

Two Paths to Oxidative C–H Amination Under Basic Conditions: A Theoretical Case Study Reveals Hidden Opportunities Provided by Electron Upconversion**

Paul Eckhardt,^[a] Quintin Elliot,^[b] Igor V. Alabugin,^{*,[b]} and Till Opatz^{*,[a]}

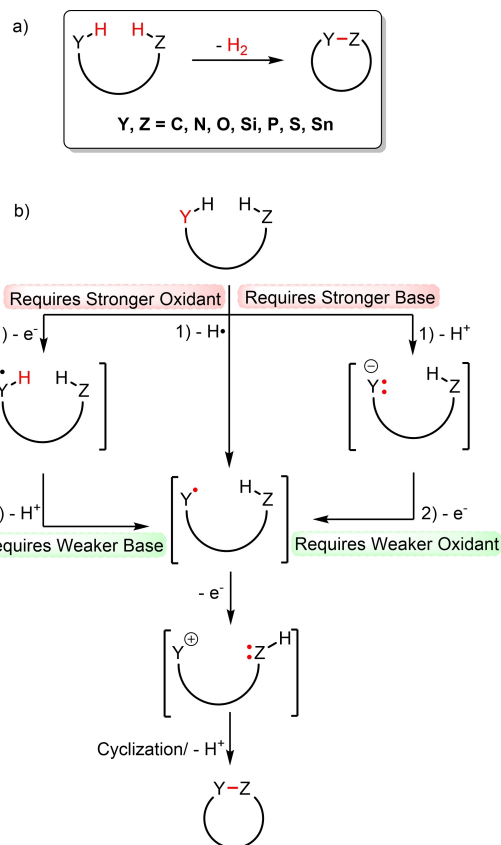
Abstract: Traditionally, cross-dehydrogenative coupling (CDC) leads to C–N bond formation under basic and oxidative conditions and is proposed to proceed via a two-electron bond formation mediated by carbenium ions. However, the formation of such high-energy intermediates is only possible in the presence of strong oxidants, which may lead to undesired side reactions and poor functional group tolerance. In this work we explore if oxidation under basic conditions allows the formation of three-electron bonds (resulting in

“upconverted” highly-reducing radical-anions). The benefit of this “upconversion” process is in the ability to use milder oxidants (e.g., O₂) and to avoid high-energy intermediates. Comparison of the two- and three-electron pathways using quantum mechanical calculations reveals that not only does the absence of a strong oxidant shut down two-electron pathways in favor of a three-electron path but, paradoxically, weaker oxidants react faster with the upconverted reductants by avoiding the inverted Marcus region for electron transfer.

Introduction

Initially, cross-dehydrogenative coupling (CDC) described transition metal catalyzed reactions developed to form C–C bonds through oxidative coupling of two C–H units.^[1] Over the years CDC has evolved into a universal tool for the construction of Y–Z bonds starting from Y–H and Z–H units (Scheme 1a). The reaction may proceed either intermolecularly or intramolecularly in the coupling of Y and Z, which include a carbon and a heteroatom (e.g., N, O, Si, P, S, Sn).^[2]

As the scope of CDC reactions widened, they have been performed under a broader variety of conditions through a number of mechanistic pathways, even under metal-free conditions.^[2a,3] In the absence of metals, a variety of scenarios for the formal loss of H₂ are possible (Scheme 1b). All of them involve a combination of several of the following elementary



Scheme 1. Schematic representation of possible pathways for CDC reactions with the overall formal loss of H₂ via a two-electron bond formation. More combinations are possible (see below).

steps: deprotonation, oxidation, H-atom abstraction, and hydride abstraction.

[a] P. Eckhardt,⁺ Prof. Dr. T. Opatz
Department of Chemistry
Johannes Gutenberg University Mainz
Duesbergweg 10–14, 55128 Mainz (Germany)
E-mail: opatz@uni-mainz.de

[b] Q. Elliot,⁺ Prof. Dr. I. V. Alabugin
Department of Chemistry and Biochemistry
Florida State University
Tallahassee, Florida 32306 (USA)
E-mail: alabugin@chem.fsu.edu

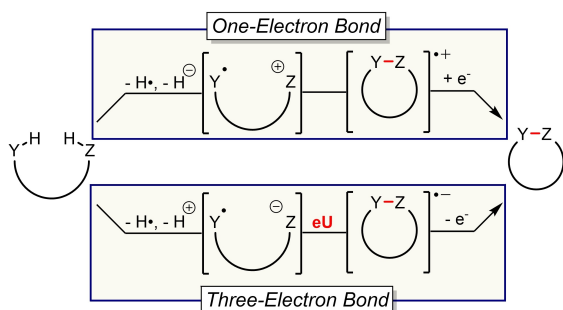
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This is accomplished by activation of Y–H in the presence of the nucleophilic Z–H moiety. Most often, this bond is suggested to form via a reaction between a cation and a nucleophile. Methods for cation formation include formal hydride loss accomplished by removal of either a hydrogen atom and an electron,^[4] or a proton and two electrons.^[5] Occasionally, the Y–Z bond can be formed via recombination of two radical centers, Y and Z.^[6] Although multiple pathways exist to reach the final products, the commonly suggested scenarios for Y–Z bond formation mostly involve formation of a classic two-electron bond as the last step to conclude the CDC cascade. However, these pathways do not paint the whole picture and there are additional approaches to consider.

Conceptually different mechanisms utilize an odd number of electrons for the Y–Z bond formation (Scheme 2).^[7] For example, a two-center one-electron bond (2c,1e) may be created by a radical and cation formed by the removal of a hydrogen atom and a hydride from the reactant.^[8] Alternatively, removal of a hydrogen atom and a proton allows for the coupling of a radical and anion to form a two-center three-electron (2c,3e) bond.^[7c,9] Such bonds, first described by Linus Pauling, are known to play many roles in chemical structure and reactivity (e.g., contribute greatly to the stability of triplet oxygen, account for the weakened BDEs of α -C–H bonds in ethers, aldehydes, and alcohols,^[10] lead to radical and oxy-anionic fragmentations, etc).^[11] An intriguing, but not always appreciated, feature of such bonds is that their formation can lead to “electron upconversion” (eU),^[12] a conceptually interest-



Scheme 2. Schematic representation of odd-electron bond formations. Top: One-electron bond formed between a radical and a cation. Bottom: Three-electron bond formed between a radical and anion.

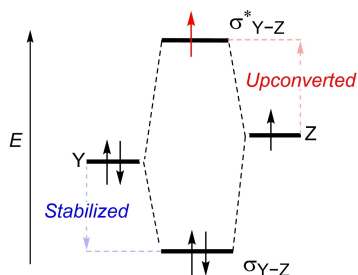


Figure 1. Schematic representation of electron upconversion. The upconverted electron is highlighted in red.^[12b,c]

ing phenomenon of direct importance for the design of “electron-catalyzed” reactions.^[13]

Electron upconversion may seem like a paradox or even a violation of the laws of thermodynamics because it transforms a relatively weak reductant into a *more* potent reductant in a thermodynamically favorable manner.^[11e,14] This behavior contrasts the course of typical exergonic redox reactions where the reactants are stronger reductants than the product. Electron upconversion is made possible by moving (“upconverting”) one electron from a *non-bonding* orbital into a higher energy *anti-bonding* orbital via a 2c,3e bond (Figure 1).^[12c] Because of its high energy, the upconverted electron can be readily removed via a facile one-electron transfer to a very mild oxidant, thus completing the process of a Y–Z bond formation

The selective amplification of reductive potential by creating the high-energy upconverted singly occupied MO (SOMO) could be used for the construction of reaction sequences which may not tolerate stronger oxidants.

