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# A concise and stereoselective synthesis of the BCDF tetracyclic ring system of C<sub>19</sub>-diterpenoid alkaloids†

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A new synthetic route for the BCDF tetracyclic ring system of  $C_{19}$ -diterpenoid alkaloids ( $C_{19}$ -DTAs) has been developed. The key step is a Pd-catalyzed transannular alkenylation that installs a functionalized bridged F ring. The overall strategy is concise and stereoselective, and it provides a valuable new tool for the synthesis of  $C_{19}$ -DTAs. The synthesis begins with a bridged [3.2.1] ring system, which is converted to a key intermediate through a series of highly regio- and stereoselective processes. The introduction of an allylic side chain with high precision is accomplished, culminating in a Pd-catalyzed transannular alkenylation that installs a functionalized bridged F ring to yield the BCDF tetracyclic analog of  $C_{19}$ -DTAs.

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### Introduction

Aconitine (1), first isolated by Manske in 1833 and its structure elucidated in 1959,1,2 is a prototypical C19 diterpenoid alkaloid (C<sub>19</sub>-DTAs). It is the major toxic component of over 250 species of Aconitum plants, commonly referred as monkshood or wolf's bane, which have long been used in traditional medicine.3 Aconitine's primary mode of action involves potent activation of sodium ion channels, preventing repolarization of muscles and neurons. As a result, it can induce ventricular arrhythmias, potentially leading to death.3 While aconitine's high toxicity has limited its clinical use, less toxic analogs and derivatives are emerging as promising candidates for the treatment of pain, inflammation, or arrhythmias.4 They share a common hexacyclic ring system (ABCDEF ring) and typically possess three to nine oxygen substituents such as hydroxy, methoxy, and acyloxy groups (Fig. 1). Their intricate structures and diverse biological activities have attracted considerable attention from synthetic chemists over the past five decades. 5 As pioneers in the field, Wiesner et al. developed four generations of fundamental synthetic strategies for the practical synthesis of talatisamine, chasmanine, and 13-desoxydelphonine in the 1970s.6 In the last decade, research groups led by Gin,7 Sarpong,8 Fukuyama,9 Inoue<sup>10</sup> and Reisman<sup>11</sup> have made significant contributions, culminating in the elegant total syntheses of weisaconitine D, liljestrandinine, cardiopetaline, talatisamine and puberuline C.

Recently, Qin and co-workers reported an ingenious total synthesis of vilmoraconitine by using an efficient hydrodealkenylative fragmentation/Mannich strategy. While other synthetic efforts exist, these remain the only eight natural  $C_{19}$  ( $C_{18}$ ) DTAs with successful total syntheses to date. Notably, all synthesized  $C_{19}$ -DTAs or  $C_{18}$ -DTAs possess only three to six oxygen functionalities. The total synthesis of  $C_{19}$ -DTAs with a more oxygenated ring system, such as aconitine (1) with its unprecedented nine oxygen functionalities, remains a significant challenge.

Due to our long-standing research interest in the chemistry of C<sub>19</sub>-DTAs, we embarked on synthetic studies of aconitine and other bioactive C<sub>19</sub>-DTAs. In previous synthetic efforts, we have constructed several important ring systems, such as the AEF, ABEF, ABF, BCD and BCDE rings of C<sub>19</sub>-DTAs. In particular, we have previously utilized a cascade oxidative dearomatization/dimerization/retro-DA/IMDA reaction and the Wagner-Meerwein rearrangement as key steps to successfully construct the tricyclic [6.2.1.0] carbocyclic BCD ring analog, providing

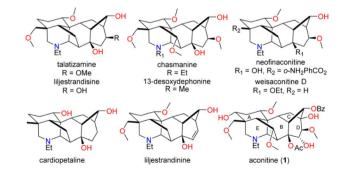


Fig. 1 Structures of  $C_{18}$ - or  $C_{19}$ -diterpenoid alkaloids.

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a potential approach for the construction of the bridged A, E, and F rings in aconitine.

However, the stereoselectivity for the preparation of the key rearrangement precursor was still undesirable and needed further improvement. Herein, we report further synthetic studies on the highly functionalized BCD ring analog 4 *via* an improved synthetic route that further introduced a functionalized bridged F ring *via* a Pd-catalyzed transannular alkenylation reaction to successfully afford the tetracyclic BCDF ring compound 2, which is an important model substrate intermediate for the total synthesis of aconitine.

