Electrosynthesis, Characterization and Electrocatalytic Behaviour of Organic-Metal Thin-Films Based on 2,2’-Bithiophene-5-carboxylic Acid and Metal Ions

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ABSTRACT

2,2’-Bithiophene-5-carboxylic acid (BTA) thin-films on platinum (Pt) electrodes were electrochemically prepared in acetonitrile solution containing 0.1 M tetrabutylammonium perchlorate (TBAP) and 0.05 M BTA. These films were complexes with several metal ions such as Cu2+, Ag+ and Co2+. Their structural characteristics were compared with those of powder complexes chemically prepared from BTA and the corresponding metal ion. IR and XPS techniques reveal that the film complexes with metal ions have the same structures as the corresponding powder complexes. The electrocatalytic activity of BTA film-metal ions has been investigated toward ascorbic acid (AA) oxidation and compared to that obtained on a free BTA film. BTA film-metal ions exhibit good catalytic proprieties and better detection of AA than a free BTA film. This new propriety allows these films to be used as electrochemical sensors. This electronic document is a “live” template. The various components of your paper [title, text, heads, etc.] are already defined on the style sheet, as illustrated by the portions given in this document.

KEYWORDS

2,2’-Bithiophene-5-carboxylic Acid; Metal Ions; Cyclic Voltammetry; IR and XPS; Electrocatalytic Activity; Ascorbic Acid

1. Introduction

Thiophene and its derivatives can be easily polymerized electrochemically or chemically to give conducting polymers with a range of useful properties that make these materials interesting candidates for a variety of potential applications [1,2], such as electro-chromic and electronic devices, electrical energy storage in rechargeable batteries [3] and electrochemical sensors [4], owing to their high chemical stability in air, their ease of preparation and the ability of controlling the doping level. However, conducting polymer systems do not have any intrinsic electrocatalytic activity, and they have been used as host matrices for different chemical species such as metal particles [5], complex ions [6] and enzymes [7], through which these systems acquire catalytic activity.

In recent years, there has been a growing interest in conducting polymer-metal systems [8,9], where the metal is directly attached to the π-conjugated backbone, and has the ability to access a continuum of redox states afforded by metal-polymer cooperativity [9].

The incorporation of metal ions into organic conducting polymers offers potential in the area of organic-inorganic hybrid materials. Polyalkynyl platinum(II) complexes were the first materials prepared in this field [10]. In general, a wide variety of metallic ions can be included by incorporation into conducting polymers. If these ions are catalytically or electrocatalytically active, new composite materials can be obtained, whose proper-
ties will be determined not only by the catalytically active ions, but also by the conducting polymer matrix [11]. This incorporation of dispersed ions into the conducting polymer matrix has two fundamentally different consequences: 1) the polymer structure becomes more porous and spongy, and eventually even at low ion concentrations, facilitates the transport processes accompanying the intrinsic redox process of the polymer; 2) a novel composite material is obtained where the properties are influenced by both components [12].

The oxidation reaction of ascorbic acid (AA) was widely tested on various modified electrodes [13-17]. The role of the AA is known for its properties and its use as an antioxidant agent in foods and drinks, and it is also important for therapeutic purposes and biological metabolism. The determination of AA is a subject of considerable interest and with the use of simple, accurate and advantageous methods. Among of them, the electrochemical determination method is very accurate with high selectivity and best sensitivity. It is recognized as one of the most approaches for the separation of biological compounds such as ascorbic acid and dopamine [18-20]. Therefore, this advantage has a significant attraction in biological and electrochemical research.

With the aim of developing more efficient and novel organic-inorganic hybrid materials to improve their electrocatalytic properties, we were interested here in preparing materials based on 2,2'-bithiophene-5-carboxylic acid (BTA) (Schema 1(a)) and metal ions such as Cu^{2+}, Ag^{+} and Co^{2+} that displayed electrocatalytic properties toward ascorbic acid oxidation (AA). BTA is selected for this study because: 1) it has an electroactive functional group (carboxylate group) able to coordinate to metal ions leading to complex materials; 2) the electro-oxidation of BTA leads to the formation of thin films on the electrodes. The BTA films chemically treated with metal ions leading to complex materials; 2) the electro-oxidation of BTA leads to the formation of thin films on the electrodes. The BTA films chemically treated with metal ions solutions can be used as electrodes for the electrocatalytic oxidation of AA. The complex materials (film or powder form) were characterized by XPS and IR spectroscopy.

2. Experimental

2.1. Materials

All chemical products and solvents (2,2'-bithiophene-5-carboxylic acid (BTA) (Acros organics); copper chloride CuCl2·2H2O (Aldrich), cobalt chloride (CoCl2) and silver nitrate (AgNO3) (Fluka) and acetonitrile (Prolabo), were analytical grade and were used without further purification.

