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Photochemical aerobic oxidation of sulfides to sulfoxides: the crucial role of wavelength irradiation†

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The sulfoxide moiety is recognized as one of the most important groups in organic and medicinal chemistry. Many efforts worldwide focus on developing novel and sustainable protocols, accessing sulfoxide-containing molecules. Photochemistry plays a unique role in sulfoxide synthesis, since this new emerging field, employing light as the energy source, is essential in developing novel and eco-friendly protocols. Herein, we report a novel, sustainable, light-driven protocol, where the impact of wavelength irradiation on sulfide aerobic photooxidation was examined. In this work, two different low-catalyst loading (0.05–0.5 mol%) protocols, utilizing anthraquinone as the photocatalyst (under CFL lamps or 427 nm irradiation) were developed and a catalyst-free protocol (under 370 nm irradiation) was also assessed. In addition, a broad scope of substrates was tested and extensive mechanistic studies were performed, in order to distinguish the mechanistic pathways that are followed in the different cases of aryl or alkyl sulfide oxidation under different wavelength irradiation. We also implemented our protocols towards the oxidation of several intermediates *en route* to Sulforaphane.

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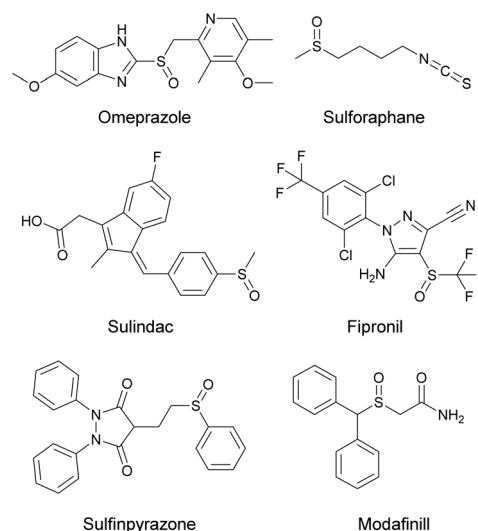
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Introduction

Sulfoxide-containing compounds display a prominent role in the field of organic synthesis,¹ pharmaceuticals,² agrochemicals³ and polymers.⁴ Several sulfoxides are known to display a salient pharmacological profile and have been introduced as marketed drugs, including omeprazole, sulindac, sulfinpyrazone and sulforaphane (Scheme 1). Over the past years, scientists have developed numerous protocols for the synthesis of sulfoxides, where a wide variety of oxidants, such as oxone,⁵ *m*-chloroperbenzoic acid,⁶ sodium periodate,⁷ chromic acid,⁸ chlorine⁹ *etc.*, were employed for the selective oxidation of aryl and/or alkyl sulfides. In all the above-mentioned protocols, the major drawback is the use of more than one equivalent of oxidant, which generates over-stoichiometric amounts of waste. Moreover, in all methodologies presented in literature, another issue is selectivity, highlighted by the formation of the corresponding sulfone, due to over-oxidation of sulfides. Even though the presence of the corresponding sulfone in some cases appears in a very small percentage, in the case of an

active pharmaceutical ingredient this is prohibitive. In addition, in most reported methodologies, heating in high temperatures narrows their potential for application in industry. A more green, sustainable, and popular approach is the use of hydrogen peroxide as the oxidant,¹⁰ since its main by-product is water. However, the activation of hydrogen peroxide



Scheme 1 Selected examples of pharmaceuticals' active ingredients or natural products containing the sulfoxide moiety.

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‡ Denotes equal contribution.



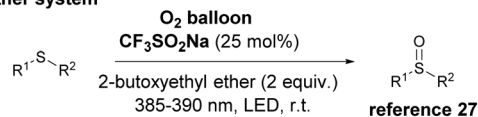
requires either heat¹¹ or metal catalysts.¹² From an environmental point of view, metal-free organocatalysts, such as flavins¹³ or activated ketones¹⁴ have been reported.

Apart from the environmentally friendly and sustainable hydrogen peroxide, another mild oxidant, widely employed in oxidation protocols is oxygen.¹⁵ It is an ambient, abundant, harmless gas, which can be considered the “greenest” approach towards oxidation reactions. The “green” character of molecular oxygen as the oxidant, either deriving from ambient air or being supplied as a gas to the reaction mixture, was very early realized.¹⁵ In recent years, different groups have reported metal-catalyzed¹⁶ or organocatalyzed¹⁷ processes that utilize oxygen as the oxidant in various processes. In a very recent example, in 2020, Petsi and Zografos designed a series of novel proline-based organocatalysts and examined their application in the oxidation of sulfides to sulfoxides, utilizing HFIP as the solvent.¹⁸

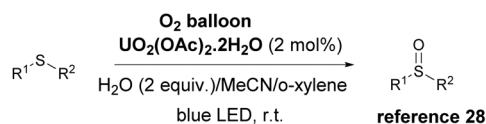
Photochemistry is an exciting and powerful area of research, which deals with electronically excited molecules and chemical reactions induced by light.^{19,20} Despite the fact, that it was more than clear that the reactivity of excited molecules differs from those in ground state and light-mediated chemistry has the potential to trigger reactions that are unavailable to ground-state pathways, synthetic photochemistry remained for many years a specialized area of research with limited practical applications. In 2008, seminal publications from MacMillan,^{20a} Stephenson,²¹ and Yoon²² changed this notion and the synthetic community quickly recognized the new opportunities created by photoredox catalysis and photochemistry.

