Biosynthesis and regulation of terpenoid indole alkaloids in *Catharanthus roseus*

Jianhua Zhu¹, Mingxuan Wang¹, Wei Wen¹, Rongmin Yu¹,²

¹Biotechnological Institute of Chinese Materia Medica, ²Department of Natural Medicinal Chemistry, College of Pharmacy, Jinan University, Guangzhou 510632, China

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**ABSTRACT**

*Catharanthus roseus* produces a wide range of terpenoid indole alkaloids (TIA). Many of them, such as vinblastine and vincristine, have significant bioactivity. They are valuable chemotherapy drugs used in combination with other drugs to treat lymphoma and leukemia. The TIA biosynthetic pathway has been investigated for many years, for scientific interest and for their potential in manufacturing applications, to fulfill the market demand. In this review, the progress and perspective of *C. roseus* TIA biosynthesis and its regulating enzymes are described. In addition, the culture condition, hormones, signaling molecules, precursor feeding on the accumulation of TIA, and gene expression are also evaluated and discussed.

**Key words:** Biosynthesis pathway, enzymes, gene expression, regulation, terpenoid indole alkaloids, vinblastine

**BIOSYNTHESIS OF BISINDOLE ALKALOIDS**

The medicinal plant *Catharanthus roseus* (L.) G. Don is of enormous pharmaceutical interest because it contains about 130 terpenoid indole alkaloids (TIAs), some of which exhibit strong pharmacological activities. Jamalicine is an antihypertensive alkaloid. Vinblastine and vincristine, which are bisindole alkaloids derived from coupling vindoline and catharanthine, were the first natural drugs used in cancer therapy and are still among the most valuable agents used in the treatment of cancer, to date. All TIAs in *C. roseus* are derived from the central precursor strictosidine, which is a fusion product of the shikimate pathway–derived tryptamine moiety and the plastidic nonmevalonate pathway–derived secologanin moiety. The anticancer agents vinblastine and vincristine are produced exclusively by *C. roseus*. More importantly, the TIA biosynthetic pathway is under strict developmental and environmental control.

**FORMATION OF STRICTOSIDINE**

Strictosidine, the central intermediate in the TIA biosynthesis of *C. roseus*, is formed by the coupling of indole glycoside secologanin and tryptamine under the catalysis of strictosidine synthase (STR).

**FORMATON OF VINDOLINE**

Strictosidine-β-D-glucosidase (SGD) may be the enzyme playing an important role in steering the monoterprenoid indole alkaloid biosynthesis in a specific direction. The removal of the glucose moiety of strictosidine by SGD leads to an unstable, highly reactive aglucon, which is thought to be converted to 4,21-dehydrogeissoschizine. The latter is believed to be converted by cathenamine synthase to cathenamine. Subsequently, the cathenamine is converted into tabersonine through several steps, which are not clearly understood. Finally, tabersonine is
transformed into vindoline by a sequence of six steps.\[4,5\] The steps include: Aromatic hydroxylation, O-methylation, hydration of the 2,3-double bond, N(1)-methylation, hydroxylation at position 4, and 4-O-acetylation. The intermediates involved are 16-hydroxytabersonine, 16-methoxytabersonine, 16-methoxy-2,3-dihydro-3-hydroxy-tabersonine, desacetoxyvindoline, and deacetylvindoline. Most enzymes in the biosynthetic pathway from tabersonine to vindoline have been identified. They are tabersonine 16-hydroxylase (T16H), O-methyltransferase (OMT), N-methyltransferase (NMT), desacetoxyvindoline 4-hydroxylase (D4H), and deacetylvindoline-4-O-acetyltransferase (DAT). However, the enzyme that catalyzes the conversion of 16-methoxytabersonine to 16-methoxy-2,3-dihydro-3-hydroxy-tabersonine is still unknown.\[4\]

**FORMATION OF CATHARANTHINE**

The information on catharanthine biosynthesis is very limited. It may be derived from strictosidine through the intermediate of geissoschizine and stemmadenine. However, the enzymes involved are not isolated and the genes are not cloned.\[5\]

