

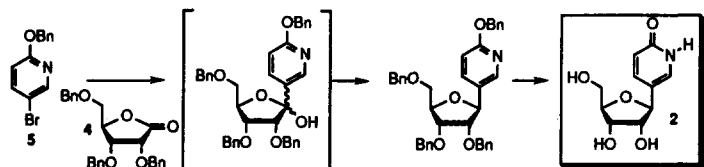
GRAPHICAL ABSTRACTS

SYNTHESIS OF 5-(β-D-RIBOFURANOSYL)-PYRIDIN-2-ONE: A "DELETION-MODIFIED" ANALOGUE OF URIDINE

Jasenska Matulic-Adamic and Leonid Beigelman,
Department of Organic Chemistry, Ribozyme Pharmaceuticals Inc., Boulder, CO 80301

Pyridine-2-one C-nucleoside **2** was prepared using several approaches. The most efficient pathway utilized condensation of 2,3,5-tri-*O*-benzyl D-ribo-1,4-lactone (**4**) and 2-(benzyloxy)-5-bromopyridine (**5**) followed by 1'-deoxygenation and removal of benzyl groups.

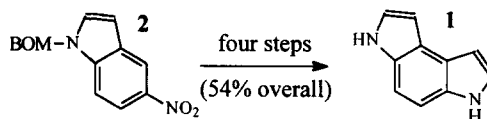
Tetrahedron Letters, 1997, 38, 1669



SYNTHESIS AND REACTIVITY OF PYRROLO[3,2-*E*]INDOLE: REMOVAL OF A N-BOM GROUP FROM AN UNACTIVATED INDOLE

John E. Macor^{*1}, James T. Forman, Ronald J. Post and Kevin Ryan, Department of Medicinal Chemistry
Central Research Division, Pfizer Inc, Groton, Connecticut 06340

A practical synthesis of pyrrolo[3,2-*e*]indole (**1**) is described. Removal of the BOM group was found to be problematic, but could be accomplished in moderate yield. Limited studies on the chemistry of **1** are also presented.

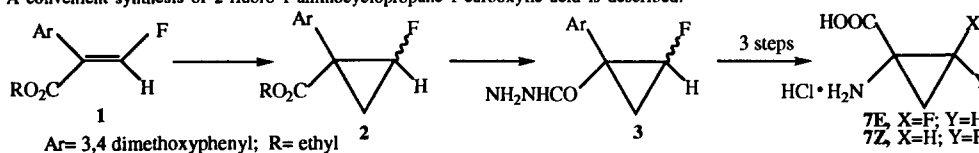


Tetrahedron Letters, 1997, 38, 1673

Synthesis of a Fluorinated Analog of 1-Aminocyclopropane Carboxylic Acid. Milton J. Sloan and Kenneth L. Kirk,*

Laboratory of Bioorganic Chemistry, National Institute of Diabetes and
Digestive and Kidney Diseases, National Institutes of Health, Bethesda, Maryland 20892 USA

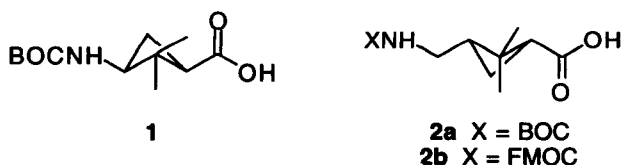
A convenient synthesis of 2-fluoro-1-aminocyclopropane-1-carboxylic acid is described.



Tetrahedron Letters, 1997, 38, 1677

CHIRAL 1,3-CYCLOBUTANE AMINO ACIDS: SYNTHESES AND EXTENDED CONFORMATIONS

Kevin Burgess*, Shiming Li, and Joe Rebenspies, Department of Chemistry, Texas A & M University
College Station, TX 77843 (email: burgess@chemvx.tamu.edu)

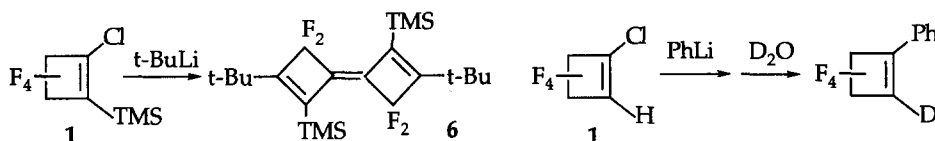


Amino acids **1** and **2** were prepared in optically active form. Crystallographic studies of derivatives of **1** show that these amino acids have extended conformations, and can pack with intermolecular hydrogen bonds in sheet-like arrays.

Tetrahedron Letters, 1997, 38, 1681

Reactions of Perfluoro-1-chloro-2-trimethylsilylcyclobutene.

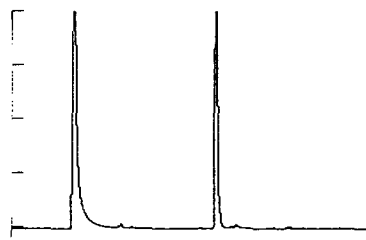
Kenneth B. Wiberg* and Manuel Marquez, Department of Chemistry,
Yale University, New Haven CT 06520-8107



CHEMILUMINESCENT NITROGEN DETECTION FOR HPLC: AN IMPORTANT NEW TOOL IN ORGANIC ANALYTICAL CHEMISTRY

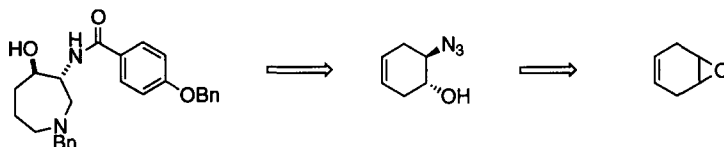
William L. Fitch* and A. Katrin Szardenings, Affymax Research Institute, 3410 Central Expressway,
Santa Clara CA 95051. Eugene M. Fujinari, Antek Instruments, 300 Bammel Westfield
Rd., Houston TX 77090

