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Fe-Ti-O Based Catalyst for Large-Chiral-Angle Single-Walled Carbon Nanotube Growth

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Abstract:

Catalyst selection is very crucial for controlled growth of single-walled carbon nanotubes (SWNTs). Here we introduce a well-designed Fe-Ti-O solid solution for SWNT growth with a high preference to large chiral angles. The Fe-Ti-O catalyst was prepared by combining Ti layer deposition onto premade Fe nanoparticles with subsequent high-temperature air calcination, which favours the formation of a homogeneous Fe-Ti-O solid solution. Using CO as the carbon feedstock, chemical vapour deposition growth of SWNTs at 800 °C was demonstrated on the Fe-Ti-O catalyst. Nanobeam electron diffraction characterization on a number of individual SWNTs revealed that more than 94% of SWNTs have chiral angles larger than 15°. *In situ* environmental transmission electron microscopy study was carried out to reveal the catalyst dynamics upon reduction. Our results identify that the phase segregation through reducing Fe-Ti-O catalyst leads to the formation of TiO_x-supported small Fe nanoparticles for SWNT growth. The strong metal-support interactions induced by partial reduction of TiO_x support promote the wettability of Fe nanoparticle, accounting for the preferential growth of large-chiral-angle SWNTs. This work opens a new avenue for chiral angle selective growth of SWNTs.

Keywords: single-walled carbon nanotube, Fe-Ti-O catalyst, environmental transmission electron microscope, chemical vapor deposition, large chiral angle

1. Introduction:

With the aim to exploit potential technological applications of single-walled carbon nanotubes (SWNTs), it is necessary to synthesize SWNTs with well-defined structures [1, 2]. This is due to the fact that the unique fundamental properties of an SWNT, like electrical and optical properties, derive from the fashion of folding carbon sheet described by the tube chirality. A pair of chiral indices (n, m) is commonly used to define the tube chirality. Highly crystalline SWNTs can be produced by high temperature techniques, such as arc discharge [3, 4] and laser ablation [5] methods. However, these

methods usually yield SWNTs with mixed properties. Moreover, SWNTs produced by these high temperature techniques typically need to be dispersed onto desirable substrates for further device fabrication. Nowadays, chemical vapour deposition (CVD) technique has become the most widespread method for directly synthesizing SWNTs on substrate/support, since the orientation [6-8], density [9, 10], diameter [11] and even chirality [12, 13] can all be controlled to a certain extent.

As a subgoal of fully control the final structure of SWNTs, controlling the chiral angles of SWNTs has been challenging [14-17]. Indeed, many growth parameters involved in CVD, including catalyst composition [12, 13, 18-20], catalyst state [17], carbon source [21, 22], reaction temperature [12, 13, 23, 24] and metal-support interactions [25, 26], all affect the SWNT growth results, making chiral-selective SWNT growth extremely difficult. Among all the above parameters, catalyst is known as the most important parameter in defining SWNT chirality distribution. Since the pioneer work of CoMoCAT catalyst [18], a number of catalyst systems have been proposed for chiral-selective SWNT growth in the past decade. However, in most instances, the chirality selectivity has been limited to low temperature growth of small diameter SWNTs [12, 18, 24]. Therefore, it is still highly desired to develop catalyst systems suitable for growing SWNTs with relatively large diameters and certain chiral-angle bias.

Solid solutions arise recently as promising catalysts for achieving SWNTs with specific chiralities [13, 19]. This is because the solid solution offers additional degree of freedom for regulating the morphology and structure of the active catalyst component. For example, by tuning the synthesis temperatures, SWNTs enriched with different tube types were obtained on a $\text{Co}_x\text{Mg}_{1-x}\text{O}$ solid solution [19]. Reduction of such a $\text{Co}_x\text{Mg}_{1-x}\text{O}$ catalyst leads to the epitaxial formation of uniform Co nanoparticles and the concomitant strong metal-support interactions (SMSI), accounting for the high SWNT selectivity to near-armchair species. SMSI is thus anticipated to favor the synthesis of SWNTs with bias chirality. It has been long recognized that TiO_2 supported metals (like Fe) possess SMSI [27]

and show impressive properties in Fischer–Tropsch [28, 29] and carbon nanotube growth [30, 31]. However, on the one hand, the structures and dynamics of Fe-Ti-O catalyst upon reduction are still unveiled; on the other hand, the development of TiO₂-based solid solution for the synthesis of SWNTs, especially SWNTs with certain chiral angle selectivity, has been scarce.

