GLUCOSE ELECTRODES BASED ON CROSS-LINKED [Os(bpy)2Cl]+/2+ COMPLEXED POLY(1-VINYLIMIDAZOLE) FILMS

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Abstract

Enzyme electrodes based on a redox hydrogel formed upon complexing water-soluble poly(1-vinylimidazole) ("PVI") with [Os(bpy)2Cl]⁺ ("Os") and cross-linked with water-soluble poly(ethylene glycol) diglycidyl ether (molecular weight 400, peg 400) are described. The properties of the electrodes depended on their polymers' osmium content, its extent of cross-linking, on the pH, and the ionic strength in which they were used. The redox hydrogels' electron diffusion coefficients (Dₑ) increased with osmium content of their polymers. The Dₑ values were 1.5 X 10⁻⁸ cm² s⁻¹, 1.3 X 10⁻⁸ cm² s⁻¹, 4.3 X 10⁻⁹ cm² s⁻¹ for PVI₃-Os, PVI₅-Os, and PVI₁₀-Os, respectively, the subscripts indicating the number of monomer units per osmium redox center. Dₑ decreased with increasing ionic strength and increased upon protonation of the polymer. In glucose electrodes, made by incorporating in the films glucose oxidase (GOX) through covalent bonding in the cross-linking step, glucose was electrooxidized at 250 mV (SCE). The characteristics of these electrodes depended on the GOX concentration, film thickness, O₂ concentration, pH, NaCl concentration, and electrode potential. The steady-state glucose electrooxidation currents were independent of the polymers' osmium content in the studied (3 - 10 osmium centers per monomer unit) range. Electrodes containing 39% GOX reached
Enzyme electrodes based on a redox hydrogel formed upon complexing watersoluble poly(1-vinylimidazole) ("PVI") with [Os(bpy)2Cl]+ ("Os") and cross-linked with water-soluble poly(ethylene glycol) diglycidyl ether (molecular weight 400, peg 400) are described. The properties of the electrodes depended on their polymers' osmium content, its extent of cross-linking, on the pH, and the ionic strength in which they were used. The redox hydrogels' electron diffusion coefficients \(D_e\) increased with osmium content of their polymers. The \(D_e\) values were \(1.5 \times 10^{-8} \text{ cm}^2 \text{s}^{-1}\), \(1.3 \times 10^{-8} \text{ cm}^2 \text{s}^{-1}\), and \(4.3 \times 10^{-8} \text{ cm}^2 \text{s}^{-1}\). (Continued)
$10^{-9}$ cm$^2$ s$^{-1}$ for PVI$_3$-Os, PVI$_5$-Os, and PVI$_{10}$-Os, respectively, the subscripts indicating the number of monomer units per osmium redox center. $D_e$ decreased with increasing ionic strength and increased upon protonation of the polymer. In glucose electrodes, made by incorporating in the films glucose oxidase (GOX) through covalent bonding in the cross-linking step, glucose was electrooxidized at 250 mV (SCE). The characteristics of these electrodes depended on the GOX concentration, film thickness, O$_2$ concentration, pH, NaCl concentration, and electrode potential. The steady-state glucose electrooxidation currents were independent of the polymers’ osmium content in the studied (3 - 10 osmium centers per monomer unit) range. Electrodes containing 39% GOX reached steady-state glucose electrooxidation current densities of 400 $\mu$A cm$^{-2}$ and, when made with thick gel films, were selective for glucose in the presence of physiological concentrations of ascorbate and acetaminophen.
steady-state glucose electrooxidation current densities of 400 μA cm⁻² and, when made with thick gel films, were selective for glucose in the presence of physiological concentrations of ascorbate and acetaminophen.

Introduction

Redox hydrogel films are unique in having both adequate electron diffusion coefficients and in being permeable to water soluble substrates and products of enzymatic reactions. When a cross-linked redox polymer network electrically "wires" an enzyme that is covalently bound to it, then the gel and the current collecting metal form enzyme electrodes. Such electrodes are potentially useful in applications where release of diffusional mediators from the electrodes is to be avoided and where small size is important. Here we show that hydrogels, based on the Os complex of PVI are adequate electron conductors and that glucose oxidase (GOX) transfers electrons through the gel's polymer network to electrodes. The hydrogels are made by cross-linking with poly(ethylene glycol) diglycidyl ether (peg 400), PVIₙ-Os, and GOX.

