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Special Issue

In Celebration of the 65th Birthday of Professor George Fleet

Guest editor:
Professor Antony Fairbanks

University of Canterbury, Private Bag 4800, Christchurch 8140, New Zealand

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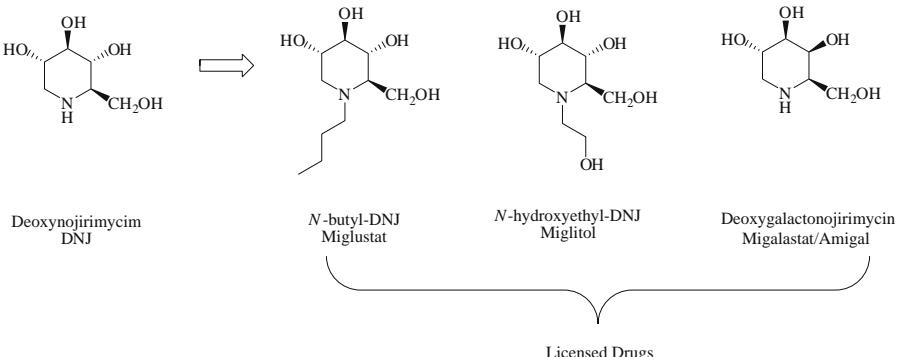
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Iminosugars: from botanical curiosities to licensed drugs

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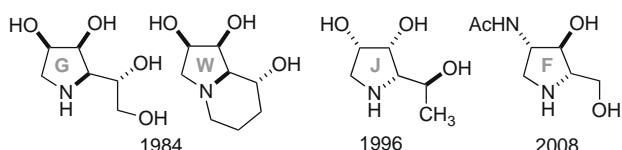
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A silver-lined anniversary of Fleet iminosugars: 1984–2009, from DIM to DRAM to LABNAC

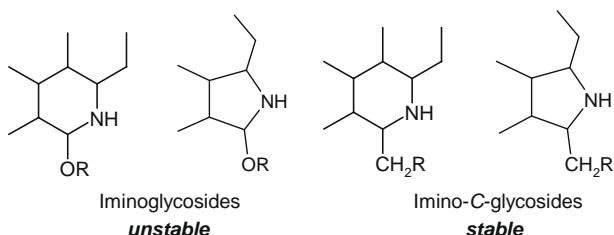
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Benjamin G. Davis *



Tactics and strategies for the synthesis of iminosugar C-glycosides: a review

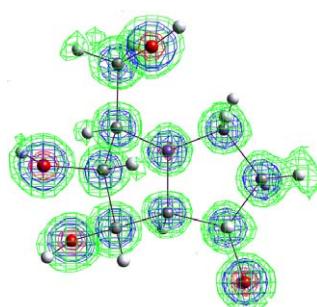
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Philippe Compain ^{*}, Vincent Chagnault, Olivier R. Martin

The purpose of this review is to provide an overview of the versatile strategies that have been developed to synthesize imino-C-glycosides as stable analogues of imino-glycosides and imino-glycoconjugates.

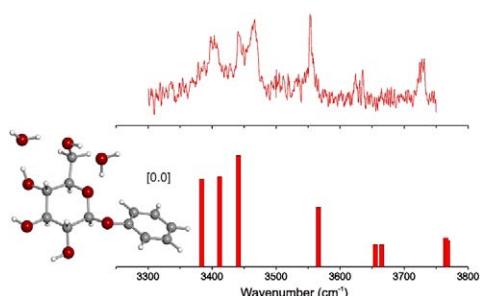
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Amber L. Thompson, David John Watkin ^{*}