The inherent advantages of photochemical protocols and aerobic (oxygen)-mediated oxidations were recognized quite early and the merger of these two areas was envisaged in the oxidation of sulfides to sulfoxides from the 1960s.^{23,24} Throughout the years, many synthetic efforts were made for the photochemical aerobic oxidation of sulfides, however most of them focused on the use of specialized apparatus for irradiation or the use of heterogeneous and in some cases homogeneous catalysts (metal-based or organocatalysts).²⁴ With the advent of Photoorganocatalysis, the use of small organic molecules for photochemical reactions,²⁵ the renewal of research interest in this photochemical aerobic protocols occurred. In 2019, Liu and co-workers, following their interest in developing a new protocol using a mild oxidant, proposed the use of molecular oxygen and the non-photochemical reaction was held in the presence of an ether, which promoted the reaction.²⁶ In that case, the use of oxygen in the presence of a flammable organic solvent limited their industrial application. To overcome these disadvantages, in 2021, they exploited molecular oxygen as the sole oxidant in a visible-light-induced sulfide oxidation using CF₃SO₂Na (25 mol%) as the photocatalyst (Scheme 2A).²⁷ However, a high catalyst loading for such a process had to be employed. Since 2017, a number of protocols have been reported for the photochemical aerobic oxidation of sulfides, such as the selective oxidation of sulfides induced by visible light in the presence of uranyl salts

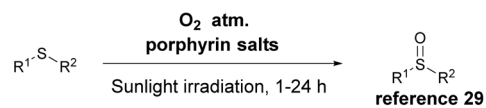
A. Sulfide oxidation employing CF₃SO₂Na / 2-butoxyethyl ether system



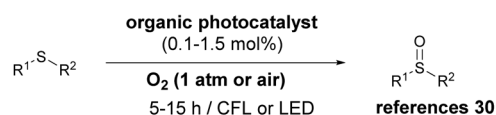
B. Sulfide oxidation mediated by uranyl salts



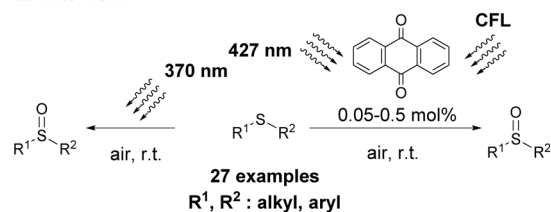
C. Sulfide oxidation using porphyrin derivatives



D. Sulfide oxidation using organic photocatalysts



E. This work



Scheme 2 Photochemical aerobic oxidation of sulfides.

(Scheme 2B)²⁸ or porphyrin derivatives as the photocatalyst (Scheme 2C).²⁹ In addition, a variety of visible-light induced oxidations, employing either naturally-occurring^{30b,c} or commercially available molecules^{30a} as the organic photocatalyst were also reported recently in literature (Scheme 2D). However, expensive uranyl salts, expensive and hazardous solvents or/and the synthesis of the photoorganocatalysts, not only increases the manufactory cost, but poses difficulties in the final purification of the product. Furthermore, in all these cases where organic photocatalysts were employed (Scheme 2D), the two generic mechanisms of aerobic oxidations (singlet oxygen or single electron transfer) were proposed to be operating, without proving if one of them is dominant and if the mechanism of action changes, depending on the nature of the employed substrate.

Apart from the use of several catalysts for either oxygen or substrate activation, many efforts have targeted the development of catalyst-free protocols. Since 1990s, a series of publications were focused on studying the behaviour of sulfides under visible light irradiation,^{24,31} however, the major drawback was the formation of undesired by-products, since the C–S fragmentation pathway prevailed the oxidation one. Lately,



Sun and coworkers,³² while testing the use of a benzothiadiazole-based conjugated microporous polymer as the photocatalyst to trigger sulfide oxidation, discovered that under blue LED irradiation (100 Watt), the oxidation proceeded in the absence of a catalyst. This novel catalyst-free approach of sulfide oxidation suffers from limited substrate scope (only aryl aryl or alkyl aryl sulfides were successfully employed) and prolonged reaction times, which in some cases is crucial for the formation of several by-products. Also, in their proposed mechanism, although they employ DFT calculations, they proposed a thiadioxirane intermediate as the active intermediate,³² however, this intermediate has been discarded by researchers as a possible intermediate in the recent past.²⁴

Following our continuous interest in developing novel light-driven methodologies,³³ herein, we report a mild, green and chemoselective direct photochemical aerobic oxidation of sulfides to the corresponding sulfoxides (Scheme 2E). From an environmental and economical point of view, this novel, green and sustainable protocol for the oxidation of sulfides using a mild oxidant, such as molecular oxygen, in the presence of a low-cost commercially available photocatalyst serves the basic principles of green chemistry, which is highly desirable on the chemical and pharmaceutical industry. We, furthermore, point out the crucial role of the wavelength irradiation source, towards the selectivity and the mechanism of the oxidation process. In this sense, we have developed a fast and green catalyst-free protocol for the photochemical aerobic oxidation of sulfides under 370 nm irradiation, while two low-catalyst loading (0.05 or 0.5 mol%) protocols (under CFL lamps or 427 nm irradiation) were also developed. Finally, in our attempt to shed light on the reaction mechanism, we examined the role and application of several known quenchers into the investigation of the reaction mechanism, and we propose a chart flow, researchers could follow in the future.

Results and discussion

We initially investigated the visible light-mediated aerobic oxidation of thioanisole (**1a**), as the model reaction to optimize the reaction parameters (Table 1). In the beginning, we studied the influence of the wavelength irradiation source in the outcome of the photochemical aerobic oxidation (Table 1). The self-excitation of sulfides is well established,³¹ however, usually harsh reaction conditions are employed (Xe or Hg lamps for the irradiation). For this purpose, the reaction was conducted in the absence of a photocatalyst, under various wavelengths of LEDs or CFL household lamps (Table 1). More specifically, under the irradiation of compact fluorescence lamps (CFL), the reaction appears to be sluggish (Table 1, entry 1). In this case, the use of an appropriate photocatalyst to facilitate the reaction is required.³⁴ On the contrary, the use of LEDs, as the irradiation source, facilitates the reaction, leading not only to the desired product, but also to the corresponding sulfone, due to over-oxidation (Table 1, entries 2–8). Also, it is obvious that upon decrease of the wavelength

Table 1 Examination of the impact of the irradiation source on the photochemical aerobic oxidation of sulfide **1a**^a

