Evidence for Photocatalyst Involvement in Oxidative Additions of Nickel-Catalyzed Carboxylate O-Arylations

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ABSTRACT: Dual photocatalysis and nickel catalysis can effect cross-coupling under mild conditions, but little is known about the in situ kinetics of this class of reactions. We report a comprehensive kinetic examination of a model carboxylate O-arylation, comparing a state-of-the-art homogeneous photocatalyst (Ir(ppy)3) with a competitive heterogeneous photocatalyst (graphitic carbon nitride). Experimental conditions were adjusted such that the nickel catalytic cycle is saturated with excited photocatalyst. This approach was designed to remove the role of the photocatalyst, by which only the intrinsic behaviors of the nickel catalytic cycles are observed. The two reactions did not display identical kinetics. Ir(ppy)3 deactivates the nickel catalytic cycle and creates more dehalogenated side product. Kinetic data for the reaction using Ir(ppy)3 supports a turnover-limiting reductive elimination. Graphitic carbon nitride gave higher selectivity, even at high photocatalyst-to-nickel ratios. The heterogeneous reaction also showed a rate dependence on aryl halide, indicating that oxidative addition plays a role in rate determination. The results argue against the current mechanistic hypothesis, which states that the photocatalyst is only involved to trigger reductive elimination.

INTRODUCTION

Palladium catalysis is key to many conceptual and practical advances in modern organic chemistry, but the scarcity of palladium bodes poorly for long-term use. As such, transferring well-defined reactivities of palladium to nickel, an isoelectronic and more abundant metal, has been an explicit goal of catalysis for more than 50 years. Low-valent nickel species are capable of undergoing oxidative additions readily, but the subsequent reductive eliminations (REs), particularly for Ni(II) complexes, are thought to be comparatively difficult. Nickel couplings are hence believed to go through higher-valent intermediates, accessed by stoichiometric redox modulators, such that the RE is more favorable.

The combination of photocatalysis with nickel catalysis has provided an elegant pathway by which nickel can turn over without adding excessive reductants or harsh conditions. Iridium polypyridyl complexes were found to act as photocatalysts (PCs) to effect C–N, C–O, C–S, and C–C couplings using nickel(II) precatalysts. Subsequent reports showed that the homogeneous iridium photocatalysts can be replaced with heterogeneous graphitic carbon nitride (g-CN) semiconductors, rendering the overall transformations noble-metal-free.

The current mechanistic proposal for a relatively well-studied carboxylate O-arylation (Figure 1A) in this family of reactions is depicted in Figure 1B. Numerous common nickel(II) precatalysts are viable, which are thought to undergo two single-electron transfers to a bipyridyl Ni(0). From this reduced species, oxidative addition to form S is facile. The catalyst resting state 6 is unable to undergo the necessary reductive elimination. To close the cycle, 6 receives an energy transfer from the excited state of the photocatalyst, which enables C–O bond formation. Stern–Volmer quenching studies, transient absorption spectroscopy, and computational studies have been used to support quenching of Ir(ppy)3 by a potential nickel(II) intermediate.

Broadly, the substrate scopes for both homogeneous (Ir(ppy)3) and heterogeneous (g-CN) variations of carbon–heteroatom couplings contain only electron-poor aryl halides and electron-neutral or rich coupling partners (HNR2, HOR, HSR, etc.). To our knowledge, these limitations have not been explicitly discussed in a mechanistic context. The Ir(ppy)3-catalyzed homogeneous carboxylate O-arylation accommodates only electron-withdrawing aryl bromides, and electron-withdrawing carboxylic acids are absent. Such a combination could be explained by arguing that an SAr-type reductive elimination is required.

However, tension between the existing mechanism and the resulting substrate scopes surfaced during our development of the heterogeneous congener using graphitic carbon nitride...
photocatalysts. During a cursory kinetic study, methyl 4-iodobenzoate 1 (96%, 4 h) coupled considerably faster than methyl 4-bromobenzoate (88%, 48 h) with N-Boc-proline 2. Once the nickel catalytic cycle is at the resting state 6, the differences between an aryl iodide and bromide are irrelevant, as only the aryl ring remains on-cycle. This discrepancy in rate must therefore arise from the oxidative addition or ligand exchange step. As even aryl chlorides show facile proclivities toward oxidative addition with low-valent nickel in non-photochemical reactions and the homogeneous analogue converted aryl bromides efficiently, we began to question whether the currently accepted mechanism is operational with the heterogeneous congener. Given that we found the same electronic substituent limitations with g-CN as for Ir(ppy)₃, albeit with iodides instead of bromides, we further wondered if there were mechanistic commonalities among the two O-arylations.

