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Tetrahedron Vol. 63, No. 38, 2007

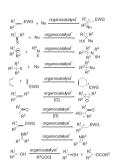
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REPORT

Asymmetric organocatalysis

Hélène Pellissier

This review is intended to update the impressive amount of recent developments of asymmetric organocatalysis in numerous reaction types, such as nucleophilic additions to electron-deficient C=C double bonds, nucleophilic additions to C=N double bonds, nucleophilic additions to unsaturated nitrogen, nucleophilic substitutions at aliphatic carbon, cycloaddition reactions, oxidations, reductions, kinetic resolutions and miscellaneous reactions, covering the literature from 2005 to 2007. This review clearly demonstrates the explosive growth and power of this new field of organic chemistry, which has become, in the last few years, the third methodology of asymmetric catalysis besides organometallic and enzymatic catalysis.



ARTICLES

A concise synthesis of (±)-pseudodeflectusin, an antitumor isochroman derivative isolated from *Aspergillus* sp.

pp 9333-9337

Makiko Tobe, Takuya Tashiro, Mitsuru Sasaki and Hirosato Takikawa*

Novel one-step synthesis of 2-carbonyl/thiocarbonyl isoindolinones and mechanistic disclosure on the pp 9338–9344 rearrangement reaction of o-phthalaldehyde with amide/thioamide analogs

Jieping Wan, Bin Wu and Yuanjiang Pan*

CHO
$$\frac{\text{CH}_3\text{CN/DMF}}{\text{or [bmim]BF}_4}$$
 $\frac{\text{R= aryl, alkyl}}{\text{TMSCI}}$
 $\frac{\text{R= aryl, alkyl}}{\text{R= alkyl, aryl or amidocyanogen}}$
 $\frac{\text{R= alkyl, aryl or amidocyanogen}}{\text{X= 0, S}}$



O-Alkylation versus C-alkylation under Mitsunobu conditions

pp 9345-9353

Catalina Gurgui-Ionescu, Loïc Toupet, Lycia Uttaro, Alain Fruchier and Véronique Barragan-Montero*

Synthesis and photoluminescence properties of BF2 complexes with 1,3-diketone ligands

pp 9354-9358

Katsuhiko Ono,* Kyohei Yoshikawa, Yujiro Tsuji, Hiroyuki Yamaguchi, Rie Uozumi, Masaaki Tomura, Keijiro Taga and Katsuhiro Saito

Reaction of β -lactam carbenes with alkyl isonitriles for a ready approach to 4-cyano and 4-carbamoyl pp 9359–9364 substituted β -lactams

Lan-Qing Cheng and Ying Cheng*

A simple and clean procedure for three-component synthesis of spirooxindoles in aqueous medium pp 9365–9372 Song-Lei Zhu, Shun-Jun Ji* and Yong Zhang

Effects of α -alkoxy substitution and conformational constraints on 6-exo radical cyclizations of hydrazones via reversible thiyl and stannyl additions

pp 9373-9381

Gregory K. Friestad* and Alex K. Mathies

Radical 6-exo cyclizations of imino compounds leading to aminosugar building blocks exhibit anomalous diastereoselectivity, attributable to dipole repulsion from an α -alkoxy substituent.



A solvent-controlled highly efficient Pd-C catalyzed hydrogenolysis of benzaldehydes to methylbenzenes via a novel 'acetal pathway'

pp 9382-9386

Lixin Xing, Xinyan Wang, Chuanjie Cheng, Rui Zhu, Bo Liu and Yuefei Hu*

(i)+

Diastereoselectivity in the Staudinger reaction: a useful probe for investigation of nonthermal microwave effects

pp 9387-9392

Libo Hu, Yikai Wang, Bonan Li, Da-Ming Du and Jiaxi Xu*

Nonthermal microwave effect in the Staudinger reaction was investigated using diastereoselectivity as a probe.