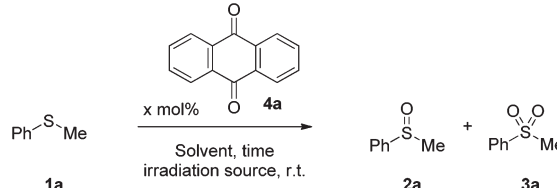
$\text{Ph-S-Me} \xrightarrow[\text{irradiation source, r.t.}]{\text{MeCN, 18 h}} \text{Ph-S(=O)-Me} + \text{Ph-S(=O)}_2\text{-Me}$				
	1a	2a	3a	
Entry	Lamps (nm)	1a ^b (%)	2a ^b (%)	3a ^b (%)
1	CFL	89	11	0
2	370	0	61	39
3	390	0	56	44
4	427	0	81	19
5	440	60	40	0
6	456	98	2	0
7	467	99	1	0
8	525	100	0	0

^a The reaction was performed with **1a** (0.20 mmol) in MeCN (1 mL) under open air irradiation for 18 h. ^b Conversion was determined by ¹H-NMR.

irradiation of the LED, the ratio of the products alters and the sulfone by-product increases significantly (Table 1, entries 2 and 3 vs. 4–8). Next, we turned our attention into optimizing the reaction conditions, employing a suitable photocatalyst when needed, to shorten the reaction time and suppress the formation of the overoxidized byproduct.³⁴ A thorough investigation on the reaction outcome based on different irradiation wavelength and reaction conditions can be found in the ESI.[†]³⁴ From these studies, we concluded that the irradiation wavelength and reaction conditions are crucial for product distribution (sulfide : sulfoxide : sulfone), and as will be described below, CFL lamp irradiation, as well as 370 nm or 427 nm, were chosen as optimum for the development of catalyst and catalyst-free protocols.³⁴

Initially, we investigated the sulfide oxidation under household lamp irradiation, employing a wide range of photocatalysts and we concluded that anthraquinone (**4a**) was the one that outperformed the others.³⁴ Then, we turned our attention to the nature of the reaction medium (Table 2). Protic solvents, *i.e.* methanol, ethanol afforded the best results (Table 2, entries 2–5), while in aprotic solvents, the reaction proved to be sluggish (Table 2, entries 7 and 8).³⁴ Addition of 10 equiv. of H₂O was essential in accelerating the reaction rate and suppressing the formation of the sulfone by-product (Table 2, entry 4 vs. entry 3).³⁴ Interestingly, when the reaction was conducted using water as the solvent, no formation of the desired product was observed, due to the insolubility of anthraquinone in water (Table 2, entry 6). Under LED irradiation (427 nm), sulfide oxidation led to a selective formation of the corresponding sulfoxide vs. the sulfone, after 5 h of irradiation in the presence of 0.05 mmol% of **4a** (Table 2, entry 10 vs. entry 9). When the reaction was performed in methanol, the desired product was obtained in quantitative conversion, suppressing the undesired sulfone by-product (Table 2, entry 11). In aprotic solvents, the reaction was incomplete (Table 2, entries 12–15). The optimum reaction conditions were also tested under sun-



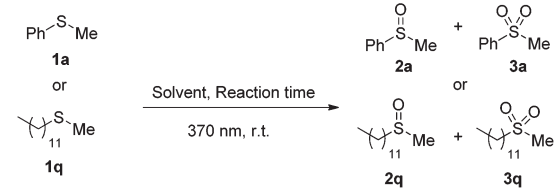
Table 2 Optimization of the reaction conditions for the photochemical aerobic oxidation of **1a**, using a photocatalyst^a


Entry	Lamp (nm)	Solvent, time (h)	4a (mol%)	1a ^b (%)	2a/3a ^b (%)
1	CFL	MeCN, 18	0.1	71	29/0
2	CFL	MeOH, 18	0.1	61	39/0
3 ^c	CFL	MeOH, 18	0.5	0	95/5
4	CFL	MeOH, 18	0.5	0	100/0
5	CFL	EtOH, 18	0.1	69	31/0
6	CFL	H ₂ O, 18	0.1	100	0/0
7	CFL	CH ₂ Cl ₂ , 18	0.1	96	4/0
8	CFL	DMF, 18	0.1	67	26/0
9 ^c	427	MeCN, 18	0.05	0	81/19
10 ^c	427	MeCN, 5	0.05	36	64/0
11 ^{c,d}	427	MeOH, 5	0.05	0	100/0
12 ^{c,d}	427	EtOAc, 5	0.05	11	89/0
13 ^{c,d}	427	CH ₂ Cl ₂ , 5	0.05	20	80/0
14 ^{c,d}	427	Benzene, 5	0.05	72	28/0
15 ^{c,d}	427	DMF, 5	0.05	50	50/0

^a The reaction was performed with **1a** (0.20 mmol) in solvent (1 mL) and H₂O (40 μL), under open air irradiation for 18 h. ^b Conversion was determined by ¹H-NMR. ^c The reaction was performed without the addition of H₂O. ^d 0.5 mL of solvent instead of 1 mL.

light irradiation, leading to a quantitative yield of **2a**.³⁴ In both cases (CFL lamps and 427 nm), low-catalyst loadings were employed, outperforming most of the known procedures highlighted in Scheme 1, while in the 427 nm protocol, the reaction required short reaction time.

We also targeted in identifying a catalyst-free protocol for the photochemical aerobic oxidation of **1a** (Table 3). To our delight, when the reaction was performed under UVA-LED irradiation (370 nm), quantitative formation of the desired product was obtained after 3 h, under catalyst-free conditions (Table 3, entry 3). Prolonged reaction time and the presence of a photocatalyst led to the formation of sulfone **3a** (Table 3, entry 1). Decreasing the reaction time and omitting the presence of the photocatalyst, led to a better yield of **2a** (Table 3, entry 2). Adding of 10 equiv. of H₂O led to an accelerated reaction, while suppressing the formation of the overoxidation by-product (Table 3, entry 3). Due to hydrogen bond formation, water inhibits the conversion of sulfoxide to sulfones.³⁵ Again, solvent optimization did not reveal any medium leading to better yields (Table 3, entries 4–8). It is important to note that this fast and mild reaction protocol (3 h and 45 W 370 nm) provides a better alternative to literature (11 h and 100 W Blue LED).³² A major drawback of the literature process³² is that it cannot be applied for dialkyl sulfides. Our desire to develop a photochemical protocol with broad scope of substrates led us to investigate the optimized reactions conditions on the less reactive dialkyl sulfides. Dodecyl methyl sulfide (**1q**) was

