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With or without a co-solvent? highly efficient ultrafast phenanthrenequinone-electron rich alkene (PQ-ERA) photoclick reactions†

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The light-induced photocycloaddition of 9,10-phenanthrenequinone (PQ) with electron-rich alkenes (ERA), known as the PQ-ERA reaction, is a highly attractive photoclick reaction characterized by its operational simplicity and high biocompatibility. One essential aspect of photoclick reactions is their high rate, however the limited solubility of PQs often requires the use of a co-solvent. Evaluating the effect of different co-solvents on the PQ-ERA reaction and their influence on the reaction rate, we discovered that sulfur-containing compounds, in particular the frequently used solubilizing co-solvent DMSO, quench the triplet state of the PQ. These experimental results, supported by nanosecond-microsecond and ultrafast transient absorption data, show that even minimal amounts of DMSO result in a decreased lifetime of the reactive triplet state, essential for the photoclick reaction. Without DMSO as co-solvent, exceptionally high photoreaction quantum yields (Φ_P up to 93% with only 1 equivalent ERA) and complete conversion in seconds can be achieved. With these outstanding efficiencies, the PQ-ERA reaction can be used without excess ERA and at low light intensities, facilitating photoclick transformations in various future applications.

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1 Introduction

The use of photochemical activation procedures in click chemistry has led to the development of a new category of light-

induced reactions, known as “photoclick chemistry”.^{1–4} Besides the advantageous properties of conventional click chemistry (operationally simple, fast, high yielding and selective), these reactions offer a high degree of spatial resolution and temporal control.^{5–10} Therefore, photoclick reactions are excellent transformations for diverse applications ranging from surface functionalization, polymer conjugation and photo-crosslinking to protein labelling and bioimaging.^{11–14}

The uniquely high efficiency of photoclick reactions is exemplified by the [4 + 2] cycloaddition between 9,10-phenanthrenequinones (PQs) and electron rich alkenes (ERAs).^{14–17} This reaction was first reported in 1944 by Schönberg,¹⁵ but was recently re-introduced by Zhang and coworkers.¹⁷ Taking advantage of ERAs (vinyl ethers, enamines) in PQ-ERA photocycloadditions, our group discovered that in this photoclick system, which simultaneously introduces a fluorescent moiety, the reaction rates, photoreaction quantum yields (Φ_P up to 65%) and fluorescence quantum yields (Φ_F up to 97%) can be drastically increased.^{14,18–20}

The PQ-ERA proceeds selectively in a range of organic solvents such as acetonitrile (MeCN), toluene, dichloromethane, and ethyl acetate, while the use of ethereal solvents and alcohols leads to formation of side products.^{14,16} The PQ-ERA can also be performed in PBS-buffer, making it relevant for biological applications.^{17,18} However, as with many other photoclick reactions, the limited solubility of its substrates (especially PQs) in polar media often requires the use of small

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amounts of solubilizing additives.^{11,17,18,21–24} Considering the previously reported effects of organic solvents on the **PQ-ERA**, we were curious to explore the effects of such co-solvents on this photoclick reaction.

Herein, we report for the first time a study of the effect of various solubilizing co-solvents on the reaction kinetics of the **PQ-ERA** photoclick reaction. Strikingly, it was found that the commonly used co-solvent dimethyl sulfoxide (DMSO) has a significant effect on these kinetics (Scheme 1). The lifetime of the **PQ**-triplet state decreases in the presence of DMSO, leading to a 5-fold decrease in reaction rate. Without DMSO as co-solvent, extraordinary photoreaction quantum yields (up to 93%) can be achieved with typical reaction times of seconds.

