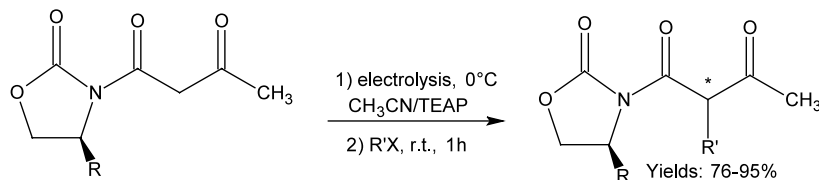


Electrochemical generation of tetraethylammonium *N*-acetoacetyl-oxazolidin-2-one enolates: an easy access to α -alkylated acetoacetic derivatives

Tetrahedron Letters 43 (2002) 2881

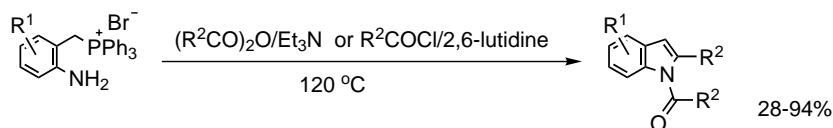
Laura Palombi,^{a,*} Marta Feroci,^b Monica Orsini,^a Leucio Rossi^a and Achille Inesi^{a,*}^aDip. di Chimica, Ingegneria Chimica e Materiali, Università degli Studi, I-67040 Monteluco di Roio, L'Aquila, Italy^bDip. di Ingegneria Chimica, Materiali, Materie Prime e Metallurgia, Università 'La Sapienza', Via Castro Laurenziano 7, I-00161 Roma, Italy

A facile synthesis of 2-substituted indoles from (2-aminobenzyl)-triphenylphosphonium salts

Tetrahedron Letters 43 (2002) 2885

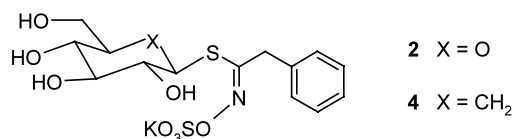
Shin'ichi Taira, Hiroshi Danjo and Tsuneo Imamoto*

Department of Chemistry, Faculty of Science, Chiba University, Yayoi-cho, Inage-ku, Chiba 263-8522, Japan



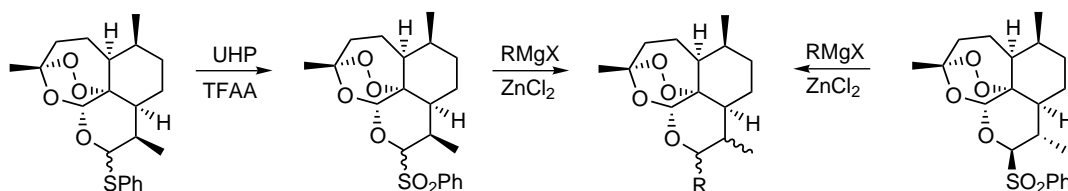
Carba-glucotropaeolin: the first non-hydrolyzable glucosinolate analogue, to inhibit myrosinase

Tetrahedron Letters 43 (2002) 2889

Myriam Lefoix,^a Arnaud Tatibouët,^{a,*} Sylvain Cottaz,^b Hugues Driguez^b and Patrick Rollin^{a,*}^aInstitut de Chimie Organique et Analytique, Université d'Orléans, BP6759, F-45067 Orléans Cedex 2, France^bCentre de Recherches sur les Macromolécules Végétales, CNRS, BP 53, F-38041 Grenoble Cedex 9, FranceA 5a-carba-analogue **4** of glucotropaeolin **2** was synthesized in racemic form and showed a good inhibiting power against myrosinase.

A simple synthesis of C-10 substituted deoxyartemisinin and 9-*epi*-deoxyartemisinin with various organozinc reagents

Tetrahedron Letters 43 (2002) 2891

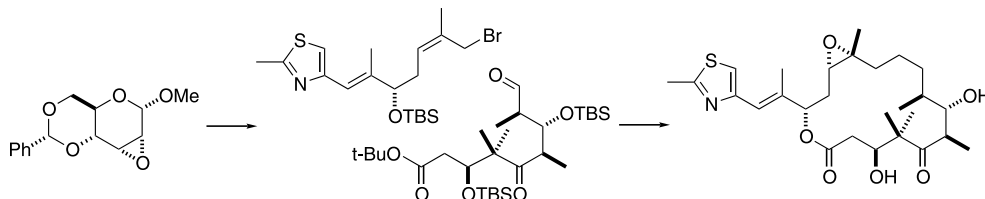
Seokjoon Lee^{a,*} and Sangtae Oh^b^aDepartment of Premedical Science, College of Medicine, Kwandong University, Gangnung 210-701, South Korea^bDepartment of Chemistry, Yonsei University, Wonju 220-710, South Korea

Synthesis of epothilones B and D from D-glucose

Mikhail S. Ermolenko* and Pierre Potier

Institut de Chimie des Substances Naturelles du CNRS, Avenue de la Terrasse, 91198 Gif-sur-Yvette, France

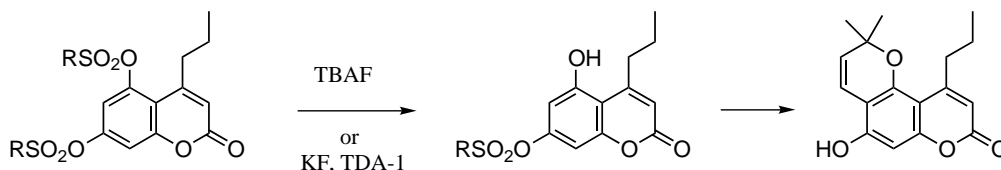
An enantiospecific total synthesis of epothilones B and D from D-glucose is reported.

**A novel synthesis of 5-hydroxy-2,2-dimethyl-10-propyl-2H-pyrano[2,3-f]chromen-8-one**

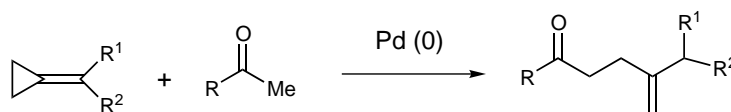
Martin E. Fox, Ian C. Lennon* and Graham Meek

Chirotech Technology Ltd, Unit 321 Cambridge Science Park, Milton Road, Cambridge CB4 0WG, UK

A novel synthesis of a key precursor for calanolide A is described.

