

REVIEW

[View Article Online](#)
[View Journal](#) | [View Issue](#)Cite this: *Chem. Sci.*, 2025, 16, 18519Received 26th July 2025
Accepted 19th September 2025

DOI: 10.1039/d5sc05603a

rsc.li/chemical-science

Electrochemical deamination functionalization via C–N bond cleavage and radical formation

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The C–N bond cleavage of amines has gained attention in the scientific community due to its numerous synthetic applications. In traditional methods, the requirements for toxic oxidants and costly catalysts affect their cost-effectiveness and sustainability. However, recent advances in synthetic organic electrochemistry allow for the *in situ* activation of C–N bonds, affording different functionalizations under mild reaction conditions with a shorter reaction time. In light of the ever-increasing importance of electrosynthesis spanning most disciplines of the chemical sciences, we review recent developments in electrochemically promoted radical deamination functionalization over the past decade (from 2015 to 2025). Special emphasis is put on various electrochemical transformation paths and proposed mechanisms.

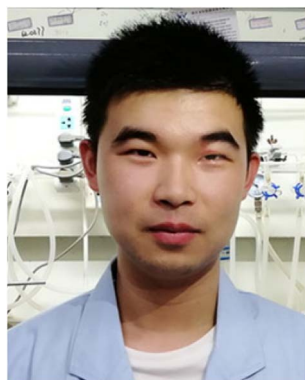
1. Introduction

Amines constitute a foundational class of organic compounds with significant structural diversity and broad applications across pharmaceuticals, agrochemicals, fine chemicals, dyes, and functional materials. The C–N bond, as one of the most prevalent chemical linkages, is ubiquitous in organic molecules and biological macromolecules. Its formation and transformation represent pivotal research domains in organic synthesis, organometallic chemistry, and biochemistry.¹ The

precise addition and removal of chemical functionalities in sophisticated molecular settings is a highly sought-after, yet challenging goal for organic chemists, essential for flexible molecular decoration. However, the high bond dissociation energy (102.6 ± 1.0 kcal mol^{−1}) of C–N bonds and the fact that NH₂ is virtually non-viable as a leaving group pose a persistent challenge for their selective cleavage.² Thus, converting the NH₂ group into a versatile and modular leaving group is highly desirable. Current strategies primarily rely on functional group activation, wherein amines are converted into activated intermediates such as diazonium salts,³ hydrazines,⁴ Katritzky salts,⁵ or ammonium salts.⁶ These modifications facilitate subsequent C–N bond cleavage and enable the construction of novel bonds. However, the majority of traditional C–N bond cleavage methods require a chemical reductant or transition metal catalyst, photocatalyst, or organocatalyst. Therefore, the search for new

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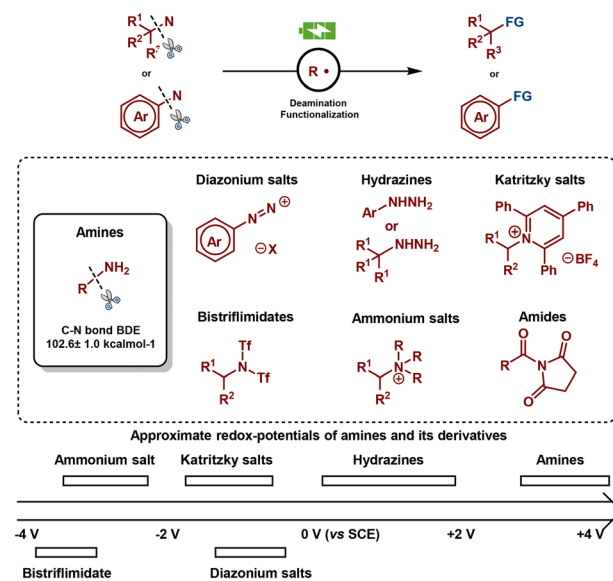
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“green” processes to achieve the cleavage of C–N bonds has become a hot topic in the development of new methods.

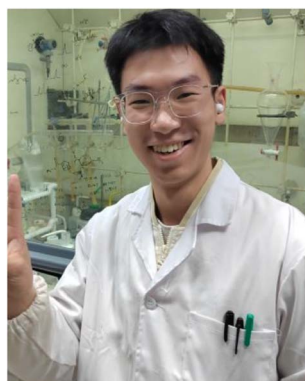
Organic electrochemistry⁷ leverages electrical energy as a renewable and clean synthetic driving force, as with wind and solar energy. By utilizing electrons and electron holes as traceless redox equivalents, it eliminates the need for stoichiometric chemical oxidants or reductants, positioning itself as an emerging environmentally sustainable paradigm in synthetic chemistry. This approach significantly enhances atom economy while diminishing dependence on fossil-derived energy resources. Through precision modulation of electrical input *via* optimized electrochemical conditions (*e.g.*, current, voltage, current density, electrode, electrolyte, and reaction temperature), electrochemical systems enable controlled reaction pathway steering. Due to the multiple redox conditions be allowed to exist simultaneously in the electrochemical reaction process, this facilitates stabilization of transient intermediates and enhanced selectivity, while occasionally unlocking unconventional mechanistic pathways.⁸ Many researchers are therefore working eagerly to discover new reaction patterns and make previously known reactions electrochemically accessible.

In fact, with significant advancements in the fields of C–N bond activation and electrochemical synthesis over recent decades, a large number of high-quality studies have been reported. However, only a few examples are compatible with



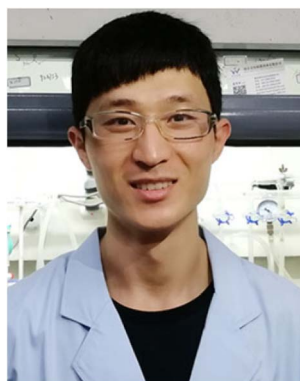
Scheme 1 Electrochemical deamination functionalization via C–N bond cleavage.

electrochemical conditions and achieve stable conversion. Additionally, radical type reagents are very easily activated through the electrode single-electron transfer redox process.



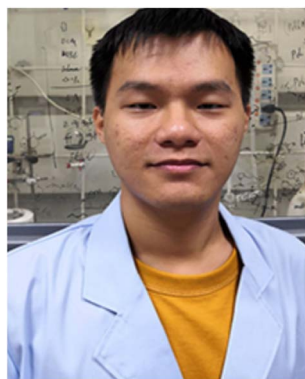
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For this reason, we focus on electrochemical synthesis strategies and review the electrochemical radical deamination functionalization methods established in the past decade (from 2015 to 2025) using a wide range of nitrogen-containing compounds, including diazonium salts, hydrazines, Katritzky salts, bistriflimidates and other nitrogen sources (Scheme 1).

2. Electrosynthetic deamination *via* aryl diazonium salts

Compared to the C-halide (Br/I) bond, the C–N bond exhibits significantly shorter bond lengths, lower polarizability, and greater chemical inertness.⁹ Consequently, chemical transformations reliant on C–N bond cleavage remain a substantial challenge in synthetic chemistry. In the initial report by Grieffs in 1858,¹⁰ a series of prominent named reactions associated with aryl diazonium salts, which are conveniently synthesized from the corresponding anilines, have been discovered and developed over the past centuries. Especially, the seminal discovery by Traugott Sandmeyer in 1884 (ref. 11) established a transformative paradigm for converting aryl diazonium salts into aryl halides, enabling regioselective substitution at the diazonium site. Moreover, those century-old transformations, including the Pschorr reaction (1896),¹² Gomberg–Bachmann reaction (1924),¹³ Balz–Schiemann reaction (1927),¹⁴ and Meerwein arylation (1939),¹⁵ are still widely used in modern organic synthesis in academic and industrial settings to this day due to their exceptional regioselectivity and broad functional group tolerance.

There are four main reaction types of aryl diazonium salts, including aryl radical transformations *via* single-electron transfer (SET) processes, aryl cation transformations *via* nucleophilic aromatic substitution processes (S_NAr), transition-metal-catalyzed processes, and transformations with retention of the dinitrogen group.¹⁶ Given the mechanistic constraints of electrochemistry, aryl diazonium cations readily undergo direct SET reduction at the cathode surface, forming diazo radicals. These intermediates spontaneously release nitrogen gas, producing aryl radicals primed for subsequent reaction transformation (Scheme 2).

