Fabrication and Properties of a Nanocomposite Multilayer Film Based on a Polyoxometalate

Jihong Liu\textsuperscript{a,b}, Tao Dong\textsuperscript{b}, Shuang Li\textsuperscript{a}, Huiyuan Ma\textsuperscript{a}, and Bo Liu\textsuperscript{a}

\textsuperscript{a} Key Laboratory of Green Chemical Engineering and Technology of College of Heilongjiang Province, College of Chemical and Environmental Engineering, Harbin University of Science and Technology, Harbin 150040, P. R. China

\textsuperscript{b} Chemistry Department, Harbin Normal University, Harbin 150025, P. R. China

Reprint requests to Huiyuan Ma. Tel.: 86-0451-86392716. Fax: 86-0451-86392716. E-mail: mahy017@163.com or Bo Liu. E-mail: liubo200400@vip.sina.com.


Received April 12, 2012

A nanocomposite multilayer film based on a polyoxometalate Na\textsubscript{16}[P\textsubscript{4}W\textsubscript{30}Mn\textsubscript{4}(H\textsubscript{2}O)\textsubscript{2}O\textsubscript{112}]·xH\textsubscript{2}O (Mn\textsubscript{4}P\textsubscript{4}W\textsubscript{30}) was fabricated by the layer-by-layer self-assembly method. It was characterized by UV/Vis spectra and atomic force microscopy (AFM). The electrochromic and electrocatalytic properties were explored using chronoamperometry (CA), UV/Vis spectroscopy and cyclovoltammetry (CV). The attractive feature is a color change of the film by the reduction of the polyoxometalate at different potentials. The film also exhibits good electrocatalytic activity toward the reduction of IO\textsubscript{3}\textsuperscript{−}.

Key words: Electrochromic Behavior, Layer-by-layer Deposition, Polyoxometalate, Electrocatalysis

Introduction

Polyoxometalates (POMs) have gained particular attention for their applications in many fields of science such as medicine, biology, catalysis, and materials due to their remarkable structural and electronic versatility. They show diverse properties such as catalytic activity for chemical transformations, molecule-based conductivity, magnetism, as well as photochromism, electrochromism, and luminescence [1 – 13]. One of the most important electronic properties of these structurally well defined polyoxometalate clusters is that they act as electron reservoirs since the reduction products are mixed-valence species with a characteristically deep-blue color (“heteropoly blues”) [14, 15]. As possible components of electrochromic devices, polyoxometalates (POMs) are promising candidates, also due to their ability to act as an electron reservoir, thereby giving rise to colored mixed-valent species while retaining their structural integrity [1, 16, 17]. In 1978, Tell and co-workers investigated the electrochromic properties of H\textsubscript{3}PW\textsubscript{12}O\textsubscript{40}·29H\textsubscript{2}O and H\textsubscript{3}PMo\textsubscript{12}O\textsubscript{40}·29H\textsubscript{2}O [18, 19], but these electrochromic cells have a disadvantage in that their bleaching is slow when the color is intensified. Recently, Kurth \textit{et al.} fabricated a multilayer film containing the POM cluster [Eu(H\textsubscript{2}O)P\textsubscript{5}W\textsubscript{30}O\textsubscript{110}]\textsuperscript{12−} by a layer-by-layer self-assembly method, which displayed good electrochromism [17]. Gao \textit{et al.} prepared electrochromic multilayer films by the combination of a copper or an iron complex and a monolacunary Dawson-type polyoxometalate of P\textsubscript{2}W\textsubscript{17}. The color changes are adjustable depending on the extent of the reduction of the P\textsubscript{2}W\textsubscript{17} units, by applying more negative potentials [15], indicating that most POMs with a variety of structures could be investigated for practical applications in electrochromic materials.

Taking into account the above, we fabricated a nanocomposite multilayer film containing the polyoxometalate Na\textsubscript{16}[P\textsubscript{4}W\textsubscript{30}Mn\textsubscript{4}(H\textsubscript{2}O)\textsubscript{2}O\textsubscript{112}]·xH\textsubscript{2}O (Mn\textsubscript{4}P\textsubscript{4}W\textsubscript{30}) with electrochromic properties by the layer-by-layer self-assembly method. It was found that the multilayer film displays a color change from yellow to blue due to the reduction of the polyoxometalate. Also, their performance of higher contrast, suitable response time and low operation potential may be promising to meet the requirement for flexible displays and electrochromic devices.

