

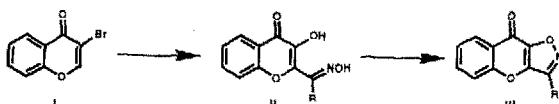
GRAPHICAL ABSTRACTS

Tetrahedron Lett. 1992, 33, 993

**THE SYNTHESIS AND CHEMISTRY OF FUNCTIONALIZED FUROCHROMONES.<sup>4</sup>  
ADDITION OF NITRONATE ANIONS TO 3-BROMOCHROMONE AND 6-BROMOFUROCHROMONE.  
AN EXPEDIENT ROUTE TO FURO(3',2':6,7)-BENZOPYRANO(2,3-d)-1ISOXAZOLONES  
AND CHROMONO(2,3-d)1ISOXAZOLONES.**

Ronald B. Gammill,\* Sharon A. Nash and William Watt  
Upjohn Laboratories, The Upjohn Company, Kalamazoo, MI 49001 USA

Addition of nitronate anions to I yield products containing functionalized  $\gamma$ -pyrone rings.

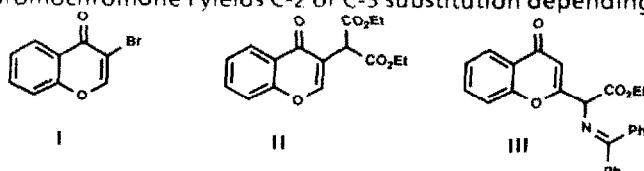


Tetrahedron Lett. 1992, 33, 997

**A NOVEL ENTRY TO SUBSTITUTED CHROMONES AND FUROCHROMONES THROUGH CYCLOPROPANE INTERMEDIATES**

Ronald B. Gammill,\* Sharon A. Nash, Larry T. Bell, and William Watt  
Upjohn Laboratories, The Upjohn Company, Kalamazoo, MI 49001 USA

Addition of carbon nucleophiles to bromochromone I yields C-2 or C-3 substitution depending on the nucleophilic species.

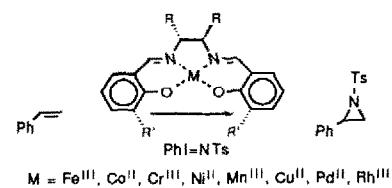


Tetrahedron Lett. 1992, 33, 1001

**ALKENE AZIRIDINATION AND EPOXIDATION CATALYZED BY CHIRAL METAL SALEN COMPLEXES.** Kenneth J. O'Connor

Shiow-Jyi Wey and Cynthia J. Burrows,\* Department of Chemistry, State University of New York at Stony Brook, Stony Brook, NY 11794-3400, USA.

Metal(salen) complexes have been compared as catalysts for alkene aziridination and epoxidation. While several metal complexes catalyzed epoxidation, only the Mn<sup>III</sup> complex catalyzed aziridination, and this occurred without enantioselectivity.

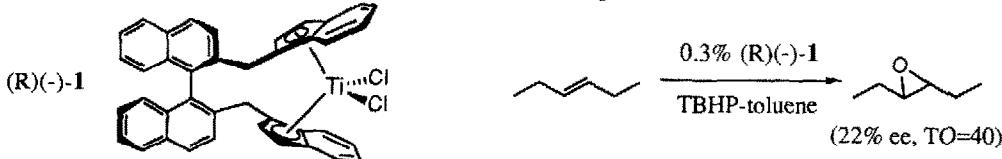


Tetrahedron Lett. 1992, 33, 1005

**Asymmetric Epoxidation of Unfunctionalized Alkenes Using the New C<sub>2</sub>-Symmetrical 1,1'-Binaphthyl-2,2'-Dimethylene-Bridged *ansa*-Bis(1-indenyl)Titanium Dichloride Catalyst**

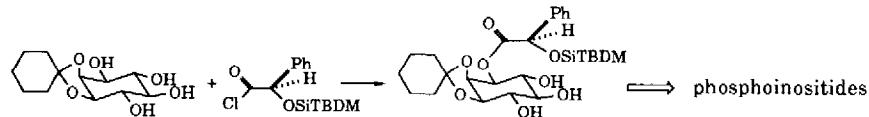
Steven L. Colletti and Ronald L. Halterman\*

Department of Chemistry, University of Oklahoma, 620 Parrington Oval, Norman, Oklahoma 73019-0370



