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## Acridinium amidate as a hydrogen-bonding photocatalyst for direct decarboxylative alkylation of native carboxylic acids

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The direct generation of carbon radicals from carboxylic acids under visible-light photocatalysis offers an appealing strategy for molecular diversification. Here, we report a decarboxylative Giese-type addition using a zwitterionic acridinium amidate as a uniquely effective photocatalyst, which enables the efficient and selective activation and functionalization of a broad range of carboxylic acids without an external base or oxidant. The catalytic process relies on the formation of a hydrogen-bonded complex between the carboxylic acid and amidate, which triggers proton-coupled electron transfer upon photoexcitation to generate the corresponding carboxyl radical. This method featuring neutral conditions demonstrates excellent functional group tolerance and is applicable to structurally demanding substrates, including complex natural products and pharmaceutically relevant compounds.

### Introduction

Carboxy groups are fundamental functional groups that are prevalent in a range of organic molecules, including biomolecules, natural products, and pharmaceuticals. Because of its large dipole moment and ability to act as a hydrogen-bond donor and acceptor, the carboxy group plays a cardinal role in biochemical processes as well as in drug design as a constituent of pharmacophores.<sup>1</sup> Given that replacement of this functional group with other fragments can profoundly alter the physicochemical properties, it offers an effective strategy for modulating molecular behaviour for specific purposes, such as understanding biological mechanisms, upgrading natural products, and improving the efficacy of drug candidates. Accordingly, direct decarboxylative functionalization has garnered particular attention as an ideal means of fulfilling the synthetic demand in this pursuit.<sup>2,3</sup>

Photocatalysis has emerged as an enabling platform for decarboxylative transformations under mild conditions through the generation of carbon radicals from carboxylic acids. The methodologies developed for this purpose rely on one of the three distinct modes of activation: single-electron transfer

(SET),<sup>4–15</sup> ligand-to-metal charge transfer (LMCT),<sup>16–20</sup> and proton-coupled electron transfer (PCET) (Fig. 1A).<sup>21–41</sup> The SET-based approach typically involves oxidation of carboxylate ions, but the relatively high oxidation potential of these species (*ca.* +1.3 V *vs.* SCE) often results in poor chemoselectivity, especially in the presence of electron-rich functional groups. While LMCT allows for the direct generation of carboxyl radicals *via* the homolytic cleavage of photoexcited metal carboxylates, it generally requires a stoichiometric amount of base and is sensitive to polar or Lewis basic functionalities, which compromise both the efficiency and substrate scope of the reaction. In this regard, methods exploiting PCET obviate the use of an external base, and recent development of azaanthraquinone catalysts significantly expanded the scope of this approach.<sup>42</sup> In parallel with this progress, the effectiveness of photoinduced hydrogen atom transfer (HAT) catalysis with ketone derivatives has been explored for the direct radical generation from carboxylic acids.<sup>43</sup> Despite these significant advances in catalytic activation of carboxy groups, efficient and chemoselective decarboxylative functionalization in the presence of multiple polar and reactive functional groups, which are common features of natural products and therapeutic agents, remains a considerable challenge. Therefore, further development of photocatalysts that operate with high reactivity and chemoselectivity in such demanding settings is a highly desirable goal.

Recently, we introduced zwitterionic acridinium amidate **1a** as a novel organic molecular photocatalyst capable of facilitating HAT for the direct functionalization of structurally unbiased aliphatic C–H bonds (Fig. 1B).<sup>44</sup> The salient feature of this catalyst is that attractive noncovalent interaction between

