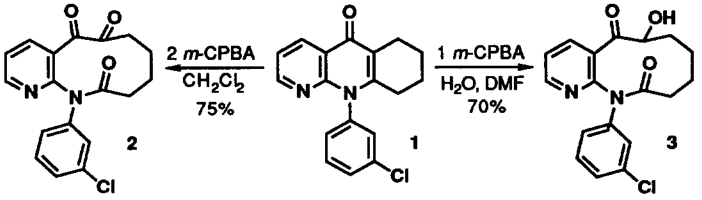
Tetrahedron, **1991**, *47*, 9981

Oxidative Cleavage of a Tricyclic Pyridone to a Bicyclic Lactam-dione

Richard J. Friary* and John H. Schwerdt

Schering-Plough Research, Schering-Plough Corp.
60 Orange Street, Bloomfield, NJ 07003, U. S. A.

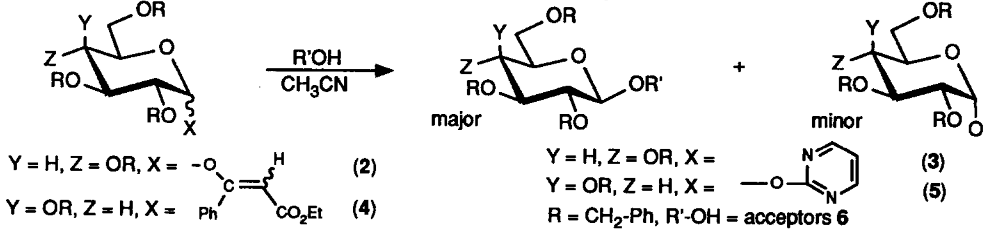
Two equivalents of *m*-chloroperbenzoic acid in dry dichloromethane cleaves the puridone ring of 1, forming lactam α-diketone 2. With one equivalent of peracid in aqueous dimethylformamide, the product is lactam α-ketol 3.



Tetrahedron, **1991**, *47*, 9985

SYNTHESIS OF β-O-GLYCOSIDES USING ENOL ETHER AND IMIDATE DERIVED LEAVING GROUPS. EMPHASIS ON THE USE OF NITRILE AS A SOLVENT

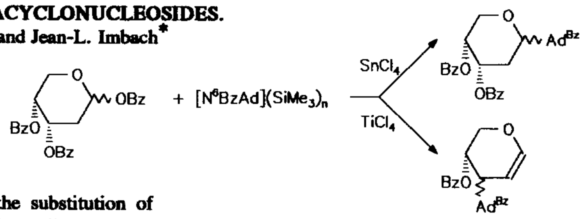
Y.D. Vankar, P.S. Vankar, M. Behrendt and R.R. Schmidt*
Fakultaet Chemie, Universitaet Konstanz, D-7750 Konstanz, Germany



Tetrahedron, **1991**, *47*, 9993

SnCl$_4$ VERSUS TiCl$_4$-MEDIATED COUPLING BETWEEN N-6-BENZOYLADENINE AND PERBENZOYLATED 2-DEOXYRIBOPYRANOSE:APPLICATION TO THE SYNTHESIS OF CERTAIN HOMOCHIRAL ACYCLO AND CARBOACYCLONUCLEOSIDES.

Marie-V. Baud, Claude Chavis, Marc Lucas and Jean-L. Imbach*
Université de Montpellier II, Sciences
et Techniques du Languedoc,
Laboratoire de Chimie Bio-Organique,
case 008, 34095 Montpellier Cedex 5
(France).

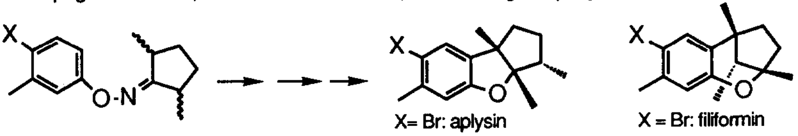


TiCl$_4$ and SnCl$_4$ mediate regioselectively the substitution of N-6-benzoyladenine on perbenzoylated 2-deoxyribopyranose. 3'-Adenylglycals are obtained with TiCl$_4$ while nucleosides are formed with SnCl$_4$.

Tetrahedron, **1991**, *47*, 10003

The Rearrangement of some Cyclopentanone-Aryloximes : Synthesis of (±)-Aplysin, (±)-Filiformin and of their Debromo Analogues.

