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Visible light-driven α -sulfonylation of ketone-derived silyl enol ethers *via* an electron donor–acceptor complex†

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The diverse utility of β -ketosulfones in pharmaceuticals and bioactive compounds has generated considerable interest in their synthesis. However, existing synthetic approaches often depend on transition-metal catalysts, which require extensive purification and result in low yields. Herein, we present a cost-effective, metal- and photocatalyst-free, visible light electron donor–acceptor (EDA) complex-mediated sulfonylation of ketone-derived silyl enol ethers with thiosulfonates (acceptor) and DABCO as an electron donor under mild conditions, offering a more efficient and straightforward approach. Our method enables the synthesis of a diverse range of β -ketosulfone derivatives, including biologically active and late-stage molecules, in good yields. Our strategy offers several significant advantages over existing techniques, which include (i) transition-metal and photoredox catalyst-free conditions; (ii) no need for an external SO_2 source; (iii) broad substrate scope; (iv) recyclable and reusable by-products; and (v) excellent atom economy, reaction mass efficiency, process mass intensity, and *E*-factor and EcoScale scores, highlighting its efficiency and economic sustainability. Detailed mechanistic studies confirm the involvement of an EDA-complex-mediated radical process that operates without a catalyst.

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Introduction

Organosulfones are essential in synthetic and medicinal chemistry, particularly in modulating and controlling drug metabolism and biotransformation rates.¹ Several clinically approved drugs containing sulfone groups are shown in Fig. 1A. Among these, β -ketosulfones have garnered significant attention from synthetic chemists due to their versatile applications in organic synthesis and their considerable pharmaceutical relevance.² These compounds demonstrate a wide array of biological activities, making them particularly valuable in organic synthesis. As a result, extensive efforts have been devoted to synthesizing β -ketosulfones, underscoring their importance in chemical and biological contexts. Several methods have been developed for the synthesis of β -ketosulfones. These include the acylation of alkyl sulfones with acid chlorides, esters, or *N*-acyl benzotriazoles,³ the oxidation of β -ketosulfides or β -hydroxysulfones with stoichiometric oxidants,⁴ and the alkylation of metallic arene sulfinates with α -halo- or α -tosyloxy ketones.⁵ All these methods have drawbacks such as the use of

costly transition metal catalysis, hazardous oxidants, and elevated temperatures. In this regard, radical sulfonylation has gained attention as an effective approach for synthesizing β -ketosulfones. This method typically involves the reaction of alkenes,⁶ alkynes⁷ or activated alkenes⁸ with various radical sulfonylation reagents. Although traditional methods are useful, they often require cumbersome multi-step prefunctionalization or preactivation of the starting materials, have limited substrate scopes, yield low results, or involve harsh reaction conditions, all of which present significant challenges. In the past decade, visible-light-promoted photoredox reactions have achieved significant milestones in organic synthesis.⁹ Amongst these, three methods involving visible light-promoted synthesis of β -ketosulfones have been reported involving the reaction between silyl enol ethers, an external sulfur dioxide (SO_2) source, and an alkyl/aryl radical source. The first of these was disclosed by Ye, Wu, and co-workers in 2019, who reported a visible light Ir-catalyzed sulfonylation of silyl enol ethers using amine-derived Katritzky salts as alkyl radical precursors with $\text{K}_2\text{S}_2\text{O}_5$ as a SO_2 source (Fig. 1Ba).¹⁰ However, the substrate scope was limited and did not include aryl radical intermediates and alkyl-substituted *O*-silyl enol ethers. Moreover, the reaction produced 2,4,6-triphenylpyridine as a non-reusable by-product. The following year, Wu and co-workers developed a Ru-catalyzed synthesis of β -ketosulfones *via* a three-component reaction using aryldiazonium tetrafluoroborates, $\text{Na}_2\text{S}_2\text{O}_5$, and 2,2-difluoroenol silyl ethers under

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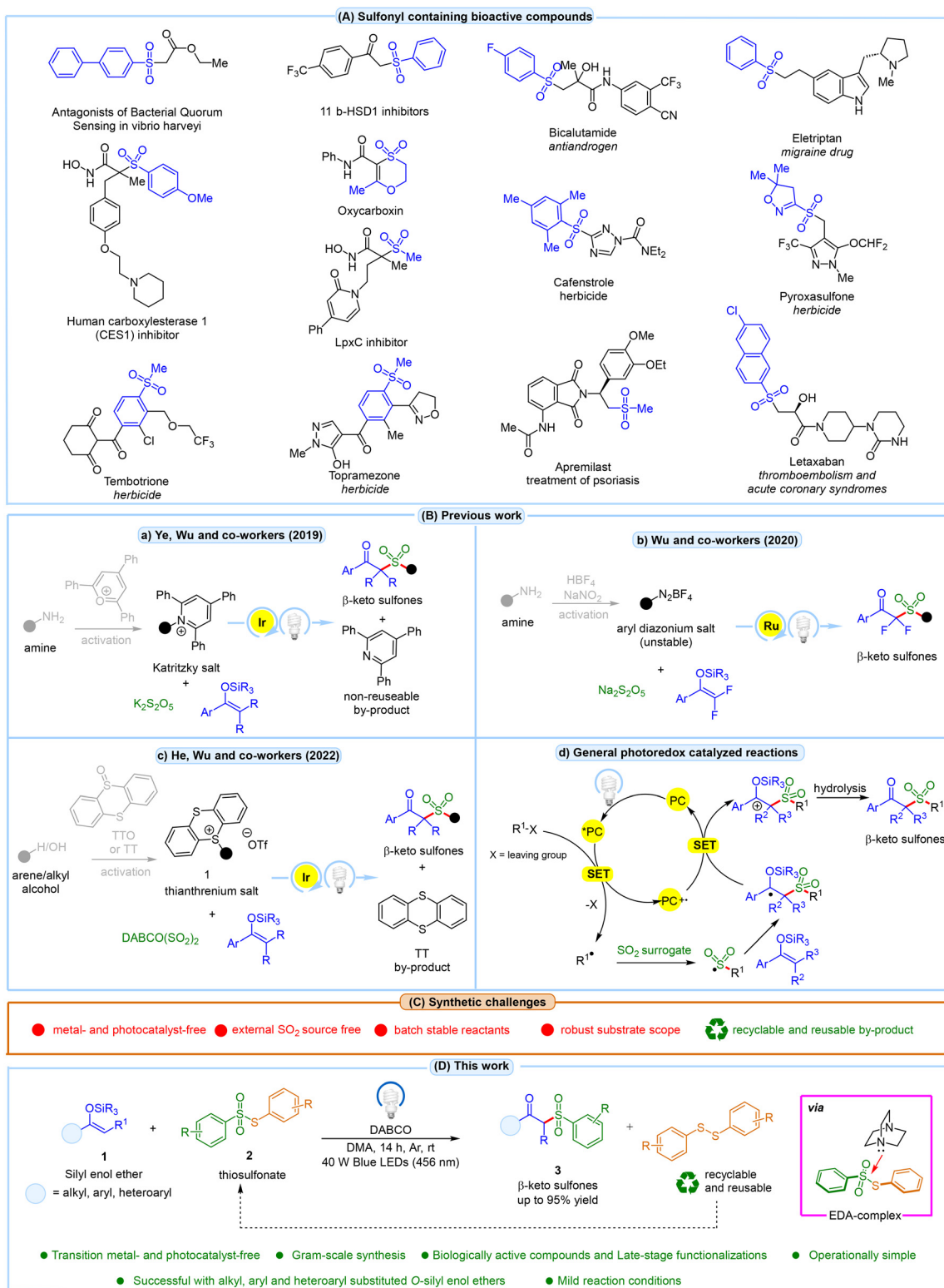


Fig. 1 (A) Sulfonyl-containing bioactive drugs; (B) previous work; (C) synthetic challenges; and (D) this work.

photochemical conditions (Fig. 1Bb).¹¹ Additionally, this method also suffers from a narrow substrate scope due to the use of unstable diazonium salts with an expensive photo-

catalyst. Later in 2022, He, Wu, and co-workers introduced an Ir-catalyzed sulfonylation of O-silyl enol ethers with DABCO-(SO_2)₂ and arene or alcohol-derived thianthrenium

salts (Fig. 1Bc).¹² Despite its innovation, the method has limitations, including its restriction to aryl-substituted *O*-silyl enol ethers, reliance on an expensive photoredox catalyst, the use of a complex and costly SO₂ surrogate, and a long reaction time of 48 h. Typically, in these methods, the photoexcited catalyst (*PC) reduces the starting substrate *via* single electron transfer (SET), generating a radical intermediate (Fig. 1Bd). This radical then reacts with the external SO₂ source, forming a sulfonyl radical intermediate, which adds to the silyl enol ethers, followed by SET reduction and subsequent hydrolysis, yielding β-ketosulfones. Consequently, developing a method for β-ketosulfone synthesis that eliminates the need for any transition metals or photocatalysts and directly leads to the sulfonyl radical intermediate would be a significant breakthrough and a major advancement in β-ketosulfone synthesis.