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**Figure 1:** Biosynthesis of Catharanthus TIAs. Solid arrows indicate confirmed enzymatic conversions, whereas, the broken arrows indicate unknown enzymatic conversions. G10H: geraniol 10-hydroxylase; TDC: Tryptophan decarboxylase; STR: strictosidine synthase; SGD: Strictosidine β-D-glucosidase; T16H: 16-hydroxylase; OMT: O-methyltransferase; NMT: N-methyltransferase; D4H: Desacetoxyvindoline 4-hydroxylase; DAT: Deacetylvindoline-4-O-acetyltransferase; AVLBS: Anhydrovinblastine synthase.
FORMATION OF BISINDOLE ALKALOIDS
The bisindole alkaloids vinblastine and vincristine are of great interest. They are synthesized from the coupling of the monomeric alkaloids catharanthine and vindoline. The product resulting from the coupling is α-3′,4′-anhydrovinblastine, which is converted into vinblastine and then further converted into vincristine. The coupling process is catalyzed by the enzyme anhydrovinblastine synthase (AVLBS).[^25] However, the enzyme catalysis the formation of vinblastine from α-3′,4′-anhydrovinblastine is still unknown. Moreover, the enzyme that catalyzes the conversion of vinblastine to vincristine is also not isolated.

REGULATION FACTORS OF DIMERIC INDOLE ALKALOIDS DIAS AND THEIR PRECURSOR BIOSYNTHESIS

Light
Light is thought to have an effect on enzyme induction and activation. It has been shown that light significantly influences the biosynthesis of vindoline and other alkaloids, as well as acidic and basic peroxidase activities. Light promotes vindoline and serpentine biosynthesis, and stimulates plastid development and peroxidase activity.[^11] After it is light-treated, the concentration of vindoline increases significantly in cultures of *C. roseus*, including in cultured cells, leaves, seedlings, and plants. The results of gene expression investigation have demonstrated that upregulation of tryptophan decarboxylase (TDC), D4H, and DAT has been observed in *C. roseus* cultures after light expression.[^12,13] Researches have also shown that the content of catharanthine, vindoline, and vinblastine is markedly increased by ultraviolet (UV)-B radiation in *C. roseus*.[^13,16]

Plant growth regulators
Plant growth regulators affect both culture growth and secondary metabolite production.

Auxin and cytokinin
Plant growth hormones inhibit the accumulation of alkaloid, while cytokinins act as accelerants. 2,4-D strongly inhibits alkaloid production, essentially during the growth phase.[^18] More importantly, genes, such as, the 1-deoxy-D-xylulose-5-phosphate synthase (DXS) gene, 1-deoxy-D-xylulose-5-phosphate reductoisomerase (DXR) gene, and TDC gene are repressed in the suspension-cultured cells of *C. roseus* with 2,4-D.[^20,21] Transferring these cell suspensions in a 2,4-D-free culture medium gradually increases the expression of these genes.[^21,22] Moreover, 2,4-D depletion also enhances the 2C-methyl-D-erythritol 2,4-cyclodiphosphate synthase (CRMECS) mRNA steady-state level in *C. roseus*. This gene remains expressed at a low rate in the cell suspensions cultured with 2,4-D.[^23] These investigations support the regulatory role of auxin on the methylerythritol phosphate (MEP) pathway, leading to the biosynthesis of the TIA terpene moiety, and auxin acts, at least in part, at the MEP pathway gene expression regulation level.[^29]

Except for 2,4-D, other auxins such as, 1-Naphthaleneacetic acid (NAA) and indole-3-acetic acid (IAA), also downregulate the TDC gene in the biosynthetic pathway of alkaloids. Omission of NAA from the growth medium results in the accumulation of TDC.[^21] Thus, auxin negatively regulates the expression of the genes associated with the terpenoid indole alkaloid biosynthesis in *C. roseus*.

