

# Understanding the influence of the biomass-derived alcohols on the activity and stability of Pt nanoparticles supported on graphene nanoribbons

*Rodrigo Teles<sup>a</sup>, Ana Arenillas<sup>b</sup>, Gabriel C. da Silva<sup>c</sup>, Pablo S. Fernández<sup>d</sup>, Eduardo S. F. Cardoso<sup>e</sup>, Gilberto Maia<sup>e</sup> and Cauê A. Martins<sup>a\*</sup>*

<sup>a</sup>Faculty of Exact Sciences and Technology, Universidade Federal da Grande Dourados, 79804-970 Dourados, MS, Brazil.

<sup>b</sup>Instituto Nacional del Carbón, CSIC, Apartado 73, 33080 Oviedo, Spain.

<sup>c</sup>Instituto de Química de São Carlos/Universidade de São Paulo, IQSC-USP, C.P. 780, São Carlos, SP, Brasil.

<sup>d</sup>Institute of Chemistry, State University of Campinas, 13083-970 Campinas, SP, Brazil.

<sup>e</sup>Institute of Chemistry, Universidade Federal de Mato Grosso do Sul, C.P. 549, 79070-900 Campo Grande, MS, Brazil.

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\*Corresponding Authors: C. A. Martins

Zip Code: 79804-970

Phone: +55-67-3410-2089. Fax: +55-67-3410-2089.

e-mail: [cauealvesmartins@gmail.com](mailto:cauealvesmartins@gmail.com), [cauemartins@ufgd.edu.br](mailto:cauemartins@ufgd.edu.br)

## Abstract

We produced Pt/GNRs by a one-step synthesis procedure and evaluated their electroactivity and stability towards glycerol electrooxidation reaction (GEOR) for the first time. We compared the electrocatalytic performance of GEOR with methanol and ethanol electrooxidation on Pt/GNRs at identical experimental conditions. The activities and stabilities for these biomass-derived alcohols electrooxidation on Pt/GNRs were compared to commercial Pt/C. The synthesis of the Pt/GNRs was confirmed by transmission electron microscopy, x-ray diffractometry, ultraviolet spectrophotometry and Raman spectroscopy. We found that the activities of Pt/GNRs for these reactions are comparable to Pt/C, with improvement in terms of current density for methanol electrooxidation. Comparing potentiostatic measurements, we found that glycerol produces lower pseudo-stationary current densities than ethanol and methanol on both catalysts, with greatest values found for methanol electrooxidation on Pt/C. Otherwise, the GNRs remarkably enhance the stability of the catalyst for all the reactions, by increasing the stability of the current density during successive potential cycles, and by preventing the loss of electrochemically active surface area by avoiding carbon corrosion and Pt detachment. Moreover, we showed that the stability of the NPs depends on the biomass-derived alcohol used. The solution containing methanol reveals itself the most aggressive electrochemical environment to the catalyst, impacting in the decrease of surface area, while glycerol is less aggressive. Hence, the different products formed at the interface electrode / solution might lead to different electrochemical environment, which plays an important role on the stability of the catalysts.

**Keywords:** Pt-modified nanoribbons, glycerol electrooxidation, ethanol electrooxidation, methanol electrooxidation, electrochemical stability.

## 1. Introduction

Biomass-derived alcohols such as methanol, ethanol, and glycerol have been investigated as potential fuels for direct alcohol fuel cells (DAFCs). These devices produce energy through a coupled electrochemical reaction, wherein the alcohol is electrooxidized at the anode while oxygen is electroreduced at the cathode [1]. The use of DAFCs is an environmentally friendly alternative to fossil fuels for the production of clean energy. However, several problems remain that pose a challenge for the commercialization of DAFCs: (i) the low activity and low tolerance to poisoning of Pt-based catalysts; (ii) the high cost of precious metal nanocatalysts; (iii) the substandard proton conductivity of the nanocatalysts; and (iv) the low stability of the nanocatalysts. Among the challenges, finding an electrochemically stable catalyst seems to be the most critical. The electrochemical stability is directly related to the electrochemical environment [2], changes in the nature of the metallic catalyst [3], and the nature of the carbon support [4] under operational conditions.

Carbon materials are used in fuel cells to anchor the metallic nanoparticles (NPs), and also work as electron collectors. The corrosion of carbon black, a commonly used commercial material, greatly contributes to the instability of both the cathodes and the anodes in fuel cells [5-9]. Alternative materials such as modified carbon black [10], mesoporous carbon [11,12], carbon nanotubes [13,14], carbon nanofibers [15], and graphenes [1] have been extensively studied as candidates to overcome the instability problem. Among these materials, metallic NPs immobilized on carbon nanotubes (CNTs), multi-walled carbon nanotubes (MWCNTs), and chemically converted graphene (CCG) have emerged as the most stable nanocomposites for fuel cell applications [1,16-19].

Compared with CNTs and CCG, amorphous carbon black is thermochemically less stable and contains a high level of impurities [20], which results in poor electrocatalytic performance. In addition, while CNTs and CCG contain well-defined surface structures in a  $sp^2$  network, carbon black has a high degree of superficial disorder [21]. Although CNTs exhibit high electrical conductivity, high durability, and low levels of impurities, the high aspect ratio and strong  $\pi$ - $\pi$  interactions cause CNTs to form highly packed structures, hindering the uniform deposition of NPs onto the surface. The two-dimensional structure of CCG has similar characteristics to that of CNTs, while allowing the uniform deposition of NPs. The main disadvantage of CCG is the enveloping of NPs, which makes the NPs inactive. The enveloping of NPs represents decay in efficiency in terms of specific activity, since part of the metallic content would not work properly, which might decrease the efficiency of the fuel cell.

Recently, we compared the electrochemical stabilities of carbon black-, CCG-, and MWCNT-modified Pd NPs in an alkaline medium by performing successive voltammetry cycles as a test protocol [17]. Both CCG and MWCNTs enhanced the stability of the nanocomposite compared with carbon black. Furthermore, Pd/CCG was found to be electrochemically more stable during the early cycles, while the Pd/MWCNTs were more stable over the long term. These results were rationalized as a competition between the availability of the support for metallic agglomeration and the corrosion of the support itself. In other words, although CCG initially seems to be more stable, its flat surface allows continuous agglomeration of the NPs over a long period of time; in contrast, the MWCNTs have a steric limitation for agglomeration, and therefore do not continuously agglomerate NPs on their surfaces. Furthermore, in a DFT investigation, Staykov et al. showed that the binding interaction between the

nanoparticles and the support is not significant on flat surfaces but significant on curved surfaces [22]. These authors reported that the chemisorption of NPs on CNTs prevented aggregation, whereas agglomeration of NPs on graphene under high temperatures caused a decrease in stability over time [22]. Considering the importance of both longitudinal curved surfaces and flat surfaces, a question emerges: why not combine the characteristics of both materials into one nanocomposite?

The Tour's group [23,24] introduced a new method to produce graphene-based supports by a simple process for synthesizing graphene nanoribbons (GNRs). Nanoribbons combine the longitudinality of nanotubes and the flat surface of graphene. Such process can be divided in two steps. First, the MWCNTs are strongly oxidized, forming the so-called oxidized graphene nanoribbons (GONRs) [25]. In this step, the walls of the oxidized nanotubes are opened longitudinally, in a process called "unzipping" [23,25]. Second, the GONRs are chemically reduced to produce GNRs. The proposed mechanism for the "unzipping" process is described elsewhere [23]. The longitudinal 2D surface (high length-to-width ratio) of the GNRs changed the nature of the support from a semiconductor to a semimetal [26].

The steric and electronic properties of GNRs lead to advantageous electrochemical features. Lima and Maia showed that a wide variety of easily synthesized large area GNRs show high activity toward the oxygen reduction reaction (ORR) [27]. In a pioneer study, Wang et al. used *in situ* chemical co-reduction to produce Pt/GNRs that showed excellent activity toward the methanol electrooxidation reaction (MEOR) [28]. However, no studies of the activity and stability of Pt-modified GNRs towards glycerol electrooxidation has been reported to date and, to the best of our knowledge, there is no work comparing the electrooxidation of methanol, ethanol,

glycerol. Moreover, the different experimental set up used in the distinct reports make the comparison of the alcohols electrooxidation very difficult. Here, we report a simple method for the one-step synthesis of Pt/GNRs for biomass-derived electrooxidation (methanol, ethanol, and glycerol) and an investigation of their electrochemical stabilities in the presence of the fuel. We present for the first time a thorough comparison of the electrooxidation properties of the alcohols on the Pt/GNRs.

