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A sustainable photochemical aerobic sulfide oxidation: access to sulforaphane and modafinil†

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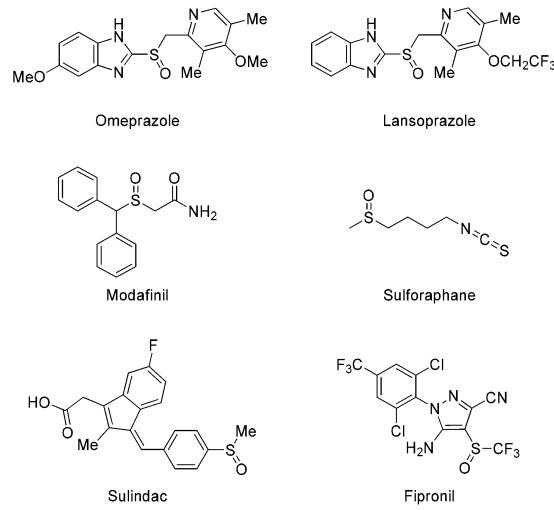
Sulfoxide-containing molecules are an important class of compounds in the pharmaceutical industry and many efforts have been made to develop new and green protocols, targeting the chemoselective transformation of sulfides into sulfoxides. Photochemistry is a rapidly expanding research field employing light as the energy source. Photochemical aerobic processes possess additional advantages to photochemistry and may find applications in the chemical industries. Herein, a 370 nm catalyst-free aerobic protocol was developed, using 2-Me-THF as the green solvent. At the same time, two low-catalyst-loading anthraquinone-based processes (under a CFL lamp or 427 nm irradiation) in 2-Me-THF were developed. Furthermore, a broad range of substrates was tested. We also implemented our protocols towards the synthesis of the pharmaceutical active ingredients (APIs) sulforaphane and modafinil.

Introduction

Historically, sulfoxides comprise an important class of molecules in the pharmaceutical industry, beginning with the rise of antibacterial ingredients at the beginning of the 20th century.¹ A number of sulfoxide-containing molecules are known to display an excellent pharmacological profile and are marketed as drugs, including omeprazole, lansoprazole, modafinil, sulforaphane, sulindac and fipronil (Scheme 1). Omeprazole and lansoprazole are proton-pump inhibitors (PPIs) and are widely used in pharmaceuticals treating gastric acid hypersecretion.² Modafinil, another sulfoxide-containing drug, is used for the treatment of narcolepsy symptoms.³ Sulforaphane is a naturally occurring product, found in broccoli and Brussels sprouts, that has been proven to possess anti-cancer properties.⁴ Sulindac acts as a prodrug, which is metabolised in the human body to the corresponding sulfide, and it is well known as an anti-inflammatory and analgesic agent,⁵ while fipronil is an insecticide, used to protect crops against specific soil and foliar pests.⁶

Over the past years, many researchers worldwide, both in industry and in academia, have put great effort into developing

a variety of industrially applicable oxidation methodologies, converting sulfides into the corresponding sulfoxides. In most of the protocols reported in the literature, hydrogen peroxide is the oxidant which holds the lion's share.⁷ The use of peroxides in general is highly beneficial due to their availability, low cost, and benign by-products. The major drawback of the use of peroxides appears to be their activation, which in most cases requires either heat or metal catalysts. Another widely used class of oxidants in the pharmaceutical industry is organic peracids,⁸ such as *m*-chloroperbenzoic acid (*m*-CPBA). The latter has been exploited in numerous sulfoxide synth-



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eses,⁹ but is lacking chemoselectivity. In all the reported sulfide syntheses, over-stoichiometric amounts of oxidants are needed to increase the reaction yield. As a consequence, in many cases, sulfide overoxidation may occur, leading to sulfone by-products, which in the case of an active pharmaceutical ingredient (API) is restrictive.