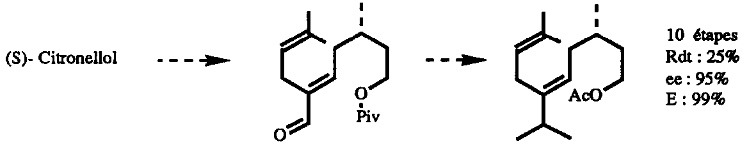
J.Yves Laronze*, Rachida El Boukili, Dominique Patigny,Seloua Dridi, Dominique Cartier and Jean Lévy.
Laboratoire de Transformations et Synthèse de Substances Naturelles, associé au CNRS, Université de Reims-Champagne-Ardenne, Faculté de Pharmacie, 51, rue Cognacq-Jay, 51096 Reims Cédex, France.



Tetrahedron, **1991**, *47*, 10015

SYNTHESE STÉRÉOSÉLECTIVE DE LA PHÉROMONE SEXUELLE D'AONIDIELLA CITRINA : LE (3S,5E)-(–)-3,9-DIMÉTHYLE-6-ISOPROPYLE DÉCA-5,8-DIENE-1-YLE ACÉTATE

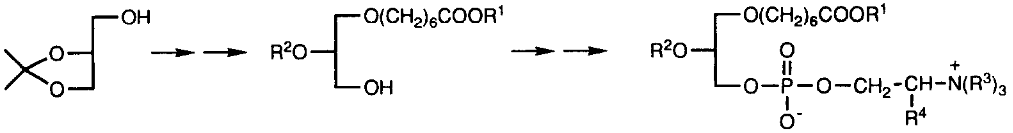
René Baudouy et Marie-Rose Sancho
Laboratoire de Synthèse Organique Appliquée et Laboratoire de Chimie Organique I - Université Claude Bernard Lyon I - ESCIL - 43, Bd du 11 Novembre 1918 - 69622 Villeurbanne Cédex - France



Tetrahedron, **1991**, *47*, 10023

SYNTHESIS OF NEW ETHER GLYCEROPHOSPHOLIPIDS STRUCTURALLY RELATED TO MODULATOR

M. L. García*, J. Pascual, L. Borràs, J. A. Andreu, E. Fos, D. Mauleón, G. Carganico and F. Arcamone[#]
Laboratorios Menarini S.A., Alfonso XII, 587, Badalona, Spain. [#]Menarini Ricerche Sud, Tito Speri, 10, Pomezia, Italy.

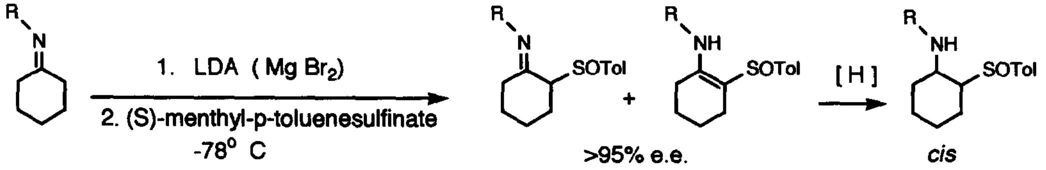


Several ether glycerophospholipids with phosphoserine or phosphocholine moiety in position *sn*-3 were synthesized in good overall yields.

Tetrahedron, **1991**, *47*, 10035

Synthesis and Hydride Reductions of Chiral Cyclic β-Iminosulfoxides.

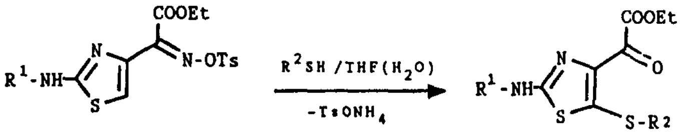
M. Carmen Carreño, Esteban Domínguez, José L. García Ruano,*
Concepción Pedregal and Jesús H. Rodríguez.
Dpto. de Química, Facultad de Ciencias, Universidad Autónoma. Cantoblanco. 28049-Madrid, SPAIN.



Tetrahedron, **1991**, *47*, 10045

Reaction of Aminothiazole-oximetosylates with Thiols

Béla Rezessy,* Lajos Toldy, Gyula Horváth, Pál Sohár[a]
Institute for Drug Research, H-1325 Budapest, POB 82, Hungary
[a] EGIS Pharmaceuticals, Spectroscopic Department, H-1475 Budapest POB 100, Hungary



R^1: H-, Tr-

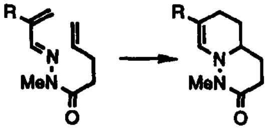
R^2: -Et, -benzyl, -CH_2COOMe

Tetrahedron, **1991**, *47*, 10053

INTRAMOLECULAR AND INTERMOLECULAR DIELS-ALDER REACTIONS OF ACYLHYDRAZONES DERIVED FROM METHACROLEIN AND ETHYLACROLEIN

Sylvia J. Allcock, Thomas L. Gilchrist and Stephen J. Shuttleworth, Chemistry Department, University of Liverpool, Liverpool L69 3BX
Frank D. King, SmithKline Beecham Pharmaceuticals, Medicinal Research Centre, The Pinnacles, Harlow CM19 5AD, U.K.

