

Vol. 69 • Issue 46 • 18 November 2013 • ISSN 0040-4020

Tetrahedron

THE INTERNATIONAL JOURNAL FOR THE RAPID PUBLICATION OF FULL
ORIGINAL RESEARCH PAPERS AND CRITICAL REVIEWS IN ORGANIC CHEMISTRY

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Tetrahedron Vol. 69, Issue 46, 2013

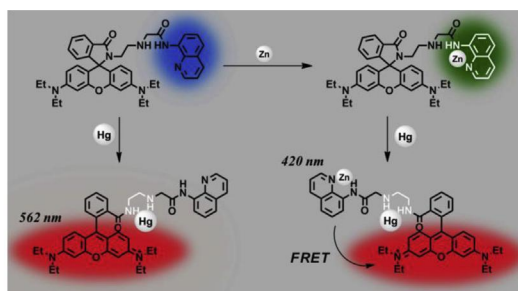
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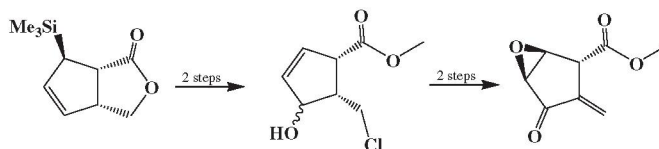
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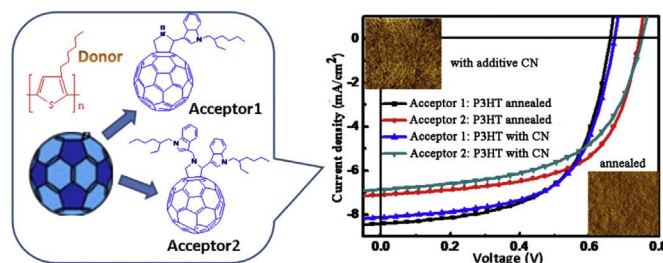
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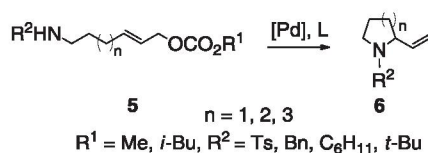
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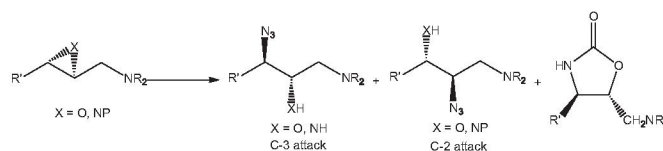
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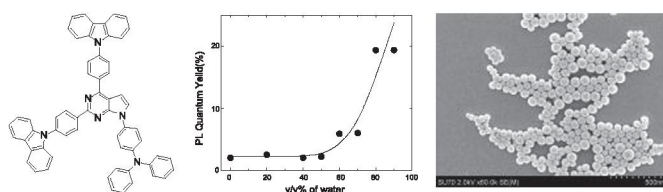
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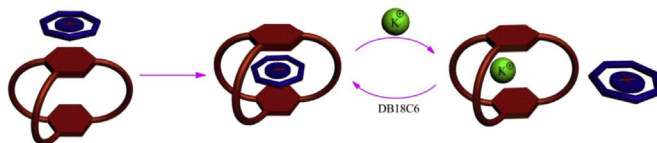
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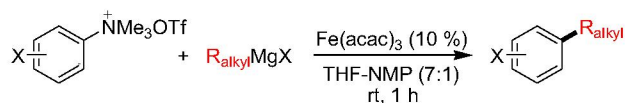
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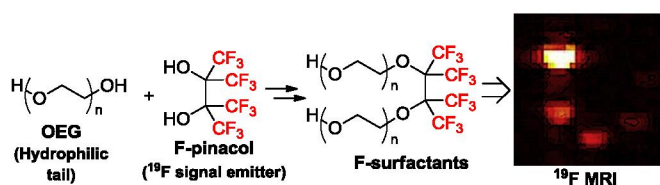
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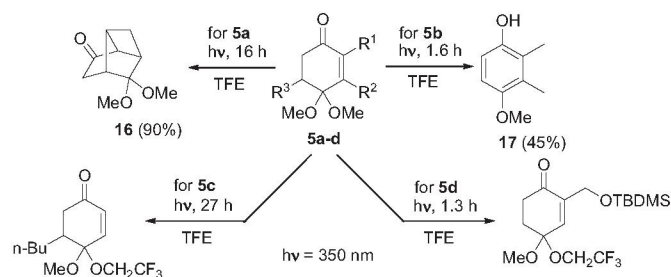
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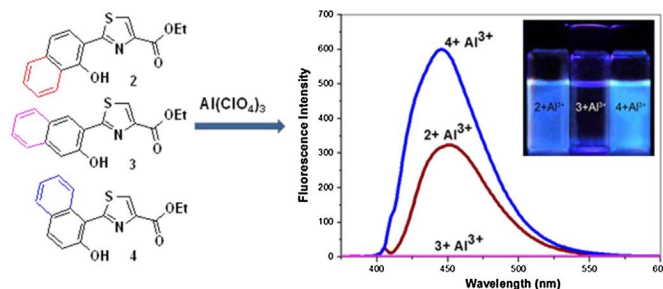
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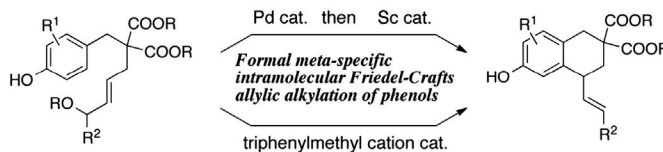
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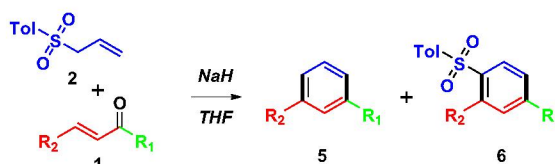
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Mariko Yoshida, Tomoyuki Nozaki, Tetsuhiro Nemoto, Yasumasa Hamada*

