SHORT COMMUNICATION

# Pharmacognostical examination of the roots and stems of *Rauwolfia mannii* Stapf

#### WILLIAM E. COURT

Pharmacognosy Laboratories, School of Pharmacy, University of Bradford, Yorks, U.K.

In previous communications the occurrence, indigenous usage, anatomy and chromatographic investigation of *R. rosea* K. Schum. have been described (Khalil, Court & Stewart, 1967; Court, Khalil & Stewart, 1967). Brief reference was made to the closely related species *R. mannii* Stapf and it was apparent that a detailed investigation was necessary to establish any differences between the species.

R. mannii was described by Stapf in 1894 and recorded with R. cardiocarpa K. Schum., R. preussii K. Schum. and R. rosea in Supplement I of the Kew Index (1886–1895). R. cardiocarpa and R. preussii are now considered synonymous with R. mannii (Feuell, 1955; Bisset, 1958). Confusion over the separate identities of R. mannii and R. rosea is related to the original morphological basis of differentiation, which depends on variations in the form of branches and peduncles (Stapf in Thiselton-Dyer, 1904). Pichon (1947) grouped both species in the section Afrovolfia of his classification of the genus. Boutique & Monseur (1955) observed close similarity between R. mannii growing in the Eastern Congo and R. rosea from the Usambaras, Tanzania, although synonymy has not been established (Bisset, 1958). Whereas Poisson (1965) does record the species as synonymous, East African botanists and the Royal Botanic Gardens, Kew (private correspondence) accept the species as distinct.

The presence of reserpine (0.04%),  $\delta$ -yohimbine (ajmalicine), reserpiline, ajmaline and serpentine has been reported in the roots of R. rosea (Korzun, St. André & Ulshafer, 1957; Court & others, 1967) whilst reserpine, rescinnamine, reserpiline, ajmaline and serpentine have been observed in R. mannii (Court & others, 1967). Other authors (Kaiser & Popelak, 1959) claim serpentine and its isomer alstonine to be absent.

### Habitat

R. mannii is a shrub attaining a height of 1-3 m at altitudes up to 2,300 m in forest and scrubland in Southern Nigeria, Cameroun, Gabon, Ruanda-Urundi and the Congo (Thiselton-Dyer, 1904; Dubois, 1955). R. preussii was reported indigenous to Southern Nigeria and French Cameroun, but no authentic specimens are available.

# Indigenous usage

R. mannii is known in West Africa as Mbara and may be occasionally used as an arrow poison supplement (Lewin, 1923). The powdered bark may be applied to wounds (de Wildemann, 1939).

## Plant material

The material used in this investigation was:

R. mannii roots and stems, herbarium specimens from Nigeria, obtained by the Tropical Products Institute, London, 1964; R. mannii roots and stems collected in Yaounde, Cameroun, 1968; R. rosea roots and stems collected in Tanzania in 1962 and verified by the East African Herbarium, Nairobi, Kenya; R. rosea roots and stems collected in Lushoto, Tanzania in 1964 and verified by the East African Herbarium.

# Microscopical examination

Measurements of cells in radial, tangential and longitudinal planes and of cell inclusions and cells isolated by maceration, were obtained as described for *R. rosea* (Khalil & others, 1967). Methods for the determination of vessel diameter index values and vessel element length were as described earlier (Court, 1967).

#### Results and discussion

The tissue arrangement and distribution in the axis of *R. mannii* is similar to that described for *R. rosea* (Khalil & others, 1967) although sclereid development is more pronounced in *R. mannii* and can be related to the greater height of this species.

Comparison of cell dimensions corresponding to those reported for R. rosea indicates fair agreement except for the root vessel measurements. For R. mannii, R = 21 to 30 to 42 to 63  $\mu$ m and T = 21 to 30 to 42 to 68  $\mu$ m and for R. rosea, R = 25 to 34 to 55 to 84  $\mu$ m and T = 21 to 29 to 50 to 67  $\mu$ m where R and R refer to measurements made in the radial and tangential planes respectively.

Quantitative microscopy confirmed that the root vessel diameters are smaller for R. mannii, although the vessel element lengths are similar (Table 1). The laborious technique of vessel diameter index differentiates the two species. For R. mannii the  $60 \mu m$  index is less than 10% and for R. rosea greater than 30%.

Stapf (1894) has recorded that young branches of R. mannii are quadrangular with more or less conspicuous decurrent raised lines; Schumann (Engler, 1895) describes

Table 1. Vessel index determinations (Vessel index = percentage of elements exceeding critical diameter) and vessel element length.