In 2004, Avrutov and coworkers described the synthesis of Lansoprazole using *tert*-butyl hydroperoxide (TBHP) activated by a vanadium catalyst (Scheme 2A).¹⁰ In their patent application, the employed oxidation system was used to suppress the formation of the sulfone by-product.¹⁰ The use of peracids towards sulfide oxidation was extensively studied in the synthesis of Cephalosporins as well (Scheme 2B).¹¹ The major drawback in their use, apart from the usual safety issues, is the need to purge the resulting organic acid, which in many cases is a difficult task in a manufacturing plant. To avoid this difficulty, inorganic oxidants were employed. In 2001, omeprazole was proposed to be produced using sodium percarbonate and a molybdenum catalyst (Scheme 2C).¹²

Another mild oxidant that lately has been widely employed in oxidation protocols is oxygen.¹³ It is an abundant and ambient gas, considered the "greenest" option for oxidation reactions. Oxygen activation can occur *via* metal catalysis¹⁴ or organocatalysis.¹⁵ In 2020, Zografas and coworker proposed the use of proline-based organocatalysts for the aerobic oxidation of sulfides to sulfoxides, using HFIP as the solvent.¹⁶ However, expensive metal-based catalysts¹⁴ or expensive and hazardous solvents (*e.g.* HFIP)¹⁶ increase significantly the

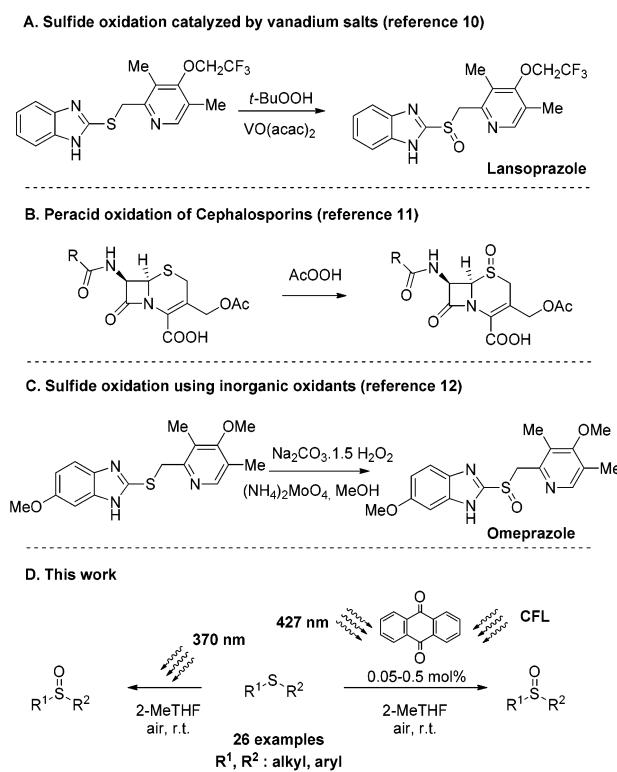
manufacturing cost, while the final purification of the product can be troublesome.

In 2008, MacMillan,¹⁷ Stephenson¹⁸ and Yoon¹⁹ relaunched an exciting and powerful area of research called photochemistry.^{20,21} This partially forgotten and rather misunderstood research field very quickly created new opportunities, unlocking many reactions not easily accessed using the conventional ground state pathways. The merger of photochemical protocols with aerobic (oxygen)-mediated oxidations was recognized quite early and the oxidation of sulfides to sulfoxides was first studied in the 1960s.^{22,23} Throughout the years, the photochemical aerobic oxidation of sulfides received some attention, but the use of specialized apparatus for irradiation or the use of heterogeneous and in some cases homogeneous catalysts (metal-based or organocatalysts) limited its applicability.²³ However, since 2017, the emerging field of photochemistry merged with organocatalysis has drawn the interest of many researchers into studying photochemical aerobic sulfide oxidation.²⁴

In our group over the past years, we have been focused on developing novel light-driven methodologies.²⁵ Recently, we reported a mild, green and chemoselective direct photochemical aerobic oxidation of sulfides to the corresponding sulfoxides.²⁶ Our experience in developing operationally simple and safe protocols easily applied in industrial scale productions²⁷ led us to upgrade our methodology to a sustainable and industry-friendly protocol. Herein, we report environmental, novel, green and sustainable protocols for the oxidation of sulfides using a mild oxidant, such as molecular oxygen, in the presence of a low-cost commercially available photocatalyst. The reaction medium comprises the use of a biomass-based solvent, serving the basic principles of green chemistry, which is highly desirable in the chemical and pharmaceutical industries (Scheme 2D).

