



 Cite this: *RSC Adv.*, 2021, **11**, 18966

 Received 14th December 2020  
 Accepted 19th May 2021

DOI: 10.1039/d0ra10517a

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# Visible-light enabled room-temperature dealkylative imidation of secondary and tertiary amines promoted by aerobic ruthenium catalysis†

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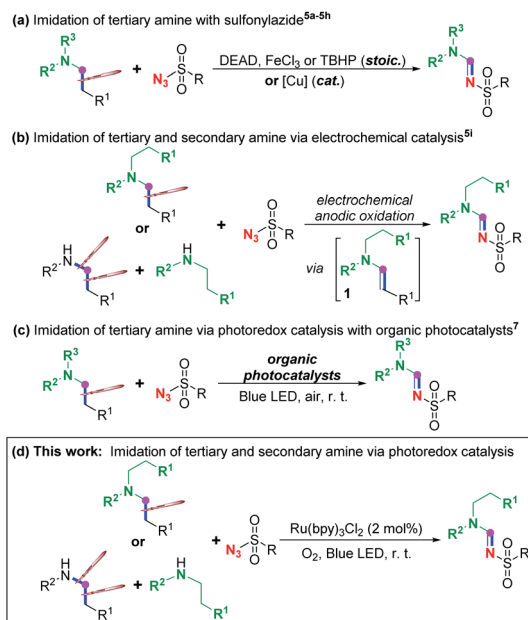
Employing sulfonyl azide as a nitrogen donor, a visible-light-enabled aerobic dealkylative imidation of tertiary and secondary amines involving C(sp<sup>3</sup>)–C(sp<sup>3</sup>) bond cleavage with moderate to excellent yields at room temperature is described. It has been demonstrated that this imidation could take place spontaneously upon visible-light irradiation, and could be facilitated considerably by a ruthenium photocatalyst and oxygen. An alternative mechanism to the previous aerobic photoredox pathway has also been proposed.

## Introduction

Imine<sup>1</sup> and, particularly, amidine<sup>2</sup> are among the most ubiquitous and pivotal moieties in miscellaneous ligands, organocatalysts, functional materials, synthetic building blocks, bioactive molecules and pharmaceuticals. Beside the most traditional approach through condensation between carbonyl and amine/amide to gain access to such significant scaffolds, various methodologies through transition-metal catalysed direct C–N cross-coupling of amine *via* C–H cleavage have been reported in the past decades.<sup>3</sup> Alternatively, it has also been widely reported that imine and imide could be acquired from azide instead of amine or amide.<sup>4</sup>

Among these methods with azide, imidation of the aliphatic chain of tertiary amine with sulfonyl azide through C(sp<sup>3</sup>)–C(sp<sup>3</sup>) bond cleavage (Scheme 1a)<sup>5</sup> has drawn our attention for C(sp<sup>3</sup>)–C(sp<sup>3</sup>) cleavage under mild conditions because it is generally challenging due to its high bond-dissociation energy and low polarity. In 2008, Li *et al.* disclosed a pioneering azo-mediated imidation of tertiary amines with sulfonyl azide at room temperature,<sup>5a</sup> followed by a couple of reports on the same transformation under mild conditions with other oxidants beside the azo compound, such as FeCl<sub>3</sub> (ref. 5b) and peroxides,<sup>5c</sup> or at high temperatures,<sup>5d</sup> while sulfamide could serve as a surrogate of sulfonyl azide in the presence of NBS<sup>5e</sup> or peroxide.<sup>5f</sup> Apart from these imidations through stoichiometric employment of oxidant, homogenous or heterogeneous copper-catalysed imidation of tertiary amine with sulfonyl azide has

also been reported.<sup>5g–j</sup> Meanwhile, such imidation could not only be achieved with tertiary amine *via* C–C cleavage as illustrated above, it can also be realized on secondary amine involving C(sp<sup>3</sup>)–N cleavage by electrochemical anodic oxidation (Scheme 1b),<sup>5k</sup> in which single electron transfer (SET) oxidation of tertiary or secondary amine to form nitrogen-centred radical cation has been proposed to initiate this imidation. Tertiary enamine **1** has been suggested to be the common intermediate for both tertiary and secondary amine, dividing the mechanism of these reaction into two processes:



Scheme 1 Imidation of amine with sulfonylazide.

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† Electronic supplementary information (ESI) available: Detailed experimental procedures, <sup>1</sup>H and <sup>13</sup>C NMR data and spectra. See DOI: 10.1039/d0ra10517a



a dehydrogenation process to give this enamine intermediate, and a 1,3-dipolar cycloaddition-decomposition process to afford the final amidine product.<sup>5</sup>

Our attention was then emphasized on other means beside electrochemical oxidation for the SET oxidation to generate nitrogen-centred radical cation that initiates such dealkylative imidation, and photoredox catalysis<sup>6</sup> which provides a potent synthetic strategy at room temperature and ambient pressure is considered to be the most efficient and green solution. Recently, Zeng<sup>7a</sup> and Pan<sup>7b</sup> disclosed two separate studies on visible-light-induced sulfonylimidation of tertiary amine using Eosin Y and acridinium salt, respectively, as organic photocatalysts in air (Scheme 1c),<sup>7</sup> based on an earlier report in which it had been proved that enamine **1** can be engendered from tertiary amine through photocatalysis.<sup>8</sup> However, employing secondary amine, such as diethylamine, instead of tertiary amine afforded sulfonylated diethylamine<sup>7a</sup> rather than the amidine product.<sup>5k</sup> Such sulfonyl azide participated imidation of secondary amine *via* photoredox catalysis is still unrealized.

Herein, as a continuation of our previous efforts on imidation<sup>9</sup> and other methodologies on the synthesis of nitrogen-containing compounds,<sup>10</sup> we would like to report our recent study on photoredox dealkylative imidation of tertiary and secondary amine with sulfonyl azide facilitated by aerobic ruthenium-catalysis to afford sulfonyl amidine at room temperature (Scheme 1d).

