Nanofabrication and properties of the highly oriented carbon nanocones

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Abstract

A process to fabricate Co-assisted carbon nanocones (CNCs) was successfully developed by using microwave plasma chemical vapor deposition (MP-CVD). The nanostructures could be manipulated by adjusting the ratios of the source gases (CH4/H2=5/80–15/80 sccm/sccm) and the substrate bias (0~150 V). The properties of deposited nanostructures were characterized by field emission scanning electron microscopy (FESEM), Raman Spectroscopy, auger electron spectroscopy (AES) spectra. The results show that the nanostructures with cone shape are resulted from competition between the ion bombardment rate and the growing speed of the nanostructures in the plasma, where ion bombardment is drove by the presence of negative substrate bias. In other words, a higher substrate negative bias (~150 V) and a lower concentration of carbon species in the plasma are favorable conditions to grow highly oriented CNCs due to a greater ion bombardment energy and lower lateral growth rate of the nanostructures, respectively. The results indicate that CNCs synthesized under the applied bias of ~300 V show an excellent performance in field emission current density (up to ~173 mA/cm² at 10 V/µm).

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Keywords: Microwave plasma (MP); Chemical vapor deposition (CVD); Carbon nanocones (CNCs); Field emission (FE)

1. Introduction

Applications of carbon nanostructures, e.g. carbon nanotubes (CNTs), carbon nanocones (CNCs), have been studied for many years due to their higher aspect ratio at the tips, excellent mechanical properties and chemical stability, especially for field emission display (FED) devices. Samsung [1] had successfully fabricated a prototype of 4.5 in. full-color CNT-field emission display with brightness reaching 1800 cd/cm² at electrical field of 3.7 V/µm. This amazing result has attracted much attention. Many researches are focused on developing a self-assembly process to fabricate the carbon nanostructures with sharp apex angle to enhance their field emission properties, e.g. carbon nanocones (CNCs) [2–8]. Many reports indicate that the turn on electric field of CNCs can be as low as 0.1–1.38 V/µm due to a larger field enhancement factor β, which are much better than CNTs [9–12]. Furthermore, the cone-shape structure provides higher mechanical and thermal stability than a narrow cylinder structure of CNTs [13]. It was reported that the CNCs can be synthesized under low substrate temperatures (~200 °C) [14], implying its IC compatible fabrication process due to lower temperatures, which are much lower than the temperatures of ~600 °C for CNTs growth. Hence, the CNCs have demonstrated a greater potential and practicability to realize the applications in electronic devices.

In this paper, we report a process to fabricate high-oriented CNCs and study their nanostructures, properties and growth mechanisms. Their field emission properties were emphasized.

2. Experimental

The high-oriented CNCs were synthesized on Si wafer by using catalyst-assisted microwave plasma chemical vapor deposition (MP-CVD) with H2 and CH4 as source gases.
Cobalt film with thickness of 10 nm was first deposited on Si wafer by physical vapor deposition (PVD) to act as catalyst. The H-plasma (H₂ flow rate=100 sccm) was utilized for 10 min to activate the Co film to become the well-distributed nano-particles with size distribution between 40 and 95 nm. Then, the wafer with nano-particles on the surface was used to synthesize the carbon nanostructures by MP-CVD at about 600 °C temperature for 10 min and varying H₂/CH₄ ratio and substrate bias. The process parameters and conditions of CNCs are listed in Table 1.

The nanostructures of the deposited CNCs were characterized by field emission scanning electron microscopy (FESEM), transmission electron microscopy (TEM) and high resolution transmission electron microscopy (HRTEM). The bonding structure of the deposited CNCs was studied by Raman spectroscopy (Jobin Yvon LAB-RAM HR) with 632.8 nm He–Ne laser. Auger electron spectroscopy (AES) (VG Microlab 310F with a Schottky field emission electron source) was used for chemical analysis of CNCs. Field emission properties of as-growth carbon nanostructures were analyzed by $I–V$ detector (Keithley 237), and the distance between anode and cathode (CNTs) was set at about 100 μm. The lifetime test of field emitters was examined by current versus time.

