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# Preparation of graphene/Au aerogel film through the hydrothermal process and application for H<sub>2</sub>O<sub>2</sub> detection

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In this paper, one-step preparation of graphene/gold nanoparticle hydrogel film through the hydrothermal method is reported. The hydrogel film could be formed on a glass substrate under hydrothermal conditions, and upon freeze-drying, the aerogel film of 40 μm thickness with satisfying flexibility and strength is obtained. The aerogel composite film is characterized by scanning/transmission electron microscopy, X-ray diffraction, Raman spectroscopy and X-ray photoelectron spectroscopy. Moreover, the aerogel film is directly used as the electrochemical electrode for sensing H<sub>2</sub>O<sub>2</sub>, and exhibits good performance with a broad linear range, low detection limit and excellent selectivity. This work provides a route for the fabrication of graphene film material with wide potential in various aspects.

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## Introduction

Graphene has gathered tremendous attention and in-depth research in various fields owing to its high surface area, excellent conductivity and high mechanical strength.<sup>1,2</sup> Multiple graphene-based materials have been prepared for potential applications, such as energy storage and conversion,<sup>3,4</sup> catalysis<sup>5</sup> and electrochemical sensing.<sup>6</sup> Electrochemical analysis is one of the most promising aspects of graphene's application, ascribed to the excellent electron transport property and high specific surface area. Graphene-involving non-enzyme electrochemical sensors have shown excellent performance in terms of sensitivity, durability and reproducibility.<sup>7,8</sup> In general, graphene-based electrochemical sensors are fabricated by the modification of a glassy carbon electrode (GCE) with the graphene suspension, while the drop-drying method would inevitably bring about the restacking of the graphene sheets and influence the properties of the sensors. Recently, with the rising research of graphene aerogels,<sup>9–12</sup> some efforts have been focused on the preparation of GCEs modified with graphene aerogels, which could utilize the superiority of the porous material and have exhibited dramatic electrochemical performance enhancement.<sup>13–15</sup> The fabrication processes also usually involve the deposition of the aerogel suspension on GCEs. However, owing to the good mechanical strength of the graphene aerogel, the manipulation is usually difficult, even requiring the aid of ultrasonication or grinding pre-treatment. Besides, some untraditional methods were also utilized to prepare graphene-aerogel modified GCEs. For example, Zhu *et al.* fabricated a hemispheric graphene/Ag aerogel on a GCE<sup>16</sup>

and Tian *et al.* covered the GCE with a phosphorus-doped graphene aerogel disk,<sup>17</sup> which both exhibited excellent performance in electrochemical sensing of hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>).

So far, most applications of graphene in electrochemical sensors are realized on GCEs, which involves a relatively tedious process of electrode pre-treatment and graphene dispersing/dropping/drying with accurate control and necessary consideration of adhesion and duration. One convenient alternative way is the direct use of graphene, especially 3D materials, as electrode sensors. For instance, Chen *et al.* prepared graphene foam by CVD method as an excellent electrochemical sensor for detection of dopamine.<sup>18</sup> Wang *et al.* reported the direct use of nitrogen-doped graphene aerogel piece cut from the hydrothermally prepared aerogel as high-performance non-enzymatic sensor.<sup>19</sup> We have reported the preparation of graphene/gold nanoparticle (GNP) aerogel under a mild condition as electrode for electrochemical sensing of H<sub>2</sub>O<sub>2</sub>.<sup>20</sup>

Hydrothermal process is a widely used and efficient method for converting the aqueous graphene oxide (GO) dispersion into reduced GO (rGO) hydrogel, which could subsequently yield rGO aerogel by freeze-drying treatment.<sup>21</sup> Under the hydrothermal condition, the rGO sheets are interconnected owing to the strong π–π, hydrophobic and electrostatic interactions.<sup>22</sup> The release of CO<sub>2</sub> in the hydrothermal process endows the rGO framework with porous structure.<sup>23</sup> In most cases, the as-prepared rGO aerogels are in the forms of 3D bulk or monolith.<sup>24</sup> Actually, the graphene aerogel films have numerous potential applications, such as electronic materials,<sup>25,26</sup> sensors,<sup>19,27</sup> batteries/supercapacitors,<sup>18,28–30</sup> and filter membranes.<sup>31,32</sup> However, the thin aerogel films, are generally fabricated by peeling or grinding-casting from the bulky aerogels, and the quality of the final film is uncontrollable.

