Visible-Light Active Sulfur-Doped Titania Nanoparticles Immobilized on a Silica Matrix: Synthesis, Characterization and Photocatalytic Degradation of Pollutants

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Abstract: The photocatalytic oxidation (PCO) of pollutants using TiO$_2$-based materials can significantly improve indoor air quality (IAQ), which in turn, has a significant impact on human health and life expectancy. TiO$_2$-based nanoparticles (NPs) are widely used as part of building materials to function as photocatalysts in PCO. In this work, a series of sulfur-doped TiO$_2$ NPs immobilized on a silica matrix were synthesized by combining a sol-gel process with ball milling. The samples were structurally characterized by X-ray diffraction (XRD), UV-Vis diffuse reflectance spectroscopy (DRS), Fourier-transform infrared spectroscopy (FT-IR) and N$_2$ adsorption-desorption isotherms. Furthermore, the morphological characteristics were determined by dynamic light scattering (DLS), scanning electron microscopy (SEM) and transmission electron microscopy (TEM). The photocatalytic activity of the as-prepared S-doped TiO$_2$/SiO$_2$ NPs in the degradation of liquid and air pollutants under visible-light irradiation was investigated. Our results show that sulfur is an effective dopant for activating TiO$_2$/SiO$_2$ photocatalysts under visible-light irradiation. Silica constitutes a “safe-by-design” approach and inhibits the aggregation of NPs during synthesis. The most efficient photocatalyst afforded 79% removal of methyl orange (5 h), 26% removal of acetaldehyde (1 h) and 12% oxidation of NO (1 h).

Keywords: photocatalysis; S-doping; TiO$_2$ nanoparticles; SiO$_2$; MO degradation; NO$_x$ oxidation; safe by design

1. Introduction

The levels of air pollutants such as nitrogen oxides (NO$_x$), volatile organic compounds (VOCs) and carbon monoxide (CO) in indoor environments can be significantly higher compared with the levels outdoors due to the contribution of indoor sources such as building materials, office equipment, consumer products and combustion by-products [1]. The photocatalytic degradation of these pollutants by nanoparticles (NPs), integrated into building materials and capable of photocatalytically oxidizing these toxic compounds to non-hazardous products, will greatly improve indoor air quality (IAQ). To this end, metal oxide semiconductors such as TiO$_2$ [2–5] and ZnO [6,7] are still considered as the most efficient photocatalysts by virtue of their high chemical stability, low toxicity and low cost.

TiO$_2$ exists in three different crystalline phases, i.e., anatase, rutile and brookite. In general, anatase shows better photocatalytic activity; however, anatase and brookite are metastable phases that transform to rutile at higher temperatures (500–600 °C). The well-documented photocatalytic oxidation (PCO) mechanism of TiO$_2$ is illustrated in Scheme 1. Light absorption by TiO$_2$ can generate electron-hole pairs. The photogenerated electrons can reduce molecular oxygen to superoxide radical anions ($\bullet$O$_2^-$) or hydrogen peroxide (H$_2$O$_2$), whereas the corresponding holes can oxidize oxygen-containing surface species...
such as adsorbed water molecules or Ti-bound hydroxyl groups to hydroxyl radicals (•OH). These highly reactive oxygen species (ROS) can readily decompose common indoor pollutants such as NOx and acetaldehyde [8].

![Scheme 1. Mechanism for the photocatalytic oxidation of pollutants over TiO2-based catalysts.](image)

However, TiO2 is a semiconductor with a relatively large band gap between 3.0 (pure rutile phase) and 3.2 eV (pure anatase phase). As a result, pure TiO2 can only absorb the UV fraction of sunlight which accounts for less than 5% of the solar spectrum. This poses significant limitations, particularly in indoor applications where the intensity of UV radiation is lower. Moreover, the relatively faster charge recombination of electron-hole pairs in pure TiO2 photocatalysts, before any type of interaction with the adsorbed species can take place, may further decrease the photocatalytic activity [9, 10].

Current challenges in the field aim at the development of novel TiO2-based photocatalysts that can operate under visible light radiation and exhibit extended charge separation. Narrowing of the band gap and prolonged charge separation can be achieved via doping of TiO2 with metallic and non-metallic species, co-doping with more than one element and coupling of TiO2 with other semiconductors [11–14].

