



WO₃ NANOWIRE BASED DIODE FOR ULTRA VIOLET LIGHT SENSING APPLICATIONS

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ABSTRACT

In this paper, we report a simple approach to fabricate UV photo-detector based on WO₃ nanowires. The WO₃ nanowires were synthesized by a simple hydrothermal method using tungsten powder as a precursor, Palladium nanoparticles as catalytic to remove excess of H₂O₂ and Na₂SO₄ as a structure-directing agent. The structural properties of the WO₃ nanowires are analyzed by the UV-spectroscopy, XRD and SEM. XRD and SEM results indicated that the WO₃ nanowires are hexagonal. Device has been made by using spray coating method keeping substrate on hot-plate and Palladium contacts has been deposited on film by using thermal evaporation method. After fabrication of the device, device has been characterized for electrical properties in dark condition and in the presence of UV light with UV illumination and Agilent B1500A Semiconductor device Analyzer.

Keywords: Tungsten Oxide nanowires, UV Photo-detector and sensor.

I. INTRODUCTION

Among the transition metal oxides, tungsten oxide has drawn great attention due to its distinctive physical and chemical properties and therefore has been extensively investigated for applications like electrochromism, photochromism, gas sensing, photo catalysis, photoluminescence, and many more [1-7]. Tungsten oxide has been used to construct photochromic smart-windows, flat panel displays, optical modulation devices, gas sensors, humidity and temperature sensors. Tungsten oxide is an indirect, wide-gap semiconductor. Upon bandgap photoirradiation, electron-hole pairs are produced. Depending on the behavior of the newly photogenerated electrons and holes, the optical absorption of tungsten oxide can be altered, leading to a color change from transparent to blue. It is well known that

nanostructures have unique physical and chemical properties and can be used as elementary units of optoelectronic devices. Tungsten oxide (WO₃) nanostructures have outstanding properties which make it valuable as functional building blocks for the fabrication of biological sensors, optoelectronic devices, optical memories and other areas [1].

Photo-detectors convert incoming optical energy into electrical energy. Si-based UV photo-detectors can be very sensitive in UV region with low noise and quick response, however it has some limitations, such as lower efficiency or the need of an ultra high vacuum environment and a very high voltage supply. To avoid these disadvantages UV photo-detectors based on wide band gap semiconductors have received more attention. ZnO based photodetectors are very popular.

The UV detector based on polycrystalline ZnO thin film shows low photoresponsivity and long response time of the order of few minutes. ZnO/Porous-silicon photovoltaic UV detectors have been demonstrated to be ultra fast and have high responsivity [8]. Near ideal characteristic ZnO thin films have been characterised for UV detector applications [9]. WO₃ is also a potential material for detecting UV radiation because it has large energy band-gap [2,3]. One dimensional (1D) nanostructure wide band-gap semi-conductors (e.g. WO₃), have high responsivity, high thermal stability and high speed. So WO₃ nanowires can be used as a sensing element for UV photo-detectors as it has a very high photoconductivity gain. UV photo-detectors are highly suitable for a wide range of industrial, military, environmental and biological applications [4].

II. EXPERIMENTAL

In experiment two steps has been followed for making the tungsten oxide nanowires. In first step WO₃ sol was prepared whereas in second step WO₃ nanowires was obtained from WO₃ sol. For preparation of WO₃ sol, Tungsten powder was dissolved into hydrogen peroxide. The aqueous solution was kept in an ice water bath under slight stirring [5]. After centrifugation, a transparent solution was formed. A small amount of Palladium nanoparticles were added to remove the excess of H₂O₂. A homogeneous and stable WO₃ sol was obtained by adding distilled water and oxalic acid to the mixture and adjusting the pH value of the solution to 1.1 by using the NaOH solution (1 mol per liter).

In second step WO₃ sol was transferred into an autoclave, and then Na₂SO₄ was added in the sol and autoclave was maintained at 180°C for 24 hours. Product was washed with ethanol and water to remove the ions and dried at 60°C and hexagonal WO₃ (h-WO₃) nanowires were obtained [6]. These nanowires were coated on p-type silicon substrate by using spray coating method keeping substrate on hot-plate. After that, annealing was done at

500°C in presence of Nitrogen. At last Palladium contacts were deposited on film by using thermal evaporation method with 12A 4 Hind hivac vacuum coating unit.

III. RESULTS AND DISCUSSION

III.1. Structural Analysis

Figure 1 a-b shows the scanning electron microscope (SEM) images of as prepared WO₃ nanowires synthesized using the hydrothermal method at different magnifications. The few nano meter wire-like structure is clearly seen uniformly across the specimen. Figure 2 shows the XRD pattern of the as-synthesized product. It can be indexed to the hexagonal phase of the WO₃ nanowires with the lattice constants $a=7.324 \text{ \AA}$, $c=7.662 \text{ \AA}$ (JCPDS 85-2459).

