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Contents

COMMUNICATIONS

Comparison of microwave-assisted and conventional preparations of cyclic imides Sunil K. Upadhyay, Subramanya R. K. Pingali, Branko S. Jursic*



Microwave-assisted preparation of cyclic imides is superior with respect to isolated yield and length of reaction time when compared to conventional methods of preparation of imides.

Effect of fluorine or oxygen atom(s) in propargylic position on the reactivity in click chemistry Danielle Grée, René Grée*



Newly designed w-diynes allow to establish, through competitive reactions, the effect of C-F, C-OH, CF₂, C=O and C(OMe)₂ substituents on the reactivity of neighbouring triple bonds in click chemistry.

Synthesis of 2-acetamido-1,2-dideoxy-p-galacto-nojirimycin [DG]NAc] from p-glucuronolactone: the first sub-micromolar inhibitor of α-N-acetylgalactosaminidases

Daniel Best, Phoom Chairatana, Andreas F. G. Glawar, Elizabeth Crabtree, Terry D. Butters, Francis X. Wilson, Chu-Yi Yu, Wu-Bao Wang, Yue-Mei Jia, Isao Adachi, Atsushi Kato, George W. J. Fleet*



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Novel oxidation of toluenes catalyzed by reusable vanadyl(IV) sulfate under mild conditions with molecular oxygen

Takeo Nakai*, Toshiyuki Iwai, Masatoshi Mihara, Takatoshi Ito, Takumi Mizuno



Efficient oxidation system using vanadyl(IV) sulfate catalyst with molecular oxygen was established. Recovered catalyst could be reused without loss of activity.

Real-time colorimetric screening of endopeptidase inhibitors using adenosine triphosphate (ATP)-stabilized gold pp 2228–2231 nanoparticles

Mi Hee Kim, Soo Suk Lee, Sang J. Chung, Hyun Hye Jang, Sujung Yi, Sudeok Kim, Suk-Kyu Chang, Min Su Han*



Palladium (II) catalyzed 5-*endo* **epoxynitrile cyclizations: total syntheses of enokipodins A and B** Jesús Armando Luján-Montelongo, José G. Ávila-Zárraga*



Stereoselective synthesis of selenosteroids

Oscar E. D. Rodrigues^{*}, Diego de Souza, Letiére C. Soares, Luciano Dornelles, Robert A. Burrow, Helmoz R. Appelt, Camila F. Alves, Diego Alves^{*}, Antonio L. Braga



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 (\mathbf{J}^{+})

Palladium-catalyzed synthesis of carbazoles from *N***-(2-halophenyl)-2,6-diisopropylanilines via C–C cleavage** Anthony R. Chianese^{*}, Scott L. Rogers, Hanna Al-Gattas



A novel practical cleavage of tert-butyl esters and carbonates using fluorinated alcohols

Jason Choy, Saul Jaime-Figueroa*, Teresa Lara-Jaime



The thermolytic cleavage of *t*-butyl esters and *t*-butyl carbonates was accomplished using TFE (2,2,2-trifluoroethanol) or HFIP (hexafluoroisopropanol) as solvent, several examples are presented.

PIDA-mediated synthesis of oxazoles through oxidative cycloisomerization of propargylamides

Akio Saito*, Asami Matsumoto, Yuji Hanzawa*



Gold-catalyzed intermolecular [4C+3C] cycloaddition reactions

Benjamin W. Gung*, Lauren N. Bailey, Josh Wonser

+ $5 \mod \% 7$ 13 + $5 \mod \% 7$ $5 \mod \% AgSbF_6$ $CH_2Cl_2, r.t.$ 95%14 + minor isomers

In the presence of the N-heterocyclic carbene gold catalyst (NHC-AuIPr, **7**), propargyl esters **1a–f** and **13** undergo a [4C+3C] cycloaddition reaction with cyclopentadiene and furan under mild conditions. The evidence suggests that the formation of the seven-membered ring occurs by a direct cycloaddition process, rather than a stepwise cyclopropanation/Cope rearrangement sequence.

