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# Facile preparation of covalently functionalized graphene with 2,4-dinitrophenylhydrazine and investigation of its characteristics

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This article reports a fast and easy method for simultaneously *in situ* reducing and functionalizing graphene oxide. 2,4-Dinitrophenylhydrazine hydrate salt molecules are reduced by graphene oxide by reacting with oxide groups on the surface and removing these groups, and 2,4-dinitrophenylhydrazone groups are replaced with oxide groups. The synthesized materials have been investigated using Fourier transform infrared spectroscopy (FTIR), Raman spectroscopy, X-ray diffraction (XRD), thermogravimetric analysis (TGA), X-ray photoelectron spectroscopy (XPS), and UV absorption. Also, the morphology has been examined with a scanning electron microscope (SEM) and Brunauer–Emmett–Teller (BET) analysis. The result of the photocurrent response and electrochemical behavior of the samples through cyclic voltammetry, galvanostatic charge/discharge, and electrochemical impedance spectroscopy (EIS) have been analyzed to investigate the effect of physical and chemical changes compared to graphene.

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## 1. Introduction

Graphene is a two-dimensional monolayer of carbon atoms with a hexagonal honeycomb-like structure in which the carbon bonds have  $sp^2$  hybridization, and the out-of-plane  $\pi$  bond creates a network of decentralized electrons that is responsible for electron conduction and also causes a weak interaction between graphene layers or between graphene and a substrate. Due to its unique structural features, graphene has shown exceptional physical properties, which has attracted the attention of many researchers in various sciences.<sup>1–4</sup>

Various methods have been used to synthesize graphene and its derivatives. Due to weak van der Waals forces among the layers in graphite, pristine graphene can be obtained from the mechanical exfoliation of graphite.<sup>5,6</sup> The growth of graphene sheets using bottom-up techniques such as chemical vapor deposition (CVD) has been used to grow single-layer and multi-layer graphene sheets on metal surfaces.<sup>7–9</sup> However, these methods are not suitable for large-scale production. Therefore, oxidation and exfoliation of graphite oxide, followed by a reduction process, have been widely used and considered to prepare reduced graphene oxide (R-GO).<sup>10–13</sup> However, the precursor of R-GO, graphite needs to be exfoliated through the reaction with a mixture of oxidants to create graphene oxide (GO) sheets.<sup>14</sup> Therefore, GO sheets are strongly oxidized and characterized by oxygen-containing groups.<sup>15,16</sup> To remove these groups and prepare R-GO, various reducing agents; such as

hydrazine,<sup>17,18</sup> strong alkaline environments,<sup>19</sup> vitamin C,<sup>20,21</sup> and bovine serum albumin (BSA);<sup>22,23</sup> and various methods; such as electrochemical,<sup>24,25</sup> photochemical,<sup>26,27</sup> and thermal reduction<sup>28,29</sup> procedures; have been used.

Reduced graphene oxide has a wide range of applications, including its use in energy storage devices,<sup>30,31</sup> electronic devices,<sup>32</sup> biomedical applications,<sup>33</sup> sensors,<sup>34</sup> membranes,<sup>35</sup> catalysts,<sup>36,37</sup> water purification,<sup>30</sup> *etc.* However, the presence of these functional groups in GO and R-GO compared to pristine graphene allows the ability to adjust the electrical and optical properties through chemical reactions. However, unfortunately, due to the incomplete reduction and the presence of multiple defects, which disrupt the  $sp^2$  lattice, R-GO exhibits lower electrical conductivity than pristine graphene.<sup>38–40</sup> In addition, the solubility of R-GO in water is minimal, so the direct dispersion of graphene sheets in water is impossible without the help of dispersing agents.<sup>41</sup> To improve these properties, various functional groups have been attached to the graphene structure by chemical modification,<sup>42,43</sup> covalent,<sup>44–48</sup> or non-covalent<sup>49–52</sup> functionalization.

The functional groups covalently attached to the carbon atoms on the graphene convert the triangular  $sp^2$  orbital to a tetragonal  $sp^3$  orbital. Such transformations drastically change the electronic properties of pristine graphene.<sup>53–56</sup> In like manner, by functionalizing graphene, the specific capacity increases by increasing the quantum capacity of these sheets due to the formation of new levels in the vicinity of the Fermi level.<sup>57,58</sup> Also, due to the creation of steric repulsion caused by the presence of functional groups, the graphene sheets' recombination rate decreases, which leads to an increase in the available surface in functionalized graphene.<sup>59,60</sup> Recent

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experimental studies have shown that the covalent functionalization of graphene using a nitrophenyl group directly affects the magnetic properties of the graphene surface.<sup>61–64</sup> During the functionalization process, converting the carbon centers from  $sp^2$  to  $sp^3$  saturates the carbon atoms. It opens a barrier band gap for electrons, which enables the production of insulating and semiconducting regions in graphene wafers.<sup>65,66</sup>

This article presents a method for preparing graphene functionalized with nitrophenyl groups, in which graphene oxide is *in situ* and simultaneously reduced and functionalized with 2,4-dinitrophenylhydrazine groups. Also, this method was performed at a pH close to neutral and without dispersing agents, while for the synthesis of reduced graphene oxide (R-GO), to avoid re-adhesion of the layers and obtain the lowest number of layers as well as the lowest amount of defects, the pH is usually set at about 10, and a scattering agent is used.<sup>67</sup> The structural and chemical differences of the material synthesized with graphene were studied. Also, the electrochemical behavior of this material with graphene was investigated to check the effectiveness of the changes made, due to the high sensitivity of electrochemical techniques to changes in the structure.

## 2. Materials and procedures

### 2.1. Materials

Graphite powder (<45  $\mu\text{m}$ ), sulfuric acid ( $\text{H}_2\text{SO}_4$ , 98%), potassium persulfate ( $\text{K}_2\text{S}_2\text{O}_8$ , 99%), sodium nitrate ( $\text{NaNO}_2$ ), potassium permanganate ( $\text{KMnO}_4$ , 98%), hydrochloric acid ( $\text{HCl}$ , 37%) and hydrogen peroxide ( $\text{H}_2\text{O}_2$ , 30 wt%) were used to synthesize GO. To prepare graphene, hydrazine was used to reduce graphene oxide. Likewise, 2,4-dinitrophenylhydrazine was used to reduce and functionalize graphene oxide. Sodium sulfate was used to make electrolytes. Graphite powder, hydrazine, and 2,4-dinitrophenyl hydrazine were bought from Merck, and the rest of the materials from Sigma-Aldrich.

### 2.2. Preparing graphene oxide (GO)

The Hummers method was used to synthesize the graphene oxide. Concisely,  $\text{K}_2\text{S}_2\text{O}_8$  (5 g),  $\text{P}_2\text{O}_5$  (5 g), and concentrated  $\text{H}_2\text{SO}_4$  (50 mL) were mixed at 90 °C, and then the temperature was lowered to 80 °C. Then, graphite powder (10 g) was added to this mixture and stirred for 5 hours. After that, distilled water (1000 mL) was added to the mixture and after 24 hours, the suspension was filtered and dried at ambient temperature. Afterward, using an ice bath, concentrated  $\text{H}_2\text{SO}_4$  (200 mL) was cooled to 0 °C and the obtained deposit was added.  $\text{KMnO}_4$  (20 g) was slowly added while the temperature was kept below 10 °C. The suspension was stirred for 2 hours at a temperature of 35 °C, and then distilled water (460 mL) was added. After 2 hours, distilled water (140 mL) and 30%  $\text{H}_2\text{O}_2$  (25 mL) were added. Then the suspension was left at rest for 24 hours, and the clear liquid was drained. The remaining sediment was washed using a centrifuge with 10%  $\text{HCl}$  solution and distilled water.

### 2.3. Reduction and functionalization of graphene oxide with DNPH (NP-GO)

200 mg of the graphene oxide obtained from the previous procedure was dispersed in distilled water (200 mL) using an ultrasonic bath model ELEMA-P30H for 1 h. The pH was adjusted to 8 by 1 M  $\text{NaOH}$  (using Istek pH meter model 240L). Then 2,4-dinitrophenylhydrazine salt (2 g) was added to the suspension and stirred for 2 hours at 80 °C. Afterward, the suspension was filtered using PTFE filter paper (0.22  $\mu\text{m}$ ), and the obtained product was washed with ethanol, acetone, as well as a mixture of acetonitrile/water and, finally, dried for 24 hours at 70 °C.

### 2.4. Synthesize of graphene (R-GO)

200 mL of GO suspension (1 mg/1 mL) was prepared to prepare graphene, and 1 w% sodium dodecyl sulfate (SDS) as a dispersing agent was added. Subsequently, the suspension was put in the ultrasonic bath for 1 h. Then, the pH of the solution was adjusted to 10 *via* 1 M  $\text{NaOH}$ . Afterward, hydrazine hydrate 60% (2 mL) was added and the suspension was stirred for 2 h at 90 °C.

