Dynamic Rheological Studies of Poly(p-phenyleneterephthalamide) and Carbon Nanotube Blends in Sulfuric Acid

Yutong Cao 1*, Zhaofeng Liu 1*, Xianghua Gao 1, Junrong Yu 1, Zuming Hu 1 and Ziqi Liang 2

1 State Key Laboratory for Modification of Chemical Fibers and Polymer Materials, Donghua University, Shanghai 201620, China
2 National Renewable Energy Laboratory, Golden, CO 80401, USA

* Author to whom correspondence should be addressed; E-Mail: lzf@dhu.edu.cn (Z.L.); caoyt@hotmail.com (Y.C.); Tel.: +86-21-6237-3934; Fax: +86-21-6237-3934.

Received: 11 February 2010; in revised form: 17 March 2010 / Accepted: 18 March 2010 / Published: 31 March 2010

Abstract: We have studied the dynamic scanning of liquid-crystalline (LC) poly(p-phenyleneterephthalamide) sulfuric acid (PPTA-H2SO4) solution, and its blend with single-walled carbon nanotubes (SWNTs), by using a flat plate rotational rheometer. The effects of weight concentration and molecular weight of PPTA, as well as operating temperature, on dynamic viscoelasticity of the PPTA-H2SO4 LC solution system are discussed. The transition from a biphasic system to a single-phase LC occurs in the weight concentration range of SWNTs from 0.1% to 0.2%, in which complex viscosity reaches the maximum at 0.2 wt% and the minimum at 0.1 wt%, respectively, of SWNTs. With increasing SWNT weight concentration, the endothermic peak temperature increases from 73.6 to 79.9 °C. The PPTA/SWNT/H2SO4 solution is in its plateau zone and storage modulus (\(G'\)) is a dominant factor within the frequency (\(\omega\)) range of 0.1–10 rad/s. As \(\omega\) increases, the \(G'\) rises slightly, in direct proportion to the \(\omega\). The loss modulus (\(G''\)) does not rise as a function of \(\omega\) when \(\omega < 1\) s\(^{-1}\), then when \(\omega > 1\) s\(^{-1}\) \(G''\) increases faster than \(G'\), yet not in any proportion to the \(\omega\).

Keywords: liquid crystalline (LC) solution; single-walled carbon nanotubes (SWNTs); rheology
1. Introduction

A liquid-crystalline (LC) solution is an orientationally ordered liquid that displays fluidic characteristics of the liquid state while the molecules still maintain a directional order. There are three main classes of LC phases, i.e., nematic, smectic and cholesteric phase, which were originally identified by Friedel et al. according to the degree of order and molecular alignment of each phase [1]. Poly(p-phenyleneterephthalamide) (PPTA), a rigid-rod polymer as shown in Scheme 1, is typical of high tensile strength, remarkable stiffness, and excellent thermal stability [2]. One of the most interesting physical properties of PPTA macromolecules is their strong affinity to self-organize in a suitable solvent such as absolute sulfuric acid (H₂SO₄) and hence form LC phases [3]. Such strong molecular arrangement is attributed to the extreme anisometry of rigid PPTA and disruption of intermolecular hydrogen bonds in PPTA caused by the super-acidity of H₂SO₄. Nematic sulfuric acid LC solution of PPTA, referred to as PPTA-H₂SO₄, with a weight concentration range of 19–20% when heated to 80–95 °C has been used to fabricate high-strength fibers by dry-jet wet-spinning process [4–6]. The unusual rheology phenomena of PPTA-H₂SO₄ LC solution have been reported [7–11], including strong shear-thinning, long relaxation time, and high viscous activation energy. These notable behaviors inspired us to investigate the dynamic rheological properties of the PPTA-H₂SO₄ LC solution system in the spinning process. We then carried out the dynamic scanning of the PPTA-H₂SO₄ LC solution system with a flat plate rotational rheometer and discussed the effects of weight concentration and molecular weight of PPTA, as well as temperature, on such viscoelastic properties as complex viscosity ($\eta^*$), storage modulus ($G'$), loss modulus ($G''$), and loss-tangent ($\tan\delta$). Our results shown in this paper provide some general fundamental insight for the LC spinning process.

