

## Supporting Information

Controlled Synthesis of Graphene-Gd(OH)<sub>3</sub> Nanocomposites and Their Applications for Detection  
of Ascorbic Acid

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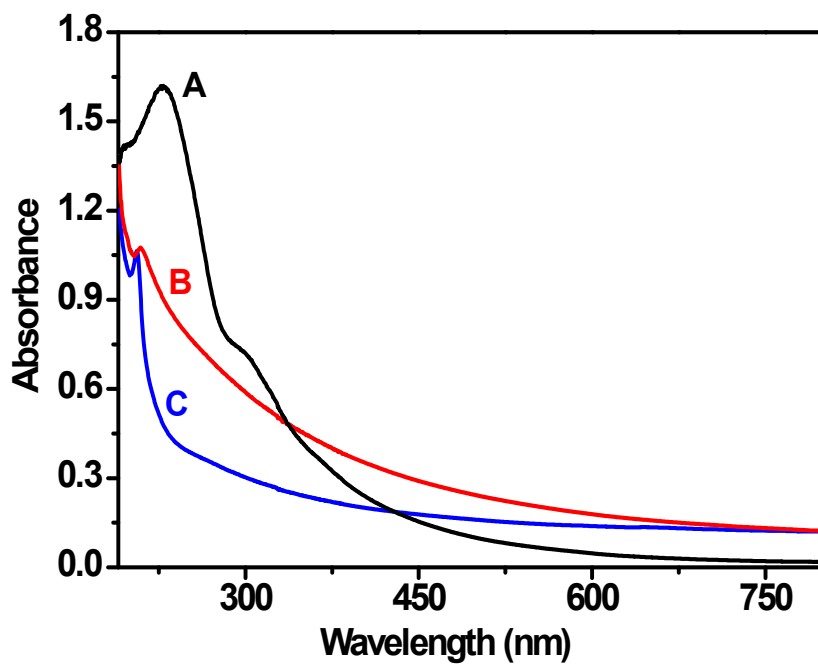


Fig. S1. UV-vis spectra of GO (A), GR-Gd(OH)<sub>3</sub>-1 (B) and GR-Gd(OH)<sub>3</sub>-2 (C).

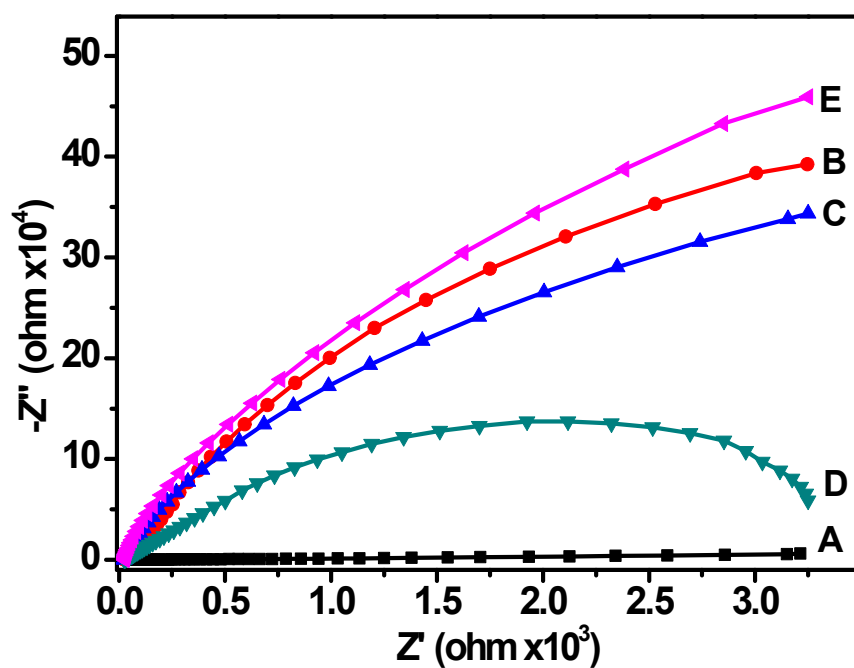
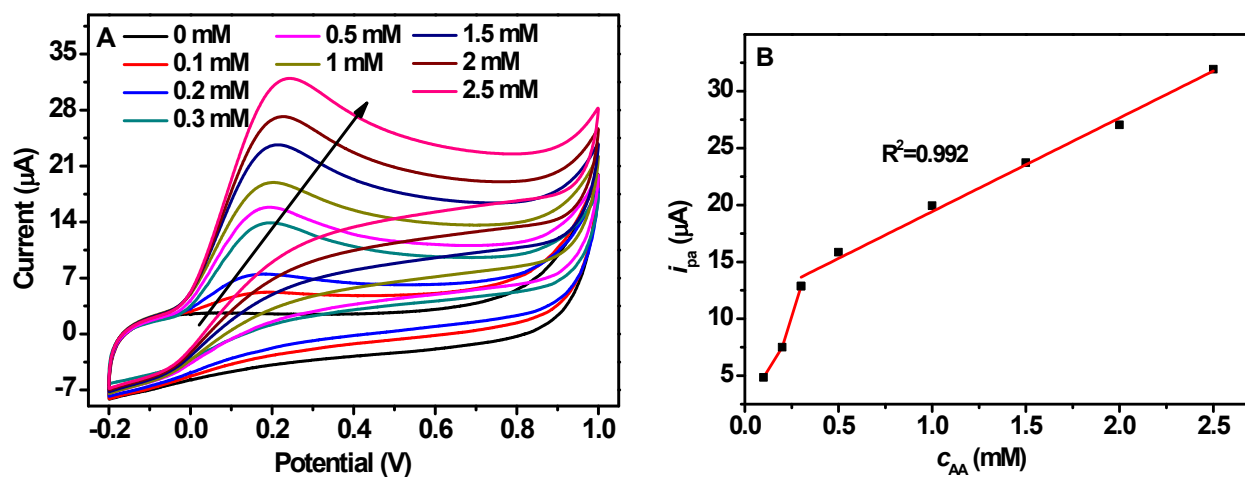


