

Tetrahedron Letters Vol. 48, No. 3, 2007

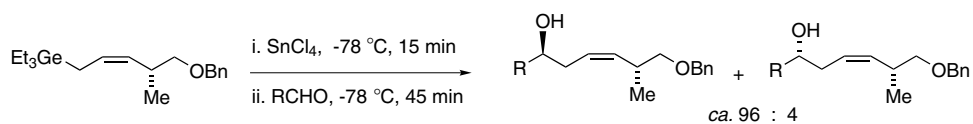
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COMMUNICATIONS

1,5-Remote stereocontrol using allylgermanes

pp 337–340

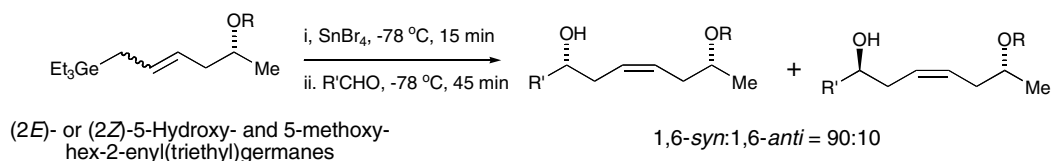
Pilar Castreno, Eric J. Thomas* and Anna P. Weston



1,6-Remote stereocontrol using allylgermanes

pp 341–343

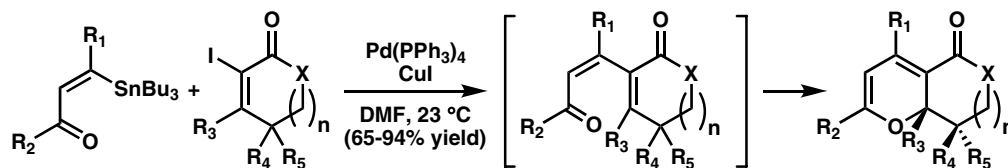
Eric J. Thomas* and Anna P. Weston



The development and scope of a versatile tandem Stille-oxa-electrocyclization reaction

pp 345–350

Uttam K. Tambar, Taichi Kano, John F. Zepernick and Brian M. Stoltz*

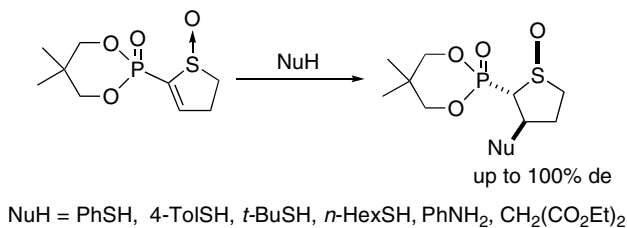


A palladium-catalyzed tandem Stille-oxa-electrocyclization reaction has been developed for the convergent preparation of highly substituted polycyclic pyran systems. The strategy presented in this letter is an alternative to the known methods for constructing similar pyran systems. The substrate scope of this diastereoselective transformation is explored.

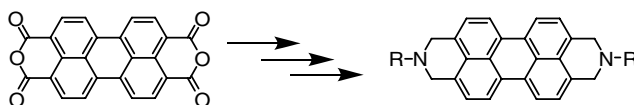


Diastereoselective Michael additions to α,β -unsaturated α -sulfinyl phosphonates in the thiolane series pp 351–355

Piotr Łyżwa, Aleksandra Jankowiak, Małgorzata Kwiatkowska, Marian Mikołajczyk, Piotr Kiełbasiński,* Alexander Betz, Paul-Alain Jaffres, Annie-Claude Gaumont and Mihaela Gulea*