The non-metal doping of TiO2 is considered as one of the most efficient methods to increase TiO2 photocatalytic activity under visible light. Doping of TiO2 with non-metals such as carbon, nitrogen and sulfur can shift the absorption band of TiO2 toward the visible region, due to either the generation of new occupied energy levels above the valence band or the formation of oxygen vacancies [15–19]. Nitrogen is the most widely used non-metal dopant due to its small ionization energy and comparable size with oxygen. Nitrogen enters the TiO2 lattice as an anion, either occupying an interstitial position or replacing an oxide anion. By contrast, sulfur can be incorporated either as an anion or a cation. Anionic sulfur doping involves the incorporation of sulfide (S^{2−}) anions into the lattice, as originally shown by Umebayashi et al. [20]. Cationic doping proceeds via the replacement of Ti^{4+} cations in the TiO2 lattice by S^{4+} or S^{6+} cations. The larger radius of sulfide compared with oxide anions makes anionic doping less thermodynamically favored than cationic doping [21–30].

Ohno et al. were the first to report the cationic S-doping of TiO2 using thiourea as the sulfur source [31]. Importantly, S-doped TiO2 exhibited stronger visible light absorption compared to C- and N-doped TiO2. Lie et al. investigated the effect of the thiourea-to-titanium nominal ratio on the photocatalytic phenyl degradation over S-doped TiO2 NPs [22]. A higher amount of S-doping hindered the anatase-to-rutile transformation at higher temperatures and enhanced visible light absorption. However, an upper limit of S-doping was identified above which the photocatalytic activity diminished. The authors concluded that higher levels of S-doping elevated the newly generated energy states to the point where they acted as charge recombination centers. Likewise, an optimal sulfur-to-titanium nominal ratio was observed in the photocatalytic phenol degradation over S-doped TiO2 NPs bearing sulfate surface groups [24].

The employment of a supporting matrix with high mechanical and thermal strength, high surface area and available anchoring sites for TiO2 NPs could promote photocatalytic...
activity. To this end, pristine or doped TiO₂ NPs have been immobilized on metal oxides, carbon nanotubes, reduced graphene oxide, glassy substrates, polymeric surfaces and silica [32–34]. Previous works have shown that particularly the employment of silica (SiO₂) as the support increased the specific surface area of the catalyst, restricted the agglomeration of TiO₂ NPs and suppressed the anatase-to-rutile phase transformation during synthesis [35–40]. The use of silica is also regarded as a “safe-by-design” approach to prevent the release of TiO₂ NPs into the environment due to strong Si–O–Ti interactions [41–47].

Nevertheless, to the best of our knowledge, there is only one literature example in which S-doped TiO₂ NPs were immobilized on silica. Specifically, Chen et al. reported in 2019 the synthesis of SiO₂-supported cationic S-doped TiO₂ NPs via the co-hydrolysis of Si- and Ti-based precursors in the presence of thiourea as the sulfur source. The synthesized catalyst was active in the photocatalytic degradation of phenol under visible light irradiation.

In this work, we present a novel and safe-by-design process that combines sol-gel synthesis and ball-milling to produce a series of S-doped TiO₂/SiO₂ photocatalysts. The effect of sulfur doping on the photocatalytic properties was probed by varying the nominal sulfur-to-titanium ratio. Generally, the sol-gel synthesis enables control of the reaction parameters and affords nanosized crystalline powders (nanopowders) of high purity and stability [48], whereas ball-milling before calcination leads to smaller NPs with a more uniform particle size distribution [49].

The synthesized photocatalysts were characterized via powder X-ray diffraction (PXRD), scanning electron microscopy (SEM), energy dispersive X-ray spectroscopy (EDS), transmission electron microscopy (TEM), selected area electron diffraction (SAED), dynamic light scattering (DLS), Brunauer-Emmett-Teller (BET) surface area analysis, diffuse reflectance (DRS) and Fourier-transform infrared (FTIR) spectroscopy. The photocatalytic activity in the degradation of methyl orange, acetaldehyde and NOₓ was investigated. Our results show that S-doped TiO₂ NPs entrained in a SiO₂ matrix exhibited substantial activity in the photocatalytic degradation of liquid and air pollutants under visible-light irradiation.

2. Experimental Section

2.1. Reagents

The following analytical-grade reagents were used as received: titanium(IV) tetraisopropoxide (TTIP, 97%, Sigma-Aldrich, St. Louis, MO, USA) as a precursor of TiO₂, tetraethyl orthosilicate (TEOS, >99%, Sigma-Aldrich, St. Louis, MO, USA) as a precursor of SiO₂, thiourea (CH₄N₂S, >99%, Penta, Prague, Czech Republic) as the sulfur source and methyl orange (MO, >85%, AppliChem GmbH, Darmstadt, Germany). Absolute ethanol (EtOH, 97%) and deionized water were used as the solvents.

