Photoinduced hydroxylperfluoroalkylation of styrenes†

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A transition metal- and organophotocatalyst-free hydroxylperfluoroalkylation of styrenes under visible-light irradiation was developed. In this protocol, an electron donor–acceptor complex (perfluoroalkyl iodide and tertiary amine) was employed for the introduction of a perfluorinated chain and molecular oxygen was used as a green oxidant for the generation of hydroxyl groups. Various styrenes successfully undergo the reaction affording the corresponding fluorine-containing alcohols in up to 96% yields.

Introduction

Over the past several years, impressive developments in the field of photoredox catalysis using visible light have occurred.[1] However, studies on the photochemistry of electron donor–acceptor (EDA) complexes are rare.[2] In some reactions, catalyst-free systems only lead to a slightly better result than that with a photocatalyst.[3] In the 1990s, we published several reports on the generation of fluoroalkyl and perfluoroaryl radicals from EDA complexes by UV irradiation.[4] UV light has a high energy, making it hard to control the chemical selectivity. On the other hand, visible-light is a viable alternative for designing a reaction with mild conditions and good functional group tolerance. Visible-light promoted fluoroalkylations without the use of a photocatalyst were reported by Yu in 2016 and later by Chen in 2017 for the synthesis of 2-fluoroalkylated quinoxalines, perfluoroalkyl-substituted phenanthridines, β-iodo-perfluoroalkylated alkenes/alkynes, and perfluoroalkylated arenes.[5] These pioneering reactions should encourage the further design of catalytic visible-light promoted reactions.

Difunctionalization of carbon–carbon double bonds is a useful strategy for organic synthesis, and there are numerous such reactions involving fluoroalkylation.[6] However, the hydroxylperfluoroalkylation of alkenes is challenging. Previous successful examples required expensive electrophilic trifluoromethylating reagents, toxic tin reagents, stoichiometric amounts of copper, or showed a lack of selectivity.[7] In 2017, Tsui reported the hydroxytrifluoromethylation of alkenes in the presence of bis(pinacolato)diboron (B_2Pin_2) using $CuCF_3$ derived from fluoroform in air.[7c] Recently, we disclosed a visible light-induced hydroxyltrifluoroethylation of styrenes with CF_3CH_2I and O_2 using $Ir(ppy)_3$ as a photocatalyst.[8] As part of our continued interest in the difunctionalization and fluoroalkylation of alkenes,[9] we applied perfluoroalkyl iodides as electron acceptors as well as fluoroalkylating reagents in photoredox reactions. We observed the "No Catalyst is Better" phenomenon which was mentioned several years ago by MacMillan,[3a] and has received more attention by Aggarwal and Wu very recently.[10] Although metal-mediated electron transfers and sulfinatodehalogenations are commonly used for the generation of perfluoroalkyl radicals, visible light-promoted reactions can occur under oxygen or air, thus facilitating the introduction of a hydroxy group, which is difficult to realize under other reaction conditions.[11,12] Herein, we report the hydroxylperfluoroalkylation of styrenes under visible light without the use of any photocatalyst or other additives besides amines under air.

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Results and discussion

The protocol was initially evaluated with styrene (**1a**) as a model substrate; it was reacted with 2 equivalents of n-C_4F_9I (**2**) and 2 equivalents of $^iPr_2NEt$ in various solvents under visible light irradiation with a 24 W fluorescent lamp under air (Table 1). Among the common solvents tested including acetonitrile, DMF, THF, and DMSO, all of them promoted the reac-

