



Cite this: *Green Chem.*, 2024, **26**, 10265

# General electron–donor–acceptor complex mediated thioesterification reaction *via* site-selective C–H functionalization using aryl sulfonium Salts†

Roshan I. Patel, Barakha Saxena and Anuj Sharma \*

Contemporary methods for synthesizing thioesters often necessitate expensive catalysts and harsh conditions, making their synthesis from chemical feedstocks challenging. Herein, we report a sustainable metal-, photocatalyst-, and oxidant-free electron donor–acceptor (EDA) mediated synthesis of thioesters *via* site-selective C–H functionalization using aryl sulfonium salts (acceptor) with potassium thioacid salts (donor) under visible light irradiation. Our approach enables rapid access to thioesters from a wide variety of arenes, including pharmaceutical and agrochemical compounds, as well as a diverse range of alkyl, aryl, and heteroaryl potassium thioacid salts with excellent efficiency and regioselectivity. Mechanistic studies supported the formation of an EDA-complex, and radical trapping experiments corroborated the involvement of a radical-based mechanism for the product formation. Moreover, our method demonstrates excellent atom economy and *E*-factor scores, which are considered excellent in terms of safety, economic and ecological yardsticks.

Received 30th July 2024,  
Accepted 2nd September 2024

DOI: 10.1039/d4gc03768e

[rsc.li/greenchem](https://rsc.li/greenchem)

## Introduction

Thioesters are an essential functional group in a wide range of complex natural products, polymers, and therapeutics.<sup>1</sup> They serve as high-energy intermediates in numerous biochemical processes, playing a critical role in cellular regulation and biosynthetic pathways such as metabolism, fatty acid synthesis, and the production of esters and polyketides in living organisms.<sup>2</sup> Due to their significant biological and pharmaceutical properties, exemplified in Fig. 1a,<sup>3</sup> thioesters are essential intermediates in organic synthesis. Consequently, the synthesis of thioesters has attracted considerable interest, prompting the development of various strategies for their preparation.<sup>4</sup> Contemporary methods for synthesizing thioesters have expanded to a wide range of starting materials, including alkenes, alkynes, amines, halides, alcohols, aldehydes, and phenol derivatives.<sup>5</sup> These methods often employ toxic carbon monoxide (CO) gas as a carbonyl group source and toxic and malodorous thiols, sulfonyl hydrazides, disulfides, or sulfonyl chlorides as S-group sources.<sup>5</sup> In addition, these approaches have other notable drawbacks, such as the

use of expensive transition metal catalysts, harsh oxidants, and high reaction temperatures.<sup>4,5</sup> In recent years, the synthesis of thioesters has focused on developing more efficient techniques that utilize carbonyl compounds such as aldehydes, acyl chlorides, ketoacids, and carboxylic acids or their activated forms with thiols, disulfides, sodium sulfinates, or elemental sulfur (S<sub>8</sub>) to construct C–S bonds.<sup>6</sup> Unfortunately, these methods also invariably require expensive transition metal catalysts under harsh reaction conditions, which result in reduced yields and necessitate extensive purification of the final products. In the past decade, visible light photoredox catalysis has become one of the most exciting fields in research among chemists, harnessing light energy for chemical transformations.<sup>7</sup> Given its extensive applications in organic synthesis, several light-induced thioesterification approaches have been developed.<sup>8</sup> For instance, Liao and co-workers (2019) demonstrated a visible light-induced base-mediated deaminative thioesterification of amino-acid-derived Katritzky salts (acceptor) using a single example of non-substituted thiobenzoic acid (donor), thus seriously limiting substrate scope of the method (Fig. 1b).<sup>9</sup> Additionally, the reaction produced 2,4,6-triphenylpyridine as a non-reusable by-product. Subsequently, the same group published a dual Ru/Cu-catalyzed decarboxylative thioesterification of carboxylic acid-derived *N*-(acetoxy) phthalimides with again a single thiobenzoic acid as a sulfur source under visible light irradiation.<sup>10</sup> In addition to the limited substrate scope, the method suffered in terms of

Green Organic Synthesis laboratory, Indian Institute of Technology Roorkee, Roorkee, 247667, India. E-mail: [anujsharma.mcl@gmail.com](mailto:anujsharma.mcl@gmail.com), [anuj.sharma@cy.iitr.ac.in](mailto:anuj.sharma@cy.iitr.ac.in)

†Electronic supplementary information (ESI) available: Detailed optimization, experimental procedure, <sup>1</sup>H, <sup>13</sup>C, and <sup>19</sup>F NMR spectra of all the synthesized compounds. See DOI: <https://doi.org/10.1039/d4gc03768e>

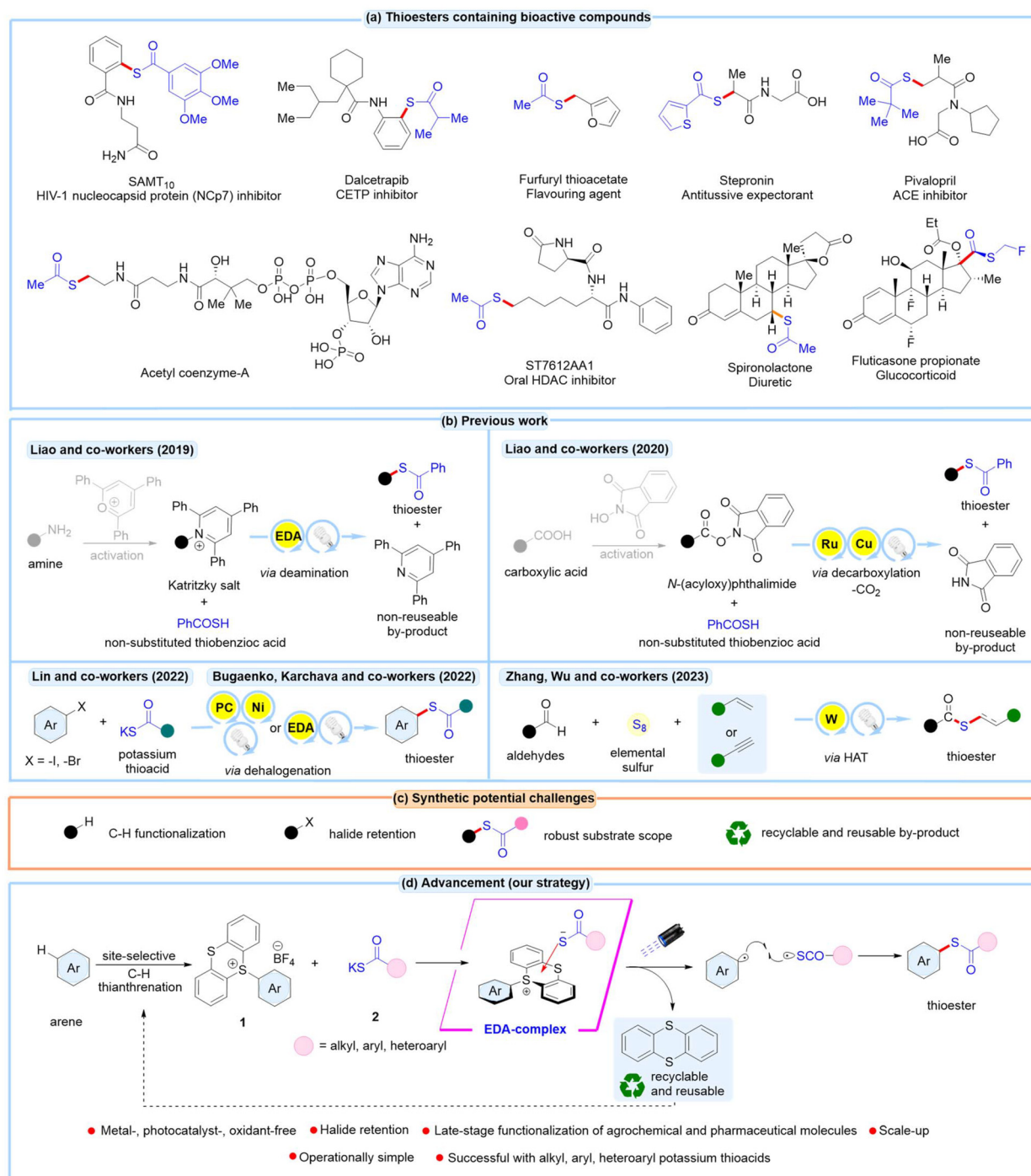


Fig. 1 (a) Thioesters containing bioactive compounds; (b) previous work; (c) synthetic potential challenges; (d) advancement (our strategy).

