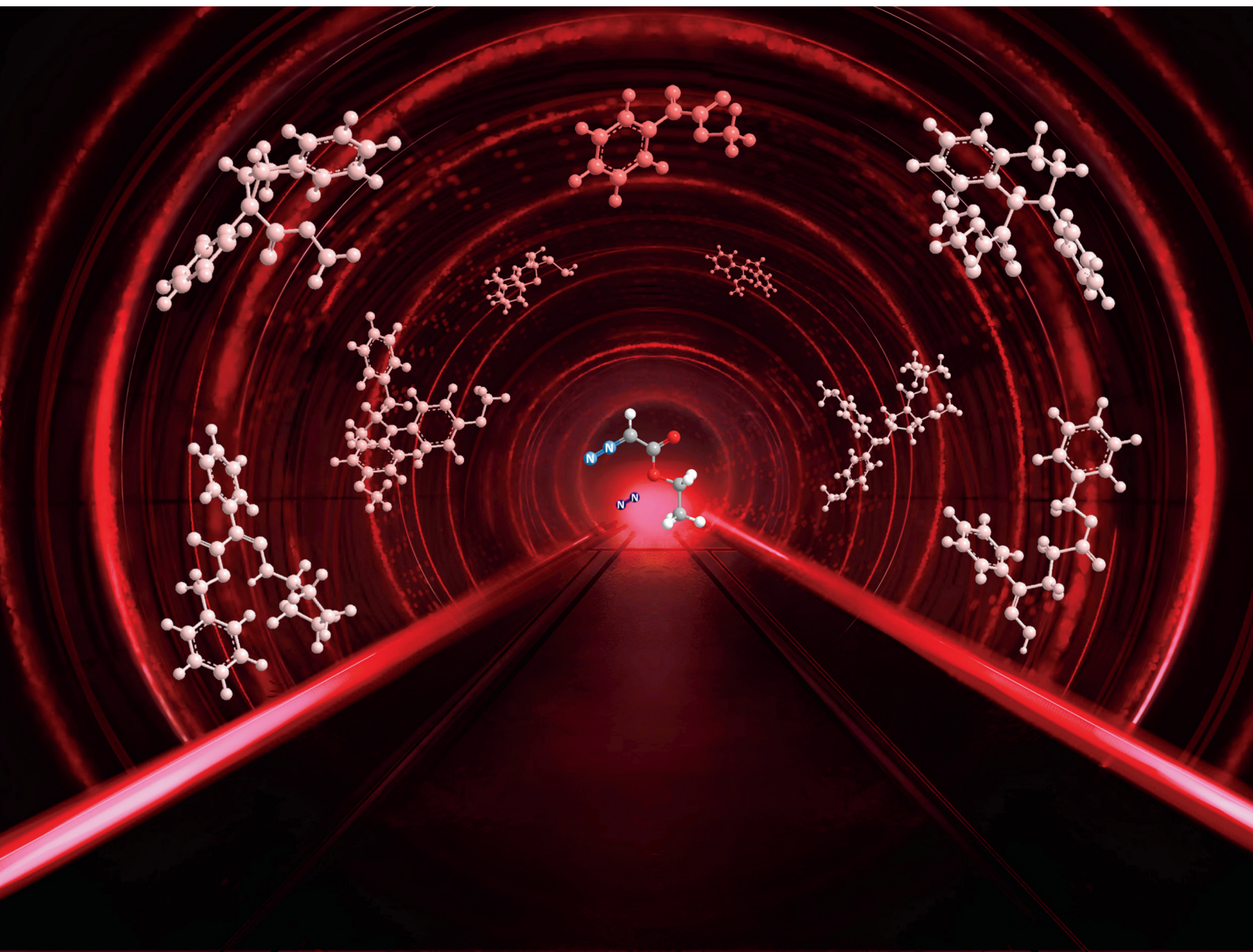


ChemComm

Chemical Communications

rsc.li/chemcomm



ISSN 1359-7345

COMMUNICATION

Katarzyna Rybicka-Jasińska, Dorota Gryko *et al.*
Unlocking the reactivity of diazo compounds in red light
with the use of photochemical tools


 Cite this: *Chem. Commun.*, 2023, 59, 14649

 Received 20th October 2023,
 Accepted 15th November 2023

DOI: 10.1039/d3cc05174a

rsc.li/chemcomm

Structurally diversified diazoalkanes can be activated under red light irradiation relying on direct photolysis, photosensitization or photoredox catalysis.

