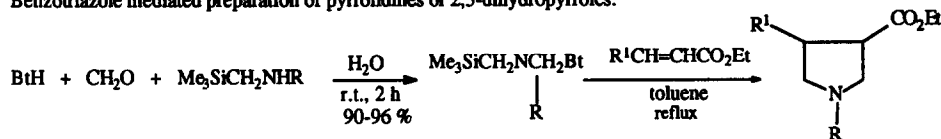


GRAPHICAL ABSTRACTS

BENZOTRIAZOLYLMETHYLAMINOSILANES: NOVEL AZOMETHINE YLIDE EQUIVALENTS

Alan R. Katritzky,* Jens Köditz and Hengyuan Lang
Center for Heterocyclic Compounds, Department of Chemistry,
University of Florida, FL 32611-7200

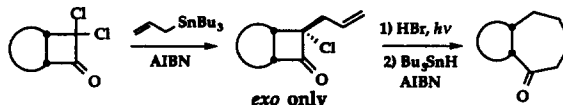
Benzotriazole mediated preparation of pyrrolidines or 2,5-dihydropyrroles.



Tetrahedron, 1994, 50, 12571

STEREOSELECTIVE ALLYLATION OF DICHLOROCYCLOBUTANONES AND SEQUENTIAL RING EXPANSIONS: CONSTRUCTION OF CIS-FUSED CYCLOHEPTANONES

Wei Zhang, Ye Hua, Steven J. Geib,
Garrett Hoge and Paul Dowd*
Department of Chemistry, University of
Pittsburgh, Pittsburgh, PA 15260, USA



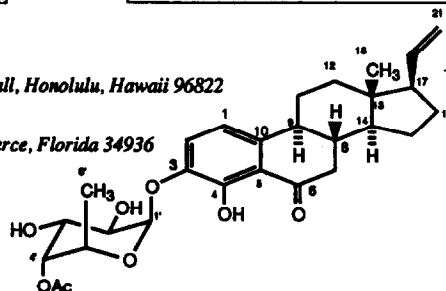
Stereoselective annulation of cyclic alkenes for the preparation of *cis*-fused seven-membered ring systems was accomplished by sequential [2 + 2] cycloaddition, *exo*-allylation, hydrohalogenation, and free radical ring expansion.

Tetrahedron, 1994, 50, 12579

HAPAIOSIDE: A 19-NORPREGNANE GLYCOSIDE FROM THE SPONGE *CRIBROCHALINA OLEMDA*.

Bryan K. S. Yeung, Mark T. Hamann, and Paul J. Scheuer*
Department of Chemistry, University of Hawaii at Manoa, 2545 The Mall, Honolulu, Hawaii 96822
Michelle Kelly - Borges
Harbor Branch Oceanographic Institution, 5600 North U. S. 1, Fort Pierce, Florida 34936

A 4-hydroxy-6-oxonorpregnane-3-glycoside with an aromatic ring A was isolated from a Pohnpei sponge, *Cribrochalina olemda*. The sugar is a 6'-deoxy-L-β-altropyranose-4'-acetate

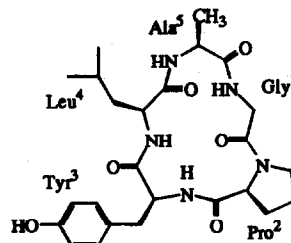


Tetrahedron, 1994, 50, 12593

Conformational Analysis of a Tyrosinase Inhibitory Cyclic Pentapeptide, Pseudostellarin A, from *Pseudostellaria heterophylla*

Hiroshi Morita, Takashi Kayashita,
Koichi Takeya and Hideji Itokawa*

Department of Pharmacognosy, Tokyo College of
Pharmacy, Horinouchi 1432-1, Hachioji, Tokyo
192-03, Japan,



Pseudostellarin A (1)

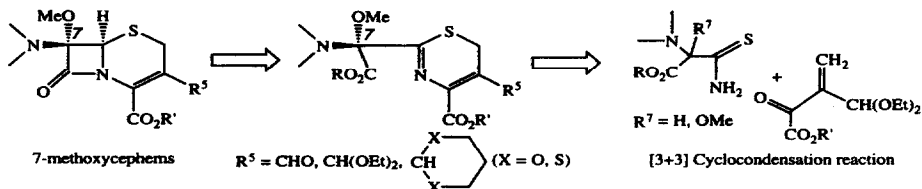
Tetrahedron, 1994, 50, 12599

α -Methoxy- α -(6H-1,3-thiazin-2-yl)glycinates precursors of 7-methoxycephems by [3+3] cyclocondensation reaction

Tetrahedron, 1994, 50, 12609

Abdelhadi Boussoufi, Jean-Luc Parrain, Pierrick Hudhomme and Guy Duguy*. *Laboratoire de Synthèse Organique, URA CNRS n°475, Faculté des Sciences et des Techniques, 2, rue de la Houssinière, 44072 Nantes Cedex 03, France.*

The synthesis of functionalised 6H-1,3-thiazin cycloadducts, versatile key intermediates in the total multistep syntheses of 7-methoxycephems, as analogues of 7-methoxycephalosporins, was described using the [3+3] cyclocondensation reaction.



