A Chemiresistor Sensor Based on Azo-Polymer and Graphene for Real-Time Monitoring of Mitochondrial Oxygen Consumption

André Olean-Oliveira,† Tiago Olean-Oliveira,‡ Ana C. R. Moreno,‡ Patricia M. Seraphim,‡ and Marcos F. S. Teixeira*†

†Department of Chemistry and Biochemistry and ‡Department of Physiotherapy, School of Science and Technology, Sao Paulo State University (UNESP), Rua Roberto Simonsen, 305, CEP 19060-900 - Presidente Prudente, São Paulo, Brazil

ABSTRACT: In the present study, a chemiresistor sensor based on a poly(Bismarck Brown Y)-reduced graphene oxide nanocomposite was developed to analyze the respiratory capacity of the constituent complexes of the electron transport chain. The sensorial platform was characterized using electrochemical impedance spectroscopy, and oxygen detection was accomplished by measuring the resistive properties of the sensor at fixed AC frequency. The impedance decreased significantly in response to small variations of the O2 concentrations tested up to saturation of the electrolyte solution with molecular oxygen. The resistive response of the sensor at 0.1 Hz was linear over the oxygen concentration range from 1.17 × 10⁻⁵ mol L⁻¹ to 1.02 × 10⁻³ mol L⁻¹, with a detection limit of 3.60 × 10⁻⁷ mol L⁻¹. Using the new O2 sensing platform, we monitored gradients in static cultures of adherent cells exposed to graded oxygen both at rest and upon metabolic stimulation. Under high dissolved oxygen conditions, the respiration of resting cells dictated that local O2 was moderately reduced, while cell metabolic stimulation triggered a major redistribution of O2. The usefulness of the developed sensor was demonstrated by continuous monitoring of mitochondrial oxygen consumption in various biologic applications.

KEYWORDS: resistive sensor, electrochemical impedance, conducting polymer, biomonitoring, mitochondrial respiratory chain activity

Mitochondria are organelles found in eukaryotic cells, and their metabolic role is crucial for cellular functioning. The main cellular function of mitochondria is energy production (ATP), but they also act on thermogenesis, steroid synthesis, and apoptosis. The regulation of energy metabolism by mitochondria occurs through the production of ATP by the oxidation of organic substrates in the Krebs cycle, and most of the energy produced comes from the electron transport chain. Each mitochondrial electron complex has a specific role so that, overall, the process results in the production of ATP through the consumption of molecular oxygen. Energy production is directly related to cellular oxygen concentration. Mitochondrial oxygen insufficiency is indicative of structural and functional alteration of cells, tissues, and organs facing pathological conditions. Furthermore, additional diseases have been investigated and found to be related to changes in the constituent complexes of the electron transport chain. Currently, there are several techniques for the study and investigation of mitochondrial performance. Clark-type electrodes have been utilized to monitor oxygen consumption in tissues and cells. Among the classes of electrochemical sensors, chemiresistor materials are the sensorial platform for chemical detection. The response mechanism of these sensors is the change of electrical resistance based on the resonant quantum tunneling effect of the sensor material in response to a change in the chemical environment. Different materials have been used as the sensorial platform in chemiresistors, such as polymers, inorganic, and organic semiconductors, and carbonaceous. The combination of two or more materials can
generate composites with new physicochemical properties taking advantage of the best properties and characteristics of each.\textsuperscript{13} Polymer−graphene nanocomposites prepared by combining the π-conjugated system of polymers with the various properties of graphene have been highlighted.\textsuperscript{14} In the case of azo-polymers, in addition to the advantages of being a conductive polymer, they still have the advantage of the presence of redox active sites due to the inclusion of diazonium groups in the polymer.\textsuperscript{15}

This work presents the first report of a mitochondrial oxygen chemiresistor sensor. The sensor response was detected by measuring the electrical conductivity change of the graphene polymer by electrochemical impedance spectroscopy. Furthermore, the effects of the applied potential, applied AC frequency, and dissolved oxygen concentration on the sensor response and efficiency were examined. The use of the proposed platform for real-time monitoring of mitochondrial oxygen in skeletal muscle cells exposed to graded oxygen both at rest and upon metabolic stimulation have been investigated in this work.

