

Growth of molybdenum sulphide ultra-thin films and their characterization



G. Papadimitropoulos^{1,*}, A. Balliou¹, D. Davazoglou¹ and D. N. Kouvatsos¹

¹Institute of Nanoscience and Nanotechnology, NCSR "Demokritos", 153 10 Aghia Paraskevi, Attiki, Greece

*E-mail: g.papadimitropoulos@inn.demokritos.gr

Abstract

In this work a simple method for depositing molybdenum sulphide films, with thicknesses from a few monolayers to some nanometers, is described. The grown films were amorphous and deposited very easily by heating a molybdenum wire under vacuum at low filament temperature (400 °C) in an ambient composed by H₂ and H₂S. The deposited films were subjected to thermal and microwave annealing in order to be crystallized and were characterized with various techniques to investigate their morphological and structural properties.

MoS₂ film deposition

The films are deposited very easily compared to other methods by a heated molybdenum filament, covered by a native oxide, at low pressure (1,5 Torr).





- MoO_3 (native oxide) yields MoS_2 according to the reaction: $2\text{MoO}_3 + 4\text{H}_2\text{S} \rightarrow 2\text{MoS}_2 + 4\text{H}_2\text{O} + \text{O}_2 \qquad \text{T}_{\text{filament}} \ge 400^{\circ}\text{C}$
- The rotated shutter system enables the deposition of MoS₂ ultra thin films with thicknesses from a few monolayers to some nanometers.
- The samples remained at room temperature rendering the deposition method suitable for using organic and plastic substrates, too.

(nm/sec) Growth Rate 2

The rotated shutter system.



MoS₂ film characterization



SEM micrographs and EDXS spectra of







AFM images of a MoS_2 film with a roughness of 1 nm.



Summarizing, a simple method for depositing MoS₂ ultra thin films at room temperature was presented. The as-deposited films were amorphous and were characterized with respect to their morphology and chemical composition. As shown by SEM and AFM measurements, the obtained samples are smooth and homogeneous. At variance with conventional thermal treatments, microwave annealing yields MoS_2 films exhibiting a better atomic ordering as shown by XRD and Raman measurements. The rotated shutter system contributes to the control of film thickness from a few monolayers to some nanometers.

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