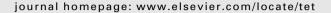


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Tetrahedron





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Contents

REPORT

Radicals in organic synthesis. Part 1

pp 8603-8655

Gareth J. Rowlands

This report offers the first part of a two part overview of recent advances in radical reactions in organic synthesis. It includes an introduction to common radical reagents and outlines the use of radicals in intermolecular additions. The report includes over 300 references.

ARTICLES

$Synthesis\ of\ amine,\ halide,\ and\ pyridinium\ terminated\ 2-alkyl-\emph{p-tert-}butyl calix [4] are nes$

pp 8657-8667

Matthew P. Hertel, Andrew C. Behrle, Samantha A. Williams, Joseph A.R. Schmidt^{*}, Jordan L. Fantini^{*}

Calix[4]arenes that have been mono-lithiated at a methylene bridge react with 1, ω -bromochloroalkanes to give 2-(ω -chloroalkyl)-calix[4]arenes. Subsequent reactivity of these species has allowed for successful synthesis of 2-(ω -iodoalkyl)-calix[4]arenes, 2-(ω -pyridiniumalkyl)-calix[4]arenes, and 2-(ω -aminoalkyl)-calix[4]arenes.

28 examples

$Stereo controlled \ one-pot \ synthesis \ of \ cycloal kane \ derivatives \ possessing \ a \ quaternary \ carbon \ using \ allyl \ phenyl \ sulfone$

pp 8668-8676

Koichiro Ota, Takao Kurokawa, Etsuko Kawashima, Hiroaki Miyaoka*

Ph SO₂Ph SO₂Ph SO₂Ph
$$\frac{^{n}\text{BuLi, THF,}}{^{-78}\,^{\circ}\text{C to } -20\,^{\circ}\text{C}}$$
 Ph $\frac{^{n}\text{BuLi, Me}_{3}\text{Al, 84\%}}{^{n}\text{BuLi, Me}_{3}\text{Al, 84\%}}$ OH α -sulfone : β -sulfone = 3.7 : 1

A facile approach towards synthesis of verbalactone and biologically active δ -lactones using p-glucose Ashish Garg, Vinod K. Singh*

pp 8677-8682

A general synthetic approach has been developed for the asymmetric synthesis of chiral δ -lactones and verbalactone using p-glucose. The key intermediate used in this approach was α -diazoketone.

One-pot, solvent-free regioselective addition reactions of propargyl bromide to carbonyl compounds mediated by Zn-Cu couple

pp 8683-8689

Xiaofang Ma, Jin-Xian Wang*, Shunxi Li, Ke-Hu Wang, Danfeng Huang

OH
$$R^{1} \longrightarrow R^{2} + P$$

$$R^{2} \longrightarrow R^{2} \longrightarrow R^{1} \longrightarrow R^{2} \longrightarrow R^{1} \longrightarrow R^{2} \longrightarrow R^{2}$$

R¹ = Ph, Aryl, Heterocyclic, Aliphatic; R² = H, Me, Et.

A simple approach to pyrazol-3-ones via diazenes

pp 8690-8696

Bojan Burja, Marijan Kočevar, Slovenko Polanc*

Synthesis of all four stereoisomers of 5-hydroxy-4-methyl-3-heptanone using plants and oyster mushrooms $Bj\ddot{o}rn\ Bohman,\ C.\ Rikard\ Unelius^*$

pp 8697-8701

Synthesis of substituted 2-aroyl-3-methylchromen-4-ones from isovanillin via 2-aroyl-3-methylchroman intermediates

pp 8702-8707

Sie-Rong Li, Chung-Jung Shu, Liang-Yeu Chen, Hsing-Ming Chen, Po-Yuan Chen, Eng-Chi Wang*

$$\begin{array}{c} \text{OH} \\ \text{CH}_3\text{O} \\ \text{CHO} \\ \end{array} \begin{array}{c} \text{CH}_3\text{O} \\ \text{OCH}_3 \\ \text{CH}_3\text{O} \\ \end{array} \begin{array}{c} \text{CH}_3\text{O} \\ \text{OCH}_3 \\ \text{OCH}_3 \\ \text{OCH}_3 \\ \end{array} \begin{array}{c} \text{CH}_3\text{O} \\ \text{OCH}_3 \\ \text{OCH}_3 \\ \text{OCH}_3 \\ \end{array} \begin{array}{c} \text{CH}_3\text{O} \\ \text{OCH}_3 \\$$

Intermediate 2-aroyl-3-methylchromans prepared from isovanillin through a carbanion-olefin 6-endo-trig cyclization reaction were oxidized with DDQ to yield the title compounds, in good over-all yields.

Facile synthesis of 2-unsubstituted benzofuran-3-carboxylates using diazo(trimethylsilyl)methylmagnesium bromide Yoshiyuki Hari*, Ryosuke Kondo, Koji Date, Toyohiko Aoyama*

pp 8708-8713

$$\begin{array}{c|c} R & CO_2Bu^t \\ \hline CO_2Bu^t & TMSC(MgBr)N_2 \\ \hline OMe & OMe \\ \end{array}$$

Addition of kinetic boron enolates generated from β -alkoxy methyl ketones to aldehydes. Density functional theory calculations on the transition structures

pp 8714-8721

Luiz C. Dias*, Sávio M. Pinheiro, Vanda M. de Oliveira, Marco A.B. Ferreira, Cláudio F. Tormena, Andrea M. Aguilar, Julio Zukerman-Schpector, Edward R.T. Tiekink



Heterocyclic synthesis using nitrilimines. Part 13: Synthesis of new 1,2,3,4-tetrahydro-s-tetrazine derivatives Hany M.M. Dalloul

pp 8722-8726

Synthesis of polyamino nitriles, en route to acylpolyamine neurotoxins, via the regioselective michael cyanoethylation of unprotected polyamines. Unusual behaviour of 1-(2-aminoethyl)piperazine

pp 8727-8732

Piotr Bałczewski*, Remigiusz Żurawiński, Maciej Mikina, Bogdan Dudziński

$$NH_2$$
 or NH_2 NH_2

AlCl₃-Nal(NaBr)-t-BuOH: mild, chemo- and stereoselective reagents for hydrohalogenation of propiolic derivatives

pp 8733-8737

Laurence Feray*, Patricia Perfetti, Michèle Bertrand*

1) AlCl₃ (1.5 equiv)

NaX (2 equiv)

$$X = I, Br$$
alcohol :solvent (1:9)

2) H_2O
 $Y = NR^1R^2$
 $Y = Ph$

Convenient and versatile synthesis of formyl-substituted benzoxaboroles

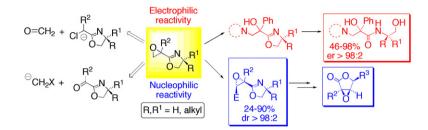
pp 8738-8744

Long Ye, Dazhong Ding, Yiqing Feng, Dongsheng Xie, Puhua Wu, Hui Guo, Qingqing Meng, Huchen Zhou*

Terminal oxazolinyloxiranes: synthesis, reaction with amines and regioselective β -lithiation

pp 8745-8755

Leonardo Degennaro, Vito Capriati, Claudia Carlucci, Saverio Florio*, Renzo Luisi, Irene Nuzzo, Corrado Cuocci





Synthesis and chromatographic resolution of conformationally constrained analogues of homotaurine

Maria Carmela Fulco, Maura Marinozzi, Burcu Çaliskan Ergün, Roccaldo Sardella, Benedetto Natalini, Roberto Pellicciari*

pp 8756-8762

*Corresponding author

(i) + Supplementary data available via ScienceDirect



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