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A practical and sustainable two-component Minisci alkylation *via* photo-induced EDA-complex activation†

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An operationally simple, open-air, and efficient light-mediated Minisci C–H alkylation method is described, based on the formation of an electron donor–acceptor (EDA) complex between nitrogen-containing heterocycles and redox-active esters. In contrast to previously reported protocols, this method does not require a photocatalyst, an external single electron transfer agent, or an oxidant additive. Achieved under mildly acidic and open-air conditions, the reaction incorporates primary-, secondary-, and tertiary radicals, including bicyclo[1.1.1]pentyl (BCP) radicals, along with various heterocycles to generate Minisci alkylation products in moderate to good yields. Additionally, the method is exploited to generate a stereo-enriched, heteroaryl-substituted carbohydrate.

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Introduction

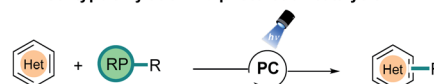
The extensive occurrence of nitrogen-containing heterocycles in biologically active natural products and pharmaceutical drugs makes them an attractive target in the development of new synthetic methods. In particular, reactions that can directly functionalize such heterocyclic substructures to generate medicinally valuable structural analogues are highly desirable.^{1–4}

One way to access these heterocyclic building blocks is to perform a C(sp²)–C(sp³) bond construction *via* an alkyl radical addition to heteroarenes.^{5–9} In this context, the venerable Minisci alkylation reaction developed by F. Minisci in 1971 has proven to be a powerful synthetic strategy for the C–H functionalization of a diverse range of heteroarenes.^{10,11} Since its discovery, extensive follow-up protocols have been identified by synthetic chemists to improve the applicability and scope of the reaction; however, the synthetic utility of this reaction has often been limited because of the use of stoichiometric, strong oxidants, highly acidic conditions, and/or high temperatures.^{12–21}

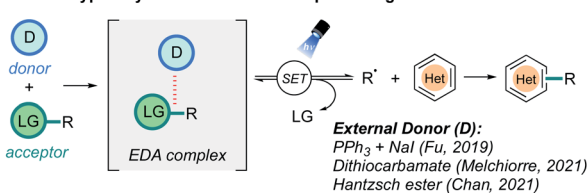
In recent years, the emergence of photoredox catalysis has transformed the Minisci reaction by offering milder conditions,

concomitantly circumventing the obligatory requirement of a strong oxidant to generate radicals under harsh conditions, thus making this process amenable to a wide range of heterocycles and radical precursors (Fig. 1a).^{22–25} However, despite contemporary advances in this field, some of the existing

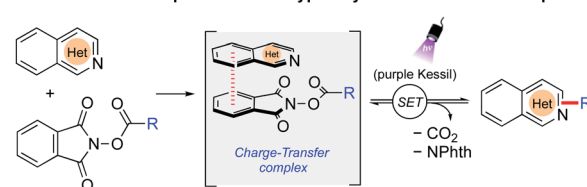
A. Minisci-type alkylation *via* photoredox catalysis



B. Minisci-type alkylation *via* an EDA complex using an external electron donor



C. This work: Two-component Minisci-type alkylation *via* self EDA complexation



- ✓ Photocatalyst free/EDA reaction
- ✓ No external donor
- ✓ No terminal oxidant
- ✓ Mildly acidic conditions (NaHSO₄)
- ✓ Open air/No precaution
- ✓ Atom-economical
- ✓ Compatible with 1°, 2°, 3°-alkyl radicals and various heterocycles (28 examples)

Fig. 1 Strategies for photoinduced Minisci alkylations. (A) *Via* photoredox catalysis (B) *via* EDA-activation using an external donor. (C) Developed two-component Minisci-alkylation *via* an EDA method.