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In this work, we report the facile preparation of graphene aerogel films through the hydrothermal method. In the autoclave, the mixed solution of GO, reductant and  $\text{HAuCl}_4$  is uniformly spreaded on a glass sheet and undergoes a hydrothermal process. The obtained rGO/GNP composite hydrogel is then freeze-dried to yield the rGO/GNP aerogel film, which could be cut and directly used as electrochemical electrode. The aerogel film sensor demonstrates satisfying electrocatalytic performances in the detection of  $\text{H}_2\text{O}_2$ .

## Experimental

The reagents used in the experiment are all analytically pure and used without further purification. The natural graphite powder was purchased from Alfa Aesar.  $\text{HAuCl}_4 \cdot 4\text{H}_2\text{O}$  and L-ascorbic acid (AA) were purchased from Sinopharm Chemical Reagent Beijing Co., Ltd. Other chemicals were purchased from Beijing Chemical Works. GO was prepared from graphite powder by a modified Hummers' method.<sup>33</sup> Briefly, 5 mL of aqueous GO solution ( $6 \text{ mg mL}^{-1}$ ,  $\text{pH} = 8.5$ , adjusted by  $\text{NaOH}$ ) was mixed with  $60 \mu\text{L}$  of aqueous solution of  $\text{Na}_2\text{B}_4\text{O}_7$  ( $5 \text{ mg mL}^{-1}$ ) as an enhancer<sup>34</sup> in a beaker.  $90 \text{ mg}$  of AA was added into the mixture with vigorous and continuous stirring. Afterwards,  $200 \mu\text{L}$  of aqueous  $\text{HAuCl}_4$  solution ( $14.34 \text{ mg mL}^{-1}$ ) was dropped into the solution. The composite aerogel was made by dropping and spreading  $500 \mu\text{L}$  of the mixed solution on a glass sheet ( $2 \text{ cm} \times 2.3 \text{ cm}$ ). Then the glass sheet was put into a hydrothermal autoclave ( $50 \text{ mL}$ ) containing  $1 \text{ mL}$  of deionized water before heated at  $180 \text{ }^\circ\text{C}$  for  $14 \text{ h}$ . Afterwards, the formed hydrogel film was freeze-dried under vacuum ( $-50 \text{ }^\circ\text{C}$ ) to obtain the resulting rGO/GNP aerogel film (thickness  $\sim 40 \mu\text{m}$ ).<sup>35</sup> Finally, the film was cut into pieces ( $2 \text{ mm} \times 3 \text{ mm}$ ) and connected with a conductive copper paste (wrapped tightly with insulating tape to prevent the contact with the solution) to make the electrode for electrochemical measurements. The preparation process of aerogel film and electrode is outlined in Scheme 1.

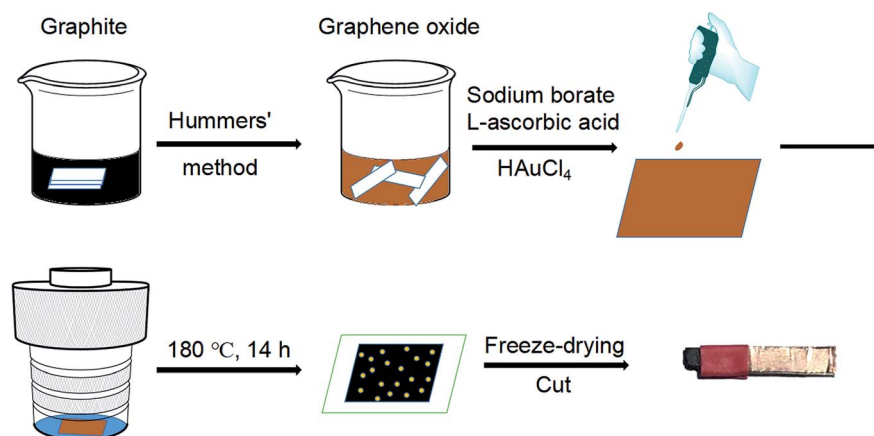
Scanning electron microscopic (SEM) characterization was conducted on a Hitachi SU8010 with the accelerating voltage at