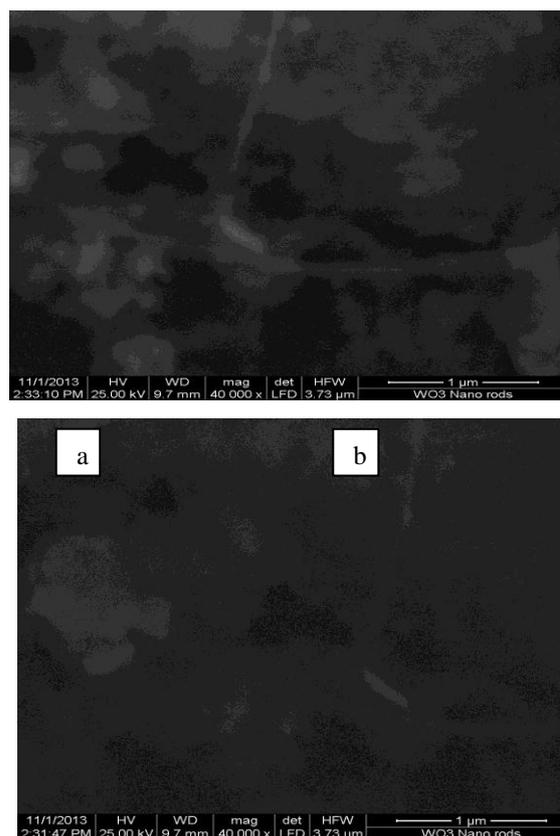


Figure 1: SEM images of synthesized WO₃ nanowires

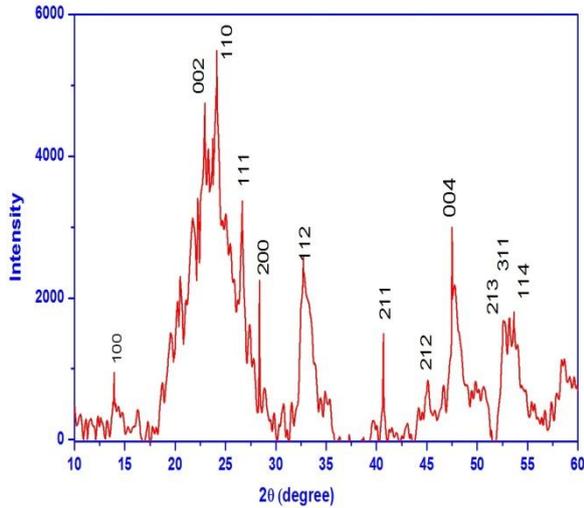


Figure 2: XRD pattern of WO₃ nanowires.

III.2. Optical and Electrical Analysis

Figure 3 shows the UV absorption spectrum of as-prepared WO₃ nanowires. Energy band gap can be found by using formula

$$\alpha h\nu = a(h\nu - E_g)^r \dots\dots(1)$$

(where α is the absorption coefficient, a is a constant which does not depend on the photon energy and r is the numerical constant which has the value of $1/2$), the band gap varies as the absorbance (α^2) [7]. Figure 4 shows the plot of $(\alpha h\nu)^2$ versus $h\nu$. The intercept with the $h\nu$ axis of the tangent to the curve of $(\alpha h\nu)^2$ plotted against $h\nu$ gives a good approximation of the energy band gap (i.e. E_g is~ 4.17 eV), seen from the figure 4

