

## GRAPHICAL ABSTRACTS

*Tetrahedron*, 1992, 48, 5941

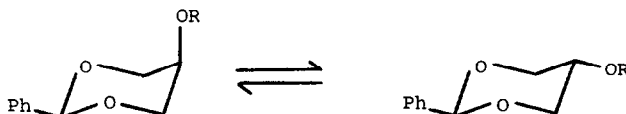
### Conformational Analysis of 5-Substituted 1,3-Dioxanes.

#### 6. Study of the Attractive Gauche Effect in O-C-C-O

##### Segments

Eusebio Juaristi\* and Sandra Antúnez, Depto. Química, CINVESTAV-IPN, Apdo. Postal 14-740, 07000 México, D.F., México.

NMR and quantum mechanical studies of several 5-O-substituted-1,3-dioxanes appear to support a stereoelectronic  $\sigma_{C-H} \rightarrow \sigma_{C-OR}^*$  interpretation of the attractive *gauche* effect.

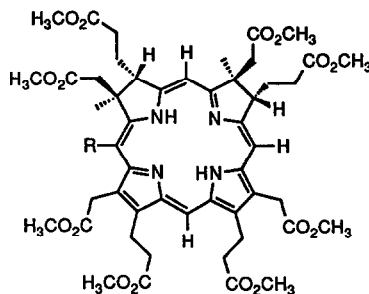


*Tetrahedron*, 1992, 48, 5951

### <sup>1</sup>H AND <sup>13</sup>C NMR STUDIES OF SIROHYDROCHLORIN (FACTOR II) AND ITS 20-METHYL DERIVATIVE (FACTOR III).

Mario D. Gonzalez, Howard J. Williams, Patricio J. Santander, Shin-ichi Ozaki, Neal J. Stolowich, and A. Ian Scott\*. Center for Biological NMR, Texas A & M University, College Station, Texas, 77843, USA.

<sup>1</sup>H and <sup>13</sup>C NMR spectra of Sirohydrochlorin octamethylester (R = H) and its 20-methyl derivative (R = CH<sub>3</sub>) have been assigned. Evidence for the presence of tautomeric forms of differing stabilities is presented.



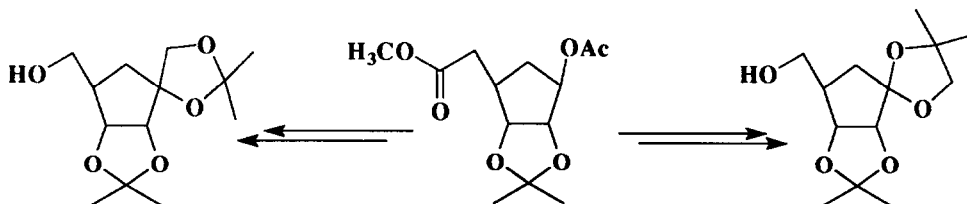
*Tetrahedron*, 1992, 48, 5961

### SYNTHESIS OF PROTECTED (±)-α- AND β-CARBA-PSICOFURANOSE

Michael Bodenteich and Victor E. Marquez\*

Laboratory of Medicinal Chemistry, DTP, DCT, NCI, NIH, Bethesda, MD 20892.

Protected racemic α- and β-psicofuranose were synthesized from a non-carbohydrate precursor which is available in both enantiomeric forms.

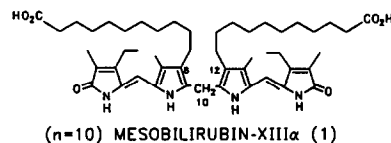


*Tetrahedron*, 1992, 48, 5969

# SYNTHESIS AND CONFORMATION OF A BILIRUBIN ANALOG WITH PROPIONIC ACID SIDE CHAINS EXTENDED TO UNDECANOIC ACID

John Chiefari, Richard V. Person and David A. Lightner\*  
Department of Chemistry, University of Nevada, Reno

A lipophilic analog of bilirubin (1) with C-8 and C-12 propionic acid groups replaced by undecanoic acids was synthesized and found to participate in conformation-determining intramolecular hydrogen bonding by <sup>1</sup>H-NMR analyses and molecular dynamics calculations.



