

GRAPHICAL ABSTRACTS

Tetrahedron Lett. 27, 5787 (1986)

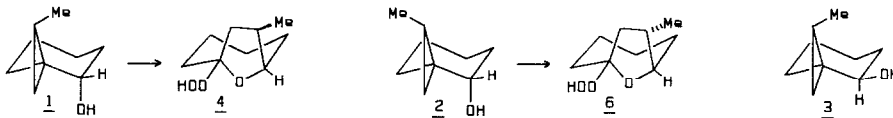
SOLVOLYTIC HYDROPEROXIDE REARRANGEMENTS III.

Stereoselective Rearrangements of Methylated Cyclopropyl Carbinols.

Robert C. Ronald* and Thomas S. Lillie

Department of Chemistry, Washington State University, Pullman, WA 99164 USA

The solvolysis of 1-3 in acidified THF-90% H₂O, through a combination of cyclopropylcarbinyl and Criegee rearrangements constitutes a novel stereoselective synthesis of 3-alkyl-4-hydroxycyclooctanone derivatives.



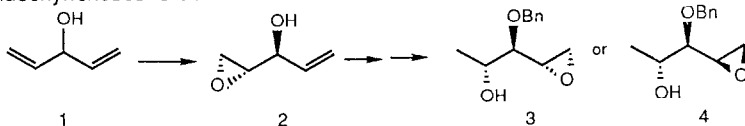
Tetrahedron Lett. 27, 5791 (1986)

ASYMMETRIC EPOXIDATION OF DIVINYL CARBINOL: A NEW APPROACH TO THE SYNTHESIS OF 2,6-DIDEOXYHEXOSE

Robert E. Babine

Infectious and Neoplastic Disease Research Section American Cyanamid Company Medical Research Division Lederle Laboratories Pearl River, New York 10965

Sharpless epoxidation of divinyl carbinol (1) gives 2 as the major product. Of special note is the ability to transform 2 selectively into either 3 or 4. The use of these reactions in the synthesis of homochiral 2,6-dideoxyhexoses is described.



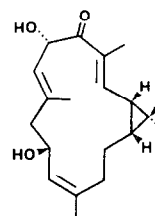
Tetrahedron Lett. 27, 5795 (1986)

AGROSTISTACHIN, A NOVEL MACROCYCLIC DITERPENE FROM AGRO-STISTACHYS HOOKERI

Y.-H. Choi, J. Kim, J.M. Pezzuto, A.D. Kinghorn*, and N.R. Farnsworth, Program for Collaborative Research in the Pharmaceutical Sciences, College of Pharmacy, Chicago, IL 60612

and H. Lotter* and H. Wagner, Institut für Pharmazeutische Biologie der Universität München, D-8000 München, W. Germany

The structure of agrostistachin, a novel cytotoxic agent from *Agrostistachys hookeri*, was determined as I.



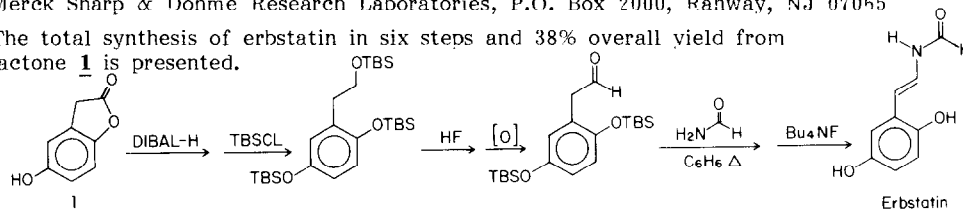
Tetrahedron Lett. 27, 5799 (1986)

TOTAL SYNTHESIS OF ERBSTATIN

David G. Hangauer

Merck Sharp & Dohme Research Laboratories, P.O. Box 2000, Rahway, NJ 07065

The total synthesis of erbstatin in six steps and 38% overall yield from lactone 1 is presented.

