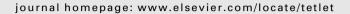


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Tetrahedron Letters





Tetrahedron Letters Vol. 50, No. 41, 2009

Contents

COMMUNICATIONS

The first total synthesis of (±)-4-methoxydecanoic acid: a novel antifungal fatty acid

Néstor M. Carballeira *, Carlos Miranda, Keykavous Parang

pp 5699-5700

The hitherto unknown (±)-4-methoxydecanoic acid (1) was synthesized in six steps and in 25% overall yield starting from 4-penten-1-ol. The title compound demonstrated 17-fold higher antifungal activity (MIC = 1.5 mM) against *Candida albicans* ATCC 60193 and *Cryptococcus neoformans* ATCC 66031 when compared to unsubstituted *n*-decanoic acid. Mid-chain methoxylation appears to be a viable strategy for increasing the fungitoxicity of fatty acids.

A gold-catalyzed [4+3]-cycloaddition of functionalized dioxines

pp 5701-5703

Michael Harmata*, Chaofeng Huang

Treatment of 5-silyloxydioxines with 5 mol % AuCl $_3$ /AgSbF $_6$ in the presence of cyclopentadiene or furan resulted in the rapid formation of [4+3]-cycloadducts at room temperature.



An expedient one-pot synthesis of novel 10-substituted 9-aminophenanthrenes

Christophe Rochais, Rodrigue Yougnia, Patrick Dallemagne *, Sylvain Rault

pp 5704-5708

An efficient synthesis of 9,10-disubstituted phenanthrenes is described. These novel useful building blocks were obtained in a one-pot reaction including Suzuki-Miyaura cross-coupling followed by a Dieckmann-Thorpe ring closure under microwave irradiation.

Convenient synthesis of novel 2,2-dialkyl-1,2-dihydronaphtho[2,1-b]furans

Doug Vaughan, Amitabh Jha *

pp 5709-5712

pp 5713-5715

A concise method for the synthesis of 2-tetralone by titanium tetrachloride-promoted cyclization of 4-aryl-2-hydroxybutanal diethyl acetal

Yung-Son Hon*, Rammohan Devulapally

$$R^{2} \xrightarrow[R^{4}]{R^{5}} \underbrace{\begin{array}{c} 1. \text{ Rieke Mg, THF;} \\ 2. \text{ oxirane 4,} \\ \text{THF} \\ \text{CI} \\ \text{R} \xrightarrow[]{CH(OEt)_{2}} \\ \text{R}^{3} \\ \text{R} = \text{H, Me} \end{array}}_{\text{R}} R^{1} \underbrace{\begin{array}{c} R^{5} \\ R^{5} \\ \text{R}^{4} \\ \text{OEt} \end{array}}_{\text{R}^{5} \underbrace{\begin{array}{c} R^{1} \\ \text{CI} \\ \text{CH}_{2}\text{Cl}_{2}(0.05 \text{ M}),} \\ \text{OH} \\ \text{OEt} \\ \text{OEt} \\ \text{OEt} \\ \text{R}^{3} \xrightarrow[]{R^{4}} \underbrace{\begin{array}{c} R^{1} \\ R^{5} \\ \text{R}^{4} \\ \text{OE} \\ \text{OET} \\ \text{OE$$

Direct 1,4-difunctionalization of isoquinoline

pp 5716-5718

Frédéric Louërat, Yves Fort, Victor Mamane

A one-pot procedure comprising a nucleophilic addition followed by an electrophilic trapping is performed in order to functionalize isoquinoline in 1- and 4-positions.



$Synthetic\ studies\ toward\ 1, 2-dioxanes\ as\ precursors\ of\ potential\ endoperoxide-containing\ antimalarials$

Sandra Gemma, Francesc Martí, Emanuele Gabellieri, Giuseppe Campiani *, Ettore Novellino, Stefania Butini

pp 5719-5722

R Me R¹ 6a, R = Me, R¹ = Et, R² = Ac 6b, R = Me, R¹ =
$$n$$
-Pr, R² = Ms OR² (6 S , R)-6c, R = Et, R¹ = H, R² = Ac (6 S , R)-6d, R = n -Pr, R¹ = H, R² = Ac

Synthesis of (+)-striatene: confirmation of its stereostructure

Paul Brémond, Nicolas Vanthuyne, Gérard Audran

pp 5723-5725

Synthesis of functionalized diaryl selenides by the first [3+3] cyclocondensations of 1,3-bis(silyloxy)-1,3-butadienes with organoselenium compounds

pp 5726-5728

Mohanad Shkoor, Olumide Fatunsin, Abdolmajid Riahi, Alexander Villinger, Peter Langer

$A\ convenient\ synthesis\ of\ 5\hbox{-}ary lamino-4 \textit{H-pyran-4-ones}\ using\ palladium-catalyzed\ amination$

pp 5729-5732

Julien Farard, Cédric Logé, Bruno Pfeiffer, Brigitte Lesur, Muriel Duflos

A concise approach to 5-arylamino-4H-pyran-4-ones is described via palladium-catalyzed amination reaction.