2.2. Materials Synthesis and Characterization

In a typical sol-gel synthesis, TEOS (SiO₂ precursor, 0.30 mL, 1 mmol) was added to an ethanol-water mixture (8 mL EtOH–37 mL H₂O). The mixture was vigorously stirred for 1 h at room temperature and then TTIP (TiO₂ precursor, 7.1 mL, 0.024 mol) was added dropwise. This addition sequence was necessary to obtain a uniform sol-gel mixture since TTIP is far more reactive than TEOS [50]. Stirring was continued for 1 h and then thiourea (3.65 g, 0.048 mol) was added. After stirring for 3 h, the mixture was transferred into a crystallization dish and dried at 80 °C overnight. Subsequently, the sample was ground for 30 min at 350 rpm by planetary ball-milling, calcined at 500 °C for 2 h in static air and finally ground again as above. The synthesized sample was denoted as S(2)-TiO₂/SiO₂. Two more samples were synthesized using a higher nominal S/Ti ratio. Specifically, 8.2 g (0.108 mol) or 12.79 g (0.168 mol) of thiourea was added to produce S(4)-TiO₂/SiO₂ or S(6)-TiO₂/SiO₂, respectively. For the sake of comparison, pure TiO₂, undoped TiO₂/SiO₂ and unsupported S-doped TiO₂ samples were also synthesized following the same synthetic protocol without adding thiourea or TEOS.
The PXRD patterns were collected on a Bruker D8 Advance (Karlsruhe, Germany) diffractometer with Cu Kα radiation (λ = 0.15418 nm). The accelerating voltage and applied current were 40 kV and 40 mA, respectively. Profiles were measured in the $20^\circ < 2\theta < 80^\circ$ range with a step of 0.04°/2 s. The crystalline phases were identified with reference to the PDF cards of the International Centre for Diffraction Data (ICDD). The average crystallite size of the anatase phase was determined from the intensity of the main (101) reflection using the Scherrer equation, as follows:

$$d (\text{nm}) = \frac{0.89 \times \lambda}{\beta \times \cos(\theta)}$$

where $\lambda$ (nm) is the X-ray wavelength, $\beta$ (rad) is the full width at half the maximum of the peak intensity and $\theta$ is the respective Bragg angle. Particle morphology was investigated by SEM on a Jeol 6380 LV instrument (Tokyo, Japan) equipped with an Oxford Instruments INCA EDS system (Abingdon, UK). The electron beam accelerating voltages were between 15–20 kV. To improve the surface conductivity of the samples, standard gold deposition was applied through vacuum evaporation. UV-Vis spectra in solution and diffuse reflectance spectra in the solid state were measured between 350–800 nm using an Agilent Carry 60 spectrometer (Santa Clara, CA, USA). The DRS measurements were performed using a Harrick Videobarrelino DRA fiber optic coupler (Pleasantville, NY, USA). The band gap was calculated using the Kubelka-Munk (K-M) model by plotting $[F(R) \times E]^{1/2}$ vs. E (eV), where $F(R) = (1 - R)^2/2R$ is the K-M function and R is the reflectance of the materials.

$N_2$ adsorption-desorption isotherms at 77 K were measured on a Quantachrome NOVA 1200 gas analyzer (Boynton Beach, FL, USA). Samples were degassed at 150 °C for 3 h before measurement. The specific surface area ($SSA_{exp}$) of the samples was determined via the BET model using the multipoint method of the Quantochrome NovaWin2 software (version 2.2). The pore size distribution was obtained by applying Barret-Joyner-Halenda (BJH) analysis for the desorption part of the isotherm. The theoretical SSA of the TiO$_2$ nanoparticles ($SSA_{calc}$) was calculated based on the average particle diameter, estimated by XRD ($d_{XRD}$) and the weighted density of anatase ($\rho_A = 3.84$ gr cm$^{-3}$), assuming spherical and non-agglomerated particles, as follows:

$$SSA_{calc} = \frac{6}{(\rho_A \times d_{XRD})}$$

Particle size distribution was also measured via DLS using an Anton Paar Litesizer 500 particle size analyzer (Graz, Austria). Particles were suspended in water via ultrasonication for 1 min prior to measurement. The FTIR spectra of the synthesized samples were measured on a Brucker Tension 27 FTIR spectrometer (Karlsruhe, Germany) equipped with a diamond ATR accessory at a spectral resolution of 4 cm$^{-1}$ in the 4000–600 cm$^{-1}$ range. IR spectra were analyzed with the Bruker OPUS software (version 5.2).

