

Phytochemistry, 1973, Vol. 12, p. 1821. Pergamon Press. Printed in England.

## MINOR ALKALOIDS OF *RAUWOLFIA CAFFRA*

MUSTAFA SALMAN HABIB and WILLIAM EDWARD COURT

Postgraduate School of Studies in Pharmacy, University of Bradford, Bradford BD7 1DP

(Received 26 February 1973. Accepted 16 March 1973)

**Key Word Index**—*Rauwolfia caffra*; Apocynaceae; root: indole alkaloids: aricine: renoxidine: sarpagine: yohimbine.

*Plant. Rauwolfia caffra* Sond. *Plant part.* Root bark. *Source.* Kokstad, Pondoland, South Africa (voucher No. RAU101-571, deposited with the Collection of Materia Medica and Herbaria, University of Bradford); Louis Trichardt, Northern Transvaal (voucher No. RAU101-574). *Previous work.* Isolation of indole alkaloids reserpine, ajmaline,<sup>1</sup> serpentine,<sup>2</sup> rescinnamine,<sup>3</sup> ajmalicine and raucaffricine<sup>4</sup> from root bark.

*Present work.* Ten alkaloids were isolated; aricine, renoxidine, sarpagine and yohimbine have not previously been isolated from *R. caffra*.

### EXPERIMENTAL

*Extraction and fractionation.* Dried powdered bark was extracted with ammoniated MeOH and fractionated as described earlier.<sup>5</sup> The weakly basic fraction was extracted successively with Et<sub>2</sub>O-EtOAc-isooctane-xylene (8:1:9:3), acetone-CCl<sub>4</sub>-isooctane-light petrol. (1:7:3:9) and acetone-CCl<sub>4</sub>-isooctane-light petrol. (7:4:3:6) to yield fractions *A*, *B*, *C* and residue *D*. Fractions *A* and *B* were further separated on silica columns to yield fractions *A*<sub>1</sub>, *A*<sub>2</sub>, *B*<sub>1</sub> and *B*<sub>2</sub>. The strongly basic fraction was extracted with HOAc-acetone-light petrol. (1:20:79) to yield fraction *E* and residue *F*.

The alkaloids were recovered from the dried fractions by crystallization using various solvents and purified by recrystallization. Using co-TLC (6 systems), chromogenic reactions, m.p., m.m.p., UV, IR and MS, aricine (*A*<sub>1</sub>), ajmalicine (*A*<sub>2</sub>), reserpine (*B*<sub>1</sub>), rescinnamine (*B*<sub>2</sub>), yohimbine (*C*), raucaffricine (*D*), ajmaline (*E*) and serpentine (*F*) were identified. *Aricine*, yellow powder, m.p. 142–146°; UV<sub>max</sub>, 228, 280, 290 nm; MS *m/e* 382 (M<sup>+</sup>), 381, 351, 255, 200, 199, 187, 95. Identical with reference material (m.m.p., co-TLC, IR). *Yohimbine*, white needles, m.p. 230–232°; UV<sub>max</sub>, 227, 282, 289 nm; MS *m/e* 355, 354 (M<sup>+</sup>), 353, 295, 184, 170, 169, 156, 144, 143. Identical with reference material (m.m.p., co-TLC, IR).

Preparative TLC on silicagel plates (layers 500 μm thick) using the solvent systems acetone-CCl<sub>4</sub>-isooctane-light petrol. (7:4:3:6) and acetone-Et<sub>2</sub>NH-MeOH (7:1:2) enabled the recovery and identification of renoxidine and sarpagine. *Renoxidine*, yellow powder, m.p. 236–239°, UV<sub>max</sub>, 226, 265, min. 242 nm; IR V<sub>max</sub><sup>KBr</sup> 3500 cm<sup>-1</sup> (NH), 3000 cm<sup>-1</sup> (NH), 2500 cm<sup>-1</sup> (ester); MS *m/e* 625, 624 (M<sup>+</sup>), 623, 608, 593, 512, 414, 198, 95. *Sarpagine*, greyish-white powder, m.p. 345–348°; UV<sub>max</sub>, 223, 274, min. 242 nm; IR V<sub>max</sub><sup>KBr</sup> 3450 cm<sup>-1</sup> (NH, OH), 1580 cm<sup>-1</sup> (Ph-N<); MS *m/e* 310 (M<sup>+</sup>), 196, 185, 183, 149, 97.

*Acknowledgement*—The authors are grateful to the Department of Forestry, South Africa for the supply of plant material.

<sup>1</sup> SCHULER, B. O. G. and WARREN, F. L. (1956) *J. Chem. Soc.* 215.

<sup>2</sup> PARIS, R. A. and DILLEMANN, G. (1956) *Ann. Pharm. Franc.* **14**, 505.

<sup>3</sup> PARIS, R. A. and DILLEMANN, G. (1957) *Ann. Pharm. Franc.* **15**, 310.

<sup>4</sup> KHAN, N. H., KHAN, M. A. and SIDDIQUI, S. (1965) *Pakistan J. Sci. Ind. Res.* **8**, 23.

<sup>5</sup> COURT, W. E., EVANS, W. C. and TREASE, G. E. (1958) *J. Pharm. Pharmacol.* **10**, 380.