Carbohydr. Res. 1997, 304, 97

NMR spectroscopy study of the complexation of p-gluconic acid with tungsten(VI) and molybdenum(VI)

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Eight and ten complexes have been identified with molybdate and tungstate ions, respectively, in aqueous solution, depending on pH and concentration conditions; the structure of an important species formed both with tungstate and molybdate ions is shown.

R=CHOHCH2OH

Carbohydr. Res. 1997, 304, 111 An efficient chemical-enzymatic synthesis of 4-nitrophenyl β-xylobioside: a chromogenic substrate for xylanases

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Xylan HO HO HO HO OR

R = 4-nitrophenyl

Autocatalytic oxidation of primary hydroxyl functions in glucans with nitrogen oxides

Carbohydr. Res. 1997, 304, 117

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Carbohydr. Res. 1997, 304, 125

Some properties and action mode of (1(ar4-α-L-guluronan lyase from *Enterobacter cloacae* M-1

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E. cloacae $(1 \rightarrow 4)$ - α -L-guluronan lyase degraded hepta(GulA) to dimer and pentamer.

0-0-0-0-0-0-0 → 0-0, Δ-0-0-0-0

Carbohydr. Res. 1997, 304, 143

Cellobiohydrolase I from Trichoderma reesei:

Identification of an active-site nucleophile and additional information on sequence including the glycosylation pattern of the core protein

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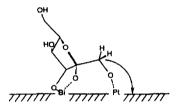
(R,S)-3,4-Epoxybutyl β -cellobioside inactivates cellobiohydrolase I from T. reesei by covalent modification of Glu^{212} , the putative active site nucleophile.

Carbohydr. Res. 1997, 304, 155

The oxidation of fructose on Pt/C catalysts. The formation of D-threo-hexo-2.5-diulose and the effect of additives

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Carbohydr. Res. 1997, 304, 165

Structural elucidation of the capsular polysaccharide from *Streptococcus pneumoniae* type 18B

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Streptococcus pneumoniae type 18B was shown to have a capsular polysaccharide with the following structure:

OPO₃H-CH₂-CHOH-CH₂OH $\begin{vmatrix}
1 & 3 \\
-34\end{pmatrix}$ \rightarrow 4)- β -D-Glcp-(1 \rightarrow 4)- β -D-Galp-(1 \rightarrow 4)- α -D-Glcp-(1 \rightarrow 3)- β -t-Rhap-(1 \rightarrow $\begin{vmatrix}
2 & 1 \\
1 & \alpha$ -D-Glcp

Carbohydr. Res. 1997, 304, 173

Enzymatic synthesis of the glycosides of calystegines B_1 and B_2 and their glycosidase inhibitory activities

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Calystegine $\mathbf{B}_1: \mathbf{R}^1 = \mathbf{H}$ 1: $\mathbf{R}^1 = \mathbf{B} \cdot \mathbf{G}$ -jucopyranosy

2: $\mathbf{R}^1 = \mathbf{G} \cdot \mathbf{G}$ -jucopyranosy

HO OH OH

Calystegine B₂ : R² = H

3 : R² = β-p-glucopyranosyl
 4 : R² = β-p-galactopyranosyl

5 : R² = α-D-galactopyranosyl

Carbohydr. Res. 1997, 304, 179

Synthesis of ginsenoside Rg₃, a minor constituent of Ginseng Radix

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Glycosylation of 12-acetoxy-protopanaxadiol, with α -acetobromosophorose under catalysis by $\mathrm{Ag_2CO_3}$ or $\mathrm{Ag_2O}$, followed by deacetylation, afforded a mixture of the α - and β -linked sophorosides. The last was identical in all respects with ginsenoside $\mathrm{Rg_3}$, the minor component of Ginseng Radix Rubra.

Carbohydr. Res. 1997, 304, 183

IR spectra and hydrogen bonding in tetritols

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The enhanced water absorption capability of D,L-threitol as compared to *meso*-erythritol is ascribed to differences in strength of H-bonds in the crystals based on IR spectral data.