

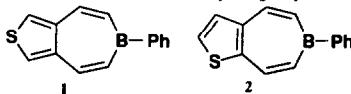
## GRAPHICAL ABSTRACTS

### Thienoborepins: Conjugation Characteristics in Boron Heterocycles

*Tetrahedron*, 1994, 50, 6495

Yoshikazu Sugihara,<sup>a\*</sup> Ryuta Miyatake,<sup>a</sup> Toshiyasu Yagi,<sup>a</sup> Ichiro Murata,<sup>a</sup> Mamoru Jinguji,<sup>b</sup> Tomoo Nakazawa,<sup>b</sup> and Akira Imamura<sup>c</sup>

*Department of Chemistry, Faculty of Science, Osaka University, Toyonaka Osaka 560, Japan,<sup>a</sup> Department of Chemistry, Medical University of Yamanashi, Tamaho, Nakakoma, Yamanashi 408-38, Japan<sup>b</sup>, Department of Chemistry, Faculty of Science, Hiroshima University, Kagamiyama, Higashi-Hiroshima 724, Japan.<sup>c</sup>*



Protonolysis, complex-formation with amines, ab initio molecular orbital calculations, redox potentials, and spectroscopic features of 1-phenylthieno-[3,4-d]borepin (**1**) and 1-phenylthieno[2,3-d]borepin (**2**) were examined.

*Tetrahedron*, 1994, 50, 6505

### Unprecedented Stereochemical Control in the Intramolecular Ene-Reactions of $\delta,\epsilon$ -Unsaturated Aldehydes Using Exceptionally Bulky Organoaluminum Reagents: Elucidation of the Transition State

Takashi Ooi, Keiji Maruoka, and Hisashi Yamamoto\*

*School of Engineering, Nagoya University Chikusa, Nagoya 464-01, Japan*

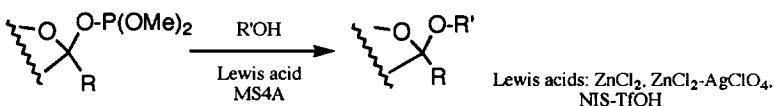


### GLYCOSYLATION USING GLYCOSYL PHOSPHITE AS A GLYCOSYL DONOR

*Tetrahedron*, 1994, 50, 6523

Yutaka Watanabe,\* Chikara Nakamoto, Takashi Yamamoto, and Shoichiro Ozaki

*Department of Applied Chemistry, Faculty of Engineering, Ehime University, Matsuyama 790, Japan*



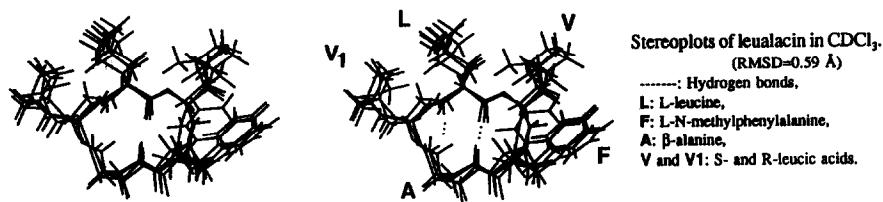
### Conformational Analysis of a New Cyclic Depsipeptide Calcium Blocker, Leualacin, by NMR Spectroscopy

*Tetrahedron*, 1994, 50, 6537

Keiko Yoda, Hideyuki Haruyama\*, Harumitu Kuwano, Kiyosi Hamano\*, Kazuhiko Tanzawa\*

Analytical and Metabolic Research Laboratories, Biomedical Research Laboratories,

\* Biological Research Laboratories, Sankyo Co., Ltd, 1-2-58 Hiromachi, Shinagawa-ku, Tokyo 140, Japan



THE INTRODUCTION OF NITRILE GROUPS INTO HETEROCYCLES AND CONVERSION OF CARBOXYLIC GROUPS INTO THEIR CORRESPONDING NITRILES WITH CHLOROSULFONYLISOCYANATE AND TRIETHYLAMINE

