

Sulfur doped Co/SiO₂ catalysts for chirally selective synthesis of single walled carbon nanotubes†

Hong Wang, Kunli Goh, Rui Xue, Dingshan Yu, Wenchao Jiang, Raymond Lau and Yuan Chen*

Cite this: *Chem. Commun.*, 2013, **49**, 2031

Received 15th December 2012,
Accepted 19th January 2013

DOI: 10.1039/c3cc38973a

www.rsc.org/chemcomm

Monochiral single-walled carbon nanotubes have many potential uses, especially in the field of electronics and medical applications. Here we demonstrate a method to activate three types of Co/SiO₂ catalysts via sulfur-doping for the synthesis of tubes with a narrow chiral distribution around (9,8) chirality.

Electronic and optical properties of single-walled carbon nanotubes (SWCNTs) are correlated with their geometric structures, *i.e.* chirality index (n,m).¹ Thus, many potential applications require the use of monochiral SWCNTs. Chirally selective synthesis of nanotubes is the ultimate goal in SWCNT production, which remains a great challenge. In common growth methods, *i.e.* chemical vapor deposition (CVD), carbon precursors decompose at high temperature to form atomic carbon species, which then nucleate into one-dimensional tubes on catalytic particles. Theoretical studies suggest that catalysts play a critical role in the (n,m) selectivity.^{2–7} To control SWCNT (n,m) structures, the size of catalytic particles need to be first confined in a suitable range.^{7–9} However, metal species are mobile and form particles of various sizes at high growth temperatures, which eventually results in the growth of multiwalled carbon nanotubes (MWCNTs), graphite and amorphous carbon.

Co catalysts supported on silica substrates (Co/SiO₂) are the most often used catalysts for (n,m) selective growth of SWCNTs.^{10–20} Several approaches have been used to control the nucleation of Co, *e.g.* stabilizing Co clusters in bimetallic alloys,^{10,12,20} incorporating Co ions into mesoporous substrates,^{11,14,16} forming dispersed surface Co silicates,¹⁷ anchoring reduced Co atoms by unreduced Co ions,¹⁸ and creating strong interfacial Co–Si interactions.¹⁹ Most of these approaches only result in the selective growth of small-diameter (6,5) tubes. Some previous studies had shown that when sulfur (S) species are added to carbon precursors, it can promote the growth rate and yield of the nanotubes, and change MWCNT diameters.^{21–24} However, to the best of our knowledge, no work has

been reported on the direct doping of S in solid catalysts to control metal particles for SWCNT growth.

In this work, we have demonstrated that non-selective Co/SiO₂ catalysts can be converted into efficient chiral selective catalysts by S doping. SWCNTs were characterized by photoluminescence (PL), UV-vis-near-infrared (UV-vis-NIR) absorption and Raman spectroscopies. Catalysts were characterized by elemental analysis, H₂ temperature programmed reduction (H₂-TPR), and UV-vis diffuse reflectance spectroscopy. The molecular structural changes of Co species on SiO₂ caused by S doping are believed to be responsible for the chiral selectivity.

Three Co/SiO₂ catalysts with 1 wt% Co were prepared by the impregnation method using Co precursors, including Co(II) acetylacetonate (Co(acac)₂, Sigma-Aldrich, 97%), Co(II) chloride (CoCl₂, Alfa Aesar, 97%), and Co(II) nitrate hexahydrate (Co(NO₃)₂·6H₂O, Sigma-Aldrich, 99.999%). They are denoted as CoACAC/SiO₂, CoCl/SiO₂, and CoN/SiO₂, respectively. To add S, the calcined Co/SiO₂ catalysts were impregnated by dilute sulfuric acid (H₂SO₄, 0.04 mol L⁻¹) at 8 mL solution per g catalyst ratio for 1 h. The resulting S doped catalysts were marked as CoACAC/SiO₂/S, CoCl/SiO₂/S, and CoN/SiO₂/S, respectively (see ESI† for details). Catalysts were introduced into a CVD reactor, and first reduced under pure H₂ (1 bar, 50 sccm) from room temperature to 540 °C at 20 °C min⁻¹, and then further heated to 780 °C under an Ar flow (1 bar, 50 sccm). At 780 °C, pressured CO (6 bar, 200 sccm) replaced Ar, and the SWCNT growth lasted for 1 h. As-synthesized SWCNTs with catalysts were first characterized by Raman spectroscopy, and then dissolved in NaOH aqueous solution (1.5 mol L⁻¹) to remove SiO₂. The carbon deposits were then dispersed in 2 wt% sodium dodecyl benzene sulfate (SDBS, Aldrich) D₂O solution. The SWCNT suspension obtained after centrifugation at 50 000 g was characterized by PL and UV-vis-NIR absorption spectroscopies (see ESI† for details).

