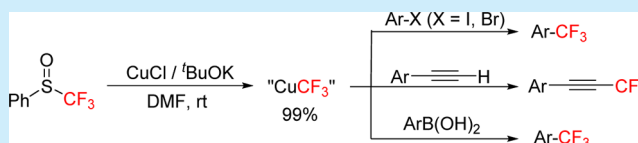


Copper-Mediated Trifluoromethylation Using Phenyl Trifluoromethyl Sulfoxide

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S Supporting Information

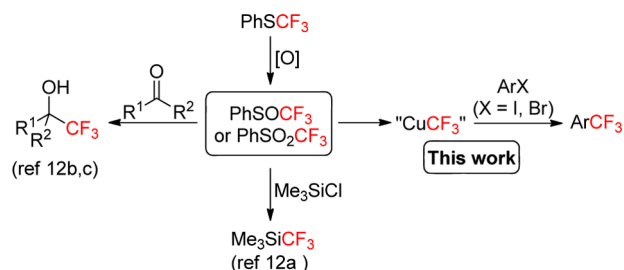
ABSTRACT: A new method for the generation of trifluoromethylcopper ("CuCF₃") species from readily available phenyl trifluoromethyl sulfoxide has been developed. The "CuCF₃" reagent can be applied in efficient trifluoromethylations of aryl iodides and activated aryl bromides in the absence of additional ligands. Furthermore, the "CuCF₃" species can also undergo oxidative cross-coupling with terminal alkynes and arylboronic acids.



Organofluorine compounds have been the subject of increasing research activity in recent years, since the incorporation of fluorine into bioactive compounds could enhance their lipophilicity and metabolic stability, among others.^{1,2} In this context, aromatic compounds bearing CF₃ group(s) are frequently used in medicinal and agricultural chemistry. Hence, it is of great interest to develop new methods for the selective introduction of trifluoromethyl groups onto aromatic rings. As a result, various synthetic methods of aromatic trifluoromethylation have been reported, including transition-metal-mediated or -catalyzed trifluoromethylation³ as well as radical⁴ and electrophilic trifluoromethylation.⁵ Among various transition-metal-assisted methods for incorporating CF₃ into arenes, copper-mediated trifluoromethylation has been most extensively studied due to its high efficiency and the relatively low cost of copper.⁶ Trifluoromethylcopper ("CuCF₃"), which is often generated prior to the desired reaction, is a relatively stable and highly useful trifluoromethylating agent.^{6f,j,l}

Me₃SiCF₃ (Ruppert–Prakash reagent),⁷ a useful nucleophilic trifluoromethylating agent, can be used to generate "CuCF₃" utilizing a copper(I) salt.⁸ A new method for the efficient generation of "CuCF₃" from CF₃H has been developed by Grushin et al. via treatment of CuCl and 2 equiv of ^tBuOK.^{6f,9} More recently, Mikami et al. have also found that PhCOCF₃ can be used to generate "CuCF₃" with a similar protocol.¹⁰ Previously, we were interested in the exploration of the synthetic application of fluorinated sulfones, sulfoxides, and sulfoximines.¹¹ Phenyl trifluoromethyl sulfone (PhSO₂CF₃) and phenyl trifluoromethyl sulfoxide (PhSOCF₃) as good nucleophilic trifluoromethylation reagents have been investigated.¹² Both PhSO₂CF₃ and PhSOCF₃ are commercially available and can be easily prepared from PhSCF₃ through oxidation (Scheme 1).¹³ However, transition-metal-mediated trifluoromethylation with PhSO₂CF₃ or PhSOCF₃ is rare. It was found that CuI-mediated trifluoromethylation of iodobenzene with PhSO₂CF₃ was

Scheme 1. Preparation and Application of Phenyl Trifluoromethyl Sulfone or Sulfoxide



inefficient, and a 26% yield of benzo-trifluoride was obtained.^{12b} Herein, we report an efficient method to generate "CuCF₃" from PhSOCF₃ and its trifluoromethylation of aryl halides, terminal alkynes, and arylboronic acids.