In this work, we present a Fe-Ti-O solid solution catalyst for synthesizing SWNTs with a high preference to large chiral angles. The Fe-Ti-O solid solution is obtained by high-temperature calcination of Ti-covered Fe nanoparticles, which are supported by porous Si₃N₄ grid. The chiralities of SWNTs crossing the grid holes will be identified by nanobeam electron diffraction (ED) and compared with those grown on monometallic Fe catalyst. In order to understand the SWNT growth mechanism, the reduction behavior of the Fe-Ti-O catalyst will be elucidated with the assistance of *in situ* environmental transmission electron microscopy (E-TEM). The SMSI is postulated to play an important role in the chiral-angle selective growth of SWNTs.

2. Materials and methods

2.1 Preparation and characterization of Fe-Ti-O catalyst

Fig. 1 depicts the preparation of the Fe-Ti-O solid solution. The FeO_x nanoparticles were prepared by hydrolysis of anhydrous ferric chloride (FeCl₃) [32]. Briefly, FeCl₃ with a mass of 0.807g was dissolved in 5 mL H₂O to form a clear solution. The aqueous solution of FeCl₃ was added dropwise into 180 mL boiling H₂O under vigorous stirring and refluxed for 2 h to form FeO_x nanoparticles. As-prepared FeO_x nanoparticles were casted onto a Si₃N₄ TEM grid (DuraSiN mesh). After that, the grid was deposited with 4 nm Ti (Edwards E306A Evaporator e-gun system). The system pressure was kept at 10⁻⁶ Pa during Ti deposition. The catalyst was finally annealed at 900 °C for 20 h in air to facilitate the formation of Fe-Ti-O solid solution.

The morphology and structure of the Fe-Ti-O catalyst were characterized by a Zeiss LIBRA 200FMC TEM operated with an accelerating voltage of 200 kV. Electron energy-loss spectroscopy (EELS) was applied to analyse the elemental components of the Fe-Ti-O particles.

Fig. 1 Schematic illustrations of preparing Fe-Ti-O solid solution catalyst for CVD growth of SWNTs.

2.2 Growth and characterization of carbon nanotubes

The Si₃N₄ supported Fe-Ti-O catalyst was loaded into a horizontal CVD reactor with a quartz tube (inner diameter 40 mm). The system was heated to 800 °C in the protection of helium with a flow rate of 200 cm³/min. After being stabilized at 800 °C, helium was replaced by CO (200 cm³/min) for SWNT growth. The feeding of CO at 800 °C lasted for 1 h. The reactor was cooled to room temperature in flowing helium.

As-prepared carbon nanotubes were characterized with Raman spectrometer with an excitation laser wavelength of 632.8 nm (Horiba Jobin–Yvon LabRam HR800) and scanning electron microscopy (SEM, JEOL JSM-7500F). A JEOL-2200FS double aberration-corrected TEM operated at 80 kV was applied to investigate the carbon nanotubes. The chiralities of SWNTs were identified from their ED patterns on the basis of a calibration-free intrinsic layer line-spacing method [33].

2.3 In situ E-TEM investigations on the catalyst dynamics upon reduction

In situ experiment was performed on an FEI Titan 80-300ST E-TEM operated at an accelerating voltage of 300 kV. The Fe-Ti-O catalyst supported by Si₃N₄ grid was loaded onto an Inconel heating

holder and inserted into the TEM chamber. Once heated up to 700 °C, CO with a flow rate of 6 cm³/min was introduced to the chamber and the pressure was stabilized at 7.0 mbar. The catalyst dynamics upon CO reduction was monitored by TEM imaging.

