

Biosensors of Laccase Based on Hydrophobic Ionic Liquids Derived from Imidazolium Cation

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Biossensores contendo lacase (*Aspergillus oryzae*) e líquidos iônicos derivados do cátion 1-butil-3-metilimidazol (BMI) associados com os ânions hexafluorofosfato (BMI·PF₆) ou bis(trifluorometilsulfonyl)imida (BMI·Tf₂N) foram construídos para determinação de adrenalina. O biossensor baseado no BMI·Tf₂N foi selecionado por apresentar maior resposta quando comparado ao BMI·PF₆. As melhores condições para otimização foram estabelecidas por voltametria de onda quadrada (amplitude 100 mV, frequência 10 Hz e incremento 4,0 mV). O melhor desempenho foi obtido em 50:20:15:15% (m/m/m/m) de pó de grafite:lacase:Nujol:LI em tampão acetato 0,1 mol L⁻¹ (pH 4,0). A curva analítica foi linear na faixa de concentração 2,49 × 10⁻⁶ a 2,27 × 10⁻⁴ mol L⁻¹ com limite de detecção de 5,34 × 10⁻⁷ mol L⁻¹. A recuperação de adrenalina em amostras injetáveis variou de 96,3 a 101,6%. Os resultados obtidos para a adrenalina usando o biossensor proposto e o procedimento da Farmacopéia Americana estão em concordância ao nível de confiança de 95%.

Biosensors based on laccase from *Aspergillus oryzae* and ionic liquids derived from the 1-butyl-3-methylimidazolium cation (BMI) associated with the anions hexafluorophosphate (BMI·PF₆) or bis(trifluoromethylsulfonyl)imide (BMI·Tf₂N) were constructed for adrenaline determination. The biosensor based on BMI·Tf₂N was selected by presenting higher response when compared with that based on BMI·PF₆. The best conditions for the optimization were established by square wave voltammetry (pulse amplitude 100 mV, frequency 10 Hz and scan increment 4.0 mV). The best performance was obtained with 50:20:15:15% (m/m/m/m) as the graphite powder:laccase:Nujol:ILs composition in 0.1 mol L⁻¹ acetate buffer solution (pH 4.0). The analytical curve was linear in the concentration range 2.49 × 10⁻⁶ to 2.27 × 10⁻⁴ mol L⁻¹ with a detection limit of 5.34 × 10⁻⁷ mol L⁻¹. The recovery of adrenaline in injectable samples ranged from 96.3 to 101.6%. The results obtained for the adrenaline using the proposed biosensor and the United States Pharmacopeia procedure were in agreement at the 95% confidence level.

Keywords: biosensor, laccase, ionic liquids, imidazolium, bis(trifluoromethylsulfonyl)imide, adrenaline

Introduction

Epinephrine, often called adrenaline, a hormone secreted from the adrenal glands, is an important catecholamine for message transfer in mammalian central nervous systems, present in nerve tissue and body fluids in the form of large organic cations. Studies show that many diseases are related to changes in its concentration. The quantitative determination of adrenaline is of great help in understanding the effects of the nervous system and

artificial substitutes, and is of great significance to research in the areas of pharmacology and the life sciences.^{1,2}

A number of methods have been applied in adrenaline determination including spectrophotometry,³ fluorimetry,⁴ liquid chromatography coupled with various detection techniques,⁵ capillary electrophoresis,⁶ chemiluminescence⁷ and electrochemiluminescence.⁸ Adrenaline is an electroactive molecule and it is important to examine its electrochemical behavior, thus electrochemical detection with various modified electrodes has also been reported.^{1,2,9-13} Brondani *et al.*⁹ have investigated the detection of adrenaline with a biosensor containing the ionic

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liquid 1-butyl-3-methylimidazolium hexafluorophosphate (BMI·PF₆) and a corn (*Zea Mays* L.) homogenate as a source of peroxidase immobilized in chitin chemically crosslinked with carbodiimide and glyoxal. The amperometric response of adrenaline was carried out at -0.23 V (vs. Ag/AgCl) via the electrochemical reduction of the product of the enzymatic reaction. Felix¹³ have developed a biosensor to determine adrenaline in pharmaceutical samples. The polyphenol oxidase enzymes present in the fibers of palm tree fruits (*Livistona chinensis*) catalyze the oxidation of adrenaline to adrenaline-quinone as a primary product. This product is then electrochemically reduced on the biosensor surface and the resulting current is used for its quantification. In addition, biosensors have some advantages, such as low cost, simplicity of construction and rapid response.

