

GRAPHICAL ABSTRACTS

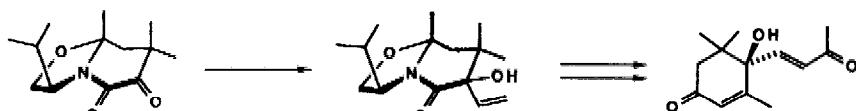
THE ASYMMETRIC TOTAL SYNTHESIS OF (-)-DEHYDROVOMIFOLIOLE.

Tetrahedron Lett. 30, 1741 (1989)

THE PENULTIMATE PRECURSOR TO (-)-ABSCISIC ACID (ABA)

A. I. Meyers* and Michael A. Sturgess

Department of Chemistry, Colorado State University, Fort Collins, CO 80523 USA



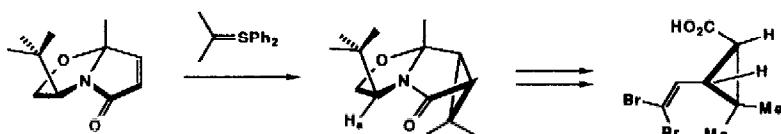
Vinyl Grignard addition gave > 100:1 selectivity

CHIRAL BICYCLIC LACTAMS. AN ASYMMETRIC SYNTHESIS OF CIS-(1S, 3R) DELTAMETHRINIC ACID

Tetrahedron Lett. 30, 1745 (1989)

A. I. Meyers* and Daniel Romo

Department of Chemistry, Colorado State University, Fort Collins, CO 80523 USA



Cyclopropanation of bicyclic lactams gives > 200:1 stereoselectivity.

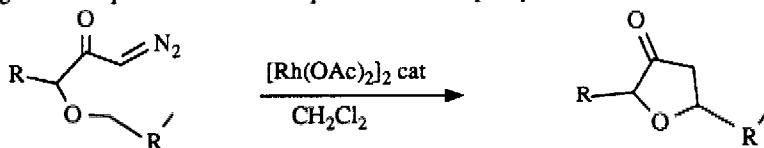
Rhodium Acetate Catalyzes the Addition of Carbenoids α - to Ether Oxygens

Tetrahedron Lett. 30, 1749 (1989)

Julian Adams*, Marc-André Poupart, Louis Grenier, Chris Schaller, Nathalie Ouimet*, and Richard Frenette*

Bio-Méga Inc. 2100 rue Cunard, Laval, Quebec, Canada H7S 2G5

Summary: Diazo-carbonyl compounds, when catalyzed by rhodium acetate, insert preferentially adjacent to ether oxygens. This phenomenon was exploited to develop a synthesis of 3(2H)-furanones.



Diastereoselectivity in the Synthesis of 3(2H)-Furanones.

Tetrahedron Lett. 30, 1753 (1989)

Total Synthesis of (+)-Muscarine

Julian Adams*, Marc-André Poupart, Louis Grenier

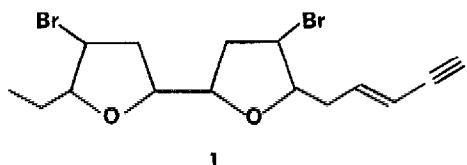
Bio-Méga Inc. 2100 rue Cunard, Laval, Quebec, Canada H7S 2G5

Summary: The carbonyl cyclization reaction to form disubstituted 2,5-3(2H)-furanones exhibited a stereoselection favoring the *cis* isomers. This phenomenon was exploited in an enantioselective synthesis of (+)-muscarine



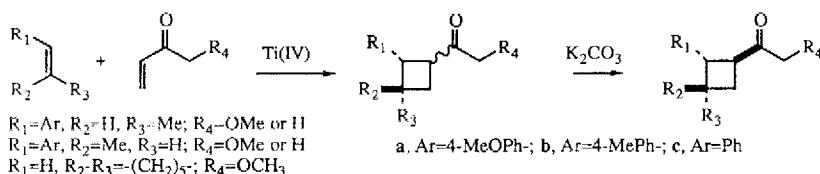
LAUROXOLANES FROM THE MARINE ALGA LAURENCIA MAJUSCULA
 In Kyu Kim, Mary R. Brennan, and Karen L. Erickson
 Jeppson Laboratory, Department of Chemistry
 Clark University, Worcester, Massachusetts 01610

Enyne 1 was isolated and its structure determined by spectroscopic means. 1 is the first 2,2'-bis tetrahydrofuran lauroxane to be found in a Laurencia species.



