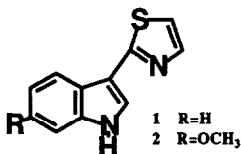


GRAPHICAL ABSTRACTS

Tetrahedron, 1991, 47, 3909

THE CAMALEXINS: NEW PHYTOALEXINS PRODUCED IN THE LEAVES OF *CAMELINA SATIVA* (CRUCIFERAE).

Lois M. Browne^{1*}, Kenneth L. Conn², William A. Ayer¹ and Jalpa P. Tewari²,
Departments of Chemistry¹ and Plant Science², University of Alberta, Edmonton, Alberta, Canada T6G 2E1



Two new thiazoyl substituted indole phytoalexins, camalexin (1) and methoxycamalexin (2), were isolated from *Camelina sativa* leaves following elicitation by the fungus *Alternaria brassicae*.

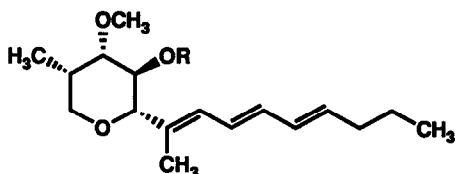
Tetrahedron, 1991, 47, 3915

STRUCTURE ELUCIDATION OF RESTRICTICIN, A NOVEL ANTIFUNGAL AGENT FROM *PENICILLIUM RESTRICTUM*

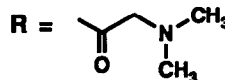
Otto D. Hensens*, Carol F. Wichmann, Jerrold M. Liesch, Frank L. VanMiddlesworth, Kenneth E. Wilson and Robert E. Schwartz

Department of Natural Products Chemistry, Merck Sharp & Dohme Research Laboratories, P. O. Box 2000, Rahway, N. J. 07065-0900, U. S. A.

The structures and absolute stereochemistry of a new class of antifungal agents were determined on the basis of spectroscopic evidence.



R = H



Tetrahedron, 1991, 47, 3925

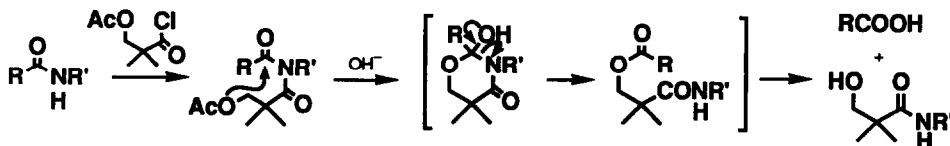
AN EFFICIENT METHOD FOR HYDROLYSIS OF *N*-MONOSUBSTITUTED AMIDES UTILIZATION OF INTRAMOLECULAR *N*-O ACYL MIGRATION IN HYDROXYPIVALIMIDES

Tetsuto Tsunoda*, Osamu Sasaki, Osamu Takeuchi, and Shô Itô

Department of Chemistry, Tohoku University, Sendai 980, and

Faculty of Pharmaceutical Sciences, Tokushima Bunri University Tokushima, 770 Japan

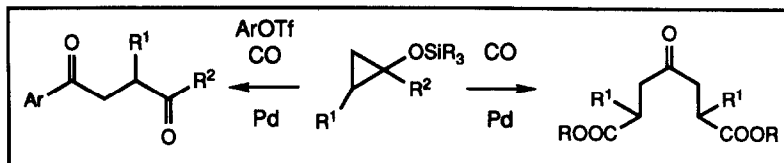
An efficient and versatile method for the hydrolysis of *N*-monosubstituted carboxamides has been developed.



