GRAPHICAL ABSTRACTS

DIELS-ALDER REACTION OF α -SUBSTITUTED ACRYLATES AND α -(METHYLENE)LACTONES: CONFORMATION OF DIENOPHILE AND ENDO/EXO SELECTIVITY

Tetrahedron, 1994, 50, 10839

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Enzymatic Resolution of 2-Acyl-3-hydroxymethyl-4-butanolide and Preparation of Optically Active IM-2, the Autoregulator from Streptomyces sp. FRI-5

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Racemic 2-acyl-3-hydroxymethyl-4-butanolide were resolved with lipases. The absolute configuration of IM-2 was deduced to be (2R, 3R, 6R).

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Novel Picrotoxane Norditerpene Lactones from Picrodendron baccatum

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Three novel picrotoxane terpenoids, picrodendrins U (1), V (2) and W(3), have been isolated from *Picrodendron baccatum*. Their structures were determined by spectral, X-ray analysis and CD spectrum.

SYNTHESIS AND REARRANGEMENT OF DISPIRO[3.1.3.2]-, DISPIRO[3.0.3.3]- AND DISPIRO[3.0.4.2]UNDECANES - NEW ENTRIES TO [3.3.3]PROPELLANES

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Dispiroundecanes 1-3 and 5-7 have been synthesized and rearranged to [3.3.3] propellanes 4 and 8, respectively. Some consequences concerning the synthesis of naturally occurring triquinanes are discussed.

SYNTHESIS AND REARRANGEMENT OF DISPIRO[2.0.3.4]-, DISPIRO[3.0.3.3]- AND DISPIRO[2.1.3.3]UNDECANES - PREFERRED C_4 - C_5 OVER C_5 - C_4 AND C_4 - C_5 OVER C_5 - C_6 REARRANGEMENTS

Tetrahedron, 1994, 50, 10879

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Dispiroundecanes 1-3 and 5-7 have been synthesized, and 2 and 5 rearranged to [3.3.3] propellanes 4 and 8, respectively. With 6 and 7, an initial C_4 - C_5 ring enlargement and C_4 - C_3 ring contraction, respectively, prevents a formation of 8.

INDOLOQUINONES, PART 3. PALLADIUM-PROMOTED

Tetrahedron, 1994, 50, 10893

SYNTHESIS OF HYDROXY-SUBSTITUTED 5-CYANO-5H-BENZOIb|CARBAZOLE-6.11-DIONES

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The synthesis of hydroxy-substituted 5-cyano-5*H*-benzo[*b*]carbazole-6,11-diones has been achieved by the following four-step sequence: addition of an arylamine to a naphthoquinone, palladium(II)-promoted oxidative cyclization, *N*-cyanation with cyanogen bromide, and chemoselective ether cleavage using pyridine hydrochloride.

ASYMMETRIC SYNTHESIS OF B-AMINOTETRALINS BY ELECTROPHILIC AMINATION

Tetrahedron, 1994, 50, 10909

Peter Gmeiner* and Bernd Bollinger; Pharmazeutisches Institut der Universität, An der Immenburg 4, 53121 Bonn, Germany

An asymmetric synthesis of 6 starting

O NR* CO₂R'

An asymmetric synthesis of 6 starting from 1a,b is reported when chiral amino ethers were used as auxiliaries. The protected hydrazino imines 5, readily obtained via electrophilic amination served as the key intermediates.

Partial Synthesis of Nitrogenous Brassinosteroid Analogues with Solanidane Skeleton

Tetrahedron, 1994, 50, 10923

Le thi Quyen, Günter Adam, and Klaus Schreiber Institute of Plant Biochemistry, Weinberg 3, D-06120 Halle/Saale, Germany

The nitrogenous brassinosteroid analogues 2α , 3α -dihydroxy- 5α , $22\alpha H$, 25BH-solanidan-6-one (2) and 2α , 3α -dihydroxy-6,7-seco- 5α , $22\alpha H$, 25BH-solanidano-6,7-lactone (3) have been synthesized from the *Solanum* steroid alkaloid solanidine (1).

Tetrahedron, 1994, 50, 10933

KINETICS AND MECHANISM OF OXIDATION

OF SOME SUBSTITUTED TRANS-CINNAMIC ACIDS BY

PYRIDINIUM CHLOROCHROMATE --- A NON-LINEAR HAMMETT PLOT

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The order with respect to [PCC] and [oxalic acid] in one each. The order in [H⁺] and [Substrate] is fraction Both electron-releasing and withdrawing substituents facilitate the Oxidation rate.

CH -CH = CH-CO H + 2C H NHCro CI + (CO H), ----> C H CHO + OHCCO H + 2CO + 2 C H NHCI + H O + Cr O

KINETICS AND MECHANISM OF Os(VIII)-CATALYSED

Tetrahedron, 1994, 50, 10945

OXIDATION OF SOME SUBSTITUTED TRANS-CINNAMIC ACIDS

BY CHLORAMINE-T IN ALKALINE MEDIUM — A NON-LINEAR HAMMETT PLOT

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Department of Chemistry, Annamalai University, Annamalainagar-608 002 (INDIA)

The reaction is first order in [CAT] and [OsO] and inverse first order in [OH]. A plot of logk verses σ gives a 'V' shaped curve.

$$C_{65}^{H}$$
-CH=CH-CO₂H + 2CH₂C₁H₂-SO₂-NCiNa + 2H₂O₂O₂O₃H₂ - > C₁H₂CHO + OHCCO₂H + 2CH₂C₁H₂-SO₂NH₂ + 2NaCl

AN EASY LEWIS ACID-MEDIATED ISOMERIZATION FROM (E)- TO (Z)-OXOINDOLIN-3-YLIDENE KETONES.

