

Tetrahedron Letters Vol. 46, No. 17, 2005

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COMMUNICATIONS

Stereodivergent radical cyclisation reactions of cyclohexa-1,4-dienes

Mark C. Elliott* and Nahed Nasser Eid El Sayed

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Either isomer can be formed with good stereocontrol depending on choice of protecting group on oxygen.

Synthesis of the enantiomer of nelarabine

pp 2961-2964

Karim Herbal,* John Kitteringham, Martyn Voyle and Andrew J. Whitehead

A synthesis of the enantiomer of nelarabine is described. Subtle changes in the protecting group strategy allow for an efficient inversion of the C-2' stereocentre.

Regioselective cleavage of the bis-benzylidene acetal of D-mannitol under oxidative and reductive conditions: a new approach to C_2 -symmetric chiral ligands

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Appu Aravind, Subhendu K. Mohanty, T. Veerabhadra Pratap and Sundarababu Baskaran*

An amphiphilic C_{60} derivative with a tris(2,2'-bipyridine)ruthenium(II) polar head group: synthesis and incorporation in Langmuir films

pp 2969-2972

François Cardinali, Jean-Louis Gallani, Stefano Schergna, Michele Maggini* and Jean-François Nierengarten*

RO
$$1 R = C_{12}H_{25}$$

Biotechnological production of two new 8,4'-oxynorneolignans by elicitation of *Echinacea purpurea* cell cultures

pp 2973-2977

Wen-Wu Li* and Wolfgang Barz

Synthesis of tetrasubstituted Zn(II)-phthalocyanines carrying four carboranyl-units as potential BNCT and PDT agents

pp 2979-2982

Francesca Giuntini, Yann Raoul, Donata Dei,* Moira Municchi, Giacomo Chiti, Clara Fabris, Paolo Colautti, Giulio Jori and Gabrio Roncucci

(i)+

Synthesis of chiral N-protected 1,2-dihydro-quinoline-2-carbonitrile and 1,2-dihydro-isoquinoline-1-carbonitrile via an asymmetric Reissert reaction

pp 2983-2987

Mickaël Pauvert, Sylvain C. Collet, Marie-Jo Bertrand, André Y. Guingant* and Michel Evain

(i)+

The asymmetric synthesis of each of the above Reissert adducts has been realised.

A simple and convenient procedure for the synthesis of 1-aminophosphonates from aromatic aldehydes

pp 2989-2991

Babak Kaboudin* and Khavar Moradi

Catalytic oxidation of catechins to p-benzoquinones with hydrogen peroxide/methyltrioxorhenium

pp 2993-2996

Roberta Bernini,* Enrico Mincione, Gianfranco Provenzano and Giancarlo Fabrizi*

Copper-catalyzed aryl amination in aqueous media with 2-dimethylaminoethanol ligand

pp 2997-3001

Zhikuan Lu and Robert J. Twieg*

$$R = \frac{\text{Cul (cat.)}}{\text{H}_2\text{NR}_1} \frac{\text{Cul (cat.)}}{\text{K}_3\text{PO}_4,\text{H}_2\text{O}} R = \text{NHR}_1$$

Copper-catalyzed amination of aryl bromides and iodides under mild conditions has been developed with 2-dimethylaminoethanol as ligand and water as solvent. A variety of hydrophilic and hydrophobic aryl halide substrates have been aminated in good yield with a variety of amino acids, amino alcohols and peptides.



Photocyclodimers from Z-ligustilide. Experimental results and FMO analysis

pp 3003-3006

Beatriz Quiroz-García, Ricardo Figueroa, J. Antonio Cogordan and Guillermo Delgado*

The irradiation of the natural phthalide Z-ligustilide (1) gave the natural dimeric phthalide riligustilide (5), and three novel dimeric phthalides 7, 8 and 9.

Solid support synthesis of 15-membered macrocycles containing a serotonin unit

pp 3007-3010

Alexander S. Kiselyov

Water-based biphasic media for exothermic reactions: green chemistry strategy for the large scale preparation of clofibric acid and analogues

pp 3011-3013

Ajay K. Bose,* Maghar S. Manhas, Subhendu N. Ganguly, Suhas Pednekar and Arun Mandadi

An eco-friendly improved procedure is described for the synthesis of clofibric acid and analogues.

