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COMMUNICATIONS

Nickel-catalyzed carbonylative cycloaddition of allyl halides and alkenes Daniel del Moral, Josep M. Moretó, Elies Molins, Susagna Ricart ^{*} pp 6947-6950



The carbonylative addition between allyl halides and alkenes is described. The [2+2+1] reaction is catalyzed by Ni(1), and takes place with different strained alkenes under very mild conditions. Changes in solvent and use of different amounts of water in the reaction produce changes on the final products obtained.

Microwave-assisted highly diastereoselective synthesis of oxazolidines derived from ketones and aminoalcohols pp 6951–6954 Philip C. Bulman Page ^{*}, Genna A. Parkes, Benjamin R. Buckley, Harry Heaney, Mostafa Gholizadeh, J. Steven Wailes



Novel light-fluorous TEMPO reagents and their application in oxidation reactions Adrian P. Dobbs ^{*}, Mark J. Penny, Peter Jones pp 6955-6958



Highly stereoselective allylic ethylation with alkoxytitanacyclopropane reagents. Synthesis of (1*R***/***S***,7***R***)-1,7-dimethylnonyl propanoate, the Western corn rootworm sex attractant** Vladimir E. Isakov, Oleg G. Kulinkovich *



MW H₂O

A mild Boc deprotection and the importance of a free carboxylate

Ali Thaqi, Adam McCluskey *, Janet L. Scott



ĊН,

Synthesis of novel annulated uracils via domino Knoevenagel-hetero-Diels–Alder reaction in aqueous media Malihe Javan Khoshkholgh, Saeed Balalaie ^{*}, Hamid Reza Bijanzadeh, Frank Rominger, Jürgen H. Gross

,OH



сно

H₃C

ĊН,

CuI H₂O, reflux

Jalil Noei, Ahmad R. Khosropour *

$$\text{RCH=NOH} \xrightarrow{\text{TiCl}_3\text{OTf}-[\text{bmim}]\text{Br}}_{(\text{NH}_4)_2\text{S}} \xrightarrow{\text{S}}_{\text{RC-NH}_2}$$

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A versatile synthesis of (+)-deoxoprosopinine and (-)-deoxoprosophylline

Enzo B. Arévalo-García *, Juan Carlos Colmenares



Eco-friendly polyethylene glycol promoted Michael addition reactions of α,β-unsaturated carbonyl compounds pp 6974-6976 Dalip Kumar^{*}, Gautam Patel, Braja G. Mishra, Rajender S. Varma^{*}



New reactivity of 1-(2-pyridyl)-2-propen-1-ol with nitro derivatives Donatella Giomi^{*}, Renzo Alfini, Alberto Brandi



The first asymmetric synthesis of all four isomers of cis- and trans-3,4-dihydroxy-3,4-dihydromollugin Naga Venkata Sastry Mudiganti, Sven Claessens, Norbert De Kimpe *



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Stereoselective synthesis of highly functionalised P-stereogenic nucleosides via palladium-catalysed P-C cross-coupling reactions

Benjamin Whittaker, Manuel de Lera Ruiz, Christopher J. Hayes *



Novel cleavage of (E)-allyl vic-diols to aldehydes using the 2nd-generation Grubbs catalyst Chunguang Han, Daisuke Uemura



Stereospecific epoxide-opening reactions of 1,1-dibromo-3,4-epoxy-1-alkenes with carbon nucleophiles Fumihiko Yoshimura, Masaki Takahashi, Keiji Tanino, Masaaki Miyashita



Imine allylation using 2-alkoxycarbonyl allylboronates as an expedient three-component reaction to polysubstituted pp 6995-6998 α-*exo*-methylene-γ-lactams

Tim G. Elford, Dennis G. Hall *



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En route to color-stable pyranoflavylium pigments—a systematic study of the reaction between 5-hydroxy-4-methylflavylium salts and aldehydes

Stefan Chassaing *, Géraldine Isorez, Marie Kueny-Stotz, Raymond Brouillard



The reaction between 4-methylflavylium cations and aldehydes was investigated and was found to be highly substrate-dependant.



Facile syntheses of 3-halo and mixed 3,5-dihalo analogues of *N*-acetyl-L-tyrosine via sulfonic acid-catalysed regioselective monohalogenation

Pakorn Bovonsombat *, Pratheep Khanthapura, Michael M. Krause, Juthamard Leykajarakul



3-Halo and mixed-3,5-dihalo analogues of N-acetyl-L-tyrosine were obtained at ambient temperature with high yields and selectivities via p-toluenesulfonic acid-catalysed regioselective electrophilic halogenation.

