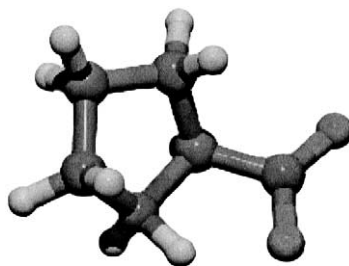


**Proline-catalyzed asymmetric reactions**

Benjamin List

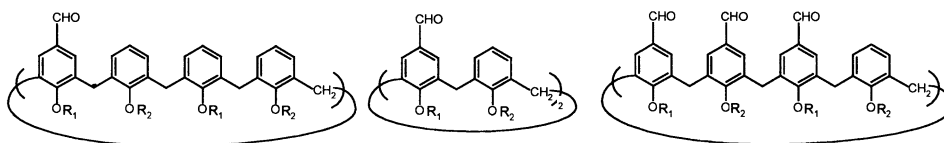
Department of Molecular Biology, The Scripps Research Institute, 10550 N. Torrey Pines Rd., La Jolla, CA 92037, USA

*Tetrahedron 58 (2002) 5573***Synthesis of selectively formylated calixarene ethers**

Vandna Arora, H. M. Chawla\* and Anuradha Santra

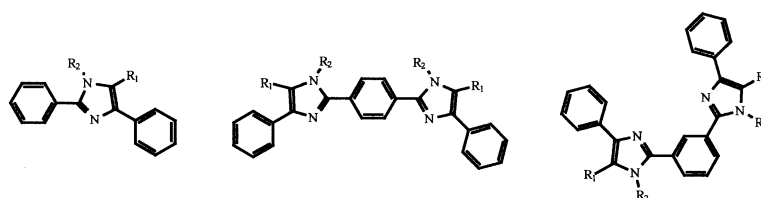
Department of Chemistry, Indian Institute of Technology, Hauz Khas, New Delhi 110016, India

Several selectively formylated calix[4]arene ethers have been synthesized.

*Tetrahedron 58 (2002) 5591***Facile two-pot syntheses of novel alternating benzene/imidazole systems**

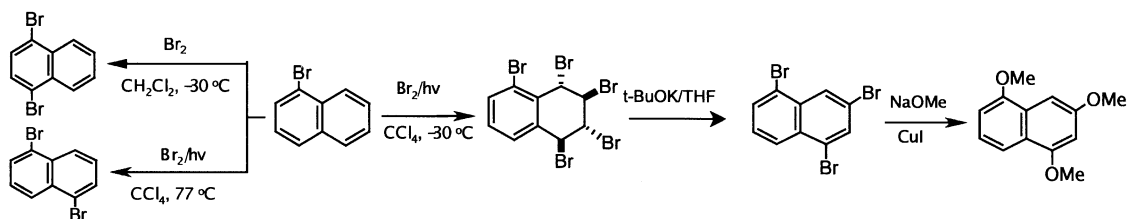
Kuangsun Sung,\* Shu-Hwa Wu and Phon-I Chen

Department of Chemistry, National Cheng Kung University, Tainan 70101, Taiwan, ROC

*Tetrahedron 58 (2002) 5599***Selective bromination of 1-bromonaphthalene: efficient synthesis of bromonaphthalene derivatives**

Osman Cakmak,\* Ibrahim Demirtas and Halis T. Balaydin

Department of Chemistry, Faculty of Science, Gaziosmanpasa University, 60240 Tokat, Turkey

*Tetrahedron 58 (2002) 5603*

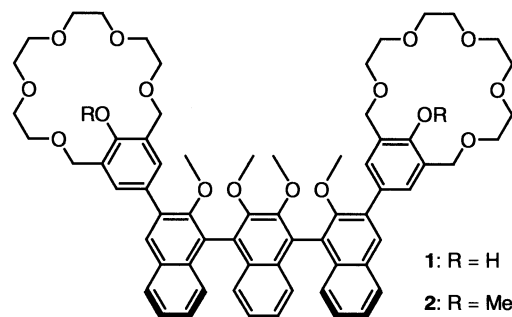
## Use of *meso*-ternaphthalene derivatives: linear recognition of the $\alpha,\omega$ -diamines by homoditopic receptors