expensive transition metal catalysts. In 2022, independent research groups revealed a visible light dual 4-CzIPN/Ni-catalyzed or photoinduced dehalogenative thioesterification of aryl halides ( $X = -\text{Br}, -\text{I}$ ) with potassium thioacid salts.<sup>11</sup> However, these methods are limited to simple aryl halides ( $X = -\text{Br}, -\text{I}$ ) with a low functional group tolerance and have yet to be tested on more complex molecules, thereby restricting its substrate scope. Additionally, these methods require long reaction times

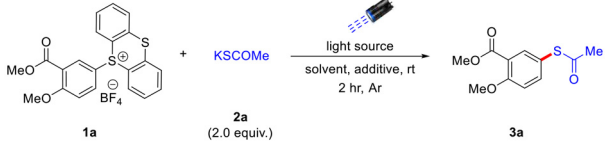
and expensive photocatalysts under light, making them less favorable. Later, Zhang, Wu, and co-workers, in 2023, made a significant advancement in the synthesis of thioesters by developing a *tetra-*n**-butylammonium decatungstate (TBADT)-catalyzed thioesterification of aldehydes with elemental sulfur ( $\text{S}_8$ ) and alkenes or alkynes *via* hydrogen atom-transfer (HAT).<sup>12</sup> The method, however, suffers from the use of toxic elemental sulfur ( $\text{S}_8$ ) and an expensive transition metal catalyst.

Around the same time, Liu, Zhao, and co-workers reported a visible light palladium-catalyzed decarboxylative thiocarbonylation of sulfonium salts with oxalic acid mono-thioethers (OAMs).<sup>13</sup> However, the method required an expensive metal catalyst with a complex ligand system for thioester synthesis. In general, a metal-, photocatalyst-, oxidant-, and base-free methods can be a significant addition to thioesterification reactions.<sup>6,8</sup> Moreover, the synthesis of thioesters *via* C–H functionalization is challenging and highly desirable, and achieving this would represent a substantial advancement in thioesterification reactions.<sup>5,6,8</sup> In this context, site-selective C–H functionalization *via* thianthrenium salt synthesis has emerged as a powerful tool in organic synthesis, enabling the introduction of various functional groups into the aromatic compound and providing a streamlined and efficient pathway toward complex molecules.<sup>14</sup> Concurrently, EDA-complex photochemistry has garnered significant interest due to its unique capability to harness visible light to activate colorless substances, generate radical intermediates, and drive subsequent reactions without the need for external catalysts.<sup>15</sup> In line with our ongoing research into photoinduced EDA-complex reactions,<sup>16</sup> herein, we present a practical method for synthesizing thioesters. We hypothesize the formation of an EDA-complex between aryl thianthrenium salts (acceptor) and potassium thioacid salts (donor) to synthesize valuable thioester products (Fig. 1d). Under visible light irradiation, potassium thioacid salt (donor) engage in a single electron transfer (SET) process with aryl thianthrenium salts (acceptor), generating an aryl radical intermediate and recyclable thianthrene. This aryl intermediate subsequently interacts with the resulting sulfur-centered radical to give the desired thioester product. By leveraging the EDA-complexes in our reaction, this method enables regioselective C–H thioesterification of arenes, a challenging feat to achieve without transition metal catalysts, aryl halides as radical precursors, and toxic, foul-smelling sulfur reagents. Additionally, the thianthrene by-product generated during the reaction is reusable and recyclable, offering further benefits in terms of sustainability and efficiency. Moreover, our method facilitates coupling complex agrochemical and pharmaceutical compounds with thioacids, offering substantial benefits to academic research and industrial applications.

## Results and discussion

Our study commenced with the use of methyl-2-methoxybenzoate-derived thianthrenium tetrafluoroborate **1a** and potassium thioacetate **2a** as representative substrates in DMSO solvent under 390 nm Kessil lamp irradiation (Table 1). We were delighted to observe the desired product obtained in 69% yield (entry 1). DMSO efficiently solubilized the reaction components (**1a** and **2a**), resulting in an intense yellow color change in the reaction mixture, which may be speculated to be an EDA-complex. Screening other solvents such as MeCN, DCM, THF, DMA, DMF, and EtOAc resulted in lower yields

**Table 1** Reaction conditions: a mixture of **1a** (0.1 mmol), and **2a** (0.2 mmol, 2.0 equiv.), additive (2 equiv.), solvent (0.8 mL) under visible light irradiation (light source) for 2 h under Ar atmosphere at rt



| Entry | Solvent | Additive                        | Light (nm) | Yield (%)              |
|-------|---------|---------------------------------|------------|------------------------|
| 1     | DMSO    | —                               | 390 nm     | 69%                    |
| 2     | MeCN    | —                               | 390 nm     | 58%                    |
| 3     | DCM     | —                               | 390 nm     | 37%                    |
| 4     | THF     | —                               | 390 nm     | 25%                    |
| 5     | DMA     | —                               | 390 nm     | 48%                    |
| 6     | DMF     | —                               | 390 nm     | 43%                    |
| 7     | EtOAc   | —                               | 390 nm     | 54%                    |
| 8     | DMSO    | SiO <sub>2</sub> (20 mg)        | 390 nm     | 72%                    |
| 9     | DMSO    | 4 Å MS (20 mg)                  | 390 nm     | 77%                    |
| 10    | DMSO    | 3 Å MS (20 mg)                  | 390 nm     | 70%                    |
| 11    | DMSO    | Basic alumina (20 mg)           | 390 nm     | 73%                    |
| 12    | DMSO    | K <sub>2</sub> CO <sub>3</sub>  | 390 nm     | 12%                    |
| 13    | DMSO    | Na <sub>2</sub> CO <sub>3</sub> | 390 nm     | 17%                    |
| 14    | DMSO    | K <sub>2</sub> HPO <sub>4</sub> | 390 nm     | 15%                    |
| 15    | DMSO    | Cs <sub>2</sub> CO <sub>3</sub> | 390 nm     | 21%                    |
| 16    | DMSO    | 4 Å MS (20 mg)                  | 427 nm     | 85% (83%) <sup>a</sup> |
| 17    | DMSO    | 4 Å MS (20 mg)                  | 440 nm     | 62%                    |
| 18    | DMSO    | 4 Å MS (20 mg)                  | 456 nm     | 50%                    |
| 19    | DMSO    | 4 Å MS (20 mg)                  | Dark       | Nr                     |
| 20    | DMSO    | 4 Å MS (20 mg)                  | 427 nm     | 72% <sup>b</sup>       |
| 21    | DMSO    | 4 Å MS (60 mg)                  | 427 nm     | 83% <sup>c</sup>       |