A reference polymer with which the PVIₙ-Os polymers is compared is poly(4-vinylpyridine) ("PVP") complexed with osmium ("POs") and partially quaternized with bromoethylamine ("POs-EA"). POs is only marginally soluble in water except at high osmium loading. POs-EA is, however, water soluble and easy to cross-link with diepoxides such as peg 400 that binds POs-EA amines and lysyl functions of enzyme proteins. The resulting redox hydrogel adheres well to the electrodes. In contrast with the PVP derived polymers, the PVIₙ-Os polymers are highly water soluble and do not require quaternization with bromoethylamine for easy cross-linking with water-soluble diepoxides. The redox potential of PVIₙ-Os is 200 mV (SCE) vs 280 mV (SCE) for POs or POs-EA.
Because the operating potentials of the resulting enzyme electrodes is also lower, currents resulting from electrooxidation of some interferants are reduced.

Experimental Section

**Chemicals.** 1-vinylimidazole (Aldrich), K$_2$OsCl$_6$ (Johnson Matthey), 2,2'-bipyridine (Aldrich), Na-HEPES (sodium 4-(2-hydroxyethyl)-1-piperazineethanesulfonate) (Aldrich), poly(ethylene glycol) diglycidyl ether (Polysciences, peg 400, cat. no. 08210), and glucose oxidase (EC 1.1.3.4) from Aspergillus niger (type X-S, 198 units) were used as received. 2,2'-azo-bis(isobutyronitrile) (Polysciences, AIBN) was purified by double recrystallization from methanol and stored at $-20^\circ$C. Os(bpy)$_2$Cl$_2$ was prepared by a reported procedure.$^6$

**Poly(1-vinylimidazole), (PVI).** Bulk polymerization of PVI was carried out by heating 6 ml of 1-vinylimidazole and 0.5 g of AIBN at $70^\circ$C for 2 hours under Ar. A dark yellow precipitate formed soon after heating. After allowing the reaction mixture to cool, the precipitate was dissolved in methanol and added dropwise to a strongly stirred solution of acetone. The filtered precipitate was a pale yellow hygroscopic solid. The molecular weight of the polymer was found, by HPLC analysis using a Synchrom Catsec 300 column with 0.1% trifluoroacetic acid and 0.2 M NaCl as the elutant, to be about 7000 g mol$^{-1}$. The flow rate in the molecular weight determination was 0.4 ml/min and poly(2-vinylpyridine) was used as the standard.

**PVI$_n$-Os, n = 3, 5, 10.** Osmium derivatized polymers were prepared by a procedure similar to that of Forster and Vos$^6$ where the appropriate amount of Os(bpy)$_2$Cl$_2$ was refluxed with PVI in ethanol for 3 days. The elemental analyses were: Calculated for C$_{35}$Cl$_2$H$_{34}$N$_{10}$Os $\cdot$ 3 H$_2$O: C, 46.2; H, 4.4; N, 15.4; Os, 20.9; Found C, 46.4; H, 4.2; N, 15.3; Os, 18.4. Calculated for
C₄₅Cl₂H₄₆N₁₄Os • 5 H₂O: C, 47.7; H, 5.0; N, 17.3; Os, 16.8. Found C, 48.2; H, 4.5; N, 16.2; Os, 15.4. Calculated for C₇₀Cl₂H₇₆N₂₄Os • 10 H₂O: C, 49.6; H, 5.7; N, 19.9; Os, 11.2. Found C, 51.1; H, 5.2; N, 18.7; Os, 11.0. The three osmium derivatized polymers are referred to as PVl₃-Os, PVl₅-Os, and PVl₁₀-Os where the subscript represents the number of vinylimidazole units per Os(bpy)₂Co.

**Electrodes.** Rotating disk electrodes were prepared by embedding vitreous carbon rods (3 mm diameter, V-10, Atomergic) in a teflon shroud using a low viscosity epoxy (Polysciences, cat. no. 01916). Electrodes were prepared by syringing a 2 μl aliquot of 5 mg ml⁻¹ PVl₇-Os solution onto the electrode surface (0.071 cm²). Next, a 2 μl volume of a 4 mg ml⁻¹ (10 mM HEPES pH = 8.1) solution of glucose oxidase was added on the electrode and stirred with a syringe needle. In the final step, 1.2 μl of a 2.5 mg/ml solution of peg 400 was added to the electrode and stirred. The electrode was allowed to cure for at least 48 hours under vacuum.