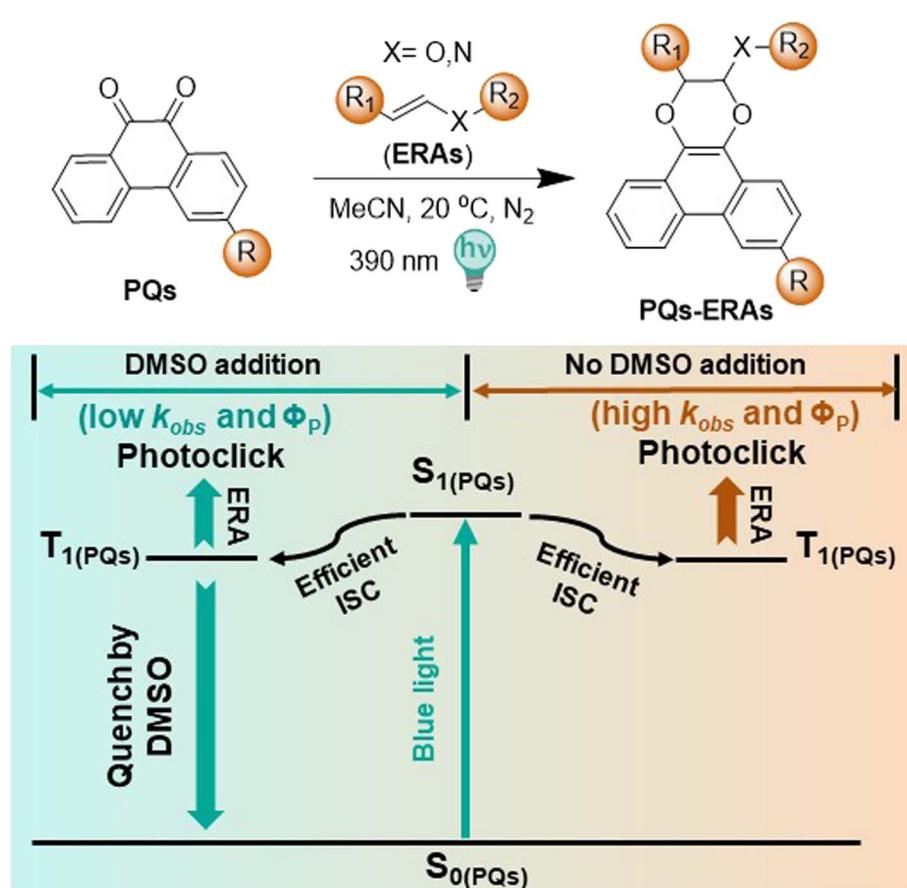
2 Results and discussion

We initiated this research by a systematic evaluation of the influence of various additives on the **PQ-ERA** reaction rate (in acetonitrile (MeCN) as the main solvent), including common alcohols (methanol (MeOH), ethanol (EtOH), isopropyl alcohol (IPA)), ethereal solvents (tetrahydrofuran (THF), dioxane), amide solvents (dimethyl formamide (DMF), dimethyl acetamide

(DMA)), sulfur-bearing additives (DMSO, dimethyl sulfide (DMS), sulfolane), and water. Unsubstituted **PQ** and *N*-boc-2,3-dihydro-1*H*-pyrrole **PY** (10 eq., Fig. 1A) were chosen as model compounds for this **PQ-ERA** reaction study, because of their accessibility and relatively fast reaction rate. It has to be noted that the reaction also proceeds efficiently with lower equivalents of **PY**.¹⁹ The progress of the reaction was monitored by UV-Vis absorption spectroscopy, which provided the reaction rate constants $k_{\text{obs}}(390)$ of the photoclick processes (Fig. 1B and C).

Remarkably, the **PQ-PY** reaction in pure MeCN showed a rate constant of 0.0297 s^{-1} , a substantial enhancement (~ 5 times) compared to previous reports (0.0062 s^{-1} with 0.5% (v/v) DMSO addition).¹⁹ This indicates that the commonly used co-solvent DMSO has a significant effect on the reaction kinetics. Intriguingly, the phenomenon observed upon DMS addition was similar to that of DMSO. Upon the addition of other solvents, including the sulfur-bearing sulfolane, minimal alterations in the $k_{\text{obs}}(390)$ were found (Fig. 1D, for detailed information see ESI, Section 2.1, Fig. S2†). The results were confirmed by HPLC analysis (see ESI, Section 3, Fig. S13–S22†).

