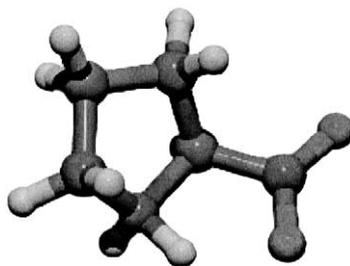


Proline-catalyzed asymmetric reactions

Benjamin List

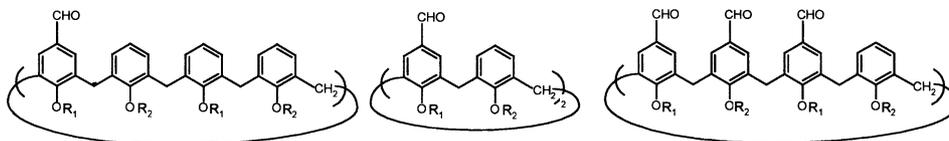
Department of Molecular Biology, The Scripps Research Institute, 10550 N. Torrey Pines Rd., La Jolla, CA 92037, USA

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

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

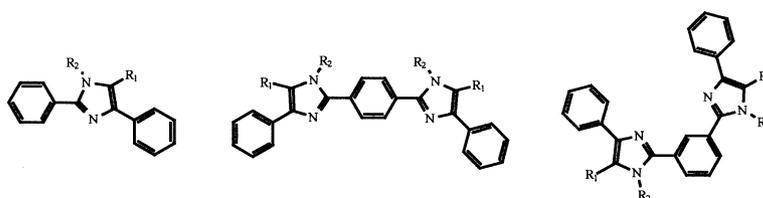
Department of Chemistry, Indian Institute of Technology, Hauz Khas, New Delhi 110016, India

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

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

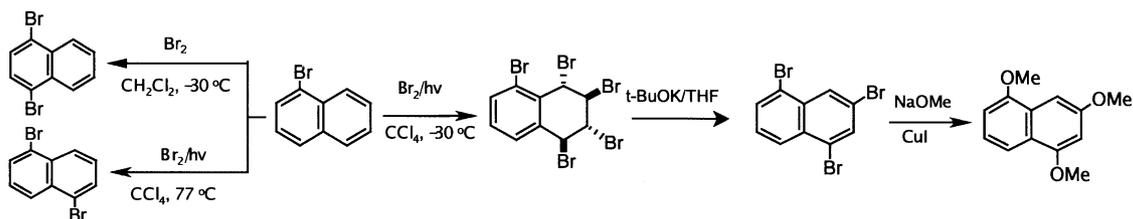
Kuangseng Sung,* Shu-Hwa Wu and Phon-I Chen

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

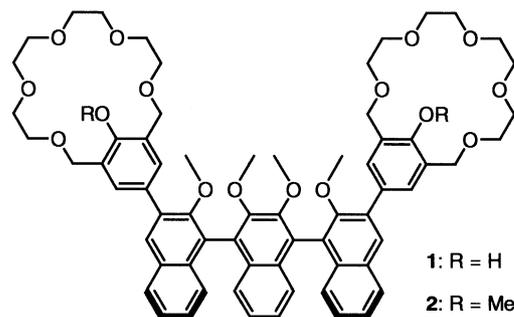
Use of *meso*-ternaphthalene derivatives: linear recognition of the α,ω -diamines by homoditopic receptors

Tetrahedron 58 (2002) 5611

Kazunori Tsubaki,^{a,*} Hiroyuki Tanaka,^a Takumi Furuta,^a Kiyoshi Tanaka,^a Takayoshi Kinoshita^b and Kaoru Fuji^{a,*}

^aInstitute for Chemical Research, Kyoto University, Uji, Kyoto 611-0011, Japan

^bExploratory Research Laboratories, Fujisawa Pharmaceutical Co., Ltd., Tokodai, Tsukuba, Ibaragi 300-2698, Japan



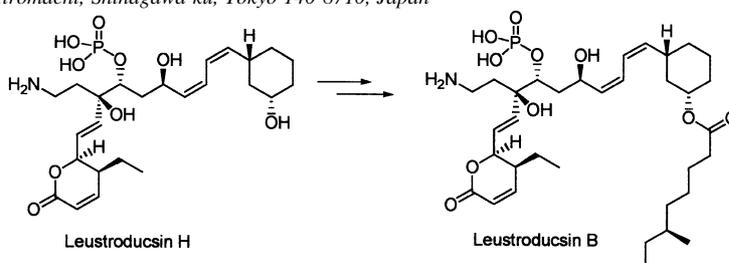
Chemical transformation of Leustroductins: synthesis of Leustroductin B