**Palladium-catalyzed addition of ketones to alkylidenecyclopropanes**

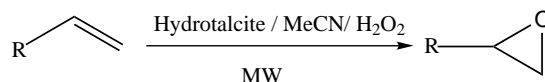
Drexel H. Camacho, Itaru Nakamura, Byoung Ho Oh, Shinichi Saito and Yoshinori Yamamoto*

*Department of Chemistry, Graduate School of Science, Tohoku University, Sendai 980-8578, Japan*Alkylidenecyclopropanes react with ketones under neutral conditions in the presence of a Pd(0) catalyst to afford the corresponding α -allylated ketone products in moderate to good yields.**Microwave-expedited olefin epoxidation over hydrotalcites using hydrogen peroxide and acetonitrile**

Unnikrishnan R. Pillai, Endalkachew Sahle-Demessie* and Rajender S. Varma

National Risk Management Research Laboratory, Sustainable Technology Division, MS 443, US Environmental Protection Agency, 26 West M. L. King Drive, Cincinnati, OH 45268, USA

Microwave-assisted epoxidation of olefins is described over hydrotalcite catalysts in the presence of hydrogen peroxide and acetonitrile.

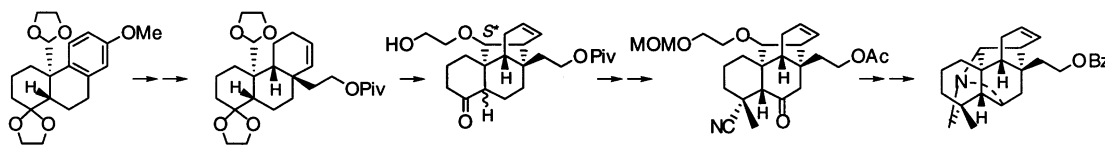


Synthesis of a compound having the essential structural unit for the hetisine-type of aconite alkaloids

Tetrahedron Letters 43 (2002) 2913

Hideaki Muratake* and Mitsutaka Natsume*

Research Foundation Itsuu Laboratory, 2-28-10 Tamagawa, Setagaya-ku, Tokyo 158-0094, Japan



Practical synthesis of chiral ligands for catalytic enantioselective cyanosilylation of ketones

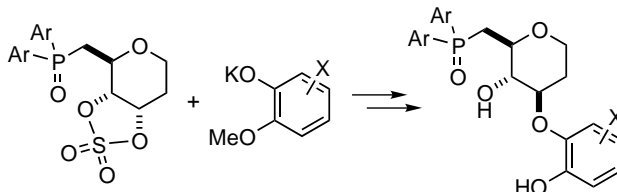
Tetrahedron Letters 43 (2002) 2919

Shuji Masumoto,^a Kazuo Yabu,^a Motomu Kanai^{a,b} and Masakatsu Shibasaki^{a,*}

^aGraduate School of Pharmaceutical Sciences, The University of Tokyo, Hongo, Bunkyo-ku, Tokyo 113-0033, Japan

^bPRESTO, Japan Science and Technology Corporation (JST), Tokyo, Japan

A range of chiral ligands that are useful for catalytic enantioselective cyanosilylation of ketones was practically synthesized using a cyclic sulfate as a key intermediate.



Studies toward practical synthesis of (20S)-camptothecin family through catalytic enantioselective cyanosilylation of ketones: improved catalyst efficiency by ligand-tuning

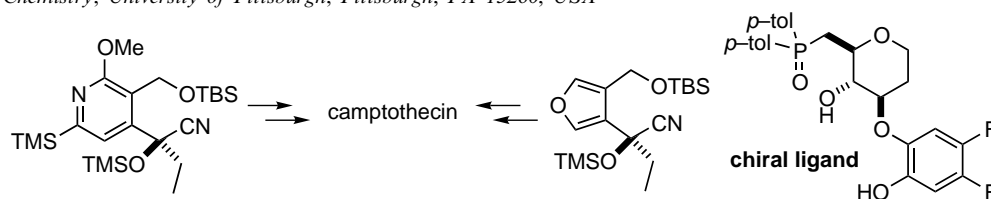
Tetrahedron Letters 43 (2002) 2923

Kazuo Yabu,^a Shuji Masumoto,^a Motomu Kanai,^{a,b} Dennis P. Curran^c and Masakatsu Shibasaki^{a,*}

^aGraduate School of Pharmaceutical Sciences, The University of Tokyo, Hongo, Bunkyo-ku, Tokyo 113-0033, Japan

^bPRESTO, Japan Science and Technology Corporation (JST), Tokyo, Japan

^cDepartment of Chemistry, University of Pittsburgh, Pittsburgh, PA 15260, USA

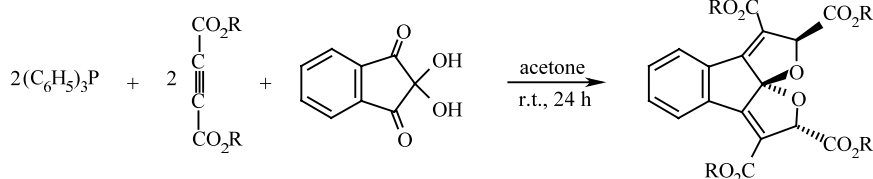


An efficient diastereoselective one-pot synthesis of dihydrofuro-[2',3':2,3]indeno[2,1-b]furan derivatives

Tetrahedron Letters 43 (2002) 2927

Issa Yavari,* Mehdi Adib and Mohammad Hosein Sayahi

Department of Chemistry, University of Tarbiat Modarres, PO Box 14115-175, Tehran, Iran



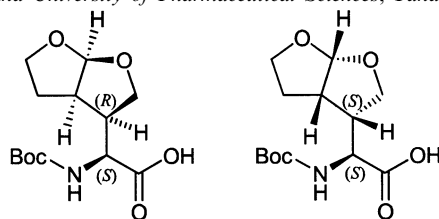
Synthesis of novel amino acids, L-bis-tetrahydrofuranylglycines

Tetrahedron Letters 43 (2002) 2931

Ei'ichi Ami,^a S. Rajesh,^a Jung Wang,^a Tooru Kimura,^a Yoshio Hayashi,^a Yoshiaki Kiso^{a,*} and Toshimasa Ishida^b