Intramolecular Diels-Alder reactions of the type illustrated have been achieved using terminally unsaturated acylhydrazones of methacrolein and ethylacrolein.



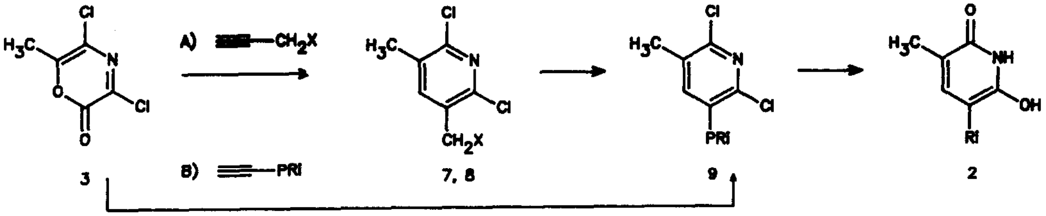
These acylhydrazones, and also some others, undergo intermolecular cycloaddition to *N*-phenylmaleimide.

Tetrahedron, **1991**, *47*, 10065

An Approach to Acyclo-1-Deazathymidine C-Nucleosides via 3,5-Dichloro-6-Methyl-2*H*-1,4-Oxazin-2-one

Lieven Meerpoel, Suzanne M. Toppet, Frans Compernolle and Georges J. Hoornaert*. Laboratorium voor Organische Synthese K.U.Leuven, Celestijnenlaan 200-F, B-3001 Leuven, Belgium

Diels-Alder reaction of oxazinone 3 with A) HC≡CCH$_2$X (X=Br, OTHP) or B) HC≡CPRi affords precursors 7 and 8 or 9, which are elaborated to the title compounds 2 (PRi and Ri = protected and deprotected acyclo sugar units).

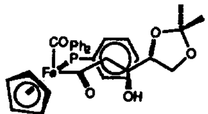


Tetrahedron, **1991**, *47*, 10077

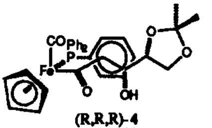
A Study of the Aldol Reaction between Enolates Derived from the Iron Acetyl Complex [(η^5-C$_5$H$_5$)Fe(CO)(PPh$_3$)-COCH$_3$] and 2,3-O-Isopropylidene-D-glyceraldehyde

Graham J. Bodwell, Stephen G. Davies* and Andrew A. Mortlock
The Dyson Perrins Laboratory, South Parks Road, Oxford, OX1 3QY, U.K.

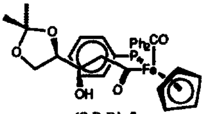
All four diastereomers of the β-hydroxy acyl complex resulting from the aldol reaction between enolates derived from the iron acetyl complex (η^5-C$_5$H$_5$)Fe(CO)(PPh$_3$)COCH$_3$ and the homochiral aldehyde 2,3-O-isopropylidene-D-glyceraldehyde have been prepared in high diastereisoomeric excess by varying the enantiomer of the chiral auxiliary and the counterion of the enolate.



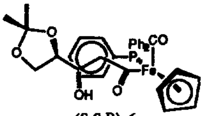
(R,S,R)-3
d.e. = >99%



(R,R,R)-4
d.e. = 90%



(S,R,R)-5
d.e. = 94%



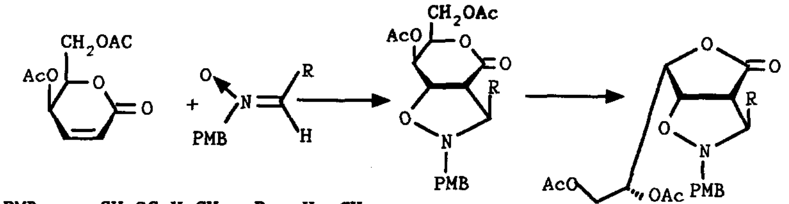
(S,S,R)-6
d.e. = 92%

Tetrahedron, **1991**, *47*, 10087

1,3-DIPOLAR CYCLOADDITION OF NITRONES TO SUGAR ENLACTONES

Irma Panfil, Czesław Bełżecki, Zofia Urbańczyk-Lipkowska, and Marek Chmielewski*
Institute of Organic Chemistry, Polish Academy of Sciences, 01-224 Warsaw, Poland

1,3-Dipolar cycloaddition of nitrones to sugar enlactones is described.