## 3. Investigation of structure and properties

### 3.1. Characterization of the samples

FTIR (SHIMADZU company, model 8400S), Raman spectroscopy (TEKSAN company model Takram), and thermogravimetric analysis (Bahr device model STA 540) were used to investigate the effect of reducing and functionalizing procedures. XRD (Bruker D8 advance) at 40 kV using  $\text{Cu K}\alpha$  radiation ( $\lambda = 1.54 \text{ \AA}$ ) was used to identify phases presented in the samples. UV absorption (spectral absorption measurements) was measured with Shimadzu model UV-160 A. X-ray photoelectron spectroscopy (XPS) was carried out *via* an ESCA2000 under the vacuum pressure of  $\sim 10^{-10}$  torr with an X-ray source of  $\text{Al K}\alpha$  (1486.6 eV). The surface features of powders were studied according to the Brunauer–Emmett–Teller (BET) method (Micrometric company model ASAP 2020). The morphology of synthesized materials was also studied by SEM (Scanning electron microscopy, TESCAN model VEGA 2).

### 3.2. Photoelectrochemical measurements

A homemade system was used to measure the photoelectrochemical behavior. As a source of radiation, a Xe lamp (500 W) with a cut-off filter ( $\lambda > 325 \text{ nm}$ ) was employed. The powder of the samples (20 mg) was dissolved in acetone and ultrasonicated for 30 minutes, then a grain of iodine was added to the suspension. The ultrasonication continued for another 30 minutes. Afterward, the prepared samples were electroplated (15 volts for 10 minutes) on the surface of ITO glass with an area of 1  $\text{cm}^2$ . The photocurrent response was measured in a 0.5 M  $\text{Na}_2\text{SO}_4$  (sodium sulfate solution) in a three-electrode configuration system containing indium tin oxide (ITO) as the working electrode, platinum wire as the counter electrode, and saturated



Ag/AgCl as the reference electrode. The interval between turning on and off the light was 4 seconds. Electrochemical impedance spectroscopy (EIS) was conducted at 100 kHz to 100 mHz in an open circuit potential (OCP).

### 3.3. Electrochemical measurements

The electrochemical behavior of the samples was investigated by cyclic voltammetry and galvanostatic charge/discharge methods, KIMIA STAT device model 126. Electrochemical impedance spectroscopy (EIS) was studied by the Autolab device model PGSTAT204. A three-electrode system including prepared samples (working electrode), a platinum electrode (counter electrode), Ag/AgCl (reference electrode), and sodium sulfate solution (0.5 M) as the electrolyte was used.

To make electrode materials for electrochemical analyses, prepared material, black acetylene and polytetrafluoroethylene (PTFE) were mixed with a weight ratio of 80:15:5, and by adding ethanol and then 15 min ultrasonication, a grout was prepared with an appropriate concentration. The prepared material for the electrode was deposited on nickel foam with an area of 1 cm<sup>2</sup> by the painting method. In the end, the prepared electrodes were dried for 24 h at 70 °C. EIS measurement was conducted in OCP at a frequency range between 100 kHz to 10 mHz. Capacities of the materials were obtained from the galvanostatic charge/discharge method by the following formula:

$$C = \frac{i \times t}{V \times m}$$

In this equation,  $i$  is the applied current,  $t$  is the discharge time,  $V$  is the potential window and  $m$  is the weight of materials deposited on the electrode (0.8 mg).

## 4. Result and discussions

### 4.1. Structural investigation

Fig. 1a illustrates the results of FTIR spectra corresponding to GO, R-GO, and NP-GO. The existence of oxygen-containing groups; C–O groups (at 1050 cm<sup>-1</sup>), ketone C=O (at 1620 cm<sup>-1</sup>), carboxyl C=O (at 1720 cm<sup>-1</sup>), O–H and CO–OH groups, as well as water molecules in GO layers (at 3500 cm<sup>-1</sup>); can be verified in the spectrum of GO. In the graphene (R-GO) sample, carboxyl, ketone, ether, and O–H peaks disappeared, indicating that most of the oxygen groups were removed due to the reaction with hydrazine. In addition, 2850 to 2920 cm<sup>-1</sup> peaks are associated with asymmetric C–H stretching. After reducing and functionalizing graphene oxide with 2,4-dinitrophenylhydrazine, peaks that appeared at 1345 and 1515 cm<sup>-1</sup> are connected to the asymmetric and symmetric stretching vibration related to NO<sub>2</sub> groups. The peak appearing at 1590 cm<sup>-1</sup> assumes to be associated with the aromatic stretch. Likewise, the peak corresponding to stretching vibrations due to the presence of C–N nitrobenzene groups appears at 850 cm<sup>-1</sup>. The broad peak in the 3300–3500 cm<sup>-1</sup> region seems to correspond to the N–H group of 2,4-dinitrophenylhydrazine.

Fig. 1b displays the results of the X-ray diffraction (XRD) analysis of the samples. In GO, bonding with groups that contain oxygen increases the spacing between the layers of

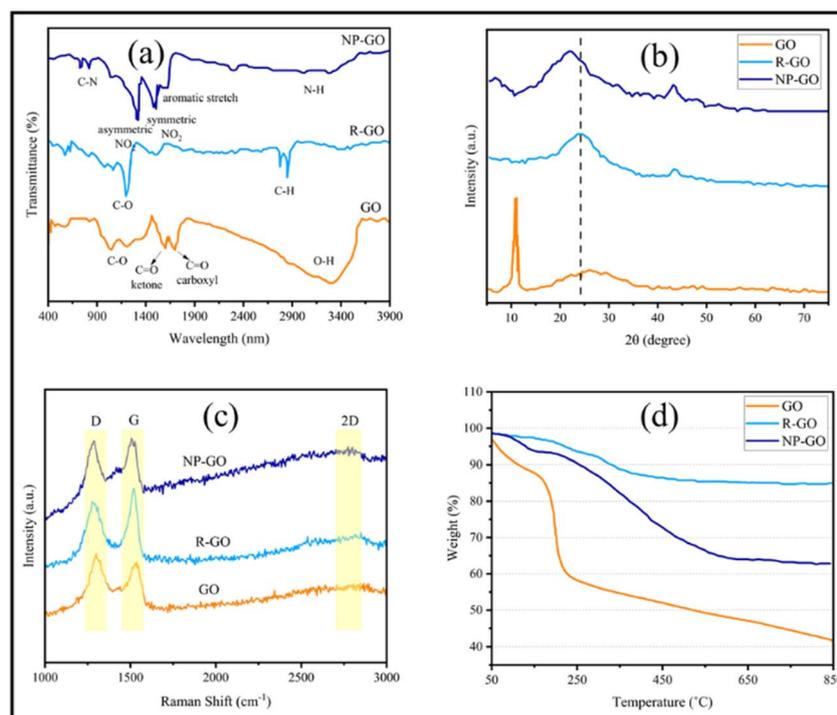


Fig. 1 (a) FTIR, (b) XRD, (c) Raman spectra, and (d) TGA plots of GO, R-GO, and NP-GO.



graphite. Therefore, the position of the peak indicates the oxidation level of graphite.<sup>68</sup> Accordingly, the peak at 11.6° denotes the 7.57 Å *d*-spacing in graphene oxide. After GO reduction, R-GO sheets approach together and result in a peak at 23.8° (3.73 Å *d*-spacing) with low intensity. In NP-GO, the reaction of GO with DNPH leads to the removal of oxygen groups in GO and the elimination of the sharp, strong peak related to GO. Furthermore, it yields a broad peak at 22.7° (3.89 Å *d*-spacing). Based on the results of the FTIR analysis that show the functional groups' existence and by comparing the *d*-spacing value for R-GO and NP-GO, it seems that the difference

in the distance of the layer is caused by the reduction and functionalizing of graphene oxide.