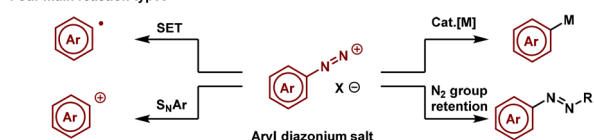
2.1 Electrochemical-mediated C(sp²)–C(sp²) bond formation *via* aryl diazonium salts

The earliest report on electrochemical synthesis involving aryl diazonium salts was published by Elofson and co-workers in

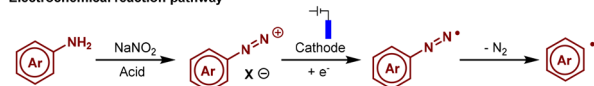
1969.¹⁷ Their study demonstrated that aryl–aryl coupling reactions could be achieved under electrochemical conditions using diazonium tetrafluoroborate as the aryl radical source, employing tetrabutylammonium perchlorate as the electrolyte and acetonitrile as the solvent with a standard calomel electrode in a divided cell. This approach successfully facilitated coupling across seven aromatic substrates, including benzene, toluene, anisole, benzonitrile, nitrobenzene, bromobenzene and naphthalene. However, this methodology exhibited limitations in regioselectivity and yields (below 33% yields) (Scheme 3A). Since then, the field of electrochemical aryl–aryl coupling *via* diazonium salts experienced a prolonged quiescent period until 2018, when Amaya's group developed a controlled electrochemical Gomberg–Bachmann reaction strategy for synthesizing heteroaromatic hydrocarbons by employing an undivided electrolytic cell with a carbon cathode and stainless steel anode.¹⁸ This methodology utilized DMSO-*d*₆ as the solvent, sodium tetrafluoroborate as the supporting electrolyte, and methanesulfonic acid as an additive. Notably, this method achieved heteroaromatic coupling with enhanced functional group tolerance toward electron-deficient substituents (*e.g.*, nitro, cyano, and halogens), yielding significantly higher efficiencies compared to electron-rich aromatic substrates (Scheme 3B).

In 2019, a new approach for the electrochemical Minisci-type arylation of various electron-deficient arenes with aryl diazonium salts was demonstrated by Lei and co-workers, providing 35 examples in yields up to 92%.¹⁹ Utilizing tetrabutylammonium tetrafluoroborate as the supporting electrolyte and a mixture of MeCN/TFA as the co-solvent, the arylation product of quinoxaline was obtained under 10 mA constant current for 4.5 hours. To prove the possible radical reaction pathway conjecture, the controlled experiments were conducted. EPR experiments were conducted in a divided cell and it was concluded that quinoxaline was reduced in the cathodic chamber, forming a quinoxaline radical that generated an EPR peak at *g* = 2.0040, 2 AN = 6 G, and 6 AH = 6 G. Moreover, a cyclic voltammetry study suggested that quinoxaline in the

Four main reaction types

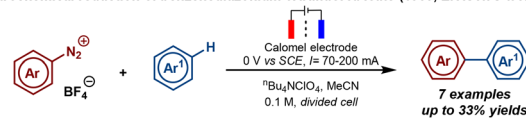


Electrochemical reaction pathway

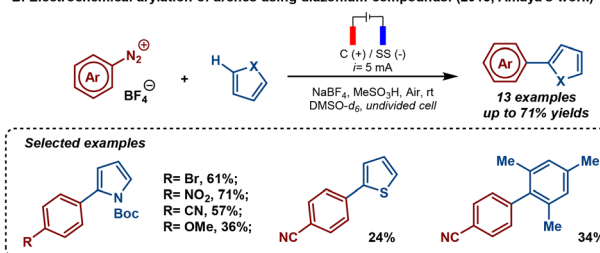


Scheme 2 The reaction pathway of aryl diazonium salts.

A. Electrochemical reduction of benzenediazonium tetrafluoroborate (1969, Elofson's work)

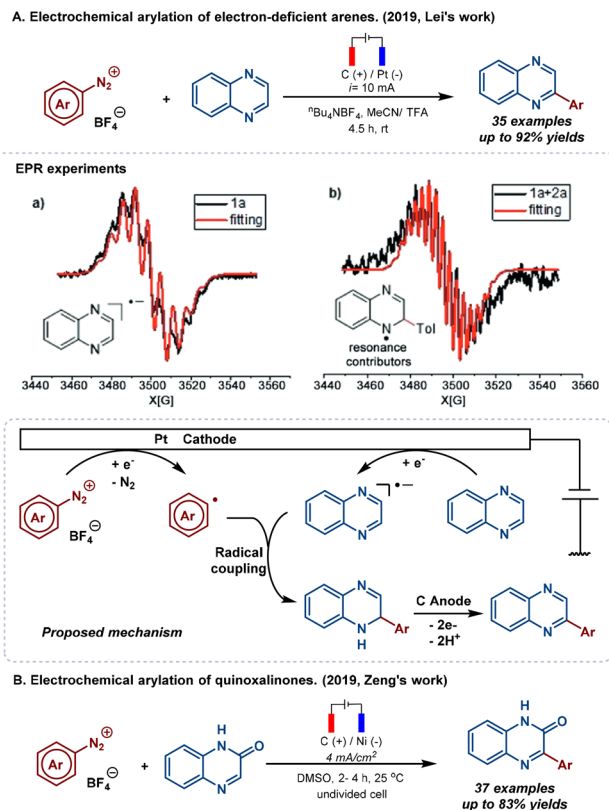


B. Electrochemical arylation of arenes using diazonium compounds. (2018, Amaya's work)



Scheme 3 Electrochemical-mediated aryl C–C bond formation with aryl diazonium salts. (A) Using aromatic substrates; (B) using heteroaromatic substrates.



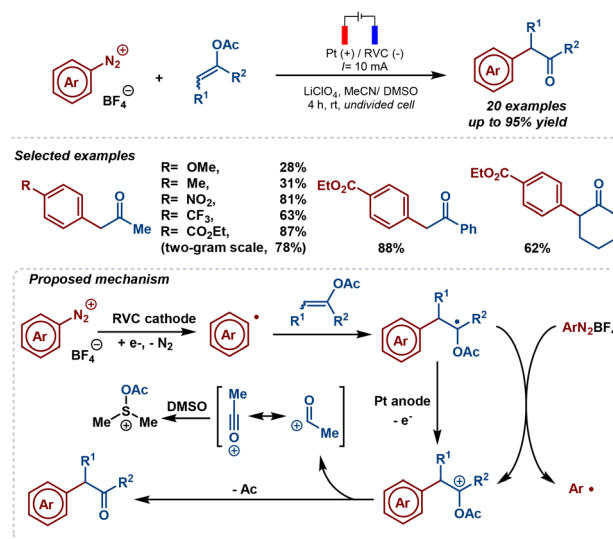


Scheme 4 Electrochemical-mediated Minisci-type arylation with aryl diazonium salts. (A) Using quinoxalines; (B) using quinoxalinones.

presence of TFA was initially protonated and then reduced at -0.34 V. Based on the above results, the author proposed a possible mechanism shown in Scheme 4A, in which quinoxaline and diazonium salts respectively undergo SET reduction at the C cathode to form the corresponding quinoxaline radical anion and aryl radical. Subsequently, a radical coupling process is carried out to obtain a coupling intermediate. Then, deprotonation and re-aromatization processes are successively performed under two anodic oxidation actions, ultimately resulting in the target product. With similar mechanisms to those proposed by Zeng²⁰ and co-workers in the same year, but the difference was that their electrochemical Minisci-type arylation of quinoxalinones with aryl diazonium salts in the absence of an external supporting electrolyte (Scheme 4B).

2.2 Electrochemical-mediated C(sp²)-C(sp³) bond formation via aryl diazonium salts

In 2022, Wang and co-workers reported a green, economical electrosynthesis process for ketone α -arylation, achieved by reacting aryl diazonium salts with enol acetates at room temperature without bases or metal catalysts.²¹ The optimized reaction conditions included lithium perchlorate as the supporting electrolyte, a mixture of CH₃CN/DMSO (v/v = 5:1) as the co-solvent and a constant current of 10 mA in an undivided cell equipped with a Pt plate anode and an RVC cathode. Enhanced yields were observed for *para*-substituted aryl



Scheme 5 Electrochemical-mediated Csp²-Csp² bond coupling of enol acetates.

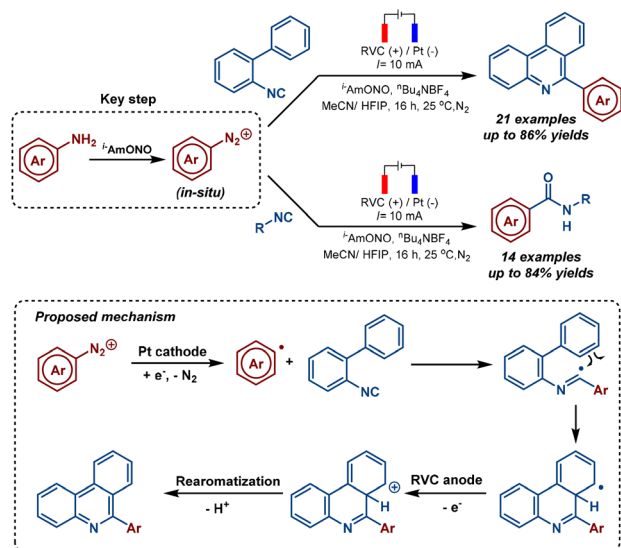
diazonium salts bearing electron-withdrawing groups (such as ester, cyano, nitro, and trifluoromethyl) compared to those with electron-donating groups (such as methoxy and methyl). Significantly, this methodology proved applicable to an *in situ* one-pot diazotization/electrochemical approach, and it achieved a 76% yield in a gram-scale synthesis of the target compound from enol acetate and aryl diazonium salt by using cheap graphite plate electrodes instead of expensive RVC electrodes. On the basis of control experiments, a reaction mechanism is proposed, which involves the initial reduction of aryl diazonium salts at the RVC cathode's surface to generate corresponding phenyl diazo radicals. Then, the aryl radical adds to enol acetate to generate a carbon radical intermediate. Subsequently, this intermediate is oxidized at the anode to afford the corresponding cation intermediate. Finally, the required α -arylation product was obtained through the departure of an acyl cation (Scheme 5).

