SECONDARY METABOLITES FROM CHILEAN BACCHARIS SPECIES

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In a search for biologically active compounds from Chilean plants, we have studied the composition of extracts from *Baccharis linearis* Ruiz et Pav, *Baccharis rhomboidalis* Remy, and *Baccharis solieri* Remy (Compositae). The ethanolic extracts afforded a novel chromene 1 along with seventeen known compounds (Table 1).

Chromenes are of common occurrence in the Compositae (1), and the methyl ester of compound 1 was recently reported from a Werneria species (2). In addition to chromene 1, oleanolic acid, stigmasterol, β -amyrin, and friedelinol were also present in varying amounts in the three Baccharis species (Table 1).

EXPERIMENTAL

PLANT MATERIALS.—The plants were collected at the base of the mountains near Santiago, Chile, in October 1984. Voucher specimens are kept at the Private Herbarium of M. Mahú, Departamento de Ecología, Universidad de Chile, Santiago, Chile.

EXTRACTION AND ISOLATION.—Finely

ground stems and leaves (500 g) were defatted with light petroleum ether followed by exhaustive extraction with EtOH at room temperature. Evaporation ofthe solvent under reduced pressure afforded the ethanolic extract, which was partitioned between CHCl₃ and H₂O-MeOH (9:1). THe CHCl₃ was then evaporated to dryness and the residue partitioned between equal volumes of MeOH-H₂O (9:1) and petroleum ether (60-80°). The petrol-soluble fractions yielded on evaporation extract A. Evaporation of the alcoholic layer afforded gummy extract B.

Repeated column chromatography of the extracts using silica gel and mixtures of petroleum ether/EtOAc or CHCl₃/MeOH as eluents allowed the separation of the reported compounds. Further purification was achieved by using preparative tlc.

Chromene (1) gave mp 191-192°; ir max (KBr) $cm^{-1} 3000 (s), 2800-3100 (br), 1670 (s), 1610$ (s), 1410, 1430, 1490, 820; ¹H nmr (80 MHz, CDCl₃) δ 1.48 (6H, s, Me), 5.58 (1H, d, J=10 Hz, H-3), 6.27 (1H, d, J=10 Hz, H-4), 6.20 (1H, d, J=16 Hz, H-10), 7.61 (1H, d, J=16)Hz, H-9), 6.69 (1H, d, J=9 Hz, H-8), 7.19 (1H, d, J=9 Hz, H-5), 7.23 (1H, dd, J=2.9)Hz, H-7); ms m/z 230 (M⁺, 63%), 215 (100%), 169 (26.4%), 144 (11.2%), 141 (11.3%), 115 (32%), 77 (11.6%); ¹³C nmr (20.15 MHz, DMSO- d_6 -CDCl₃) δ 173.0 (C-11), 155.6 (s, C-8a), 146.9 (d, C-9), 131.4 (d, C-4), 129.9 (d, C-7), 126.8 (s, C-4), 126.5 (d, C-5), 121.7 (C-3), 116.9 (d, C-10), 114.6 (d, C-8), 77.2 (s, C-2), 28.3 (q, C-12, C-13).

The known compounds were identified by direct comparison with authentic samples or comparisons with literature data using the usual techniques (uv, ir, ¹H nmr, ¹³C nmr, ms, mp). Full details of the isolation and identification of the compounds are available on request.

TABLE 1. Compounds Isolated from Baccharis Species^a

B. linearis	spathulenol (3), lachnophyllum ester (4), chromene (2), erythrodiol (5), axillarin (6-8).
B. rhomboidalis	* * *
B. solieri	
	tettamethoxynavone (10).

^{*}Oleanolic acid, stigmasterol, \beta-amyrin, and friedelinol were also present in all three species.

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