Design and synthesis of thioether-imidazolium chlorides as efficient ligands for palladium-catalyzed Suzuki-Miyaura coupling of aryl bromides with arylboronic acids

pp 9393-9400

Masami Kuriyama,* Rumiko Shimazawa and Ryuichi Shirai*

Practical synthesis and applications of benzoboroxoles

pp 9401-9405

Dinara S. Gunasekera, Dennis J. Gerold, Nathan S. Aalderks, J. Subash Chandra, Christiana A. Maanu, Paul Kiprof,* Viktor V. Zhdankin* and M. Venkat Ram Reddy*

A convenient one-pot synthesis of benzoboroxoles has been developed via the reaction of o-bromobenzyl alcohols with NaH, ⁿBuLi, and B(OⁱPr)₃ followed by acidic hydrolysis. Applications of these benzoboroxoles have been demonstrated in Pd-catalyzed cross-coupling reactions and the protocol has been extended for the synthesis of a chiral benzoboroxole. An exceptionally short synthesis of a potent antifungal agent AN2690 and several of its analogs has also been realized.

Methylene-2-ethynylcyclopropanes: synthesis and biological activity of (Z)- and (E)-9-{[2-ethynyl-2- pp 9406–9412 (hydroxymethyl)cyclopropylidene]methyl}adenine and -guanine

Shaoman Zhou, Mark N. Prichard and Jiri Zemlicka*

R = t-BuMe₂Si, B = adenine and guanine, Z- and E-isomers

Oxidation of acyclic monoterpenes by P450 BM-3 monooxygenase: influence of the substrate E/Z-isomerism on enzyme chemo- and regioselectivity

pp 9413-9422

Yomi Watanabe, Sabine Laschat, Michael Budde, Olena Affolter, Yuji Shimada and Vlada B. Urlacher*

The Z-isomers of monoterpenes were oxidized by a P450 BM-3 mutant to several products, including allylic alcohols and epoxides. *E*-isomers were epoxidized exclusively. Geranylacetone was converted with high activity and enantioselectivity to 9,10-epoxygeranylacetone.

Efficient synthesis of functionalized spiro-2,5-dihydro-1,2-λ⁵-oxaphospholes

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Issa Yavari,* Zinatossadat Hossaini, Maryam Sabbaghan and Majid Ghazanfarpour-Darjani

gem-Difluoromethylation of α - and γ -ketoesters: preparation of gem-difluorinated α -hydroxyesters and γ -butyrolactones

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Isoquinolone derivatives via a furan recyclization reaction

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Artem S. Dmitriev, Vladimir T. Abaev, Wolfgang Bender and Alexander V. Butin*

Expeditious synthesis of 5,6,7,8-tetrahydro-imidazo[1,2-a]pyrimidin-2-ones and 3,4,6,7,8,9-hexahydro-pyrimido[1,2-a]pyrimidin-2-ones

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Richa Pathak and Sanjay Batra*

Toward new camptothecins. Part 4: On the reactivity of nitro and amino precursors of aza analogs pp of 5-methoxycarbonyl camptothecin

pp 9456-9464

Laurent Gavara, Benoît Rigo,* Daniel Couturier, Laurence Goossens and Jean-Pierre Hénichart*

Synthesis of benzo analogs of oxoarcyriaflavins and caulersine

pp 9465-9475

Aurélie Bourderioux, Sylvain Routier,* Valérie Bénéteau* and Jean-Yves Mérour

$$\begin{array}{c} Z' \\ Z \\ Z \\ Z \\ Z' = (COXCO), X = 0, NH, NCH_3 \\ or Z = H, Z' = CO_2CH_3 \\ R1 = H, Br \\ R2 = H, CH_2CH_2N(CH_3)_2 \\ (R3, R4) = (=0), (H, H) \end{array}$$

