BAKUCHICIN, A NEW SIMPLE FURANOCOUMARIN FROM PSORALEA CORYLIFOLIA L.

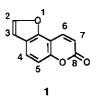
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Abstract - A new simple furanocoumarin, bakuchicin, has been isolated from the hexane-extract of seeds of Psoralea corylifolia L. (Leguminosae) alone with stigmasterol, psoralen The structure of bakuchicin has been shown as and bakuchiol. 8-oxo-8<u>H</u>-furo[2,3-<u>f</u>][1]benzopyran (1) by spectral means.

The seeds of Psoralea corylifolia L. (Leguminosae, Sanskrit name: Bakuchin) is used as a tonic or an aphrodisiac against impotence and menstruation disorders. The seed-oil of this plant also is used externally for the treatment of leucodermy, psoriasis, and leprosy in Indian folkloric remedy.¹ Many studies on the constituents of the seeds have been reported.² Now the presence of a new simple furanocoumarin was proved.

The n-hexane extract of the powdered seeds of P. corylifolia yielded an oily fraction which on silica gel chromatography gave a 0.1% of yield of a crystalline component (1), mp 138 °C, besides the known compounds, stigmasterol, psoralen (2), and bakuchiol (3). 1 is a new compound, for which we propose the name bakuchicin.



J (Hz) _δ (ppm) 2.0 C2-H 7.70 (d) C3-H 7.14 (dd) 2.0,1.25 24-H 4.44 (dd) 8.75,1.25 C5-H 7.38 (d) 8.75 C6-H 7.82 (d) 9.5 C7-H 6.40 (d) 9.5

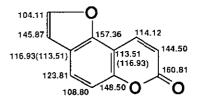
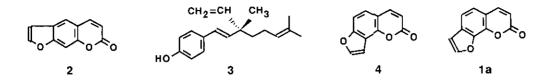
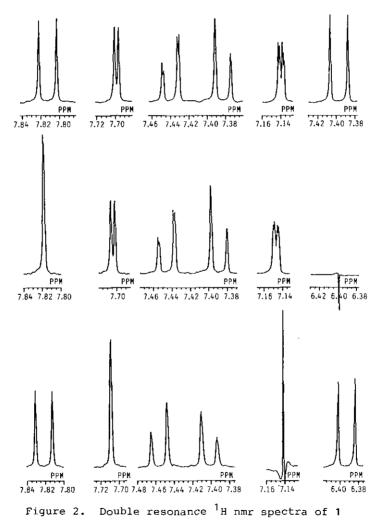


Table 1. ¹H Nmr spectral data of 1 Figure 1. ¹³C Nmr spectral data of 1



Bakuchicin (1) analyzed for $C_{11}H_6O_3$, a formula which confirmed by high-resolution mass spectrum molecular ion peak 186.0334 (Calcd 186.0317). The ultraviolet absorption spectrum (in 95% EtOH) showed maxima at 242 nm (infl.) (log ϵ 4.47), 248 nm (log ϵ 4.48) and 297 nm (log ϵ 4.13), while the infrared spectrum (in CHCl₃) displayed bands at 1722 and 1619 cm⁻¹, suggesting the presence of a furanocoumarin skeleton. This suggestion is supported by spectral data of ¹H nmr (Table 1), ¹³C nmr (Figure 1), and 2D nmr.



The COSY spectrum of bakuchicin showed that the signal at 6 7.14 (1H, dd, J=2.0 Hz, 1.25 Hz) is correlated with the signals at 6 7.44 (1H, dd, J=8.75 Hz, 1.25 Hz) and 6 7.70 (1H, d, J=2.0 Hz), but no correlation between the signals 6 7.82 and other aromatic protons. These findings, combined with a double-resonance experiment (Figure 2), showed that another structure (1a) can be ruled out and the structure of bakuchicin is elucidated as 8-0x0-8H-furo[2,3-f][1]benzopyran (1). Interestingly, previous investigators have been reported isolation of angelicin (4) from seeds of <u>P. corylifolia</u>,³ however, we proved the presence of bakuchicin (1), a new isomeric simple furanocoumarin, in this paper.

EXPERIMENTAL

Melting points were taken on a Yamato MP-2 melting point apparatus and are uncorrected. Ultraviolet spectra were recorded with a Hitachi U-3200 spectropohotometer. Infrared spectra were determined as $CHCl_3$ solutions on a Jasco A-100S infrared spectrophotometer. ¹H Nmr and ¹³C nmr spectra were recorded in $CDCl_3$ on a JEOL FX-500 spectrometer. TMS was used as an internal standard; chemical shifts are reported in δ ppm units. Mass spectra were determined with a JEOL DX-303 double focusing mass spectrometer operating at 70 eV.

<u>Isolation of substances</u> Dried and powdered seeds of <u>P. corylifolia</u> (1000 g) were exhaustively extracted with <u>n</u>-hexane (2 l x 4) at room temperature for 24 h. The <u>n</u>-hexane solution was evapolated and then subjected to column chromatography on silica gel (6 x 50 cm). Elution was accomplished with <u>n</u>-hexane and increasing quantities of AcOEt. Fractions of about 50 ml were collected and monitored by thin-layer chromatography (tlc). The four major compounds, bakuchiol **3** (52.127 g), stigmasterol (0.050 g), bakuchicin 1 (0.992 g), and psoralen **2** (0.884 g), were obtained from the corresponding fractions 10-16, 28-30, 31-35, and 37-41, respectively.

<u>Bakuchiol (3)</u> Colorless oil, bp 146-147 °C/0.8 mmHg. Ms $\underline{m}/\underline{z}$: 256 (M⁺) (Calcd 256). The identity of this compound was established by comparison of ¹H nmr spectral data.⁴ <u>Bakuchiol 3,5-dinitrobenzoate</u>: Pale yellow plates (acetone-MeOH), mp 130 °C (lif.,⁴ mp 136 °C). <u>Anal.</u> Calcd for $C_{25}H_{26}N_2O_6$: C, 66.56; H, 5.82; N, 6.22. Found: C, 66.61; H, 5.97; N, 6.14. Fdms $\underline{m}/\underline{z}$: 450 (M⁺). <u>Stigmasterol</u> Colorless plates (MeOH), mp 170 °C. <u>Anal.</u> Calcd for $C_{29}H_{48}O$: C, 84.40; H, 11.72. Found: C, 84.59; H, 11.77. Ms $\underline{m}/\underline{z}$: 412 (M⁺). <u>Bakuchicin (1)</u> Colorless needles (acetone-n-hexane), mp 138 °C. <u>Anal.</u> Calcd for $C_{11}H_6O_3$: C, 70.97; H, 3.25. Found: C, 70.97; H, 3.46. Ms <u>m/z</u>: 186.0334 (Calcd 186.0317). Ir (CHCl₃) cm⁻¹: 1722 (α , β -unsaturated lactone), 1619 (aromatic). Uv (95% EtOH) nm (log ϵ): 242 infl. (4.47), 248 (4.48), 297 (4.13). ¹H Nmr: Table 1. ¹³C Nmr: Figure 1.

<u>Psoralen (2)</u> Colorless needles (MeOH), mp 155 °C. <u>Anal.</u> Calcd for $C_{11}H_6O_3$: C, 70.97; H, 3.25. Found: C, 71.11; H, 3.29. Ms <u>m/z</u>: 186 (M⁺). The identity of this compound with an authentic sample of psoralen was confirmed by comparison of ¹H nmr, ir, uv spectra, and tlc behavior.

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