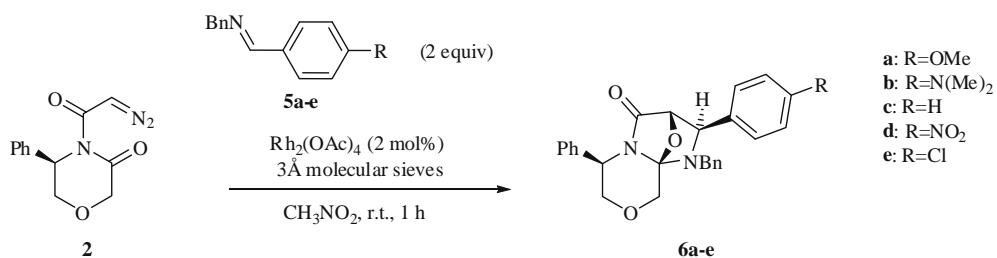
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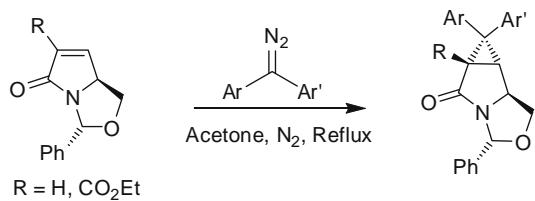
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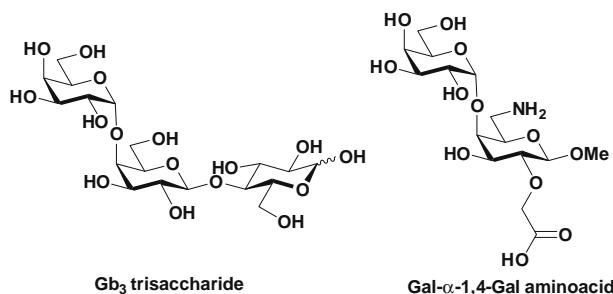
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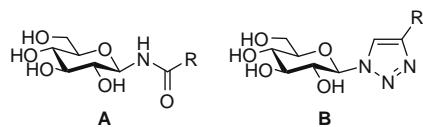
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Evangelia D. Chrysina *, Éva Bokor, Kyra-Melinda Alexacou, Maria-Despoina Charavgi, George N. Oikonomakos, Spyros E. Zographos, Demetres D. Leonidas, Nikos G. Oikonomakos, László Somsák *

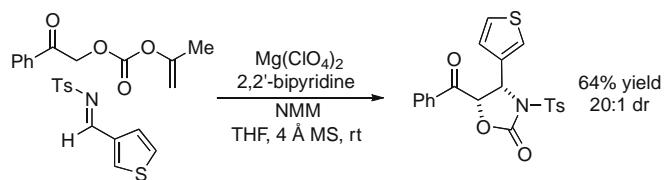


Kinetic measurements and X-ray crystallographic studies of rabbit muscle glycogen phosphorylase *b* enzyme-inhibitor complexes prove bioisosteric relationship of *N*-acyl- β -D-glucopyranosylamines (A) and 1-(β -D-glucopyranosyl)-4-substituted-1,2,3-triazoles (B).

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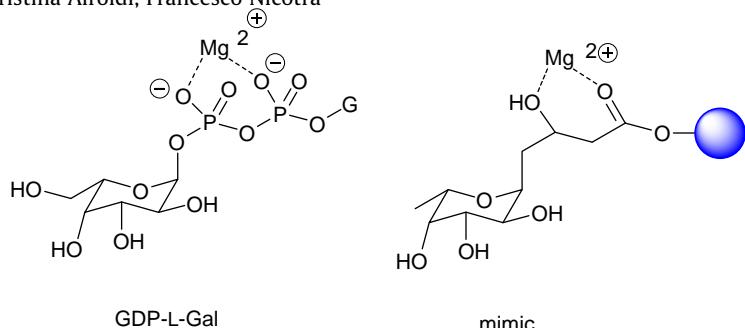
Nikki E. Stainforth, Gary A. Cutting, Matthew P. John, Michael C. Willis *



Solid-phase supported mimic of GDP-L-galactose

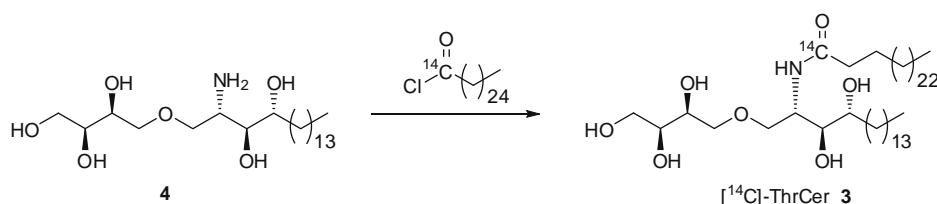
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**Articles****Synthesis of threitol ceramide and [¹⁴C]threitol ceramide, non-glycosidic analogues of the potent CD1d antigen α -galactosyl ceramide**

