

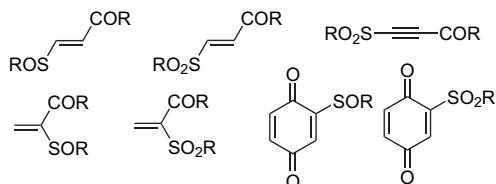
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The chemistry of sulfinyl and sulfonyl enones

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Valentine G. Nenajdenko,* Arkady L. Krasovskiy and Elizabeth S. Balenkova

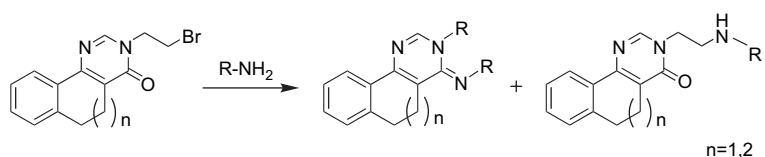


ARTICLES

Polycyclic N-heterocyclic compounds. Part 58: Rearrangement reactions of fused 3-(2-bromoethyl)pyrimidin-4(3H)-ones with primary amines and antidepressive evaluation of the products

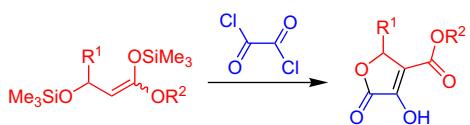
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Hiromi Ohtomo, Tsuyoshi Tagata, Kenji Sasaki, Takashi Hirota* and Kensuke Okuda*

**Synthesis of 4-alkoxycarbonyl-butenolides by uncatalyzed one-pot cyclization of 1,3-bis(silyloxy)alk-1-enes with oxalyl chloride**

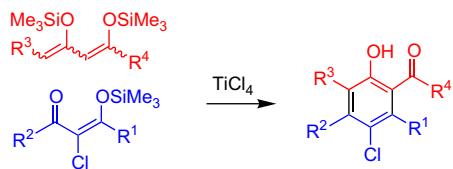
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Rüdiger Dede, Lars Michaelis, Dilver Fuentes, Mirza Arfan Yawer, Ibrar Hussain, Christine Fischer and Peter Langer*



Regioselective synthesis of 4-chlorophenols, 10-chloro-7-hydroxy-6H-benzo[c]chromen-6-ones, and 4-chloro-1-hydroxy-9H-fluoren-9-ones based on [3+3] cyclizations of 1,3-bis(silyloxy)-1,3-dienes with 2-chloro-3-silyloxy-2-en-1-ones

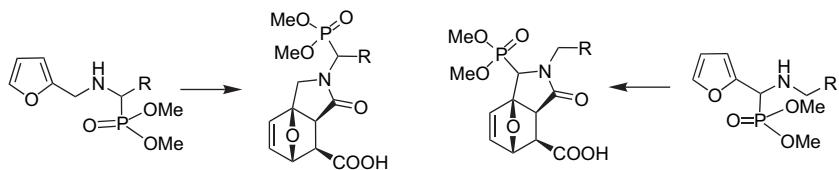
Mirza Arfan Yawer, Ibrar Hussain, Stefanie Reim, Zafar Ahmed, Ehsan Ullah, Inam Iqbal, Christine Fischer, Helmut Reinke, Helmar Görls and Peter Langer*



α -Acylaminophosphonates possessing epoxyisoindolone moiety

Georgiy O. Kachkovskyi and Oleg I. Kolodiazhnyi*

pp 12576–12582

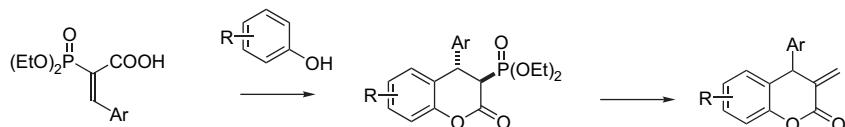


α -Acylaminophosphonates **6** possessing an epoxyisoindolone moiety were prepared with a good stereoselectivity ($de \geq 80\%$) by a tandem acylation/[4+2]-cycloaddition reaction between maleic anhydride and α -aminophosphonates derived from a furfurylamine.