Electron deficiency of an aryl halide can accelerate oxidative additions, as the aryl halide is itself formally reduced. In this vein, aryl halide limitations in this class of reactions could be explained by scission of the aryl halide bond that pushes at the energetic boundaries of photocatalytic reductions. Reduction potentials of common aryl halides range from −1.0 to −3.5 V vs SCE, with electron-donating groups and higher-row halides ranking as more difficult. The iridium polypyridyl complex Ir(ppy)₃, used in the homogeneous O-arylation has a reduction potential of −1.73 V, and substrate limitations could be a marker of the Ir(III)/Ir(IV) potential within the aforementioned range. Such logic has been invoked for the thioetherification of aryl iodides as well as dehalogenations with this exact photocatalyst. Stern–Volmer and transient absorption studies showed that aryl iodides could in fact quench the excited triplet state of Ir(ppy)₃ forming an aryl radical.

With these considerations in mind, we set out to answer three basic questions: (1) Do the heterogeneous and homogeneous reactions proceed by the same mechanism? If so, (2) do they both conform to the extant hypothesis, in which reductive elimination is turnover-limiting, and (3) are the turnover-limiting steps and the steps which require a photocatalyst the same?

**RESULTS**

**Heterogeneous (Graphitic Carbon Nitride) PC. Photon-Limited (3.33 mg/mL g-CN, 5 mM Ni L, 50% Lamp Power).** First, the kinetics of the carboxylate O-arylation of aryl iodides using the graphitic carbon nitride CN-OA-m as a photocatalyst in photon-limited conditions were analyzed. These photocatalytic reactions can be tracked by attaching a transient absorption studies showed that aryl iodides could in fact quench the excited triplet state of Ir(ppy)₃ forming an aryl radical. Data obtained in this regime allow us to both evaluate the intrinsic kinetics of the nickel cycle, as well as enable a more rigorous comparison between data sets of different photocatalysts.

As no in situ studies exist to date for any dual nickel-catalyzed and photocatalyzed cross-couplings, comparison of the photon-limited and photon-unlimited regimes with each other for the two reactions would be of general interest. We anticipated that we would see differences between the photon-limited and photon-unlimited regimes, potentially in the magnitude of the reagents' coefficients in the rate laws.

Most importantly, the intrinsic, photon-unlimited kinetics of both reactions should be compared with each other and to the extant mechanistic hypothesis. The kinetics in photon-unlimited regimes of the homogeneous and heterogeneous reactions should be identical since the PC should drop out of the rate law. The extant hypothesis for the dual photo- and nickel-catalyzed carbonate O-arylation, congruent with the accrued understanding of nickel catalysis, is that the PC enables a turnover-limiting reductive elimination. Thus, we predicted that the photon-unlimited regime for both reactions would give rate $-k_1[\text{Ni}]$.
From a cursory, qualitative glance, these reactions at the native conditions display no unusual behavior, except for exhibiting a short induction period. A series of experiments were conducted to assess which reagents were responsible for this induction (Figure S5). Delaying the injection of NiCl₂-glyme and its bipyridyl ligand or aryl iodide 1 resulted in immediate productive catalysis. Delayed injection of carboxylic acid 2 retained the induction period. Conspicuously, delaying addition of the base N-tert-butylisopropylamine (BIPA) resulted in a reaction profile slightly suggestive of catalyst activation, indicating that the secondary amine plays a role more significant than simply that of a Brønsted base.

Varying the nickel concentration from 5 to 20 mM unsurprisingly resulted in no rate change (Figure S6). At 5 mM Ni, no catalyst deactivation or product inhibition was observed (Figure S8), which is expected at such a high metal catalyst concentration. Surprisingly, modulation in [1] resulted in a fractionally positive order (Figure S9) for the aryl iodide. Carboxylic acid 2 showed a zero-order dependence (Figure S10). Reagent orders obtained in this regime should be considered with some measure of caution as the interactions between PC and Ni are ill-defined. However, we were curious if the reagent order of the aryl iodide would persist in the photon-unlimited regime, as it would be a significant clue to help resolve the questions posed above.