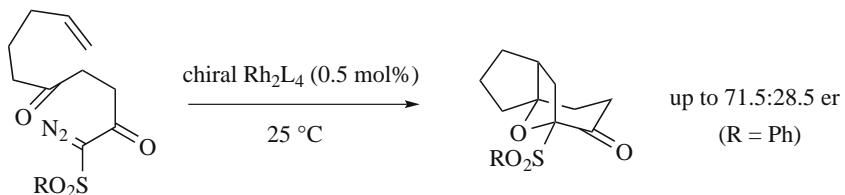
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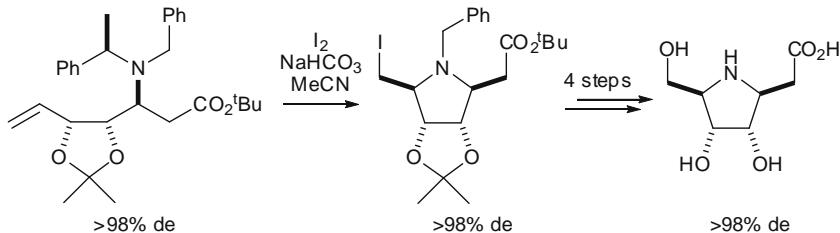
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**Iodine-mediated ring-closing iodoamination with concomitant N-debenzylation for the asymmetric synthesis of polyhydroxylated pyrrolidines**

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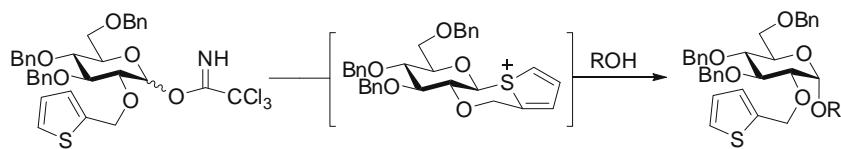
Stephen G. Davies *, Rebecca L. Nicholson, Paul D. Price, Paul M. Roberts, Angela J. Russell, Edward D. Savory, Andrew D. Smith, James E. Thomson



A novel ring-closing alkene iodoamination with concomitant N-debenzylation protocol provides a direct route for the asymmetric synthesis of polyhydroxylated pyrrolidines from homochiral β -amino esters.

Stereoselective synthesis of α -glucosides by neighbouring group participation via an intermediate thiophenium ion
 Daniel J. Cox, Antony J. Fairbanks *

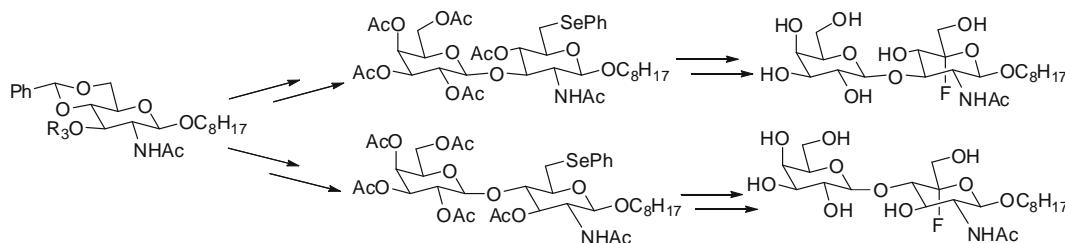
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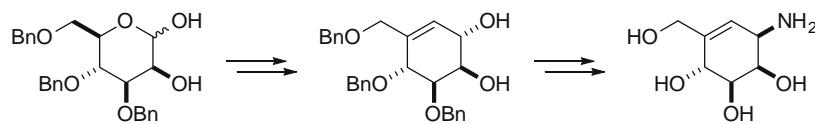
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β -Mannosidase and β -hexosaminidase inhibitors: synthesis of 1,2-bis-*epi*-valienamine and 1-*epi*-2-acetamido-2-deoxy-*valienamine from D-mannose*

