

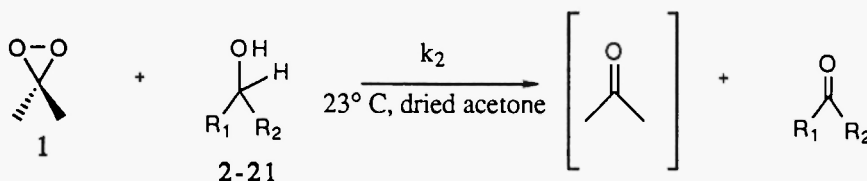
## Graphical Abstracts

Heterocycl. Commun. 4 (1998) 201-204

### OXIDATION OF SECONDARY ALIPHATIC ALCOHOLS BY DIMETHYLDIOXIRANE: KINETICS AND SELECTIVITY

Mark A. Cunningham, P.C. Vasquez, Paul J. Franklin and A.L. Baumstark,\* Department of Chemistry, Center of Biotechnology and Drug Design, Georgia State University, Atlanta, GA 30303-3083, USA

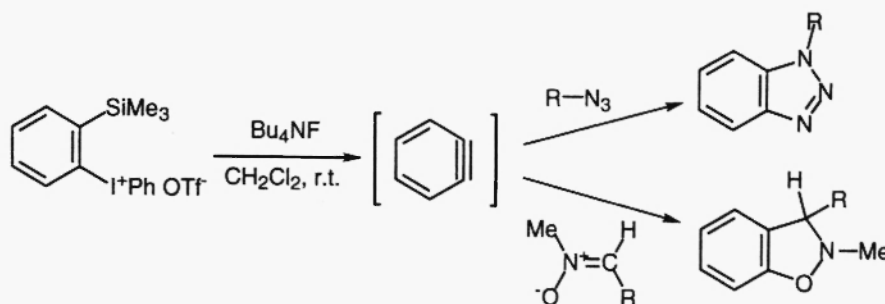
The oxidation of acyclic (2-10) and cyclic (11-21) secondary aliphatic alcohols by dimethyldioxirane in acetone at 23 °C produced the corresponding ketones in very good to excellent yields. Kinetic data ( $k_2$ 's) were found to be similar for the acyclic series except for the most hindered compounds. The  $k_2$ 's for the cyclic series were found to vary with ring size and to be generally larger than those for the acyclic cases.



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### A CONVENIENT AND EFFICIENT SYNTHESIS OF BENZOTRIAZOLES AND BENZISOXAZOLINES USING A NEW HYPERVALENT IODINE-BENZYNE PRECURSOR

Tsugio Kitamura,\* Mitsuru Todaka, Ichiro Shin-machi, and Yuzo Fujiwara  
Department of Chemistry and Biochemistry, Graduate School of Engineering, Kyushu University, Hakozaki, Fukuoka 812-8581, Japan

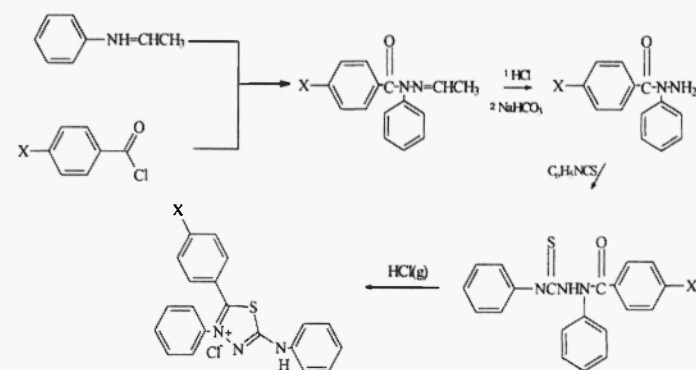


### On the synthesis of mesoionic 1,3,4-thiadiazolium-2-aminide and precursors

Marcelo M. Britto<sup>1</sup>, Carlos A. Montanari<sup>1\*</sup>, Claudio L. Donnici<sup>1</sup> and Quézia B. Cass<sup>2</sup>

<sup>1</sup>Núcleo de Estudos em Química Medicinal, Departamento de Química - Universidade Federal de Minas Gerais, Campus da Pampulha, 31270-901, Belo Horizonte, MG, Brazil

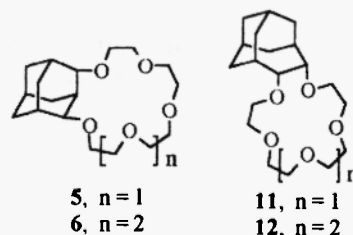
<sup>2</sup>Departamento de Química - Universidade Federal de São Carlos - São Carlos - SP - Brazil



### SYNTHESIS OF CROWN ETHERS EMBODIED ADAMANTANE AND HOMOADAMANTANE SKELETONS

Shoji Eguchi,\* Hideo Miyake, Archana Gupta, and Takashi Okano  
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Adamantano[2,4]-16-crown-5 and -19-crown-6 ethers (**5**, **6**), and homoadamantano[4,5]-15-crown-5 and -18-crown-6 ethers (**11**, **12**) have been prepared and their some cation binding properties have been discussed based on the solvent extraction and PM3 calculation results.

