

#### Carbohydrate Research Vol. 339, No. 4, 2004

#### **Contents**

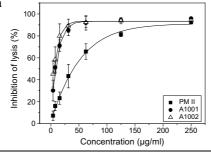
#### **FULL PAPERS**

Bioactive polysaccharides from the stems of the Thai medicinal plant *Acanthus ebracteatus*: their chemical and physical features

pp 753–762

Sanya Hokputsa,\* Stephen E. Harding, Kari Inngjerdingen, Kornelia Jumel, Terje E. Michaelsen, Thomas Heinze,

Andreas Koschella and Berit S. Paulsen



Structure of oligosaccharide side chains of an intestinal immune system modulating arabinogalactan isolated from rhizomes of *Atractylodes lancea* DC

pp 763-770

Ikue Taguchi, Hiroaki Kiyohara, Tsukasa Matsumoto and Haruki Yamada\*

$$\rightarrow$$
3)-β-D-Gal-(1 $\rightarrow$ 3)-β-D-Gal-(1 $\rightarrow$ 3)-β-D-Gal-(1 $\rightarrow$ 6)-Gal-(1 $\rightarrow$ 6

Complexation of vanadium(V) oxyanions with hexopyranose- and mannopyranoseuronic acid-containing polysaccharides: stereochemical considerations

pp 771-775

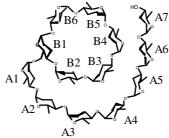
Cesar A. Tischer, Maria Benigna M. Oliveira, Fany Reicher, Marcello Iacomini, Carmem L. O. Petkowicz, Maria Eugênia R. Duarte and Philip A. J. Gorin\*

<sup>51</sup>V NMR spectroscopy showed that in  $D_2O$ , NaVO<sub>3</sub> complexed with *cis*-3,4-diols of α- and β-Galp, and *cis*-2,3-diols of α-Manp units, present in various polysaccharides, but barely with β-Manp and β-Manp A units, as with experiments on related methyl glycosides. Stereochemical factors are involved in ester formation.

#### <sup>1</sup>H NMR spectra of branched-chain cyclomaltohexaoses (α-cyclodextrins)

pp 777-785

Yasuko Ishizuka,\* Tadashi Nemoto, Kenji Kanazawa and Hiroshi Nakanishi



 $^{1}H$  NMR spectra of seven branched-chain cyclomaltohexaoses ( $\alpha$ -CDs) were analyzed intensively using a high-resolution NMR spectrometer.

#### Structural studies on the R-type lipopolysaccharide of Aeromonas hydrophila

pp 787-793

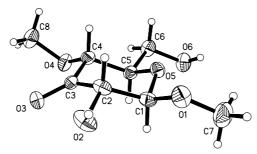
Yuriy A. Knirel,\* Evgeny Vinogradov, Natalia Jimenez, Susana Merino and Juan M. Tomás

$$\begin{array}{c} \beta\text{-D-Gal-}(1\rightarrow 4) \\ \rceil \\ \text{D-}\alpha\text{-D-Hep-}(1\rightarrow 6)\text{-D-}\alpha\text{-D-Hep-}(1\rightarrow 6)\text{-}\beta\text{-D-Glc-}(1\rightarrow 4)\text{-L-}\alpha\text{-D-Hep-}(1\rightarrow 5)\text{-}\alpha\text{-Kdo-}(2\rightarrow \alpha\text{-D-GlcN-}(1\rightarrow 7)\text{-L-}\alpha\text{-D-Hep-}(1\rightarrow 2)\text{-L-}\alpha\text{-D-Hep-}(1\rightarrow 3) } \end{array}$$

#### Crystal and molecular structure of methyl 4-O-methyl-β-D-ribo-hex-3-ulopyranoside

pp 795-799

Immanuel Adorjan, Thomas Rosenau, Antje Potthast, Paul Kosma,\* Kurt Mereiter, Jutta Pauli and Christian Jäger

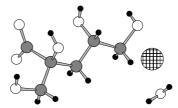


Crystal structure, <sup>13</sup>C CPMAS data, and hydrogen bonding of the title compound are discussed.

