Preparation and characterization of sulfonic acid-functionalized single-walled carbon nanotubes

Luqman Adams\textsuperscript{a}, Aderemi Oki\textsuperscript{a,*}, Tony Grady\textsuperscript{a}, Hylton McWhinney\textsuperscript{a}, and Zhiping Luo\textsuperscript{b}

\textsuperscript{a} Department of Chemistry, Prairie View A&M University, Prairie View, TX 77446, USA
\textsuperscript{b} Microscopy and Imaging Center, Biological Sciences Building West 119, Texas A&M University, College Station, TX 77843-2257, USA

Abstract
A strategy for the functionalization of single-walled carbon nanotubes is reported. The synthesis involved the conversion of fluorinated single-walled carbon nanotubes to the thiolated derivative assisted by phosphorous pentasulfide. The thiol group is then quantitatively oxidized to the sulfonic acid group. The extent of oxidation of the thiol precursor is confirmed using X-ray photoelectron spectroscopy, which proved to be immensely useful to discriminate between the –SH and –SO\textsubscript{3}H with a chemical shift for the sulfur 2p (approx. 5 eV). The functionalized carbon nanotubes were further characterized by infrared spectroscopy, thermogravimetric analysis, and transmission electron microscopy which revealed a significant change in morphology between the fluoro carbon nanotubes, the thiol and sulfonic acid-modified carbon nanotubes.

Keywords
Carbon nanotubes; Sulfonic acid

1. Introduction
The extraordinary properties of carbon nanotubes [1–5] and the vast areas of potential applications, in medicine, based on their ability to interact with biomolecules, such as proteins and oligosaccharides [6,7], as scanning microscope probe tip [8], as hydrogen storage materials [9], as fillers for ultra-high strength composites [10], and as electron field emitters [11] are some of potential areas of applications under exploration. In order to fully realize potential applications of carbon nanotubes in these and other areas, modification of the surfaces either by molecular self-assembly or chemical functionalization is usually required [12–14]. The covalent functionalization appears to offer strategic advantages over noncovalent approaches, such as polymer wrapping [15], bimolecule binding [16], and metal ion binding [17]. The improvement in solubility and better control of the surface chemistry to impart new characteristics on nanoscale materials are easily accomplished through covalent functionalization of carbon nanotubes [18].

Thiol-functionalized multi-wall carbon nanotubes (MWCNTS) have been prepared by exposing the MWCNTs to harsh acid treatment, which induced surface defect sites such as hydroxyl and carboxyl groups that are further reacted with \textsubscript{P}_\textsubscript{4}\textsubscript{S}\textsubscript{10} to afford reaction products such as MWCNT-SH, -CSOH, and -CSSH [19]. Other methods reported in the literature includes modification of surface oxidized groups with SOCl\textsubscript{2} and subsequent reaction with...
2-mercapto ethylbromide [20], or introduction of surface oxidized groups by mechano-
chemical modification followed by reaction with H₂S [21]. Single-walled carbon nanotubes
were oxidized and thiolated by means of amidation reaction activated by SOCl₂ [22], or
converted to the sulfonic acid derivative by the hydrothermal method [23].

In this paper, we report a strategy that does not entail an initial surface oxidation protocol
with neither an acid, nor a mechano-chemical approach to single-walled carbon nanotubes
functionalized with thiol and sulfonic acid groups. This simple and efficient surface-
substitution approach furnishes sulfur containing functionality covalently bonded to the
surface of single-walled carbon nanotubes that is set up for other practical applications.

2. Experimental

The fluorinated single-walled carbon nanotubes (CNT-F) produced by the elemental fluorine
method was provided by Dr. Valery Khabashesku (RICE University). The CNT-F was used
as received after characterization by elemental analysis and thermogravimetric analysis. The
elemental analysis shows a C₄F, formula. The following commercially available reagents
P₄S₁₀, ethanol, sodium metal, sulfuric acid 98%, and HPLC grade isopropyl alcohol were
obtained from Sigma-Aldrich, while hydrogen peroxide 30% (H₂O₂) obtained from
Mallinckrodt was used in this research.

Fourier transform infrared spectroscopy (FTIR, IR-200 Thermo-Nicolet 2.2) (KBr) in the
range 400–4000 cm⁻¹ was used to confirm the covalently bound thiol and sulfonic acid
groups on the CNT surface. Transmission electron microscopy (TEM) with a conventional
15 kV electron microscope was used to analyze the surface morphology of the
functionalized carbon nanotubes. The thermal gravimetric analysis was performed under
nitrogen on approximately 10 mg samples on a Universal V3.4C TA instruments. The run
consisted of a ramp at a steady rate of 10 °C/min from 40 to 800 °C in air. The X-ray
photoelectron spectroscopy (XPS) powder samples were collected with a Perkin Elmer PHI
5600 XPS for the surface analysis using the Mg Kα X-ray line of 1253.6 eV excitation
energy, 300 W, 15 kV. The vacuum base pressure was approximately 1 × 10⁻⁸ Torr. The
adventitious carbon 284.6 eV binding energy line was used as an internal standard.
Spectrometer pass energy of 58.7 eV was utilized. Carbon nanotube samples were set on
indium foil, and then placed on the sample holder for analysis in the XPS surveys and
multiplex. Carbon, oxygen, sodium, fluorine and sulfur analysis were carried out on each
sample. An electron flood gun was used to minimize any charging effects during analysis.

