

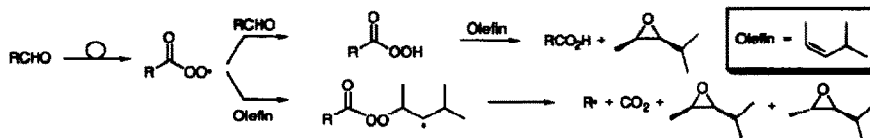
GRAPHICAL ABSTRACTS

Tetrahedron Letters, 1994, 35, 8077

Aldehyde/Olefin Cooxidations: Parallel Epoxidation Pathways and Concerted Decomposition of the Peroxyacyl-Olefin Adduct

Kevin R. Lassila*, Francis J. Waller, Steven E. Werkheiser, and Amy L. Wressell
Corporate Science and Technology Center, Air Products and Chemicals, Inc., Allentown, PA 18195

Aldehyde-mediated olefin epoxidations appear to proceed by not only a peracid path, but also a parallel radical addition path in which the peroxyacyl-olefin adduct decomposes concertedly to form alkyl radical, CO₂ and epoxide.



Tetrahedron Letters, 1994, 35, 8081

A RAPID DEALKYLATION OF PHOSPHONATE DIESTER FOR THE PREPARATION OF 4-PHOSPHONOMETHYLPHENYLALANINE-CONTAINING PEPTIDES.

O.M. Green

ARIAD Pharmaceuticals Inc., 26 Landsdowne St., Cambridge, MA 02139, U.S.A.

Phosphonomethylphenylalanine-containing peptides are rapidly and conveniently prepared from the corresponding phosphonate diesters using TMSI/MeCN.

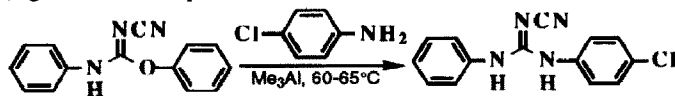


Tetrahedron Letters, 1994, 35, 8085

TRIMETHYLALUMINUM PROMOTED SYNTHESIS OF CYANOGUANIDINES

Karnail S. Atwal*, Francis N. Ferrara and Syed Z. Ahmed
The Bristol-Myers Squibb Pharmaceutical Research Institute,
P. O. Box 4000, Princeton, N.J. 08543-4000

The synthesis of cyanoguanidines from N-cyano-O-phenylisoureas and relatively nonbasic amines (e.g., aniline) is reported.

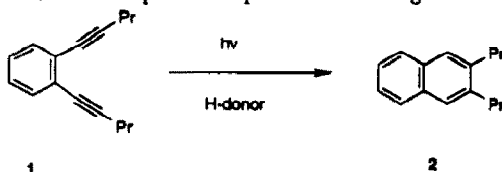


Tetrahedron Letters, 1994, 35, 8089

Photochemical Analogue of the Bergman Cycloaromatization Reaction. Nicholas J. Turro and Ariella Evenzahav,

Dept. of Chemistry, Columbia University, New York, New York 10027;
K. C. Nicolaou, Scripps Research Institute, La Jolla, CA, 92037

When irradiated, 1 forms a product expected from a Bergman rearrangement reaction.

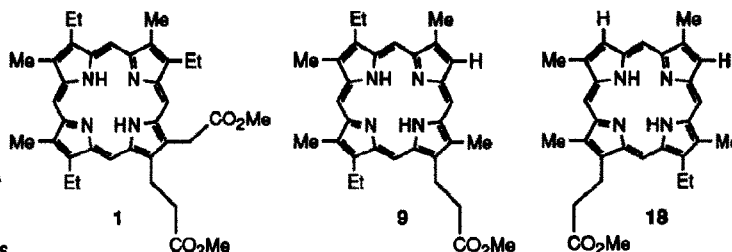


TOTAL SYNTHESSES OF NEW PORPHYRINS

ISOLATED FROM THE CORAL SEA
DEMOSPONGE *CORALLISTES* SP.

Ravindra K. Pandey, Sam H. Leung
and Kevin M. Smith, Department
of Chemistry, University of California,
Davis, CA 95616, USA and Department
of Radiation Biology, Roswell Park
Cancer Institute, Buffalo, NY 14263, USA

Syntheses, from pyrroles, of **1**, **9**, and **18**,
using MacDonald and a,c-biladiene routes.

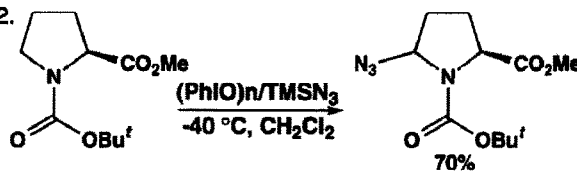


Tetrahedron Letters, 1994, 35, 8093

**Oxidation of L-Proline Methyl Ester Derivatives with the
Iodosylbenzene/Trimethylsilylazide Reagent Combination**

Philip Magnus* and Christopher Hulme
Department of Chemistry and Biochemistry,
University of Texas at Austin, Austin, Texas 78712.

Treatment of a variety of L-proline derivatives
with $(\text{PhIO})_n/\text{TMSN}_3/\text{CH}_2\text{Cl}_2$ at -40°C gave the
 α -azidoamides as the major product.



Tetrahedron Letters, 1994, 35, 8097

**SYNTHESIS OF DEPHOSTATIN, A NOVEL PROTEIN TYROSINE
PHOSPHATASE INHIBITOR.**

Pierre Hamel* and Yves Girard, Merck
Frost Centre for Therapeutic Research, P.O. Box 1005 Pointe-Claire - Dorval, Québec H9R 4P8 Canada

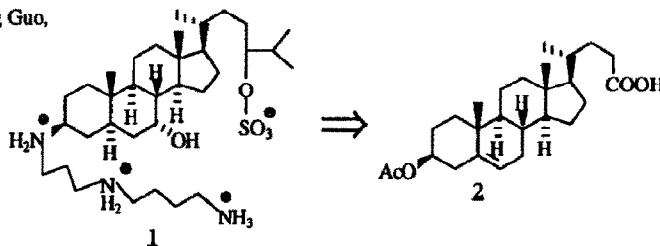
The novel protein tyrosine phosphatase inhibitor dephostatin was synthesized in 6 steps with an overall yield of 35% from
2-nitrohydroquinone



