

Chemical constituents of *Xylopia excellens* (ANNONACEAE).

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Abstract

In the present study we report the isolation of two new ent-kaurene diterpene glycosides (1-2) and eleven isoquinoline alkaloids and their derivatives, from *Xylopia excellens* (Annonaceae): one benzyltetrahydroisoquinoline (3), two aporphines (4-5), eight oxoaporphines (6-13).

Introduction

Xylopia excellens (Annonaceae) is an amazon plant commonly known as “envireira”. Previous phytochemical and pharmacological investigation on *Xylopia* species revealed the presence of several bioactive compounds^{1,2}. Phytochemical study of the leaves and stems of *Xylopia excellens* afforded two new ent-kaurene diterpene glycosides named ent-kaur-16-en-7 β -D-glucose (1) and ent-kaur-16-en-7 β -D-galactose (2) and eleven alkaloids were isolated, and characterized as reticuline(3) (benzyltetrahydroisoquinoline), anonaine (4) and normantenine (5) (aporphines), liriodenine (6), lysicamine (7), isomochastoline (8), oxoglucine (9), O-methylmoschatoline (10), lanuginosine (11), oxonantenine (12) and oxophoebine (13) (oxoaporphines). All structures were identified through 1D and 2D NMR techniques along with mass spectrometry and by comparison with literature.

of the leaves (100 mg) was subjected to purification by HPLC-DAD in semi-preparative scale, using a C18 semi-preparative column eluted with methanol / water with 5% (v / v) trifluoroacetic acid in the proportion of 80:20, flow 5 mL / min and detection at UV₁=280nm and UV₂ = 305 nm affording three alkaloids (3,4 and 9). The alkaloidal fraction of the stems (100 mg) was subjected to purification by HPLC-DAD in semi-preparative scale, the conditions were the same described previously, yielding six fractions. These subfractions were subjected to a new purification by HPLC-DAD, using a phenyl-hexyl semiprep column eluting with methanol/water with 5% (v/v) trifluoroacetic acid in the proportion of 60:40 flow of 5 ml/min and the wavelengths used were the same as described, yielding (5, 6, 7, 8, 10, 11, 12 and 13). Compounds 1-2 were reported for the first time in the literature. Compounds 3-13 were reported for the first time in this species. These class of compounds are very common in species of the genus *Xylopia*.⁴

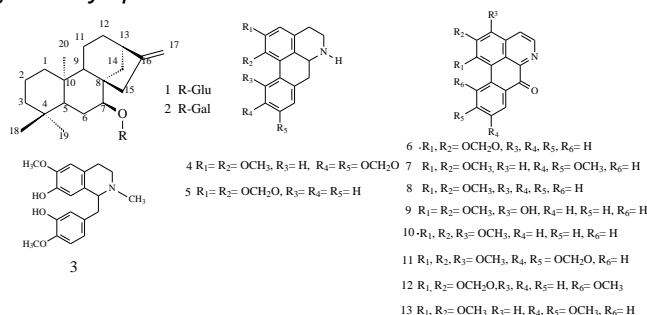


Figura 1. Isolated chemical constituents of *Xylopia excellens*

Resultados e Discussão

The powdered air-dried leaves and stems of *X. excellens* was extracted successively with hexane and MeOH. The hexane extract (5 g) was partitioned with hexane/10% aqueous metanol (1:1), yielding the hydroalcoholic fraction (1.7g). This fraction was supported over silica gel and eluted initially with n-hexane, followed by a gradient of EtOAc and methanol, yielding 1 and 2. The MeOH extract of the leaves and stems were redissolved in CHCl₃ and subjected to extraction with 3% aqueous HCl. This aqueous solution was adjusted with NH₄OH_{conc.} to pH 10, and extracted with CHCl₃ to yield CHCl₃ alkaloid fraction³. The alkaloidal fraction

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Conclusões

This is the first report of the chemical composition of *Xylopia excellens*. The results of this study contribute to the chemotaxonomic knowledge of the family and stimulate the continuation of investigations of this species for the identification of bioactive compounds.

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- ¹Quintans JSS, Soares BM, Ferraz RPC, Oliveira ACA, Silva TB, Menezes LRA, Sampaio MFC, Prata APN, Moraes M, Pessoa C, Antonioli A, Costa E, Bezerra D. *Planta Medica* 2013, 79: 123-130.
- ²Nishiyama Y, Moriyasu M, Ichimaru M, Iwasa K, Kato A, Mathenge SG, Chalo-Mutiso PB, Juma FD. *Phytochemistry* 2006, 67: 2671-2675.
- ³Chang, F. R., Wei, J.L., Teng, C. M., Wu, Y.C., *Phytochemistry*. 1998, 49, 2015.
- ⁴Moreira IC, Roque NF, Vilegas W, Zalewski CA, Lago JHG, Funasaki M. *Chemistry & Biodiversity* 2013, 10: 1921-1943.