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RUTACEAE

CONSTITUENTS OF THE LEAF AND PEEL OILS OF CITRUS HYSTRIX, D.C.

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Plant. Citrus hystrix D.C.

Source. Thailand (local name 'makrud').

Uses. As a perfume for hair dressings and shampoos.^{1,2}

Previous Work. The presence of citronellal (84%) and terpene alcohols (leaf oil)3 and citronellal (26%) and terpene hydrocarbons (peel oil)4 have been previously reported.

Leaf. The fresh leaves were steam distilled yielding a volatile oil which had the following properties: spec. grav. 0.8580; r.i, 1,4500 and specific rotation $-10^{\circ}44'$ (all at 20°). The oil was analysed by a combination of fractional distillation, alumina-column chromatography and GLC. Compounds isolated by preparative GLC were characterized by i.r. spectroscopy and retention time comparison.^{5,6} The oil was found to contain: α-pinene (0.2%), camphene (trace), β -pinene and sabinene (4.9), myrcene (0.6), limonene (0.6), trans-ocimene (0·3), y-terpinene (0·2), p-cymene (0·1), terpinolene (0·2), citronellal (65·4), copaene (0·1), linalool (2·9), β -cubebene (0·1), isopulegol (4·9), caryophyllene (0·4), citronellyl acetate (5·1), citronellol, geranyl acetate and δ-cadiene (6 4). All percentages were calculated from disc integration measurements using a flame ionization detection of a pressure and temperature programmed capillary gas chromatographic analysis using a column of Carbowax 6000.

Peel. The oil, obtained from a cold press of the peel of fresh fruit which had the following properties: spec. grav. 0.8680; r.i, 1,4729; specific rotation $+26^{\circ}17'$, was analysed as described above. The following compounds were found to be present: α -pinene (2.5%), camphene (0·2), β -pinene (30·6), sabinene (22·6), myrcene (1·4), limonene (29·2), 1,8 cineol (1·3), y-terpinene (0·1), p-cymene (0·1), terpinolene (0·1), trans-sabinene hydrate (0.6), citronellal (4.2), copaene (0.6), linalool (0.5), β -cubebene (0.5), terpinen-4-ol and β -elemene (0.2), caryophyllene (0.3), citronellyl acetate (0.2), α -terpineol (0.7), geranial

¹ I. H. Burkhill, A Dictionary of the Economic Products of the Malay Peninsula, 2nd Ed., Vol. 1, p. 574, Ministry of Agriculture and Co-operatives, Kuala Lumpar, Malaysia (1966).

W. H. Brown, Useful Plants of the Philippines, Technical Bull., Vol. 2, p. 200, No. 10, Department of Agriculture and Natural Resources, Manila, Philippines (1964).

³ M. IGOLEN, Parfum Cosmet Savons 1, 51 (1958).

⁴ S. S. TANCHICO and A. P. WEST, *Philipp J Sci* 52, 263 (1933).
⁵ B. M. LAWRENCE, J. W. HOGG and S. J. TERHUNE, *Perfume. Essent. Oil Rec.* 60, 88 (1969)

⁶ B. M. LAWRENCE, J. W. HOGG and S. J. TERHUNE, J. Chromatog. 50, 59 (1970).

(0·1), geranyl acetate and citronellol (0·4), δ -cadinene (0·3), geraniol (0·1), nerolidol (0·1) and elemol (0·3).

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TROCHODENDRACEAE

CONSTITUENTS OF THE WOOD OF TROCHODENDRON ARALIOIDES

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Plant. Trochodendron aralioides Sieb. et Zucc.-Trochodendraceae

Uses. Not known.

Previous work. Resin.1

Wood. Extracted with MeOH. Chromatographed using S₁O₂.

Acetyloleanolic aldehyde. C₃₂H₅₀O₃, m.p. 226-227°. Reduction with LiAlH₄ followed by acetylation to erythrodiol diacetate, m.p., mixed m.p., superposable i.r. spectra.

Betulin. M.p., mixed m.p., i.r.

Unidentified compounds. A: m.p. 198–204°, i.r. ν^{KBr} 3400, 1700, 1660 cm⁻¹. B: m.p. 128–130°, u.v. λ_{max} 251 nm (ϵ 10000). i.r. ν^{KBr} 1732, 1652 cm⁻¹. C: m.p. 134–135°, i.r. ν^{KBr} 3400 cm⁻¹.

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¹ K. YAGISHITA, Bull. Agri. Chem. Soc. Japan 21, 77 (1957).