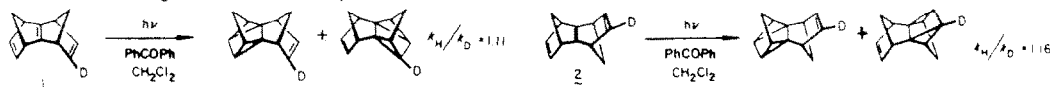


KINETIC DEUTERIUM ISOTOPE EFFECTS OPERATIVE DURING TRIPLET-SENSITIZED [2+2] PHOTOCYCLIZATION OF *syn*- AND *anti*-SESQUINORBORNATRIENES

Tetrahedron Lett. 27, 5803 (1986)

Leo A. Paquette,* Hermann Künzer, and Liladhar Waykole
Department of Chemistry, The Ohio State University, Columbus, Ohio 43210 USA

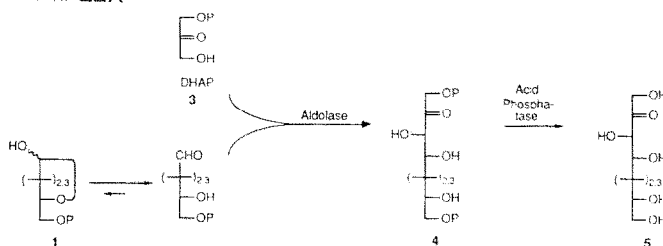
Product distributions in the triplet-sensitized photorearrangements of 1 and 2 are shown to be contrary to ground state expectations.



ADLOLASE-CATALYZED SYNTHESIS OF COMPLEX C₈ AND C₉ MONOSACCHARIDES

Mark D. Bednarski, Herbert J. Waldmann,
and George M. Whitesides*
Department of Chemistry
Harvard University
Cambridge, MA, 02138

Tetrahedron Lett. 27, 5807 (1986)



REGIOSPECIFIC SYNTHESIS OF α -ARYLSULFONOXY KETONES FROM KETONE DERIVATIVES

Robert V. Hoffman* and C. Sean Carr
Department of Chemistry, New Mexico State University, Las Cruces, NM 88003

Tetrahedron Lett. 27, 5811 (1986)

Isomeric enol ester, enamine, and silyl enol ether derivatives of ketones are converted regiospecifically to α -arylsulfonyl ketones.



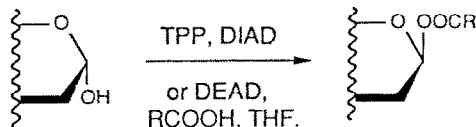
AN EFFICIENT SYNTHESIS OF GLYCOSYL ESTERS EXPLOITING THE MITSUNOBU REACTION.

Amos B. Smith III*, Karl J. Hale and Ralph A. Rivero.

Department of Chemistry, The Laboratory for Research on the Structure of Matter and The Monell Chemical Senses Center, University of Pennsylvania, Philadelphia, Pennsylvania 19104.

Tetrahedron Lett. 27, 5813 (1986)

The anomeric hydroxyl group of pyranose hemiacetals can be esterified with inversion of configuration using the Mitsunobu protocol.

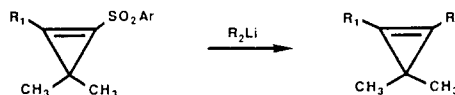


Tetrahedron Lett. 27,5817 (1986)

NUCLEOPHILIC SUBSTITUTION REACTIONS OF
1-SULFONYL SUBSTITUTED CYCLOPROPENES WITH
ALKYL LITHIUM REAGENTS

Albert Padwa* and M. Woods Wannamaker
Department of Chemistry, Emory University, Atlanta, GA 30322 USA

Sulfonyl substituted cyclopropenes react with
a variety of alkyl lithium reagents to give
disubstituted cyclopropenes in good yield.



ATOM TRANSFER CYCLIZATION OF SIMPLE HEXENYL IODIDES.
A CAUTION ON THE USE OF ALKENYL IODIDES AS PROBES FOR
THE DETECTION OF SINGLE ELECTRON TRANSFER PROCESSES

Tetrahedron Lett. 27,5821 (1986)

Dennis P. Curran*¹ and Dooseop Kim, Department of Chemistry, University of Pittsburgh, Pittsburgh, PA 15260

The trialkyltin initiated isomerization of 1-iodo-5-hexene to (iodomethyl)cyclopentane is reported. Rapid and reversible iodine atom transfer between alkyl radicals is the key chain transfer step. The results suggest an important caveat in the use of hexenyl iodides as free radical probes: a good mechanism exists whereby the yield of rearranged products may be much greater than the amount of radicals produced by the reaction under study!

