



Cite this: *Org. Biomol. Chem.*, 2022, **20**, 6707

Received 10th June 2022,
Accepted 1st August 2022

DOI: 10.1039/d2ob01080a

rsc.li/obc

Electrosynthetic C–F bond cleavage

Johannes L. Röckl,^a Emma L. Robertson^b and Helena Lundberg  *^a

Fluorinated organic compounds are common among pharmaceuticals, agrochemicals and materials. The significant strength of the C–F bond results in chemical inertness that, depending on the context, is beneficial, problematic or simply a formidable synthetic challenge. Electrosynthesis is a rapidly expanding methodology that can enable new reactivity and selectivity for cleavage and formation of chemical bonds. Here, a comprehensive overview of synthetically relevant electrochemically driven protocols for C–F bond activation and functionalization is presented, including photoelectrochemical strategies.

1. Introduction

Fluorinated organic compounds have widespread use as pharmaceutical motifs, agrochemicals and materials due to their unique physical, chemical, and biological properties. For example, incorporation of fluorine in molecules can result in increased lipophilicity, bioavailability and metabolic stability.¹ Fluorine has the highest electronegativity of all elements in the periodic table² and forms highly polarized bonds with carbon, similar to its heavier halogen congeners. However, the C–F bond is significantly stabilized by the electrostatic attraction between

the positively polarized carbon and the negatively polarized fluoride.³ This stabilization is reflected in the considerably higher dissociation energies of C–F bonds^{4,5} compared to other carbon-halide bonds (Fig. 1) and affects the kinetics for their cleavage, resulting in low reactivity of fluorocarbons.⁶ This relative chemical inertness is evident in everything from classic polar nucleophilic substitution reactions of alkyl halides³ to formation of carbon centered radicals *via* reductive dissociative electron transfer in *e.g.* cross-electrophile coupling (XEC) reactions⁷ and halogen atom transfer (XAT) processes⁸ as well as in transfer halogenation of alkenes.⁹ Depending on the context, the chemical inertness of the C–F bond can be viewed as beneficial (*e.g.* metabolic stability of pharmaceuticals), problematic (*e.g.* persistence of per- and polyfluoroalkyl substances – PFAS) or simply an intriguing synthetic challenge.

^aDepartment of Chemistry, KTH Royal Institute of Technology, SE-100 44 Stockholm, Sweden. E-mail: hellundb@kth.se

^bNovogensys Ltd, Dunnet, Kirrenleigh, Thurso KW14 8YD, UK



Johannes L. Röckl

Johannes L. Röckl is currently a Senior Scientist at RISE, Research Institutes of Sweden in Södertälje, Sweden. He received his Ph.D. under supervision of Prof. Dr Siegfried R. Waldvogel at Johannes Gutenberg University Mainz in Germany and Prof. Dr Bill Morandi at ETH Zurich in Switzerland, working mainly on novel electro-synthetic and catalytic transformations. In 2021, he joined the group of Dr Helena Lundberg

at KTH Royal Institute of Technology in Stockholm in Sweden, focusing on kinetic analysis and novel electroreductive reactions.



Emma L. Robertson

Emma L. Robertson studied medicinal and biological chemistry at Edinburgh University, Scotland and ETH Zurich, Switzerland. After internships in medicinal chemistry at Roche in Basel, Switzerland and agrochemistry at BASF SE in Ludwigshafen, Germany, Emma worked as an Editor for Wiley VCH, Weinheim in Germany. She pursues a career as a publications specialist and medical writer in the pharmaceutical industry.



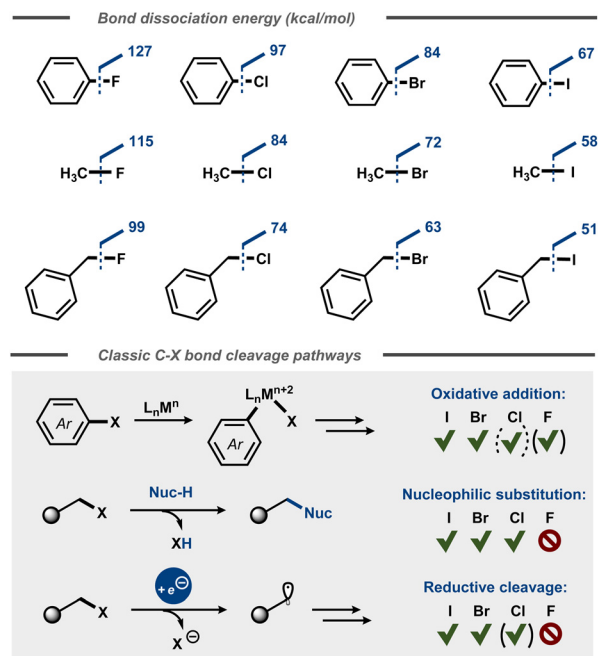


Fig. 1 Top: Bond dissociation energies for aromatic and aliphatic C–X bonds.⁴ Bottom: Oxidative addition, nucleophilic substitution, and reductive cleavage of C–X bonds.

While fluorides are typically introduced into a compound for the macroscopic properties they give rise to, selective defluorination can be of high synthetic benefit. For example, such a strategy can enable the use of trifluoromethyl groups as precursors for difluoromethyl (CF₂H) groups that have received significant attention for medicinal chemistry applications.¹⁰ Despite the low reactivity of C–F bonds, synthetic strategies are available for their cleavage. The most common approach relies on elimination through an E1cB mechanism, including

nucleophilic aromatic substitutions.^{3,11} In these processes, an inductively stabilized anion is formed adjacent to a fluoride and subsequently undergoes an irreversible elimination step to form a double bond with release of the halide. In addition, C–F bond activation *via* a variety of mechanisms has been reported using *e.g.* transition metal catalysis,¹² Lewis acids based on main group elements,^{6,13} early transition metals and lanthanides^{13,14} and low-valent metal reagents.^{10,15} Recent developments in radical synthesis have also delivered new methods for C–F bond cleavage, including photo- and electrochemical strategies.¹⁶ The latter is becoming an increasingly popular synthetic approach, due to the prospects of new reactivity and selectivity, as well as resource friendly synthesis that follows with electricity as terminal reagent.¹⁷ While excellent reviews summarizing the principles of and synthetic possibilities for electrochemistry have been published lately,¹⁸ a comprehensive overview of electrochemical protocols for C–F bond cleavage is missing in the literature to date. To address this gap, this review discusses synthetically relevant protocols for electrochemically promoted C–F bond cleavage from the 1950s to present, along with mechanistic details, with the overarching aim of promoting further developments in the field. Electrochemically driven degradation of PFAS was recently surveyed and is thus not included in the present review.^{19,20} In all figures throughout this review, the anode is found to the left and the cathode to the right when depicted above a reaction arrow.

2. Cleavage of C(sp²)–F bonds

Electroanalytical studies of C–F bond cleavage in fluorobenzenes were first reported more than 50 years ago,^{21,22} establishing that the standard reduction potential for fluorobenzene was close to –3 V vs. SCE.^{23,24} One of the first synthetic protocols for hydrodefluorination of fluoroarenes was reported by Kariv-Miller *et al.*²⁵ Using a Hg pool cathode in the presence of dimethylpyrrolidinium tetrafluoroborate ((DMP)BF₄), selective monodefluorination of 1,3-difluorobenzene to fluorobenzene was achieved in 85% yield using a divided cell setup (Fig. 2, top). A constant current of 1.25 mA cm^{–2} at 2 °C and 2 F mol^{–1} was found optimal for the monodefluorination, whereas higher current densities resulted in over-reduction to benzene. While the selectivity for monodefluorination was kept intact, the concentration of both water and (DMP)BF₄ was found to influence the yield due to competing hydrogen evolution. The optimized conditions were also amenable for reduction of fluorobenzene to benzene in 75% yield. The mechanism was probed using cyclic voltammetry, revealing an increase of cathodic current and decrease of anodic current for (DMP)BF₄ in the presence of substrate, indicative of catalysis. Supported by separate studies,²⁶ the authors proposed that a catalytically active “amalgam” of DMP⁺ and mercury forms upon single electron reduction at the cathode. This mediating surface was claimed to react with the fluorinated substrate to form the corresponding radical anion that, in turn, decomposes to an



Helena Lundberg

Helena Lundberg is employed as Assistant Professor in Organic Chemistry at KTH Royal Institute of Technology in Stockholm, Sweden. She received her Ph.D. degree under the guidance of Professor Hans Adolfsson at Stockholm University, after which she carried out postdoctoral research at the same institution with Professor Fahmi Himo. In 2017, she joined Professors Donna G. Blackmond and Phil Baran as a postdoctoral

fellow at Scripps Research in La Jolla, USA. Helena's research group at KTH focuses on activation and functionalization of strong polarized σ -bonds using catalysis and electrochemistry.



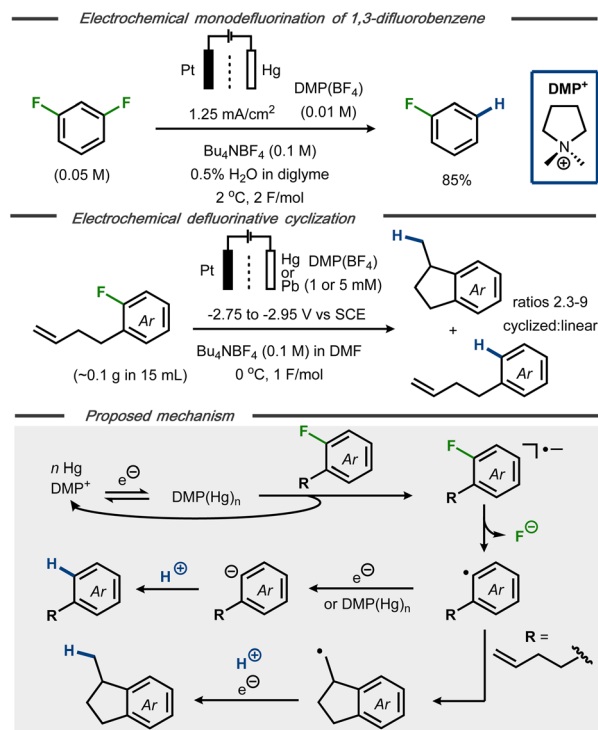


Fig. 2 Top: Selective monodefluorination of 1,3-difluorobenzene.²⁵ Middle: Defluorinative reductive cyclization.²⁷ Bottom: Mechanistic proposal for reductive hydrodefluorination.^{25–27}

aryl radical with release of fluoride (Fig. 2, bottom). A second single electron transfer to the aryl radical followed by protonation would thereafter form the hydrodefluorinated arene product. This mechanistic proposal was further supported by studies on defluorinative cyclization of *o*-(3-butenyl)fluorobenzene at Hg and Pb cathodes in the presence of catalytic amounts of DMP⁺.²⁷ From experiments with cyclic voltammetry and preparative electrolysis, it was found that the DMP⁺ ion enabled a shift in reduction potential from -2.9 V to -2.75 V vs. SCE for reduction of the substrate, thus supporting the idea of its mediating effect for the electron transfer. In addition, a selectivity enhancement for cyclization product 1-methylindane over hydrodefluorination product 3-butenylbenzene was observed in the presence of DMP⁺ (Fig. 2, middle). The cyclization pathway was proposed to proceed *via* the same aryl radical intermediate as the hydrodefluorination product (Fig. 2, bottom), followed by rapid intramolecular cyclization and subsequent single electron reduction/protonation to afford the defluorinated indane product. The solvent, residual moisture or the tetrabutylammonium cation of the supporting electrolyte was hypothesized to serve as proton source.

