Thermal-heating CVD synthesis of BN nanotubes from trimethyl borate and nitrogen gas

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Abstract

Boron nitride (BN) is one of III–V compounds widely applied on the electrical industry. It has been fabricated by numerous techniques, but so far there is no reliable method to produce pure and high-yielding BN nanotubes at relatively lower temperature. Therefore, the exploration on its synthesis is still a challenging subject.

In the study, the BN nanotube would be synthesized by thermal-heating chemical vapor deposition (TH-CVD) with trimethyl borate evaporated at 60 ◦C and nitrogen gas flown into reaction chamber as the source of B and N, respectively. 434 stainless steel wires will be coiled as an entangled wire scaffold with pore size of 1 mm and then placed in the middle part of reaction chamber. The metallic ions contained in the stainless steel will serve as the catalysts for of BN nanotube in situ growth.

From the results of SEM, HRTEM, FTIR and XRD analysis, hexagonal-BN (h-BN) and orthorhombic-BN (o-BN) nanotubes were successfully synthesized at relatively low temperature between 1000 and 1200 ◦C. All the nanotubes prepared in the system were identified as h-BN and o-BN. At reaction temperature of 1200 ◦C, several types of BN morphology appeared. The BN nanotubes could be obtained at the temperature between 1000 and 1100 ◦C. However, BN nanotubes for the latter temperature grow into larger size tube. The optimum reaction temperature for BN nanotube synthesis is 1000 ◦C. The reproduction property of synthesized BN nanotube by this method is very promising. The method should have a great potential to prepare BN nanotube in the future.

Keywords: Nitride; Nano-structure; Chemical vapour deposition

1. Introduction

Boron nitride (BN) is one of the most interesting III–V compounds due to its unique properties, such as low density, high thermal conductivity, excellent mechanical strength wear resistance, stability at high temperatures, and possibility of easy doping with silicon (n-type) and beryllium (p-type). Thus, the material appears as a good alternative for carbon-related materials in several applications [1–3]. BN cannot only form hardy compound but also have excellent optics, electric, thermal conduction and mechanical properties because the negative electricity property is similar to carbons. In addition, carbon nanotube demonstrates both conducting and semiconducting property but BN nanotube behaves as a semiconductor [4] which is more convenient to component development and application in the future.

BN nanotubes have been fabricated by different techniques, such as arc-discharge, laser ablation, carbon nanotubes template-confinement and some specifically designed methods [5–9]. Unfortunately, these methods have not been effectively to synthesize a great amount of BN nanotubes. Very recently, a hazardous procedure of producing high-yield BN nanotubes via chemical vapor deposition (CVD) of the precursor borazine on nickel boride catalysts has been reported [10]. However, CVD method produced many types of BN structure and contaminated by the nickel boride or catalysts [11,12]. So far there is no reliable method to produce pure and high-yield BN nanotubes therefore, the exploration on its synthesis is still a challenging...
subject. Furthermore, very high reactant temperature is needed to prepare BN nanotubes from the mentioned methods, such as over 2000 °C for arc-discharged and over 1200 °C for traditional CVD method [13,14].

In this study, thermal-heating chemical vapor deposition (TH-CVD) will be used to synthesize the BN nanotube by vaporizing trimethyl borate at 60 °C and providing nitrogen gas to the reaction chamber as the source of B and N, respectively. 434 stainless steel will be coiled as entangled wire scaffold with pore size of 1 mm. This wire scaffold, thereafter, placed in the middle part of TH-CVD reaction chamber. Cr, Fe, Mo contained in the stainless steel will serve as the catalysts to provide in situ growth of BN nanotube on the coil surface. We hope the method can produce a great amount of pure BN nanotube under relatively low temperature. The produced BN nanotube will be characterized by scanning electron microscope (SEM), high-resolution field-emission transmitted electron microscope (HRTEM), thin-film X-ray diffractometer (TF-XRD) and Fourier transformation infrared (FTIR).

2. Materials and methods

2.1. Materials preparation

BN nanotube is prepared by TH-CVD method with auto-catalyzed process, which is briefly described as follows. 434 stainless steel wire with 0.5 mm in diameter is coiled in a disk-like shape and placed in the center area of TH-CVD chamber. The chamber is a pure Al2O3 tube with 52 mm in diameter and 60 cm in length. The tube is sealed on both sides by three-way connector with thermal-resistant O-ring. Ultra pure nitrogen gas and trimethyl borate (TMB) are introduced as the source of N and B, respectively. TMB was vaporized in a water bath at the temperature of 60 °C and flow to the TH-CVD chamber via a corrosion-resistant tube. The flow rate of nitrogen gas is 20 cm3 min−1. A rotary pump is used for ventilation. The ventilation goes through a water tank to remove water-soluble gas for environmental protection. TH-CVD chamber was heated to a temperature in the range of 800–1300 °C with the heating rate of 3 °C min−1 and the maintained for 2 h. The reaction gases will form a stagnant flow to complete the reaction on the surface of 434 stainless steel coil. BN nanotube can be harvested from the coil surface. The design of the TH-CVD device is schemed as Fig. 1 [14].