Fig. 1c shows the results of the Raman spectra of samples. The Raman device was equipped with a 532 nm laser and a thermoelectrically cooled CCD detector. A 60×-objective microscope lens was used to focus on the surface, and the exposure time was 5 s. As it is apparent, there are two bands. The first one relates to band D which signifies the graphene flakes' defect and irregularity, and the second corresponds to the G band that represents the first scattering of E<sup>2g</sup> phonon associated with carbon-carbon bonds with the hybridization of sp<sup>2</sup>. The 2D band is derived from the second-order double-

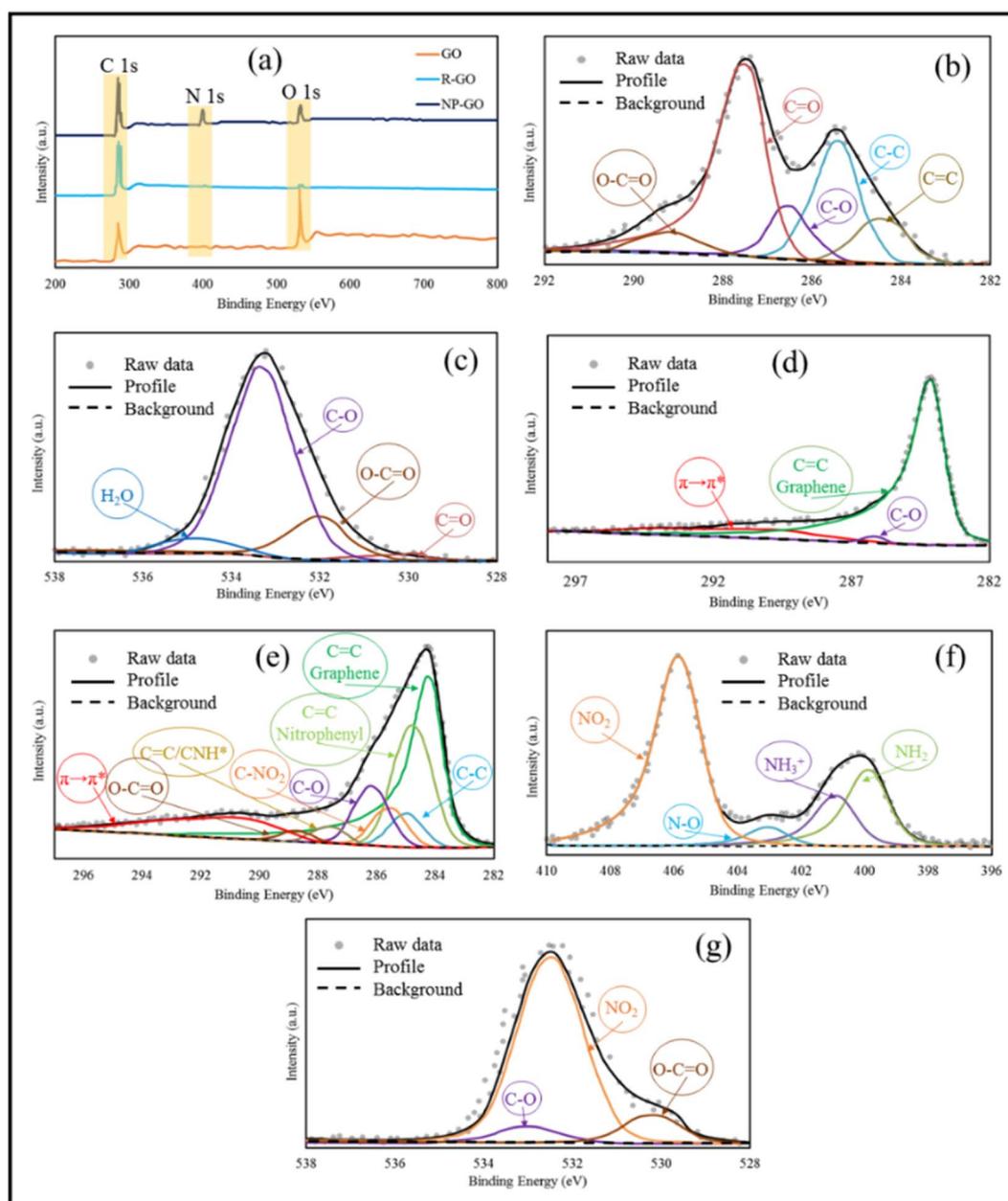


Fig. 2 (a) XPS spectra of GO, R-GO, and NP-GO, (b) high-resolution C 1s spectra for GO, (c) high-resolution O 1s spectra for GO, (d) high-resolution C 1s spectra for R-GO, (e) high-resolution C 1s spectra for NP-GO, (f) high-resolution N 1s spectra for NP-GO, and (g) high-resolution O 1s spectra for NP-GO.



resonant process between the  $K$  and  $K'$  points in the BZ involving two iTO phonons. The 2D band is highly sensitive to the Fermi velocity and iTO phonon band slope due to the unique electronic band structure of graphene.<sup>69</sup> The increased intensity of the D and 2D bands after functionalization indicates the attachment of functional groups.<sup>70</sup> The level of reduction and defect is determined through the ratio between  $I_D$  and  $I_G$  which in GO, R-GO, and NP-GO are 0.89, 1.32, and 1.02, respectively. After the reduction of graphene oxide, the band D intensity increases due to the formation of defects in graphene sheets due to the reaction with hydrazine.<sup>71</sup> It has been shown that the  $I_{2D}/I_G$  ratio of the Raman spectra can indicate the number of layers in graphene materials.<sup>72</sup> This ratio for GO, R-GO, and NP-GO are 0.81, 0.75, and 0.94, respectively, which indicates that these materials are multilayered. Considering these increases in the ratio of  $I_{2D}/I_G$  and  $I_D/I_G$  after the reaction of GO with DNP, it can be concluded that these changes are attributable to the reduction and replacing oxygen-containing groups in GO with 2,4-dinitrophenylhydrazone groups. In addition, the lower ratio of  $I_D/I_G$  compared to the graphene suggests that there are fewer irregularities and defects in the NP-GO structure.

Fig. 1d exhibits the thermogravimetric analysis (TGA) of the samples. During this process, the samples were heated to a temperature of 850 °C at a rate of 10 °C min<sup>-1</sup> in an Ar atmosphere, and the weight changes of the samples were for the temperature. The initial weight loss of graphene oxide (GO) at 100 to 200 °C is associated with water and residual decomposition. The gradual weight loss from 200 to 650 °C is due to the decomposition of oxygen-containing groups on the surface and edge of GO sheets. Graphene (R-GO) shows a much lower reduction than GO, which may be related to removing a high percentage of oxidizing groups during reduction by hydrazine. In the sample of NP-GO, reducing 5% of the original weight under 200 °C indicates that there are no more than 5% unstable oxide groups in this sample. The lower weight loss of NP-GO compared to GO and R-GO can be a confirmation of reducing and functionalizing of graphene oxide with 2,4-dinitrophenylhydrazone.

Using XPS, the chemical composition of the samples was investigated more closely. Fig. 2a represents the XPS spectra of GO, R-GO, and NP-GO. In addition, the binding energy and percentage of compounds from C 1s, O 1s, and N 1s are listed in Table 1. The results showed carbon, oxygen, and nitrogen, which shows that the nitrophenyl groups are present on the surface of the NP-GO. In Fig. 2b it is obvious that the C 1s spectrum of graphene oxide consists of peaks at 284.6 eV (sp<sup>2</sup> carbon), 286.6 eV (epoxide), and 288.5 eV (carboxyl groups).<sup>73</sup> From the O 1s spectrum of GO (Fig. 2c), the oxygen-containing groups are evident as C–O, C=O, C–O–C, O–H, and H<sub>2</sub>O groups. The C 1s spectrum of R-GO (Fig. 2d) shows a band at 284.6. The intensity of epoxide, carboxyl, or other bands is greatly reduced compared to graphene oxide, which means that the C=C bonds relative percentage increased and it decreased for C–O bonds. The XPS spectrum of R-GO results shows that this sample only contains C and O, which designates that there are no impurities in the sample. The spectrum of C 1s related to NP-GO (Fig. 2e)

shows the presence of the C=C peak related to graphene, the C=C bond corresponds to nitrophenyl groups (284.8 eV), the graphene C–C defect (at the site of bonding, 285 eV), and the peak related to C–NO<sub>2</sub> (285.6 eV). The ratio of O/C in GO, R-GO, and NP-GO samples is 1.8, 0.04, and 0.2, respectively, showing that most oxygen-containing groups are removed and GO is reduced. The N 1s spectrum of the NP-GO (Fig. 2f) indicates that some nitro groups were reduced to other groups such as amines, and protonated amines due to the reduction by the irradiation of X-ray.<sup>74</sup> The O 1s spectrum of NP-GO (Fig. 2g) shows three peaks at 531.3 eV (carboxyl groups), 532.7 (nitro groups), and 533.1 eV (epoxy groups) in the NP-GO sample. The nitro groups come from 2,4 dinitrophenylhydrazine, while the carboxyl and epoxy groups come from non-reduced groups in GO.

