Evaluation of Tensile Properties of Alcohol-Dispersed Carbon Nanotube/Low-Density Polyethylene Composites

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Abstract-The agglomeration of carbon nanotubes makes the formation of composites difficult as it hinders uniform distribution of carbon nanotubes within the matrix. This paper investigates the effects of alcohol as a de-agglomerating chemicals i.e surfactant or dispersing agent on the carbon nanotubes. The carbon nanotubes were purified, dried and reconstituted with the dispersing agents followed by mechanical agitation using sonicator. The formed composites, after embedding the dispersed carbon nanotubes in the Low Density Poly-Ethylene (LDPE) were subjected to mechanical test. The Ultimate Tensile Strength of 4.6MPa and percentage elongation of 25.75% of 0.5%wt CNT/LDPE-alcohol-dispersed composites were achieved. The interfacial interaction of the carbon nanotube reinforcement in Low Density Polyethylene was studied using scanning electron microscope. The tubes were seen evenly distributed and perfectly embedded within the low density polyethylene.

Keywords: Carbon, nanotube, alcohol, dispersion, polyethylene

I. INTRODUCTION
Carbon nanotubes (CNT) are known to have exceptional mechanical, thermal and physical properties and this makes researchers to explore and harness it to improve the properties of polymer matrix [1]. CNT carbon atoms are covalently bonded, wrapped with graphene sheets to form seamless cylinders through sp² hybridization with planes of bond stronger than sp³ hybridization, as in diamond. CNTs are axially strong with Young’s Modulus and tensile strength in the range 270 – 950 GPa and 11 – 63 GPa respectively [2].

CNTs possess unique electronic configuration such that single walled CNT chiral structure can be varied to be metallic or semi-conducting. Single-walled CNT exhibit current-carrying strength of about 10,000 cm² V⁻¹ s⁻¹, more than silicon current-carrying strength. CNTs have 1000 times a metal’s electrical conductivity. CNT has been applied as novel constituent materials in electrode for super-capacitor, fuel cell, dye-sensitized solar cell and light-emitting diode [1–2].

The electrical conductivities of disentangled multi-walled carbon nanotube (MWCNT) are 50 times higher than the entangled MWCNTs which may be associated to straight MWCNTs having little defects [3–4].

The entangled MWCNTs have less electrical conductivity which may be as a result of the magnetic field interference in the passing of electric current whereby the field’s interference can increase the current impedance [4].

The dispersion of CNT within the matrix is difficult due to interfacial energy and Van der Waal’s forces within the tubes. Research has shown that with surface-active agents or surfactants which reduces the interfacial energy and increases the repulsive forces within the tubes, CNT can be unbounded and disentangled [1-3].

Hilding et al., 2003 [5], reported that the usual procedure of carbon nanotube dispersion is that surfactant is adsorbed on the surface of the carbon nanotube, followed by ultrasonication for minutes in order to electrostatically separate the carbon nanotubes. But recently, research has shown that surfactant is able to adhere only to the external nanotubes but the internal nanotubes remain agglomerated [6-10]. Therefore, in order to adequately unbundle the agglomerate, mechanical exfoliation prior to surface treatment of the nanotube should be done. It is on this note that ionic surfactants and non-ionic surfactants are proposed for CNT/water soluble solutions and CNT/organic solvents respectively [11-13]. Carbon nanotubes, when coated with adsorbed surfactants having high hydrophyle-lyphophyle balance can be easily dispersed in water. This is a non-covalent method which can be used for organic and in-organic dispersion of particles in aqueous solutions [12]. Sodium dodecyl sulphate and sodium dodecyl benzene sulfonic acid are known among the ionic surfactants to decrease the
tendency of CNT agglomeration in water [14–17]. The presence of benzene ring in the surfactant is believed to make it have a better dispersive power. It has also been shown that apart from aromatic-group containing surfactants, naphthenic groups do efficiently disperse CNT [9]. An example of such is aerosol-OS (sodium disopropyl-naphthalene sulfonate) which shows a high surfactant-tube affinity as revealed by spectroscopy analysis [18]. Multiwalled carbon nanotube has been efficiently dispersed in polyethylene glycol using polyethylene oxide-20- sorbitan mono-oleate, a non-ionic surfactant. This dispersion was observed to be as a result of the dispersant tail having carbon double bond [19].