Hydrodefluorination of a limited number of 7-piperazinoquinolones under electroreductive conditions was reported by Albini and co-workers (Fig. 3).²⁸ Similar to Kariv-Miller's findings,²⁵ selective monodefluorination of position 8 in 6,8-difluorinated lomefloxacin was achieved after 2 F mol⁻¹ and defluorination of the monofluoride-containing enoxacin was

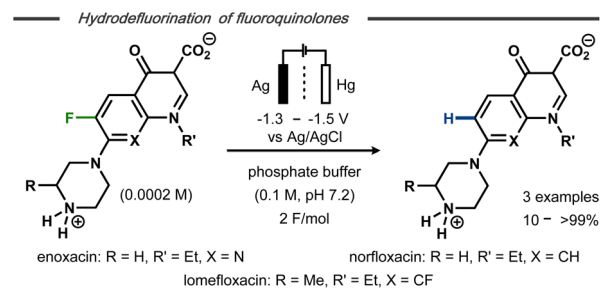


Fig. 3 Electrochemical hydrodefluorination of fluoroquinolones.²⁸

observed. In contrast, norfloxacin resulted in a mere 10% yield of defluorinated product along with side-products. The higher yields and selectivities for enoxacin and lomefloxacin compared to norfloxacin correlated with the reduction potentials of the compounds observed by differential pulse voltammetry. While distinct reduction waves were observed for the former two, the cathodic wave of the latter was found very close to that of the electrolyte solution. This difference was argued to result from the substitution pattern of the heterocyclic compounds, resulting in greater or lesser stabilization of the hypothesized intermediate aryl radical and, hence, a more or less selective reaction. Selective monodefluorination of methyl-2,6-difluorobenzoate was also reported by Périchon and co-workers in the presence of 10 mol% SmCl₃.²⁹ While no mechanistic details were conveyed, the catalyst afforded a selectivity switch from ester reduction to defluorination. Defluorination of the monofluoride product was not observed under the applied conditions.

Wu *et al.* described electrochemical hydrodefluorination of fluoroaromatic compounds in the presence of borohydride reagents.³⁰ Using platinum electrodes in an undivided cell, the electrolysis was carried out in diglyme or *N*-methylpyrrolidine (NMP) at constant current to provide 15 examples of the corresponding hydrodefluorinated products of mono- and perfluorinated arenes in good to excellent yields (up to 98%) (Fig. 4). Labelling experiments revealed that deuterium incorporation

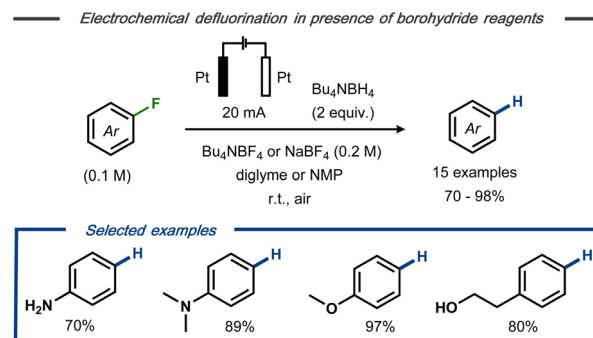


Fig. 4 Electrochemical hydrodefluorination of aryl fluorides in the presence of borohydride reagents.³⁰



occurred to only a minor extent in the presence of sodium borodeuteride, whereas GC-MS analysis indicated the presence of tributylamine in the crude reaction mixture when Bu_4NBF_4 was used as electrolyte. Based on these findings, it was hypothesized that the tetrabutylammonium cation serves as hydrogen source rather than the hydride reagent. Similar to Kariv-Miller's proposal,^{25,26} the cathodic defluorination was suggested to proceed *via* stepwise dissociative electron transfer with a radical anion intermediate that decomposes to the corresponding fluoride and an open shell aryl species. Reductive radical-polar crossover³¹ by a second SET to the aryl radical forms the corresponding aryl anion and, subsequently, the final product upon protonation.

The formation of aryl radicals *via* stepwise dissociative electron transfer with an intermediate radical anion is a commonly invoked mechanism for electrochemical $\text{C}(\text{sp}^2)\text{-F}$ bond cleavage in aryl fluorides (Fig. 5, top).^{25,28,30,32} Nevertheless, alternative mechanisms have been proposed in certain cases, suggesting that an initially formed arylfluoride radical anion can undergo chemical steps prior to C-F bond cleavage.^{22,33} An interesting case of a substrate-dependent mechanistic switch was reported by Muthukrishnan and Sangaranarayanan for electrochemical defluorination of fluorobenzoates (Fig. 5, bottom).³⁴ Voltammetric studies of methyl-2-fluorobenzoate at high scan rates revealed a reversible electron transfer, indicative of a stable radical anion with decomposition to the corresponding aryl radical and fluoride being rate limiting. In contrast, methyl-4-fluorobenzoate was found to undergo electro-dimerization rather than hydrodefluorination. After systematic voltammetric studies and with support by analogous dimerization of 4-fluorobenzonitrile³⁵ and pentafluoronitrobenzene,³⁶ the authors concluded that the dimerization likely occurs between initially formed radical anions, followed by C-F bond cleavage. DFT calculations for both fluorobenzoate substrates indicated that the π^* orbital of the extended aromatic system is the initial electron acceptor, with the charge subsequently

being transferred to the C-F σ^* bond and resulting bond cleavage.

Electroreductive polymerization of polyhalogenated arenes with concomitant C-X bond cleavage was recently reported by Ekinici and co-workers.³⁷ In this reductive synthesis, polyaromatic compounds were formed on the surface of silicon or gold cathodes from polyhalogenated starting materials, including hexafluorobenzene, and were subsequently treated thermally to form graphene-like films. Mechanistically, it was proposed that the coupling reaction either occurs between two radical anions, similar to the mechanism invoked for dimerization of methyl-4-fluorobenzoate,³⁴ or *via* dimerization of neutral aryl radicals. The latter coupling type is commonly deemed unfavorable due to the propensity of aryl radicals to undergo further electron transfers or hydrogen atom transfer events.^{38,39} However, voltammetric studies of hexafluorobenzene suggested that the dissociative electron transfer may take place in a concerted rather than a stepwise manner under the applied conditions. Alternative pathways, such as intermediate aryl anions taking part in nucleophilic aromatic substitutions ($\text{S}_{\text{N}}\text{Ar}$), were not discussed.

Senboku *et al.* reported a defluorinative electrocarboxylation protocol for polyfluorinated arenes with carbon dioxide.⁴⁰ Constant current electrolysis in an undivided cell equipped with a Pt cathode and a Mg anode at -40 °C resulted in mono-carboxylated polyfluorobenzoic acids as products in moderate to good yields and high regio- and chemoselectivity (Fig. 6, top). Lower yields were obtained using either higher reaction temperature or higher current density. Methoxy and acetoxy groups, as well as a benzylic alcohol protected as the tetrahydropyran (THP) acetal, were tolerated under the applied conditions and resulted in selective carboxylation in *para* position (Fig. 6, bottom). The transformation was proposed to proceed *via* stepwise dissociative electron transfer, followed by radical-polar crossover to form the aryl anion (see Fig. 5, top). This species would, in turn, react with CO_2 to form the magnesium carboxylate by interaction with the ions from the dissolving metal anode and, eventually, the polyfluorinated carboxylic acid products upon acidic workup.

Electrocarboxylation with $\text{C}(\text{sp}^2)\text{-F}$ bond cleavage was reported by Xie *et al.* using *gem*-difluoroalkenes as starting

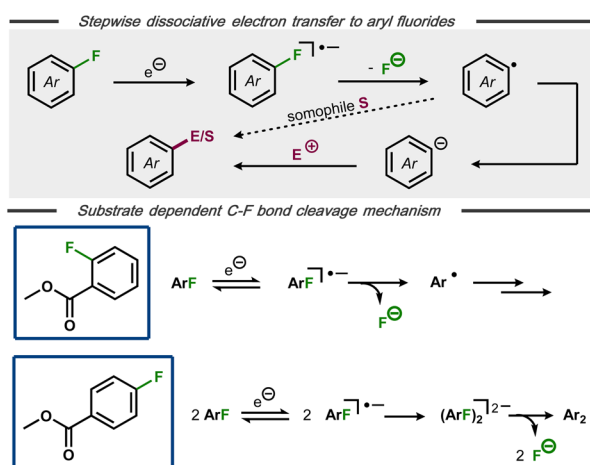


Fig. 5 Top: Mechanism for stepwise defluorination of aryl fluorides. Bottom: Substrate dependent pathways for electroreductive defluorination of methylfluorobenzoates.³⁴

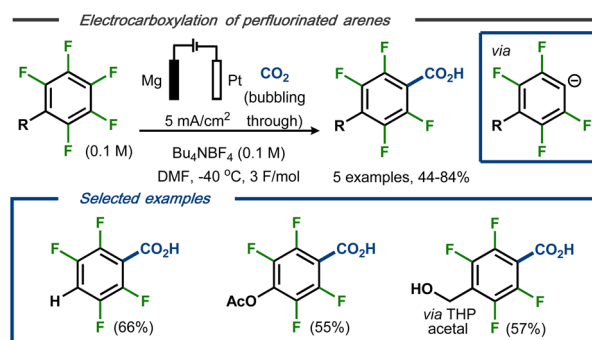


Fig. 6 Defluorination/carboxylation of polyfluorinated arenes.⁴⁰



material.⁴¹ Under constant current electrolysis in an undivided cell, 20 α -fluoroacrylic acids were demonstrated with up to 83% yield and 20 : 1 Z : E selectivity (Fig. 7, top). The protocol displayed tolerance towards functional groups such as carboxylic esters and amides, sulfones, nitriles, and alkynes and allowed for selective dehalogenation of the olefinic fluorides, whereas both aromatic and benzylic fluorides remained intact. Based on cyclic voltammetry studies, and in contrast to Senboku's anionic electrocarboxylation mechanism,⁴⁰ it was proposed that the initially formed radical anion is involved in the C-C bond formation prior to C-F bond cleavage (Fig. 7, bottom).^{22,33-36} Single electron reduction of the resulting C-centered radical followed by fluoride displacement *via* an E1cB mechanism would thereafter furnish the product as carboxylate. Dissolution of the nickel anode was proposed to serve as the oxidative counter reaction in the process. For synthetic ease, the carboxylate products were converted to the corresponding methyl esters as the final step to facilitate isolation and analysis.