$5 \text{ kV}$ . Transmission electron microscopic (TEM) images were captured *via* a JEOL 2010 operating with the accelerating voltage at  $200 \text{ kV}$ . A Gatan model 780 CCD camera is used to confirm the sizes and dispersions of nanoparticles. X-ray diffraction (XRD) analysis was examined by Shimadzu XRD-7000 with  $\text{Cu K}\alpha$  radiation in the  $2\theta$  range from  $5^\circ$  to  $80^\circ$ , which had a scan rate of  $2^\circ \text{ min}^{-1}$  at  $40 \text{ kV}$  and  $30 \text{ mA}$ . Raman spectra were obtained by a Horiba XploRA PLUS confocal microscope with  $532 \text{ nm}$  laser excitation. X-ray photoelectron spectroscopy (XPS) was investigated by a Kratos AXIS Ultra at  $300 \text{ W Al K}\alpha$  radiation. The electrochemical analysis was conducted on a CHI660E electrochemical analyzer (Chenhua instruments, Shanghai) using a conventional three-electrode system composed by a platinum wire as the counter electrode, a  $\text{KCl}$  saturated  $\text{Ag/AgCl}$  electrode as the reference electrode and the aerogel film electrode with copper paste as working electrode.

## Results and discussion

The preparation of the graphene hydrogel film refers to the traditional hydrothermal method except that small dose of GO solution ( $500 \mu\text{L}$ ) on the glass sheet replaces the relatively large amount of solution (dozens of  $\text{mL}$ ) in the autoclave. To create a hydrothermal condition and avoid the drying of GO solution, extra water is added around the glass sheet. During the hydrothermal process, the reduction of GO and  $\text{Au}^{3+}$  by AA occurs simultaneously to form the rGO/GNP hydrogel film on the glass. Just like the shrinkage of common graphene hydrogel, the area of the obtained rGO/GNP hydrogel film is about half of the precursor solution. Subsequent freeze-drying treatment produces the smooth aerogel composite film with the thickness of  $\sim 40 \mu\text{m}$ . The rGO/GNP aerogel film possesses a satisfying tensile stress of  $2.5 \text{ MPa}$ , which facilitates the following manipulation and ensures its stability in application.

The microstructure of the rGO/GNP aerogel film was investigated by SEM (Fig. 1a–c) and TEM (Fig. 1d). As seen from a rough review of the surface (Fig. 1a), the film demonstrates the smooth morphology with pores within the interconnected graphene sheets. In higher magnification of the side view (Fig. 1b),



Scheme 1 Illustration for the preparation of rGO/GNP aerogel film electrode.