The large change in reaction kinetics, caused by the addition of a small amount of DMSO, is striking, as DMSO is often used



Scheme 1 Comparison of two conditions for photocycloaddition of 9,10-phenanthrenequinone (PQ) with electron-rich alkenes (ERAs), cyan: with DMSO, brown: without DMSO. One of the possible isomers formed is shown as an example for clarity. k_{obs} : observed first-order reaction rate; and Φ_p : photoreaction quantum yield.



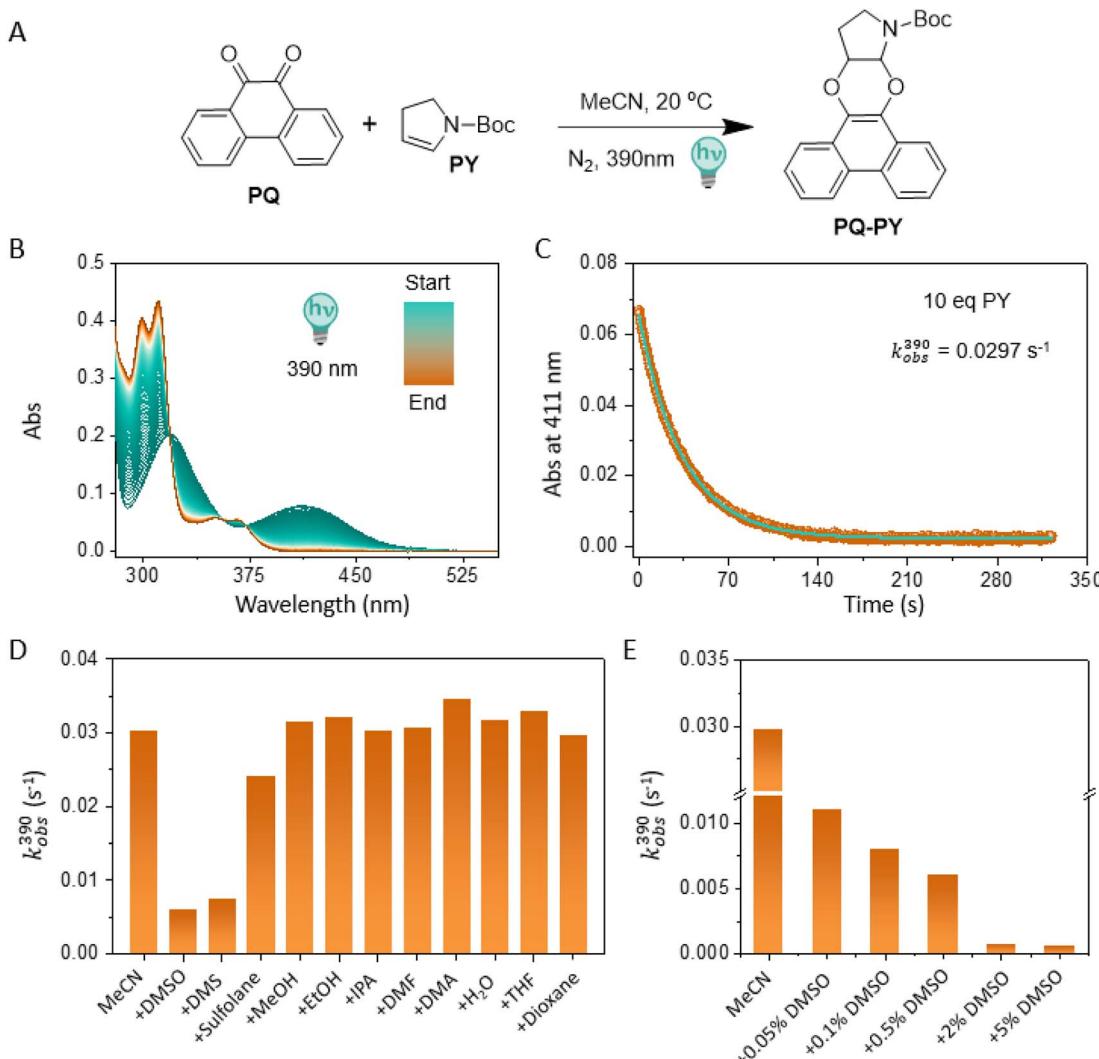


Fig. 1 Photocycloaddition reaction between 9,10-phenanthrenequinone (PQ) and 2,3-dihydropyrrrole (PY) in the presence of co-solvents and additives. (A) Reaction scheme; (B) time-resolved UV-Vis absorption spectra ($0\text{ s} \rightarrow 320\text{ s}$); (C) UV-Vis kinetic traces (at $\lambda = 411\text{ nm}$) of the photocycloaddition between PQ and PY in pure MeCN; (D) kinetic analysis of photocycloaddition between PQ and PY with various additives. These include DMSO, DMS, sulfolane, MeOH, EtOH, IPA, DMF, DMA, H_2O , THF and dioxane, respectively and, (E) concentration-dependent kinetic analysis of PQ-PY photoclick reaction in MeCN with addition of different amounts of DMSO. For all of the measurements, the sample was irradiated with a 390 nm LED and the absorbance at 411 nm was followed over time by UV-Vis absorption spectroscopy (1 cm cuvette, 2.5 mL MeCN sample volume, sample interval 1 s). PQ-PY formation was fitted to an exponential rise to the maximum equation, $y = (y_0 - a)e^{k_{\text{obs}} \times t} + b$, to give k_{obs} .