## **2. Experimental**

### ***2.1 Synthesis of graphene oxide nanoribbons (GONRs)***

The GONRs were produced by a method adapted from Hummers and Offeman [29], as reported by Lima and Maia [27]. Briefly, 0.70 g MWCNTs ( $10\pm 1$  nm O.D.,  $4.5\pm 0.5$  nm I.D., 3–6  $\mu\text{m}$  long; 6–8 tubes; Aldrich, used as received) were dispersed in 20 mL of concentrated  $\text{H}_2\text{SO}_4$ . Next, 0.37 g of  $\text{K}_2\text{S}_2\text{O}_8$  and 0.37 g  $\text{P}_2\text{O}_5$  were added to the solution, which was heated at  $80^\circ\text{C}$  for 6 h under stirring. Then, the solution was cooled to room temperature and immersed in a water bath at  $0^\circ\text{C}$ . The product was filtered under vacuum through a 0.22  $\mu\text{m}$  Nylon membrane while washing with water to achieve a neutral pH. The resulting pre-oxidized graphene nanoribbons were dried at room temperature.

Next, 0.70 g of the pre-oxidized graphene nanoribbons were re-oxidized [30] by adding 40 mL of concentrated  $\text{H}_2\text{SO}_4$ , 0.70 g of  $\text{NaNO}_3$ , and 2.10 g of  $\text{KMnO}_4$  to the reaction flask at  $0^\circ\text{C}$ . The mixture was held at  $20^\circ\text{C}$  under stirring, and then heated to  $35^\circ\text{C}$  for 2 h. Next, we added 120 mL of water at  $0^\circ\text{C}$  and 200 mL of water at  $25^\circ\text{C}$ . Then 40 mL 30%  $\text{H}_2\text{O}_2$  was added, and the mixture was stirred for at least 20 min. The solution was cooled to room temperature and transferred to a 3500-mL container of

water at 0°C. After 24 h, the precipitate was centrifuged and washed three times with 1) water, 2) aqueous HCl, and 3) water. The GONRs were obtained by filtration and washed with water on a Nylon membrane. Finally, the GONRs were dried in an oven at 30°C, forming a powder.

## ***2.2 Synthesis of Pt nanoparticles immobilized on graphene nanoribbons (Pt/GNRs)***

The Pt/GNR NPs were produced by a one-pot synthesis from the GONRs, in a process based on previous reports [28]. First, 12 mg of the GONRs was dispersed in 12 mL water in an ultrasonic bath for 45 min. Then, 11.5 mg poly(acrylic acid) salt (PA,  $M_w = 2100$ ) was added, and the dispersion was sonicated for at least 5 min. Any variation of this protocol should be performed while taking into account the ratio  $PA/Pt_{atoms} = 3$ . Next, 2 mL of  $H_2PtCl_2$  solution (10.6 mg/mL) was added to produce Pt/GNRs with a 40% metal loading, and the mixture was sonicated for 1 min.

The mixture was heated to 95°C and purged with ultrapure  $N_2$  for 10 min. Next,  $N_2$  was removed from the solution and kept in the atmosphere of the reaction bath. Then 32 mL of  $NaBH_4$  0.5 mol  $L^{-1}$  was added, and the reaction bath was held at 95°C in a  $N_2$  atmosphere for 60 min. The reaction bath was allowed to spontaneously reach room temperature, and then the dispersion was washed with ultrapure water and centrifuged three times. Finally, the dispersion was dried for 24 h at 60°C to produce the nanoparticle powder.

## ***2.3 Electrode preparation and electrochemical measurements***

The dispersions of Pt/GNRs (ink) were prepared by adding 2 mg of powder to 1 mL water. To prepare the electrodes, 15  $\mu L$  of the ink was deposited onto a 0.24  $cm^2$  Au electrode, in order to produce a catalytic surface with 50  $\mu g$   $cm^{-2}$  loading (taking into

account the mass of Pt on the Au electrode). To prepare the Pt/C (E-TEK, 40%) electrodes, we followed the same protocol, adding 10  $\mu\text{L}$  of highly diluted Nafion<sup>®</sup> (1 mL 5% Nafion<sup>®</sup> in 20 mL 2-propanol) to the NPs immobilized on Au; this step was taken because the Pt/C NPs do not adhere as strongly to Au as do the Pt/GNRs. Prior to immobilizing the NPs on Au and start the measurements, we carefully followed an electrochemical protocol to guarantee reproducibility and reliability, as detailed in Supplementary Information, Section I. Briefly, the Au disk was electrochemically cleaned and its electrochemical profile was registered. Fig. S1 shows the characteristic profile of the bare Au. After the immobilization process, the Pt/C NPs were also electrochemically cleaned [32] and their profile recorded, as shown in Fig. S2.

The electrochemical experiments were carried out in a conventional three-electrode cell in an oxygen-free 0.5 mol L<sup>-1</sup> H<sub>2</sub>SO<sub>4</sub> solution using a potentiostat system (Potentiostat/Galvanostat  $\mu$ Autolab with current integration). A Pt plate with high surface area was used as counter electrode. The modified Au disk was used in a meniscus configuration as the working electrode. All potentials were measured against a reversible hydrogen electrode (RHE). The current densities ( $j$ ) were calculated as the measured current divided by the electrochemically active surface area (ECSA), assuming 210  $\mu\text{C}$  per square centimeter of Pt surface as the charge released by desorption of a hydrogen monolayer. All electrochemical experiments were performed in triplicate at 25°C and all measurements were performed at least three times, and all parameters are given as averages with standard deviations presented as error bars to provide reliable information. To assess the electrocatalytic performance of the catalysts, we determined the onset potential and the potential peak from the first derivative of the voltammetry signal [31] and the current density peak directly from the voltammogram.



#### ***2.4 Evaluation of biomass-derived electrooxidation and stability tests***

We investigated methanol (MeOH), ethanol (EtOH), and glycerol (GIOH) electrooxidation on the Pt/GNRs and on 40% Pt/C E-TEK. The most important experimental detail is that the concentrations of the alcohols were selected to contain the same number of carbons ( $0.6 \text{ mol L}^{-1}$  MeOH,  $0.3 \text{ mol L}^{-1}$  EtOH, and  $0.2 \text{ mol L}^{-1}$  GIOH). This experimental feature guarantees a reliable comparison among the output current densities, potentials (onset and peak) and electrochemical stabilities. It is well-known that the chemical environment, as pH, temperature and concentration of reactants highly influences the pathways of surface electrochemical reaction. In this sense, we followed exactly the same condition, at least the maximum as possible, to compare the electrocatalytic parameters of the MeOH, EtOH and GIOH electrooxidation on Pt/GNRs and Pt/C.

The electrochemical responses were measured in a solution containing  $0.5 \text{ mol L}^{-1}$   $\text{H}_2\text{SO}_4$  and  $X \text{ mol L}^{-1}$  alcohol (where X is the alcohol concentration) at  $0.05 \text{ V s}^{-1}$  between 0.05 and 1.0 V. The starting potential was set to 0.12 V for all experiments to avoid the reduction of GIOH to propane [33] during the standby period. The representative feature of electrooxidation is shown as the fifth cycle, referred here as the stationary behavior. The onset potential determined from the first derivative [31] (not shown) was averaged from three independent measurements. Chronoamperometry experiments were performed by polarizing the electrodes at 0.12 V and stepping the potential to 0.6 V for 1800 s in the presence of the alcohol. The pseudo-stationary current densities are showed as an average of three independent measurements.

No standard protocol exists for evaluating the stability of potential anodes for direct alcohol fuel cells. A number of different methods have been reported in the

literature. Chronoamperometry is often used to evaluate the stability of a catalyst toward the production of an electrochemical current over time [34], such method does not mimic the start-up and shut-down processes found in fuel cells. To overcome this issue, one option is to measure the decay in electrochemically active surface area in the presence of the electrolyte during several cyclic voltammetry cycles over a wide range of potentials [9,17,35]. Although reliable, this approach is not ideal because it is usually performed in absence of fuel. We believe that the current density of the anodic process must be measured [36] concomitantly with the surface area. Here, we measured the current densities during the alcohol electrooxidation at a given potential over 1000 cyclic voltammetry cycles between 0.3 and 1.0 V at 0.3 V s<sup>-1</sup> (in alcohol and electrolyte). Moreover, we recorded a profile (between 0.05 and 1.2 V at 0.05 V s<sup>-1</sup>) in the presence of only electrolyte before and after the stability tests, which allowed us to determine the surface changes caused by the application of electric potential in the presence of fuel.