Bioorthogonal chemistry represents chemical transformations that proceed selectively in biological environments without perturbing the structure of biomolecules or interfering with biochemical pathways.^{1–6} Several photoactivated methods have been designed, yet most of them employ short-wavelength light emitting sources.^{7–10} The phototoxicity of high energetic photons makes them inappropriate for biological applications. Therefore, switching to less energetic light is desirable. Along this line, tetrazole bioorthogonal chemistry was performed under NIR radiation *via* two-photon excitation or upconversion processes.^{11,12} Dihydropyridazine oxidation *in vivo* can also be achieved with less energetic photons.¹³ Yet, red light-induced reactions even in synthetic chemistry call for in depth studies.¹⁴

Diazoalkanes are versatile reactants for photochemical synthesis of small/complex structures^{15–17} and functionalization of bioactive compounds.^{18–21} They have been utilized in enzymatic cyclopropanation, ring expansion, cyclopropanation, or insertion reactions.^{22–27} So far, however, generation of carbenes in biological systems is mostly limited to diazirines that are activated in UV/violet light.^{28–30} In view of benefits arising from the application of low energetic photons, red light-induced diazo chemistry is highly desirable. Given the structural diversity of diazoalkanes, they can be directly photolyzed or activated *via* photocatalytic processes under visible light (even red, Fig. 1A). We wondered whether it is possible to unlock the potential of red light toward the generation of reactive species from structurally diversified diazo compounds utilizing various photochemical modes. While studying the photocatalytic activity of porphyrins under red-light irradiation,

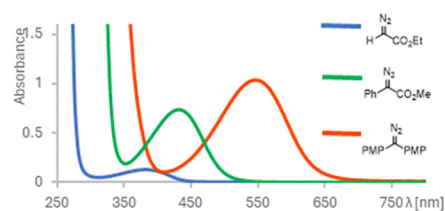
Unlocking the reactivity of diazo compounds in red light with the use of photochemical tools†

 Katarzyna Orłowska, Klaudia Łuczak, Piotr Krajewski,^{ib} João V. Santiago, Katarzyna Rybicka-Jasińska* and Dorota Gryko^{ib}*

we found that they catalyze photoalkylation of aldehydes with ethyl diazoacetate.³¹ Herein, we present our comprehensive study on the red light-induced photolysis, photosensitization, and photoredox-driven generation of reactive intermediates from diazo reagents (Fig. 1B).

Photolysis – Direct photolysis of diazoalkanes enables carbene generation with no catalyst required. Although acceptor-only and acceptor/acceptor diazo compounds exhibit light absorption beyond the visible range, replacing H/one of the acceptor groups with an aryl substituent bathochromically shifts the λ_{max} toward the visible spectrum.^{16,32} By increasing the donating character of the phenyl ring, λ_{max} is shifted even further (for –OMe, λ_{max} = 543 nm),³³ and has an impact on the carbene spin state. Given the ubiquity of free hydroxy-, amino-, and thio-groups in natural compounds, we focused on red light-induced photolysis of diaryldiazoalkanes in the presence of alcohols, amines, and thiols (Scheme 1). The light-induced method works well for primary alcohols efficiently affording ethers 1–5. Incorporation into the phenolic O–H bond, a

A. UV/Vis spectra of diazo compounds



B. Our work

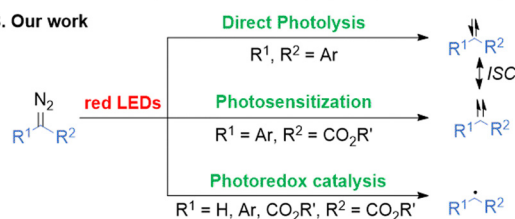


Fig. 1 Red light-induced reactions of diazoalkanes.

