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Tetrahedron Letters

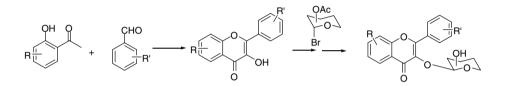
Tetrahedron Letters Vol. 49, No. 51, 2008

Contents

COMMUNICATIONS

Synthesis of a library of glycosylated flavonols

Zhitao Li^{*}, George Ngojeh, Paul DeWitt, Zhi Zheng, Min Chen, Brendan Lainhart, Vincent Li, Peter Felpo



Flavonols are an important class of natural products isolated from plants. Some glycosylated flavonols showed very interesting biological activities. A library of flavonols has been made through Algar–Flynn–Oyamada reaction from 2'-hydroxyacetophenones and benzaldehydes. Glycosylation of these flavonols with various glycosyl donors affords a library of glycosylated flavonols. These compounds are potentially useful pharmacologically active compounds and will be studied for biological activities.

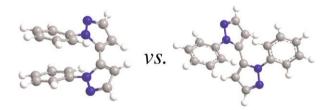
The use of a molecular balance derived from 5,5'-bipyrazole to calculate π - π stacking interactions lbon Alkorta *, Fernando Blanco, José Elguero

pp 7246-7249

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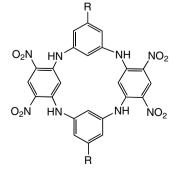
pp 7243-7245

IDOII AIROITA , FEITIAIIUO DIAIICO, JOSE EIGUEIO



Metal-free synthesis of azacalix[4]arenes

Mounia Touil, Mohammed Lachkar, Olivier Siri *

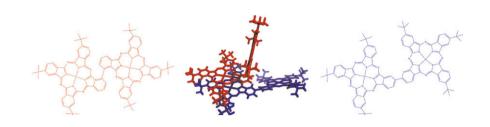


The facile preparation of new N(H)-bridged azacalix[4]arenes has been achieved by stepwise nucleophilic aromatic substitutions assisted by hydrogen bonding interactions.

Synthesis and properties of C-C conjugated phthalocyanine dimers

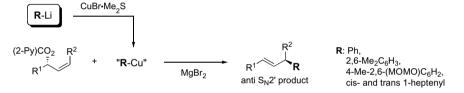
Hasrat Ali, Pierre Baillargeon, Johan E. van Lier

pp 7253-7255



New protocol for allylic substitution with aryl and alkenyl copper reagents derived from organolithiums Yohei Kiyotsuka, Yuichi Kobayashi ^{*}

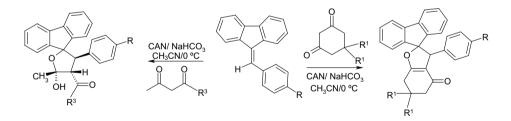
pp 7256-7259



An efficient one-pot synthesis of spiro dihydrofuran fluorene and spiro 2-hydroxytetrahydrofuran fluorene derivatives via [3+2] oxidative cycloaddition mediated by CAN

pp 7260-7263

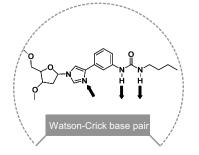
G. Savitha, R. Sudhakar, P. T. Perumal *



Synthesis and DNA duplex recognition of a triplex-forming oligonucleotide with an ureido-substituted 4-phenylimidazole nucleoside

pp 7264-7267

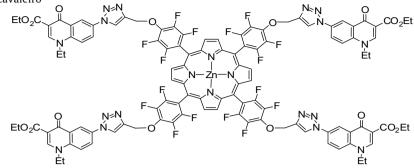
Falk Wachowius, Michael Rettig, Gottfried Palm, Klaus Weisz *



Synthesis of porphyrin-quinolone conjugates

pyridinecarboxaldehyde, under mild reaction conditions.

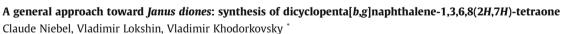
Fernanda da C. Santos, Anna C. Cunha, Maria Cecília B. V. de Souza, Augusto C. Tomé, Maria G. P. M. S. Neves, Vitor F. Ferreira, José A. S. Cavaleiro *



Bischler–Napieralski cyclocondensation in the synthesis of new 11*H***-pyrimido**[4,5-*b*][1,4]benzodiazepines Justo Cobo ^{*}, Manuel Nogueras, John N. Low, Ricaurte Rodríguez

A synthetic strategy based on nitrosation-aminolysis-nitroso reduction and Bischler-Napieralski cyclocondensation has been developed for the synthesis of a family of 2-amino-4-methoxy-11H-pyrimido[4,5-b][1,4]benzodiazepines.

