

Volume 125, Issue 11, November 2004



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Graphical Abstracts/J. Fluorine Chem. 125 (2004) 1553-1559



J. Fluorine Chem., 125 (2004) 1579

New class of coordination compounds with noble gas fluorides as ligands to metal ions G. Tavčar, M. Tramšek, T. Bunič, P. Benkič, B. Žemva *Jožef Stefan Institute, Jamova 39, SI-1000 Ljubljana, Slovenia* $M^{n+}(AF_6)_n + zXeF_2 \xrightarrow{aHF}_{22 \circ C} [M^{n+}(XeF_2)_p](AF_6)_n + (z-p)XeF_2$ A=P, As, Sb, BiM = Mg, Ca, Sr, Ba, Cd, Pb, lanthanides

 $Mg(AsF_6)_2 + nXeF_4 \xrightarrow{aHF} [Mg(XeF_4)](AsF_6)_2 + (n-1)XeF_4$ $M(AsF_6)_2 + nXeF_6 \xrightarrow{aHF} 22 \circ C \rightarrow MF_2 + 2Xe_2F_{11}AsF_6 + (n-4)XeF_6$ M = Mg, Ca, Sr

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A computational study of $\text{Sb}_n \text{F}_{5n}$ (<i>n</i> =1–4) Implications for the fluoride ion affinity of <i>n</i> SbF ₅	
H. Donald Brooke Jenkins ^a , Ingo Krossing ^b , Jack Passmore ^c , Ines Raabe ^b	$Sb_nF_{5n} \xrightarrow{\Delta_r H^o_{depoly.} = ?} n SbF_5$
^a Department of Chemistry, University of Warwick, Coventry CV4 7AL, West Midlands, UK ^b Universität Karlsruhe, Institut für Anorganische Chemie, Engesserstr. Geb. 30.45, 76128 Karlsruhe, Germany ^c Chemistry Department, University of New Brunswick, Fredericton, NB, Canada E3B6E2	$Sb_{n}F_{5n(g)} + F'_{(g)} \xrightarrow{FIA = ?} [Sb_{n}F_{5n+1}]'_{(g)}$ $FIA = ? \qquad (Sb_{n}F_{5n+1})^{T}$
nF_{5n} (n=1-4) as well as the fluoride ion affinities of $Sb_nF_{5n}(g)$, $nSbF_5(g)$ and $nSbF_{5(1)}$ were calculated.	$n \operatorname{SbF}_{5(1)} + F_{(g)} \longrightarrow [\operatorname{Sb}_n F_{5n+1}]_{(g)}$

Claisen rearrangements based on vinyl fluorides Frank Tranel, Günter Haufe Organisch-Chemisches Institut, Westfälische Wilhelms-Universität Münster, Corrensstr. 40, D-48149 Münster, Germany 2-Fluoroalk-1-en-3-ol esters prepared in two steps from 2-fluoroalk-1-enes undergo Claisen rearrangements to form 2-substistuted 4fluoroalk-4-enecarboxylic acids. J. Fluorine Chem., 125 (2004) 1609 Syntheses, structures and bioactivities of fluorine-containing phenylimino-thia(oxa)zolidine derivatives as agricultural bioregulators Xuhong Qian^a, Xiaoyong Xu^b, Zhibin Li^b, Zhong Li^b, Gonghua Song^b ^aState Key Laboratory of Fine Chemicals, Dalian University of Technology, Dalian 116012, China ^bShanghai Key Laboratory of Chemical Biology, Institute of Pesticides and Pharmaceuticals, East China University of Science and Technology, Shanghai 200237, China From insight into the structure of trehazolin as trehalase inhibitor, six series of fluorine-containing phenyliminothiazolidines (oxazolidines) derivatives were designed and prepared through a convenient synthesis of fluoroaryl isothiocyanate and a one-pot facile synthesis in high yield of fluorophenyl aminobenzoxazoles by cyclodesulfurization. The structures of the target compounds were confirmed with using IR, NMR, MS and elemental analysis. Their X-ray crystal analysis suggested that there were novel intermolecular (sp^2 CF...H₃C-) and intramolecular (sp^2 CF...HN) hydrogen bonds between the fluorine atom on benzene ring and hydrogen atom of methyl group or amino group on five-membered heterocycle. Their fungicidal activities against Rhizoctonia solani and Pyricuraria oryzae at 100 ppm were determined. J. Fluorine Chem., 125 (2004) 1621 Reactions of quadricyclane with fluorinated nitrogen-containing compounds. Synthesis of 3-aza-4-perfluoroalkyl-tricyclo [4.2.1.0^{2,5}]non-3,7-dienes C₂F₅N=CFCF Viacheslav A. Petrov, Frederic Davidson, Will Marshall DuPont Central Research and Development, Experimental Station, P.O. Box 80328, Wilmington, DE 19880-0328, USA **B**₄CN The reaction of quadricyclane with perfluoroazaalkenes and perfluorinated nitriles leads to high yield formation of the corresponding cycloadducts. 56-81%

