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Stereoselective Access to Trisubstituted Fluorinated Alkenyl Thioethers

Indira Fabre,^{a,b} Thomas Poisson,^c Xavier Pannecoucke,^c Isabelle Gillaizeau,^d Ilaria Ciofini,^{*b} Laurence Grimaud^{*a}

We report the first copper-catalyzed olefinic ethoxy carbonyl difluoromethylation of alkenyl thioethers *via* direct C-H bond functionalization using BrCF₂COOEt. The developed methodology allows the preparation of trisubstituted olefins with a controlled stereochemistry. A mechanistic study is reported and reveals a radical mechanism.

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1. INTRODUCTION

Since the last decade, organofluorine chemistry has known a very fast expansion since fluorinated molecules became popular in the quest for drugs and agrochemicals. This is reflected in the continuing increase of the number of fluorinated drugs already approved or drug candidates entering clinical trials.^{1–3} Indeed, the presence of a fluorine atom or a fluorinated group strongly affects the physical and chemical properties of a molecule. These features result from the particular properties of the fluorine atom, like its electronegativity, its small size and the strength of the carbon fluorine bond, for instance.⁴ In that context the development of new methodologies to access these strategic molecules is of high demand toward the discovery of new bioactive molecules. For this purpose, the direct C-H bond functionalization was recently recognized as an outstanding and efficient method for a straightforward access to the desired fluorine-containing molecules.⁵

Besides, sulfur-containing molecules are prevalent in a broad range of pharmaceuticals and natural products, and often display interesting various biological activities.⁶

As part of these sulfur-containing molecules, alkenyl thioethers⁷ are a versatile chemical platform and represent an interesting

motif to access complex sulfur-containing molecules. It is noteworthy, that Julia and coworkers developed the introduction of alkyl, allyl or silyl groups on 1-*tert*-butylthio-3-methoxy-1-alkenes using metallating agents (Scheme 1a).⁸ However, most efforts to functionalize this motif focused on the development of palladium-catalyzed arylation of alkenyl thioethers (Scheme 1b).^{9–11} Concerning the synthesis of fluorinated alkenyl mercaptans, Marquet and coworkers described an approach based on the anodic functionalization of alkenyl thioethers allowing the introduction of a fluorine atom at the α or β position (Scheme 1c).^{12,13} More recently, β -trifluoromethyl- α -functionalized-vinyl sulfides were synthesized under anionic conditions by Hanamoto and coworkers.^{14,15} As a complementary approach, Zard and coworkers reported the synthesis of tri- and tetra-substituted functionalized alkenyl thioethers by radical allylation, including examples bearing a CF₃ motif.¹⁶ The fluorination of thiophenes and benzothiophenes, which have an aromatic scaffold, have been reported with different metal catalysts (Ag,¹⁷ Pd,¹⁸ Fe¹⁹ and Ir²⁰ or Ru^{21,22} with light irradiation) and under photochemical conditions²³ (Scheme 1d). These reactions are selective for the C₂ position. To the best of our knowledge, no example of alkenyl thioether fluorination has been reported so far using functionalized fluorinated building blocks as the CF₂CO₂R group. Stimulated by our previous works dealing with the copper-catalyzed ethoxy carbonyl difluoromethylation of various scaffolds through a direct C-H functionalization,^{24–27} we report herein a successful application to the functionalization of alkenyl thioethers (Scheme 1e). In addition, significant insights regarding the mechanism of this reaction were also disclosed, demonstrating the existence of a radical pathway.