Single-walled carbon nanotubes (SWNTs) have been shown to possess extraordinary mechanical, electrical and thermal properties [12–15]. Polymers have been shown to be significantly reinforced with SWNTs [16,17], and some typical examples include pitch-based/SWNT composite carbon fibers [18], PBO/SWNT [19], and polycrylonitrile/SWNT composite fibers [20]. The major challenge, however, remains to achieve uniform and stable dispersions of SWNTs in the polymer matrix with significant nanotube weight-fractions. It is known that SWNTs can be successfully dispersed in sulfuric acid. The nematic structure of SWNT/H₂SO₄ solution has been evidenced by birefringence and rheological curves [21]. It is thus of special interest to investigate whether rigid rod-like polymers (e.g., PPTA) can be reinforced with SWNTs in a large loading ratio, both in the LC solution, which will be of great significance to wet-spinning production of high-performance fibers. For this purpose, LC solution of PPTA/SWNT/H₂SO₄ was prepared, and its thermal and rheological behaviors were studied accordingly. Our study aims to assess the effects of SWNTs on the properties of the spinning solution. Through our studies, we are able to identify the optimal range of SWNTs concentration and spinning temperature for dry-jet wet-spinning process of PPTA fibers.

Scheme 1. Molecular structure of poly(p-phenylenetetraphthalamide) (PPTA).
2. Results and Discussion

2.1. Effect of Weight Concentration of PPTA on Rheological Behaviors

Effects of the variation of weight concentrations of PPTA (PPTA wt%: 18.5, 19 and 19.5%) and temperatures (75, 80, 85, 90 °C) on rheological curves of the PPTA-H$_2$SO$_4$ LC solution are shown in Figure 1. The solution exhibits a typical shear-thinning behavior even at a low shearing frequency ($\omega$), i.e., its $\eta^*$ decreases as $\omega$ increases, which results in the alignment of PPTA chains in parallel. With increasing $\omega$, the density of junction points decreases since the crosslinks break significantly faster than they form owing to their shear-thinning behavior [22]. Such changes of chain configuration and orientation under shearing force indicate that both $G''$ and $\tan\delta$ of PPTA-H$_2$SO$_4$ LC solution would increase as $\omega$ rises. At a high shearing rate, nearly all macromolecular chains orient, implying that $G'$ hardly follows the variation of frequency, since in this regard the first normal stress difference hardly follows frequency variation [23]. However, when weight concentration of the solution increases, density of entanglement crosslinks and intermolecular forces both increase, and then macromolecular chains become more difficult to orient and diffuse, leading to the increase in all the rheological characteristics including $\eta^*$, $G'$, $G''$, and $\tan\delta$. On the other hand, as the temperature increases, thermal motion of macromolecular chains increases, flow resistance decreases, and rheological curves of $\eta^*$, $G'$, $G''$, and $\tan\delta$ move down.

Figure 1. Rheological curves dependences of PPTA-H$_2$SO$_4$ LC solution on shearing frequency ($\omega$), PPTA wt% at 18.5, 19 and 19.5%, and temperatures of (a) 75 °C, (b) 80 °C, (c) 85 °C and (d) 90 °C.
2.2. Effect of Molecular Weight of PPTA on Rheological Behaviors

Supramolecular structure is key to affecting the rheological characterization of the polymer system. Figure 2 displays the rheological behaviours of PPTA-H_2SO_4 LC solution at 19.5 wt% of PPTA as a function of M_w (i.e., 28169, 56170, 68407, and 80135 g/mol) and temperature (i.e., 80, 85, and 90 °C). All four rheological parameters (η*, G', G'', and tanδ) increase with increasing M_w of polymers. Also, the larger M_w is, the longer relaxation time (λ) appears, meaning that oriented molecular chains upon shearing cannot resile, thus showing more distinct shear-thinning behaviour. This is greatly advantageous to obtain high degree of orientation in spinning process and hence obtain fibers with higher tenacity and modulus. Moreover, the larger M_w is, the more entanglement crosslinks are generated, meaning that the dynamic process of disentanglement and entanglement would again take place at less shear rate [24,25]. Hence, M_w of PPTA would be ideally neither high (difficult for molding) nor low (accelerating the relaxation).
Figure 2. Effect of PPTA molecular weight of PPTA-H$_2$SO$_4$ LC solution on rheological curves versus shearing frequency ($\omega$).
2.3. Effect of Weight Concentration of SWNT on Rheological Behaviors

