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The 1-N-(methoxycarbonyl-2-phenylethyl)imino-2,2,2-trifluoroethanephosphonate systems are not stable enols of carboxylic esters

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The claim that 1-N-(1-methoxycarbonyl-2-phenylethyl)imino-2,2,2-trifluoroethane phosphonates 2 exist as their stable enols of carboxylic esters tautomers,

was refuted on three grounds, reexamination of NMR spectra, experimental results, and B3LYP calculations.

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A study of the thermal decarboxylation of three perfluoropolyether salts

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The thermal decarboxylation of KOOCCF2-Rf-CF2COOK, $R_f = -O-$, $-OCF_2O-$, $-OCF_2CF_2O-$, has been investigated and the products and kinetics of the main reactions have been defined. The experimental data showed that the two consecutive decarboxylation reactions have similar rate constants and activation energies.

$$\begin{array}{ccc} \text{KOOCCF}_2\text{R}_1\text{CF}_2\text{COOK} & \frac{k_1}{\text{H}_2\text{O}} & \text{HCF}_2\text{R}_1\text{CF}_2\text{COOK} + \text{KHCO}_3 \\ & \frac{k_2}{\text{H}_2\text{O}} & \text{HCF}_2\text{R}_1\text{CF}_2\text{H} & + \text{KHCO}_3 \end{array}$$

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Poly-silafluoroalkyleneoligosiloxanes: a class of fluoroelastomers with low glass transition temperature

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$$HO \left\{ \begin{matrix} CH_{3} & CH_{3} & R \\ -(Si-C_{n}H_{2n}-R"_{F}-C_{n}H_{2n}-Si-O)_{w} & -(Si-O)_{v} \\ R" & R" & R' \end{matrix} \right\} H$$

Synthesis of unsaturated poly-silafluoroalkyleneoligosiloxanes which, after crosslinking, gave elastomeric materials characterized by good flexibility at low temperature, glass transition temperature below -45°C and good thermooxidative stability, over 250°C, that are complementary to polyfluoroolefin elastomers.

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Synthesis of fluoro-substituted monomers bearing a functionalised lateral chain. Part 2. Preparation of sulfoxides and sulfones containing monomers

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^bAtofina, Centre de Recherches et Développement de l'Est, B.P. 61005, 57501 Saint-Avold Cedex, France

O SOy q C_nF_{2n+1} p = 2, 3, 4, 11 q = 2, 11 n = 6, 6-8*, 8

^cLaboratoire de Chimie et Applications (LCA), Groupe Synthèse Organique EA-3471, Université de Metz, Ile du Saulcy, 57012 Metz Cedex 01, France The oxidation of ω-(ω-perfluoroalkylalkyl-sulfanyl)-alkyl acrylates to corresponding sulfoxides and sulfones is described in this paper.

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Fluoride-assisted trifluoromethylation of aromatic thiones with (trifluoromethyl)trimethylsilane

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Bd du 11 Novembre 1918, F-69622 Villeurbanne Cedex 69622, France

$$\begin{array}{c} \text{Ar} & \text{CF}_{3}\text{TMS} \\ \text{Ar} & \text{S} \\ \text{TBAF.3 H}_{2}\text{O} \\ \text{Ar} & \text{C}_{2 \text{ eq}}\text{)} \\ \text{Ar} & \text{C}_{6}\text{H}_{5} \\ \text{4-MeO-C}_{6}\text{H}_{4} \end{array} \begin{array}{c} \text{Ar} & \text{SCF.} \\ \text{Ar} & \text{H} \\ \text{(35-37\%)} \\ \text{+} \\ \text{Ar} & \text{SH} \\ \text{Ar} & \text{CF}_{3} \\ \text{(13-14\%)} \end{array}$$

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N-Bis(methylthio)methylen-trifluoromethanesulfonylamide CF₃SO₂N=C(SCH₃)₂: new reagent for the preparation of *N*-trifluoromethylsulfonylimino carbonic and thiocarbonic acids derivatives

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Synthesis and NMR studies of 2- and 3-fluorosubstitued five-membered heterocycles

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$$\begin{array}{c|c} H(Br) & & & \\ & & 1) \text{ BuLi, THF} \\ & & 2) \text{ FN(SO}_2\text{Ph)}_2 \end{array}$$

A full set of 2- and 3-fluoro-substituted thiophenes, pyrroles and furans has been synthesized for the first time. The $^1J_{\rm CC}$ couplings measured for these compounds are the largest among those determined for five-membered heterocycles so far.

