Nitrogen doped TiO\textsubscript{2} photoactive in visible light

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ABSTRACT

In this research TiO\textsubscript{2} modified with nitrogen has been prepared. The synthesis method used was sol gel from the hydrolysis of titanium tetrabutoxide with urea solution, combined with a low temperature treatment at atmospheric pressure. The theoretical mole ratio N/Ti was equal to 0.5 and the theoretical mole ratio HNO\textsubscript{3}/TiO\textsubscript{2} was equal to 0.5. The solid obtained was calcined at different temperatures in static air with high heating rate and residence time of 1 hour. The samples were characterized by X-ray diffraction, UV-Vis diffuse reflectance spectroscopy, FTIR Raman and X-ray diffraction revealed for all the heat treatments, the presence of crystalline phases corresponding to anatase and brookite polymorphism. The photocatalytic activity of the catalysts was tested in degradation of NOx. Asahi at al. reported that TiO\textsubscript{2} doped with different anions (C, N, S and F) concluded that the presence of substitutional nitrogen was necessary to reduce the band gap and to display photocatalytic activity under visible light.

Keywords: TiO\textsubscript{2}-N, synthesis. Characterization, photocatalysis

1. INTRODUCTION

Heterogeneous photocatalysis is considered one of the best alternatives to solve problems of energy and environmental pollution [1,2]. Titanium dioxide (TiO\textsubscript{2}) has been employed as an efficient photocatalyst because it is relatively inexpensive, chemical and thermal stability, optics and electronics properties, non-toxicity and high photoactivity [2-4]. However, its high value of band gap hinders the use of solar radiation as primary energy source. To perform photocatalysis under visible irradiation, several methods have been applied in order to improve the visible response of TiO\textsubscript{2}.

In 1986, Sato [5] reported for the first time the synthesis of nitrogen-doped TiO\textsubscript{2} (N-TiO\textsubscript{2}) based on calcination of titanium hydroxide (H\textsubscript{2}TiO\textsubscript{3}). The response to visible light was attributed to sensitization by impurities NO\textsubscript{x}. Asahi at al. [6] obtained N-TiO\textsubscript{2} by sputtering of TiO\textsubscript{2} under atmosphere of N\textsubscript{2}-Ar. The solid showed an absorption edge shift to the visible light region. Based on the calculation of the density of states for TiO\textsubscript{2} doped with different anions (C, N, S and F) concluded that the presence of substitutional nitrogen was necessary to reduce the band gap and to display photocatalytic activity under visible light.

Since this work, TiO\textsubscript{2} doped with nitrogen has been synthesized by different methods, including reduction in the presence of gaseous NH\textsubscript{3} [7,8], oxidation of TiN [9,10], treatment of mixtures of urea and TiO\textsubscript{2} [11], sputtering of TiO\textsubscript{2} in atmosphere of N\textsubscript{2} [12] and hydrolysis of precursors containing Ti [13]. All these methods require treatments at high temperatures to obtain crystalline solids, thus reducing the surface of materials by sintering and in some cases removing the doping element. In order to solve this problem, the synthesis of doped TiO\textsubscript{2} by hydrothermal or solvothermal treatment of TiO\textsubscript{2} and N precursors, allows to produce crystalline solids at low temperature and with important textural properties.

The present work reports the synthesis of nitrogen-doped TiO\textsubscript{2} prepared by sol-gel in combination with thermal treatment at atmospheric pressure. The catalysts were characterized by UV-Vis diffuse reflectance spectroscopy, X-ray diffraction, FTIR and Raman spectroscopy, thermogravimetric and thermodiffer-
ential analysis and specific surface. The photocatalytic activity under visible light was also investigated.

2. MATERIALS AND METHODS

2.1 Reagents
Titanium tetrabutoxide TBT (Ti[O(CH₃)₂CH₃]₄ 97%) from Aldrich and urea (NH₂CONH₂) from Cicarelli laboratories were used as TiO₂ and nitrogen precursors respectively. Nitric acid (HNO₃), tartrazine (an azo dye) and n-butyl alcohol of analytical reagent grade were used. All solutions and experiments were carried out using bidistilled water.

