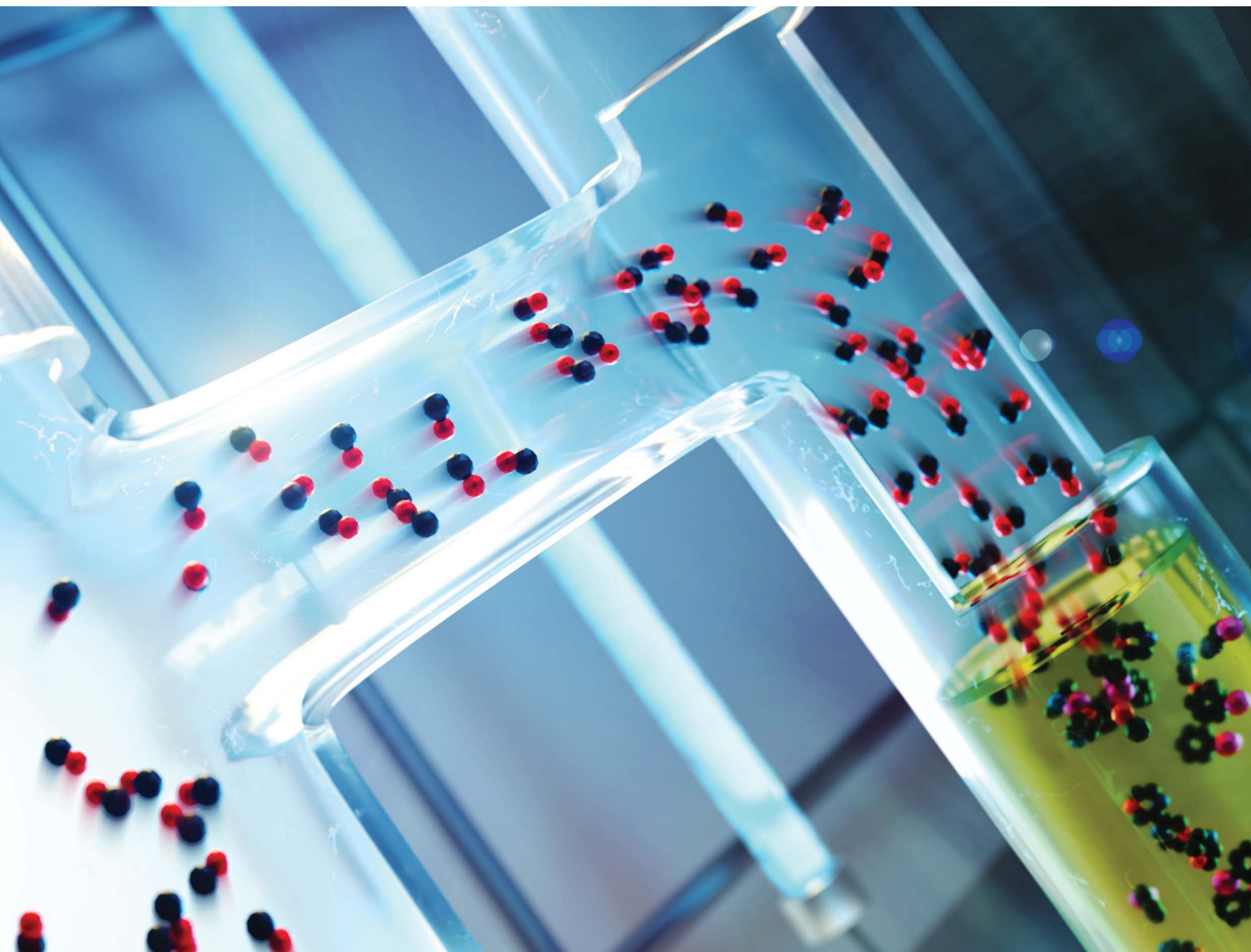


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Metal-free visible-light carbonylation of alkyl iodides to amides *via* consecutive photoinduced electron transfer

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A visible-light-driven, metal-free carbonylation of unactivated alkyl iodides is reported, enabling the direct synthesis of 35 structurally diverse amides in good to excellent yields. The reaction shows broad functional-group tolerance toward both amines and alkyl iodides, including bioisosteric motifs and complex natural product derivatives, underscoring its potential for late-stage functionalization. Mechanistic investigations combining flash photolysis, spectroelectrochemistry, irradiated thin-layer cyclic voltammetry, and EPR spectroscopy reveal a consecutive photoinduced electron transfer (ConPET) mechanism that diverges from conventional single-photon photoredox catalysis. DFT calculations elucidate the key radical carbonylation steps governing reactivity and selectivity. This sustainable and operationally simple method offers a transition-metal-free approach to carbonylation, expanding the toolbox for highly reducing transformations under mild conditions.

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Introduction

The direct conversion of simple alkyl halides into complex molecules remains a defining challenge in synthetic chemistry. Their strong C–X bonds and highly negative reduction potentials render both oxidative addition and single-electron transfer intrinsically difficult, limiting the catalytic strategies capable of activating these feedstock reagents under mild conditions.^{1–5} Surmounting this kinetic and thermodynamic barrier would unlock a wide spectrum of bond-forming transformations—from cross-couplings to carbonylations and beyond—based on readily available building blocks.

