### **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)

ASC
5-Year Impact Factor 2007
5.193
The Cutting Edge that Stays Sharp!

2009, 351, 9, Pages 1169-1452

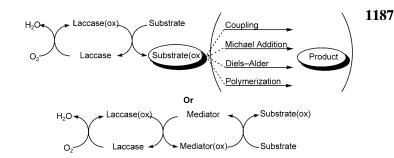
Issue 7 + 8/2009 was published online on May 12, 2009

#### REVIEW

Synthetic Applications of Laccase in Green Chemistry

Adv. Synth. Catal. 2009, 351, 1187-1209

Suteera Witayakran, Arthur J. Ragauskas\*

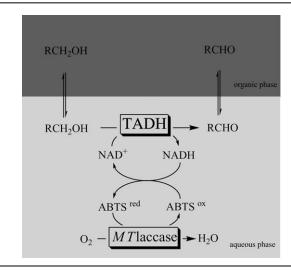


# COMMUNICATIONS

**1211** A New Regeneration System for Oxidized Nicotinamide Cofactors

Adv. Synth. Catal. 2009, 351, 1211-1216

Seda Aksu, Isabel W. C. E. Arends,\* Frank Hollmann\*



1217 Remarkable Electronic Effect on the Diastereoselectivity of the Heck Reaction of Methyl Cinnamate with Arenediazonium Salts: Formal Total Synthesis of  $(\pm)$ -Indatraline and  $(\pm)$ -Sertraline

Adv. Synth. Catal. 2009, 351, 1217-1223

Julio Cezar Pastre, Carlos Roque Duarte Correia\*

1224 Highly Enantioselective Construction of the  $\alpha$ -Chiral Center of Amides via Iridium-Catalyzed Hydrogenation of  $\alpha,\beta$ -Unsaturated Amides

Adv. Synth. Catal. 2009, 351, 1224-1228

☐ Wei-Jing Lu, Xue-Long Hou\*

$$R^{2} \xrightarrow{H} \frac{H_{2} \text{ (50bar)}}{\text{catalyst (2 mol\%)}} \xrightarrow{R^{2}} \frac{H}{N} \xrightarrow{R^{3}}$$

$$R^{1} = \text{Ar, alkyl; } R^{2} = \text{alkyl, H;}$$

$$R^{3} = i \cdot \text{Bu, Bn}$$

$$R^{2} \xrightarrow{H} \frac{100\% \text{ conversion}}{84 - 98\% \text{ ee}}$$

 $\frac{1}{2}$ 

catalyst

Fe

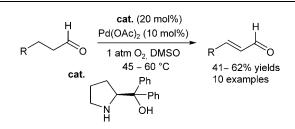
Ir COD

BAR<sub>F</sub>

1229 A Direct Amine-Palladium Acetate Cocatalyzed Saegusa Oxidation Reaction of Unmodified Aldehydes to  $\alpha,\beta$ -Unsaturated Aldehydes

Adv. Synth. Catal. 2009, 351, 1229-1232

☐ Jin Zhu, Jie Liu, Ruoqun Ma, Hexin Xie, Jian Li,\*
Hualiang Jiang, Wei Wang\*



1233

1238

1250

1255

1263

Cobalt/Rhodium Heterobimetallic Nanoparticle-Catalyzed Oxidative Carbonylation of Amines in the Presence of Carbon Monoxide and Molecular Oxygen to Ureas

Adv. Synth. Catal. 2009, 351, 1233-1237

☐ Ji Hoon Park, Jae Chun Yoon, Young Keun Chung\*

The Use of Samarium Enolates, A Novel Alternative in the Addition Reactions to Imines. Synthesis of 3-Amino Esters, Amides and Enantiopure 3,4-Diamino Esters

Adv. Synth. Catal. 2009, 351, 1238-1242

Carmen Simal

José M. Concellón,\* Humberto Rodríguez-Solla,

Copper-Catalyzed Stereoselective Hydroarylation of 3-Aryl-2-propynenitriles with Arylboronic Acids

