Electrochemical Method for Heavy Metals Detection by Inhibition of Acetylcholinesterase Immobilized on Pt-nanoparticles Modified Graphite Electrode

G. L. Turdean¹ and I. C. Popescu¹

Department of Physical Chemistry, University “Babes-Bolyai”, RO-400028 Cluj-Napoca, ROMANIA, gturdean@chem.ubbcluj.ro

Abstract. The optimization and the characterization of a new amperometric biosensor based on acetylcholinesterase (AChE), immobilized on a graphite electrode modified with Pt-nanoparticles (PtNP), are reported. The G/PtNP-AChE biosensor was used for heavy metals detection. The degree of inhibition (%I), the kinetic constants of the inhibition process, as well as the influence of the PtNP presence on these parameters were estimated.

Key words: Heavy metals, acetylcholinesterase, inhibition, wastewater.

Introduction

Heavy metal detection in environmental control is usually performed by spectroscopic, electrochemical or chromatographic methods (Malitesta and Gauscito, 2005), which are able to identify single elements and/or species at low concentrations, but can hardly be used for in-situ analysis and/or low-cost screening methods. Also, these tasks can be successful accomplished using biosensors. Thus, the enzyme biosensors used for the detection of commonly analytes in clinical and food analysis are based on the direct specific interaction between enzyme and substrate. Contrarily, in environmental analysis pollutants (e.g. pesticides, heavy metals etc.) the detection caused by the toxic substances is based on the inhibition of the enzymatic activity. The reduced specificity of inhibition phenomenon makes possible only the estimation of the sum parameters, i.e. total concentration of the substances belonging to a certain class. Because of their high specificity, it is expected that biosensor detects only contaminant ions, which are pollutant to life.

The aim of this work is to build-up and the subsequent application of a new amperometric biosensor based on acetylcholinesterase (AChE) immobilized on a graphite electrode modified with Pt-nanoparticles (PtNP) for heavy metals detection by using voltammetric techniques.

Materials and Methods

Acetylcholinesterase (EC. 3.1.1.7) and acetylthiocholine chloride were obtained from Sigma. Na₂HPO₄, NaH₂PO₄, EDTA disodium salt, copper (II) sulphate, cadmium (II) nitrite, chromium (III) nitrite, Zn(II) sulphate were from Reactivul-Bucuresti. The buffer solution and the heavy metals standard solutions were prepared by dissolving the appropriate amount of salts in distilled water. The Pt-nanoparticles (PtNP) were kindly provided by a nearby galvanisation factory.

Electrochemical experiments were carried out with a voltammetric analyser AUTOLAB PGSTAT 10 (Eco Chemie, Holland), connected to a PC computer. A graphite (inner diameter 3 mm from Ringsdorff, Germany) or platinum support electrode was modified by immobolizing the enzyme and the PtNP into a polymer matrix and were used as working electrode. An Ag/AgCl, KCl sat (Radiometer) and a platinum wire were used as reference and counter electrode, respectively.

All voltammetric measurements were performed in the presence of dissolved oxygen. The working electrode was immersed into a 10 ml cell containing 0.1 M...
phosphate buffer. A background voltammogram was recorded prior to the addition of corresponding aliquots of heavy metals standard solution or of wastewater sample. The preconcentration step consist in applying a deposition potential (usually -1.5 V) for a 120 s period and after in recording a voltammogram when a positive going square wave voltammetric (SWV) scan is applied with a frequency of 50 Hz, potential step of 5 mV and amplitude of 50 mV. All experiments were performed at room temperature.

The amperometric experiments were carried out in a stirred solution of phosphate buffer (pH 7) at a + 0.4 V vs. Ag/AgCl, KCl sat, to the working electrode.

Results and Discussion

On recorded SWVs well-defined peaks were observed together with low background currents, although no sample de-aeration was undertaken. Linear calibration curves having good correlation coefficients were obtained. The reproducibility and the robustness of the biosensor were also investigated (Legeai and Vittori, 2006).

The kinetic analysis of the results obtained with amperometric method is based on the experimental data collected by recording the current intensity of the biosensor corresponding to the produced thiocholine before and after addition of heavy metals solution. The thiocholine is the product of the hydrolysis reaction of acetylthiocholine catalysed by immobilized acetyleholinesterase on the surface of the working electrode (figure 1) corresponding to the following reaction (Stoytcheva and Sharkova, 2002):

\[(\text{CH}_3)_2\text{N}^+(\text{CH}_2)_2\text{SOCH}_3 + \text{H}_2\text{O} \rightarrow (\text{CH}_3)_2\text{N}^+(\text{CH}_2)_2\text{SH} + \text{CH}_3\text{COOH}\]

Fig. 1. Schematic diagram representing the exchange of electrons during the oxidation of thiocholine (RSH) on PtNP-AChE/G electrode

The results of inhibition of enzyme by the presence of heavy metals are the decrease of the generated thiocholine concentration.

The effect of inhibition exerted by the heavy metals was quantitatively estimated by the degree of inhibition (I, %) which was calculated by using the following equation: \(I = 100(I_0/I)\), where: \(I\) is the difference between the current intensities recorded in the presence and in the absence of heavy metals and \(I_0\) is the current intensity in the absence of heavy metals.

The graphic analysis of the experimental data following the Hanes-Woolf linearization method demonstrated the type of inhibition of acetyleholinesterase by the tested heavy metals. The inhibition process of the immobilised enzyme can be described by the following kinetic scheme:

\[E + S \xrightleftharpoons[k_1]{k_{cat}} ES \xrightarrow{k_{inh}} E + P\]

where: E is the immobilised enzyme; S is the free substrate, here ASChCl; P is the product, here thiocholine; E-S is the enzyme-substrate complex; E-Inh is the enzyme-inhibitor complex; E-S-Inh is the ternary complex containing enzyme-substrate-inhibitor; \(K_i\) and \(K_i\) are the equilibrium dissociation constants of the E-S-Inh complex and the E-Inh complex, respectively.

Thus, if the secondary plots of the slope vs. [Inh] and the 1/I axis intercepts vs. [Inh] (i.e. 1/I_max,inh vs. [Inh]) and \(K_{M,inh}/I_{max,inh}\) vs. [Inh], respectively) (Trevor Palmer, 1987) are linear, they allow to estimate the values of \(K_i\) and \(K_i\), respectively, from a single set of experimental data.

The influence of the PtNP presence on the inhibition response was also evaluated.

Conclusion

The proposed Pt-NP-AChE/G electrode was applied to detect heavy metals (Cu, Cd, Cr, Zn) in synthetic and real samples of wastewater. The bioelectrochemical methods for acetyleholinesterase inhibitor analysis, due to their simplicity, rapidity and sensitivity could be an alternative to the sophisticated classical tests applied for environmental monitoring.

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References


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