

Thermal and Electrochemical Properties of Polymannuronate-polyaniline Nanocomposites

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New types of conducting polyaniline-polymannuronate (PANI-PM) composites were synthesized by *in situ* deposition techniques in an aqueous media. By dissolving different weight percentage of polymannuronate (PM) (5, 10, 15, and 25%), the oxidative polymerization of aniline was carried out using ammonium per sulfate as an oxidant. The obtained composites were studied for their thermal stability and electrochemical behavior. The thermal stability of PANI-PM composites is lower than PANI, which supports a strong interaction between PANI and PM. However, the composites show an appreciable electrochemical behavior. Based on these observation the PANI-PM composites can be explored in different fields such as electric devices, sensors, functional coatings, etc.

Key Words: Conducting polymer, Polyaniline, Polymannuronate, Nanocomposites, Thermal stability

Introduction

The development of intrinsically conducting polymers (ICP) has attracted great attention by the scientific community owing to the increased potential for technological applications in electrochromic devices, sensors, electrolytic capacitors, rechargeable batteries, etc.¹ Among the polymers, polyaniline (PANI) generates a special interest owing to its relatively low cost, physico-chemical stability, and a good combination of optical, catalytic, conductive, and sensor properties.^{2,3} However, a major problem still exists in the practical use of PANI and other ICP because of their infusibility, insolubility to an aqueous solution, and poor mechanical characteristics. An alternative approach is the use of PANI in conducting composites or blends with conventional polymers.^{4,5} The obtained composites combine good conductive properties with mechanical and even optical properties of the polymer matrix. This allows the development of polymeric composites with specific properties and shapes.⁶⁻¹⁰ For synthesizing such composite material, various attempts have been made to blend PANI with insulating polymers that have good mechanical strength.¹⁰⁻¹³ Many effective efforts have been directed toward enhancing the electroactivity of PANI^{11,12} either by introducing acidic groups into the PANI chains¹⁴ or doping PANI with negatively charged polyelectrolytes.¹⁵ Recent advances in the development of fabrication technology to the micro or nano scale to obtain an inexpensive signal processing systems have made a variety of novel biomedical sensors possible.^{10,11} Thus it is a very demanding process to synthesize new PANI containing appropriate material suitable for improved electrical and processable properties better than PANI. The introduction of biopolymers especially in the polymer matrix of PANI makes the process of synthesis easier because of the high solubility of biopolymers either partially or completely in aqueous solvents.¹⁶

Polymannuronate (PM) is a good biopolymer for the purpose since it has a well-defined chemical composition; it

has therefore attracted special interest in biomedical research. PM is a component of alginate and has been shown to possess many medical and pharmaceutical applications.^{17,18} The administration of PM has been shown to protect mice from lethal X-ray irradiation and also stimulates the generation of murine myeloid progenitor system.¹⁹ *In vivo* animal models have now revealed the immunologic potential of PM in such diverse areas as the protection against lethal bacterial infections and irradiation, and increasing non-specific immunity.²⁰ In this report, PANI-PM conducting composites have been synthesized with different weight percentages of PM (5, 10, 15, and 25 %). The PANI-PM conducting composites were studied for their thermal behavior by thermogravimetric (TGA), and the electrochemical properties of these composites were studied by cyclic voltammetric measurements.

Experimental

Materials. All the chemicals used for the synthesis of PANI and its composites were purchased from Sigma-Aldrich and used without further purification.

