

Analysis of Pure and Ag-Doped TiO₂ Thin Films Deposited by Field Assisted Spray Pyrolysis

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Original Research Article

ABSTRACT

This paper reports the structural and optical properties of Undoped TiO₂ and 5% Ag-doped TiO₂ nanoparticles under visible- light irradiation. These films were synthesised using Electrohydrodynamic Spray Pyrolysis Technique, deposited at 250°C and later annealed at 400°C few minutes. The films were characterised by Surface Profilometry, Ultraviolet – Visible light Spectroscopy (UV-Vis Spectroscopy), Optical Microscope and X-ray diffraction (XRD). The result from surface profilometry showed that the thickness of all the samples is thin films. XRD results showed that all the films have crystalline nature of TiO₂ and annealed films result in an increase of crystallinity. The crystal phase observed was trigonal for both TiO₂ and 5% Ag doped TiO₂. The unannealed Undoped TiO₂ exhibited low absorbance within the visible spectrum but increases as we dope with 5% Ag-doped TiO₂ and effect of annealing increases the absorbance. Extinction coefficient increases within the visible region as we doped and thermal annealing also increases its effect within the visible region. The study of the microstructural properties using Optical Microscopy showed an increase in the hardness of the film for annealed films and they have micro-grains that are closely packed compared to unannealed samples. The direct band-gap for unannealed films range from 4.70 eV- 3.10 eV while annealed films was between 3.20 eV and 2.80 eV. Doped Annealed films showed reduced band-gap.

Keywords: Ag doped TiO₂; field assisted spray pyrolysis; structural properties; optical properties and wide band gap semiconductor.

1. INTRODUCTION

Titanium dioxide (TiO₂) also known as titania is the most promising electron acceptor and a very useful wide band-gap semiconducting transition metal oxide which has attracted much interest among researchers due to its unique characteristics such as non-toxicity, affordability, easy handling, resistance to photochemical and chemical erosion (Hoffmann, *et al.*, 1995). It is a wide band-gap semiconductor with energy of 3.0–3.4 eV and is widely used in applications such as hydrogen production (Zaleska, 2008), gas sensor (Sayilkan, *et al.*, 2005), (Gomez & Meghraoui, 2003), photo catalytic activities (Yogeswaran & Chen, 2008), dye-sensitised

solar cells and photo electrochemical cells because of its relative high efficiency and high stability. TiO₂ exists in both crystalline and amorphous forms. In its existence in three crystalline polymorphs, namely, anatase, rutile and brookite, anatase and rutile have a tetragonal structure, whereas brookite has an orthorhombic structure (Mahshid, *et al.*, 2007).

There are different routes that can be used to synthesise titanium dioxide. These include a conventional solid state route (Dong, 2004), (Guimarães *et al.*, 2003), precipitation (Joo, *et al.*, 2005), (Lee & Yang, 2005), Sonochemical (Guimarães *et al.*, 2003), hydrothermal (Hidalgo, *et al.*, 2007), (Deshpandea, 2011), microwave

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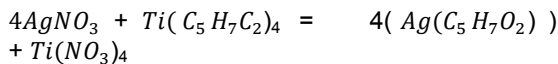
hydrothermal (Murugan, *et al.*, 2006) and sol-gel methods (Pookmanee & Phanichphant, 2009), (Tiberiu, *et al.*, 2014). Titanium dioxide (TiO₂) also exhibits useful electrical, optical, and photocatalytic behaviour due to the powerful oxidation and reduction properties that arise in the absorption of photon energy (Kim & Kwak, 2007).

Due to TiO₂ wide band-gap which ranges from 3.0eV- 3.4eV, between the valence and conduction bands, there is a significant limitation with regards to the photocatalytic applications. Hence several attempts have been made by researchers to improve its absorbance within the visible region thus enhancing its photocatalytic activities through the process of doping (Ambrus, *et al.*, 2008).

In this work, we carried out a comparative analysis of the different deposited samples to identify the effect of concentration of dopant and the effect of thermal annealing on the samples using Electrohydrodynamic Spray Pyrolysis Technique. The films deposited were characterised by Surface Profilometry, Ultraviolet – Visible light Spectroscopy (UV-Vis Spectroscopy), Optical Microscopy and X-ray diffraction (XRD).

