

## GRAPHICAL ABSTRACTS

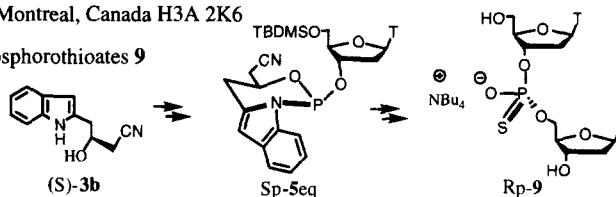
### A Stereoselective Synthesis of Dinucleotide Phosphorothioates, Using Chiral Indol-oxazaphosphorine Intermediates

*Tetrahedron Letters*, 1997, 38, 3797

Jian-Chao Wang and George Just\*

Department of Chemistry, McGill University, Montreal, Canada H3A 2K6

A stereoselective synthesis of dinucleotide phosphorothioates **9** with a de of > 98%, using (S) and (R)-**3b** as chiral auxiliaries, is reported. The procedure may be adaptable to solid phase synthesis.

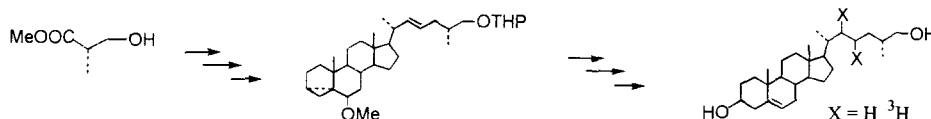


### Synthesis of (25R)-Cholest-5-ene-3 $\beta$ , 26-diol and its Radiolabeled Analog.

*Tetrahedron Letters*, 1997, 38, 3801

Thomas E. D'Ambra<sup>a</sup> Norman B. Javitt,<sup>b</sup> Koji Nakanishi,<sup>c</sup> Tadeusz Warchol<sup>c,\*</sup>

<sup>a</sup> Albany Molecular Research, Inc., 21 Corporate Circle, Albany, N. Y. 12203, <sup>b</sup>New York University, Medical Center, 550 First Avenue, N. Y. 10016, <sup>c</sup>Columbia University in the City of New York, Department of Chemistry, N.Y. 10027.



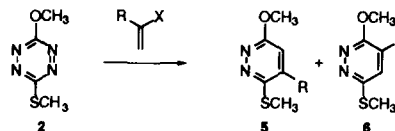
### Preparation and Inverse Electron Demand Diels-Alder

*Tetrahedron Letters*, 1997, 38, 3805

#### Reactions of 3-Methoxy-6-methylthio-1,2,4,5-tetrazine.

Subas M. Sakya\*, Kelley K. Groskopf<sup>†</sup>, Chemical Sciences, Infectious Disease Subdivision, Wyeth-Ayerst Research, Pearl River, New York 10965; Dale L. Boger, Department of Chemistry, The Scripps Research Institute, 10550 North Torrey Pines Road, La Jolla, California 92037.

Synthesis and preliminary investigation into the participation of **2** in [4+2] cycloaddition reactions with electron rich and neutral dienophiles along with the resulting regioselective formation of the products (**5**) are discussed.



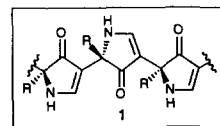
### A SECOND-GENERATION SYNTHESIS OF SCALEMIC 3,5,5-TRISUBSTITUTED PYRROLIN-4-ONES: INCORPORATION OF FUNCTIONALIZED AMINO ACID SIDE-CHAINS

*Tetrahedron Letters*, 1997, 38, 3809

Amos B. Smith, III,\* Andrew B. Benowitz, David A. Favor, Paul A. Sprengeler, and Ralph Hirschmann\*

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

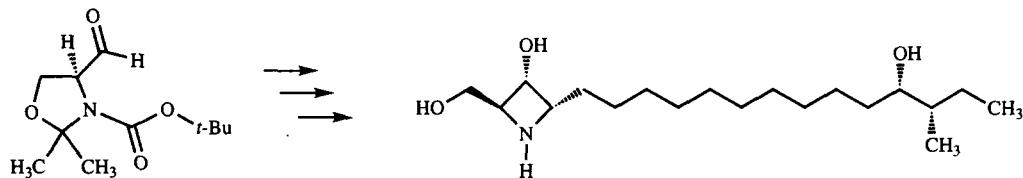
**Abstract:** To access mimics of peptidal  $\beta$ -strands (**1**), scalemic 3,5,5-trisubstituted pyrrolin-4-ones bearing the tyrosine, serine, and lysine side-chains have been generated via cyclization of metalated imino esters and deprotection. The functionalized imino esters were prepared by asymmetric alkylation of a common oxazolidinone precursor (**2**) derived from L-prenylglycine.



*Tetrahedron Letters*, 1997, 38, 3813

**STEREOSELECTIVE SYNTHESIS OF PENARESIDIN A AND RELATED**

**AZETIDINE ALKALOIDS.** Spencer Knapp\* and Yaohua Dong, Department of Chemistry, Rutgers  
The State University of New Jersey, New Brunswick, New Jersey 08903 USA

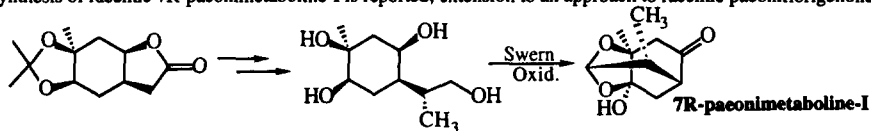


*Tetrahedron Letters*, 1997, 38, 3817

**STEREOSELECTIVE SYNTHESIS OF 7R-PAEONIMETABOLINE-I;  
STEREOSELECTIVE ROUTE TOWARDS PAEONIFLORIGENONE**

David P. Richardson,\* Priscilla W. Carr, Jared N. Cumming, Warren G. Harbison, Nina D. Raoof, Michelle S. Sanders, Eliza Shin,  
Thomas E. Smith, and Thomas H. Wintner, Department of Chemistry, Williams College Williamstown, MA 01267 USA

A synthesis of racemic 7R-paeonimetaboline-I is reported; extension to an approach to racemic paeoniflorigenone is also summarized.

