Electronic inhomogeneities in graphene: the role of the substrate interaction and chemical doping

Inhomogeneidades electrónicas en grafeno: el rol de la interacción con el substrato y el dopaje químico

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Abstract
We probe the local inhomogeneities of the electronic properties of graphene at the nanoscale using scanning probe microscopy techniques. First, we focus on the study of the electronic inhomogeneities caused by the graphene-substrate interaction in graphene samples exfoliated on silicon oxide. We find that charged impurities, present in the graphene-substrate interface, perturb the carrier density significantly and alter the electronic properties of graphene. This finding helps to understand the observed device-to-device variation typically observed in graphene-based electronic devices. Second, we probe the effect of chemical modification in the electronic properties of graphene, grown by chemical vapour deposition on nickel. We find that both the chemisorption of hydrogen and the physisorption of porphyrin molecules strongly depress the conductance at low bias indicating the opening of a bandgap in graphene, paving the way towards the chemical engineering of the electronic properties of graphene.

Resumen
Hemos estudiado las inhomogeneidades locales de las propiedades electrónicas del grafeno a escala nanométrica utilizando técnicas de microscopía de sonda próxima. En primer lugar, nos centramos en el estudio de las inhomogeneidades electrónicas causadas por la interacción del grafeno con el sustrato en muestras de grafeno exfoliado sobre óxido de silicio. Encontramos que las impurezas cargadas, presentes en la interfaz entre el grafeno y el sustrato, perturban considerablemente la densidad de portadores y alteran las propiedades electrónicas del grafeno. Este hallazgo ayuda a comprender la gran variabilidad entre distintos dispositivos que se observa típicamente en dispositivos electrónicos basados en grafeno. En segundo lugar, investigamos el efecto de la modificación química de las propiedades electrónicas de grafeno, crecido sobre níquel por depósito por vapor químico. Encontramos que tanto la quimisorción de hidrógeno como la fisisorción de moléculas de porfirina logran reducir fuertemente la conductancia a bajo voltaje, lo que indica la apertura de un gap en el grafeno, allanando el camino hacia el diseño químico de las propiedades electrónicas de grafeno.

1. Introduction
Since the first experimental realization of graphene [1], its unique properties [2-4] have boosted the research in this novel material. Due to the high sensitivity of graphene to external electric fields, it can be used for sensing applications or as the channel in field effect transistors. This high sensitivity, however, makes graphene very vulnerable to charged impurities in the surroundings and to chemical doping [5, 6], leading to a large device-to-device variation in the electronic performance [6]. This effect is especially critical because graphene is all surface and thus when it lays on a surface or it is covered by a layer of adsorbates almost every single carbon atom of the graphene layer can be altered by the graphene-substrate interaction or the chemical doping.

In this work we review experimental results on the local electronic properties of graphene, focused first on the study of the electronic inhomogeneities caused by the graphene-substrate interaction (based on Ref. [7]) and second on the variation of the local electronic properties of graphene by chemical modification of the surface (based on Ref. [8]).

2. Graphene-substrate interaction: exfoliated graphene on SiO2
When graphene is exfoliated on insulating substrates, the presence of charged impurities typically generates electric fields strong enough to change the doping level of the graphene layer at the nanoscale [9, 10]. As a consequence, graphene-based electronic devices show a large device-to-device variation in electrical performance and reproducibility [5, 6].

We have developed a combined scanning tunnelling microscopy and atomic force microscopy (STM/AFM) to characterize the electronic properties of graphene layers even when they are deposited on top of insulating substrates. This scanning probe microscopy (SPM) tool can operate as an AFM, without the need of a conductive substrate, to locate the graphene flake. The microscope relies on an STM which has been supplemented with a piezoelectric quartz tuning fork force sensor [11, 12] in the so-called qPlus configuration [13] (spring constant k ~ 12500 N/m, resonance frequency f0 ~ 32.1 kHz and quality factor Q ~ 4200). The use of a carbon fiber tip has found to optimize the performance of this type of combined STM/AFM microscopes [14-16].

