumulations of (S)-reticuline 14 instead of various canonical amino acids [33]. These contradictory results are surprisingly common in the metabolic engineering of alkaloids in plants and cell cultures and provide us with the impetus to understand these pathways in greater detail, paying specific attention to their biochemical and molecular regulatory elements.

Reconstituting BIA biosynthesis into microbial systems

Many pathways in BIA biosynthesis are fully characterized, which opens the possibility of transplanting entire alkaloid pathways into microbial hosts. Though relatively difficult, the reconstitution of entire metabolic pathways into microbial hosts confers a number of advantages, including rapid biomass accumulation, facile purification and access to the host of tools available for workhorse

Figure 2



(a) The benzylisoquinoline alkaloid (BIA) pathway. NCS, norcoclaurine synthase; 6-OMT, norcoclaurine 6-O-methyltransferase; CNMT, coclaurine-N-methyltransferase; 4'-OMT, 3'-hydroxy-N-methylcoclaurine-4'-O-methyltransferase; DRS, 1,2-dehydroreticulene synthase; DRR, 1,2-dehydroreticulene reductase; BBE, berberine bridge enzyme; SMT, scoulerine 9-O-methyltransferase; MAO, bacterial monoamine oxidase; CYP2D6, human cytochrome P450 enzyme; CYP80G2, CYP719B1 and CYP719A1, plant cytochrome P450 enzymes. **(b)** The morphinan alkaloid pathway. CODM, codeine O-demethylase; T6ODM, thebaine 6-O-demethylase; COR, codeinone reductase.

organisms like *E. coli* and *S. cerevisiae*. A number of recent reports have successfully reconstituted portions of BIA pathways into *S. cerevisiae*, *E. coli* and combinations thereof in co-culture systems [34–36]. For example,

Hawkins *et al.* were able to produce reticuline 14 as well as sanguinarine/berberine-type and morphinan-type BIAs in yeast by overexpressing genes from mixed plant sources and human [35]. Notably, they were also able

to tune enzyme expression levels through use of a glucocorticoid-inducible promoter and *in situ* promoter titration [35]. This level of tuning enables maximal pathway flux and minimal enzyme expression. The expression system is nominally taxed under these conditions, since valuable cellular resources are not used on the biosynthesis of supernumerary proteins and nucleic acids.