HPLC/CLND is a technique for quantifying yields and purities in organic chemistry, which will be especially useful in solid phase synthesis and combinatorial chemistry



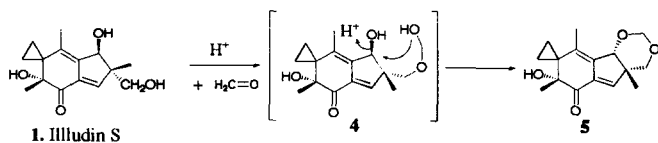
AN EFFICIENT FORMAL SYNTHESIS OF BALANOL VIA THE ASYMMETRIC EPOXIDE RING OPENING REACTION

Michael H. Wu and Eric N. Jacobsen*
Department of Chemistry and Chemical Biology
Harvard University, Cambridge, MA 02138, USA



AN ACETAL DERIVATIVE OF ILLUDIN S WITH IMPROVED ANTITUMOR ACTIVITY

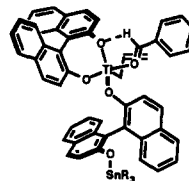
Trevor C. McMorris^{#*}, Jian Yu[#], Peter K. Gantzel[#], Leita A. Estes[^] and Michael J. Kelner[^]
[#]Department of Chemistry and Biochemistry, [^]Department of Pathology, University of California, San Diego,
La Jolla, CA 92093-0506



The Formyl C-H...O Hydrogen Bond As a Key to Transition-State Organization in Enantioselective Allylation, Aldol and Diels-Alder Reactions Catalyzed by Chiral Lewis Acids.

E. J. Corey,* David Barnes-Seeman and Thomas W. Lee
Department of Chemistry and Chemical Biology
Harvard University, Cambridge, Massachusetts 02138

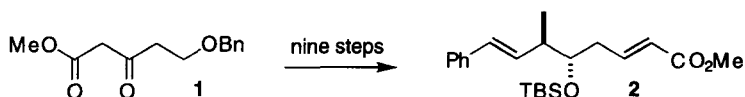
Tetrahedron Letters, 1997, 38, 1699



Keck catalytic allylation complex

Formal Syntheses of Cryptophycin A and Arenastatin A

Syed M. Ali and Gunda I. Georg, Department of Medicinal Chemistry, University of Kansas, Lawrence, KS 66045

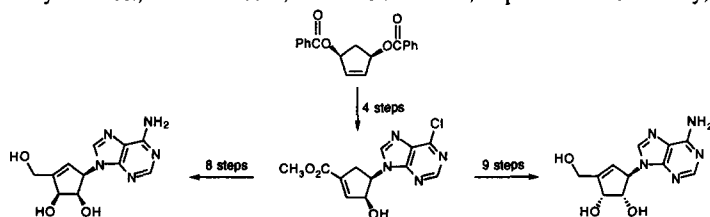


An efficient synthesis of the key intermediate **2** for the syntheses of cryptophycin A and arenastatin A from β -keto ester **1** is detailed. Asymmetry is introduced through an asymmetric reduction of β -keto ester **1**. Frater alkylation and subsequent functional group transformations provided the targeted methyl octadienoate **2**.

Tetrahedron Letters, 1997, 38, 1703

AN ENANTIO- AND DIASTEREOCONTROLLED SYNTHESIS OF (-)-NEPLANOCIN A AND ITS 2,3-DI-EPI ISOMER

Barry M. Trost, Robert Madsen, and Simon D. Guile, Department of Chemistry, Stanford University, Stanford, CA 94305



Tetrahedron Letters, 1997, 38, 1707

Catalysis of the Oxidation of Triphenylphosphine and of Trimethyl Phosphite by Hydrogen Peroxide in the Presence of Fe^{III} Compounds

Derek H. R. Barton,* David R. Hill and Bin Hu, Department of Chemistry, Texas A & M University, College Station, TX 77843-3255

The oxidations of triphenylphosphine and trimethyl phosphite to the corresponding oxides were studied in pyridine by ³¹P-NMR. They are faster in the presence of Fe^{III} compounds.

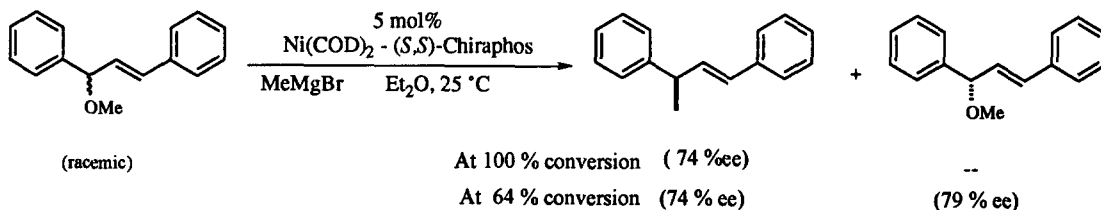


Tetrahedron Letters, 1997, 38, 1711

Tetrahedron Letters, 1997, 38, 1713

Nickel-Catalyzed Asymmetric Allylation of Alkyl Grignard Reagents.

Effect of Ligands, Leaving Groups and a Kinetic Resolution with a Hard Nucleophile. Nobuyoshi Nomura and T. V. RajanBabu,* Department of Chemistry, The Ohio State University, 100 W. 18th Ave., Columbus, Ohio 43210 USA



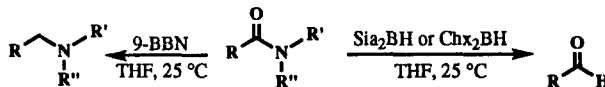
Tetrahedron Letters, 1997, 38, 1717

CONTROLLED REDUCTION OF TERTIARY AMIDES TO THE CORRESPONDING ALDEHYDES OR AMINES USING DIALKYLBORANES

Gayane Godjoian and Bakthan Singaram*

Department of Chemistry and Biochemistry, University of California, Santa Cruz, California 95064

Reduction of various tertiary amides with two equivalents of 9-BBN gave the corresponding tertiary amine. However, sterically more demanding dialkylboranes, such as dicyclohexylborane(Chx_2BH) and disiamylborane(Sia_2BH), reduced tertiary amides to the corresponding aldehydes.