For example, Alabugin and co-workers have demonstrated two variants of electron upconversion in C–H amination of unprotected anilines and amides. In these examples, the (2c,3e) half-bond is formed by a nitrogen-anion and a carbon radical center.^[12b,c] Because the intermediate containing the (2c,3e) half-bond has a much higher reduction potential than other species present in the reaction, it can be selectively oxidized by a relatively mild oxidant, for example, molecular oxygen. Additional examples of redox chains mediated by highly reducing upconverted radical anion intermediates continue to appear, for example, a photochemical thiol-yne click reaction reported by Ananikov et al.,^[15] hydroxylation reaction of aryl halides with oximes by James et al.^[16] and a number of other transformations.^[17] Recently, Jui and co-workers have demonstrated a facile transformation of formate into upconverted $\text{CO}_2^{\bullet-}$ species (a traditionally challenging reductant to make, $E_{1/2} = -2.2$ V vs. SCE) capable of reducing a broad range of substrates.^[18]

Herein, we will use quantum-mechanical calculations to investigate whether a three-electron bond formation with electron upconversion can be a viable alternative for these transformations. In particular, we will evaluate literature C–N bond forming reactions proposed to proceed via a two-electron pathway (e.g., a nitrogen nucleophile attacking a carbenium ion). This analysis is guided by a hypothesis that the preferred reaction path can be switched from a two-electron pathway to a three-electron pathway when a mild oxidant is used instead of a stronger one.

Computational methods

Density-functional theory (DFT) calculations were carried out using the GAUSSIAN 16 Revision A.03 program package.^[19] Geometry optimizations and frequency calculations were carried out at the (U)M06-2X/6-31 + G(d,p) level of theory^[20] including Grimme’s D3 correction for empirical dispersion^[21] and the implicit SMD solvation model^[22] to simulate the effects of acetonitrile (MeCN), dibutyl ether (DBE), *N,N*-dimethylformamide (DMF) or dimethyl sulfoxide (DMSO). DBE (ϵ around 3.0)^[23] was chosen as the solvent because of the

similar dielectric constant to methyl tert-butyl ether (ϵ around 2.6),^[24] which was used as the main solvent in the work of Xu et al. (see below) but is presently not implemented in the GAUSSIAN program package.^[25] DMF (ϵ 37.2)^[26] was chosen based on the earlier experimental work.^[12b,c] Tight cutoffs on forces and step size were used (opt = tight). Frequency calculations were used to identify the structures as minima or transition states (TS). Unless otherwise stated, the presented results were produced at the (U)M06-2X(D3)/6-31+G(d,p)/(opt=tight)/(SMD) level of theory. CAS(3,3) and DLPNO-CCSD(T)^[27] single point calculations were carried out using the ORCA 4.2.1 program package,^[28] utilizing the 6-31+G(d,p) basis set^[20d] (CAS(3,3)) and the cc-pVTZ basis set^[29] (DLPNO-CCSD(T)) with the TightSCF and TightPNO keywords and the optimized geometries obtained from DFT calculations. Three-dimensional depictions and spin density plots were produced with Chemcraft 1.8.^[30] The procedure for the calculations regarding Marcus theory and further details can be found in the Supporting Information.

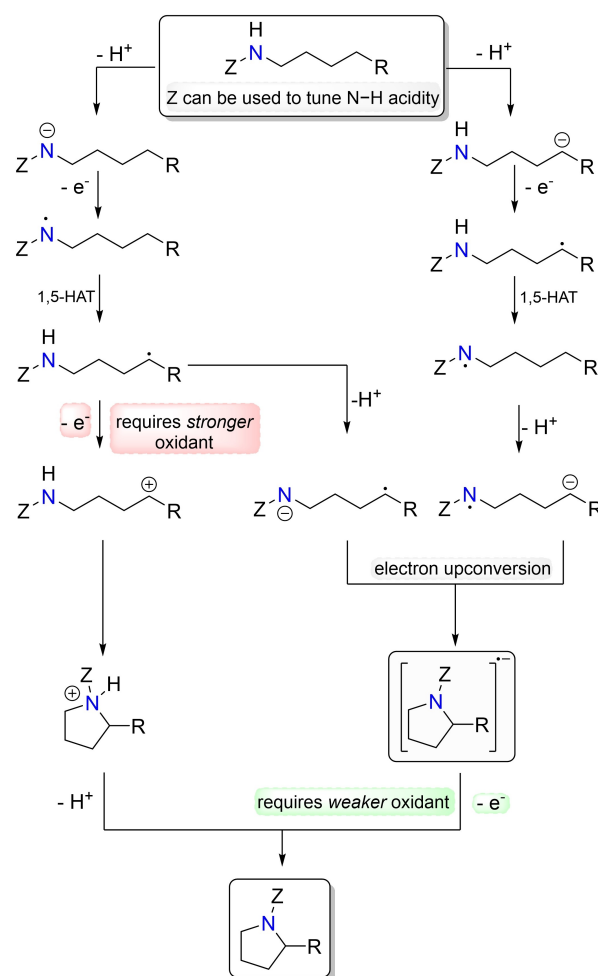
Results and Discussion

Our focus will be on representative C–N bond-forming cyclization reactions under basic conditions in the presence of an oxidant. Commonly, the proposed C–N bond forming steps either involve formation of a high energy carbenium ion (a step that requires a strong oxidant) or alternatively, a vague “bond forming step”. To address such mechanistic ambiguities, we sought to evaluate the relative thermodynamic profiles for the two main C–N bond forming paths, i.e., the cation/nucleophile (two-electron) and the anion/radical (three-electron) pathways (Scheme 3).

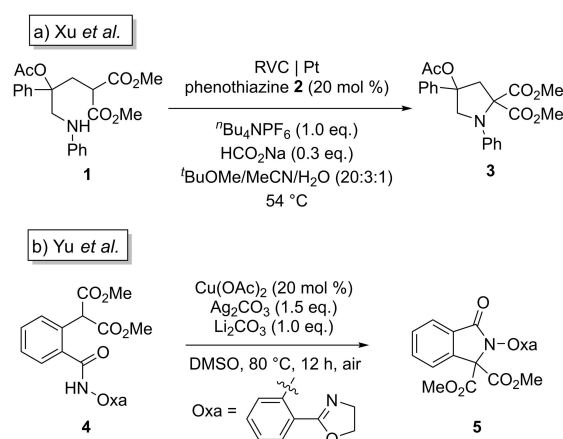
Oxidative C–N bond formation has to involve a formal loss of H_2 . Under basic conditions, this process is usually accomplished via the loss of two protons and two electrons.^[25,31] However, the sequence of these four elementary steps can be accomplished *in different orders*. A typically proposed reaction mechanism involves the sequence of deprotonation/oxidation followed by oxidation/deprotonation.^[5e] This is the sequence that leads to the formation of a carbenium ion which is then intercepted by a nitrogen nucleophile (left pathway of Scheme 3).^[32] Although this process is plausible, it has a drawback because the conversion of a radical into a cation generally requires a strong oxidant.^[33]

An alternative radical pathway proceeds via the sequence of deprotonation/H-atom abstraction with cyclization/oxidation as the final step. In this process, the C–N bond is formed at an intermediate stage via the reaction of an anion and a radical. An interesting feature of this step is that it upconverts an electron, i.e., transforms a *weakly reducing anion* into a *strongly reducing radical anion* (right pathway of Scheme 3).^[12a] By taking advantage of upconversion, one can perform the same reaction with milder oxidants.^[34] This feature can be valuable from a practical point of view. For example, the use of less aggressive oxidative conditions may allow for the preservation of sensitive functional groups.

Scheme 4 shows examples of C(sp³)–H/N–H cross-coupling reactions under basic and oxidative conditions. Both examples have malonate moieties reacting as pronucleophiles.^[25,35] No



Scheme 3. General mechanism for C–N bond formation under basic and oxidative conditions. Left pathway shows a commonly proposed carbenium ion intermediate. Right pathway shows a conceptually different approach (eU mechanism) that requires milder oxidative conditions to reach the same product.



Scheme 4. The selection of literature cyclization reactions which may proceed via eU.^[25,35]

specific C–N bond forming step was proposed, opening the doors for the mechanistic analysis including a potential 2c,3e cyclization.