*Tetrahedron* 58 (2002) 5611

Kazunori Tsubaki,<sup>a,\*</sup> Hiroyuki Tanaka,<sup>a</sup> Takumi Furuta,<sup>a</sup> Kiyoshi Tanaka,<sup>a</sup> Takayoshi Kinoshita<sup>b</sup> and Kaoru Fuji<sup>a,\*</sup>

<sup>a</sup>Institute for Chemical Research, Kyoto University, Uji, Kyoto 611-0011, Japan

<sup>b</sup>Exploratory Research Laboratories, Fujisawa Pharmaceutical Co., Ltd., Tokodai, Tsukuba, Ibaragi 300-2698, Japan



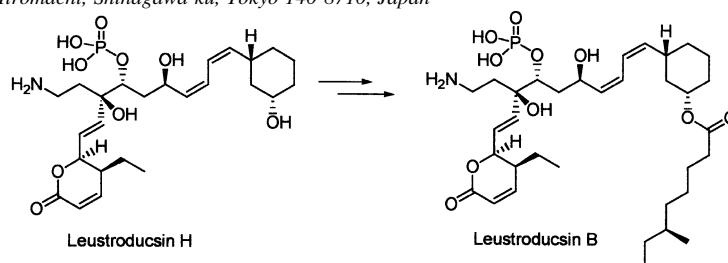
## Chemical transformation of Leustroductins: synthesis of Leustroductin B

*Tetrahedron* 58 (2002) 5619

Hayao Matsuhashi\* and Kousei Shimada

Medicinal Chemistry Research Laboratories, Sankyo Co. Ltd, Hiromachi, Shinagawa-ku, Tokyo 140-8710, Japan

Chemical transformation of Leustroductins is studied to synthesize Leustroductin B from a key intermediate Leustroductin H.



## Synthesis of chiral oxacyclic dienes via ruthenium-catalyzed enyne metathesis: useful building blocks for chiral tricyclic oxygen derivatives

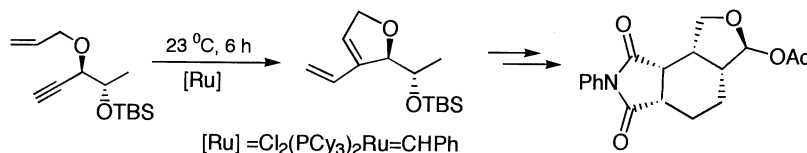
*Tetrahedron* 58 (2002) 5627

Hongyun Guo,<sup>a</sup> Reniguntala J. Madhushaw,<sup>a</sup> Fwu-Ming Shen<sup>b</sup> and Rai-Shung Liu<sup>a,\*</sup>

<sup>a</sup>Department of Chemistry, National Tsing-Hua University, Hsinchu 30043 Taiwan, ROC

<sup>b</sup>Department of Medical Technology, Yuanpei Institute of Science and Technology, Hsinchu, Taiwan, ROC

Various chiral oxacyclic dienes were prepared via ruthenium-catalyzed enyne metathesis and these dienes are useful building blocks for enantiopure tricyclic furan derivatives.

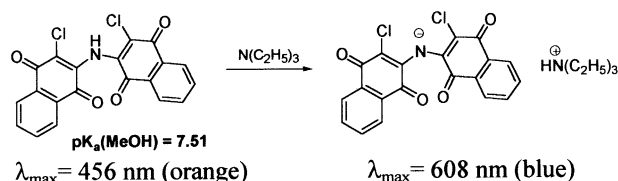


## Acidic properties of *N,N*-naphthoquinonylamines

*Tetrahedron* 58 (2002) 5639

Galina Temtsin, Sofia Gorohovsky and Shmuel Bittner\*

Department of Chemistry, Ben-Gurion University of the Negev, Beer Sheva 84105, Israel



## Synthesis of new pyridazino[4,5-c]isoquinolinones by Suzuki cross-coupling reaction

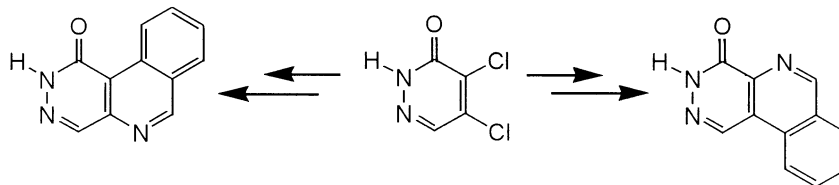
*Tetrahedron 58 (2002) 5645*

Zsuzsanna Riedl,<sup>a</sup> Bert U. W. Maes,<sup>b</sup> Katrien Monsieurs,<sup>b</sup> Guy L. F. Lemière,<sup>b</sup> Péter Mátyus<sup>c</sup> and György Hajós<sup>a,\*</sup>

