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# *In situ* generation of exfoliated graphene layers on recycled graphite rods for enhanced capacitive performance of Ni–Co binary hydroxide

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A functionalized exfoliated graphite rod (FEGR), with a high surface area, is produced for use as a promising substrate for supercapacitors, *via* controlled oxidative treatment of a recycled graphite rod of exhausted zinc–carbon batteries. SEM, EDX, XPS, FT-IR, Raman, and contact angle measurements are carried out to disclose the surface characteristics of the FEGR. The surface of the FEGR is characterized by *in situ* generated grooves, together with graphene layers which are directly attached to the underlying graphite base. The FEGR electrodes enhance the capacitive performance of Ni(OH)<sub>2</sub> and binary Ni–Co(OH)<sub>2</sub>. The Ni–Co(OH)<sub>2</sub>/FEGR electrode displays a superb specific capacity value (2552.6 C g<sup>−1</sup>) at a current density of 5 A g<sup>−1</sup> and this value is retained to 70.8% at a high current density of 50 A g<sup>−1</sup> indicating the outstanding rate performance of this electrode material. This enhanced behavior is attributed to the facile interaction of electrolyte species, even at high current density, with the active sites of the redox catalyst layer (distributed over a larger fraction of the underlying substrate with enhanced hydrophilicity). Moreover, the excellent electrical conductivity of the *in situ* surface generated graphene layers is another promoting factor.

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## 1. Introduction

Nowadays, energy storage technology has attracted great attention in research due to the increase of various renewable energy sources in the world to cover the rapid decrease in fossil fuels. Also, increasing the development in many applications *e.g.*, mobile phones, energy recovery systems, electric vehicles and electric tramcar, energy harvesting, and uninterruptible power supply requires efficient energy storage devices.<sup>1–5</sup>

Supercapacitors (SCs) are among the most efficient energy storage devices. Recently, SCs have received considerable attention for storing energy due to their unique advantages such as high-power density, fast charge–discharge rate, long-life cycles, and being eco-friendly.<sup>6,7</sup>

SCs storage mechanisms can be classified into two main types according to the main nature of the electrode material; non-faradic and faradic electrodes. Also, they can be categorized into four types (electrochemical double-layer capacitors (EDLCs), pseudocapacitive, battery, and recently intercalation pseudocapacitive electrodes) according to the main nature of the electrode material, potential evaluation profile during charge and discharge, and the reversibility of the electrode material.<sup>8–10</sup> The EDLCs store energy through the accumulation

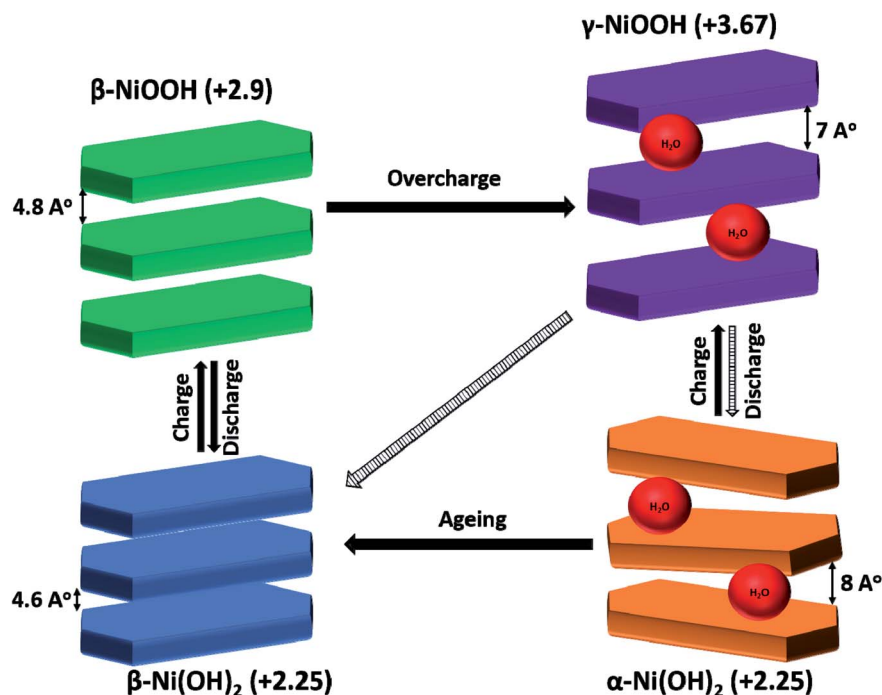
of the electrolyte ions on the accessible surface area of electrode materials *via* electrostatic attraction. Unfortunately, this type of electrode material has a low energy density which is used only in the low energy applications.<sup>8</sup> So, the research is directed to investigate the battery type electrode in SCs devices which stores energy chemically *via* faradaic reactions that take place at the electrode material at characteristic potentials giving higher energy density than EDLCs electrode materials.<sup>8,10,11</sup>

Interestingly, Ni(OH)<sub>2</sub> is one of the most investigated battery-type materials in the SCs devices due to their inspired properties; high theoretical specific capacity (*C*) (1728 C g<sup>−1</sup>), being environmentally friendly, natural abundance, and low cost which make it available for commercialization.<sup>12–14</sup> It is worth noting that Ni(OH)<sub>2</sub> has two polymorphic crystal structures,  $\alpha$ -Ni(OH)<sub>2</sub> phase with hydrotalcite-like structure and  $\beta$ -Ni(OH)<sub>2</sub> with brucite-like structure (see Scheme 1). The former has a higher capacitive performance than the latter because of  $\alpha$ -Ni(OH)<sub>2</sub> has a higher *d*-spacing value with the turbostratically crystallized structure which facilitates the diffusion of electrolyte inside the hydroxide layers and utilize almost active materials with a higher rate than  $\beta$ -Ni(OH)<sub>2</sub>. So,  $\alpha$ -Ni(OH)<sub>2</sub> has a higher *C* value, rate capability, and power density.<sup>15,16</sup> Besides, the energy resulting from the conversion of  $\alpha$ -Ni(OH)<sub>2</sub> to  $\gamma$ -NiOOH during the charging process is greater than that resulting from the conversion of  $\beta$ -Ni(OH)<sub>2</sub> to  $\beta$ -NiOOH. This is due to the difference between the average oxidation state between  $\gamma$ -NiOOH (+3.67) and  $\alpha$ -Ni(OH)<sub>2</sub> (+2.25) compared with

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Scheme 1 Schematic illustration of  $\text{Ni(OH)}_2/\text{NiOOH}$  phase transformation (Bode diagram).<sup>42</sup>

the difference from  $\beta\text{-NiOOH}$  (+2.9) to  $\beta\text{-Ni(OH)}_2$  (+2.25).<sup>15,17</sup> Despite the high theoretical  $C$  and high capacitive properties of  $\text{Ni(OH)}_2$ , these properties cannot be achieved practically due to its low electrical conductivity ( $\sim 10^{-17} \text{ S cm}^{-1}$ ) and low ionic conductivity.<sup>14,18,19</sup> Also,  $\text{Ni(OH)}_2$  has low structure stability in alkaline media due to mechanical stresses during cycling.<sup>17,18</sup> So, researchers directed their efforts to discover strategies to overcome these obstacles. One of these strategies is to develop a bimetallic structure by substituting Ni with some metals such as Al, Zn, Mn, and Co.<sup>15,20</sup> Importantly, Ni-Co bimetallic hydroxides are among the well-known binary hydroxides in SCs field. Ni and Co atoms have a very close ionic size which enables their easy substitution during the synthesizing process of the binary hydroxide.<sup>21–25</sup>

The coexistence of cobalt in the crystal lattice of  $\text{Ni(OH)}_2$  displayed many advantages. Firstly, the total electronic conductivity of Ni-Co binary hydroxide is increased which in turn increases the  $C$  and rate capability.<sup>8,26,27</sup> Jian Wu *et al.*<sup>28</sup> demonstrated that the incorporation of Co into Ni lattice decreases the band gap between valence and conduction band by quantum calculations using density functional theory. The four prepared samples  $\text{Ni(OH)}_2$ ,  $\text{Co}_{0.33}\text{Ni}_{0.77}(\text{OH})_2$ ,  $\text{Co}_{0.5}\text{Ni}_{0.5}(\text{OH})_2$ , and  $\text{Co}_{0.77}\text{Ni}_{0.33}(\text{OH})_2$  displayed band gap 3.43 eV, 3.13 eV, 3.06 eV, and 3.01 eV, respectively.<sup>28</sup>

Secondly, the binary hydroxides had higher structural stability than individual hydroxides due to Co metal preventing mechanical failure of  $\text{Ni(OH)}_2$  during cycling by two approaches. The first approach is enhancing the stability of  $\beta\text{-Ni(OH)}_2$  by arresting the formation of  $\gamma\text{-NiOOH}$  from  $\beta\text{-NiOOH}$  by lowering the oxidation potential which prevents the overcharging of  $\beta\text{-NiOOH}$  to  $\gamma\text{-NiOOH}$ .<sup>20,29</sup> The second approach is

improving the stability of  $\alpha\text{-Ni(OH)}_2$  by inhibiting the transformation of  $\gamma\text{-NiOOH}$  to  $\beta\text{-Ni(OH)}_2$  during the discharge process.<sup>15,23</sup>

On the other hand, the co-existence of Co into  $\text{Ni(OH)}_2$  crystal lattice changes the morphology of electrode material that results in reinforcing the kinetics of the electrode. Firstly, by increasing the rate of diffusion of electrolyte ions inside the active materials due to the unique nano-porous structure that results by mixing two metals during the preparation process such as nanosheets,<sup>30</sup> nanoarrays,<sup>31</sup> flower-like morphology.<sup>32</sup> Secondly, by decreasing the charge transfer resistance of  $\text{Ni(OH)}_2$ .<sup>33</sup> But, the kinetics of the binary hydroxide is still far enough than EDLCs electrode materials. So, the research efforts are directed to investigate the substrate materials which are characterized by high electrical conductivity to overcome this problem.<sup>34</sup>

