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Metal-Free Amidation of Ether sp^3 C–H Bonds with Sulfonamides using $\text{PhI}(\text{OAc})_2$

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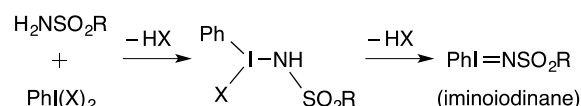
A selective protocol for the metal-free α -C–H amidation of ethers using sulfonamides and hypervalent iodine oxidants has been developed. The absence of precious metals and the conditions employed make the method environmentally attractive. A number of cyclic and acyclic, linear and branched ethers have been successfully amidated, and a broad sulfonamide scope has been demonstrated. Two unusual reactions, namely the amidation of an unactivated *tert*-butyl group and a tandem C–C coupling reaction, are also described.

Introduction

Direct amidation of sp^3 C–H bonds remains an important and challenging task with applications ranging from production of valuable organic targets to enhancing the utility of readily available but inert aliphatic substrates. The classical strategy for these atom-economical transformations involves transition metal catalysts that are able to mediate nitrene insertion into C–H bonds.^{1–11} The many pharmaceutical applications for oxidative coupling of C–H and N–H bonds require rigorous removal of trace amounts of these catalysts, which are often based on toxic and expensive precious metals. Eliminating the need for the metal catalysts is thus desirable from a standpoint of simplification and economy.^{12–14}

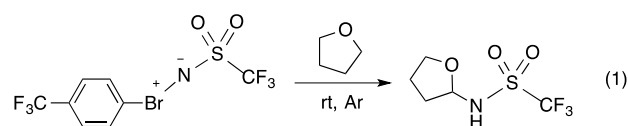
A variety of amine and oxidant combinations have been used for metal-free amination, amidation and imidation of C–H bonds. The oxidants in these systems include peroxides (*tert*-butyl hydroperoxide,^{15–18} di-*tert*-butyl peroxide,¹⁹ H_2O_2 ²⁰), benzoquinones,²¹ chloramine-T,²² and hypervalent halogen species.^{23–35} Additional organic catalysts (e.g., I_2 ^{15,24,29} and *n*- Bu_4NI)^{17,18} and stoichiometric additives (e.g., I_2 ,^{16,22,35–37} N-iodosuccinimide,^{16,20} $n\text{-Bu}_4\text{NI}$,¹⁶ KI ,^{16,28} TsOH ¹⁹ and AcOH ^{15,20}) are often involved and perform various functions. Hypervalent iodine species, particularly iodosobenzene derivatives such as $\text{PhI}(\text{OAc})_2$, are prevalent in both metal-catalyzed and metal-free processes for C–H amidation.^{1–10,23–32,35} Owing to their high

oxidation potential, these species are documented to activate sulfonamides for in situ generation of iminoiodinanes ($\text{PhI}=\text{NSO}_2\text{R}$), thus expanding the scope of the prior strategies based on preformed iminoiodinanes (Scheme 1).^{7–10}



Scheme 1 In situ formation of an iminoiodinane from a hypervalent iodine oxidant.

While a majority of the metal-free processes for C–H amidation and amination have been applied to oxidative coupling of N–H bonds to *sp* and sp^2 C–H bonds of unsaturated alkynes,³⁸ arenes,^{23,35,37,39,40} alkenes,^{25–27,41} and azoles,^{20,31,32,42,43} several examples of sp^3 C–H amidation of benzylic^{16–19,22–24,41} and allylic^{18,21,30} bonds have also been reported. In addition, Muñiz and coworkers described the amidation of an α -methyl group of a ketone.²⁵ Metal-free amidation is rare for sp^3 C–H bonds unactivated by adjacent double bonds. The I_2 -catalyzed oxidative coupling of purines with tetrahydrofuran (THF) oxidized by $\text{PhI}(\text{OAc})_2$ is known,²⁹ and a similar system utilizing $\text{PhI}(\text{OAc})_2$ as oxidant but involving stoichiometric I_2 has been reported for intramolecular amidation of alkyl sulfonamides.³⁶ Most prominently, Ochiai and coworkers discovered a highly active hypervalent bromine reagent, *p*-(CF_3)(Ph) BrNTf ($\text{Tf} = \text{CF}_3\text{SO}_2$), for α -C–H amidation of alkyl ethers (eq 1)³⁴ and also, in a striking example, for regioselective amidation of unfunctionalized alkanes.³³



