

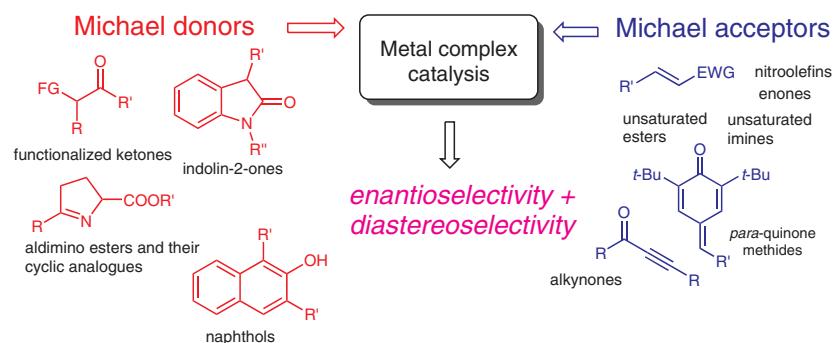
Synthesis

Synthesis 2020, 52, 781–795
DOI: 10.1055/s-0039-1690044

A. N. Reznikov*
Y. N. Klimochkin
Samara State Technical
University, Russian Federation

Recent Developments in Highly Stereoselective Michael Addition Reactions Catalyzed by Metal Complexes

Review
781



Synthesis

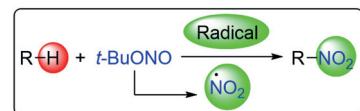
Synthesis 2020, 52, 796–806
DOI: 10.1055/s-0039-1690789

S.-Z. Song
Y. Dong*
G.-P. Ge*
Q. Li
W.-T. Wei*
Ningbo University, P. R. of China

Recent Advances in Radical Nitration Using *tert*-Butyl Nitrite

Short Review

796



Synthesis

Synthesis 2020, 52, 807–818
DOI: 10.1055/s-0039-1690046

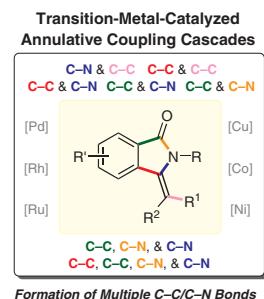
S. W. Youn*

Hanyang University,
Republic of Korea

Transition-Metal-Catalyzed Annulative Coupling Cascade for the Synthesis of 3-Methyleneisoindolin-1-ones

Short Review

807

**Synthesis**

Synthesis 2020, 52, 819–833
DOI: 10.1055/s-0039-1691561

J.-C. Hsieh*

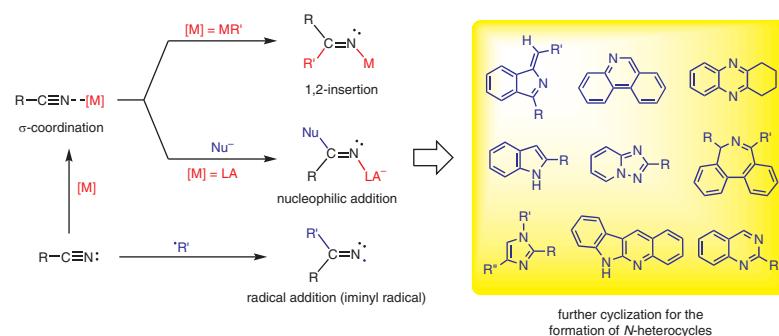
H.-L. Su

Tamkang University, Taiwan

Synthesis of *N*-Heterocycles via Transition-Metal-Catalyzed Tandem Addition/Cyclization of a Nitrile