Amides exemplify the impact such activation strategies can have. They are among the most pervasive functional groups in chemistry, forming the structural backbone of peptides and proteins and occurring in countless pharmaceuticals, agrochemicals, and advanced materials.^{6–11} Despite their ubiquity, amide synthesis still largely depends on stoichiometric condensation of carboxylic acids and amines with coupling reagents such as carbodiimides or phosphonium salts, generating significant waste (Scheme 1a₁).¹² Catalytic alternatives have improved atom economy through organo- and transition-metal-catalyzed amidations or dehydrogenative couplings.^{12–14}

Aminocarbonylation—the three-component coupling of CO, an amine, and an organic halide—offers an elegant catalytic entry to amides in a single step (Scheme 1a₂).^{15–21} While well established for aryl, benzyl, and allyl halides, the extension to unactivated alkyl halides remains difficult because oxidative addition to Pd is slow and often followed by β-hydride elimination.²² Copper catalysis can circumvent these issues through single-electron transfer (SET) to generate alkyl radicals, enabling carbonylation of otherwise inert alkyl iodides, though typically under strongly reducing or forcing conditions.^{23–25} Ni(0) complexes supported by strongly σ-donating ligands such as N-heterocyclic carbenes or bulky phosphines were also reported to react with primary and secondary alkyl iodides under relatively mild conditions. These cases, however, remain limited in scope because the resulting metal-alkyl intermediates are also prone to rapid β-hydride elimination, and because SN₂-

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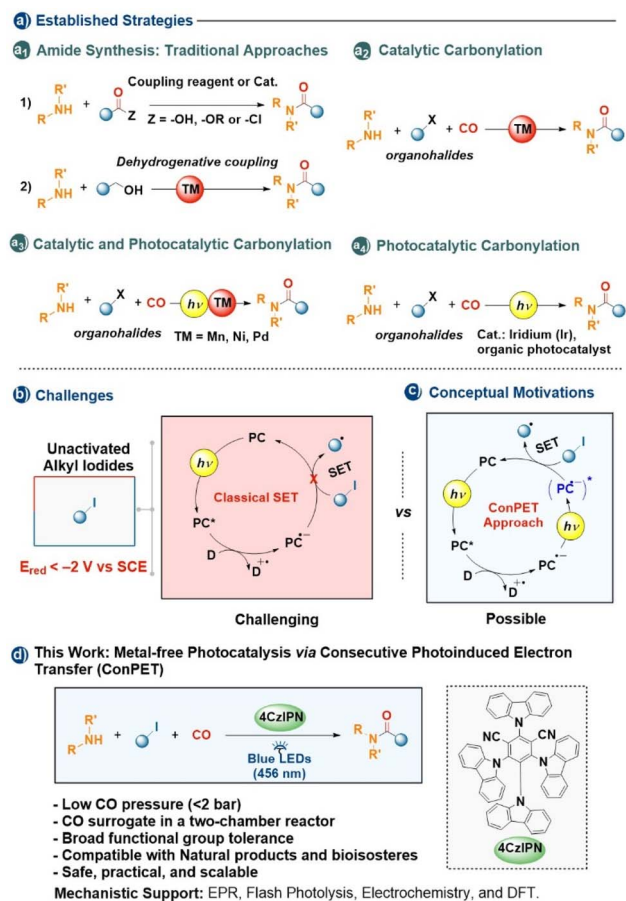
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Scheme 1 Overview of strategies for amide bond formation. (a) Traditional approaches: stoichiometric or catalytic condensation of carboxylic acids (or acyl halides) with amines, and transition-metal-catalyzed dehydrogenative coupling of alcohols. (b) Challenges in activating unactivated alkyl iodides by single-photon photoredox catalysis due to their highly negative reduction potentials. (c) Conceptual motivations: the consecutive photoinduced electron-transfer (ConPET) approach, in which a photocatalyst radical anion absorbs a second photon to reach an excited state of enhanced reducing power. (d) This work: metal-free visible-light aminocarbonylation of alkyl iodides using **4CzIPN** and an *in situ* CO source under mild conditions; mechanistic studies combine spectroscopy, electrochemistry, and DFT.

type oxidative addition is strongly disfavoured by steric and electronic factors. In addition, first-row transition metals generally favour single-electron pathways over the higher-energy two-electron oxidative addition.^{26–30} Co and Fe complexes predominantly operate through SET pathways, which can enable carbonylative aminocarbonylations. However, these systems generally require high CO pressures, typically in the range of 6 to 40 bar.^{31,32}

Photoredox catalysis has brought new momentum to this field by merging light-driven radical generation with transition-metal catalysis (Scheme 1a₃).^{33–39} Visible-light excitation has been shown to promote radical-assisted oxidative addition and reductive elimination at Pd under mild conditions, and related strategies have been extended to nickel and manganese catalysis.^{37–39} More recently, tandem two-photon cycles based on

fac-Ir(ppy)₃-type photocatalysts have enabled the cleavage of demanding C(sp³)-I bonds even under visible light irradiation (Scheme 1a₄).⁴⁰ Recent advances have demonstrated that metal-free photocatalytic strategies, employing organic photocatalysts or photochemical approaches, enable the carbonylative transformation of unactivated alkyl iodides using aryl formates as CO surrogates.^{41,42} This process generates phenolate while promoting the reduction of the iodoalkane to the corresponding alkyl radical.⁴¹

Nonetheless, the direct reduction of unactivated alkyl iodides ($E_{red} < -2 \text{ V vs. SCE}$) remains beyond the capability of most single-photon photocatalysts (Scheme 1b), limiting their engagement in mild radical transformations.^{1,43}

To transcend this fundamental energetic barrier, the concept of consecutive photoinduced electron transfer (ConPET) has emerged as a transformative strategy (Scheme 1c).^{44–50} In this two-photon process, a photocatalyst radical anion absorbs a second photon to access an excited state of dramatically enhanced reducing power, enabling reactions far beyond the thermodynamic reach of conventional photoredox catalysis. Building on these advances, we envisioned that combining ConPET activation with *in situ* CO generation could enable a metal-free aminocarbonylation of unactivated alkyl iodides under visible light.