Tetrahedron Letters, 1997, 38, 1721

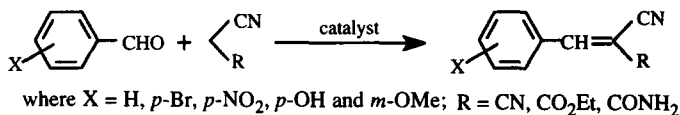
RARE-EARTH (RE) EXCHANGED NaY ZEOLITE PROMOTED KNOEVENAGEL CONDENSATION

T. Indrasena Reddy¹ and Rajender S. Varma^{*1, 2}

¹Department of Chemistry and ²Texas Regional Institute for Environmental Studies (TRIES)

Sam Houston State University, Huntsville, TX 77341-2117, U. S. A.

The condensation of aromatic aldehydes with active methylene compounds is promoted by rare-earth cation-exchanged NaY zeolites that affords olefinic products in moderate to good yields.



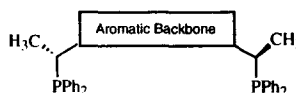
Tetrahedron Letters, 1997, 38, 1725

Synthesis of Chiral Phosphine Ligands with Aromatic Backbones and Their Applications in Asymmetric Catalysis

James M. Longmire, and Xumu Zhang*

Department of Chemistry, Pennsylvania State University, University Park, PA 16802

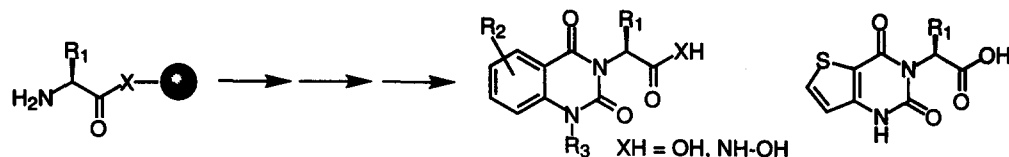
A general strategy for the synthesis of new chiral phosphine ligands has been discovered. A common feature of these ligands is that they contain rigid aromatic backbones which can be used to restrict conformational flexibility.



A General and Efficient Solid Phase Synthesis of Quinazoline-2,4-diones

Mikhail F. Gordeev,* Hon C. Hui, Eric M. Gordon, and Dinesh V. Patel
Versicor, Inc., 270 East Grand Ave., South San Francisco, CA 94080, U.S.A.

Tetrahedron Letters, 1997, 38, 1729

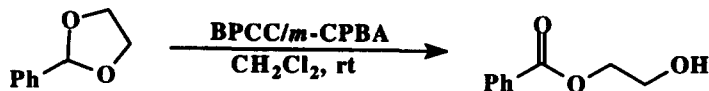


2,2'-BIPYRIDINIUM CHLOROCHROMATE/*m*-CHLOROPERBENZOIC ACID MEDIATED CLEAVAGE OF CYCLIC ACETALS TO ESTERS.

Frederick A. Luzzio* and Rhiana A. Bobb, Department of Chemistry, University of Louisville, Louisville, Kentucky 40292.

Tetrahedron Letters, 1997, 38, 1733

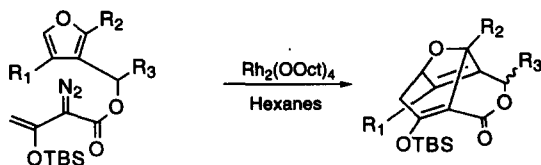
Benzylidene acetals are converted to hydroxyesters using a reagent system composed of 2,2'-bipyridinium chlorochromate and *m*-chloroperbenzoic acid.



TYPE II INTRAMOLECULAR ANNULATIONS BETWEEN VINYL CARBENOIDS AND FURANS

Huw M. L. Davies,* Rebecca Calvo and Gulzar Ahmed
Department of Chemistry, State University of New York at Buffalo, Buffalo, New York 14260-3000

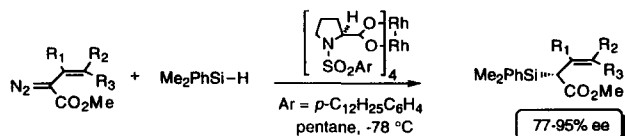
Tetrahedron Letters, 1997, 38, 1737



RHODIUM(II) (S)-N-(ARYLSULFONYL)PROLINATE CATALYZED ASYMMETRIC INSERTIONS OF VINYL- AND PHENYLACETENES INTO THE Si-H BOND

Huw M. L. Davies,*^a Tore Hansen,^a James Rutberg,^b and Paul R. Bruzinski^a
a) Department of Chemistry, State University of New York at Buffalo, Buffalo, NY 14260-3000
b) Department of Chemistry, Wake Forest University, Box 7486, Winston-Salem, NC 27109

Tetrahedron Letters, 1997, 38, 1741

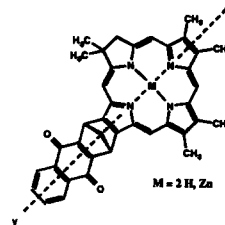


Tetrahedron Letters, 1997, 38, 1745

**Synthesis of a Novel Chlorin-Quinone System
for the Investigation of Light Induced Electron Transfer**

Yvonne Abel and Franz-Peter Montforts*, Institut für Organische Chemie, FB2,
Universität Bremen, Leobener Str. NW2, D-28359 Bremen.

In order to investigate the effect of the molecular symmetry on the light induced electron transfer in chlorin-quinone dyades we aimed at the synthesis of chlorin-quinone models **18a** and **18b** in which the quinone is situated at ring D adjacent to the reduced pyrrole ring A and thus orientated along the y-axis.