as co-solvent in biomedical studies because of its low toxicity and effective solubilizing properties.^{25,26} Therefore, we continued by conducting a detailed study on the effect of DMSO in the **PQ-ERA** photoclick reaction. Increasing amounts (0% to 5%, v/v) of DMSO were introduced, followed by the determination of $k_{\text{obs}}(390)$ within this concentration range. The results unveiled a notable trend in decreasing $k_{\text{obs}}(390)$ upon increasing the DMSO content (Fig. 1E, for detailed information see ESI, Section 2.1, Fig. S3†). For instance, with a DMSO addition of 5%, the $k_{\text{obs}}(390)$ plummeted to 0.0006 s^{-1} , a stark reduction of approximately 50-fold in comparison to the pure MeCN solvent system (0.0297 s^{-1}). Although the effects of increasing amounts of DMS were similar to that of DMSO (for

detailed information, see ESI, Section 2.1, Fig. S4†), we focused on DMSO due to its common use as co-solvent.

Reported indications of DMSO influencing the triplet states of *p*-quinones made us realize that potentially such an effect could also occur with the *o*-quinone **PQ** studied here.^{27,28} To investigate the generality of this DMSO-effect on the **PQ-ERA** photoclick reaction, we examined a range of **PQ** derivatives with varying substituents (depicted in Fig. 2A, ranging from electron-withdrawing groups to electron-donating groups). The $k_{\text{obs}}(390)$ for these photocycloaddition reactions was quantified (for detailed information, see ESI, Section 2.1, Fig. S5†) under both DMSO and DMSO-free conditions. Notably, as illustrated in Fig. 2B, the $k_{\text{obs}}(390)$ values for all **PQ** derivatives were

