Photocatalytic Studies of TiO$_2$/SiO$_2$ Nanocomposite Xerogels

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Abstract

The use of titania-silica materials in photocatalytic processes has been proposed as an alternative to the conventional TiO$_2$ catalysts, in order to facilitate the separation of products after the reaction. However, despite the large number of research in this field, the mechanism governing the photocatalytic activity of the mixed TiO$_2$/SiO$_2$ oxides is not clear. Titania-Silica nanocomposite xerogels were prepared by sol-gel method. This work has been used to describe the synthesis and the photocatalytic properties of TiO$_2$/SiO$_2$ nanocomposite xerogel. The nanocomposite xerogels were prepared by keeping the molar ratio of TEOS:TTIP:MOH:DIW at 1:1:6:14 respectively and the catalysts used were HCl and NH$_4$OH. After the preparation xerogels were characterized by FTIR, XRD, UV and LLS. All these techniques show the amorphous nature of Titania-silica xerogel.

Keywords: Photocatalysis; TiO$_2$, SiO$_2$, mixed oxide; LLS; FTIR; UV; XRD

Introduction

TiO$_2$ is well recognized as a valuable material with application as a white pigment in paints, as filler in paper, textile and in rubber/plastics [1]. Due to low cost, non-toxicity, stability and other best characteristics TiO$_2$ attracts a great attention. TiO$_2$ has wide applications in various fields like antirefection opticals, coatings, waste water purifications, catalyst supporting, ceramics sensor element, as a photocatalyst, in electric devices like (in lithium based battery), as a base in high quality paints, paper, plastics [2]. Titania has excellent biocompatibility with respect to bones implants and applications in electrochromic devices [3]. Titania shows good photocatalytic applications due to which it gained tremendous demands and green energy and environmental protection. Many other oxides like iron oxides, zinc oxides etc. also shows the similar behavior due to photocatalytic activity of titanium dioxide it play a wide role in different fields like air, waste water purification, good UV blocking properties weakening of the organic fibers [4]. Silica doped in to the titania matrix increase the photocatalytic activity because the silica doping decrease particle size and also increase the specific surface area and thermal stability of titania particle towards anatase to rutile phase conversion [5]. SiO$_2$-TiO$_2$ materials are used in different fields like as catalyst supporting materials, acidic catalyst for many reactions, selective reduction, as an anti-reflective materials for coatings or sensing nanoimprints photonic crystals [6-9]. Dielectric mirrors and low loss waveguides solids of low thermal expansion coefficient, bioactive solids self-cleaning coatings solids of controlled acidity and photocatalysts [10-14]. The TiO$_2$-SiO$_2$ mixed oxides catalytic activity was studied and observed that the TiO$_2$-SiO$_2$ have better photocatalytic activity as compared to TiO$_2$ and SiO$_2$ which was confirmed through LLS and UV results.

Experimental

Sample preparation

The xerogels can be synthesized by Sol-Gel process in which metal alkoxide is used as a precursor source that undergoes catalyzed hydrolysis and condensation to get nano scale materials of that metal [15]. TTIP was used as a precursor and TEOS was added as an organic solvent. In this synthesis HCl and NaOH were used as catalysts.

TiO$_2$-SiO$_2$ mixed oxide

TiO$_2$-SiO$_2$ xerogels was synthesized by Sol-Gel process. TEOS (Tetraethylorthosilicate) and TTIP (Titanium tetra isopropoxide) were used as precursor. It was observed that the TTIP hydrolysis rate is much faster than TEOS [16]. The synthesis consists of two steps. The first step is the synthesis of SiO$_2$ sol, in which 1:6 TEOS and Methanol were mixed together i.e. 7 ml of TEOS and 43 ml of methanol. TEOS were added drop wise slowly to the methanol with continuous stirring. Then 0.05 mol L$^{-1}$ HCl was added drop wise to the sol in order to adjust the acidic pH at 2. Then the sol was allowed to stir for about 2 h in order to get the homogeneous sol. In the second step TiO$_2$ Sol synthesis, 1:14 solution of TTIP and DIW were mixed with continuous stirring until the homogeneous sol of TiO$_2$ were obtained. Then in 1:1 of TiO$_2$ and SiO$_2$ sols were mixed with continuous stirring in order to get homogeneous sol and then 0.05 mol L$^{-1}$ NH$_4$OH solution was added to this mixture drop wise for the adjustment of pH. The pH of the homogeneous mixtures (sol) was observed by the pH-meter continuously. At last the mixtures (sol) were allowed to stir for some time to get the homogeneous mixture (sol).

Gel and xerogel preparation

In this method 3ml of TEOS, the silica precursor was taken in the reaction beaker and 6ml of ethanol was added dropwise with continue stirring. Further 3 ml of water, 4 ml of acetic acid and 3 ml of TTIP (Titania precursor) were added with continuous stirring for 15 min at room temperature. The prepared sol changed into a gel which was placed in the oven at 65 for 1 h which resulted in the conversion of gel in to xerogel.

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Characterization

Structure, quality, photocatalytic studies and morphological characteristics of Titania-silica gel were studied by different characterization techniques like, LLS, FTIR, UV and XRD etc.