\section*{EXPERIMENTAL SECTION}

\textbf{Material and Methods.} All experiments were conducted in a conventional electrochemical cell containing three electrodes: a saturated calomel electrode (SCE) as the reference electrode, a platinum wire as the counter electrode, and a fluoro-doped tin oxide (FTO) electrode coated with a poly(azo-BBY)-reduced graphene oxide (rGO) as the working electrode (1.0 cm\textsuperscript{2}). The electrodes were connected to a PalmSens3 potentiostat and managed by PSTrace 5.4 software. All reagents used were of high purity ≥98\% (Sigma-Aldrich).

\textbf{Preparation of the Chemiresistor Oxygen Sensor.} Before coating, the FTO electrode was cleaned by immersion in an alkaline solution of 6:1 \( \text{H}_2\text{O} : \text{H}_2\text{O}_2 \) (30\% v/v), and \( \text{NH}_4\text{OH} \) (conc.) for 5
Table 1. Parameters Determined from Fitting of the Electrochemical Impedance Spectra in Figure 1A and B, in the Absence and Presence of O₂ (1.28 mmol L⁻¹), Using Electrodes Coated with rGO, poly(azo-BBY), or poly(azo-BBY)-rGO.

<table>
<thead>
<tr>
<th>Interface</th>
<th>$R_{ct}$</th>
<th>$R'_{ct}$</th>
<th>CPE$\alpha$</th>
<th>$\alpha_{dl}$</th>
<th>CPE$\alpha_{film}$</th>
<th>$\alpha_{film}$</th>
<th>Equivalent circuit</th>
</tr>
</thead>
<tbody>
<tr>
<td>rGO/FTO</td>
<td>N₁</td>
<td>92.4</td>
<td>27.3</td>
<td>0.9</td>
<td>500</td>
<td>0.7</td>
<td></td>
</tr>
<tr>
<td></td>
<td>O₁</td>
<td>17.0</td>
<td>28.7</td>
<td>0.9</td>
<td>500</td>
<td>0.7</td>
<td></td>
</tr>
<tr>
<td>pBBY/FTO</td>
<td>N₁</td>
<td>48.4</td>
<td>23.9</td>
<td>0.9</td>
<td>18.3</td>
<td>0.4</td>
<td></td>
</tr>
<tr>
<td></td>
<td>O₁</td>
<td>12.8</td>
<td>23.2</td>
<td>0.9</td>
<td>496</td>
<td>0.4</td>
<td></td>
</tr>
<tr>
<td>pBBY-rGO/FTO</td>
<td>N₁</td>
<td>0.08</td>
<td>49.6</td>
<td>93.7</td>
<td>0.9</td>
<td>486</td>
<td></td>
</tr>
<tr>
<td></td>
<td>O₁</td>
<td>3.05</td>
<td>25.5</td>
<td>22.8</td>
<td>0.9</td>
<td>312</td>
<td></td>
</tr>
</tbody>
</table>

$F_{applied} = -0.30$ V vs. SCE. $R_0 = 43$ Ω cm$^2$. The fitting error is ≤2%.

RESULTS AND DISCUSSION

Impedance Study of the Poly(azo-BBY)-rGO Nanocomposite Film. The development of the chemiresistor sensor based on poly(azo-BBY)-rGO nanocomposite was designed to include the desired properties of conductive azopolymers and graphene. Azo-polymers exhibit high sensitivity to molecular oxygen, which selectively reacts through the diazo active sites present in its polymeric structure.15,18 Moreover, azo-polymers exhibit extensive π-conjugation in which small perturbations at specific points of the polymer chain are reflected in the electronic properties of the material.19 These disturbances, in the case of the poly(azo-BBY), occur through the “off” and “on” possibilities of the diazo active sites in the π-conjugated polymer chain system (Figure S1). Graphene was introduced into the material to speed the rate of the electronic response, as well as to strengthen the chemical and mechanical resistance.20