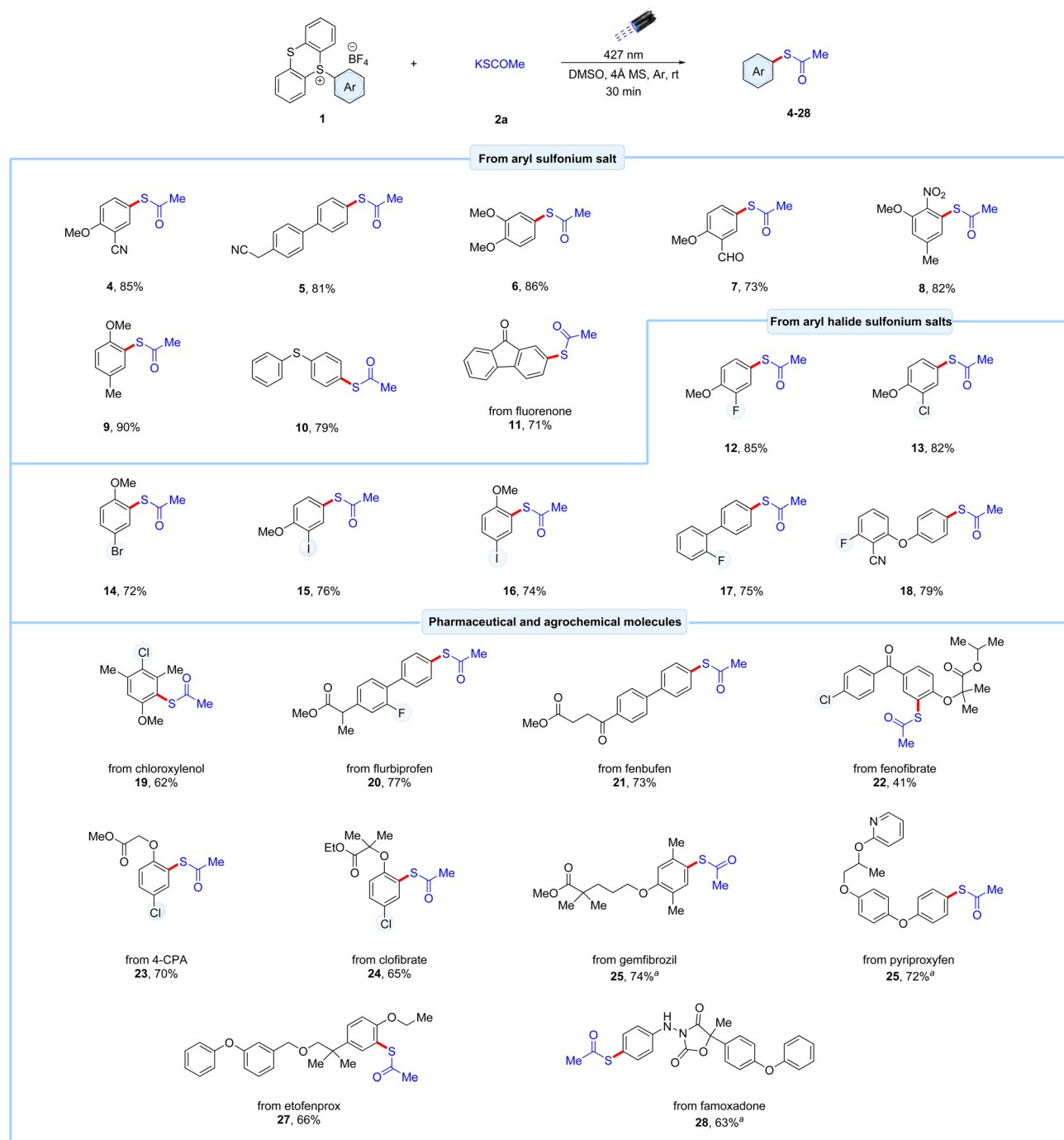
<sup>a</sup> **1a** (0.3 mmol), and **2a** (0.6 mmol, 2.0 equiv.), 4 Å MS (60 mg), dry DMSO (1.5 mL). <sup>b</sup> Open-to-air. <sup>c</sup> **1a** (0.3 mmol), and **2a** (0.6 mmol, 2.0 equiv.), dry DMSO (1.5 mL), 4 Å MS (60 mg), 30 min; nr = no reaction.

(entries 2–7). We then investigated the effect of different additives (entries 8–15). To our delight, using 4 Å MS as an additive slightly increased the yield to 77% (entry 9). We speculate that 4 Å MS functions as an ion-trap reagent, which captures the potassium ion of thioacetate **2a**, which may result in the formation of a stable thiolate anion.<sup>17</sup> This can then give rise to a more efficient interaction between the electron acceptor **1a** and the electron donor **2a**. Notably, using an inorganic base such as K<sub>2</sub>CO<sub>3</sub> resulted in reduced product yield (entry 12). Screening of other inorganic bases as additives, such as Na<sub>2</sub>CO<sub>3</sub>, K<sub>2</sub>HPO<sub>4</sub>, and Cs<sub>2</sub>CO<sub>3</sub> did not improve the reaction yield either (entries 13–15). We next explored using different light sources (entries 16–18). Remarkably, under 427 nm, visible light irradiation led to an 85% yield of the required product (entry 16). No product formation was observed when the reaction was conducted in the dark, underscoring the necessity of light irradiation (entry 19). Moreover, performing the reaction in an open-to-air condition resulted in a lower yield of 72%, indicating the importance of an inert atmosphere for optimal product formation (entry 20). Furthermore, the optimized time of 30 minutes resulted in no additional change in yield (entry 21).

With the optimized reaction conditions established, various aromatic feedstock substrates were subjected to the C–H

thioesterification using potassium thioacetate **2a** as the coupling partner in the photochemical reaction. Aryl thianthrenium salts bearing electron-donating as well as electron-withdrawing substituents such as cyano-, trifluoromethyl-, nitro-, methoxy-, and methyl groups, reacted well under the optimized conditions, yielding the desired thioester products **4–9** in 73–90% yields (Scheme 1). Additionally, diphenyl sulfide-derived thianthrenium salt reacted efficiently under this proto-

col, producing the thioacetate product **10** in 79% yield. The bioactive fluorenone-derived thioacetate **11** was also obtained with a 71% yield. The retention of halides is problematic under reported transition metal and light-induced reactions. To evaluate the compatibility of this method, a series of aryl halide-derived thianthrenium salts were subjected to the EDA protocol. Pleasingly, the protocol effectively retained the halide groups to give the corresponding thioester products **12–18**.



**Scheme 1** Reaction conditions: aryl sulfonium salt **1** (0.3 mmol), potassium thioacetate **2a** (2 equiv., 0.6 mmol), 4 Å MS (60 mg), in 1.5 mL DMSO, were irradiated with 427 nm Kessil Lamp (40 W) at room temperature under argon for 30 min. <sup>a</sup> 2 h.