Results and discussion

In early 2022, we reported a general, mild and green protocol to convert sulfides into sulfoxides.²⁶ Firstly, we explored the role of wavelength irradiation and proposed two low-catalyst-loading (0.05–0.5 mol%) anthraquinone-mediated methods (CFL lamps or 427 nm) and one photocatalyst-free protocol (370 nm) for the photochemical aerobic oxidation of sulfides into sulfoxides, having the advantage of short reaction times and no overoxidation reactions. Having in our minds the importance of this transformation in the pharmaceutical industry, we envisaged an expansion of those protocols to gain a more industrial and sustainable character. Initially, we turned our attention to the nature of the reaction medium, since the industrial character of any process is directly linked to the amount of waste produced (Table 1). We examined the anthraquinone-mediated reaction output of thioanisole (**1a**) oxidation under CFL irradiation in a variety of green solvents (Table 1).²⁸ Protic solvents, *i.e.* isopropanol or glycerol, proved to be sluggish (Table 1, entries 1 and 2), while in water the



Scheme 2 Sulfide oxidation in API synthesis.

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

Entry	Lamp (nm)	Solvent, time (h)	4 (mol%)	1a ^b (%)	2a/3a ^b (%)	1a	2a	3a	
						Ph-S-Me	x mol%	Ph-S(=O)-Me	Ph-S(=O ₂)-Me
1	CFL	Glycerol, 18	0.5	75	25/0				
2	CFL	i-PrOH, 18	0.5	88	12/0				
3	CFL	H ₂ O, 18	0.5	100	0/0				
4	CFL	EtOAc, 18	0.5	95	5/0				
5	CFL	GVL, 18	0.5	75	25/0				
6	CFL	Cyrene, 18	0.5	100	0/0				
7	CFL	MTBE, 18	0.5	86	14/0				
8	CFL	2-Me-THF, 18	0.5	0	70/30				
9 ^c	CFL	2-Me-THF, 18	0.5	0	99/1				
10 ^d	427	Glycerol, 3	0.05	94	6/0				
11 ^d	427	i-PrOH, 3	0.05	79	21/0				
12 ^d	427	H ₂ O, 3	0.05	91	9/0				
13 ^d	427	EtOAc, 3	0.05	11	89/0				
14 ^d	427	GVL, 3	0.05	61	39/0				
15 ^d	427	Cyrene, 3	0.05	100	0/0				
16 ^d	427	2-Me-THF, 3	0.05	1	99/0				

^aThe reaction was performed with **1a** (0.20 mmol) in solvent (1 mL), under open air irradiation for 18 h. ^bConversion was determined by ¹H-NMR. ^cThe reaction was performed with the addition of H₂O (40 μ L). ^d0.5 mL of solvent was used instead of 1 mL.

reaction did not occur due to anthranquinone's insolubility (Table 1, entry 3). In solvents bearing a carbonyl group, *i.e.* ethyl acetate or γ -valerolactone (GVL), the reaction was sluggish, while trying to use a novel biobased solvent called Cyrene, the reaction did not take place (Table 1, entries 4–6). Turning to ethereal solvents, *i.e.* *t*-butyl methyl ether, afforded poor results (Table 1, entry 7), while 2-methyltetrahydrofuran (2-Me-THF) was the solvent that outperformed all the others (Table 1, entry 8). 2-Me-THF is a commercially available solvent, derived from furfural or levulinic acid, both products being easily derived from renewable lignocellulosic biomass.²⁹ 2-Me-THF has a strong chemical resemblance to tetrahydrofuran, but it features different physical properties. Its low miscibility with water and its good stability to acids and bases, compared with THF, are some of the advantages offered.³⁰ Thus, its use in the pharmaceutical industry is highly appreciated. The addition of H₂O (10 equiv.) was essential for the acceleration of the reaction rate and the suppression of the formation of the sulfone by-product (Table 1, entry 9 *vs.* entry 8).³¹ Due to hydrogen bond formation, water inhibits the conversion of sulfoxide to sulfones.³² In our previous study under LED irradiation (427 nm), photochemical aerobic sulfide oxidation led to the selective formation of the corresponding sulfoxide *vs.* sulfone after 5 h of irradiation in the presence of 0.05 mmol% of **4**.²⁶ When the reaction was performed in

2-Me-THF, the desired product was afforded in quantitative conversion, suppressing the undesired sulfone by-product, while the reaction time was reduced to 3 h (Table 1, entry 16). Apart from ethyl acetate, which afforded high sulfoxide conversion (Table 1, entry 13), all the other green solvents mentioned above afforded lower yields (Table 1, entries 10–15). Indeed, 2-Me-THF was the solvent of choice, reducing the reaction time at the same time, in comparison with our previous results.²⁶