SYNTHESIS OF 1,4-DICARBONYL COMPOUNDS AND 4-KETO PIMELATES BY PALLADIUM-CATALYZED CARBONYLATION OF SILOXYCYCLOPROPANES

Satoshi Aoki and
Eiichi Nakamura*

Department of Chemistry,
Tokyo Institute of Technology
Meguro, Tokyo 152 Japan

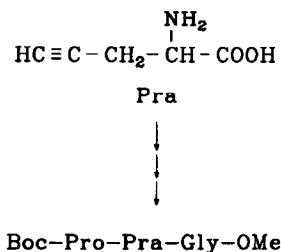


SYNTHESIS, PROPERTIES AND CRYSTAL STRUCTURE OF THE TRIPEPTIDE BOC-L-PROYL-L-PROPARGYL-GLYCYL-GLYCINE METHYL ESTER

Hans Willisch, Wolfgang Hiller, Bahram Hemmasi, Ernst Bayer.*
Institute of Organic Chemistry, University of Tübingen, Auf der
Morgenstelle 18, D-7400 Tübingen, F.R. Germany

Propargylglycine (Pra), as a powerful inhibitor of microbial growth, was built into a protected tripeptide with the sequence Pro-Pra-Gly. The peptide was employed to study its effects on the activity of prolyl 4-hydroxylase and the collagen biosynthesis. The Boc-protected tripeptide methyl ester was identified by mass spectrometry and NMR spectroscopy and its crystal structure was established by X-ray diffraction analysis.

Tetrahedron, 1991, 47, 3947

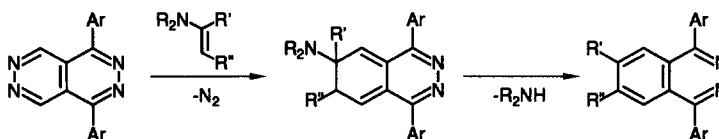


INVERSE-ELECTRON-DEMAND DIELS-ALDER REACTIONS OF CONDENSED PYRIDAZINES, PART 1. SYNTHESIS OF PHTHALAZINE DERIVATIVES FROM PYRIDAZINO[4,5-d]PYRIDAZINES.

Norbert Haider
Institute of Pharmaceutical Chemistry, University of Vienna, Währinger Straße 10, A-1090 Vienna, Austria

1,4-Diarylpyridazino[4,5-d]pyridazines were found to undergo [4+2] cycloaddition reactions with a variety of electron-rich dienophiles like enamines and ketene acetals to afford phthalazine derivatives.

Tetrahedron, 1991, 47, 3959

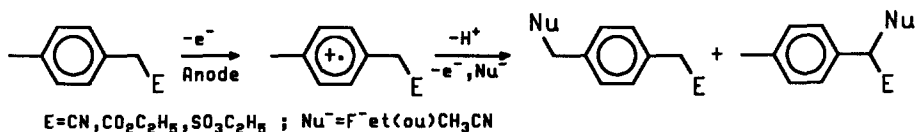


ETUDE DE LA REGIOSELECTIVITE EN FLUORATION
ANODIQUE DE DERIVES BENZYLQUES

E. Laurent, B. Marquet et R. Tardivel

UCB-Lyon I, Lab. de Chimie Organique 3, URA CNRS 467

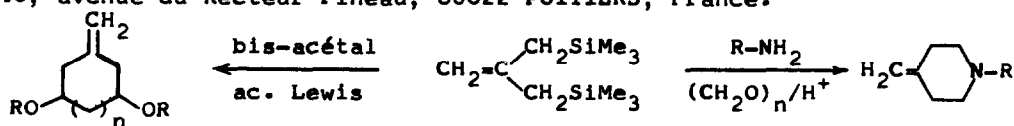
43, Bd du 11 Novembre 1918 69622 VILLEURBANNE Cedex (France)



Le 2-triméthylsilylméthyl allyltriméthylsilane,
précurseur de carbocycles et hétérocycles à
groupe méthylénique.

B. GUYOT, J. PORNET et L. MIGINIAC,

Laboratoire de Synthèse Organique, UA 574 CNRS, Université de Poitiers,
40, avenue du Recteur Pineau, 86022 POITIERS, France.