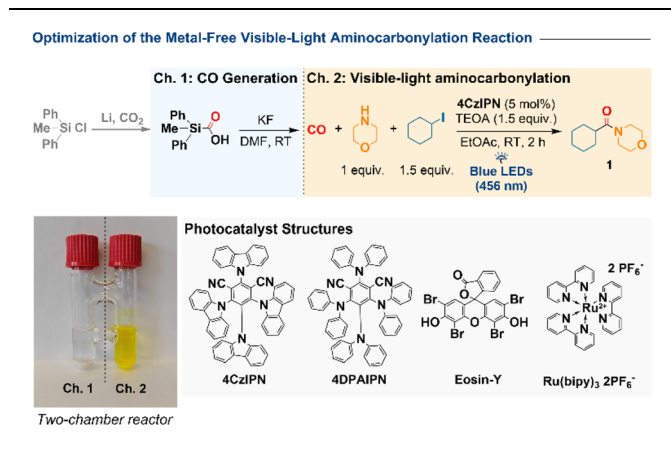
Here we describe such a system employing the organic photocatalyst **4CzIPN** (Scheme 1d). Carbon monoxide is produced safely *in situ* (1–2 bar) from methylphenylsilane carboxylic acid (“Silacogen”) in a sealed two-chamber reactor and consumed directly under visible light and ambient conditions. This operationally simple protocol affords a broad range of amides—including complex natural products and bioisosteres—in up to 85% yield within 2–8 h. Mechanistic investigations (*in situ* UV-vis spectroscopy, flash photolysis, spectroelectrochemistry, irradiated thin-layer cyclic voltammetry, and DFT) support a ConPET mechanism in which photoexcited **4CzIPN** is reduced to **4CzIPN^{•-}** and, upon re-excitation, activates demanding C-I bonds. Compared to single-photon photocatalysts such as **4DPAIPN**, this system enables rapid, mild, and operationally safe aminocarbonylation under low CO pressure.⁴²

Results and discussion

We commenced our investigation with the aminocarbonylation of iodocyclohexane and morpholine in a sealed two-chamber reactor (Table 1). Carbon monoxide was generated in chamber 1 from SilaCOgen—a bench-stable CO surrogate readily obtained from diphenylmethylchlorosilane and CO₂—by treatment with KF in DMF.^{51,52} In chamber 2, iodocyclohexane, morpholine, **4CzIPN** (5 mol%), and triethanolamine (TEOA) as a sacrificial electron donor were dissolved in ethyl acetate under argon and irradiated with a 50 W blue LED (456 nm, 50% intensity) at room temperature. After 2 h, amide **1** was obtained in 97% yield (GC-MS, hexadecane internal standard; entry 1). Control experiments highlighted the critical role of each component. Omission of TEOA reduced the yield to 37% (entry 2), and substitution with triethylamine or tributylamine led to



Table 1 Optimization and Control Experiments for the Carbonylative Amidation of Iodocyclohexane **1a**^a



Entry	Deviation from above reaction	Yield (%) ^b
1	None	97
2	No TEOA	37
3	Et ₃ N instead of TEOA	79
4	<i>n</i> -Bu ₃ N instead of TEOA	67
5	EtOH as solvent	13
6	CH ₃ CN as solvent	28
7	DMSO as solvent	40
8	DMF as solvent	45
9	Acetone as solvent	71
10	1,4-Dioxane as solvent	84
11	Eosin-Y instead of 4CzIPN	19
12	Ru(bpy) ₃ (PF ₆) ₂ instead of 4CzIPN	0
13	4DPAIPN instead of 4CzIPN	60
14	No photocatalyst (LED light only)	0
15	No light	0
16	Bromocyclohexane instead of iodocyclohexane	0
17	Chlorocyclohexane instead of iodocyclohexane	0

^a Chamber 1: SilaCOgen (0.60 mmol), KF (0.66 mmol), DMF (1.0 mL); chamber 2: morpholine (0.10 mmol), iodocyclohexane (0.15 mmol), **4CzIPN** (5 mol%), TEOA (0.15 mmol), AcOEt (1.0 mL); 456 nm Kessil® blue LED, RT, 2 h. ^b GC-MS yield vs. hexadecane (see SI).

diminished efficiency (79% and 67%, entries 3–4). The solvent proved decisive: EtOH, CH₃CN, DMSO, and DMF afforded only 13–45% yield (entries 5–8), whereas acetone and 1,4-dioxane performed significantly better (71% and 84%, entries 9–10). Among the photocatalysts evaluated, **4CzIPN** emerged as optimal. **Eosin-Y** delivered only 19% yield (entry 11), **Ru(bpy)₃**²⁺ was inactive (entry 12), and **4DPAIPN** provided 60% (entry 13). No product was detected in the absence of photocatalyst or light (entries 14–15). Finally, when using substrates with very high negative reduction potentials, such as bromo- and chlorocyclohexane, no products were obtained (entries 16–17).