 $R_{f} = CF_{3}, C_{2}F_{5}, n-C_{3}F_{7}$



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Surface modification of several carbon-based materials: comparison between CF₄ rf plasma and direct F₂-gas fluorination routes

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In fluorinated carbon-based materials, the fitting of F 1s XPS envelopes - as those of C 1s ones - allows to assign the components to the different types of C-F bonds formed at the surface of the materials, as shown below in the case of F2-treated nitrile-butadiene elastomer.



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Fluorothiazynes, 50 years old and still exciting: electrophilic attack at the thiazyl nitrogen of $NSF_2NS(O)F_2$

Rüdiger Mews^a, Tobias Borrmann^b, Reinhard Hoppenheit^c, Enno Lork^a, Simon Parsons^d, Jan Petersen^a, Markus Schröter^a, Wolf-Dieter Stohrer^b, Alfred Waterfeld^e, Paul G. Watson^a ^aInstitute of Inorganic and Physical Chemistry, University of Bremen, Leobener Str. NW2, D-28334 Bremen, Germany ^bInstitute of Organic Chemistry, University of Bremen, Leobener Str. NW2, D-28334 Bremen, Germany ^cLindenstrasse 31, D-37181 Hardegsen, Germany ^dSchool of Chemistry, The University of Edinburgh, King's Buildings, West Mains Road, Edinburgh EH9 3JJ, Scotland ^eDepartment of Chemistry, University of Alabama, Tuscaloosa, USA

J. Fluorine Chem., 125 (2004) 1657 Surface fluorination of the cathode active materials for lithium secondary battery Susumu Yonezawa, Masahiro Yamasaki, Masayuki Takashima Li/Li⁺ 4.2 Untreated sample Department of Materials Science and Engineering, Faculty of Engineering, 4 S University of Fukui, Bunkyo 3-9-1, Fukui 910-8507, Japan 3.8 The electrochemical properties of LiMn2O4 as a cathode active material Potential 3.6 of lithium secondary battery was improved by its fluorination with F2 gas. 3.4 3.2 (fluorinated sample) 3 20 40 60 80 100 120 Capacity / mA h g-1 J. Fluorine Chem., 125 (2004) 1663

Synthesis, vibrational and NMR spectroscopic characterization of $[N(CH_3)_4][IO_2F_2]$ and X-ray crystal structure of [N(CH₃)₄]₂[IO₂F₂][HF₂]

Michael Gerken^{a,b}, Johnathan P. Mack^a, Gary J. Schrobilgen^b, Reijo J. Suontamo^c

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 $[N(CH_3)_4][IO_3F_2]$ was synthesized and characterized by vibrational and ¹⁹F NMR spectroscopy. The crystal structure of $[N(CH_3)_4]_2[IO_2F_2][HF_2]$ provides an unique example of a symmetric bifluoride bridge. The geometry and vibrational frequencies of IO2F2- were calculated at the MP2 and SVWN levels of theory.