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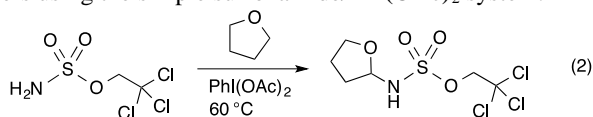
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[‡]Electronic Supplementary Information (ESI) available: experimental procedures, ¹H and ¹³C NMR data for all new compounds, additional optimization data, and crystallographic details for compounds **2a**, **2d**, **8** and **12**. See DOI: 10.1039/x0xx00000x

The present work began with an initial finding that reaction of 2,2,2-trichloroethoxysulfonamide in neat THF in the presence of $\text{PhI}(\text{OAc})_2$ selectively produces the α -C–H amidation product (eq 2). The simple sulfonamide/ $\text{PhI}(\text{OAc})_2$ system offers some advantages over the specialized p -(CF_3)(Ph)BrNTf reagent reported for amidation of alkyl ethers with sulfonamides.³⁴ The air- and moisture-stable and commercially available $\text{PhI}(\text{OAc})_2$ is more convenient than p -(CF_3)(Ph)BrNTf, which must be handled under inert atmosphere and whose two-step synthesis involves BrF_3 and liberates HF.^{41,44} Additionally, extending the reaction scope to new sulfonamides while using preformed reagents is limited by the necessity to synthesize a different reagent for each case. In this work, we sought to explore the potentially broader amide substrate scope offered for direct α -C–H amidation of alkyl ethers using the simple sulfonamide/ $\text{PhI}(\text{OAc})_2$ system.



Results and Discussion

We first optimized the experimental conditions for the α -CH amidation of THF using $\text{PhI}(\text{OAc})_2$ /2,2,2-trichloroethoxysulfonamide (**1a**) (Table 1). Although amidation takes place even at room temperature (entry 1), mild heating (60 °C, entry 4) leads to faster reactions and near-quantitative conversion of the sulphonamide to the functionalized compound **2a**. The first step in our screening was to test different oxidants. Whereas the hypervalent iodine oxidants $\text{PhI}(\text{OAc})_2$ and $\text{PhI}(\text{OPiv})_2$ (OPiv = pivalate) worked equally well and led to almost quantitative formation of **2a** (entry 6), the fluorinated $\text{PhI}(\text{TFA})_2$ (TFA = trifluoroacetate) yielded poor results (entry 7). We focused on $\text{PhI}(\text{OAc})_2$ due to its lower price and higher stability compared to its pivalate analogue. Although chloramine-T has been shown to be an excellent nitrene source for the α -CH amidation of ethers using a copper catalyst,⁴⁵ product formation was barely detected under our metal-free conditions (entry 8). Acetoxylation of activated C–H bonds has been previously identified as an undesired side reaction in the amidation of C–H bonds using $\text{PhI}(\text{OR})_2$ /sulfonamide systems.^{24,46} With the aim of avoiding this deleterious pathway and, at the same time, looking for cheaper and greener alternatives,^{47,48} we tested a series of iodine-containing salts^{49,50} (entries 9–12), although with unsuccessful results. In addition, the same product selectivity was achieved using this method in several common organic solvents but with diminished reaction rates (entries 13–16).

