Tetrahedron Letters 43 (2002) 1359



Palladium-catalyzed intramolecular arylation of an anilide enolate, application to an efficient formal total synthesis of physovenine

Tetrahedron Letters 43 (2002) 1363

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An expedient formal total synthesis of the calabar alkaloid physovenine was reported. The key step involves an oxindole synthesis via palladium-catalyzed intramolecular arylation of o-bromoanilide.



Facile selective cleavage of a myo-inositol trans-isopropylidene acetal in the presence of a *cis*-isopropylidene acetal

Tetrahedron Letters 43 (2002) 1367

K. S. Ravikumar and David Farquhar*

Department of Experimental Therapeutics, The University of Texas M.D. Anderson Cancer Center, Houston, TX 77030, USA



R = Bn, Ts, TBDPSi, Bz R' = Bn, Bz

Preparation of active esters on solid support for aqueous-phase

Tetrahedron Letters 43 (2002) 1369

Andrew D. Corbett and James L. Gleason*

peptide couplings

Department of Chemistry, McGill University, 801 Sherbrooke W., Montreal, QC, Canada H3A 2K6

TentaGel supported tetrafluorophenyl esters are found to be useful for the formation of dipeptides in water. Minimal hydrolysis of the active ester is observed.



Total synthesis of bengamide E

Tetrahedron Letters 43 (2002) 1373

Wenming Liu,* Joanna M. Szewczyk, Liladhar Waykole, Oljan Repič and Thomas J. Blacklock

Process R&D, Chemical and Analytical Development, Novartis Institute for Biomedical Research, One Health Plaza, East Hanover, NJ 07936, USA

A total synthesis of bengamide E is reported. The synthesis includes the utilization of D-tartrate as the chiral building block, construction of the E-olefin by the Julia protocol, an *anti*-aldol reaction to generate C-2 and C-3 stereocenters, and coupling of the thioester with caprolactam hydrochloride using sodium 2-ethylhexanoate.





The use of Sonogashira coupling for the synthesis of modified uracil peptide nucleic acid

Tetrahedron Letters 43 (2002) 1381

Robert H. E. Hudson,* Ge Li and Joseph Tse

Department of Chemistry, The University of Western Ontario, London, Ontario, Canada N6A 5B7



Facile access to aryltellurium compounds from arylboronic acids

Tetrahedron Letters 43 (2002) 1387

Aaron R. Clark,^a Rashmi Nair,^a Frank R. Fronczek^b and Thomas Junk^{a,*}

^aDepartment of Chemistry, University of Louisiana at Monroe, Monroe, LA 71203, USA ^bDepartment of Chemistry, Louisiana State University, Baton Rouge, LA 70803, USA

Arylboronic acids react with tellurium tetrachloride to generate aryltellurium trichlorides, which were reduced to diaryl ditellurides without prior isolation. Similarly, asymmetrical diaryl tellurides are accessible from arylboronic acids and aryltellurium tribromides.





Lower primary alkanols and their esters in a Ritter-type reaction with nitriles. An efficient method for obtaining N-primary-alkyl amides

Mikhail Y. Lebedev* and Mark B. Erman

Millennium Specialty Chemicals, PO Box 389, Jacksonville, FL 32201, USA

N-Primary-alkyl amides were obtained by a Ritter-type reaction of nitriles with lower primary alkanols or their esters in the presence of acids.

Various aromatic nitriles are reduced to the corresponding aldehydes by platinum(IV) oxide in aqueous formic acid with

yields ranging from 76 to 94%. This mild method may be generally applied to multi-step organic synthesis.

Tetrahedron Letters 43 (2002) 1401

Tetrahedron Letters 43 (2002) 1397

Tetrahedron Letters 43 (2002) 1395

The structure of desmosdumotin C (1), a novel cytotoxic compound isolated

Desmosdumotin C, a novel cytotoxic principle from **Desmos dumosus**

Jiu-Hong Wu,^{a,*} Andrew T. McPhail,^b Kenneth F. Bastow,^c Hiroaki Shiraki,^c Junko Ito^c and Kuo-Hsiung Leec.*

^aDepartment of Pharmacy, 306 Hospital of PLA, Beijing 100101, China ^bDepartment of Chemistry, Paul M. Gross Chemical Laboratory, Duke University, MeO Durham, North Carolina 27708, USA

^cNatural Products Laboratory, School of Pharmacy, University of North Carolina, Chapel Hill, North Carolina 27599, USA

from the roots of Desmos dumosus, has been established from spectral data and X-ray crystallographic analysis.