#### Crystal structure of sodium isosaccharate, NaC<sub>6</sub>H<sub>11</sub>O<sub>6</sub>·H<sub>2</sub>O

pp 801-805

Ranko P. Bontchev and Robert C. Moore\*



### Crystal structure, solid state and solution conformation of 1D-1,4-di-*O*-[(*S*)-*O*-acetylmandeloyl]-2,3:5,6-di-*O*-isopropylidene-*myo*-inositol

pp 807-811

Kana M. Sureshan, Tomomi Miyasou and Yutaka Watanabe\*

The association of water to cellulose and hemicellulose in paper examined by FTIR spectroscopy

pp 813-818

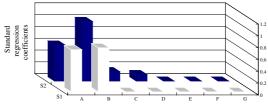
Anne-Mari Olsson\* and Lennart Salmén

The nature of water sorption to cellulose and hemicellulose is examined by FTIR spectroscopy. The moisture was found to be adsorbed homogeneously to all sorbing sites. The dynamic sorption process was also evaluated.

### Quantitative estimate of the effect of cellulase components during degradation of cotton fibers

pp 819-824

Lu-Shan Wang, Yu-Zhong Zhang, Hong Yang and Pei-Ji Gao\*



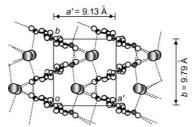
(A) CBHI+EGI, (B) (CBHI+EGI)\*Glucosidase, (C) EGI, (D) EGI+Glucosidase, (E) CBHI, (F) Glucosidase, (G) CBHI+Glucosidase

The results demonstrate that the effects of each of three cellulase components (CBHI, EGI, and  $\beta$ -glucosidase) either alone or in combination in two heterogeneous processes, glucose formation and cellulose solubilization, can be quantitatively distinguished by multivariate stepwise regression analysis.

#### Molecular and crystal structures of chitosan/HI type I salt determined by X-ray fiber diffraction

pp 825-833

Amornrat Lertworasirikul, Shingo Yokoyama, Keiichi Noguchi, Kozo Ogawa and Kenji Okuyama\*

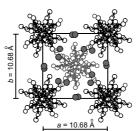


X-ray fiber diffraction analysis indicated that chitosan/HI type I salt adopted an extended 2/1-helix with a fiber repeat of 10.33 Å. The helix was strengthened by O-3···O-5 intramolecular hydrogen bonds. The salt was stabilized by hydrogen bonds between iodide ions and N-2, O-6 of polymer chains together with an electrostatic interaction.

#### Plausible molecular and crystal structures of chitosan/HI type II salt

pp 835-843

Amornrat Lertworasirikul, Keiichi Noguchi, Kozo Ogawa and Kenji Okuyama\*



X-ray fiber diffraction analysis revealed that chitosan/HI type II salt adopted 4/1-helical conformation having a disaccharide as a helical asymmetric unit. One of the O-3···O-5 intramolecular hydrogen bonds is weakened by interacting with iodide ions. This seems to cause the polymer to take the newly found conformation rather than a typical extended 2/1 helix.

### Synthesis and characterization of a brush-like copolymer of polylactide grafted onto chitosan

pp 845-851

Y. Liu,\* F. Tian and K. A. Hu

A brush-like poly(DL)-lactide is grafted onto chitosan as the backbone using Et<sub>3</sub>Al as the catalyst.

