

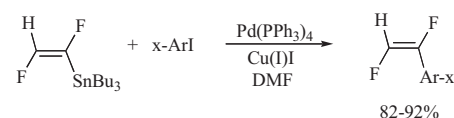
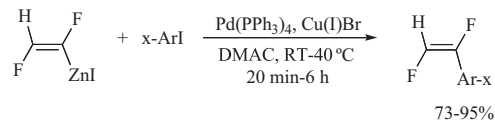
Graphical Abstracts/J. Fluorine Chem. 132 (2011) 75–77

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The stereoselective synthesis of (*Z*)-HFC=CFZnI and stereospecific preparation of (*E*)-1,2-difluorostyrenes from (*Z*)-HFC=CFZnI via an unusual Pd(PPh₃)₄-Cu(I)Br co-catalysis approach or (*Z*)-HFC=CFSnBu₃

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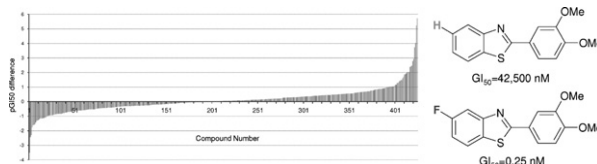


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Extreme modulation properties of aromatic fluorine

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Aromatic fluorine is a unique modulator of biological properties of organic compounds. In some rare cases, introduction of aromatic fluorine improved biological activity more than 100,000 times and even higher.

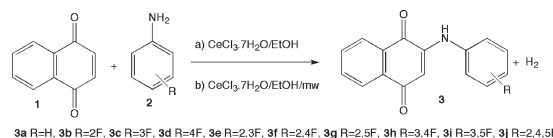


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Synthesis, spectral and electrochemical characterization of novel 2-(fluoroanilino)-1,4-naphthoquinones

Elisa Leyva^a, Lluvia I. López^b, Silvia E. Loredó-Carrillo^a, Margarita Rodríguez-Kessler^a, Antonio Montes-Rojas^a^aFacultad de Ciencias Químicas, Universidad Autónoma de San Luis Potosí, Av. Manuel Nava No. 6, Zona Universitaria, C.P. 78210, San Luis Potosí, S.L.P., Mexico^bFacultad de Ciencias Químicas, Universidad Autónoma de Coahuila, Blvd. V. Carranza e Ing. José Cárdenas s/n, Col. República, C.P. 25250, Saltillo, Coahuila, Mexico

Novel 2-(fluoroanilino)-1,4-naphthoquinones were prepared reacting the corresponding fluoroaniline and 1,4-naphthoquinone in the presence of a Lewis acid catalyst with strong oxidation properties such as CeCl₃·7H₂O. This preparation was also investigated under microwave irradiation. All 1,4-naphthoquinone derivatives were characterized by UV-Vis, IR, ¹H and ¹⁹F NMR, MS and cyclic voltammetry.



3a R=H, 3b R=2F, 3c R=3F, 3d R=4F, 3e R=2,3F, 3f R=2,4F, 3g R=2,5F, 3h R=3,4F, 3i R=3,5F, 3j R=2,4,5F

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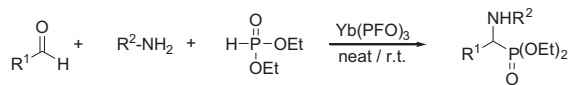
A facile synthesis of α -aminophosphonates catalyzed by ytterbium perfluorooctanoate under solvent-free conditions

Jun Tang^a, Limin Wang^{ab}, Wenbo Wang^a, Liang Zhang^a, Shengying Wu^a, Dan Mao^a

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^bKey Laboratory of Organofluorine Chemistry, Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, 354 Fenglin Road, Shanghai 200032, PR China

An efficient and one-pot protocol for the synthesis of α -aminophosphonates promoting by ytterbium perfluorooctanoate [Yb(PFO)₃] under solvent-free conditions is described.



17 samples, up to 95% yield

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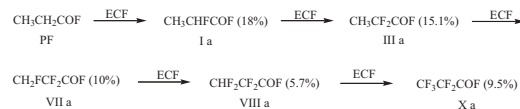
Products formed at intermediate stages of electrochemical perfluorination of propionyl and n-butyryl chlorides. Further evidence in support of NiF₃ mediated free radical pathway

T.M. Rangarajan^a, S. Sathyamoorthi^a, D. Velayutham^a, M. Noel^a, R.P. Singh^b, Raju Brahma^b

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The partially fluorinated HF soluble intermediates formed during the electrochemical perfluorination of propionyl chloride (PC) and n-butyryl chloride (n-BC) were analyzed after passing 0%, 25%, 50%, 75% and 100% of theoretical charge required for the fluorination of PC and n-BC. Free radical mechanism appears to be the major pathway as indicated by formation of large number of partially fluorinated compounds at every stage of electrochemical fluorination. For both PC and n-BC this mechanism also explains radical coupling, C–C bond cleavage and carbon chain isomerization.