^aDepartment of Medicinal Chemistry, Center for Frontier Research in Medicinal Science, Kyoto Pharmaceutical University, Yamashina-ku, Kyoto 607-8412, Japan

^bDepartment of Physical Chemistry, Osaka University of Pharmaceutical Sciences, Takatsuki, Osaka 569-1604, Japan

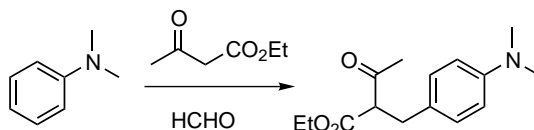


Novel Mannich-type nucleophilic substitution reaction with tertiary aromatic amines

Tetrahedron Letters 43 (2002) 2935

Hiroyasu Takahashi, Nobuyuki Kashiwa, Yuichi Hashimoto and Kazuo Nagasawa*

Institute of Molecular and Cellular Biosciences, University of Tokyo, 1-1-1 Yayoi, Bunkyo-ku, Tokyo 113-0032, Japan

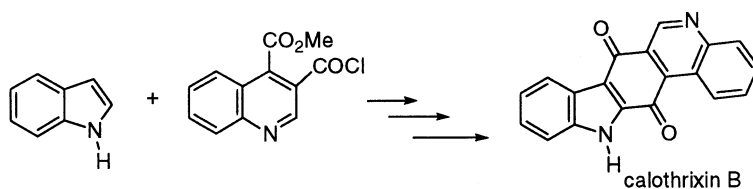


A simple and concise route to calothrixin B

Tetrahedron Letters 43 (2002) 2939

Paul H. Bernardo, Christina L. L. Chai* and John A. Elix

Department of Chemistry, Australian National University, Canberra, ACT 0200, Australia



Lotthanongine, an unprecedented flavonoidal indole alkaloid from the roots of Thai medicinal plant, *Trigonostemon reidioides*

Tetrahedron Letters 43 (2002) 2941

Tripetch Kanchanapoom,^{a,b} Ryoji Kasai,^a Phannipha Chumsri,^c Krisana Kraintintu^d and Kazuo Yamasaki^{a,*}

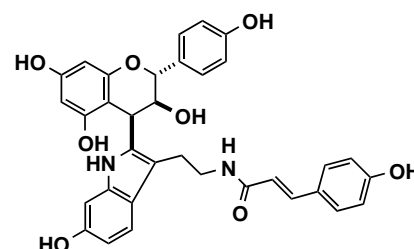
^aInstitute of Pharmaceutical Sciences, Faculty of Medicine, Hiroshima University, 1-2-3 Kasumi, Minami-ku, Hiroshima 734-8551, Japan

^bDepartment of Pharmaceutical Botany and Pharmacognosy, Faculty of Pharmaceutical Sciences, Khon Kaen University, Khon Kaen 40002, Thailand

^cDepartment of Pharmacognosy, Faculty of Pharmacy, Mahidol University, Bangkok 10400, Thailand

^dResearch and Development Institute, Government Pharmaceutical Organization, Bangkok 10400, Thailand

Lotthanongine was isolated from the roots of *Trigonostemon reidioides*.



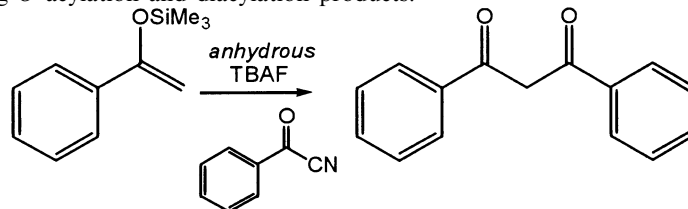
The regioselective preparation of 1,3-diketones

Charlotte Wiles,^a Paul Watts,^a Stephen J. Haswell^{a,*} and Esteban Pombo-Villar^b

^aDepartment of Chemistry, Faculty of Science and the Environment, University of Hull, Cottingham Road, Hull HU6 7RX, UK

^bNervous Systems Research, WSJ-386.07.15, Novartis Pharma Ltd., CH4002, Basel, Switzerland

We illustrate a simple method for the regioselective preparation of 1,3-diketones from both Li enolates and silyl enol ethers, free from competing *O*-acylation and diacylation products.



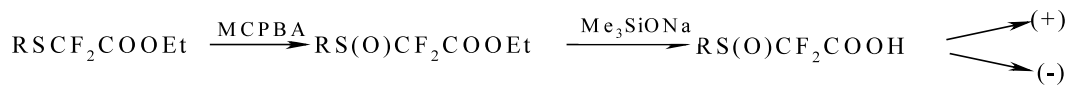
Tetrahedron Letters 43 (2002) 2945

Synthesis and optical activity of difluoro(organylsulfinyl)acetic acids and their esters

Andrej V. Matsnev, Nataliya V. Kondratenko, Yurii L. Yagupolskii* and Lev M. Yagupolskii

Institute of Organic Chemistry, National Academy of Sciences of Ukraine, 5 Murmanskaya St., 02094 Kiev, Ukraine

Representatives of a new type of optically active sulfur compound, arylsulfinyldifluoroacetic acids, have been prepared.



Tetrahedron Letters 43 (2002) 2949

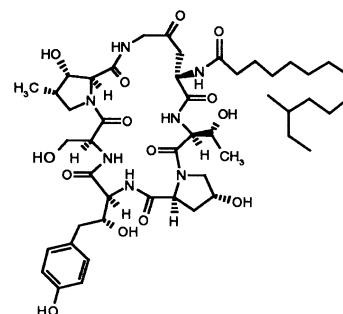
Synthesis of new echinocandin derivatives via a diol-keto transposition

Jozsef Aszodi,^a Patrick Fauveau,^a Dominique Melon-Manguer,^a Eberhard Ehlers^b and Laurent Schio^{a,*}

^aMedicinal Chemistry, Aventis Pharma, 102 Route de Noisy, F-93235 Romainville Cedex, France

^bProcess Development Biochemistry, Aventis Pharma, Biologika Süd, H780, D-65956 Frankfurt am Main, Germany

The synthesis of an original carbonyl analogue of deoxymulundocandin has been realised via a new transposition reaction performed in smooth acid conditions.

