

# Photooxidation of 4-chlorophenol sensitised by iron *meso*-tetrakis(2,6-dichloro-3-sulfophenyl)porphyrin in aqueous solution †

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The photosensitised degradation of 4-chlorophenol (4-CP) by iron *meso*-tetrakis(2,6-dichloro-3-sulfophenyl)porphyrin (FeTDCPPS) has been studied in aerated aqueous solution, and is shown to lead to formation of *p*-benzoquinone (BQ) and *p*-hydroquinone (HQ) as main photoproducts. In deaerated solution no *p*-benzoquinone was formed. The photolysis products were identified by high performance liquid chromatography (HPLC) and UV-visible spectroscopy. The photodegradation in aerated solution was also carried out in the presence of sodium azide (NaN<sub>3</sub>) as a singlet oxygen [<sup>1</sup>O<sub>2</sub>(<sup>1</sup>Δ<sub>g</sub>)] quencher, and showed a significant decrease in the rate of photolysis, suggesting under these conditions, that Type II sensitisation is one of the dominant mechanisms of 4-CP degradation. Support for this comes from laser flash photolysis and time-resolved singlet oxygen phosphorescence measurements. However, these also show direct reaction between the excited porphyrin and 4-CP, indicating that there are two mechanisms involved in the chlorophenol photodegradation.

## 1. Introduction

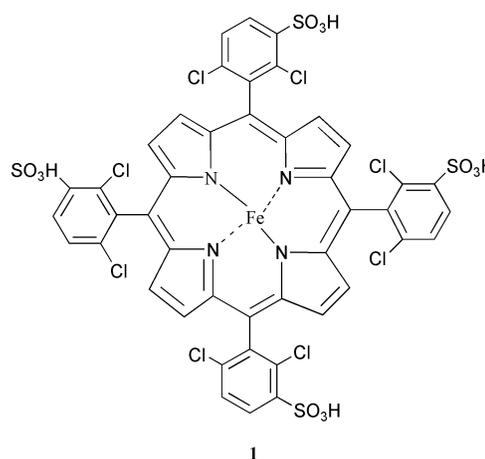
Chlorinated phenolic compounds are frequently found as pollutants in wastewater,<sup>1-3</sup> and originate from sources as diverse as the wood-pulp industry, water purification and production of synthetic pesticides, polymers, detergents, *etc.* The impact caused on environment by these compounds has led to an increasing interest not only in developing new methods for promoting the decomposition of such materials in aqueous medium but also in the identification of their photoproducts. The degradation of phenolic compounds has been extensively studied,<sup>4-10</sup> although in several systems the catalysts and the main products of oxidation are more toxic than the starting materials.

Increased emphasis is being placed on advanced oxidation processes (AOP)<sup>11</sup> for water treatment, and for environmental remediation. Photochemical oxidation, particularly using molecular oxygen, is undoubtedly one of the most important of these methodologies, since it does not liberate any additional pollutant. Singlet oxygen is a particularly good candidate for these applications since it is a very reactive species, as can be seen from its applications in areas as diverse as photodynamic therapy for treatment of cancer,<sup>12</sup> involvement in processes of oxidative stress,<sup>13</sup> photodegradation of dyes, polymers, alkenes and phenolic compounds.<sup>14</sup> The development of new synthetic substances capable of efficient production of singlet oxygen, [<sup>1</sup>O<sub>2</sub>(<sup>1</sup>Δ<sub>g</sub>)], is currently of great interest. Porphyrins, metalloporphyrins<sup>14</sup> and phthalocyanines<sup>15,16</sup> have been shown to be highly effective triplet state photosensitisers capable of producing singlet oxygen with high quantum yields.<sup>17,18</sup> Further, it is well documented that the presence of chlorine atoms in *meso* phenyl groups or the presence of central metal atoms in the porphyrin structure increases the quantum yields of S<sub>1</sub> → T<sub>1</sub> intersystem crossing (isc), and hence triplet state formation of the sensitiser.<sup>19,20</sup> For example, it has been reported that the triplet quantum yield, Φ<sub>T</sub>, for *meso*-tetrakis(2,6-dichloro-

phenyl)porphyrin has a value of 0.995, while the corresponding singlet oxygen quantum yield, Φ<sub>Δ</sub>, is 0.98. This porphyrin has proved to be an efficient sensitiser for the photooxidation of 1,5-dihydroxynaphthalene to 5-hydroxynaphthoquinone.<sup>21</sup>