Next, we turned our attention to the catalyst-free protocol for the photochemical aerobic oxidation of **1a** (Table 2). To our delight, when the reaction was performed under UVA-LED irradiation (370 nm) in 2-Me-THF, quantitative formation of the desired product was obtained after 2 h, under catalyst-free conditions (Table 2, entry 8). Again, this is a slightly decreased reaction time compared with our previous studies (2 h *vs.* 3 h).²⁶ Quantitative formation of the desired sulfoxide was also obtained when GVL or water was used as the solvent (Table 2, entries 3 and 5); however, lower yields were obtained upon isolation (60% and 68%, respectively). Furthermore, GVL and water might pose difficulties when the catalyst-free protocol is used in the pharmaceutical industry. The use of GVL will add an extra purification step to the procedure, which raises the manufacturing cost, while the use of water as the reaction medium will narrow down the substrate scope, excluding intermediates bearing easily hydrolysed functional groups. Again, solvent optimization did not reveal any medium leading to better yields, compared with 2-Me-THF (Table 2, entries 1, 2, 4, 6, 7 *vs.* 8).

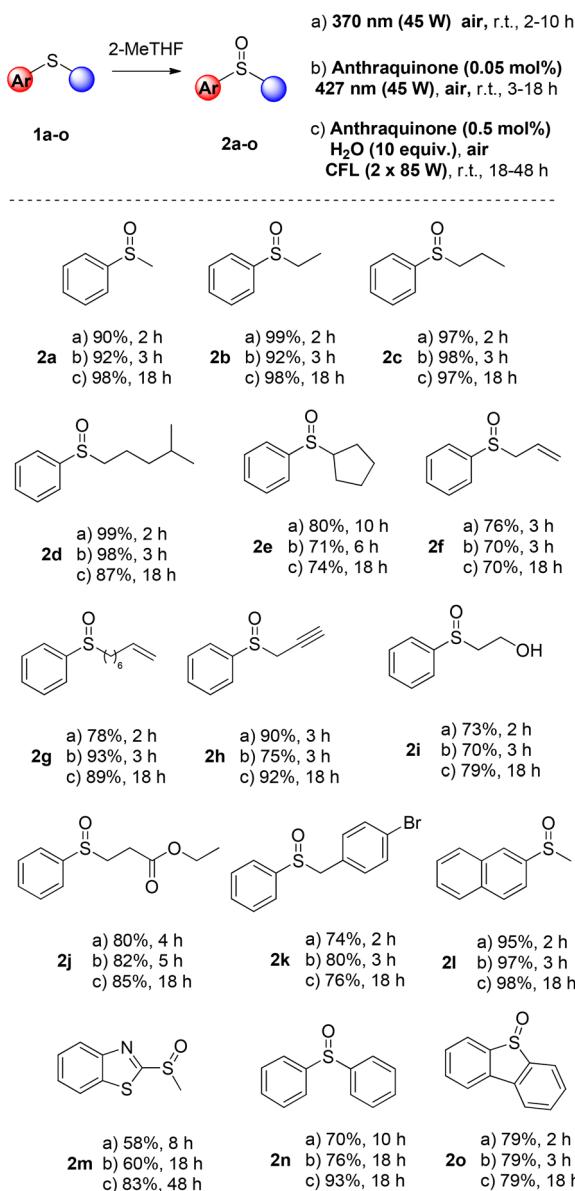
Having found the optimum reaction conditions, we studied the substrate scope (Scheme 3). We probed all three approaches, using air as the sole oxidant. The first process is the catalyst-free approach, using irradiation at 370 nm in 2-Me-THF (45 W, protocol A). Also, the anthraquinone-mediated photochemical reaction, using a low catalyst loading (0.05 mol%) under 427 nm irradiation, was examined (45 W, protocol B). Finally, a CFL-mediated approach in 2-Me-THF was envisaged, using

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

Entry	Solvent	1a ^b (%)	2a/3a ^b (%)	1a	Solvent, 2 h	370 nm, r.t.	2a	3a
				Ph-S-Me	x mol%	Ph-S(=O)-Me	Ph-S(=O ₂)-Me	
1	Glycerol	72	28/0					
2	Isopropanol	17	83/0					
3	H ₂ O	0	100/0					
4	EtOAc	80	20/0					
5	GVL	0	100/0					
6	Cyrene	100	0/0					
7	MTBE	87	13/0					
8	2-Me-THF	0	100/0					

^aThe reaction was performed with **1a** (0.20 mmol) in solvent (1 mL), in open air under 370 nm irradiation for 2 h. ^bConversion was determined by ¹H-NMR.