In terms of the sulfonamide scope, we obtained excellent yields with both aromatic and aliphatic sulfonamides (Table 2). This constitutes an important benefit compared to the previous metal-free amidation of ethers reported by Ochiai, which, despite its excellent activity, was limited to the highly reactive sulfonylimino- λ^3 -bromane.³⁴ In our system, α -CH amidation of THF takes place with excellent yields for substituted and unsubstituted aromatic sulfonamides (entries 1–5), although no conversion was detected in the case of the –OMe and –NH₂ derivatives (entries 6 and 7). Aliphatic sulfonamides also gave excellent conversions and high selectivity for the α -CH position (entries 8–10). The broad sulfonamide scope exhibited by our method is especially significant considering the useful biological and pharmacological properties of molecules based on the *N*-(methoxyalkyl)sulfonamide framework.^{51–54}

Table 1 Conditions screening for the amidation of THF.^a

| Entry | Oxidant | Solvent | T (°C) | t (h) | Yield (%) ^b |
|-------|------------------------------|-----------------------|--------|-------|------------------------|
| 1 | $\text{PhI}(\text{OAc})_2$ | neat | 25 | 10 | 53 |
| 2 | $\text{PhI}(\text{OAc})_2^c$ | neat | 60 | 2 | 77 |
| 3 | $\text{PhI}(\text{OAc})_2^d$ | neat | 60 | 2 | 89 |
| 4 | $\text{PhI}(\text{OAc})_2$ | neat | 60 | 2 | 91 |
| 5 | $\text{PhI}(\text{OAc})_2^e$ | neat | 60 | 2 | 94 |
| 6 | $\text{PhI}(\text{OPiv})_2$ | neat | 60 | 2 | 93 |
| 7 | $\text{PhI}(\text{TFA})_2$ | neat | 60 | 2 | 23 |
| 8 | chloramine-T ^f | neat | 60 | 2 | 3 ^g |
| 9 | NH_4IO_3 | neat | 60 | 2 | < 1 |
| 10 | KIO_3 | neat | 60 | 2 | < 1 |
| 11 | KIO_4 | neat | 60 | 2 | < 1 |
| 12 | $n\text{Bu}_4\text{NIO}_4$ | neat | 60 | 2 | < 1 |
| 13 | $\text{PhI}(\text{OAc})_2$ | DCE/THF (1:1) | 60 | 4 | 71 |
| 14 | $\text{PhI}(\text{OAc})_2$ | MeCN/THF (1:1) | 60 | 4 | 83 |
| 15 | $\text{PhI}(\text{OAc})_2$ | toluene/THF (1:1) | 60 | 4 | 42 |
| 16 | $\text{PhI}(\text{OAc})_2$ | cyclohexane/THF (1:1) | 60 | 4 | 56 |

^aConditions: **1a** (0.2 mmol, 46 mg), oxidant (0.4 mmol, 2 equiv), THF or mixed solvent (2 mL), reactions carried out under N₂ atmosphere. ^bYields determined by ¹H NMR spectroscopy using trimethoxybenzene as internal standard. ^c1 equiv of $\text{PhI}(\text{OAc})_2$. ^d1.5 equiv of $\text{PhI}(\text{OAc})_2$. ^e3 equiv of $\text{PhI}(\text{OAc})_2$. ^fChloramine-T was used instead of the pair **1a**/ $\text{PhI}(\text{OAc})_2$. ^gThe –NTs adduct was detected in this case.