**Measurements.** Electrochemical measurements were performed with a Princeton Applied Research 175 universal programmer, a Model 173 potentiostat and a Model 179 digital coulometer. The signal was recorded on a Kipp and Zonen X-Y-Y' recorder. Rotating disk electrode experiments were performed with a Pine Instruments AFMSRX rotator with an MSRS speed controller. The three-electrode cell contained 0.1 M NaCl buffered with phosphate (20 mM, pH = 7.2). The current response of the glucose electrodes were found to have a sigmoidal dependence on potential resulting in a maximum oxidation current at potentials ≥300 mV (SCE). In the constant potential experiments, the working electrode was poised at 0.4 V (SCE) to ensure that the maximum oxidation current to glucose was observed. The chronoamperometric measurements were performed with a Princeton Applied Research Model 273 potentiostat. The
potential was initially held at 0.0 V (SCE) for 15 s, then stepped to 0.8 V for 0.3 s. The slopes of the resulting $i$ vs $t^{-1/2}$ plots were unaffected upon varying the residence time (0.1 to 0.5 s). Three hundred data points were recorded. The reported values are averages of either four or five measurements.

Results and Discussion

Cyclic Voltammetry of PVI$_n$-Os-Peg 400 Coated Electrodes.

The effect of the extent of cross-linking on the electrochemical behavior of PVI$_n$-Os ($n = 3, 5, 10$) films cross-linked with 5% to 20% peg 400 was determined for electrodes in which the quantity of redox polymer was held constant, with the films cured for $>48$ hours. Coulometry (by cyclic voltammetry) showed that approximately 50% of the osmium was immobilized and that it did not vary with the extent of cross-linking. The cross-linked films adhered well to the vitreous carbon electrodes and retained about 95% of their electroactive osmium when soaked in a stirred phosphate buffer at room temperature for 72 hours.

The separation of the oxidation and the reduction peaks ($\Delta E_P$) of the voltammograms at 100 mV s$^{-1}$ scan rate remained constant at $\sim 100$ mV through the cross-linking range studied. In contrast to POs-EA films, $\Delta E_P$ increased with the cross-linker (peg 400) concentrations.$^3$

The peak width at half-height ($E_{\text{fwhm}}$) (for the oxidation wave measured at 1 mV s$^{-1}$) increased with the extent of cross-linking from 60 mV to 110 mV for PVI$_n$-Os ($n = 3, 5, 10$). While in the POs-EA system the $E_{\text{fwhm}}$ of the oxidation peak was narrower than that of the reduction peak, in the PVI$_n$-Os-peg 400 systems the two were precisely equal.$^3$

Diffusion Coefficients. $D_e C_p^2$ values ($D_e$ being the electron diffusion coefficient and $C_p$ the concentration of the redox couple in the film) were measured for the cross-linked films by potential step chronoamperometry.$^7$ The
film thickness was calculated by assuming that all three osmium polymers had a density of 1 g cm\(^{-3}\). There was no discernible trend in the dependence of \(D_e C_p^2\) on cross-linking (Table 1). \(D_e\) was, however, higher in the PVI5-Os polymer films than in the cross-linked POs-EA films by a factor of five.\(^3\) \(D_e\) was higher and nearly identical for PVI3-Os and PVI5-Os films, and lower for PVI10-Os films.

**Effects of Ionic Strength and pH on the Electron Diffusion Coefficients.** As seen in Figure 1, \(D_e C_p^2\) decreased with increasing NaCl concentration through the 88 mM to 2000 mM range. Specifically, \(D_e C_p^2\) decreased by about 50% when the NaCl concentration was raised from 88 mM to 1000 mM NaCl.

Figure 2 shows the pH dependence of \(D_e C_p^2\) in 0.2 M NaCl. The observed decrease in \(D_e C_p^2\) at pH = 4 coincides with the pKa of PVI.\(^8\) As the PVI is deprotonated the repulsive forces within the network are decreased and the mobility of chain segments that apparently controls electron transport diminishes. The pH dependence of \(D_e\) was similar to that observed in the POs-EA-peg 400 system\(^3,9\) where charging of the network and absence of screening also facilitated electron transfer. In PVI\(_n\)-Os networks swelling at low pH should decrease \(C_p\). Nevertheless, the increase in \(D_e\) was so large that \(D_e C_p^2\) still increased.