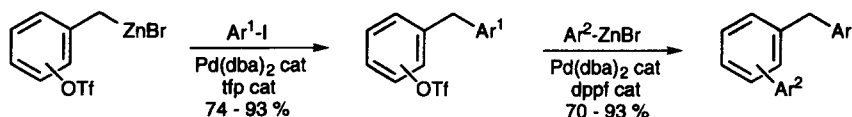


**New Multi-Coupling Benzylic Zinc Reagents for the Preparation
of Flexible Aromatic Compounds**

Mario Rottländer and Paul Knochel*

Fachbereich Chemie der Philipps-Universität Marburg, 35032 Marburg, Germany

Tetrahedron Letters, 1997, 38, 1749

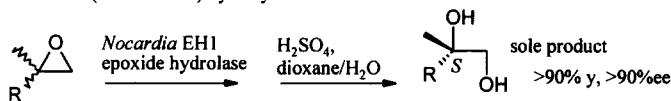


**DERACEMIZATION OF (±)-2,2-DISUBSTITUTED EPOXIDES VIA
ENANTIOCONVERGENT CHEMOENZYMATIC HYDROLYSIS USING
BACTERIAL EPOXIDE HYDROLASE AND SULFURIC ACID**

Romano V. A. Orru, Wolfgang Kroutil, Kurt Faber, Institute of Organic Chemistry, Graz University of Technology,
Stremayrgasse 16, A8010 Graz, Austria.

Complete deracemization of 2,2-disubstituted oxiranes was achieved by the combination of biocatalytic (*Nocardia* EH1) and acidic (sulfuric acid) hydrolysis.

Tetrahedron Letters, 1997, 38, 1753

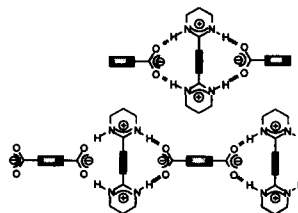


**Molecular Tectonics IV : Molecular Networks Based on Hydrogen
Bonding and Electrostatic Interactions**

Olivier Félix, Mir Wais Hosseini*, André De Cian, Jean Fischer
Institut de Chimie, Université Louis Pasteur, 4, rue Blaise Pascal,
F-67000 Strasbourg, France

Whereas diprotonated **1** (1,2-Bis(2'-tetrahydropyrimidyl)ethane) forms a discrete exobinuclear complex with a dihapto mode of hydrogen bonding with 4-methylbenzoate anion **2**⁻ in the solid state, with 4,4'-biphenyldicarboxylate **3**²⁻ an α -network composed of **1**-2H⁺ dication and **3**²⁻ dianion interconnected through strong hydrogen bonds and arranged in an alternating manner was obtained.

Tetrahedron Letters, 1997, 38, 1755



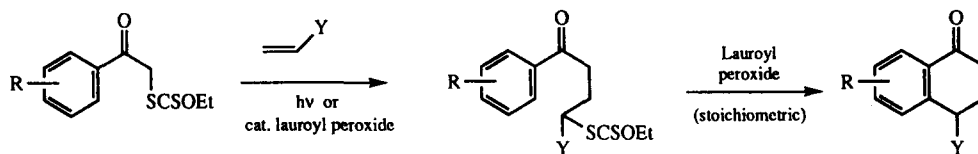
Tetrahedron Letters, 1997, 38, 1759

A NEW SYNTHESIS OF α -TETRALONES.

Annie Liard^a, Béatrice Quiclet-Sire^a, Radomir N. Saicic^a, and Samir Z. Zard^{a,b*}

a) Institut de Chimie des Substances Naturelles, 91198 Gif-Sur-Yvette, France.

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

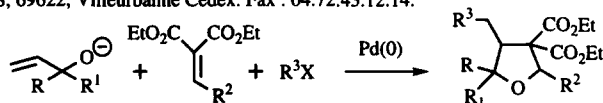


Tetrahedron Letters, 1997, 38, 1763

THREE PARTNERS FOR A ONE POT PALLADIUM-MEDIATED SYNTHESIS OF VARIOUS TETRAHYDROFURANS.

M. Cavicchioli, E. Sixdenier, A. Derrey, D. Bouysi, G. Balme*

Laboratoire de Chimie Organique 1, associé au CNRS, Université Claude Bernard, CPE, 43 Bd du 11 Novembre 1918, 69622, Villeurbanne Cédex. Fax : 04.72.43.12.14.



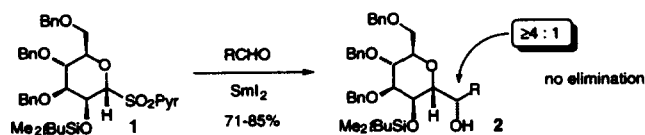
Tetrahedron Letters, 1997, 38, 1767

FURTHER STUDIES IN α -C-MANNOSYLATION PROMOTED

BY SAMARIUM DIODIDE. Olivier Jarreton, Troels Skrydstrup,*

and Jean-Marie Beau,* Laboratoire de Synthèse de Biomolécules associé au CNRS, Université de Paris-Sud, F-91405 Orsay Cedex (France).

The samarium diiodide-promoted coupling of mannosyl pyridylsulfone **1** with aldehydes afforded the C-glycoside **2** without any elimination.

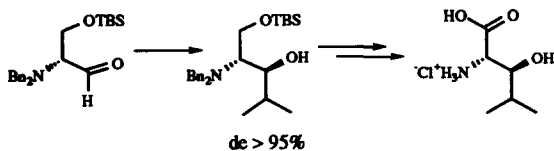


Tetrahedron Letters, 1997, 38, 1771

A HIGHLY STEREOSELECTIVE SYNTHESIS OF (2S, 3S)- β -HYDROXYLEUCINE

Taoues Laïb, Jacqueline Chastanet, Jieping Zhu*

Institut de Chimie des Substances Naturelles, CNRS, 91198 Gif-Sur-Yvette, France



Enantiomerically pure (2S, 3S)- β -hydroxy-leucine was synthesized featuring a diastereoselective nucleophilic addition of Grignard reagent to D-N,N-dibenzyl-O-TBS serinal as a key step.

