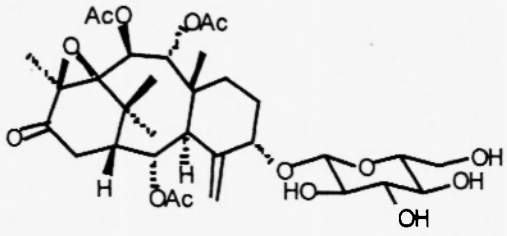
Graphical Abstracts

Heterocycl. Commun. 11 (2005) 211 - 214

First example of 11,12-epoxytaxane-glucoside from the needles of taxus cuspidata

Chun-Lin Wang[1], *Man-Li Zhang*[1], *Cong-Mei Cao*[1], *Qing-Wen Shi**[1], *Hiromasa Kiyota**[2]
[1]School of Pharmaceutical Sciences,
Hebei Medicinal University, China
[2]Graduate School of Agricultural Science, Tohoku University, Japan



A new 11,12-epoxytaxane-glucoside, 2α,9α,10β-triacetoxy-11, 12-epoxytax-4 (20)-en-13-one-5α-*O*-β-D-glucopyranoside was isolated from the needles of *Taxus cuspidata*.

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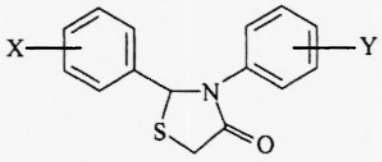
A preliminary study on predicting the ^{13}C chemical shifts for a series of disubstituted 2,3-diphenyl-1,3-thiazolidin-4-ones

John Tierney,[a] *Douglas Sheridan,*[a] *Linda Mascavage,*[b] *Daniela Gorbecheva,*[b] *Michelle Ripp*[b] *and Sonjoo Son*[b]
[a] Department of Chemistry, Pennsylvania State University, Delaware County Campus, Media, Pennsylvania 19063, USA
[b] Department of Chemistry, Arcadia University, Glenside, Pennsylvania, 19038, USA



Series 1: X = *p*-NO_2, *m*-NO_2, *p*-F, *m*-F, *p*-Cl, *p*-Br, *m*-Br, H, *p*-CH_3, *m*-CH_3, *p*-OCH_3, *m*-OCH_3; Y = H

Series 2: Y = *p*-NO_2, *m*-NO_2, *p*-F, *m*-F, *p*-Cl, *p*-Br, *m*-Br, H, *p*-CH_3, *m*-CH_3, *p*-OCH_3, *m*-OCH_3; X = H

Series 3: X = Y = *p*-NO_2, *m*-NO_2, *p*-F, *m*-F, *p*-Cl, *p*-Br, *m*-Br, *p*-CH_3, *m*-CH_3, *p*-OCH_3, *m*-OCH_3

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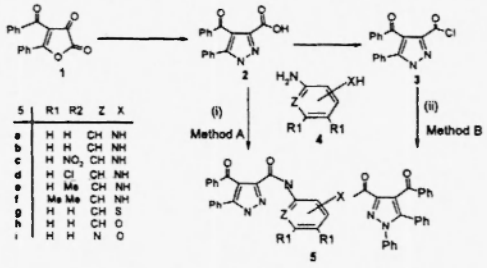
Synthesis and theoretical calculations of the 1*h*-pyrazole-3-carboxamide and -3-carboxylate derivatives

İsmail Yıldırım[1] *and Fatma Kandemirli**[2]
[1]Department of Chemistry, Erciyes University, TR -38039, Kayseri, Turkey
[2]*Department of Chemistry, Kocaeli University, TR -41300, Izmit, Turkey



5	R1	R2	Z	X
a	H	H	CH	NH
b	H	H	CH	NH
c	H	NO_2	CH	NH
d	H	Cl	CH	NH
e	H	Me	CH	NH
f	Me	Me	CH	NH
g	H	H	CH	S
h	H	H	CH	O
i	H	H	N	O

