Influence of the growth parameters on TiO$_2$ thin films deposited using the MOCVD method

(Influência dos parâmetros de crescimento em filmes finos de TiO$_2$ depositados pelo método MOCVD)

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Abstract

In this work we report the synthesis of TiO$_2$ thin films by the Organometallic Chemical Vapor Deposition (MOCVD) method. The influence of deposition parameters used during the growth in the obtained structural characteristics was studied. Different temperatures of the organometallic bath, deposition time, temperature and type of the substrate were combined. Using Scanning Electron Microscopy associated to Electron Dispersive X-Ray Spectroscopy, Atomic Force Microscopy and X-ray Diffraction, the strong influence of these parameters in the thin films final microstructure was verified.

Keywords: organometallic compounds, thin films, optical properties.

Resumo

A síntese de filmes finos de TiO$_2$ por meio do método MOCVD é descrita. Foi estudada a influência de parâmetros de deposição usados durante o crescimento nas características estruturais obtidas. Foram combinados diferentes temperaturas do banho organometálico, tempo de deposição, temperatura e tipo do substrato. A forte influência destes parâmetros na microestrutura final dos filmes foi estudada por meio de diferentes técnicas: microscopia eletrônica de varredura associada com espectroscopia eletrônica de dispersão de raios X, microscopia de força atômica e difração de raios X.

Palavras-chave: compostos organometálicos, filmes finos, propriedades ópticas.

INTRODUCTION

Titanium dioxide has several advantageous properties that make it a candidate for different applications. Reports of its high dielectric constant, chemical stability, high refraction index and semiconductor properties motivated research in the growth of high-quality thin films by a variety of techniques including MOCVD [1-3], pulsed laser deposition [4], sputtering [5-7], sol-gel [8] and filtered arc deposition (FAD)[9]. There are three types of TiO$_2$ crystalline structures: anatase, rutile, and brookite. Rutile presents the highest refractive index and is the most thermodynamically stable structure. It is also well known that titanium dioxide presents two isomorphous crystal phases: the anatase and the rutile [10]. The anatase structure is obtained at temperatures of around 350 °C, which renders it more useful for industrial applications. At temperatures between 400 and 800 °C, the rutile phase is also present, while at higher temperatures, only the rutile structure is present.

The method known as Organometallic Chemical Vapor Deposition (MOCVD) consists of heating an organometallic solution, which evaporates and is deposited on a heated substrate. The films grown by this method usually require expensive, sophisticated apparatus.

In this letter, we report on the sintering procedures for growing TiO$_2$ thin films using a simple, low-cost deposition apparatus especially built in our laboratory. The films were characterized by X-ray diffraction (XRD), scanning electron microscopy (SEM) and atomic force microscopy (AFM).

EXPERIMENTAL PROCEDURES

TiO$_2$ thin films were deposited on Si(100) and glass substrates using the organometallic chemical vapor deposition (MOCVD) system shown in Fig. 1. Titanium isopropoxide [Ti(OCH(CH$_3$)$_2$)$_3$] which is liquid at room temperature (melting point 20 °C), was used as the organometallic (OM) precursor. The titanium isopropoxide was stored in a glass bubbler whose temperature was controlled by a hot plate. The vapor of the OM precursor was transported by high purity oxygen gas to the reactor. Pure oxygen was used as oxidant. Single-layer films were grown using Si (100) and glass substrates and different deposition times, substrate temperature and the organometallic...
Each pair of thin film samples (A1G-A1S and A2G-A2S) was obtained from the same deposition run (same conditions) but using different substrates; G stands for glass and S for silicon(100), as shown in Table II.

The structural properties of the deposited films were studied by X-ray diffraction, and the measurements were carried out with a Siemens D5000 diffractometer with CuK\textsubscript{α} radiation. The geometry of the diffractometer was the same for all the samples studied (grazing incidence diffraction - incidence angle = 2°, step time = 7s, step scan = 0.007°, 2θ = 20 - 50°, U = 40 kV and I = 40 mA). The surface morphology of the TiO\textsubscript{2} thin films was analyzed by scanning electron microscopy using a Zeiss DSM940A microscope. Film thickness was determined by analyzing the cross section images. More detailed information on morphology and roughness was obtained by atomic force microscopy (AFM) (Digital Instruments Multi-Mode Nanoscope III A).

