Comunications/Brief Reports

ISOLATION AND IDENTIFICATION.-Bark (150 g) from freshly collected roots was subjected to steam distillution and the distillate multiply extracted with CH₂Cl₂. After drying over Na₂SO₄, the CH₂Cl₂ was removed at room temperature in a nitrogen stream to yield 140 mg of a pale yellow liquid. Capilliary gc (22m × 0.25mm DBI, 70°-250° at 5°/min, flow 0.7 ml/min, He) demonstrated the liquid to be an essentially pure (>99.9%) compound. The retention time (8.14 min), mass spectrum, 60 MHz 1 Hnmr, and ir of the substance were identical with those of authentic 2-hydroxyacetophenone (Aldrich Chemical Co., Inc.).

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A FLAVONE WITH ANTIINFLAMMATORY ACTIVITY FROM THE ROOTS OF RHUS UNDULATA¹

THEUNIS G. FOURIE* and FRIEDRICH O. SNYCKERS

Central Research Laboratory, Noristan Limited, Private Bag X516, Silverton, 0127, Pretoria, Republic of South Africa

In the course of our search for biologically active compounds from indigenous South African flora, we have investigated the roots of Rhus undulata Jacq. var. undulata (Anacardiaceae, Kuni-bush), a tree widespread in southern Africa (1). Plant material and extracts were worked up in accordance with our normal procedures (see Experimental section) (2) and yielded 5-hydroxy-4',7-dimethoxyflavone (apigenin dimethylether) in addition to some nonpolar compounds that were not characterized.

Apigenin dimethylether showed a 25% inhibition (75 mg/kg dose) [phenylbutazone (reference), 81% inhibition at 75 mg/kg dose] of the phlogistic response (carrageenan-induced edema) (3) in the rat, an activity reported (4-6) for several closely related flavonoids (Table 1). The presence of antiinflammatory activity may provide an explanation for the claims made with regard to the therapeutic value of R. undulata roots in infective disorders of the gastrointestinal tract.

TABLE 1.	Antiinflammatory A	ctivity of Some	Flavonoids	Compared v	vith Apigenin	Dimethylether

Test Compound	ED ₂₅ (mg/kg)	Potency (Antiinflammatory units/g)
Naringin	Inactive	0
Nobiletin	20	50
Hydrocortisone phosphate (reference)	13.5	74
Apigenin dimethylether	75	13

¹Part 3 in the series "Studies of South African Medicinal Plants." For Part 2, see S. Afr. J. Chem., 36, 114 (1983).

EXPERIMENTAL

PLANT MATERIAL.—The roots of *R. undulata* were collected April 8, 1976, at Hennopspride near Pretoria. Voucher specimen (no. 882) is deposited in the Botanical Research Institute, Pretoria.

EXTRACTION AND FRACTIONATION.—Air-dried, milled, roots of *R. undulata* (3.5 kg) were successively extracted with C_6H_6 (235 g extract), EtOAc (83 g extract) and MeOH (279 g extract) at room temperature for 48 h. After removal of the solvents, the crude extracts were fractionated separately over silica gel (Kieselgel 60, 70-230 mesh; Merck). Elution was conducted with mixtures of petroleum ether, EtOAc, and MeOH of increasing polarity. Fractions with corresponding Rf values on the tlc [petroleum ether-EtOAc (1:1)] were combined into three groups. Of the groups obtained, group 2 was found to exhibit antiinflammatory activity.

ISOLATION OF APIGENIN DIMETHYLETHER.—The active group was chromatographed over silica gel and elution with C_6H_6 gave the title compound that crystallized from EtOAc as fine yellow needles (1.1 g; 0.18% of total extract), mp 171°-172° [Lit (7) mp 170°-171°]; ir ν max (KBr) 3450, 1665, 1605, 1510, 1338, 1310, 1270, 1215, 1190, 1185, 1160, 1022, 1012, 830, 815, and 760 cm⁻¹; ¹H-nmr (CDCl₃) δ 12.67 (1H, s, disappears on deuteration, OH), 7.83 (2H, dd, J=2.5 Hz and J=8.5 Hz, H-2', 6'), 6.99 (2H, dd, J=2.5 Hz and J=8.5 Hz, H-3', 5'), 6.55 (1H, s, H-3), 6.46 (1H, d, J=2.5 Hz, H-8), 6.33 (1H, d, J=2.5 Hz, H-6), 3.8 (6H, s, 2×OMe); ms m/z (%) 398 M+ (100).

IDENTIFICATION OF APIGENIN DIMETHYLETHER.—The physical data of apigenin dimethylether are in agreement with those reported in the literature (7,8).

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FIVE COUMARINS AND A CARBAZOLE ALKALOID FROM THE ROOT BARK OF CLAUSENA HARMANDIANA

J.D. WANGBOONSKUL, S. PUMMANGURA,* and C. CHAICHANTIPYUTH

Department of Pharmaceutical Chemistry, Faculty of Pharmaceutical Sciences, Chulalongkorn University, Bangkok 10500, Thailand

Clausena harmandiana Pierre (Rutaceae) is a reputed folk medicine, decoctions of the roots being used as a stomachic and antipyretic. The root bark of this species has yielded five known coumarins and a carbazole alkaloid, which are reported for the first time from this species.

EXPERIMENTAL

GENERAL EXPERIMENTAL PROCEDURES.—Spectral data were obtained with the following instruments: a Perkin-Elmer 283 grating infrared spectrophotometer; a JEOL FX90Q (90 MHz nmr spectrometer); a Shimadzu UV-180 spectrophotometer; and a JEOL DX 300 mass spectrometer.