Electrochemical Sensor Based on the Clay-Carbone Paste Electrode Modified by Bacteria-Polymer for Elimination of Phenol

R Maallah*, MA Smaini, C Laghlimi, J El Mastour and A Chtaini
Molecular Electrochemistry and Inorganic Materials Team, Sultan Moulay Slimane University, Morocco

*Corresponding author: R Maallah, Molecular Electrochemistry and Inorganic Materials Team, Sultan Moulay Slimane University, Morocco.

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Abstract
A Cyclic voltammetry (VC), Square Wave Voltammetry and Impedance Spectroscopy methods for the determination of trace amounts of phenol at bacteria-polymer-Clay-CPE bioelectrode is proposed. The results showed that the bacteria-polymer-Clay-CPE exhibited excellent electro catalytic activity to phenol. This electrochemical biosensor shows an excellent performance for detecting phenol.

Keywords: Modified electrodes; Cyclic voltammetry; Clay; impedance spectroscopy; Bacteria; Phenol

Introduction
Phenol and related compounds are widely used in industry for the manufacture of a wide variety of aromatic compounds, including rubber, fertilizers, paints, drug preparations, petroleum and agricultural industries [1,2]. Phenol is considered carcinogenic and exposure to phenol causes several symptoms such as convulsions, dizziness and irregular breathing [1,3]. Phenol is one of 129 compound chemicals considered important pollutants listed by the Environmental Protection Agency (EPA) [4,5]. Due to severe legislation, prohibit the release of toxic products into the environment. Current research has focused on the development of a simple, cost-effective, fast and reliable technology that can be used for the immediate detection of heavy metals and toxic products in the environment. In this context, a technology seems to fully correspond to the requirements necessary for sustainable development and the preservation of the ecosystem: "bio electrochemistry". This discipline can be defined as a science involving the principles of electrochemistry associated with the field of living. Here we will focus on the electrochemistry / bacteria coupling. Indeed, the conversion of the catalytic activity of a bacterium into an electric current has paved the way for the development of various and varied bioelectrodes. Their fields of application are currently numerous and particularly in fields such as energy (biopiles) and analytical applications (bacterial biosensors) [6,7].

With regard to electro-analytical techniques, procedures involving the oxidation of phenol on solid electrodes [8,9] have been reported. In addition, chemically modified carbon paste electrodes have proved very useful for analytical applications [10]. In previous work [11,12], the electrochemical oxidation of phenol has been studied. We present a simple and sensitive method of determining these compounds based on their reaction. The purpose of the work presented here was to study the electrochemical properties of phenol on the bacterial-polymer-clay modified carbon paste electrode as well as the electrochemical characterization of electrodes by the cyclic voltammetric technique, Square Wave Voltammetry and Impedance Spectroscopy.

Experimental
Apparatus and Software
Voltammetric experiments were performed using a voltalab potentiostat (model PGSTAT 100, Eco Chemie B.V., Utrecht, The Netherlands) driven by the general-purpose electrochemical systems data processing software (voltalab master 4 software) run under windows 2007. The three-electrode system consisted of a chemically modified carbon paste electrode as the working electrode a saturated calomel electrode (SCE) serving as reference electrode, and platinum as an auxiliary electrode.
Electrodes

Modified electrodes were prepared by mixing a carbon powder and the desired weight of clay. The body of the working electrode for voltammetric experiments was a PTFE cylinder that was tightly packed with carbon paste. The geometric area of this electrode was 0.1256cm². Electrical contact was made at the back by means of a bare carbon.

Procedure for Immobilization of Bacteria in a Polymer Film

The modified electrodes were immersed in a cell 100 ml of the 0.1 M electrolyte of NaCl contains 1 ml of ε-caprolactone monomer; the electro polymerization is carried out by cyclic voltammetry. Then the modified electrodes were immersed in a cell 20 ml of distilled water, 0.1M NaCl and the bacterial suspensions and then characterized as a function of time by linear cyclic voltammetry and by electrochemical impedance spectroscopy (Figure 1).

Results and Discussion

Surface characteristics
The morphology of the electrode surface of Clay was observed by scanning electron microscopy (Figure 2). We find that the matrix is formed by compact particles fractions between 1 and 15 μm. Clay treaty has the following chemical composition given by transmission electron microscopy (TEM): O (22%), Mg (5.4%), Al (22.4%), K (2.7%), Ca (1%), Ti (1.8%) Fe (17.1%), Si (27.8%) and more metals order ppm (Figure 3). An examination of clay modified carbon paste electrode indicates some kind of agglomeration [13].

Electrodeposition of Polymer and Bacteria

Figure 4 represents the cyclic voltammograms corresponding to the polymerization of ε-caprolactone on the CPE-Ar electrode, recorded with a scanning speed equal to 80 mV / S in the NaCl solution (0.1 mol / l) containing 1 ml. - caprolactone in a range of potential between -2 V and 2 V. It can be seen that the presence of two anodic peaks at about -0.35 V and 0.46 V. These peaks are attributed to the oxidation of the monomer (ε-caprolactone). The decrease in the intensity of the currents during the sweeps accounts for the growth of the film deposited on the surface of the electrode with the number of cycles, the densities current of the voltammograms tend to 0 indicating that the developed polymer has an insulating character [14,15] (Figure 5,6).