**1,3,8,10-Tetrahydro-2,9-diazadibenzo[*cd,lm*]perylene: synthesis of reduced perylene bisimide analogues** pp 357–359

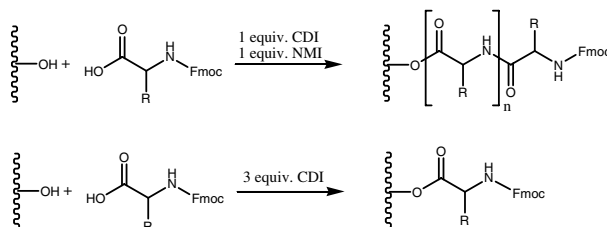
Masaki Takahashi,* Yousuke Suzuki, Yasunori Ichihashi, Mitsuji Yamashita and Hideki Kawai



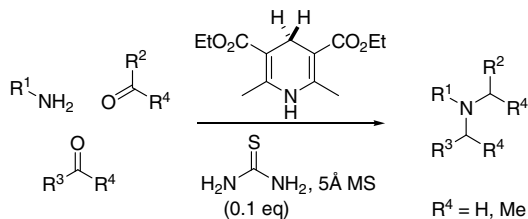
A new class of reduced perylene bisimide analogues was prepared from commercially available perylene-3,4,9,10-tetracarboxylic dianhydride.

**Synthesis of cleavable peptides with authentic C-termini: an application for fully automated SPOT synthesis** pp 361–364

Bernhard Ay, Rudolf Volkmer and Prisca Boisguerin*

Direct coupling of CDI activated amino acids on cellulose membranes allows the synthesis of cleavable peptides with free C-termini in yields of at least 50 $\mu\text{g}/\text{cm}^2$ which are enough for any biological assays.**Synthesis of hindered tertiary amines by a mild reductive amination procedure** pp 365–369

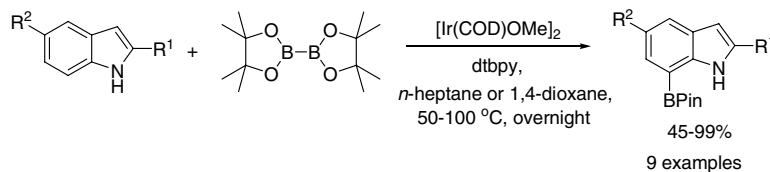
Dirk Menche,* Stefanie Böhm, Jun Li, Sven Rudolph and Wiebke Zander

An efficient metal-free procedure for the synthesis of tertiary amines by the reductive amination of carbonyl compounds is reported, which allows a convergent and mild access to amines of the general formulas NR(R')₂ and NRR'R''.

A highly selective Ir-catalyzed borylation of 2-substituted indoles: a new access to 2,7- and 2,4,7-substituted indoles

pp 371–375

Wei Fun Lo, Hanns Martin Kaiser, Anke Spannenberg, Matthias Beller and Man Kin Tse*

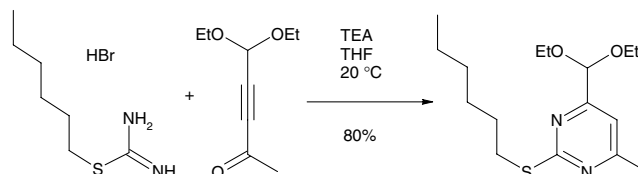


The iridium-catalyzed borylation of 2-substituted indoles has been investigated. Direct CH-activation is observed highly selective at the 7-position of the indole. The scope and limitation of the mono- and di-borylation reaction are presented.

Highly flexible and efficient synthesis of the GABA_B enhancer 4-(2-hexylsulfanyl-6-methyl-pyrimidin-4-ylmethyl)-morpholine

pp 377–380

Julien Verron, Paricher Malherbe, Eric Prinssen, Andrew W. Thomas, Nadine Nock and Raffaello Masciadri*

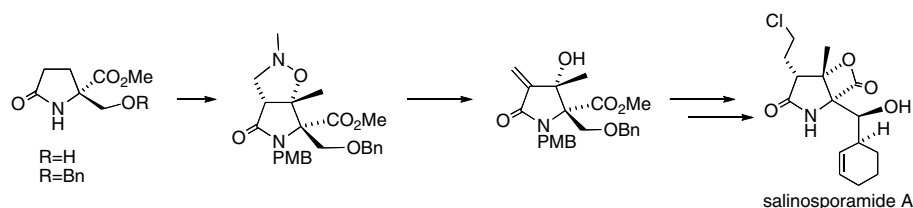


The key step involved a cyclocondensation of a thionium salt with an acetylenic ketone harbouring an acetal protected aldehyde function.