# Synthesis of oligosaccharides as potential inhibitors of mycobacterial arabinosyltransferases. Di- and trisaccharides containing C-5 modified arabinofuranosyl residues

pp 853-865

Oana M. Cociorva and Todd L. Lowary\*

Synthesis and biological activities of octyl 2,3,4-tri-O-sulfo- $\alpha$ -L-fucopyranosyl- $(1 \rightarrow 3)$ -2,4-di-O-sulfo- $\alpha$ -L-fucopyranosyl- $(1 \rightarrow 3)$ -2,4-di-O-sulfo- $\beta$ -L-fucopyranoside Yuxia Hua, Guofeng Gu and Yuguo Du\*

pp 867–872

### Synthesis of cluster mannosides via a Ugi four-component reaction and their inhibition against the binding of yeast mannan to concanavalin A

pp 873-879

Yingxia Li,\* Xiaoru Zhang, Shidong Chu, Kunyu Yu and Huashi Guan

#### **NOTES**

### Structures of two O-polysaccharides of the lipopolysaccharide of *Citrobacter youngae* PCM 1538 (serogroup O9)

pp 881-884

Olga G. Ovchinnikova, Nina A. Kocharova, Ewa Katzenellenbogen, George V. Zatonsky, Alexander S. Shashkov, Yuriy A. Knirel,\* Tomasz Lipiński and Andrzej Gamian

Polysaccharide chain I:  $\rightarrow$  2)-α-D-Rhap4NAc-(1  $\rightarrow$  Polysaccharide chain II:  $\rightarrow$  3)-α-D-Rhap4NAc-(1  $\rightarrow$  3)-β-D-Rhap4NAc-(1  $\rightarrow$  3)-β-D-Rhap4NAC-(1 p4NAC-(1 p4NAC-(1 p4NAC-(1 p4NAC-(1 p4NAC-(1 p4NAC-(1

## Chloramine-T-mediated chemoselective hydrolysis of thioglycosides into glycosyl hemiacetals under neutral conditions

pp 885-890

Anup Kumar Misra\* and Geetanjali Agnihotri

$$\begin{array}{c} \text{PO} \\ \text{PO} \\ \text{1} \end{array} \begin{array}{c} \text{OP} \\ \text{SR} \\ \hline \\ \text{CH}_{3}\text{CN} \\ \end{array} \begin{array}{c} \text{PO} \\ \text{PO} \\ \text{2} \end{array} \begin{array}{c} \text{OP} \\ \text{OP} \\ \end{array}$$

A mild, efficient and chemoselective hydrolysis of thioalkylglycosides (1) into their corresponding 1-hydroxy sugars (2) using chloramine-T without affecting other functional groups is reported.

### Two new flavonol glycosides from $\it Gymnema\ sylvestre\ and\ \it Euphorbia\ ebracteolata$

pp 891-895

Xin Liu, Wencai Ye,\* Biao Yu,\* Shouxun Zhao, Houming Wu and Chuntao Che

**1**: R = H, R, = β-D-Glcp-(1 $\rightarrow$ 4)-  $\alpha$ -L-Rhap-(1 $\rightarrow$ 6)-  $\beta$ -D-Galp **2**: R = OH, R, = 6-Q-(3-OH-3-Me-glutaroyl)- $\beta$ -D-Glcp

## Synthesis of 2,3,4,6-tetra-O-benzyl-D-glucal on the gram scale. A convenient method for its facile synthesis and subsequent stereoselective cyclopropanation

pp 897–899

Corin M. Storkey, Anna L. Win and John O. Hoberg\*

A new, facile synthesis of 2,3,4,6-tetra-*O*-benzyl-D-glucal and its subsequent cyclopropanation has been accomplished by bromination and then base-induced elimination of pent-4-enyl 2,3,4,6-tetra-*O*-benzyl-D-glucopyranoside. Yields and selectivities are excellent in addition to the ease of formation of the glycal.

\*Corresponding author

#### **COVER**

Well-defined glycoforms of glycoproteins can easily be obtained by oxidative coupling of synthetic thioaldoses with proteins that have a cysteine moiety in lieu of an asparagine residue carrying natural N-linked oligosaccharides. In vitro glycosylation offers several advantages such as quantitative conjugation, incorporation of oligosaccharides that display high bioactivities and the possibility of using convenient bacterial or yeast protein expression systems. The figure is related to Geert-Jan Boons' *Carbohydrate Research Award* paper, Carbohydr. Res., **2004**, *339*, 181–193.



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