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N-Monoarylated dihydrophenazines in reduced and oxidized states as efficient organo-photocatalysts†

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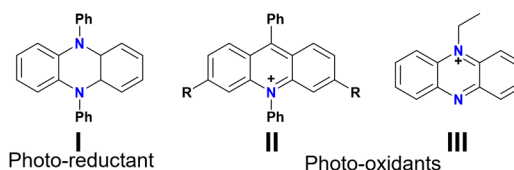
In this work, the synthesis of an *N*-monoarylated dihydrophenazine is reported together with its interconversion to its oxidized mono-cationic form. While the reduced state was employed for the dechlorination of aromatic substrates, the oxidized mono-cationic one was exploited for the formation of C–N bonds between aryl rings and azoles, which was achieved with high yields and very low catalyst loadings (down to 0.5 mol%).

N,N-Diphenyl-dihydrophenazines belong to a class of aromatic compounds that have undergone a renaissance in the last few years, due to their great potential for sensing,^{1–3} light emission,^{4,5} molecular switching^{6,7} and catalysis.^{8–11} In particular, dihydrophenazines have been extensively investigated, especially as photo-reducing agents (Fig. 1, I).^{9,10} In this perspective, we recently demonstrated that through π -extension of the dihydrophenazine scaffolds, it is possible not only to tune the optoelectronic properties, but also to introduce hydroxyl functional groups on the π -extended scaffold, which enables efficient photocatalysis.¹² Moreover, π -extension coupled with the design of the aromaticity patterns, resulted in much more accessible and stable dicationic states.^{7,13} Given their electron-depleted nature and generally intense absorption in the visible region, dicationic oxidized states of dihydrophenazines should present great potential as organo-photo oxidizers, leading to the potential use of both states (neutral and dicationic) as active photocatalysts. This is of course highly interesting since it would enable the use of the same scaffold for different photo-redox reactions, resulting in a great versatility and practicality. Despite this, while many

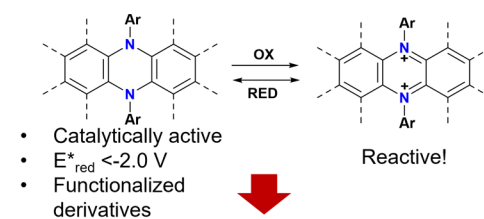
examples of photocatalytic applications of the neutral dihydrophenazine have been reported, the use of dicationic dihydrophenazine derivatives for this application has been hindered by their reactivity¹⁴ and general instability towards moisture and nucleophiles which persisted, albeit to a lesser extent, also in π -extended derivatives.⁷ One possible solution to this, could be the preparation of *N*-monoarylated dihydrophenazines, which upon oxidation should form mono-cationic phenazine derivatives (1 and 1^{ox}, Fig. 1).

The latter should present an ideal balance between stability and reactivity aiming at catalytic applications, possibly with

Previous works



Our group 2022 and 2023



This work

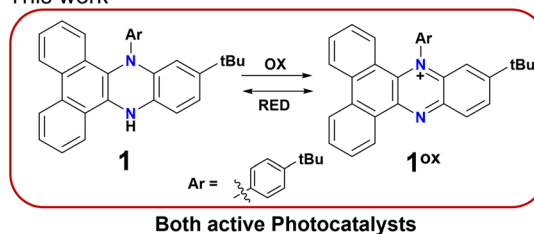


Fig. 1 Previously reported photocatalytic systems and recent work on π -extended *N,N*-diphenyl dihydrophenazine systems.

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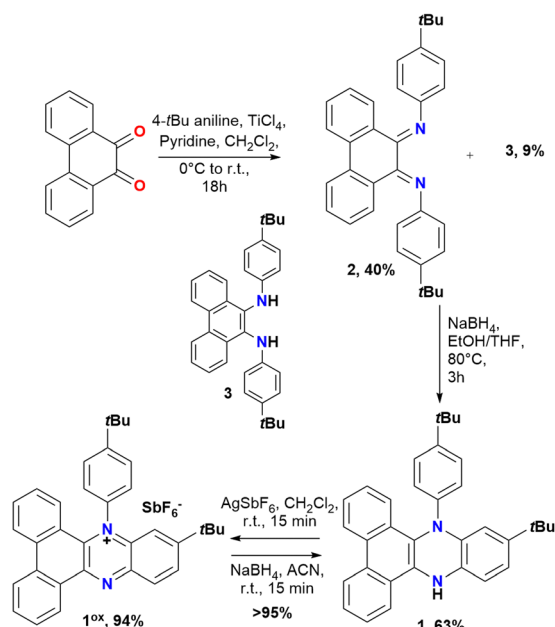