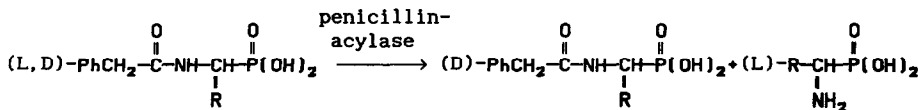


PREPARATION OF OPTICALLY ACTIVE 1-AMINOALKYL-
PHOSPHONIC ACIDS BY STEREOSELECTIVE ENZYMIC
HYDROLYSIS OF RACEMIC N-ACYLATED 1-AMINOALKYLPHOSPHONIC ACIDS

V. A. Solodenko,^a T. N. Kasheva,^a V. P. Kukhar,^a E. V. Kozlova,^b D. A. Mironenko^b
and V. K. Švedas^b,

^aInstitute of Bioorganic Chemistry of the Ukrainian Academy of
Sciences, (USSR),

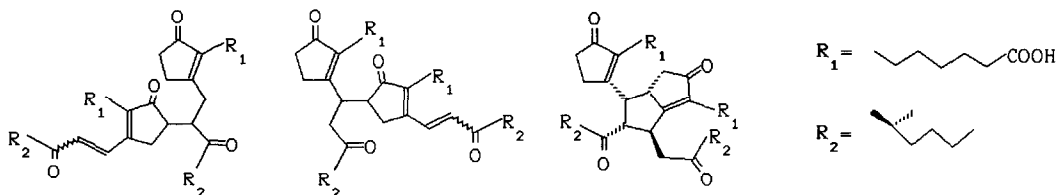
^bA. N. Belozersky Laboratory, Moscow State University, (USSR)



THE SYNTHESIS OF 16,16-DIMETHYL-15-KETO-PGB OLIGOMERS. THE CHEMICAL STRUCTURES OF DIMERS.

I. Martin^{*}, J. Anvelt, T. Pehk¹, Ü. Lille

Department of Prostanoid Chemistry, Institute of Chemistry, Akadeemia tee 15, Tallinn

200108, Estonia. ¹Department of Physics, Institute of Chemical Physics and Biophysics, Lenini pst. 10, Tallinn 200001, Estonia

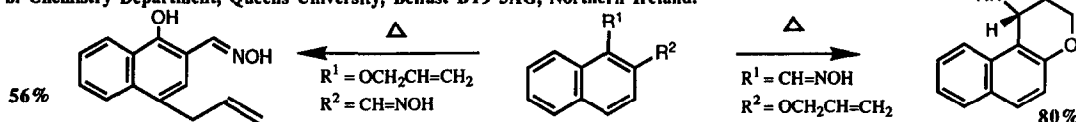
X=Y-ZH Systems as Potential 1,3- Dipoles. Part 31. Generation of Nitrones from Oximes. Background and Scope of the Tandem

1,2- Prototropy-Intramolecular Cycloaddition Sequence.

Ronald Grigg^a, Frances Heaney^b, Jasothara Markandu^a, Sivagnanasundram Surendrakumar^a, and William J. Warnock^a.

a. School of Chemistry, Leeds University, Leeds LS2 9JT.

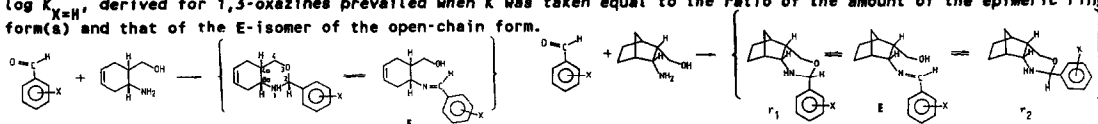
b. Chemistry Department, Queens University, Belfast BT9 5AG, Northern Ireland.



RING-CHAIN TAUTOMERISM AND THREE- TO FOUR-COMPONENT EQUILIBRIA IN FUSED-RING TETRAHYDRO-1,3-OXAZINES

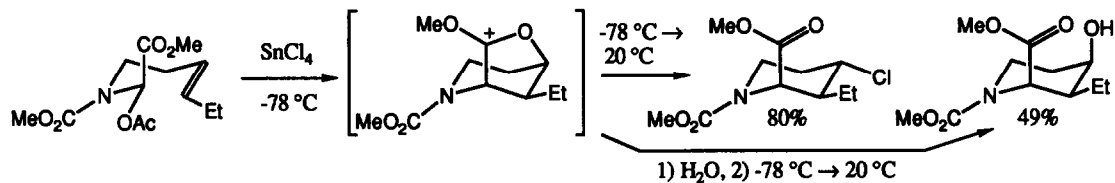
Kalevi Pihlaja,^{*,†} Aija Parkkinen,[†] Ferenc Fülöp,^{*,†} Jorma Mattinen,[†] and Gábor Bernáth[†][†]Department of Chemistry, University of Turku, SF-20500 Turku, Finland, and [†]Institute of Pharmaceutical Chemistry, Albert Szent-Györgyi Medical University, H-6701 Szeged, POB 121, Hungary