The glucosinolates

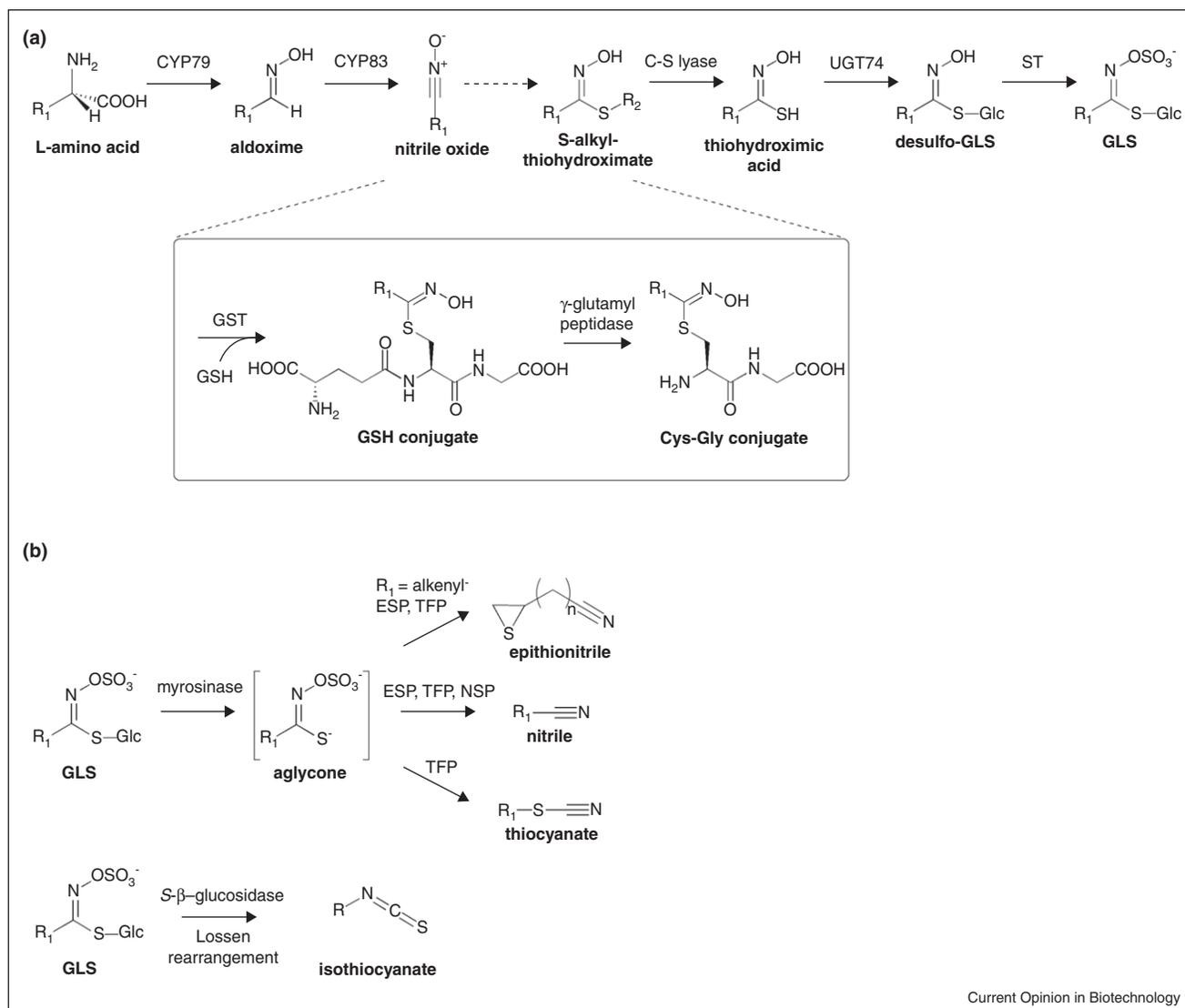
Introduction

The glucosinolates are not classified as alkaloids, although, along with the alkaloids, these compounds are amino acid-derived, nitrogen-containing small molecules of plant origin. The glucosinolates are included in this review because the recent and creative metabolic engineering studies performed on this class of compounds

will undoubtedly inform the forward engineering of all plant natural products, including the alkaloids.

Glucosinolates are sulfur-containing and nitrogen-containing compounds that are derived from glucose and various amino acids (Figure 3a) [37]. They are found in cruciferous vegetables (the Brassicaceae plant family) and have been shown to possess a range of bioactivities [37]. The glucosinolates occupy an essential space in the chemical ecology of their host organisms by attracting specialist crucifer pollinators and insects and deterring predatory herbivores [38]. Specifically, crucifers employ myrosinases (hydrolases) to cleave the glucose moiety of glucosinolates in response to predation and herbivory (Figure 3b) [39]. The myrosinases and glucosinolates are physically separated, coming into contact only upon

Figure 3



(a) The glucosinolate (GLS) pathway. GSH, glutathione. (b) Glucosinolate hydrolysis to form epithionitrile, nitrile, thiocyanate and isothiocyanate.

disruption of the plant tissue (Figure 3b) [39]. Upon hydrolysis, the resultant unstable aglycone intermediate spontaneously rearranges into the corresponding isothiocyanate via a Lossen-type rearrangement [39]. The three known types of specifier proteins, Thiocyanate-Forming Proteins (TFPs), Nitrile-Forming Proteins (NFPs) and Epithiospecifier Proteins (ESPs)—which can be found *in planta* or in various specialist insects—can redirect glucosinolate hydrolysis from isothiocyanate products toward thiocyanate, simple nitrile and epithionitrile products, respectively (Figure 3b) [39]. Notably, many specifiers can direct glucosinolate hydrolysis to more than one product [39]. Early workers on this plant defense and pollination system dubbed it ‘The Mustard Oil Bomb’ [38]. To date, over 120 glucosinolates have been identified [40].

