Enhanced Performance of GCE/N-Reduced Graphene Oxide-Au Nanocomposite in Dopamine Sensing †

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Abstract: The reduced graphene oxide (rGO) and nitrogen-reduced graphene oxide (N-rGO) were electrochemically coated with gold nanoparticles and compared as the modifier of a glassy carbon electrode (GCE) for dopamine (DA) electrochemical sensors. The rGO and N-rGO were prepared by reduction of graphene oxide (GO) under hydrothermal conditions. Urea was applied as a nitrogen dopant. The modified GCEs were tested as working electrodes in the electrochemical DA detection in the presence of ascorbic and uric acids. Initially, the working parameters of sensors in a pure dopamine solution were determined. The GCE/N-rGO-Au electrode exhibited a lower limit of detection compared with the GCE/rGO-Au (385 vs. 700 nM). Both sensors had a wide linear range of 1-100 µM and high sensitivity of 0.78 µA/µM for GCE/N-rGO-Au and 1.78 µA/µM for GCE/rGO-Au. The presence of ascorbic and uric acids in the solution resulted in a decrease in the intensity of DA oxidation peak, maintaining acceptable limit of detection. This study showed that the surface modification of the graphene materials with gold nanoparticles allows to obtain satisfactory working parameters of DA sensors. The GCE/N-rGO-Au demonstrated the superior performance in DA sensing due to a homogeneous distribution of gold nanoparticles on the surface of the N-doped graphene material.

Keywords: dopamine; gold nanoparticles; reduced graphene oxide; nitrogen-doped reduced graphene oxide; electrochemical sensor

1. Introduction

Nowadays, great scientific efforts are put into the development of electrode materials in order to obtain sensitive, fast and effective dopamine (DA) detection [1]. Reduced graphene oxide (rGO) and N-doped reduced graphene oxide (N-rGO) are promising materials in the electrochemical DA detection due to their high electrical conductivity and a high surface to volume ratio [2,3]. The introduction of nitrogen atoms into the structure of rGO leads to elimination of oxygen groups and enhancement of electrical conductivity [3]. The main problem of the DA detection is that DA coexists with uric and ascorbic acids in biological samples [4]. Voltammetric responses of these compounds are similar, therefore it is important to use materials which ensure selective DA detection in the presence of different interferents. Increasingly popular, the modification of graphene materials by noble metal nanoparticles, such as gold nanoparticles (AuNPs), results in an enhancement of DA sensing and its selectivity [5]. AuNPs can catalyze red-ox reactions and enhance electron transfer [5]. Deposition of AuNPs on the surface of carbon materials can increase the active surface and enhance the affinity between the electrode material and the analyte such as DA.

In this work, we demonstrate the influence of rGO and N-rGO modified with AuNPs on the electrochemical DA detection. The synthesized graphene materials exhibited different nanoparticles distribution and size ranges on their surfaces. rGO and N-rGO were used for modification of a glassy carbon electrode (GCE) for dopamine (DA) electrochemical sensors. The modified GCEs were tested as working electrodes in the electrochemical DA detection in the presence of ascorbic and uric acids. Initially, the working parameters of sensors in a pure dopamine solution were determined. The GCE/N-rGO-Au electrode exhibited a lower limit of detection compared with the GCE/rGO-Au (385 vs. 700 nM). Both sensors had a wide linear range of 1-100 µM and high sensitivity of 0.78 µA/µM for GCE/N-rGO-Au and 1.78 µA/µM for GCE/rGO-Au. The presence of ascorbic and uric acids in the solution resulted in a decrease in the intensity of DA oxidation peak, maintaining acceptable limit of detection. This study showed that the surface modification of the graphene materials with gold nanoparticles allows to obtain satisfactory working parameters of DA sensors. The GCE/N-rGO-Au demonstrated the superior performance in DA sensing due to a homogeneous distribution of gold nanoparticles on the surface of the N-doped graphene material.
carbon electrode (GCE), which was applied as a working electrode in the DA detection. GCE/N-rGO-Au exhibited a lower limit of DA detection when compared to GCE/rGO-Au (385 vs 700 nM).