3. Results and Discussion

3.1. Preparation and characterization of Fe-Ti-O solid solution

Supporting Information Fig. S1a presents a TEM overview of FeO_x nanoparticles prepared by hydrolysis of FeCl₃. After their dispersion onto the surface of Si₃N₄ TEM grid, a layer of 4 nm Ti was deposited onto the entire surface. When the Ti coated FeO_x nanoparticles were heated in air at 900 °C, the mobility of Ti atoms increases, leading to the formation of Fe-Ti-O particles due to surface immigration [34]. A TEM image of the catalyst after air calcination is presented in **Supporting Information Fig. S1b**. Clearly, most of the particles after air calcination have sizes of several tens of nm. **Fig. 2a** presents a TEM image of an ellipse-shaped catalyst particle, showing a perfect single crystalline structure. The major and minor axes of the particle are 28 nm and 19 nm, respectively. EELS (**Fig. 2b**) acquired on as-prepared catalyst particle distinctively show the Ti L₂ and L₃ major edges at 455 eV and 461 eV, respectively. Besides, the O_K edge at 532 eV, the L₂ (702 eV) and L₃ (715 eV) edges of Fe are also present.

Fig. 2 (a) TEM image of a typical Fe-Ti-O catalyst particle. (b) Electron energy loss spectrum taken from a Fe-Ti-O particle. (c) TEM image of another Fe-Ti-O particle and (d) its corresponding fast Fourier transformation digital diffractogram.

Fig. 2c and **Fig. 2d** present high resolution TEM image of a large Fe-Ti-O particle and its corresponding fast Fourier transformation (FFT) pattern, respectively. Our high resolution TEM image fully confirms a single crystalline structure of the Fe-Ti-O catalyst. Combined with its FFT pattern, the crystal structure was assigned as FeTiO_3 with a trigonal rhombohedral structure ($a = b = 0.508 \text{ nm}$, $c = 1.42 \text{ nm}$). The respective interplanar spacing of (102) and (110) are 0.37 nm and 0.25 nm (**Fig. 2d**). All the above results suggest that particles of Fe-Ti-O solid solution are formed on the surface of Si_3N_4 grid.

3.2. Synthesis and characterizations of SWNTs grown on Fe-Ti-O catalyst

Ambient pressure CVD growth of carbon nanotube was performed at 800 °C using CO as the carbon source. **Supporting Information Fig. S2a** presents radial breath modes of Raman spectra acquired at 632.8 nm on the carbon nanotube products, indicating the synthesis of SWNTs. The relatively low density of D mode (**Supporting Information Fig. S2b**) suggests that the quality of produced SWNTs is high. It is noted that Raman scattering is a resonant process. That is, when using a fixed laser wavelength, only a small portion of SWNTs with certain bandgaps can be detected by Raman spectroscopy [35]. In addition, the Raman frequencies of SWNTs are sensitive to environments, challenging the precise assignment of the chiral indices of SWNTs.

Fig. 3a and **Supporting Information Fig. S3a** presents a TEM overview and a SEM overview of as-prepared carbon nanotubes grown on Fe-Ti-O catalyst, respectively. High-resolution TEM (**Supporting Information Fig. S3b**) and ED characterization results confirm that the synthesized nanotubes are single-walled. **Fig. 3b** shows the diffraction pattern of an individual SWNT (inset of **Fig. 3a**). The diffraction layer lines are separated by certain distances from the equatorial layer line at the centre. The period of the equatorial line is related to the diameter of the tube [33, 36]. Coupled with

the spacing of non-equatorial lines, the chirality of an SWNT can be well assigned. For example, **Fig. 3b** gives the SWNT chirality of (10, 8), indicating that the SWNT is a semiconducting tube with a chiral angle of 26.3°. Additional confirmation of the assignment is provided by the simulated ED pattern of a (10, 8) tube (right of **Fig. 3b**).

Fig. 3 (a) A TEM overview of as synthesized carbon nanotubes grown on Fe-Ti-O catalyst at 800 °C using CO as the carbon source. (b) The ED pattern of an individual SWNT (inset of Figure (a)). The right is the simulated diffraction pattern of a (10, 8) tube. (c) Diameter distribution of metallic and semiconducting SWNTs grown on Fe-Ti-O. (d) The distributions of SWNTs grown from Fe-Ti-O and Fe as a function of chiral angles.

Nanobeam ED characterization was performed on a total of 52 randomly isolated SWNTs. The deduced (n , m) indices, chiral angles and diameters of SWNTs are all listed in **Supporting Information Table S1**. There is no preference to tubes with metallic or semiconducting indices. 19 out of 52 SWNTs were identified to be metallic species, comprising 36% of the total, which is close to the theoretically predicted amount (33%). **Supporting Information Fig. S4a** presents the diameter

distribution of SWNTs. Most of SWNTs (96%) have diameters in the range of 1.0 – 2.4 nm, as also demonstrated in **Fig. 3c**. Although no high chiral selectivity is observed, (14, 8) tube with a chiral angle of 21.1° , is the most abundant species detected in the product. Besides, SWNTs with diameters near 1.2 and 1.3 nm, including (10, 8), (10, 9), (11, 7) and (11, 8) were also observed more frequently than other tube species, occupying 23% of all the SWNTs.