TEM images were collected on a Jeol 2100 HR (Tokyo, Japan) microscope operating at 200 kV. The TEM samples were prepared as follows: a small amount of the photocatalyst was dispersed in ethanol and the suspension was sonicated for 10 min. Afterwards, a single drop of the suspension was placed on a carbon-coated grid and was allowed to dry at ambient temperature. SAED patterns were analyzed with ImageJ software (LOCI, Madison, WI, USA).

### 2.3. Photocatalytic Evaluation Methods

The synthesized powders were evaluated as photocatalysts in the decomposition of pollutants in an aqueous medium using methyl orange (MO) as a model compound, based on the relevant ISO 10678:2010 standard procedure [31]. In a typical experiment, 50 mg of the powder sample was combined with 150 mL of an aqueous solution of MO (2 mg L$^{-1}$). The mixture was sonicated for 15 min, stirred in the dark for 40 min and afterwards irradiated with visible light for 5 h using 4 parallel Daylight 18 W lamps and a
UV cut-off filter (99% UV cut-off capability). Aliquots (3 × 1 mL) were collected at 60 min intervals and centrifuged. The concentration of MO in the supernatant was determined by UV-Vis spectroscopy. A control experiment was also run during which an identical system was kept in the dark for 5 h in order to correct for the amount of MO adsorbed on the surface of the photocatalyst.

Regarding the PCO of air pollutants, nitric oxide (NO) and acetaldehyde (CH₃CHO) were selected as representative airborne inorganic and organic pollutants, respectively. The photocatalytic activity of all samples was investigated following the ISO 22197–1:2007 and ISO 22197-2:2011 standard procedures [52,53]. A detailed description of the experimental set up and parameters are provided in the literature [54]. A constant flow rate of 3 L min⁻¹ and a relative humidity of 50% were maintained during all the experiments. Visible light irradiation (~7000 lux) was applied for 60 min. A short dark period of 5 min preceded irradiation to allow the mixture to reach equilibrium and correct for adsorption of pollutants on the photocatalyst.

The concentration of acetaldehyde was determined by a Shimadzu Tracera High Sensitivity GC 2010 (Kyoto, Japan). The concentrations of NO, NO₂ and NOₓ (NOₓ = NO + NO₂ combined concentration) were determined by a HORIBA 370 analyzer (Kyoto, Japan) by integrating the respective peak area over time. The activity of the photocatalysts was calculated using Equations (3)–(5):

\[
\text{Effectiveness} (\text{NO}) = \frac{\int_{t_0}^{t_1} (\text{NO initial} - \text{NO measured}) dt}{(\text{NO initial}) \times T} \tag{3}
\]

\[
\text{Effectiveness} (\text{NO₂}) = \frac{\int_{t_0}^{t_1} (\text{NO₂ measured} - \text{NO₂ initial}) dt}{(\text{NO₂ initial}) \times T} \tag{4}
\]

\[
\text{Effectiveness} (\text{NOₓ}) = \frac{\int_{t_0}^{t_1} (\text{NOₓ initial} - \text{NOₓ measured}) dt}{(\text{NOₓ initial}) \times T} \tag{5}
\]

where \( T = t_1 - t_0 \) corresponds to the period of visible irradiation, during which the lamp was switch on at \( t = t_0 \) and switched off at \( t = t_1 \). The effectiveness of each photocatalyst per compound corresponds to the change in the compound’s concentration at the end of the irradiation period, expressed as a percentage.

3. Results and Discussion

3.1. Powder XRD and Porosity Analysis

The powder XRD patterns of the TiO₂, TiO₂/SiO₂ and S(4)-TiO₂/SiO₂ samples (Figure 1a) showed diffractions peaks characteristic of anatase TiO₂ (JCPDS 21-1272), whereas peaks corresponding to the rutile or brookite phases were not observed. Therefore, powder XRD indicated that the synthetic protocol adopted herein led to the formation of a single crystalline TiO₂ phase, anatase. The average crystallite size (d_{XRD}) was determined using the Scherrer equation (Equation (1)) on the (101) main diffraction peak of anatase at 2θ = 25.28°. All the diffraction peaks in the XRD patterns of TiO₂/SiO₂ and S(4)-TiO₂/SiO₂ were broader compared with those of pure TiO₂, suggesting that the growth of the TiO₂ particles was restricted when SiO₂ was introduced. This was reflected by the average crystallite size which was significantly reduced from 20 nm for pure TiO₂, to 7.6 nm for TiO₂/SiO₂ and 6.7 nm for S(4)-TiO₂/SiO₂ (Table 1). Therefore, S-doping resulted in even smaller particles, probably due to the substitution of Ti⁴⁺ by S⁶⁺ and S⁴⁺ cations, which can further restrict the growth of TiO₂ particles [28].