Variation of complex viscosity as a function of SWNT weight concentration (SWNT wt%) in PPTA/SWNT/H₂SO₄ dopes is plotted in Figure 3, showing the nonmonotonic characteristic as a function of frequency and temperature. The $\eta^*$ first gradually increases with increasing SWNT wt%, and then decreases abruptly when beyond ~0.1 SWNT wt%, and arises again when beyond 0.2 SWNT wt% to reach about the same viscosity for three temperature-dependent curves. In general, this concentration dependent viscosity behavior of PPTA/SWNT/H₂SO₄ solution resembles that of SWNT/H₂SO₄ solution. As the SWNT concentration was below 0.1 wt%, the solution viscosity increased accordingly with increasing SWNT wt%. This can be considered as normal behavior of an isotropic solution. When SWNT concentration was exceeded 0.1 wt%, the viscosity decrease at increasing SWNT concentration can be considered as a unique behavior of anisotropic solution. It is caused by the liquid crystal behavior where SWNT are aligned upon shear and form domains of highly ordered structures.

Figure 3. Dependence of complex viscosity ($\eta^*$) of PPTA/SWNT/H₂SO₄ dopes on temperature and SWNT wt% at a shear frequency ($\omega$) of (a) 10 s⁻¹, (b) 1 s⁻¹ and (c) 0.1 s⁻¹.

It has been revealed in rheology and microscopy studies that SWNTs in H₂SO₄ exhibit roughly similar phase behavior as rod-like polymer solutions [21,26]. We have shown in optical microscopy studies that SWNTs in sulfuric acid solution tend to self-assemble into extremely long strandlike structures, which is typically characteristic of the LC phase [21]. The LC transition from a biphasic
phase to a single phase occurs in the SWNT wt% range from 2.7 wt% (maximum viscosity) to 4 wt% (minimum viscosity) [21]. For a blend of SWNT with such a rigid-rod polymer as PPTA, SWNT appears easier to adopt a higher LC order as the transition occurs within the lower SWNT wt% range [27,28]. Thus, 0.1 wt% was the critical concentration of SWNT for PPTA/SWNT/H₂SO₄ system. A biphasic region of mixed isotropic and anisotropic phase existed between 0.1 wt% and 0.2 wt%. Beyond 0.2 wt%, the solution became an increasingly anisotropic-rich phase, and the solution viscosity increased again with increasing SWNT wt%.

2.4. Temperature Dependence of Rheological Behaviors

Temperature effect on complex viscosity (η*) as opposed to shearing frequency (ω) of PPTA/SWNT/H₂SO₄ dope is shown in Figure 4.

Figure 4. Temperature dependence of complex viscosity (η*) of (a) PPTA/H₂SO₄, (b) PPTA/SWNT (98/2 wt%)/H₂SO₄, and (c) PPTA/SWNT (97/3 wt%)/H₂SO₄ as a function of shearing frequency (ω). PPTA is kept at 19.5 wt% in all above cases.

All three samples composed of pure PPTA and PPTA coupled with different portion of SWNT exhibit an obvious shear-thinning behavior through the entire ω range. As ω increases, η* decreases in direct proportion to ω. The slope of such decreasing line on the logarithmic plot remains similar, at about −0.91, for all three samples. This shear-thinning behavior is thought to arise from entanglements...
between macromolecular chains. For rigid-rod polymers such as PPTA, shearing decreases the number of entanglements and thus affords macromolecules to rearrange the configurations readily. By comparison, PPTA/SWNT/H₂SO₄ dopes are more highly shear-sensitive. (1) The η* of PPTA/H₂SO₄ dope decreases as temperature increases from 75 to 85 °C. Such behavior is often observed among conventional polymers. (2) The η* of PPTA/SWNT (98/2 wt%)/H₂SO₄ dope decreases as temperature increases from 80 to 85 °C, while it becomes nearly independent of temperature ranging between 75 °C and 80 °C. (3) The η* of PPTA/SWNT (97/3 wt%)/H₂SO₄ dope decreases slightly as temperature increases from 80 to 85 °C, but increases slightly as temperature increases from 75 to 80 °C.

