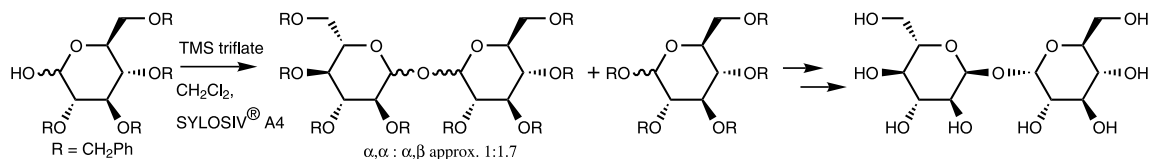


Synthesis of L-trehalose and observations on isomer and by-product formation

Carbohydr. Res. **2003**, 338, 813

Alan H. Haines

Centre for Carbohydrate Chemistry, School of Chemical Sciences and Pharmacy, University of East Anglia, Norwich NR4 7TJ, UK

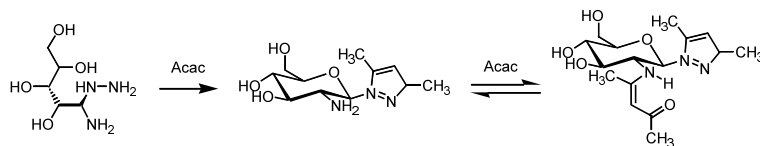


The reaction of acetylacetone with amino sugars: implications for the formation of glycosylpyrazole derivatives

Carbohydr. Res. **2003**, 338, 819

Warren C. Kett

Department of Chemistry, Division of Environmental and Life Sciences, Macquarie University, North Ryde, NSW 2109, Australia



Synthesis of a typical *N*-acetylglucosamine-containing saponin, oleanolic acid 3-yl α -L-arabinopyranosyl-(1 \rightarrow 2)- α -L-arabinopyranosyl-(1 \rightarrow 6)-2-acetamido-2-deoxy- β -D-glucopyranoside

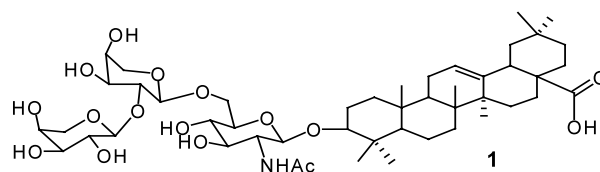
Carbohydr. Res. **2003**, 338, 827

Jiansong Sun,^a Xiuwen Han,^a Biao Yu^b

^a*State Key Laboratory of Catalyst, Dalian Institute of Chemical Physics, Chinese Academy of Sciences, Dalian 116023, PR China*

^b*State Key Laboratory of Bio-organic and Natural Products Chemistry, Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, Shanghai 200032, PR China*

A member of a small structurally conservative family of plant saponins **1** was synthesized employing glycosyl trifluoroacetimidates as key donors.

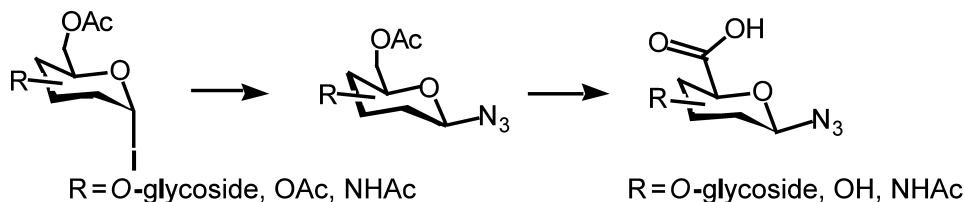


General methods for the synthesis of glycopyranosyluronic acid azides

Carbohydr. Res. **2003**, 338, 835

Laiqiang Ying, Jacquelyn Gervay-Hague

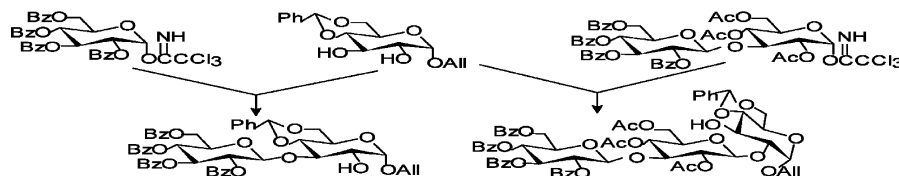
Department of Chemistry, University of California, One Shields Avenue, Davis, CA 95616, USA



Regioselective glycosylation of 4,6-*O*-benzylidenated glucopyranosides

Ying Zeng, Fanzuo Kong

Research Center for Eco-Environmental Sciences, Academia Sinica, Chinese Academy of Sciences, P.O. Box 2871, Beijing 100085, China