Short Review

819

**Synthesis**

Alkyl Tosylates as Alkylation Reagents in the Catellani Reaction

Feature

834

Synthesis 2020, 52, 834–846
DOI: 10.1055/s-0039-1690801

Q. Gao

Z.-S. Liu

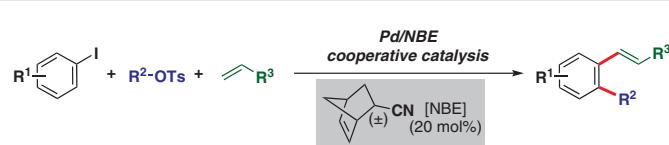
Y. Hua

S. Zhou

H.-G. Cheng*

Q. Zhou*

Wuhan University, P. R. of China



- NBE-CN as the catalytic mediator
- easily accessible starting materials

- broad substrate scope
- scalable protocol

45 examples
up to 97% yield

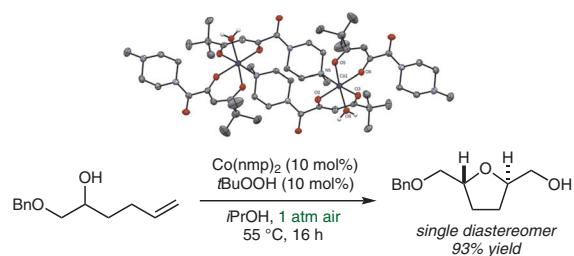
Synthesis

Synthesis 2020, 52, 847–852
DOI: 10.1055/s-0039-1690730

B. Morra
N. A. Morra
D. G. MacDonald
B. L. Pagenkopf*
The University of Western Ontario, Canada

Gram-Scale Synthesis of the Co(nmp)₂ Catalyst to Prepare *trans*-2,5-Disubstituted Tetrahydrofurans by the Aerobic Oxidative Cyclization of Pent-4-en-1-ols

PSP
847

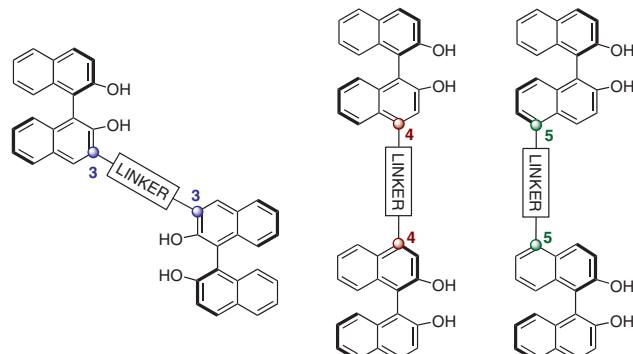
**Synthesis**

Synthesis 2020, 52, 853–860
DOI: 10.1055/s-0039-1690763

M. Kohlhaas
F. Lutz
N. Paransothy
F. Octa-Smolín
C. Wölper
J. Niemeyer*
University of Duisburg-Essen, Germany

Synthesis of Bis-BINOL Derivatives: Linking via the 3-, 4-, or 5-Position by Generation of Suitable C₁-Symmetric Precursors

Paper
853

**Synthesis**

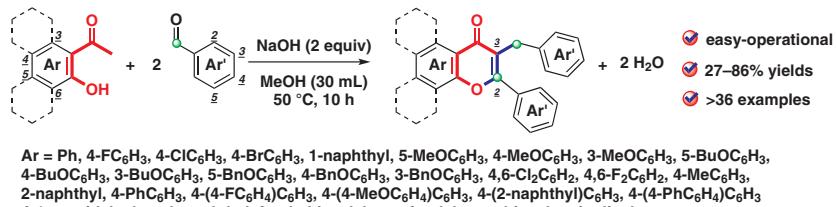
Synthesis 2020, 52, 861–872
DOI: 10.1055/s-0039-1690760

M.-Y. Chang*
K.-T. Chen
Y.-L. Tsai
H.-Y. Chen
Kaohsiung Medical University, Taiwan
Kaohsiung Medical University Hospital, Taiwan