## Results and discussions

Our investigation commenced with the optimization of the reaction conditions employing triethylamine (TEA, **2a**) as the tertiary amine substrate, as summarized in Table 1.

Table 1 Optimization of the reaction conditions<sup>a</sup>

Entry	Changes to standard conditions <sup>a</sup>	Yield <sup>b</sup> (%)
1	None	91
2	In EtOAc (2 mL)	25
3	In DCE (2 mL)	73
4	In DME (2 mL)	76
5	In THF (2 mL)	70
6	In toluene (2 mL)	32
7	In MeNO <sub>2</sub> (2 mL)	25
8	In H <sub>2</sub> O (2 mL)	18
9	With Ir(ppy) <sub>3</sub> (2 mol%)	48
10	In open air	76
11	In Ar (1 atm)	49
12	No photocatalyst	29
13	No photocatalyst in Ar (1 atm)	27
14	No light	0

<sup>a</sup> Reaction conditions: TEA **2a** (0.5 mmol), TsN<sub>3</sub> (1.5 mmol), Ru(bpy)<sub>3</sub>Cl<sub>2</sub> (0.01 mmol) in 1,4-dioxane (2 mL) at room temperature in O<sub>2</sub> (1 atm) under 12 W blue LED irradiation for 6 hours. <sup>b</sup> Isolated yields of **4a** based on **2a**.

Table 2 Scope of the photoredox imidation of tertiary amine with azide<sup>a</sup>

Entry	2	R	4	Yield <sup>b</sup> (%)
1		Me ( <b>3a</b> )		91
2		H ( <b>3b</b> )		80
3		<sup>t</sup> Bu ( <b>3c</b> )		86
4		OMe ( <b>3d</b> )		67
5		F ( <b>3e</b> )		97
6		Cl ( <b>3f</b> )		82
7		CF <sub>3</sub> ( <b>3g</b> )		90
8		Me ( <b>3a</b> )		89
9		Me ( <b>3a</b> )		85
10		Me ( <b>3a</b> )		82
11		Me ( <b>3a</b> )		48
12		Me ( <b>3a</b> )		67
13		Me ( <b>3a</b> )		70

<sup>a</sup> Reaction conditions: see entry 1, Table 1. <sup>b</sup> Isolated yields of **4** based on **2**.



When **2a** and tosyl azide (**3a**) was subjected to the photocatalyst  $\text{Ru}(\text{bpy})_3\text{Cl}_2$  in 1,4-dioxane under  $\text{O}_2$  at room temperature, amidine **4a** was obtained in an excellent yield of 91% after 6 hours (entry 1). A series of widely used solvents were tested instead of 1,4-dioxane (entries 2–8), but none of them was found to provide **4a** in competent yields comparing to 1,4-dioxane. The yield of **4a** declined dramatically if the photocatalyst was replaced with  $\text{Ir}(\text{ppy})_3$  (entry 9). When the reaction was conducted in open air (entry 10), **4a** was acquired in a decreased yield of 76%, while the yield of **4a** further declined to 49% under argon (entry 11), demonstrating that the molecular oxygen could considerably promoted this imidation, although it is not prerequisite. Interestingly, the desired amidine product could still be generated, albeit in relatively low yields, in the absence of the photocatalyst either in  $\text{O}_2$  (entry 12) or under argon (entry 13), whereas no **4a** was engendered in the absence of LED irradiation even with the photocatalyst and  $\text{O}_2$  (entry 14), suggesting that this imidation with azide might spontaneously take place upon visible-light irradiation and neither photocatalyst nor molecular oxygen is prerequisite.

**Table 3** Scope of the photoredox imidation of secondary amine with azide<sup>a</sup>

Entry	2	4	Yield <sup>b</sup> (%)
1			92
2			86
3			90
4			54
5			48
6 <sup>c</sup>			61
7 <sup>c</sup>			72

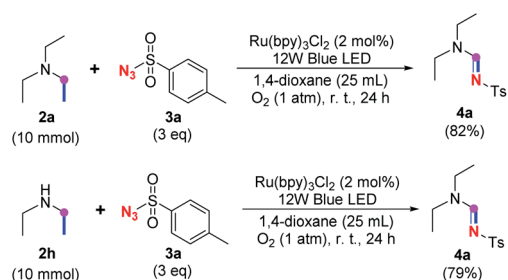
<sup>a</sup> Reaction conditions: see entry 1, Table 1. <sup>b</sup> Isolated yields of **4** according to their theoretical yields of 0.25 mmol from 0.5 mmol of **2h–2x**. <sup>c</sup> The reaction time was prolonged to 24 h.

Subsequently, the scope of amine **2** and sulfonyl azide **3** was explored, as shown in Table 2 and 3. With regard to the photoredox imidation of tertiary amine (Table 2), a series of benzenesulfonyl azides **3a–3g** with diverse *para*-substitutions on the benzene ring were tested under the optimized conditions (entries 1–7), and the excellent yields of corresponding amidines were generally maintained, except for amidine **4d** obtained from the electron-rich sulfonyl azide **3d** with strongly electron-donating methoxyl substituent. Reactions between other acyclic aliphatic tertiary amines and tosyl azide **3a** also provided amidines **4h–4j** in excellent yields, whereas dialkyl aniline **2e** afforded amidine **4k** in a moderate yield. In addition, two cyclic tertiary amines **2f** and **2g** could generate the desired product **4l** and **4m**, respectively, in synthetically acceptable yields.

Meanwhile, several secondary amines were investigated, as illustrated in Table 3. Since two molecules of secondary amine could provide only one molecule of the amidine product according to the mechanism proposed in previous work on the electrochemical imidation,<sup>5k</sup> the photoredox imidation of 0.5 mmol of the secondary amines could theoretically afford 0.25 mmol of corresponding amidines at most, and the yields in Table 3 were calculated accordingly. Results shown in Table 3 suggested that a series of secondary amines **2h–2n** were well tolerated with the optimized conditions established on tertiary amines, providing amidines **4a**, **4i**, **4k**, **4l** and **4n–4p** in moderate to excellent yields which are comparable to their tertiary amine siblings, although some of them required prolonged reaction time to guarantee the good yields. In addition, two gram-scale reactions with triethylamine (**2a**) and diethylamine (**2h**), respectively, under the optimized conditions have been conducted as shown in Scheme 2, and the good yields could be generally maintained.

