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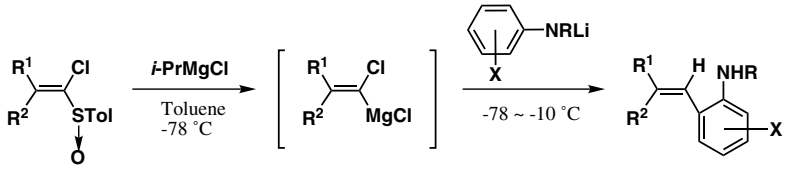
Tetrahedron Letters

Tetrahedron Letters Vol. 45, No. 30, 2004

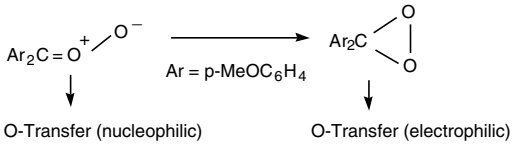
Contents

COMMUNICATIONS

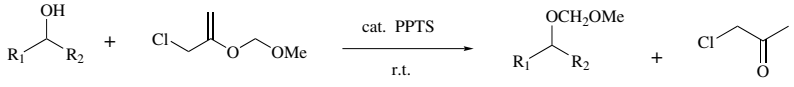
Direct alkenylation of arylamines at the *ortho*-position **pp 5785–5789**
Tsuyoshi Satoh,* Yumi Ogino and Masatomo Nakamura



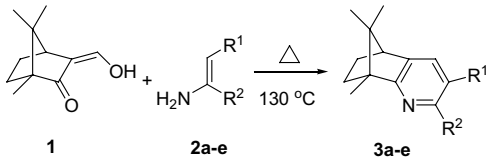
Direct observation for the cyclization of a diarylcarbonyl oxide **pp 5791–5793**
Hiroshi Hanaki, Yuta Fukatsu, Masaki Harada and Yasuhiko Sawaki*



New efficient method of alkoxymethyl etherification of secondary alcohols **pp 5795–5797**
Yosuke Watanabe and Tetsuya Ikemoto*

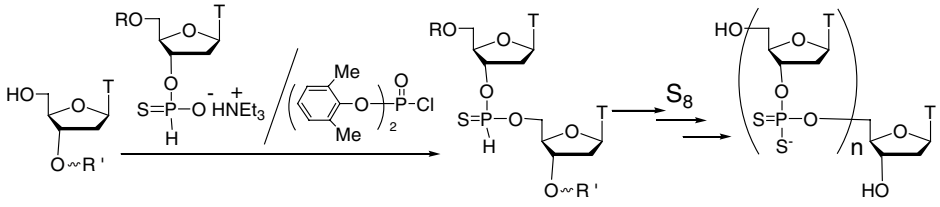


Synthesis of various camphor-based chiral pyridine derivatives **pp 5799–5801**
Cihangir Tanyeli,* İdris M. Akhmedov and Murat Işık



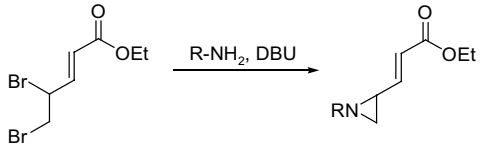
(+)-β-Hydroxymethylenecamphor **1** and enamines **2a**–**e** were transformed into chiral camphor-based pyridine derivatives **3a**–**e** via a tandem condensation reaction in good yields.

Synthesis of oligonucleoside phosphorodithioates by the *H*-phosphonothioate method **pp 5803–5806**
Kazuo Kamaike,* Kumi Hirose, Yoshihiro Kayama and Etsuko Kawashima

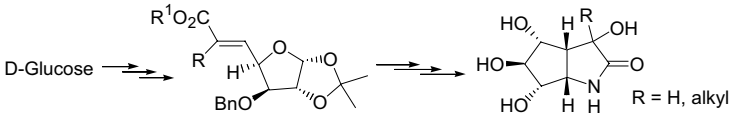


The phosphorodithioate oligomers were efficiently synthesized using bis(2,6-dimethylphenyl) phosphorochloridate as a coupling agent by application of the *H*-phosphonothioate method.