2.2 Synthesis of N-TiO₂
First 370 mg of urea, to obtain a molar ratio N / Ti of 0.5, were dissolved in water (solution A). On the other hand, titanium tetrabutoxide was dissolved in n-butyl alcohol to obtain a 1 mol L⁻¹ solution. 25.04 mL of this solution was transferred to a beaker and 88 μL of concentrated HNO₃ were added dropwise under vigorous stirring during one hour (solution B).

The solution A was then added to solution B under slow stirring. Then the new solution was placed into a water bath at 40°C and kept under stirring for two hours to allow the hydrolysis process and the doping with nitrogen. Finally it was cooled up to room temperature. The obtained powder (NTiO₂) was calcined at 200, 250 and 300°C in static air with high heating rate and a residence time of 1 hour. The catalysts were referred as NTiO₂-x where x is the calcining temperature.

2.3 Characterization
XRD patterns of the prepared NTiO₂-x samples were recorded on a Philips PW 1710 diffractometer using Cu-Kα radiation (λ=1.5406 Å) over the 20 ranges of 20-80 degree. FTIR and Raman spectra were collected on a Perkin Elmer GX FT-IR equipped with a laser as excitation source. Diffuse reflectance spectra were measured on a UV-vis spectrophotometer equipped with an integrating sphere with BaSO₄ as the reference sample. The analysis range was 200 to 800 nm. Kubelka Munk function was used to transform data from diffuse reflectance into absorption coefficients and for obtaining the Band Gap value. Thermodifferential and thermogravimetric analysis were performed in static air on a Rigaku TAS 100 equipment. Specific surface area was measured on a Micromeritics FlowSorb II, for these samples were outgassed at 100°C for 30 min.

2.4 Photodegradation experiment
The photocatalytic activities of the catalysts were evaluated by measuring the decomposition of 1 x 10⁻⁵ mol L⁻¹ solution of tartrazine. Experiments were carried out at room temperature in a batch reactor thermostatically controlled. A 250 W halogen lamp was used as the light source in the home made photo-reactor (Fig. 1), a sodium nitrite solution was used to cut off the light having wavelength below 400 nm.

Figure 1: Photography of home made photo-reactor.
In a typical run, 0.02 g of the catalyst was weighed and added into a 100 mL of the above mentioned tartrazine solution. Prior to illumination, the suspension was magnetically stirred for 30 min to establish the adsorption/desorption equilibrium between the photocatalyst and tartrazine. At specific time intervals, about 5 mL aliquots were sampled. The filtrates were analyzed by recording variations in the absorption in UV–vis spectra of tartrazine, using a UV–vis spectrophotometer.

3. RESULTS AND DISCUSSION

3.1 DTA-TG

The results of thermogravimetric and thermodifferential analysis TG- DTA of NTiO$_2$ are shown in Fig. 2.

The TG curves can be separated in several areas. The first area, from room temperature to 180ºC where the mass loss is 8%, corresponds to endothermic process showed in DTA curve. It is due to the removal of n-butyl alcohol and water, physically adsorbed and decomposition of nitrate groups [14]. The second area, from 180ºC to 236ºC, corresponds to a mass loss of around 8% and it is attributed to the decomposition of urea (Tb 235ºC) in according to spectrum IR. This process is associated to the endothermic peak at 200ºC.

The third area, from 236ºC to 260ºC, corresponds to a mass loss of 11%, attributed to the combustion and the decomposition of the non-hydrolyzed groups of TBT [15]. This process corresponds to the exothermic peaks at 245ºC. Between 260ºC and 345ºC, slow oxidation processes take place associated with a broad exothermic heat effect, which stops at 345ºC. The peak at 388ºC is attributed to the crystallization of amorphous TiO$_2$ into anatase with a mass loss (1%) due to the removal of the hydroxyl groups trapped inside the particles [16].