Trifluoromethanesulfonic acid mediated Friedel–Crafts reaction of (*E*)-3-aryl-2-(diethoxyphosphoryl)acrylic acids with electron-rich hydroxyarenes. A convenient approach to α -methylene- δ -valerolactones

Henryk Krawczyk,* Łukasz Albrecht, Jakub Wojciechowski and Wojciech M. Wolf

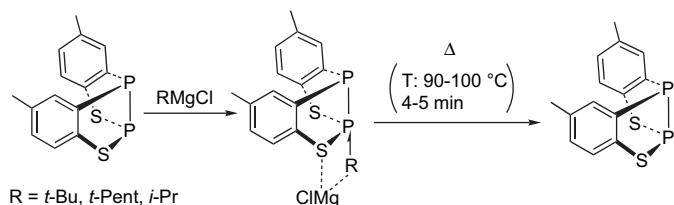
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Identification of a four-center intermediate in a Grignard addition reaction to a P–S bond

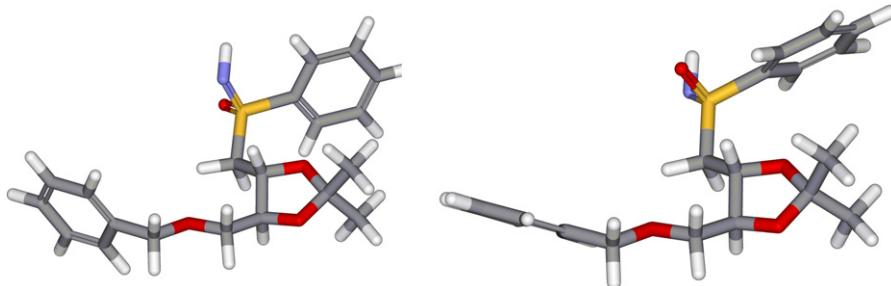
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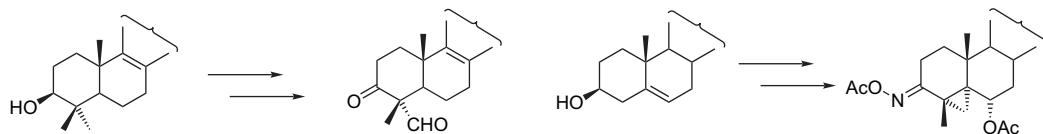
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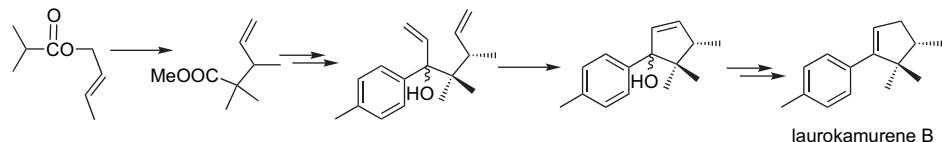
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The first total synthesis of (\pm)-laurokamurene B

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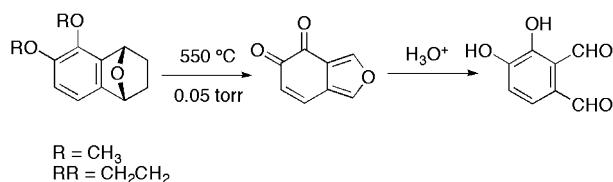
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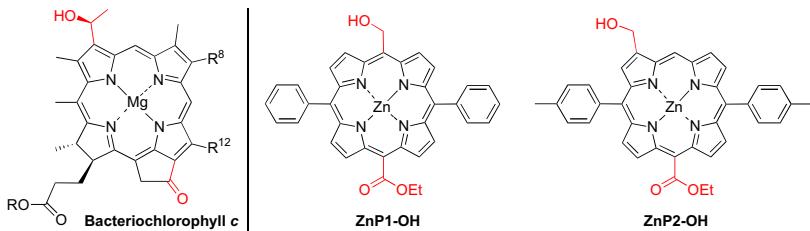
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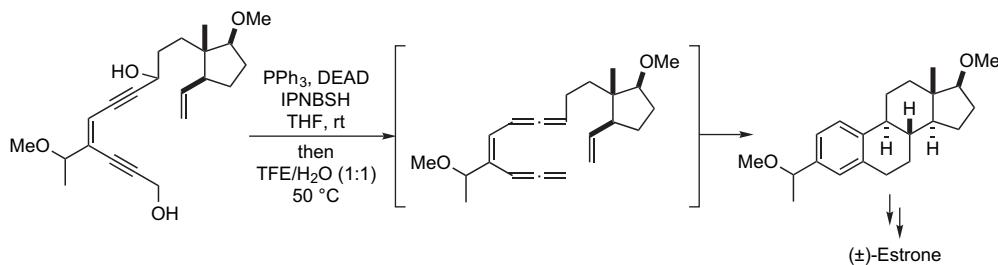
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A porphyrin (**ZnP2-OH**) has been prepared and found in the crystalline state to exhibit infinite coordination polymers that are distinct from those of the analogue **ZnP1-OH**.

