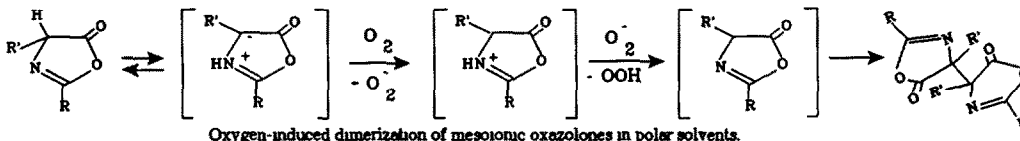


OXIDATION OF MESOIONIC OXAZOLONES BY OXYGEN

H. Rodriguez* (1), A. Marquez (1), C.A. Chuaqui* (2) and B. Gomez (1)

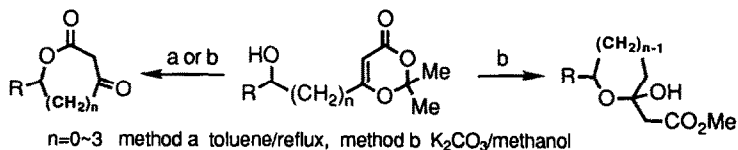
(1) Departamento de Quimica Organica, Facultad de Ciencias Quimicas y Farmaceuticas, Universidad de Chile, Casilla 233, Santiago, Chile; (2) Radiation Applications Research Branch, Whiteshell Laboratories, AECL Research, Pinawa, Manitoba, ROE 1L0, Canada



TWO LACTONE FORMATION REACTIONS FROM 1,3-DIOXIN-4-ONES HAVING HYDROXYALKYL GROUP AT THE 6-POSITION: DIFFERENCE IN RING OPENING AND CLOSURE

Masayuki Sato*, Jun-ichi Sakaki, Yoshiaki Sugita, Sanae Yasuda, Hiroko Sakoda, and Chikara Kaneko*
Pharmaceutical Institute, Tohoku University, Aobayama, Sendai 980, Japan

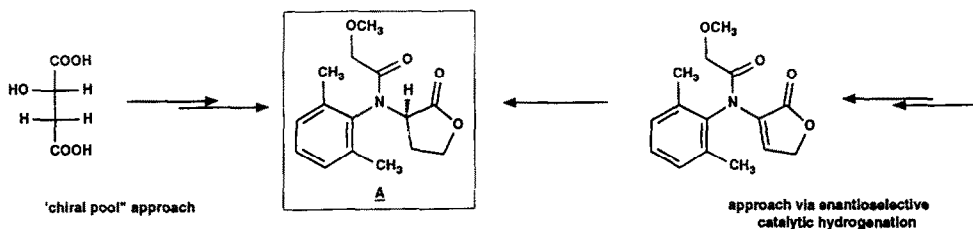
Scope and mechanism of methods a and b are clarified



TWO ENANTIOSELECTIVE SYNTHESSES OF A PRECURSOR OF THE BIOLOGICALLY MOST ACTIVE ISOMER OF CGA 80000 (CLOZYLACON)

H P Buser*, B Pugin, F Spindler, M Sutter
Agricultural Division and Central Research Laboratories, CIBA-GEIGY Ltd, CH-4002 Basel, Switzerland

The aminobutyrolactone A was synthesized enantioselectively via the following two conceptually different routes

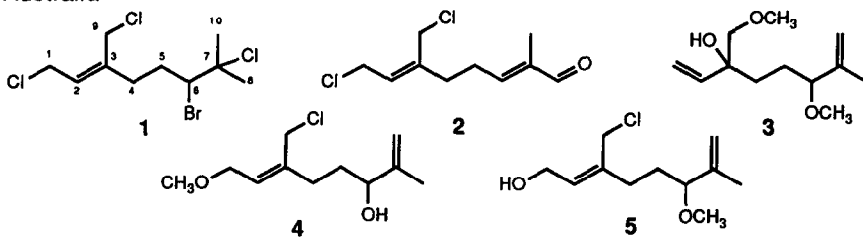


FIVE NEW MONOTERPENES FROM THE MARINE RED ALGA *PORTIERIA HORNEMANNII*

Anthony D Wright¹, Gabriele M König¹, Otto Sticher¹ and Rocky de Nys²

¹Department of Pharmacy, Swiss Federal Institute of Technology(ETH) Zurich,

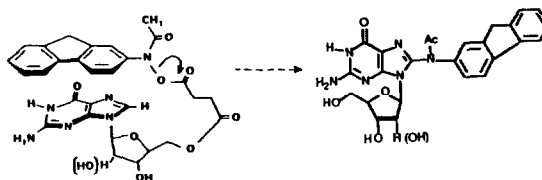
²Department of Chemistry and Biochemistry, James Cook University, Townsville, Australia