2. EXPERIMENTAL

TiO₂ was synthesised using 0.2mol of titanium acetyl acetate [Ti(C₅H₇O₂)₄] and 20mol of propanol [C₃H₈O] as the precursor while 0.68g of silver nitrate [AgNO₃], mixed with 20mol of propanol in a conical flask, placed on a magnetic stirrer and was stirred with the magnetic stirring bar, until the AgNO₃ was properly dissolved served as the precursor for Silver(Ag). The precursor delivery system containing string pump was formatted using a laptop to get accurate atomization or deposition. The flow rate was set at 4200microlitres/ hour, the volume was set at 400microlitres/hour and the time of deposition was set at 10mins. The Electrohydrodynamic Spray Pyrolysis Technique was set at the voltage of 3.5KV and the distance between the nozzle and the substrate (pro cleaned glass) was 5.95mm. From the precursors we obtained



5% Ag doped TiO₂ was deposited at 250°C and was duplicated to get two samples. Undoped TiO₂ was also deposited at 250°C and also duplicated to two samples. Hence we have a total of four(4) samples. One 5% Ag doped TiO₂

and one undoped TiO₂ were later annealed at 400°C in few minutes using master chef annealing machine with Model No: P70B17L-Ds(7014336), in order to expel the water of crystallisation from the samples.

The thickness of the synthesised samples (both doped and undoped) was measured using surface profilometry with a VeecoDektak 150 profiler machine. Using UV-Vis spectroscopy, the absorption spectra of the deposited undoped TiO₂ and Ag- doped TiO₂ thin films were recorded within the wavelength range of (UV) region of (230 nm to 390 nm), visible light (VIS) region of (400 nm to 700 nm) and near infrared (NIR) region of (714 nm to 1080 nm). The reflectance values of the work were obtained using a relation between the absorbance (A), transmittance (T) and reflectance (R), according to the law of conservation of energy given by

$$A + T + R = 1 \quad (1.1)$$

From equation 1.1, reflectance (R) was given as

$$R = 1 - T - A \quad (1.2)$$

The values of refractive index of were obtained using the equation of refractive index according to (Ali *et al.*, 2015),

$$n = (1 + \sqrt{R}) / (1 - \sqrt{R}) \quad (1.3)$$

Where R is the reflectance and n is the refractive index.

The values of extinction coefficient of this work were calculated using the equation of extinction coefficient according to (Mahrov, *et al.*, 2004)

$$K = \alpha\lambda/4\pi \quad (1.4)$$

Where k is called the extinction coefficient (k), α is the absorption coefficient, λ is the wavelength, π is equal to 3.142.

The microstructural properties of both doped and undoped TiO₂ were determined using Light Microscope; Celestron LCD digital microscope. (Model no: 44348), under the magnification of 800. The XRD pattern was recorded on a Rigaku Ultima IV X-ray diffractometer equipped with a graphite- monochromated Cu Kα radiation source (40kv, 30mA) and wave length 1.54Å. A diffractogram was collected in the 2θ range between 3° and 90° with a step size of 0.01° and a scan speed of 1°/min. the XRD Pattern was processed using JCPDS card numbers. We used

Debye Scherer equation to calculate for the crystallite size.

$$D = k\lambda / \beta \cos\theta \quad (1.5)$$

Where D is the crystallite size, K= shape factor, which is constant (0.89), λ = wavelength (1.54Å), β = Full Width Half Maximum (FWHM), θ = 2 Theta, which to obtain θ , we divided by 2.

3. RESULTS AND DISCUSSION

The thickness of the synthesised samples (both doped and undoped) was measured using surface profilometry with a VeecoDektak 150 profiler machine; the result showed that all the samples are thin films.

3.1 Optical Properties of the Deposited Films

The absorbance spectra of the deposited films were shown in Fig. 1a and 1b. It is found that absorbance spectra of 5% Ag doped TiO₂ has high absorbance within the visible region than the undoped TiO₂. Comparing the annealed and unannealed films, it also showed that annealed films have the best absorbance within the visible region than unannealed films.