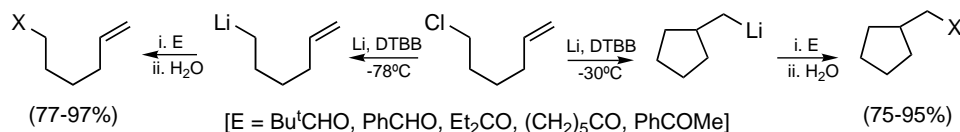


Tetrahedron Letters 43 (2002) 2953

DTBB-catalysed lithiation of 6-chloro-1-hexene and related systems: synthetically useful temperature-dependent behaviour

Miguel Yus,* Rosa Ortiz and Fernando F. Huerta

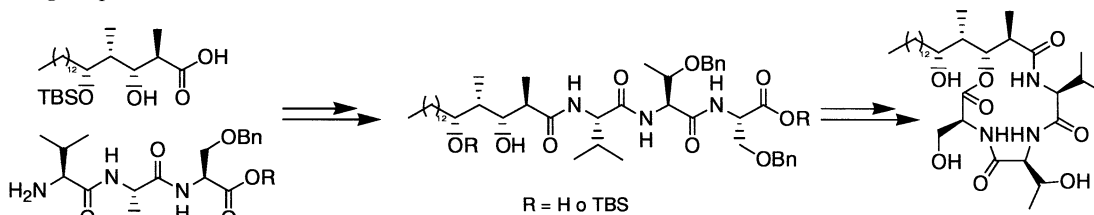
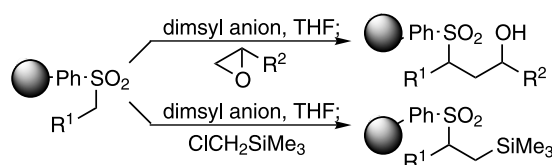
Departamento de Química Orgánica, Facultad de Ciencias, Universidad de Alicante, Apdo. 99, 03080 Alicante, Spain



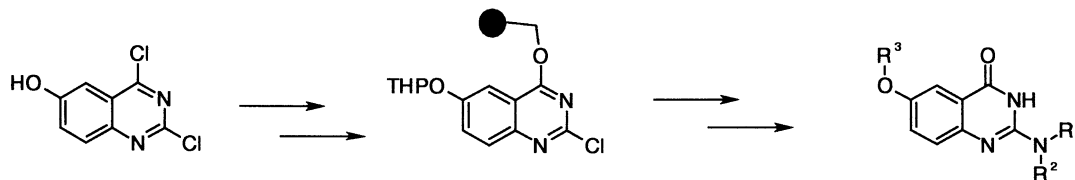
Tetrahedron Letters 43 (2002) 2957

A macrolactonization approach to the stevastelins

Francisco Sarabia,* Samy Chammaa and F. Jorge López-Herrera

Departamento de Bioquímica, Biología Molecular y Química Orgánica, Facultad de Ciencias, Universidad de Málaga, 29071 Málaga, Spain**Dimethyl anion in the monoalkylation of solid-phase alkyl sulfones**Wei-Chieh Cheng,^a Chu-Chung Lin^b and Mark J. Kurth^{a,*}^a*Department of Chemistry, University of California, One Shields Avenue, Davis, CA 95616-5295, USA*^b*Taigen Biotechnolog 7F, 138 Shin Ming Rd., Neihu Dist., Taipei 114, Taiwan, ROC***Novel solid-phase synthesis of 2,6-disubstituted 4(3H)-quinazolinones for combinatorial library generation**

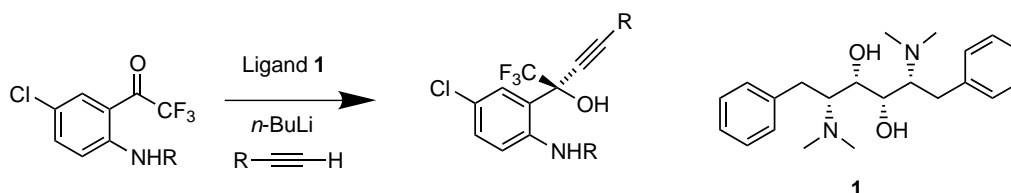
Csaba Wéber,* Attila Bielik, Györgyi I. Szendrei and István Greiner

Chemical and Biotechnological Research and Development, Gedeon Richter Ltd, PO Box 27, H-1475 Budapest, Hungary

THP: tetrahydropyran-2-yl, R³: primary or secondary alkyl,
 R¹, R²: primary alkyl or NR¹R²: cyclic amine

Enantioselective alkylation of a prochiral ketone catalyzed by C₂-symmetric diamino diols

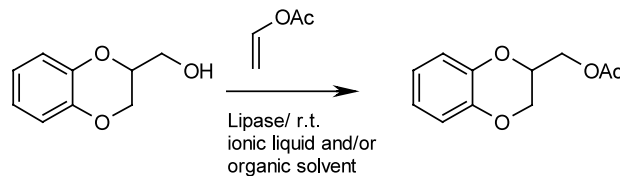
Biao Jiang* and Yan Feng

The State Key Laboratory of Organometallic Chemistry, Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, 354 Fengling Road, Shanghai 200032, PR China

Lipase-catalysed transesterification in ionic liquids and organic solvents: a comparative study

Susheel J. Nara, Jitendra R. Harjani and Manikrao M. Salunkhe*

Department of Chemistry, The Institute of Science, 15-Madam Cama Road, Mumbai 400 032, India

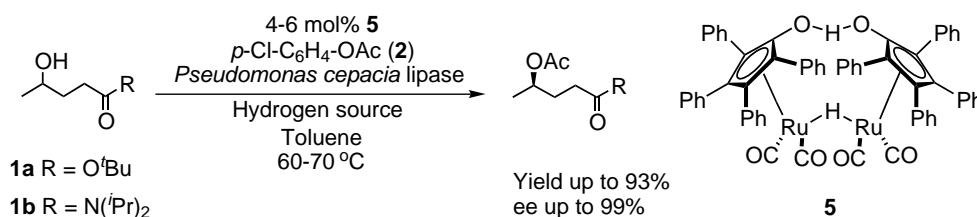


Dynamic kinetic resolution of γ -hydroxy acid derivatives

Ann-Britt L. Runmo,^a Oscar Pàmies,^a Kurt Faber^b and Jan-E. Bäckvall^{a,*}

^aDepartment of Organic Chemistry, Arrhenius Laboratory, Stockholm University, SE-10691 Stockholm, Sweden

^bDepartment of Organic & Bioorganic Chemistry, University of Graz, A-8010 Graz, Austria