**RESULTS AND DISCUSSION**

The results showed that the amorphous glass structure lead to titania in the amorphous phase. Otherwise, the crystalline silicon substrate, induced the anatase crystal structure for the film.

It was possible to observe that an obtained result, using the same deposition procedure, was always different depending on the used substrate. An example of this fact can be observed in Fig. 2 and Table II, for films grown over glass and Si (100) substrates following the same conditions. As can be seen from the XDR diffraction patterns, the structures are different, showing the influence of the nature of the substrate in each case. This figure also shows that the films deposited on Si (100) present a polycrystalline structure with the anatase phase. The films grown on glass, on the other hand, do not show...
Figure 3: SEM micrographs of the cross section of TiO$_2$ thin films deposited on different substrates, under the same conditions: a) glass and b) Si(100).

[Figura 3: Micrografias MEV de secção transversal de filmes finos de TiO$_2$ depositados em diferentes substratos sob as mesmas condições: a) vidro e b) Si (100).]
crystalline structure, preserving their amorphous character.

The nature of the substrate, crystalline or amorphous, provides different degrees of packing and, consequently, different stages of density and thickness in the morphological structure of the obtained films [11]. This may be attributed to the fact that the mobility of the atoms on the substrate surface, which is responsible for the degree and type of nucleation on the substrate, depends on its nature.

The morphological analyses were of key importance to choose the substrates and bath precursors temperature. The collection of SEM micrographs presented in Fig. 3 shows surface and cross section views for the obtained TiO₂ thin films. In Table II are listed the obtained films nomenclature.

Comparing the micrograph studies for A1 (S and G) with the A3 (S and G), growth using substrate temperature $T_s = 400 ^\circ C$ and $T_s = 500 ^\circ C$, respectively, it was observed that the A6G thin film presents a distinct structure than other films. This special feature has already been observed in thin films obtained by the microwave growth technique [12]. For the A3S film, on the other hand, it was observed that the growth occurred in order to form plane plates with low adherence. Analyzing both A1S and A1G films, one can observe that these films present a homogeneous surface, with no dendritic or plate-form arrangements and uniform thickness. On the contrary, for the growth of A3 films, the determination of the thickness was not possible due to morphology irregularities. These results indicate that $T_s$, the optimum temperature of the substrate for deposition of TiO₂ films using the built apparatus is 400 °C.

A comparison of the results of A1(S and G) with the A4(S and G) growth in the bath temperature $(T_B)$ of 90 °C and 120 °C, respectively, was done. These temperatures are thoroughly related to the partial pressure of the organomettalic compound, and the higher is the temperature, the vapor partial pressure is so high that the deposition is inadequate, leading to a non-homogeneous film. Hence, it is possible to conclude that the temperature of 90 °C favors the growth of clean and more homogeneous films, since for films grown at 120 °C, A4 films, agglomerates are formed in the film surface.

In order to establish the chemical composition of the observed agglomerates, a chemical element mapping for the A4G film was also performed using the energy dispersive X-ray electron spectroscopy (EDS), as can be observed in Fig. 9. The elements found in such agglomerates are Au, from the recovering, Si, Na and Ca. These elements are present in the composition of the substrate and the occurrence of Na indicates a possible migration from the substrate surface to the film at this temperature.

After these analyses, it was established that the best temperature conditions were substrate temperature $T_s = 400 ^\circ C$ and bath temperature $T_B = 90 ^\circ C$. Using these parameters, it was possible to study the influence of the deposition time. The results, shown in Fig. 3, present just an increase in the thickness of the film. One can observe that the films deposited on the glass substrates are thicker than those grown on the Si (100) under the same conditions. It is also possible to observe a variation of