Biosensors are sensors modified with biological material (enzymes, antibodies, organelles, microorganisms and animal and plant tissues) integrated to the electronic transducer that converts a biochemical signal into a quantifiable analytical response.¹⁴ One of the most important enzymes in terms of applicability and versatility in industry is laccase. This is a type of copper-containing polyphenol oxidase, one of the extracellular glycoprotein enzymes obtained from white-rot fungi and other natural sources. This enzyme acts on substrates such as polyphenols, aminophenols and methoxy-substituted phenols by catalyzing the oxidation of their phenolic hydroxyl groups to phenoxy radicals while molecular oxygen is reduced to water.¹⁵ Numerous biosensors based on laccase have been described in the literature.¹⁶⁻²²

Ionic liquids (ILs) are organic salts, which are liquid at ambient temperature. Unlike traditional solvents, which can be described as molecular liquids, ILs are composed of ions. They possess a unique association of properties including non-volatility, non-flammability and excellent chemical and thermal stability.^{23,24} The class of ILs based on imidazolium cations exhibits a wider liquid range and an electrochemical window (up 7.0 V) and these have therefore become ideal electrolytes in electrochemical devices like batteries, capacitors and electrochemical sensors and biosensors.²⁴⁻²⁹ Nujol and paraffin are the most commonly used binder components of carbon electrode pastes. Recently, these traditionally used binders have been totally or partially replaced with ILs, indicating that they offer an attractive and efficient alternative, due to the high ionic conductivity and sensitivity of the resulting electrodes.^{9,10,17,18,30-36} There are several known hydrophobic ILs and those containing the bis(trifluoromethylsulfonyl)imide (Tf₂N⁻) anion offer several advantages for use in electrochemistry. In relation to the analogous ILs based on the PF₆⁻ anion they are less

viscous, water stable and have good conductivity, excellent thermal stability and a large electrochemical window.²⁴ Therefore, these hydrophobic ILs are promising candidates for the construction of novel biosensors.

In this paper the electrochemical behavior of adrenaline was investigated on biosensors containing the enzyme laccase from *Aspergillus oryzae* using ILs (BMI·Tf₂N or BMI·PF₆) and Nujol mixtures as the binder. The biosensors were then applied to adrenaline determination. The excellent properties of ILs effectively facilitate the fast electron transfer on the biosensor surface and improve sensitivity in the determination of this catecholamine in injectable samples. The experimental conditions, such as Nujol:ILs ratio, enzyme percentage, solution pH and square-wave voltammetry parameters (frequency, pulse amplitude and scan increment) for the analytical performance of the biosensors were optimized. The results obtained for the adrenaline determination in injectable samples using the proposed biosensor and those obtained using the spectrophotometric method are in agreement at the 95% confidence level.

Experimental

Reagents and solutions

All reagents were of analytical grade and used without further purification. Ultrapure water (18.2 MΩ cm) was obtained from a Milli-Q purification system and used throughout the experiments. Laccase from *Aspergillus oryzae* was purchased from Novozymes (Denilite® II BASE) in the form of microspheres. Graphite powder (grade #38) and Nujol were supplied by Fisher Scientific and Aldrich, respectively. Adrenaline, carbidopa, methyl dopa, levedopa, sodium chloride, sucrose, lactose, magnesium stearate, poly(ethylene glycol) and starch were purchased from Sigma. The ionic liquids 1-butyl-3-methylimidazolium hexafluorophosphate (BMI·PF₆) and 1-butyl-3-methylimidazolium bis(trifluoromethylsulfonyl)imide (BMI·Tf₂N) were synthesized as previously described in the literature.^{37,38} 0.1 mol L⁻¹ Acetate buffer solution (pH 4.0) was used as the supporting electrolyte. Injectable samples containing adrenaline were acquired from the University Hospital in Florianópolis (Santa Catarina, Brazil) and analyzed using the proposed biosensor and a standard spectrophotometric method.

Biosensor construction

To prepare 200.0 mg of paste, initially 40.0 mg (20%, m/m) of laccase and 100.0 mg (50%, m/m) of graphite

powder were hand mixed in a mortar for 10 min to ensure uniform dispersion. After this, 30.0 mg of Nujol (15%, m/m) and 30.0 mg of BMI·PF₆ or BMI·Tf₂N (15%, m/m) were added to this mixture which was mixed for at least 20 min to produce the final paste. A portion of the resulting paste was packed firmly into the cavity of a 1.0 mL plastic syringe and a copper wire was inserted, on the opposite side, to obtain the external electrical contact. The biosensors were stored at room temperature when not in use. For comparison, the carbon paste electrode (CPE) without laccase was prepared following the same steps.