Figs. 2a and 2b below showed the extinction coefficient of unannealed samples and annealed samples. It showed that doping increases the extinction coefficient of the film within the visible region.

using Tauc's relation

$$\alpha h\nu = A(h\nu - E_g)^n \quad (1.6)$$

where $h\nu$ is the photon energy, h is Planck's constant, α is the absorption Coefficient, A is the constant, for direct transitions $n=2$, and E_g is the optical band gap, which is given as

$$E_g = h\nu = hc / \lambda \quad (1.7)$$

C is the velocity of light = $3 \times 10^8 \text{ ms}^{-1}$ λ is the wavelength in meters (m). ν is the transition frequency, h is the Planck's constant = $6.62 \times 10^{-34} \text{ Js}$. Since $1\text{J} = 1.602 \times 10^{-19} \text{ eV}$

We substituted these variables into equation 1.7 and obtained

$$\begin{aligned} h\nu(\text{eV}) &= \frac{hc}{\lambda_0} = \left(\frac{6.63 \times 10^{-34} \text{ Js} \times 3 \times 10^8 \text{ ms}^{-1}}{\lambda \times 1.602 \times 10^{-19} \text{ c}} \right) \\ &= \left(\frac{6.63 \times 10^{-34} \text{ Js}^{-1} \times 3 \times 10^8 \text{ ms}^{-1}}{\lambda(\text{nm}) \times 10^{-9} \times 1.602 \times 10^{-19} \text{ c}} \right) \\ h\nu(\text{eV}) &= 1.241 / \lambda(\mu\text{m}) \end{aligned} \quad (1.8)$$

From equation 1.8 we calculated the photon energies in (eV) for various wavelengths λ in (μm). A graph of $(\alpha h\nu)^2$ versus $h\nu$ for both unannealed and annealed samples were plotted and the direct band gap was extracted by extrapolating the straight portion of the graph on $h\nu$ axis at $(\alpha h\nu)^2 = 0$ which gives the band gap of both unannealed and annealed films. For unannealed, the undoped TiO₂ has the band gap of 4.70eV while 5% -doped Ag has band gap of 3.10eV which shows that doping brings about reduction of the energy band gap from the graph below.

