JETIR.ORG



ISSN: 2349-5162 | ESTD Year : 2014 | Monthly Issue JOURNAL OF EMERGING TECHNOLOGIES AND INNOVATIVE RESEARCH (JETIR) An International Scholarly Open Access, Peer-reviewed, Refereed Journal

Chemical Study of Natural Product Obtained from Very Important medicinal plant Chrotolaria alata

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Abstract:-

A water-soluble and non-ionic D-galactomannan has been isolated from the seeds of Chrotolaria alata of Indian origin, containing D-galactose and D-Mannose in 8:2 molar ratio. Acid catalyzed fragmentation, periodate oxidation, methylation and enzyme hydrolysis showed that the mucilage has a branched structure consisting of a linear chain of β -D-(1 4) linked mannopyranosy units, some of which are substituted at (1 6) by α -D-galactopyranosyl anits, glycosidically. This galactomannan have similarity with ghee gaur plants gum.

Key words: Oligosaccharides, Mucilage,

Introduction:-

Polysaccharides are polymer obtained by stracts of seets of chroloraria alata sucking in Acoh overnight. A water soluble and non- ionic D-galactomann has been isolated from the seed of Chrotolaria alata of Indian origin¹. Polysaccharide was JETIR2208385 Journal of Emerging Technologies and Innovative Research (JETIR) www.jetir.org d781 conveniently extracted from the crushed, defatted and decolorized seeds by extracting with 1% aqueous acetic acid and by repeated precipitation² from its solution therein with ethanol. It was purified and tested for homogeneity by usual methods. The white amorphous polysaccharide had $[\alpha]_{D^{25}}$ + 68° (in water), an ash content³ of (0.8%) and a negligible percentage of methoxy, acetu1 and uronic acid contents.

Experiment and analysis:-

after complete acid-hydrolysis the polysaccharide yielded (Dgalactose and D-mannose I 2.7 molar ratio). Graded acid hydrolysis resulted in the preferential removal of α -linked D-galactose units⁴ on the periphery as end groups. To determine the position of linkages between the building units of the galalctomannan⁵, it was exhaustively methylated by Haworth-Purdie method^{6,7}, to afford a brown, semisolid glassy mass and had [α] $_{D}$ ²⁵ +41° (chloroform). Hydrolysis of the methylated seed-gum gave 2,3,46-tetra-0-methy1-D-galactose⁸ (4 mol), 2,3,6-tri-0-methyl-D-mannose⁹ (5 mol) and 2,3-di-0-methyl-D mannose¹⁰ (3 mol).

The identity of these methylated monosaccharides¹¹ was established on the basis of their R_{TMG} values, optical rotations and crystalline derivatives. The percentage of end groups calculated from methlation studies was 24.9%. Oxidation of the mucilage with sodium metaperiodate consumed 945 mM of the oxidant with the liberation of 283 mm of formic acid per 100g of the polysaccharide indicating 25.5% end-groups (*cf.* methylation).

Conclusion and Result:-

Acid catalyzed partial hydrolysis of the mucilage gave, two disaccharides: α -D-Galp (1 6)-D-Manp, β -D-Manp+(1 4)-D-Manp¹² and two disaccharides α -D-Galp (1 6)- β -D-Manp (1 4)-D-Manp¹³ and β -D-Manp (1 4)- β -D-Manp along with the component sugars. All of the oligosaccharides were characterized. These results corroborated in the earlier findings. The foregoing date accord with the following structure.1 6 attested galactose and mannose 1 4 attested mannose and mannose in main chan. $\begin{array}{c} \alpha \text{-D-Galp} & \alpha \text{-D-Galp} \\ 1 & 1 \\ \downarrow \\ 6 & 6 \end{array}$

 $(4 \rightarrow (\beta)_D \operatorname{Manp} - 1 \rightarrow 4(\beta)_D \operatorname{Manp} - 1 \rightarrow 4(\beta)_D \operatorname{Manp} - 1 \rightarrow 4(\beta)_D \operatorname{Manp} \rightarrow)n$

Paper chromatography¹⁴ was conducted on Whatmann filter paper no. 1 and 3 mm papers by descending technique using the following systems (v/v). A-1-butanol-ethanol-water (5:1:4), B-1-butanol-isopropanol-water¹⁵ (11:6:3), C-ethylecetate-pyridine-water (2:1:2).

Solution were concentrated at diminished pressure and at low temperature. All residues were dried in accuo over anhydrous $Cacl_2$ melting points are uncorrected and $[\alpha]_D$ values are for equilibria.

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