 Institute of Organic Chemistry, Polish Academy of Sciences,
 Kasprzaka 44/52 01-224, Warsaw, Poland. E-mail: dorota.gryko@icho.edu.pl

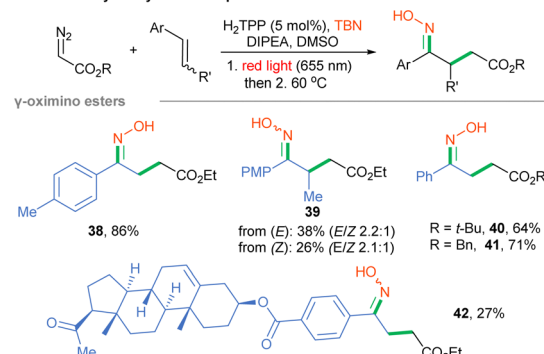
 † Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3cc05174a>


various aryldiazoesters leading to products **27–30**. Electron-poor aryldiazo ester reached the highest cyclopropanation productivity (**34**, 90%). The method is suitable for both electron-rich and -poor styrenes, with a better outcome for *p*-methoxy-styrene-derived product **35** (70%). A modest yield was observed when the internal olefin was subjected to the reaction conditions giving cyclopropane **37** (55%). α -Diazo esters, diazo-malonates, and aryldiazoketones possessing higher E_T values than porphyrin (calculated $E_T = 158 \text{ kJ mol}^{-1}$ for EDA) cannot, in principle, be activated under the developed conditions. Intuitively, the reaction rate for diazoalkane transformation depends on the carbene rate formation, which for the red light-mediated EnT-approach occurs slower than *via* direct photolysis under blue light (see ESI†).

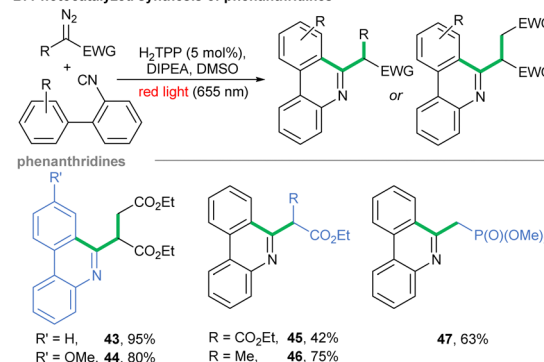
Photoredox catalysis – To unlock the red-light mediated reactivity of yet unconquered α -diazo esters, we screened the possibilities offered by photoredox catalysis. These acceptor-only types of diazoalkanes are reduced to alkyl radicals *via* proton coupled electron transfer (PCET, $E_{\text{RED}} = -1.28 \text{ V vs. SCE}$ for EDA).⁴³ In this view, numerous blue light-induced methodologies utilizing diazoesters as surrogates of alkyl radicals have been reported.^{43–48} Recently, we have proved that porphyrins are suitable photo-oxidants and photo-reductants for red light-mediated organic transformations.³¹ Therefore, we harnessed their photoredox abilities to tune already reported blue light-induced, radical-based transformations of α -diazoesters and applied them on red illumination instead. Our studies were initiated with the redesign of the photocatalyzed synthesis of γ -oximino esters, originally performed by Li under blue light with the use of α -diazoester, styrene and TBN as starting materials.⁴³ Optimization studies substantially shortened the reaction time (reported on blue: 60 h) to 37 h by thermally accelerating the isomerization of the nitroso compound to the final product **38** (see ESI†). Our method works well for various α -diazoesters giving esters **38**, **40** and **41** in yields comparable to those reported by Li (Scheme 3A). A slight yield decrease was observed for *trans*-anethole, though with a similar *E/Z* ratio (product **39**). Due to solubility problems, the synthesis of pregnenolone-derived ester **42** was less efficient. For the Ru-catalyzed reaction a key step relies on the reduction of diazo ester by the photocatalyst in the excited state. In our case, as the reduction potential of the porphyrin in the excited state (-0.91 vs. SCE)³¹ is higher than that of EDA (-1.28 V vs. SCE), we assume that the excited porphyrin oxidizes DIPEA, thus generating a strongly reducing porphyrin radical anion, similar to the mechanism reported for the generation of radicals from aminopyridinium salts.⁴⁹