Tetrahedron Letters, 1997, 38, 1773

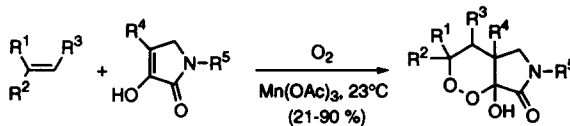
**Mn(III)-INDUCED MOLECULAR OXYGEN TRAPPING
REACTION OF ALKENES WITH 2,3-PYRROLIDINEDIONE
DERIVATIVES. A NOVEL ENTRY TO 1-HYDROXY-8-AZA-2,3-DIOXABICYCLO[4.3.0]NONAN-9-ONES**

Van-Ha Nguyen,[†] Hiroshi Nishino,* and Kazu Kurosawa

[†]Department of Environmental Science, Graduate School of Science and Technology, Kumamoto University

Department of Chemistry, Faculty of Science, Kumamoto University, Kurokami 2-39-1, Kumamoto 860, Japan

Manganese(III) oxidation of alkenes and 2,3-pyrrolidinediones with molecular oxygen gave bicyclic peroxides in good yield.

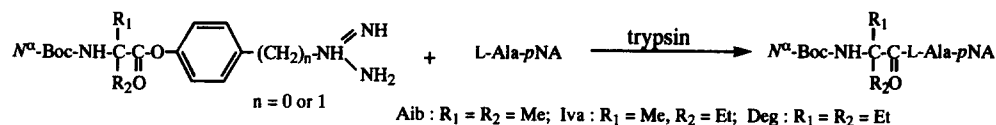


Tetrahedron Letters, 1997, 38, 1777

**ENZYMATIC COUPLING OF α,α -DIALKYL AMINO ACIDS
USING INVERSE SUBSTRATES AS ACYL DONORS¹**

Haruo Sekizaki, Kunihiko Itoh, Eiko Toyota and Kazutaka Tanizawa*, Faculty of Pharmaceutical Sciences, Health Sciences University of Hokkaido, Ishikari-Tobetsu, Hokkaido 061-02, Japan.

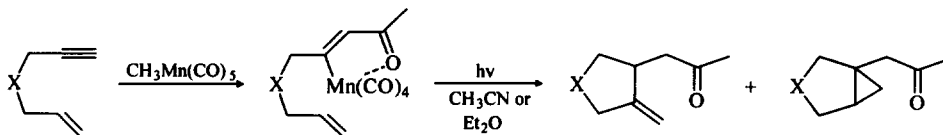
Streptomyces griseus trypsin was a more efficient catalyst than bovine trypsin.



Tetrahedron Letters, 1997, 38, 1781

**Synthesis of Cyclopentanoids via Enyne Cycloaddition
Reaction Using Methylmanganese Carbonyl Complex.**

Joo Eun Lee, Soon Hyeok Hong, and Young Keun Chung*, Department of Chemistry and Center for Molecular Catalysis, College of Natural Sciences, Seoul National University, Seoul 151-742, Korea

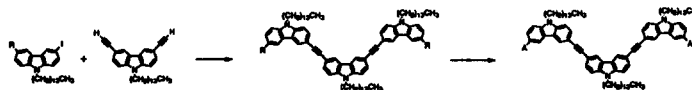


Tetrahedron Letters, 1997, 38, 1785

**A NOVEL APPROACH TO THE SYNTHESIS OF CONJUGATED
CARBAZOLE TRIMERS AS MULTIFUNCTIONAL CHROMOPHORES**

FOR PHOTOREFRACTIVE MATERIALS. Yadong Zhang,[†] Tatsuo Wada,^{†,‡} Liming Wang[‡] and Hiroyuki Sasabe,^{†,‡} [†]Core Research for Evolution Science and Technology (CREST), JST, [‡]Frontier Research Program, The Institute of Physical and Chemical Research (RIKEN), Hirosawa 2-1, Wako, Saitama, 351-01 Japan

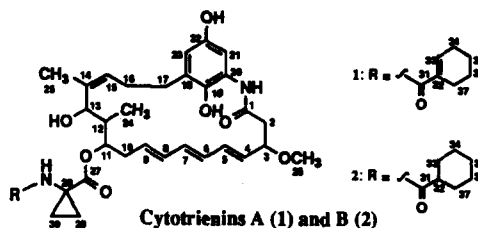
Conjugated carbazole trimers for photorefractive Chromophores were synthesized by Pd-catalytic coupling reaction.



NOVEL TRIENE-ANSAMYCINS, CYTOTRIENINS A AND B, INDUCING APOPTOSIS ON HUMAN LEUKEMIA HL-60 CELLS

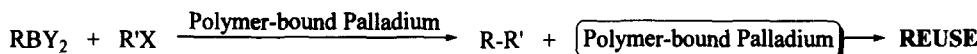
H.-p. Zhang, H. Kakeya, H. Osada,*
Antibiotics Laboratory, Institute of Physical and Chemical Research (RIKEN),
Hirosawa 2-1, Wako-shi 351-01, Japan

Cytotrienins A (1) and B (2) were two novel triene-ansamycins containing a unique 1-aminocyclopropane carboxylic acid moiety, and exhibited a potent apoptosis-inducing activity on HL-60 cells.


Polymer-bound Palladium-catalyzed Cross-coupling of Organoboron Compounds with Organic Halides and Organic Triflates Su-Bum Jang*

*Research & Development Center, DaeWoong Pharmaceutical Co. Ltd., 223-23Sangdaewon-Dong, Joongwon-Gu, Sungnam 462-120, Kyunggi-Do, Korea

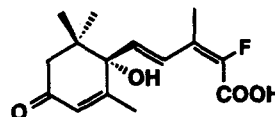
The Polymer-bound palladium-catalyzed cross-coupling reaction of electrophiles (i.e., halides and triflates) with organoboron compounds to form carbon-carbon bonds was achieved at mild conditions with very high activity in the Suzuki coupling reaction. The polymeric catalyst can be easily separated from a reaction mixture and reused more than 10 times with no decrease in activity.