NON-PHOTOCHEMICAL 2+2 CYCLOADDITIONS OF ACYCLIC ENONES AND ALKENES

Thomas A. Engler,* Mohammed Hashmat Ali and David Vander Velde, Department of Chemistry, University of Kansas, Lawrence, KS 66045.

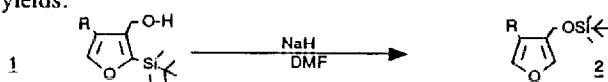


THE 1,4 C→O SILYL MIGRATIONS OF VARIOUS FURAN AND THIOPHENE SYSTEMS

Patrick G. Spinazzé and Brian A. Keay*

Department of Chemistry and Biochemistry, University of Windsor, Windsor, Ontario, Canada, N9B 3P4

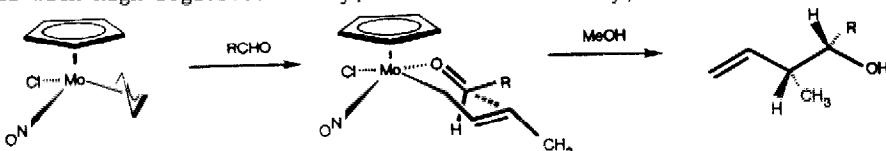
2-Trialkylsilyl-3-hydroxymethyl-furans (**1**) and -thiophenes undergo a 1,4 C→O silyl migration when treated with bases containing potassium or sodium counterions to produce 3-[trialkylsilyloxy]methyl-furans (**2**) and -thiophenes in excellent yields.



CONTROLLING STEREOCHEMISTRY IN CROTYL ADDITIONS TO ALDEHYDES WITH CROTYLMOLYBDENUM COMPLEXES

J.W. Faller*, J.A. John and M.R. Mazzieri, Dept. Chemistry, Yale University, New Haven, CT

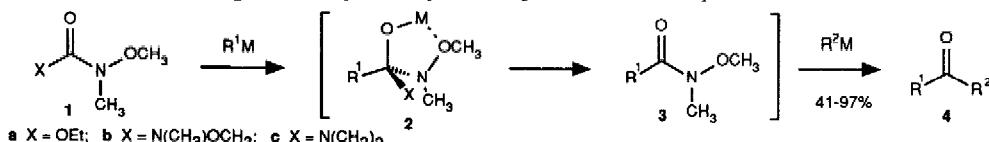
(Cyclopentadienyl)Mo(NO)(Cl)(π -crotyl) complexes add to aldehydes to yield homoallylic alcohols with high regioselectivity, diastereoselectivity, and enantioselectivity.



**TANDEM ORGANOMETALLIC ADDITION REACTIONS
TO N-METHOXY-UREAS AND URETHANES IN THE**

PREPARATION OF UNSYMMETRICAL AND SYMMETRICAL KETONES, Dennis J. Hlasta* and John J. Court, Sterling Research Group, Department of Medicinal Chemistry, Rensselaer, New York 12144

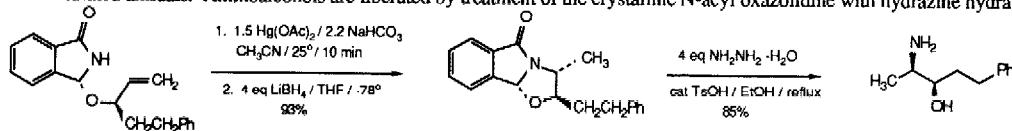
The use of the novel reagents **Ia-c** for the synthesis of ketones in a one pot reaction is described.



A Removable Auxiliary for Amidomercuration Reactions: The Stereocontrolled Preparation of Vicinal Aminoalcohols.

James M. Takacs*, Mark A. Helle, and Linrong Yang
Department of Chemistry, University of Nebraska-Lincoln, Lincoln, NE 68588-0304

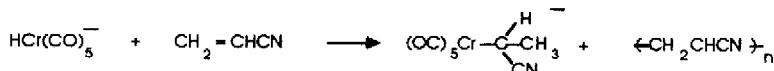
A phthalimide-derived auxiliary provides the dominant influence in directing the stereochemical course of the amidomercuration cyclization of certain unsaturated amidals. Aminoalcohols are liberated by treatment of the crystalline N-acyl oxazolidine with hydrazine hydrate.



