

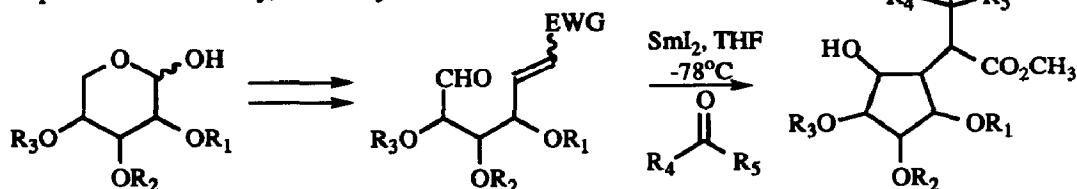
GRAPHICAL ABSTRACTS

SEQUENTIAL SmI₂-PROMOTED ONE- AND TWO-ELECTRON REACTIONS OF CARBOHYDRATES

Eric J. Enholm* and Antigone Trivellas

Department of Chemistry, University of Florida, Gainesville, FL 32611, USA

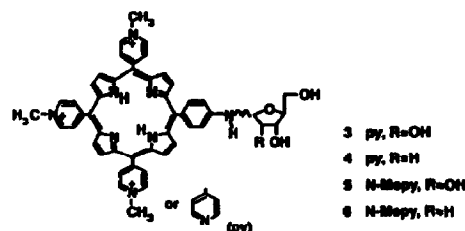
Tetrahedron Letters, 1994, 35, 1627



RIBOFURANOSIDES N-SUBSTITUTED WITH meso-PORPHYRIN AS NUCLEOSIDE-LIKE COMPOUNDS. Handong Li and Leszek Czuchajowski*, Department of Chemistry, University of Idaho, Moscow, ID 83844, USA

meso-Tri(4-pyridyl)porphyrin-p-phenylene-1'-amino-D-ribose and -D-2-deoxyribose were synthesized together with their N-methyl-4-pyridinium derivatives.

Tetrahedron Letters, 1994, 35, 1629



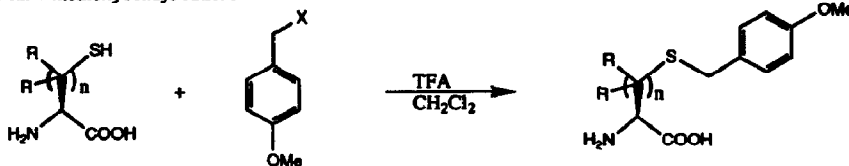
Two New Procedures for the Introduction of Benzyl-type Protecting Groups for Thiols.

Lutz S. Richter, James C. Marsters, Jr. and Thomas R. Gadek

Department of Bioorganic Chemistry, Genentech, Inc. 460 Point San Bruno Boulevard, South San Francisco, CA 94080

Tetrahedron Letters, 1994, 35, 1631

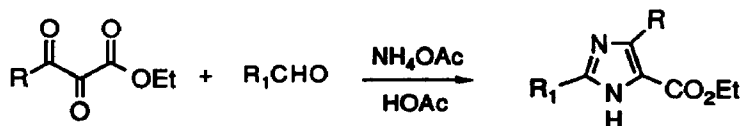
S-alkylations with 4-methoxybenzyl cations



An Efficient and Mild Synthesis of Highly Substituted Imidazoles

Marcus F. Brackeen,* Jeffrey A. Stafford, Paul L. Feldman, and Donald S. Karanewsky
Glaxo Research Institute, 5 Moore Drive, Research Triangle Park, NC 27709

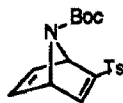
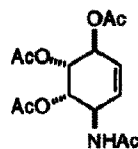
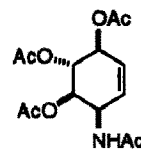
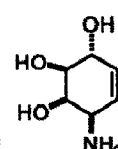
Tetrahedron Letters, 1994, 35, 1635



SYNTHESIS OF (±)-CONDURAMINES FROM

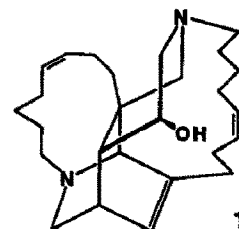
PYRROLE Regis Leung-Toung, Yanzhou Liu, Joseph M. Muchowski* and Yu-Lin Wu. Syntex Research, Institute of Organic Chemistry, 3401 Hillview Ave., Palo Alto, CA 94304.

The Diels-Alder product **1b**, of tosylacetylene and N-tert-Boc-pyrrole, was converted into (±)-conduramine C-1 (**22**) and the tetraacetates of (±)-conduramines A-1 (**9b**) and F-1 (**15b**).

**1b****9b****15b****22****INGENAMINE, A NOVEL PENTACYCLIC ALKALOID FROM THE MARINE SPONGE XESTOSPONGIA INGENS**

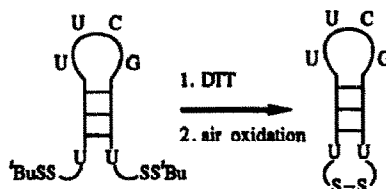
Fangming Kong and Raymond J. Andersen*, Departments of Chemistry and Oceanography, University of British Columbia, Vancouver, BC, CANADA V6T 1Z4
Theresa M. Allen, Department of Pharmacology, University of Alberta, Edmonton, Alberta, CANADA T6G 2H7.