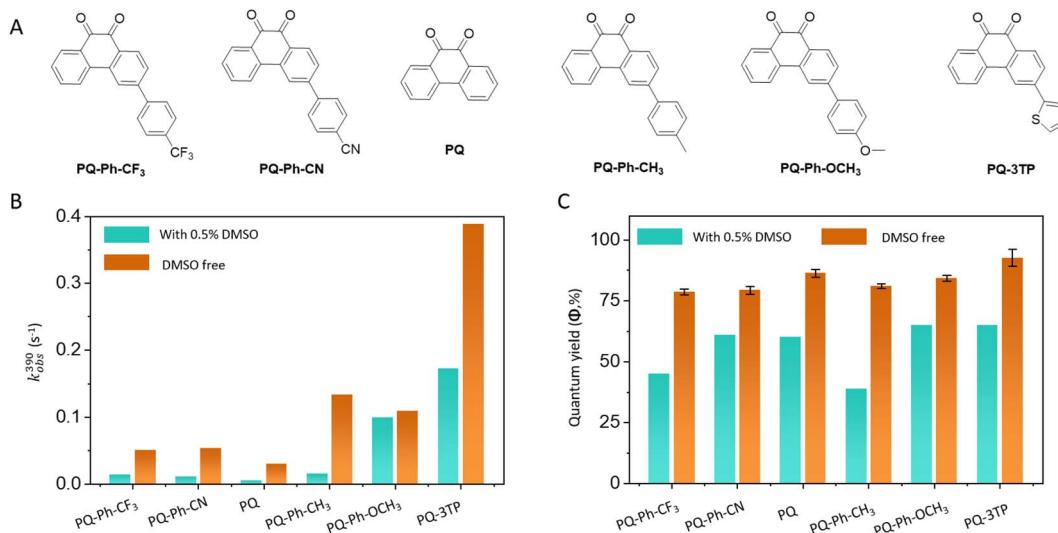


Fig. 2 Overview of the first order reaction rate (k_{obs}) and photoreaction quantum yield (Φ_p) of the PQ-ERA photoclick reaction system. (A) Structures of the investigated PQs; (B) The $k_{obs}(390)$ and (C) Φ_p of PQ-ERA photoclick reaction. Cyan: with DMSO, orange: DMSO-free, respectively. NB: k_{obs} and Φ_p are measured under different conditions (for more detailed information see ESI,† Section 2.1 and 2.2).

significantly higher without DMSO co-solvent, aligning with the trend observed for the unsubstituted **PQ**.

These results are also reflected by the extraordinary high photoreaction quantum yields (Φ_p) of these **PQ-ERA** reactions. The outcomes were evident across diverse **PQ** derivatives, engaged in reactions with **PY**, as illustrated in Fig. 2A (with and without DMSO co-solvent, respectively). Particularly noteworthy is the Φ_p value demonstrated by **PQ** in the DMSO-free context, achieving an impressive 86% with the addition of 10 equivalents of **PY** (Fig. 2C; for detailed information, see ESI, Section 2.2, Fig. S7–S12†).

Interestingly, the Φ_p of the **PQ-ERA** photoclick reactions in the absence of DMSO co-solvent become largely independent on electron-donating or electron-withdrawing nature of the substituents on the **PQ** molecule, showcasing uniformly high values of approximately 80%. It has to be noted that the effect of DMSO seems smaller for **PQ-Ph-OMe**, which also has a high $k_{obs}(390)$ and Φ_p in the presence of DMSO (for detailed information see ESI, Section 2.1, Fig. S6†).

The high Φ_p values reported here, surpass the efficacy of our previously introduced fast **PQs-VEs/PY** systems, with Φ_p (**PQs-VEs/PY**) ranging from 0.6% to 65%,^{14,18} with an exceptional case of 98%,¹⁹ as well as other documented photoclick reactions.⁴ In comparison to other photoclick processes such as *o*-naphthoquinone methides (*o*-NQMs)-ene hetero-Diels-Alder cycloaddition ($\Phi_p = 20\%$),²⁹ photoinduced tetrazole-alkene cycloaddition ($\Phi_p = 0.29\%$ to 24%),^{30,31} and photoinduced sydnone-alkene/alkyne cycloadditions ($\Phi_p = 10\%$ to 25%),^{21,32} the present system with a Φ_p of 83% establishes a ground-breaking benchmark.

2.1 Transient absorption spectroscopy

Intrigued by these results, we set out to investigate the cause of these large changes in kinetics upon introduction of DMSO as

a co-solvent. It is known that the formation of the triplet state of **PQ** is essential for the photocycloaddition to occur.¹⁹ Our previous studies demonstrated that unsubstituted **PQ** undergoes a rapid intersystem crossing (ISC), producing triplet states with a high quantum yield.¹⁹ We hypothesized that the reduced reaction rate upon addition of DMSO could be related to either the formation or the lifetime of the triplet state of **PQ**.

