AIMS AND SCOPE

While 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 catalytic methods based on metal complexes or enzymes 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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Synthesis &
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2006, 348, 10+11, Pages 1113-1316

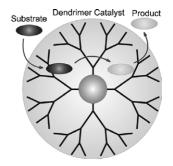
Issue 9/2006 was published online on June 28, 2006

REVIEW

The Dendrimer Effect in Homogeneous Catalysis

Adv. Synth. Catal. 2006, 348, 1125-1148

Brett Helms, Jean M. J. Fréchet*



COMMUNICATIONS

Activation of Nucleophilic Fluorination by Salts in Ionic Liquids and in Sulfolane

Adv. Synth. Catal. 2006, 348, 1149-1153

Stéphane Anguille, Maxime Garayt, Vincent Schanen, René Grée*

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1125

1149

1154 Potassium Fluoride/Basic Alumina as Far Superior Heterogeneous Catalyst for the Chemoselective Conjugate Addition of Nitroalkanes to Electron-Poor Alkenes Having Two Electron-Withdrawing Groups in α- and β-Positions

Adv. Synth. Catal. 2006, 348, 1154-1156

- Roberto Ballini,* Alessandro Palmieri
- **1157** Asymmetric Hydrogenation of β-Keto Esters Using Chiral Diphosphonites

$$R^1$$
 OR^2 R_0 OR_2 R_1 OR_2 R_1 OR_2

Adv. Synth. Catal. 2006, 348, 1157-1160

Manfred T. Reetz,* Xiaoguang Li

1161 Catalytic Enantioselective Conjugate Addition of Nitromethane to α'-Hydroxy Enones as Surrogates of α,β-Unsaturated Carboxylic Acids and Aldehydes

Adv. Synth. Catal. 2006, 348, 1161-1164

Claudio Palomo,* Raquel Pazos, Mikel Oiarbide, Jesús M. García

1165 Enzymatic Transformations; 61. Preparation of Enantiopure Trifluoromethyl-Substituted Aromatic Epoxides and Vicinal Diols using the *Aspergillus niger* Epoxide Hydrolase-Catalysed Resolution

Adv. Synth. Catal. 2006, 348, 1165-1169

Justine Deregnaucourt, Alain Archelas, Fabien Barbirato, Jean-Marc Paris, Roland Furstoss*

$$\label{eq:resolvent} \begin{split} \mathbf{R} &= \mathbf{H}, \, \mathbf{C} \mathbf{H}_3 \\ \mathbf{X} &= \, \, \mathbf{C} \mathbf{F}_3, \, \mathbf{O} \mathbf{C} \mathbf{F}_3, \, \mathbf{S} \mathbf{C} \mathbf{F}_3 \end{split}$$

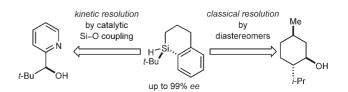
ee >98% 95% > ee > 60%

FULL PAPERS

1171 Preparation of a Privileged Silicon-Stereogenic Silane: Classical *versus* Kinetic Resolution

Adv. Synth. Catal. 2006, 348, 1171-1182

Sebastian Rendler, Gertrud Auer, Manfred Keller, Martin Oestreich*



1183 The 3-(3-Pyridine)propionyl Anchor Group for Protease-Catalyzed Resolutions: *p*-Toluenesulfinamide and Sterically Hindered Secondary Alcohols

Adv. Synth. Catal. 2006, 348, 1183-1192

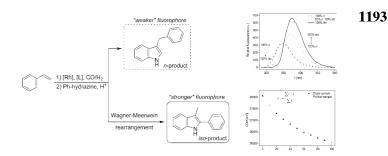
Christopher K. Savile, Romas J. Kazlauskas*

- increases solubility in water
- enhances binding to subtilisin, chymotrypsin
- simplifies separation by acid extraction

A Rapid and Reliable Assay for Regioselectivity Using Fluorescence Spectroscopy

Adv. Synth. Catal. 2006, 348, 1193-1199

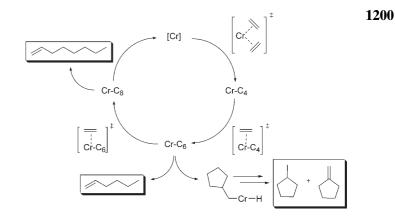
Goran Angelovski,* Mark D. Keränen,* Petra Linnepe, Stefan Grudzielanek, Peter Eilbracht*



Influence of Elevated Temperature and Pressure on the Chromium-Catalysed Tetramerisation of Ethylene

Adv. Synth. Catal. 2006, 348, 1200-1206

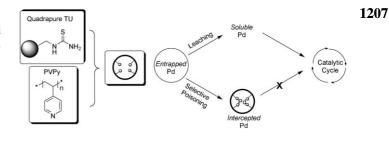
Sven Kuhlmann, John T. Dixon, Marco Haumann, David H. Morgan, Jimmy Ofili, Oliver Spuhl, Nicola Taccardi, Peter Wasserscheid*



Poly(4-vinylpyridine) and Quadrapure TU as Selective Poisons for Soluble Catalytic Species in Palladium-Catalyzed Coupling Reactions – Application to Leaching from Polymer-Entrapped Palladium