ADDITION OF ACRYLONITRILE TO ANIONIC TRANSITION METAL HYDRIDES. AN IONIC MODEL FOR HOMOGENEOUS OLEFIN PROCESSES

Marcetta Y. Dahrensbourg*, Barbara Floris, and Kay A. Youngdahl
Department of Chemistry, Texas A&M University, College Station, Texas 77843

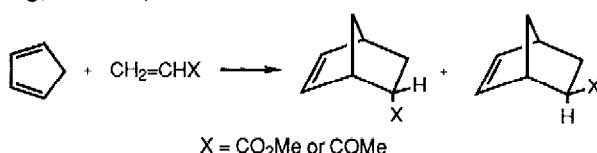
Kinetic and substituent effects suggest the following (and other activated olefins) to react by an associative hydride transfer mechanism.



**DIELS-ALDER REACTIONS IN ETHYLAMMONIUM NITRATE,
A LOW-MELTING FUSED SALT**

David A. Jaeger* and Charles E. Tucker
Department of Chemistry, University of Wyoming, Laramie, WY 82071

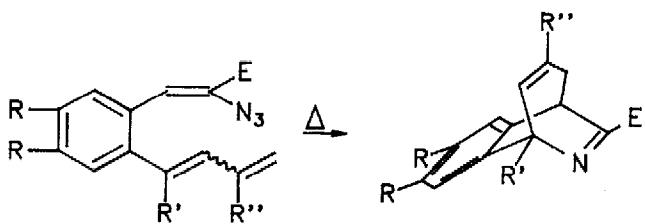
Relative to nonpolar organic solvents, ethylammonium nitrate gave endo selectivity enhancements for the illustrated reactions.



A SYNTHESIS OF THE 6-AZA-BICYCLO(3.2.2)NONANE SKELETON

Claus Vogel* and Paul Delavrier
Inst. f. Organische Chemie, Hagenring 30, D-3300 Braunschweig, FRG

The title skeleton is formed by an intramolecular reaction of vinyl-azides with 1,3-dienes.

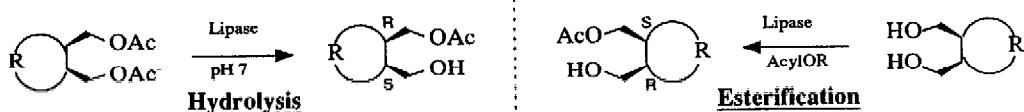


ENZYMATIC ESTER HYDROLYSIS AND SYNTHESIS -

TWO APPROACHES TO CYCLOALKANE DERIVATIVES OF HIGH ENANTIOMERIC PURITIES

Ulrich Ader, Detlef Breitgoff, Peter Klein, Kurt E. Laumen, Manfred P. Schneider*

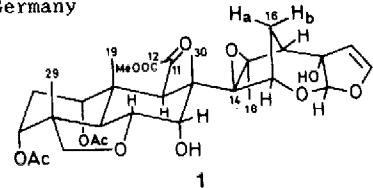
Fb 9 - Bergische Universität, D-5600 Wuppertal 1, Germany



1,3-DIACETYL-11,19-DEOXA-11-OXO-MELIACARPIN, A POSSIBLE PRECURSOR OF AZADIRACHTIN, FROM *AZADIRACHTA INDICA* A. JUSS (MELIACEAE)

W. Kraus*, H. Gutzeit, and M. Bokel, Institut für Chemie der Universität Hohenheim, Lehrstuhl für Organische Chemie, Garbenstr. 30, 7000 Stuttgart 70, West Germany

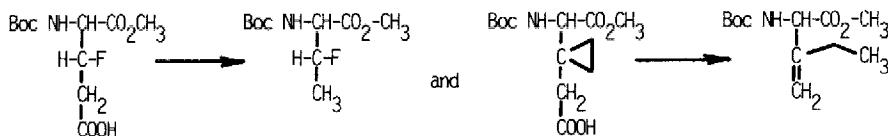
1,3-Diacetyl-11,19-deoxa-11-oxo-meliacarpin (1), a possible intermediate in the biosynthesis of azadirachtin, was isolated from methanolic extracts of *Azadirachta indica* seeds. Structure 1 is proposed on the basis of ¹H and ¹³C NMR data.



