Development of an Amperometric Biosensor Based on Peroxidases from *Brassica napus* for the Determination of Ochratoxin a Content in Peanut Samples

Eduardo A. Ramírez¹, Adrián M. Granero², María A. Zón** and Héctor Fernández**

¹Laboratorio de Nanoscopias y Fisicoquímica de Superficies, Instituto de Investigaciones Fisicoquímicas Teóricas y Aplicadas (INIFTA), Diagonal 113 y calle 64 - La Plata, Argentina
²Departamento de Química, Facultad de Ciencias Exactas, Físico-Químicas y Naturales, Universidad Nacional de Río Cuarto, Agencia Postal No. 3 (5800) - Río Cuarto, Argentina.

**Abstract**

An amperometric biosensor based on *Brassica napus* hairy roots peroxidases to determine ochratoxin A is proposed by the first time. The enzymatic reaction rate was studied under different experimental conditions and the kinetics parameters were determined. The method employs a dialysis membrane covered, peroxidases entrapped and ferrocene-embedded carbon paste electrode (PBHR-Fc-CP) and is based on the fact that the decreased amount of H₂O₂ produced by the action of PBHR is proportional to the oxidised amount of ochratoxin A in the solution. The PBHR-Fc-CP biosensors showed a very good stability during at least five days. The reproducibility and the repeatability were 7.7% and 5.4%, respectively, showing a good biosensor performance. The calibration curve was linear in the ochratoxin A concentration range from 1 x 10⁻⁸ to 1.4 x 10⁻⁴ mol L⁻¹. The lowest concentration value measured experimentally for a signal to noise ratio of 3:1 was 5.7 x 10⁻⁹ mol L⁻¹.

**Keywords**: Ochratoxin A; *Brassica napus* hairy roots peroxidases; Amperometric biosensor

**Abbreviations**: OTA: Ochratoxin A; PBHR: *Brassica napus* Hairy Roots Peroxidases; CP: Carbon Paste; HRP: Horseradish Peroxidase; Fc: Ferrocene; PBS: Phosphate Buffer Solution

**Introduction**

The ochratoxins are a group of highly toxic fungal secondary metabolites that are produced by several species of the *Aspergillus* and *Penicillium* genera [1,2]. Ochratoxin A (OTA) is the most toxic and prevalent among these toxins, and is receiving increasing attention because of the hazard imposed on both human and animal health [3].

OTA or 7-carboxyl-5-chloro-8-hydroxyl-3,4-dihydro-3R-methyl-isocumarin-7-L-β-phenylalanine, is a colorless crystalline compound that belongs to a group of closely related derivatives of isocumarin linked to L-phenylalanine and classified as pentaketides [4] (Structure 1).

![Structure 1](image)

The toxin is potently nephrotoxic, teratogenic and an immunosuppressive agent [5-7]. It has been considered by the International Agency for Research on Cancer to be a potential carcinogen (group 2B) for human [8]. For this reason, many countries have restricted OTA levels in foods, with upper limits of 1 to 10 ppb depending on the type and quality of the foodstuff [9].

Peanuts are one of the most important agricultural products in Argentina. The central-south region of Córdoba Province produces 94% of the country’s peanuts. The peanut industry exports 90% of its product, being Argentina the second largest exporter in the world [10]. Each year, a significant portion of the peanuts produced can not be marketed because of fungal disease at the postharvest stage and mycotoxin contamination [11]. With imports and exports of agricultural commodities increasing in recent years, monitoring of OTA is essential for food safety in the world-wide market.

Many research groups have been studying methods for monitoring naturally occurring OTA in a number of agricultural commodities [9,12]. Many different analytical methods are now available for the determination of OTA in a variety of matrices [9]. Chromatographic methods such as thin-layer chromatography (TLC), gas chromatography (GC) and high-performance liquid chromatography (HPLC) with immunooaffinity columns and fluorescence detection [13-17] provide sensitive and specific techniques, but generally require multiple steps prior to detection. Today, the electroanalytical methods are very valuable alternatives to other methods because generally no separation procedure is required prior to substrate determination. Furthermore, the required instrumentation is typically less expensive, smaller amounts of solvent are necessary and analysis time is shorter for electroanalytical methods over chromatographic measurements.