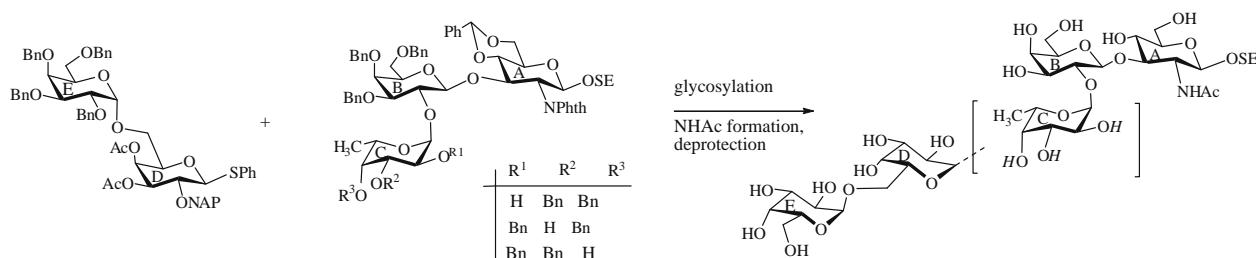
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Clinton Ramstadius, Omid Hekmat, Lars Eriksson, Henrik Stålbrand, Ian Cumpstey *



Synthesis of three regioisomers of the pentasaccharide part of the Skp1 glycoprotein of *Dictyostelium discoideum*
 Zoltán B. Szabó, Mihály Herczeg, Anikó Fekete, Gyula Batta, Anikó Borbás, András Lipták *, Sándor Antus

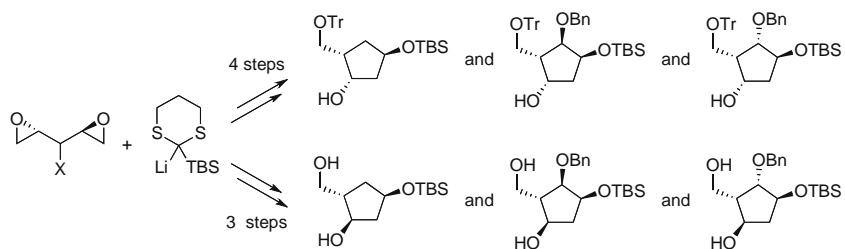
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An enantioselective synthesis of carbafuranose sugars based on a linchpin carbacyclisation approach

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Leo M.H. Leung, Mark E. Light, Vicky Gibson, Bruno Linclau *

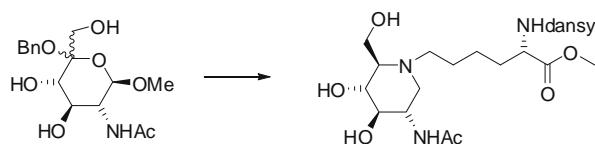


Enantiopure carbafuranose derivatives were synthesised via a linchpin carbacyclisation process starting from 1,4-bisepoxides. Both 2-deoxy and 2-deoxy-6-hydroxycarbafurano derivatives were obtained, which were converted to suitably protected precursors for carbanucleoside synthesis.