2. EXPERIMENTAL

2.1 Materials and methods
As-synthesized multi-walled carbon nanotube was obtained from Centre for Genetic Engineering and Biotechnology (CGEB), Federal University of Technology Minna. Granules of translucent low density polyethylene of 0.916g/cm³ density was purchased from Nigeria Institute of Leather and Science Technology (NILEST). Ethanol of 0.01ml N/1% acidity/alkalinity was purchased from a chemical shop in Minna, Nigeria. Compression moulding machine and the two-roll mill were obtained in NILEST, Zaria, Nigeria. TEM and SEM were carried out in South Africa.

2.2 Purification of carbon nanotube
Multi-walled carbon nanotubes was purified using 1:100 solid to liquid mixture of CNT and aqueous solution of 3:1 concentrated acetic acid and hydrochloric acid respectively. The mixture was refluxed at 70°C for 24 hours and continuously stirred using a magnetic stirrer at 400rpm. Then, it was cooled, the CNT was filtered and washed using distilled water and dried in oven at 60°C for 12 hours.

2.3 Functionalization of carbon nanotube
The CNTs were functionalized using 1.3 HNO₃/H₂SO₄. The mixture was mechanically agitated on a sonicator for 18 hours. Then, the carbon nanotube was filtered and rinsed repeatedly using distilled water till pH 7 was attained. After attaining neutrality pH, the moist CNT was dried in an oven at 60°C for 12 hours.

2.4 Dispersion of CNT in alcohol
2g of functionalized carbon nanotube was added into a 3.35M alcohol and then sonicated for 2 hours. The dispersed solution of CNT in alcohol was filtered and oven-dried at 60°C for 12 hours.

2.5 Fabrication of CNT/LDPE
Two composite samples of 50g low-density polyethylene with each melt-blended with 0.5g and 1g weight percent alcohol-dispersed CNT were prepared in a two-roll mill operating at 120°C and 50rpm for 10 minutes. The third sample was without CNT. The blended composites were compressed in a square mould at about 100°C in a compressing machine.

Tensile test specimens in a dog-bone shape according to ASTM-D3039 were cut from the samples and clamped in a tensometer. Upon application of force, the load-strain curve was being pen-plotted till the specimen failed and later traced for computation.

2.6 Surface morphology
Transmission electron microscope was used to analyse a drop of alcohol-dispersed CNT placed on TEM grids, (200 mesh) coated with carbon after the CNT-alcohol solution has been ultrasonicated for 20minutes to achieve a better image. Scanning electron microscope (SEM, Leo Supra 55-Zeiss Inc., Germany) was used to determine the level of dispersion of the CNTs in the composites after the tensile test. The scanning was carried out at room temperature after polishing the surface of the specimen.

3. RESULT AND DISCUSSION

3.1 Purification and functionalization
Purification of carbon nanotube became necessary to avoid the interaction of unwanted impurities like amorphous carbon, residual catalyst, carbon nanoparticles that are attached to the carbon nanotube during synthesis by contaminating the process of effective dispersion of carbon nanotube in matrix.

A micro-scale inspection of the carbon nanotubes is presented in the TEM images below.: as-synthesized, purified and functionalized. The bundled size of entangled CNT is more and the unbundled size less in as-synthesized, this is expected because of its pristine nature. The bundled size is reduced in purified and further reduced in functionalized carbon nanotube but the unbundled size increased in both. The metallic impurities, amorphous and carbonaceous materials attached to the CNT have been washed away during the purification and a functional group added during the functionalization for the CNT to react with other materials [20].
3.2 Dispersion of CNT in alcohol.
During the sonication process of the functionalized CNT in alcohol, it was observed with bare-eye that there was a change in the colour of the solution. The longer the sonication time the darker the solution, that is, exfoliation of the carbon nanotubes in the alcohol till it was stable for the period of the two hours. The same nature of observation in the work of Dujardeen et al., 2012 [21].