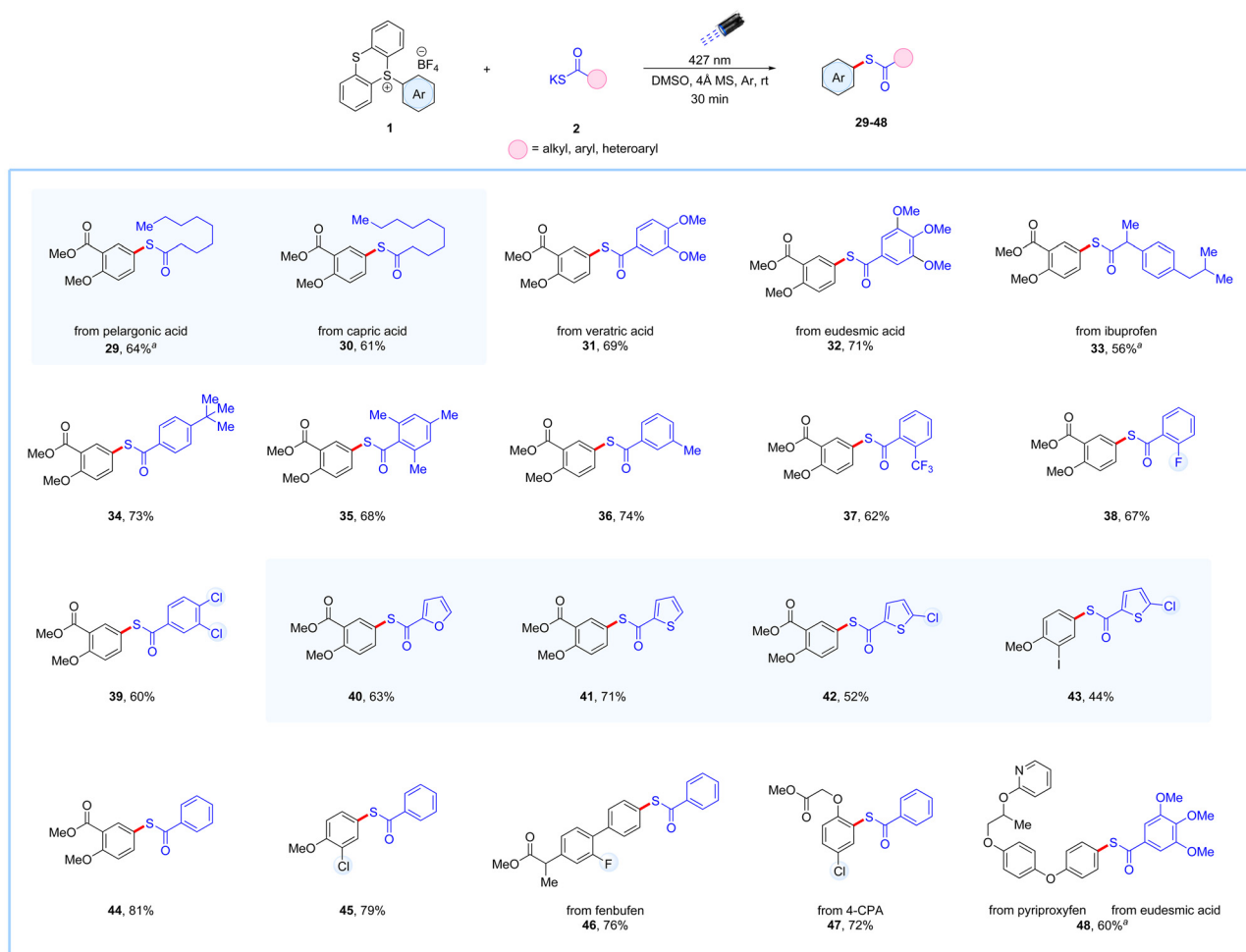
Notably, aryl thianthrenium salts with fluoro- and chloro-group reacted exceptionally well (**12–13**), giving higher yields than their bromo- and iodo-substituted counterparts (**14–16**). Additionally, *meta*-substituted aryl iodo-substituted thianthrenium salts reacted smoothly under this method, achieving a yield of 74% (**16**). This protocol was also extended to diaryl-fluoro-substituted thianthrenium salts, resulting in the isolation of their corresponding thioester products **17** and **18** in good yields (75 and 79% yield, respectively). With the excellent site-selectivity achieved through the synthesis of thianthrenium salts,<sup>14a</sup> combined with the effectiveness of our EDA-complex protocol, makes this method a powerful tool for executing precise late-stage thioesterification of complexed agrochemical and pharmaceutical compounds, as illustrated in the Scheme 1. The thioesterification of thianthrenium salt derived from the polysubstituted antiseptic and disinfectant agent chloroxyleneol proceeded smoothly, yielding the desired thioacetate product **19** in 62% yield. Moreover, using this protocol, aryl ester bearing thianthrenium salts derived from the anti-inflammatory drugs flurbiprofen and fenbufen underwent successful thioesterification, yielding the desired products **20** and **21** in 77% and 73% yield, respectively. Next, a series of aryl ether-substituted agrochemical and pharmaceutical-derived thianthrenium salts were subjected to the optimized conditions. These included thianthrenium salts derived from the hypertriglyceridemia drugs fenofibrate and gemfibrozil, the plant growth regulator *p*-chlorophenoxyacetic acid (4-CPA), antilipidemic drug clofibrate, the insecticidal agents pyriproxyfen, and etofenprox, as well as the amide, ester and ether-bearing fungicidal agent famoxadone. These reactions were well tolerated, affording the corresponding thioester products as single regioisomers in satisfactory to good yields (**22–28**). Notably, the hypertriglyceridemia drug fenofibrate **22**, which contains a highly bulky ether group on the corresponding carbon, resulted in a lower yield of 41%, likely due to steric hindrance. In general, the protocol exhibited excellent compatibility with numerous functional groups, including halides (F and Cl), esters, ethers, amides, and heteroarenes.

To further diversify our EDA-complex strategy, we employed various potassium thioacid salts with alkyl, aryl, and hetero-aryl functionalities with the aryl thianthrenium salt. We were delighted that various potassium thioacid salts reacted smoothly with the aryl thianthrenium salts to yield the corresponding thioester products in moderate to good yields, as illustrated in Scheme 2. Potassium thioacid salts derived from naturally occurring long-chain saturated fatty acids such as pelargonic and capric acid, performed well under the established conditions, yielding the corresponding thioester products **29** and **30** with 64% and 61% yield, respectively. Additionally, potassium thioacid salts derived from the anti-proliferative agent veratric acid and antioxidant agent eudesmic acid salts proved to be efficient electron donors in this photochemical strategy, producing the desired thioester products **31** and **32** in 69% and 71% yield, respectively. The potassium thioacid derived from the anti-inflammatory drug ibuprofen was also compatible under this protocol, yielding the

thioester product **33** in 56% yield. Aryl potassium thioacid salts bearing alkyl functionalities were found to be suitable coupling partners with aryl thianthrenium salts, yielding the corresponding thioester products **34–36** in yields ranging from 68–74% yield. Furthermore, aryl potassium thioacids bearing trifluoromethyl- and fluoro-, and dichloro-groups were also well tolerated, yielding the required thioester product **37**, **38**, and **39** with yields of 62%, 67%, and 60% yield, respectively. Additionally, potassium thioacids derived from various O- and S-containing heterocycles were also effective in this photochemical process, yielding the desired thioester products **40–43** in satisfactory to good yields. Moreover, non-substituted potassium benzothioate proved to be a suitable coupling partner under this protocol, efficiently reacting with different aryl thianthrenium salts to produce the required thioester products **44–47** in good yields. Lastly, the compatibility of this method was demonstrated in the thioesterification of the insecticidal agent pyriproxyfen with eudesmic acid-derived potassium thioacid salt, affording the required thioester product **48** in 60% yield.