### Table 1

<table>
<thead>
<tr>
<th>Specimen designation</th>
<th>Ratio of H₂/CH₄ (sccm/sccm)</th>
<th>Substrate temperature (°C)</th>
<th>Deposition time (min)</th>
<th>Bias (V)</th>
</tr>
</thead>
<tbody>
<tr>
<td>B1</td>
<td>80/5</td>
<td>624</td>
<td>10</td>
<td>0</td>
</tr>
<tr>
<td>B2</td>
<td>644</td>
<td>644</td>
<td>50</td>
<td></td>
</tr>
<tr>
<td>B3</td>
<td>637</td>
<td>640</td>
<td>150</td>
<td></td>
</tr>
<tr>
<td>B4</td>
<td>640</td>
<td>646</td>
<td>200</td>
<td></td>
</tr>
<tr>
<td>B5</td>
<td>646</td>
<td>645</td>
<td>300</td>
<td></td>
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<tr>
<td>B6</td>
<td>80/1</td>
<td>645</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>B7</td>
<td>80/5</td>
<td>640</td>
<td>50</td>
<td></td>
</tr>
<tr>
<td>B8</td>
<td>80/10</td>
<td>622</td>
<td>150</td>
<td></td>
</tr>
<tr>
<td>B9</td>
<td>80/15</td>
<td>617</td>
<td>200</td>
<td></td>
</tr>
</tbody>
</table>

Substrate: 10 nm cobalt-coated silicon [(100) P-type] wafer. Other CNCs deposition conditions: pressure, 9 Torr; microwave power, 800 W.

Fig. 1. Typical SEM micrographs of the as-grown CNCs synthesized under different negative substrate bias: (a) 0 V, (b) –50 V, (c) –150 V, (d) –200 V and (e) –300 V (ref. to Table 1).
(I–t curve) measurement, where 900 V bias was applied during the 3600 s of operation.

3. Results and discussion

3.1. Effects of applied bias and H$_2$/CH$_4$ ratio on the growth of CNCs

The effect of the substrate bias on carbon nanostructures was explored by varying bias from 0 to $-300$ V under the same H$_2$/CH$_4$ ratio ($=80/5$ sccm/sccm) of the gas sources. The SEM morphologies of the as-grown nanostructures under five different bias values are shown in Fig. 1(a) to (e), respectively. It indicates that the nanostructures gradually change from polyhedral carbon clusters to needle-like CNCs by increasing the negative substrate bias, and become the well-aligned CNCs at bias $>-150$ V (Specimen B4 and B5). In other words, the substrate bias is one of the key parameters to form the aligned CNCs. Besides, the average height of the CNCs increases from 20–80 to 500 nm as the applied bias increasing from $-150$ to $-300$ V. It implies that a greater negative substrate bias can enhance the growth rate of CNCs perpendicular to the substrate. Furthermore, higher substrate bias may promote to form...
the CNCs with sharper tip due to the effect of ions bombardment. This phenomenon is in agreement with the observation of the smaller apex angle at the tips of CNCs, where the apex angles are 15–20° and 9–12° under –200 V (specimen B4) and –300 V (specimen B5) applied bias, respectively.

The SEM morphologies of the as-grown CNCs synthesized under different H₂/CH₄ ratios of 80/1, 80/5, 80/10, and 80/15 sccm/sccm are shown in Fig. 2(a) to (d), respectively. It indicates the CNCs with a greater average apex angle and larger bottom diameters are deposited under a lower H₂/CH₄ ratio in gas sources. In other words, the higher concentration of CH₄ gas may lead to an increase of the lateral growth rate of CNCs to form a blunt apex angle. The results imply that the nanostructures of CNCs are determined essentially by the competition among etching rate of plasma species, the lateral growth rate and the upward deposition rate of carbon along the surface or through the interior of the catalysts. It is interesting to note that CH₄ concentration adopted to the as-grown CNCs in these cases are between the concentrations for growth of diamond film and CNTs, where diamond film deposition require a lower concentration (H₂/CH₄=~100/5 sccm/sccm) [15,16]. It is concluded that highly oriented CNCs are more likely to be formed under the appropriate concentration of carbon species in the plasma and higher negative substrate bias.

