

ELECTROOXIDATION OF 2,4,6-TRICHLOROPHENOL ON GLASSY CARBON ELECTRODES MODIFIED WITH COMPOSITE Ni(OH)₂-Co(OH)₂ FILMS

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ABSTRACT

Glassy carbon electrodes coated with a film of electrodeposited cobalt and nickel hydroxides were prepared in order to determine its activity to the 2,4,6-trichlorophenol (TCP) oxidation. The modified electrodes were characterized by cyclic voltammetry (CV), electrochemical impedance spectroscopy (EIS), X-ray photoelectron spectroscopy and Scan Electron Microscopy (SEM) with Energy Dispersive X-ray Spectroscopy (EDS). The results indicate that both hydroxides are homogeneously distributed in the electrode surface. From the viewpoint of electrochemical oxidation of TCP, the presence of cobalt and nickel hydroxides on the electrode surface (i) promotes a more complete oxidation of TCP and (ii) decreases the fouling of the electrode surface in comparison with GC and GC modified with Ni or Co separately.

Keywords: Modified electrodes, (Ni-Co) hydroxides, chlorophenol oxidation

1.- INTRODUCTION

Waste from domestic, agricultural and industrial activities such as cellulose treatment and the manufacturing of plastics, adhesives and petrochemicals may pollute watercourses [1]. The widely used halogenated compounds are not amenable to biological degradation, and can accumulate in living organisms, this being the reason why they are used as wood preservatives, antibacterial agents, fungicides, insecticides and herbicides [2,3].

Au and Pt show high catalytic activity for the electrooxidation of chlorophenols (CP) over short times, but they are quickly fouled up by small oligomers, generated by the coupling of two phenoxy radicals or by reaction of a radical with unreacted CP molecules [4,5]. Electrodes modified with redox mediators could increase the activity and specificity for a given reaction and avoid fouling [6]. So, glassy carbon (GC) electrodes modified with Co(II) phthalocyanine (CoPc) show higher currents and less fouling in the oxidation of 2-chlorophenol (2-CP) and 4-chlorophenol (4-CP) in 1 M NaOH [7]. Also Alatorre et al. [8] found that GC electrodes modified with electropolymerized nickel complexes showed less fouling in the oxidation of 4-CP and 4-nitrophenol in 1 M NaOH. Our laboratory has some experience in this field [4, 5, 9, 10, 11]. The goal is to decrease the oligomerization rate and to promote full oxidation of the phenolic compound.

The redox mediator activity in alkaline media of electrodes modified with Ni(II) compounds is due to the quasi reversible properties of the Ni(III)/Ni(II) process [9,10,11], which may be represented as follows [12]:



Actually nickel hydroxide can have more than one stable structure [13,14,15]. Van der Ven et al. [16] reported the formation of two types of nickel hydroxide, α - and β -Ni(OH)₂, and two kinds of nickel oxyhydroxides, α - and γ -NiOOH, which, due to their different structures and degrees of hydration, have different electrochemical properties.

Other hydroxide, Co(OH)₂, has been widely used as an additive to Ni(OH)₂ in order to improve alkaline secondary batteries [17,18,19], these improvements being attributed to an increase of both ionic and electronic conductivities. Jafarian et al. [20] have reported a good oxidation of methanol at glassy carbon electrodes modified with cobalt hydroxide. Nickel-cobalt hydroxides electrodes can oxidize methane [21] and indirectly chlorophenols [22]. Vidotti et al. [23] reported that ITO electrodes modified with nanostructured hydroxides of nickel and cobalt were very active for the oxidation of urea and Yan et al. [24] found that the overpotential for urea oxidation of a (Ni-Co)-hydroxide electrode was 150 mV lower than that of a Ni hydroxide electrode.

In this work we have prepared and characterized glassy carbon electrodes modified with both nickel and cobalt hydroxides, and have determined their electrocatalytic activity for the oxidation of 2,4,6-trichlorophenol (TCP). As far

as we know, no similar studies have been reported.