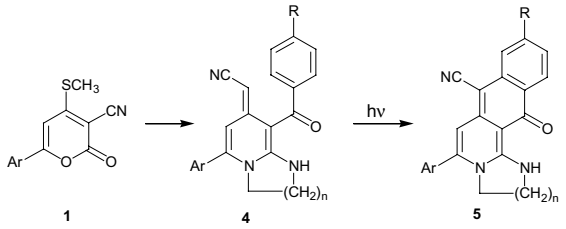
Aziridination of γ,δ-dibromoethyl-2-pentenoate with primary amines: extension of the Gabriel–Cromwell reaction **pp 5807–5810**
Rachel L. Weller and Scott R. Rajski*



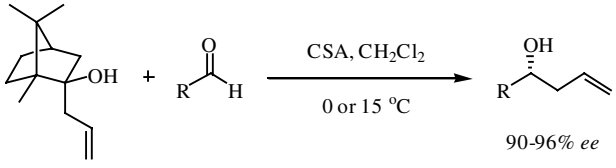
[3+2] Cycloaddition reactions: a simple entry to the 1-aza-2-oxo-3,4,5,6-tetrahydroxybicyclo[3.3.0]octane ring system **pp 5811–5814**
Ashim Roy, Biswajit G. Roy, Basudeb Achari and Sukhendu B. Mandal*



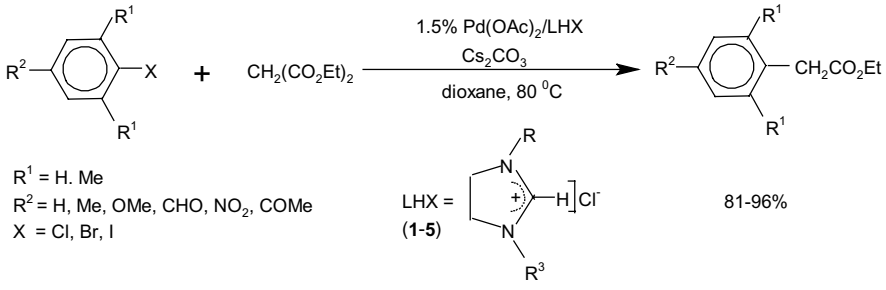
An innovative approach to the synthesis of annelated [*a*]diaza-anthracenones through tandem cyclization **pp 5815–5818**
Ashoke Sharon, Prakas R. Maulik, Raja Roy and Vishnu Ji Ram*



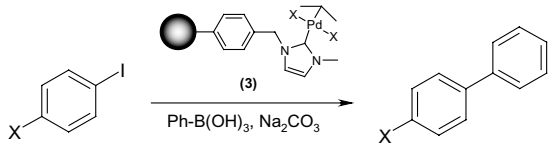
A highly enantioselective allyl-transfer through suppression of epimerization **pp 5819–5822**
Cheng-Hsia Angeline Lee and Teck-Peng Loh*



Synthesis of arylacetic acid derivatives from diethyl malonate using in situ formed palladium(1,3-dialkylimidazolidin-2-ylidene) catalysts **pp 5823–5825**
Ismail Özdemir, Murat Yiğit, Engin Çetinkaya and Bekir Çetinkaya*

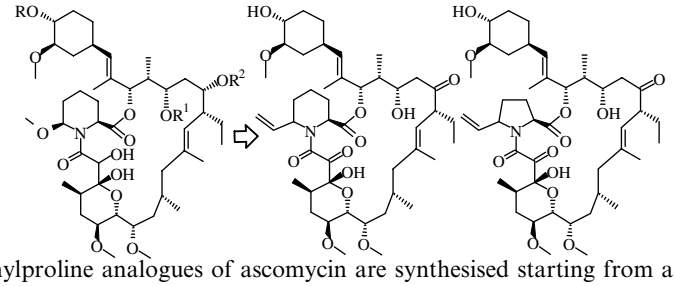


***N*-Heterocyclic carbene–palladium complex on polystyrene resin surface as polymer-supported catalyst and its application in Suzuki cross-coupling reaction** **pp 5827–5831**
Jong-Ho Kim, Bong-Hyun Jun, Jang-Woong Byun and Yoon-Sik Lee*



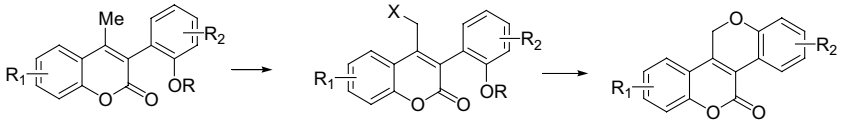
A poly(imidazoliummethyl styrene)-*surface grafted*-polystyrene resin was prepared by suspension polymerization. This was used as the polymer-supported carbene precursor for the palladium complex, which efficiently catalyzed the Suzuki cross-coupling of aryl halides and phenylboronic acid.