Substrate scope

With the optimized conditions in hand, we explored the scope of the visible-light aminocarbonylation (Scheme 2). Cyclic secondary amines such as morpholine, piperidine, tetrahydroisoquinoline, and pyrrolidine participated readily, furnishing the corresponding amides **1–4** in 61–85% yield. A Boc-

protected piperazine delivered **5** in 59%, while a mono-substituted piperazine and diethylamine provided **6** and **7** in 51% and 57% yield, respectively. Primary amines were also suitable coupling partners, affording the corresponding amides **8–12** in up to 85% yield. Functionalized amines such as 2-aminoethanol and benzylamine were well tolerated, delivering amides **11** and **12**, respectively, whereas propargylamine afforded product **10** in 67% NMR yield (40% isolated).

Bioisosteric amines, including bicyclo[1.1.1]pentan-1-amine hydrochloride and its methyl-ester analogue, were compatible after *in situ* deprotonation with K₂CO₃, affording **16** and **17** in 50–78%. Strained heterocycles such as 2-oxa-6-azaspiro[3.3]heptane and 3-aminooxetane also underwent clean conversion, producing **18** and **19** in 74% yield.

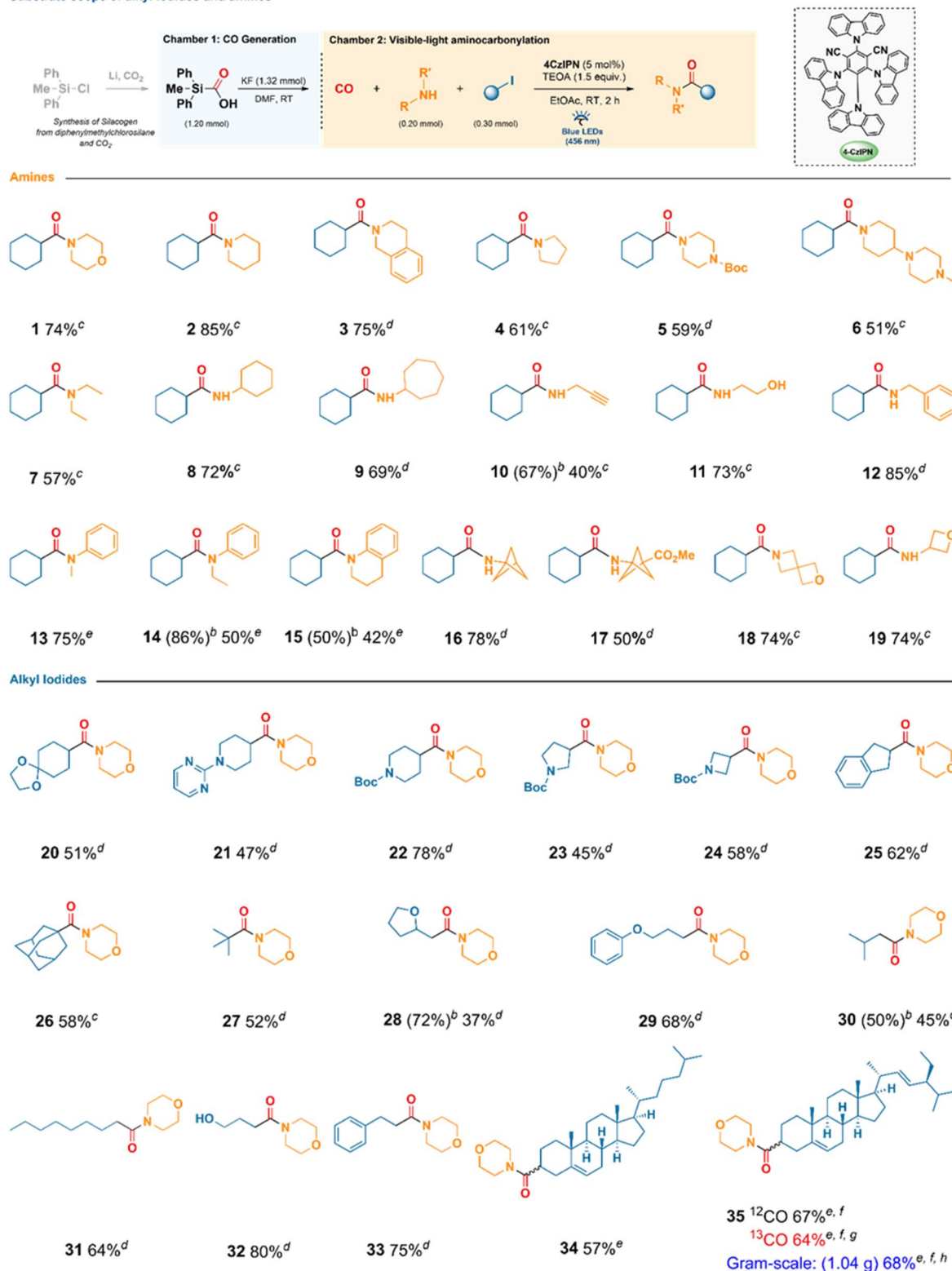
Aromatic amines displayed lower reactivity: aniline remained unreactive, whereas *N*-alkylanilines required extended irradiation to furnish **13–15** in 42–75% isolated yield (up to 86% NMR yield).