One-Pot Access to 2-Aryl-3-(aryl methyl)chromones

Paper

861



Synthesis

Synthesis 2020, 52, 873–881
DOI: 10.1055/s-0039-1690766

J. Skotnitzki

A. Kremsmair

B. Kicin

R. Saeb

V. Ruf

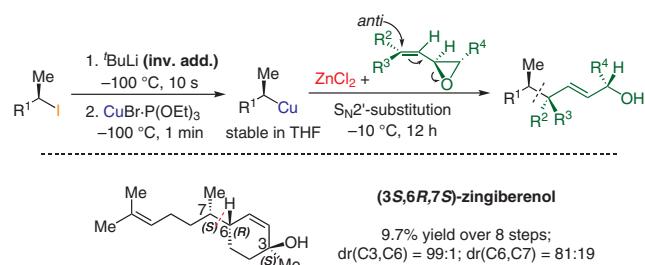
P. Knochel*

Ludwig-Maximilians-Universität,
Germany

Stereoselective *anti*- S_N2' -Substitutions of Secondary Alkylcopper-Zinc Reagents with Allylic Epoxides: Total Synthesis of (3*S*,6*R*,7*S*)-Zingiberenol

Paper

873

**Synthesis**

Synthesis 2020, 52, 882–892
DOI: 10.1055/s-0039-1691487

L. Pruschinski

A.-L. Lücke

T. Freese

S.-R. Kahnert

S. Mummel

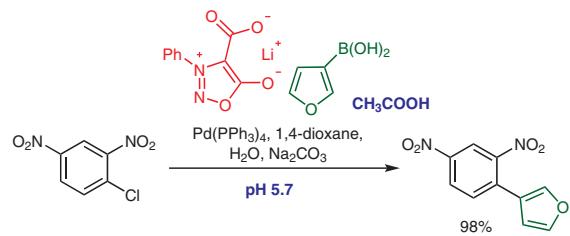
A. Schmidt*

Clausthal University of Technology,
Germany

Suzuki–Miyaura Cross-Couplings under Acidic Conditions

Paper

882

**Synthesis**

Synthesis 2020, 52, 893–900
DOI: 10.1055/s-0039-1690765

H.-L. Lu

F.-H. Guo

T.-L. Wang

X.-C. Wang

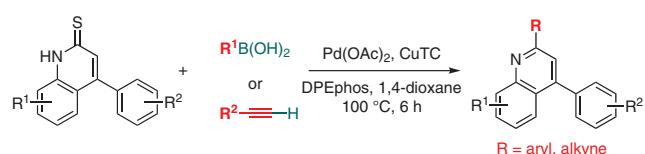
Z.-J. Quan*

Northwest Normal University,
P. R. of China
Gansu International Scientific
and Technological Cooperation
Base of Water-Retention Chemical
Functional Materials,
P. R. of China

Palladium-Catalyzed/Copper-Mediated Desulfurization and Arylation of Quinoline-2-(1*H*)-thione for Rapid Access to Quinoline Derivatives

Paper

893



● C–C bonds formation
● 29 examples, up to 85% yield
● without an inert atmosphere
● base-free

Synthesis

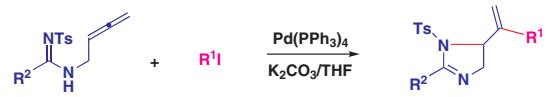
Synthesis 2020, 52, 901–908
DOI: 10.1055/s-0037-1610742

Synthesis of 2-Imidazolines via Palladium-Catalyzed Cyclization Reaction of 2,3-Allenyl Amines and Aryl Iodides**Paper**

901

Y. Liu
C. Zhang
X. Liang
X. Zeng
R. Lu
Z. Fang
S. Wang
Y. Liu
J. Hu*

Weifang Medical University,
P. R. of China



19 examples: 48–78%

* Low toxicity * form C–C and C–N * Readily available substrates

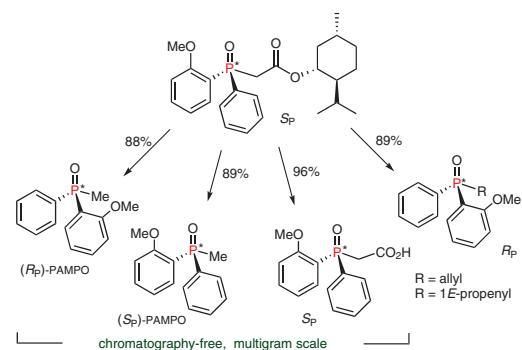
Synthesis

Synthesis 2020, 52, 909–916
DOI: 10.1055/s-0039-1691531

Enantiodivergent Synthesis of Both PAMPO Enantiomers Using L-Menthyl Chloroacetate and Stereomutation at P in Classical Quaternisation Reactions**Paper**