Tetrahedron Letters, 1994, 35, 8101

**SYNTHESIS OF SQUALAMINE. A STEROIDAL ANTIBIOTIC FROM
THE SHARK**

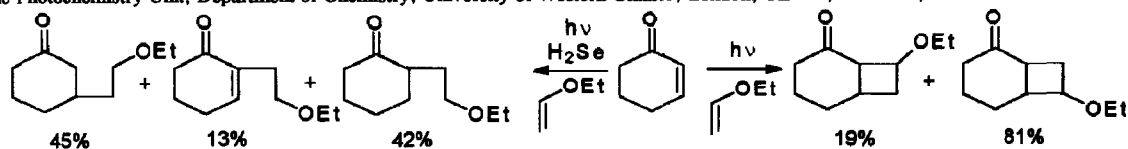
Robert M. Moriarty, Sudersan M. Tuladhar, Liang Guo,
and Suzanne Wehrli
Steroids, Ltd
2201 W. Campbell Park Drive,
Chicago, IL 60612
Children's Hospital of Philadelphia,
University of Pennsylvania School of Medicine,
34th & Civic Center Blvd.,
Philadelphia, PA 19104



Tetrahedron Letters, 1994, 35, 8103

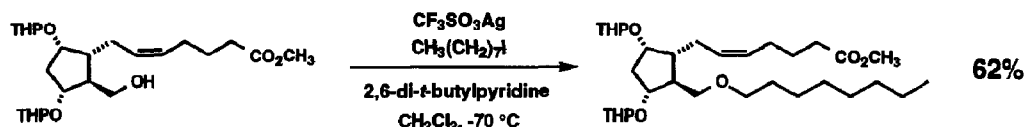
THE PHOTOCHEMICAL CYCLOADDITION REACTION OF 2-CYCLO-HEXENONE WITH ALKENES: TRAPPING OF TRIPLET 1,4-BIRADICAL INTERMEDIATES WITH HYDROGEN SELENIDE. David J. Maradyn and Alan C. Weedon.*

The Photochemistry Unit, Department of Chemistry, University of Western Ontario, London, Ontario, N6A 5B7, Canada.



A MILD PROCEDURE FOR ETHERIFICATION OF ALCOHOLS WITH PRIMARY ALKYL HALIDES IN THE PRESENCE OF SILVER TRIFLATE

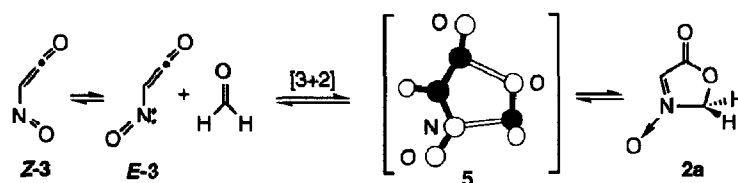
Robert M. Burk,* Todd S. Gac and Michael B. Roof
Department of Chemical Sciences, Allergan Inc., 2525 Dupont Drive, Irvine, California 92713



1°, 2° and even 3° alcohols were etherified with primary alkyl halides in the presence of $\text{CF}_3\text{SO}_3\text{Ag}$ and 2,6-di-*tert*-butylpyridine in CH_2Cl_2 .

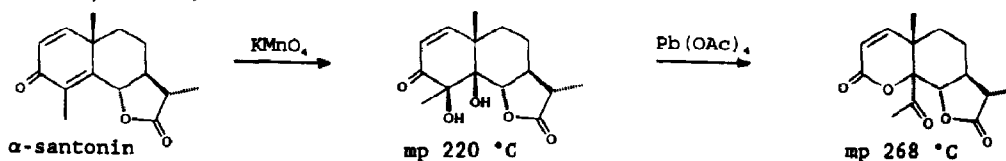
AN AB INITIO STUDY OF THE REACTIVITY OF NITROSOKETENE WITH FORMALDEHYDE. Sihyun Ham and David M. Birney,* Department of Chemistry and Biochemistry, Texas Tech University, Lubbock, TX 79409 USA

Ab initio calculations predict that *E*-nitrosoketene (*E*-3) adds to formaldehyde in a concerted, pseudopericyclic [3+2] reaction (5) with a barrier of only 3.5 kcal/mol.



STEREOCHEMISTRY OF 4,5-DIHYDROXY- α -SANTONIN AND STRUCTURE OF A NEW SANTONIN OXIDATION PRODUCT.

S. K. Paknikar,* B. L. Malik, Department of Chemistry, Goa University, P.O. 403202, Goa, India. R. B. Bates,† S. Caldera and T. V. Wijayaratne, Department of Chemistry, University of Arizona, Tucson, Arizona 85721, USA



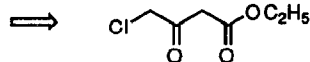
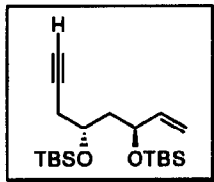
Tetrahedron Letters, 1994, 35, 8119

A Practical Asymmetric Synthesis of a 1,7-Enyne A-Ring Synthon En Route Toward the Total Synthesis of Vitamin D₃ Analogues

Barry M. Trost* and Paul R. Hanson

Department of Chemistry, Stanford University, Stanford, CA 94305

A new synthesis of a key A-ring synthon en route to vitamin D₃ analogues has been achieved in 8 steps (overall yield of 20.4%) starting from commercially available 4-chloroacetoacetate

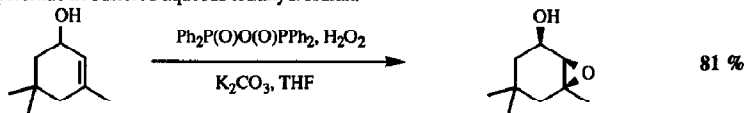


**A NEW PARADIGM FOR ALKENE EPOXIDATION
ACTIVATION OF HYDROGEN PEROXIDE BY
ORGANOPHOSPHORUS ELECTROPHILES**

Andrew S. Kende,* Philippe Delair, Benjamin E. Blass

Department of Chemistry, University of Rochester, Rochester, New York, 14627, USA.

Diphenylphosphinic anhydride and certain other organophosphorus electrophiles mediate the high-yield conversion of alkenes to epoxide by hydrogen peroxide in buffered aqueous tetrahydrofuran.