2-Acetamino-1,2-dideoxyojirimycin–lysine hybrids as hexosaminidase inhibitors

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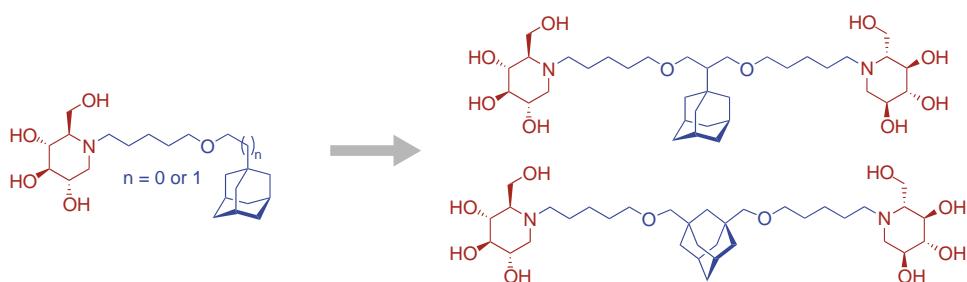
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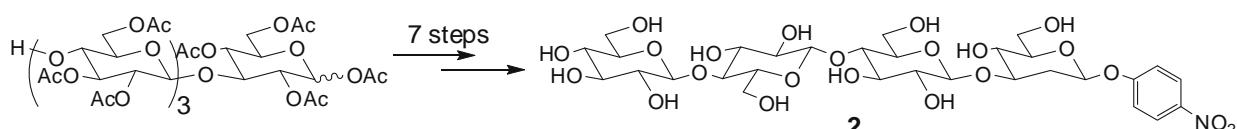
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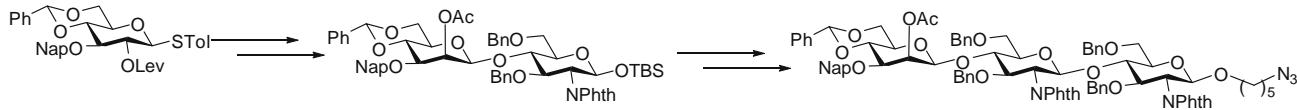
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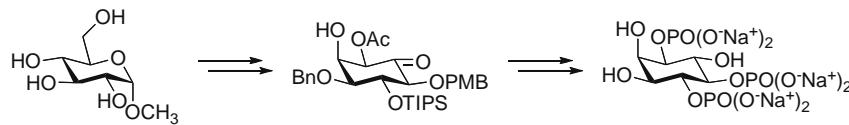
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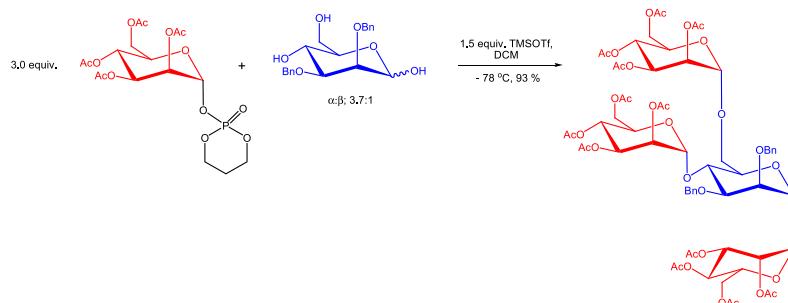
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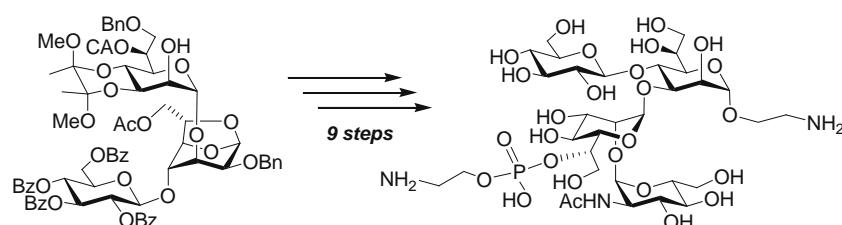
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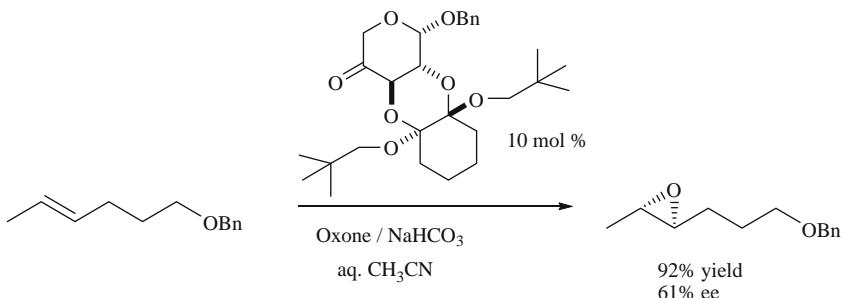
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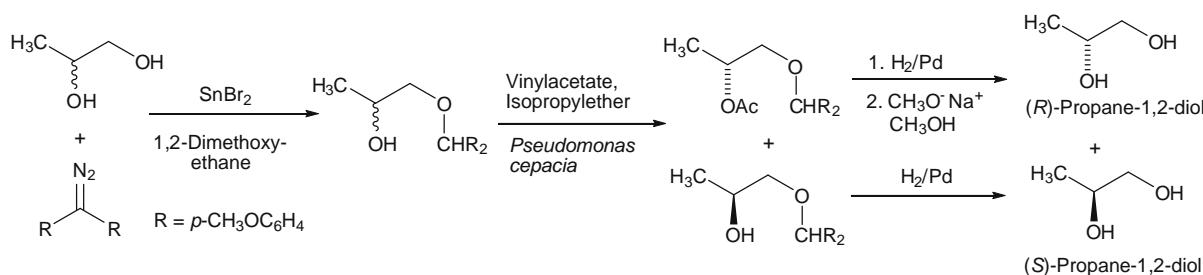
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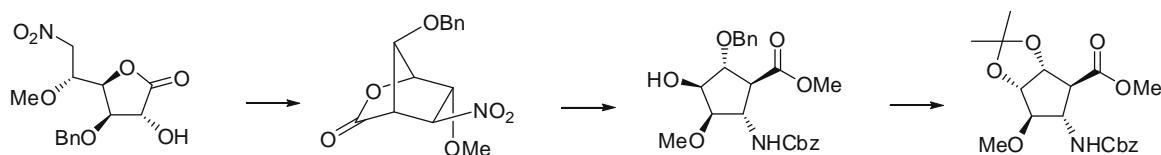
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Stereocontrolled transformation of nitrohexofuranoses into cyclopentylamines via 2-oxabicyclo[2.2.1]heptanes. IV: Synthesis of enantiopure methyl (1*S*,2*R*,3*R*,4*R*,5*S*)-5-benzoyloxycarbonylamino-2,3-isopropylidenedioxy-4-methoxycyclopentanecarboxylate

