

Fragment Couplings via CO₂ Extrusion–Recombination: Expansion of a Classic Bond-Forming Strategy via Metallaphotoredox

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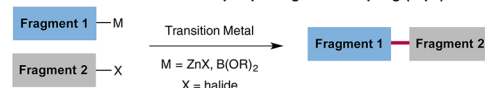
S Supporting Information

ABSTRACT: In this study we demonstrate that molecular fragments, which can be readily coupled via a simple, in situ RO–C=OR bond-forming reaction, can subsequently undergo metal insertion–decarboxylation–recombination to generate C_{sp²}–C_{sp³} bonds when subjected to metallaphotoredox catalysis. In this embodiment the conversion of a wide variety of mixed anhydrides (formed in situ from carboxylic acids and acyl chlorides) to fragment-coupled ketones is accomplished in good to high yield. A three-step synthesis of the medicinal agent edivoxetine is also described using this new decarboxylation–recombination protocol.

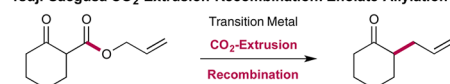
Metal-catalyzed intermolecular C–C bond formation has long been established as the predominant technology for fragment coupling in chemical synthesis. In particular, organometallic nucleophiles and organic halide/pseudohalide electrophiles have become the mainstay coupling partners for the transition metal-catalyzed production of hetero C_{sp²}–C_{sp³} bonds in a highly efficient and selective fashion (eq 1).¹ Moreover, the allylation and benzylation of enolates via the decarboxylative formation of π-allyl systems from β-keto-allyl esters have long been established as an important variant of the classic Tsuji–Trost mechanism (eq 2).² The recent merger of photoredox and transition metal catalysis (termed metallaphotoredox catalysis) has gained momentum as a strategy for unique cross-coupling protocols,³ mainly due to the capacity to employ naturally occurring functional groups as traceless activation handles and the ability to achieve fragment couplings that readily build challenging C_{sp²}–C_{sp³} bonds. In this context, our lab recently disclosed a light-enabled decarboxylative cross-coupling strategy that employs a diverse range of carboxylic acids in lieu of organometallic nucleophiles in combination with Ni catalysis.⁴ These methodologies utilize abundant and easily accessible starting materials to build a diverse array of C_{sp²}–C_{sp³} bonds at room temperature while producing CO₂ as a traceless byproduct.

Recently, we became interested in establishing a heretofore unknown fragment-coupling reaction that employs a CO₂ extrusion–recombination strategy (CO₂ExR) that in a general sense bears the hallmarks of the classic Tsuji allylation reaction. Specifically, we hoped to demonstrate that two fragments that can be readily coupled via a simple C–O bond-forming step (e.g., in situ formation of an anhydride, ester, carbamate, etc.) might subsequently undergo metal insertion–decarboxylation–recombination under metallaphotoredox conditions to enable the production of relatively complex C–C bonds (e.g., C_{sp²}–C_{sp³},

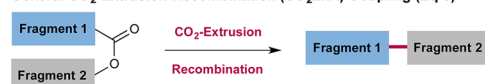
Conventional Metal-mediated sp²–sp² Fragment Coupling (Eq 1)



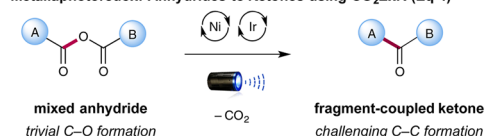
Tsuji-Saegusa CO₂-Extrusion-Recombination: Enolate Allylation (Eq 2)



General CO₂-Extrusion-Recombination (CO₂ExR) Coupling (Eq 3)



Metallaphotoredox: Anhydrides to Ketones using CO₂ExR (Eq 4)