Karol S. Bruzik\*, Jeffrey Myers and Ming-Daw Tsai

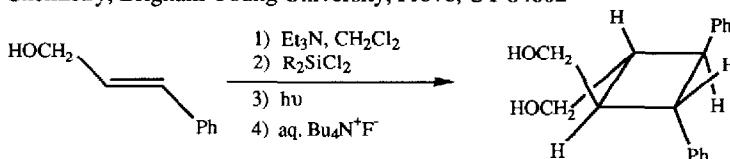
Department of Chemistry, The Ohio State University, Columbus, OH 43210

Enantiomerically pure *myo*-inositol derivatives are synthesized using regiospecific protection of cyclohexylidene-*myo*-inositol with a chiral mandelic acid-derived acyl group.

## STEREOCONTROLLED PHOTOCHEMICAL [2 + 2] CYCLOADDITION

Steven A. Fleming\* and Susan C. Ward

Department of Chemistry, Brigham Young University, Provo, UT 84602



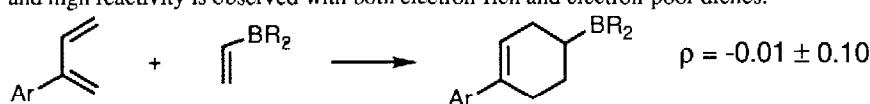
## VINYLBORANES ARE OMNIPHILIC DIENOPHILES.

## SOME UNUSUAL AND USEFUL PROPERTIES OF VINYLBORANES IN DIELS-ALDER REACTIONS

Daniel A. Singleton,\* Jose P. Martinez, and José V. Watson

Department of Chemistry, Texas A&amp;M University, College Station, Texas 77843, USA

The rate of Diels-Alder reactions with vinyl-9-BBN is uniquely insensitive to diene substituent effects, and high reactivity is observed with both electron-rich and electron-poor dienes.

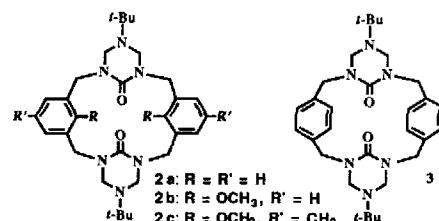
Synthesis and Structures of a New Class of Calixarene Analogs Derived from 5-*tert*-Butyltetrahydro-1,3,5-triazine-2(1H)one.P.R. Dave,\*<sup>a</sup> G. Doyle,<sup>a</sup> T. Axenrod,<sup>b</sup> H. Yazdekhasti<sup>b</sup> and H.L. Ammon<sup>c</sup><sup>a</sup>GEO-CENTERS, INC. at ARDEC,

762 Route 15 South, Lake Hopatcong, NJ 07849

<sup>b</sup>Department of Chemistry, The City College of CUNY, New York, NY 10031<sup>c</sup>Department of Chemistry and Biochemistry, Univ. of Maryland,

College Park, MD 20742

Calixarene analogs 2a-c and 3 have been synthesized and their structures investigated by x-ray crystallography and NMR spectroscopy.