Table 2 Sulfonamide scope on the α -CH amidation of THF.^a

| Entry | Substrate | –R | Yield (%) ^b |
|-------|-----------|--|------------------------|
| 1 | 1b | –C ₆ H ₅ | 89 |
| 2 | 1c | –C ₆ H ₄ (<i>p</i> -Me) | 90 |
| 3 | 1d | –C ₆ H ₄ (<i>p</i> -Cl) | 92 |
| 4 | 1e | –C ₆ H ₄ (<i>p</i> -Br) | 95 |
| 5 | 1f | –C ₆ H ₄ (<i>p</i> -COOH) | 71 |
| 6 | 1g | –C ₆ H ₄ (<i>p</i> -OMe) | < 1 |
| 7 | 1h | –C ₆ H ₄ (<i>p</i> -NH ₂) | < 1 |
| 8 | 1i | –(CH ₂)Ph | 92 |
| 9 | 1j | –CH ₃ | 89 |
| 10 | 1k | –CF ₃ | 96 |

^aConditions: **1b–k** (0.2 mmol), oxidant (0.3 mmol, 1.5 equiv), neat THF (2 mL), 60 °C, 2 h, reactions carried out under N₂ atmosphere. ^bYields were determined by ¹H NMR spectroscopy using trimethoxybenzene as internal standard.

The new functionalized THF compounds **2a-k** were characterized by ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectroscopy as well as high-resolution mass spectrometry (see Experimental Section). A characteristic ^1H NMR resonance in the range 5.20 and 5.42 ppm is assigned to the O-CH-NH proton and is common to all compounds **2a-k**; the corresponding $^{13}\text{C}\{^1\text{H}\}$ signal appears at *ca.* 85 ppm. The molecular structures of two of these species (**2a** and **2d**) were further confirmed by X-ray diffraction studies (Fig. 1). Their C–N bond distances are identical within the experimental error, with values of 1.472(8) Å and 1.470(4) Å for **2a** and **2d**, respectively, which are consistent with single bond character. Different rotameric orientations of the THF ring are present in each structure, likely due to packing forces.

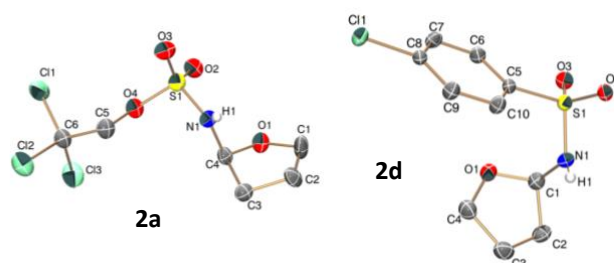


Fig. 1 ORTEPs of compounds **2a** and **2d**. 50% thermal ellipsoids are shown. Most hydrogen atoms have been omitted for clarity.

Table 3 α -CH amidation of ethers.^a

| $\text{R-O-CH}_2\text{-R}' + \text{H}_2\text{N-SO}_2\text{-O-CH}_2\text{-C(Cl)}_2\text{Cl} \xrightarrow[\text{MW, 120 }^\circ\text{C, 10 min}]{\text{PhI(OAc)}_2} \text{R-O-CH(R')-NH-SO}_2\text{-O-CH}_2\text{-C(Cl)}_2\text{Cl}$ | | | | | |
|--|-----------------------------|---------|-----------------------|------------------------|--------------------------|
| Entry | Substrate | Product | Additive (equiv.) | Yield (%) ^b | Selectivity ^c |
| 1 | | | - | 62 (56) | 0.82 (0.79) |
| 2 | | | I ₂ (0.02) | 59 | 0.76 |
| 3 | | | I ₂ (0.5) | 3 | 0.20 |
| 4 | Et-O-Et | | - | 60 (51) | 0.83 (0.78) |
| 5 | | | I ₂ (0.02) | 70 | 0.89 |
| 6 | $^n\text{Pr-O-}^n\text{Pr}$ | | - | 27 (27) | 0.49 (0.44) |
| 7 | | | I ₂ (0.02) | 68 | 0.84 |
| 8 | $^n\text{Bu-O-}^n\text{Bu}$ | | - | <1 | 0.10 |
| 9 | | | I ₂ (0.5) | 12 (12) | 0.21 (0.24) |
| 10 | $^i\text{Bu-O-}^i\text{Bu}$ | | - | 14 | - |
| 11 | | | I ₂ (0.02) | 26 | - |
| 12 | $^t\text{Bu-O-Me}$ | | - | 79 (85) | 0.98 (0.94) |
| 13 | | | I ₂ (0.02) | 60 | 0.83 |
| 14 | | | I ₂ (0.5) | 15 | 0.30 |
| 15 | $^t\text{Bu-O-Et}$ | | - | 14 (13) | 0.61 (0.54) |
| 16 | | | I ₂ (0.02) | 3 | 0.10 |
| 17 | Bz-O-Me | | - | 10, 1.5 ^d | - |
| 18 | | | - | 11, 1.9 ^d | - |
| 19 | Bz-O-Bz | | - | 10, 5.6 ^d | - |
| | | | | 11, 3.5 ^d | |