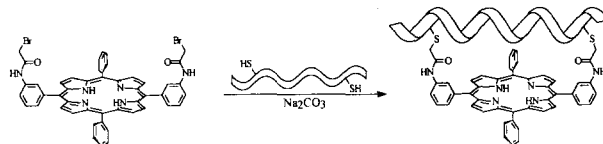


*Tetrahedron Letters*, 1997, 38, 3821

**THE DESIGN, SYNTHESIS, AND CHARACTERIZATION OF A  
PORPHYRIN-PEPTIDE CONJUGATE.**

G. Richard Geier III and Tomikazu Sasaki,\* Department of Chemistry, University of Washington, Seattle, WA 98195 USA

The design and synthesis of a novel peptide strapped porphyrin assembly is described. CD spectroscopy and <sup>1</sup>H NMR are supportive of the designed structure.

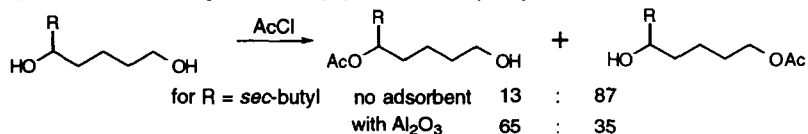


*Tetrahedron Letters*, 1997, 38, 3825

**ACETYLATION OF UNSYMMETRICAL DIOLS IN THE  
PRESENCE OF Al<sub>2</sub>O<sub>3</sub>.**

Gary W. Breton\*, Melissa J. Kurtz, Sharyn L. Kurtz  
Berry College, Department of Chemistry, Mount Berry, GA 30149-5016 USA

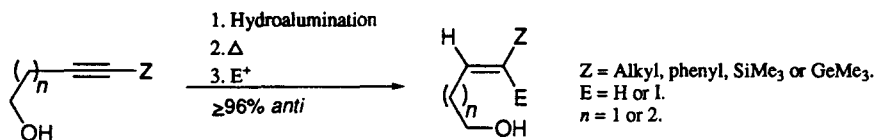
Acetylation of unsymmetrical diols in the presence of Al<sub>2</sub>O<sub>3</sub> affords sterically congested monoacetates.



**Anti-Hydroalumination of Homo- and Bishomopropargyl Alcohols**

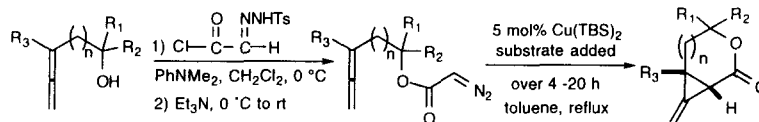
Shengming Ma, Fang Liu, and Ei-ichi Negishi

Department of Chemistry, Purdue University, W. Lafayette, IN 47907, USA

**SYNTHESIS AND REACTIONS OF 1,2-DISUBSTITUTED METHYLENOCYCLOPROPANES PREPARED VIA INTRA-MOLECULAR CYCLOPROPANATION OF ALLENIC DIAZO-ACETATES.**

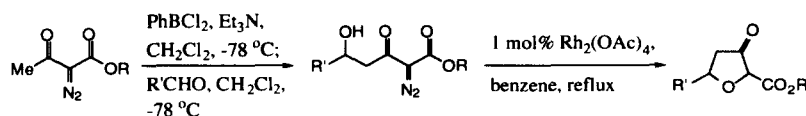
Mark Lautens,\* Christophe Meyer, and Arjan van Oeveren Department of Chemistry, University of Toronto, Toronto, Ontario, Canada M5S 3H6

1,2-Disubstituted methylenecyclopropanes were prepared via a copper catalyzed diazo decomposition reaction.

**ALDOL-CYCLIZATION REACTION SEQUENCE FOR THE SYNTHESIS OF TETRAHYDROFURANS.**

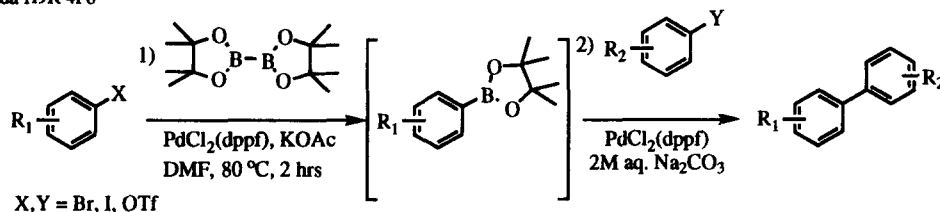
Michael A. Calter\*, Priyantha M. Sugathapala and Cheng Zhu, Department of Chemistry, Virginia Tech, Blacksburg, VA 24061-0212

The combination of an aldol and an insertion reaction affords a novel synthesis of tetrahydrofurans.

**ONE POT BIARYL SYNTHESIS VIA IN SITU BORONATE FORMATION**

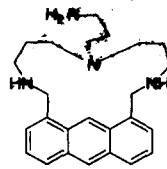
André Giroux\*, Yongxin Han and Petpiboon Prasit

Department of Medicinal Chemistry, Merck-Frosst Centre for Therapeutic Research P.O. Box 1005, Pointe-Claire-Dorval, Québec, Canada H9R 4P8

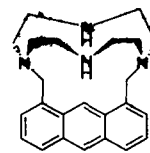


**A Fluorescent Chemosensor Signaling Only Single and Double Bond Cleavage**, Norman S. Opler, David M. Vance, and Christopher J. Chang, Department of Chemistry, Ohio State University, Columbus, OH 43210

Two fluorescent chemosensors **5** and **7** are described. Chemosensor **5** reacts with aldehydes and ketones (R<sub>1</sub>, R<sub>2</sub>) in 100% aqueous solution. On the other hand, a rigid-bridged cyclic derivative (**7**) showed no effect on fluorescence by any metal ion examined. A novel approach to the attainment of binding discrimination in chemosensors, based upon the rigid immobilization of polyamine ligands onto a fluorophore framework, is discussed.



5



7

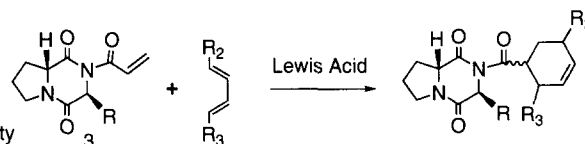
**(2,5-dihetopiperazine) New Chiral Auxiliaries for Asymmetric Diels-Alder Reactions**

Thuy X. H. Le, Jacqueline C. Bussolari and William V. Murray\*

The R.W. Johnson Pharmaceutical Research Institute

1000 Route 202, Raritan NJ 08869

Cyclo-S-phenylalanine -R-proline has been shown to be an excellent chiral auxiliary for Diels-Alder reactions. Reactions of dienes with **3** proceed in high chemical yield, high endo selectivity and excellent diastereoselectivity.