<sup>a</sup>Institute of Chemistry, Chemical Research Center, Hungarian Academy of Sciences, P.O. Box 17, H-1525 Budapest, Hungary

<sup>b</sup>Department of Chemistry, University of Antwerp (RUCA), Groenenborgerlaan 171, B-2020 Antwerpen, Belgium

<sup>c</sup>Institute of Organic Chemistry, Semmelweis University, H-1092 Budapest, Hőgyes E. u. 7., Hungary

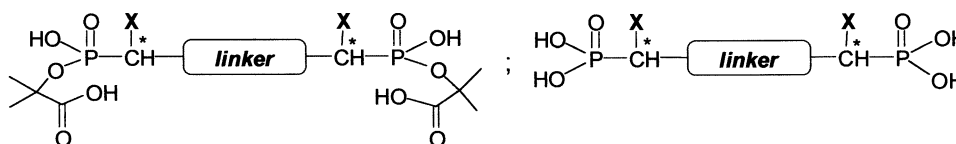


## Efficient synthesis of bolaform- and gemini-type alkyl-bis-[( $\alpha$ -amino)phosphonocarboxylic or phosphonic acid] surfactants

*Tetrahedron 58 (2002) 5651*

Karine Verduyts-Moreira, Christophe Déjugnat\* and Guita Etemad-Moghadam

Laboratoire des IMRCP (UMR 5623), Université Paul Sabatier, 118 route de Narbonne-Bât. 2R1, 31062 Toulouse cedex 04, France



## Regio- and stereo-selective biotransformation of 2 $\alpha$ ,5 $\alpha$ ,10 $\beta$ ,14 $\beta$ -tetra-acetoxy-4(20), 11-taxadiene by *Ginkgo* cell suspension cultures

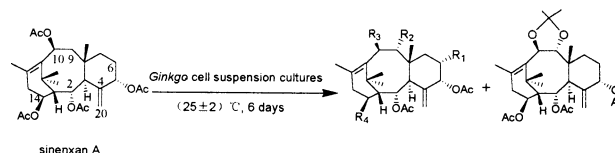
*Tetrahedron 58 (2002) 5659*

Jungui Dai,<sup>a</sup> Min Ye,<sup>a</sup> Hongzhu Guo,<sup>a</sup> Weihua Zhu,<sup>b</sup> Dayong Zhang,<sup>b</sup> Qiu Hu,<sup>b</sup> Junhua Zheng<sup>a</sup> and Dean Guo<sup>a,\*</sup>

<sup>a</sup>The State Key Laboratory of Natural and Biomimetic Drugs, School of Pharmaceutical Sciences, Peking University, Xueyuan Road #38, Beijing 100083, People's Republic of China

<sup>b</sup>Institute of Materia Medica, Chinese Academy of Medical Sciences and Peking Union Medical College, Xiannong Tan Street #1, Beijing 100050, People's Republic of China

2 $\alpha$ ,5 $\alpha$ ,10 $\beta$ ,14 $\beta$ -Tetra-acetoxy-4(20),11-taxadiene was regio- and stereo-selectively biotransformed by *Ginkgo* cell suspension cultures, and eight products were obtained, among which five are new compounds.