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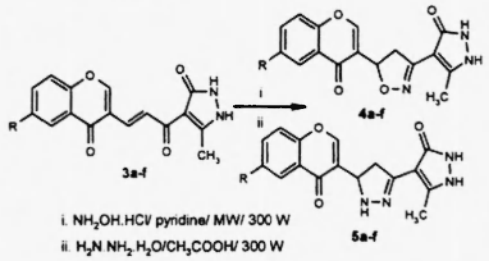
Synthesis of new hetero aroyl chromen-4-ones

*P Narasimha Reddy, Y Thirupathi Reddy, V Naveen Kumar & B Rajitha**

Synthesis of 5-methyl-4-[3-(4-oxo-4*H*-chromene-3-yl)-acryloyl)-1,2-dihydro-pyrazol-3-ones **3a-f**, synthesis of 5-methyl-4-[5-(4-oxo-4*H*-chromene-3-yl)-4,5-dihydro-isoxazol-3-yl]-1,2-dihydro-pyrazol-3-ones **4a-f** and 5′-methyl-5-(4-oxo-4*H*-chromen-3-yl)-4,5,1′,2′-tetrahydro-1*H*-[3,4′]bipyrazole-3′-ones **5a-f** in good yields is described under microwave irradiation conditions.



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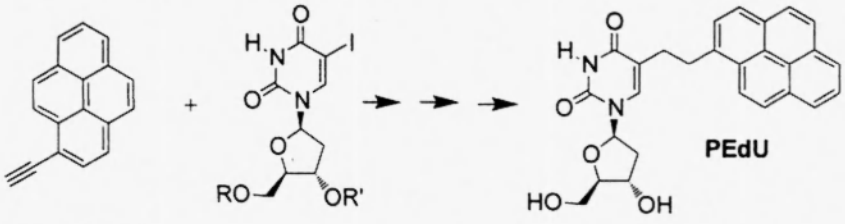
Synthesis of 5-(2-pyren-1-yl-ethylenyl)-2'-du as a fluorescent probe for studying electron transfer in DNA

*Samir T. Gaballah[1] and Thomas L. Netzel**
Department of Chemistry, Georgia State University, P.O. Box 4098, Atlanta, GA 30302-4098
[1]Current Address: Department of Photochemistry, National Research Center, Dokki, Cairo 12311, Egypt



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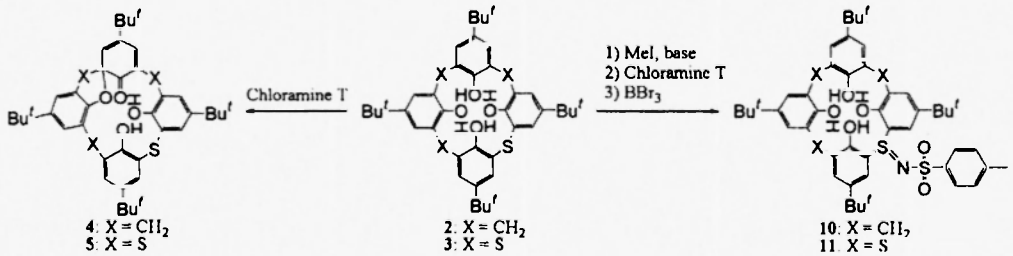
Conversion of mono- and tetra-thiacalix[4]arenes to sulfilimine derivatives and unexpected formation of monospirodienone derivatives

*Naoya Morohashi, Makoto Kojima, Akihiro Suzuki, Yoshihiro Ohba**
Department of Chemistry and Chemical Engineering, Faculty of Engineering, Yamagata University, Yonezawa 992-8510, Japan



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Synthesis, reactions and antimicrobial activity of some new 1,3,4-oxadiazoles, 1,2,4-triazoles and 1,3,4-thiadiazines derived from pyrazole.