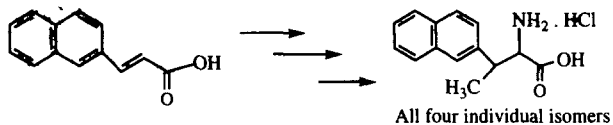


**Asymmetric Synthesis of Unusual Amino Acid: Synthesis of Four Isomers of β-Methyl-3-(2'-Naphthyl)alanine**

Wei-Yuan and Victor J. Stoltz\*

Department of Chemistry, The University of Arizona, Tucson, Arizona 85721

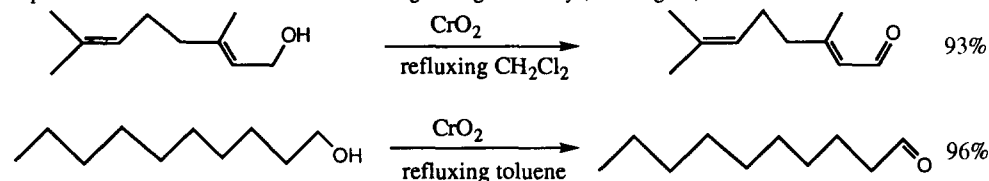
The asymmetric synthesis of all four individual isomer of 3-methyl-3-(2'-naphthyl)alanine guided by an Evans-type chiral auxiliary is reported.



All four individual isomers

**MAGTRIEVE™: AN EFFICIENT, MAGNETICALLY RETRIEVABLE AND RECYCLABLE OXIDANT.** Ross Lee<sup>a</sup> and Dennis Donald<sup>a,b</sup>,

Dupont Films<sup>a</sup> and DuPont Central Science and Engineering Laboratory<sup>b</sup>, Wilmington, Delaware

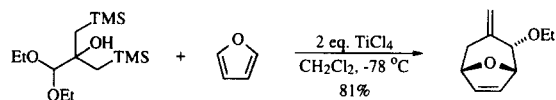
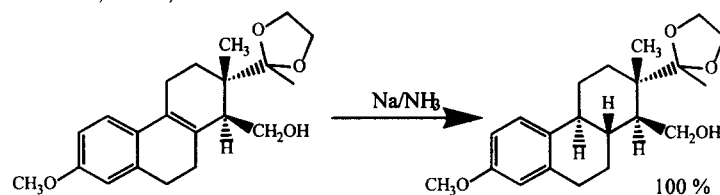


**HETEROATOM-STABILIZED ALLYLIC CATIONS IN 4+3  
CYCLOADDITIONS. A TANDEM PETERSON OLEFINATION/4+3 CYCLOADDITION REACTION**

Michael Harmata\* and Darin E. Jones

Department of Chemistry, University of Missouri-Columbia, Columbia, Missouri 65211

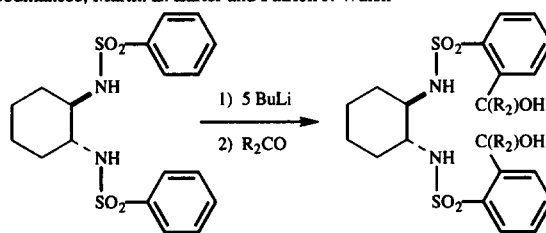
A tandem Peterson olefination/4+3 cycloaddition reaction has been established.


**Neighbouring Group Directed Protonation in the Birch**
**Reduction of Styrene Double Bonds** Zhen Lin,<sup>§</sup> Jinhua Chen,<sup>§\*</sup>
and Zdenek Valenta<sup>†</sup> Department of Chemistry, University of New Brunswick, Bag Service 45222, Fredericton, New Brunswick, Canada, E3B 6E2
**Synthesis of New Chiral Tetradentate Sulfonamide Based Ligands.**

David E. Ho, Juan M. Betancort, David H. Woodmansee, Martin L. Larter and Patrick J. Walsh\*

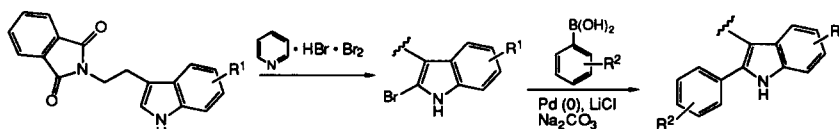
Department of Chemistry, San Diego State University, San Diego, Ca 92182-1030, U.S.A.

The key step in the synthesis of novel tetradentate bisulfonamide based ligands is a directed metallation followed by quenching with either ketones or carbon dioxide.


**Synthesis of 2-Aryltryptamines with Palladium Catalyzed  
Cross-Coupling of 2-Bromotryptamines and Arylboronic Acids**

Lin Chu\*, Michael H. Fisher, Mark T. Goulet, and Matthew J. Wyvratt, Department of Medicinal Chemistry, Merck Research Laboratories, P.O. Box 2000, Rahway, New Jersey 07065 USA

A high-yielding synthesis of 2-aryltryptamines employing palladium(0) catalyzed cross-coupling of 2-bromotryptamines and arylboronic acids is reported.

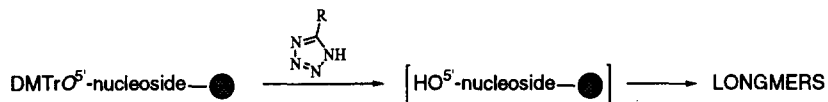


*Tetrahedron Letters*, 1997, 38, 3875

### ON THE FORMATION OF LONGMERS IN PHOSPHOROTHIOATE OLIGODEOXYRIBONUCLEOTIDE SYNTHESIS

Achim H. Krotz\*, Patrick G. Klopchin, Kathleen L. Walker, G. Susan Srivatsa, Douglas L. Cole and Vasulinga T. Ravikumar, Isis Pharmaceuticals, Inc., 2292 Faraday Avenue, Carlsbad, CA 92008, USA

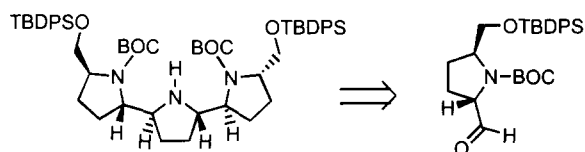
The formation of longmers [(n+1)-, (n+2)-mers, etc.] in phosphorothioate oligodeoxyribonucleotide synthesis through phosphoramidite coupling chemistry occurs in a base-, activator- and contact time-dependent manner.