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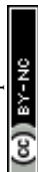
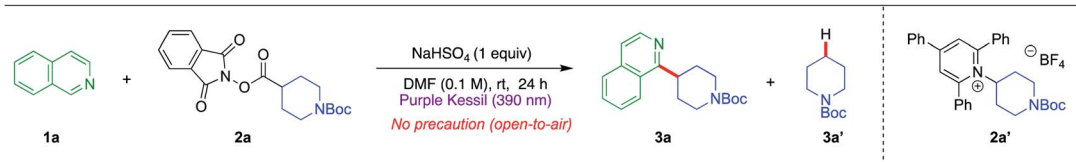


Table 1 Optimization of reaction conditions^a

Entry	Deviation from standard condition	%Yield ^b
1	None	62(58)
2	Pyridinium salt 2a' instead of RAE 2a	0
3	MeCN instead of DMF	15
4	DMA instead of DMF	46
5	THF instead of DMF	20
6	CH ₂ Cl ₂ instead of DMF	0
7	0.5 equiv. NaHSO ₄	49
8	No NaHSO ₄	24
9	Blue Kessil instead of purple	55
10	Argon condition	65
11	Dark	0

^a Reaction conditions: isoquinoline **1a** (0.1 mmol, 1 equiv.), RAE **2a** (0.15 mmol, 1.5 equiv.), NaHSO₄ (0.1 mmol, 1 equiv.) in DMF (1.0 mL, 0.1 M), 24 h irradiation with purple LEDs (390 nm). ^b Yields were determined by ¹H NMR analysis using trimethoxybenzene as internal standard.

methods still require strong acids for the activation of heterocycles. In addition, a stoichiometric strong oxidant is often needed in conjunction with the photocatalyst, particularly when the reaction is not net-redox neutral. External electron transfer agents (*e.g.*, Hantzsch ester) have also been required in some instances, diminishing the convenience and sustainability of the methods developed. Thus, further improvements are desirable to make Minisci C–H alkylation reactions more tolerant of sensitive functional groups, more convenient, and more efficient.

As an alternative, the photoinduced electron donor–acceptor (EDA) complex strategy is gaining considerable attention because of its ability to manifest photocatalyst-free, oxidant-free reaction conditions.^{26–30} The radicals are generated *via* a synergistic non-covalent interaction between an electron-rich donor and an electron-deficient acceptor. Although there are already many transformations reported based on this paradigm to construct new C–C, C–N, C–S, and C–B bonds efficiently, only a handful of reports are available in the realm of Minisci reactions (Fig. 1b). In all of these reported to date, an external single electron transfer agent is required. Thus, in 2019, Fu and coworkers disclosed an enantioselective variant of the Minisci alkylation by identifying a unique combination of catalytic sodium iodide and triphenylphosphine (NaI/PPh₃) as an effective electron donor to participate in an EDA-complex formation with redox-active esters (RAEs).³¹ Another report by Melchiorre *et al.* (2021) demonstrated that a catalytic dithiocarbamate anion possesses a similar capability as an electron donor toward the same acceptor, a redox-active ester.³² During the course of this investigation, Chan *et al.* reported that the commercially available Hantzsch ester, an established organo-reductant in EDA chemistry, could be used in the presence of trifluoroacetic

acid to perform Minisci alkylations on heterocycles by generating alkyl radicals under photocatalyst-free conditions.³³