In this context, electron donor–acceptor (EDA) complex-based reactions have garnered significant interest in organic synthesis.¹³ Such EDA complexes, formed in the ground state, can absorb visible light, triggering an intermolecular single electron transfer (SET) event and eliminating the critical need for a customary redox potential matching in such reactions. Building on our ongoing research into green and sustainable photoinduced EDA-complex reactions,¹⁴ we present a practical method for the synthesis of β-ketosulfones using silyl enol ethers with thiosulfonates (acceptor) and DABCO (donor) under visible light irradiation (Fig. 1D).

We hypothesize that an EDA-complex forms between thiosulfonates (acceptor) and DABCO (donor), leading to the generation of an S-centered radical intermediate. This intermediate

reacts with the silyl enol ether to yield the desired β-ketosulfone product and an aryl disulfide by-product. The use of an inexpensive base as an electron donor and thiosulfonates as a sulfonyl source under photochemical conditions facilitates the synthesis of β-ketosulfones, a challenging feat to achieve without the need for expensive photoredox catalysis and an external SO₂ source. Additionally, the aryl disulfide by-product generated in the reaction is recyclable and reusable, offering additional advantages in terms of economic sustainability. Our method also enables the synthesis of various bioactive molecules and late-stage modification of pharmaceutical complexes, thus significantly benefiting industrial and academic research.

Results and discussion

We initiated our study using the trimethylsilyl enol ether of acetophenone **1a** and *S*-phenyl 4-methylbenzenesulfonothioate **2a** as model substrates in MeCN solvent, with DBU as a base, under 456 nm Kessil lamp irradiation. To our satisfaction, the desired product was obtained in 52% yield (entry 1). Further screening of other bases proved unproductive (entries 2–6) except for DABCO, which emerged as the optimal base, delivering the highest yield of 67% (entry 6). We then tested various solvents, including DMSO, acetone, DCM, DMF, DCE, EtOH, and MeOH (entries 7–15). Among these, the reaction achieved an 81% yield in DMA solvent (entry 11), likely due to the efficient solubilization of the reaction components (**1a** and **2a**).

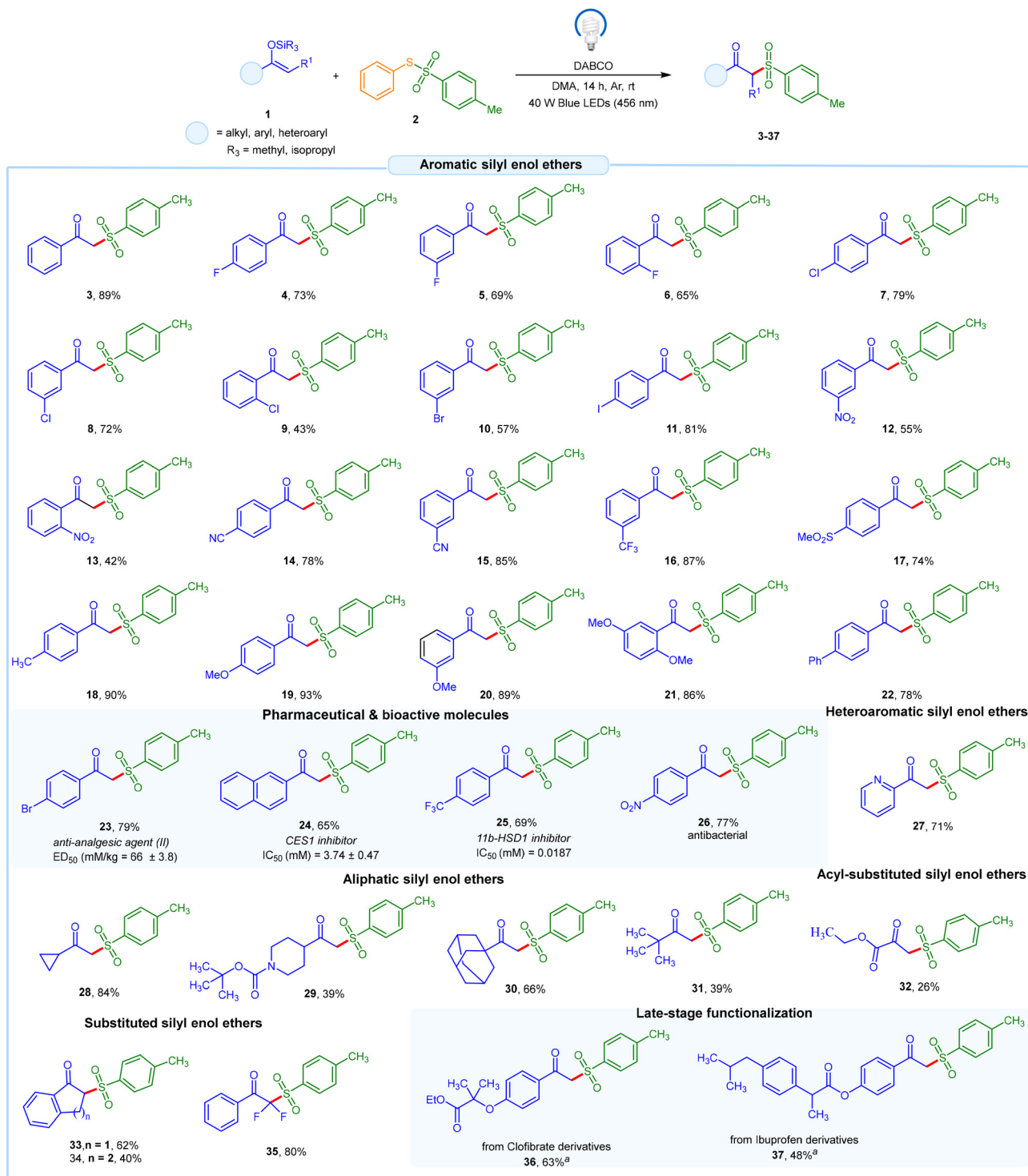
Table 1 Optimization of the reaction conditions

Entry	Base (2.0 equiv.)	Solvent	Light	Yield (%) ^a	Entry	Base (2.0 equiv.)	Solvent	Light	Yield (%)
1	DBU	MeCN	456	52	14	DABCO	THF	456	42
2	TMEDA	MeCN	456	48	15	DABCO	MeOH	456	N.D
3	DMAP	MeCN	456	30	16 ^b	DABCO	DMA	456	55
4	Et ₃ N	MeCN	456	Trace	17 ^c	DABCO	DMA	456	89
5	Triphenylamine	MeCN	456	Trace	18 ^d	—	DMA	456	N.D
6	DABCO	MeCN	456	67	19 ^e	DABCO	DMA	Dark	N.D
7	DABCO	DMSO	456	72	20	DABCO	DMA	23 W CFL bulb	40
8	DABCO	Acetone	456	38	21	DABCO	DMA	427	65
9	DABCO	DCM	456	52	22	DABCO	DMA	390	55
10	DABCO	DMF	456	70	23	DABCO	DMA	Green LED	72
11	DABCO	DMA	456	81	24 ^f	DABCO (TIPS)	DMA	456	52
12	DABCO	EtOH	456	N.D	25 ^g	DABCO (TES)	DMA	456	0
13	DABCO	DCE	456	55	26 ^h	DABCO (TBS)	DMA	456	0

^a Reaction conditions: a mixture of *O*-silyl enol ether [Si] = TMS (trimethylsilyl) **1a** (0.2 mmol), *S*-phenyl 4-methylbenzenesulfonothioate **2a** (0.24 mmol, 1.2 equiv.), base (0.4 mmol, 2.0 equiv.) and solvent (0.5 mL) under visible light irradiation (light source) for 14 h under an Ar atmosphere at rt. ^b Base (0.2 mmol 1.0 equiv.). ^c Base (0.6 mmol 3.0 equiv.). ^d No base. ^e In the dark. ^f [Si] = TIPS (triisopropylsilyl). ^g [Si] = TES (triethylsilyl). ^h [Si] = TBS (*tert*-butyldimethylsilyl).

