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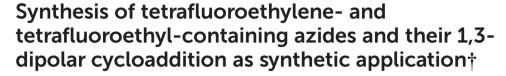
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Tetrafluoroethylene-containing azides are accessed in two steps (one pot) from tetrafluoroalkyl bromides by metalation and reaction with electrophilic azides. Subsequent copper(i)-catalyzed azide—alkyne cycloaddition afforded N-tetrafluoroethyl and N-tetrafluoroethylene 4-substituted 1,2,3-triazoles. In addition, the protocol for the synthesis of 4,5-disubstituted 1,2,3-triazoles is presented.

Tetrafluoroethyl- and tetrafluoroethylene-containing compounds have been recently intensively investigated in agrochemistry, medicinal chemistry, and materials science as alternatives to well established fluorinated functional groups such as trifluoromethyl, perfluoroalkyl, and difluoromethyl. 1-9 For instance, tetrafluorinated sugar analogues were found to be significantly less hydrophilic than the parent carbohydrates, and thus exhibit enhanced protein-carbohydrate binding capabilities. 10-14 The tetrafluoroethylene motif is found in insecticides^{15,16} and liquid crystals.^{17,18} The tetrafluoroethyl- and triazole-containing molecule tetraconazole displays fungicidal activity. 19 Despite important recent advances in synthetic methods toward these compounds, 20-26 protocols affording CF2CF2-containing molecules of the unprecedented structure are still in demand. Our contribution to the field has been in the design and application of nucleophilic, 27-29 electrophilic 30 and radical 11 reagents for the CF₂CF₂ transfer which provided straightforward synthetic access to a variety of tetrafluoroethyl- and tetrafluoroethylenecontaining compounds. In the latest report, tetrafluoroethyl bromides were reacted with i-PrMgCl·LiCl to afford metalated species stable at low temperature and displaying excellent reactivity with a range of electrophiles with one example of an

In this article, the synthesis of tetrafluoroethylene-containing azides is disclosed by metalation of tetrafluoroethyl bromides through organomagnesium or silicon species and their application in CuAAC is shown in the preparation of tetrafluoroethyl- and tetrafluoroethylene-containing triazoles.‡

Magnesiation of tetrafluoroethyl bromides (1) was performed with i-PrMgCl·LiCl (turbo Grignard reagent) in THF at -78 °C. We have earlier found that the metalated species are stable at this temperature and metalation times range from less than 5 min for R¹ = PhS and alkyl to half an hour or slightly longer for R¹ = imidazolyl, pyrazolyl and ArO.³² An electrophilic azide source (tosyl or nonaflyl azide working equally well) was then added to the generated Grignard reagent and the temperature was allowed to rise to ambient temperature over a period of 3 hours. In this way, good yields of tetrafluoroethyl azides (2) were obtained with electronneutral, -withdrawing or -donating groups on the aryl ring of the aryloxy substituent $(R^1 = ArO)$ (Table 1). The 4-bromopyrazolyl derivative (1h) afforded the corresponding azide 2h in moderate yield; however, the reaction was highly regioselective (only the bromine atom on the fluorinated carbon reacted). In contrast, the phenylthio derivative 1i, under the same reaction conditions, provided selectively and in high conversion PhSCF₂CF₂SPh instead of the expected azide 2i. This outcome can be explained by high susceptibility of the Grignard intermediate to oxidation to radicals, partial fragmentation giving PhS radicals, which prefer to react with electrophilic PhSCF2CF2 radicals. Therefore, for the synthesis of 2i, an alternative method was used starting from PhSCF₂CF₂SiMe₃.⁴⁷ Nucleophilic fluoroalkyl transfer, after activation with CsF, to either tosyl or nonaflyl azide afforded high yields of 2i (Scheme 1).

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electrophilic azide. ³² Azides are highly valuable compounds in organic synthesis – some fluoroalkyl azides are known, ^{33–40} including $CF_3N_3^{41,42}$ and perfluoroalkyl analogues for which we have recently reported a new and practical synthetic method. ⁴³ Perfluoroalkyl azides were investigated in copper(ι)-catalyzed azide–alkyne cycloadditon (CuAAC) ^{44–46} furnishing *N*-perfluoroalkyl-1,2,3-triazoles. ⁴³

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Entry 1

2 3

5

6

Table 1 Synthesis of tetrafluoroethylene-containing azides 2

	1 PringCitud R1 F THF, -78 °C 2. R2SO ₂ N ₃ (2 THF, -78 °C	equiv)	F ─N ₃
7	R^1	R^2	2, Yield (%)
	PhO	<i>p</i> -Tol	2a , 92
	$4-FC_6H_4O$	<i>p</i> -Tol	2b , 79
	4-BrC ₆ H ₄ O	<i>p</i> -Tol	2c, 76
	$4-(MeO)C_6H_4O$	n-C ₄ F ₉	2d , 84
	$3-(Me_2N)C_6H_4O$	n-C ₄ F ₉	2e , 87
	$3-(CF_3)C_6H_4O$	n-C ₄ F ₉	2f , 73
	$2,4-(t-Bu)_2C_6H_3O$	n-C ₄ F ₉	2g, 71
	N-W	n-C ₄ F ₉	2h, 44

