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PEDOT:PSS/Fe₂O₃ as hybrid composite film for tuning colour in electrochromism

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Abstract

Electrochromism is known as a modulation of the optical properties under an applied voltage. Among conductive polymers, poly(3,4-ethylenedioxythiophene):poly(styrene sulfonate) (PEDOT:PSS) switching from a neutral to reduced blue state, existing as commercial ink, allows to address displays applications. Aiming at enlarging the color panel, we recently developed an approach based on the addition of oxides leading to PEDOT-PSS+Oxides composites. Particularly, in this work, the influence of the incorporation of Fe₂O₃ on the electrochromic properties of the PEDOT:PSS based films deposited by bar coater are studied. The electrochemical and optical characterizations of the hybrid composite films PEDOT:PSS/Fe₂O₃ by chronoamperometric (CA) coupled with *in situ* UV-visible spectroscopy, indicate that the hybrid composite film with 420 nm thickness shows a reversible switch from blue to red associated with a coloration efficiency of (CE=37 cm²C⁻¹) at 650 nm and transmittance modulation (ΔT=22%) in lithium based electrolytes with switching speeds in the order of the sec. All these results suggest that the hybrid composite film, combining the benefit of mixing inorganic and organic materials, are promising materials for electrochromic devices.

Keywords: electrochromic materials; conductive polymers, PEDOT-PSS, Fe₂O₃, color tuning

Introduction

Electrochromism is generally defined as a reversible change of the optical properties upon applied voltage [1-3]. The combination of various materials both organic and inorganic in different device architectures, has opened up the variety of application areas for electrochromic materials aside to the long standing ones of rear view mirrors and smart windows [4-8]. Electrochromism is known to exist in various types of materials [9-12]. Among these, conducting organic polymers such as poly(3,4-ethylenedioxythiophene):poly(styrenesulfonate) (PEDOT:PSS) have received considerable attention owing to their greater durability, fast switching in redox states, environmental stability, improved processability, and color tunability through easy inks modification or formulation [13, 14]. Composites based on oxides and PEDOT:PSS mixtures can exhibit properties that are superior to those of the pure PEDOT and pure oxide. For instance, Cigdem Dulgerbaki et al [15] reported the synthesis of the hybrid electrochromic material PEDOT/WO₃, in which the tungsten addition leads to significantly enhanced durability under electrochemical cycling and improved coloration efficiency.

Recently, our group [16] has produced a serie of oxides and pigments-substituted PEDOT:PSS that are soluble in common organic solvents and therefore easily processable by doctor blading, serigraphy or bar coating. The electrochromic properties of these electroactive PEDOT:PSS can be adjusted by choosing the appropriate oxides of which nature will in particular play a role on the achieved color. For example we recently reported [17] that the electrochromic properties of PEDOT:PSS based devices were very sensitive to the addition of Fe₂O₃, switching in between a bluish to a reddish state. More precisely, the highest stability and color switching were achieved for small quantity of the Fe₂O₃ (2% Fe₂O₃). Aiming at further improvement, herein the influence of the thickness on the EC properties for single 2% Fe₂O₃ PEDOT:PSS composite films, cycled in three electrodes cell, is discussed.

1. 2. Experiment

2.1. Hybrid composite inks formulation

The hybrid electrochromic inks were formulated from the mixture of the commercial PEDOT:PSS ink Agfa Orgacon EL-P5015 and commercial based Fe_2O_3 pigments (Rouge 110 purchased from Ogres de France, Fe_2O_3 content > 97.1 %). The commercial PEDOT:PSS ink characterized by high-viscosity commercial paste (>100,000 mPa·s) developed for screen printing was first homogenized with a three-roll mill and then diluted with ethanol to lower the viscosity. The various hybrid inks were prepared by mixing PEDOT:PSS, Fe_2O_3 and ethanol stirred for 15 min at room temperature, then dispersed using an ultrasonic bath for 30 min at room temperature. The Fe_2O_3 weight percentage was calculated from the PEDOT:PSS paste mass. Eight identical premixes were prepared: weight ratio of (PEDOT:PSS paste)/(Fe_2O_3) = (100-x)/x, with x = 2%. These preparations were then diluted with ethanol with a mixing ratio of 40 wt % of (PEDOT:PSS + Fe_2O_3) paste and 60 wt % of ethanol. The resulting dilute was dispersed using an ultrasonic bath for 15 min and then stirred again for 5h at room temperature.