The measurement starts by positioning the carbon fiber tip on top of a graphene flake (prepared by cleavage of highly oriented pyrolytic graphite on a SiO2/Si substrate using silicone stamps [17]) with the help of a long working distance optical microscope (see Figure 1a). Then the AFM capability of our combined STM/AFM is used to scan the region under study (see Figure 1b) and thus to determine thickness of the different areas of the flake. Before starting the STM measurements the tip is positioned onto the flake, which is electrically contacted by a gold electrode (deposited by shadow mask evaporation), and the scan range is reduced (dashed square in Figure 1b) in order to avoid the tip reaching the insulator substrate which would result in a tip crash.
that degrades the sample and/or tip irreversibly. Figure 1c shows the topography of a mechanically exfoliated graphene flake deposited on a 285 nm SiO$_2$/Si substrate obtained in the constant current STM mode. All STM measurements were acquired under ambient conditions.

In order to gain a deeper insight in the role of charged impurities on the electronic properties of graphene, we have also measured (simultaneously with the topography) the variations of the tunneling barrier height (see the supplementary information of Ref. [7] for technical details) because the tunnelling current decay constant ($\beta$) is strongly influenced by the presence of subsurface charges [18, 19]. In fact, the presence of negatively (positively) charged impurities causes an electric field which effectively shifts upwards (downwards) the energy of the bottom of the band by an amount $\Delta E$. This effect reduces (increases) the apparent tunneling barrier height, $\Phi_{\text{app}}$. This barrier height change can be modeled as $\Phi_{\text{app}} = (\Phi_{\text{graphene}} - \Phi_{\text{tip}} + \Delta E)$ where $\Phi_{\text{graphene}}$ and $\Phi_{\text{tip}}$ are the graphene and the tip work function respectively. The associated tunneling decay constant then changes according to $\beta = 2\sqrt{2m \cdot \Phi_{\text{app}}} / h$ ($m$ is the electron mass and $h$ is the reduced Planck constant).

Figure 2a shows the spatial variation of $\beta$ for a multilayer graphene simultaneously measured with the STM topography (Figure 1d). The $\beta$ image for a multilayer shows an almost constant value ($5.3 \pm 0.5$ nm$^{-1}$), typical of STM operation in air [20]. The spatial variation of $\beta$ for single-layer graphene, on the other hand, shows strongly localized inhomogeneities (identified as dips in the $\beta$ image) caused by local doping induced by negative charged impurities in the substrate (Figure 2b). These inhomogeneities in the tunneling decay constant can be also observed by measuring tunnelling vs. distance traces at different positions in the single layer graphene region (Figure 2d).

Considering that each dip in the $\beta$ image (Figure 2b) is due to the presence of one individual negative charge we can estimate the density of charged impurities by tentatively counting the number of depressions in a given area, resulting in an impurity density $\sigma = (2.9 \pm 0.6) \times 10^{11}$ cm$^{-2}$. A more objective procedure to statistically analyse the short-range ordering [21] and spatial variation of $\beta$ is to obtain the radially-averaged autocorrelation function $g(r)$, shown in Figure 5c. The typical radius $r_{FWHM}$ at half minimum of the localized inhomogeneities is obtained from the radial distance at which the value of $g(r)$ is
0.5, which yields $F_{WHM} = 4.7 \pm 1.1 \text{ nm}$. The inhomogeneities do not show long range ordering and the mean spacing between them is $d = 22 \pm 2 \text{ nm}$, determined from the position of the first maximum of $grf$[21], which corresponds to a charge density $\sigma = \frac{\pi}{4}(d^2/2) = (2.6 \pm 0.5) \times 10^{11} \text{ cm}^{-2}$. As the radius of the $\beta$ dips presents a well-defined average value, it indicates that the charged inhomogeneities are localized at a well-defined distance from the graphene layer. If the charged impurities would be located randomly in the SiO$_2$ layer (300 nm thick) the distribution of radius would be much broader. Therefore, it is reasonable to assume that the charged impurities are located at the interface between the SiO$_2$ layer and the graphene flake.

