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*, 11+12, **Pages 1645-1908**

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DEDICATED CLUSTER – COMMENTARY

In Honor of Professor Chi-Huey Wong on the Occasion of His 60^{th} Birthday

Adv. Synth. Catal. 2008, 350, 1659-1660

Michael D. Burkart, William A. Greenberg*

DEDICATED CLUSTER – REVIEW

Metathesis in Peptides and Peptidomimetics

Adv. Synth. Catal. 2008, 350, 1661-1675

Ashraf Brik*

InterScience®

1659

1661

DEDICATED CLUSTER - COMMUNICATIONS

1677 Efficient Synthesis of a Novel Resorcyclide as Anticancer Agent Based on Hsp90 Inhibition

Adv. Synth. Catal. 2008, 350, 1677-1681

☐ Xiaoguang Lei, Samuel J. Danishefsky*

- Me HO Me H Click Me O OH HO Me H TriazoleCI OMOM
- 1682 Combined Chemical-Enzymatic Assembly of Aminoglycoside Derivatives with N-1-AHB Side Chain

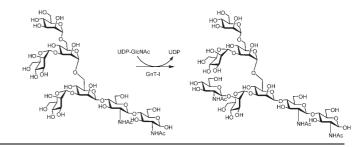
Adv. Synth. Catal. 2008, 350, 1682-1688

☐ Igor Nudelman, Lilach Chen, Nicholas M. Llewellyn, El-Habib Sahraoui, Marina Cherniavsky, Jonathan B. Spencer,* Timor Baasov*

- 1689 Enzyme-Catalyzed Synthesis of a Hybrid N-Linked Oligosaccharide using N-Acetylglucosaminyltransferase I

Adv. Synth. Catal. 2008, 350, 1689-1695

Rui Chen, Mark A. Pawlicki, Brian S. Hamilton, Thomas J. Tolbert*



1696 Bromodimethylsulfonium Bromide-Silver Triflate: A New Powerful Promoter System for the Activation of Thioglycosides

Adv. Synth. Catal. 2008, 350, 1696-1700

De-Cai Xiong, Li-He Zhang, Xin-Shan Ye*

PO
$$R^2 = Et$$
, Tol SR^1 BDMS, AgOTf, CH_2Cl_2 $R^2 = OMe$, $STol$ R^2 armed and disarmed

thioglycosides 19 examples

DEDICATED CLUSTER - FULL PAPERS

1701 Studies toward the Synthesis of (–)-Zampanolide: Preparation of the Macrocyclic Core

Adv. Synth. Catal. 2008, 350, 1701-1711

Dawn M. Troast, Jiayi Yuan, John A. Porco, Jr.*

- **1712** Chemoenzymatic Approach to Enantiomerically Pure (*R*)-3-Hydroxy-3-methyl-4-pentenoic Acid Ester and Its Application to a Formal Total Synthesis of Taurospongin A

Adv. Synth. Catal. 2008, 350, 1712-1716

🖳 Aya Fujino, Takeshi Sugai*

BnO 1) BSEH BnO OH (R)

HO

1) R. rhodochrous
NBRC 15564
2) PMBCI

PMBO₂C HO (R), 100% ee

1717

1736

1761

1771

251-membered

compound librar

 $Nu = C_{-}, N_{-}, O_{-}, S_{-}$ nucleophiles

Chemoenzymatic Syntheses of Tumor-Associated Carbohydrate Antigen Globo-H and Stage-Specific Embryonic Antigen 4

Adv. Synth. Catal. 2008, 350, 1717-1728

Zhen Wang, Michel Gilbert, Hironobu Eguchi, Hai Yu, Jiansong Cheng, Saddam Muthana, Luyuan Zhou, Peng George Wang, Xi Chen, Xuefei Huang*

One-Pot, Regioselective Synthesis of Substituted Arylglycines for Kinetic Resolution by Penicillin G Acylase

Adv. Synth. Catal. 2008, 350, 1729-1735

Peter Grundmann, Wolf-Dieter Fessner*

Solid-Phase Synthesis of [5.5]-Spiroketals

Adv. Synth. Catal. 2008, 350, 1736-1750

Stefan Sommer, Marc Kühn, Herbert Waldmann*

Large-Scale Synthesis of New Pyranoid Building Blocks Based on Aldolase-Catalysed Carbon-Carbon Bond Formation