Electrochemical and spectrophotometric measurements

Electrochemical measurements, using square-wave voltammetry, were performed on an Autolab PGSTAT12 potentiostat/galvanostat (Eco Chemie Utrecht The Netherlands) connected to data processing software (GPES, version 4.9.006, Eco Chemie). The electrochemical cell was assembled with a conventional three-electrode system: biosensors based on the different ILs BMI·Tf₂N-laccase and BMI·PF₆-laccase as the working electrodes (surface area 0.8 mm²), a platinum wire as the auxiliary electrode and an Ag/AgCl (3.0 mol L⁻¹ KCl) as the reference electrode. Voltammetric measurements were carried out in an unstirred, non de-aerated acetate buffer solution (pH 4.0) at 25.0 ± 0.5 °C and all potentials were measured and reported vs. Ag/AgCl. In a typical run, 10 mL of the acetate buffer solution was transferred to a clean, dry cell and the required volume of the adrenaline or injectable sample was added by micropipette. After a stirring period of 60 s to homogenize the solution, square-wave voltammograms were recorded.

To compare the results obtained by the methodology here proposed, we followed the UV spectrophotometric method established in the United States Pharmacopoeia.³⁹ A Hewlett-Packard (Boise, ID, USA) UV-Visible spectrophotometer, model 8452A, with a quartz cell (optical path of 1.00 cm) was used as the standard method for the adrenaline determination.

Determination of Michaelis-Menten constant

The Michaelis-Menten kinetic constant (K_m^{app}) was determined using the Lineweaver-Burk equation and employing the proposed biosensor in the concentration range of 2.49×10^{-6} to 5.18×10^{-4} mol L⁻¹ adrenaline in 0.1 mol L⁻¹ acetate buffer solution (pH 4.0) at 25.0 ± 0.5 °C. The current response was measured by square-wave voltammetry at a frequency of 10 Hz, pulse amplitude of 100 mV and scan increment of 4.0 mV.

Results and Discussion

Biosensors based on imidazolium ILs: voltammetric response to adrenaline

ILs are generally divided into two groups according to their solubility in water. The first type, water-miscible (hydrophilic) ILs, such as BMI·BF₄ have received less attention, because they have so far been considered unstable in aqueous solution. However, the second type, water-immiscible (hydrophobic) ILs, such as those based on PF₆⁻ and Tf₂N⁻ anions have proved to be promising for the construction the biosensors due to their hydrophobic nature, forming non-polarizable interfaces when in contact with water.²⁴ The imidazolium-based IL molecular structures are shown in Figure 1. In addition, the enzyme laccase is a multicopper oxidase which catalyze the four-electron reduction of molecular oxygen to water and the oxidation of a variety of aromatic compounds.¹⁵ Laccase immobilized on the biosensor catalyzes the oxidation of adrenaline present in the solution to the corresponding *o*-quinone. This product is electrochemically reduced back to adrenaline on the surface of the biosensor as shown in Figure 1. The resulting reduction current is used as the analytical response for adrenaline determination.

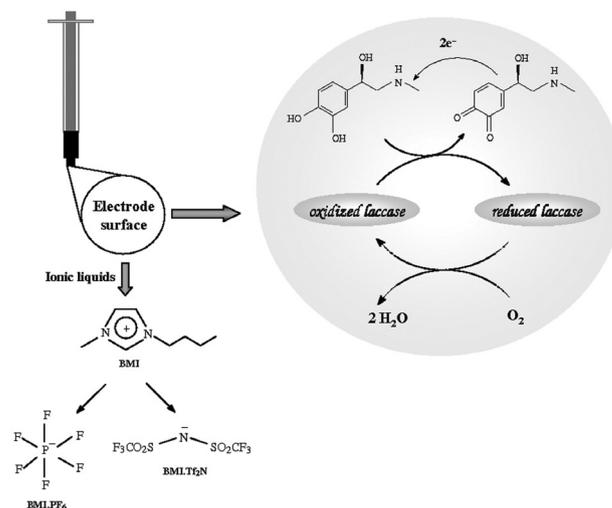


Figure 1. Enzymatic reaction involving adrenaline and laccase on biosensor surface and structure of the ILs derived from the 1-butyl-3-methylimidazolium cation (BMI) associated with the anions hexafluorophosphate (BMI·PF₆) or bis(trifluoromethylsulfonyl)imide (BMI·Tf₂N).

The square-wave voltammograms using 6.10×10^{-5} mol L⁻¹ adrenaline in 0.1 mol L⁻¹ acetate buffer solution (pH 4.0) were recorded between +0.4 and -0.3 V vs. Ag/AgCl (BMI·PF₆) and +0.4 and -0.5 V vs. Ag/AgCl (BMI·Tf₂N). The electrochemical behavior of adrenaline

was investigated in the presence of the biosensors based on laccase with BMI·PF₆:Nujol or BMI·Tf₂N:Nujol as binders and also, for comparison, in the presence the bare CPEs. Figure 2 shows the voltammograms obtained using (a') BMI·PF₆-CPE, (a) BMI·PF₆-biosensor, (b') BMI·Tf₂N-CPE, and (b) BMI·Tf₂N-biosensor. It can be noted that the BMI·PF₆- and BMI·Tf₂N-biosensors gave higher responses when compared with BMI·PF₆-CPE and BMI·Tf₂N-CPE, respectively. This is attributed to the presence of the laccase which catalyzes the oxidation of adrenaline to the corresponding *o*-quinone which is then electrochemically reduced back to adrenaline on the surface of the biosensor. The laccase enzyme favors the redox reaction, generating higher response for the reduction wave of *o*-quinone to adrenaline at an approach potential of -0.12 V for the biosensors. The peak observed around +0.1 V can be attributed to the neighboring group OH of the phenolic ring, however, the increase in this peak is not significant when compared with the peak at -0.12 V (attributed to the reduction of *o*-quinone to regenerate the initial compound).