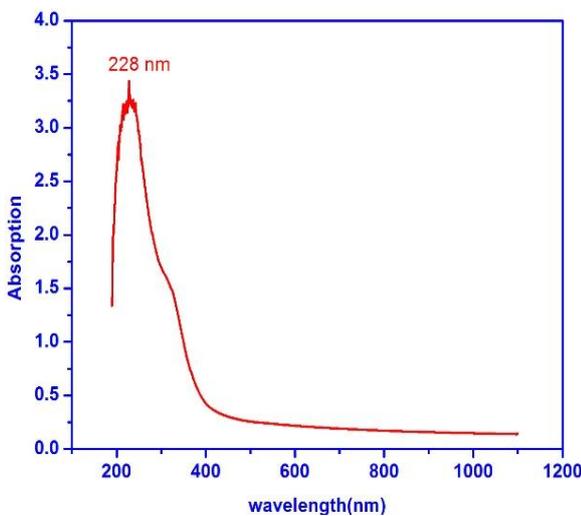


Figure 3: WO₃ nanowire UV absorption spectrum

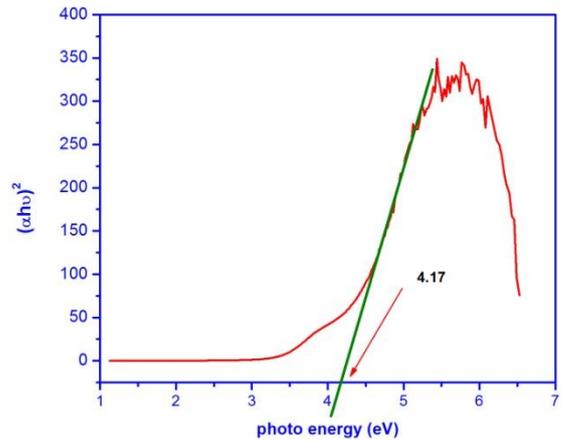


Figure 4: Plot of $(\alpha h\nu)^2$ versus photon energy ($h\nu$) .

The I-V characterization was performed with UV illumination and an Agilent B1500A Semiconductor Device Analyzer. Figure 5 shows the rectifying nature of the device in both the condition i.e. dark condition and illuminated condition. From the I-V characteristic shown in figure 5 it is obtained that the forward current increases after the exposure of the light. Under UV light illumination, electron-hole (e-h) pairs are generated in WO₃ film and holes are migrated to the WO₃ surface which then became neutralized by reacting with the oxygen ions. In this manner, the unpaired electron density in WO₃ film increases, which is responsible for an increase in forward and reverse current under UV light illumination. I-V curves are obtained when the nanodevices was exposed to the UV light with wavelength 365 nm at a constant light intensity of 650 μ W. At 2 V photo current was 47.5 μ A and dark current was 16 μ A.

Figure 6 is the plot of current density versus voltage. Current density can be found using formula,

$$current\ density = \frac{current}{area}$$

Diameter of the contact was 2 mm.

Responsivity(R) of the photo-detector relates the electric current flowing in the device to the

incident optical power. R has units of A/W and It is expressed as,

$$R = \frac{I_P}{P} \dots\dots\dots(2)$$

where I_P is the photocurrent, and P is the intensity of the optical power.

Responsivity of the device is .0730 A/W at wavelength of 365 nm and with light intensity 650 μ W.

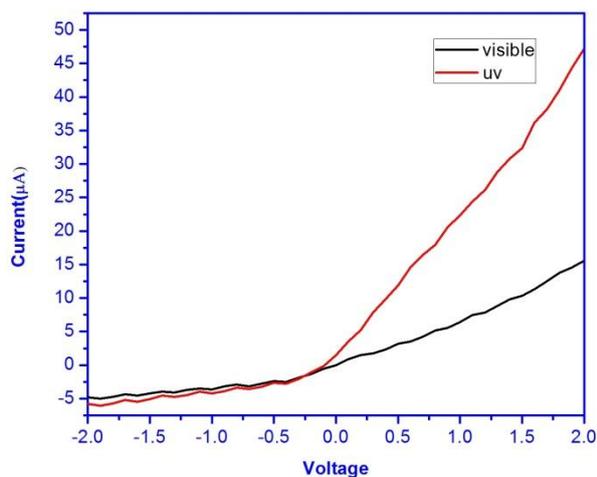


Figure 5: I-V Characteristics of UV Photo-detector.

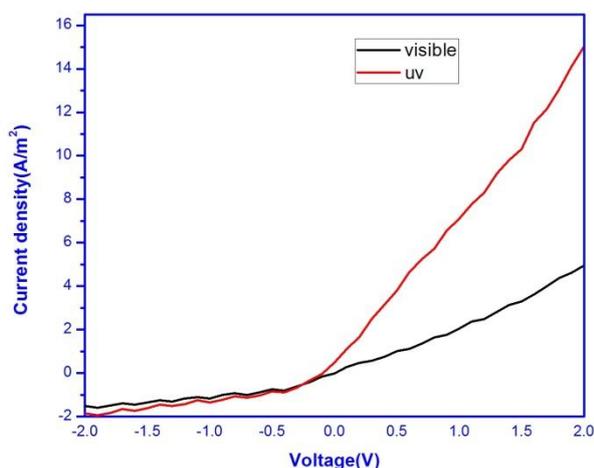


Figure 6: Current density versus Voltage plot of UV Photo-detector.

IV. CONCLUSION

WO₃ nanowires have been synthesized using hydrothermal method. This WO₃ nanowires based UV photo-sensor is very sensitive to UV illumination; an increase in current is obtained under UV illumination at 365 nm with optical

power of 650 μ W. Adsorption of oxygen molecules on the surface of the WO₃ nanowires has a significant influence.

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