Improving yield in non-native hosts

The reconstitution of entire metabolic pathways into heterologous plant hosts requires the use of efficient and facile ‘gene stacking’ methodologies. A spectacularly successful example is the engineering of benzylglucosinolate biosynthesis into *Nicotiana benthamiana*. Benzylglucosinolate was reconstituted in *N. benthamiana* using a transient expression system. In this study, Geu-Flores *et al.* identified a γ -glutamyl peptidase bottleneck, suggesting that reduced sulfur is incorporated into glucosinolates via glutathione conjugation (Figure 3a) [37]. The co-expression of this peptidase augmented the yield of benzylglucosinolate 5.7-fold, indicating how consideration of primary metabolite resources can impact natural product yield [37]. In a separate metabolite analysis, Møldrup *et al.* monitored the accumulation of desulfo-benzylglucosinolate, the penultimate product in the benzylglucosinolate pathway [41]. Directing sulfur from primary to secondary metabolism through the co-expression of adenosine 5-phosphosulfate kinase—which provides the 3'-phosphoadenosine-5'-phosphosulfate (PAPS) co-substrate necessary for the final step of benzylglucosinolate biosynthesis (Figure 3a)—in the *N. benthamiana* expression system alleviated the subsequent bottleneck and increased the benzylglucosinolate yield by 16-fold [41]. In yeast, Mikkelsen *et al.* were able to reconstitute the biosynthesis of indolylglucosinolate [42]. This example was a proof-of-concept study for a technology that enables the stacking of large numbers of genes, a requirement for total pathway reconstitution.

Notably, the benzylglucosinolate biosynthetic pathway has also been stably transformed into *Nicotiana tabacum*, another non-cruciferous plant, which does not normally produce glucosinolates [43^{*}]. This reengineered plant has been shown to attract the diamondback moth (*Plutella xylostella*) and encourage oviposition (the deposition of eggs), highlighting its potential utility as a dead-end trap crop to deter predatory insects and prevent billion-dollar damages to cruciferous crops worldwide [43^{*}].

Future directions and conclusions – alkaloids and beyond

Historically, altering metabolic pathways in plants to achieve a given end has been difficult. Metabolic engineering in plants is still in its infancy and has largely been confined to single-gene expression or silencing events in the background of endogenous plant cell metabolism. The complexity of the plant host's metabolism has been shown, in many cases, to effectively mute the engineering effort or lead to unpredictable results (Table 1). However, in recent years, a wealth of new approaches has expanded the capabilities of multi-gene pathway expression in both plants and microbes and has highlighted our increasing ability to engineer the production of plant natural products in both native and heterologous systems. The increase in available and reliable sequencing and expression data enables the (relatively) facile discovery of gene, transporter and regulatory elements, the identification of which is often a prerequisite for multi-step metabolic engineering efforts. The three case studies in this review (MIAs, BIAs and glucosinolates) exemplify the challenges and progress in metabolically engineering plant-derived natural products. While we have made a special effort to highlight the advantages and pitfalls of individual techniques and efforts throughout this Opinion, a number of grand challenges for plant metabolic engineering remain to be tackled in the coming years.

Effective mining strategies

Effective mining strategies, such as those employed by Giddings *et al.* [12], Hagel *et al.* [27^{*}], Winzer *et al.* [29^{**}] and Liscombe *et al.* [44], are required to sift through the mounting data of the sequencing age. Hanson *et al.* (this issue) provides a more comprehensive review of effective mining strategies and phylogenetic analyses. Traditionally, plant enzyme discovery methods have relied heavily upon time-intensive reverse genetics. Bioinformatic techniques that engage co-expression analyses and comparative metabolite profiling to limit the gene space to be investigated are improving the discovery process. Moreover, a suite of new silencing tools, including VIGS [7], RNAi [23] and the IL-60 system [45], can provide rapid insight into the physiological function of plant enzymes.