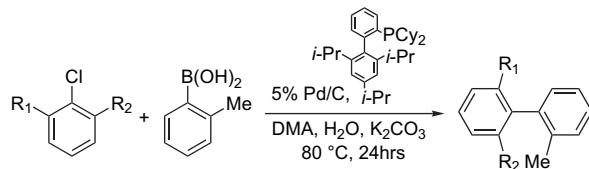
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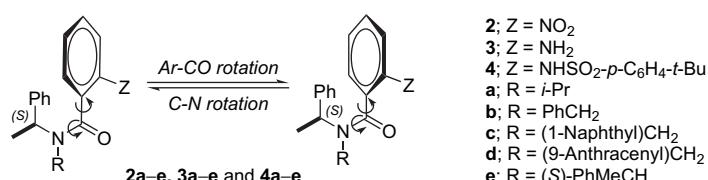
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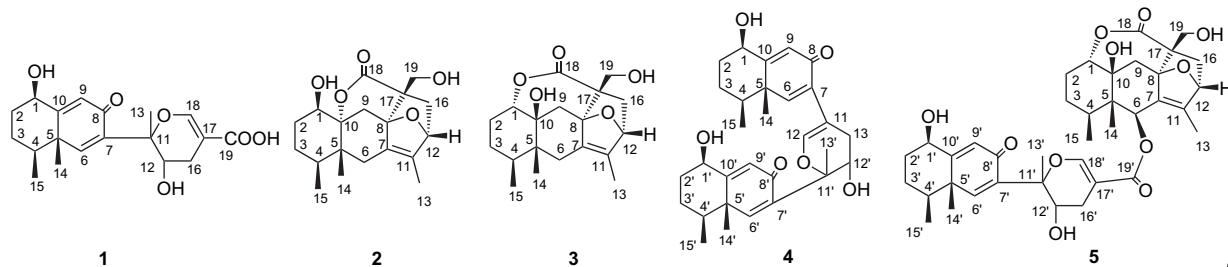
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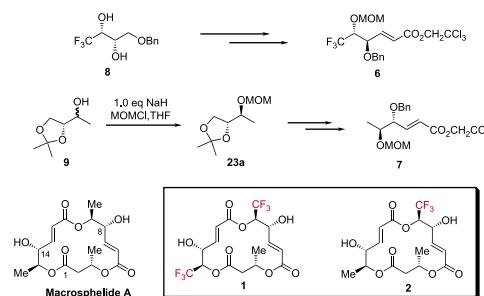
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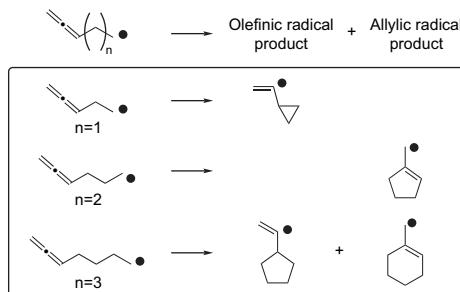


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Regiochemistry in radical cyclization of allenes