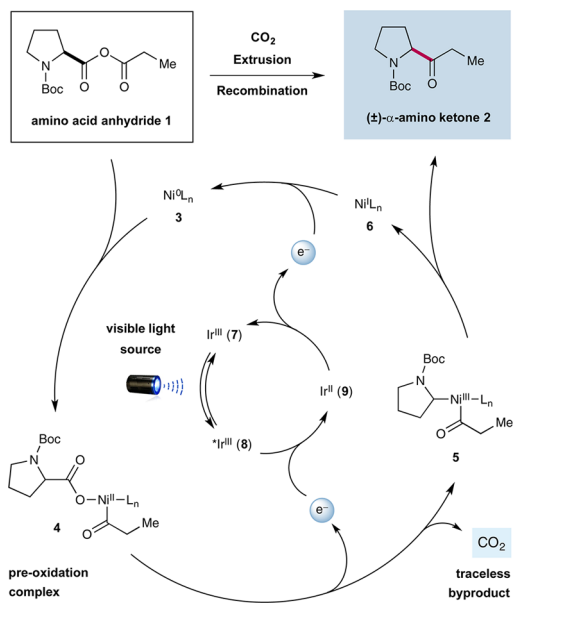
C_{sp³}–C_{sp³}, eq 3). While the strategy of CO₂ExR has long been established in organometallic catalysis for enolate allylation or benzylation,^{2,5} we hoped this new metallaphotoredox mechanism would provide an expansion in the types of organic fragments or motifs (e.g., nucleophiles and electrophiles) that can be linked via a simple RO–C=OR bond-forming step prior to CO₂ExR.⁶ As an initial proof of concept, we chose to examine a protocol that would selectively combine and convert acid chlorides and carboxylic acids to fragment-coupled ketones via the intermediacy of a mixed anhydride (formed in situ, eq 4).

Here we present the successful implementation of these ideals and disclose the first application of this expanded CO₂ExR concept toward the production of fragment coupled ketones and C_{sp²}–C_{sp³} bonds.

Design Plan. As shown in Scheme 1, our proposed mechanism begins with oxidative insertion of Ni⁰ complex 3 to acid anhydride 1 (generated in situ from carboxylic acid and acyl chloride coupling partners) to form the corresponding acyl-carboxylate–Ni^{II} complex 4.⁷ Concurrently, Ir^{III} photocatalyst Ir[dF(CF₃)ppy]₂(dtbbpy)PF₆ (7) [dF(CF₃)ppy = 2-(2,4-difluorophenyl)-5-(trifluoromethyl)pyridine, dtbbpy = 4,4'-di-tert-butyl-2,2'-bipyridine] is known to undergo photoexcitation in the presence of visible light to yield the corresponding *Ir^{III} complex 8. This long-lived excited state (τ = 2.3 μs)⁸ possesses a high oxidizing power (E_{1/2}^{red}[*Ir^{III}/Ir^{II}] = +1.21 V vs SCE in MeCN)⁸ and should rapidly accept an electron from the Ni^{II}

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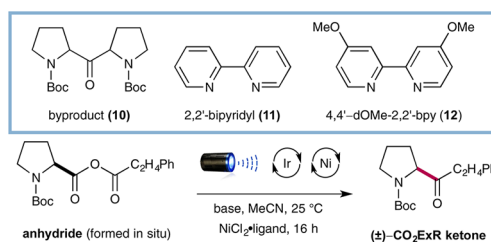
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Scheme 1. Mechanism of CO₂ Extrusion–Recombination

anhydride-insertion species **4**, thereby inducing oxidative decarboxylation to forge the corresponding alkyl acyl Ni^{III} complex **5**.⁹ Rapid reductive elimination should then deliver ketone product **2** and the corresponding Ni^I species **6**. Finally, completion of both catalytic cycles would occur simultaneously via single electron transfer (SET) between the highly reducing Ir^{II} complex **9** ($E_{1/2}^{\text{red}}[\text{Ir}^{\text{III}}/\text{Ir}^{\text{II}}] = -1.37$ V vs SCE in MeCN)⁸ and the transient Ni^I species **6** to reconstitute the ground state of photocatalyst **7** and Ni⁰ catalyst **3** ($E_{1/2}^{\text{red}}[\text{Ni}^{\text{II}}/\text{Ni}^{\text{0}}] = -1.20$ V vs SCE in DMF).¹⁰