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Ring opening of aziridines with tetranitromethane in the presence of triethylamine. Efficient synthesis of **β-tosylamino** nitrates

Yuliya A. Volkova, Elena B. Averina*, Tamara S. Kuznetsova, Nikolai S. Zefirov



Preparation, characterization and application of a stationary chromatographic phase from a new (+)-tartaric acid pp 2258-2261 derivative

Sacha Legrand*, Harri Heikkinen, Ian A. Nicholls, Andrew Root, Johan Svenson, C. Rikard Unelius

Synthesis of 9-alkyl-6-amino[1,2,4]triazolo[3,4-c]-5-azaquinoxalines. Mild and effective S_NAr amination of highly pp 2262-2264 electron-poor heterocycles

O OMe

Asier Unciti-Broceta*, María José Pineda-de-las-Infantas, Miguel Ángel Gallo, Antonio Espinosa



Due to the notable electrophilic character of the C-6 position of the [1,2,4]triazolo[3,4-c]-5-azaquinoxaline tricyclic system, direct S_NAr amination was performed by reacting the corresponding 6-chloro derivative with ammonia-saturated acetonitrile in a sealed reaction vessel, using microwave-mediated or conventional heating.

The ionic liquid [bmim]Br as an alternative medium for the catalytic cleavage of aromatic C-F and C-Cl bonds Sergey A. Prikhod'ko, Nicolay Yu. Adonin*, Valentin N. Parmon





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A novel cascade Kröhnke condensation—an intramolecular nucleophilic cyclization approach toward annulated chromenes

Leonid G. Voskressensky^{*}, Larisa N. Kulikova, Anna V. Listratova, Roman S. Borisov, Muhamadsho A. Kukaniev, Alexey V. Varlamov



Cycloaddition of benzynes and nitrile oxides: synthesis of benzisoxazoles James A. Crossley, Duncan L. Browne*



A simple and effective synthesis of activated vinylphosphonates from 3-methoxy-2-diethoxyphosphorylacrylate pp 2274–2276 Tomasz Janecki^{*}, Anna Albrecht, Jacek F. Koszuk, Jakub Modranka, Dominika Słowak



when NuH = 3,5-dimethoxyphenol

PS-SNAP, a practical polymer-supported nitrosation reagent in organic synthesis Didier Roche^{*}, Claude Lardy, Lucie Tournier, Marc Prunier, Eric Valeur

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Palladium-catalyzed benzylation of N-Boc indole boronic acids

Aaron M. Kearney, Adrienne Landry-Bayle, Laurent Gomez*



Copper-promoted rearrangement of 1,3-cyclohexadiene-acylnitroso cycloadducts Stefano Crotti, Ferruccio Bertolini, Franco Macchia, Mauro Pineschi*



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Carbon tether rigidity as a stereochemical tool directing intramolecular radical cyclizations

Gagik G. Melikyan^{*}, Erin Voorhees, Christopher Wild, Ryan Spencer, Justin Molnar



The phenyl group incorporated into a carbon tether provides for the synthesis of 1,5-cyclodecadiynes with 95–100% d_i -diastereoselectivity due to intrinsic conformational constraints and preorganization of the Co₂(CO)₆-complexed propargyl intermediates.

The allylic nucleophilic substitution of Morita–Baylis–Hillman acetates with isocyanides: a facile synthesis of trisubstituted olefins

J. S. Yadav*, B. V. Subba Reddy, Ashutosh Pratap Singh, Nilanjan Majumder



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PTSA-catalyzed tandem cyclization protocol for the stereoselective total synthesis of obolactone

Palakodety Radha Krishna*, Palabindela Srinivas



A stereoselective total synthesis of obolactone is reported.