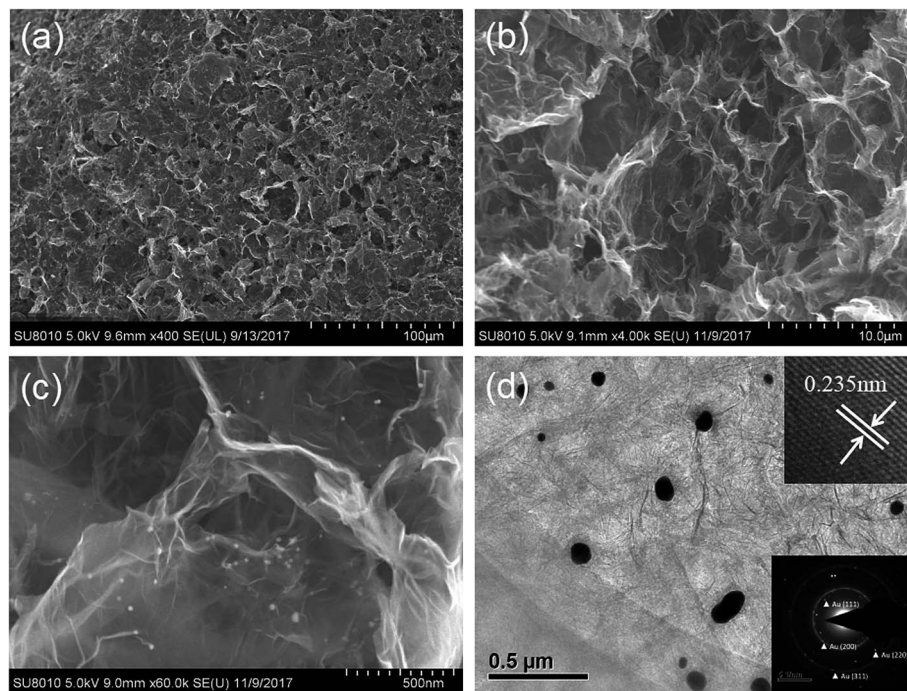


Fig. 1 SEM images of the rGO/GNP aerogel at (a) front view and (b and c) side view. (d) TEM image of the rGO/GNP aerogel with the insets showing the (111) planes and SAED pattern of Au.

the porous graphene structure could be obviously observed, while the GNPs could hardly be seen. Fig. 1c clearly displays the homogeneously distributed GNPs on the graphene sheet. The GNPs exhibit the relatively large size of  $\sim 40$  nm, as shown in TEM of Fig. 1d, with the inset exhibiting the (111) planes of crystalline gold observed in the high resolution. The selected area electron diffraction (SAED) shows typical reflexes of ordered regions of Au (Fig. 1d inset), which is consistent with the following XRD result.

The rGO/GNP aerogel film was further characterized by XRD and Raman measurements. As seen in Fig. 2a, the XRD patterns

of the aerogel film and pristine GO show distinct characteristics. Compared with the peak of GO sample at  $10^\circ$ , the graphene peak of rGO/GNP aerogel appears at  $25.3^\circ$ , indicating the stacking of graphene sheets owing to the reduction of GO to rGO. The peaks at  $38.35^\circ$ ,  $44.54^\circ$ ,  $64.81^\circ$  and  $77.71^\circ$  are assigned to face-centered cubic bulk Au (111), (200), (220) and (311), respectively, which match well with the standard values of Au (JCPDS 04-0784).<sup>36</sup> The Raman spectra (Fig. 2b) of rGO/GNP and GO samples display the characteristic peaks at  $1257\text{ cm}^{-1}$  and  $1500\text{ cm}^{-1}$ , which can be attributed to the D band and G band of graphene, respectively. The D band stands for the disordered