Among various substrate materials, graphene is considered as the pioneer support for the redox-active materials due to its ultra-high electrical conductivity, very high charge carrier mobility ( $\sim 200\,000 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ ), and huge gravimetric surface area ( $\sim 2600 \text{ m}^2 \text{ g}^{-1}$ ).<sup>35,36</sup> So by incorporation of the active materials (*e.g.* metal hydroxide or oxide) on the graphene layers, the kinetics and the capacitive performance of the electrode is boosted largely due to: (i) the active materials become accessible to interact with the electrolyte ions because of the uniform distributions of the active materials on the huge surface area of graphene, (*i.e.* decreasing of ionic diffusion pathways), and (ii) decreasing the electronic pathways.<sup>37</sup> Also, graphene can be produced *via* many synthetic methods, *e.g.* micromechanical exfoliation, thermal decomposition of SiC, chemical vapor deposition (CVD), chemical exfoliation, and electrochemical



exfoliation.<sup>35</sup> Electrochemical partial exfoliation is considered the best method due to the following characteristics: (a) it is an available method that carried out in one simple step without using harsh materials,<sup>38,39</sup> (b) it is a controllable method because the degree of oxidation can be controlled by oxidation potential or current, time of oxidation, and type and concentration of the electrolyte,<sup>40</sup> (c) it is a low cost and eco-friendly method,<sup>39</sup> (d) it is a binder-free method due to the partially exfoliated graphite is consisting of two layers. While the bottom layer is composed of the electrically conductive graphite which serves as a current collector, the upper layer is formed of the functionalized graphene sheets as active material,<sup>37</sup> (e) the resulting three dimensional (3D) highly conductive material with a very high surface-functionalized area possesses a high specific capacitance ( $C_s$ ) due to ease interaction between the electrolyte and active material,<sup>41</sup> and (f) the highly electrical 3D conductive network can be used as a support to accommodate the redox-active materials to enhance the electrical conductivity of redox materials.<sup>37</sup>

The objective of this study is to synthesize the graphene layers as a support material for the Ni-Co(OH)<sub>2</sub> from the low-cost carbon source (exhausted zinc-carbon battery rods) with simple, controllable, and eco-friendly methods. Graphene layers are successfully produced from the recycled graphite rod obtained from the exhausted zinc-carbon batteries with a nominal voltage of 1.5 V using a simple oxidation step. The developed binary Ni-Co(OH)<sub>2</sub> supported on the functionalized exfoliated graphite rods (FEGR) displays a higher capacitive performance with an ultra-high  $C$  reaches to 2552.6 C g<sup>-1</sup> at 5 A g<sup>-1</sup> that retained to 70.8% at 50 A g<sup>-1</sup>.

## 2. Experimental

### 2.1. Chemicals and electrodes preparation

**2.1.1. Chemicals.** All the used chemicals were of analytical grade (purchased from Sigma Aldrich and Merck) and were used as received without any further purification. All solutions were prepared using second distilled water.

**2.1.2. Electrode preparation.** (i) Preparation of the RGR and FEGR electrodes

Recycled graphite rod (RGR) with 4 mm diameter is obtained from the exhausted zinc-carbon batteries with nominal voltage 1.5 V. RGR is cleaned by the following procedure: removing the large residual particles on the surface *via* mechanical polishing using very smooth emery paper and then is immersed in the boiling distilled water for 3 min with continuous stirring to remove any fine residual particles on its surface.

The functionalized exfoliated graphite rod (FEGR) is prepared in the three-electrode configuration cell [RGR served as a working electrode (immersed length 1 cm), graphite rod and saturated silver electrode (Ag/AgCl/KCl sat.) are used as the counter and reference electrodes, respectively] by a simple electrochemical anodic oxidation step *via* applying an oxidation potential equal to +2 V and for 10 min in a solution containing 1 M H<sub>2</sub>SO<sub>4</sub>.

(ii) Preparation of modified RGR and FEGR electrodes

Two Ni(OH)<sub>2</sub> films are prepared, *i.e.*, RGR/Ni(OH)<sub>2</sub>, and FEGR/Ni(OH)<sub>2</sub> by cathodic reduction at deposition potential of -1 V with a controlled amount of charge equal to 200 mC to pass in 20 mM NiSO<sub>4</sub> deposition bath using RGR or FEGR, respectively, as a working electrode for the two films. While graphite rod and (Ag/AgCl/KCl sat.) are used as the counter and reference electrodes, respectively. FEGR-Ni-Co(OH)<sub>2</sub> film is prepared by using sequential deposition of 100 mC of Co(OH)<sub>2</sub> from deposition bath containing [20 mM CoSO<sub>4</sub> + 20 mM KIO<sub>3</sub>] then depositing 100 mC of Ni(OH)<sub>2</sub> from 20 mM NiSO<sub>4</sub> solution with the same three-electrode configuration cell of the FEGR/Ni(OH)<sub>2</sub> film.<sup>43</sup>

### 2.2. Material characterization

Surface morphology and chemical composition of the various modified electrodes are examined by field emission scanning electron microscope (FE-SEM, QUANTA FEG 250) coupled with an energy dispersive X-ray spectrometer (EDX) unit. X-ray photoelectron spectroscopy (XPS), using monochromatic X-ray Al K-alpha radiation, Thermo Fisher Scientific. Additionally, the crystalline structures are characterized using X-ray diffraction (XRD) with Cu K $\alpha$  radiation, STOE STADI. Moreover, surface wettability is probed by contact angle measurements *via* Attention Biolin Scientific (Version 2.7). While Raman spectra are recorded by LabRAM HR Evolution and are used to follow the degree of the crystallinity of the graphite material before and after the electrochemical exfoliation process. Besides, the Fourier transform infrared (FT-IR) spectra are recorded with an FT-IR spectrometer (Nicolet series 670-FTIR) to investigate the surface functional groups.

### 2.3. Electrochemical measurements

The electrochemical measurements were investigated under a three-electrode cell configuration at room temperature (25 °C  $\pm$  1) in 1 M H<sub>2</sub>SO<sub>4</sub> for RGR and FEGR electrodes and 0.5 M NaOH for modified RGR and FEGR electrodes. The cyclic voltammetry (CV), galvanostatic charge-discharge (GCD), and electrochemical impedance spectroscopy (EIS) (at open circuit potential (OCP) from 100 kHz to 10 mHz) measurements are performed using Biologic potentiostat (model VSP-300).

The  $C_s$  was calculated from the CV and GCD curves using eqn (1) and (2), respectively and  $C$  was calculated using eqn (3):

$$C_s = \frac{\int Idt}{S\Delta V} = \frac{\int IdV}{Sv\Delta V} \quad (1)$$

$$C_s = \frac{I\Delta t}{S\Delta V} \quad (2)$$

$$C = \frac{I\Delta t}{S} \quad (3)$$

where  $C_s$  is the specific capacitance in F g<sup>-1</sup> when the  $S$  is the mass of the active material in the electrode in g (which is estimated from Faraday's law assuming 100% efficiency) or in the unit of F cm<sup>-2</sup> when  $S$  is the area of the electrode in cm<sup>2</sup>,  $\int IdV$  is the integrated oxidation or reduction current-potential area,  $v$  is the scan rate (V s<sup>-1</sup>),  $I(A)$  is the discharge current,  $\Delta t(s)$  is the

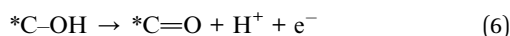
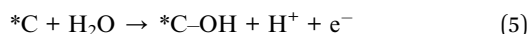
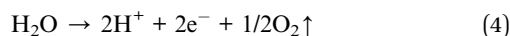


discharge time,  $\Delta V(V)$  is the operating potential window,  $C$  is specific capacity in unit  $C\ g^{-1}$  or  $C\ cm^{-2}$ .<sup>6,12,43</sup>

### 3. Results and discussion

#### 3.1. Modification of the substrate

**3.1.1. Mechanism of graphene synthesis by using partially electrochemical exfoliation method.** Synthesis of functionalized graphene layers was done by using a simple electrochemical oxidation step done on RGR (grey rod with smooth surface) with an oxidation potential of 2 V for 10 min as shown in Scheme 2. Firstly, the applied potential is larger than the water oxidation potential which stimulates the large evolution of oxygen bubbles (eqn (4)) which in turn makes defects in the entire external surface of RGR that simplify the insertion of water and sulfate anions ( $SO_4^{2-}$ ) inside the RGR layers.<sup>44,45</sup> The inserted water molecules and  $SO_4^{2-}$  anions produce the  $O_2$  and  $SO_2$  gases inside the graphite layers due to the  $H_2O$  molecules' self-oxidation and  $SO_4^{2-}$  anions reduction, respectively. These gases force the graphite layers to overcome the van der Waals forces and expand away from each other to yield the exfoliated graphene layers.<sup>44,46</sup> Also, the inserted  $H_2O$  molecules and  $SO_4^{2-}$  anions are modifying the graphene layers during the expansion by functionalizing its surface with hydroxyl ( $*C-OH$ ), carbonyl ( $*C=O$ ), and carboxylic ( $*COOH$ ) groups as shown in eqn (5)–(7).<sup>44,47</sup>

