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TMSN₃

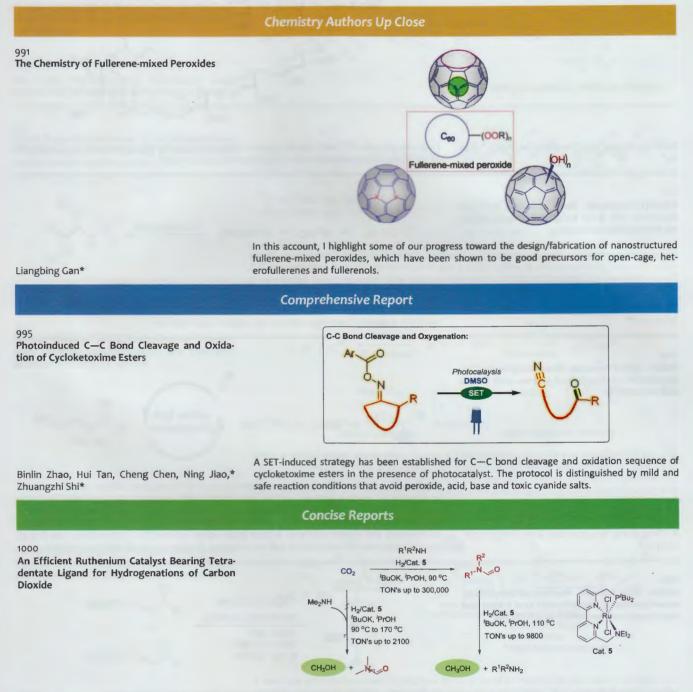
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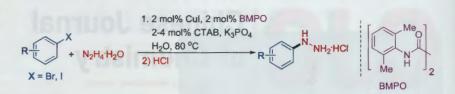
Feng-Hua Zhang, Chong Liu, Wei Li, Gui-Long Tian, Jian-Hua Xie, Qi-Lin Zhou* A ruthenium catalyst with tetradentate bipyridine ligand gave TONs up to 300 000, 9800, and 2100 for the hydrogenations of CO_2 to formamides, formamides to methanol and amines, and the direct hydrogenation of CO_2 to methanol, respectively.



Content

1003

Synthesis of Aryl Hydrazines via Cul/BMPO Catalyzed Cross-Coupling of Aryl Halides with Hydrazine Hydrate in Water

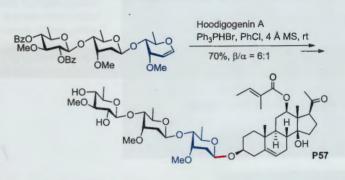


Siripuram Vijay Kumar, Dawei Ma

A practical method for preparing aryl hydrazines from aryl halides is developed.

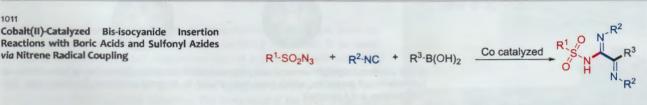
1007

A Glycal Approach to the Synthesis of Pregnane **Glycoside P57**



Chao Liu, Yuyong Ma, Chengfeng Pei, Wei Li,* Biao Yu*

Pregnane glycoside P57 with potent appetite suppressant activity was synthesized via a β -selective TPHB-promoted glycosylation with a trisaccharide glycal as donor.



Zheng-Yang Gu, Rong Zhang, Shun-Yi Wang,* Shun-Jun Ji⁴

A Co(II)-catalyzed effective synthesis of amidinium imine derivatives with isocyanides and boric acids using organic azides as nitrene source has been developed. This protocol provides a new, environment-friendly and simple strategy to effective synthesis of the amidinium imine derivatives with a range of substrates without any oxidants and additives.

1017

1011

Visible Light-Promoted Three-Component Carboazidation of Unactivated Alkenes with TMSN₃ and Acrylonitrile



Visible light-promoted three-component carboazidation of unactivated alkenes using TMSN₃ and acrylonitrile as reaction partners without any stoichiometric peroxides was reported. The intramolecular reaction of 1,6-diene could also undergo smoothly to afford the cyclization product. The obtained δ-azido alkylnitriles could be readily converted into valuable building blocks for medicinal chemistry and organic synthesis. A facile azido radical initiated [3+2] cycloaddition reaction of vinylcyclopropane with acrylonitrile was also observed to deliver a substituted cyclopentane.