Ingenamine (**1**), a novel cytotoxic pentacyclic alkaloid, has been isolated from the marine sponge *Xestospongia ingens* collected in Papua New Guinea. The structure of ingenamine has been solved by extensive spectroscopic analysis.

**1****SYNTHESIS OF A DISULFIDE STABILIZED RNA HAIRPIN**

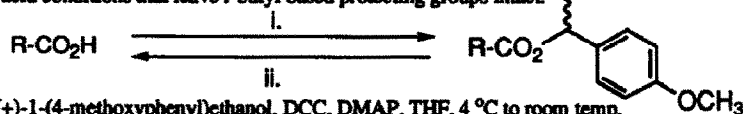
Jay T. Goodwin and Gary D. Glick*
Department of Chemistry, University of Michigan
Ann Arbor, Michigan, 48109-1055, U.S.A.

An N-3 ethylthiol-modified uridine has been synthesized and site-specifically incorporated through solid-phase phosphoramidite chemistry at the 5'- and 3'-termini of an RNA hairpin to provide increased conformational stability via a disulfide cross-link.

**THE (±)-1-(4-METHOXYPHENYL)ETHYL ESTER AS A CARBOXYL PROTECTING GROUP**

Michael S. Bernatowicz, Hann-Guang Chao, and Gary R. Matsueda, The Bristol-Myers Squibb Pharmaceutical Research Institute, P.O.Box 4000, Princeton, NJ 08543-4000 USA

(±)-1-(4-Methoxyphenyl)ethyl (Mpe) esters were prepared and shown to be quantitatively cleavable under mild acid conditions that leave *t*-butyl based protecting groups intact.



i. (±)-1-(4-methoxyphenyl)ethanol, DCC, DMAP, THF, 4 °C to room temp.

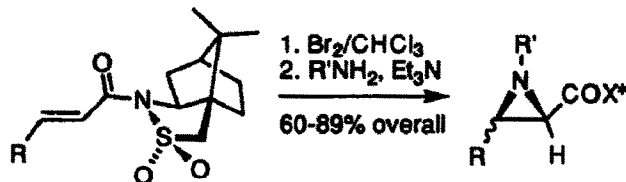
ii. 1 % (v/v) TFA (5 min) or 10 % (v/v) DCA (15 min) in CH₂Cl₂, room temp.

Auxiliary Mediated Synthesis of Aziridine-2-carboxylic Acid Derivatives

Philip Garner*, Ozdemir Dogan, and Satish Pillai

Department of Chemistry, Case Western Reserve University, Cleveland, OH 44106-7078

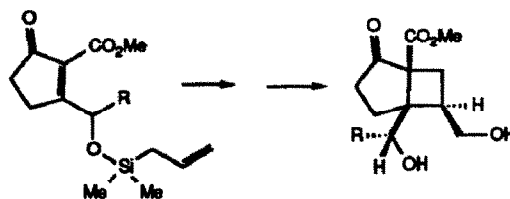
Synthetically useful aziridine-2-carboxylic acid derivatives can be prepared in a stereodefined manner from readily available α,β -unsaturated acyl camphor sultams via an asymmetric Gabriel-Cromwell reaction sequence.

**Siloxanes As Temporary Tethers in [2+2] Photocycloadditions**

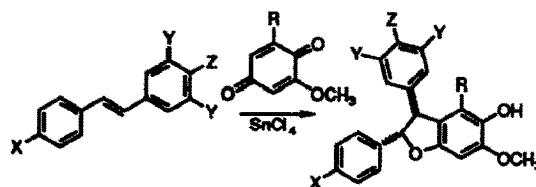
Michael T. Crimmins* and Lisa E. Guise

Department of Chemistry, University of North Carolina, Chapel Hill, NC 27599-3290

The use of siloxanes as temporary tethers in intramolecular [2+2] photocycloadditions has been investigated. High levels of regio and stereocontrol are achieved due to the geometric and steric constraints in the intramolecular transition state. Removal of the siloxane bridge provides the product of an *intermolecular* photocycloaddition with excellent regio and stereoselectivity.

**EVALUATION OF A SYNTHETIC ROUTE TO ϵ -VINIFERIN BASED ON A NEW METHOD FOR THE STEREoseLECTIVE PREPARATION OF 2,3-DIARYL-2,3-DIHYDROBENZOFURANS**Thomas A. Engler,* Bill W. Drancy and Gregory A. Gfesser
Department of Chemistry, University of Kansas, Lawrence, KS 66045-0046

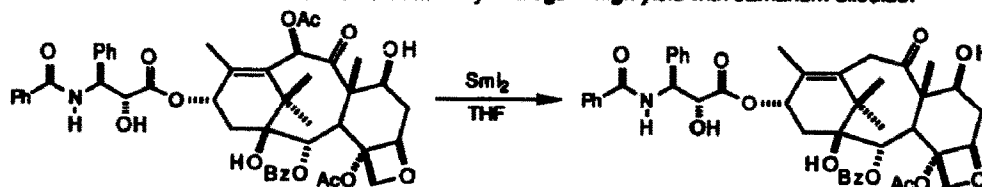
Lewis acid-promoted reactions of unsymmetrically substituted (*E*)-stilbenes with 2-methoxy-1,4-benzoquinones regio- and stereoselectively produce *trans* 2,3-diaryl-2,3-dihydrobenzofurans.