Stereoselective formal synthesis of the potent proteasome inhibitor: salinosporamide A

pp 381–384

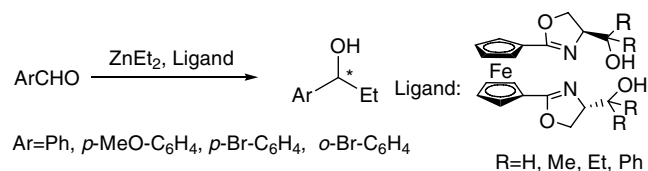
Virginie Caubert, Julien Massé, Pascal Retailleau and Nicole Langlois*



The synthesis and application of novel C₂-symmetric chiral N,N,O,O bisoxazoline ligands with a ferrocene backbone

pp 385–388

Genghong Hua, Delong Liu, Fang Xie and Wanbin Zhang*

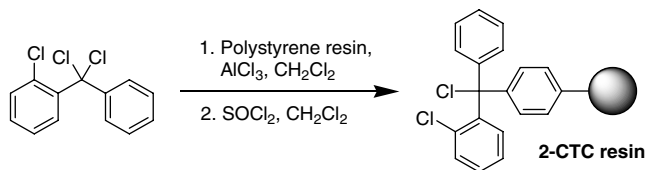


Novel C₂-symmetric bisoxazoline ligands **5** with a ferrocene backbone and a hydroxyl group at the substituent of the oxazoline ring were designed and prepared. With these ligands, up to 94% ee was obtained for the alkylation of arylaldehyde with diethylzinc.

A new method for the preparation of 2-chlorotrityl resin and its application to solid-phase peptide synthesis

pp 389–391

Tae-Kyung Lee, Sun-Jong Ryoo and Yoon-Sik Lee*



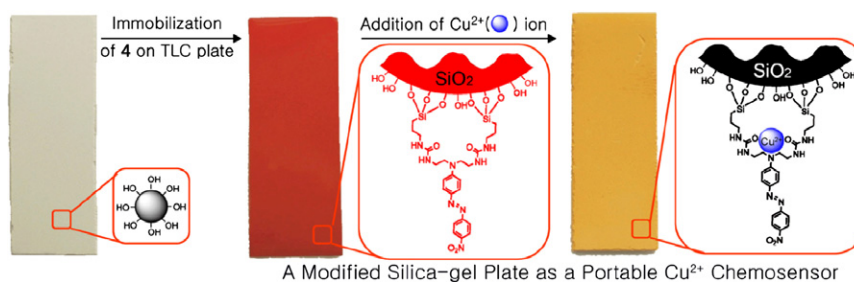
2-Chlorotritylchloride (CTC) resin was prepared efficiently from 1% DVB-crosslinked polystyrene resin and 1-chloro-2-(dichloro(phenyl)methyl)benzene. This 2-CTC resin showed excellent properties as a support for solid-phase synthesis. Four peptide fragments were obtained in high purity using the resin.



Azobenzene coupled chromogenic receptors for the selective detection of copper(II) and its application as a chemosensor kit

pp 393–396

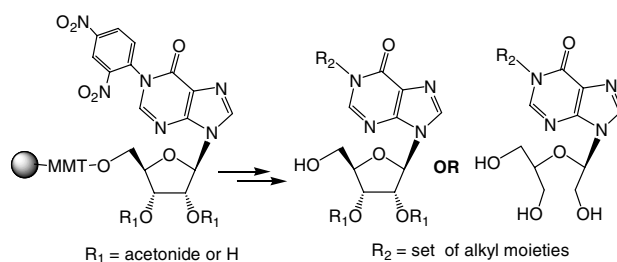
Soo Jin Lee, Shim Sung Lee, Il Yun Jeong, Jin Yong Lee and Jong Hwa Jung*