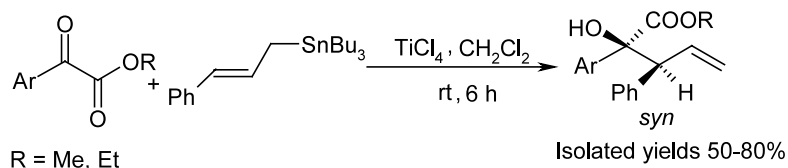


Highly diastereoselective TiCl₄ mediated addition of (*E*)-cinnamyl(tributyl)tin to α -keto esters

Deevi Basavaiah* and Bandaru Sreenivasulu

School of Chemistry, University of Hyderabad, Hyderabad 500 046, India

High *syn*-(87–98%) diastereoselective addition of (*E*)-cinnamyltin to α -keto esters under the influence of titanium tetrachloride, leading to the isolation of pure alkyl *syn*-2-aryl-2-hydroxy-3-phenylpent-4-enoates in 50–80% yields, is described.



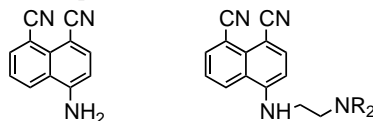
4-Amino-1,8-dicyanonaphthalene derivatives as novel fluorophore and fluorescence switches: efficient synthesis and fluorescence enhancement induced by transition metal ions and protons

Xuhong Qian^{a,*} and Yi Xiao^{a,b}

^aState Key Laboratory of Fine Chemicals, Dalian University of Technology, PO Box 40, Dalian 116012, China

^bShanghai Key Laboratory of Chemical Biology, PO Box 544, Shanghai 200237, China

These fluorosensors containing a novel fluorophore exhibit considerable fluorescence enhancement in the presence of transition metal ions and H⁺.



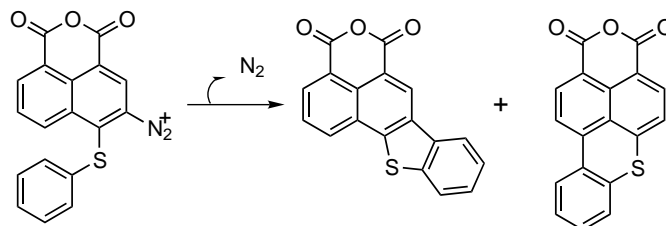
Synthesis and properties of benzothioxanthene dicarboximide hydroperoxide: an efficient 'time-resolved' DNA photocleaver with long-wavelength

Xuhong Qian,^{a,*} Ping Mao,^{b,c} Wei Yao^{b,c} and Xiangfeng Guo^a

^aState Key Laboratory of Fine Chemicals, Dalian University of Technology, PO Box 40, 158 Zhongshan Road, Dalian 116012, China

^bInstitute of Pesticides and Pharmaceuticals, East China University of Science and Technology, Shanghai 200237, China

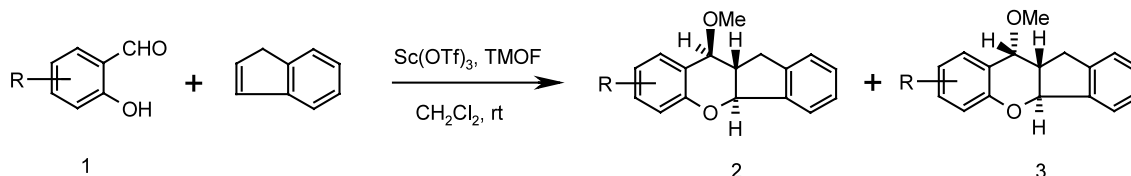
^cShanghai Key Laboratory of Chemical Biology, PO Box 544, Shanghai 200237, China



A facile synthesis of indeno[1,2-b]chromanes catalyzed by scandium triflate

Jhillu S. Yadav,^{*} B. V. Subba Reddy, Celine Parisse, Peter Carvalho and T. Prabhakar Rao

Division of Organic Chemistry, Indian Institute of Chemical Technology, Hyderabad 500007, India



Synthesis of a new chelating agent derived from phenylenediamine for application in radioimmunotherapy

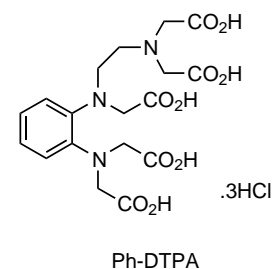
Sébastien G. Gouin,^{a,b} Jean-François Gestin,^{b,c} Alain Reliquet,^a Jean Claude Meslin^a and David Deniaud^{a,*}

^aLaboratoire de Synthèse Organique, UMR CNRS 6513, Faculté des Sciences et des Techniques, 2 rue de la Houssinière, BP 92208, 44072 Nantes Cedex, France

^bINSERM U463, Chimie des bioconjugués, 9 quai Moncoussu, 44093 Nantes Cedex, France

^cCHELATEC SAS, Institut de Biologie, 9 quai Moncoussu, 44093 Nantes Cedex, France

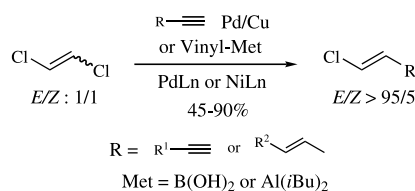
We report the synthesis of a precursor of a bifunctional chelating agent possessing an aromatic rigid skeleton.



Stereoselective synthesis of (E)-chloroenynes or (E,E)-chlorodienes starting from a stereoisomeric mixture of 1,2-dichloroethylenes

Mouâd Alami,^{*} Jean-François Peyrat and Jean-Daniel Brion

Laboratoire de Chimie Thérapeutique associé au CNRS (ESA 8076, BIOCIS), Faculté de Pharmacie, rue J.B. Clément, 92296 Châtenay Malabry Cedex, France



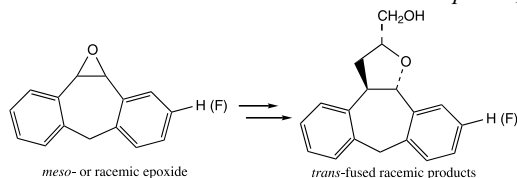
Stereoselective synthesis of *trans*-fused tetrahydrofuran derivatives of 5*H*-dibenzo[*a,d*]cycloheptene