© 2012 Verlag der Zeitschrift für Naturforschung, Tübingen · http://znaturforsch.com
Experimental

Materials

Tris(1,10-phenanthroline)ruthenium(II) chloride Ru(phen)$_2$Cl$_2$· (Ru(phen)$_3$)$_2$, polyethyleneimine (PEI MW 750,000), polystyrene sulfonate (PSS MW 70,000) and (3-aminopropyl)-trimethoxysilane were purchased from Aldrich Chemical Co. and used without further purification. The polyoxometalate Na$_{16}$P$_6$W$_{30}$M$_n$(H$_2$O)$_2$O$_{112}$·$x$H$_2$O (M$_n$P$_3$W$_{30}$) was synthesized according to the literature procedure [20]. The water used in all experiments was deionized to a resistivity of 16 – 18 MΩ cm$^{-1}$. All other reagents were of reagent grade.

Instrumentation

UV/Vis spectra of quartz-supported films were recorded on a U-3010 UV/Vis spectrophotometer made in Japan. All electrochemical experiments were carried out at a CHI 660B instrument at room temperature. A conventional three-electrode system was used, with a bare ITO electrode or a {PSS/PEI/[Mn$_3$P$_4$W$_{30}$/Ru(phen)$_3$]$_n$} multilayer film coated on an ITO electrode as a working electrode, platinum foil as a counter electrode, and Ag/AgCl as a reference electrode.

Preparation of the layer-by-layer film

Quartz substrates and ITO-coated glass were used for the preparation of the film by self-assembly. The substrates were cleaned according to the literature [21], which made their surface hydrophilic, rinsed with deionized water, and dried under a nitrogen stream. A precursor film was deposited on the hydrophilized substrate slide by immersing it into (3-aminopropyl)-trimethoxysilane, PSS and PEI solution for 20 min, respectively, followed by rinsing with deionized water and drying in a gentle nitrogen stream after each immersion. The precursor films were then alternately dipped into 2 × 10$^{-3}$ M Mn$_3$P$_4$W$_{30}$ and 2 × 10$^{-3}$ M Ru(phen)$_3$ for 20 min. Deionized water-rinsing and nitrogen-drying steps were performed after each dipping. The multilayer film {PSS/PEI/[Mn$_3$P$_4$W$_{30}$/Ru(phen)$_3$]$_n$} was formed on the polymer matrix.

Results and Discussion

UV/Vis spectra

UV/Vis spectroscopy has proved to be a useful and facile technique to evaluate the growth process of multilayer films [22–25] and was used in the present work to monitor the assembling process of the {PSS/PEI/[Mn$_3$P$_4$W$_{30}$/Ru(phen)$_3$]$_n$} films. Fig. 1 shows the UV/Vis spectra of the multilayer film {PSS/PEI/[Mn$_3$P$_4$W$_{30}$/Ru(phen)$_3$]$_n$} ($n = 0–10$) deposited on quartz substrates (from bottom to top). Inset: The plots of the absorbance values at 199, 221, 265, 292, and 314 nm of the multilayer film ($n = 1 – 10$).

Atomic force microscopy

The three-dimensional AFM image of the {PSS/PEI/[Mn$_3$P$_4$W$_{30}$/Ru(phen)$_3$]$_3$} film was taken to obtain detailed information about the surface morphology and the homogeneity of this deposited film. As seen in Fig. 2, a mass of uniform particles with a mean grain size of ca. 37 nm was observed on the surface of the film. The film presents a granular texture surface, with a root-mean-square roughness of 3.1 nm calculated over an area of 2.0 × 2.0 μm$^2$.