**Permeability of PVI\(_n\)-Os-Peg 400 Hydrogels.** Permeability of the hydrogels to water soluble substrates and products is important in biosensor applications. For the analysis of permeability p-benzoquinone was chosen as a model compound, because of its known electrochemistry and its non-ionic nature. The diffusional and kinetic characteristics of benzoquinone partitioning through an electrode film has been analyzed by Saveant et al\(^{10}\), who measured limiting current densities for rotating disk electrodes and obtained Koutecky-Levich plots for polymer film electrodes. Benzoquinone is reduced at pH = 7 at
-0.2 V negative of the redox potential of PVIn-OsII/III, i.e., at a potential where benzoquinone can not be catalytically reduced by PVIn-OsII. The benzoquinone electroreduction current densities were measured as a function of film thickness. The current densities decreased upon increasing film thickness more rapidly than they did in POs-EA films. A cross-linked PVIn-Os film having $6.6 \times 10^{-8}$ mol/cm$^2$ of osmium reached a current density of 1.2 mA/cm$^2$ per mM benzoquinone, while a POs-EA film with $9.7 \times 10^{-8}$ mol/cm$^2$ Os had a current density of 2 mA cm$^{-2}$ per mM benzoquinone. The relatively lesser permeability of PVIn-Os may be a result of ethyl amine spacer groups in POs-EA films, which may loosen the cross-linked structure.

**Steady-State Amperometric Glucose Response of Cross-linked PVIn-Os Films Containing GOX.** The steady-state electrooxidation current was measured at 1000 RPM as a function of the amount of enzyme in the films at 48 mM glucose concentration, well above the $K_m$ of the electrodes (figure 3). In electrodes prepared with a fixed amount of 10 µg PVIn-Os (n = 3, 5, 10), 11% peg 400 (by weight), and 0.5 µg to 60µg GOX, the current densities did not vary with the osmium content of the polymers. They were highest in electrodes containing 8 µg GOX, corresponding to 39 weight %. In a series of electrodes made with 2% to 22% peg 400, the highest current densities were observed in electrodes made with 11% of the cross-linker.

**Glucose Diffusion Through Cross-linked Films.** Increasing the rotation rate from 100 RPM to 2500 RPM at 10 mM and 48 mM glucose concentrations did not substantially affect the glucose electrooxidation currents. At 1 mM glucose concentration under a nitrogen atmosphere the current decreased only by 14% when the rotation rate was reduced from 2500 RPM to 100 RPM. As will be discussed later, this was not the case when the solution was O2 saturated. The absence of greater dependence of current density on rotation rate suggests
that the currents were controlled primarily by a process within the films and not by mass transport to the films at $\geq 1$ mM glucose concentration. Thus, from the measured 1.2 mA cm$^{-2}$ current density for benzoquinone, and assuming that glucose and benzoquinone permeation rates do not differ greatly, one can conclude that the current density is limited either by the rate of electron transfer to the redox polymer from the enzyme's FADH$_2$ centers, or by electron diffusion through the cross-linked polymer.

**O$_2$ Effects on Glucose Response.** The steady-state glucose response is shown in figure 4 for a typical cross-linked electrode (10 $\mu$g PVI$_3$-Os, 8 $\mu$g GOX, and 2.5 $\mu$g peg 400) under N$_2$, air, and O$_2$ at 1000 RPM. Evidently, O$_2$ competes effectively with PVI$_3$-Os, in the oxidation of FADH$_2$ centers.

At 48 mM glucose concentration, the glucose electrooxidation current decreased by 45% when the bubbled gas was switched from N$_2$ to O$_2$. At 2 mM glucose, the decrease was 76%. Apparently, at high glucose concentrations the O$_2$ flux is consumed in the outer layer of the film, while a substantial inbound glucose flux survives oxidation by O$_2$ and penetrates the film. Consistently, the loss in current upon switching the atmosphere from N$_2$ to O$_2$ was reduced not only at high glucose concentrations but also when thicker cross-linked films were employed or when the films were heavily loaded with GOX. Furthermore, upon decreasing the O$_2$ flux through stopping the 1000 RPM rotation of the electrodes, the O$_2$ associated loss diminished (figure 5). The glucose electrooxidation current at $\geq 6$ mM glucose concentration was greater when the electrode rotation was stopped because of the decrease in O$_2$ flux. Below 6 mM glucose the behavior was, however normal; i.e. the currents were higher for the rotating electrode, being dominated by glucose flux.

