

Effect & Growth of TiO₂ Thin Films for Solar Cell Applications

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Abstract

Titanium di-oxide (TiO₂) has many excellent physical properties such as high dielectric constant, a strong mechanical and chemical stability. TiO₂ has high refractive index and good insulating properties, as a result it is widely used as a protective layer for very large scale integrated (VLSI) circuits and for the manufacture of optical elements. Additionally, TiO₂ films have potential uses for a number of electronic device applications such as dye-sensitized photovoltaic cells as well as anti-reflective (AR) coatings, gas sensors, electrochromic displays, and planar waveguides.

The nanomaterials were characterized using x-ray diffraction, UV-visible reflectance spectroscopy. In this paper, we have discussed the growth of the high quality Titanium di-oxide (TiO₂) material and study their optical and electrical properties.

Keywords: TiO₂, XRD, PLD, UV/Vis, Thin Film, DSSC

I. Introduction

This Titanium dioxide (TiO₂) nanomaterials are used in a wide range of applications such as (photo) catalysis, separations, sensor devices, paints, and dye-sensitized solar cells [1–4]. The material properties of TiO₂ nanoparticles are a function of the crystal structure, nanoparticle size, and morphology and, hence, are strongly dependent on the method of synthesis [5–15]. TiO₂ exists in three main phases: anatase, brookite and rutile. As a bulk material, rutile is the stable phase; however, solution-phase preparation methods for TiO₂ generally favour the anatase structure [8]. Titanium-di-oxide (TiO₂) can be grown mainly with anatase (tetragonal), rutile (tetragonal) and brookite (orthorhombic) crystalline structures; among this rutile is the most stable phase. The actual efficiency of Titanium-di-oxide (TiO₂) depends not only on its phase composition but also in its microstructure, Particle size, morphology and porosity which is in turn controlled by the synthesis method employed. Nanocrystalline TiO₂ usually exhibits a wider bandgap than that of the bulk (3.03 eV for rutile and 3.20 eV for anatase). Moreover, anatase becomes more stable than rutile when the particle size is decreased below 14 nm. Generally speaking, the functional properties of nano-TiO₂ are influenced by a large number of factors which include particle size, surface area, synthesis method and conditions, and crystalline. All the specific properties of nanosized particles have led to the exploitation of nano-TiO₂ for a wide variety of applications in which nano-TiO₂ is essentially preferred over conventional TiO₂ particles. Such applications include self-cleaning surfaces and textiles, UV-resistant coatings and paints, disinfectant sprays, sunscreens, water treatment agents, anticancer treatments [5]. TiO₂ is indeed one of the most widely used nanoscale materials. Amongst the multiple uses of nano-TiO₂, two major applications can be highlighted in the field of clean energy, namely photo catalytic water splitting and solar cells. Photo catalytic water splitting into H₂ and O₂

using nanostructured TiO₂ electrodes is thoroughly investigated as it is an environmentally friendly way to produce hydrogen. Another promising application of the TiO₂ semi conductivity is as electrode in dye-sensitized solar cells (DSSCs) in which the high surface-to-volume ratio of the nanostructured semiconductor is required to obtain an acceptable power conversion efficiency.

In this paper, we have discussed the Preparation of nanostructured Titanium-dioxide (TiO₂) thin film using laser ablation on silicon substrate kept in vacuum. Pulse laser Deposition (PLD) is a simple low cost method to grow oxide films, effect of film deposition conditions and structural, electrical and optical properties of films have been discussed. The measurement of the Band Gap is also discussed using the UV/Vis Spectroscopy Technique of nanostructured Titanium-dioxide (TiO₂) thin film using laser ablation on quartz substrate kept in vacuum is studied.