These observations were supported by N₂ adsorption-desorption measurements. The BET plots and the BJH pore size distributions for TiO₂, TiO₂/SiO₂ and S(4)-TiO₂/SiO₂ are presented in Figure 1b,c, respectively. The BET surface area and the pore volume increased from SSA_{exp} = 60 m² g⁻¹ and V_p = 0.198 cm³ g⁻¹ for TiO₂ to SSA_{exp} = 195 m² g⁻¹ and V_p = 0.315 cm³ g⁻¹ for TiO₂/SiO₂ (Table 1), in good agreement with the calculated values (SSA_{calc}).
The higher porosity was accompanied by a decrease in pore diameter from 7.87 nm for TiO\textsubscript{2} to 4.67 nm for TiO\textsubscript{2}/SiO\textsubscript{2}. This, in turn, suggested that the \(d_{XRD}\) values calculated from the (101) diffraction peak of the respective powder XRD patterns nearly represented the true size of the TiO\textsubscript{2} nanoparticles, as verified by TEM images (see below Section 3.4). S-doping of TiO\textsubscript{2}/SiO\textsubscript{2} had no effect on the BET surface area \((\text{SSA}_{\text{exp}} = 195 \text{ m}^2 \text{ g}^{-1})\) compared with TiO\textsubscript{2}/SiO\textsubscript{2}. However, the pore volume \((V_P = 0.277 \text{ cm}^3 \text{ g}^{-1})\) and pore diameter (3.67 nm) decreased compared with undoped TiO\textsubscript{2}/SiO\textsubscript{2}. This indicated that S-doping resulted in partial pore blocking, in agreement with literature reports \cite{22,24,55}.

**Table 1.** Textural properties of TiO\textsubscript{2}, TiO\textsubscript{2}/SiO\textsubscript{2} and S(4)-TiO\textsubscript{2}/SiO\textsubscript{2} samples.

| Sample          | \(d_{XRD}\) [\text{nm}] | SSA\textsubscript{calc} [\text{m}^2 \text{ g}^{-1}] | SSA\textsubscript{exp} [\text{m}^2 \text{ g}^{-1}] | \(V_P\) [\text{cm}^3 \text{ g}^{-1}] | Pore Diameter (nm) |
|-----------------|--------------------------|--------------------------|--------------------------|--------------------------|---------------------|
| TiO\textsubscript{2} | 20                       | 78                       | 60                       | 0.198                    | 7.85                |
| TiO\textsubscript{2}/SiO\textsubscript{2} | 7.6                      | 206                      | 195                      | 0.315                    | 4.67                |
| S(4)-TiO\textsubscript{2}/SiO\textsubscript{2} | 6.7                      | 233                      | 195                      | 0.277                    | 3.67                |

\[^{[a]}\text{Particle size based on powder XRD,}^{[b]}\text{Calculated specific surface based on} \(d_{XRD},^{[c]}\text{Experimental specific surface based on BET model,}^{[d]}\text{Pore volume.}\]

### 3.2. FTIR Spectroscopy

FTIR analysis was performed to identify the nature of the chemical bonds in the synthesized samples. The FTIR spectra of TiO\textsubscript{2}, TiO\textsubscript{2}/SiO\textsubscript{2} and S(x)-TiO\textsubscript{2}/SiO\textsubscript{2} (\(x = 2, 4, 6\)) are presented in Figure 2. The peaks at 3300–3500 cm\(^{-1}\) and 1640 cm\(^{-1}\) were observed in all spectra. These peaks were assigned to the stretching and bending vibrations of adsorbed water molecules and surface hydroxyl groups, as previously reported \cite{24}. Two more peaks
at 1028 cm\(^{-1}\) and 907 cm\(^{-1}\) were observed in the FTIR spectrum of undoped TiO\(_2\)/SiO\(_2\), which have previously been assigned to the stretching vibration of the Si–O–Si and Ti–O–Si groups, respectively [30,56]. The detection of the Ti–O–Si vibration at 907 cm\(^{-1}\) was consistent with the incorporation of Ti into the silica framework in undoped TiO\(_2\)/SiO\(_2\).

![FTIR spectra](image)

Figure 2. FTIR spectra of (as prepared) TiO\(_2\), TiO\(_2\)/SiO\(_2\) and S-doped TiO\(_2\)/SiO\(_2\) samples.