![Figure 2: DTA and TG of NTiO$_2$.](image)

3.2 XRD

Fig. 3 shows the XRD patterns of the prepared samples before and after being heat treated. The patterns indicate that all samples exhibit the presence of anatase (A) and brookite (B) phases. The existence of brookite is discernable from its diffraction line (121) located at 30.8º (20) in XRD patterns, which is not overlapped with any peak of anatase or rutile phases. The phase content was estimated from equation (1) and (2).

\[
W_A = \frac{K_{A1A}}{K_{A1A} + K_{B1B}}
\]

\[
W_B = \frac{K_{B1B}}{K_{A1A} + K_{B1B}}
\]
Where $I_A$ is the area of diffraction of the peak of anatase (101) and $I_B$ is the area of diffraction of the peak of brookite (121). $K_A$ and $K_B$ are constants equal to 0.886 and 2.721, respectively [17].

\[
D = \frac{K\lambda}{\beta \cos \theta}
\]

(3)

Where $D$ is the crystallite size, $K$ is an adimensional constant, $2\theta$ is the diffraction angle, $\lambda$ is the X-ray wavelength corresponding to the Cu Kα irradiation and $\beta$ is the half-height width of the diffraction peaks.

In Table 1, the values of crystal size and the percentage of the two crystalline phases are presented.

**Table 1:** Crystal size and content of anatase and brookite phases.

<table>
<thead>
<tr>
<th>SAMPLE</th>
<th>$D_A$(NM)</th>
<th>$D_B$(NM)</th>
<th>$W_A$</th>
<th>$W_B$</th>
</tr>
</thead>
<tbody>
<tr>
<td>NTiO₂ 70</td>
<td>4</td>
<td>6.1</td>
<td>0.90</td>
<td>0.10</td>
</tr>
<tr>
<td>NTiO₂ 200</td>
<td>4.1</td>
<td>6.5</td>
<td>0.91</td>
<td>0.09</td>
</tr>
<tr>
<td>NTiO₂ 250</td>
<td>5.1</td>
<td>6.5</td>
<td>0.87</td>
<td>0.13</td>
</tr>
<tr>
<td>NTiO₂ 300</td>
<td>5.7</td>
<td>6.6</td>
<td>0.86</td>
<td>0.14</td>
</tr>
</tbody>
</table>

All samples contain a small fraction of brookite, not exceeding 14%. The size of the crystallites increases 42% and 7% for anatase and brookite after heat treatment at 300 °C. The crystal sizes are in the range of 4 to 5.7 nm for anatase, and from 6.1 to 6.6 nm for brookite, in agreement with results reported by Alphonse [16].

The diffractograms were also analyzed by Rietveld method. The unit cell parameters $a$, $b$ and $c$ determined for anatase phase are presented in Table 2.
Table 2: Cell parameters and volume for the anatase phase of each of the samples.

<table>
<thead>
<tr>
<th>SAMPLE</th>
<th>a(Å)</th>
<th>b(Å)</th>
<th>c(Å)</th>
<th>VOL(Å³)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NTiO₂ 70</td>
<td>3.7941</td>
<td>3.7941</td>
<td>9.4797</td>
<td>136.4598</td>
</tr>
<tr>
<td>NTiO₂ 200</td>
<td>3.7887</td>
<td>3.7887</td>
<td>9.4539</td>
<td>135.7050</td>
</tr>
<tr>
<td>NTiO₂ 250</td>
<td>3.7854</td>
<td>3.7854</td>
<td>9.4688</td>
<td>135.6820</td>
</tr>
<tr>
<td>NTiO₂ 300</td>
<td>3.7848</td>
<td>3.7848</td>
<td>9.4724</td>
<td>135.6877</td>
</tr>
</tbody>
</table>

The parameters a, b and c for all prepared catalysts show small differences as compared to TiO₂ anatase phase reference values (JCPDS No. 21-1272, a=b=0.37852 nm and c=0.95139 nm). These differences are due to lattice distortion as a result of nitrogen doping.

It is observed that a parameter decrease when the temperature thermal treatment increased. This is due to the fact that during heat treatment the nitrogen is removed, decreasing its concentration in the solid framework [18] and therefore the value of $a$ for doped TiO₂ tends to the value of undoped TiO₂. While, $c$ shows an opposite behaviour, but also tends to the reference value [19].

3.3 Raman

Raman spectra of NTiO₂ and NTiO₂ 250 are showed in Fig. 4. NTiO₂ presents Raman peaks at 649, 519 and 408 cm⁻¹, whereas that of NTiO₂ 250 shows peaks at 644, 519 and 401 cm⁻¹. These are very similar to the Raman active fundamental modes of anatase phase (639, 519, and 394 cm⁻¹), the first is assigned to vibrational mode $E_g$ and the last two to the vibrational modes $B_{1g}$.