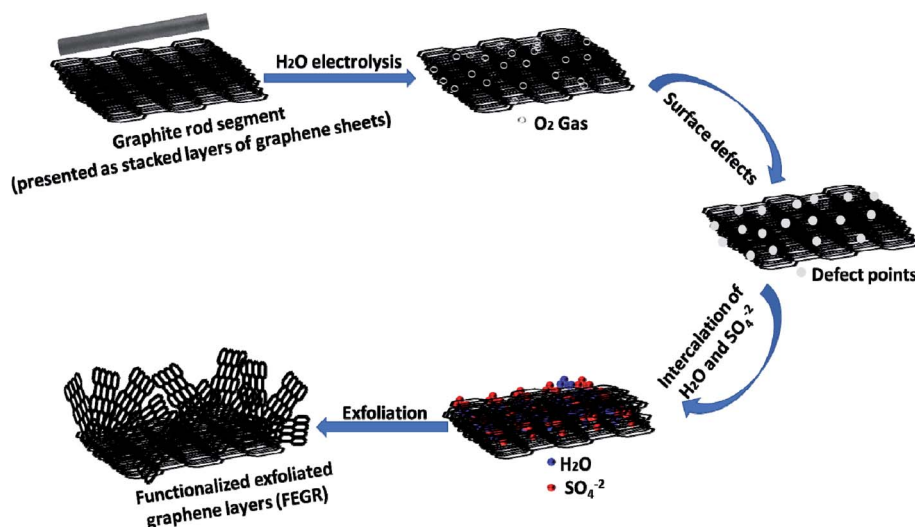


**3.1.2. Physical characterization of the RGR and FEGR.** The surface morphology of RGR and FEGR was inspected using SEM

analysis. The surface of the RGR displayed two different regions. The first one consists of a smooth surface with the presence of small wrinkles as clear from Fig. 1A. While the second region indicates the presence of some defects in the surface which is composed of densely stacked graphite layers as shown in Fig. 1B. On the other hand, the entire external surface of FEGR is fully oxidized and defected with the formation of interconnected graphene layers on the top surface of RGR that is supported by the internal surface of RGR as displayed in Fig. 1C. The synthesized graphene sheets with the open porous structure facilitate the insertion of electrolyte ions by lowering the diffusion pathways indicating the better electrochemical performance of this electrode in comparison with RGR, see Fig. 4(A–F). Not only the characteristic morphology is the main reason behind enhancing the electrochemical performance, but also the intensely inserted oxygen functional groups in the FEGR play an important role.

The EDX analysis was performed (see Fig. 1(D and E)) to show the chemical composition of the prepared FEGR and to display the change of oxygen content between RGR and FEGR electrodes due to the inserted functional groups during the electrooxidation. As displayed in Fig. 1D, the chemical composition of RGR is composed mainly of pure graphitic carbon. EDX analysis of FEGR displays a presence of small oxygen peaks (mass percent 14.5%) which indicate the successful insertion of oxygen functional groups on its surface. Also, the presence of the small sulfur peak in the EDX analysis of FEGR is evidence of the intercalation of  $SO_4^{2-}$  anions during the exfoliation of graphene layers.<sup>44</sup>

Also, XPS was done to investigate the accurate oxygen content and the type of inserted functional groups after the modification. From the XPS survey spectra, as shown in Fig. 2A, there are two main peaks centered around 285 and 532 eV corresponding to carbon and oxygen elements which indicate that the two electrodes are composed only of carbon and oxygen.<sup>46,48</sup> By comparing the enlarged view of C 1s peaks of RGR and FEGR that depicted in Fig. 2B, the presence of the



Scheme 2 Schematic illustration of the partial electrochemical exfoliation of RGR.





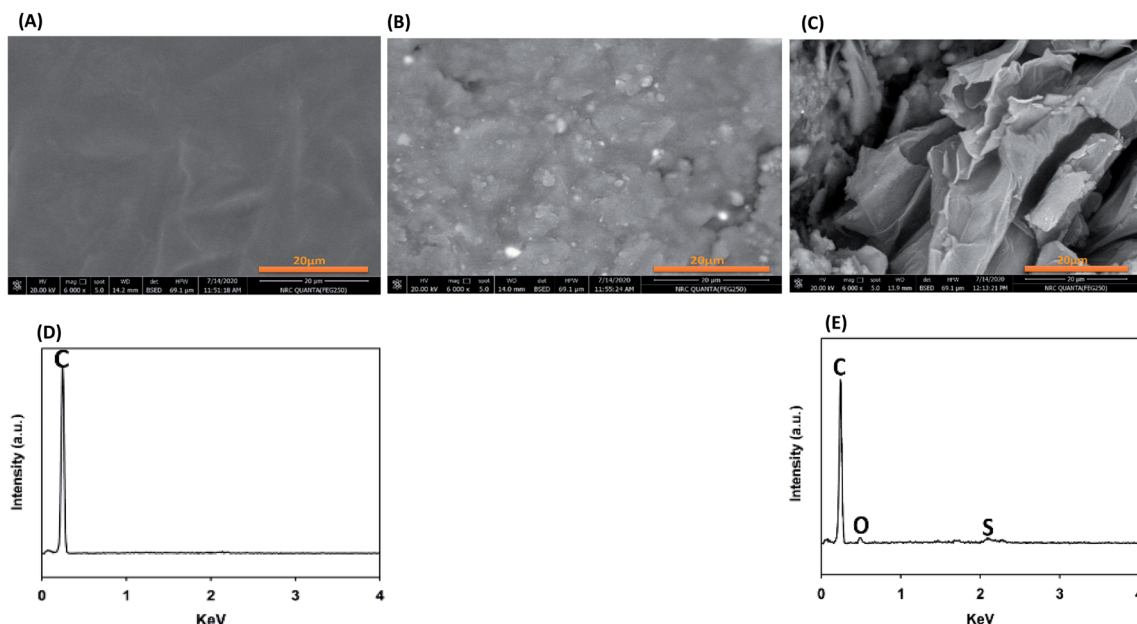


Fig. 1 (A–C) SEM images of (A) RGR' smooth region, (B) RGR' rough region, and (C) FEGR electrodes. (D and E) EDX analysis of RGR' smooth region (D) and FEGR (E) electrodes, respectively.

broad peak in the FEGR of C 1s is evidence for the successful oxidation and the insertion of more functional groups on the graphite surface which is obviously clear from the deconvolution of FEGR C 1s peak into four peaks which are corresponding to C=C, \*C–OH, \*C=O and \*COOH.<sup>46–48</sup> Also, the enlarged view of O 1s of both electrodes shown in Fig. 2C displays that the FEGR peak has a higher intensity compared to that of RGR indicating the higher oxygen content on the FEGR electrode surface (7.48%) compared to the RGR surface (4.94%). Also, the deconvolution of O 1s of FEGR indicates the formation of surface functional groups which is consistent with the deconvolution of its C 1s.<sup>48–50</sup>

Further investigation was done to confirm the presence of the functional groups and defects that are introduced to the surface of the FEGR during the electrochemical partial exfoliation using the FT-IR spectra, Raman spectroscopy, and contact angle measurements. Fig. 3A displays the FTIR spectra of the FEGR electrode to further ensure the presence of the functional

groups on the surface as depicted from the XPS analysis. While the presence of the peak around  $1700\text{ cm}^{-1}$  indicates the presence of carbonyl group (C=O), the presence of an unsmooth broad peak started from  $\sim 3370$  to  $3750\text{ cm}^{-1}$  refers to the presence of both hydroxyl group (–OH) and carboxylic group (–COOH). Also, (C=C) group stretching was observed at wave number around  $1600\text{ cm}^{-1}$ .<sup>51</sup> It is clear that FT-IR spectra agree with the deconvoluted peaks in the XPS analysis.

Moreover, Raman spectra are presented in Fig. 3B for RGR and FEGR electrodes to investigate the change in the crystallinity that happened during the electrochemical exfoliation by comparing the characteristic vibrational mode of carbon materials before and after the oxidation step. Generally, there are three characteristic bands G, 2D, and D bands are observed in Raman spectra at wavenumbers  $1586\text{ cm}^{-1}$ ,  $2696\text{ cm}^{-1}$ , and  $1348\text{ cm}^{-1}$  respectively which are corresponding to the carbonaceous materials. While the G and 2D band are arising due to  $E_{2g}$  vibrational mode of the  $sp^2$  hybrid graphitic carbon and the

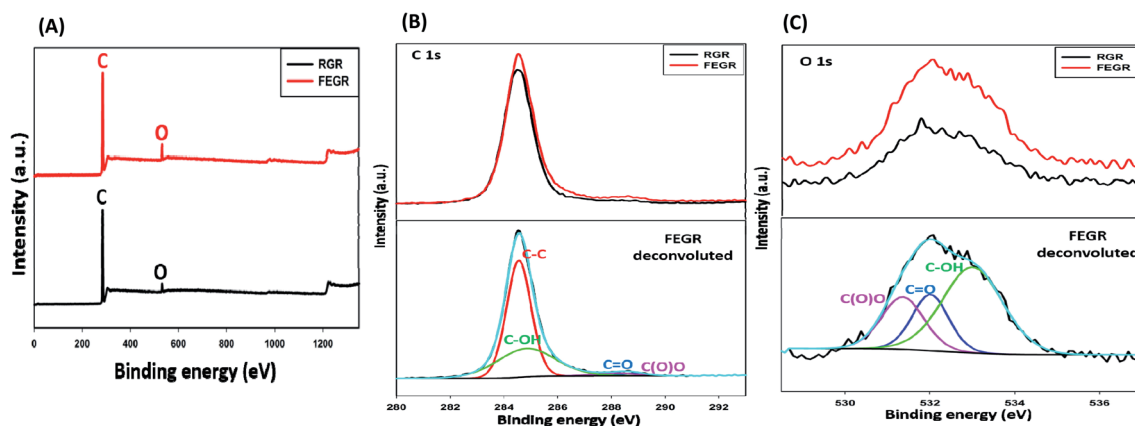


Fig. 2 XPS analysis of RGR (black line), and FEGR (red line) electrodes: (A) survey, (B) enlarged view of C 1s, and (C) enlarged view of O 1s.