**Photon-Unlimited (3.33 mg/mL g-CN, 0.2 mM NiL, 100% Lamp Power).** To switch from the photon-limited to the photon-unlimited regime, we needed to increase the ratio of photons and/or PC to nickel. Although numerous photocatalytic reactions with metal centers have been tracked with *in situ* apparatuses, to our knowledge only one study has described finding a region in which the metal’s access to excitation is carefully controlled to be unimpeded. In the work of Lehnherr et al., light directly excited a metal catalyst, with no photocatalytic intermediary. To find a region in which their metal catalyst was not limited by throughput of photons, initial rate was plotted against catalyst concentration, and at lower concentrations a roughly linear correlation is apparent. To make a comparable plot, we retained a high PC loading and maximum lamp power (Figure 2A).

Moving from high to low nickel concentrations along this plot surprisingly did not provide an identical shape to that of the literature precedent. Specifically, we observed an unexpected “peak” where catalysis has a much higher rate, around 1 mM. Although further studies are required to account for this behavior, it is possible that, above a certain threshold, the metal catalyst engages in some combination of disproportionation and/or higher-order nickel species thought to be catalytically relevant. The area at very low concentration of nickel would likely provide a regime in which the nickel catalytic cycle was not limited by access to an excited photocatalytic species. To confirm this was the case, at low concentrations we used variable time normalization analysis (VTNA) to find a first-order dependence on nickel between 0.15–0.25 mM (Figure S11). Kinetic studies in this area are expected to produce data that reflect the intrinsic characteristics of the nickel cycle.

To probe the robustness of the nickel catalyst, two “same-excess” experiments were conducted under photon-unlimited conditions and the less concentrated was time-shifted on the x-axis (Figure 2B). Overlay here indicates that the product does not inhibit catalysis, nor does meaningful or disproportionate catalyst deactivation occur.

Experiments to determine the reagent orders of the substrates revealed that aryl iodide again exhibited a positive order dependence (Figure 2C). Due to the induction period, first attempts at VTNA overlay were found to be problematic (Figure S12). Elegant mathematical solutions to this common issue have been disclosed, but the most rigorous would require spectroscopic determination of the active catalyst concentration. Repeating the different excess experiments with delayed catalyst injections (Figure S13), which bypass the induction period as described above, cleanly leads to a reagent order of 0.3 under 150 mM [1].

![Figure 2](https://dx.doi.org/10.1021/jacs.0c02848)
Examination of the dependence on reagents’ concentrations for 2, the base, and the photocatalyst provided a much less straightforward picture. All three reagents exhibit positive order dependence at lower concentrations and inhibitory effects at higher concentrations (Figures S14–S16).

**Homogeneous (Ir(ppy)₃) PC.** We began the homogeneous studies by attempting to determine a proper photocatalyst loading that would enable photon-limited conditions. Initial experiments with 0.1 mM Ni·L and the standard (1 mM) photocatalyst concentration gave unsatisfactory yields. We observed dehalogenation accompanying productive catalysis (Table 1). Formation of the phenol side product 9 appears to be independent of an unfavorable PC-to-Ni ratio.

Table 1. Screening to Determine Photocatalyst Loading for Photon-Limited Regimes of the Homogeneous Reaction

<table>
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<tr>
<th>mM PC</th>
<th>3</th>
<th>1</th>
<th>8</th>
<th>9</th>
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<td>1</td>
<td>62</td>
<td>0</td>
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<td>6</td>
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<td>10</td>
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<td>94</td>
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<tr>
<td>0.125</td>
<td>90</td>
<td>0</td>
<td>5</td>
<td>5</td>
<td>100</td>
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</tbody>
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*Final concentrations of product and side products were determined by ¹H NMR using 1,3,5-trimethoxybenzene as an internal standard.*

High concentrations of Ir(ppy)₃ accelerate the consumption of 1 but not significantly the rate of product formation (Figure 3A). This finding is in striking contrast to the homogeneous analogue, where slower product formation is a function of slower consumption of 1 at higher PC loading (Figure S16). Experiments with 0.125 mM PC loading (Figure 3A, blue) appear to be limited by the relay of energy from PC to Ni (Figure S17), and are the basis for the photon-limited studies.