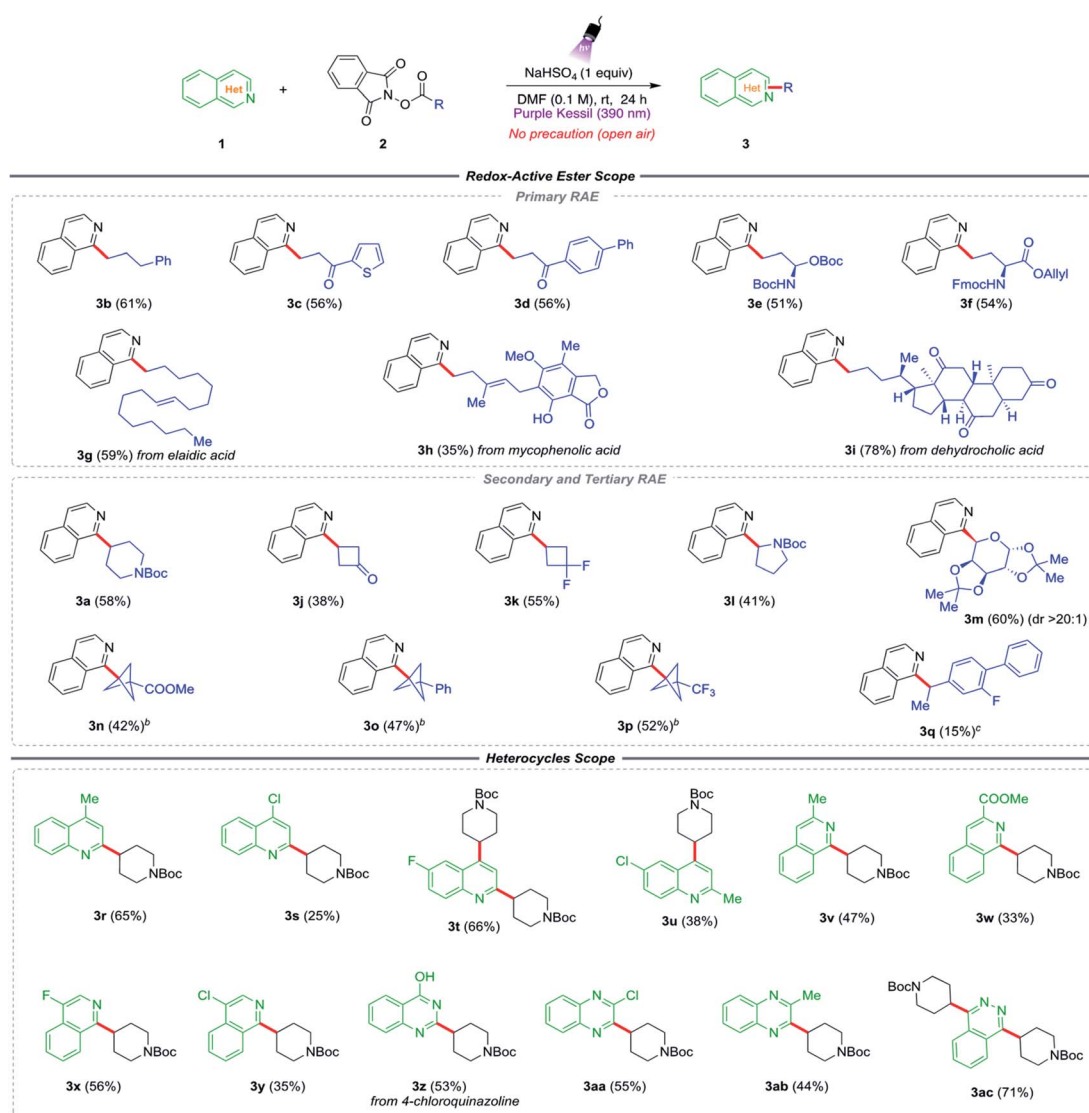
Discussion

In an effort to make the Minisci alkylation reactions operationally simpler and applicable in the presence of acid-sensitive groups, we envisioned exploring a two-component Minisci reaction (with no photocatalyst and no exogenous electron transfer agents) by investigating the feasibility of having nitrogen-containing heterocycles such as isoquinoline **1a** themselves serve as electron donors in a charge-transfer complex with suitable electron-accepting partners, using a very mild acid, NaHSO₄, as an activating agent (Fig. 1c). Both redox-active ester **2a** and pyridinium salt **2a'** were probed in optimization studies, owing to their pervasive utilization as effective electron-acceptor partners in EDA chemistry (Table 1).^{31–38} Gratifyingly, treatment of isoquinoline **1a** (1 equiv.) with redox-active ester **2a** (1.5 equiv.) generated the desired alkylation product **3a** in good yield (62% NMR, 58% isolated) after 24 h of purple light irradiation in DMF solvent (0.1 M) in the presence of NaHSO₄ (1 equiv.) under open-to-air conditions (entry 1). On the other hand, the reaction with pyridinium salt **2a'** under the same conditions failed to produce any desired product (entry 2). Notably, in entry 1, the yield of the Minisci alkylation product **3a** is somewhat compromised because of the formation of the decarboxylative hydrogenation side product **3a'** (33% NMR yield). A quick examination of other reaction solvents such as MeCN, DMA, and THF resulted in lower yields of the alkylation product **3a** and a higher amount of the side product **3a'** (entries 3–5), while dichloromethane did not support the desired reaction (entry 6). Typically, there is a strict