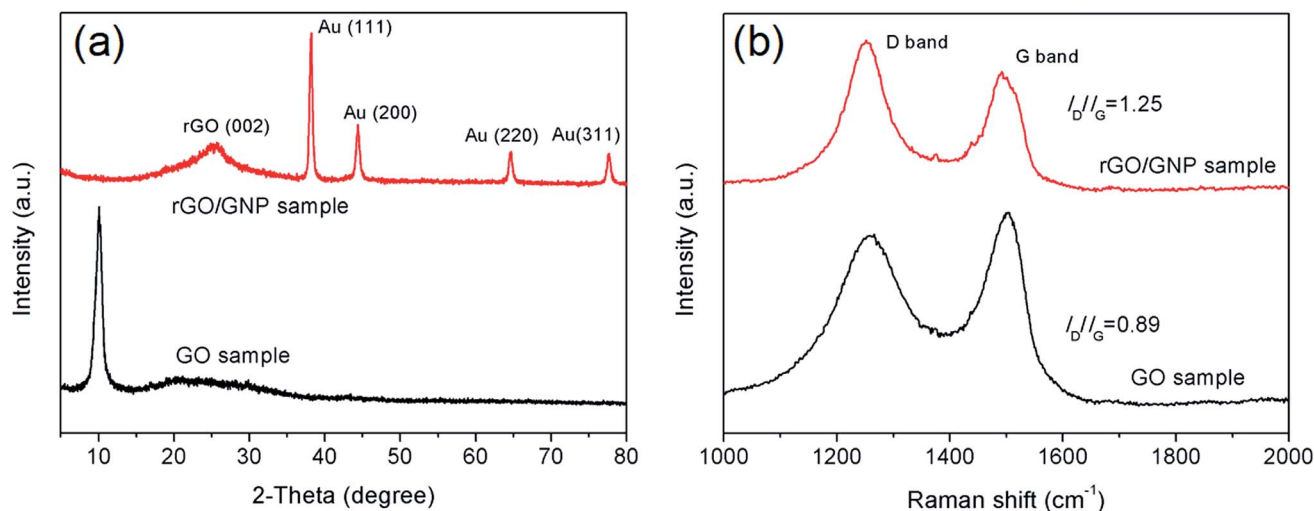


Fig. 2 (a) XRD patterns and (b) Raman spectra of rGO/GNP and GO samples.



structural defects and the G band represents the first-order scattering of the  $E_{2g}$  mode observed for  $sp^2$  carbon domains. The intensity ratio of two bands ( $I_D/I_G$ ) indicates the structural integrity of the graphene. Generally, the value of  $I_D/I_G$  would decrease after the reduction of GO due to removal of the oxygen-containing functional groups and the recovery of  $sp^3$  carbon. However, the  $I_D/I_G$  value of rGO/GNP aerogel increases to 1.25, much higher than 0.89 of GO sample. This phenomenon demonstrates the increasing of the disorder for the graphene carbon in the rGO/GNP sample, which could be ascribed to the breaking of graphene sheets into smaller pieces after reduction.<sup>35,37</sup>

The XPS spectra of rGO/GNP and GO samples clearly demonstrate the variation of the compositions. As shown in Fig. 3a, both of the samples exhibit the features of C1s and O1s at the binding energies of 284 eV and 532 eV, respectively. The spectrum of rGO/GNP sample exhibits the peaks of Na Auger at 497 eV corresponding to the residual byproduct. The peak of Au could be divided into two peaks corresponding to  $Au4f_{7/2}$  and  $Au4f_{5/2}$  in the metallic state with the binding energies of 84.2 eV

and 87.8 eV, respectively (Fig. 3b).<sup>38</sup> The deconvoluted C1s spectra for GO and rGO/GNP samples are shown in Fig. 3c and d, respectively. The peak of 284.5 eV is ascribed to the C=C bonds and the peaks at 286.6 eV, 285.6 eV and 287.9 eV are assigned to C-O-C, C-OH and O-C=O, respectively.<sup>39,40</sup> Obviously, the peak intensity associated with oxygen-containing bonds dramatically decreases for the rGO/GNP aerogel compared with that for GO, indicating that GO in the aerogel is almost completely reduced.

The aerogel film could be conveniently tailored owing to its good flexibility and strength. Herein the rGO/GNP film is utilized directly as the electrode for electrochemical analysis. The electrocatalytic performance of rGO/GNP electrode in detecting  $H_2O_2$  in phosphate buffer solution (PBS, pH = 7) are investigated. As shown in Fig. 4a, the cyclic voltammograms (CVs) exhibit obvious redox couple peaks upon 0.5 mM  $H_2O_2$ , and the currents increase with the increase of the scan rate. The inset reveals that there is a linear relationship between the reduction current and the square root of the scan rate, which suggests the diffusion-controlled electrochemical process for

