Phytochemical and Pharmacological Investigations of Virola oleifera Leaves

Kátia N. Kuroshima^{a,c}, Fátima de Campos^b, Márcia M. de Souza^b, Rosendo A. Yunes^c, Franco Delle Monache^d and Valdir Cechinel Filho^b*

- ^a Centro de Ciências Tecnológicas da Terra e do Mar CTTMar, Universidade do Vale do
- Itajaí, UNIVALI

 b Núcleo de Investigações Químico-Farmacêuticas NIQFAR/CCS, Universidade do Vale do Itajaí (UNIVALI), Rua Uruguai, 458. Cx.P. 360, CEP 88302–202, Itajaí-SC-Brasil. Fax: 55 021 47 341 7664.
- E-mail: cechinel@mbox1.univali.br
- Departamento de Química, Universidade Federal de Santa Catarina (UFSC), Florianópolis-SC Brasil
- d Centro Chimica Recettori, C.N.R., Rome, Italy
- * Author for correspondence and reprint requests

Z.Naturforsch. **56c**, 703–706 (2001); received October 17, 2000/May 28, 2001

Virola oleifera, Analgesic Activity, Oleiferin-C

A methanolic extract and two fractions (n-hexane and ethyl acetate) from *Virola oleifera* leaves and some compounds (one lignan and two flavonoids) were investigated to verify the analgesic activity by using the writhing test in mice. The crude methanolic extract showed a moderate analgesic effect (about 40% of inhibition in this test at 10 mg/kg), whereas n-hexane and ethyl acetate fractions caused inhibition of $51.3 \pm 5.9\%$ and $50.5 \pm 6.3\%$, respectively. Oleiferin-C (1), a lignan isolated from the n-hexane fraction, showed an interesting analgesic potential in this model when compared to two standard drugs, paracetamol (4-acetamidophenol) and aspirin (acetylsalicylic acid). The ID₅₀ calculated for this compound was $17.25~\mu$ mol/kg, with confidence interval between 13.7~and $21.3~\mu$ mol/kg, being about 8 times more potent than the standard drugs. The mixture of two glycoside-flavonoids, identified as astilbin (2) and quercitrin (3), also exhibited good analgesic activity, causing 63% of reduction of abdominal constriction in mice. These results suggest beneficial effect of this plant to treat dolorous processes.