### Mechanistic insights

Next, a series of mechanistic studies were conducted to elucidate the reaction mechanism as illustrated in Fig. 2. The UV/Vis-absorption analysis of individual components and the reaction mixture (**1a** + **2a**) in DMSO is shown in Fig. 2a. The DMSO solutions of methyl-2-methoxybenzoate-derived thianthrenium salt **1a** (red line), and potassium thioacetate **2a** (green line) displayed a small absorption band in the visible light region (>400 nm). Moreover, a clear bathochromic shift (blue band) was observed of the reaction mixture (**1a** + **2a**) in DMSO, which was visible by the intense yellow color of the reaction mixture, as shown in Fig. 2a. It indicates the formation of an electron donor–acceptor (EDA) aggregate (blue band). Additionally, a Job's plot using UV-visible absorption experiments was performed to determine the stoichiometry of the EDA-complex between **1a** and **2a**. The maximum absorption of the reaction mixture [**1a** + **2a**] at a 50% molar fraction indicated a 1 : 1 ratio of **1a** and **2a** (Fig. 2b). Moreover, <sup>1</sup>H NMR titrations were conducted to provide further evidence of an EDA-complex formation in DMSO-*d*<sub>6</sub> (Fig. 2c). The <sup>1</sup>H NMR signal of C1–H proton in methyl-2-methoxybenzoate-derived thianthrenium salt **1a**, shifted downfield along with the increasing amount of potassium thioacetate **2a**. In contrast, C4–H, C3–H, C2–H, and C5–H proton shifted upfield, thus indicating the formation of an EDA-complex between **1a** and **2a**.<sup>18</sup> Under standard conditions, the photochemical reaction was subjected to a radical trap experiment with TEMPO (2,2,6,6-tetramethyl-1-piperidinyloxy), butylated hydroxytoluene (BHT) and 1,1-diphenylethylene (Fig. 2d). The reactions were quenched, and the radical trapped adducts were analyzed and detected by high-resolution mass spectrometry (HRMS). These results support the generation of an aryl radical as well as thiyl radical intermediate in the photochemical transformation, likely initiated by the photoactive EDA-complex between methyl-2-methoxy



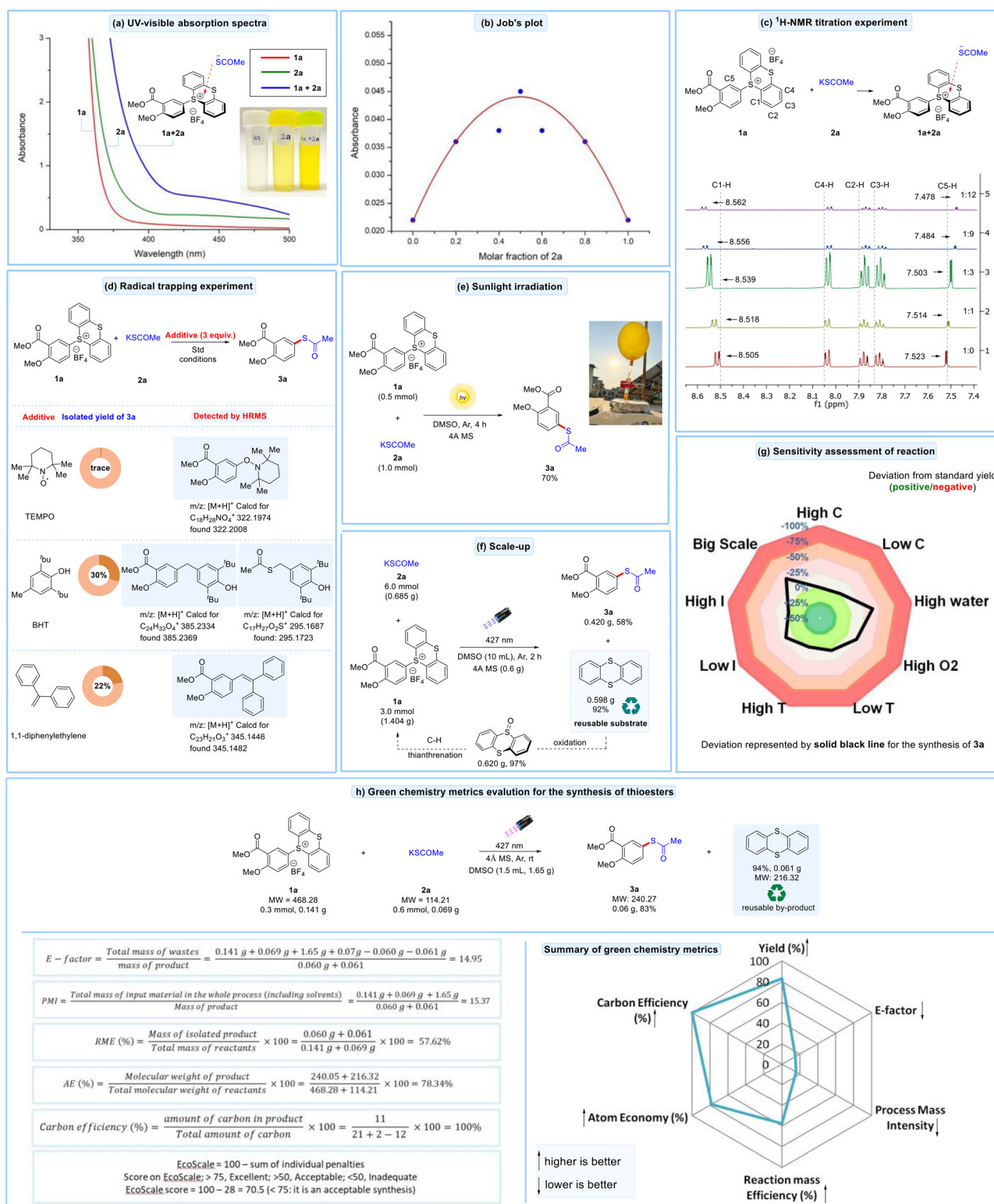
**Scheme 2** Reaction conditions: aryl sulfonium salt **1** (0.3 mmol), potassium thioacid salt **2** (2 equiv., 0.6 mmol), 4 Å MS (60 mg), in 1.5 mL DMSO, were irradiated with 427 nm Kessil Lamp (40 W) at room temperature under argon for 30 min. <sup>a</sup> 2 h.

benzoate-derived thianthrenium salt **1a** and potassium thioacetate **2a**.

A quantum yield experiment was also performed to determine a possible reaction pathway for this transformation. The quantum yield was measured as  $\Phi = 44$ , indicating a radical chain pathway (see ESI† for detailed discussion). Furthermore, the reaction also proceeded successfully under natural sunlight irradiation, yielding the targeted product **3a** in 70% yield (Fig. 2e). A gram-scale synthesis was conducted using methyl-2-methoxy benzoate-derived thianthrenium salt **1a** (1.404 g, 3 mmol) and potassium thioacetate **2a** (0.685 g, 6.0 mmol, 2 equiv.) (Fig. 2f). The reaction was carried out under visible light irradiation for 2 h, yielding the desired product **3a** in 58% yield (0.420 g). Moreover, 0.598 g (92% yield) of thianthrene was also recovered *via* chromatographic separation and was recycled back to thianthrene-*S*-oxide (0.620 g, 97%). This thianthrene-*S*-oxide was further converted back to **1a** (0.957 g, 77%) (see ESI† for detailed discussion). Next, the efficiency of our reaction protocol for the synthesis of **3a** was investigated under various sensitivity assessment parameters.<sup>19</sup> This EDA-complex

transformation was found sensitive to water, oxygen concentration, and low light intensity (Fig. 2g). However, the protocol showed good tolerance towards substrate concentration, temperature, and high light intensity variations. Further attempts were made to synthesize the thioester compound *via* one-pot, but the reaction failed to yield the required product.