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

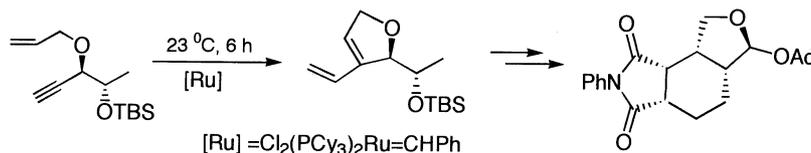
Tetrahedron 58 (2002) 5627

Hongyun Guo,^a Reniguntala J. Madhushaw,^a Fwu-Ming Shen^b and Rai-Shung Liu^{a,*}

^aDepartment of Chemistry, National Tsing-Hua University, Hsinchu 30043 Taiwan, ROC

^bDepartment 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.

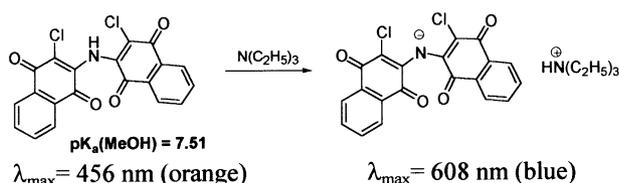


Acidic properties of *N,N*-naphthoquinonylamines

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

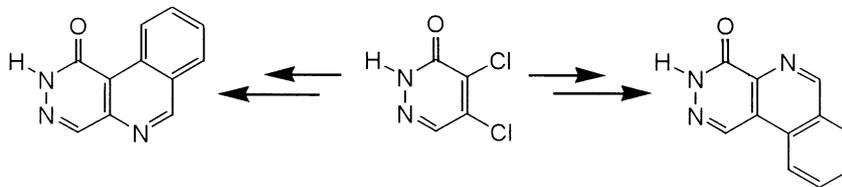
Tetrahedron 58 (2002) 5645

Zsuzsanna Riedl,^a Bert U. W. Maes,^b Katrien Monsieurs,^b Guy L. F. Lemière,^b Péter Mátyus^c and György Hajós^{a,*}

^aInstitute of Chemistry, Chemical Research Center, Hungarian Academy of Sciences, P.O. Box 17, H-1525 Budapest, Hungary

^bDepartment of Chemistry, University of Antwerp (RUCA), Groenenborgerlaan 171, B-2020 Antwerpen, Belgium

^cInstitute of Organic Chemistry, Semmelweis University, H-1092 Budapest, Hőgyes E. u. 7., Hungary

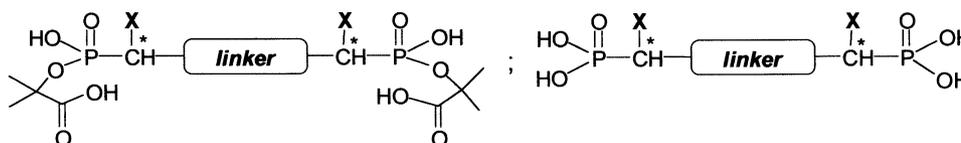


Efficient synthesis of bolaform- and gemini-type alkyl-bis-[(α -amino)phosphonocarboxylic or phosphonic acid] surfactants

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 α ,5 α ,10 β ,14 β -tetra-acetoxy-4(20), 11-taxadiene by *Ginkgo* cell suspension cultures

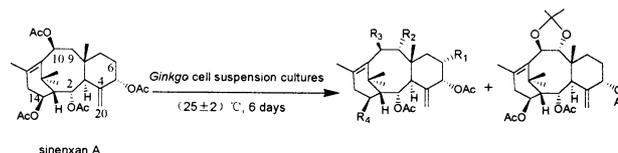
Tetrahedron 58 (2002) 5659

Jungui Dai,^a Min Ye,^a Hongzhu Guo,^a Weihua Zhu,^b Dayong Zhang,^b Qiu Hu,^b Junhua Zheng^a and Dean Guo^{a,*}

^aThe State Key Laboratory of Natural and Biomimetic Drugs, School of Pharmaceutical Sciences, Peking University, Xueyuan Road #38, Beijing 100083, People's Republic of China

^bInstitute 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 α ,5 α ,10 β ,14 β -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

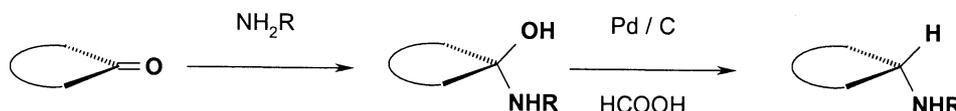
Tetrahedron 58 (2002) 5669

Valerio Berdini,^a Maria C. Cesta,^b Roberto Curti,^b Gaetano D'Anniballe,^b Nicoletta Di Bello,^b Giuseppe Nano,^b Luca Nicolini,^b Alessandra Topai^b and Marcello Allegretti^{b,*}

^aAstex Technology, 250 Cambridge Science Park, Milton Road, Cambridge CB4 0WE, UK

^bChemistry 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.