2. EXPERIMENTAL SECTION

2.1 Electrode preparation

Two kinds of glassy carbon electrodes were used as supports, a GC disc (CHI Instruments, 0.071 cm² geometric area) and serigraphied electrodes (DropSens, 0.130 cm² geometric area). A one-compartment electrochemical cell with a Pt auxiliary electrode and an Ag/AgCl/KCl_{sat} reference electrode with a Luggin connection were used. The GC disc electrode was cleaned as previously described [11], whereas the GC-serigraphied electrodes were only washed with bidistilled water. Both types of electrodes were stabilized by repetitive cyclic voltammetry (RCV) at 0.1 V s⁻¹ in 0.5 M H₂SO₄ until achieving a constant double layer current density and then activated by RCV (10 cycles) in Britton-Robinson buffer of a given pH. The (Ni-Co)(OH)₂/GC electrode was formed indirectly by reduction of the nitrate anion in a 50 mM (Ni + Co) total concentration of a Ni(NO₃)₂ + Co(NO₃)₂ solution and with an 80/20 or 50/50 mol/mol Ni/Co ratio, using 1.0 M NaNO₃, pH 7.2 as support electrolyte, under nitrogen and at room temperature, as reported previously for Ni(OH)₂ [25]. Deposition was carried out by RCV (6 cycles) at 0.005 V s⁻¹ between -0.4 and 1.2 V. Electrodes of pure nickel hydroxide (Ni(OH)₂/GC) and pure cobalt hydroxide (Co(OH)₂/GC) were obtained by RCV (6 scans) at 0.005 V s⁻¹ between -0.4 and 1.2 V in 0.1 M NaNO₃ with Ni(NO₃)₂ and Co(NO₃)₂, respectively.

2.2 Electrode characterization

The modified GC-DropSens electrode, coated with a gold film, was characterized by Scanning Electron Microscopy (SEM) with an S440 Leica Microscope, using Energy Dispersive X-ray Spectroscopy (EDS) in selected areas. X-Ray Photoelectron Spectroscopy (XPS) was carried out with a channeltron analyzer CLAM2, at pressures from 10⁻⁹ Torr to ultra high vacuum.

Cyclic voltammetry was carried out at 0.005 V s⁻¹ in a Britton-Robinson buffer, pH 9.3. A CHI Instruments CHI660C was used for impedance measurements at constant potential with a 5 mV rms sinusoidal modulation in the 10 kHz–10 MHz frequency range, the frequency being decreased in a logarithmic mode with 10 steps per decade over a period of about 30 min, after which no significant impedance changes had occurred. The impedance data were analyzed with the Autolab, Nova 1.7 software.

3. RESULTS AND DISCUSSION

3.1 Preparation, stabilization and characterization of the Ni(OH)₂-Co(OH)₂ films

3.1.1. Preparation and stabilization of the films.

Although two Ni(NO₃)₂/Co(NO₃)₂ ratios were used, no great differences were observed, and therefore in Figure 1 only the results for an 80/20 Ni/Co

ratio are shown. The hydroxide film, formed by RCV at 0.005 V s⁻¹, produced a current increase over an extended potential range (Figure 1A). The film-coated electrode was washed with abundant distilled water and stabilized by RCV at 0.1 V s⁻¹ in 0.1 M NaOH (Figure 1B). This stabilization, followed by 5 CVs at 0.005 V s⁻¹ in a pH 9.3 Britton–Robinson buffer (Figure 1C), yields a CV with clearly defined peaks (Figure 1D). Three anodic peaks appear: *a1* at 0.42 V, due to the formation of Co(III); *a2* at 0.62 V, due to the formation of Co(IV); and *a3* at 0.73 V, with its corresponding cathodic peak, *c3*, due the Ni(II)/Ni(III) process. This peak *c3* masks the reduction peaks of Co(IV) and Co(III) species. The assignment of the processes is based on the CVs at 0.005 V s⁻¹ in a 9.3 buffer electrolyte of a Ni(OH)₂/GC electrode (Figure 2A) and of a Co(OH)₂/GC electrode (Figure 2B). The characteristic non-Nernstian Ni(III)/Ni(II) couple [25], with peaks at 0.71 and 0.58 V, corresponds to equation (1) above [16].