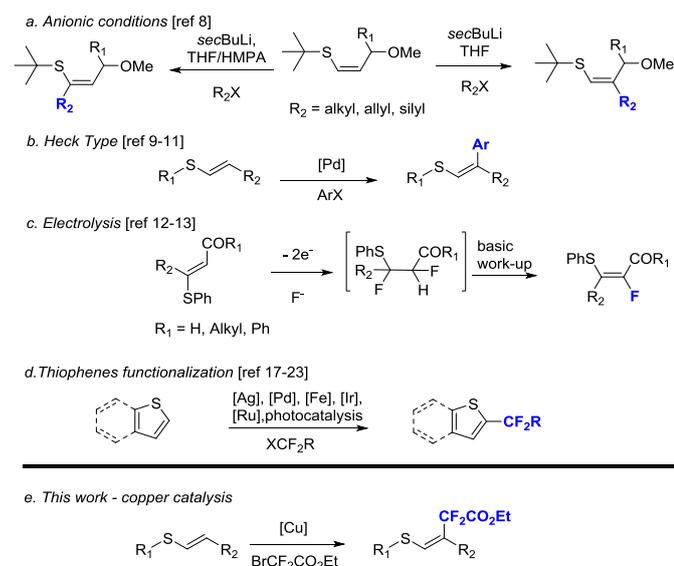
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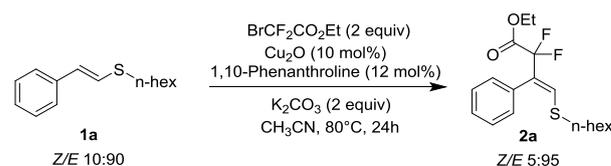
Scheme 1 Schematic illustration of previous and present works

2. RESULTS AND DISCUSSION

2.1 Optimization of the Reaction

The reaction conditions were optimized using *n*-hexyl(styryl)sulfane **1a** as a model substrate. Standard screening of solvent, catalysts, bases, ligands, temperature and reagent stoichiometry confirmed that the best yield is obtained using the conditions previously settled for the ethoxy carbonyl difluoromethylation of enamides,²⁶ with longer reaction time. A brief overview of the optimization of the reaction is displayed in Table 1 (for a more complete study, see Supporting Information). The desired product **2a** was isolated in 82% yield (Table 1, entry 1) when using Cu₂O (10 mol%), 1,10-phenanthroline (12 mol%) as a ligand in the presence of K₂CO₃ (2 equiv) and BrCF₂CO₂Et (2 equiv), in acetonitrile at 80 °C for 24 h. Several copper sources were also found to be efficient (Table 1, entries 2-4); however, the use of different ligands (Table 1, entries 5-8) or bases (Table 1, entries 9-13) gave very low to no conversion. Unfortunately, the catalyst loading could not be reduced since the yield significantly dropped when using 2 mol% of copper salt (Table 1, entry 14). Similarly, we could perform the reaction with 2 equiv of BrCF₂CO₂Et but lowering its amount (1.3 equiv) greatly impacted the reaction yield (Table 1, entry 15). The reaction was carried out under an air atmosphere, but with a lower yield (65%). Control experiments showed no reactivity in the absence of ligand, catalyst or base, and that no conversion occurred at room temperature. Under this optimized set of experimental conditions, the reaction turned out to be highly regio- and stereoselective. Indeed, the trisubstituted alkene **2a** was isolated as the major compound with a 5:95 *Z/E* ratio starting from alkenyl thioether **1a** as a *Z/E* (10:90). A NOE between the alkenyl proton and the alkyl chain on the sulfur atom allowed us to unambiguously confirm the regioselectivity of the reaction as shown in Table 2.

