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Self-organized growth of bamboo-like carbon nanotube arrays for field emission properties

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Abstract Well-aligned nitrogen-doped carbon nanotube (N-CNTs) film was fabricated on silicon substrate by thermal chemical vapor deposition process with varying the growth temperature. The effect of growth temperature on morphology, microstructure and crystallinity for the growth of N-CNTs was studied. At all growth temperatures, the bamboo-like morphology of graphene layers with compartments in CNTs were observed in transmission electron microscope micrographs. The doping level and the type of nitrogen-related moieties were determined by X-ray photoelectron spectroscopy analysis. The compartment distance decreases with increase in nitrogen doping level in hexagonal graphite network. The increase in nitrogen doping level in N-CNTs will lead to decrease in crystallinity and in-plane crystallite size. Field emission study of nitrogen-doped carbon nanotubes grown at optimum parameters showed that they are good emitters with a turn-on and threshold field of 0.3 and 1.6 V/µm, respectively. The maximum current density was observed to be 18.8 mA/cm² at the electric field of 2.1 V/µm. It is considered that the enhanced field emission performance of doped nanotube is due to the presence of lone pairs of electrons on nitrogen atom that supplies more electrons to the conduction band.

Keywords Bamboo-like carbon nanotubes · Field emission · Turn-on field · Threshold field

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Introduction

Miniaturization of nano-scale devices forced researchers to discover advanced materials with unique bulk and surface properties. Carbon nanotube (CNT) is one of the promising materials having exceptional properties for potential use in scanning probe sensors (Hafner et al. 1999), cold cathode flat panel display (Choi et al. 1999), fuel cells (Reddy et al. 2009), supercapacitors (Niu et al. 1997) and vacuum nanoelectronics. Compared to pristine CNTs, the synthesis of nitrogen-doped carbon nanotubes (N-CNTs) have gained growing interest to tailor the electronic (Terrones et al. 2002; Ghosh et al. 2010; Lim et al. 2006), electrical (Ibrahim et al. 2010) and mechanical properties (Ganesan et al. 2010) with the controlled process parameters aid. The electrical and electronic properties of the CNTs can be modified to a large extent by doping the CNTs. It has been predicted that the N-CNTs have n-type conductivity, low resistance and excellent electron field emission property (Czerw et al. 2001). Incorporation of N in carbon network will lead to the shift of the localized state to the Fermi level, and the significant contribution of the coupled state near the Fermi level (Czerw et al. 2001). In wide range of applications of CNTs, field emission is of paramount interest in which electrons are emitted from a solid material by quantum mechanical tunneling when electric field is applied in high vacuum. CNTs are considered to be one of the best candidates for the field emission due to their atomically sharp tips with very high aspect ratio. The field emission property of the material is dependent on electronic properties. The electronic property of the N-CNTs depends on the dopant concentration and it is a relatively facile method to tune this property by changing the dopant concentration into the bulk material.

This paper mainly focused on feasible process to develop the organized growth of highly aligned N-CNT



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arrays with nitrogen content modulated to tailor the field emission property.

Experimental procedure

Nitrogen-doped CNTs were synthesized by injection chemical vapor deposition (CVD) on silicon substrate. The silicon substrate was cleaned with acetone and placed at the center of the furnace. The chamber was evacuated and then filled with argon gas to create the inert atmosphere inside the chamber before injecting the precursors. The precursors used were acetonitrile (Trident Fine Chemicals and laboratories, 99.8 %) and Ferrocene (Merck, 98 %). Acetonitrile was the dual source for carbon and nitrogen while ferrocene, on pyrolysis, provided iron which acts as the catalyst for CNT growth. The catalyst and precursor solution was made with the catalyst concentration of 10 mg/ml and the solution was injected at the rate of 0.5 ml/min. The precursor was vaporized at 170 °C and carried towards the silicon substrate by the carrier gas. The synthesis was carried out at the three different growth temperatures 800, 850 and 900 °C with argon gas flow rate of 180 sccm.