In 2021, Sharma *et al.* utilized the electrochemical method to synthesize phenanthridines by the coupling of amines and 2-isocyanobiphenyls in a simple undivided cell.²² From the mechanistic perspective, the key step to the success of the reaction lies in the *in situ* formation of the aryl diazonium salt of aniline and amyl nitrite in the reaction system. Subsequently, the aryl diazonium salts are reduced at the Pt cathode to release nitrogen gas and aryl radicals. Notably, the aryl radical undergoes preferential capture by aryl isonitrile, followed by anodic oxidation and re-aromatization, affording phenanthridines. Conversely, alkyl isonitrile capture diverts the pathway toward amide formation (Scheme 6).

2.3 Electrochemical-mediated C-X bond formation via aryl diazonium salts

The classical Sandmeyer reaction represents a fundamentally important method to convert an aryl amine to an aryl halide *via* the intermediacy of a diazonium salt, including a copper metal



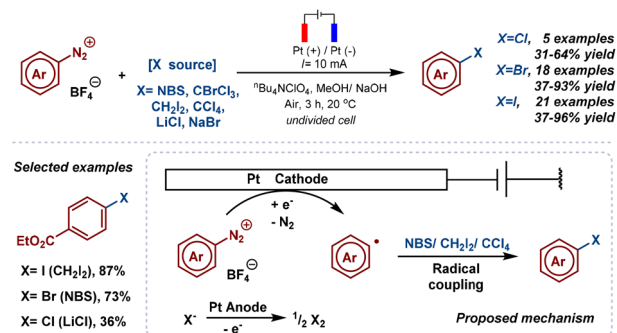


Scheme 6 Electrosynthesis of phenanthridines via 2-isocyanobiphenyls and aromatic amines.

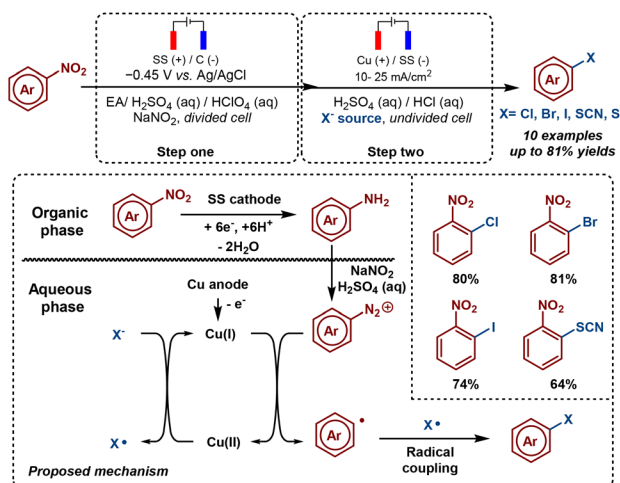
mediated process (Gattermann reaction),²³ diazotization in organic phases (Doyle diazotization),²⁴ a Cu(I)-catalyzed process²⁵ and acetate-facilitated metal-free halogenation.²⁶ Critically, the incompatibility of transition metal reagents or halogenated reagents with electrochemical conditions has greatly limited the development of electrochemical Sandmeyer reactions.

In 2018, Mo and co-workers reported a general electrochemical strategy for the Sandmeyer reaction, using a simple and inexpensive halogen source, such as NBS, CBrCl₃, CH₂I₂, CCl₄, LiCl and NaBr, for the halogenation of aryl diazonium salts.²⁷ Due to the advantage of electrochemical strategies, aryl diazonium salts can easily generate the key aryl radicals from cathode electrons and capture halogen radicals directly from halogen sources. However, aryl iodides and aryl bromides are more readily available than aryl chlorides, which may be affected by the differences in the activity of halogenation reagents and operating conditions. To prove the scalability of this electrochemical strategy, a gram-scale reaction was successfully performed and its potential for future industrial applications was demonstrated. Importantly, mechanistic studies, including *in situ* EPR, support a single-electron reduction pathway for the electrochemical halogenation of diazonium salts. This method complements Sandmeyer reactions, potentially paving the way for other metal-free transformations (*e.g.* noble metal-free trifluoromethylation) in the near future (Scheme 7).

An uncommon one-pot two-phase electrochemical reduction of aryl diazonium salts derived from nitrobenzenes to aryl halides was reported by Nematollahi in 2022 (Scheme 8).²⁸ To ensure success, the author adopted a step-by-step reaction strategy. First, the author respectively used a carbon electrode and a stainless-steel electrode as the cathode and anode, which were respectively located in the divided cells of an ethyl acetate/water (0.25 M HClO₄, 1.5 equiv. NaNO₂, and 1.2 M H₂SO₄) co-



Scheme 7 Electrochemical-mediated Sandmeyer reaction with aryl diazonium salts.

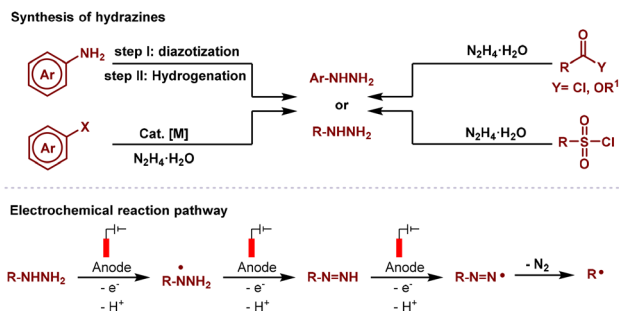


Scheme 8 Electrochemical-mediated heterogeneous Sandmeyer reaction of nitroaromatics.

solvent system, with a cell voltage of -0.45 V. This process ensures the *in situ* conversion of the nitro group to the diazo group and transfer from the organic phase to the aqueous phase. Next, an aqueous solution containing HCl, HBr, KI or NaSCN was added to the above solution and electrolysis was performed under constant current conditions at a copper anode/stainless steel cathode. Key to this strategy is the use of an available copper rod as an anode, which serves as an electrode and a catalyst by electrogenerating copper(I) ion species, which then oxidize aryl diazonium salts to generate the corresponding aryl radicals. Notably, this unique work synthesizes haloaryl compounds in a two-phase system at room temperature using either galvanic or electrolytic cells, achieving higher yields than previous approaches.

3. Electrosynthetic deamination via hydrazines

Hydrazines are characterized by their unique C–N bond architecture and have emerged as promising reagents in contemporary coupling chemistry. These reactions exploit C–N bond

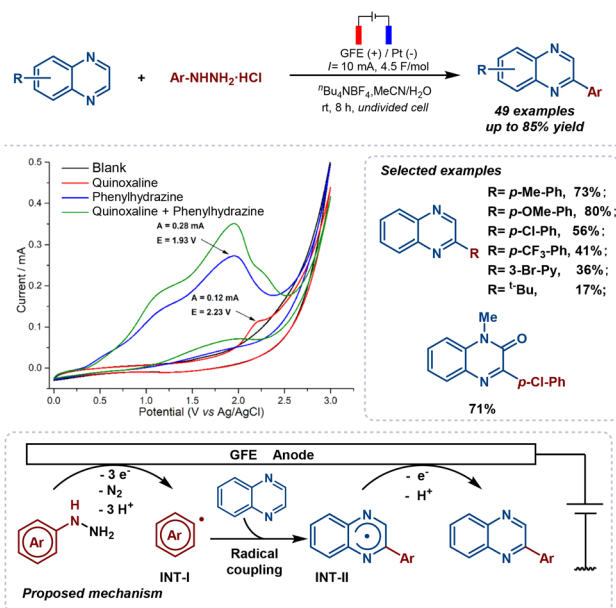


Scheme 9 The synthesis and electro-transformations of hydrazines.

cleavage to achieve flexible and synthetically diverse reaction systems. Relative to traditional electrophilic partners such as halogenated hydrocarbons, hydrazines present superior operational advantages, such as low cost, easy accessibility (derived from aniline, aryl halide, ester or acyl halide), and minimized environmental impact. In particular, the corresponding carbon radicals and nitrogen are released by electrochemical oxidation activation (Scheme 9). There is a growing demand for the development of strategies to achieve the transformation and utilization of hydrazines.