requirement for a stoichiometric strong acid in Minisci alkylations, mainly because of the sluggishness in the addition of alkyl radicals possessing nucleophilic character to electrophilic heterocycles in their non-protonated form. However, under the conditions developed, the reaction occurs under relatively benign acidic conditions, permitting incorporation of some highly acid-sensitive functional groups. Thus, making a direct comparison with the most-utilized Brønsted acids in Minisci alkylation, *p*-toluenesulfonic acid ($pK_a = -2.8$) or trifluoroacetic acid ($pK_a = 0.23$), the current method presents much milder acidic conditions with the use of NaHSO_4 ($pK_a = 1.99$).³⁹ Although the process works best when NaHSO_4 is used in a stoichiometric (1.0 equiv.) quantity, it also produces a significant amount of product with catalytic NaHSO_4 (entry 7).⁴⁰

Surprisingly, we still observed the desired product formation, albeit in low yield, when the acid (NaHSO_4) was completely removed from the reaction conditions (entry 8). This might be attributed to the formation of an isoquinoline radical cation species after an electron transfer event between isoquinoline **1a** and redox-active ester **2a** during the EDA complex formation, which is sufficiently electrophilic to react with the nucleophilic radical species. We also investigated the effect of light sources on the reaction. Replacing the purple Kessil lamp ($\lambda = 390$ nm) by a blue Kessil lamp ($\lambda = 427$ nm) delivered the Minisci alkylation product in a slightly lower yield (entry 9), although longer reaction times with a blue Kessil lamp compensated for the low yields. Next, changing the reaction environment from an open-air to an inert (argon) atmosphere provided only

Table 2 Substrate scope^a

^a Reaction conditions: isoquinoline **1** (0.3 mmol, 1 equiv.), RAE **2** (0.45 mmol, 1.5 equiv.), NaHSO_4 (0.3 mmol, 1 equiv.) in DMF (3.0 mL, 0.1 M), 24 h irradiation with purple Kessil (390 nm). ^b Trifluoroacetic acid (TFA) was used instead of NaHSO_4 . ^c Yield was determined by ¹H NMR analysis using trimethoxybenzene as internal standard.



a slight improvement in the yield, suggesting the method to be tolerant of oxygen and moisture (entry 10). Finally, as expected, the exclusion of a light source from the reaction conditions resulted in no conversion of the starting materials to the products (entry 11).

With suitable conditions in hand, the generality of the two-component Minisci alkylation method was evaluated (Table 2). First, the scope of redox-active esters was examined by employing various primary-, secondary-, and tertiary radical precursors. The method works well with relatively unstable primary radicals embellished with various functional groups. Redox-active esters with phenyl-, thienyl-, and biphenyl ketones on the side chain generated the desired products in good yields (**3b–3d**). Notably, because of the mildly acidic conditions, the reaction efficiently tolerated a substrate containing an acid-sensitive amine protecting group (**3e**). Further, the primary radical scope was extended to the modification of some biologically active and medicinally relevant compounds such as substrates derived from *L*-glutamate (**3f**), long-chain fatty acid elaidic acid (**3g**), an immunosuppressive drug mycophenolic acid (**3h**), and a steroid, dehydrocholic acid (**3i**).