The alignment and microstructure of as-grown CNTs were characterized using scanning electron microscope (SEM, Model S-4300S E/N, Hitachi, Japan) and transmission electron microscope (TEM, Model FEI Technai G² S-Twin). Raman spectroscopy (Lab Ram HR 800 Raman spectrometer) was carried out using 514.5 nm green line of Ar+ ion laser to study the crystallinity and the disorder in CNTs. The measurement of elemental composition was carried out using X-ray photoelectron spectroscopy (XPS, Omicron with Al K_{α} 1,486.8 eV radiation source). N-CNTs arrays thin film deposited on silicon wafer was used as the cold cathode and stainless steel plate as counter electrode was placed at a distance of 1 mm for field emitters. The field emission properties of the deposited N-CNTs were characterized in high vacuum environment with a base pressure of about 6.3×10^{-7} mbar. Programmable Keithley 248 high voltage power supply with low output ripple for precision sourcing and flexible operation was used for sourcing the voltage (up to 3,000 V) and measuring the current (with pA sensitivity), allowing the characterization of the current-voltage (I-V) behavior.

Results and discussion

Microstructure analysis

Figure 1 shows the SEM and TEM images of N-CNT arrays grown at different growth temperatures. It can be observed that the average diameter of the N-CNTs grown



at 800, 850 and 900 °C are 28, 65 and 33 nm, respectively. N-CNT arrays grown at the growth temperature 850 °C are well aligned having the length up to 50 µm. At 800 and 900 °C the CNTs are short and not highly aligned. It indicates that the carbon source that decomposed at 850 °C was able to provide the more amount of precursor for the nucleation of the carbon nanotubes leading to the increase in length as compared to other growth temperatures. Similar phenomenon was observed by other groups (He et al. 2005). The change in their microstructure by nitrogen incorporation was investigated by TEM studies. Nitrogendoped CNTs exhibit very distinct morphologies, significantly different from undoped counterparts. The undoped CNTs are continuously hollow throughout their length. Doped CNTs exhibited bamboo-shaped morphology and each carbon nanotube is separated into a series of compartments with corrugated walls (Jang et al. 2004). This is caused by nitrogen atoms embedded in graphitic lattice, including the curvature of nanotubes due to the different bond length of C-N and C-C. The difference in the bond length will induce the strain in the hexagonal graphite network and the graphene sheets will try to bend due to pulsating effect. The introduction of such defects disrupts the planar hexagonal arrangement of carbon atoms in CNTs and results in the rough surface of doped CNTs. The increased nitrogen concentration induces more defective graphene sheets. As a result the graphene sheets are easily buckled, facilitating cross-linking between the graphene planes through sp³ coordinated carbons. Through the N-sp² C bond, nitrogen is incorporated in the graphene sheets without changing the graphitic structure. TEM micrographs depicted that the length of the compartment in N-CNTs keeps on decreasing as the nitrogen content in hexagonal network increases. As the nitrogen content increases, the buckling of the graphene sheets will take place and try to stabilize themselves by framing the curved layers, i.e., internal capping. The distance between internal capping of CNTs grown at 800, 850 and 900 °C was found to be 43, 60 and 20 nm, respectively.

X-ray photoelectron spectroscopy

The composition and the functional moieties present in N-CNTs were analyzed by using the XPS. It provides detailed information on the occupied electronic levels and chemical binding states of individual atoms. The Gaussian multi-peak reveals the individual nitrogen moieties bonded to the carbon atoms in CNTs. XPS-wide spectrum showed strong peak at 285, 401 and 530 eV which can be assigned to C1s from sp² hybridized carbon in the nanotubes; the carbon atoms on the edge plane can be easily oxidized to oxygen-containing functionalities and nitrogen incorporated in carbon network, respectively. The asymmetric N1s



Fig. 1 Typical SEM and TEM micrographs of NCNT arrays grown at different growth temperatures at 800 °C (\mathbf{a} , \mathbf{b}), at 850 °C (\mathbf{c} , \mathbf{d}) and at 900 °C (\mathbf{e} , \mathbf{f})

peak in narrow spectrum centered at 401 eV can be deconvoluted into four peaks at 398.5, 401, 403 and 405 eV (Liu et al. 2010). They are ascribed as pyridine-like nitrogen (pN), graphitic-like nitrogen (gN), oxidized nitrogen and molecular nitrogen. 398.5 eV peak is assigned to a tetragonal nitrogen phase bonded to a sp³ hybridized carbon, 401 eV assigned to a trigonal nitrogen phase bonded to a sp² coordinated carbon, 403 eV is assigned to oxidized N species or oxygen molecules physisorbed onto the rough surface and 405 eV is assigned to molecular

nitrogen molecules adsorbed or intercalated at the carbon walls or trapped in the compartments.