The CV of Co(OH)₂/GC (Figure 2B) shows two anodic peaks at 0.33 and 0.59 V, with the respective cathodic peaks at 0.31 and 0.56 V, respectively. Zhou et al. [26] attributed these processes to

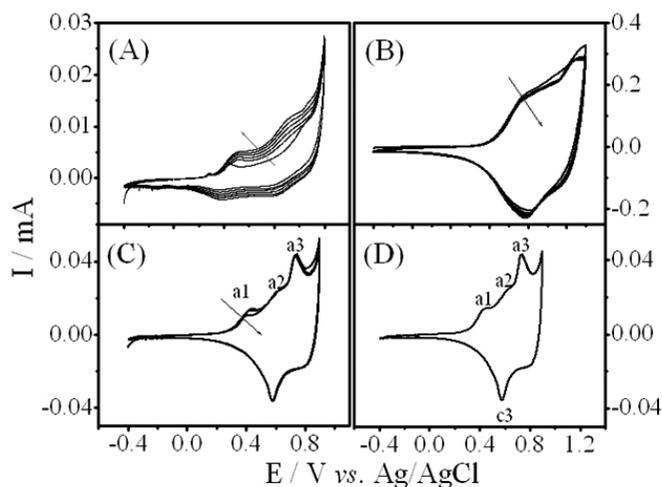
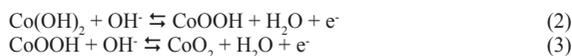


Figure 1: Preparation of the (Ni-Co)(OH)₂/GC electrode. (A) RCVs at 0.005 V s⁻¹ in a 50 mM (Ni + Co) total concentration of a Ni(NO₃)₂ + Co(NO₃)₂ solution and with an 80/20 or 50/50 mol/mol Ni/Co ratio, using 0.1 M NaNO₃, pH 7.2 as support electrolyte. (B) Stabilization of the (Ni-Co)(OH)₂ film by RCV at 0.1 V s⁻¹ in 0.1 M NaOH. (C) Stabilization of the (Ni-Co)(OH)₂ film in a pH 9.3 Britton-Robinson buffer by RCV at 0.005 V s⁻¹. (D) Stabilized CV under the conditions in (C).

It becomes apparent that the CV of a mixture of Co and Ni hydroxides is simply the sum of the individual CVs of the two pure hydroxides.

3.1.2. Electrochemical Impedance Spectroscopy of the films.

Nyquist and Bode plots of electrodes of (Ni-Co)(OH)₂/GC (Figure 3 A and D), Co(OH)₂/GC (Figure 3 B and E), Ni(OH)₂/GC (Figure 3 C and F) were obtained at three potentials, 0.45 V (squares), 0.63 V (circles) and 0.83 V (triangles). The Nyquist diagrams (Figure 3A→C) show that at 0.45 V all the electrodes behave as a constant phase element, this behaviour obtaining also at 0.63 V for (Ni-Co)(OH)₂/GC (Figure 3A) and for Co(OH)₂/GC (Figure 3B) electrodes, whereas in Ni(OH)₂/GC it is clear that the Ni(II)/Ni(III) process has already started at this potential (Figure 3C). At 0.83 V charge transfer occurs in all the electrodes, the lower charge-transfer resistance (R_p) corresponding to the Co/GC electrode. It is very interesting that the addition of Co to the Ni electrode reduces by a factor of ten its R_p, which so becomes similar to that of Co(OH)₂/GC. The Bode plots (Figure 3 D→F) show that at 0.45 V both the Co(OH)₂/GC and the (Ni-Co)(OH)₂/GC electrodes have a time constant at high frequencies, which disappears at 0.63 V, when the Co(III)/Co(IV) process is occurring.

