



Physical Investigations of Titanium Dioxide Nanorods Film Prepared by Hydrothermal Technique

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Abstract

In this work, well-oriented and homogeneous titanium dioxide (TiO₂) nanorods (NRs) was synthesized by hydrothermal method. In this method, vertically aligned arrays of TiO₂ were built on the conductive Fluorine–Tin–Oxide (FTO) glass substrate. Nanoparticles (NRs) of TiO₂ showed a tetragonal shape with a square top face, according to the image of the field emission scanning electron microscope (FE-SEM). The TiO₂ NRs are polycrystalline, having two phases: rutile and anatase, according to X-ray diffraction (XRD) analysis. The optical properties of a TiO₂ nanorods arrays were examined, including transmittance, absorption coefficient, and energy bandgap. An optical energy band gap of 3.18 eV was obtained. According to the photoluminescence emission measurement, the energy bandgap was 3.3 eV. For further study of the optical properties of the TiO₂ Nanorods films, reflectance spectrum was used as a function of wavelength to estimate the value of the energy bandgap and its value was 3.45 eV. By comparing the values obtained from the three methods, it is found that they are closely alike, which confirms the formation of the TiO₂ nanostructure.

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1. Introduction

Titanium dioxide has been extensively researched over the last few decades for its photovoltaic applications, sensors, photocatalysis, dye-sensitized solar cells, integrated circuits, and environmental devices [1,2]. Chemical inertness, nontoxicity, efficiency, physical stability, a strong refractive index, a direct bandgap, and a high melting point are only a few of TiO₂'s physical and chemical features. [3,4]. TiO₂ thin films exist in three crystalline phases: brookite, rutile, and anatase [5]. They take different structures; however, their chemistry is the same [2]. The chemical and physical properties of TiO₂ depend on the phase structure [6]. Because of its varied morphologies, such as hollow spheres, nanorods, nanotubes, and nanoparticles, titanium dioxide has been researched for a range of uses. Titanium dioxide nanorods (1D) have a one-dimensional shape that provides quick and direct transfer channels for injected electrons, as well as a longer electron diffusion length. [7]. In comparison to its particulate forms, the structure of the TiO₂ nanorod has a larger surface area [8]. Chemical bath deposition (CBD), hydrothermal approach, solvothermal, chemical vapor deposition (CVD), and DC reactive magnetron sputtering are some of the synthetic procedures for generating TiO₂ nanorod structures that have been documented [9,10]. The high reactivity of reactants, facile regulation of solution or interfacial reactions, creation of metastable and

distinctive condensed phases, decreased air pollution, and low energy consumption all contributed to the growing interest in hydrothermally synthesized compounds. [11]. Due to oxygen vacancies, titanium dioxide TiO_2 is an n-type metal oxide semiconductor. Its conductivity arises in proportion to the amount of oxygen lost. It has a high visible transmission, a high refractive index, and high photocatalytic effectiveness [3]. It has a higher charge distribution constant and a larger energy band gap (rutile energy bandgap = 3.05 eV). Because of its unique chemical/physical properties, (TiO_2) is used in photovoltaic and electromechanical devices [12]. TiO_2 NRs have narrow grain boundaries, allowing for quick charge carrier transfer. Previous research indicated that TiO_2 NRs had superior photocatalytic activity than other nanoparticles due to a large number of active sites. TiO_2 film has greater surface activity, making it suitable for antibacterial and self-cleaning applications in the industry [13]. TiO_2 nanorod arrays can be used as a photocatalytic coating to improve the antifungal and antibacterial efficiency of Ti substrates [14]. In addition to the previously described applications, when the nanostructured TiO_2 substrate is used under AM 1.5 illumination, photovoltaic performance reaches a power conversion efficiency of 0.39 percent and a fill factor of 0.47 [15].

In this work, We'll use a hydrothermal approach to make TiO_2 NRs array films on FTO substrates and investigate their optical characteristics. By adjusting the growth parameters, reaction time, temperature, precursor concentration, and substrate orientation. The diameter, length, and density of the nanorods could be altered. The nucleation and development of TiO_2 nanorods on FTO are aided by the epitaxial interaction between the FTO substrate and TiO_2 with a modest lattice mismatch [16].