Hydrogen peroxide mediated formation of heteroaryl ethers from pyridotriazol-1-yloxy heterocycles and arylboronic acids

pp 5733-5736

Sujata Bardhan, Keiko Tabei, Zhao-Kui Wan, Tarek S. Mansour *



Aqueous Wittig reactions of semi-stabilized ylides. A straightforward synthesis of 1,3-dienes and 1,3,5-trienes James McNulty *, Priyabrata Das

pp 5737-5740

$$Et_3P$$
 + Br \longrightarrow Et_3P - $CH_2CH=CH_2$ $\xrightarrow{NaOH, H_2O}$ R -

 Br $RCHO$ $Sa-5t$

Phenyl esters, preferred reagents for mono-acylation of polyamines in the presence of water

pp 5741-5743

Kyrie Pappas, Xiang Zhang, Wei Tang, Shiyue Fang

An expedient synthesis of poly-substituted pyrroles from γ -ketonitriles via indium-mediated Barbier reaction strategy

pp 5744-5747

Sung Hwan Kim, Se Hee Kim, Ka Young Lee, Jae Nyoung Kim *

A novel carbanion-olefin intramolecular cyclization: synthesis of substituted 2-aroyl-3,4-dihydro-2*H*-benzopyrans from salicylaldehydes

pp 5748-5750

Liang-Yeu Chen, Sie-Rong Li, Po-Yuan Chen, Ian-Lih Tsai, Chia-Ling Hsu, Hsin-Ping Lin, Tzu-Pin Wang, Eng-Chi Wang

$$\begin{array}{c} R_1 \\ \downarrow CHO \\ R_2 \\ \downarrow CHO \\ OH \end{array} \longrightarrow \begin{array}{c} R_1 \\ \downarrow CHO \\ R_2 \\ \downarrow CHO \\ R_3 \\ O \end{array} \longrightarrow \begin{array}{c} R_4 \\ \downarrow CHO \\ R_2 \\ \downarrow CHO \\ R_3 \\ O \end{array} \longrightarrow \begin{array}{c} R_4 \\ \downarrow CHO \\ R_2 \\ \downarrow CHO \\ R_3 \\ O \end{array} \longrightarrow \begin{array}{c} R_4 \\ \downarrow CHO \\ R_2 \\ \downarrow CHO \\ R_3 \\ O \end{array} \longrightarrow \begin{array}{c} R_4 \\ \downarrow CHO \\ \downarrow CHO$$

Through a sequence of a Wittig reaction, O-alkylation, and carbanion-olefin intramolecular cyclization, salicylaldehydes were converted into a series of new 2-aroyl-3,4-dihydro-2*H*-benzopyrans in two steps or in one-pot reaction.



Synthesis of 9-oxa-noradamantane derivative, an aesthetically pleasing 'oxa-basket'

pp 5751-5753

Faiz Ahmed Khan *, Ch. Nageswara Rao

Dichloro bis- γ -lactone readily undergoes acid mediated mixed ketal formation to furnish a 9-oxa-noradamantane derivative. Replacement of chlorines with hydrogens or allyl groups results in the formation of hydrolysis product rather than cyclization.

Halohydroxylation of alkylidenecyclopropanes using *N*-halosuccinimide (NXS) as the halogen source: an efficient synthesis of halocyclopropylmethanol and 3-halobut-3-en-1-ol derivatives

pp 5754-5756

Yewei Yang, Chenliang Su, Xian Huang *, QingYang Liu

A variety of halocyclopropyl-methanol and 3-halo-but-3-en-1-ol derivatives were prepared in moderate to excellent yields via the simple halohydroxylation reaction of alkylidenecyclopropanes with convenient and mild sources of electrophilic halogen NXS (X = I, Br, CI, F) and H_2O . A plausible mechanism for the halohydroxylation has been proposed.



Arylations of allylic acetates with triarylbismuths as atom-efficient multi-coupling reagents under palladium catalysis

pp 5757-5761

Maddali L. N. Rao *, Debasis Banerjee, Somnath Giri

OAc
$$R^1$$
 R^2 R^2 R^2 R^3 R^4 R^2 R^2 R^2 R^2 R^2 R^2

Chemo- and regioselective halogenation of 4-(pyrazol-4-yl)-pyrimidines

pp 5762-5764

Young Shin Cho^{*}, Ying Hou, Christine H.-T. Chen, Moo Je Sung

 $A convenient \ and \ selective \ halogenation \ of \ 4-(pyrazol-4-yl)-pyrimidines \ is \ described \ herein. \ This \ method \ allows \ quick \ access \ to \ a \ diverse \ set \ of \ pyrazolyl-pyrimidine \ derivatives.$

*Corresponding author

** Supplementary data available via ScienceDirect

Abstracted/indexed in: AGRICOLA, Beilstein, BIOSIS Previews, CAB Abstracts, Chemical Abstracts, Chemical Engineering and Biotechnology Abstracts, Current Biotechnology Abstracts, Current Contents: Life Sciences, Current Contents: Physical, Chemical and Earth Sciences, Current Contents Search, Derwent Drug File, Ei Compendex, EMBASE/Excerpta Medica, Medline, PASCAL, Research Alert, Science Citation Index, SciSearch. Also covered in the abstract and citation database SCOPUS[®]. Full text available on ScienceDirect[®]



