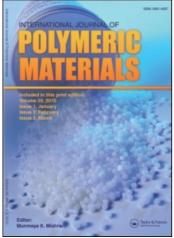
This article was downloaded by: On: *18 January 2011* Access details: *Access Details: Free Access* Publisher *Taylor & Francis* Informa Ltd Registered in England and Wales Registered Number: 1072954 Registered office: Mortimer House, 37-41 Mortimer Street, London W1T 3JH, UK



International Journal of Polymeric Materials

Publication details, including instructions for authors and subscription information: http://www.informaworld.com/smpp/title~content=t713647664

Investigation on Molybdenum Thin Films Deposited by DC-Sputtering on Polyethylene Terephthalate Substrate

M. G. Faraj^a; K. Ibrahim^a; M. H. Eisa^{ab}; M. K. M. Ali^a; F. Azhari^a

^a School of Physics, Universiti Sains Malaysia, Penang, Malaysia ^b Department of Physics, College of Science, Sudan University of Science and Technology, Sudan

Online publication date: 29 June 2010

To cite this Article Faraj, M. G., Ibrahim, K., Eisa, M. H., Ali, M. K. M. and Azhari, F.(2010) 'Investigation on Molybdenum Thin Films Deposited by DC-Sputtering on Polyethylene Terephthalate Substrate', International Journal of Polymeric Materials, 59: 8, 622 – 627

To link to this Article: DOI: 10.1080/00914031003760733 URL: http://dx.doi.org/10.1080/00914031003760733

PLEASE SCROLL DOWN FOR ARTICLE

Full terms and conditions of use: http://www.informaworld.com/terms-and-conditions-of-access.pdf

This article may be used for research, teaching and private study purposes. Any substantial or systematic reproduction, re-distribution, re-selling, loan or sub-licensing, systematic supply or distribution in any form to anyone is expressly forbidden.

The publisher does not give any warranty express or implied or make any representation that the contents will be complete or accurate or up to date. The accuracy of any instructions, formulae and drug doses should be independently verified with primary sources. The publisher shall not be liable for any loss, actions, claims, proceedings, demand or costs or damages whatsoever or howsoever caused arising directly or indirectly in connection with or arising out of the use of this material.

International Journal of Polymeric Materials, 59:622–627, 2010 Copyright © Taylor & Francis Group, LLC ISSN: 0091-4037 print/1563-535X online DOI: 10.1080/00914031003760733



Investigation on Molybdenum Thin Films Deposited by DC-Sputtering on Polyethylene Terephthalate Substrate

M. G. Faraj,¹ K. Ibrahim,¹ M. H. Eisa,^{1,2} M. K. M. Ali,¹ and F. Azhari¹

¹School of Physics, Universiti Sains Malaysia, Penang, Malaysia ²Department of Physics, College of Science, Sudan University of Science and Technology, Sudan

Molybdenum (Mo) films were prepared by DC sputtering on a polyethylene terephthalate (PET) substrate with different thicknesses. The molybdenum finds use in a very broad spectrum of applications in widely different forms. The obtained results of thin films of molybdenum deposited on PET are characterized by atomic force microscopy (AFM) and X-ray diffraction (XRD) and (EDX). It was found that the thickness increases with the time of deposition and reduces the resistivity and sheet resistance. The lowest resistivity value we found for the Mo films was $1.3 \times 10^{-5} \Omega \cdot \text{cm}$ at thickness (210 nm).

Keywords DC sputtering, molybdenum, polyethylene terephthalate

This work was supported by the Nano-optoelectronics Research Laboratory, School of Physics; Universiti Sains Malaysia under Grant No. 305/PFIZIK/613321.

Received 20 December 2009; in final form 7 March 2010.