Tetrahedron Letters, 1994, 35, 8123

A Short Synthesis of the Tricyclo[3.3.2^{1,4}.0]decane Ring System

Scott McN. Sieburth* and Elena Doval Santos, Department of Chemistry, State University of New York at Stony Brook, Stony Brook, New York 11794-3400 USA

Acyloin condensation of endo, endo diester I yields the tricyclo[3.3.2^{1,4}.0]decane ring system II containing a 1,2-disilyloxy alkene of surprising stability.

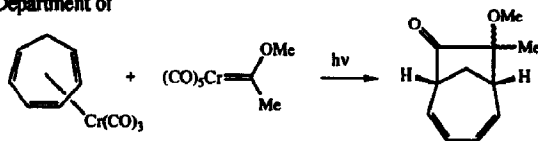


Tetrahedron Letters, 1994, 35, 8127

**TRANSITION METAL PROMOTED HIGHER-ORDER
CYCLOADDITION REACTIONS. [6+2] CYCLOADDITION OF A
FISCHER CARBENE COMPLEX WITH
TRICARBONYL(CYCLOHEPTATRIENE) CHROMIUM(0)**

James H. Rigby*, F. Christopher Pigge, and Mark D. Ferguson, Department of Chemistry, Wayne State University, Detroit, Michigan 48202

Photolytic reaction of pentacarbonyl [(methoxy)(methyl)carbene] chromium(0) with tricarbonyl(cycloheptatriene) chromium(0) afforded an isomeric mixture of bicyclo[4.2.1]nonanone products.



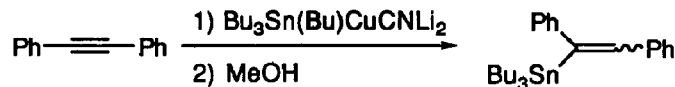
Tetrahedron Letters, 1994, 35, 8131

Tetrahedron Letters, 1994, 35, 8133

Cis-Trans Stereoselectivity in the Stannylation of Diphenylacetylene

Clark H. Cummins* and Eva J. Gordon, Dow Chemical Company, Bldg. 1707, Midland, MI 48674, USA

Stereospecific *cis*- or *trans*- stannylation of diphenylacetylene can be obtained by temperature control during vinylcuprate hydrolysis.



cis-addition: Temp = -78
39%

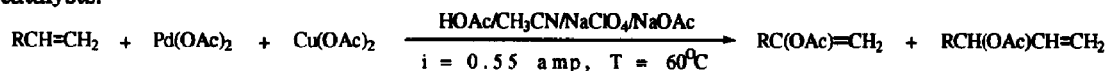
trans-addition: Temp = 25
61%

PALLADIUM CATALYZED INDIRECT ELECTROCHEMICAL ACETOXYLATION OF OLEFINS.

Tetrahedron Letters, 1994, 35, 8137

Frederick W. Hartstock*, Department of Chemistry, Wilfrid Laurier University, Waterloo, Ontario, Canada, N2L 3C5; and Danial D. M. Wayner, Steacie Institute of Molecular Sciences, National Research Council of Canada, Ottawa, Ontario, Canada, K1A 0R6.

Olefins are catalytically acetoxylation by Pd(OAc)₂/Cu(OAc)₂ in combination with anodic re-oxidation of the catalysts.



The Stereochemistry of Carbonyl Oxides From Ozonolysis of

Vinyl Ethers. William H. Bunelle* and Sang-gi Lee,
Department of Chemistry, University of Missouri, Columbia, Missouri 65211

Tetrahedron Letters, 1994, 35, 8141

E-Vinyl ethers produce mainly *syn*, and Z-vinyl ethers mainly *anti*-carbonyl oxides, as determined by intramolecular trapping.

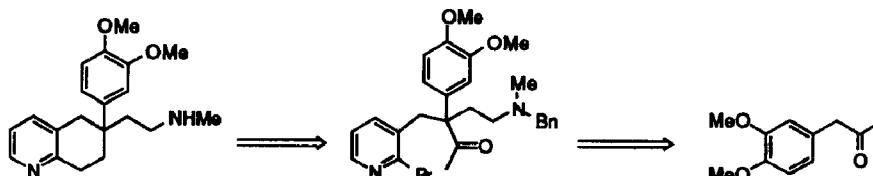


A SHORT SYNTHESIS OF (±)-TORTUOSAMINE

R. Richard Goehring

Department of Chemistry, North Carolina State University, Raleigh, NC 27695-8204

Tetrahedron Letters, 1994, 35, 8145

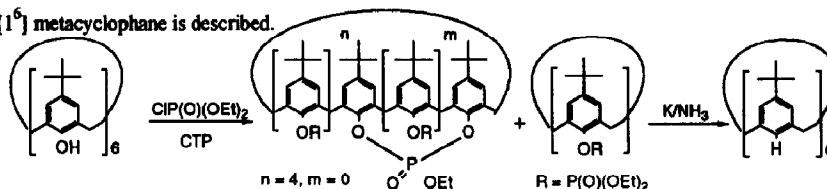


SYNTHESIS OF A FULLY OH-DEPLETED *p*-tert-BUTYL-CALIX[6]ARENE

Tetrahedron Letters, 1994, 35, 8147

J.-B. Regnouf de Vains, S. Pellet-Rostaing, R. Lamartine; Laboratoire de Chimie Industrielle, Université Claude Bernard, 69622 Villeurbanne cedex.

The synthesis of the *p*-tert-butyl-[1⁶] metacyclophane is described.

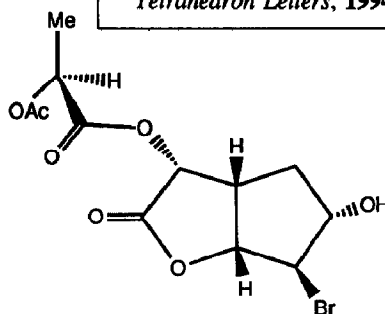


Convenient Chemical Resolution of a Bicyclic Hydroxylactone of Synthetic Interest

Tetrahedron Letters, 1994, 35, 8151

Fabienne Burlina, Pascale Clivio, Jean-Louis Fourrey*, Claude Riche and Martial Thomas
Institut de Chimie des Substances Naturelles, C.N.R.S., 91198 Gif-sur-Yvette, France.

