LITERATURE CITED

CORRÊA, D. DE B.; FONSECA E SILVA, L.G.; GOTTLIEB, O.R. & GONÇALVES, S.J.

1970 — Quinone and xanthone constituents of Kielmeyera rupestris. Phytochemistry, 9:447-541.

CRONQUIST, A.

1968 — The Evolution and Classification of Flowering Plants, p. 196, Nelson & Sons Ltda., London.

Drake, N.L. & Campbell, W.P. 1936 — Cerin and friedelin. III. Study of the oxidative degradation of friedelin. J. Amer. Chem. Soc., 58:1681-1684.

HEGNAUER, R.

1968 — Chemotaxonomie der Pflanzen, v. 5, p. 416, Birkhäuser Verlag, Basel.

LAVIE, D. & KAYE, I.A.

1963 — Isolation of β -sitostenone from Quassia amara, J. Chem. Soc., p. 5001-5002.

WEINGES, K.

1961 — Über einige neue Lignane und stereochemische Zusammenhänge in der Lignangruppe. Ber., 94:2522-2533.

The Chemistry of Brazilian Lauraceae. XLVII. Ferulic esters from Endlicheria and Ocotea species

Aura M. P. de Diaz (1), Pedro P. Dias D. (1), Zenaide S. Ferreira (2), Otto R. Gottlieb (2), Roberto A. de Lima (3) e Sérgio de H. Cavalcante (3)

A specimen from the Torquato-Tapajós Highway, km 133, Amazonas, voucher Herbarium INPA 48255, was identified tentatively with Endlicheria anomala Nees (Lauraceae) by Dr. W. A. Rodrigues. A trunk wood sample (1.2 kg) was extracted with ethanol. The chloroform-benzene 1:1 soluble part (15 g) of the extract (38 g) was chromatographed on a silica (360 g) column. Elution with C6H6 gave. in order, fatty ester (2 g, oil), n-tetracosyl ferulate [347 mg, mp 65-67° (C6H6)], n-tetracosanol [93 mg, 71-73° (C_6H_6)] and sitosterol [1230 mg, 138-140° (EtOH)]. Elution with $C_6H_6\text{-}CHCI_3$ 9:I, $C_6H_6\text{-}CHCI_3$ 8:2 to 0:10, $CHC!_3\text{-}$ -AcOEt 9:1 and AcOEt-MeOH 9:1 gave respectively stearic acid [70 mg, mp 68-69° (C₆H₆-EtOH)], a stearate of a fatty alcohol [58 mg, mp 86-88° (EtOH-CHCl3)], a ferulate of a fatty alcohol [90 mg, mp 77-78° (EtOH-CHCI₃)] and a glycoside [28 mg, mp 291-293° (EtOH-CHCI₃)].

n-Tetracosyl ferulate was identified by spectral data (IR, 'H NMR, MS) and direct comparison with an authentic sample [Franca et al., 1975]. n-Tetracosanol was identified by direct comparison with the alcohol obtained by saponification of the ferulate. Sitosterol and stearic acid were identified by direct

comparison with authentic samples. Saponification of the additional esters, mp 86-88° and 77-78°, gave respectively stearic acid, mp. 69-70°, and ferulic acid, 169-170°.

The simplicity of composition of *E. anomala* contrasts with the diversity and complexity of cinnamate derived metabolites of Lauraceae species described in all previous reports of the present series [for part XLVI see Diaz *et al.*, 1977]. In this respect, the species, nevertheless, does not occupy a peculiar position. Absence of detectable quantities of cinnamate derived metabolites and accumulation of ferulic acid was noted additionally for the following species.

Ocotea canaliculata Mez from the Manaus-Itacoatiara Highway, km 140, Amazonas, cf. voucher Herbarium INPA 16896, identified by Dr. W. A. Rodrigues.

- O. guianensis Aubl. from the Manaus-Ponta Negra Road, Amazonas, voucher Herbarium INPA 50121, identified by B. Albuquerque.
- O. neesiana (Miq.) Kosterm. from the Manaus-Itacoara Highway, km 69, voucher Herbarium INPA, Manaus 47243, identified by Dr. W. A. Rodrigues

^{(1) —} Instituto Nacional de Pesquisas da Amazônia, Manaus.

