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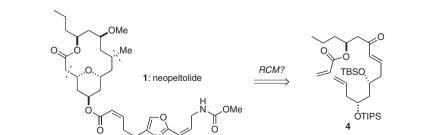
Liebeskind-Srogl cross coupling mediated synthesis of verbenachalcone pp 5753-5756 Srinivasa Reddy Dandepally, Alfred L. Williams* OH SR CO₂H 2 steps MOMO HO nн 6 stens OH нΟ SR MeO MeO) 0 ö ÓН Verbenachalcone A flexible and scalable total synthesis of verbenachalcone is achieved in eight linear steps from commercially cheap available starting material, 3-(4-hydroxyphenyl)propanoic acid.

Synthesis of hemigossypol and its derivatives

Jun Wei, David L. Vander Jagt, Robert E. Royer, Lorraine M. Deck*

Studies towards the synthesis of neopeltolide: synthesis of a ring-closing metathesis macrocyclization precursor pp 5761-5763 Gordon J. Florence*, Romain F. Cadou

HO HO CHO OH





COMMUNICATIONS

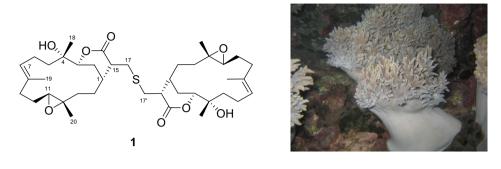






A novel symmetric sulfur-containing biscembranoid from the Formosan soft coral Sinularia flexibilis

Bo-Wei Chen, Chih-Hua Chao, Jui-Hsin Su, Chiung-Yao Huang, Chang-Feng Dai, Zhi-Hong Wen, Jyh-Horng Sheu*



 $\begin{array}{c} & & & \\ & & & & \\ & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\$

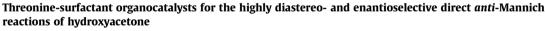
R¹ CuI, L-proline NaIO₄, AcOH, 80 °C

Electroreductive intramolecular coupling of aliphatic cyclic imides with ketones and O-methyloximes Naoki Kise*, Kazuaki Fukazawa, Toshihiko Sakurai

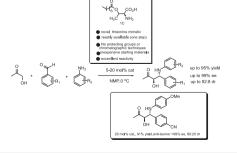
Cul/L-proline-catalyzed selective one-step mono-acylation of styrenes and stilbenes

m = 1,2; n = 1~3

P. Sai Prathima, C. Uma Maheswari, K. Srinivas, M. Mohan Rao*



Chuanlong Wu, Xiangkai Fu*, Xuebing Ma, Shi Li, Chao Li



ОН

R¹=H (or) Ar 14 Examples Upto 85% yield

ÓAc

pp 5767-5770

pp 5771-5774

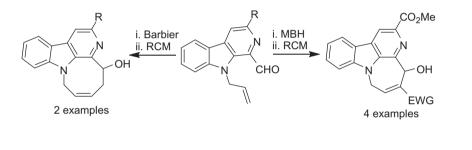
pp 5764-5766

Enantioselective Darzens reaction using organoselenide–lithium hydroxide complexes Shin-ichi Watanabe^{*}, Risa Hasebe, Jun Ouchi, Hideko Nagasawa, Tadashi Kataoka

Ph Br + RCHO $\xrightarrow{\text{Catalyst (0.3 eq)}}_{\text{LiOH}}$ R $\xrightarrow{\text{O}}$ Ph catalyst = CHCl₃ rt, 5 h up to 62% ee

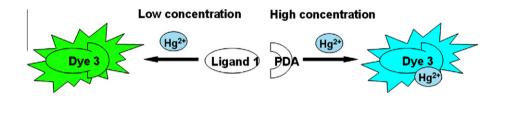
RCM-based approach to seven- and eight-member ring-fused $\beta\text{-carbolines}$

Samiran Hutait, Sanjay Batra*



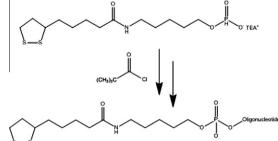
A new fluorogenic chemodosimetric system for mercury ion recognition