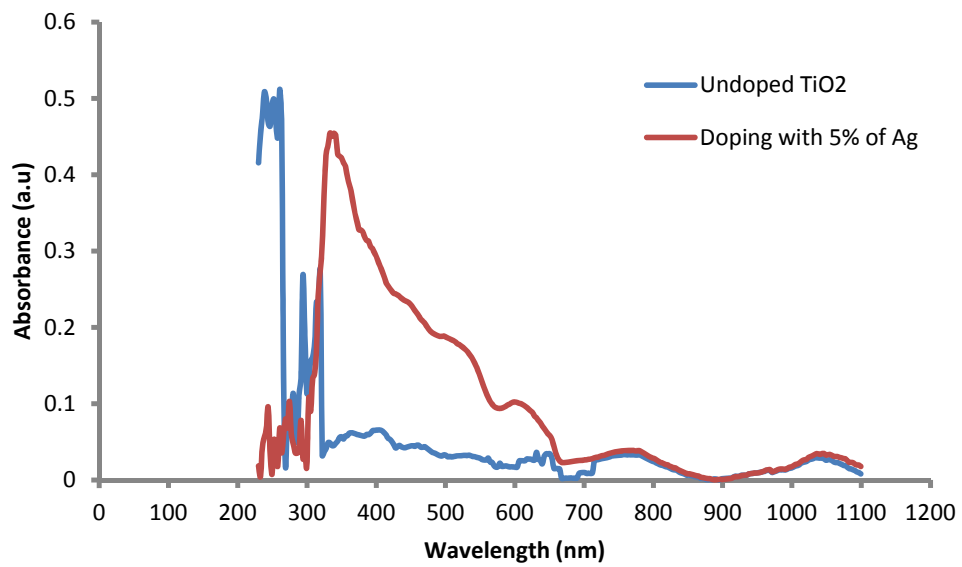


Fig. 1a. Absorbance spectra for the undoped and doped unannealed samples

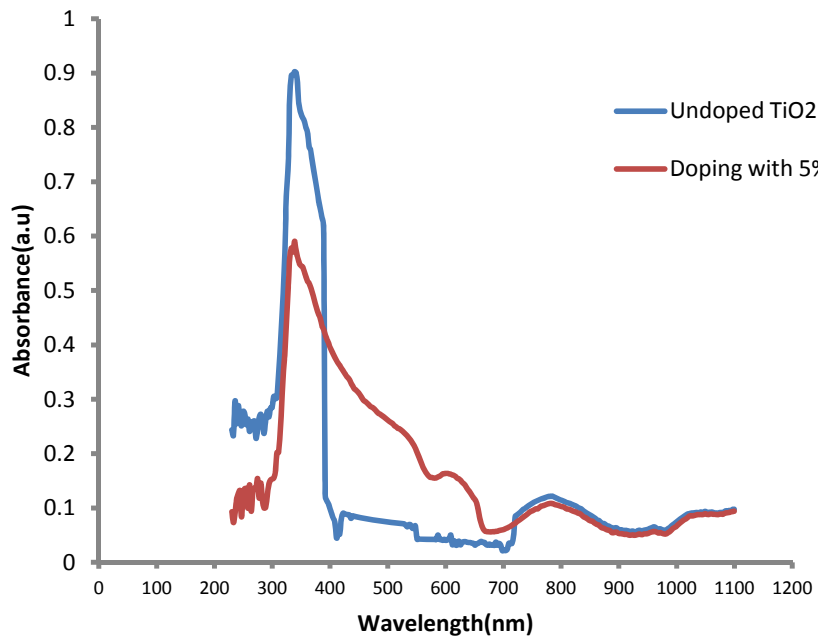


Fig. 1b. Absorbance spectra for the undoped and doped annealed samples

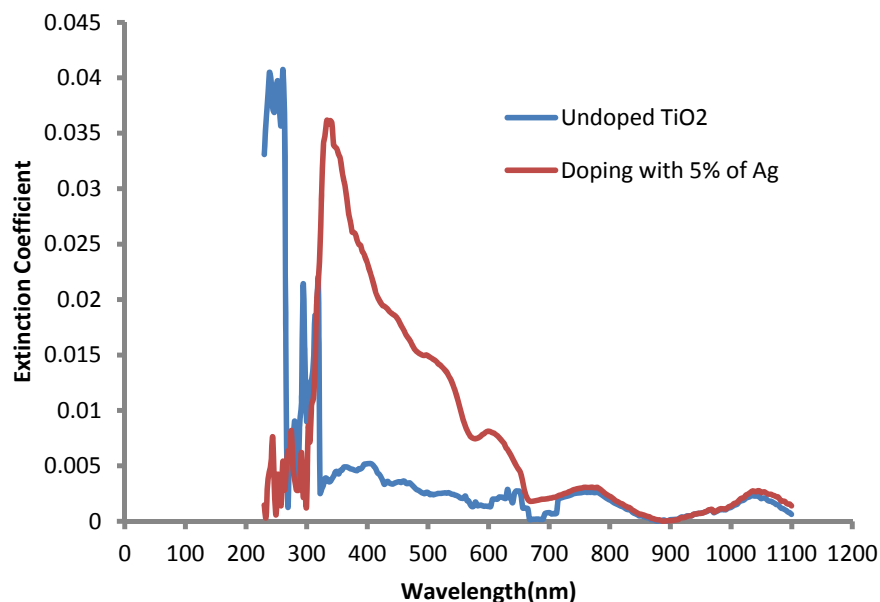


Fig. 2a. Dependence of extinction coefficient on the wavelength for undoped and doped unannealed sample

