

RESEARCH ARTICLE



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Total synthesis of (–)-deglycocadambine†

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The first total synthesis of the monoterpene indole alkaloid (–)-deglycocadambine is achieved in 12 steps with (+)-genipin as the chiral starting material. The reported synthetic approach is characterized by an orchestrated cascade annulation between tryptamine and the highly functionalized dialdehyde precursor, rapidly introducing the unique 6/5/6/7/6-fused pentacyclic skeleton and the ketone functional group at C19 in a convergent manner. The successful implementation of transannular oxidative cyclization at C3 for bridged oxazolidine formation in the late-stage synthetic campaign ensured the final total synthesis of this molecule.

Introduction

(–)-Deglycocadambine (Fig. 1) is a member of the monoterpene indole alkaloids, a large family of metabolites that have attracted much attention from both synthetic and pharmaceutical chemists due to their high structural diversity and significant biological activities.¹ First isolated from the twigs and leaves of *Emmenopterys henryi* by Song and Zhao in 2013,² (–)-deglycocadambine has a characteristic 6/5/6/5/6/6 hexacyclic ring system, with its 6/5/6/7 tetracyclic nucleus containing a unique hydroptyrido[1,2-*a*]azepine subunit, which can also be found in (–)-rubenine, (–)-kopsiyunnanine K, (–)-voacacines A and B and other cadambine-type alkaloids, as shown in Fig. 1.³ Although preliminary studies revealed no promising activity, scarcity in nature precludes its systematic biological evaluation.² To the best of our knowledge, no synthetic route has been reported thus far for this molecule. Of note, in 1991, Brown *et al.* reported an elegant biologically inspired synthesis of (–)-cadambine from (–)-secologanin,^{4,5} wherein a well-

designed chemo- and regioselective reaction between tryptamine and a highly functionalized epoxide was conducted to efficiently introduce the seven-membered azepine ring of (–)-cadambine. Theoretically, hydrolysis of (–)-cadambine should give the desired aglycone (–)-deglycocadambine. Accordingly, the exploration of a general synthetic route for the total synthesis of (–)-deglycocadambine is required to pursue the novel molecular architecture and investigate the potential molecular functionality. Based on a rationally designed two-fold annulation sequence for the rapid installation of the 6/5/6/7/6-fused pentacyclic core, we report our synthetic efforts toward (–)-deglycocadambine.

Retrosynthetically, as shown in Scheme 1, (–)-deglycocadambine could be properly obtained from the pentacyclic precursor A by the formation of the bridged oxazolidine ring *via* the orchestrated transannular oxidative cyclization.⁶ The stereospecific hydroxyl group at C19 of A could theoretically be

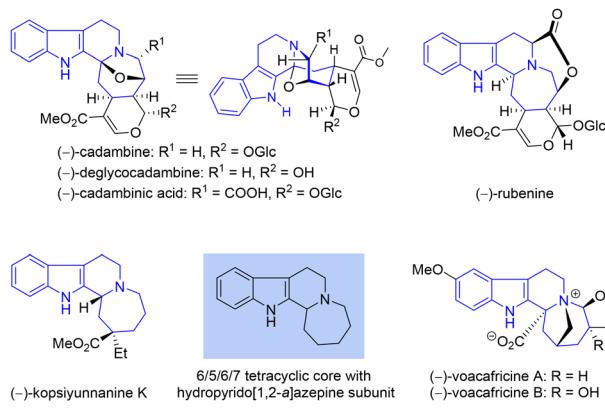


Fig. 1 Representative indole alkaloids with hydroptyrido[1,2-*a*]azepine subunits embedded in the 6/5/6/7-fused tetracyclic core.