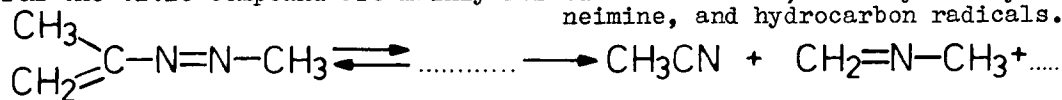


MATRIX ISOLATION AND IR SPECTROSCOPIC STUDY OF
PYROLYSIS PRODUCTS OF 2-METHYL-3,4-DIAZAPENTA-
1,3-DIENE. CLEAVAGE OF N=N-BOND IN UNSATURATED AZO COMPOUNDS.

Tetrahedron Lett. 27,5825 (1986)

U. Ziegler, G. Zimmermann, B. Ondruschka, V.V. Volkova, I.E. Gusel'nikov,
V.I. Smirnov, N.S. Nametkin

From the title compound are mainly formed acetonitrile, N-methyl methyleneimine, and hydrocarbon radicals.

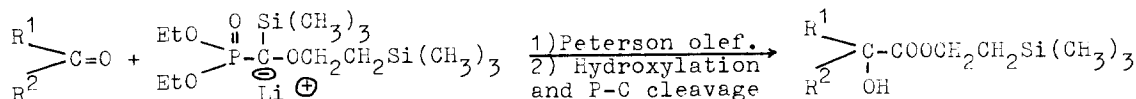


A NEW PROCEDURE FOR HOMOLOGATION OF CARBONYL
COMPOUNDS TO α -HYDROXY-CARBOXYLIC ESTERS BY
MEANS OF DIETHYL-[TRIMETHYLSILYLETHOXYMETHYL]PHOSPHONATE

Tetrahedron Lett. 27,5829 (1986)

Josef Binder and Erich Zbiral*

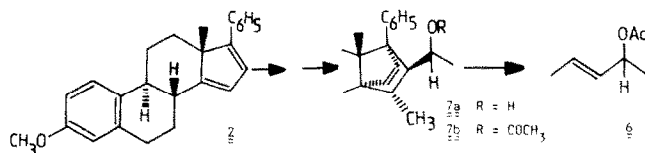
Institut für Organ.Chemie d.Univ.Wien,A-1090 Wien, Währinger Straße 38,Austria



DIENES AS CHIRAL TEMPLATES

Dietmar Schomburg⁺, Magion Thielmann⁺⁺, and Ekkehard Winterfeldt⁺⁺⁺; + GBF Gesellschaft für Biotechnologische Forschung mbH, Mascheroder Weg 1, D-3300 Braunschweig-Stückheim. ++ Institut für Organische Chemie der Universität Hannover, Schneiderberg 1 B, D-3000 Hannover 1

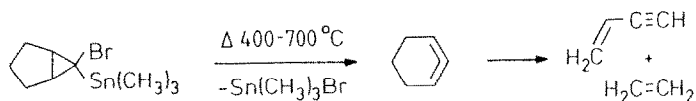
Stereoselective and regioselective cycloadditions to the chiral cyclopentadiene **2** are reported. Diastereoselective transformations of the adducts followed by a thermal retro reaction yields pure enantiomers.



Tetrahedron Lett. 27, 5833 (1986)

6-Bromo-6-(trimethylstannyl)bicyclo[3.1.0]hexane as a Thermal Precursor of 1,2,-Cyclohexadiene

Andreas Runge und Wolfram Sander*
Organisch-Chemisches Institut der Universität, D-6900 Heidelberg



Tetrahedron Lett. 27, 5835 (1986)

THERMAL REARRANGEMENT OF SOME [1.1.1]PROPELLANES

Johannes Belzner and Günter Szeimies
Institut für Organische Chemie der Universität München, Karlstr. 23, D-8000 München 2



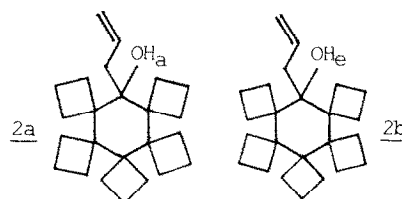
Tetrahedron Lett. 27, 5839 (1986)

CONFORMATIONAL ISOMERISM IN A FULLY SUBSTITUTED CYCLOHEXANE

Detlef Wehle and Lutz Fitjer*
Institut für Organische Chemie der Universität Göttingen,
Tammannstr. 2, D-3400 Göttingen, Germany

Conformational isomerism has been observed with **2a** and **2b**. Due to the highest barriers of inversion known so far [$\Delta G_{413}^\ddagger = 134.9$ kJ/mol (**2a**) and 136.9 kJ/mol (**2b**)] both conformers are indefinitely stable at room temperature in solution.