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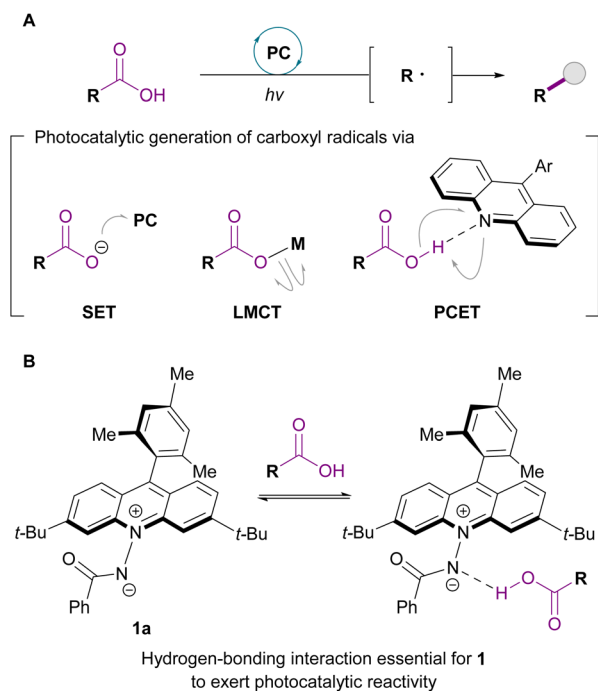


Fig. 1 Background and summary. (A) Representative strategy for photocatalytic activation of carboxylic acids. (B) Acridinium amidate 1 as a hydrogen-bonding photocatalyst.

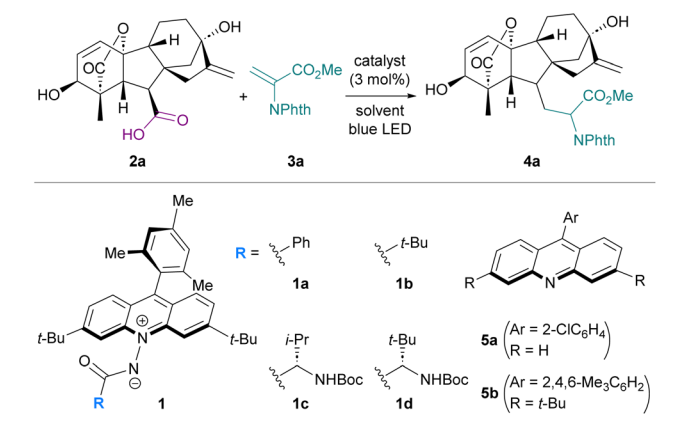
the pertinent hydrogen-bond donor and the amidate moiety is crucial for exerting productive photocatalytic reactivity. Considering the basicity of **1a** ( $pK_a$  of **1a**·H = 17.0 in acetonitrile (MeCN)), we reasoned that carboxylic acids could serve as hydrogen-bond donors, which would not only ensure the precise recognition of the carboxy group but also set a reactivity basis for the generation of the corresponding carbon radicals *via* PCET with **1** upon photoexcitation followed by decarboxylation, thereby allowing precisely chemoselective transformation. Herein, we report the decarboxylative Giese-type addition of carboxylic acids to electron-deficient olefins in the presence of acridinium amidate **1** as the sole catalyst under visible-light irradiation. This protocol enables the direct decarboxylative alkylation of a broad range of carboxylic acids, including those with polar and/or redox-sensitive functional groups, using an equimolar amount of olefins with high efficiency, demonstrating the vast synthetic utility of hydrogen-bond-induced photocatalysis of **1** as a practical tool for unlocking the long-standing limitations in carboxy group activation.

## Results and discussion

To assess the feasibility of carboxy group-targeted photocatalytic functionalization by harnessing the characteristic attributes of **1**, we chose gibberellic acid **A<sub>3</sub>** (**2a**), an important plant hormone with an *ent*-gibberellane skeleton decorated with a carboxy group, two hydroxy groups, internal and terminal alkenes, and a lactone as a model substrate. With *N*-phthaloyl dehydroalanine ester **3a** as the radical acceptor, our