Next, we examined an analogous PCET-based approach toward phenanthridines utilizing isocyanobiphenyls and diazoalkanes.⁵⁰ Scheme 3B shows the optimization of the red light-mediated protocol-enabled synthesis of heterocycles **43–47** with better productivity or comparable to the Xuan methodology. Finally, there are methodologies involving diazo reagents in which the diazo moiety remains intact or does not generate reactive intermediates. To fill the picture of the photochemistry of diazo compounds under red-light

A. Photocatalyzed synthesis of γ -oximino esters



B. Photocatalyzed synthesis of phenanthridines

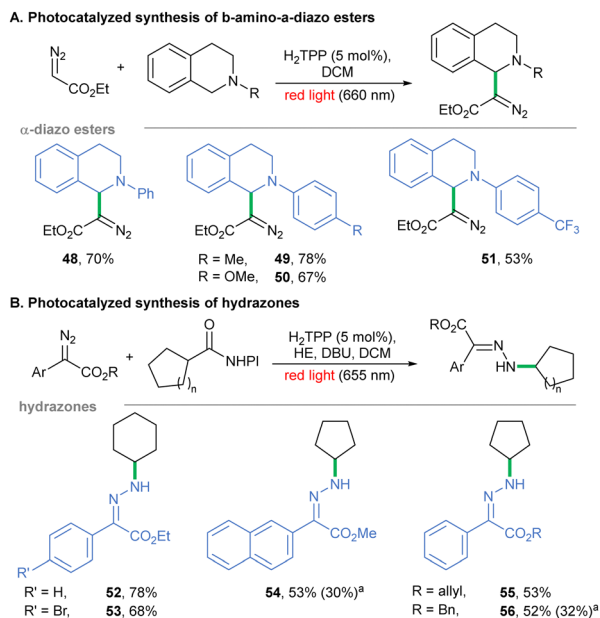


Scheme 3 Diazo compounds as radical precursors in red light-mediated photocatalyzed transformations.

irradiation, such transformations were studied. Given that sole H_2TPP is unable to photoreduce EDA, we tested H_2TPP as a photo-oxidant of diversely substituted tetrahydroisoquinolines in the presence of EDA and red light, similar to Zhou's report.⁵¹ In fact, products **48–51** were obtained in decent yields (Scheme 4A). Furthermore, diazo compounds have been shown to react with radicals generated under photochemical conditions, including alkyl radicals generated from NHPI esters in the presence of Rose Bengal on yellow LEDs.⁵² We performed this transformation with the H_2TPP catalyst instead, under red light irradiation. A wide range of donor/acceptor diazoalkanes reacted under the developed conditions to give hydrazones **52–56** (Scheme 4B).

In summary, this study demonstrates that photochemistry provides tools for red light-driven activation of various diazo compounds. A proper structural modification of diazoalkane results in a bathochromic shift of the absorption maxima allowing for direct photolysis under low-energetic, red-light irradiation. If this pathway is not possible, we induce transformations of diazo compounds taking advantage of nature-inspired dyes, established as safe and effective for photodynamic therapy and artificial photosynthesis. The triplet energy level of the porphyrin excited state is sufficient for productive EnT to aryl-diazo esters giving access to triplet carbenes. Other diazoalkanes may be activated through porphyrin-mediated photoredox processes by undergoing reduction to alkyl radicals or by serving as radical acceptors. Therefore, three-modes of





Scheme 4 Diazo compounds as radical acceptors in red light-mediated photocatalyzed transformations. ^aReaction set under blue light irradiation (25 W, 455 nm).

activation of diazo compounds under red-light irradiation have been unlocked.

Financial support for this work was provided by National Science Center, Poland, grant number Maestro UMO-2020/38/A/ST4/00185.

Conflicts of interest

There are no conflicts to declare.