However, this porphyrin is only soluble in organic solvents, and for environmental applications it is desirable to have water-soluble sensitisers. The use of water-soluble sensitisers such as sulfonated zinc, aluminium, gallium and iron complexes of phthalocyanines<sup>14</sup> as photosensitisers (P) has been reported for the photodegradation of 4-chlorophenol<sup>14,22</sup> and atrazine.<sup>23</sup> In these photosensitised oxidation reactions, two possible mechanisms may be involved: Type I mechanism involving electron or hydrogen transfer<sup>24</sup> or Type II mechanism involving formation of singlet oxygen, <sup>1</sup>O<sub>2</sub>, by energy transfer from the triplet state of the sensitiser, <sup>3</sup>P\* to molecular oxygen, <sup>3</sup>O<sub>2</sub>, and subsequent reaction of singlet oxygen with the substrate.

In this work we report the synthesis of water-soluble iron *meso*-tetrakis(2,6-dichloro-3-sulfophenyl)porphyrin (FeTDCPPS), **1**, and its efficient application as photosensitiser in the photooxidation of 4-chlorophenol, taken as a model compound. Kinetic and mechanistic details will be discussed.



† Dedicated to Professor Sebastião Formosinho on the occasion of his 60th birthday.

## 2. Experimental

### Chemicals

4-Chlorophenol (4-CP) (Aldrich) **2**, *p*-benzoquinone (BQ) **3** (Aldrich Chem, p.a.), *p*-hydroquinone (HQ) **4** (Merck, p.a.), sodium azide (NaN<sub>3</sub>) (Aldrich) and methanol (HPLC analysis grade, Merck) were used as received without further purification. All solvents used in the synthesis of the photosensitiser were purified before use. Solutions were prepared with doubly distilled water, either equilibrated with air or bubbled with argon for 30 min at room temperature.

### Synthesis of the photosensitiser

The photosensitiser iron *meso*-tetrakis(2,6-dichloro-3-sulphophenyl)porphyrin, **1**, was synthesised according to the method described by Gonsalves and Pereira.<sup>25,26</sup> In this, the dichlorobenzaldehyde (10 mmol) was dissolved in a mixture of acetic acid (75 ml) and nitrobenzene (50 ml) and the temperature was raised to 120 °C. Pyrrole (0.7 ml) was then added. The temperature was kept constant for 1 h. The solution was then allowed to cool to room temperature to give crystals of the product porphyrin, which were filtered off and dried, resulting in 5% of *meso*-tetrakis(2,6-dichlorophenyl)porphyrin (TDCPP). <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ: 8.62 (s, 8H), 7.68 (m, 12H), -2.59 (s, 2H); MS (FAB), *m/z* 887–905 ([M + H]<sup>+</sup>, Cl isotopes). Further details of the characterisation has been reported elsewhere.<sup>25</sup>

TDCPP (100 mg) was then stirred with neat chlorosulfonic acid (6 ml) at 100 °C for 3 h. After separation by precipitation with ice, *meso*-tetrakis(2,6-dichloro-3-chlorosulfonylphenyl)porphyrin was obtained in 85% yield. <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ: 8.65 (s, 8H), 8.60 (d, 4H, *J* = 8.6 Hz), 8.05 (d, 4H, *J* = 8.6 Hz), -2.4 (s, 2H); MS (FAB), *m/z* 1279 ([M + H]<sup>+</sup>). Further details of the characterisation has been reported elsewhere.<sup>26</sup>

Metallation of this porphyrin was achieved by refluxing it with iron(II) sulfate in an inert atmosphere in *N,N*-dimethylformamide.<sup>27</sup> The reaction was monitored by UV-visible absorption spectroscopy and was stopped when the characteristic four Q bands of the free porphyrin disappeared and were replaced by the metalloporphyrin spectrum.