## A modified palladium catalysed reductive amination procedure

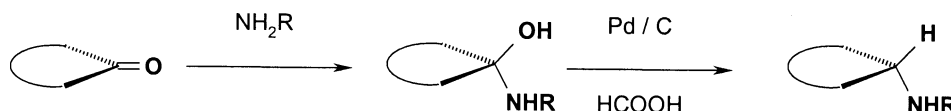
*Tetrahedron 58 (2002) 5669*

Valerio Berdini,<sup>a</sup> Maria C. Cesta,<sup>b</sup> Roberto Curti,<sup>b</sup> Gaetano D'Anniballe,<sup>b</sup> Nicoletta Di Bello,<sup>b</sup> Giuseppe Nano,<sup>b</sup> Luca Nicolini,<sup>b</sup> Alessandra Topai<sup>b</sup> and Marcello Allegretti<sup>b,\*</sup>

<sup>a</sup>Astex Technology, 250 Cambridge Science Park, Milton Road, Cambridge CB4 0WE, UK

<sup>b</sup>Chemistry Department, Dompé S.p.A. Research and Development Centre, V. Campo di Pile, 67100 L'Aquila, Italy

New, extended applications of a modified palladium catalysed reductive amination procedure are described; a mechanistic hypothesis alternative to the common imine pathway is proposed.

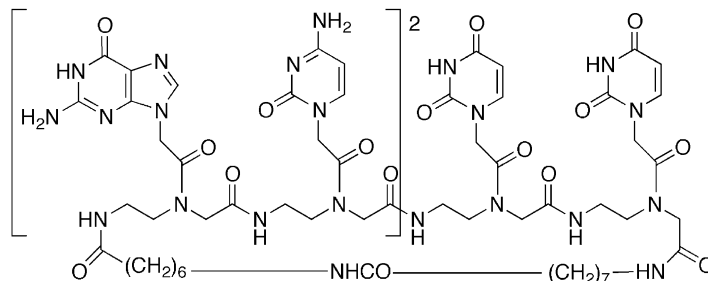


## Cyclic PNA hexamer-based compound: modelling, synthesis and inhibition of the HIV-1 RNA dimerization process

*Tetrahedron* 58 (2002) 5675

Caroline Schwergold,<sup>a</sup> Geoffrey Depecker,<sup>a</sup> Christophe Di Giorgio,<sup>a</sup> Nadia Patino,<sup>a</sup> Fabrice Jossinet,<sup>b</sup> Bernard Ehresmann,<sup>b</sup> Raphael Terreux,<sup>c</sup> Daniel Cabrol-Bass<sup>c</sup> and Roger Condom<sup>a,\*</sup>

<sup>a</sup>Laboratoire de Chimie Bio-organique, Université de Nice-Sophia Antipolis, UMR UNSA-CNRS 6001, F-06108 Nice cedex 2, France  
<sup>b</sup>Institut de Biologie Moléculaire et Cellulaire, CNRS UPR 9002, 15 rue Descartes, F-64084 Strasbourg cedex, France  
<sup>c</sup>Laboratoire A.S.I. Equipe Chimométrie et Modélisation, Université de Nice-Sophia Antipolis, F-06108 Nice cedex 2, France

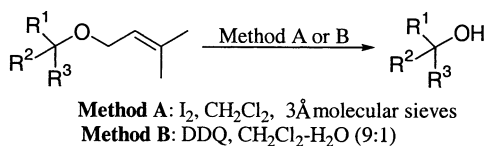


## The prenyl group: a versatile hydroxy protecting group, removable chemoselectively under mild conditions

*Tetrahedron* 58 (2002) 5689

Jean-Michel Vatèle

Laboratoire de Chimie Organique 1, UMR 5622 CNRS, Domaine Scientifique de la Doua, CPE 3 rue Victor Grignard, 69616 Villeurbanne cedex, France



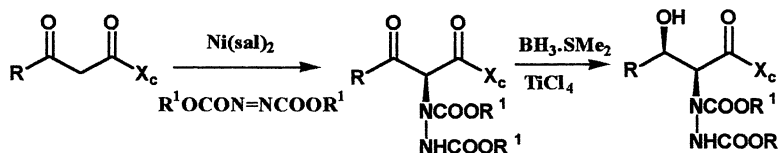
## Ni(II)-catalyzed Michael additions. Part 2: Dynamic kinetic resolution in the reduction of chiral α-hydrazino-β-ketoacid derivatives

*Tetrahedron* 58 (2002) 5699

Caroline Marchi,<sup>a</sup> Elisenda Trepát,<sup>a</sup> Marcial Moreno-Mañas,<sup>a</sup> Adelina Vallribera<sup>a,\*</sup> and Elies Molins<sup>b</sup>

<sup>a</sup>Department of Chemistry, Universitat Autònoma de Barcelona, Cerdanyola, 08193 Barcelona, Spain