Synthesis of 6-vinyl and 5-vinylproline analogues of ascomycin **pp 5833–5836**
Murty A. R. C. Bulusu,* Peter Waldstätten, Thomas Tricotet, Christophe Rochais, Andrea Steck and Markus Bacher

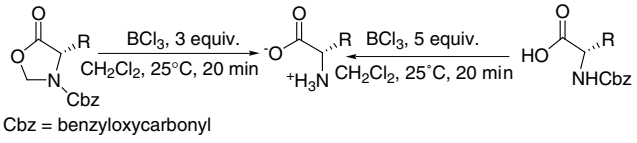


6-Vinyl, and, (5*R*)- and (5*S*)-vinylproline analogues of ascomycin are synthesised starting from a photoproduct obtained through irradiation of a suitably protected ascomycin derivative.

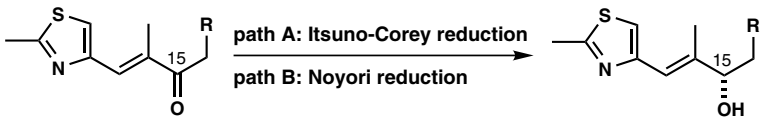
A facile synthesis of an unsymmetric benzopyranobenzopyran ring system **pp 5837–5839**
Ramesh M. Kanojia, Nareshkumar Jain, Jiayi Xu and Zhihua Sui*



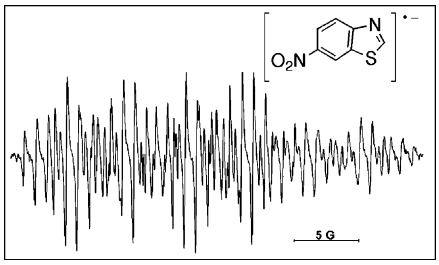
Facile and rapid regeneration of free amino acids from *N*-benzyloxycarbonyl-5-oxazolidinones and from *N*-benzyloxycarbonylamino derivatives by treatment with BCl_3 in dichloromethane **pp 5841–5843**
Pietro Allevi,* Riccardo Cribiù and Mario Anastasia



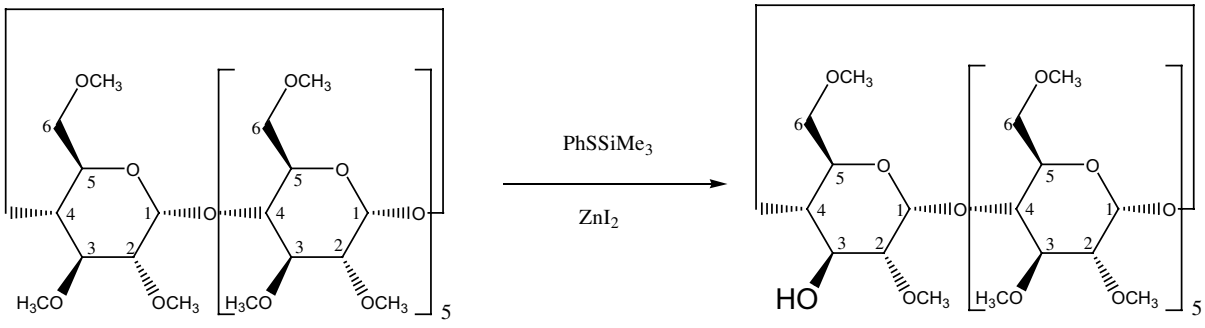
Practical syntheses of the C12–C21 epothilone subunit via catalytic asymmetric reductions: Itsuno–Corey oxazaborolidine reduction and asymmetric Noyori hydrogenation **pp 5845–5847**
Emily A. Reiff, Sajiv K. Nair, B. S. Narayan Reddy, Jun Inagaki, John T. Henri, Jack F. Greiner and Gunda I. Georg*