Tetrahedron Lett. 27, 5843 (1986)



LASER INDUCED SYNTHESIS:

CONDENSATION OF ACRYLONITRILE ON MESITYL OXIDE

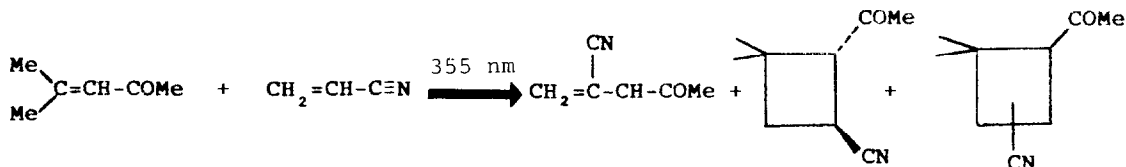
F.Lemaire, R.Stringat, R.Fellous & C.Bertaina

Laboratoire d'Optique Atomique et Moléculaire

Université de Nice, Parc Valrose 06034 NICE Cedex France

This photoaddition is realized with high quantum yield (15%) using the second harmonic of a pulsed Nd-YAG laser (intensity= 9 MW/cm²)

Tetrahedron Lett. 27,5847 (1986)



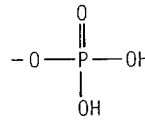
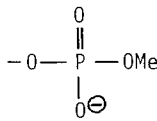
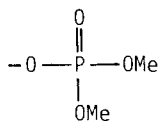
THE TAFT σ_I CONSTANTS OF SOME PHOSPHORYLOXY SUBSTITUENTS.

Marc Julia and Jean-Maurice Mallet

Laboratoire de Chimie, Ecole Normale Supérieure, 24, rue Lhomond, 75231 PARIS CEDEX 05 - FRANCE

Tetrahedron Lett. 27,5851 (1986)

The Taft constants of the following groups were determined by ¹⁹F NMR spectroscopy.



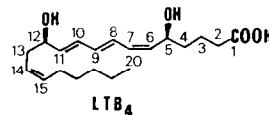
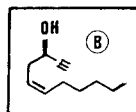
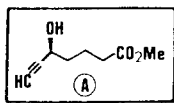
Tetrahedron Lett. 27,5853 (1986)

Optically Active Propargylic Alcohols from D-Xylose. Useful Precursors for LTB₄ Synthesis

P. Pianetti, P. Rollin and J.R. Pougny*

Laboratoire de Biochimie Structurale, UA 499, UFR de Sciences, 45067 Orléans Cédex 2, France

An efficient synthesis of the optically active propargylic alcohols **A** and **B**, chiral precursors for the LTB₄ synthesis, has been achieved. The main transformation uses the homologation of the α -acyloxy aldehyde to the corresponding α -acyloxy alkyne.



STEREOSELECTIVE SYNTHESIS OF POLYENIC ALCOHOLS.

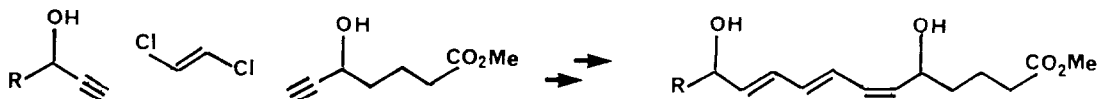
A NEW ROUTE TO THE LEUKOTRIENES B.

D. Guillerm and G. Linstrumelle

Laboratoire de Chimie, Ecole Normale Supérieure, 24 Rue Lhomond, 75231 Paris Cedex 05, France

A stereoselective synthesis of 14,15 dihydro LTB₄ and analogues is described.

Tetrahedron Lett. 27,5857 (1986)

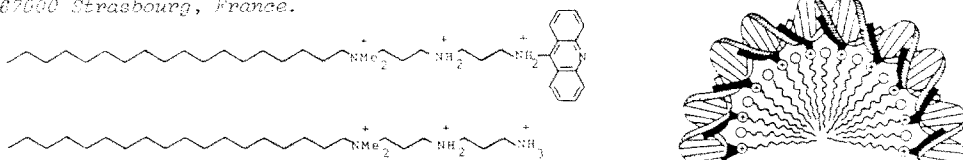


DNA STRONGLY BINDS TO MICELLES AND VESICLES CONTAINING LIPOLYAMINES OR LIPOINTERCALANTS.