Production of PM from bacteria. The production of PM from the bacterial strain (E1) is derived from *Pseudomonas* sp KL 28, which was previously isolated as n-alkyl phenol degrading bacterium.²¹ The modified King's medium (PG) used for the production of PM consists of (g/L) peptone (20), glycerol (10), and MgSO₄ (1.5). The medium was prepared by phosphate buffer of 50 mM with pH 6.0 sterilized by autoclaving at 15 lb for 20 min. Inoculum was prepared by growing a single colony in 5 mL LB medium for 10 - 12 h at 30 °C. The 50 μL of the inoculum was used to inoculate a 250 mL Erlenmeyer flask containing 50 mL of medium. After then, the flasks were incubated at 30 °C on a rotary shaker at 160 rpm for 3 days. The PM in culture supernatant was precipitated by the addition of an equal volume of isopropyl alcohol and the precipitate was removed by a glass rod. The precipitate was redissolved in water and reprecipitated with

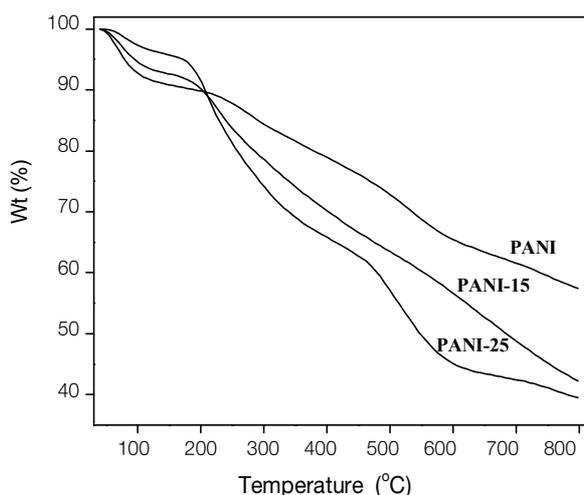


Figure 2. TGA of PANI, PANI-15, and PANI-25 composites.

controlled redox process, which in turn indicates a diffusion-controlled mechanism.

Figure 4 shows the voltammograms of PANI, PANI-15, and PANI-25 at the scan rate 10 mV s^{-1} . The linear dependence of the peak currents on scan rates indicates that the redox process is fast. The electrochemical behavior of PANI-PM composites may originate from the fact that both PANI and PM are conducting, which facilitates a fast charge transfer across the solution. With an increase in the scan rate for PANI-15 and PANI-25, the anodic and cathodic peak potentials shift towards more positive and negative direction, respectively, as shown in Figures 3(a) and 3(b), and this behavior is similar to that of PANI/carbon nanotube multilayer films prepared via the layer-by-layer method.²⁵ Through close comparisons of the CV curves between PANI-15 and PANI-25 and pure PANI in Figure 4, it is observed that not only does PANI-PM display a higher background current in the potential sweep but also there exist faradaic currents, which are believed to arise from the contribution of the loaded PM. It has been reported that PANI was considered as a promising material for electrochemical capacitors because of the existence of different oxidation states. Owing to the higher current in the voltammograms of the PANI-PM electrode than that in a pure PANI,²⁶ a larger capacitance for capacitors equipped with PANI-PM electrode can be anticipated.

The molecular dispersion of PM in solution forms a micro-gel state, which makes the particles a continuous electron path that generates more active sites for the charge transfer through the interface inside the electrode by making good contacts with the PANI matrix. In the cyclic voltammetric curves, the PANI-PM composite core-shell shows appreciable redox activity. The molecular chain of PM is flat and ribbon-like, whose conformation is stabilized by the formation of an intramolecular hydrogen bond in the solution. By the result, the PANI-PM composite macromolecules show very rigid and extended patterns. This makes a continuous electron path by charge transfer at the interface between the electrodes. Thus PANI particles located far from the electrode surface can effectively take part in the redox reaction.^{27,28} The electrochemical activity of these composites is enhanced by a slight

