Preparation of NiO/Ni Nanocomposites Under Microwave Irradiation and Its Catalysis for Reduction of 4-Nitrophenol

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Nickel(II) hydroxide nanocomposites were synthesized by a reaction of nickel(II) acetate tetrahydrate $[\text{Ni}(\text{CH}_3\text{COO})_2 \cdot 4\text{H}_2\text{O}]$ and sodium hydroxide in ethanol under microwave irradiation for 5 min. The NiO/Ni nanocomposites were then prepared by calcining the $\text{Ni}(\text{OH})_2$ nanocomposites in an electric furnace at 700 °C for 2 h. The resulting heated products were examined by X-ray diffraction, scanning electron microscopy and transmission electron microscopy. In addition, the reduction of 4-nitrophenol with sodium borohydride and the NiO/Ni nanocomposites as a catalyst was examined by UV-visible spectrophotometer.

Keywords: NiO/Ni nanocomposites, Microwave irradiation, Reduction of 4-nitrophenol, UV-visible spectrophotometer.

INTRODUCTION

Development of nanotechnology presents many new chemical subjects and provides new opportunities¹. The synthesis of nanocomposites has been studied extensively over the last three decades. Nano-sized composites exhibit novel mechanical, magnetic, electronic and optical properties compared to their bulk counterparts²⁻⁶. Metal-metal oxide nanoparticles, such as $\text{Cu}/\text{Cu}_2\text{O}$ ⁷, Zn/ZnO ⁸ and Sn/SnO_2 ^{9,10}, have potential in magnetic materials, gas sensors and catalysis¹¹. Because of the easy reduction of cationic copper, zinc and tin species, the first preparation of the metallic nanocomposites is normally to reduce the corresponding metal cations with reducing agents under mild conditions, followed by the controlled oxidation of the outer layer with air or oxygen to form the metal/metal oxide nanocomposites^{7,8,11}. The NiO/Ni nanocomposites are of great technical importance because nickel is used as a major component in numerous alloys for high temperature applications¹²⁻¹⁴. NiO, a semiconductor metal oxide with a band gap of 3.6-4.0 eV, has potential applications, such as battery cathode^{15,16}, catalyst¹⁷, gas sensors¹⁸, magnetic materials^{19,20} and electrochromic films²¹. A range of methods have been developed for the synthesis of NiO nanocomposites, such as sol-gel, hydrothermal, precipitation, emulsion, electrodeposition and sputtering²²⁻²⁵. In this study, the $\text{Ni}(\text{OH})_2$ nanocomposites were prepared by nickel(II) acetate tetrahydrate and sodium hydroxide in a liquid phase environment and the resulting $\text{Ni}(\text{OH})_2$ was calcined to prepare the NiO/Ni nano-

composites. The NiO/Ni nanocomposites were characterized by X-ray diffraction (XRD), scanning electron microscopy (SEM) and transmission electron microscopy (TEM). The reduction of 4-nitrophenol with NiO/Ni nanocomposites as a catalyst in the presence of sodium borohydride was examined by UV-visible spectrophotometer.

EXPERIMENTAL

$\text{Ni}(\text{CH}_3\text{COO})_2 \cdot 4\text{H}_2\text{O}$, NaOH and $\text{C}_2\text{H}_5\text{OH}$ were obtained from Samchun Chemicals. 4-Nitrophenol was supplied by Sigma-Aldrich Co., Inc. NaBH_4 was purchased by Kanto Chemical Co., Inc. The structures of nanomaterials were examined by XRD (Bruker, D8 Advance). The surfaces of NiO/Ni nanocomposites were observed by SEM (JEOL Ltd, JSM-6510) at an accelerating voltage of 0.5 to 30 kV. The morphology and crystallite size of samples were examined by TEM (JEOL Ltd, JEM-2010) at an acceleration voltage of 200 kV. UV-visible spectroscopy of samples was performed using a UV-visible spectrometer (Shimadzu UV-1691PC).

