AIMS AND SCOPE

Although total synthesis reached extraordinary levels of sophistication in the last century, the development of practical and efficient synthetic methodologies is still in its infancy. The goal of achieving chemical reactions that are economical, safe, environmentally benign, resource- and energy-saving will demand the highest level of scientific creativity, insight and understanding in a combined effort by academic and industrial chemists.

Advanced Synthesis & Catalysis is designed to stimulate and advance that process by focusing on the development and application of efficient synthetic methodologies and strategies in organic, bioorganic, pharmaceutical, natural product, macromolecular and materials chemistry. The targets of synthetic studies can range from natural products and pharmaceuticals to macromolecules and organic materials. While metal catalysis, biocatalysis and organocatalysis play an ever increasing role in achieving synthetic efficiency, all areas of interest to the practical synthetic chemist fall within the purview of Advanced Synthesis & Catalysis, including synthesis design, reaction techniques, separation science and process development.

Contributions from industrial and governmental laboratories are highly encouraged. It is the goal of the journal to help initiate a new era of chemical science, based on the efforts of synthetic chemists and on interdisciplinary collaboration, so that chemistry will make an even greater contribution to the quality of life than it does now.

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2007, 349, 4+5, Pages 477-768

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DEDICATED CLUSTER - COMMENTARY

In Honor of Professor Masakatsu Shibasaki on the Occasion of His $60^{\rm th}$ Birthday

Adv. Synth. Catal. 2007, 349, 491-492

Shū Kobayashi*

DEDICATED CLUSTER – REVIEW

Metallocene Sulfoxides as Precursors of Metallocenes with Planar Chirality

Adv. Synth. Catal. 2007, 349, 493-507

Benoit Ferber, Henri B. Kagan*

Fe Li 2 steps Fe S'''O i) t-BuLi Fe R^1 Fe = S'''O = i Fe = S'''O

491

493

DEDICATED CLUSTER - COMMUNICATIONS

509 Catalytic Asymmetric Aldol Reactions of Enolizable Carbon Pronucleophiles with Formaldehyde and Ethyl Glyoxylate

Adv. Synth. Catal. 2007, 349, 509-512

Isao Fukuchi, Yoshitaka Hamashima, Mikiko Sodeoka*

$$R^{1} = R^{2} = R^{2} = R^{1} = R^{2} = R^{1} = R^{2} = R^{1} = R^{2} = R^{2$$

513 Asymmetric 1,4-Addition of Organoboron Reagents to Quinone Monoketals Catalyzed by a Chiral Diene/Rhodium Complex: A New Synthetic Route to Enantioenriched 2-Aryltetralones

Adv. Synth. Catal. 2007, 349, 513-516

☐ Norihito Tokunaga, Tamio Hayashi*

517 Highly Efficient Asymmetric Hydrogenation of α,β Unsaturated Carboxylic Acids Catalyzed by Ruthenium(II)Dipyridylphosphine Complexes

Adv. Synth. Catal. 2007, 349, 517-520

Liqin Qiu, Yue-Ming Li, Fuk Yee Kwong, Wing-Yiu Yu, Qing-Hua Fan, Albert S. C. Chan*

$$R^{2} \xrightarrow{R^{1}} \frac{H_{2}, Ru(II)-L}{MeOH} \xrightarrow{R^{2}} \overset{R^{1}}{\underset{\longleftarrow}{R^{0}}}$$

 $\label{eq:Xyl-P-Phos} \mbox{ Xyl-P-Phos, Ar} = 3.5 - (\mbox{CH}_3)_2 \mbox{C}_6 \mbox{H}_3$ $\mbox{L} = \mbox{Xyl-P-Phos, conv.} > 99\%; \mbox{ $ee} \mbox{ up to } 97\%$

521 Catalytic Enantioselective Tandem Carbonyl Ylide Formation/1,3-Dipolar Cycloaddition Reactions of α-Diazo Ketones with Aromatic Aldehydes using Dirhodium(II) Tetrakis[*N*-benzene-fused-phthaloyl-(*S*)-valinate]

Adv. Synth. Catal. 2007, 349, 521-526

Hideyuki Tsutsui, Naoyuki Shimada, Takumi Abe, Masahiro Anada, Makoto Nakajima, Seiichi Nakamura, Hisanori Nambu, Shunichi Hashimoto*

527 Tin-Free Radical Carbonylation: Synthesis of Acylated Oxime Ethers Using Alkyl Allyl Sulfone Precursors, Carbon Monoxide, and Phenylsulfonyl Oxime Ether

Adv. Synth. Catal. 2007, 349, 527 – 530

Sangmo Kim, Kyoung-Chan Lim, Sunggak Kim,* Ilhyong Ryu

531

Polymer Incarcerated Ruthenium Catalyst for Oxidation of Alcohols with Molecular Oxygen

