

Wet-Spinning Multifunctional Carbon Nanotube/Polymer Composite Fibers

José Antonio Benedico¹, Rebeca Marcilla², Eneko Azaceta³, Andrés Seral-Ascaso¹, Enrique García-Bordejé¹, Vicente L. Cebolla¹, Rosa Garriga⁴, Edgar Muñoz¹

¹ Instituto de Carboquímica ICB-CSIC, C/Miguel Luesma Castán 4, 50018 Zaragoza, Spain

² Electrochemical Process Unit, Institute IMDEA Energy, Avda. Ramón de la Sagra 3, 28935 Móstoles, Madrid, Spain

³ IK4-CIDETEC, Parque Tecnológico de San Sebastián, Paseo Miramón 196, 20009 Donostia-San Sebastián, Spain

⁴ Departamento de Química Física, Universidad de Zaragoza, 50009 Zaragoza, Spain
jabenedico@icb.csic.es

Abstract

Remarkable progress in the fabrication of carbon nanotube composite fibers has resulted through the development of the wet-spinning technique pioneered by Vigolo *et al.* [1] This process implies the fabrication of gel fibers as a result of the collapse of surfactant-stabilized carbon nanotube dispersions when injected into a coagulation bath. When dried, those gel fibers become solid fibers with high carbon nanotube contents (≥ 50 wt.%), significantly higher than those achieved by other fiber spinning technologies, such as melt-spinning or electrospinning.

We here report how this wet-spinning method provides carbon nanotube composite fibers with tunable properties, which mainly depend on the composition of the coagulation bath used. Polymers such as polyvinyl alcohol (PVA) [2-5], polyethylenimine (PEI) [6], and a polymeric ionic liquid [7] have been here investigated as coagulants. Remarkable transport-, supercapacitor- and electrochemical actuation properties are demonstrated for these coagulation wet-spun fibers, which offer promise for a variety of applications [2-4].

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References

- [1] B. Vigolo, A. Pénicaud, C. Coulon, C. Sauder, R. Pailler, C. Journet, P. Bernier, P. Poulin, *Science*, **290** (2000) 1331.
- [2] A. B. Dalton, S. Collins, E. Muñoz, J. M. Razal, V. H. Ebron, J. P. Ferraris, J. N. Coleman, B. G. Kim, R. H. Baughman, *Nature*, **423** (2003) 703.
- [3] A. B. Dalton, S. Collins, J. Razal, E. Muñoz, V. H. Ebron, B. G. Kim, J. N. Coleman, J. P. Ferraris, R. H. Baughman, *J. Mater. Chem.*, **14** (2004) 1.
- [4] E. Muñoz, A. B. Dalton, S. Collins, M. Kozlov, J. Razal, J. N. Coleman, B. G. Kim, V. H. Ebron, M. Selvidge, J. P. Ferraris, R. H. Baughman, *Adv. Eng. Mater.*, **6** (2004) 801.
- [5] J.M. Razal, J.N. Coleman, E. Muñoz, B. Lund, Y. Gogotsi, H. Ye, S. Collins, A.B. Dalton, R.H. Baughman, *Adv. Funct. Mater.*, **17** (2007) 2918.
- [6] E. Muñoz, D.-S. Suh, S. Collins, M. Selvidge, A. B. Dalton, B. G. Kim, J. M. Razal, G. Ussery, A. G. Rinzler, M. T. Martínez, R. H. Baughman, *Adv. Mater.*, **17** (2005) 1064.
- [7] R. Marcilla et al., submitted.