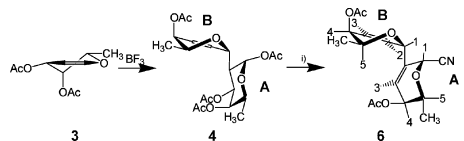
C–H-Deprotonation mediated by a remote *syn*-axial acetoxy group—an unprecedented double bond formation upon cyanation of the dimer from L-fucal

Andreas H. Franz,^a Vyacheslav V. Samoshin,^a Chris Myers,^b Allen D. Hunter,^b Paul H. Gross^a

^aDepartment of Chemistry, University of the Pacific, 3601 Pacific Avenue, Stockton, CA 95211, USA

^bDepartment of Chemistry, Youngstown State University, YSU Structure Center, 1 University Plaza, Youngstown, OH 44555, USA

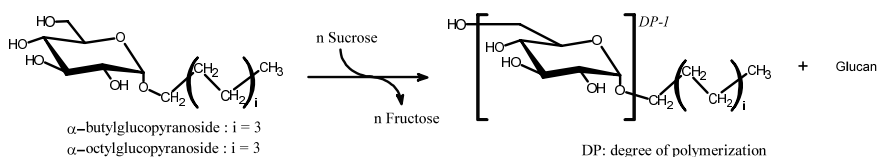
Replacement of the anomeric acetate by a cyanide group in the dimer of di-*O*-acetyl-L-fucal by the action of mild Lewis acid ($\text{Hg}(\text{CN})_2$ – HgBr_2 – Me_3SiCN), resulted not only in the desired transformation but also in an additional double bond between C-2A and C-3A.



Glucosylation of α -butyl- and α -octyl-D-glucopyranosides by dextran sucrose and alternansucrase from *Leuconostoc mesenteroides*

Gaëtan Richard, Sandrine Morel, René-Marc Willemot, Pierre Monsan, Magali Remaud-Simeon

Département de Génie Biochimique et Alimentaire, Centre de Bioingénierie Gilbert Durand, UMR CNRS 5504, UMR INRA 792, INSA, 135 Avenue de Rangueil, 31077 Toulouse 4, France



On the reaction pathways and determination of transition-state structures for retaining α -galactosyltransferases

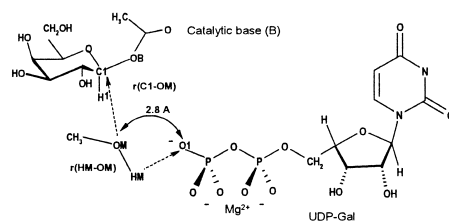
Isabelle André,^a Igor Tvaroška,^{a,b} Jeremy P. Carver^a

^aGlycoDesign Inc., 480 University Avenue, Suite 900, Toronto, ON, Canada M5G 1V2

^bInstitute of Chemistry, Slovak Academy of Sciences, Bratislava, Slovak Republic

The catalytic mechanism of retaining glycosyltransferases via a double-displacement mechanism was investigated using high level ab initio calculations up to the DFT/B3LYP/6-311++G**//DFT/B3LYP/6-31G* level. Results show that a mechanism using UDP as a general base in the second step of the reaction appears to be the most consistent with experimental data.

Covalent Glycosyl-Enzyme intermediate (CGE)



Synthesis and structural analysis of five novel oligosaccharides prepared by glucosyltransfer from β -D-glucose 1-phosphate to isokestose and nystose using *Thermoanaerobacter brockii* kojibiose phosphorylase

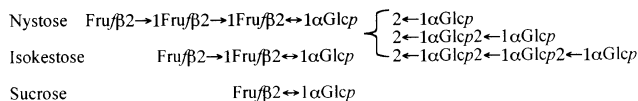
Carbohydr. Res. **2003**, 338, 879

Hideki Okada,^a Eri Fukushi,^b Shuichi Onodera,^a Tomoyuki Nishimoto,^c Jun Kawabata,^b Masanori Kikuchi,^a Norio Shiomi^a

^aDepartment of Food Production and Utility Development, Graduate School of Dairy Science Research, Rakuno Gakuen University, Ebetsu 069-8501, Japan

^bGraduate School of Agriculture, Hokkaido University, Sapporo 060-8589, Japan

^cHayashibara Biochemical Laboratories, Inc., Okayama 700-0834, Japan

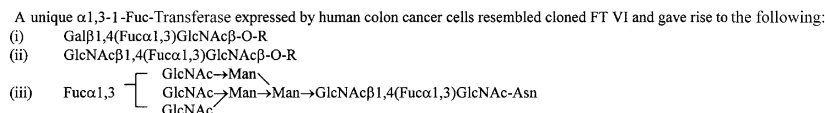


The binding characteristics and utilization of *Aleuria aurantia*, *Lens culinaris* and few other lectins in the elucidation of fucosyltransferase activities resembling cloned FT VI and apparently unique to colon cancer cells