A. A. Farghaly[a], P. Vanelle[b] and H. S. El-Kashef[a]
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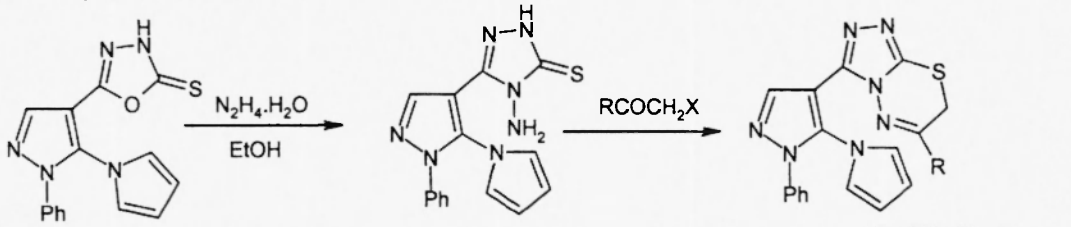
[b] Laboratoire de Chimie Organique Pharmaceutique LCOP-UMR CNRS 6517, Faculte de Pharmacie, 27 Boulevard Jean Moulin, 13385 Marseille Cedex 5, France, <patrice.vanelle@pharmacie.univ-mrs.fr>

The interaction of the oxadiazole **8** with hydrazine hydrate afforded the aminotriazolethione compound **13** which could be cyclized into the thiadiazine derivatives **14a-c**.



8 **13** **14a-c** a, R = CH_3, X = Cl; b, R = C_6H_5, X = Br; c, R = C_6H_4Br-*p*, X = Br

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Synthesis and antibacterial activities of fused pyranoquinoline derivatives

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P.O. Box 16222, Saudi Arabia

Ethyl 2-amino-4-aryl-6-chloro-4-H-pyrano[3,2-h]quinoline-3-carboxylate **1a-e** was converted into ethyl 2-(1-pyrrolyl)-4-aryl-6-chloro-4H-pyrano[3,2-h]quinoline-3-carboxylate **2a-e**. Several derivatives of the latter compound have been synthesized. Also, the synthesis of 7-aryl-5-chloropyrrolo[1",2":1',2']pyrazino[5,6:5',6']pyrano-[3,2-h]quinoline and other related heterocycles are described.

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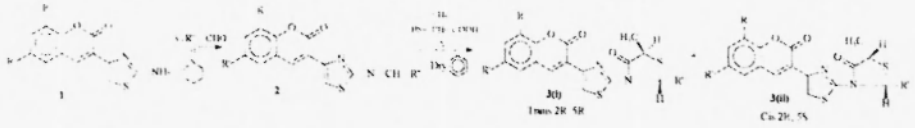
Synthesis and evaluation of anticancer and antiviral activity of some 2-aryl-3-(4-(2H-1-benzopyran-2-one-3-yl)-2-thiazolyl)-5-methyl-4-thiazolidinones

V. Rajeswar Rao, P. Vijaya Kumar, V. Ravinder Reddy and K.M. Reddy*
Department of Chemistry, National Institute of Technology,
Warangal - 506 004, India.



2-Aryl-3-(4-(2H-1-benzopyran-2-one-3-yl)-2-thiazolyl)-5-methyl-4-thiazolidinones (3), have been prepared by the reaction of Schiff's bases (2) with 2-mercaptopropionic acid. The Schiff's bases in turn are obtained by the reaction of various aldehydes with 3-(2-amino-4- thiazolyl)coumarins. The compounds 3 have been evaluated for their anticancer and antiviral activity.

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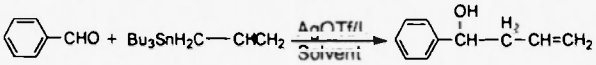
Chiral (S)-(+) 1-substituted aryl-4- (1-phenyl) ethylformamido-5-amino-1, 2, 3-triazole: a new class of chiral ligands for the silver(I)-promoted enantioselective allylation of aldehydes

Mindong Chen [a, c] *Shengli Guo* [b], *Youfei Zheng* [a], *Liang Chen* [a], *Feng Tang* [c], *Weiyi Hua* [c]
[a] Department of Environmental Sciences and Engineering, Nanjing University of Information Science & Technology; Nanjing, 210044
[b] Department of Physics, Nanjing University of Information Science & Technology; Nanjing, 210044
[c] College of Medicine, China Pharmaceutical University, Nanjing, 210009