HOMOLYTIC DECARBOXYLATION OF GLUTAMATE ANALOGUES

Anne Vidal-Cros, Sonia Bory, Michel Gaudry and Andrée Marquet

Laboratoire de Chimie Organique Biologique - Univ. P. & M. Curie - 4, place Jussieu - 75252 PARIS CEDEX 05 - France



Probes designed to study vitamin K-dependent carboxylation

ASYMMETRIC INDUCTION IN PALLADIUM CATALYZED [3+2] CYCLOADDITION
REACTION OF TRIMETHYLENEMETHANE WITH HOMOCHIRAL VINYL SULFOXIDES

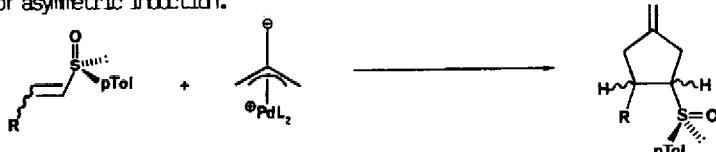
Tetrahedron Lett. 30, 1803 (1989)

Frantz Chaigne, Jean-Pierre Gotteland and Max Malacia*

Laboratoire de Chimie Organique I, U.A 467 du CNRS, Université Claude Bernard

ESCL - 43 Bd du 11 Novembre 1918, 69622 VILLEURBANNE, France.

Asymmetric [3+2] cycloaddition using homochiral vinylsulfoxides proceed in good to excellent chemical yields and with a good level of asymmetric induction.



STRUCTURES OF ISOMBAMICHALCONE AND LOPHIROCHALCONE,
BI- AND TETRA-FLAVONOIDS FROM *LOPHIRA LANCEOLATA*

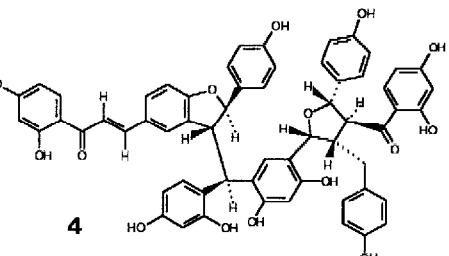
Tetrahedron Lett. 30, 1807 (1989)

R. GHOGOMU TIH, B.L. SONDENGAM, M.T. MARTIN* and B. BODO*

Département de Chimie organique, Université de Yaoundé, B.P. 812, Yaoundé, Cameroun.

*Laboratoire de Chimie, Muséum national d'Histoire naturelle, 63, rue Buffon,
75231 Paris Cedex 05, France.

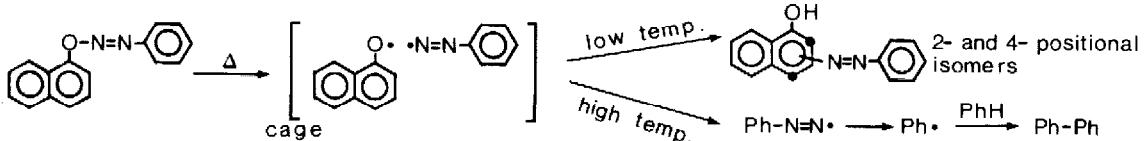
From the stem bark of *Lophira lanceolata* two new flavonoids, a chalcone dimer, isombamichalcone 1, and a tetraflavonoid, lophirochalcone 4 have been isolated and their structures established from spectral and chemical evidences.



THE RADICAL COUPLING MECHANISM IN THE DIAZO COUPLING REACTION. MIGRATION VS. DECOMPOSITION OF PHENYLAZO RADICAL GENERATED FROM PHENYLAZO 1-NAPHTHYL ETHER IN THE SOLVENT CAGE

Tetrahedron Lett. 30, 1811 (1989)

Takahiro Tezuka, Hiroharu Tanikawa, Katsunori Sasaki, and Harumi Tajima
Department of Chemistry, University of Tsukuba, Tsukuba, Ibaraki 305, Japan