The bicyclic lactone 1 was resolved by treatment with (S)-O-acetylactyl chloride to give a separable pair of diastereomers (-)-3 and (-)-4 whose configuration was determined following X-ray crystallography performed on the bromo derivative 5.

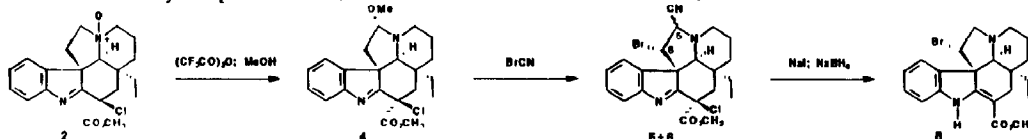


NOUVELLE FONCTIONNALISATION DE LA CHAÎNE TRYPTAMINE DE LA (-) VINCADIFFORMINE ALCALOÏDE À SQUELETTE ASPIDOSPERMANE

Tetrahedron Letters, 1994, 35, 8153

Guy Lewin*, Laboratoire de Pharmacognosie, Faculté de Pharmacie, Bld Bequereel, 14032 Caen France et Jacques Poisson, Laboratoire de Chimie des Substances Thérapeutiques Naturelles, Faculté de Pharmacie, av. J.B. Clément, 92296 Châtenay-Malabry France.

Starting from 16-chloro-1-dehydrovincadifformine-N4-oxyle 2, a two-steps sequence afforded through stereospecific bromination on C-5, 5 and 6. Further reduction of this mixture provided (-) 6S-bromovincadifformine 8.

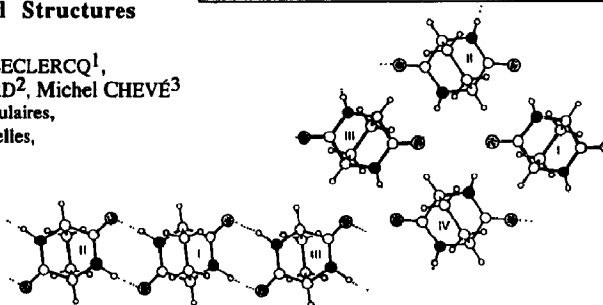


Chirality Directed Self-Assembly.— Resolution of 2,5-Diazabicyclo[2.2.2]octane-3,6-dione and Crystal Structures of its (±) Racemic and (-) Enantiomeric Forms

Tetrahedron Letters, 1994, 35, 8157

Marie-Josèphe BRIENNE¹, Jacqueline GABARD¹, Martine LECLERCQ¹, Jean-Marie LEHN^{1*}, Michèle CESARIO², Claudine PASCARD², Michel CHEVÉ³ and Gilles DUTRUC-ROSSET³. ¹Chimie des Interactions Moléculaires, Collège de France, Paris, ²Institut de Chimie des Substances Naturelles, Gif-sur-Yvette, ³Département de Chimie Pharmaceutique, Rhône-Poulenc Rorer, Vitry-sur-Seine, France

The self-assembly of the racemic mixture (±)1 and of the enantiomer (-)1 of the bicyclic bis-lactam 1 generates different supramolecular architectures.



FORMATION OF RADICALS BY IRRADIATION OF ALKYL HALIDES IN THE PRESENCE OF TRIETHYLAMINE

Janine Cossy*, Jean-Luc Ranaivosata, Véronique Bellost

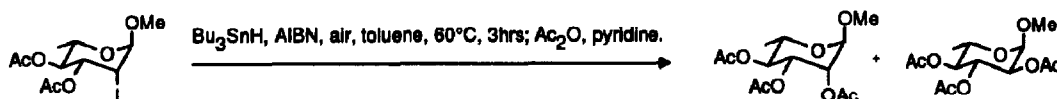
Laboratoire de Chimie Organique, Associé au CNRS., ESPCI, 10 rue Vauquelin, 75231 Paris Cedex 05 - France

The irradiation at 254 nm of alkyl halides in the presence of triethylamine leads, with high yields, to the corresponding reduction products or to the cyclized products when an unsaturation is present.

RADICAL OXYGENATION OF 2-DEOXY-2-iodo-HEXOPYRANOSIDES WITH MOLECULAR OXYGEN.

Stéphane Moutel and Jacques Prandi*, Université d'Orléans, Laboratoire de Biochimie Structurale, associé au CNRS, BP 6759, 45067 Orléans Cedex 2, France.

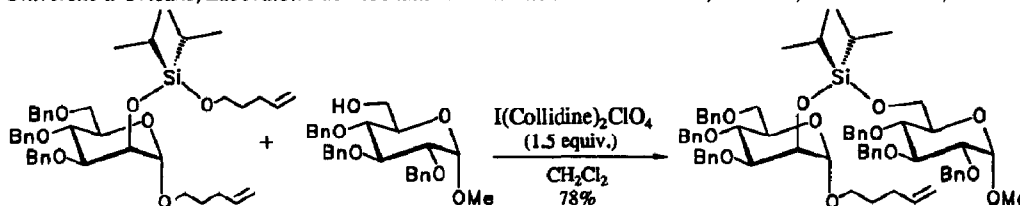
2-Deoxy-2-iodo hexopyranosides react with molecular oxygen and tributylstannane to give the epimeric C-2 alcohols in high yield and moderate selectivity.



NOVEL SILYLATING AGENTS EMPLOYING 4-PENTENYL SILYL ETHERS.