^aConditions: **1a** (0.5 mmol), PhIO(Ac)₂ (0.55 mmol, 1.1 equiv), neat ether (2 mL), microwave radiation (MW): 120 °C, 10 min. Reactions carried out under N₂ atmosphere. ^bYields were determined by ^1H NMR spectroscopy using trimethoxybenzene as internal standard. Values between parentheses correspond to yields obtained under identical conditions as for MW experiments but heating in conventional oil bath. ^cSelectivities were calculated for sulfonamide as the limiting reagent. ^dThese yields are calculated based on the ether (instead of the sulfonamide) as the limiting reagent, since the reaction also proceeds in the absence of **1a**.

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Under our optimized conditions (60 °C, 2 h, 1.5–2 equiv $\text{PhI}(\text{OAc})_2$), a series of aliphatic and aromatic, cyclic and acyclic ethers were tested as substrates for amidation with **1a**/ $\text{PhI}(\text{OAc})_2$; however, the results were unsatisfactory in terms of yields (from 5 to 10%) and selectivity, except for the cyclic tetrahydropyran (THP, 50% yield). Both the less polar character of acyclic ethers compared to THF and their slightly higher $\alpha\text{-C-H}$ bond dissociation energies⁵⁵ might contribute to the decreased activity. Further optimization of reaction conditions for diethyl ether (Table S1 in the Supporting Information) resulted in a maximum yield of 35% for the desired amidated product, but its formation was accompanied by a number of other unidentified species (60% overall sulfonamide conversion). Interestingly, we observed improved activity and considerably enhanced selectivity under microwave irradiation (MW, 120 °C, 10 min) in the neat ethers (Table 3, entries 4, 5; and Table S2). We carried out the same reactions in pressurized microwave vials but heated in a conventional oil bath (Table 3, yields in parentheses).^{56–58} The results were almost identical, suggesting a thermal origin for the improved yields and selectivity under microwave irradiation. For the sake of comparison, we undertook the $\alpha\text{-CH}$ amidation of THF under MW conditions and obtained yields comparable to those observed under mild thermal heating.

The addition of iodine in catalytic or stoichiometric amounts in conjunction with hypervalent iodine oxidants has been widely and successfully employed in other C–H amidation reactions.^{24,29,35–37} We observed an interesting effect with the addition of variable amounts of iodine. In the presence of catalytic amounts (2 mol% I_2 relative to sulfonamide) the yields for linear acyclic ethers increased considerably. For instance, the yield for di-*n*-butyl ether increased from 12 to 54% (entries 9 and 10), whereas that of di-*n*-propyl ether increased from 27 to 68% (entries 6 and 7). However, amidation of cyclic ethers (THF and THP) was not affected by the presence of trace amounts of iodine, which even had a deleterious effect in the case of *tert*-butyl ethers. In all cases, higher amounts of iodine (0.5 equiv) greatly suppressed product formation. In contrast, 0.5 equiv of I_2 was previously found to be the optimal loading for benzylic amidations by Fan and coworkers who obtained lower yields at both 1 equiv of I_2 and 0.05 to 0.2 equiv of I_2 .²⁴ These results highlight the importance of testing a wide range of I_2 concentrations in optimizing this type of reaction.

