A CHALCONE GLYCOSIDE FROM THE SEEDS OF BAUHINIA PURPUREA

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Abstract—A new glycoside, 3,4-dihydroxychalcone 4-O- β -L-arabinopyranosyl-O- β -D-galactopyranoside has been identified from the seeds of *Bauhinia purpurea*.

Bauhinia purpurea L. (Leguminosae, subfamily Caesalpinioideae) is of some medicinal importance [1], especially the roots and bark in the treatment of diarrhoea and its decoction is useful as a wash for ulcers. We have previously reported one new chalcone glycoside from this plant [2].

In the present study we have found another new glycoside, $C_{26}H_{30}O_{12}$, mp 365°, which gave the characteristic colour reactions of a chalcone [3-5] and gave 3,4-dihydroxychalcone [6], galactose and arabinose on acid hydrolysis (8% ethanolic H_2SO_4 for 12 hr). The identity of sugars was confirmed by co-chromatography with authentic samples and by the preparation of their sazzones.

The bright yellow aglycone, C₁₅H₁₂O₃, mp 203-205° was identified as 3,4-dihydroxychalcone from its R_c values: 0.96 in n-BuOH-HOAc-H₂O (4:1:5) and 0.72 in EtOAc-conc HCl-H₂O (5:1:5) and by alkaline cleavage [7] to give the corresponding acetophenone; oxime, mp 57° (lit. 59°); phenylhydrazones, mp 103°; (lit. 105°) and protocatechuic acid, mp 190° (lit. 192°). These results indicate the presence of a 3,4-dihydroxy group in ring B which was confirmed by spectral studies. λ_{max} (MeOH) values were 264, 367 nm and a bathochromic shift of 78 nm on addition of NaOMe showed the presence of a free hydroxyl group at position 4. The aglycone also gave a bathochromic shift of 36 nm with NaOAc-H₃BO₃ indicating two free hydroxyl groups at positions 3 and 4, which were confirmed by a bathochromic shift (75 nm) of band I on the addition of anhydrous AlCl₃.

Periodate oxidation of the glycoside indicated that both sugar components of the disaccharide have the pyranose configuration, 4 mol of periodate being consumed and 2.1 mol of formic acid liberated. The sugar moiety is thought to be attached at position 4 because the aglycone gave a bathochromic shift of 78 nm while the glycoside gave no shift. On methylation of the glycoside followed by hydrolysis with Kiliani's reagent (HCl-HOAc-H₂O) (7.1:5.1), 2,3,6-tri-O-methyl-D-galactose and 2,3,5-tri-O-methyl-L-arabinose were identified in the aqueous hydrolysate. The identity of the

methylated sugars was confirmed by comparison of their RG values with 2,3,4,6-tetra-O-methyl-D-glucose (TMG). RG found for 2,3,6-tri-O-methyl-D-galactose was 0.70 (lit. 0.71) and for 2,3,5-tri-O-methyl-L-arabinose, 0.95 (lit. 0.96) in n-BuOH-EtOH-H₂O (5:1:4). This result indicates that C-1 of the galactose is linked with the aglycone at position 4 and that C-4 of the galactose is linked with arabinose at position 1. The methylated aglycone was identified as 4-hydroxy-3-O-methoxy-chalcone which indicates that the sugar moiety is attached at position 4 of the aglycone.

Arabinose was identified in the aqueous hydrolysate obtained from partial acid hydrolysis of the glycoside with formic acid in cyclohexanol [8], while galactose was not detected. This confirms the disaccharide as an arabinosyl galactoside. Complete enzymic hydrolysis of the glycoside with emulsin indicates that both sugars are β -linked. Thus from the above data the chalcone is identified as 3,4-dihydroxychalcone 4-O- β -L-arabinopyranosyl-O- β -D-galactopyranoside.

Seeds were collected from the Department of Chemistry, University of Allahabad, India and identified by the Botanical Survey of India, Allahabad.

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