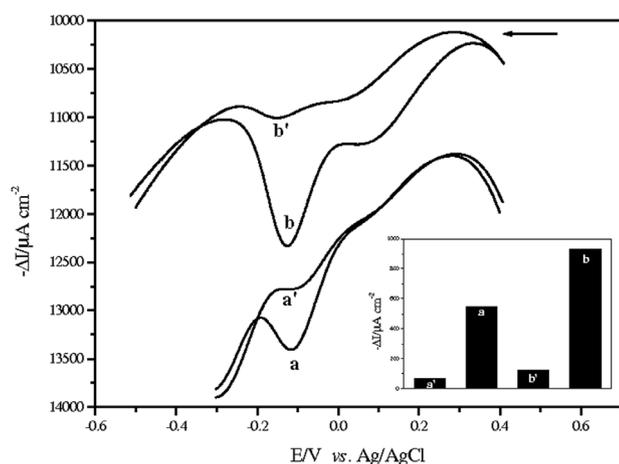


Figure 2. Square-wave voltammograms obtained using (a') BMI·PF₆-CPE, (a) BMI·PF₆-biosensor, (b') BMI·Tf₂N-CPE and (b) BMI·Tf₂N-biosensor with 6.10×10^{-5} mol L⁻¹ adrenaline in 0.1 mol L⁻¹ acetate buffer solution (pH 4.0); (a' and a) pulse amplitude 100 mV, frequency 20 Hz and scan increment 1.0 mV and (b' and b) pulse amplitude 100 mV, frequency 10 Hz and scan increment 4.0 mV. Inset: current values for each CPE and biosensor.

In addition, the BMI·Tf₂N:Nujol binder mixture led to an increase in the biosensor response and a large difference in the value of the cathodic current was observed when compared with the BMI·PF₆-biosensor. Due to hydrolysis of PF₆⁻, the imidazolium salts with Tf₂N⁻ are increasingly used as hydrophobic ILs.²⁴ Thus, the Tf₂N⁻ anion is preferred as the counter ion for the *N,N*-dialkyl substituted imidazolium cation because it is less viscous, more stable in water, and has good electrochemical conductivity and a wide electrode

potential window in aqueous solutions.²⁴ Therefore, the BMI·Tf₂N-biosensor was chosen for the development of this study, and also biosensors based on the laccase and ILs based on BMI·Tf₂N for the determination of adrenaline have not been described in the literature. Our research group has previously developed a biosensor containing laccase and ILs for the determination of phenolic compounds.^{17,18} As can be seen, for biosensor BMI·Tf₂N-laccase, a higher response for the reduction wave of *o*-quinone reduction to rutin at a potential of +0.31 V was observed when compared with DMI·Tf₂N-laccase and TDMI·Tf₂N-laccase. This behavior shows that the IL structure (*N*-alkyl chain length) is inversely proportional to the conductivity, that is, the biosensor based on TDMI·Tf₂N[(*Cn*MI)⁺], where *n* = 14 carbon atoms) gives the lowest response to rutin.¹⁷ The BMI·PF₆-biosensor showed a higher response for the reduction wave of *o*-quinone to rosmarinic acid when compared with the BMI·BF₄-biosensor. The presence of the hydrophobic IL·BMI·PF₆ offers a greater stability, good electrochemical conductivity and a wide electrode potential window in aqueous solutions.¹⁸ There are also three reports on the use of biosensors containing laccase and an IL·BMI·PF₆ for the inhibition of laccase activity by halide ions and as catalysts of the cathode of biofuel cells.⁴⁰⁻⁴²

Optimization of experimental parameters of the BMI·Tf₂N-biosensor

Various experimental conditions for the biosensor response, such as Nujol:BMI·Tf₂N ratios, percentage of laccase, pH, and square-wave voltammetry parameters, were investigated in order to obtain the best performance. The biosensor response was based on the cathodic peak current for a 6.10×10^{-5} mol L⁻¹ adrenaline solution using square-wave voltammetry. Table 1 summarizes the range over which each parameter was investigated and the optimal values found.