UV-vis spectroscopy was used to probe the interactions on the surface of the samples. As can be seen in Fig. 3, graphene oxide shows a peak at 235 nm, corresponding to  $\pi$ - $\pi^*$  transitions related to the presence of the remaining C=C bonds with sp<sup>2</sup> hybridization,<sup>75</sup> which were shifted to 265 nm (longer wavelengths) after reduction, rebuilding the  $\pi$ -bond network.<sup>76</sup> With the increase in  $\pi$ -bonding, the absorption peak shifts to a longer wavelength because less energy is needed for transition. The UV-visible spectrum of 2,4-dinitrophenylhydrazine showed two peaks at 222 nm related to the  $\pi$ - $\pi^*$  and 360 nm corresponding to n- $\pi^*$  transitions related to the aromatic

Table 1 The binding energy and percentage of compounds from C 1s, O 1s, and N 1s were obtained from the XPS spectra of the samples

Sample	Peak	Assignment	Binding energy (eV)	Atomic (%)
GO	C 1s	C=C	284.5	11.41
		C–C	285.4	24.58
		C–O	286.6	9.64
		C=O	287.9	48.48
		COOH	289.2	5.89
	O 1s	C=O	530.1	4.34
		C=O, O–H, C–O–C	532.3	13.87
		C–O	533.8	74.14
		H <sub>2</sub> O	535.2	7.65
R-GO	C 1s	C=C graphene	284.5	90.62
		C–O	286.4	3.75
		$\pi \rightarrow \pi^*$	291.3	5.63
NP-GO	C 1s	C=C graphene	284.1	42.84
		C=C nitrophenyl	284.7	25.89
		C–C	285.3	5.26
		C–NO <sub>2</sub>	285.8	4.76
		C–O	286.4	8.55
		C=O, C–NH <sub>3</sub> <sup>+</sup>	287.8	1.87
	N 1s	O–C=O	288.7	1.25
		$\pi \rightarrow \pi^*$	290.7	9.94
		NH <sub>2</sub>	399.9	19.54
		NH <sub>3</sub> <sup>+</sup>	401.3	12.13
		NO	403.5	4.92
		NO <sub>2</sub>	406.1	63.41
O 1s	O–C=O	531.4	14.28	
	NO <sub>2</sub>	532.8	74.35	
	C–O	533.4	11.37	



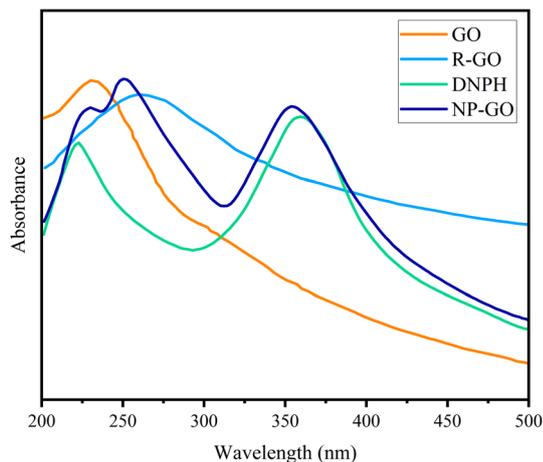


Fig. 3 UV spectra of GO, R-GO, DNP, and NP-GO.

compound.<sup>77</sup> The absorption band of NP-GO species corresponding to the  $\pi$ - $\pi^*$  transition becomes broad and also shows a redshift from 222 to 229 nm, while the  $n$ - $\pi^*$  transition peak shifts from 360 to 354 nm. Also, the peak of GO had a redshift to 247 nm, indicating the recovery of the  $\pi$  bond network as a result of the reduction. However, due to the presence of

functional groups, this displacement is lower than that of the R-GO sample. These changes and shifts in absorption peaks indicate the existence of  $\pi$ - $\pi^*$  interactions, electronic connections among the functional groups, and reduced graphene oxide sheets.

#### 4.2. Morphological characterizations

Fig. 4 shows the SEM images of R-GO and NP-GO. It is obvious from Fig. 4a and b that the morphology of R-GO is wrinkled and crumpled, due to agglomeration through van der Waals forces and  $\pi$ - $\pi$  interactions among sheets. As can be seen, the flakes of R-GO are in layers. Small R-GO flakes aggregate adjacent to each other and seem to have a bigger mass. The obtained layer structure seems to have formed during the filtering of the suspension.

Fig. 4c and d shows the morphology of NP-GO. Due to the presence of 2,4-dinitrophenylhydrazine groups and, as a result, reducing van der Waals forces and forming spatial repulse among functionalized sheets, aggregation is reduced in comparison to the R-GO, thus there is a more accessible active surface in this sample.

The binding effect of organic molecules on the surface of NP-GO can be measured and compared with graphene through the specific surface area (Fig. 5a) and the distribution of pore size

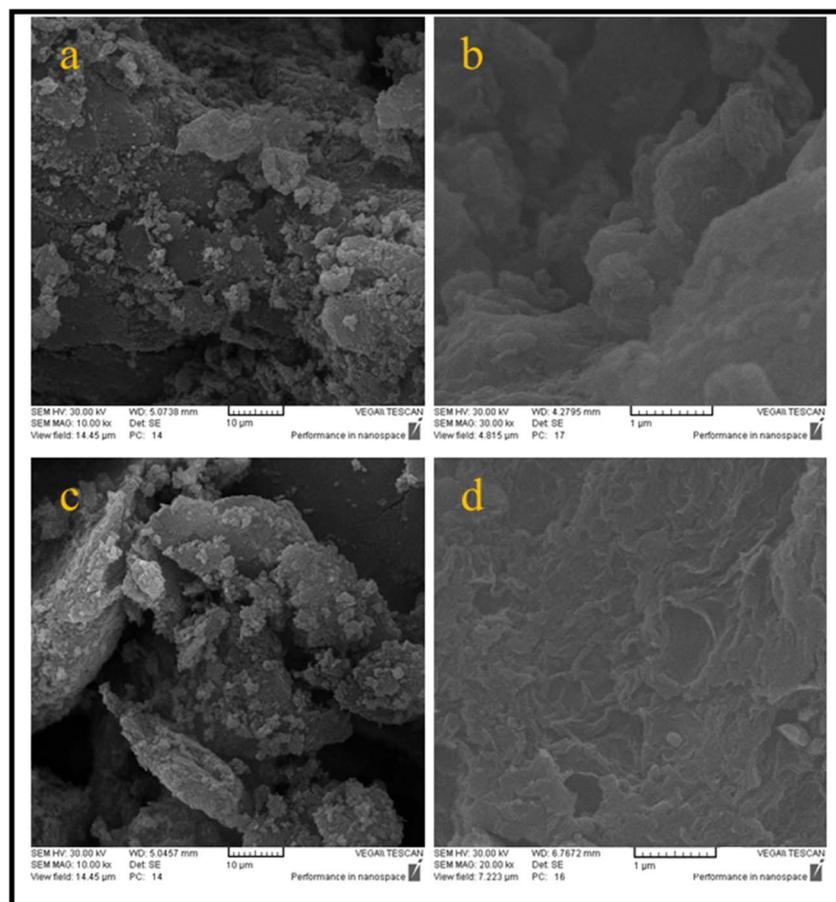


Fig. 4 SEM images related to the morphology and structure of (a and b) R-GO and (c and d) NP-GO in different magnifications.



(Fig. 5b). Fig. 5a show that the specific surface area of R-GO is near to  $500 \text{ m}^2 \text{ g}^{-1}$ . It contains small mesoporous (2–50 nm), indicating that the reduction of graphene oxide by hydrazine with the loss of oxygen functional groups causes porosity. However, the value lower than the theoretical specific surface area for a single graphene sheet ( $2600 \text{ m}^2 \text{ g}^{-1}$ ) is due to the sheets' accumulation and overlying after the drying process.<sup>78</sup> After the grafting of 2,4-dinitrophenylhydrazine groups, the surface area for NP-GO increased to  $650 \text{ m}^2 \text{ g}^{-1}$ . This increase indirectly suggests the binding 2,4-dinitrophenylhydrazine on the surface of NP-GO. The removal of oxygen-containing groups

and the replacement of functional groups causes the creation of pores in the NP-GO sheets and as a result, causes a significant increase in the surface area and a significant difference in the distribution of pore size. Interestingly, the attachment of groups in NP-GO, which may be caused by the decarboxylation of groups in graphene oxide,<sup>79</sup> leads to smaller pores than graphene.

Fig. 5c displays the results of the  $\text{N}_2$  adsorption and desorption isotherm for R-GO and NP-GO at 77 K. Both samples show type I and II mixing isotherms of relative pressure ( $P/P_0$ ) in low and high regions.<sup>80</sup> Low absorption volume at low  $P/P_0$  (0–0.5) for samples is representative of mesoporous materials, confirmed by an H3 hysteresis loop and a plateau.<sup>81</sup> As it is apparent, absorb volume increases after the reduction and functionalization of graphene oxide with DNPH.