*Tetrahedron Letters*, 1997, 38, 3879

### Stereoselective Synthesis of a Terpyrrolidine Unit, a Potential Building Block for Anion Recognition.

Hans-Dieter Arndt, Kurt Polborn and Ulrich Koert\*, Institut für Organische Chemie der LMU, Karlstraße 23, D-80333 München.

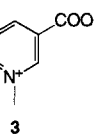
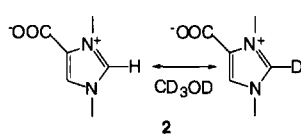
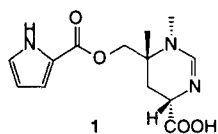


Bidirectional acetylide coupling of two *trans*-pyrrolidine aldehydes derived from *S*-pyroglutamic acid delivers enantiomerically pure terpyrrolidine via a 1,4-cyclic sulfate-ring closure sequence.

*Tetrahedron Letters*, 1997, 38, 3883

### MANZACIDIN D: AN UNPRECEDENTED SECONDARY METABOLITE FROM THE "LIVING FOSSIL" SPONGE *ASTROSCLEERA WILLEYANA*

Thorsten Jahn, Gabriele M. König\*, Anthony D. Wright, Institute for Pharmaceutical Biology, Technical University of Braunschweig, Mendelssohnstraße 1, 38106 Braunschweig, Germany, Gert Wörheide and Joachim Reitner, Institute and Museum for Geology and Palaeontology, University of Göttingen, Goldschmidt Straße 3, 37077 Göttingen, Germany



The methanol extract of *Astrosclera willeyana* was found to contain the unprecedented natural product manzacidin D (1), norzoanemonin (2), and trigonelline (3).

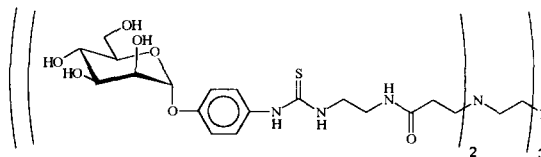
*Tetrahedron Letters*, 1997, 38, 3885

### GLYCODENDRIMER SYNTHESIS WITHOUT USING

PROTECTING GROUPS. Christoffer Kieburg and Thisbe K. Lindhorst\*,

University of Hamburg, Department of Organic Chemistry, Martin-Luther-King-Platz 6, D-20146 Hamburg, Germany.

Thiourea-bridged glycodendrimers were obtained by the reaction of unprotected NCS-functionalized carbohydrate derivatives and branched oligoamines in aqueous solution.



*Tetrahedron Letters*, 1997, 38, 3889

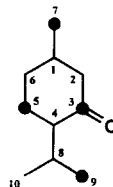
### Monoterpene Essential Oils are Not of Mevalonoid Origin

Wolfgang Eisenreich<sup>§</sup>, Silvia Sagner<sup>\*</sup>, Meinhard H. Zenk<sup>\*</sup> and Adelbert Bacher<sup>§\*</sup>

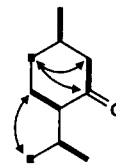
<sup>§</sup>Institut für Organische Chemie und Biochemie, Technische Universität München, Lichtenbergstraße 4, D-85747 Garching, Germany and <sup>\*</sup>Lehrstuhl für Pharmazeutische Biologie, Universität München, Karlstraße 29, D-80333 München, Germany

Incorporation of [1-<sup>13</sup>C] and [U-<sup>13</sup>C]<sub>6</sub>]glucose shows that monoterpene essential oils (geraniol, menthone, pulegone, thymol) are biosynthesised in plants by a pathway which is different from the established mevalonic acid route.

[1-<sup>13</sup>C]glucose



[U-<sup>13</sup>C]<sub>6</sub>]glucose

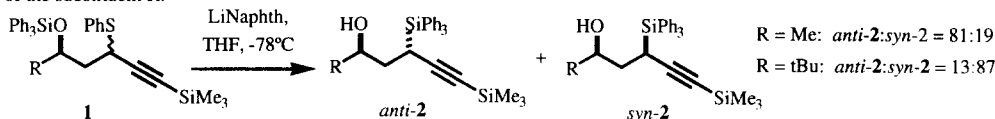


*Tetrahedron Letters*, 1997, 38, 3893

### 1,3-ASYMMETRIC INDUCTION IN RETRO-[1,4]-BROOK REARRANGEMENTS OF PROPARGYLLITHIUM COMPOUNDS TO PROPARGYLSILANES

Frank Samtleben<sup>§</sup>, Mathias Noltemeyer<sup>#</sup>, and Reinhard Brückner<sup>\*§</sup> <sup>§</sup>Institut für Organische Chemie der Georg-August-Universität, Tammannstr. 2, D-37077 Göttingen, Germany <sup>#</sup>Institut für Anorganische Chemie der Georg-August-Universität, Tammannstr. 4, D-37077 Göttingen, Germany

$\gamma$ -Siloxyated propargyl phenyl sulfides **1** were lithiated reductively to provide propargyllithium intermediates which rearranged to propargylsilanes **2**. Whether the  $\gamma$ -C-O bond and the newly formed C-Si bond are preferentially *anti*- or *syn*-oriented depends on the size of the substituent R.



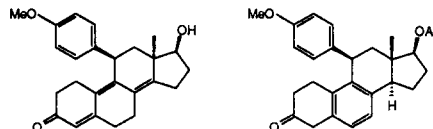
*Tetrahedron Letters*, 1997, 38, 3897

### SYNTHESIS AND SPECTROSCOPIC PECULIARITIES OF 11 $\beta$ -ARYL-ESTRA-4,8(14),9-TRIENES AND 11 $\beta$ -ARYL-ESTRA-5(10),6,8-TRIENES

Günter Neef,<sup>\*</sup> Arwed Cleve, Gisbert Depke, and Emil Eckle

Research Laboratories of Schering AG, D-13342 Berlin, Germany

Two new types of 11 $\beta$ -aryl-substituted steroids are synthesized. The 11 $\beta$ -phenyl ring shows signs of restricted rotation at ambient temperature.

