Pyrolytic formation and photoactivity of reactive oxygen species in a SiO<sub>2</sub>/carbon nanocomposite from kraft lignin [version 1; referees: 1 approved]

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Abstract
SiO<sub>2</sub> and carbon produced by kraft lignin pyrolyzed at 600°C can generate stable reactive oxygen species (ROS) by reaction with atmospheric oxygen. In this study, we systematically investigate the photochemistry of peroxyl radicals in carbon-supported silica (PCS) and assess its effects on the methylene blue (MB) photodegradation. Characterization revealed that the higher ROS generation ability of SiO<sub>2</sub>/carbon under UV light irradiation was attributed to its abundant photoactive surface-oxygenated functional groups.

Keywords
ROS, photochemistry, methylene blue, degradation, UV

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Introduction

Consistent access to clean water has come into focus this millennium due to high pollution; a reduced amount of drinkable water could be the next challenge for the future due to overpopulation[1-3]. The application of photocatalytic technology using semiconductors to solve the environmental problems, like the degradation of organic effluents has been received much attention[4-6]. Heterogeneous photocatalysis using semiconductors is an interesting method falling into advance oxidation processes (AOPs)[7-11] that can produce highly reactive species containing oxygen (ROS). In fact, with this method is possible to produce oxidizing molecules like hydrogen peroxide and singlet oxygen ($O_2^*$) together with radicals like hydroxyl radical (OH) and superoxide radical anion ($O_2^-$)[12-13]. These reactants can decompose organic pollutants in wastewater giving harmless compounds[14].

Recently, N. Chen et al. reported that reactive oxygen species generation in hydrochar and photochemistry of Sulfadimidine degradation in water[15]. Y. Chen et al. reported the photo degradation of tetracycline in aqueous solution under simulated sunlight irradiation through the singlet oxygen[16]. Li et al. reported that the degradation of ibuprofen by UV-visible light irradiation included direct photolysis and self-sensitization via ROS[17]. Wang et al. reported that when a simpler molecule without visible-light absorption is degraded, the Fe-hydroxyl complexes still produce highly reactive species that can react with MB, giving rise to photobleaching (Equation 3).

\[ \text{Dye} + \text{photon} = \text{Dye}^* \]  
\[ \text{Dye}^* + \text{O}_2^+ = \text{Dye} + \text{O}_2 \]  
\[ \text{O}_2^+ + \text{Dye} = \text{oxidation products} \]

In literature are present many methods for photoassisted AOPs like photo-electrochemical cells composed by an anode made with boron-doped carbon and cathode in carbon nanotubes; with this system, a model azo dye was depleted[18]. Also exfoliated graphene, decorated with titanium dioxide and nanoparticles, is effective for photo-catalytic water treatment[19-21].

In our current scenario, stable peroxy radicals in carbon-supported silica (PCS) are prepared from cheap starting materials. The method used is the pyrolysis under vacuum of kraft lignin deposited onto silica. Vacuum pyrolysis produced defective carbon bearing carbon radicals. These radicals are quickly transformed into peroxy radicals by reaction with oxygen molecules present in the atmosphere.

Methods

The materials and methods to produce PCS using high-vacuum pyrolysis are clearly explained and characterized previously[22]. In brief, kraft lignin was absorbed onto silica and pyrolyzed under vacuum at 600 °C. For the kinetic data analysis, linear quadratic fitting and other kinetic fitting (reaction order checking) were performed by using Origin v6.0.

Degradation of MB dye procedures and analyses

100-ml of air-equilibrated 10^{-6} M solutions of MB (Sigma Aldrich, India) in water containing 100 mg (1 mg/ml) of neat SiO$_2$ or PCS were poured in quartz cylindrical reactors (90 mm diameter x 25 mm height). Solutions were magnetically stirred in the dark for 10 min before irradiation and kept under stirring during the experiment. The light source consisted of two 15-W phosphor-coated lamps (center of emission, 366 nm). Aliquots (4 ml) were withdrawn at 5-min intervals (for a total of 10-12 samples) during the irradiation until the disappearance of the color. Solids were removed by syringe filtration with a 0.4-μm pore size, and the filtrates immediately examined by UV-visible absorption spectroscopy in 1-cm quartz cuvettes using a JASCO V-630 UV-visible spectrophotometer. The absorbance was normalized by dividing the absorbance at 668 nm of the sample (A) with the absorbance of the initial solution ($A_0$).

Results and discussion

Degradation of MB

To assess the respective photocatalytic activity of PCS and of neat SiO$_2$, we carried out competitive experiments with MB (Figure 1). PCS did not react with MB, in fact, solutions left for 24 hours in the dark does not show a decrease of MB concentration. Nonetheless, under dark conditions the dye was absorbed by PCS to a nearly tenfold greater extent than with pristine SiO$_2$ (dark region between −10 and 0 min, Figure 1b).