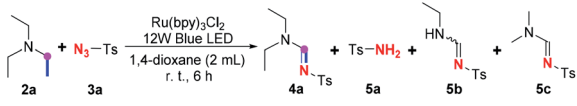
Furthermore, the spontaneity of this visible-light enabled imidation in the absence of oxygen and photocatalyst (entry 13, Table 1) attracted our attentions, for such observation suggested that an anaerobic pathways to realize this imidation probably coexist with the aerobic photoredox pathway proposed by Zeng and Pan,<sup>7</sup> where oxygen was proposed to serve a vital role in dehydrogenation. To gain more insights into the mechanistic picture, some control experiments were conducted, especially focusing on identification of byproducts generated along with the amidine product (Table 4).

It is discovered that sulfonamide **5a**, dealkylated<sup>11</sup> amidine **5b**, and amidine **5c** with an exchanged amino group<sup>12</sup> could be



**Scheme 2** Gram scale reactions.



Table 4 Control experiments on tertiary amine 2a<sup>a</sup>


Entry	O <sub>2</sub> /Ar (1 atm)	[Ru] (mol%)	Yield <sup>b</sup> (%)			
			4a	5a	5b	5c
1	O <sub>2</sub>	2	86	89	7	6
2	Ar	2	52	90	5	8
3	O <sub>2</sub>	0	25	45	6	0
4	Ar	0	26	49	5	0
5 <sup>c</sup>	O <sub>2</sub>	2	0	0	0	0
6 <sup>d</sup>	O <sub>2</sub>	2	0	0	0	0
7 <sup>e</sup>	O <sub>2</sub>	2	0	0	0	0

<sup>a</sup> Reaction conditions: TEA 2a (0.5 mmol), TsN<sub>3</sub> (1.5 mmol), Ru(bpy)<sub>3</sub>Cl<sub>2</sub> (0.01 or 0 mmol) in 1,4-dioxane (2 mL) at room temperature in O<sub>2</sub> or argon (1 atm) under 12 W blue LED irradiation for 6 hours. <sup>b</sup> Isolated yields of 4a and 5a–c based on 2a. <sup>c</sup> Without LED irradiation. <sup>d</sup> With TEMPO (2.5 mmol). <sup>e</sup> Without 2a.

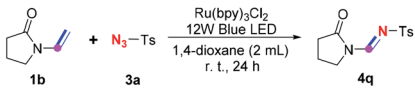
obtained under standard conditions (entry 1). However, when the imidation was conducted without oxygen, the yield of amidine 4a dramatically declined, whereas the yields of all of the three byproducts 5a–c generally maintained (entry 2). On the contrary, in the absence of the photocatalyst, 4a and 5a could only be acquired in very low yields regardless if the reactions were conducted in oxygen or argon, while no 5c was obtained, suggesting that the ruthenium catalyst is crucial to the generation of these compounds. Furthermore, no amidine 4a was engendered without visible-light irradiation (entry 5) or with TEMPO (entry 6), and neither were all of the three byproducts. A blank control experiment in the absence of amine 2a also suggested that none of these byproducts were converted from the solvent, 1,4-dioxane (entry 7).

Previous studies on the imidation of amine with sulfonyl azide have almost reached a consensus that essential measures or reagents, such as heating, external oxidant, transition-metal catalyst, electrochemical anodic oxidation and/or photoredox process with photocatalyst, are necessary to the dehydrogenation of amine to form the transient enamine intermediate 1.<sup>5,7</sup> Nevertheless, Zhu<sup>13</sup> has discovered that perfluoroalkanesulfonyl azide could readily decompose even at 0 °C in the presence of either tertiary or secondary amine *via* an intermolecular single electron transfer (SET) process between the azide and the amine, giving corresponding sulfonamide and *N*-substituted enamine equivalently. The formation of byproduct 5a, therefore, suggested that, merely induced by visible-light irradiation at room temperature (entry 4, Table 4), a similar intermolecular SET process between sulfonylazide and amine could take place spontaneously without any photocatalyst and oxidant. It has also been demonstrated that such SET process can be facilitated by the ruthenium catalyst without oxygen, due to the amount of 5a acquired in entry 2 that indicated most of 2a could be converted to the enamine intermediate 1a with the assistance of the ruthenium catalyst in the absence of oxygen. On the contrary, it

seems that oxygen cannot promote the spontaneous formation of enamine 1a (entry 3), and the previously proposed aerobic photocatalysis pathway<sup>7</sup> seems to contribute little in this ruthenium-promoted visible-light induced imidation, since the yield of 5a indicated that most of 1a was engendered through the above anaerobic SET process even in aerobic environment (entry 1). According to the above observations, it could be concluded that oxygen might be unnecessary to the conversion from amine 2a to enamine 1a, while it might enhance the following cycloaddition and ring-opening decomposition process, comparing the yields of 4a in entries 1 and 2.

To better understand the roles of visible-light, oxygen and the photocatalyst in the subsequent 1,3-dipolar cycloaddition and instantaneous ring-opening process to give the final product, a series of control experiments using a less active but stable and commercially available *N*-substituted enamine 1b as the substrate were performed (Table 5). Enamine 1b was converted to corresponding amidine product 4q in 31% yield under the standard conditions (entry 1), indicating that this visible-light enabled imidation is probably achieved through enamine intermediate as previous studies proposed.<sup>5,7</sup> The yield of 4q significantly declined when this transformation took place in the absence of oxygen (entry 2), confirming the crucial role of oxygen in the cycloaddition and decomposition process which is suggested by the results of control experiments shown in Table 4. However, only trace amount of amidine 4q could be observed on TLC when the reaction was conducted in the absence of the ruthenium photocatalyst or without visible-light irradiation (entries 3–6), demonstrating that the cycloaddition and decomposition process, although it might occur spontaneously with a extremely low rate (entry 6),<sup>14</sup> could be considerably facilitated by aerobic ruthenium photocatalysis (entry 1).