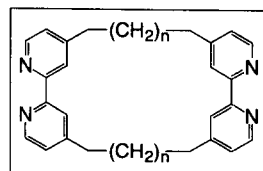


*Tetrahedron Letters*, 1997, 38, 3901

### BIPYRIDINE BASED EXODITOPIC LIGANDS : SYNTHESIS AND STRUCTURAL ANALYSIS OF HOMOBINUCLEAR RUTHENIUM COMPLEXES

Christian Kaes, Mir Wais Hosseini<sup>\*</sup>, André De Cian, Jean Fischer  
Institut de Chimie, Université Louis Pasteur, 4, rue Blaise Pascal, F-67000 Strasbourg, France

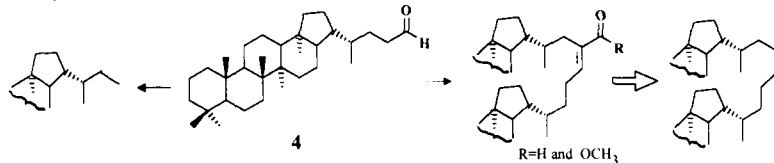
The synthesis of new macrocyclic ligands based on 2,2'-bipyridine unites interconnected at the 4 and 4' positions by alkyl chains was achieved. Homobinuclear Ru<sup>II</sup> complexes were formed and the solid state structure of one of the diastereoisomers was investigated by X-ray analysis.



**GEOMIMETIC AUTOXIDATION OF BIOHOPANOIDS :  
A ROUTE TO BIS-HOPANOIDS, POTENTIAL NEW SEDIMENTARY MOLECULAR FOSSILS.**

Philippe Bisseret,\* Anne Baron, Claude Rodier, Serge Neunlist, Ecole Nationale Supérieure de Chimie de Mulhouse, 3 rue A. Werner, 68093 Mulhouse, France; Michel Rohmer, Institut Le Bel, Université Louis Pasteur, 4 rue Blaise Pascal, 67070 Strasbourg, France.

Representative biohopanoid derivative **4** was submitted to geomimetic autoxidations in the presence of molecular sieves in benzene, giving access to bis-hopanoids, precursors to potential new molecular fossils, as well as under heating without solvent to give the saturated *nor*-hydrocarbon.



**THE BIS-LINKING OF TETRATHIAFULVALENE (TTF) TO C<sub>60</sub> :  
TOWARDS THE CONTROL OF ELECTRON TRANSFER BETWEEN  $\pi$ -DONORS AND C<sub>60</sub>**

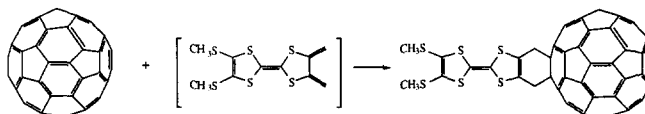
C. Boulle,<sup>ab</sup> J.M. Rabreau,<sup>a</sup> P. Hudhomme,<sup>a\*</sup> M. Cariou,<sup>b</sup> M. Jubault,<sup>b</sup> A. Gorgues,<sup>b\*</sup> J. Orduna,<sup>c</sup> and J. Garin<sup>c</sup>

<sup>a</sup> Laboratoire de Synthèse Organique, associé au CNRS, Université de Nantes, 2 rue de la Houssinière, F-44322 Nantes

<sup>b</sup> Ingénierie Moléculaire et Matériaux Organiques, associé au CNRS, Université d'Angers, 2 Boulevard Lavoisier, F-49045 Angers

<sup>c</sup> Departamento de Química Orgánica, Universidad de Zaragoza-CSIC, E-50009 Zaragoza

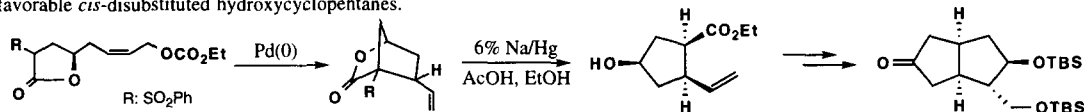
The bis-linking of a tetrathiafulvalene (TTF) to C<sub>60</sub> is carried out using the [4+2] cycloaddition between a 2,3-dimethylene-TTF and C<sub>60</sub>.



**THE HIGHLY DIASTEREOSELECTIVE PALLADIUM-CATALYZED  
CYCLIZATIONS. STEREOSELECTIVE SYNTHESSES OF *cis* AND**

***trans*-DISUBSTITUTED HYDROXYCYCLOPENTANES.** Young-Ger Suh\*, Jae-Kyung Jung, Soon-Ai Kim, Dong-Yun Shin and Kyung-Hoon Min. College of Pharmacy, Seoul National University, Seoul, 151-742, Korea.

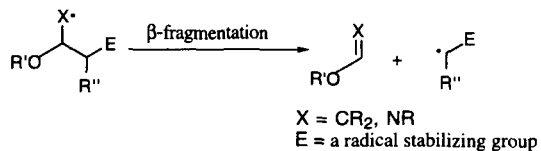
A new variant of the palladium-catalyzed cyclization provides high 1,2-diastereocontrol of two new stereogenic centers as well as 1,3-asymmetric induction. Desulfonylation of the cyclized product also affords the thermodynamically less favorable *cis*-disubstituted hydroxycyclopentanes.



**THE EFFECT OF  $\alpha$ -ALKOXY GROUP IN RADICAL-MEDIATED  
 $\beta$ -FRAGMENTATION REACTIONS**

Sunggak Kim,\* Kwan Hee Kim, and Jin Rai Cho, Department of Chemistry, Korea Advanced Institute of Science and Technology, Taejeon 305-701, Korea

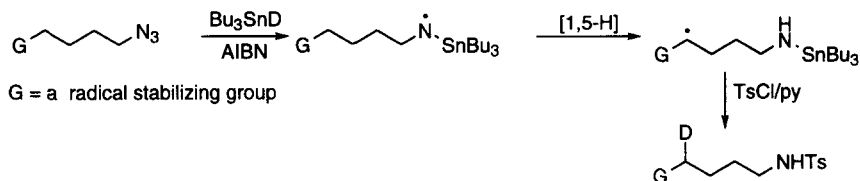
$\beta$ -Fragmentation reactions of alkyl and aminyl radicals were greatly facilitated by the presence of  $\alpha$ -alkoxy groups.