Notably, mixing **1a** and **2a** led to an intense yellow color change in the reaction mixture, suggesting the formation of an EDA-complex. The reaction yield was 55% when 1.0 equiv. of DABCO was used (entry 16). However, the yield increased to

89% when 3.0 equiv. of DABCO was used (entry 17). The reaction did not proceed without DABCO, underscoring its critical role in product formation (entry 18). Additionally, the reaction failed without light, confirming the necessity of light

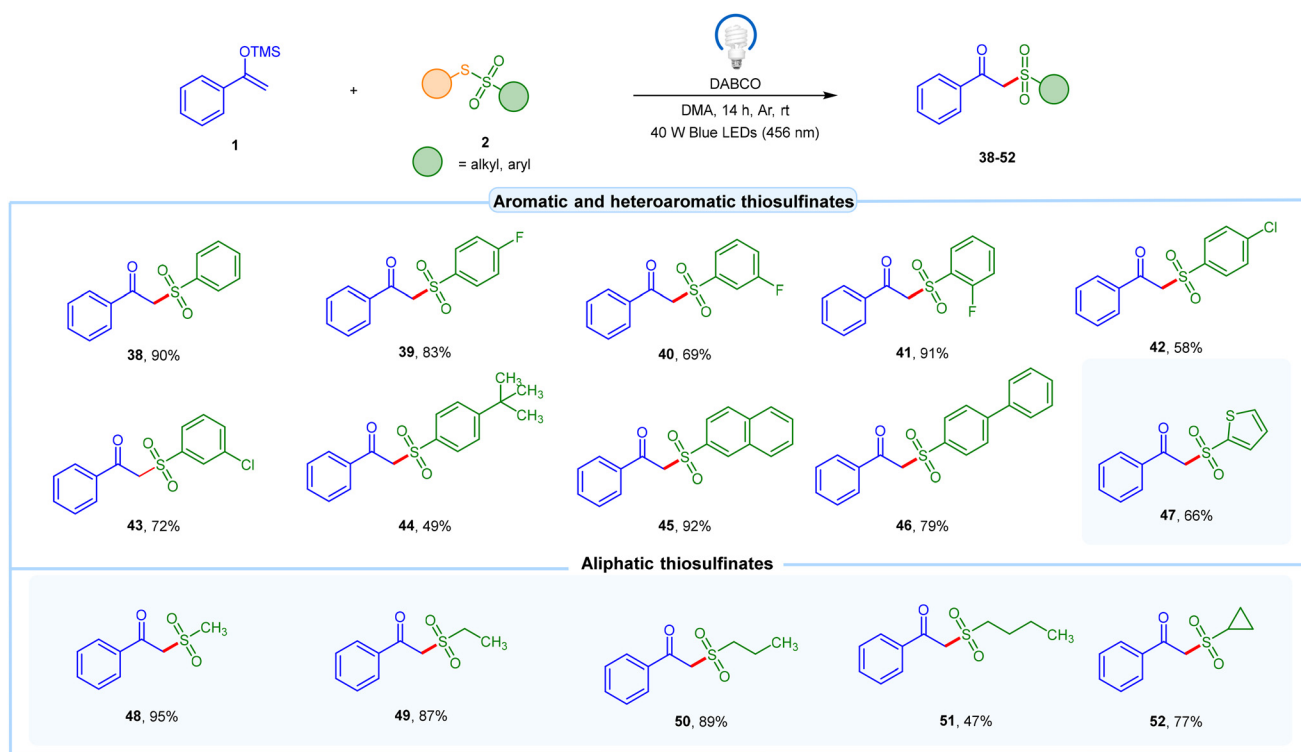


Scheme 1 Reaction conditions: *O*-silyl enol ethers **1** (0.2 mmol, where R = Me), *S*-phenyl 4-methylbenzene sulfonothioate **2a** (1.2 equiv., 0.24 mmol), and DABCO (3 equiv., 0.6 mmol) in 0.5 mL DMA were irradiated with a 456 nm Kessil lamp at room temperature under argon for 14 h. ^aR = isopropyl.

irradiation (entry 19). Screening of other light sources resulted in reduced yields (entries 20–23), indicating that 456 nm Kessil lamp irradiation is optimal for product formation. Lastly, we investigated compound **1**, which features sterically more demanding triisopropylsilyl (TIPS), triethylsilyl (TES), and *tert*-butyldimethylsilyl (TBS) groups in place of the trimethylsilyl (TMS) group of the acetophenone-derived enol ether under the optimized conditions, leading to reduced yields or complete failure (entries 24–26) and indicating that the TMS group in enol ethers is crucial for a high yield of this reaction. In a condition-based sensitivity assessment, our EDA-protocol transformation was sensitive to high oxygen concentration, low temperature, and low light intensity. However, it generally exhibited tolerance to variations in substrate concentration, high reaction temperature, high light intensity, and water (see the radar diagram in Table 1 and the ESI† for details).

After establishing the optimized reaction conditions, we applied the protocol to various *O*-silyl enol ethers of acetophenone bearing alkyl, aryl, and heteroaryl functionalities, using *S*-phenyl 4-methylbenzenesulfonothioate **2a** as the coupling partner with DABCO. The *O*-silyl enol ether of acetophenone bearing halide groups, such as fluoro-, chloro-, bromo-, and iodo-substituents, reacted smoothly, producing the desired products in moderate to good yields (**4–11**). Additionally, aryl substituted *O*-silyl enol ethers with both electron-withdrawing and electron-donating substituents, such as nitro-, cyano-, trifluoromethyl-, methylsulfonyl-, methyl, and methoxy groups,

were well tolerated under the optimized conditions, resulting in moderate to excellent yields (**12–21**). Additionally, biphenyl β -keto sulfone **22** was obtained with a 78% yield. We were pleased to discover that this mild protocol demonstrated good reactivity towards the synthesis of pharmaceutical and bio-active molecules under the optimized conditions. For example, β -keto sulfone **23**, known for its excellent anti-analgesic properties, was synthesized with a 79% yield. Furthermore, β -keto sulfones **24** and **25**, which function as carboxylesterase 1 and 11 β -hydroxysteroid dehydrogenase type I inhibitors, were obtained with yields of 65% and 69%, respectively. Additionally, β -keto sulfone **26**, which exhibits anti-bacterial properties, was synthesized with a 77% yield. Furthermore, a heteroaryl *O*-silyl enol ether containing a pyridine group reacted well, producing the desired product **27** in 71% yield. To further assess the versatility of this method, a series of alkyl-substituted *O*-silyl enol ethers was subjected to the EDA-protocol, resulting in the desired products in moderate to good yields (**28–31**). Additionally, an acyl-substituted *O*-silyl enol ether was evaluated, resulting in the targeted product **32** in 26% yield. Next, an internal silyl enol ether derivative was successfully applied under the reaction protocol, yielding the desired products **33–34** in 62% and 40% yields, respectively. Finally, the utility of this method was further demonstrated through the late-stage functionalization of silyl enol ether derivatives derived from the hypertriglyceridemia drug clofibrate and the anti-inflammatory drug ibuprofen, yielding the



Scheme 2 Reaction conditions: *O*-silyl enol ether **1** (0.2 mmol), thiosulfonate **2a** (1.2 equiv., 0.24 mmol), and DABCO (3.0 equiv., 0.6 mmol) in 0.5 mL DMA were irradiated with a 456 nm Kessil lamp at room temperature under argon for 14 h.

desired products in 63% and 48% yields, respectively (Scheme 1).