investigation was initiated by subjecting a mixture of **2a**, **3a**, and *N*-benzoyl acridinium amidate **1a** (3 mol%) in 1,2-dichloroethane (DCE) to blue-light irradiation for 6 h, affording the desired product **4a** in 13% nuclear magnetic resonance (NMR) yield as a mixture of diastereomers (Table 1, entry 1). This low efficiency can be ascribed to the poor solubility of **2a**, which is rich in polar functional groups, in DCE. In fact, changing the solvent to MeCN significantly improved the product yield (entry 2), and the use of an MeCN/H<sub>2</sub>O (4 : 1) co-solvent system, which completely dissolved **2a**, led to the formation of **4a** in 62% yield (entry 3). Additional screening of polar solvents revealed that acetone was optimal for eliciting the full performance of **1a** in this transformation (entries 4–6). We then examined the effect of the catalyst structure, specifically the substituent of the amidate carbonyl, on the reactivity profile, as its steric and electronic properties would affect the hydrogen-bonding ability

Table 1 Optimization of reaction conditions and control experiments<sup>a</sup>



Entry	Catalyst	Solvent	Yield <sup>b</sup> (%)
1	<b>1a</b>	DCE	13
2	<b>1a</b>	MeCN	52
3	<b>1a</b>	MeCN/H <sub>2</sub> O (4 : 1)	62
4	<b>1a</b>	DMF	49
5	<b>1a</b>	DMSO	28
6	<b>1a</b>	Acetone	65
7 <sup>c</sup>	<b>1b</b>	Acetone	77 (72)
8	<b>1c</b>	Acetone	52
9	<b>1d</b>	Acetone	62
10 <sup>d</sup>	TiO <sub>2</sub>	Acetone	Trace
11 <sup>d</sup>	TiO <sub>2</sub>	MeCN	Trace
12 <sup>e</sup>	<b>5a</b> (10 mol%)	Acetone	12
13 <sup>e</sup>	<b>5a</b> (10 mol%)	DCM	8
14 <sup>e</sup>	<b>5b</b> (10 mol%)	Acetone	67
15 <sup>f</sup>	<b>5b</b> (3 mol%)	Acetone	40

<sup>a</sup> Unless otherwise noted, the reactions were conducted with 0.10 mmol of **2a** and 0.11 mmol of **3a** in the presence of catalyst (3 mol%) in solvent (0.2 mL) for 6 h under an argon atmosphere with 450 nm light-emitting diode (LED) irradiation and a fan to maintain the temperature. <sup>b</sup> NMR yield using trimethoxybenzene as an internal standard. Values in parentheses are isolated yields. <sup>c</sup> 0.20 mmol scale. <sup>d</sup> With 10 mg of anatase TiO<sub>2</sub> instead of **1** with 390 nm LED irradiation. <sup>e</sup> With 10 mol% of **5a/b** instead of **1** with 390 nm LED irradiation. <sup>f</sup> With 3 mol% of **5b** instead of **1** with 390 nm LED irradiation. Abbreviations: DMF, *N,N*-dimethylformamide; DMSO, dimethyl sulfoxide; DCM, dichloromethane.



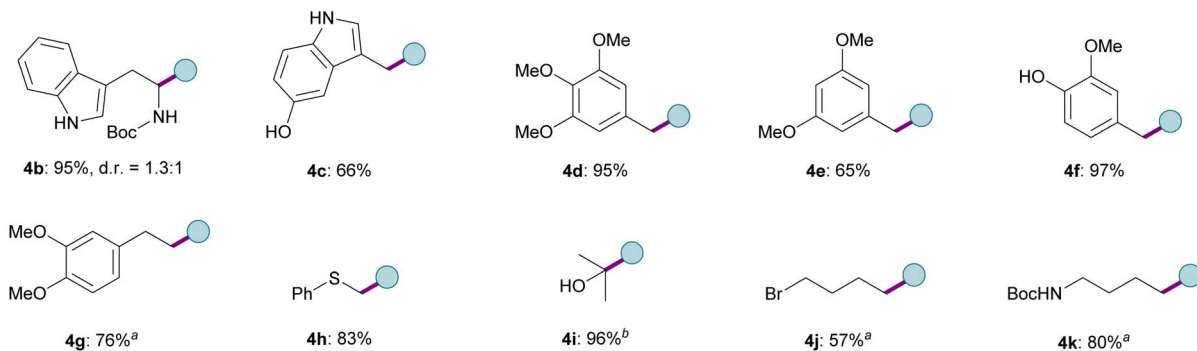


Fig. 2 Decarboxylative Giese-type addition of functionalized carboxylic acids to benzalmalononitrile **3b**. Isolated yields are shown. Diastereomeric ratio (d.r.) was determined by  $^1\text{H}$  NMR analysis of the crude reaction mixture. <sup>a</sup>Using 3 mol% of **1a**. <sup>b</sup>Isolated yield of the corresponding cyclized product **4i'** is shown. See the SI for details.