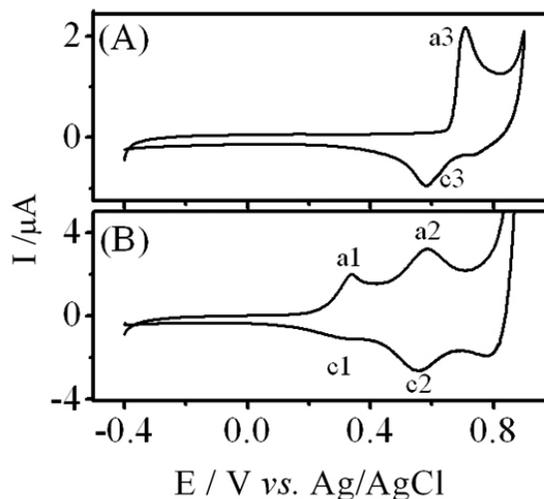


Figure 2. CV at 0.005 V s⁻¹ in a pH 9.3 Britton-Robinson buffer of: (A) Ni(OH)₂/GC electrode. (B) Co(OH)₂/GC electrode.

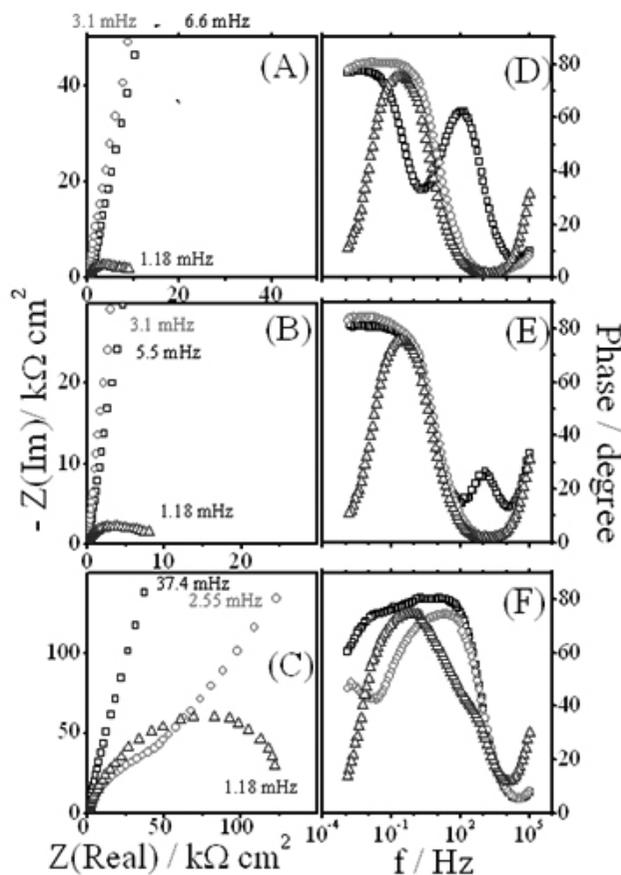


Figure 3. Nyquist plots (A, B and C) and corresponding Bode plots (D, E and F) at 0.45 V (squares), 0.63 V (circles) and 0.83 V (triangles) in a pH 9.3 buffer electrolyte of GC electrodes coated with a film of: (A, D) (Ni-Co)(OH)₂; (B, E) Co(OH)₂; (C, F) Ni(OH)₂.

3.1.3. SEM of the films.

All the samples were coated with gold in order to increase their conductivity. SEM images of serigraphed GC electrodes (Figure 4 A) show that their surface is made up of flakes, and therefore very rough. The hydroxide films appear as sand spread on these flakes (Figure 4 B→D). EDS analysis of different parts of the deposits showed the presence of Co and Ni (Figure 5). The atomic compositions of the surfaces of the different electrodes are given in Table 1.