Table 1 Optimization of the reaction conditions



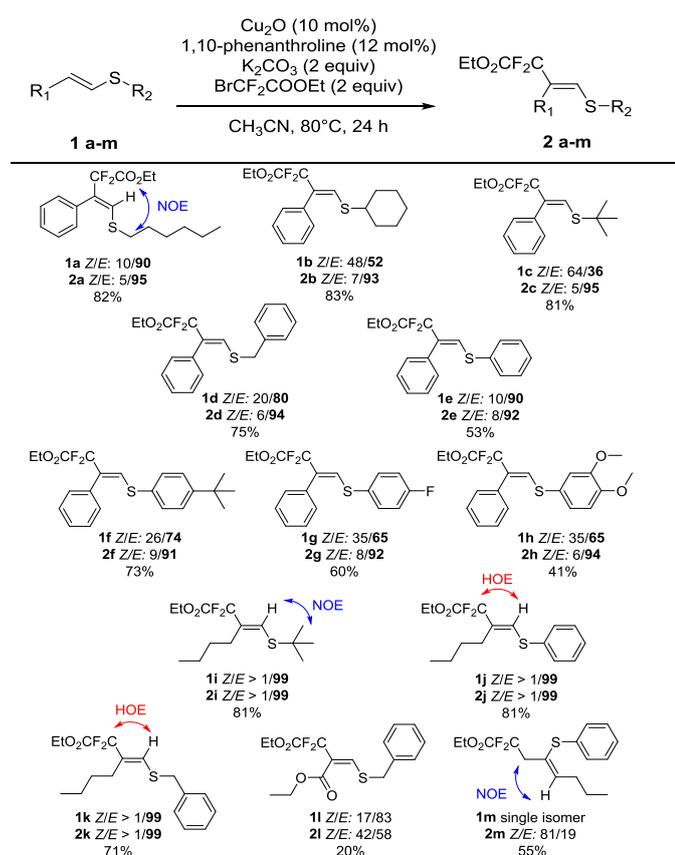
Entry	Deviation from the standard conditions	Yield
1	None	82 % ^a
2	Catalyst: Cu(OTf) ₂	60 % ^a
3	Catalyst: CuI	59 % ^a
4	Catalyst: Cu(CH ₃ CN) ₄ PF ₆	70 % ^a
5	Ligand: Neocuproine	0 %
6	Ligand: TMEDA	0 %
7	Ligand: TMHD	0 %
8	Ligand: 2,2'-bipyridine	35 % ^a
9	Base: Cs ₂ CO ₃	28 % ^b
10	Base: Na ₂ CO ₃	12 % ^b
11	Base: Et ₃ N	< 5 % ^b
12	Base: 2,6-lutidine	0 %
13	Base: <i>n</i> Bu ₄ NOAc	0 %
14	Catalyst: Cu ₂ O 2 mol%	24 % ^b
15	BrCF ₂ CO ₂ Et: 1.3 equiv	30 % ^b

Default catalyst, ligand and base are Cu₂O, 1,10-phenanthroline and K₂CO₃.^a Isolated yield. ^b ¹⁹F NMR yield, α,α,α-trifluorotoluene was used as an internal standard.

2.2 Scope of the Reaction

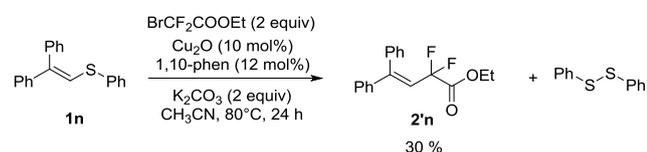
Having established the optimal reaction conditions, the scope of the reaction was further examined with a wide array of alkenyl thioether derivatives (see Table 2). Different styryl thioethers were successfully functionalized under these conditions. The reaction turned out to be of similar efficiency when performed with *S*-secondary (**2b**) or *S*-tertiary (**2c**) alkyl substituents as well as with a benzyl group (**2d**). Aryl styryl thioethers gave slightly lower yields of isolated adducts (**2e** to **2h**) whatever the nature of the substituent on the aromatic ring. When replacing the styryl moiety with an alkyl-substituted olefin, the reaction behaved similarly affording excellent yields of the trisubstituted alkenes **2i-2k**. However, the electron-deficient sulfanyl acrylate **1l** was poorly reactive giving **2l** in a modest yield (20%). When the *gem*-disubstituted alkenyl thioether **1m** was used, traces of the expected product were obtained (yield about 8%) along with the product **2m** (55% yield) resulting from a subsequent isomerization.