3.1 Electrochemical applications of aryl hydrazines

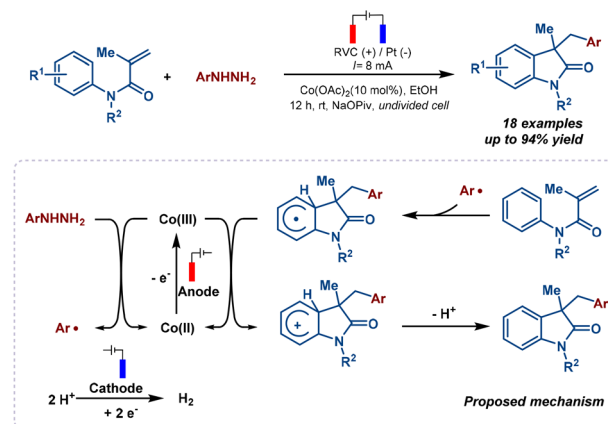
Radical oxidative coupling reactions are interesting alternatives to well-established methods for the formation of C–C and C–X bonds, contributing numerous significant innovations to synthetic chemistry. In 2022, Li and Zhou *et al.* revealed a practical and scalable protocol for electrochemical arylation of quinoxalin(on)es with arylhydrazine hydrochlorides in an undivided cell, providing 45 corresponding products in moderate to good yields (Scheme 10).²⁹ Electron-rich alkenes

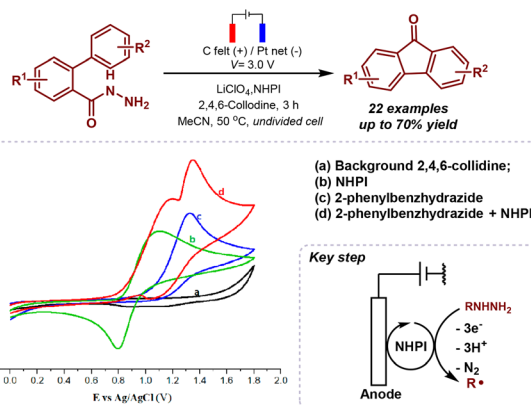


Scheme 10 Electrochemical oxidative C–H arylation of quinoxalines and quinoxalinones.

afforded high yields, while moderate yields were observed for electron-deficient alkenes. The reaction is significantly accelerated using microwave irradiation. Although this method exhibits high efficiency, easy scalability, and broad functional group tolerance, alkyl hydrazines are not suitable for this condition due to the tendency of alkyl radicals to oxidize to cations. Cyclic voltammetry experiments found that quinoxaline presented an oxidation peak at $E = +2.23$ V vs. Ag/AgCl, which is much higher than that of aryl hydrazines ($E = +1.93$ V vs. Ag/AgCl), indicating that aryl hydrazines are preferentially oxidized by the electrode. The authors proposed a reaction mechanism initiating with the oxidation of arylhydrazine at the graphite anode, which forms the aryl radical intermediate **INT-I** via deprotonation, three-electron loss, and N₂ release. This radical subsequently adds to quinoxaline to yield intermediate **INT-II**. Following further single-electron oxidation and deprotonation, the arylation product is afforded. Concurrently, hydrogen ions are reduced at the Pt cathode to form H₂. It should be noted that since the radical–radical coupling between azo and aryl radicals is inevitable, the use of an excess of arylhydrazine is proposed to suppress this side reaction and improve the arylation efficiency.

In 2018, Yu and co-workers developed a series of substituted oxindoles facily synthesized *via* electrochemical cobalt-catalyzed C–H or N–H oxidation between *N*-arylacrylamides and arylhydrazines or potassium alkyltrifluoroborates under mild conditions (Scheme 11).³⁰ The optimized reaction conditions included Co(OAc)₂ (10 mol%) as the catalyst, NaOPiv as the additive, EtOH as the effective solvent system and 8 mA constant current between a RVC anode and a Pt cathode in an undivided cell over 12 h at room temperature. To gain a deeper insight into the mechanistic pathways of these oxidative annulations, control experiments (such as radical inhibition experiments and gas-chromatographic headspace analysis) were conducted, demonstrating the radical pathway and the existence of molecular nitrogen and hydrogen as byproducts. Mechanistically, anodic oxidation converts Co(II) species to Co(III) species, which act as an oxidant to indirectly generate aryl radicals from aryl hydrazines. The generated aryl radical adds to

Scheme 11 Electrochemical co-catalyzed coupling of *N*-arylacrylamides with arylhydrazines.

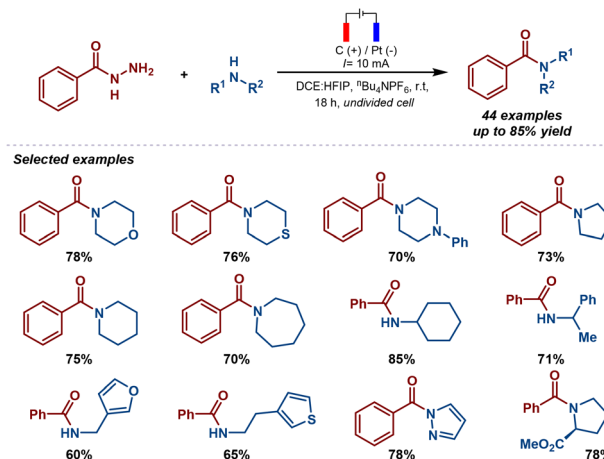


Scheme 13 Electrochemical synthesis of fluorenone derivatives from arylhydrazides.

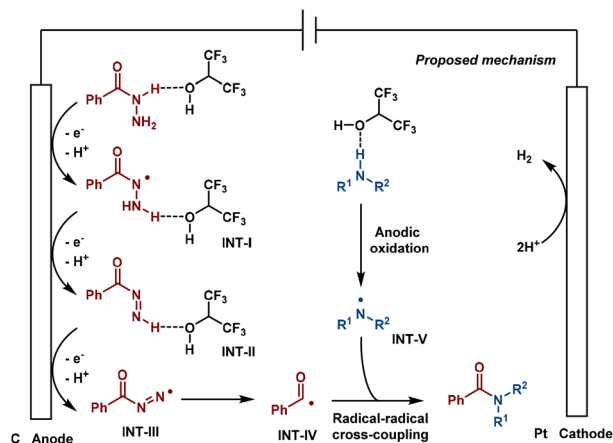
In addition to radical addition with an alkene moiety, an electrooxidative carbonylative Sonogashira-type cross-coupling reaction of arylhydrazines and alkynes has also been realized, achieving the synthesis of functionalized ynones under a relatively low CO atmosphere with moderate to excellent yields (Scheme 12).³¹ The proposed reaction pathway includes the precursor of the active catalyst Pd(II) formed by electro-oxidation of the Pd(0) species, which then reacts with arylhydrazines to yield the **INT-I** species, along with the release of N₂ and the insertion of CO into the C–Pd bond of the **INT-I** species, generating the **INT-II** species. Then, the **INT-II** intermediate reacted with terminal alkynes to yield the **INT-III** intermediate. Ultimately, reductive elimination occurred to afford the corresponding ynones and Pd(0) species. Notably, another possible initial source of Pd(0) species might be generated from the Pd(II) catalyst by the reduction of NH₄I in the presence of CO. This transformation would contribute significantly to the development of electrochemical carbonylative Sonogashira-type reactions.

attractive for the synthesis of fluorenones. To prove the possible radical reaction pathway conjecture, controlled experiments were conducted. A cyclic voltammetry study suggested that NHPI was initially oxidized at +1.05 V *vs.* Ag/AgCl (curve b) and arylhydrazines showed an oxidation potential of 1.38 V *vs.* Ag/AgCl (curve c). These results reveal that the oxidation of NHPI is preferable to anodic oxidation of arylhydrazines. However, the CV of the 1 : 1 mixture of NHPI and aroylhydrazides showed a catalytic current, while the reductive current of NHPI disappeared (curve d), which results from the hydrogen atom transfer (HAT) process between phthalimide *N*-oxyl (PINO) and arylhydrazines to regenerate NHPI. Based on the above results, a plausible mechanism was proposed, in which the reaction success hinged critically on the preferential electrochemical oxidation of acyl hydrazines to generate acyl radicals.