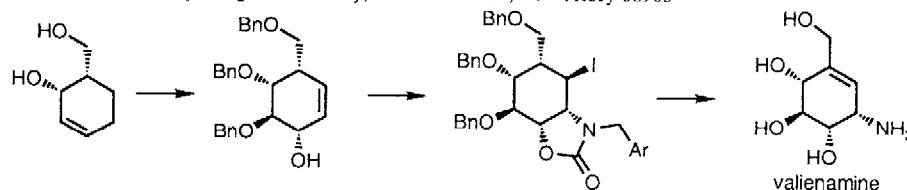
INTRAMOLECULAR AMINO DELIVERY REACTIONS

Tetrahedron Lett. 1992, 33, 1025

FOR THE SYNTHESIS OF VALIENAMINE AND ANALOGUES

Spencer Knapp,\* Andrew B. J. Naughton, and T. G. Murali Dhar

Department of Chemistry, Rutgers University, New Brunswick, New Jersey 08903

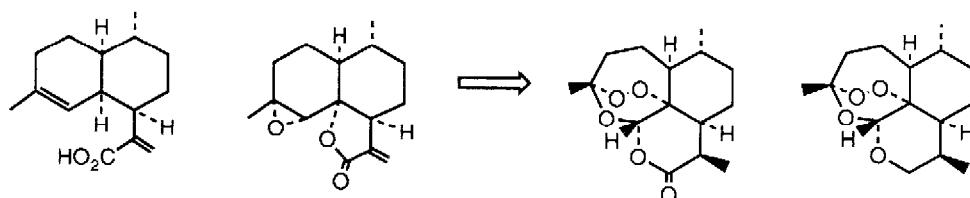


AN EFFICIENT PARTIAL SYNTHESIS OF  
(+)-ARTEMISININ AND (+)-DEOXOARTEMISININ

Tetrahedron Lett. 1992, 33, 1029

Peter T. Lansbury\* and Deanne M. Nowak

Department of Chemistry, State University of New York at Buffalo, Buffalo N. Y. 14214



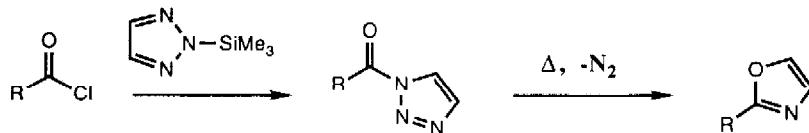
A GENERAL, ONE-POT METHOD FOR THE  
SYNTHESIS OF 2-SUBSTITUTED OXAZOLES

Tetrahedron Lett. 1992, 33, 1033

Eric L. Williams

Monsanto Company, Mail Zone T4J, 800 N. Lindbergh, St. Louis, MO 63167

The high yield conversion of an acid chloride to a 2-substituted oxazole has been accomplished through the intermediate 1,2,3-triazole amide.



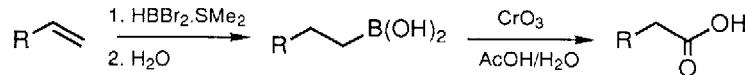
A CONVENIENT PROCEDURE FOR THE DIRECT  
CONVERSION OF TERMINAL ALKENES INTO  
CARBOXYLIC ACIDS

Tetrahedron Lett. 1992, 33, 1037

Uday S. Racherla, Vijay V. Khanna and Herbert C. Brown\*

H. C. Brown and R. B. Wetherill Laboratories of Chemistry, Purdue University, West Lafayette, IN 47907

The oxidation of alkylboronic acids with  $\text{CrO}_3$  in 90 % aqueous acetic acid directly provides carboxylic acids in 80-97 % isolated yields.

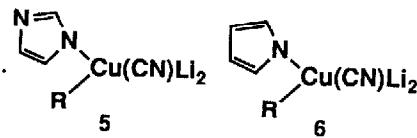


HIGHER ORDER, MIXED CYANOCUPRATES DERIVED FROM N-LITHIO-IMIDAZOLE AND PYRROLE: NEW "DUMMY" LIGAND ALTERNATIVES IN ORGANOCOPPER CHEMISTRY

*Tetrahedron Lett.* 1992, 33, 1041

B.H. Lipshutz,\* P. Fatheree, W. Hagen, and K.L. Stevens  
Department of Chemistry, University of California, Santa Barbara, CA 93106

Reactions and virtues of cuprates **5** and **6** are discussed.

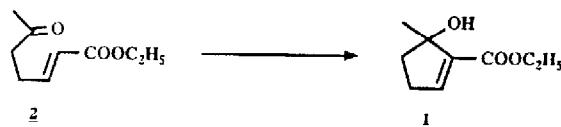


### AN INTRAMOLECULAR BAYLIS-HILLMAN REACTION

*Tetrahedron Lett.* 1992, 33, 1045

Fides Roth, Peter Gygax, Georg Fräter\*  
GIVAUDAN-ROURE, Dübendorf, Switzerland

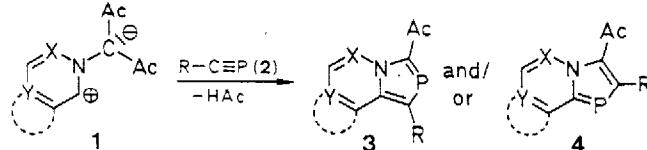
By means of an intramolecular Baylis-Hillman reaction the cyclopentenol derivative **1** has been formed from **2**



### A NEW ACCESS TO PHOSPHAINDOLIZINES BY [3+2] CYCLOADDITION OF AZOMETHINE YLIDES ONTO PHOSPHAALKYNES

*Tetrahedron Lett.* 1992, 33, 1049

Uwe Bergsträßer, Andreas Hoffmann and Manfred Regitz,  
Fachbereich Chemie der Universität Kaiserslautern,  
Erwin-Schrödinger-Str.,  
D-6750 Kaiserslautern, Deutschland