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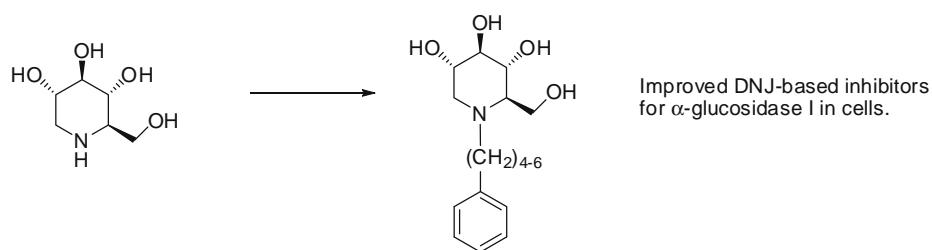
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Improved cellular inhibitors for glycoprotein processing α -glucosidases: biological characterisation of alkyl- and arylalkyl-N-substituted deoxynojirimycins

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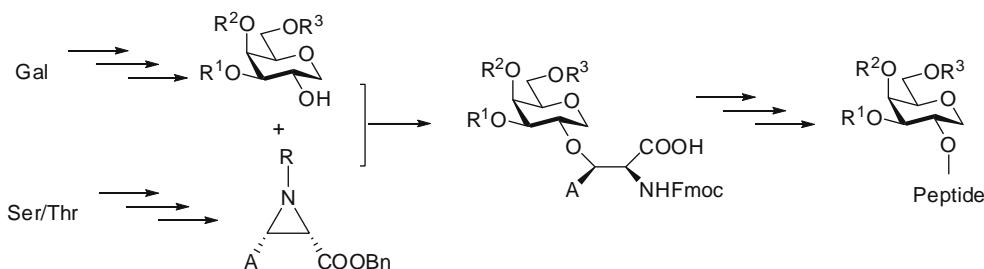
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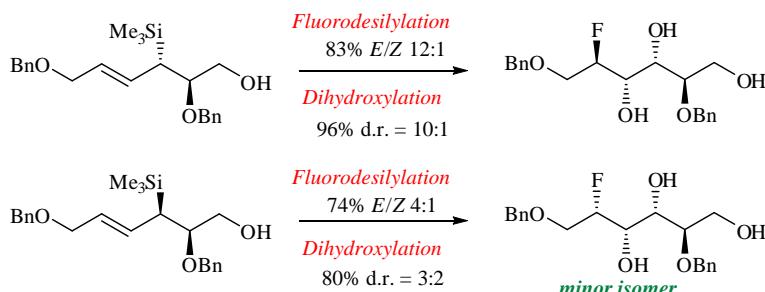
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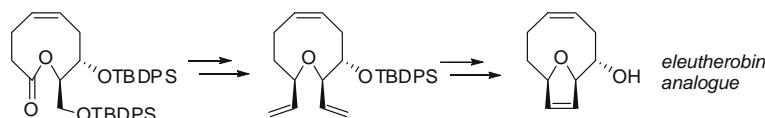
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Synthesis of simplified analogues of eleutherobin via a Claisen rearrangement/RCM strategy