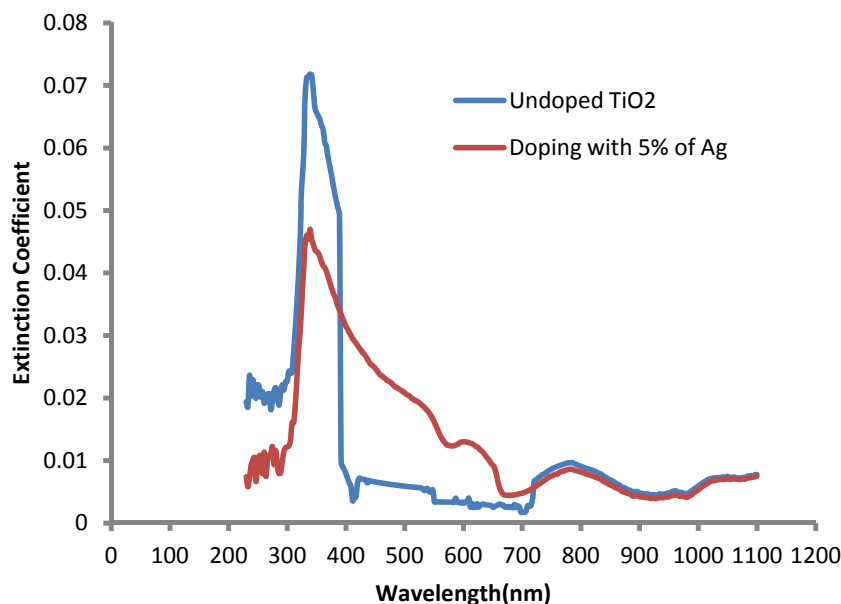


Fig. 2b. Dependence of extinction coefficient on the wavelength for undoped and doped annealed samples

For the annealed films, the undoped TiO₂ has the band gap of 3.20eV which is in agreement with undoped TiO₂ result obtained by (Hassan *et al.*, 2008) while the 5%-doped Ag has 2.80eV similar to result obtained by (Ayieko *et al.*, 2012). The annealed films have narrowed energy band gap than the unannealed films.

3.2 The Micrograph Analysis

Figs. 4a- 4b show the micrograph of unannealed and annealed samples of TiO₂ using Celestron LCD digital microscope, model #44348 with magnification of 800. Fig. 4a shows the micrograph of undoped unannealed TiO₂ sample which reviewed rough surface morphologies and larger pores on the film, which may be due to the aggregation of tiny crystals and the time of deposition. Using imageJ app, we obtained an

average particle size of 6.00 while Fig. 4b, the micrograph of undoped annealed TiO_2 sample reviewed an increase in the number of orderly stacked grains. There was reduction in the size

of pores and there was increase in the hardness of the film compared to unannealed sample. The average particle size of 8.00 was obtained using imageJ app.