Elaboration and characterization of bio- electrodes

The Figures 7 & 8 represent the cyclic voltammograms recorded respectively for the polymer-Clay - CPE and bacteria-polymer-Clay
CPE electrodes, in 0.1M NaCl medium at 100 mV/s. We find that the presence of bacteria in the mineral matrix leads to a relative increase in the current densities of the voltammogram in both anodic and cathodic scanning directions [16]. This effect appears clearly in square wave voltammetry. We find that the presence of bacteria in the mineral matrix leads to a relative increase in the current densities of the voltammogram in both anodic and cathodic scanning directions [16]. This effect appears clearly in square wave voltammetry.

Activity of the Bacteria-Polymer-CPE Electrode with respect to the Oxidation of Phenol

The oxidation of phenol on the bacteria-polymer-Clay-CPE electrode was studied by cyclic voltammetry (Figure 9). As we can see the recorded voltammogram, in electrolytic medium in the presence and absence of phenol. No peak is observed on the voltammogram in the absence of phenol, however, in the presence of phenol, there is the appearance of two anodic peaks corresponding to the oxidation of phenol, respectively at 0.5 V and 1.5 V.

Influence of Phenol Concentration

The effect of the scanning rate on the intensities of the phenol oxidation peaks was studied by cyclic voltammetry, in 0.1M NaCl (pH=7), containing 8 mM phenol, at scanning speeds included between 40 and 120 mV.s⁻¹ (Figure 10). The current densities vary linearly with the scanning speed (Figure 11), which shows that the system is in a diffusion mode.

Effect of Scanning Rate
The correlation equations and the limits of detections and quantifications calculated for the two electrodes polymer-Clay-CPE and bacteria-polymer-Clay-CPE are given as follows:

**Polymer-Clay-CPE**

\[ di = 0.260x + 1.431 \]

**Bacteria-polymer-Clay-CPE**

\[ di = 0.228x + 2.963 \]

The variation of anodic peak intensity as a function of phenol concentration was followed by cyclic voltammetry (Figure 12), square wave voltammetry (Figure 13), and impedance spectroscopy. We find that peak current densities increase with concentration, suggesting that the modified matrix has a large number of active sites. This result is confirmed by the impedance measurement, which shows a decrease in the electron transfer resistance [16].

**Evaluation of the Activity of the Bacteria-Polymer-Clay-CPE Modified Electrode for Phenol Detection**

**Calibration Curves**

Calibration curves were plotted from the square wave voltammetry at the surface of the polymer-Clay-CPE and bacteria-polymer-Clay-CPE electrodes (Figure 14). These experiments are carried out in the concentration range which varies between 4 and 12 mmol l\(^{-1}\). Table 1 groups together the correlation equations and the detection and quantification limits calculated for the two electrodes polymer-Clay -CPE and bacteria-polymer-CPE electrodes. We find that the presence of bacteria improves the sensitivity of the electrodes.

**Table 1:** The correlation equations and the limits of detections and quantifications calculated for the two electrodes polymer-Clay-CPE and bacteria-polymer-Clay-CPE in the presence of phenol.

| Electrode                   | Equation                        | \( R^2 \)   | D.L (mol. L\(^{-1}\)) | Q.L (mol. L\(^{-1}\)) |
|----------------------------|---------------------------------|-------------|------------------------|------------------------|
| Polymer-Clay-CPE+ Phenol   | \( di = 0.260x + 1.431 \)       | 0.853       | 2.041.10\(^{-5}\)     | 3.837.10\(^{-4}\)     |
| Bacteria-polymer-CPE+phenol| \( di = 0.228x + 2.963 \)       | 0.882       | 0.952.10\(^{-6}\)     | 1.863.10\(^{-5}\)     |

**The Activity of the Immobilized Bacteria**

The activity of bacteria immobilized on the polymer-Clay-CPE electrode surface calculated for the oxidation of phenol is:

\[ \alpha = \left(1 - \frac{I_{bact}}{I}\right) \times 100 \]

\[ \alpha = (1 - (4.25/2.62)) \times 100 \]

\[ = 62.21\% \]

**Conclusion**

In this work, we have developed an electrode, bacteria-polymer-CPE. This electrode has shown acceptable activity for the electrochemical oxidation of phenol. The presence of the polymer on the surface of the organic matrices causes an increase in the porosity of the electrode surfaces, which facilitates the adsorption of the electroactive molecule. The polymer developed on the surface of the electrodes facilitates the adhesion and development of a biofilm on the surface. Bacteria-polymer-CPE electrode activity was tested for phenol oxidation. The presence of a biofilm incorporated in the organic matrix facilitates the degradation of the phenol. The methods used, in this case the cyclic and square wave voltammetries, the impedance diagrams, have allowed a global and efficient characterization of the elaborated electrodes.

**References**

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