A novel and convenient transformation of nitriles to aldehydes

MedImmune, Inc., 35 West Watkins Mill Road, Gaithersburg, MD 20878, USA

Fred Xi,* Fred Kamal and Mark A. Schenerman



Tetrahedron Letters 43 (2002) 1391

Electrochemical reduction of diheteroaryl-1,2-diketones in the presence of carbonimidoyl dichlorides. First synthesis of 2-arylimino-4,5-di-2-furyl-1,3-dioxoles and (E)-1,2-di-2-furylvinylene bis(N-arylchloroformimidates)

Antonio Guirado,* Andrés Zapata, Raquel Andreu and Bruno Martiz

Departamento de Química Orgánica, Facultad de Química, Universidad de Murcia, Campus de Espinardo, 30071 Murcia, Apartado 4021, Spain





^aUniversité Louis Pasteur/CNRS, Institut Le Bel, 4 rue Blaise Pascal, F-67070 Strasbourg Cedex, France ^bDepartament de Bioquímica i Biologia Molecular, Facultat de Química, Universitat de Barcelona, Martí i Franquès 1, ES-08028 Barcelona, Spain



Tetrahedron Letters 43 (2002) 1417Tetrahedron Letters 43 (2002) 1417Tetrahedron Letters 43 (2002) 1417aminopropanoic acid suitably protected for peptide synthesisWang-Qing Liu, Catherine Olszowy, Laurent Bischoff and Christiane Garbay*Laboratoire de Pharmacochimie Moléculaire et Structurale INSERM et CNRS, Faculté des Sciences Pharmaceutiques et Biologiques, 4, avenue de l'Observatoire, 75270 Paris Cedex 06, FranceCedex 06, FranceColspan="2">Colspan="2">OOOOColspan="2">OOOOOColspan="2">Cedex 06, France



C-Backbone branched peptides via reductive amination of cyanomethyleneamino pseudopeptides

Instituto de Química Médica (CSIC), Juan de la Cierva 3, E-28006 Madrid, Spain

Susana Herrero, M. Luisa Suárez-Gea, M. Teresa García-López and Rosario Herranz*



Laboratoire de Pharmacognosie, associé au CNRS (BIOCIS), Université Paris-Sud, Faculté de Pharmacie, rue Jean-Baptiste Clément, 92296 Châtenay-Malabry, France

Acaterin was prepared via a Baylis–Hillman reaction between lactone 2 and octanal. This is the first example of a Baylis–Hillman reaction with α , β -unsaturated lactones.













The isolation and synthesis of two novel *N*-acetyl glucosamine derivatives from *Dictyostelium* cellular slime molds which exhibit neurite outgrowth activity

Tetrahedron Letters 43 (2002) 1477

Haruhisa Kikuchi, Jun Komiya, Yoshinori Saito, Jun-ichi Sekiya, Shigeyoshi Honma, Norimichi Nakahata and Yoshiteru Oshima*

Graduate School of Pharmaceutical Sciences, Tohoku University, Aoba-yama, Aoba-ku, Sendai 980-8578, Japan

Dictyoglucosamine A (1) $R = CH_3CH_2$ Dictyoglucosamine B (2) $R = (CH_3)_2CHCH_2$



A novel photoreaction of 1,2-diarylcyclopropanes with 9-cyanophenanthrene: the formation of (3+2) photocycloadducts Hajime Maeda, Yasuo Miyata and Kazuhiko Mizuno* Department of Applied Chemistry, Graduate School of Engineering, Osaka Prefecture University, 1-1 Gakuen-cho, Sakai, Osaka 599-8531, Japan $\int (CN + An^{V} - An - benzene + An^{V} - An^{V}$



Oxiduati : Third

Tetrahedron Letters 43 (2002) 1495

Synthesis of tetrahydrofurans by regio- and stereoselective cyclization of epoxyalcohols using magnesium halide

Michinori Karikomi,* Shigeru Watanabe, Yukino Kimura and Tadao Uyehara Department of Applied Chemistry, Faculty of Engineering, Utsunomiya University, Yoto 7-1-2, Utsunomiya 321-8585, Japan



Synthesis of different ring-size heterocycles from the same propargyl alcohol derivative by ligand effect on Pd(0)

Tetrahedron Letters 43 (2002) 1499

Yuji Kozawa and Miwako Mori*

Graduate School of Pharmaceutical Sciences, Hokkaido University, Sapporo 060-0812, Japan



Electroorganic synthesis under solvent-free conditions. Highly regioselective anodic monofluorination of cyclic ethers, lactones, and a cyclic carbonate

Tetrahedron Letters 43 (2002) 1503

Masaru Hasegawa, Hideki Ishii and Toshio Fuchigami*

Department of Electronic Chemistry, Tokyo Institute of Technology, Nagatsuta, Midori-ku, Yokohama 226-8502, Japan



Synthesis of selenol esters: palladium-catalyzed coupling of phenyl tributylstannyl selenide with aryl iodides and carbon monoxide



Yutaka Nishiyama,* Keiji Tokunaga, Hiroaki Kawamatsu and Noboru Sonoda*

Department of Applied Chemistry, Faculty of Engineering, Kansai University, 3-3-35 Yamate Chou Suita, Osaka 564-8680, Japan



