Directed self-assembly of carbon nanotubes into microspheres

Mi hye Choi¹ and Young nam Cho¹,²*

¹Biomarker Branch, National Cancer Center, Goyang, South Korea
²Cancer Biomedical Science, Graduate School of Cancer Science and Policy, Goyang, South Korea

Abstract

Carbon nanotube microspheres (CNMs) were prepared using the inherent self-organizing characteristics of the nanotubes that primarily rely on electrostatic interaction, not the assistance of any polymers or surfactants. They permit significant linkage with a variety of chemical species such as biomolecules, metal nanoparticles, and fluorescent dyes. The cationic species of guest molecules facilitate individual nanotubes to coalesce and merge together into new colloidal structures. These novel CNMs create new technical opportunities for a wide range of applications. In particular, the capability of conjugating with a broad number of chemical species would be a fundamental basis for development of bio-electronics and biomedical applications.

Introduction

There has been great interest in developing novel strategies regarding fabrication, characterization, and possible applications of carbon nanotubes (CNTs) [1-3]. Recent advancements extend their applications to areas as diverse as electronic devices, environmental fields, and the biotechnology industry given their unique electrical, mechanical, and thermal characteristics as well as high surface area [4-6]. However, the difficulties associated with poor solubility and self-aggregation of pristine nanotubes present limitations in further processing and often restrict the practical use of CNT, especially for micro / nanoscale devices. A number of studies have focused on manipulating the self-assembly nature of nanotubes. The most common techniques involve the use of dispersants such as surfactants, solvents, polymers, or biomolecules with the hope of facilitating stable dispersion through covalent or non-covalent functionalization of CNTs [7-9]. Another approach involves processing highly aligned nanotubes in the form of bundles with morphological diversity to fabricate mechanically strong and conductive composites such as fibers, pellets, sheets, and yarns [10-15]. Such densely packed and well-aligned microscopic forms of CNTs possess large reactive areas and are durable, while simultaneously retaining fundamental properties of individual nanotubes. A recent advancement was the development of CNT hybrid materials with planar or spherical geometries [16-10]. In fact, the addition of the nanotubes into polymers usually alters the physicochemical properties of the films, endowing a remarkable combination of conductivity and mechanical robustness [19,20]. Several groups have pursued CNT-based capsules by adopting layer-by-layer (LBL) deposition techniques [21-24]. CNT capsules can be produced through the self-assembly of oppositely charged polymers and CNTs in a water-in-oil emulsion system or with the use of colloidal templates. The size and flexible functionalities of the microcapsules rely on the diameter of the template particles and chemical species used. Notably, hollow microcapsules with large interior spaces are useful for the efficient loading and release of a variety of guest molecules [21,25]. Despite these advancements, very significant challenges remain, including complications in fabrication and the collapse of the spherical architecture after the removal of core templates. In previous study, our group has demonstrated the formation of mechanically robust, stable, surfactant-mediated carbon nanotube microspheres using a series of surfactants with cationic, anionic, and non-ionic charges [26]. In parallel with these advances, we provide a simple approach for the large-scale construction of spherical, water-stabilized CNT microspheres (CNMs) with a mean diameter of ~ 17 µm as illustrated in Figure 1a.

Additionally, we report the formation of CNM complexes incorporating various chemical species including biomolecules and metallic nanoparticles, Figure 1b. Our novel CNMs were obtained in a ‘green’ water-based process since surfactants, polymers, harsh chemicals, or solid templates were not used during fabrication.

