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α-Difluoroalkylation of Benzyl Amines with Trifluoromethylarenes

Wen-Jun Yue and Ruben Martin

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37 examples yield up to 93%

Chemoselective Three-Component Geminal Cross Couplings of Dihaloalkanes with Cr Catalysis: Rapid Access to Tertiary and Quaternary Alkanes via a Metal–Carbene Intermediate

Sha Wang, Linhong Long, Xiaoyu Zhang, Liang Ling, Hui Chen, and Xiaoming Zeng

Angew. Chem. Int. Ed. 2023, 62, e202312856 https://doi.org/10.1002/anie.202312856



53 examples yield up to 91%

Overcoming a Radical Polarity Mismatch in Strain-Release Pentafluorosulfanylation of [1.1.0]Bicyclobutanes: An Entryway to Sulfone- and Carbonyl-Containing SF5-Cyclobutanes

Yannick Kraemer, Jón Atiba Buldt, Wang-Yeuk Kong, Alexander M. Stephens, Abbey N. Ragan, Soojun Park, Zane C. Haidar, Ansh Hiten Patel, Rachel Shey, Roee Dagan, Connor P. McLoughlin, James C. Fettinger, Dean J. Tantillo, Cody Ross Pitts

Angew. Chem. Int. Ed. 2024, 63, e202319930 https://doi.org/10.1002/anie.202319930



Controllable Double Difluoromethylene Insertions into S-Cu Bonds: (Arylthio)tetrafluoroethylation of Aryl lodides with TMSCF₂B

Shitao Pan, Qiqiang Xie, Xiu Wang, Rumin Huang, Yuhao Lu, Chuanfa Ni, and Jinbo Hu*

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43 examples yield up to 96%

Synthetic Advantages of Defluorinative C-F Bond Functionalization (Rewiew)

Leidy V. Hooker and Jeffrey S. Bandar

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Much progress has been made in the development of methods to both create compounds that contain C-F bonds and to functionalize C-F bonds. As such, C-F bonds are becoming common and versatile synthetic functional handles. This review summarizes the advantages of defluorinative functionalization reactions for small molecule synthesis. The coverage is organized by the type of carbon framework the fluorine is attached to for monoand polyfluorinated motifs. The main challenges, opportunities and advances of defluorinative functionalization are discussed for each class of organofluorine. Most of the text focuses on case studies that illustrate how defluorofunctionalization can improve routes to synthetic targets or how the properties of C F bonds enable unique mechanisms and reactions. The broader goal is to showcase the opportunities for incorporating and exploiting C F bonds in the design of synthetic routes, improvement of specific reactions and advance of new methods.

Journal of the American Chemical Society

Copper-Mediated Cyanodifluoromethylation of (Hetero)aryl lodides and Activated (Hetero)aryl Bromides with TMSCF₂CN

Jeremy Nicolai, Tommaso Fantoni, Trevor W. Butcher, Sophie I. Arlow, Serhiy V. Ryabukhin, Dmytro M. Volochnyuk, John F. Hartwig

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