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Divya Tripathi, Pradeep Kumar *



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The concept of internal solubilization in peptide synthesis: ethylene glycol-based protecting groups László Kocsis, Thomas Bruckdorfer, György Orosz

An Fmoc compatible side chain protecting group family is developed, which inhibits peptide chain aggregation. Protected amino acid derivatives can easily be incorporated into existing automated and manual peptide synthesis protocols.



Synthesis of the C12-C24 fragment of peloruside A by silyl-tethered diastereomer-discriminating RCM

Emma M. Casey, Paul Teesdale-Spittle, Joanne E. Harvey



First synthesis of (+)-myxothiazol A

Yuki Iwaki, Masahiro Kaneko, Hiroyuki Akita *



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Et_3 N-Promoted reaction of acetylenic ketones with *N*-(diphenylmethylene)glycinates: an efficient synthesis of α , β -dehydroamino acid derivatives

Qing-Fa Zhou, Quan-Ping Wu, Song Xue *



Monosubstituted 1,2,3-triazoles from two-step one-pot deprotection/click additions of trimethylsilylacetylene James T. Fletcher^{*}, Sara E. Walz, Matthew E. Keeney

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General reaction conditions were developed enabling the regioselective synthesis of 1-substituted-1,2,3-triazoles with both aliphatic and aromatic substitution. Product yields and distributions were sensitive to both alcohol and base identity.

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Three-component synthesis of 2-substituted quinazolin-4(3H)-ones is investigated via rapid decomposition of formamide.

Homogeneous silicone modified primary amine-Brønsted acid salt catalyzed aldol reaction: unexpected synergistic pp 7037–7041 effect of polysiloxane with remarkable improvement of efficiency and stereoselectivity Li-Wen Xu^{*}, Ya-Dong Ju, Li Li, Hua-Yu Qiu, Jian-Xiong Jiang, Guo-Qiao Lai^{*}



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Synthesis and structure revision of tyroscherin, a growth inhibitor of IGF-1-dependent tumor cells Ryo Katsuta, Chié Shibata, Ken Ishigami, Hidenori Watanabe ^{*}, Takeshi Kitahara



Gold-catalyzed transesterification of *ortho*-alkynylbenzoic acid esters: a novel protecting group for alcohols and phenols phenols protecting group for alcohols and phenols phenols

Kazuteru Umetsu, Naoki Asao *



One-dimensional porphyrin H-aggregates induced by solvent polarity Myung-Seok Choi



The linear-shape porphyrin derivatives bearing two pentameric thiophenes showed the significant spectral changes of Soret absorption bands in both blue-shift and band broadening in *n*-hexane, indicating the formation of a relatively larger H-aggregate.

Stereoselective Wittig olefination reactions employing a novel *ortho-P-aryl* **alkoxide effect** James McNulty ^{*}, Kunal Keskar

 $\begin{array}{c} \begin{array}{c} CH_{3} \\ Ph \\ Ph \\ X \end{array} + RCHO \\ (Z):(E) \text{ from 95:5 to 8:92} \\ X = -H, -OMe, -O^{(\cdot)}, -CH_{2}O^{(\cdot)} \end{array}$

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Spiro[1*H*-azulenium-1,9′-fluorene] perchlorate. Intramolecular charge-transfer interaction between orthogonally pp 7058–7061 arranged units of the azulenium cation and fluorene

Mitsunori Oda^{*}, Nobue Nakajima, Nguyen Chung Thanh, Shigeyasu Kuroda



Unprecedented SnCl₂·2H₂O-mediated intramolecular cyclization of nitroarenes via C–N bond formation: a new entry pp 7062–7065 to the synthesis of cryptotackieine and related skeletons

Sunil Sharma, Bijoy Kundu ^{*}



Song Qin, Jing-Yu Liang, Yu-Cheng Gu, Yue-Wei Guo^{*}



Proline-catalyzed facile access to Mannich adducts using unsubstituted azoles Nagarapu Srinivas, Kalpana Bhandari *

> DMSO/60 °C 1) (HCHO)_n 1) (HCHO)_n B 2) Azole 2) Azole Ô (7)n 3) L-proline Base / Normal Mannich solvent conditions (acid or ö n= 0-2 base catalyzed) H₂O/100 °C X= CH; imidazole X= N; 1, 2, 3-triazole

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Highly stereoselective iminopinacol coupling of chiral aromatic imines derived from di- and tripeptides Naoki Kise ^{*}, Takashi Iwasaki, Yuko Yasuda, Toshihiko Sakurai pp 7074-7077



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Corresponding author () Supplementary data available via ScienceDirect

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