Studies toward the proposed CO₂ExR of mixed anhydrides began with the coupling of hydrocinnamoyl chloride and Boc-L-proline in the presence of photocatalyst **7**, NiCl₂·glyme, 2,2'-bipyridyl (**11**), Cs₂CO₃, and blue LEDs as the light source (Table 1). As a critical design element, we recognized that in situ formation of the requisite anhydride would eliminate the need for an intermediate isolation step, thereby rendering the overall transformation operationally simple. To our delight, our initial experiment furnished the desired fragment-coupled ketone in a promising 40% yield (Table 1, entry 1) albeit with 20% yield of undesired homodimeric ketone **10**. We recognized that production of this latter symmetrical dialkyl ketone likely arises from anhydride metathesis (metal or base catalyzed) prior to the oxidative decarboxylation step. Indeed, variation of the reaction base from Cs₂CO₃ to 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) provided a significant increase in the yield of the desired ketone (entry 2, 70% yield). Moreover, we were pleased to find that implementation of more electron-donating ligand **12** on the Ni^{II}Cl₂·L_n complex afforded the desired fragment-coupled ketone in 84% yield while limiting the formation of symmetrical ketone **10** to <3% yield (entry 3). The necessity of each catalytic component was demonstrated via a series of control experiments (entries 4–7, 0% yield). Finally, the direct implementation of purified anhydride gave comparable results (entry 8, 73% yield), demonstrating that the in situ O–C=O formation step does not impact the efficiency of the subsequent C–C fragment coupling.

With the optimized conditions in hand, we began to explore the scope of this transformation with respect to the acyl chloride component. As shown in Table 2, CO₂ExR is successful with a range of α -methylene-bearing acid chlorides (**14**–**18**, 77–86%

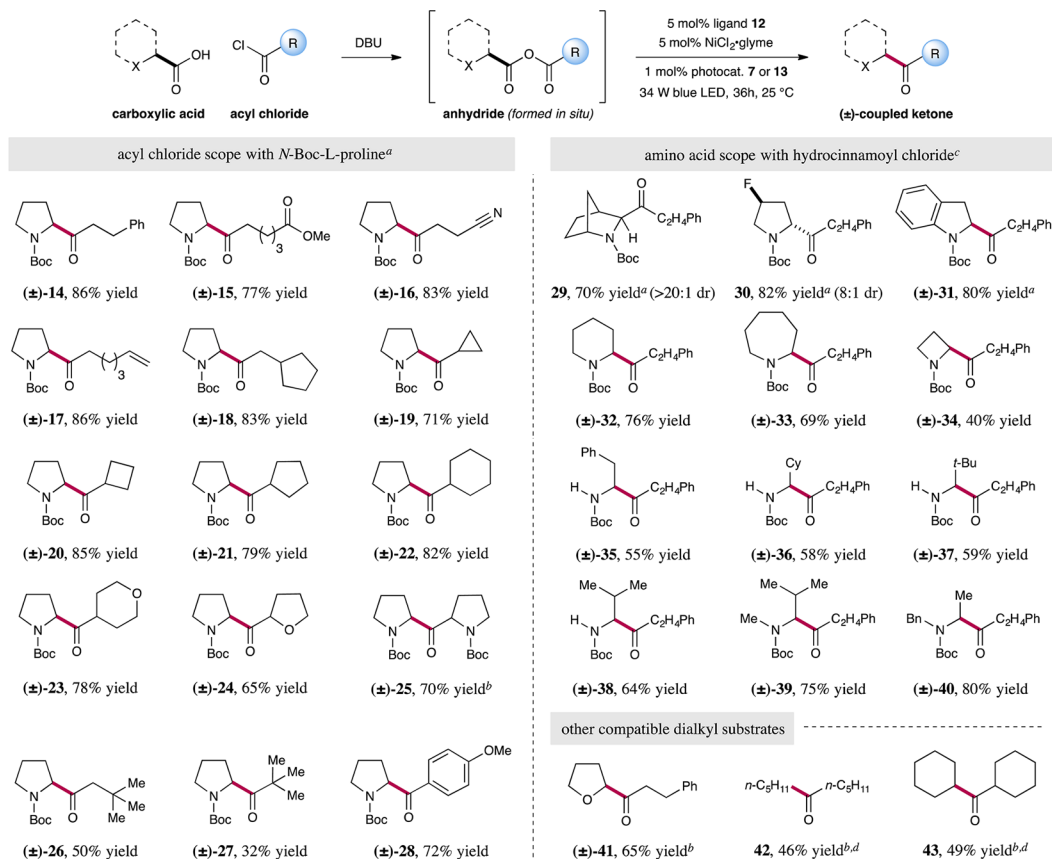
Table 1. Optimization of Anhydride CO₂ Extrusion–Recombination^a