Adv. Synth. Catal. 2009, 351, 1243-1249

Naohiro Kirai Yoshihiko Yamamoto,\* Tsuyoshi Asatani, Naohiro Kirai

17 examples, 68 - 97% isolated yields

Copper(I)-Catalyzed Asymmetric Addition of Terminal Alkynes to  $\beta$ -Imino Esters: An Efficient and Direct Method in the Synthesis of Chiral  $\beta^3$ -Alkynyl  $\beta^{2,2}$ -Dimethyl Amino Acid Derivatives

Adv. Synth. Catal. 2009, 351, 1250-1254

Jun Wang, Zhihui Shao, Kai Ding Wing Yiu Yu\* Albert S. C. Chan\*

Versatile Supramolecular Copper(II) Complexes for Henry and Aza-Henry Reactions

Adv. Synth. Catal. 2009, 351, 1255-1262

Guoqi Zhang, Eiji Yashima, Wolf-D. Woggon\*

$$R + CH_3NO_2$$

$$X = O, N-Boc; XH: OH, NH-Boc$$
up to 99% ee

Iron-Catalyzed Cross-Coupling Reactions of Terminal Alkynes with Vinyl Iodides

Adv. Synth. Catal. 2009, 351, 1263-1267

Xin Xie, Xiaobing Xu, Hongfeng Li, Xiaolei Xu, Jingyu Yang, Yanzhong Li\*

R<sup>1</sup>— 
$$R^2$$
  $\xrightarrow{\text{Cat. FeCl}_3, \text{ Ligand}}$   $R^1$   $\xrightarrow{\text{R}^2}$   $R^2$   $\xrightarrow{\text{Toluene. 110 °C}}$   $R^2$ 

 $\begin{array}{c|c} R^1 \\ \hline \\ R^2 \end{array}$ 

1268 Iron-Catalyzed Cross-Coupling Reaction of Vinyl Bromides or Chlorides with Imidazoles in the Absence of Ligands and Additives

Adv. Synth. Catal. 2009, 351, 1268-1272

- Jincheng Mao,\* Guanlei Xie, Jiaming Zhan, Qiongqiong Hua, Daqing Shi
- (X = Br, CI)10 - 20% FeCl<sub>3</sub> PO4, DMSO, 120 °C X = BrX = CIyield up to 87% Z/E up to 10:90 yield up to >99% Z/E up to >99:1
- 1273 Pyrrolidine-Camphor Derivative as an Organocatalyst for Asymmetic Michael Additions of α,α-Disubstituted Aldehydes to β-Nitroalkenes: Construction of Quaternary Carbon-Bearing Aldehydes under Solvent-Free Conditions

Adv. Synth. Catal. 2009, 351, 1273-1278

- Chihliang Chang, Ssu-Hsien Li, Raju Jannapu Reddy, Kwunmin Chen\*
- (20 mol%) PhCO<sub>2</sub>H (20 mol%)  $R^1$ *Neat*, 0 °C, 0.5 – 3 d yield up to 99% ee up to 95%
- 1279 Simple and Fast Synthesis of New Axially Chiral Bipyridine N,N'-Dioxides for Highly Enantioselective Allylation of Aldehydes

Adv. Synth. Catal. 2009, 351, 1279-1283

- Aneta Kadlčíková, Radim Hrdina, Irena Valterová, Martin Kotora\*
- (R,R)-1 or (S,R)-1OH (1 mol%) THF –78 °C, 1 h R = MeO, Me, H, Cl, F, CF<sub>3</sub> ee 91 - 96% Ō (S,R)-1
  - (R,R)-1
- 1284 Highly Enantioselective Organocatalytic syn- and anti-Aldol Reactions in Aqueous Medium

Adv. Synth. Catal. 2009, 351, 1284-1288

- Monika Raj, Gopal S. Parashari, Vinod K. Singh\*
- **1a/**TFA, H₂O R¹ **1a/**TFA, DMF:H₂O de = 98:2 ee = >99% ee = >99%  $R^2 = H$  $R^2$  = H, COOR Ph
- 1289 Sulfur-Participated Nazarov-Type Cyclization: A Simple and Efficient Synthesis for 3-Thio-1*H*-indenes