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Table 1 Screening of reaction conditions
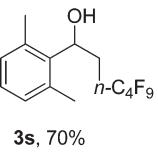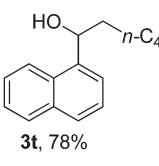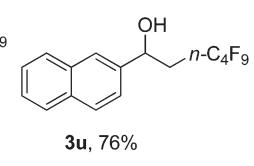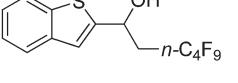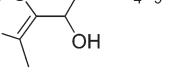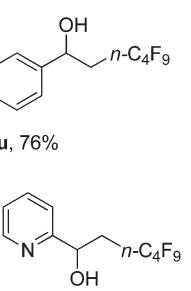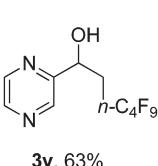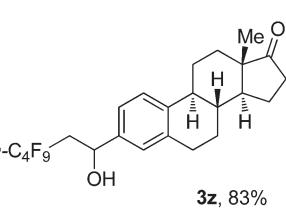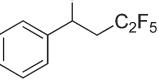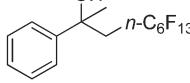
Entry[a]	Light	Solvent	Base	Yield[b] (%)
1	White light	MeCN	^{i}Pr$_2$NEt	48
2	White light	DMF	^{i}Pr$_2$NEt	60
3	White light	THF	^{i}Pr$_2$NEt	37
4	White light	DMSO	^{i}Pr$_2$NEt	75
5	Blue light	DMSO	^{i}Pr$_2$NEt	81
6	Green light	DMSO	^{i}Pr$_2$NEt	13
7	Blue light	DMSO	TMEDA	91
8[c]	Blue light	DMSO	TMEDA	99
9	Dark	DMSO	TMEDA	0
10	Blue light	DMSO	—	0
11[d]	Blue light	DMSO	TMEDA	0

[a] Reaction conditions: **1a** (0.2 mmol), **2** (0.4 mmol), base (0.4 mmol), anhydrous solvent (2 mL), visible light, 24 h, under air. [b] Yields were determined by ^{19}F NMR spectroscopy using benzotrifluoride as the internal standard. [c] DMSO (1 mL). [d] The reaction was carried out under a nitrogen atmosphere.

tion and DMSO was the best, giving product **3a** in 75% yield (Table 1, entries 1–4; detected by ^{19}F NMR spectroscopy). A blue light (15 W × 2) was applied, and gave the product in 81% yield, while irradiation with green light (15 W × 2) provided product **3a** in a low yield (13%) (Table 1, entries 5 and 6). To our delight, employing N,N,N',N'-tetramethylethane-1,2-diamine (TMEDA) as a base under irradiation with blue light provided **3a** in 91% yield and increasing the concentration from 0.1 M to 0.2 M enhanced the yield to 99% (Table 1, entries 7 and 8). Notably, **3a** could not be obtained in the dark, in the absence of the base, or under a nitrogen atmosphere (Table 1, entries 9–11).

With the optimized conditions in hand (Table 1, entry 8), we began to explore the substrate scope of the hydroxylper-fluoroalkylation. Inspiringly, a wide array of aromatic alkenes were subjected to the protocol and numerous functional groups were tolerated (Table 1). Styrenes bearing different halogens produced the corresponding products (**3b**, **3c**, **3d**, **3n**, **3o**, and **3q**) in good yields (from 85 to 92%). Styrenes with electron-donating substituents (**3e**, **3f**, **3g**, **3h**, **3p**, **3r**, and **3s**) afforded the corresponding products in moderate to excellent yields

Table 2 Substrate scope[a,b]

3a: R^1 = H, R^2 = H, 91% (84%)[e]
3b: R^1 = F, R^2 = H, 89%
3c: R^1 = Cl, R^2 = H, 90%
3d: R^1 = Br, R^2 = H, 92%
3e: R^1 = tBu, R^2 = H, 93%
3f: R^1 = OMe, R^2 = H, 88%
3g: R^1 = OAc, R^2 = H, 86%
3h: R^1 = CH$_2$OH, R^2 = H, 96%
3i: R^1 = CN, R^2 = H, 70%
3j: R^1 = H, R^2 = Me, 92%
3k: R^1 = F, R^2 = Me, 88%
3l: R^1 = Cl, R^2 = Me, 89%
3m: R^1 = H, R^2 = Ph, 93%