To investigate if DMSO influences the ISC process, we measured the transient absorption spectra of **PQ** in the sub-picosecond to nanosecond timescale (ultrafast setup) with and without addition of DMSO. The triplet absorption spectrum of **PQ** presents two positive signatures (a peak at about 460 nm and a double peak at 650–680 nm) that have been attributed to two different triplet states with $^3n\pi^*$ and $^3\pi\pi^*$ nature reaching fast equilibration.¹⁹ After excitation with 400 nm light, a transient spectrum attributed to the triplet state rises in about 9 ps (τ_2 , see

Table 1 Lifetimes obtained by fitting the transient absorption data measured for **PQ** in MeCN with different amounts of DMSO using a global analysis procedure^a

DMSO content (v/v) in MeCN	τ_1 ^b	τ_2 ^b	τ_3 ^c	τ_k
0%	1.0 ps	8.8 ps	3.8 μ s	85 μ s
0.05%	2.1 ps	8.7 ps	126 ns	—
0.1%	1.0 ps	9.1 ps	79.5 ns	—
0.5%	2.0 ps	8.6 ps	20.9 ns	—
1%	1.5 ps	8.6 ps	10.9 ns	—
2%	2.3 ps	8.5 ps	5.9 ns	—
5%	1.9 ps	9.0 ps	2.2 ^d ns	—

^a The fastest constant (τ_1) refers to a singlet excited state relaxation process; τ_2 represents the ISC rate, while τ_3 is the triplet lifetime. τ_k is associated with the formation of ketyl radical. ^b From ultrafast measurements (sub-ps to 1.5 ns). ^c From ns to μ s measurements. ^d From ultrafast measurements.

Table 1). The rise of this triplet state was not influenced by addition of different amounts of DMSO, up to 5% (Table 1 and ESI, Section 4.1, Fig. S23†). From these data, it was concluded that the rate of ISC is not affected by the addition of DMSO.

Next, we measured the transient absorption spectra of **PQ** in the nanosecond to millisecond timescale (ns setup), to examine the potential effect of DMSO on the lifetime of the triplet state. Our previous research confirmed that excitation of **PQ** leads to the presence of two long-lived species, the lowest-lying triplet state, which decays on a time scale of a few microseconds (τ_3), and a ketyl radical, which disappears on a time scale of about 80 μ s (τ_k).¹⁹ Upon the addition of 0.05% (v/v) DMSO we observed a 20-fold decrease in the lifetime of the triplet state as compared to DMSO-free conditions, lowering τ_3 from 3.8 μ s to 126 ns. Interestingly, formation of the ketyl radical was not observed anymore. Increasing the DMSO concentration resulted in a further reduction of τ_3 , to only several nanoseconds upon the addition of 2% (v/v) DMSO. The decreased triplet lifetime upon the addition of >0.5% (v/v) DMSO is already noticeable from measurements performed with the ultrafast setup (<1 ps to 1.5 ns), as shown in Fig. 3. Further details about the transient absorption measurements are reported in ESI (Section 4.1, Fig. S24†).

Additionally, we have found that τ_3 lowered with increasing excitation power, indicating occurrence of quenching at high irradiation energies. Interestingly, upon addition of 0.05% (v/v)

DMSO, the τ_3 did not depend on the excitation power anymore (ESI, Section 4.1, Fig. S25†). This indicates that the triplet does not exist long enough to recombine *via* triplet–triplet annihilation.

Plotting of the triplet state lifetimes according to the Stern–Volmer quenching equation showed a good correlation and the corresponding quenching rate constant was determined to be $5.97 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ (ESI, Section 4.2, Fig. S27†). This is in the same order of magnitude as other quenching mechanisms and indicates an intermolecular deactivation process upon the addition of DMSO. This quenching rate constant explains the differences in the experimentally observed quantum yields and rate constants.