2. Theoretical Part

2.1. Hydrothermal Technique

A hydrothermal operation is any heterogeneous reaction that takes place in the presence of aqueous solvents under high pressure and temperature to dissolve and recrystallize materials that are generally intractable under normal conditions. [17]. The technique is referred to as hydrothermal when water is used as the solvent [18]. Because of its low cost, quick reaction velocity, and simplicity, hydrothermal synthesis has been considered as a popular method for producing titanium dioxide NRs [19]. Also, as shown in Figure 1, Hydrothermal methods could be used to directly prepare well-crystallized and pure nanostructured materials processes, leading to more chemical treatments being avoided [20].

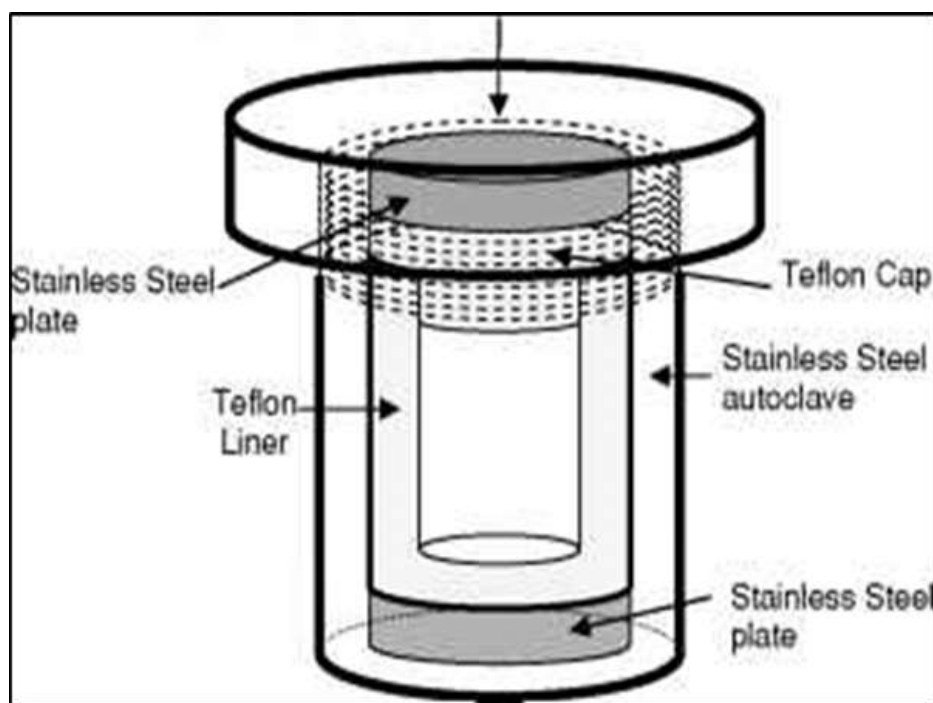
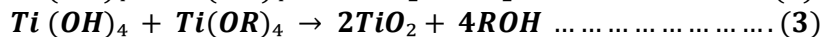
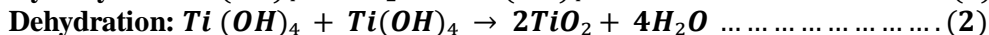
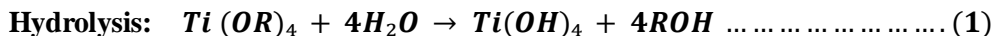


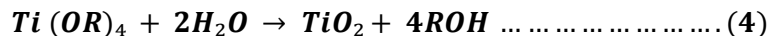
Figure 1: Hydrothermal autoclave parts use to prepare TiO_2 nanorods at 180 °C for 3 h.

2.2. Growth Mechanism

In general, a hydrothermal approach for the synthesis of TiO₂ NRs is carried out in hydrochloric acid (HCl) diluted with deionized water (DI), with Ti (IV) butoxide (TBO) having the structural formula Ti(OCH₂CH₂CH₂CH₃)₄ (or [Ti(RO)₄]) as a reactant. Several ways to describing TiO₂ growth mechanisms have been proposed, all of which are intriguing and useful in our study. When the four-fold Ti precursor ([Ti(RO)₄]) combines with water, it becomes [Ti(OH)₄] [21].



