

## NOTE

## A New Anthraquinone Glycoside from the Heartwood of *Cassia auriculata* Linn

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The present paper deals with the isolation and structural elucidation of a new anthraquinone glycoside, 3-hydroxy-6, 8-dimethoxy-2-methyl anthraquinone-1-O-β-D-galactoside from the heartwood of *Cassia auriculata* Linn.

In course of our studies on plant pigments of the genus *Cassia*<sup>1-3</sup>, systematic chemical investigation of heartwood of *Cassia auriculata* (Leguminosae) has been undertaken. The plant is highly reputed for its medicinal value and tanning material.<sup>4</sup> The present communication reports the isolation of a new glycoside on the basis of chemical and spectral evidences.

Air-dried and crushed heartwood (10 kg) of the plant (collected from Kauakol botanical garden, Nawada and identified in Botany Department, Magadh University, Bodh-Gaya) was successively extracted with hexane, CHCl<sub>3</sub> and alcohol. The concentrated alcoholic extract was treated with ice-cold water and fractionated into water-soluble and water-insoluble parts.

The ethyl acetate extract of aqueous alcoholic fraction after concentration under reduced pressure was subjected to column chromatography on silica gel and eluted with hexane → CHCl<sub>3</sub> → EtOAc → MeOH gradient as eluent. The EtOAc—MeOH (8 : 2 v/v) eluate gave a compound which was crystallised from EtOAc—pet. ether mixture as light brown amorphous solid (950 mg). It responded to characteristic colour reactions of anthraquinones<sup>5,6</sup> and glycoside.<sup>7</sup> The compound analysed for (M<sup>+</sup> 476) gave m.p. 235–37°C; (Found: C, 56.88; H, 4.76. Calcd. for C<sub>23</sub>H<sub>24</sub>O<sub>11</sub>: C, 57.98; H, 5.04%); λ<sub>max</sub>(MeOH) 255, 265, 288, 418 nm.; ν<sub>max</sub>(KBr) 3350 ν(OH), 2950 and 2885 ν(OMe), 1765, 1730, 1645, 1570, 1460, 1250, 1185, 1080, 960, 840, 810, 730 cm<sup>-1</sup>; <sup>1</sup>H NMR (DMSO-d<sub>6</sub>, 300 MHz) δ 2.38 (s, 3H, CH<sub>3</sub>), 3.25–3.70 (br, sugar protons), 3.89 (s, 6H, OCH<sub>3</sub>), 5.20 (d, 1H, J = 8 Hz, C—1' galactosyl), 6.86 (d, 1H, J = 8 Hz, C—7), 7.25 (s, 1H, C—4), 7.68 (d, 1H, J = 8 Hz, C—5) and a phenolic hydroxyl at 7.55 (s, 1H, D 20

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exchangable, C<sub>3</sub>—OH). Acid hydrolysis gave an aglycone, C<sub>17</sub>H<sub>14</sub>O<sub>6</sub>, m.p. 272–74°C, which was identified as 1,3-dihydroxy-6,8-dimethoxy-2-methyl anthraquinone by UV, NMR, mass spectra and direct comparison with an authentic sample<sup>8</sup> and galactose (CO—PC, osazone, 183°C) in the hydrolysate. The galactose was found to be linked at position-1 of the aglycone<sup>9</sup> which was supported by specific colour reaction.<sup>10</sup> The permethylated glycoside on hydrolysis yielded 2,3,4,6-tetra-O-methyl-D-galactose which was identical to the methyl galactose obtained by D-galactose. Thus, on the basis of the above observations, structure of the new glycoside was assigned as 3-hydroxy-6,8-dimethoxy-2-methyl anthraquinone-1-O-β-D-galactoside.

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