Next, we explored the viability of secondary- and tertiary redox-active esters under the reaction conditions. Secondary radical precursors such as *N*-Boc piperidine (**3a**), cyclobutanone (**3j**), and 1,1-difluorocyclobutane (**3k**) facilitated the alkylation reaction, with products being formed in moderate to good yields. Some heteroatom-containing secondary radicals were also examined, such as an α -amino alkyl radical from *N*-Boc pyrrolidine RAE (**3l**), which provided the desired product in modest yield. More importantly, the Minisci alkylation product (**3m**) from α -D-galactopyranose sugar was isolated with excellent diastereoselectivity (*dr* > 20 : 1, NMR analysis). This particular result points to another feature of the developed method, in that in some cases high diastereoselectivity can be achieved at radical reaction centers in carbohydrate substrates. Because of the importance of bicyclo[1.1.1]pentanes (BCPs) in drug development programs, there was an interest in exploring the feasibility of incorporating these substructures in the current protocol. Being identified as sp^3 -rich aryl bioisosteres, the incorporation of BCPs into drug candidates has increased considerably in recent years as they exhibit enhanced pharmacokinetic properties, including improved aqueous solubility and membrane permeability over that of their aryl congeners.⁴¹ Stimulated by this, BCP-redox active esters containing $-\text{COOMe}$, $-\text{Ph}$, and $-\text{CF}_3$ groups were examined, all of which provided the desired alkylated products (**3n–3p**) in reasonable yields. Lastly, the addition of the benzylic radical derived from a flurbiprofen precursor to the isoquinoline core under the standard conditions provided the expected product in a slightly diminished yield (**3q**).

Next, attention was focused on the scope of nitrogen-containing heterocycles. Some functionally elaborated quinolines were assessed in the presence of the *N*-Boc piperidine radical precursor (**2a**). Lepidine (**3r**) and 4-chloroquinolines (**3s**) led to good yields of the mono-alkylated Minisci products by substituting the most electrophilic 2-position of the quinoline. However, when both the 2- and 4-positions were unoccupied,

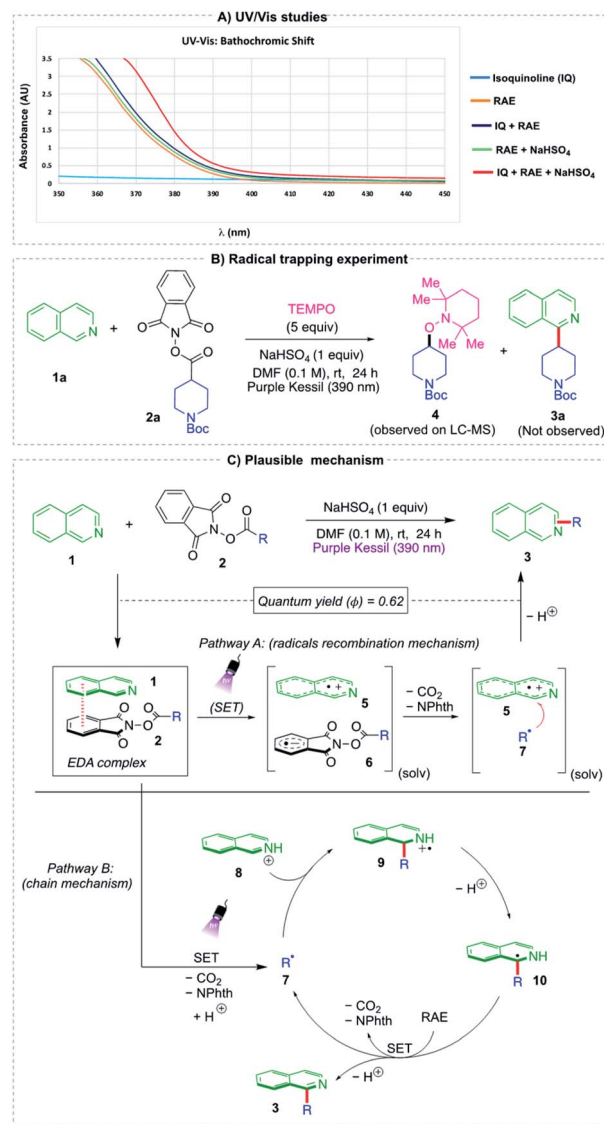


Fig. 2 (A) Chromatogram of UV-vis absorption study (B) TEMPO-radical trapping experiment (C) our proposed mechanisms.