2.2. Materials analysis

The morphology of BN growing on the surface of the stainless steel coil is examined by scanning electron microscope (SEM, FE-SEM HITACHI S4100). The specimen is coated with Au-film by sputtering CVD. The film should intensify the back-scattering electron and secondary electron for better resolution. Thin-film X-ray diffraction (TF-XRD) (Rigaku MPA-2000, Japan) is used to analyze the crystal structure of the BN. BN are carefully collected from the coil surface and mounted on the clay cake for scanning. The sample is scanned in a range of 10–90°(2θ) with a scanning rate of 4° min−1. The functional group and vibration model of the BN nanotube are analyzed by FTIR (JAPAN JASCO FT/IR 410 Series spectrometer). The investigated spectral range was between 400 and 4000 cm−1. A high-resolution transmission field emission electron microscopy (HRTEM Hitachi model HF-2000, 200 keV) is utilized to analyze the crystal structure and microstructure of BN nanotubes [15]. The stainless steel coil with BN nanotube deposition is placed in an ultrasonic tank for 15 min in 95% ethanol. Ethanol with BN nanotube is dripped onto a carbon-coated-copper grid and dried in air for TEM examination. Selected area electron diffraction is performed on HRTEM.

3. Results

3.1. Surface morphology of BN formation at different temperatures

The BN crystal observed under SEM shows different images in different reaction temperatures. From the morphological examination, they are not all to demonstrate as a tubular structure. At lower temperature, it forms needle-like structure. At higher temperature, it shows a tube-like structure and/or a granular-like structure.

Fig. 2(a) is the image of BN crystal produced at the temperature of 800 °C. BN crystal sporadically appears on the surface of
the stainless steel coil. The morphology of the BN is not a tube structure but a plate structure with nano-scale in size as shown in Fig. 2(b). The thickness of the plate is less than 10 nm. No nano-tube or tube-like BN can be observed on coil surface. We believe that the reaction temperature is too lower to crystallize BN in a larger scale due to low driving force. Higher reaction temperature may solve matter.

When the reaction temperature up to 900 °C (Fig. 3(a)), the BN crystal cover the entire area of the stainless steel coil surface. SEM images indicate a needle-like structure, which are 1 μm length and 70 nm in diameter in average (Fig. 3(b)). The length of BN crystal at 900 °C is much longer than that of at 800 °C. When the temperature increased, the driving force for crystallization should be increased where the BN needle-like crystals should be a favorable structure.

When the reaction temperature is up to 1000 °C (Fig. 3(a)), the BN crystal cover the entire area of the stainless steel coil surface. SEM images indicate a needle-like structure, which are 1 μm in length and 70 nm in diameter in average (Fig. 3(b)). The length of BN crystal at 900 °C is much longer than that of at 800 °C. When the temperature increased, the driving force for crystallization should be increased where the BN needle-like crystals should be a favorable structure.

When the reaction temperature is up to 1000 °C, BN is crystallized like a tubular structure with 70 nm in diameter and 4 μm in average length (Fig. 4(a)). The microstructure of BN crystal at a reaction temperature of 1100 °C is similar to that of 1000 °C (Fig. 4(b)).

When the reaction temperature is up to 1200 °C, there are many types of BN appeared on the stainless steel coil surface such as wire, bead string structure, and bamboo like tube as shown in Fig. 5.

The XRD patterns of BN deposited at 1200 and 1000 °C are shown in Fig. 7(a) and (b), respectively. Fig. 7(c) and (d) is the patterns for clay and 434 stainless steel, respectively. If the peaks corresponding to clay and 434 stainless steel subtracted, the XRD patterns of BN deposited at 1200 and 1000 °C are identified as orthorhombic boron nitride (o-BN) and hexagonal boron nitride (h-BN), respectively.

3.2. XRD characteristics of BN nanotubes

The XRD patterns of BN deposited at 1200 and 1000 °C are shown in Fig. 7(a) and (b), respectively. Fig. 7(c) and (d) is the patterns for clay and 434 stainless steel, respectively. If the peaks corresponding to clay and 434 stainless steel subtracted, the XRD patterns of BN deposited at 1200 and 1000 °C are identified as orthorhombic boron nitride (o-BN) and hexagonal boron nitride (h-BN), respectively.

3.3. Characteristics of BN crystal

FTIR was used to analyze the functional groups of the BN deposited on stainless steel coil. Fig. 8 shows the FTIR pattern of BN nanotubes produced at the temperature of 1000 °C. The
Fig. 5. The image of BN nanotubes produced at the temperature of 1200 °C (a) many types of BN morphology; (b) bead string structure; (c) bamboo like tube; (d) bead like structure.

Fig. 6. The morphology of BN crystal at the temperature of 1300 °C (a) BN crystal on coil surface; (b) the tube grows out to form a facet crystal.

absorption bands at the wavenumber of 1360 and 785 cm⁻¹ are attributed to the B–N stretching and B–N–B bending of BN, respectively. The FTIR patterns at 1100 and 1200 are similar to that of 1000 °C. This result further proves that the deposited crystal in this study is BN nanotubes.