Tetrahedron Lett. 27, 5861 (1986)

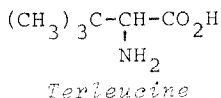
Jean-Paul Behr

Institut Le Bel, UA 422 CNRS, Université Louis Pasteur, 4, rue Blaise Pascal, 67000 Strasbourg, France.



SIMPLE OPTICAL RESOLUTION OF TERLEUCINE
Joëlle Viret, Heiko Patzelt and André Collet
Chimie des Interactions Moléculaires,
Collège de France, 75005 Paris, France

Tetrahedron Lett. 27, 5865 (1986)



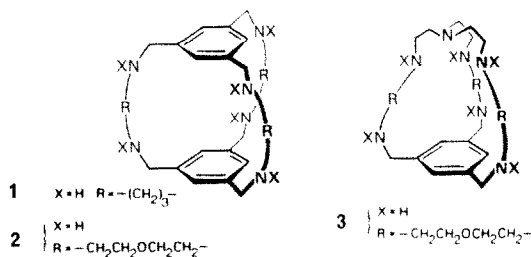
Terleucine can be easily resolved into L- and D-enantiomers by crystallization of its diastereoisomeric 10-camphorsulfonate salts.

ANION COORDINATION CHEMISTRY - SYNTHESIS AND ANION BINDING FEATURES OF CYCLOPHANE TYPE MACROBICYCLIC ANION RECEPTOR MOLECULES.

Tetrahedron Lett. 27, 5869 (1986)

Dennis Heyer and Jean-Marie Lehn*
Institut Le Bel, UA 422 CNRS,
4 rue Blaise Pascal,
67000 Strasbourg, France.

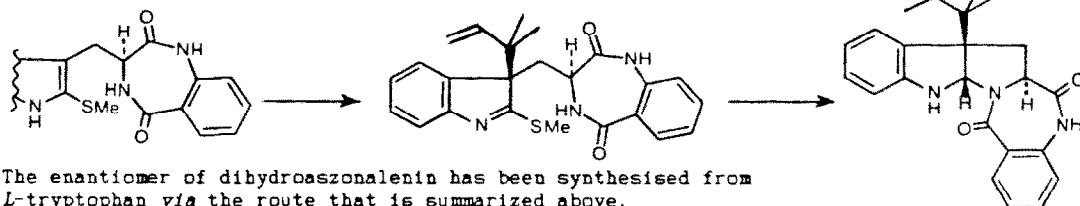
The three newly synthesized macrocyclic polyamines 1-3 strongly bind a variety of anions when protonated.



THE TOTAL SYNTHESIS OF (-)-DIHYDROASZONALENIN AND THE STEREOCHEMISTRY OF ASZONALENIN

Tetrahedron Lett. 27, 5873 (1986)

B. Bhat and D. M. Harrison, Dept. of Chemistry, University of Ulster.



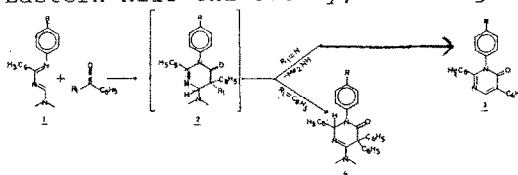
The enantiomer of dihydroaszonalenin has been synthesised from L-tryptophan via the route that is summarized above.

DIELS-ALDER CYCLOADDITION REACTION OF 1,3-DIAZABUTADIENES WITH KETENES

Tetrahedron Lett. 27,5875 (1986)

Sujit Nandy Mazumdar, Ibrahim Ibnusaud and Mohinder Pal Mahajan*
Department of Chemistry, North-Eastern Hill University, Shillong 793 003
Meghalaya, INDIA.

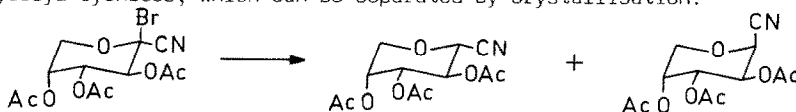
First report of (4+2)
cycloaddition of acyclic
1,3-diazabutadienes with
ketenes.