De novo synthesis of galacto-sugar δ -lactones via a catalytic osmium/palladium/osmium reaction sequence

pp 3015-3019

Md. Moinuddin Ahmed and George A. O'Doherty*

The photochemical isomerization of cross-linked 1,3,5-tristyrylbenzene dendrimer with hula-twist mechanism

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Mayuko Uda, Atsuya Momotake and Tatsuo Arai*

Efficient one-pot ring-opening/aldol reactions using (cyclopropyl)methylstannanes Bernard Leroy

pp 3025-3028

Lewis acid catalysed rearrangements of unsaturated bicyclic [2.2.n] endoperoxides in the presence of vinyl silanes; access to novel Fenozan BO-7 analogues

pp 3029-3032

Paul M. O'Neill,* Sarah L. Rawe, Richard C. Storr, Stephen A. Ward and Gary H. Posner

TMSOTf
Ar
O
$$Me_3Si$$

TMSOTf
 Me_3Si

TMSOTf
 Me_3Si

TMSOTf
 Me_3Si

TMSOTf
 Me_3Si

TMSOTf
 Me_3Si

TMSOTf

Five- and six-membered bicyclic [2.2.n] endoperoxides can readily be transformed into silyl containing endoperoxides in good yield and high stereoselectivity by treatment with allyltrimethylsilane and either TMSOTf or SnCl₄ as Lewis acid.

Synthesis of PI-88 analogue using novel O-glycosidation of exo-methylenesugars

pp 3033-3036

Rie Namme, Takashi Mitsugi, Hideyo Takahashi and Shiro Ikegami*

Design and synthesis of novel oxa-bridged isoxazolidines and 1,3-aminoalcohols

pp 3037-3040

Krishna P. Kaliappan,* Prasanta Das and Nirmal Kumar

A simple and efficient method for α -bromination of carbonyl compounds using N-bromosuccinimide in the presence of silica-supported sodium hydrogen sulfate as a heterogeneous catalyst

pp 3041-3044

Biswanath Das,* Katta Venkateswarlu, Gurram Mahender and Ibram Mahender

$$R^{1} \xrightarrow{\text{NBS}} R^{2} \xrightarrow{\text{NaHSO}_{4} \cdot \text{SiO}_{2}} R^{1} \xrightarrow{\text{R}^{1}} R^{2}$$

A total synthesis of the epoxyquinone based antifungal natural product (±)-ambuic acid Goverdhan Mehta* and Subhas Chandra Pan

pp 3045-3048

 $Stereoselective \ synthesis \ of \ (-)-gabosine \ C \ using \ a \ Nozaki-Hiyama-Kishi \ reaction \ and \ RCM$

pp 3049-3051

G. Venkata Ramana and B. Venkateswara Rao*

Petasis reaction of activated quinoline and isoquinoline with various boronic acids

pp 3053-3056

Yu Mi Chang, Seung Hwan Lee, Mi Hye Nam, Min Young Cho, Young Sang Park and Cheol Min Yoon*

or
$$RB(OH)_2$$
 $R = Aryl \text{ or vinyl},$ $R = COOEt$

Photoalkylation of 2,3-dicyanonaphthalene by methoxy-substituted 1,2-diarylcyclopropanes

pp 3057-3060

Hajime Maeda, Naohiro Matsukawa, Koichiro Shirai and Kazuhiko Mizuno*

Diels-Alder reactions of bis(spirodienone) derivatives of calix[4]arene with acetylenes: highly regio- and stereoselective synthesis of bisbicyclo[2.2.2]octenone derivatives

pp 3061-3063

R. Luxmi Varma,* V. B. Ganga and E. Suresh

$$Bu'$$

$$Bu'$$

$$E = CO_2Me$$

$$Bu'$$

$$A = CO_2Me$$

Highly regio- and stereoselective Diels-Alder reactions of bis(spirodienone) derivatives of calix[4]arene with acetylenes that provide easy access to bisbicyclo[2.2.2]octenones are described.