Biosensors have been proposed as an efficient analytical tool for the determination of polyphenolic compounds, exhibiting advantages...
such as the minimal preparation of the sample, selectivity, sensitivity, reproducibility, rapid time of response and simple use for continuous on-site analysis [18-25].

Here, we report on the development of an amperometric biosensor based on peroxidases obtained from *Brassica napus* hairy roots (PBHR) to determine OTA content in solutions composed by commercial reagent and peanut samples.

The method employs carbon paste (CP) electrodes filled up with PBHR and ferrocene (Fc) at a given composition (PBHR–Fc–CP). The biosensor was covered externally with a dialysis membrane, which was fixed at the electrode body side part with a Teflon laboratory film and an O-ring. It is well known that phenolic and/or polyphenolic compounds can work as electron-donors for peroxidases in the catalytic reduction of H$_2$O$_2$ [26,27]. This approach allows detecting the decrease in H$_2$O$_2$ concentration in a solution after the oxidation of phenolic and/or polyphenolic compounds produced by the PBHR or horse radish peroxidase (HRP) in the presence of H$_2$O$_2$, given that HRP acts in cascade in the solution and PBHR at the electrode surface. The separation of the electrode surface from the solution by a semipermeable membrane allows minimizing the electrical noises as well as the fouling of the electrode surface.

**Materials and Methods**

**Reagents and materials**

*Brassica napus* hairy roots, obtained “in vitro” in our laboratory according to a procedure previously described by Agostini et al. [28] were used as the enzyme source.

Hydrogen peroxide (30%, v/v), pH 8.00 phosphate buffer solutions (PBS), sodium acetate, acetic acid, NaCl, HCl, NaOH and Na$_2$CO$_3$ were Merck p.a.; ochratoxin A (OTA), ferrocene (Fc) and o-dianisidine (PBS), sodium acetate, acetic acid, NaCl, HCl, NaOH and Na$_2$CO$_3$ were used as the enzyme source.

**Inoculation and Incubation conditions:** Peanuts were conditioned with an appropriate amount of water far from the $a_w$ levels. The substrate was kept at 4°C for 48 h with manual periodic shaking to allow absorption and to reach equilibrium. 20 g of peanuts were placed as a monolayer into sterile Petri dishes. Peanuts were inoculated centrally with 5 μL of a spore suspension (1 x 10$^6$ spores/mL) from a 7-day-old culture growing on 2% malt extract agar (MEA) [35]. Inoculated peanuts plates with the same aW were sealed in plastic containers. Each container had beakers with a glycerol/water solution at the same aW as the peanuts, to maintain a constant relativity humidity. Temperatures tested were 18 and 25°C for a period of 30 days.

**Ochratoxin A extraction:** OTA production was analyzed after 30 days of incubation. Five grams of a finely ground peanut sample was added to a 250 mL Erlenmeyer flask along with 50 mL of an acetonitrile-water mixture (84:16). The mixture was shaken in an orbital shaker for 30 min and filtered through filter paper (Whatman No. 1). The extract (7 mL) was added to a clean up column (MycoSep $^\textregistered$229 Ochra column, MFC, Romer Labs$^\text{®}$, Inc., MO., USA), and acidified with 70 μL of acetic acid. 4 mL of the purified extract was removed, evaporated to dryness, re-dissolved in the 200 μL of mobile phase (acetonitrile-water-acetic acid, 57: 41: 2), and injected into the HPLC.

**Electrochemical instruments and experimental measurements**

Cyclic voltammograms and amperometric measurements were performed with an Epsilon (BAS) potentiostat controlled by...
electrochemical analysis software. Electrochemical measurements were carried out in a 2 mL Pyrex cell. The working electrode was a CP disk of 1.6 mm diameter obtained from Bioanalytical System (BAS), Inc. The counter electrode was a platinum foil of large area (~2 cm²). An aqueous SCE was used as the reference electrode. Aliquots of 5 to 25 µL were added to the electrochemical cell for the determination of OTA content in solutions composed by commercial reagent and peanut samples contaminated with the mycotoxin, respectively. Amperometric measurements were performed at a potential of ~0.050 V vs SCE in solutions stirred at 1600 rpm. This operational applied potential was previously optimized by Granero et al. [36].