Some novel chiral ligands (S)-(+) 1-substituted aryl-4- (1-phenyl) ethylformamido -5-amino -1, 2, 3-triazole were prepared starting from 1-substituted aryl-4- ethoxycarbonyl -5-amino-1, 2, 3- triazoles and other reagents. They were used as catalytic chiral ligands in the silver (I)-promoted enantioselective allylation reaction of aldehydes with allyltributyltin



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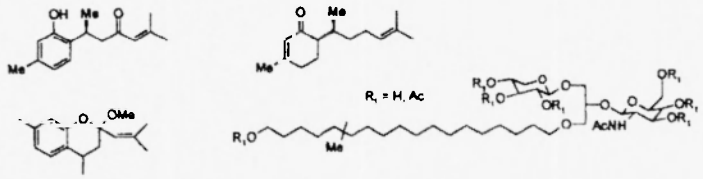
Isolation and characterization of new tetrahydropyranyl substituted sesquiterpene and myrmekiodermin glycolipid ether isolated from the marin e sponge *myrmekioderma*

Vyes Letourneux, Jean Michel Brunel, Rogelio Fernandez, Michel Dherbomez and Cecile Debitus*
*Laboratoire de Synthese et Etude de Substances Naturelles à Activités Biologiques (SESNAB), IMRN UMR INRA 1111, Case 342, Faculte de St Jerôme, Av. Escadrille Normandie Niemen, F-13397 Marseille, Cedex 20, France. e-mail : vyesletourneux@yahoo.fr



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Synthesis of some new 3-(2-oxo-2h-chromen-3-yl)-5h-[1,3]thiazolo[3,2-a] pyrimidine-5-ones

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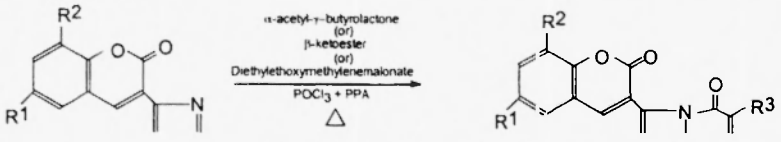
Department of Chemistry, National Institute of Technology, Warangal (A.P.) 506 004, India
e-mail: vrajesw@yahoo.com, vrajesw@nite.ernet.in



Condensation of 3-(2-amino-4-thiazolyl)coumarins (I) with α-acetyl-γ-butyrolactone in a mixture of polyphosphoric acid and $POCl_3$ affords 6-(2-chloromethyl)-7-methyl-3-(2-oxo-2H-chromen-3-yl)-5H-[1,3]-thiazolo[3,2-a] pyrimidine-5-one (IIa-f), while condensation of I with β-ketoesters gives 7-methyl-3-(2-oxo-2H-chromen-3-yl)-5H-[1,3]thiazolo [3,2-a]pyrimidine-5-ones (IIIa-g). Reaction between I and diethyl methoxy methylene malonate under solvent free conditions gives 3-(2-oxo-2H-chromen-3-yl)-5H-[1,3]thiazolo[3,2-a]pyrimidine-5-one-6-carboxylic acid ethyl ester (IV). The structures of newly synthesized compounds have been established by elemental analysis and spectral data.

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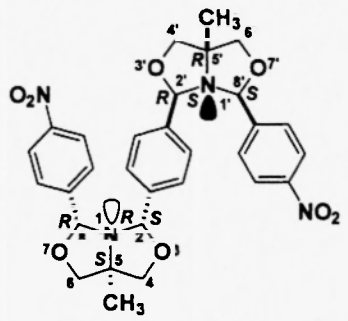
First example of long distance stereo controlled synthesis in 1-aza-3,7-dioxabicyclo[3.3.0.]octane series

Carmen Maiereanu,[a] Mircea Darabantu,[a] Eric Condamine,[b] Gérard Plé,[b] Yvan Ramondenc,[b] Marijana Fazekas,[a] Monica Pintea[a] and Camelia Berghian[c]