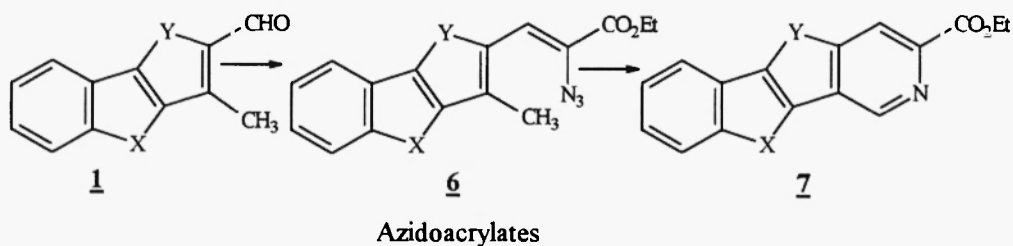


## New Heterocyclic analogues of pyridocarbazoles from azidoacrylates

Nachwa Jarkas<sup>1</sup>, Gilbert Kirsch<sup>\*a</sup> and Pierre Seck<sup>b</sup>

a) Laboratoire de Chimie Organique, Groupe de Synthèse Organique et Hétérocyclique, Université de Metz, Ile du Saulcy, 57045 Metz cedex 01, France

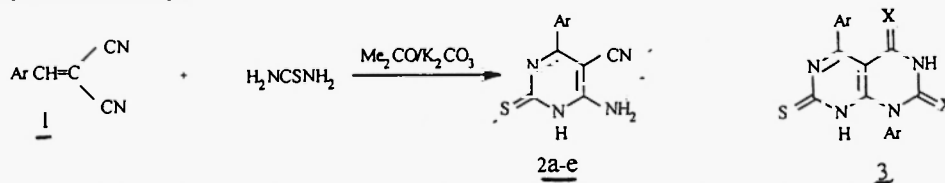
b) Centre Universitaire de Luxembourg, 162A Avenue de la Faïencerie, L-1511 Luxembourg

Analogues of isoellipticine **7** were synthesized starting from aldehydes **1** via azidoacrylates **6**.

## SYNTHESIS OF PYRIMIDO [4,5-d] PYRIMIDINETHIONE DERIVATIVES AS BIOCIDAL AGENTS

Mohamed Abdel - Megid

Department of Chemistry, Faculty of Education Ain-Shams University, Roxy, Cairo, Egypt.

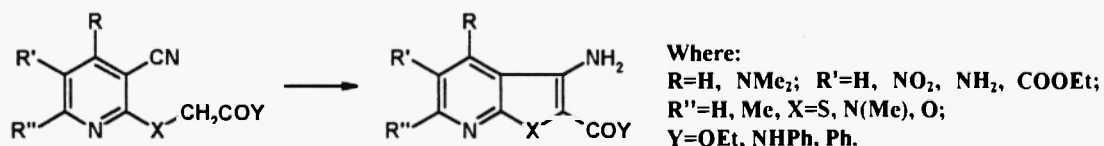
Several new pyrimido [4,5-d] pyrimidinethiones have been synthesized from  $\beta$ -arylidene malononitriles with thiourea followed by cyclocondensation with nitrogen and oxygen compounds. And they were tested for antibacterial in the comparison with antibiotics.

## THE INVESTIGATION OF PYRROLO-, THIENO- AND FURO[2,3-b]PYRIDINE SYNTHESIS BASED ON THORPE-ZIEGLER REACTION.

Mikhail Yu. Yakovlev, Alexander V. Kadushkin, Natalya P. Solov'eva, Vladimir G. Granik\*

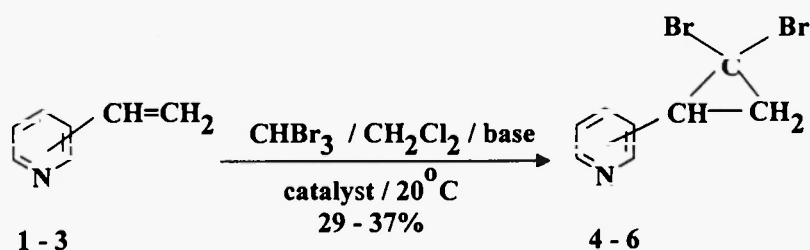
Center for Drug Chemistry, Moscow, 119815, Russian Federation

Synthesis of pyrrolo-, thieno- and furo[2,3-b]pyridines was studied under basic reaction conditions. Significant role of base catalysis as well as substituent effects in these reactions are reported.