PREPARATION OF 1,2-CIS-GLYCOSYL CYANIDES BY THE
STERESELECTIVE REDUCTION OF ACETYLATED 1-BROMO-D-GLYCOSYL CYANIDES

Tetrahedron Lett. 27,5877 (1986)

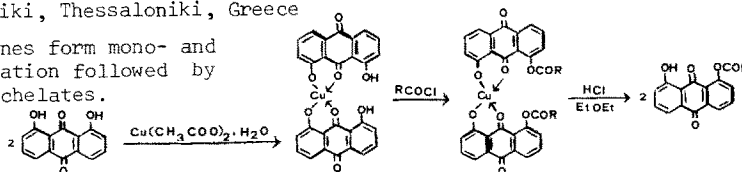
L. Somsák, Gy. Batta, and I. Farkas
Inst. Org. Chem., L. Kossuth Univ., Debrecen, P.O.B. 20. H-4010, Hungary
Stereoselective reduction of the title compounds gives epimeric mixtures of
glycosyl cyanides, which can be separated by crystallisation.



NOVEL METHOD FOR SELECTIVE ESTERIFICATION
OF POLYHYDROXY-ANTHRAQUINONES

Antonios S. Mellidis and Vassilios P. Papageorgiou*
Laboratory of Organic Chemistry, Department of Chemical Engineering,
University of Thessaloniki, Thessaloniki, Greece

Polyhydroxy-anthraquinones form mono- and
diesters via metal chelation followed by
acid hydrolysis of the chelates.



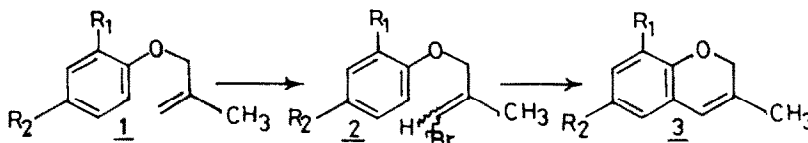
Tetrahedron Lett. 27,5881 (1986)

A FACILE SYNTHESIS OF 3-METHYL-2H-1-BENZOPYRANS VIA CLAISEN REARRANGEMENT

Tetrahedron Lett. 27,5883 (1986)

D.Gopal and K.Rajagopalan†, Department of Organic Chemistry, University
of Madras, Guindy Campus, Madras 600025, INDIA.

Several 3-methyl-2H-1-benzopyrans were prepared via Claisen rearrangement
using PEG-400 as a solvent in the presence of N,N-diethylaniline.

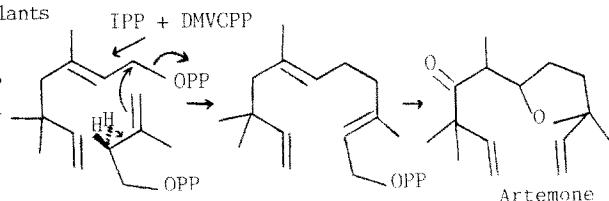


Tetrahedron Lett. 27, 5885 (1986)

A NOVEL BIOSYNTHESIS OF IRREGULAR SESQUITERPENE
ARTEMONE IN ARTEMISIA PALLENS

A.AKHILA, P.K.SHARMA and R.S.THAKUR
Central Institute of Medicinal and Aromatic Plants
Lucknow, India 226 016

A novel condensation between IPP and DMVCPP has been reported for the first time within the plant system to biosynthesise artemone, a sesquiterpene.



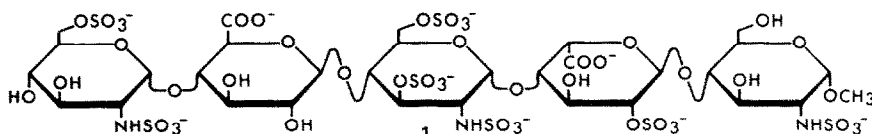
SYNTHESIS OF AN ANTITHROMBIN BINDING HEPARIN-LIKE
PENTASACCHARIDE LACKING 6-O SULPHATE AT ITS REDUCING END.

Tetrahedron Lett. 27, 5889 (1986)

T. Beetz and C.A.A. van Boeckel

Organon Scientific Development Group, P.O. Box 20, 5340 BH Oss, The Netherlands

Synthesis of pentasaccharide 1 of the structure of the antithrombin (AT-III) binding sequence, but lacking the 6-O sulphate group at the reducing end is described.



FREE RADICAL ANNULATION OF CYCLOPENTANE RING

Tetrahedron Lett. 27, 5893 (1986)

Živorad Čeković* and Radomir Sačić

Department of Chemistry, University of Belgrade, P.O. Box 550, 11001 Belgrade, Yugoslavia

Annulation of cyclopentane ring was achieved by combination of intermolecular conjugated homoallyl radical addition and intramolecular 5-nexenyl type radical cyclization