With respect to the S(x)-TiO\(_2\)/SiO\(_2\) samples, the intensity of the \(v\)(O–H) peaks at 3300–3500 cm\(^{-1}\) and \(\delta\)(O–H) at 1640 cm\(^{-1}\) increased with the increasing nominal S/Ti ratio, indicative of a gradually greater number of surface hydroxyl groups at higher levels of S-doping [30]. Moreover, three peaks were observed at 2050, 1400 and 1060 cm\(^{-1}\), which were absent in the FTIR spectrum of undoped TiO\(_2\)/SiO\(_2\). Based on previous reports on S-doped TiO\(_2\) [23,24,30,57], the higher energy peak at 2050 cm\(^{-1}\) was assigned to the C–N stretching vibration of the isothiocyanate group –NCS, whereas the two lower energy peaks at 1400 and 1060 cm\(^{-1}\) were attributed to Ti–S and Ti–O–S stretching vibrations. It should be underlined that the intensity of the above peaks increased when larger quantities of thiourea were added in the reaction mixture. These observations were consistent with the presence of Ti–NCS and Ti–S bonds in the synthesized photocatalysts.

3.3. UV–Vis Diffused Reflectance Spectroscopy: Band Gap Analysis

The DR spectra for all samples are presented in Figure 3a. As expected, the colorless semiconductors TiO\(_2\) and TiO\(_2\)/SiO\(_2\) showed a single absorption edge in the UV region of the spectrum. By contrast, a second absorption edge in the visible region was observed for all the S-doped TiO\(_2\)/SiO\(_2\) nanopowders, suggesting the presence of a localized band above the main valence band of TiO\(_2\), as previously reported [22,30,58,59]. The respective Tauc plots (Figure 3b), based on the Kubelka-Munk model, revealed that the energy band gap values corresponding to each edge—Eg1 and Eg2, respectively—decreased as the amount of thiourea in the sol-gel mixture increased (Table 2). As a result, the lowest Eg2 value of 2.22 eV was found for S(6)-TiO\(_2\)/SiO\(_2\), the sample with the higher level of S-doping.
3.3. UV–Vis Diffused Reflectance Spectroscopy: Band Gap Analysis

The DR spectra for all samples are presented in Figure 3a. As expected, the colorless semiconductors TiO2 and TiO2/SiO2 showed a single absorption edge in the UV region of the spectrum. By contrast, a second absorption edge in the visible region was observed for all the S-doped TiO2/SiO2 nanopowders, suggesting the presence of a localized band above the main valence band of TiO2, as previously reported [22,30,58,59]. The respective Tauc plots (Figure 3b), based on the Kubelka-Munk model, revealed that the energy band gap values corresponding to each edge—Eg1 and Eg2, respectively—decreased as the amount of thiourea in the sol-gel mixture increased (Table 2). As a result, the lowest Eg2 value of 2.22 eV was found for S(6)-TiO2/SiO2, the sample with the higher level of S-doping.

![Figure 3. (a) Diffuse reflectance spectra and (b) Tauc plots for TiO2, TiO2/SiO2 and S-doped TiO2/SiO2 samples.](a)

![Figure 3. (a) Diffuse reflectance spectra and (b) Tauc plots for TiO2, TiO2/SiO2 and S-doped TiO2/SiO2 samples.](b)

Table 2. Band gap values (Eg1 and Eg2) of TiO2, TiO2/SiO2 and S(x)-TiO2/SiO2 samples.

| Sample               | Eg1 (eV) | Eg2 (eV) |
|----------------------|----------|----------|
| TiO2                 | 3.11     | -        |
| TiO2/SiO2            | 3.17     | -        |
| S(2)-TiO2/SiO2       | 3.08     | 2.33     |
| S(4)-TiO2/SiO2       | 3.07     | 2.29     |
| S(6)-TiO2/SiO2       | 3.04     | 2.22     |

3.4. Morphological Analysis

Figure 4 shows characteristic SEM images, EDS analysis and DLS particle size distribution of TiO2, TiO2/SiO2 and S(4)-TiO2/SiO2. Aggregates with particle sizes ranging from 400 nm to 40 μm were observed for all samples. However, the S(4)-TiO2/SiO2 powder had a more porous morphology and the particle size was relatively confined. Importantly, EDS analysis verified the presence of silicon in TiO2/SiO2, as well as sulfur and silicon in S(4)-TiO2/SiO2. To probe the effect of silica and S-doping on particle size, DLS measurements were also performed after dispersing the NPs in water. Supporting TiO2 on silica led to a decrease in particle size from \( d = 1003 \pm 218 \) nm for TiO2 to \( d = 605 \pm 323 \) for TiO2/SiO2. S-doping led to even smaller particles with a narrow size distribution, i.e., \( d = 416 \pm 119 \) nm for S(4)-TiO2/SiO2.
Figure 4. (a) SEM images, (b) EDS analysis and (c) DLS analysis of TiO$_2$, TiO$_2$/SiO$_2$ and S(4)-TiO$_2$/SiO$_2$ samples.