Synthesis of N-1 and ribose modified inosine analogues on solid support

pp 397–400

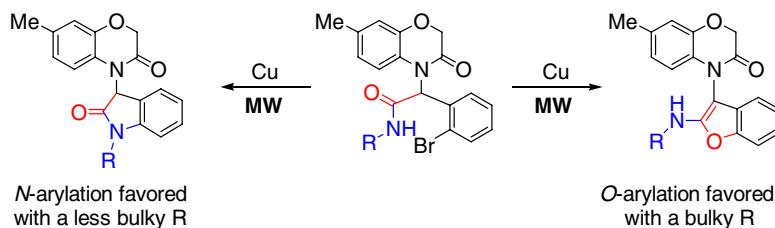
Giorgia Oliviero, Jussara Amato, Nicola Borbone, Stefano D'Errico, Gennaro Piccialli* and Luciano Mayol



Isolation and characterization of 2-alkylaminobenzo[b]furans. Evidence for competing O-arylation in Cu-catalyzed intramolecular amidation

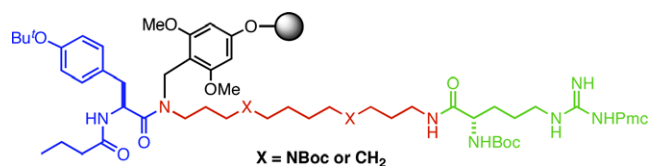
pp 401–404

Gaofeng Feng, Jinlong Wu and Wei-Min Dai*



Solid-phase synthesis of neuroactive spider–wasp hybrid toxin analogues using a backbone amide linker pp 405–408

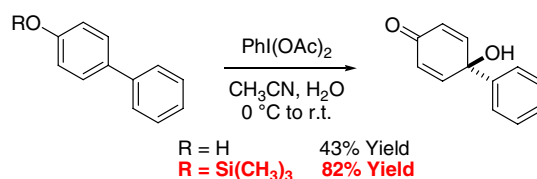
Christian A. Olsen,* Matthias Witt, Henrik Franzyk and Jerzy W. Jaroszewski



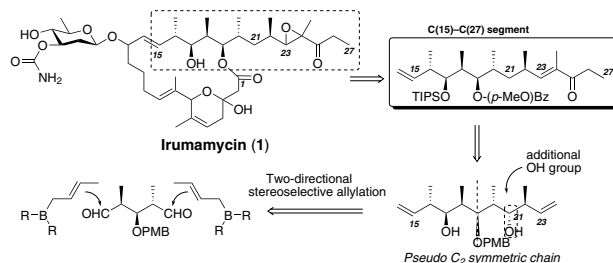
The efficient solid-phase synthesis (SPS) of novel hybrid toxins using a BAL resin is described. This strategy enables bidirectional construction of toxin molecules and has a potential in SPS of chemically diverse libraries of toxin analogues for structure–activity relationship (SAR) studies.

Oxidation of 4-arylphenol trimethylsilyl ethers to *p*-arylquinols using hypervalent iodine(III) reagents pp 409–412

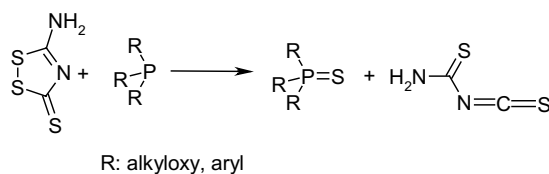
François-Xavier Felpin

**Studies toward the total synthesis of irumamycin: stereoselective preparation of the C(15)–C(27) segment via two-directional chain synthesis** pp 413–416

Tomoyasu Hirose, Toshiaki Sunazuka, Daisuke Yamamoto, Mutsumi Mouri, Yoshiaki Hagiwara, Takanori Matsumaru, Eisuke Kaji and Satoshi Omura*