FSO₂CF₂CO₂Me

DMAc or DMF rt, N₂, 12-30 h

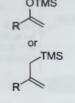
fac-lr(ppy)3 (3 mol%) visible light

1024

Zhan Lu*

Visible Light-Induced Methoxycarbonyldifluoromethylation of Trimethylsilyl Enol Ethers and Allyltrimethylsilanes with FSO₂CF₂CO₂Me

Bo Yang, Xiang Ren, Xuzhong Shen, Tongtong Li,

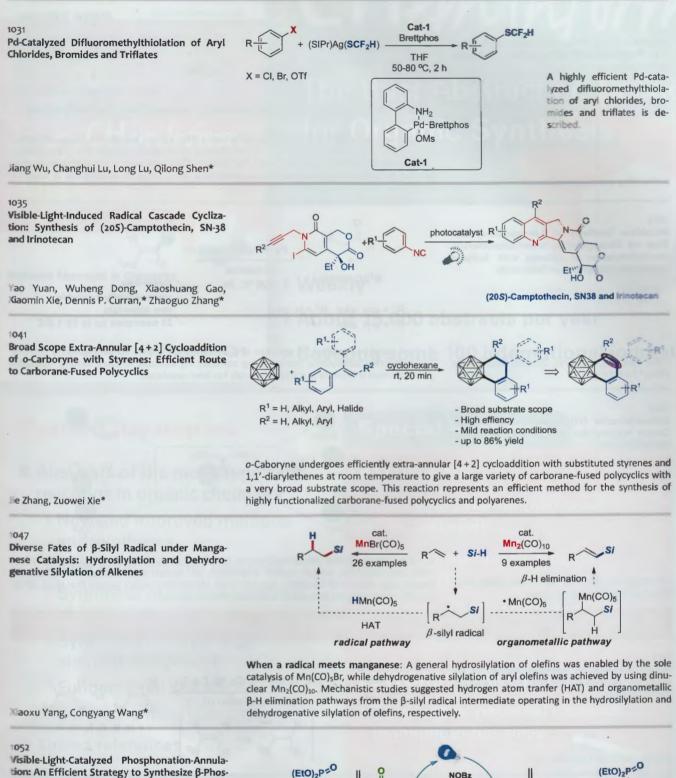


F2CO2Me

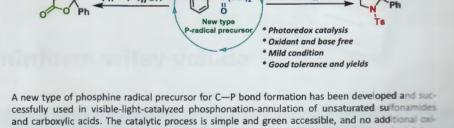
A photoredox catalytic method, using FSO2CF2CO2Me as a CF2CO2Me radical source, has been developed for the conversion of silvl enol ethers and allylsilanes to CF2CO2Me substituted ketones and allylic compounds, respectively.

Wei Yu, Yao Ouyang, Xiu-Hua Xu, Feng-Ling Qing*

Chin. J. Chem.



tion: An Efficient Strategy to Synthesize β-Phosphonopyrrolidines and β-Phosphonolactones



dant or base is needed. Mechanistic studies suggest that the reaction proceeds via a single even

P(OEt)

Chong Li, Zhi-Chao Qi, Qiang Yang, Xiao-Yue Qiang, Shang-Dong Yang*

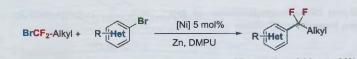
tron transfer pathway.

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Content

1059

Nickel-Catalyzed Difluoroalkylation of (Hetero)aryl Bromides with Unactivated 1-Bromo-1,1difluoroalkanes



27 examples, yield up to 88%

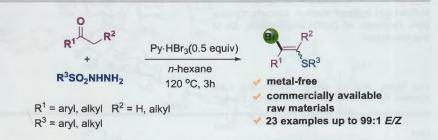
*Broad substrate scope *Excellent functional group compatibility *Synthetic simplicity

A title reaction has been developed (see the Scheme). The reaction enables the difluoroalkylations of a variety of (hetero)arylbromides with unactivated difluoroalkylbromides with high efficiency and excellent functional group tolerance. The feature of this approach is the synthetic simplicity without preformation of arylmetals, thus providing a facile route for applications in medicinal chemistry.

Xu He, Xing Gao, Xingang Zhang*

1063

Metal-Free Synthesis of β-Bromoalkenyl Sulfides via Deoxygenative Bromination/Olefination/Sulfenylation of Ketones with Sulfonyl Hydrazides and Pyridinium Tribromide

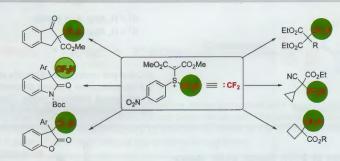


Yishu Bao, Lingyu Zhong, Qiaodan Hou, Qingfa Zhou,* Fulai Yang* tion/olefination/sulfenylation proand pyridinium tribromide as star

1069

Carbon-Selective Difluoromethylation of Soft Carbon Nucleophiles with Difluoromethylated Sulfonium Ylide





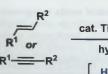
Jiansheng Zhu, Hanliang Zheng, Xiao-Song Xue,* fonil Yisa Xiao, Yafei Liu, Qilong Shen* rom

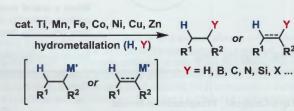
A highly carbon-selective difluoromethylation of soft carbon nucleophiles including β -ketoesters, malonates, oxindoles, benzofuranones and ketene silyl acetals with a difluoromethylated sulfonium ylide under mild conditions was described. Mechanistic studies suggest that these difluoromethylating reactions proceed *via* a difluorocarbene pathway.

Critical Review



Recent Advances in Hydrometallation of Alkenes and Alkynes via the First Row Transition Metal Catalysis





Jianhui Chen, Jun Guo, Zhan Lu*