Furthermore, **Fig. 3c** depicts that there is no simple correlation between the tube diameter and tube conductivity. Interestingly, most of the tubes (up to 94%) have chiral angle larger than 15° and the mean angle is 24° with a standard deviation of 5° . As suggested by **Supporting Information Fig. S4b** and **Table S1**, only 3 SWNTs, namely one (15, 2), one (23, 7) and one (33, 7) tube, have chiral angles smaller than 15° . **Fig. 3d** compares the percentages of Fe-Ti-O grown SWNTs in different chiral angle ranges with those of SWNTs grown on monometallic Fe catalyst [22]. Clearly, SWNTs with large chiral angles (20° - 30°) were more frequently detected in the Fe-Ti-O grown products (82%).

To help understanding of the above SWNT growth results, nucleation and growth mechanisms of SWNTs should be considered. The growth of an SWNT starts with the nucleation of an SWNT cap, while the position of the sixth pentagon in the cap determines the chiral angle of an SWNT. A few parameters, including gas environments, catalyst composition and metal-support interaction, all could influence the formation of tube cap. In our CVD experiment, pure CO, which has been demonstrated to favour the selective nucleation of certain SWNT caps [22, 37], was applied as the carbon precursor. However, as shown in **Fig. 3d**, the portion of large-chiral-angle SWNTs grown on Fe-Ti-O is higher than that of monometallic Fe-grown SWNTs [22]. Therefore, the use of CO is not fully responsible for the preferential growth of large-chiral-angle SWNTs on Fe-Ti-O catalyst.

3.3. In situ E-TEM studies on the behaviours of Fe-Ti-O catalyst upon reduction

Alloying with high melting temperature metal/metal oxide could affect the catalytic performances of Fe-group transition metal (Fe, Co and Ni). For example, CVD growth of SWNTs with high chirality

selectivity was reported on solid solutions like $\text{Co}_x\text{Mg}_{1-x}\text{O}$ [19, 26] and Co-W [13]. However, the catalytic performances of the above two catalysts are quite different. When reducing the Co-MgO catalyst, Co cations in the solid phase are reduced and migrated into the surface to form metallic Co nanoparticles [19] which catalyse the subsequent growth of SWNTs. While Co-W catalyst nanoparticles appeared to be very stable and no phase separation occurs when heated up to 1100 °C[13].

Fig. 4 (a) TEM image of a Fe-Ti-O particle. (c) TEM images of the same particle shown in (a) after *in situ* reduction in CO at 700 °C. Inset is the overview of the particle. (c) Schematic illustration of the enhanced wettability of Fe nanoparticles on TiO_x support, resulting in less Laplace pressure of Fe nanoparticles.

In order to figure out what is the active composition for growing SWNTs and address why Fe-Ti-O particles can lead to the growth of SWNTs with large chiral angles, *in situ* E-TEM experiments were performed to investigate the reductive behaviours of the Fe-Ti-O catalyst. **Fig. 4a** presents a TEM image of a representative Fe-Ti-O particles before reduction. No small particles anchor on the surfaces of Fe-Ti-O solid solution. After *in situ* reduction by CO at 700 °C, Fe particles with small diameters were observed to form on the surfaces of Fe-Ti-O particles (**Fig. 4b**). Such observations are in

agreement with the work reported by Campos *et al.* [38], who detected the formation of metallic Fe on reduced Fe-Ti-O solid solution using XRD. All the above results suggest that the catalyst behaviours of Fe-Ti-O are similar to those of $\text{Co}_x\text{Mg}_{1-x}\text{O}$ type solid solution, where phase separation occurs upon reduction at high temperature.