Carbohydr. Res. **2003**, 338, 887

E.V. Chandrasekaran, Ram Chawda, John M. Rhodes, Robert D. Locke, Conrad F. Piskorz, Khushi L. Matta

Department of Molecular and Cellular Biophysics, Roswell Park Cancer Institute, Elm and Carlton Streets, Buffalo, NY 14263, USA



Isolation, structural characterization, and immunological evaluation of a high-molecular-weight exopolysaccharide from *Staphylococcus aureus*

Carbohydr. Res. **2003**, 338, 903

Joseph G. Joyce,^a Chitrananda Abeygunawardana,^a Qiuwei Xu,^a James C. Cook,^a Robert Hepler,^a Craig T. Przysiecki,^a Karen M. Grimm,^a Keith Roper,^a Charlotte C. Yu Ip,^a Leslie Cope,^a Donna Montgomery,^a Mason Chang,^a Sherilyn Campie,^a Martha Brown,^a Tessie B. McNeely,^a Julie Zorman,^a Tomas Maira-Litrán,^b Gerald B. Pier,^b Paul M. Keller,^a Kathrin U. Jansen,^a George E. Mark III^a

^aDepartments of Virus and Cell Biology, Bioprocess and Bioanalytical Research, and Microbial Vaccine Research, Merck Research Laboratories, WP16-107, P.O. Box 4, West Point, PA 19486, USA

^bDepartment of Medicine, Channing Laboratory, Brigham and Women's Hospital, Harvard Medical School Boston, Boston, MA 02115, USA

A β -(1→6)-linked glucosamine-containing exopolysaccharide isolated from *Staphylococcus aureus* MN8m ($M_r > 300,000$ Da) and shown to be essentially devoid of succinylate and containing 45–60% *N*-acetyl is active in hemagglutination assays and is immunogenic in mice when coupled to a protein carrier.

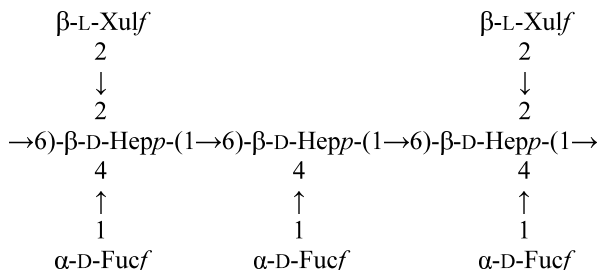
The structure of the antigenic polysaccharide produced by *Eubacterium saburreum* T15

Carbohydr. Res. **2003**, 338, 923

Naomi Sato,^a Futoshi Nakazawa,^a Teiichiro Ito,^a Tsutomu Hoshino,^b Etsuro Hoshino^a

^aDepartment of Oral Health Science, Division of Oral Ecology in Health and Infection, Course for Oral Life Science, Niigata University Graduate School of Medical and Dental Sciences Gakkocho-dori 2, Niigata 951-8514, Japan

^bDepartment of Applied Biological Chemistry, Faculty of Agriculture, Niigata University, Ikarashi, Niigata 950-2181, Japan



Fungal chitin–glucan derivatives exert protective or damaging activity on plasmid DNA

Grigorij Kogan,^a Peter Rauko,^b Eva Machová^a

^aInstitute of Chemistry, Slovak Academy of Sciences, 845 38 Bratislava, Slovakia

^bCancer Research Institute, Slovak Academy of Sciences, 833 91 Bratislava, Slovakia

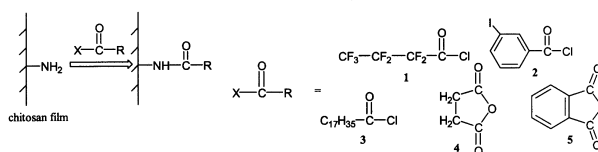
Using carboxymethylation, sulfoethylation, and subsequent ultrasonic treatment, lower molecular weight water-soluble derivatives of crude fungal chitin–glucan complex were prepared and evaluated for their effects on DNA strand breakage.