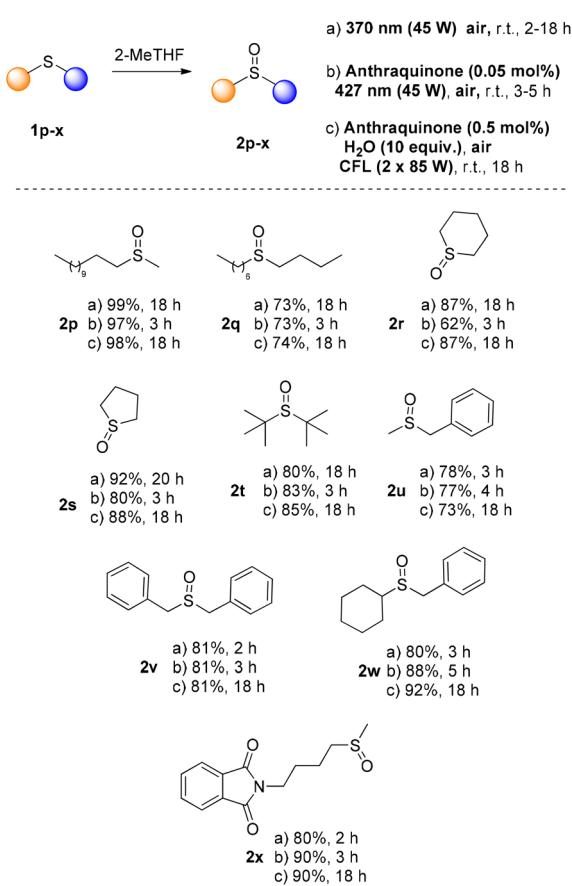


Scheme 3 Substrate scope: aryl or heteroaryl substituted sulfides.

anthraquinone at 0.5 mol% (protocol C). Initially, we examined a variety of alkyl aryl sulfides, employing catalyst-free conditions upon irradiation at 370 nm. The oxidation step proceeded smoothly, affording the corresponding sulfoxides in good to excellent yields (Scheme 3). Along with thioanisole (**1a**), other phenyl-substituted sulfides having an alkyl moiety afforded the desired sulfoxides in high to quantitative yields (Scheme 3, protocol A, **2a–e**). Aryl or substituted aryl sulfides with an alkyl group having various functional groups, *i.e.* double or triple bonds, hydroxy or ester moieties, afforded in all cases the desired products in good to excellent yields (Scheme 3, protocol A, **2f–j**). Phenyl benzyl, naphthyl methyl, heteroaryl methyl, diaryl or fused aryl sulfides also afforded the corresponding sulfoxides in good to excellent yields

(Scheme 3, protocol A, **2k-o**). Additionally, NMR monitoring of the crude reaction mixture did not reveal any by-product (either from overoxidation or from the C-S fragmentation).

When the same substrates were subjected to LED irradiation (427 nm), the presence of a catalyst was required. The catalyst loading of **4** was 0.05 mol% and the reaction time varied from 3 to 18 h (Scheme 3, protocol B). When alkyl aryl sulfides were employed, the reaction time was usually 3 h, while good to excellent yields were obtained in all cases (Scheme 3, protocol B, **2a–o**). Deviation in the reaction time was observed in the cases of phenyl secondary alkyl sulfide **1e**, alkyl phenyl sulfide bearing an ester moiety (**1j**), heteroaryl methyl (**1m**) and diphenyl sulfide (**1n**). A prolonged reaction time was required (up to 18 h) for the reaction to take place. In the cases of sulfides **1f** and **1h**, no oxidation by-products on the double or the triple bond were detected. However, if a higher catalyst loading of **4** was employed, then a mixture of other oxidized products was obtained in both cases. Similarly, under the optimised reaction conditions, in the case of **1m**, no other oxidation product was isolated. Next, we examined the substrate scope under CFL irradiation, increasing the catalyst loading (0.5 mol%) (Scheme 3, protocol C). Alkyl aryl sulfides afforded the corresponding sulfoxides in good to excellent yields in 18 h (Scheme 3, protocol C, **2a–e**). Furthermore, func-



Scheme 4 Substrate scope: dialkyl sulfides.

tional groups, like double or triple bonds, hydroxy and ester moieties or aryl aryl sulfides were well tolerated (Scheme 3, protocol C, 2f-o).