Table 3 Optimization of the reaction conditions for the photochemical oxidation of alkyl sulfides^a


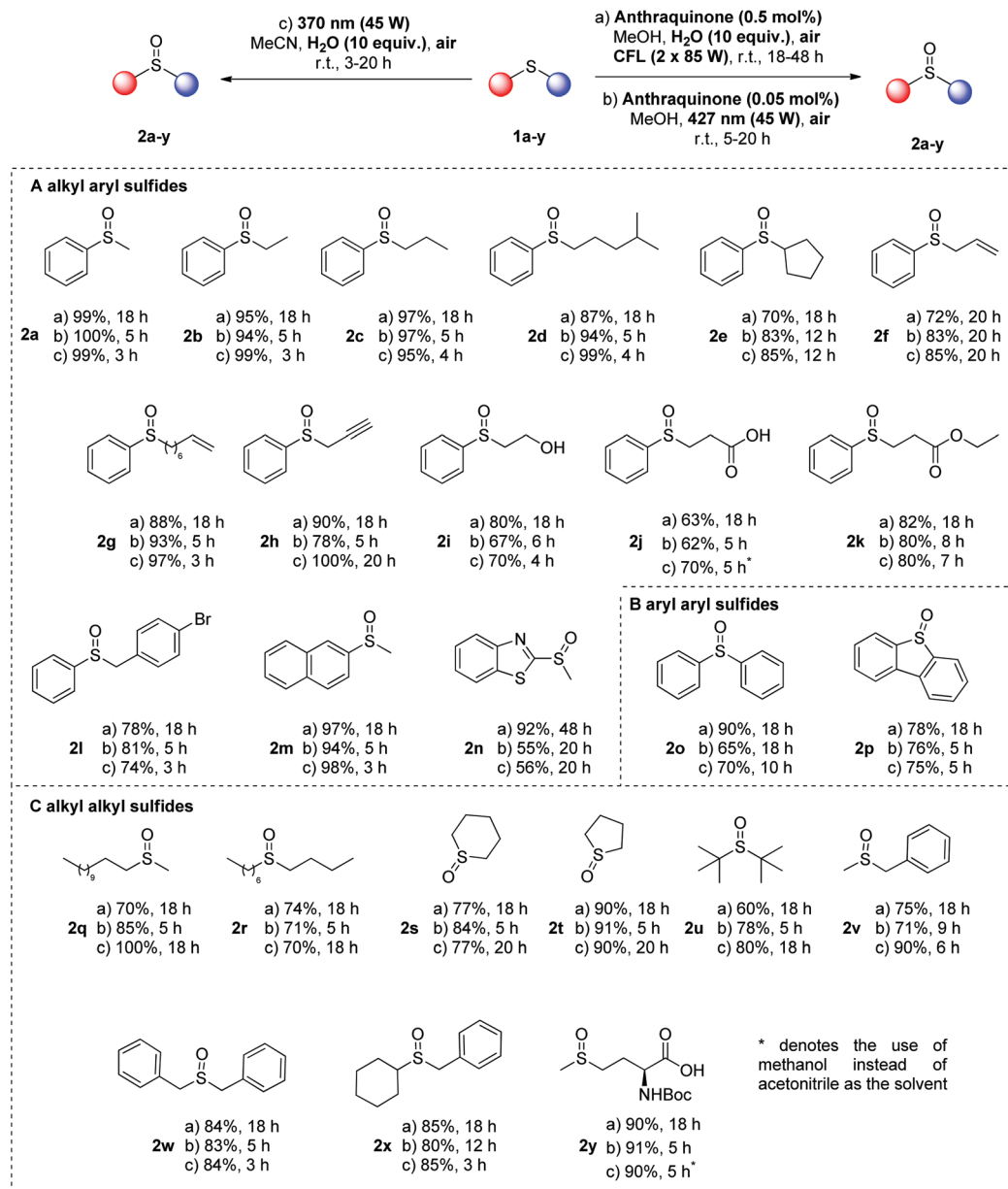
Entry	Sulfide	Solvent, time (h)	1 ^b (%)	2/3 ^b (%)
1 ^c	1a	MeCN, 18	0	29/71
2	1a	MeCN, 2	11	89/0
3 ^d	1a	MeCN, 3	0	99/1
4	1a	MeOH, 2	20	80/0
5	1a	EtOH, 2	11	89/0
6	1a	iPrOH, 2	17	83/0
7	1a	Benzene, 2	93	17/0
8	1a	DMF, 2	72	28/0
9	1q	MeCN, 18	0	100/0
10 ^c	1q	MeCN, 3	0	100/0

^a The reaction was performed with **1a** or **1r** (0.20 mmol) in solvent (1 mL), open air under 370 nm irradiation. ^b Conversion was determined by ¹H-NMR. ^c The reaction was performed with 0.5 mol% anthraquinone (**4a**). ^d 40 μL of H₂O were added.

chosen as a model substrate to apply the optimum reaction conditions (Table 3). We were pleased to observe that our optimum conditions were successfully implemented. The major difference was observed in the reaction time and in order to obtain the desired product **2q**, the reaction time was prolonged from 3 h to 18 h (Table 3, entry 9). For the cases of sensitive substrates, the presence of 0.5 mol% of anthraquinone (**4a**) was used and the desired product was formed quantitatively after 3 h of irradiation at 370 nm (Table 3, entry 10).

