4-Chromenesulphones: synthesis and transformation to isoflavonoid models

Tetrahedron Letters 43 (2002) 6893

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Universidade Federal do Rio de Janeiro, Núcleo de Pesquisas de Produtos Naturais, Ilha da Cidade Universitária, CCS, bloco H, Rio de Janeiro, RJ, 21941-590, Brazil

Isoflavonoid models (carrying basic skeleton 1) were constructed from chromenes 2 through novel sulphones 3.

Simultaneous assembly of the $\beta\mbox{-lactam}$ and thiazole moiety by a new multicomponent reaction

Tetrahedron Letters 43 (2002) 6897

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^bMorphochem AG, Gmunder Str. 37-37a, 81379 München, Germany

How does it work? Three acyclic starting materials assemble according to a molecular program to form a highly strained β -lactam ring as well as an aromatic thiazole ring in the products. The increase in molecular complexity here is dramatic as in addition to the two rings, 2 C–N, 2 C–S and 1 C–C bonds are formed in a new 'one-pot' multicomponent reaction.

$$H_2N$$
 $COSH + R^2$
 CHO
 CHO

Structural correction of the 3-methylindole oxidatively-coupled dimer

Tetrahedron Letters 43 (2002) 6903

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^bCenter for Supramolecular Science and Department of Chemistry, University of Miami, Coral Gables, FL 33124-0431, USA

A novel four component one-pot access to pyrindines and tetrahydroquinolines

Tetrahedron Letters 43 (2002) 6907

N. A. M. Yehia, K. Polborn and T. J. J. Müller*

Organisch-Chemisches Institut, Ruprecht-Karls-Universität Heidelberg, Im Neuenheimer Feld 270, D-69120 Heidelberg, Germany

A coupling-isomerization-Stork-enamine alkylation-cyclocondensation sequence to dihydropyrindines and tetrahydroquinolines. Ar¹

Ar1 = electron deficient (het)aryl

Ar²= (het)aryl

Stereoselective formation of optically active 2-oxy-1,3-oxazolidin-4-ones and an efficient synthesis of optically active secondary 2-pyrrolidones

Yoji Omata, a Akikazu Kakehi, Masashi Shiraic and Akio Kamimuraa,*

^aDepartment of Applied Chemistry, Faculty of Engineering, Yamaguchi University, Ube 755-8611, Japan

^bDepartment of Chemistry and Material Engineering, Shinshu University, Nagano 380-8553, Japan

^cUbe Laboratory, Ube Industries Ltd, Ube 755-8633, Japan

Establishment of *Camellia sinensis* cell culture with high peroxidase activity and oxidative coupling reaction of dibenzylbutanolides

Tetrahedron Letters 43 (2002) 6915

Masumi Takemoto, a.* Youichi Aoshima, b Nikolay Stoynov and James Peter Kutnev

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^bShizuoka Agricultural Experiment Station, 678-1 Tomigaoka Toyoda-cho, Iwata-gun, Shizuoka 438-0803, Japan

^cDepartment of Chemistry, University of British Columbia, 2036 Main Mall, Vancouver, B.C., Canada V6T 1Z1

Peroxidase-catalyzed oxidative coupling of racemic 1a-c to cyclic products

(2a-c) proceeded with Camellia sinensis cell culture quantitatively in the absence of foreign hydrogen peroxide.

$\begin{array}{c} R_1 \\ R_2 \\ H_3CO \\ OH \\ (\pm)\textbf{-1a-c} \end{array} \qquad \begin{array}{c} R_1 \\ R_2 \\ R_2 \\ R_3 \\ R_2 \\ R_2 \\ OH \\ OH \\ (\pm)\textbf{-2a-c} \end{array}$

The first total synthesis and neurotrophic activity of clusiparalicoline A, a prenylated and geranylated biaryl from *Clusia paralicola*

Tetrahedron Letters 43 (2002) 6919

Shigeki Takaoka, Kousuke Nakade and Yoshiyasu Fukuyama* Institute of Pharmacognosy, Faculty of Pharmaceutical Sciences, Tokushima Bunri University, Tokushima 770-8514, Japan

Clusiparalicoline A, synthesized by the sequential Stille and Suzuki reactions, has been found to exhibit a potent neurite outgrowth promoting activity at 1.0 μ M in a primary culture of fetal rat cortical neurons.