To further diversify our strategy, we employed various thiosulfonates **2a** with alkyl and aryl functionalities under the optimized conditions with *O*-silyl enol ether **1**. These thiosulfonates **2a** reacted efficiently with *O*-silyl enol ether **1**, yielding the corresponding β -ketosulfones **38–52** in moderate to excellent yields, as illustrated in Scheme 2. Simple, non-substituted thiosulfonates **2** reacted well with *O*-silyl enol ether **1** to yield the desired product **38** in 90% yield. Furthermore, thiosulfonates **2** bearing halide groups (F and Cl) were well tolerated

under the reaction conditions, yielding the corresponding β -ketosulfones **39–43** in 58–91% yield. Notably, thiosulfonates **2** with a *t*-butyl group on the benzene ring afforded the β -ketosulfone product **44** in 49% yield. This method was also compatible with polyaromatic thiosulfonates, such as those with a naphthalene ring, affording the required product **45** in 92% yield. Conversely, biphenyl derived β -ketosulfone **46** was obtained in 79% yield. Impressively, heteroaromatic thiosulfonates were also well tolerated under the optimized conditions, resulting in the desired product **47** in 66% yield. Moreover, aliphatic thiosulfonates **2** were subjected to our EDA-complex

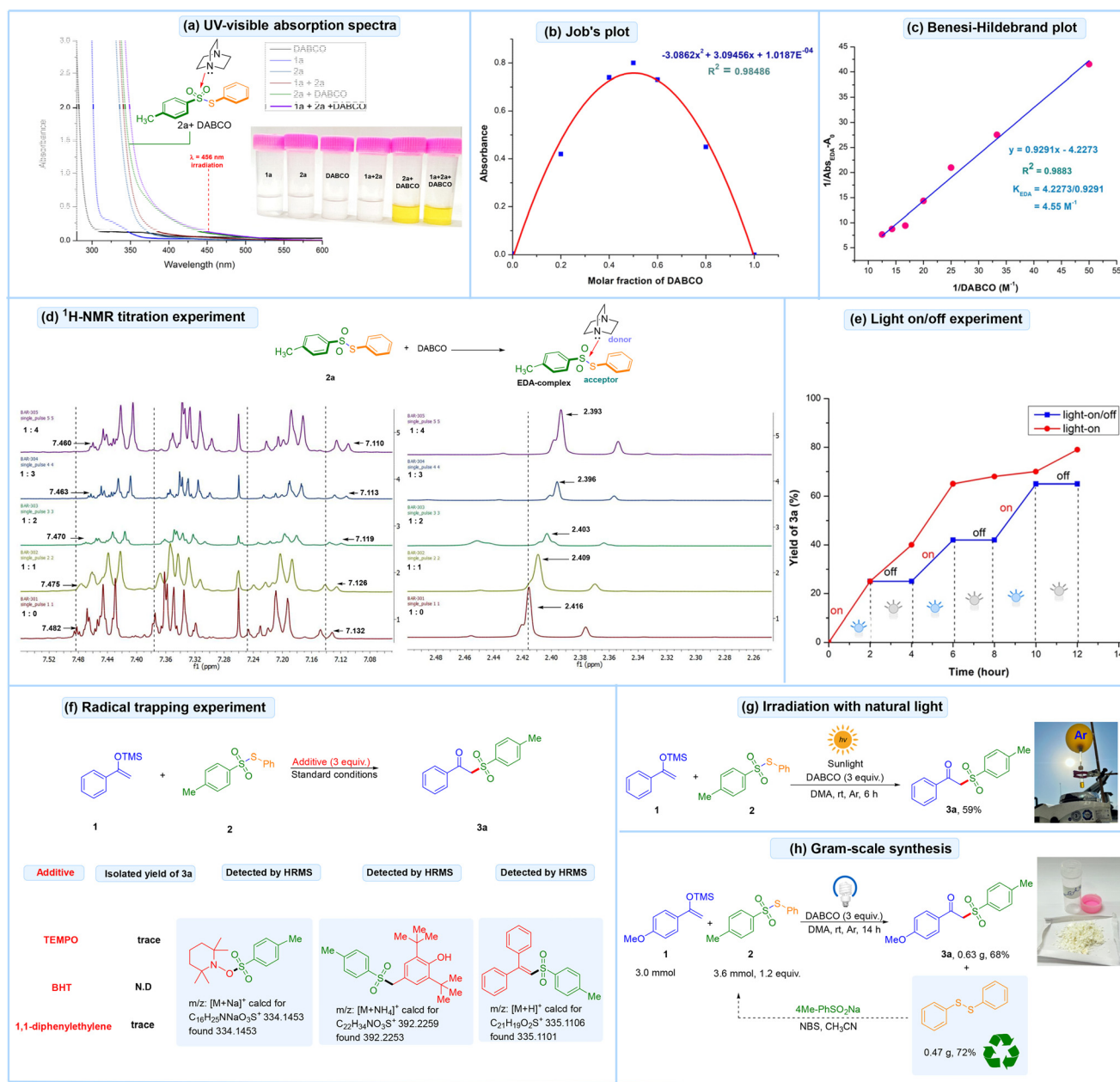


Fig. 2 (a) UV-visible absorption spectra; (b) Job's plot; (c) Benesi–Hildebrand plot; (d) ¹H-NMR titration experiment; (e) light on/off experiment; (f) radical trap experiment; (g) irradiation with natural light; and (h) gram-scale synthesis.

protocol with *O*-silyl enol ethers **1**. Aliphatic thiosulfonates **2** with primary alkyl chains (C1 to C4) were well tolerated under the reaction protocol, producing the desired products **48–51** in

47–95% yields. Additionally, thiosulfonates **2** with secondary alkyl chains proved compatible, yielding the target product **52** in 77% yield.