PL maps in Fig. 1a–f show that two undoped Co/SiO₂ catalysts (CoACAC/SiO₂ and CoCl/SiO₂) resulted in small-diameter tubes (<0.9 nm), such as (6,5), (7,5), (7,6) and (8,4). CoN/SiO₂ is not active for SWCNT growth. This is in agreement with previous studies using various SiO₂ supported Co catalysts.^{15,17,25} In contrast, after doping with S, the major (n,m) products are large-diameter tubes (>1.1 nm), such as (9,8), (9,7), (10,6), and (10,9). The abundance of these

School of Chemical and Biomedical Engineering, Nanyang Technological University, Singapore 637459, Singapore. E-mail: chenyuan@ntu.edu.sg; Tel: +65-63168939

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c3cc38973a



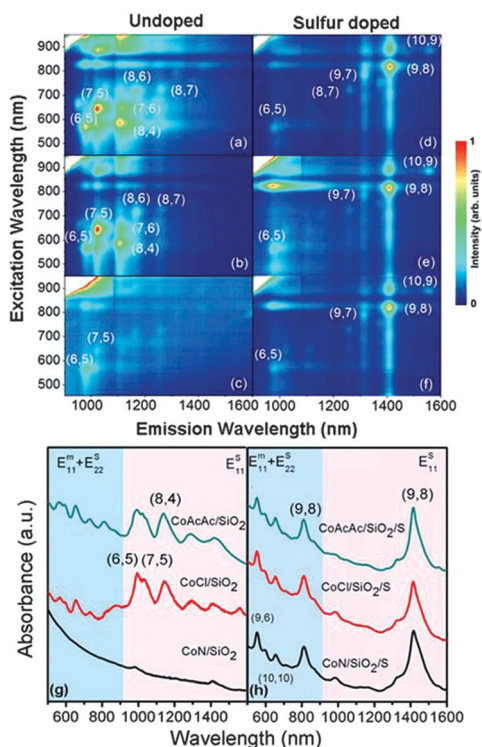


Fig. 1 PL maps and UV-vis-NIR absorption spectra of SDBS-dispersed SWCNTs grown on undoped and S doped Co/SiO₂ catalysts. PL maps: (a) CoACAC/SiO₂, (b) CoCl/SiO₂, (c) CoN/SiO₂, (d) CoACAC/SiO₂/S, (e) CoCl/SiO₂/S, and (f) CoN/SiO₂/S. Some major (*n,m*) species identified on PL maps are marked. Absorption spectra: undoped (g) and S doped (h). The shaded pink indicates the E_{11}^S absorption band and the shaded blue shows the overlapping E_{22}^S and E_{11}^M bands.

four species calculated using their PL intensity is 52.4–69.1% of all semiconducting species identified, out of which, 32.7–40.5% is (9,8) (see Tables S1–S3 in the ESI†). PL results were corroborated by UV-vis-NIR absorption spectra. Fig. 1g shows that SWCNTs from CoACAC/SiO₂ and CoCl/SiO₂ have intense absorption peaks at 992, 1025 and 1137 nm from (6,5), (7,5), (7,6), and (8,4). SWCNTs grown on CoN/SiO₂ have weak absorption peaks. In contrast, Fig. 1h shows that SWCNTs grown on S doped catalysts all have strong absorption peaks at 1420 and 818 nm, corresponding to the E_{11}^S and E_{22}^S transitions of (9,8). A few other absorption peaks below 700 nm can be assigned to either the E_{11}^M transition of metallic (9,6) (1.02 nm) and (10,10) (1.36 nm) or the E_{22}^S transition of semi-conducting (6,5). Since the intensity of (6,5) PL peaks in Fig. 1d–f is low, the absorption peaks below 700 nm in Fig. 1h are likely to be from those metallic tubes. SWCNTs were further characterized by Raman spectroscopy under two excitation lasers (785 and 514 nm). Raman results (see Fig. S1 in the ESI†) agree with the findings from PL and UV-vis-NIR. Overall, the three spectroscopic techniques have demonstrated that S doping can shift the (*n,m*) selectivity of Co/SiO₂ catalysts from small-diameter tubes near (6,5) to large-diameter tubes with a narrow distribution around (9,8).