of the amidate moiety (entries 7–9). While introduction of a substituent derived from an  $\alpha$ -amino acid, such as L-valine (**1c**) or L-tert-leucine (**1d**), was not beneficial, pivaloyl-substituted **1b** delivered a slight improvement in the reaction efficiency, giving rise to **4a** in 72% isolated yield (diastereomeric ratio = 7 : 3.5 : 1 : 1) (entry 7). Control experiments under the influence of  $\text{TiO}_2$  or 9-(2-chlorophenyl)acridine (**5a**) with violet-light irradiation resulted in markedly diminished yields (entries 10–13). To further clarify the relationship between acridinium amidates

and conventional acridine photocatalysts, we evaluated the performance of 3,6-di-*tert*-butyl-9-mesitylacridine (**5b**). Under 390 nm light irradiation, the reaction with 10 mol% of **5b** afforded **4a** in 67% yield, demonstrating the ability of the acridine-type catalysts to promote decarboxylative alkylation of **2a** under high-energy shorter-wavelength light irradiation (entries 14 and 15). UV-vis absorption spectroscopic analysis showed that **1b** exhibits a pronounced absorption band centered at 420 nm, extending well into the visible-light region,

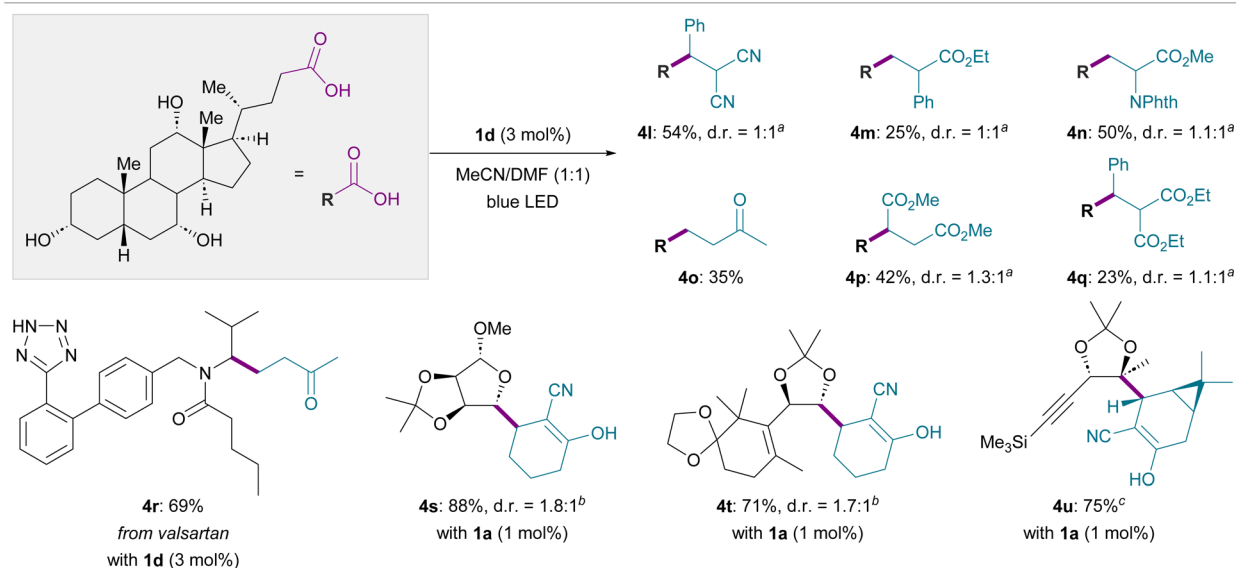
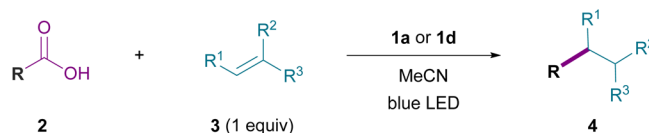


