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MAJOR FLAVONOIDS OF TEPHROSIA NUBICA

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Tephrosia Pres. (Leguminosae-Papilionoideae) is a large tropical and subtropical genus estimated to contain about three hundred species (1). Tephrosia has been used medicinally and as a fish poison (2). Chemical studies on a number of species have revealed the presence of rotenoids (3,4) and a range of isoflavones (5-8), flavanones/chalcones (9-13), flavonols (14-16), and flavones; prominent among the flavones is a group of 5,7-oxygenated (17-19) and 7-oxygenated (20-24) compounds characterized by the occurrence of a C-8 prenyl unit and prenylated flavan (25).

We undertook the chemical investigation of Tephrosia nubica (Boiss) Baker.

EXPERIMENTAL

PLANT MATERIAL.—*T. nubica* was collected from Gabal Elba at the boundry between Egypt and Sudan and authenticated by Dr. Lofty Boulos, Professor of Taxonomy at the National Research, Cairo, Egypt.

EXTRACTION AND ISOLATION OF THE FLAVONOIDS.—Air-dried and powdered herb material of T. nubica (150 g) was defatted in a continuous extraction apparatus with petroleum ether. The defatted powder was then exhaustively extracted with MeOH in a Soxhlet apparatus. The alcoholic extract was concentrated under reduced pressure, and the resulting gum was extracted successively with CHCl₃, EtOAc, and n-BuOH. The CHCl₃ residue (1.8 g) was subjected to flash column chromatography (54 g of silica gel), eluting with C_6H_6 , C_6H_6 -MeOH (99:1), (98:2), (97:3), (96:4), and (95:5). Fractions (50 ml) were collected by utilizing the distinctive fluorescence of the components as shown by tlc [CHCl₃-MeOH (19:1) and C_6H_6 -EtOAc (8:3) as solvent systems]. The least polar compounds obtained from the above column were further purified by silica gel ptlc using the Chromatotron with CHCl₃-MeOH (99:1) and (98:2) as solvent systems. Complete purification of the compounds was achieved by semipreparative hplc (10×250 mm silica gel column, 5μ Supelco, hexane-EtOH, 4:1) which gave 10 mg of semiglabrin (21), 5 mg of pseudosemiglabrin (23), 30 mg of apollinine (23), and 50 mg of laneolatin (20). The structures of all compounds were determined by spectral analysis (ir, ms, and 1 H nmr) as well as comparison with published data.

Full details of the isolation and identification of the compounds are available on request from BBJ.

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