2.2. Synthesis of BTA-Cu^{2+} Complex

The complex was prepared following the procedure described in the literature [21]. BTA (50 mg, 0.39 mmol) was dissolved in methanol (0.8 mL) and NaOH (15.6 mg, 0.39 mmol) was added to this solution. After 30 min stirring, CuCl2·2H2O (26.1 mg, 0.19 mmol) in methanol (0.5 mL) was added dropwise. The reaction mixture was stirred for an additional 30 min. The green solution was filtered, washed with methanol several times and dried at 50°C in vacuum.

2.3. Electrosynthesis of BTA-M Films

Thin films of BTA were first prepared electrochemically (cyclic voltammetry or chronocoumometry) in solution of CH3CN/TBAP 0.1 M, containing 5 × 10^{-2} M of BTA, then the film was rinsed with acetone to remove all traces of monomer and transferred in aqueous metallic solutions (M = Cu, Co, Ag) for concentration of 10^{-2} M each one for 10 min separately.

2.4. Measurements

Electrochemical measurements were carried out using potentiostat/galvanostat type EG & G Princeton Applied Research model 273 under Chem software, in three electrode and a one-compartment cell. In most of our experiments the working electrode was a platinum (Pt) disc 0.03 cm² in area. The counter-electrode was a steel grid and the reference was a conventional saturated aqueous potassium chloride-calomel electrode (SCE). The working electrode was polished before each experiment by cloth polishing with 0.1 μm alumina slurry, and rinsed with acetone and ethanol. The supported electrolyte was 0.1 M tetrabutylammonium perchlorate (TBAP) dissolved in argon-purged acetonitrile.

Infrared spectra of the complex and its precursor in the form of pressed KBr pellets were obtained on a Nicolet Magna-860 Fourier transform infrared spectrometer.

XPS spectra were recorded using a Thermo VG ESCALAB 250 system equipped with a micro-focused, monochromatic Al Kα X-ray source (1486.6 eV) and a magnetic lens which increases the sensitivity of the apparatus. The specimens were pressed against double-sided adhesive tapes mounted on sample holders, and then pumped overnight in the fast-entry lock at 5 × 10^{-8}
mbar before introduction into the analysis chamber. A 650 µm X-ray beam was used at a power of 10 mA × 15 kV. The spectra were acquired in the constant analyser energy mode, with pass energies of 150 and 40 eV for the survey and the narrow regions, respectively. All samples (powders and plates) were subjected to charge compensation achieved with an electron flood gun combined with an argon ion gun. The argon partial pressure was 2 × 10⁻⁸ mbar in the analysis chamber. 

**Figure 1.** Cyclic voltammograms of BTA 5 × 10⁻² M in 0.1 M TBAP/CH₃CN at Pt electrode. Scan rate = 50 mV·s⁻¹. (a) first cycle recorded in 0 - 1600 mV range, (b) repetitive scans between 400 and 1600 mV.

**Figure 2.** Cyclic voltammogram of BTA-Cu²⁺ complex in 0.1 M TBAP/CH₃CN at a Pt electrode at 50 mV·s⁻¹.

3. Results and Discussion

3.1. Electrochemical Behaviour of BTA and a BTA-Cu²⁺ Complex

By applying an anodic polarization to the platinum electrode in a 0.1 M TBAP acetonitrile solution of 5 × 10⁻² M BTA, we obtained cyclic voltammetric curves between 0 and +1600 mV with an oxidation peak at about 1460 mV (Figure 1(a)). This peak corresponds to the oxidation potential of BTA. As can be seen, only the anodic current intensity increases during the successive potential scans (Figure 1(b)), with a slight shift of the oxidation peak towards positive potentials. This behaviour is due to the formation of a thin film having a short conjugation length (BTA dimer). A similar film can be obtained by applying a constant potential at 1350 mV for 240 s. It is also worth noting, in both the above cases, that the films formed on the electrodes are very adherent, stable, and have a coppery colour.

A similar study was undertaken with the BTA-Cu²⁺ complex (chemically synthesized its proposed structure is shown in Schema 1(b)) under the same conditions. The voltammetric behaviour of this complex is characterized, on the first scan, by a shoulder about 1100 mV and a peak at +1600 mV (Figure 2). Compared to the peak of free BTA oxidation (Figure 1), the last peak at 1360 mV is better defined and a negative shift about 100 mV was observed. These results may be attributed to the oxidation of a BTA-Cu²⁺ complex. In the back-sweep, a wave appears at about 160 mV (see figure insert), tentatively assigned to copper reduction in the BTA-Cu²⁺ complex. On the second scan, a large wave at about 350 mV corresponds to copper oxidation and the oxidation peak observed at 1360 mV disappears. Surprisingly, during successive potential cyclic sweeps between −100 and 1600 mV (Figure 2) in a BTA-Cu²⁺ complex solution, no film is formed on the Pt electrode, contrary to the case of free BTA.