To expand the industrial feasibility of our process, we also performed the photooxidation of thioanisole **1a**, replacing the CFL lamps with sunlight (April 27, 2022, 10:00–14:00, Athens Greece, 37.97° N, 23.72° E, temperature range 23–25 °C), leading to a similarly excellent yield (97%) after 4 h. We also checked the scalability of our protocol, performing a gram-scale reaction (June 25, 2022, 09:00–17:00, Athens Greece, 37.97° N, 23.72° E, temperature range 28–35 °C), leading to an excellent yield (82%) after 8 h.³¹ Since the environmental conditions vary from time to time, we also performed aerobic photooxidation, applying our photocatalyst-free protocol and increasing the reaction scale (10 g). The reaction time was prolonged and 2 × LED 370 nm lamps were employed; however the formation of the corresponding sulfoxide was excellent (81% yield) after 13 h.³¹

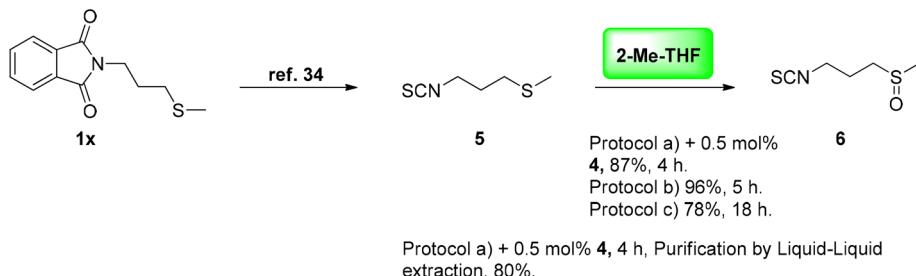
Next, we examined our protocols in a broad range of dialkyl sulfides (Scheme 4). In this case (for the 370 nm protocol), prolonged reaction times were required (18 h) due to the reactivity of the substrates. Dialkyl sulfoxides were easily accessed in good to excellent yields when the photocatalyst-free aerobic oxidation protocol in 2-Me-THF was employed (Scheme 4, protocol A, 2p-x). The anthraquinone-mediated protocols were also applied to sulfide photooxidation, affording in all cases the corresponding sulfoxides in excellent yields. Upon CFL irradiation and using 0.5 mol% anthraquinone, the photooxi-

dation reaction time was 18 h. Altering the irradiation source, using LED irradiation (427 nm), the amount of anthraquinone was 0.05 mol% and the oxidation reaction was completed after 3–5 h.

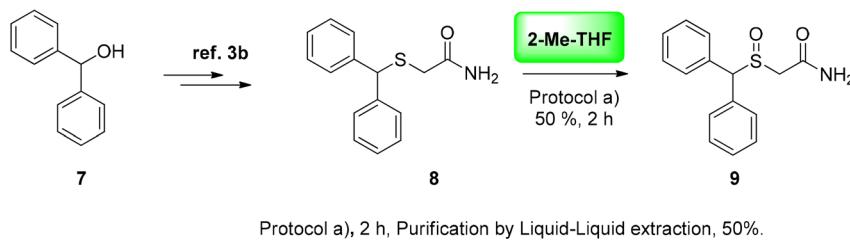
Synthetic application

Our desire to demonstrate the synthetic utility of our protocol in real-life applications led us to use our optimized reaction conditions in the oxidation of sulfide intermediates that appear in pharmaceutical active ingredients, *i.e.* sulforaphane and modafinil (Scheme 5). In 1992, Talalay and coworkers isolated sulforaphane [4-(methylsulfinyl)butylisothiocyanate] **6**.³³ Sulforaphane is the product of the enzymatic hydrolysis of glucophanin, which is found in the Brassicaceae family and in broccoli, in particular, and was recently found to exhibit anti-cancer properties.⁴ Recently, sulforaphane was also recognized as a potential inhibitor of the *in vitro* replication of SARS-CoV-2.³⁴ Sulforaphane interacts synergistically with remdesivir, inhibiting coronavirus infection *in vitro*. We commenced our synthesis from intermediate **1x**, which can be easily synthesized from 1,4-dibromobutane in two steps.³⁵ Deprotection of the phthalimide group and subsequent treatment of the corresponding amine with carbon disulfide afforded isothiocyanate **5**, which is the most common intermediate for sulforaphane synthesis (Scheme 5, top).³⁵ Applying our photochemical aerobic oxidation protocols in the oxidation of sulfide **5** afforded sulforaphane in good to excellent yields (Scheme 5, top). Contrary to all the reported