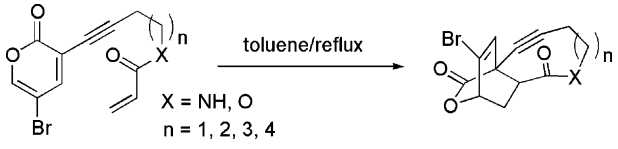
Radical anions of nitrobenzothiazoles: EPR study of conjugative properties of benzothiazolyl systems **pp 5849–5852**
Francesco Ciminale



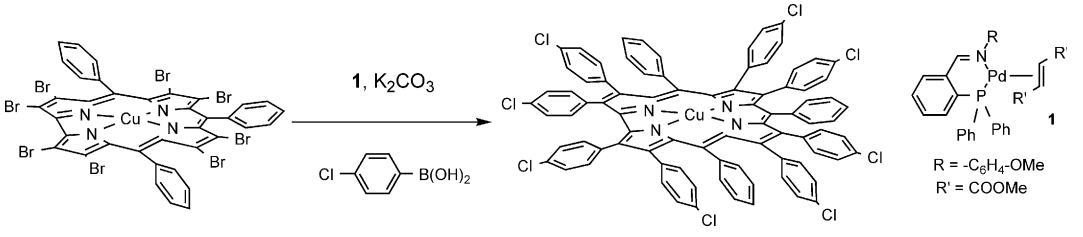
Direct and convenient access to mono 3-hydroxy per-O-methylated α-cyclodextrin **pp 5853–5856**
Thomas Chaise, Eric Bourgeaux, Pascal Cardinael* and J. C. Combret



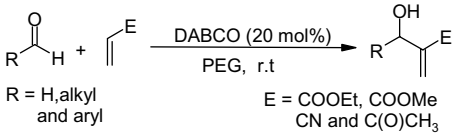
IMDA cycloadditions of 3-alkynyl tethered 2-pyrones for the synthesis of medium-sized macrocycles **pp 5857–5860**
Jeong-Taek Shin, Seunghoon Shin and Cheon-Gyu Cho*



Iminophosphine–palladium(0) complexes as highly active catalysts in the Suzuki reaction. Synthesis of undecaaryl substituted corroles **pp 5861–5864**
Alberto Scrivanti,* Valentina Beghetto, Ugo Matteoli, Simonetta Antonaroli, Alessia Marini, Federica Mandoj, Roberto Paolesse and Bruno Crociani*

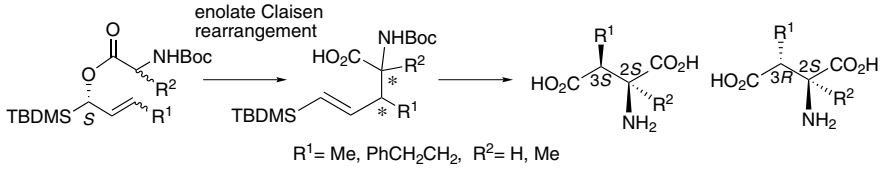


Poly(ethyleneglycol) (PEG): a rapid and recyclable reaction medium for the DABCO-catalyzed Baylis–Hillman reaction **pp 5865–5867**
S. Chandrasekhar,* Ch. Narsihmulu, B. Saritha and S. Shameem Sultana

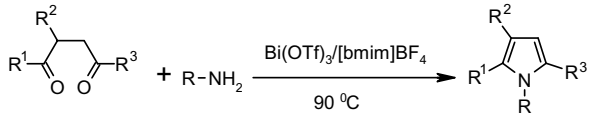


PEG (400) has been used as a rapid and recyclable reaction medium for the Baylis–Hillman reaction with the conventional basic catalyst DABCO (20 mol %) with very good yields of products. Recyclability is achieved with no further addition of DABCO to the reaction medium over four runs without substantial loss in yields. Incidentally, DABCO is recycled for the first time in this transformation.