With decreasing the growth temperature, the XPS peak intensities for the N-sp² C bond decreased, whereas those for the N-sp³ C bond increased, accounting for the deteriorated crystallinity of the graphene sheets. The N-sp³ C bonds were found to be more abundant than the N-sp² C bonds around 4 % nitrogen concentration. The percentage of nitrogen in CNTs was calculated by taking into account the individual areas of C1s and N1s. The formula used is



 $N \% = [N \times 100/(C + N)]$. The nitrogen content was found to decrease from 4.30 to 3.44 % when the growth temperature was increased from 800 to 850 °C, whereas, the N % was increased to 12.34 % at 900 °C growth temperature. The abnormal behavior in increase in nitrogen content was observed as the growth temperature increased and the reason for this is yet to be understood. The graphite type nitrogen (N-sp² C) was found to be more abundant than the pyridine type (N-sp³ C bonds) bonds at higher growth a temperature was showed in Fig. 2b, d, f.

Raman spectroscopy

Raman spectroscopy is a frequently used technique to characterize the carbon nanomaterials. It is a non-destructive tool applied to the identification and characterization of a wide variety of carbon nanomaterials, and the technique has been shown to be an excellent tool to investigate the crystallinity and graphitic nature of CNTs. Raman spectra of CNTs showed a strong band around 1,585 cm^{-1} , originating from the Raman active E2 g mode which is referred to as the G band and a D band at about 1.350 cm^{-1} which is normally explained as a disorder-induced feature. The D band originates from defects in the curved graphene sheets. In N-CNTs, the formation of pentagons and heptagons due to the doping of N atoms leads to a distortion in the graphite sheets. Thus, the intensity ratio of the D to G band in the N-CNTs will be affected by the number of defects originating from nitrogen atom incorporation in hexagonal network. As the concentration of the N atoms increases, the D band becomes stronger and broader.

The intensity ratio I_d/I_g is thus useful in estimating the defect concentration of N in the N-CNTs; as the I_d/I_g ratio decreases the CNTs will have a more ordered structure. The relative intensity ratio of the D band to G band (i.e., I_d / Ig) can be used as a rough measure of N-CNTs quality. Nitrogen content will have the profound effect on the G band (Liu et al. 2010; Yang et al. 2005). Raman analysis of N-CNTs grown at different growth temperature was illustrated in Fig. 3a, b showed the ratio of D and G band and the in-plane crystallite size of the samples. N-CNTs, differences in chemical bond lengths and atomic masses as well as the formation of pentagons due to the doping of N atoms lead to local distortion in the graphite sheets. The I_d/I_g ratio gives the measure of crystallinity of the CNTs. The lower its value, higher is the crystallinity. The in-plane crystallite size strongly depends on I_d/I_g ratio (Cancardo et al. 2006) and it will decrease with the increase in I_d/I_g ratio. In-plane crystallite size can be calculated as

$$L_{\rm a} = (2.4 \times 10^{-10}) \lambda^4 / (I_{\rm d}/I_{\rm g})$$

where $L_a =$ in-plane crystallite size (nm), $\lambda =$ wavelength of the laser source (nm).