II. Thin Film Technology

The field of material science and engineering community's ability to conceive the novel materials with extraordinary combination of chemical, physical and mechanical properties has changed the modern society. There is an increasing technological progress. Modern technology requires thin films for different applications [1]. Thin film technology is the basic of outstanding development in solid state electronics. The usefulness of the optical properties of metal films, and scientific curiosity about the behavior of two-dimensional solids has been responsible for the immense interest in the study science and technology of the thin films. Thin film studies have directly or indirectly advanced many new areas of research in solid state physics and chemistry which are based on phenomena uniquely characteristic of the thickness, geometry, and structure of the film. When we consider a very thin film of some substance, we have a situation in which the two surfaces are so close to each other that they can have a decisive influence on the internal physical properties and processes of the substance, which differ, therefore, in a profound way from those of a bulk material. The decrease in distance between the surfaces and their mutual interaction can result in the rise of completely new phenomena. Here the one dimension of the material is reduced to an order of several atomic layers which creates an intermediate system between macro systems and molecular systems, thus it provides us a method of investigation of the microphysical nature of various processes. Thin films are especially appropriate for applications in microelectronics and integrated optics [6]. However the physical properties of the films like electrical resistivity do not substantially differ from the properties of the bulk material. For a thin film the limit of thickness is considered between tenths of nanometer and several micrometers.

Thin film materials are the key elements of continued technological advances made in the fields of optoelectronic, photonic, and magnetic devices. The processing of materials into thin films allows easy integration into various types of devices. The properties of material significantly differ when analyzed in the form of thin films. Most of the functional materials are rather applied in thin film form due to their specific electrical, magnetic, optical properties or wear resistance. Thin film technologies make use of the fact that the properties can particularly be controlled by the thickness parameter. Thin films are formed mostly by deposition, either physical or chemical methods. Thin films, both crystalline and amorphous, have immense importance in the age of high technology. Few of them are: microelectronic devices, magnetic thin films in recording devices, magnetic sensors, gas sensor, photoconductors, IR detectors, interference filters, solar cells, polarizer's, temperature controller in satellite, superconducting films, anticorrosive and decorative coatings.

III. Experimental Details

Thin films were deposited using pulsed laser deposition unit by employing a Q switched: KrF laser at wavelength 248 nm with 220 mJ of laser energy, pulse width 8 ns and repetition frequency 10 Hz. Uniform ablation was ensured by rotating the target at a constant speed. Thoroughly cleaned fused amorphous quartz plates were used as substrates. Time of deposition was fixed to be 30 min. Substrate heating was provided in the temperature range $T_s = 600$ C. Pressure inside the deposition chamber was reduced to $\sim 10^{-6}$ mbar before deposition. The deposition process was repeated by varying substrate-target (DS-T) distance from 4 to 6 cm. The structure and crystalline of the films were analyzed by X-ray diffraction (XRD) technique using D8 Advance diffractometer operated with a monochromatic $\text{Cu K}\alpha$ radiation source ($\lambda = 0.15418$ nm). Optical measurements were conducted in the wavelength range 300 to 900 nm using a PerkinElmer US Lambda 950, UV-Visible spectrophotometer.

IV. Results and Discussion:

A. XRD Pattern of Pure Silicon (Si) Substrate

The XRD pattern of the Pure Silicon (Si) Substrate is shown in Figure 1.

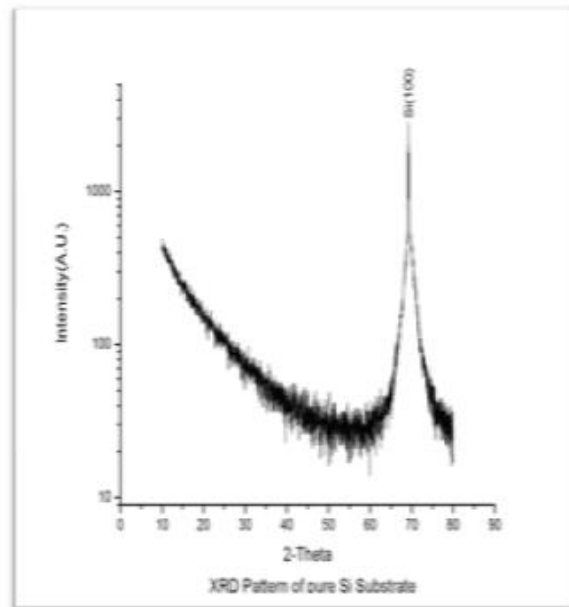


Figure 1. XRD Pattern of Pure Silicon(Si)