Adv. Synth. Catal. 2009, 351, 1289-1292

Hongwei Zhou,\* Yongfa Xie, Lianjun Ren, Kai Wang

$$R^{1}$$
  $R^{2}$   $R^{2}$   $R^{2}$   $R^{2}$   $R^{2}$   $R^{3}$   $R^{2}$   $R^{2}$   $R^{3}$   $R^{3$ 

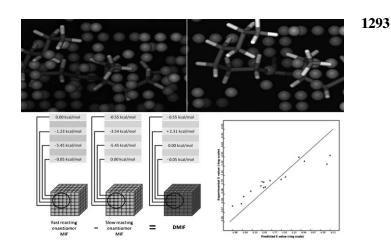
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### **FULL PAPERS**

A Three-Dimensional Quantitative Structure-Activity Relationship (3D-QSAR) Model for Predicting the Enantioselectivity of *Candida antarctica* Lipase B

Adv. Synth. Catal. 2009, 351, 1293-1302

Paolo Braiuca, Knapic Lorena, Valerio Ferrario, Cynthia Ebert, Lucia Gardossi\*



One-Pot Multienzymatic Synthesis of 12-Ketoursodeoxycholic Acid: Subtle Cofactor Specificities Rule the Reaction Equilibria of Five Biocatalysts Working in a Row

Adv. Synth. Catal. 2009, 351, 1303-1311

Daniela Monti,\* Erica Elisa Ferrandi, Ilaria Zanellato, Ling Hua, Fausto Polentini, Giacomo Carrea, Sergio Riva\* Double oxidation

To-HSDH
12α-HSDH
12α

Poly(trimethylene carbonate) from Biometals-Based Initiators/Catalysts: Highly Efficient Immortal Ring-Opening Polymerization Processes

Adv. Synth. Catal. 2009, 351, 1312-1324

Marion Helou, Olivier Miserque, Jean-Michel Brusson, Jean-François Carpentier,\* Sophie M. Guillaume\*

$$n \quad O \quad N(SiMe_3)_2$$

$$ROH (m = 1-200 \text{ equiv. vs. Zn})$$

$$bulk, 60 - 110 \, ^{\circ}\text{C}, < 2 \text{ h}$$

$$n \quad M$$

n up to 50,000 equiv. (20 ppm Zn catalyst)

TON up to 46,500 mol(TMC)·mol(Zn) $^{-1}$ TOF up to 31, 000 mol(TMC)·mol(Zn) $^{-1}$ ·h $^{-1}$ 

Catalytic Asymmetric Cycloaddition of Carbon Dioxide and Propylene Oxide Using Novel Chiral Polymers of BINOL-Salen-Cobalt(III) Salts

Adv. Synth. Catal. 2009, 351, 1325-1332

Peng Yan, Huanwang Jing\*

1325

1303

1312

**1333** Stereoselectivity in the Rhodium-Catalysed Reductions of Non-Conjugated Dienes

Adv. Synth. Catal. 2009, 351, 1333-1343

Bao Nguyen, John M. Brown\*

- OBu-t

  2 1

  for H<sub>2</sub> addition; Rh cat.
- 1344 Recyclable Polymer- and Silica-Supported Ruthenium(II)-Salen Bis-pyridine Catalysts for the Asymmetric Cyclopropanation of Olefins

Adv. Synth. Catal. 2009, 351, 1344-1354

- Christopher S. Gill, Krishnan Venkatasubbaiah, Christopher W. Jones\*
- Phi COOEt + Phi COOEt + Phi COOE (1R,2R)
- 1355 Primary Amine-Thioureas based on *tert*-Butyl Esters of Natural Amino Acids as Organocatalysts for the Michael Reaction