At the onset of our investigation, we employed phenyl trifluoromethyl sulfone (PhSO₂CF₃, **1a**) as the CF₃ source. Into the mixture of CuCl (0.3 mmol) and ^tBuOK (2 equiv) in DMF at rt for 30 min, **1a** was added dropwise at the same temperature for 30 min under an argon atmosphere. As expected, a 17% yield of "CuCF₃" ($\delta = -24.1$ ppm) was detected by ¹⁹F NMR spectroscopy (Table 1, entry 1), which matched well with the previous reported results.^{6f,l,8e,9,10} The results obtained under various reaction conditions are listed in Table 1. The use of another nucleophilic base ^tBuONa gave a lower yield (Table 1, entry 2), while the addition of ^tBuOLi and MeONa did not give the "CuCF₃" species at all (Table 1, entries 3–4). There are two possible reasons. First of all, the formed dialkoxycuprate [K(DMF)][(^tBuO)₂Cu] (as reported by Grushin^{6f}) has a high nucleophilicity to activate **1a**. Second, the potassium ion plays a

Received: November 23, 2014

Table 1. Screening of Formation of “CuCF₃”^a

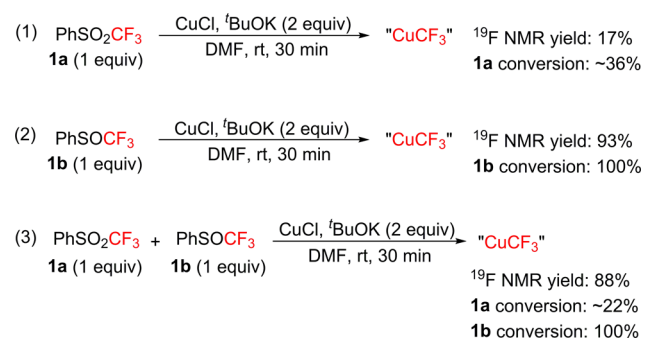
PhSO ₂ CF ₃ or PhSOCF ₃		CuCl, initiator		"CuCF ₃ "
1a	1b	solvent, rt		
entry	CF ₃ source	initiator (equiv)	solvent	"CuCF ₃ " ^b (%)
1	1a	^t BuONa (2)	DMF	17
2	1a	^t BuOK (2)	DMF	7
3	1a	^t BuOLi (3)	DMF	0
4	1a	MeONa (2)	DMF	0
5	1a	^t BuOK (3)	DMF	72
6	1a	^t BuOK (3.5)	DMF	56
7	1a	^t BuOK (4)	DMF	48
8 ^c	1a	^t BuOK (3)	DMF	52
9	1a	^t BuOK (3)	NMP	39
10	1a	^t BuOK (3)	DMPU	49
11	1a	^t BuOK (2)	DMSO	13
12	1b	^t BuOK (3)	DMF	50
13	1b	^t BuOK (2)	DMF	93
14 ^d	1b	^t BuOK (2)	DMF	96
15 ^e	1b	^t BuOK (2)	DMF	99

^aUnless otherwise noted, the reactions were performed by adding CuCl (0.3 mmol) and initiator into solvent (1 mL) at rt, and then 1a (entries 1–11) or 1b (entries 12–15) was added into the solution at the same temperature for 30 min under argon atmosphere. ^bYields were determined by ¹⁹F NMR spectroscopy using PhOCF₃ as an internal standard. ^cCuI (0.3 mmol) was used. ^dPhSOCF₃ (0.33 mmol) was used. ^ePhSOCF₃ (0.39 mmol) was used. “[Cu(CF₃)₂][−]” (10%, ¹⁹F NMR) was produced.

key role in the cupration reaction.¹⁵ When the amount of ^tBuOK was increased to 3 equiv, the formation of “CuCF₃” was enhanced to a 72% yield (Table 1, entry 5). However, a greater excess of ^tBuOK led to a lower yield because of some side reactions (Table 1, entries 6–7). Furthermore, the employment of CuI failed to improve the yield of “CuCF₃” (Table 1, entry 8). Other aprotic polar solvents were also tested, but with less success (Table 1, entries 9–11). It is worth noting that the formation of a small amount of CF₃H was always observed in these reactions,¹⁴ which may be caused by the moisture from solvent and/or reagent. This result indicates that CF₃[−] is produced in the process of generating “CuCF₃”.