activities similar to or better than those of the well-established acridinium catalysts (Fig. 1, **II**).^{15–18} From this point of view, a few examples of mono-cationic phenazinium cations have been reported in the literature, and have been generally exploited for DNA binding¹⁹ or, in a notable example, photocatalytic oxidation of aldehydes to amides.²⁰ In the former example, π -extended systems were used,¹⁹ while in the latter the simpler phenazine ethosulfate (Fig. 1, **III**) was employed with low (1–2 mol%) catalyst loadings resulting in promising high yields.²⁰ Some more examples of similar mono-cationic systems employed as photocatalysts exist, but are generally based on functionalized safranin dyes presenting excited state oxidation potentials much lower than acridinium systems.^{21–23}

Based on this, to prepare target molecule **1**, it was decided to start by synthesising phenanthrene diimine **2**, presenting *t*Bu groups for enhancing the solubility of the system. To achieve this, phenanthrenequinone was treated with 4-*t*Bu aniline in the presence of TiCl₄ and pyridine, resulting in the formation of **2** in a 40% yield (Scheme 1). As correctly reported by Hoyer and co-workers, also diamine **3** was isolated in a modest 9% yield. Subsequently, **2** was refluxed in a THF/EtOH mixture. As a result, after 3 hours, the formation of the desired product could be observed, which proved to be **1** (see Fig. S32–S37, ESI[†]) in accordance with the expected 6 π -electrocyclization.²⁴ Since **1** proved to be partially oxidized during the reaction, this was repeated with the addition of 1 equivalent of NaBH₄ resulting in a cleaner outcome with isolation of **1** in a 63% yield. Derivative **1** proved to be stable for weeks in the solid state and could be completely characterised *via* nuclear magnetic resonance spectroscopy (NMR). Given that the two-electron oxidation of **1** should result in the formation of phenazinium system **1^{ox}**, the former was treated with an excess of AgSbF₆ (3 equivalents) resulting in an immediate colour change from faint yellow to

brown/orange associated with the presence of a new species. After filtration to remove silver salts and reprecipitation from Et₂O, the new compound was isolated and characterised by NMR and high-resolution mass spectroscopy (HRMS) confirming the structure of **1^{ox}**. Interestingly, the conversion between **1** and **1^{ox}** proved to be chemically reversible by treating the latter with NaBH₄, as demonstrated by NMR experiments (see Fig. S23–S25, ESI[†]), thus enabling the interconversion between the two species. To prove the usefulness of derivatives **1** and **1^{ox}** as organo-photocatalysts, the optoelectronic properties of both derivatives were analysed. Derivative **1** presents an absorption with a $\lambda_{\text{max}} = 354$ nm, tailing in the visible region (up to *ca.* 450 nm), associated with a broad emission $\lambda_{\text{max}} = 590$ nm (Fig. S1, ESI[†]). On the other hand, **1^{ox}** presents a more red-shifted absorption with $\lambda_{\text{max}} = 462$ nm in the blue region of the spectra, resembling similar derivatives reported in the literature.¹⁹ Also in this case, the emission is broad with $\lambda_{\text{max}} = 630$ nm associated with a less marked Stokes shift compared to **1** (Fig. S7, ESI[†]). For both species the quantum yields proved to be modest, with a value <5% for **1** and of 5% for **1^{ox}**. Cyclic voltammetry measurements were carried out for both compounds in CH₂Cl₂. In the case of **1** only a single quasi reversible oxidative event was visible at $E_{1/2} = +0.24$ V vs. SCE (Fig. S17, ESI[†]). On the other hand, for **1^{ox}** two reversible reductive events were observed, at $E_{1/2} = -0.21$ and -0.84 V vs. SCE (Fig. S18, ESI[†]) while no oxidations were detected. Based on these values, using the Rehm–Weller equation,²⁵ it was possible to estimate an excited state oxidation potential of -2.25 V for **1** and an excited state reduction potential of $+2.13$ V for **1^{ox}**,²⁶ suggesting that indeed the two derivatives are highly promising in terms of photo-reduction and oxidation of organic substrates. In particular, in the case of **1^{ox}**, the observed excited state oxidation potential is close to that of some of the most oxidant acridinium ions reported in the literature.^{15,16} To evaluate the photocatalytic activity of the **1/1^{ox}** couple, it was decided to start with derivative **1** as a photo-reducing agent, by using the dehalogenation of 4-chlorobenzonitrile as the test reaction. As such, 10 mol% of **1** in DMF [0.1 M] was irradiated together with the substrate and 1 equivalent of DIPEA, with two 456 nm kessil lamps for 20 h. As a result, the dehalogenation product could be obtained in a 39% yield (Table S1, ESI[†]). Increasing DIPEA to 10 equivalents only resulted in a slight increase in the yield. Substituting DIPEA with 10 equivalents of DBU resulted in a marked improvement (84%, entry 3, Table S1, ESI[†]). While increasing or decreasing the DBU equivalents did not enhance the reaction outcome (entries 4 and 5, Table S1, ESI[†]), the addition of 1 equivalent of DIPEA (in the presence of 10 equivalents of DBU) resulted in a quantitative conversion (>95%). As a control, a reaction was carried out in the absence of **1**, resulting in the complete loss of reactivity. The same happened when the reaction was carried out in the dark, suggesting that the reaction is indeed photocatalytic.

