Determination of hydrogen peroxide with an enzymeless amperometric sensor based on poly(vinylferrocene)-supported Ag nanoparticles

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Abstract: An enzymeless electrochemical sensor for detection of low amounts of $\text{H}_2\text{O}_2$ with the aid of Ag nanoparticles supported on conducting poly(vinylferrocene) (PVF) film was developed. Experimental results revealed that contribution of Ag nanoparticles led to remarkable improvement by means of reduction potential and reduction current. Influence of experimental parameters (i.e. polymeric film thickness, concentration of Ag precursor, immersion time in precursor solution, reduction time, and reduction potential) were investigated. The Ag/PVF-modified electrode system was characterized physically by scanning electron microscopy. The results revealed that the sensor developed was easy-to-prepare, economic, selective, and sensitive, with a fast response time of 3 s. The linear concentration range of the sensor was 0.1–50 mM, with a sensitivity of 14.1 $\mu\text{A} \text{mM}^{-1}$ and a limit of detection of 0.94 $\mu\text{M}$. Finally, interference effects of uric acid, ascorbic acid, dopamine, and glucose molecules were studied and no significant interference was observed at physiological levels.

Key words: Hydrogen peroxide determination, electrochemical sensor, poly(vinylferrocene), silver nanoparticles, modified electrode

1. Introduction

Hydrogen peroxide ($\text{H}_2\text{O}_2$)—a very small compound in nature—is an interesting molecule because of its importance in both industrial processes and biological processes. It is not only involved in many processes such as food processing, textile industry, pulp and paper bleaching, and minerals processing, but also utilized in diverse areas such as pharmaceutical research and organic synthesis.\(^{1,2}\) It is even used as an oxidant in fuel cell systems. Last but not least, it is the by-product of several reactions that involve enzymes such as glucose oxidase, cholesterol oxidase, and lactate oxidase. It is also a substrate for the enzyme horseradish peroxidase. Therefore, the reliable, accurate, sensitive, rapid, and low-cost determination of $\text{H}_2\text{O}_2$ has been practically important and widely investigated.

Conventional techniques for $\text{H}_2\text{O}_2$ determination, such as chemiluminescence,\(^{3,4}\) fluorescence,\(^{5,6}\) and spectrophotometry\(^{7,8}\) are complex, costly, and time-consuming. In comparison, electrochemistry can offer simple, rapid, sensitive, and cost-effective means, since $\text{H}_2\text{O}_2$ is an electroactive molecule.\(^{9,10}\) However, electrochemical applications are limited by slow electrode kinetics, high overpotential, and possible interferences from other existing electroactive species in real samples. Thus, current research on electrochemical $\text{H}_2\text{O}_2$ detection is mainly focused on electrode modifications in order to decrease the overpotential and increase the electron

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transfer kinetics. For these considerations, a large range of materials such as redox proteins, dyes, transition metals, metal oxides, redox polymers, and carbon nanotubes are studied. On the other hand, in recent years, nanomaterials, especially nanoparticles, have been successfully applied to electrochemical sensors improving the sensitivity of the sensor due to large surface area and porous structure leading to a decrease in the overpotential for many analytes.

The main objective of the current study was to prepare an enzymeless electrochemical sensor for detection of low amounts of \( \text{H}_2\text{O}_2 \) with the aid of Ag nanoparticles supported on conducting poly(vinylferrocene) (PVF) film. Recently, the relatively high cost and lack of long-term stability of enzyme biosensors has led to an effort to develop new sensors that mimic the function of enzymes. Avoiding use of an enzyme during the construction of a sensor is also encouraged because of the lack of protein denaturation and subsequent decrease in sensitivity. For these purposes, nanoparticles are considered excellent substitutes for enzymes. Many nanoparticle-based electrochemical \( \text{H}_2\text{O}_2 \) sensors have been reported in the literature, including Ag, Au, Pt, and Pd. Among these, Ag nanoparticles have been successfully applied as an effective catalyst for amperometric \( \text{H}_2\text{O}_2 \) detection without use of an enzyme. Ag nanoparticles are known to be responsible for the sensor response due to the electrochemical reduction of \( \text{H}_2\text{O}_2 \).