Xu *et al.* reported on electrocatalytic hydrodefluorination protocols for fluorinated aromatic compounds.^{42,43} In a divided cell equipped with either a Rh- or a Rh-Pd alloy-modified Ni foam cathode with phosphate buffer (pH ~3) and a graphite anode at constant current electrolysis, hydrodefluorinated products were formed in high yields. In some cases, the aromatic hydrodefluorinated products underwent further electroreduction to afford saturated counterparts. For example, 4-fluorophenol was transformed into phenol (~5%), cyclohexanone (~25%) and cyclohexanol (~70%) as end products using the Rh-Pd alloy-modified Ni cathode (Fig. 8).⁴² Such reduction

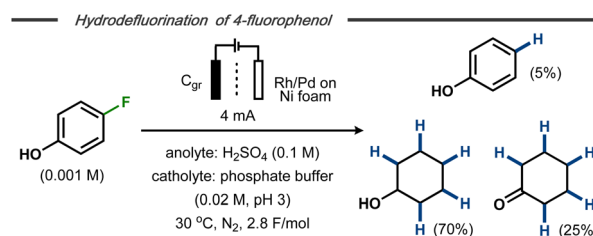


Fig. 8 Hydrodefluorination/hydrogenation of fluorophenol on transition metal modified cathode.⁴³

of the aromatic ring had previously been reported by Trnková and co-workers for fluorobenzene using a polycrystalline Pt cathode in aqueous H_2SO_4 .⁴⁴ Mechanistic studies and control experiments indicated that the hydrodefluorination on the modified Ni foam was likely to follow an indirect hydrogenation mechanism with *in situ* formed metal hydrides on the cathode surface, similar to hydrogenations using late transition metal catalysts and hydrogen gas.⁴⁵

Lambert and Huang recently developed an electrophotocatalytic S_NAr reaction of non-activated aryl fluorides and nucleophiles to forge C-N and C-O bonds.⁴⁶ Using 10 mol% 2,3-dichloro-5,6-dicyanoquinone (DDQ) as photocatalyst under blue light irradiation at constant potential, 40 examples of defluorinated coupling products were formed in moderate to high yields (Fig. 9, top). In contrast to fully electrochemical protocols, this photoelectrocatalytic protocol was claimed to undergo an oxidative pathway to afford the C-F bond cleavage (Fig. 9, bottom). Mechanistically, photoexcitation of DDQ results in an excited catalyst species able of oxidizing the fluoroarene. Nucleophilic attack on the resulting radical cation intermediate furnishes a radical that, in turn, undergoes a single-electron reduction to the corresponding anion. Re-aromatization with expulsion of the fluoride results in the S_NAr product. Due to the poor reducing power of the radical anion of DDQ, this species was deemed insufficient to mediate the reduction of the aromatic radical intermediate and, hence, that this step as well as the re-oxidation of the photocatalyst are accomplished electrochemically.

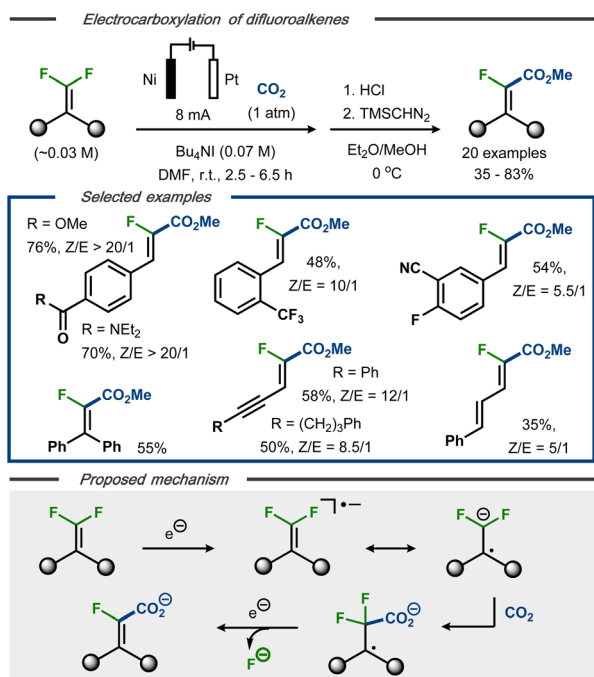


Fig. 7 Top: Defluorination/carboxylation of *gem*-difluoroalkenes.⁴¹ Bottom: Proposed mechanism.

3. Cleavage of C(sp³)-F bonds

C(sp³)-F bonds have lower bond dissociation energies (BDE) compared to their C(sp²)-F counterparts, with C-F bonds in π -activated positions being weaker compared to fully aliphatic counterparts (Fig. 1). The latter property is reflected in the number of synthetic protocols available for cleavage of the different aliphatic fluorides using any defluorination method, including electrosynthesis. The BDE of C(sp³)-F bonds increase with the degree of fluorination of the carbon. For example, the BDE has been determined to increase from 99 kcal mol⁻¹ to ~115 kcal mol⁻¹ for the C-F bond when going from α -fluorotoluene to α,α,α -trifluorotoluene.⁴⁷ As a result, selective partial defluorination of polyfluorinated sites



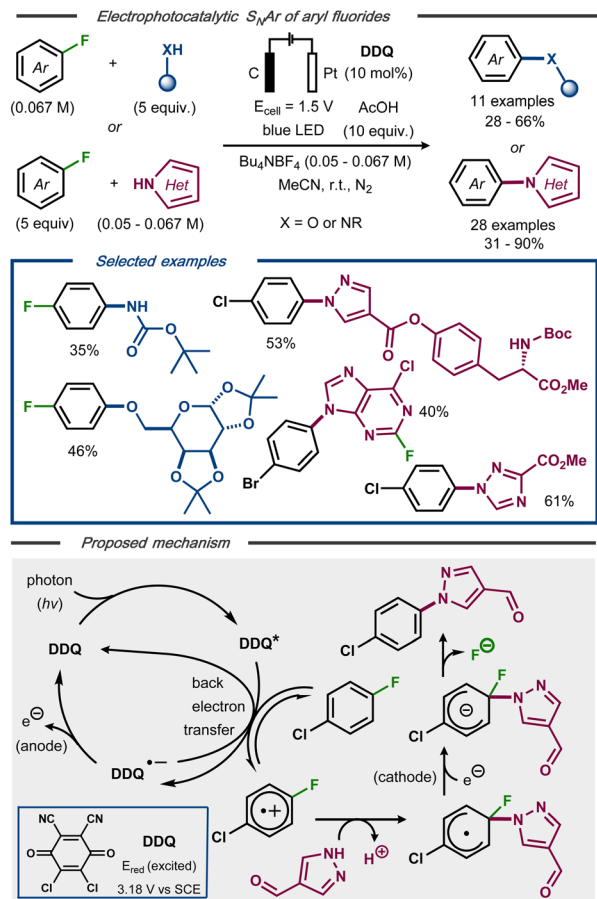


Fig. 9 Top: Electrophotocatalytic nucleophilic aromatic substitution of aryl fluorides.⁴⁶ Bottom: Proposed mechanism.

is challenging since the C–F bonds in the products of the sequential defluorination are more easily cleaved compared to those in the starting material.

3.1. Benzylic and allylic C(sp³)–F bonds

An early electrochemical reduction of a benzylic CF₃-group was reported by Lund *et al.* with polarography studies as well as preparative electrolysis with a dropping mercury cathode.⁴⁸ The potentiostatic hydrodefluorination of 1,1-dioxo-6-(trifluoromethyl)-3,4-dihydro-2H-1λ⁶,2,4-benzothiadiazine-7-sulfonamide was performed in 30% aqueous methanol and resulted in 72% yield of the desired product after 6.3 F of charge at a potential of –1.70 V vs. SCE (Fig. 10).