**Evidence for the formation of isothiocyanate during sulfurisation of phosphines and phosphites using xanthane hydride** pp 417–419

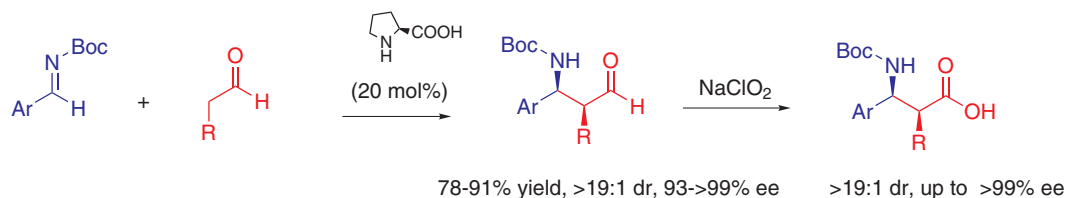
Jiří Hanusek, Mark A. Russell, Andrew P. Laws and Michael I. Page*



Highly enantioselective organocatalytic addition of unmodified aldehydes to *N*-Boc protected imines: one-pot asymmetric synthesis of β -amino acids

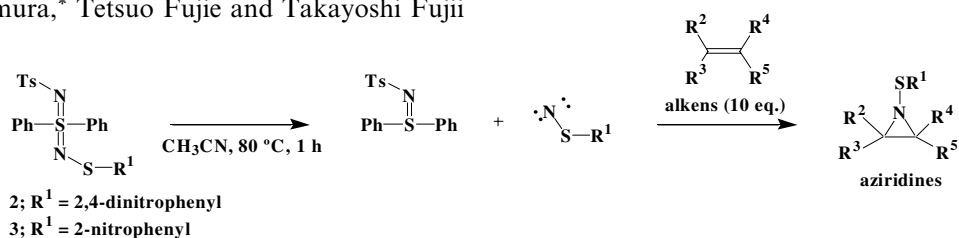
pp 421–425

Jan Vesely, Ramon Rios, Ismail Ibrahem and Armando Córdoba*


Moderate generation of sulfenylnitrenes from novel *N*-sulfenylsulfodiimides

pp 427–430

Toshiaki Yoshimura,* Tetsuo Fujie and Takayoshi Fujii

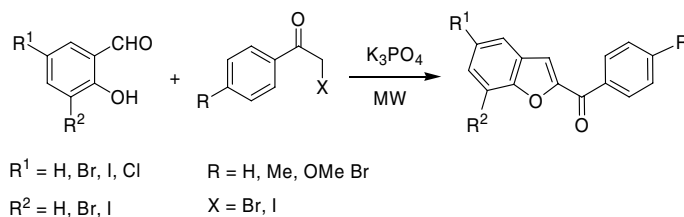


Thermolysis of *S,S*-diphenyl-*N*-(2-nitrophenylsulfenyl)- and *S,S*-diphenyl-*N*-(2,4-dinitrophenylsulfenyl)-sulfodiimides in the presence of olefins proceeded at 50–80 °C to give the corresponding deiminated *S,S*-diphenyl-*N*-tosylsulfimide and *N*-sulfenylaziridines in very good yields.

Microwave-mediated solvent free Rap–Stoermer reaction for efficient synthesis of benzofurans