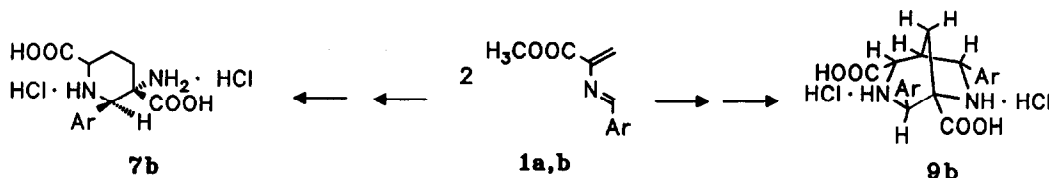
*Tetrahedron*, 1992, 48, 5985

# Preparation of Novel Heterocyclic Amino Acids from N-(Arylmethylene)dehydroalanine Methyl Esters

Günter Wulff\*, Heinz Theo Klinken

Institute of Organic Chemistry and Macromolecular Chemistry, Heinrich-Heine University Düsseldorf, Universitätsstr. 1, 4000 Düsseldorf 1 F.R.G.

From N-(arylmethylene)dehydroalanine methyl esters by dimerization and further transformations the non-proteinogenic amino acids derived from 3-amino-piperidine-3,6-dicarboxylic acids or 2,6-diazabicyclo[3.2.1]octane-1,5-dicarboxylic acids are formed.



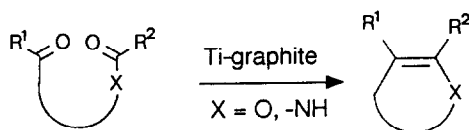
*Tetrahedron*, 1992, 48, 5991

# Titanium-Induced Syntheses of Furans, Benzofurans and Indoles

A. Fürstner,\* D.N. Jumbam

Institute of Organic Chemistry, Technical University A-8010 Graz, Austria

Intramolecular McMurry-type reactions of acyloxy- and acylamido carbonyl compounds

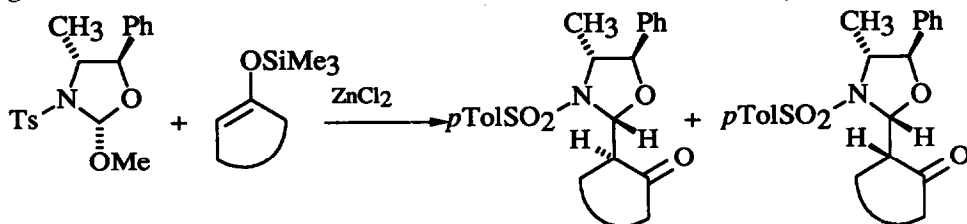


*Tetrahedron*, 1992, 48, 6011

**Norpseudoephedrine Derived 2-Methoxy-3-sulfonyl-1,3-oxazolidines:  
Chiral, Highly Diastereoselective Formyl Cation Equivalents**

Kilian Conde-Frieboes, Dieter Hoppe\*

Institut für organische Chemie der Universität Kiel, Olshausenstr. 40, D-2300 Kiel, Germany



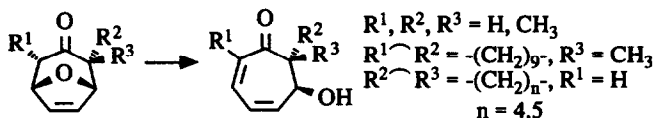
*Tetrahedron*, 1992, 48, 6021

**SYNTHESIS OF NONENOLIZABLE  
6-HYDROXY-2,4-CYCLOHEPTADIEN-1-ONES**

I. Stohrer, H.M.R. Hoffmann

Department of Organic Chemistry, University of Hannover, Schneiderberg 1 B, 3000 Hannover, Germany

8-Oxabicyclo[3.2.1]oct-6-en-3-ones with a quaternary centre adjacent to the carbonyl group have been converted into 6-hydroxy-2,4-cycloheptadien-1-ones with zirconium tetrachloride/piperidine (1:1).