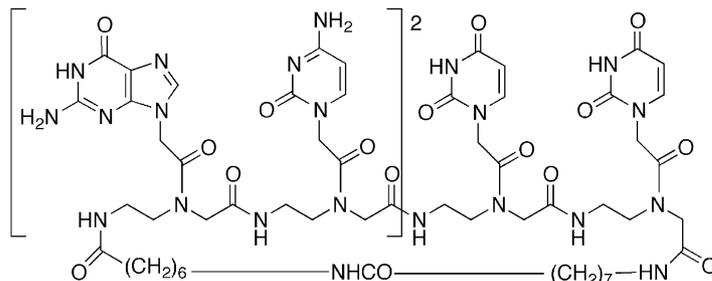


Cyclic PNA hexamer-based compound: modelling, synthesis and inhibition of the HIV-1 RNA dimerization process

Tetrahedron 58 (2002) 5675

Caroline Schwergold,^a Geoffrey Depecker,^a Christophe Di Giorgio,^a Nadia Patino,^a Fabrice Jossinet,^b Bernard Ehresmann,^b Raphael Terreux,^c Daniel Cabrol-Bass^c and Roger Condom^{a,*}

^aLaboratoire de Chimie Bio-organique, Université de Nice-Sophia Antipolis, UMR UNSA-CNRS 6001, F-06108 Nice cedex 2, France
^bInstitut de Biologie Moléculaire et Cellulaire, CNRS UPR 9002, 15 rue Descartes, F-64084 Strasbourg cedex, France
^cLaboratoire A.S.I. Equipe Chimométrie et Modélisation, Université de Nice-Sophia Antipolis, F-06108 Nice cedex 2, France

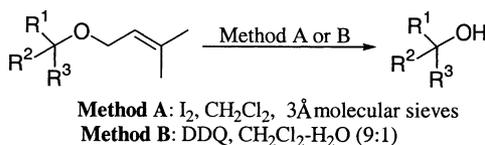


The prenyl group: a versatile hydroxy protecting group, removable chemoselectively under mild conditions

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



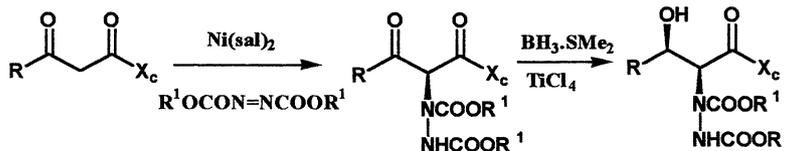
Ni(II)-catalyzed Michael additions. Part 2: Dynamic kinetic resolution in the reduction of chiral α-hydrazino-β-ketoacid derivatives

Tetrahedron 58 (2002) 5699

Caroline Marchi,^a Elisenda Trepát,^a Marcial Moreno-Mañas,^a Adelina Vallribera^{a,*} and Elies Molins^b

^aDepartment of Chemistry, Universitat Autònoma de Barcelona, Cerdanyola, 08193 Barcelona, Spain

^bInstitut 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

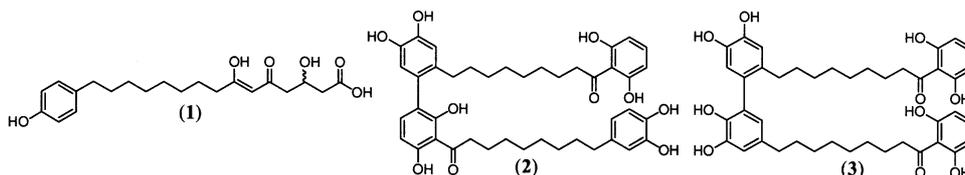
Tetrahedron 58 (2002) 5709

Van Cuong Pham,^a Akino Jossang,^a Thierry Sévenet^b and Bernard Bodo^{a,*}

^aLaboratoire de Chimie des Substances Naturelles, ESA 8041 CNRS, Muséum National d'Histoire Naturelle, 63 rue Buffon, 75005 Paris, France