The vibrational modes of the synthesized (uncalcined and calcined) samples present significant shift to blue region. According to literature, the blue shift of the Raman peaks could be attributed to quantum size effect because of the small size of the particles [20], which implies that the solid with smaller size presents the highest shift. The blue shift could also be attributed to oxygen vacancies [21].

The bands at 245, 326, 366, 449 and 589 cm⁻¹ are attributed to brookite phase [22].

The band intensity of the two crystalline phases increases with heat treatment, in agreement with the results observed in XRDs.

![Raman spectra](image)

**Figure 4**: Raman spectra of NTiO₂ (black) and NTiO₂ 250 (gray).

3.4 Diffuse Reflectance

Generally speaking, the narrower is the band gap energy of a semiconductor, the higher is the photocatalytic activity. In order to investigate the optical response of uncalcined and calcined NTiO₂ samples, the UV–visible absorption spectras were obtained by applying the Kubelka Munk function to the data of diffuse reflectance. These results are shown in Fig. 5.
The absorption edge for NTiO\textsubscript{2} and NTiO\textsubscript{2} 200 is shifting more towards the visible as compared to NTiO\textsubscript{2} calcined at 250 and 300°C, as is observed in the magnification of Fig 3. N doping could lead to lattice deformation and to form vacancies, this is probably the result of an impurity state in TiO\textsubscript{2} band gap. The existing impurity state improves the absorption of visible light by narrowing the band gap of TiO\textsubscript{2}. However, for NTiO\textsubscript{2} 250 and NTiO\textsubscript{2} 300, the absorption edge is lower because during the heat treatment the carbon and nitrogen were removed.

Unlike of the others samples, NTiO\textsubscript{2} 250 absorbs throughout all the visible light range, which may be attributed to the electron transition from energy levels of N surface species to the conduction band of TiO\textsubscript{2} \cite{23, 24}. Also this effect is probably due to the presence of carbon from residual groups O-R of the alkoxide due to incomplete hydrolysis reaction of titanium butoxide.

The band gap energy of the solids were obtained by extrapolating the linear zone of the graph \( [F(R^{\infty})]^{1/2} \) to the axis of the abscissa (photon energy \( h\nu \)), where \( F(R^{\infty}) \) is the remission function of Kubelka Munk \cite{25}. The band gap value and the corresponding wavelength are presented in Table 3.

<table>
<thead>
<tr>
<th>SAMPLE</th>
<th>( E_g ) (eV)</th>
<th>( \lambda ) (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NTiO\textsubscript{2} 70</td>
<td>3.10</td>
<td>400</td>
</tr>
<tr>
<td>NTiO\textsubscript{2} 200</td>
<td>3.12</td>
<td>398</td>
</tr>
<tr>
<td>NTiO\textsubscript{2} 250</td>
<td>3.20</td>
<td>388</td>
</tr>
<tr>
<td>NTiO\textsubscript{2} 300</td>
<td>3.17</td>
<td>391</td>
</tr>
</tbody>
</table>

3.5 FTIR

Fig. 6 shows the FTIR spectra of the synthesized TiO\textsubscript{2} at different calcination temperatures in the wave-number range of 4000–400 cm\textsuperscript{-1}.
Figure 6: FTIR spectra of NTiO$_2$ uncalcined and calcined.

In all the spectra, the intense broad band observed in the region around 3000–3500 cm$^{-1}$ is due to O-H stretching vibration of surface hydroxyl groups and adsorbed water molecules. The corresponding bending vibration appears close to 1625 cm$^{-1}$ [26-28].

For fresh NTiO$_2$ the bands at 3350 cm$^{-1}$ and 3179 cm$^{-1}$ can be attributed to N-H stretching modes and the bands at 1647 cm$^{-1}$ and 1569 cm$^{-1}$ are due to stretching C=O and bending N-H modes of urea. The intense broad band observed at 1382 cm$^{-1}$ corresponds to stretching modes of NO$_3^-$ and 1037 cm$^{-1}$ are indicative of the presence of residual species of urea and non-hydrolyzed groups of TBT [29].