Synthesis of the NiO/Ni nanocomposites and catalytic reduction of 4-nitrophenol: The NiO/Ni nanocomposites were prepared by a reaction of 50 mM nickel(II) acetate tetrahydrate $[\text{Ni}(\text{CH}_3\text{COO})_2 \cdot 4\text{H}_2\text{O}]$ and 10 mM sodium hydroxide. The chemicals were dissolved in the 25 mL ethanol as a liquid phase environment at room temperature (the mole ratio of nickel(II) acetate tetrahydrate: sodium hydroxide was 1:2). The mixture was stirred using magnetic stirrer for 4 h at room temperature. After stirring, the mixture in the beaker was placed

in a water bath and irradiated with microwave at room temperature for 5 min (15 s, 20 times). A green precipitate formed as a result. The precipitates were washed 5 times with an ethanol solution (the volume ratio of ethanol: water = 1:1) to remove the impurities. The green powders were dried for 6 h at 80 °C in an electric oven, followed by calcination in an electric furnace at 700 °C for 2 h. In the catalysis experiment, 0.025 mM 4-nitrophenol was added to 15 mL of distilled water. Subsequently, NaBH₄ and the NiO/Ni nanocomposites were added to a 4-nitrophenol solution. The catalytic effect was observed by UV-visible spectroscopy.

RESULTS AND DISCUSSION

The purity and phase of NiO/Ni nanocomposites were analyzed by XRD (Fig. 1). Cubic phase, bunsenite-nanostructured, NiO/Ni nanocomposites with high purity were obtained by calcining the as-prepared sample at 700 °C for 2 h in an electric furnace^{25,26}. Five obvious XRD peaks located at 37.2°, 43.3°, 62.8°, 75.4° and 79.3° 2θ, were assigned to the (101), (012), (110), (113) and (202) peaks, respectively, for cubic NiO (JCPDS44-1159). The three plane indices of Ni, (111), (200) and (220), were also observed at 44.5°, 51.9° and 76.4° 2θ, respectively (JCPDS4-850). Overall, XRD confirmed the synthesis of the NiO/Ni nanocomposites with crystallinity²⁷.

Fig. 1. XRD pattern of the synthesized NiO/Ni nanocomposites; * denotes NiO nanoparticles, Δ denotes Ni nanoparticles

SEM and TEM: Fig. 2 presents a typical SEM image of the NiO/Ni nanocomposites. The image revealed a small rock salt-like morphology²⁵. In addition, the product revealed homogeneous spherical shaped particles²⁸. The formation of these homogeneous particles was examined by microwave irradiation. TEM image showed that the NiO/Ni nanocomposites were spherical and cubic shape with a mean diameter of 20-60 nm (Fig. 3). The particle sizes measured from TEM were in good agreement with those estimated from the XRD patterns which had calculated by Scherrer's equation²⁹.

Fig. 2. SEM image of the synthesized NiO/Ni nanocomposites

Fig. 3. TEM image of the synthesized NiO/Ni nanocomposites

UV-visible spectroscopy: UV-visible spectroscopy is an important method for revealing the optical properties of semiconductor nanocrystals³⁰. In the present study, the UV-visible spectra were used to examine the reduction of 4-nitrophenol to 4-aminophenol. In this reaction, NaBH₄ was used as a reducing agent for 4-nitrophenol to change 4-aminophenol at the catalytic reaction of NiO/Ni nanocomposites. The change of peak from 320 to 400 nm was observed after adding the NaBH₄, this was attributed to the formation of 4-nitrophenolate ion under alkaline condition³¹. Because of conversion from 4-nitrophenol to 4-aminophenol, the color of the solution also turned from yellow to colorless. As the reaction proceeded, the absorbance peak at 400 nm lessened gradually and eventually disappeared. At the same time, a new peak at 300 nm appeared and increased due to the production of 4-aminophenol. Fig. 4 shows the reduction of 4-nitrophenol to 4-aminophenol with NaBH₄ using the NiO/Ni nanocomposites as a catalyst.

Fig. 4. UV-visible spectrum of 4-nitrophenol reduction in the presence of NaBH_4 with heated NiO/Ni nanocomposites

Conclusion

NiO/Ni nanocomposites were synthesized by a reaction with nickel (II) acetate tetrahydrate and sodium hydroxide under microwave irradiation followed by calcination in an electric furnace at 700 °C for 2 h. XRD confirmed that the prepared products were NiO/Ni nanocomposites. SEM and TEM showed that the NiO/Ni nanocomposites had a homogeneous spherical and cubic shape with a mean size of 20–60 nm. In addition, the NiO/Ni nanocomposites, as a catalyst, promoted the reaction of 4-nitrophenol to 4-aminophenol in the presence of sodium borohydride.