Adv. Synth. Catal. 2008, 350, 1751-1759

Michael Wolberg,* Ben H. N. Dassen, Martin Schürmann, Stefan Jennewein, Marcel G. Wubbolts, Hans E. Schoemaker, Daniel Mink*

DEDICATED CLUSTER – UPDATE

Optimization of Biocatalyst Specific Activity for Glycolic Acid Production

$$\begin{array}{c} \text{HCHO} + \text{HCN} \xrightarrow{\hspace{1cm} > 99 \%} & \begin{array}{c} \text{HO} \\ \text{H_2C-C=N} \end{array} & \begin{array}{c} \text{nitrilase} \\ \text{> 99 \%} \end{array} & \begin{array}{c} \text{HO} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{IEX} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{HO} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{IEX} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{HO} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{IEX} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{HO} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{IEX} \\ \text{H_2C-C-O-NH_4}^+ \end{array} & \begin{array}{c} \text{HO} \\ \text{H_2C-C-O-NH_4$$

Adv. Synth. Catal. 2008, 350, 1761 – 1769

Arie Ben-Bassat, Alison M. Walls, Matthew A. Plummer, Amy E. Sigmund, William L. Spillan, Robert DiCosimo*

COMMUNICATIONS

Highly Efficient Heterogeneous Aqueous Kharasch Reaction with an Amphiphilic Resin-Supported Ruthenium Catalyst

Adv. Synth. Catal. 2008, 350, 1771-1775

Yohei Oe, Yasuhiro Uozumi*

S/C = 2,000, in water; TOF = 1445 h^{-1} (at initial 15 min) S/C = 100, in toluene; TOF < 1 (no reaction)

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1649

1776 Chiral Lithium Salts of Phosphoric Acids as Lewis Acid–Base Conjugate Catalysts for the Enantioselective Cyanosilylation of Ketones

Adv. Synth. Catal. 2008, 350, 1776-1780

- Manabu Hatano, Takumi Ikeno, Tokihiko Matsumura, Shinobu Torii, Kazuaki Ishihara*
- **1781** Enantioselective Michael Addition of Dicyanoolefins to α,β-Unsaturated Aldehydes in Aqueous Medium

Adv. Synth. Catal. 2008, 350, 1781-1784

☐ Jun Lu, Feng Liu, Teck-Peng Loh*

- **1785** A Novel Bis-Thiourea Organocatalyst for the Asymmetric Aza-Henry Reaction

Adv. Synth. Catal. 2008, 350, 1785-1790

Constantinos Rampalakos, William D. Wulff*

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

1791 A Novel Pyridine-Bridged Bis-benzimidazolylidene Pincer Palladium Complex: Synthesis and Catalytic Properties

Adv. Synth. Catal. 2008, 350, 1791-1795

Tao Tu,* Jagadeesh Malineni, Karl Heinz Dötz*

TON: up to 2.48 $\times 10^8$

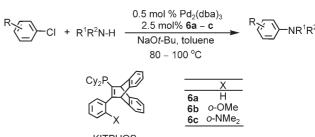
1796 Organocatalytic and Enantioselective Direct Vinylogous Michael Addition to Maleimides

Adv. Synth. Catal. 2008, 350, 1796-1800

- Jun Lu, Wei-Juan Zhou, Feng Liu, Teck-Peng Loh*
- NC CN O Cinchona alkaloids A R = Bn An R = 2-Cr₁₀H₇CH₂OH
- 1801 Electron-Rich, Bicyclic Biaryl-Like KITPHOS
 Monophosphines *via* [4+2] Cycloaddition between
 1-Alkynylphosphine Oxides and Anthracene: Highly
 Efficient Ligands for Palladium-Catalysed C-N and C-C
 Bond Formation

Adv. Synth. Catal. 2008, 350, 1801-1806

Simon Doherty,* Julian G. Knight,* Catherine H. Smyth,* Graeme A. Jorgenson



KITPHOS

1807

1816

1823

1830

1835

1841

FULL PAPERS

Aerobic Oxidations Catalyzed by Zeolite-Encapsulated Cobalt Salophen

Adv. Synth. Catal. 2008, 350, 1807-1815

Mikael Johansson, Byron W. Purse, Osamu Terasaki, Jan-E. Bäckvall*

Enhanced Cooperativity in Hydrolytic Kinetic Resolution of Epoxides using Poly(styrene) Resin-Supported Dendronized Co-(Salen) Catalysts