 ^{(2) —} Instituto de Química, Universidade de São Paulo, São Paulo.
 (3) — Departamento de Química, Universidade Federal de Alagoas, Maceió.

O. opifera Mart. from the vicinity of Manaus, Amazonas, cf. voucher Herbarium INPA 9210, identified by M. Freitas.

O. sp. from the Ducke Forest Reserve, near Manaus, voucher Herbarium INPA 42208.

It must be concluded that all these species lack the enzymes necessary to channel cinnamates into more complex end products. The ferulic acid, which is thus accumulated, inhibits L-phenylalanine ammonia-lyase activity [Havir and Hanson, 1968] and, as a consequence, the formation of benzylisoquincline alkaloids should be favoured. Indeed, the above mentioned unclassified *Ocotea* species contains two papaverine type alkaloids [Franca et al., 1975)]. This mechanism may explain the fairly general substitutive presence of benzylisoquinolines vs. arylpropanoids in the Magnoliidae [Gottlieb, 1972], i.e. the phe-

nomenon by which a species contains predominantly one of these classes of metabolites, producing, if at all, only traces of the other

LITERATURE CITED

HAVIR, E.A. & HANSON, K.R.

1968 — L-Phenylalanine ammonia-lyase. II.

Mechanism and kinetic properties of
the enzyme from potato tubers. Biochemistry, 7(5):1904-1914.

Franca, N.C.; GIESBRECHT, A.M.; GOTTLIEB, O.R.; MAGALHÃES, A.F.; MAGALHÃES, E.G. & MAIA, J.G.S.

1975 — Benzylisoquinolines from Ocotea species. Phytochemistry, 14:1671-1672.

DIAZ, A.M.P. DE; GOTTLIEB, O.R.; MAGALHÃES, A.F.; MAGALHÃES, E.G. & MAIA, J.G.S.

1977 — Notes on Aniba species. Acta Amazonica, 7(1): 41-43.

GOTTLIEB, O.R.

1972 — Chemosystematics of the Lauraceae.

Phytochemistry, 11: 1537-1570.

On the Vectors of Cutaneous Leishmaniasis in the Central Amazon of Brazil. I. Preliminary Findings (*)

Jorge R. Arias Rui A. de Freitas Instituto Nacional de Pesquisas da Amazônia, Manaus

It is of prime importance in the study of the epidemiology of vector borne diseases to identify the vector species. This problem often becomes complicated by the fact that the vector species is a part of more intricate "species-complexes" composed of several different species and subspecies. The problem of "species complexes" has been the cause of considerable confusion in such groups as the Anopheles gambiae "complex" (malaria) and the Simulium damnosum "complex" (filariasis) in Africa, and the Simulium amazonicum "complex" (filariasis) situation in Brazil. The "species complex" concept has not been widely used for many groups of New World sandflies, but we are becoming so involved with such subtle morphological differences in sandflies that we are forced to consider if we too are not faced with the "species complex" problem in our leishmaniasis research.

The search for the vectors of cutaneous leishmaniasis in Latin America has undergone much progress in the last decade, particularly in South America. Wijers & Linger (1966) found that Lutzomyia anduzei (Rozeboom) (1) was a possible vector of Leishmania braziliensis in Surinam, and Lainson & Shaw (1968) showed that Lu. llaviscutellata is the vector of Leishmania mexicana amazonensis in silvatic rodents and marsupials in the lower Amazon. In southern Brazil, Forattini et al. (1972) incriminated Lu. intermedia and Lu. pessoai as the vectors of Le. braziliensis. Lainson et al. (1973) incriminated Lu. wellcomei as the vector of Le. braziliensis braziliensis in the Serra dos Carajás, Pará State, also in the Amazon basin. Lainson et al. (1976) also showed that Lu. anduzei (Floch & Abonnenc) was the major vector of leishmaniasis in the Jarí River area of the State of Pará.

^{(*) —} This research was partly sponsored by the CNPq grant no SIP/08-131 and by INPA Project no 2017/103.
(1) — For the difference between **Lu. anduzei** (Rozeboom) and **Lu. anduzei** (Floch & Abonnenc) see Lainson **et al.** (1976).