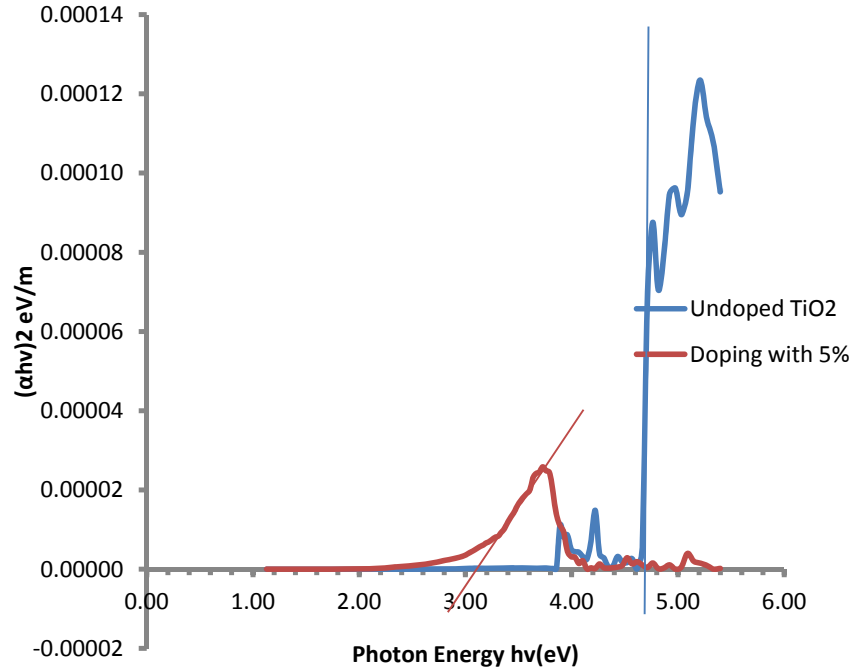


Fig. 3a. Direct band gap for unannealed undoped TiO_2 and 5%-doped Ag films

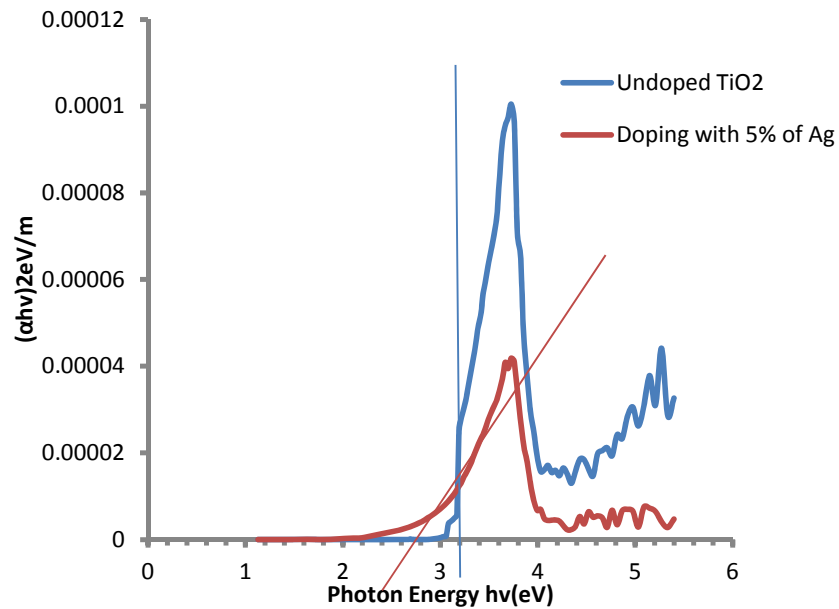


Fig. 3b. Direct band gap for annealed undoped TiO_2 and 5%-doped Ag films

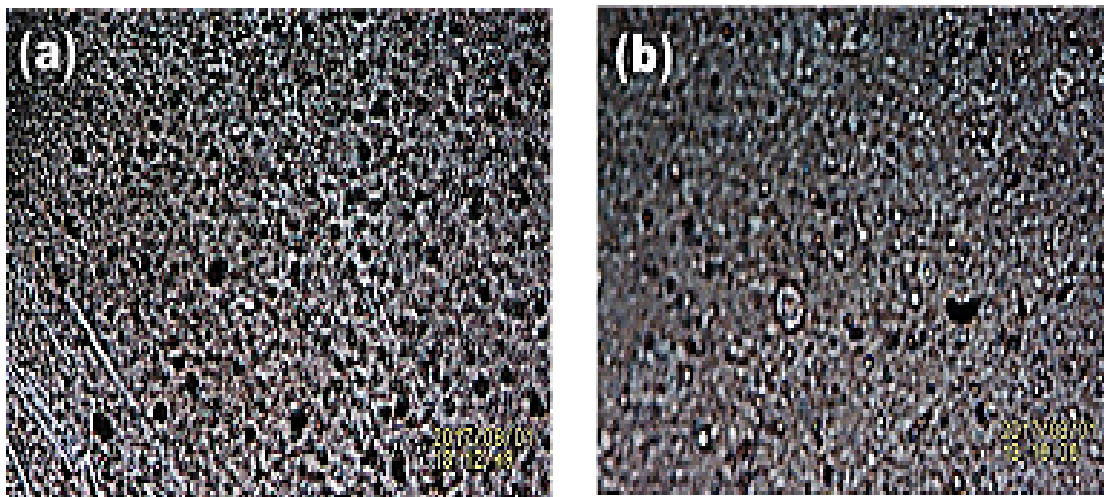


Fig. 4. Micrograph of Undoped TiO_2 , (a) unannealed sample (b) annealed sample

Figs. 5a- 5b show the micrograph of unannealed and annealed samples of 5% of Ag/TiO_2 using Celestron LCD digital microscope, model #44348 with magnification of 800. Fig. 5b shows rough surface morphology of the film; larger pores and larger grains were evident. Using imageJ app, we obtained an average particle size of 7.00 while Fig. 5b shows the micrograph of annealed 5% Ag/TiO_2 sample. Close observation reviewed that micro grains are packed closely compared to unannealed sample. There was increase in the hardness of the film and using imageJ app, we obtained an average particle size of 9.00.