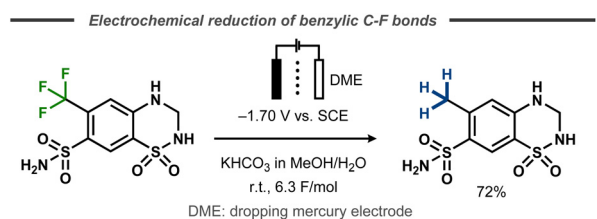


Fig. 10 Defluorination of benzylic fluorides.⁴⁸

In contrast to the stepwise reduction of other trihalomethyl compounds,⁴⁸ Lund *et al.* proposed that the cleavage of the three C–F bonds takes place in one single step. This claim was not supported by subsequent studies, that instead suggest that benzylic C–F bonds in *p*-substituted trifluoromethyl benzenes take place in a stepwise manner in polar aprotic solvents like DMF with cathode materials such as Hg, Pt and Pb.^{49–55} Similar to the postulated stepwise dissociative electron transfer for many aryl fluorides (Fig. 5, top), initial single electron reduction of the trifluoromethyl benzene is generally believed to form a stabilized radical anion that decomposes to a benzylic radical with release of fluoride. Reductive radical-polar crossover upon a second electron transfer results in a benzylic anion, which can be quenched by a proton (Fig. 11)^{50–52} or other electrophiles.^{56,57} In the case of 4-(trifluoromethyl)acetophenone, hydrodefluorinated pinacol products were observed by Liotier *et al.* along with the hydrodefluorinated alcohol in aqueous alkaline medium (pH 10).⁵⁸

One example of benzylic defluorination was reported by Périchon and co-workers using a SmCl₃-catalyzed electrochemical hydrodehalogenation protocol in an undivided cell using a nickel foam cathode and a sacrificial magnesium anode.²⁹ In this case, α,α-trifluorotoluene was transformed into a mixture of α,α-difluorotoluene, α-fluorotoluene and toluene and in 50% yield (Fig. 12). Addition of D₂O did not result in deuterium incorporation for a benchmark chloride

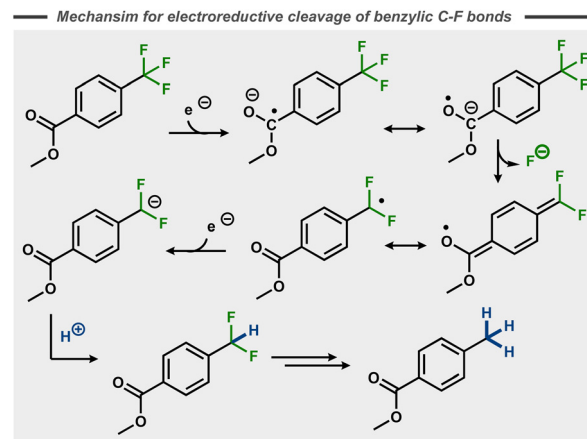


Fig. 11 Reductive C–F bond cleavage mechanism for a trifluoromethyl arene.⁵⁰

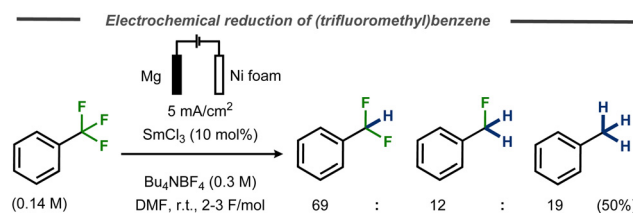


Fig. 12 Samarium-catalyzed C–F bond cleavage in trifluoromethyl benzene.²⁹



substrate and it was concluded that the hydrogen in the product originates from either the tetraalkylammonium ion in the supporting electrolyte or the solvent. While cyclization experiments using 1-allyloxy-2-chlorobenzene suggested that a radical mechanism was operating in the presence of samarium, no mechanistic details or discussion on the valence state of the catalyst were provided.

Périchon and co-workers demonstrated that electroreductive defluorination could be coupled with C–C bond formation using electrophiles such as CO₂, DMF or acetone to form carboxylic acids, aldehydes, or tertiary alcohols.⁵⁷ The electrolyses were carried out at room temperature in DMF containing the corresponding trifluoromethyl aryl compound, using a stainless steel cathode and a sacrificial anode (magnesium or aluminium rod) (Fig. 13, top). For synthesis of tertiary alcohols, acetone was used as a co-solvent (10 vol%) to furnish the products in yields up to 80% yield, whereas up to 62% yield was achieved in formylation reactions with DMF. In these cases, the crude reactions were poured into acetic anhydride and hydrolyzed to furnish diacetate products. For electrocarboxylation, carbon dioxide was bubbled through the solution at atmospheric pressure to provide benzoic acid products in up to 70% yield. The same strategy was used by Senboku and co-workers to trap benzylic anionic intermediates bearing benzylic pentafluoroethyl groups in the synthesis of tetrafluorinated analogues of the NSAIDs fenoprofen and ketoprofen (Fig. 13, bottom).⁵⁹ In this case, the electrochemical reduction was conducted in DMF using a Pt cathode and a Mg anode in an undivided cell with CO₂ bubbled through the reaction medium.

Benzylic C–F bond cleavage with trapping of the intermediate anion has also been achieved with TMSCl as electrophile,

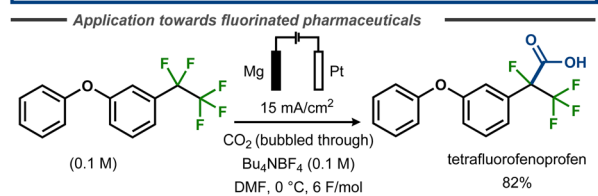
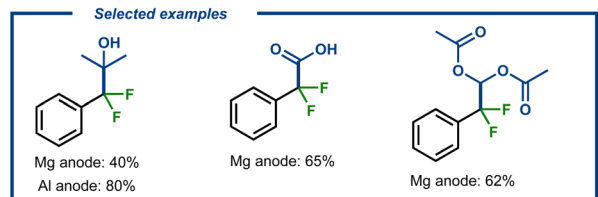
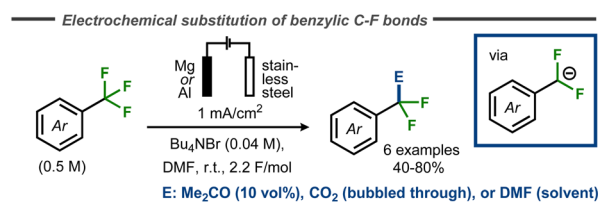


Fig. 13 Top: Defluorinative functionalization of benzyl fluorides.⁵⁷ Bottom: Defluorinative electrocarboxylation for polyfluorinated pharmaceutical analogues.⁵⁹

forming trimethylsilyldifluoromethylbenzenes.^{55,56} Marzouk *et al.* demonstrated that fluorides in α,α,α -trifluorotoluene substituents can subsequently be replaced by TMS groups in a THF/DMPU mixture, thus avoiding carcinogenic HMPA, by simply increasing the equivalents of charge applied.⁵⁶ As such, the synthesis of corresponding mono-, bis- or tris-trimethylsilyl derivatives using aluminium as a sacrificial anode was enabled in yields up to 70%, 60% and 50% respectively (Fig. 14). This reaction was scaled up to 1 mol of starting material, using a tubular flow cell.

Recently, a facile electroreductive system for the removal of various functional groups was developed by Xia and co-workers, using triethylamine (Et₃N) as sacrificial reductant in combination with inert platinum electrodes in an undivided cell at constant current.⁶⁰ Amongst examples such as reduction of cyano groups and azoles, the authors reported complete reduction of benzylic CF₃ – groups in yields up to 89% (Fig. 15). Similarly, Huang and co-workers provided one example of C–F bond cleavage in trifluoromethylbenzene in a moderate yield of 35% at low conversion using platinum electrodes in diglyme and Bu₄NBF₄ (0.2 M) under air at room temperature with NaBH₄ as additive.⁶¹

3.2. C–F bond cleavage in α -CF₃ alkenes

Recently, Cheng and co-workers developed a defluorinative method to convert α,α,α -trifluoromethyl cinnamates to *gem*-difluorostyrenes in moderate to good yields, using an electroreductive approach (Fig. 16).⁶² The reactions were performed in

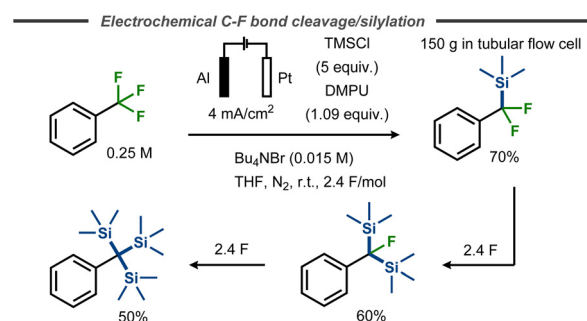


Fig. 14 Defluorinative silylation of benzyl fluorides with TMSCl.⁵⁶

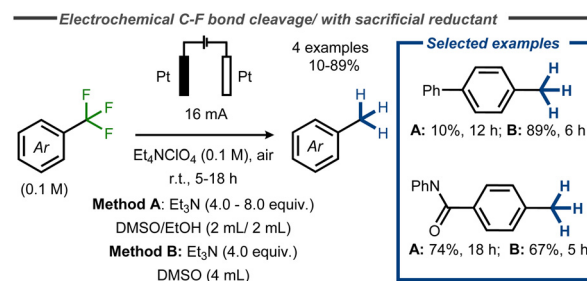


Fig. 15 Electroreductive C–F bond cleavage in benzyl fluorides with Et₃N as sacrificial reductant.⁶⁰



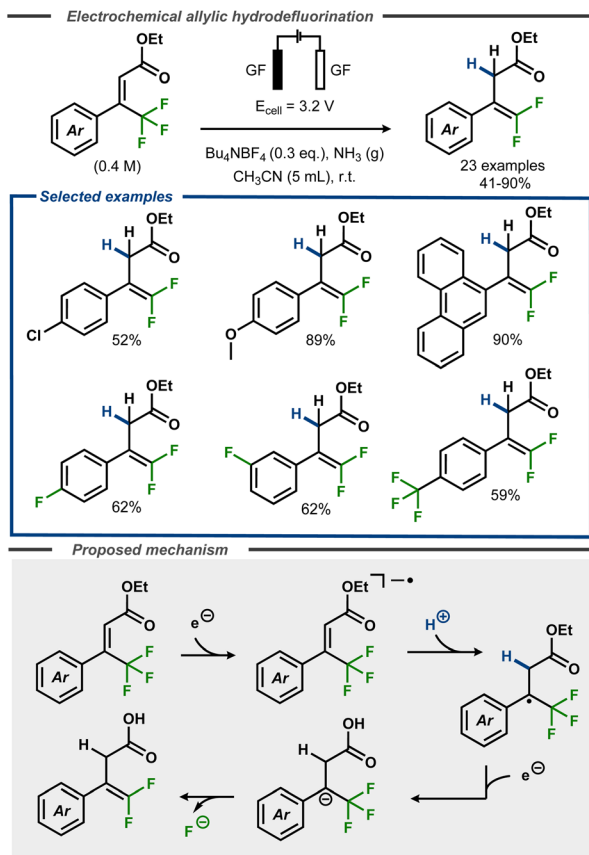


Fig. 16 Electroreductive formation of *gem*-difluorostyrenes from α,α,α -trifluoromethyl cinnamates.⁶²

acetonitrile with controlled potential, using graphite felt electrodes under an ammonia atmosphere that acted as proton donor. The method was demonstrated to proceed well with electron-rich substrates *e.g.* methoxy groups (89%) or large π -systems (90%), whereas moderate yields were observed for substrates with functional groups such as chloride (52%). Selective defluorination was observed for substrates with fluoride and trifluoromethyl substituents, furnishing the difluoroalkene products in moderate yields. The mechanism was investigated using cyclic voltammetry and square wave voltammetry, indicating a net two-electron transfer. The first electron transfer was proposed to convert the substrate to the corresponding radical anion that, upon protonation, forms the corresponding radical. A second electron transfer furnishes the benzylic anion, which undergoes an E1cB elimination of fluoride to form the desired product (Fig. 16, bottom).