Sulforaphane Synthesis



Modafinil Synthesis



a) 370 nm (45 W) air, r.t

b) Anthraquinone (0.05 mol%)
427 nm (45 W), air, r.t.

c) Anthraquinone (0.5 mol%)
H₂O (10 equiv.), air
CFL (2 × 85 W), r.t.

Scheme 5 Synthetic application: late-stage photochemical aerobic oxidation for the synthesis of sulforaphane and modafinil.

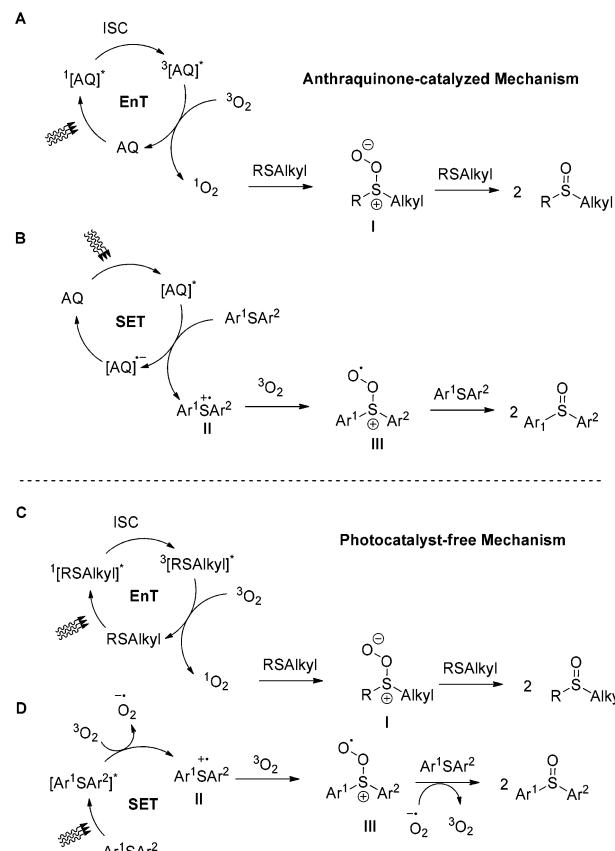


methods described in the literature, where the oxidation was performed either using hydrogen peroxide or *m*-CPBA, the sulfone derivative was not detected. Applying our photocatalyst-mediated protocols, the reaction proceeded smoothly, affording sulforaphane in good to excellent yields. Under LED irradiation using (427 nm), sulforaphane was obtained in 96% in 5 h. One of the major issues in the preparation of sulforaphane is its decomposition under open air. We faced this problem when the photooxidation took place under household lamp irradiation, when we attained sulforaphane in a moderate yield (78%). To circumvent this problem in our catalyst-free conditions, we employed 0.5 mol% of anthraquinone (**4**) to minimize the reaction time, increasing the yield to 87%. To avoid purification by column chromatography, a liquid–liquid extraction–purification afforded sulforaphane in 80% yield (reaction performed at 370 nm).³¹ Thus, we believe this constitutes an even more industry-friendly approach for the synthesis of sulforaphane.

Our initial intention in developing an industrially applicable photochemical protocol led us to search for more challenging substrate targets. For this reason, we questioned whether our photochemical protocols could be applied towards the synthesis of modafinil (Scheme 5, bottom). Modafinil **9** acts as a CNS stimulant, which promotes vigilance and wakefulness without any side effects.^{3b} It is generally used for the treatment of narcolepsy and ADHD. As presented in the literature, sulfide **8** appears to be the most common intermediate for the synthesis of modafinil.^{3b} Sulfide **8** contains an active benzylic position, which poses some difficulties in applying a chemoselective photochemical oxidation protocol. Indeed, when we examined the application of our photochemical protocols in the oxidation of sulfide **8**, we observed that in both anthraquinone-mediated protocols the decomposition of sulfide **8** took place. Probably, the photocatalyst-mediated process initiates a C–S fragmentation in the highly reactive benzylic position. Careful monitoring of the reaction showed that the parasitic decomposition pathway is much faster than the oxidation one, since we did not observe any product formation. On the contrary, under catalyst-free conditions, the photochemical aerobic oxidation of sulfide **8** proceeded smoothly, affording the desired product in 50% yield. Similar to before, a liquid–liquid extraction–purification was also possible.