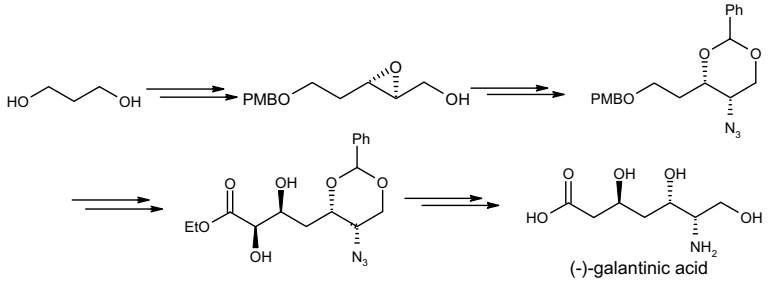
Synthesis of optically active β-alkyl aspartate via [3,3] sigmatropic rearrangement of α-acyloxytrialkylsilane **pp 5869–5872**
Kazuhiko Sakaguchi,* Masahiro Yamamoto, Tetsuo Kawamoto, Takeshi Yamada, Tetsuro Shinada, Keiko Shimamoto and Yasufumi Ohfune*



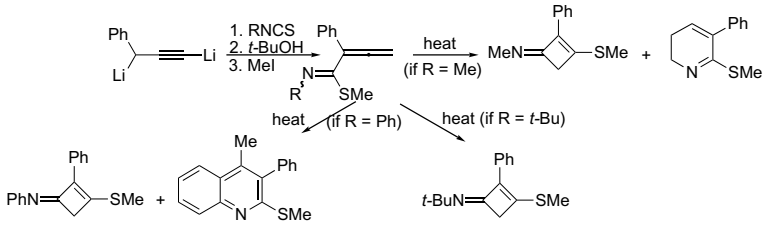
$Bi(OTf)_3$/[bmim]BF_4 as novel and reusable catalytic system for the synthesis of furan, pyrrole and thiophene derivatives **pp 5873–5876**
J. S. Yadav,* B. V. S. Reddy, B. Eeshwaraiah and Manoj Kumar Gupta



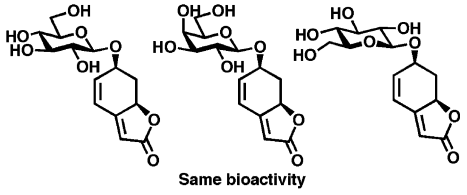
Enantioselective synthesis of (−)-galantinic acid **pp 5877–5879**
Satyendra Kumar Pandey, SubbaRao V. Kandula and Pradeep Kumar*



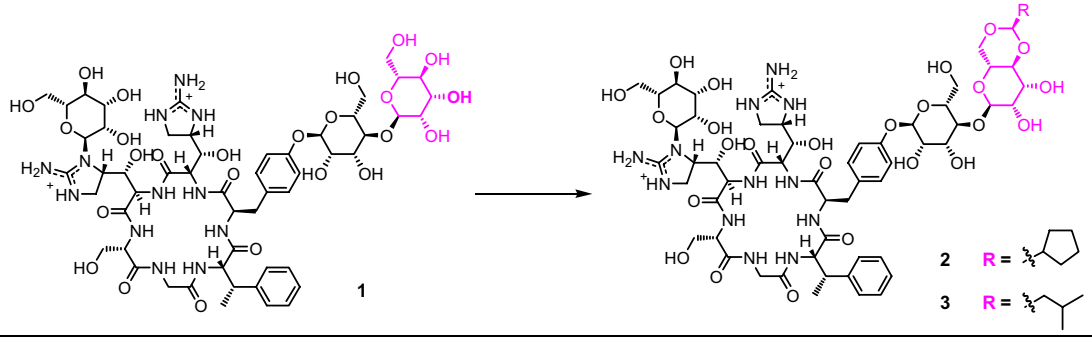
A novel thermal rearrangement of allenic imidothioates. Formation of iminocyclobutenes **pp 5881–5883**
Ol'ga A. Tarasova, Nina A. Nedolya, Lambert Brandsma* and Alexander I. Albanov



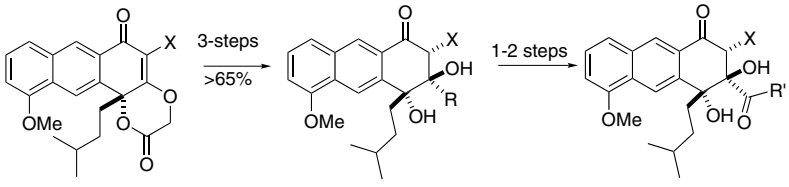
Enantioselective synthesis of phyllanthurinolactone, a leaf-closing substance of *Phyllanthus urinaria* L., and its analogs toward the development of molecular probes **pp 5885–5888**
Yoshifumi Urakawa, Takanori Sugimoto, Hirotaka Sato and Minoru Ueda*