H. Vorbrüggen and K. Krolikiewicz  
Research Laboratories, Schering AG, D-13342 Berlin, Germany



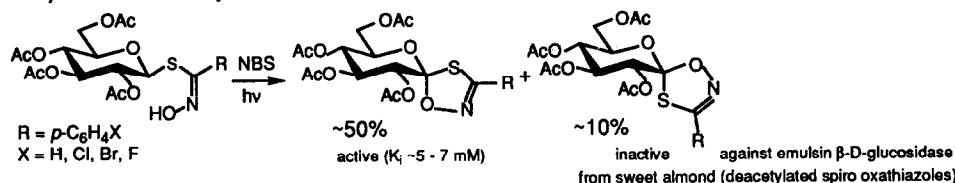
Nucleophilic unsaturated systems as well as carboxylic acids react acc. to R. Graf with chlorosulfonylisocyanate to N-chlorosulfonylamides, which are readily converted by triethylamine in high overall yields into the corresponding nitriles.

Synthesis, Structure and Enzymatic Evaluation of New Spiro Oxathiazole Sugar Derivatives

J.-P. Praly<sup>a</sup>, R. Faure<sup>b</sup>, B. Joseph<sup>c</sup>, L. Kiss<sup>d</sup> and P. Rollin<sup>c</sup>

a, b - Université Lyon I, 69622-Villeurbanne, France; c - Université d'Orléans, 45067- Orléans, France

d - Lajos Kossuth University of Debrecen, POB 55, H-4010 - Debrecen, Hungary

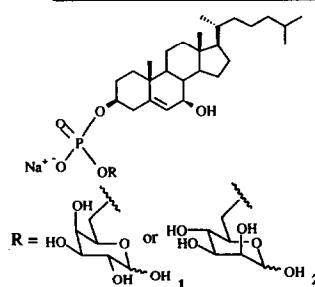


Synthesis of Phosphoric Acid Diesters of 7 $\beta$ -hydroxy-cholesterol and of Carbohydrates.

X. Pannecoucke, G. Schmitt and B. Luu\*.

Laboratoire de Chimie Organique des Substances Naturelles, URA CNRS,  
Centre de Neurochimie, 5 rue Blaise Pascal, 67084 Strasbourg, France.

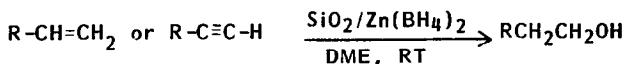
The sodium salt of 3 $\beta$ (7 $\beta$ -hydroxycholesterol) 6(carbohydrate) monophosphate (1 or 2) has been synthesized using both the phosphoramidite and the hydrogen phosphonate methods.



SILICA GEL SUPPORTED ZINC BOROHYDRIDE. A NOVEL REAGENT FOR HYDRATION OF UNACTIVATED ALKENES AND ALKYNES

Brindaban C. Ranu, Arunkanti Sarkar, Manika Saha and Rupak Chakraborty  
Department of Organic Chemistry, Indian Association for the  
Cultivation of Science, Jadavpur, Calcutta - 700 032, India.

Regioselective hydration of unactivated alkenes and alkynes has been achieved by the action of zinc borohydride on the corresponding alkene or alkyne on silica gel support.

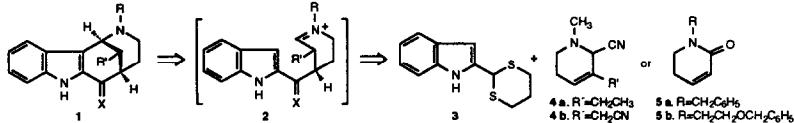


SYNTHETIC APPLICATIONS OF 2-(1,3-DITHIAN-2-YL)INDOLES. IV. NEW SYNTHESIS  
OF THE TETRACYCLIC ABED RING SYSTEM OF *STRYCHNOS* ALKALOIDS

*Tetrahedron*, 1994, 50, 6585

Anna Diez,<sup>a</sup> Josep Castells,<sup>a</sup> Pilar Forme,<sup>a</sup> Mario Rubiralta,<sup>a\*</sup> David S. Grierson,<sup>b</sup>  
Henri-Philippe Husson,<sup>b</sup> Xavier Solana,<sup>c</sup> and Mercè Font-Bardia<sup>c</sup>

a. Laboratory of Organic Chemistry, Faculty of Pharmacy, University of Barcelona, 08028 Barcelona, Spain. b. Institut de Chimie des Substances Naturelles du CNRS, F-91198 Gif-sur-Yvette, France. c. Departament de Cristallografia, Mineralogia i Dipòsits Minerals, Universitat de Barcelona, 08028 Barcelona, Spain