Beside the proposed aerobic photoredox mechanism,<sup>7</sup> a plausible alternative mechanism was therefore proposed according to the experimental observations discussed above and previously reported literatures.<sup>5,7,13,14</sup> This mechanism

Table 5 Control experiments on *N*-substituted enamine intermediate 1b<sup>a</sup>


Entry	O <sub>2</sub> /Ar (1 atm)	[Ru] (mol%)	LED irradiation	Yield <sup>b</sup> (%)
1	O <sub>2</sub>	2	Yes	31
2	Ar	2	Yes	11
3	O <sub>2</sub>	0	Yes	Trace
4	Ar	0	Yes	Trace
5	O <sub>2</sub>	2	No	Trace
6	Ar	0	No	Trace

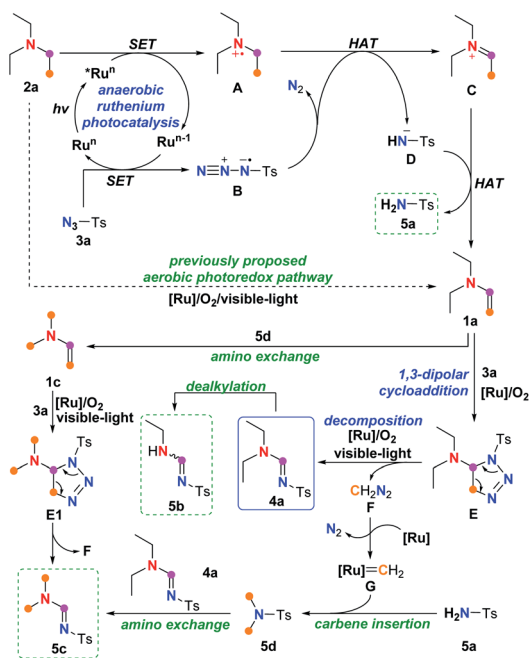
<sup>a</sup> Reaction conditions: 1b (0.5 mmol), 3a (1.5 mmol), Ru(bpy)<sub>3</sub>Cl<sub>2</sub> (0.01 or 0 mmol) in 1,4-dioxane (2 mL) at room temperature in O<sub>2</sub> or argon (1 atm) under 12 W blue LED irradiation or in dark for 24 hours. <sup>b</sup> Isolated yields of 4q based on 1b.



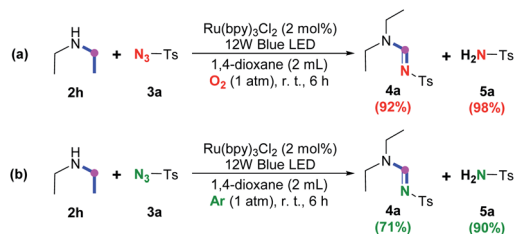
comprises two processes: the anaerobic ruthenium-promoted photoredox SET dehydrogenation process to form enamine intermediate; and the cycloaddition and ring-opening decomposition process facilitated by aerobic ruthenium photocatalysis to afford the final amidine product (Scheme 3).

Initially, SET oxidation of amine **2a** by the excited state  $^*Ru^{II}$  produced *via* photoexcitation of the higher valent ruthenium catalyst forms tertiary amine radical cation **A**,<sup>7</sup> along with the reduced ruthenium species  $Ru^{II}$ , which undergoes another SET process with the azide  $N_3-Ts$ , giving azide radical anion **B**<sup>5c,13</sup> and closing the ruthenium catalytic cycle. In the absence of the ruthenium catalyst, the above SET process from amine **2a** to sulfonyl azide could take place spontaneously upon visible-light irradiation (entry 4, Table 4), albeit with lower efficiency, and such automatic SET process could also be realized at high reaction temperature<sup>5c</sup> or with perfluoroalkanesulfonyl azide.<sup>13</sup> Then, hydrogen atom transfer (HAT) from the amine radical cation **A** to the azide radical anion **B**,<sup>13</sup> along with the loss of  $N_2$  from radical anion **B**, generates iminium **C**<sup>7</sup> and sulfonamide anion **D**<sup>5c,13</sup>. The iminium **C** would be deprotonated again by the sulfonamide anion **D** to provide the vital enamine intermediate **1a**, along with equivalent sulfonamide byproduct **5a**. Considering that approximately 90% of the starting amine **2a** was converted to **1a** and **5a** no matter if the reaction was conducted under aerobic or anaerobic conditions (entries 1 and 2, Table 4), the previously proposed aerobic photoredox pathway in which oxygen was required both in deprotonation of amine **2a** and in reoxidation of the organic photocatalysts,<sup>7</sup> might contribute little to this ruthenium-promoted visible-light enabled imidation.

Eventually, 1,3-dipolar addition of enamine **1a** with tosyl azide **3a** engenders a triazoline intermediate **E**, followed by



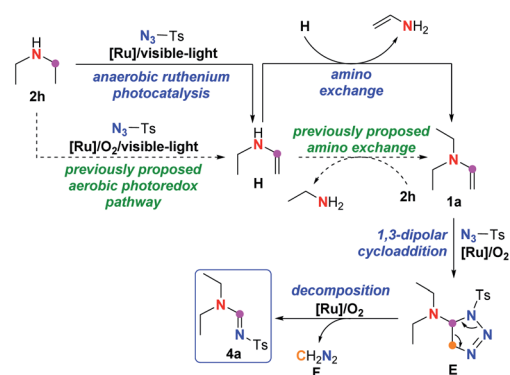
Scheme 3 Plausible alternative mechanism for imidation of tertiary amine beside the aerobic photoredox pathway.