While linear cyclic and acyclic ethers gave the expected α -amidated products, both aromatic and branched aliphatic ethers exhibited a different reactivity. The branched di-*i*-propyl ether could not be amidated under our experimental conditions; instead we observed the formation of *iso*-propyl acetate (**7**) as the major product, due to direct acetoxylation, determined by ^1H NMR after addition of an authentic sample of **7**. In contrast, acetoxylation was barely detected with the other ether substrates. Surprisingly, reaction of *tert*-butyl methyl ether and *tert*-butyl ethyl ether gave completely different outcomes. The former yielded compound **8** where the *O*-methyl fragment is doubly functionalized and the *tert*-butoxy groups released. The molecular structure of **8** is consistent with the reduced number of resonances appearing in its ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra

and was confirmed by X-ray diffraction (Fig. 2). On the other hand, *tert*-butyl ethyl ether was converted into compound **9** after $\alpha\text{-CH}$ amidation of the ethyl group and unexpected $\beta\text{-CH}$ amidation of the *tert*-butyl fragment. This species was unambiguously characterized by ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectroscopy, as well as HRMS (FT-ICR). It constitutes a rare example of amidation of unactivated primary C–H bonds. A somewhat related reaction has been recently reported by Che and coworkers, where the C–H bond of a *tert*-butyl fragment was also aminated, although in low yield as a result of competition with the more reactive benzylic C–H bond.⁵⁹ Finally, and in contrast to the related copper-catalyzed process,⁶⁰ we observed that benzylic ethers cannot be amidated using the present method; instead they experience C–O cleavage and further oxidation to yield benzaldehyde and benzoic acid. The same transformation has previously been reported under somewhat related conditions and mainly investigated as a protocol for the synthesis of aldehydes from benzylic ethers.^{61–63} Although these reactions were accelerated by the addition of the sulfonamide and especially by the presence of the hypervalent iodine oxidant, they proceed even in the absence of these additives (see Supporting Information for more details, Table S3).

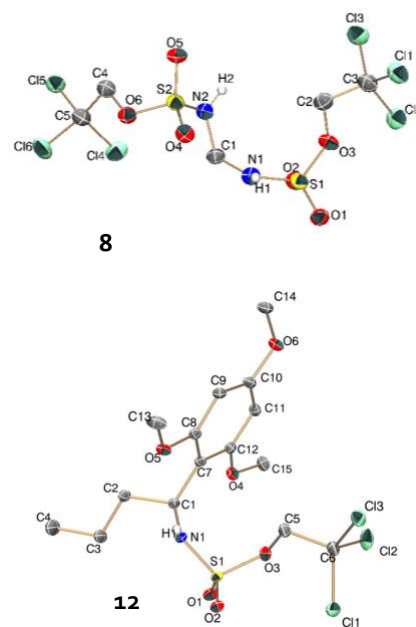
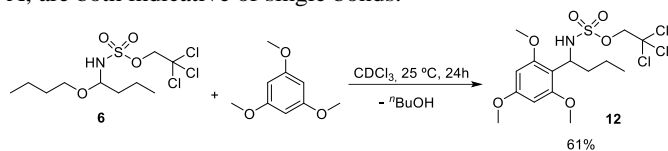


Fig. 2 ORTEPs of compounds **8** and **12**. 50% thermal ellipsoids are shown. Most hydrogen atoms have been omitted for clarity.