Catalyst-free regioselective synthesis of benzopyran-annulated thiopyrano[2,3-*b*]thiochromen-5-(4*H*)-one derivatives by domino-Knoevenagel-hetero-Diels–Alder reaction of terminal alkynes with 4-hydroxy dithiocoumarin in aqueous medium

K. C. Majumdar*, Abu Taher, Sudipta Ponra



Strategic utilization of catalytic metathesis and photo-thermal metathesis in caged polycyclic frames Sambasivarao Kotha^{*}, Vittal Seema, Kuldeep Singh, Kodand Dinkar Deodhar



Domino Knoevenagel-hetero-Diels-Alder reactions: a stereoselective synthesis of sugar-annulated furo[3,2-b] pyrano[4,3-d]pyran derivatives

J. S. Yadav*, B. V. Subba Reddy, A. V. Hara Gopal, R. Nageshwar Rao, R. Somaiah, P. Purushotham Reddy, A. C. Kunwar



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Fe nano particles mediated C-N bond-forming reaction: Regioselective synthesis of 3-[(2-chloroquinolin-3-yl)methyl]pyrimidin-4(3H)ones

Selvaraj Mohana Roopan, Fazlur Rahman Nawaz Khan*, Badal Kumar Mandal



O₂Me

A regio and diastereoselective transformation of 3-dienyl-2-azetidinones to novel pyrroloxazine Amit Anand, Gaurav Bhargava, Vipan Kumar, Mohinder P. Mahajan*



Pan-Lin Shao, Xiang-Yu Chen, Li-Hui Sun, Song Ye*



Goutam Brahmachari*, Sujay Laskar



A very simple, highly efficient and cost-effective method for N-formylation of primary and secondary amines in excellent yield using catalytic amount of sodium formate in formic acid under solvent-free conditions is reported.

OH (10 mol%) toluene, 0°C or rt up to 96% ee

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New polymer-supported phosphonate reagents for the synthesis of Z- α , β -unsaturated esters

Kaori Ando*, Yusaku Suzuki



New polymer-supported phosphonate reagents have been prepared and evaluated for the synthesis of *Z*-α,β-unsaturated esters. High *Z*-selectivity was obtained using the reagent having two *o*-*t*-BuC₆H₄ groups.

$Dy(OTf)_3/Pybox-catalyzed enantioselective Friedel-Crafts alkylation of indoles with <math>\alpha,\beta$ -unsaturated trifluoromethyl ketones

Shigeru Sasaki, Takayasu Yamauchi, Kimio Higashiyama*



The first catalytic enantioselective Friedel–Crafts alkylation of indoles with α , β -unsaturated trifluoromethyl ketones has been accomplished using the Dy(OTf)₃/Pybox complex.

Intramolecular Pauson-Khand reaction of optically active aza-Baylis-Hillman adducts

Shingo Ishikawa, Fumiaki Noguchi, Hidemitsu Uno, Akio Kamimura*



The intramolecular Pauson-Khand reaction of aza-Baylis-Hillman adducts, which were prepared through the thio-Michael/imino-aldol domino reaction of optically active sulfinimines, was examined.

Addition of alkyl radicals, generated from carboxylic acids via photochemical decarboxylation, to glyoxylic oxime pp 2332–2334 ether: a mild and efficient route to α-substituted α-aminoesters

Yasuharu Yoshimi*, Kosuke Kobayashi, Hayato Kamakura, Keisuke Nishikawa, Yoshiki Haga, Kousuke Maeda, Toshio Morita, Tatsuya Itou, Yutaka Okada, Minoru Hatanaka*

$$R - C - OH + BnON CO_{2}CH_{3} \xrightarrow{\text{Phen, DCB}} BnOHN CO_{2}CH_{3}$$

$$R - C - OH = primary, secondary, tertiary alkyl carboxylic acids, gulonic acid, N-Boc α -amino acids$$

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An expedient synthesis of ellipticine via Suzuki-Miyaura coupling