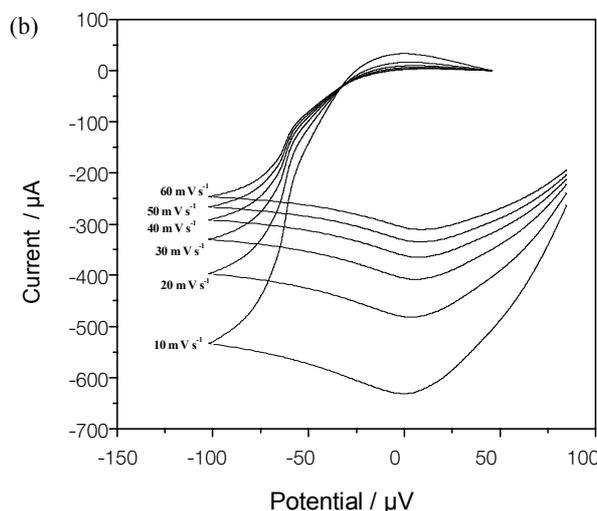
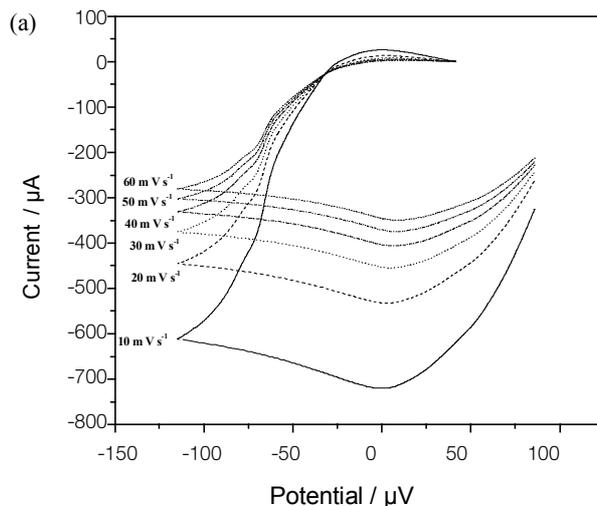


Figure 3. Cyclic voltammograms for (a) PANI-15 and (b) PANI-25 at scan rates of 10 - 60 mV s^{-1} .

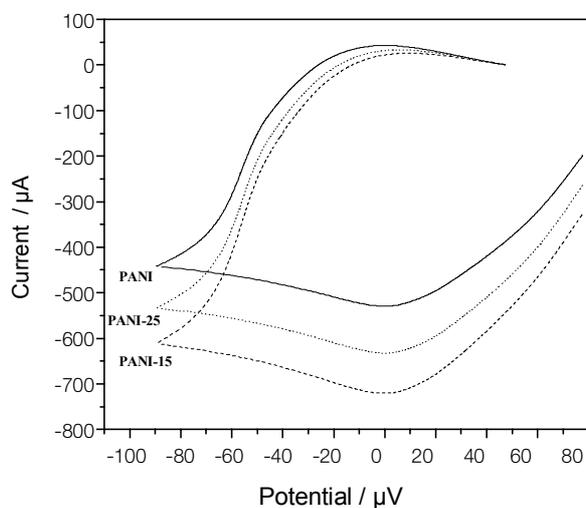


Figure 4. Cyclic voltammograms of PANI, PANI-15 and PANI-25 at the scan rate of 10 mV s^{-1} .

modification in the composite structure of PANI. Continuous studies are in progress to elucidate the mechanism responsible for the increased electroactivity of PANI-PM composites.

Conclusion

The PANI-PM composite was successfully synthesized by the chemical polymerization of aniline containing well-dissolved PM. The aligned PM chains were encapsulated in the cores of the growing PANI polymer chains, resulting in the formation of a PANI-PM hybrid material. The high viscosity and complex nature of PM revealed an improvement in physical and chemical properties of PANI such as conductivity and charge carrier mobility. The electrochemical activity of these composites was enhanced with a slight modification in the composite structure of PANI. The improvements made in the various properties of the present PM are expected to enhance the application potential of the conducting polymer without hampering its chemical properties. The method described here may be useful for developing new applications of these nano-composite films in molecular electronics and other fields. It may be expected that the attractive potentialities offered by this PANI-PM host matrix will be exploited for the development of biosensors based on microorganisms.

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