Reaction of P-Phenyl C-Aminophosphaalkenes with p-Chlorobenzonitrile Oxides. Thermal behaviour of the adducts

Tetrahedron, 1994, 50, 12625

Mustapha Rahmouni ¹, Yeung Yat Cheng Yeung Lam Ko ¹, Robert Carrié ^{1*} et François Tonnard ²

1) Groupe de Physicochimie Structurale, U.R.A. C.N.R.S. n° 704, Université de Rennes I, Avenue du Général Leclerc, F 35042 RENNES Cedex.

2) Groupe Matière Condensée et Matériaux, U.R.A. C.N.R.S. n° 804, Université de Rennes I, Avenue du Général Leclerc, F 35042 RENNES Cedex.

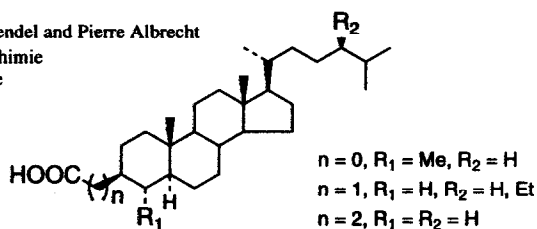
At -20°C, the title reaction leads to 1,2,3-oxazaphospholines. Thermal behaviour of these adducts is studied (experimental and theoretical approach).

CHARACTERISATION OF NOVEL 3-CARBOXYALKYL-STERANES OCCURRING IN GEOLOGICAL SAMPLES

Tetrahedron, 1994, 50, 12633

Philippe Schaeffer, Fabienne Fache-Dany, Sylvie Trifilieff, Jean M. Trendel and Pierre Albrecht
Laboratoire de Géochimie Organique, URA 31 du CNRS, Faculté de Chimie
Université Louis Pasteur, 1 rue Blaise Pascal, 67000 Strasbourg, France

Three novel series of 3-carboxyalkyl-steranes have been identified in sediments by synthesis. They may derive from yet unknown microbial precursors.



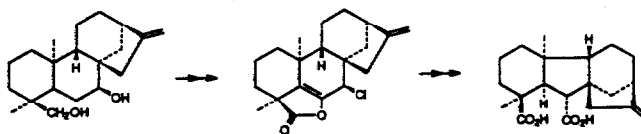
THE TRANSFORMATION OF EPICANDICANDIOL INTO A GIBBERELLIN A₁₂ ISOMER

Tetrahedron, 1994, 50, 12643

Braulio M. Fraga*, Melchor G. Hernández, Jacinto D. Arraez, and Javier G. Luis

Instituto de Productos Naturales y Agrobiología, CSIC, and Instituto Universitario de Bio-Organica, Univ. La Laguna, Avda. Astrofisico F. Sanchez 2, 38203-La Laguna, Tenerife, Spain

A gibberellin A₁₂ isomer has been obtained by Favorskii rearrangement of a chloro-enol-lactone, prepared from epicandicandiol.

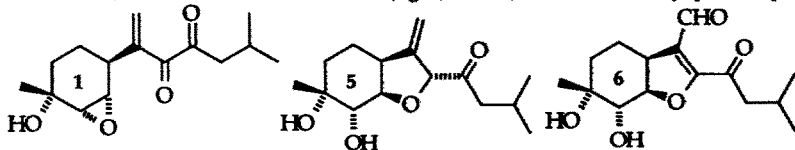


SIX NEW ANTIMICROBIAL AND NEMATICIDAL BISABOLANES FROM THE BASIDIOMYCETE CHEIMONOPHYLLUM CANDIDISSIMUM

Tetrahedron, 1994, 50, 12649

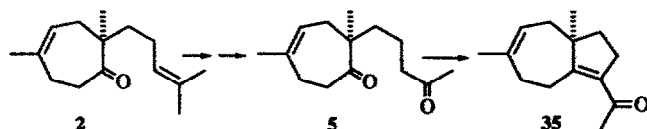
Marc Stadler, Heidrun Anke and Olov Sterner, Universities of Kaiserslautern (FRG) and Lund (Sweden)

The structures of six new bioactive bisabolanes (e.g. 1, 5 and 6) were determined by spectroscopic methods.