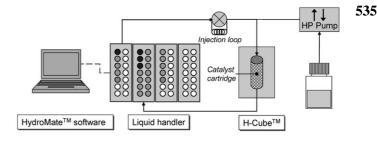
Adv. Synth. Catal. 2007, 349, 531-534

Tsutomu Matsumoto, Masaharu Ueno, Juta Kobayashi, Hiroyuki Miyamura, Yuichiro Mori, Shū Kobayashi*

Optimisation of Conditions for *O*-Benzyl and *N*-Benzyloxycarbonyl Protecting Group Removal using an Automated Flow Hydrogenator

Adv. Synth. Catal. 2007, 349, 535-538

Kristian Rahbek Knudsen, John Holden, Steven V. Ley,* Mark Ladlow



Zirconium-Catalyzed Asymmetric Carboalumination of Alkenes: ZACA–Lipase-Catalyzed Acetylation Synergy

Adv. Synth. Catal. 2007, 349, 539-545

Zhihong Huang, Ze Tan, Tibor Novak, Gangguo Zhu, Ei-ichi Negishi*

Simple Ruthenium Precatalyst for the Synthesis of Stilbene Derivatives and Ring-Closing Metathesis in the Presence of Styrene Initiators

Adv. Synth. Catal. 2007, 349, 546-550

Cheikh Lo, Renan Cariou, Cédric Fischmeister,* Pierre H. Dixneuf*

Dehydrative Cyclization Catalyzed by the Combination of Molybdenum(VI) Oxides and Benzoic Acids: First Synthesis of the Antitumour Substance BE-70016

Adv. Synth. Catal. 2007, 349, 551-555

Akira Sakakura, Shuhei Umemura, Rei Kondo, Kazuaki Ishihara*

556 Effects of Aromatic Substituents on Binaphthyl-Based Chiral Spiro-Type Ammonium Salts in Asymmetric Phase-Transfer Reactions

Adv. Synth. Catal. 2007, 349, 556-560

Taichi Kano, Quan Lan, Xisheng Wang, Keiji Maruoka*

561 Characterization of Dimeric and Tetrameric μ-Hydroxide Ytterbium(III) Binaphtholate Complexes

Adv. Synth. Catal. 2007, 349, 561-565

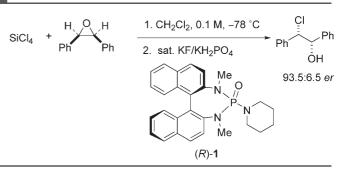
Alfred J. Wooten, Luca Salvi, Patrick J. Carroll, Patrick J. Walsh*

DEDICATED CLUSTER - FULL PAPERS

Enantioselective Ring Opening of Epoxides with Silicon Tetrachloride in the Presence of a Chiral Lewis Base: Mechanism Studies

Adv. Synth. Catal. 2007, 349, 567-582

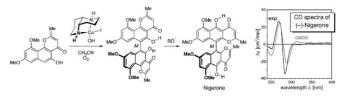
Scott E. Denmark,* Paul A. Barsanti, Gregory L. Beutner, Tyler W. Wilson



583 Asymmetric Total Synthesis of Nigerone and *ent*-Nigerone: Enantioselective Oxidative Biaryl Coupling of Highly Hindered Naphthols

Adv. Synth. Catal. 2007, 349, 583-594

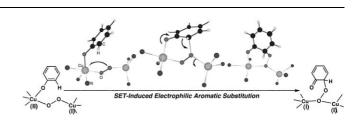
Marisa C. Kozlowski,* Elizabeth C. Dugan, Evan S. DiVirgilio, Katja Maksimenka, Gerhard Bringmann*



Theoretical Studies on *ortho*-Oxidation of Phenols with Dioxygen Mediated by Dicopper Complex: Hints for a Catalyst with the Phenolase Activity of Tyrosinase

Adv. Synth. Catal. 2007, 349, 595-600

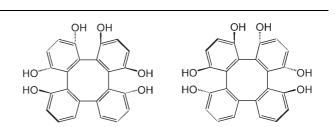
Hiroshi Naka,* Yoshinori Kondo, Shinya Usui, Yuichi Hashimoto, Masanobu Uchiyama*



601 Synthesis of Enantiopure (S,R,S)- and (R,S,R)-1,4,5,8,9,16-Hexahydroxytetraphenylenes

Adv. Synth. Catal. 2007, 349, 601-608

An-Hui Wu, Chun-Kit Hau, Henry N. C. Wong*



609

Chiral Ionic Phosphites and Diamidophosphites: A Novel Group of Efficient Ligands for Asymmetric Catalysis

Adv. Synth. Catal. 2007, 349, 609-616

Konstantin N. Gavrilov,* Sergey E. Lyubimov, Oleg G. Bondarev, Marina G. Maksimova, Sergey V. Zheglov, Pavel V. Petrovskii, Vadim A. Davankov, Manfred T. Reetz*