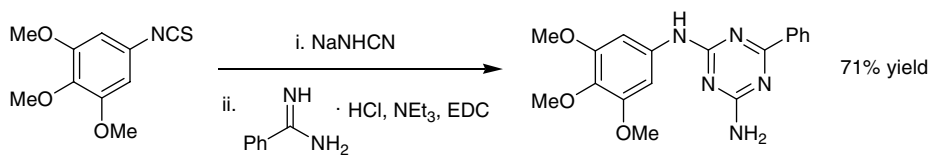
pp 431–434

Maddali L. N. Rao,* Dheeraj K. Awasthi and Debasis Banerjee

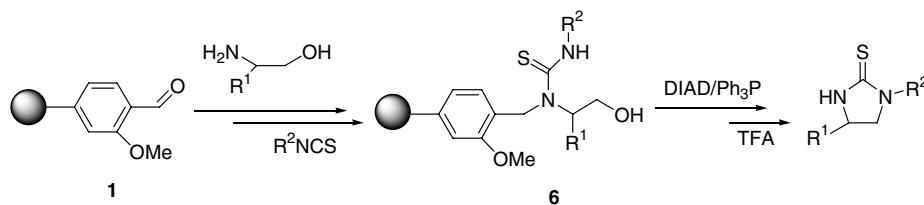

A novel one-pot synthesis of *N*,6-disubstituted 1,3,5-triazine-4,6-diamines from isothiocyanates and amidines

pp 435–437

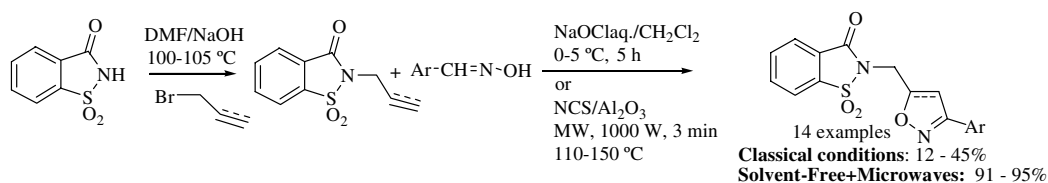
Chunjian Liu,* James Lin and Katerina Leftheris



Solid-phase synthesis of 2-imidazolidinethiones via Mitsunobu reaction of *N*-(2-hydroxyethyl)thioureas pp 439–441
Hyun Suk Jeon, Je Hwa Yoo, Jae Nyoung Kim and Taek Hyeon Kim*

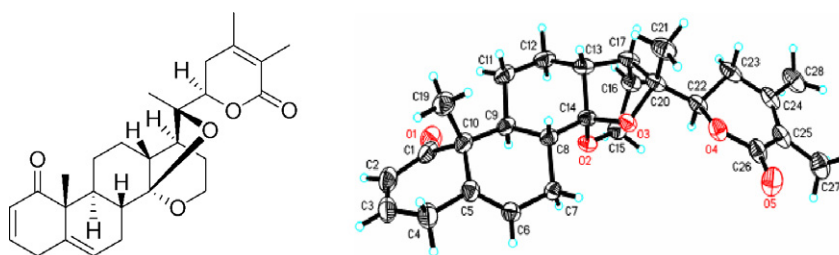


An efficient one-step regiospecific synthesis of novel isoxazolines and isoxazoles of *N*-substituted saccharin derivatives through solvent-free microwave-assisted [3+2] cycloaddition pp 443–447
Mahmoud Mabrou, Khalid Bougrin,* Rachid Benhida, André Loupy and Mohamed Soufiaoui

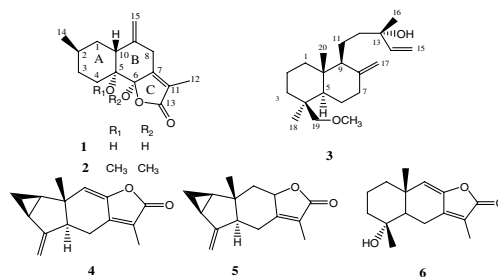


Withaphysanolide A, a novel *C*-27 norwithanolide skeleton, and other cytotoxic compounds from *Physalis divericata* pp 449–452

Lei Ma, Mumtaz Ali, Mohammad Arfan, Li-Guang Lou* and Li-Hong Hu*



Sesquiterpenoid with new skeleton from *Chloranthus henryi* pp 453–456
Bin Wu, Shan He and Yuanjiang Pan*