The entire reaction via the hydrothermal process between TBO precursor and water is [21]



3. Experimental Procedure

The first stage in the experimental part is cleaning the substrates. we use ethanol alcohol with purity of 99.9% to clean the substrate with ultrasound for 15 minutes. After that, we use deionized water to clean the sample in the second stage for 15 minutes in the same device and then let it dry well before use. The hydrothermal technique in a chemical solution was used to make titanium dioxide nanorods arrays in a Teflon-lined (100 ml) autoclave.

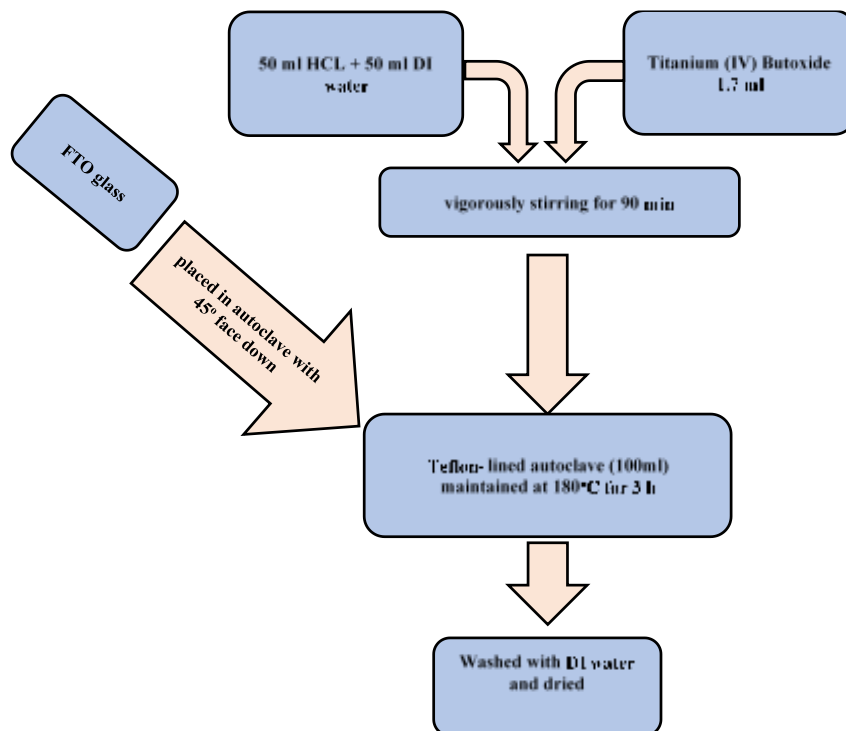


Figure 2: Illustrate the flowchart of experimental work.

Figure 2 shows the flow chart of experimental work. The concentrated hydrochloric acid (35-38 wt. percent) from the Central Drug House (P) Ltd Group in New Delhi was diluted in 50 ml of deionized water (DI). Then, for 90 minutes, aggressively swirl 1.7 ml titanium butoxide (C₁₆H₃₆O₄Ti) (TBOT) purity 99.9% (Germany). The solution was then placed into a Teflon-lined autoclave to begin the hydrothermal process. At 180 °C, the hydrothermal reaction takes 3 hours to complete. The autoclave was then taken out of the oven and left to cool to room temperature. After that, DI water was used to rinse the created film.

4. Results and Discussion

4.1. X-ray Diffraction

Figure 3 shows the X-ray diffraction of a TiO₂ sample prepared on a substrate of FTO using the hydrothermal method. The figure shows the appearance of two phases of rutile and Anatase TiO₂ (110) (27.4460°), (101) (36.0850°), (211) (54.3216°), (221) (65.7480°) for rutile phase respectively, anatase peak (004) (37.9658°). After comparing these peaks with the standard of TiO₂, for Rutile ICDD card (021-1276) and anatase ICDD card no. (046-1088), we have noticed that the peaks are identical. The benefit of a biphasic TiO₂ structure is that the different positions of the valence/conduction bands energy levels of anatase and rutile lead to charge carrier separation [22]. When these two phases are present at the same time, they have a higher Photoelectrochemical activity than when they are present separately. Meanwhile, the biphasic TiO₂ structure allows for the use of both phases' properties. For example, rutile has higher chemical stability, refractive index, and bandgap energy than anatase [23], but anatase has a larger specific surface area. The measurement of crystallites size was estimated using the Scherrer equation.

$$D = \frac{k\lambda}{\beta \cos\theta} \dots\dots\dots (5)$$

where D is a crystallite size, K is the shape factor (0.9), β is the Full-width at half maxima (FWHM) (in radians), λ is the X-ray wavelength (0.15418 nm) and θ is the Bragg angle of diffraction. The calculated crystallite size by equation 5 was about 30 nm.