To determine possible phase changes from 75 to 85 °C, we carried out thermal analysis on PPTA/SWNT/H₂SO₄ dopes. The DSC thermogram (Figure 5) reveals a broad endothermic peak for all samples, implying that multiple transitions occur including monothetic to the liquid state and also isotropic to nematic phase [25,29]. We also noted that with increasing SWNT wt%, the endothermic peak temperature increases from 73.6 to 79.9 °C, possibly because the curve of η* vs. SWNT wt% becomes nearly independent of temperature [30].

**Figure 5.** DSC heating curves of PPTA/SWNT/H₂SO₄ dope with different weight ration of PPTA and SWNT.

![Figure 5](image)

Figure 6 presents the temperature dependent plots of the dynamic modulus of PPTA/SWNT/H₂SO₄ dopes over ω. As SWNT wt% increases, G’ and G’’ arise sharply regardless of temperature. For any temperature and SWNT wt%, elastic characteristic such as G’ of PPTA/SWNT/H₂SO₄ dope is a dominant factor within the frequency range from 0.1 to 10 rad/s. As ω increases, the G’ rises slightly in direct proportion to ω. The slope of G’ versus ω on the logarithmic plot is 0.098, 0.085 and 0.063 as PPTA/SWNT is 100/0, 98/2 and 97/3, respectively. In a low ω region (ω < 1 s⁻¹), the G’ does not rise, while in other regions (ω > 1 s⁻¹), G’’ rises faster than G’, but not in proportion to frequency.
Figure 6. Dynamic modulus ($G'$ and $G''$) of PPTA/SWNT/H$_2$SO$_4$ dope as a function of shearing frequency ($\omega$) at temperatures of (a) 75 °C, (b) 80 °C, and (c) 85 °C.

According to the relative magnitude of $G'$ and $G''$, a logarithmic plot of $G'$ and $G''$ as a function of frequency comprising three zones: terminal, plateau and transition zone, has been proposed [31]. Referred to the literature, PPTA/SWNT/H$_2$SO$_4$ dope is in its plateau zone within the frequency range from 0.1 to 10 rad/s [32]. Rod-like macromolecules like PPTA are thought to exhibit longer relaxation times than flexible polymers due to the difficulty of molecular reorientation arising from the stiff molecular structure and close-packing. The dynamic behavior of Vectra®, for example, supports this argument [33]. As SWNT wt% increases, PPTA/SWNT/H$_2$SO$_4$ dope seems to exhibit more intermolecular interaction and chain rigidity, thus giving rise to the plateau-zone behavior [34].

3. Experimental Section

3.1. Materials and Preparation

Sulfuric acid with 0.2 wt% excess SO$_3$ was prepared by mixing the fuming sulfuric acid (25 wt%) with concentrated sulfuric acid (98 wt%). The concentration of sulfuric acid was measured by acid-base titration. The SWNTs used in this study was acquired from Shenzhen Nanotechnologies Co., Ltd. and an average diameter of <2 nm. Finally, SWNTs were dried under vacuum overnight at 80–100 °C. Dispersions were prepared by mixing with a magnetic stir bar for 3 days in an anhydrous environment in a glovebox which was nitrogen atmosphere at room temperature [35]. The PPTA
polymer was synthesized following the literature methods [36,37]. All glasswares were dried before use.

The PPTA-H$_2$SO$_4$ LC solutions were prepared with four different molecular weight polymers ($M_w = 28169, 56170, 68407, 80135$ g/mol), at different temperatures (75, 80, 85, 90 °C) and with different weight concentrations (18.5, 19, 19.5 wt%). For example, into a 250 mL three-neck flask equipped with a mechanical stirrer and a nitrogen inlet/outlet, were placed 34.01 g of PPTA and 140.4 g of H$_2$SO$_4$ at 75 °C. The mixture was then stirred for 2.5 h and opalescence was observed during this step. Likewise, the LC solutions of PPTA/SWNT/H$_2$SO$_4$, in which PPTA was kept at 19.5 wt% and PPTA/SWNT is 99/1 wt%, were prepared in a 250 mL three-neck flask. SWNTs (0.3401 g) were dispersed in H$_2$SO$_4$ (140.4 g) under an inert and anhydrous atmosphere with mechanical stirring and at room temperature for at least 3 days [21]. Into this flask were then added 34.01 g of PPTA at 80 °C. The mixture was stirred for 2.5 h and stir opalescence was observed during this step.