The final step consisted of the hydrolysis of the chlorosulfonated porphyrin (80 mg) with water (120 ml) in the presence of oxygen for 20 h to give the corresponding iron *meso*-tetrakis(2,6-dichloro-3-sulphophenyl)porphyrin, **1**. Although we were not able to specifically determine the oxidation state of iron, indications that it is iron(III) come from changes in the UV-visible absorption spectrum when hydroxide or cyanide ions are added, and also the fact that in flash photolysis studies its triplet state is able to oxidise 4-chlorophenol. Support comes from the literature on reactions between iron(II) salts and free-base porphyrins in aerated solutions leading to the corresponding iron(III) porphyrins.<sup>28</sup>

### Continuous photolysis

Continuous irradiation of an aerated solution of FeTDCPPS **1** and 4-CP **2** was carried out using monochromatic light from a xenon lamp (1600 W) filtered with a grating monochromator set at 340 nm. Where necessary, solutions were deaerated by bubbling with argon for 30 min before irradiation. The incident light intensity of the xenon lamp was determined by chemical actinometry using a solution of potassium ferrioxalate<sup>29</sup> (*I*<sub>0</sub> = 1.97 × 10<sup>15</sup> photons cm<sup>-2</sup> s<sup>-1</sup>). In a typical experiment an aerated aqueous solution of **2** (6.1 × 10<sup>-4</sup> M) and **1** (3.2 × 10<sup>-4</sup> M) was added to a quartz cell (1 cm pathlength), which was then irradiated. The reaction was monitored by UV-visible spectroscopy using a Shimadzu UV-2010 spectrophotometer and also by HPLC analysis. The pH of solutions was carefully controlled during the irradiation time using a Crison micro-pH 2000 apparatus. In addition, for mechanistic studies, an aerated

solution of photosensitiser **1** (3.2 × 10<sup>-4</sup> M) and 4-CP **2** (6.1 × 10<sup>-4</sup> M) was irradiated at 340 nm in the presence of sodium azide (1 × 10<sup>-3</sup> M).

### HPLC analysis

Analysis of degradation products of 4-CP was made using a HPLC Waters 540 liquid chromatography system equipped with a UV-visible photodiode array detector (Waters 990) with detection set at 280 nm. The formation of the photoproducts BQ **3** and HQ **4** was monitored against external standards. A reverse phase Waters Spherisorb 5 μm ODS 4.6 × 250 mm column (Analytical Cartridge) was used with a mixture of MeOH–H<sub>2</sub>O (60 : 40 v/v) as eluent in isocratic mode with a flow rate of 1 ml min<sup>-1</sup>.

### Time-resolved flash photolysis studies

The transient absorption spectra of the triplet state of the photosensitiser **1**, the kinetics of its decay, and quenching with oxygen were measured on an Applied Photophysics LKS 60 nanosecond laser flash photolysis kinetic spectrometer using the third harmonic (355 nm) of a Quanta Ray GCR 130–01 Nd/YAG laser for excitation. The transient absorbance spectra at preselected wavelengths were monitored by a detection system consisting of a pulsed xenon lamp (150 W), monochromator and photomultiplier. The signal from the photomultiplier was digitalised by an oscilloscope. Full details are given elsewhere.<sup>38</sup> The concentration of the photosensitiser **1** was calculated to have maximum absorbance around 0.2–0.4 at the laser wavelength. For studying the effect of oxygen as quencher of the excited state of the photosensitiser **1**, aqueous solutions of FeTDCPPS were studied with air ([O<sub>2</sub>] = 2.8 × 10<sup>-4</sup> M),<sup>30</sup> saturated with oxygen ([O<sub>2</sub>] = 1.4 × 10<sup>-3</sup> M) and deaerated with argon for 30 min. The effect of 4-CP as a quencher of the excited state of the photosensitiser **1** was also studied in deaerated solutions. The singlet oxygen quantum yield was measured by Dr S. Navaratnam (North East Wales Institute, UK) using a previously described time-resolved phosphorescence system, which uses a Nd/YAG laser for excitation and photodiode for detection.<sup>31</sup> The standard used was an aqueous solution of *meso*-tetrakis(4-sulphophenyl)porphyrin, TPPS (*φ* = 0.62).