Synthesis and structural studies of peptides containing a glucose-derived furanoid sugar amino acid

pp 3065-3070

Tushar K. Chakraborty,* Saumya Roy, S. Kiran Kumar and Ajit C. Kunwar*

An efficient synthesis of alkyl α -(hydroxymethyl)acrylates induced by DABCO in an aqueous medium

pp 3071-3072

Taoufik Turki, Jean Villiéras and Hassen Amri*

Alkyl α -(hydroxymethyl)acrylates are prepared in high yields by coupling alkyl acrylates with 30% aqueous formaldehyde using DABCO as the catalyst and DME or THF as solvent.

Lewis acid catalyzed asymmetric halohydrin reactions of chiral α,β -unsaturated carboxylic acid derivatives with N-halosuccinimide (NXS) as the halogen source

pp 3073-3077

Saumen Hajra,* Manishabrata Bhowmick and Ananta Karmakar

$$R^{1} = \text{alkyl, aryl}$$

$$R^{2} = \text{H, Me}$$

$$R^{1} \text{Me}$$

$$R^{1} \text{Me}$$

$$R^{1} \text{Me}$$

$$R^{1} \text{Me}$$

$$R^{1} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{1} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{3} \text{Holosofth} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{3} \text{Holosofth} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{3} \text{Holosofth} \text{Me}$$

$$R^{4} \text{Holosofth} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{2} \text{Holosofth} \text{Me}$$

$$R^{3} \text{Holosofth} \text{Me}$$

$$R^{4} \text{Holosofth} \text{Me}$$

$$R^{4} \text{Holosofth} \text{Me}$$

$$R^{4} \text{Holosofth} \text{Me}$$

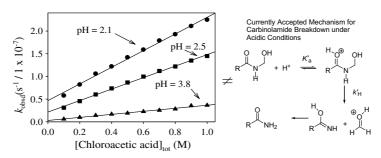
$$R^{4} \text{Holosofth} \text{Me}$$

$$R^{5} \text{Holosofth} \text{Me}$$

General-buffer catalysis of the reaction of N-(hydroxymethyl)benzamide: a new pathway for the aqueous reaction of carbinolamides

pp 3079-3083

Amanda G. Mennenga, Amy L. Johnson and Richard W. Nagorski*



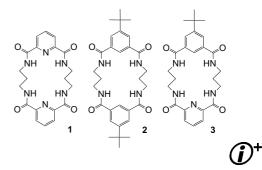
(i)+

A hybrid macrocycle containing benzene and pyridine subunits is a better anion receptor than both its homoaromatic congeners

pp 3085-3088

Michał J. Chmielewski and Janusz Jurczak*

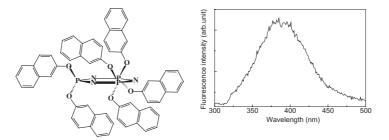
The hybrid tetraamide receptor 3 benefits from both suitable preorganization provided by the 2,5-diamidopyridine unit and good anion binding properties of the 1,3-diamidobenzene moiety and therefore is a better anion receptor than its homoaromatic counterparts 1 and 2.



Photophysical studies on multichromophoric cyclotriphosphazenes. Trinuclear excimer formation in hexakis(2-naphthyloxy)cyclotriphosphazene

pp 3089-3092

Nitin Chattopadhyay,* Basudeb Haldar, Arabinda Mallick and Saumitra Sengupta*



Cyclisation at very high temperature. Thermal transformations of N-alkyl and N,N-dialkyl cinnamic amides into pyrrolidin-2-ones under FVT conditions

pp 3093-3095

Stanisław Leśniak* and Beata Pasternak

An easy access to unsymmetric trisubstituted methane derivatives (TRSMs)

pp 3097-3102

Sajal Kumar Das, Shagufta and Gautam Panda*

MgBr

Conc.
$$H_2SO_4$$
 or

Anhyd. $AICl_3$

OCH₃

OCH₃
 X (ortho or para)

