

# Synthesis and Characterization of Uniformly-Aligned MoO<sub>3</sub> Nanobelts

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**Abstract**—We have been synthesized MoO<sub>3</sub> nanobelts by thermal chemical vapor deposition. The growth products will be studied by field emission scanning electron microscope (FESEM), X-ray photoemission (XPS), Raman Spectroscopy, X-rays diffraction (XRD). SEM images showed the belts-like nanostructures. XPS and Raman shift showed the consisting of MoO<sub>3</sub> structure. XRD patterns indicated orthorhombic MoO<sub>3</sub> structures phase of planes (020), (110), (040), (021), (060) and (010).

**Keywords**—MoO<sub>3</sub> nanobelts; thermal chemical vapor deposition; raman spectroscopy; XPS; XRD

## I. INTRODUCTION

Molybdenum trioxide (MoO<sub>3</sub>), a wide band gap n-type semiconductor is interested due to its structures crystal [1-3]. It was used in many applications such as hydrogen evolution [4], field emission [5], thin film transistor [6], capacitor [7] and gas sensor [8]. MoO<sub>3</sub> nanostructures, belts-like, wires-like, rods-like, nanodiscs, nanoflowers and nanosheets have been synthesized from many processes [2-5, 7-9]. MoO<sub>3</sub> nanowires with uniform in side can be formed on silicon substrates without using any catalyst by providing in two methods, thermal evaporation and oxidation [5]. The ordering MoO<sub>3</sub> nanosheets have been prepared via hydrothermal route using MoO<sub>3</sub> powder and hydroquinone as starting materials, for studying the conductivity mechanism of complex impedance and modulus formalism [8]. MoO<sub>3</sub> nanorods have been prepared using a simple ultrasonic method, for gas sensing high response to NO<sub>2</sub> [9]. Recently, the MoO<sub>3</sub> nanobelts can be synthesized by using ammonium molybdate tetrahydrate mixed with ethylene glycol as a starting solution, centrifugation and sintering of the precipitate, the result showed the  $\alpha$ -MoO<sub>3</sub> nanobelts grew with a strongly preferred orientation [10]. In this paper, the MoO<sub>3</sub> nanobelts have been synthesized using MoS<sub>2</sub> powder as starting materials by chemical thermal vapor deposition.

## II. EXPERIMENTAL PROCEDURE

The mixing of MoS<sub>2</sub> and S powder (SIGMA-ALDRICH) and SiO<sub>2</sub>/Si substrate were put in the alumina boat. The alumina boat was placed on the middle of furnace quartz tube and heated at 700 °C under mixing atmosphere of N<sub>2</sub> and O<sub>2</sub> gas with flow rate of 6 sccm for 1 h. The furnace quartz tube was cooled down to room temperature, nationally. Then, the

alumina boat and substrate were taken out. We can observe the different color of formed products on substrate. The heated substrate was investigated by optical microscope (OM), field emission scanning electron microscope (FESEM), X-ray photoemission (XPS), Raman spectroscopy, atomic force microscope (AFM) and X-ray diffraction (XRD).

## III. RESULTS AND DISCUSSIONS

After, the furnace tube was cooled down to room temperature, nationally. The formed products on substrate were characterization by OM, FESEM, XPS, Raman spectroscopy, AFM and XRD. The results show as follow.

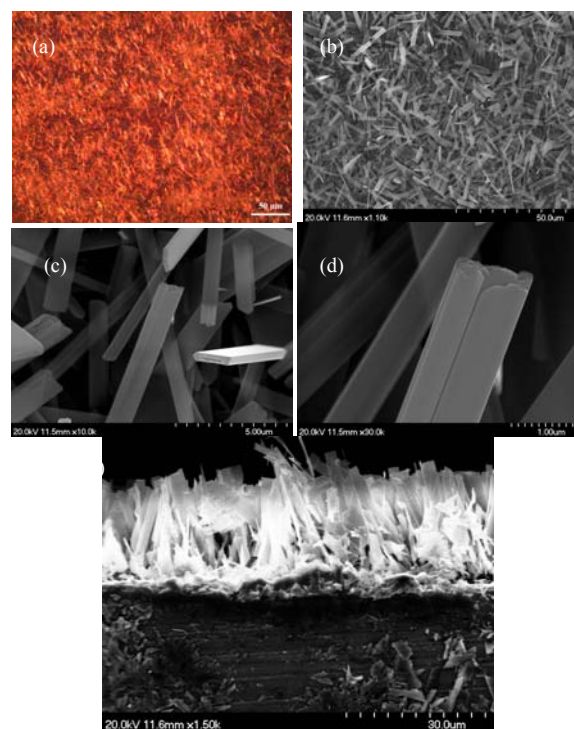


FIGURE 1. OPTICAL MICROSCOPE (a), FESEM IMAGES OF MoO<sub>3</sub> NANOBELTS (b-d) AND CROSS-SECTION (e)

The figure 1 show the pictures of OM, FESEM and cross-section of substrate. We can see belts-like of formed products with length of more a few micrometers and width of around

200-500 nanometers. The synthesized materials grew from the bottom of substrate with height of about 21.7 micrometers. The XPS patterns showed the elements consisting of MoO<sub>3</sub> nanobelts.