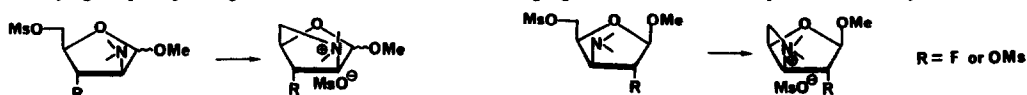


*Tetrahedron*, 1992, 48, 6033

**FORMATION COMPETITIVE D'IONS AZETIDINIUM, PYRROLIDINIUM ET AZIRIDINIUM LORS DE LA FLUORATION DE QUELQUES DIMETHYLAMINOPENTOFURANOSIDES DI-O-MESYLES**

M.B. Giudicelli, D. Picq et D. Anker\*, UCB-Lyon I, Lab. de Chimie Organique 3, associé au CNRS, 43 Boulevard du 11 Novembre 1918, 69622 VILLEURBANNE Cedex (France)

Furanosidic dimethylaminomesylates easily give stable corresponding pyrrolidinium or azetidinium ions even if R is a O-mesyl group; opening of these ions with fluorinating agents seems to be impossible or very difficult.

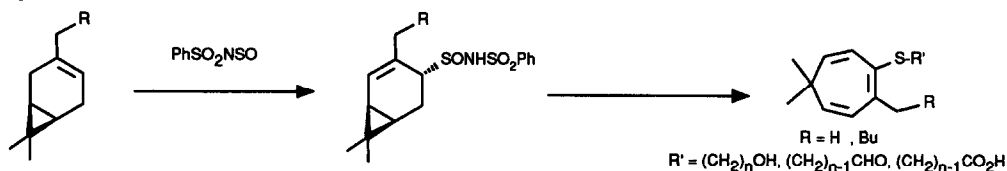


**A NEW ENE REACTION MEDIATED ACCESS TO CYCLOHEPTATRIENE ENOL-THIOETHERS**

Sylvie GEHANNE, Françoise RAYNAUD, Alain GADRAS and Gérard DELERIS

Laboratoire de Chimie BioOrganique INSERM U8, Université de Bordeaux II, 33076 Bordeaux, France.

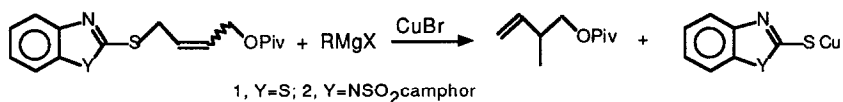
A synthesis of cycloheptatriene enol-thioethers via an ene reaction with sulfinyl sulfonamides and subsequent Grignard alkylation and ring enlargement.



**REGIOCONTROL BY ANCHIMERIC CO-ORDINATION IN THE REACTIONS OF ORGANOCOPPER REAGENTS WITH ALLYLIC SUBSTRATES BEARING TWO LEAVING GROUPS. SYNTHESIS OF OPTICALLY ACTIVE HOMOALLYLIC ALCOHOLS**

Vincenzo Calò<sup>a</sup>, Concetta De Nitti<sup>a</sup>, Luigi Lopez<sup>a</sup> and Antonio Scilimati<sup>b</sup>

<sup>a</sup>Centro CNR di Studio sulle Metodologie Innovative di Sintesi Organiche, Dipartimento di Chimica, Università di Bari, via Amendola n.173-70126 Bari-Italy. <sup>b</sup>Dipartimento Farmacochimico, Università di Bari, Trav.200 Re David n.4-70126 Bari-Italy.



**SPIROHETEROCYCLES FROM REACTION OF NITRILE OXIDES WITH HETEROCYCLIC α-QUINODIMETHANES**

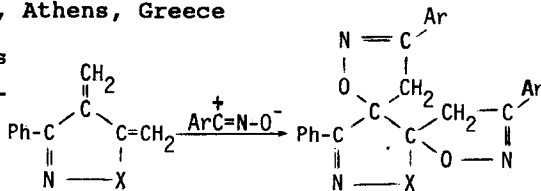
Sophia Mitkidou and Julia Stephanidou-Stephanatou\*

Laboratory of Organic Chemistry, University of Thessaloniki, Greece

Aristides Terzis and Demetrios Mentzafos

X-Ray Laboratory, NCR "Democritos", Athens, Greece

1,3-Dipolar cycloaddition reactions of stable nitrile oxides to heterocyclic α-quinodimethanes leading to the formation of dispiro-isoxazolines are described.