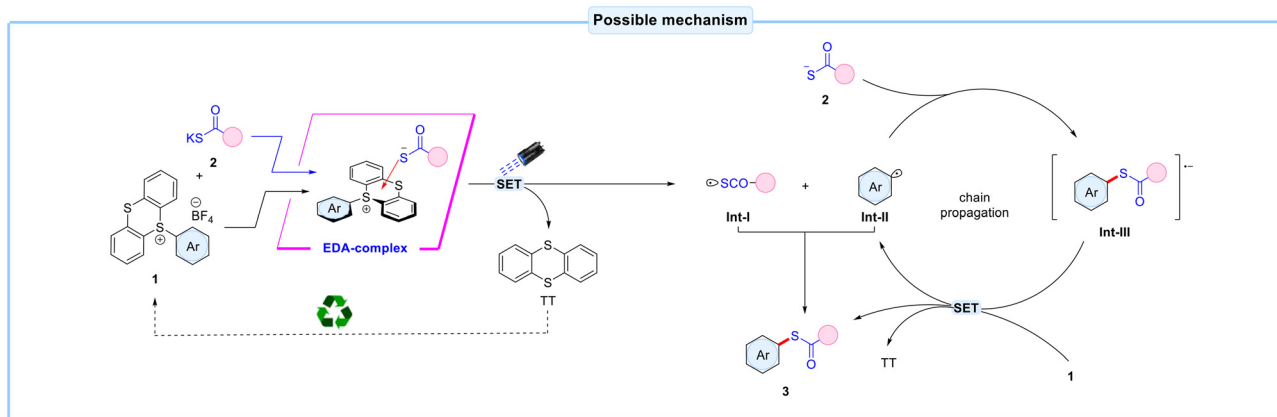
To evaluate the eco-friendliness and greenness of our developed strategy, we assessed the green chemistry metrics for the synthesis of compound **3a** (0.06 g, 83%) from methyl-2-methoxybenzoate-derived thianthrenium salt **1a** (0.3 mmol, 0.141 g) and potassium thioacetate **2a** (0.6 mmol, 0.069 g). The reaction was carried out using 4 Å molecular sieve (0.06 g) in DMSO (1.65 g, 1.5 mL) under visible light conditions. The results, as shown in Fig. 2h, demonstrate that the green chemistry metrics are excellent, the E-factor is 14.95, process mass intensity (PMI) is 15.37%, reaction mass efficiency is 57.62%, atom economy (AE) is 78.34%, and carbon efficiency is 100%.<sup>20</sup> Additionally, the EcoScale score was calculated to be 70.5, which is considered acceptable in terms of sustainability (see ESI† for detailed discussion).



**Fig. 2** (a) UV-visible absorption spectra; (b) Job's plot; (c) <sup>1</sup>H NMR titration experiment; (d) radical trap experiment; (e) sunlight irradiation; (f) scale-up; (g) sensitivity assessment of reaction; (h) green chemistry metrics (our method).

In light of all the above experimental data, a plausible mechanism for this EDA-complex mediated thioesterification reaction is depicted in Scheme 3. Initially, an EDA-complex

aggregate is formed between **1** and potassium thioacid salt **2**. Under visible light irradiation, this EDA-complex undergoes a single-electron transfer (SET) event from thiolate anion to the



Scheme 3 Possible mechanism.

aryl thianthrenium salt **1**, generating thiyl radical intermediate **Int-I**, aryl radical intermediate **Int-II**, and recyclable thianthrene by-product. The generated thiyl radical intermediate **Int-I** and aryl radical intermediate **Int-II**, can undergo subsequent radical–radical coupling, leading to the formation of the desired product **3**. Moreover, the aryl radical intermediate **Int-II** interacts with thiolate anion **2** to give **Int-III**. This intermediate **Int-III** undergoes SET with **1** to give the desired product **3**, thianthrene by-product, and **Int-II**, which propagates the chain again.

## Conclusion

In summary, we have developed an EDA-complex mediated thioesterification reaction *via* site-selective C–H functionalization using thianthrenium salts. This protocol is compatible with a wide range of aromatic feedstock, and a diverse range of alkyl, aryl, and heteroaryl potassium thioacid salts, yielding the desired thioester products in good yields. Additionally, we demonstrated the versatility of the method through late-stage thioesterification of agrochemical and pharmaceutical compounds. Our method offers several key advantages over previously known methods, including (I) C–H functionalization; (II) halide retention; (III) compatibility with alkyl, aryl, and heteroaryl potassium thioacid salts; and (IV) reusable and recyclable by-products, and (V) excellent atom economy, and *E*-factor scores, which are considered excellent in terms of safety, economic and ecological yardsticks, making it a practical solution for both academic research and industrial applications.

## Author contributions

R. I. Patel optimized the reaction conditions and synthesized all the derivatives. B. Saxena synthesized the potassium thioacid salts. R. I. Patel and B. Saxena performed the mechanistic studies and wrote the manuscript with the helpful insights of

Prof. A. Sharma. Prof. A. Sharma supervised the whole work, interpreted the results, and edited the manuscript. All the authors have given their approval to the final version of the manuscript.

## Data availability

The data supporting this article have been included as part of the ESI.†

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

Financial support from SERB (CRG/2022/002691), India is gratefully acknowledged. We also acknowledge DST-FIST (SR/FST/CSII/2018/72(C)) for the NMR and HRMS facilities in the Chemistry Department, IIT Roorkee. R. P. and B. S. thank UGC and CSIR for the SRF Fellowship, respectively.

## References

- (a) K. Taori, V. J. Paul and H. Luesch, *J. Am. Chem. Soc.*, 2008, **130**, 1806–1807; (b) X.-L. Liu, Y. Shi, J. S. Kang, P. Oelschlaeger and K.-W. Yang, *ACS Med. Chem. Lett.*, 2015, **6**, 660–664; (c) Y. Kanda, T. Ashizawa, S. Kakita, Y. Takahashi, M. Kono, M. Yoshida, Y. Saitoh and M. Okabe, *J. Med. Chem.*, 1999, **42**, 1330–1332; (d) S. Aksakal, R. Aksakal and C. R. Becer, *Polym. Chem.*, 2018, **9**, 4507–4516; (e) N. Martina, V. Neelz and H.-E. Spinnler, *Food Qual. Prefer.*, 2004, **15**, 247–257.
- (a) K. Chandru, A. Gilbert, C. Butch, M. Aono and H. J. Cleaves, *Sci. Rep.*, 2016, **6**, 29883; (b) J. Franke and C. Hertweck, *Cell Chem. Biol.*, 2016, **23**, 1179–1192.