![TEM image of purified CNT](image1)

![TEM image of functionalized CNT](image2)

![Exfoliated carbon nanotube](image3)

3.3 Tensile test
The table below is the result of tensile test carried out on the samples

<table>
<thead>
<tr>
<th>% CNT</th>
<th>0%</th>
<th>0.5%</th>
<th>1%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Width (mm)</td>
<td>9.6</td>
<td>9.65</td>
<td>9.7</td>
</tr>
<tr>
<td>Thickness (mm)</td>
<td>2.85</td>
<td>2.80</td>
<td>2.8</td>
</tr>
<tr>
<td>Area (mm²)</td>
<td>27.36</td>
<td>27.02</td>
<td>27.16</td>
</tr>
<tr>
<td>Force (Newton)</td>
<td>92.25</td>
<td>123.00</td>
<td>97.75</td>
</tr>
<tr>
<td>UTS (MPa)</td>
<td>3.37</td>
<td>4.55</td>
<td>3.60</td>
</tr>
<tr>
<td>Strain</td>
<td>0.2756</td>
<td>0.2650</td>
<td>0.2250</td>
</tr>
<tr>
<td>% Elongation</td>
<td>27.56</td>
<td>26.50</td>
<td>22.50</td>
</tr>
<tr>
<td>Young’s Modulus (MPa)</td>
<td>12.23</td>
<td>17.17</td>
<td>16.90</td>
</tr>
</tbody>
</table>

3.3.1 Effect of CNTs on the Ultimate Tensile Strength of LDPE/CNT composite
The UTS value of LDPE increased to 135% and 107% when 0.5wt% and 1wt% alcohol-dispersed CNT were added. A similar trend is observed in the work of Nazlia et al., 2007 [23], where styrene butadiene stress value or tensile strength was increased to 121% and 170.26% upon adding 1wt% and 10wt% CNT respectively. Xiao et al., 2006 [24], had 56% increment in the ultimate tensile strength of ordinary LDPE when 10wt% carbon nanotube was added.

![Ultimate Tensile Strength of Alcohol-dispersed CNT/LDPE Composites](image4)

3.3.2. Effect of CNTs on the Strain of LDPE/CNT composite
Strain reduction was experienced in alcohol-dispersed CNT/composite to 96.2% and 81.64% when 0.5wt% and 1wt% CNT were added respectively. It agrees with the trend reported in literatures [23-24].

The more the CNT content, the lower the strain value. This may be due to the rigid nature of CNT which hinders the elongation of the composites. Also, much of the elongation is from the polymer matrix. The measured elongation is an indication that composites stiffness can be improved by CNT.

The entanglement of MWNT is high as a result of the strong cohesive energy between the tubes which is in the order of 36 kT [22].
3.3.3. Effect of CNTs on the Young’s Modulus of LDPE/CNT composite

The Young’s modulus of ordinary LDPE has increased by 40% and 31% when 0.5% and 1% alcohol-dispersed carbon nanotube were added respectively.

3.4 Surface Morphology of the CNT/LDPE Composites

The figure below is a result of the scanning electron microscope. The result shows that there is a cordial interfacial bonding between the particles of the carbon nanotube and the LDPE as it can be observed that the tubes are effectively embedded in the LDPE without bundles of CNT randomly distributed in the image as shown. This is the effect of the two-roll mill being able to uniformly distribute the carbon nanotube within the LDPE and the efficient disentangling through various processes as outlined above.

4. CONCLUSION

In summary, the effect of carbon nanotube dispersed with alcohol in low density polyethylene has been investigated. The excellent properties of carbon nanotube were harnessed to reinforce the composites. Ordinary low density poly-ethylene properties like young’s modulus, ultimate tensile strength have been improved. This has proven that alcohol may also serve as a dispersing agent. The ultimate tensile strength and the young’s modulus have been increased by 35% and 40% respectively.

REFERENCE