ARYLAMIDATION DE LA GUANOSINE PAR UN AGENT CANCEROGENE, LE 2-AMINOFLOURENE. APPROCHE INTRAMOLECULAIRE.

E DEFRANCO, A LETERME, N PELLOUX, M F LHOMME et J LHOMME*

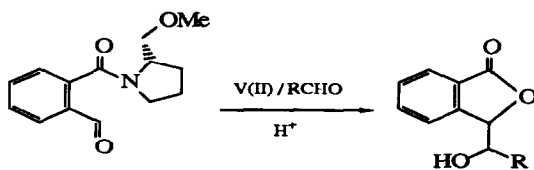
Laboratoire d'Etudes Dynamiques et Structurales de la Sélectivité URA 332, Université Joseph Fourier, BP 53 X, 38041 GRENOBLE CEDEX - FRANCE



DIASTEREO AND ENANTIOSELECTIVE SYNTHESIS OF 1,2-DIOLS BY VANADIUM (II) PROMOTED PINACOL CROSS COUPLING

R Annunziata, M Cinquini, F Cozzi, P Giaroni, and M Benaglia
Università di Milano (Italy) - Dipartimento di Chimica Organica e Industriale

The V(II) promoted pinacol coupling of chiral aromatic aldehyde **1** with aliphatic and alkoxy aldehydes afford syn-diols with good diastereo- and enantioselection

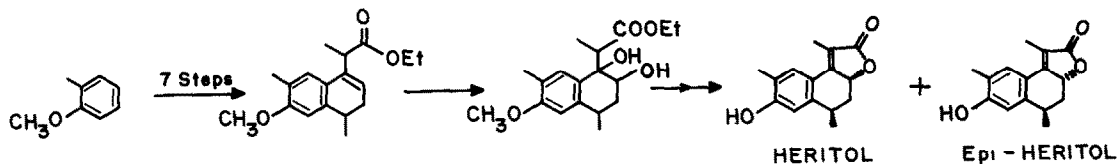


R = Pr-n, PhCH₂, Pr-i, C₆H₁₁-c, Bu-t, PhCH₂OCH₂, (S)- and (R)-CH(OR')R''

SYNTHESIS OF (±)HERITOL

P.K. Zubaidha, Subhash P. Chavan, Uday S. Racherla and Nagaraj R. Ayyangar*
National Chemical Laboratory, Pune 411 008, INDIA.

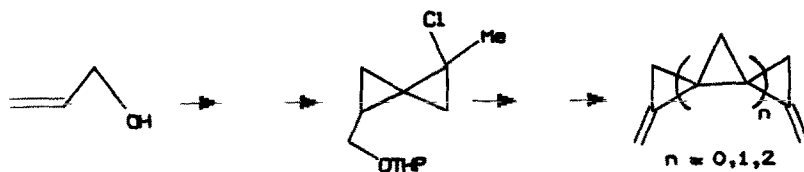
Synthesis of (±)Heritol and (±) epi-Heritol has been achieved by osmylation of the unsaturated ester as the key step to construct the butenolide moiety of the title compound.