## 3. Results and discussion

### Synthesis of the photosensitiser

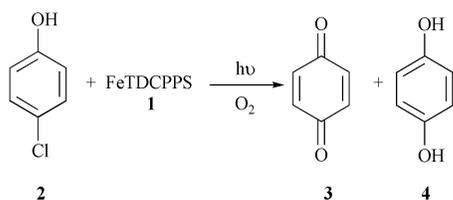
Although water-soluble sulfonated *meso*-aryl porphyrins have frequently been synthesised by direct reaction of strong sulfuric acid and the desired precursor porphyrin,<sup>32</sup> the use of chlorosulfonic acid in the present study has the advantage that it is more efficient, and the chlorosulfonyl porphyrin formed is very easy to isolate from the reaction medium, simply by addition of ice. The precipitate formed was extracted with dichloromethane and purified by silica gel column chromatography using dichloromethane as eluent. The iron complex of the *meso*-tetrakis(2,6-dichloro-3-sulphophenyl)porphyrin was prepared using the *N,N*-dimethylformamide method,<sup>27</sup> where the porphyrin was dissolved in DMF and iron(II) sulfate was added (1 : 1). The formation of the complex was monitored by measuring the absorption spectrum and the bands at 429, 516, 555, 581 and 636 nm change to 417, 518 and 586 nm. The final step involved hydrolysis of the sulfonyl chloride by refluxing the iron(III) complex with water for 20 h, giving after solvent evaporation, the photosensitiser **1** in 80% yield. A typical metalloporphyrin visible spectrum was observed: solvent (H<sub>2</sub>O) λ/nm (ε/M<sup>-1</sup> cm<sup>-1</sup>): 417 (3.3 × 10<sup>5</sup>), 518 (8.0 × 10<sup>3</sup>), 586 (4 × 10<sup>3</sup>).

### Continuous irradiation

On irradiation of an aerated solutions of 4-CP **2** (6.1 × 10<sup>-4</sup> M) in the presence of the photosensitiser **1** (3.2 × 10<sup>-4</sup> M) at

340 nm significant changes in the UV-visible spectra were observed. New bands were observed in the 240–300 nm region. However, 4-CP and the photoproducts absorb in the same wavelength range, such that it is not possible to follow the extent of transformation simply from changes in the UV spectra. For quantitative measurements of photoconversion and product formation HPLC analysis was used.

When irradiation of 4-CP and the photosensitiser **1** was performed in the presence of oxygen the main photooxidation photoproduct was *p*-benzoquinone **3** ( $\lambda_{\text{max}} = 245 \text{ nm}$ ) (33%) although some *p*-hydroquinone **4** ( $\lambda_{\text{max}} = 288 \text{ nm}$ ) (6%) was also observed (Scheme 1). This is similar to what is reported for the degradation of 4-CP and related phenols in the presence of phthalocyanines and other photosensitisers.<sup>4,33–39</sup>



Scheme 1

The formation of the dimeric photoproducts 2,4'-dihydroxy-5-chlorobiphenyl ( $\lambda_{\text{max}} = 257$  and  $295 \text{ nm}$ ) and 2,2'-dihydroxy-5,5'-dichlorobiphenyl ( $\lambda_{\text{max}} = 280 \text{ nm}$ ) was also observed, in agreement with previous literature results for photodegradation of 4-CP with photooxidants.<sup>37,38</sup> However, the dimeric species suffer further oxidation. Although their oxidation products were not characterised, increase light scattering in the UV-visible absorption spectra, at long irradiation times, confirms the formation of insoluble compounds.

Upon photolysis of deaerated solutions under the same conditions, the main photoproduct obtained was *p*-hydroquinone **4**, in agreement with previous reports on photolysis of 4-CP using different sensitiser.<sup>15,39</sup>

Upon direct photolysis at 340 nm of aqueous solutions of 4-CP ( $3.2 \times 10^{-4} \text{ M}$ ), in the presence of oxygen and without sensitiser, no photodegradation was observed after 3 h of irradiation.

We believe that the formation of *p*-benzoquinone, **3**, is mainly due to the reaction of singlet oxygen with 4-CP while the formation of hydroquinone, **4**, and dimeric photoproducts can be explained by electron transfer from 4-CP to the triplet state of the sensitiser with concomitant formation of the 4-chlorophenoxy radical followed by HO addition or dimerisation, respectively.

