Stemocurtisine, the first pyrido[1,2-a]azapine Stemona alkaloid

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Stemocurtisine, the First Pyrido[1,2-a]azepine Stemona Alkaloid

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A new pentacyclic stemona alkaloid, stemocurtisine (2), with a novel pyrido[1,2-
\textit{a}]azepine A,B-ring system, has been isolated from a root extract of \textit{S. curtisii} Hook. f.
The structure and relative stereochemistry was determined by spectral data
interpretation and X-ray crystallography.
The stemona group of alkaloids includes more than 40 different natural products that have been structurally classified into five different groups. The pyrrolo[1,2-a]azepine (1-azabicyclo[5.3.0]decane) nucleus is common to all compounds in these groups. For example, the relatively complex *Stemona* alkaloid, stemofoline (1), has the typical pyrrolo[1,2-a]azepine A,B-ring structure, characteristic of this class of alkaloids. A few of these alkaloids do not fit these five structural groups and have a more complex bridged structure or ring structures that most likely arises from initial oxidative cleavage of the pyrrolo[1,2-a]azepine ring system. Phytochemical studies on the Stemonaceae family have been limited to eight of the recorded 30 or more species. Extracts of the roots of *Stemona* species have been used in traditional Chinese medicine for the treatment of various respiratory diseases and as anthelmintic agents for domestic animals. More recently, extracts and the pure alkaloids derived from the extracts of the leaves and roots of *Stemona collinsae* and *S. tuberosa* were shown to have insect toxicity, antifeedent, and repellent activities. We report here the isolation and structure determination of a novel *Stemona* alkaloid (2) from the root extracts of *S. curtisii* Hook.f. growing in the southern region of Thailand. This plant is known in the Thai vernacular as “non tai yak” and has received no phytochemical and little biological attention apart from studies of its effect on the action potential of the frog sciatic nerve and its toxicity on house fly larvae (*Culex p. fatigans* and *Aedes aegypti*).

A crude ethanol extract of the roots of *S. curtisii* was partitioned between 5% hydrochloric acid solution and chloroform. The aqueous solution was made basic with aqueous ammonia and extracted with chloroform. The crude residue was subjected to flash column chromatography and preparative TLC to provide 2 in 0.5%
yield (w/w) based on the dry weight of root material. Examination of the crude ethanol extract by TLC analysis showed the presence of 2, indicating that this compound was not being produced via an acid catalysed reaction during the acid extraction process. We have named this compound stemocurtisine, based on its botanical origin. Compound 2 was obtained as colorless prismatic crystals (mp 149-151 °C) by careful and slow evaporation of a solution of 2 in ethyl acetate and diethyl ether. HRMS (CI +ve, m/z [MH]+ 347.1727, calcd 347.1733) indicated that 2 has the molecular formula C_{19}H_{25}NO_5. The ¹H and ¹³C NMR spectra of 2 indicated the presence of the C- and D-ring system that is typically found in the stemoamide group of alkaloids, including stemofoline (1). The ¹³C/DEPT NMR spectra of 2 indicated four methine carbons, and six methylene carbons and, unlike the other Stemona alkaloids, except 1 and its didehydro and 2'-hydroxy derivatives, a quaternary carbon at δ 120.4 was apparent, indicative of an acetal-like structure (C-9). Indeed the corresponding acetal carbon (C-8) in 1 occurs further upfield at δ 112.7.² The X-ray structural analysis confirmed the molecular formula of 2 and revealed its connectivity and relative stereochemistry and showed that this alkaloid has a novel pentacyclic structure based on a unique pyrido[1,2-a]azepine A,B-ring system (that is a 6,7-bicyclic A,B-ring system) and not the typical pyrrolo[1,2-a]azepine A,B-ring system (5,7-bicyclic A,B-ring system). This is the first Stemona alkaloid to have this type of base structure. The absolute configuration of 2 is not known but is assumed based on the known configurations of Stemona alkaloids with similar C,D-ring structures. The X-ray structure of 2 also showed that the piperidine A-ring adopts a chair-like conformation and is connected to the B- and C-rings through an ether bridge between C-1 and the quaternary acetal carbon C-9 readily identified in the ¹³C NMR spectrum.
The full $^1$H and $^{13}$C NMR spectral assignments for 2, based on extensive COSY, TOCSY, NOESY, HMQC, and HMBC experiments, are shown in Table 1. Of significance in the NOESY spectra of 2 were cross peaks between H-10 and H-6β, H-6β and H-8β, H-7α and H-3α, H-3β and H-2β, and H-2β and H-4β (Figure 2). These cross peaks allowed the unequivocal assignment of all methylene protons in the $^1$H NMR spectrum of 2 (Table 1) and indicated that the solution-structure of 2 is similar to the solid-state structure. That is, the A-ring adopts a chair-like conformation in which the axial proton H-3α is in close proximity to H-7α in the seven-membered ring on the concave face of the molecule (Figures 1 and 2).

In conclusion, a new pentacyclic stemona alkaloid, stemocurtisine (2), with a novel pyrido[1,2-$a$]azepine A,B-ring system has been isolated from a root extract of $S$. curtisii.

Experimental Section

General Experimental Procedures. Melting points were determined by a Reichert hot-stage melting point apparatus and are uncorrected. Optical rotations were measured using a JASCO DIP-370 polarimeter. IR spectra were obtained on a Nicolet AVATAR 300 FTIR spectrophotometer. $^1$H (500 MHz), $^{13}$C (75 MHz), and 2D NMR spectra were recorded on Varian Unity 300 and 500 spectrometers. HRCIMS were recorded on a Fison/VG Autospec-TOF-oa Mass Spectrometer (70 eV).