entry	conditions	base	ligand	byproduct 10 ^b	ketone
1	as shown	Cs ₂ CO ₃	11	20%	40%
2	as shown	DBU	11	14%	70%
3	as shown	DBU	12	3%	84%
4	no photocatalyst	DBU	12	0%	0%
5	no Ni catalyst	DBU	–	0%	0%
6	no base	–	12	0%	0%
7	no light	DBU	12	0%	0%
8 ^c	as shown	DBU	12	5%	73%

^aReactions performed using photocatalyst **7** (1 mol%), NiCl₂·glyme (5 mol%), bipyridine ligand (5 mol%), hydrocinnamoyl chloride (0.10 mmol), *N*-Boc-L-proline (0.13 mmol), and base (0.13 mmol). Yields determined by GC analysis using an internal standard. ^bMajor byproduct. ^cAnhydride was synthesized and isolated prior to reaction.

yield). Moreover, various carbo- and heterocyclic-substituted anhydrides may be employed to rapidly generate ketones that incorporate three- through six-membered rings at the α -carbonyl position (**19**–**25**, 65–85% yield). Notably, many of these ketones would not be readily accessible using conventional ketone forming technologies such as Weinreb amide-Grignard additions.^{11,12} We have found that acid chlorides that contain sterically demanding groups, such as *tert*-butyl or neopentyl moieties, deliver appreciable levels of efficiency (**26** and **27**, 50 and 32% yield). Finally, CO₂ExR provides a new strategy for the production of aryl-substituted ketones as exemplified by the formation of adduct **28** in good yield.

We next examined the generality of anhydride CO₂ExR with respect to the carboxylic acid component. Given their widespread availability, abundance, and diverse structural complexity, we were delighted to find that α -amino acids were exemplary substrates for this new decarboxylation fragment-coupling protocol (Table 2). For example, various five-membered ring amino acids rapidly undergo CO₂ExR with hydrocinnamoyl chloride to form the corresponding α -amino ketone adducts in good to excellent yield (**29**–**31**, 70–82% yield). Interestingly, the bridged bicyclic ketone **29** was formed as a single diastereomer while maintaining good levels of reaction efficiency (70% yield). For amino acid systems beyond proline derivatives, we found that the photocatalyst Ir[dF(OMe)ppy]₂(dtbbpy)PF₆ (**13**) [dF(OMe)ppy = 2-(2,4-difluorophenyl)-5-(methoxy)pyridine] was more effective (products **32**–**40**). As an example of the useful levels of complexity that can be generated in this new CO₂ExR protocol, direct access to six- and seven-membered cyclic α -amino ketones is readily accomplished (**32** and **33**, 76 and 69% yield). We were also pleased to find that smaller strained rings, such as azetidine **34**, can be tolerated, albeit in slightly diminished yield (40% yield). Gratifyingly, this transformation does not appear to be overly influenced by the steric constraints of the amino acid substrate (e.g., valine and *tert*-leucine systems are readily employed, **35**–**38**, 55–64% yield). Indeed, *N*-alkylated *N*-Boc acids were found to exhibit superior efficiency over *N*-H-

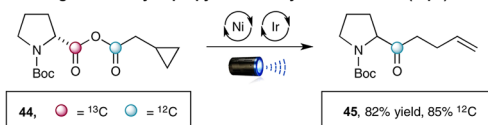
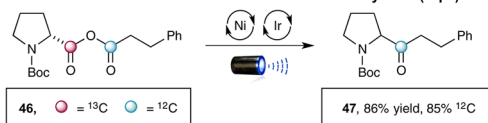
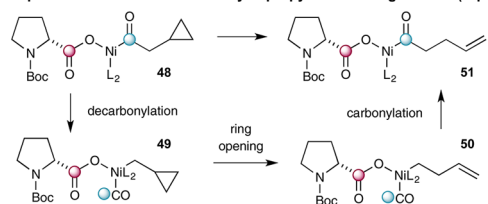
Table 2. Mixed Anhydride CO₂ Extrusion–Recombination: Scope of Acyl Chloride and Carboxylic Acid Components

