In Situ and Ex Situ Preparations of ZnO/Poly-\{trans-[RuCl_2(vpy)_4]/styrene}\) Nanocomposites

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As nanopartículas de semicondutores são de grande interesse científico e tecnológico devido ao fato das suas propriedades eletrônicas e ópticas dependerem do tamanho das partículas. Os nanocristais de semicondutores têm sido intensamente investigados, por exemplo, como nanocargas em diversas matrizes poliméricas. Neste trabalho é apresentada a síntese de nanocompósitos obtidos pela incorporação de nanocristais (NCs) de ZnO no copolímero poli-\{trans-[RuCl_2(vpy)_4]/sty\} (sty = estireno e vpy = 4-vinilpiridina). Numa primeira etapa, o copolímero foi preparado por meio da reação entre os monômeros trans-[RuCl_2(vpy)_4] e estireno. Em seguida, NCs de ZnO passivados organicamente foram misturados com o copolímero poli-\{trans-[RuCl_2(vpy)_4]/sty\}, em CH_2Cl_2, para produzir filmes poliméricos do nanocompósito após evaporação do solvente (método ex situ). As propriedades destes materiais foram comparadas com as de nanocompósitos análogos obtidos a partir da polimerização in situ na presença de NCs de ZnO.

Nanosized semiconductor particles are of great scientific and technological interest because of their size-dependent electronic and optical properties. Semiconductor nanocrystals have been investigated, for example, as nano-fillers for diverse polymer matrices in order to produce new optically active materials. Here we report, for the first time, results concerning the preparation of nanocomposites made of ZnO nanocrystals (NCs) incorporated in the co-polymer poly-\{trans-[RuCl_2(vpy)_4]/sty\} (sty = styrene and vpy = 4-vinylpyridine). In a first step, the co-polymer was prepared by reaction of trans-[RuCl_2(vpy)_4] with styrene. Then, organically capped ZnO NCs and poly-\{trans-[RuCl_2(vpy)_4]/sty\} were mixed in CH_2Cl_2 to produce cast films of the nanocomposites after evaporation of the solvent (ex situ method). The properties of these materials were compared to those of nanocomposites obtained by in situ polymerization in the presence of ZnO NCs.

Keywords: ruthenium(II) complexes, nanocomposites, ZnO nanocrystals

Introduction

Semiconductor nanocrystals (NCs) show size-dependent optical properties and have been investigated as new optical materials in the field of nanotechnology. For example, these nanosized particles have been used together with polymers to produce new optically active inorganic/polymer nanocomposites. This approach takes advantage not only of the intrinsic properties of the starting materials but might also result in new properties due to synergistic effects arising from the combination of the inorganic and organic components. Depending on the nanoparticles’ characteristics and the synthesis-processing of the polymer matrices, there are several synthetic strategies to prepare polymer nanocomposites. Available approaches include sol-gel methods, melt-processing and in situ polymerization. The homogeneous distribution of inorganic nanoparticles within the polymer matrix and the strong interface adhesion between the matrix and nano-fillers are important aspects to be considered because of their influence in the nanocomposites’ performance. In this context, comparative studies concerning the in situ and ex situ preparation of new nanocomposite systems are particularly relevant.

Ruthenium complexes have been extensively investigated both from the fundamental and technological points of view. For example, it is well known that Ru
complexes are of great relevance in the fields of catalysis and solar energy conversion. In addition, Ru complexes have been reported to exhibit properties with strong impact on biological systems. Examples include their role as antioxidants and their antitumoral activity. However, nanoscale research on the use of Ru complexes as precursors for new composite materials has been scarce. In this work, we report for the first time the preparation of nanocomposites containing ZnO nanocrystals (NCs) dispersed in the co-polymer poly-, trans-[RuCl₂(vpy)₄]/sty}, with styrene (sty) using methodologies described previously. In order to assess the synthetic requirements for the preparation of homogeneous ZnO/poly-, trans-[RuCl₂(vpy)₄]/sty} nanocomposites, both ex situ and in situ techniques have been employed and investigated.