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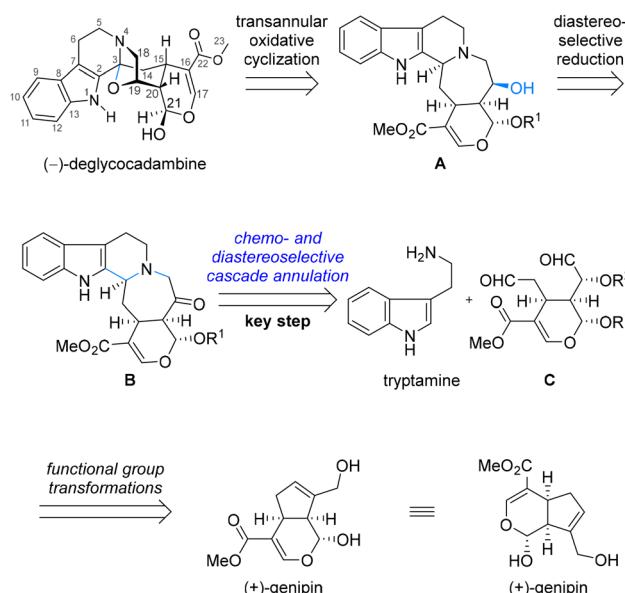
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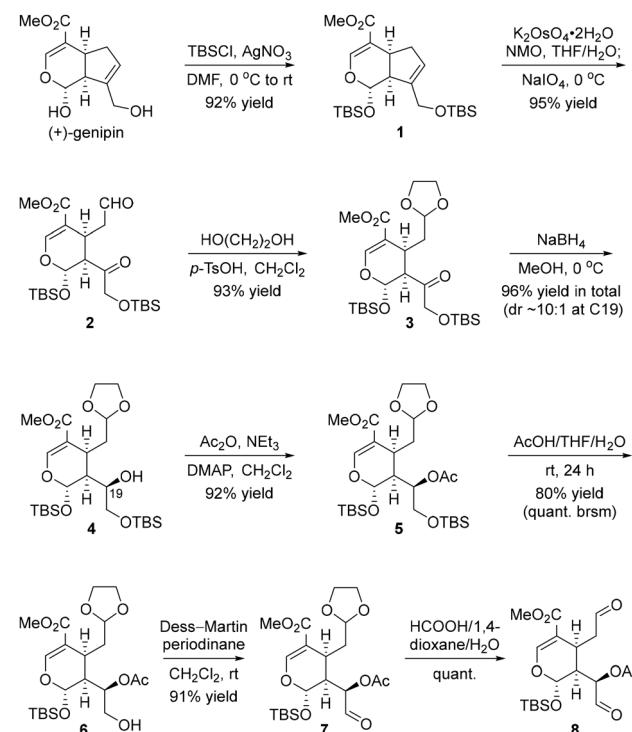


Scheme 1 Retrosynthetic analysis of (-)-deglycocadambine.

introduced *via* the diastereoselective reduction of the corresponding ketone in synthon **B**. The critical hydroptyrido[1,2-*a*]azepine subunit embedded in **B** could be tentatively assembled by a rationally designed twofold annulation reaction between tryptamine and the highly functionalized dialdehyde **C**,⁷ the latter of which could be formally derived from the commercially available (+)-genipin by means of continuous functional group transformations.