The kinetic profiles were studied of photodegradation of 4-CP with photosensitiser **1** and of photoproduct **3** and **4** formation, in aerated solution, following continuous irradiation with monochromatic 340 nm light. The disappearance of 4-CP with irradiation time follows first order kinetics with a rate constant of  $k = 9.5 \times 10^{-5} \text{ s}^{-1}$  and was accompanied by formation of benzoquinone and hydroquinone, Fig. 1.

Some metalloporphyrin photodegradation was observed. This, together with consumption of oxygen in the reaction mixture, may explain the fact that only 65% conversion of the chlorophenol was observed. Other metalloporphyrins, with higher photostability are currently under study.

Upon photolysis of aerated solution of the photosensitiser **1** and 4-CP **2** in the presence of sodium azide ( $1 \times 10^{-3} \text{ M}$ ), no formation of the photoproduct **3** (BQ) was observed. Azide ion is known to be a selective  $^1O_2$  ( $^1\Delta_g$ ) singlet oxygen quencher,<sup>40</sup> strongly suggesting that BQ formation results from reaction with singlet oxygen.

### Laser flash photolysis experiments

The kinetic behaviour of transient species was monitored by nanosecond laser flash photolysis. Following excitation at 355

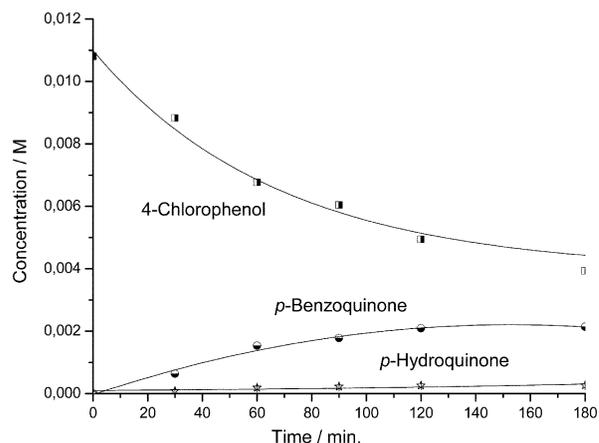


Fig. 1 HPLC analysis of the photosensitised degradation of 4-CP and the formation of the photoproducts *p*-benzoquinone and *p*-hydroquinone as a function of time.

nm of an aqueous solution of **1** ( $6.2 \times 10^{-4} \text{ M}$ ) in the presence of oxygen, a transient absorption was observed between 300 and 600 nm, with a maximum at 460 nm, Fig. 2. This is identified as the sensitiser triplet state (T–T) absorption, in agreement with flash photolysis studies on related systems.<sup>41</sup> No other transient species were found.

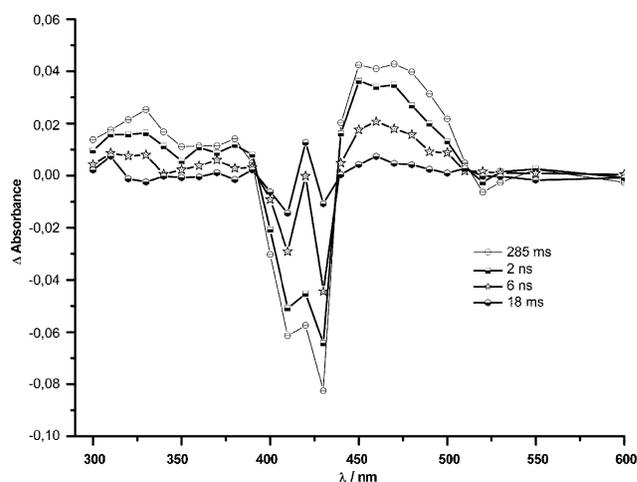
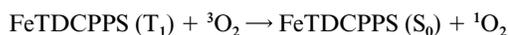


Fig. 2 Transient absorption spectra observed following laser flash photolysis at 355 nm of a nitrogen-saturated solution of sensitiser **1** in water.