2.2. Hybrid composite film deposition and characterization

The films were deposited with a bar coater (K control from RK Print Coat Instruments, Erichsen) onto ITO-coated ($\text{In}_2\text{O}_3:\text{Sn}$) glass substrates (commercialized by SOLEMS with a resistance of 30 Ω/sq) and then dried at 120 °C/5 min on a hot plate. The thickness of the (PEDOT:PSS/ Fe_2O_3) films, measured using a Dektak mechanical profilometer. The surface morphology was characterized by JEOL JSM-840 (operating at 15 kV) microscope.

2.3. Electrochromic Measurements

Electrochemical measurements of hybrid composite (PEDOT:PSS/ Fe_2O_3) films deposited on ITO/glass were carried out in a three electrodes cell configuration using a BioLogic SP50 potentiostat/ galvanostat apparatus. The counter electrode and reference electrode consisted of a platinum foil and saturated calomel electrode, SCE ($E_{\text{SCE}} = 0.234 \text{ V/ENH}$), respectively. The operating voltage was controlled between -1 V and +0.3 V, in lithium-based electrolyte, namely, lithium bis-trifluoromethanesulfonimide (LiTFSI, Solvionic, purity N 99.99%) in 1-ethyl-3 methylimidazoliumbis(trifluoromethanesulfonyl)-imide (EMITFSI, purity N 99.99%). All the electrochemical measurements were performed at room temperature. The *in-situ* optical transmittance of PEDOT:PSS/ Fe_2O_3 hybrid composite films was measured using a Varian Cary 5000 UV-vis-NIR spectrophotometer between 300 and 750 nm. Colorimetry analysis was carried out using a Konica Minolta CM-700D spectrophotometer, allowing the direct determination of colorimetric parameters of the CIE ($L^*a^*b^*$) color space.

3. Results and Discussion

To optimize the electrochromic performance of the hybrid composite film based 98% PEDOT:PSS+ 2% Fe_2O_3 , on which composition has been earlier optimized [13], the thickness of the film was tuned in between 100 nm to over 1 μm using a set of eight bars. **Figure 1** shows the photographs of the various as-deposited films by bar coater, and in the reduced (- 1.0 V) and oxidized (+0.3 V) states as detailed below. An increase in thickness is associated with a significant color change. The pronounced color gradient is supported by the corresponding measured $L^*a^*b^*$ parameters mentioned beside each picture. The addition of 2 wt% of Fe_2O_3 , that remains a low percentage, has a drastic effect on the color of as deposited state evolving from a blue (PEDOT-PSS) to a reddish (PEDOT:PSS+ 2% Fe_2O_3) appearance. From a light reddish color for the 133 nm as-deposited film to a dark-black color for the thickest film (1266 nm) while the composite films adopt a red color in between (420 nm).

When cycled in lithium electrolyte using a three-electrode cells, an optical contrast from blue (-1 V) to reddish-pink (+0.3 V), is visible for thickness below 420 nm while thicker films exhibit limited contrast due to rather dark reddish to black appearance.

Figure 1. Photographs of the hybrid composite films (98% PEDOT:PSS+2% Fe_2O_3) with different thicknesses and corresponding visual appearance of films in the reduced (- 1 V) and oxidized (+0.3 V) states.

In CIE colorimetric space, the color is represented by three parameters, the luminance axis (L^*) and two hue axes (a^*) and (b^*), which can be used to define and compare quantitatively the colors. The evolution of the relative luminance (L^*), the hue (a^*) and (b^*) values of all hybrid composite films with different thicknesses were measured. **Figure 2** shows the evolution of the $L^*a^*b^*$ parameters by adjusting thickness. L^* luminosity parameter gradually decreases versus the thickness from 75 for 133 nm to 25 for the 1450 nm (this thickness is not displayed in Figure 2

as the visible appearance looks alike the one of 1266 nm film) (**Figure 2a**). The hue color, which is directly characterized by the a^* and b^* values shows a saturation maximum around 1200 nm (**Figure 2b and c**). The Chroma of the color C^* was calculated by the following equation:

$$C^* = (a^{*2} + b^{*2})^{1/2}$$

The increase of the thickness from 133 nm to 1450 nm increases the C^* value from 7 to 19.