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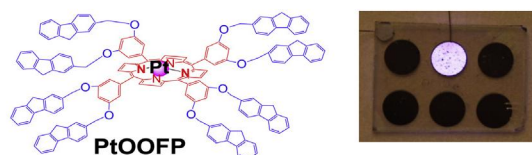
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Meng-Yang Chang*, Chieh-Kai Chan, Shin-Ying Lin, Ming-Hao Wu

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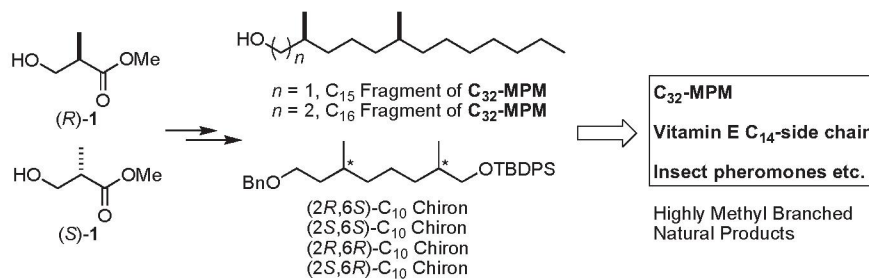


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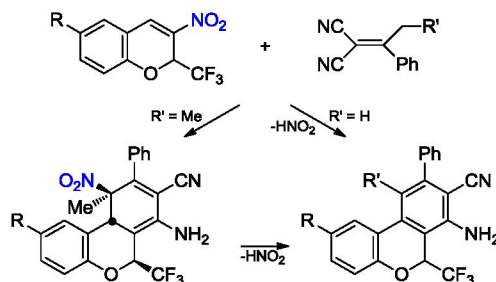
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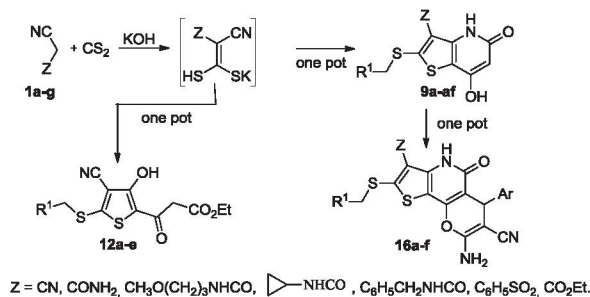
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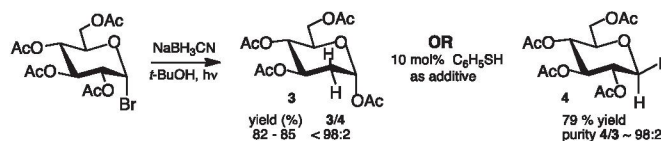
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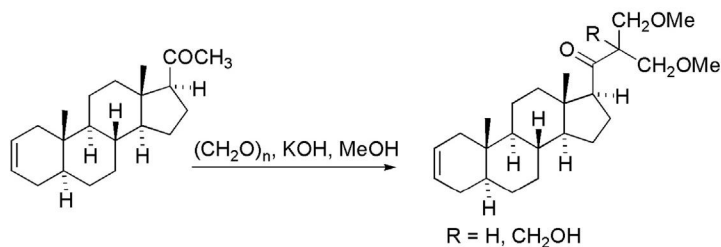
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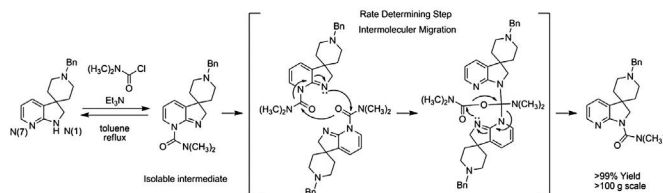
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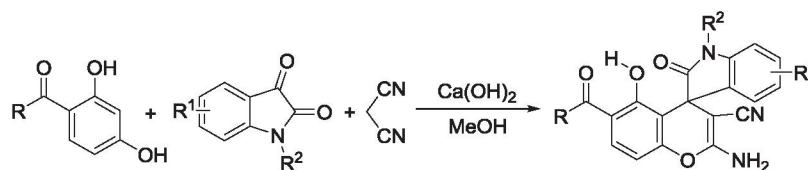
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**A novel synthesis of diverse 2-amino-5-hydroxy-4*H*-chromene derivatives with a spirooxindole nucleus by Ca(OH)₂-mediated three-component reactions of substituted resorcinols with isatins and malononitrile**