3.2. Rheological Properties

Subsequently, the PPTA-H$_2$SO$_4$ LC and PPTA/SWNT/H$_2$SO$_4$ dope solutions were performed with dynamical scanning on an ARES rotational rheometer (TA Instruments, New Castle, DE, USA) at four temperatures (75, 80, 85, 90 °C) in the frequency range of 0.01–100 rad·s$^{-1}$ and 0.1–10 rad·s$^{-1}$, respectively. Testing fixtures parallel plates (50 mm) were made of stainless steel 316 to avoid corrosion. In order to prevent water exchange between the sample and environment, the sample surface between the plate and the cone was coated with a mineral oil [38,39]. The effects of dynamic frequency ($\omega$) on rheological behaviors such as $\eta^*$, $G'$, $G''$ and $\tan\delta$ have been examined.

3.3. Characterization

Thermal analysis on the PPTA/SWNT/H$_2$SO$_4$ dope was carried out on a differential scanning calorimeter (DSC) (Mettler Toledo 822) under a N$_2$ atmosphere at a heating rate of 5 °C/min ranging from 25 to 90 °C.

4. Conclusions

We have shown that PPTA-H$_2$SO$_4$ LC solution exhibits a typical shear-thinning behavior. Our rheology studies are summarized as follows. (1) As PPTA wt% increases, rheological curves of $\eta^*$, $G'$, $G''$ and $\tan\delta$ all move up. By contrast; (2) as temperature rises, rheological curves of $\eta^*$, $G'$, $G''$ and $\tan\delta$ all move down. Finally, (3) $\eta^*$, $G'$, $G''$ and $\tan\delta$ all increase as $M_w$ of PTTA increase.

When SWNTs are added into PPTA-H$_2$SO$_4$ LC solution, the protonation of SWNTs in superacids enables their good dispersion to form homogeneous PPTA/SWNT/H$_2$SO$_4$ dope solution. The dope solution exhibits a remarkable nematic LC phase. Our findings are multi-folds. (1) For all temperature and concentration of SWNT, the complex viscosity of PPTA/SWNT/ H$_2$SO$_4$ dope shows an obvious shear-thinning behavior, and the slope of complex viscosity versus frequency on a logarithmic plot is about the identical at ca. –0.91. The DSC thermogram displays multiple phase transitions between monotetic and the liquid state, as well as isotropic and nematic phase. As concentration of SWNT increase, $G'$ and $G''$ rise sharply at various temperatures. (2) For all temperature and concentration of
SWNT, the elastic characteristic such as $G'$ of PPTA/SWNT/H$_2$SO$_4$ dope is a dominant factor within the frequency range from 0.1 to 10 rad/s. At sufficiently high concentrations of SWNT, approximately 0.2 wt%, and at 85 °C, a novel type of single-phase nematic liquid crystal is formed. This allows the dope solution to produce superior-performance fibers by dry-jet wet-spinning process.

Acknowledgements

We are grateful for valuable discussions with Zhusheng Zhou, Qingji Wu, Xingsheng Zhu. This work was supported by Industrial Technology Development Foundation of National Development and Reform Commission of China.