Results and discussion

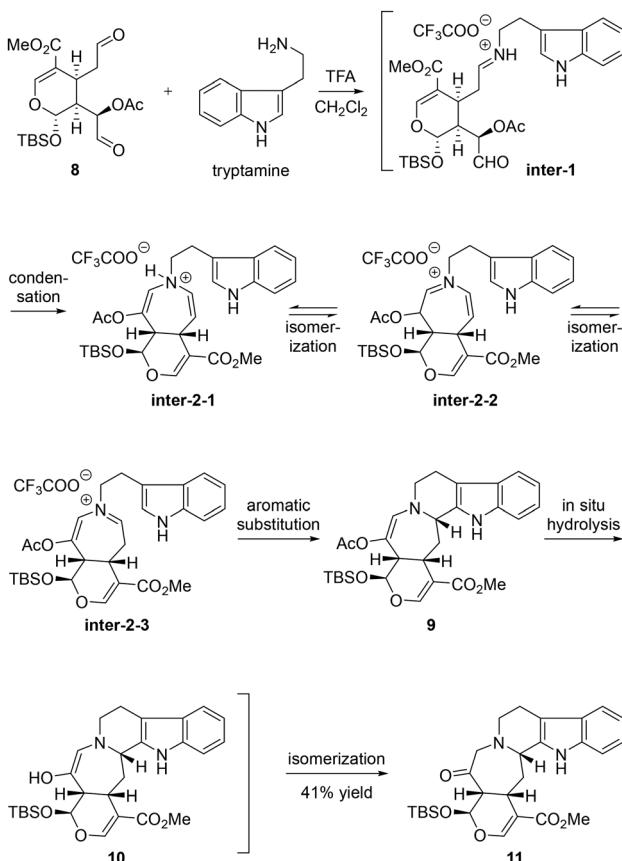
Initially, to access fragment **8** (Scheme 2), we selected commercially available iridoid (+)-genipin⁸ (~\$5 per g) from the chiral pool to start our synthetic campaign. Protection of both hydroxyl groups as TBS ethers was conducted in the presence of TBSCl and AgNO₃, giving **1** in 92% yield. The following chemoselective dihydroxylation/oxidative cleavage cascade reaction of the isolated alkene in the five-membered ring was performed, affording compound **2** in 95% yield, with the left-hand alkene embedded in the conjugated system remaining intact. After the two-step operation involving the chemoselective protection of the aldehyde in **2** as the acetal and the diastereoselective reduction of the ketone in **3**, alcohol **4** was produced as the major epimer (dr ~ 10 : 1) in an overall yield of 81%. Notably, for the reductive transformation from **3** to **4**, the observed stereochemistry at C19 could be explained by the Felkin–Anh model,⁹ wherein the *Si* face attack was preferred. With **4** in hand, esterification of the secondary alcohol in the presence of Ac₂O gave, after chemoselective deprotection of the right-hand TBS ether, the primary alcohol **6**. Finally, oxidation of the hydroxyl group in **6** using Dess–Martin periodinane followed by chemoselective hydrolysis of the ethylene acetal in **7** delivered the dialdehyde **8** without incident.

Scheme 2 Synthesis of dialdehyde **8**.

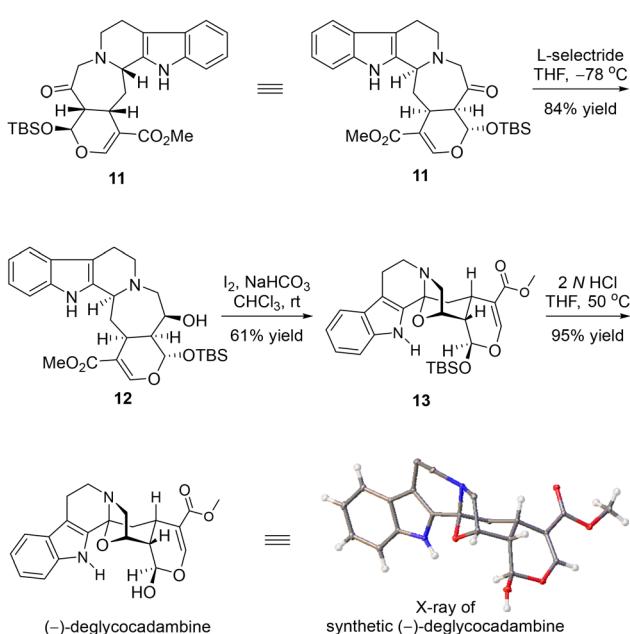
For the key cascade annulation of dialdehyde **8** and tryptamine, the pentacyclic ketone **11** was obtained as the sole identified product in 41% yield when the reaction mixture was exposed to TFA in CH₂Cl₂.^{10,11} In this process, as shown in Scheme 3, it was postulated that tryptamine firstly condensed chemoselectively with the sterically more accessible aldehyde **8** to form **inter-1**, followed by the second condensation, delivering **inter-2-1**. **Inter-2-1** might isomerize to **inter-2-2** and further to **inter-2-3**. Among these isomers existing in dynamic equilibration, **inter-2-3** might be the most structurally favored for further aromatic substitution. The resulting pentacyclic compound **9** was further *in situ* hydrolyzed to give enol **10**, which was spontaneously isomerized to the more stable ketone **11**. The stereochemical outcome at C3 in **11** might be attributed to the much easier accessibility of the *Re* face of the iminium ion in **inter-2-3** as molecular modelling of **inter-2-3** revealed that the *Si* face attack presumably led to severe steric repulsion between the indole ring and H-15. Strategically, by using this elaborated twofold annulation protocol, we not only incorporated the synthetically challenging hydroptyrido[1,2-*a*]azepine substructure but also finished the installation of ketone at C19 for further selective transformation at the same time. It should be noted that this is the first example for this cascade annulation wherein a complex and asymmetric dialdehyde was used to react with tryptamine.^{7,12}