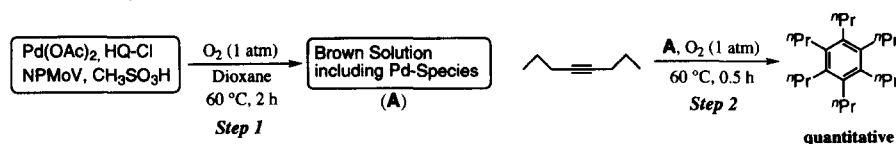
### 1,5-HYDROGEN TRANSFERS FROM CARBON TO *N*-TRIBUTYLTIN SUBSTITUTED NITROGEN

Sunggak Kim,\* Kyu Man Yeon, and Kwang Sub Yoon, Department of Chemistry, Korea Advanced Institute of Science and Technology, Taejeon 305-701, Korea



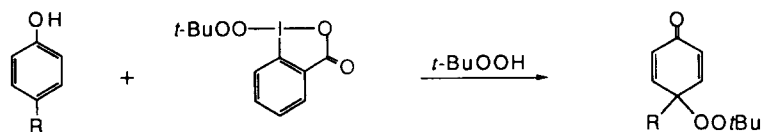
**Cyclotrimerization of Alkynes Using a Multicatalytic System, Pd(II) / Chloroquinone / NPMoV, Under Dioxigen.** Takahiro Yokota, Yasunori Sakurai, Satoshi Sakaguchi, Yasutaka Ishii\*, Department of Applied Chemistry, Faculty of Engineering & High Technology Research Center, Kansai University, Suita Osaka 564, Japan

Cyclotrimerization of alkynes was performed using a new triple catalytic system, Pd(II) / chloroquinone / NPMoV, under atmospheric oxygen.



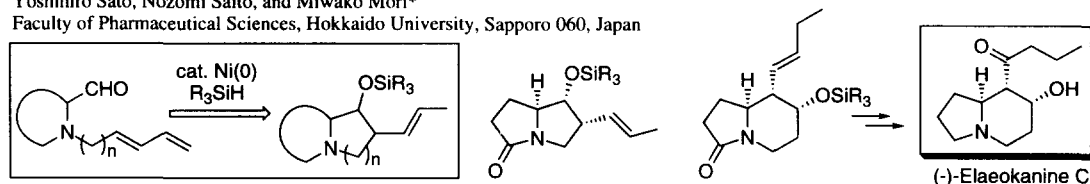
### FREE-RADICAL OXIDATION OF PARA-SUBSTITUTED PHENOLS BY HYPERVALENT TERT-BUTYLPEROXYIODANE AND TERT-BUTYL HYDROPEROXIDE: SYNTHESIS OF 4- (TERT-BUTYLPEROXY)-2,5-CYCLOHEXADIEN-1-ONES

Masahito Ochiai,\* Akinobu Nakanishi, and Akiko Yamada  
Faculty of Pharmaceutical Sciences, University of Tokushima, 1-78 Shomachi, Tokushima 770, Japan



### EFFICIENT SYNTHESIS OF PYRROLIZIDINE AND INDOLIZIDINE DERIVATIVES USING NICKEL-CATALYZED CYCLIZATION OF 1,3-DIENE AND ALDEHYDE: FORMAL TOTAL SYNTHESIS OF (-)-ELAEOKANINE C

Yoshihiro Sato, Nozomi Saito, and Miwako Mori\*  
Faculty of Pharmaceutical Sciences, Hokkaido University, Sapporo 060, Japan

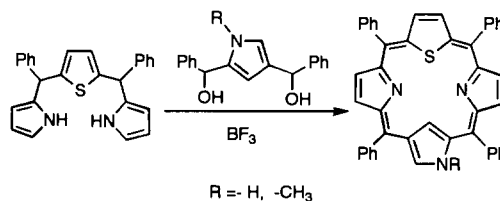


*Tetrahedron Letters*, 1997, 38, 3935

### Synthesis of meso-Tetraphenylthiaporphyrins Bearing One Inverted Pyrrole

Han-Je Kim and Chang-Hee Lee\*  
Department of Chemistry, Kangwon National University,  
Chun-Cheon 200-701, Korea

2-Aza-21-thia-23-carba-5,10,15,20-tetraphenylporphyrin and its N-methylated analogue were synthesized and characterized. The synthesis utilizes [3+1] condensation of thiatripyrryn with 2,4-bis( $\alpha$ -hydroxy- $\alpha$ -phenylmethyl)pyrrole or 2,4-bis( $\alpha$ -hydroxy- $\alpha$ -phenylmethyl)-N-methylpyrrole in the presence of acid catalyst. This convenient process gives predominantly the titled porphyrin. Reversion of the pyrrole unit is observed by prolonging the reaction time.

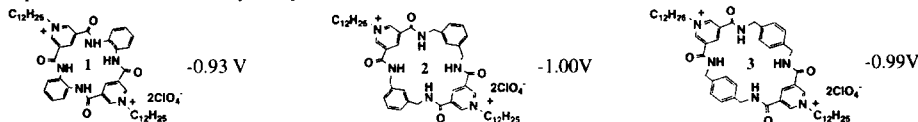


*Tetrahedron Letters*, 1997, 38, 3939

### SYNTHESES AND REDOX BEHAVIOR OF NOVEL CYCLIC HOSTS HAVING MULTIPLE REDOX CENTERS OF NAD<sup>+</sup> ANALOGUE

Yasuhisa Kuroda,\*<sup>†</sup> Hidenori Seshimo, Toshio Kondo, Masana Shiba and Hisanobu Ogoshi  
<sup>†</sup>Department of Polymer Science, Kyoto Institute of Technology, Sakyo-ku, Matsugasaki, Kyoto 606, Japan  
Department of Synthetic Chemistry and Biological Chemistry, Kyoto University, Sakyo-ku, Kyoto 606-01, Japan

A new series of cyclophanes having 3,5-dicarbamoylpyridinium moieties, **1**, **2** and **3**, were synthesized and their one-electron reduction potentials are determined by the cyclic voltammetric method.

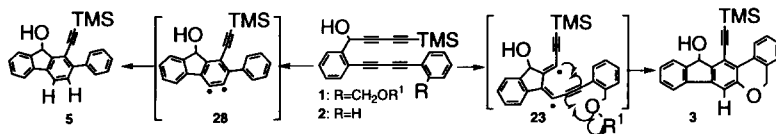


*Tetrahedron Letters*, 1997, 38, 3943

### Cycloaromatization of a Non-Conjugated Polyenyne System: Synthesis of 5H-Benzo[d]fluoreno[3,2-b]pyrans via Diradicals Generated from 1-[2-(4-(2-Alkoxyethylphenyl)butan-1,3-diynyl)]phenyl-pentan-2,4-diyn-1-ols and Trapping Evidence for the 1,2-Didehydrobenzene Diradical

Kazuhiro Miyawaki, Riho Suzuki, Tomikazu Kawano, and Ikuro Ueda\*, The Institute of Scientific and Industrial Research, Osaka University, Mihogaoka, Ibaraki, Osaka 567, Japan

Tetraynes **1** and **2** undergo thermal intramolecular cyclization at 25 °C to provide new polycyclic aromatic compounds **3** and **5**.