Along the same lines, Gao *et al.* recently developed a regioselective electrochemical γ -carboxylation of α -CF₃ alkenes using CO₂ under constant current conditions.⁶³ The transformation was carried out in DMF with Bu₄NClO₄ as supporting electrolyte, using platinum electrodes (Fig. 17). The method proceeded with a good functional group tolerance, with yields up to 84%, including examples such as sugars and steroids. Supported by cyclic voltammetry studies, radical trap experi-

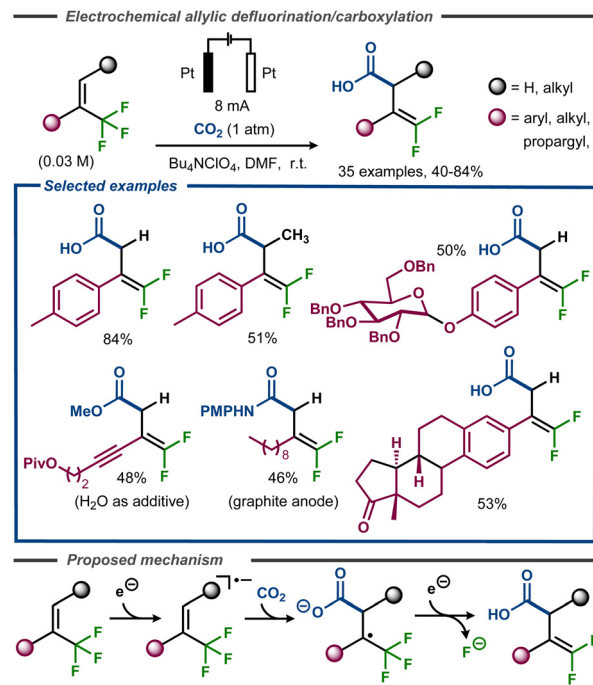


Fig. 17 Electroreductive formation of carboxylated *gem*-difluorostyrenes from α -trifluoromethyl styrenes.⁶³

ments, as well as DFT calculations, the reaction mechanism was believed to be of a similar type of that proposed by Cheng and co-workers (Fig. 16).⁶² After an initial reduction of the trifluoromethyl alkene to the corresponding radical anion, addition of CO₂ would take place. The resulting radical carboxylate would thereafter undergo a second electron transfer to the corresponding enolate that decomposes to the *gem*-difluoroalkene product with release of fluoride (Fig. 17, bottom). The anodic counter reaction was proposed to be oxidation of either DMF or water.

Electroreductive C–F bond cleavage and concomitant functionalization of α -trifluoromethyl styrenes to *gem*-difluoroalkenes have recently been reported using radical precursors such as alkyl halides, redox active esters and Katritzky salts, as well as aryl halides (Fig. 18).^{64–68} Ni, Guo and Wang and co-workers demonstrated that Katritzky salts could be successfully used as coupling partners in DMSO under galvanostatic conditions with a sacrificial Zn anode and a Ni foam cathode, furnishing products in up to 85% yield (Fig. 18, Method A).⁶⁴ Masson, Claraz and co-workers demonstrated coupling of Katritzky salts as well as NHP esters in yields up to 94%, including amino acid derived alkyl NHP esters (Fig. 18, Method B).⁶⁵ Similarly, Xia, Guo and co-workers demonstrated that α -trifluoromethyl styrene substrates react with NHP esters and Katritzky salts, as well as alkyl halides, in yields up to 76% (Fig. 18, Method C).⁶⁶ In the cases where a Fe sacrificial anode was not used, amine bases such as triethylamine or DABCO were employed as sacrificial reductants, similar to what was previously reported for electroreduction of trifluoromethyl arenes.⁶⁰ Qi, Huang and Lei and co-workers reported yields up



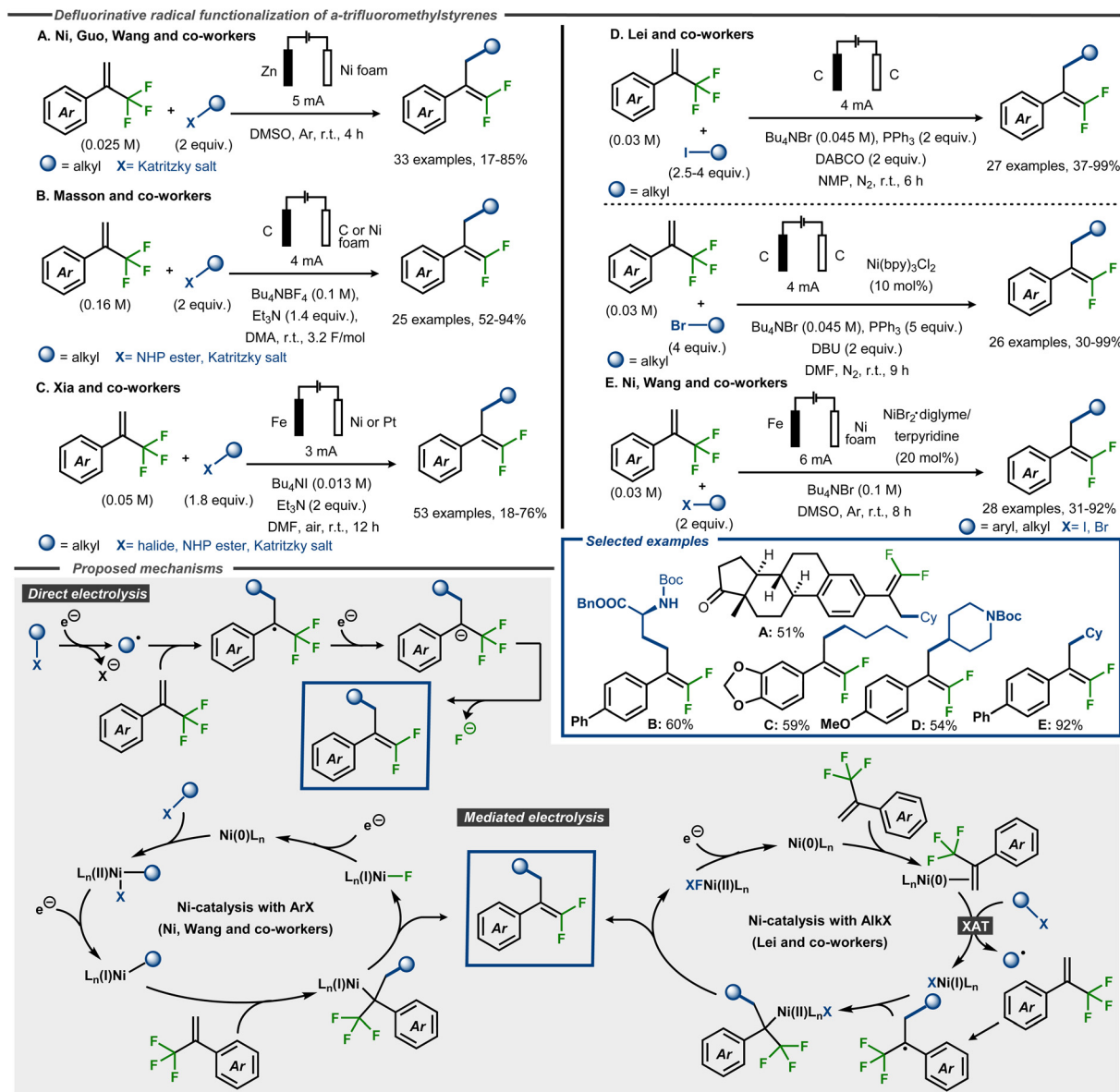


Fig. 18 Electrocyclic formation of γ -functionalized *gem*-difluorostyrenes from α -trifluoromethyl styrenes and radical precursors.^{64–68}

to 99% with alkyl halides as coupling partners, using a direct electrolysis method for alkyl iodides (Fig. 18, Method D) or an indirect electrolysis method with a Ni(bpy)-complex in catalytic amounts for alkyl bromides.⁶⁷ In addition, the former method was also demonstrated using one Katritzky salt and one NHP ester as coupling partner to furnish the desired product in moderate to good yield, whereas the latter method resulted in moderate yield using an alkyl chloride as partner. Oxidation of the phosphine or amine additives were postulated as the oxidative counter reactions for the process. A fully catalytic system was reported by Ni and Wang and co-workers, who demonstrated that alkyl and aryl halides react with α -trifluoromethyl styrene substrates using a Ni catalyst with a terpyridine ligand, furnishing products in yields up to 93% (Fig. 18, Method E).⁶⁸

Mechanistically, the direct electrolytic methods are thought to be initiated by cathodic dissociative electron transfer to the radical precursor to provide an aliphatic carbon-centered radical intermediate. This radical would thereafter attack the α -trifluoromethyl alkene to form a C–C bond and a benzylic stabilized α -CF₃ carbon radical. Cathodic reduction of this radical furnishes α -CF₃ carbanion that undergoes a β -fluoride elimination to afford the *gem*-difluoromethyl alkene product (Fig. 18). For the Ni-catalyzed protocols, the Lei group and Ni and Wang and co-workers propose different mechanisms. The former group utilizes alkyl bromides as coupling partners and suggest, supported by DFT calculations, that the Ni(0) catalyst initially coordinates the alkene to thereafter interact with the alkyl bromide in an XAT event to form an alkyl radical and a Ni(I)–Br complex. This open-shell species adds to the CF₃-



alkene substrate analogous to the indirect mechanism, whereas the Ni(I) species traps the resulting benzylic radical to form a Ni(II)-complex. After a two-electron reduction and β -fluoride elimination, the product is released and the Ni(0) catalyst is regenerated (Fig. 18, bottom). In contrast, Ni and Wang and co-workers do not speculate about the mechanism for alkyl halides but focuses on aryl halides as coupling partners. In this case, the Ni(0) catalyst is proposed to undergo an oxidative addition with the aryl halide, thereby forming a Ni(II) ArX complex. This species undergoes a single electron reduction at the cathode to furnish the corresponding Ni(I)Ar complex. Migratory insertion of the trifluoromethyl styrene derivative generates a Ni(I)R species that is expected to undergo rapid β -fluoride elimination to deliver the desired product. Another cathodic reduction would regenerate the Ni(0) species from Ni(I)F to close the catalytic cycle (Fig. 18, bottom).