With the optimized conditions in hand, we turned our attention in examining the substrate scope (Scheme 3). We decided to follow three different approaches towards the identification of broad and easily-replicated photochemical protocols, using air as the sole oxidant. The first protocol is the catalyst-free approach, using irradiation at 370 nm (45 W, protocol c). In the same time, we opted for an anthraquinone-mediated photochemical reaction, using a low catalyst loading (0.05 mol%) under 427 nm irradiation (45 W, protocol b). Finally, if one cannot have access to LED irradiation source, a CFL-mediated approach was developed, using anthraquinone at 0.5 mol% (protocol a). A variety of sulfides were tested, affording the corresponding sulfoxides in good to excellent yields (Scheme 3). Initially, the easier type of sulfoxides, according to literature, was probed, using alkyl aryl sulfides (Scheme 3A). Same as thioanisole (**1a**), other phenyl-substituted sulfides bearing an alkyl group afforded the corresponding sulfoxides in high to quantitative yields, when the reaction took place under household lamp irradiation for 18 h (Scheme 3, protocol a, **2a–e**). Furthermore, phenyl-substituted sulfides with an alkyl group decorated with different functional groups, *i.e.* double or triple bonds, hydroxy or carboxyl





Scheme 3 Substrate scope.

moiety, afforded in all cases the desired products in good to excellent yields (Scheme 3, protocol a, **2f–k**). Similarly, benzyl phenyl sulfides afforded the corresponding sulfoxides in excellent yields (Scheme 3, protocol a, **2l**). Naphthyl or heteroaryl methyl sulfides were also employed successfully, although prolonged reaction time was required in some cases (Scheme 3, protocol a, **2m** and **2n**). In the case of aryl aryl sulfides, both diphenyl sulfide (**1o**) and fused-aryl sulfide **1p** were tested, leading to high yields of the desired product (Scheme 3, protocol a, **2o** and **2p**). Finally, a broad scope of dialkyl sulfides were employed under the optimum conditions leading to the dialkyl sulfoxides in good to excellent yields (Scheme 3, protocol a, **2q–y**). The obtained lower yields in the cases of dialkyl

sulfoxides **2s**, **2t** and **2u** are attributed to the difficulties posed during the purification process and not to the formation of undesired byproducts (Scheme 3, protocol a).

When the same substrates were employed under LED irradiation (427 nm), the catalyst loading of **4a** can be reduced to 0.05 mol% and the reaction time varied from 5–20 h, while similar results were obtained (Scheme 3, protocol b). In the cases of alkyl aryl sulfides, the reaction time was usually 5–6 h, leading to excellent yields in all cases (Scheme 3, protocol b, **2a–n**). Only in the cases of phenyl secondary alkyl sulfide **1e**, allyl phenyl **1f** and heteroaryl methyl **1n**, an extended reaction time was required (up to 20 h), in order to achieve similar high yields. In the same vein, aryl aryl sulfides led to high yields,



under these conditions, as well (Scheme 3, protocol b, **2o** and **2p**). Finally, dialkyl sulfide **1q-y** led to excellent yields in short reaction time (Scheme 3, protocol b, **2q-y**).

The substrate scope was also examined under catalyst-free conditions upon irradiation at 370 nm (Scheme 3, protocol c). Sulfides bearing an aryl group afforded the corresponding sulfoxides in good to excellent yields in short reaction times (Scheme 3, protocol c, **2a-e**). Furthermore, functional groups, *i.e.* double or triple bonds, hydroxy and carboxyl moiety, were well tolerated, as well as aryl aryl sulfides (Scheme 3, protocol c, **2f-p**). Interestingly, the less reactive dialkyl sulfides under the catalyst-free conditions were converted into sulfoxides in good to excellent yields (Scheme 3, protocol c, **2q-y**). In this case, prolonged reaction times were required, due to the reactivity of the substrates. Furthermore, in all cases, NMR monitoring of the crude reaction mixture did not reveal any byproduct, derived either from overoxidation or from the C-S fragmentation pathway. In the last case of the catalyst-free protocol (Scheme 3, protocol c), for the cases of 3-phenyl-thio-propionic acid (**1j**) and Boc-protected methionine **1y**, the solvent medium was switched to methanol, instead of acetonitrile, due to the insolubility of the substrates in acetonitrile.

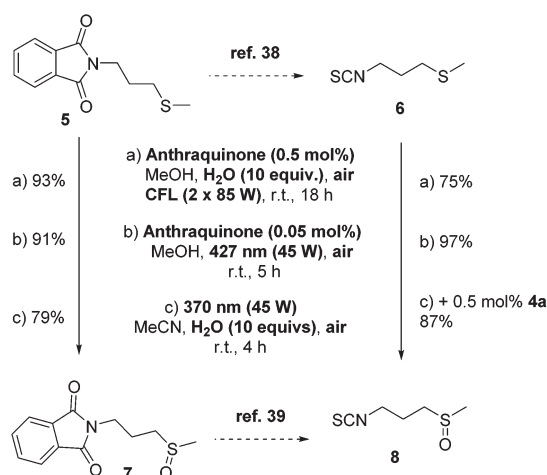
In addition, to demonstrate the synthetic utility of our protocol in a real-life application, we attempted to use our optimized reaction conditions into the oxidation of sulfide intermediates that appear in the synthetic preparation of sulforaphane **8** (Scheme 4). Sulforaphane [4-(methylsulfinyl)butyl-isothiocyanate] **8** was isolated in 1992 by Talalay and coworkers.³⁶ Sulforaphane is the product of the enzymatic hydrolysis of glucoraphanin by the enzyme myrosinase, which is most represented in broccoli. Recently, it has been found to exhibit excellent anti-cancer properties.³⁷ In literature, sulforaphane is synthesized from **5** and in two steps, isothiocyanate **6** is obtained (Scheme 4, top).³⁸ Then, a *m*-CPBA-mediated oxidation leads to sulforaphane **8**.³⁸ Alternatively, sulfoxidation of **5** leads to **7**, which can be converted into sulforaphane **8** (Scheme 4).³⁹ Our photochemical aerobic oxidation protocol

was successfully implemented in the oxidation of sulfides **5** or **6** to afford the corresponding sulfoxides **7** or **8**, respectively, in good to excellent yields (Scheme 4). In the case of **5**, photochemical aerobic oxidation under all three different protocols led to excellent yields of sulfoxide **7**. Similar good yields were obtained in the case of the oxidation of **6** to sulforaphane **8** (Scheme 4). One of the major issues in the preparation of sulforaphane is its decomposition under open air conditions. We came across this problem, when the photooxidation took place under household lamp irradiation, when we attained sulforaphane in a moderate yield. To avoid this issue in our catalyst-free conditions, we added 0.5 mol% of anthraquinone (**4a**) to decrease the reaction time, increasing the yield up to 87%.

