NAPHTHAQUINONES AND TRITERPENOIDS OF EUCLEA DIVINORUM

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Key Word Index—Euclea divinorum; Ebenaceae; naphthaquinones; triterpenoids; diosindigo A; 2-methylnaphthazarin; 7-methyl-juglone; diospyrin; lupeol; betulin.

Plant. Euclea divinorum Hiern was collected 38 km from Lourenço Marques, Mozambique, on the route to Namaacha. Voucher specimens are deposited in the Herbarium of the Laboratory of Botany, University of Lourenço Marques. Plant material examined. Roots. Previous work. A sample of the plant, from South Africa, shown to contain 7-methyljuglone, mamegakinone, diospyrin and isodiospyrin in the roots but the stems and fruits contained no quinones [1].

Present work. Dried powdered roots of E. divinorum were extracted (Soxhlet) with petrol (bp 50-70°). The extract was concentrated under red pres and submitted to preparative TLC on silica in petrol saturated with MeOH. Three bands were formed, I, II, III, by order of decreasing R_f values. The silica corresponding to each band was removed from the plates and extracted with CHCl₃, the chloroform extracts then being rechromatographed under the same conditions. Band I yielded diosindigo A and band II 2-methylnaphthazarin. The CHCl₃ extract corresponding to band III was chromatographed on silica in CHCl₃, yielding two bands (IIIa, IIIb). Band IIIa yielded 7-methyljuglone and band IIIb diospyrin. All substances were identified by direct comparison with authentic samples (mp, TLC, UV, IR). It is interesting to note that 2-methylnaphthazarin, a somewhat rare substance [2], was found earlier by us in E. lanceolata E. Mey ex D C: [3] and in E. pseudebenus E. Mey ex DC. [4]. Diosindigo A [5], a common constituent of Diospyros spp. [6], has also been isolated from four Euclea spp. [1,7].

Another portion of the petrol extract was chromatographed over silicic acid, and the column eluted with solvents of increasing polarity. Petrol-C₆H₆ (8:2)

removed triterpenoid compounds. The solvent was evaporated under red pres and residue obtained was chromatographed over alumina. Elution with petrol– C_6H_6 (3:7; 2:8) gave lupeol, $M^+=426$, mp 214–215° (MeOH and Me₂CO), $[\alpha]_D^{20}+27.3^\circ$; acetate, $M^+=468$, mp = 212° (Me₂CO), $[\alpha]_D^{20}+47.1^\circ$ (CHCl₃) while C_6H_6 –CHCl₃ (2:8) and CHCl₃ yielded betulin, $M^+=442$, mp 258–260° (Me₂CO) $[\alpha]_D^{20}+19^\circ$ (pyridine); diacetate $M^+=526$, mp 219–220° (Me₂CO), $[\alpha]_D^{20}+21.9$ (CHCl₃). The identifications were confirmed in all cases by direct comparison with authentic samples (TLC, IR, NMR).

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STEROLS AND TRITERPENES FROM THE FRUIT OF ARTOCARPUS ALTILIS

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Key Word Index—Artocarpus altilis; Moraceae; breadfruit; sterols; triterpenes; cycloartenol; α -amyrin; cycloart-23-ene-3 β ,25-diol; cycloart-25-ene-3 β ,24-diol.

Plant. Artocarpus altilis. Source. Hunts Point Fruit Market, Bronx, New York. Use. Foodstuff. Previous work. Isolation of cyclopropane sterols from the bark and sister species [1].

Present work. The fresh fruit (1.4 kg) was extracted with CHCl₃. The solvent evaporated to give a residue (6.8 g) which was saponified. The non-saponifiable extract (3.3 g) in Et₂O was subjected to preparative layer