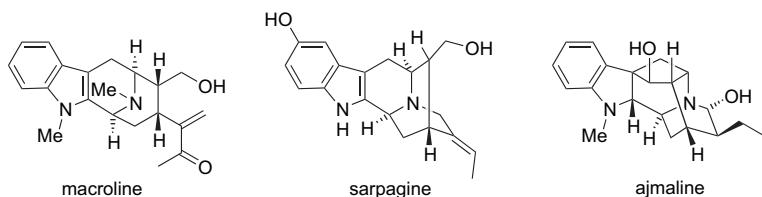


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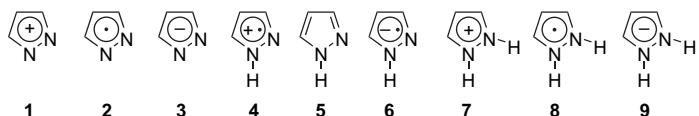


## ARTICLES

**The calculated enthalpies of the nine pyrazole anions, cations, and radicals: a comparison with experiment**

Ibon Alkorta\* and José Elguero

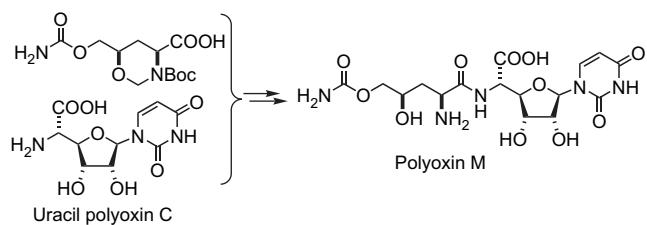
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## First synthesis of polyoxin M

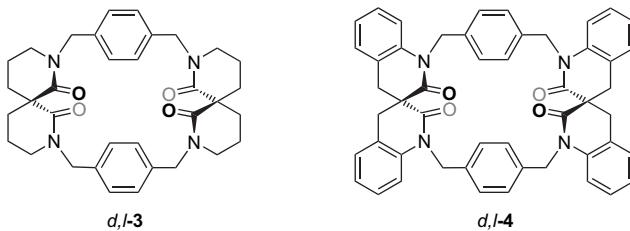
Yuuichi Shiro, Keisuke Kato, Mikio Fujii, Yoshiteru Ida and Hiroyuki Akita\*

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**A  $D_2$  symmetric tetraamide macrocycle based on 1,1',4,4'-tetrahydro[3,3'(2H,2'H)-spirobiquinoline]-2,2'-dione: synthesis and selectivity for lithium over sodium and alkaline earth ions** pp 8696–8701

Heung-Jin Choi,\* Yeon Sil Park, Moon Goo Kim, Yang Jin Park, Nam Sik Yoon and Thomas W. Bell

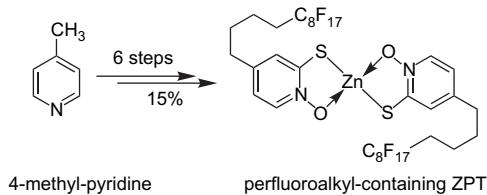


Macrocyclic ionophores *d,l*-3 and *d,l*-4 with four amide carbonyl ligands were synthesized and investigated in lithium ion-selective electrodes.

**Design and synthesis of novel perfluoroalkyl-containing zinc pyrithione biocide**

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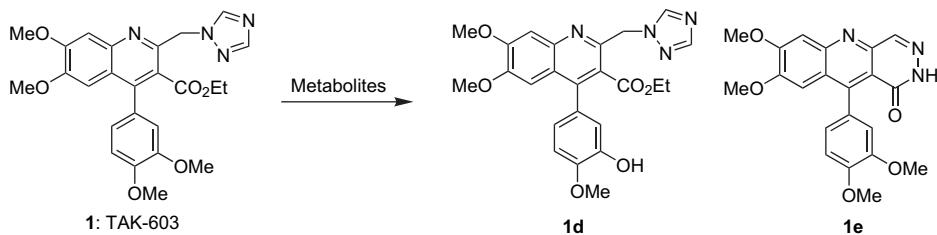
Jian-Ying Sun, Xiao-Long Qiu, Wei-Dong Meng and Feng-Ling Qing\*