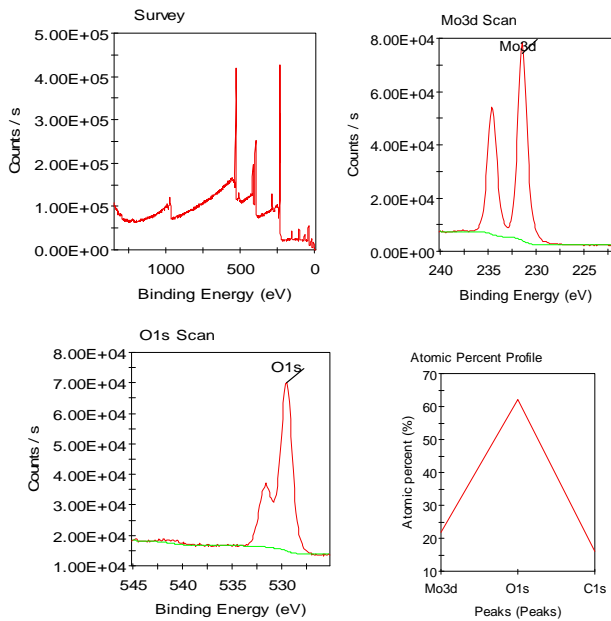


FIGURE II. XPS PATTERNS OF MoO<sub>3</sub> NANOBELTS

The XPS curves in Figure 2 revealed the elements consisting of prepared MoO<sub>3</sub> nanobelts with Mo and O atoms, the S atom no appear.

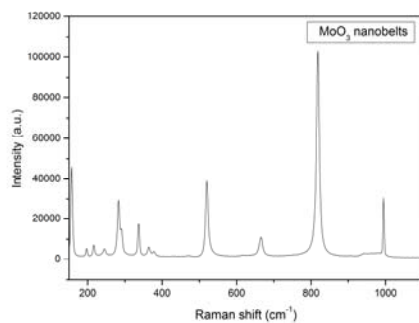


FIGURE III. RAMAN SHIFT OF MoO<sub>3</sub> NANOBELTS

The operating of Raman spectroscopy of MoO<sub>3</sub> nanobelts-like showed principal peaks of 994.2, 818.7, 666.4 and 284.5 cm<sup>-1</sup>, respectively, and we can observed the interference peak of SiO<sub>2</sub> of SiO<sub>2</sub>/Si substrate around 525 cm<sup>-1</sup>.

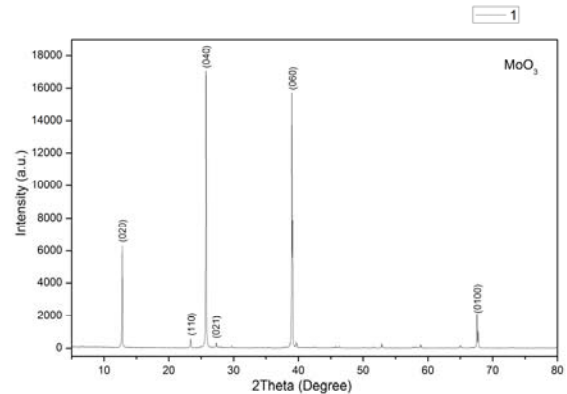


FIGURE IV. XRD PATTERNS OF MoO<sub>3</sub>

Figure 4 showed XRD patterns of MoO<sub>3</sub> nanobelts with plane of (020), (110), (040), (021), (060) and (010). All of peaks correspond with MoO<sub>3</sub> (JCPDS reference card no. 05-0508) [11].

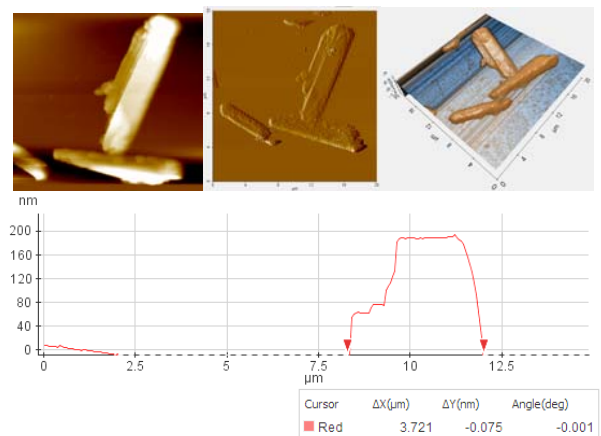


FIGURE V. AFM IMAGES OF SELECTED ANY MoO<sub>3</sub> NANOBELTS

The AFM images in FIGURE 5 show some selected MoO<sub>3</sub> nanobelts. This image reveals the morphology side of formed materials on substrate. The thickness of MoO<sub>3</sub> nanobelts are about 20-60 nanometer.

#### IV. CONCLUSION

In summary, The MoO<sub>3</sub> nanobelts can be synthesized by thermal chemical vapor deposition. The XRD, Raman shift and XPS were confirming that MoO<sub>3</sub> nanobelts as-synthesized, and OM, FE-SEM and AFM images showed the length, width and thickness of MoO<sub>3</sub> nanobelts with few micrometers, 200-500 and 20-60 nanometers, respectively.

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