Finally, performing the reaction in air resulted in only 15% yield, highlighting the radical nature of the process. Based on these results, several substrates were evaluated (Fig. 2),



Scheme 1 Synthetic procedure for the preparation of **1** and **1^{ox}**.



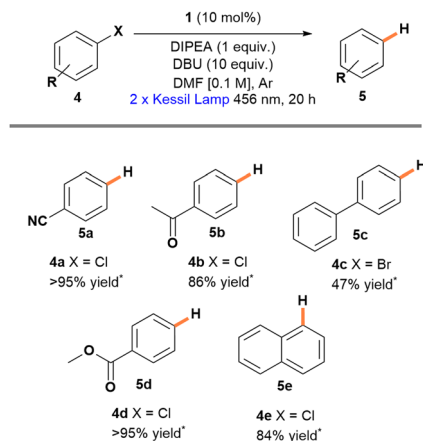


Fig. 2 Substrate scope for photo-reductive dehalogenation using **1** as a photocatalyst. *NMR yields obtained with trichloroethylene (TCE) as an internal standard.

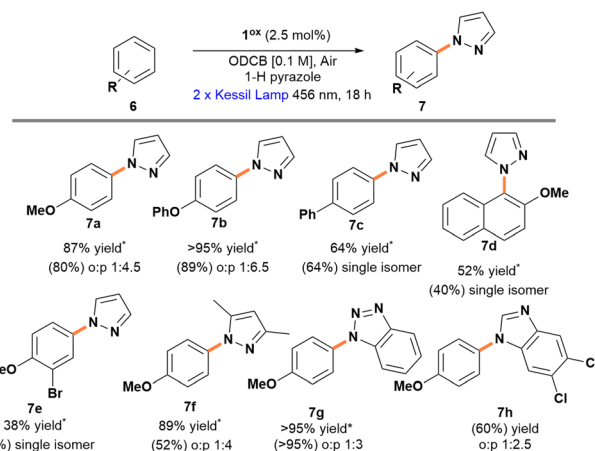


Fig. 3 Substrate scope for the photo-oxidation reaction using **1^{ox}** as a photocatalyst. *NMR yields obtained with TCE as an internal standard. Yields in parentheses are isolated yields.

demonstrating that it is possible to dehalogenate systems containing esters (**4d**), ketones (**4b**), polyaromatics (**4e**) and also non-activated biphenyls (**4c**). To complete the mechanistic investigation, a Stern–Volmer titration was performed with **4a** as a quencher on a 1.0×10^{-5} M solution of **1** in DMF resulting in a visible quenching of the emission (Fig. S4, ESI[†]). Also, no EDA complex formation was observed in different mixtures of 4-chlorobenzonitrile, DBU and **1** in DMF (Fig. S3, ESI[†]), suggesting that the reaction is based on a SET event from **1** to **4a**. Overall, these results and the marked improvement of the reaction outcome in the presence of a base (DBU) led us to hypothesise a mechanism involving a proton-coupled electron-transfer plausibly similar to the one reported by Bortolato *et al.* (Fig. S14, ESI[†]).²⁷ This is consistent with the presence of a hydrogen bonding interaction between DBU and the NH proton in **1** as highlighted by an NMR titration performed in DMF (Fig. S20, ESI[†]). The catalytic behavior of **1^{ox}** was then studied in the photo-oxidation of anisole (**6a**), followed by trapping of the resulting radical with 1H-pyrazole (Table S3, ESI[†]).¹⁷ A first reaction was carried out by irradiating a 0.1 M solution of anisole in dichloroethane (DCE) with 10 mol% of **1^{ox}** under air for 18 h. The desired product **7a** was formed in a 73% yield as a 4.6:1 mixture of *p*:*o* isomers (Table S3, ESI[†]). Building on this encouraging result, the reaction was repeated twice adding 10 and 20 mol%, respectively, of TEMPO as additive,¹⁷ which however did not result in improvement of the reaction yield (entries 2 and 3, Table S3, ESI[†]). At this point *o*-dichlorobenzene (ODCB) was used in place of DCE (entry 4, Table S3, ESI[†]), resulting in a marked increase of the yield up to 92%. Also in this case, performing the reaction in the absence of catalyst resulted in a complete loss of reactivity, as did carrying out the reaction in the absence of light (entries 5 and 6, Table S3, ESI[†]). Interestingly, decreasing the catalyst content from 10 mol% to 2.5 mol% did not affect significantly the reaction outcome, while moving down to 1.0 and 0.5 mol% only resulted in a moderate decrease in yield (entries 9–11, Table S3, ESI[†]). Also, irradiating the reaction at less energetic