Scheme 4 Control experiments on secondary amine **2h**.

decomposition of triazoline **E** to release diazomethane **F**,<sup>14</sup> along with the final amidine product **4a**, which might undergo oxidative dealkylation<sup>11</sup> to generate the de-ethylated byproduct **5b**. Results shown in Table 5 indicated that such cycloaddition-decomposition process, which might take place spontaneously albeit in very low efficiency (entry 6), could be promoted by aerobic ruthenium photocatalysis (entry 1). Additionally, in the presence of the ruthenium catalyst (entries 1 and 2, Table 4), diazomethane **F** could be converted to ruthenium carbene species **G**, which might be trapped by sulfonamide **5a** to form a *N,N*-dimethyl sulfonamide **5d** *via* N-H carbene insertion.<sup>15</sup> Amino group exchange<sup>12d-f</sup> between **5d** and the amidine product **4a** could directly afford the dimethyl amidine byproduct **5c**. Alternatively, such amino exchange might also take place on enamine **1a**,<sup>12a-c</sup> giving dimethyl enamine **1c** that also leads to **5c** through triazoline **E1** by cycloaddition-decomposition. The observation of byproduct **5c**, thus, could be considered as an indicator for the generation of diazomethane **F**.

With regard to secondary amine, control experiments on amine **2h** (Scheme 4) suggested that almost all of the secondary amine could be efficiently dehydrogenated through the anaerobic ruthenium promoted photoredox SET process either in the presence (Scheme 4a) or in the absence (Scheme 4b) of oxygen, indicating that the previously proposed aerobic photoredox pathway<sup>7</sup> also contributes little to the conversion of secondary amine to give corresponding enamine. Meanwhile, the yield of **4a** is higher in the presence of oxygen (Scheme 4a), demonstrating the significance of oxygen to the following cycloaddition-decomposition process as observed in the imidation of tertiary amine.



Scheme 5 Plausible mechanism for imidation of secondary amine.



The mechanism of imidation on secondary amine was therefore proposed as shown in Scheme 5. First, the secondary amine **2h** is converted to an enamine intermediate **H** through the same route proposed in Scheme 3. Unlike previous literature<sup>5d,5k</sup> in which the amino exchange between enamine **H** and amine **2h** was proposed to lead to the formation of enamine **1a**, it might be the amino exchange between two molecules of enamine **H** that is responsible for the generation of **1a**, since most of amine **2h** has been converted to enamine **H** either in aerobic or anaerobic environment according to the control experiments in Scheme 4, leaving little **2h** available for the following amino exchange. Finally, enamine **1a** undergoes the same cycloaddition-decomposition pathway shown in Scheme 3, leading to the amidine product **4a** though triazoline **E**, along with release of diazomethane **F**.

## Conclusions

To summarize, a photo-induced and aerobic ruthenium-promoted dealkylative imidation of tertiary and secondary amine with sulfonyl azide through C(sp<sup>3</sup>)-C(sp<sup>3</sup>) bond cleavage at room temperature has been developed, providing a concise and efficient route toward the significant amidine scaffold under mild conditions. This protocol avoids the employment of metal complexes as the catalyst described previously,<sup>5f</sup> though both reactions could be realized at room temperature. To our knowledge, this is the first example for photocatalysed imidation of secondary amine with azide. Furthermore, unlike previously proposed mechanism for visible-light-induced imidation with nonmetal organic photocatalysts, a plausible mechanism for this ruthenium photocatalyst participated imidation, featuring the anaerobic ruthenium-promoted photo-redox SET dehydrogenation of amine to form the key enamine intermediate and subsequent aerobic ruthenium photocatalysis facilitated cycloaddition-decomposition of this intermediate, have been proposed. Further studies to fully uncover the precise roles of visible-light, oxygen and photocatalyst, as well as application of this mild imidation to the preparation of amidine-based functional molecules, are undergoing.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

Financial support from the National Natural Science Foundation of China (No. 21801177), as well as the NMR technical assistance from the Engineering Experimental Teaching Centre, School of Chemical Engineering, Sichuan University, are highly appreciated. We also thank Zhong Shao and Fang Wang in this group for reproducing the results of **4b**, **4h**, **4i-l** and **4n-p**.

## Notes and references

- 1 (a) J. Brawley, E. Etter, D. Heredia, A. Intasiri, K. Nennecker, J. Smith, B. M. Welcome, R. K. Brizendine, T. W. Gould,

T. W. Bell and C. Cremona, Synthesis and evaluation of 4-hydroxycoumarin imines as inhibitors of class II myosins, *J. Med. Chem.*, 2020, **63**, 11131; (b) Z. Dong, Z. Wang, Z.-Q. Guo, S. Gong, T. Zhang, J. Liu, C. Luo, H. Jiang and C.-G. Yang, Structure-activity relationship of SPOP inhibitors against kidney cancer, *J. Med. Chem.*, 2020, **63**, 4849; (c) K. E. Archer, E. E. Reid, M. Shizuka, J. Woods, L. Harris, E. K. Maloney, L. M. Bartle, O. Ab, A. Wilhelm, Y. Setiady, J. F. Ponte, R. Singh, T. A. Keating, R. V. J. Chari and M. L. Miller, Synthesis of highly potent N-10 amino-linked DNA-alkylating indolinobenzodiazepine antibody-drug conjugates (ADCs), *ACS Med. Chem. Lett.*, 2019, **10**, 1211; (d) R. G. Almeida, W. O. Valença, L. G. Rosa, C. A. de Simone, S. L. de Castro, J. M. C. Barbosa, D. P. Pinheiro, C. R. K. Paier, G. G. C. de Carvalho, C. Pessoa, M. O. F. Goulart, A. Kharmá and E. N. da Silva Junior, Synthesis of quinone imine and sulphur-containing compounds with antitumor and trypanocidal activities: redox and biological implications, *RSC Med. Chem.*, 2020, **11**, 1145.