Notes and references

- E. M. Sletten and C. R. Bertozzi, *Angew. Chem., Int. Ed.*, 2009, **48**, 6974.
- C. R. Bertozzi, *Acc. Chem. Res.*, 2011, **44**, 651.
- N. K. Devaraj, *ACS Cent. Sci.*, 2018, **4**, 952.
- S. Jia and E. M. Sletten, *ACS Chem. Biol.*, 2022, **17**, 3255.
- J. Li, H. Kong, C. Zhu and Y. Zhang, *Chem. Sci.*, 2020, **11**, 3390.
- B. D. Fairbanks, L. J. Macdougall, S. Mavila, J. Sinha, B. J. Kirkpatrick, K. S. Anseth and C. N. Bowman, *Chem. Rev.*, 2021, **121**, 6915.
- G. S. Kumar and Q. Lin, *Chem. Rev.*, 2021, **121**, 6991.
- W. Song, Y. Wang, J. Qu, M. M. Madden and Q. Lin, *Angew. Chem., Int. Ed.*, 2008, **47**, 2832.
- W. Song, Y. Wang, J. Qu and Q. Lin, *J. Am. Chem. Soc.*, 2008, **130**, 9654.
- P. An, T. M. Lewandowski, T. G. Erbay, P. Liu and Q. Lin, *J. Am. Chem. Soc.*, 2018, **140**, 4860.
- Z. Yu, T. Y. Ohulchanskyy, P. An, P. N. Prasad and Q. Lin, *J. Am. Chem. Soc.*, 2013, **135**, 16766.
- P. Lederhose, Z. Chen, R. Muller, J. P. Blinco, S. Wu and C. Barner-Kowollik, *Angew. Chem., Int. Ed.*, 2016, **55**, 12195.
- C. Wang, H. Zhang, T. Zhang, X. Zou, H. Wang, J. E. Rosenberger, R. Vannam, W. S. Trout, J. B. Grimm, L. D. Lavis, C. Thorpe, X. Jia, Z. Li and J. M. Fox, *J. Am. Chem. Soc.*, 2021, **143**, 10793.