Remarkable rate acceleration of the solvent-free Baeyer-Villiger reaction on the surface of NaHCO₃ crystals for sterically congested cyclic and acyclic ketones

Tetrahedron Letters 43 (2002) 6925

Takayuki Yakura, Tomoko Kitano, Masazumi Ikeda and Jun'ichi Uenishi* Kyoto Pharmaceutical University, Misasagi, Yamashina, Kyoto 607-8412, Japan

An efficient synthesis of (+)-aureol via boron trifluoride etheratepromoted rearrangement of (+)-arenarol

Tetrahedron Letters 43 (2002) 6929

Masahiko Nakamura, Akiyuki Suzuki, Mari Nakatani, Takamasa Fuchikami, Munenori Inoue and Tadashi Katoh*

Sagami Chemical Research Center, Hayakawa 2743-1, Ayase, Kanagawa 252-1193, Japan

Enzyme-mediated enantioselective hydrolysis of cyclic carbonates bearing an unsaturated substituent

Tetrahedron Letters 43 (2002) 6933

Kazutsugu Matsumoto, a,* Yasuhide Nakamura, Megumi Shimojoc and Minoru Hatanakab

^aDepartment of Chemistry, Meisei University, Hodokubo 2-1-1, Hino, Tokyo 191-8506, Japan

^bDepartment of Applied Chemistry and Biotechnology, Fukui University, Bunkyo 3-9-1, Fukui 910-8507, Japan

Department of Biosciences and Informatics, Keio University, Hiyoshi 3-14-1, Yokohama 223-8522, Japan

Pinacol rearrangement for constructing asymmetric centers adjacent to heterocycles

Tetrahedron Letters 43 (2002) 6937

Tomoichi Shinohara and Keisuke Suzuki*

Department of Chemistry, Tokyo Institute of Technology, and CREST, Japan Science and Technology Corporation (JST), O-okayama, Meguro-ku, Tokyo 152-8551, Japan

Versipelostatin, a novel GRP78/Bip molecular chaperone downregulator of microbial origin

Tetrahedron Letters 43 (2002) 6941

Hae-Ryong Park,^a Kazuo Furihata,^b Yoichi Hayakawa^a and Kazuo Shin-ya^{a,*}

^aInstitute of Molecular and Cellular Biosciences,

The University of Tokyo, Bunkyo-ku, Tokyo 113-0032, Japan

^bGraduate School of Agricultural and Life Sciences,

The University of Tokyo, Bunkyo-ku, Tokyo 113-8657, Japan

Oxathioacetalization, thioacetalization and transthioacetalization of carbonyl compounds by N-bromosuccinimide: selectivity and scope

Tetrahedron Letters 43 (2002) 6947

Ahmed Kamal,* Gagan Chouhan and Kaleem Ahmed

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

$$\overset{O}{\underset{H}{ }} \quad \text{or} \quad \overset{X}{\underset{X}{ }} \quad \overset{2\text{-mercaptoethanol/thiol/dithiol}}{\underset{NBS \ (cat.)}{ \text{CH}_2\text{Cl}_2, r.t.}} \quad \overset{O}{\underset{S}{ }} \quad \text{or} \quad \overset{SEt}{\underset{SEt}{ }} \quad \text{or} \quad \overset{S}{\underset{S}{ }} \overset{S}{\underset{n}{ }} \quad \overset$$