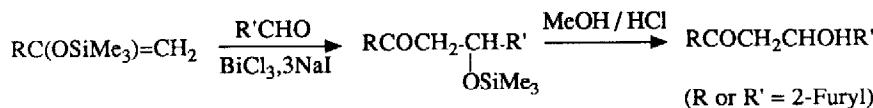


Azomethine ylides of type **1** add onto the triple bond of phosphaalkynes **2** followed by elimination of HAc to yield the phosphaindolizines **3** and/or **4**.

*Tetrahedron Lett.* 1992, 33, 1053

### REACTION D'ALDOLISATION CROISEE EN SERIE FURANIQUE PAR L'INTERMEDIAIRE D'ENOLS SILICIES.

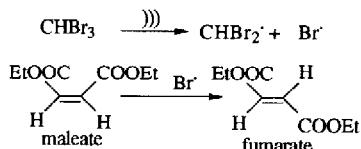
C. Le Roux, M. Maraval<sup>o</sup>, M.E. Borredon<sup>o</sup>, H. Gaspard-Iloughmane et J. Dubac\*  
Laboratoire des Organométalliques, Université Paul-Sabatier, 118, route de Narbonne,  
31062 Toulouse Cedex et <sup>o</sup>Laboratoire de Chimie des Agroressources, E.N.S.C.T.,  
118, route de Narbonne, 31077 Toulouse Cedex (FRANCE).



## PERFLUOROCARBONS AS INERT GASES IN HOMOGENEOUS SONOCHEMISTRY.

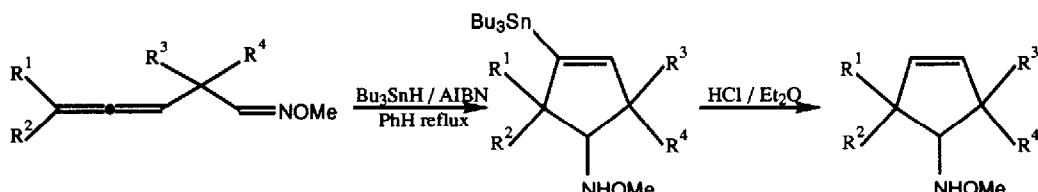
Th. Lepoint<sup>a\*</sup>, F. Mullie<sup>a</sup>, N. Voglet<sup>a</sup>, D.H. Yang<sup>b</sup>, J. Vandercammen<sup>b</sup>, J. Reisse<sup>b\*</sup><sup>a</sup>Institut Meurice, CERIA, 1, avenue E. Gryson, B-1070 Bruxelles; <sup>b</sup>Université Libre de Bruxelles, (CP165), 50, avenue Franklin Roosevelt, B-1050 Bruxelles, Belgique.

The polyatomic inert gases  $\text{CF}_4$  and  $\text{C}_2\text{F}_6$  are efficient in homogeneous sonochemistry. This observation leads to the conclusion that some analogies could exist between sonochemistry and plasma chemistry.

RADICAL CYCLIZATION OF  $\beta$ -ALLENIC OXIME ETHERS

J. Hatem\*, C. Henriet-Bernard, J. Grimaldi and R. Maurin

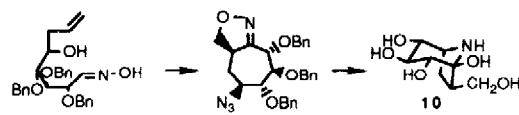
Laboratoire de Chimie Organique de Synthèse, associé au CNRS, Université de Provence, Case 541, 13397 Marseille Cedex 13, France



## Access to Polyhydroxylated Cycloheptane Derivatives by Stereoselective Nitrile Oxide Intramolecular Cycloaddition. Synthesis of an Analogue of Calystegine B2

O. Duclos, A. Duréault\*, J.C. Depazay. Université René Descartes, Laboratoire de Chimie et Biochimie Pharmacologiques et Toxicologiques, associé au CNRS, 45 rue des Saints-Pères, 75270 Paris cedex 06, France.