B. XRD Pattern of Pure Target Titanium-di-oxide (TiO₂)

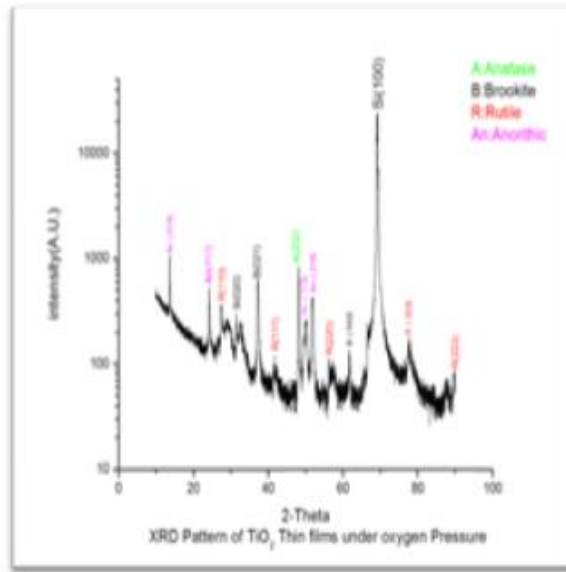


Figure 2. XRD Pattern of Pure Target Titanium-di-oxide (TiO₂)

The XRD pattern of the Pure Target Titanium-di-oxide is shown in Figure 2.

C. XRD Pattern of TiO₂ Thin films under oxygen pressure

The XRD pattern of the TiO₂ films deposited by Pulse laser Deposition under different conditions is shown in Figure 3.

All these films were indexed and Matched with TiO₂ Phase as in JCPDS Card (No. 89-4920), JCPDS Card (No. 89-4921), JCPDS Card (No. 76-1937), JCPDS Card (No. 85-1060). The fig XRD Pattern indicates that these films have Polyphase contribution with weak peak indicating the (R) rutile phase formation, (A) Anatase phase formation, (B) Brookite Phase Formation and Another Phase (An) Anorithic Phase Formation also Present.

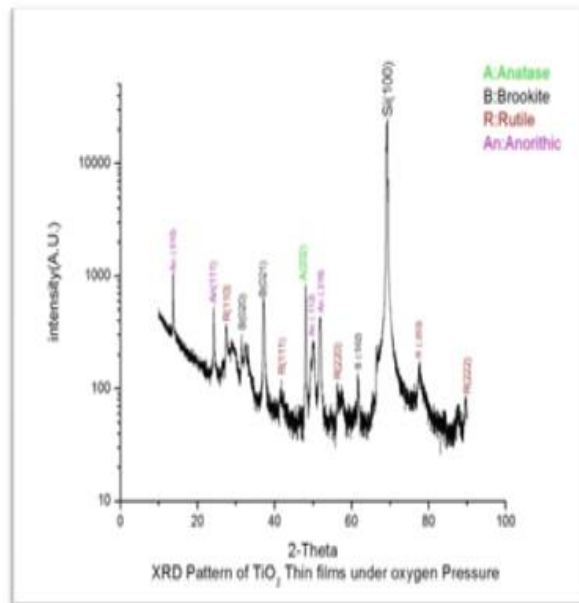


Figure 3. XRD Pattern of TiO₂ Thin Films Under Oxygen Pressure

D. XRD Pattern of TiO₂ Thin films without oxygen pressure

The XRD pattern of the TiO₂ films deposited by Pulse laser Deposition under different conditions is shown in Figure 4.

All these films were indexed and Matched with TiO₂ Phase as in JCPDS Card (No. 89-4920), JCPDS Card (No. 89-4921), JCPDS Card (No. 76-1937), JCPDS Card (No. 85-1060). The fig XRD Pattern indicates that these films have Polyphase contribution with a weak peak indicating the (R) rutile phase formation, (A) Anatase phase formation, (B) Brookite Phase Formation and Another Phase (An) Anorithic Phase Formation also Present.