One year later, An electrochemical amidation of benzoyl hydrazine/carbazate and 1°/2° amine as coupling partners *via* concomitant cleavage and formation of C(sp²)-N bonds has been achieved by Patel and co-workers (Scheme 14).³³ This reaction generated both acyl and *N*-centered radicals from



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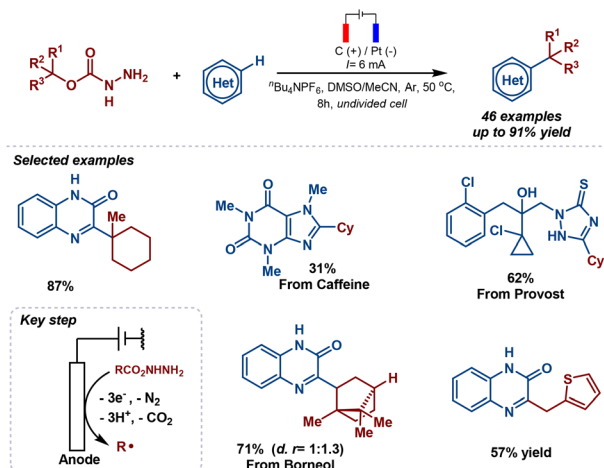
Scheme 15 Proposed mechanism.

benzoyl hydrazines and amines *via* the simultaneous cleavage and formation of C(sp²)-N bonds. A broad substrate scope was observed for both the benzoyl hydrazine/carbazate and amine (aromatic or aliphatic) components in this catalytic system. Significantly, the process produced only environmentally benign nitrogen and hydrogen gas as by-products. To further demonstrate the synthetic utility of this methodology, a gram-scale (10 mmol) reaction of benzoyl hydrazine with morpholine proceeded for 32 hours to yield morpholino(phenyl) methanone in 70% isolated yield. Remarkably, the practical applicability of this electrochemical radical coupling strategy was successfully extended to the synthesis of bezafibrate, a commercially available fibrate drug widely used for the treatment of hyperlipidemia.

The authors propose a plausible reaction mechanism for crucial bifunctional hexafluoroisopropanol (HFIP). First, the reaction begins with HFIP-solvent hydrogen-bonding activation of benzoyl hydrazine, enabling anodic electrochemical oxidation to form the *N*-centered diazanyl radical **INT-I**. Subsequently, intermediate **INT-I** then undergoes a two-step anodic oxidation sequence, progressing through intermediate **INT-II** to yield the diazene radical intermediate **INT-III**. Cleavage of the C-N bond within **INT-III** releases molecular nitrogen (N₂) and produces the benzoyl radical **INT-IV**. Simultaneously, an H-bonding interaction and anodic oxidation generate the *N*-centered radical intermediate **INT-V**. Finally, radical-radical cross-coupling between **INT-IV** and **INT-V** delivers the desired products (Scheme 15).

3.3 Electrochemical conversion of hydrazine carbamates

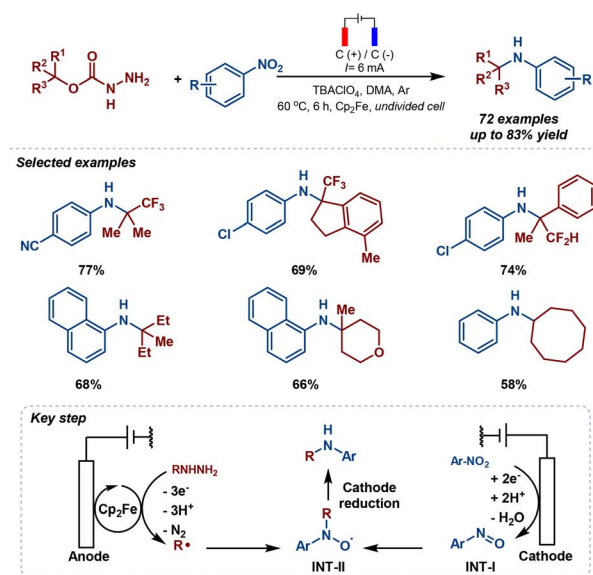
Compared with aryl hydrazines, there are only a few reports and applications of alkyl hydrazines under electrochemical conditions. In 2020, Wang and co-workers established that electrochemical oxidation efficiently affords key alkyl radical intermediates concurrent with N₂ and CO₂ elimination (Scheme 16).³⁴ This green electrochemical approach leverages sequential anodic oxidative fragmentation to access primary/secondary/tertiary alkyl radicals, enabling direct functionalization of diverse nitrogen-containing heteroarenes and some



Scheme 16 Electrochemical Minisci reaction.

natural products (*e.g.* benzoquinoxalinones, pyrazinones, quinazolinones, isoquinolines, phthalazines, quinazolines, phenanthridines, caffeine, provost, and borneol) in moderate to good yields with high stereoselectivity. Key to this reaction's success is the low oxidation potential of alkyl carbazates ($E_{\text{ox}} = +1.4$ V vs. SCE), facilitating their anodic oxidation to release electrons, protons, N₂ and CO₂. The generated radicals then undergo Minisci-type coupling with nitrogen-containing heterocycles.

More recently, the same group achieved a general electrochemical deoxygenative C-N bond coupling of alkyl carbazates with nitroarenes (Scheme 17).³⁵ This strategy enabled the conversion of diverse primary, secondary, and tertiary alkyl carbazates into valuable alkylamines, including substrates bearing α -CF₃, α -CF₂H, and benzyl groups. The reaction



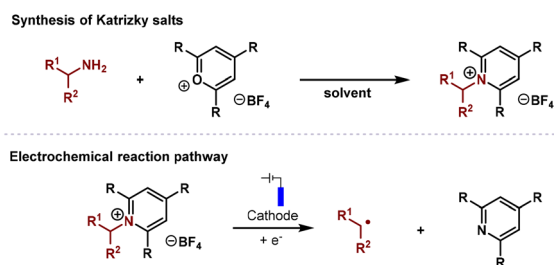
Scheme 17 Electrochemical C-N bond coupling of hydrazine carbamates with nitroaromatics.



proceeded with a broad substrate scope and mild conditions, demonstrating significant potential for sustainable synthesis. With the combination of flow chemistry strategy, the gram-scale electrochemical C–N bond coupling of alkyl carbazates with nitroarenes was realized under continuous-flow conditions, enabling overall reaction times of 20 h. To expand the application range of the substrate and enhance the conversion efficiency, the authors added ferrocene as an oxidation medium to the reaction system. The paired electrolysis mechanism of this reaction is shown in Scheme 17. Initially, some of the alkyl carbazates were multistep oxidized at the anode, and the rest were initially oxidized at the anode involving Cp_2Fe (0.42 V), regenerating Cp_2Fe^+ . The generated alkyl radicals are captured by an *in situ*-generated nitroso species **INT-I** (from cathodic nitrobenzene reduction), forming intermediate **INT-II**. Subsequent cathodic reduction of intermediate **INT-II** ultimately delivers the amine products. In fact, the mismatched reactivity of alkyl radicals and nitrogen sources has been addressed by paired electrolysis, providing a powerful and versatile tool for streamlined access to a wide array of amine compounds on a preparative scale, especially for advancements in the synthesis of α -fluorinated amines.

4. Electrosynthetic deamination via Katritzky salts

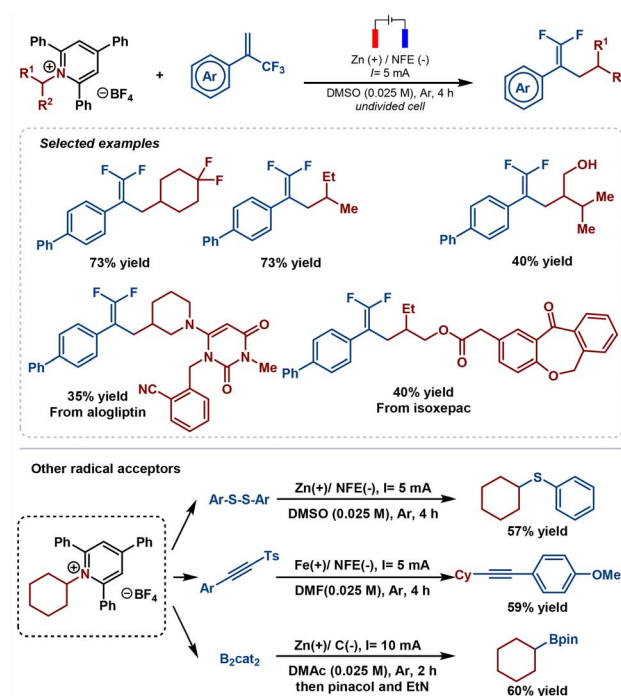
Due to the low redox potentials, Katritzky salts ($E_{\text{red}} = -0.92$ V vs. SCE) are more reactive compared to NHP-ester ($E_{\text{red}} = -1.28$ V vs. SCE) and halogens ($E_{\text{red}} = -1.7$ to -2.2 V vs. SCE), and they provide new suite precursors for selective radical alkylation. As early as the 1970s, Katritzky³⁶ and co-workers first found that 2,4,6-trimethylpyrylium tetrafluoroborate can convert alkyl amines into bench-stable, bulky pyridinium salts in a single step. Katritzky salts effectively reduce the energy of the $\text{C}(\text{sp}^3)\text{--N}$ bond, opening up the prospect of functionalization by the cleavage of the $\text{C}(\text{sp}^3)\text{--N}$ bond under mild reaction circumstances. Primary amine-derived Katritzky pyridinium salts have been employed as carbon radical surrogates for transition-metal catalysis and photoredox chemistry. Previous approaches³⁷ on reductive cross-coupling of Katritzky salts required expensive photocatalysts, electron donor–acceptor (EDA) complexes or heavy metal catalysts for the single electron transfer process. The analogous SET outcome could be achieved by cathodic reduction without chemical catalyst loading (Scheme 18).