Tetrahedron Letters 43 (2002) 3011

Frans Compennolle,^a Hua Mao,^a Abdellah Tahri,^a Tomasz Kozlecki,^a Erik Van der Eycken,^a Bart Medaer^b and Georges J. Hoornaert^{a,*}

^aLaboratorium voor Organische Synthese, K.U.Leuven, Celestijnenlaan 200F, B-3001 Leuven, Belgium

^bDrug Evaluation, Johnson & Johnson Pharmaceutical Research & Development, Turnhoutse steenweg 30, B-2340 Beerse, Belgium



Amination of α,β -unsaturated (2-trimethylsilylmethyl) carboxylic esters

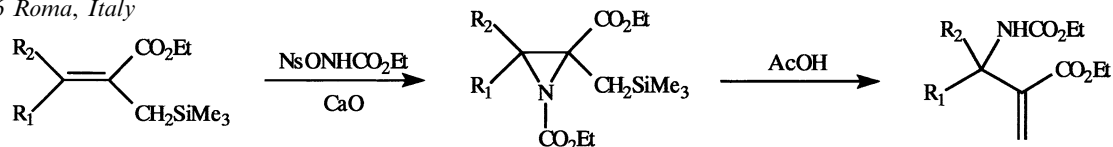
Tetrahedron Letters 43 (2002) 3017

Tecla Gasperi,^a M. Antonietta Loreto,^{a,b,*} Paolo A. Tardella^a and Augusto Gambacorta^c

^aDipartimento di Chimica, Università 'La Sapienza', P. le Aldo Moro 5, I-00185 Roma, Italy

^bIstituto C.N.R. di Chimica Biomolecolare – Sezione Roma – Dipartimento di Chimica, Università 'La Sapienza', Roma, Italy

^cDipartimento di Ingegneria Meccanica e Industriale, Università degli studi Roma Tre, via della Vasca Navale 79, I-00146 Roma, Italy

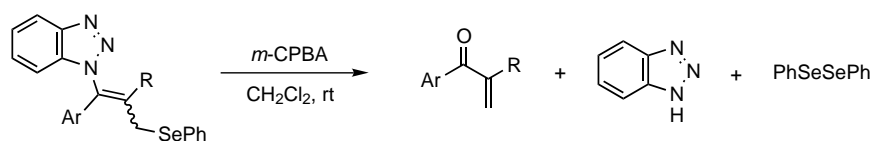


Synthesis of 2-alkyl (and aryl)-1-aryl-2-propen-1-ones via *m*-CPBA mediated oxidation of γ -(benzotriazol-1-yl)allylic selenides

Tetrahedron Letters 43 (2002) 3021

Taehoon Kim and Kyongtae Kim*

School of Chemistry and Molecular Science, Seoul National University, Seoul 151-742, South Korea



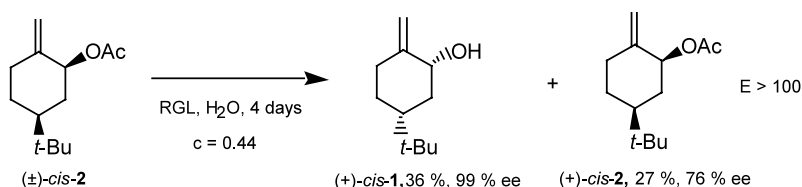
Diastereoselective and enantioselective lipase-catalyzed hydrolysis of stereoisomeric 2-methylene, 5-*t*-butylcyclohexyl acetates

Tetrahedron Letters 43 (2002) 3025

Chahra Bidjou,^a Louisa Aribi-Zouioueche^{a,*} and Jean-Claude Fiaud^{b,*}

^aDepartment of Organic Chemistry, University of Annaba, B.P. 12, 23000 Annaba, Algeria

^bInstitut de Chimie Moléculaire d'Orsay, Université Paris-Sud, F-91405 Orsay cedex, France

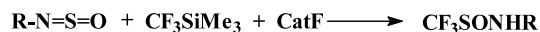


Perfluoroalkylation of heterocumulenes with trimethyl(perfluoroalkyl)silanes in the presence of fluoride ions: synthesis of perfluoroalkanesulfinyl amides from *N*-organylsulfinyl amines

Yurii L. Yagupolskii,^{a,*} Nataliya V. Kirij,^a Aleksey V. Shevchenko,^a Wieland Tyrra^b and Dieter Naumann^b

^a*Institute of Organic Chemistry, National Academy of Sciences of Ukraine, 5 Murmanskaya St., UA-02094 Kiev 94, Ukraine*

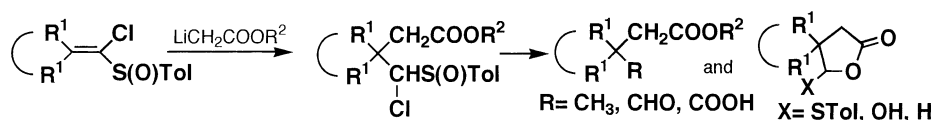
^b*Institut für Anorganische Chemie, Universität zu Köln, Greinstr. 6, D-50939 Cologne, Germany*



A novel synthesis of carboxylic acid derivatives having a quaternary carbon at 3-position and functional groups at 4-position from 1-chlorovinyl *p*-tolyl sulfoxides and acetic acid esters

Tsuyoshi Satoh,^{*} Shimpei Sugiyama and Hiroyuki Ota

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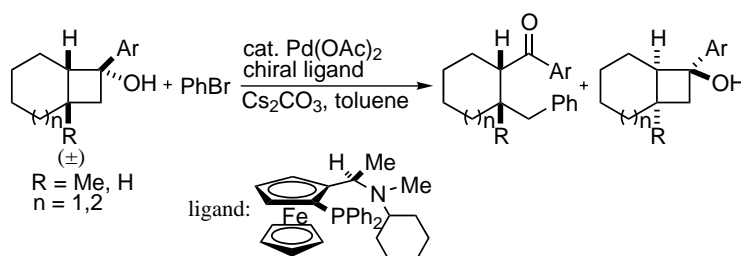


Palladium-catalyzed kinetic resolution of *tert*-cyclobutanols via C–C bond cleavage

Takahiro Nishimura, Satoshi Matsumura, Yasunari Maeda and Sakae Uemura^{*}

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Palladium-catalyzed kinetic resolution of *tert*-cyclobutanols with bromobenzene affords chiral ketones via C–C bond cleavage as well as chiral *tert*-cyclobutanols with moderate to good enantioselectivity.