**Syntheses of metabolites of ethyl 4-(3,4-dimethoxyphenyl)-6,7-dimethoxy-2-(1,2,4-triazol-1-ylmethyl)quinoline-3-carboxylate (TAK-603)**

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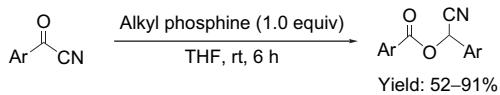
Masahiro Mizuno,\* Makoto Yamashita, Yasuhiro Sawai, Koji Nakamoto and Mitsutaka Goto



**Alkyl phosphines promoted reductive coupling of acyl cyanides: formation of *O*-acyl cyanohydrins**

pp 8715–8719

Wen Zhang and Min Shi\*

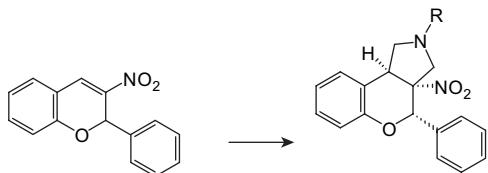


In the presence of alkyl phosphines, reductive coupling of acyl cyanides proceeded smoothly at room temperature in THF to give the corresponding *O*-acyl cyanohydrins in moderate to high yields. The possible mechanism was discussed on the basis of deuterium labeling and control experiments, indicating that one hydride transfer took place from alkyl phosphine to cyanohydrin.



**3-Nitrochromene derivatives as  $2\pi$  components in 1,3-dipolar cycloadditions of azomethine ylides**  
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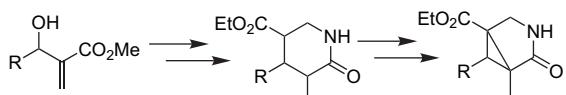
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**Studies toward the construction of substituted piperidine-2-ones and pyridine-2-ones from Baylis–Hillman adducts: discovery of a facile synthesis of 5-methyl-4-oxo-6-aryl-3-aza-bicyclo[3.1.0]hexane-1-carboxylates**

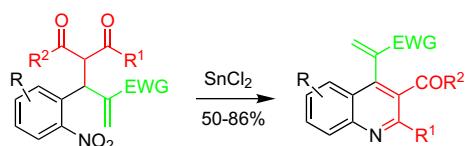
V. Singh, G. P. Yadav, P. R. Maulik and S. Batra\*

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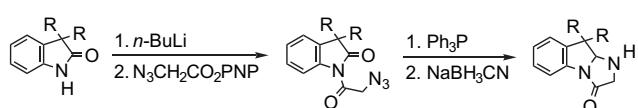
**An alternate approach to quinoline architecture via Baylis–Hillman chemistry:  $\text{SnCl}_2$ -mediated tandem reaction toward synthesis of 4-(substituted vinyl)-quinolines**  
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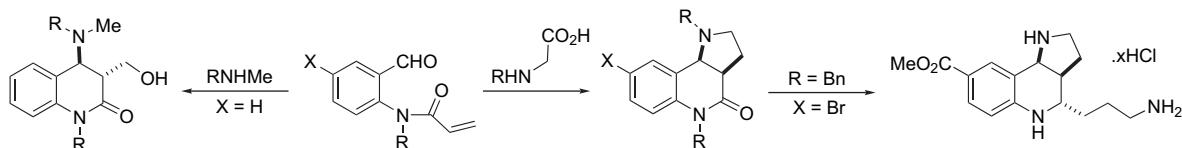
Two methods for annulation of glycine to the 1 and 2 positions of oxindoles are described. One method introduces *p*-nitrophenyl  $\alpha$ -azidoacetate as an N-acylation reagent. The other method involves a selective  $\text{BF}_3\text{-Et}_3\text{SiH}$  imide reduction.