To improve the yield of “CuCF₃”, phenyl trifluoromethyl sulfoxide (PhSOCF₃, 1b) was employed as the trifluoromethyl precursor. Disappointedly, only a 50% yield of “CuCF₃” was obtained under the optimized conditions on the basis of 1a (Table 1, entry 12). However, when 2 equiv of ^tBuOK were used, to our delight, the yield of “CuCF₃” was improved to 93% (Table 1, entry 13). As a result, the generation of CF₃H was also dramatically reduced. A higher yield of “CuCF₃” was obtained when the loading of 1b was increased to 1.1 equiv (Table 1, entry 14). It is noteworthy that a little excess of 1b results in a new species (−30.6 ppm in ¹⁹F NMR spectroscopy), which is assigned as “[Cu(CF₃)₂][−]” on the basis of the literature data.¹⁶ Consequently, when 1.3 equiv of 1b was used, the total yield of “CuCF₃” species was increased to 99% (Table 1, entry 15), with 89% contribution from L-CuCF₃ (−24.3 ppm in ¹⁹F NMR spectroscopy) and 10% from [Cu(CF₃)₂][−] (−30.6 ppm in ¹⁹F NMR spectroscopy).

These results indicate that PhSOCF₃ is an efficient precursor to generate the “CuCF₃” species, and the competing reaction to produce CF₃H can be inhibited. Furthermore, we also compared

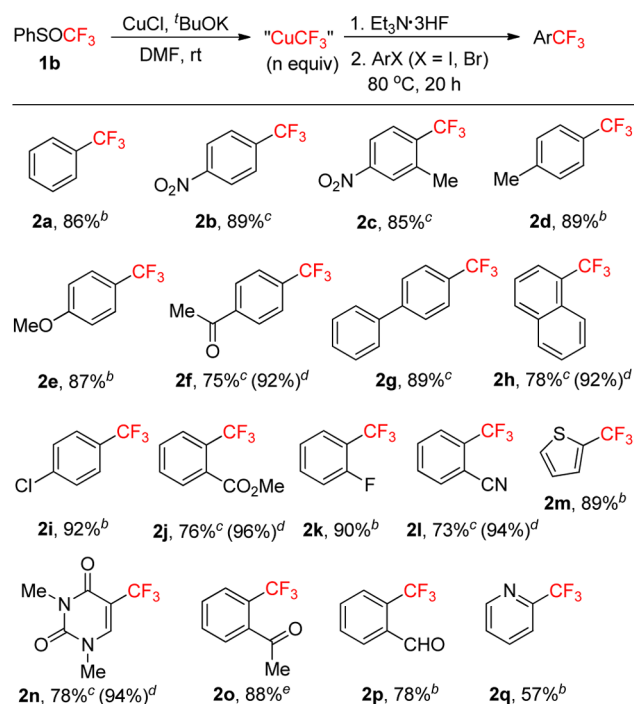
Scheme 2. Comparison of PhSO₂CF₃ and PhSOCF₃ in the Generation of “CuCF₃”

the reactivity of PhSOCF₃ and PhSO₂CF₃ in generating “CuCF₃” under the optimal conditions (Scheme 2). The results indicate that PhSOCF₃ is much more reactive than PhSO₂CF₃ to generate “CuCF₃”, and that the formation of “CuCF₃” generated from PhSOCF₃ is less than 30 min (see Supporting Information (SI), section 3 for details). It should be noted that *this is the first report that PhSOCF₃ is more efficient than PhSO₂CF₃ in the generation of the “CuCF₃” species*, which is in sharp contrast to the magnesium-mediated reductive trifluoromethylation of chlorosilanes.^{12a} The low reactivity of PhSO₂CF₃ for generating the “CuCF₃” species could be attributed to the steric hindrance of sulfone (compared to sulfoxide) during the nucleophilic attack of an alkoxide. Indeed, when we used PhSOCF₃ as the precursor to generate the “CuCF₃” species in the presence of ^tBuOK, benzenesulfinate (PhSO₂Bu) was produced quantitatively.