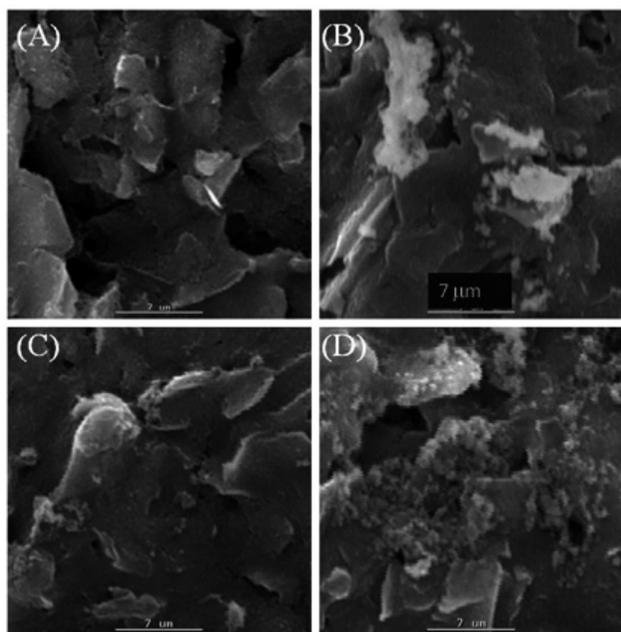


Figure 4. SEM images of: As-received GC serigraphed electrode (A). As-prepared $\text{Co(OH)}_2/\text{GC}$ (B), $\text{Ni(OH)}_2/\text{GC}$ (C), and $(\text{Ni-Co})(\text{OH})_2/\text{GC}$ electrodes (D).

Table 1 Atomic composition (atom %) of the surfaces of bare GC and of GC modified with Ni(OH)_2 , Co(OH)_2 and $(\text{Ni-Co})(\text{OH})_2$ films, as obtained by EDS.

Atomic element	Electrode			
	Bare GC	$\text{Ni(OH)}_2/\text{GC}$	$\text{Co(OH)}_2/\text{GC}$	$(\text{Ni-Co})(\text{OH})_2/\text{GC}$
C	80	73	1.9	49
Cl	20	10	13	8.8
O	-	6.3	16	7.7
Ni	-	11	-	6.1
Co	-	-	69	28

The Co/Ni ratio in the $(\text{Ni-Co})(\text{OH})_2$ films is 4.6. This cannot be due to solubility differences, since the pK_{sp} constant of Co(OH)_2 (14.8) is only slightly lower than that of Ni(OH)_2 (15.3), and consequently the solubility of the former is only 50% higher than that of the latter [27]. Therefore, the high Co/Ni ratio should be due to a higher deposition rate of Co(OH)_2 .

3.1.4. XPS of the films.

The wide-scan XPS spectra of both bare and $(\text{Ni-Co})(\text{OH})_2$ -coated GC electrodes are given in Figure 6A. The spectrum of bare GC shows a strong carbon signal (85%) and a moderate Cl contribution (12%), which, according to the binding energy of the main Cl 2p_{3/2} core level, corresponds to chloride ions. A tiny oxygen contribution (3%) is also observed. The spectrum of the modified electrode shows additional Co and Ni photoemission and Auger lines. The surface concentrations of the different elements estimated from this spectrum are: C, 63%; O, 33%; Co, 2.5%; Ni, 0.5%; and Cl, 1%. The sparse coverage of hydroxides agrees with the SEM micrographs of Figure 4. The Co/Ni ratio is within 10% of that evaluated by SEM-EDS.

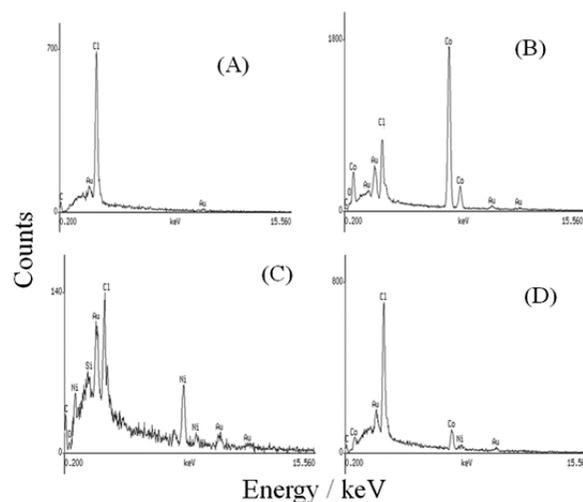


Figure 5. EDS analysis of the samples in Figure 4.

