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INDOLE ALKALOIDS AND QUASSIN FROM QUASSIA AFRICANA

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In our continuing phytochemical investigation of biologically active constituents from Simaroubaceae spp., we report here the occurrence in root barks of *Quassia africana* Baill. (syn. *Simaba africana* Baill.) of three alkaloids (canthin-6-one, 4,5-dimethoxycanthin-6-one, β -carboline-1-propionic acid) and quassin, an additional quassinoid different from those previously isolated from this plant (1). Their structures were established by spectroscopy and direct comparison with authentic samples.

EXPERIMENTAL

PLANT MATERIAL.—Root samples of *Q. africana* were collected in northeastern Zaïre (October 1983) and in the south Congo (February 1984). A voucher specimen has been deposited at the Botanical Institute of the Free University of Brussels.

ISOLATION OF CANTHINONE ALKALOIDS.—The crude chloroformic extract obtained after alkalinization of the dried powder was extracted with 0.1 N HCl. The extraction of the acid aqueous phase with CHCl₃ led to isolation of canthin-6-one (70 mg, 0.07%). 4,5-Dimethoxycanthin-6-one (40 mg, 0.04%) was further isolated by extraction with CHCl₃ after adjustment to pH 10 of the aqueous phase. Purification of both alkaloids was achieved by preparative tlc on Si gel with toluene-Me₂CO-EtOH-NH₄OH (50:30:4:1) (2).

Isolation of β -carboline-1-propionic acid (10 mg, 0.02%).—The dried powder was extracted with MeOH acidified with HOAc. The methanolic extract was concentrated after dilution with H₂O; the pH value was adjusted to 5 and the alkaloid was precipitated by addition of picric acid. The picrate was dissolved in Me₂CO-H₂O (1:1), chromatographed through an Amberlite IRA 400(Cl⁻), and purified by preparative tlc on Si gel with toluene-EtOH-EtOAc-NH₄OH (4:4:2:1) (3).

Extraction of quassin (25 mg, 0.05%).—The dried powder was extracted with MeOH and the residue was fractionated on a short Si gel column with CHCl₃ containing increasing amounts of MeOH. The fraction which contained quassin was further purified by preparative tlc on Si gel using CHCl₃-MeOH (7:3) (4).

Full details of the isolation and identification of the reported compounds are available upon request.

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