**A Simple Synthesis of 10-Deacetoxytaxol Derivatives**

Robert A. Holton,* Carmen Somoza, and Ki-Byung Chai,

Dittmer Laboratory of Chemistry, Florida State University, Tallahassee, Florida 32306 USA

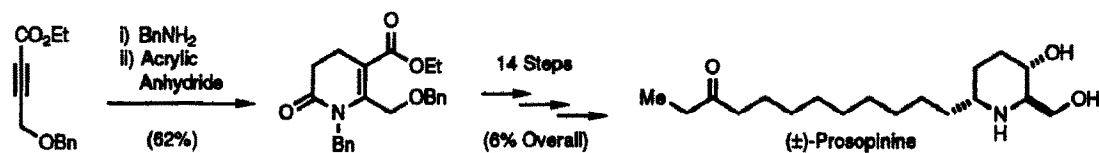
Taxol and baccatin III are reduced to the 10-deacetoxy analogs in high yield with samarium diiodide.



AZA-ANNULATION AS A ROUTE TO HYDROXYLATED ALKALOID LIPIDS. THE SYNTHESIS OF (±)-PROSOPININE.

Gregory R. Cook, Lars G. Beholz, and John R. Stille*

Department of Chemistry, Michigan State University, East Lansing, MI 48824-1322



REGIOSELECTIVE ENZYMIC DEACETYLATION OF SUCROSE OCTAACETATE IN ORGANIC SOLVENTS

David C. Palmer* and Fernand Terradas

McNeil Specialty Products Co., PO Box 800001, 1440 Olympic Drive, Athens, GA 30608-8001

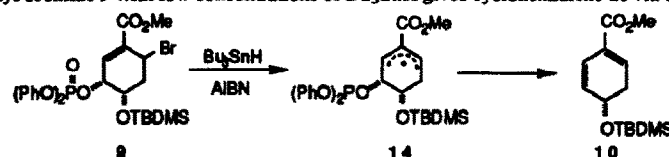
Sucrose octaacetate **1** can be regioselectively deacetylated using lipases and proteases in organic solvents to produce sucrose hepta-, hexa- and pentaacetates.



1,4-Radical Elimination in Cyclohexene Systems: A Model for the Chorismate Synthase Reaction

B. Giese*, N.G. Almstead, Institute of Organic Chemistry, University of Basel, 4056 Basel, Switzerland.

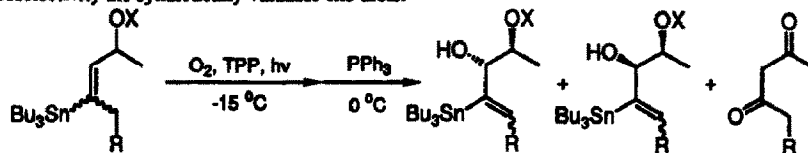
Treatment of cyclohexenyl bromide **9** with low concentrations of Bu₃SnH gives cyclohexadiene **10** via allylic radical **14** as intermediate.



Photooxygenation of γ -Hydroxy Vinylstannanes and their Acyl Derivatives: Mechanistic Insight into the Hydroxy-Directing Effect Waldemar Adam*

Olaf Gevert* and Peter Klug, Institute of Organic Chemistry, University of Würzburg, Am Hubland, D-97074 Würzburg, Germany.

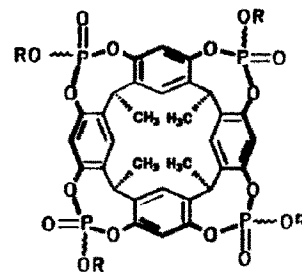
The Schenck reaction of ¹O₂ with γ -hydroxy vinylstannanes and their acyl derivatives, followed by Ph₃P reduction affords in high regio- and diastereoselectivity the synthetically valuable ene diols.



Synthesis and Configurational Analysis of Phosphorus Bridged Cavitanes. Tino Lippmann and Gerhard Mann*, Department of Chemistry, University of Leipzig, D-04103 Germany
 Enrico Dalcanale*, Department of Organic and Industrial Chemistry, University of Parma, I-43100 Italy

Tetrahedron Letters, 1994, 35, 1685

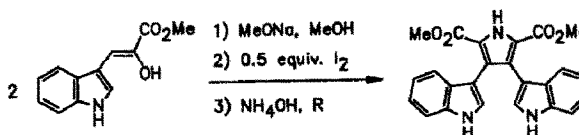
A novel family of diastereomeric cavitanes, obtained by incorporation of four phosphate groups on a resorcinol derived calix[4]arene has been studied.



Isolation and Synthesis of 3,4-Bis(indol-3-yl)pyrrole-2,5-dicarboxylic Acid Derivatives from the Slime Mould *Lycogala epidendrum*

Tetrahedron Letters, 1994, 35, 1689

R.Fröde, C. Hinze, I. Josten, B. Schmidt, B. Steffan and W. Steglich*
 Institut für Organische Chemie der Universität,
 Karlstr. 23, D-80333 München

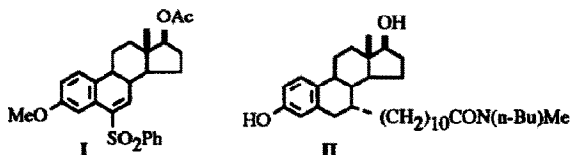


A one-pot synthesis of the title compounds is described.