### ***2.5 Physical characterization of Pt/GNRs***

The chemical composition of the Pt/GNRs was determined using a scanning electron microscope (Phenom ProX) connected to an energy dispersive X-ray spectrometer (EDS). The porosities of the GONRs and the Pt/GNR NPs were measured by physical adsorption of N<sub>2</sub> at -196°C with an adsorption analyzer (Micromeritics Tristar 3020) after degassing the samples at 120°C and 0.1 mbar for 8 h using a degasser (Micromeritics VacPrep 061). The following parameters were obtained: surface area ( $S_{\text{BET}}$ ) and micropore volume ( $V_{\text{micro}}$ ), calculated with the Brunauer-Emmett-Teller (BET) [37] and Dubinin-Radushkevich (DR) equations [38] respectively; total pore volume ( $V_{\text{p}}$ ), estimated from the amount of N<sub>2</sub> adsorbed at  $p/p^\circ$

= 0.99; and pore size distribution (PSD), determined by a density functional theory (DFT) calculation.

The crystallographic structures of the GONRs and Pt/GNRs were investigated by X-ray diffraction (XRD), using a Bruker D8 powder diffractometer equipped with a monochromatic Cu-K $\alpha$  X-ray source; scans were performed with a step size of 0.02° in the 5-90° range and an interval of 2 s between steps. The disorder of the carbon supports was determined by Raman spectroscopy using a LabRAM HRUV 800 (JYV-Jobin Yvon) with a microscope (Olympus BXFM-ILHS) and an argon laser (CDPS532M, 532 nm at 24.3 mW). The analysis was performed between 800 and 3500 cm<sup>-1</sup> with two 12-s accumulations.

The reduction of the GONRs to form GNRs was analyzed by ultraviolet spectroscopy. For the measurement, 50  $\mu$ L of the 2 mg/mL dispersion was diluted to 5 mL, and the ultraviolet absorption was measured between 200 and 800 nm; the response is shown as an average of six measurements.

The morphologies and sizes of the nanoparticles and carbon supports were analyzed using a transmission electron microscope (TEM) (JEOL JEM2100) with a LaB<sub>6</sub> filament operating at 200 kV. Samples were prepared for TEM analysis by dispersion in 2-propanol and deposition onto a 400-mesh copper grid. The images were evaluated using the software Axio Vision SE64 Rel.4.8.

The wettability of the catalysts, which is related to the hydrophobic or hydrophilic interaction between the electrolyte + fuel solution and the carbon supports, was investigated by measuring the contact angle. We used a 4  $\mu$ L droplet of the solution containing the electrolyte and fuel, holding constant the total number of carbons in

solution for MeOH, EtOH, and GIOH to avoid a significant contribution from the viscosity of the alcohols. The droplet was placed on a thin film (made with 9  $\mu$ L of the inks) and dried overnight on a glassy carbon plate (previously polished to a mirror finish). The detailed experimental procedure is described in Fig. S3.

### 3. Results and Discussion

#### 3.1 Physical characterization: Production of Pt/GNRs

The chemical composition of the Pt/GNRs determined from an average of eight measurements was  $38\pm 3\%$  metal loading on carbon (% weight of Pt). Hence, the synthesized and the commercial material have approximately the same amount of Pt on carbon, which diminishes the contribution of the metal loading on the comparison of activity and stability.

The production of the Pt NPs-modified GNRs from the GONRs via one-step synthesis was evaluated by Raman and ultraviolet spectroscopy, as shown in Fig. 1. As depicted in Fig. 1A, both the GONRs and the Pt/GNRs exhibit D and G bands at  $\sim 1332$  and  $\sim 1595$   $\text{cm}^{-1}$  respectively. The D band can be attributed to either disorder or  $\text{sp}^3$  carbon hybridization, and the G band is attributed to energy level splitting associated with the  $\text{E}_{2g}$  stretching mode of graphite [39]. Since a defect-free graphite structure should not have a D band, the structures of the GONRs and Pt/GNRs must exhibit some disorder. The increase in the degree of disorder was determined by measuring the change in the ratio  $I_D/I_G$  [28,40-42]. Fig. 1A clearly shows that the  $I_D/I_G$  ratio decreases from 1.35 to 1.05, which indicates augmentation of the G network after the reduction of superficial groups containing oxygen. The second harmonic of the D band, referred to

as the 2D band [43], appears as a shoulder at  $\sim 2680\text{ cm}^{-1}$  in both materials due to the graphite-like characteristics.

### Figure 1

The ultraviolet absorption spectrum shows a bathochromic shift from 239 nm for the GONRs to 258 nm for the Pt/GNRs (Fig. 1B), indicating recovery of the electronic conjugation of the nanoribbons. The ultraviolet absorption spectrum is consistent with the literature [23] and with the results from Raman spectroscopy. The results of the structural characterization by XRD are shown in Fig. 2. The two diffraction peaks at  $25.0^\circ$  and  $45.0^\circ$  correspond to the (002) and (100) planes of the GONRs. The strong diffraction peaks at  $\sim 40.0^\circ$ ,  $45.5^\circ$ ,  $67.5^\circ$ ,  $81.5^\circ$ , and  $86.0^\circ$  for the Pt/GNR material are assigned to the (111), (200), (220), (311), and (222) facets of the face-centered structure of the Pt crystallites, respectively. The reduction of the GONRs to flat carbon surfaces containing metal nanoparticles hinders the C (100) diffraction peak and both decreases the intensity and broadens the width of the C (002) peak. Broad and low-intensity peak between  $10.0^\circ$  and  $37.0^\circ$  was previously reported for GONRs treated with  $\text{NaBH}_4$  [28] and for electrochemically produced Pt NPs shell decorating graphene (Pt NPs@G) from Pt NPs@GO [44]. The presence of Pt oxides [45,46] and residual graphene oxides results in several low intensity peaks between  $10.0^\circ$  and  $37.0^\circ$ .

### Figure 2

The  $\text{N}_2$ -adsorption isotherms of the catalysts are shown in Figs. S4 and S5. Both GONR and Pt/GNR materials can be classified as type IV, according to BDDT (Brunauer, Deming, Deming, and Teller) classification [47], with a hysteresis loop at high relative pressures indicating the presence of mesopores (see Fig. S4). Both isotherms have similar shapes at high relative pressures, indicating similar pore sizes, as

shown by the pore diameters ( $d_{\text{pore}}$ ) in Table 1. However, the materials adsorb different volumes of  $\text{N}_2$ , especially at low relative pressures. These observations are corroborated by the porosity parameters shown in Table 1.

**Table 1.** Porous characteristics of GONRs and Pt/GNRs.

Sample	$S_{\text{BET}}$ ( $\text{m}^2 \text{g}^{-1}$ )	$V_{\text{micro}}$ ( $\text{cm}^3 \text{g}^{-1}$ )	$V_{\text{p}}$ ( $\text{cm}^3 \text{g}^{-1}$ )	$V_{\text{meso}}$ ( $\text{cm}^3 \text{g}^{-1}$ )	$d_{\text{pore}}$ (nm)
Pt/GNRs	26.00	0.01	0.11	0.10	27.00
GONRs	205.00	0.07	0.41	0.34	25.00

Among all the parameters investigated here, the most affected by the reduction process and the deposition of the Pt NPs was the specific area ( $S_{\text{BET}}$ ). The GONRs are composed of structures covered by functional groups that cause them to repel each other. After reduction, the loss of these groups produces flat structures that tend to stack on top of each other due to strong van der Waals interactions, diminishing the surface area available for  $\text{N}_2$  adsorption. Another factor that contributes to the decrease in  $S_{\text{BET}}$  is the deposition of the Pt NPs. The NPs cover the ribbons, which blocks the penetration of  $\text{N}_2$  into some regions of the ribbons.