<sup>b</sup>Institut de Ciència de Materials de Barcelona (CSIC), Campus de la UAB, E-08193 Cerdanyola, 08193 Barcelona, Spain



## Novel cytotoxic acylphenol dimers of *Myristica gigantea*; enzymatic synthesis of giganteones A and B

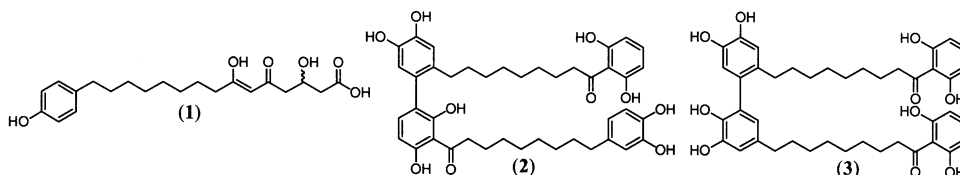
*Tetrahedron* 58 (2002) 5709

Van Cuong Pham,<sup>a</sup> Akino Jossang,<sup>a</sup> Thierry Sévenet<sup>b</sup> and Bernard Bodo<sup>a,\*</sup>

<sup>a</sup>Laboratoire de Chimie des Substances Naturelles, ESA 8041 CNRS, Muséum National d'Histoire Naturelle, 63 rue Buffon, 75005 Paris, France

<sup>b</sup>Institut de Chimie des Substances Naturelles, 1206 CNRS, 91198 Gif sur Yvette Cedex, France

The structure determination of **1**, alkylphenol precursor of malabaricone, and of **2** and **3**, acylphenol dimers is described. An enzymatic synthesis of **2** and **3** from malabaricone C was performed.



## Xanthates and solid-phase chemistry. A new soluble polymer analogue of Wang resin

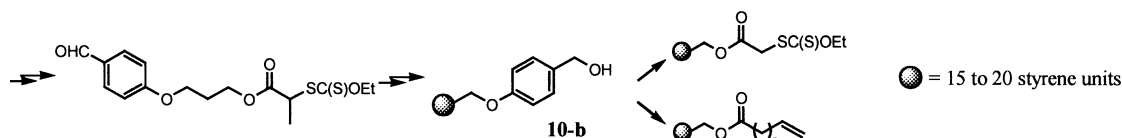
*Tetrahedron 58 (2002) 5715*

Anne-Claude Dublanchet,<sup>a</sup> Marie Lusinchi<sup>a,b,\*</sup> and Samir Z. Zard<sup>b</sup>

<sup>a</sup>Department of Chemistry, Pfizer Global Research and Development, Fresnes Laboratories, 3-9 rue de la loge, F-94265 Fresnes, France

<sup>b</sup>Laboratoire de synthèse Organique associé au CNRS, Ecole Polytechnique, 91128 Palaiseau Cedex, France

Inter-molecular radical additions of xanthates onto olefins were performed with the new resin **10-b** in comparison with the classical Wang resin.



## Dioxane-type (2-naphthyl)methylene acetals of glycosides and their hydrogenolytic transformation into 6-O- and 4-O-(2-naphthyl)methyl (NAP) ethers

*Tetrahedron 58 (2002) 5723*

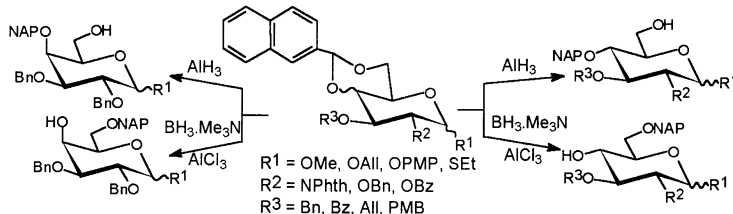
Anikó Borbás,<sup>a</sup> Zoltán B. Szabó,<sup>a</sup> László Szilágyi,<sup>b</sup> Attila Bényei<sup>c</sup> and András Lipták<sup>a,d,\*</sup>

<sup>a</sup>Research Group for Carbohydrates of the Hungarian Academy of Sciences, P.O. Box 55, Debrecen H-4010, Hungary

<sup>b</sup>Department of Organic Chemistry, Faculty of Science, University of Debrecen, P.O. Box 20, Debrecen H-4010, Hungary