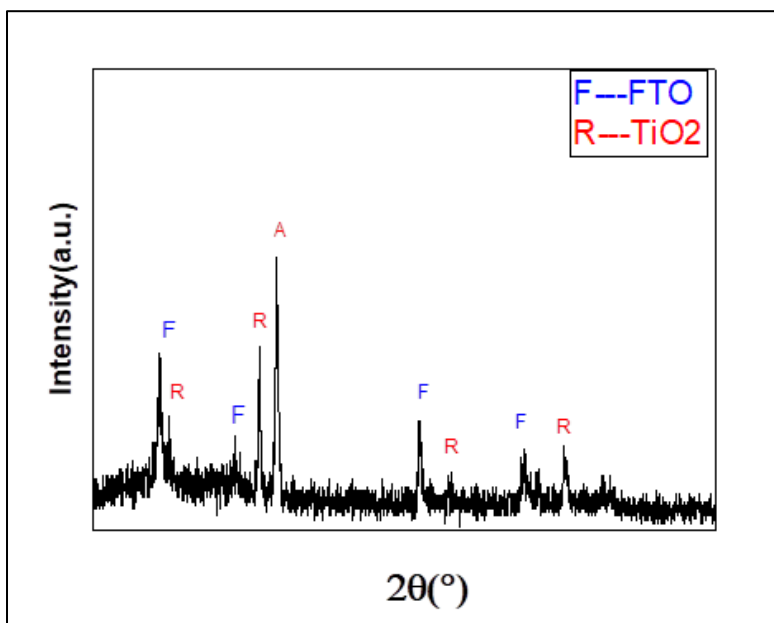


Figure 3. XRD pattern of TiO₂ NRs film prepared on FTO by hydrothermal method at 180 °C for 3h.

4.2. Field Emission Scanning Electron Microscopy (FE-SEM)

Figure 4 exhibits the FE-SEM images of the surface morphology of TiO₂ NRs arrays synthesized by the hydrothermal method. The images show that the entire surface of the FTO substrate is covered uniformly with ordered TiO₂ nanorods, with an average diameter of 65.24 nm calculated by the software (Image J). The nanorods appeared as nanowire bundles.