References

1. Friedel, G. Mesomorphic states of matter. *Ann. Phys.* **1922**, *18*, 273–474.
2. Yang H.H. *Aromatic High-Strength Fibers*, 1st ed.; Wiley-VCH: New York, NY, USA, 1989; pp. 207–289.
3. Zhou, M.; Frydman, V.; Frydman, L. On the Molecular organization of PPTA in sulfuric acid: an NMR study. *J. Phys. Chem.* **1996**, *100*, 19280–19288.
4. Kiyo-Oglu, V.N.; Rozhdestvenskaya, T.A.; Serova, L.D. Rheological properties of liquid-crystalline solutions of poly(p-phenylene terephthalamide) and behavior of the jet in spinning through an air space. *Fibre Chem.* **1997**, 29, 81–86.
5. Picken, S.J.; Zwaag, S.V.; Northolt, M.G. Molecular and macroscopic orientational order in aramid solutions: a model to explain the influence of some spinning parameters on the modulus of aramid yarns. *Polymer* **1992**, *33*, 2998–3006.
6. Slugin, I.V.; Sklyarova, G.B.; Kashirin, Al, Tkacheva, L.V.; Komissarov, S.V. Rusar microfilament yarn for ballistic protection. *Fibre Chem.* **2006**, *38*, 22–24.
7. Picken, S.J.; Aerts, J.; Doppert, H.L.; Reuvers, A.J.; Northolt, M.G. Structure and rheology of aramid solutions: transient rheological and rheo-optical measurements. *Macromolecules* **1991**, *24*, 1366–1375.
8. Doppert, H.L.; Picken, S.J. Rheological properties of aramid solutions: transient flow and rheo-optical measurements. *Mol. Cryst. Liq. Cryst.* **1987**, *153*, 109–116.
9. Picken, S.J. Phase transitions and rheology of aramid solutions. *Liq. Cryst.* **1989**, *5*, 1635–1643.
10. Picken, S.J.; Aerts, J.; Visser, R. Northolt, MG. Structure and rheology of aramid solutions: X-ray scattering measurements. *Macromolecules* **1990**, *23*, 3849–3854.
11. Baird, D.G.; Ballman, R.L. Comparison of the rheological properties of concentrated solutions of a rodlike and a flexible chain polyamide. *J. Rheol.* **1979**, *23*, 505–524.
12. Baughman R.H.; Zakhidov A.A.; De Heer W.A. Carbon nanotubes--the route toward applications. *Science* **2002**, *297*, 787–792.
13. Wong, E.W.; Sheehan, P.E.; Lieber, C.M. Nanobeam mechanics: elasticity, strength, and toughness of nanorods and nanotubes. *Science* **1997**, *277*, 1971–1975.
14. Tans, S.J.; Devoret, M.H.; Dai, H.J.; Thess, A.; Smalley, R.E.; Geerligs, L.J.; Dekker, C. Individual single-wall carbon nanotubes as quantum wires. *Nature* **1977**, *386*, 474–477.
15. Treact, M.M.J.; Ebbesen, T.W.; Gibson, J.M. Exceptionally high Young's modulus observed for individual carbon nanotubes. *Nature* 1996, 381, 678–680.

16. Lee, H.K.; Pejanovic, S.; Mondragon, I.; Mijovic, J. Dynamics of single-walled carbon nanotube (SWNT)/polyisoprene (PI) nanocomposites in electric and mechanical fields. *Polymer* 2007, 48, 7345–7355.

17. Kinloch, I.A.; Roberts, S.A.; Windle, A.H. A rheological study of concentrated aqueous nanotube dispersions. *Polymer* 2002, 43, 7483–7491.

18. Andrews R.; Jacques, D.; Rao, A.M.; Rantell, T.; Derbyshire, F. Nanotube composite carbon fibers. *Appl. Phys. Lett.* 1999, 75, 1329–1331.

19. Kumar, S.; Dang, T.D.; Arnold, F.E.; Bhattacharyya, A.R.; Min, B.G.; Zhang, X.; Vaia, R.A.; Park, C.; Adams, W.W.; Hauge, R.H.; Smalley, R.E.; Ramesh, S.; Willis, P.A. Synthesis, structure, and properties of PBO/SWNT composites. *Macromolecules* 2002, 35, 9039–9043.

20. Streekumar, T.V.; Liu, T.; Min, B.G.; Gao, H.; Kumar, S.; Hauge, R.H.; Smalley, R.E. Polyacrylonitrile single-walled carbon nanotube composite fibers. *Adv. Mater.* 2004, 16, 58–62.

21. Davis, V.A.; Ericson, L.M.; Parra-Vasquez, A.N.G.; Fan, H.; Wang, Y.; Prieto, V.; Longoria, J.A.; Ramesh, S.; Saini, R.K.; Kittrell, C.; Billups, W.E.; Adams, W.W.; Hauge, R.H.; Smalley, R.E.; Pasquali M. Phase behavior and rheology of SWNTs in superacids. *Macromolecules* 2004, 37, 154–160.

22. Lin, J.P.; Wu, H.R.; Li, S.J. Kinetics of phase transition in lyotropic liquid crystalline polymers. *Polym. Int.* 1993, 32, 339–342.