- 2 (a) R. G. Doveston, R. Steendam, S. Jones and R. J. K. Taylor, Total synthesis of an oxepine natural product, (±)-janoxepin, *Org. Lett.*, 2012, **14**, 1122; (b) M. Y. Lee, M. H. Kim, J. Kim, S. H. Kim, B. T. Kim, I. H. Jeong, S. Chang, S. H. Kim and S.-Y. Chang, Synthesis and SAR of sulfonyl- and phosphoryl amidine compounds as anti-resorptive agents, *Bioorg. Med. Chem. Lett.*, 2010, **20**, 541; (c) G. Brasche and S. L. Buchwald, C-H functionalization/C-N bond formation: copper-catalyzed synthesis of benzimidazoles from amidines, *Angew. Chem., Int. Ed.*, 2008, **47**, 1932; (d) Y.-F. Wang, X. Zhu and S. Chiba, Copper-catalyzed aerobic [3 + 2]-annulation of *N*-alkenyl amidines, *J. Am. Chem. Soc.*, 2012, **134**, 3679; (e) S. Li, Z. Li, Y. Yuan, Y. Li, L. Zhang and Y. Wu, Gold(I)-catalyzed aminohalogenation of fluorinated *N'*-aryl-*N*-propargyl amidines for the synthesis of imidazole derivatives under mild conditions, *Chem.-Eur. J.*, 2013, **19**, 1496; (f) S. H. Oakley, D. B. Soria, M. P. Coles and P. B. Hitchcock, Structural diversity in the coordination of amidines and guanidines to monovalent metal halides, *Dalton Trans.*, 2004, 537.
- 3 (a) J. Kim and S. S. Stahl, Cu-Catalyzed aerobic oxidative three-component coupling route to *N*-sulfonyl amidines via an ynamine intermediate, *J. Org. Chem.*, 2015, **80**, 2448; (b) I. Bae, H. Han and S. Chang, Highly efficient one-pot synthesis of *N*-sulfonylamidines by Cu-catalyzed three-component coupling of sulfonyl azide, alkyne, and amine, *J. Am. Chem. Soc.*, 2005, **127**, 2038; (c) J. Kim, S. Y. Lee, J. Lee, Y. Do and S. Chang, Synthetic utility of ammonium salts in a Cu-catalyzed three-component reaction as a facile coupling partner, *J. Org. Chem.*, 2008, **73**, 9454; (d) S. Chang, M. Lee, D. Y. Jung, E. J. Yoo, S. H. Cho and S. K. Han, Catalytic one-pot synthesis of cyclic amidines by virtue of tandem reactions involving intramolecular hydroamination under mild conditions, *J. Am. Chem. Soc.*, 2006, **128**, 12366; (e) Q. Gou, Z. Liu, T. Cao, X. Tan, W. Shi, M. Ran, F. Cheng and J. Qin, Copper-catalyzed coupling of sulfonamides with alkylamines: synthesis of (*E*)-*N*-