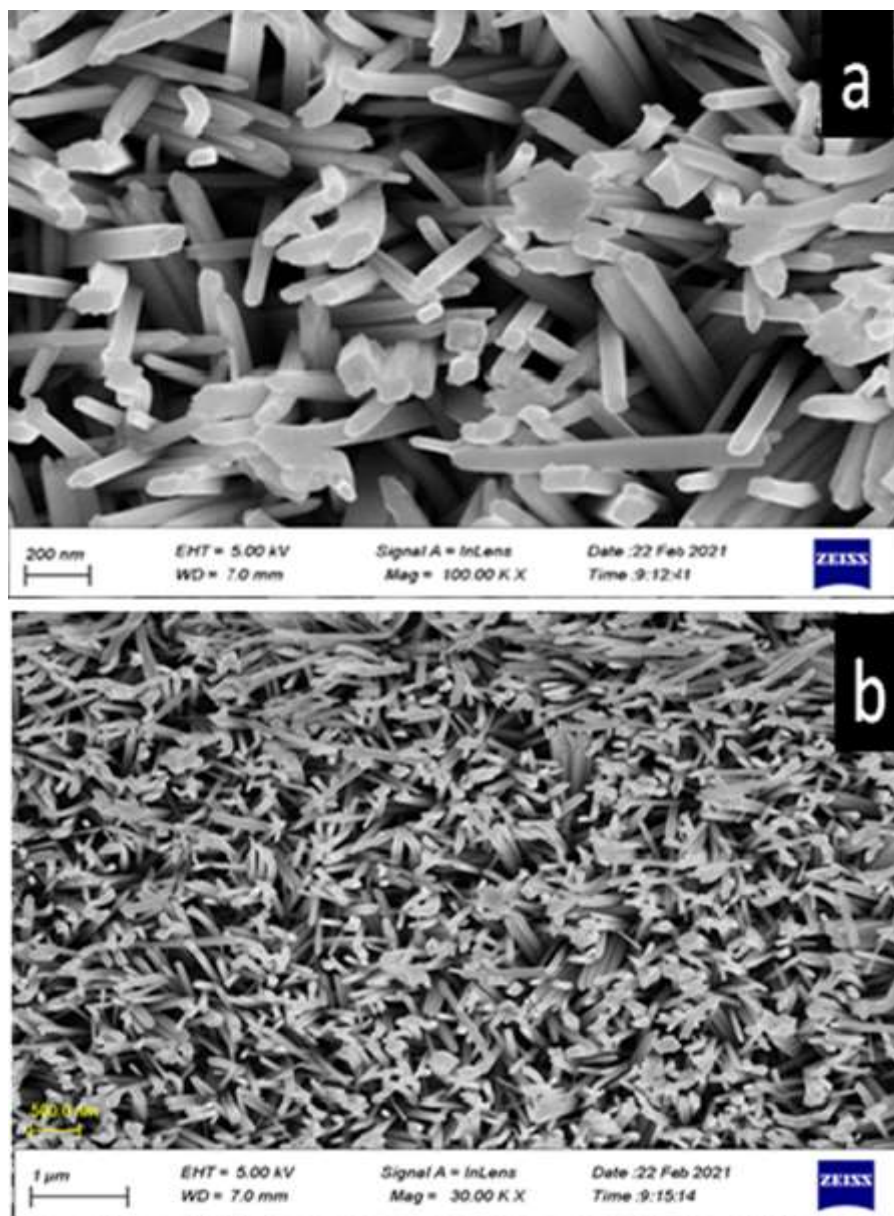


Figure 4: FE-SEM image for the surface of TiO₂ NRs with two magnifications (a)100 KX (b) 30 KX prepared by hydrothermal method 180 °C for 3 h.

4.3. Optical Properties

4.3.1 UV-Vis Measurement

The UV-Visible transmission spectrum and absorption coefficient of the TiO₂ NRs array grown on FTO substrate by hydrothermal technique in the range (380–1100) nm are shown in Figure 5 a, b, and c. The absorption coefficient spectrum of TiO₂ peaks at 380 nm and then gradually decreases with increasing wavelength, whereas the transmission spectrum increases with increasing wavelength. The optical energy band gap measurement of 3.18 eV confirms the identity of the material [24]. The following equation was used to calculate the energy bandgap [25].

$$\alpha h\nu = A(h\nu - E_g)^{1/2} \dots\dots\dots (6)$$

where A is constant and measures the disorder of the material, α absorption coefficient, ($h\nu$) (where h is Plank's constant ν is the photon frequency), (E_g) bandgap.