Wen Xiu Ren, Sankarprasad Bhuniya, Jun Feng Zhang, Young Hoon Lee, Suk Joong Lee, Jong Seung Kim*



Thioctic acid modification of oligonucleotides using an H-phosphonate Jennifer A. Dougan, Andrew. K. Reid, Duncan Graham*

ennifer A. Dougan, Andrew. K. Reid, Duncan Granam



A convenient route for 5'-modification of an oligonucleotide with a cyclic disulfide, thioctic acid, is reported using the H-phosphonate method.

pp 5778-5780

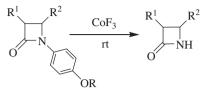
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Synthesis of N-unsubstituted β-lactams from N-alkoxyphenyl-β-lactams with cobalt(III) fluoride

Maaroof Zarei, Aliasghar Jarrahpour*



Stereospecific benzylic dehydroxyfluorination reactions using Bio's TMS-amine additive approach with challenging pp 5795-5797 substrates

DAST, Deoxo-Fluor™

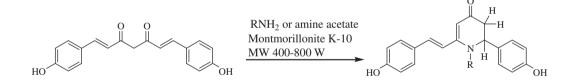
Stefano Bresciani, David O'Hagan*

Stereospecific benzylic fluorinations are carried out on particularly challenging substrates.

Synthesis of novel 2,3-dihydro-4-pyridinones from bisdemethoxycurcumin under microwave irradiation

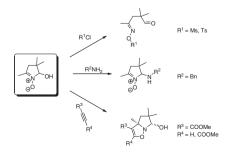
т́мs ŌН





A novel synthesis of 2,3-dihydro-4-pyridinones via the reaction of bisdemethoxycurcumin and primary amines or amine acetates is demonstrated.

Diverse chemical behaviour of 2-hydroxy-functionalized pyrroline-1-oxide Marian Buchlovič, Stanislav Man, Milan Potáček*



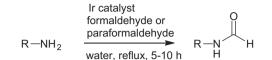


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Iridium-catalyzed formylation of amines with paraformaldehyde

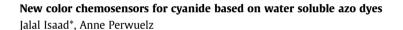
Ourida Saidi, Mark J. Bamford, A. John Blacker, James Lynch, Stephen P. Marsden, Pawel Plucinski, Robert J. Watson, Jonathan M. J. Williams*

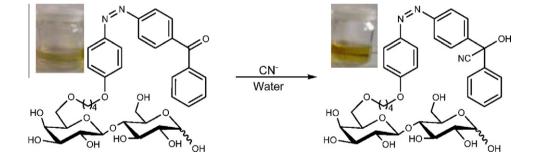


The reaction of amines with either formaldehyde or paraformaldehyde in water in the presence of $[Cp*IrI_2]_2$ affords the corresponding formamides in good yields.

A novel one-pot and efficient procedure for the synthesis of 3*H*-spiro[isobenzofuran-1,6′-pyrrolo[2,3-*d*]pyrimidine]-2′,3,4′,5′-tetraones

Mohammad Reza Mohammadizadeh*, Mojtaba Bahramzadeh, S. Zainabkhatoon Taghavi

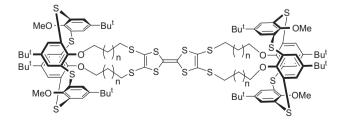




1. AcOH, r.t., 1 h

2. Pb(OAc)₄ r.t. 4 h

Synthesis and electrochemical behavior of a model redox-active thiacalix[4]arene-tetrathiafulvalene assembly Bang-Tun Zhao*, Zhen Zhou, Zhen-Ning Yan, Esmah Belhadj, Franck Le Derf, Marc Sallé*



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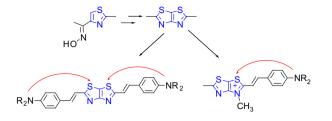
pp 5815-5818

 $(\mathbf{j})^{\dagger}$

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Thiazolo[4,5-d]thiazole—a new domain for potential optoelectronic application

Peter Zahradník, Peter Magdolen^{*}, Pavol Zahradník



A new route to 2,5-dimethyl thiazolo [4,5-d] thiazole has been developed. The condensation products of this compound represent dipolar D- π -A or quadrupolar D- π -A- π -D structures with intramolecular charge transferability.