The quenching effect of DMSO on **PQ-Ph-OCH₃** seems to be less pronounced than for that of other substituted **PQs** (for detailed information see ESI, Section 2.1, Fig. S6†). To explain this observation, we measured the transient absorption spectra of the various **PQs** (for detailed information see ESI, Section 4.1, Fig. S26†). In all cases, increased occupation of the $^3\pi\pi^*$ triplet state was observed, likely due to the asymmetric nature of these substituted **PQs**. However, compared to native **PQ**, the absorption band of this $^3\pi\pi^*$ was either blue-shifted (**PQ-Ph-CN**, **PQ-Ph-CF₃**), comparable (**PQ-Ph-CH₃**) or red-shifted (**PQ-Ph-OCH₃**). As an electron transfer mechanism is suggested for DMSO quenching of *para*-quinones,²⁷ which is consistent with the minimal effect of sulfolane on the photoclick, we hypothesize that the redox potential of the $^3\pi\pi^*$ of **PQ-Ph-OCH₃** is decreased

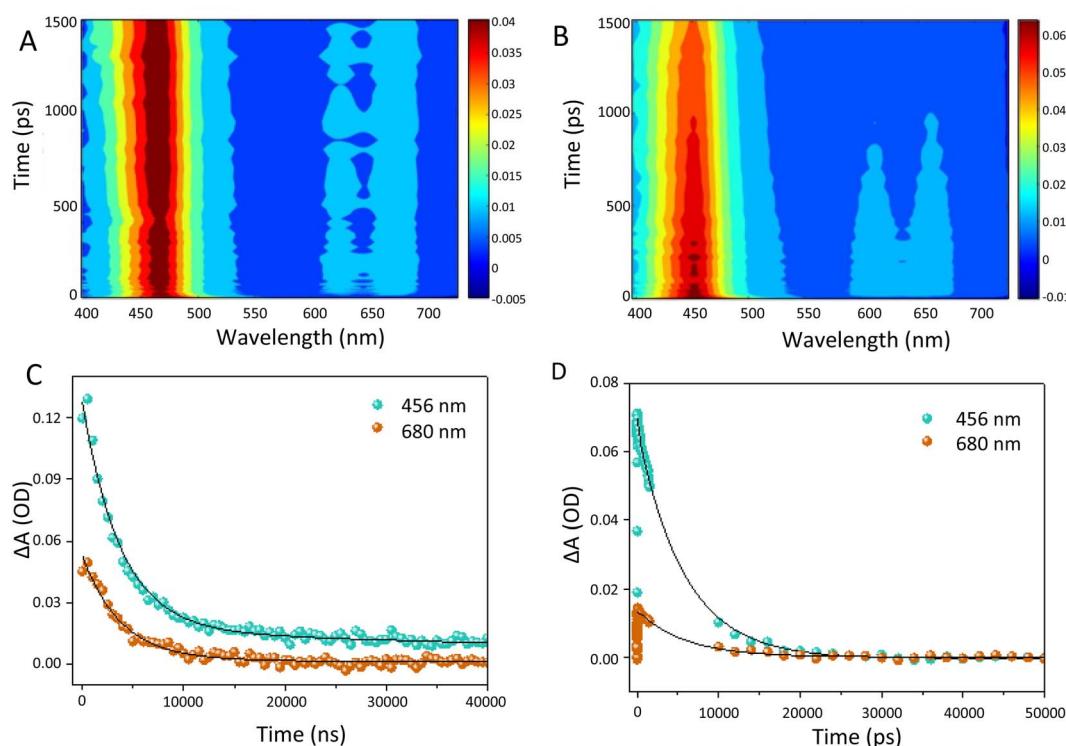


Fig. 3 Transient absorption measurements performed in the sub-ps to 1.5 ns time interval upon excitation at 400 nm on **PQ** dissolved in (A) MeCN and, (B) MeCN with addition of 2% (v/v) DMSO. (C) Kinetic traces recorded at 456 nm and 680 nm for **PQ** dissolved in MeCN, measured with the ns setup (10 ns to 40 μ s time interval). (D) Kinetic traces recorded at 456 nm and 680 nm for **PQ** dissolved in MeCN with addition of 2% (v/v) DMSO. Time points up to 1.5 ns have been acquired using the ultrafast setup, time points in the 10–50 ns range have been acquired using the ns setup (see ESI† Section 4 for further information).



compared to that of other **PQs**, making it less sensitive towards triplet quenching by DMSO.