3.2. Characterization of Free BTA Film and BTA Film-Cu²⁺ Complex

3.2.1. Electrochemical Characterization

Thin films freshly prepared either by cyclic voltammetry or by the potentiostatic method were cycled in CH₃CN containing 0.1 M TBAP or in H₂O containing 0.1 M LiClO₄ to study their electrochemical behaviour. In both cases, these films present no redox peaks in the range of potential from 0 to 1600 mV (Figure 3(a)), indicating that there is no electroactivity corresponding to the doping-dedoping of these films.

Despite the lack of electroactivity in both the above solutions, the BTA dimer films having carboxylic functional groups are able to complex electroactive species such as Cu²⁺, Co²⁺ and Ag⁺, which allow them to be used as sensors. These films were immersed in aqueous solutions containing either Cu²⁺ or Co²⁺ for 10 min, then

**Figure 3.** (a) Cyclic voltammograms of BTA 5 × 10⁻² M in 0.1 M TBAP/CH₃CN at Pt electrode. Scan rate = 50 mV·s⁻¹. (b) Cyclic voltammograms of BTA-Cu²⁺ complex in 0.1 M TBAP/CH₃CN at Pt electrode at 50 mV·s⁻¹.
rinsed with water and transferred to an electrochemical cell containing either CH₃CN + 0.1 M TBAP or water + 0.1 M LiClO₄.

In the first case, the cyclic voltammogram (CV) displayed oxidation and reduction peaks at 590 and 340 mV (Figure 3(b)), respectively. In addition, the difference between the oxidation and reduction potentials, ΔEₚ = 255 mV, showed that the process is irreversible and the ratio of anodic to cathodic peak currents (Ipc/Ipa = 1) corresponds to a one-electron process. These results can be attributed to the Cu²⁺/Cu⁺ redox couple incorporated (trapped or complexed) [22] into the BTA films.

In the second case, similar observations to those of the first case were observed, but the two redox peaks shifted to more negative potentials, at 100 and −100 mV, respectively (Figure 4). Moreover, when the electroactivity of BTA thin-films after immersion in CuCl₂ solution was tested in an aqueous solution of 0.1 M LiClO₄, a drastic change in the CV features was observed. The related CV shows behaviour similar to that observed for CuCl₂ in aqueous solution, consisting of electrochemically irreversible redox peaks Cu²⁺/Cu⁺ at ca. +100 and −100 mV. It is interesting to note that these peaks shift to more negative potentials compared to that obtained in CH₃CN and have good electrochemical stability under repetitive cyclic voltammetry. These results, in aqueous solution, imply that the BTA film-Cu²⁺ has rather good electrochemical stability, is more electroactive and has high porosity, which is necessary to allow the oxidation-reduction of Cu²⁺ complexing or trapping with BTA dimer films.

A similar study was carried out by immersing BTA thin-films in an aqueous solution of CoCl₂ or AgNO₃. The redox properties of BTA thin-film-Co²⁺ or BTA thin-film-Ag⁺ in aqueous solution exhibit features similar to those observed for BTA thin-film-Cu²⁺ (data not shown), which confirms the presence of metallic ions in BTA thin-films.

3.2.2. XPS Spectra Analysis
To verify and confirm the presence of metal ions in the BTA thin-films, XPS measurements were carried out on the free BTA thin-film, BTA thin-film-Cu²⁺ and BTA-Cu²⁺ complex powder chemically prepared.

The survey spectra of a BTA thin-film and after its immersion in a 0.1 M CuCl₂ solution for 10 min are shown in Figure 5. All the expected elements such as C₁s, O₁s, S₂p in the polymer film can be seen (Figure 5(a)). Additional peaks can be detected in BTA thin-film-Cu²⁺ at 935.2 and 955.0 eV corresponding to Cu 2p₃/₂ and 2p₁/₂ (Figure 5(b)), respectively, which indicates that Cu²⁺ species present in the BTA-Cu²⁺ complex-modified surface. However, it is interesting to note that similar peaks at the same binding energies are observed in the case of a BTA-Cu²⁺ complex powder. These results suggest that Cu²⁺ is complexed by BTA thin-film. In addition, when comparing the binding energy (BE) values of the O₁s in BTA and BTA film-Cu²⁺, we note that the oxygen BE is shifted about 0.8 eV (from 532.5 to 531.7 eV) due to O-Cu bonding [26], resulting probably from the complexation of a -COO− functional group of BTA with Cu²⁺. This observation was confirmed below by IR spectroscopy.