- sulfonylformamidines, *J. Org. Chem.*, 2020, **85**, 2092; (f) B. Huang, C. Yang, J. Zhou and W. Xia, Electrochemically generated *N*-iodoaminium species as key intermediates for selective methyl sulphonylimination of tertiary amines, *Chem. Commun.*, 2020, **56**, 5010.
- 4 (a) S. Y. Chow and L. R. Odell, Synthesis of *N*-sulfonyl amidines and acyl sulfonyl ureas from sulfonyl azides, carbon monoxide, and amides, *J. Org. Chem.*, 2017, **82**, 2515; (b) Z. Y. Gu, Y. Liu, F. Wang, X. Bao, S. Y. Wang and S. J. Ji, Cobalt(II)-catalyzed synthesis of sulfonyl guanidines via nitrene radical coupling with isonitriles: a combined experimental and computational study, *ACS Catal.*, 2017, **7**, 3893; (c) J. Y. Kim, S. H. Kim and S. Chang, Highly efficient synthesis of  $\alpha$ -amino amidines from ynamides by the Cu-catalyzed three-component coupling reactions, *Tetrahedron Lett.*, 2008, **49**, 1745.
- 5 (a) X. Xu, X. Li, L. Ma, N. Ye and B. Weng, An unexpected diethyl azodicarboxylate-promoted dehydrogenation of tertiaryamine and tandem reaction with sulfonyl azide, *J. Am. Chem. Soc.*, 2008, **130**, 14048; (b) S. Wang, Z. Wang and X. Zheng, Facile synthesis of sulfonylamidines via carbon-nitrogen bond formation mediated by FeCl<sub>3</sub>, *Chem. Commun.*, 2009, 7372; (c) A. Rouzi, R. Hudabaierdi and A. Wusiman, Synthesis of *N*-sulfonylformamidines by *tert*-butyl hydroperoxide-promoted, metal-free, direct oxidative dehydrogenation of aliphatic amines, *Tetrahedron*, 2018, **74**, 2475; (d) B. Kaboudin, S. Torabi, F. Kazemi and H. Aoyama, Transition metal- and catalyst-free one-pot green method for the synthesis of *N*-sulfonyl amidines via direct reaction of sulfonyl azides with amines, *RSC Adv.*, 2020, **10**, 26701; (e) J. Chen, Y.-P. Guo, M.-H. Sun, G.-T. Fan and L. Zhou, Bromoform reaction of tertiary amines with *N,N*-dibromosulfonamides or NBS/sulfonamides, *Chem. Commun.*, 2014, **50**, 12367; (f) Y. Zheng, J. Mao, J. Chen, G. Rong, D. Liu, H. Yan, Y. Chi and X. Xu, Unexpected C=N bond formation via NaI-catalyzed oxidative de-tetrahydrogenative cross-couplings between *N,N*-dimethyl aniline and sulfamides, *RSC Adv.*, 2015, **5**, 50113; (g) X. Xu, Z. Ge, D. Cheng, L. Ma, C. Lu, Q. Zhang, N. Yao and X. Li, CuCl/CCl<sub>4</sub>-Promoted convenient synthesis of sulfonyl amidines from tertiary amines and sulfonylazides, *Org. Lett.*, 2010, **12**, 897; (h) N. Liu, B.-Y. Tang, Y. Chen and L. He, Catalyzed imidation of tertiary amines by simple copper salts, *Eur. J. Org. Chem.*, 2009, 2059; (i) S. Shojaei, Z. Ghasemi and A. Shahrissa, Three-component synthesis of *N*-sulfonylformamidines in the presence of magnetic cellulose supported *N*-heterocyclic carbene-copper complex, as an efficient heterogeneous nanocatalyst, *Tetrahedron Lett.*, 2017, **58**, 3957; (j) W.-Z. Bi, W.-J. Zhang, Z.-J. Li, X.-Y. Xia, X.-L. Chen, L.-B. Qu and Y.-F. Zhao, Air-induced one-pot synthesis of *N*-sulfonylformamidines from sulfonyl chlorides, NaN<sub>3</sub>, and tertiary/secondary amines, *Eur. J. Org. Chem.*, 2019, 6071; (k) L. Zhang, J.-H. Su, S. Wang, C. Wan, Z. Zha, J. Du and Z. Wang, Direct electrochemical imidation of aliphatic amines via anodic oxidation, *Chem. Commun.*, 2011, **47**, 5488.
- 6 (a) D. M. Yan, J. R. Chen and W. J. Xiao, New roles for photoexcited Eosin Y in photochemical reactions, *Angew. Chem., Int. Ed.*, 2019, **58**, 378; (b) Q. Q. Zhou, Y. Q. Zou, L. Q. Lu and W. J. Xiao, Visible-light-induced organic photochemical reactions through energy-transfer pathways, *Angew. Chem., Int. Ed.*, 2019, **58**, 1586; (c) H. Jiang and A. Studer,  $\alpha$ -Aminoxy-acid-auxiliary-enabled intermolecular radical  $\gamma$ -C(sp<sup>3</sup>)-H functionalization of ketones, *Angew. Chem., Int. Ed.*, 2018, **57**, 1692; (d) X. Zhang and D. W. C. MacMillan, Direct aldehyde C-H arylation and alkylation via the combination of nickel, hydrogen atom transfer, and photoredox catalysis, *J. Am. Chem. Soc.*, 2017, **139**, 11353; (e) S. Mukherjee, R. A. Garza-Sanchez, A. Tlahuext-Aca and F. Glorius, Alkynylation of Cu<sub>sp2</sub>(O)-H bonds enabled by photoredox-mediated hydrogen-atom transfer, *Angew. Chem., Int. Ed.*, 2017, **56**, 14723; (f) T. Niwa, H. Yorimitsu and K. Oshima, Palladium-catalyzed 2-pyridylmethyl transfer from 2-(2-pyridyl)ethanol derivatives to organic halides by chelation-assisted cleavage of unstrained C<sub>sp3</sub>-C<sub>sp3</sub> bonds, *Angew. Chem., Int. Ed.*, 2007, **46**, 2643.
- 7 (a) J. Gui, H. Xie, H. Jiang and W. Zeng, Visible-light-mediated sulfonylimination of tertiary amines with sulfonylazides involving C<sub>sp3</sub>-C<sub>sp3</sub> bond cleavage, *Org. Lett.*, 2019, **21**, 2804; (b) R. Ding, H. Chen, Y.-L. Xu, H.-T. Tang, Y.-Y. Chen and Y.-M. Pan, Photoinduced cascade reaction of tertiary amines with sulfonyl azides: synthesis of amidine derivatives, *Adv. Synth. Catal.*, 2019, **361**, 3656.
- 8 G. Q. Xu, J. T. Xu, Z. T. Feng, H. Liang, Z. Y. Wang, Y. Qin and P. F. Xu, Dual C(sp<sup>3</sup>)-H bond functionalization of *N*-heterocycles through sequential visible-light photocatalyzed dehydrogenation/[2 + 2] cycloaddition reactions, *Angew. Chem., Int. Ed.*, 2018, **57**, 5110.
- 9 (a) Q. Miao, Z. Shao, C. Shi, L. Ma, F. Wang, R. Fu, H. Gao and Z. Li, Metal-free C-H amination of arene with *N*-fluorobenzenesulfonimide catalysed by nitroxyl radicals at room temperature, *Chem. Commun.*, 2019, **55**, 7331; (b) X. Wang, B. Lei, L. Ma, H. Jiao, W. Xing, J. Chen and Z. Li, Iron-catalyzed C(5)-H imidation of azole with *N*-fluorobenzenesulfonimide, *Adv. Synth. Catal.*, 2017, **359**, 4284; (c) B. Lei, X. Wang, L. Ma, Y. Li and Z. Li, NFSI-participated intermolecular aminoazidation of alkene through iron catalysis, *Org. Biomol. Chem.*, 2018, **16**, 3109; (d) B. Lei, Q. Miao, L. Ma, R. Fu, F. Hu, N. Ni and Z. Li, Efficient metal-free aminoiodination of alkenes with *N*-fluorobenzenesulfonimide under mild conditions, *Org. Biomol. Chem.*, 2019, **17**, 2126.
- 10 (a) Z. Li, X. Huang, F. Chen, C. Zhang, X. Wang and N. Jiao, Cu-Catalyzed concise synthesis of pyridines and 2-(1*H*)-pyridones from acetaldehydes and simple nitrogen donors, *Org. Lett.*, 2015, **17**, 584; (b) Z. Li, X. Wang, L. Ma and N. Jiao, Copper-catalyzed aerobic oxidation and oxygenation of anilines and acetaldehydes with dioxygen for the concise synthesis of 2-arylquinolines, *Synlett*, 2017, **28**, 1581; (c) C. Shi, Q. Miao, L. Ma, T. Lu, D. Yang, J. Chen and Z. Li, Room-temperature C-H bromination and iodination with sodium bromide and sodium iodide