23. Gupta, R.K. *Polymer and Composite Rheology*, 2nd ed.; Marcel Dekker: New York, NY, USA, 2000.

24. Schaefgen, J.R.; Foldi, V.S.; Logullo, F.M.; Good, V.H.; Gulrich, L.W.; Killian, F.L. Viscosity-molecular weight relationshios in stiff-chain aromatic polyamides. *Polym. Prepr.* 1976, 17, 69–74.

25. Arpin, M.; Strazielle, C. Conformation de polyamides aromatiques en solution diluée dans l'acide sulfurique concentré. I. Étude par viscosité et diffusion de la lumière. *Makromol. Chem.* 1976, 177, 581–584.

26. Ramesh, S.; Ericson, L.M.; Davis, V.A.; Saini, R.K.; Kittrell, C.; Pasquali, M.; Billups, W.E.; Adams, W.W.; Hauge, R.H.; Smalley, R.E. Dissolution of pristine single walled carbon nanotubes in superacids by direct protonation. *J. Phys. Chem. B* 2004, 108, 8794–8798.

27. Rommel, H.; Forster, G. Topology of the ternary phase system poly(p-phenylene terephthalamide)-sulfuric acid-water. *Macromolecules* 1994, 27, 4570–4576.

28. Picken, S.J. Orientational order in aramid solutions determined by diamagnetic susceptibility and birefringence measurements. *Macromolecules* 1990, 23, 464–470.

29. Li, L.; Li, C.Y.; Ni, C.; Rong, L.; Hsiao, B. Structure and crystallization behavior of Nylon 66/multi-walled carbon nanotube nanocomposites at low carbon nanotube contents. *Polymer* 2007, 48, 3452–3460.

30. Vlasveld, D.P.N.; Fischer, H.R.; Swierenga, E.; Picken, S.J. Interaction of SWCNT and PPTA with sulfuric acid—compatibilization of two materials in a common solvent. *J. Polym. Sci. Part B: Polym. Phys.* 2008, 46, 1914–1922.
31. Ferry, J.D. Viscoelastic Properties of Polymers, 3rd ed.; Wiley-VCH: New York, NY, USA, 1980.
32. Piau, J.M.; Agassant, J.F. Rheology for Polymer Melt Processing, 1st ed.; Elsevier: Amsterdam, The Netherland, 1996.
33. Hsieh, T.; Tiu, C.; Simon, G.P. Rheology and miscibility of thermotropic liquid crystalline polymer blends. J. Non-Newtonian Fluid Mech. 1999, 86, 15–35.
34. Kim, J.Y.; Han, S.I.; Hong, S. Effect of modified carbon nanotube on the properties of aromatic polyester nanocomposites. Polymer 2008, 49, 3335–3345.
35. Chiang, I.W.; Brinson, B.E.; Huang, A.Y.; Willis, P.A.; Bronikowski, M.J.; Margrave, J.L.; Smalley, R.E.; Hauge, R.H. Purification and characterization of single-wall carbon nanotubes (SWNTs) obtained from the gas-phase decomposition of CO (HiPco Process). J. Phys. Chem. B 2001, 105, 8297–8301.
36. Morgan, P.W. Synthesis and properties of aromatic and extended chain polyamides. Macromolecules 1977, 10, 1381–1390.
37. Kwolek, S.L.; Morgan, P.W.; Schaefgen, J.R.; Gulrich, L.W. Synthesis, anisotropic solutions, and fibers of poly(1,4-benzamide). Macromolecules 1977, 10, 1390–1396.
38. Frey, M.W.; Cuculo, J.A.; Khan, S.A. Rheology and gelation of cellulose/ammonia/ammonium thiocyanate solutions. J. Polym. Sci. Part B: Polym. Phys. 1996, 34, 2375–2381.
39. Kim, S.O.; Shin, W.J.; Cho, H.; Kim, B.C.; Chung, I.J. Rheological investigation on the anisotropic phase of cellulose–MMNO/H2O solution system. Polymer 1999, 40, 6443–6450.

© 2010 by the authors; licensee Molecular Diversity Preservation International, Basel, Switzerland. This article is an open-access article distributed under the terms and conditions of the Creative Commons Attribution license (http://creativecommons.org/licenses/by/3.0/).