Table 1. Optimization of the experimental conditions

Parameters investigated	Range studied	Optimal value
Nujol:BMI·Tf ₂ N (% , m/m)	100:0-0:100	50:50
Laccase (% , m/m)	10-30	20
pH	4.0-8.0	4.0
Frequency (Hz)	10-100	10
Pulse amplitude (mV)	10-100	100
Scan increment (mV)	0.5-5.0	4.0

The effect of Nujol:BMI·Tf₂N compositions of (a) 100:0; (b) 75:25; (c) 50:50; (d) 25:75 and (e) 0:100%,

(m/m) on the biosensor response was investigated. It was observed that the analytical response increased with an increase in the percentage of BMI·Tf₂N used up to 50% and decreased at higher percentages (data not shown). Thus, the Nujol:IL ratio of 50:50% m/m was used for further construction of the biosensor. The influence of laccase in percentages from 10 to 30% (m/m) on the biosensor was investigated. The analytical signals (cathodic peak currents) became larger with increases in the enzyme composition up to 20% (m/m). Consequently, this is the most suitable amount of enzyme and it was selected for further studies. The composition of 50:20:15:15% (m/m/m/m) graphite powder:laccase:Nujol:BMI·PF₆, respectively, was used in the construction of the BMI·Tf₂N-biosensor. These results (Nujol:BMI·Tf₂N ratio 50:50% and laccase percentage of 20%) are in agreement with that obtained using the BMI·Tf₂N- and BMI·PF₆-biosensors in the rutin and rosmarinic acid determination, respectively.^{17,18}

With respect to effect of pH (4.0-8.0), the current response resulting from the laccase-catalyzed reaction achieved a maximum value at pH 4.0, decreasing at other pH values and this pH was therefore selected for further experiments. Electrochemical responses of the biosensor to adrenaline are dependent on the square-wave parameters. The effect of varying frequency (10 to 100 Hz), pulse amplitude (10 to 100 mV) and scan increment (1.0 to 5.0 mV) were investigated and the best analytical signals obtained for the BMI·Tf₂N-biosensor were 10 Hz, 100 mV and 4.0 mV, respectively. These experimental conditions were therefore selected for subsequent experiments.

Square-wave voltammograms and analytical curve of adrenaline

Under the optimal conditions mentioned above, the analytical curve obtained for the adrenaline was linear from 2.49×10^{-6} to 2.27×10^{-4} mol L⁻¹ and the regression equation found was $\Delta I = -8.40 + 9.19 \times 10^6[\text{adrenaline}]$ (ΔI : $\mu\text{A cm}^{-2}$; [adrenaline]: mol L⁻¹ and $r = 0.9999$). The detection limit (three times the blank signal/slope) was calculated using this calibration curve and was found to be 5.34×10^{-7} mol L⁻¹. Figure 3 shows these voltammograms, when the cathodic current density increased with adrenaline concentration, and the corresponding analytical curve can be seen in the inset.

The analytical features of the BMI·Tf₂N-biosensor developed in this study was compared with other biosensors for adrenaline determination.^{1,2,9-13} Our proposed BMI·Tf₂N-biosensor exhibited excellent performance in terms of wide linear range, good correlation coefficient and low limit of detection, and these results are comparable,

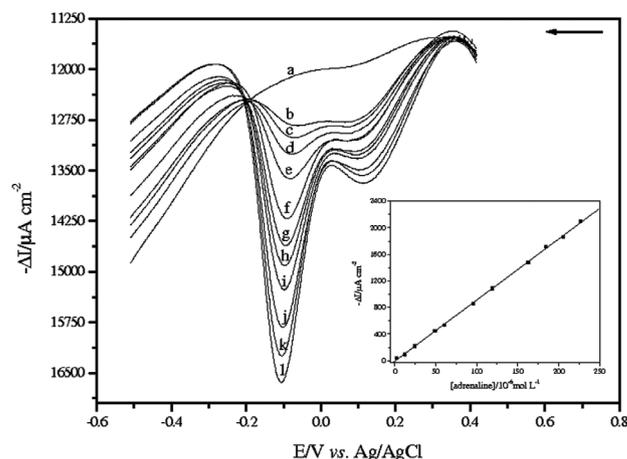


Figure 3. Square wave voltammograms obtained using the BMI·Tf₂N-biosensor for (a) blank in 0.1 mol L⁻¹ acetate buffer solution (pH 4.0) and adrenaline solutions at the following concentrations: (b) 2.49×10^{-6} ; (c) 1.24×10^{-5} ; (d) 2.48×10^{-5} ; (e) 4.91×10^{-5} ; (f) 6.10×10^{-5} ; (g) 9.61×10^{-5} ; (h) 1.19×10^{-4} ; (i) 1.63×10^{-4} ; (j) 1.85×10^{-4} ; (k) 2.06×10^{-4} and (l) 2.27×10^{-4} mol L⁻¹ at pulse amplitude 100 mV, frequency 10 Hz and scan increment 4.0 mV. Inset: the analytical curve of adrenaline.

or better than, those obtained for the biosensors described in the literature. The good analytical performance for the proposed biosensor can be attributed to the synergism of enzyme laccase and IL BMI·Tf₂N to the fast electron transfer between adrenaline and biosensor surface.