The reaction turned out to be highly regio- and stereoselective. Indeed, when starting with a nearly equimolar *Z/E* mixture of alkenyl thioether (eg. **1b** or **1c**), the trisubstituted alkenes **2b** and **2c** were isolated in a diastereomeric ratio > 90:10 in favor of the *E* isomer. Except for **2l**, no isomerization was observed when starting with pure *E* substrates as observed for **1i** and **1k**. NMR experiments were performed to determine the configuration of the trifunctionalized thioethers. Heteronuclear NOESY experiments clearly demonstrated a correlation between fluorine and the olefinic proton on **2j** and **2k**, enabling us to establish the *E* selectivity of the reaction.ⁱⁱ

Table 2 Scope of the copper-catalyzed ethoxy carbonyl difluoromethylation of various alkenyl thioethers **1**

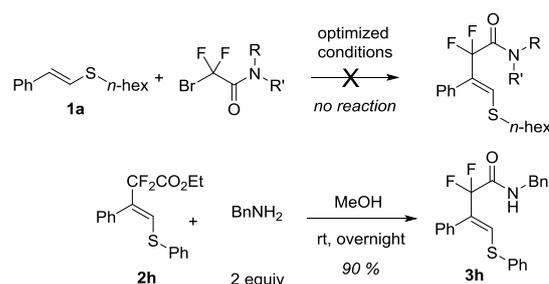
The major isomer is represented. NOE stands for Nuclear Overhauser Effect (between ^1H and ^1H) HOE stands for Heteronuclear Overhauser Effect (between ^{19}F and ^1H).

The regioselectivity of the reaction was further confirmed using the β -disubstituted styryl thioether **1n**. In this case, the compound **2'n** was isolated in a modest 30% yield along with diphenyldisulfide. The latter could result from an α -addition-desulfination sequence due to the steric hindrance at the β -position (see Scheme 2). It is worth noting that this byproduct has been observed in 0 to 15 % yield also with **1a** to **1l**.

**Scheme 2** Copper-catalyzed difluoroacetylation of **1n**

Surprisingly, the replacement of $\text{BrCF}_2\text{CO}_2\text{Et}$ with the bromodifluoroacetamide or bromodifluorophosphonate failed to give the corresponding fluorinated alkenyl thioethers. However, the post-functionalization of product **2h** turned out to be very efficient as shown in Scheme 3. The ester residue can be readily converted into the corresponding amide by treating compound **2e** with an

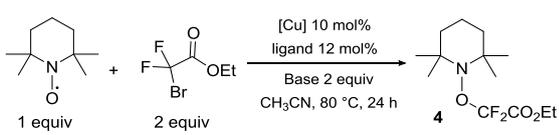
excess of benzylamine (Scheme 3). This transformation enlarges even more the molecular diversity accessible with this protocol.

**Scheme 3** Transformation of fluorinated alkenyl thioether **2h**

2.3 Mechanistic Studies

To get more insights into the mechanism, selected reactions were performed in the presence of radical scavengers.

When **1a** was reacted under standard reaction conditions in the presence of Galvinoxyl (1 equiv), no product was formed, while in the presence of TEMPO (1 equiv, Table 3, entry 1), traces of product were obtained along with 53 % of the TEMPO-adduct **4**, characterized both by ^{19}F NMR and Mass Spectroscopy. These observations along with the formation of diphenyldisulfide previously reported suggest that a radical mechanism is involved in the process. These conclusions are consistent with the recent studies of fluorination of hydrazones with analogous copper systems^{28,29} and with similar radical reactions mediated by copper.³⁰⁻³² To identify the key players in the radical formation, the trapping by TEMPO was further examined under various conditions (Table 3). The yield of **4** is only slightly lower (44 %, Table 1, entry 2) when the reaction is performed in the absence of the alkenyl thioether **1a**, suggesting that **1a** is not essential for the formation of the difluoroacetate radical. Note that in the absence of copper, no radical was generated (Table 3, entry 6). As already observed during the optimization process, tetrakis(acetonitrile)copper(I) hexafluorophosphate behaved similarly (Table 3, entry 4). This copper source allowed us to test the effect of the ligand toward the formation of **4**, indeed the use of Cu_2O without ligand was not conclusive due to solubility issues. In the absence of phenanthroline, no traces of **4** were detected (Table 3, entry 3), revealing its crucial role for the generation of the radical species, and explaining the failure of the reaction in the absence of ligand. Without a base the formation of **4** dropped significantly (Table 3, entry 5), unveiling its contribution in the generation of the reactive species. These experimental observations suggested that the combination of copper, 1,10-phenanthroline and K_2CO_3 might be responsible for the formation of a radical species in our reaction conditions.