To evaluate competition between the two mechanistic pathways, we have chosen to explore the reaction energies for the cyclization disclosed earlier by Xu et al. (Scheme 4a).^[25] The reaction was performed under electrochemical conditions with phenothiazine **2** as an electron transfer mediator and sodium hydroxide (formed in situ) as a base.

A simplified model (**6**) of compound **1** devoid of the phenyl and the acetoxy group was chosen as a starting point for a computational analysis. We compared the change in Gibbs free energy of three (commonly proposed) possible reaction pathways and utilized Marcus theory (see Supporting Information) to calculate the transition states for electron transfer reactions (shown in different colors in Scheme 5).

All three pathways start from a common sequence. After the initial deprotonation of **6** by hydroxide ($\Delta G = -32 \text{ kcal mol}^{-1}$), oxidation of the carbanion **7** by phenothiazine **2** radical-cation ($E_{p/2} = 0.52 \text{ V vs. SCE}^{[25,36]}$) has no barrier and results in the formation of radical **8** ($\Delta G = -36 \text{ kcal mol}^{-1}$).

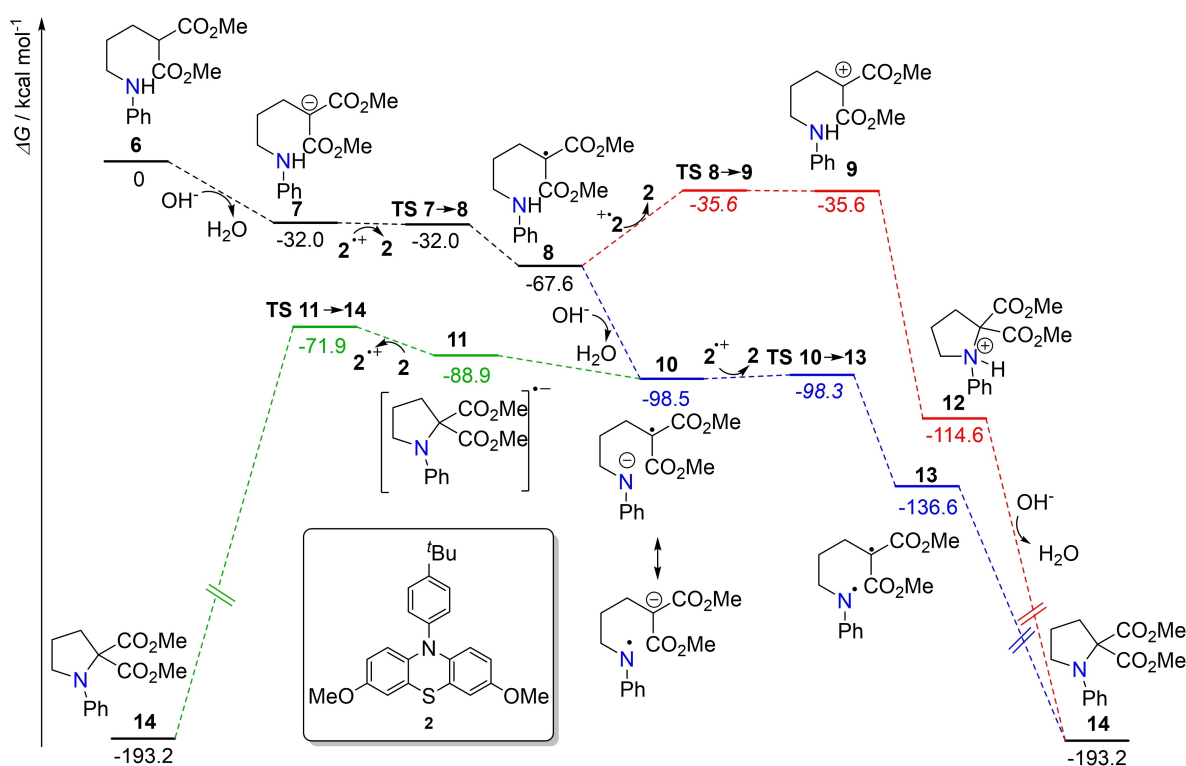
It is at this point that the reaction pathway diverges. The first (red) pathway is oxidation of the carbon-centered radical **8** to carbenium ion **9** ($\Delta G^\ddagger = 32 \text{ kcal mol}^{-1}$, $\Delta G = 32 \text{ kcal mol}^{-1}$). This step is thermodynamically unfavorable due to the formation of a carbenium ion next to two ester groups. Alternatively, the second electron may be removed from the lone pairs on nitrogen, thus, forming a carbon radical and

nitrogen radical-cation. Once the cationic species **9** is formed, the intramolecular nucleophilic attack by the amine is highly favored ($\Delta G = -79 \text{ kcal mol}^{-1}$). Subsequent deprotonation to form the final product **14** is downhill by $-79 \text{ kcal mol}^{-1}$.

The overall sequence of steps in the red pathway is the consecutive removal of $\text{H}^+/\text{e}^-/\text{e}^-/\text{H}^+$. The second oxidation has to proceed immediately after the first one, therefore requiring an oxidant that is strong enough to form a high energy cationic intermediate. Furthermore, the two sequential oxidations may require two separate oxidants to come into proximity to the reactant. When oxidants are present in low concentration, such sequential oxidation (while potentially thermodynamically favorable) may be slow. A portion of the electron transfer mediator will be present in the reduced form and unable to accept an electron. Additionally, oxidation of an anion is thermodynamically favored over the oxidation of a radical.

During the initial stages of the reaction, the concentration of anions will be higher than the concentration of radicals. As a result, it is more likely that an oxidant will come into proximity of an anion instead of a radical. Of course, the likelihood of an oxidant coming into proximity to a radical may increase as the concentration of anions dwindles and the concentration of radicals grows. However, radicals are often less persistent than anions. Hence, instead of patiently waiting for an oxidant to take an electron, they may undergo alternative reactions.

The second (blue) pathway proceeds via N–H deprotonation of **8** with the in situ-generated hydroxide anion to form the acyclic radical anionic species **10**. Such deprotonation is even



Scheme 5. Change in free energy and transition states (in kcal mol^{-1}) starting from **6** for three different reaction pathways. Red pathway: Oxidation resulting in carbenium ion formation, followed by cyclization, and deprotonation. Blue pathway: Deprotonation, oxidation, resulting in diradical formation, followed by cyclization. Green pathway: Radical anion cyclization and oxidation of the upconverted electron. Calculations performed for $T = 327.15 \text{ K}$, SMD = DBE.

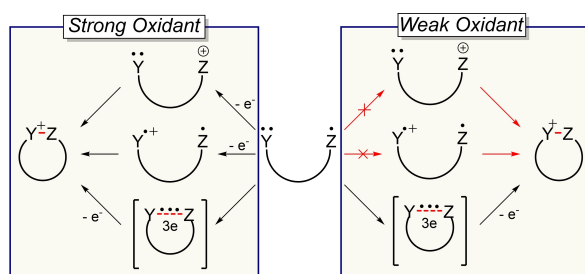
more thermodynamically favorable under these reaction conditions ($\Delta G = -31 \text{ kcal mol}^{-1}$) than the formation of carbenium ion **9** ($\Delta G = 32 \text{ kcal mol}^{-1}$). Upon formation of the acyclic radical anion, all that is needed to reach the final product is a one-electron oxidation and C–N bond formation (cyclization). The two subsequent (blue and green) pathways are different in the order of these steps.

Continuing along the blue path, a direct oxidation of **10** by another molecule of 2^{*+} gives rise to a biradical **13** which can be formed either as a singlet or a triplet ($\Delta G^\ddagger = 0.2 \text{ kcal mol}^{-1}$, $\Delta G = -38 \text{ kcal mol}^{-1}$ for the triplet product). If the triplet is formed, the diradical has to undergo intersystem crossing (ISC) into a singlet diradical with a subsequent highly favorable radical-radical recombination ($\Delta G = -56 \text{ kcal mol}^{-1}$) that yields the final product **14**.