R = aryl, allyl, cyclohexyl; X = OMe, OEt, OAc; n = 1,2

Ytterbium(III) trifluoromethanesulfonate for specific activation of *n*-pentenyl orthoesters in the presence of acid-sensitive functionalities

Tetrahedron Letters 43 (2002) 6953

K. N. Jayaprakash, K. V. Radhakrishnan and Bert Fraser-Reid*

Natural Products and Glycotechnology Research Institute, Inc. (NPG), 4118 Swarthmore Road, Durham, NC 27706, USA

A general method for the highly diastereoselective, kinetically controlled alkylation of (+)-nopinone

Tetrahedron Letters 43 (2002) 6957

Kevin R. Campos,* Sandra Lee, Michel Journet, Jason J. Kowal, Dongwei Cai, Robert D. Larsen and Paul J. Reider

Department of Process Research, Merck Research Laboratories, PO Box 2000, Rahway, NJ 07065, USA

Solid-phase synthesis of phosphine-oxazoline peptides

Tetrahedron Letters 43 (2002) 6961

Scott R. Gilbertson* and Ping Lan

Department of Chemistry, Washington University, St. Louis, MO 63130, USA

Studies on the tetramerization of substituted monopyrroles to type I porphyrins

Tetrahedron Letters 43 (2002) 6967

C. Pichon-Santander and A. I. Scott*

Center for Biological NMR, Department of Chemistry,

Texas A&M University, PO Box 30012, College Station,

TX 77842-3012, USA

Tetramerization of pyrroles bearing two different electron-donating groups as substituents under slightly acidic conditions afforded porphyrins with a high ratio of type I isomer in good yields.

Efficient highly stereoselective synthesis of olefinic macrocyclic crown-formazans with the Z-configuration via ring-closure metathesis

Tetrahedron Letters 43 (2002) 6971

Yehia A. Ibrahim,* Haider Behbehani, Maher R. Ibrahim and Nada M. Abrar

Chemistry Department, Faculty of Science, Kuwait University, PO Box 5969, Safat 13060, Kuwait

Towards a total synthesis of guanacastepene A: construction of fully functionalized AB and BC ring segments

Tetrahedron Letters 43 (2002) 6975

Goverdhan Mehta,* Jayant D. Umarye and Vanessa Gagliardini

Department of Organic Chemistry, Indian Institute of Science, Bangalore 560 012, India

$$AcO$$
 A
 B
 C
 AcO
 A
 B
 C
 AcO
 A
 B
 C
 CH_3

The room temperature preparation of the 1-chloro-2,2-difluorovinylzinc reagent from HCFC-133a (CF_3CH_2Cl).

Tetrahedron Letters 43 (2002) 6979

The first ambient, high yield, one-flask preparation of α -chloro- β , β -difluorostyrenes

R. Anilkumar and Donald J. Burton*

Department of Chemistry, University of Iowa, Iowa City, IA 52242, USA

The first total synthesis of goniothalesdiol

Tetrahedron Letters 43 (2002) 6983

Matej Babjak, Peter Kapitán and Tibor Gracza*

Department of Organic Chemistry, Slovak University of Technology, Radlinského 9, SK-812 37 Bratislava, Slovakia

A total synthesis of goniothalesdiol and its epimer is described.

Cyclopropylboronic acid: synthesis and Suzuki cross-coupling reactions

Tetrahedron Letters 43 (2002) 6987

Debra J. Wallace* and Cheng-yi Chen

Department of Process Research, Merck Research Laboratories, PO Box 2000, Rahway, NJ 07065, USA

An efficient synthesis of cyclopropylboronic acid is reported. This compound undergoes efficient Suzuki-type cross-coupling reactions with a range of aryl and heteroaryl bromides.