Scheme 18 The reaction pathway of Katritzky salts.

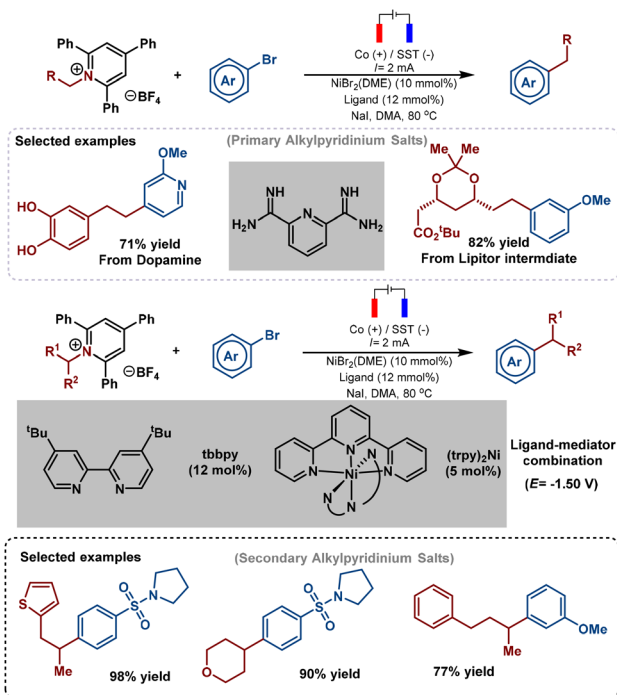
In 2021, Wang's team first demonstrated a metal-free electrochemical reductive deaminative $\text{C}(\text{sp}^3)\text{--C}(\text{sp}^3)$ bond cross-coupling of Katritzky salts with various radical acceptors.³⁸ In that work, many *gem*-difluoroalkenes were synthesized under electrochemical reductive conditions. The results showed that a diversity of secondary Katritzky salts were verified for this electrolytic protocol, providing cross-coupling products in moderate to excellent yields. It is worth mentioning that no Ni catalyst was involved in this reaction and C–N scission in this conversion could be mediated by the zinc anode serving as the sacrificial electrode, which significantly improved the atomic economy and reduced the consumption of fossil fuels. Moreover, $\text{C}(\text{sp}^3)\text{--C}(\text{sp}^3)$, $\text{C}(\text{sp}^3)\text{--C}(\text{sp}^2)$, $\text{C}(\text{sp}^3)\text{--C}(\text{sp})$, C–S, and C–B bonds were also successfully constructed. This deaminative functionalization, facilitated by rapid molecular diffusion across microfluidic channels, demonstrates practicality that outperforms conventional electrochemistry setups (Scheme 19).

Nickel-catalyzed coupling reactions are important synthetic tools for the construction of carbon–carbon and carbon–heteroatom bonds, owing to advantages, such as low cost, wide substrate compatibility, and mild reaction conditions. However, the favorable reduction potential of Katritzky salts ($E_{1/2} \approx -1.4$ V vs. Fc/Fc^+) poses a particular challenge for electroreductive Ni catalysis (-1.7 V vs. Fc/Fc^+) that Katritzky salts reduce faster than Ni intermediates or typical alkyl electrophiles (e.g., alkyl halides and redox-active esters), generating off-cycle byproducts *via* dihydropyridyl trapping or alkyl radical dimerization/reduction. Moreover, successful catalysis critically depends on balancing the rate of aryl halide oxidative addition with the alkylpyridinium salt activation step.



Scheme 19 Electrochemical deaminative functionalization of Katritzky salts with radical acceptors.

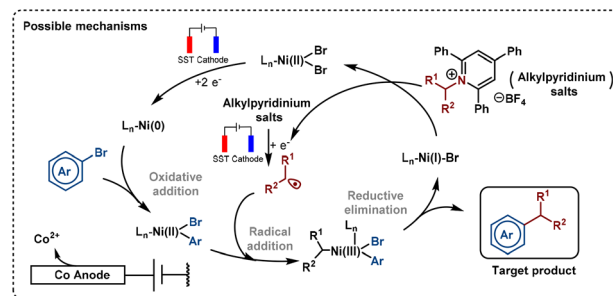




Scheme 20 Nickel-catalyzed electroreductive coupling of Katritzky salts with aryl halides.

In 2023, the first electroreductive coupling of alkylpyridinium salts with aryl bromides, leveraging state-of-the-art high-throughput experimentation (HTE) for electrochemical reaction development, was disclosed by Sevov, Watson, and Kalyani (Scheme 20).³⁹ First, the authors optimized the conditions with the help of high-throughput technology and determined that such telescoped methods effectively leverage the amine building blocks for library synthesis in the context of medicinal chemistry applications (over 50 drug examples). Second, they embarked on the elucidation of the scope and generality of electrochemical reductive cross-coupling by studying the reactions of alkylpyridinium salts with diverse aryl and heteroaryl bromides, furnishing the desired products in good to excellent isolated yields. Next, they re-optimized conditions for reactions of secondary alkylpyridinium salts in detail and ligand mediators were found to be critical to achieving high product yields. Finally, they conducted a multi-dimensional library synthesis of 48 distinct products *via* the reductive cross-coupling of 6 aryl bromides against 4 primary and 4 secondary alkylpyridinium salts using HTE-Chem. Excitingly, microscale HTE further demonstrated the synthetic value of this method by enabling the efficient construction of extensive libraries of cross-coupled products, a key advantage for material-sparing discovery in medicinal chemistry, and this strategy highlights the power of HTE technologies for enabling new pharmaceutically relevant electrochemical transformations that provide complementary efficiency or chemical space access to their nonelectrochemical counterparts.

As shown in Scheme 21, the proposed mechanism starts with the formation of a carbon radical by single-electron transfer



Scheme 21 Proposed mechanism.

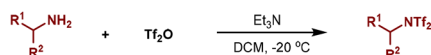
(SET) from the excited state of the SST cathode to alkylpyridinium salts, or alkylpyridinium salts oxidise the Ni(i) reactive species to obtain the corresponding carbon radical intermediates and Ni(II) intermediates, which are cathodically reduced to Ni(0) intermediates and then oxidatively adducted with bromobenzene to obtain the Br-Ni(II)-Ar intermediates, which trap free carbon radicals to form unstable Ni(III) intermediates and undergo a rapid reductive elimination, resulting in C(sp²)-C(sp³) coupling products and Ni(i) reactive species. Notably, this one-step paired electrolysis can avoid the isolation of frequently toxic alkyl halides and the use of stoichiometric Zn, Mg or Mn in reductive cross-couplings. Regarding substrate generality, this methodology demonstrates broad applicability across three key dimensions: (i) facilitating coupling reactions between diverse primary/secondary amines and aryl bromides, (ii) enabling efficient synthesis of structurally complex natural product derivatives, and (iii) permitting precise modifications of pharmacologically active scaffolds. The protocol exhibits moderate to excellent functional group compatibility while maintaining satisfactory stereochemical control.

5. Electrosynthetic deamination *via* bistriflimidates

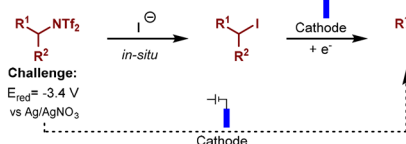
Although bistriflimidates could be readily synthesized on a gram scale using amines and trifluoromethanesulfonic anhydride as starting materials, with triethylamine as the base and dichloromethane as the solvent at -20 °C, its direct electrochemical transformation *via* cathodic reduction presents a significant challenge. This difficulty arises from its exceptionally high reduction potential ($E_{\text{red}} = -3.4$ V *vs.* Ag/AgNO₃), which prevents direct radical generation.⁴⁰ Consequently, the electrochemical pathway for bistriflimidates involves a two-step sequence: (i) priority conversion to the corresponding alkyl iodide ($E_{\text{red}} = -3.1$ V *vs.* Ag/AgNO₃) and (ii) subsequent cathodic reduction of alkyl iodide to generate the target radical species for further reaction (Scheme 22).