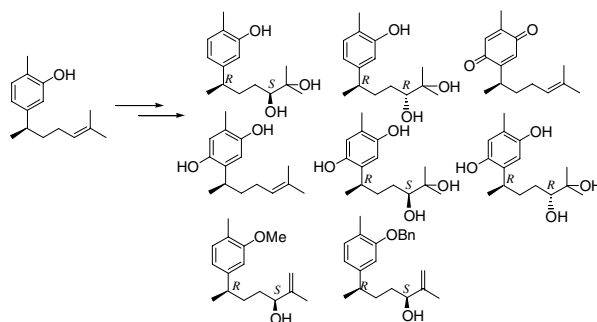


Dayejijiol (1), a novel sesquiterpene with a new carbon skeleton, and a novel labdane-type diterpenoid together with three known compounds were isolated.

Chemistry of xanthorrhizol: synthesis of several bisabolane sesquiterpenoids from xanthorrhizol

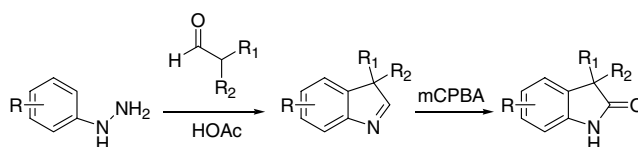
pp 457–460

Hasnah M. Sirat,* Ngai Mun Hong and Muhd Haffiz Jauri

**Synthesis of 3,3-disubstituted oxindoles**

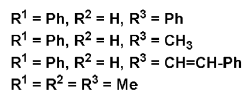
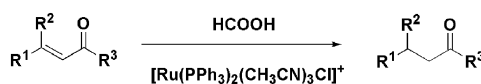
pp 461–463

Kevin G. Liu* and Albert J. Robichaud

**Regiospecific solvent-free transfer hydrogenation of α,β -unsaturated carbonyl compounds catalyzed by a cationic ruthenium(II) compound**

pp 465–467

Sipra Naskar and Manish Bhattacharjee*

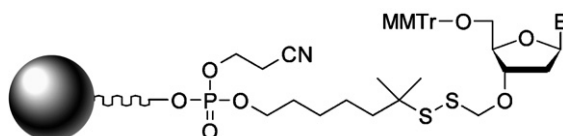


$[\text{Ru}(\text{PPh}_3)_2(\text{CH}_3\text{CN})_3\text{Cl}][\text{BPh}_4]$ catalyzes regiospecific transfer hydrogenation of the double bond in α,β -unsaturated carbonyl compounds.

A base-stable dithiomethyl linker for solid-phase synthesis of oligonucleotides

pp 469–472

Andrey Semenyuk and Marek Kwiatkowski*

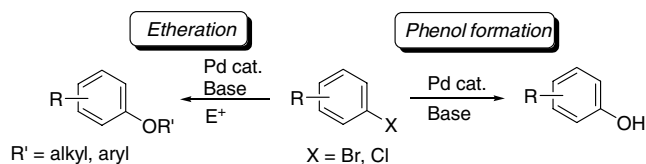


A novel linkage, useful for the synthesis of oligonucleotides is described. The linking function is compatible with all conditions used for oligonucleotide synthesis, orthogonal to all other protecting groups, but regenerates 3'-OH rapidly upon mild reduction under aqueous conditions. This method is employed in the removal of depurinated fragments during the synthesis of oligonucleotides.

Palladium-catalyzed C–O bond formation: direct synthesis of phenols and aryl/alkyl ethers from activated aryl halides

pp 473–476

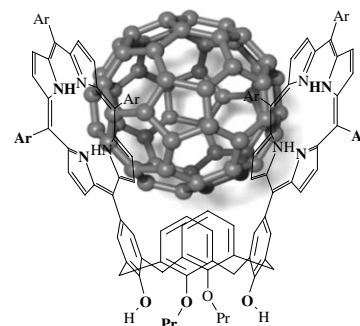
Guoshu Chen, Albert S. C. Chan and Fuk Yee Kwong*

