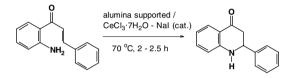


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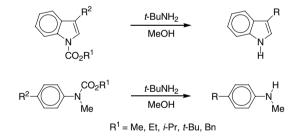
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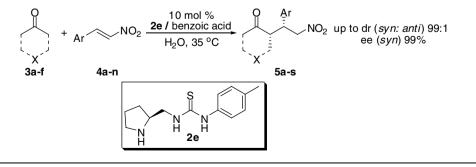


Cleavage of alkoxycarbonyl protecting groups from carbamates by t-BuNH2pp 17–20Oscar R. Suárez-Castillo,* Luis Alberto Montiel-Ortega, Myriam Meléndez-Rodríguez and
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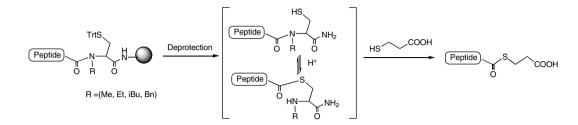
Michael additions in water of ketones to nitroolefins catalyzed by readily tunable and bifunctional pyrrolidine-thiourea organocatalysts

Yi-Ju Cao, Yuan-Yuan Lai, Xiang Wang, Yong-Jian Li and Wen-Jing Xiao*



N-Alkyl cysteine-assisted thioesterification of peptides

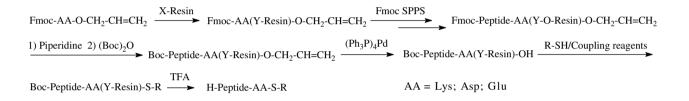
Hironobu Hojo,* Yuko Onuma, Yuri Akimoto, Yuko Nakahara and Yoshiaki Nakahara*



N-Alkyl cysteine at the C-terminus of peptides effectively promotes thioesterification by 3-mercaptopropionic acid.

Transfer allyl esters to thioesters in solid phase condition: synthesis of peptide thioesters by Fmoc pp 29–32 chemistry

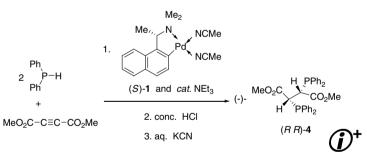
Lei Li and Pu Wang*



Asymmetric synthesis of dimethyl-1,2-bis-(diphenylphosphino)-1,2-ethanedicarboxylate by means of a pp 33–35 chiral palladium template promoted hydrophosphination reaction

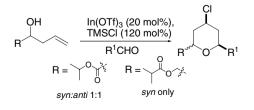
Lulu Tang, Yi Zhang, Luo Ding, Yongxin Li, Kum-Fun Mok, Wee-Chuan Yeo and Pak-Hing Leung*

An optically pure C_2 -symmetrical diphosphine ligand containing two ester functional groups at the two chiral carbon stereogenic centres was prepared efficiently from the asymmetric hydrophosphination reaction between diphenylphosphine and dimethyl acetylenedicarboxylate in the presence of an organopalladium(II) complex derived from (S)-N,N-dimethyl-1-(1-naphthyl)ethylamine.



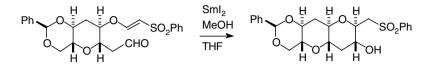
Stereochemical Prins cyclization: electronic versus steric effects on the synthesis of 2,4,6-trisubstituted pp 37–41 tetrahydropyran rings

Kok-Ping Chan, Ai-Hua Seow and Teck-Peng Loh*





SmI₂-induced reductive cyclization of (*E*)- and (*Z*)- β -alkoxyvinyl sulfones with aldehyde Tomohiro Kimura and Tadashi Nakata^{*}

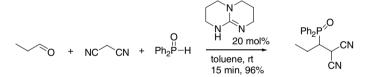


Utilization of 2,6-bis(2-benzimidazolyl)pyridine to detect toxic benzene metabolites Bolin Chetia and Parameswar K. Iyer*

It is demonstrated that 2,6-bis(2-benzimidazolyl)pyridine can act as a sensor to detect toxic metabolites of benzene such as phenol, hydroquinone, resorcinol, catechol and *p*-benzoquinone.