The Fc was used as a redox mediator. Biosensors were constructed by using the following procedure: PBHR, Fc and CP in different weight ratios were checked in order to obtain the best biosensor composition. Then, the optimal composition was used to fill up the CP electrodes. The biosensors were stored at 4°C in a dry container when they were not in use. Experiments performed with and without bubbling pure nitrogen in solutions did not show any significant difference. Therefore, measurements were carried out in non-deoxygenated solutions. Experiments were performed at 25 ± 0.2°C.

Results and Discussion

Response of PBHR-FC-CP biosensor versus H₂O₂ concentration

Paste electrodes were prepared by mixing different amounts of PBHR, Fc and CP. A cyclic voltammogram recorded in unstimred pH 8.00 PBS (blank solution) for the PBHR-Fc-CP biosensor showed an increase in the anodic current at potentials higher than 0.16 V vs SCE, which corresponds to the oxidation of Fc to Fc+. A cathodic peak with a peak potential at about 0.2 V was observed when the potential sweep direction was reversed at 0.6 V, which can be assigned to the reduction of Fc⁺ to Fc at the electrode surface (Figure 1). On the other hand, a cyclic voltammogram recorded when H₂O₂ was added to the reaction medium showed a cathodic current at potentials more negative than -0.3 V vs SCE, which corresponds to the reduction of H₂O₂ at the biosensor surface. Therefore, the base current was practically zero between -0.3 and 0.2 V in pH 8.00 PBS. This behavior was similar to that previously found by our group when a similar biosensor was employed to determine t-Res using the comercial reagent [36].

On the other hand, four biosensors were constructed to optimize

de amount of PBHR in the biosensor, for which the ratio of the other two components remained constant, i.e., 3 mg Fc and 45 mg CP. These biosensors were used to study their responses towards H₂O₂. Therefore, when only H₂O₂ was added to the stirred reaction medium, the enzymatic reaction took place between PBHR on the electrode surface and H₂O₂ penetrated the inner layer between the semi permeable membrane and the electrode surface. H₂O₂ was reduced to H₂O by PBHR and the enzyme was reduced to its native form by Fc, which was oxidized to Fc⁺. The Fc⁺ was then immediately reduced to Fc at the electrode surface held at a potential of -0.050 V vs SCE. Steady-

Figure 1: Cyclic voltammogram of PBHR-Fc-CP biosensor recorded in unstirred pH 8.00 PBS. V = 0.050 V s⁻¹.

Figure 2: Steady-state current responses on the addition of different H₂O₂ concentrations at the stirred pH 8.00 PBS reaction medium measured with a dialysis membrane covered PBHR-Fc-CP (3-6-45 mg) biosensor. c_H₂O₂: (1) 61.8 µmol L⁻¹, (2) 163.5 µmol L⁻¹, (3) 743 µmol L⁻¹ and (4) 1443 µmol L⁻¹.

Figure 3: Differences between the base and the steady-state currents, Δi_ss, as a function of H₂O₂ concentration.
state reduction currents ($I_s$) obtained after the addition of different aliquots of H$_2$O$_2$ are shown in Figure 2. The differences between the initial base current and the corresponding $I_s$ ($\Delta I_s$) were proportional to the H$_2$O$_2$ bulk concentration ($\varepsilon_{\text{H}_2\text{O}_2}$), showing a Michaelis-Menten type saturation (Figure 3). Plots of $I_s$ vs $I_{ss}$ (Eadie-Hofstee plots) were linear and from them we calculate the kinetic parameters of the PBHR-Fc-CP biosensor for H$_2$O$_2$ which are summarized in Table 1. Average values of H$_2$O$_2$ maximum current ($I_{\text{max,} \text{H}_2\text{O}_2}$) and Michaelis-Menten apparent constant ($K_{\text{A,} \text{H}_2\text{O}_2}$) were calculated from the intercept and the slope of Eadie-Hofstee plots for five replicate measurements. Therefore, from results shown in Table 1, biosensors constructed using 8–3–45 mg of PBHR, Fc and CP, respectively, were employed in the next experiments, considering that this electrode composition gave the maximum current value. Then, similar experiments were performed to optimize the Fc amount. Thus, four biosensors were constructed for which the ratio of the other two components remained constant, i.e., 8 mg PBHR and 45 mg CP. The differences between the initial base current and the corresponding $I_s$ ($\Delta I_s$) were proportional to the H$_2$O$_2$ bulk concentration ($\varepsilon_{\text{H}_2\text{O}_2}$), showing a Michaelis-Menten type saturation (Figure 4). Plots of $I_s$ vs $I_{ss}$ (Eadie-Hofstee plots) were linear. From those plots kinetic parameters of the PBHR-Fc-CP biosensor for H$_2$O$_2$ were calculated. They are summarized in Table 2. Average values of H$_2$O$_2$ maximum current ($I_{\text{max,} \text{H}_2\text{O}_2}$) and Michaelis-Menten apparent constant ($K_{\text{A,} \text{H}_2\text{O}_2}$) were calculated from the intercept and the slope of Eadie-Hofstee plots for five replicate measurements. Therefore, from results shown in Table 2, biosensors constructed using 8–6–45 mg of PBHR, Fc and CP, respectively, were employed in the next experiments, considering that this electrode composition gave the maximum current value. It should be noted that when the biosensor was built with HRP instead of PBHR as the recognition element, it showed less affinity by H$_2$O$_2$, as has been previously reported [36]. For this reason PBHR was selected as the recognition element for the assembly of the biosensor.