## REGIOSELECTIVE NUCLEOPHILIC RING OPENING

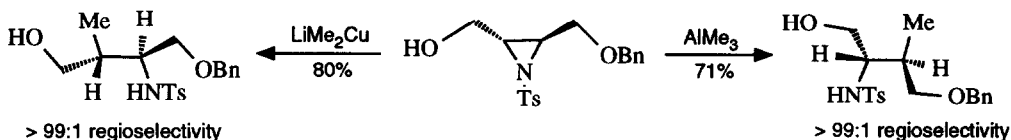
## OF 2,3-AZIRIDINO ALCOHOLS

David Tanner\* and Hua Ming He, Dept. of Organic Chemistry, University of Uppsala,

Box 531, S-751 21 Uppsala, Sweden

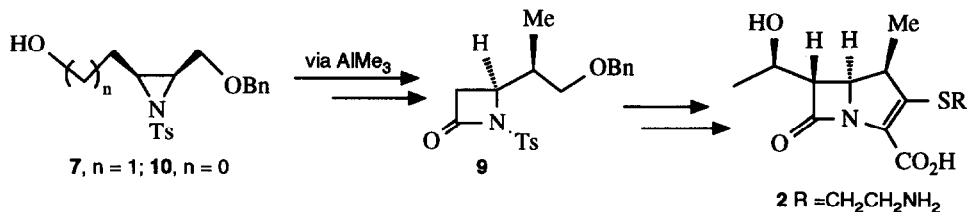
Peter Somfai, Organic Chemistry 2, Lund Institute of Technology, Box 124, S-221 00 Lund, Sweden

Excellent regioselectivity can be attained in the ring-opening of the title compounds, e.g.:

ENANTIOSELECTIVE ROUTES TOWARD 1 $\beta$ -METHYLCARBAPENEMS

## FROM CHIRAL AZIRIDINES

David Tanner\* and Hua Ming He, Dept. of Organic Chemistry, University of Uppsala, Box 531, S-751 21 Uppsala, Sweden

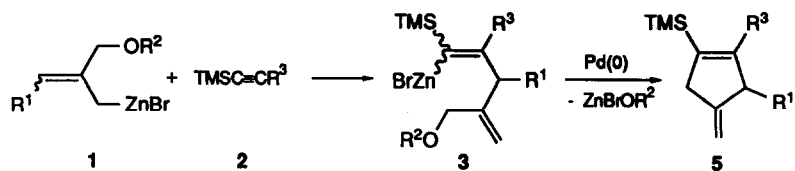
Enantioselective routes toward 1 $\beta$ -methylcarbapenems, e.g. **2**, from the chiral aziridines **7** or **10** are described.A key step is the completely regioselective ring-opening of the aziridines by  $\text{AlMe}_3$ .

## ALLYLMETALLATION OF 1-SILYLALKYNES BY

## 2-(BROMOZINC METHYL)-2-ALKENYL ETHERS FOLLOWED BY Pd(0)-CATALYZED CYCLIZATION: A ONE-POT SYNTHESIS OF 4-METHYLENOCYCLOPENTENES

J. van der Louw, J.L. van der Baan, F.J.J. de Kanter, F. Bickelhaupt, G.W. Klumpp

Scheikundig Laboratorium, Vrije Universiteit, De Boelelaan 1083, 1081 HV Amsterdam, The Netherlands

4-Methylenecyclopentenones **5** were prepared by carbocationization of **2** by **1** followed by Pd(0)-catalyzed cyclization of **3**.

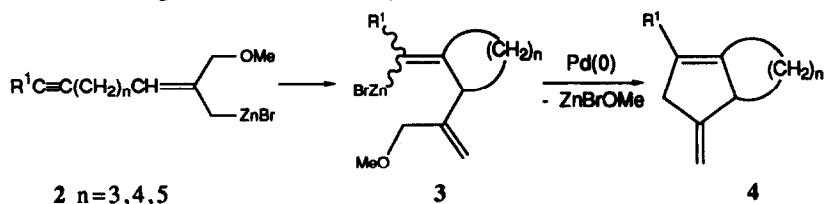
*Tetrahedron*, 1992, 48, 6105

#### INTRAMOLECULAR ZINC-ENE REACTIONS OF ALKYNES;

#### PREPARATION OF 1,5-ANNULATED 4-METHYLENECYCLOPENTENES

J. van der Louw, J.L. van der Baan, C.M.D. Komen, A. Knol, F.J.J. de Kanter, F. Bickelhaupt, G.W. Klumpp  
Scheikundig Laboratorium, Vrije Universiteit, De Boelelaan 1083, 1081 HV Amsterdam, The Netherlands

Zinc-ene reaction of 2 gave 3, which on Pd(0)-catalyzed cyclization was converted to 4.