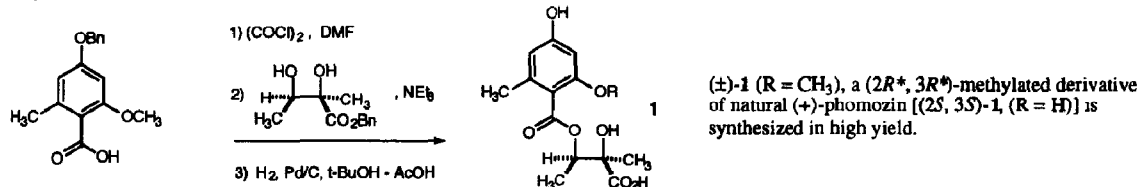
Caroline Colombier, Troels Skrydstrup* and Jean-Marie Beau,

Université d'Orléans, Laboratoire de Biochimie Structurale associé au CNRS, BP 6759, 45067 Orléans, France



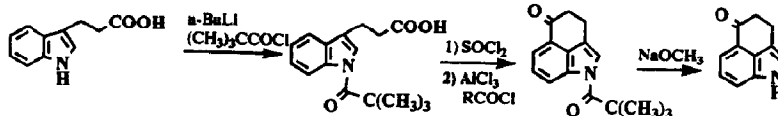
Diastereoselective Synthesis of a Methylated Derivative of Phomozin, a Phytotoxin Isolated from *Phomopsis helianthi*, a Phytopathogenic Fungus of Sunflowers

Robert Nougier, Michèle P. Bertrand, Philippe Picon and Patricia Perfetti.- Laboratoire de Chimie Moléculaire Organique, Faculté Saint Jérôme, Av. Normandie-Niemen, 13397 Marseille cedex 20, France.



**A FACILE BIOMIMETIC SYNTHESIS OF UHLE'S
KETONE BY THE REGIOSELECTIVE FRIEDEL-CRAFTS
CYCLIZATION OF INDOLE-3-YLPROPIONYL CHLORIDE**

Katsunori Teranishi[†], Syunzi Hayashi, Shin-ichi Nakatsuka^{††*} and Toshio Goto
School of Agriculture, Nagoya University, Chikusa, Nagoya 464-01, Japan

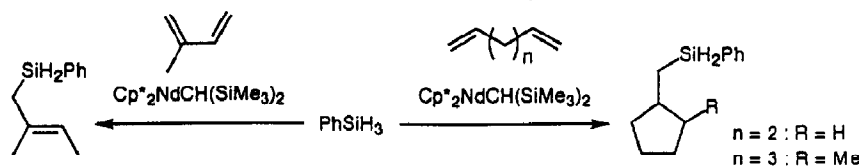


Hydrosilation of Dienes Catalyzed by $Cp^*_2NdCH(SiMe_3)_2$

Shun-ya Onozawa,^{a)} Toshlyasu Sakakura^{*a)} and Masato Tanaka^{*a,b)}

a) National Institute of Materials and Chemical Research, Higashi, Tsukuba, Ibaraki 305, Japan

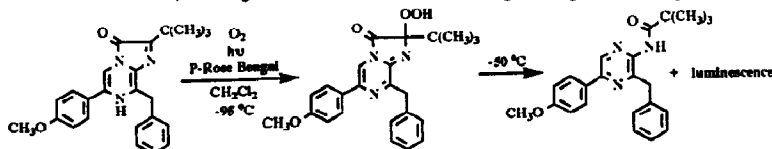
b) Department of Chemistry, University of Tsukuba, Tsukuba, Ibaraki 305, Japan



**SYNTHESIS OF HYDROPEROXIDE VIA
PHOTOXYGENATION FOR A MODEL
AEQUORIN BIOLUMINESCENCE**

Katsunori Teranishi,^{*} Kazuo Ueda,[†] Hidekazu Nakao, Makoto Hisamatsu and Tetsuya Yamada
School of Bioresources, Mie University, Kamihama-chou, Tsu 514, Japan

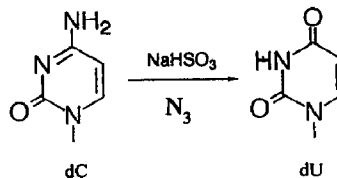
[†]Aburahi Laboratories, Shionogi & Co., Ltd., Koka-cho, Koka-gun, Shiga 520-34, Japan



**Catalysis of Diethylenetriamine for Bisulfite-Induced
Deamination of Cytosine in Oligodeoxyribonucleotides**

Makoto Komiyama^{*} and Shinji Oshima
Department of Chemistry and Biotechnology, Faculty of Engineering,
University of Tokyo, Hongo, Tokyo 113 Japan

Diethylenetriamine (N_3) catalyzes bisulfite ion-induced deamination of 2'-deoxycytidine in DNA to 2'-deoxyuridine at pH 5, much more efficiently than ammonia, ethylenediamine, 3,3'-diaminodipropylamine, and spermine.



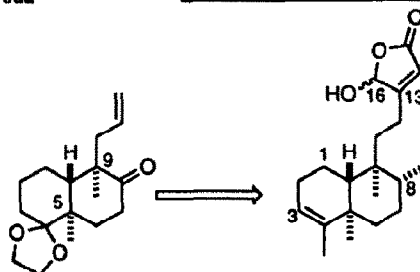
Tetrahedron Letters, 1994, 35, 8189

A Total Synthesis of Antibacterial Clerodane, 16-Hydroxycleroda-3,13(14)Z-dien-15,16-olide

Hisahiro Hagiwara,* Kazuhiro Inome and Hisashi Uda

*Institute for Chemical Reaction Science, Tohoku University
Katahira, Aoba-ku, Sendai 980, Japan*

Abstract: The total synthesis of an antibacterial clerodane, 16-hydroxycleroda-3,13(14)Z-dien-15,16-olide, is described and its absolute stereochemistry has been determined.



Tetrahedron Letters, 1994, 35, 8193

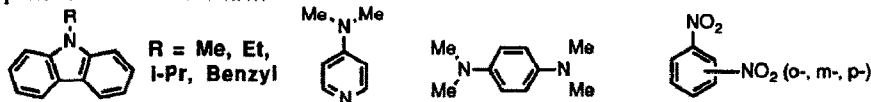
Bimolecular Solid-State Photoreactions in the Adduct Crystals of an Aromatic Nitro Compound with an Aromatic Amine

Yoshikatsu Ito,* Sadayuki Asaoka, and Isao Saito

*Department of Synthetic Chemistry, Faculty of Engineering, Kyoto University, Kyoto 606, Japan
Shigeru Ohba*

Department of Chemistry, Faculty of Science and Technology, Keio University, Yokohama 223, Japan

Of the twelve charge-transfer-type adduct crystals studied, only three which involve m-dinitrobenzene as the acceptor are photoreactive in the solid state.