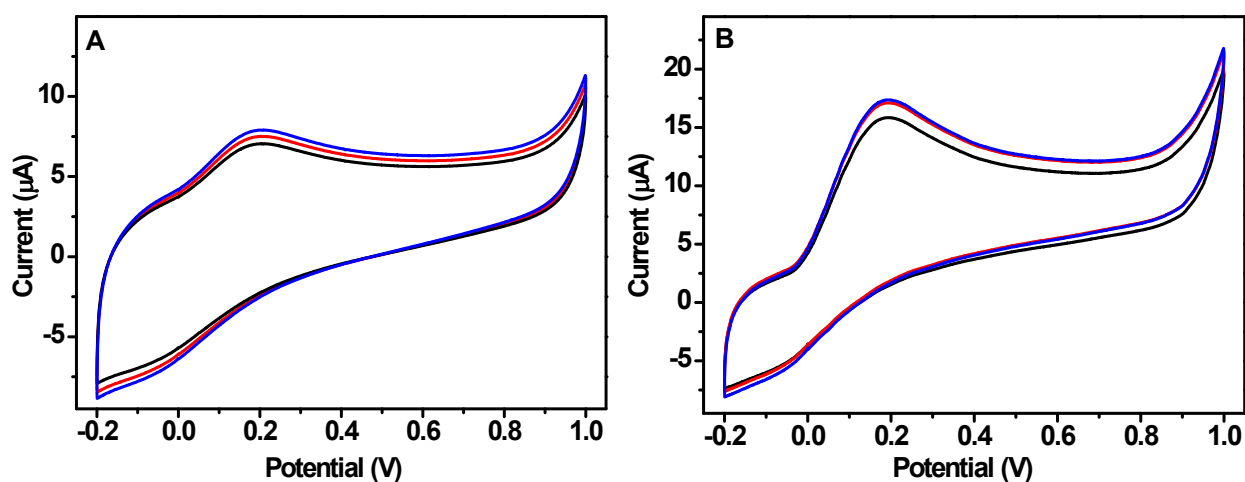
Fig. S2. EIS of GCE/GR (A), GCE/Gd(OH)<sub>3</sub> (B), GCE/GR-Gd(OH)<sub>3</sub>-1 (C), GCE/GR-Gd(OH)<sub>3</sub>-2 (D) and bare GCE (E) in 0.1 M PBS-0.5 mM AA aqueous solution recorded in the frequency range of 0.0-4.0 x 10<sup>4</sup> Hz.



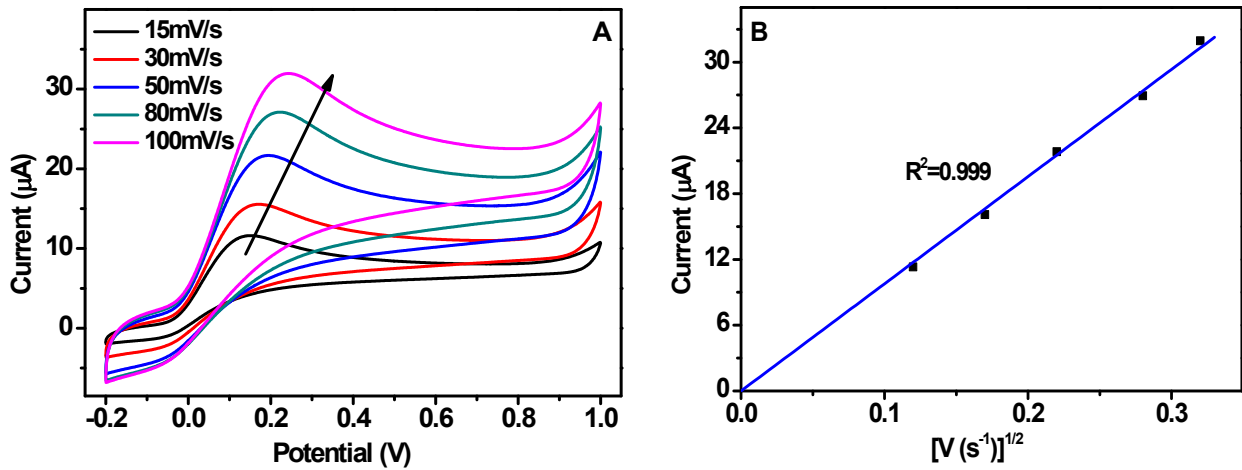
**Fig. S3** A) CV of GCE/GR-Gd(OH)<sub>3</sub>-2 in 0.1 M PBS (pH = 7.0) at varying concentration of AA (0.1-2.5 mM) and B) the relationship between  $i_{pa}$  and  $c_{AA}$ .