<table>
<thead>
<tr>
<th>Film Labels</th>
<th>Thickness (nm)</th>
<th>Color of the film</th>
<th>Roughness (nm)</th>
<th>Lattice Parameters</th>
<th>Crystallite Size (Å)</th>
<th>Cell Volume (Å³)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A1G</td>
<td>130</td>
<td></td>
<td>3.4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A1S</td>
<td>60</td>
<td>Light blue</td>
<td>4.8</td>
<td>3.7700 9.4548</td>
<td>325</td>
<td>134.38</td>
</tr>
<tr>
<td>A2G</td>
<td>230</td>
<td></td>
<td>0.9</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A2S</td>
<td>110</td>
<td>Green</td>
<td>2.0</td>
<td>3.7788 9.4970</td>
<td>723</td>
<td>135.62</td>
</tr>
<tr>
<td>A3G</td>
<td>varied</td>
<td></td>
<td>7.9</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A3S</td>
<td>varied</td>
<td>Colored*</td>
<td>10.4</td>
<td>3.7774 9.4970</td>
<td>643</td>
<td>135.51</td>
</tr>
<tr>
<td>A4G</td>
<td>varied</td>
<td></td>
<td>2.2</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A4S</td>
<td>varied</td>
<td>Colored*</td>
<td>-</td>
<td>3.7774 9.4916</td>
<td>420</td>
<td>135.44</td>
</tr>
</tbody>
</table>

(*) Colored means that the films have more than just one color.
color patterns with the increase of thickness, as shown in Table III. This effect of color variation is more evident for TiO$_2$ films grown on Si (100) substrates; it could also be observed on the glass ones, fact already reported elsewhere [12].

The films were visible at naked eye, once their color change, observed by reflection. TiO$_2$ layers on glass showed different colors (see Table III). These colored films were well red. The layer homogeneity was also visible, due to the interference phenomenon, also previously reported [13].

Studies of Atomic Force Microscopy were also performed to investigate the morphology of the films. In Fig. 5 it is possible to inspect an example of the surface structure and roughness.

Figure 5: AFM surface morphology of TiO$_2$ thin films deposited on different substrates, under the same conditions: A1G and A2G, glass (amorphous) and A1S and A2S, Si(100) (crystalline).

[Figura 5: Morfologia obtida por AFM de filmes finos de TiO$_2$ depositados sobre diferentes substratos nas mesmas condições: A1G e A2G, vidro (amorfo) e A1S e A2S, Si(100) (cristalino).]
listed in Table III. The data shows different crystallization patterns for both Si (100) and glass substrates. While the films grown on glass were composed of poorly defined grains, the films on Si (100) presented grain formations, in agreement with XRD data. It is necessary to point out that XRD experiments yields long and medium range order analysis, and on the contrary, the AFM is able to detect short range order. In our opinion, the AFM images of thin films on glass substrates here reported are just exhibiting the initial stage of crystallization, not detectable by X-ray diffraction analysis. Besides, observing the data presented in Table III, one can notice that the film roughness on glass substrates was always lower than those presented for the Si (100). This seems to confirm our results, since the roughness of the glass substrate (0.92 nm) is higher than that of Si (100), 0.20 nm.

The roughness values indicate the good quality and homogeneity of the films, showing that the increase of the deposition time leads to a decrease in the roughness, as reported for SnO2 films obtained using the Pechini method [14].

A first explanation is certainty related to the fact that the titania vapor is deposited over an organized structure resulting in a crystalline phase, or oppositely, the deposition over a random structure of those products leads to an amorphous specimen. Not just the crystal structure of the substrate, but also its chemical bonding affects the growth of the film, conditioned to the energy competition between the volume and the interface, and furthermore, to the growth kinetics, that is dependent on substrate temperature (ST) and deposition rate (DR).

CONCLUSIONS

It was possible to obtain good quality thin films of TiO2 with acceptable homogeneity, adherent, speculating and with minor roughness using a simple and non-expensive equipment. The films showed a strong dependence of the kind of substrate on the structural characteristics, and consequently to a variation of optical properties. The results suggested that the obtained film thickness is directly related to the deposition time. Through the analysis of the films grown under different conditions, it was chosen the temperature of 90 °C for the organometallic bath and 400 °C for the substrate temperature. Using these conditions, it was possible to obtain crystalline thin films using Si (100) substrates and amorphous thin films using glass substrates.

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