Adv. Synth. Catal. 2009, 351, 1355-1362

Christoforos G. Kokotos, George Kokotos\*

f-BuO  $\bigvee_{O}$   $\bigvee_{H}$   $\bigvee_{H}$   $\bigvee_{N}$   $\bigvee_{N$ 

$$\label{eq:R} \begin{split} R &= CH_3, CH_2C_6H_5, CH(CH_3)_2, CH_2CO_2Bu-t, CH_2CH_2CO_2Bu-t, CH_2OBu-t, CH(CH_3)OBu-t, CH_2C_6H_4OBu-t \end{split}$$

1363 Anionic Bridged Bis(amidinate) Lithium Lanthanide Complexes: Efficient Bimetallic Catalysts for Mild Amidation of Aldehydes with Amines

Adv. Synth. Catal. 2009, 351, 1363-1370

- ☐ Junfeng Wang, Junmei Li, Fan Xu, Qi Shen\*
- 1371 Synthesis of Trisubstituted Pyrroles from Rhodium-Catalyzed Alkyne Head-to-Tail Dimerization and Subsequent Gold-Catalyzed Cyclization

Adv. Synth. Catal. 2009, 351, 1371-1377

- Hong Mei Peng, Jing Zhao,\* Xingwei Li\*
- 1378 Atom-Efficient, Palladium-Catalyzed Stille Coupling Reactions of Tetraphenylstannane with Aryl Iodides or Aryl Bromides in Polyethylene Glycol 400 (PEG-400)

Adv. Synth. Catal. 2009, 351, 1378-1382

- Wen-Jun Zhou, Ke-Hu Wang, Jin-Xian Wang\*
- R + SnPh<sub>4</sub> PdCl<sub>2</sub>(PPh<sub>3</sub>)<sub>2</sub> R PEG-400, NaOAc, 100 °C

X = I, Br $R = H, 4-CI, 4-NO_2 4-CH_3 4-OCH_3 4-CHO, 4-CN, 4-COCH_3 etc.$ 

1383

1390

1395

Efficient, Nickel-Catalysed Kumada-Tamao-Corriu Cross-Coupling with a Calix[4]arene-Diphosphine Ligand

Adv. Synth. Catal. 2009, 351, 1383-1389

Laure Monnereau, David Sémeril,\* Dominique Matt,\* Loïc Toupet, Antonio J. Mota

PhMgBr + Ar-X 
$$\frac{[(\eta^5-C_5H_5)Ni(L)]BF_4 (2)}{100 °C, dioxane} Ph-Ar$$
ArX·Ni = 50000

TOF up to 21,250 mol(ArBr)  $\cdot$ mol(Ni)<sup>-1</sup>  $\cdot$ h<sup>-1</sup>

Hydrosilylation of Ketone and Imine over Poly-N-Heterocyclic Carbene Particles

Adv. Synth. Catal. 2009, 351, 1390-1394

MeiXuan Tan, Yugen Zhang,\* Jackie Y. Ying\*

$$R^{1}$$
  $R^{2}$  +  $Ph_{2}SiH_{2}$   $Ph_{2}HSi$   $Ph_{2}$ 

Improved Protocols for Molybdenum- und Tungsten-Catalyzed Hydrostannations

Adv. Synth. Catal. 2009, 351, 1395-1404

Alexander O. Wesquet, Uli Kazmaier\*

AcO

TFAHN

COO-
$$t$$
-Bu

Bu<sub>3</sub>SnH

M(CO)<sub>3</sub>(CNR)<sub>3</sub>

TFAHN

AcO

SnBu<sub>3</sub>

TFAHN

COO- $t$ -Bu

M = Mo, W

90%, 91% rs

An Efficient Solvent-Free Route to Silyl Esters and Silyl Ethers

Adv. Synth. Catal. 2009, 351, 1405-1411

[RuCl<sub>2</sub>(p-cymene)]<sub>2</sub>

🖳 Yuko Ojima, Kazuya Yamaguchi, Noritaka Mizuno\*

1405 [RuCl<sub>2</sub>(p-cymene)]<sub>2</sub> R-OH + HSiR'3 R-OSiR'

Effective Chiral Ferrocenyl Phosphine-Thioether Ligands in Enantioselective Palladium-Catalyzed Allylic Alkylations