Jing Shi, Miao Zhang, Yao Fu,* Lei Liu and Qing-Xiang Guo*

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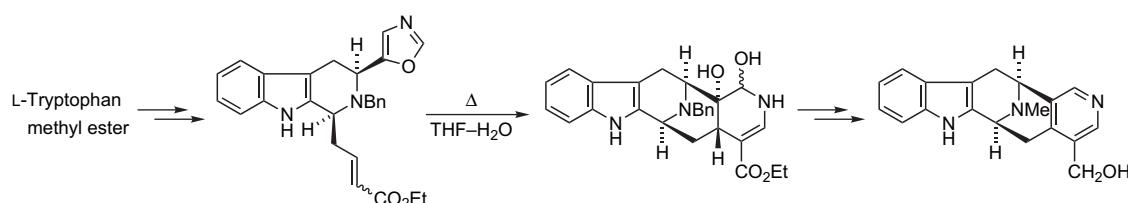


i+

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Masashi Ohba* and Itaru Natsutani

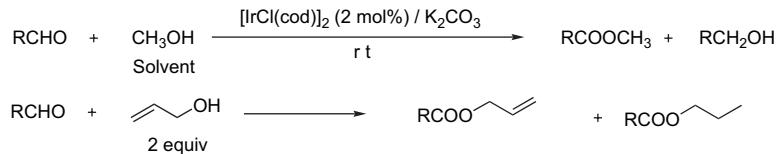
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[IrCl(cod)]₂-catalyzed direct oxidative esterification of aldehydes with alcohols

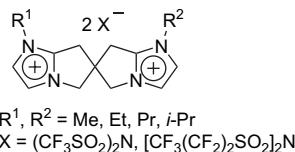
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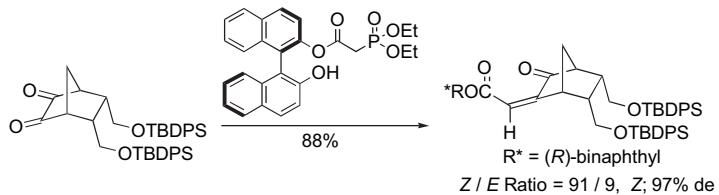
Mahesh L. Patil,* C. V. Laxman Rao, Shinobu Takizawa, Kazuhiro Takenaka, Kiyotaka Onitsuka and Hiroaki Sasai*

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**Discrimination of carbonyl groups of meso- α -diketones with Horner–Wadsworth–Emmons reagent of chiral binaphthyl esters**

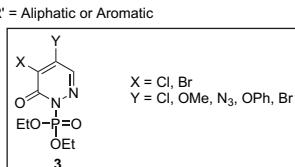
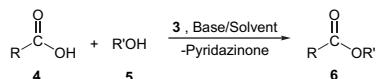
Daiki Monguchi, Yoshihisa Ohta, Tatsuya Yoshiuchi, Toshiyuki Watanabe, Takumi Furuta,* Kiyoshi Tanaka and Kaoru Fuji

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**Effective esterification of carboxylic acids using (6-oxo-6H-pyridazin-1-yl)phosphoric acid diethyl ester as novel coupling agents**

Ju-Eun Won, Ho-Kyun Kim, Jeum-Jong Kim, Heong-Seup Yim, Min-Jung Kim, Seung-Beom Kang, Hyun-A Chung, Sang-Gyeong Lee* and Yong-Jin Yoon*

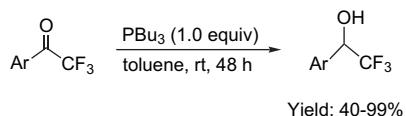
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Reduction of 2,2,2-trifluoro-1-arylethanones with alkyl phosphines

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Min Shi,* Xu-Guang Liu, Ying-Wen Guo and Wen Zhang

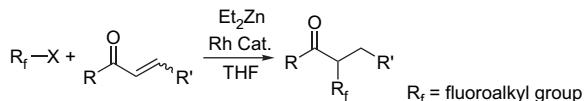


In the presence of alkyl phosphines, reduction of 2,2,2-trifluoro-1-arylethanones proceeded smoothly to give the corresponding reduction products in moderate to high yields at room temperature. The possible mechanism was discussed on the basis of deuterium labeling and control experiments, indicating that one hydrogen transfer took place from alkyl phosphine to the carbonyl group activated by a strongly electron-withdrawing trifluoromethyl group.