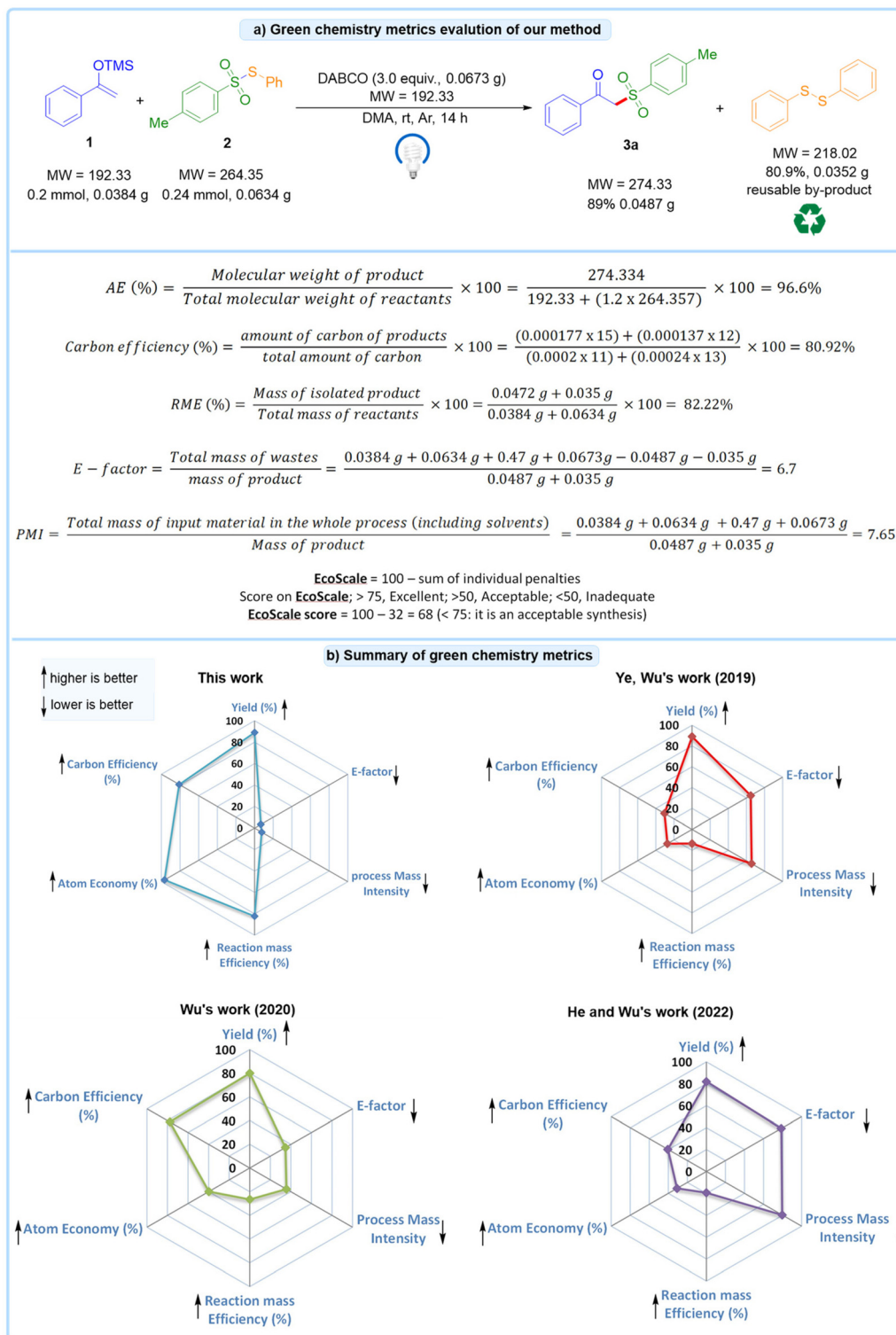


Fig. 3 Green chemistry metrics analysis: (a) green chemistry metrics evaluation of our method and (b) summary of the green chemistry metrics of our method compared to previous methods (see the ESI† for detailed calculation). Note: atom economy (AE), reaction mass efficiency (RME), process mass intensity (PMI), carbon efficiency, and *E*-factor. (↑) Higher is better and (↓) lower is better.

Mechanistic insights

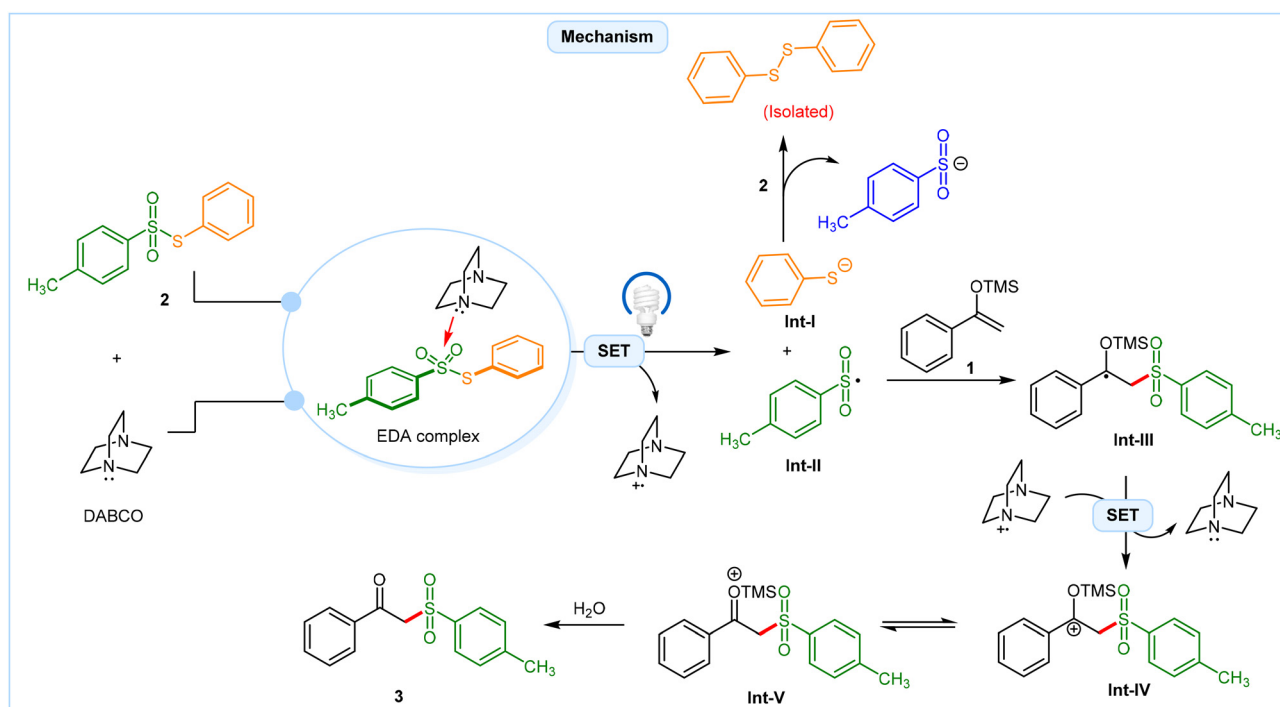
Next, a series of mechanistic studies were conducted to elucidate the reaction mechanism, as illustrated in Fig. 2. UV/Vis-absorption analysis of individual components and the reaction mixture in DMA was performed, as shown in Fig. 2a. Neither silyl enol ether **1a** nor DABCO displayed an absorption band beyond 400 nm. However, the mixture of **1a** and **2a** exhibited a significant bathochromic shift (brown line). Additionally, the mixture of **2a** and DABCO (green line), as well as the reaction mixture of **1**, **2a**, and DABCO (purple line), displayed a notable bathochromic shift with visible light absorption in the range of 425–500 nm, indicating the formation of an EDA-complex. The intense yellow color observed in the reaction mixture further suggested the formation of an EDA-complex aggregate, as depicted in Fig. 2a.

Additionally, a molar absorption ratio of 1:1 for the mixture of **2a** and DABCO was demonstrated by Job's plot study (Fig. 2b). Subsequently, a Benesi-Hildebrand experiment¹⁵ was performed, yielding an association constant of 4.55 M^{-1} in DMA, which confirmed the formation of a visible light-active EDA complex (Fig. 2c). Next, a ¹H-NMR titration experiment between *S*-phenyl 4-methylbenzenesulfonothioate **2a** and DABCO in CDCl₃ revealed that the ¹H-proton signals of **2a** shifted upfield, thus indicating the formation of an EDA-complex between **2a** and DABCO (Fig. 2d). Thereafter, light-on and on/off experiments were performed, demonstrating that continuous visible-light irradiation is necessary for the reaction to proceed (Fig. 2e). Following this, a radical trapping experiment was performed under standard conditions using

TEMPO (2,2,6,6-tetramethyl-1-piperidinyloxy), butylated hydroxytoluene (BHT) and 1,1-diphenylethylene. The reactions were quenched, and the resulting radical trapped adducts were successfully detected by high-resolution mass spectrometry (HRMS), indicating the involvement of a radical pathway (Fig. 2f). Next, the reaction mixture was irradiated with natural sunlight, resulting in the formation of the desired product **3a** in 59% yield (Fig. 2g). The reaction was scaled up to the gram scale, achieving a 68% yield of the target product **3a** (Fig. 2h). Lastly, a quantum yield experiment was performed and the yield was found to be $\Phi = 0.82$, which indicates a closed-chain pathway (see the ESI† for detailed discussion).

Green chemistry metrics

To assess the environmental impact of our EDA strategy,¹⁶ we evaluated the green chemistry metrics for the synthesis of compound **3a** (0.0487 g, 89%), starting from **1a** (0.2 mmol, 0.0384 g) and *S*-phenyl 4-methylbenzenesulfonothioate **2a** (0.24 mmol, 0.0634 g), using DABCO (3.0 equiv., 0.0673 g) under visible light conditions. The results are depicted in Fig. 3a. Our method demonstrates outstanding green chemistry metrics. Specifically, the atom economy was calculated to be 96.6%, the reaction mass efficiency was 82.22%, and the process mass intensity was 7.65, all of which indicate a highly efficient process. Additionally, the *E*-factor was determined to be 6.7, the lowest among comparable methods. The EcoScale score was calculated to be 68 (see the ESI† for detailed calculation), which is acceptable and reflects a favorable balance of safety, economic, and ecological considerations. Overall, the



Scheme 3 Possible reaction mechanism.