3.2. Raman and AES analyses of the as-grown CNCs

The Raman spectrum of the as-grown CNCs is shown in Fig. 3 (Specimen B5). The peak position of the D band and G band are around 1334 and 1598 cm⁻¹, respectively, where the \( \frac{I_G}{I_D} \) ratio is ~ 0.81. Huang’s group proposed that a shift of the E₂g G mode from its normal value of 1580 to 1598 cm⁻¹ is an indication of the existence of nanocrystalline graphite or sp² clusters [17–19]. The peak at 1334 cm⁻¹ is assigned as A₁g mode due to the presence of aromatic rings in disordered graphite rather than diamond peak at 1332 cm⁻¹. Therefore, the bonding structure of the as-grown CNCs can be considered as sp² clusters in an amorphous carbon (a:C) matrix.

The AES spectra and the corresponding SEM image of the as-grown CNCs at the tip (P1) and bottom (P2) sites are shown in Figs. 4 and 5, respectively (Specimen B5). The results show detectable signals of carbon, nitrogen, oxygen, cobalt and silicon both on the tip and bottom surfaces of the CNCs. It speculates that the signals of oxygen and nitrogen

<table>
<thead>
<tr>
<th>Specimen designation</th>
<th>Applied bias (V)</th>
<th>( E_{\text{turn-on}} ) at 10 ( \mu )A/cm² (V/µm)</th>
<th>( E_{\text{threshold}} ) at 10 mA/cm² (V/µm)</th>
<th>Current density at 10 V/µm (mA/cm²²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>B2</td>
<td>–50</td>
<td>7.6</td>
<td>not reach</td>
<td>0</td>
</tr>
<tr>
<td>B3</td>
<td>–150</td>
<td>8.3</td>
<td>not reach</td>
<td>0</td>
</tr>
<tr>
<td>B4</td>
<td>–200</td>
<td>5.8</td>
<td>8.1</td>
<td>59</td>
</tr>
<tr>
<td>B5</td>
<td>–300</td>
<td>5.0</td>
<td>7.0</td>
<td>173</td>
</tr>
</tbody>
</table>
3.3. Field emission properties

The field emission properties of the as-grown CNCs deposited with different negative substrate biases (Specimens B2, B3, B4 and B5) are shown in Table 2. It shows that the best FE properties belong to the CNCs by the greatest negative bias (−300 V, Specimen B5), which is essentially the CNCs of the greatest aspect ratio; where the turn on electric field ($E_{\text{turn-on}}$), threshold electric field ($E_{\text{th}}$) and the current density (at 10 V/$E_{\text{th}}$) are 5.0 V/$\mu$m, 7.0 V/$\mu$m and 173 mA/cm², respectively. It is interesting to note that the CNCs at a relative lower negative bias than −300 V show the agglomeration phenomenon of CNCs (Specimen B4 at −200 V bias), which may further decline the FE properties due to screening effect among neighbor CNCs. The current versus time ($I$–$t$) curve of the as-grown CNCs at 900 V is used to examine the stability of CNCs during operation, as shown in Fig. 8 (Specimen B5). Except the first 5 min, it indicates that the current remains constant for a testing period of 1 h, though it can be stand longer, implying good adhesion between CNCs with the substrate. The rising of emission current during the first 5 min is due to removal of the loose nanostructures and annealing effect by the intense electric field.

4. Conclusions

By manipulating the H₂/CH₄ ratio and applying sufficient applied negative bias (≥−200 V), the well-aligned CNCs were successfully synthesized by Co-assisted MP-CVD method. The results show that CNCs are basically consisting of mixture of Co particles, graphene planes, disordered graphite, amorphous Si and amorphous carbon. It is found that a greater negative substrate bias can give rise to better-aligned CNCs with sharper apex angle at the tips, better FE properties, greater adhesion with the substrate and better structure stability in electric field. It is also found that the CNCs under −200 V bias has agglomeration phenomenon, which will have a negative effect to FE properties due to screening effect. Under the present conditions, the well-aligned CNCs synthesized under H₂/CH₄ (80/5 sccm/sccm) ratio, −300 V applied bias and 10 min deposition time exhibit good life testing stability and best FE properties: $E_{\text{turn-on}}$ ~ 5.0 V/$\mu$m, $E_{\text{th}}$ ~ 7.0 V/$\mu$m, and $J$ ~ 173 mA/cm² at 10 V/$\mu$m.

Acknowledgements

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References