

**DEGRADATION OF ERIOCHROME BLACK-T USING SnO<sub>2</sub> NANOPARTICLES****A. FATHIMA BEEVI<sup>a</sup>, G. SREEKALA<sup>b</sup> AND B. BEENA<sup>c1</sup>**<sup>abc</sup>Nanoscience Research Lab, Department of Chemistry, K.S.M.D.B. College, Sasthamcotta, Kollam, India**ABSTRACT**

SnO<sub>2</sub> nanoparticles were synthesized by sonochemical method using Tin Tetrachloride and Ammonium Hydroxide as precursors. The prepared SnO<sub>2</sub> nanoparticles were characterized by PXRD, FTIR, UV-Vis and SEM- EDX. The dye degradation activity of the prepared sample was investigated by degradation of Eriochrome Black-T in aqueous medium under natural sunlight using UV-Vis spectrophotometer. The prepared SnO<sub>2</sub> nanoparticles are efficient for removing water soluble dyes.

**KEYWORDS:** SnO<sub>2</sub> Nanoparticles; Sonochemical Method; Degradation; Eriochrome Black-T

In recent decades, the semiconductor photocatalysts with high performances for water contaminant degradation have attracted great interest to solve environmental issues. Among the various nanocrystalline materials, tin oxide is a versatile oxide having wider applications. Because of its strong physical and chemical interaction with adsorbed species, low operating temperature and strong thermal stability SnO<sub>2</sub> is widely used as a common 'n' type direct wide band gap semiconductor and photocatalyst. There are a variety of techniques that have been developed to prepare nanoparticles. They are sol-gel, solvothermal, microwave, chemical co-precipitation, hydrothermal and sonochemical methods. In the present work, the sonochemical method was employed to prepare tin oxide nanoparticles.

Eriochrome black-T is an azo dye used in textile industry. The complex chemical structure of Eriochrome black-T offers a greater resistance to photodegradation as well as certain chemical reagents, impairing the removal or reduction of its colour during the waste water treatment (Barka et al;2011).

**MATERIALS AND METHODS****Synthesis of SnO<sub>2</sub> nanoparticles**

Typically, 50 mL of 0.4M Ammonium hydroxide solution was first added drop wise to 50 mL of 0.1M tin tetrachloride [(SnCl<sub>4</sub>.5H<sub>2</sub>O), 97%, Sigma- Aldrich] solution and kept under continuous stirring for 30 minutes. Then the solution was sonicated for 1 hour (230 volts, 50Hzs). Thus obtained precipitate was washed, dried and calcined at 300°C for 3hours.

**Characterisation of SnO<sub>2</sub> Nanoparticles**

The crystalline quality and grain size of the samples were evaluated using Powder X-ray diffraction (PXRD) measurements. The morphology of sample was examined using Scanning Electron Microscopy (SEM).

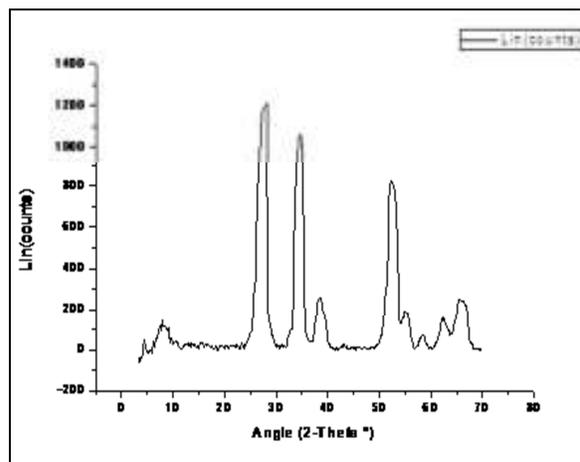
Chemical nature of the synthesized SnO<sub>2</sub> was analysed by FT-IR spectrum. The band gap and optical properties of the nanoparticles were studied using UV-Visible spectrum.

**Preparation of Eriochrome Black-T Solution**

The accurately weighed 0.01gm quantity of the Eriochrome black-T dye was dissolved in 100mL of distilled water.

**Dye Degradation using Nano SnO<sub>2</sub>**

The photodegradation activity of nano SnO<sub>2</sub> is carried out by mixing 0.015gm of the sample and dye solution at 10minute intervals and recording the UV-Visible absorption spectra.