The electrical resistivity properties of electrodes coated with rGO, poly(azo-BBY), and poly(azo-BBY)-rGO at an applied −0.30 V (vs SCE) were investigated by electrochemical impedance spectroscopy (EIS) in 0.05 mol L⁻¹ PBS (pH 7.4) in both the absence and presence of O₂, as represented in Figure 1A and B, respectively. This applied potential was chosen because it is coincident with reduction of the azo groups of the polymer by cyclic voltammogram (Figure S2). Table 1 shows the impedance parameters obtained according to the electrical circuit described for each platform (inside table). The plot of phase vs frequency was used to guide the proposal of equivalent circuit models for each studied platform (Figures 1C and D).

In general, the equivalent circuit models assigned to the platforms exhibited two constant phase elements (CPE), corresponding to the double electric (CPE$\alpha$) and deposited film (CPE$\alpha_{film}$) layers, with one charge transfer resistance ($R_{ct}$). The rGO film exhibited a high value of CPE$\alpha_{film}$ with a phase...
angle close to 90° (see Figure 1C), which agrees with previous studies reporting the charge storage capacity of graphene in the development of supercapacitors.21 The poly(azo-BBY)-rGO film also displayed a high value of CPE compared to the electrode coated with poly(azo-BBY)-rGO in the presence of dissolved oxygen (Table 1). In the case of the electrode coated with poly(azo-BBY)-rGO, the presence of oxygen in moderate led to a decrease in the impedance value, with the film resistance decreasing from 49.6 to 2.55 kΩ cm², indicating the smaller interfacial electron transfer resistance. This indicates the existence of the synergetic effect between graphene and poly(azo-BBY), where the space and gaps between graphene moieties were filled with polymer in the direct network formation.22

Comparing spectra in saturated oxygen solution, as in Figure 1B, differences are most noticeable in the radius of the semicircle arc. In all studied electrodes, the film resistance values were significantly lower than the values obtained in the absence of dissolved oxygen (Table 1). In the case of the electrode coated with poly(azo-BBY)-rGO film, the presence of oxygen in moderation led to a decrease in the impedance value, with the film resistance decreasing from 49.6 to 2.55 kΩ cm². This decrease clearly indicates the contribution of the reaction between polymer and oxygen.

To verify the greater sensitivity of the electrode coated with poly(azo-BBY)-rGO to that of graphene and polymer only, the real impedance variation as a function of frequency was studied. Figure 2 shows the real impedance values for the rGO, poly(azo-BBY), and poly(azo-BBY)-rGO films in the presence of solutions saturated with N2 and O2.

When compared with imaginary impedance, real impedance more directly contributed to the values associated with charge transfer resistance. The rGO, poly(azo-BBY), and poly(azo-BBY)-rGO films exhibited decreasing Zreal values at 0.1 Hz in the solution containing dissolved O2, as observed previously. However, the greatest difference in impedance response (∆Zreal = 22.2 kΩ cm²) was attributed to the electrode coated with poly(azo-BBY)-rGO (Figure 2C). The decrease in the resistance of poly(azo-BBY)-rGO nanocomposite was approximately 60% and 64% in comparison to rGO and poly(azo-BBY), respectively. The poly(azo-BBY)-rGO material provided real impedance change over the frequency range from 0.1 to 1 Hz. As observed, the change in real impedance is visible only at low frequencies, whereas the phase shift is more distinctive—especially in the presence of dissolved oxygen (see Figure 1D).