Fig. 3 Decarboxylative Giese-type addition of complex carboxylic acids. Isolated yields are shown. <sup>a</sup>Diastereomeric ratios (d.r.) were determined by  $^1\text{H}$  NMR analysis of the crude reaction mixture. <sup>b</sup>Isolated yields after the acetylation of enol moiety are shown; d.r. values were determined after acetylation and isolation. <sup>c</sup>Obtained as a 1 : 4.7 : 1.4 mixture of the enol and diastereomeric keto nitriles.



whereas **5b** displays negligible absorption above 400 nm, even in the presence of carboxylic acid (Fig. S4). Consistent with these absorption profiles, comparison of the catalytic activity of **5b** with that of **1b** in the reaction of **2a** with **3a** under carefully controlled blue-light irradiation conditions revealed that the product formation was not detected with **5b** as the catalyst, while **1b** facilitated the reaction in a time-dependent manner (Fig. S3). These results demonstrate that acridinium amidates operate in a distinct photophysical and catalytic regime, enabling efficient PCET-driven decarboxylation under longer-wavelength visible-light irradiation.

Next, we explored the general applicability of this photocatalytic protocol for the decarboxylative alkylation of various carboxylic acids bearing diverse functional groups using benzalmalononitrile (**3b**) as the radical acceptor (Fig. 2). When simple aliphatic carboxylic acids were employed, the Giese-type addition proceeded efficiently with 1 mol% of **1a** as a catalyst in MeCN to give the corresponding alkylated products **4** in high yields, irrespective of the substitution pattern at the  $\alpha$ -position (see the SI for optimization details). Notably, even with a redox-active indole component in the side chain, whose oxidation potential is lower than that of the carboxylate, *N*-Boc-tryptophan underwent smooth transformation to afford **4b** in nearly quantitative yield. Furthermore, decreasing the loading of **1a** to 0.1 mol% still provided **4b** in good yield (Table S1).<sup>45</sup> Likewise, a carboxylic acid with a more electron-rich 5-hydroxyindole unit appeared to be a viable substrate, furnishing **4c** in good yield. In addition, substrates with electron-rich aromatic substituents, such as poly-methoxylated benzenes and methoxyphenols, were well tolerated, leading to the formation of **4d–4g** without any observable inhibition. The broad functional-group compatibility of this method was also demonstrated in the reactions with carboxylic acids incorporating sulfide, hydroxy, bromo, and carbamate functionalities, which produced **4h–4k** in uniformly good yields.

Moreover, this catalytic system was reliably applicable to more complex substrate settings by adjusting the reaction conditions, as exemplified by the selective functionalization of cholic acid, a bile acid featuring a fused polycyclic framework with multiple hydroxy groups (Fig. 3). In the presence of **1d** (3 mol%) as a catalyst, decarboxylative radical addition to **3b** occurred in MeCN/DMF (1 : 1), affording the desired product **4l**. As radical acceptors for this decarboxylative alkylation, not only olefins with two electron-withdrawing groups but also simple vinyl ketones,  $\alpha$ -phenyl acrylate, and dehydroalanine derivatives were all employable, and the corresponding alkylated products **4m–4q** were obtained in moderate to good yields. In contrast, the reactions with less activated acceptors, such as simple acylates and acrylamides, gave only trace amounts of the desired products under the standard conditions (see SI, Section "Unsuccessful examples of radical acceptors"). The possibility of the chemoselective transformation of functionalized carboxylic acids was also showcased by the effective conversion of valsartan into **4r**. Base-labile 2-cyano-2-cyclohexenone was applicable to the coupling reaction to yield **4s**, accentuating the neutral nature of the present conditions.<sup>20</sup> The versatility of this protocol was further demonstrated by its application to complex

molecules bearing diverse reactive functionalities. Specifically, substrates containing acetal, cyclopropane, alkene, and alkyne groups were irradiated with **1a** to realize the intermolecular formation of the sterically hindered linkages of **4t** and **4u**, which were previously utilized as pivotal intermediates for the total synthesis of taxol and euphorbialoid A.<sup>46,47</sup>