3.3 The X-ray Diffraction

Fig. 6 show the x – ray diffraction patterns for both Annealed and Unannealed undoped TiO_2 samples with peaks at $2\theta = 25.30^\circ, 27.56^\circ, 37.99^\circ, 48.25^\circ, 54.22^\circ, 62.65^\circ$ and 74.80° which correspond to miller indices [101], [110], [103], [200], [105], [213] and [107] respectively. These results show that the samples are of trigonal crystalline phase of TiO_2 which correspond to the Crystallography Open Database (COD) entry file number of 96 – 153 – 7225. Using Debye-Scherrer equation the average crystallite

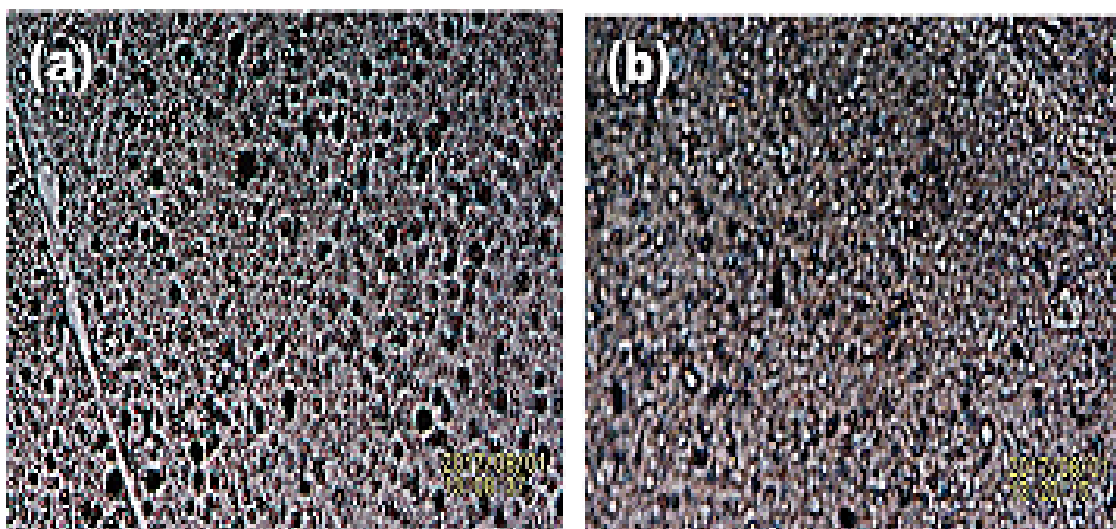


Fig. 5. Micrograph of 5% $\text{Ag}:\text{TiO}_2$, (a) unannealed sample (b) annealed sample

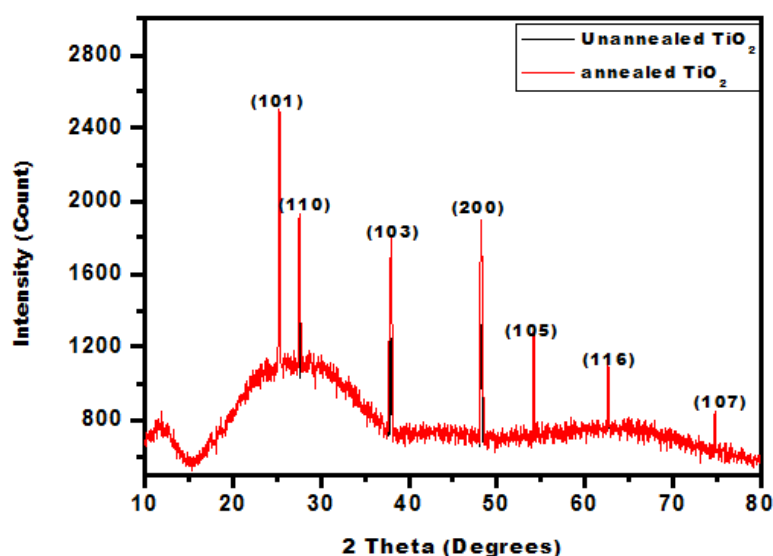


Fig. 6. X- ray Diffraction Pattern for Undoped Unannealed TiO_2 and Undoped Annealed TiO_2

was 8.78nm and 8.76nm respectively. Comparing the two, the annealed TiO_2 pattern had well defined or high intensity of peaks than the undoped unannealed TiO_2 sample.