As the active component of the Fe-Ti-O solid solution for nucleating SWNTs is Fe nanoparticles, it seems reasonable to understand that SWNTs grown on Fe-Ti-O display no metallic property tuning compared with Fe-grown SWNTs [22]. Further considering both the carbon precursor and the reaction temperature are same, we tentatively attribute the pronounced growth of large-chiral-angle SWNTs on Fe-Ti-O to the effects of the support materials. On the one hand, the acid-base character of the support surface could influence the carbon source chemistry. TiO_2 supported Fe catalyst has been explored in Fischer-Tropsch synthesis where CO is used as one of the starting materials. Analyzed by a gas chromatograph, Arai *et al.*[28] discovered that Fe/TiO_2 catalyst showed a high selectivity to $\text{C}_1\text{-C}_5$ hydrocarbons and no C_{6+} hydrocarbons were produced. Compared with Fischer-Tropsch synthesis [28], the temperature (800 °C) used in our CVD process is higher. Such a high temperature not only facilitates faster reaction and higher conversion rates but favours the production of C_1 hydrocarbon. Such short carbon chains could act as intermediates and be directly attached to the open ends of SWNTs. More importantly, the carbon fragments generated when using TiO_2 as the support could affect the carbon dissolution in the Fe nanoparticles and therefore, the subsequent nucleation of a carbon nanotube.

On the other hand, metal-support interaction could regulate the morphology of supported particles [39], which influences the formation of a SWNT cap. By controlling the rearrangement of catalyst particles, growth of SWNTs with different structures was reported by Harutyunyan *et al.*[40] They demonstrated that there exist correlations between catalyst morphology and the resulted nanotube electronic structure. Our recent work also shows that the carbon incorporation depends on the

nanoparticle structures [41], thus affecting the nucleation of SWNTs. In the case of Fe-Ti-O catalyst, as evidenced by E-TEM experiments, reduced Fe anions migrate onto the surface to form metallic Fe nanoparticles. Different from oxides such as SiO₂, Al₂O₃ and MgO, which usually show weak metal-support interactions, TiO₂ support could exhibit SMSI with reduced Fe nanoparticles [27]. The occurrence of SMSI correlates with the partial reduction of surface TiO₂ to “TiO_x” which converts Ti⁴⁺ with a d⁰ configuration to Ti³⁺ with a d¹ configuration [42]. In the support surface, a local anion deficiency allows the reduced Fe atoms migrate to freshly reduced region of the adjacent TiO_x. The strong strength of the interaction between the metal and the support not only inhibits agglomeration of the metal particles under reaction condition, but induces the metallic particle to adopt a pillbox-like morphology (**Fig. 4b**) that maximizes the number of metal atoms in contact with the support [43]. The SMSI-induced pillbox-like morphology should increase the proportion of low-index planes [42] and eventually result in the growth of SWNTs different from those grown on Si₃N₄ or SiO₂ supported Fe nanoparticles. In addition, the SMSI promotes the wettability of Fe nanoparticles on the substrate and leads to the reduced contacted angle of the nanoparticles. The curvature of Fe nanoparticles decreases accordingly, which results in less Laplace pressure of liquid Fe nanoparticles (**Fig. 4c**) and thus limits their entrance into the carbon nanotubes. As a result, the tube cap structure formed on the Fe nanoparticle differs from that produced on SiO₂ supported Fe, where exists a large contact angle between Fe and substrate [44]. Such an effect could also be induced by modulating carbon content inside metal particles, which has been demonstrated to be of great importance for carbon nanotube nucleation and growth [41, 45]. Indeed, previous work also revealed that metallic Co nanoparticles, which have strong interactions with the underlying MgO support and favor the growth of near-armchair SWNTs [19, 26].

4. Conclusions

In conclusion, a facile deposition and calcination technique was developed to prepare a Fe-Ti-O solid solution for catalysing the growth of SWNTs. ED characterizations on the SWNTs revealed a preferential growth of SWNTs with large chiral angles. *In situ* E-TEM investigations revealed that the active catalyst for growing SWNTs is Fe nanoparticles, generated from migration of Fe ions in the solid solution upon high temperature reduction. The presence of SMSI induced by TiO_x support lead to the formation of metallic Fe nanoparticles with unique morphologies and enhanced wettability of Fe nanoparticle on support, accounting for such SWNT growth results. This work extends the development of solid solution catalysts for CVD synthesis of SWNTs and sheds more light on chiral-selective growth of SWNTs.

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Supplementary data: Extensive figures and results related to catalyst and carbon nanotube characterizations.

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