Surface modification of chitosan films Effects of hydrophobicity on protein adsorption

Varawut Tangpasuthadol,^{a,b} Noppong Pongchaisirikul,^a Vipavee P. Hoven^{a,b}

^aDepartment of Chemistry, Organic Synthesis Research Unit, Faculty of Science, Chulalongkorn University, Bangkok 10330, Thailand

^bCenter for Chitin/Chitosan Biomaterials, Chulalongkorn University, Bangkok 10330, Thailand



Unusual structures in the polysaccharides from the red seaweed *Pterocladia capillacea* (Gelidiaceae, Gelidiales)

María I. Errea, María C. Matulewicz

Departamento de Química Orgánica, CIHIDECAR-CONICET, Facultad de Ciencias Exactas y Naturales, Universidad de Buenos Aires, Pabellón 2-Ciudad Universitaria, 1428 Buenos Aires, Argentina

Sequential extraction of *Pterocladia capillacea* with water at room temperature and then at 50 °C gave two products, which by thorough fractionation and further structural analysis, afforded evidence of the presence of 3-substituted, 4-linked D-galactopyranosyl units in polysaccharides from this seaweed.

NMR and modelling studies of disaccharide conformation

Norman W.H. Cheetham,^a Paramita Dasgupta,^a Graham E. Ball^b

^aSchool of Chemical Sciences, The University of New South Wales, Sydney 2052, Australia

^bNMR Facility, School of Chemical Sciences, The University of New South Wales, Sydney 2052, Australia

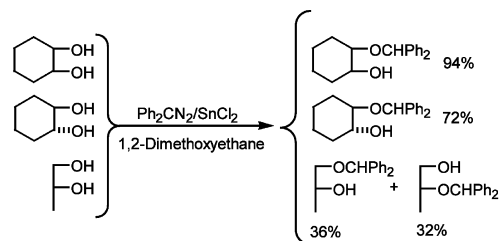
Multiple ¹³C site-selective excitation experiments using ¹H decoupling in conjunction with pulsed field gradient-enhanced spectroscopy were used to determine ³J_{C,H} values across disaccharide glycosidic linkages. These were compared with couplings calculated from molecular dynamics simulations with the explicit inclusion of water.

Tin(II) chloride catalyzed reactions of diazodiphenylmethane with vicinal diols in an aprotic solvent

The reactions with *cis*- and *trans*-1,2-cyclohexanediols and 1,2-propanediol

Sigthor Petursson

Faculty of Natural Resource Sciences, University of Akureyri,
600 Akureyri, Iceland



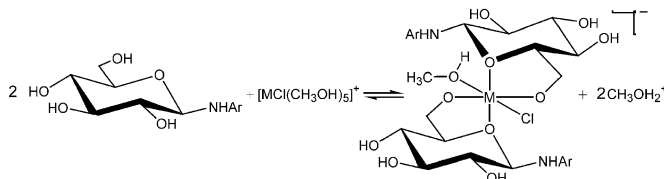
Carbohydr. Res. **2003**, 338, 963

Mechanism of the transition-metal-catalyzed

mutarotation reaction of *N*-(*p*-chlorophenyl)- β -D-glucopyranosylamine in methanol

Kazimiera Smiataczowa, Jarosław Kosmalski, Teresa Widernik, Zygmunt Warnke

Faculty of Chemistry, University of Gdańsk, Sobieskiego 18, PL-80-952 Gdańsk, Poland



Carbohydr. Res. **2003**, 338, 969

X-ray structure of the cyclomaltohexaicosaoose triiodide

inclusion complex provides a model for amylose–iodine at atomic resolution

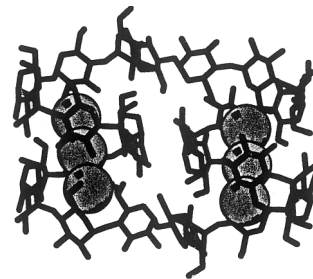
O. Nimz,^a K. Geßler,^a I. Usón,^b S. Laetig,^c H. Welfle,^c G.M. Sheldrick,^b
W. Saenger^a

^aInstitut für Kristallographie, Freie Universität Berlin, Takustrasse 6, D-14195 Berlin, Germany

^bInstitut für Anorganische Chemie, Universität Göttingen, Tammannstrasse 4,
D-37077 Göttingen, Germany

^cMax-Delbrück-Centrum für Molekulare Medizin, Robert-Rössle-Str. 10, D-13092 Berlin-Buch,
Germany

The V-amylose channels of cyclomaltohexaicosaoose accommodate linear I_3^- chains as guest molecules and I^- in interstices. C–H \cdots I contacts are the dominant interaction of the I_3^- polyiodide subunits with the carbohydrate.