**RESULTS AND DISCUSSION**

**Figure 1: PXRD spectrum of SnO<sub>2</sub>nanoparticles**

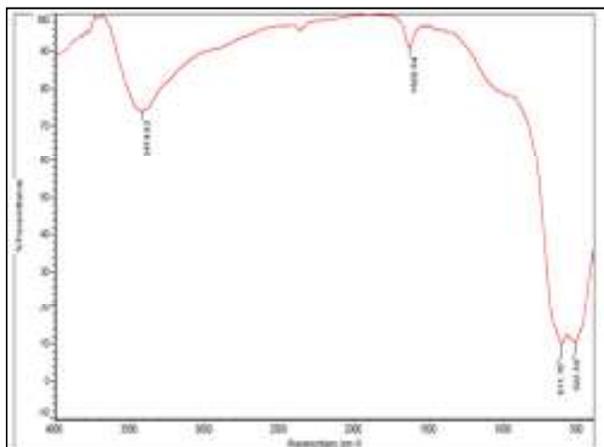


Figure 2: FT-IR spectrum of SnO<sub>2</sub> nanoparticles

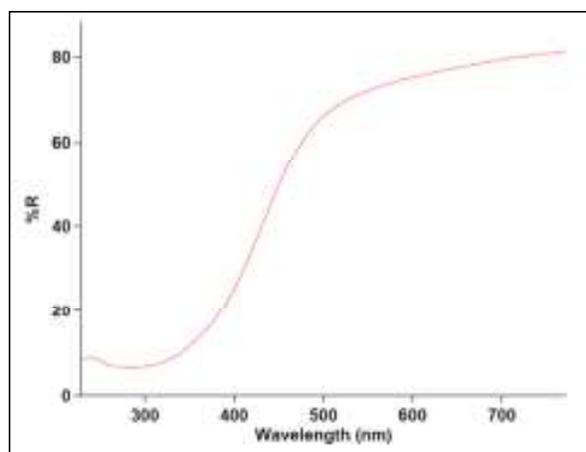


Figure 3: Reflectance spectrum of SnO<sub>2</sub> nanoparticles

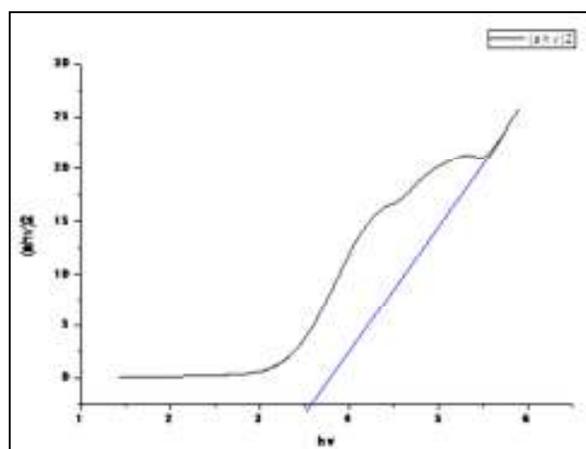


Figure 4: Tauc's plot of SnO<sub>2</sub> nanoparticles

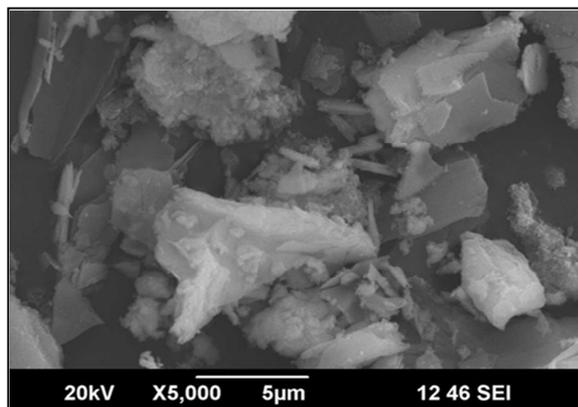


Figure 5: SEM image of SnO<sub>2</sub> nanoparticles

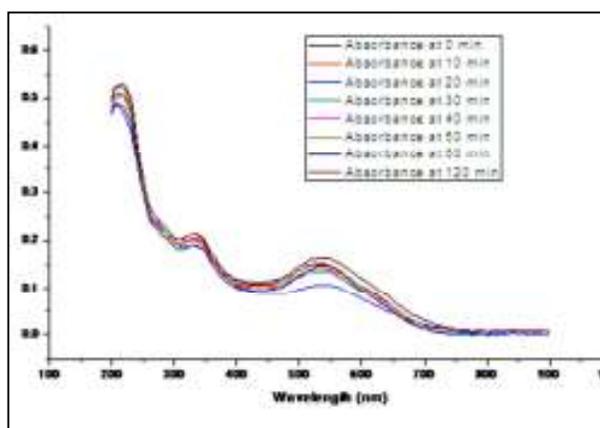


Figure 6: UV-Visible Absorption spectra of EBT at various time intervals

The X-ray diffraction is used to identify crystalline structure and particle size of the samples. X-ray diffraction patterns of our sample of SnO<sub>2</sub> are shown in Figure.1. The diffraction peaks readily indexed to the tetragonal rutile phase of SnO<sub>2</sub> (JCPDS No.77-0450). From the crystallite size calculation, using Scherrer equation,  $t=0.9\lambda/\beta\cos\theta$ , the average size of prepared sample of SnO<sub>2</sub> is between 10-20 nm.

The FTIR transmission spectrum of the synthesised SnO<sub>2</sub> is shown in Figure.2. The peaks at 3414.53cm<sup>-1</sup> and 1629.54cm<sup>-1</sup> were due to stretching and bending vibrations of water molecules or hydroxyl groups adsorbed at the surface of the tin oxide.(S.Y.Ho,et al 2009.) The bands at 611.18 cm<sup>-1</sup> and 522.58cm<sup>-1</sup> refers to Sn-O stretching modes of Sn-O-Sn.(Farukh et al. 2010)

SEM image shows the flakes like morphology of the prepared sample in Figure 5.

From UV-Vis Spectra and Tauc's relation,  $\alpha hv = A (hv - E_g)^n$ , the band gap obtained is 3.6eV [Figure.3, Figure.4 and Figure.6]. The SnO<sub>2</sub> prepared by sonication

method is a good adsorbent for the removal of Eriochrome Black-T and points out the potential application of the material for waste water treatment .

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