The transformation was general with respect to the alkyl partner. Secondary iodides furnished the corresponding amides in 45–78% yield, encompassing cyclic, spirocyclic, and benzylic substrates (**20–25**). Tertiary iodides—including 1-iodo-adamantane and *tert*-butyl iodide—underwent smooth coupling to give **26** and **27** (52–58%). Primary iodides exhibited substitution-dependent reactivity: heteroatom-substituted 2-(iodomethyl)tetrahydrofuran gave **28** in 72% NMR yield (37% isolated, the isolated yield being limited by purification), whereas (3-iodopropoxy)benzene, isobutyl iodide, and 1-iodooctane afforded **29–31** in 45–68%. Notably, substrates bearing unprotected hydroxyl or benzylic groups were well tolerated; 3-iodo-1-propanol and (2-iodoethyl)benzene yielded **32** and **33** in 80% and 75%, respectively.

The method was further extended to complex molecular scaffolds. Cholesterol- and stigmaterol-derived iodides underwent efficient aminocarbonylation under visible light (10 mol% **4CzIPN**, 8 h irradiation), providing **34** and **35** in 57% and 67% yield, respectively. Carbon-13 labelling was achieved by replacing Silacogen with Sila¹³COgen, affording the labelled stigmaterol amide in 64% yield; ¹³C incorporation was confirmed by FT-ICR-MS (see the SI). A gram-scale experiment using the stigmaterol substrate further highlighted the practicality and scalability of the protocol, furnishing **35** in 68% isolated yield.

This method demonstrates the efficiency of this metal-free reaction, which proceeds without the need for a glove box, within a short reaction time (ranging from 2 to 8 hours) at room temperature. It is compatible with a broad range of alkyl iodides (primary, secondary, and tertiary), including complex molecules such as cholesterol and stigmaterol derivatives, as well as various amines “cyclic, acyclic, aromatic, and bioisosteric” thereby overcoming the substrate limitations observed in previous studies. Notably, the incorporation of isotopic ¹³C using pure Sila¹³COgen achieves exceptional isotopic enrichment in natural products, reaching up to 99.7% (see SI). This represents a significant improvement over prior methods, which achieved enrichments of only 89% to 95%. Additionally, the CO source is derived exclusively from Sila¹³COgen, in contrast to previous approaches that relied on bubbling CO gas.⁴²



Substrate scope of alkyl iodides and amines^a

Scheme 2 Substrate scope of alkyl iodides and amines. (a) Chamber 1: SilaCOgen@ (1.20 mmol), KF (1.32 mmol), DMF (2.0 mL); chamber 2: amine (0.20 mmol), alkyl iodide (0.30 mmol), 4CzIPN (5 mol%), TEOA (0.30 mmol), ethyl acetate (2.0 mL); 456 nm Kessil@ blue LED, RT. Isolated yields unless otherwise noted. (b) NMR yields in parentheses (¹H NMR with tetrachloroethane internal standard). (c) 2 h. (d) 4 h. (e) 8 h. (f) 4CzIPN (10 mol%). (g) Sila¹³COgen. (h) gram-scale (3 mmol of morpholine).



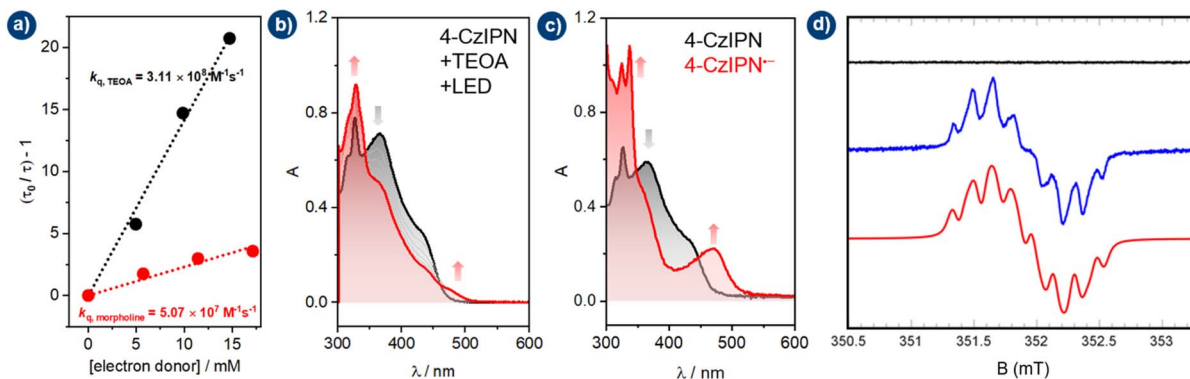


Fig. 1 (a) Stern–Volmer plot of $[(\tau_0/\tau) - 1]$ vs. concentration to determine dynamic reductive quenching by TEOA and morpholine. Data: quenching of TADF emission at 514 nm of 30 μM **4CzIPN** in Ar-saturated ethyl acetate, laser flash photolysis ($\lambda_{\text{exc}} = 430 \text{ nm}$, $E = 11 \text{ mJ}$). (b) *In situ* spectroscopy of Ar-saturated MeCN, 45 μM **4CzIPN** with 45 mM TEOA under blue irradiation (463 nm; black \rightarrow red: 0 \rightarrow 80 s). (c) Spectroelectrochemical generation of **4CzIPN** $^{\cdot-}$ in Ar-saturated MeCN at -1.40 V vs. SCE. (d) Experimental EPR spectra of TEOA and **4CzIPN** in the dark (black spectrum), under illumination ($\lambda = 475 \text{ nm}$, blue spectrum), and corresponding simulation (red spectrum), taking into account six ^{14}N nuclei with hyperfine coupling constants of 0.19, 0.17, 0.13, 0.08, 0.02, and 0.017 mT, respectively. Worthy of note, the two weakest values only contribute to the linewidth of the simulated spectra and are consistent with the optimized structures and SOMO distributions obtained by DFT (Fig. S3).