such as in the case of 6-fluoroquinoline (**3t**), the reaction afforded disubstituted alkyl product. With 6-chloro-2-methylquinoline (**3u**), the alkylation occurred at the 4-position. Similarly, isoquinoline-based heterocycles with methyl-, ester-, and halide substitutions at the 3- and 4- positions also furnished the desired products in good yields (**3v–3y**). Some other medicinally important heterocycles, such as substituted quinazolines and quinoxalines, also proved to be equally effective under the established reaction conditions (**3z–3ac**).

To investigate the reaction mechanism of the two-component Minisci alkylation, ultraviolet/visible (UV/vis) and radical trapping experiments were performed (Fig. 2a). The UV/vis absorption spectrum of each individual reaction component was measured, along with that of the combined reaction mixture to provide evidence for the formation of an electron donor–acceptor (EDA) complex. When isoquinoline **1a** and redox-active ester **2a** were mixed along with NaHSO_4 in DMF



(0.1 M), the obtained absorption spectrum clearly depicted a red (bathochromic) shift, with a visible-light absorption tailing to the 410–420 nm region, indicating the formation of a charge-transfer molecular aggregate. To probe the generation of an alkyl radical species, a radical trapping agent, TEMPO (2,2,6,6-tetramethyl-1-piperidinyloxy), was added under standard conditions (Fig. 2b). This additive inhibited the alkylation reaction, with none of the Minisci alkylated product observed. Instead, TEMPO adduct **4** formed as confirmed *via* LC-MS analysis (see ESI†).

Based on these findings, two mechanisms appear reasonable (Fig. 2). Thus, considering pathway **A**, the heterocycle **1** and redox-active ester **2** form an EDA complex in the ground state. Upon photoirradiation of this molecular aggregate at 390 nm, a single-electron transfer (SET) event from **1** to **2** occurs, which leads to the formation of radical ion pair **5/6** in a solvent cage. Fragmentation of reduced RAE **6** generates phthalimide ion, CO₂, and alkyl radical. This newly generated radical **7** subsequently undergoes a radical recombination with the aromatic radical cation **5**,⁴² followed by deprotonation to afford the desired product **3**. Alternatively, a chain mechanism pathway **B** could also be operative, where the alkyl radical **7** escapes from the solvent cage and reacts with protonated *N*-heterocyclic intermediate **8** to yield an alkyl-substituted, nitrogen-centered radical cation **9**. The adduct formed subsequently undergoes deprotonation followed by oxidation *via* another molecule of redox-active ester **2** (ref. 24 and 33) to furnish the desired product **3** along with another alkyl radical species **7** to participate in the radical chain propagation step. Given that the experimentally determined quantum yield is high, but <1, ($\Phi = 0.62$, see ESI†), it seems likely that the two pathways coexist, with the radical ion pair recombination mechanism prevailing.

Conclusions

In summary, an operationally simple yet highly practical two-component Minisci alkylation method has been developed. This protocol avoids the requirement for expensive metal-based photocatalysts, strong oxidants, an external single electron transfer agent, and harsh acidic conditions, enabling the formation of medicinally relevant alkylated heterocycles from commercially available and/or readily synthesizable starting materials. Consistent with the spectroscopic studies and radical trapping experiment, a photoactivated EDA complexation between heterocycles and redox-active esters forms the basis of the reaction mechanism for the generation of alkyl radicals under open-air-conditions.

Data availability

Source/synthesis of *N*-(acyloxy)phthalimides and heterocycles used. General procedure for the two-component Minisci reactions. Description of mechanistic investigations. Quantum yield measurements. NMR spectra of all materials synthesized.

