GRAPHICAL ABSTRACTS

Cyclodextrins, Cyclomannins, and Cyclogalactins with five and six (1—4)-linked Sugar Units: a comparative Assessment of their Conformations and Hydrophobicity Potential Profiles

Tetrahedron: Asymmetry 1994, 5, 2045



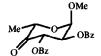
Institut für Organische Chemie Technische Hochschule Petersenstraße 22 D-64287 Darmstadt (Germany)



Tetrahedron: Asymmetry 1994, 5, 2061

THE SYNTHESIS OF NOVEL MIMICS OF THE SIALYL LEWIS X DETERMINANT Nigel. M. Allanson. Alan H. Davidson, Christopher D. Floyd, Fionna M. Martin British Biotech Pharmaceuticals Ltd, Watlington Rd, Oxford, OX4 5LY, UK

Dissacharide mimics of the sLex ligands have been prepared as antagonists against binding to E-selectin. The stereochemistry of nucleophilic addition to the keto group of the fucosyl ketone 7 is determined by the conditions employed. Lewis acid mediated conditions give rise to attack from the Re face while the addition of metal promoted dianions proceeds via attack from the Si face.



7

Synthesis of α -D-(1 \rightarrow 3) and α -D-(1 \rightarrow 4)-C-linked Galactosides of D-Mannose Derivatives. Conformation of α -C-Galactosides.

Tetrahedron: Asymmetry 1994, 5, 2077

Rafael Ferritto and Pierre Vogel*, Section de Chimie de l'Université de Lausanne, BCH Dorigny, CH-1015 Lausanne, Switzerland.

Tetrahedron: Asymmetry 1994, 5, 2093

Sialyl Le^x Structures In O-Glycans Attached to Lysosomal Membrane Glycoproteins, Lamp-1 and Lamp-2. Comparison to Those In N-Glycans.

Kentaro Maemura and Minoru Fukuda

the Glycobiology Program, La Jolla Cancer Research Foundation 10901 North Torrey Pines Road, La Jolla, CA 92037

The structures of O-glycans and N-glycans were determined on lysosomal membrane glycoproteins, lamp-1 and lamp-2 isolated from HL-60 cells. The O-glycans were found to express as much sialyl Le^x structure as N-glycans.

The Use of O-Glycosyl Trichloroacetimidates in

Tetrahedron: Asymmetry 1994, 5, 2109

The Synthesis of Unsymmetrical Trehalose Analogues.

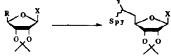
Tor E.C.L. Rønnow, Morten Meldal, and Klaus Bock*.

Carlsberg Laboratory, Department of Chemistry, Gamle Carlsberg Vej 10, DK-2500 Valby (Denmark).

Stereocontrolled Radical Reactions in Carbohydrate and Nucleoside Chemistry.

Tetrahedron: Asymmetry 1994, 5, 2123

Derek H.R. Barton, ^a Stephane D. Géro, ^b Béatrice Quiclet-Sire ^b, Mohammad Samadi ^b. ^aDepartment of Chemistry, Texas A&M University, College Station, Texas 77843, U.S.A. ^b Institut de Chimie des Substances Naturelles, C.N.R.S., 91198 Gif-sur-Yvette, France.

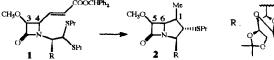


X= methoxy, uracil, adenine COOH Y= CO₂Me, SO₂Ph,... The radicals, generated by photolysis of 2,3-dimethyl ketals of N-hydroxy-2-thiopyridone uronic esters, reacted stereoselectively with electron deficient olefins leading to highly functionalised chain-elongated pentofuranosides, hexapyranosides and pentafuranosyl-nucleosides through the 4,5 and 4' radicals, respectively.

THE USE OF RADICAL DECARBOXYLATION IN THE PREPARATION OF 1-METHYLCARBAPENEM ANTIBIOTIC PRECURSORS FROM D-GLUCOSAMINE.

Tetrahedron: Asymmetry 1994, 5, 2137

J. Anaya**, D.H.R.Bartonb, M.C. Caballero*, S.D.Gero*, M. Grande*, N.M. Laso** and J.I.M. Hernando*. *Departamento de Química Orgánica. Universidad de Salamanca, 37008 Salamanca, Spain. *Department of Chemistry, Texas A&M University, College Station, Texas 77843, USA. *Institut de Chimie des Substances Naturelles, C.N.R.S., 91198 Gif-sur-Yvette, France



Radical cyclization and radical decarboxylation were used for the transformation of β -lactams 1 into 1-methylcarbapenem antibiotic precursors 2 starting with D-glucosamine.

RADICAL CYCLIZATIONS ON SUGAR TEMPLATES: STEREOSELECTIVE SYNTHESIS OF FUSED Y-BUTYROLACTONES OF CARBOHYDRATES. Tetrahedron: Asymmetry 1994, 5, 2141

Sonsoles Velázquez and María-José Camarasa

Instituto de Química Médica (C.S.I.C.), Juan de la Cierva 3, 28006-Madrid, Spain

x

The Use Of Free Radical Cyclization In The Synthesis Of Compounds Related To The Mannostatins. A. H. Ingall, P. R. Moore^b and S. M. Roberts.^b

- a Fisons Pharmaceuticals, Bakewell Road, Loughborough, LE11 ORH.
- b Department of Chemistry, Exeter University, Exeter, Devon, EX4 4QD.