THE SYNTHESIS OF BISMETHYLENE DERIVATIVES OF TRIANGULANES

Kirill A. Lukin, Anna Yu. Masunova, Bogdan I. Ugrak,
Nikolai S. Zefirov*

Department of Chemistry Moscow State University, Moscow,
119899, USSR

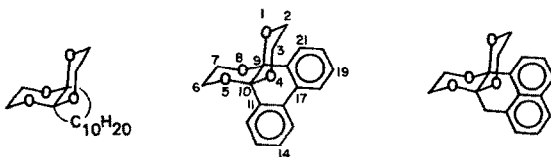


STRUCTURE, CONFORMATION AND STEREOELECTRONICS OF 9,10-ANNELATED-1,4,5,8-TETRAOXADICALINS A MULTIDISCIPLINARY PROBE

Pinchas Aped^a, Benzion Fuchs^a, Israel Goldberg^{a*}, Hanoch Senderowitz^a, Leah Schleifer^a, Evgeny Tartakovsky^a, Marc Anteunis^b and Frans A. M. Borremans^b

^aSchool of Chemistry, Tel-Aviv University, Ramat-Aviv, 69978 Tel-Aviv, Israel

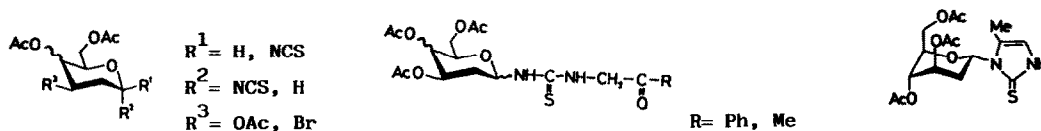
^bDepartment of Organic Chemistry, State University of Gent, B-9000 Gent, Belgium



PREPARATION AND PROPERTIES OF 2-DEOXYGLYCOSYL ISOTHIOCYANATES

J. Fuentes, M. A. Pradera and I. Robina, Departamento de Química Orgánica, Facultad de Química, Universidad de Sevilla, Sevilla, Spain

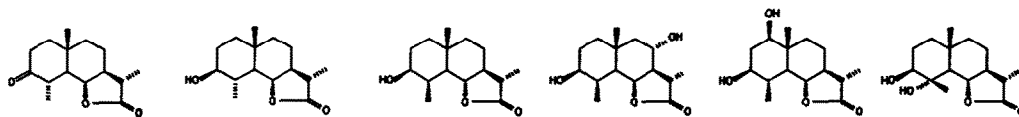
The syntheses and spectroscopic data of 2-deoxy- α and β -D-arabino-(D-lyxo)hexopyranosyl isothiocyanates, 2-deoxy hexopyranosylthiureas, and 2-deoxyhexopyranosyl-4-imidazoline-2-thione are reported.



BIOTRANSFORMATION OF 6 β -Eudesmanolides FUNCTIONALIZED AT C-3 WITH *CURVULARIA LUNATA* AND *RHIZOPUS NIGRICANS* CULTURES

Y Amate^a, J.L. Bretón^b, A. García-Granados^{a*}, A. Martínez^a, M.E. Onorato^a, A. Sáenz de Buruaga^a and J.M. Aras^c; ^aDepartamento de Química Orgánica, ^bDepartamento de Microbiología, Facultad de Ciencias, Universidad de Granada, 18071-Granada, ^cInstituto de Productos Naturales Orgánicos (C.S.I.C.), La Laguna (Tenerife), Spain.

Biotransformation of substrates by *Rhizopus nigricans* gave these metabolites in acceptable yield. Epimerization at C-4 was produced.



SUBSTRATES

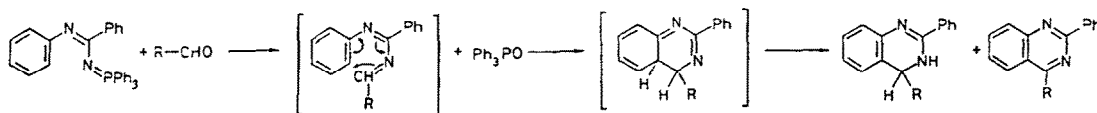
METABOLITES

A CONVENIENT SYNTHESIS OF QUINAZOLINE RING BY TANDEM AZA-WITTIG REACTION/ELECTROCYCLIC RING CLOSURE

Elisabetta Rossi^{*}, Daniela Calabrese and Francesca Farma

Istituto di Chimica Organica, Facoltà di Farmacia, Viale Abruzzi, 42 - 20131 - Milano - Italy

The reaction mechanism was proven by an independent and original synthesis of the 1,3-diazabuta-1,3-diene intermediate