Repeatability, reproducibility and stability of the BMI·Tf₂N-biosensor

The study of the BMI·Tf₂N-biosensor was evaluated by investigating its performance for five parallel determinations of 6.10×10^{-5} mol L⁻¹ adrenaline solution in 0.1 mol L⁻¹ acetate buffer solution (pH 4.0). The relative standard deviation was calculated as 1.94%. Four biosensors were constructed using the same procedure and were independently used for the adrenaline determination under the conditions described above. All the biosensors revealed an acceptable reproducibility with a relative standard deviation of 2.83%, indicating that the results obtained with the proposed biosensor have a good reproducibility.

The lifetime and stability of the biosensor are very important features, and thus these parameters were also investigated and the response was recorded in a 6.10×10^{-5} mol L⁻¹ adrenaline solution (pH 4.0). To examine the long-term stability, the biosensor was tested by repeated measurements without surface renewal over 6 months. When the biosensor was stored at room temperature and measurements taken every 1-2 days in solutions of the same composition, no notable changes were observed in its response, indicating good stability and long lifetime.

Selectivity, interference, recovery study and determination of the adrenaline in injectable samples

The selectivity of the BMI-Tf₂N-biosensor was investigated by voltammetric method and in this experiment were used different compounds, such as, adrenaline, carbidopa, methyldopa, levedopa, sodium chloride, sucrose, lactose, magnesium stearate, poly(ethylene glycol) and starch. The responses were obtained from solutions containing 6.10×10^{-5} mol L⁻¹ of the substrates in acetate buffer solution. The sensitivity of the proposed biosensor followed the order: adrenaline > carbidopa > methyldopa > levedopa. No response was observed for sodium chloride, sucrose, lactose, magnesium stearate, poly(ethylene glycol) and starch. Thus, adrenaline was selected for the optimization of the biosensor and application to injectable samples.

In order to evaluate the effect of possible interferents in the determination of adrenaline (compounds mentioned above) were also investigated. Ratios of adrenaline concentration to that each substance was fixed at 1:1. Carbidopa, methyldopa, levedopa caused a positive interference (lower 5.0%) when present at the same concentration as the analyte and none of the other tested substances interfered in the adrenaline determination with the proposed procedure. In this way, the biosensor was able to determine adrenaline and it was evident that the influence of interfering species tested on the adrenaline response was negligible, indicating a high selectivity of the proposed biosensor.

Recovery experiments of adrenaline were applied using the proposed biosensor, by the standard addition method. Recoveries of 96.3 to 101.6% of adrenaline, given in Table 2, from injectable samples ($n = 3$) were obtained using the biosensor, the results being considered satisfactory, indicating the absence of matrix effects in these determinations. 8.70, 16.73 and 23.95 mg L⁻¹ adrenaline solutions were added to each sample and the cathodic peak current was registered for the pharmaceutical samples.

The proposed method was validated through the applicability of the BMI-Tf₂N-biosensor in the determination of adrenaline in injectable samples (A and B). The results of the analysis using the biosensor were compared with those obtained using the standard method³⁹ (Table 3). *T*-test (95%

Table 2. Recoveries of adrenaline (mg L⁻¹) standard solution in injectable samples using the BMI-Tf₂N-biosensor ($n = 3$)

Sample	Adrenaline/(mg L ⁻¹)		Recovery (%)
	added	Found	
A	8.7	8.4 ± 0.1	96.5
	16.7	16.1 ± 0.2	96.3
	23.9	24.3 ± 0.1	101.6
B	8.7	8.7 ± 0.1	100.0
	16.7	16.5 ± 0.1	98.8
	23.9	24.0 ± 0.2	100.4

confidence level) performed on these data showed that there was no significant difference between the results obtained using the two methods. It can thus be concluded that the proposed method is suitable for adrenaline determination in these samples.

Determination of the Michaelis-Menten constant

For the determination of the kinetic parameters of the enzyme in the modified electrodes, the apparent Michaelis-Menten constant have been calculated by fitting the variation of the peak current versus concentration of the adrenaline to the electrochemical Michaelis-Menten equation 1:

$$I = \frac{I_{\max}[S]}{K_m^{app}} \quad (1)$$

where [S] is the substrate concentration, I_{\max} is the maximum current and K_m^{app} represents the apparent Michaelis-Menten constant.⁴³ Figure 4 shows the variation in cathodic peak current versus adrenaline concentration (2.49×10^{-6} to 5.18×10^{-4} mol L⁻¹) and the Lineweaver-Burk graph is shown in the inset. As can be seen, the current response deviated from linearity and approached a constant value at high adrenaline concentrations. This is attributed to the maximum rate of the enzymatic reaction under adrenaline saturating conditions. The apparent Michaelis-Menten (K_m^{app}) was estimated using the Lineweaver-Burk equation and a value of $K_m^{app} = 3.25$ mmol L⁻¹ was obtained. This kinetic parameter is in agreement with the values reported for a biosensors using laccase from *Aspergillus oryzae*²¹ and *Trametes versicolor*.⁴⁴