Experimental

The fabrication of CNMs

The fabrication of water-soluble CNMs was prepared as follows. Four different types of multi-walled carbon nanotubes (MWCNT; Nanolab Inc, MA) were used: (1) pristine; (2) amine-modified nanotubes (diameters of 30 nm and lengths of 1 ~ 5 µm); (3) carboxyl (COOH) groups-decorated (diameter of 30 nm and lengths in the range of 1 ~ 5 µm); and (4) short COOH-decorated nanotubes (diameters of 30 nm and lengths in the range of 500 nm ~ 1 µm). According to the company, carboxylated nanotubes have 2-7 wt % COOH by titration after functionalization and metal catalyst impurities would be around 3-4%. Pristine and functionalized nanotube dispersion with typically

*Correspondence to: Cho Y, Biomarker Branch, National Cancer Center, Goyang, Gyeonggi-do 410-769, South Korea, E-mail: yncho@ncc.re.kr

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0.1 ml/mL were prepared by adding MWCNTs into milli-Q water and then sonicated using high power ultrasonics for 5 min. The dispersion was subsequently treated in an ultrasonic bath for 1 hr and then subjected to high-power sonication again for 5 min. After this, the pH of samples was adjusted to 2, 7, or 11 by adding 0.1 M HCl or NaOH to allow unstable bundles to be modulated into spherical morphology. The final samples were further purified with DI water via centrifugation and redispersion cycles. After repeating the process three to five times, resulting materials were dried in a vacuum oven at 40 °C. Through a series of these steps, only carboxyl (COOH) groups-decorated (diameter of 30 nm and lengths in the range of 1 ~ 5 µm) nanotubes have allowed obtaining stable spherical agglomerates without any damage to their resulting structure during the washing and drying treatments that is further demonstrate by SEM. Continually, the conjugation of poly-L-lysine (PLL; Mw=70,000 ~ 150,000; Aldrich), bovine serum albumin (BSA), or amine-functionalized gold nanoparticles (~ 100 nm Au-NH₂; Nanopartz, Inc) within CNMs was achieved by mixing the aqueous solution containing COOH-decorated nanotubes (diameter of 30 nm and lengths in the range of 1 ~ 5 µm) of 0.1 mg/mL with 100 µM PLL, 100 µM BSA, or 50 µM Au-NH₂ at various pH to allow spontaneous chemical adsorption. Unbound materials were thoroughly removed by centrifugation. The fluorescent dye-labeled CNMs were obtained by mixing 1 mM FITC (Aldrich), 1 mM rhodamine 123 (Aldrich), 100 µM of FITC-PLL (Nanocs), or 100 µM of Rhodamine-BSA (Nanocs) with 0.1 mg/mL carboxylic acid-decorated nanotubes solution at various pH. The solution was then centrifuged and reconstituted until no fluorescence was detected in the supernatant. Finally, fluorescent CNMs were air-dried at room temperature.

The characterization of CNMs

Sample of CNMs were dried onto glass coverslips, and sputter-coated with AuPd (Hummer II sputter-coater, Anatech USA, Union City, CA) prior to imaging with a FEI NOVA nanoSEM (FE Company, Hillsboro OR). Imaging was done with an Everhart-Thornley (ET) detector or Through-the-Lens high resolution detector (TLD) using 5kV accelerating voltage. The focused ion beam (FIB) ablation, used to analyze the inner structure of CNMs, was done with an FEI Quanta 3D FEG dual beam FESEM. Pt deposition protected the CNM surface during FIB milling. Milling parameters were 30kV, 10 mm WD and 520 till. Images were captured using the ET detector at 5kV accelerating voltage.

Results and Discussion

The intertube interactions as a consequence of van der Waals attraction and hydrophobic bonds cause them self-aggregate in both organic and an aqueous medium. Moreover, electrostatic interaction through excess charges present on nanotube surfaces induces an additional aggregation which might involve their processing in stable spherical assembly with significant structural integrity. Indeed, our results suggest that electrostatic forces seem to be critical for determining controllable colloidal shapes as microsphere stabilization depends on the transition of the carboxyl group (COOH) present on the nanotubes into ionized carboxylic acid species (COO⁻). Notably, a subtle difference in the density of the carboxyl group on the nanotubes has caused a significant impact on the geometric structure. Greater attractive interaction was observed at high pH, where increased deprotonation of –COOH groups preferentially react with adjacent positive ions derived from the basic aqueous solution. Scanning electron microscopy (SEM) was used to monitor the CNM formation. High magnification images show CNMs with tightly packed but relatively smooth exterior surfaces (Figures 2a and 2b). The structures retain the rigid form while the aggregates of CNTs making up the structure still provide for large surface area. The presence of reactive surfaces provides several technical or scientific advantages given that: i) increasing the active surface area would be beneficial for supporting a variety of molecular species and ii) a randomly-connected nanotube network permits the free flow of gas or liquids through the CNMs.