A convenient synthetic route to a useful synthon: 4-bromo-2-pyridinecarboxaldehyde Nicolas Zaman, Régis Guillot, Katell Sénéchal-David ^{*}, Marie-Laure Boillot



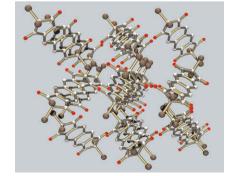
We have developed a novel four-step method to synthesise the versatile synthon, 4-bromo-2-pyridinecarboxaldehyde, from 2-picoline-N-oxide via 4-nitro-2-

pp 7276-7278



pp 7271-7273





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7237

Microwave-assisted Pd(OH)₂-catalyzed direct C-H arylation of free-(NH₂) adenines with aryl halides Sophian Sahnoun, Samir Messaoudi, Jean-François Peyrat, Jean-Daniel Brion, Mouad Alami

49-86%

k'n

12 examples

Preparation of a cationic bisoxazolinic nickel pincer catalyst and its applications to Michael addition and Mizorokipp 7287-7289 **Heck reaction**

Koichi Mitsudo *, Tatsuhiko Imura, Takashi Yamaguchi, Hideo Tanaka *

NC

Synthesis of benzimidazoles from 1,1-dibromoethenes

Wang Shen *, Todd Kohn, Zice Fu, XianYun Jiao, Sujen Lai, Michael Schmitt

0°C - room temp. quant. R = Me, Et (3.0 equiv) $R' = (CH_2)_2COMe$ 1 (5 mol%) Na₂CO₃ (3.0 equiv) CO₂Bu Ph—I CO₂Bu DMF (5.0 equiv)

Access to chiral tertiary amines via the iridium-catalyzed asymmetric hydrogenation of enamines Pradeep Cheruku, Tamara L. Church, Anna Trifonova, Thomas Wartmann, Pher G. Andersson

(o-tol)2 R^1 R^2 R², R³ = alkyl or aryl up to 99% conv. [BAr_F] up to 87% ee

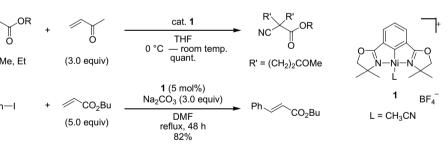


Pd(OH)₂/C / Cul, Cs₂CO₃, NMP, 160 °C

15 min to 1 h



 R^{1} H^{N} H^{N}



DABCO NMP, 100 °C R

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pp 7284-7286

pp 7290-7293

Copper-free Sonogashira coupling in amine-water solvent mixtures Anna Komáromi, Gergely László Tolnai, Zoltán Novák

+ <u></u>_R' ^sBuNH₂/H₂O 1:1,

Effect of stabilizing ligands bearing ferrocene moieties on the gold nanoparticle-catalyzed reactions of arylboronic pp 7299-7302 acids

-B(OH)

B(OH)

Au(III):Ligand with Ferrocene MoietiesinTHF

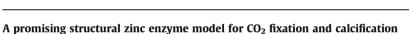
K₂CO₃(3eq), H₂O, air, RT

High Activities

25 °C, 90 min

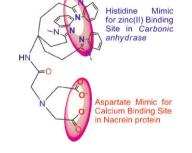
Laksamee Chaicharoenwimolkul, Ampaporn Munmai, Sanoe Chairam, Udomchai Tewasekson, Sarawut Sapudom, Yuthana Lakliang, Ekasith Somsook

The homocoupling reaction of phenylboronic acid and demetalation reaction of ferrocenylboronic acid was inhibited and highly active, respectively, in the presence of gold nanoparticles stabilized by ligands containing ferrocene moieties.



Mohamed M. Ibrahim^{*}, Shaban Y. Shaban, Kazuhiko Ichikawa

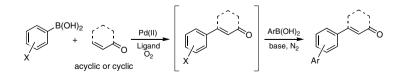
A new ligand: N-{tris([2-[(1-methylbenzimidazol-2-yl)ethyl]methyl]amino)-2-oxoethyl} iminodiacetic acid has been synthesized and characterized, and was used to prepare a zinc complex as a promising model for the active site of the nacreous protein in mollusc shells. The CO_2 fixation and calcification have been studied in light of the influence of the pK_a value of the coordinated water molecule and the carboxylate groups.