The decrease of the real impedance is mainly due to the decreasing charge transfer resistance (Rct) of the poly(azo-BBY)-rGO as oxygen concentration increases. The Rct results from the reaction between molecular oxygen and hydrazobenzene groups present in the polymer.15 The reaction generates the "on" \( \pi \)-conjugated system in the polymer, increasing the electronic conductivity between the conjugated network of polymer and graphene.23

**Optimization of Measurement Parameters for the Oxygen Chemiresistor Sensor.** Because the impedance parameters of the poly(azo-BBY)-rGO material depend on the O2 concentration of the solution, a precise determination of the applied frequency is very important in the context of possible applications for sensing small amounts of oxygen. Figure 3A shows the real impedance response of the sensor as a function of the dissolved oxygen concentration (0.17–1.03 mmol L⁻¹). The maximum changes in the impedance are observed at frequencies of less than 0.5 Hz, which is the region that most reflects the charge resistance. It is evident that the resistance tended to reach a minimum when O2 concentration in the solution reached saturation. We have previously seen that the impedance of the poly(azo-BBY)-rGO is dependent on the applied frequency range. As illustrated in Figure 3B, the response range of the sensor decreased gradually with increasing applied frequency. The sensitivity of the sensor (see Figure 3C) extracted from the calibration curve was also dependent on applied frequency. The applied frequency of 0.1 Hz was chosen as a better operational parameter for the next study step.

The use of the chrono-impedance technique for real time monitoring of dissolved oxygen variation in phosphate buffer (pH 7.4) was evaluated. The chrono-impedance response to small variations of oxygen concentration was achieved by first saturating the electrolyte solution with O2, then consuming small amounts of O2 by injecting aliquots of sulfite. A typical response of the sensor, recorded at different O2 concentrations, is shown in Figure 4A. The appreciable change in the impedance response reflects the possibility of sensor application in monitoring mitochondrial oxygen consumption.
Figure 4B shows a linear relationship between the impedance magnitude and dissolved oxygen concentration in the range of 0.0117 mmol L\(^{-1}\) to 1.02 mmol L\(^{-1}\):

\[
\text{Impedence (k}\ \Omega\text{)} = (28.4 \pm 0.3) - (16.8 \pm 0.4)[O_2] \text{ (mmol L}^{-1}\text{)}
\]

with a detection limit of 0.360 \(\mu\text{mol L}^{-1}\).

The dynamic impedance response of the poly(azo-BBY)-rGO sensor (Figure S3) was examined in PBS solutions, saturated with \(O_2\) and \(N_2\), with an applied frequency of 0.1 Hz. The sensor exhibited good reversibility and reproducibility with a relative variation of 4.1\% in the overall response of the chemiresistor per regeneration cycle. The response time to dissolved oxygen was quite rapid, with an apparent response time of \(\pm 3\) min. This time meets expectations for the applied purpose of the chemiresistor sensor in biological systems, since similar measurements require several minutes to stabilize after each mitochondrial stimulus-response.\(^{16}\)

Chemiresistor Sensor Applications in Real-Time Monitoring of Mitochondrial Oxygen. To address the application of the sensor on mitochondrial respiration, mitochondrial \(O_2\) consumption corresponding to the response of each electron transport chain complex was measured during a sequential substrate addition protocol.\(^{16,24}\) The competition for these substrates (pyruvate, ADP, succinate, and cytochrome c) was ascertained by the direct response of sensor as shown by chrono-impedance measurements of the dissolved \(O_2\) (1.00 mmol L\(^{-1}\)). Figure S4 shows the interference study in the impedance response of the sensor. The analyzed substrates were not considered to interfere when the relative response in the measured impedance was less than 5\%.