Asymmetric cyclization-carbonylation of cyclic-2-methyl-2propargyl-1,3-diols

Tetrahedron Letters 43 (2002) 1511

Keisuke Kato,^{a,*} Maki Tanaka,^a Yasuhiro Yamamoto^b and Hiroyuki Akita^{a,*}

^aSchool of Pharmaceutical Sciences, Toho University, 2-2-1 Miyama, Funabashi, Chiba 274-8510, Japan ^bDepartment of Chemistry, Faculty of Science, Toho University, 2-2-1 Miyama, Funabashi, Chiba 274-8510, Japan

Palladium-catalyzed asymmetric cyclization-methoxycarbonylation of cyclic-2-methyl-2-propargyl-1,3-diols 1 under mild conditions afforded (*E*)-bicyclic- β -alkoxyacrylates 2 in good yields with moderate enantioselectivity.



Combination of silyl carbamate and amino acid fluoride for solid-phase peptide synthesis

Kimitoshi Sakamoto,^a Yoshiaki Nakahara^{a,b} and Yukishige Ito^{a,*}

Tetrahedron Letters 43 (2002) 1515

^aRIKEN (The Institute of Physical and Chemical Research), 2-1 Hirosawa, Wako-shi, Saitama 351-0198, Japan ^bDepartment of Applied Biochemistry, Tokai University, Hiratsuka-shi, Kanagawa 259-1292, Japan

$$^{i}Pr_{3}Si-OTN + N + N + F$$

 $OR = 1$ R^{2} $R^{$



Highly chemoselective Pummerer reactions of sulfinyldiacetic

Tetrahedron Letters 43 (2002) 1519

The first use of chiral oxazoline ligands in the highly
enantioselective diethylzinc addition to diphenylphosphinoyl iminesTetrahedron Letters 43 (2002) 1535Xiaomei Zhang,^a Wenqing Lin,^a Liuzhu Gong,^{a,*} Aiqiao Mi,^{a,*} Xin Cui,^a Yaozhong Jiang,^aMichael C. K. Choi^b and Albert S. C. Chan^b

^aUnion Laboratory of Asymmetric Synthesis, Chengdu Institute of Organic Chemistry, Chinese Academy of Sciences, Chengdu 610041, China

^bOpen Laboratory of Chirotechnology and Department of Applied Biology and Chemical Technology, The Hong Kong Polytechnic University, Hong Kong, China







A mild and efficient method for cleavage of C=N using $Mg(HSO_4)_2$ in the presence of wet SiO₂

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^aDepartment of Chemistry, Faculty of Science, Guilan University, Rasht, Iran

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

^cOrganic Polymer Chemistry Research Laboratory, College of Chemistry, Isfahan University of Technology, Isfahan, Iran ^dDepartment of Chemistry, Isfahan University, Isfahan, Iran

An efficient and convenient conversion of oximes, hydrazones and semicarbazones to the corresponding carbonyl compounds with $Mg(HSO_4)_2$ in the presence of wet SiO₂ is reported.

$$-\overset{I}{C} = \overset{N-Y}{\underset{Y= \text{ OH, NHPh, NHC}_{6}H_{4}-p-NO_{2}, \text{ NHCONH}_{2}}{\overset{Mg(HSO_{4})_{2}/\text{wet SiO}_{2}}} - \overset{I}{C} = O$$

Direct Cu(I)-catalysed coupling of a carborane to a *meso*-tetraphenylporphyrin

Christophe Frixa,^a Mary F. Mahon,^b Andrew S. Thompson^a and Michael D. Threadgill^{a,*} ^aDepartment of Pharmacy & Pharmacology, University of Bath, Bath BA2 7AY, UK ^bDepartment of Chemistry, University of Bath, Bath BA2 7AY, UK

Coupling of Cu-carborane with iodo-TPPZn gives the carboranylporphyrin.



Tetrahedron Letters 43 (2002) 1557





Synthesis and Ag⁺-catalyzed cyclization of 2,3-dienamides

Tetrahedron Letters 43 (2002) 1569

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Organic Division I, Indian Institute of Chemical Technology, Hyderabad 500 007, India



The use of enantiomerically pure *N*-sulfinimines in asymmetric Baylis–Hillman reactions

Tetrahedron Letters 43 (2002) 1577

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11-(Tetrahydro-3 and 4-pyridinyl)dibenzo[*b*,*e*][1,4]diazepines undergo novel rearrangements on treatment with concentrated HBr

Tetrahedron Letters 43 (2002) 1583

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Convenient synthesis of disulfide substrates for trypanothione reductase using polymer-supported reagents

Tetrahedron Letters 43 (2002) 1587

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Coupling of the cystine derivative with 1 using polymer-supported reagents gave disulfides in high yields.