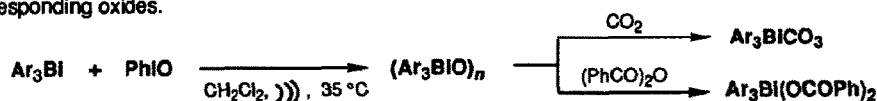
Tetrahedron Letters, 1994, 35, 8197

Ultrasonic Reaction of Triarylbiomuthines and Triarylstibines with Iodosylbenzene. Mild Oxidizing Ability of the Organo-bismuth Oxide Function for Organic Substrates.

Hitomi Suzuki,* Tohru Ikegami and Yoshihiro Matano

Department of Chemistry, Faculty of Science, Kyoto University, Sakyo-ku, Kyoto 606-01, Japan

Sonochemical reaction of triarylbiomuthines and triarylstibines with iodosylbenzene produces the corresponding oxides.



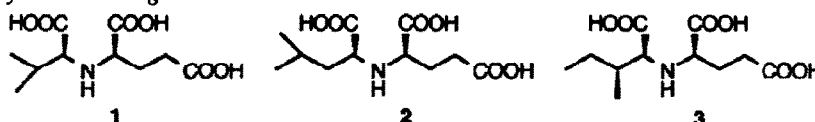
Tetrahedron Letters, 1994, 35, 8201

THREE NEW AMINO ACIDS FROM A POISONOUS MUSHROOM,

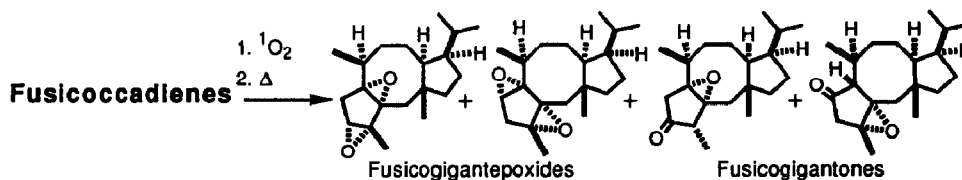
CLITOCYBE ACROMELALGA. Shinji Fushiya, Shohei Yamada,

Mamoru Matsuda and Shigeo Nozoe, Pharmaceutical Institute, Tohoku University, Aobayama, Aoba-ku, Sendai 980-77, Japan

Three new amino acids, valinopine (1), *epi*-leucinopine (2) and isoleucinopine (3), were isolated from a poisonous mushroom, *Clitocybe acromelalga*.



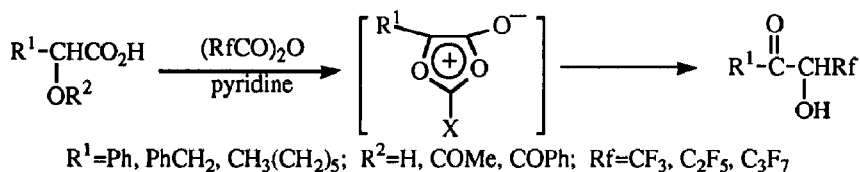
Total Synthesis of Fusicogigantones A and B and Fusicogigantepoxide via the Singlet Oxygen-Oxidation of Fusicoccadienes. "Fusicogigantepoxide B", a Missing Congener Metabolite. Nobuo Kato, Kohji Nakanishi, Xue Wu, Hideyuki Nishikawa, and Hitoshi Takeshita,* Institute of Advanced Material Study, 86, Kyushu University, Kasuga-koen, Kasuga, Fukuoka 816 Japan



A Convenient Synthesis of α -Trifluoromethylated and α -Perfluoroalkylated Acyloins from α -Hydroxy Acids

Masami Kawase ^a and Teruo Kurihara ^b

Faculty of Pharmaceutical Sciences ^a and Faculty of Science, ^b Josai University, Sakado-shi, Saitama 350-02, Japan



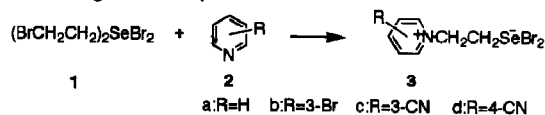
Reactivities of Bis(2-bromoethyl)selenium Dibromide and Its Related Compounds: Formation of Hypervalent T-shaped

Coordinated Selenium Compounds by Reaction with Pyridine and Its Derivatives.

Masatsugu Miura, Yoshinori Takanohashi, Yoichi Habata, and Sadatoshi Akabori*

Department of Chemistry, Faculty of Science, Toho University, Funabashi, Chiba 274, Japan

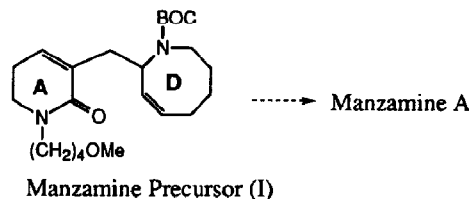
The reaction of 1 with 2 gave T-shaped coordinated compounds 3 in moderate to good yields.



Synthetic Studies on Manzamine A: Construction of a Key Bicyclic AD Ring Subunit I

S. Li, S. Kosemura, and S. Yamamura
Dept. of Chemistry, Faculty of Science and Technology,
Keio University, Hiyoshi, Yokohama, Japan

Synthesis of manzamine A precursor has successfully been accomplished.

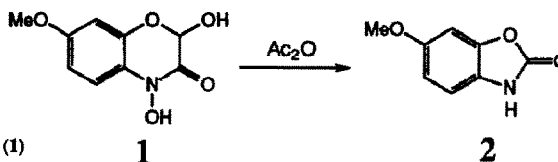


Chemical Studies on 2,4-Dihydroxy-7-methoxy-2*H*-1,4-benzoxazin-3(4*H*)-one
in Connection with 6-Methoxy-2-benzoxazolinone,
an Auxin-inhibiting Substance of *Zea mays* L.

Tetrahedron Letters, 1994, 35, 8221

S. Kosemura, S. Yamamura,* Toyoaki Anai, and K. Hasegawa
Dept of Chem. Faculty of Science and Technology,
Keio University, Hiyoshi, Yokohama 223, Japan
Institute of Applied Biochemistry, University of Tsukuba,
Tennodai 1-1-1, Tsukuba 305, Japan

In the presence of acetic anhydride, the reaction rate of DIMBOA (1)
was remarkably accelerated to afford MBOA (2) in high yield.