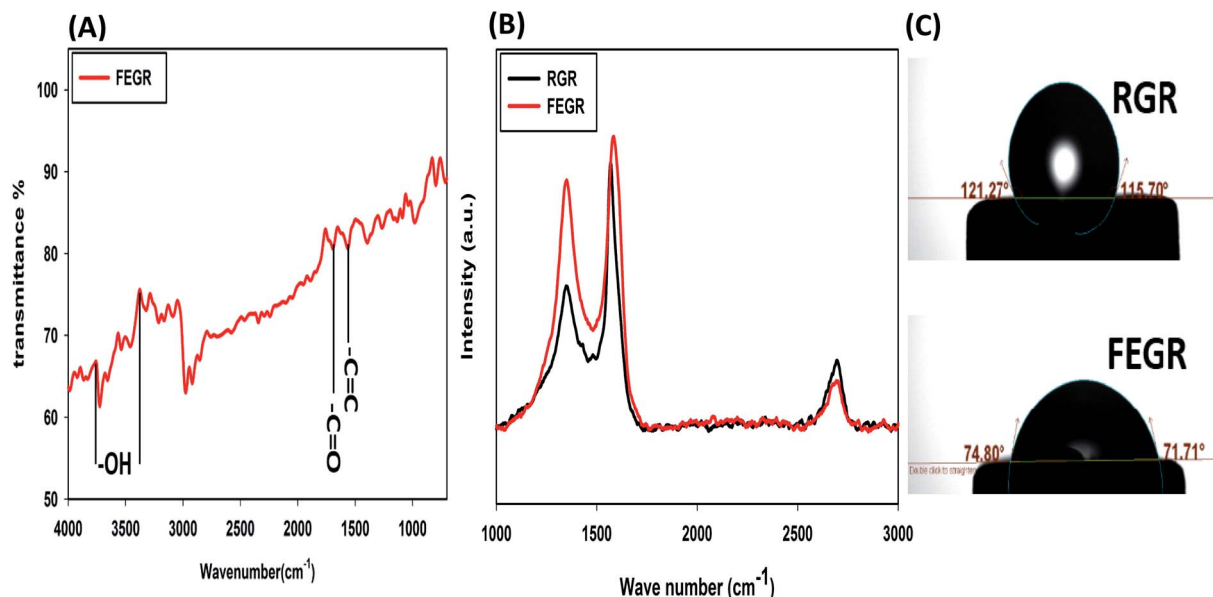


Fig. 3 (A) FT-IR spectrum of FEGR electrode. (B) Raman spectra of RGR (black line) and FEGR electrode (red line). (C) Contact angle test for RGR and FEGR electrode.

double resonance scattering within the crystal lattice of carbon materials, respectively, the D band is related to the presence of the oxygenated, defective, and edged carbon atoms of carbon materials.<sup>44,46,50,52,53</sup> Also, the degree of disorder of carbon materials can be noticed from three important factors; band intensity, band shape, and band position. By increasing the intensity ratio  $I_D/I_G$  and decreasing the intensity ratio  $I_{2D}/I_G$ , the degree of crystallinity is decreased, and the degree of disorder is increased. While the calculated  $I_D/I_G$  value is increased from 0.5454 to 0.8535 corresponding to RGR and FEGR electrodes, respectively, the  $I_{2D}/I_G$  value is decreased from 0.2713 (RGR) to 0.1797 (FEGR). It is clearly obvious from these results that there is an increase in the degree of disorder (roughness) of FEGR electrode due to the inserted functional groups and defects that are generated during the production of graphene on the surface of bulk graphite by electrochemical exfoliation.<sup>41,44,46,49,53–55</sup> Also, the more broadening and increase in the wavenumber of G band in FEGR electrode compared to the RGR electrode confirms the more defects formed at the surface of FEGR electrode.<sup>52</sup> The decrease of full width at half a maximum of the 2D band of FEGR electrode compared to RGR is a good indication of the successful exfoliation of graphite into graphene layers as obtained from the SEM analysis, see Fig. 1(A–C).<sup>55</sup>

Finally, the water droplet method was used to investigate the contact and accessibility of electrolyte ions into the surface of FEGR and RGR electrodes. It is clear that FEGR electrode displays good wetting properties than RGR electrode as a consequence of the electrochemical exfoliation step which increases: (i) the roughness of the electrode surface, (ii) the functional groups on the electrode surface, (iii) the number of graphene layers, and also (iv) the porosity of the electrode material as evidenced from the previous analysis. So, all these factors enhance the contact between electrolyte and the surface of the electrode and also increase the diffusion behavior of

electrolyte.<sup>37,45</sup> The contact angle of FEGR electrode is smaller than that of RGR electrode as shown in Fig. 3C which indicates the higher hydrophilicity nature of FEGR electrode compared to RGR electrode.

**3.1.3. Electrochemical performance of RGR and FEGR electrodes.** As predicted from the physical analysis of both FEGR and RGR electrodes, FEGR electrode has unique properties. Consequently, these properties are reflected in the electrochemical performance. FEGR electrode has a superior electrochemical performance than RGR electrode and this is observed firstly from the integrated surface area obtained from CVs of both electrodes as displayed in Fig. 4A. FEGR electrode has a much larger surface area than the RGR electrode, which indicates that the FEGR electrode has higher  $C_s$  than the RGR electrode. The larger  $C_s$  of FEGR electrode is attributed to, the large activated surface of this electrode to ingest a large number of electrolyte ions on its surface by supplying a rough surface, the large number of the holes on the surface which make the adsorption of electrolyte species available for the internal part of the electrode, and finally the large produced number of expanded graphene sheets. All these factors contribute to increasing the  $C_s$  of FEGR electrode by increasing the electrostatic attraction site of the electrolyte ions. Also, the inserted functional groups are another important factor that contributes to the enhancement of the  $C_s$  of the FEGR electrode by increasing the redox sites on its surface as observed by the presence of redox peaks between 0.35 and 0.45 V which is attributed to the hydroquinone or phenol sites.<sup>44,54,56–58</sup>

Remarkably, the ideal capacitor is characterized by very fast adsorption–desorption at the electrode surface and this can be demonstrated electrochemically by using eqn (8) which describes the relationship between current ( $I$ ) at a characteristic potential with potential scan rate ( $\nu$ ).



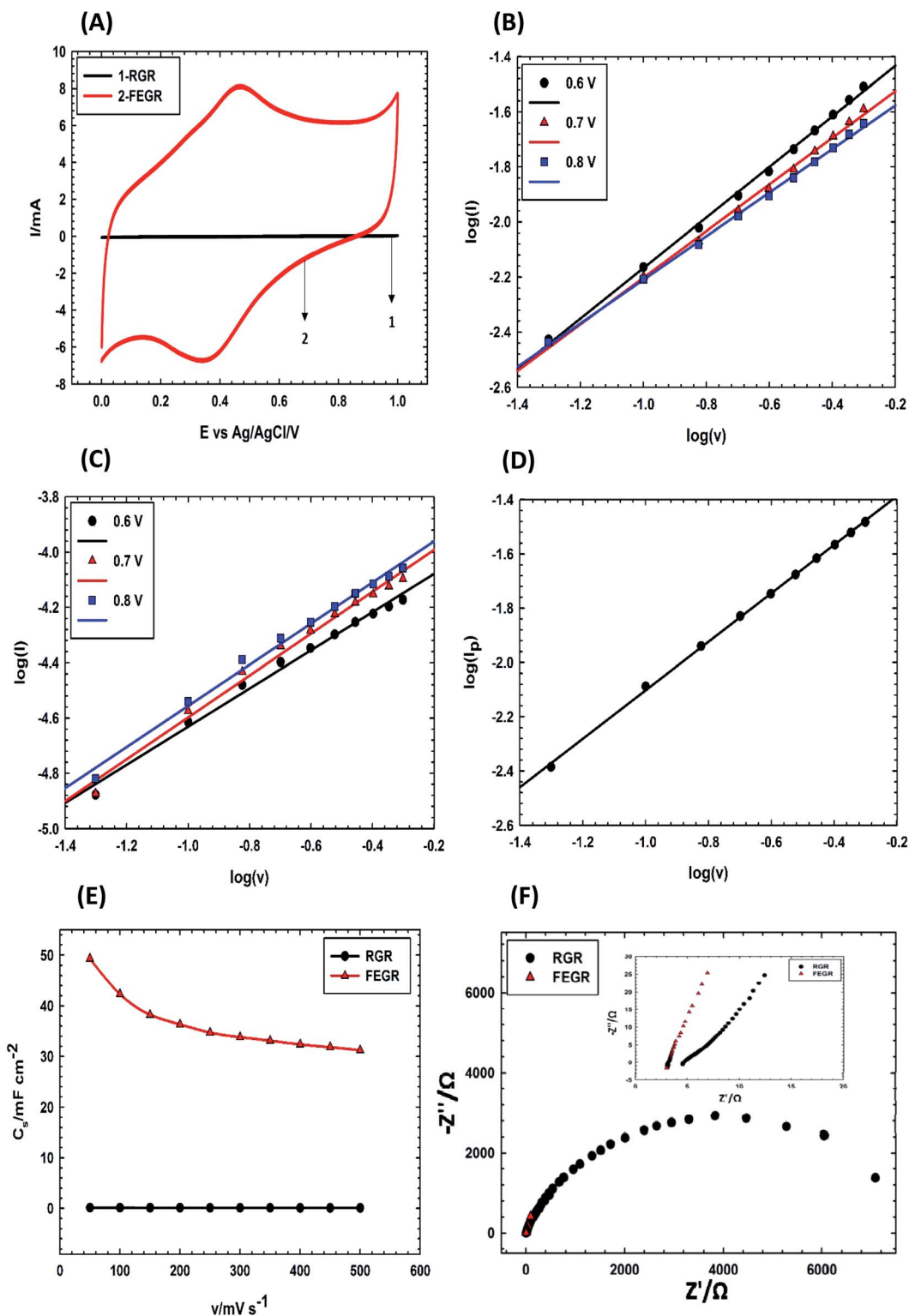


Fig. 4 (A) CVs of RGR (black line), and FEGR (red line) electrodes at a potential scan rate of  $100\ mV\ s^{-1}$ . The variation of ( $\log I$ ) with ( $\log v$ ) at different potentials (0.6, 0.7, and 0.8 V) for (B) RGR electrode, and (C) FEGR electrode. (D) The variation of ( $\log I_p$ ) with ( $\log v$ ) of FEGR electrode. (E) The variation  $C_s$  with  $v$  of RGR (black line) and FEGR (red line) electrodes. (F) Nyquist plots of RGR (black line) and FEGR (red line) electrodes at OCP and its inset at high-frequency region.