Normally photocatalysts produce radicals able to degrade organics but in the case of PCS the catalyst already possesses reactive radicals.

Simple mechanism of established photocatalysts in MB

The net effect of PCS on the photodegradation of MB is a threefold increase in the kinetics of photodegradation (Table 1). Without the assistance of an active photocatalyst, the only reaction mechanism that is applicable is the generation of singlet oxygen by sensitization (Equation 2) via the excited state of the dye. The singlet oxygen can react with MB, giving rise to photobleaching (Equation 3).

\[ \text{Dye} + \text{photon} = \text{Dye}^* \]  
\[ \text{Dye}^* + \text{O}_2^+ = \text{Dye} + \text{O}_2 \]  
\[ \text{O}_2^+ + \text{Dye} = \text{oxidation products} \]

With PCS, MB is strongly absorbed onto the pyrolytic carbon present on the catalyst surface. Moreover, pyrolytic carbon possesses a high concentration of peroxy radicals. The enhancement on the reaction kinetic could be due to a local increase of concentration of dye and active oxygen. Since the oxygen is reversibly absorbed on the carbon giving peroxy radicals[22], the surface of the catalyst is never depleted due to the presence of oxygen in solution.

In fact, in these conditions, we can have, together with Equation 1–Equation 3, a possible reaction of the excited state of the reactant with peroxy radicals or adsorbed oxygen on PCS (Equation 4).

\[ \text{Dye}^* + \text{PCS-OO} = \text{PCS} + \text{dye oxidation} \]  
\[ \text{PCS} + \text{O}_2^+ = \text{PCS-OO} \]

The peroxy radicals are reversibly formed by capture of atmospheric oxygen due to the presence of highly active pyrolytic carbon on PCS:
Figure 1. Normalized spectral intensity of the 668 nm band of methylene blue (MB) during (a) the UV-irradiation of the MB/SiO$_2$ suspension at 366 nm at different time intervals, and (b) the same process for the MB/peroxyl radicals in carbon-supported silica (PCS) suspensions under otherwise identical conditions. The region between −10 and 0 min refers to the extent of adsorption of the MB dye under dark conditions. It shows the first-order kinetics of the photodegradation of the MB dye by MB/PCS. 3 repeats performed.

Table 1. Extent of adsorption and first-order kinetics of photodegradation of methylene blue (MB) (1.0 μM) on pristine SiO$_2$ and on SiO$_2$/graphene in aqueous media under ambient atmospheric conditions and under UV irradiation at 366 nm.

<table>
<thead>
<tr>
<th>Dye</th>
<th>k (min$^{-1}$)</th>
<th>Adsorption, %</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>SiO$_2$</td>
<td>PCS</td>
</tr>
<tr>
<td>MB</td>
<td>0.027 ± 0.005</td>
<td>0.092 ± 0.006</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>91</td>
</tr>
</tbody>
</table>

Another possibility is the transfer of energy (or sensitization) of the excited state of the absorbed dye directly to the defective pyrolytic carbon, giving rise to formation of ROS. All these mechanism lead to an enhancement on the degradation of MB.

600°C did not affect the crystalline state of silica when it was coated with carbon. The photocatalytic activity was measured against pristine SiO$_2$ through an examination of the kinetics of degradation of MB by UV-vis spectroscopy. Under UV light irradiation, the degradation was threefold greater for the MB-PCS compared with MB-silica.

Data availability
Dataset 1: Raw data for the article ‘Pyrolytic formation and photoactivity of reactive oxygen species in a SiO$_2$/carbon nanocomposite from kraft lignin’ are presented, [10.5256/f1000research.16080.d21890](https://doi.org/10.5256/f1000research.16080.d21890)

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This manuscript is focused on the application of supported stable peroxyl radicals for the photo-degradation of organic materials. In this work the authors systematically investigated the photochemistry of peroxyl radicals in carbon-supported silica (PCS) and then they evaluated the effects of PCS on the methylene blue photodegradation as a model for a generic organic effluent. The manuscript is clearly written with few errors (e.g. “N. Chen et al.” and “Y. Chen et al.” refer to the same reference article). However, the authors have extracted some interesting data that well supports the discussion and the appropriate conclusions. Furthermore, problems such as overpopulation and the lack of drinking water are unfortunately a plague that afflicts our entire planet. I encourage the authors to continue with their research, thus contributing to increase the impact of their study.

Is the work clearly and accurately presented and does it cite the current literature?
Yes

Is the study design appropriate and is the work technically sound?
Yes

Are sufficient details of methods and analysis provided to allow replication by others?
Yes

If applicable, is the statistical analysis and its interpretation appropriate?
Yes

Are all the source data underlying the results available to ensure full reproducibility?
Yes

Are the conclusions drawn adequately supported by the results?
Yes

Competing Interests: No competing interests were disclosed.

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