Mechanistic studies

To gain mechanistic insights into the visible-light ConPET aminocarbonylation, a combination of photophysical, electrochemical, and spectroscopic techniques was employed. Stern–Volmer analysis revealed that both TEOA and morpholine can reductively quench the excited state of **4CzIPN** to generate the radical anion **4CzIPN** $^{\cdot-}$ (Fig. 1a). This observation accounts for the residual reactivity observed in the absence of TEOA (Table 1, entry 2), when morpholine is present at high concentration. The dynamic quenching rate constant, however, is an order of magnitude higher for TEOA ($k_{q,\text{TEOA}} = 3.11 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$) than for morpholine ($k_{q,\text{morpholine}} = 5.07 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$). At concentrations of 150 mM TEOA and 100 mM morpholine, the apparent quenching rates are $4.66 \times 10^7 \text{ s}^{-1}$ and $5.06 \times 10^6 \text{ s}^{-1}$, respectively, rationalizing the superior performance of TEOA, which efficiently generates **4CzIPN** $^{\cdot-}$ while preserving morpholine as a nucleophile.

Under catalytically relevant conditions, *in situ* absorption spectroscopy of **4CzIPN**/TEOA in Ar-saturated MeCN revealed the growth of a new species upon 463 nm irradiation, characterized by diagnostic bands at 324, 336, and 470 nm (Fig. 1b; similar results in ethyl acetate, Fig. S4). UV-vis spectroelectrochemistry at -1.40 V vs. SCE, slightly beyond the first reduction wave, reproduced the same spectral features (Fig. 1c), assigning this species to the radical anion **4CzIPN** $^{\cdot-}$.

Continuous wave electron paramagnetic resonance (EPR) spectroscopy performed at room temperature under irradiation at 475 nm further corroborated the formation of **4CzIPN** $^{\cdot-}$ (Fig. 1d). While no EPR signal was observed in the dark, a characteristic EPR fingerprint was detected as soon as the light was switched on (Fig. 1d, blue spectrum). The observed spectra are in agreement with those previously reported for **4CzIPN** $^{\cdot-}$ and derivatives,⁴⁴ obtained with hyperfine coupling to six nitrogen ^{14}N nuclei (Fig. 1d, red spectrum). The observed signal also supports the role of morpholine acting as

a sacrificial agent by transferring an electron to the photocatalyst (Fig. S2), explaining its partial consumption and the associated reduction in yield.

Single-photon photoredox cycles, as established for related catalysts such as **4DPAIPN**, typically proceed through a radical anion ($E_{\text{red}} \approx -1.5 \text{ V}$ vs. SCE) that is not sufficiently reducing to activate unactivated alkyl iodides ($E_{\text{red}} < -2 \text{ V}$ vs. SCE).^{40,42} In contrast, our results support a consecutive photoinduced electron transfer (ConPET) pathway,^{46,47,53} in which **4CzIPN** $^{\cdot-}$ absorbs a second photon to form a highly reducing excited radical anion (**4CzIPN** $^{\cdot-*$), capable of cleaving $\text{C}(\text{sp}^3)\text{-I}$ bonds and initiating the aminocarbonylation.

To exclude the possibility that ground-state **4CzIPN** $^{\cdot-}$ acts as the active reductant, we performed irradiated thin-layer cyclic

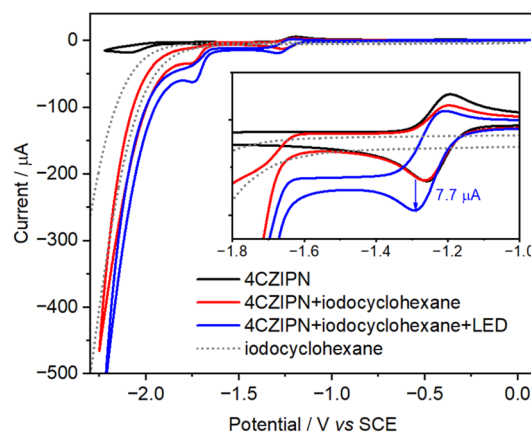


Fig. 2 Cyclic voltammograms (CVs) of 0.5 mM **4CzIPN** in Ar-saturated MeCN with 0.1 M NBu_4PF_6 (black), with addition of 50 mM iodo-cyclohexane in the dark (red), and under irradiation of blue LED (blue). Dotted gray line shows background reduction of 50 mM iodo-cyclohexane on the glassy carbon electrode. Inset shows a magnified part of the changes in the first redox peak.