**Table 1** The values of  $b$  of RGR and FEGR electrodes at different potentials

Potential (V)	$b$ -Value	
	FEGR electrode	RGR electrode
0.6	0.92	0.69
0.7	0.85	0.76
0.8	0.79	0.74

$$I = av^b \quad (8)$$

If the  $b$  value is equal to 1, the kinetics of the electrode will have ideally EDLC behavior.<sup>50,59</sup>

Fig. 4(B and C) describes the electrode kinetics of RGR, and FEGR RGR electrodes, respectively, at three characteristic potentials 0.6, 0.7, and 0.8 V. The  $b$ -values of FEGR electrode as observed from Table 1 is higher than that of RGR electrode at any potential which indicates that FEGR electrode has higher kinetics behavior than RGR electrode. This can be attributed to the following reasons: (i) the fast diffusion of the electrolyte ions inside the holes which provide a good channel for insertion and desertion without any hindrance,<sup>53,60</sup> (ii) the presence of the functional groups which facilitates the adsorption-desorption by making the contact between electrode and electrolyte be very fast as revealed by contact angle measurements,<sup>37,59</sup> and (iii) the presence of a large number of the graphene sheets which has

a very high electrical conductivity due to the low HOMO–LUMO gap, and also the graphene sheets are away from each other which make the diffusion of the electrolyte within the sheets is very fast.<sup>53–55</sup>

Moreover, the  $b$ -value of the redox reaction due to the presence of the functional groups can be obtained using the same equation (eqn (8)) by plotting ( $\log I_{pa}$ ) vs. ( $\log v$ ) as displayed in Fig. 4D. The  $b$ -value of the redox reaction equals 0.89 which indicates the fast redox surface reactions. As a result, the kinetics of FEGR is faster than that over RGR electrode due to the enhancement of the ionic and electronic pathways.

Also, FEGR electrode exhibited a higher rate performance than RGR electrode as shown in Fig. 4E. Obviously, the calculated  $C_s$  of FEGR electrode at  $50 \text{ mV s}^{-1}$  using eqn (1) is equal to  $49.33 \text{ mF cm}^{-2}$  and this value can be retained to 63.27% at a very high scan rate of  $500 \text{ mV s}^{-1}$ . On the other hand, the RGR electrode displayed a retention value equal to 57.59% at scan rate from 50 to  $500 \text{ mV s}^{-1}$  and this is due to the poor kinetics of this electrode compared to FEGR electrode.

To further ensure the favorable kinetics and capacitive behavior of FEGR electrode over RGR electrode, EIS measurements were carried out at OCP on both electrodes. According to the ideal theoretical capacitive behavior, the electrode must possess a vertical line that makes an angle equal to  $90^\circ$  with the real impedance axis in the Nyquist impedance plot.<sup>61</sup> As observed from the Nyquist impedance plot of the two electrodes shown in Fig. 4F and its inset, the FEGR electrode exhibited the steepest line which indicates that it has a higher capacitive

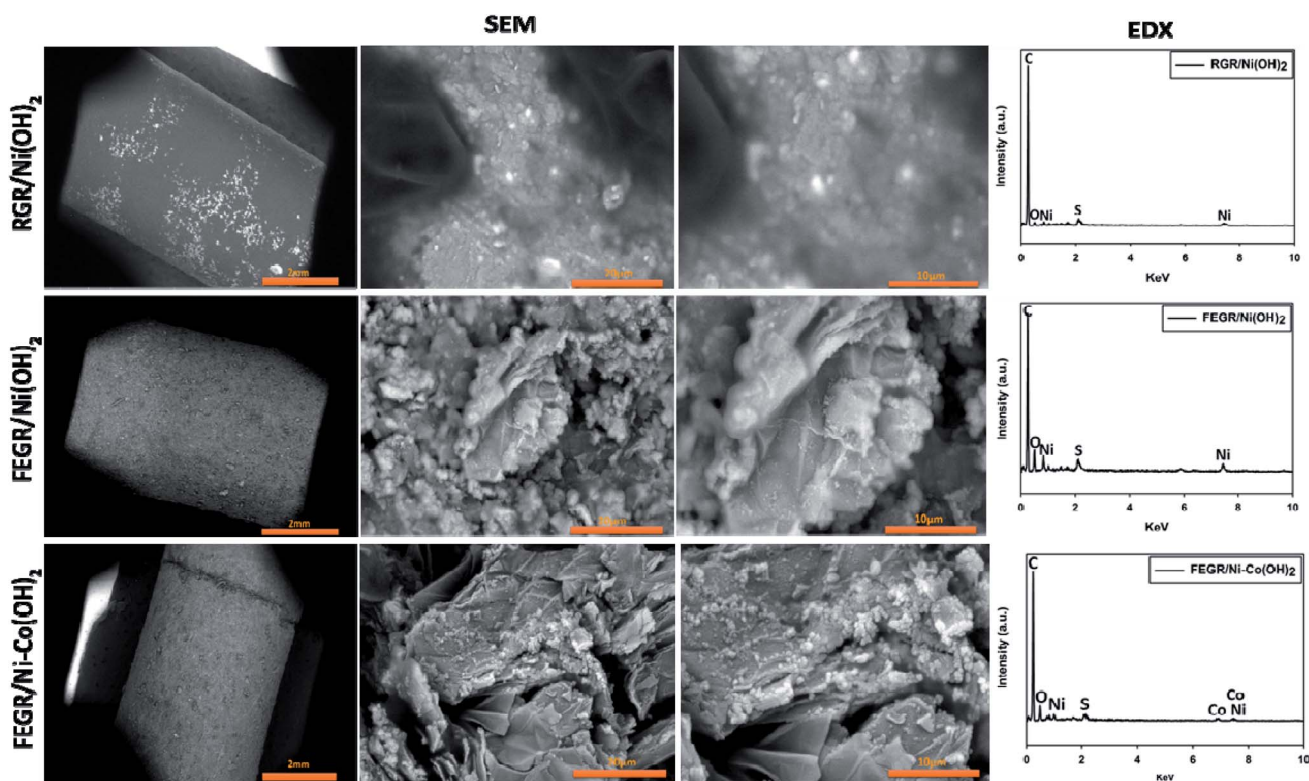


Fig. 5 SEM images with different magnifications and EDX analysis of RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni–Co(OH)<sub>2</sub> film electrodes.





behavior than RGR electrode due to the facilitation of the ions transportation on the surface of the electrode which is in agreement with the kinetic study.<sup>37,45,60</sup> Also, the electronic conductivity of the electrode is enhanced, and this is obvious from the value of ESR that can be determined by the intersection of the impedance curve with *x*-axis at a higher frequency region from the Nyquist plot as clear in the inset of Fig. 4F. FEGR electrode has a lower ESR value and this is reflecting the successful exfoliation of the graphite rod to the graphene sheets.<sup>46,55,60</sup>

As a result of the higher electrochemical performance of FEGR electrode due to its 3D porous structure with the presence of the graphene sheets in the external surface and the presence

of more functional groups, this electrode can be used as a promising substrate to enhance redox activity of the redox-active materials as will be discussed in the next Section 3.2.