### 4.3. Photoelectrochemical descriptions

The efficiency of charge transfer/separation is very important in optical nonlinearity. Therefore, Photoelectrochemical investigations were conducted to study the electronic interactions on the samples. Fig. 6a shows the photocurrent performances of GO, R-GO, and NP-GO *versus* the time of the virtual on-and-off cycles of sunlight. For each on/off cycle, increases and decreases in photocurrent are obvious. When simulated sunlight irradiation is employed, photocurrent is produced *via* transferring and separating light charges to the working electrodes.<sup>82</sup> At off cycle, the working electrode's responses are weak and keep their dark current state. However, when the switch is on, the photocurrent's intensity increases due to the rapid separation of charge carriers that are sensitive to light.<sup>83</sup> Obviously, GO and R-GO exhibit weak photocurrent responses under simulated radiation of sunlight. At the same time, NP-GO, owing to the creation of paths for direct conduction through covalent bonds originating between 2,4-dinitrophenylhydrazine and reduced graphene oxide sheets, shows better performance. As a result, the separation and transportation of light-generated carriers were enhanced by removing oxygen-containing groups and introducing 2,4-dinitrophenylhydrazine groups on the GO surface.<sup>84</sup>

The Nyquist diagram of electrochemical impedance spectroscopy (EIS) was studied to further describe the charge carrier migration of the prepared samples. The results are shown in Fig. 6b. Subsequently, the proposed  $R_s(R_{ct}Q_{ct})$  model is used to study the resistance behavior of the oxide layer in terms of applied potential at a constant frequency range. In the resultant equivalent model,  $R_s$ ,  $R_{ct}$ , and  $Q_{ct}$  are the solution resistance, charge transfer resistance, and constant phase element (CPE), correspondingly. Since the EIS capacitor element does not designate steady and regular behavior owing to the concurrent complex corrosion procedure at the border, the constant phase element (CPE) as an expressive deviation from the ideal capacitor-based circuit was used.<sup>85</sup> The best-fitted parameters for the reached equivalent circuit are listed in Table 2. As can be seen, NP-GO shows a smaller charge transfer resistance than GO and R-GO, which indicates the reduction of the resistance of the solid-state interface layer

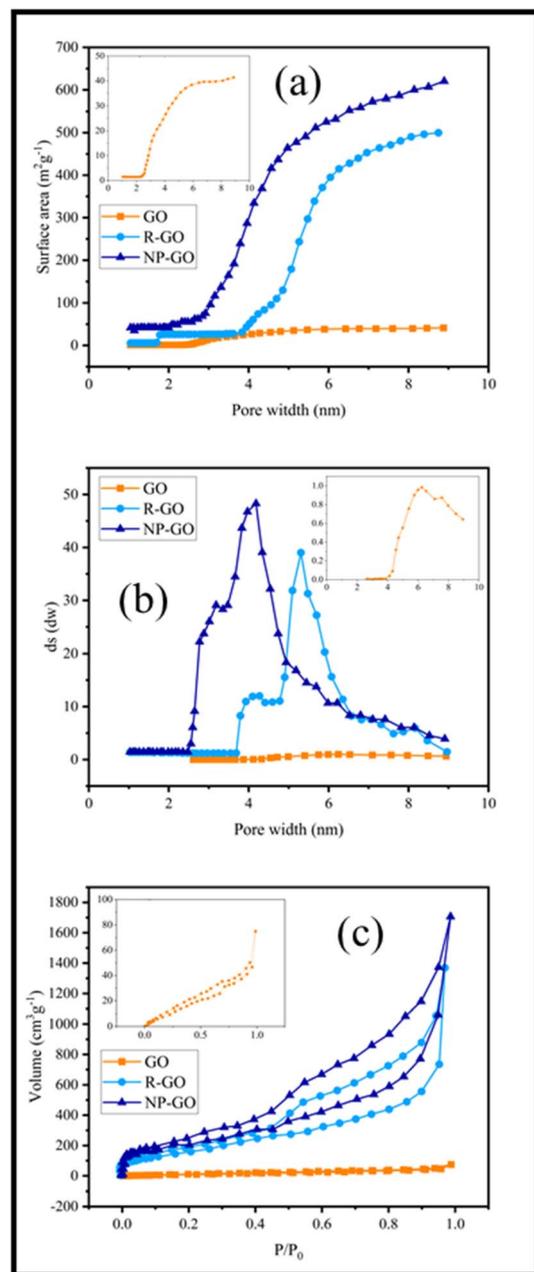


Fig. 5 (a) Surface area *versus* pore width in R-GO and NP-GO, (b) the distribution of pore size of R-GO and NP-GO, and (c)  $\text{N}_2$  adsorption isotherms in R-GO and NP-GO.



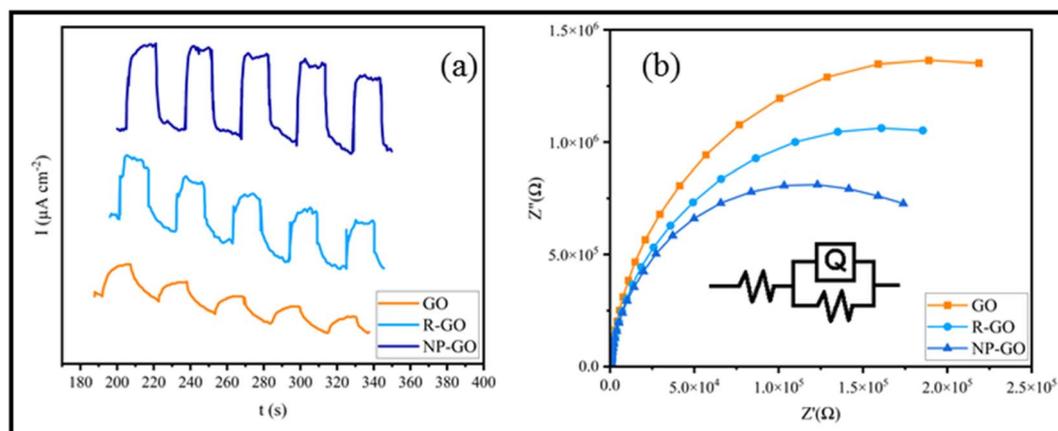


Fig. 6 (a) The photocurrent performances of GO, R-GO, and NP-GO and (b) EIS spectra of GO, R-GO, and NP-GO under simulated sunlight.

Table 2 Obtained EIS parameters by the model of  $R_s(R_{ct}Q_{ct})$  for samples

Samples	$R_s$ ( $\Omega$ cm $^2$ )	$R_{ct}$ (K $\Omega$ cm $^2$ )	$Q_{ct}$	
			$Y_0$ ( $\mu$ F cm $^{-2}$ )	$n$
GO	82.54	56.43	0.132	0.8952
R-GO	108.51	49.28	0.205	0.9601
NP-GO	89.68	39.78	0.263	0.9522

and charge transfer through the electrode/electrolyte boundary by forming covalent bonds among 2,4-dinitrophenylhydrazine groups and reduced graphene oxide

sheets. These results are in good agreement with the measurements of photocurrent. They indicate that photo-induced carriers become separate and more effective, and charges transfer more rapidly on the surface of NP-GO.

#### 4.4. Electrochemical measurements

Fig. 7a displays the cyclic voltammetry of the electrodes made of R-GO and NP-GO at a scan rate of  $5 \text{ mV s}^{-1}$ . For R-GO, an almost rectangular curve which is a characteristic of carbon materials, and a very small peak at  $-400 \text{ mV}_{\text{Ag}/\text{AgCl}}$  related to the reduction of unreduced oxide groups, can be seen.

In the curve related to NP-GO, there is an anodic peak in the range of  $-100 \text{ mV}_{\text{Ag}/\text{AgCl}}$  and a cathodic one in  $-300 \text{ mV}_{\text{Ag}/\text{AgCl}}$ .