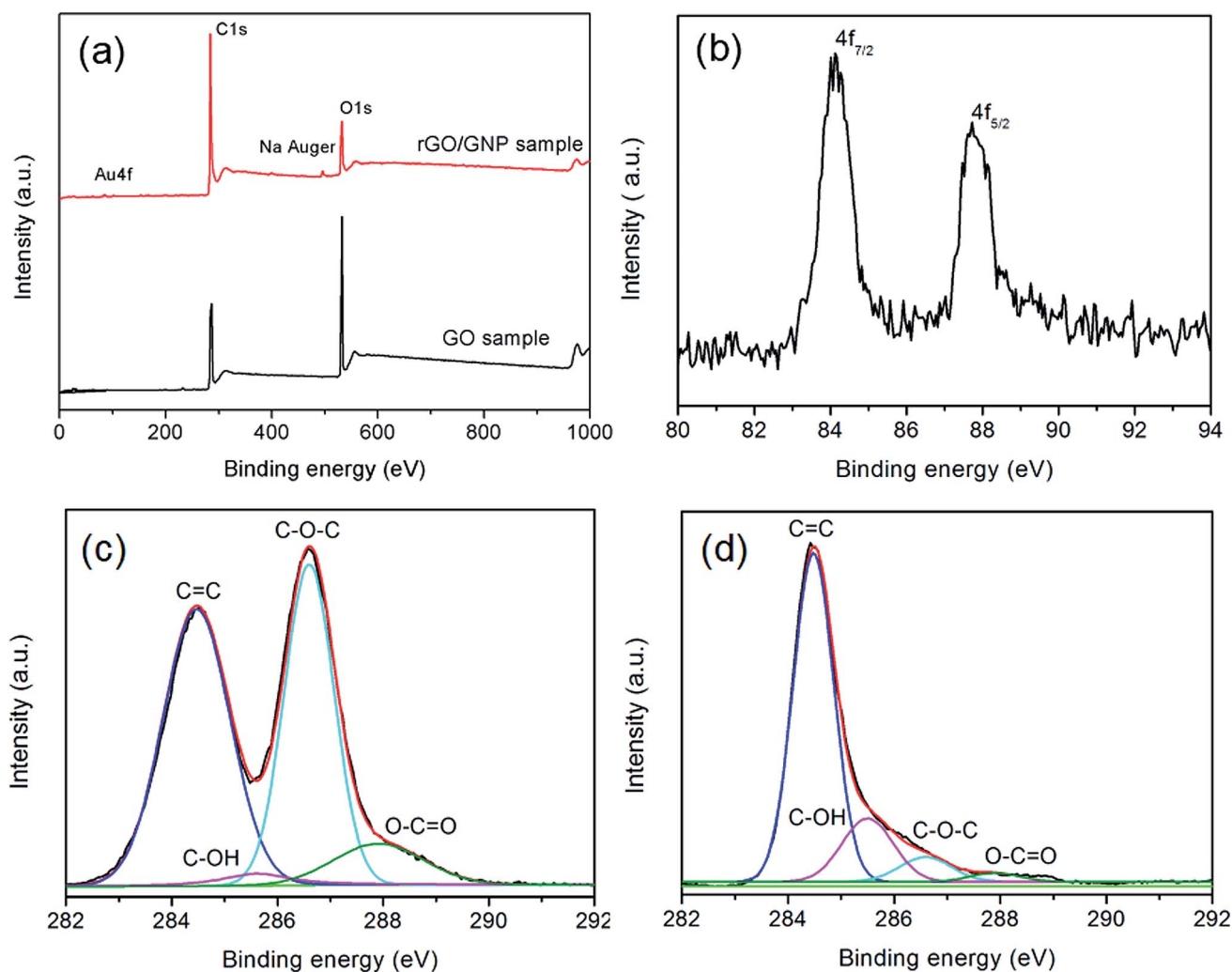


Fig. 3 (a) XPS spectra of rGO/GNP and GO samples and deconvoluted XPS peaks for (b) Au4f of GO/GNP sample, (c) C1s of GO sample and (d) C1s of rGO/GNP sample, with a correction of 0.3 eV for sample charging.