Dynamic Kinetic Resolution for the Catalytic Asymmetric Total Synthesis of Antithrombotic Agents M58163 and M58169

Adv. Synth. Catal. 2007, 349, 617-628

Fumihiko Saitoh,* Hidemitsu Nishida, Takafumi Mukaihira, Kohsuke Aikawa, Koichi Mikami*

Peptidic Amidomonophosphane Ligand for Copper-Catalyzed Asymmetric Conjugate Addition of Diorganozincs to Cycloalkenones

Adv. Synth. Catal. 2007, 349, 629-635

Takahiro Soeta, Khalid Selim, Masami Kuriyama, Kiyoshi Tomioka*

Reaction of Tetrahydropyran Ethers with Triethylsilyl Trifluoromethanesulfonate–2,4,6-Collidine Combination: Speculation on the Intermediate, Efficient Deprotection, and Application to Efficient Ring-Closing Metathesis as a Tether

Adv. Synth. Catal. 2007, 349, 636-646

Hiromichi Fujioka,* Takashi Okitsu, Takuya Ohnaka, Yoshinari Sawama, Ozora Kubo, Kazuhisa Okamoto, Yasuyuki Kita*

$$\begin{array}{c} \text{TESOTf} \\ \text{Z.4.6-collidine} \\ \text{CH}_2\text{Cl}_2, 0 \, ^{\circ}\text{C} \\ \end{array} \\ \begin{array}{c} \text{R. Alkyl and aryl units} \\ \text{some of which include} \\ \text{acid-labile functional} \\ \text{groups} \\ \text{TES} = \text{Et}_{\text{IS}}\text{i} \\ \text{Tf} = \text{CF}_3\text{SO}_2 \\ \end{array} \\ \begin{array}{c} \text{R. Alkyl and aryl units} \\ \text{R. Alkyl aryl units} \\ \text{R. Alkyl and aryl units} \\ \text{R. Alkyl aryl$$

Synthesis of Biaryls *via* Palladium-Catalyzed [2+2+2] Cocyclization of Arynes and Diynes: Application to the Synthesis of Arylnaphthalene Lignans

Adv. Synth. Catal. 2007, 349, 647-661

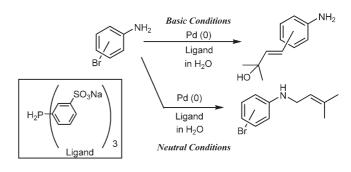
Yoshihiro Sato,* Takayuki Tamura, Atsushi Kinbara, Miwako Mori

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662 Chemoselective Palladium-Catalyzed Reaction in Aqueous Media: Selectivity in the Reaction of Haloanilines with 1,1-Dimethylallyl Alcohol

Adv. Synth. Catal. 2007, 349, 662-668

Yuusaku Yokoyama,* Noriko Takagi, Hidemasa Hikawa, Satoru Kaneko, Natsume Tsubaki, Hiroaki Okuno



669 Generation and Application of *o*-Quinone Methides Bearing Various Substituents on the Benzene Ring

Adv. Synth. Catal. 2007, 349, 669-679

Hiromichi Sugimoto, Satoshi Nakamura, Tomohiko Ohwada*

- 680 Suppression of β-Hydride Elimination in the Intramolecular Hydrocarboxylation of Alkynes leading to the Formation of Lactones

 $\begin{array}{c}
\text{Pd}(PPh_3)_4/(o\text{-tol})_3P \\
-COOH
\end{array}$ Toluene, 100 °C

Adv. Synth. Catal. 2007, 349, 680-684

Zhibao Huo, Nitin T. Patil, Tienan Jin, Nirmal K. Pahadi, Yoshinori Yamamoto*

Yoshinori Yamamoto*

3-piperidenes, Promising Chiral Building Blocks, by Palladium-Catalyzed Deracemization of Their Alkyl

A New Preparation of Homochiral N-Protected 5-Hydroxy-

Pd₂(dba)₃-CHCl₃ (2 mol %)

ONH HN

8 mol %

PPh₂ Ph₂P

CH₂Cl₂/H₂O (9:1)

P = Ts 93% (99% ee)

P = Boc 94% (94% ee)

P = Cbz 88% (87% ee)

P = COOMe 90% (99% ee)

Adv. Synth. Catal. **2007**, 349, 685–693

Carbonates

Hiroki Takahata,* Yumiko Suto, Erina Kato, Yuichi Yoshimura, Hidekazu Ouchi

A Direct Organocatalytic Entry to Selectively Protected Aldopentoses and Derivatives