3. Effect of chemical doping on the local electronic properties of CVD graphene

Chemical functionalization of graphene is a very promising route to achieve large scale production of semiconducting graphene for industrial applications. Indeed, the application of graphene in microelectronic devices is hampered by the lack of a bandgap which is mandatory in order to fabricate electrically switchable devices with large current on/off ratio. Up to date several strategies have been developed to open a band gap in graphene. Here we have explored two different options. First, the exposure of the graphene surface to hydrogen plasma [22-24] that leads to the chemisorption of hydrogen atoms, which produces the sp$^2$ hybridization of carbon network, reduces the number of delocalized sp$^2$ electrons and consequently opens a band gap. Second, the non-covalent stacking of aromatic organic molecules on graphene through $\pi-\pi$ interaction.

3.1. Hydrogen chemisorption

The hydrogenation of CVD graphene has been done using Ar:H$_2$ plasma (composition of 85:15) in a reactive ion etching system with a high frequency generator operating at 13.56 MHz, capacitively coupled to the bottom electrode [25]. The gas flow is kept constant at 200 sccm and the pressure in the chamber is 0.05 mbar. We chose the lowest plasma ignition power, $P = 3 \text{ W}$ (power density is ~4 mW/cm$^2$), and tuned the circuit impedance to reduce the built-in DC self-bias between the bottom electrode and the plasma down to zero. Each exposure is done for 40 min, which leads to moderate hydrogen coverage and is directly followed by STM/STS measurements.

Figure 3 shows an 9 x 9 nm$^2$ topographic image acquired in the constant current STM mode of the surface of graphene grown on nickel by CVD.[26] Before the plasma treatment, the CVD graphene exhibits a Moiré pattern superimposed to the honeycomb lattice of graphene (Figure 3a). This is due to the lattice parameter mismatch between the graphene and the nickel surfaces and it is the characteristic of most of epitaxial graphene samples [27-29]. On the other hand, for the hydrogenated CVD graphene, the expected structural changes are twofold. First, the chemisorption of hydrogen atoms will change the sp$^2$ hybridization of carbon atoms to tetragonal sp$^3$ hybridization, modifying the surface geometry.[30, 31] Second, the impacts of heavy Ar ions, present in the plasma, could also modify the surface by inducing geometrical displacement of carbon atoms (rippling graphene surface) or creating vacancies and other defects. Figure 3(b) shows the topography image of the surfaces CVD graphene after the extended plasma treatment. The corrugation increases after the treatment and there are brighter regions in where the atomic resolution is lost or strongly distorted.

We have also found that these bright regions present a semiconducting behaviour while the rest of the surface remains conducting (see Figure 3(c)-(d)). Although the room temperature thermal drift makes it challenging to spatially resolve the electronic properties of the samples by STS with atomic accuracy, it is possible to analyze statistically thousands of tunneling spectroscopy measurements to infer changes on the electronic properties of the graphene surface (see Ref. [8] for more details).

3.2. Porphyrin physisorption

Iron porphyrins (FePP) are deposited onto the CVD graphene surface by drop casting. The porphyrins were used without any further purification from Aldrich. A solution 1.63 mM of FePP in uvasol grade chloroform was prepared. Further, a droplet of 10 µL of the porphyrin solution was dropcasted onto CVD-graphene surface and the solvent was left to evaporate for 3h in open air condition.

Figure 4a shows the STM topography of the CVD graphene surface after the deposition of FePP molecules. We observe six-pointed star shape protuberances with 0.26 nm in height and approximate dimension of 13 Å x 19 Å. These lateral
dimensions are in agreement with the dimensions of a single porphyrin molecule obtained by first principle calculations (16 Å x 16 Å) (Hyperchem 6.0). The resolved six-star shape, however, does not perfectly match the expected geometrical structure of porphyrin molecules (four-star shape). The additional features in the observed star shape can be due to the splitting of the local density of the state (LDOS) of one phenyl ring. Another interesting finding is that FePP molecules are preferable physisorbed at the valleys of the rippled structure of graphene (Fig. 4a).