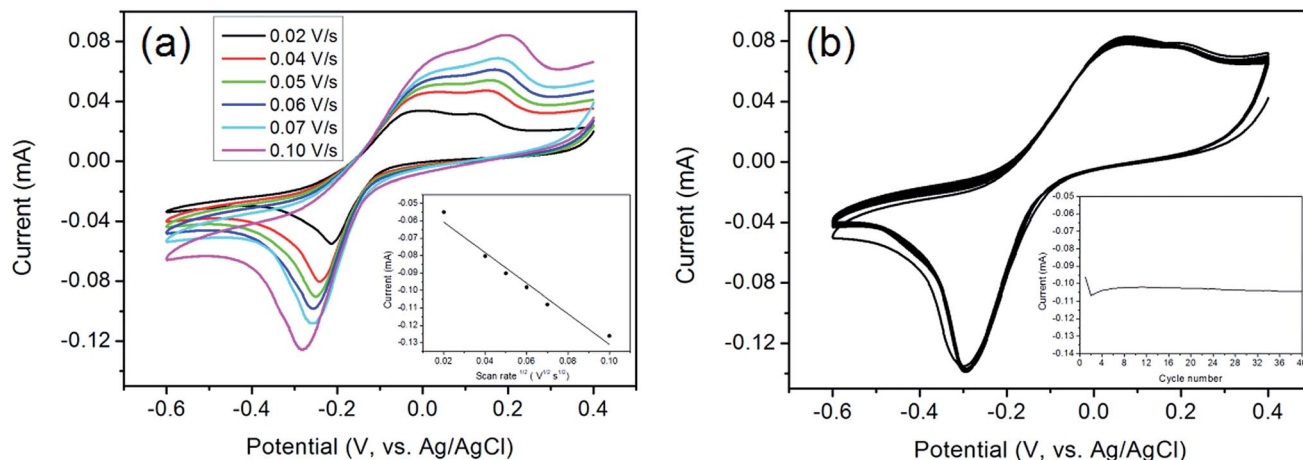


Fig. 4 CVs of rGO/GNP aerogel electrode in PBS solutions (pH = 7) with the presence of 0.5 mM  $\text{H}_2\text{O}_2$  (a) at different scan rates (0.02, 0.04, 0.05, 0.06, 0.07 and 0.10  $\text{V s}^{-1}$ ) with the inset showing the linear relationship of the response current and the square root of scan rates, and (b) at the scan rate of 0.10  $\text{V s}^{-1}$  for 40 cycles with the presence of 0.5 mM  $\text{H}_2\text{O}_2$  with the inset showing the current variation upon the cycle number.

the rGO/GNP aerogel electrode. In addition, to investigate the repeatability of rGO/GNP aerogel electrode, CV measurements are carried out at the scan rate of 0.1  $\text{V s}^{-1}$  with the presence of 0.5 mM  $\text{H}_2\text{O}_2$  for 40 cycles, as seen in Fig. 4b, and the inset shows the variation of the current along with the cycle number. Obviously, except the first two cycles for electrode activation, the current almost stays in a straight line with only 2% current loss at the end, which means the excellent repeatability of the rGO/GNP aerogel electrode.

The typical current–time ( $I-t$ ) plot of rGO/GNP electrode in PBS solution (pH = 7) on consecutive step change of  $\text{H}_2\text{O}_2$  concentrations is shown in Fig. 5a. The instant increase of the reductive current responding to the stepwise dropping of  $\text{H}_2\text{O}_2$  suggests the rapid amperometric response of the rGO/GNP electrode regarding  $\text{H}_2\text{O}_2$ . The inset demonstrates the calibration curve of the response current against the  $\text{H}_2\text{O}_2$  concentration, which contains a broad linear range of 0.4–6.4 mM ( $R^2$

= 0.992) and a low detection limit of 12  $\mu\text{M}$  according to  $3S_b/S$ , in which 3 is signal-to-noise ratio,  $S_b$  is the standard deviation of the current response in the absence of  $\text{H}_2\text{O}_2$  and  $S$  is the sensitivity.<sup>16,41</sup> The sensitivity of the rGO/GNP electrode towards  $\text{H}_2\text{O}_2$  is  $2.4 \times 10^{-2}$   $\text{mA mM}^{-1}$  according to the slope of the  $I-t$  calibration curve.