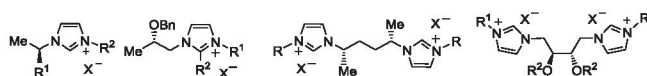
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Ji Hyang Park, Yong Rok Lee*, Sung Hong Kim

**Imidazolium-based chiral ionic liquids: synthesis and application**

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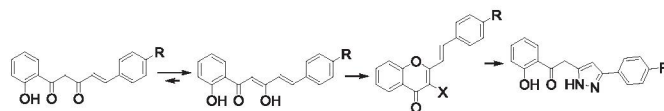
Yumiko Suzuki*, Junichiro Wakatsuki, Mariko Tsubaki, Masayuki Sato



(E)-3-Halo-2-styryl-4H-chromen-4-ones: synthesis and transformation to novel pyrazoles

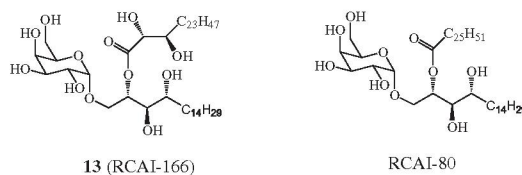
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Joana P.A. Ferreira, Vera L.M. Silva*, José Elguero, Artur M.S. Silva*

**Synthesis and biological activity of hydroxylated analogs of RCAI-80**

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Masao Shiozaki*, Takuya Tashiro, Hiroyuki Koshino, Tomokuni Shigeura, Hiroshi Watarai, Masaru Taniguchi, Kenji Mori

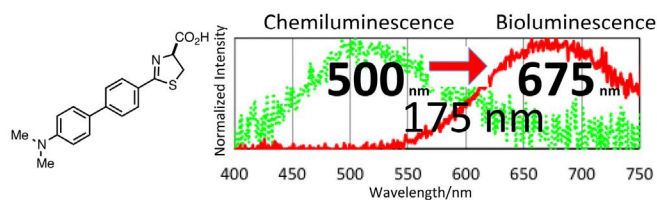


This (2*R*,3*R*)-dihydroxylated analogue (**13**) of RCAI-80 showed moderate activity for EAE suppression.