Products formed after 50% of theoretical charge passed

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The effect of column and eluent fluorination on the retention and separation of non-fluorinated amino acids and proteins by HPLC

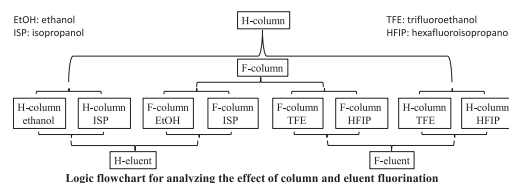
Katherine Joyner^a, Weizhen Wang^b, Yihua Bruce Yu^{ac}

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Fluorocarbon column and fluorocarbon eluents are compared side-by-side with their hydrocarbon counterparts to extract the effect of column and eluent fluorination on analyte retention and separation. Statistical analyses of retention time data revealed that optimal retention and separation is achieved when the fluorocarbon column is paired with ethanol.

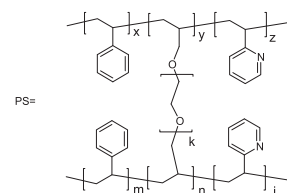
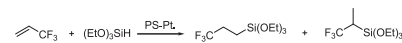
*J. Fluorine Chem.*, 132 (2011) 123

Synthesis of a novel functional polymer immobilized platinum complex and its application in the catalytic hydrosilylation of 3,3,3-trifluoropropene with triethoxysilane

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The novel polymer synthesized immobilized platinum and applied in the catalytic hydrosilylation of 3,3,3-trifluoropropene with triethoxysilane. The catalysts show the excellent activity and the high selectivity of β -adduct.

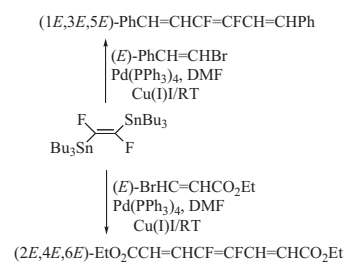


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Stereospecific preparation of (1*E*,3*E*,5*E*)-3,4-difluoro-1,6-diphenylhexatriene and (2*E*,4*E*,6*E*)-diethyl-4,5-difluoroocta-2,4,6-trienedioate

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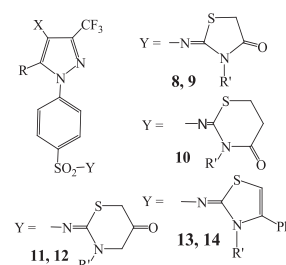


J. Fluorine Chem., 132 (2011) 131

Synthesis and biological evaluation of new 3-trifluoromethylpyrazolesulfonyl-urea and thiourea derivatives as antidiabetic and antimicrobial agents

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Fluorinated pyrazoles were prepared as hypoglycemic and antibacterial agents by condensation and subsequent cyclization of the thiourea derivatives of 3-trifluoromethylpyrazoles with ethyl bromoacetate, ethyl β -bromopropionate, 1,3-dichloroacetone and α -bromoacetophenone to give compounds **2–18**.

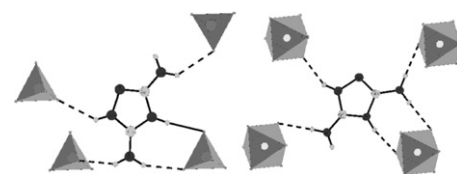


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Synthesis, crystal structure, IR-spectral data and some properties of 3,5-diamino-1,2,4-triazolium tetrafluoroborate and hexafluorosilicate

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3,5-Diamino-1,2,4-triazolium tetrafluoroborate (LH)BF₄ and hexafluorosilicate (LH)₂SiF₆ have been isolated and characterized by single-crystal X-ray structure determination, IR spectroscopy, mass spectrometry, TGA, solubility data, potentiometry.



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Unusual β,β' -coupling and β -alkylation of methyl 2,3,3-trifluoropropenoate by lithium diorganocuprates

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Lithium dimethyl- or di(*tert*-butyl)cuprate afforded the product of unusual coupling, while dibutyl- or diphenylcuprate gave the product of β -fluorine substitution in the substrate.

