



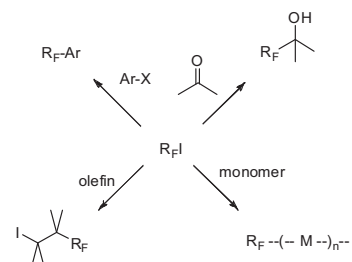
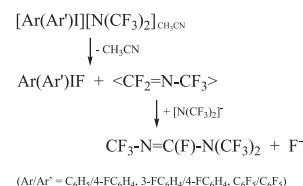
Graphical Abstracts/J. Fluorine Chem. 138 (2012) 1–2

Syntheses utilizing *n*-perfluoroalkyl iodides [R_FI, C_nF_{2n+1}-I]
2000–2010

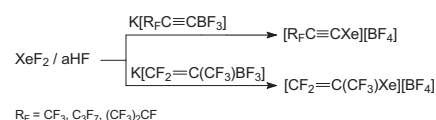
Peter M. Murphy, Christopher S. Baldwin, Robert C. Buck

E. I. DuPont de Nemours and Company, Inc., Wilmington, DE, USA

A review of the synthetic utility of *n*-perfluoroalkyl iodides from 2000 to 2010 presenting a compilation of the scientific literature describing chemical syntheses and a review of the applications and uses of the materials synthesized utilizing *n*-perfluoroalkyl iodides.

J. Fluorine Chem., 138 (2012) 3[Ar(Ar')I][N(CF₃)₂], effect of fluorine substituents on aryl groups of
iodonium cations and of coordinating bases on the fluoride transfer
from the anion to the cationMarkus E. Hirschberg^a, André Wenda^a, Hermann-Josef Frohn^b, Nikolai V. Ignat'ev^c^a*Inorganic Chemistry, Bergische Universität Wuppertal, Gaußstr. 20, D-42097 Wuppertal, Germany*^b*Inorganic Chemistry, Universität Duisburg-Essen, Lotharstr. 1, D-47048 Duisburg, Germany**J. Fluorine Chem.*, 138 (2012) 24Perfluoroorganylxenonium(II) salts from reactions of XeF₂ or
[FXe][SbF₆] with selected perfluorinated alk-1-yn-1-yl- and alk-
1-en-1-yltrifluoroborate salts in aHFVadim V. Bardin^a, Hermann-Josef Frohn^b^a*N.N. Vorozhtsov Novosibirsk Institute of Organic Chemistry, SB RAS, Acad. Lavrentjev Avenue 9, 630090 Novosibirsk, Russian Federation*^b*Inorganic Chemistry, University of Duisburg-Essen, Lotharstrasse 1, D-47048 Duisburg, Germany*

Perfluoroorganyltrifluoroborate salts do not react with XeF₂ in halocarbons or CH₃CN, but in aHF, HF activates XeF₂ and enables electrophilic attack on the nucleophilic *ipso*-carbon of the C–BF₃ moiety. This “xenodeboration” approach is a new method for the synthesis of Xe–C onium salts.

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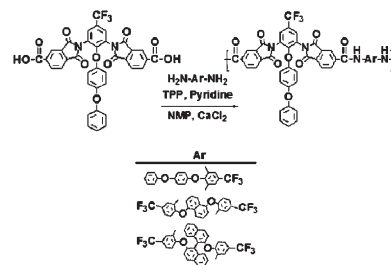
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Fluoro-containing poly(amide-imide)s with sterically hindered pendants: Synthesis and characterization

Hossein Behniafar, Farzin Zardoozi, Ali Rastkar

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These fluoro-containing poly(amide-imide)s displayed excellent organo-solubility, low crystallinity, reasonable thermal stability and glass transition temperatures, making them suitable for thermoforming processing.



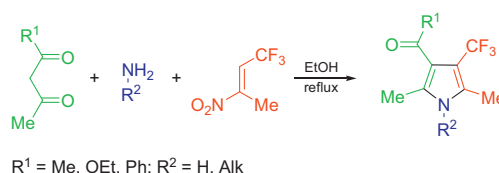
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Three-component synthesis of substituted β -(trifluoromethyl)pyrroles via Grob cyclization of 1,1,1-trifluoro-3-nitrobut-2-ene with 1,3-dicarbonylic compounds and ammonia or primary amines

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A variety of substituted β -(trifluoromethyl)pyrroles were easily synthesized in good yields by a one-pot, three-component Grob cyclization of 1,1,1-trifluoro-3-nitrobut-2-ene with 1,3-dicarbonylic compounds (ethyl acetoacetate, acetylacetone, and benzoylacetone) and ammonia or primary aliphatic amines.



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Fast and reliable method for the preparation of ortho- and para-[^{18}F]fluorobenzyl halide derivatives: Key intermediates for the preparation of no-carrier-added PET aromatic radiopharmaceuticals

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Conversion of various [^{18}F]fluorobenzaldehydes into the corresponding [^{18}F]fluorobenzyl halides (X = Cl, Br, I) is described. The aldehydes were reduced and halogenated on line on a solid support by using an aqueous solution of NaBH_4 and an aqueous concentrated acid solution (HCl, HBr, HI). This two step synthesis path proceeds fast and delivers the nca radiofluorinated aromatic products in high yields.