The high-resolution spectra of the Co 2p, Ni 2p and O 1s core levels are given in Figure 6B-D. The Co 2p spectrum (Figure 6B) is rather complex, showing two spin-orbit doublets and several shake-up satellites (the dashed and dotted lines are the contributions of Co^{3+} and Co^{2+} , respectively). The spin-orbit doublet characterized by binding energies of the Co 2p_{3/2} and Co 2p_{1/2} core levels of 779.9 eV and 795.0 eV, together with the small shake-up satellite peak at 790.0 eV, is characteristic of Co^{3+} [28,29]. The broader spin-orbit doublet with binding energies of 781.5 eV (Co 2p_{3/2}) and 796.7 eV (Co 2p_{1/2}) and the two strong shake-up satellites at 785.4 eV and 802.3 eV are assigned to Co^{2+} [28,29]. The Co^{2+} and Co^{3+} relative concentrations obtained from the fitting of this spectrum are 62% and 38%, respectively.

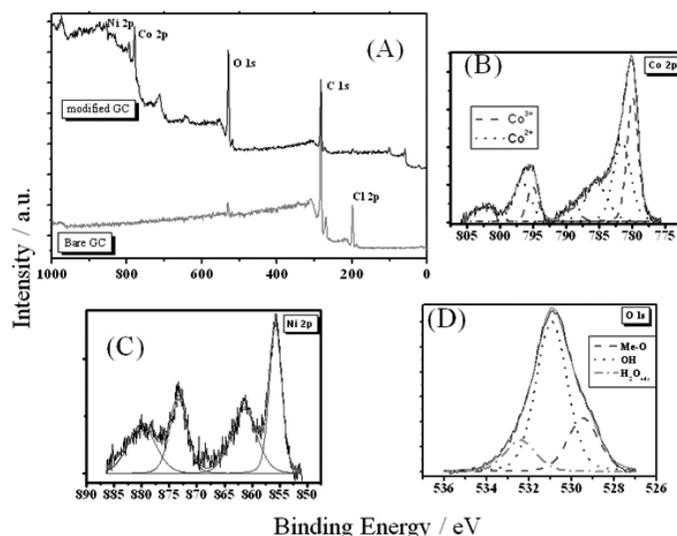


Figure 6. (A) Ex situ XPS spectra of an untreated GC serigraphed electrode and of the former coated with an as-prepared $(\text{Ni-Co})(\text{OH})_2$ film. Expanded regions of the XPS spectrum of the $(\text{Ni-Co})(\text{OH})_2/\text{GC}$ electrode: (B) Expanded Co2p region, (C) Expanded Ni2p region, and (D) Expanded O1s region.

The Ni 2p spectrum (Figure 6C) shows only one spin-orbit doublet with binding energies of 855.8 eV (Ni 2p_{3/2}) and 873.2 eV (Ni 2p_{1/2}) and two intense shake-up satellites at approximately 6 eV above these main photoemission lines. All these spectral features are compatible with the presence of Ni(OH)_2 [30, 31]. In this respect, it is interesting to note that the O 1s spectrum (Figure 6D) of the modified electrode shows three contributions.

The most intense one (64%) corresponding to metal-OH bonds (530.9 eV), which supports the assignment to Ni(OH)₂ (and, most likely, supports also the presence of Co(OH)₂). The binding energy at 529.5 eV (23%) is characteristic of a metal=O double bond [28,29] and, if we take into account the above results, it could be due to CoOOH. The third contribution at 532.4 eV (13%) is typically associated with adsorbed water [28,29].

Finally, both SEM-EDS (mass analysis) with XPS (surface analysis) yielded nearly the same Co/Ni ratio, supporting a homogeneous distribution of the two hydroxides in the film.