Results and Discussions

FTIR result

The FTIR spectra of the synthesized nanocomposite (TiO$_2$/SiO$_2$) were recorded by Perkin Elmer series 100 FTIR spectrometer with a 5 cm$^{-1}$ resolution. FTIR spectrum was recorded at 4000-450 cm$^{-1}$. The absorption at 1074 cm$^{-1}$ (Figure 1) is the characteristics peak for Si-O-Si. The peak observed at 801 cm$^{-1}$ is due to Si-O-Si symmetric stretching. The broad absorption at 1633 cm$^{-1}$ match with OH bending vibrations and is attributed to chemisorbed water. The well defined peak at 3441 cm$^{-1}$ shows OH stretching vibrations. The peak observed at 923 cm$^{-1}$ corresponded Si-O-Ti vibrations. The band observed at 450-610 cm$^{-1}$ is due to Ti-O Stretching.

Many characteristic FTIR peaks were observed in Figure 2. The bands observed at 3391 cm$^{-1}$ and 1557cm$^{-1}$ were due to the OH bending and stretching vibrations respectively. The peak observed at 1165 cm$^{-1}$ correspond to Si-O-Si antisymmetric stretching vibration. The peak obtained at 923 cm$^{-1}$ correspond to Si-O-Ti vibration. The peak observed at 801 cm$^{-1}$ is due to the Si-O-Si symmetric stretching vibration. The absorption at 1410 cm$^{-1}$ is due to C-H interaction of Si-R structure unit (Figure 3).

XRD measurement

The XRD pattern of synthesized nanocomposites (TiO$_2$/SiO$_2$) was collected in the range of 10-60 2θ (degree) shown in Figure 4. The XRD patterns of the synthesized materials indicate that the TiO$_2$-SiO$_2$ nanocomposites are essentially non-crystalline and have amorphous structure which can be accomplished from the broad characteristic diffraction peak between 2θ ~ 20° and 30 θ°.

The Figures 5 and 6 also show the same XRD results of the synthesized TiO$_2$/SiO$_2$ nanocomposites. These samples were also synthesized by sol-gel method with different concentrations of precursors and solvents used. The hump in XRD pattern indicate that the TiO$_2$-SiO$_2$ nanocomposites are essentially non-crystalline and have amorphous structure which can be accomplished from the broad characteristic diffraction peak between 2θ ~ 20° and 30 θ° while the Figure 2 shows very low crystallinity at 2θ and 25 θ°.

Catalytic activity

To check the catalytic activity of TiO$_2$-SiO$_2$ the p-nitrophenol was reduced by NaBH$_4$ to p-aminophenol in aqueous medium. The procedure adopted was used as a given amount of mixed oxides was added to 1 ml of p-nitrophenol (0.08 mmol L$^{-1}$) for initiation of reduction. For mixture preparation 1 ml of aqueous solution of NaBH$_4$ (1.5 mmol L$^{-1}$) was added to reaction chamber. The p-nitrophenol reduction was determined by studying the absorbance at 440 nm with respect to time. This reaction was selected due to simplicity and formation of single product. By using time dependence uv-vis spectra the reduction process was checked. The appearance of a new peak at 300 nm was observed which confirm the p-nitrophenol reduction to p-aminophenol which is shown in Figure 6.

LLS result (hydrodynamic radii)

The LLS results (hydrodynamic radii) of the three samples are given in Table 1 below. All the three samples have different hydrodynamic radii. The two samples TSX$_2$ and TSX$_3$ have same precursors and solvent but used at different ratio. The sample TSX$_1$ have same precursors but different solvents and precursor ratio were used. The LLS results (hydrodynamic radii) are different because the LLS result
Figure 4: XRD results of TiO$_2$-SiO$_2$ nanocomposite xerogels of sample TSX$_2$.

Figure 5: XRD results of TiO$_2$-SiO$_2$ nanocomposite xerogels of sample TSX$_3$.

Figure 6: Catalytic reduction of P-nitrophenol to P-aminophenol.

Figure 7: SEM results of TiO$_2$-SiO$_2$.

Figure 8: EDX results of TiO$_2$-SiO$_2$.

The XRD results confirmed that the synthesized TiO$_2$-SiO$_2$ mixed oxide xerogel is non-crystalline and having amorphous nature. In the TiO$_2$/SiO$_2$ nanocomposites, materials, in the amorphous SiO$_2$ matrix TiO$_2$ nanocrystals are present in highly dispersed form. The amorphous SiO$_2$ and Ti–O–Si bond formation and TiO$_2$-SiO$_2$ mixed oxides give rise effectively stability of TiO$_2$ anatase form. It also bound crystallites growth, significantly increase surface area. So, the increase in surface area caused the improvement in photocatalytic activity of TiO$_2$-SiO$_2$ mixed oxides nanocomposites. FTIR spectra show the presence of Ti-O-Si crosslinks and revealed interactions between TiO$_2$ and SiO$_2$ at a molecular scale. Ti-O-Si bonds and interactions may...
enhance surface properties, catalytic and photoactivity. The TiO$_2$-SiO$_2$ mixed oxides catalytic activity was also studied and observed that the TiO$_2$-SiO$_2$ have better photocatalytic activity as compared to TiO$_2$ and SiO$_2$ which was confirmed through LLS and UV results.

Acknowledgements

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References


Table 1: LLS Results (hydrodynamic radii) of the three samples prepared by sol gel process.

<table>
<thead>
<tr>
<th>S No</th>
<th>Sample</th>
<th>Hydrodynamic radius (nm)</th>
</tr>
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<tbody>
<tr>
<td>1</td>
<td>TSX$_1$</td>
<td>170.3</td>
</tr>
<tr>
<td>2</td>
<td>TSX$_2$</td>
<td>138.49</td>
</tr>
<tr>
<td>3</td>
<td>TSX$_3$</td>
<td>52.6</td>
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</tbody>
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