More detailed morphological and structural characterization of the samples was carried out using TEM (Figure 5). The TEM micrographs of all samples revealed numerous aggregates and ill-defined nanostructures. However, the TiO$_2$/SiO$_2$ and S(4)-TiO$_2$/SiO$_2$ samples demonstrated better dispersion and smaller crystallite size compared with pure TiO$_2$. Specifically, the crystallite size calculated from the TEM images was approximately 20 nm for pure TiO$_2$ (Figure 5a), in accordance with the XRD results. The introduction of SiO$_2$ and S-doping resulted in considerably smaller particles, between 5–10 nm. Importantly, the TEM images of the composites TiO$_2$/SiO$_2$ and S(4)-TiO$_2$/SiO$_2$ revealed that the TiO$_2$ nanoparticles were almost completely surrounded by a silica layer (Figure 5b,c). As has already been reported, the opposite surface charge of TiO$_2$ and SiO$_2$ could promote the encapsulation of TiO$_2$ nanoparticles into the SiO$_2$ matrix through electrostatic interactions [43,44]. It is, therefore, likely that encapsulation of TiO$_2$ inside the SiO$_2$ matrix restricted the accumulation of larger TiO$_2$ nanoparticles during the synthesis of the photocatalysts.
Furthermore, SAED patterns showed a sequence of spots for TiO$_2$ and rings for undoped TiO$_2$/SiO$_2$ (inset—Figure 5a,b). The detection of distinct diffraction spots in the case of TiO$_2$ suggested a more crystalline sample and a larger crystallite size. On the contrary, the observation of diffraction rings in the case of TiO$_2$/SiO$_2$ indicated a less crystalline sample and a smaller particle size. Therefore, the TEM images were in accordance with the powder XRD analysis (Section 3.1). Moreover, the interplanar spacings derived from the diffraction rings were $d_{(215)} = 0.35$ nm, $d_{(004)} = 0.24$ nm, $d_{(200)} = 0.19$ nm, $d_{(211)} = 0.17$ nm, $d_{(204)} = 0.15$ nm and $d_{(211)} = 0.13$ nm. These values corresponded to the anatase TiO$_2$ crystalline phase (PDF no 21-1272), as also suggested by the powder XRD pattern.

![Figure 5. TEM images and SAED patterns (inset) of (a) TiO$_2$, (b) TiO$_2$/SiO$_2$ and (c) S(4)-TiO$_2$/SiO$_2$ samples.](image)

3.5. Photocatalytic Evaluation

3.5.1. Liquid Pollutant Degradation

The photocatalytic activity of the as-synthesized samples was initially benchmarked against the photocatalytic oxidation of MO under visible light irradiation. The results are presented in Figure 6a alongside the related control experiments, which were carried out in the dark. As expected, pure TiO$_2$ showed negligible activity, removing only 4.2% of MO in solution after 5 h. A minor increase of activity was observed for undoped TiO$_2$/SiO$_2$, which afforded 14.6% MO removal, most likely due to the smaller TiO$_2$ particle size and the larger porosity of the TiO$_2$/SiO$_2$ nanocomposite, as shown above (Table 1).
Figure 6. (a) MO removal (%) in aqueous media over TiO2, TiO2/SiO2, S(4)-TiO2 and S(x)-TiO2/SiO2 samples under visible light irradiation for 5 h; (b) Overlay of absorbance spectra for MO degradation over S(4)-TiO2/SiO2, collected at 1 h intervals.

The S-doped samples showed significantly higher activity. S(4)-TiO2/SiO2 was the most efficient photocatalyst, affording 79.0% MO removal after 5 h. Figure 6b shows the corresponding UV-Vis absorption spectra, collected at 1 h intervals. The intensity of the characteristic absorption peak of MO at 470 nm rapidly decreased in 5 h. The S(2)-TiO2/SiO2 and S(6)-TiO2/SiO2 photocatalysts afforded 31.4% and 72.2% MO removal, respectively. Therefore, an optimal thiourea-to-titanium nominal ratio was required to maximize the photocatalytic activity in MO degradation. As previously reported [22,24,60], lower levels of S-doping limit the ability of the catalyst to operate under visible light, whereas higher levels of S-doping enhance charge recombination. Finally, it should be noted that the unsupported S(4)-TiO2 photocatalyst afforded 73.9% MO removal in 5 h. Therefore, the use of silica as a matrix did not significantly improve the photocatalytic activity in MO degradation despite the higher porosity of the SiO2-supported samples, most likely due to the rather large size of MO, which in turn, slows down its diffusion within the pores.