Metabolic engineering in native versus non-native hosts

Many efforts aimed at improving the yield of alkaloids in native hosts have focused on feeding precursors and overexpressing transcription factors or enzymes positioned at metabolic bottlenecks. While these efforts often result in modest improvements to yield (Table 1), many are often accompanied by adverse morphological effects that may significantly stunt the growth of plant and tissue cultures, highlighting the tight regulation of metabolic processes within highly organized plant cells and tissues. In native systems especially, the slow growth morphologies may result from the depletion of cellular resources used in synthesizing a surfeit of transcripts and

Table 1

Examples of recent alkaloid engineering studies and their outcomes

Class	Engineering strategy	System	Observations	Reference
MIAs	Overexpression of TDC	<i>C. roseus</i> crown gall	•Did not significantly alter alkaloid yield	[53]
MIAs	Overexpression of transcription factor CrWRKY1	<i>C. roseus</i> hairy root cultures	•3-fold increase in serpentine •10-fold increase in ajmalicine	[54]
MIAs	Overexpression of feedback insensitive anthranilate synthase alpha subunit	<i>C. roseus</i> hairy root cultures	•2-fold decrease in catharanthine •300-fold increase in tryptophan •10-fold increase in tryptamine •1.8-fold in lochnericine	[55]
MIAs	Overexpression of alpha or alpha and beta subunits of anthranilate synthase and feeding of 10-deoxy-D-xylulose, loganin and secologanin	<i>C. roseus</i> hairy root cultures	•2.3-fold increase in horhammericine •1.5-fold increase in cathenamine •1.3-fold increase in catharanthine •1.8-fold increase in ajmalicine •2.1-fold increase in lochnericine •4.5-fold increase in tabersonine	[56]
MIAs	Methyl jasmonate elicitation Fed loganin and/or tryptamine	<i>C. roseus</i> hairy root tissue cultures	•3.2-fold increase in strictosidine •8.8-fold increase in ajmalicine •8.4-fold increase in tabersonine •Substrate feeding did not increase yield in elicited cultures	[57]
MIAs	Feeding with unnatural tryptamine precursors	<i>C. roseus</i> tissue cultures	•Observed $\mu\text{g/g}$ fresh weight quantities of many modified MIAs	[58]
MIAs	Mutagenesis	Fed tryptamine isomers to TDCi-silenced periwinkle hairy roots	•Observed $\mu\text{g/g}$ fresh weight quantities of many modified MIAs •Achieved 'alkaloid free' background	[21]
MIAs	Substrate feeding to roots harboring STS with expanded substrate specificity	<i>C. roseus</i> hairy root cultures	•Observed $\mu\text{g/g}$ fresh weight quantities of previously metabolically inaccessible modified products	[23]
MIAs	Overexpression of RebH/ RebF and PyrH/RebF in planta	<i>C. roseus</i> hairy root cultures	•Observed the <i>de novo</i> production of 12-chloro-19,20-dihydroakuammicine (25 $\mu\text{g/g}$ fresh weight) •Observed accumulation of 7-chlorotryptophan (50 $\mu\text{g/g}$ fresh weight)	[24]
MIAs	Overexpression of RebF and reengineered RebH in planta	<i>C. roseus</i> hairy root cultures	•Alleviated TDC bottleneck (no accumulation of 7-chlorotryptophan) •Observed the <i>de novo</i> production of 12-chloro-19,20-dihydroakuammicine (2.7 $\mu\text{g/g}$ fresh weight) •Overexpression of TDC led to slow-growing hairy roots	[25]
BIAs	heterologous expression system for production of S-reticuline and downstream alkaloids	Artificial pathway in yeast	•Produced S-reticuline in yields up to 150 mg/L •Demonstrated production of (S)-tetrahydroberberine ($\ll 5$ mg/L) •Demonstrated production of (S)-scoulerine •Demonstrated production of (S) tetrahydrocolumbamine (60 mg/L)	[35]
BIAs	heterologous expression system for production of S-reticuline	Artificial pathway in <i>E. coli</i>	•Produced (S)-reticuline in yields up to 40 mg/L	[36]
BIAs	Suppression of codeinone reductase expression	RNAi in Opium poppy	•Accumulation of (S)-reticuline (7 metabolic steps upstream)	[30]
BIAs	Overexpression of BBE in <i>E. californica</i>	<i>E. californica</i> root cultures	•5.8-fold increase of total downstream alkaloids •3.2-fold decrease in leucine concentration •2.4-fold decrease in threonine concentration •2.1-fold decrease in valine concentration •Tyrosine levels unaltered	[31]