2.2 Optimized **PQ-ERA** photoclick conditions

Having established that DMSO quenches the triplet state of **PQ**, we were able to rationalize the significant accelerations observed in our **PQ-ERA** photoclick process without the use of DMSO as co-solvent. This inspired us to investigate the kinetics and Φ_p of the **PQ-ERA** reaction using **PQ-3TP** (see Scheme 1, R = 2-thienyl), which we found to be the most efficient **PQ**-derivative so far. Our prior research indicated its enhanced reaction kinetics compared to standard **PQs** due to the advantageous generation of the reactive triplet state. We initially reported a Φ_p of 65% using 1 equivalent of **PY**. Remarkably, without DMSO as co-solvent, this Φ_p could be increased to 93% (For detail information see ESI, Section 2.2, Fig. S12†). This is near-quantitative conversion with equivalent amounts of **PQ** and **PY**, which further underscores the essential role of a long-lived triplet state. To the best of our knowledge, this achievement stands as the highest Φ_p among all known photoclick reactions.^{18,19}

Turning our attention to the fundamental work conducted by Schönberg and colleagues almost 80 years ago,^{15,33} we aimed for a comparison to benchmark the advancements offered by our newly devised system (For detail information, see ESI, Movie 1†). We compared our optimized system featuring **PQ-3TP**, **PY** (1 eq), and pure MeCN as solvent, against the original conditions involving **PQ**, stilbene, and benzene as solvent. The initial reaction requires 9 days of exposure to sunlight, in the intense Egyptian sun.³³ In contrast, we performed our reaction behind a glass window, utilizing the (limited) sunlight as the driving force on a February day in Groningen (The Netherlands). We were excited to observe the completion of the reaction in a mere 10 seconds (>77000 times faster than the original report, see ESI, Movie 1†), unequivocally highlighting that contemporary limitations in reaction time or light intensity are no longer barriers to advance **PQ-ERA** photoclick chemistry.

3 Conclusion

In summary, we have explored the impact of various solubilizing additives on the reaction kinetics of the **PQ-ERA** photoclick reaction. Notably, we found a substantial influence of the commonly employed co-solvent DMSO on these kinetics. The use of minimal quantities of DMSO leads to a remarkable decrease of the reaction speed (up to 5 times slower), accompanied by a drop in photoreaction quantum yield (Φ_p), from 80% to 65%. Employing transient absorption spectroscopy, we established that the DMSO quenches the triplet state of **PQ**. Using this knowledge, we evaluated the use of **PQ-3TP** in the **PQ-ERA** system without co-solvents. Addition of only 1 equivalent of **PY** leads to an outstanding Φ_p of 93% and reaction times of merely seconds to completion. These findings are testimony of the extraordinary characteristics of the **PQ-ERA** photoclick reaction system. We anticipate that the ultrafast **PQ-ERA** photoclick reaction, showcased in this work, holds promise beyond the depicted chemical

scenarios. Its potential advantages are expected to extend to diverse applications, including surface photo-patterning, biomacromolecule labeling, and photochemical crosslinking. Studies along these lines are now ongoing in our laboratories.

Data availability

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

Author contributions

A. M. D., Y. F., M. D. D., G. L., P. H. E., W. J. B., W. S., and B. L. F. conceived the project and designed phenanthrenequinone derivatives. B. L. F. and W. S. guided the research. Y. F. and A. M. D. synthesized all phenanthrenequinone derivatives. A. M. D. and Y. F. performed UV-Vis experiments. M. D. D. performed the Femtosecond Transient Absorption Spectroscopy measurements. A. M. D., Y. F., M. F. H., and W. J. B. performed the Nanosecond Transient Absorption Spectroscopy measurements. A. M. D., Y. F., M. D. D., G. L., P. H. E., W. J. B., W. S., and B. L. F. wrote the manuscript with support and contributions from all authors.

Conflicts of interest

There are no competing conflicts of interests to declare.

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