2. Materials and Methods

GO was synthesized by the modified Hummers method from the graphite (TimCal). rGO and N-rGO were obtained by treatment of 100 ml of GO in an autoclave under hydrothermal conditions at 180 °C for 5 h (rGO) and 12 h in the presence of urea (N-rGO) [6]. A dispersion of the graphene material (4 mg) in a mixture of 1 mL Mili-Q water and DMF as a dispersion stabilizer (1:1 v/v) was used to modify the GCE by drop-casting. After that, the AuNPs were electrodeposited on the electrode surface from 1 mM HAuCl4 aqueous solution using cyclic voltammetry (CV).

The electrochemical measurements were performed in a three electrode cell using GCE/(N)-rGO-Au as a working electrode, a platinum wire as a counter and a saturated calomel electrode as a reference electrode, respectively. The measurements were carried out in a phosphate buffer solution (PBS, pH 7.4) containing 100 µM DA as an electrolyte. After each measurement the working electrodes were purified by CV in 7.4 PBS buffer solution to remove DA.

3. Results and Discussion

The DA oxidation-reduction reaction was investigated by CV and differential pulse voltammetry (DPV) which enable to determine the working parameters of the DA sensors (Table 1). As shown on Figure 1, the modification of GCE with N-rGO and N-rGO/Au results in an enhancement of the DA oxidation peak. The oxidation potential for both electrodes was comparable (Table 1). GCE/N-rGO-Au exhibits a lower detection limit and sensitivity when compared with GCE/rGO-Au (385 vs. 700 nM and 0.78 vs. 1.78 µA/µM, respectively).

![Figure 1](image-url) CVs of the examined electrodes in 0.1 M PBS containing 100 µM of DA.

<table>
<thead>
<tr>
<th>Working Electrode</th>
<th>LOD, nM</th>
<th>Linear Range, µM</th>
<th>Sensitivity, µA/µM</th>
<th>Oxidation Potential, mV</th>
</tr>
</thead>
<tbody>
<tr>
<td>GCE/N-rGO-Au</td>
<td>385</td>
<td>1-100</td>
<td>0.78</td>
<td>210</td>
</tr>
<tr>
<td>GCE/rGO-Au</td>
<td>700</td>
<td>1-100</td>
<td>1.78</td>
<td>190</td>
</tr>
<tr>
<td>GCE</td>
<td>5000</td>
<td>10-100</td>
<td>0.12</td>
<td>190</td>
</tr>
</tbody>
</table>

The morphology of the graphene materials was examined by the field emission-scanning electron microscopy (FESEM) (Figure 2). N-rGO-Au exhibited a high concentration and homogenous distribution of AuNPs on its entire surface, despite fewer cycles of electrodeposition than for rGO (20 vs. 30 cycles). Moreover, the Au nanoparticles deposited on the N-rGO were smaller than those of rGO-Au (20-260 nm vs. 30-450 nm). This suggests that AuNPs are preferentially deposited on the N-rich regions of the graphene material.
To determine the surface elemental composition of nanocomposites, the X-ray photoelectron spectroscopy (XPS) was applied. It was revealed that a 7.6 % at. of Au was electrodeposited on the surface of N-rGO, which was followed by a decrease in the nitrogen content from 6.4 % at. for N-rGO to 2.7 % at. for N-rGO-Au. The lower nitrogen amount on the surface of N-rGO proves that AuNPs anchor on the N-rich regions covering nitrogen atoms, which leads to a decrease of the nitrogen content on the surface of N-rGO-Au determined by XPS.

4. Conclusions

This study shows that N-rGO is a promising graphene material to obtain Au-containing nanocomposites with a homogeneous distribution of AuNPs, which result in enhanced working parameters of DA sensors. Despite a lower number of cycles of the electrodeposition, the efficiency of the AuNP's deposition on the graphene nanosheets was higher for N-rGO than for rGO.

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Conflicts of Interest: The authors declare no conflict of interest.

References


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