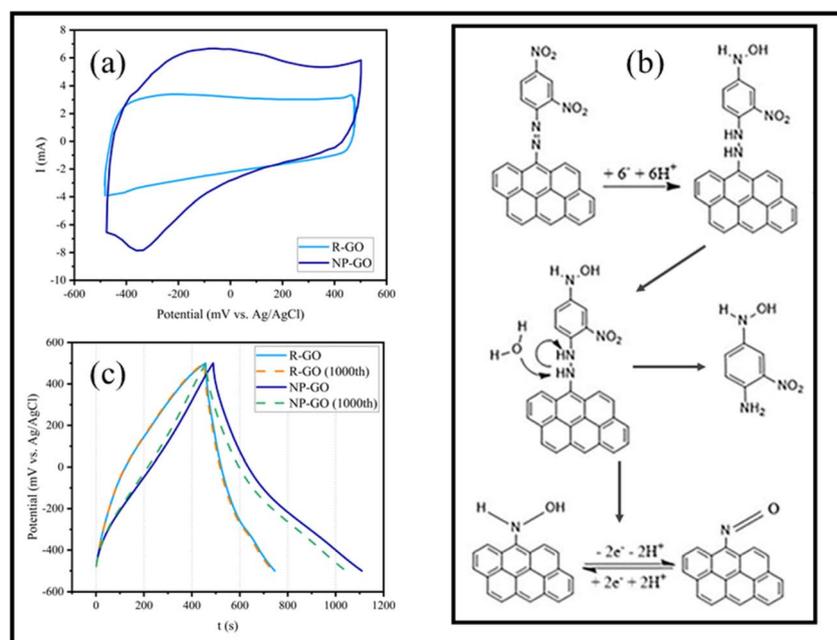


Fig. 7 (a) Cyclic voltammetry of electrodes made of R-GO and NP-GO at the scan rate of  $5 \text{ mV s}^{-1}$ , (b) the mechanism of the reduction of 2,4-dinitrophenylhydrazine presented on the surface of NP-GO in an aqueous environment and the redox reaction of the nitroso group, (c) First and 1000th galvanostatic charge/discharge of R-GO and NP-GO at  $0.5 \text{ A g}^{-1}$ .



Table 3 The reported capacitance of some graphene and nitrophenyl-functionalized graphene electrodes in neutral electrolytes

Electrode materials	Electrolyte	Current density or scan rate	Capacitance	Reference
R-GO, CNT, MnO <sub>2</sub>	Na <sub>2</sub> SO <sub>4</sub>	50 mV s <sup>-1</sup>	275 F g <sup>-1</sup>	88
R-GO, CNT, PVDF	Na <sub>2</sub> SO <sub>4</sub>	0.1 A g <sup>-1</sup>	129 F g <sup>-1</sup>	89
R-GO, CNT, MnO <sub>2</sub> , PTFE, acetylene black	Na <sub>2</sub> SO <sub>4</sub>	1 A g <sup>-1</sup>	120 F g <sup>-1</sup>	90
R-GO, CNT, MnO <sub>2</sub> , PTFE, carbon black	Na <sub>2</sub> SO <sub>4</sub>	0.5 A g <sup>-1</sup>	202 F g <sup>-1</sup>	91
Graphene foil produced by exfoliating graphite	Na <sub>2</sub> SO <sub>4</sub>	0.1 A g <sup>-1</sup>	106 F g <sup>-1</sup>	92
R-GO, PTFE, carbon black	Na <sub>2</sub> SO <sub>4</sub>	0.5 A g <sup>-1</sup>	144 F g <sup>-1</sup>	This work
NP-R-GO (diazonium salt) NP-GO	K <sub>2</sub> SO <sub>4</sub>	0.1 A g <sup>-1</sup>	315 F g <sup>-1</sup>	93
		0.1 A g <sup>-1</sup>	290 F g <sup>-1</sup>	
NP-GO	Na <sub>2</sub> SO <sub>4</sub>	0.5 A g <sup>-1</sup>	310 F g <sup>-1</sup>	This work

At the potential close to the initial reduction of NO<sub>2</sub> groups from the 4-position of the 2,4-dinitrophenylhydrazine and before the reduction at the 2-position, reduction of the azo bond occurs, separating 1-amino-2-nitro-4-nitrophenyl and forming aryl nitroso group on NP-GO<sup>86</sup> (Fig. 7b). The redox peak observed in the cyclic voltammetry of the NP-GO sample is related to the aryl nitroso groups' redox reaction.

Fig. 7c shows the results of the galvanostatic charge and discharge test related to R-GO and NP-GO electrodes in the initial cycle and after 1000 cycles at a current density of 0.5 A g<sup>-1</sup>. The discharging time of the electrodes made of R-GO and NP-GO are 290 and 620 seconds, respectively, and their respective specific capacities are 145 and 310 F g<sup>-1</sup>. Table 3 presents the reported capacities of some graphene-based electrodes and those made of functionalized graphene by nitrophenyl in neutral electrolytes. It can be seen that the prepared electrode materials show suitable performances compared to the literature. The results of this test show that R-GO has better stability than NP-GO so that after 1000 charge and discharge cycles, its capacity remains almost constant and equals 144 F g<sup>-1</sup>, while in the functionalized sample, the capacity has decreased by about 8% and reaches 291 F g<sup>-1</sup>. It should be noted that the high stability of carbon materials is one of the most important reasons for their use as an electrode material. For this reason, the higher stability of the R-GO electrode is not unconventional. However, the reduction in the capacity of the NP-GO electrode can be attributed to the reduction of the active surface or electrical conductivity due to the loss of layers or the breaking of the bond of the functional groups during the charge and discharge cycles.<sup>87</sup>

Fig. 8a shows the Nyquist diagram of R-GO and NP-GO electrodes. The obtained data were fitted using the Randles model as an equivalent electric circuit and their values are presented in Table 4. In this equivalent circuit,  $R_s$  represents the electrolyte resistance,  $R_{dl}$  is the charge transfer resistance,  $C_{dl}$  is the electric double-layer capacitance,  $Z_W$  is the Warburg impedance, and  $C_{ct}$  is the electrode's pseud-capacitance. It should be noted that lower  $R_{ct}$  and  $Z_W$  indicate lower charge transfer resistance and higher ion transfer rate.<sup>94</sup> The electrolyte resistance ( $R_{ct}$ ) in R-GO and NP-GO is 2.23, and 1.07  $\Omega$ , respectively, which suggests that the functionalization process leads to improved electrical conductivity and easy electron transfer. Establishing internal connections between the plain and functional groups in NP-GO can be the reason for that. In addition, the presence of  $Z_W$  with a higher slope in NP-GO than R-GO specifies better ion diffusion, and it can be the result of the existence of more active area in the morphology in this sample. It can also be seen in the Bode-phase diagrams (Fig. 8b) that the maximum value for phase changes is  $-73$  and  $-84$  for R-GO and NP-GO, respectively. Therefore, it can be concluded that NP-GO shows better capacitive features than R-GO due to its morphology and the presence of functional groups. The

Table 4 The obtained values of  $R_s$ ,  $R_{ct}$ , and  $Z_W$  from the electrical equivalent circuit

Sample	$R_s$ ( $\Omega$ )	$R_{dl}$ ( $\Omega$ )	$C_{dl}$ (F g <sup>-1</sup> )	$Z_W$	$C_{ct}$ (F g <sup>-1</sup> )
R-GO	4.46	2.46	$3.27 \times 10^{-4}$	0.016	0.0167
NP-GO	3.43	1.84	$6.53 \times 10^{-4}$	0.021	0.0639

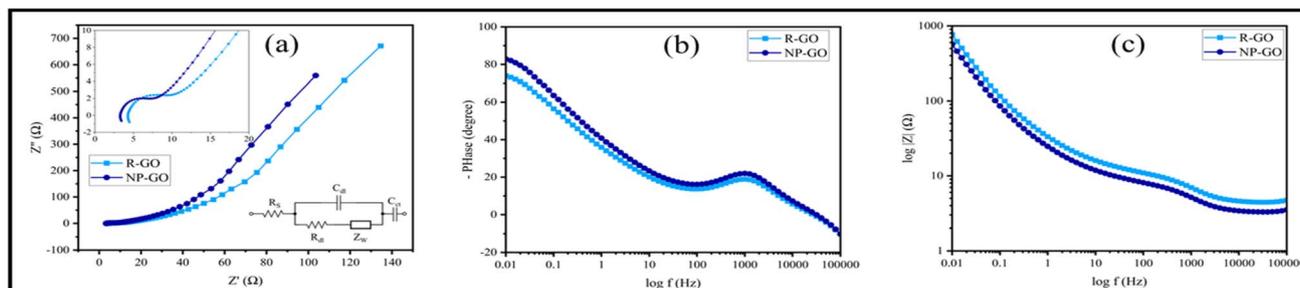


Fig. 8 (a) Nyquist plots of R-GO, NP-GO, and the electrical equivalent circuit, (b) Bode-phase diagrams, and (c) Bode-magnitude diagrams.



lower resistance of NP-GO compared to R-GO can be seen in the Bode-magnetite diagrams (Fig. 8c).