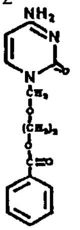
PMB - p-CH$_3$OC$_6$H$_4$CH$_2$; R - H, CH$_3$

Tetrahedron, **1991**, *47*, 10095

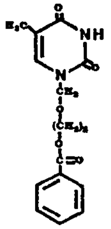
Selective Intracomplex Reaction of Pendant Groups in a Purine Pyrimidines Base Pair.

Salim Hadad, Bernard S. Green, Jehoshua Katzhendler*
Department of Pharmaceutical Chemistry, School of Pharmacy,
The Hebrew University of Jerusalem 91120, ISRAEL.

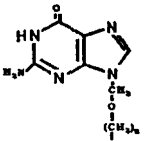
Association between guanine (G-NH_2) derivatives and cytosine benzoate ester (C-Benz) exhibit enhanced intracomplex reactions. The reaction rates depends on the chain length of G-NH_2. The best spatial orientation of G-NH_2 is achieved when n=4.



C-Benz



T-Benz



GNH$_2$(2) n=2
GNH$_2$(4) n=4
GNH$_2$(6) n=6

Tetrahedron, **1991**, *47*, 10101

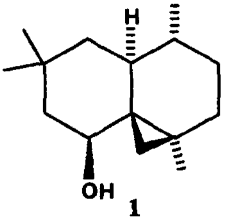
THE STRUCTURE OF LAUROBTUSOL, A NEW REARRANGED SESQUITERPENOID FROM THE MEDITERRANEAN RED ALGA *LAURENCIA OBTUSA*

Salvatore Caccamese[a], Vincenzo Amico[a], Placido Neri[b] and Mario Foti[b]

[a]Dipartimento di Scienze Chimiche, Università Catania, Italy
[b]Istituto C.N.R. Sostanze Naturali, Catania, Italy

The new tricyclic humulane derivative laurobtusol (1) was isolated from the red alga *Laurencia obtusa*.

The structure was established mainly by 2D NMR methods and the relative configuration was assigned by a quantitative computer simulation of the lanthanide induced shifts in the ^1H NMR spectrum and Molecular Mechanics Calculation (MM2).



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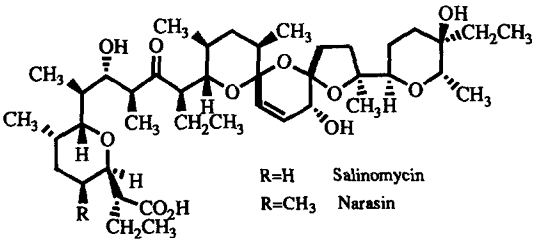
Tetrahedron, **1991**, *47*, 10109

THE COMPLETE ASSIGNMENT OF THE ^1H AND ^{13}C NMR SPECTRA OF THE ALKALI METAL SALTS OF SALINOMYCIN AND NARASIN

Frank G Riddell and Stephen J. Tompsett, Department of Chemistry, The University of St Andrews, St Andrews KY16 9ST, Scotland.

The preparation of and the assignments of the ^1H and ^{13}C nmr spectra of the alkali metal salts of salinomycin and narasin is described.



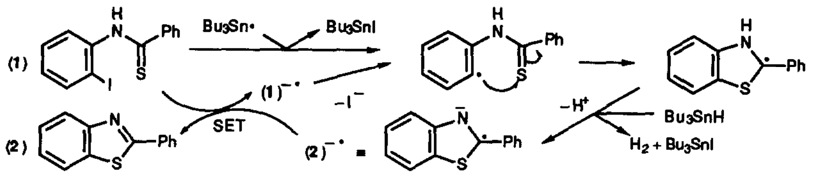
R=H Salinomycin
R=CH_3 Narasin

Tetrahedron, **1991**, *47*, 10119

OXIDATION DURING REDUCTIVE CYCLISATIONS USING Bu_3SnH

W. Russell Bowman*, Harry Heaney*, and Benjamin M. Jordan
Department of Chemistry, University of Technology, Loughborough, Leics. LE11 3TU, Great Britain

Reductive cyclisations using Bu_3SnH include an "oxidation" step if the removal of an acidic proton from the intermediate cyclised radical, by Bu_3SnH acting as a base, is favourable. A "pseudo $S_{RN}1$ mechanism is proposed, e.g



Tetrahedron, **1991**, *47*, 10129

SOME CHEMICAL TRANSFORMATIONS OF THE *NEO*-CLERODANE DITERPENE TEUBOTRIN

P. Y. Malakov, M. C. de la Torre, B. Rodríguez* and G. Y. Papanov; Instituto de Química Orgánica, CSIC, Madrid, Spain, and Department of Organic Chemistry, University of Plovdiv, Bulgaria

The title compound (1) was transformed, among other substances, into the derivatives 5-7, 9 and teuscordinon (12), a naturally occurring diterpenoid.