Co species deposited on SiO₂ would first be partially reduced in H₂, and then nucleated into Co nanoparticles to initiate SWCNT growth. The change in (*n,m*) selectivity shown in Fig. 1 may be attributed to the changes in Co species caused by S doping. Firstly, we conducted an elemental analysis of S doped catalysts.

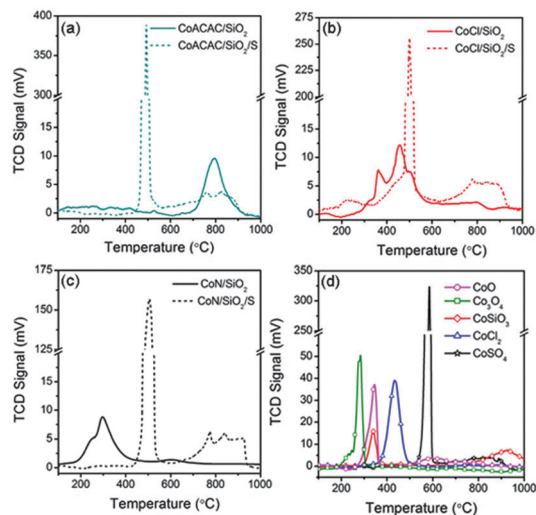


Fig. 2 H₂-TPR profiles of undoped and S doped Co/SiO₂ catalysts and several Co references (CoO, Co₃O₄, CoSiO₃, CoCl₂ and CoSO₄·7H₂O).

The S content in CoACAC/SiO₂/S, CoCl/SiO₂/S, and CoN/SiO₂/S was found to be 0.91, 1.17 and 0.83 wt%, respectively. This confirms the existence of S. Next, H₂-TPR was employed to study the reducibility of Co species. Fig. 2 shows that CoACAC/SiO₂ displays a peak around 797 °C, which is due to the surface Co silicate.¹⁷ CoCl/SiO₂ has multiple peaks at 360–800 °C, which may come from the reduction of CoO, CoCl₂, and surface Co silicate. CoN/SiO₂ possesses a broad peak around 290 °C which can be attributed to Co₃O₄ and CoO. In contrast, all the TPR profiles of the three S doped Co/SiO₂ catalysts have a sharp peak at 493–506 °C, similar to our recently reported chiral selective CoSO₄/SiO₂ catalyst.²⁶ In addition, it is observed that the CoO_x peaks of undoped catalysts become significantly smaller after S doping, and new peaks around 800 °C appear on CoCl/SiO₂/S and CoN/SiO₂/S. This observation suggests the formation of Co hydrosilicate or surface Co silicate,²⁷ while the 797 °C peak of CoACAC/SiO₂ becomes smaller. Lastly, UV-vis diffuse reflectance spectroscopy was used to probe the surface chemistry of the catalysts. As shown in Fig. S2 in the ESI†, CoACAC/SiO₂ has two peaks around 570 and 650 nm, suggesting the formation of surface Co silicates.²⁸ The spectrum of CoCl/SiO₂ shows two broad peaks at 550 and 720 nm, indicating the presence of CoO_x and CoCl₂. The spectrum of CoN/SiO₂ is similar to that of Co₃O₄, having two broad peaks at ~400 and 720 nm, which can be assigned to the transitions of octahedral configured Co³⁺ ions.²⁹ In contrast, all the three S doped Co/SiO₂ catalysts have a broad peak around 535 nm, similar to that of CoSO₄, and this suggests the existence of Co species bonded to SO₄²⁻.

Doping sulfate ions into metal oxides has created various solid acid catalysts, such as SO₄²⁻/ZrO₂, SO₄²⁻/TiO₂, and SO₄²⁻/Fe₂O₃.^{30,31} Based on our characterization of the Co/SiO₂ catalysts, we propose the following mechanism to explain their (*n,m*) selectivity in SWCNT growth. As shown in Fig. 3, undoped Co/SiO₂ catalysts contain CoO_x, Co hydrosilicate, and surface Co silicates, which are evident from their H₂-TPR profiles and UV-vis spectra. Surface Co silicates on CoACAC/SiO₂ and CoCl/SiO₂ would be reduced and nucleated into small Co nanoparticles,



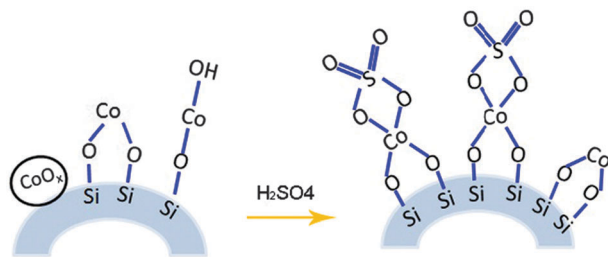


Fig. 3 Schematic illustration of changes in Co species on Co/SiO₂ catalysts caused by S doping.