Novel chemical cyclization routes to prepare ladder-type conjugated molecules

Inja Kim, Minji Yoo and Tae-Hyun Kim*

Convenient synthesis of photochromic symmetrical or unsymmetrical bis(heteroaryl)maleimides via pp 9482-9487 the Suzuki-Miyaura cross-coupling reaction

A. El Yahyaoui, G. Félix, A. Heynderickx, * C. Moustrou and A. Samat

O N O Suzuki-Miyaura cross-coupling O N O Ar = Ar' or Ar
$$\neq$$
 Ar'

Synthesis of photochromic bis(heteroaryl)maleimides, involving Suzuki-Miyaura cross coupling sequence, was developed on the basis of the reaction of diiodomaleimide with cyclic boronate esters. Behavior of the new products was examined.

Laccase-catalyzed conversion of green tea catechins in the presence of gallic acid to epitheaflagallin and pp 9488-9492 epitheaflagallin 3-O-gallate

Nobuya Itoh,* Yuji Katsube, Keiichi Yamamoto, Noriyuki Nakajima and Kenzaburo Yoshida

Epitheaflagallin (5) and epitheaflagallin 3-O-gallate (6), which are minor components of black tea, were preferentially synthesized from EGC (3) and EGCG (4) in green tea extracts using laccase and gallic acid. This biooxidation process could be applicable to the production of epitheaflagallin derivatives to improve the functionality of green tea.

pp 9476-9481

Novel peptidomimetic macrocycles showing exciplex fluorescence

pp 9493-9501

M. Isabel Burguete, Francisco Galindo,* M. Angeles Izquierdo, Santiago V. Luis* and Laura Vigara

EXCIPLEX FLUORESCENCE: 1a ~ 1b > 2a > 1c > 2b > 2c

Total synthesis of miraziridine A and identification of its major reaction site for cathepsin B Hiroyuki Konno,* Kanako Kubo, Hidefumi Makabe, Emi Toshiro, Naoyuki Hinoda, Kazuto Nosaka and Kenichi Akaji

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$Synthesis\ of\ all\ -cis\ -3\ -(2\ -diphenylphosphinoethyl)\ -1\ ,2\ ,4\ -tris(diphenylphosphinomethyl)\ cyclopentane\ (Ditricyp)\ from\ dicyclopentadiene$

pp 9514-9521

Isabelle Kondolff, Marie Feuerstein, Henri Doucet* and Maurice Santelli*

$$\begin{array}{c} \text{HO} \\ \text{OH} \end{array} \begin{array}{c} \text{OH} \\ \text{Ph}_2 \\ \text{Ph}_2 \\ \text{Ph}_2 \end{array} + \begin{array}{c} \text{Ph}_2 \\ \text{Ph}_2 \\ \text{Ph}_2 \\ \text{PPh}_2 \end{array}$$

The unique properties observed for the unsymmetrical macrocyclic compounds with the highly distorted structure

pp 9522-9530

Junko Morita, Shinji Tsuchiya, Nao Yoshida, Nirei Nakayama, Sayaka Yokokawa and Shojiro Ogawa*

A new class of aza-macrocycles with the highly distorted structure reacts with various lithium salts to form lithium complexes and their lithium complexation reactions depend on a substituent on the macrocyclic ring; slower rates and larger equilibrium constants were observed for the macrocycle with a bulkier substituent. The irradiation of these macrocycles by UV light was found to lead to the isomerization, and the photoisomerization rate of macrocycle with the bulky substituent was much faster.

A novel one-pot oxidative cyclization of 2'-amino and 2'-hydroxychalcones employing $FeCl_3 \cdot 6H_2O$ -methanol. Synthesis of 4-alkoxy-2-aryl-quinolines and flavones

pp 9531-9535

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In vivo transformations of dihydro-epi-deoxyarteannuin B in Artemisia annua plants Geoffrey D. Brown* and Lai-King Sy

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In vivo transformations of artemisinic acid in *Artemisia annua* plants Geoffrey D. Brown* and Lai-King Sy

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ISSN 0040-4020