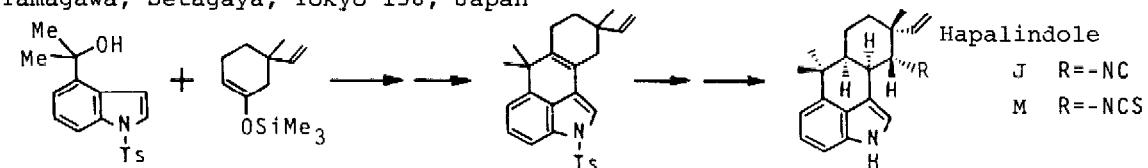


TOTAL SYNTHESIS OF MARINE ALKALOIDS
 (\pm) -HAPALINDOLES J AND M

Tetrahedron Lett. 30, 1815 (1989)

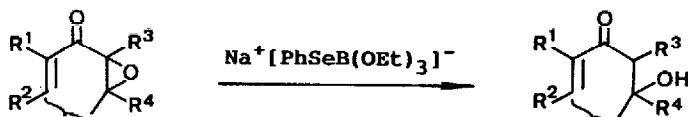
Hideaki Muratake and Mitsutaka Natsume*

Research Foundation Itsuu Laboratory
Tamagawa, Setagaya, Tokyo 158, Japan



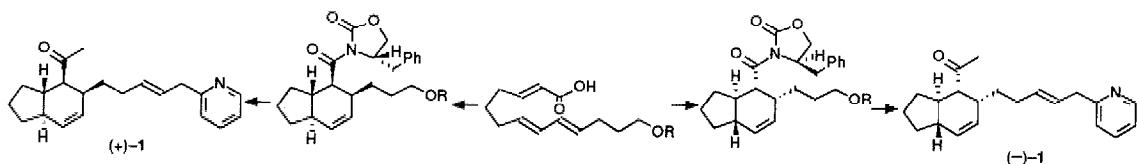
CHEMOSELECTIVE REDUCTION OF AN α,β -EPOXY KETONE MOIETY COEXISTING WITH AN ENONE FUNCTION

Masaaki Miyashita, Toshio Suzuki, and Akira Yoshikoshi*
Chemical Research Institute of Non-Aqueous Solutions,
Tohoku University, Sendai 980, Japan



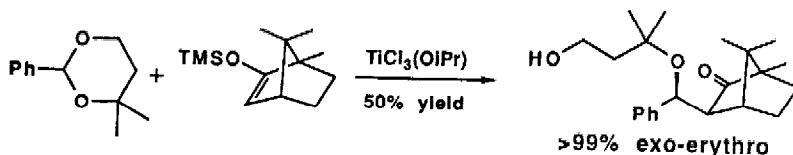
ASYMMETRIC TOTAL SYNTHESSES OF (+)- and (-)-PULO'UPONE

Tsutomu Sugahara, Takashi Iwata, Miyako Yamacka, and Seiichi Takanishi
Pharmaceutical Institute, Tohoku University, Aobayama, Sendai 980, Japan



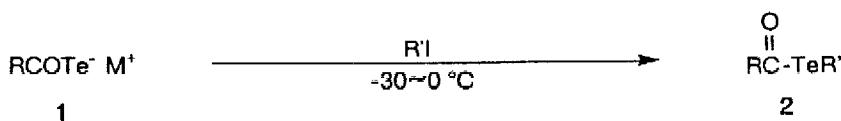
DIASTEREOSELECTIVE ALDOL SYNTHESIS USING ACETAL TEMPLATES

Kazuaki Ishihara and Hisashi Yamamoto*, Department of Applied Chemistry, Nagoya University, Chikusa, Nagoya 464-01, Japan; Clayton H. Heathcock, Department of Chemistry, University of California, Berkeley, California 94720



Te-ALKYL TELLUROCARBOXYLATES — ISOLATION AND CHARACTERIZATION

Takahiro Kanda, Shoho Nakaiida, Toshiaki Murai, and Shinzi Kato*
Department of Chemistry, Faculty of Engineering, Gifu University,
Yanagido, Gifu 501-11 Japan



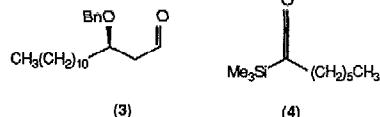
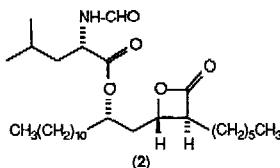
A SYNTHESIS OF (-)-TETRAHYDROLIPSTATIN