In the present work, a modified electrode system was developed and used as the working electrode of an amperometric \( \text{H}_2\text{O}_2 \) sensor. PVF, a redox polymer, was used as support material on a Au electrode for incorporation of Ag nanoparticles, with a view to combine the features of PVF and Ag nanoparticles for enhanced sensor response. To the best of our knowledge, this is the first study that combines use of Ag nanoparticles with PVF polymer, and the sensor has the advantage of easy preparation together with a low limit of detection (LOD) compared with similar studies in the literature. One of the most important benefits of using PVF is the ease of modification of the electrode and control of the modification parameters during the procedure. It is well known from previous studies that PVF provides an excellent medium for biological compounds and has been employed in many electrochemical sensors. In order to obtain maximum current for \( \text{H}_2\text{O}_2 \) reduction, the influence of experimental parameters was investigated. The Ag/PVF modified electrode system was characterized physically by scanning electron microscopy (SEM).

2. Results and discussion

PVF-supported Ag nanoparticles were prepared following a very simple, three-step procedure: (i) coating the Au electrode with polymer film, (ii) immersion of the film coated electrode in aqueous solution of Ag\(^+\) at open circuit, (iii) electrochemical reduction of Ag\(^+\) ions by constant potential electrolysis.

During the procedure, the polymer film was deposited onto the working electrode by electrooxidation of methylene chloride solution of PVF containing tetra-n-butylammonium perchloride (TBAP) as the supporting electrolyte. The thickness of the polymer film was controlled by the charge passed during electrolysis. As an example, a charge of \( 1 \times 10^{-3} \) C corresponds to \( 1.32 \times 10^{-6} \) mol of the oxidized PVF per cm\(^2\) (dry thickness of approximately 300 \( \mu \)m, which corresponds to about \( 3 \times 10^5 \) layers). The resulting film has a porous structure containing ClO\(_4^-\) ions as the counter ion, ferrocene, and ferrocenium groups. From previous experience, we know that the PVF-coated electrode is readily capable of immobilizing Ag\(^+\) ions from an aqueous solution. Therefore, in the second step, the PVF-coated electrode was immersed in a stirred aqueous solution of AgNO\(_3\) in order to immobilize the Ag\(^+\) ions in the polymeric matrix prior to electrochemical reduction at constant
potential to obtain PVF-supported Ag nanoparticles. The resulting Ag/PVF coated electrode was used as an enzymeless sensor for $\text{H}_2\text{O}_2$ detection.

In order to evaluate the influence of Ag nanoparticles on the response of the modified electrode system towards $\text{H}_2\text{O}_2$ reduction, cyclic voltammograms (CVs) of $\text{H}_2\text{O}_2$ solution were recorded and compared with those of an uncoated Au electrode, a PVF-coated Au electrode, and the Ag/PVF modified Au electrode (Figure 1). In the uncoated Au and PVF-coated Au electrodes, reduction of $\text{H}_2\text{O}_2$ was observed to a smaller extent at higher potentials (Figures 1a and 1b). However, when the PVF-coated electrode was decorated with Ag nanoparticles, both the reduction potential and the reduction peak current values were improved considerably (Figure 1c). CVs recorded with the Ag/PVF modified Au electrode in $\text{N}_2$-saturated phosphate buffer system (PBS, pH = 7.0) in the absence and presence of 1.0 mM $\text{H}_2\text{O}_2$ are also presented as an inset in Figure 1.

![Figure 1](image1.png)

Figure 1. CVs recorded with (a) uncoated Au electrode, (b) PVF-coated Au electrode, (c) Ag/PVF-modified Au electrode in $\text{N}_2$-saturated PBS (pH = 7.0) in the presence of 1.0 mM $\text{H}_2\text{O}_2$. Inset: CVs recorded with Ag/PVF-modified Au electrode in $\text{N}_2$-saturated phosphate buffer system (PBS, pH = 7.0) in the absence (---) and presence (-- of 1.0 mM $\text{H}_2\text{O}_2$. Scan rate: 100 mV s$^{-1}$.

The effect of scan rate during electrocatalytic reduction of $\text{H}_2\text{O}_2$ with the Ag/PVF-modified Au electrode was investigated. As seen from the CVs in Figure 2a, cathodic peak potential shifts gradually towards negative potentials as the scan rate increases, which confirms kinetic limitation in the electrochemical reaction. Additionally, a plot of peak current versus the square root of scan rate ($v^{1/2}$) displayed a linear relationship, indicating that the process is diffusion-controlled rather than surface-controlled (Figure 2b).