SYNTHESIS OF A 1-ACETYL-3 α ,6-DIMETHYL-HEXAHYDROAZULENE. VERSATILE INTERMEDIATE FOR THE PREPARATION OF TERPENOIDS WITH BICYCLE[5.3.0]DECANE SYSTEM. Isidro S. Marcos, Isabel M. Oliva, Rosalina F. Moro, David Díez and Julio G. Urones*
Dpto. Química Orgánica, Pza. de los Caídos 1-5, 37008, Salamanca, Spain

Tetrahedron, 1994, 50, 12655



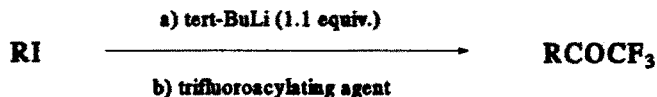
The transformation of cycloheptenone 2, synthesised from *Z/E* (\pm) nerolidol, into enone 35 through the dione 5 has been setted up in 7 steps with an overall yield of 53%. The enone 35 is a versatile precursor for the preparation of terpenoids either sesquiterpenes or diterpenes with a bicycle[5.3.0]decane system.

AN EFFICIENT AND EXPEDITIOUS SYNTHESIS OF FUNCTIONALIZED TRIFLUOROMETHYL KETONES THROUGH LITHIUM-IODINE EXCHANGE REACTION

Tetrahedron, 1994, 50, 12673

Isabel Villuendas, Alfredo Parrilla and Angel Guerrero*

Department of Biological Organic Chemistry, C.I.D., C.S.I.C., Jordi Girona 18-26. 08034 Barcelona, Spain

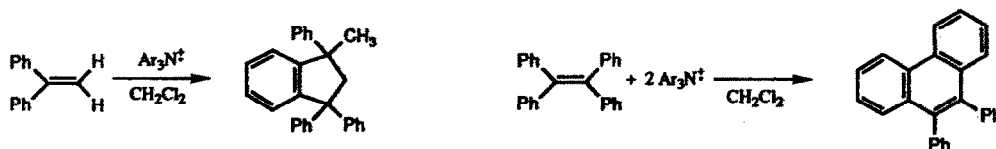


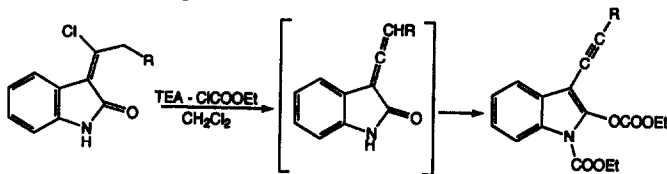
REACTIONS ON AROMATIC OLEFINS INDUCED BY AMINIUM SALTS: PROTIC-ACID OR RADICAL-CATION CATALYZED PROCESSES

Tetrahedron, 1994, 50, 12685

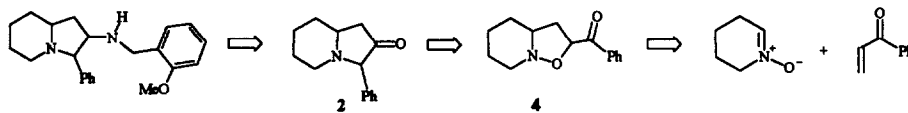
Francesco Ciminale, Luigi Lopez,* Giuseppe Mele

Centro C.N.R. "M.I.S.O.", Dipartimento di Chimica, Università di Bari, Via Amendola 173, Bari, Italy 70126