P-C Bond formation via direct and three-component conjugate addition catalyzed by 1,5,7-triazabicyclo[4.4.0]dec-5-ene (TBD)

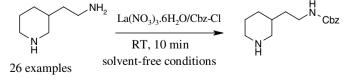
Zhiyong Jiang, Yan Zhang, Weiping Ye and Choon-Hong Tan*



The direct addition of P(O)–H bonds (dialkyl phosphites and diphenyl phosphonite) across various activated alkenes was catalyzed effectively by 1,5,7-triazabicyclo[4.4.0]dec-5-ene (TBD). This is a mild, rapid and efficient protocol to generate P–C bonds. This simple procedure allows a series of dialkyl alkylphosphonates and trisubstituted phosphine oxides to be prepared in high yields. Further investigation resulted in a convenient one-pot, three-component reaction involving diphenylphosphonite, malononitrile and an aldehyde.

A mild and efficient chemoselective protection of amines as N-benzyloxy carbonyl derivatives in the presence of La(NO₃)₃·6H₂O under solvent-free conditions

K. Chinni Mahesh, M. Narasimhulu, T. Srikanth Reddy, N. Suryakiran and Y. Venkateswarlu*



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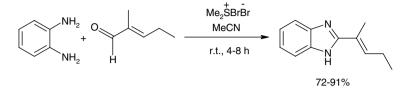
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pp 55–59

Efficient (bromodimethyl)sulfonium bromide mediated synthesis of benzimidazoles Biswanath Das,* Harish Holla and Yallamalla Srinivas



Enantioselective synthesis of (-)-cytoxazone and (+)-*epi*-cytoxazone via Rh-catalyzed diastereoselective pp 65–68 oxidative C–H aminations

R = CONH₂

ÖTBS R = SO₂NH₂

OB OTBS

(-)-cytoxazone

(+)-epi-cytoxazone

MeO

MeC

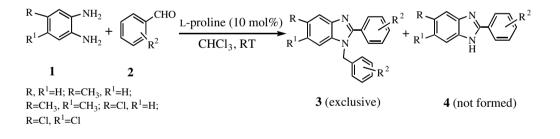
Srinivasarao V. Narina, Talluri Siva Kumar, Shyla George and Arumugam Sudalai*

сно

MeC

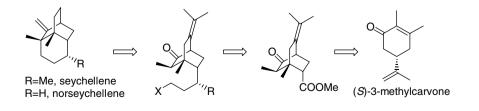
L-Proline catalyzed selective synthesis of 2-aryl-1-arylmethyl-1*H*-benzimidazoles

Ravi Varala, Aayesha Nasreen,* Ramu Enugala and Srinivas R. Adapa*



The first enantiospecific total synthesis of (+)-seychellene

A. Srikrishna,* G. Ravi and G. Satyanarayana



pp 61-64

pp 69–72

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Electroorganic synthesis of benzathine

Shaik Lateef, Srinivasulu Reddy Krishna Mohan and Srinivasulu Reddy Jayarama Reddy*



A mild and simple regioselective iodination of activated aromatics with iodine and catalytic ceric pp 81-83 ammonium nitrate

CAN (10 mol%) r. t., 3 h

Pd(II)-catalyzed oxidative cvclization

Biswanath Das,* Maddeboina Krishnaiah, Katta Venkateswarlu and V. Saidi Reddy

Synthesis of tribromobenzofuran and tribromobenzopyran derivatives from methyl 2-allyl-4,5,6tribromo-3-hydroxybenzoate

ÇO₂Me

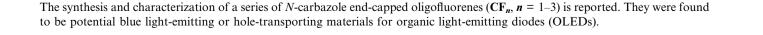
Faiz Ahmed Khan* and Laxminarayana Soma

OMe

Synthesis and characterization of N-carbazole end-capped oligofluorenes Vinich Promarak,* Sayant Saengsuwan, Siriporn Jungsuttiwong, Taweesak Sudyoadsuk and

ОН

Tinnagon Keawin



CF_n (n = 1-3)

 C_6H_{13}

 C_6H_{13}

CO_Me

ÇO₂Me

Br

Br

94%



pp 85-88

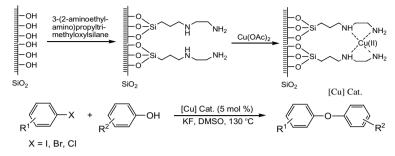


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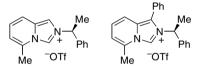
Immobilization of copper in organic-inorganic hybrid materials: a highly efficient and reusable catalyst pp 95–99 for the Ullmann diaryl etherification

