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Graphical Abstracts/J. Fluorine Chem. 190 (2016) iv-vii

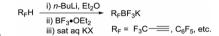
Preparation of potassium fluoroorganotrifluoroborates under non-etching conditions *via* a Li-K exchange

P. Veeraraghavan Ramachandran, Wataru Mitsuhashi

Herbert C. Brown Center for Borane Research, Department of Chemistry, Purdue University, 560 Oval Drive, West Lafayette, IN, 47907, USA

- A non-etching preparation of potassium trifluoropropynyl- and fluoroaryltrifluoroborates.
- BF_3OEt_2 served as the fluoroboron source. Aqueous saturated potassium salt solutions, other than KHF_2 , were used as the potassium source. Facile Li-K exchange generates potassium organotrifluoroborates.

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Carbon nanodots as fluorescent platforms for recognition of fluoride ion via the inner filter effect of simple arylboronic acids. Experimental and theoretical investigations

Zahra Mohammadpour^a, Afsaneh Safavi^a, Akbar Omidvar^a, Afshan Mohajeri^a, Nabiollah Mobaraki^a, Mojtaba Shamsipur^b

^aDepartment of Chemistry, Faculty of Science, Shiraz University, Shiraz, 7194684795, Iran ^bDepartment of Chemistry, Razi University, Kermanshah, Iran

• Design of a simple and selective water soluble nanosensor for assay of fluoride ion. • Use of carbon nanodots as fluorescent sensors for F[−] ion via IFE of arylboronic acids. • Experimental and computational study of substituents on anythoronic acids on consequence. • Theoretical

f

A new strategy designed for flooride sensing in superox media through the inner
filter effect of replement sold on floorection modes.

All prediction of effect of fluoridation reaction

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and computational study of substituents on arylboronic acids on sensor response. • Theoretical prediction of effect of fluoridation reaction on λ_{ex} of carbon dots via IFE. • Fulfillment of EPA requirement for water safety by the minimum detectable value for F^- ion.

Radical reactivity of α,β,β -trifluoroacrylic ester: Facile approach to γ -silyloxy- α,β,β -trifluorobutyric acid derivatives involving regioselective nucleophilic radical addition reaction

Ken Tamamoto, Shigeyuki Yamada, Masao Higashi, Tsutomu Konno, Takashi Ishihara Faculty of Molecular Chemistry and Engineering, Kyoto Institute of Technology, Matsugasaki, Sakyo-ku, Kyoto 606-8585, Japan

● A radical addition of cyclic ethers towards trifluoroacrylate proceeded smoothly. ● We succeeded in the radical coupling of trifluoroacrylate with carbonyl compounds. ● A variety of aldehydes and ketones could be successfully applied for the reaction.

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Convergent synthesis and cytotoxicity of novel trifluoromethylsubstituted (1*H*-pyrazol-1-yl)(quinolin-4-yl) methanones

Helio G. Bonacorso^a, Pablo A. Nogara^a, Fernanda D'A. Silva^b, Wilian C. Rosa^a, Carson W. Wiethan^a, Nilo Zanatta^a, Marcos A.P. Martins^a, João B.T. Rocha^b

^aNúcleo de Química de Heterociclos (NUQUIMHE), Departamento de Química, Universidade Federal de Santa Maria. 97105-900 Santa Maria. Brazil

bLaboratório de Bioquímica Toxicológica, Departamento de Bioquímica e Biologia Molecular, Universidade Federal de Santa Maria, 97105-900 Santa Maria, Brazil

• A convergent synthesis of new trifluoromethylated (1*H*-pyrazol-1-yl)(quinolin-4-yl) methanones (16 examples), is presented. • High regiosectective [3+2] cyclocondensation reactions are successful performed to obtain pyrazoles under green conditions. • Significant cytotoxicity in human leukocytes only at high concentrations for (di-heteroaryl) methanones, is observed. • Most of new (1*H*-pyrazol-1-yl) (quinolin-4-yl) methanones were in agreement with Lipinski's rule of five.

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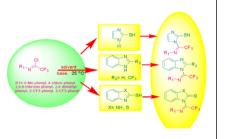
Synthesis of new α -trifluoromethyl substituted formamidines framework by using N-nucleophiles and N,S bidentate nucleophiles

Ali Darehkordi, Fariba Rahmani

Department of Chemistry, Faculty of Science, Vali-e-Asr University of Rafsanjan, Rafsanjan 77176, Iran

- New trifluoromethylated formamidines framework was prepared without using any catalyst. Simple procedure and purification. Catalyst free reactions with excellent yields.
- New bioactive trifluoromethylated benzimidazoles, imidazoles and benzothiazoles.

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Study on thermal decomposition properties and its decomposition mechanism of pentafluoroethane (HFC-125) fire extinguishing agent

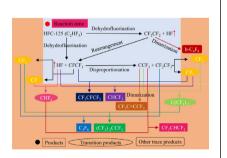
Wang Tinga,b, Hu Ying-jiea,c, Zhang Pina, Pan Ren-minga

^aNanjing University of Science and Technology, Nanjing, 210094, Jiangsu, China ^bJiangsu Police Institute, Nanjing, 210031, Jiangsu, China

^cNanjing Xiaozhuang University, 211171, Nanjing, Jiangsu, China

- A combination of two-approach study for thermal decomposition of pentafluoroethane.
- A possible decomposition mechanism is proposed. The mechanism relies on stabilization changes of reaction energies and barriers. Strictly confirm gas products with different pyrolysis temperature.

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Strong influence of the trifluoromethyl group on the chemoselectivity of [3+2]-cycloadditions of thiocarbonyl S-methanides with α,β -unsaturated ketones

Grzegorz Mlostoń^a, Paulina Grzelak^a, Heinz Heimgartner^b

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ullet Selected fluorinated enones were reacted with thiocarbonyl S-methanides. ullet Type of the obtained [3+2]-cycloadduct depends on the location of the CF $_3$ unit. ullet Tetrahydrothiophene derivatives were obtained from the CF $_3$ CH=CH enones. ullet 1,3-Oxathiole derivatives were obtained from the CF $_3$ C=O enones.

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