Table 1 (Continued)

Class	Engineering strategy	System	Observations	Reference
BIAs	Suppression of BBE expression	<i>E. californica</i> root cultures	<ul style="list-style-type: none"> •Elevated levels of (S)-reticuline to 310 $\mu\text{g/g}$ cell fresh weight •(S)-reticuline levels at 6 mg/20 mL in the media 	[33]
BIAs	Systematic knockdown of morphinan pathway enzymes through VIGS	Opium poppy seedlings	<ul style="list-style-type: none"> •Suppression of SalSyn, SalR, T6ODM, CODM resulted in concomitant increase of direct precursors •Suppression of SalAT resulted in the accumulation of salutaridine (not direct precursor) •Suppression of COR resulted in increases of (S)-reticuline (not direct precursor) 	[59]
Glucosinolates	Benzylglucosinolate pathway reconstitution	Transient expression in <i>N. benthamiana</i>	<ul style="list-style-type: none"> •Produced benzylglucosinolate in yields up to 0.57 nmol/mg freshweight leaf tissue •Coexpression of peptidase raised yield 5.7-fold 	[37]
Glucosinolates	Benzylglucosinolate pathway reconstitution	Coexpression of gamma-glutamyl peptidase Transient expression in <i>N. benthamiana</i>	<ul style="list-style-type: none"> •Production of benzylglucosinolate in yields up to 1.8 nmol/mg freshweight •Expression of kinase elevated yield by 16-fold 	[41]
Glucosinolates	Indolylglucosinolate pathway reconstitution	Coexpression of adenosine 5-phosphosulfate kinase Yeast	<ul style="list-style-type: none"> •Reached indolylglucosinolate titers as high as 1.07 mg/ml •Product excreted into the media 	[42]
Glucosinolates	Benzylglucosinolate pathway reconstitution	Stable expression in <i>N. tabacum</i>	<ul style="list-style-type: none"> •Benzylglucosinolate titers as high as 0.5 nmol/mg fresh weight •Increase oviposition on non-cruciferous plants 	[43]

enzymes or from the accumulation of intermediates that negatively impact growth and development. Engineering in native hosts or heterologous plant species is attractive because not having to build the starting substrates and supply the co-factors greatly simplifies engineering efforts. However, with this strategy, maintaining the balance between primary metabolism and the engineered metabolism—a feat that will probably improve growth morphologies—is complicated precisely because endogenous primary metabolite pools are expropriated for the overproduction of selected secondary metabolites. Also, despite the advantage of minimal gene stacking, the often uncharacterized and unanticipated complex metabolism and regulatory elements of native systems can lead to engineering outcomes that are particularly difficult to control and predict. The industrial scale production of plant natural products will probably require more comprehensive engineering efforts than single-gene overexpression or silencing events in the context of native plant hosts. Engineering in faster-growing and ‘chemically silent’ heterologous hosts may increase biomass accumulation and simplify purification.

Controlling metabolic flux through new expression constructs, scaffolds and tunable regulatory elements

The intricate relationship between primary metabolism (i.e. glycolysis, the TCA cycle) and the native or heterologous secondary metabolism (i.e. isoprenoid and alkaloid

pathways) must be considered. In plants, Park *et al.* alluded to this interplay by demonstrating that BBE expression levels vastly affect amino acid levels [31,32]. As metabolic engineering strategies in plants become more sophisticated, we should also begin to consider flux analyses, taking into account that natural product pathways are evolutionarily optimized to channel intermediates toward product through a highly choreographed system of protein-protein interactions, localization and regulation [46]. The overall goal is to maximally channel metabolic resources to the desired products without over taxing the host system.