Holding the potential of oxidation peak at 0.185 V with the  $c_{AA}$  ranged from 0.3-2.5mM, a linear relationship between  $i_{pa}$  and  $c_{AA}$  was found. The resulting calibration plot is a straight line, with a coefficient of 0.992, given by the equation:

$$i_{pa}(\mu A) = 8.23c_{AA}(\text{mM}) + 11.19 \quad (\text{S1})$$



**Fig. S4** CV curves of GCE/GR-Gd(OH)<sub>3</sub>-1 (A) and GCE/GR-Gd(OH)<sub>3</sub>-2 (B) in 0.1 M PBS with 0.5mM AA at 100 mV s<sup>-1</sup>. In each case the measurements were performed for three times to check the reproducibility.



**Fig. S5** (A) CV of GCE/GR-Gd(OH)<sub>3</sub>-2 in 2.5 mM AA aqueous solution at varying scan rate (15-100 mV/s) and (B) the relationship between  $i_{pa}$  and  $V^{1/2}$ .

**Table S1.** BET specific surface area ( $S_{BET}$ ) and total pore volume of Gd(OH)<sub>3</sub>, GR-Gd(OH)<sub>3</sub>-1 and GR-Gd(OH)<sub>3</sub>-2.

Sample	Gd(OH) <sub>3</sub>	GR-Gd(OH) <sub>3</sub> -1	GR-Gd(OH) <sub>3</sub> -2
$S_{BET}$ (m <sup>2</sup> ·g <sup>-1</sup> )	3.23	18.02	32.38
Total pore volume (cm <sup>3</sup> ·g <sup>-1</sup> )	0.29	0.36	0.41

### Calculation of the lowest limit of detection (LOD)

The LOD of GCE/GR-Gd(OH)<sub>3</sub>-1 and GCE/GR-Gd(OH)<sub>3</sub>-2 can be calculated to be 0.06 mM and 0.05 mM, respectively. Nine blank experiments of GCE/GR-Gd(OH)<sub>3</sub>-1 and GCE/GR-Gd(OH)<sub>3</sub>-2 were carried out in 0.1 M PBS without AA, from which nine oxidation peak currents were obtained. Then LOD of modified GCE can be calculated by the equations shown below (based on S/N=3) :

$$SD = \sqrt{\frac{\sum (x_i - \bar{x})^2}{n-1}}$$

$$LOD = \frac{3SD}{s}$$

where  $x_i$  is the oxidation peak current of the nine experiments,  $\bar{x}$  is the averaged value of  $x_i$ ,  $n$  is the times of experiments. SD is standard deviation.  $s$  is the slope determined by the curves of a linear relationship between  $i_{pa}$  and  $c_{AA}$  of electrodes. Taking the calculation process of LOD of GCE/GR-Gd(OH)<sub>3</sub>-1 for example,  $s$  is 7.34, determined by equation 1 shown in the maintext. The detailed calculation of the LOD is as bellow:

$x_1/\mu A$	$x_2/\mu A$	$x_3/\mu A$	$x_4/\mu A$	$x_5/\mu A$	$x_6/\mu A$	$x_7/\mu A$	$x_8/\mu A$	$x_9/\mu A$	$\bar{x} / \mu A$
2.683	2.598	2.817	2.528	2.833	2.741	2.479	2.475	2.849	2.667

### Calculation of the relative standard deviation (RSD)

Taking the calculated method of RSD of GCE/GR-Gd(OH)<sub>3</sub>-1 for example, the detailed calculation is shown below:

$$SD = \sqrt{\frac{\sum (x_i - \bar{x})^2}{n-1}}$$

$$RSD = \frac{SD}{\bar{x}} = \frac{\sqrt{\frac{\sum (x_i - \bar{x})^2}{n-1}}}{\bar{x}}$$

where  $x_i$  is the oxidation peak current of GCE/GR-Gd(OH)<sub>3</sub>-1,  $\bar{x}$  is the averaged value of  $x_i$ ,  $n$  is the number of the experiment, SD is the standard deviation.

$x_1/\mu\text{A}$	$x_2/\mu\text{A}$	$x_3/\mu\text{A}$	$\bar{x}/\mu\text{A}$
7.511	7.902	7.045	7.486