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n-1-ALKANOLS OF *HYPERICUM PERFORATUM*ILIA BRONDZ,¹* TYGE GREIBROKK,

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Several species of the genus *Hypericum* have been shown to produce antibacterial constituents (1-5), and in the case of *Hypericum perforatum* L. (St. John's wort), extracts have been utilized clinically in Russia to treat infections (6) and in the United States as a food preservative (3). Recently, we reported on the stereochemistry (7) of the antibiotic hyperforin, which is present in *H. perforatum* (6) and on the *n*-alkanes occurring in this plant (8). As part of our study on the constituents of *H. perforatum*, which is commercially available as dried plant material from Scandinavian drugstores, we have examined the saturated long-chain alcohols present in an acetone extract of this plant.

Previously, dodecanol (9), 1-tetracosanol (10), 1-hexacosanol (10-11), and 1-octacosanol (10) have been identified as constituents of *H. perforatum*. By gc-ms and co-chromatography with authentic *n*-alkanols, the present study showed that, based on the acetone extract, the content of long-chain alkanols was 4.3 g in 1 kg of dried plant material. The mixture of alkanols consisted of 1-tetracosanol (9.7%), 1-hexacosanol (27.4%), 1-octacosanol (39.4%), and 1-triacontanol (23.4%). The amount of 1-triacontanol is noteworthy because this constituent had not previously been identified in *H. perforatum*. The discrepancy between the former and the present investigations could possibly be explained by the development in gc-columns. The high-temperature fused silica column used in the present work would be expected to be better suited for compounds of low volatility.

EXPERIMENTAL

PLANT MATERIAL.—Dried leaf material of *H. perforatum* (*Herba hyperici*) was purchased from Norsk Medisinaldepot, Oslo. A voucher specimen is deposited at the Department of Pharmacy, University of Oslo.

EXTRACTION AND IDENTIFICATION.—Dried, powdered plant material (1 kg) was extracted with acetone (3 liters) in a Soxhlet apparatus for 3 h. The acetone extract was stored at -10° for 24 h. The precipitate that appeared was collected by filtration and was washed with methanol until coloured material could no longer be removed. The remainder of the precipitate was fractioned on a silica gel column (120 g) and, on elution with benzene, a fraction (4.3 g) corresponding to saturated alkanols was obtained. This fraction was examined using gas chromatography with a Chrompack CP-Sil 5 column (25 m, id 0.22 mm) and gas chromatography in combination with mass spectroscopy. The identity of the *n*-1-alkanols was confirmed by observing no separation on co-chromatography with authentic *n*-1-alkanols.

Full details of the isolation and identification are available on request to the senior author.

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Several species of the genus *Hypochaeris* have been shown to produce antihypertensive constituents (1-5), and in the case of *H. perforata* L. (Sc. John's wort), extracts have been utilized clinically in Russia on heart infections (6) and in the United States as a food preservative (7). Recently, we reported on the stereochemistry (7) of the antihypertensive hypochlorogenic acid (8) and on the α -alkenes occurring in this plant (8). As part of our study on the constituents of *H. perforata*, which is common-ly available as dried plant material from Scandinavian drugstores, we have examined the saturated long-chain alcohols present in an acetone extract of this plant. Previously, dodecanol (9), 1-tetradecanol (10), 1-hexacosanol (10-11), and 1-octacosanol (10) have been identified as constituents of *H. perforata*. By gc-ms and co-chromatography with authentic α -alkenes, the present study showed that, based on the acetone extract, the content of long-chain alcohols was 4.3 g in 1 kg of dried plant material. The mixture of alcohols consisted of 1-tetradecanol (9.7%), 1-hexacosanol (37.4%), 1-octacosanol (39.4%), and 1-triacontanol (13.5%). The amount of 1-triacontanol is noteworthy because this constituent had not previously been identified in *H. perforata*. The discrepancy between the former and the present investigation could possibly be explained by the development in gc-column. The high-boiling long-chain alcohols used in the present work would be expected to be better suited for compounds of low volatility.

EXPERIMENTAL

PLANT MATERIAL.—Dried leaf material of *H. perforata* (L.) was purchased from Norsk Medicinalpflanze, Oslo. A voucher specimen is deposited at the Department of Pharmacy, University of Oslo.

EXTRACTION AND IDENTIFICATION.—Dried, powdered plant material (1 kg) was extracted with acetone (7 liters) in a Soxhlet apparatus for 3 h. The acetone extract was concentrated to 10% on a rotary evaporator and was washed with methanol until acetone-soluble material could no longer be removed. The residue of the precipitate was extracted on a silica gel column (150 g) and, on elution with benzene, a fraction (4.3 g) corresponding to saturated alcohols was obtained. This fraction was examined using gas chromatography with a Chrompack CP-581 2 column (25 m, 0.032 mm) and gas chromatography in combination with mass spectrometry. The identity of the α -alkenes was confirmed by observing no separation on co-chromatography with authentic α -1-alkenes.

Full details of the isolation and identification are available on request to the senior author.

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