

Alkaloids of *Siparuna griseo-flavescens*

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and identification procedures, including spectroscopic data

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Literature describing the indigenous medicine of Central and northern South America contains numerous references to various plant species of the genus *Siparuna* (Monimiaceae) that have been used in the therapy of snakebite, fever, headache, rheumatism, and various infections (1). To our knowledge, the literature is devoid of references to the constituents of *Siparuna griseo-flavescens* Perk. However, the following species are sources of various benzylisoquinoline-derived alkaloids as noted (2, 3): *S. gilgiana* (liriodenine, oxonantenine), *S. dresslerana* (flavinantine, *O*-methylflavinantine), *S. guianensis* (cassamedine, liriodenine), *S. nicaraguensis* (liriodenine), *S. pauciflora* (nantenine, noroliveroline, *N*-methyllaurotetanine), and *S. tonduziana* (laurotetanine, *N*-methyllaurotetanine, norantenine, nantenine, anonaine, asimilobine, liriodenine, oxoantenine, reticuline).

Siparuna griseo-flavescens Perk. was collected on December 5, 1990 at Reserva Forestal de San Ramon, Alajuela, and identified by (voucher deposited) Dr. Jorge Gomez Laurito, Herbarium of the National Museum, San Jose. The air-dried, ground stems (1.2 kg) were extracted with EtOH and the resulting extract fractionated via accepted methods (2) to afford a basic fraction. Repeated gradient column chromatography (CHCl₃-MeOH mixtures) (silica gel) and preparative TLC (C₆H₆-Me₂CO-NH₄OH mixtures) (silica gel) afforded the aporphine alkaloids (+)-isocorydine (2 mg) (3), (+)-nantenine (10 mg) (4), (+)-*N*-methyllaurotetanine (5 mg) (4), and (+)-asimilobine (45 mg) (4). The alkaloids were characterized by direct comparison (UV, IR, ¹H-NMR, EIMS, m.p., optical activity) with authentic samples and/or published data (3, 4).

References

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- Lopez, J. A., Laurito, J. G., Lin, F.-T., Duah, F. K., Sharaf, M., Aly, Y., Wong, L. K., Schiff, P. L., Jr. (1990) *Planta Med.* 56, 492.

Erratum

Ammon, H. P. T., Anazodo, M. I., Safayhi, H., Dhawan, B. N., and Srimal, R. C. (1992) *Planta Med.* 58, 226.

The following text should be added: Curcumin inhibited the calcium/calcium ionophore-stimulated formation of LTB₄ with an EC₅₀ of 27 × 10⁻⁶ M.