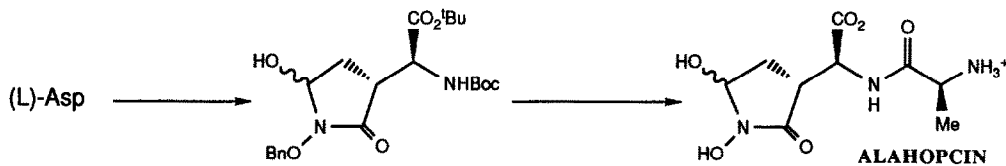
THE STEREOSPECIFIC SYNTHESIS OF ALAHOPCIN

Jack E Baldwin, Robert M Adlington, Christopher R A Godfrey,^a

David W Gollins, and Christopher J Schofield

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

^aICI Agrochemicals, Jealott's Hill Research Station, Bracknell, Berkshire, RG12 6EY



We report the first total synthesis of the dipeptide antibiotic alahopcin, from (L)-aspartic acid, via a previously reported intermediate

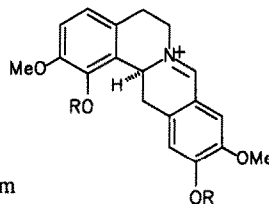
A NOVEL OXIDATION STAGE IN THE CHEMISTRY OF PROTOBERBERINE ALKALOIDS. SYNTHESIS OF 7,8-DEHYDROBERBINES.

Rafael Suau^{*}, M Victoria Silva and María Valpuesta

Departamento de Química Orgánica

Universidad de Málaga, Málaga, SPAIN

Cis B/C fused berbines are partially oxidized by iodine allowing the preparation of the 7,8-dehydroberbines derived from (-)-Caseamine, (-)-Caseadine and (-)-O-Methylcaseadine



STRUCTURAL STUDIES ON 1-(1-DEOXY- β -D-PSICOFURANOSYL)

THYMINE

J Plavec¹, V Buet², A Grouiller², L. Koole¹ & J Chattopadhyaya^{1*}

¹Department of Bioorganic Chemistry, Box 581, Biomedical Center, University of Uppsala, S-751 23 Uppsala, Sweden

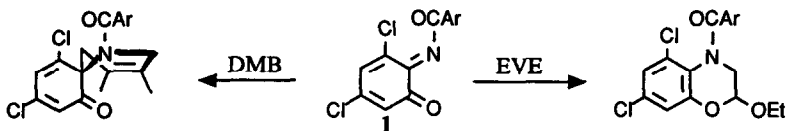
²Laboratoire de Chimie Organique II, Université Lyon I, ESCIL, 43, Boulevard du 11 Novembre 1918, 69622 Villeurbanne, France

Conformational analysis of a novel nucleoside analogue, 1-(1-Deoxy- β -D-psicofuranosyl)thymine (**3**), is described basing (i) on the vicinal proton-proton J-coupling constants by 500 MHz ¹H-NMR spectroscopy at different sample temperatures, (ii) the data analysis by pseudorotation concept, and (iii) finally an explicit model of the preferred conformation of **3** in solution has been defined through a set of molecular mechanics calculation using Allinger's MM2 force field and molecular modelling.

SOLVENT EFFECT AS THE RESULT OF FRONTIER MOLECULAR ORBITAL INTERACTION. VI. THE DIELS ALDER REACTIONS OF AN N-ACYL-o-QUINONE MONOIMIDE BEHAVING AS DIENE OR DIENOPHILE.

G Desimoni, G Fatta, and P P Righetti.

Dipartimento di Chimica Organica dell'Università, V le Taramelli 10, 27100 Pavia, Italy

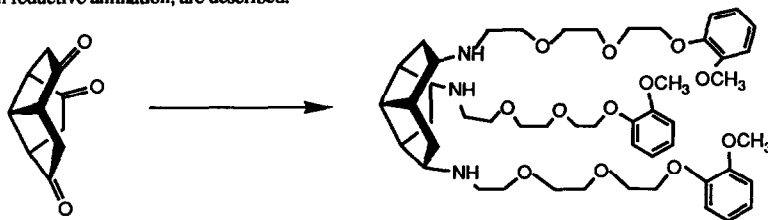


o-Quinone monoimide **1** behaves as heterodienophile or heterodiene in the Diels-Alder reactions with dimethylbutadiene (DMB) and ethyl vinyl ether (EVE). The solvent effect of these two reactions was investigated.