**Experimental**

All chemicals were supplied by Aldrich, except ethyl acetate (Lab-Scan), trioctylphosphine oxide (topo, 97% purity) and dichloromethane (Fluka). Styrene (Sigma-Aldrich) was treated over a column of neutral aluminium oxide and stored at 4 °C. The polymerization precursors (the “ruthenium blue solution” and the monomer trans-[RuCl₂(vpy)₄], vpy = 4-vinylpyridine) were obtained as described elsewhere. The co-polymer was prepared by reaction of trans-[RuCl₂(vpy)₄] with styrene (sty) using methodologies previously described. The ZnO NCs were prepared, at room temperature, by dropwise addition of a dimethylsulfoxide (dmso) solution 0.101 mol L⁻¹ in Zn(CH₃COO)₂·2H₂O. The as prepared ZnO NCs were then dispersed in the co-polymer poly-, trans-[RuCl₂(vpy)₄]/sty} under stirring. The ZnO NCs prepared as described above were then dispersed in an equal amount of CH₂Cl₂ under stirring. The ZnO NCs prepared as described previously. In order to assess the synthetic requirements for the preparation of homogeneous ZnO/poly-, trans-[RuCl₂(vpy)₄]/sty} nanocomposites, both ex situ and in situ techniques have been employed and investigated.

**Results and Discussion**

Nanocomposites obtained by the simple mixture of poly-, trans-[RuCl₂(vpy)₄]/sty} and ZnO NCs were initially prepared to investigate the effects of ZnO NCs on the polymeric matrix. Additionally, nanocomposite samples prepared by the simple blend of the components (ex situ method) were also used for comparative purposes with the analogous nanocomposites prepared in situ. Both preparation methodologies, ex situ and in situ, were employed to achieve a nominal load of 1% (m/m) of ZnO/topo in the co-polymer poly-, trans-[RuCl₂(vpy)₄]/sty}, here used as matrix. Figure 1 shows the absorption (UV-Vis) spectra of the starting ZnO/topo NCs and the derived poly-, trans-[RuCl₂(vpy)₄]/sty} nanocomposites. The UV-Vis spectrum of the original ZnO colloid (Figure 1, insert), i.e., before any surface modification step, shows an optical feature peaked at 324 nm that is due to nanosized ZnO. This absorption band is assigned to the first electronic transition of the ZnO exciton and is shifted to higher energies when compared to the typical bulk band gap value for ZnO (Eg = 3.37 eV or 368 nm at room temperature, r.t.), showing that the average diameter of the ZnO nanoparticles is in the quantum regime. In fact, this has been explained as consequence of quantum size effects occurring in nanosized particles with dimensions comparable to the Bohr radius (a₀) of the exciton of the bulk semiconductor. For the case of ZnO, the enlargement of the band gap due to particle size decrease is expected to occur for diameters comparable (or less) than 4 nm.
After chemical surface modification of the ZnO colloid using the coordinating solvent topo, the resulting sample showed a clear red shift (Figure 1a) of the ZnO excitonic peak in relation to the original colloid (Figure 1, insert). This red shift was attributed to a slight increase of the ZnO NCs average size resulting from the coalescence of the smaller nanoparticles during the surface modification step. This optical feature is still observed in the spectrum of the nanocomposite ZnO/poly-(trans-[RuCl2(vpy)4]/sty) prepared in situ (Figure 1c) but it is not noticeable for the nanocomposite synthesized by the ex situ method (Figure 1d). This is probably related to a less dispersion of the ZnO NCs in the polymer for the ex situ preparation, hence leading to a decrease in the homogeneous distribution of the ZnO NCs within the polymeric matrix. As will be discussed later, the presence of ZnO NCs within the polymer was further confirmed through EDX mapping of both nanocomposites (i.e., prepared in situ and ex situ). Finally the most intense band observed at 440 nm (Figure 1) is characteristic of the trans-[RuCl2(vpy)4] complex, as described in a previous work.19

Figure 2 shows the FTIR spectra of the as prepared ZnO NCs and of ZnO NCs after surface modification with the high boiling point solvent topo. As expected, the FTIR spectrum of the original ZnO NCs (Figure 2a) presents the diagnostic bands at 1576 and 1403 cm⁻¹, related to the symmetrical and asymmetrical stretching modes of the carboxylate groups of acetate coordinated to the surface of the ZnO NCs. Note that the ZnO colloid was synthesized using zinc acetate as the ZnII source. Therefore the ZnO NCs surfaces end up terminated with acetate groups, which confer long-term colloidal stability to the ZnO particles in dimethoxysulfoxide (DMSO)/ethanol.17 The IR spectrum of the ZnO powder collected after topo treatment of the original ZnO colloid shows a slight shift (1409 and 1577 cm⁻¹) and a decrease in the intensity of the characteristic carboxylate vibrational bands in relation to the diagnostic band of ZnO (441 cm⁻¹). Moreover, there is the appearance of new bands at 2924-2854 and at 1120 cm⁻¹, that are assigned respectively to the stretching vibrational modes of the CH₂ and P=O groups of the topo molecules. This is an evidence of partial exchange of acetate groups at the ZnO NCs surfaces by topo molecules and thus accounts for the hydrophobic character of the surface-modified ZnO powders.24