Next, we examined the stability of “CuCF₃” under various conditions. The “CuCF₃” species generated from PhSOCF₃ was gradually decomposed under air. Furthermore, the yield of “CuCF₃” decreased from 93% to 72% after 24 h under an argon atmosphere at rt. Similar to Grushin’s report,⁹ we found that the addition of Et₃N·3HF could stabilize the “CuCF₃” species generated from PhSOCF₃ (see SI, section 4.1), but the ¹⁹F NMR spectroscopy signal of the stabilized “CuCF₃” species shifted slightly from −24.5 to −26.5 ppm, and the signal of [Cu(CF₃)₂][−] at −30.6 ppm was increased.

With an efficient method for generating “CuCF₃” in hand (Table 1, entry 15), we further employed the “CuCF₃” species for the trifluoromethylation of various aryl halides. Initially, we attempted the reaction of iodobenzene with the stabilized trifluoromethylcopper at rt for 24 h. However, only a trace amount of trifluoromethylated product was detected by ¹⁹F NMR spectroscopy. When the reaction was carried out at 50 °C, to our delight, the product benzotrifluoride was formed in 50% yield after 28 h, but no obvious improvement was found when we prolonged the reaction time. Finally, the trifluoromethylation of iodobenzene proceeded smoothly at 80 °C under the standard conditions. Furthermore, it was found that the trifluoromethylation of aryl iodides proceeded well in the absence of any ligands, which is different from the previous reports.^{16b–d,17} The representative results are summarized in Scheme 3. Generally, both electron-rich and -deficient aryl iodides reacted well under the optimal conditions. For electron-deficient iodoarenes, nearly quantitative conversions (monitored by GC-MS) were achieved even within 16 h (Scheme 3, 2b, 2c, 2f, 2j, 2l). Ether, acetyl, halide, ester, and nitrile were tolerated under the standard reaction conditions. In addition, trifluoromethylated heteroarenes were also obtained in good yields (Scheme 3, 2m–2n). It is noted that thiophene was not observed by GC-MS in the

Scheme 3. Trifluoromethylation of Aryl Halides with “CuCF₃” Generated from PhSOCF₃^a



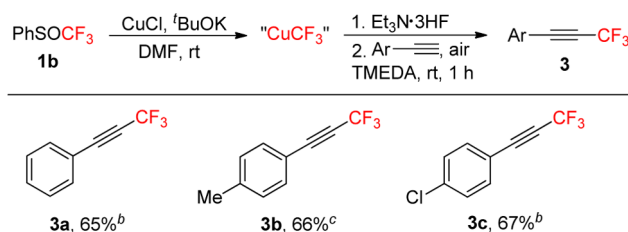
^aAll reactions were performed by adding ArX (X = I, Br) into the pregenerated “CuCF₃” species stabilized by Et₃N·3HF. For **2a**–**2n**, the substrates were ArI; for **2o**–**2q**, the substrates were ArBr. ^bReactions were performed on 0.1 mmol scale (n = 2) in DMF (1 mL) under an argon atmosphere. Yields were determined by ¹⁹F NMR spectroscopy using PhOCF₃ as an internal standard. ^cReactions were performed on 0.5 mmol scale (n = 1.5) in DMF (2.5 mL) under an argon atmosphere. Yields were of isolated products. ^dThe yields were determined by ¹⁹F NMR spectroscopy using PhOCF₃ as an internal standard. ^eThe reaction was performed on 0.1 mmol scale (n = 2) in DMF (1 mL) under an argon atmosphere, and the yield was determined by ¹⁹F NMR spectroscopy using PhCF₃ as an internal standard.

trifluoromethylation of 2-iodothiophene, which was shown as a side product reported by Grushin.⁹ Compared to iodoarenes, trifluoromethylation of bromoarenes with the copper reagent is known to be more challenging. In our case, only some electron-deficient bromoarenes reacted smoothly with the “CuCF₃” species to afford the trifluoromethylated products in moderate to good yields (Scheme 3, **2o**–**2q**).