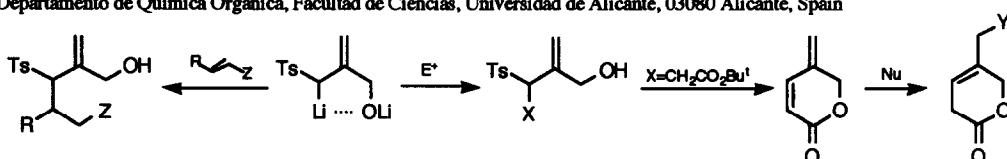


*Tetrahedron*, 1994, 50, 6603

**TOSYLATED LITHIUM 2-(LITHIOMETHYL)-2-PROPEN-1-Olate:  
A  $\gamma$ -ALKOXIDE ALLYL SULFONE ANION IN ORGANIC SYNTHESIS**

Diego A. Alonso, Carmen Nájera and José M. Sansano

Departamento de Química Orgánica, Facultad de Ciencias, Universidad de Alicante, 03080 Alicante, Spain

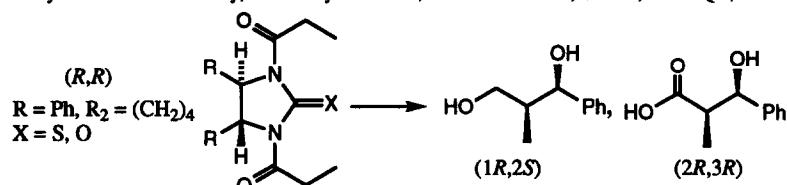


*Tetrahedron*, 1994, 50, 6621

**BIFUNCTIONAL CHIRAL AUXILIARIES 7: ALDOL REACTIONS OF ENOLATES DERIVED FROM 1,3-DIACYLIMIDAZOLIDINE-2-THIONES AND 1,3-DIACYLIMIDAZOLIDINE-2-ONES**

S.G. Davies\*, A.J. Edwards, G.B. Evans and A.A. Mortlock,

The Dyson Perrins Laboratory, University of Oxford, South Parks Road, Oxford, OX1 3QY, UK.

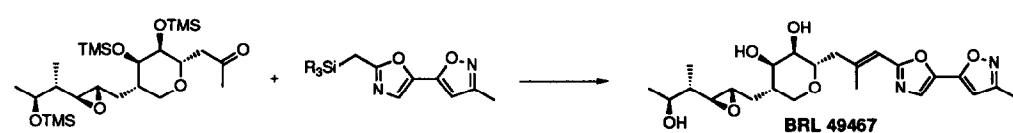


*Tetrahedron*, 1994, 50, 6643

**STEREOSELECTIVITY IN THE PETERSON REACTION -  
APPLICATION TO THE SYNTHESIS OF BRL 49467**

David Bell, Eleanor A. Crowe, Nicholas J. Dixon, Graham R. Geen\*, Inderjit S. Mann and Mark R. Shipton  
SmithKline Beecham Pharmaceuticals, Coldharbour Road, The Pinnacles, Harlow, Essex, CM19 5AD, U.K.

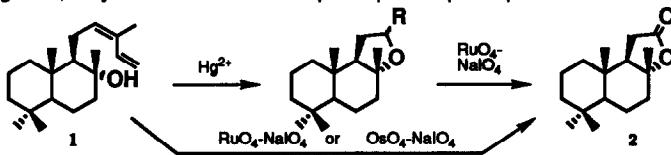
The *E*:*Z* selectivity in the introduction of the tri-substituted double bond in BRL 49467 could be controlled by the choice of conditions used in a Peterson reaction.