In the absence of a strong external oxidant, the radical anion **10** can follow the final (green) pathway. In this scenario, the cyclization proceeds first to form the cyclic radical anionic species **11**. Although this step is mildly endergonic ($\Delta G = +9.6 \text{ kcal mol}^{-1}$), the sequential oxidation of the upconverted electron via 2^{*+} is highly favored ($\Delta G = -104 \text{ kcal mol}^{-1}$). However, due to the highly exergonic nature of this electron transfer, the transition state resides in the Marcus inverted region ($\Delta G^\ddagger = 17 \text{ kcal mol}^{-1}$).

This last reaction pathway is intriguing as it provides the most exergonic oxidation. Removing the upconverted electron is a staggering $136 \text{ kcal mol}^{-1}$ more favorable than carbenium ion formation and 66 kcal mol^{-1} more favorable than formation of a triplet biradical.

Depending on the reactants used, formation of a carbenium ion and/or diradical are only possible in the presence of a strong oxidant. In the absence of such an oxidant, these reactive intermediates may not form (see below). Utilizing such potent oxidants opens the door for undesired oxidation processes, possibly leading to side reactions. Conversely, oxidation of the upconverted electron becomes the only viable pathway in the absence of a strong oxidizing agent (Scheme 6). For further insight into the mechanism, we continued our studies with a weaker oxidant, molecular oxygen, and by searching for possible transition states for the cyclization steps.



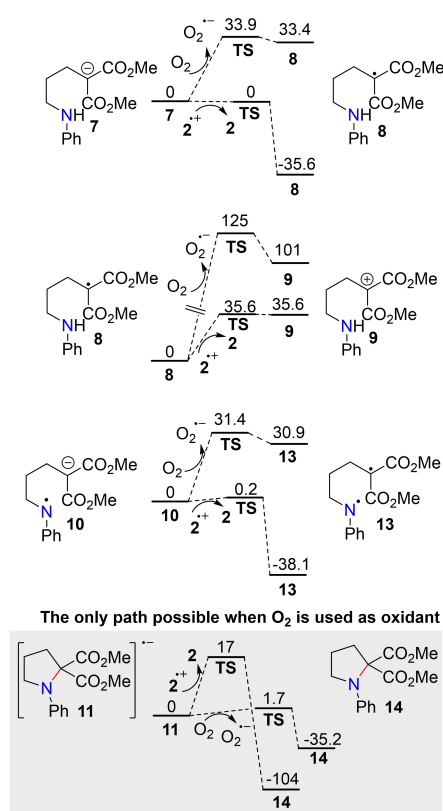
Scheme 6. Left: A strong oxidant may be used to form two-electron bonds (via carbenium ions/diradicals) or may be used to oxidize an upconverted reductant. Right: Use of a weak oxidant shuts down those paths which proceed via a two-electron bond formation. Removal of an upconverted electron is still possible.

Achieving selectivity via a three-electron path

The qualitative scenarios suggested above were confirmed by computational data. An illustrative comparison for the formation of a radical, carbenium ion, and diradical via oxidation using 2^{*+} and O_2 as alternative oxidant is given in Scheme 7.

While previously favorable with 2^{*+} , the use of O_2 as an oxidant makes the oxidation of the carbanion **7** to a carbon-centered radical unfavorable by 33 kcal mol^{-1} ($\Delta G^\ddagger = 34 \text{ kcal mol}^{-1}$). In the absence of a strong oxidant, the oxidation of the carbon radical **8** to form carbenium ion **9** is even more uphill ($\Delta G^\ddagger = 125 \text{ kcal mol}^{-1}$, $\Delta G = 101 \text{ kcal mol}^{-1}$). An important alternative to consider is the possible oxidation of acyclic radical anion **10** to form diradical **13**. However, the calculated free energy of the oxidation of **10** with molecular oxygen to the corresponding biradical (triplet) is still highly unfavorable ($\Delta G^\ddagger = 31 \text{ kcal mol}^{-1}$, $\Delta G = 31 \text{ kcal mol}^{-1}$). This rather gloomy thermodynamic landscape renders the highly exergonic electron transfer from the cyclic radical anion of **11** to the oxidant O_2 ($\Delta G^\ddagger = 1.7 \text{ kcal mol}^{-1}$, $\Delta G = -35 \text{ kcal mol}^{-1}$) particularly striking! The contrast in the reducing power between the *upconverted cyclic* radical anion **11** and the distonic *acyclic* radical anion **10** is dramatic ($\sim 66 \text{ kcal mol}^{-1}$ difference!).

In summary, when O_2 is used as an oxidant, only the 3e-path is thermodynamically viable. Furthermore, as an added bonus, the rate of electron transfer to the milder oxidant is predicted to be much faster under these conditions than the



Scheme 7. Direct comparison of oxidation steps between 2^{*+} and O_2 as oxidants. SMD = DBE, all energies reported in kcal mol^{-1} .

rate of electron transfer to the stronger oxidant (1.7 vs. 17 kcal mol⁻¹ free energy barriers for O₂ and 2^{•+}, respectively). This trend may seem counterintuitive as the much more exergonic electron transfer reaction is slower. Such observation suggests that the use of weaker oxidants for the upconverted reactants moves the electron transfer potential energy surface from the kinetically unfavorable Marcus inverted region into a more normal Marcus region.^[37]

However, there is a question of whether this path will be kinetically accessible. In order to address this question, we have calculated representative activation barriers for such reactions. Such calculations are extremely interesting from a fundamental perspective because they provide insight into the electronic and steric effects for the formation of 3e-bonds, a fundamentally important but underutilized process.

Full energy profile of the three-electron pathway

The thermodynamic landscape for the transformation of **6** to **14** was reinvestigated using previously reported conditions (Scheme 8)^[12c] DMF, *tert*-butoxide, and O₂ have been shown to initiate radical anion formation via N–H deprotonation and hydrogen atom transfer (HAT) mediated by the DMF radical. Experimental and computational evidence suggests the DMF radical to be formed in small concentrations under these conditions.^[12c]

In the first step, the deprotonation ($\Delta G = -21$ kcal mol⁻¹) of the C–H acidic dimethyl malonate (pK_a = 15.9 in DMSO)^[38] is favored because aniline is less acidic (pK_a = 30.7 in DMSO).^[39] Hydrogen atom abstraction by a DMF radical^[12b,c] allows for the formation of a nitrogen-centered radical ($\Delta G = -5.2$ kcal mol⁻¹). The acyclic radical anion **10** can undergo an endothermic cyclization ($\Delta G = +8.7$ kcal mol⁻¹) via **TS2** with a relatively low