We envisioned the synthesis of the tetracyclic aconitine BCDF ring analog 2 (Scheme 1). Initially, compound 3, featuring a [3.2.1] bridged ring structure, could be transformed from compound 2 through a transannular alkenylation reaction. <sup>15</sup> Compound 2 could be obtained from ketone 4 *via* regioselective acylation and stereoselective alkylation reactions. Ketone 4, in turn, could be derived from the 5/6/6 tricyclic ketone 5 *via* a Wagner–Meerwein rearrangement. <sup>16</sup> Finally, compound 5 could be synthesized in a single step through an intramolecular oxidative dearomatization/IMDA cascade reaction of phenol 6 with a dienophile side chain. Although we have previously synthesized analogous tricyclic compounds to 4, developing a rapid, highly stereoselective, and gram-scalable method for the synthesis of the BCD tricyclic compound 4 is still necessary to further expand our synthetic work.

#### Results and discussion

Our synthetic studies began with the preparation of vinyl arene 13 as shown in Scheme 2. First, commercially available 2,3-dihydroxybenzaldehyde 7 was converted on a 10 gram scale to 3-hydroxy-2-methoxybenzaldehyde 8 *via* a known regioselective phenolic hydroxyl-methylation reaction. To Compound 8 was then converted to the TBS phenol ether 9 under standard conditions. Next, the Wittig reaction of compound 9 with a phosphonium salt generated the enol ether intermediate 10, which was subsequently subjected to *in situ* acid-promoted hydrolysis, affording the one-carbon homologated aldehyde 11 with an overall yield of 71% across two steps. Finally, aldehyde 11 was subjected to a Barbier allylation reaction with allyl bromide in the presence of Zn/NH<sub>4</sub>Cl to introduce the terminal olefin side chain, yielding highly functionalized phenol 12.

Scheme 1 Synthetic plan for the BCDF ring system 2.

Scheme 2 Synthesis of vinyl arene 13.

Selective protection of the resulting homoallylic alcohol with TBS then afforded phenol 13.

After successfully obtaining vinyl phenol 13, we proceeded with the oxidative dearomatization/intramolecular Diels-Alder reaction (IMDA) and the subsequent planned Wagner-Meerwein rearrangement (Scheme 3). Fortunately, phenol 13 with bis(acetoxy)iodobenzene (PIDA) under standard oxidative conditions, uniquely produced the relatively stable *o*-quinone monoacetal 14 at 0 °C in methanol and, notably, the typically observed *o*-quinone monoacetal dimerization at room

Scheme 3 Synthesis of BCDF analogue 29.

temperature<sup>20</sup> did not occur. Rapid column chromatographic purification of compound 14, followed by reflux in toluene, easily afforded a separable mixture of endo diastereoisomers 15a and 15b, with a small amount of the undesired exo addition product (15%),<sup>21</sup> achieving the expected endo stereoselectivity with a combined yield of 72% for the two steps. The  $\alpha$ -stereochemistry of the OTBS substituent was determined in 15a by observing the NOE correlation between the vinyl proton signal ( $\delta = 6.24$ , m) and the *tert*-butyl proton signal ( $\delta = 0.88$ , s) of the OTBS substituent. Notably, 15a or 15b showed a resemblance in the position of the carbonyl and acetal substituents adjacent to the [2.2.2] bicyclooctane bridge portion with our previously synthesized tricyclic analog 15x,14 which could potentially enhance the stereoselectivity of the key Wagner-Meerwein rearrangement precursor. It was encouraging to find that the ketone carbonyl reduction of 15a proceeded exclusively from the more hindered β-face. This selectivity may have been influenced by the potential torsional steering effect of the neighboring gem-dimethoxy group.8 By using sodium borohydride (NaBH<sub>4</sub>) for the attack from the more crowded β-face, we were able to synthesize compound 16 with high stereoselectivity. Subsequent treatment with sodium hydride (NaH) and iodomethane (MeI) converted the newly formed methylene hydroxy group into the methoxy ether compound 17, achieving a 99% yield. After removal of the dimethyl ketal functional group under acidic conditions to give the ketone 18, a stereoselective reduction with NaBH<sub>4</sub> occurred from the β-side again (due to the steric hindrance of the  $\alpha$ -side OMe substituent), leading to the formation of alcohol 19 with the hydroxy group oriented to the α-position in a combined yield of 82% in two steps. Subsequently, by activating 19 with trifluoromethanesulfonic anhydride (Tf<sub>2</sub>O) in DCM and using methanol as nucleophile, the key Wagner-Meerwein rearrangement was performed in a one-pot manner without the need to isolate the potential intermediate triflate, directly yielded the allyl methyl ether isomers 20a and 20b in yields of 72% and 8%, respectively.22 We attribute the predominance of 20a as the major product to the absence of a strained bridgehead double bond. In contrast, 20b has a strained bridgehead olefin, making its formation less favored. When water is used instead of methanol to trap the allyl carbocation, the C8-hydroxy substituted rearranged product is generated, along with a small amount of the