Fig. 7 shows the x – ray diffraction patterns for 5% Ag of both annealed and unannealed samples. They have diffraction peaks at $2\theta = 25.33^\circ, 27.49^\circ, 37.98^\circ, 48.31^\circ, 54.28^\circ, 62.47^\circ$ and 74.89° which correspond to miller indices [101], [110], [103], [200], [105], [213] and [107] respectively. The results show that the Ag doped TiO_2 are crystalline in nature. The x-ray spectra

were matched with the standard data using Match3 software. The result corresponds to the Crystallography Open Database (COD) entry file number of 96 – 153 – 7225. There was no significant shift in the peak position of diffraction patterns of both annealed and unannealed 5% doped with Ag, except the changes in the intensity of these peaks. Annealed sample had well defined or high intensity of peaks than the unannealed sample due to annealing of the sample. Using Debye-Scherrer equation the average crystallite was 8.80nm and 8.76nm respectively.

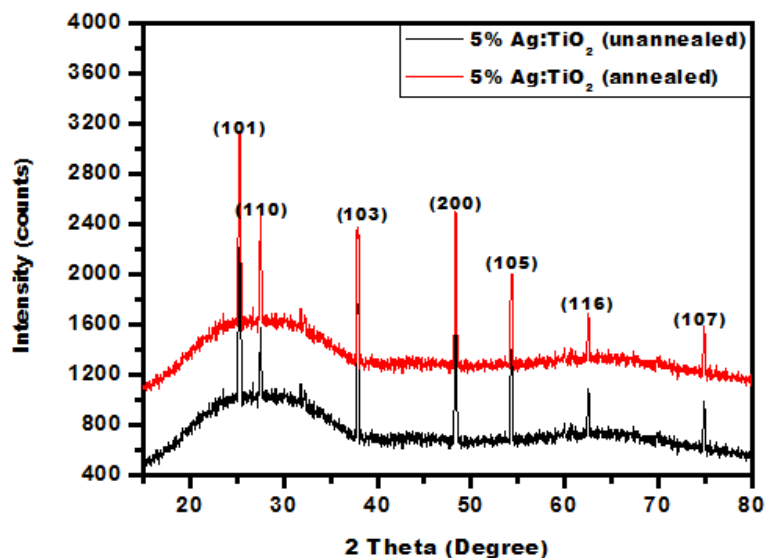


Fig. 7. X- ray diffraction pattern for both 5% of Ag/ TiO_2 annealed and unannealed samples

4. CONCLUSION

In this paper we have reported the structural, optical properties of TiO₂ and Ag-doped TiO₂ synthesised using Electrohydrodynamic spray pyrolysis equipment. The XRD pattern showed crystalline nature of TiO₂. There was increase in the crystallinity of the films in the case of annealed. UV- vis spectroscopy revealed the shifting of absorption edge of silver-doped TiO₂ to the visible region compared to that of Undoped TiO₂ and the annealed films showed higher absorbance than unannealed films. The Extinction coefficient showed that as the dopants are been added to the parent material, the extinction coefficient values of the films increases within the visible region. The direct band gap for annealed showed more decrease in energy band gap than the unannealed doped samples. Using Optical Microscopy showed increase in the hardness of the film for annealed films and they have micro-grains that are closely packed compared to unannealed samples. For the unannealed, the direct band gap ranges from 4.70eV – 3.10eV and the annealed the direct band gap ranges from 3.20eV – 2.80eV. The undoped TiO₂ nanoparticles showed poor photocatalytic activity, while doping of silver ions improves the efficiency under the visible –light irradiation.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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