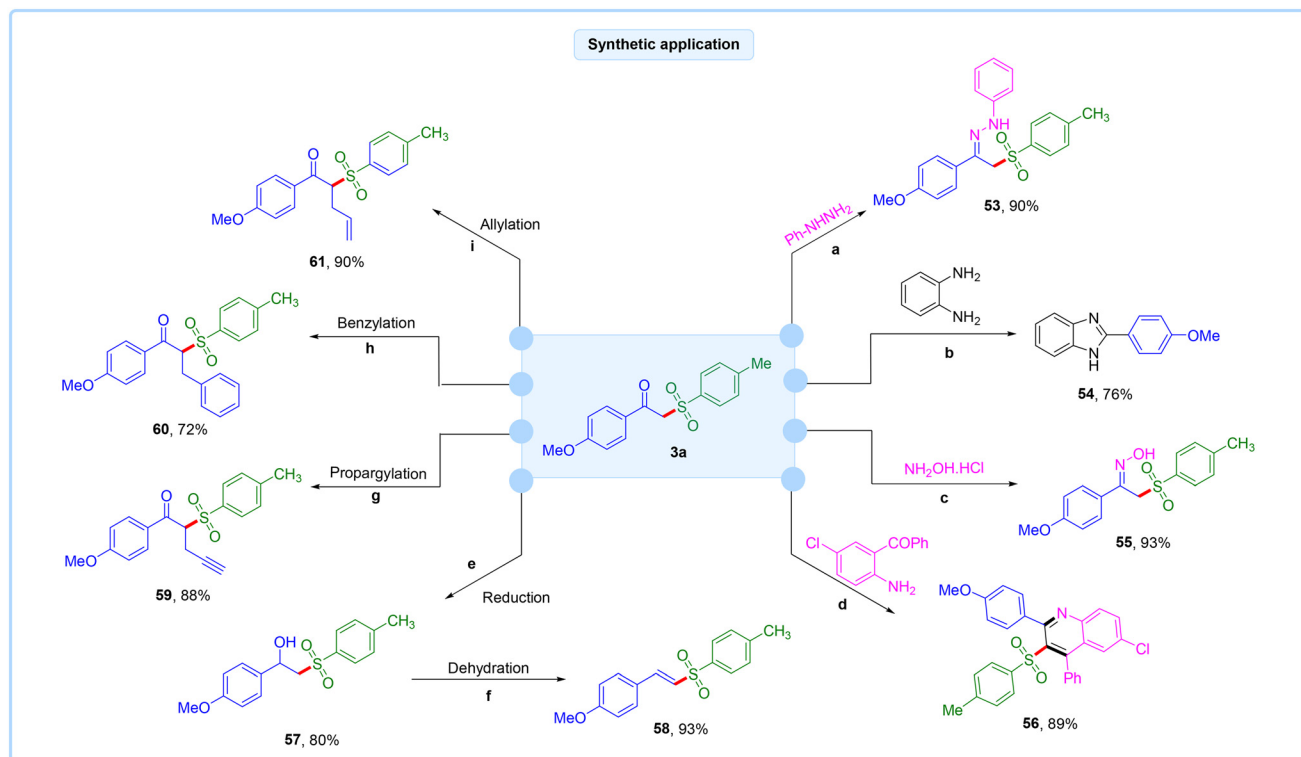


Fig. 4 Diversification of β -Keto sulfones. Reaction conditions: (a) **3a** (0.2 mmol), phenylhydrazine (1.0 equiv.), acetic acid (cat. amount), ethanol (1.0 ml), reflux, and 1 h; (b) **3a** (0.2 mmol), *o*-phenylenediamine (1 equiv.), AcOH, 120 °C, and 10 h; (c) **3a** (0.2 mmol) hydroxylamine hydrochloride (1.5 equiv.) and NaOAc (1.5 equiv.) ethanol (1 mL), water (3 ml), reflux, and 5 h; (d) **3a** (0.2 mmol), (2-amino-5-chlorophenyl)(phenyl)methanone (1.5 equiv.), TfOH (20 mol%), PhCl, 150 °C, and 10 h. (e) **3a** (0.2 mmol), NaBH₄ (1.5 equiv.), methanol (2.0 mL), 25 °C, and 1 h; (f) **57** (0.1 mmol), BF₃·Et₂O (0.2 mmol), DCM, 25 °C, and 20 h; (g) **3a** (0.2 mmol), K₂CO₃ (2.0 equiv.), propargylic bromide (1.1 equiv.), acetone (2.0 mL), reflux, and 12 h; (h) **3a** (0.2 mmol), K₂CO₃ (3.5 equiv.), BnBr (1.1 equiv.), acetone, and reflux; and (i) **3a** (0.2 mmol), K₂CO₃ (2.0 equiv.), allyl bromide (1.1 equiv.), acetone (2.0 mL), reflux, and 16 h.

green chemistry metrics of our method indicate a strong alignment with sustainability goals. The comparison between the green chemistry metrics of our strategy and other reports is shown in Fig. 3b (see the ESI† for detailed calculation).

Based on the experimental observations and prior literature,¹⁷ a plausible mechanism for this photochemical EDA-complex mediated transformation is depicted in Scheme 3. Initially, an EDA-complex aggregate is formed between **2** and DABCO. In the presence of visible light irradiation, this EDA complex undergoes a single-electron transfer (SET) event from DABCO to *S*-phenyl 4-methylbenzenesulfonylthioate **2**, generating the thiyl anion intermediate **Int-I**, the *S*-centered radical intermediate **Int-II**, and the DABCO radical cation. The generated sulfonyl radical intermediate **Int-II** reacts with silyl enol ether **1**, leading to the formation of the radical intermediate **Int-III**, which subsequently undergoes SET with the DABCO radical cation to generate the cation intermediate **Int-IV**. Lastly, tautomerization, followed by hydrolysis, yields the desired product **3**.

To broaden the synthetic application, various reactions were conducted using compound **3a**, as shown in Fig. 4. The reaction of **3a** with phenylhydrazine produced the required product **53** in 90% yield.¹⁸ Additionally, the reaction with *o*-phenylenediamine yielded the corresponding product **54** in

76% yield.¹⁹ Similarly, the reaction with hydroxylamine hydrochloride gave the oxime product **55** in 93% yield.¹⁸ Conversely, the reaction with (2-amino-5-chlorophenyl)(phenyl)methanone yielded a six-membered ring product **56** in 89% yield.²⁰ The carbonyl group of the β -ketosulfone was smoothly reduced to compound **57** using NaBH₄ in methanol.²¹ The resulting hydroxyl group was then dehydrated to produce vinyl sulfone **58**, with an approximate yield of 93%.²² Finally, we also demonstrated the propargylation,²³ benzylation,²⁴ and allylation²⁵ of compound **3a**, resulting in the corresponding products (**59–61**) in good to excellent yields.

Conclusion

In summary, we reported a transition metal- and photo-catalyst-free method for the visible light-induced synthesis of β -ketosulfones *via* the sulfonylation of ketone-derived silyl enol ethers with thiosulfonates (electron acceptor) and DABCO (electron donor) under mild conditions. This method facilitates the synthesis of a wide range of β -ketosulfone derivatives, including biologically active and late-stage molecules, from structurally diverse silyl enol ethers, including alkyl, aryl, and heteroarylsilyl enol ethers. Extensive mechanistic studies,

including UV-visible absorption studies, Job's plot, $^1\text{H-NMR}$ titration experiment, Benesi-Hildebrand plot, radical trapping experiment, light on/off experiment, and quantum yield experiment, confirm the involvement of an EDA complex-mediated radical process. Additionally, our method offers several advantages over the existing literature. This includes (i) being transition-metal and photoredox catalyst-free; (ii) being external SO_2 -source free; (iii) accommodating a diverse range of substrates; (iv) yielding recyclable and reusable by-products; and (v) demonstrating excellent atom economy, reaction mass efficiency, process mass intensity, *E*-factor, and EcoScale score, highlighting its efficiency and economic sustainability. Moreover, the versatility of β -ketosulfones was demonstrated by their successful conversion into various aromatic and heteroaromatic compounds, showcasing their broad utility as intermediates and making them a practical solution for academic research and industrial applications.

Author contributions

B. Saxena optimized the reaction conditions and synthesized all the derivatives. R. I. Patel synthesized the thiosulfonates. B. Saxena and R. I. Patel performed the mechanistic studies and wrote the manuscript with helpful insights from 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.†

Experimental data have been provided as the ESI.†

Conflicts of interest

The authors declare the following competing interests: one patent has been registered (no: 202411058425) in India.