which are selective toward small-diameter SWCNTs,¹⁷ as shown in Fig. 1. CoO_x on CoN/SiO₂ is reduced to form large Co particles, which are not selective to SWCNTs. Doping S through H₂SO₄ leads to the formation of chelating bidentate SO₄²⁻, where one S atom is linked to one Co atom through two O atoms, a common structure found in sulfate promoted metal oxide catalysts.³⁰ This is supported by the sharp peaks at 493–506 °C in the TPR profiles and the broad peak around 535 nm in the UV-vis spectra. Windle *et al.* showed that adding thiophene or carbon disulfide in gas phase would restrict the growth of Fe particles for selective growth of (9,9).³² Similarly, we propose that the co-existence of S atoms near Co atoms may limit the nucleation of Co and/or form Co–S compounds, which changes the selectivity of the catalysts to favor the formation of (9,8). As discussed in our previous study,²⁶ the selectivity toward (9,8) may be attributed to the close match between carbon caps and the most stable Co particles in their size range, as well as the higher growth rate of high chiral angle tubes.³³ As active Co nanoparticles are embedded under or near SiO₂ surface, we are still unable to quantify their size and composition in transmission electron microscopy analysis (see Fig. S3 in the ESI†). Furthermore, we propose a reaction between H⁺ ions and CoO_x, releasing Co ions to form well dispersed Co hydrosilicate and surface Co silicate on SiO₂, which would increase the selectivity toward SWCNTs. This is suggested by the increased SWCNT selectivity of CoN/SiO₂/S. To further verify the proposed mechanism, we doped CoN/SiO₂ using (NH₄)₂SO₄. We expected the same effect from SO₄²⁻, but selectivity toward SWCNTs may be compromised since NH₄⁺ is less reactive than H⁺. As shown in Fig. S4–S7 in the ESI,† (NH₄)₂SO₄ doped CoN/SiO₂ can result in the growth of (9,8) nanotubes because of S doping. However, it is less selective to SWCNTs as compared to CoN/SiO₂/S. This provides strong credibility to our proposed mechanism.

In conclusion, we have demonstrated a method to convert three types of Co/SiO₂ catalysts, which are either inactive for the SWCNT growth or only selective to small-diameter nanotubes, into chirally selective catalysts to grow SWCNTs enriched with large-diameter (9,8) tubes (up to 40.5%) by doping catalysts with S. We also proposed in the mechanism that S atoms near Co atoms assist the formation of Co nanoparticles which are selective to (9,8) tubes. Moreover, H⁺ ions may react with CoO_x to form well dispersed Co hydrosilicate and surface Co silicate on SiO₂, which increases the selectivity to SWCNTs.

This work is supported by National Research Foundation (NRF-CRP2-2007-02) and Ministry of Education, Singapore

(MOE2011-T2-2-062). We thank Prof. Qiang Zhang of Tsinghua University for the TEM analyses of our samples.