On the contrary, cytokinin remarkably enhances the accumulation of alkaloids in Catharanthus cultures. Genes experiments show that cytokinin greatly enhances the expression of the geraniol 10-hydroxylase (*G10H*) gene.[^25]

Other plant growth regulators
Enhancement of catharanthine in the hairy roots and accumulation of vindoline in shoot cultures are observed after ethylene treatment.[^13] In addition, ethylene applications promote the pathways toward ajmalicine, serpentine, and tabersonine.[^24] Gibberellic acid (GA), similar to 2,4-D, has a strongly negative influence on the accumulation of vinblastine, vindoline, and catharanthine.[^25] Our previous study has found that artemisinic acid, a cadinane-type of sesquiterpene, can stimulate the biosynthesis of catharanthine, vindoline, and vinblastine in *C. roseus* cultured cells. In addition, upregulation of *G10H*, *SGD*, *TDC*, *T16H*, and *D4H* was also observed.[^26,27]

Signaling molecules

Jasmonates
Jasmonates are plant signaling molecules that play key roles in protection against certain pathogens and insects, by switching on the expression of gene-encoding defense proteins, including enzymes involved in the biosynthesis of toxic secondary metabolites.[^28]

In the hairy roots, jasmonic acid (JA) is found to be a unique elicitor leading to an enhancement in the flux to several branches in the indole alkaloid pathway.[^29] The accumulation of ajmalicine, serpentine, lochnericine, and hörhammercine are significantly increased after the addition of jasmonic acid.[^29]

Methyl jasmonate (MJ) induces several alkaloids in *C. roseus* cultures. After treatment with MJ, ajmaline is increased in the cultured plant cells, while both ajmalicine and catharanthine are increased in the hairy roots. For shoot cultures, vindoline accumulation, induced by MJ, is observed, which is about 6.5-fold compared to the control.[^28]

In *C. roseus* seedlings, the TIA genes exhibit a significant variation in the magnitude and timing of induction by MJ. ORCA3, a jasmonate-responsive APETALA2 (AP2)-domain transcription factor gene, exhibits the greatest increase in the transcript levels after MJ treatment.[^30,31] MJ-induced increases in the transcript levels of the TIA genes occur in the following order: ORCA3, D4H, STR, TDC, G10H, and cytochrome P-450 reductase (CPR).[^30]
Salicylic acid

Salicylic acid (SA) has been known to ameliorate the adverse effects of salinity by improving plant productivity through protecting the photosynthetic pigments and producing antioxidative compounds and enzymes. It enhances vincristine and vinblastine alkaloid production in Catharanthus roseus by improving the antioxidant defense system. In seeding, salicylic acid treatment increased the production of tabersonine and a higher concentration of salicylic acid induced vindoline accumulation. Meanwhile, the activity of alkaline peroxidase increased 5-fold.

Nitric oxide

Nitric oxide (NO) is known as a signaling molecule involved in the elicitor-induced defense responses of plants. Sodium nitroprusside (SNP), a donor of NO, stimulates catharanthine formation in Catharanthus roseus cells. However, the yield of catharanthine decreases when treated with the JA inhibitor (such as ibuprofen). Therefore, NO stimulates the accumulation of catharanthine, which is JA-dependent.

Precursor feeding

Feeding of precursors is one of the most effective strategies to increase the production of important secondary metabolites in cells and organ cultures. The results of the addition of some precursors show positive results on the accumulation of alkaloids. For instance, the accumulation of ajmalicine and strictosidine is significantly enhanced when treated with tryptamine or loganic acid.

CONCLUSION

The biosynthesis of the Catharanthus roseus terpenoid indole alkaloid has been studied for decades. Although there is tremendous progress, there are many steps that are still not profiled. To date, Catharanthus roseus is the only nature resource for antitumor medicinal compounds. However, their content is low. The unclear biosynthesis pathway of the TIAs is the main reason that blocks the progress. Therefore, elucidation of the TIA biosynthesis pathway must be emphasized on greatly and some new strategies, such as plant and microbial production, may also be employed for the manufacture of these valuable medicinal compounds.

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