In fact, tailoring a surface-to-volume ratio within the microspheres would be essential for extending the application of CNMs through the incorporation of reactive species. Thus, it is also necessary to understand the inner structure of CNMs. Focused ion beam (FIB) milling was used to ablate a portion of single CNM so as to observe transverse CNM in cross-section. This technique provided a unique view inside the spheres revealing the distribution of the nanotubes, the presence of defects that might exist, and the impurities that might be created during the nucleation and growth of the microspheres. As shown in Figures 2d and 2e, the interior analysis of microspheres confirmed the absence of crystalized chemical impurities and apparent structural defects. However, these colloidal aggregates tend to collapse, flatten, or become stretched into the substrate surface when pH is decreased [Figure S1] in Supporting Online Materials (SOM)]. Indeed, protonation / deprotonation process of negatively charged carboxylic acid species is essentially responsible for the dissociation/assembly of CNMs.

The current fabrication method for producing the CNM results in a size distribution. There are a number of factors that are crucial for governing the fabrication process, including the concentration, length, and surface charge of the nanotubes as well as reaction time and temperature. The length of the nanotubes appears to be more relevant in determining the final diameter and morphology of the microspheres than the concentration and experimental conditions.
Using short COOH-decorated nanotubes (500 nm ~ 1µm) resulted in a definite trend toward a decrease in the CNM average diameter (3.3 µm) with a variety of different morphologies. If the density of negatively charged species (that is, carboxylic acid groups) is more dependent on nanotube length, this could result in a decrease in ionic strength when short CNTs are used. This may account for the smaller CNM size and less consistent spherical morphology (Figure S2) in Supporting Online Materials (SOM). Neutral charged nanotubes and amine-modified MWCNTs (30 nm diameter, 1-5 µm length, >95% purity; Nanolab Inc, MA) were also used for the formation of microspheres. The lack of a specific chemical affinity resulted in the construction of a thinly spread nanotube film rather than the spherical particulate structures produced at various pHs (Figure S3) in Supporting Online Materials (SOM). On the other hand, amine (NH₂) functionalized nanotubes also showed an entangled flat structure instead of forming the spherical assembly (Figure S4) in Supporting Online Materials (SOM). The reason is likely attributed to the insufficient charge due to the lack of active amine moieties on the nanotubes. As a derivative of our COOH functionalized nanotubes, amine-decorated nanotubes were produced through the reaction of the carboxyl COOH group on the nanotubes with ethylene diamine (H₂N-CH₂-CH₂-NH₂) as a coupling agent that would be responsible for the decrease in the amine concentration. Currently, the mechanism underlying these microspheres is not discovered yet. However, CNM formation is closely related to the surface charge densities around the nanotube. On the other hand, Figure 2f showed the size distributions of CNMs that were obtained by measuring individual particles by SEM analysis. A statistical histogram revealed a wide range of size distribution, with the average diameter...
of 17.6 µm. To elucidate the mechanism behind the effect on the particle formation and size, further detailed experimental studies are currently underway in our group. It was recently reported that novel constructs possessing a comparable size to biological structures can readily interact with diverse organic samples – including both chemical and biological moieties [27-30]. Therefore, we investigated whether our novel CNMs would provide a useful and efficient matrix for the simultaneous encapsulation of chemical species within the interior and attached to the exterior of the particles. Indeed, CNMs would make promising candidates for application in broad areas if they were shown to effectively entrap and confine molecules. CNMs composed of nanotubes would be expected to offer significant advantages, especially in biomedical applications: i) relatively long nanotube arrays with an enormous surface area are ideally suited for adsorption of multiple cargo molecules, ii) biocompatible coatings composed of these spheres would offer ideal micro-environment for cell growth and tissue formation, iii) chemical adsorption with biomolecules would produce significantly greater storage stability preventing undesirable degradation or denaturation, and iv) CNMs could be utilized as novel diagnostic tools when combined with imaging agents, such as luminophore or magnetic functionalities. Indeed, the simple and uncomplicated fabrication of CNM using individual nanotubes as molecular building blocks offers finer control over the hybrid compositions and a more standardized production of nanomaterials. As a proof-of-concept experiment in this vein, we functionalized CNMs with representative fluorescent derivatives such as both fluorescein (FITC) and rhodamine (Rho) that were initially conjugated with poly-L-lysine (PLL) and bovine serum albumin (BSA) respectively. The objective was to enhance electrostatic stabilization and simultaneously achieve co-loading of these desired molecules to form large protein complexes. The decorated CNMs were visualized by SEM, conventional bright-field, and fluorescence microscopy (Figures 3 a-f).