3.2. Electrooxidation of 2,4,6-trichlorophenol (TCP)

3.2.1. Cyclic voltammetry.

We had already reported [10] that the active species in the oxidation of TCP at glassy carbon electrodes is the phenolate anion, and therefore pHs higher than the pKa of 2,4,6-TCP, 7.7 [27], should be used. Previous results indicate that in cyclic voltammetry experiments there is diffusional control in the first electron transfer but that phenolic oligomers formed afterwards foul the electrode. Therefore, the goal is to favor the second charge transfer, so avoiding the formation of oligomers. Using Britton-Robinson buffers of different pH values, it was found that the optimum pH for the oxidation of 1 mM TCP on Ni-Co/GC electrodes was 9, because at this pH TCP reacts at the potentials of the Co(III)/Co(IV) and of Ni(II)/Ni(III) reactions (Figure 7). In the negative scan two new cathodic peaks are observed, *c4* and *c5*, due to the reduction of two different quinones to the respective hydroquinones, as confirmed using commercial 2,6-dichloro-1,4-hydroquinone and 2-chloro-1,6-hydroquinone.

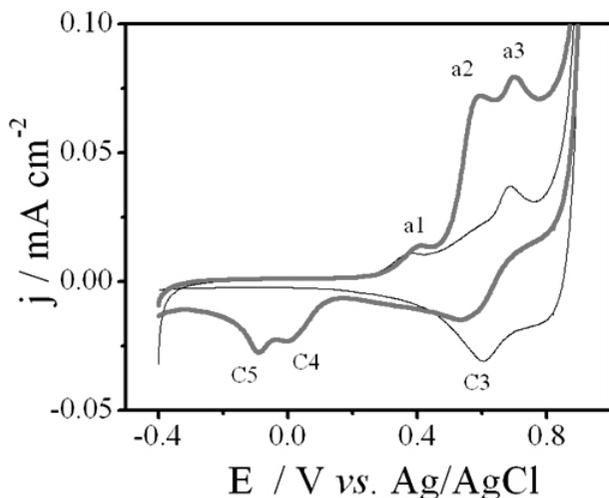


Figure 7. Cyclic voltammograms at 0.005 V s⁻¹ of a (Ni-Co)(OH)₂/GC electrode in a pH 9.3 Britton-Robinson buffer (thin solid line) and in the presence of 1 mM TCP (thick solid line).

The CVs at 0.005 V s⁻¹ of a (Ni-Co)(OH)₂/GC electrode in pH 9.3 buffer electrolyte, before (thin solid line) and after (thick solid line) 5 CVs in the presence of 1 mM TCP, are shown in Figure 8. Although some reaction products with a quinone/hydroquinone structure remain adsorbed on the electrode surface (peaks *c5* and *a5*), the activity of the Co and Ni processes remain practically unaltered, indicating that the electrode had not become coated with phenolic oligomers, or that, if present, these are very porous.

The onset potentials at 0.005 V s⁻¹ in 1 mM TCP in a pH 9 buffer electrolyte are similar for Co(OH)₂/GC (0.44 V) and Ni(OH)₂/GC (0.45 V) electrodes (not shown). The onset potential at the (Ni-Co)(OH)₂ electrode is the same, 0.46 V, but then the current increases much more rapidly.

3.2.2. Electrochemical Impedance Spectroscopy.

The Nyquist (A→C) and Bode (D→F) plots at 0.451 V in pH 9.3 buffer (filled symbols) and in the presence of 1 mM TCP (open symbols) are shown in Figure 9 for the three electrodes. In all cases the charge-transfer resistance decreases in the presence of TCP, which evidences its oxidation. The lowest charge-transfer resistance was that of the Co/GC electrode (Figure 9B). The R_c of the (Ni-Co)(OH)₂/GC electrode (Figure 9A) is lower than that of the Ni(OH)₂/GC electrode (Figure 9C), which agrees with the hypothesis that the presence of Co favours the conductivity across the Ni(OH)₂ films, thereby improving the performance. Please note that at the higher frequencies the Bode

plot of the Co(OH)₂/GC electrode changes dramatically in the presence of TCP, whereas those of (Ni-Co)(OH)₂/GC and Ni(OH)₂/GC electrodes do not. The system is so complex that a reasonable equivalent circuit cannot be obtained.