<sup>c</sup>Faculty of Science, Institute of Physical Chemistry, University of Debrecen, P.O. Box 7, Debrecen H-4010, Hungary

<sup>d</sup>Department of Biochemistry, Faculty of Science, University of Debrecen, P.O. Box 55, Debrecen H-4010, Hungary



## Reaction of 2-hetarylacetonitriles with ethyl 2-alkylsulfanyl-4-chloro-5-pyrimidinecarboxylates. Synthesis of new condensed pyrimidines

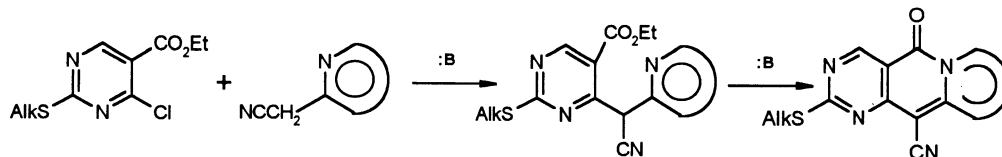
*Tetrahedron 58 (2002) 5733*

E. V. Blyumin,<sup>a</sup> Yu. M. Volovenko,<sup>a</sup> Hans Neunhoffer,<sup>b,\*</sup> S. V. Shishkina,<sup>c</sup> R. A. Zubatyuk<sup>c</sup> and Oleg V. Shishkin<sup>c</sup>

<sup>a</sup>Chemical Department, Kiev Taras Shevchenko University, Volodymyrska 64, Kiev 01033, Ukraine

<sup>b</sup>Institute of Organic Chemistry, Darmstadt University of Technology, Petersenstraße 22, D-64287 Darmstadt, Germany

<sup>c</sup>Scientific Research Department of Alkali Halide Crystals, National Academy of Sciences of Ukraine, 60 Lenina ave., Khar'kov 310001, Ukraine



## Solid and solution phase syntheses of the 2-cyanopyrrolidide DPP-IV inhibitor NVP-DPP728

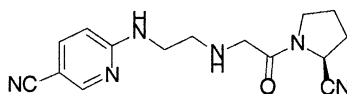
*Tetrahedron 58 (2002) 5741*

Nicolas Willand,<sup>a</sup> Jurgen Joossens,<sup>a</sup> Jean-Claude Gesquière,<sup>a</sup> André L. Tartar,<sup>a</sup> D. Michael Evans<sup>b</sup> and Michael B. Roe<sup>b,\*</sup>

<sup>a</sup>Laboratoire de chimie organique, UMR 8525, Faculté des sciences pharmaceutiques et biologiques, 3 rue du Pr. Laguesse, F-59006 Lille Cedex, France

<sup>b</sup>Ferring Research Limited, Chilworth Research Centre, Southampton SO16 7NP, UK

Alternative synthetic approaches to NVP-DPP728 are reported.



## Conformationally constrained 1,4-DHPs. A convenient route to bis-1,4-DHPs as a novel class of nitrogen compounds

Tetrahedron 58 (2002) 5747

Štefan Marchalín,<sup>a</sup> Miloslav Chudík,<sup>a</sup> Katarína Cvopová,<sup>a</sup> Jozef Kozíšek,<sup>b</sup> Ján Leško<sup>c</sup> and Adam Daich<sup>d,\*</sup>

<sup>a</sup>Department of Organic Chemistry, Faculty of Chemical Technology, Slovak University of Technology, SK-812 37 Bratislava, Slovak Republic

<sup>b</sup>Department of Physical Chemistry and Central Laboratory, Faculty of Chemical Technology, Slovak University of Technology, SK-812 37 Bratislava, Slovak Republic

<sup>c</sup>Central Laboratory, Faculty of Chemical Technology, Slovak University of Technology, SK-812 37 Bratislava, Slovak Republic

<sup>d</sup>Laboratoire de Chimie de l'Université du Havre, Faculté des Sciences and Techniques, URCOM, EA 3221, 25 rue Philippe Lebon, B.P.: 540, F-76058 Le Havre Cedex, France

Various functionalised indolizines and corresponding bis-1,4-DHPs were synthesised efficiently in an acidic medium from 2-formyl-1,4-DHP derivatives **2** in one pot procedure.