$$\begin{array}{c|c} & \text{i. } (\text{MeO})_3 \text{B} \\ \hline & & \\ & \text{ii. } \text{H}_3 \text{O}^+ \\ \hline \end{array} \begin{array}{c|c} & \text{B(OH)}_2 \\ \hline & \text{Toluene, H}_2 \text{O, K}_3 \text{PO}_4 \\ \hline & \text{Pd(OAc)}_2, \text{Pcy}_3 \\ \hline \end{array}$$

The homologation of histidine

Tetrahedron Letters 43 (2002) 6991

Amit Kumar, Stephanos Ghilagaber, Jamie Knight and Peter B. Wyatt*

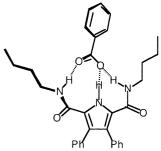
Department of Chemistry, Queen Mary, University of London, Mile End Road, London E1 4NS, UK

$$\begin{array}{c} \text{Mts} \\ \text{N} \\ \text{N} \\ \text{N} \\ \text{Mts} \\ \text{Mts} \\ \text{H} \\ \text{CO}_2 \\ \text{Mts} \\ \text{Protected histidine} \\ \end{array} \begin{array}{c} \text{(i) NaBH}_4 \text{/MeOH} \\ \text{(ii) NaCN} \\ \text{(iv) HBr/H}_2 \\ \text{O/PhOH} \\ \text{H}_2 \\ \text{N} \\ \text{Histidine } \beta \\ \text{-homologue} \\ \end{array} \begin{array}{c} \text{Me} \\ \text{Mts} \\ \text{Mts} \\ \text{Mts} \\ \text{Mts} \\ \text{Mts} \\ \text{N} \\ \text{N$$

Confirmation of a 'cleft-mode' of binding in a 2,5-diamidopyrrole anion receptor in the solid state

Tetrahedron Letters 43 (2002) 6995

Salvatore Camiolo, Philip A. Gale,* Michael B. Hursthouse and Mark E. Light *University of Southampton, Department of Chemistry, Highfield, Southampton SO17 1BJ, UK* The crystal structure of benzoate bound to a 2,5-diamidopyrrole anion receptor has been elucidated revealing the formation of a 'cleft conformation'.



Preparation of 1,2,4-trisubstituted imidazoles by ammonolysis of N-(2-oxoalkyl)oxazolinium salts

Tetrahedron Letters 43 (2002) 6997

Matthew P. John, Stephen A. Hermitage* and James R. Titchmarsh

GlaxoSmithKline, Chemical Development Division, Medicines Research Centre, Gunnels Wood Road, Stevenage, Hertfordshire SG1 2NY, UK

A variety of aryl and alkyl substituted imidazoles have been prepared by the ring opening of N-(2-oxoalkyl)oxazolinium salts with ammonia.

SmI₂-mediated facile one-pot preparation of 2,4-diarylquinolines from 3-aryl-2,1-benzisoxazoles

Tetrahedron Letters 43 (2002) 7001

Xuesen Fana,c and Yongmin Zhanga,b,*

^aDepartment of Chemistry, Zhejiang University (Campus Xixi), Hangzhou 310028, PR China

^bState Key Laboratory of Organometallic Chemistry, Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, Shanghai 200032, PR China ^cDepartment of Chemistry, Henan Normal University, Xinxiang 453002, PR China

THF, r.i. Z-C-C-C-(3) x 1

On treatment with SmI₂, 3-aryl-2,1-benzisoxazoles undergo reductive cleavage of the N–O bond leading to 2-aminobenzophenones in high yields upon protonation. If aryl methyl ketones are added to the reaction mixture prior to protonation, 2,4-diarylquinolines can be obtained in moderate yields.

Microwave assisted synthesis of 5-hydroxy-5-trichloromethyl-4,5-dihydroisoxazoles

Tetrahedron Letters 43 (2002) 7005

Marcos A. P. Martins,* Paulo Beck, Wilson Cunico, Claudio M. P. Pereira, Adilson P. Sinhorin, Rogério F. Blanco, Rodrigo Peres, Helio G. Bonacorso and Nilo Zanatta

Núcleo de Química de Heterociclos (NUQUIMHE), Departamento de Química, Universidade Federal de Santa Maria, 97.105-900 Santa Maria, RS, Brazil

A series of 13 5-hydroxy-5-trichloromethyl-4,5-dihydroisoxazoles have been synthesized by environmentally benign microwave (MW) induced techniques.