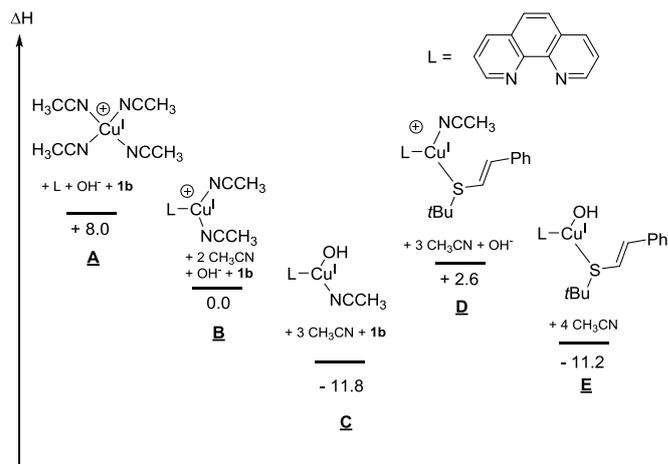
Table 3 Mechanistic experiments performed in the presence of TEMPO


Entry	[Cu]	Base	Ligand	Yield of 4
1 ^a	Cu ₂ O	K ₂ CO ₃	1,10-Phen	53 %
2	Cu ₂ O	K ₂ CO ₃	1,10-Phen	44 %
3 ^b	Cu(CH ₃ CN) ₄ PF ₆	K ₂ CO ₃	-	0 %
4	Cu(CH ₃ CN) ₄ PF ₆	K ₂ CO ₃	1,10-Phen	61 %
5	Cu ₂ O	-	1,10-Phen	15 %
6	-	K ₂ CO ₃	1,10-Phen	0 %

Yields were evaluated by ¹⁹F NMR using α,α,α -trifluorotoluene as an internal standard ^a In the presence of 1 equiv of **1a**, 2 % of **2a** was formed. ^b The copper source was changed for solubility issues in the absence of ligand.

For a better understanding of the elementary steps involved in this transformation, further investigations on the copper(I) complex involved in the reduction of BrCF₂CO₂Et were performed at the theoretical level using density functional theory (DFT). Bulk solvent effects were included by means of a polarizable continuum model (PCM) (see the Supporting Information for details). Two different mechanisms of activation of BrCF₂CO₂Et were compared: a single electron transfer (SET) and a halogen atom transfer (HAT).ⁱⁱⁱ Though the approach used here cannot provide a good description of the entropic effects that can play a role in the present case, the energetic –ie enthalpies– associated to these reaction can be trustfully estimated and it is here discussed. An estimate of the associated reaction free energies is reported in SI.

As starting complexes models, **A** to **E** were considered, where solvent molecules, alkenyl thioether **1b**, 1,10-phenanthroline and hydroxide anion (the base formed in the reaction medium) were included as possible ligands (see Scheme 4 and Table 4, details in the Supporting Information).^{iv}