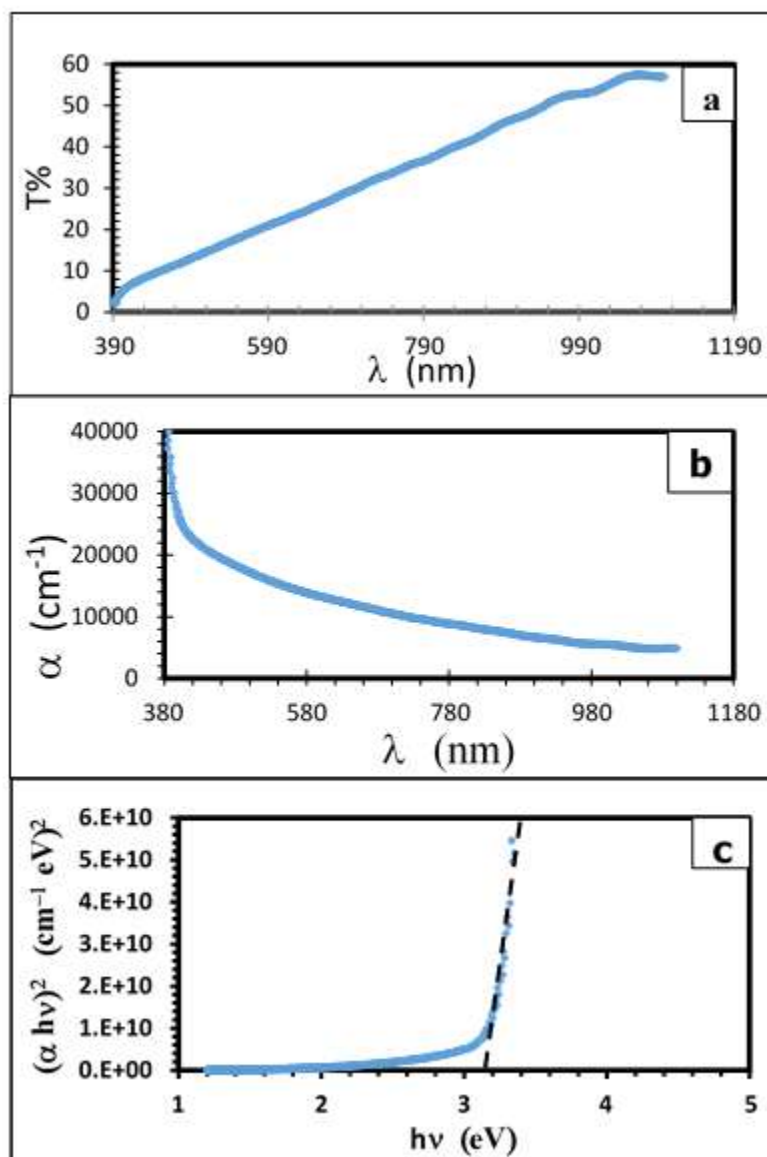


Figure 5: (a) Transmission curves for pure TiO₂ NRs (b) absorption coefficient variation with λ for TiO₂ NRs (C). energy gab for TiO₂ NRs.

4.4. Photoluminescence Emission Spectra Properties (PL)

The interband recombination peak formed at roughly 360 nm when the TiO₂ nanorods film was activated with exciting wavelengths less than 320 nm [26]. Due to oxygen vacancies and hydroxyl groups, another peak emerged in the visible region of the spectrum. The most prevalent places for trapped electrons and holes are oxygen vacancies and surface hydroxyl groups. These trapped carriers, which are captured by oxygen vacancies and surface hydroxyl groups, create visible luminescence in samples. [27]. The interband recombination peak formed at roughly 360 nm when the TiO₂ nanorods film was activated with exciting wavelengths less than 320 nm [26], and another peak appeared in the visible portion of the spectrum due to oxygen vacancies and hydroxyl groups. For trapped electrons and holes, oxygen vacancies and surface hydroxyl groups are the most common locations. The visible luminescence in samples is caused by these trapped carriers, which are caught by oxygen vacancies and surface hydroxyl groups [28].

