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
2-Year Impact Factor 2008
5.619
A Record High
for Organic Chemistry

2009, 351, 16, Pages 2533-2732

Issue 14+15/2009 was published online on October 16, 2009

COMMUNICATIONS

Asymmetric Hydrogenation of Quinoxalines Catalyzed by Iridium/PipPhos

Adv. Synth. Catal. 2009, 351, 2549-2552

Nataša Mršić, Thomas Jerphagnon, Adriaan J. Minnaard,* Ben L. Feringa,* Johannes G. de Vries*

Axial Chirality Control by 2,4-Pentanediol for the Alternative Synthesis of C₃*-TunePhos Chiral Diphosphine Ligands and Their Applications in Highly Enantioselective Ruthenium-Catalyzed Hydrogenation of β-Keto Esters

Adv. Synth. Catal. 2009, 351, 2553-2557

☐ Xianfeng Sun, Wei Li, Guohua Hou, Le Zhou, Xumu Zhang*

$$R^{1} \longrightarrow OR^{2} \longrightarrow \frac{1 \text{ mol}\% \text{ RuL*Cl}_{2}(\text{DMF})_{n}}{H_{2} (50 \text{ atm}), 80 \text{ °C}, 12 \text{ h}} \longrightarrow R^{1} \longrightarrow OR^{2}$$

$$ee \text{ up to } 99\%$$

$$L^{1} = \bigvee_{i=1}^{n} O \longrightarrow PAr_{2}$$

$$Ar = 3,5-Me_{2}C_{6}H_{3}$$

2549

2553

2558 Palladium on Charcoal as a Recyclable Catalyst for C–S Cross-Coupling of Thiols with Aryl Halides under Ligand-Free Conditions

Adv. Synth. Catal. 2009, 351, 2558-2562

....,

Zheng Jiang, Jin She, Xufeng Lin*

2 mol% Pd/C 1.5 equiv. KOH DMSO 110 °C, 9 h

R = functional group; R' = Ar, alkyl

up to 99% yield

2563 Enantioselective Organocatalytic Conjugate Addition of Aldehydes to α,β -Unsaturated Thiol Esters

Adv. Synth. Catal. 2009, 351, 2563-2566

Shaolin Zhu, You Wang, Dawei Ma*

- O O Catalyst O SR²

 | A Catalyst O SR² | 2 20 mol% Catalyst O SR² | SR
 - catalyst: Ph Ph Ph $R^1 = CO_2R^3$, $CONR^4R^5$ OTMS
- **2567** Highly Efficient Synthesis of Quaternary α-Hydroxy Phosphonates *via* Lewis Acid-Catalyzed Hydrophosphonylation of Ketones

Adv. Synth. Catal. 2009, 351, 2567-2572

- Xin Zhou, Yanling Liu, Lu Chang, Jiannan Zhao, Deju Shang, Xiaohua Liu, Lili Lin, Xiaoming Feng*
- $R^{1} \stackrel{\bigcirc{}}{ \stackrel{}{ \coprod}} R^{2} \stackrel{+}{ \stackrel{}{ \coprod}} H \stackrel{\bigcirc{}^{O}}{\stackrel{}{ \stackrel{}{ \longrightarrow}} OMe} \underbrace{\frac{\text{Ti}(O\text{-}i\text{-Pr})_{4}}{\text{Solvent-free}}}_{up \ to \ 98\% \ yield} \stackrel{\bigcirc{}}{ \stackrel{}{ \longleftarrow}} R^{1} \stackrel{\bigcirc{}}{ \stackrel{}{ \longleftarrow}} R^{2}$

aromatic, diaryl, heteroaromatic, condensed ring, conjugated, aliphatic, functionalized ketones

2573 Synthesis of 2,5-Diaminoquinones by One-Pot Copper-Catalyzed Aerobic Oxidation of Hydroquinones and Addition Reaction of Amines

Adv. Synth. Catal. 2009, 351, 2573-2578

Sungjin Kim, Daehwan Kim, Jaiwook Park*

- **2579** An Organocatalytic Synthesis of *cis-N*-Alkyl- and *N*-Arylaziridine Carboxylates

Adv. Synth. Catal. 2009, 351, 2579-2588

- Sean P. Bew,* Rachel Carrington, David L. Hughes, John Liddle, Paolo Pesce
- R³O