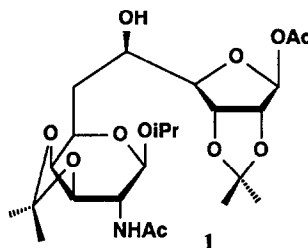
*Tetrahedron*, 1992, 48, 6123

#### NEW CONVENIENT SYNTHESIS OF TUNICAMINE

J. Ramza and A. Zamojski

*Institute of Organic Chemistry, Polish Academy of Sciences, 01-224 Warsaw*

A ten-step synthesis of tunicamine derivative 1, starting from isopropyl 2-azido-2-deoxy-3,4-*O*-isopropylidene- $\beta$ -D-galactopyranoside, is described.

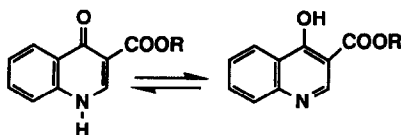


*Tetrahedron*, 1992, 48, 6135

#### TAUTOMERISM AND ACIDITY IN 4-QUINOLONE-3-CARBOXYLIC ACID DERIVATIVES

Angeles de la Cruz, José Elguero, Pilar Goya, Ana Martinez  
Instituto de Química Médica (CSIC), Juan de la Cierva 3, 28006 Madrid (Spain)  
Wolfgang Pfeleiderer  
Fakultat für Chemie, Universität Konstanz, D-7750 Konstanz (Germany)

Prototropic tautomerism in 4-quinolone-3-carboxylic acid derivatives has been studied. The techniques used include UV,  $^1\text{H-NMR}$ ,  $^{13}\text{C-NMR}$  (solution and solid state) and semiempirical and *ab initio* calculations.

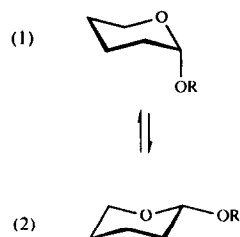


**EXPERIMENTAL STUDIES OF THE ANOMERIC EFFECT. PART V THE INFLUENCE OF SOME SOLVENTS ON THE CONFORMATIONAL EQUILIBRIA IN 2-METHOXY- AND 2-(2',2',2'-TRIFLUOROETHOXY)-TETRAHYDROPYRAN**

Harold Booth,\* J. Mark Dixon and Simon A. Readshaw

Department of Chemistry, The University, Nottingham, NG7 2RD

$\Delta H_{1 \rightarrow 2}^{\circ}$  and  $\Delta S_{1 \rightarrow 2}^{\circ}$  values are measured for ( $1 \rightleftharpoons 2$ ; R=Me,  $\text{CH}_2\text{CF}_3$ ) in ether/toluene and ether/methanol, and for ( $1 \rightleftharpoons 2$ ; R=Me) in  $\text{CD}_2\text{Cl}_2$  and ether/ $\text{CD}_2\text{Cl}_2$ . Replacement of toluene by methanol causes increases in  $\Delta H_{1 \rightarrow 2}^{\circ}$  and  $\Delta S_{1 \rightarrow 2}^{\circ}$  values, consistent with a preferential H-bonding to the exo-cyclic O of axial conformations



**EXPERIMENTAL STUDIES OF THE ANOMERIC EFFECT. PART VI RING INVERSION EQUILIBRIA IN CYCLOHEXANE, TETRAHYDROPYRAN AND PIPERIDINE RINGS SUBSTITUTED BY A CARBOMETHOXY OR A CYANO GROUP**

Harold Booth,\* J. Mark Dixon and Khedhair A. Khedhair

Department of Chemistry, The University, Nottingham, NG7 2RD

A variable temperature  $^{13}\text{C}$  nmr study of ( $1 \rightleftharpoons 2$ ; Y= $\text{CH}_2$ , O, NH; X= $\text{CO}_2\text{Me}$ , CN) gives values of  $\Delta H_{1 \rightarrow 2}^{\circ}$  which can be rationalised in terms of the steric effects of X and the endo-anomeric effects of Y.