Synthetic Study of Marine Macrolide Swinholide A.
Stereocontrolled Synthesis of the C11 - C23 Segment.

Tetrahedron Letters, 1994, 35, 8225

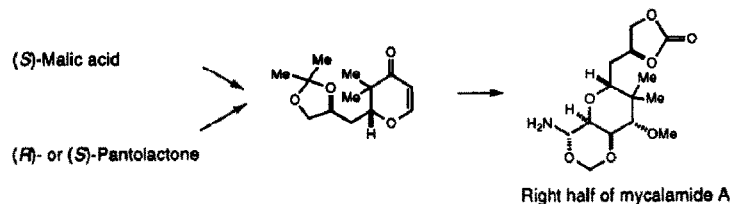
Tadashi Nakata,* Toshiya Komatsu, Kazuo Nagasawa, Haruo Yamada,† and Takashi Takahashi†
The Institute of Physical and Chemical Research (RIKEN), Wako-shi, Saitama 351-01, Japan
† Tokyo Institute of Technology, Meguro, Tokyo 152, Japan



SYNTHESIS OF THE RIGHT HALF OF MYCALAMIDE A.
A FORMAL TOTAL SYNTHESIS

Tetrahedron Letters, 1994, 35, 8229

Tadashi Nakata,* Hiroko Matsukura, Dunlong Jian, and Hiroaki Nagashima
The Institute of Physical and Chemical Research (RIKEN), Wako-shi, Saitama 351-01, Japan

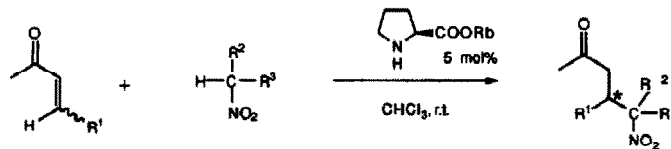


Catalytic Asymmetric Michael Addition of Nitroalkane to Enone and Enal

Tetrahedron Letters, 1994, 35, 8233

Masahiko Yamaguchi,* Tai Shiraiishi, Yoshihiro Igarashi., and Masahiro Hirama
Department of Chemistry, Faculty of Science, Tohoku University, Aoba, Sendai, 980-77 JAPAN

L-Proline rubidium salt catalyzes the asymmetric Michael addition of nitroalkane to enone and enal.

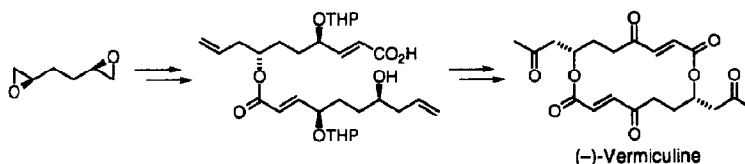


Total Synthesis of (-)-Vermiculine

Tetrahedron Letters, 1994, 35, 8237

Atsushi Noda, Sakae Aoyagi, Nobuo Machinaga, and Chihiro Kibayashi*
Tokyo College of Pharmacy, Horinouchi, Hachioji, Tokyo 192-03, Japan

Macrolide (-)-vermiculine has been synthesized via intramolecular Mitsunobu reaction utilizing a C₂ symmetrical diepoxide chiral synthon.



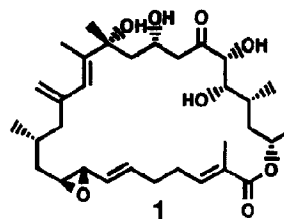
Absolute Stereochemistry of Amphidinolide B

Tetrahedron Letters, 1994, 35, 8241

Masami Ishibashi, Haruaki Ishiyama, and Jun'ichi Kobayashi*

Faculty of Pharmaceutical Sciences,
Hokkaido University, Sapporo 060, Japan

The absolute stereochemistry of amphidinolide B (1), a potent cytotoxic 26-membered macrolide isolated from the cultured marine dinoflagellate *Amphidinium* sp., has been established as 8*S*, 9*S*, 11*R*, 16*R*, 18*S*, 21*R*, 22*S*, 23*R*, and 25*S*, on the basis of enantiospecific synthesis of a degradation product.

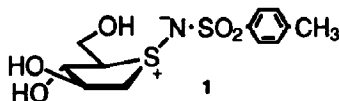


SYNTHESIS OF IMINOTHIASUGAR AS A POTENTIAL TRANSITION-STATE ANALOG INHIBITOR OF GLYCOSYL-TRANSFER REACTIONS

Hideya Yuasa,[§] Tetsuya Kajimoto,^{*§} and Chi-Huey Wong,^{*§,¶} § Frontier Research Program, The Institute of Physical and Chemical Research, 2-1 Hirosawa, Wako, 351-01, JAPAN, and ¶ The Scripps Research Institute, 10666 North Torrey Pines Road, La Jolla, CA 92037, USA

Tetrahedron Letters, 1994, 35, 8243

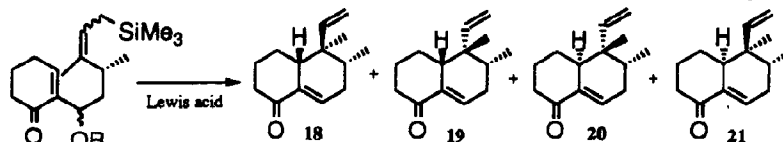
Iminothiasugar 1 was synthesized in 10 steps from D-xylose and was considered to be a potentially useful transition-state analog inhibitor of glycosidases.



ALTERNATION OF STEREOCONTROL MODE IN THE CYCLIZATION OF 2-(3',4'-DIMETHYL-6'-TRIMETHYLSILYL-4'-HEXENYL)-2-CYCLOHEXENONES BY THE AUXILIARY CONTROLLING SUBSTITUENTS AT 1'- AND 2'-POSITIONS OF THE SIDE CHAIN.

Takashi Tokoroyama,* Masashi Kato, Takehiko Aoto, Taeko Hattori, Hideo Iio and Yoshihiko Odagaki. Faculty of Science, Osaka City University, Sumiyoshi-ku, Osaka 558 and Minase Research Institute, Ono Pharmaceutical Co. Ltd., Shimamoto-cho, Osaka 618, Japan

By the choice with respects to kinds and stereochemistry of the oxy-substituents, and Lewis acids, modes of the cyclization can be controlled so as to get one of three isomers 18, 20 and 21 with good selectivities.