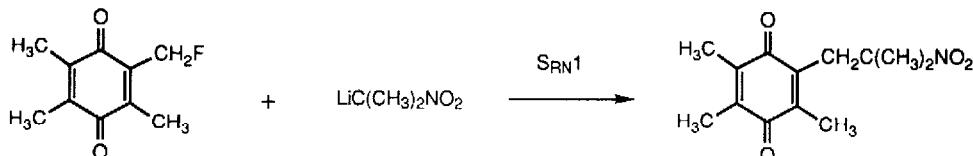
The polyhydroxylated nortropane 10 is synthesized from an olefinic C-8 acyclic carbohydrate oxime.

FLUORIDE AS LEAVING GROUP IN  $S_{RN}1$  REACTIONS

## OF A TETRASUBSTITUTED-1,4-BENZOQUINONE

Michel P. CROZET\*, Luc GIRAUD, Jean-François SABUCO and Patrice VANELLE.

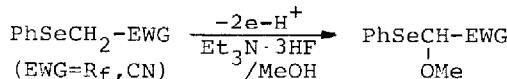
Laboratoire de Chimie Organique B, associé au CNRS, BP 562, 13397 Marseille Cedex 13, France

The C-alkylation reaction of 2-fluoromethyl-3,5,6-trimethyl-1,4-benzoquinone by 2-nitropropane anion is shown to proceed by the  $S_{RN}1$  mechanism.

Kazimierz Surowiec and Toshio Fuchigami\*

Department of Electronic Chemistry, Tokyo Institute of Technology,  
Midori-ku, Yokohama 227, Japan

Anodic  $\alpha$ -methoxylation of 1,1-dihydroperfluoroalkyl and cyanomethyl selenides was successfully carried out in the presence of  $\text{Et}_3\text{N} \cdot 3\text{HF}$  as a supporting electrolyte.

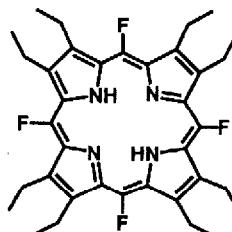


### ***meso*-Perfluorination of Porphyrins with *N*-Fluoropyridinium Triflate**

Y. Naruta, F. Tani, K. Maruyama

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

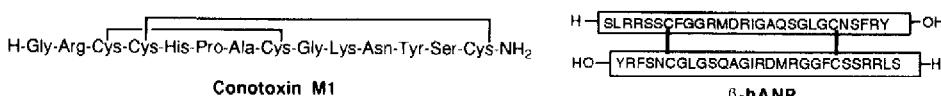
Octaethylporphyrin was efficiently perfluorinated at its *meso*-positions with *N*-fluoropyridinium triflate.



## REGIOSELECTIVE DOUBLE DISULFIDE FORMATION USING SILYLCHLORIDE-SULFOXIDE SYSTEM

Kenichi Akaii, Kenji Fujino, Tadashi Tatsumi, and Yoshiaki Kiso

**Department of Medicinal Chemistry, Kyoto Pharmaceutical University, Yamashina-ku, Kyoto 607, Japan.**

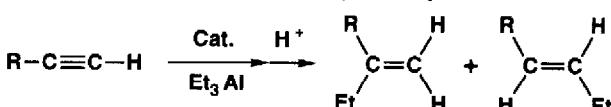


Two intramolecular or intermolecular disulfide bonds of conotoxin M1 or  $\beta$ -hANP are formed regioselectively by the combination of silylchloride-sulfoxide method with conventional oxidation methods.

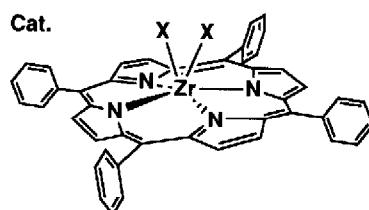
## Zirconium Porphyrins as Novel Active Catalysts for Highly Regio- and Stereo-selective Ethylalumination of Alkynes

Kenji Shibata, Takuzo Aida and Shohei Inoue\*

Kenji Sasaki, Takuji Aida and Shigeru Inoue  
Department of Synthetic Chemistry, Faculty of Engineering  
University of Tokyo, Hongo, Bunkyo-ku, Tokyo 113, Japan

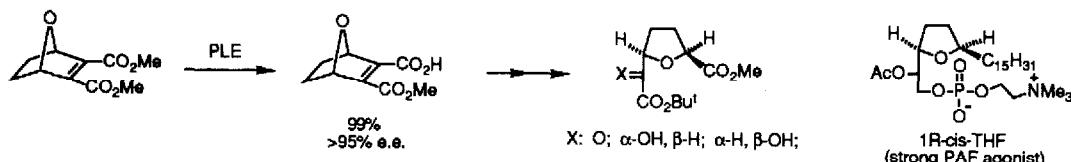


More than 99 % regioselectivities with satisfactorily high turnover numbers were attained when R =  $n$ C<sub>3</sub>H<sub>7</sub>, tC<sub>4</sub>H<sub>9</sub> and  $n$ C<sub>5</sub>H<sub>11</sub>.