**Dependence of the Glucose Response on Film Thickness.** Electrodes with films having 1.1 X 10$^{-9}$ mol/cm$^2$ to 9.7 X 10$^{-8}$ mol/cm$^2$ electroactive osmium
were studied. A plot of the steady-state glucose electrooxidation current at 48 mM as a function of the surface density of the osmium centers is shown in figure 7. The current density increased with the surface density of osmium sites through a 100-fold range. The trend seen in figure 7 is similar to that observed for POs-EA.

pH and NaCl Concentration Dependence of the Glucose Electrooxidation Currents. The dependence of the current density on pH is shown in figure 8. As in the case of POs-EA the catalytic current peaked at and was nearly independent of pH from pH 7.5 to pH 10, in contrast with the GOX catalyzed oxidation of glucose by O2, where a narrow pH optimum near 5.5 has been observed. Current was lost irreversibly at pH > 10, while below pH 7.5 the changes in current with pH were reversible.

The dependence of the current density on NaCl concentration is shown in Figure 9 for a 10 mM phosphate buffer solution (pH = 7.2). The highest glucose current is observed at 50 mM NaCl with the current density decreasing as the NaCl concentration is increased, dropping to approximately half of its maximum value at 300 mM NaCl.

Selectivity of PVIn-Os Films in the Presence of Interferants. Glucose sensors are often insufficiently selective when operated at potentials where urate, ascorbate, and acetaminophen are also electrooxidized. The resulting errors can be reduced by membranes that preferentially transport glucose. Examples of such membranes are cellulose acetate and Nation. The effects of interferants can also be reduced by poising the electrodes at more negative potentials, where the rate of interferant electrooxidation is reduced. Since the redox potential of PVIn-Os is cathodic relative to that of POs-EA by 80 mV, the PVIn-Os electrodes are more selective. When required, glucose selectivity can
be further improved by non-electrochemical catalytic pre-oxidation of the interferants by H$_2$O$_2$ in an outer film of horseradish peroxidase.$^{15}$

Figure 10 shows a glucose calibration curve for a thick film electrode (prepared with 560 $\mu$g cm$^{-2}$ PVI$_3$-Os, 450 $\mu$g cm$^{-2}$ GOX, 140 $\mu$g cm$^{-2}$ peg 400) in the absence and presence of 0.1 mM ascorbate. Although in the absence of glucose the ascorbate electrooxidation current was high, this current was small with > 2 mM glucose. The current was increased by the 0.1 mM ascorbate at 6 mM glucose by 9%, corresponding to a $+0.5$ mM error in the glucose reading. The error was reduced when the glucose concentration was increased. Thus, at 20 mM glucose, the error was only 2%.

Interferants can be electrooxidized at the electrode surface ($I_{\text{surf}}$) and at osmium redox sites ($I_{\text{Os}}$). In electrodes with thin permeable PVI$_3$-Os films interferants were oxidized both directly at the carbon surface and at Os$^{\text{III}}$ redox sites, whereas, in thick PVI$_3$-Os films they were oxidized predominantly at the Os$^{\text{III}}$ redox sites.

When the glucose electrodes were made with thick PVI$_3$-Os films the ascorbate and acetaminophen caused currents decreased. This reduction in interference is interpreted as resulting from a lesser flux of ascorbate to the carbon surface i.e. from a reduction in $I_{\text{surf}}$. In the bulk of the polymer, where there is competition between ascorbate and FADH$_2$ for Os$^{\text{III}}$ sites, $I_{\text{Os}}$ decreases upon increasing the glucose concentration. At 0 mM glucose, the concentration of Os$^{\text{III}}$ sites is the highest and $I_{\text{Os}}$ is largest. Upon increasing the glucose concentration, the FADH$_2$ centers compete with ascorbate for the Os$^{\text{III}}$ redox centers and $I_{\text{Os}}$ decreases. Note that if ascorbate rather than a FADH$_2$ center is oxidized by an Os$^{\text{III}}$ center, then the glucose electrooxidation current ($I_G$) is decreased, and the increase in $I_{\text{Os}}$ is offset by the decrease in $I_G$. 

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In thin film electrodes the situation is quite different. A glucose response curve is shown in figure 11 for an electrode with such a thin film (140 μg cm\(^{-2}\) PVl3-Os, 110 μg cm\(^{-2}\) GOX, 35 μg cm\(^{-2}\) peg 400) in the absence and in the presence of 0.1 mM ascorbate. The ascorbate-related current increment is large, causing a 100% error at 6 mM glucose. The error slightly decreases, however, at high glucose concentration. Evidently, in thin film electrodes \(I_{\text{surf}}\) is large at high convection. The oxidation of ascorbate is strongly rotation rate dependent, in contrast with the glucose electrooxidation rate, that is nearly independent of the rotation rate.