3.3. Cleavage of C(sp³)-F bonds in α -position to carbonyls

The first mechanistic insights into cleavage of C-F bonds in α -position to carbonyl carbons were reported by Elving and Leone in 1956.⁶⁹ In this case, polarographic studies of the reduction of 2-fluoroacetophenone indicated two cathodic waves that were interpreted as initial cleavage of the C-F bond, and subsequent reduction of the resulting acetophenone to the corresponding alcohol. Stocker and Jenevein reported that initial defluorination of α,α,α -trifluoroacetophenone to acetophenone was followed by pinacolization (Fig. 19, top) with the selectivity being dependent on applied potential and reaction time.⁷⁰ However, later cyclic voltammetry studies by Yang *et al.* suggested that the fluorinated pinacol forms first and is subsequently reduced (Fig. 19, bottom).⁷¹ Mechanistically, it was suggested that the pinacol formation occurs *via* radical-radical coupling between one protonated and one unprotonated electrochemically formed ketyl radical, whereas certain substrates were proposed to undergo a second electron transfer and two protonations to form the corresponding alcohol.

Evans and Kopilov reported that defluorination of α -fluoroacetanilide took place during their attempts to electro-

chemically remove the electron-poor amide protecting group.⁷² Instead of furnishing the free aniline, potentiostatic electrolysis of α,α,α -trifluoroacetanilide in a divided cell with a mercury pool cathode resulted in a mixture of 65% α -fluoroacetanilide and 35% acetanilide (Fig. 20).

Uneyama *et al.* demonstrated successful trapping of the intermediate after the initial hydrodefluorination of trifluoromethyl ketones as well as trifluoroketimines with silyl chloride reagents, isolating difluoro silyl enol ethers and difluoro silyl enamines in yields up to 85% and 78%, respectively (Fig. 21).^{73,74} In the former case, a Pb cathode and carbon anode was utilized to achieve electroreductive defluorination of a range of trifluoromethyl ketones in anhydrous acetonitrile using an H-type divided cell (Fig. 21, top). The reaction was reported to proceed efficiently with aromatic substrates, resulting in yields up to 85%, whereas aliphatic substrates furnished the corresponding products in yields up to 55%. Similar conditions were used for the electroreductive defluorination of

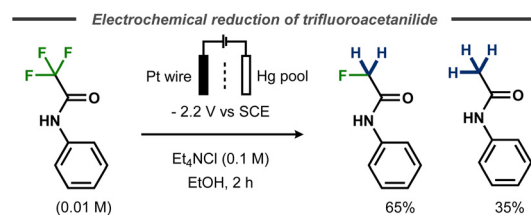


Fig. 20 Electrochemical defluorination of α,α,α -trifluoroacetanilide.⁷²

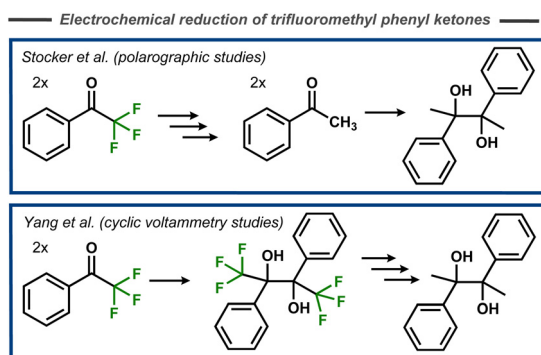
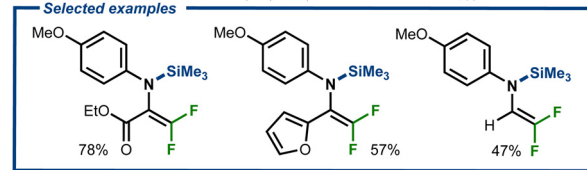
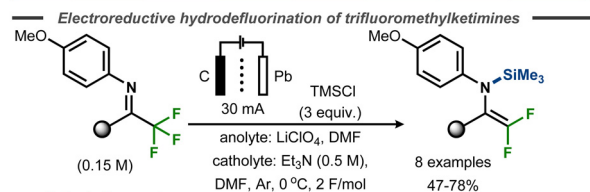
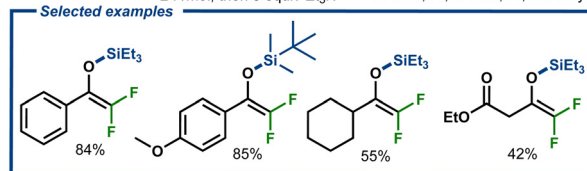
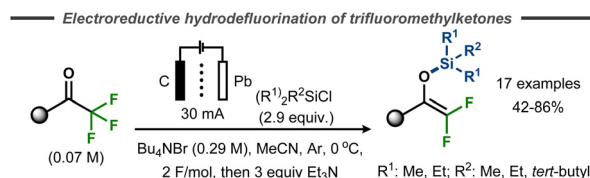


Fig. 19 Electroreductive pinacolization of α,α,α -trifluoroacetophenone.^{70,71}

Fig. 21 Electroreductive defluorinative silylation of trifluoromethyl ketones.⁷³



ketimines (Fig. 21, bottom). The authors did not provide any mechanistic details for the transformations.

The methodology was expanded to trifluoroacetic acid derivatives to yield 2,2-difluoro-2-trimethylsilylacetate and difluoroketene silyl (*O,O*-, *O,S*-, and *O,N*-) acetals.⁷⁵ These findings stood in contrast to those of Stepanov *et al.* who reported Claisen adducts under similar conditions.⁷⁶ After a thermal isomerization with silyl migration from oxygen to carbon, the corresponding products were successfully coupled with a range of electrophiles (aldehydes, ketones, imines, acyl- and alkylhalides) to give α -alkylated- α,α -difluoroacetates in good to excellent yields (Fig. 22). This isomerization could either be carried out in a stepwise procedure with electrolysis carried out at low reaction temperature, or direct by electrolysis at 50 °C.

Recently, Lennox *et al.* demonstrated selective monodefluorination of α,α,α -trifluoromethylketones in a single step to access a broad scope of difluoromethylketones (Fig. 23).⁷⁷ Using a divided cell setup, the reaction was reported to proceed efficiently on platinum or nickel cathodes with TMSCl present as a radical anion trapping agent *via* a proposed silyl enol ether intermediate of Uneyama-type.⁷³ The transformation tolerated heterocyclic groups such as indoles as well cyclopropanes or even aryl fluorides incorporated in the substrates. The reaction performs best with electron-rich and simple alkyl substrates (up to 95%), whereas unprotected indoles as well as protected indoles bearing cyano groups resulted in lower yields (45 and 42%, respectively). Similar to other electrochemical hydrodefluorination protocols, tetraalkylammonium salts were found to act as reductively stable,

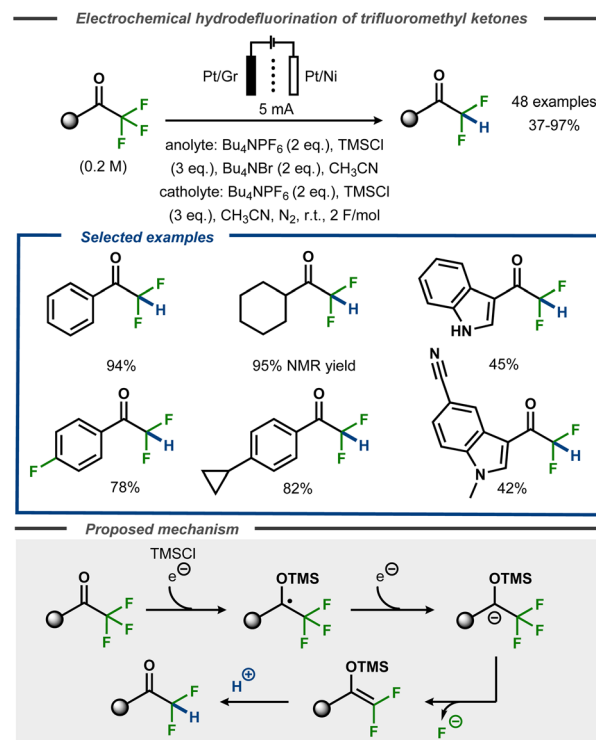


Fig. 23 Selective monoreduction of α,α,α -trifluoromethyl ketones.⁷⁷

masked proton sources upon Hoffmann elimination. Supported by DFT calculations, the transformation was proposed to proceed *via* cathodic reduction of the substrate to furnish a transient ketyl radical anion that is trapped with TMSCl to furnish an O-TMS protected benzylic radical. A second electron transfer furnishes the corresponding anion that undergoes loss of fluoride to form the O-TMS protected *gem*-difluoroalkene that, in turn, furnishes the hydrodefluorinated product upon protonation. Voltammetric studies suggested that oxidation of bromide to tribromide was the anodic counter reaction.

