Hot-Filament Metal Oxide Deposition (HFMOD): A Novel Method for Depositing Thin Films of Metallic Oxides

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O presente artigo descreve um novo método para a síntese de filmes finos de óxidos metálicos. Um filamento de metal, que pode ser aquecido por uma fonte de corrente alternada é instalado numa câmara de vácuo. Por meio de um fluxômetro de massa, oxigênio pode ser admitido no interior da câmara. Da reação entre o oxigênio e o metal do filamento aquecido, espécies de óxidos voláteis MeOy, onde Me é o metal, são formadas, se condensando num substrato colocado próximo ao filamento, formando o filme. Foi observado que os filmes finos de WxOy e MoxOy podem ser depositados de forma satisfatória por este novo método. Embora várias outras técnicas de análise tenham sido usadas para caracterizar os óxidos, esta nota enfatiza os resultados obtidos por espectroscopia de fotoelétrons de raio-X (XPS).

This short report describes a novel method for the synthesis of metal oxide thin films. The experimental setup consists of a metal filament installed inside a vacuum chamber. The filament can be heated by an ac power supply while oxygen is admitted into the chamber using a mass flowmeter. From reactions between oxygen and the heated metal filament, volatile oxide species, MeOy, where Me is the metal, can be formed, condensing on a nearby substrate. We have observed that thin films of WxOy and MoxOy can be satisfactorily deposited by this novel method. Although several techniques were used to characterize the oxides, this note emphasizes the results obtained by X-ray Photoelectron Spectroscopy (XPS).

Keywords: WO3, MoO3, thin films, XPS, CVD

Introduction

Metallic oxides are important materials from the standpoint of both fundamental and applied science. Particularly, tungsten and molybdenum oxide films have been the focus of extensive scientific investigations due to their prospective technological applications in (i) electrochromic devices,1,2 (ii) gasochromic sensors,3,4 and (iii) electrocatalysis.5,6 However, their most intensively investigated property so far is the electrochromism. WO3 films are considered one of the most viable options in emerging electrochromic technology,1 being applicable for regulating the throughput of radiant energy in smart windows and antidazzling mirrors. These oxides have also been used as “templates” in the synthesis of composites of the type transition metal oxide/conducting polymer, generating promising results in the area of cathodic materials in secondary lithium batteries.7

Several deposition techniques such as sputtering,8-10 thermal evaporation,3,11 plasma-enhanced chemical vapor deposition12 and sol-gel13,14 have been used to obtain tungsten and molybdenum oxide films. This report describes the development of a new deposition method, which we have called hot filament metal oxide deposition (HFMOD), using a metallic filament heated in a rarefied oxygen atmosphere. The film is formed on a substrate positioned near the filament and the deposition rate is controlled by the filament temperature and the oxygen pressure. Both the thermochemistry of the process and the kinetics of film formation are currently under investigation. It is clear, however, that the film is formed from volatile MeOy precursors, where Me is the metal, generated on the heated tungsten surface from reactions between oxygen and tungsten. The investigations so far carried out in our laboratory show that the films can be deposited with a good stoichiometry control, with relatively high deposition rates and present good adhesion to both metallic and dielectric substrates. Tests on the electrochromical...
properties carried out on samples of WO₃ show that their optical efficiency is higher than those of WO₃ films obtained by the above-mentioned techniques. It is also important to remark that this technique differs from a deposition technique called hot filament chemical vapor deposition (HFCVD), which have been used to deposit siloxane¹⁵ and diamond-like films,¹⁶ because the filament used here is not just a “catalyst” used to activate chemical species; it is also a reactant in the reaction.

**Experimental**

Figure 1 depicts the experimental deposition setup that was designed in our laboratory for the deposition of transition metal oxides. The filament (F), which is made of the metal of the oxide to be deposited, is resistively heated by an ac current supply. Oxygen is admitted to the chamber via an electronic mass flowmeter. Pressure measurements are made using a capacitance manometer. During the depositions the chamber is continuously pumped by a Roots pump and the oxygen pressure is adjusted using the flowmeter. The chamber base pressure is about 2.0 x 10⁻² Pa. Substrate temperatures are measured using a chromel-alumel thermocouple and filament temperatures are determined with an optical pyrometer through a glass viewpoint in the wall of the deposition chamber.

The XPS data were gathered for a WₓOₓ and a MoₓOₓ sample deposited with the conditions given in Table 1.

The tungsten 4f XPS spectrum (squares) for sample WO is shown in Figure 2. As can be seen from that figure, the 4f profile can be fit by two Gaussian peaks centered at 37.7 and 35.5 eV which are, respectively, the binding energy of electrons in the 4f⁵/₂ and 4f⁷/₂ levels of tungsten in the W⁺⁶ valence state,¹⁷ indicating that the film is composed of stoichiometric WO₃.

**Results and Discussion**

The XPS spectrum of the Mo3d core levels for sample MoO is presented in this work. The most intense pair is centered at 235.8 and 232.7 eV which indicate the possible tungsten or molybdenum valence states, the 4f-doublet peak or the 3d-doublet peak, respectively, were fitted with Gaussian peaks corresponding to known bonding states of tungsten and oxygen.
are, respectively, the binding energy of electrons in the 3d_{3/2} and 3d_{5/2} levels of molybdenum in the Mo^{6+} valence state. The other pair is centered at 234.6 and 231.6 eV corresponding to the binding energy of electrons in the same 3d levels, but in MoO^{5+} valence state. For both valence states, these binding energies are in close agreement with literature values. \textsuperscript{18,19} Therefore, it was concluded that the Mo atoms were in mixed valence states, Mo^{6+} and Mo^{5+}, with a high predominance of the former over the latter. Thus the film was in the overall MoO_x stoichiometry, with x smaller but close to 3. The observation that the film stoichiometry is close to that of MoO_3, is consistent with the expectation that the film is formed from MoO_3 and MoO_2 species desorbed from the Mo filament and that the desorption rate of the former is greater than that of the latter.

The spectra of Figures 2 and 3 are representative of other spectra of samples prepared with oxygen flow rates in the interval between 6.0 and 21 sccm and the same filament temperatures. Thus the chemical composition of the films does not significantly change in this oxygen flow rate range.

Conclusions

By means of the unexpensive and simple deposition technique described in this note, it was possible to deposit W_xO_y and MoxO_y films with x close to 3. Several details about the role played by the deposition parameters on the overall structure and properties of the films are underway in our laboratory.

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