In summary, S-doping of TiO2 resulted in a significantly higher photocatalytic activity in terms of MO degradation under visible light irradiation. This could be attributed to the narrower band gap induced by S-doping (Figure 3b), as well as the higher number of surface hydroxyl groups in the S-doped TiO2 samples, as indicated by FTIR spectroscopy (Figure 2) [61].

3.5.2. Air Pollutant Degradation

The photocatalytic activity of the as-synthesized samples was also evaluated in the photocatalytic degradation of CH3CHO and NO(g). Figure 7 presents a comparison of the effectiveness of the photocatalysts, i.e., the change in concentration of each compound, expressed as a percentage. As also observed for MO degradation, the S-doped catalysts were significantly more active under visible light irradiation compared with TiO2, which exhibited negligible activity. Specifically, the acetaldehyde concentration was reduced by 6.0% over unsupported S(4)-TiO2, by 14.5% over S(2)-TiO2/SiO2 and by 26.4% over S(4)-TiO2/SiO2 in 1 h under visible light irradiation (Figure 7a). Therefore, entraining S-doped TiO2 NPs in a silica matrix, clearly had a beneficial effect in the case of acetaldehyde degradation, a small molecule that can readily diffuse through the pores.
With respect to the photocatalytic degradation of NO(g) over TiO$_2$-based materials, it is widely accepted that NO is initially oxidized to NO$_2$(g) and subsequently to NO$_3^-$ by photogenerated hydroxyl (•OH) rather than superoxide (•O$_2^-$) radicals [62]. Along these lines, Todorova et al. reported that N,S co-doping of TiO$_2$ using thiourea as the source had a detrimental effect on the photocatalytic degradation of NO$_x$ under visible-light irradiation because the formation of •OH radicals was suppressed [25].

On the contrary, the TiO$_2$-based photocatalysts synthesized in this work using thiourea as the source and silica as a supporting matrix were moderately active in NO$_x$ decomposition under visible-light irradiation. In particular, the S(2)-TiO$_2$/SiO$_2$ catalyst afforded 5.9% NO oxidation to NO$_2$ and 2.2% combined NO$_x$ removal in 1 h. The catalytic activity significantly increased over S(4)-TiO$_2$/SiO$_2$, which afforded 12.3% NO oxidation and 4.5% total NO$_x$ removal in 1 h. Clearly, the higher porosity and the smaller particle size of the photocatalysts associated with the use of silica as a matrix (see Table 1) promoted the adsorption of •OH radicals on the catalytic surface, which in turn, resulted in the enhancement of NO to NO$_2$ oxidation. Nevertheless, NO$_2$ was only partially further oxidized to NO$_3^-$, resulting in lower efficiency values with respect to total NO$_x$ removal (NO + NO$_2$). This suggested that NO$_3^-$ products could partially block active sites on the catalytic surface.

4. Conclusions

This study demonstrated the feasibility of producing silica-supported S-doped TiO$_2$ nanopowders with substantial activity in the photocatalytic degradation of liquid (MO) and air (CH$_3$CHO and NO$_x$) pollutants under visible light irradiation. S-doping using thiourea as the source shifted the absorption towards the visible region of the spectrum to enable photocatalytic activity under visible light. Moreover, the ‘safe-by-design’ strategy of entraining the S-doped TiO$_2$ NPs within a silica layer improved the photocatalytic activity by increasing the surface area, reducing the particle size and facilitating the diffusion of small-sized pollutants toward the catalytically active sites.

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![Figure 7](https://example.com/figure7.png)

**Figure 7.** (a) Acetaldehyde photocatalytic degradation (%) in air over TiO$_2$, S(4)-TiO$_2$ and S(x)-TiO$_2$/SiO$_2$ samples; (b) NO$_x$ photocatalytic degradation (effectiveness, %) in air over TiO$_2$, S(2)-TiO$_2$/SiO$_2$ and S(4)-TiO$_2$/SiO$_2$. Conditions: flow rate of 3 L min$^{-1}$, RH of 50% and visible light irradiation (~7000 lux) for 1 h.
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