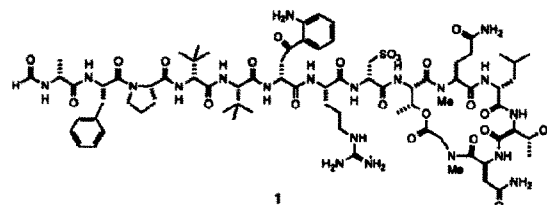


Tetrahedron Letters, 1994, 35, 8247

Discodermin E, a Cytotoxic and Antimicrobial Tetradecapeptide, from the Marine Sponge *Discodermia kilensis*.

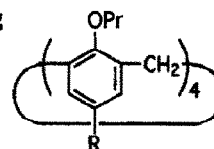
Geonseok Ryu, Shigeki Matsunaga, and Nobuhiro Fusetani,*
 Laboratory of Marine Biochemistry, Faculty of Agriculture,
 The University of Tokyo, Bunkyo-ku, Tokyo 113, Japan

A new cytotoxic and antimicrobial tetradecapeptide, discodermin E (1), has been isolated from the marine sponge *Discodermia kilensis*. The structure of 1 was determined by spectral and chemical methods.

**Self-Assembled Molecular Capsule Based on the Hydrogen-Bonding Interaction between Two Different Calix[4]arenes**

Kwangnak Koh, Koji Araki, and Seiji Shinkai*

Department of Chemical Science & Technology, Faculty of Engineering, Kyushu University, Fukuoka 812, Japan



1 R = COOH

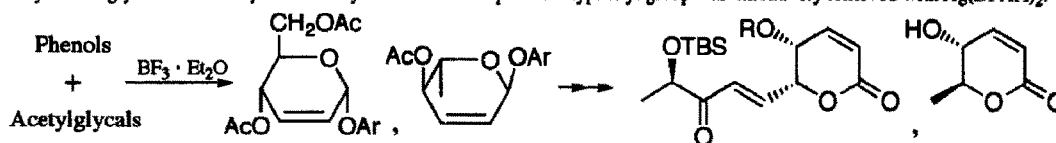
2 R =

Intermolecularly hydrogen-bonded molecular container constructed from two calix[4]arene derivatives having carboxylic acids (1) or pyridyl groups (2) on their upper rim was designed for the first time.

PHENOLIC FERRIER REACTION AND ITS APPLICATION TO THE NATURAL PRODUCT SYNTHESIS.

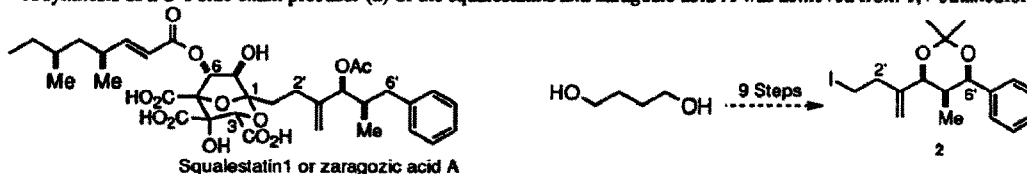
Toshiro Noshita, Takeyoshi Sugiyama*, Yoshiharu Kitazumi, and Takayuki Oritani, Department of Applied Biological Chemistry, Faculty of Agriculture, Tohoku University, Sendai 981, Japan.

Aryl *O*- Δ^2 -glycosides were synthesized by Ferrier reaction. *p*-Methoxyphenyl group was oxidatively removed with Ag(DPAH)₂.

**SYNTHESIS OF THE C-1 SIDECHAIN OF THE SQUALESTATINS AND ZARAGOZIC ACID A.**

Jack G. Parsons and Mark A. Rizzacasa*
 School of Chemistry, The University of Melbourne, Parkville, Victoria 3052, Australia.

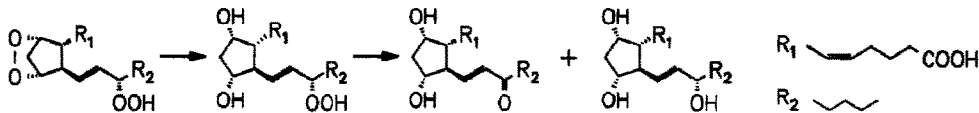
A synthesis of a C-1 side chain precursor (2) of the squalestatin and zaragozic acid A was achieved from 1,4-butanediol.



ENDOPEROXIDE PATHWAY IN PROSTAGLANDIN BIOSYNTHESISIN THE SOFT CORAL *GERSEMIA FRUTICOSA* K. Varvas, R. Koljak,

I. Jarving, T. Pehk', N. Samel', Institute of Chemistry, Akadeemia tee 15,

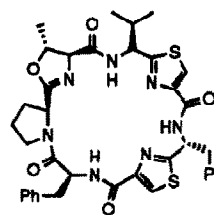
EE0026 Tallinn, Estonia, *Institute of Chemical Physics and Biophysics, R vala pst. 10, EE0100, Tallinn, Estonia.

PGG₂ has been found to be a key intermediate in the prostaglandin biosynthesis in the coral *Gersemia fruticosa*.**TOTAL SYNTHESIS OF LISSOCLINAMIDE 5, A CYTOTOXIC CYCLIC PEPTIDE FROM THE TUNICATE *LISSOCLINUM PATELLA***

Christopher Boden and Gerald Pattenden*

Department of Chemistry, The University, Nottingham NG7 2RD, England

A total synthesis of lissoclinamide 5, and some of its stereoisomers, shows that its stereostructure should be revised.

**ALKENYLSULFENYLCHLORIDES : SYNTHESIS AND Ad_E REACTIONS OF 2-ALKOXY-2-OXO-3-R-4-CHLOROTHIO-1,2-OXAPHOSPHOL-3-ENES.**Igor V. Alabugin^a, Valery K. Brel^{*a}, Anatoly N. Chekhlov^a, Nikolai S. Zefirov^{*a},Peter J. Stang^b.

Institute of Physiologically Active Compounds, Russian Academy of Science, Chernogolovka, Moscow Region, 142432,

Russia. FAX: (095) 939 0290^a ; Department of Chemistry, University of Utah, Salt Lake City, U84112, USA^b.