Synthesis and luminescence properties of biphenyl-type firefly luciferin analogs with a new, near-infrared light-emitting bioluminophore

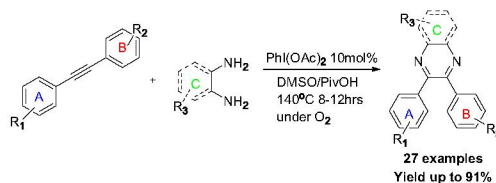
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Chihiro Miura, Masahiro Kiyama, Satoshi Iwano, Kazuto Ito, Rika Obata, Takashi Hirano, Shojiro Maki*, Haruki Niwa*

**Efficient synthesis of quinoxalines with hypervalent iodine as a catalyst**

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Chung-Yu Chen, Wan-Ping Hu, Mei-Chun Liu, Pi-Cheng Yan, Jeh-Jeng Wang*, Mei-Ing Chung*

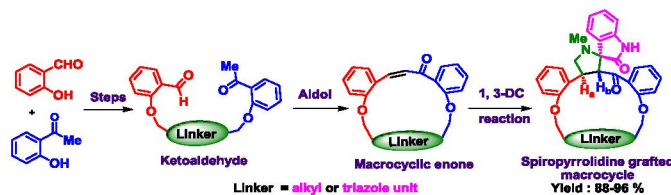


The one-pot metal free synthesis of quinoxaline derivatives with hypervalent iodine as a catalyst.

Regioselective synthesis of spiro pyrrolidine/spiropyrrrolizidine/spirothiazolidine-grafted macrocycles through 1,3-dipolar cycloaddition methodology

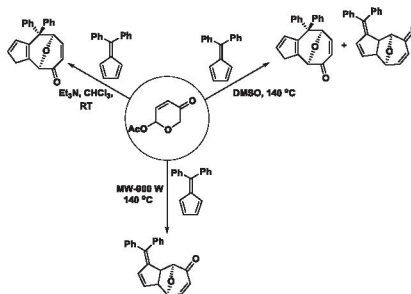
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S. Purushothaman, R. Prasanna, R. Raghunathan*


Cycloaddition profile of pentafulvenes with 3-oxidopyrylium betaine: experimental and theoretical investigations

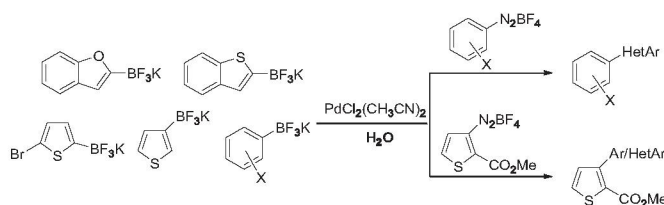
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Jinesh M. Kuthanapillil, A. Nijamudheen, Nayana Joseph, Praveen Prakash, E. Suresh, Ayan Datta*, K.V. Radhakrishnan*


A simple catalytic system based on PdCl₂(CH₃CN)₂ in water for cross-coupling reactions using diazonium salts

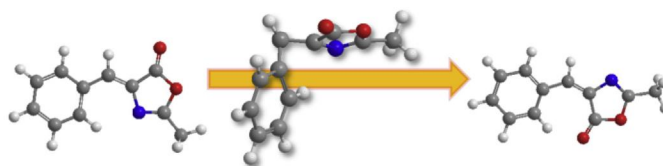
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Ouissam El Bakouri, Martí Fernández, Sandra Brun, Anna Pla-Quintana*, Anna Roglans*


Benzylidene–oxazolones as photoswitches: photochemistry and theoretical calculations

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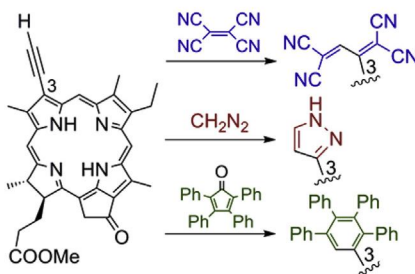
Ignacio Funes-Ardoiz, Marina Blanco-Lomas, Pedro J. Campos, Diego Sampedro*