Adv. Synth. Catal. 2006, 348, 1207-1216

John M. Richardson, Christopher W. Jones*



Subsequent Enzymatic Galactosylation and Sialylation Towards Sialylated Thomsen-Friedenreich Antigen Components

Adv. Synth. Catal. 2006, 348, 1217-1227

Lars Kröger, Agnes Scudlo, Joachim Thiem*

Efficient, Enantioselective Organocatalytic Synthesis of Trichostatin A

Adv. Synth. Catal. 2006, 348, 1228-1234

Shilei Zhang, Wenhu Duan,* Wei Wang*

1228

1217

1235 Efficient Preparation of New Rhodium- and Iridium-[Bis(oxazolinyl)-3,5-dimethylphenyl] Complexes by C-H Bond Activation: Applications in Asymmetric Synthesis

Adv. Synth. Catal. 2006, 348, 1235-1240

Jun-ichi Ito, Takushi Shiomi, Hisao Nishiyama*

1241 Iron-Salen Complexes as Efficient Catalysts in Ring Expansion Reactions of Epoxyalkenes

Adv. Synth. Catal. 2006, 348, 1241-1247

Gerhard Hilt,* Christian Walter, Patrick Bolze

1248 Improved Enantioselectivity of Immobilized Chiral Bisoxazolines by Partial Precapping of the Siliceous Mesocellular Foam Support with Trimethylsilyl Groups

Adv. Synth. Catal. 2006, 348, 1248-1254

Su Seong Lee, Sukandar Hadinoto, Jackie Y. Ying*

1255 Efficient Carbonylation of Aryl and Heteroaryl Bromides using a Palladium/Diadamantylbutylphosphine Catalyst

Adv. Synth. Catal. 2006, 348, 1255-1261

Helfried Neumann, Anne Brennführer, Peter Groß, Thomas Riermeier, Juan Almena,* Matthias Beller*

Pd(OAc)_{2,} CO

cataCXium[®] A, R'OH

R

1262 Palladium-Catalyzed Stereoselective Synthesis of (*E*)-Stilbenes *via* Organozinc Reagents and Carbonyl Compounds

Adv. Synth. Catal. 2006, 348, 1262-1270

Jin-Xian Wang,* Kehu Wang, Lianbiao Zhao, Hongxia Li, Ying Fu, Yulai Hu

FG-Ar
$$Ar^{1}$$
 Ar^{1}
 Ar^{1}
 Ar^{1}
 Ar^{1}

FG-Ar

FG-Ar

FG-Ar

 Ar^{1}

FG-Ar

 Ar^{1}

FG-Ar

 Ar^{1}
 Ar^{1}

1271 Asymmetric Hydrogenation of α,β-Unsaturated Carboxylic Acids Catalyzed by Ruthenium(II) Complexes of Spirobifluorene Diphosphine (SFDP) Ligands

Adv. Synth. Catal. 2006, 348, 1271-1276

Xu Cheng, Jian-Hua Xie, Sheng Li, Qi-Lin Zhou*

COOH
$$H_2/Cat.$$
 (R)-1b $COOH$ Ar_2 Ar_2 P $Ru(OAc)_2$ R Ar_2 R Ar_3 Ar_4 R Ar_4 R Ar_5 R Ar_5 R Ar_6 Ar_6 R Ar_6 $Ar_$

1277

1283

1289

1301

1306

Ruthenium-Catalyzed Enantioselective Reduction of Electron-Rich Aryl Alkyl Ketones

Adv. Synth. Catal. 2006, 348, 1277-1282

Jenny Wettergren, Anders Bøgevig, Maud Portier, Hans Adolfsson*

Heterogeneous Gold-Catalysed Synthesis of Phenols

Adv. Synth. Catal. 2006, 348, 1283-1288

Silvio Carrettin, M. Carmen Blanco, Avelino Corma,* A. Stephen K. Hashmi*

Asymmetric Reductive Amination: Convenient Access to Enantioenriched Alkyl-Alkyl or Aryl-Alkyl Substituted α -Chiral Primary Amines

Adv. Synth. Catal. 2006, 348, 1289-1299

Thomas C. Nugent,* Abhijit K. Ghosh, Vijay N. Wakchaure, Rashmi R. Mohanty

$$R_L$$
 Overall yields 64, 71 – 78% $\stackrel{\text{NH}_2}{=}$ $\stackrel{\text{NH}_2}{=}$ or $\stackrel{\text{NH}_2}{=}$ $\stackrel{\text{NH}$

Alkyl alkyl or aryl alkyl prochiral ketones (acyclic and cyclic examples)

UPDATES

2,3-Disubstituted Indoles through the Palladium-Catalyzed Reaction of Aryl Chlorides with *o*-Alkynyltrifluoro-acetanilides

R
+ ArCI
Pd₂(dba)_{3,} Xphos
Cs₂CO₃

MeCN, 120 °C

Ar

N
H

Sandro Cacchi,* Giancarlo Fabrizi, Antonella Goggiamani

Facile Synthesis of a Monosulfonated Triphenylphosphane (TPPMS) Derived Ligand having Strong π -Acceptor Character

Adv. Synth. Catal. 2006, 348, 1306-1310

Henrik Gulyás, Zoltán Bacsik, Áron Szöllősy, József Bakos*

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

stAuthor to whom correspondence should be addressed.