Octamethoxynaphthalene from Cașsia nodosa

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A hitherto unreported compound, octamethoxynaphthalene has been isolated from Cassia nodosa stems. It was been characterised from the spectral data.

ASSIA nodosa belongs to the family Leguminosae, subfamily Caeselpiniaceae. It is grown for its ornamental flowers and foliage¹. Roots are used as a soap for washing clothes². Ethanolic extract of the plant has antiinflammtory activity³. Leaves of the plant function as a purgative⁴. It has also been reported to possess insecticidal activity⁵. We report here the isolation and characterisation of a new compound from the stems of this plant.

The methanolic extract of the stems was subjected to silica gel column chromatography. Compound 1 was obtained on elution with hexane and it crystallised from hexane as colourless crystals. It was assigned the molecular formula C₁₈H₂₄O₈ on the basis of elemental analysis and the GCMS fragmentation pattern. The 1H NMR of the compound in CDCI3 exhibited no signal for the aromatic proton. A singlet at δ 3.89 integrating to 24 protons could be due to eight aromatic methoxyls. The data fits for a fully substituted naphthalene derivative and the compound could be octamethoxynaphthalene (1). The ¹³C NMR and DEPT-135 of the compound are consistent with the proposed structure. Aromatic carbons appeared as singlets at δ 150.00 and δ 127.60. The methoxyl carbons appeared at δ 60.84. The daughter ions in the MS lend further support to the proposed structure.

Stems were collected from the Landscape, CCS HAU, Hisar. ¹H NMR and ¹³C NMR were recorded at 300 MHz and 75 MHz respectively in CDCl₃; UV in CHCl₃; IR in KBr. The dried stems (2 Kg) were crushed and extracted 4 times with hot methanol. Extractives were concentrated under reduced pressure and the viscous material was mixed with

silica gel (60-120 mesh), dried on a water bath and subjected to silica gel column chromatography. Elutions afforded octamethoxynaphthalene (1, 100 mg), chrysophanol⁶ (25 mg), lupenone⁷ (20 mg), lupane-3 β, 11α , 20-triol⁸ (20 mg) and 1,2-dihydroxy-3-·methylanthraquinone9 (25 mg). The compound 1 has m.p. 166°; (Found C, 58.65; H, 6.50. C₁₈H₂₄O₈ requires : C, 58.69; H, 6.52%); IR v_{max} KBr cm⁻¹: 1580, 1460, 1200, 1160, 1005, 820, 710; UV λ_{max} CHCl₃ nm: 212.8, 228.8, 247.2, 256.0 and 296.8; ¹H NMR (300 MHz, CDCl₃, δ): 3.89 (24H, s, 8 x OMe); ¹³C NMR (75MHz, CDCl₃, δ): 150.5 (s, C-1, C-2, C-3, C-4, C-5, C-6, C-7 and C-8), 127.6 (s, C-4a and C-8a), 60.8 (OCH₃); GCMS m/z (rel. int.) 280 (6.89), 278 (32.48), 276(72.63), 274(42.72), 265(9.25), 263(44.88), 261(100.00), 246(0.10), 233(2.19), 222(0.44), 220(1.90), 216(3.25), 190(3.09).

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MS FRAGMENTATION PATTERN

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