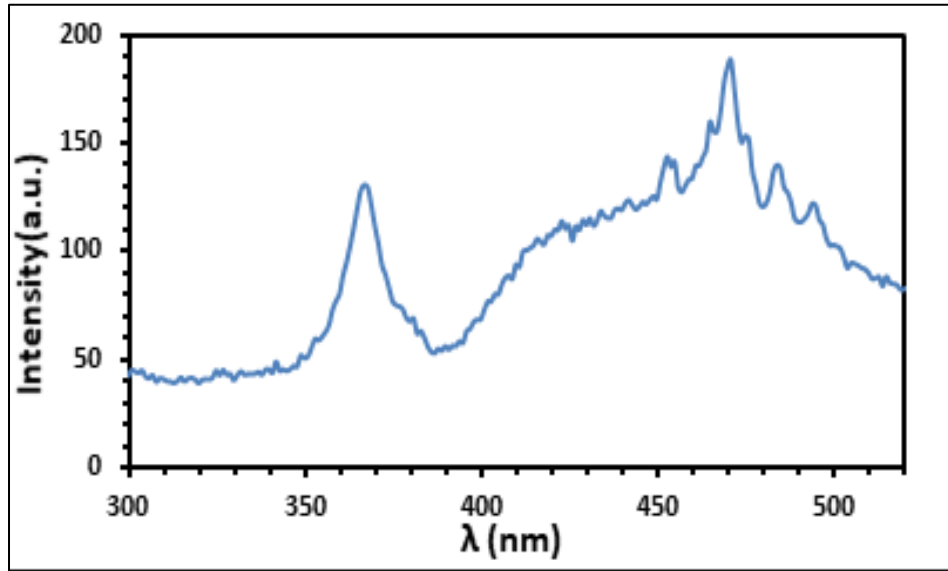


Figure 6: Photoluminescence (PL) for TiO₂ NRs on FTO substrate by the hydrothermal method at 180 °C for 3h.

4.5. Reflectance

For analyzing semiconducting materials, spectroscopic techniques are quite useful. In this study, we used a different method to determine the energy bandgap. As can be observed in Figure 7, the TiO₂ reflection begins to rise around 350 and peaks at 390. The energy bandgap can be used to confirm the identity of the material [29]. As in the absorption spectrum, we plot a graph between $h\nu$ (abscissa) and the square of $\ln\left(\frac{R_{max}-R_{min}}{R-R_{min}}\right)$ to get a straight line. The extrapolation of a straight line to $\left[\ln\left(\frac{R_{max}-R_{min}}{R-R_{min}}\right)\right]^2 = 0$ axis gives the value of the optical energy band gap, as in the following equation: [29].

$$\ln\left(\frac{R_{max}-R_{min}}{R-R_{min}}\right) = A(h\nu - E_g)^{1/2} \dots\dots\dots (7)$$

where: R is the reflectance and is given by the ratio of the intensity of reflected beam (I) to incident intensity (I₀).

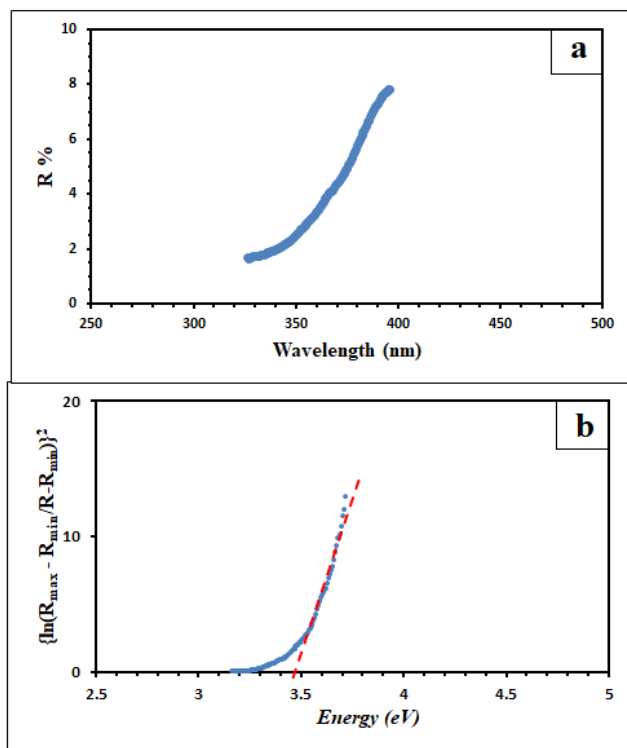


Figure 7: (a) Reflection spectra of TiO₂ nanorods by hydrothermal method and (b) energy bandgap.

It is obvious from equation 7 that $\ln[(R_{\max} - R_{\min}) / (R - R_{\min})]$ and, where R_{\max} and R_{\min} are the greatest and minimum reflectance in reflection spectra, respectively, and R is the reflectance for any intermediate energy photons measured.

5. Conclusions

Physical investigations of titanium dioxide nanorods film have been fabricated via hydrothermal method using TBOT as a precursor. XRD results show a two-phase structure i.e., rutile and anatase. The whole surface of the FTO substrate is uniformly coated with TiO₂ nanorods, according to FE-SEM pictures. According to the optical measurement results, there is a high agreement in the values of the optical energy band gap calculated by the three measurements i.e., UV-Vis transmittance, photoluminescence, and reflectance, and its values were 3.18, 3.3, and 3.45 eV, respectively.

Conflict of Interest

There are no conflicts of interest.

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