R\$\frac{2}{3} OR R\$\frac{2}{3} R^1

2a-m

CeCl₃·7H₂O: a novel reagent for the synthesis of 2-deoxysugars from D-glycals

Tetrahedron Letters 43 (2002) 7009

J. S. Yadav,* B. V. S. Reddy, K. Bhaskar Reddy and M. Satyanarayana

1a-m

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

Regioselective acylation at the 5- or 6-position of L-tryptophan derivatives

Yongwen Jiang^a and Dawei Ma^{b,*}

^aDepartment of Chemistry, Fudan University, Shanghai 200433, China

^bState Key Laboratory of Bioorganic and Natural Products Chemistry, Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, 354 Fenglin Lu, Shanghai 200032, China

$$\begin{array}{c|c} Z & H \\ \hline N & RCOCI/AICI_3 \\ \hline N & NHCO_2Me \end{array} \xrightarrow{RCOCI/AICI_3} \begin{array}{c} Z & H \\ \hline RCOCI/AICI_3 \\ \hline CICH_2CH_2CI \\ \hline ROC \end{array} \xrightarrow{N} \begin{array}{c} Z & H \\ \hline N & ROC \\ \hline N & NHCO_2Me \end{array} \xrightarrow{N} \begin{array}{c} Z & H \\ \hline N & NHCO_2Me \end{array}$$

Sterol methyl transferase. Evidence for successive C-methyl transfer reactions generating $\Delta^{24(28)}$ - and $\Delta^{25(27)}$ -olefins by a single plant enzyme

Tetrahedron Letters 43 (2002) 7017

Allen L. Dennis and W. David Nes*

Department of Chemistry and Biochemistry, Texas Tech University, Lubbock, TX 79409, USA

Recombinant soybean sterol methyl transferase converted [27- 13 C]-cycloartenol to (25R)-[27- 13 C]24(28)methylenecycloartanol and 24(28)-methylenecycloartanol to a mixture of (24E)- and (24Z)-24-ethylidenecycloartanol and 24 β -ethyl-25(27)-dehydrocycloartanol.

Nucleus (Nu)
$$^{13}C$$
 SMT AdoMet Nu ^{13}C H SMT AdoMet Nu SMT AdoMet Nu Nu

Semi-pinacol strategy for constructing B-ring of pradimicin—benanomicin antibiotics

Tetrahedron Letters 43 (2002) 7023

Ken Ohmori, Mitsuru Kitamura, Yuji Ishikawa, Hirohisa Kato, Mami Oorui and Keisuke Suzuki* Department of Chemistry, Tokyo Institute of Technology, and CREST, Japan Science and Technology Corporation (JST), O-okayama, Meguro-ku, Tokyo 152-8551, Japan

Reducing oligosaccharides via glycal assembly: on the remarkable stability of anomeric hydroxyl groups to global deprotection with sodium in liquid ammonia

Tetrahedron Letters 43 (2002) 7027

Ulrich Iserloh, a Vadim Dudkin, a Zhi-Guang Wang and Samuel J. Danishefsky b, a Zhi-Guang Wang and Samuel J. Danishefsky

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^bDepartment of Chemistry, Columbia University, Havemeyer Hall, New York, NY 10027, USA

Chelation-assisted intramolecular hydroacylation: synthesis of medium ring sulfur heterocycles

Tetrahedron Letters 43 (2002) 7031

Holly D. Bendorf,* Christine M. Colella, Elizabeth C. Dixon, Melissa Marchetti, Alicia N. Matukonis, Jeffrey D. Musselman and Tara A. Tiley