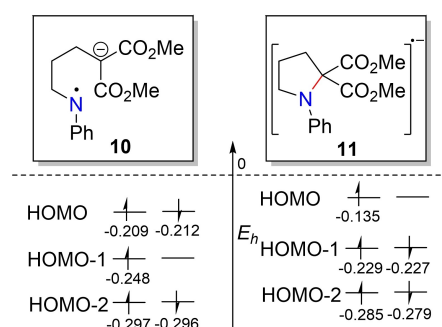


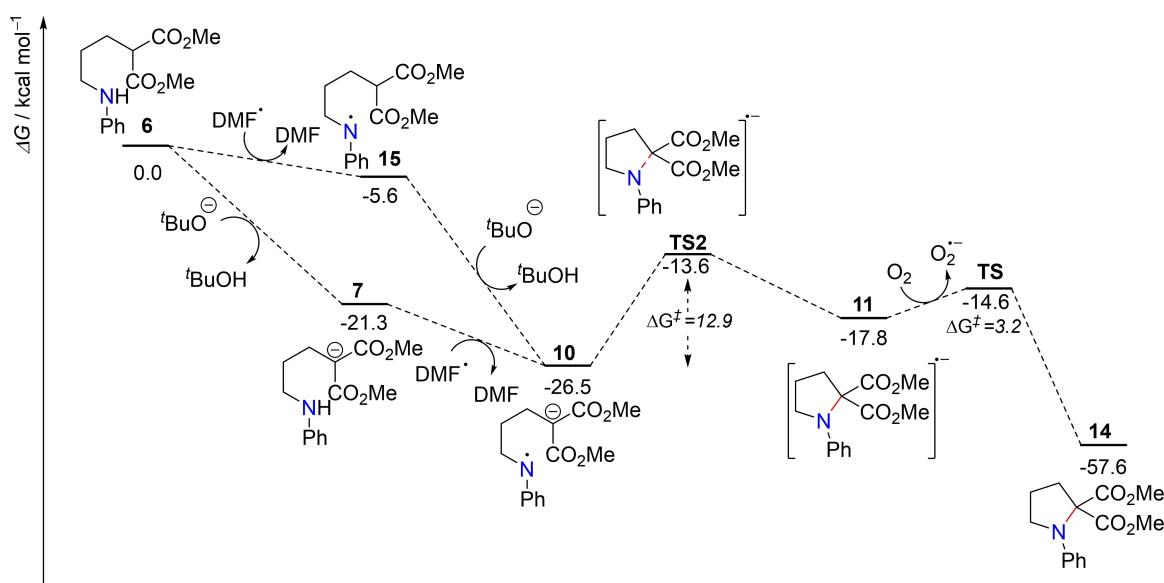
Figure 2. Molecular orbital energies in hartree (E_h) for acyclic and cyclic radical anions **10** and **11**.

activation barrier of 12.9 kcal mol⁻¹. The cyclic radical anion **11** can be readily oxidized by molecular oxygen to form the final product **14** ($\Delta G^\ddagger = 3.2$ kcal mol⁻¹, $\Delta G = -40$ kcal mol⁻¹) and superoxide.

Electronic structures

To gain additional insight into why acyclic radical anion **10** is a weaker reductant than cyclic radical anion **11**, it is illustrative to consider their respective molecular orbital energies (Figure 2) and to evaluate how well the simple model presented earlier for atomic orbitals (Figure 1) is transferrable to the more complex systems.

We found that the energetically highest alpha and beta electrons in **10** have an energy of -0.209 and -0.212 hartrees (E_h), respectively. HOMO-1 was found to only contain an alpha electron with an orbital energy of $-0.248 E_h$. Therefore, a HOMO-SOMO inversion is present in the acyclic radical anion



Scheme 8. Changes in free energy when using *tert*-butoxide as a base, DMF radical as a hydrogen atom transfer reagent, and O₂ as the oxidant^[12c] (U)M06-2X(D3)/6-31 + G(d,p)/SMD = *N,N*-dimethylformamide/opt = tight.

10.^[40] On the other hand, radical anion **11** was found to have all of its orbitals destabilized (when compared to acyclic radical anion **10**). The HOMO underwent notable destabilization and only contained a single (alpha) electron with an orbital energy of $-0.135 E_h$ (a difference of $\sim 46 \text{ kcal mol}^{-1}$ compared to the HOMO of **10**). HOMO-1 contained both an alpha and a beta electron at -0.229 and $-0.227 E_h$, respectively.

This direct comparison (Figure 2) provides some quantitative insight into the nature of electron upconversion. Both **10** and **11** are radical anions, the difference however is that the cyclic product **11** contains an electron in a high energy antibonding orbital. As a consequence, the molecular orbital containing this upconverted electron is raised in energy, rendering the reactant a more potent reducing agent. Meanwhile, **10** only contains electrons in lower lying non-bonding orbitals. As a result, the acyclic radical anion is a weaker reducing agent (compared to **11**).

Flow of spin density during the cyclization

The evolution of spin density (green) throughout the cyclization (Figure 3) reveals the change in localization of the unpaired electron. In **10**, spin density is located on the nitrogen and the aromatic ring. In **TS2**, spin density shifts to nitrogen and one of the ester carbonyl groups. Specifically, spin density is located in the carbonyl group with π orbitals that are parallel to the C–N bond being formed. Meanwhile, the carbonyl with a π orbital that is orthogonal to the newly formed C–N bond does not participate in spin delocalization.

In the cyclized species **11**, the entire spin density is located at the carbonyl group, the π orbital of which was parallel to the newly formed C–N bond in **TS2**, producing a ketyl-type radical. Accordingly, the SOMO of **11** has a strong contribution from the carbonyl group. In other words, the upconverted electron resides in a π^* orbital of one of the ester groups.

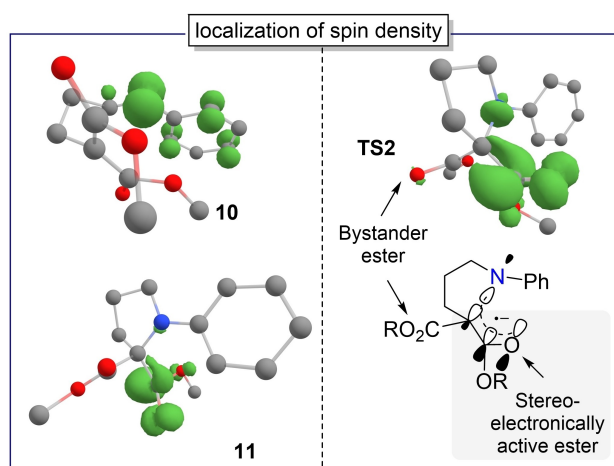


Figure 3. Spin density (green) plot obtained from DFT calculations of **10**, **TS2**, and **11**. Allylic type spin delocalization in **TS2**. Contour value: 0.008. Hydrogen atoms are omitted for clarity.

The π -system of the “bystander” ester is orthogonal to the incipient C–N bond and is stereoelectronically inactive in radical stabilization. In contrast, the second ester is stereoelectronically activated by rotation which aligns it with the forming C–N bond. As result, its carbonyl moiety participates in an allylic type delocalization of spin density at the α -carbon.

The spin density was also compared to the spin densities obtained from CAS(3,3) and DLPNO-CCSD(T) single point calculations and found to be in agreement.

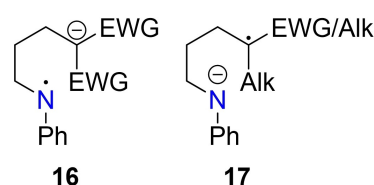
Substitution effects on the free energy of the cyclization

To evaluate the effect of substituents, the reaction energies of the cyclization of radical anions related to **10** were calculated.

Depending on the substituents on the δ -carbon and on nitrogen, two different electron arrangements (i.e., the preferred resonance structures **16** and **17**) were found (Scheme 9). The HOMO-SOMO inversion mentioned in Figure 2 depends on the nature of the substituents.

Electron withdrawing groups (EWGs) at the δ -carbon favor the carbanion **16**, whereas alkyl groups (Alk) or a combination of one EWG and one alkyl group (e.g., **17a–e**) favor **17**. When considering dimethyl malonate and varying the *N*-substituent, an electron distribution similar to **16** was observed in most cases.

As shown in Table 1, the reaction free energies are only negative for two entries (7 and 8), which have two different



Scheme 9. Effects of substituents on the preferred electron arrangements in acyclic radical anions.

Table 1. Enthalpy and free energy for cyclization reactions of 16 . All energies reported in kcal mol^{-1} .					
#	Substituent	16	18	ΔH	ΔG
1	$R^1 = R^2 = \text{COMe}$	a	a	14.0	16.2
2	$R^1 = R^2 = \text{CO}_2\text{Me}$	10	11	8.6	8.6
3	$R^1 = R^2 = \text{CO}_2\text{Et}$	b	b	9.0	9.4
4	$R^1 = R^2 = \text{CO}_2^i\text{Pr}$	c	c	9.4	10.0
5	$R^1 = R^2 = \text{CO}_2^t\text{Bu}$	d	d	11.8	12.8
6	$R^1 = R^2 = \text{CO}_2\text{CF}_3$	e	e	10.4	13.0
7	$R^1 = \text{CO}_2\text{Me}$ $R^2 = \text{CO}_2\text{CF}_3$	f	f	−1.2	−0.1
8	$R^1 = \text{CO}_2\text{Me}$ $R^2 = \text{H}$	g	g	−2.6	−1.7

substituents at the reacting carbon atom. Changes in reaction enthalpy in Table 1 are largely dependent upon the relative abilities of R¹ and R² to stabilize the carbanion in the reactant and the extra electron in the product. For instance, in the model substrate **10**, the two ester groups in the (CO₂Me)₂CH₂ (pK_a = 15.9,^[41] DMSO) moiety can provide synergistic stabilization to its conjugate base. However, communication between the two carbonyls is broken in both the transition state and the product where the extra electron resides in only one of the carbonyl groups (Figure 3). In this scenario, presence of two EWGs leads to unproductive and excessive reactant stabilization.