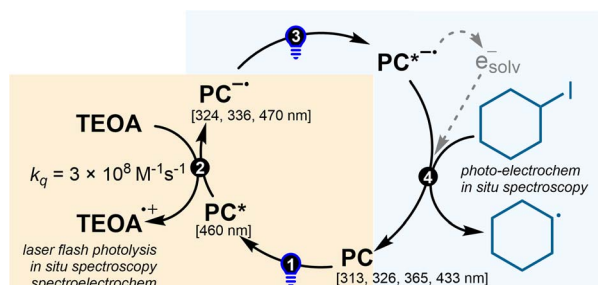


voltammetry (CV)—to our knowledge, its first application in a ConPET aminocarbonylation (Fig. 2). CV of **4CzIPN** (Fig. 2, black CV) exhibited a reversible first reduction with $E_{1/2} = -1.23$ V vs. SCE (**4CzIPN**/**4CzIPN^{-•}**). Addition of iodo-cyclohexane in the dark (Fig. 2, red CV) produced no enhancement of current at this wave, indicating that the electrochemically generated **4CzIPN^{-•}** alone is insufficiently reducing. Current growth due to halide reduction appeared only beyond -1.70 V vs. SCE. Under 463 nm irradiation (Fig. 2, blue CV), however, a pronounced current increase at the first wave was observed, consistent with *in situ* photoexcitation of **4CzIPN^{-•}** to its excited state (**4CzIPN^{-•*}**) or formation of a functionally equivalent solvated electron. Although the excited radical anion is ultrashort-lived and nonemissive,⁴⁶ the light-induced catalytic wave provides compelling electrochemical evidence for its enhanced reducing character.⁵⁴ Photoinduced detachment to solvated electrons,^{46,48,55} previously proposed for similar systems, may also contribute; our CV experiments cannot distinguish between these parallel pathways.

Together, the photophysical and electrochemical results support the ConPET sequence depicted in Scheme 3: (1) single-photon excitation of **4CzIPN**; (2) reductive quenching by TEOA (or morpholine) to yield **4CzIPN^{-•}**; (3) subsequent photoexcitation of **4CzIPN^{-•}** to its excited state (**4CzIPN^{-•*}**) (or, alternatively, generation of a solvated electron); and (4) reduction of the alkyl iodide to produce the corresponding alkyl radical.

This result contrasts with the recently reported mechanism involving a single photon absorption by the **4DPAIPN** photocatalyst for the carbonylation of alkyl halides.⁴²

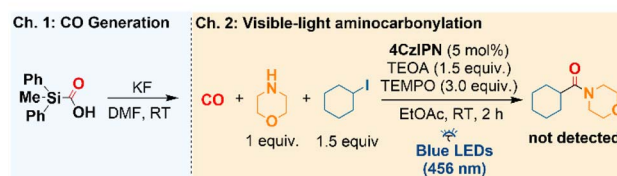
The use of **Eosin-Y** resulted in only a 19% yield, which can be attributed to its insufficient reducing potential ($E_{\text{red}} \approx -1.1$ V).⁵⁶ The calculated energy difference (ΔE) of -54.9 kcal mol⁻¹ for **Eosin-Y** indicates a relatively low electron affinity, consistent with the inefficient formation of a stable radical anion. This suggests that **Eosin-Y** does not effectively participate in the ConPET cycle, thereby explaining the observed low yield. In contrast, **4CzIPN** exhibits a more negative energy difference ($\Delta E = -73.9$ kcal mol⁻¹), reflecting a higher electron affinity and enabling the formation of a more stable radical anion. This stability facilitates its participation in the ConPET cycle, leading to efficient re-excitation and a higher quantum yield in the



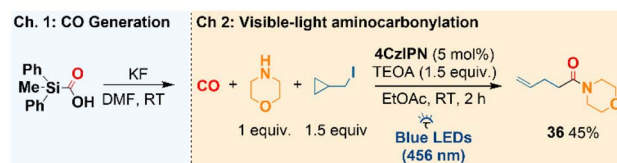
Scheme 3 Proposed ConPET mechanism for alkyl radical generation by **4CzIPN** (PC). Reductive quenching of photoexcited **4CzIPN** by TEOA forms **4CzIPN^{-•}**, which, upon second-photon excitation, reaches an excited state capable of reducing unactivated alkyl iodides to alkyl radicals.

Probing the Radical Mechanism

(a) Radical Trapping with TEMPO



(b) Cyclopropylmethyl Iodide Ring Opening



Scheme 4 Probing the radical mechanism. (a) TEMPO-trapping experiment: addition of TEMPO suppresses product formation. (b) Cyclopropylmethyl iodide undergoes ring-opening aminocarbonylation to yield the unsaturated amide **36**, consistent with a radical pathway.

photocatalytic process. For **4DPAIPN**, the calculated ΔE of -66.9 kcal mol⁻¹ indicates some capacity to form a radical anion; however, this species is less stable than the one derived from **4CzIPN**. This lower stability correlates with its limited efficiency in the ConPET cycle compared to **4CzIPN** (see SI).

The radical pathway of the carbonylation step was established through diagnostic control experiments. Addition of the radical scavenger TEMPO completely suppressed product formation (Scheme 4a). Furthermore, the use of cyclopropylmethyl iodide afforded the ring-opened amide **36** (Scheme 4b), a typical radical-clock outcome. These findings confirm that, once formed, the alkyl radical engages in a free-radical carbonylation sequence rather than a closed-shell pathway.

Density-functional theory (DFT) calculations provided additional insight into the reaction profile following alkyl-radical formation (Fig. 3). The initially generated cyclohexyl radical (**1a**) rapidly adds to carbon monoxide to give the corresponding acyl radical (**1b**). This step proceeds with an activation barrier of 11.9 kcal mol⁻¹ (**TS1**) and is slightly exergonic ($\Delta G = -0.8$ kcal mol⁻¹). Subsequently, (**1b**) undergoes iodine-atom transfer from iodo-cyclohexane to regenerate the cyclohexyl radical (**1a**), passing through **TS2** with a barrier of 16.4 kcal mol⁻¹ to afford the acyl iodide (**1c**), located +3.2 kcal mol⁻¹ above (**1b**). This sequence suggests a radical chain-propagation mechanism in which the alkyl radical is continuously regenerated. Next, a nucleophilic attack of morpholine on acyl iodide **1c**, accompanied by iodide departure, proceeds through **TS3** ($\Delta G^\ddagger = 11.3$ kcal mol⁻¹) to give the cationic intermediate **1d**, which subsequently undergoes hydrogen abstraction (Int1) to yield the final amide (**1e**). The computed free-energy profile indicates that (**1e**) is 7.7 kcal mol⁻¹ lower in energy than the starting radical (**1a**), confirming both the thermodynamic driving force and the overall feasibility of the transformation.