[a]"Babes-Bolyai" University, Department of Organic Chemistry, 11 Aranyi Jànos str., RO-3400 Cluj-Napoca, Romania
[b]Université de Rouen, Institut de Recherche en Chimie Organique Fine (I.R.C.O.F.),76131 Mont Saint-Aignan, France
[c]"Babes-Bolyai" University, Department of Inorganic Chemistry, 11 Aranyi Jànos str., RO-3400 Cluj-Napoca, Romania



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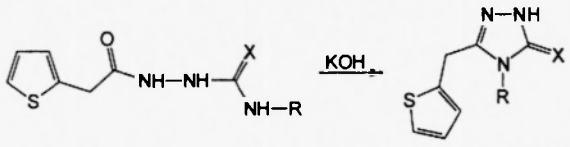
Organic sulfur compounds. vi. 5-(2-thienylmethyl)-1,2,4-triazole derivatives

Gheorghe Roman[a], Ileana Manciulea[a], Rodinel Ardeleanu[b], Lucia Dumitrescu[a]
[a] Department of Chemistry, Transilvania University, 29 Eroilor Blvd., Braşov, 500036, Romania
[b] P. Poni Institute for Macromolecular Chemistry, 41A Gr. Ghica Vodă Alley, Iaşi, 700487, Romania

New 1,2,4-triazole-3-thiones and 1,2,4-triazole-3-ones bearing a 2-thienylmethyl moiety in position 5 have been synthesized through the cyclization in the presence of aqueous KOH of the corresponding 1-(2-thienylacetyl)thiosemicarbazides and 1-(2-thienylacetyl)semicarbazides respectively. Potassium N'-(2-thienylacetyl)hydrazinecarbodithioate led to 4-amino-3-mercapto-5-(2-thienylmethyl)-1,2,4-triazole on treatment with hydrazine hydrate.



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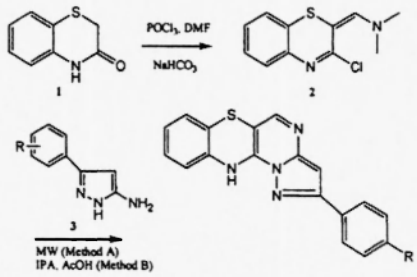
Reactions of 5-aminopyrazoles: Synthesis of pyrazolo [5′,1′: 2,3]pyrimido[5,4-*b*][1,4]-benzothiazines. a new tetracyclic ring system under microwave irradiation conditions

K. N. Jayaveera and S. Sailaja
Oil Technological Research Institute, Jawaharlal Nehru Technological University, Anantpur - 515001
and
*G. Jagath Reddy * and K. Srinivasa Rao*
R & D Laboratories, Dr. Jagath Reddy's Heterocyclics, 81, S.V.Co-op Industrial Estate,
Balanagar, Hyderabad – 500 037, India.
E-mail: jagathreddy@usa.net; Fax # 91-40-23773487.



A series of pyrazolo[5′,1′:2,3]pyrimido[5,4-*b*][1,4]-benzothiazines (**4a-f**), a new tetracyclic ring system have

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Reactions of acylglycines with heteroarylhydrazines

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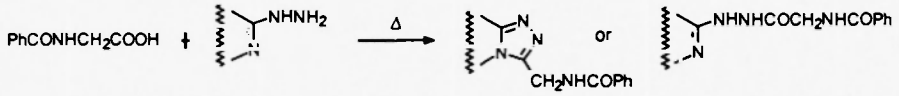
Faculty of Chemistry and Chemical Technology, University of Ljubljana, 1000 Ljubljana, Aškerčeva 5, Slovenia

Reactions of hippuric acid with heteroarylhydrazines leading to *N*-(heteroarylmethyl)benzamides or *N*-[2-oxo-2-(heteroarylhydrazino)ethyl]benzamides are described.