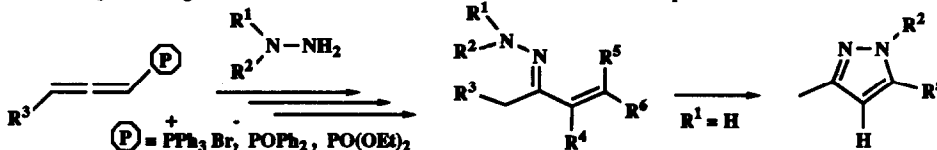


2-ETHOXYCARBOXYLOXY-3-ETHYNYLINDOLES FROM INDOL-2(3H)-ONES*Tetrahedron, 1994, 50, 12697*E.M. Beccalli,^a A. Marchesini,^a T. Pilati^b^aIstituto di Chimica Organica, Facolta' di Farmacia, Universita' degli Studi di Milano, via Venezian 21 - 20133 Milano - Italy ^bCNR Centro Studio delle Relazioni tra Struttura e Reattivita' Chimica, via Golgi 19 - 20133 Milano - Italy**SYNTHESES OF 3-PHENYL SUBSTITUTED INDOLIZIDIN-2-ONES AND A PYRROLIZIDIN-2-ONE ON THE ROUTE TO CONSTRAINED POTENTIAL NK₁ RECEPTOR ANTAGONIST***Tetrahedron, 1994, 50, 12713*Franca M. Cordero,^a Alberto Brandi,^{a*} Simone Cristini,^a Francesco De Sarlo,^a Giovanni Viti^b^aDipartimento di Chimica Organica "U. Schiff" and Centro CNR dei Composti Eterociclici, Università di Firenze, Via G. Capponi 9, I-50121 Firenze (Italy);^bMenarini Industrie Farmaceutiche Riunite S.r.l., Via Sette Santi 3, I-50131 Firenze (Italy)

New indolizidine NK₁ receptor antagonists have been synthesized starting from 3-phenyl-indolizidin-2-one (2), obtained from a benzoylisoxazolidine 4 by reductive ring cleavage followed by intramolecular condensation.

**AN EFFECTIVE STRATEGY FOR THE PREPARATION OF α,β -UNSATURATED HYDRAZONES AND PYRAZOLE DERIVATIVES. SYNTHETIC APPLICATIONS OF β -FUNCTIONALIZED PHOSPHORUS COMPOUNDS.***Tetrahedron, 1994, 50, 12727*Francisco Palacios^a, Domitila Aparicio, Jesús M. de los Santos.

Departamento de Química Orgánica, Facultad de Farmacia, Universidad del País Vasco, Apartado 450, 01006 Vitoria, SPAIN.

**Preparation of Phosphono Peptides Containing Phosphoramidate Bond***Tetrahedron, 1994, 50, 12743*

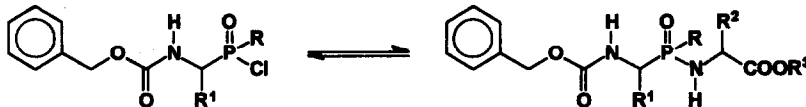
Artur Mucha, Pawel Kafarski

Institute of Organic Chemistry, Biochemistry and Biotechnology, Technical University of Wrocław, Poland

Francoise Plenat, Henri-Jean Cristau

Laboratoire de Chimie Organique, Ecole Nationale Supérieure de Chimie, Montpellier, France

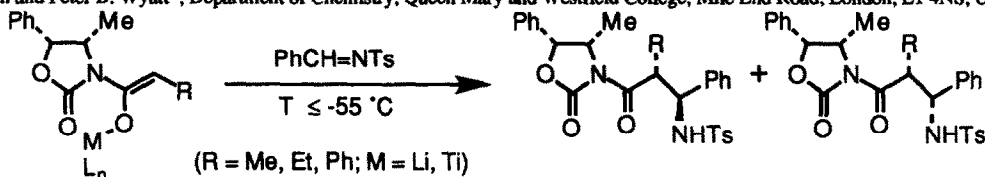
Factors influencing the formation of phosphoramidate bond during phosphono peptide synthesis were studied.



ADDITION TO ACTIVATED IMINES OF ENOLATES FROM CHIRAL N-ACYLOXAZOLIDINONES Isaac Abrahams, Majid Motevalli, Andrew J.

Robinson and Peter B. Wyatt*, Department of Chemistry, Queen Mary and Westfield College, Mile End Road, London, E1 4NS, UK.

Tetrahedron, 1994, 50, 12755



Typically only two out of the four possible addition products were formed, implying that reaction is through chelated (Z)-enolates.

ELECTROCHEMICAL OXIDATION OF DIHYDROINDOLO-QUINAZOLINES AND DIHYDROQUINAZOLINES

James Y. Becker* and Elias Shakkour, Department of Chemistry, Ben-Gurion University of the Negev, Beer Sheva 84120, Israel

Tetrahedron, 1994, 50, 12773

Abstract: Cyclic voltammetry of five tetracyclic dihydroindoloquinazolines (1) and five bicyclic dihydroquinazolines (2) have been studied in dichloromethane. Anodic oxidation of type 1 derivatives yields mostly benzophenone and quinazolinone 4 (the X-ray crystal structure of 4c has been determined).