3n: R^3 = Cl, R^4 = H, 88%
3o: R^3 = Br, R^4 = H, 90%
3p: R^3 = Me, R^4 = H, 91%
3q: R^3 = H, R^4 = Br, 85%
3r: R^3 = H, R^4 = OMe, 89%

3s, 70%
3t, 78%
3u, 76%
3v, 87%
3w, 74%
3x, 68%
3y, 63%
3z, 83%
3aa, 90%[c]
3ab, 81%[d]

[a] Reaction conditions: **1** (1 mmol), **2** (2 mmol), TMEDA (2 mmol), anhydrous DMSO (5 mL), blue light (15 W × 2), 48 h, under an air atmosphere. [b] Isolated yield. [c] n-C$_4$F$_9$I was replaced by C$_2$F$_5$I. [d] Reaction with α-methylstyrene **1j** and n-C$_6$F$_{13}$I. [e] 5 mmol scale.

(from 70 to 96%). Some disubstituted alkenes underwent the desired reaction rendering the corresponding products (**3j**, **3k**, **3l**, and **3m**) effectively. Remarkably, styrenes with electron-withdrawing groups efficiently gave the corresponding products (**3i**) in 70% yield. In addition, this reaction could be applied to naphthalene and heterocyclic aryl alkenes such as vinylnaphthalenes (**3t** and **3u**), a benzothiophene derivative (**3v**), a thiazole derivative (**3w**), 2-vinylpyridine (**3x**), 2-vinylpyrazine (**3y**), and a steroid derivative (**3z**) to give the corresponding products in moderate to good yields (from 63 to 87%). A short or perfluorinated chain (C_2F_5 or n-C_6F_{13}) was successfully introduced in products **3aa** and **3ab**. The reaction could also be scaled up to 5 mmol to give 1.41 g (84%) of product **3a** in 72 h (Table 2).

Next, we evaluated the effect of a photocatalyst (Table 3). We chose two reaction conditions for comparison and compared the results with or without the photocatalyst. Condition A represents the optimized conditions (1 equivalent of **1a**, 2 equivalents of **2**, 2 equivalents of TMEDA, anhydrous DMSO, blue light (15 W × 2), 48 h, and under an air atmosphere). Under condition A, we found that the yield was more than 90% on 0.1 mmol, 0.5 mmol, and 1 mmol scales without the use of a photocatalyst (Table 3, entries 1–3). However, if the photocatalyst $Ir(ppy)_3$ was added to the reaction, interesting results were observed (Table 3, entries 4–6). On the 0.1 mmol scale, the yield did not change much (Table 3, entry 4 *vs.* entry 1). However, the yield decreased dramatically when the scale was 0.5 mmol (44% yield), and almost no product was detected in the 1 mmol scale reaction (Table 3, entries 5 and 6). In these experiments, we concluded that the photocatalyst had a negative effect on the visible light-induced hydroxylperfluoroalkylation reaction of styrenes, especially when the reaction was conducted on a large scale (1 mmol).

In order to further confirm the above conclusion, we carried out the reaction under condition B. Condition B was the optimal conditions in the presence of a photocatalyst (for the condition screening for reactions with a photocatalyst, see ESI Tables S3 and S4†). 2 equivalents of perfluorobutyl iodide

2 (to styrene **1a**) were used and 2 equivalents of Hünig's base were used. The reaction was carried out in DMSO under irradiation with a 24 W white fluorescent lamp under air (Table 3, entries 7–12). Very interestingly, when the reaction was carried out on the 0.1 mmol scale, both conditions (with or without catalyst) gave excellent yields of the products (95% in entry 7 *vs.* 97% in entry 10). Surprisingly, the yield decreased when the reaction was conducted on a larger scale (55% yield at 0.5 mmol scale in entry 11 or 43% yield at 1.0 mmol scale in entry 12) when the reaction was carried out in the presence of a photocatalyst. In contrast, the yields of the reaction did not fluctuate largely when the reaction was conducted on a larger scale without a catalyst (entries 8 and 9). Therefore, we explored the substrate scope using condition B on the 0.5 mmol scale (see ESI Table S5†). Most of the substrates gave the products in 40 to 85% yields. However, pure **3h** and **3i** could not be obtained under condition B. Therefore, when the reaction was conducted without a photocatalyst, it was more efficient and had a wider substrate scope as compared to that with a photocatalyst.