The decrease in the mesopore volume (or increase in the size of the pores) is related to the decrease in the surface area of carbon in contact with the electrolytic solution, which is a crucial parameter in electrochemistry. Both materials are highly mesoporous, as shown by the micropore volume ( $V_{\text{micro}}$ ) and mesopore volume ( $V_{\text{meso}}$ ). The GONRs show considerable microporosity (pores  $< 2$  nm) which leads to a large specific area ( $\sim 205 \text{ m}^2/\text{g}$ ). The pore size distribution indicates that the porosities of the surfaces of both the GONRs and the Pt/GNRs are mostly in the mesoporous range (2-50 nm), and also indicates that the pore volume of the GONRs is high (Fig. S5). During reduction of the GONRs and deposition of the NPs, the same factors that diminish the specific area also diminish the porosity.

### ***3.2 Microscopic analysis: From MWCNTs to Pt/GNRs***

The synthesis of the Pt/GNRs from the precursor MWCNTs was observed by TEM. The unzipping of the nanotubes is expected to increase the average diameter of the support as the nanoribbons begin to form. Figs. 3A and 3D show the TEM image and average diameter ( $9.4\pm 2.3$  nm) for the MWCNTs. The value for the average diameter is in line with the value reported by the company providing the nanotubes ( $10\pm 1$  nm). The average diameter of the support particles increased to  $17.4\pm 3.9$  nm after the oxidation process, when the GONRs were formed (Figs. 3B and 3E). The  $\sim 8.0$  nm increase in the average diameter of the support proves the efficiency of the method for unzipping the nanotubes. The relatively large standard deviation is a consequence of the non-uniform nanoribbons, since the outer and inner nanotubes of the MWCNTs have different diameters. Additionally, the partial unzipping of some nanotubes results in an increase in the heterogeneity, and thereby increases the standard deviation.

Fig. 3C shows the Pt NPs immobilized on the GNRs. The NPs had an average diameter of  $5.0\pm 0.9$  nm, and mostly formed in small clusters. Hence, the average size and size distribution (Fig. 3F) were calculated from well-defined NPs, which could be either isolated NPs or NPs formed in clusters. The deposition of the Pt NPs on the flat surfaces of the nanoribbons generates some defects on the surfaces, which work as nucleation centers for clustering (NCCs), as previously described for Pd deposition on chemically reduced graphenes [17]. As a result, the continuous deposition of Pt on the NCCs leads to the formation of small clusters. The reduced nanoribbons had an average diameter of  $17.0\pm 2.5$  nm, which indicates that the size of the support is not affected by the reduction process (Fig. S6). Fig. S6 shows that the Pt NPs are more uniformly distributed on the carbon support than they are on the GNRs with few regions of

agglomeration. Moreover, the average size of the Pt particles was  $3.6 \pm 0.7$  nm, with sizes ranging from 2 to 7 nm.

### Figure 3

#### 3.3 Contact angle (wettability) measurements

The wettability of the carbon material is determined by the nature (chemical composition) and structure (roughness) of the material [48]. Hence, the wettability is intrinsically related to the electroactivity of a catalyst, since the reaction depends on the fuel and electrolyte having access to the active sites.

In general, a surface is defined as hydrophobic if the contact angle is  $> 90^\circ$ , or hydrophilic if the contact angle is  $< 90^\circ$ . In this study, we minimized the contribution of the viscosity to the contact angle measurements by maintaining a constant number of carbons in the solution for all fuels. The average contact angles of the Pt/C and Pt/GNR catalysts for MeOH, EtOH, and GIOH in  $H_2SO_4$  solution are given in Fig. 4.

### Figure 4

Both the Pt/C and Pt/GNR catalysts are hydrophilic in all the alcohols solutions, as shown by the  $< 90^\circ$  angles. As expected, the hydrophilicity in MeOH was higher than that in EtOH due to the smaller organic chain of methanol and as a consequence of the non-oxidized carbon in ethanol ( $-CH_3$ ). Comparing GIOH with EtOH, the contact angles reveal that the three hydrated carbons in GIOH ( $HOCH_2-HCOH-H_2COH$ ) facilitate the interaction with the catalysts, increasing their hydrophilicity. Pt/C surface in GIOH is slightly less hydrophilic than that in MeOH and more hydrophilic than Pt/GNR surface in GIOH. This observation suggests that GIOH (in  $H_2SO_4$  solution) interacts well with GNRs compared with MeOH and EtOH. Interestingly, the Pt/GNRs had a higher contact angle than that of Pt/C for all fuels. The low hydrophilicity of the GNRs is a



consequence of the extended  $C_{sp^2}$  network containing only a small amount of oxides on the surface, as well as the low surface area ( $26 \text{ m}^2 \text{ g}^{-1}$ ) compared to that of the commercial Pt/C material, which is  $\sim 150 \text{ m}^2 \text{ g}^{-1}$ .

Now that the physical-chemical characterization of Pt/GNRs was presented, we dedicate the next sections to thoroughly investigate the activity and stability of Pt/GNRs compared to Pt/C for the alcohols electrooxidation.

### ***3.4. Comparing the activities of biomass-derived electrooxidation on Pt/GNRs***

As can be seen in the stable methanol oxidation electrochemical profile in Fig. 5A, the Pt/GNRs produced more current density than the Pt/C catalyst at potentials higher than 0.73 V during the forward scan and over the entire range of potentials during the reverse scan. The peak current of the Pt/GNRs was  $\sim 2.3$  times higher than that of the Pt/C. Chronoamperometry measurements (not shown) during 1800 s of polarization at 0.6 V (with a starting potential of 0.12 V) in the presence of MeOH showed that the average pseudo-stationary current density of the Pt/C was 1.5 times higher than that of the Pt/GNRs ( $0.057$  and  $0.038 \text{ mA cm}^{-2}$ , respectively). The potentiostatic experiments corroborate the potentiodynamic measurements that the Pt/GNRs are more active only in higher potentials, since it was performed at 0.6 V. The average onset potential calculated from the first derivative of the voltammetry signal [31] ( $dj/dE$ , not shown) was 0.66 V for the Pt/GNRs and 0.60 V for the Pt/C.

The increase in the current density of MEOR on the Pt/GNRs in the potentiodynamic experiment is in line with previous works investigating MEOR on graphene-based surfaces [28,34,35,44]. Looking at the hydrogen under potential deposition  $H_{UPD}$  region (blank profiles) in Fig. 7, we can conclude that the NPs are

essentially polycrystalline, and the Pt surface arrangements are very similar for both the Pt/GNR and Pt/C materials. Therefore, the important differences in the potentiodynamic behavior after 0.73 V are not caused by a different surface orientation of the metallic catalyst. The contribution of the substantial length of the nanoribbons to the electronic nature of Pt is probably the reason for the increase in the current densities, which is evidenced at potentials higher than 0.7 V. This result is in line with previous investigations of flat supports [34,35,44], although the onset potential is slightly delayed.

The same experimental protocol used to investigate MEOR was used for the EEOR analysis, but the alcohol concentration was adjusted to maintain the same number of carbons in solution (Fig. 5B). Both onset and peak potential are almost the same. The peak potentials in the forward scans were virtually the same for both catalysts. During the reverse scan, the anodic current is larger for the Pt/GNRs.

Chronoamperometric measurements (at 0.6 V for 1800 s) showed average pseudo-stationary current densities of 0.043 and 0.033 mA cm<sup>-2</sup> for the Pt/GNRs and the Pt/C, respectively. The slightly increased activity of the Pt/GNRs in the potentiostatic experiments is a consequence of the slight lower onset potential, since the voltammograms did not show any remarkable improvement during the forward potential scan.

### Figure 5

Cyclic voltammograms in GLOH show comparable features for both catalysts (Fig 5C). The onset potentials were virtually equal, 0.66 V and 0.65 V for the Pt/GNRs and the Pt/C, respectively. The average pseudo-stationary current densities were also approximately equal (0.018 and 0.017 mA cm<sup>-2</sup> for the Pt/GNRs and the Pt/C

respectively, after 1800 s at 0.6 V). The most relevant difference is in the peak potential of GEOR during the forward scan, which for the Pt/GNR material was located at a potential 40 mV lower than that of the Pt/C.

### ***3.5. Comparing the stabilities of biomass-derived electrooxidation on Pt/GNRs***

The stabilities of the catalysts, determined by measuring the current peak during 1000 potential cycles, are shown in Fig. 6. The Pt/GNRs are remarkably stable toward the MEOR reaction (Fig. 6A). After the degradation test protocol, the Pt/GNRs lose 15% of their initial activity, whereas the commercial material loses 43%. This result indicates a 28% improvement in the performance of the Pt/GNRs compared to Pt/C.

