GRAPHICAL ABSTRACTS

Carbohydr. Res. **1997**, *299*, 1

Enantiomer separation of permethylated monosaccharides and 1,5-anhydroalditols and simultaneous determination of linkage positions and absolute configuration in the galactan of *Helix pomatia*

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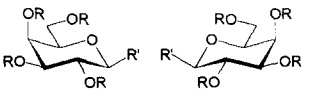
The enantiomers of permethylated monosaccharides and 1,5-anhydroalditols were resolved by enantioselective GLC using modified cyclodextrin as chiral stationary phase. The 1,5-anhydro D- and L-galactitols obtained from the permethylated galactan from *Helix pomatia* could be separated, allowing the simultaneous determination of linkage position and absolute configuration of galactose residues in snail galactan.



Carbohydr. Res. **1997**, *299*, 7

A convenient synthesis of disaccharides containing furanoside units

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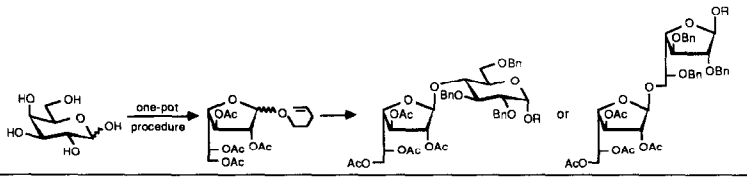
Laboratoire de Chimie Organique et des Substances Naturelles, associé au CNRS, Ecole Nationale Supérieure de Chimie de Rennes, Avenue du Général Leclerc, F-35700 Rennes, France



Carbohydr. Res. **1997**, *299*, 23

Synthesis of α-tocopheryl oligosaccharides

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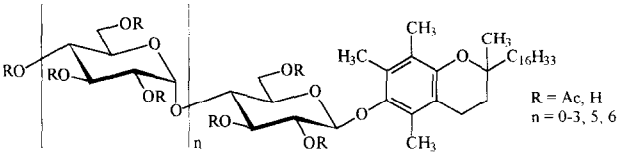
α-Tocopheryl alto-oligosaccharides were prepared as antioxidants water soluble.



Carbohydr. Res. **1997**, *299*, 33

Synthesis and characterisation of highly glycosylated glycopeptides with T^n-antigenic structures corresponding to human glycophorin A^N

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Two highly glycosylated glycopeptides corresponding to the human glycophorin A^N with T^n-antigenic structures were synthesised. The solid phase synthesis was realised by using carbohydrate containing building blocks. The synthesised substances were characterised by NMR spectroscopic techniques.

H-T-S-T-S-S-S-V-T-K H-Ṫ-Ṡ-Ṫ-Ṡ-Ṡ-Ṡ-V-T-K

L-Ṡ-Ṫ-Ṫ-E-V-A-M-H-Ṫ-Ṡ-Ṫ-Ṡ-Ṡ-Ṡ-V-T-K * α-D-GalNAc

Carbohydr. Res. **1997**, *299*, 49

Synthesis and surface-active properties of amphiphilic 6-aminocarbonyl derivatives of D-glucose

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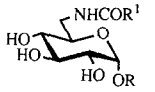
Several 6-amido-6-deoxy derivatives of methyl α-D-glucopyranoside and D-glucopyranose were prepared from 6-azido-6-deoxy intermediates. The Staudinger reaction (PPh_3, carboxylic acid or acid chloride) or the addition of *n*-heptanol on the 6-deoxy-6-isocyano derivatives, obtained from the 6-azido analogues, completed the synthesis. The Krafft temperatures and surfactant properties of these compounds are reported.



Carbohydr. Res. **1997**, *299*, 59

Synthesis of 4-cyanophenyl 2-deoxy-1,5-dithio-β-D-*threo*-pentopyranoside

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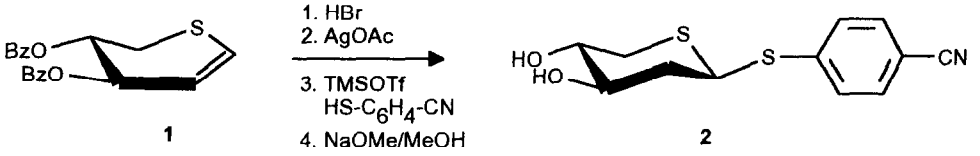
5-Thio-D-xylose was converted via its 1-ene derivative **1** into the title compound **2** possessing oral antithrombotic activity.



Carbohydr. Res. **1997**, *299*, 85

Chemoenzymatic synthesis of the Thomsen–Friedenreich antigen determinant

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