The condensation of *cis*- and *trans*-2-hydroxymethyl-4-cyclohexenyl-1-amine and 2,3-*diexo*- and 2,3-*diendo*-3-hydroxymethylbicyclo[2.2.1]heptylamine with aromatic aldehydes led to ring-chain tautomeric equilibria between epimeric tetrahydro-1,3-oxazines and open-chain Schiff bases. For each system studied a simple equation, $\log K_X = 0.76\sigma^+$ + $\log K_{X=H}$, derived for 1,3-oxazines prevailed when K was taken equal to the ratio of the amount of the epimeric ring form(s) and that of the E-isomer of the open-chain form.



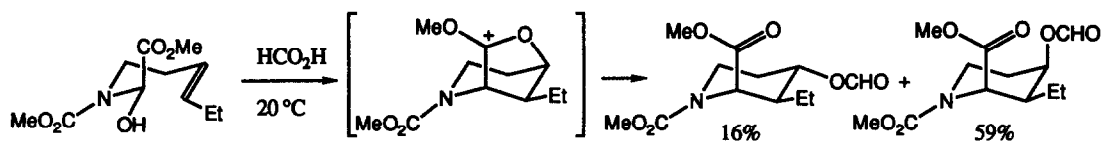
**TIN TETRACHLORIDE INDUCED π -CYCLIZATIONS OF
GLYCINE CATION EQUIVALENTS TO SUBSTITUTED PIPECOLIC ACID DERIVATIVES**

Peter M. Esch, Ilona M. Boska, Henk Hiemstra*, Richard F. de Boer, and W. Nico Speckamp*
Department of Organic Chemistry, University of Amsterdam, Nieuwe Achtergracht 129, 1018 WS Amsterdam,
The Netherlands



**FORMIC ACID INDUCED π -CYCLIZATIONS OF
GLYCINE CATION EQUIVALENTS TO SUBSTITUTED PIPECOLIC ACID DERIVATIVES**

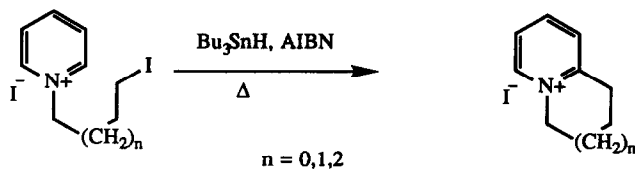
Peter M. Esch, Richard F. de Boer, Henk Hiemstra*, Ilona M. Boska, and W. Nico Speckamp*
Department of Organic Chemistry, University of Amsterdam, Nieuwe Achtergracht 129, 1018 WS Amsterdam,
The Netherlands



INTRAMOLECULAR FREE-RADICAL SUBSTITUTION OF PYRIDINIUM RINGS.

John A. Murphy* and Michael S. Sherburn, Department of Chemistry, University of Nottingham, Nottingham NG7 2RD.

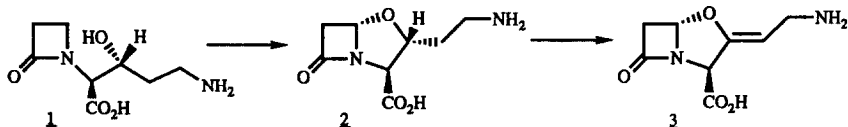
Intramolecular free radical substitution of pyridinium salts has been accomplished giving good yields of [6,5]-, [6,6]- and [6,7]- bicyclic compounds.