Co-localization through scaffolding is a proven way to channel metabolites in *E. coli*. These systems attempt to mock natural megasynthases, which efficiently shuttle metabolites between active sites. Essentially, scaffolding increases the local metabolite and enzyme concentrations and effectively lowers the K_m of the substrate. These systems are widely modular and are known to improve titers, alleviate metabolic bottlenecks and reduce metabolic loads by preventing carbon from exiting the pathway. Under conditions of low enzyme expression (decreased metabolic load), Dueber *et al.* successfully achieved a 77-fold enhancement in mevalonate production by building a scaffold based on the protein-protein interactions of GBD, SH3 and PDZ domains and their cognate ligands [47]. They built the scaffold on hydroxymethylglutaryl-CoA reductase, the enzymatic

bottleneck of mevalonate production [47]. Notably, these scaffolding systems require that the enzyme at the metabolic bottleneck, the subsequent enzyme and the substrate be co-localized [47], underscoring why they may be untenable for some highly compartmentalized systems. Nonetheless, the prospect of engineering metabolons into plants is exciting.

The effective metabolic engineering of plant natural products will inevitably require advanced, but easy-to-use, gene stacking techniques. Traditionally, multi-gene expression in plants has been plagued by inadvertent silencing events, the incomplete incorporation of all genes and lengthy and technically challenging procedures [48]. A number of new technologies, however, are being developed to assemble and transplant large fragments of DNA easily [48]. Golden Gate cloning and USER fusion have been used to clone multiple pathway elements [48,49]. Additionally, synthetic plant chromosomes and the universal expression and silencing IL-60 platform both have the demonstrated capability of introducing multiple plant pathway elements into plants [48,50]. For example, under the transformation-free IL-60 platform, Mozes-Koch *et al.* expressed an entire bacterial operon in tomato and produced pyrrolinitrin, which they observed after only two days [50*].

A number of RNA-based silencing systems, including RNAi, have also been engineered and applied to medicinal plants [21]. Notably, RNAi, which provides a permanent phenotype or chemotype, has been employed to block shunt pathways and channel metabolic resources toward a desired product, enhancing our ability to engineer in multiple dimensions [21].

Lastly, promoter libraries, engineered untranslated regions (e.g. 5' untranslated regions and intergenic regions), genetic circuit designs and biosensor regulators have been tremendously helpful in microbial engineering. Applying these design principles to the metabolic engineering of plants may greatly enrich our efforts to produce valuable and chemically diverse alkaloids. Notably, a variety of constitutive and inducible plant promoters and expression systems are now widely available. Synthetic RNA elements, ribosome binding site elements and a combination of different strength promoters strategically placed in front of stacked pathway genes could theoretically enable tunable protein expression [35,51]. These elements could potentially limit the expression of toxic activities or the accumulation of toxic metabolites until the stationary phase (or an appropriate stage) of growth, thereby absolving the system of unsustainable metabolic burden.

Localization and transport – engineering in multiple dimensions

Many alkaloid biosynthetic pathways are highly compartmentalized at both the intercellular and intracellular

levels. For example, at least three cell types are required for the biosynthesis of many MIAs [6]. The forward engineering of plant natural products will require sifting through the increasing amount of available sequencing and expression data and untangling the complexity of the plant cell and different tissue types. In addition to the linear design and channeling of metabolic pathways, the successful metabolic engineering of plant natural products will require engineering in the ‘third dimension,’ namely at the level of localization and cell type [52].

Combinatorial biosynthesis in plants – mixing and matching pathways and engineering new enzyme specificities

De novo combinatorial biosynthesis in plant systems has gone largely underexplored. Most of the few efforts to engineer unnatural natural products have utilized precursor feeding or mutasynthesis-based approaches, which can be costly and time intensive. The *de novo* biosynthesis of unnatural natural products will require that constituent enzymes have reengineered or broad specificity. Notably, directed evolution has been successfully used to alter enzyme specificity. However, enzyme engineering efforts are greatly enhanced if the protein structure is known and the mechanism is well understood. Then, the enzyme can be subjected to structure-guided techniques, such as site-directed mutagenesis and domain swapping, which create smaller protein libraries enriched with functional mutants.

As we seek further to convert plants and microbes into the chemical factories to meet our medicinal needs, we should remember that, although many plant natural products are bioactive and serve as important lead compounds, they often require modification before making it to the clinic. Therefore, the forward engineering of ‘unnatural’ or ‘new-to-nature’ natural products must also be a grand challenge if plant natural products are to be shuttled from the annals of human tradition into the drug development programs and clinics of tomorrow.

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