SEM images verified the general construction of the CNM-PLL/FITC complexes but did not reveal whether the proteins were contained within the interior or attached to exterior surfaces of CNMs. However, the complementary evidence provided by fluorescent imaging presented strong fluorescence from FITC-PLL associated with the CNMs (Figure 3c). Like with FITC-PLL loading, the cationic characteristics of Rho-BSA complexes resulted in their binding to the negatively charged surface of the nanotubes by electrostatic forces at physiological pH and confirmed the co-adsorption of negatively charged BSA molecules as shown in Figure 3f. In control experiments, the conjugation with only negatively charged FITC molecules without the PLL led to a very weak signal. As another specific application of such interesting microspheres, we detailed the functionalization of CNMs with positively charged particles, particularly amine-terminated gold nanoparticles (Au-NH₂) with ~100 nm in diameter. SEM images have verified the existence of highly concentrated and uniformly dispersed gold particles at pH 11 (Figure 4). However, the cross-sectional images revealed that most of gold particles were presented on the surface rather than inside of CNM microsphere, assuming that the accumulation of more gold particles inside CNMs might be interrupted by their relatively large size (Figure 4d). And more importantly, novel CNM complexes conjugated with gold nanoparticles exhibited the variation in loading efficiency that can be tuned by altering pH. These results suggested that positively charged moieties derived from guest molecules are definitely responsible for enhancing spherical stability but do not seem to be a critical factor in initiating the microsphere formation.

In parallel experiments, negatively charged silica particles (150 nm in a diameter) were used as a control (data not shown). As expected, no detectable quantity of silica particles was adsorbed. Although colloidal stability was not affected, the charge repulsion of these particles inhibited the particle attachment.

Conclusion

In summary, we offer a convenient method for preparing carbon nanotube microspheres made of only carbon nanotubes. This procedure is in sharp contrast to conventional methods which focus on breaking bundles into individual nanotubes. Rather we exploit the spontaneous self-aggregation characteristics of nanotubes to merge them into colloidal structures by manipulation of the pH. Such thermodynamically and kinetically efficient fabrication offers new promises in discovery and design of CNM hybrids by simply engineering the structural geometry or chemical integrity. In addition,
the fact that our approach is accessible to innumerable guest molecules extends this new variation of CNM complex to diverse endeavors, especially chemistry, biotechnology, electronics, and of great importance - therapeutic approaches.

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Compliance with Ethical Standards

Conflict of Interest

The authors declare that they have no conflict of interest.

Supporting Information Available

SEM images of CNMs and nanotube surface produced from pristine multiwalled carbon nanotubes and the nanotube surface produced from amine.

References


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