2.1. Optimization of experimental parameters

Several experimental parameters, such as polymeric film thickness and AgNO$_3$ concentration, were investigated to obtain the maximum peak current for $\text{H}_2\text{O}_2$ reduction for best sensor response. Optimization of these parameters was performed according to the reduction peak current values recorded during CVs of 1.0 mM $\text{H}_2\text{O}_2$ in 50 mM PBS buffer (pH = 7.0).

Thickness of the polymer coating on the working electrode is an important parameter because it affects the porosity of the polymer film. As mentioned above, the thickness of the polymer film was controlled by the
Figure 2. (a) CVs recorded with Ag/PVF-modified Au electrode in N₂-saturated PBS (pH = 7.0) in the presence of 5.0 mM H₂O₂ at different scan rates: (A) 10, (B) 25, (C) 50, (D) 75, and (E) 100 mV s⁻¹. (b) Plot of peak potential versus square root of scan rate.

charge passed during electrolysis. To evaluate the influence of polymer film thickness, PVF films of different thicknesses were coated on the working electrode and immersed in stirred 1.0 mM AgNO₃ solution for 10 min. The electrode was then washed with distilled water and transferred into a cell containing 50 mM PBS (pH = 7.0). Then constant potential electrolysis was applied at −0.2 V (versus saturated calomel electrodes [SCE]) for 300 s in order to reduce the adsorbed Ag⁺ ions to metallic Ag, and CVs of H₂O₂ solution were recorded and compared. Polymer film corresponding to 3.0 mC showed the best performance and therefore the optimum thickness was determined (Figure 3a).

Concentration of AgNO₃ solution was also an important parameter that probably controlled the size of the Ag nanoparticles, along with the immersion time in Ag⁺ solution. We studied with Ag⁺ solutions within the concentration range 1.0 to 6.0 mM, keeping all other parameters constant throughout the trials (3.0 mC charge passed during polymer coating, 10 min immersion time in Ag⁺ solution, 300 s electrolysis at −0.2 V versus SCE). As can be seen from Figure 3b, H₂O₂ reduction peak current was significantly higher for the 4.0 mM AgNO₃ solution. We also studied the influence of immersion time in stirred AgNO₃ solution and obtained a maximum at 15 min for H₂O₂ reduction peak current (Figure 3c).

Probably the most outstanding feature of our work is the ease of the nanoparticle synthesis compared with similar studies in the literature, because synthesis of metal nanoparticles generally requires complex and time-consuming procedures. However, we managed to obtain Ag nanoparticles dispersed on the PVF matrix simply by using bulk electrolysis at moderate potentials through electroreduction of the adsorbed ionic species. After immobilizing Ag⁺ ions in the polymer matrix via a simple dip-coating process, we applied constant potential electrolysis in 50 mM PBS (pH = 7.0) to obtain Ag nanoparticles. According to the experimental results, only 4 min of electrolysis at −0.2 V (versus SCE) was enough to obtain improved catalytic activity from the Ag/PVF system towards H₂O₂ reduction (Figures 3d and 3e).

2.2. Physical characterization of the modified electrode

The morphology and structure of the polymer film before and after incorporation of Ag nanoparticles were characterized by SEM. For SEM studies, the modified electrode system was prepared using pencil graphite electrodes (PGEs) with identical surface area. As clearly seen in Figure 4a, the polymer film has a porous structure that supports adsorption of the Ag⁺ ions, and the Ag nanoparticles are well dispersed in the polymeric
Figure 3. Influence of experimental parameters on reduction peak current of \( \text{H}_2\text{O}_2 \) (a) polymeric film thickness, (b) \( \text{AgNO}_3 \) concentration, (c) immersion time in \( \text{AgNO}_3 \), (d) reduction potential of \( \text{Ag}^+ \) ions, (d) reduction time of \( \text{Ag}^+ \) ions. (Data obtained from CVs of \( \text{N}_2 \)-saturated PBS (pH = 7.0) in the presence of 1.0 mM \( \text{H}_2\text{O}_2 \)).

matrix (Figure 4b) with an average particle size of 12 nm. The corresponding energy-dispersive X-ray (EDX) spectra of the PVF film before and after incorporation of Ag nanoparticles and the elemental mapping of the Ag/PVF catalyst system is presented in Figures 4c and 4d, indicating the presence of Ag atoms well dispersed in the polymer matrix.