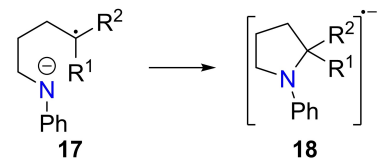
If one of the ester groups is removed (as seen in entry 8), the reactant carbanion is less stabilized (CH₃CO₂Et: pK_a = 29.5,^[42] DMSO). By removing unproductive reactant stabilization, more favorable reactions can be designed ($\Delta H = 8.6 \text{ kcal mol}^{-1}$ vs. $\Delta H = -2.6 \text{ kcal mol}^{-1}$).

A similar effect can be achieved when one of the EWGs is a much stronger acceptor than the other (i.e., R¹ = CO₂Me, R² = CO₂CF₃, entry 7). In this situation, the trifluoro-substituted group plays a dominant role in stabilizing both the reactant and the product. Hence, the loss of contribution from the less important CO₂Me substituent in stabilizing the negative charge along the reaction path is less penalizing.

The consequences of over-stabilizing the conjugate base can additionally be seen in entries 1 and 2. When the two ester groups R¹ and R² are swapped for the stronger ketone C(O)Me acceptors (pK_a = 13.3^[43]), the reaction becomes even less favorable ($\Delta H = 14.0 \text{ kcal mol}^{-1}$).

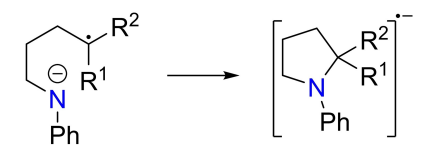
Entries 2–5 show that the size and steric effects of alkyl groups in the ester moiety impact reactivity. When R¹ = R² = CO₂R, an increase in the size of R from Me → Et → ⁱPr → ^tBu results in a decrease in reaction favorability ($\Delta H = 8.6, 9.0, 9.4,$ and $11.8 \text{ kcal mol}^{-1}$, respectively).

Table 2. Enthalpy and free energy for cyclization reactions of **18**. R¹ and R² participate in stabilization of both reactant and product. All energies reported in kcal mol⁻¹.



#	Substituent	17	18	ΔH	ΔG
1	R ¹ = COMe R ² = H	a	h	0.1	0.1
2	R ¹ = COMe R ² = Me	b	i	2.9	5.6
3	R ¹ = CO ₂ Me R ² = Me	c	j	7.5	10.2
4	R ¹ = CO ₂ Me R ² = Et	d	k	8.4	8.9
5	R ¹ = CO ₂ Me R ² = ^t Bu	e	l	12.2	13.1
6	R ¹ = CO ₂ Me R ² = Ph	f	m	5.5	7.4

Table 3. Enthalpy and free energy for cyclization reactions of **18**. R¹ and R² only participate in stabilization of reactant. All energies reported in kcal mol⁻¹.



#	Substituent	17	18	ΔH	ΔG
1	R ¹ = R ² = H	g	n	0.2	1.1
2	R ¹ = R ² = Me	h	o	0.9	4.3
3	R ¹ = Et R ² = Me	i	p	3.3	7.2
4	R ¹ = ^t Bu R ² = Me	j	q	6.7	9.7
5	R ¹ = R ² = ^t Bu	k	r	16.2	19.0

For the reactants in both Tables 2 and 3, the preferred localization of the radical moves from nitrogen to carbon and, hence, the negative charge is localized on nitrogen (Scheme 9).

In Table 2, entries 1 (R¹ = COMe, R² = H) and 2 (R¹ = COMe, R² = Me), additionally show that moving from a less stable (secondary) to a more stable (tertiary) radical adds a thermodynamic penalty ($\Delta H = 0.1 \text{ kcal mol}^{-1}$ vs. $\Delta H = 2.9 \text{ kcal mol}^{-1}$). Entries 3–5 (R¹ = CO₂Me, R² = Me, Et, ^tBu) continue to show that increasing the size and steric of alkyl groups near the reactive centers further increases the reaction enthalpy. Finally, the cyclic radical anion **18 m** is different from previous examples in that the upconverted electron is additionally delocalized over the aromatic system of the pendant phenyl group (R²) in the cyclic product providing some additional stabilization in comparison to the unsubstituted analog **11**.

As can be seen in Table 3, R¹ and R² provide direct stabilization for the radical in the reactant but not the product. Entry 1 is the most favored cyclization ($\Delta H = 0.2 \text{ kcal mol}^{-1}$) as it starts from the least stabilized primary radical. The cyclization of a more stable tertiary carbon radical (R¹ = R² = Me) is less favorable ($\Delta H = 0.9 \text{ kcal mol}^{-1}$). Similar to what was observed in Table 1, an increase in alkyl group size continuously renders the cyclization more endergonic as it provides additional radical stabilization in the reactant (through hyperconjugation) and destabilizes the product by steric effects.

Figure 4 shows the change in enthalpy for the cyclization of different *N*-aryl reactants **19** to the cyclic radical anions **20** versus the Hammett parameter (σ) of the related *N*-substituent. We explored a total of 23 aromatic rings substituents Hammett sigma constants^[44] and found them to show a good correlation with the change in reaction enthalpy ($R^2 = 0.89$). We found that reactions with EWGs on the *N*-aryl group mold the reaction more energetically favored. Thus, a more electron deficient nitrogen center favors the cyclization. Electron donating groups (EDGs) have a less deactivating effect in the meta- than in the para-position. In contrast, EWGs favor the para- over the meta-position. In other words, EDG and EWGs favor positions which

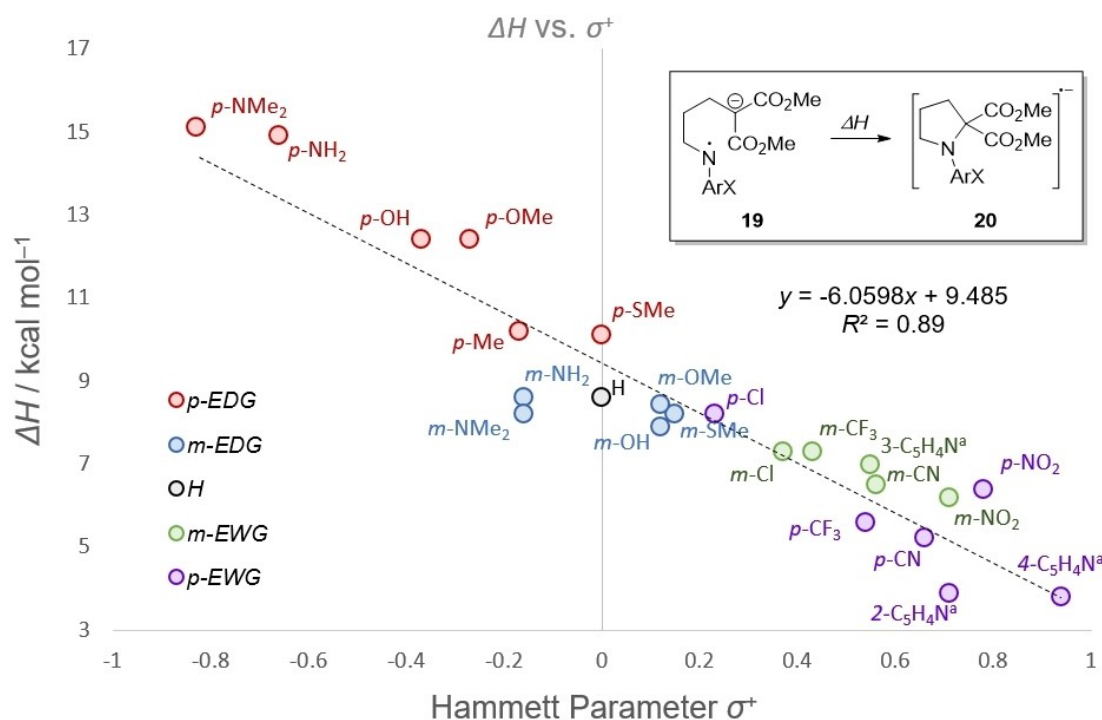


Figure 4. Plot of Hammett parameters vs. ΔH in kcal mol^{-1} for the conversion of compound **19** to **20**. A total of 23 different aromatic units were analyzed. The para and meta electron withdrawing groups are presented in purple and green, respectively. The para and meta electron donating groups are presented in blue and red, respectively. [a] The Ar group is pyridine (2-, 3-, or 4-Py) and not a phenyl ring with a pyridyl substituent.