### 3.2. Effect of modified substrate on electrochemical performance of Ni(OH)<sub>2</sub> and Ni-Co(OH)<sub>2</sub>

**3.2.1. Physical characterization of the modified RGR and FEGR electrodes.** The as-prepared films, RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub>, were inspected morphologically using SEM analysis. As clear from Fig. 5, the active particles of Ni(OH)<sub>2</sub> were electrodeposited on the different spots of RGR surface with aggregations, and this reflecting the smooth surface properties and the hydrophobicity nature of RGR

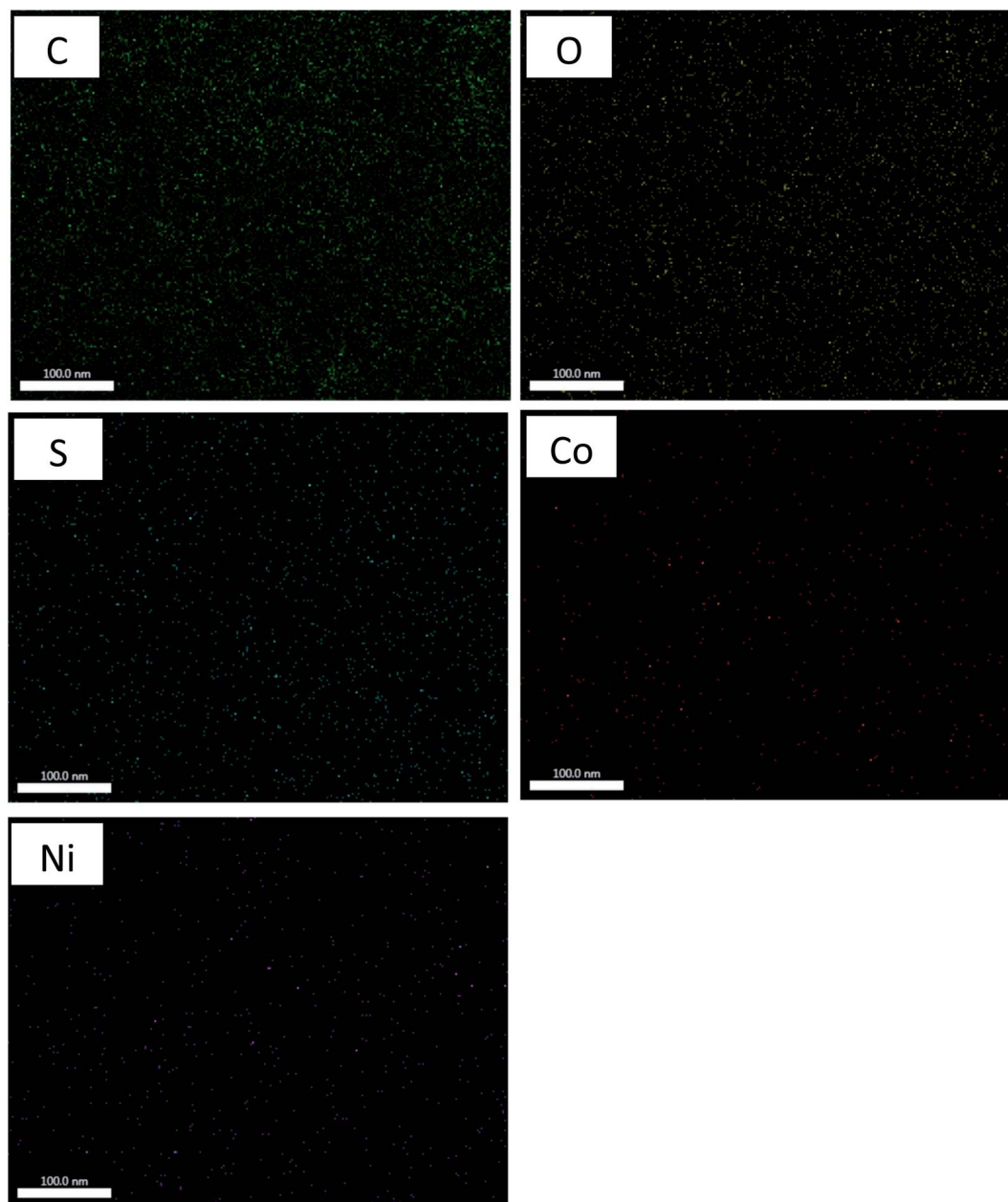


Fig. 6 Elemental mapping EDX analysis of FEGR/Ni-Co(OH)<sub>2</sub> film.



substrate which makes the distribution of the active materials is very poor and this suggests that the electrochemical performance of this electrode is not good that will be discussed later and is shown in Fig. 7 and 8.

On the other hand, the distribution of the electrodeposited Ni(OH)<sub>2</sub> and Ni-Co(OH)<sub>2</sub> binary hydroxide, see Fig. 5, on the FEGR is very well which is distributed uniformly inside the pores and within the graphene layers of the external surface of FEGR electrode that makes almost the active material acts as outer active surface and is available to interact with the electrolyte ions very easily resulting in higher energy produced during the charging and discharging process. Additionally, the enhanced distribution behavior of the FEGR/Ni-Co(OH)<sub>2</sub> film as results of synthesis graphene layers, high surface area, and inserted O functions groups is observed by the successful distribution of elemental O, Ni, and Co over FEGR substrate from mapping EDX analysis (cf. Fig. 6).

Also, EDX analysis was performed to investigate the surface chemical composition of the as-prepared three films. From Fig. 5, recognized peaks of Ni and O appear in all three prepared films which confirm the successful electrodeposition of Ni in all films. Additionally, Fig. 5 has an extra peak for the presence of Co which confirms the successful formation of Co in the binary hydroxide film (FEGR/Ni-Co(OH)<sub>2</sub>). Generally, the presence of sulfur peak in all prepared films is due to the incorporation of SO<sub>4</sub><sup>2-</sup> from the deposition bath between the layers' structure of the metal hydroxide which suggests the appearance of some facets corresponding to the alpha metal hydroxide phase in the XRD patterns of all films as will be shown in Fig. 7. Specifically for the FEGR/Ni(OH)<sub>2</sub> and FEGR/Ni-Co(OH)<sub>2</sub> films, there is another contribution of the appearance of the sulfur peak in their EDX analysis that comes from the incorporation of SO<sub>4</sub><sup>2-</sup> between the graphite layers during the expansion to produce the expanded graphene layers during the electrochemical exfoliation of the RGR to produce the FEGR electrode.<sup>44</sup> Interestingly, the mass percent of the oxygen atoms in all films is 7.61%, 23.29%, and 21.67% for the RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films, respectively. The higher oxygen content in the FEGR/Ni(OH)<sub>2</sub> and FEGR/Ni-Co(OH)<sub>2</sub>

films is due to the insertion of more oxygen functional groups during the preparation of the FEGR substrate.

XRD patterns of the as-prepared films, *i.e.*, RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> are depicted in Fig. 7. As described elsewhere,<sup>62–66</sup> the similarity of the crystalline structure of Ni(OH)<sub>2</sub> and Co(OH)<sub>2</sub> make the diffraction peaks very close to each other and one can't distinguish between them as seen from the XRD patterns of the proposed films. While the four peaks observed in XRD patterns around 21°, 36°, 37°, and 63° indicate the formation of β-Ni(OH)<sub>2</sub> and β-Co(OH)<sub>2</sub>, then the two peaks around 12° and 23° are corresponding to α-Ni(OH)<sub>2</sub> and α-Co(OH)<sub>2</sub> due to the incorporation of SO<sub>4</sub><sup>2-</sup> inside the crystalline structure as predicted from the EDX analysis.<sup>67</sup> Also, the presence of peaks around 25°, 42°, 43°, 77°, and 83° are due to the hexagonal graphitic structure of RGR and FEGR substrate.<sup>6,68</sup> So, one can conclude from the similarity of the patterns of all films that the enhancement of capacitive behavior of the active materials on the FEGR substrate over the RGR substrate (cf. Fig. 8, and 9) is attributed to a better distribution of active materials on the surface of the substrate as evident from SEM images of all films which means that the phase difference has no role in this regard because all films have the same crystalline structure (*i.e.* same phases and facets).

**3.2.2. Electrochemical performance of the three prepared films.** As indicated in Fig. 8A, the integrated surface area of both RGR and FEGR substrate is very small compared to that produced from prepared films (RGR/Ni(OH)<sub>2</sub> and FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub>) which confirm that substrates have insignificant contribution in the total capacity of the prepared films. Additionally, CVs of the prepared films show characteristic redox peaks within the selected potential window resulting from the transformation of the α- and β-M-OH to β- and γ-M-OOH. This is due to the intercalation and deintercalation of the H<sup>+</sup> within the crystal structure of metal hydroxide according to the following eqn (9) and (10).<sup>34</sup>



where the M represents Ni for the RGR/Ni(OH)<sub>2</sub> and FEGR/Ni(OH)<sub>2</sub> films and the Ni and Co for prepared binary hydroxide film FEGR/Ni-Co(OH)<sub>2</sub>, indicating that all films have the battery type behavior. This behavior is confirmed by fitting the linear relation between log *I*<sub>pa</sub> and log *I*<sub>pc</sub> with log *ν* according to eqn (8) where the *b*-value is equal to 0.5.<sup>59</sup> As observed from Fig. 8B, log(*I*<sub>pa</sub>) and log(*I*<sub>pc</sub>) for all as-prepared films have a linear relation with log *ν* where *b*-value equal 0.5 with *R*<sup>2</sup> value equal 0.999 which confirms the kinetics of the battery type like-electrode for all films.

As the indication of the large integrated surface area of the charging or discharging branch of the FEGR/Ni(OH)<sub>2</sub> and FEGR/Ni-Co(OH)<sub>2</sub> compared with RGR/Ni(OH)<sub>2</sub>, the *C* value of the active materials on modified FEGR electrode is greatly enhanced. This firstly is due to FEGR electrode provides homogeneous and uniform distribution of the active materials on its surface due to its high surface area. Secondly, the more

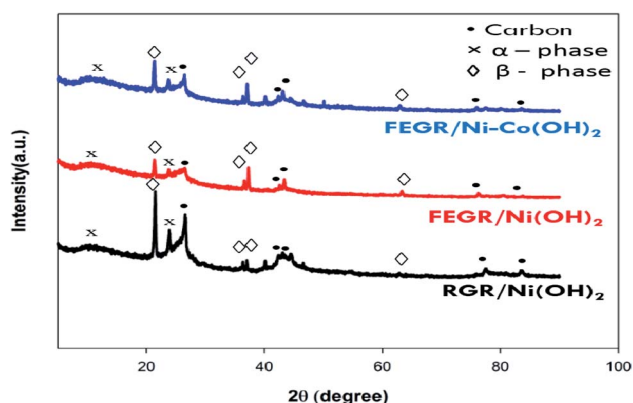


Fig. 7 XRD patterns for RGR/Ni(OH)<sub>2</sub> (black line), FEGR/Ni(OH)<sub>2</sub> (red line), and FEGR/Ni-Co(OH)<sub>2</sub> (blue line) films.