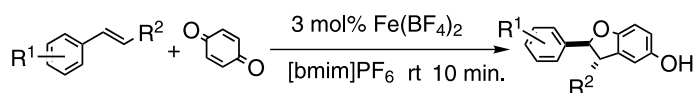


Cycloaddition of styrene derivatives with quinone catalyzed by ferric ion; remarkable acceleration in an ionic liquid solvent system

Hiroyuki Ohara,^b Hiromi Kiyokane^b and Toshiyuki Itoh^{a,*}

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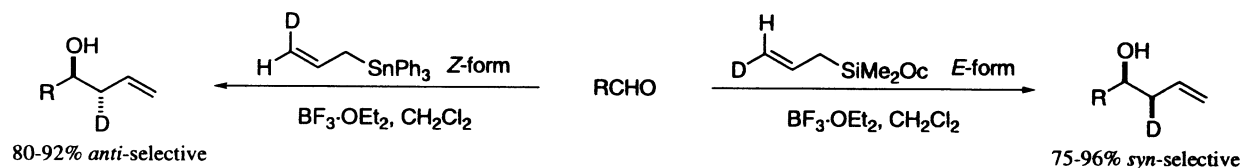
^b*Department of Chemistry, Faculty of Education, Okayama University, Okayama 700-8530, Japan*



Stereospecific formation of deuterated homoallyl alcohols by Lewis acid-promoted reactions of allyltin and allylsilicon reagents toward aldehydes

Yutaka Nishigaichi* and Akio Takuwa

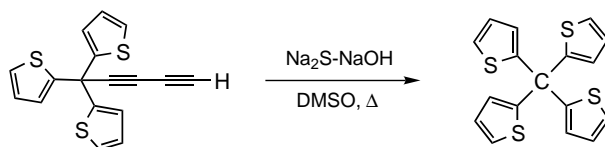
Department of Material Science, Faculty of Science and Engineering, Shimane University, 1060 Nishikawatsu-cho, Matsue, Shimane 690-8504, Japan



Synthesis of tetrakis(2-thienyl)methane

Kouzou Matsumoto, Hiromichi Nakaminami, Mitsufumi Sogabe, Hiroyuki Kurata and Masaji Oda*

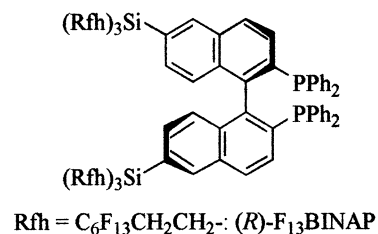
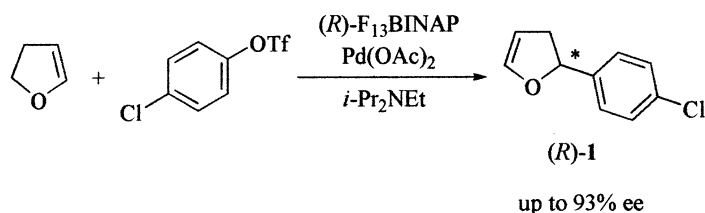
Department of Chemistry, Graduate School of Science, Osaka University, Toyonaka 560-0043, Japan



Preparation of a fluoros chiral BINAP and application to an asymmetric Heck reaction

Yutaka Nakamura,* Seiji Takeuchi,* Songlin Zhang, Kazuo Okumura and Yoshiaki Ohgo

Niigata College of Pharmacy, 5-13-2 Kamishin'ei cho, Niigata 950-2081, Japan

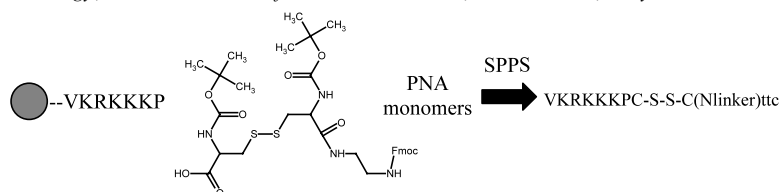


Optimised solid phase synthesis of a cystine-linked peptide-PNA chimera

Enrico Millo,^a Raffaele Nicolai,^a Sonia Scarfi,^a Carlo Scapolla,^a Barbara Biasotti,^b Umberto Benatti^a and Gianluca Damonte^{a,*}

^aDepartment of Experimental Medicine, Section Biochemistry, University of Genoa, Viale Benedetto XV, 1, 16132 Genoa, Italy

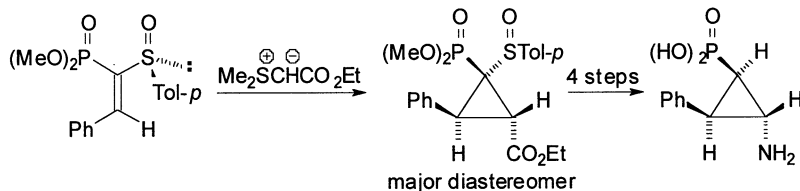
^bLaboratory of Experimental Oncology, National Institute for Cancer Research, 16132 Genoa, Italy



Asymmetric cyclopropanation of chiral (1-dimethoxyphosphoryl-2-phenyl)vinyl *p*-tolyl sulfoxide: a new synthesis of enantiomerically pure 2-amino-3-phenyl-1-cyclopropane-phosphonic acid—a constrained analog of phaclofen

Wanda H. Midura and Marian Mikołajczyk*

Centre of Molecular and Macromolecular Studies, Department of Heteroorganic Chemistry, Polish Academy of Sciences, 90-363 Łódź, Sienkiewicza 112, Poland

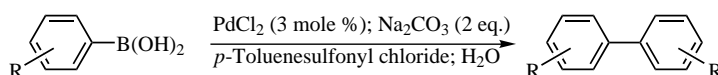


Ligandless palladium chloride-catalyzed homo-coupling of arylboronic acids in aqueous media

George W. Kabalka* and Lei Wang

Departments of Chemistry and Radiology, The University of Tennessee, Knoxville, TN 37996-1600, USA

In the presence of *p*-toluenesulfonyl chloride, ligandless palladium chloride catalyzes the homo-coupling of arylboronic acids to afford symmetrical biaryls in excellent yields.