C

J. Fluorine Chem., 125 (2004) 1671 New fluoropolymer materials Colin D. Wood^a, Udo Michel^a, Jason P. Rolland^a, 100 um 100 µm 100 µm Joseph M. DeSimone^{a,b} ^aDepartment of Chemistry, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA B Α. ^bDepartment of Chemical Engineering, North Carolina State University, Raleigh, NC, USA A solvent resistant photocurable "liquid Teflon" which can be used for microfluidic device fabrication, actuation of the valve is shown. J. Fluorine Chem., 125 (2004) 1677

Fluorinations, chlorinations and brominations of organic compounds in micro reactors

Patrick Löb, Holger Löwe, Volker Hessel

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An overview of the application of micro reactors for fluorination (e.g. of toluene derivatives), chlorination and bromination of organic compounds using the elemental halogens is given.

+ F₂; - HF micro reactor

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Ternary and tetrahedral symmetry in hybrid fluorides, fluoride carbonates and carbonates

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J. Fluorine Chem., 125 (2004) 1723

Recent achievements in the synthesis and characterization of metal hexafluorantimonates and hexafluoroaurates $MF_2 + 2AuF_3$

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CuSbF₆, a rare example of Cu(I) compound in a pure fluorine environment was synthesized. In the reactions of $MF_2/2AuF_3$ with KrF_2 or UV-irradiated F_2 in aHF, the synthesis of new Au(V)-salts was achieved. Previously reported syntheses of M(AuF₆)₂ (M = Mg, Ca, Sr, Ba) were reinvestigated.

 $MF_{2} + 2AuF_{3}$ $KrF_{2} \quad \text{or UV-irradiated } F_{2}$ aHF, 298 K $M(AuF_{6})_{2}$ (M = Ni, Cu, Ag, Zn, Cd, Hg)





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Synthesis of fluorine-containing compounds under operationally convenient conditions

Vadim A. Soloshonok, Dmitrii O. Berbasov

Department of Chemistry and Biochemistry, University of Oklahoma, Norman, OK 73019, USA

This paper describes a concept of "operationally convenient conditions", gives a short overview of the synthetic applications of biomimetic reductive amination under such conditions and discusses new kinetic data on the mechanism of 1,3-proton shift transfer.



J. Fluorine Chem., 125 (2004) 1765 Preparation of fluorine compounds of groups 13 and 14: a study case for the diagonal relationship of aluminum and germanium Herbert W. Roesky

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J. Fluorine Chem., 125 (2004) 1771

Polyfluoroorganotrifluoroborates and -difluoroboranes: interesting materials in fluoroorgano and fluoroorgano-element chemistry

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> $C_6F_{13}C_2H_4BF_2$ $K[(C_2F_5)_2BF_2]$ $[C_{e}F_{5}(4-C_{5}F_{4}N)][BF_{4}]$ [p-FC₆H₄(trans-CF₃CF=CF)I][BF₄]



Dissolution behavior of tetrafluoroethylene-based fluoropolymers for 157-nm resist materials

Takuji Ishikawa^a, Tesuhiro Kodani^a, Tomohiro Yoshida^a, Tsukasa Moriya^a, Tsuneo Yamashita^a, Minoru Toriumi^a, Takayuki Araki^a, Hirokazu Aoyama^a, Takuya Hagiwara^b, Takamitsu Furukawa^b, Toshiro Itani^b, Kiyoshi Fujii^b

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We have synthesized various main-chain fluorinated polymers and studied their transparency, dry-etch durability and solubility.

J. Fluorine Chem., 125 (2004) 1801

Environmental assessment of CFC alternatives. Rate constants for the reactions of OH radicals with fluorinated compounds

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The rate constants for reactions of OH radicals with CF₃OCHFCF₃, and CF₃CHFCF₃ have been measured using absolute rate method. The Arrhenius rate constants have been determined from these kinetic data as: $k(CF_3OCHFCF_3) = (4.39 \pm 1.38) \times 10^{-13} exp[-(1780 \pm 100)/T]cm^3$ molecule⁻¹ s⁻¹, and $k(CF_3CHFCF_3) = (6.19 \pm 2.07) \times 10^{-13} exp[-(1830 \pm 100)/T]cm^3$ molecule⁻¹ s⁻¹.