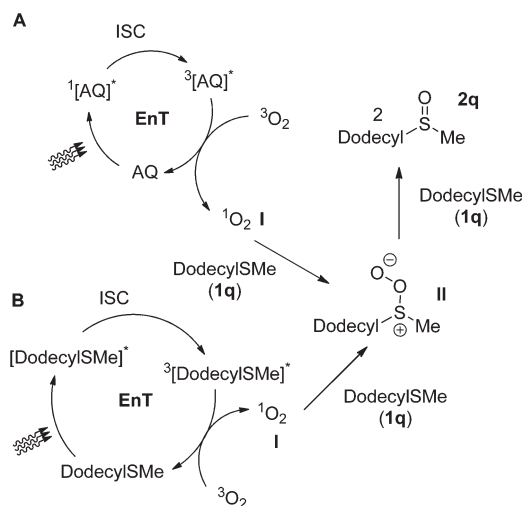
Mechanistic studies

One of the most challenging tasks in photooxidation protocols is the complete understanding of the reaction mechanism. In general, two different mechanistic pathways for the photochemical aerobic sulfide oxidation are reported in literature.²⁴ The first mechanism is promoted by singlet oxygen ($^1\text{O}_2$, **I**), which is generated in the presence of a photosensitizer (Scheme 5A). The generated singlet oxygen mediates the oxygenation of the sulfide to afford persulfoxide intermediate **II** (Scheme 5A). The latter reacts with a second molecule of sulfide to afford two molecules of the desired sulfoxide. In the second mechanistic pathway, a single electron transfer (SET) event occurs between the excited photocatalyst and the sulfide, affording a sulfide radical cation (like **IV** in Scheme 7A). Reaction with either the superoxide radical anion or triplet state oxygen $^3\text{O}_2$ leads to the desired sulfoxide.²⁴

In many literature reports, both mechanisms are proposed to be involved.²⁴ The use of different irradiation sources in our protocols, along with the use of alkyl aryl, aryl aryl or dialkyl sulfides may suggest that different reaction mechanism may



Scheme 4 Photochemical aerobic synthesis of sulforaphane **8**.



Scheme 5 Proposed mechanistic pathways for photochemical aerobic oxidation of dodecyl methyl sulfide (**1q**).



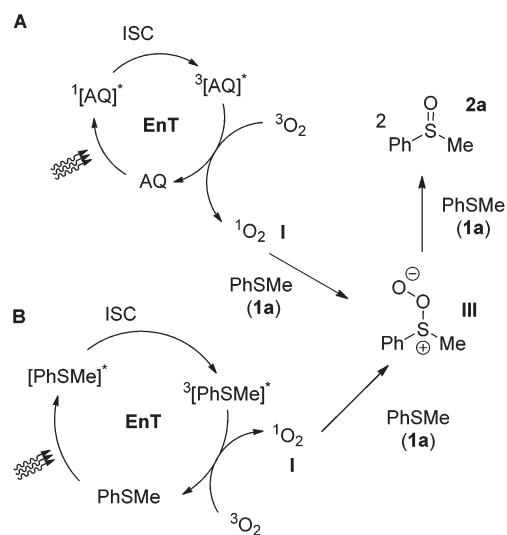
be in place in each case. In general, anthraquinone (**4a**) is known in literature that can generate singlet oxygen *via* energy transfer.⁴⁰ Furthermore, upon excitation, anthraquinone (**4a**) possesses the appropriate potential ($E^{\text{red}} = 1.77 \text{ V vs. SCE}$)⁴¹ to oxidize sulfides ($E^{\text{ox}} = 1.34 \text{ V vs. SCE}$, $E^{\text{ox}} = 1.43 \text{ V vs. SCE}$ and $E^{\text{ox}} = 1.63 \text{ V vs. SCE}$ for thioanisole, diphenyl sulfide and dibutyl sulfide, respectively).⁴² However, fluorescence quenching experiments of excited anthraquinone did not show any quenching either by thioanisole (**1a**) (alkyl aryl sulfide) or dodecyl methyl sulfide (**1q**) (dialkyl sulfide).³⁴ Not following the same trend, diphenyl sulfide (**1o**) (aryl aryl sulfide) does quench excited anthraquinone.³⁴ In addition, it is also reported that sulfides, without the use of an external photocatalyst, upon irradiation, may either generate singlet oxygen or mediate an electron transfer event.³¹

To clarify the accurate mechanistic pathway in a photooxidation protocol, we examined the behavior of three representative examples of sulfides. Thioanisole (**1a**), dodecyl methyl sulfide (**1q**) and diphenyl sulfide (**1o**) were chosen as representative examples of an alkyl aryl sulfide, a dialkyl sulfide and a diaryl sulfide, respectively. First of all, in the absence of oxygen (reaction performed under argon), no oxidation took place.³⁴ Then, a variety of known additives, able to quench either mechanistic pathways, have been reported in literature.²⁴ Initially, to commence our investigations in establishing the exact mechanistic pathway of our protocols, we conducted a series of fluorescence quenching studies.³⁴ For this purpose, we examined the impact of singlet oxygen quenchers, such as sodium azide, DABCO or $\text{Co}(\text{acac})_3$ on the fluorescence of anthraquinone (**4a**).³⁴ Similar experiments were performed, using SET quenchers, such as 1,4-dimethoxybenzene (DMB) or benzoquinone.³⁴ Our studies revealed that among all, benzoquinone and $\text{Co}(\text{acac})_3$ appears to decrease greatly the anthraquinone fluorescence and thus, their use is not indicated for distinguishing the sulfide photooxygenation mechanism, when anthraquinone (**4a**) is employed as the photocatalyst.³⁴ In the same vein, it was found that benzoquinone also quenches the fluorescence of thioanisole (**1a**).³⁴ Thus, the remarkable suppression of the oxidation processes, both in the case of the anthraquinone-mediated oxidation protocols or the catalyst-free conditions by benzoquinone can be attributed to the fluorescence quenching of the catalyst or **1a** and by no means can be assigned to electron transfer events.³⁴ Thus, it is clearly supported that benzoquinone is *not a suitable additive* for the investigation of the anthraquinone-mediated or the catalyst-free thioanisole photooxygenation.³⁴ Similarly, $\text{Co}(\text{acac})_3$ is *not a suitable additive* for the investigation of the anthraquinone-mediated photooxidations.³⁴ On the other hand, DABCO, sodium azide and 1,4-dimethoxybenzene do not interact with anthraquinone.³⁴