 1 10 mol% of a protonated heterocyclic salt

 +

 R²

 1 10 mol% of a protonated heterocyclic salt

 R¹

 24 examples

 aryl or aliphatic substituents

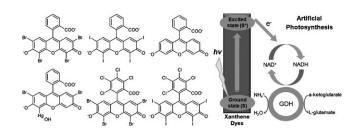
 R²

 or heterocyclic substituents

 ethyl or *tert*-butyl substituents
- 2589 Screening Xanthene Dyes for Visible Light-Driven Nicotinamide Adenine Dinucleotide Regeneration and Photoenzymatic Synthesis

Adv. Synth. Catal. 2009, 351, 2589-2594

Sahng Ha Lee, Dong Heon Nam, Chan Beum Park*



2595

2605

Room Temperature Lewis Base-Catalyzed Alumination of Terminal Alkynes

Adv. Synth. Catal. 2009, 351, 2595-2598

🖳 Yuhan Zhou, Thomas Lecourt, Laurent Micouin*

Iron-Catalyzed Regioselective Hydroaryloxylation of C≡C Triple Bonds: An Efficient Synthesis of 2H-1-Benzopyran Derivatives

Adv. Synth. Catal. 2009, 351, 2599-2604

Xiaobing Xu, Jun Liu, Linfeng Liang, Hongfeng Li, Yanzhong Li*

Silver-Catalyzed Intramolecular Cyclization of o-(1-Alkynyl)benzamides: Efficient Synthesis of (1H)-Isochromen-1-imines

Adv. Synth. Catal. 2009, 351, 2605-2610

Guannan Liu, Yu Zhou, Deju Ye, Dengyou Zhang, Xiao Ding, Hualiang Jiang, Hong Liu*

$$R^{2} \xrightarrow{\text{II}} H \xrightarrow{\text{AgSbF}_{6}} R^{2} \xrightarrow{\text{II}} O \xrightarrow{\text{N}} R^{3}$$

R¹ = aryl, alkyl, heterocyclic;

 $R^2 = H, CH_3;$

R³ = aryl, alkyl, heterocyclic.

Preparation of 2H-1,4-Benzoxazin-2-one Derivatives under Heterogeneous Conditions via Domino Process

2611

Adv. Synth. Catal. 2009, 351, 2611-2614

Roberto Ballini,* Alessandro Palmieri,* Mohammad AbdulKarim Talaq, Serena Gabrielli

Palladium-Catalyzed Annulation of 2-(1-Alkynyl)benzenamines with Disulfides: Synthesis of 3-Sulfenylindoles

Adv. Synth. Catal. 2009, 351, 2615-2618

🖳 Yan-Jin Guo, Ri-Yuan Tang, Jin-Heng Li,* Ping Zhong, Xing-Guo Zhang*

Direct Preparation of 7-Allyl- and 7-Arylindolines

Adv. Synth. Catal. 2009, 351, 2619-2623

Daniele Leonori, Iain Coldham*

2619

2633

FULL PAPERS

2625 Aerobic Oxidation of Benzylic Alcohols in Water by 2,2,6,6-Tetramethylpiperidine-1-oxyl (TEMPO)/Copper(II) 2-N-Arylpyrrolecarbaldimino Complexes

Adv. Synth. Catal. 2009, 351, 2625-2632

Paweł J. Figiel, Ahlam Sibaouih, Jahir Uddin Ahmad, Martin Nieger, Minna T. Räisänen, Markku Leskelä, Timo Repo*

Reusable Copper-Aluminum Hydrotalcite/rac-BINOL System for Room Temperature Selective Aerobic Oxidation

Adv. Synth. Catal. 2009, 351, 2633-2637

alcohols: primary, secondary, alicyclic, heterocyclic alcohols

2638 Metal-Free: An Efficient and Selective Catalytic Aerobic Oxidation of Hydrocarbons with Oxime and *N*-Hydroxyphthalimide

