

B149 Cyanolipids from *Paullinia cupana* var. *sorbilis* (Mart.) DuckeP. Avato^a, M.A. Pesante^a, F.P. Fanizzi^b and C.A.M. Santos^c^a Dipartimento Farmaco-Chimico, Università, Via Orabona 4, I-70125 Bari, Italy. ^b Dipartimento di Scienze e Tecnologie Biologiche ed Ambientali, Università, Via Monteroni, I-73100 Lecce, Italy. ^c Lab. de Farmacognosia, Jardim Botânico, 80.210-170 Curitiba, PR, Brasil.

Paullinia cupana var. *sorbilis* (Mart.) Ducke, commonly known as guaraná, is a plant native to the Amazonian forest, belonging to the Sapindaceae family (1). Seeds from this plant are known to contain high amounts of caffeine and are used to prepare a powder recommended as an energy reconstituent. Nevertheless, the seeds of many species of Sapindaceae are rich in oils that contain acylglycerols and an unusual class of plant lipids, the cyanolipids (2).

The chemical composition of the oil extracted from the seeds of *P. cupana* has been investigated with particular reference to the content of cyanolipids and data are reported in the present communication.

Cyanolipids amounted to 3% of the total oil from guaraná seeds. Generally, four types of cyanolipid structures, with fatty acids esterified to a mono- or di-hydroxy-nitrile moiety, have been reported as occurring in plants. ¹H and ¹³C NMR analyses indicated that cyanolipids of the type I (1-cyano-2-hydroxymethylprop-2-ene-1-ol diesters) are present in the oil extract from *P. cupana*. Moreover, the GC analysis of the 4,4-dimethylxazoline derivatives from those metabolites showed that *cis*-13-eicosenoic acid (paullinic acid) was the main fatty acid (38%) esterified to the nitrile group. Vaccenic acid (21%) and *cis*-15-eicosenoic acid (16%) were other abundant constituents. Identification of these fatty acids as the major components of the cyanolipid fraction from the guaraná seeds was also confirmed by GC/MS. To the best of our knowledge, only one paper has been previously published on the occurrence of cyanolipids in guaraná seed oils (3). Our data contribute to improve earlier findings.

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B150 Scale-up of isolation of oxindole alkaloids from *Uncaria tomentosa* by HPLC using chromatographic modelJ.L. Mazzei^a, S.L. Rosario^b, R. de Souza e Silva^c, A.C. Siani^b, L.M.M. Valente^c and L.A. d'Ávila^a^a Depto. Processos Orgânicos, Escola de Química, Universidade Federal do Rio de Janeiro, E-204 Centro de Tecnologia, 21949-900 Rio de Janeiro, Brazil. ^b Far-Manguinhos, Fundação Oswaldo Cruz, Sizenando Nabuco 100, Manguinhos, 21041-250 Rio de Janeiro, Brazil. ^c Depto. Química Orgânica, Instituto de Química, Universidade Federal do Rio de Janeiro, 620-A Centro de Tecnologia, 21949-900 Rio de Janeiro, Brazil.

The species *Uncaria tomentosa* (Rubiaceae), known as Cat's Claw, is a large woody vine indigenous to the Amazon rainforest. In herbal medicine, it is employed mainly for the treatment of immunological diseases and inflammations. Pentacyclic oxindole alkaloids present in the species have been considered biochemical markers and essential to standardize the commercial herbal medicines.

Semipreparative and preparative HPLC have been used to produce high purity compounds from natural sources. In this work we have applied chromatographic models for scale-up prediction aimed to optimize the isolation of the alkaloids isopteropodine, pteropodine, uncarine F, myrtraphylline, isomyrtraphylline and speciophylline found in *U. tomentosa*.

From the ethanol extract of the stalk bark of *U. tomentosa* an alkaloid-rich fraction was obtained through a classic acid-base partition. The parameters related to retention and separation efficiency of the oxindole alkaloids in analytical reverse-phase HPLC were determined varying the stationary phase, modifier content and temperature. The runs were performed in LiChrospher RP-18, LiChrosorb RP-18 and Shimpack MRC-ODS columns using acetonitrile-water 54:46, 46:54 and 38:62, at 30, 50 and 80 °C. The effects of the cited variations on the parameters were significant.

A model based on statistical moment analysis was used as a tool to simulate chromatograms of the studied alkaloids. Uncarine F, myrtraphylline and isopteropodine were separated using a Shimpack column, under different optimal conditions, 38:62 (30 °C), 38:62 (50 °C), and 50:50 acetonitrile-water (30 °C), respectively. The predicted and experimental separations were similar revealing the applicability of the methodology.

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