Author contributions

Lisa Marie Kammer, Matthias Krumb, Alexander Lipp, and Eugénie Romero carried out initial optimization studies, which were subsequently refined by Mohammed Sharique, Jadab Majhi, and Roshan K. Dhungana. All of the examples presented in the paper were carried out by Mohammed Sharique, Jadab Majhi, and Roshan K. Dhungana, with Mohammed Sharique leading the effort. Mohammed Sharique, Jadab Majhi, and Roshan K. Dhungana wrote the original draft of the manuscript with input from Gary A. Molander. All authors contributed to reaction development and discussion of results.

Conflicts of interest

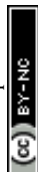
There are no conflicts to declare.

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Notes and references

- 1 J. A. Joule, in *Advances in Heterocyclic Chemistry*, ed. E. F. V. Scriven and C. A. Ramsden, Academic Press, 2016, vol. 119, pp. 81–106.
- 2 J. C. Lewis, R. G. Bergman and J. A. Ellman, *Acc. Chem. Res.*, 2008, **41**, 1013–1025.
- 3 J. Liu, J. Jiang, L. Zheng and Z.-Q. Liu, *Adv. Synth. Catal.*, 2020, **362**, 4876–4895.
- 4 J. Wencel-Delord and F. Glorius, *Nat. Chem.*, 2013, **5**, 369–375.
- 5 J.-R. Chen, X.-Q. Hu, L.-Q. Lu and W.-J. Xiao, *Acc. Chem. Res.*, 2016, **49**, 1911–1923.
- 6 F. O'Hara, D. G. Blackmond and P. S. Baran, *J. Am. Chem. Soc.*, 2013, **135**, 12122–12134.
- 7 J.-T. Yu and C. Pan, *Chem. Commun.*, 2016, **52**, 2220–2236.
- 8 Y. Tan, J. Wang, H.-Y. Zhang, Y. Zhang and J. Zhao, *Front. Chem.*, 2020, **8**, 582.
- 9 Y. Ji, T. Brueckl, R. D. Baxter, Y. Fujiwara, I. B. Seiple, S. Su, D. G. Blackmond and P. S. Baran, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 14411–14415.
- 10 F. Minisci, R. Bernardi, F. Bertini, R. Galli and M. Perchinummo, *Tetrahedron*, 1971, **27**, 3575–3579.
- 11 F. Minisci, F. Fontana and E. Vismara, *J. Heterocycl. Chem.*, 1990, **27**, 79–96.
- 12 M. A. J. Dunton, *MedChemComm*, 2011, **2**, 1135–1161.