Purification of the functionalized ethers reported in this work proved difficult. In most cases extensive decomposition

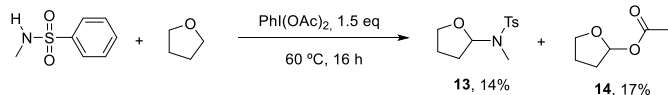
occurred during attempted column chromatography on both silica gel and alumina. In some cases degradation of the amidated compounds was observed after mild work-up or even upon standing in air for several days. The intrinsic reactivity of these species gave rise to an interesting derivative (**12**, Scheme 2), which is formed from Friedel-Crafts alkylation of electron-rich trimethoxybenzene (employed as an internal standard for ^1H NMR spectroscopy analysis). The Friedel-Crafts coupling of sulfonamides with aromatic compounds typically employs N-sulfonyl aldimines,⁶⁴⁻⁶⁷ but the direct use of N,O-acetals has been less explored. In an elegant study, Du Bois and co-workers developed a related rhodium-catalyzed one-pot amidation/nucleophilic addition process using ethers, sulphonamides and nucleophiles such as allylsilanes, silyl enol ethers, and ketene acetals.⁶⁸ Another interesting related process for the synthesis of 3-oxyindoles via tandem amidation/acetoxylation/Friedel-Crafts reaction has been recently described.⁶⁹ The molecular structure of **12** was elucidated by X-ray diffraction studies of suitable crystals grown by slow diffusion of pentane into a dichloromethane solution of the compound (Fig. 2). The length of the new C–C bond, 1.512(3) Å, and that of the adjacent C–N bond, 1.490(3) Å, are both indicative of single bonds.



Scheme 2 Formation of compound **12** by C–C coupling between **6** and trimethoxybenzene.

Attempts to perform amination reactions of primary and secondary alkyl amines with THF using $\text{PhI}(\text{OAc})_2$ led to acetoxylation product (compound **14** in Scheme 3) as evidenced by ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectroscopy in comparison with literature data.⁷⁰ This reaction is presumably base-mediated as the yields were dependent on the amount of amine present, and the same product could be formed using inorganic base K_2CO_3 in place of the amine.

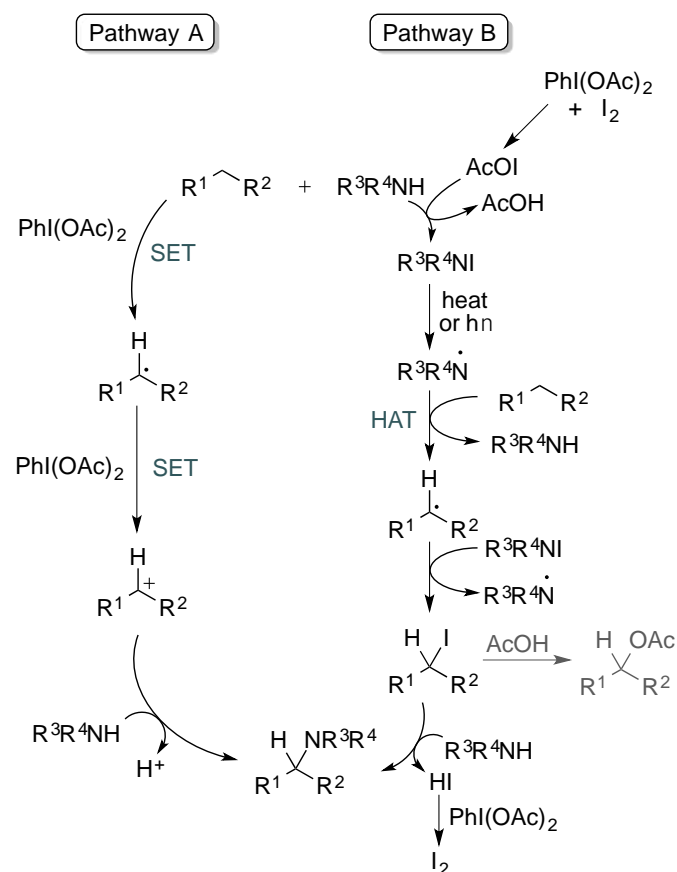
To investigate the feasibility of the classic mechanism for in situ nitrene formation (Scheme 1), which requires a primary sulfonamide,^{6,10,11} we tested the reactivity of methyl tosylamide with THF. Interestingly, we found that amidated product is formed in 14% yield along with 17% of the acetoxylation product (Scheme 3). Compound **13** displays ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra similar to a previously reported and related product derived from cyclohexyl tosylamide.⁷¹ This result suggests that in situ nitrene formation is not the only available pathway in the present system, and there is at least one other mechanism implicated that would account for the amidation using secondary sulfonamides.