The Pt/GNRs displayed an increased electrochemical stability relative to that of the Pt/C for EEOR (Fig. 6B). The Pt/GNRs lose 21% of their initial activity, while the Pt/C loses 32%. However, the large standard deviation (error bars) in Fig. 6B suggests that similar performance might be achieved using Pt/C for EEOR, especially in short-term. Regarding the GEOR, we found that the Pt/GNRs lose 38% of their initial activity compared to 49% of loss for Pt/C, as shown in Fig. 6C. It is worth noting that the stability of the Pt/GNRs is remarkably reproducible (as shown by the small standard deviation).

The stability was also evaluated by monitoring the loss of ECSAs before (solid lines) and after (dashed lines) the successive cycles of biomass-derived electrooxidation, as shown in Fig. 7. The ECSAs obtained from the  $H_{UPD}$  region (Figs. 7A and 7B) change significantly after the stability tests in presence of MeOH. ECSAs decrease  $\sim 20\%$  for both Pt/GNR and Pt/C materials. Interestingly, the increase in the size of Pt ( $\Delta d(Pt)$ ) was  $\sim 0.7$  and  $\sim 0.6$  nm for the Pt/C and Pt/GNRs respectively (Figs.

7C and 7D). Histograms of the mean diameters calculated after the stability tests are shown in Fig. S7.

The increase in the size of the Pt NPs is not necessarily associated with the decrease in ECSA, since the particles are agglomerated from the beginning. Another important factor that can contribute to the decrease in ECSA is the low stability of carbon Vulcan compared to nanoribbons. It is important to note that even though both electrodes lose about 20% of their ECSA after MEOR, the decrease in the activity of the Pt/C was larger than that of the Pt/GNRs (Fig. 6A). In other words, the current density was negatively affected to a greater degree for the Pt/C than for the Pt/GNRs, which indicates that the degradation process not only affects the ECSA but also modifies the carbon surfaces as a consequence of carbon corrosion.

### Figure 6

Figs. 7E and 7F show the ECSA of the catalysts before and after EEOR. It is clear that the decrease in ECSA is greater for the Pt/C than for the Pt/GNRs, as a consequence of the low average stability. As observed in the MEOR experiments, the  $\Delta d(\text{Pt})$  values of the Pt/GNR and Pt/C materials in the EEOR experiments were similar (Figs. 7G and 7H). However, the values were larger than that for MEOR. This evidence suggests that the composition of the solution has an influence on the stability of the catalyst. Since the  $\Delta d(\text{Pt})$  values are similar, the decrease in ECSA can be attributed to the detachment of the Pt NPs from the carbon surface, as was previously reported by identical-location TEM in absence of fuel [2,49].

The loss of area was also greater for the commercial material after GEOR, as shown in Figs. 7I and 7J. The micrographs reveal that the  $\Delta d(\text{Pt})$  of the Pt/GNRs was 1.7 nm, while the  $\Delta d(\text{Pt})$  of the Pt/C was 0.7 nm (Figs. 7K and 7L). The agglomeration

effect in the Pt/GNRs causes the decrease in electrochemical stability. Here we note that even though the particles were more agglomerated on the GNRs than on the carbon at the beginning of the experiment, the agglomeration was more intense on the GNRs, indicating that the presence of GIOH in the electrolyte has a different effect on the stability of Pt immobilized on different supports.

### Figure 7

Considering the low  $\Delta d(\text{Pt})$  of Pt/C, we infer that the detachment of Pt NPs, which was mainly caused by oxidation of the carbon support, added to the strong poisoning of the well-distributed Pt over the carbon black, were the main reasons for such a high loss of activity (49%) in the long-term experiments.

Here, we point out that we tentatively estimated the average width of the nanoribbons after the degradation test protocol for all reactions investigated, as shown in Fig. S7. Even though our estimative is quite rough as evidenced by the large standard deviation, we can infer that the widths of the nanoribbons were not significantly affected by the tests, although the chemical nature of the nanoribbons might be affected to some extent. Nevertheless, understanding the change in the electronic state and chemical composition of the GNRs is beyond the scope of this work. In the next section, we summarize the results of the Pt/GNRs for biomass-derived alcohol electrooxidation.

### ***3.6 Evaluating the activity and stability of Pt/GNRs toward biomass-derived alcohols***

Fig. 8 shows the parameters related to the activity and electrochemical stability of the Pt/C and the Pt/GNRs toward MeOH, EtOH, and GIOH electrooxidation reactions under exactly the same experimental condition. Comparing the onset potentials for both catalysts in biomass-derived electrooxidation (Fig. 8A), we find that

the EtOH electrooxidation reaction is clearly the least favorable. The non-oxidized  $-CH_3$  makes C-C cleavage difficult. Moreover, the hydrophobicity caused by  $-CH_3$  might obstruct the penetration into the active sites of the catalyst. Among all the alcohols, MeOH is the most likely to be oxidized. Comparing the Pt/C and the Pt/GNRs, we only found a slight difference in presence of MeOH.

Another electrochemical parameter that helps to identify the activity of materials is the potential peak [31]. The potential peaks show similar values for MEOR and EEOR on Pt/GNRs and Pt/C NPs (Fig. 8B). GEOR shows lower potential peaks and the Pt/GNRs shift the potential peak towards more negative values; although the onset potential is higher.

The current density peaks ( $j$ ) determined from the first oxidation peak (Fig. 8C) indicate the activity under the influence of potential transients ( $dE/dt$ ). The lowest values were found for GEOR, for which the Pt/GNRs and the Pt/C showed similar features. EEOR showed a higher  $j$  value than GEOR, also with similar values. Although more electrons might be extracted from EtOH than from EtOH, the sequential pathways of GEOR, involving several partially oxidized compounds, decrease the  $j$  value for GEOR on Pt surfaces, even considering that the same amount of carbon participates in the reaction [50].

MEOR exhibits the highest current densities, as a consequence of the diminished number of partially oxidized compounds and due to the more facile oxidation, since the fuel contains only one carbon in the main. Interestingly, a very large increase in the  $j$  value was observed in the potentiodynamic experiment for MEOR on the Pt/GNRs compared to that for MEOR on the Pt/C, as a consequence of the improvements under elevated applied potential, since this peak appears at 0.95 V. This observation indicates

that MEOR produces more output current than EEOR and GEOR in high potentials by using Pt/GNRs as catalyst.

The interpretation of the potentiostatic experiments using the pseudo-stationary  $j$  values, in Fig 8D, reveals the weak performance of both catalysts for GEOR, which is consistent with the potentiodynamic experiments. EEOR displays higher  $j$  values than GEOR, and the Pt/GNRs showed higher  $j$  values than the Pt/C for EEOR. MEOR showed the highest  $j$  values, and the Pt/C was found to be the best catalyst in terms of the pseudo-stationary  $j$  value. Thus, with regard to MEOR, although the Pt/GNRs showed the highest  $j$  values during the potential sweep in high potentials, Pt/C showed better performance when the material experienced conditions similar to those of a fuel cell working under a low potential (onset).

Since the  $\text{H}_2\text{SO}_4$  concentration was the same for all reactions, the decay of the ECSA was less dramatic when the catalysts were subjected to potential cycles in the presence of GIOH, as shown in Fig. 8E. MEOR represents the most unfavorable electrochemical environment, while EEOR presents intermediate values. The different products and co-products produced in the different reactions change the interfacial pH accordingly [51], thereby modifying the electrochemical environment, which leads to different stabilities in terms of ECSA loss. Thus, the stability of a material highly depends on the surface reaction.

Theoretical calculations predict weak interaction between flat surfaces and metallic NPs [22] similar to those expected in the Pt/nanoribbons, so it is reasonable that the  $d$  band of Pt is more available for react than the Pt immobilized on carbon black. This prediction, in addition to the high conductivity of the long double-bonded network, leads us to expect an improvement of activity in alcohols electrooxidized by Pt

NP-modified nanoribbons. However, as reported by Brownson *et al.*, graphenes might not be as electrochemically advantageous as they seem [52]; some characteristics might be desirable, but the overall performance might not be adequate for use in fuel cells.

The effectiveness of EtOH as a fuel is more strongly dependent than MeOH on the improvement in performance by multi-metallic catalysts. Assessing effectiveness of GIOH in fuel cells remains complicated, but the possibility of producing high-value compounds concomitantly with energy through the use of adequate catalysts [53,54] increases the level of interest in such alcohol.