A NEW, STEREOSELECTIVE APPROACH TO C(7)-ALKYLATED ESTRA-1,3,5(10)-TRIENE DERIVATIVES

Tetrahedron Letters, 1994, 35, 1691

H. Künzer,* M. Thiel, G. Sauer, and R. Wiechert
 Research Laboratories, Schering AG-Berlin,
 Müllerstr. 170-178,
 1000 Berlin 65,
 Germany



A study on the conjugate addition of simple organolithium reagents to I led to a short, stereoselective synthesis of the estrogen receptor antagonist II (ICI 164384).

INDOLOQUINONES, PART 2.

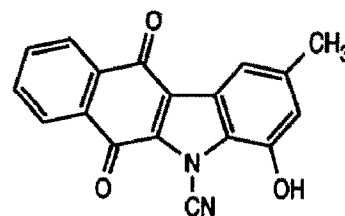
PALLADIUM-PROMOTED SYNTHESIS OF A 7-DEOXYPREKINAMYCIN ISOMER

Tetrahedron Letters, 1994, 35, 1695

H.-J. Knölker* and N. O' Sullivan

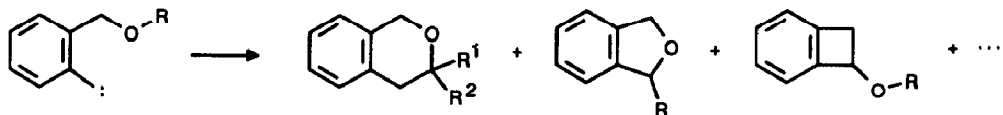
Institut für Organische Chemie, Universität Karlsruhe
 Richard-Willstätter-Allee, 76131 Karlsruhe, Germany

We have developed a four-step synthesis of a 7-deoxyprekinamycin isomer based on a palladium-promoted oxidative coupling as the key step.

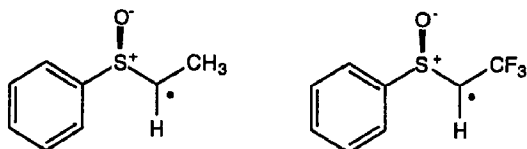


**INTRAMOLECULAR REACTIVITY OF ARYL CARBENES:
2-(ALKOXYMETHYL)PHENYL CARBENES.**

Wolfgang Kirmse* and Dirk Schnitzler, Fakultät für Chemie, Ruhr-Universität, 44780 Bochum, Germany
Stereochemistry and mechanism of the formal insertion of the title carbenes into C-H and C-O bonds of the side chain have been investigated.


STEREOCHEMICAL STUDY OF THE ALLYLATION OF 1-PHENYLSULFINYLETHYL AND 1-PHENYLSULFINYL-2,2,2-TRIFLUOROETHYL RADICALS.

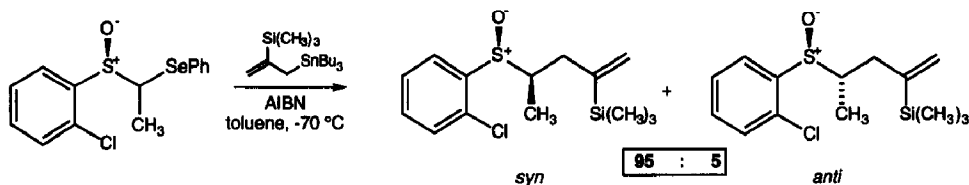
Philippe Renaud,*^{a)} Pierre-Alain Carrupt^{b)}, Michèle Gerster^{a)} and Kurt Schenk^{c)}, a) Université de Fribourg, Institut de Chimie Organique, Pérolles, CH-1700 Fribourg, Switzerland; b) Université de Lausanne, Ecole de Pharmacie (BEP), c) Université de Lausanne, Institut de Cristallographie (BSP), CH-1015 Lausanne-Dorigny, Switzerland.



Allylation reactions of the two title radicals produce preferentially compounds of opposite relative configuration in 61-64 % (methyl) and 81-82 % (trifluoromethyl) diastereoselectivity.

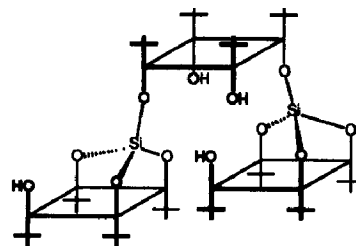
ORTHO-SUBSTITUTED ARYL SULFOXIDES DESIGNED FOR HIGHLY DIASTEREOSELECTIVE RADICAL REACTIONS.

Philippe Renaud,* and Thierry Bourquard, Université de Fribourg, Institut de Chimie Organique, Pérolles, CH-1700 Fribourg Switzerland.



MULTICAVITANDS III: SYNTHESIS AND NMR STUDIES OF A TRI-DIRECTIONAL KOILAND COMPOSED OF THREE *p*-TERT-BUTYLCALIX[4]ARENE UNITS FUSED BY TWO SILICON ATOMS. Xavier Delaigue, Mir Wais Hosseini,* Roland Graff, Jean-Pierre Kintzinger, Jesus Raya. Institut de Chimie, Université Louis Pasteur, 4, rue Blaise Pascal, F-67000 Strasbourg, France

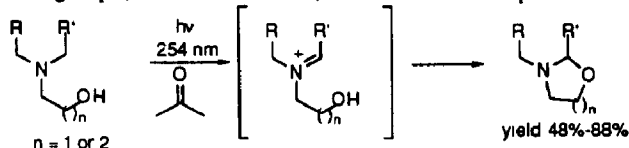
The high yield synthesis and NMR structural investigation of a divergent trireceptor obtained by fusion of three *p*-tert-butylcalix[4]arenes with two silicon atoms are described.



