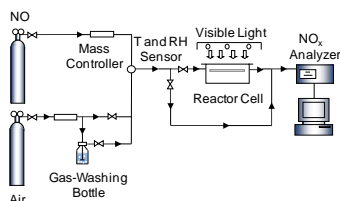


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Different commercial, visible light response, TiO₂ powders are studied in order to assess their application in indoor air decontamination. To do that the TiO₂ powders were immobilized on borosilicate glass plates according to a dip coating method. Then, the photocatalytic performance of these plates under visible light was evaluated in a continuous gas flat plate photoreactor using nitrogen oxide gas (NO) as the pollutant. The photocatalytic NO_x degradation by immobilized TiO₂ powders was related with their optical properties, finding a clear correlation between them.

These results are useful to decide which TiO₂ will be more efficient in a full scale air decontamination process under indoor conditions.

Indoor air quality has received much attention because of the very important role indoor environment plays on human health. Nitrogen oxides (NO and NO₂), sulfur dioxide (SO₂) and volatile organic compounds (VOCs), as typical inorganic and organic indoor air pollutants, can be emitted from cooking, combustion, exhaust gases, tobacco smoke, furniture, building materials, and even traffic pollutants from outside of the building and can cause serious health problems like drowsiness, headache, sore throat, and mental fatigue. The US Environmental Protection Agency pointed out that indoor air pollution poses greater risk than outdoor air pollution due to the fact that people spend more time in indoor environments. So, it is of vital importance to remove these pollutants in order to improve the indoor air quality for people's health. In addition, some studies have shown that when certain contaminants were removed from a working places, the performance of some office tasks improved.

Heterogeneous photocatalysis represents an emerging environmental control option for the efficient removal of chemical pollutants and it can be applied to water and air purification. However, normal TiO₂ can only be activated by UV radiation (280-400 nm) that represents 4% of the total energy of the sun; meanwhile the visible light constitutes 45% of the solar radiation. On the other hand, the UV radiation amounts to only 0.001-0.05 W/m² in indoor lighting [1]. To extend the use of photocatalysis to visible light region, it is necessary to prolong TiO₂ radiation absorption to wavelengths corresponding to visible spectrum (400-700 nm). So far, several modification

methods of photocatalyst to amplify its absorption spectrum to visible radiation have been investigated and therefore, the applicability of heterogeneous photocatalysis has been extended [2,3]. Even several commercial types of modified TiO₂ can be already found in the market for indoor visible light applications.

In the present work, different visible light TiO₂ commercially available, one doped with carbon and the other one doped with nitrogen, are studied. The main objective of this work is to determine which photocatalyst will present the best pollution abating ability in a full scale application under real indoor conditions.

The immobilization of the TiO₂ powders was carried out over borosilicate glass plates by a dip coating procedure employing a coating suspension of 75 g L⁻¹ of TiO₂ powder in deionised water. Every coating cycle, the glass pieces were dried at 110°C for 24 hours and calcined at 500°C for 2 hours. This procedure was performed four times for each TiO₂ powder.

The experimental setup to assess the photocatalytic performance is described in the Graphical Illustration. This consists of a continuous flat photoreactor where the glass pieces with the immobilized TiO₂ are placed inside of it. The photoreactor is fed by NO gas stabilized in air and irradiated with fluorescence visible light lamps. The outlet contaminant concentration from the reactor is analyzed with a chemiluminescence NO_x analyzer. Table 1 shows the main characteristics, dimensions and operating conditions of the employed test setup to execute the NO_x degradation experiments.

The optical properties of the immobilized TiO₂

in the glass plates were determined as a function of wavelength between 300 and 600 nm in an Optronic OL Series 750 spectroradiometer equipped with an OL 740-70 integrating sphere reflectance attachment. To evaluate the fraction of energy absorbed (A) by the coated glass with TiO₂, the spectral diffuse reflectance (R) and the spectral diffuse transmittance (T) were experimentally determined. The mathematical expression between the absorbed fraction, transmittance and reflectance is given by [4]:

$$1 = A + T + R$$

Table 1. Experimental setup characteristics and operating conditions.

Reactor	Length	2 dm
	Width	1 dm
	Height	0.03 dm
	Volume	0.06 dm ³
Photocat. Samples	Length	2 dm
	Width	1 dm
Visible Lamps: Philips Master TLD 865 x 3	Input Power	18 W
	Emission Wavelength	380-720 nm
Flow Rate		3 L min ⁻¹
Relative Humidity		50%
Irradiance Flux		10 W m ⁻²
NO Inlet Concentration		1 ppm
NO ₂ Inlet Concentration		0 ppm

Figure 1 shows the conversion results of total NO_x by the carbon and nitrogen doped TiO₂. Clearly, the carbon doped TiO₂ has shown a better photocatalytic performance for the NO_x degradation than the nitrogen doped TiO₂ powder. Meanwhile the total NO_x conversion by the TiO₂ - C is about 18%, the TiO₂ - N presents only 6% of NO_x degradation.

Figure 2 shows the spectral fraction of energy absorbed by both immobilized TiO₂ powders calculated from the experimentally determined diffuse reflectance and transmittance of the coated glass plates. In addition, the spectral emission

distribution (E_{λ}/E_{total}) of the employed lamps in the reacting experimental setup is shown in this figure. Despite that these lamps do not emit radiation in the UV region, where the TiO₂ presents a significant light absorption, the energy emitted in the visible light range is enough to activate this kind of modified photocatalysts.

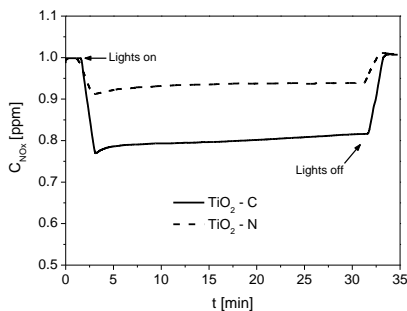


Figure 1. NO_x outlet concentration evolution for the carbon and nitrogen doped TiO₂.

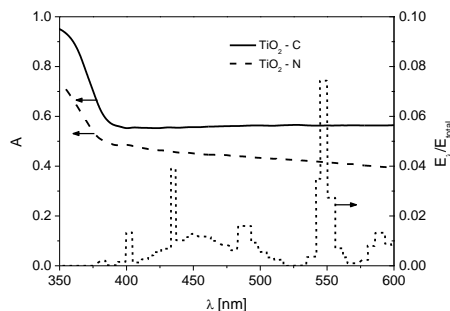


Figure 2. Spectral fraction of energy absorbed by the immobilized carbon and nitrogen doped TiO₂; and the spectral emission distribution of the visible light lamps employed in the photoreactor.

A correlation between the obtained results for the photocatalytic performance and the fraction of energy absorbed by the immobilized TiO₂ powders can be observed. At the same time that the TiO₂ - C absorbs about 20% more energy than TiO₂ - N, it can convert 66% more NO_x. However, it is worth to mention that other photocatalyst characteristics can influence in its performance, for instance the specific surface.

Indeed, the carbon modified photocatalyst promises a better performance for a real application of indoor purification.

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