Tetrahedron, 1994, 50, 10955

G. Faita, M. Mella, P.P. Righetti*, and G. Tacconi.

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

(E)-2-oxoindolin-3-ylidene ketones can be easily isomerized to their (Z)-isomers by AlCl₃ at r.t. in CH₂Cl₂. The behaviour of the unsaturated dicarbonyl framework as a bidentate ligand can be the key-step of the isomerization.

PHENALENONE-TYPE PHYTOALEXINS FROM MUSA ACUMINATA. SYNTHESIS OF 4-PHENYL-PHENALENONES

Tetrahedron, 1994, 50, 10963

J. G. Luis', W. Q. Fletcher', F. Echeverri', T. A. Grillo'.

 C.P.N.O. "Antonio González", Instituto de Bio-Orgánica, Universidad de La Laguna, Carretera de La Esperanza 2, La Laguna, 38206, Tenerife, Canary Islands, Spain; Universidad de Antioquia, Medellín, Colombia.

The structure of two new phenalenone-type phytoalexins from Musa acuminata, elicited by the fungus Fusarium oxysporum, were established. The synthesis of these compounds, unknown until now either as natural or synthetic substances was acomplished by a simple and direct route which constitutes the first synthetic approach to 2-hydroxy-4-phenyl-phenalen-1-ones.

Preparation of a 3-Phenyl-4(3H)-isoquinolinone and its Transformation in 12(11H)-Benzo[c]phenanthridinone Derivatives. Crystal Structure Determinations.

Tetrahedron, 1994, 50, 10971

Ana M. González-Cameno. Dolores Badía. Esther Domínguez. M.Karmele Urtiaga, M. Isabel Arriortua. and Xavier Solans ¹Dpt. Química Orgánica and ²Dpt.Mineralogía-Petrología, Facultad de Ciencias, Universidad del País Vasco, P.O.Box 644-48080 Bilbao. Spain. 3 Dpt. Cristalografía, Mineralogía y Depósitos Minerales, Universidad de Barcelona, Martí i Franqués s/n, 08028 Barcelona. Spain

A procedure for the preparation of 12(11H)benzo[c]phenanthridinone derivatives involving the initial formation of 3-phenyl-4(3H)-isoquinolinone is described. The structures of these compounds are established by X-ray diffraction analyses

Reactions of Ninhydrin with Activated Anilines: Formation of Indole

Tetrahedron, 1994, 50, 10983

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In benzene, ninhydrin undergoes electrophilic substitution at C2 of 3,5-dimethoxyaniline leading to the indeno[1,2-b]indole (7): the corresponding reaction in water undergoes electrophilic substitution at C4 to give compound (11).

6 R,R1=CH2

CHEMISTRY OF ZAMORANIC ACID. PART V. HOMOCHIRAL SEMISYNTHESES OF ACTIVE DRIMANES: PERENIPORIN B.

Tetrahedron, 1994, 50, 10995

Tetrahedron, 1994, 50, 11013

POLYGODIAL AND WARBURGANAL. Julio G. Urones*, Isidro S. Marcos, Belén Gómez Pérez, David Díez, Anna M. Lithgow, Patricio M. Gómez, Pilar Basabe and N. M. Garrido. Dpto. Química Orgánica, Pza. de los Caídos 1-5, 37008, Salamanca, Spain

3 R=OH

The key intermediate for the preparation of pereniporin B 4, polygodial 2 and warburganal 3 is the epoxide 22 that was obtained from diene 6 by chemoselective epoxidation. The latter was prepared from zamoranic acid methyl ester 1.

SYNTHESIS OF [1-[3',5'-BIS-O-(t-BUTYLDIMETHYLSILYL) -B-D-ARABINO- AND B-D-RIBOFURANOSYLICYTOSINE1-2'-SPIRO-5"-(4"-AMINO-1",2"-OXATHIOLE-2",2"-DIOXIDE). ANALOGUES OF THE HIGHLY SPECIFIC ANTI-HIV-1 AGENT TSAO-T.

Sonsoles Velázquez, María-Luisa Jimeno, Jan Balzarini and María-José Camarasa

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PERFLUOROACYLATION OF ALKENES

Tetrahedron, 1994, 50, 11023

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$$\begin{array}{c|cccc}
R_1 & & & R_1 \\
\hline
 & CH_2 & & (CF_3CO)_2O \\
\hline
 & S(CH_3)_2, BF_3 & R_2
\end{array}$$
CHCOCF₃

SYNTHESIS AND TAUTOMERISM OF 9-AZABICYCLO-[42.1]NONAN-1-OLS (NORHOMOTROPAN-1-OLS), N-ALKYL AND 7,8-DEHYDRO- DERIVATIVES, AND OXABICYCLIC ANALOGUES

Craig R. Smith, David E. Justice and John R. Malpass, Department of Chemistry, University of Leicester, Leicester LE1 7RH, U.K.

Routes to the title compounds are described and the ratios of monocyclic and bicyclic tautomers are determined using NMR spectroscopy

Tetrahedron, 1994, 50, 11039

HNY

Y = COPh, CO2CH2Ph

X = NCH₃, NH, NCH₂Ph, NCOPh, NCO₂CH₂Ph

Chiral Acetals in Organic Synthesis: Regioselective Synthesis of 2- and 3-Hydroxy Acetals from

Tetrahedron, 1994, 50, 11057

2,3-Olefinic Acetals. Reinvestigation and Further Applications. Yashwant D. Vankar, M. Venkatram Reddy and Narayan C. Chaudhuri

Department of Chemistry, Indian Institute of Technology, Kanpur-208016, India Reactions of chiral and achiral 2,3-olefinic acetals have been described.