**Photon-Limited (0.125 mM Ir(ppy)₃, 0.1 mM Ni·L, 100% Lamp Power).** The first noteworthy qualitative observation of iridium-catalyzed O-arylations is the lack of induction period. Catalysis begins immediately in a positive-order fashion. Increasing concentration of nickel leads to increased rate, which was determined to be a first-order dependence by VTNA (Figure S18). A same-excess experiment shows no deactivation of the nickel throughout the course of the reaction (Figure S19). In contrast to the heterogeneous congener, no rate dependence was found for 1 or 2 (Figure S20). Interestingly, increasing concentration of base increases the rate (Figure S21). Again, some caution is advised in mechanistic interpretation from these data, as these conditions are designed to maximize yield. However, the lack of rate dependence on aryl iodide could be a significant finding if it persisted in photon-unlimited conditions.

**Photon-Unlimited (1 mM Ir(ppy)₃, 0.2 mM Ni·L, 100% Lamp Power).** In order to draw meaningful comparisons between the two-photon-limited data sets, we sought to use the same [Ni]₀ as used for the heterogeneous system. As such, finding the photon-unlimited regime for the homogeneous reaction occurred through a reverse process from the heterogeneous case. A short series of experiments was conducted in which the nickel concentration was kept constant and the photocatalyst concentration was raised, similar to...
heterogeneous reaction would have been observed, so we can attribute this behavior to catalyst deactivation.

Experiments varying the amounts of substrate and coupling partner were conducted to obtain the reagent orders of our reactants. In contrast to the heterogeneous reaction, we found the rate to be essentially independent of the concentration of 1 (Figure 3C). The dependence on [2] was not found to be significant (Figure S25). Last, we modified the amount of base while keeping all other reagents constant. Similar to the observation from the photon-limited data set, this change was the sole factor that accelerates the homogeneous reaction (Figure S26).

**DISCUSSION**

The most conspicuous finding of this study was the rejection of the hypothesis that the kinetics of both systems would give identical rate laws. In other words, as particular care was taken such that the nickel cycle was saturated with excited photocatalyst species for both systems, the intrinsic kinetics of a nickel cross-coupling cycle should have been identical between the two had the mechanisms been the same.

The chief quantitative finding of this study is the fractional dependence of aryl iodide 1 in the heterogeneous rate law. To some degree in the heterogeneous congener, oxidative addition is turnover-limiting. We propose that this is a result of direct photocatalytic activation of the substrate (Figure 4B). The fractional order arises as this activation occurs off of the nickel cycle. Nickel is first-order as it intercepts the radical species on the nickel catalytic cycle.

This process may have been overlooked because it is not necessarily turnover-limiting. The reduction potential of Ir(III)*/Ir(IV), −1.73 V, is sufficient to donate an electron to the aryl halide 1,25,42,43 this step is photocatalyst-dependent but fast for aryl iodides (Figure 4B). In contrast, the analogous excited state of the graphitic carbon nitride used in this study is less reducing (up to −1.65 V vs SCE). The capacity for electron transfer is worse, and therefore even in conditions that can be considered photon-unlimited, formation of oxidative addition intermediate 5 is still limited by the kinetically relevant halide fragmentation.

We reject the hypothesis that photocatalysts are required only as a late-cycle trigger for RE. We contend that the photocatalyst is required for at least one additional point on the catalytic cycle, namely for single-electron transfer to the aryl halide. Substrate limitations for dual photo- and nickel-catalyzed carbon−heteroatom cross-couplings derive from this elementary step. The most obvious example is the far superior conversion of aryl bromides in this homogeneous O-arylation compared to the heterogeneous congener. Such a framework also harmonizes the mechanisms by which these transformations occur with the existing body of photocatalytic dehalogenations and radical-based transformations.25,26

To augment our understanding of the heterogeneous reaction, we created a Hammett plot examining rate dependencies on aryl iodides with substituents of varying electronic character (Figure 4C). Qualitatively, this shape is also seen for the oxidative addition of tris(triphenylphosphine)nickel(0) to aryl chlorides.20 Our ρ value (2.1) is far below that of this more traditional oxidative addition (8.8), thought to proceed through an S_NAr-type mechanism. This could be explained by differences in ligand systems and halides, but also could point to a lowered dependence on the electronic substituent due to a non-concerted oxidative addition.