In 2024, Wang and Hong collaborated to develop new alkyl bistriflimides, which were unprecedentedly employed for C-N bond activation (Scheme 23).⁴¹ Alkyl amines are efficiently converted to alkyl bistriflimides *via* trifluoromethanesulfonic anhydride, serving as superior electrophiles over traditional Katritzky salts or redox-active imines. This replaces toxic

Synthesis of bistriflimides



Electrochemical reaction pathway

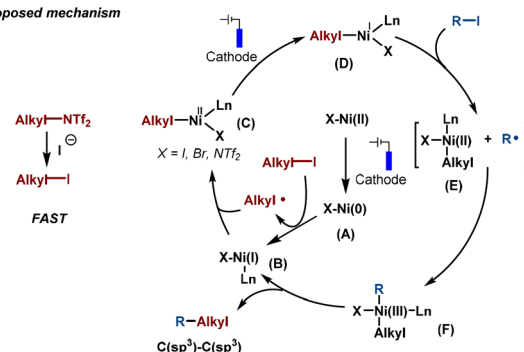


Scheme 22 The reaction pathway of bistriflimides.

stoichiometric metal reductants (e.g., Mn and Zn) with electricity, aligning with green chemistry principles. The methodology enables the formation of diverse $C(sp^3)-C(sp^3)$ and $C(sp^3)-C(sp^2)$ bonds with broad compatibility. It couples alkyl bistriflimides with alkyl halides, aryl/heteroaryl halides, alkenyl triflates, Katritzky salts and NHPI esters, achieving moderate to excellent yields. The broad substrate scope, excellent functional group tolerance, and mild reaction conditions collectively underscore the practicality and effectiveness of this methodology. Beyond expanding the toolbox for C–C bond construction, this work establishes a sustainable strategy for employing amine-derived precursors in organic synthesis.

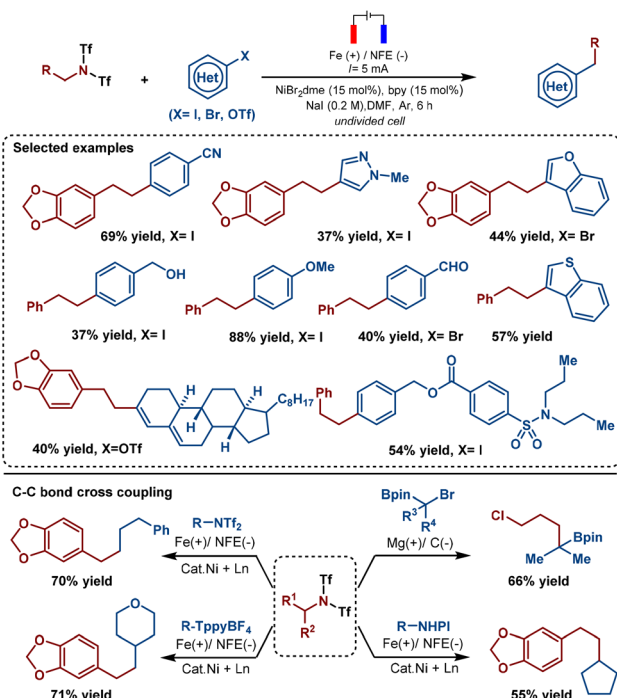
This approach avoids stoichiometric oxidants, expensive catalysts (e.g., Pd and Ru) and neutral by-product interference. Reactions proceed at room temperature under constant current (5 mA) using inexpensive electrodes (Mg anode/Ni foam cathode) and NaI as a critical electrolyte. Alkyl bistriflimides

Proposed mechanism

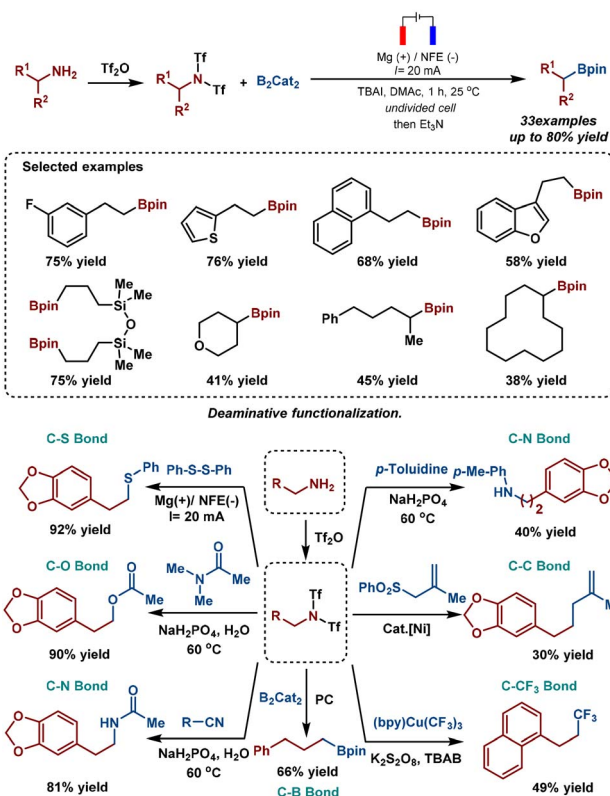


Scheme 24 Proposed mechanism.

react rapidly *via* an S_N2 mechanism to generate key alkyl iodide intermediates in the presence of iodide ions. The proposed catalyst pathway initiates with the electrochemical reduction of Ni(II) to generate a low-valent Ni(0) species **A**. This Ni(0) complex subsequently undergoes single-electron transfer (SET) with alkyl iodides, yielding alkyl radicals while oxidizing to Ni(I) species **B**. Species **B** participates in radical addition to form alkyl-Ni(II) intermediate **C**. A second electroreduction event then reduces **C** to Ni(I) species **D**. Crucially, **D** engages in oxidative addition with an additional halide substrate, forming Ni(III) species **F**. The cycle concludes with reductive elimination from



Scheme 23 Electrochemical C–C bond coupling of alkyl bistriflimides with aryl halides.

Scheme 25 Electrochemical C–B bond coupling of alkyl bistriflimides with B_2cat_2 .

F, releasing the C(sp³)-C(sp³) cross-coupled product and regenerating the Ni(I) catalyst (Scheme 24).

Later, the same groups demonstrated that the more efficient C-B bond cross-coupling reaction could be extended to alkyl bistrimides and B₂cat₂.⁴⁰ After detailed investigations, the author found that the desired alkyl boronate could be obtained in 78% yield under an undivided cell set-up with a magnesium anode and a nickel foam cathode at a working current of 20 mA. This protocol exhibited good efficiency toward the borylation of alkyl bistrimides, showcasing wide functional tolerance and generating borylation products in moderate to high yields. Notably, the reaction time only requires 1 hour. In addition to C-B bond formation through electrochemistry, subsequent interception by various nucleophiles provides products with remote C-C, C-S, C-O, and C-N bonds, and bistrimides have also been explored as radical initiators to allow the regioselective cleavage of C-N bonds and the preparation of trifluoromethylated derivatives in the presence of CuCF₃. The author proposed that this strategy employs an undivided cell without the use of transition metal or photo-catalysts and exhibits high conversion and stability in flow reactors (Scheme 25).

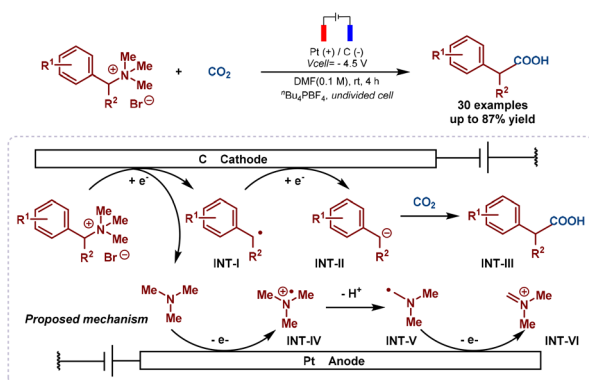
6. Electrosynthetic deamination via other nitrogen sources

Apart from the above-mentioned deoxygenation reagents, other nitrogen sources (such as quaternary ammonium salts and amides) have made significant contributions to the field of electrochemical deoxygenation functionalization.

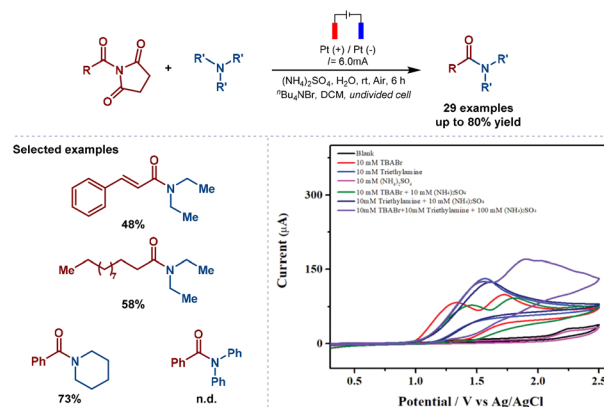
In 2019, Manthiram and co-workers provided a new design strategy for electrochemical carboxylation which utilizes the benzyltrimethylammonium bromide intermediate as a substrate for carboxylation (Scheme 26).⁴² The optimized conditions involved performing the reaction in an undivided cell with a Pt anode and a C cathode in the presence of benzylammonium salt (0.15 mmol), CO₂ bubbling, and ⁿBu₄-PBF₄ as electrolyte in DMF (0.1 M) at -4.5 V cell voltage and room temperature for 4 h. Compared to previous transition-metal-catalyzed carboxylations, this electrochemical method

avoids using stoichiometric metals as reducing agents. By employing trimethylamine (*in situ* generated from the substrate) as an anodically oxidizable sacrificial agent, this method avoids traditional sacrificial anodes while achieving high current efficiency, suppressed overoxidation, and broader substrate compatibility. The proposed mechanism involves concurrent electrode processes: cathodic reduction of an ammonium salt generates benzyl radical **INT-I**, which undergoes further reduction to anion **INT-II** followed by carboxylation to 2-phenylacetate **INT-III**. Simultaneously, trimethylamine **INT-IV** liberated through reductive C-N cleavage diffuses to the anode, oxidizing to radical cation **INT-V**, which further loses electrons to form iminium cation **INT-VI**. Taken together, this method proceeds through C-N bond cleavage and subsequent CO₂ insertion, eliminating the need for stoichiometric metal reductants, sacrificial anodes, or purification by column chromatography. Both primary and secondary benzylammonium substrates were carboxylated with high selectivity and afforded products in moderate to excellent yields. The reaction also demonstrated excellent functional group tolerance. Given its user-friendly nature, this system holds promise for application in the synthesis of diverse aliphatic carboxylic acids and dicarboxylic acids.