Figure 2. Evolution of the $L^*a^*b^*$ parameters of 98% PEDOT:PSS+2% Fe_2O_3 films with different thickness.

According to the electrochromic performance of the different films with different thickness, it can be concluded that the most visible optical contrast in between red to blue, instead of light blue to dark blue for pure PEDOT:PSS film, is observed for a thickness of 420 nm. Thus in the following, we focus our attention on the hybrid composite film with 420 nm thickness.

The SEM analysis of the different films before and after Fe₂O₃ addition (**Figure 3**) demonstrates that the films morphology is quite different. Indeed, the PEDOT:PSS film appears homogenous with smooth surface (Figure 3a) while a homogenous dispersion of the Fe₂O₃ particles in the PEDOT:PSS matrix is visible for the PEDOT:PSS/Fe₂O₃ (**Figure 3b**).

Figure 3. SEM images of the surface morphology of (a) PEDOT:PSS film with an average thickness of 550 nm and (b) 98% PEDOT:PSS+2% Fe₂O₃ film with an average thickness of 420 nm.

The electrochromic activity of the hybrid composite films (98% PEDOT:PSS+2% Fe₂O₃) was recorded in a lithium-based ionic liquid. **Figure 4a** shows the chronoamperometric (CA) with voltage steps of (−1 V/30 s) and (0.3 V/30 s) in a three electrodes cell using ionic liquid based LiTFSI in EMITFSI as a supporting electrolyte and Saturated Calomel Electrode as reference electrode (SCE). The cycling of the hybrid composite film shows good reversibility, and fast switching time in between the redox processes. Both response times are in the same order of 1 s in oxidation and 1 s in reduction. The hybrid composite films switch reversibly between a reduced blue state (−1 V) and an oxidized red state (+0.3 V) (**Figure 4b**).

The *in-situ* transmittance upon cycling of 98% PEDOT:PSS+2% Fe₂O₃/LiTFSI in EMITFSI/Pt/SCE is displayed in **Figure 4b**. The reduced-blue state (−1 V) and the oxidized red state at (0.3 V) are associated with transmittance values of about $T_c \approx 8\%$ and $T_b \approx 30\%$ at 650 nm, respectively, corresponding to a total optical transmittance modulation of $\Delta T \approx 22\%$. Such value may appear rather limited in terms of optical contrast but it should be kept in mind that the goal of those systems are the color change and their further use in displays favouring the integration of opaque films.

Nevertheless in regards of the EC community, the evaluation of the coloration efficiency remains a key criteria of merit. The coloration efficiency (CE) is defined as the change in optical density (OD) per unit charge (Q) inserted into or extracted from the EC films and can be calculated from the following formula [18]:

$$CE = \Delta OD / Q$$

where $\Delta OD = \log(T_{red}/T_{ox})$, Q is the electrochemical capacity injected into the electrochromic material per unit area, T_{red} is the transmittance in the blue state, and T_{ox} is the transmittance in the oxidized state, a coloration efficiency 37 cm²C^{−1} is obtained.

Figure 4. Electrochromic response of the 98% PEDOT:PSS+2% Fe_2O_3 with an average thickness of 420 nm. (a) Chronoamperograms cycled in 98% PEDOT:PSS+2% Fe_2O_3 /Li-electrolyte/Pt vs SCE with voltage steps of (-1 V/30 s) and (0.3 V/30 s), and (b) *In-situ* transmittance spectra of the film cycled in 98% PEDOT:PSS+2% Fe_2O_3 /Li-electrolyte/Pt vs SCE, in reduction at -1 V and re-oxidized at 0.3 V.

Conclusion

In this paper, we have successfully elaborated and deposited hybrid composite film 98%PEDOT-PSS+2% Fe_2O_3 using a bar coater with thickness ranging from 133 to 1450 nm. The investigation of the relationship between the PEDOT:PSS/ Fe_2O_3 layer thickness and the electrochromic performances shows that the hybrid composite film with 420 nm exhibits superior reversible electrochromic property with nice color change between reduced blue state and oxidized red state associated with good coloration efficiency and significant transmittance modulation in lithium based electrolytes. The combination of conductive polymers and oxide/pigments is promising to address color issues in displays applications that are of interest to the scientific and industrial communities.

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