2.3. Effect of pH

In order to apply the modified electrode to amperometric \( \text{H}_2\text{O}_2 \) detection, we controlled the catalytic activity of the Ag/PVF system towards \( \text{H}_2\text{O}_2 \) reduction at the physiological pH (7.4). For this purpose, we studied
the effect of pH of the H$_2$O$_2$ solution on the response of the Ag/PVF modified electrode, which was prepared at the optimum conditions. It is clearly seen from Figure 5 that the activity was highest at around pH 7.0 and was also active between pH values 7.0 and 7.5.

Figure 4. SEM images of (a) PVF-modified, (b) Ag/PVF-modified PGE. (c) EDX spectra and composition of PVF- and Ag/PVF-modified PGE. (d) Elemental mapping of Ag/PVF-modified PGE.
3.0 4.0 5.0 6.0 7.0 8.0

Figure 4. Continued.

Figure 5. Influence of pH on the response of the Ag/PVF-modified electrode prepared under optimum conditions. (Data obtained from CVs of N₂-saturated PBS (pH = 7.0) in the presence of 1.0 mM H₂O₂.)

2.4. Chronoamperometry studies and amperometric detection of H₂O₂

Chronoamperometry is a useful method for determining the catalytic rate constant ($k_{cat}$) using the equation

$$I_{cat}/I_L = \pi^{1/2}(k_{cat}C_0t)^{1/2},$$

(1)
where $I_{\text{cat}}$ is the catalytic current of the Ag/PVF catalyst in the presence of H$_2$O$_2$, $I_L$ is the limiting current in the absence of H$_2$O$_2$, $C_0$ is the bulk concentration of H$_2$O$_2$ (M), and $t$ is the elapsed time (s). For this purpose, we sketched $I_{\text{cat}}/I_L$ versus $t^{1/2}$ using chronoamperometric data obtained from 10 mM solution of H$_2$O$_2$ at –0.5 V versus SCE (not shown). From the slope of $I_{\text{cat}}/I_L$ versus $t^{1/2}$ plot, $k_{\text{cat}}$ was $2.4 \times 10^3$ M$^{-1}$ s$^{-1}$. This value is in the same order of magnitude with H$_2$O$_2$ sensors in the literature.$^{54,55}$

Chronoamperometric data can also be used for estimation of the diffusion coefficient ($D$) for an electroactive material using Cottrell’s equation:

$$I = nFAC(D/\pi t)^{1/2},$$

(2)

where $n$ is the number of electrons, $F$ is the Faraday constant, $A$ is the electrode area, $C$ is the bulk concentration of the analyte, and $t$ is the elapsed time.$^{56}$ In order to determine the specific surface area ($A$) of the Ag/PVF-modified Au electrode, cyclic voltammetry experiments were performed with the modified electrode in 1.0 mM K$_3$[Fe(CN)$_6$] + 50 mM KCl at different scan rates (between 10 and 100 mV s$^{-1}$). From the slope of anodic peak current ($I_{\text{pa}}$) versus square root of scan rate ($v^{1/2}$) plot, the estimated value for $A$ was calculated as 0.123 cm$^2$ using the following equation:

$$I_{\text{pa}} = (2.69 \times 10^5)n^{2/3}AD^{1/2}v^{1/2}C_0$$

(3)

using $7.6 \times 10^{-6}$ cm$^2$ s$^{-1}$ as the diffusion coefficient for K$_3$[Fe(CN)$_6$].$^{56,57}$ According to the chronoamperometric data of 10 mM H$_2$O$_2$ solution, the diffusion coefficient of H$_2$O$_2$ was $7.40 \times 10^{-6}$, which is reasonable compared with the literature.$^{53,58}$

The performance of the Ag/PVF-modified Au electrode as an enzymeless amperometric H$_2$O$_2$ sensor was evaluated by chronoamperometry. Figure 6a shows the typical current–time curve of the sensor prepared under optimized conditions after successive additions of certain concentrations H$_2$O$_2$ to a continuously stirred N$_2$-saturated PBS solution (pH = 7.4). The applied potential was –0.5 V versus SCE. The results revealed that the sensor had a short response time (under 5 s) and was sensitive to the addition of H$_2$O$_2$ aliquots in a wide concentration range.