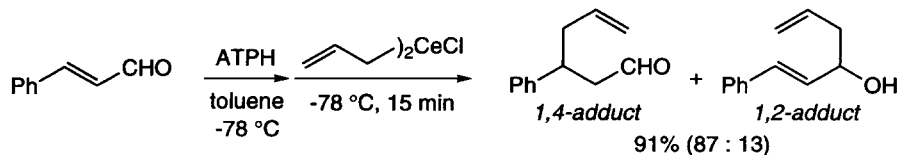


*Tetrahedron Letters*, 1997, 38, 3947

### CONJUGATE ALLYLATION TO $\alpha,\beta$ -UNSATURATED ALDEHYDES WITH AMPHIPHILIC ALKYLATION SYSTEM, ATPH/ALLYLCERIUM REAGENTS

Takashi Ooi, Tomoya Miura, Yuichiro Kondo, and Keiji Maruoka\*  
Department of Chemistry, Graduate School of Science, Hokkaido University, Sapporo 060, Japan

Conjugate allylation to  $\alpha,\beta$ -unsaturated aldehydes has been successfully accomplished with a new amphiphilic alkylation system, aluminum tris(2,6-diphenylphenoxy) (ATPH)/allylcerium reagents.

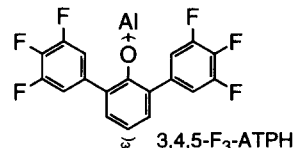
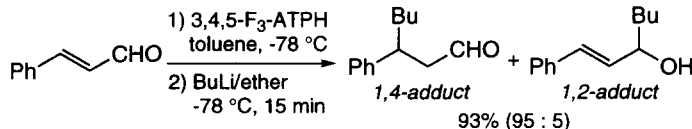


**EVALUATION OF SEVERAL FLUORINATED ATPH DERIVATIVES AS FUNCTIONALIZED LEWIS ACID RECEPTOR FOR CONJUGATE ALKYLATION TO  $\alpha,\beta$ -UNSATURATED ALDEHYDES WITH ALKYL LITHIUM NUCLEOPHILES**

Takashi Ooi, Yuichiro Kondo, Tomoya Miura, and Keiji Maruoka\*

Department of Chemistry, Graduate School of Science, Hokkaido University, Sapporo 060, Japan

Several fluorinated aluminum tris(2,6-diphenylphenoxide) (ATPH) derivatives have been synthesized to evaluate the conjugate alkylation ability to  $\alpha,\beta$ -unsaturated aldehydes by the combined use of alkyl lithium nucleophiles.

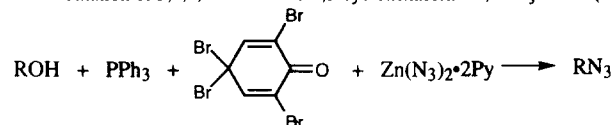


**AN EFFICIENT METHOD FOR CONVERTING ALCOHOLS TO AZIDES WITH 2,4,4,6-TETRABROMO-2,5-CYCLOHEXADIENONE/PPh<sub>3</sub>/Zn(N<sub>3</sub>)<sub>2</sub>·2Py**

Akiko Saito, Kazumi Saito, Akira Tanaka,\* and Takayuki Oritani

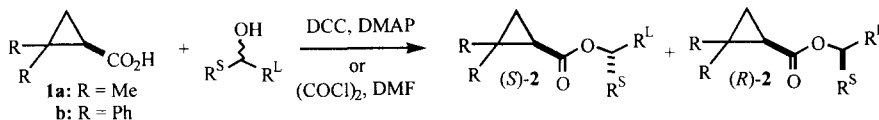
Division of Environmental Bioremediation, Graduate School of Agriculture, and Department of Applied Biological Chemistry, Faculty of Agriculture, Tohoku University, 1-1 Tsutsumidori-amamiyamachi, Aoba-ku, Sendai 981, Japan

A combination of 2,4,4,6-tetrabromo-2,5-cyclohexadienone, PPh<sub>3</sub> and Zn(N<sub>3</sub>)<sub>2</sub>·2Py converted alcohols to azides in excellent yields.



**CHIRAL 2,2-DISUBSTITUTED CYCLOPROPANECARBOXYLIC ACIDS: EFFECTIVE DERIVATIZING AGENTS FOR ANALYSIS OF ENANTIOMERIC PURITY OF ALCOHOLS AND FOR RESOLUTION OF 1,1'-BI-2-NAPHTHOL.**

Hyun Chang Kim, Soojin Choi, Hongdoo Kim, Kwang-Hyun Ahn,\* Department of Chemistry, Kyunghee University, Yongin 449-701, Republic of Korea; Jeong Hwan Koh, and Jaiwook Park\* Department of Chemistry, Center for Biofunctional Molecules, POSTECH, Pohang 790-784, Republic of Korea

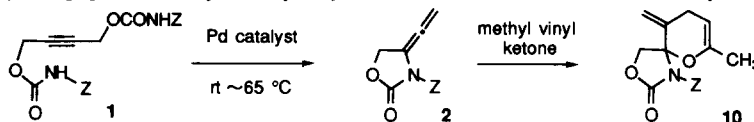


**EFFICIENT SYNTHESIS OF 4-ETHENYLIDENE-2-OXAZOLIDINONES VIA PALLADIUM-CATALYZED AMINOCYCLIZATION OF 2-BUTYN-1,4-DIOL BISCARBAMATES**

Masanari Kimura, Yoshinori Wakamiya, Yoshikazu Horino, and Yoshinao Tamaru\*

Department of Applied Chemistry, Faculty of Engineering, Nagasaki University, 1-14 Bunkyo, Nagasaki 852, Japan

The enamine moiety of 2, prepared from 1 by Pd-catalyzed cyclization, is reactive toward unsaturated carbonyls and provides, e.g., 10.