**REACTIONS OF DIBROMOCARBENE GENERATED FROM  
BROMOFORM WITH VINYLPIRIDINES UNDER PHASE TRANSFER  
CATALYSIS CONDITIONS**

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Latvian Institute of Organic Synthesis, 21 Aizkraukles str., Riga,  
LV - 1006, Latvia

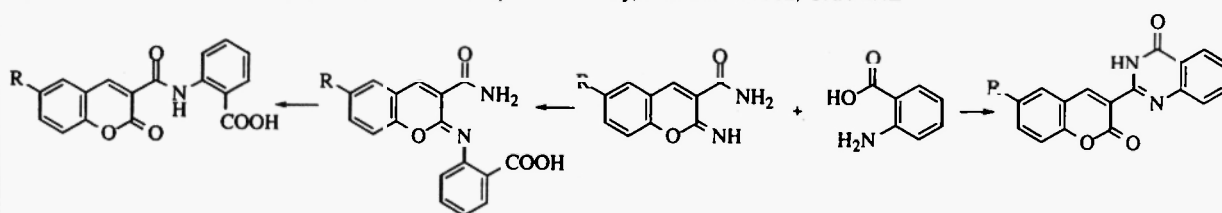


Phase transfer catalytic reactions of 2-vinylpyridine (1) with dibromocarbene generated from bromoform in the system liquid / liquid and liquid / solid were studied. The PTC system 50% aq. KOH / TEBA / CH<sub>2</sub>Cl<sub>2</sub> was found to be most active and was used for preparative synthesis of vinylpyridine dibromocarbene adducts (4 - 6).

**REARRANGEMENTS OF 2-IMINO-2H-1-BENZOPYRAN-3-CARBOXAMIDES  
UNDER ACTION OF ANTHRANILIC ACID AS N-NUCLEOPHILE**

Yaroslav V. Bilokin (Belokon),\*† Sergey N. Kovalenko, Igor E. Bylov, and Valentin P. Chernykh

*Department of Organic Chemistry, Ukrainian Academy of Pharmacy, Kharkov 310002, UKRAINE*



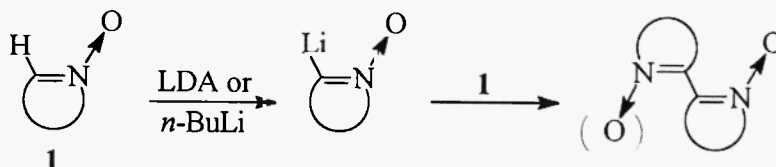
New rearrangements of 2-imino-2H-1-benzopyran-3-carboxamides under action of anthranilic acid as N-nucleophile have been revealed. These rearrangements have provided new methods for synthesis of 2-(2-oxo-2H-1-benzopyran-2-yl)-3H-quinazolin-4-ones and 2-oxo-2H-1-benzopyran-3-(N-2-carboxyphenyl)-carboxamides.

**DIPOLE-STABILIZED CARBANIONS IN SERIES OF CYCLIC ALDONITRONES.  
PART I. ALDONITRONES METALLATION AND DIMERIZATION IN LDA AND  
*n*-BuLi SOLUTIONS.**

Maxim A. Voinov\*, Igor A. Grigor'ev and Leonid B. Volodarsky

Institute of Organic Chemistry, Ave. akad. Lavrent'eva 9, 630090 Novosibirsk, Russia.

Cyclic aldonitrones of pyrroline, 3-imidazoline and 2*H*-imidazole series metallation can be used as a method of their activation towards electrophilic reagents.

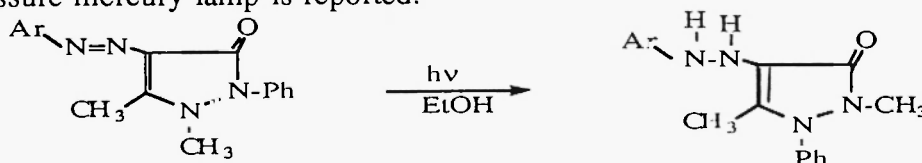


**THE PHOTOREARRANGEMENT OF SOME 4-ARYLAZOPYRAZOLIN-5-ONE  
DERIVATIVES.**

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(1) Photochemistry Dept., National Research Center, Dokki, Egypt. (2) Chemistry Dept., Faculty of Science, Cairo University, Giza, Egypt.

Photorearrangement and reduction of some 4-arylazopyrazolin-5-one derivatives by the use of high pressure mercury lamp is reported.



**SYNTHESIS OF FLUOROPHENOTHIAZINES BY SMILES REARRANGEMENT**

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Synthesis of fluorophenothiazines by Smiles rearrangement is reported. The spectral data of the synthesized compounds are also included.