Scheme 3 Amidation of THF with methyl tosylamide.

Interestingly, the amidation of THF by sulfonamide **1a** was completely shut down by addition of the radical trap TEMPO, suggesting a mechanism involving radicals. Two distinct radical-based mechanisms for $\text{PhI}(\text{OAc})_2$ -based systems have been proposed in the literature for oxidative C–N functionalization of sp^3 benzylic C–H bonds. In an account by

Cho and Chang involving imidation of benzylic C–H bonds using $\text{PhI}(\text{OAc})_2$ alone, a carbon-centered radical $\text{R}^1(\cdot\text{CH})\text{R}^2$ at the benzylic position was proposed from a single-electron transfer (SET) reaction between the benzylic substrate $\text{R}^1(\text{CH}_2)\text{R}^2$ ($\text{R}^1 = \text{Ar}$) and $\text{PhI}(\text{OAc})_2$ (Scheme 4, Pathway A).²³ A second SET to form a carbocation $\text{R}^1(\text{CH}^+)\text{R}^2$; followed by attack by a sulfonimide nitrogen and removal of a proton would then produce the final amidation product. Accounts describing amidation and amination of sp^3 carbons using $\text{PhI}(\text{OAc})_2/\text{I}_2$ systems are in general agreement in proposing mechanisms involving formation of a nitrogen-centered $\text{R}^3\text{R}^4\text{N}\cdot$ radical resulting from reaction of the oxidant/ I_2 combination with substrate either thermally or by photoactivation.^{24,29,36,37} Subsequent hydrogen atom transfer (HAT) produces a carbon-centered radical $\text{R}^1(\cdot\text{CH})\text{R}^2$, which abstracts iodine from $\text{R}^3\text{R}^4\text{NI}$ to propagate the production of the $\text{R}^3\text{R}^4\text{N}\cdot$ radical. Substitution of the $\text{R}^1(\text{CH})\text{R}^2\text{I}$ species with either $\text{R}^3\text{R}^4\text{NH}$ or AcOH can lead to formation of amination/amidation product or the undesired acetoxylation byproduct.



Scheme 4 Possible radical-based pathways for the amidation reaction based on literature reports.

In our system, a substantial KIE of $k_{\text{H/D}} = 9.3$ is observed for the reaction of **1a** in a 1:1 mixture of THF and THF- d_8 with 2 equiv of $\text{PhI}(\text{OAc})_2$. Under identical conditions after addition of 2 mol% I_2 , the $k_{\text{H/D}}$ is decreased to 2.2. The large difference in KIE between these two cases implies a change in mechanism which may involve a shift from Pathway A operative with $\text{PhI}(\text{OAc})_2$ alone to Pathway B upon addition of I_2 (Scheme 4). Future studies will focus on further elucidating the reaction mechanism(s) implicated and understanding the dependence of the role of I_2 on the identity of the ether substrate.

Conclusions

In summary, we have developed a metal-free protocol for the selective amidation of sp^3 C–H bonds in ethers using inexpensive and commercially available reagents. The resulting functionalized ethers are valuable nitrogen-containing compounds with potential biological and pharmacological properties. The reactions proceed without the need of precious-metal catalysts, with minimal use of organic solvents and under mild conditions. Higher reaction temperatures, achieved most conveniently by microwave irradiation, allowed for reaction times in the scale of several minutes, thus reducing energetic costs and making the process environmentally attractive. Preliminary studies suggest several radical mechanistic pathways in competition.

Acknowledgements

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