Figure 3 shows the FTIR spectra for the nanocomposites prepared by the in situ and ex situ approaches investigated in this work. For the sake of comparison, the FTIR spectrum of the polymeric matrix is also shown. When compared with the spectrum of the polymeric matrix, the FTIR spectra recorded for both ex situ and in situ nanocomposites show a broad band in the 445-475 cm⁻¹ region as a distinctive spectral feature that is due to ZnO NCs dispersed in the polymer.18,25 All other vibrational bands are typical of the poly-(trans-[RuCl2(vpy)4]/sty) used as the matrix.14,19 Although the FTIR spectra shown in Figure 3 can only provide a qualitative analysis of the nanocomposites, it clearly reveals that there were no detrimental effects of the ZnO NCs on the polymer, either using the in situ or ex situ methods for nanocomposite preparation.

The TEM images of the nanocomposite prepared by in situ polymerization show ZnO NCs (dark spots) evenly dispersed in the polymer (Figure 4a); conversely, for the ex situ prepared nanocomposites, the ZnO NCs are not homogeneously dispersed in the matrix (Figure 4c). However, in both cases the TEM images and size analyses

Figure 1. Optical absorption spectra of NCs ZnO/topo (a); in situ ZnO/ topo NCs nanocomposite (b); ex situ ZnO/topo NCs nanocomposite (c) and (d) co-polymer. The insert shows the optical spectrum of the original ZnO colloid.

Figure 2. FTIR spectra of powders collected from the original ZnO colloid (a) and after treatment of the ZnO colloid with topo (b).
show that the ZnO/topo NCs remain morphologically intact after the polymerization process. X-ray powder diffraction confirmed the presence of nanocrystalline ZnO (wurtzite) in the polymeric matrix (see insert in Figure 4c for a typical result). The electron diffraction patterns (insert) obtained for these samples appear as diffuse rings (see insert in Figure 4a for a typical result) due to the nanosized dimensions of the crystalline phase, but are still indicative of ZnO (wurtzite). SEM analysis (Figure 5) confirmed a better dispersion of the ZnO NCs within the poly-[trans-[RuCl2(vpy)4]/sty] for the in situ prepared nanocomposites as compared to the ex situ counterpart. In fact, the sample obtained by blending poly-[trans-[RuCl2(vpy)4]/sty] and ZnO NCs appears as a thin film containing large ZnO agglomerates (Figure 5b), while in Figure 5c the ZnO phase appears as more dispersed particulates.

In light of the above discussion, we used EDX mapping to inquire qualitatively the dispersion degree of the ZnO NCs in the polymer, for both the nanocomposites prepared by ex situ and in situ methods. In the typical SEM images shown in Figure S1 the nanocomposite areas enriched in Zn are labelled in blue while the areas assigned in red are due to the presence of Ru. It is clear that in both nanocomposites the ZnO NCs are well dispersed in the polymeric matrix. However, the in situ prepared sample seems more morphologically homogeneous as revealed by a higher density of ZnO NCs (blue regions) throughout the matrix. This result corroborates the discussion above in suggesting that the in situ method produces nanocomposites with better homogeneity. There is also a homogeneous distribution of the RuII complex in the polymeric matrix under analysis, as assigned by the red regions in the EDX mapping.
In summary, nanocomposites containing ZnO and poly-\{trans-[RuCl\(\text{vpy}\)]\}/styrene\} have been prepared by in situ and ex situ strategies and investigated. The results showed that the in situ polymerization led to more homogeneous nanocomposites as compared to the blend of the components. This study allows further development of this synthetic strategy to obtain homogeneous nanocomposites of variable composition that can find practical interest in ruthenium-based sensor devices.

**Supplementary Information**

Supplementary data are available free of charge at http://jbcs.sbq.org.br, as PDF file.

**Acknowledgments**

K. S. thanks CNPq for financially supporting this project and for a research fellowship. The authors would also like to acknowledge the financial support from the Universidade de Aveiro and FCT (Project PTDC/QUI/67712/06).