R'

Chemoselective three-component coupling via a tandem Pd-catalyzed boron-Heck and Suzuki reactions

Justin O'Neill, Kyung Soo Yoo, Kyung Woon Jung



pp 7303-7306

ОН

pp 7294-7298

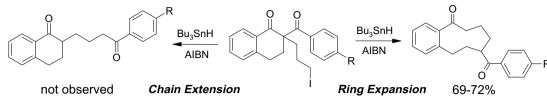
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pp 7307-7310

Free radical ring expansion and spirocyclization of 1,3-diketone derivatives

Wei Xu, Jian-Ping Zou *, Xue-Jun Mu, Wei Zhang *

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Free radical-promoted three-carbon ring expansions of 1,3-diketones to form corresponding nine-membered 1,6-diketones and associated spirocyclization reactions are described.

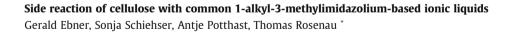
On the structure of the bioactive constituent from ayurvedic medicine Salacia reticulata: revision of the literature pp 7315-7317 Osamu Muraoka^{*}, Weijia Xie, Genzoh Tanabe, Mumen F. A. Amer, Toshie Minematsu, Masayuki Yoshikawa

to be de-O-sulfated kotalanol 4 on the basis of the detailed analysis of the spectral data.

tert-Butyl isocyanide revisited as a convertible reagent in the Groebke-Blackburn reaction

Mikhail Krasavin^{*}, Sergey Tsirulnikov, Mikhail Nikulnikov, Yuri Sandulenko, Konstantin Bukhryakov

TFA reflux, 3 h



lonic liquids with 1-alkyl-3-methyl-imidazolium cations react at C-2 with the reducing ends of cellulose forming a carbon-carbon bond, and are thus no inert cellulose solvents. The reaction is catalyzed by bases, such as commonly present impurities in ILs.

AcO 3% cellulose in IL r.t., 8 h R = H, 2-naphthyl

67-100%

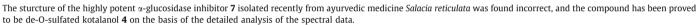
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KOH (3 eq.)

MeOH-H₂O (1:1) 60 °C, 8 h

ÓН

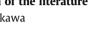


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pp 7318-7321



pp 7311-7314

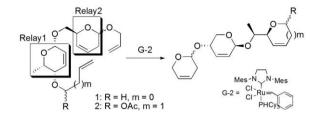


NH,

35 - 78% 10 examples

Extended RCM-ROM sequences: a novel approach to polyunsaturated trisaccharides

Morgan Donnard, Théophile Tschamber, Jacques Eustache



37% KF/Al₂O₃ 18-crown-6 microwave 180 °C, DMSO

EWG

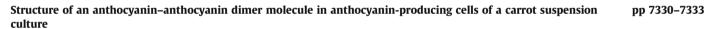
OH

Xylose / Galactose

Glucose Sinapic acid

The first examples of controlled RCM-ROM-RCM-ROM-RCM sequences involving non-strained heterocyclic relays are described.

Investigation of the N-arylation of various substituted indoles using microwave-assisted technology Gregory L. Frayne, Gary M. Green *



H₃C-Ċ-H

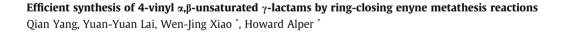
ÓН

> Galactose Xvlose

Glucose

он но

ĖWG



 $\begin{array}{c} O \\ R^{1} \\ \hline \\ O \\ R^{2} \\ \hline \\ R^{3} \\ \end{array} \begin{array}{c} Cl \\ Cl \\ PCy_{3} \\ Cl \\ PCy_{3} \\ PCy_{3} \\ Cl \\ PCy_{3} \\ PC$

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pp 7325-7327

7241

pp 7328-7329



2-Aminobenzimidazoles as potent ITK antagonists: de novo design of a pyrrole system targeting additional hydrogen pp 7337–7340 bonding interaction

Ho Yin Lo^{*}, Jörg Bentzien, Andre White, Chuk C. Man, Roman W. Fleck, Steven S. Pullen, Hnin Hnin Khine, Josephine King, Joseph R. Woska Jr., John P. Wolak, Mohammed A. Kashem, Gregory P. Roth, Hidenori Takahashi

De novo design of a series of 2-aminobenzimidazole-pyrrole type compounds as ITK inhibitors.

 R^1 = H or CH₃ R^2 = H, acetyl R^3 = nitro, bromo, acetyl, benzoyl

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