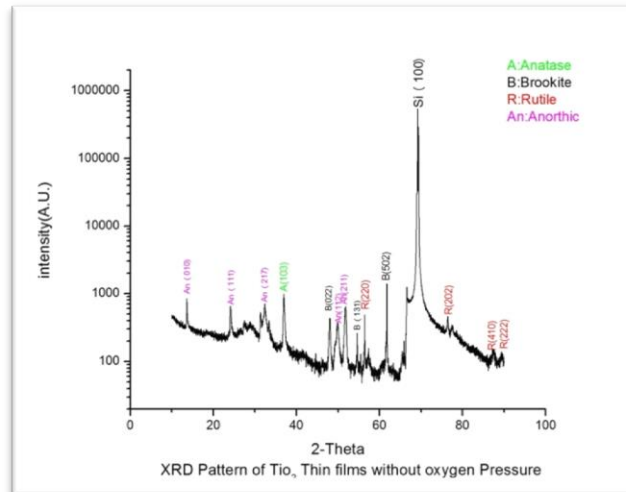


Figure 4. XRD Pattern of TiO₂ Thin Films without Oxygen Pressure

E. Band Gap Measurement of TiO₂ ThinFilm Deposited on Quartz Crystal (SiO₂)

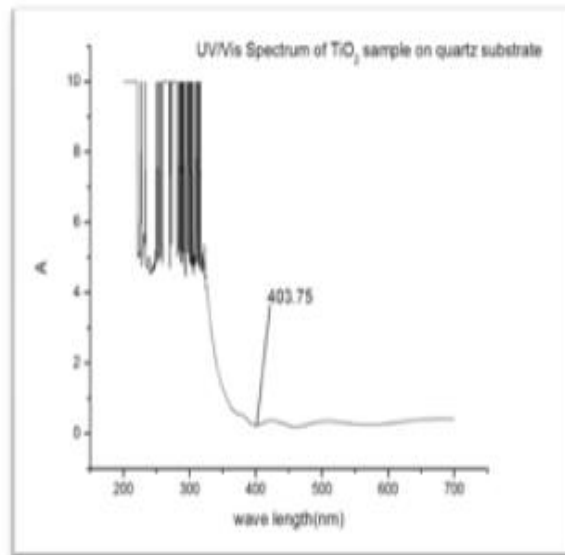


Figure 5. TiO₂ UV/Vis Spectrum Obtained in this Work

Calculations:

$$\text{BandGap Energy (E)} = h \cdot C / \lambda$$

$$h=6.626 \times 10^{-34} \text{ Joules sec}$$

$$C = 3.0 \times 10^8 \text{ meter/sec}$$

$$\lambda = 403.75 \times 10^{-9} \text{ meters}$$

$$E = 0.049233 \times 10^{-17} = 3.07 \text{ eV}$$

V. Conclusions

Semitransparent and nanostructured highly conducting TiO₂ (Titanium-di-oxide) thin films were prepared using Pulse laser deposition (PLD) technique, in this Study. The Thin film has been deposited by Pulse laser deposition (PLD) technique under different oxygen fluence. The Variation of oxygen pressure indicates the different phases or different growth of TiO₂ thin films are present. The Substrate used is silicon [si (100)]. The Structure is Characterized by the Bruker D-8 Advance X-ray diffractometer. The X-Ray Pattern suggested that it is crystalline in nature. The Film growth is found Polycrystalline in nature is A (Anatase), B (Brookite), R (Rutile) Phases are well matched by Standard data base files in the vacuum condition. The Phase Anorthic has are going to increasing the oxygen Pressure. The TiO₂ Thin films phases Anatase and Anorthic has been dominant. This is clearly visible in X-Ray Diffractometer Pattern .The Tio₂ Thin films growth is the Polycrystalline. TiO₂ thin films as they are prepared using the Pulse laser Deposition (PLD) Technique. The Tio₂ Thin films growth is the Polycrystalline. The energy Band gap is found 3.07eV. The quality of TiO₂ also can be determined. The XRD characteristic clearly show the Thin film is applicable for the fabricating a solar cell.

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