^bInstitut 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

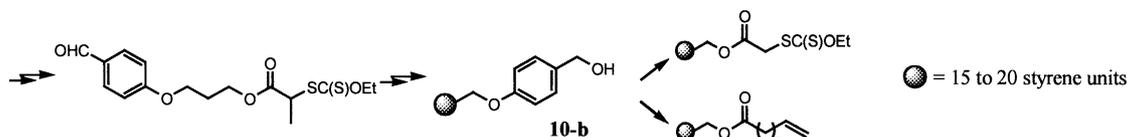
Tetrahedron 58 (2002) 5715

Anne-Claude Dublanchet,^a Marie Lusinchi^{a,b,*} and Samir Z. Zard^b

^aDepartment of Chemistry, Pfizer Global Research and Development, Fresnes Laboratories, 3-9 rue de la loge, F-94265 Fresnes, France

^bLaboratoire 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

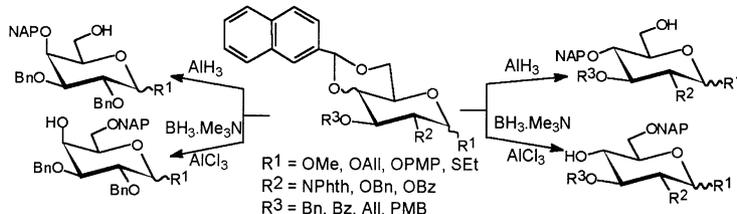
Anikó Borbás,^a Zoltán B. Szabó,^a László Szilágyi,^b Attila Bényei^c and András Lipták^{a,d,*}

^aResearch Group for Carbohydrates of the Hungarian Academy of Sciences, P.O. Box 55, Debrecen H-4010, Hungary

^bDepartment of Organic Chemistry, Faculty of Science, University of Debrecen, P.O. Box 20, Debrecen H-4010, Hungary

^cFaculty of Science, Institute of Physical Chemistry, University of Debrecen, P.O. Box 7, Debrecen H-4010, Hungary

^dDepartment 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

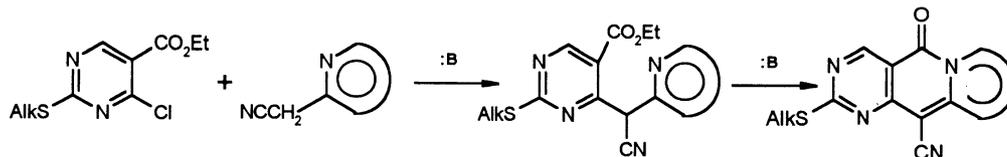
Tetrahedron 58 (2002) 5733

E. V. Blyumin,^a Yu. M. Volovenko,^a Hans Neunhoffer,^{b,*} S. V. Shishkina,^c R. A. Zubatyuk^c and Oleg V. Shishkin^c

^aChemical Department, Kiev Taras Shevchenko University, Volodymyrska 64, Kiev 01033, Ukraine

^bInstitute of Organic Chemistry, Darmstadt University of Technology, Petersenstraße 22, D-64287 Darmstadt, Germany

^cScientific 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

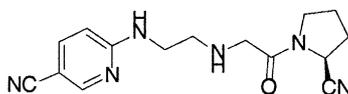
Tetrahedron 58 (2002) 5741

Nicolas Willand,^a Jurgen Joossens,^a Jean-Claude Gesquière,^a André L. Tartar,^a D. Michael Evans^b and Michael B. Roe^{b,*}

^aLaboratoire de chimie organique, UMR 8525, Faculté des sciences pharmaceutiques et biologiques, 3 rue du Pr. Laguesse, F-59006 Lille Cedex, France

^bFerring 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,^a Miloslav Chudík,^a Katarína Cvopová,^a Jozef Kozíšek,^b Ján Leško^c and Adam Daich^{d,*}

^aDepartment of Organic Chemistry, Faculty of Chemical Technology, Slovak University of Technology, SK-812 37 Bratislava, Slovak Republic

^bDepartment of Physical Chemistry and Central Laboratory, Faculty of Chemical Technology, Slovak University of Technology, SK-812 37 Bratislava, Slovak Republic

^cCentral Laboratory, Faculty of Chemical Technology, Slovak University of Technology, SK-812 37 Bratislava, Slovak Republic

^dLaboratoire 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.