### Figure 8

EtOH and GIOH seem not affected by the Pt-support interaction. Thus, the low activity of Pt/GNRs in EEOR and GEOR in acid media might be caused by the limited penetration of the fuel into the nanoribbon structures, caused by the  $sp^2$  network as well as some wrapped ribbons attached to the small clusters of Pt on the nanoribbons.

In general, the stability of the Pt/GNRs in long-term experiments is improved compared to that of the Pt/C for all the biomass-derived alcohols. Nanoribbons assist Pt NPs in maintaining the activity through several cycles of use by preventing corrosion of the support and loss of surface area.

Summarizing, Pt/GNRs are advantageous materials to be used as anodes in direct alcohol fuel cells due to the long-term stability of the support, independent of the alcohol used. Therefore, the use of multi-metallic Pt-based catalysts immobilized on nanoribbons is a potential alternative for improving the electroactivity while maintaining the high electrochemical stability of the anode.

## 4. Conclusions



We successfully synthesized Pt-modified graphene nanoribbons (Pt/GNRs) from graphene oxide nanoribbons (GONRs) by a  $\text{NaBH}_4$  one-step method assisted by poly(acrylic acid) salt. The Pt nanoparticles (NPs) were distributed in small clusters on the long flat surfaces. The average size of the individual NPs was approximately  $5.0 \pm 0.9$  nm (with sizes ranging from 2-9 nm), and the NPs were dispersed over nanoribbons with an average width of  $17.4 \pm 3.9$  nm.

Comparing the electroactivity of the Pt/GNRs with that of a commercial Pt/C in acid media, taking into account the same experimental conditions including the number of carbons in the solution, we found that the ethanol and glycerol electrooxidation reactions performed equivalently on both catalysts. Only the methanol electrooxidation reaction showed improved activity to some extent.

On the other hand, graphene nanoribbons substantially improve the ability of the catalyst to maintain an electrochemical response, and also improve the electrochemical stability, since GNRs prevent loss of electrochemically active surface area by preventing carbon corrosion and Pt detachment. Moreover, we found that the stability of the nanoparticles depends on the biomass-derived alcohol used, which is a variable rarely explored in the field. Among the reactions investigated, methanol electrooxidation in acid media appears to be the most aggressive environment for Pt nanoparticles supported either on carbon or on nanoribbons while glycerol media is less aggressive. Therefore, the different products formed at the interface electrode / solution might lead to different electrochemical environment, which plays an important role on the stability of the catalysts.

Considering that the electroactivity toward biomass-derived electrooxidation can be tuned by alloying Pt with ad-atoms, GNRs are promising supports for multi-metallic

nanoparticles used as anodes in direct alcohol fuel cells fed by MeOH, EtOH or GlOH. The use of an ad-atom might improve the activity while GNRs improve the catalyst stability.

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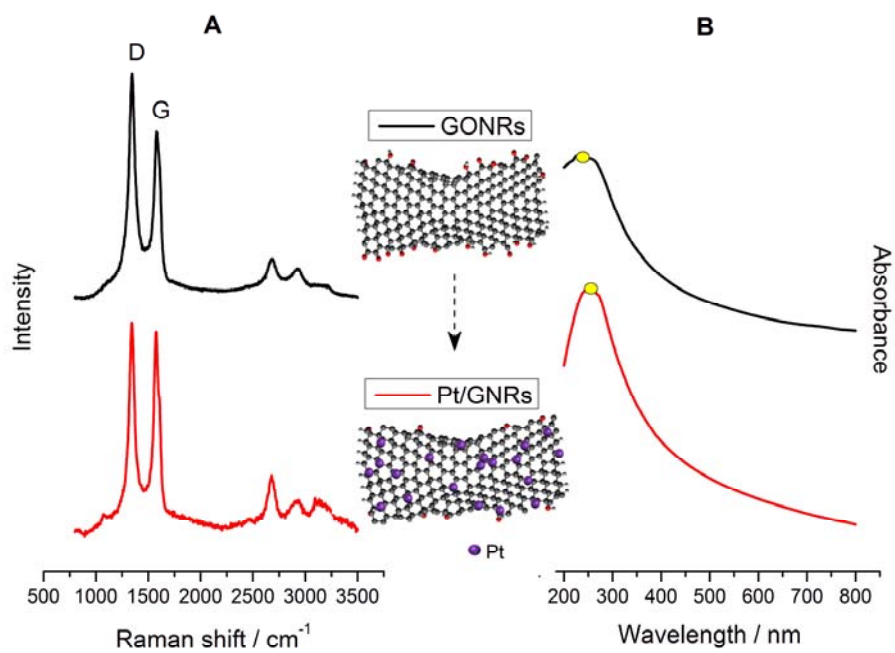
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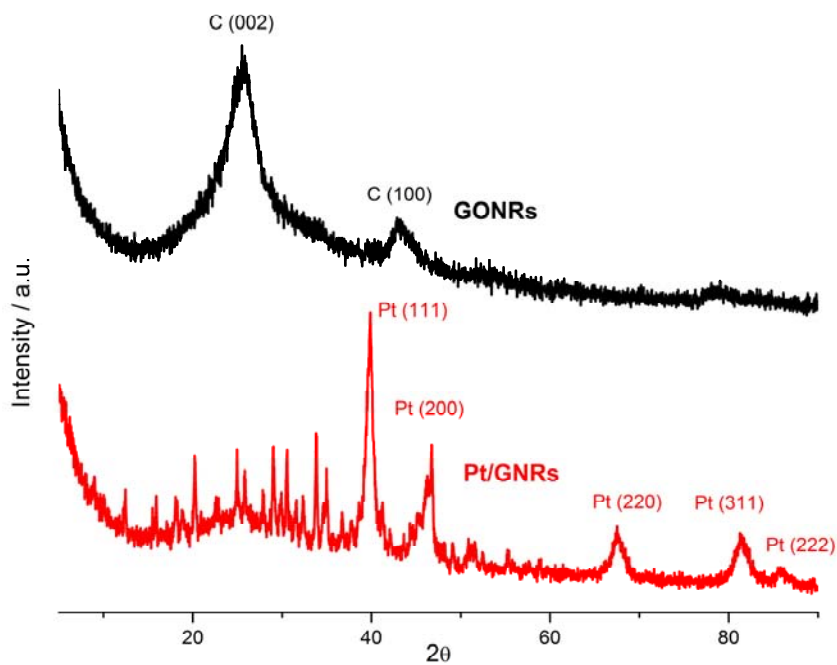
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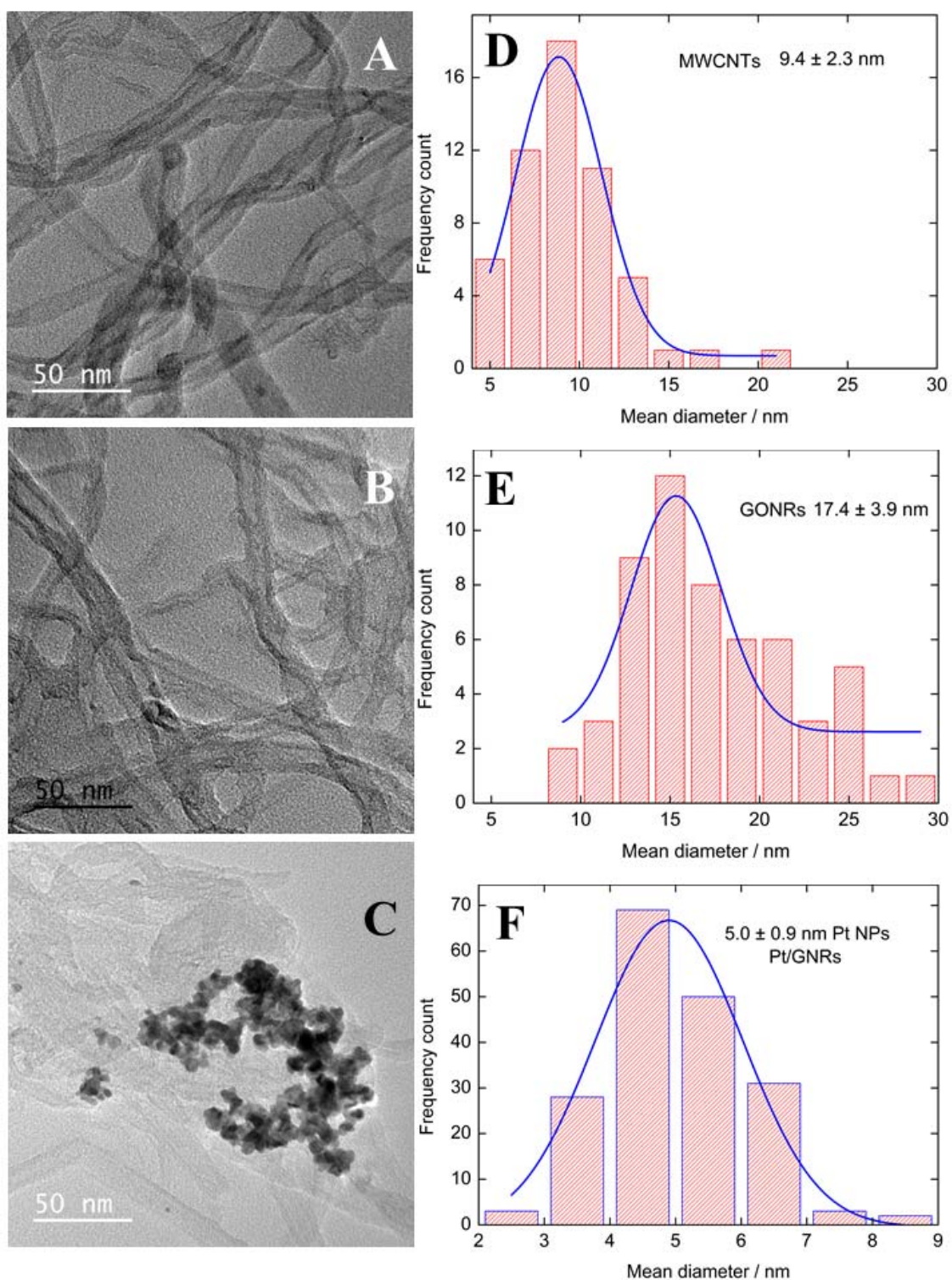
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**Figure 1.** (A) Raman spectra of GONRs and Pt/GNRs and (B) Ultraviolet spectrum of an aqueous solution of GONR (black,  $\lambda_{\text{max}}$  239 nm) and Pt/GNRs (red,  $\lambda_{\text{max}}$  258 nm).