Department of Chemistry, Lycoming College, Williamsport, PA 17701, USA

$$\begin{array}{c} O \\ H \\ S \\ \end{array} \begin{array}{c} 0.05\text{-}0.1 \text{ eq Rh}(PPh_3)_3Cl} \\ \hline CH_2Cl_2, \text{ rt} \\ 54\text{-}92\%, \text{ } n = 1\text{-}2 \\ \end{array}$$

A novel one-pot synthesis of 3-acetyl- and 3-benzoylisoxazole derivatives using ammonium cerium nitrate (CAN)

Tetrahedron Letters 43 (2002) 7035

Ken-ichi Itoh,^a Shigeo Takahashi,^a Tetsuya Ueki,^a Takashi Sugiyama,^b T. Tomoyoshi Takahashi^c and C. Akira Horiuchi^{a,*}

^aDepartment of Chemistry, Rikkyo (St. Paul's) University, 3-34-1 Nishi-Ikebukuro, Toshima-ku, Tokyo 171-8501, Japan ^bInstitute for Chemical Research, Kyoto University, Uji, Kyoto 611-0011, Japan

^cDepartment of Chemistry, The Jikei University School of Medicine, Kokuryo-cho Chofu, Tokyo 182-8570, Japan

Tandem Michael-aldol reaction via 6-endo-dig cyclization of ynone-chalcogenides: synthesis of 2-unsubstituted 3-(hydroxyalkyl)-chalcogenochromen-4-ones

Tetrahedron Letters 43 (2002) 7039

Tadashi Kataoka,* Hironori Kinoshita, Sayaka Kinoshita and Tatsunori Iwamura

Gifu Pharmaceutical University, 6-1 Mitahora-higashi 5-chome, Gifu 502-8585, Japan

RCHO
$$BF_3 \cdot Et_2O$$
 OH O CH_2Cl_2 , $-20^{\circ}C$, 20 h CH_2Cl_2 , $-20^{\circ}C$

Asymmetric synthesis of a C1-C19 fragment of ulapualide A

Tetrahedron Letters 43 (2002) 7043

Cassandra A. Celatka and James S. Panek*

Department of Chemistry and Center for Streamlined Synthesis, Metcalf Center for Science and Engineering, Boston University, 590 Commonwealth Avenue, Boston, MA 02215, USA

4,5-Dehydrooctafluoro[2.2]paracyclophane: facile generation and extraordinary Diels-Alder reactivity

Tetrahedron Letters 43 (2002) 7047

Merle A. Battiste,* Jian-Xin Duan, Yi-An Zhai, Ion Ghiviriga, Khalil A. Abboud, Adrian Roitberg, G. Robert Shelton and William R. Dolbier, Jr.*

Department of Chemistry, University of Florida, PO Box 117200, Gainesville, FL 32611-7200, USA

Synthesis of 2,5-dihalothiazole-4-carboxylates

Tetrahedron Letters 43 (2002) 7051

John F. Okonya and Fahad Al-Obeidi*

Aventis Combinatorial Technologies Center, Selectide, 1580 E. Hanley Blvd., Tucson, AZ 85737, USA

The silyl-Prins reaction: a novel method for the synthesis of dihydropyrans

Tetrahedron Letters 43 (2002) 7055

Adrian P. Dobbs* and Saša Martinović

School of Chemistry, University of Exeter, Stocker Road, Exeter EX4 4QD, UK

TMS
$$R = H, Me$$

$$R^{1} = Alkyl, aryl$$

$$R = H, Me$$

$$R^{2} = Alkyl, aryl$$

$$R = H, Me$$

$$R^{3} = Alkyl, aryl$$

$$R = H, Me$$

$$R^{3} = Alkyl, aryl$$

$$R = H, Me$$

$$R^{4} = Alkyl, aryl$$

$$R = H, Me$$

$$R^{5} = Alkyl, aryl$$

$$R = H, Me$$

$$R^{6} = Alkyl, aryl$$

Effect of hydroxyl groups in receptors bearing disulfonamide on anion recognition in acetonitrile- d_3