^aReactions performed using photocatalyst 7 (1 mol%) in acetonitrile, NiCl₂·glyme (5 mol%), ligand 12 (5 mol%), acyl chloride (0.10 mmol), *N*-Boc-L-proline (0.13 mmol), and DBU (0.13 mmol). Yield of isolated product. ^bSee Supporting Information for experimental procedure. ^cReactions performed using photocatalyst 13 (1 mol%) in dioxane using hydrocinnamoyl chloride (0.10 mmol), amino acid (0.10 mmol), and DBU (0.13 mmol). Yield of isolated product. ^dGC yield.

bearing substrates (cf. 38, 39, and 40). We speculate that the presence of additional electron-donating groups on the nitrogen group might lower the barrier to SET in the oxidative decarboxylation step, resulting in improved rates of formation of the requisite Ni^{III} species (i.e., 5, Scheme 1). Beyond amino acids, we were pleased to find that α -oxy and aliphatic acids can be utilized in this new coupling protocol to forge the corresponding ketones in moderate to good yield (41–43, 46–65% yield).^{13,14}

During the examination of the substrate scope, an interesting rearrangement was observed when β -cyclopropyl anhydride 44 (formed in situ) was subjected to this decarboxylation–recombination strategy. As shown in eq 5, the product obtained was not the expected β -cyclopropyl ketone (formed in <3% as determined by ¹H NMR and GC analysis), but instead homoallylic ketone 45 (formed in 82% yield). Control experiments have demonstrated that this rearrangement occurs during the CO₂ExR pathway (see Supporting Information). Moreover, labeling experiments involving ¹³C-labeled amino acids have shown that CO₂ loss occurs mainly from the proline subunit with 85% of the acid chloride carbonyl being retained in the ketone product (eqs 5 and 6).¹⁵ Given that the Ni(0) anhydride-insertion step (Scheme 1, 3→4) is likely reversible and nonregioselective (outside of small electronic and steric perturbations), we presume that the relative activation barriers of the two possible decarboxylation steps most likely dictate the observed 85:15 ratio of isotopically labeled products. This would seem reasonable given that proline carboxylate undergoes

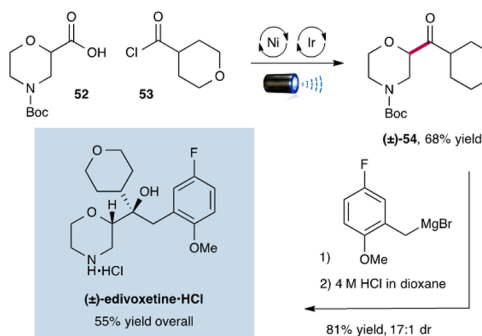
Rearrangement of Cyclopropylacetic Anhydride Substrate (Eq 5)

Extrusion Recombination with ¹³C-labeled Mixed Anhydride (Eq 6)Proposed Mechanism Based on Cyclopropyl ¹³C-labeling Studies (Eq 7)

oxidation–decarboxylation much faster than the corresponding hydrocinnamate, due to the relative stabilities of the resulting radical intermediates.¹⁶ As such, we presume that cyclopropyl ring-opened product 47 must arise from a Ni^{II}-mediated decarboxylation–recombination process, wherein formation of Ni^{II} complex 49 enables cyclopropyl ring opening prior to recombination.¹⁷ Given the wealth of previously reported Ni-

catalyzed carbonylation and decarbonylation reactions,^{18,19} we feel this mechanism best supports the results of our labeling experiments.

Finally, we applied our CO₂ExR technology to a three-step synthesis of (±)-edivoxetine-HCl, a medicinal agent in development for the treatment of ADHD.²⁰



As shown, commercial acid **52** and acyl chloride **53** were readily coupled using the optimized metallaphotoredox conditions to generate ketone **54** in good yield (68%). The synthesis was thereafter completed via a Grignard addition, followed by HCl-mediated Boc removal to afford (±)-edivoxetine-HCl in 55% yield over three steps.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.5b08304.

Experimental procedures, structural proofs, and spectral data for all new compounds (PDF)

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Notes

The authors declare no competing financial interest.

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