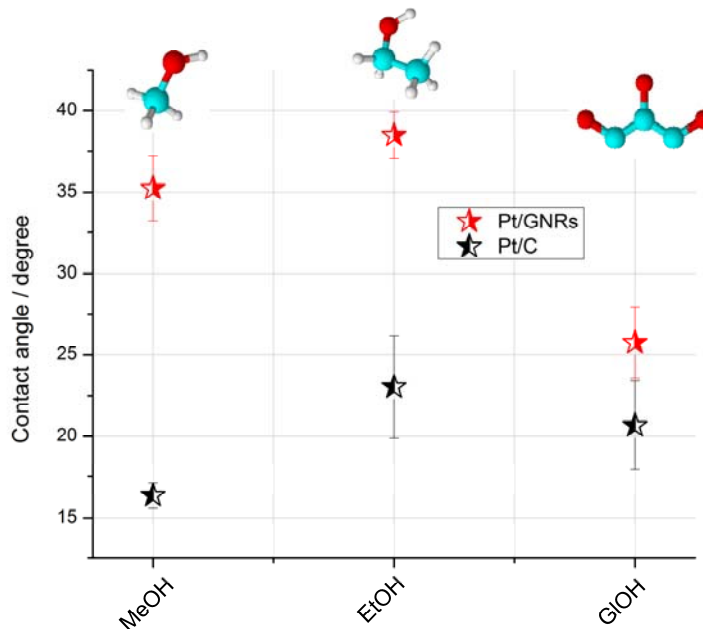
**Figure 2.** XRD patterns of GONRs (black) and Pt/GNRs (red).



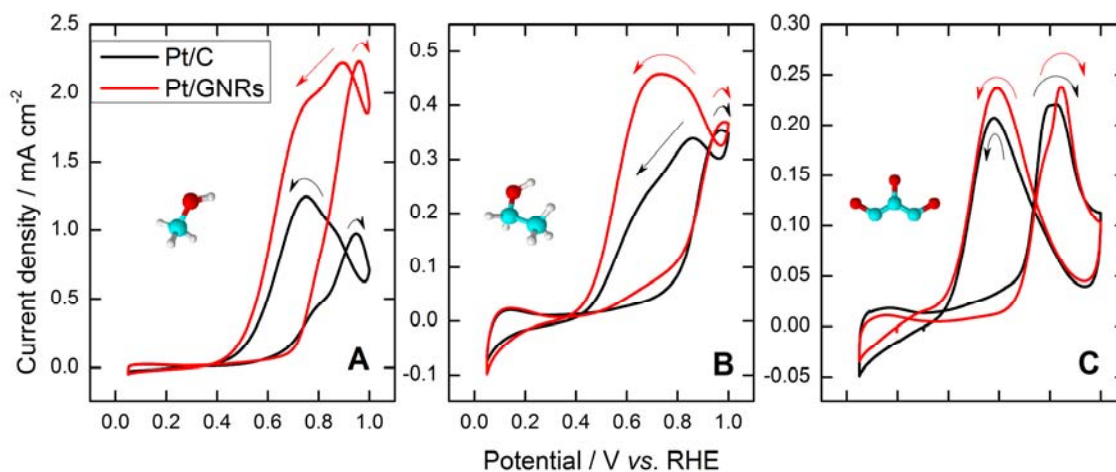
**Figure 3.** TEM investigation of the synthesis of Pt/GNRs NPs. Micrographs and mean size diameter distribution of MWCNTs (A and D respectively). Micrographs and mean



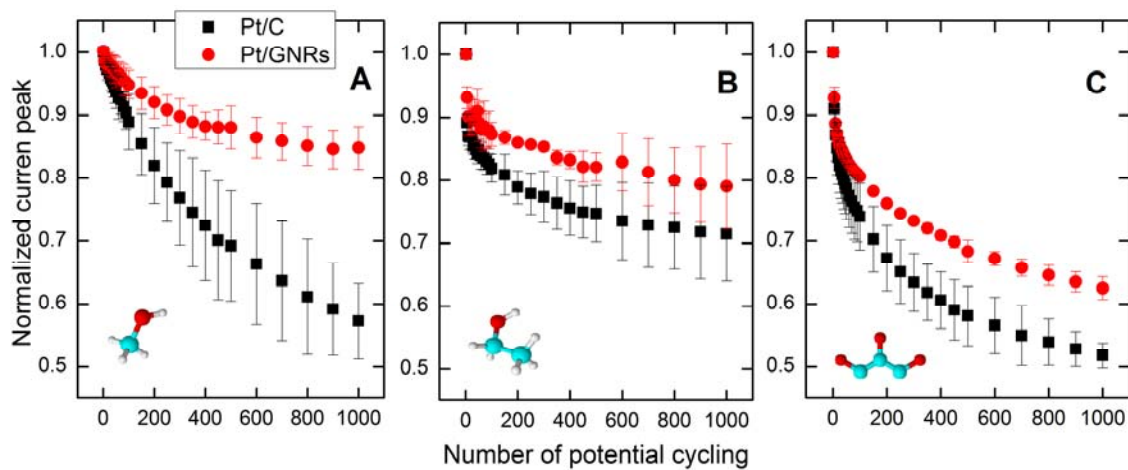
diameter distribution of GONRs(B and E respectively). Micrograph of Pt/GNRs (C) and mean size distribution of Pt immobilized on GNRs (F).