Tetrahedron: Asymmetry 1994, 5, 2163

DIBUTYLSTANNYLENE ACETALS: USEFUL INTERMEDIATES FOR THE REGIOSELECTIVE SULFATION OF GLYCOSIDES.

Bénédicte Guilbert, Nicola J. Davis, Melanie Pearce, Robin T. Aplin and Sabine L. Flitsch*, The Dyson Perrins Laboratory, South Parks Road, Oxford OX1 3OY, Great Britain

Asymmetric Induction in [2+2]Cycloaddition of Chlorosulfonyl Isocyanate to 1,2-O-Isopropylidene -3-O-vinyl-glycofuranoses. Zbigniew Kałuża*, Bartłomiej Furman,

Tetrahedron: Asymmetry 1994, 5, 2179

Madhumeeta Patel[†] and Marck Chmiclewski. Institute of Organic Chemistry, Polish Academy of Sciences, 01-224 Warsaw, Poland. [†]Stevens Institute of Technology, Hoboken, N.J. 07030, USA

FACILE PREPARATION OF GLYCOSYL DONORS FOR OLIGOSACCHARIDE

Tetrahedron: Asymmetry 1994, 5, 2187

SYNTHESIS 2-AZIDO-2-DEOXYHEXOPYRANOSYL BUILDING BLOCKS Therese Buskas, Per J. Garegg, Peter Konradsson*, and Jean-Luc Maloisel

Department of Organic Chemistry, Arrhenius Laboratory, Stockholm University, S-106 91 Stockholm, Sweden

(i) RSSiMe₃, Znl₂, ClCH₂CH₂Cl, (ii) NaOH 1M (iii) N₃OTf, DMAP, MeOH

Also with 4-pentenyl β-D-glucoside. Also in galactose and mannose series.

Sphingosine and phytosphingosine from D-Threose

Synthesis of a 4-Keto-Ceramide

Robert Wild and Richard R. Schmidt

Fakultät Chemie, Universität Konstanz, Postfach 5560 M 725, D-78434 Konstanz, Germany

Sphingosine Ph
$$O$$
 Phytosphingosine Phytosphingosine 1 Sa: $R^1 = H$, $R^2 = OH$ Sx: $R^1 = OH$, $R^2 = H$

SYNTHESIS OF A SIALIC ACID ANALOG WITH THE ACETAMIDO GROUP AT C-4

Tetrahedron: Asymmetry 1994, 5, 2209

A. Dondoni,* A. Boscarato, and A. Marra

An isomer of N-acetylneuraminic acid with the acetamido group at C-4 has been synthesized through stereoselective 1,4-conjugate addition of trimethylsilyl azide to a 2-thiazolyl α,β -enone bearing a protected D-mannose moiety at C- β .

Diastereoselective Cleavage of β -Glucosylsulfoxides by β -Glucosidase Olaf Karthaus, Shin-Ichiro Shoda, Shiro Kobayashi Tohoku University, Aoba, Sendai , Japan

Tetrahedron: Asymmetry 1994, 5, 2213

$$CH_2OH$$
 O B B -glucosidase CH_2OH OH CH_2OH O

A SYNTHETIC STUDIES ON THE TRICHOTHECENE FAMILY FROM D-GLUCOSE

Tetrahedron: Asymmetry 1994, 5, 2217

Jun Ishihara, Rie Nonaka, Yuki Terasawa, Kin-ichi Tadano,* and Seiichiro Ogawa

Departnment of Applied Chemistry, Keio University, Hıyoshi, Yokohama 223, Japan

Compounds 1 and 2 as promising intermediates for total synthesis of the trichothecenes were synthesized from D-glucose both in enantiomerically pure forms.

Stereocontrolled Synthesis of Spirodihydrouracil Nucleoside Hiromi Sano,* Shigeru Mio,* Junko Kitagawa, and Soji Sugai Agroscience Research Laboratories, Sankyo Co. Ltd., 1041 Yasu-cho, Yasu-gun, Shiga-ken 520-23, Japan

DIFRUCTOSE DIANHYDRIDES AS SYNTHETIC INTERMEDIATES. A SYNTHESIS OF 3,6-ANHYDRO-keto-D-FRUCTOSE

Tetrahedron: Asymmetry 1994, 5, 2241

Jacques Defaye and José M. García Fernández, CNRS and CEA, Département de Recherche Fondamentale sur la Matière Condensée/SESAM, Centre d'Etudes de Grenoble, F-38054 Grenoble, France

A convenient synthesis of the title α -hydroxyketone 3 from readily available α -D-fructofuranose β -D-fructofuranose 1,2',2,1'-dianhydride (1) is described.