n-C₄F₉

2i, 0

$$\begin{array}{c} \text{PhS} & \begin{array}{c} \text{F} \\ \text{F} \\ \text{F} \end{array} & \begin{array}{c} \text{CsF (1.28 equiv)} \\ \text{R}^2 \text{SO}_2 \text{N}_3 \ (2 \text{ equiv}) \\ \text{DMF, } -60 \ ^{\circ}\text{C, } 1 \ \text{h} \\ -60 \ ^{\circ}\text{C to rt, } 3 \ \text{h} \end{array} & \begin{array}{c} \text{PhS} \\ \text{F} \\ \text{F} \end{array} & \begin{array}{c} \text{F} \\ \text{N}_3 \\ \text{Zi} \end{array} \\ \\ \text{R}^2 = p \text{-Tol} & 89\% \\ \text{R}^2 = n \text{-C}_4 \text{Fg} & 72\% \end{array}$$

Scheme 1 Synthesis (2-azido-1,1,2,2-tetrafluoroethyl)(phenyl) sulfane (2i)

With azides 2, CuAAC was investigated. Our earlier work with azidotrifluoromethane and other azidoperfluoroalkanes⁴³ showed that copper(1) 3-methylsalicylate (CuMeSal) was a preferred catalyst in terms of catalytic activity, good solubility in organic solvents and high stability. Therefore, it was favored over other Cu(I) sources. In the reaction, equimolar amounts of aryl, heteroaryl or alkyl acetylenes and 1 mol% of CuMeSal afforded good to high yields of N-tetrafluoroalkylated 4-substituted 1,2,3-triazoles 4 (Table 2). The examples showing moderate yields resulted from isolation difficulties rather than a lack of reactivity or side reactions. Exclusive formation of 1,4-disubstituted triazoles was observed in all cases. Compounds of the type 4 are unknown in the literature with the exception of a triazole formed by thermal cycloaddition of N₃CF₂CF₂CO₂Me. 48 Reductive cleavage of the phenylsulfanyl group provided N-tetrafluoroethyl-containing triazole 5 in good yield (Scheme 2). However, attempts to prepare HCF2CF2N3 from 2i under the same reaction conditions met with limited success partially due to product volatility and also due to some unknown side reactions. Nevertheless, the in situ formed HCF₂CF₂N₃ transformed to N-tetrafluoroethyl triazole 5 (Scheme 2).

Finally, N-tetrafluoroethyl 4,5-disubstituted 1,2,3-triazoles were accessed in two steps from azides 2a and 2d and copper acetylides 6j and 6m, respectively. The reaction with iodine and a base afforded iodotriazoles 7aj and 7dm, respectively, in

Table 2 Synthesis of N-tetrafluoroalkyl triazoles 4

Scheme 2 Preparation of N-tetrafluoroethyl 4-substituted 1,2,3-triazole 5.

good to high yields which were amenable to Sonogashira or Suzuki-Miyaura cross-coupling reactions giving substituted triazoles 8 and 9, respectively (Scheme 3).

Crystallographic analysis of triazoles 4dm, 4hm, 4ij, 5 and 7dm confirmed their molecular structures (see the ESI†) and also showed an interesting difference in conformation at the N-CF₂CF₂-X unit. While in compounds 4hm (X = N) and 4ij (X = S) the N and X atoms are approximately anti-periplanar, in compounds 4dm, 7dm (X = O) and 5 (X = H) the N and X atoms are gauche as illustrated in Fig. 1. The reasons for this difference is unclear; however, an interplay between the steric

PhS ^a PhSCF₂CF₂SPh formed instead of 2i.

Scheme 3 Synthesis of 4,5-disubstituted 1,2,3-triazoles 8 and 9

Fig. 1 Schematic representation of the N-C-C-X conformation in the crystal structures of *N*-tetrafluoroethyl 1,2,3-triazoles.

repulsion of N and X atoms favoring *anti-periplanar* conformation and favorable vicinal fluorine-fluorine *gauche* conformation (three *vs.* two F, F *gauche* arrangements) should be taken into consideration.

In conclusion, magnesiation of tetrafluoroethyl bromides with i-PrMgCl·LiCl and subsequent reaction with tosyl or non-aflyl azides afforded tetrafluoroethylated azides. (2-Azido-1,1,2,2-tetrafluoroethyl)(phenyl)sulfane was prepared by fluoride-initiated nucleophilic fluoroalkyl transfer from the corresponding silane. The azides proved to be competent partners in copper(i)-catalyzed azide alkyne cycloadditions with terminal alkynes to generate *N*-tetrafluoroalkylated 4-substituted 1,2,3-triazoles. Reductive cleavage of the PhS group in the tetrafluoroethyl unit afforded *N*-tetrafluoroethyl azide. 4,5-Disubstituted triazoles were accessed by CuAAC of the triazoles with copper acetylides in the presence of iodine followed by transition metal-catalyzed cross-coupling reactions.

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