Tetrahedron Letters 43 (2002) 7059

Shin-ichi Kondo,* Takashi Suzuki and Yumihiko Yano

Department of Chemistry, Faculty of Engineering, Gunma University, Kiryu, Gunma 376-8515, Japan

To elucidate the roles of hydroxyl group on anion-recognition chemistry, receptors bearing disulfonamide and hydroxyl groups were prepared and their anion-binding properties were evaluated in acetonitrile.

Engineering reactions in crystals: *gem*-dialkoxy substitution enables the photodecarbonylation of crystalline 2-indanone

Danny Ng, Zhe Yang and Miguel A. Garcia-Garibay*

Department of Chemistry, University of California, Los Angeles, CA 90095-1569, USA

Synthesis of polypropionate motifs containing the anti-anti unit

Tetrahedron Letters 43 (2002) 7067

Philippe Mochirian,^a Benoit Cardinal-David,^a Brigitte Guérin,^a Michel Prévost^a and Yvan Guindon^{a,b,*}

^aInstitut de recherches cliniques de Montréal (IRCM), Bio-organic Chemistry Laboratory, 110, avenue des Pins Ouest, Montréal, Québec, Canada H2W 1R7

^bDepartment of Chemistry and Department of Pharmacology, Université de Montréal, CP 6128, succursale Centre-Ville, Montréal, Québec, Canada H3C 3J7

Reported herein is the iteration of a strategy employing a Mukaiyama reaction in tandem with a hydrogen transfer reaction for the elaboration of four polypropionate motifs containing the *anti–anti* unit. In this process, Lewis acid acts as the key element in controlling the diastereoselectivity of each step, the outcome of which is >20:1 for all of the reactions performed.

One step facile synthesis of bromo calix[n] arenes

Satish Kumar, H. M. Chawla and R. Varadarajan*

Department of Chemistry, Indian Institute of Technology, New Delhi, India

Bromination of p-tert-butylcalix[n]arenes under different reaction conditions can provide either methylene bridge or ring substituted calix[n]arenes that are usually only amenable through long circuitous routes.

Tetrahedron Letters 43 (2002) 7073

The reaction of aryl triflates and aryl pivalates with electrophiles. The triflate as a *meta*-directing group

Tetrahedron Letters 43 (2002) 7077

George A. Kraus,* Wenge Cui and Young Ho Seo

Department of Chemistry, Iowa State University, Ames, IA 50011, USA

Electrophilic reactions of substituted aryl triflates yield products wherein the substituents direct the regioselectivity of electrophilic substitution.

$$R_1$$
 R_1
 R_2
 E^+
 R_1
 R_2
 R_2

Synthesis of furanochromones: a new step in improvement of fluorescence properties

Andrey S. Klymchenko, a,b,* Turan Ozturka and Alexander P. Demchenko a,c

^aTUBITAK RIGEB and TUBITAK MRC, Gebze-Kocaeli 41470, Turkey

^bDepartment of Chemistry, Kyiv National Taras Shevchenko University, 01033 Kyiv, Ukraine ^cA. V. Palladin Institute of Biochemistry, 9 Leontovicha str., 02030 Kyiv, Ukraine

An improvement in the procedure for the preparation of 3-hydroxychromones resulted in the synthesis of furanochromones. They exhibit the highest fluorescence quantum yields among all 3-hydroxychromones known to date and show larger separation between the two emission bands. This makes them promising for designing two-band fluorescence sensors.

