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. Achieving chemical reactions that are highly selective, economical, safe, resource- and energy-efficient, and environmentally benign is a primary challenge to chemistry in this century. Realizing this goal will demand the highest level of scientific creativity, insight and understanding in a combined effort by academic, government and industrial chemists and engineers.

Advanced Synthesis & Catalysis promotes that process by publishing high-impact research results reporting the development and application of efficient synthetic methodologies and strategies for organic targets that range from pharmaceuticals to organic materials. Homogeneous catalysis, biocatalysis, organocatalysis and heterogeneous catalysis directed towards organic synthesis are playing an ever increasing role in achieving synthetic efficiency. Asymmetric catalysis remains a topic of central importance. In addition, Advanced Synthesis & Catalysis includes other areas that are making a contribution to green synthesis, such as synthesis design, reaction techniques, flow chemistry and continuous processing, multiphase catalysis, green solvents, catalyst immobilization and recycling, separation science and process development.

Practical processes involve development of effective integrated strategies, from an elegant synthetic route based on mechanistic and structural insights at the molecular level through to process optimization at larger scales. These endeavors often entail a multidisciplinary approach that spans the broad fields chemistry, biology, and engineering and involve contributions from academic, government, and industrial laboratories.

The unique focus of *Advanced Synthesis & Catalysis* has rapidly made it a leading organic chemistry and catalysis journal. The goal of *Advanced Synthesis & Catalysis* is to help inspire 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.



succeeding Journal für praktische Chemie (founded in 1828)

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2008, 350, 4, Pages 521-636

Issue 3/2008 was published online on February 19, 2008

COMMUNICATIONS

Highly Efficient Molybdenum(II)-Catalyzed Intramolecular Allylic Alkylation of Arenes

Adv. Synth. Catal. 2008, 350, 531-536

Marco Bandini,* Astrid Eichholzer, Peter Kotrusz, Achille Umani-Ronchi*

OCO₂Me
$$[Mo(CO)_4Br_2]_2$$

$$(2.5 \text{ mol}\%)$$

$$E = CO_2Et$$

$$[Mo(CO)_4Br_2]_2$$

$$(2.5 \text{ mol}\%)$$

$$x \text{ if } E$$

$$E$$

$$yield 48 - 97\%$$

$$14 \text{ examples}$$

Synthesis of 2,3,4-Trisubstituted Thiochromanes using an Organocatalytic Enantioselective Tandem Michael-Henry Reaction

Adv. Synth. Catal. 2008, 350, 537-541

Rajasekhar Dodda, Joshua J. Goldman, Tanmay Mandal, Cong-Gui Zhao,* Grant A. Broker, Edward R. T. Tiekink

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531

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542 Recyclable Carbon Supported Copper-Manganese Oxide for Selective Aerobic Oxidation of Alcohols in Combination with 2,2,6,6-Tetramethylpiperidyl-1-oxyl under Neutral

Condition 30 - 80 °C Cu-Mn oxide supported on active carbon Adv. Synth. Catal. 2008, 350, 542-546

- Guanyu Yang,* Weimin Zhu, Panke Zhang, Huazhen Xue, Wei Wang, Junshan Tian, Maoping Song*
- 547 Cyclization of Allenyne-1,6-diols Catalyzed by Gold and Silver Salts: An Efficient Selective Synthesis of Dihydrofuran and Furan Derivatives

Adv. Synth. Catal. 2008, 350, 547-551

Sundae Kim, Phil Ho Lee*

2:3 = 2.2 - 7.5:1 $R = n-Pr, c-C_6H_{11}, Ph, 4-Cl-C_6H_4$ $R = n-Pr, c-C_6H_{11}, Ph, 4-Cl-C_6H_4$ 3-MeO-C₆H₄, 4-Ac-C₆H₄ 3-MeO-C₆H₄, 4-MeO-C₆H₄

4-Ac-C₆H₄, 2-furyl

552 Palladium-Catalyzed/Lewis Acid-Promoted Alkene Dimerization and Cross-Coupling with Alcohols via C-H **Bond Activation**