**References**

1. Trindade, T.; O’Brien, P.; Pickett, N. L.; Chem. Mater. 2001, 13, 3843.
2. Sanchez, C.; Julian, B.; Belleville, P.; Popall, M.; J. Mater. Chem. 2005, 15, 3559.
3. Caruso, R. A.; Antonietti, M.; Chem. Mater. 2001, 13, 3272.
4. Vaia, R. A.; Ishii, H.; Giannelis, E. P.; Chem. Mater. 1993, 5, 1694.
5. Vaia, R. A.; Vasudevan, J.; Krawie, W.; Scanlan, L. G.; Giannelis, E. P.; Adv. Mater. (Weinheim, Ger.) 1995, 7, 154.
6. Huang, X. Y.; Brittain, W. J.; Macromolecules 2001, 34, 3255.
7. Esteves, A. C. C.; Barros-Timmons, A.; Monteiro, T.; Trindade, T.; J. Nanosci. Nanotechnol. 2005, 5, 766.
8. Zhang, F.; Zhang, H.; Su, Z.; Polyim. Bull. (Heidelberg, Ger.) 2008, 60, 251.
9. Hagfeldt, A.; Graetzel, M.; Acc. Chem. Res. 2000, 33, 269.
10. Garber, S. B.; Kingsbury, J. S.; Gray, B. L.; Hoveyda, A. H.; J. Am. Chem. Soc. 2000, 122, 8168.
11. Paula, M. M. S.; Pich, C. T.; Petronilho, F.; Drei, L. B.; Rudnicki, M.; Oliveira, M. R.; Moreira, J. C. F.; Henriques, J. A. P.; Franco, C. V.; Dal Pizzol, F.; Redox Rep. 2005, 10, 139.
12. Phillips, A. D.; Gonsalvi, L.; Ronnerosa, A.; Vizza, F.; Peruzzini, M.; Coord. Chem. Rev. 2004, 248, 955; Ravera, M.; Bagni, G.; Mascini, M.; Osella, D.; Bioinorg. Chem. Appl. 2007, 91078.
13. Segala, K.; Dutra, R. L.; Oliveira, E. M. N.; Rossi, L. M.; Matos, J. R.; Paula, M. M. S.; Franco, C. V.; J. Braz. Chem. Soc. 2006, 17, 1679.
14. Paula, M. M. S.; Moraes Jr., V. N.; Mocellin, F.; Franco, C. V.; J. Mater. Chem. 1998, 8, 2049.
15. Bandeira, M. C. E.; Prochnow, F. D.; Noda, L. K.; Gonçalves, N. S.; Costa, I.; Melo, H. G.; Crayston, J. A.; Franco, C. V.; J. Solid State Electrochem. 2004, 8, 244.
16. Radovanovic, P. V.; Norberg, N. S.; McNally, K. E.; Gamelin, D. R.; J. Am. Chem. Soc. 2002, 124, 15192.
17. Schwartz, D. A.; Norberg, N. S.; Nguyen, Q. P.; Parker, J. M.; Gamelin, D. R.; J. Am. Chem. Soc. 2003, 125, 13205.
18. Pereira, A. S.; Peres, M.; Soares, M. J.; Alves, E.; Neves, A.; Monteiro, T.; Trindade, T.; Nanotechnology 2006, 17, 834.
19. Franco, C. V.; Paula, M. M. S.; Goulart, G.; De Lima, L. F. C.; Noda, L. K.; Gonçalves, N. S.; Mater. Lett. 2006, 60, 2549.
20. Dantas, N. O.; Cardoso, W. A.; Fanyao, Q.; Monte, A. F. G.; Morais, P. C.; Brito Madurro, A. G.; Brito Madurro, J. M.; Microelectronics J. 2005, 36, 234.
21. Qu, F.; Santos Jr., D. R.; Dantas, N. O.; Monte, A. F. G.; Morais, P. C.; Physica E 2004, 23, 410.
22. Peres, M.; Costa, L. C.; Neves, A.; Soares, M. J.; Monteiro, T.; Esteves, A. C.; Barros-Timmons, A.; Trindade, T.; Kholkin, A.; Alves, E.; Nanotechnology 2005, 16, 1969.
23. Cheng, H-M.; Lin, K-F.; Hsu, H-C.; Lin, C-J.; Lin, L-J.; Hsieh, W-F.; J. Phys. Chem. B 2005, 109, 18385.
24. Kim, C. G.; Sung, K. W.; Chung, T. M.; Jung, D. Y.; Kim, Y.; Chem. Commun. (Cambridge, U. K.) 2003, 16, 2068.
25. Yang, Y.; Li, Y-Q.; Fu, S-Y.; Xiao, H-M.; J. Phys. Chem. C 2008, 112, 10553.

Submitted: April 27, 2009
Published online: July 22, 2010