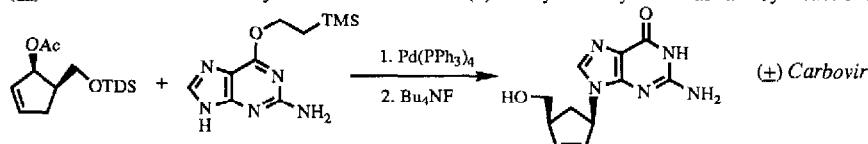
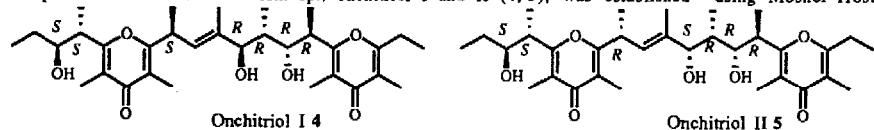
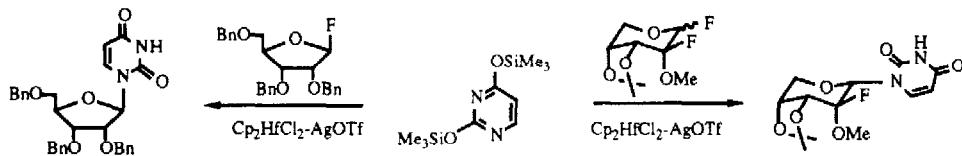


**Development of Tetrahydrofuran Chiral Synthons by Enzymatic Approach:****Improved Synthesis of a Strong Agonist of Platelet Activating Factor**

S. Kobayashi\*, M. Sato, Y. Eguchi, and M. Ohno, Faculty of Pharmaceutical Sciences, University of Tokyo, Hongo, Bunkyo-ku, Tokyo 113, Japan

**Pd(0)-Catalyzed Allylic Alkylation in the Synthesis of (±)Carbovir.**Lise-Lotte Gundersen, Tore Benneche\* and Kjell Undheim  
Department of Chemistry, University of Oslo, N-0315 Oslo 3, Norway.

(±) Carbovir has been synthesized with a Pd(0)-catalyzed allylation as a key reaction.

**NEW MARINE CYTOTOXIC BISPYRONES. ABSOLUTE STEREOCHEMISTRY OF THE ONCHITRIOLS I AND II**Jaime Rodríguez, Ricardo Riguera\*  
Depart. de Química Orgánica, Fac. de Química, Univ. de Santiago de Compostela, Santiago de Compostela. 15706. España  
Cécile Debitus  
Centre ORSTOM, B. P. A5. Nouméa Cedex. Nouvelle Calédonie.The complete absolute stereochemistry of two new cytotoxic marine polypropionates isolated from the saponified extract of the pulmonate mollusc *Onchidium* sp., onchitriol I and II (4, 5), was established using Mosher-Trost's methodology.**STEREOSELECTIVE SYNTHESIS OF NUCLEOSIDES BY METALLOCENE PROMOTED ACTIVATION OF GLYCOSYL FLUORIDES**M.I. Matheu, R. Echarri, S. Castillón\*  
Departament de Química, Facultat de Ciències Químiques de Tarragona, Universitat de Barcelona  
Pça. Imperial Tarraco 1. 43005 Tarragona. Spain

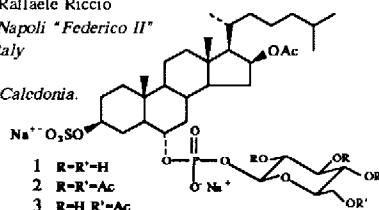
The First Occurrence of Polyhydroxylated Steroids with Phosphate  
Conjugation from the Starfish *Tremaster novaecaledonicae*

Francesco De Riccardis, Maria Iorizzi, Luigi Minale\* and Raffaele Riccio  
Dipartimento di Chimica delle Sostanze Naturali, Università di Napoli "Federico II"  
Via Domenico Montesano 49, 80131 Napoli, Italy

Cecile Debitus

ORSTOM, Centre de Nouméa, BP. A5, Nouméa, New Caledonia.