We begin with the case of dodecyl methyl sulfide (**1q**), supporting that dialkyl sulfide quenching studies indicate that in both type of conditions (photocatalyst-free and anthraquinone-mediated), the reaction follows the singlet oxygen mechanistic pathway (Scheme 5). Photochemical aerobic oxidation of **1q**, in the presence of DABCO, sodium azide or $\text{Co}(\text{acac})_3$, was totally suppressed (0–5%), while the presence of DMB decreased slightly the yield of the sulfoxide (88%).³⁴

These results indicate that under the photocatalyst-free conditions, the singlet oxygen mechanism is in place (Scheme 5B). Upon irradiation, triplet excited $^3[1q]^*$ generates singlet oxygen **I**, *via* an energy transfer event (Scheme 5B). Singlet oxygen reacts with a ground state **1q**, leading to **II**, which leads to **2q**. In the case of the anthraquinone-mediated photooxygenations, upon 427 nm irradiation, the photochemical aerobic oxidation of **1q**, in the presence of DABCO or sodium azide, was also suppressed, while the presence of DMB, no change in the yield of the sulfoxide was observed (88%).³⁴ These results indicate that under 427 nm irradiation in the presence of anthraquinone, the singlet oxygen mechanism is in place (Scheme 5A). Upon irradiation, triplet excited $^3[AQ]^*$ generates singlet oxygen **I**, *via* an energy transfer event (Scheme 5A). Singlet oxygen reacts with a ground state **1q**, leading to **II**, which leads to **2q**. Upon household lamps irradiation, similar results with DABCO, sodium azide or DMB were obtained.³⁴ Similarly as before, in this case, the energy transfer mechanism (Scheme 5A) is the major pathway.

We then moved to the cases of alkyl aryl sulfide, using thioanisole as the benchmark study. When the reaction is performed under the photocatalyst-free conditions (370 nm), the impact of the substrate into the mechanism of the reaction is more easily distinguished. Photochemical aerobic oxidation of **1a**, in the presence of DABCO, sodium azide or $\text{Co}(\text{acac})_3$, led to diminished yields (12–28%), while the presence of DMB decreased slightly the yield of the sulfoxide (84%).³⁴ These results indicate that under the photocatalyst-free conditions, the singlet oxygen route is dominant (Scheme 6B). In conclusion, thioanisole (**1a**), upon excitation, leads to triplet $^3[1a]^*$, which generates singlet oxygen **I**, through an energy transfer event. Singlet oxygen reacts with a ground state **1a**, leading to **III**, which leads to **2a**. This appears to be the major mechanistic pathway.



Scheme 6 Proposed mechanistic pathways for the photochemical aerobic oxidation of thioanisole (**1a**).



tic pathway of the photocatalyst-free aerobic oxidation of aryl alkyl sulfides at 370 nm (Scheme 6B). This is in accordance with the results of Bonesi and coworkers for the reaction of thioanisoles at 310 nm.³¹ In the case of the anthraquinone-mediated photooxygenations, except from benzoquinone, Co(acac)₃ cannot be employed as a reliable quencher, since it decreases the fluorescence of anthraquinone (**4a**).³⁴ When the anthraquinone-mediated processes were performed in the presence of DABCO or sodium azide (singlet oxygen quenchers), decreased yields (23–31%) were obtained, thus singlet oxygen is involved. Similarly, when the anthraquinone-mediated processes were performed in the presence of DMB (SET quencher), high yields (75–89%) were also obtained, indicating that the singlet oxygen pathway is again the dominant mechanism of action (Scheme 6A).

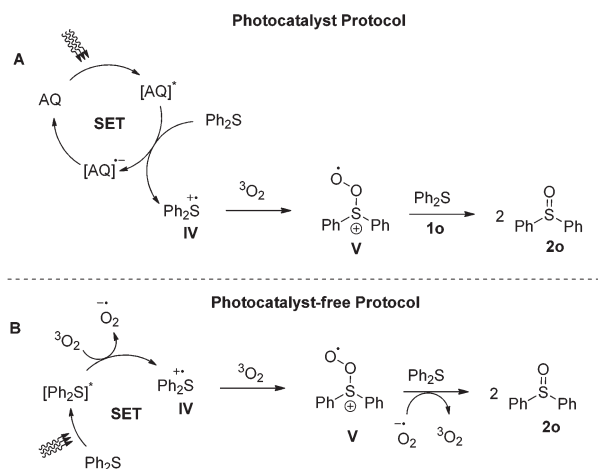
The investigation of the mechanistic pathway when diphenyl sulfide is employed, proved to be rather intricate, since fluorescence quenching studies reveals an interaction between diphenyl sulfide and excited anthraquinone.³⁴ Considering the triplet energy of anthraquinone (E_T 260 kJ mol⁻¹)⁴³ with the E_T of diphenyl sulfide (280 kJ mol⁻¹),⁴⁴ an energy transfer event is unlikely to occur. However, the energy transfer mechanism from anthraquinone to oxygen is viable.⁴⁰ On the other hand, based on the potentials of excited anthraquinone and diphenyl sulfide, (1.77 V vs. SCE and 1.43 V vs. SCE, respectively),⁴² an electron transfer event is more likely to occur. Photocatalyst-free aerobic oxidation of **1o**, in the presence of DABCO, sodium azide or Co(acac)₃, suppressed the reaction (0–8%), while the presence of DMB also suppressed the generation of sulfoxide (6%).³⁴ These results could indicate that under the photocatalyst-free conditions, both mechanistic pathways (singlet oxygen and electron transfer) are involved. However, careful consideration of a variety of parameters is required. The mechanism of the reaction at 370 nm involves excitation of diphenyl sulfide. An electron transfer event takes place, generating sulfide radical cation **IV** (Scheme 7B), which reacts with ground state oxygen to afford the corresponding sulfox-