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REFERENCES

1. Z.F. Liu, T. Zhu and J. Zhang, *Univ. Chem.*, **16**, 1 (2001).
2. D.Y. Han, H.Y. Yang, C.B. Shen, X. Zhou and F.H. Wang, *Powder Technol.*, **147**, 113 (2004).

3. R.N. Bhargava, *J. Lumin.*, **70**, 85 (1996).
4. E.T. Goldburt, B. Kulkarni, R.N. Bhargava, *J. Taylor and M. Libera, J. Lumin.*, **72**, 190 (1997).
5. R.W. Siegel, *Nanostruct. Mater.*, **3**, 1 (1993).
6. E.F. Hilinski, P.A. Lucas and Y. Wang, *J. Chem. Phys.*, **89**, 3435 (1988).
7. S.U. Son, I.K. Park, J. Park and T. Hyeon, *Chem. Commun.*, **7**, 778 (2004).
8. X.Y. Kong, Y. Ding and Z.L. Wang, *J. Phys. Chem. B*, **108**, 570 (2004).
9. C. Nayral, T. Ould-Ely, A. Maisonnat, B. Chaudret, P. Fau, L. Lescouzères and A. Peyre-Lavigne, *Adv. Mater.*, **11**, 61 (1999).
10. Y. Wang, J.Y. Lee and T.C. Deivaraj, *J. Mater. Chem.*, **14**, 362 (2004).
11. L.G. Teoh, K.D. Li and Y.H. Liu, *Nanosci. Nanotechnol. Lett.*, **3**, 798 (2011).
12. D. Monceau and B. Pieraggi, *Oxid. Met.*, **50**, 477 (1998).
13. A.M. Huntz, *J. Mater. Sci. Lett.*, **18**, 1981 (1999).
14. A.M. Huntz, B. Lefevre and F. Cassino, *Mater. Sci. Eng. A*, **290**, 190 (2000).
15. D. Levin and J.Y. Ying, *Stud. Surf. Sci. Catal.*, **110**, 367 (1997).
16. M. Yoshio, Y. Todorov, K. Yamato, H. Noguchi, J.- Itoh, M. Okada and T. Mouri, *J. Power Sources*, **74**, 46 (1998).
17. M.L. Curri, A. Agostiano, F. Mavelli and M. Della Monica, *Mater. Sci. Eng. C*, **22**, 423 (2002).
18. H.X. Yang, Q.F. Dong, X.H. Hu, X.P. Ai and S.X. Li, *J. Power Sources*, **79**, 256 (1999).
19. Y. Wu, G. Wu and X. Ni, *J. Vac. Sci. Technol.*, **19**, 228 (1999).
20. Y. Wang and J. Ke, *High Technol. Lett.*, **3**, 92 (1997).
21. E.L. Miller and R.E. Rocheleau, *J. Electrochem. Soc.*, **144**, 3072 (1997).
22. Y. Wu, Y. He, T. Wu, T. Chen, W. Weng and H. Wan, *Mater. Lett.*, **61**, 3174 (2007).
23. S. M. Karadeniz, A. E. Ekinci, F. N. Tuzluca and M. Ertugrul, *Asian J. Chem.*, **24**, 1765 (2012).
24. C. Natarajan, H. Matsumoto and G. Nogami, *J. Electrochem. Soc.*, **144**, 121 (1997).
25. R. Mariappan and S. Mahalingam, *Asian J. Chem.*, **25**, 3081 (2013).
26. T. Sreethawong, S. Chavadej, S. Ngamsinlapasathian and S. Yoshikawa, *Colloids Surf. A Physicochem. Eng. Asp.*, **296**, 222 (2007).
27. X.F. Li, A. Dhanabalan and C.L. Wang, *J. Power Sources*, **196**, 9625 (2011).
28. K. Karthik, G.K. Selvan, M. Kanagaraj, S. Arumugam and N.V. Jaya, *J. Alloys Comp.*, **509**, 181 (2011).
29. S. Farhadi and Z. Roostaei-Zaniyani, *Polyhedron*, **30**, 1244 (2011).
30. S.K. Hong, J.H. Lee, J.M. Kim, M.H. Kwon and W.B. Ko, *J. Nanosci. Nanotechnol.*, **11**, 593 (2011).
31. J.H. Lee, S.K. Hong and W.B. Ko, *Asian J. Chem.*, **23**, 2347 (2011).