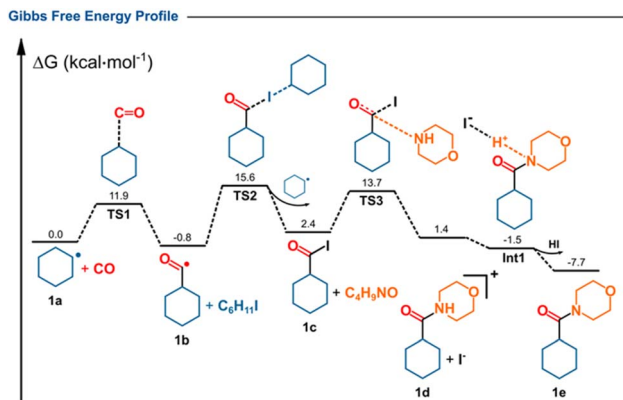


Fig. 3 Gibbs free energy (ΔG) profile for the proposed reaction path. DFT calculations performed at the M06-2X/6-311++G** level of theory, considering acetonitrile as the solvent within the Polarizable Continuum Model (PCM).

Unlike morpholine, aniline fails to yield any carbonylation product. While aniline exhibits lower nucleophilicity (N_{aniline} : 12.6–13.0) compared to morpholine ($N_{\text{morpholine}}$: 15.6),⁵⁷ this difference alone does not fully account for its lack of reactivity relative to other amines. DFT calculations reveal that the computed activation free energies (ΔG^\ddagger) for nucleophilic addition to the carbonyl group are comparable across the four amines investigated: aniline (12.9 kcal mol⁻¹), ethylaniline (13.2 kcal mol⁻¹), methylaniline (14.1 kcal mol⁻¹), and morpholine (11.3 kcal mol⁻¹). However, the stark contrast in observed reactivity is primarily governed by the thermodynamic stability of the resulting tetrahedral intermediates.

Morpholine demonstrates both a moderately low activation barrier and a product/intermediate free energy slightly lower than that of the reactants ($\Delta G = -1.0$ kcal mol⁻¹), indicating a kinetically accessible and thermodynamically favorable reaction pathway. Conversely, aromatic amines exhibit positive free energies of reaction ($\Delta G = +8.9$, $+4.0$, and $+6.9$ kcal mol⁻¹ for aniline, ethylaniline, and methylaniline, respectively), signifying thermodynamically disfavored intermediate formation that inhibits the reaction under standard conditions. This effect is most pronounced for aniline: despite a comparable activation barrier, the tetrahedral intermediate is highly unstable ($\Delta G = +8.9$ kcal mol⁻¹), effectively precluding product formation. Collectively, these findings underscore that both nucleophilicity and the stability of the tetrahedral intermediate dictate the reactivity of amines in this transformation.

Conclusions

We have developed a metal-free, visible-light-driven amino-carbonylation of unactivated alkyl iodides that affords a broad range of amides under mild conditions. The transformation tolerates diverse primary and secondary amines, sterically hindered and functionalized alkyl iodides, and complex scaffolds derived from bioisosteres and natural products, rendering it particularly attractive for late-stage functionalization and medicinal chemistry applications. Key practical advantages include the use of an inexpensive organic photocatalyst (**4CzIPN**) and a bench-

stable, low-pressure CO source, which enhance operational safety and scalability compared to pressurized CO systems. Mechanistic studies combining *in situ* UV-vis spectroscopy, flash photolysis, spectroelectrochemistry, and irradiated thin-layer cyclic voltammetry support a consecutive photoinduced electron transfer (**ConPET**) pathway. Following initial single-photon reduction of **4CzIPN**, a second excitation of the radical anion generates an exceptionally reducing excited state capable of activating demanding C(sp³)-I bonds. These observations exclude a simple single-photon process and establish **ConPET** as a general strategy for achieving extreme photoreductive potentials under mild, metal-free conditions. Overall, this work demonstrates that organic photoredox catalysis can rival and even surpass transition-metal systems in challenging carbonylation chemistry, offering a safe, scalable, and operationally simple route to amides while broadening the conceptual scope of highly reducing radical transformations.

Author contributions

F. A., G. C., M. E., and N. I. conducted the photocatalytic experiments and performed the associated data analysis. J. H. carried out the FT-ICR-MS and HRMS analyses. B. V. and N. L. carried out the EPR experiments and analyzed the results. M. B. and M. P. V. performed the DFT calculations. A. V. provided technical support for the execution of experiments and data analysis. P. G. conducted the flash photolysis and electrochemistry analyses, including mechanistic studies. Z. H., C. W., and I. A. conceived the project and oversaw the experimental work. All authors contributed to the writing of the manuscript and have approved its final version.

Conflicts of interest

The authors declare no conflicts of interest.

Data availability

The data supporting this article are included in the supplementary information (SI). Supplementary information: experimental and syntheses procedures, characterisations including NMR, EPR, electrochemistry, spectroelectrochemistry, *in situ* UV-vis spectroscopy, DFT calculations. See DOI: <https://doi.org/10.1039/d5sc09778a>.

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