(through an inductive effect) lower electron density at the nitrogen-radical.

Exploratory transition states

In addition to the thermochemical data depicted thus far, further radical anionic transition states have been identified and are shown in Table 4. The same trends observed for the thermochemical data apply to the activation free energies. EWGs lower the activation barrier while steric hindrance on the reacting centers leads to an increase. The *N*-phenyl moiety leans to the same side as the ester group, where the spin density is located. This observation can be seen for transition states **TS2**–**5**. In the absence of any EWGs, transition state energies are further destabilized **TS6**–**8**. The broad range of the incipient C–N distances ($r_{\text{C-N}}$) indicate the C–N bond to be sensitive to the strength and the number of acceptors.

Outlook

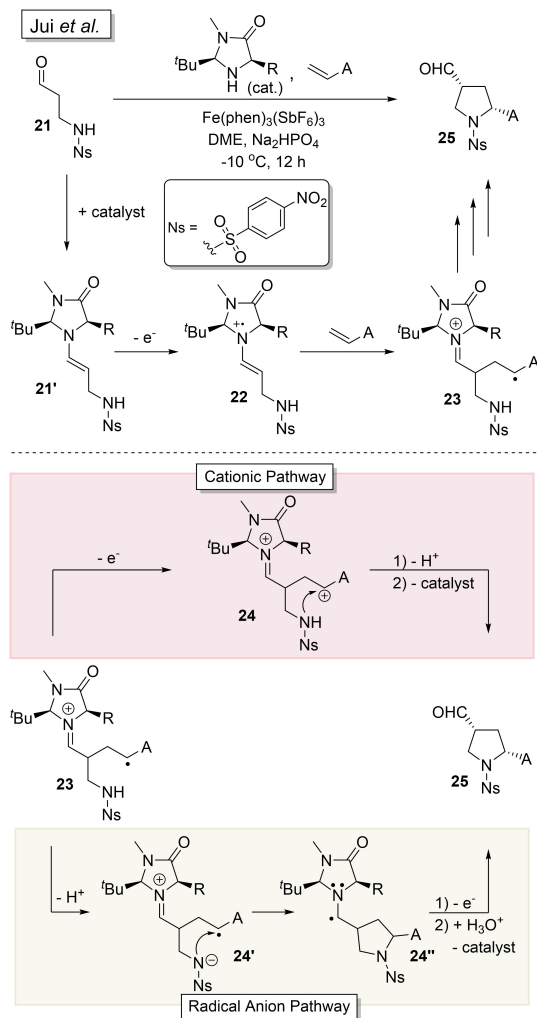
The potential utility of hidden 3e-pathways extends to cascades where such pathways are coupled with other reactions. For example, Jui et al. described an elegant method for combining aldehydes and alkenes to generate pyrrolidines (Scheme 10).^[5b] Here, the C–N bond formation is the final step in the stereospecific formation of pyrrolidine-3-carbaldehydes starting from a transient enamine and an olefin.

Table 4. Calculated radical anionic transition states. All energies reported in kcal mol^{-1} .

TS	Substituent	ΔG^\ddagger	ΔG	$r_{\text{C-N}}/\text{\AA}$
TS2	$R^1 = R^2 = \text{CO}_2\text{Me}$	12.9	8.6	1.7
TS3	$R^1 = \text{CO}_2\text{Me}$ $R^2 = \text{CO}_2\text{CF}_3$	11.2	-0.1	1.8
TS4	$R^1 = R^2 = \text{CO}_2\text{Et}$	14.8	9.4	1.7
TS5	$R^1 = R^2 = \text{CO}_2^t\text{Bu}$	18.0	12.8	1.8
TS6	$R^1 = R^2 = \text{H}$	20.8	1.1	1.9
TS7	$R^1 = R^2 = \text{Me}$	20.4	4.3	2.0
TS8	$R^1 = \text{Me}$ $R^2 = \text{Et}$	22.5	7.2	2.0

The authors suggest a plausible mechanism where, after the in situ formation of the enamine **21'**, oxidation forms a nitrogen radical-cation **22**. This intermediate adds to an alkene to produce a distonic radical-cation **23** with a separated carbon radical and an iminium cation center. The newly formed carbon radical is proposed to be oxidized to carbenium ion **24**. The carbenium ion is captured by the pendant nitrogen nucleophile to form the C–N bond of the pyrrolidine core.

The above-mentioned mechanism is reasonable but, perhaps, not general, as one can envision an alternative scenario for the key cyclization step. When comparing a cationic molecule (such as **23**) with its neutral counterpart, one can

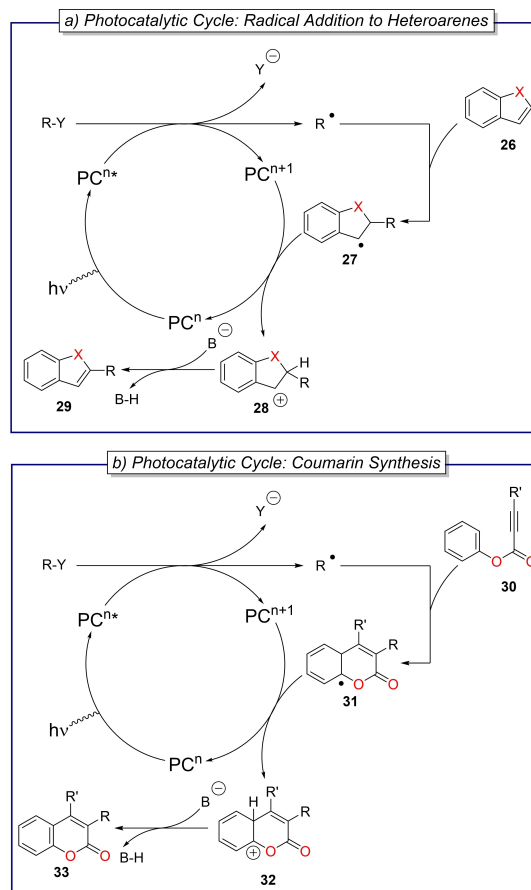


Scheme 10. Top: Proposed mechanistic pathway for the conversion of aldehydes and alkenes into pyrrolidines. Bottom: Possible deprotonation-mediated cyclization pathways.

anticipate that the cation would be harder to oxidize.^[45] Note that the proposed oxidation of distonic radical-cation **23** would form a dication (the carbenium ion **24**).

On the other hand, the difficult oxidation of the radical-cation **23** can be bypassed if one could trap the carbon radical via formation of a 2c,3e bond with the nitrogen lone pair in the *N*-deprotonated intermediate **24'** (neutral but shown as a zwitter-ionic resonance structure in Scheme 10). The resulting upconverted neutral radical product **24''** would be easier to oxidize than the cation **23**. In fact, it is possible that the N–H bond of the sulfonamide (pro)nucleophile would be already deprotonated under the reaction conditions well before to the cyclization step.^[12c,46]

Furthermore, these mechanistic questions are not limited to C–N bond formation but can also involve the processes that create other bond types via radical/polar cross-over scenarios. The examples of radical addition to heteroarenes^[47] and the synthesis of coumarin^[48] in a photocyclization show how it can apply to the formation of C–C bonds as well (Scheme 11).