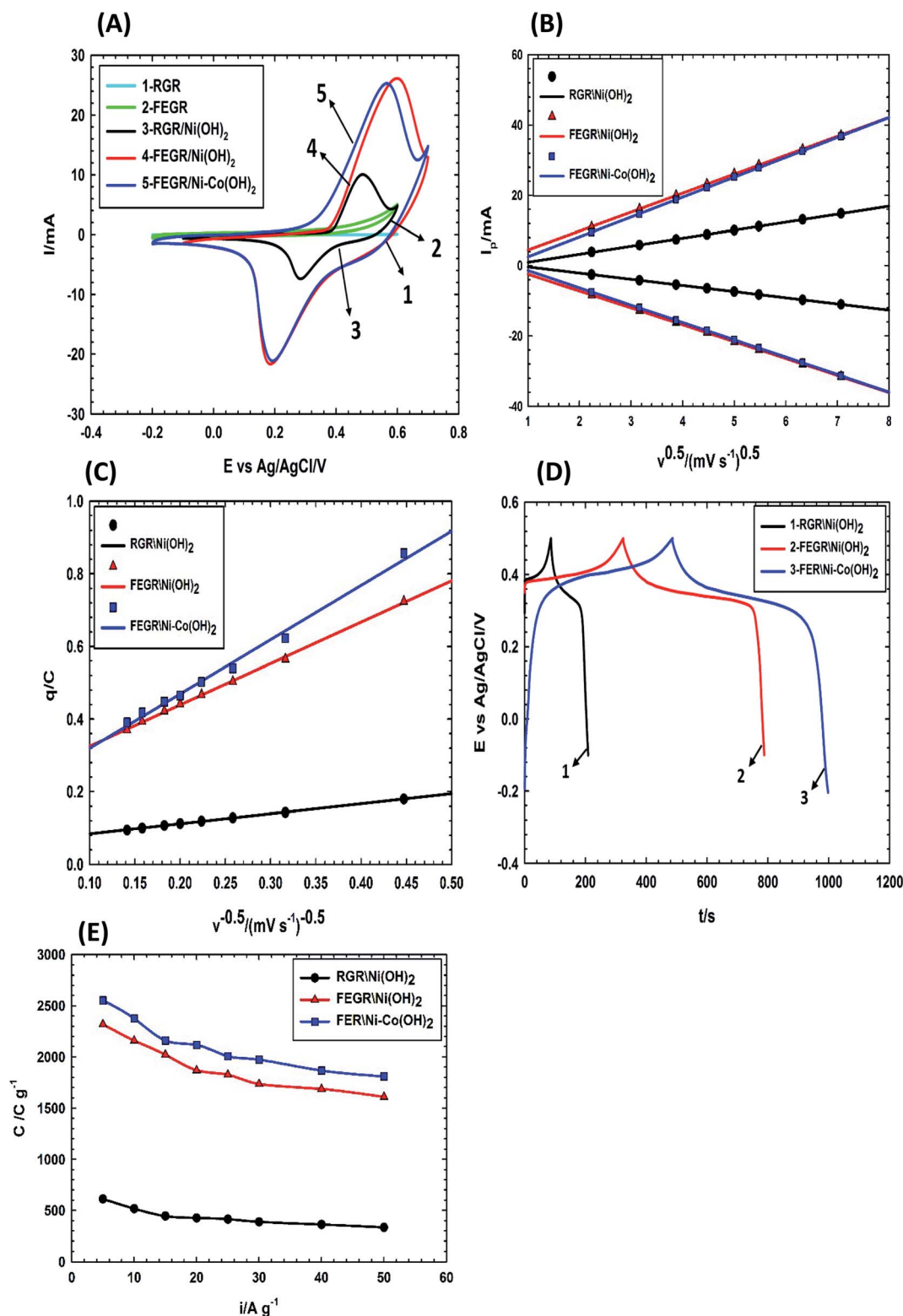


Fig. 8 (A) CVs obtained at (1) bare RGR, (2) bare FEGR substrates and (3) RGR/Ni(OH)<sub>2</sub>, (4) FEGR/Ni(OH)<sub>2</sub>, and (5) FEGR/Ni-Co(OH)<sub>2</sub> films measured in 0.5 M NaOH solution at a potential scan rate of 25 mV s<sup>-1</sup>. (B) The variation of anodic and cathodic peak current with the square root of potential scan rate ( $\nu^{0.5}$ ) for RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films. (C) The variation of  $q$  with reciprocal of the square root of the potential scan rate ( $\nu^{-0.5}$ ). (D) GCD curves of RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films at a current density of 5 A g<sup>-1</sup>. (E) The rate capability curves of RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films at various current densities.



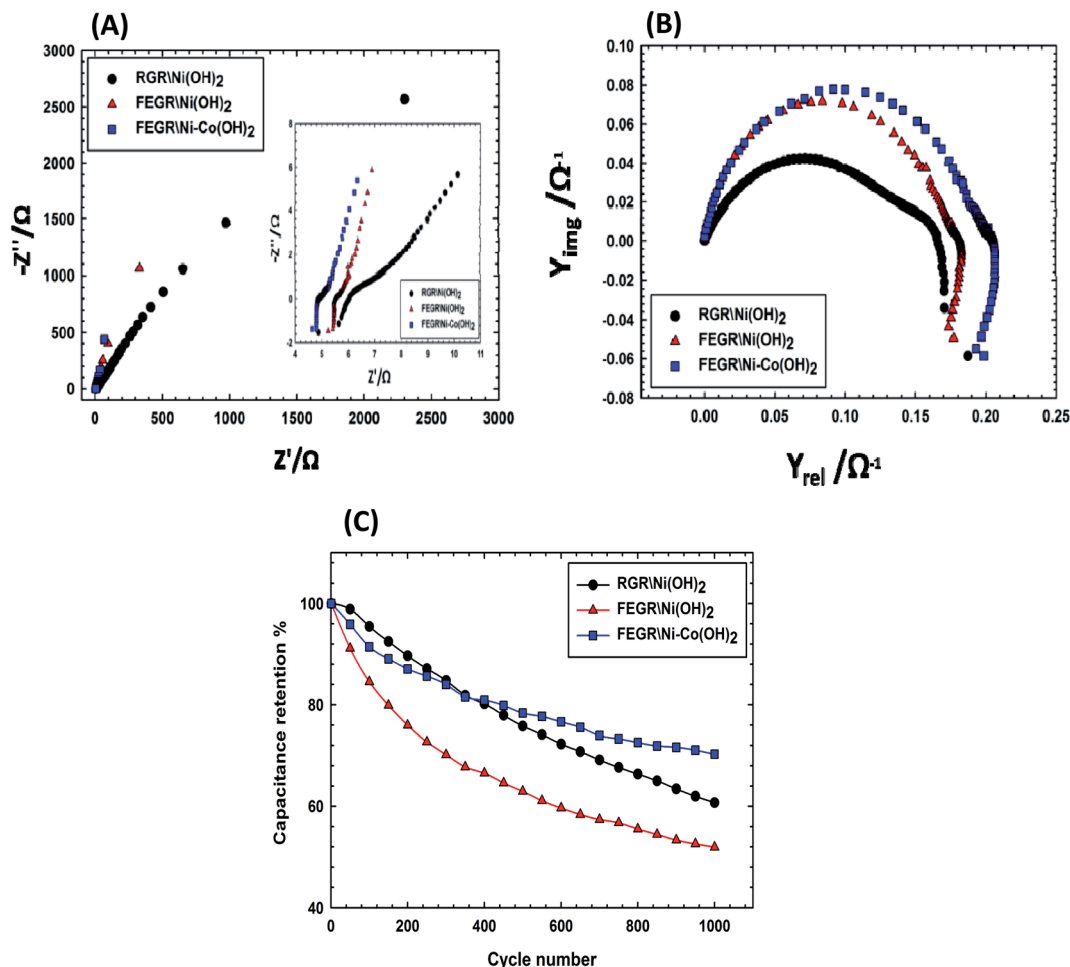


Fig. 9 (A) Nyquist plots of RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films at OCP and their insets at high frequencies. (B) Admittance plots of RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films. (C) Stability curves of RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films at a current density of 50 A g<sup>-1</sup>.

inserted and distributed functional groups on FEGR surface makes almost the active materials act as outer active surface, more accessible area, and this can be demonstrated easily by determining the amount of the outer active site which can be estimated from the intercept value of the following simple empirical relation of eqn (11).<sup>59,69–71</sup>

$$q = q_{\infty} + av^{-0.5} \quad (11)$$

where the  $q$  is the charge (Coulomb) at different potential scan rates obtained by integrating CVs,  $q_{\infty}$  is the maximum amount of the charge related to the “outer” surface of active material,  $a$  is a constant, and  $v$  is the potential scan rate (mV s<sup>-1</sup>). The value of the intercept of all films is obtained from Fig. 8C is 0.057, 1.14, and 1.49 C for RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films, respectively. The values are consistent with SEM images that displayed homogeneous distribution of the active materials on the FEGR electrode surface compared with RGR electrode. Moreover, the FEGR also increases the diffusion efficiency of the electrolyte ions to reach an active site by very high speed due to its hydrophilic nature and unique 3D

porous structure.<sup>50,59,71</sup> In addition, the FEGR increases electron transportation to the active materials *via* the developed graphene layers during the exfoliation process which is characterized by excellent electronic properties.<sup>59,71</sup>