Notes and references

- R. H. Baughman, A. A. Zakhidov and W. A. de Heer, *Science*, 2002, **297**, 787–792.
- S. Irlé, Y. Ohta, Y. Okamoto, A. J. Page, Y. Wang and K. Morokuma, *Nano Res.*, 2009, **2**, 755–767.
- H. Dumlich and S. Reich, *Phys. Rev. B*, 2010, **82**, 085421.
- E. C. Neyts, A. C. T. van Duin and A. Bogaerts, *J. Am. Chem. Soc.*, 2011, **133**, 17225–17231.
- Q. Yuan, Z. Xu, B. I. Yakobson and F. Ding, *Phys. Rev. Lett.*, 2012, **108**, 245505.
- D. A. Gómez-Gualdrón, G. D. McKenzie, J. F. J. Alvarado and P. B. Balbuena, *ACS Nano*, 2012, **6**, 720–735.
- M. F. C. Fawoo, A. M. Bonnot, H. Amara, C. Bichara, J. Thibault-Pénisson and A. Loiseau, *Phys. Rev. Lett.*, 2012, **108**, 195503.
- C. L. Cheung, A. Kurtz, H. Park and C. M. Lieber, *J. Phys. Chem. B*, 2002, **106**, 2429–2433.
- A. G. Nasibulin, P. V. Pikhitsa, H. Jiang and E. I. Kauppinen, *Carbon*, 2005, **43**, 2251–2257.
- S. M. Bachilo, L. Balzano, J. E. Herrera, F. Pompeo, D. E. Resasco and R. B. Weisman, *J. Am. Chem. Soc.*, 2003, **125**, 11186–11187.
- S. Lim, D. Ciuparu, Y. Chen, L. Pfefferle and G. L. Haller, *J. Phys. Chem. B*, 2004, **108**, 20095–20101.
- Y. Miyauchi, S. Chiashi, Y. Murakami, Y. Hayashida and S. Maruyama, *Chem. Phys. Lett.*, 2004, **387**, 198–203.
- B. Wang, C. H. P. Poa, L. Wei, L. J. Li, Y. Yang and Y. Chen, *J. Am. Chem. Soc.*, 2007, **129**, 9014–9019.
- Y. Chen, L. Wei, B. Wang, S. Y. Lim, D. Ciuparu, M. Zheng, J. Chen, C. Zoican, Y. H. Yang, G. L. Haller and L. D. Pfefferle, *ACS Nano*, 2007, **1**, 327–336.
- B. Wang, Y. Yang, L. J. Li and Y. Chen, *J. Mater. Sci.*, 2009, **44**, 3285–3295.
- H. Wang, B. Wang, X. Quek, L. Wei, J. Zhao, L. Li, M. Chan-Park, Y. Yang and Y. Chen, *J. Am. Chem. Soc.*, 2010, **132**, 16747–16749.
- N. Li, X. M. Wang, S. Derrouiche, G. L. Haller and L. D. Pfefferle, *ACS Nano*, 2010, **4**, 1759–1767.
- M. S. A. He, A. I. Chernov, P. V. Fedotov, E. D. Obraztsova, E. Rikkinen, Z. Zhu, J. Sainio, H. Jiang, A. G. Nasibulin, E. I. Kauppinen, M. Niemela and A. O. I. Krause, *Chem. Commun.*, 2011, **47**, 1219–1221.
- M. Fouquet, B. C. Bayer, S. Esconjauregui, R. Blume, J. H. Warner, S. Hofmann, R. Schlogl, C. Thomsen and J. Robertson, *Phys. Rev. B*, 2012, **85**, 235411.
- B. Liu, W. Ren, S. Li, C. Liu and H. M. Cheng, *Chem. Commun.*, 2012, **48**, 2409–2411.
- L. Ci, J. Wei, B. Wei, J. Liang, C. Xu and D. Wu, *Carbon*, 2001, **39**, 329–335.
- W. Ren, F. Li, S. Bai and H. M. Cheng, *J. Nanosci. Nanotechnol.*, 2006, **6**, 1339–1345.
- J. Wei, H. Zhu, Y. Jia, Q. Shu, C. Li, K. Wang, B. Wei, Y. Zhu, Z. Wang and J. Luo, *Carbon*, 2007, **45**, 2152–2158.
- B. Yu, C. Liu, P. X. Hou, Y. Tian, S. Li, B. Liu, F. Li, E. I. Kauppinen and H. M. Cheng, *J. Am. Chem. Soc.*, 2011, **133**, 5232–5235.
- W. E. Alvarez, B. Kitiyanan, A. Borgna and D. E. Resasco, *Carbon*, 2001, **39**, 547–558.
- H. Wang, L. Wei, F. Ren, Q. Wang, L. D. Pfefferle, G. L. Haller and Y. Chen, *ACS Nano*, 2013, **7**, 614–626.
- E. v. Steen, G. S. Sewell, R. A. Makhothe, C. Micklethwaite, H. Manstein, M. de Lange and C. T. O'Connor, *J. Catal.*, 1996, **162**, 220–229.
- A. Khodakov, J. Girardon, A. Griboval-Constant, A. Lermontov and P. Chernavskii, *Stud. Surf. Sci. Catal.*, 2004, **147**, 295–300.
- Y. Brik, M. Kacimi, M. Ziyad and F. Bozon-Verduraz, *J. Catal.*, 2001, **202**, 118–128.
- T. Yamaguchi, *Appl. Catal.*, 1990, **61**, 1–25.
- T. Yamaguchi, T. Jin, T. Ishida and K. Tanabe, *Mater. Chem. Phys.*, 1987, **17**, 3–19.
- R. M. Sundaram, K. K. K. Koziol and A. H. Windle, *Adv. Mater.*, 2011, **23**, 5064–5068.
- F. Ding, A. R. Harutyunyan and B. I. Yakobson, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 2506–2509.