SYNTHESIS OF 2-FLUOROABSCISIC ACID: A POTENTIAL PHOTO-STABLE ANALOG OF ABSCISIC ACID

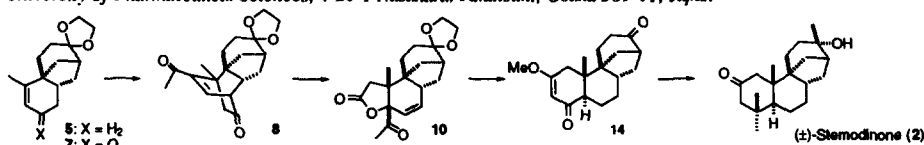
Bum Tae Kim^{a†} Yong Ki Min^a, Tadao Asami^b, No Kyun Park^a,
Oh Young Kwon^c, Kwang Yun Cho^c and Shigeo Yoshida^b

^aKorea Research Institute of Chemical Technology, P.O.Box 107, Yusong, Taejeon 305-606, Korea; ^bThe Institute of Physical and Chemical Research (RIKEN), 2-1 Hirosawa, Wako 351-01, Japan; ^cDepartment of Chemistry, Soong-Sil University, 1-1, Sang Do 5 Dong, Dong Jak Gu, Seoul 156-743, Korea

Abstract: 2-Fluoroabscisic acid (ABA) was synthesized by introducing fluorine through the Wittig reaction of α -ionone derivative with triethyl phosphono-2-fluoroacetate. Molecular orbital calculations showed that the introduction of fluorine at the 2 position stabilized the configuration of the side chain.


TOTAL SYNTHESIS OF (±)-STEMODINONE

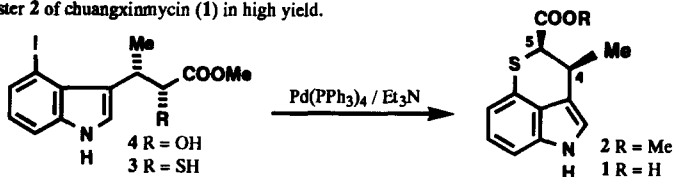
Tetsuaki Tanaka,^a Kazuo Murakami,^a Atsushi Kanda,^a Debasis Patra,^a Sachiko Yamamoto,^a Norifumi Satoh,^a Sang-Won Kim,^b Toshimasa Ishida,^c Yasuko In^c and Chuzo Iwata^{a,*} ^aFaculty of Pharmaceutical Sciences, Osaka University, 1-6 Yamadaoka, Suita, Osaka 565, Japan; ^bFaculty of Pharmaceutical Sciences, Josai University, 1-1 Keyakidai, Sakado, Saitama 350-02, Japan; ^cOsaka University of Pharmaceutical Sciences, 4-20-1 Nasahara, Takatsuki, Osaka 569-11, Japan



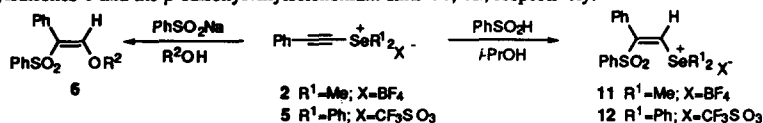
NEW TOTAL SYNTHESIS OF (±)-CHUANGXINMYCIN

Keisuke Kato, Machiko Ono and Hiroyuki Akita*

School of Pharmaceutical Science, Toho University, 2-2-1 Miyama, Funabashi, Chiba, 274, Japan

Palladium-catalysed cyclisation of 4'-iodo-thioindolmycenate 3 derived from 4'-iodoindolmycenate 4 with retention of C₂-stereochemistry gave the (±)-*cis* methyl ester 2 of chuangxinmycin (1) in high yield.*Tetrahedron Letters*, 1997, 38, 1805**REACTIONS OF ALKYNYLSELENONIUM SALTS WITH SODIUM BENZENESULFINATE.** Tadashi Kataoka,* Yoshihiro Banno, Shin-ichi Watanabe, Tatunori Iwamura and Hiroshi Shimizu
Gifu Pharmaceutical University, 6-1, Mitahora-higashi 5-chome, Gifu 502 Japan*Tetrahedron Letters*, 1997, 38, 1809

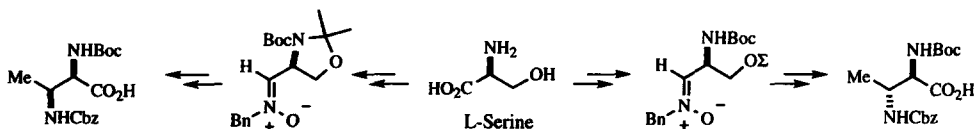
The reactions of alkynylselenonium salts 2, 5 with sodium benzenesulfinate and with benzenesulfinic acid afforded the (Z)-β-alkoxyvinylsulfones 6 and the β-sulfonylvinylselenonium salts 11, 12, respectively.

**STEREOCONTROLLED SYNTHESIS OF 2,3-DIAMINO BUTANOIC ACIDS**

P. Merino,* A. Lanaspá, F.L. Merchán and T. Tejero.

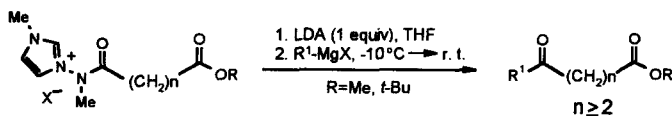
Departamento de Química Orgánica, Facultad de Ciencias, ICMA. Universidad de Zaragoza. E-50009 Zaragoza. Aragon. Spain.