**SYNTHESIS OF *nor*-AMBREINOLIDE FROM (+)-*cis*-ABIENOL**  
 Alejandro F. Barrero,\* Juan F. Sánchez, Enrique J. Alvarez-Manzaneda, Joaquín Altarejos, Manuel Muñoz and Ali Haidour  
 Departamento de Química Orgánica, Facultad de Ciencias, Universidad de Granada, 18071 Granada (Spain)

Tetrahedron, 1994, 50, 6653

*nor*-Ambreinolide (2) has been prepared from *cis*-abienol (1) through induced cyclization with  $Hg(OAc)_2$  and oxidative treatment of the resulting crude, or by direct reaction with  $RuO_4-NaIO_4$  or  $OsO_4-NaIO_4$ .



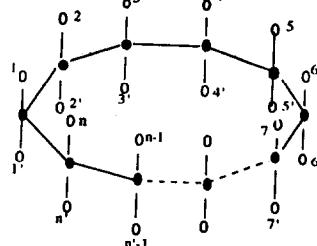
**ON THE ENUMERATION OF CHIRAL AND ACHIRAL SKELETONS OF POSITION ISOMERS OF HOMOSUBSTITUTED MONOCYCLIC CYCLOALKANES WITH A RING SIZE  $n$  (odd or even).**

Tetrahedron, 1994, 50, 6663

R. M. NEMBA\*, F. NGOUHOOU, Laboratory of Physical chemistry, Faculty of Science, University of Yaounde I, P.O. Box 812 Yaounde, Cameroun.

Stereograph of a monocyclic cycloalkane whose cycle index is taken as a basis for applying Polya's theorem to the enumeration of chiral and achiral skeletons of position isomers of  $C_nH_{2n-k}X_k$  systems.

Stereograph of a monocyclic cycloalkane  $(CH_2)_n$



**STEREOSELECTIVE CONTROL IN 1,3-DIPOLAR CYCLOADDITION OF NITRONES TO SUBSTITUTED STYRENES**

Tetrahedron, 1994, 50, 6671

Ugo Chiacchio,<sup>a</sup> Franco Casuscelli,<sup>a</sup> Antonino Corsaro,<sup>b</sup> Antonio Rescifina,<sup>b</sup> Giovanni Romeo,<sup>a,\*</sup> and Nicola Uccella<sup>c</sup>

<sup>a</sup>Dipartimento Farmaco-chimico, Università, 98168 Messina, Italy

<sup>c</sup>Dipartimento di Scienze Chimiche, Università, 95125 Catania, Italy

<sup>c</sup>Dipartimento di Chimica, Università, 87036 Arcavacata di Rende, Italy

The stereochemistry of 1,3-dipolar cycloaddition of C-methyl-*N*-phenylnitrone with substituted styrenes has been investigated. The presence of an hydroxyl function at the ortho position in the dipolarophile completely controls the stereochemical course of the reaction with the exclusive formation of *cis* cycloadduct.

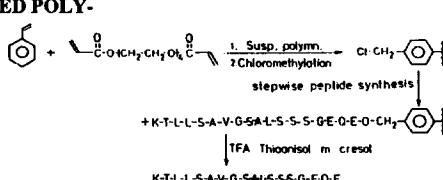


**GEL-PHASE PEPTIDE SYNTHESIS ON A NEW HIGH-CAPACITY TETRAETHYLENEGLYCOL DIACRYLATE-CROSSLINKED POLYSTYRENE SUPPORT: SYNTHESIS OF PARDAXIN 16-33**

Tetrahedron, 1994, 50, 6681

M. Renil, R. Nagaraj, V.N. Rajasekharan Pillai\*

School of Chemical Science,  
 Mahatma Gandhi University,  
 Kottayam 686 560, India.