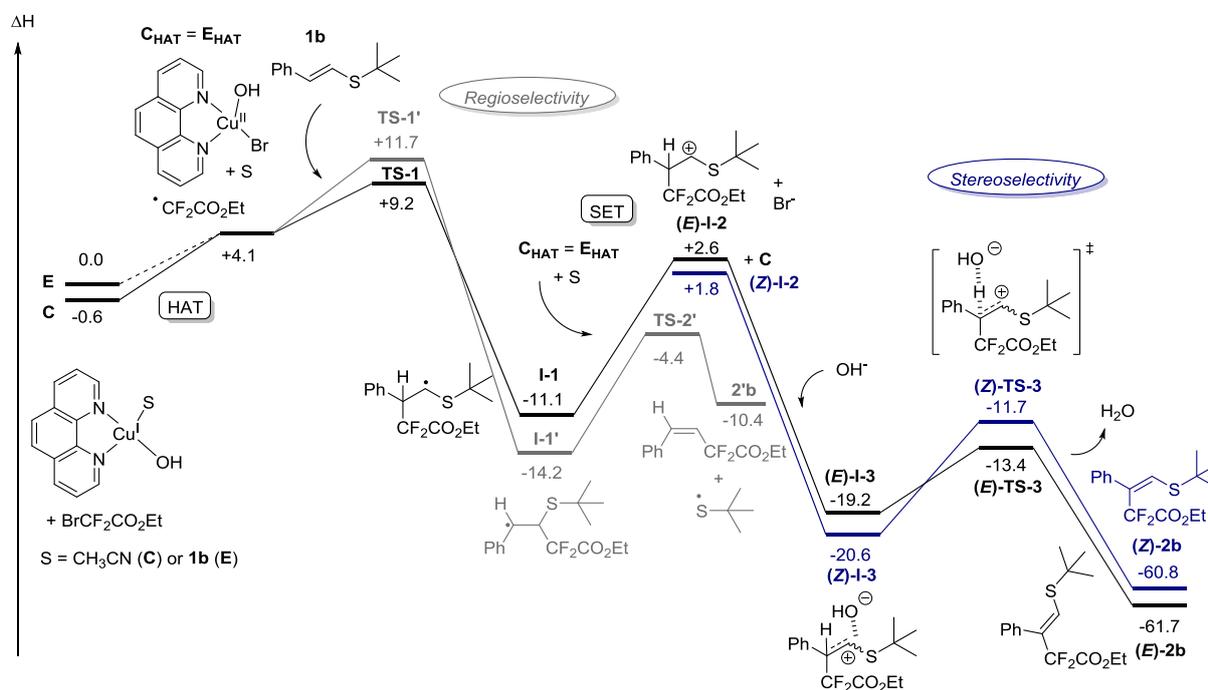
**Scheme 4** Relative stability of possible starting complexes computed (in kcal.mol⁻¹)**Table 4** Reaction energies for the different complexes (Enthalpy in kcal.mol⁻¹)

	A	B	C	D	E
SET	[Cu ^I] $\xrightarrow{\text{BrCF}_2\text{CO}_2\text{Et}}$ [Cu ^{II}] ⁺ + [BrCF ₂ CO ₂ Et] ⁻				
ΔE SET	55.2	39.9	11.7	39.1	16.7
HAT	[Cu ^I] $\xrightarrow{\text{BrCF}_2\text{CO}_2\text{Et}}$ [Cu ^{II}]-Br + \cdot CF ₂ CO ₂ Et				
ΔE HAT	-	28.4	4.7	25.8	4.1

Both SET and HAT were computed to be endothermic processes ($\Delta H > 0$). In the absence of the ligand (**A**), the energy required for the SET is very high ($\Delta H = 55.2$ kcal.mol⁻¹). This energy is lowered by complexation with 1,10-phenanthroline (**B**: $\Delta H = 39.9$ kcal.mol⁻¹ and **D**: $\Delta H = 39.1$ kcal.mol⁻¹), and even more favorable when copper is bound to a hydroxide anion (**C**: $\Delta H = 11.7$ kcal.mol⁻¹ and **E**: $\Delta H = 16.7$ kcal.mol⁻¹). The presence of alkenyl thioether **1b** has only a tiny effect, as one can see by comparing the energies computed for **B** to the ones for **D** and the energies computed for **C** to the ones for **E**. In all cases, the reaction energies for HAT reactions were lower than those calculated for SET. A similar behavior was observed whatever the presence of the alkenyl thioether; in the absence of phenanthroline, no radical was formed; and the absence of base significantly impacted the radical formation. These findings are consistent with experimental data from Table 3.