Regioselective ring opening of epoxides by nucleophiles mediated by lithium bistrifluoromethanesulfonimide

Tetrahedron Letters 43 (2002) 7083

Janine Cossy, a,* Véronique Bellosta, a Claire Hamoir and Jean-Roger Desmurs

^aLaboratoire de Chimie Organique associé au CNRS, ESPCI, 10 rue Vauquelin, 75231 Paris Cedex 05, France ^bRhodia, 190 avenue Thiers, 69457 Lyon Cedex 06, France

$$\begin{array}{c|c} O & LiNTf_2 & HO \\ \hline R' & Nu & R & R' \\ \hline CH_2Cl_2, 20 \text{ h, rt} & \\ \end{array}$$

 $Nu = HNR_2$, H_2NR , PhSH, $H_2N-N(Me)_2$

Palladium-catalyzed hydrodehalogenations by fluoride activated polymethylhydrosiloxane

Tetrahedron Letters 43 (2002) 7087

Robert E. Maleczka, Jr.,* Ronald J. Rahaim, Jr. and Robson R. Teixeira Department of Chemistry, Michigan State University, East Lansing, MI 48824, USA

$$R = X = \begin{cases} 6 \text{ eq. PMHS,} \\ 12 \text{ eq. KF,} \\ 1 \text{ mol } \% \text{ (Ph}_3\text{P)}_2\text{PdCl}_2, \\ \text{THF, rt-70}^6\text{C} \end{cases} \qquad R = \text{aryl, styryl,} \\ x = \text{Br, I} \end{cases}$$

Palladium-catalyzed carbon-carbon coupling reactions using aryl Grignards

Tetrahedron Letters 43 (2002) 7091

Christine Gottardo* and Andrea Aguirre Department of Chemistry, Lakehead University, Thunder Bay, Ont., Canada P7B 5E1

Grignard reagents were used in the Sonogashira coupling of alkynes and aryl halides.

R = EWG, EDG

Cross-metathesis and ring-closing metathesis of olefinic monosaccharides

Tetrahedron Letters 43 (2002) 7095

Maarten H. D. Postema* and Jared L. Piper

Department of Chemistry, Wayne State University, Detroit, MI 48202, USA

Addition of trialkylaluminum reagents to glyconolactones. Synthesis of 1-C-methyl GlcNAc oxazoline and thiazoline

Tetrahedron Letters 43 (2002) 7101

Spencer Knapp,* Chunhua Yang and Thomas Haimowitz

Department of Chemistry & Chemical Biology, Rutgers—The State University of New Jersey, 610 Taylor Road, Piscataway, NJ 08854-8087, USA

A novel, facile methodology for the synthesis of N,N'-bis(tert-butoxycarbonyl)-protected guanidines using polymer-supported carbodiimide

Tetrahedron Letters 43 (2002) 7105

Olga Guisado, Sonia Martínez and Joaquín Pastor*

High Throughput Chemistry Group, Johnson & Johnson Pharmaceutical Research and Development, a division of Janssen-Cilag, S.A., Centro de Investigación Química. C/ Jarama s/n, Toledo 45007, Spain

A novel methodology for the synthesis of guanidines has been developed using PS-carbodiimide as activating agent for N,N'-bis(tert-butoxycarbonyl)thiourea and PS-trisamine as scavenger, followed by deprotection with trifluoroacetic acid.

Palladium-catalyzed cross coupling of Grignard reagents with in situ-derived enol phosphates

Tetrahedron Letters 43 (2002) 7111

Joseph A. Miller*

DSM Pharmaceuticals, Inc., 5900 NW Greenville Boulevard, Greenville, NC 27834, USA

$$R^{1} = \frac{R^{2}MgX}{ClPO(OPh)_{2}} = \begin{bmatrix} OPO(OPh)_{2} \\ R \end{bmatrix} \xrightarrow{R^{3}MgX} R^{1}$$

$$R^{3} = \frac{R^{3}MgX}{Pd Catalyst} = \frac{R^{3}}{R^{3}MgX}$$

Heck reactions in a non-aqueous ionic liquid using silica supported palladium complex catalysts

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