**Novel fullerene receptors based on calixarene–porphyrin conjugates**

pp 477–481

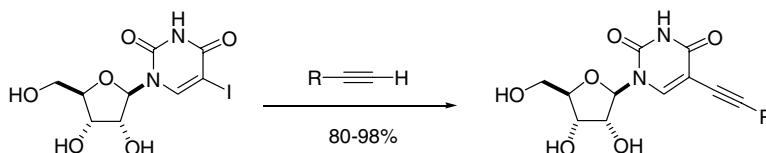
Martin Káš, Kamil Lang, Ivan Stibor and Pavel Lhoták*

Novel calix[4]arene–porphyrin conjugates, possessing a pronounced selectivity towards fullerene C₇₀, have been prepared using several different synthetic approaches.

**Efficient coupling of low boiling point alkynes and 5-iodonucleosides**

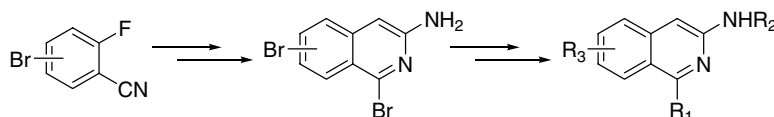
pp 483–486

Stephanos Ghilagaber, William N. Hunter and Rodolfo Marquez*

**An efficient synthesis of 1,6- and 1,7-dibromo-3-aminoisoquinolines: versatile templates for the preparation of functionalized isoquinolines**

pp 487–489

Mike Frohn,* Roland W. Bürli, Bobby Riahi and Randall W. Hungate

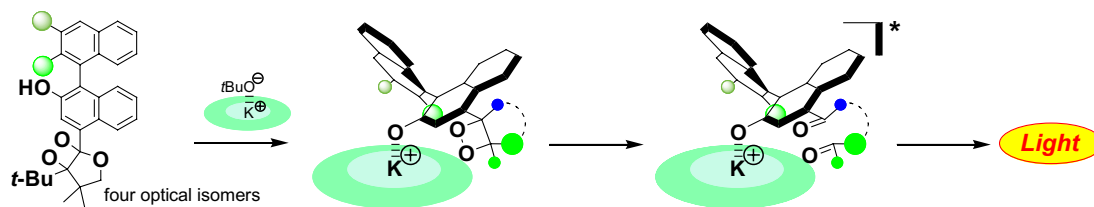


An efficient synthetic route to 1,6- and 1,7-dibromo-3-aminoisoquinoline was devised. These intermediates served as ideal templates for the preparation of 3-aminoisoquinoline analogues functionalized at C(6) or C(7).

Chemiluminescence in anisotropic microenvironment: splitting of chemiluminescence efficiency for charge-transfer-induced decomposition of optically active bicyclic dioxetanes bearing a 2-hydroxy-1,1'-binaphthyl-4-yl moiety under chiral recognition

pp 491–496

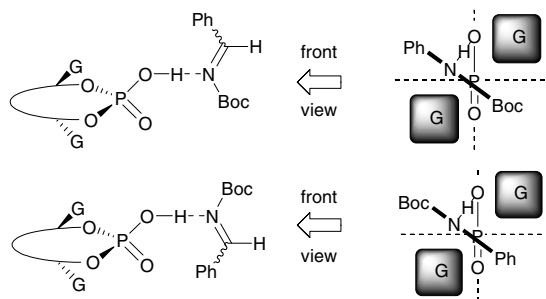
Masakatsu Matsumoto,* Hidetoshi Maeda, Naoyuki Hoshiya, Nobuko Watanabe and Hisako K. Ijuin



On the mechanism of stereoselection in direct Mannich reaction catalyzed by BINOL-derived phosphoric acids

pp 497–500

Ilya D. Gridnev,* Mitsuhiro Kouchi, Keiichi Sorimachi and Masahiro Terada*



OTHER CONTENTS

Corrigendum

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*Corresponding author

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