Total Synthesis of the Lycorenine-type Amaryllidaceae Alkaloid (±)-Clivonine via a Biomimetic Ring-switch from a Lycorine-type Progenitor

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Abstract

A fully diastereoselective total synthesis of the lycorenine-type Amaryllidaceae alkaloid (±)-clivonine (19) is reported via a route that employs for the first time a biomimetic ring switch from a lycorine-type progenitor, thereby corroborating experimentally the biogenetic hypothesis first expounded for these compounds by Barton in 1960.

Introduction
The Amaryllidaceae alkaloids are a large class of naturally occurring bases isolated from herbaceous perennials such as daffodils that can mostly be classified as belonging to one of eight skeletally distinct sub-classes.\textsuperscript{1} All these alkaloids derive from a common bisphenol biosynthetic precursor, norbelladine (I, itself derived from Phe and Tyr).\textsuperscript{2} This biogenetic scheme was first enunciated by Barton in 1957.\textsuperscript{3} He used the Amaryllidaceae alkaloids to illustrate his thesis that intramolecular phenolic oxidative coupling constituted a critical diversifying step in alkaloid biosynthesis; an idea that proved correct and revolutionized our understanding of alkaloid biogenesis.\textsuperscript{2} Initially, Barton was unable to account for the tazettine\textsuperscript{4} and lycorenine sub-classes within this regime, proposing that these compounds were possibly derived from intermolecular phenolic coupling,\textsuperscript{3} but in 1960 he revised his proposal to encompass their formation by rearrangement of haemanthamine (2) and lycorine-type (I) progenitors respectively.\textsuperscript{5} Interconversion was proposed to involve benzylic oxidation (→ lactamols 3 and II) then ring-opening/bond rotation/ring closure/N-methylation (→ lactols 4 and III), a process we will refer to as ‘ring–switching’. An intramolecular crossed-Cannizzaro rearrangement (during isolation)\textsuperscript{6,7c} accounts for the conversion of pretazettine (4) to tazettine (5) whereas lactol III to lactone IV oxidation occurs in the lycorenine series (Scheme 1).

\textbf{Scheme 1.} Outline biosynthesis of tazettine and lycorenine type alkaloids from norbelladine (1) via ‘ring switching’. \textit{NB}. Precise structures and non-essential stereochemistry omitted from I-IV.
Wildman subsequently corroborated these hypotheses by tritium feeding experiments in *Sprekelia formosissima* for tazettine (5)\(^8\) and in *Narcissus* ‘King Alfred’ for lycorenine.\(^9\) Moreover, Wildman developed a biomimetic protocol for the synthesis of pretazettine (4) from haemanthidine (3)\(^6\) which has been employed in all but two\(^10\) subsequent total syntheses of tazettine\(^7\) and pretazettine.\(^11\) However, Wildman was unable to develop a corresponding protocol for biomimetic conversion of lycorine to lycorenine-type ring systems (I \(\rightarrow\) IV), noting that this conversion requires a \(\sim 180^\circ\) rotation and minimal relief of strain as compared to a \(\sim 90^\circ\) rotation accompanied by significant relief of strain in the haemanthidine/pretazettine series (3 \(\rightarrow\) 4).\(^6,7c,9\) Consequently, although Mizukami and Kotera have developed a multi-step, non-biomimetic synthetic sequence for this type of interconversion based on the von Braun reaction,\(^12\) Barton’s original hypothesis remains synthetically unverified. Herein we describe a concise, fully diastereoselective total synthesis of the lycorenine-type Amaryllidaceae alkaloid (±)-clivonine (19) from a lycorine-type progenitor 17 in which this key transformation has finally been accomplished.

**Results and Discussion**

Clivonine (19) was isolated and characterized from *Clivia miniata* Regel in 1956 by Wildman\(^13\) and its relative and absolute stereochemistry was established by Jeffs *et al.* in 1971.\(^14,15\) To date, the only synthesis of (±)-clivonine has been that of Irie in 1973 (17 steps, 0.43% overall yield from piperonal).\(^16\)

The synthesis of (±)-clivonine progenitor 15 parallels our previous synthesis of (+)-trianthine (16), employing a retro-Cope elimination\(^17\) (11 \(\rightarrow\) 12) as the key step (Scheme 2).\(^18\)
Scheme 2. Synthesis of clivonine progenitor 15.

Although trianthine (16) and clivonine progenitor 15 both have trans B-C/cis C-D ring-junctions they are diastereomeric with respect to the ring C cis-diol motif. Consequently, following 1,2-addition of aryl lithium reagent to the convex face of bicyclic enone (±)-619,20 and trapping as acetate 7 (92% yield), a one-pot Ireland-Claisen rearrangement/CH2N2 esterification was employed to relay the stereochemistry at C11b to C3a with retention of configuration (→ 8, 85% yield; cf. the vinyl cuprate S_N2' displacement with inversion of configuration employed for trianthine)18a. Ester to aldehyde reduction (DIBAL-H) then oximation (NH2OH.HCl, 82% yield, 2 steps) and oxime reduction (NaCNBH3) then afforded retro-Cope elimination substrate 11 (83% yield). Hydroxylamine 11 cyclized smoothly upon heating as a 0.014 M solution in degassed toluene at 80 °C for 17 h to provide N-hydroxyhydrindole 12 as a single stereoisomer in 98% yield.18a Hydrogenolysis of the N-O bond (Raney-Ni, 94% yield), N-formylation (HCO2COMe, 93% yield) and then Bischler-Napieralski ring B closure with concomitant acetonide deprotection (POCl3) gave water soluble iminium salt 15 after purification by ion-exchange then C18 reverse-phase solid phase extraction (SPE) (42% yield).