**Responses of PBHR-Fc-CP biosensors versus OTA concentration**

When both HRP and OTA were added to the stirred reaction medium composed by pH 8.00 PBS and a given H$_2$O$_2$ bulk concentration, the enzymatic catalytic cycle also took place in the solution bulk. The oxidized HRP, produced as a result of its reaction with H$_2$O$_2$, is reduced back to its native state by OTA. The decrease of H$_2$O$_2$ in the solution bulk was detected as a decrease in the Fc+ reduction current at the biosensor surface [36]. The optimum HRP concentration in solution was 10 nmol L$^{-1}$. Therefore, the addition of H$_2$O$_2$ to the pH 8.00 PBS + 10 nmol L$^{-1}$ HRP reaction medium produced a steady limiting current ($I_{ss}$) at -0.050 V vs SCE after 180 s, which corresponds to the reduction of Fc+ generated by HRP catalyzed reduction of H$_2$O$_2$ to H$_2$O (Figure 5). The difference between Is,lim with and without OTA ($\Delta I_{ss}$) corresponds to the H$_2$O$_2$ concentration decrease given that the enzymatic reaction was taking place in the bulk solution. The steady state

![Figure 4](https://example.com/figure4.png)

**Figure 4:** Differences between the base and the steady-state currents, $\Delta I_s$, as a function of $c_{\text{H}_2\text{O}_2}$ under the same experimental conditions as Figure 2. The PBHR-Fc-CP biosensor compositions were: (A) 8-1.5-45 mg (●) 8-3-45 mg (●) 8-6-45 mg (●) 8-9-45 mg.

| PBHR-Fc-CP biosensor composition (mg) | $I_{\text{max,} \text{H}_2\text{O}_2}$ (µA) | $K_{\text{A,} \text{H}_2\text{O}_2}$ (mmol L$^{-1}$) |
|--------------------------------------|-----------------|-----------------|
| 8-1.5-45                            | 1.64            | 0.7             |
| 8-3-45                              | 0.98            | 0.25            |
| 8-6-45                              | 2.17            | 0.8             |
| 8-9-45                              | 1.72            | 0.79            |

**Table 2:** $I_{\text{max,} \text{H}_2\text{O}_2}$ and $K_{\text{A,} \text{H}_2\text{O}_2}$ kinetics parameters obtained for different biosensor compositions shown in Figure 4 after adding different aliquots of H$_2$O$_2$ to pH 8.00 PBS.

![Figure 5](https://example.com/figure5.png)

**Figure 5:** Steady-state current responses on the addition of different OTA concentrations in a reaction medium under stirring containing 10 nmol L$^{-1}$ HRP + 100 µm L$^{-1}$ H$_2$O$_2$ in pH 8.00 PBS with a dialysis membrane covered PBHR-Fc-CP (8-6-45 mg) biosensor. OTA: (1) 30 µmol L$^{-1}$; (2) 73 µmol L$^{-1}$; (3) 113 µmol L$^{-1}$ and (4) 150 µmol L$^{-1}$.
currents obtained after the addition of OTA were reached at about 60 s.