Figure 4. (A) Key findings of this study. (B) Updated mechanistic proposal. Secondary amine base is proposed to close the photoredox cycle by serving as single electron donor (SED). (C) Hammett plot for heterogeneous congener supports that oxidative addition is to some degree turnover-limiting.
With a rate law proportional only to the nickel concentration, data for the homogeneous reaction could reasonably fit three turnover-limiting steps: a unimolecular rearrangement, ligand dissociation, or reductive elimination. We have observed nothing to support or reject the first two. Given that the incumbent hypothesis is that RE is turnover-limiting, our data support the existing proposals. As we had originally predicted, our kinetic data here likely reflects the limitations inherent to nickel catalysis.

Whether or not 6 requires a photocatalyst to trigger RE likely depends on the oxidation state of this intermediate. It has been clearly demonstrated that a Ni(II) resting state cannot undergo RE without photocatalytic excitation.17-19 However, extensive literature precedent points to the viability of multiple nickel oxidation states in cross-coupling reactions across very different conditions,35,44,45 and the oxidation state of the on-cycle intermediate for these photochemical reactions has yet to be convincingly resolved. Should this intermediate require activation via energy/electron transfer in the homogeneous reactions, it would not be turnover-limiting, given the photon-unlimited environment.

Of note is the fact that the only way to accelerate the homogeneous reaction was by addition of more secondary amine base, which is already in large excess. We offer three potential explanations. First, and most likely, the secondary amine donates an electron to [Ir(ppy)3]+ to regenerate the photocatalytic cycle. We can assume that the concentration of 2 requires equimolar amounts of base for deprotonation, and extra amine beyond that point fulfills various roles including weak ligand and electron source. Second, the base is potentially required activation via energy/electron transfer in the homogeneous reactions, it would not be turnover-limiting, given the photon-unlimited environment.

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Qualitatively, two main observations stand out for iridium-catalyzed carboxylate O-arylations. First, this system inherently tends to be less selective. In the photon-unlimited regime, at peak catalytic activity was found at a much lower concentration of nickel than used in method 6. Additionally, peak catalytic activity was found at a much lower concentration of nickel than used in method 6. The choice of photocatalyst may, among other things, be determined by the electronic character of the substrate, as less dehalogenation was observed in the g-CN case. We believe that photocatalysts with weaker oxidation potentials, such as graphitic carbon nitrides, are less likely to participate in a Ni(0)/Ni(II) cycle.

**CONCLUSION**

Efforts to swap photocatalysts in dual photocatalytic and metal catalytic systems may not preserve the inherent reactivity of the metal catalytic cycle. Homogeneous and heterogeneous esterifications differ primarily in that the homogeneous congener showed a rate dependence on the aryl halide substrate. We ascribe this to the nickel trapping an off-cycle radical anion generated by the photocatalyst and the aryl halide. In this sense, the requirements for electron-withdrawing substrates may be ascribed to an energetic barrier that is independent of the nickel catalytic cycle. Overcoming such limitations likely must occur through photocatalyst development and tuning to maximize the reduction potential of the excited PC. Alternatively, methods to front-load the energetic relay from external source to photocatalyst to substrate show great promise. However, as our PC and nickel screenings to find different regimes show, using a PC that is too reducing for the chosen substrate and/or metal catalyst can lead to over-reduction and catalyst deactivation. Countering this requires careful modulation of PC and metal catalyst concentrations. Hence, for the Ir(ppy)3-catalyzed O-aryl shown here, a photon-limited approach is desirable. Optimization of the graphitic carbon nitride-catalyzed arylation shown here, a photon-limited approach is desirable. Optimization of the graphitic carbon nitride-catalyzed arylation shown here, a photon-limited approach is desirable. Optimization of the graphitic carbon nitride-catalyzed arylation shown here, a photon-limited approach is desirable.
Complete contact information is available at: https://pubs.acs.org/10.1021/jacs.0c02848

Notes

The authors declare no competing financial interest.

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(22) All redox potentials in this article are given in reference to the saturated calomel electrode (SCE).

(23) These values may not serve as a hard limit and are often found to be surpassed in reaction conditions. See the following for a seminal discussion, and ref.s 24 and 25 for examples: King, K. A.; Spellane, P. J.; Watts, R. J. Excited-State Properties of a Triply Ortho-metalated Iridium(III) Complex. J. Am. Chem. Soc. 1985, 107, 1431–1432.


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