Cycloaddition to a C3-ethynylated chlorophyll derivative and self-aggregation of zinc chlorin–pyrazole/triazole conjugates

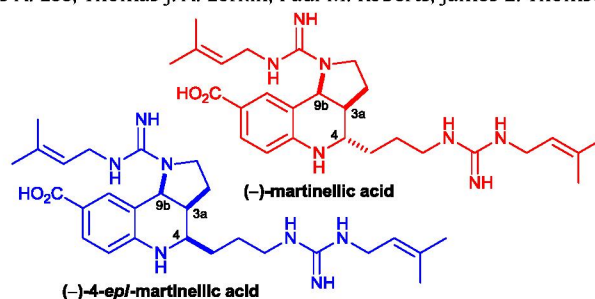
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Shin-ichi Sasaki, Keisuke Mizutani, Michio Kunieda, Hitoshi Tamiaki*

**A diastereodivergent strategy for the asymmetric syntheses of (–)-martinellic acid and (–)-4-*epi*-martinellic acid**

pp 9779–9803

Stephen G. Davies*, Ai M. Fletcher, James A. Lee, Thomas J. A. Lorkin, Paul M. Roberts, James E. Thomson

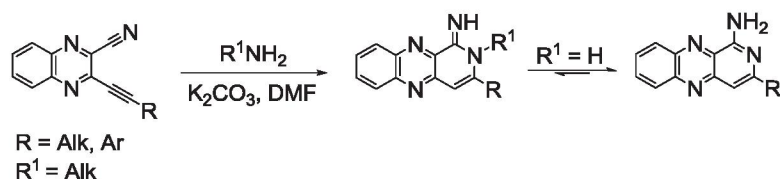


The asymmetric syntheses of (–)-martinellic acid and (–)-4-*epi*-martinellic acid were achieved in 20 steps from commercially available starting materials using a diastereodivergent strategy.

**Nucleophilic cyclization of 3-alkynylquinoxaline-2-carbonitriles into pyrido[3,4-*b*]quinoxalines**

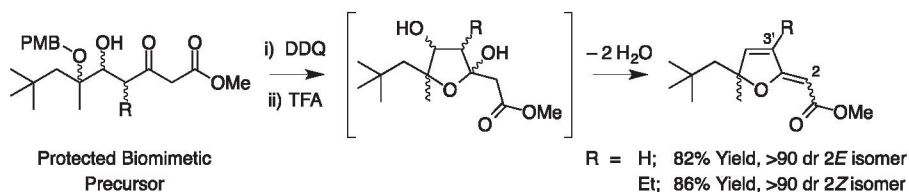
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Alexander S. Tyaglivy, Anna V. Gulevskaya*, Alexander F. Pozharskii, Olga I. Askalepova

**A biomimetic cascade for the formation of the methyl [2(5*H*)-furanlydene]ethanoate core of spongosoritin A and the gracilioethers**

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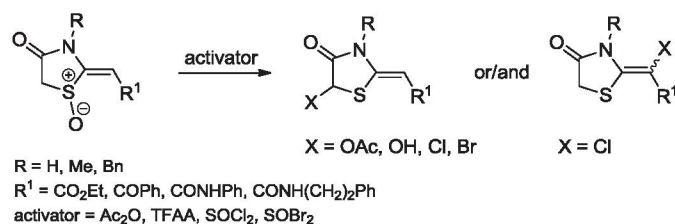
Matthew D. Norris, Michael V. Perkins*



Unusual mode of reactivity of 2-alkylidene-4-oxothiazolidine S-oxides under the Pummerer reaction conditions

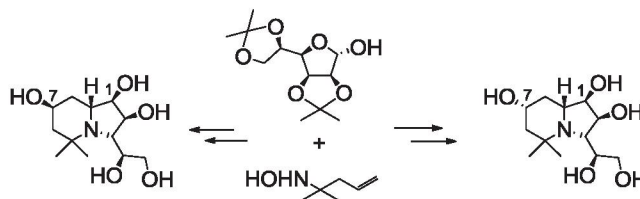
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Zdravko Džambaski, Đorđe Toljić, Bojan Bondžić, Rade Marković, Marija Baranac-Stojanović*