Table 3. Adrenaline determination in injectable samples using the biosensor and the standard spectrophotometric method

Sample	Label value	Standard method ³⁹	BMI-Tf ₂ N-biosensor	R _{e1} / (%)	R _{e2} / (%)
A*	1000.0	1020.0 ± 0.1	1005.8 ± 0.1	+0.6	-1.4
B*	1000.0	970.0 ± 0.2	993.0 ± 0.1	-0.7	+2.3

R_{e1} = BMI-Tf₂N-biosensor vs. label value; R_{e2} = BMI-Tf₂N-biosensor vs. standard method. * mg L⁻¹; $n = 3$; confidence level of 95%.

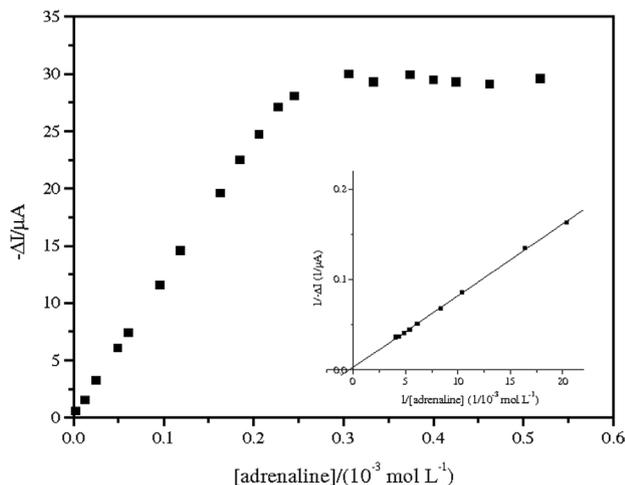


Figure 4. Cathodic current response of the biosensor as a function of adrenaline concentration. Inset: the Lineweaver-Burk graph.

Conclusions

In this paper, we present novel biosensors based on laccase from *Aspergillus oryzae* and the ILs, BMI·PF₆ or BMI·Tf₂N, used as the binders, together with Nujol. The hydrophobic ILs derived from the imidazolium cation associated with the PF₆⁻ and Tf₂N⁻ were used for the determination of adrenaline in injectable samples and were found to be promising candidates for biosensor construction. The BMI·Tf₂N-based biosensor had more favorable analytical features of high sensitivity, linear calibration range, low detection limit, rapid response, good reproducibility and long-term stability. In addition, the low cost, renewability, simplicity and fast construction of the biosensor, using square-wave voltammetry as the analysis method, make it superior to other techniques for adrenaline determination. The results indicate that this catecholamine was successfully determined and the results obtained were satisfactory when compared with the spectrophotometric method.

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References

- Zeng, Y.; Yang, J.; Wu, K.; *Electrochim. Acta* **2008**, *53*, 4615.
- Amorim, C. G.; Araújo, A. N.; Montenegro, M. C. B. S. M.; *Talanta* **2007**, *72*, 1255.
- Solich, P.; Polydorou, C. K.; Koupparis, M. A.; Efstathiou, C. E.; *J. Pharm. Biomed. Anal.* **2000**, *22*, 781.