Fig. 8D shows the GCD curves of all as-prepared films at current density equal to 5 A g<sup>-1</sup>. The maximum  $C$  of each film can be calculated from the discharge time at current density of 5 A g<sup>-1</sup> using eqn (3) and it is estimated to be 612.5, 2318.4, and 2552.6 C g<sup>-1</sup> corresponding to RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni-Co(OH)<sub>2</sub> films, respectively. Note that the values of  $C$  for the FEGR-based electrodes is much improved compared to the similar RGR-based electrodes. Moreover, these values exceed the theoretically-calculated  $C$  values (*ca.* 1730 C g<sup>-1</sup>) based on the redox phenomenon of the metal hydroxide ( $C = nF/M_w$ ).<sup>12–14</sup> This could be possibly attributed to (i) the enhanced redox behavior of the metal hydroxide to higher oxidation states with average number of electron transfer ( $n$ ) approaching 1.7,<sup>42</sup> (ii) better distribution of the metal hydroxide in the case of FEGR-substrate (see SEM images in Fig. 5 and the corresponding mapping EDAX in Fig. 6), thus leading to a higher electrochemical active surface area, and (iii) a considerable





**Table 2** Comparison between the capacitive performance of Ni–Co(OH)<sub>2</sub>/FEGR electrode with other reported binary Ni–Co hydroxide in SCs applications

Method	Electrolyte	<i>C</i> (C g <sup>−1</sup> )	Rate performance	Cyclic performance	Ref.
Precipitation	1 M KOH	309.6	~55.6% (0.5 to 5 A g <sup>−1</sup> )	75% (500 cycle at 1 A g <sup>−1</sup> )	75
Precipitation	6 M KOH	1307	85.9% (1 to 10 A g <sup>−1</sup> )	86.4% (1000 cycle at 5 A g <sup>−1</sup> )	76
Hydrothermal	3 M KOH	1194.7	48% (1 to 10 A g <sup>−1</sup> )	77% (1500 cycle at 10 A g <sup>−1</sup> )	77
Hydrothermal	6 M KOH	794.8	59.5% (2 to 40 A g <sup>−1</sup> )	40% (3000 cycle at 30 A g <sup>−1</sup> )	78
Solvothermal	3 M KOH	860	76.9% (1 to 20 A g <sup>−1</sup> )	74% (1000 cycle at 20 A g <sup>−1</sup> )	79
Solvothermal	1 M KOH	1084.1	~22% (9.1 to 31.8 A g <sup>−1</sup> )	67.8% (2000 cycle at 31.8 A g <sup>−1</sup> )	80
Microwave	6 M KOH	849.4	62.9% (1 to 10 A g <sup>−1</sup> )	99.7% (3000 cycle at 8 A g <sup>−1</sup> )	81
CBD	2 M KOH	923.5	47% (5 to 80 A g <sup>−1</sup> )	64% (1000 cycle at 80 A g <sup>−1</sup> )	82
Reflux	6 M KOH	693	88% (1 to 10 A g <sup>−1</sup> )	80% (1000 cycle at 10 A g <sup>−1</sup> )	83
Electrodeposition	1 M KOH	1052.5	56.6% (2 to 20 A g <sup>−1</sup> )	89.5% (2000 cycle at 10 A g <sup>−1</sup> )	84
Electrodeposition	6 M KOH	541.8	91.8% (1 to 10 A g <sup>−1</sup> )	80.1% (1000 cycle at 10 A g <sup>−1</sup> )	22
Electrodeposition	3 M NaOH	959.9	37.7% (4 to 24 A g <sup>−1</sup> )	91% (1000 cycle at 12 A g <sup>−1</sup> )	33
Electrodeposition	1 M KOH	1665.4	73.5% (2 to 50 A g <sup>−1</sup> )	94% (2000 cycle at 20 A g <sup>−1</sup> )	34
<b>Electrodeposition</b>	<b>0.5 M NaOH</b>	<b>2552.6</b>	<b>70.8% (5 to 50 A g<sup>−1</sup>)</b>	<b>70.3% (1000 cycle at 50 A g<sup>−1</sup>)</b>	<b>This work</b>

contribution of the double layer of the underlying FEGR substrate. The active materials on FEGR electrodes' surface exhibit ultra-high *C* compared to RGR substrate and it is also one of the highest reported *C* for Ni(OH)<sub>2</sub> and binary Ni–Co(OH)<sub>2</sub> active materials as reported in Table 2. Interestingly, this value is higher than other active materials loaded on the exfoliated carbon materials such as FEG/MnO<sub>2</sub> (1061 F g<sup>−1</sup>),<sup>37</sup> WO<sub>3</sub>/Ex-GF (495.8 F g<sup>−1</sup>),<sup>72</sup> and PANI–WO<sub>3</sub>/Ex-GF (408 F g<sup>−1</sup>).<sup>73</sup>

Fig. 8E illustrates the variation of *C* with different discharge current densities for the investigated films from 5 to 50 A g<sup>−1</sup>. Capacitive retention for each film at the current density of 50 A g<sup>−1</sup> is 44.6%, 64.9%, and 70.8% which is corresponding to RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni–Co(OH)<sub>2</sub> films, respectively. Generally, the active materials loaded on the FEGR electrode have better rate performance than that loaded on RGR, and this is due to the enhancement of the electronic and ionic diffusion pathways and the homogeneous distribution of active materials as mentioned above.

More specifically the binary hydroxide FEGR/Ni–Co(OH)<sub>2</sub> film has a higher retention value than single hydroxide FEGR/Ni(OH)<sub>2</sub> film. This can be explained by the increment of the total electronic conductivity of Ni(OH)<sub>2</sub> in the presence of Co(OH)<sub>2</sub> because the charging and discharging of Co(OH)<sub>2</sub> leads to the formation of highly conductive Co(OOH) which speeds up the transportation of the electrons which make the utilization of the active materials at high scan speed is more efficient. Also, the energy gap between the valence and conduction bands of Ni(OH)<sub>2</sub> can be decreased by the presence of the Co ions inside its crystal structure.<sup>8,30,74</sup>

For furthermore confirmation of the above-mentioned conclusion, EIS measurements were carried out on the proposed films. Fig. 9A displayed the impedance spectra of the films showing firstly both films FEGR/Ni–Co(OH)<sub>2</sub> and FEGR/Ni(OH)<sub>2</sub> exhibited the excellent capacitive behavior more than RGR/Ni(OH)<sub>2</sub> due to they have the steep line in the high-frequency region compared to the inclined line of RGR/Ni(OH)<sub>2</sub>.<sup>37,61</sup> Secondly, from the low-frequency region, see inset of Fig. 9A, the loaded active materials on FEGR film has a lower

ESR value than that loaded on RGR film and this can be attributed to the improved conductivity due to the synthesized graphene layers.<sup>37,55</sup> Interestingly, the ESR value of FEGR/Ni–Co(OH)<sub>2</sub> film is smaller than that of RGR/Ni(OH)<sub>2</sub> film and this demonstrates the role of the Co(OH)<sub>2</sub> in increasing the total electronic conductivity of the electrode. Also, the admittance plot, see Fig. 9B, displayed the similar behavior of ESR value that is in the order that RGR/Ni(OH)<sub>2</sub> > FEGR/Ni(OH)<sub>2</sub> > FEGR/Ni–Co(OH)<sub>2</sub> film accompanied with the increment of the conductivity value in the order of RGR/Ni(OH)<sub>2</sub> < FEGR/Ni(OH)<sub>2</sub> < FEGR/Ni–Co(OH)<sub>2</sub> because the admittance, as known, is the inverse of the impedance.

Moreover, the stability test of the assigned films was also examined by GCD test for 1000 cycles at 50 A g<sup>−1</sup> current density, see Fig. 9C. Capacitive retention for each film is 60.7%, 52%, and 70.3% for RGR/Ni(OH)<sub>2</sub>, FEGR/Ni(OH)<sub>2</sub>, and FEGR/Ni–Co(OH)<sub>2</sub> films, respectively. It is clearly obvious from the capacitive retention values that the stability of binary metallic hydroxide film is better than that of the individual Ni hydroxide film. This is due to the incorporation of Co ions into Ni hydroxide layers preventing the mechanical failure that can be happen during the charging process due to the transformation of β-NiOOH to γ-NiOOH.<sup>20,29</sup>

## 4. Conclusion

In summary, the electrochemical partial exfoliation was done on the RGR that obtained from exhausted zinc–carbon batteries to produce FEGR electrode that is characterized by high functionalized surface area with the presence of holes and exfoliated graphene layers. The inserted functional groups on the defected surface of the FEGR electrode are confirmed by the EDX, XPS, FT-IR, and contact angle measurements. Also, the presence of the graphene layers on the top surface is observed from the SEM images. This leads to providing a higher degree of distribution (homogeneous deposition) of the Ni(OH)<sub>2</sub>, and Ni/Co binary hydroxide and also shortening the diffusion pathway for electrolyte ions to efficiently contact more electroactive sites for



faradaic energy storage, even at high current densities, leading to enhanced electrochemical properties. Also, the observed decrease in the equivalent series resistance of the active materials on the FEGR substrate is due to the presence of the graphene layers that can transport the electrons very fast due to its excellent electronic conductivity which is considered another enhancement of the electrochemical performance of the modified FEGR over the modified RGR electrode. Interestingly, FEGR/Ni-Co(OH)<sub>2</sub> film displayed an ultra-high specific capacity (more than 2500 C g<sup>-1</sup> at 5 A g<sup>-1</sup>) which is considered one of the highest reported values based on the binary Ni/Co hydroxide and the value is retained to 70.8% at a high current density of 50 A g<sup>-1</sup> indicating the superb rate performance of this electrode material.

## Conflicts of interest

There are no conflicts to declare.

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