A stereodivergent synthesis of the title compounds is achieved using L-serine as the only starting compound.

Tetrahedron Letters, 1997, 38, 1813**CHEMOSELECTIVE ADDITION OF GRIGNARD REAGENTS TO ALKOXY-CARBONYLALKYL-N-IMIDAZOLIUM-N-METHYL AMIDES: SYNTHESIS OF 4-OXO AND HOMOLOGOUS ESTERS.** María A. de las Heras, Juan J. Vaquero, José L. García Navio, Julio Alvarez-Builla.*

Departamento de Química Orgánica, Universidad de Alcalá, 28871-Alcalá de Henares, Madrid. Spain.

The reaction of alkoxy carbonylalkyl-N-imidazolium-N-methyl amides with 1 equiv of LDA followed by addition of 1 equiv of a Grignard reagent provides a general method for the synthesis of 4-oxo and homologous esters.

Tetrahedron Letters, 1997, 38, 1817

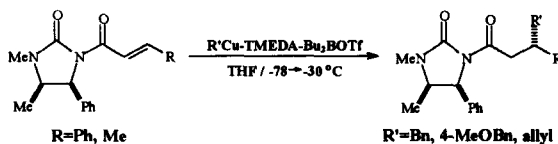
Tetrahedron Letters, 1997, 38, 1821

Dibutylboron Triflate Promoted Conjugate Addition of Benzylic and Allylic Organocopper Reagents to Chiral α,β -Unsaturated N-Acyl Imidazolidinones

Pieter S. van Heerden, Barend C.E. Bezuidenhout and Daneel Ferreira

Department of Chemistry, University of the Orange Free State, P.O. Box 339, Bloemfontein, 9300 South Africa

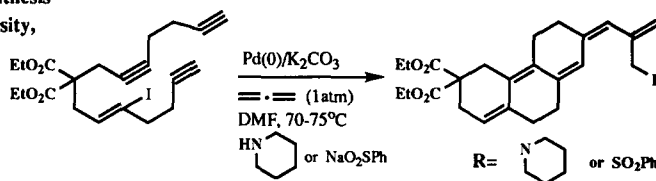
The organocopper-Lewis acid system, $\text{RCu-TMEDA-Bu}_2\text{BOTf}$, is useful for conjugate addition to highly constrained chiral α,β -unsaturated N-acyl imidazolidinones. Bu_2BOTf exhibits a dramatic increase in reactivity during 1,4-addition of benzylic and allylic organocopper reagents.



PALLADIUM CATALYSED TRISCYCLISATION-ANION CAPTURE QUEUING CASCADES

Ronald Grigg,^{*} Rukhsana Rasul and Vladimir Savic
Molecular Innovation, Diversity and Automated Synthesis (MIDAS) Centre, School of Chemistry, Leeds University, Leeds LS2 9JT

Two- and three-component polycyclisation-anion capture processes employing vinyl and allenyl starter species occur regio- and stereo-specifically and in good yield.



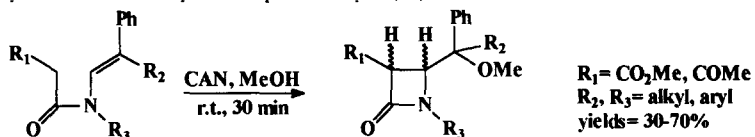
Tetrahedron Letters, 1997, 38, 1829

CERIC AMMONIUM NITRATE PROMOTED FREE RADICAL CYCLIZATION REACTIONS LEADING TO β -LACTAMS

Andrea D'Annibale, Antonella Pesce, Stefano Resta and Corrado Trogolo

Centro C.N.R. di Studio per la Chimica delle Sostanze Organiche Naturali, Dipartimento di Chimica, Università "La Sapienza", P.le Aldo Moro 5, 00185 Roma, ITALIA.

A 4-*exo-trig* radical cyclization of N-alkenyl amides promoted by Ce(IV) affords azetidion-2-ones.



Tetrahedron Letters, 1997, 38, 1833

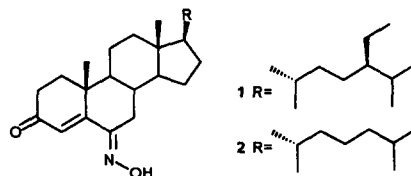
Isolation and Synthesis of the First Natural 6-Hydroximino 4-en-3-one Steroids from the Sponges *Cinachyrella* spp.

Jaime Rodríguez¹, Lucia Nuñez¹, Solange Peixinho² and Carlos Jiménez^{2*}

¹Departamento de Química Fundamental e Industrial, Universidade da Coruña, Campus A Zapateira s/n, A Coruña 15071, Spain.

²Departamento de Zoologia, Instituto de Biologia, Universidade Federal de Bahia, Campus Universitário de Ondina, 40170-290, Salvador, Bahia, Brazil.

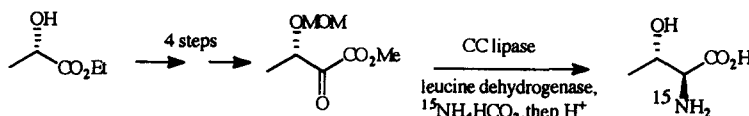
The first natural 6-hydroximino 4-en-3-one steroids (1-2) are described from a mixture of two morphospecies of the Brazilian sponges *Cinachyrella alloclada* and *C. apion*. Synthetic and spectroscopic methods corroborated the proposed structures.



ENANTIOSELECTIVE SYNTHESIS OF α -AMINO- β -HYDROXY ACIDS, [^{15}N]-L-ALLOTHREONINE AND [^{15}N]-L-THREONINE

Andrew Sutherland and Christine L. Willis*

School of Chemistry, University of Bristol, Cantock's Close, Bristol BS8 1TS, UK.