- Tzontcheva, A.; Denikova, N.; *Clin. Chim. Acta* **2000**, *297*, 217.
- Sabbioni, C.; Saracino, M. A.; Mandrioli, R.; Pinzauti, S.; Furlanetto, S.; Gerra, G.; Raggi, M. A.; *J. Chromatogr., A* **2004**, *1032*, 65.
- Sänger-van de Griend, C. E.; Ek, A. G.; Widahl-Näsman, M. E.; Andersson, E. K. M.; *J. Pharm. Biomed. Anal.* **2006**, *41*, 77.
- Du, J.; Shen, L.; Lu, J.; *Anal. Chim. Acta* **2003**, *489*, 183.
- Su, Y.; Wang, J.; Chen, G.; *Talanta* **2005**, *65*, 531.
- Brondani, D.; Dupont, J.; Spinelli, A.; Vieira, I. C.; *Sens. Actuators, B* **2009**, *138*, 236.
- Brondani, D.; Scheeren C. W.; Dupont, J.; Vieira, I. C.; *Sens. Actuators, B* **2009**, *140*, 252.
- Qiao, J. X.; Luo, H. Q.; Li, N. B.; *Colloids Surf., B* **2008**, *62*, 31.
- Beitollahi, H.; Ardakani, M. M.; Ganjipour, B.; Naeimi, H.; *Biosens. Bioelectron.* **2008**, *24*, 362.
- Felix, F. S.; Yamashita, M.; Angnes, L.; *Biosens. Bioelectron.* **2006**, *21*, 2283.
- Grieshaber, D.; Mackenzie, R.; Vörös, J.; Reimhult, E.; *Sensors* **2008**, *8*, 1400.
- Widsten, P.; Kandelbauer, A.; *Enzyme Microb. Technol.* **2008**, *42*, 293.
- Wang, Z.; Xu, Q.; Wang, J. H.; Yang, Q.; Yu, J. H.; Zhao, Y. D.; *Microchim. Acta* **2009**, *165*, 387.
- Franzoi, A. C.; Migowski, P.; Dupont, J.; Vieira, I. C.; *Anal. Chim. Acta* **2009**, *639*, 90.
- Franzoi, A. C.; Dupont, J.; Spinelli, A.; Vieira, I. C.; *Talanta* **2009**, *77*, 1322.
- Fernandes, S. C.; Oliveira, I. R. W. Z.; Fatibello-Filho, O.; Spinelli, A.; Vieira, I. C.; *Sens. Actuators, B* **2008**, *133*, 202.
- Zhang, Y.; Zeng, G. M.; Tang, L.; Huang, D. L.; Jiang, X. Y.; Chen, Y. N.; *Biosens. Bioelectron.* **2007**, *22*, 2121.
- Santhiago, M.; Vieira, I. C.; *Sens. Actuators, B* **2007**, *128*, 279.
- Jarosz-Wilkolazka, A.; Ruzgas, T.; Gorton, L.; *Talanta* **2005**, *66*, 1219.
- Dupont, J.; *J. Braz. Chem. Soc.* **2004**, *15*, 341.
- Wei, D.; Ivaska, A.; *Anal. Chim. Acta* **2008**, *607*, 126.
- Galiński, M.; Lewandowski, A.; Stepniak, I.; *Electrochim. Acta* **2006**, *51*, 5567.
- Hayashi, K.; Nemoto, Y.; Akuto, K.; Sakurai, Y.; *J. Power Sources* **2005**, *146*, 689.
- Liu, H.; He, P.; Li, Z.; Liu, Y.; Li, J.; *Electrochim. Acta* **2006**, *51*, 1925.
- Suarez, P. A. Z.; Selbach, V. M.; Dullius, J. E. L.; Einloft, S.; Piatnicki, C. M. S.; Azambuja, D. S.; de Souza, R. F.; Dupont, J.; *Electrochim. Acta* **1997**, *42*, 2533.
- Dai, J.; Cheng, J.; Jin, J.; Li, Z.; Kong, J.; Bi, S.; *Electrochem. Commun.* **2008**, *10*, 587.
- Sun, W.; Li, Y.; Gao, H.; Jiao, K.; *Microchim. Acta* **2009**, *165*, 313.
- Li, Y.; Liu, X.; Zeng, X.; Liu, Y.; Liu, X.; Wei, W.; Luo, S.; *Microchim. Acta* **2009**, *165*, 393.

32. Musameh, M. M.; Wang, J.; *Anal. Chim. Acta* **2008**, *606*, 45.
33. Sun, W.; Jiang, Q.; Yang, M.; Jiao, K.; *Bull. Korean Chem. Soc.* **2008**, *29*, 915.
34. Sun, W.; Li, Y.; Yang, M.; Li, J.; Jiao, K.; *Sens. Actuators, B* **2008**, *133*, 387.
35. Ding, C.; Zhang, M.; Zhao, F.; Zhang, S.; *Anal. Biochem.* **2008**, *378*, 32.
36. Zhang, Y.; Zheng, J. B.; *Electrochim. Acta* **2007**, *52*, 7210.
37. Suarez, P. A. Z.; Dullius, J. E. L.; Einloft, S.; de Souza, R.F.; Dupont, J.; *Polyhedron* **1996**, *15*, 1217.
38. Cassol, C. C.; Ebeling, G.; Ferrera, B.; Dupont, J.; *Adv. Synth. Catal.* **2006**, *348*, 243.
39. United States Pharmacopeia, *The Official Compendia of Standards*, Twinbrook Parkway:Rockville, 2007.
40. Liu, Y.; Huang, L.; Dong, S.; *Biosens. Bioelectron.* **2007**, *23*, 35.
41. Liu, Y.; Dong, S.; *Electrochem. Commun.* **2007**, *9*, 1423.
42. Liu, Y.; Dong, S.; *Biosens. Bioelectron.* **2007**, *23*, 593.
43. Haghighi, B.; Gorton, L.; Ruzgas, T.; Jönsson, L. J.; *Anal. Chim. Acta* **2003**, *487*, 3.
44. Freire, R. S.; Durán N.; Kubota, L. T.; *Talanta* **2001**, *54*, 681.

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