With the broad applicability in mind, our investigation focused on elucidating the activation mechanism of the carboxy group in this photocatalytic transformation. The excited-state reduction potential of acridinium amidate **1a** was estimated to be +1.63 V vs. SCE,<sup>44</sup> which exceeds the oxidation potential of typical carboxylate ions (*ca.* +1.3 V vs. SCE). Nevertheless, when the reaction was conducted in the presence of hydroxide ions to ensure the *in situ* generation of carboxylate ions, little to no product formation was observed (Fig. 4A). This outcome indicates that direct oxidation of free carboxylate ions is unlikely to be a dominant pathway under the standard reaction conditions. <sup>1</sup>H NMR titration study in CD<sub>3</sub>CN revealed a significant association between **1a** and the carboxylic acid, consistent with the formation of a ground-state complex under catalytically relevant

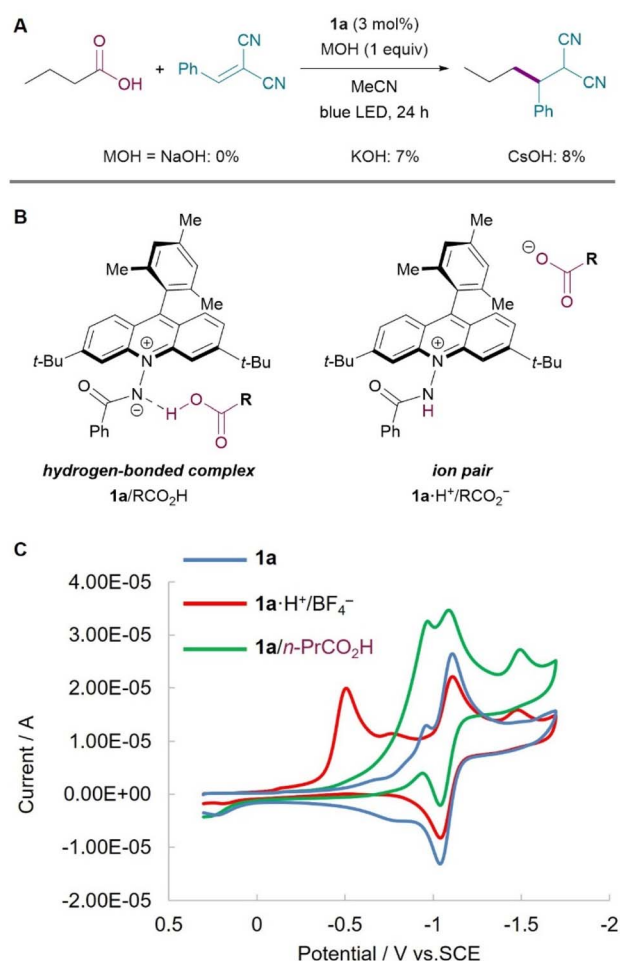


Fig. 4 Mechanistic investigations. (A) Control experiments with Brønsted base additives. (B) Structures of hydrogen-bonded complex and amide-acridinium/carboxylate ion pair. (C) Cyclic voltammetry of **1a** (blue line), the mixture of **1a** and HBF<sub>4</sub> (**1a**-H<sup>+</sup>/BF<sub>4</sub><sup>-</sup>, red line), and the mixture of **1a** and butyric acid (**1a**/*n*-PrCO<sub>2</sub>H, green line).