909

K. Dziuba
M. Łubańska
K. M. Pietrusiewicz*
Maria Curie-Skłodowska University, Poland



chromatography-free, multigram scale

Synthesis

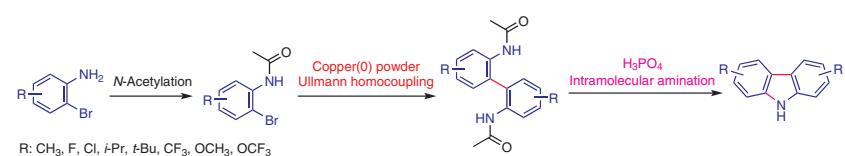
Synthesis 2020, 52, 917–927
DOI: 10.1055/s-0039-1690759

Site-Specific Synthesis of Carbazole Derivatives through Aryl Homocoupling and Amination**Paper**

917

J. Ban
M. Lim
S. Shabbir
J. Baek
H. Rhee*

Hanyang University,
Republic of Korea



R: CH3, F, Cl, i-Pr, t-Bu, CF3, OCH3, OCF3

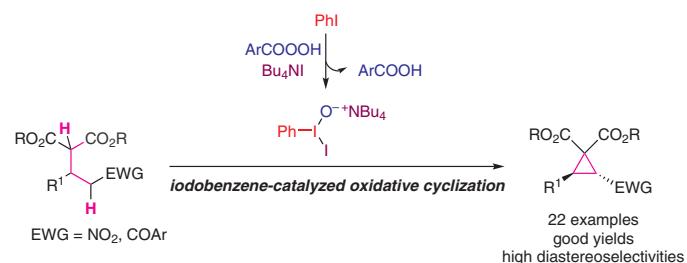
Synthesis

Synthesis 2020, 52, 928–932
DOI: 10.1055/s-0039-1690809

Y. Li
H. Guo*
R. Fan*
Fudan University, P. R. of China

Iodobenzene-Catalyzed Oxidative Cyclization for the Synthesis of Highly Functionalized Cyclopropanes**Paper**

928

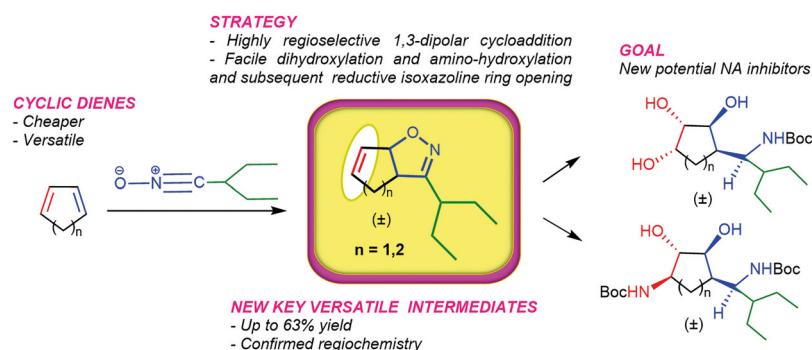
**Synthesis**

Synthesis 2020, 52, 933–941
DOI: 10.1055/s-0039-1690039

R. Bartolotta
C. La Rosa*
D. Nava
Università degli Studi di Milano,
Italy

New Strategy of Synthesis of Peramivir Analogues as Potential Neuraminidase Inhibitors**Paper**

933

**Synthesis**

Synthesis 2020, 52, 942–948
DOI: 10.1055/s-0039-1691522

K. Ohsawa
H. Zhao
T. Tokunaga
C. Thomas
A. Ganeshan
Y. Masuda
T. Doi*
Tohoku University, Japan

Stereoselective Synthesis of Protected L-*allo*-Enduracididine and L-Enduracididine via Asymmetric Nitroaldol Reaction**Paper**

942