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S.Y. Frankie Mak, Gary C.H. Chiang, James E.P. Davidson, John E. Davies, Andrew Ayscough, Gilles Pain, Jonathan W. Burton *, Andrew B. Holmes *

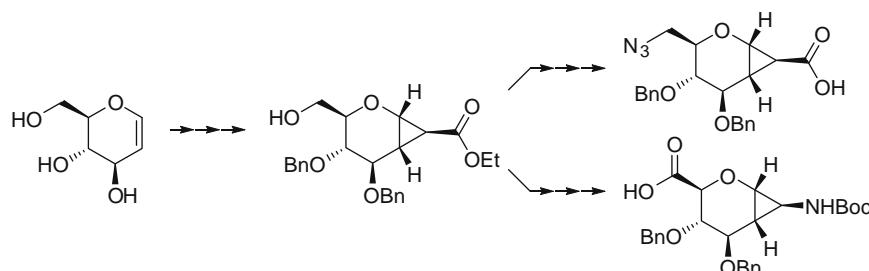


The enantioselective synthesis of a number of simplified analogues of the cytotoxic natural product eleutherobin is reported.

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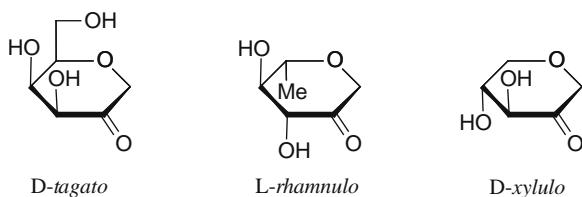
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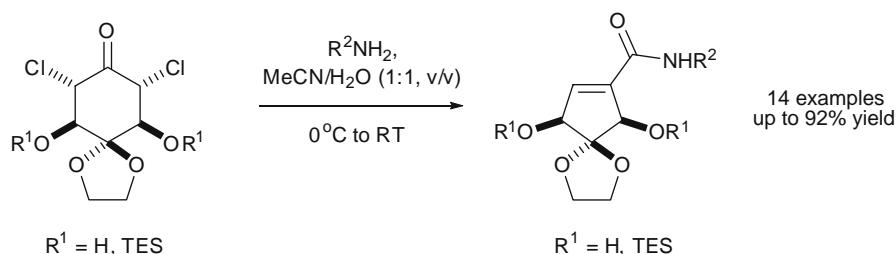
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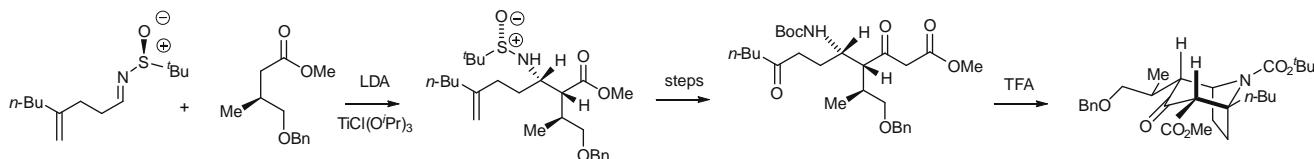
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Eric J. Thomas *, Clare F. Vickers



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*Corresponding author

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