X = O, S, N-Me

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Synthesis of amino terminated semifluorinated long-chain alkanethiols

C. Amato, P. Calas

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$$I(CF_2)_{n1} \xrightarrow{\text{H}_2\text{C}=\text{CH}-(CH_2)_9-\text{OH}} I(CF_2)_n \xrightarrow{\text{N}-\text{allylphthalimide}} I(CF_2)_n \xrightarrow{\text{N}-\text{a$$

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Selective mono- and di{(perfluoroalkyl)acylation} of ferrocene

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$$X$$
 C_6F_{13}
 F_e
 F_e
 C_6F_{13}
 C_6F_{13}

$$X = C(O), CH_2$$

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A novel perfluoromonomer: perfluoro-2,3-dihydro-1,4-benzodioxin

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Synthesis and solution properties of sulfate-type hybrid surfactants with a benzene ring

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FmPHnOS aqueous solution showed low critical micelle concentration and high surface activity. Aggregation number of FmPHnOS micelles ranged from 6 to 45 and the hydrodynamic radius of the micelles was in a range of 1.4–3.1nm.

$$\begin{array}{c|c} \mathsf{OSO_3Na} \\ \mathsf{F}(\mathsf{CF_2})_m & -\mathsf{CH} \\ \mathsf{(CH_2)_nH} \end{array}$$

m = 4, 6, 8, n = 3, 5, 7

FmPHnOS

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Novel fluorinated polymer from 18-crown-6 by radical polyaddition

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^bDepartment of Applied Chemistry, Faculty of Engineering, Saitama Institute of Technology, 1690 Fusaiji, Okabe, Saitama 369-0293, Japan

$$\begin{array}{c} \begin{array}{c} CF_3 \\ CF_2-CH-O-C \\ O \end{array} \begin{array}{c} CF_3 \\ CF_2-CH-O-C \\ O \end{array} \begin{array}{c} CF_3 \\ CF_2-CH-CF_2 \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \\ O \\ O \\ O \\ O \\ O \end{array} \begin{array}{c} O \\ O \end{array} \begin{array}{c} O \\$$

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Studies on the reaction of unsymmetrical trifluoromethyl 1,2-phenylenediamine with various ketones leading to novel fluorinated heterocycles

G. Venkat Reddy, V.V.V.N.S. Rama Rao, D. Maitraie, S. Ravikanth, R. Yadla, S.N. Reddy, B. Narsaiah,

P. Shanthan Rao

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Synthesis of polyfluorophenyl substituted-4,5-dihydropyrazole derivatives via 1,3-dipolar cycloaddition of nitrile imine with ethyl acrylate

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^aSchool of Chemistry and Pharmaceutics, East China University of Science and Technology, Shanghai 200237, China

^bLaboratory of Organofluorine Chemistry, Shanghai Institute of Organic Chemistry, Chinese Academy of Science, Shanghai 200032, China F F NHN=CHR+ CH_2 = $CHCO_2Et$ OD_2 OD_2 OD_2 OD_3 OD_4 OD_4 OD_5 OD_4 OD_5 OD_5 OD_5 OD_6 OD_6 OD_7 OD_8 OD_8 OD_8 OD_9 OD_9

3-Substituted-1-(4-chloro-2,3,5,6-tetrafluorophenyl)-5-ethoxycarbonyl-4,5-dihydropyrazoles (3) were synthesized by reaction of aldehyde 4-chloro-2,3,5,6-tetrafluorophenylhydrazones (1) with [bis(acetoxy)iodo]benzene in the presence of ethyl acrylate.

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Kinetic aspects involved in the simultaneous enzymatic synthesis of (S)-3-fluoroalanine and (R)-3-fluorolactic acid

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1-Fluoro-2,4,6-trichloro-1,3,5-triazinium tetrafluoroborate: synthesis, characterization, and ability to effect electrophilic aromatic substitution

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The synthesis of 1-fluoro-2,4,6-trichloro-1,3,5-triazinium tetrafluoroborate,

 $[(ClCN)_3F]^+[BF_4]^-$ (1), and its application of 1 as an electrophilic fluorinating agent are reported.