Scheme 11. a) A commonly proposed photocatalytic cycle for radical addition into heteroarenes. b) A commonly proposed photocatalytic cycle for the transformation of phenyl propiolates into coumarin derivatives.

Commonly, photocatalytic cycles involving radical addition to heteroarenes are initiated by promoting the photocatalyst (PC) to an excited state. Subsequently, the PC injects an electron into an R–Y bond resulting in the formation of a Y[−] anion and an R-radical. The R-radical adds to the π -system of the heteroarene **26** resulting in the formation of resonance-stabilized radical **27**. The next commonly proposed step is the oxidation of the newly formed radical and the reduction of the PC. The result is the formation of carbenium ion **28** and the regeneration of the PC, thus completing the catalytic cycle. After formation of a cation, the loss of a proton restores aromaticity and yields the final product **29**.

A similar sequence can be suggested for the synthesis of coumarins from phenyl propiolates **30**. Under photocatalytic conditions, the excited PC initiates the reaction via reduction of an R–Y bond. The newly formed R-radical then attacks the triple bond to initiate a 6-*exo*-dig cyclization that forms carbon radical **31**. Subsequent oxidation of the carbon radical forms carbenium ion **32** which can be deprotonated to restore aromaticity in **33**.

Considering the photocatalytic cycle mentioned above, the transformation from **27** to **28** (and **31** to **32**) is not unreasonable and is likely to occur under suitable reaction conditions.

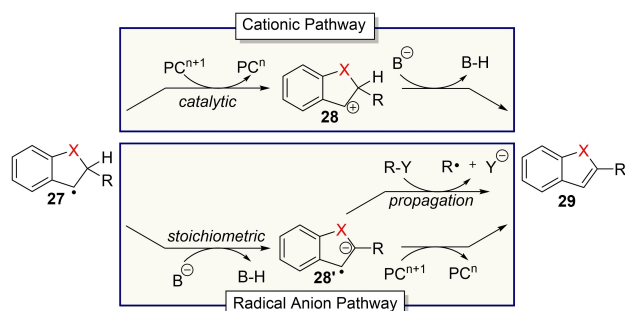
However, an alternative reaction pathway is worth considering, especially in the absence of strong oxidants (Scheme 12).

One should consider that the oxidation of the carbon radical to the carbenium ion requires the open-shell species to come in proximity to the oxidized PC. However, the PC is only present in *catalytic* amounts and this bimolecular electron transfer may not necessarily be a fast process.

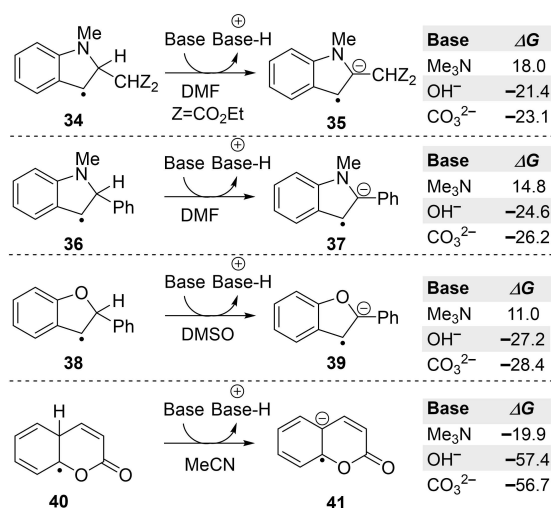
Alternatively, photocatalytic reactions often use base in *stoichiometric* (or even excess) quantities. Because the concentration of base is much greater than that of the PC, it is conceivable that, instead of the commonly proposed e^-/H^+ loss, the order of operations may be to remove H^+/e^- .

Consider the deprotonation of **27** and the subsequent formation of radical anion **28'**. Not only will **28'** be easier to oxidize by the PC than **27** but situationally, **28'** may be capable of injecting an electron into R–Y (instead of the PC), thus propagating the reaction.

We conclude by evaluating the possibility of electron upconversion via radical addition to heteroarenes (e.g., benzofuran and indoles) and in coumarin synthesis (Scheme 13). In



Scheme 12. Schematic representation for the final steps in a radical addition to heteroarenes. Top: Cationic pathway, loss of e^- then H^+ . Bottom: Radical anion pathway, loss of H^+ then e^- .



Scheme 13. Reaction energies for radical anion formation. Four separate reactants **34**, **36**, **38** and **40** were looked at, along with three bases (Me_3N , OH^- and CO_3^{2-}). Three common solvents for photocyclization were explored: DMF, DMSO, MeCN.

order to simulate the common reaction conditions, we explored a selection of frequently used solvents (DMF, DMSO, and MeCN) and three bases, i.e., trimethylamine (TMA), hydroxide, and carbonate that can assist in the formation of a radical anion via deprotonation of a radical.

In the presence of a strong oxidant and/or a weak base, carbenium ion formation is certainly feasible. The role of base is important. For example, with TMA as a base, the formation of radical anions **35**, **37**, and **39** was found to be thermodynamically unfavorable (Scheme 13).

However, in the presence of a suitable base, deprotonation and formation of radical anion prior to oxidation becomes feasible. This preference is illustrated by the deprotonation of radicals **34**, **36**, and **38** (Scheme 13) which becomes exergonic in the reactions with hydroxide and carbonate. Furthermore, as the acidity of the C–H bond increases, the formation of radical anions becomes more favorable. This is clearly observed in the deprotonation of radical **40**, in which all three bases were found to be suitable for radical anion formation **41**.

Conclusion

Alternative routes to C–H amination that involve intramolecular (2c,3e)C–N bond formation have been investigated with DFT calculations. Dialkyl 2-(3-(arylamino)propyl)malonates similar to the one used in the electrochemical cyclization of Xu et al. are predicted to be capable of a thermodynamically feasible cyclization from a radical anionic species.^[25] Additionally, we found that a more electron deficient nitrogen center should favor the cyclization. In contrast, an over-stabilization of the reactive center on the δ -carbon and sterically demanding groups are predicted to have a negative impact on the energetics of the cyclization.

In particular, an interesting stereoelectronic feature for the reaction of *N*-radicals and malonate anions is that only one of the ester groups is involved in the extra electron delocalization in the cyclized radical anion product. In contrast, the acyclic reactants have two esters that can stabilize the anionic center. Such unproductive excessive reactant stabilization can be removed when enolates derived from monoesters are used as the starting materials, leading to much more favorable thermodynamics for the respective three-electron cyclization.

Seven transition states were located for the model substrates with activation barriers ranging from 11 kcal mol⁻¹ to 22 kcal mol⁻¹. While the upper extreme of this range is likely to be less relevant for practical work due to potential side reactions, the high exergonicity of the overall process when combined with a mild terminal oxidant such as dioxygen should provide attractive opportunities for the design of new reactions based on electron upconversion.

Theoretical examinations of additional C–N and C–C bond forming reactions indicate that the concept of electron upconversion is likely to be relevant to a wider range of reactions (e.g., organo- and photocatalytic reactions). This alternative mechanism might be favored in situations in which a weaker oxidants and stronger bases are utilized.

Data Availability Statement

A detailed version of Scheme 5 can be found in the Supporting Information. Furthermore, the results of a benchmark for the free energy of the radical anion cyclization against different levels of theory are included in the Supporting Information. The DFT optimized geometries for all compounds and for unrestricted compounds the spin-squared expectation value $\langle S^2 \rangle$ related to the calculation can be found in the Supporting Information.

Author contributions

T. O. and I. V. A. conceived and supervised the project. [†]P. E. and Q. E. contributed equally to this work. All authors contributed to manuscript edits.

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Conflict of Interest

The authors declare no conflict of interest.

Keywords: C–N bond formation · cyclization · Marcus inverted region · radical anion · two-center three-electron bond

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