Fluoroform (CHF₃)

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Introduction

Fluoroform, which is generated in ~20 kilotons/year as a side product of Teflon manufacture,¹² is a low-boiling (-82 °C), non-toxic and non-ozone-depleting gas.¹³ However, fluoroform is a potent greenhouse agent and there is great interest in methods for use of the gas as a synthetic reagent. The direct use of trifluoromethyl anion as a nucleophile has been challenging due to facile α-elimination to fluoride and difluorocarbene (Scheme 1). The activation of Me₃SiCF₃ (often described as the Ruppert–Prakash reagent) with Lewis bases (most commonly catalytic fluoride) to afford pentavalent silicon species as nucleophiles has enabled trifluoromethylation of a number of electrophiles.⁴ The trifluoromethyl group has great importance in medicinal, agrochemical and materials science.⁵ This spotlight describes recently reported methods for tri- and difluoromethylation based upon fluoroform.

![Scheme 1](image)

**Scheme 1** α-Elimination of trifluoromethyl anion to difluorocarbene and fluoride.

<table>
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<th>Table 1 Use of Fluoroform (CHF₃)</th>
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(A) Prakash and co-workers have demonstrated successful deprotonation of CHF₃ in etheral solvents using KHMDS; the method was applied to the synthesis of trialkyl trifluoromethyl silane, trifluoromethyl fluoroborate and trifluoromethane sulfonic acid (Eq 1–3). Nucleophilic trifluoromethylation was also demonstrated for 1,2-additions to non-enolizable ketones and esters, chalcones, formate esters, and displacement of benzyl bromide and methyl benzoate was also demonstrated to proceed in moderate yields (illustrated for non-enolizable carbonyl compounds, Eq 4). Attempts to apply sodiated and lithiated bases were unsuccessful, revealing that potassium is essential for stable :CF₃ generation.¹

(B) Dolbier and co-workers utilized CHF₃ as a difluorocarbene source, preparing difluoromethyl ethers and thioethers from the corresponding phenols and thiophenols in the presence of KOH.⁶ The method, which involves biphasic conditions and can be conducted at atmospheric pressure, furnishes moderate to good yields of ethers.

(C) Mikami and co-workers discovered that lithium enolates react with CF₃H to achieve α-difluoromethylation via displacement of a C–F bond; the method has been applied thus far only to lactone and lactam enolates.⁷
(D) The Shibata group explored activation of fluoroform with organ-ic super bases and demonstrated that P4-t-Bu can be used to generate a stable CF3 anion at –30 °C in THF solvent; under these conditions, the anion was stable at up to –10 °C. The anion was applied to the trifluoromethylation of aromatic carbonyl compounds.5

(E) Nucleophilic trifluoromethylation using CHF3 in the presence of P4-t-Bu has also been applied to reactions with acid chlorides, esters, carbon dioxide, and epoxides.8 Acid chlorides react to provide mixtures of trifluoromethyl ketones and 3° alcohols (Eq 1). In contrast, methyl benzoate reacted to give only phenyl trifluoromethyl ketone (Eq 2). The trifluoromethyl anion reacts with carbon dioxide to produce trifluoroacetic acid (Eq 3). Curiously, reaction with terminal epoxides occurs at the more hindered carbon to give a 3° alcohol (Eq 4).

(F) Grushin and co-workers synthesized CuCF3 by cupration of fluoroform using CuCl and t-BuOK in DMF.3a The reagent was applied to trifluoromethylation of aryl and heteroaryl halides,3b aryl boronates,9 and α-halo ketones.10

(G) Potash and Rozen demonstrated the formation of trifluoromethyl thioethers through reaction of CuCF3 with alkyl, aryl, and heteroaryl thiocyanates.2 Similarly, fluoroform-derived CuCF3 was applied to the synthesis of trifluoromethyl selenides from selenocyanates.11

References