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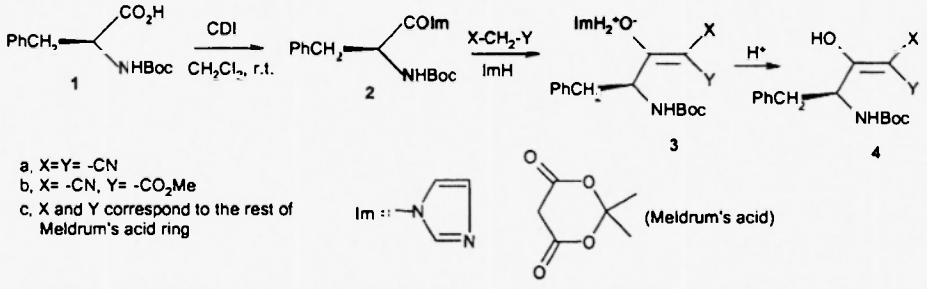
Synthesis of acylated active methylene compounds with *n*-boc-l-phenylalanine and their heterocyclization, both achieved enantioselectively

*Stylianos Hamilakis and Athanase Tsolomitis**
The Laboratory of Organic Chemistry, The School of Chemical Engineering,
The National Technical University of Athens, Athens 157 80, Greece
E-mail : tsolom@chemeng.ntua.gr

An overall enantioselective simple synthesis of acylated active methylene compounds with *N*-Boc-L-Phenylalanine and their transformation to heterocycles.



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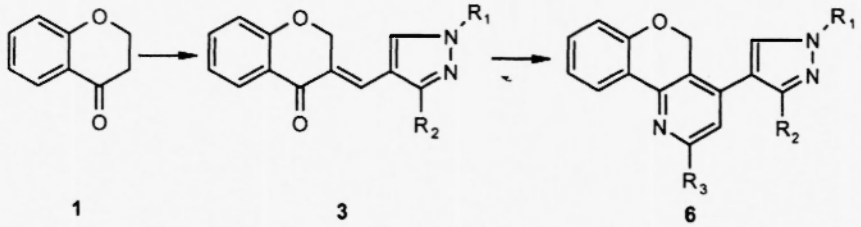
Synthesis of 2-Aryl-4-(1,3-diarylpyrazol-4-yl)-benzopyrano[4,3-*b*]pyridines

*G. Jagath Reddy * and K. Srinivasa Rao*
R & D Laboratories, Dr. Jagath Reddy's Heterocyclics, 81, S.V.Co-op Industrial Estate, Balanagar, Hyderabad -- 500 037, India. E-mail: jagathreddy@usa.net; Fax # 91-40-23773487.
and
Md. Khalilullah, D. Latha and C. Thirupathaiah
Department of Chemistry, Jawaharlal Nehru Technological University, Hyderabad - 500 072, India

A series of 2-Aryl-4-(1,3-diarylpyrazol-4-yl)benzopyrano [4,3-b]pyridines (**6a-h**) have been synthesized.



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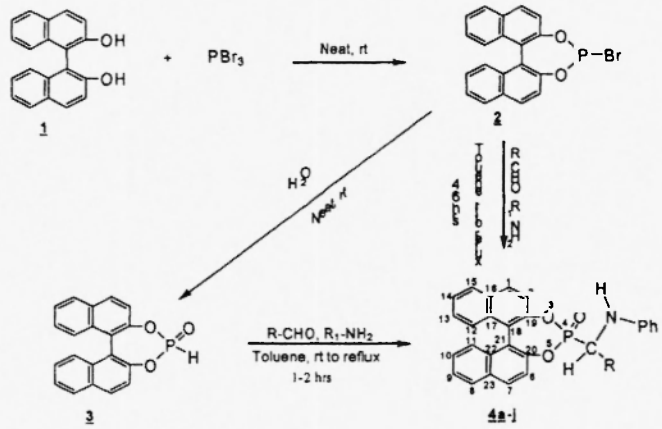
Synthesis and Antimicrobial activity of 2, 10 - dibromo dibenzo [*d,g*] [1,3,6,2] dioxathiaphosphocin 6-sulfido-6-amino acid esters

*P. Haranath, U. Anasuyamma, C.Devendranath Reddy and C. Suresh Reddy**
Department of Chemistry, Sri Venkateswara University, Tirupati, A.P., India.
E-mail : csureshsvu@yahoo.com

4-α-Arylamino benzyl dinaphtho [2,1-d:1',2'-f][1,3,2] dioxaphosphepin 4-oxides **4a-j**) were synthesized in excellent yield from three-component one pot reaction of aldehydes, anilines and dinaphtho [2,1-d:1',2'-f][1,3,2] dioxaphosphorobromodite **2** / corresponding hydrogen-phosphite **3** intermediates and characterized by IR, ^{1}H, ^{13}C and ^{31}P NMR spectral data and the title compounds were screened for their antimicrobial activity.