ISOLATION OF DIHYDROCLAVAMINIC ACID AN INTERMEDIATE IN THE BIOSYNTHESIS OF CLAVULANIC ACID

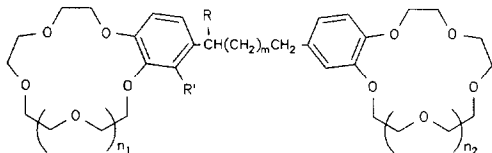
Jack E. Baldwin*, Robert M. Adlington, Justin S. Bryans, Alain O. Bringham, Janice B. Coates, Nicholas P. Crouch, Matthew D. Lloyd, and Christopher J. Schofield. The Dyson Perrins Laboratory and the Oxford Centre for Molecular Sciences, South Parks Road, Oxford, OX1 3QY, U.K., Stephen W. Elson*, Keith H. Baggeley, Robert Cassels, and Neville Nicholson, SmithKline Beecham Pharmaceuticals, Brockham Park, Betchworth, Surrey RH3 7AJ, U.K.

A primary isotope effect was used in an *in vitro* study to allow the isolation of an intermediate (2), between proclavaminic acid (1) and clavaminic acid (3) in clavulanic acid biosynthesis.



BIS(BENZOCROWN ETHER)S WITH POLYMETHYLENE BRIDGES AND THEIR APPLICATION IN ION-SELECTIVE ELECTRODES

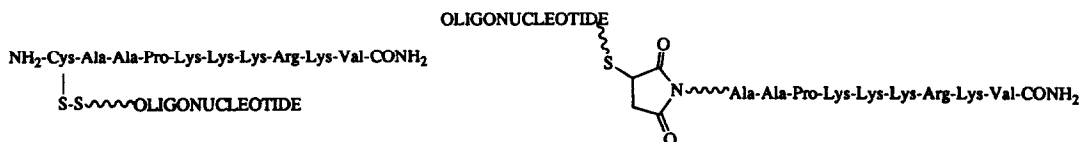
Elżbieta Luboch*, Andrzej Cygan and Jan F. Biernat
Faculty of Chemistry, Technical University of Gdańsk
80-952 Gdańsk, Poland

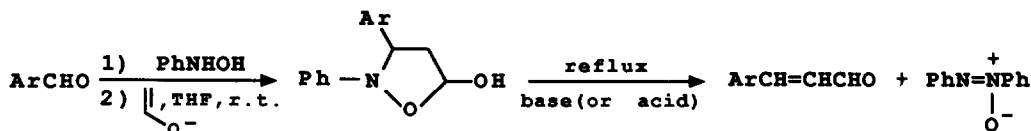


Synthetic procedures based on the condensation of benzocrown ether derivatives leading to bis(benzocrown ether)s with polymethylene bridges have been elaborated. Methods for introducing lipophilic substituents into these compounds also have been described. The reported compounds have been tested in ion-selective electrodes.

SYNTHESIS OF DEFINED PEPTIDE-OLIGONUCLEOTIDE HYBRIDS CONTAINING A NUCLEAR TRANSPORT SIGNAL SEQUENCE.

Ramon Eritja*, Anna Pons, Mónica Escarceller, Ernest Giralt[§], and Fernando Albericio[§]. Department of Molecular Genetics. CID-CSIC. Jordi Girona 18-26., 08034 Barcelona. Spain. [§]Department of Organic Chemistry. University of Barcelona. 08028 Barcelona, Spain.

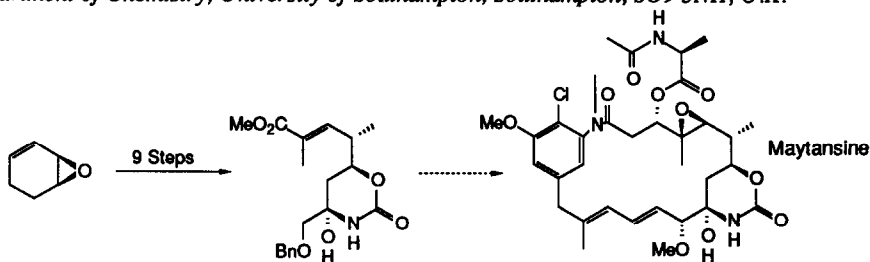
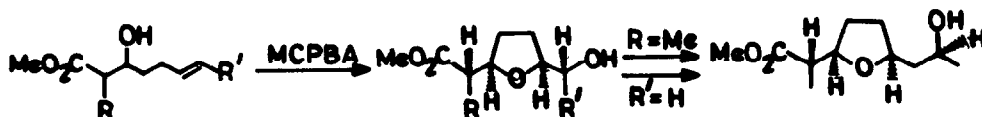