Electrochromic properties

The UV/Vis spectra of the {PSS/PEI/[Mn$_3$P$_4$W$_{30}$/Ru(phen)$_3$]$_5$} film modified on an
ITO electrode in 0.2 M NaAc + HAc (pH = 3.50) buffer solution were recorded under different potentials from 0 to −0.9 V (Fig. 3). When the applied potentials shifted toward negative, the absorbance at 567 nm was gradually increased, and the {PSS/PEI/[Mn₄P₄W₃0/Ru(phen)₃]₁₃/[Mn₄P₄W₃0]} film was gradually reduced to a different extent of green-blue color, which results from the charge transfer-type (W⁵⁺−O−W⁶⁺ or W⁶⁺−O−W⁵⁺) optical absorption, indicating that the film was electrochromic. The response time of the film was investigated by double-potential experiments with absorbance measurements at 567 nm. The coloration and bleaching times are 5.5 and 7.0 s, respectively, for 90% ΔA (difference between maxima) (see Fig. 4), which is faster than that of a similar [P₂W₁₇][Cu²⁺(Phen)₂]₃₀ film previously reported by Gao (25 and 15 s for the coloration and bleaching times, respectively) [15]. At the same time, the electrochromic reversibility of the films was evaluated by performing repetitive double potential steps from −0.9 to 0 V (Fig. 4). The response time for coloration and bleaching as well as the absorbance of the electrochromic film did not change noticeably even after 200 cycles, which demonstrates a stable electrochromic behavior of the self-assembled films during double potential cycles.
Electrocatalytic activity

Our interest in the multilayer films is also related to its electrocatalytic behavior. Here, we use the \([\text{PSS/PEI}]/[\text{Mn}_4\text{P}_4\text{W}_{30}/\text{Ru(phen)}_3]/\text{Mn}_4\text{P}_4\text{W}_{30}]\) film as the working electrode, and iodate as the test species. Fig. 5 presents CVs of the multilayer film in 0.5 M \(\text{Na}_2\text{SO}_4 + \text{H}_2\text{SO}_4\) (pH = 3.04) buffer solutions containing IO\(_3^−\) at various concentrations. In the range \(-0.85\) to 0.1 V, the cathodic peak currents of the three redox peaks of the \([\text{PSS/PEI}]/[\text{Mn}_4\text{P}_4\text{W}_{30}/\text{Ru(phen)}_3]/\text{Mn}_4\text{P}_4\text{W}_{30}]\) film increased substantially with the addition of iodate, while the relative anodic peak currents decreased, which indicated that iodate was reduced by four-electron, eight-electron, twelve-electron steps of \(\text{Mn}_4\text{P}_4\text{W}_{30}\) [29]. The electrocatalytic efficiency can be calculated as defined by the equation [30]:

\[
\text{CAT} = 100\% \times \left[ \frac{\Delta P(\text{POM, substrate}) - \Delta P(\text{POM})}{\Delta P(\text{POM})} \right],
\]

where \(\Delta P(\text{POM, substrate})\) and \(\Delta P(\text{POM})\) are the peak currents of the POM with and without the presence of substrate IO\(_3^−\), respectively. When 0.6 mM IO\(_3^−\) is employed, the electrocatalytic efficiency of the \([\text{PSS/PEI}]/[\text{Mn}_4\text{P}_4\text{W}_{30}/\text{Ru(phen)}_3]/\text{Mn}_4\text{P}_4\text{W}_{30}]\) film is 259%.

Conclusion

For the first time, the polyanion \(\text{Mn}_4\text{P}_4\text{W}_{30}\) was incorporated into a multilayer film by a layer-by-layer self-assembly method, and the electrochromic properties of the film were investigated. This film exhibited electrochromism with good reversibility and stability. When more negative potentials were applied, the color of the film was gradually changed into deep blue. The coloration and bleaching times are 5.5 and 7.0 s, respectively, for 90% \(\Delta A\). This film also exhibited electrocatalytic activity toward iodate, and thus has the potential for application in electrochromic and electrocatalytic materials.

Acknowledgement

This work was financially supported by the National Science Foundation of China (no. 21071038 and 21101045), the Science and Technology Innovation Foundation of Harbin (no. 2010RFJXG004), the National Science Foundation of Heilongjiang Province (no. 201103), the Foundation of Educational Committee of Heilongjiang (no. 12511082, 12521072), and the Excellent Academic Leader Program of Harbin University of Science and Technology.