ide, through intermediate **V** (Scheme 7B). The alternative well-accepted mechanism that leads to the desired sulfoxide involves the reaction of sulfide radical cation **IV** with superoxide radical ion,²⁴ however, for the specific case of diphenyl sulfide, Bonesi and coworkers have proven through kinetic studies that superoxide radical anion does not react directly with **IV**, but plays a secondary role in the process, more likely mediating the conversion of **V** to **2o**.³¹ Also, the energy transfer mechanism from excited **1o** to oxygen, in order to generate singlet oxygen is also not operative, as Bonesi and coworkers has also proven for the case of **1o**.³¹ However, our quenching experiments with DABCO or sodium azide delivered minimum yields of the reaction.³⁴ Herein, it is necessary to highlight the need to be fully be aware of the oxidation potentials of the compounds involved in the process. Taking into consideration the potential of diphenyl sulfide ($E^{ox} = 1.43$ V vs. SCE),⁴² as well as the potentials of DABCO ($E^{ox} = 0.57$ V vs. SCE)⁴⁵ and sodium azide ($E^{ox} = 1.29$ V vs. NHE,⁴⁶ ~approx. 1.05 V vs. SCE), then it can easily be deduced that the reduced yields observed in the presence of DABCO or sodium azide are due to redox reactions between sulfide radical cation **IV** with either DABCO or NaN₃ and not due to quenching of singlet oxygen. The singlet oxygen pathway is also not viable, since it is known that product **2o** constitutes a quencher of persulfoxide intermediates (like intermediates **II** or **III** of Schemes 5 and 6).²⁴ In the photocatalyst-mediated protocols, the reaction in the presence of DABCO or sodium azide or DMB, was suppressed (0–19%).³⁴ As before though, upon anthraquinone irradiation, an electron transfer event occurs, generating sulfide radical cation **IV**, which reacts with oxygen to afford the corresponding sulfoxide, through intermediate **V** (Scheme 7A). All other potential mechanisms are not viable, due to the same reasons.

Based on the literature, Cismesia and Yoon employed the quantum yield as a mechanistic tool.⁴⁷ Using potassium ferrioxalate as the actinometer, we calculated the quantum yield of the thioanisole photooxidation under the anthraquinone-mediated conditions.³⁴ The achieved quantum yields ($\Phi = 2.0$ and 2.1 for 427 nm and CFL irradiation) are impressive, since the maximum value based on the proposed mechanism is 2.0, regardless the pathway followed.

Conclusions

Photochemical aerobic oxidations merge two interesting fields, photochemistry, and air-mediated processes. Although for many years, both areas received limited attention, with the uprise of modern synthetic photochemistry and increasing number of literature reports on oxygen-mediated oxidations, the last five years, the photochemical aerobic oxidation of sulfides to sulfoxides has attracted exponential attention. Herein, we provide a general, fast, mild, green and easy-to-operate procedure to perform these reactions. Initially, we explored the role of wavelength irradiation towards sulfides aerobic photooxygenation to the corresponding sulfoxides. Compared to all the previous methods presented in the literature, two low-cata-



Scheme 7 Proposed mechanistic pathways for the photochemical aerobic oxidation of diphenyl sulfide (**1o**).



lyst loading (0.05–0.5 mol%) anthraquinone-mediated protocols (CFL lamps or 427 nm) and one photocatalyst-free aerobic protocol (370 nm) are reported, combining short reaction time and overcome the unwanted overoxidation reaction. These conditions outperform most known literature procedures as far as reaction time and catalyst loading are concerned. Furthermore, expensive photocatalysts or complicated non-natural occurring catalysts, which are commonly employed in sulfide oxidation, are replaced by a cheap and commercially available photocatalyst, like anthraquinone. A broad scope of substrates was successfully tested, under the optimum conditions. We also applied our photooxidation protocols towards the synthesis of Sulforaphane, a promising anti-cancer agent. The mechanism of the reaction was extensively studied and for this purpose a series of quenching studies were performed, and we herein provide a flowchart for future researchers. Firstly, after UV-Vis studies, the potential of developing a photocatalyst-free protocol should be defined. When the appropriate photocatalyst is identified, *via* literature reports, the potential mechanistic scenarios should be identified. Utilizing fluorescence quenching studies (photocatalyst with substrates, photocatalyst with quenchers and sulfides with quenchers), the appropriate probes (quenchers) can be identified, in order to correctly recognize which probes can be used and which cannot, since they provide false positive or negative results. In this case, for the photocatalyst-free process, *via* the appropriate quenchers, we identified that dodecyl methyl sulfide (dialkyl sulfides) are oxidized *via* singlet oxygen, while when anthraquinone is employed, the singlet oxygen pathway is the dominant. In the case of thioanisole (alkyl aryl sulfides), in the photocatalyst-free protocol and under anthraquinone-mediated protocols, the singlet oxygen mechanism is the major pathway. Finally, in the case of diphenyl sulfide (aryl aryl sulfides), in the photocatalyst-free protocol and under the use of anthraquinone, the sulfide radical cation mechanism is operative.

Author contributions

C. G. K conceived the study, E. S., P. L. G., N. F. N. and C. G. K. designed the experiments and analyzed the data. E. S., P. L. G. and N. F. N. performed the experiments. P. L. G. and C. G. K. prepared the draft of the manuscript and C. G. K. performed the final editing. The manuscript was written through contributions of all authors.

Conflicts of interest

There are no conflicts to declare.

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