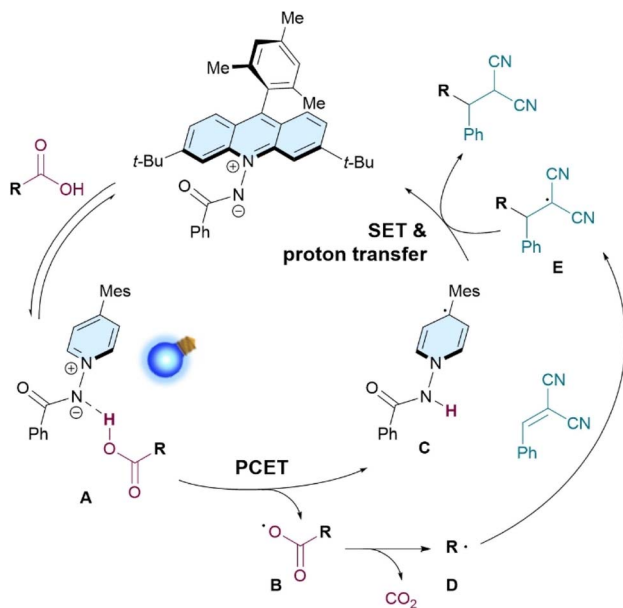


Fig. 5 Proposed catalytic cycle.

conditions (Fig. S7). Since the NMR data alone is insufficient to distinguish between a hydrogen-bonded assembly ( $1/\text{RCO}_2\text{H}$ ) and an ion-pairing interaction ( $1 \cdot \text{H}^+/\text{RCO}_2^-$ ), cyclic voltammetry (CV) measurements were performed to clarify the nature of this interaction (Fig. 4B). The CV of a solution containing **1a** and butyric acid showed the reduction wave corresponding to **1a** without additional signals attributable to the protonated form of the acridinium amidate, amide acridinium  $1\text{a} \cdot \text{H}^+/\text{BF}_4^-$  (Fig. 4C). This result suggests that **1** and carboxylic acid predominantly form a hydrogen-bonded assembly rather than an ion pair.

Based on these experimental findings, a plausible mechanism for the acridinium amidate **1**-catalyzed decarboxylative Giese-type addition is illustrated in Fig. 5. Initially, **1** engages in a hydrogen-bonding interaction with carboxylic acid to form ground-state complex **A**. Upon blue-light excitation, **A** undergoes PCET to generate carboxyl radical **B**, accompanied by the formation of **1**-derived acridinyl radical **C**. Decarboxylation of the resultant **B** readily occurs to generate carbon radical **D**, which adds to the electron-deficient olefin to afford radical intermediate **E**. Subsequent SET between **E** and **C**, followed by protonation, gives the desired Giese-type adduct and concurrently regenerates ground-state catalyst **1**.

## Conclusions

In summary, we developed a chemoselective decarboxylative Giese-type addition by exploiting the unique behaviour of a zwitterionic acridinium amidate as a photocatalyst. This transformation proceeds with remarkable efficiency under essentially neutral, visible-light-driven conditions and exhibits a broad substrate scope, tolerating a wide range of polar and redox-sensitive functional groups. The advantage of this distinct feature is underscored by its application to the direct

remodelling of natural product scaffolds and reactions of multiply oxygenated carboxylic acids with highly electron-deficient olefins, which constitute relevant bond-forming processes in the total synthesis of natural products. Mechanistic investigations indicated that the key to this prominent selectivity lies in the formation of a hydrogen-bonded complex between the catalyst and carboxylic acid, which triggers PCET to generate the requisite carboxyl radical without requiring pre-activation or external additives. This study demonstrates the immense potential of rationally designed zwitterionic photocatalysts for expanding the versatility of decarboxylative radical transformations, particularly for addressing the reactivity and selectivity challenges in high-complexity structural settings.

## Author contributions

The manuscript was prepared through the contributions of all authors. All the authors approved the final version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Data availability

The data supporting the findings of this study are available within the supplementary information (SI). Supplementary information: the exploratory investigation, experimental procedures, and characterization data. See DOI: <https://doi.org/10.1039/d6sc00657d>.

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