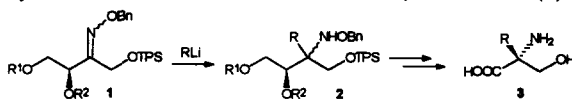
A chemo-enzymatic approach to the syntheses of [^{15}N]-L-threonine from methyl (*R*)-lactate and [^{15}N]-L-allothreonine from ethyl (*S*)-lactate is described.

Diastereoselective Additions Of Organolithium Reagents to the C=N Bond of Protected Erythrulose Oxime Ethers. Synthesis of Enantiopure α,α -Disubstituted α -Aminoacids.

J.A. Marco,*^a M. Carda,*^b J. Murga,^b F. González,^b and E. Falomir.^b

^aDpt. Q. Orgánica, Univ. Valencia, E-46100 Burjassot. ^bDpt. Q. Inorg. y Orgánica, Univ. Jaume I, E-12080 Castellón, Spain.

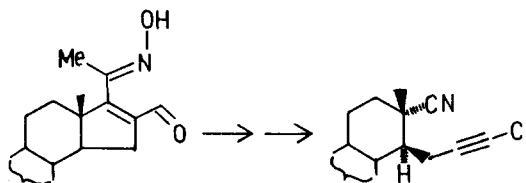
The stereoselective addition of organolithium reagents to erythrulose oxime ethers of general formula 1 (either configuration of the C=N bond) has yielded aminopolyols 2. Two of them have been converted into α,α -disubstituted (*R*)- α -aminoacids 3 (*R*=Me, Ph).



A NOVEL SYNTHESIS OF STEROIDAL HALOMETHYLENES AND THEIR RING OPENING REACTION TO ALKYNES

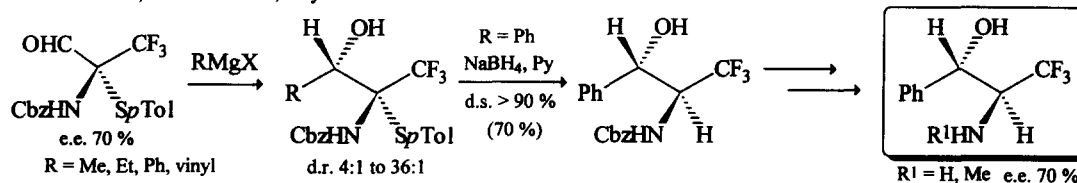
Shahadat Ahmed and Romesh Chandra Boruah*
Organic Chemistry Division, Regional Research Laboratory,
Jorhar-785006, India

Abstract: γ -Formyl conjugated steroidal oximes under Vilsmeier condition afforded (*E*)-chloromethylene as potential precursor of steroidal alkynes.



***N*-Cbz-TRIFLUOROPYRVALDEHYDE *N,S*-KETAL: ABSOLUTE STEREOCHEMISTRY AND ADDITION OF GRIGNARD REAGENTS.**

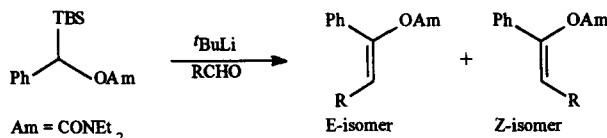
HIGHLY STEREOSELECTIVE ENTRY TO TRIFLUORO ANALOGUES OF *EPHEDRA* ALKALOIDS. Alessandro Volonterio, Pierfrancesco Bravo*, Silvia Capelli, Stefano V. Meille, and Matteo Zanda*, Dipartimento di Chimica del Politecnico, C.N.R. - C.S.S.O.N., Via Mancinelli 7, I-20131 Milano, Italy.



THE PETERSON OLEFINATION OF BENZYL CARBAMATES¹

L. Frances van Staden, Birgit Bartels-Rahm and Neville D. Emslie,
Department of Chemistry, University of Natal, Private Bag X01, Scottsville, 3209,
Republic of South Africa.

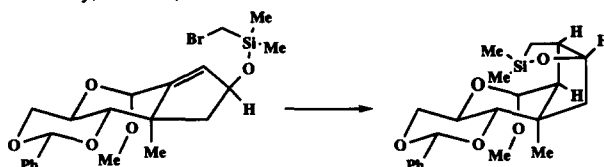
The stereoselective synthesis of substituted vinyl carbamates from α -silylbenzyl carbamates is described.



THE FIRST EXAMPLE OF A HIGHLY STEREOSELECTIVE INTRAMOLECULAR

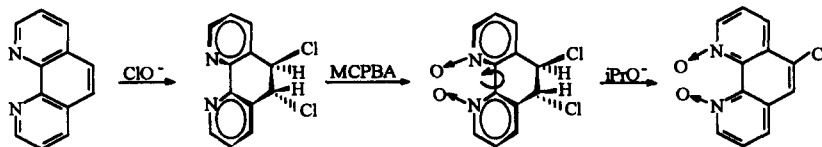
RADICAL CYCLISATION OF A CYCLOPENTENOL DERIVATIVE. Paul R Jenkins* and Andrew J Wood, Department of Chemistry, Leicester University, Leicester, UK LE1 7RH.

Silyl methylene radical cyclisation of a β -allylic cyclopentaannulated derivative of glucose leads to a single *cis* fused tricyclic ring system.



IS THE FORMATION OF 1,10-PHENANTHROLINE DI-N-OXIDE POSSIBLE ?

Róża Antkowiak and Wiesław Z. Antkowiak*
Faculty of Chemistry, Adam Mickiewicz University, Grunwaldzka 6, 60-780 Poznań, Poland



REACTIONS OF DIAZOACETATES WITH PHOSPHORUS TRIESTERS AND THIOPHOSPHATE TRIESTER: >P⁺-O-C< AND >P⁺-S-C<

INTERMEDIACY FORMATION Konstantin A. Popov, Alexander M. Polozov* and Sergei V. Tcherezov
A.M. Butlerov Research Chemical Institute, Kazan State University, Lenin Str. 18, Kazan 420008, Russian Federation.

