## PYRROLIZIDINE ALKALOIDS FROM FIVE SENECIO SPECIES

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In previous papers we have described the pyrrolizidine alkaloids isolated from roots of Senecio subulatus (1), Senecio glandulosus (1), Senecio uspallatensis (2), and Senecio gilliesiano (3).

As a part of our continuing investigation of Argentine Senecio species, we now report the isolation and identification of the pyrrolizidine alkaloids obtained from the roots of Senecio argentino Baker (vira-vira Hieron), Senecio filaginoides (H. et A.) DC, Senecio seratophiloides Griseb, Senecio leucostachys Baker, and Senecio ragonesei Cabr.

The alkaloidal content of the studied species as well as those reported previously (1-3) is summarized in Table 1. The common alkaloids, senecionine or retrorsine, have been isolated from almost all the species studied. Uspallatine, which was recently reported (2), has been isolated from four of the species studied. The rare alkaloid, senecivernine, previously isolated only from *Senecio vernalis* (4), has been tentatively identified from *Senecio seratophiloides*.

#### **EXPERIMENTAL**

GENERAL EXPERIMENTAL PROCEDURES. — Mps were recorded on a Leitz hot-stage apparatus, the ms were recorded in a Varian MAT 112 S, the <sup>1</sup>H-nmr spectra were recorded on a Varian EM 360 A (60 MHz) in CDCl<sub>3</sub>, the <sup>13</sup>C-nmr spectra were recorded in a Bruker WR-80 (20 MHz) in CDCl<sub>3</sub>. Column chromatography was carried out on silica gel 60 H. The solvent system used was CHCl<sub>3</sub>-MeOH-NH<sub>3</sub> (85:14:1). Tlc was carried out on silica gel 60 HF<sub>254</sub> with the solvent system, CHCl<sub>3</sub>-MeOH-NH<sub>3</sub> (85:14:1).

PLANT MATERIAL.—S. argentino was collected in September 1984 by M.J. Pestchanker near San Luis City, Argentina, and identified by L.A. Del Vitto (MERL No. 848).

Species	Isolated Compound	Percentage <sup>a</sup>
S. argentino (roots)	senecionine	0.0010
	uspallatine	0.10
S. filaginoides ((roots)	senecionine	0.0435
	retrorsine	0.0643
S. seratophiloides (roots)	senecionine	0.035
	senecivernine	0.0070
	usaramine	0.0036
	retrorsine	0.037
	uspallatine	0.0185
S. leucostachys ((roots)	senecionine	0.0038
	integerrimine	0.0024
	uspallatine	0.0432
S. ragonesei (roots)	senecionine	0.021
	integerrimine	0.018
	retrorsine	0.019
S. subulatus (1)	1,2-dihydroretrorsine	0.0018
	senecionine	0.0030
	retrorsine	0.0376
S. glandulosus (1)	integerrimine	0.0191
	retrorsine	0.0352
	usaramine	0.0325
S. uspallatensis (2)	uspallatine	0.0078
	retrorsine	0.0090
S. gilliesiano (3)	retrorsine	0.0306
	senecionine	0.0206

TABLE 1. Pyrrolizidine Alkaloids Isolated from Nine Senecio Species

<sup>a</sup>Based on dry weight of the plant material.

## **Brief Reports**

S. filaginoides was collected in June 1984 by L.A. Del Vitto in La Crucecita, Mendoza, Argentina and identified by L.A. Del Vitto (MERL No. 897).

S. seratophiloides was collected in November 1984 by M.J. Pestchanker in El Volcan, San Luis, Argentina and identified by D.L. Anderson (INTA, Villa Mercedes, San Luis No. 2379-B).

*S. leucostachys* was collected in November 1984 by L.A. Del Vitto in Lujan Mendoza, Argentina, and identified by L.A. Del Vitto (MERL No. 592).

S. ragonesei was collected in November 1984 near Cacheuta, Mendoza, Argentina, by L.A. Del Vitto and identified by L.A. Del Vitto (MERL No. 918).

EXTRACTION AND ISOLATION.—Air-dried roots of all five species were extracted three times with hot MeOH. The resulting extract was concentrated in vacuo and the residue taken up in 30% citric acid solution and extracted several times with *n*-hexane and CHCl<sub>3</sub>. The acidic extract was made alkaline to pH 10.5 with NH<sub>4</sub>OH and extracted several times with CHCl<sub>3</sub>. After drying and evaporation, the combined CHCl<sub>3</sub> extracts gave a viscous mass containing the total alkaloids. The aqueous alkaline solution was carried to pH 2 with HCl, zinc dust was added, and the mixture was stirred for 24 h to reduce *N*-oxides that might be present. The extraction procedure of alkaloids thus obtained was carried out as described above.

The isolated alkaloids were identified by comparison with authentic samples and by comparison with previously reported data (4-8).

# ACKNOWLEDGMENTS

The authors wish to thank Ing. D.L. Anderson, INTA, Villa Mercedes, San Luis, and Ing. L.A. Del Vitto, IADIZA, Medoza, for their collaboration in the collection of plant species and their botanical identification. The authors are also indebted to Dr. M. Gonsalez Siera, IQUIOS, Rosario, for the <sup>13</sup>C-nmr spectral determinations and Lic. F.H. Guidugli for much help in measuring hr ms. This work was supported by a grant from CONICET and SUBCYT. M.J.P. wishes to thank CONICET for a fellowship.

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Received 11 December 1985