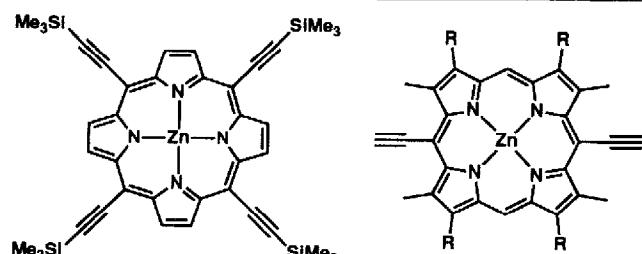
The first occurrence of three polyhydroxylated steroids with phosphate conjugation is reported from the starfish *Tremaster novaecaledonicae*. The structures were characterized by  $^1\text{H}$ ,  $^{13}\text{C}$  and  $^{31}\text{P}$  NMR and chemical degradation.



*Meso*-Alkynyl Porphyrins

Harry L. Anderson

University Chemical Laboratory,  
Lensfield Road, Cambridge,  
CB2 1EW (UK)

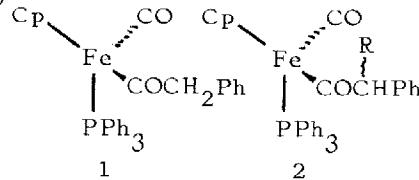


BASE-DEPENDENT STEREOSELECTIVITY IN  
REACTIONS OF ACYL LIGAND IN PHENYLACETYL-  
IRON COMPLEX ( $\eta^5\text{-C}_5\text{H}_5$ )Fe(CO)(PPh<sub>3</sub>)(COCH<sub>2</sub>Ph)

Zhong-Wu Guo and Aleksander Zamojski

Institute of Organic Chemistry, Polish Academy of  
Sciences, 01-224 Warszawa, Poland

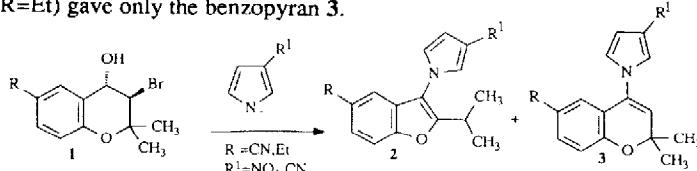
Proportion of stereoisomeric products 2 obtained from  
the complex 1 depends on the base used for generation  
of the anion.



AN UNUSUAL BASE-MEDIATED RING CONTRACTION REACTION OF  
BENZOPYRANS TO BENZOFURANS

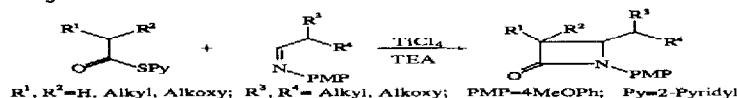
D.R.Buckle,\* A.Faller, I.L.Pinto and D.G.Smith SmithKline Beecham Pharmaceuticals, Yew Tree  
Bottom Road, Epsom, Surrey, KT18 5XQ.

**1** (R=CN) with electron deficient pyrrole and pyrazole anions gave benzofurans **2**. Corresponding reaction of **1** (R=Et) gave only the benzopyran **3**.



**HIGHLY DIASTEREOSELECTIVE SYNTHESIS OF  $\beta$ -LACTAMS BY  
ADDITION OF TITANIUM ENOLATES OF 2-PYRIDYL THIOESTERS TO CHIRAL IMINES.**  
R. Annunziata, M. Cinquini\*, F. Cozzi\*, P.G. Cozzi  
Dipartimento di Chimica Organica e Industriale - Università di Milano - Italy.

Addition of titanium enolates of 2-pyridyl thioesters to chiral imines gives  $\beta$ -lactams with good diastereofacial control.



**BIFUNCTIONAL CHIRAL AUXILIARIES 4: ALKYLATION OF  
ENOLATES DERIVED FROM 1,3-DIACYL-*TRANS*-4,5-TETRAMETHYLENEIMIDAZOLIDIN-2-ONES**

S.G. Davies\* and A.A. Mortlock, The Dyson Perrins Laboratory, South Parks Road, Oxford, OX1 3QY, U.K.