The calibration curve obtained from chronoamperometric data indicated a good linear relationship between the peak current and H$_2$O$_2$ concentration over a wide concentration range (0.1 to 50 mM) with a correlation coefficient of 0.9964 (Figure 6b). LOD was calculated as 0.94 µM according to Borgmann et al. ($N = 6$) with a signal-to-noise ratio of 3, which is lower than many of the previously reported sensors.$^{59}$ Sensitivity of the sensor was 14.1 µA mM$^{-1}$. As a comparison, analytical performances of some enzymeless H$_2$O$_2$ sensors based on Ag nanoparticles is presented in Table 1 in terms of LOD and linear range.

2.5. Interference study

The influence of common interfering species such as uric acid (UA), ascorbic acid (AA), dopamine (DA), and glucose (Glu) was investigated. As clearly seen in Figure 7, none of these species demonstrated a significant interference to the signal of 0.5 mM H$_2$O$_2$ at a concentration of 0.5 mM under optimum working conditions. Thus it can be concluded that H$_2$O$_2$ can be determined selectively with the prepared sensor in the presence of these interferents.
2.6. Real sample analysis

The Ag/PVF-modified Au electrode was applied to the determination of H$_2$O$_2$ in hair dye oxidant cream. An oxidant cream sample (3.0 mL) with a reported H$_2$O$_2$ concentration of 9% (v/v) was diluted to 100 mL. From this solution, three different samples were prepared (1.0 mM, 5.0 mM, and 10.0 mM) and analyzed using standard addition. As seen from the analysis results in Table 2, the modified electrode presented a poor recovery (%) response for determination of H$_2$O$_2$ in concentrated real samples. However, in samples with moderate concentrations, the results indicate that the Ag/PVF-modified Au electrode is applicable for real sample analysis.

2.7. Reproducibility of Ag/PVF-modified Au electrode

In order to investigate the reproducibility of the Ag/PVF-modified Au electrode, reduction current obtained by chronoamperometry using 10 mM H$_2$O$_2$ solution was monitored daily for 10 days. When not in use, the modified electrode was kept at 4 ºC, open to air, and no visual change was observed on the electrode surface. The current response was the same for the first two days and decreased gradually afterwards to 29% of the initial current in 10 days (Figure 8).
Table 1. Comparison between performances of different enzymeless \( \text{H}_2\text{O}_2 \) sensors containing Ag nanoparticles.

<table>
<thead>
<tr>
<th>Modified electrode</th>
<th>LOD (( \mu )M)</th>
<th>Linear range (mM)</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>MWCNT/Ag nanohybrids/Au</td>
<td>0.5</td>
<td>0.05–17</td>
<td>[34]</td>
</tr>
<tr>
<td>Ag NPs/ZnO/GCE</td>
<td>0.42</td>
<td>0.002–5.5</td>
<td>[31]</td>
</tr>
<tr>
<td>AgNP/rGO/GCE</td>
<td>7.1</td>
<td>0.1–80</td>
<td>[35]</td>
</tr>
<tr>
<td>AgNP/F-SiO(_2)/GO/GCE</td>
<td>4</td>
<td>0.1–260</td>
<td>[36]</td>
</tr>
<tr>
<td>AgNPs/rGO/GCE</td>
<td>3.6</td>
<td>0.1–100</td>
<td>[20]</td>
</tr>
<tr>
<td>AgNP/TiO(_2) NWs/GCE</td>
<td>1.70</td>
<td>0.1–60</td>
<td>[30]</td>
</tr>
<tr>
<td>AgNPs/rGO/GCE</td>
<td>1.80</td>
<td>0.1–60</td>
<td>[32]</td>
</tr>
<tr>
<td>Ag NW array</td>
<td>29.2</td>
<td>0.1–3.1</td>
<td>[19]</td>
</tr>
<tr>
<td>AgNPs/MWCNTs-IL/GCE</td>
<td>0.0039</td>
<td>( 1.2 \times 10^{-5} )–( 4.8 \times 10^{-3} )</td>
<td>[21]</td>
</tr>
<tr>
<td>AgNP/rGO/GCE</td>
<td>10</td>
<td>0.05–5</td>
<td>[22]</td>
</tr>
<tr>
<td>AgNP-PmPD/GCE</td>
<td>0.88</td>
<td>0.1–10</td>
<td>[10]</td>
</tr>
<tr>
<td>AgNPs/Ox-pTTBA/MWCNT/GCE</td>
<td>0.24</td>
<td>0.010–0.260</td>
<td>[11]</td>
</tr>
<tr>
<td>Ag/PVF/Au</td>
<td>0.94</td>
<td>0.1–50</td>
<td>This work</td>
</tr>
</tbody>
</table>

Figure 7. Current–time responses of Ag/PVF-modified Au electrode exposed to UA, AA, DA, and Glu (0.5 mM each) in the presence of 0.5 mM \( \text{H}_2\text{O}_2 \) at an applied potential of \(-0.5 \) V versus SCE.