Address correspondence to M. G. Faraj, School of Physics, Universiti Sains Malaysia, Penang 11800, Malaysia. E-mail: mohphysics_79@yahoo.com

INTRODUCTION

Molybdenum is a transition metal. The pure metal is silvery white in color and very hard. Molybdenum is used in some electronic applications as the conductive metal layers in thin-film transistors (TFTs) [1]. In many electronic applications, depositing a metal layer on a substrate can be performed by different techniques, such as thermal evaporation, electron beam evaporation, or sputtering [2–6]. Molybdenum has a high melting point (2623°C) and a low vapor pressure (3.47 Pa at 30°C) which makes molybdenum ideal for sputtering. However, the deposition of a molybdenum film as a back contact is not by itself an assurance of a high-efficiency solar cell. The deposition parameters and process play a key role in obtaining a layer with the appropriate properties. Extensive research has been done on the deposition of molybdenum thin films by DC sputtering [7,8]. The polyethylene terephthalate is an excellent commercial thermoplastic polymer resin of the polyester family. PET substrate has attracted interest in a wide array of fields because of its low cost, good thermal stability, surface inertness, good spin ability and excellent moisture resistance [9,10]. In this study, different thicknesses of Mo films deposed on PET substrates have been prepared by the DC sputtering method using low deposition temperature with different deposition conditions. The structural, optical and electrical properties of the obtained films depending on deposition parameters, such as sputtering power and working pressure, were investigated.

EXPERIMENTAL DETAILS

The PET substrate with thickness of $250 \,\mu\text{m}$ and dimensions $20 \,\text{mm} \times 20 \,\text{mm}$ were washed with alcohol and then ultrasonically cleaned for $10 \,\text{min}$. The deionized water was used to rinse the PET substrate. The PET substrate was dried by nitrogen gas. The molybdenum target with thickness of $0.3 \,\text{mm}$, diameter 75 mm and purity of 99.95% was deposited with PET ($T_{\text{melting}} = 265^{\circ}\text{C}$) using DC sputtering (model AUTO306). The residual gas pressure in the chamber was evacuated by a rotary and diffusion pump arrangement. The target was pre-sputtered for $10 \,\text{min}$ to remove contamination.

The shutter was displaced to expose the substrates in the sputtering plasma for 30 min. The sputtering deposition was carried out in a pure argon atmosphere at a pressure from 0.05 to 2 m torr and the sputtering power at 220 W. The sputtering was done in room temperature of 25°C and the distance between the target and PET substrate was approximately 6 cm. The thickness of the Mo film deposited on PET was determined for each time by using optical reflects of meter (Model: Filmetric F20). The deposition times were (5, 10, 15, 20 and 25 min). In this work, different thicknesses of Mo film (49 nm, 89 nm,

No.	Thicknesses (nm)	Time deposition (min)
1	49	5
2	89	10
3	150	15
4	180	20
5	210	25

Table 1: The thicknesses of molybdenum deposited on PET.

150 nm, 180 nm and 210 nm) were deposited. The surface morphology of each Mo films was performed by AFM (model: Ultra Objective) and SEM (model: JSM-6460 LV). The crystallographic structure of Mo deposited on PET was determined using high-resolution X-ray diffractometer system (model: Panalytical X'Pert PRO MRD PW3040). The sheet resistance and the resistivity of the Mo deposited on PET were measured with a four-point probe (Model: Changmin Tech CMT-SR2000N).

RESULTS AND DISCUSSION

The results of films of molybdenum deposited on PET are characterized by different techniques. The samples used in the experiment consisted of a thin film of molybdenum and PET. The measured values of films thicknesses range from 49 nm to 210 nm. The thicknesses of these samples are given in Table 1.

Structural Characterization

AFM images for surface morphologies of molybdenum deposited on PET are given in Figure 1(a–e). Figure 1 shows the variation of the film roughness with thickness which correlated with deposition time. The surfaces of the product Mo thin films obviously were smooth. The evaluated root mean square (rms) surface roughness of the films, were 9.35 nm and 16.79 nm for films with a thickness of 49 nm and 210 nm, respectively. These results indicate that the surface quality of Mo on PET thin films improves with a decrease of the film thickness. In all cases conical features clearly seen on the film surface smoothness is a highly desired parameter for the coatings that are used for optical applications in order to reduce the reflection loss due to roughness-induced surface scattering.