After successfully synthesizing bicyclo [3.2.1] octane vinyl ether isomers **20a** and **20b**, we investigated their further transformation into the BCDF tetracyclic target molecule. Treatment of a mixture of **20a** and **20b** with 40% aqueous HF resulted in both TBS deprotection and removal of the allylic methyl ether, forming the diol **21**. This demonstrates that the vinyl isomer **20b** can also generate the same allylic alcohol under acidic conditions, probably due to the formation of a more stable allylic tertiary carbocation. The addition of the hydroxy group is controlled by the relative stability of the olefin isomers. Subsequent oxidation of the C-17 secondary alcohol in **21** with Dess-Martin periodinane (DMP) afforded the alcohol–ketone **22**. After TES protection of the exposed C-8 tertiary alcohol, deprotonation of ketone **23** with lithium hexamethyldisilazane

(LiHMDS) and subsequent introduction of a methoxycarbonyl group using Mander's reagent<sup>23</sup> (methyl cyanoformate) afforded enol ester 24 in high regioselectivity (98% yield) due to the steric hindrance caused by the OTES group at the adjacent C-7 position. C-Alkylation of enol ester 24 at C-11 with 2,3-dibromopropene to introduce the F ring side chain initially yielded significant O-alkylation by-products. However, the addition of KI as an additive<sup>24</sup> facilitated the regioselective formation of the C-alkylated product 25 in 90% yield. Based on HSAB25 theory, this improved regioselectivity is attributed to the in situ replacement of the allyl bromide atom with a more nucleophilic iodide ion, forming a softer allyl iodine intermediate that preferentially reacts with the carbon nucleophile. Although the stereochemistry of the newly installed allylic side chain was initially unclear, subsequent experiments revealed an 11\beta configuration, likely due to the greater accessibility of the exo face of the ester enolate intermediate. Transannular alkenylation<sup>26</sup> with Pd(PPh<sub>3</sub>)<sub>4</sub> efficiently completed the F ring construction, affording the BCDF tetracyclic product 26. Initially, low yields were observed, we speculated this may attributed to the degradation of catalyst Pd(PPh3)4 during the reaction. Adding PPh3 ligand to the reaction inhibited ligand dissociation from the Pd(0) center, ultimately leading to a successful 94% yield under milder conditions. Finally, reduction of the C-17 ketone in 26 with NaBH<sub>4</sub> provided the βhydroxy compound 27 in a highly stereoselective manner. NOE experiments on 27 confirmed the 11ß configuration of the allyl side chain, as evidenced by the spatial correlations between the C-17 hydroxy methine protons and both the C-5 methylene and ester methoxy methylene protons. The whole stereochemical structure of 27 was further unambiguously established on the basis of the X-ray crystallography analysis of its ketone derivative 29, which was prepared by sequential sulfonylation of hydroxyl group to give 28 and oxidative cleavage of the exocyclic olefin by treatment with osmium tetraoxide/NMO followed by lead tetraacetate.

## Conclusions

In summary, we have developed an efficient synthetic route that, for the first time, assembles the tetracyclic BCDF ring analog of C<sub>19</sub>-DTAs in 14 steps with an overall yield of 10.8%. This synthetic route is based on the preparation of the gramscale tricyclic BCD compound 24, with key steps including the highly stereoselective reduction of the ketone intermediate 15a, followed by a Wagner–Meerwein rearrangement, then introduce an allylic side chain with high regioselectivity and stereoselectivity. Finally, a Pd-catalyzed transannular alkenylation was used to further introduce a functionalized bridged F-ring, efficiently providing the BCDF tetracyclic analog of C<sub>19</sub>-DTAs. Further experiments are in progress.

# Data availability

The data supporting this article have been included as part of the ESI. Crystallographic data for compound 29 has been

allyl isomer, although in lower yield.

Paper

deposited at the CCDC under [accession number 2362943] and can be obtained from https://deposit.ccdc.cam.ac.uk/.†

#### Conflicts of interest

There are no conflicts to declare.

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