Jean-Marc Pons and Philip Kocienski

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

Tetrahedron Lett. 30, 1833 (1989)

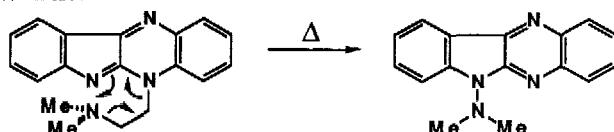
Cycloaddition of n-hexyl trimethylsilyl ketene (4) to (3R)-3-(benzyloxy)-tetradecanal (3) is the key step in a synthesis of the anti-obesity agent (-)-tetrahydrolipstatin (2).



**Formation of N-N Bonds by Thermolysis of
5-(2-Dimethylaminoethyl)-5H-indolo[2,3-b]quinoxaline.**

Jan Bergman, Royal Institute of Technology, Department of Organic Chemistry,
S-100 44 Stockholm, SWEDEN

Tetrahedron Lett. 30, 1837 (1989)

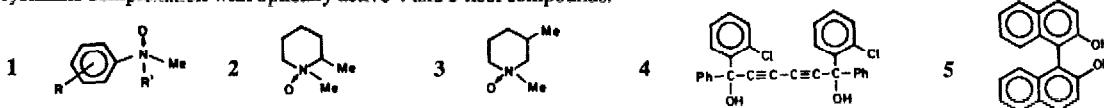


**OPTICAL RESOLUTION OF AMINE N-OXIDE BY DIASTEREOMERIC
COMPLEX FORMATION WITH OPTICALLY ACTIVE HOST COMPOUND**

Fumio Toda* and Koji Mori, Department Of Industrial Chemistry, Ehime University, Matsuyama 790, Japan
Zafra Stein and Israel Goldberg*, School of Chemistry, Tel Aviv-University, 69978 Ramat-Aviv, Israel

Tetrahedron Lett. 30, 1841 (1989)

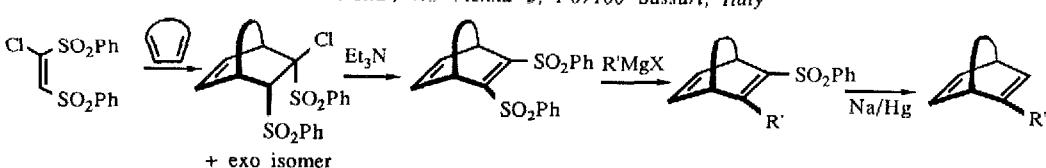
Enantiomers of 1 and diastereoisomers of 2 and 3 were resolved, and their absolute configuration was determined, by selective crystalline complexation with optically active 4 and 5 host compounds.



**1,2-BIS(PHENYLSULFONYL)ALKENES AS VERSATILE GROUPS
IN ORGANIC SYNTHESIS: THE PREPARATION OF ALKYL- AND
ARYL-SUBSTITUTED NORBORNADIENES VIA THE DIELS-ALDER CYCLOADDITION - GRIGNARD REACTION
- DESULFONYLATION SEQUENCE**

Ugo Azzena, Sergio Cossu, Ottorino De Lucchi,* and Giovanni Melloni
Dipartimento di Chimica dell'Università, via Vienna 2, I-07100 Sassari, Italy

Tetrahedron Lett. 30, 1845 (1989)

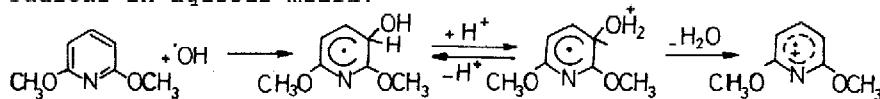


A CATION RADICAL FROM 2,6-DIMETHOXYPYRIDINE
INVESTIGATIONS BY IN-SITU-RADIOLYSIS ESR

Siddik İcli

Department of Chemistry, Ege University, Izmir, TURKEY

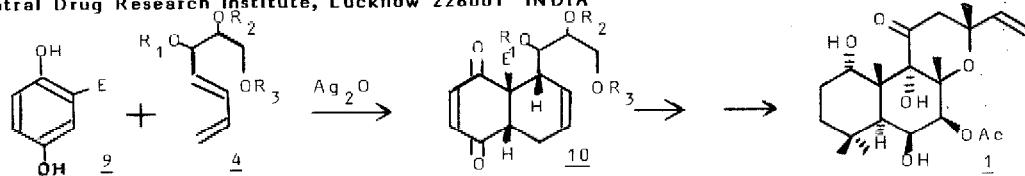
Formation and detection of a heteroaromatic cation radical in aqueous media.