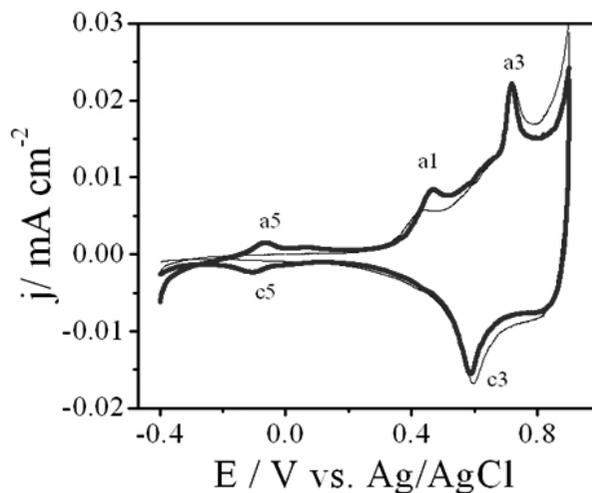


Figure 8. Cyclic voltammetry at 0.005 V s⁻¹ of a (Ni-Co)(OH)₂/GC electrode in a pH 9.3 Britton-Robinson buffer before (thin solid line) and after (thick solid line) 5 RCVs in the presence of 1 mM TCP.

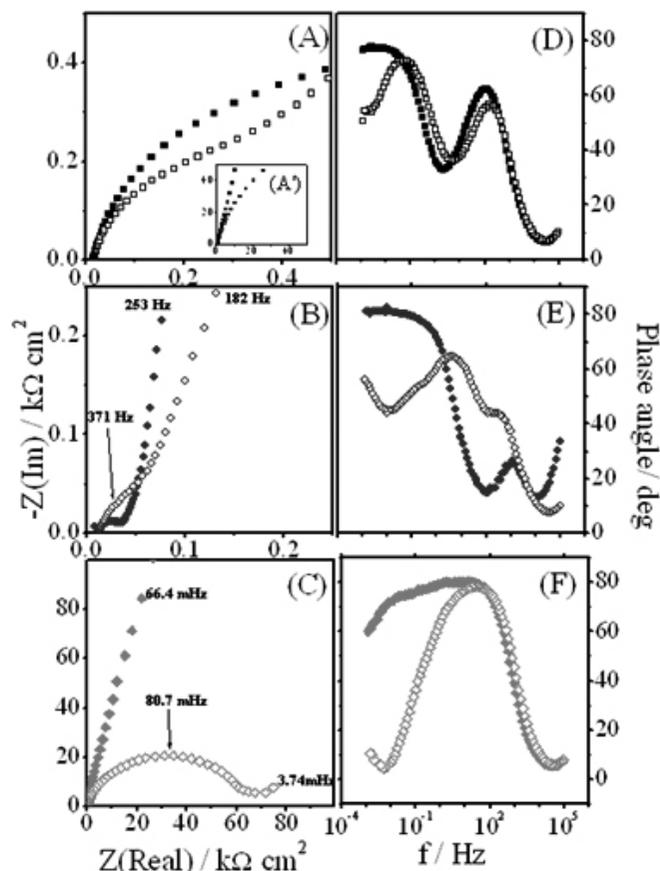


Figure 9. Nyquist (A→C) and Bode (D→F) plots at 0.451 V in a pH 9.3 buffer (closed symbols) and in the presence of 1 mM TCP (open symbols) for the following electrodes: (A) and (D), (Ni-Co)(OH)₂/GC; (B) and (E), Co(OH)₂/GC; (C) and (F), Ni(OH)₂/GC.

4.- CONCLUSION

4.1 Surface characterization

Both XPS and SEM show that in the mixed film Ni(OH)₂-like and Co(OH)₂-like compounds coexist. Independently of the initial Ni²⁺/Co²⁺ ratio in the electrolyte, the Co/Ni ratio in the film is about 4.6, indicating that the Co(OH)₂ precipitation is much faster. It was found by CV that in (Ni-Co)(OH)₂ films each hydroxide behaves as in its pure state.

4.2. Oxidation of 2,4,6-TCP

The mixed (Ni-Co)(OH)₂ film promotes a more complete oxidation of TCP, facilitating the formation of the corresponding quinones, which substantially decreases the fouling of the electrode surface.

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