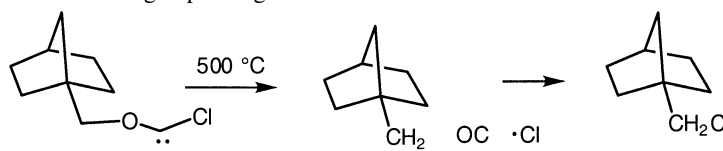
Fragmentation of alkoxychlorocarbenes in the gas phase

Michael E. Blake,^a Maitland Jones, Jr.,^{a,*} Fengmei Zheng^b and Robert A. Moss^{b,*}

^aDepartment of Chemistry, Princeton University, Princeton, NJ 08544, USA

^bDepartment of Chemistry, Rutgers University, New Brunswick, NJ 08903, USA

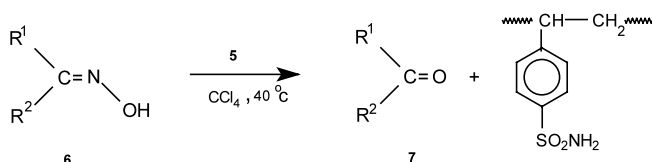
In contrast to photolysis or thermal decomposition in solution, which is dominated by ionic reactions, flash vacuum pyrolysis of alkylchlorodiazirines in the gas phase generates radicals.



Facile regeneration of carbonyl compounds from oximes using poly[4-vinyl-*N,N*-dichlorobenzenesulfonamide]

Ardeshir Khazaei* and Ramin Ghorbani Vaghei

Department of Chemistry, Faculty of Science, Bu-Ali Sina University, Hamadan, Iran

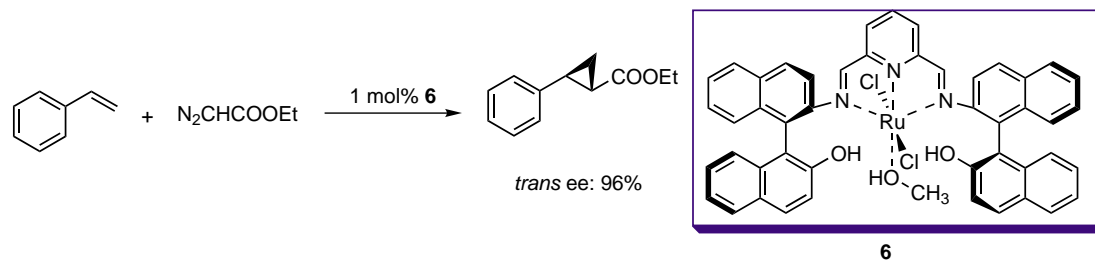


A new chiral ruthenium complex for catalytic asymmetric cyclopropanation

Tetrahedron Letters 43 (2002) 3075

Wenjun Tang, Xinquan Hu and Xumu Zhang*

Department of Chemistry, The Pennsylvania State University, University Park, PA 1680, USA



A new pentacyclic sulfated hydroquinone from the marine sponge *Haliclona* sp.

Tetrahedron Letters 43 (2002) 3079

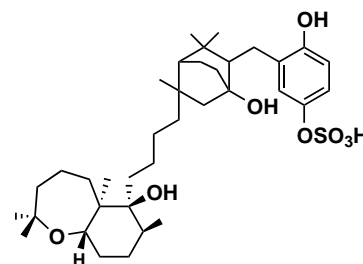
Heidi R. Bokesch,^b Amy C. Stull,^a Lewis K. Pannell,^c Tawnya C. McKee^a and Michael R. Boyd^{a,*}

^aMolecular Targets Drug Discovery Program, Center for Cancer Research, National Cancer Institute at Frederick, Frederick, MD 21702-1201, USA

^bIntramural Research Support Program, SAIC Frederick, Frederick, MD 21702-1201, USA

^cLaboratory of Bioorganic Chemistry, NIDDK, NIH, Bethesda, MD 20892, USA

A new pentacyclic sulfated hydroquinone, phuklona sulfate, has been isolated from the marine sponge *Haliclona* sp. Phuklona sulfate contains a novel triterpenoid carbon skeleton linked to a sulfated hydroquinone moiety.



Synthesis of diselenides and selenides from elemental selenium

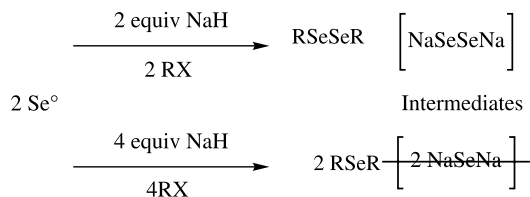
Tetrahedron Letters 43 (2002) 3083

Alain Krief^{a,*} and Michel Derock^{a,b}

^aLaboratoire de Chimie Organique de Synthèse, Département de Chimie, Facultés Universitaires Notre-Dame de la Paix, 61 rue de Bruxelles, Namur B-5000, Belgium

^bFonds pour la Formation à la Recherche dans l'Industrie et dans l'Agriculture, 5 Rue d'Egmont, Bruxelles B-1050, Belgium

Sodium hydride is able to reduce elemental selenium to sodium diselenide, but not to sodium selenide.

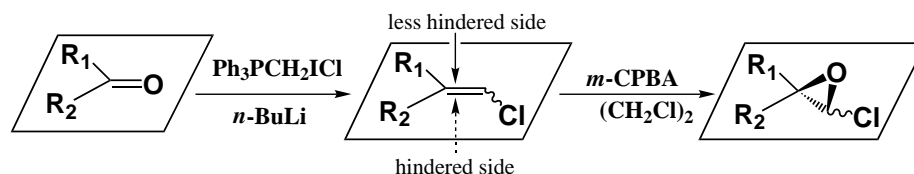


Improved synthetic method for preparing spiro α -chloroepoxides

Tetrahedron Letters 43 (2002) 3087

Ken-ichi Sato,* Takao Sekiguchi, Tetsuya Hozumi, Toshiya Yamazaki and Shoji Akai

Laboratory of Organic Chemistry, Faculty of Engineering, Kanagawa University, 3-27-1, Rokkakubashi, Kanagawa-ku, Yokohama 221-8686, Japan



Copper-promoted C–N bond cross-coupling with phenylstannane

Patrick Y. S. Lam,* Guillaume Vincent, Damien Bonne and
Charles G. Clark

Bristol-Myers Squibb Co., Experimental Station, PO Box 80500, Wilmington, DE 19880-0500, USA

Copper-promoted C–N bond cross-coupling of NH-containing substrates with phenylstannane at room temperature was accomplished with the addition of TBAF.