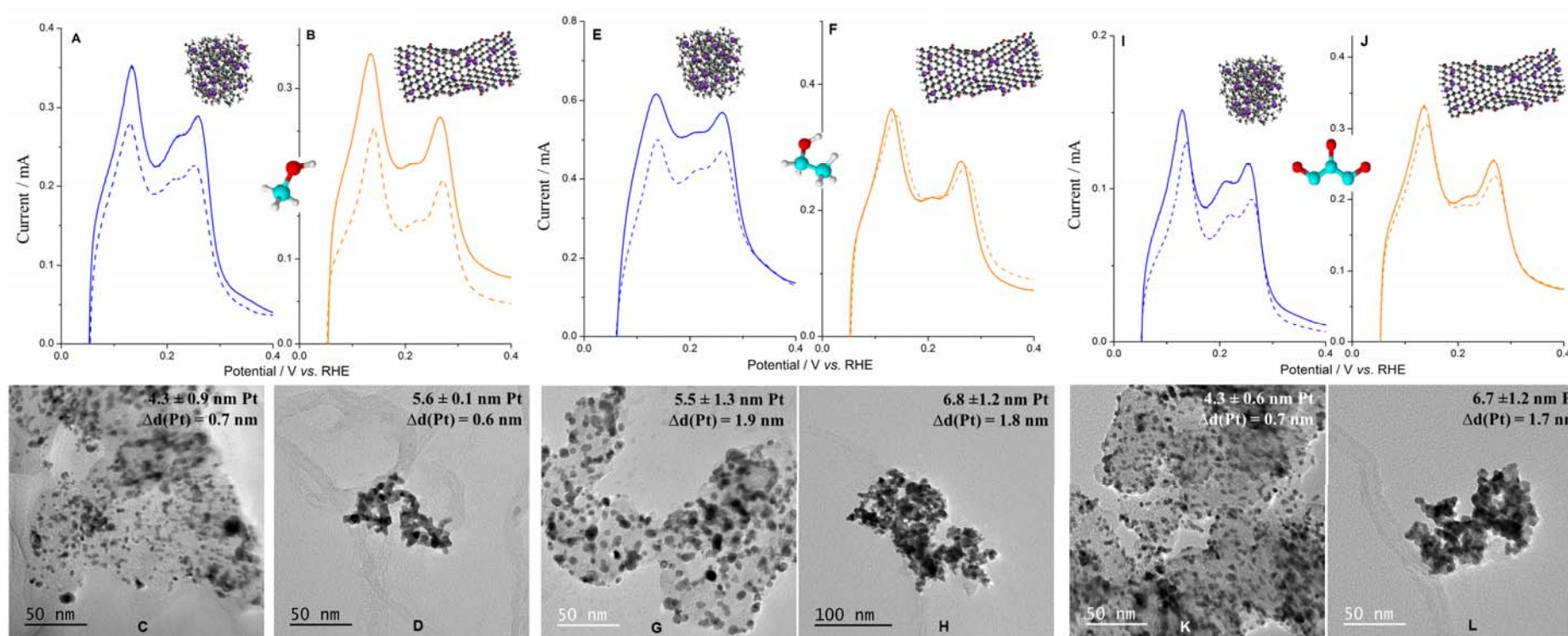
**Figure 4.** Average contact angle of  $0.6 \text{ mol L}^{-1}$  methanol (MeOH),  $0.3 \text{ mol L}^{-1}$  ethanol (EtOH) and  $0.2 \text{ mol L}^{-1}$  glycerol (GIOH) in  $0.5 \text{ mol L}^{-1} \text{ H}_2\text{SO}_4$  on Pt/C and Pt/GNRs.



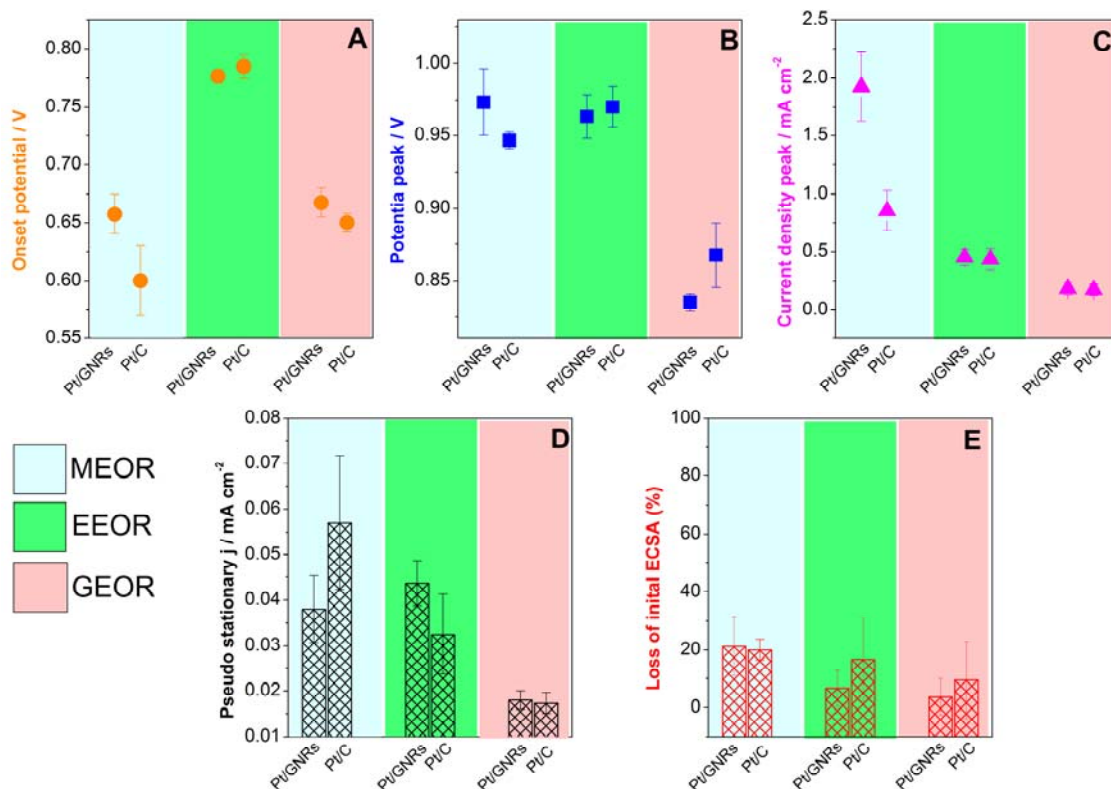
**Figure 5.** Stable cyclic voltammograms of Pt/C and Pt/GNRs in  $0.5 \text{ mol L}^{-1} \text{ H}_2\text{SO}_4$  + (A)  $0.6 \text{ mol L}^{-1}$  methanol, (B)  $0.3 \text{ mol L}^{-1}$  ethanol and (C)  $0.2 \text{ mol L}^{-1}$  glycerol between 0.05 and 1.0 V at  $0.05 \text{ V s}^{-1}$ .



**Figure 6.** Normalized current peak of the anodic current of the positive scan during 1000 potential cycles between 0.3 and 1.0 V at  $0.3 \text{ V s}^{-1}$ . Voltammograms obtained in  $0.5 \text{ mol L}^{-1} \text{ H}_2\text{SO}_4$  + (A)  $0.6 \text{ mol L}^{-1}$  methanol, (B)  $0.3 \text{ mol L}^{-1}$  ethanol and (C)  $0.2 \text{ mol L}^{-1}$  glycerol.



**Figure 7.** Investigation of the electrochemical stability of Pt/C and Pt/GNRs (indicated by lustration) after 1000 potential cycles in 0.5 mol L<sup>-1</sup> H<sub>2</sub>SO<sub>4</sub> + (A-D) 0.6 mol L<sup>-1</sup> methanol, (E-H) 0.3 mol L<sup>-1</sup> ethanol and (I-L) 0.2 mol L<sup>-1</sup> glycerol. The hydrogen desorption regions of voltammeteries recorded in 0.5 mol L<sup>-1</sup> H<sub>2</sub>SO<sub>4</sub> were obtained before (solid lines) and after (dashed lines) the degradation protocols. The micrographs, average sizes and NPs size increment after the tests.



**Figure 8.** The comparison of the electrochemical parameters regarding the methanol (MEOR,  $0.6 \text{ mol L}^{-1}$ ), ethanol (EEOR,  $0.3 \text{ mol L}^{-1}$ ) and glycerol (GEOR,  $0.2 \text{ mol L}^{-1}$ ) electrooxidation reaction on Pt/C and Pt/GNRs nanoparticles. All measurements are shown as an average of three experiments with standard deviation (error bars). (A) Onset potential, (B) potential peak and (C) current density peak obtained from the stable cyclic voltammogram of each alcohol in  $0.5 \text{ mol L}^{-1} \text{ H}_2\text{SO}_4$  between 0.05 and 1.0 V at  $0.05 \text{ Vs}^{-1}$ . (D) Pseudo stationary current density taken from current-time curves after 1800 s of polarization at 0.6 V (start potential of 0.2 V). (E) Average loss of electrochemically active surface area (ECSA) after 1000 potential cycles between 0.3 and 1.0 V at  $0.3 \text{ V s}^{-1}$ .