R

Y

OCH₃

OCH₃

OCH₃
 $X = OH, OCH_3, NHC_2H_5, N(CH_3)_2, NH_2; Y = S$

R= 2-Thienyl, 3-Pyridyl, (5-Methyl)-2-furyl

(R)-2,3-Cyclohexylideneglyceraldehyde, a novel template for facile and simple entry into chiral hydroxy γ -lactones: synthesis of (–)-muricatacin

pp 3103-3105

Bhaskar Dhotare and Angshuman Chattopadhyay*

Solid-phase synthesis of [1,2,4]triazolo[3,4-a]phthalazine and tetrazolo[5,1-a]phthalazine derivatives Jong Yeon Hwang, Hyung-Sub Choi and Young-Dae Gong*

pp 3107-3110

Rotational isomerism of a resveratrol tetramer, shoreaketone, in Shorea uliginosa

pp 3111-3114

Tetsuro Ito,* Miyuki Furusawa, Ibrahim Iliya, Toshiyuki Tanaka, Ken-ichi Nakaya, Ryuichi Sawa, Yumiko Kubota, Yoshikazu Takahashi, Soedarsono Riswan and Munekazu Iinuma

A new resveratrol tetramer, shoreaketone, was isolated from the stem bark of Shorea uliginosa (Dipterocarpaceae). The structure has a novel framework of heptacyclic ring system including an α,β-unsaturated carbonyl group. In NMR spectra shoreaketone is observed as two different conformers due to rotational isomerism.

Catalytic enantioselective fluorination of α -cyano acetates catalyzed by chiral palladium complexes Hye Ran Kim and Dae Young Kim*

pp 3115-3117

Regioselective synthesis of triiodo-o-carboranes and tetraiodo-o-carborane

pp 3119-3122

Hiroto Yamazaki, Kiminori Ohta and Yasuyuki Endo*

suitable building block for supramolecular construction

Dynamic kinetic resolution during iron carbonyl promoted [6+2] ene-type reactions

pp 3123-3126

Anthony J. Pearson* and Xiaolong Wang



Total synthesis of the alkaloid (-)-codonopsinine from L-xylose

pp 3127-3129

S. Chandrasekhar,* V. Jagadeshwar and S. Jaya Prakash

$$\begin{array}{c} \text{HO} \quad \text{OH} \\ \text{HO} \quad \text{OOH} \\ \text{HO} \quad \text{OOH} \\ \text{OCHO} \end{array} \begin{array}{c} \text{OMe} \\ \text{H}_3\text{C} \quad \text{N} \\ \text{CH}_3 \\ \text{OMe} \\ \text{(-)-Codonopsinine} \end{array}$$

The total synthesis of (–)-codonopsinine is described from commercially available L-xylose in 20% overall yield. The key steps included Julia *trans* olefination and cascade epoxidation–cyclisation strategies.



pp 3131-3135

Solid-phase synthesis of α-substituted proline hydantoins and analogs

Jordi Alsina, William L. Scott* and Martin J. O'Donnell*

$$\begin{array}{c} CI \\ N \\ O \\ O \\ O \\ \end{array}$$

$$\begin{array}{c} R_1 \\ N \\ O \\ \end{array}$$

$$\begin{array}{c} R_1 \\ O \\$$

Ionic liquid $[omim][PF_6]$ as an efficient and recyclable reaction media for the cyanosilylation of aldehydes without Lewis acid or any special activation

pp 3137-3139

Zhi-Liang Shen, Shun-Jun Ji* and Teck-Peng Loh*

$$\begin{array}{c} O \\ R \\ H \end{array} + \begin{array}{c} TMSCN \\ \hline 2. \ 1M \ HCl/THF \\ \end{array} \begin{array}{c} OH \\ R \\ \hline CN \\ \end{array} \\ \begin{array}{c} OH \\ CN \\ \end{array}$$

Ionic liquid [omim][PF₆] has been demonstrated as an efficient, environmentally friendly and recyclable reaction media as well as a promoter for the cyanosilylation of aldehydes under mild conditions. The recovered ionic liquid could be reused for subsequent runs with only a gradual decrease in activity.

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*Corresponding author

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