DNA-, RNA- and self-pairing properties of a pyrrolidinyl peptide nucleic acid with a (2'*R*,4'S)-prolyl-(1*S*,2*S*)-2-aminocyclopentanecarboxylic acid backbone

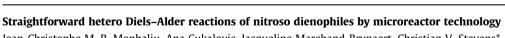
Jaru Taechalertpaisarn, Pitchanun Sriwarom, Chalotorn Boonlua, Nattawut Yotapan, Chotima Vilaivan, Tirayut Vilaivan*

A pyrrolidinyl peptide nucleic acid (PNA) with an alternating nucleobase-modified proline at the 4' position with (2'R.4'S) configuration and a (1S,2S)-2-aminocyclopentane-carboxylic acid backbone was synthesized and its DNA-, RNA- and self-pairing properties evaluated.

CI Ru=

Removal of an olefin metathesis catalyst using 4-nitrophenyl acrylate based polymer supports

Irena Pulko, Martina Sandholzer, Mitja Kolar, Christian Slugovc*, Peter Krajnc*



Jean-Christophe M. R. Monbaliu, Ana Cukalovic, Jacqueline Marchand-Brynaert, Christian V. Stevens*

 $\begin{bmatrix} 0 \\ N-R^{1} \end{bmatrix} \xrightarrow[0.2-1 \text{ mL.min}^{-1} \\ \underbrace{ 0.2-1 \text{ mL.min}^{-1} \\ 0.95^{\circ}\text{C} \\ 1-100 \text{ bar} \\ R^{2} \xrightarrow[---++]{n} \\ R^{2} \xrightarrow[---++]{n} \\ 65-96\% \end{bmatrix}$



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pp 5822-5826

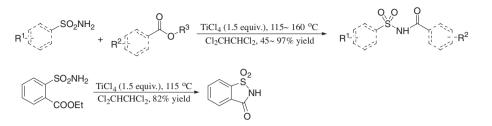


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TiCl₄-promoted direct N-acylation of sulfonamide with carboxylic ester

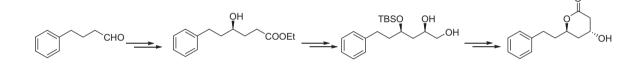
Shaomin Fu, Xiaoyan Lian, Tongmei Ma, Wenhua Chen, Meifang Zheng, Wei Zeng*



Several Lewis acids were investigated as promoters in the intermolecular or intramolecular direct N-acylation reaction of sulfonamides using carboxylic ester as an acylating agent. TiCl₄ was found to possess the highest activity and enhanced efficiently sulfonamide to form *N*-acylsulfonamides under optimized conditions. This method provides a novel approach to make *N*-acylsulfonamides from ester via an easy work-up procedure.

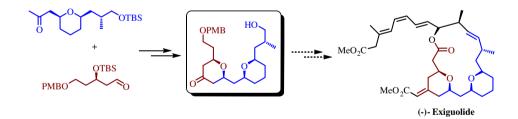
An organocatalytic route to the synthesis of lactone moiety of compactin and mevinolin

Pradeep Kumar*, Menaka Pandey, Priti Gupta, Dilip D. Dhavale



Synthesis of the methylene bis-tetrahydropyran motif of (-)-exiguolide

Ch. Raji Reddy*, N. Narsimha Rao



Novel two-step, one-pot synthesis of primary acylureas

Zili Xiao, Michael G. Yang*, Andrew J. Tebben, Michael A. Galella, David S. Weinstein

A new procedure for the synthesis of primary acylureas from cyanamide and a variety of carboxylic acids is described. Under mild reaction conditions, the products were obtained in good yield from commercially available starting materials.

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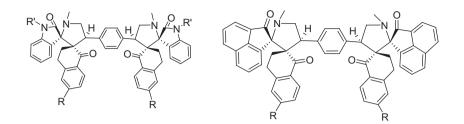


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Regio- and stereoselective synthesis of novel tetraspiro-bispyrrolidine and bisoxindolopyrrolidine derivatives through 1,3-dipolar cycloaddition reaction

R. Rajesh, R. Raghunathan*



OTHER CONTENT

Corrigendum

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*Corresponding author

()+ Supplementary data available via ScienceDirect

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