It was observed that the intensity of 1384 cm$^{-1}$ peak decreased when the calcination temperature increased from 250 to 300ºC and the complete disappearance of this peak takes place at 400ºC (not showed here). The peaks attributed to species of nitrogen and titanium precursors disappear at 250ºC in accordance with the results of DTA.

At temperatures above 300ºC, the intensity of 3400 cm$^{-1}$ and 1625 cm$^{-1}$ bands decreased due to removal of the remaining hydroxyl groups which are probably trapped inside the crystallites [14].

3.6 Specific Surface Area

The values of specific surface area of the synthesized solids and after heat treatment are showed in Table 4.

Table 4: Specific surface area values of the samples.

<table>
<thead>
<tr>
<th>SAMPLE</th>
<th>S(m$^2$/g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NTiO$_2$ 70</td>
<td>33</td>
</tr>
<tr>
<td>NTiO$_2$ 200</td>
<td>149</td>
</tr>
<tr>
<td>NTiO$_2$ 250</td>
<td>165</td>
</tr>
<tr>
<td>NTiO$_2$ 300</td>
<td>146</td>
</tr>
</tbody>
</table>

The surface area of the solid after the hydrothermal treatment at 40ºC is the lower, due to the presence of O-R residual groups at the surface. During the calcination process, residual groups from nitrogen and titanium precursors are removed, thus increasing the surface area.

The surface reaches a maximum at 250ºC, temperature at which it is produced the combustion of the organic residual groups, according to the DTA results. With the subsequent increase of temperature until 300ºC the surface decreased to 146 m$^2$/g produced by sintering.

3.7 Photocatalytic Activity

Adsorption equilibrium is reached for all the tested photocatalysts, after stirring for 30 minutes in the dark. Under the visible light irradiation, the colour of the solutions gradually fade out and the absorbance of tartra-
zine in the visible light region significantly decreases.

In Fig. 7 the degradation curves of tartrazine for NTiO$_2$ 250, NTiO$_2$ 300 and undoped TiO$_2$ calcined at 250°C are shown. These are expressed as degradation percentages of dye as time function and were calculated with the equation 4.

$$ Deg\% = \frac{C_0 - C}{C_0} \times 100 $$

(4)

Where $C_0$ is the initial concentration of the solution and $C$ is the solution concentration at a specific time.

It was observed that the tartrazine concentration hardly changes when the solution is irradiated with visible light for 180 min in the absence of any catalyst, indicating that the photolysis of tartrazine is negligible. The degradation rate of tartrazine on different TiO$_2$ catalyst samples follows the order: NTiO$_2$ 250 > TiO$_2$ 250 > NTiO$_2$ 300.

NTiO$_2$ 200, not showed here, does not present photocatalytic activity despite that is the solid with smallest bang gap. This can be due to the presence of carbon residues from incomplete hydrolysis of the titanium precursor that decreases the efficiency, because it increased the recombination rate of charge carriers.

The activity of NTiO$_2$ 250 was about 4 times of that of the undoped TiO$_2$ calcined at the same temperature (TiO$_2$ 250) while NTiO$_2$ 300 degraded only 4% of dye. It is attributed to absorption in the visible range because the undoped sample absorbs more visible light than NTiO$_2$ 300 due to the presence of carbon.

Regarding to dye adsorption, NTiO$_2$ 250 adsorbs 57% of the initial concentration, while NTiO$_2$ 300 adsorbs only 8%. Even if these results are in agreement with the observed decrease of the surface area, the great difference may be due to the presence of surface groups that interact differently with the dye.

The largest photoactivity showed by the NTiO$_2$ 250 solid in the tartrazine degradation can be attributed to the capacity of energy absorption in the visible region, high specific surface area and presence of two crystalline phases [30–32].

4. CONCLUSION

The technique of synthesis used in this work allows to successfully incorporating nitrogen in the titania lattice, developing solids with a crystalline structure well defined and composed by anatase and brookite phases.

The results of photocatalysis allow confirming that the incorporation of nitrogen in the TiO$_2$ effectively produces a narrowing of the band gap in the catalysts allowing obtaining a good photoactivity under visible light radiation.

Temperature of thermal treatment is a critical variable, because best catalytic behaviour was obtained
for solids calcined at 250 °C. Higher heat temperatures results in a deactivation due to nitrogen lost from the titania lattice.

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6. BIBLIOGRAPHY