- N. Sellet, M. Cormier and J. P. Goddard, *Org. Chem. Front.*, 2021, **8**, 6783.
- Z. Yang, M. L. Stivanin, I. D. Jurberg and R. M. Koenigs, *Chem. Soc. Rev.*, 2020, **49**, 6833.
- J. Durka, J. Turkowska and D. Gryko, *ACS Sustainable Chem. Eng.*, 2021, **9**, 8895.
- L. W. Ciszewski, K. Rybicka-Jasińska and D. Gryko, *Org. Biomol. Chem.*, 2019, **17**, 432.
- H. M. L. Davies and J. R. Denton, *Chem. Soc. Rev.*, 2009, **38**, 3061.
- S.-S. Ge, B. Chen, Y.-Y. Wu, Q.-S. Long, Y.-L. Zhao, P.-Y. Wang and S. Yang, *RSC Adv.*, 2018, **8**, 29428.
- S. Harada, *Chem. Pharm. Bull.*, 2021, **69**, 1170.
- K. A. Mix, M. R. Aronoff and R. T. Raines, *ACS Chem. Biol.*, 2016, **11**, 3233.
- Z. Liu and F. H. Arnold, *Curr. Opin. Biotechnol.*, 2021, **69**, 43.
- B. J. Wittmann, A. M. Knight, J. L. Hofstra, S. E. Reisman, S. B. J. Kan and F. H. Arnold, *ACS Catal.*, 2020, **10**, 7112.
- R. Mao, D. J. Wackelin, C. S. Jamieson, T. Rogge, S. Gao, A. Das, D. M. Taylor, K. N. Houk and F. H. Arnold, *J. Am. Chem. Soc.*, 2023, **145**, 16176.
- D. C. Miller, R. G. Lal, L. A. Marchetti and F. H. Arnold, *J. Am. Chem. Soc.*, 2022, **144**, 4739.
- K. Chen, X. Huang, S. B. Jennifer Kan, R. K. Zhang and F. H. Arnold, *Science*, 2018, **360**, 71.
- R. K. Zhang, K. Chen, X. Huang, L. Wohlschlagler, H. Renata and F. H. Arnold, *Nature*, 2019, **565**, 67.
- A. Blencow and W. Hayes, *Soft Matter*, 2005, **1**, 178.
- S. W. Huth, J. V. Oakley, C. P. Seath, J. B. Geri, A. D. Trowbridge, D. L. Parker Jr., F. P. Rodriguez-Rivera, A. G. Schwaib, C. Ramil, K. A. Ryu, C. H. White, O. O. Fadeyi, R. C. Oslund and D. W. C. MacMillan, *J. Am. Chem. Soc.*, 2023, **145**, 16289.
- J. B. Geri, J. V. Oakley, T. Reyes-Robles, T. Wang, S. J. McCarver, C. H. White, F. P. Rodriguez-Rivera, D. L. Parker Jr., E. C. Hett, O. O. Fadeyi, R. C. Oslund and D. W. C. MacMillan, *Science*, 2020, **367**, 1091.
- K. Rybicka-Jasińska, T. Wdowik, K. Łuczak, A. J. Wierzbka, O. Drapała and D. Gryko, *ACS Org. Inorg. Au*, 2022, **2**, 422.
- I. D. Jurberg and H. M. L. Davies, *Chem. Sci.*, 2018, **9**, 5112.
- S. Jana, C. Pei, C. Empel and R. M. Koenigs, *Angew. Chem., Int. Ed.*, 2021, **60**, 13271.
- R. W. R. Humphreys and D. R. Arnold, *Can. J. Chem.*, 1979, **57**, 2652.
- F. Strieth-Kalthoff and F. Glorius, *Chem*, 2020, **6**, 1888.
- F. Strieth-Kalthoff, M. J. James, M. Teders, L. Pitzer and F. Glorius, *Chem. Soc. Rev.*, 2018, **47**, 7190.
- K. Orłowska, J. V. Santiago, P. Krajewski, K. Kisiel, I. Deperasińska, K. Zawada, W. Chaładaj and D. Gryko, *ACS Catal.*, 2023, **13**, 1964.
- R. Costa e Silva, L. O. da Silva, A. de Andrade Bartolomeu, T. J. Brocksom and K. T. de Oliveira, *Beilstein J. Org. Chem.*, 2020, **16**, 917.
- J. Kou, D. Dou and L. Yang, *Oncotarget*, 2017, **8**, 81591.
- D. Gust, T. A. Moore and A. L. Moore, *Acc. Chem. Res.*, 2009, **42**, 1890.
- R. Seely, *Photochem. Photobiol.*, 1978, **27**, 639.
- L. G. Arnaut, Design of porphyrin-based photosensitizers for photodynamic therapy, in *Advances in Inorganic Chemistry*, ed. R. van Eldik and G. Stoichel, Academic Press, 2011, ch. 5, vol. 63, pp. 187–233.
- Y. Liu, K. Zhu, J. Zhao and P. Li, *Org. Lett.*, 2022, **24**, 6834.
- F. Li, S. Zhu and R. M. Koenigs, *Chem. Commun.*, 2022, **58**, 7526.
- L. W. Ciszewski, J. Durka and D. Gryko, *Org. Lett.*, 2019, **21**, 7028.
- X. Huang, R. D. Webster, K. Harms and E. Meggers, *J. Am. Chem. Soc.*, 2016, **138**, 12636.
- Y.-L. Su, G. X. Liu, L. De Angelis, R. He, A. Al-Sayyed, K. S. Schanze, W.-H. Hu, H. Qiu and M. P. Doyle, *ACS Catal.*, 2022, **12**, 1357.
- Y. L. Su, G. X. Liu, J. W. Liu, L. Tram, H. Qiu and M. P. Doyle, *J. Am. Chem. Soc.*, 2020, **142**, 13846.
- K. Golszewska, K. Rybicka-Jasińska, J. A. Clark, V. I. Vullev and D. Gryko, *ACS Catal.*, 2020, **10**, 5920.
- H. B. Ye, X. Y. Zhou, L. Li, X. K. He and J. Xuan, *Org. Lett.*, 2022, **24**, 6018.
- T. Xiao, L. Li, G. Lin, Z. W. Mao and L. Zhou, *Org. Lett.*, 2014, **16**, 4232.
- C. M. Chan, Q. Xing, Y. C. Chow, S. F. Hung and W. Y. Yu, *Org. Lett.*, 2019, **21**, 8037.