DIELS-ALDER REACTION OF IN-SITU GENERATED
2-METHOXYCARBONYL- β -QUINONE WITH D-GLUCOSE
BASED DIENES: A NEW APPROACH TO FORSKOLIN

Aloka Mukhopadhyay, S.M. Ali, Mashkoor Husain, S.N. Suryawanshi * and D.S. Bhakuni *

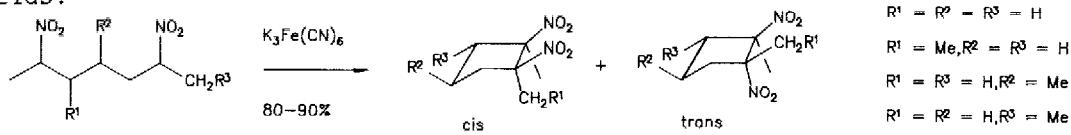
Central Drug Research Institute, Lucknow 226001 INDIA



RADICAL CYCLISATION OF 2,6-DINITROALKANES

W. Russell Bowman* and Stuart W. Jackson
Department of Chemistry, University of Technology, Loughborough, Leics.

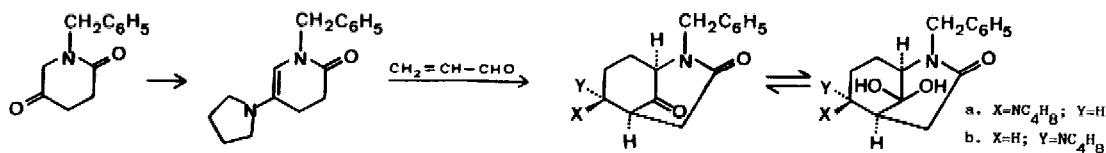
The dinitronate dianions of 2,6-dinitroalkanes were oxidised to 1,2-dinitro-cyclopentanes via α -nitroalkyl radicals by stereoselective cyclisation in high yields.

FUNCTIONALIZED 2-AZABICYCLO[3.3.1]NONANES. X. α , α' -ANNELATION

OF 1-BENZYL-2,5-PIPERIDINEDIONE AND ISOLATION OF AN AZABICYCLIC ADDUCT AS STABLE HYDRATE

Josep Bonjoch*, Josefina Quirante, Isabel Serret, and Joan Bosch*

Laboratory of Organic Chemistry, Faculty of Pharmacy, University of Barcelona, 08028-Barcelona, Spain.



SYNTHESIS OF 2,2-DIALKYL-1-AMINOCYCLOPROPANE-CARBOXYLIC ACIDS FROM α -CHLOROIMINES

N. De Kimpe*, P. Sulmon, P. Brunet, F. Lambein, N. Schamp,
Fac. Agric. Sciences, Univ. of Gent, Gent, Belgium

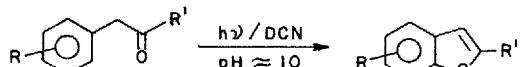
The first examples of 2,2-dialkyl-ACC analogues were synthesized by cyclopropanation of α -chloroimines.



A ONE STEP SYNTHESIS OF 2-SUBSTITUTED BENZOFURANS FROM 2-ARYL-1-SUBSTITUTED ETHANE-1-ONES BY PHOTO-INDUCED SET REACTIONS

G. Pandey*, A. Krishna and U.T. Bhalerao
Organic Division, RRL, Hyderabad 500 007, India

An efficient one step synthesis of 2-substituted benzofurans from the enolate of 2-Aryl-1-substituted ethane-1-ones have been reported.



STEREOSELECTIVE SYNTHESIS OF TILIVALLINE

Tatsuo Nagasaka, * Yuji Koseki, and Fumiko Hamaguchi
Tokyo College of Pharmacy
Horinouchi, Hachioji, Tokyo 192-03, Japan

A convenient and stereoselective synthesis of tilivalline, a metabolite from Klebsiella pneumoniae var. oxytoca, is described.