4. Conclusions

In contrast to their heavier congeners, fluorocarbons are challenging from a reactivity perspective for any synthetic strategy. This challenge is present also using an electrochemical approach, as evident from the limited number of synthetically relevant examples of C–F bond cleavage for both sp^2 - and sp^3 -hybridized carbon centers. With only a few exceptions, available electrochemical protocols are only amenable for cleaving C–F bonds in close proximity to π -systems *via* similar reductive mechanisms. While a limited number of chemical protocols are at hand for C(sp^3)–F bond activation in fully aliphatic positions, electrochemical variants for selective activation/functionalization are yet to be developed. In depth mechanistic understanding, alternative activation modes and mediated (photo)electrolysis are likely keys to enable new fluoroselective

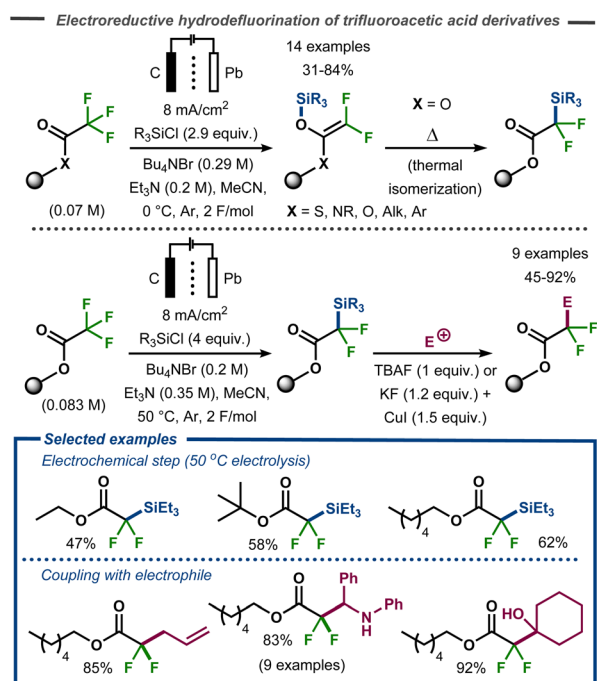


Fig. 22 Electrochemical defluorinative C–Si bond formation and subsequent functionalization.⁷⁵



synthetic methodology that can open new reactivity space and retrosynthetic disconnections, as well as facilitate efficient remediation of fluorinated organic pollutants.^{19,20,78}

Author contributions

All authors contributed to the writing of this review.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Financial support from the Swedish Research Council (grant 2021-05551), Formas (grant 2021-00678), the Swedish Foundation for Strategic Research (grant FFL21-0005), Stiftelsen Olle Engkvist Byggmästare, Magnus Bergvalls stiftelse, Stiftelsen Lars Hiertas Minne, and KTH Royal Institute of Technology is gratefully acknowledged.

References

- (a) M. Shimizu and T. Hiyama, *Angew. Chem., Int. Ed.*, 2005, **44**, 214–231; (b) M. Schlosser, *Angew. Chem., Int. Ed.*, 2006, **45**, 5432–5446; (c) K. Müller, C. Faeh and F. Diederich, *Science*, 2007, **317**, 1881–1886; (d) J. Wang, M. Sánchez-Roselló, J. L. Aceña, C. del Pozo, A. E. Sorochinsky, S. Fustero, V. A. Soloshonok and H. Liu, *Chem. Rev.*, 2014, **114**, 2432–2506; (e) E. P. Gillis, K. J. Eastman, M. D. Hill, D. J. Donnelly and N. A. Meanwell, *J. Med. Chem.*, 2015, **58**, 8315–8359; (f) Y. Ogawa, E. Tokunaga, O. Kobayashi, K. Hirai and N. Shibata, *iScience*, 2020, **23**, 101467–101467; (g) P. Jeschke, *ChemBioChem*, 2004, **5**, 570–589; (h) R. Berger, G. Resnati, P. Metrangolo, E. Weber and J. Hulliger, *Chem. Soc. Rev.*, 2011, **40**, 3496–3508; (i) R. Ragni, A. Punzi, F. Babudri and G. M. Farinola, *Eur. J. Org. Chem.*, 2018, 3500–3519.
- L. Pauling, *The Nature of the Chemical Bond and the Structure of Molecules and Crystals: An Introduction to Modern Structural Chemistry*, Cornell University Press, Ithaca, NY, 1939.
- D. O'Hagan, *Chem. Soc. Rev.*, 2008, **37**, 308–319.
- S. J. Blanksby and G. B. Ellison, *Acc. Chem. Res.*, 2003, **36**, 255–263.
- C. Cao, *Sci. China, Ser. B: Chem.*, 2009, **52**, 943–951.
- T. Stahl, H. F. T. Klare and M. Oestreich, *ACS Catal.*, 2013, **3**, 1578–1587.
- D. A. Everson and D. J. Weix, *J. Org. Chem.*, 2014, **79**, 4793–4798.
- F. Juliá, T. Constantin and D. Leonori, *Chem. Rev.*, 2022, **122**, 2292–2352.
- X. Dong, J. L. Röckl, S. R. Waldvogel and B. Morandi, *Science*, 2021, **371**, 507–514.
- F. Zhao, W. Zhou and Z. Zuo, *Adv. Synth. Catal.*, 2022, **364**, 234–267.
- T. Fujita, K. Fuchibe and J. Ichikawa, *Angew. Chem., Int. Ed.*, 2019, **58**, 390–402.
- (a) H. Amii and K. Uneyama, *Chem. Rev.*, 2009, **109**, 2119–2183; (b) H. Torrens, *Coord. Chem. Rev.*, 2005, **249**, 1957–1985; (c) M. K. Whittlesey and E. Peris, *ACS Catal.*, 2014, **4**, 3152–3159; (d) H. Kameo, H. Yamamoto, K. Ikeda, T. Isasa, S. Sakaki, H. Matsuzaka, Y. García-Rodeja, K. Miqueu and D. Bourissou, *J. Am. Chem. Soc.*, 2020, **142**, 14039–14044.
- (a) Q. Shen, Y.-G. Huang, C. Liu, J.-C. Xiao, Q.-Y. Chen and Y. Guo, *J. Fluor. Chem.*, 2015, **179**, 14–22; (b) H.-J. Ai, X. Ma, Q. Song and X.-F. Wu, *Sci. China: Chem.*, 2021, **64**, 1630–1659; (c) K. Balaraman and C. Wolf, *Sci. Adv.*, 2022, **8**, eabn7819.
- (a) W. D. Jones, *Dalton Trans.*, 2003, 3991–3995; (b) A. G. M. Barrett, M. R. Crimmin, M. S. Hill, P. B. Hitchcock and P. A. Procopiu, *Angew. Chem., Int. Ed.*, 2007, **46**, 6339–6342; (c) M. Klahn and U. Rosenthal, *Organometallics*, 2012, **31**, 1235–1244; (d) T. L. Gianetti, R. G. Bergman and J. Arnold, *Chem. Sci.*, 2014, **5**, 2517–2524; (e) A. M. Träff, M. Janjetovic, L. Ta and G. Hilmersson, *Angew. Chem., Int. Ed.*, 2013, **52**, 12073–12076; (f) W. Huang and P. L. Diaconescu, *Organometallics*, 2017, **36**, 89–96.
- (a) H. Amii, T. Kobayashi, Y. Hatamoto and K. Uneyama, *Chem. Commun.*, 1999, 1323–1324; (b) K. Matsubara, T. Ishibashi and Y. Koga, *Org. Lett.*, 2009, **11**, 1765–1768; (c) C. J. Burns and R. A. Anderson, *J. Chem. Soc., Chem. Commun.*, 1989, 136–137; (d) G. B. Deacon and P. I. MacKinnon, *Tetrahedron Lett.*, 1984, **25**, 783–784; (e) P. L. Watson, T. H. Tulip and I. Williams, *Organometallics*, 1990, **9**, 1999–2009; (f) M. Janjetovic, A. M. Träff, T. Ankner, J. Wettergren and G. Hilmersson, *Chem. Commun.*, 2013, **49**, 1826–1828.
- (a) S. Li and W. Shu, *Chem. Commun.*, 2022, **58**, 1066–1077; (b) G. Yan, K. Qiu and M. Guo, *Org. Chem. Front.*, 2021, **8**, 3915–3942; (c) J. Wang, H. Gao, C. Shi, G. Chen, X. Tan, X. Chen, L. Xu, X. Cai, B. Huang and H. Li, *Chem. Commun.*, 2021, **57**, 12203–12217; (d) G. Yan, *Chem. – Eur. J.*, 2022, **28**, e202200231; (e) Z. Wang, Y. Sun, L.-Y. Shen, W.-C. Yang, F. Meng and P. Li, *Org. Chem. Front.*, 2022, **9**, 853–873.
- (a) E. J. Horn, B. R. Rosen and P. S. Baran, *ACS Cent. Sci.*, 2016, **2**, 302–308; (b) B. A. Frontana-Urbe, R. D. Little, J. G. Ibanez, A. Palma and R. Vasquez-Medrano, *Green Chem.*, 2010, **12**, 2099–2119; (c) A. Wiebe, T. Gieshoff, S. Mohle, E. Rodrigo, M. Zirbes and S. R. Waldvogel, *Angew. Chem., Int. Ed.*, 2018, **57**, 5594–5619; (d) A. Shatskiy, H. Lundberg and M. D. Kärkäs, *ChemElectroChem*, 2019, **6**, 4067–4092; (e) C. Margarita and H. Lundberg, *Catalysts*, 2020, **10**, 982–1006.
- (a) M. Yan, Y. Kawamata and P. S. Baran, *Chem. Rev.*, 2017, **117**, 13230–13319; (b) R. D. Little and K. D. Moeller, *Chem. Rev.*, 2018, **118**, 4483–4484; (c) C. Sandford, M. A. Edwards, K. J. Klunder, D. P. Hickey, M. Li, K. Barman, M. S. Sigman,