3.4. Transmission and electric diffraction patterns of BN nanotubes

Fig. 9(a) shows the TEM image of BN nanotube produced at 1000 °C. The diameter of nanotube is about 50 nm.

Fig. 7. The XRD patterns of BN nanotube deposited at the reaction temperature of (a) 1200 °C; (b) 1000 °C; (c) clay; (d) 434 stainless steel.
Fig. 8. The FTIR pattern of BN nanotubes produced at the temperature of 1000 °C.

Fig. 9(b) is the diffraction pattern of the TEM image which shows a ring pattern with four clear rings on the pattern. The four rings from inner part to the outer are identified as (0 0 2), (1 0 1), (0 0 4) and (1 1 2) of h-BN, respectively.

Fig. 10. (a) A multi-walled BN nanotube produced at the temperature of 1000 °C; (b) the diffraction pattern of the image.

Fig. 11(a) is the TEM image of BN nanotube produced at 1200 °C. Fig. 11(b) is the lattice image of Fig. 11(a). The lattice distance of the pattern is about 3.345 Å that is corresponding to (0 0 2) plane of h-BN. As observed in SEM, BN also demonstrate several types of morphology as shown in Fig. 12(a). Most of the tip of the BN nanotubes produced at 1200 °C is closed with the axial sequence of inverted conical void (Fig. 12(b)). Although different types of morphology shown, BN nanotubes produced at 1200 °C are h-BN.
4. Discussions

Crystal growth of one-dimensional structure requires a pathway to continuously providing gas- or liquid-phase reactants to a specified site, in which an anisotropic driving force exists to assure an oriented growth [15,16]. When gas reactants and liquid impurity (catalyst) are used, the vapor–liquid–solid (VLS) growth mechanism is responsible for the synthesis of wire-like structure [17]. Morales and Lieber synthesized Si and Ge nanowires by combining laser ablation and a V–L–S method [18]. This technique can produce nano-scale liquid catalysts, which determine the diameter of nanowires.

In the study, the overall reaction to produce BN crystal is as follows:

\[
\text{C}_3\text{H}_9\text{BO}_3(\text{g}) + \text{N}_2(\text{g}) \rightarrow \text{BN}(\text{s}) + \text{NH}_3(\text{g}) + 3\text{CO}(\text{g})
\]

The residual gases of NH\(_3\)(g) + 3CO(g) are removed from the reaction system through a water tank by a rotary pump. BN crystals formed on the coil surface and catalyzed by the Fe, Cr, Mo elements provided from the surface of the coil scaffold.

At lower temperature, catalysts are difficult to combine with absorbed reactant gases and reach a liquid eutectic composition. In this condition, the BN crystal deposited on the 434 stainless steel coil surface is epitaxial growth. The crystal orientation should be matched to the crystal orientation of coil substrate. A plate-like morphology is the most favorable as shown in Fig. 2. When the temperature increased to the 900 °C, the crystal growth is also based on epitaxial growth but the growth rate is much faster than that of at 800 °C due to higher driving force. This leads to a needle-like structure not a hollow tube, as shown in Fig. 3.

When the temperature is up to 1000 °C, Fe, Cr, or Mo will combine with the reactant gases to reach a eutectic composition to form a partial liquid on the surface. The V–L–S mechanism to form the BN nanotube occurs at 1000 °C as shown in Fig. 4. If the temperature increased to 1100 °C, BN nanotube also appeared on the coil surface as the same mechanism as 1000 °C. Nevertheless, the nanotubes decreased in size are due to coalescence phenomena to eliminate excessive surface energy.

When temperature increases to 1200 °C, Fe Cr, or Mo partial pressure in the reaction chamber will increase. BN crystals growth mechanism at this temperature is not only V–L–S but also V–S or L–S that causes many kinds of BN morphology to present on the coil surface (Fig. 5). In addition, the growth process of the nanotube in the study is the axial sequence of inverted conical void (Fig. 12). The mechanism has been proposed for BN nanotube growth by catalytic capillary [19–22]. When temperature is up to 1300 °C, there are no nanotubes or wires to form...
on the coil surface. It becomes granular structures with hollow inside due to tube fast growth to minimize the surface energy.

From the study of XRD analysis (Fig. 7), the BN deposited on the substrate is identified as two types of BN crystal structure (h-BN and o-BN). However, all the nanotubes are identified as h-BN by FTIR, TEM, and HR-TEM. The h-BN nanotubes can be obtained between the temperatures of 1000 and 1100 °C, which is relatively low for the BN nanotubes preparation.

5. Conclusions

In this study, BN nanotubes were successfully synthesized at relatively low temperature between 1000 and 1100 °C. All the nanotubes prepared in the system were identified as h-BN that was formed on the o-BN template. When the reaction temperature is 1200 °C, there are many types of h-BN morphology appeared. BN nanotubes formed at 1100 °C are larger than the one at 1000 °C. The optimum temperature for BN nanotube synthesis is 1000 °C. The reproduction property of synthesized BN nanotube by this method is very stable comparing with conventional methods. This method should have a great potential to prepare BN nanotube in the future.

References