**9-ADAMANTYLIDENE-10-ALKYL-9,10-DIHYDROACRIDINE OXIDE AS A NOVEL AND POTENTIAL CHEMILUMINESCENT PRECURSOR: PROXIMITY EFFECT OF THE NITROGEN ATOM**

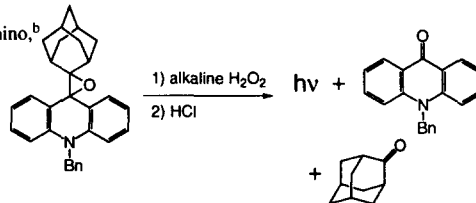
*Tetrahedron Letters*, 1997, 38, 3967

Takeshi Imanishi,<sup>a\*</sup> Yohko Ueda,<sup>a</sup> Masatoshi Minagawa,<sup>a</sup> Nobuhiro Hoshino,<sup>b</sup> and Kazuyuki Miyashita<sup>a</sup>

<sup>a</sup> Faculty of Pharmaceutical Sciences, Osaka University, 1-6 Yamadaoka, Suita, Osaka 565, Japan

<sup>b</sup> Iatron Laboratories, Inc., 1460-6 Mitodai, Takocho, Katori-gun, Chiba 289-22, Japan

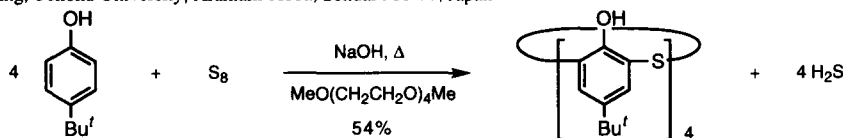
The oxirane was found to show chemiluminescence by treatment with alkaline H<sub>2</sub>O<sub>2</sub> and subsequent with HCl.



**FACILE SYNTHESIS OF P-TERT-BUTYLTHIACALIX[4]ARENE BY THE REACTION OF P-TERT-BUTYLPHENOL WITH ELEMENTAL SULFUR IN THE PRESENCE OF A BASE.**

*Tetrahedron Letters*, 1997, 38, 3971

Hitoshi Kumagai,<sup>a\*</sup> Mitsuharu Hasegawa,<sup>a</sup> Setsuko Miyanari,<sup>a</sup> Yoshihiro Sugawa,<sup>a</sup> Yoko Sato,<sup>a</sup> Takashi Hori,<sup>a</sup> Sanae Ueda,<sup>a</sup> Hiroki Kamiyama,<sup>a</sup> and Sotaro Miyano,<sup>a,b</sup> <sup>a</sup>Chemical Technology Laboratory, Cosmo Research Institute, Satte, Saitama, 340-01, Japan, <sup>b</sup>Department of Biochemistry and Engineering, Faculty of Engineering, Tohoku University, Aramaki-Aoba, Sendai 980-77, Japan

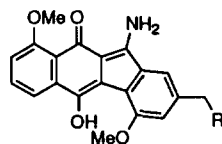


**TOTAL SYNTHESIS OF O<sup>6,9</sup>-DIMETHYL STEALTHINS A AND C**

*Tetrahedron Letters*, 1997, 38, 3973

Hiroya Koyama and Tadao Kamikawa\*, Faculty of Science and Technology, Department of Chemistry, Kinki University, Kowakae, Higashi-osaka, Osaka 577, Japan

O<sup>12</sup>-Acetyl-O<sup>4,9</sup>-dimethylstealthin A and O<sup>4,9</sup>-dimethylstealthin C were synthesized using the Suzuki coupling reaction as a key step.



O<sup>4,9</sup>-dimethylstealthin C R = H

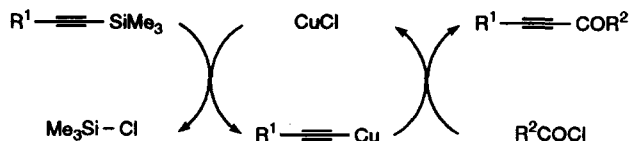
O<sup>12</sup>-acetyl-O<sup>4,9</sup>-dimethylstealthin A R = OAc

**DIRECT ALKYNYL GROUP TRANSFER FROM SILICON TO COPPER: NEW PREPARATION METHOD OF ALKYNYL COPPER (I) REAGENTS**

*Tetrahedron Letters*, 1997, 38, 3977

Hajime Ito, Kikuo Arimoto, Hiro-omi Sensui, and Akira Hosomi\*  
Department of Chemistry, University of Tsukuba, Tsukuba, Ibaraki 305, Japan

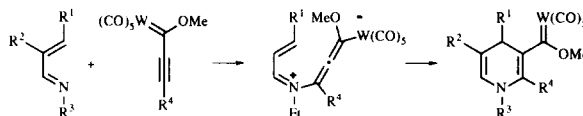
A first observation of the direct alkyne group transfer from silicon to copper is reported. The silicon group of alkynylsilanes was smoothly replaced by copper (I) chloride in DMI to give the corresponding copper (I) acetylides. This transformation was applied to the synthesis of alkyne ketones catalyzed by CuCl.



**FIRST [4+2] CYCLOADDITION OF ALKYNYL FISCHER CARBENE COMPLEXES WITH HETERODIENES. FACILE SYNTHESIS OF 1,4-DIHYDROPYRIDINES FROM 1-AZADIENES.**

José Barluenga,\* Miguel Tomás, José A. López-Peigrín and Eduardo Rubio. Instituto Universitario de Química Organometálica "Enrique Moles", Universidad de Oviedo, Julián Clavería 8, 33071- Oviedo

Neutral 1-azadienes react in a [4 + 2] fashion with alkynyl Fischer carbene complexes to afford regioselectively substituted 1,4-dihydropyridines.

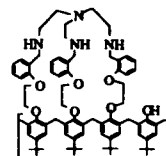


**Synthesis of the Tripodal-Amine Capped Benzo Crown *p-tert*-Butylcalix[4]arene and Its Host-Guest Chemistry**

Thawatchai Tuntulani\*, Vithaya Ruangpornvisuti, Nonglak Tantikunwathana, Onanong Ngampaiboonsombut and Ratana Seangprasertkij-Magee, Department of Chemistry, Chulalongkorn University, Bangkok 10330, Thailand

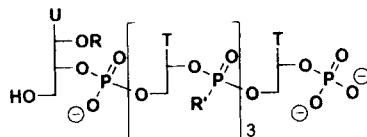
Zouhair Asfari and Jacques Vicens, E.C.P.M., 1 rue Blaise Pascal, Strasbourg, France

The tripodal-amine capped benzo crown *p-tert*-butylcalix[4]arene has been synthesised, and complexation studies of the compound with Zn(II) salts are also reported.



**A NOVEL SOLID SUPPORT FOR SYNTHESIS OF 3'-PHOSPHORYLATED CHIMERIC OLIGONUCLEOTIDES CONTAINING INTERNUCLEOSIDIC METHYL PHOSPHOTRIESTER AND METHYLPHOSPHONATE LINKAGES.**

Andrei Guzaev and Harri Lönnberg, Department of Chemistry, University of Turku, FIN-20014 Turku, Finland



R = H, Fmp  
R' = Me, OMe