Adv. Synth. Catal. 2009, 351, 2638-2642

Adv. Synth. Catal. 2009, 351, 2643-2649

NHPI, O_{2,} 80 °C

Conversion 82.1%

NOOH

O

Selectivity 94.9%

Gengxiu Zheng,* Chunhong Liu, Qiufen Wang, Mingyu Wang, Guanyu Yang*

2643 Direct Amide Synthesis from Alcohols and Amines by

Phosphine-Free Ruthenium Catalyst Systems

Subhash Chandra Ghosh, Senthilkumar Muthaiah, Yao Zhang, Xiangya Xu, Soon Hyeok Hong*

2650 Magnetic Nanoparticle Supported Second Generation
Hoveyda–Grubbs Catalyst for Metathesis of Unsaturated

Adv. Synth. Catal. 2009, 351, 2650-2656

Fatty Acid Esters

Zhu Yinghuai,* Loo Kuijin, Ng Huimin, Li Chuanzhao, Ludger Paul Stubbs, Chia Fu Siong, Tan Muihua, Ship Chee Peng

2657 Rhodium-Catalysed Coupling of Allylic, Homoallylic, and Bishomoallylic Alcohols with Aldehydes and *N*-Tosylimines: Insights into the Mechanism

Adv. Synth. Catal. 2009, 351, 2657-2666

🖳 Nanna Ahlsten, Belén Martín-Matute*

OH R^2 + R^2 + R

via H O RhL_n

2667

2683

Synthesis of Biaryls and Aryl Ketones *via* Microwave-Assisted Decarboxylative Cross-Couplings

Adv. Synth. Catal. 2009, 351, 2667-2674

Lukas J. Gooßen,* Bettina Zimmermann, Christophe Linder, Nuria Rodríguez, Paul P. Lange, Jens Hartung

Efficient Functionalisation of Cubic Monovinylsilsesquioxanes *via* Cross-Metathesis and Silylative Coupling with Olefins in the Presence of Ruthenium Complexes

Adv. Synth. Catal. 2009, 351, 2675-2682

Patrycja Żak, Cezary Pietraszuk, Bogdan Marciniec,* Grzegorz Spólnik, Witold Danikiewicz

Synthesis of 2,3-Diarylbenzo[b]thiophenes via Nickel-Catalyzed Suzuki-Miyaura Cross-Coupling and Palladium-Catalyzed Decarboxylative Arylation

Adv. Synth. Catal. 2009, 351, 2683-2688

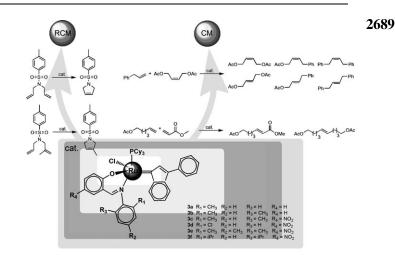
Mitsuru Miyasaka, Koji Hirano, Tetsuya Satoh, Masahiro Miura*

 $R = aryl, CH_2Ph, CH_2SiMe_3, O-t-Bu, O-n-Bu,$

New Indenylidene-Schiff Base-Ruthenium Complexes for Cross-Metathesis and Ring-Closing Metathesis

Adv. Synth. Catal. 2009, 351, 2689-2701

Ana M. Lozano Vila, Stijn Monsaert, Renata Drozdzak, Stanislaw Wolowiec, Francis Verpoort*



Multicatalytic Tandem Reactions of 2-Alkynylbenzaldoximes with Isocyanides

Adv. Synth. Catal. 2009, 351, 2702-2708

Zhiyuan Chen, Xingxin Yu, Mingchao Su, Xiaodi Yang, Jie Wu*

$$R^{1} \stackrel{\text{II}}{\stackrel{\text{II}}}{\stackrel{\text{II}}{\stackrel{\text{II}}}{\stackrel{\text{II}}}{\stackrel{\text{II}}{\stackrel{\text{II}}}{\stackrel{\text{II}}{\stackrel{\text{II}}{\stackrel{\text{II}}{\stackrel{\text{II}}{\stackrel{\text{II}}{\stackrel{\text{II}}}{\stackrel{\text{II}}{\stackrel{\text{II}}}{\stackrel{\text{II}}}{\stackrel{\text{II}}}{\stackrel{\text{II}}}}{\stackrel{\text{II}}}}\stackrel{\text{II}}{\stackrel{\text{II}}}}\stackrel{\text{II}}{\stackrel{\text{II}}{\stackrel{\text{II}}}{\stackrel{\text{II}}{\stackrel{\text{II}}}{\stackrel{\text{II}}}{\stackrel{\text{II}}}}}\stackrel{\text{II}}{\stackrel{\text{II}}}}\stackrel{\text{II}}}}\stackrel{\text{II}}}\stackrel{\text$$