- 13 R.-J. Tang, L. Kang and L. Yang, *Adv. Synth. Catal.*, 2015, **357**, 2055–2060.
- 14 D. R. Sutherland, M. Veguillas, C. L. Oates and A.-L. Lee, *Org. Lett.*, 2018, **20**, 6863–6867.
- 15 D. N. Mai and R. D. Baxter, *Org. Lett.*, 2016, **18**, 3738–3741.
- 16 A. P. Antonchick and L. Burgmann, *Angew. Chem., Int. Ed.*, 2013, **52**, 3267–3271.
- 17 I. B. Seiple, S. Su, R. A. Rodriguez, R. Gianatassio, Y. Fujiwara, A. L. Sobel and P. S. Baran, *J. Am. Chem. Soc.*, 2010, **132**, 13194–13196.
- 18 G. A. Molander, V. Colombel and V. A. Braz, *Org. Lett.*, 2011, **13**, 1852–1855.
- 19 M. Pisset, N. Fleury-Brégeot, D. Oehlich, F. Rombouts and G. A. Molander, *J. Org. Chem.*, 2013, **78**, 4615–4619.
- 20 Y. Fujiwara, J. A. Dixon, R. A. Rodriguez, R. D. Baxter, D. D. Dixon, M. R. Collins, D. G. Blackmond and P. S. Baran, *J. Am. Chem. Soc.*, 2012, **134**, 1494–1497.
- 21 Á. Gutiérrez-Bonet, C. Remeur, J. K. Matsui and G. A. Molander, *J. Am. Chem. Soc.*, 2017, **139**, 12251–12258.
- 22 R. S. J. Proctor and R. J. Phipps, *Angew. Chem., Int. Ed.*, 2019, **58**, 13666–13699.
- 23 R. S. J. Proctor, P. Chuentragool, A. C. Colgan and R. J. Phipps, *J. Am. Chem. Soc.*, 2021, **143**, 4928–4934.
- 24 L. M. Kammer, A. Rahman and T. Opatz, *Molecules*, 2018, **23**, 764–780.
- 25 M. Krumb, L. M. Kammer, S. O. Badir, M. J. Cabrera-Afonso, V. E. Wu, M. Huang, A. Csakai, L. A. Marcaurrelle and G. A. Molander, *Chem. Sci.*, 2022, **13**, 1023–1029.
- 26 Z. Yang, Y. Liu, K. Cao, X. Zhang, H. Jiang and J. Li, *Beilstein J. Org. Chem.*, 2021, **17**, 771–799.
- 27 Y.-q. Yuan, S. Majumder, M.-h. Yang and S.-r. Guo, *Tetrahedron Lett.*, 2020, **61**, 151506.
- 28 G. E. M. Crisenza, D. Mazzarella and P. Melchiorre, *J. Am. Chem. Soc.*, 2020, **142**, 5461–5476.
- 29 L. Zheng, L. Cai, K. Tao, Z. Xie, Y.-L. Lai and W. Guo, *Asian J. Org. Chem.*, 2021, **10**, 711–748.
- 30 C. G. S. Lima, T. de M. Lima, M. Duarte, I. D. Jurberg and M. W. Paixão, *ACS Catal.*, 2016, **6**, 1389–1407.
- 31 M.-C. Fu, R. Shang, B. Zhao, B. Wang and Y. Fu, *Science*, 2019, **363**, 1429–1434.
- 32 E. de Pedro Beato, D. Spinnato, W. Zhou and P. Melchiorre, *J. Am. Chem. Soc.*, 2021, **143**, 12304–12314.
- 33 J. Li, S. Siang Tan, S. H. Kyne and P. Wai Hong Chan, *Adv. Synth. Catal.*, 2022, **364**, 802–810.
- 34 V. C. Polites, S. O. Badir, S. Keess, A. Jolit and G. A. Molander, *Org. Lett.*, 2021, **23**, 4828–4833.
- 35 M. Elkhalfi, M. B. Elbaum, D. M. Chenoweth and G. A. Molander, *Org. Lett.*, 2021, **23**, 8219–8223.
- 36 L. M. Kammer, S. O. Badir, R.-M. Hu and G. A. Molander, *Chem. Sci.*, 2021, **12**, 5450–5457.
- 37 M. Yang, T. Cao, T. Xu and S. Liao, *Org. Lett.*, 2019, **21**, 8673–8678.
- 38 F. J. R. Klauck, M. J. James and F. Glorius, *Angew. Chem., Int. Ed.*, 2017, **56**, 12336–12339.
- 39 International Union of Pure and Applied Chemistry, E. P. Serjeant and B. Dempsey, *Ionisation constants of organic acids in aqueous solution*, Pergamon Press, Oxford, 1979.
- 40 W.-M. Cheng, R. Shang and Y. Fu, *ACS Catal.*, 2017, **7**, 907–911.
- 41 M. R. Bauer, P. Di Fruscia, S. C. C. Lucas, I. N. Michaelides, J. E. Nelson, R. I. Storer and B. C. Whitehurst, *RSC Med. Chem.*, 2021, **12**, 448–471.
- 42 S. R. Kandukuri, A. Bahamonde, I. Chatterjee, I. D. Jurberg, E. C. Escudero-Adan and P. Melchiorre, *Angew. Chem., Int. Ed.*, 2015, **54**, 1485–1489.