- 3 (a) Y. Yang, J. Zhu, M. Hassink, L. M. M. Jenkins, Y. Wan, D. H. Appella, J. Xu, E. Appella and X. Zhang, *Emerging Microbes Infect.*, 2019, **6**(1), 1–8; (b) H. Shinkai, K. Maeda, T. Yamasaki, H. Okamoto and I. Uchida, *J. Med. Chem.*, 2000, **43**, 3566–3572; (c) K. Yamada, M. Satoh, S. Shimura, T. Sasaki, T. Takishima and K. Shirato, *Respiration*, 1994, **61**, 42–47; (d) J. T. Suh, J. W. Skiles, B. E. Williams, R. D. Youssefyeh, H. Jones, B. Loev, E. S. Neiss, A. Schwab and W. S. Mann, *J. Med. Chem.*, 1985, **28**, 57–66; (e) D. M. Peter, B. Vögeli, N. S. Cortina and T. J. Erb, *Molecules*, 2016, **21**(4), 517.
- 4 (a) A. Pal, P. P. Mondal, F. Niloofar and B. Sahoo, *Eur. J. Org. Chem.*, 2022, e202201159; (b) V. Hirschbeck, P. H. Gehrtz and I. Fleischer, *Chem. – Eur. J.*, 2018, **24**, 7092–7107; (c) M. Kazemi and L. Shiri, *J. Sulfur Chem.*, 2015, **36**, 613–623.
- 5 For selected examples see: (a) V. Hirschbeck, P. H. Gehrtz and I. Fleischer, *J. Am. Chem. Soc.*, 2016, **138**, 16794–16799; (b) H.-J. Ai, W. Lu and X.-F. Wu, *Angew. Chem., Int. Ed.*, 2021, **60**, 17178–17184, (*Angew. Chem.*, 2021, **133**, 17315–17321); (c) M. Kim, S. Yu, J. G. Kim and S. Lee, *Org. Chem. Front.*, 2018, **5**, 2447–2452; (d) Y. Kim, K. H. Song and S. Lee, *Org. Chem. Front.*, 2020, **7**, 2938–2943; (e) Y. Li, G. Bao and X.-F. Wu, *Chem. Sci.*, 2020, **11**, 2187–2192; (f) S. Murakami, T. Nanjo and Y. Takemoto, *Org. Lett.*, 2021, **23**, 7650–7655; (g) Y.-W. Huo, X. Qi and X.-F. Wu, *Org. Lett.*, 2022, **24**, 4009–4013; (h) W. Wang, X. Qi and X.-F. Wu, *Adv. Synth. Catal.*, 2021, **363**, 2541–2545; (i) H.-J. Ai, J. Rabeah, A. Brückner and X.-F. Wu, *Chem. Commun.*, 2021, **57**, 1466–1469; (j) F. Zhao and X.-F. Wu, *Org. Chem. Front.*, 2021, **8**, 670–675; (k) J. Luo, M. Rauch, L. Avram, Y. Diskin-Posner, G. Shmul, Y. Ben-David and D. Milstein, *Nat. Catal.*, 2020, **3**, 887–892; (l) X. Qi, Z.-P. Bao, X.-T. Yao and X.-F. Wu, *Org. Lett.*, 2020, **22**, 6671–6676.
- 6 For selected examples see. (a) S.-H. Guo, X.-L. Zhang, G.-F. Pan, X.-Q. Zhu, Y.-R. Gao and Y.-Q. Wang, *Angew. Chem., Int. Ed.*, 2018, **57**, 1663–1667, (*Angew. Chem.*, 2018, **130**, 1679–1683); (b) S.-H. Guo, M.-Y. Wang, G.-F. Pan, X.-Q. Zhu, Y.-R. Gao and Y.-Q. Wang, *Adv. Synth. Catal.*, 2018, **360**, 1861–1869; (c) B. Xu, D. Li, L. Lu, D. Wang, Y. Hu and Q. Shen, *Org. Chem. Front.*, 2018, **5**, 2163–2166; (d) Y.-D. Kwon, M. T. La and H.-K. Kim, *New J. Chem.*, 2018, **42**, 10833–10841; (e) P. Gopinath, R. S. Vidyarini and S. Chandrasekaran, *J. Org. Chem.*, 2009, **74**, 6291–6294; (f) Y.-M. Xiao, Y. Zhao, J.-Q. Li, J.-W. Yuan, L.-R. Yang, P. Mao and W.-P. Mai, *New J. Chem.*, 2023, **47**, 17092–17097; (g) I. Tiefenbrunner, B. R. Brutiu, T. Stopka and N. Maulide, *J. Org. Chem.*, 2023, **88**, 3941–3944; (h) J. Xu, F. Lu, L. Sun, M. Huang, J. Jiang, K. Wang, D. Ouyang, L. Lu and A. Lei, *Green Chem.*, 2022, **24**, 7350–7354; (i) G. Liu, S. Xu, Y. Yue, C. Su and W. Song, *Chem. Commun.*, 2024, **60**, 6154–6157; (j) M.-C. Wang, X.-Y. Yang, J.-F. Zhou, W.-X. Zhang and B.-J. Li, *Chem. Commun.*, 2024, **60**, 6671–6674.
- 7 (a) C. R. J. Stephenson, T. Yoon and D. W. C. Macmillan, in *Visible light photocatalysis in organic chemistry*, Wiley-VCH, German, 2018; (b) L. Marzo, S. K. Pagire, O. Reiser and B. König, *Angew. Chem., Int. Ed.*, 2018, **57**, 10034–10072, (*Angew. Chem.*, 2018, **130**, 10188–10228); (c) D. S. Hamilton and D. A. Nicewicz, *J. Am. Chem. Soc.*, 2012, **134**, 18577–18580; (d) D. Ravelli, S. Protti and M. Fognoni, *Chem. Rev.*, 2016, **116**, 9850–9913; (e) R. I. Patel, S. Sharma and A. Sharma, *Org. Chem. Front.*, 2021, **8**, 3166–3200; (f) N. Kvasovs and V. Gevorgyan, *Chem. Soc. Rev.*, 2020, **50**, 2244–2259; (g) K. L. Skubi, T. R. Blum and T. P. Yoon, *Chem. Rev.*, 2016, **116**, 10035–10074; (h) T. Constantin, F. Julia and D. Leonori, *Chem. Rev.*, 2022, **122**, 2292–2352; (i) L. Capaldo, D. Ravelli and M. Fagnoni, *Chem. Rev.*, 2022, **122**, 1875–1924; (j) F. Mo, D. Qiu, L. Zhang and J. Wang, *Chem. Rev.*, 2021, **121**, 5741–5829; (k) R. I. Patel, A. Sharma, S. Sharma and A. Sharma, *Org. Chem. Front.*, 2021, **8**, 1694–1718; (l) R. I. Patel, J. Singh and A. Sharma, *ChemCatChem*, 2022, **14**, e202200260; (m) B. Saxena, R. I. Patel, J. Tripathi and A. Sharma, *Org. Biomol. Chem.*, 2023, **21**, 4723–4743; (n) B. Saxena, R. I. Patel and A. Sharma, *RSC Sustainability*, 2024, **2**, 2169–2189; (o) P. Ranjan, S. Pillitteri, G. Coppola, M. Oliva, E. V. Van der Eycken and U. K. Sharma, *ACS Catal.*, 2021, **11**, 10862–10870; (p) G. A. Coppola, S. Pillitteri, E. V. Van der Eycken, S.-L. You and U. K. Sharma, *Chem. Soc. Rev.*, 2022, **51**, 2313–2382.
- 8 For selected examples, see. (a) G. Liu, N. Zheng, X. Duan, X. Sun and W. Song, *Green Chem.*, 2023, **25**, 5035–5040; (b) H. Wang, Z. Liu, A. Das, P. Bellotti, S. Megow, F. Temps, X. Qi and F. Glorius, *Nat. Synth.*, 2023, **2**, 1116–1126; (c) G. Bogonda, D. V. Patil, H. Y. Kim and K. Oh, *Org. Lett.*, 2019, **21**, 3774–3779; (d) S. Murakami, T. Nanjo and Y. Takemoto, *Org. Lett.*, 2021, **23**, 7650–7655; (e) V. J. Roy, P. P. Sen and S. R. Roy, *J. Org. Chem.*, 2021, **86**, 16965–16976; (f) J. Su, A. Chen, G. Zhang, Z. Jiang and J. Zhao, *Org. Lett.*, 2023, **25**, 8033–8037; (g) S. Mukherjee, T. Patra and F. Glorius, *ACS Catal.*, 2018, **8**, 5842–5846; (h) M.-u.-S. Bhat, M. A. Ganie, S. Kumar, M. A. Rizvi, S. Raheem and B. A. Shah, *J. Org. Chem.*, 2024, **89**, 4607–4618; (i) G.-Q. Hu, W.-Y. Zhang, Y.-X. Liu, J.-H. Liu and B. Zhao, *J. Org. Chem.*, 2023, **88**, 14351–14356; (j) K. Sun, T. Sun, Y. Jiang, J. Shi, W. Sun, Y. Zheng, Z. Wang, Z. Li, X. Lv, X. Zhang, F. Luo and S. Liu, *Chem. Commun.*, 2024, **60**, 5755–5758.
- 9 M. Yang, T. Cao, T. Xu and S. Liao, *Org. Lett.*, 2019, **21**, 8673–8678.
- 10 T. Xu, T. Cao, M. Yang, R. Xu, X. Nie and S. Liao, *Org. Lett.*, 2020, **22**, 3692–3696.
- 11 (a) W. Zheng, Y. Xu and L. Lin, *ChemPhotoChem*, 2022, **6**, e202100264; (b) A. A. Volkov, D. I. Bugaenko, A. V. Bogdanov and A. V. Karchava, *J. Org. Chem.*, 2022, **87**, 8170–8182.
- 12 H. Tang, M. Zhang, Y. Zhang, P. Luo, D. Ravelli and J. Wu, *J. Am. Chem. Soc.*, 2023, **145**, 5846–5854.
- 13 G.-Q. Hu, W.-Y. Zhang, Y.-X. Liu, J.-H. Liu and B. Zhao, *J. Org. Chem.*, 2023, **88**, 14351–14356.
- 14 For selected examples see. (a) F. Berger, M. B. Plutschack, J. Riegger, W. Yu, S. Speicher, M. Ho, N. Frank and T. Ritter, *Nature*, 2019, **567**, 223–228; (b) P. S. Engl,