INTRAMOLECULAR QUENCHING OF IMINIUM IONS GENERATED BY PHOTOOXIDATION OF AMINOALCOHOLS WITH KETONES. A NEW SYNTHESIS OF OXAZINES AND OXAZOLES

Janine Cossy*, Madhumita Guha

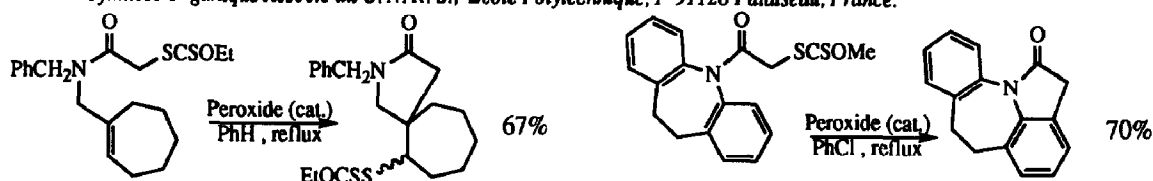
Laboratoire de Chimie Organique, Associé au C.N.R.S., E.S.P.C.I. 10 rue Vauquelin 75231 - PARIS Cédex 05 - France



A NEW RADICAL BASED SYNTHESIS OF LACTAMS AND INDOLONES FROM DITHIOCARBONATES (XANTHATES).

Jonathan Axon, Laurent Boiteau, Jean Boivin, Judith E. Forbes and Samir Z. Zard*

Laboratoire de Synthèse Organique Associé au C. N. R. S., Ecole Polytechnique, F-91128 Palaiseau, France.

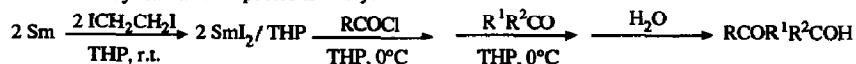


SAMARIUM DIODIDE IN TETRAHYDROPYRAN: PREPARATION

AND SOME REACTIONS IN ORGANIC CHEMISTRY. Jean-Louis Namy*,

Marielle Colomb and Henri B. Kagan*, Laboratoire de Synthèse Asymétrique (URA CNRS 1497), Institut de Chimie Moléculaire d'Orsay, Université Paris-Sud, 91405 Orsay Cedex, France.

The preparation of SmI_2 in tetrahydropyran (THP) is reported. In this solvent, reactions of SmI_2 with acid chlorides are cleaner than in THF. Formation of stable acyl samarium species is likely.

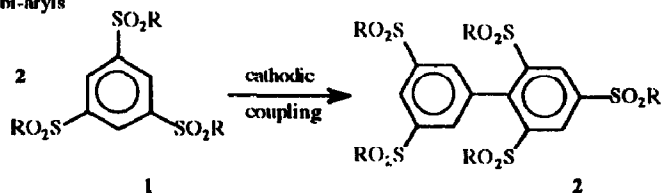


CATHODIC COUPLING OF 1, 3, 5 - TRI (ARYL AND ALKYL)SULFONYL BENZENES INTO TETRA AND PENTA-SUBSTITUTED BIARYLS

Mohamed Benaskar and Jacques Simonet *

Laboratoire d'Electrochimie, associé au CNRS, Université de Rennes I, Beaulieu, 35042 Rennes (France).

Trisulfones yield under non-aqueous conditions to unexpected bi-aryls



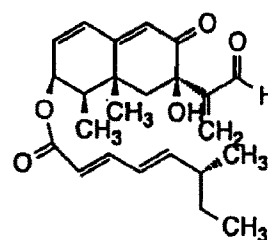
A BRASSINOLIDE-INHIBITOR KM-01, ITS ISOLATION AND STRUCTURE ELUCIDATION FROM A FUNGUS *DRECHSLERA AVENAE*.

Tetrahedron Letters, 1994, 35, 1731

Kim, S.K., Mizuno, K., Hatori, M. and Marumo, S.

Department of Agricultural Chemistry, Nagoya University, Chikusa-ku, Nagoya, Japan

A first brassinolide-inhibitor named KM-01 was isolated from a fungus, *Drechslera avenae*, and its structure with absolute stereochemistry determined as the fatty acid ester of bipolaroxin.

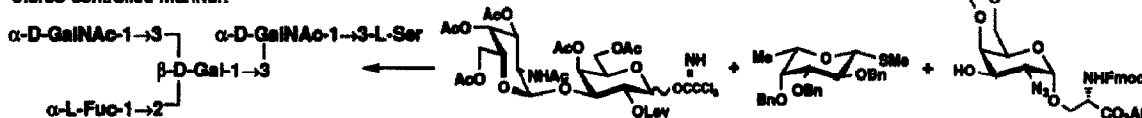


SYNTHESIS OF A GLYCOTETRAOSYL SERINE, A PARTIAL STRUCTURE OF AN OVARIAN CYST MUCIN GLYCOPROTEIN OF BLOOD GROUP A ACTIVITY

Tetrahedron Letters, 1994, 35, 1735

Wallace M. Macindoe^a, Hiroyuki Iijima^a, Yoshiaki Nakahara^a, and Tomoya Ogawa^{a,b}
^aThe Institute of Physical and Chemical Research (RIKEN), Wako-shi, Saitama, 351-01 Japan
^bFaculty of Agriculture, University of Tokyo, Yayoi, Bunkyo-ku, Tokyo, 113 Japan

A synthesis of a glycotetraosyl serine with blood group A determinant structure is described in a regio- and stereo-controlled manner.

