Synthesis and Characteristics of SnO$_2$ Nanorods on Pd-Coated Substrates

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We have fabricated tin oxide (SnO$_2$) nanorods on palladium-coated substrates by carrying out the thermal evaporation of solid Sn powders. We have employed X-ray diffraction, scanning electron microscope, transmission electron microscope and photoluminescence (PL) spectroscopy to characterize the synthesized products. The obtained 1D nanomaterials with a jagged surface were SnO$_2$ with rutile structure. PL spectra exhibited visible light emission.

I. INTRODUCTION

Tin oxide (SnO$_2$) is an important metal oxide. As a transparent conducting oxide, it has a long history of use as an electronic conductor, primarily in the form of thin films. In addition, the material has received a great deal of attention due to its applications in optoelectronic devices including flat panel displays and solar energy cells [1].

Inorganic materials with different morphologies and size can exhibit different properties [2], even if they are made up of the same elements. Accordingly, various structural and morphological forms of SnO$_2$ materials have been fabricated over the past several years, including nanowires [3,4], nanoribbons or nanobelts [5-8], nanorods [9], nanotubes [10], nanodiskettes [11], and nanocomposites [12,13].

Although carbon nanotubes have been synthesized using the palladium (Pd) catalyst [14-16], to our knowledge, synthesis of any inorganic nanomaterial including SnO$_2$, on the Pd substrate has not been reported to date. Furthermore, Pd is a platinum-group metal with good corrosion resistance and has been studied as the VLSI materials [17]. In this paper, we report a thermal evaporation approach, by which we have produced the SnO$_2$ nanorods on Pd-coated Si substrates. We have investigated the samples with respect to their structural and photoluminescence (PL) characteristics.

II. EXPERIMENTAL

Pure Sn powders were placed in an alumina boat, which was put in the middle of quartz tube inserted in a horizontal tube furnace. We used Si(001) as a starting material onto which a layer of Pd was deposited. On top of the boat, a piece of the substrate was placed with the Pd-coated side downwards. The vertical distance between the alumina boat and the substrate was approximately 10 mm. After evacuating the wards, the vertical distance between the alumina boat and the substrate was approximately 10 mm. After evacuating the furnace for analysis. A white layer was found on the surface of the substrate.

Glancing angle X-ray diffraction (XRD: CuK$_\alpha_1$ radiation) patterns were obtained on a Philips X’pert MRD diffractometer with an incidence angle of 0.5$^\circ$. Scanning electron microscopy (SEM) studies were carried out on a Hitachi S-4200. Further structural analysis using transmission electron microscopy (TEM) was performed on a Philips CM-200. For TEM observation, the products were ultrasonically dispersed in acetone, and then a drop of the suspension was placed on amorphous carbon films supported by copper grids and dried in air. PL spectrum was measured at room temperature by a 325 nm He-Cd laser (Kimon, 1K, Japan).

III. RESULTS AND DISCUSSION

Figure 1a shows the typical side-view SEM image of the product, indicating that this wool-like material consists of aggregates of one-dimensional (1D) nanostructures. The SEM image reveals that the growth direction of the nanostructure is randomized. Fig. 1b shows the SEM image of the products with a closer view, revealing that the product comprises the straight, bent, and crooked rod-like structures. Statistical analysis of many SEM images shows that average diameter of these nanostructures varies from 30 to 300 nm and length reaches up to several tens of micrometers. The upper right inset in Fig. 1b reveals that no catalyst particle can be seen at tips of the 1D structures. Fig. 1c shows the high magnification SEM image enlarging an area of Fig. 1b. The jagged-edged or saw-like structure is apparent. Although similar structure has been observed in the case of MgO [18], to the best of our knowledge, no similar SnO$_2$ structure was reported in the literature previously.

Figure 2 shows a typical XRD pattern recorded from the products. The Miller indices are indicated on each diffraction peak. The diffraction peaks of the (110), (101), (200), (211), (220), (002), (310), (112), (301), (202), and (321) planes can be readily indexed to the tetragonal rutile structure of SnO$_2$ with lattice constants of $a=4.738\,\text{Å}$ and $c=3.187\,\text{Å}$ (JCPDS File No. 41-1445). No obvious reflection peaks from the impurities, such as unreacted Sn or other tin oxides, were detected, indicating the high purity of the products. In the XRD measurements, the angle of the incident beam to the substrate surface was approximately 0.5$^\circ$, and the detector was rotated to scan the samples. Therefore, we surmise that the peaks are mainly from the products.

TEM analysis of individual nanostructures provide further insight into the structure of these materials. Low magnifica-
FIG. 1: (a) Low magnification side-view and (b) closer-view SEM images of the products. Inset shows the end of a nanostructure. (c) High magnification SEM image.

FIG. 2: XRD pattern of the products.

higher oxygen pressure of 1000 mTorr (compared to the air pressure of 150 mTorr in the present study), we obtained the cluster-like structures without any thin nanorod. The variation of SnO\(_2\) structural morphology with O\(_2\) content indicates that O\(_2\) gas plays an important role in controlling nucleation and growth of SnO\(_2\) nanorods. It is generally agreed that the SnO(g), which forms at the initial stage of oxidation of Sn (Sn(s) + 1/2O\(_2\) (g) → SnO(g)), is metastable and decomposes according to the reaction [20]: 2SnO(g) → SnO\(_2\)(s) + Sn(l). The additional oxygen may facilitate large supersaturation of O\(_2\) gas and thus SnO gas in the gaseous state, resulting in fast condensation. Under low supersaturation, narrow 1D structures are easy to grow. Under high supersaturation, however, it may activate the secondary growth sites and heterogeneous nucleation on the side of the 1D structures, tending to produce the thicker structures. In the present synthesis route, while elevating the temperature with no gas flowing, we surmise that Pd layer stays pure or preferentially transforms to stable Pd silicide compounds. Therefore, it may provide negligible amount of additional oxygen to the growing SnO\(_2\) nanostructures at the synthesis temperature of 900°F, keeping the thin 1D nanostructures from being thicker. Our experiments under the same condition using other substrate materials which comprise oxygen, resulted in the generation of the cluster-like or thick structures, agreeing with the above speculation. Efforts are now in progress to pursue these routes in more detail to reveal the synthesis mechanism.

Room-temperature PL spectra of the as-synthesized products were measured and shown in Fig. 4. Visible emissions with a peak wavelength position of around 591 nm (corresponding to 2.10 eV) is dominantly observed. The visible light emission is known to be related to defect levels within the band gap of SnO\(_2\), associated with O vacancies or Sn interstitials that have formed during the synthesis process [21-23]. A similar emission has been reported in the case of SnO\(_2\) nanoribbons synthesized by laser ablation [21] and SnO\(_2\) nanorods synthesized by solution phase growth [22].
IV. CONCLUSIONS

In summary, we have demonstrated the production of SnO$_2$ nanorods on Pd-coated substrate by heating Sn powders at 900°C. SEM image and TEM analysis coincidentally indicate that the product consists of nanorods with a jagged surface. The results of XRD, TEM and SAED pattern indicate that the SnO$_2$ nanorods have a tetragonal rutile-type crystalline structure. We speculate about the growth mechanism and need deep study in the future. The PL measurement shows an apparent visible light emission band centered at around 591 nm. Compared with SnO$_2$ nanostructures reported previously, the SnO$_2$ nanorods in the present study exhibit different structural characteristics, which may lead to novel applications in miniaturized devices.

FIG. 4: Room temperature PL spectrum with an excitation wavelength at 325 nm.

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