- A. P. Häring, F. Berger, G. Berger, A. Pérez-Bitrián and T. Ritter, *J. Am. Chem. Soc.*, 2019, **141**, 13346–13351; (c) F. Ye, F. Berger, H. Jia, J. Ford, A. Wortman, J. Börgel, C. Genicot and T. Ritter, *Angew. Chem., Int. Ed.*, 2019, **58**, 14615–14619; (d) R. Sang, S. E. Korkis, W. Su, F. Ye, P. S. Engl, F. Berger and T. Ritter, *Angew. Chem., Int. Ed.*, 2019, **58**, 16161–16166; (e) J. Li, J. Chen, R. Sang, W.-S. Ham, M. B. Plutschack, F. Berger, S. Chhabra, A. Schnegg, C. Genicot and T. Ritter, *Nat. Chem.*, 2020, **12**, 56–62; (f) F. Juliá, J. Yan, F. Paulus and T. Ritter, *J. Am. Chem. Soc.*, 2021, **143**, 12992–12998; Y. Cai, S. Chatterjee and T. Ritter, *J. Am. Chem. Soc.*, 2023, **145**, 13542–13548; (g) A. Granados, M. J. Cabrera-Afonso, M. Escolano, S. O. Badir and G. A. Molander, *Chem. Catal.*, 2022, **2**, 898–907; (h) M. J. Cabrera-Afonso, A. Granados and G. A. Molander, *Angew. Chem.*, 2022, **134**, e202202706; (i) K. Sun, A. Shi, Y. Liu, X. Chen, P. Xiang, X. Wang, L. Qua and B. Yu, *Chem. Sci.*, 2022, **13**, 5659–5666; (j) E. M. Alvarez, Z. Bai, S. Pandit, N. Frank, L. Torkowski and T. Ritter, *Nat. Synth.*, 2023, **2**, 548–556; (k) P. Sarró, A. Gallego-Gamo, R. Pleixats, A. Vallribera, C. Gimbert-Suriñach and A. Granados, *Adv. Synth. Catal.*, 2024, **366**, 2587; (l) A. Dewanji, L. van Dalsen, J. A. Rossi-Ashton, E. Gasson, G. E. M. Crisenza and D. J. Procter, *Nat. Chem.*, 2023, **15**, 43–52.
- 15 (a) A. K. Wortman and C. R. J. Stephenson, *Chem*, 2023, **9**, 1–26; (b) G. E. M. Crisenza, D. Mazzarella and P. Melchiorre, *J. Am. Chem. Soc.*, 2020, **142**(12), 5461–5476; (c) C. G. S. Lima, T. D. M. Lima, M. Duarte, I. D. Jurbery and M. W. Paixão, *ACS Catal.*, 2016, **6**, 1389–1407; (d) Y. Q. Yuan, S. Majumder, M. H. Yang and S. R. Guo, *Tetrahedron Lett.*, 2020, **61**, 151506; (e) P. Garra, J. P. Fouassier, S. Lakhdar, Y. Yagci and J. Lalevée, *Prog. Polym. Sci.*, 2020, **107**, 101277; (f) M. Mondal, S. Ghosh, D. Lai and A. Hajra, *ChemSusChem*, 2024, e202401114.
- 16 (a) B. Saxena, R. I. Patel and A. Sharma, *Adv. Synth. Catal.*, 2023, **365**, 1538–1564; (b) A. Sharma, N. Singh, R. Kumar and A. Sharma, *Adv. Synth. Catal.*, 2024, **366**, 2735–2741; (c) N. Singh, S. Sharma and A. Sharma, *Adv. Synth. Catal.*, 2023, **365**, 3505–3511; (d) A. Monga, S. Bagchi, R. K. Soni and A. Sharma, *Adv. Synth. Catal.*, 2020, **362**, 2232–2237.
- 17 K. Shin, Y. Zheng, F. Zhang, S. Wu and Y. Tang, *ACS Appl. Energy Mater.*, 2020, **3**, 7030–7038.
- 18 W. Liu, H. Hou, H. Jing, S. Huang, W. Ou and C. Su, *Org. Lett.*, 2023, **25**, 8350–8355.
- 19 L. Pitzer, F. Schäfers and F. Glorius, *Angew. Chem., Int. Ed.*, 2019, **58**, 8572–8576.
- 20 (a) K. Van Aken, L. Streckowski and L. Patiny, *Beilstein J. Org. Chem.*, 2006, **2**, 3; (b) A. Beillard, X. Bantreil, T.-X. Métro, J. Martinez and F. Lamaty, *Green Chem.*, 2018, **20**, 964–968; (c) B. Saxena, R. I. Patel, S. Sharma and A. Sharma, *Green Chem.*, 2024, **26**, 2721–2729; (d) Y. Zhang, J. Mao, Z. Wang, L. Tang and Z. Fan, *Green Chem.*, 2024, **26**, 9371–9377.