Plant Material. The roots of $Stemona curtisii$ were collected at Tumbol Kaunmao, Amphor Rasda, in the North of Trang Province, Thailand, in May 2002. The plant material was identified by Mr. James F. Maxwell from the
Extraction and Isolation. The dry ground root of *S. curtisii* (0.4 kg) was percolated with 95% ethanol (3 x 800 mL) over 3 days at room temperature. The ethanolic solution was evaporated to give a dark residue (50 g) that was partitioned between water and chloroform. The chloroform extract was extracted with 5% hydrochloric acid solution and the aqueous solution was made basic with aqueous ammonia and extracted with chloroform to afford 0.9 g of crude alkaloid material. This material was chromatographed on silica gel (100 mL) using gradient elution from 100% dichloromethane to 50% methanol/dichloromethane containing 1% concentrated aqueous ammonia as eluent. A total of 1500 mL of eluent was collected in test tubes of 25 mL. On the basis of TLC analysis these fractions were pooled to give 18 fractions. Fractions 12 (62.2 mg) and 13 (86.2 mg) were combined and re-chromatographed by preparative TLC (dichloromethane-methanol-aqueous ammonia, 94:5:1) to give 20.8 mg of pure stemocurtisine (2).

Stemocurtisine (2): *R*$_f$ 0.46 (Merck, silica gel 60 F254 TLC plates on aluminium backing using dichloromethane-methanol-aqueous ammonia, 94:5:1 as eluent) with detected using a UV-lamp and/or iodine vapour; colorless prismatic crystals (from EtOAc/diethyl ether); mp 149-151 °C; [*α*]$^D_{25}$ +334° ($c$ 0.66, CHCl$_3$); IR (film) $\nu_{\text{max}}$ 2932, 1741, 1618, 1021 cm$^{-1}$; $^1$H, $^{13}$C, DEPT and HMBC NMR data are presented in Table 1; HREIMS $m/z$ 347.1727 [M]$^+$, calcd for C$_{19}$H$_{25}$NO$_5$ 347.1733.
**X-ray Structure Determination:** \( \text{C}_{19}\text{H}_{25}\text{NO}_{5}, \ M = 347.4. \) Trigonal, space group \( P3_1 \ (C^3_3, \text{No. 144}), \ a = 9.046(6) \ \text{Å}, \ c = 18.222(1) \ \text{Å}, \ V = 1291.5(2) \ \text{Å}^3. \) \( D_c (Z = 3) = 1.340 \ \text{g cm}^{-3}. \) \( \mu_{\text{Mo}} = 0.97 \ \text{cm}^{-1}; \) specimen: 0.48 x 0.43 x 0.38 mm; 'T' min/max (multiscan absorption correction) = 0.95. \( 2\theta_{\text{max}} = 70^\circ; \ N_{\text{total}} = 23659, \ N_{\text{unique}} = 3751 \) \( (R_{\text{int}} = 0.037), \ N_{\text{obs}} (F > 4\sigma(F)) = 3294; \ R = 0.041, \ R_w = 0.042 \) (weights: \( \sigma^2(F) + 0.0002F^2)^{-1}) \). \( |\Delta\rho_{\text{max}}| = 0.31(3) \ \text{e Å}^{-3}. \) Bruker AXS instrument, \( \omega \)-scans; monochromatic Mo K\( \alpha \) radiation, \( \lambda = 0.71073 \ \text{Å} ; \ T \) ca. 153 K. Chirality adopted from chemical expectations, being indeterminate in the present experiment, with \( (x, y, z, U_{\text{iso}})_H \) refined throughout. CCDC # 199384.

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References and Notes


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(14) Sakata, K.; Aoki, K.; Chang, C.-F.; Sakurai, A.; Tamura, S.; Murakoshi, S. 


Table 1. $^{13}$C NMR (75 MHz) and $^1$H NMR (500 Mz) Spectral Data of 2 in CDCl$_3$ Solution.

<table>
<thead>
<tr>
<th>Position</th>
<th>$\delta_C$ (DEPT)</th>
<th>$\delta_H$ (mult., $J$ (Hz), assign.)</th>
<th>HMBC</th>
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<tr>
<td>1</td>
<td>75.5 (CH)</td>
<td>4.01 (s)</td>
<td>H-2β, H-3β, H-10, H-10a</td>
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<td>2</td>
<td>26.9 (CH$_2$)</td>
<td>1.62 (m, β) 2.21 (d, 14.5, α)</td>
<td>H-3α, H-3β, H-4α, H-4β</td>
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<tr>
<td>3</td>
<td>18.9 (CH$_2$)</td>
<td>1.21 (d, 13.5, β) 1.82 (m, α)</td>
<td>H-1, H-2β, H-4α, H-4β</td>
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<tr>
<td>4</td>
<td>53.6 (CH$_2$)</td>
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<td>3.38 (t, 13, β) 2.96 (m, α)</td>
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<td>27.0 (CH$_2$)</td>
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<td>9</td>
<td>120.4 (C)</td>
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<td>10</td>
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<td>10a</td>
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<td>124.9 (C)</td>
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CAPTION FOR X-RAY STRUCTURE:

**Figure 1.** Molecular projection of stemocurtisine (2) showing 50% probability amplitude ellipsoids for C,N,O, hydrogen atoms having arbitrary radii of 0.1 Å.
Figure 2. Molecular model (AM1, Spartan) of the A,B,C-ring substructure of 2 showing significant NOESY cross peaks (curved lines).