Mechanistic studies

The most intriguing part of photooxidation is understanding the reaction mechanism. In the literature, two different mechanistic pathways for photochemical aerobic sulfide oxidation are reported.²³ In the first mechanism singlet oxygen, $^1\text{O}_2$, is generated using a photosensitizer (Scheme 6). Singlet oxygen performs the oxygenation of the sulfide, leading to per-sulfoxide intermediate **I** (Scheme 6). This intermediate reacts with another sulfide, affording two molecules of the desired sulfoxide. In the second mechanistic pathway, a single electron transfer (SET) occurs between the excited photocatalyst and the sulfide, affording sulfide radical cation **II** (Scheme 6).



Scheme 6 Proposed mechanistic pathways for the photochemical aerobic oxidation of sulfides.

Reaction with either the superoxide radical anion or triplet state oxygen $^3\text{O}_2$ leads to the sulfoxide. In our work, the use of different irradiation sources in our protocols, along with the use of alkyl aryl, aryl aryl or dialkyl sulfides, may suggest that a different reaction mechanism may be in place in each case. In general, anthraquinone (**4a**) is known from the literature to be able to generate singlet oxygen *via* energy transfer.³⁶ In our previous studies, we performed a detailed study of the reaction mechanism.²⁶ Utilizing fluorescence quenching studies and the appropriate quenchers shed light on the reaction mechanism.²⁶ In the photocatalyst-free process (370 nm), *via* the appropriate quenchers, we identified that dodecyl methyl sulfide (dialkyl sulfides) are oxidized *via* singlet oxygen (Scheme 6C), while when anthraquinone is employed (427 nm or CFL), the singlet oxygen pathway is dominant (Scheme 6A). In the case of thioanisole (alkyl aryl sulfides), in the photocatalyst-free protocol (370 nm), the singlet oxygen mechanism is in place (Scheme 6C). Under the use of anthraquinone (427 nm or CFL), the singlet oxygen pathway generated by anthraquinone is followed (Scheme 6A). Finally, in the case of diphenyl sulfide (aryl aryl sulfides), in the photocatalyst-free protocol (370 nm), the sulfide radical cation pathway is in place (Scheme 6D). With the use of anthraquinone (427 nm or CFL), the sulfide radical cation is produced (Scheme 6B).



Conclusions

The basic principles of green chemistry have been known since 1990, but real application in the pharmaceutical industry is rather limited. During the last decade, the photochemical aerobic oxidation of sulfides to sulfoxides has attracted exponential amounts of attention. Herein, we married our previously developed photochemical oxidation protocols with the basic principles of green chemistry, providing a general, fast, mild, green and easy-to-operate industry-friendly procedure to perform these reactions. We exploited 2-Me-THF, a commercially available green solvent, derived from furfural or levulinic acid, both products being easily derived from renewable lignocellulosic biomass, in our low-catalyst-loading (0.05–0.5 mol%) anthraquinone-mediated protocols (CFL lamps or 427 nm) and in our photocatalyst-free aerobic protocol (370 nm). Among the different protocols developed, the photocatalyst-free aerobic protocol (370 nm) appears to be the most efficient, since it is fast, it can be scaled up and it can be easily applied to more challenging substrates bearing easily oxidized C–H bonds; see the case of modafinil. However, the 427 nm protocol can be employed in cases where organic compounds bear chromophores below 400 nm, since in this case, they do not interfere with the oxidation process. Finally, the cheapest alternative is the CFL-mediated procedure. A broad range of substrates was successfully tested under the optimum conditions. In some cases, the reaction proved to be faster than those performed in common organic solvents. We also applied our photooxidation protocols towards the synthesis of two APIs: sulforaphane, a promising anti-cancer agent and a potential SARS-CoV-2 inhibitor, and modafinil, a drug used to treat narcolepsy symptoms. We sincerely hope these processes can find application in real life.

Author contributions

C. G. K. conceived the study, E. S., P. L. G. and C. G. K. designed the experiments and analyzed the data. E. S. and P. L. G. 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 from all authors.

Conflicts of interest

There are no conflicts to declare.

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