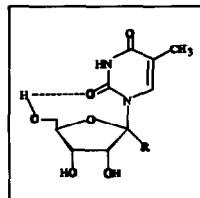


INTRAMOLECULAR HYDROGEN BONDING IN PRIMARY HYDROXYL OF THYMINE 1-(1-DEOXY- $\beta$ -D-PSICOFURANOSYL) NUCLEOSIDE

Tetrahedron, 1994, 50, 6689

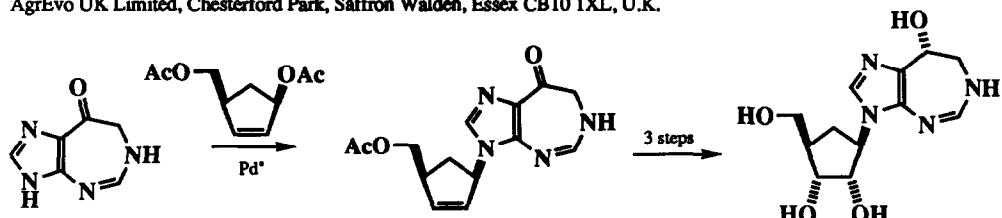
X. Martin\*, M. Moreno\*, J. M. Lluch\* and A. Grouiller\*. a)Departament de Química, Universitat Autònoma de Barcelona, 08193 Bellaterra (Barcelona), Spain. b) Laboratoire de Chimie Organique II, Université Lyon I, ESCIL, 43, Boulevard du 11 Novembre 1918, 69622 Villeurbanne, France.

By using the semiempirical AM1 method, it is concluded that a clear intramolecular hydrogen bond exists in 1-(1-Deoxy- $\beta$ -D-Psicofuranosyl) Thymine.



TOTAL SYNTHESIS OF CARBOCYCLIC ANALOGUES OF COFORMYCIN

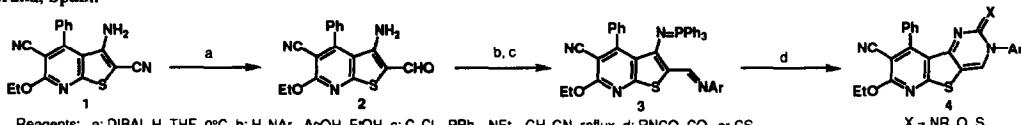
E. A. Saville-Stones, R. M. Turner, S. D. Lindell,\* N. S. Jennings, J. C. Head and D. S. Carver, AgrEvo UK Limited, Chesterford Park, Saffron Walden, Essex CB10 1XL, U.K.



AN EFFICIENT IMINOPHOSPHORANE-MEDIATED SYNTHESIS FOR PYRIDO[3',2':4,5]THIENO[3,2-d]PYRIMIDINE DERIVATIVES

Tetrahedron, 1994, 50, 6705

Carlos Peinador, María J. Moreira and José M. Quintela\*  
Departamento de Química Fundamental e Industrial, Facultad de Ciencias, Universidad de La Coruña, Campus de A Zapateira, E-15071, La Coruña, Spain.



Reagents: a: DIBAL-H, THF, 0°C; b: H2NAr, AcOH, EtOH; c: C2Cl6, PPh3, NEt3, CH3CN, reflux; d: RNCO, CO2 or CS2

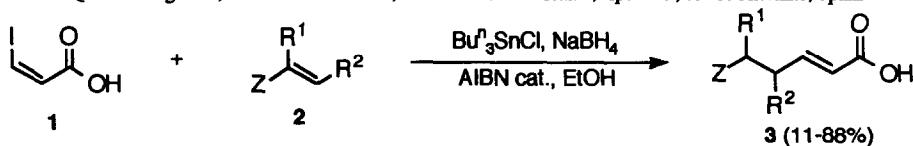
X = NR, O, S

Pyridothienopyrimidines bearing various substituents at position 2 of the pyrimidine ring are reported. The aza Wittig-type reaction of iminophosphoranes 3 with isocyanates, carbon dioxide and carbon disulfide leads to 2,3-dihydropyrido[3',2':4,5]thieno[3,2-d]pyrimidines 4.

$\beta$ -ACYLVINYL INTERMOLECULAR RADICAL ADDITIONS TO DOUBLE BONDS: STEREOSELECTIVE SYNTHESIS OF FUNCTIONALISED (*E*)- $\alpha$ , $\beta$ -UNSATURATED CARBOXYLIC ACIDS

Tetrahedron, 1994, 50, 6715

F. Foubelo, F. Lloret and M. Yus\*  
Departamento de Química Orgánica, Facultad de Ciencias, Universidad de Alicante, Apdo. 99, 03080 Alicante, Spain



(R<sup>1</sup>= or ≠R<sup>2</sup>=H, Me, Cl; Z=CO<sub>2</sub>Me, CN, CONMe<sub>2</sub>, Cl, CO<sub>2</sub>CH<sub>2</sub>CH=CH<sub>2</sub>)