- using *N*-fluorobenzenesulfonimide as an oxidant, *ChemistrySelect*, 2019, **4**, 6043; (d) Z. Li, L. Ma, J. Xu, L. Kong, X. Wu and H. Yao, Pd(II)-catalyzed direct C5-arylation of azole-4-carboxylates through double C–H bond cleavage, *Chem. Commun.*, 2012, **48**, 3763.
- 11 For dealkylation of tertiary amine, see: (a) S. Kim, J. W. Ginsbach, J. Y. Lee, R. L. Peterson, J. J. Liu, M. A. Siegler, A. A. Sarjeant, E. I. Solomon and K. D. Karlin, Amine oxidative *N*-dealkylation via cupric hydroperoxide Cu–OOH homolytic cleavage followed by site-specific Fenton chemistry, *J. Am. Chem. Soc.*, 2015, **137**, 2867; (b) C. X. Zhang, H.-C. Liang, E.-I. Kim, Q.-F. Gan, Z. Tyeklar, K.-C. Lam, A. L. Rheingold, S. Kaderli, A. D. Zuberbuhler and K. D. Karlin, Dioxygen mediated oxo-transfer to an amine and oxidative *N*-dealkylation chemistry with a dinuclear copper complex, *Chem. Commun.*, 2001, 631; (c) J. P. Ferris, R. D. Gerwe and G. R. Gapski, Detoxication mechanisms. III. Scope and mechanism of the iron-catalyzed dealkylation of tertiary amine oxides, *J. Org. Chem.*, 1968, **33**, 3493; (d) S. B. Karki, J. P. Dinnocenzo, J. P. Jones and K. R. Korzekwa, Mechanism of oxidative amine dealkylation of substituted *N,N*-dimethylanilines by cytochrome P-450: application of isotope effect profiles, *J. Am. Chem. Soc.*, 1995, **117**, 3657; (e) J. R. L. Smith and D. N. Mortimer, Model systems for cytochrome P450-dependent mono-oxygenases. Part 5. Amine oxidation. Part 17. Oxidative *N*-dealkylation of tertiary amines by metalloporphyrin-catalysed model systems for cytochrome P450 mono-oxygenases, *J. Chem. Soc., Perkin Trans. 2*, 1986, 1743; for dealkylation of imine, see: (f) D. E. Fogg and B. R. James, Net amine dealkylation at a diruthenium center: dehydrogenation of a secondary amine and hydrolysis of a coordinated imine, *Inorg. Chem.*, 1995, **34**, 2557.
- 12 For amino exchange occurring on enamine, see: (a) R. F. Abdulla, K. H. Fuhr and J. C. Williams Jr, New synthetic approaches to 4(1*H*)-pyridinone derivatives. 3. 2-Bromoacetylated enamines as pyridine ring synthons, *J. Org. Chem.*, 1979, **44**, 1349; (b) Y. Zhang, S. Xie, M. Yan and O. Ramstrom, Dynamic covalent chemistry of aldehyde enamines: Bi<sup>III</sup>- and Sc<sup>III</sup>-catalysis of amine–enamine exchange, *Chem.–Eur. J.*, 2017, **23**, 11908; (c) J. T. Gupton, N. Telang, X. Jia, B. C. Giglio, J. E. Eaton, P. J. Barelli, M. Hovaizi, K. E. Hall, R. S. Welden, M. J. Keough, E. F. Worrall, K. L. Finzel, E. J. Kluball, R. P. F. Kanters, T. M. Smith, S. Q. Smith, S. R. Nunes, M. T. Wright and J. M. Birnstihl, Further studies on vinamidinium salt amine exchange reactions, borohydride reductions, and subsequent transformations, *Tetrahedron*, 2010, **66**, 8485; for amino exchange occurring on amidine, see: (d) M. dF. Capela, N. J. Mosey, L. Xing, R. Wang and A. Petitjean, Amine exchange in formamidines: an experimental and theoretical study, *Chem.–Eur. J.*, 2011, **17**, 4598; (e) N. T. Tran and C. S. Cho, An efficient synthesis of benzimidazoles *via* palladium-catalyzed amine exchange reaction from trialkylamines to *o*-phenylenediamine in an aqueous medium, *Bull. Korean Chem. Soc.*, 2012, **33**, 4188; (f) D. D. Diaz, W. G. Lewis and M. G. Finn, Acid-mediated amine exchange of *N,N*-dimethylformamidines: preparation of electron-rich formamidines, *Synlett*, 2015, 2214.
- 13 Y. Xu and S. Zhu, The reaction of per(poly) fluoroalkanesulfonyl azides with tertiary and secondary amines: generation and trapping of enamines, *Tetrahedron*, 2001, **57**, 4337.
- 14 X. Zheng and J.-P. Wan, The C=C bond decomposition initiated by enamine–azide cycloaddition for catalyst- and additive-free synthesis of *N*-sulfonyl amidines, *Adv. Synth. Catal.*, 2019, **361**, 5690.
- 15 For carbene insertion into N–H bond, see: (a) D. Padin, J. A. Varela and C. Saa, Ruthenium-catalyzed tandem carbene/alkyne metathesis/*N*–H insertion: synthesis of benzofused six-membered azaheterocycles, *Org. Lett.*, 2020, **22**, 2621; (b) W.-S. Huang, Z. Xu, K.-F. Yang, L. Chen, Z.-J. Zheng and L.-W. Xu, Modular construction of multifunctional ligands for the enantioselective ruthenium-catalyzed carbenoid *N*–H insertion reaction: an enzyme-like and substrate-sensitive catalyst system, *RSC Adv.*, 2015, **5**, 46455; (c) L. Chen, H. Cui, Y. Wang, X. Liang, L. Zhang and C.-Y. Su, Carbene insertion into *N*–H bonds with size-selectivity induced by a microporous ruthenium–porphyrin metal–organic framework, *Dalton Trans.*, 2018, 47, 3940.