Tao Miao and Lei Wang*



Synthesis of optically active imidazopyridinium salts and the corresponding NHCs Michael A. Schmidt and Mohammad Movassaghi*

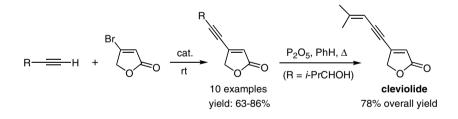




A convergent synthesis of optically active imidazo-[1,5-a]-pyridinium salts and the corresponding NHCs is described.

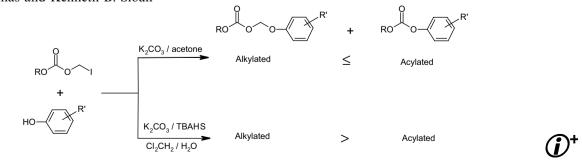
Facile access to 4-(1-alkynyl)-2(5H)-furanones by Sonogashira coupling of terminal acetylenes withpp 105–107β-tetronic acid bromide: efficient synthesis of cleviolideLaka Baukanunkas* Séhastian Câté and Brune Nidri

John Boukouvalas,* Sébastien Côté and Bruno Ndzi



Overcoming steric effects in the coupling reaction of alkyloxycarbonyloxymethyl (AOCOM) halides with phenols: an efficient synthesis of AOCOM phenolic prodrugs

Joshua D. Thomas and Kenneth B. Sloan*



Regioselective synthesis of 2,4,6-triaminopyridines

Rupa Shetty,* Duyan Nguyen, Dietmar Flubacher, Franziska Ruggle, Andreas Schumacher, Martha Kelly and Enrique Michelotti

A regioselective synthesis of 2,4,6-trisubstituted pyridine is described starting from 2,6-dibromo-4-nitropyridine. All three different regioisomers of the 2,4,6-triamino substituted pyridine have been synthesized in four to five steps. The method described is a general route to unsymmetrical 2,4,6-trisubstituted amino pyridines.

Synthesis of bis(alk-3-en-1-ynyl)benzene with either *E*- or *Z*-configuration via a one-pot three-component pp 119–124 coupling reaction and its optical properties

PdCl₂(dppf)/Cul/n-Bu₄NOH

Masayuki Hoshi,* Souichi Suzuki, Shingo Saitoh, Mitsuhiro Okimoto and Kazuya Shirakawa

Homochiral tripodal imidazolium receptors: structural and anion-receptor studies Nameer Alhashimy, Dermot J. Brougham, Joshua Howarth, Alan Farrell, Brid Quilty and

Me₃SiC ≣ CBr

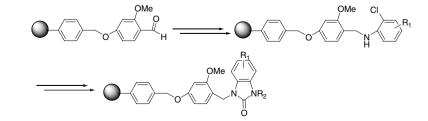
Cu(acac)₂/1M-NaOMe

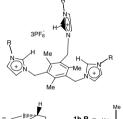
Kieran Nolan*

Two homochiral tripodal receptors have been characterised by X-ray crystallography, a first for this class of imidazolium receptor. These receptors were also screened for anion recognition. Both receptors demonstrated selectivity towards chloride and bromide with binding constants as high as 16,000.

C = C

Microwave-assisted traceless synthesis of benzimidazolones Xin-Jian Xu* and Ying-Xiao Zong





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pp 129-132

pp 125-128

 $(CH_2)_n$ $(CH_2)_n$ $(CH_2)_n$ $(CH_2)_n$

R-alcohols

predominate

v

85-95%

The stereochemistry of fatty acid hydroxylation by cytochrome P450_{BM3} Max J. Cryle, Nick J. Matovic and James J. De Voss^{*}

n = 8, 9, 10

- D2

A selective and convenient ruthenium mediated method for the synthesis of mixed acetals and orthoesters pp 137–140 Stanisław Krompiec,* Robert Penczek, Nikodem Kuźnik, Jan Grzegorz Małecki and Marek Matlengiewicz

$$R^{1}O$$

$$R^{3}OH$$

$$R^{3}OH$$

$$R^{3}OH$$

$$R^{2}OR^{3}$$

$$R^{$$

 $R^1 = n$ -Bu, PhCH₂ $R^2 = H$, Pr Y = H $R^1, Y = -CH_2CH_2CH_2O R^3 = n$ -Bu, Ph, PhCH₂, PhCH₂CH₂, *m*-MeC₆H₄, *p*-HOC₆H₄

A new method for the synthesis of symmetrical or mixed acetals and orthoesters via inter- or intramolecular addition of an OH group (alcoholic or phenolic) to O-allylic systems (ethers or acetals), catalyzed by $[RuCl_2(PPh_3)_3]$ is presented.