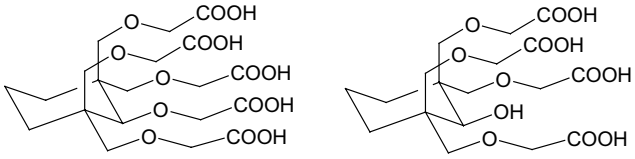
Structural determination of mannopeptimycin cyclic acetals **pp 5889–5893**
Haiyin He,* Ting-Zhong Wang, Russell G. Dushin, Xidong Feng, Bo Shen, Joseph S. Ashcroft, Frank E. Koehn and Guy T. Carter



A short diastereoselective synthesis of the (±)-rishirilide B core structure **pp 5895–5899**
Junhua Wang and Thomas R. R. Pettus*

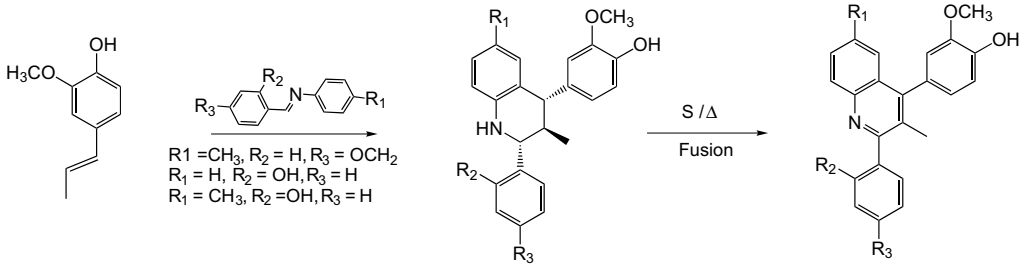


Synthesis of new polyoxapolycarboxylic ligands for lanthanide(III) ions complexation **pp 5901–5903**
Silvio Aime, Camilla Cavallotti, Giancarlo Cravotto, Giovanni B. Giovenzana* and Giovanni Palmisano*

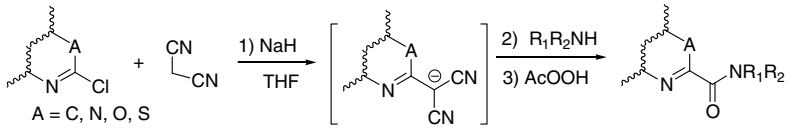


The synthesis of new polyoxapolycarboxylic ligands and the evaluation of their complexing ability towards lanthanide(III) ions is reported. The ligands, obtained in two-steps from easily available chemicals, show interesting complexing properties.

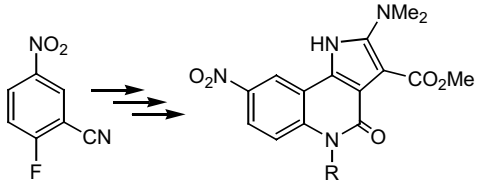
Synthèse de nouveaux dérivés tétrahydroquinoléines et quinoléines via la réaction d'aza-Diels–Alder suivie d'aromatisation **pp 5905–5908**
Fouzia Fadel, Soumia Lafquih Titouani,* Mohamed Soufiaoui, Hafida Ajamay and Ahmed Mazzah



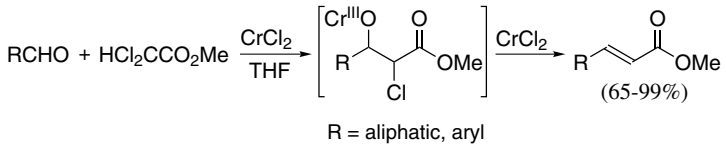
Malononitrile as a carbonyl synthon: a one-pot preparation of heteroaryl amide via a S_NAr-oxidation–displacement strategy **pp 5909–5911**
Juliang Zhu, Henry Wong, Zhongxing Zhang, Zhiwei Yin, John F. Kadow, Nicholas A. Meanwell and Tao Wang*



Concomitant ring contraction cyclization strategy for the synthesis of novel 4-oxo-4,5-dihydro-pyrroloquinolines **pp 5913–5916**
Gabriel Thia Manh, Hicham Bakkali, Lucie Maingot, Muriel Pipelier, Uday Joshi, Jean Paul Pradère, Stéphane Sabelle, Remy Tuloup and Didier Dubreuil*