AN EFFICIENT C₂-HOMOLOGATION OF AROMATIC ALDEHYDES VIA 5-HYDROXYISOXAZOLIDINEL. Di Nunno ^{a,b} and A. Scilimati ^b^aCentro CNR di Studio sulle Metodologie Innovative di Sintesi Organiche, Dipartimento di Chimica, Università, Trav. 200 Re David 4, 70126 Bari-Italy.^bDipartimento Farmaco-Chimico, Università, Trav. 200 Re David 4, 70126 Bari-Italy.

A SHORT AND EFFICIENT SYNTHESIS OF THE C-3 TO C-10 PORTION OF THE MAYTANSINOIDS

David M. Hodgson,[†] Philip J. Parsons,^{†*} and Peter A. Stones

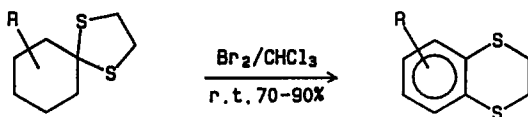
Department of Chemistry, University of Southampton, Southampton, SO9 5NH, U.K.

METACHLOROPEROXYBENZOIC ACID PROMOTED STEREOSELECTIVE SYNTHESIS OF 2,5-DISUBSTITUTED TETRAHYDROFURANS FROM α OR γ -ALLYL- β -HYDROXY ESTERS: A FORMAL SYNTHESIS OF (\pm) METHYL NONACTATEJaved Iqbal^{*}, Anu Pandey and Bhanu P.S. Chauhan
Department of Chemistry, Indian Institute of Technology, Kanpur 208016, India

REACTIVITY OF ETHANEDIYL S,S-ACETALS - 3.
RING AROMATIZATION IN CYCLOHEXANONE DERIVATIVES:
A NOVELTY SYNTHESIS OF 1,4-BENZODITHIANS

ROMUALDO CAPUTO, CARLA FERRERI, GIOVANNI PALUMBO*, FRANCESCO RUSSO
Dipartimento di Chimica Organica e Biologica dell'Università
Via Mezzocannone, 16 I-80134 Napoli (Italy)

A ready and reliable synthesis of 1,4-benzodithians with substituents on the benzenoid ring is reported for the first time.

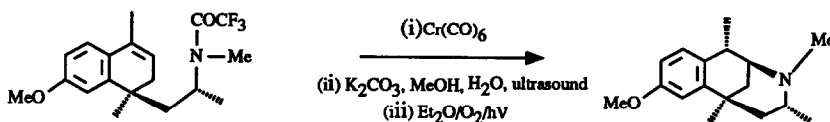


A CHIRAL SYNTHESIS OF A 6,7-BENZOMORPHAN

Malcolm Sainsbury*, Mary F. Mahon, and Colin S. Williams,

School of Chemistry, University of Bath, Claverton Down, Bath BA27AY, U.K. Alan Naylor and David I.C. Scopes, Medicinal Chemistry Department, Glaxo Group Research Ltd., Ware, Herts. SG12ODJ

The synthesis of (-)-(S),2(S),4(R),6(R)-1,2,3,4,5,6-hexahydro-2,6-methano-8-methoxy-1,3,4,6-tetramethyl-3-benzazocine is achieved through the chromium hexacarbonyl mediated cyclisation of 1(S),1'(R)-1,2-dihydro-7-methoxy-1,4-dimethyl-1-(N-methyl-N-trifluoroacetamido-1'-methyl-ethan-2'-yl)naphthalene



ON THE REACTION OF TRISUBSTITUTED OLEFINS
WITH PHENYLSELENYNYL CHLORIDE IN METHANOL.

P. Ceccherelli,* M. Curini, M.C. Marcotullio, O. Rosati; Istituto di Chimica Organica, Facoltà di Farmacia, Università di Perugia, Italy.

The interaction of cholesterol and *p*-menthene with excess phenylselenenyl chloride in methanol produced the corresponding *trans* chloromethoxy and dimethoxy-derivatives. The mechanism of these transformations has been investigated.