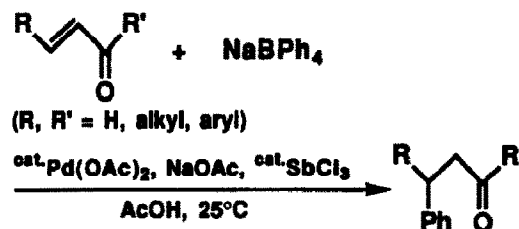


ANTIMONY(III) CHLORIDE AS AN EFFICIENT CATALYST FOR PALLADIUM-CATALYZED HYDROPHENYLATION OF α,β -UNSATURATED KETONES AND ALDEHYDES

Tetrahedron Letters, 1994, 35, 1739

Chan Sik Cho, Shin-ichi Motofusa, and Sakae Uemura^{*}
 Division of Energy and Hydrocarbon Chemistry, Graduate School of Engineering, Kyoto University, Sakyo-ku, Kyoto 606-01 (Japan)

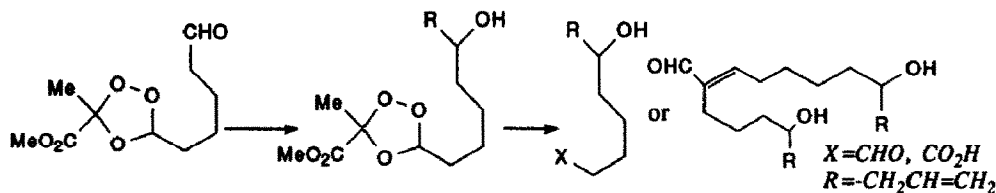
A remarkable catalytic effect of antimony(III) chloride was disclosed in palladium-catalyzed hydrophenylation of α,β -unsaturated ketones and aldehydes (Michael-type conjugate addition) with sodium tetraphenylborate in acetic acid at 25°C.



CHEMOSELECTIVE REACTIONS OF A COMPOUND CONTAINING BOTH ALDEHYDE AND OZONIDE GROUPS

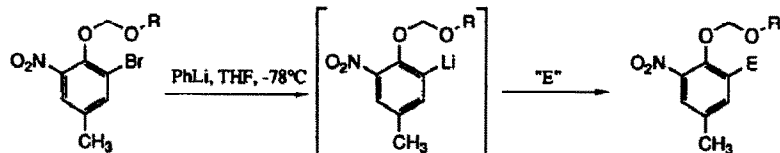
Tetrahedron Letters, 1994, 35, 1743

Yung-Son Hon^a and Jiann-Long Yan^b ^aInstitute of Chemistry, Academia Sinica, Nankang, Taipei, Taiwan 11529, R.O.C.
^bInstitute of Applied Chemistry, Chinese Culture University, Taipei, Taiwan, R.O.C.

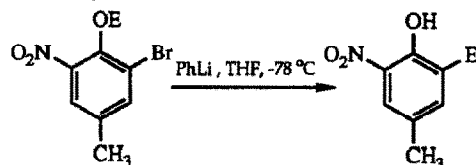


PREPARATION AND REACTIONS OF STABLE 2-LITHIO-6-NITROPHENOL DERIVATIVES. Ian R. Hardcastle, Peter Quayle* and E. Lucy M. Ward
Department of Chemistry, The Victoria University of Manchester, Manchester, M13 9PL.

2-Bromo-6-nitrophenyl MOM and SEM ethers form stable lithio compounds on treatment with phenyllithium.



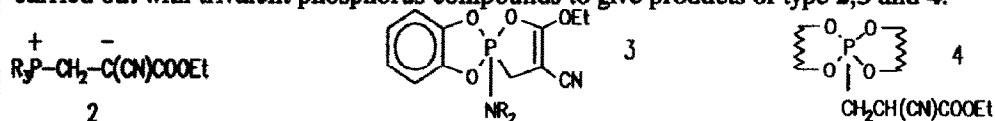
"Metallo-Fries" Rearrangements of 2-Lithio-6-nitrophenol Derivatives. I. R. Hardcastle and P. Quayle*. Department of Chemistry, The Victoria University of Manchester, Manchester M13 9PL. UK.



UNUSUAL TRANSFORMATIONS OF ETHYL-2-CYANOACRYLATE IN REACTIONS WITH TRIVALENT PHOSPHORUS COMPOUNDS

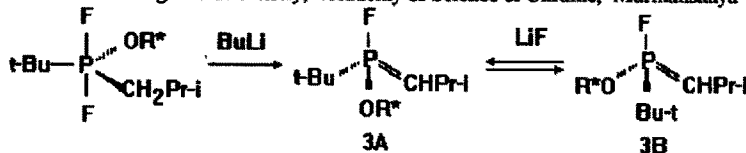
Yu.G.Gololobov, G.D.Kolomnikova, T.O.Krylova
A.N.Nesmeyanov Institute of Organo-Element Compounds RussAcad.ofSci.
Vavilov str.,28,Moscow V-334,117813 GSP-1 Russia

Various transformations of ethyl-2-cyanoacrylate - $\text{CH}_2=\text{C}(\text{CN})\text{COOEt}$ **1** was been carried out with trivalent phosphorus compounds to give products of type **2**, **3** and **4**.