Wang and Luo have developed a novel and efficient electrochemical oxidative transamidation of tertiary amines with *N*-acyl imides, offering advantages over traditional metal catalysis or emerging visible light catalysis by eliminating the need for metal catalysts and oxidants.⁴³ After extensive optimization, ⁿBu₄NBr as the supporting electrolyte, (NH₄)₂SO₄ as an additive, and DCM/H₂O as the effective solvent system furnished the desired product by employing a constant current of 6 mA between Pt (anode) and Pt (cathode) over 6 hours at room temperature in an undivided cell. The reaction proceeded smoothly to generate the desired transamidation products under transition metal-free and photocatalyst-free conditions with good functional group compatibility (including alkyl, halogen, alkene and heteroaromatic-ring-based *N*-acyl-imides) in moderate to good yields. Unfortunately, triarylamines cannot serve as an ideal nitrogen source to complete the conversion under these conditions (Scheme 27).

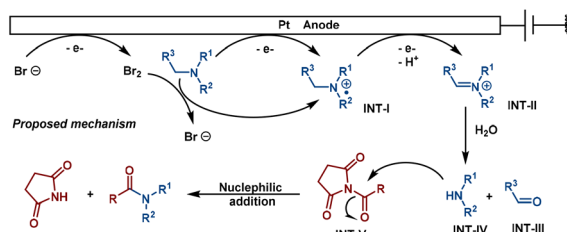


Scheme 26 Electrochemical carboxylation of benzylic C-N bonds with CO₂.



Scheme 27 Electrochemical oxidative transamidation of tertiary amines with *N*-acyl imides.

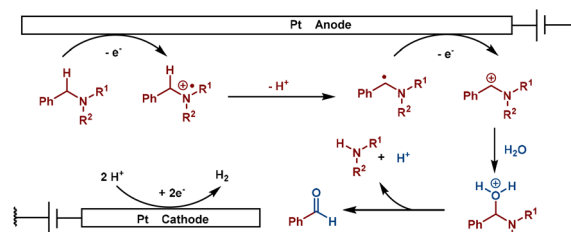




Scheme 28 Proposed mechanism.

Moreover, cyclic voltammetry (CV) analysis under standard conditions produced oxidation peaks at +1.29 V, +1.57 V, and +1.75 V *vs.* Ag/AgCl, indicating that bromine anions are first oxidized to molecular bromine and then involved in the oxidation of triethylamine. The mechanism commences with anodic bromide oxidation generating Br₂, which migrates to the organic phase oxidizing trimethylamine to intermediate **INT-I** while regenerating Br[−] to sustain a catalytic bromine cycle. Concurrently, trimethylamine undergoes direct anodic oxidation *via* single-electron transfer and α -hydrogen loss at the tertiary amine center to yield **INT-I**. This intermediate sequentially transforms into unstable iminium ion **INT-II**, which undergoes hydrolytic fragmentation to aldehyde **INT-III** and secondary amine **INT-IV**. Finally, nucleophilic attack by **INT-IV** on activated tertiary amide **INT-V**, followed by *N*-deprotection, furnishes the transamidation product. It presents a novel approach for the cleavage and conversion of C–N bonds, with the potential to pave the way for the design and improvement of synthetic pathways (Scheme 28).

Apart from the above reports, an uncommon efficient electrochemical method for the oxidative cleavage of C–N bonds under mild conditions using water as the oxygen source was furnished by Liu and Xia (Scheme 29).⁴⁴ The optimized reaction conditions include Et₄NBr as the electrolyte, a 10:1 (v/v) mixture of MeOH/H₂O as the effective solvent system, a constant current of 2.0 mA and a duration of 24 h in an undivided cell setup assembly with Pt as the anode and cathode. With the optimal conditions in hand, the substrate scope of this protocol was investigated, and it was found that benzylamines



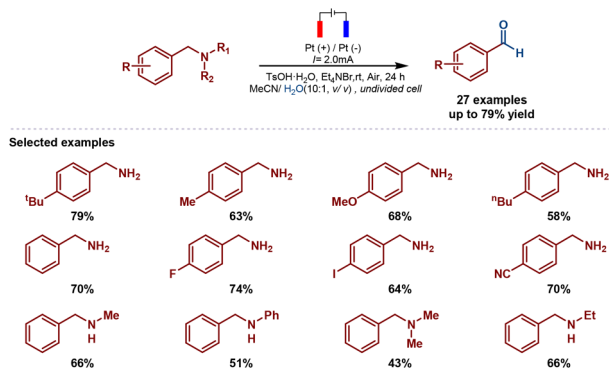
Scheme 30 Proposed mechanism.

bearing electron-donating groups and electron-withdrawing groups at the para position of the benzene ring are converted to the corresponding aldehyde compounds with moderate to good yields. Additionally, disubstituted benzylamines, *N*-substituted and *N,N*-disubstituted benzylamines also tolerate this transformation to afford the corresponding compounds. In order to further explore the reaction mechanism, butylhydroxytoluene (BHT) was added under standard conditions, and the yield of the product was significantly reduced, which implied the possible involvement of radical species in the reaction. In CV analysis, the addition of TsOH/H₂O and H₂O was able to cause a significant emergent oxidation peak at +3.2 V, proving that the reaction could proceed *via* the oxidation of benzylamine. The transformation mechanistically involves a single-electron transfer (SET) between benzylamine salts and an *in situ* generated nitrogen radical cation. This initiates concurrent deprotonation and radical 1,2-migration to afford an α -aminoalkyl radical intermediate. Subsequent anodic oxidation of this radical generates a benzyl cation, which undergoes nucleophilic trapping by H₂O. TsOH/H₂O-mediated dissociation of the resulting aminol liberates the target carbonyl product (Scheme 30).

7. Conclusions and perspectives

Radical reactions are the heart of modern synthetic chemistry. The generation of radicals *via* electrochemical strategies is efficient and mild, compared with the traditional methods with the use of stoichiometric chemical oxidants or reductants. However, many aspects of electrochemical deamination functionalization *via* C–N bond cleavage and radical formation are still not purely understood and are still far from being well developed. The discussion of the electrochemical deamination functionalization reaction, in particular, the electro-redox generation of radical intermediates from redox-active amines and the subsequent reactions, as well as the reaction mechanism, has been presented in this review.

Despite significant advancements and achievements being made in this area, there is still room for further exploration, as follows: (a) electrochemical construction of diverse C–Y (*e.g.*, C–Si, C–P, and C–N₃) bonds based on redox-active amines and established reaction types would undoubtedly expand the research horizon in this area. (b) Development and exploration of the deamination properties and electrochemical applications of new deamination reagents, such as imines, amides, azo



Scheme 29 Electrochemical selective oxidative cleavage of the benzyl C–N bond.



compounds, and azides. (c) There are no reports on chiral control through redox-active amines using electrochemistry. (d) Electrochemical three-component reactions using redox-active amines as a radical precursor are underexplored. (e) Reports involving direct electrochemical deamination of amines are scanty. Overall, we believe that electrochemical deamination functionalization will play a major role in complementing the growing repertoire of residue-specific modifications and bi-conjugation approaches, which are driving innovation in organic synthesis, medicinal chemistry, and chemical biology in the near future, and hope that this review will receive significant attention and contribute to further achievements in this area.

Author contributions

Investigation: J. D., Y. S. and W. C.; writing – review & editing: Z. Z.; supervision: W. Z. and Y. W.

Conflicts of interest

There are no conflicts to declare.

Data availability

No primary research results, software or code have been included and no new data were generated or analysed as part of this review.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (No. 22071101, 22271147, 22471123 and 22401144), the Foundation of the Hexi University (ZDSYS.2023-5), the Science Foundation of Tongling University (2023tlxyrc15 and 2024tlxykjZD008), and the Innovation Project of Anhui Province (S202410383041).

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