Field emission study

Field emission consists in the extraction of electrons from a conducting solid under the application of an electric field and the mechanism for the emission of electron was illustrated in Fig. 4a. Unlike thermion emission, no heat is required for obtaining field emission. Very high electric fields are, however, required for the electrons to tunnel through the surface potential barrier. When the solid is shaped as the tip, the electric fields lines are concentrated around the tip and the local electric field is enhanced. This geometrical enhancement of the electric field is used in field emitters to allow extraction of electrons from sharp tips at relatively low macroscopic electric fields. Maximizing the current output for a given applied voltage requires a material with a low work function that is shaped as sharp as possible to offer the highest field enhancement factor. The emission characteristics of emitters are strongly dependent on the tip radii, and preservation of the emission characteristics requires the tip sharpness remain unaltered. Figure 4b showed the measured I-V characteristics of the carbon nanotube pillar arrays. It indicated that the emission current increases monotonically with the applied field and the turn-on field value required to obtain 2 µA current for cold cathode was observed at ~ 500 V. Good field emission from the doped CNTs is attributed to the presence of nitrogen atoms in the hexagonal graphite network. In doping mechanism, the nitrogen substitutes the carbon atom on the graphite layer and modifies the conductance property of N-CNTs by donating electrons. The iron particles were used as the catalyst for the growth of the CNTs and were encapsulated. These catalyst particles may also contribute additional electron density that leads to improved field emission current. Field emission study of as-grown nitrogen-doped carbon nanotubes suggests that they are good emitters with a turn-on and threshold field of 0.9 and 1.6 V/µm, respectively. The maximum current density was observed to be 18.8 mA/cm² at electric field of 2.1 V/µm. The Fowler-Nordheim model was used to describe the field emission effectiveness. Figure 4b shows the corresponding Fowler-Nordheim (F-N) plots. The F-N plots of the N-CNTs arrays almost straight line, showing that the electron emission is due to the tunneling effect. According to the F-N theory, the slope of the F-N plot is $-6.83 \times 10^9 \beta^{3/2} d/\beta$, where β is the work function of the emitter and d is the electrode distance and β is the geometrical enhancement factor. The field amplification factor is defined as $F = \beta V/d$, where F is the field just above the surface of the tip, and V the applied voltage. It is considered that the good field emission performance of doped nanotube is due to the presence of lone pairs of electrons on nitrogen atom that supplies more electrons to the conduction band and also the increased local density of states in



Fig. 2 XPS wide spectrum and N1s narrow spectrum for NCNT arrays grown at different growth temperature at 800 °C (\mathbf{a} , \mathbf{b}), at 850 °C (\mathbf{c} , \mathbf{d}) and at 900 °C (\mathbf{e} , \mathbf{f})





Fig. 3 Typical Raman spectrum (a) and the in-plane crystallite size (b) of N-CNT arrays grown at different growth temperatures



the conduction band close to the Fermi level through nitrogen incorporation.

The nitrogen incorporation into the nanotubes hexagonal network enhances the electrical conductance of nanotube films which in turn lowers the turn-on field and threshold fields in the field emission study. When the graphite sheet of CNTs bends (internal capping), which resulted in the deterioration in crystallinity of CNTs, the atomic bonding must lose some of its sp² character and gain some sp³ character. The transformation of sp² into sp³ characteristics will enhance the emission properties by lowering of the effective work function, increase of the field enhancement factor and the number of the emission sites. Along with the lone pairs of electrons supplied by the nitrogen atom, the curve at the tip of the N-CNTs consist of defects responsible for the concentration of degree of states (DOS) at the tip and are the prime source to improve the field emission. The disordered structure of the tip and the high defect



density may also have an influence on the field emission properties.

Conclusions

Nitrogen-content-modulated growth of CNTs was developed by controlling the growth temperature to tailor the field emission properties. The additional lone pair of electrons on nitrogen atom that acts as a donor with respect to the delocalized π system of the hexagonal framework which can enhance the conducting property of doped CNTs in turn will enhance the emission properties by lowering the effective work function. The maximum current density was observed to be 18.8 mA/cm² at 2.1 V/µm electric field and linear Fowler–Nordheim plot verifies that such field electron emission characteristics can be mainly caused by the quantum tunneling effect. Acknowledgments We are thankful to director Dr. G. Sundararajan for permitting to publish these results. We are also thankful to G.V. Reddy and K. Subbarao for their technical support in present work. We are thankful to Dr. B.V. Sharada for Raman spectroscopy, Dr. Neha Hebalkar for XPS analysis, Mr. L. Venkatesh for SEM micrographs.

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