The copper-mediated oxidative cross-coupling of trifluoromethylcopper with nucleophilic substrates have proven to be powerful trifluoromethylation strategies.¹⁸ We envisioned that the “CuCF₃” species generated from PhSOCF₃ may also work well under the standard conditions. Initially, we attempted the oxidative trifluoromethylation of terminal alkynes. We were delighted to find that the corresponding products were obtained in moderate yields when a 2-fold excess of the “CuCF₃” reagent was used (Scheme 4). It should be noted that, to inhibit the homocoupling of alkynes, alkynes were added slowly by a syringe pump to the “CuCF₃” species in DMF under an air atmosphere. Furthermore, the addition of tetramethylethylenediamine (TMEDA) as a ligand was also the key to enhancing product yields.

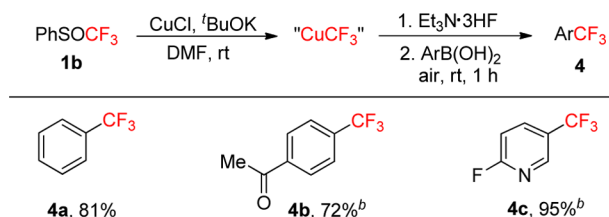
It was also encouraging to find that the trifluoromethylcopper reagent also reacted with arylboronic acids under an air

Scheme 4. Trifluoromethylation of Terminal Alkynes with “CuCF₃” Generated from PhSOCF₃^a



^aAll reactions were performed by adding alkyne (0.1 mmol) and TMEDA (2 equiv) into the pregenerated “CuCF₃” species (2 equiv) stabilized by Et₃N·3HF. ^bYields were determined by ¹⁹F NMR spectroscopy using PhCF₃ as an internal standard. ^cThe yield was determined by ¹⁹F NMR spectroscopy using PhOCF₃ as an internal standard.

Scheme 5. Trifluoromethylation of Arylboronic Acids with “CuCF₃” Generated from PhSOCF₃^a



^aAll reactions were performed by adding arylboronic acid (0.1 mmol) into the pregenerated “CuCF₃” species (2 equiv) stabilized by Et₃N·3HF. Yields were determined by ¹⁹F NMR spectroscopy using PhOCF₃ as an internal standard. ^bTMEDA (0.2 mmol) was used.

atmosphere at rt to give the trifluoromethylated products in up to 95% yield. Some representative results are summarized in Scheme 5.

In summary, we have shown the synthetic applications of phenyl trifluoromethyl sulfones and sulfoxides in copper-mediated trifluoromethylation and developed a new method for the synthesis of the “CuCF₃” species from PhSOCF₃. It is worth noting that PhSOCF₃ is more reactive than PhSO₂CF₃ in the formation of the “CuCF₃” species (up to 99% yield). The formed “CuCF₃” reagent has been found to be remarkably efficient in the trifluoromethylation of aryl iodides in the absence of additional ligands. Some activated aryl bromides can also be trifluoromethylated in moderate to good yields. Furthermore, the oxidative cross-coupling of terminal alkynes or arylboronic acids with the “CuCF₃” species has afforded the corresponding products in good yields.

■ ASSOCIATED CONTENT

Supporting Information

Experimental procedures and characterization data for products. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This work was supported by the National Basic Research Program of China (2015CB931900, 2012CB215500), National Natural Science Foundation of China (21421002, 21372246, 21102163, 21272069, and 20825209). We also thank Professor Shizheng Zhu (SIOC) for helpful discussions.

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