Overall, complexes **C** and **E** are the most likely to initiate the formation of the difluoroacetate radical by HAT (**C**: $\Delta H = 4.7$ kcal.mol⁻¹ and **E**: $\Delta H = 4.1$ kcal.mol⁻¹). In addition, the complexation of one molecule of **1a** to the copper/phenanthroline complex in the absence of base (complex **D**) was evidenced by ¹H NMR, but not in its presence (complex **E**, see the Supporting Information). After HAT, both **C** and **E** formed a copper (II) complex, bearing a phenanthroline

ligand, a hydroxide and a bromide. Surprisingly, the alkenyl thioether has low affinity for the copper complex and is expelled from the coordination sphere of copper at this step for **E**.

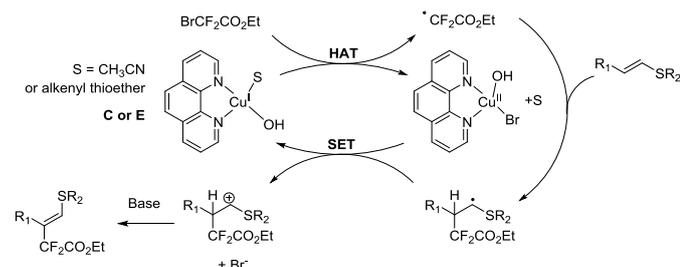


Scheme 5 Computed reaction pathways (in kcal.mol⁻¹). Solid line: initiation with **C**. Dashed line: initiation with **E**. The last part of the reaction path after regeneration of **E** by SET is omitted for clarity. This part is the same as for **C**. Grey: path for the formation of the main byproduct. Blue: path for the formation of the minor stereoisomer.

The formed ^oCF₂CO₂Et radical can then add to the alkenyl thioether **1b**. The addition at the α position with regards to the S atom (ΔH_a = 7.6 kcal.mol⁻¹, **TS-1'**) had a higher activation barrier than at the β position (ΔH_a = 5.1 kcal.mol⁻¹, **TS-1**), but appeared achievable under the reaction conditions (Scheme 5). This small difference allowed us to explain the observed regioselectivity in some cases with the formation of the disulfide as byproduct. The addition of the ^oCF₂CO₂Et radical on the alkenyl thioether **1b** on the copper (I) complex **D** has a higher energy barrier (ΔH_a = 8.2 kcal.mol⁻¹, see the Supporting Information), and is thus less favorable. The so formed radical **I-1** can reduce the copper (II) complex to regenerate copper (I) (ΔH = 12.9 kcal.mol⁻¹ to regenerate **C**, ΔH = 13.5 kcal.mol⁻¹ to regenerate **E**). The resulting carbocation **I-2** can then be deprotonated by the base to form the final product **2b**. The different conformations of the carbocation **I-2** are very close in energy (ΔH = 0.8 kcal.mol⁻¹) and are in equilibrium. These conformers are similar whatever the configuration of the starting alkenyl thioether (**Z** or **E**). Both can undergo a deprotonation step: after interaction with the base (OH⁻) leading to two stable and closely lying intermediates **(E)-I-3** and **(Z)-I-3** (ΔH = 1.4 kcal.mol⁻¹), the **E** and **Z** products can be formed. In agreement with the experiments, the transition state corresponding to the deprotonation step for the **E** form (that is **(E)-**

TS-3) and leading to the most stable **(E)-2b** product is computed to be lower in energy than the one of the corresponding **Z** form. Similar conclusions can be drawn by considering the values for the Gibbs free energies. The interaction with the base leads to the two stable intermediates **(E)-I-3** and **(Z)-I-3** (ΔG = 2.7 kcal.mol⁻¹). The transition state for the deprotonation leading to the major product **E** is the lowest in energy (ΔG = 1.7 kcal.mol⁻¹) and this major product is the most stable one (ΔG = 0.9 kcal.mol⁻¹). The complete Gibbs free energy profile can be found in Supporting Information.