	V	essel index det	erminations		
			Critical diameter	•	Number of
Species		30 μm	60 μm	90 μm	specimens
R. mannii (Cameroun) R. mannii (Nigeria) R. rosea (Tanzania, 1962) R. rosea (Tanzania, 1964)	   (Ar	$94.7 \pm 0.4$ $94.7 \pm 0.3$ $99.8 \pm 0.1$ $99.8 \pm 0.2$ ithmetic mean	$1.5 \pm 0.4$ $5.9 \pm 0.8$ $44.4 \pm 2.3$ $41.0 \pm 2.5$ $\pm$ Standard error	$\begin{array}{c} 0 \\ 0 \\ 2.4 \pm 0.2 \\ 2.2 \pm 0.6 \end{array}$	10 10 9 9
		Vessel eler	nent length		
					Mumban

Species	Range	Mean	Standard deviation	Number of samples	of observa- tions
R. mannii (Cameroun) R. mannii (Nigeria) R. rosea (Tanzania, 1962) R. rosea (Tanzania, 1964)	58 <b>-274-618-</b> 840 μm 116 <b>-341-661-</b> 971 μm 50 <b>-296-566-</b> 994 μm 85 <b>-355-709-</b> 1,022 μm	501 μm 421 μm	172 μm 160 μm 145μ m 177 μm	6 6 6	900 900 300 300

the young branches of R. rosea as terete without decurrent lines. Examination of the available specimens confirms this differentiation.

Thin layer chromatographic examination of the specimens available using the methods described earlier (Court & others, 1967) again indicated the presence of rescinnamine in *R. mannii* but not *R. rosea*.

Khalil & others (1967) commented on the similarity between R. rosea and R. volkensii Stapf (now known to be R. oreogiton Mgf.) and R. obscura K. Schum. roots. These species exhibit small vessels and limited sclereid development and R. mannii must be considered in this group. The main features are compared in Table 2.

Table 2. Comparison of the main features of four species of rauwolfia.

	Normal		,	
Species	height	Vessel diameter	Sclereid development	Reference
R. mannii	2–3 m	R 21 <b>-30-42</b> -59 μm T 21 <b>-30-42</b> -68 μm	Isolated or groups of up to 12	This work
R. rosea	0·5–2 m	R 25 <b>-34-55</b> -84 μm T 21 <b>-29-50</b> -67 μm	Isolated or small groups. Rare in roots below 1 cm diameter	Khalil & others, 1967
R. obscura	1–1·5 m	R 52–56 μm T 35–50 μm	Absent	Paris & Dillemann, 1956
R. oreogiton	2 m	R 26–44–56–96 μm T 26–37–48–78 μm	Isolated or small groups of up to 10. Rare except in specimens exceeding 4 cm diameter	Court, 1961

It is concluded that although similar in structure, the roots of *R. mannii* and *R. rosea* may be identified by careful investigation of sclereid development, vessel diameter index and by thin layer chromatography.

## REFERENCES

BISSET, N. G. (1958). Ann. Bogor., 3, Part 1, 233.

BOUTIQUE, R. & MONSEUR, X. (1955). Bull. agric. Congo belge, 46, 271-280.

COURT, W. E. (1961). J. Pharm. Pharmac., 13, 422-434.

COURT, W. E. (1967). Can. J. pharm. Sci., 2, 68-71.

COURT, W. E., KHALIL, A. A. & STEWART, A. F. (1967). Planta Med., 15, 173-178.

Dubois, L. (1955). Bull agric. Congo Belge, 46, 567-595.

ENGLER, A. (1895). Pflanzenw. Ost. Afr. C., 317.

FEUELL, A. J. (1955). Colon. Pl. Anim. Prod., 5, 1-33.

Kaiser, F. & Popelak, A. (1959). Chem. Ber., 92, 278-287.

KHALIL, A. A., COURT, W. E. & STEWART, A. F. (1967). Planta Med., 15, 104-117.

Korzun, B. P., St. André, A. F. & Ulshafer, P. R. (1957). J. Am. pharm. Ass. (Sci. Edn.), 46, 720-723.

LEWIN, L. (1923). Die Pfeilgifte (Leipzig: J. A. Barth), 259.

Paris, R. & Dillemann, G. (1956). Ann. pharm. fr., 14, 505-518.

PICHON, M. (1947). Bull. Soc. bot. Fr., 94, 31-39.

Poisson, J. (1965). Ibid., 112, 162-174.

STAPF, O. (1894). Kew Bull., 21.

THISELTON-DYER, W. T. (1904). Flora of Tropical Africa Vol. IV, i, 113-114. London: Lovell Reeve and Co., Ltd.

DE WILDEMAN, E. (1939). Mém. Inst. r. colon. belge, Sect. Sci. nat. méd., 9, 330.