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References

- (a) N. S. Simpkins, in *Sulfones in Organic Synthesis*, ed. J. E. Baldwin, Pergamon, Oxford, 1993; (b) B. M. Trost, *Comprehensive Organic Chemistry*, Pergamon, Oxford, 1991; (c) D. C. Meadows and S. Gervay-Hague, *Med. Res. Rev.*, 2006, **26**, 793–814; (d) A. El-Awa, M. N. Noshi, X. Mollat du Jourdin and P. L. Fuchs, *Chem. Rev.*, 2009, **109**, 2315–2349; (e) M. Nielsen, C. B. Jacobsen, N. Holub, M. W. Paixão and K. A. Jørgensen, *Angew. Chem., Int. Ed.*, 2010, **49**, 2668–2679; (f) A. N. R. Alba, X. Companyö and R. Rios, *Chem. Soc. Rev.*, 2010, **39**, 2018–2033.
- (a) R. E. Swenson, T. J. Sowin and H. Q. Zhang, *J. Org. Chem.*, 2002, **67**, 9182–9185; (b) J. Xiang, M. Ipek, V. Suri, W. Masefski, N. Pan, Y. Ge, M. Tam, Y. Xing, J. F. Tobin and X. Xu, *Bioorg. Med. Chem. Lett.*, 2005, **15**, 2865–2869; (c) C. Curti, M. Laget, A. O. Carle, A. Gellis and P. Vanelle, *Eur. J. Med. Chem.*, 2007, **42**, 880–884; (d) R. K. Saini, Y. C. Joshi and P. Joshi, *Heterocycl. Commun.*, 2007, **13**, 219–222; (e) X. Huang and J. Xue, *J. Org. Chem.*, 2007, **72**, 3965–3968; (f) J. Xiang, M. Ipek, V. Suri, M. Tam, Y. Xing, N. Huang, Y. Zhang, J. Tobin, T. S. Mansour and J. McKew, *Bioorg. Med. Chem.*, 2007, **15**, 4396–4405; (g) H. Yang, R. G. Carter and L. N. Zakharov, *J. Am. Chem. Soc.*, 2008, **130**, 9238–9239; (h) N. Pokhodylo, V. Matychuk and M. Obushak, *Synthesis*, 2009, 2321–2323; (i) Z. Ding, J. Yang, T. Wang, Z. Shen and Y. Zhang, *Chem. Commun.*, 2009, 571–573; (j) A. Kumar, S. Sharma, V. D. Tripathi and S. Srivastava, *Tetrahedron*, 2010, **66**, 9445–9449; (k) A. Kumar and M. K. Muthyala, *Tetrahedron Lett.*, 2011, **22**, 1287; (l) A. F. Eweas, G. Allam, A. S. A. Abuelsaad, A. H. AlGhamdi and I. A. Maghrabi, *Bioorg. Chem.*, 2013, **46**, 17–25.
- A. R. Katritzky, A. A. Abdel-Fattah and M. Y. Wang, *J. Org. Chem.*, 2003, **68**, 1443–1446.
- (a) B. M. Trost and D. P. Curran, *Tetrahedron Lett.*, 1981, **22**, 1287–1290; (b) C. R. Holmquist and E. J. Roskamp, *Tetrahedron Lett.*, 1992, **33**, 1131–1134; (c) C. Lai, C. Xi, Y. Jiang and R. Hua, *Tetrahedron Lett.*, 2005, **46**, 513–515; (d) T. Zweifel, M. Nielsen, J. Overgaard, C. B. Jacobsen and K. A. Jørgensen, *Eur. J. Org. Chem.*, 2011, 47–52; (e) N. Samakkanad, P. Katrun, T. Techajaronjit, S. Hlekhilai, M. Pohmakotr, V. Reutrakul, T. Jaipetch, D. Soorukram and C. Kuhakarn, *Synthesis*, 2012, 1693–1699.
- (a) G. E. Vennstra and B. Zwaneburg, *Synthesis*, 1975, 519–520; (b) Y.-Y. Xie and Z.-C. Chen, *Synth. Commun.*, 2001, **31**, 3145–3149.
- (a) W. Wei, C.-L. Liu, D.-S. Yang, J.-W. Wen, J.-M. You, Y.-R. Suo and H. Wang, *Chem. Commun.*, 2013, **49**, 10239–10241; (b) Y. Jiang and T.-P. Loh, *Chem. Sci.*, 2014, **5**, 4939–4943; (c) H. Wang, Q. Lu, C. Qian, C. Liu, W. Liu, K. Chen and A. Lei, *Angew. Chem., Int. Ed.*, 2016, **55**, 1094–1097; (d) D. Yang, B. Huang, W. Wei, J. Li, Y. Liu, J. Ding, P. Sun and H. Wang, *Green Chem.*, 2016, **18**, 5630–5634; (e) F. Chen, N.-N. Zhou, J.-L. Zhan, B. Han and W. Yu, *Org. Chem. Front.*, 2017, **4**, 135–139; (f) A. Shao, M. Gao, S. Chen, T. Wang and A. Lei, *Chem. Sci.*, 2017, **8**, 2175–2178; (g) X. Yang, J. Yang, K. Yan, H. Qin, W. Dong, J. Wen and H. Wang, *Eur. J. Org. Chem.*, 2020, 3456–3461;