Based on these findings, we suggested the mechanism depicted in Scheme 6. In the presence of 1,10-phenanthroline and a base, the copper(I) hydroxo complex reduced BrCF₂CO₂Et to form a copper (II) complex and an ethoxy carbonyl difluoromethyl radical. Then, the addition of this radical on the alkenyl thioether **1** proceeded according to an outsphere process and the observed stereoselectivity is related to the relative stability of the two possible isomers after the deprotonation step.



Scheme 6 Proposed reaction mechanism

These findings can be generalized. Indeed, for the initiation step, we could rationalize the absence of reactivity of $\text{BrCF}_2\text{CONR}_2$, $\text{BrCF}_2\text{PO}(\text{OEt})_2$ and $\text{BrCH}_2\text{CO}_2\text{Et}$ with the higher barrier energies computed for the HAT as well as with their higher reduction potential measured by cyclic voltammetry (see the Supporting Information). Moreover, the fact that the radical initiation did not depend on the presence of the alkenyl thioether showed that this method is broadly applicable to the generation of $^\circ\text{CF}_2\text{CO}_2\text{Et}$ radicals using copper catalyst. This hypothesis was confirmed by performing a radical trapping experiment with TEMPO in the presence of enamide (Table 5, entry 1) and benzofuranes (Table 5, entry 2 and 3) in previously reported conditions.^{25,26} For example the adduct **4** was isolated in 30% yields in the case of ene-carbamates without traces of the desired addition product, revealing a similar mechanism for these substrates. In a previous report,²⁶ a non-radical mechanism was proposed based on stoichiometric reactions performed in the absence of the base. After ruling out the direct oxidative addition of the bromo compound with the cationic copper(I) complex (by both NMR and electrochemistry performed at RT)²⁶, a pre-complexation of the latter with enamide was established via cyclic voltammetry as evidenced here by ^1H NMR in the case of the alkenyl thioether (in the absence of the base). This emphasized the important role of the base that transformed the cationic copper(I) complex to neutral copper(I) hydroxo complex. Finally, the radical mechanism established herein turned out to be quite general for these systems.

Table 5 Experiments in the presence of TEMPO for enamide and benzofuranes

Entry	Substrate	Solvent, time	Yield of 4 ^a
1		CH ₃ CN, 6 h	30 %
2		DMF, 20 h	10 %
3		DMF, 20 h	10 %

^a ^{19}F NMR yield, α,α,α -trifluorotoluene was used as an internal standard.

3. CONCLUSIONS

We report herein a general regio- and stereoselective method for the ethoxy carbonyl difluoromethylation of alkenyl thioethers. The developed methodology was applied to a broad range of substrates with good yields and an excellent selectivity in favor of the *E* isomer. Mechanistic studies supported by experimental observations and theoretical calculations gave evidences for a radical mechanism. The generation of a difluoroacetate radical promoted by a well-defined copper complex was established by these studies. Compared to other strategies to introduce a fluorine atom or a fluorinated group,^{19–21,23,33–39} this method allowed the generation of a radical under mild conditions, without requirement of light irradiation, expensive metal or radical initiator, which constitutes a real asset. It allows an easy access to fluorinated chemical platform of interest, since post-functionalizations of these substrates are possible thanks to the versatility of the ester function.

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Notes and references

- i The complete elucidation of the structure of the major isomer was established further by HOESY experiments.
 - ii All other configurations were assigned by analogy (see the Supporting Information)
 - iii For comparison purposes, a mechanistic path involving an oxidative addition (OA), which does not involve the generation of radicals, was also calculated and is reported in the Supporting Information.
 - iv We chose $\text{Cu}(\text{CH}_3\text{CN})_4\text{PF}_6$ as a simpler starting complex whose observed reactivity was similar to Cu_2O (see the Supporting Information and Table 3).
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