Adv. Synth. Catal. 2008, 350, 552-556

- 🖳 Yi-Jun Jiang, Yong-Qiang Tu,* En Zhang, Shu-Yu Zhang, Ke Cao, Lei Shi

557 Eight-Step Synthesis of Routiennocin

Adv. Synth. Catal. 2008, 350, 557-560

Kenji Matsumoto, Sergey A. Kozmin*

- Routiennocin
- 561 Novel Chiral Biheteroaromatic Diphosphine Oxides for Lewis Base Activation of Lewis Acids in Enantioselective Allylation and Epoxide Opening

Adv. Synth. Catal. 2008, 350, 561-564

Valentina Simonini, Maurizio Benaglia,* Tiziana Benincori*

524

565

571

577

591

602

Catalytic Dehydrogenation of o-Alkylated or o-Alkoxylated Iodoarenes with Concomitant Hydrogenolysis

Adv. Synth. Catal. 2008, 350, 565-569

Elena Motti, Marta Catellani*

FULL PAPERS

Titanium-Catalyzed Enantioselective Synthesis of α-Ambrinol

Adv. Synth. Catal. 2008, 350, 571-576

José Justicia,* Araceli G. Campaña, Btissam Bazdi, Rafael Robles, Juan M. Cuerva,* J. Enrique Oltra*

Hybrid Organic-Inorganic Materials from Di-(2pyridyl)methylamine-Palladium Dichloride Complex as Recoverable Catalysts for Suzuki, Heck and Sonogashira Reactions

Adv. Synth. Catal. 2008, 350, 577-590

Montserrat Trilla, Roser Pleixats,* Michel Wong Chi Man,* Catherine Bied, Joël J. E. Moreau

Catalysis of Salicylaldehydes and Two Different C-H Acids with Electricity: First Example of an Efficient Multicomponent Approach to the Design of Functionalized Medicinally Privileged 2-Amino-4H-Chromene Scaffold

Adv. Synth. Catal. 2008, 350, 591-601

Michail N. Elinson,* Alexander S. Dorofeev,* Fedor M. Miloserdov, Alexey I. Ilovaisky, Sergey K. Feducovich, Pavel A. Belyakov, Gennady I. Nikishin

Mn (cat) Ti (cat) ³.Ĥ (-)- α -ambrinol

SiO_{1.5} •n SiO₂ n = 10, 20, 40

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0.1-0.2 F/mol

stronger C-H acid

Сĺ ĊΙ

weaker C-H acid

15 examples, 65-86%

Simple Methodology for Heck Arylation at C-8 of Adenine Nucleosides

Adv. Synth. Catal. 2008, 350, 602-608

Pallavi Lagisetty, Li Zhang, Mahesh K. Lakshman*

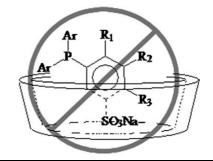
 NH_2 TBDMSC TBDMSO R = OTBDMS TBDMSO R = HTBDMSO

UPDATES

609 Biphasic Aqueous Organometallic Catalysis Promoted by Cyclodextrins: How to Design the Water-Soluble Phenylphosphane to Avoid Interaction with Cyclodextrin

Adv. Synth. Catal. 2008, 350, 609-618

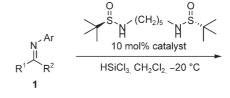
Michel Ferreira, Hervé Bricout, Adlane Sayede, Anne Ponchel, Sophie Fourmentin, Sébastien Tilloy, Eric Monflier*



Rationally-Designed S-Chiral Bissulfinamides as Highly Enantioselective Organocatalysts for Reduction of Ketimines

Adv. Synth. Catal. 2008, 350, 619-623

Dong Pei, Yu Zhang, Siyu Wei, Meng Wang, Jian Sun*



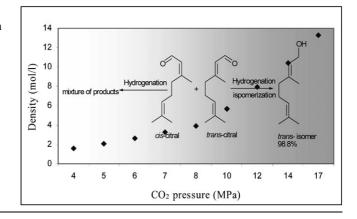
 R^{1} R^{2}

up to 95% yield up to 96% ee

624 Density-Dependent Formation of the Pure *trans*-Isomer of Unsaturated Alcohol by Selective Hydrogenation of Citral in Supercritical Carbon Dioxide

Adv. Synth. Catal. 2008, 350, 624-632

M. Chatterjee,* Y. Ikushima, T. Yokoyama, M. Sato



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

*Author to whom correspondence should be addressed.