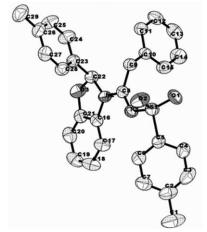
2702

2709 Copper-Catalyzed Efficient Multicomponent Reaction: Synthesis of Benzoxazoline-Amidine Derivatives

Adv. Synth. Catal. 2009, 351, 2709-2713

Yongjia Shang,* Xinwei He, Jinsong Hu, Jianwei Wu, Min Zhang, Shuyan Yu, Qianqian Zhang

$$R^{1}$$
-SO₂N₃
 R^{2}
 $+$
 $+$
 $CH=N$
 HO
 $Cat. Cul$
 TEA
 $THF, 8 h$
 $0-5 °C$
 R^{2}
 $N-SO_{2}R^{2}$



UPDATES

2715 First Copper-Catalyzed Intramolecular Amidation in Substituted 4-Iodopyrazoles Leading to the Synthesis of Pyrazolo[4,3-*b*]-pyridin-5-ones[§]

Adv. Synth. Catal. 2009, 351, 2715-2723

Somnath Nag, Maloy Nayak, Sanjay Batra*

2724 Ruthenium Catalysts for Controlled Mono- and Bis-Allylation of Active Methylene Compounds with Aliphatic Allylic Substrates

Adv. Synth. Catal. 2009, 351, 2724-2728

Hui-Jun Zhang, Bernard Demerseman, Zhenfeng Xi,* Christian Bruneau*

CORRIGENDA

In the full paper by Feng-Quan Li, Shi Zhong, Gui Lu and Albert S. C. Chan in Issue 11+12, 2009, pp. 1955-1960 (DOI: 10.1002/adsc.200900177), the third and fourth sentences in the third paragraph on page 1955 should be corrected as follows:

old: Although several catalytic asymmetric reactions using acylsilanes have been developed, [20,21] to the best of our knowledge, the enantioselective alkynylation of acylsilanes has not been reported so far. [22] Herein we describe for the first time the direct catalytic alkynylation of alkyl acylsilane with both aliphatic and aromatic alkynes.

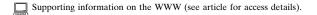
new: Although several catalytic asymmetric reactions using acylsilanes have been developed,^[20,21] few reports were on the enantioselective alkynylation of acylsilanes.^[19,22] For instance, Johnson et al. described a tandem alkynylation of silylglyoxylate using chiral amino alcohol to mediate chirality transfer,^[19a] Scheidt and co-workers described the asymmetric alkyne addition to acylsilane in 74% ee with tridentate Schiff base ligand.^[19b] Herein we describe an effective direct alkynylation of alkyl acylsilane with both aliphatic and aromatic alkynes.

Reference 19 should be as follows:

[19] a) D. A. Nicewicz, J. S. Johnson, *J. Am. Chem. Soc.* **2005**, *127*, 6170–6171; b) T. E. Reynolds, A. R. Bharadwaj, K. A. Scheidt, *J. Am. Chem. Soc.* **2006**, *128*, 15382–15383.

In the communication by Thomas Boddaert, Yoann Coquerel, and Jean Rodriguez in Issue 11+12, 2009, pp. 1744–1748 (DOI: 10.1002/adsc.200900292), there were mistakes in several references. The corrected references are as follows.

- [5a] A. Michaut, T. Boddaert, Y. Coquerel, J. Rodriguez, Synthesis 2007, 2867 2871.
- [5b] T. Boddaert, Y. Coquerel, J. Rodriguez, C. R. Chim. 2009, 12, 872-875.
- [7b] S. Gessler, S. Randl, S. Blechert, *Tetrahedron Lett.* **2000**, *41*, 9973–9976.
- [13f] E. M. Phillips, M. Wadamoto, H. S. Roth, A. W. Ott, K. A. Scheidt, Org. Lett. 2009, 11, 105-108.



*Author to whom correspondence should be addressed.