**Table 3** Comparison of reaction without or with a photocatalyst

$$1a + n\text{-}C_4F_9I \xrightarrow[\text{without or with photocatalyst}]{\text{visible light condition A or B}} 3a$$

Entry	Condition	Scale	Catalyst	Yield[a]
1	Condition A: blue light	0.1 mmol	—	99%[b]
2	TMEDA DMSO (c = 0.2 M)	0.5 mmol	—	90%
3		1 mmol	—	91%
4		0.1 mmol	$Ir(ppy)_3$ (1%)	94%[b]
5		0.5 mmol	$Ir(ppy)_3$ (1%)	41%
6		1 mmol	$Ir(ppy)_3$ (1%)	Trace[b]
7	Condition B: white light	0.1 mmol	—	95%[b]
8	^{i}Pr$_2$NEt DMSO (c = 0.1 M)	0.5 mmol	—	84%
9		1 mmol	—	83%
10		0.1 mmol	$Ir(ppy)_3$ (1%)	97%[b]
11		0.5 mmol	$Ir(ppy)_3$ (1%)	55%
12		1 mmol	$Ir(ppy)_3$ (1%)	43%

[a] Isolated yield. [b] Yields were determined by ^{19}F NMR spectroscopy using benzotrifluoride as the internal standard.

(a) Inhibition experiments

Additive	^{19}F NMR yield (%)
without additive	81
HO—⟨⟩—OH (1 equiv.)	52
NO_2—⟨⟩—NO_2 (1 equiv.)	0

(b) Proposed mechanism

Scheme 1 Mechanistic study of the visible light-induced hydroxylperfluoroalkylation of styrenes under air.

Next we concentrated on the reaction mechanism (Scheme 1). In the ^{19}F NMR spectra of pure perfluorobutyl iodide and its mixtures with TMEDA in DMSO, a 1.7 ppm downfield shift in the signal at around −70 ppm, which assigned for the terminal –CF$_2$I group of n-C$_4$F$_9$I, was observed. Consequently, we proposed that there was an interaction between the nitrogen of TMEDA and the iodine atom of n-C$_4$F$_9$I. Analogous chemical shift changes have been reported in other cases between fluoroalkyl iodides and nitrogen-containing molecules.[4,13] Notably, aromatic amines did not promote the reaction, but most tertiary aliphatic amines worked (see the ESI, Table S1†). On the other hand, the use of hydroquinone and 1,4-dinitrobenzene partially or completely inhibited the reaction (Scheme 1a). Based on the aforementioned phenomena, a mechanism is proposed in Scheme 1b. A perfluoroalkyl iodide (n-CF$_3$(CF$_2$)$_n$I n = 1, 3 or 5) and TMEDA would produce electron donor–acceptor complex **A** where the C–I bond is activated. **A** reacts with styrene subsequently under visible light irradiation generating benzyl radical **B**. The capture of O$_2$ from air by radical intermediate **B** gives peroxide radical **C**, which provides the final product **3**. To shed further light on the mechanism, we measured the quantum yields (Φ) of the reaction over 2 h. Quantum yields of 0.20% and 0.23% were observed, which suggested that a chain propagation pathway was unlikely.[14] A color change of the reaction was not observed when the reactants were mixed, which was also confirmed by UV-visible light analysis.

In summary, we developed a transition metal- and organophotocatalyst-free hydroxylperfluoroalkylation reaction that occurs under visible light irradiation. In this reaction, an electron donor–acceptor complex (R$_F$I-amine) acts as an inexpensive fluoroalkylating reagent and oxygen serves as an environmentally friendly oxidant. Various styrenes undergo the reaction to afford the corresponding products in up to 96% yields.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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