FLUOROALKOXYPHOSPHONIUM YLIDS. EPIMERIZATION AND TRANSFORMATIONS. O.I. Kolodiazhnyi*, S.Ustenko, O.Golovaty.

Institute of Bioorganic Chemistry, Academy of Science of Ukraine, Murmanskaya Street, 1, KIEV - 253094 - UKRAINE

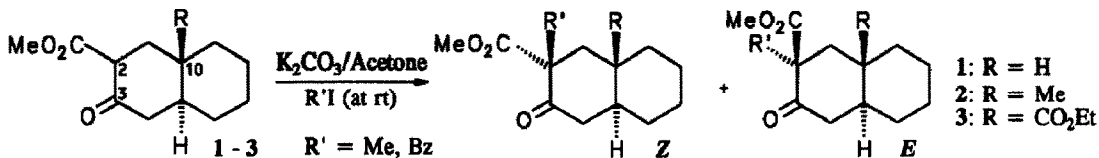


Dehydrofluorination of alkoxydifluorophosphoranes passes under conditions of kinetic control to afford diastereomers of phosphorus ylids **3A** and **3B**. However in the presence of lithium fluoride the most thermodynamic stable diastereomer of **3** is formed.

STEREOSELECTIONS IN CYCLIC β -KETOESTER ALKYLATIONS

Tetrahedron Letters, 1994, 35, 1759

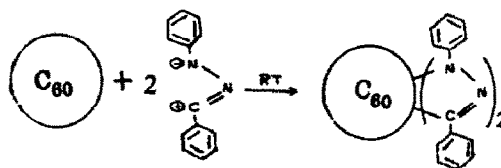
H. Surya Prakash Rao, K. Subba Reddy, Department of Chemistry, Pondicherry University, Pondicherry 605 014, India, and S.N. Balasubrahmanyam, Department of Organic Chemistry, Indian Institute of Science, Bangalore 560 012, India.
A hyperconjugative effect from the C-10 *syn*-axial ester in the β -ketoester 3 could contribute to the Z/E C-2 alkylation ratio being nearly the inverse of that in 2 and closely similar to that in 1.



REACTION OF BUCKMINSTERFULLERENE WITH 1,3-DIPHENYLNITRILIMINE: SYNTHESIS OF PYRAZOLINE DERIVATIVES OF FULLERENE.

Tetrahedron Letters, 1994, 35, 1763

S. Muthu and P. Maruthamuthu, Department of Energy, R. Ragunathan, Department of Organic Chemistry, University of Madras, A.C.College Campus, Madras 600 025, India. P.R. Vasudeva Rao and C.K. Mathews*, Chemical Group, IGCAR, Kalpakkam 603 102, India.



C₆₀ reacts with 1,3-diphenylisobutyl nitrilimine to form a diadduct

(R,S) 2-Fluoro (Chloro) -4'-Carboxy-Triphenyl Methanol. Novel Acid

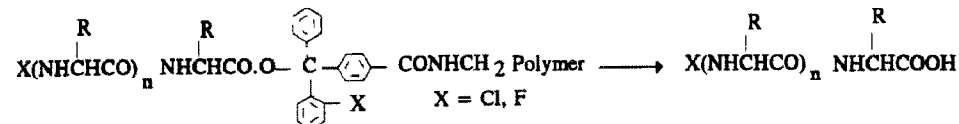
Tetrahedron Letters, 1994, 35, 1767

Labile Trityl Type Handles For Solid Phase Peptide Synthesis.

C. C. Zikos and N. G. Ferderigos.

Department of Chemistry, University of Athens, Panepistimioupolis, Zografou, 15771 Athens, Greece.

Synthesis of linkers for solid phase peptide synthesis which allows release by very mild acidic conditions of the product peptide.

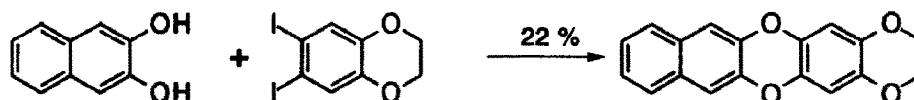


SYNTHESIS OF ANNULATED DIOXINS AND THEIR USE AS DONORS FOR CATION RADICAL SALTS.

Tetrahedron Letters, 1994, 35, 1769

Jonas Hellberg* and Margit E. Pelcman, Organic Chemistry, Royal Institute of Technology, S-100 44 Stockholm SWEDEN

The synthesis of a series of new alkoxyated annulated dioxins is described together with their cyclovoltammetric behaviour.



GRAPHICAL ABSTRACTS

Tetrahedron Letters, 1994, 35, 1773

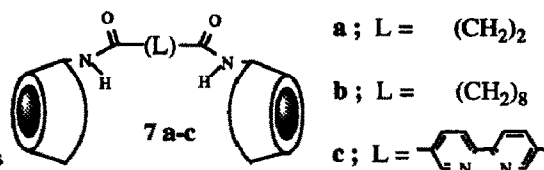
SYNTHESIS AND BINDING PROPERTIES OF NOVEL CYCLODEXTRIN DIMERS.