Takeo Konakahara*, Y. B. Kiran, Yuri Okuno, Reiko Ikeda, Norio Sakai





Synthesis of 2-substituted quinones, vitamin K₃, and vitamin K₁ from *p*-cresol. BF₃·OEt₂-catalyzed methyl migration of 4-tert-butyldioxycyclohexadienones

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Shun-Ichi Murahashi*, Akiko Fujii, Yasutaka Inubushi, Naruyoshi Komiya



o-Benzenedisulfonimide as a reusable Brønsted acid catalyst for an efficient and facile synthesis of quinolines via pp 2342-2344 Friedländer annulation

Margherita Barbero, Stefano Bazzi, Silvano Cadamuro, Stefano Dughera*



o-Benzenedisulfonimide as catalyst in quinoline synthesis.

A BINOL-terpyridine-based multi-task catalyst for a sequential oxidation and asymmetric alkylation of alcohols pp 2345-2347 Xi Chen, Qiang Liu, Hong-Bao Sun, Xiao-Qi Yu*, Lin Pu*

 $\mathsf{R}-\mathsf{CH}_2\mathsf{OH} \xrightarrow{(R)-\mathbf{6}} [\mathsf{R}-\mathsf{CHO}] \xrightarrow{\mathsf{Ti}(\mathsf{O}^{\mathsf{i}}\mathsf{Pr})_4} \mathsf{R} \xrightarrow{\mathsf{OH}} \mathsf{R}$ ОΗ ΩН (R)-6

Exploring the *one-pot* C-acylation of cyclic 1,3-diones with unactivated carboxylic acid

Sylvie Goncalves, Marc Nicolas, Alain Wagner, Rachid Baati*



The use of DCC/Et₃N/4-DMAP/DCM provides a general and standard *one-pot* procedure for the smooth C-acylation of cyclic 1,3-diones with unactivated carboxylic acids, giving rise to β -triketones.

Effective syntheses of per-2,3-di- and per-3-O-chloroacetyl-β-cyclodextrins: A new kind of ATRP initiators for star pp polymers

Zhizhang Guo, Xingyu Chen, Xiao Zhang, Jianyu Xin, Jianshu Li*, Huining Xiao*



Stereoselective formal synthesis of (–)-mesembrane by intramolecular condensation of chiral amide and 1,3-cyclohexanedione moiety

Le Anh Tuan, Guncheol Kim*

Conversion of cellulose to glucose and levulinic acid via solid-supported acid catalysis Jessica Hegner, Kyle C. Pereira, Brenton DeBoef^{*}, Brett L. Lucht^{*}





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Rapid access to morphinones: removal of 4,5-ether bridge with Pd-catalyzed triflate reduction

Christopher D. Hupp, John L. Neumever*



Chemoselective sulfide oxidation mediated by bridged flavinium organocatalysts

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Barrie J. Marsh, David R. Carbery*



The chemoselective oxidation of sulfides to sulfoxides, catalysed by bridged, tetracyclic flavinium catalysts is presented. The flavinium catalysts are easily prepared via a telescoped three-step process. A range of sulfoxides is accessed in excellent yield and chemoselectivity.

Highly efficient dynamic kinetic resolution of secondary aromatic alcohols with low-cost and easily available acid pp 2366-2369 resins as racemization catalysts

Yongmei Cheng, Gang Xu, Jianping Wu, Chensheng Zhang, Lirong Yang*



Efficient synthesis of triarylmethanes via bisarylation of aryl aldehydes with arenes catalyzed by silica gel-supported sodium hydrogen sulfate

Yixin Leng, Fang Chen, Li Zuo*, Wenhu Duan*



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Lewis acid-catalyzed one-pot sequential reaction for the synthesis of α -halogenated β -keto esters Han-Feng Cui, Ke-Yan Dong, Jing Nie, Yan Zheng, Jun-An Ma^{*}

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*Corresponding author

(*D*⁺ Supplementary data available via ScienceDirect

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