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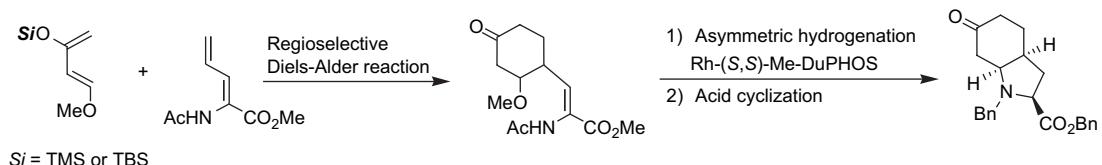
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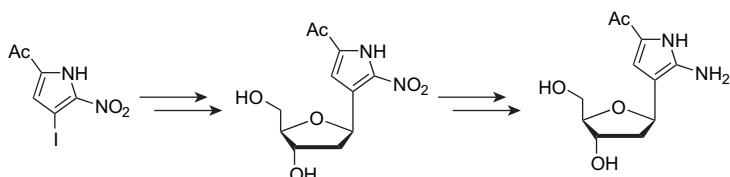
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Yoichiro Hoshina, Takayuki Doi and Takashi Takahashi*

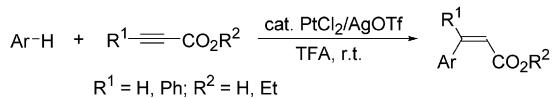
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 Jozo Oyamada and Tsugio Kitamura*

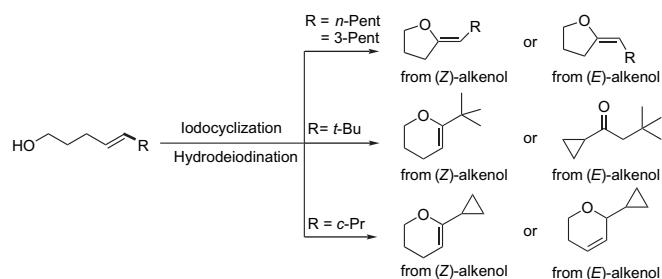


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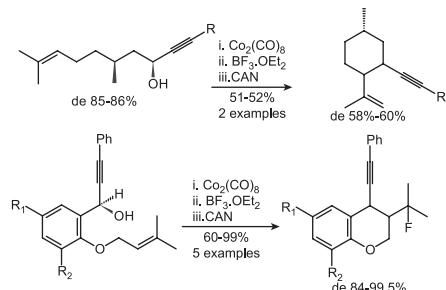
Claudio Paolucci* and Paolo Righi



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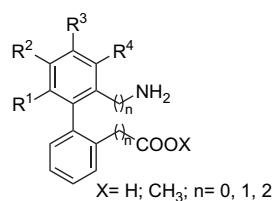
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Elisa Perissutti, Francesco Frecentese, Antonio Lavecchia, Ferdinando Fiorino, Beatrice Severino, Francesca De Angelis, Vincenzo Santagada* and Giuseppe Caliendo



Synthesis of an indole containing KDR kinase inhibitor by tandem Sonogashira coupling-5-endo-dig-cyclization as a key step

Sanjay S. Palimkar, Vijaykumar S. More, P. Harish Kumar and Kumar V. Srinivasan*

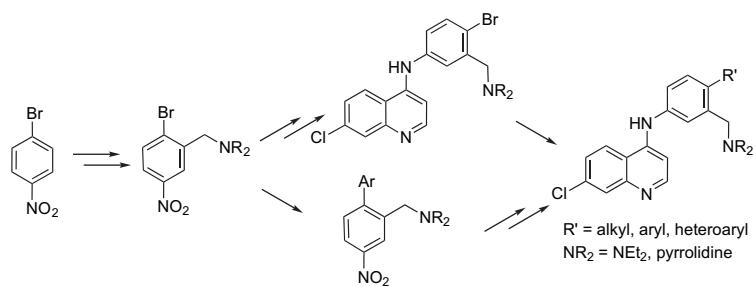
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Suzuki–Miyaura cross-coupling reaction as the key step for the synthesis of some new 4'-aryl and alkyl substituted analogues of amodiaquine and amopyroquine

Emilia Paunescu, Nicolas Matuszak and Patricia Melnyk*

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*Corresponding author

ⓘ⁺ Supplementary data available via ScienceDirect



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