wavelengths (525 nm) resulted in a good reactivity (entry 8, Table S3, ESI[†]) with a 63% yield of the desired product. Performing the reaction under an inert atmosphere (argon) resulted in a marked yield decrease (entry 12, Table S3, ESI[†]), highlighting the important role of oxygen in the reaction mechanism. Since **1** is known to be oxidized in air, the reaction was repeated using 2.5 mol% of **1** as the catalyst, resulting in a 24% yield of **7a**. This suggests that **1** can be oxidized in the reaction conditions to the active species resulting in reactivity albeit with lower yield than with isolated **1^{ox}**. Increasing the amount of **1** resulted in an enhancement in the yield of **7a** up to 40% (entry 14, Table S3, ESI[†]). To understand if **1^{ox}** was responsible for this reactivity, a solution of **1** was irradiated for 2 hours at 456 nm and upon UV-Vis analysis the absorption of **1^{ox}** was clearly visible (Fig. S13, ESI[†]). Having defined the best conditions for the reaction, different substrates were tested (Fig. 3). From this point of view, performing the reaction on diphenyl ether resulted in a quantitative conversion to **7b**, while using less activated biphenyl as a substrate produced a 64% yield of **7c** as a single isomer. Naphthalene derivative **7d** was also obtained in a good 52% yield, while halo derivative **7e** resulted in a lower 38% yield.

Finally, different azole derivatives were tested: in the case of methylated 1H-pyrazoles, derivative **7f** was obtained with an 89% yield, and when benzotriazole was used instead, **7g** was formed quantitatively, whereas the use of the halogenated 5, 6-dichloro benzimidazole resulted in the formation of **7h** in a 60% yield. To elucidate the reaction mechanism, also in this case UV-Vis experiments were carried out. As for **1** no EDA complexes were visible by adding 100 or even 1000 eq. of **6a** to **1^{ox}** (Fig. S11, ESI[†]). Instead, when a Stern–Volmer titration was performed on 8.3×10^{-6} M of **1^{ox}** using anisole **6a** as a quencher, a clear decrease in the emission was visible, suggesting that the photooxidation is occurring *via* a SET event from **6a** to **1^{ox}** (Fig. S9, ESI[†]). This result, together with the previous experiments suggests that the reaction should occur following a similar mechanism to the one reported by Nicewitz and



co-workers for acridinium ions (Fig. S15, ESI†).¹⁷ In conclusion, in this work, the synthesis of a novel *N*-monoaryl dihydrophe-nazine is reported along with its oxidation to the corresponding monocationic derivative **1^{ox}**. The compounds were completely characterised, also demonstrating the reversibility of the oxidation/reduction process. Moreover, both forms proved to behave as effective photocatalysts. In particular, **1^{ox}** was shown to promote the oxidative coupling between electron rich aromatics and azoles at low catalyst loadings (from 2.5 to 0.5 mol%) and without the need for additives or pure O₂ atmospheres. As such, the present work is a relevant imple-mentation of the recent studies on π -extended dihydrophena-zine systems, which are expected to be further developed for catalytic and material applications in both their reduced and oxidized forms.

Maurizio Prato: conceptualization, reviewing and editing of manuscript Jacopo Dosso: synthesis, characterisation, concep-tualization, writing of original draft.

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Data availability

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

Conflicts of interest

There are no conflicts to declare.

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