**Cyclization reactions of *N*-acryloyl-2-aminobenzaldehyde derivatives: formal total synthesis of martinellic acid**

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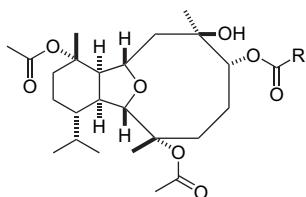
Yong He, Hossen Mahmud, Remond Moningka, Carl J. Lovely\* and H. V. Rasika Dias\*



**A homologous series of eunicellin-based diterpenes from *Acalycigorgia* sp. characterised by tandem mass spectrometry**

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Kwaku Kyeremeh, Thomas C. Baddeley, Bridget K. Stein and Marcel Jaspars\*

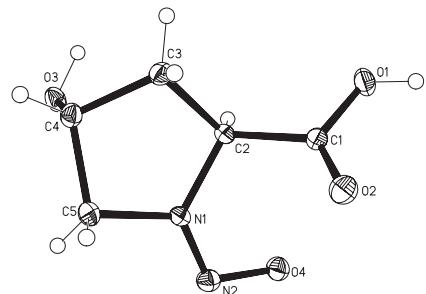


The discovery and structure determination of a homologous series of eunicellin-based diterpenes from the gorgonian *Acalycigorgia* sp. by NMR, MS and MS/MS methods are described.

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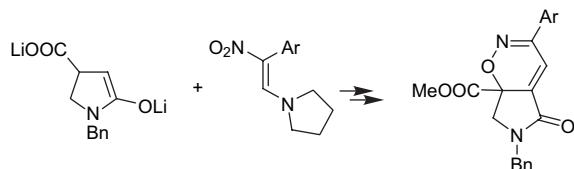
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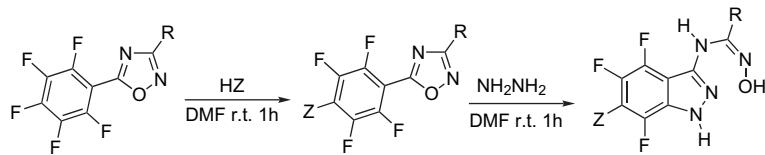
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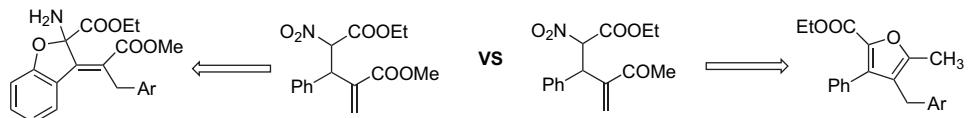
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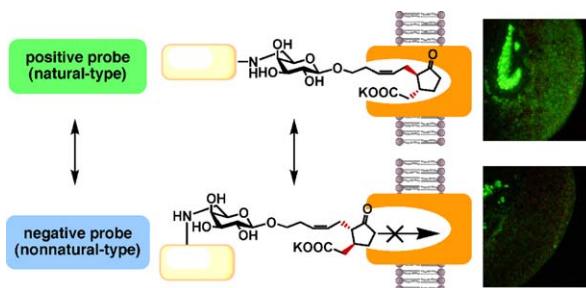
Ka Young Lee, Saravanan Gowrisankar, Young Ju Lee and Jae Nyong Kim\*



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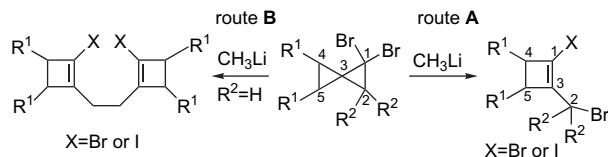
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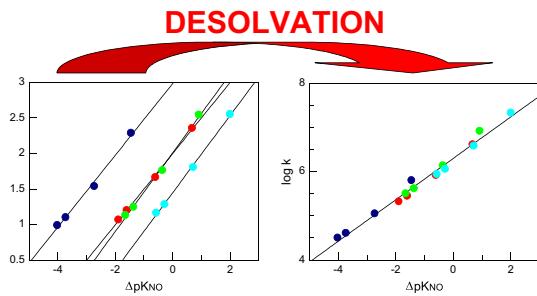
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Elena B. Averina, Rashad R. Karimov, Kseniya N. Sedenkova, Yurii K. Grishin, Tamara S. Kuznetzova\* and Nikolai S. Zefirov