To investigate the anti-interference performance of the rGO/GNP sensor in detecting  $\text{H}_2\text{O}_2$ , several common interfering species including glucose (Glu), citric acid (CA) and AA are used in the  $I-t$  measurement at  $-0.3$  V. As shown in Fig. 5b, with successive injection of  $\text{H}_2\text{O}_2$  (1.2 mM) and three interfering species (Glu, CA, AA, 1.0 mM), the rGO/GNP electrode only presents obvious response to  $\text{H}_2\text{O}_2$  and the current keeps steady upon the addition of the interfering species. Such satisfactory selectivity could be attributed to the low potential of  $-0.3$  V used in the  $\text{H}_2\text{O}_2$  detection and the uniform dispersion of GNPs in the composite aerogel film, which indicates that the aerogel

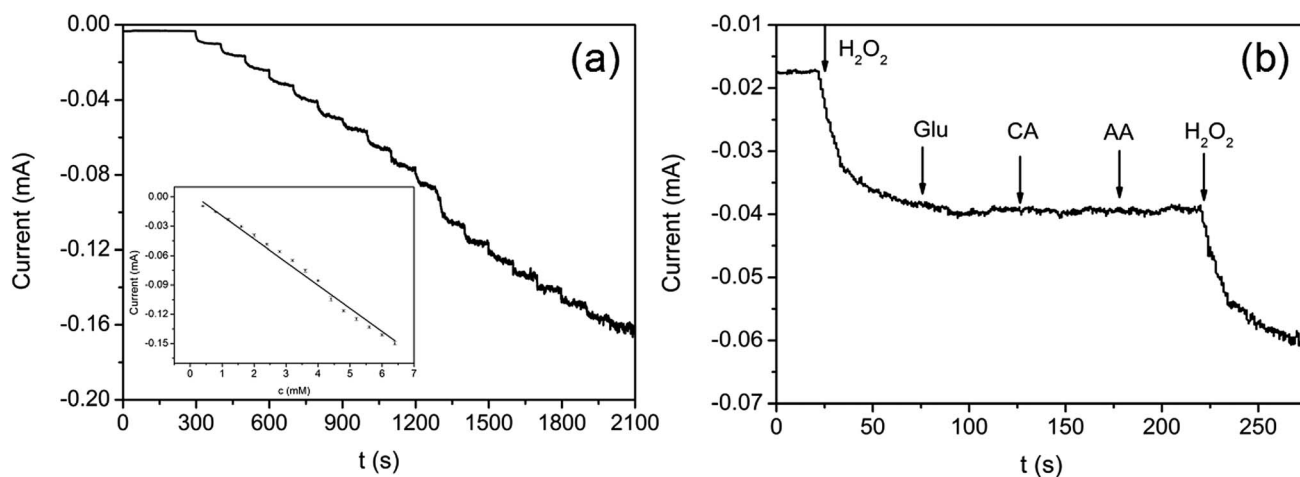


Fig. 5 (a) Amperometric  $I-t$  curve for the rGO/GNP aerogel electrode with successive injection of  $\text{H}_2\text{O}_2$  in PBS (pH = 7) at  $-0.3$  V (inset: calibration plot of current vs.  $\text{H}_2\text{O}_2$  concentration). (b) Amperometric  $I-t$  curve for the rGO/GNP electrode measured at  $-0.3$  V with the successive addition of  $\text{H}_2\text{O}_2$ , Glu, CA, AA and  $\text{H}_2\text{O}_2$  in PBS (pH = 7).



sensor possesses potential for sensing H<sub>2</sub>O<sub>2</sub> in practical application.

## Conclusions

In this study, we report the development of hydrothermal method for the one-step preparation graphene hydrogel film. The subsequent freeze-drying process yields thin aerogel film that could be tailored for direct use of electrochemical electrode. The Au-decorated aerogel film sensor shows good behaviours in H<sub>2</sub>O<sub>2</sub> detection, with a broad linear range at 0.4–6.4 mM, low detection limit of 12 μM and remarkable selectivity against common interfering species. Our work may provide new strategy for the preparation of graphene-based functional hydrogel/aerogel film materials, which could be used in various fields, such as sensor, catalysis and energy storage.

## Conflicts of interest

There are no conflicts to declare.

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