Adv. Synth. Catal. 2008, 350, 1816-1822

Poorva Goyal, Xiaolai Zheng, Marcus Weck*

(rac)-epoxide 0.7 eq
$$(0.04 \text{ mol}\%)$$

Highly Efficient and Stereoselective Julia–Kocienski Protocol for the Synthesis of α -Fluoro- α , β -unsaturated Esters and Weinreb Amides Employing 3,5-Bis(trifluoromethyl)phenyl (BTFP) Sulfones

Adv. Synth. Catal. 2008, 350, 1823-1829

Diego A. Alonso,* Mónica Fuensanta, Enrique Gómez-Bengoa, Carmen Nájera* F₃C CF_3 CF_3

Copper-Catalyzed Cycloaddition of Sulfonyl Azides with Alkynes to Synthesize *N*-Sulfonyltriazoles 'on Water' at Room Temperature

Adv. Synth. Catal. 2008, 350, 1830-1834

Feng Wang, Hua Fu,* Yuyang Jiang, Yufen Zhao

Iron-Catalyzed Aziridination Reactions

Adv. Synth. Catal. 2008, 350, 1835-1840

Masafumi Nakanishi, Anne-Frederique Salit, Carsten Bolm*

for R = OSiMe₃

$$R'''$$

Fe(OTf)₂ (cat.), Arl=NX
(preformed or in situ generated)

for R = H or Me
(also asymmetric)

 R'''
 R''''
 R''''

Palladium-Catalyzed Trimerizations of Terminal Arylalkynes: Synthesis of 1,3-Diaryl-2-arylethynyl-1,3butadienes

Adv. Synth. Catal. 2008, 350, 1841-1849

Yao-Ting Wu,* Wei-Chih Lin, Chia-Ju Liu, Chuan-Yi Wu

1850 Access to Functionalized Isoquinoline *N*-Oxides *via* Sequential Electrophilic Cyclization/Cross-Coupling Reactions

- Qiuping Ding, Jie Wu*
- 1855 New Functionalized Vinyl Monomers by Ethylene-(Functionalized)-Norbornene Hetero-Trimerization Catalyzed by Cobalt(II)-(Imino)pyridine Complexes

Adv. Synth. Catal. 2008, 350, 1855-1866

Alessandro Toti, Giuliano Giambastiani,* Claudio Bianchini,* Andrea Meli, Lapo Luconi

R = H, CH_2OH , COOH; R' = i-Pr, Me Ar = Ph, 2-thienyl, 2-benzothienyl, 2-furanyl

1867 Iminium Salt-Catalysed Asymmetric Epoxidation using Hydrogen Peroxide as Stoichiometric Oxidant

Adv. Synth. Catal. 2008, 350, 1867-1874

Philip C. Bulman Page,* Phillip Parker, Gerasimos A. Rassias, Benjamin R. Buckley, Donald Bethell

1875 Organocatalytic Asymmetric Hydrophosphination of α,β-Unsaturated Aldehydes: Development, Mechanism and DFT Calculations

Adv. Synth. Catal. 2008, 350, 1875-1884

☐ Ismail Ibrahem, Peter Hammar, Jan Vesely, Ramon Rios, Lars Eriksson, Armando Córdova*

1885 Stereoselective Synthesis of γ-Butyrolactones *via* Organocatalytic Annulations of Enals and Keto Esters

Adv. Synth. Catal. 2008, 350, 1885-1890

☐ Yi Li, Zhuo-An Zhao, Hu He, Shu-Li You*

1891 Domino Catalysis in the Direct Conversion of Carboxylic Acids to Esters

Adv. Synth. Catal. 2008, 350, 1891-1900

☐ I. Held, P. von den Hoff, D. S. Stephenson, H. Zipse*

Catalytic Deprotonative Functionalization of Propargyl Silyl Ethers with Imines

OSiMe₃ +
$$R^4$$
 + R^2 H cat. t -Bu-P₄ base toluene, r.t., 3 h

Adv. Synth. Catal. 2008, 350, 1901-1906

Hiroshi Naka,* Daiki Koseki, Yoshinori Kondo*

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

stAuthor to whom correspondence should be addressed.