The performance of the sensor in real-time monitoring of mitochondrial oxygen consumption was assessed in tissue samples collected from the skeletal muscle from 4 different groups of rats: control (C), smokers (F), smokers exercised (FE), and exercised (E). Mitochondrial respiration can be evaluated, in general, according to the respiration type. There are three main types of respiration:\(^{24,25}\) (1) the ROUTINE (R) respiration state is the demand for cellular oxygen at steady-state without any additions to the cell medium; (2) the LEAK (L) respiration state is an acronym for the resting oxygen flux compensating for proton leak, proton slip, and cation cycling and is related to the generation of reactive oxidative species (ROS); (3) the OXPHOS (P) respiration state is the mitochondrial respiratory capacity to produce ATP. Figure 5 illustrates real-time measurements of cellular oxygen consumption by studied tissue in respiratory buffer using the proposed sensor. The decrease in the oxygen concentration (red line) was detected by the sensor for each injection of tissue (R) and pyruvate (L). The addition of ADP (P) and succinate (P) substantially increased oxygen consumption by activating mitochondrial oxidative phosphorylation. This
indicated that respiration was substrate dependent. Cytochrome c was added at the end of the measurements to verify mitochondrial integrity, and no significant effect was detected across all samples. The oxygen consumption rates (Table 2) were calculated for each substrate injected into the system in the four test groups, according to the literature descriptions.6b,26

The evaluation of the functional performance of mitochondria is not the main focus of this manuscript, but the results obtained may indicate applicability of the proposed sensor. These results clearly show the ability of the proposed sensor to monitor the oxygen consumption with respiratory chain substrates confirms the maintenance of an intact and functional respiratory chain when tissues are suspended in the respiratory buffer.

**CONCLUSION**

The chemiresistor platform based on the poly(azo-BBY)-rGO nanocomposite film was demonstrated to be a simple, sensitive, and selective sensor for O2 detection in mitochondrial respiration. The application of the proposed sensor for the real-time monitoring of mitochondrial oxygen consumption during a sequential substrate addition protocol was consistent with similar techniques in the biological area.

**ASSOCIATED CONTENT**

* Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acssensors.8b01013.

Schematic representation of the conducting azo-polymer; electrochemical response of the azo-polymer; transient responses of the oxygen sensor and the selectivity of the sensor (PDF)

**AUTHOR INFORMATION**

**Corresponding Author**

E-mail: marcos.fs.teixeira@unesp.br. Tel.: +55 18 3229 57 49. Fax: + 55 18 3221 56 82.

**ORCID**

Marcos F. S. Teixeira: 0000-0001-9355-2143

**Notes**

The authors declare no competing financial interest.

**ACKNOWLEDGMENTS**

The authors acknowledge FAPESP (2016/09017-1 and 13/07296-2) for financial support. A.O.O. thanks CAPES-Brazil for a master fellowship. GADU and NSA.

**REFERENCES**


---

**Table 2. Mitochondrial Oxygen Consumption**

<table>
<thead>
<tr>
<th>Test group</th>
<th>ROUTINE</th>
<th>LEAK</th>
<th>OXPHOS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>17.2 ± 2.1</td>
<td>25.4 ± 1.0</td>
<td>132.2 ± 3.9</td>
</tr>
<tr>
<td>Exercised</td>
<td>74.2 ± 5.7</td>
<td>25.7 ± 2.1</td>
<td>165.3 ± 4.1</td>
</tr>
<tr>
<td>Smoker</td>
<td>10.0 ± 2.3</td>
<td>28.3 ± 1.9</td>
<td>143.0 ± 4.4</td>
</tr>
<tr>
<td>Smoker/Exercised</td>
<td>19.3 ± 3.1</td>
<td>27.8 ± 1.8</td>
<td>104.0 ± 6.2</td>
</tr>
</tbody>
</table>

"The rate of oxygen consumption in different states of mitochondrial respiration. Values represent the mean for a set of three animals."
(9) (a) Koudehi, M. F.; Pourmortazavi, S. M. Polyanil Vinyl Alcohol/ Polypyrrole/Molecularly Imprinted Polymer Nanocomposite as Highly Selective Chemiresistor Sensor for 2,4-DNT Vapor Recognition. *Electroanalysis* 2018, 30 (10), 2302−2310. (b) Alizadeh, T.; Rezaloo, F. Toluene chemiresistor sensor based on nano-porous toluene-imprinted polymer. *Int. J. Environ. Anal. Chem.* 2013, 93 (8), 919−934.