PERHYDROTRIQUINACENIC HOSTS. 1. SYNTHESIS, COMPLEXATION AND TRANSPORT PROPERTIES OF TRIPODANDS OF C₃ SYMMETRY.

Carmen Almansa, Albert Moyano and Félix Serratosa. Departament de Química Orgànica, Universitat de Barcelona, Facultat de Química, C Martí i Franquès, 1-11, E-08028 Barcelona (Spain)

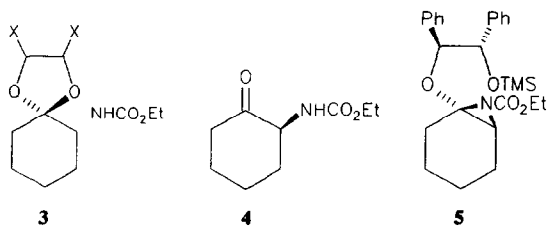
The synthesis, complexation and transport properties of tripodands of C₃ symmetry, derived from tricyclo[5.2.1.0^{4,10}]decane-2,5,8-trione through reductive amination, are described.



ASYMMETRIC FORMATION OF C-N BONDS IN CHIRAL ENOL ETHERS

Stefania Fioravanti, M Antonietta Loreto, Lucio Pellacani, Paolo A. Tardella

Dipartimento di Chimica, Università "La Sapienza", Roma, Italy



3 and/or **4** are the main products of the diastereoselective addition of EtO₂CN (generated at 120 °C or at room temperature from different precursors) to chiral enol ethers

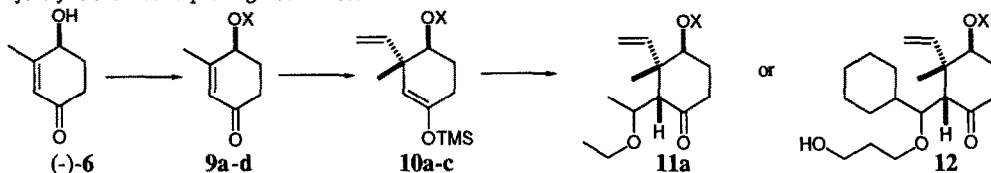
In one case the intermediate aziridine **5** is obtained in high diastereomeric purity

**SYNTHESIS OF OPTICALLY ACTIVE CYCLOHEXENOL DERIVATIVES
VIA ENZYME CATALYZED ESTER HYDROLYSIS OF 4-ACETOXY-3-METHYL-2-CYCLOHEXENON.**

Magnus Polla and Torbjorn Frejd*

Org Chem 2, Chemical Center, The Lund Institute of Technology, P O Box 124, S-221 00 Lund, Sweden

The optically active cyclohexenol derivatives **9a-d**, **10a-c**, **11a** and **12** are synthesized from (-)-**6**, which is obtained by PLE catalyzed hydrolysis of the corresponding racemic acetate.



**Diels-Alder Reactions of Protoanemonin with
Heterosubstituted Dienes. Synthesis of
Polyfunctional Oxaspiro 4.5 decanes.**

D. Alonso,^a J. Font,^a R. M. Ortuño,^a J. d'Angelo,^b

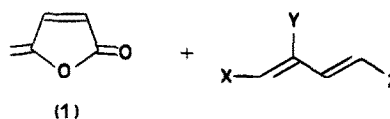
A. Guingant,^b C. Bois.^c

a: Universitat Autònoma de Barcelona, Bellaterra (Barcelona), Spain;

b: ESPCI, Unité Associée au CNRS, N°476, Paris, France;

c: Université Pierre et Marie Curie, Paris, France.

Cycloadditions between protoanemonin (**1**) and several heterosubstituted dienes have been investigated; (**1**) has shown to be a good dienophile towards electron-rich dienes giving adducts that can be easily transformed into useful synthetic building blocks.



INTRAMOLECULAR S_N2' MACROCYCLISATIONS

Jack E Baldwin, Robert M. Adlington, Mark B. Mitchell and Jeremy Robertson

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

The synthesis of 10-15 membered α -methylene macrocyclic lactones from functionalised allylstannanes is described. Attempts to synthesise analogous 6-9 membered lactones proved unsuccessful, resulting instead in the production of dilactones and AIBN derived adducts.