Fokke Venema,^a Chantal M. Baselier,^a Erik van Dienst,^b Bianca H.M. Ruël,^b Martinus C. Feiters,^a Johan F.J. Engbersen,^b David N. Reinhoudt,^b and Roeland J.M. Nolte.^{a*}

^a Department of Organic Chemistry, NSR-Center, University of Nijmegen, Toernooiveld, 6525 ED Nijmegen, The Netherlands

^b Laboratory of Organic Chemistry, University of Twente, P.O. Box 217, 7500 AE Enschede, The Netherlands.

The synthesis of cyclodextrin dimers **7a-c** and some binding constants for TNS are reported.

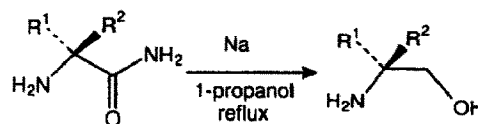


Tetrahedron Letters, 1994, 35, 1777

SYNTHESIS OF ENANTIOMERICALLY PURE 2,2-DISUBSTITUTED 2-AMINO-ETHANOLS BY DISSOLVING METAL REDUCTION OF α,α -DISUBSTITUTED AMINO ACID AMIDES.

Harold M. Moody, Bernard Kaptein, Quirinus B. Broxterman, Wilhelmus H. J. Boesten and Johan Kamphuis
 DSM Research, Bio-organic Chemistry Section, PO Box 18, 6160MD Geleen, The Netherlands

Enantiomerically pure 2,2-disubstituted-2-amino-ethanols are prepared in 65-99% yield by reduction of α,α -disubstituted amino acid amides using sodium in refluxing 1-propanol.

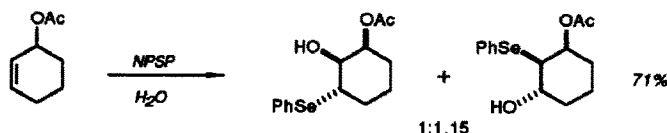


Tetrahedron Letters, 1994, 35, 1781

HYDROXYSELENATION OF ACETOXYCYCLOHEX-2-ENE

Alan F. Haughan, J.R. Knight and J.B. Sweeney,^{*}
 School of Chemistry, University of Bristol, Cantock's Close, Bristol BS8 1TS, UK

Addition of "HOSePh" to 1-acetoxycyclohex-2-ene is not regioselective, in contrast to the addition of PhSeCl



Tetrahedron Letters, 1994, 35, 1785

THE ASYMMETRIC SYNTHESIS OF ALLYLIC ALCOHOLS USING A RECOVERABLE CHIRAL SULPHOXIDE.

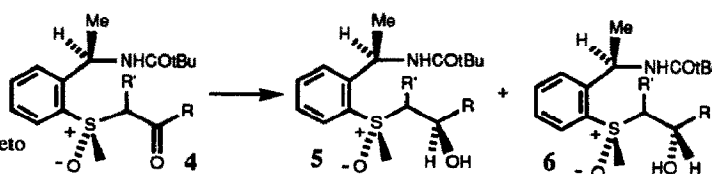
Ian D. Linney, Heather Tye and Martin Wills.^{*}

School of Chemistry, University of Bath,
 Claverton Down, Bath, BA2 7AY.

Roger J. Butlin

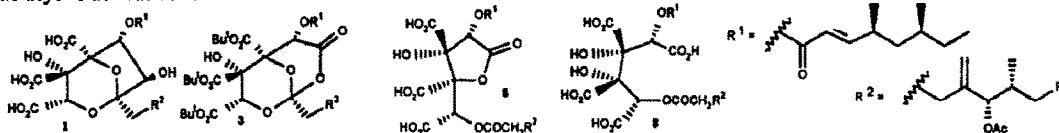
Zeneca Pharmaceuticals, Mereside,
 Alderley Park, Cheshire, SK10 4TG.

The synthesis and diastereoselective reductions of β -keto sulphoxides **4** are described.



The Squalostatins: Cleavage of the Bicyclic Core via the Novel 2, 8, 9-Trioxa-bicyclo[3.3.1]nonane Ring System. Daniele Andreotti, Panayiotis A. Procopiou*, Nigel S. Watson, *Department of Medicinal Chemistry, Glaxo Group Research Ltd., Greenford Road, Greenford, Middlesex UB6 0HE, United Kingdom.*

Squalestatin 1 was converted into the γ -lactone analogue 5, via hydrolysis of the novel ortho ester anhydride 3, and thence into the acyclic derivative 8.



σ -ASSISTANCE OF C_4-C_7 BOND IN THE SOLVOLYSIS OF 1-NORBORNYL TRIFLATES. A. García Martínez,* J. Osío Barcina, M. E. Rodríguez Herrero, M. Iglesias de Dios, E. Teso Vilar, L. R. Subramanian, *Departamento de Química Orgánica, Facultad de Ciencias Químicas, Universidad Complutense, Ciudad Universitaria, 28040 Madrid, Spain.*

The solvolysis of 4,7,7-trimethyl-1-norbornyl triflate proceeds under σ -participation of the C_4-C_7 bond, with formation of the σ bridged cation 22 as intermediate.



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