## 5. Conclusion

Graphene oxide was synthesized by the hummer method and *in situ* and simultaneously reduced and functionalized with 2,4-dinitrophenylhydrazine. Also, graphene is synthesized through the reaction of GO with hydrazine for comparing it with the functionalized sample. FTIR, Raman, TGA, and XRD of the samples were conducted to investigate the presence and effect of functional groups in samples. In addition, for further study, XPS of the samples was employed. According to the obtained results, due to the reaction with 2,4-dinitrophenylhydrazine, groups containing oxygen in GO are reduced and replaced with 2,4-dinitrophenylhydrazone groups. SEM and BET results display that the morphology of NP-GO has less agglomeration and, as a result, a more accessible active surface compared to graphene. The results of UV absorption show that shifts in absorption peaks of the NP-GO sample indicate the existence of electronic connections and  $\pi$ - $\pi^*$  interactions among the functional groups and reduced graphene oxide sheets. The photocurrent response of the sample exhibits that the separation/transportation of light-induced carriers was enhanced by replacing oxygen-containing groups with 2,4-dinitrophenylhydrazine groups. Galvanostatic charge and discharge show that the electrode made of NP-GO has a much higher capacity but lower stability compared to the electrode made of reduced graphene oxide. Based on the EIS results, the NP-GO electrode has a lower charge transfer resistance and better electrochemical performance than R-GO.

## Conflicts of interest

There are no conflicts of interest to declare.

## References

- W. Yu, L. Sisi, Y. Haiyan and L. Jie, *RSC Adv.*, 2020, **10**, 15328–15345.
- G. Yang, L. Li, W. B. Lee and M. C. Ng, *Sci. Technol. Adv. Mater.*, 2018, **19**, 613–648.
- X. Li, W. Cai, J. An, S. Kim, J. Nah, D. Yang, R. Piner, A. Velamakanni, I. Jung and E. Tutuc, *Science*, 2009, **324**, 1312–1314.
- J. Hass, W. De Heer and E. Conrad, *J. Phys.: Condens. Matter*, 2008, **20**, 323202.
- P. Avouris, M. Freitag and V. Perebeinos, *Nat. Photonics*, 2008, **2**, 341–350.
- P. Blake, E. Hill, A. Castro Neto, K. Novoselov, D. Jiang, R. Yang, T. Booth and A. Geim, *Appl. Phys. Lett.*, 2007, **91**, 063124.
- Z. Chen, Y. Qi, X. Chen, Y. Zhang and Z. Liu, *Adv. Mater.*, 2019, **31**, 1803639.
- K. Xia, W. Wu, M. Zhu, X. Shen, Z. Yin, H. Wang, S. Li, M. Zhang, H. Wang and H. Lu, *Sci. Bull.*, 2020, **65**, 343–349.

- Š. Meškiniš, A. Vasilias, A. Guobienė, M. Talaikis, G. Niaura and R. Gudaitis, *RSC Adv.*, 2022, **12**, 18759–18772.
- J.-Y. Moon, M. Kim, S.-I. Kim, S. Xu, J.-H. Choi, D. Whang, K. Watanabe, T. Taniguchi, D. S. Park and J. Seo, *Sci. Adv.*, 2020, **6**, eabc6601.
- Q. He, S. Wu, S. Gao, X. Cao, Z. Yin, H. Li, P. Chen and H. Zhang, *ACS Nano*, 2011, **5**, 5038–5044.
- X. Qi, K. Y. Pu, H. Li, X. Zhou, S. Wu, Q. L. Fan, B. Liu, F. Boey, W. Huang and H. Zhang, *Angew. Chem., Int. Ed.*, 2010, **49**, 9426–9429.
- H. G. Sudibya, Q. He, H. Zhang and P. Chen, *ACS Nano*, 2011, **5**, 1990–1994.
- X. Cao, Q. He, W. Shi, B. Li, Z. Zeng, Y. Shi, Q. Yan and H. Zhang, *Small*, 2011, **7**, 1199–1202.
- W. Gao, L. B. Alemany, L. Ci and P. M. Ajayan, *Nat. Chem.*, 2009, **1**, 403–408.
- A. Lerf, H. He, M. Forster and J. Klinowski, *J. Phys. Chem. B*, 1998, **102**, 4477–4482.
- X. Lv, Y. Huang, Z. Liu, J. Tian, Y. Wang, Y. Ma, J. Liang, S. Fu, X. Wan and Y. Chen, *Small*, 2009, **5**, 1682–1687.
- L. Zhang, X. Li, Y. Huang, Y. Ma, X. Wan and Y. Chen, *Carbon*, 2010, **48**, 2367–2371.
- X. Fan, W. Peng, Y. Li, X. Li, S. Wang, G. Zhang and F. Zhang, *Adv. Mater.*, 2008, **20**, 4490–4493.
- V. Dua, S. P. Surwade, S. Ammu, S. R. Agnihotra, S. Jain, K. E. Roberts, S. Park, R. S. Ruoff and S. K. Manohar, *Angew. Chem., Int. Ed.*, 2010, **49**, 2154–2157.
- S. Sravya, D. RamaDevi, N. Belachew, K. E. Rao and K. Basavaiah, *RSC Adv.*, 2021, **11**, 12030–12035.
- J. Liu, S. Fu, B. Yuan, Y. Li and Z. Deng, *J. Am. Chem. Soc.*, 2010, **132**, 7279–7281.
- A. Baral, L. Satish, D. P. Das, H. Sahoo and M. K. Ghosh, *J. Biomol. Struct. Dyn.*, 2020, **38**, 2038–2046.
- H.-L. Guo, X.-F. Wang, Q.-Y. Qian, F.-B. Wang and X.-H. Xia, *ACS Nano*, 2009, **3**, 2653–2659.
- Z. Wang, X. Zhou, J. Zhang, F. Boey and H. Zhang, *J. Phys. Chem. C*, 2009, **113**, 14071–14075.
- X. Huang, X. Zhou, S. Wu, Y. Wei, X. Qi, J. Zhang, F. Boey and H. Zhang, *Small*, 2010, **6**, 513–516.
- X. H. Tai, S. W. Chook, C. W. Lai, K. M. Lee, T. C. K. Yang, S. Chong and J. C. Juan, *RSC Adv.*, 2019, **9**, 18076–18086.
- W. Chen and L. Yan, *Nanoscale*, 2010, **2**, 559–563.
- C. Mattevi, *Adv. Funct. Mater.*, 2009, **19**, 2577–2583.
- A. Amiri, M. Baghayeri, F. Karimabadi, F. Ghaemi and B. Maleki, *Microchim. Acta*, 2020, **187**, 1–8.
- H. Kumar, R. Sharma, A. Yadav and R. Kumari, *J. Energy Storage*, 2021, **33**, 102032.
- R. Tarcan, O. Todor-Boer, I. Petrovai, C. Leordean, S. Astilean and I. Botiz, *J. Mater. Chem. C*, 2020, **8**, 1198–1224.
- P. Sandhya, J. Jose, M. Sreekala, M. Padmanabhan, N. Kalarikkal and S. Thomas, *Ceram. Int.*, 2018, **44**, 15092–15098.
- S. M. Majhi, A. Mirzaei, H. W. Kim and S. S. Kim, *Sensors*, 2021, **21**, 1352.
- A. P. Straub, D. S. Bergsman, B. A. Getachew, L. M. Leahy, J. J. Patil, N. Ferralis and J. C. Grossman, *Nano Lett.*, 2021, **21**, 2429–2435.