Carbohydr. Res. **2003**, 338, 977

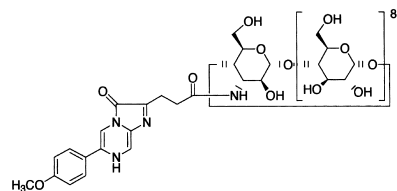
Enhanced chemiluminescence of 6-(4-methoxyphenyl)-

imidazo[1,2-*a*]pyrazin-3(7*H*)-one by attachment of a cyclomaltooligosaccharide (cyclodextrin). Attachment of cyclomaltononaose (δ -cyclodextrin)

Katsunori Teranishi,^a Tatsuya Nishiguchi,^a Haruhisa Ueda^b

^aFaculty of Bioresources, Mie University, 1515 Kamihama, Tsu, Mie 514-8507, Japan

^bFaculty of Pharmaceutical Sciences, Hoshi University, 4-41, Ebara 2-chome,
Shinagawa-ku, Tokyo 142-8501, Japan



Monoesterification of di-*O*-isopropylidene and di-*O*-cyclohexylidene *chiro*-inositols

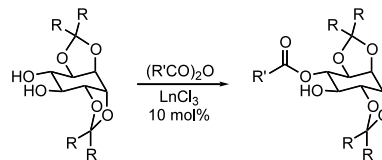
Carbohydr. Res. **2003**, *338*, 995

Ghislaine Cousins,^a Andrew Falshaw,^b John O. Hoberg^a

^a*School of Chemical and Physical Sciences, Victoria University of Wellington, Box 600, Wellington, New Zealand*

^b*Industrial Research Limited, PO Box 31-310, Lower Hutt, New Zealand*

Monoesterification of D- or L-*chiro*-inositols protected as diacetals proceeds in excellent selectivity and yields. The metal-catalyzed, one-step reaction proceeds at room temperature under an air atmosphere and has been developed using a range of examples.



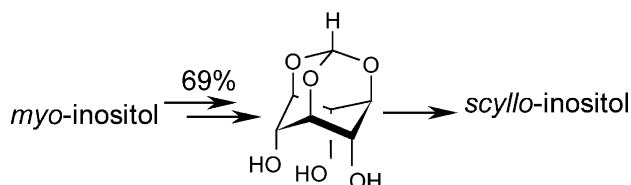
Sulfonate protecting groups.

Improved synthesis of *scyllo*-inositol and its orthoformate from *myo*-inositol

Carbohydr. Res. **2003**, *338*, 999

Manash P. Sarmah, Mysore S. Shashidhar

National Chemical Laboratory, Division of Organic Synthesis, Pune 411 008, India



Fungal β -*N*-acetylhexosaminidases with high

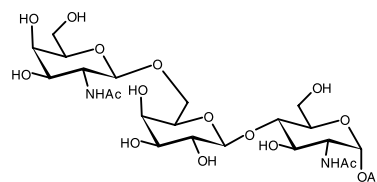
β -*N*-acetylgalactosaminidase activity and their use for synthesis of β -GalNAc-containing oligosaccharides

Carbohydr. Res. **2003**, *338*, 1003

Lenka Weignerová,^a Petra Vavrušková,^a Andrea Pišvejcová,^a Joachim Thiem,^b Vladimír Křen^a

^a*Laboratory of Biotransformation, Institute of Microbiology, Academy of Sciences of the Czech Republic, Vídeňská 1083, Prague 4, CZ 142 20, Czech Republic*

^b*Institute of Organic Chemistry, University of Hamburg, Martin-Luther-King-Platz 6, D 20146 Hamburg, Germany*



Structure of the O-specific polysaccharide of *Providencia*

rustigianii O14 containing *N*^ε-[(*S*)-1-carboxyethyl]-*N*^α-(D-galacturonoyl)-L-lysine

Carbohydr. Res. **2003**, *338*, 1009

Nina A. Kocharova,^a George V. Zatonsky,^a Agnieszka Torzewska,^b Zuzanna Maciejka,^b Olga V. Bystrova,^a Aleksander S. Shashkov,^a Yuriy A. Knirel,^a Antoni Rozalski^b

^a*N.D. Zelinsky Institute of Organic Chemistry, Russian Academy of Sciences, Leninsky Prospekt 47, 119991 Moscow, GSP-1, Russia*

^b*Department of Immunobiology of Bacteria, Institute of Microbiology and Immunology, University of Lodz, Banacha 12/16, 90-237 Lodz, Poland*

The O-specific polysaccharide of *Providencia rustigianii* O14 was obtained by mild acid degradation of the LPS and studied by chemical methods and NMR spectroscopy, including 2D ¹H, ¹H COSY, TOCSY, NOESY, and ¹H, ¹³C HSQC experiments.