Oxygen is a good quencher of triplet excited states. The lifetimes of the triplet state of the photosensitiser **1** were determined at 460 nm in aerated ( $4.70 \mu\text{s}$ ) and nitrogen saturated ( $378 \mu\text{s}$ ) solutions. The bimolecular quenching rate constant ( $k_q$ ) for quenching of the triplet state of FeTDCPPS by molecular oxygen ( $^3O_2$ ) in aqueous solutions is  $7.25 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ , which is comparable to literature values for related porphyrins.<sup>42</sup> This is considerably lower than the diffusion controlled rate in water ( $9.8 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$ ), in agreement with the studies of Porter and co-workers,<sup>43</sup> who suggested that for quenching of triplet states by molecular oxygen *via* an energy transfer mechanism,  $k_q \approx 1/9k_{\text{diffusion}}$ . This supports the idea that quenching of the triplet state ( $T_1$ ) of the photosensitiser **1** by molecular oxygen occurs by energy transfer with formation of the reactive species singlet oxygen ( $^1O_2$ )



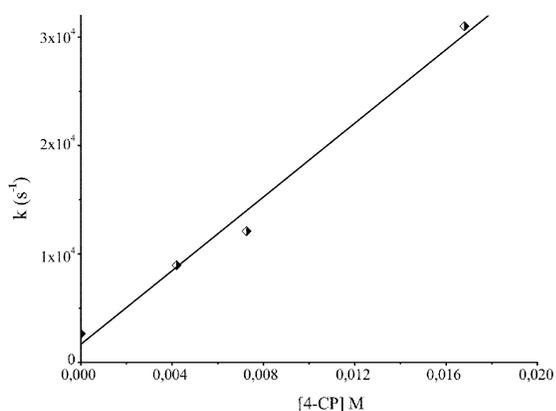
To confirm this, time-resolved measurements were carried out to detect the singlet oxygen phosphorescence. Upon excitation at 355 nm of aerated solutions of sensitiser **1** in water, singlet oxygen phosphorescence was observed at 1270 nm.

Using TPPS as standard, a singlet oxygen yield of  $\Phi_{\Delta} \approx 1$  was observed. More details will be reported elsewhere. Singlet oxygen is known to react with 4-CP<sup>44</sup> with a rate  $6.0 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ , and the above results support the idea that the photosensitised degradation of 4-CP by **1** involves, at least in part, reaction with singlet oxygen.

Formation of quinones has previously been reported in the reaction between singlet oxygen and phenols or naphthols.<sup>21,36</sup> This reaction is suggested to proceed *via* 1,4-cycloaddition followed by breakdown of the intermediate endoperoxide.<sup>36</sup>

However, the reaction in the absence of oxygen suggests another pathway may also be involved. To evaluate the direct reaction between 4-CP and the triplet state of the photosensitiser **1**, flash photolysis experiments were carried out with 355 nm excitation on deaerated aqueous solutions of FeTDCPPS with different concentrations of 4-CP, with sensitizer absorption monitored at 460 nm. Decay of the triplet absorption was more rapid in the presence of the phenol, and from the study of the pseudo-first-order rate constant ( $k_{\text{obs}}$ ) as a function of 4-CP concentration, Fig. 3, the kinetics were found to follow the rate law

$$k_{\text{obs}} = k_0 + k_q[4\text{-CIPh}]$$



**Fig. 3** Kinetic plot of the rate of decay of the triplet state of sensitizer **1** as a function of 4-chlorophenol concentration.

From the slope of this, a bimolecular quenching rate constant, of reaction of the triplet state of the photosensitiser **1** with 4-CP of  $k_q = 1.68 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$  was determined.

This confirms that reaction between the triplet state of **1** and 4-CP occurs in parallel with singlet oxygen formation. However, for the reaction conditions employed in this study, this is likely to be less important than the sensitised photodegradation of 4-CP by singlet oxygen.

## Conclusions

A water soluble metal complex, iron *meso*-tetrakis(2,6-dichloro-3-sulfophenyl)porphyrin, has been synthesised, and used as photosensitiser for the degradation of 4-chlorophenol. Benzoquinone and hydroquinone have been identified as the main products in aerated aqueous solution. Studies using azide as quencher, direct observation of the quenching of the porphyrin triplet state by molecular oxygen, and detection of singlet oxygen phosphorescence suggest that the dominant mechanism of quinone formation involves reaction of singlet oxygen with the chlorophenol. However, the triplet state of the quencher is also quenched by the phenol, and this process may provide a second, but less important, route to 4-chlorophenol degradation

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