Prior studies in which we had been unable to obtain lactamol 17 cleanly, via lactam half-reduction (LiEtBH3) or via Polonovski reactions from the amine-N-oxide (Ac2O or TFAA), had taught us that lactamol 17 was extremely sensitive to Cannizzaro disproportionation to give a 1:1 mixture of the corresponding amine and lactam, particularly under basic conditions. Attempts to transform iminium salt 15 into the corresponding N-methyl aldehyde according to a procedure developed by Rozwadowska
for hydrastinine using MeI in MeOH,\textsuperscript{21,22} and into lactamol 17 according to procedures developed by Dostál for sanguinarine using NaOD in $d_3$-MeCN/D$_2$O\textsuperscript{23} or Na$_2$CO$_3$/D$_2$O,\textsuperscript{24} also induced substantial disproportionation. However, treatment of a solution of iminium salt 15 in $d_6$-DMSO/D$_2$O (5:1 v/v) with a solution of Cs$_2$CO$_3$ in D$_2$O (0.77 M, 1.3 equiv.) reproducibly gave clean conversion to a single, unassigned epimer of lactamol 17 in ~5 min, as evidenced by $^1$H NMR spectroscopy (15 iminium methine: $s$ @ $\delta$ ~9.07 ppm $\rightarrow$ 17 lactamol methine: $s$ @ $\delta$ ~4.92 ppm) (Scheme 3).

Scheme 3. Biomimetic ring-switch of lycorine-type progenitor 17 into clivonine (19) and the molecular structure of 19$\cdot$HCl (X-ray).

Next, we explored $N$-methylation. Wildman\textsuperscript{6} described two protocols for conversion of haemanthidine (3) to pretazettine (4): $N$-methiodide salt formation (MeI in MeOH) then careful basification of an aqueous acidic solution of this salt with K$_2$CO$_3$ and extraction into CHCl$_3$ was the method adopted (with modifications)$\textsuperscript{7,11}$ in subsequent syntheses, but Eschweiler-Clarke reductive methylation (HCO$_2$H/H$_2$CO) then basification and extraction was reportedly equally efficient. In our hands, the Eschweiler-Clarke method returned only the corresponding amine when applied to lactamol 17 whereas treatment with methanolic MeI gave a complex mixture of products containing methyl ether/acetal signals by $^1$H NMR spectroscopy. Extensive experimentation established that addition of just 1 equiv. of a dilute solution of MeI in $d_6$-DMSO to freshly prepared solution of lactamol 17/Cs$_2$CO$_3$ (in $d_6$-DMSO/D$_2$O) afforded a mixture of species of which the major component was tentatively assigned as $N$-methyl aldehyde 18 by $^1$H NMR spectroscopy (aldehyde proton: $s$ @ $\delta$ ~9.67 ppm, N-Me: $s$ @ $\delta$ ~2.35 ppm). Further optimization was confounded by the formation of what appeared to be quaternised
salts which were also formed to a greater extent when employing alternative methylating agents (e.g. Me₂SO₄, MeOTf). However, freeze-drying of this mixture, suspension of the residue in toluene and treatment with Fetizon’s reagent reproducibly afforded (±)-clivonine (19) in 32% yield after chromatography from iminium salt 15 (12 steps, 6.1% overall yield from enone 6). All spectroscopic data matched that reported for the natural material¹⁴,¹⁵,²⁵ and its molecular structure was confirmed by a single-crystal X-ray structure determination on its hydrochloride (Scheme 3).

**Conclusion**

In conclusion, we have reported the total synthesis of the lycorenine-type Amaryllidaceae alkaloid (±)-clivonine (19) via a route that employs, for the first time, a biomimetic ring switch from a lycorine-type progenitor, thereby finally corroborating experimentally the biogenetic hypothesis first expounded for these compounds by Barton 50 years ago.

We are currently exploring this approach for the synthesis of hippeastrine from lycorine¹ and investigating whether there is a causal relationship between ring-switching and the life-cycle of the herbaceous perennials in which these alkaloids are found.²⁶

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**Supporting Information Available**

Full experimental details, NMR spectra and details of the crystallographic analysis, including CIF file, of structure 19•HCl. This material is available free of charge via the internet at [http://pubs.acs.org](http://pubs.acs.org).

**References.**

† Deceased 15th March 1996.

In 1957, tazettine was thought to be a natural product. Later, Wildman showed it to be an artifact of isolation during which an intramolecular crossed-Cannizzaro rearrangement takes place; pretazettine is the natural product (refs. 6 and 7c).


(20) Enone 6 can also be prepared in enantiomerically pure form [(5S, 6S)-(+) -6] via *Pseudomonas putida* microbial oxidation of chlorobenzene, see: (a) Spivey, A. C.; Giró Mañas, C.; Mann, I. *Chem. Commun.* **2005**, *4426*-4428, (b) ref. 18a.


(26) The isolated yields of these alkaloid constituents show strong seasonal dependence (e.g. ref. 8).

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