Similar XPS analysis were performed on BTA thin-film-Ag⁺, BTA thin-film-Co²⁺ and on the corresponding complexes under form of powders. These analyses reveal the presence of Ag⁺ and Co²⁺ in the BTA thin-films and powder complexes (data not shown). In our study, the detection of an O-Cu bond by XPS spectrum confirmed the formation of the BTA-Cu²⁺ complex [27].

3.2.3. IR Spectra
IR measurements were performed on the ligand (BTA), BTA film-Cu²⁺ deposited on an ITO electrode and a
BTA-Cu$^{2+}$ powder. By comparing the IR frequencies of the free ligand with those of BTA film-Cu$^{2+}$ and BTA-Cu$^{2+}$ powder can be highlighted the formation of the complex (Figure 6). It is interesting to note that there are differences between the main characteristic absorption peaks of these materials and that of the ligand. The strong band at 1673 cm$^{-1}$ corresponding to $\nu$ (C=O) stretching vibration of COOH in the free ligand (Figure 6(a)) disappears and new very strong and sharp bands in the 1550 - 1510 and 1450 - 1400 cm$^{-1}$ regions corresponding to the asymmetric and symmetric vibrations of the -COO$^-$ group, respectively, appear in the BTA film-Cu$^{2+}$ and BTA-Cu$^{2+}$ powder (Figure 6(b)). These results clearly show COOH functional group deprotonation and coordination to a Cu$^{2+}$ [28-30]. In addition, a broad, weak band at 3358 cm$^{-1}$ observed in BTA film-Cu$^{2+}$ and BTA-Cu$^{2+}$ powder is attributed to $\nu$OH of the methanol or water involved in the BTA-Cu$^{2+}$ structure [21,31,32]. The bands at 813 and 777 cm$^{-1}$ are assigned to C-S bond vibrations.

### 3.3. Electrocatalytic Behaviour of BTA Thin-Film-Metal Ion Complexes toward Ascorbic Acid

Many compounds have been found to be active electron transfer mediators of the electrocatalytic oxidation of ascorbic acid (AA) [13,33,34]. In our case also, the electrocatalytic behaviour of BTA thin-film modified by ion metals has been studied toward ascorbic acid oxidation in a 0.1 M phosphate buffer at pH 7.4.

BTA thin-films were immersed for 10 min in various solutions containing CuCl$_2$, CoCl$_2$ and AgNO$_3$; then the electrode was removed, rinsed with distilled water and
transferred to a freshly prepared ascorbic acid solution. Cyclic voltammetry was used to examine the electroactivity of BTA thin-film-metal ion complexes deposited on Pt electrodes. The cyclic voltammograms of 0.1 M phosphate buffer containing 5 mM of AA clearly show (Figure 7) that BTA thin-film-metal ion complexes present good electrocatalytic activity toward AA compared to that obtained on a free BTA thin-film.

As shown in Figure 7, there is no peak corresponding to the oxidation of AA on a free BTA film (Figure 7(a)). However, irreversible oxidation peaks at ca. 126, 285 and 306 mV were obtained on BTA thin-film-Cu$^{2+}$, BTA thin-film-Ag$^+$ and BTA thin-film-Co$^{2+}$, respectively, corresponding to AA oxidation. These results suggest that the metal ion plays an intermediate to transport the electron transfer between AA and the BTA thin-film elec-
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Figure 7. Comparison of the CV responses for 5 mM AA in phosphate buffer solution (0.1 M PBS, pH 7.4) at various modified electrodes: (a) BTA thin-film; (b) BTA thin-film-Cu$^{2+}$; (c) BTA thin-film-Ag$^+$ and (d) BTA thin-film-Co$^{2+}$; $\nu$ = 100 mV·s$^{-1}$.

4. Conclusion

In this work, we have prepared and characterized BTA films deposited electrochemically on Pt electrodes. We have shown that these films after immersing in metal ion solutions such as Cu$^{2+}$, Ag$^+$ or Co$^{2+}$ had an electrocatalytic process for AA oxidation as analyte. This electrocatalytic activity toward such an analyte is particularly interesting owing to the fundamental importance for several applications, especially in the pharmaceutical and food fields. Moreover, we have shown by spectroscopic techniques (IR, XPS) that BTA can form a complex with Cu$^{2+}$, Ag$^+$ or Co$^{2+}$, and that these metallic ions remain in the film after its use as an electrode in the reaction of AA oxidation. These electrodes may be used as sensors for the detection of AA.

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