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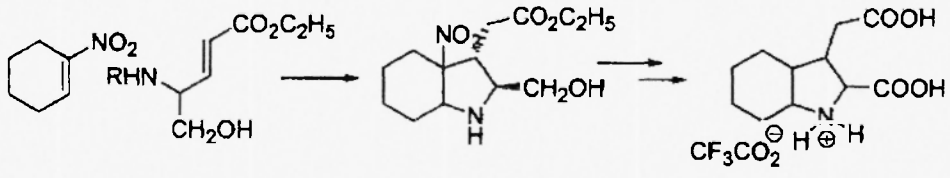
Synthesis of an analog perhydroindole of kanaic acid

*Matheieu Danel, Akram Hijazi, Roland Barret**
Laboratoire de Chimie Thérapeutique, EA 3741,
Faculté de pharmacie, Université Claude Bernard,
8 Avenue Rockefellr, F69373 Lyon Cedex 08



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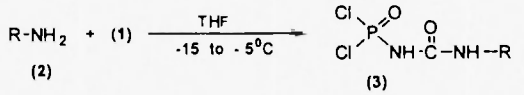
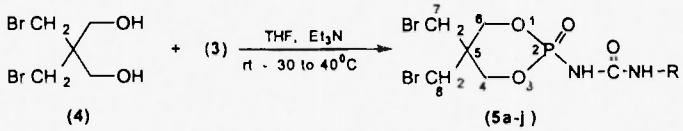
Synthesis of *N*-(substituted)-*N'*- [5,5'-bis(bromomethyl)-2-oxido-1,3,2-dioxaphosphorinane-2yl] ureas

*M.F. Stephen Babu, Y.B. Kiran, K.Ananda Kumar, C.Devendranath Reddy and C. Suresh Reddy**

The title compounds have been synthesized by condensation of equimolar quantities of chlorides of various carbamidophosphoric acids **3** with 2,2'-bis(bromomethyl)1,3-propanediol **4** in the presence of triethylamine in dry tetrahydrofuran at 30-40°C.





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Synthesis of new fluorescent β-cyclodextrin sensor

*Mathieu Becuwe,[a] François Delattre,[a] Georgiana G Surpateanu,[b] Francine Cazier,[a] Patrice Woisel,[a] Guillaume Garçon,[b] Pirouz Shirali,[b] Gheorghe Surpateanu[a]**

[a] Laboratoire de Synthèse Organique et Environnement, EA 2599, Université du Littoral Côte d'Opale,
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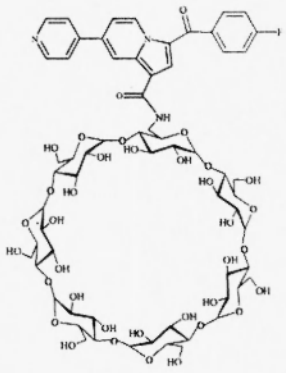
[b] Laboratoire de Recherche en Toxicologie Industrielle et Environnementale, Université du Littoral Cote d'Opale, 189A avenue Maurice Schumann, 59140 Dunkerque

Fax : (33) 03 28 23 76 05 ; e-mail : surpatea@univ-littoral.fr



The synthesis of a new fluorescent sensor incorporating a fluoro pyridine indolizinic unit and a β-cyclodextrin fragment by two different synthetic ways is described.

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Synthesis and spectral studies of some novel dialkyl/diaryl β-diketones and β-ketoesters from diazonium salt of 3n-propyl-4-amino-5-carboxamido-N-methyl pyrazole

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Various novel β-diketones and β-ketoesters (4a-4e) has been prepared by the condensation of diazonium salt of 3n-propyl-4-amino-5-carboxamido-N-methyl pyrazole with β-diketones and β-ketoesters (3a-3e).