**Convenient synthesis of epimeric indolizidines by the intramolecular 1,3-dipolar cycloaddition of a sugar derived N-(3-alkenyl)nitron**

pp 9826–9831

Ewa Mironiuk-Puchalska*, Tomasz Rowicki, Wojciech Sas, Mariola Koszytkowska-Stawińska

**Isocyanide-based multicomponent reactions: synthesis of 2-(1-(alkylcarbamoyl)-2,2-dicyanoethyl)-N-alkylbenzamide and 1,7-diazaspiro[4,4]nonane-2,4-dione derivatives**

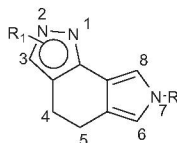
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Ebrahim Soleimani*, Mohsen Zainali, Neda Ghasemi, Behrouz Notash

**Convenient synthesis of pyrrolo[3,4-g]indazole**

pp 9839–9847

Virginia Spanò, Alessandra Montalbano, Anna Carbone, Barbara Parrino, Patrizia Diana, Girolamo Cirrincione, Paola Barraja*



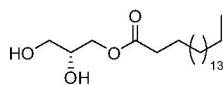
The synthesis of a novel class of tetrahydropyrrolo[3,4-g]indazoles is reported, by annelation of the pyrrole ring on the isoindole moiety by means of 5-hydroxymethylene tetrahydroisoindole-4-ones key intermediates, with good regioselectivity. Dihydroderivatives were also obtained by oxidation with DDQ of the corresponding tetrahydropyrrolo[3,4-g]indazoles. The growth inhibitory effect was evaluated at the National Cancer Institute of Bethesda and some derivatives showed modest activity.



On the optical rotation of 1 (or 3)-stearoyl-*sn*-glycerol

Hui-Jun Chen, Chao-Yuan Chen, Po Gao, Yikang Wu*

pp 9848–9851

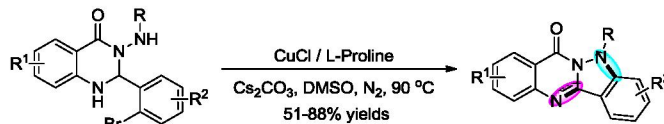
**What a confusion!**

$[\alpha]_D = +1.15$ ($c = 10.0$, CHCl_3)
 $[\alpha]_D = -0.43$ ($c = 1.25$, CHCl_3)
 $[\alpha]_D = -35.0$ ($c = 0.25$, CHCl_3)
 $[\alpha]_D = -36.3$ ($c = 0.055$, CHCl_3)

**Copper-catalyzed intramolecular C–N bond formation reaction of 3-amino-2-(2-bromophenyl)dihydroquinazolines: synthesis of indazolo[3,2-*b*]quinazolines**

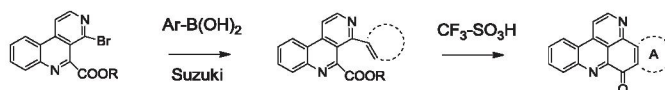
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Weiguang Yang, Leping Ye, Dayun Huang, Miaochang Liu, Jinchang Ding*, Jiuxi Chen*, Huayue Wu

**A novel approach to ring A analogues of the marine pyridoacridine alkaloid ascididemin**

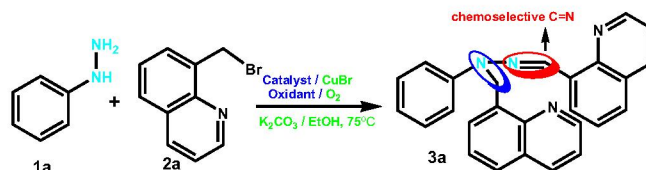
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Alois Plodek, Stephan Raeder, Franz Bracher*