Thick film PVIn-Os based glucose electrodes were also selective against acetaminophen. The glucose response curve with 1 mM acetaminophen was similar to that shown for 0.1 mM ascorbate. The addition of urate caused, however, an initial increase in oxidation current, followed by a rapid decay in current output, showing that an intermediate in the electrooxidation of urate damaged at least one group of centers involved in relaying electrons from glucose to the electrode. Cyclic voltammetry confirmed urate electrooxidation by Os\(^{III}\) sites and a resulting loss in the surface density of these sites.

Conclusions

The polyvinylimidazole derived redox polymers PVIn-Os form with GOX, upon cross-linking with a water-soluble diepoxyde, hydrogels that are permeable to glucose and through which electrons diffuse. The polymer is simpler and easier to make than the earlier reported polyvinyl pyridine derived POs-EA, and does not require modification with primary amines for cross-linking with the diepoxyde at ambient temperature in an aqueous solution. Electrons diffuse through the gel through a chain flexing dependent mechanism, i.e. the rate of
electron diffusion is not controlled by hopping between neighboring redox sites within chains, but by collisions between segments of redox polymer chains. Amperometric glucose sensors made with thick PVI\textsubscript{n}-Os and GOX based gels are reasonably selective in their glucose response in the presence of the electrooxidizable interferants ascorbate and acetaminophen.

Acknowledgement

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Table I. Effects of the extent of cross-linking and redox site density on $D_{eCp^2}$.

Figure 1. Dependence of $D_{eCp^2}$ on the concentration of NaCl for electrodes coated with cross-linked PVI10-Os film with 9.1% peg 400.

Figure 2. Dependence of $D_{eCp^2}$ on the pH for electrodes coated with PVI10-Os with 9.1% peg 400.

Figure 3. Dependence of the limiting catalytic current density on the GOX weight fraction for PVI$_n$-Os [$n = 3$ (circles), 5 (squares), 10 (triangles)] cross-linked with 11% peg 400. 1000 RPM, 20 mM, pH 7.2, air.

Figure 4. Dependence of the steady-state current density on the glucose concentration for the same electrode under N$_2$ (circles), air (triangles), and O$_2$ (squares). Electrode coated with 10 $\mu$g PVI$_3$-Os, 8 $\mu$g GOX, and 2.5 $\mu$g peg 400. At 1000 RPM

Figure 5. Dependence of the steady-state current density on the glucose concentration under N$_2$ (circles), air (triangles), and O$_2$ (squares). Electrode coated as in figure 4. Stagnant solution (no rotation).

Figure 6. Dependence of the steady-state current density on the glucose concentration under O$_2$ at 1000 RPM (squares) and under stagnant conditions (circles). Electrode coated as in figure 4.
Figure 7. Dependence of the limiting catalytic current density on the surface density of electroactive osmium sites. Conditions as in figure 3 with 48 mM glucose.

Figure 8. Dependence of the steady-state current density on pH. Conditions as in figure 3.

Figure 9. Dependence of the steady-state current density on NaCl concentration. Conditions as in figure 3.

Figure 10. Dependence of the steady-state current density on glucose concentration in the presence of 0.1 mM ascorbate (squares) and without ascorbate (circles). Electrode coated with 40 µg PVI3-Os, 32 µg GOX, and 10 µg peg 400.

Figure 11. Dependence of the steady-state current density on glucose concentration in the presence of 0.1 mM ascorbate (squares) and without ascorbate (circles). Electrode coated with 10 µg PVI3-Os, 8 µg GOX, and 2.5 µg peg 400.
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### Table I

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$D_e C_p^2 \times 10^{16}$ mol$^2$ cm$^{-4}$ s$^{-1}$

$D_e \times 10^8$ cm$^2$ s$^{-1}$
\( D_{eCp^2} \times 10^{16} \text{ mol}^2 \text{ s}^{-1} \text{ cm}^{-4} \)

**Figure 1**
$D_\theta C_p^2 \times 10^{16} \text{ mol}^2 \text{ s}^{-1} \text{ cm}^{-4}$
Figure 3
Current Ratio

NaCl Concentration (mM)

0.0 0.2 0.4
0 1.0
0.0
2000
1000

Current Ratio
Current Density (μa cm$^{-2}$)

Glucose Concentration (mM)