With **11** in hand, as shown in Scheme 4, a chemo- and diastereoselective reduction of a ketone using *L*-selectride as the reducing reagent gave alcohol **12** in 84% yield. The I₂-mediated transannular oxidative cyclization of the tertiary





Scheme 3 Synthesis of pentacyclic ketone 11.



Scheme 4 Late-stage synthesis of (-)-deglycocadambine.

benzylic amine occurred smoothly to furnish compound 13, with the last bridged oxazolidine ring characterized by the *N*,*O*-substituted quaternary stereocenter at C3 assembled

efficiently.⁶ When subjected to HCl in THF at elevated temperature, the TBS ether of 13 was directly deprotected to provide (−)-deglycocadambine, the absolute stereochemistry of which was unambiguously confirmed by X-ray crystallographic analysis.¹³ The ¹H and ¹³C NMR spectroscopic data of the synthetic sample are in accordance with the reported ones.^{2,11}

Conclusions

In conclusion, a convergent synthetic strategy has been developed based on the ingenious use of a twofold annulation reaction between the dialdehyde and tryptamine to rapidly install the highly functionalized 6/5/6/7/6-fused pentacyclic ring system and a late-stage transannular oxidative cyclization to directly incorporate the bridged oxazolidine ring for the total synthesis of (−)-deglycocadambine, which was obtained in an overall yield of 9.5% over a longest linear sequence of 12 steps from (+)-genipin. Strategically, the bond-forming logic demonstrated in our protocol for installing the multicyclic ring system characterized by a unique hydroxypyrido[1,2-*a*]azepine subunit of (−)-deglycocadambine should enlighten the upcoming efforts towards chemical synthesis of other structurally related monoterpenoid alkaloids in our laboratory.

Author contributions

F.-X. W. and Y. R. C. directed the project. F.-X. W. and Y. R. C. conceived the synthetic route. F.-X. W., Y.-T. C. and H. L. conducted the synthetic work. F.-X. W., Y. R. C., Y.-T. C. and H. L. analyzed the results. F.-X. W. and Y. R. C. wrote the manuscript. All authors commented on this manuscript and gave approval to the final version of this manuscript.

Data availability

Electronic supplementary information (ESI) available: Experimental procedures, NMR spectra, IR, and HRMS data. CCDC 2358655. For ESI and crystallographic data in CIF or other electronic format see <https://doi.org/10.1039/D4QO01122H>.

Conflicts of interest

There are no conflicts to declare.

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13 CCDC 2358655† for (–)-deglycocadambine contains the supplementary crystallographic data for this paper.