**A one-pot synthesis of bisarylhydrazones by Cu(I)-catalyzed aerobic oxidation**

pp 9865–9869

Jiu-Rong Hu*, Wan-Jia Zhang, Da-Gui Zheng



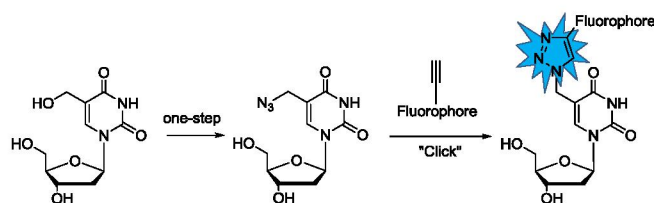
An efficient one-pot sequential synthesis of *N*-substituted (or *NH*) bisarylhydrazones based on Cu(I)-catalyzed aerobic oxidative coupling reaction has been developed. A further cyclization reaction could occur towards the synthesis of benzimidazoles or triazoles with elevated temperature. A plausible alkylation–oxidation–alkylation mechanism is proposed based on the control experiments.



One-step to get 5-azidomethyl-2'-deoxyuridine from 5-hydroxymethyl-2'-deoxyuridine and detection of it through click reaction

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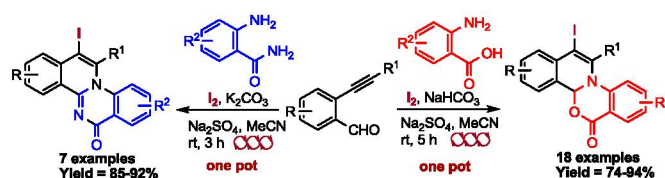
Xiaowei Xu, Shengyong Yan, Jianlin Hu, Pu Guo, Lai Wei, Xiaocheng Weng, Xiang Zhou*



Iodine-mediated electrophilic tandem cyclization of 2-alkynylbenzaldehydes with anthranilic acid leading to 1,2-dihydroisoquinoline-fused benzoxazinones

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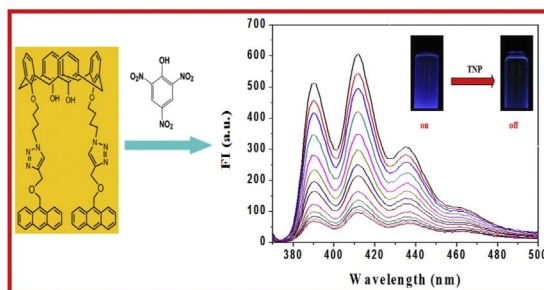
Shashikant U. Dighe, Sanjay Batra*



Synthesis of a novel fluorescent anthryl calix[4]arene as picric acid sensor

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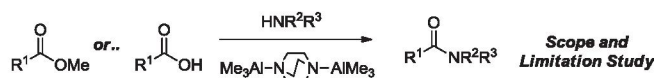
Fan Zhang, Li Luo, Yue Sun, Fajun Miao, Jiahai Bi, Shiliang Tan, Demei Tian*, Haibing Li*



On DABAL-Me₃ promoted formation of amides

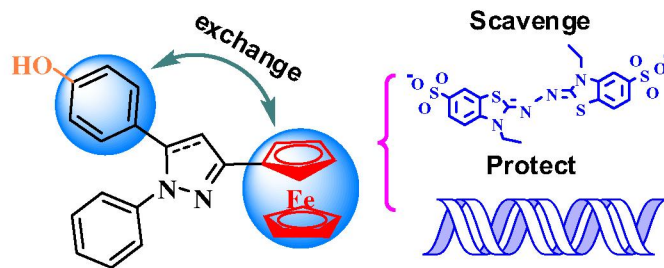
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Nathalie Dubois, Daniel Glynn, Thomas McInally, Barrie Rhodes, Simon Woodward*, Derek J. Irvine, Chris Dodds




Ferrocenyl-contained dendritic-like antioxidants with dihydropyrazole and pyrazole as the core: investigations into the role of ferrocenyl group and structure–activity relationship on scavenging radical and protecting DNA pp 9898–9905

Pei-Ze Li, Zai-Qun Liu*



Capacities of ferrocenyl dendritic antioxidants to scavenge ABTS^{•+} radical and to protect DNA against AAPH-induced oxidation.

*Corresponding author

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