Adv. Synth. Catal. 2007, 349, 694-702

Christoph Grondal, Dieter Enders*

R = aryl, alkyl; n = 1, 2

685

703

709

DEDICATED CLUSTER - UPDATES

Diarylmethanols by Catalyzed Asymmetric Aryl Transfer Reactions onto Aldehydes Using Boronic Acids as Aryl Source

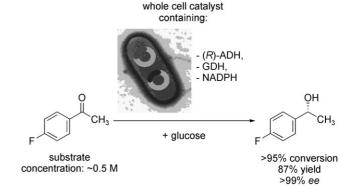
Adv. Synth. Catal. 2007, 349, 703-708

Frank Schmidt, Jens Rudolph, Carsten Bolm*

Enantioselective Reduction of 4-Fluoroacetophenone at High Substrate Concentration using a Tailor-Made Recombinant Whole-Cell Catalyst

Adv. Synth. Catal. 2007, 349, 709-712

Harald Gröger,* Claudia Rollmann, Francoise Chamouleau, Isabelle Sebastien, Oliver May, Wolfgang Wienand, Karlheinz Drauz



COMMUNICATION

Dioxomolybdenum(VI)-Catalyzed Reductive Cyclization of Nitroaromatics. Synthesis of Carbazoles and Indoles

Adv. Synth. Catal. 2007, 349, 713-718

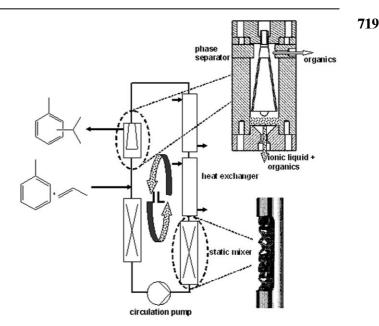
Roberto Sanz,* Jaime Escribano, María R. Pedrosa, Rafael Aguado, Francisco J. Arnáiz*

FULL PAPERS

Continuous, Ionic Liquid-Catalysed Propylation of Toluene in a Liquid-Liquid Biphasic Reaction Mode using a Loop Reactor Concept

Adv. Synth. Catal. 2007, 349, 719-726

Viktor Ladnak, Norbert Hofmann, Nicole Brausch, Peter Wasserscheid*



727 A Comparative Study of the Synthesis of 3-Substituted Catechols using an Enzymatic and a Chemoenzymatic Method

Adv. Synth. Catal. 2007, 349, 727-739

V. Berberian, C. C. R. Allen, N. D. Sharma, D. R. Boyd,* and C. Hardacre*

740 A New Imidazole-Containing Imidazolidinone Catalyst for Organocatalyzed Asymmetric Conjugate Addition of Nitroalkanes to Aldehydes

Adv. Synth. Catal. 2007, 349, 740-748

Leila Hojabri, Antti Hartikka, Firouz Matloubi Moghaddam, Per I. Arvidsson*

749 Diversity-Oriented Preparation of Enantiopure Spirocyclic 2-Azetidinones from α-Oxo-β-lactams through Barbier-Type Reactions followed by Metal-Catalyzed Cyclizations

Adv. Synth. Catal. 2007, 349, 749-758

Benito Alcaide,* Pedro Almendros,* Teresa Martínez del Campo, Raquel Rodríguez-Acebes

$$R^{4n} \xrightarrow{X} R^{2} \longrightarrow R^{5} \xrightarrow{HQ} H R^{2} \longrightarrow 0 \xrightarrow{R^{1}} R^{2} \longrightarrow 0 \xrightarrow{R^{1}} R^{2}$$

 $n = 0, 1; X = O, H_2; Y = NH, CH_2$ $R^5 =$ allyl, propargyl, allenyl, 1,3-butadien-2-yl

UPDATE

759 Metal-Free, Enantioselective Strecker Reactions Catalyzed by Chiral BINOL and TADDOL Catalysts

Adv. Synth. Catal. 2007, 349, 759-764

Magnus Rueping,* Erli Sugiono, Stefan A. Moreth

CORRIGENDUM

In the paper by A. Gheorghe, A. Matsuno, and O. Reiser in Issue 9, 2006, pp. 1016-1020 (DOI: 10.1002/adsc.200606043), the molar amounts of co-catalysts Mn(NO₃)₂·4 H₂O and CoNO₃·6 H₂O are incorrect: in Table 3 (p 1018) and in the *General Procedure for the Aerobic Oxidation* (p 1019), the correct amounts of co-catalysts are 0.02 mmol instead of 0.2 mmol; in Table 4 (p. 1018) and in the procedure *Aerobic Oxidation of 4-Bromobenzyl Alcohol PS-CLICK-TEMPO; Recycling Experiments* (page 1019), 0.06 mmol of the co-catalysts instead of 0.6 mmol were employed. The absolute amounts employed were stated correctly.

Supporting information on the WWW (see article for access details).

*Author to whom correspondence should be addressed.