Adv. Synth. Catal. 2009, 351, 1412-1422

Hong Yee Cheung, Wing-Yiu Yu, Terry T. L. Au-Yeung, Zhongyuan Zhou, Albert S. C. Chan\*

Asymmetric Hydrogenation of α,β-Unsaturated Ester-Phosphonates

Adv. Synth. Catal. 2009, 351, 1423-1430

Yange Huang, Florian Berthiol, Bart Stegink, Michael M. Pollard, Adriaan J. Minnaard\*

1423

## **UPDATES**

**1431** Palladium-Catalyzed Three-Component 1:2:1 Coupling of Aryl Iodides, Alkynes, and Alkenes to Produce 1,3,5-Hexatriene Derivatives

Adv. Synth. Catal. 2009, 351, 1431-1436

Hakaru Horiguchi, Koji Hirano, Tetsuya Satoh,\*
Masahiro Miura\*

**1437** Iridium-Catalysed Asymmetric Hydrogenation of Enamides in the Presence of 3,3'-Substituted H8-Phosphoramidites

Adv. Synth. Catal. 2009, 351, 1437-1441

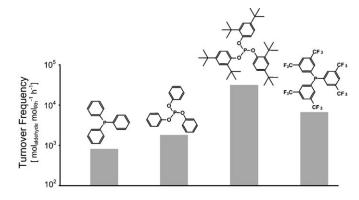
Giulia Erre, Stephan Enthaler, Kathrin Junge, Daniele Addis, Matthias Beller\*

$$L = \begin{array}{c} & & & \\ & &$$

1442 Efficient Hydroformylation in Dense Carbon Dioxide using Phosphorus Ligands without Perfluoroalkyl Substituents

Adv. Synth. Catal. 2009, 351, 1442-1450

Ard C. J. Koeken,\* Nieck E. Benes, Leo J. P. van den Broeke, Jos T. F. Keurentjes



### CORRIGENDA

In the paper by Yong-Chua Teo in Issue 5, 2009, pp. 720–724 (DOI: 10.1002/adsc.200800746), the author failed to make direct reference to very similar work published by Correa and Bolm in 2007 (reference 4r: A. Correa, C. Bolm, *Angew. Chem.* 2007, 119, 9018; *Angew. Chem. Int. Ed.* 2007, 46, 8862). The paper by Correa and Bolm had already reported very similar screening for the same reaction in toluene with the same starting materials and had already disclosed the best catalyst system, FeCl<sub>3</sub> (10 mol%), dmeda (20 mol%), K<sub>3</sub>PO<sub>4</sub> (2 equiv.), the same optimized catalyst system reported by Teo for the reaction in water. Tables 3 and 4 in the paper by Correa and Bolm report almost all the substrates and products subsequently reported in the Teo paper in Tables 2 and 3; again the difference being that Teo used water as solvent instead of toluene. The tolerance of the cross-coupling process to water had also been disclosed in the paper by Correa and Bolm.

In addition, the author also failed to cross reference very similar work using a cobalt catalyst system, which he simultaneously published in Chemistry, a European Journal (Y.-C. Teo, G.-L. Chua, *Chem. Eur. J.* **2009**, *15*, 3072). The cobalt paper and the iron paper report similar goals, reactions, ligand scanning, starting materials, products, and conclusions.

The author apologizes for these oversights and agrees that the record be set straight by means of this corrigendum.

Finally, in Scheme 1 on page 723, the structure at the top of the catalytic cycle should be without "X". The correct scheme is as follows:

In the paper by Jeanne L. Bolliger and Christian M. Frech in Issue 6, 2009, pp. 891–902 (DOI: 10.1002/adsc.200900112), the graphic abstract in the table of contents of the issue on page 809 incorrectly shows two methyl groups bound to palladium in the pincer complex, which should be removed. The correct graphic abstract is as follows:

 $NR_2$  = piperidinyl; Y = NH or O X = I or Br

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

\*Author to whom correspondence should be addressed.