In a typical experiment, 70.0 mg of P₄S₁₀ was stirred for 15 min in a previously prepared
solution of sodium 60.0 mg/ethanol 7 ml to generate hydrogen sulfide which undergoes
further reaction with the sodium ethoxide to produce soluble sodium hydrogen sulfide in the
suspension [24]. To this suspension 22 mg of fluoro carbon nanotubes (CNT-F) was added
and heated at 70 °C for 3 h. The black residue obtained after filtration with a 0.2 μm
membrane was washed with three portions of 10 ml ethanol and dried under vacuum to
afford the sidewall thiolated carbon nanotubes, CNT-SH, 1. Thereafter, the sulfonic acid-
functionalized carbon nanotube was prepared by oxidation of 1 obtained with 30% H₂O₂
(5.0 ml) at 60 °C for 1 h [25]. The product obtained was filtered through a 0.2 μm
membrane, washed with ethanol and re-suspended with stirring in 10% H₂SO₄ (10.0 ml) for
1 h to ensure complete protonation. The oxidation product after filtering through a 0.2 μm
membrane was washed with deionized water until neutral pH and dried at 80 °C for 12 h to
give 2.

Control experiments were designed as follows: (i) 10.0 mg of CNT-F was stirred with 10%
H₂SO₄ (10.0 ml) for 1 h at ambient temperature. The work-up was performed as for 2 above
and (ii) 10.0 mg of CNT-F was reacted with 30% H₂O₂ (5.0 ml) at 60 °C for 1 h. The product was filtered through a 0.2 μm membrane and dried at 80 °C for 12 h.

3. Results and discussion

The thiol 1, and sulfonic acid 2, functionalized CNT were prepared according to the strategy described in Scheme 1.

XPS analysis was performed on pristine CNT, and the following derivatives: CNT-F, CNT-SH, and CNT-SO₃H with the aim of obtaining information on the functional groups on the carbon nanotube surface (Fig. 1(a–d)). XPS data for the control experiments (Fig. 5(a) and (b)) were also analyzed.

The X-ray photoelectron spectrum in the carbon 1s region is shown in Fig. 1(a). The main peaks appear at 284.6 eV with a non-symmetrical edge at the higher binding energy side of the peak. This has been generally referred as the sp² C=C/sp³ C–C bonding [26]. The pristine CNT shows a peak at ~291 eV. This is typical of the position for carbon π–π* shake up satellite peak of the sp²-hybridized carbon atoms which is more prominent in CNT-SH, 1 compared to the CNT-SO₃H, 2 functionalized carbon nanotube. The peak at ~288.5 eV confirms the C–F bond [26].

In Fig. 1(b), the fluorine 1s region further distinguishes the C–F bond of the CNT-F from the pristine CNT. The absorption peak in this region for the CNT-SH, 1 shows a strong band at ~685.2 eV associated with sodium (NaF) [27] byproduct distinguishable from the CNT-F at ~687.2 eV [26], which is further evidence for sidewall fluoro group displacement. The F 1s XPS spectra for 1 and 2, functionalized CNT shows very weak peaks centered at ~687 eV suggesting only trace amounts of starting CNT-F compound particularly in the product CNT-SH, 1.

In Fig. 1(c), the bands in the sulfur 2p region as expected indicate no sulfur moiety for the CNT and CNT-F compounds. Interestingly, the CNT-SH shows two separate peaks centered at ~163.5 and ~169.5 eV. The peak at 169.5 eV is assigned to a higher oxidation state of sulfur, CNT-SO₃H, 2 [25], while the peak at 163.5 eV shows a reduced form of sulfur as in a mercaptan, CNT-SH, 1 [21,25].

The oxygen 1s region, Fig. 1(d), shows that all the peaks occur between 535 and 530 eV with peak maxima at ~532.5 eV. The interpretation suggests that the oxygen is in the –2 oxidation state for the entire specimen.

Thermal degradation of the samples was monitored with TGA as shown in Fig. 2(a) and (b). The CNT-SH, 1 analysis, Fig. 2(a), display a gradual trend in decomposition from 40 to 430 °C with a weight loss of about 20%, probably coming from the loss of the adsorbed volatiles on the functionalized carbon nanotubes. This is followed by a sharp weight loss between 430 and 480 °C probably due to the loss of the covalently bonded thiol side group on the CNT framework with a weight loss of about 28%. In the CNT-SO₃H, 2 analysis, Fig. 2(b), a gradual trend in weight loss from 40 to 300 °C, arising from adsorbed volatiles molecules on the carbon nanotubes. This is followed by another weight loss between 300 and 560 °C due to decomposition of covalently bonded SO₃H side groups on the carbon nanotubes. The SO₃H may have further decomposed to release oxygen to the environment which ultimately led to sharp increase in weight loss between 560 and 640 °C attributed to the CNT framework decomposition in the presumably oxygen-rich nitrogen atmosphere.