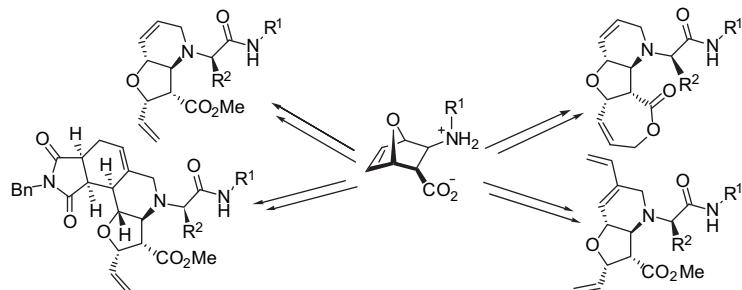
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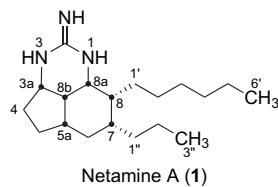
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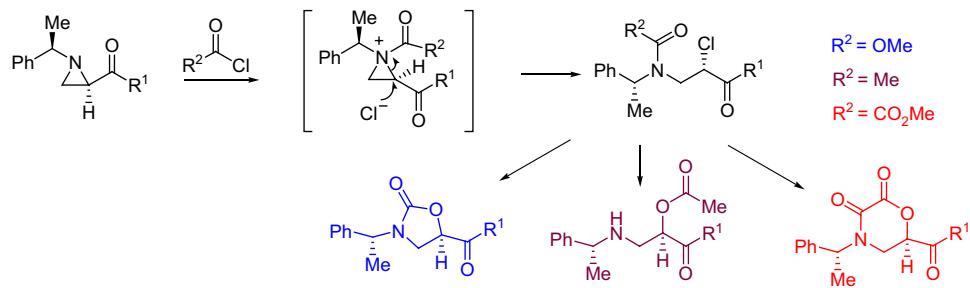
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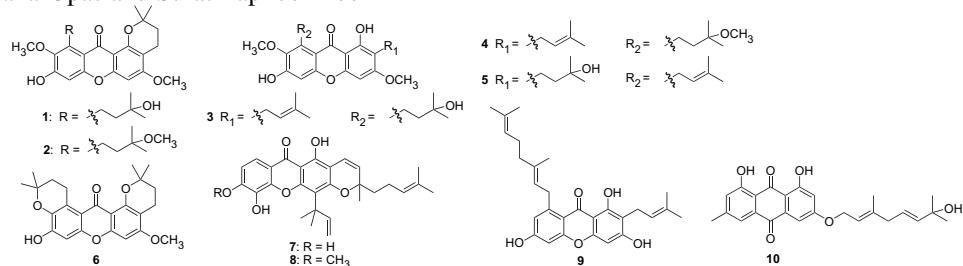
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Yongeon Kim, Hyun-Joon Ha,\* Hoseop Yun, Baeck Kyoung Lee and Won Koo Lee\*



**Bioactive prenylated xanthones and anthraquinones from *Cratoxylum formosum* ssp. *pruniflorum*** pp 8850–8859

Nawong Boonnak, Chatchanok Karalai,\* Suchada Chantrapromma, Chanita Ponglimanont, Hoong-Kun Fun,  
Akkharawit Kanjana-Opas and Surat Laphookhieo



Nine new prenylated xanthones (**1–9**) and a new anthraquinone (**10**) were isolated from the roots and barks of *Cratoxylum formosum* ssp. *pruniflorum*. In addition, antibacterial and cytotoxic activities of the isolates were also evaluated.

\*Corresponding author

**i** Supplementary data available via ScienceDirect



Full text of this journal is available, on-line from **ScienceDirect**. Visit [www.sciencedirect.com](http://www.sciencedirect.com) for more information.

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