

COUMARINS FROM *PEREZIA CARTHAMOIDES*

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As part of a general phytochemical investigation of the Argentinian region of Cuyo, we studied *Perezia carthamooides* (D. Don) Hook. et Arn. (Compositae), this being the third species of the South American section *Perezia* (1) of the genus *Perezia* that has been investigated.

Chromatographic separation of the partitioned methanolic extracts of the dried roots yielded 3,4,8-trimethoxy-5-methylcoumarin and 8-hydroxy-4-methoxy-5-methylcoumarin. Both compounds were identified by ir, <sup>1</sup>H nmr, <sup>13</sup>C nmr, ms, and direct comparison. The trimethoxymethylcoumarin was previously isolated from *Perezia multiflora* (2,3) collected in Ecuador, while the other was isolated from *Perezia coerulescens* (4) collected in Perú. Highly oxidized coumarins may be chemotaxonomically significant metabolites of South-American *Pereziae*.

## EXPERIMENTAL

**PLANT MATERIAL.**—The plant was collected in the vicinity of the Laguna Horcones, Mendoza, Argentina in March 1986. A voucher specimen (Ambrosetti 1665) is deposited in the Ruiz Leal herbarium (MERL), Mendoza, Argentina.

**EXTRACTION AND ISOLATION.**—The air-dried roots (1.2 kg) were ground and extracted at room temperature with MeOH (3 times × 24 h), and the solvent was removed under reduced pressure. The crude extract was partitioned between MeOH-H<sub>2</sub>O (9:1) and *n*-hexane and, after removal of the hexane soluble part, between MeOH-H<sub>2</sub>O (7:3) and CHCl<sub>3</sub>. The CHCl<sub>3</sub> layer was adsorbed on Si gel packed in C<sub>6</sub>H<sub>6</sub> and eluted with C<sub>6</sub>H<sub>6</sub>/EtOAc mixtures of increasing polarity. The early fractions contained 3,4,8-trimethoxy-5-methylcoumarin, and the later fractions gave 8-hydroxy-4-methylcoumarin. Complete purification of the compounds was achieved by rechromatography followed by crystallization to afford 17 and 22 mg, respectively.

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## LITERATURE CITED

1. B.S. Vuilleumier, "The Systematics and Evolution of *Perezia* Section *Perezia*," The Gray Herbarium of Harvard, Cambridge, 1970.
2. P. Joseph-Nathan, J. Hidalgo, and D. Abramo-Bruno, *Phytochemistry*, **17**, 583 (1978).
3. F. Bohlmann and C. Zdero, *Phytochemistry*, **16**, 239 (1977).
4. L.R. Angeles, O. Look de U., I.C. Salkeld, and P. Joseph-Nathan, *Phytochemistry*, **23**, 2094 (1984).

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