- (h) T.-F. Niu, J. Cheng, C.-L. Zhuo, D.-Y. Jiang, X.-G. Shu and B.-Q. Ni, *Tetrahedron Lett.*, 2017, **58**, 3667–3671; (i) Q. Liu, F. Liu, H. Yue, X. Zhao, J. Li and W. Wei, *Adv. Synth. Catal.*, 2019, **361**, 5277–5282; (j) J. Wen, X. Yang, Z. Sun, J. Yang, P. Han, Q. Liu, H. Dong, M. Gu, L. Huang and H. Wang, *Green Chem.*, 2020, **22**, 230–237; (k) J. Wu, Y. Zhang, X. Gong, Y. Menga and C. Zhu, *Org. Biomol. Chem.*, 2019, **17**, 3507–3513; (l) Z. Peng, Y.-Y. Hong, S. Peng, X.-Q. Xu, S.-S. Tang, L.-H. Yang and L.-Y. Xie, *Org. Biomol. Chem.*, 2021, **19**, 4537–4541; (m) Y. Fang, D. Xu, Y. Yu, R. Tang, S. Dai, Z. Wang and W. Zhang, *Eur. J. Org. Chem.*, 2022, e202200091; (n) W.-H. Rao, C. Gao, L.-L. Jiang, F.-Y. Zhou, J.-F. Liu and G.-D. Zou, *J. Org. Chem.*, 2024, **89**, 12681–12692; (o) Y. Xia, X. Chen, L. Qu, K. Sun, X. Xia, W. Fu, X. Chen, Y. Yang, Y. Zhao and C. Li, *Asian J. Org. Chem.*, 2016, **5**, 878–881; (p) R. Chen, Y. Tang, X. He, K.-K. Wang, L. Ding and L. Liu, *Org. Lett.*, 2023, **25**, 5454–5458.
- 7 (a) Q.-Q. Lu, J. Zhang, G.-L. Zhao, Y. Qi, H.-M. Wang and A.-W. Lei, *J. Am. Chem. Soc.*, 2013, **135**, 11481–11484; (b) S. Handa, J. C. Fennewald and B. H. Lipshutz, *Angew. Chem., Int. Ed.*, 2014, **53**, 3432–3435; (c) A. K. Singh, R. Chawla and L. D. S. Yadav, *Tetrahedron Lett.*, 2014, **55**, 2845–2848; (d) R. Chawla, A. K. Singh and L. D. S. Yadav, *Eur. J. Org. Chem.*, 2014, 2032–2036; (e) A. K. Singh, R. Chawla and L. D. S. Yadav, *Tetrahedron Lett.*, 2014, **55**, 4742–4746; (f) X. Shi, X. Ren, Z. Ren, J. Li, Y. Wang, S. Yang, J. Gu, Q. Gao and G. Huang, *Eur. J. Org. Chem.*, 2014, 5083–5088; (g) N. Kumar and A. Kumar, *ACS Sustainable Chem. Eng.*, 2019, **7**, 9182–9188; (h) X. Chen, S. Lu, Y. Zheng, J. Wang, L. Yang and P. Sun, *Adv. Synth. Catal.*, 2022, **364**, 1305–1312; (i) S.-W. Yu, Z.-J. Chen, H.-Q. Li, W.-X. Li, Y. Li, Z. Li and Z.-Y. Wang, *Molecules*, 2024, **29**, 3559; (j) H.-L. Wan, Z. Guan and Y.-H. He, *Asian J. Org. Chem.*, 2021, **10**, 3406–3410; (k) V. K. K. Pampana, V. P. Charpe, A. Sagadevan, D. K. Das, C.-C. Lin, J. R. Hwu and K. C. Hwang, *Green Chem.*, 2021, **23**, 3569–3574; (l) T. Song, Y. Zhang, C. Wang, Y. Li and Y. Yang, *Chin. J. Chem.*, 2022, **40**, 2618–2624; (m) S. Cai, D. Chen, Y. Xu, W. Weng, L. Li, R. Zhang and M. Huang, *Org. Biomol. Chem.*, 2016, **14**, 4205–4209; (n) B. Dam, A. K. Sahoo and B. K. Patel, *Green Chem.*, 2022, **24**, 7122–7130; (o) I. Yavari and S. Shaabanzadeh, *Org. Lett.*, 2020, **22**, 464–467; (p) X. Chen, X. Chang, S. Zhang, S. Lu, L. Yang and P. Sun, *Environ. Chem. Lett.*, 2022, **20**, 2773–2779.
- 8 (a) Y. Tang, Y. Zhang, K. Wang, X. Li, X. Xu and X. Du, *Org. Biomol. Chem.*, 2015, **13**, 7084; (b) H. Wang, G. Wang, Q. Lu, C.-W. Chiang, P. Peng, J. Zhou and A. Lei, *Chem. – Eur. J.*, 2016, **22**, 14489; (c) T. Liu, D. Zheng, Y. Ding, X. Fan and J. Wu, *Chem. – Asian J.*, 2017, **12**, 465; (d) W. Chen, X. Liu, E. Chen, B. Chen, J. Shao and Y. Yu, *Org. Chem. Front.*, 2017, **4**, 1162; (e) H. Jiang, Y. Cheng, Y. Zhang and S. Yu, *Eur. J. Org. Chem.*, 2013, 5485–5492.
- 9 For recent reviews on photoredox reaction, see: (a) D. Ravelli, M. Fagnoni and A. Albini, *Chem. Soc. Rev.*, 2013, **42**, 97; (b) Y. Xi, H. Yi and A. Lei, *Org. Biomol. Chem.*, 2013, **11**, 2387; (c) C. K. Prier, D. A. Rankic and D. W. C. MacMillan, *Chem. Rev.*, 2013, **113**, 5322; (d) D. M. Schultz and T. P. Yoon, *Science*, 2014, **343**, 985; (e) J. Xie, H. Jin, P. Xu and C. Zhu, *Tetrahedron Lett.*, 2014, **55**, 36; (f) E. Meggers, *Chem. Commun.*, 2015, **51**, 3290; (g) J. J. Douglas, M. J. Sevrin and C. R. J. Stephenson, *Org. Process Res. Dev.*, 2016, **20**, 1134; (h) I. Ghosh, L. Marzo, A. Das, R. Shaikh and B. König, *Acc. Chem. Res.*, 2016, **49**, 1566; (i) B. Saxena, R. I. Patel and A. Sharma, *RSC Sustainability*, 2024, **2**, 2169–2189; (j) B. Saxena, R. I. Patel, J. Tripathi and A. Sharma, *Org. Biomol. Chem.*, 2023, **21**, 4723–4743; (k) J. Singh, B. Saxena and A. Sharma, *Catal. Sci. Technol.*, 2024, **14**, 5143–5160; (l) R. I. Patel, S. Sharma and A. Sharma, *Org. Chem. Front.*, 2021, **8**, 3166–3200; (m) R. I. Patel, A. Sharma, S. Sharma and A. Sharma, *Org. Chem. Front.*, 2021, **8**, 1694–1718; (n) R. I. Patel, J. Singh and A. Sharma, *ChemCatChem*, 2022, **14**, e202200260.
- 10 X. Wang, Y. Kuang, S. Ye and J. Wu, *Chem. Commun.*, 2019, **55**, 14962–14964.
- 11 F.-S. He, Y. Yao, W. Xie and J. Wu, *Chem. Commun.*, 2020, **56**, 9469–9472.
- 12 F.-S. He, P. Bao, Z. Tang, F. Yu, W.-P. Deng and J. Wu, *Org. Lett.*, 2022, **24**, 2955–2960.
- 13 (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; (g) Z.-Y. Cao, T. Ghosh and P. Melchiorre, *Nat. Commun.*, 2018, **9**, 3274.
- 14 (a) R. I. Patel, B. Saxena and A. Sharma, *Green Chem.*, 2024, **26**, 10265–10274; (b) B. Saxena, R. I. Patel and A. Sharma, *Adv. Synth. Catal.*, 2023, **365**, 1538–1564; (c) A. Sharma, N. Singh, R. Kumar and A. Sharma, *Adv. Synth. Catal.*, 2024, **366**, 2735–2741; (d) N. Singh, S. Sharma and A. Sharma, *Adv. Synth. Catal.*, 2023, **365**, 3505–3511; (e) A. Monga, S. Bagchi, R. K. Soni and A. Sharma, *Adv. Synth. Catal.*, 2020, **362**, 2232–2237.
- 15 H. A. Benesi and J. Hildebrand, *J. Am. Chem. Soc.*, 1949, **71**, 2703–2707.
- 16 (a) N. Mukherjee and T. Chatterjee, *Green Chem.*, 2021, **23**, 10006–10013; (b) A. N. V. Satyanarayana, N. Mukherjee and T. Chatterjee, *Green Chem.*, 2023, **25**, 779–788; (c) K. Van Aken, L. Strekowski and L. Patiny, *Beilstein J. Org. Chem.*, 2006, **2**, 3; (d) A. Beillard, X. Bantreil, T.-X. Métro, J. Martinez and F. Lamaty, *Green Chem.*, 2018, **20**, 964–968; (e) B. Saxena, R. I. Patel, S. Sharma and A. Sharma, *Green Chem.*, 2024, **26**, 2721–2729.
- 17 (a) J. Tong, H. Li, Y. Zhu, P. Liu and P. Sun, *Green Chem.*, 2022, **24**, 1995–1999; (b) K. Cao, N. Zhang, L. Lin, Q. Shen, H. Jiang and J. Li, *Adv. Synth. Catal.*, 2024, **366**, 207–213;

- (c) K. Sun, A. Shi, Y. Liu, X. Chen, P. Xiang, X. Wang, L. Qua and B. Yu, *Chem. Sci.*, 2022, **13**, 5659–5666; (d) D.-L. Zhang, Z.-G. Le, Q. Li, Z.-B. Xie, W.-W. Yang and Z.-Q. Zhu, *Chem. Commun.*, 2024, **60**, 2958–2961.
- 18 H. A. A. Aziz, K. A. A. Rashood, K. H. El-Tahir and G. M. Suddek, *Eur. J. Med. Chem.*, 2014, **80**, 416–422.
- 19 M.-Y. Chang, C.-K. Chan and Y.-C. Chen, *Heterocycles*, 2014, **89**, 1237.
- 20 C.-K. Chan, C.-Y. Lai, W.-C. Lo, Y.-T. Cheng, M.-Y. Chang and C.-C. Wang, *Org. Biomol. Chem.*, 2020, **18**, 305–315.
- 21 Y.-S. Lin, Y.-C. Kuo, C.-H. Kuei and M.-Y. Chang, *Tetrahedron*, 2017, **73**, 1275–1282.
- 22 Y. Wang, Y. Zhao, C. Cai, L. Wang and H. Gong, *Org. Lett.*, 2021, **23**, 8296–8301.
- 23 M.-Y. Chang, Y.-C. Cheng and Y.-J. Lu, *Org. Lett.*, 2015, **17**, 1264–1267.
- 24 M.-Y. Chang, Y.-C. Chen and C.-K. Chan, *Synlett*, 2014, 1739–1744.
- 25 M.-Y. Chang, Y.-C. Cheng and Y.-J. Lu, *Org. Lett.*, 2014, **16**, 6252–6255.