 $(NH + R^2 \xrightarrow{R^1}_X \xrightarrow{H_2O}_{rt} X \xrightarrow{R^2}_X X$

 R^1 , $R^2 = H/alkyl$; $X = CO_2R$, COR, CONH₂, CN

Significant rate acceleration of the aza-Michael reaction in water

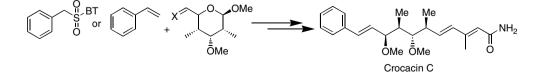
Brindaban C. Ranu* and Subhash Banerjee

Remarkable rate acceleration is reported for the aza-Michael reaction in water at room temperature in the absence of any catalyst or co-organic solvent.

20-50 min

A formal total synthesis of crocacin C

J. S. Yadav,* P. Venkatram Reddy and L. Chandraiah



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hh 141-142

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pp 149-151

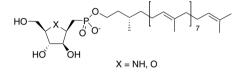
A high yielding, one-pot synthesis of O,S-dialkyl dithiocarbonates from alcohols using Mitsunobu's reagent

Devdutt Chaturvedi* and Suprabhat Ray

$$R^{2} \xrightarrow[R^{3}]{} OH + HO \xrightarrow[R^{6}]{} R^{5} \xrightarrow[CS_{2}, R.T., 4-8 h, 76-98\%]{} R^{2} \xrightarrow[R^{3}]{} S \xrightarrow[R^{6}]{} O \xrightarrow[R^{6}]{} R^{5}$$

Synthesis of 2',3'-dihydrosolanesyl analogues of β-D-arabinofuranosyl-1-monophosphoryldecaprenol with promising antimycobacterial activity promising antimycobacterial activity

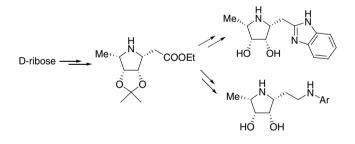
Michaël Bosco, Philippe Bisseret,* Patricia Constant and Jacques Eustache*



Stereoselective synthesis of novel five-membered homoazasugars. A convenient route to all-cis tetrasubstituted pyrrolidines

pp 159-162

Elena Moreno-Clavijo, Ana T. Carmona,* Antonio J. Moreno-Vargas and Inmaculada Robina*



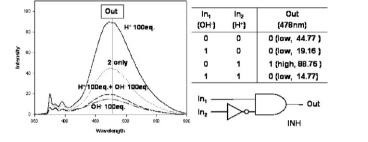
N-Phenylurea as an inexpensive and efficient ligand for Pd-catalyzed Heck and room-temperature pp 163–167 Suzuki reactions

Xin Cui, Yuan Zhou, Na Wang, Lei Liu* and Qing-Xiang Guo*

 $Ar - X + R \xrightarrow{Pd(OAc)_2} Ar \xrightarrow{R} TON = 10^3$ $Ar - X + R \xrightarrow{B(OH)_2} \frac{Pd(OAc)_2}{1-phenylurea} Ar \xrightarrow{R} TON = 10^4$

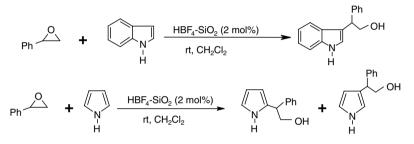
New imidazolium systems bearing two pyrene groups as fluorescent chemosensors for anions and anion pp 169–172 induced logic gates

Ha Na Lee, N. Jiten Singh, Sook Kyung Kim, Ji Young Kwon, Yoo Young Kim, Kwang S. Kim * and Juyoung Yoon *



Fluoroboric acid adsorbed on silica gel catalyzed regioselective ring opening of epoxides with nitrogen pp 173–176 heterocycles

B. P. Bandgar* and Abasaheb V. Patil



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