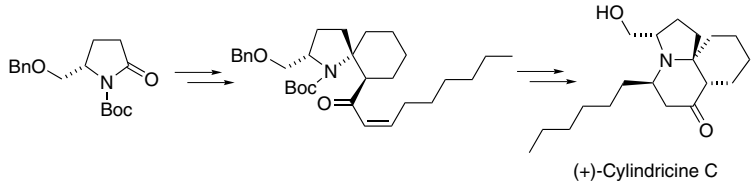


A highly stereospecific synthesis of (*E*)-α,β-unsaturated esters **pp 5917–5920**
Deb K. Barma, Asish Kundu, Anish Bandyopadhyay, Abhijit Kundu, Bhavani Sangras, Anne Briot, Charles Mioskowski* and J. R. Falck*

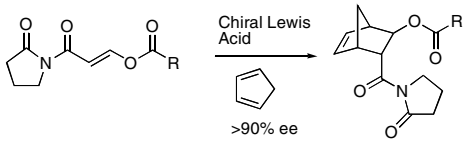


Olefination of aldehydes using methyl dichloroacetate and $CrCl_2$ generates (*E*)-α,β-unsaturated esters exclusively. Using limited $CrCl_2$, the intermediate α-chloro-β-hydroxy adducts can be isolated in good yields.

Total synthesis of (+)-cylindricine C **pp 5921–5924**
Toshiharu Arai, Hideki Abe, Sakae Aoyagi and Chihiro Kibayashi*

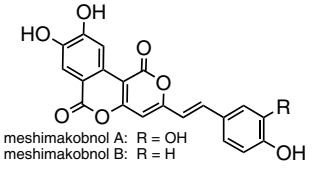


Enantioselective Diels–Alder reactions of 3-(acyloxy)acrylates **pp 5925–5929**
Mukund P. Sibi* and Hirofumi Matsunaga



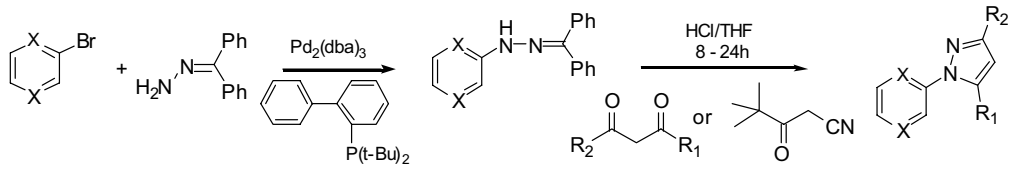
We have successfully carried out Diels-Alder reactions with 3-acyloxyacrylates using chiral Lewis acid catalysts. These reactions proceed with high enantioselectivity when a chiral Lewis acid derived from $Cu(OTf)_2$ and a bisoxazoline is used. The facility of the reaction is dependent on the nature of the acyloxy group in the dienophile.

Identification of novel substituted fused aromatic compounds, meshimakobnol A and B, from natural *Phellinus linteus* fruit body **pp 5931–5933**
Akito Nagatsu, Shizue Itoh, Rie Tanaka, Setsuko Kato, Mitsumasa Haruna, Keiichi Kishimoto, Hideki Hirayama, Yukihiro Goda, Hajime Mizukami and Yukio Ogihara*



Novel 1*H*,6*H*-pyranyl[4,3-*c*][2]benzopyrane-1,6-diones, meshimakobnol A and B, were isolated from natural *Phellinus linteus* fruit body. The structure elucidation of these fused aromatic compounds was achieved by a spectroscopic method including the measurement of FG-HMBC with various delay times.

Application of the palladium-catalyzed N-arylation of hydrazones to deactivated heteroaryl halides in the synthesis of pyrazoles **pp 5935–5937**
Nizar Haddad,* Annette Salvagno and Carl Busacca



OTHER CONTENTS

Contributors to this issue **p I**
Instructions to contributors **pp III–VI**

*Corresponding author
ⓘ+ Supplementary data available via ScienceDirect

COVER

A total synthesis of (+)-cylindricine C has been carried out. Key steps include a spirocyclization via enamine formation and an intramolecular Michael addition to form the tricyclic core. Details can be found in *Tetrahedron Letters*, **2004,** *45,* 5921–5924.

Indexed/Abstracted 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

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