The Utility of 2,5-Dideoxy-2,5-imino-D-mannitol as a PFP Enzyme Inhibitor

Tetrahedron: Asymmetry 1994, 5, 2251

Mukund S. Chorghade ¹*, Csaba T. Cseke² and Paul S. Liu³†

¹ Abbott Laboratories, North Chicago, IL 60064, ² Dow Elanco, Indianapolis, IN 46268, ³ Marion Merrell Dow Research Center, Cincinnati, OH 45215

2,5-Dideoxy-2,5-imino-D-mannitol was synthesized from an arabinofuranose derivative. Mercuric acetate cyclization of an ene-carbamate was the key step. The compound proved to be an inhibitor of the pyrophosphate-fructose-6-phosphate-1-phosphotransferase (PFP) enzyme and has potential utility in the biorational design of herbicides.

TWO CATIONIC ANALOGUES OF DISACCHARIDE HETEROLYSIS: POTENTIAL EPITOPES FOR PRE-

Tetrahedron: Asymmetry 1994, 5, 2255

PARING ANTIBODIES WITH SACCHARIDASE ACTIVITY.

Jochen Lehmann' and Beatrice Rob, Institut für Organische Chemie und Biochemie, Universität Freiburg, D 79104 Freiburg, Germany

Two disaccharide analogues containing the structural element of a guanidinium ion have been prepared.

STEREOCONTROLLED HYDROXYMETHYLATION OF CARBOHYDRATE IMINES. FORMAL SYNTHESIS OF DESTOMIC ACID AND LINCOSAMINE

F.L. van Delft, M. de Kort, G.A. van der Marel and J.H. van Boom' Leiden Institute of Chemistry, Gorlaeus Laboratories,

P.O. Box 9502, 2300 RA, Leiden, The Netherlands

Addition of [(dimethylphenylsilyl)methyl]magnesium chloride to 6-benzylimino-6-deoxy-1,2;3.4-di-O-isopropylidene-α-D-galactopyranose mediated by Ce(III)Cl₃ or Cul/BF₃·Et₂O proceeds with excellent diastereoselectivity to give the syn or anti-adduct 1 and 2, respectively.

SYNTHESIS OF NOVEL FUSED RING C-GLYCOSIDES

Tetrahedron: Asymmetry 1994, 5, 2265

Daniel E. Levy,* Falguni Dasgupta, and Peng Cho Tang Glycomed, Inc. 860 Atlantic Ave. Alameda, CA 94501

The Lewis acid mediated addition of olefins to O-benzyl protected sugars is studied. The results show the formation of C-glycosides with α selectivity and a propensity to cyclize with loss of a benzyl group.

The Synthesis of Difluoromethylene-Linked C-Glycosides and C-Disaccharides

Tetrahedron: Asymmetry 1994, 5, 2269

Timothée F. Herpin, William B. Motherwell, * Matthew J. Tozer

- a. Department of Chemistry, University College, 20 Gordon Street, London, WC1H 0AJ, UK.
- b. James Black Foundation, 68 Half Moon Lane, Dulwich, London, SE24 9JE, UK.

SYNTHESIS OF TETRASACCHARIDE FROM SERRATIA MARCESCENS

Tetrahedron: Asymmetry 1994, 5, 2283

Jian Zhang, Jianmin Mao, Hongming Chen, Mengshen Cai* Department of Organic chemistry, School of Pharmaceutical Sciences, Beijing medical University, Beijing 100083, China

A glycosylation of oligosaccharide via Schmidt method.

SYNTHESIS OF HYALURONIC ACID RELATED DI- AND TETRA-

Tetrahedron: Asymmetry 1994, 5, 2291

Tetrahedron: Asymmetry 1994, 5, 2303

SACCHARIDES HAVING A GLUCURONIC ACID AT THE REDUCING END

Ted M. Slaghek^a, Teija K. Hyppönen^a, Tomoya Ogawa^b, Johannis P. Kamerling^a, and Johannes F. G. Vliegenthart^{a,*}

^a Bijvoet Center, Department of Bio-Organic Chemistry, Utrecht University, Utrecht, Netherlands; ^b The Institute of Physical and Chemical Research (RIKEN), Wako-shi, Saitama, Japan

The syntheses of the 4-methoxyphenyl (MP) glycosides of β -D-GlcpNAc-(1 \rightarrow 4)- β -D-GlcpA and β -D-GlcpNAc-(1 \rightarrow 4)- β -D-GlcpNAc-(

HOLDH AND HOLDH

HOLDH OH OH OH OH OH OH OH OH OH

Mild Stereoselective Syntheses of Thioglycosides Under PTC Conditions and their Use as Active and Latent Glycosyl Donors.

S. Cao, S.J. Meunier, F.O. Andersson, M. Letellier and R.Roy*,

Department of Chemistry, University of Ottawa, Ottawa, ON., Canada K1N 6N5.

"Active" and "latent" throglycosyl donors and acceptors were synthesized with complete anomeric inversions from glycosyl halides under mild PTC conditions The strategy was applied to sialic acid containing disaccharides.

AcO OAc COME
SRn4-X

BzC

N.

NIS, TIOH ACC

Acc COMe

SPh4NO

X = H, Me, OMe, NHAc (No reaction when $X = NO_2$)