Table 2. Determination of \( \text{H}_2\text{O}_2 \) in hair dye oxidant cream.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Added (mM)</th>
<th>Found (mM)</th>
<th>Recovery (%)</th>
<th>RSD (%, n = 3)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1.00</td>
<td>1.04</td>
<td>104.0</td>
<td>2.7</td>
</tr>
<tr>
<td>2</td>
<td>5.00</td>
<td>4.82</td>
<td>96.4</td>
<td>1.4</td>
</tr>
<tr>
<td>3</td>
<td>10.00</td>
<td>29.8</td>
<td>298.0</td>
<td>6.0</td>
</tr>
</tbody>
</table>
2.8. Conclusions

A modified electrode system was developed based on Ag nanoparticles supported on conducting PVF film. As expected, the synergetic effect of combining PVF and the Ag nanoparticles resulted in enhanced sensor response even without using an enzyme for $\text{H}_2\text{O}_2$ detection. One of the most important features of the sensor is the ease of preparation, together with the low LOD compared with similar studies in the literature. The developed sensor exhibited a very good linearity within the concentration range of 0.1 mM to 50 mM, with a detection limit of 0.94 $\mu$M. Sensitivity of the sensor was calculated as 14.1 $\mu$A mM$^{-1}$. The interference study indicated that the enzymeless sensor was not affected by common interferents such as UA, AA, DA, and Glu. These characteristics make the Ag/PVF system an excellent amperometric sensor for detection of low levels of $\text{H}_2\text{O}_2$.

3. Experimental

3.1. Reagents and instruments

Poly(vinylferrocene) was synthesized according to the procedure described by Aso et al.\textsuperscript{60} Vinylferrocene was obtained from Sigma-Aldrich (St. Louis, MO, USA). 2,2-Azo-bis(2-methyl-propionitrile) was obtained from Alfa (Ronkonkoma, NY, USA). TBAP ($\geq$99.0%) was purchased from Fluka (Munich, Germany). Methylene chloride (HPLC grade) was obtained from Sigma-Aldrich. Hydrogen peroxide solution was diluted from 30% aqueous stock solution (Sigma-Aldrich). AgNO$_3$, NaH$_2$PO$_4$, Na$_2$HPO$_4$, UA, AA, and DA were purchased from Sigma-Aldrich and used as received.

All solutions were deoxygenated by purging high-purity nitrogen gas prior to use in the electrochemical experiments and all experiments were carried out at ambient temperature.

Electrochemical experiments were performed using a CHI 600E electrochemical workstation. SEM images were recorded using a JEOL model JSM-7001F (Tokyo, Japan).

3.2. Electrodes

Electrochemical experiments were carried out in a three-electrode system glass cell. A Au disc electrode ($r = 1 \text{ mm}$) was used as the working electrode. This electrode was polished using alumina (first 1.0 $\mu$m, then 0.3 $\mu$m), then rinsed with triple-distilled water, cleaned in an ultrasonic bath, and dried. Ag/AgCl and SCE were used as reference electrodes in methylene chloride and aqueous media, respectively. A Pt wire was used as the counter electrode. In order to record SEM images, the PVF coating and the Ag/PVF catalyst system were prepared on disposable PGEs.
3.3. Preparation of the Ag/PVF-modified Au electrode

During the procedure, the polymer film was first deposited onto the working electrode by electrooxidation of 1.0 mg mL\(^{-1}\) PVF solution in methylene chloride containing 0.1 M TBAP as the supporting electrolyte. The potential was held constant at 0.7 V versus Ag/AgCl throughout the electrolysis. In the second step, the PVF-coated electrode was immersed in a stirred aqueous solution of AgNO\(_3\) in order to immobilize the Ag\(^+\) ions in the polymeric matrix. After this step, the electrode was washed thoroughly with triple-distilled water and placed in an electrochemical cell containing 50 mM phosphate buffer system (pH = 7.0) in order to reduce the Ag\(^+\) ions at constant potential to obtain PVF-supported Ag nanoparticles.

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References