- 36 M. Tarahomi, H. Alinezhad and B. Maleki, *Appl. Organomet. Chem.*, 2019, **33**, e5203.
- 37 H. Boroumand, H. Alinezhad, B. Maleki and S. Peiman, *Polycyclic Aromat. Compd.*, 2022, 1–17.
- 38 G. Eda, Y. Y. Lin, C. Mattevi, H. Yamaguchi, H. A. Chen, I. S. Chen, C. W. Chen and M. Chhowalla, *Adv. Mater.*, 2010, **22**, 505–509.
- 39 A. Hashimoto, K. Suenaga, A. Gloter, K. Urita and S. Iijima, *nature*, 2004, **430**, 870–873.
- 40 S. Stankovich, R. D. Piner, X. Chen, N. Wu, S. T. Nguyen and R. S. Ruoff, *J. Mater. Chem.*, 2006, **16**, 155–158.
- 41 D. Li, M. B. Müller, S. Gilje, R. B. Kaner and G. G. Wallace, *Nat. Nanotechnol.*, 2008, **3**, 101–105.
- 42 S. Park, J. An, I. Jung, R. D. Piner, S. J. An, X. Li, A. Velamakanni and R. S. Ruoff, *Nano Lett.*, 2009, **9**, 1593–1597.
- 43 S. Park, J. An, R. D. Piner, I. Jung, D. Yang, A. Velamakanni, S. T. Nguyen and R. S. Ruoff, *Chem. Mater.*, 2008, **20**, 6592–6594.
- 44 T. A. Strom, E. P. Dillon, C. E. Hamilton and A. R. Barron, *Chem. Commun.*, 2010, **46**, 4097–4099.
- 45 X. Wu, H. Cao, B. Li and G. Yin, *Nanotechnology*, 2011, **22**, 075202.
- 46 H.-P. Cong, X.-C. Ren, P. Wang and S.-H. Yu, *Energy Environ. Sci.*, 2013, **6**, 1185–1191.
- 47 G. Pognon, C. Cougnon, D. Mayilukila and D. Bélanger, *ACS Appl. Mater. Interfaces*, 2012, **4**, 3788–3796.
- 48 W. Ai, W. Zhou, Z. Du, Y. Du, H. Zhang, X. Jia, L. Xie, M. Yi, T. Yu and W. Huang, *J. Mater. Chem.*, 2012, **22**, 23439–23446.
- 49 H. Bai, C. Li, X. Wang and G. Shi, *Chem. Commun.*, 2010, **46**, 2376–2378.
- 50 S. Liu, J. Tian, L. Wang, H. Li, Y. Zhang and X. Sun, *Macromolecules*, 2010, **43**, 10078–10083.
- 51 Q. Su, S. Pang, V. Alijani, C. Li, X. Feng and K. Müllen, *Adv. Mater.*, 2009, **21**, 3191–3195.
- 52 W. Tu, J. Lei, S. Zhang and H. Ju, *Chem. - Eur. J.*, 2010, **16**, 10771–10777.
- 53 D. K. Samarakoon, Z. Chen, C. Nicolas and X. Q. Wang, *Small*, 2011, **7**, 965–969.
- 54 D. K. Samarakoon and X.-Q. Wang, *ACS Nano*, 2009, **3**, 4017–4022.
- 55 D. K. Samarakoon and X.-Q. Wang, *ACS Nano*, 2010, **4**, 4126–4130.
- 56 U. K. Wijewardena, S. E. Brown and X.-Q. Wang, *J. Phys. Chem. C*, 2016, **120**, 22739–22743.
- 57 F. Chen, Q. Qing, J. Xia, J. Li and N. Tao, *J. Am. Chem. Soc.*, 2009, **131**, 9908–9909.
- 58 E. Paek, A. J. Pak, K. E. Kweon and G. S. Hwang, *J. Phys. Chem. C*, 2013, **117**, 5610–5616.
- 59 M. Mousavi-Khoshdeld, E. Targholi and M. J. Momeni, *J. Phys. Chem. C*, 2015, **119**, 26290–26295.
- 60 S. M. Mousavi-Khoshdeld and E. Targholi, *Carbon*, 2015, **89**, 148–160.
- 61 R. N. Gunasinghe, D. G. Reuven, K. Suggs and X.-Q. Wang, *J. Phys. Chem. Lett.*, 2012, **3**, 3048–3052.
- 62 T. R. Nanayakkara, U. K. Wijewardena, A. B. Arampath, K. Suggs, N. Ravi and X.-Q. Wang, in *Theoretical and Computational Chemistry*, Elsevier, 2022, vol. 21, pp. 131–155.
- 63 X. Tian, J. Gu and J.-b. Xu, *J. Chem. Phys.*, 2014, **140**, 044712.
- 64 Y. Feng, H. Liu, W. Luo, E. Liu, N. Zhao, K. Yoshino and W. Feng, *Sci. Rep.*, 2013, **3**, 1–8.
- 65 E. Bekyarova, S. Sarkar, S. Niyogi, M. Itkis and R. Haddon, *J. Phys. D: Appl. Phys.*, 2012, **45**, 154009.
- 66 J. R. Lomeda, C. D. Doyle, D. V. Kosynkin, W.-F. Hwang and J. M. Tour, *J. Am. Chem. Soc.*, 2008, **130**, 16201–16206.
- 67 C. Bosch-Navarro, E. Coronado, C. Martí-Gastaldo, J. Sánchez-Royo and M. G. Gómez, *Nanoscale*, 2012, **4**, 3977–3982.
- 68 K. Krishnamoorthy, M. Veerapandian, K. Yun and S.-J. Kim, *Carbon*, 2013, **53**, 38–49.
- 69 Y. Liu, Y. Shi, W. Zhou, W. Shi, W. Dang, X. Li and B. Liang, *Opt. Laser Technol.*, 2021, **139**, 106960.
- 70 B. Vinayan, R. Nagar, V. Raman, N. Rajalakshmi, K. Dhathathreyan and S. Ramaprabhu, *J. Mater. Chem.*, 2012, **22**, 9949–9956.
- 71 B. Rajagopalan and J. S. Chung, *Nanoscale Res. Lett.*, 2014, **9**, 1–10.
- 72 H. D. Le, T. T. T. Ngo, D. Q. Le, X. N. Nguyen and N. M. Phan, *Adv. Nat. Sci.: Nanosci. Nanotechnol.*, 2013, **4**, 035012.
- 73 F. T. Johra, J.-W. Lee and W.-G. Jung, *J. Ind. Eng. Chem.*, 2014, **20**, 2883–2887.
- 74 K. Roodenko, M. Gensch, J. Rappich, K. Hinrichs, N. Esser and R. Hunger, *J. Phys. Chem. B*, 2007, **111**, 7541–7549.
- 75 V. Kumar, R. K. Gupta, R. K. Gundampati, D. K. Singh, S. Mohan, S. H. Hasan and M. Malviya, *RSC Adv.*, 2018, **8**, 619–631.
- 76 Y. Zhou, Q. Bao, L. A. L. Tang, Y. Zhong and K. P. Loh, *Chem. Mater.*, 2009, **21**, 2950–2956.
- 77 K. Bijudas, *J. Appl. Chem.*, 2019, **8**, 1887–1892.
- 78 M. D. Stoller, S. Park, Y. Zhu, J. An and R. S. Ruoff, *Nano Lett.*, 2008, **8**, 3498–3502.
- 79 A. Le Comte, D. Chhin, A. Gagnon, R. Retoux, T. Brousse and D. Bélanger, *J. Mater. Chem. A*, 2015, **3**, 6146–6156.
- 80 A. Gambou-Bosca and D. Bélanger, *J. Electrochem. Soc.*, 2015, **162**, A5115.
- 81 T. Ohkubo, J. Miyawaki, K. Kaneko, R. Ryoo and N. A. Seaton, *J. Phys. Chem. B*, 2002, **106**, 6523–6528.
- 82 W. Zhao, C. Li, A. Wang, C. Lv, W. Zhu, S. Dou, Q. Wang and Q. Zhong, *Phys. Chem. Chem. Phys.*, 2017, **19**, 28696–28709.
- 83 W. Zhao, A. Wang, Y. Wang, C. Lv, W. Zhu, S. Dou, Q. Wang and Q. Zhong, *J. Alloys Compd.*, 2017, **726**, 164–172.
- 84 J. Zhang, A. Wang, W. Zhao, C. Li, X. Chen, Y. Wang, W. Zhu and Q. Zhong, *Dyes Pigm.*, 2018, **153**, 241–247.
- 85 J. Yang, J. Wu, C. Zhang, S. Zhang, B. Yang, W. Emori and J. Wang, *J. Alloys Compd.*, 2020, **819**, 152943.
- 86 C. A. Thorogood, G. G. Wildgoose, J. H. Jones and R. G. Compton, *New J. Chem.*, 2007, **31**, 958–965.
- 87 L. Zhang, K. Hui, K.-S. Hui, X. Chen, R. Chen and H. Lee, *Int. J. Hydrogen Energy*, 2016, **41**, 9443–9453.
- 88 Y. Cheng, S. Lu, H. Zhang, C. V. Varanasi and J. Liu, *Nano Lett.*, 2012, **12**, 4206–4211.
- 89 O. Okhay, A. Tkach, P. Staiti and F. Lufrano, *Electrochim. Acta*, 2020, **353**, 136540.



Paper

- 90 L. Deng, Z. Hao, J. Wang, G. Zhu, L. Kang, Z.-H. Liu, Z. Yang and Z. Wang, *Electrochim. Acta*, 2013, **89**, 191–198.
- 91 B. Ding, D. Guo, Y. Wang, X. Wu and Z. Fan, *J. Power Sources*, 2018, **398**, 113–119.
- 92 A. Ambrosi and M. Pumera, *Chem. - Eur. J.*, 2016, **22**, 153–159.
- 93 S. Alipour and S. M. Mousavi-Khoshdel, *Electrochim. Acta*, 2019, **317**, 301–311.
- 94 Y. Zhu, W. Chu, N. Wang, T. Lin, W. Yang, J. Wen and X. Zhao, *RSC Adv.*, 2015, **5**, 77958–77964.