Plots of $\Delta I_{\text{s,lim}}$ vs OTA concentration ($c_{\text{OTA}}$) also showed a Michaelis-Menten saturation. Figure 6 displays the maximum current ($I_{\text{max,OTA}}$) and the apparent Michaelis Menten constant ($K_{a,\text{OTA}}$) values obtained from the addition of different aliquots of OTA to the solution composed by pH 8.00 PBS + 10 nmol L$^{-1}$ HRP in the presence of different H$_2$O$_2$ bulk concentration. For the sensitivity ($S = I_{\text{max,OTA}} / K_{a,\text{OTA}}$ vs $c_{\text{OTA}}$, see insert of Figure 6) [37], the plateau was reached at 100-150 μmol L$^{-1}$ of H$_2$O$_2$. This means that at these peroxide concentrations, the bioelectrode responses are limited by the enzymatic kinetics rate. Therefore, a concentration of 100 μmol L$^{-1}$ of H$_2$O$_2$ was chosen as the best peroxide level in the reaction medium. It should be noted that when PBHR is used as biological material in solution, after the addition of OTA there were no appreciable changes in the amperometric response, which would indicate that OTA is not a good substrate for this enzyme. On the other hand, it has been found that OTA behave as a good substrate for HRP. Thus, we decided to use HRP as biological material in solution.

**Biosensor statistical parameters**

The PBHR-Fc-CP biosensor reproducibility was tested by measuring the calibration curve slope for OTA of five different bioelectrodes in a solution of pH 8.00 PBS + 10 nmol L$^{-1}$ HRP + 100 μmol L$^{-1}$ H$_2$O$_2$ as the reaction media. Percent relative standard deviations (%RSD) of 7.7% were obtained for OTA. The repeatability assays were performed carrying out six consecutive amperometric measurements on the same solution of pH 8.00 PBS + 10 nmol L$^{-1}$ HRP + 100 μmol L$^{-1}$ H$_2$O$_2$. The stability of PBHR-Fc-CP biosensor was tested by using the same biosensor to determine the slopes from several calibration curves (n = 5) for OTA. The slopes obtained were practically constant in the order of the experimental error until about five days, showing a good stability of PBHR biosensor. A noticeable decrease in the slope started from the fifth day and about 40% decrease was obtained on the seventh day. A linear variation was found for $\Delta I_{\text{s,lim}}$ vs OTA concentration ($c_{\text{OTA}}$) in the range from 1x10$^{-5}$ to 1x10$^{-4}$ mol L$^{-1}$. The lowest concentration value measured with PBHR-Fc-CP biosensor for a signal to noise ratio of 3:1 was 5.7 x 10$^{-7}$ mol L$^{-1}$ (2.3 ppb).

In order to achieve another criterion to test the accuracy of the biosensor method for OTA quantification in peanut samples the results were also investigated by the HPLC method [34]. Five samples were used to perform this analysis. The concentrations determined by HPLC-immunoaffinity column cleanup were plotted vs those obtained from the biosensor method. We found a linear variation with slope = 0.931 and correlation coefficient close to 1 (r = 0.9973) indicating a good correspondence between methods (Figure 7).

Experimental results obtained demonstrate that our enzymatic bioelectrode method is a very useful technique to detect and quantify OTA in peanut samples. The correlation between biosensor and HPLC method [34] indicates the effectiveness of the enzymatic electrode for OTA determination. While the detection limit determined is higher than values in some previous reports [38,39] still this electrode shows important advantages by using it for screening purposes due to its very simple construction and rapid measurements.

**Conclusions**

It is concluded that the PBHR biosensor exhibited a good performance, stability, reproducibility, repeatability, detection limit and linear range for the quantification of OTA. This good analytical performance allowed us to estimate the OTA in peanut samples using a very simple experimental procedure. The contents of OTA in samples of peanuts determined with the biosensor show a good correlation with values obtained with HPLC method. In addition, this method offers some advantages over the HPLC method, such as: short detection time (~1 min), small sample volumes, reduced use of solvents and low cost. These advantages indicate that a PBHR biosensor can be used as a useful tool for a rapid screening in the determination of OTA in peanuts samples.

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