Infrared spectroscopy analysis, Fig. 3(a) and (b), shows a mode centered at about 1540 cm⁻¹ assigned to stretching mode of the C=C double bond that forms the framework of the carbon
nanotubes sidewall. Fig. 3(a) shows characteristic modes centered at about 700 and 1145 cm\(^{-1}\) for the \(-\text{SH}\) group in 1, while the \(-\text{SO}_3\text{H}\) group in 2, Fig. 3(b) is shown by the band centered at 1202 cm\(^{-1}\). The broad absorptions centered between 3000 and 3500 cm\(^{-1}\) may also be attributed partly to adsorbed solvents.

TEM analysis assists with direct imaging of the sidewall modification of the CNT-F. Fig. 4(a–c) shows the TEM images of the CNT-F, CNT-SH, and CNT-SO\(_3\)H, respectively, placed on carbon-coated copper grid. In all three functionalized single-walled carbon nanotubes, the TEM images show evidence of small kinks on the sidewalls, which appears more pronounced in the thiolated CNT (<0.5 nm). In addition, the CNT-SH 1 carbon nanotubes also appears to be aligned together probably through the thiol groups. This alignment appears disrupted when \(-\text{SH}\) in 1 is oxidized to the \(-\text{SO}_3\text{H}\) group in 2.

Control experiments of CNT-F with 10% H\(_2\)SO\(_4\) and 30% H\(_2\)O\(_2\), respectively, were performed to elucidate if the former mild acid medium can introduce on the CNT surface SO\(_3\)H groups, while the later is to investigate the fate of the fluoro substituent in the event of successful sulfonation under the mild acid control conditions. The carbon 1s region in Fig. 5(a) shows a band centered at about 288.5 eV for the starting CNT-F (similar to Fig. 1(a)) suggesting that under the control experimental conditions the fluoro group is not displaced. Furthermore, sulfur 2p region in the control, Fig. 5(b), does not show evidence of introduction of sulfur-related peaks under the mild 10% H\(_2\)SO\(_4\) conditions. Although there is an apparent offset in the maxima of the binding energy between 1 and 2 in Fig. 1(c), the peaks are however centered at about the same point seen also in Fig. 5(b).

4. Conclusions

A route utilizing milder conditions starting from CNT-F offers the opportunity for substitution with other groups. XPS analysis is highly informative in tracking the surface chemistry occurring in these reactions, and has assisted with assignment of the \(-\text{SH}\) and \(-\text{SO}_3\text{H}\) peaks [21,25]. The result of this study suggests that the NaSH generated \textit{in situ} can be used to introduce a thiol group by displacement of fluoro group on carbon nanotubes. Of significance is the alignment of thiolated CNT-SH as seen from the TEM in Fig. 5(b) which also suggests the potential and possibility of exploiting such surface to deposit proteins, enzymes, and other biomolecules.

Acknowledgments

The PIs acknowledge the support from NIH-NIAMSD Grant #AR049172, NIH-NIGMS RISE Grant #1 R25 GM078361-01, the Welch Foundation and the US Air Force Research Laboratory-Minority Leaders Nanocomposite project.

References

Fig. 1.
XPS spectra in the regions of (a) C 1s, (b) F 1s, (c) S 2p, and (d) O 1s for SWCNT, CNT-F, CNT-SH, and CNT-SO$_3$H.
Fig. 2.
TGA analysis for (a) CNT-SH and (b) CNT-SO$_3$H.
Fig. 3.
FTIR analysis for sulfur-functionalized nanotubes (a) CNT-SH, 1 and (b) CNT-SO$_3$H, 2.
Fig. 4.
TEM images of (a) CNT-F, (b) CNT-SH, and (c) CNT-SO$_3$H.
Fig. 5.
XPS spectra in the regions of (a) C 1s and (b) S 2p for CNT-F control reaction (in blue and green) overlaid with actual CNT-SH, 1 (purple) and CNT-SO$_3$H, 2 (black). (For the interpretation of the reference to colour in this figure legend, the reader is referred to the web version of this article.)
Scheme 1.

\[
\begin{align*}
\text{CNT-F} + \text{NaSH} &\rightarrow \text{CNT-SH} + \text{NaF} \\
\text{CNT-SH} + 3\text{H}_2\text{O}_2 &\rightarrow \text{CNT-}\text{SO}_3\text{H} + 3\text{H}_2\text{O}
\end{align*}
\]