From the X-ray diffraction spectra, the intensity of the main peaks is determined as a quantitative measure of the crystallinity. Thus, for Mo films on the PET substrate, a broad diffraction peak corresponding to the PET substrate was observed at 2θ angle 26° . Mo belonging to the cubic system with

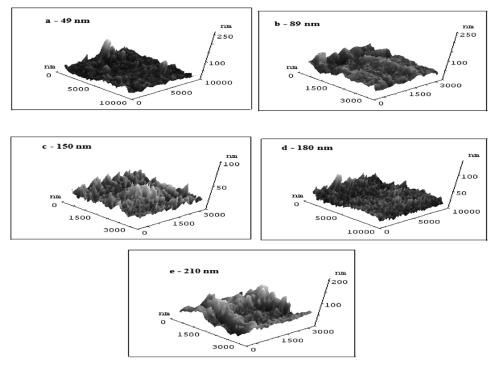


Figure 1: Variation of the Mo films roughness with thickness.

the (1 1 0) preferred orientation has been identified, as shown in Figure 2. This orientation is the most commonly presented in the literature [5]. The vertical lines below indicate the corresponding reflection peaks for Mo thin films. Figure 3 showed that the chemical components of the products were Mo, O. The presence of the oxygen small peak in the spectrum is due to the PET substrate.

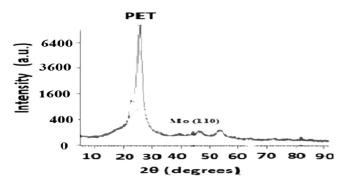


Figure 2: XRD spectrum of Mo on PET substrates.

626 M. G. Faraj et al.

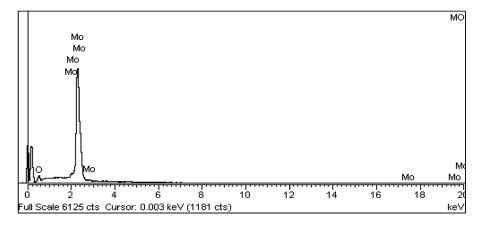


Figure 3: EDX of Mo deposited on PET.

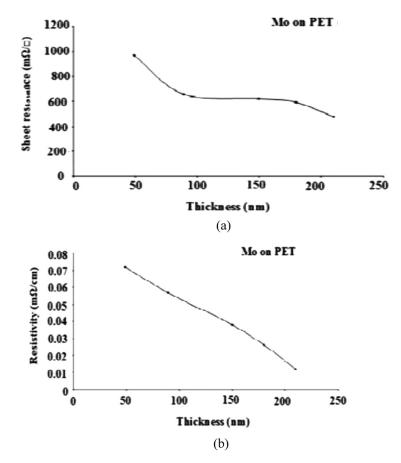


Figure 4: Variation thickness with (a) sheet resistance and (b) resistivity.

Electrical Characterization

Figure 4 shows results indicating that increasing the film thickness can also reduce the resistivity and sheet resistance of the Mo on PET films. This behavior could also be attributed to an increase of the carrier mobility induced by increasing the grain size. The lowest resistivity value we found for the Mo films was $1.3 \times 10^{-5} \Omega \cdot cm$ at thickness (210 nm).

CONCLUSION

Mo thin films have been deposited onto PET substrates by using DC sputtering kept at room temperature. The thickness of the films varied from 49 nm to 210 nm by changing deposition times. From AFM images it was found that the root mean square roughness of film surface increased as the film thickness increased. The variations of electrical parameters such as the resistivity and sheet resistance with film thickness are correlated to the changes in the films microstructure.

REFERENCES

- Park, J. H., Kim, D. Y., Ko, J. K., Chakrabarty, K., and Yi, J. *Thin Solid Films* 427, 303–308 (2003).
- [2] Obata, T., Sato, K., Chiba, M., Mohri, M., Yamashina, T., and Yabe, K. *Thin Solid Films* 87, 207 (1982).
- [3] Guillén, C., and Herrero, J. Journal of Materials Processing Technology 143, 144 (2003).
- [4] Gordillo, G., Grizalez, M., Hernandez, L. Solar Energy Materials and Solar Cells 51, 327 (1998).
- [5] Martinez, M. A., and Guillen, C. Surface and Coatings Technology 110, 62 (1998).
- [6] Scofield, J. H., Duda, A., and Albin, D. Thin Solid Films, 260, 26 (1995).
- [7] Martínez, M. A., and Guillén, C. Materials Processing Technology 143, 326 (2003).
- [8] Orgassa, K., Schock, H. W., and Werner, J. H. Thin Solid Films 431-432, 387 (2003).
- [9] Ajji, A., and Chapleau, N. Journal of Materials Science 37, 3893 (2002).
- [10] Bach, C., Dauchy1, X., Etienne, S. Materials Science and Engineering 5, 012005 (2009).