The *n*-BuOH concentrate was chromatographed on cellulose (preparative tlc) with *n*-BuOH-HOAc- H_2O (4:1:5, upper phase). The above four flavonoids were isolated. Schaftoside and isoschaftoside were purified by preparative tlc on cellulose with 15% HOAc. All flavonoids were identified by comparison of uv and ms of their permethylated ethers with published values (5-7). They have been isolated from any sources (8).

The number of Rutaceae species that have been investigated form C-glycosylflavones is too few to determine whether the compounds identified from M. trifolia may have taxonomic significance. In the Cuspariae, most of the C-glycosylflavones identified are apigenin derivatives (9, 10).

Details of the isolation and identifications are available from the senior author.

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DITERPENES FROM VIGUIERA PORTERI

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Approximately 25 species of the large genus Viguiera (Compositae, tribe Heliantheae, subtribe Helianthinae) have been studied chemically (1-19). Characteristic constituents are heliangolides incorporating a furanone ring and diterpene acids of the ent-kaurane and ent-trachelobane series, although not all species elaborate both types of compounds. The sole representative of the genus in the southeastern U.S. is Viguiera porteri (A. Gray) Blake, which has a limited distribution in the Piedmont plateau of Georgia and southeastern Alabama. In keeping with the chemistry of other representatives of Viguiera and the closely related genus Helianthus, our examination of V. porteri has furnished the diterpene acids 1a-1d and 2. Acids 1c and 1d were isolated in the form of the methyl esters 1e and 1f. Other compounds present were β-sitosterol; stigmasterol; and linoleic, linolenic, and stearic acids. Sesquiterpene lactones were not detected.

1a R, R', R"=H

1b R, R"=H, R'=OH

1c R=OH, R'=OAng, R''=H

1d R, R'=H, R'=OEpoxyang

1e R=OH, R'=OAng, R''=Me

1f R=H, R'=OEpoxyang, R"=Me

2

EXPERIMENTAL

PLANT MATERIAL.—V. porteri was collected by Dr. R.K. Godfrey on September 13, 1980, in the vicinity of the Dialtown Community on the Gum Creek Road near Oxford, Newton Co., Georgia (RKG #78141 on deposit in the Florida State University herbarium).

EXTRACTION AND ISOLATION.—Dried aerial parts (0.75 kg) were extracted with CHCl₃. The usual work up (20) gave 9 g of gum, which was adsorbed on 15 g of Si gel (EM Reagent, 70-230 mesh) and chromatographed over 250 g of the same adsorbent set in C₆H₆, 250-ml fractions being collected as follows: fractions 1-12 (C₆H₆), fractions 13-24 (C₆H₆-CHCl₃, 1:1), fractions 25-32 (CHCl₃), fractions 33-36 (CHCl₃-MeOH, 49:1), fractions 37-40 (CHCl₃-MeOH, 19:1), and fractions 41-48 (CHCl₃-MeOH, 9:1).

Fractions 9-16 (0.72 g) were mainly ${\bf 1a}$. Purification of fractions 17-18 (0.12 g) by tlc (C_6H_6 -EtOAc, 19:1) afforded from the upper band 55 mg of ${\bf 1a}$, identical with authentic material (21), and 47 mg of a 1:1 mixture of β -sitosterol and stigmasterol from the lower band. Fraction 28 (1.6 g) showed several spots on tlc. Esterification of ca. 0.2 g of this mixture with CH_2N_2 followed by tlc (C_6H_6 -EtOAC, 19:1, three developments) gave a mixture (60 mg) of methyl linoleate, linolenate, and stearate (by 1H nmr and ms); 26 mg of ${\bf 1f}$, identified by comparison with authentic material (21); and 32 mg of ${\bf 1e}$, identified by comparison with authentic material (21). Fraction 31, upon standing in $CHCl_3$, furnished 14 mg of a 1:1 mixture of ${\bf 1b}$ (21) and ${\bf 2}$ (21).

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