- H. S. White and S. D. Minter, *Chem. Sci.*, 2019, **10**, 6404–6422; (d) R. Francke and R. D. Little, *ChemElectroChem*, 2019, **6**, 4373–4382; (e) R. D. Little, *J. Org. Chem.*, 2020, **85**, 13375–13390; (f) C. Kingston, M. D. Palkowitz, Y. Takahira, J. C. Vantourout, B. K. Peters, Y. Kawamata and P. S. Baran, *Acc. Chem. Res.*, 2020, **53**, 72–83; (g) E. C. R. McKenzie, S. Hosseini, A. G. C. Petro, K. K. Rudman, B. H. R. Gerroll, M. S. Mubarak, L. A. Baker and R. D. Little, *Chem. Rev.*, 2022, **122**, 3292–3335.
- 19 S. Sharma, N. P. Shetti, S. Basu, M. N. Nadagouda and T. M. Aminabhavi, *Chem. Eng. J.*, 2022, **430**, 13289–132911.
- 20 E. Mousset and K. Doudrick, *Curr. Opin. Electrochem.*, 2020, **22**, 221–227.
- 21 S. H. Langer and S. Yurchak, *J. Electrochem. Soc.*, 1969, **116**, 1228–1229.
- 22 B. H. Campbell, *Anal. Chem.*, 1972, **44**, 1659–1663.
- 23 C. P. Andrieux, C. Blocman and J. M. Savéant, *J. Electroanal. Chem. Interfacial Electrochem.*, 1979, **105**, 413–417.
- 24 A. Kakhr, Y. Mugnier and E. Laviron, *Electrochim. Acta*, 1983, **28**, 1897–1898.
- 25 E. Kariv-Miller and Z. Vajtner, *J. Org. Chem.*, 1985, **50**, 1394–1399.
- 26 E. Kariv-Miller and R. Andruzzi, *J. Electroanal. Chem. Interfacial Electrochem.*, 1985, **187**, 175–186.
- 27 D. M. Loffredo, J. E. Swartz and E. Kariv-Miller, *J. Org. Chem.*, 1989, **54**, 5953–5957.
- 28 A. Profumo, E. Fasani and A. Albini, *Heterocycles*, 1999, **51**, 1499–1502.
- 29 H. Hebri, E. Duñach and J. Périchon, *Synth. Commun.*, 1991, **21**, 2377–2382.
- 30 W. B. Wu, M. L. Li and J. M. Huang, *Tetrahedron Lett.*, 2015, **56**, 1520–1523.
- 31 L. Pitzer, J. L. Schwarz and F. Glorius, *Chem. Sci.*, 2019, **10**, 8285–8291.
- 32 A. Muthukrishnan and M. V. Sangaranarayanan, *Chem. Phys. Lett.*, 2007, **446**, 297–303.
- 33 D. B. Denney, D. Z. Denney and S. P. Fenelli, *Tetrahedron*, 1997, **53**, 9835–9846.
- 34 A. Muthukrishnan and M. V. Sangaranarayanan, *Electrochim. Acta*, 2010, **55**, 1664–1669.
- 35 K. J. Houser, D. E. Bartak and M. D. Hawley, *J. Am. Chem. Soc.*, 1973, **95**, 6033–6040.
- 36 C. P. Andrieux, A. Battle, M. Espin, I. Gallardo, Z. Jiang and J. Marquet, *Tetrahedron*, 1994, **50**, 6913–6920.
- 37 Z. Kudaş, E. Gür and D. Ekinçi, *Langmuir*, 2018, **34**, 7958–7970.
- 38 H. Lund and O. Hammerich, *Organic Electrochemistry*, Marcel Dekker, New York, 4th edn, 2001.
- 39 C. P. Andrieux, C. Blocman, J. M. Dumas-Bouchiat and J. M. Saveant, *J. Am. Chem. Soc.*, 1979, **101**, 3431–3441.
- 40 H. Senboku, K. Yoneda and S. Hara, *Electrochemistry*, 2013, **81**, 380–382.
- 41 S. L. Xie, X. T. Gao, H. H. Wu, F. Zhou and J. Zhou, *Org. Lett.*, 2020, **22**, 8424–8429.
- 42 Y. H. Xu, H. X. Ma, T. J. Ge, Y. Q. Chu and C. A. Ma, *Electrochem. Commun.*, 2016, **66**, 16–20.
- 43 Y. Xu, T. Ge, H. Ma, X. Ding, X. Zhang and Q. Liu, *Electrochim. Acta*, 2018, **270**, 110–119.
- 44 N. Musilova-Kebrlova, P. Janderka and L. Trnkova, *Collect. Czech. Chem. Commun.*, 2009, **74**, 611–625.
- 45 M. K. Whittlesey and E. Peris, *ACS Catal.*, 2014, **4**, 3152–3159.
- 46 H. Huang and T. H. Lambert, *Angew. Chem., Int. Ed.*, 2020, **59**, 658–662.
- 47 H. Dang, A. M. Whittaker and G. Lalic, *Chem. Sci.*, 2016, **7**, 505–509.
- 48 H. Lund, I. Fischmeister, E. Stenhagen, G. Andersson and H. Palmstierna, *Acta Chem. Scand.*, 1959, **13**, 192–194.
- 49 A. I. Cohen, B. T. Keeler, N. H. Coy and H. L. Yale, *Anal. Chem.*, 1962, **34**, 216–219.
- 50 J. P. Coleman, H. G. Gilde, J. H. P. Utley and B. C. L. Weedon, *J. Chem. Soc., Chem. Commun.*, 1970, **12**, 738–739.
- 51 J. P. Coleman, Naser-ud-din, H. G. Gilde, J. H. P. Utley, B. C. L. Weedon and L. Ebersson, *J. Chem. Soc., Perkin Trans. 2*, 1973, **14**, 1903–1908.
- 52 H. Lund, N. J. Jensen, S.-O. Almqvist, C. R. Enzell, A. Taticchi and B. Mannervik, *Acta Chem. Scand.*, 1974, **28b**, 263–265.
- 53 C. Combellas, F. Kanoufi and A. Thiébault, *J. Electroanal. Chem.*, 1996, **407**, 195–202.
- 54 C. P. Andrieux, C. Combellas, F. Kanoufi, J.-M. Savéant and A. Thiébault, *J. Am. Chem. Soc.*, 1997, **119**, 9527–9540.
- 55 P. Clavel, G. Lessene, C. Biran, M. Bordeau, N. Roques, S. Trevin and D. de Montauzon, *J. Fluor. Chem.*, 2001, **107**, 301–310.
- 56 P. Clavel, M. P. Léger-Lambert, C. Biran, F. Serein-Spirau, M. Bordeau, N. Roques and H. Marzouk, *Synthesis*, 1999, 829–834.
- 57 C. Saboureaux, M. Troupel, S. Sibille and J. Perichon, *J. Chem. Soc., Chem. Commun.*, 1989, **16**, 1138–1139.
- 58 E. Liotier, G. Mousset and C. Mousty, *Can. J. Chem.*, 1995, **73**, 1488–1496.
- 59 Y. Yamauchi, K. Sakai, T. Fukuhara, S. Hara and H. Senboku, *Synthesis*, 2009, 3375–3337.
- 60 B. B. Huang, L. Guo and W. J. Xia, *Green Chem.*, 2021, **23**, 2095–2103.
- 61 W. B. Wu, M. L. Li and J. M. Huang, *Tetrahedron Lett.*, 2015, **56**, 1520–1523.
- 62 J. Sheng, N. Wu, X. Liu, F. Liu, S. Liu, W. Ding, C. Liu and X. Cheng, *Chin. J. Org. Chem.*, 2020, **40**, 3873–3880.
- 63 X. T. Gao, Z. Zhang, X. Wang, J. S. Tian, S. L. Xie, F. Zhou and J. Zhou, *Chem. Sci.*, 2020, **11**, 10414–10420.
- 64 Y. Liu, X. Tao, Y. Mao, X. Yuan, J. Qiu, L. Kong, S. Ni, K. Guo, Y. Wang and Y. Pan, *Nat. Commun.*, 2021, **12**, 6745–6752.
- 65 A. Claraz, C. Allain and G. Masson, *Chem. – Eur. J.*, 2022, **28**, e202103337.
- 66 H. Zhang, M. Liang, X. Zhang, M.-K. He, C. Yang, L. Guo and W. Xia, *Org. Chem. Front.*, 2022, **9**, 95–101.



- 67 X. Yan, S. Wang, Z. Liu, Y. Luo, P. Wang, W. Shi, X. Qi, Z. Huang and A. Lei, *Sci. China: Chem.*, 2022, **65**, 762–770.
- 68 W. Chen, S. Ni, Y. Wang and Y. Pan, *Org. Lett.*, 2022, **24**, 3647–3651.
- 69 P. J. Elving and J. T. Leone, *J. Am. Chem. Soc.*, 1957, **79**, 1546–1550.
- 70 J. H. Stocker and R. M. Jenevein, *Chem. Commun.*, 1968, **16**, 934–935.
- 71 J.-S. Yang, K.-T. Liu and Y. O. Su, *J. Phys. Org. Chem.*, 1990, **3**, 723–731.
- 72 J. Kopilov and D. H. Evans, *J. Electroanal. Chem. Interfacial Electrochem.*, 1990, **280**, 435–438.
- 73 K. Uneyama, K. Maeda, T. Kato and T. Katagiri, *Tetrahedron Lett.*, 1998, **39**, 3741–3744.
- 74 K. Uneyama and T. Kato, *Tetrahedron Lett.*, 1998, **39**, 587–590.
- 75 K. Uneyama, G. Mizutani, K. Maeda and T. Kato, *J. Org. Chem.*, 1999, **64**, 6717–6723.
- 76 A. A. Stepanov, T. V. Minyaeva and B. I. Martynov, *Tetrahedron Lett.*, 1999, **40**, 2203–2204.
- 77 J. R. Box, A. P. Atkins and A. J. J. Lennox, *Chem. Sci.*, 2021, **12**, 10252–10258.
- 78 (a) J. Niu, H. Lin, J. Xu, H. Wu and Y. Li, *Environ. Sci. Technol.*, 2012, **46**, 10191–10198; (b) H. Lin, J. Niu, J. Xu, H. Huang, D. Li, Z. Yue and C. Feng, *Environ. Sci. Technol.*, 2013, **47**, 13039–13046; (c) Y. Su, U. Rao, C. M. Khor, M. G. Jensen, L. M. Teesch, B. M. Wong, D. M. Cwiertny and D. Jassby, *ACS Appl. Mater. Interfaces*, 2019, **11**, 33913–33922; (d) E. Mousset, N. Oturan and M. A. Oturan, *Appl. Catal., B*, 2018, **226**, 135–146; (e) Y. Su, U. Rao, C. M. Khor, M. G. Jensen, L. M. Teesch, B. M. Wong, D. M. Cwiertny and D. Jassby, *ACS Appl. Mater. Interfaces*, 2019, **11**, 33913–33922; (f) J. X. Zhu, Y. H. Chen, Y. R. Gu, H. Ma, M. Y. Hu, X. L. Gao and T. Z. Liu, *J. Hazard. Mater.*, 2022, **422**, 126953–126962.