After the deposition of FePP onto the graphene surface, we measured the $I$-$V$ traces directly on top of six porphyrin molecules (as the one indicated by the red cross in Figure 4a). The tunneling differential conductance shows a strong reduction in comparison to pristine graphene. In fact, the tunneling differential conductance traces present zero conductance (below the experimental resolution, 10 pS) for a wide range in bias, a characteristic of semiconducting behaviour while for pristine graphene the low bias differential conductance value is significantly larger than zero. Surprisingly, when this measurement is repeated on graphene several angstroms away from a single porphyrin molecule (at the topmost point of Moiré as indicated by the black cross in Fig 4a) the characteristic $I$-$V$ traces (Fig. 4c) are different than that of pristine graphene (Fig. 4b). The tunnelling differential conductance at low bias is still strongly reduced. This indicates that the non-covalent modification of the graphene surface can be used to engineer its electronic properties.

Conclusions
We have studied the local electronic properties of graphene using scanning probe microscopy techniques. We found that when graphene is deposited on a SiO₂ surface, the charged impurities present in the graphene-substrate interface produce strong inhomogeneities of the electronic properties of graphene. On the other hand, we show how homogeneous graphene grown by CVD can be altered by chemical modification of its surface. We found that both the chemisorption of hydrogen and the physisorption of porphin molecules strongly depress the local conductance at low bias indicating the opening of a bandgap in graphene.

Figure 4. (a) 5,10,15,20-Tetraphenyl-21H,23H-porphine iron(III) chloride (abbreviated as FePP), molecules decorating the terraces of CVD graphene ($I = 0.1$ nA; $V = 0.5$ V). FePP can be found at the valleys of the Moiré pattern of graphene. (b)-(d) Comparison of the differential conductance measured on the graphene sample after FePP deposition. The data is presented as 2D histograms, each one built from 1000 individual STS traces. (b) STS spectra measured on top of the Moiré pattern far from the molecules. (c) $dI/dV$ traces acquired on top of the porphyrin molecule marked with a red cross in Fig. 4a. (d) STS measured on a hill of the Moiré close to a porphyrin molecule (black cross in Fig. 4a).

Figura 4. (a) 5,10,15,20-tetrafenil-21H,23H-porphirina de cloruro de hierro (III) (abreviado como FePP), decorando las terrazas de grafeno CVD ($I = 0.1$ nA; $V = 0.5$ V). Las FePP se pueden encontrar en los valles del patrón de Moiré del grafeno. (b) - (d) Comparación de la conductancia diferencial medida en la muestra después de la deposición de FePP en grafeno. Los datos se presentan como histogramas 2D, cada uno construido a partir de 1000 trazas STS individuales. (b) espectros STS medida en la parte superior del patrón de Moiré lejos de las moléculas. (c) Curvas $dI/dV$ adquiridas en la parte superior de la molécula de porfirina marcada con una cruz roja en la Figura 4a. (d) STS medido en la parte superior del patrón de Moiré cercano a una molécula de porfirina (cruz de color negro en la figura. 4a).

Acknowledgements
A.C.-G. acknowledges fellowship support from the Comunidad de Madrid (Spain) and the Universidad Autónoma de Madrid (Spain). M.W., and N.T. acknowledge financial support from the Ubo Emmius program of the Groningen Graduate School of Science, the Zernike Institute for Advanced Materials and the Netherlands Organization for Scientific Research (NWO-CW) through a VENI grant. A. thanked financial support from the Foundation for Fundamental Research on Matter (FOM G-08). This work was supported by MINECO (Spain) through the programs MAT2008-01735, MAT2011-25046 and CONSOLIDER-INGENIO-2010 1‘Nanociencia Molecular’ GSD-2007-00010, Comunidad de Madrid through program Nanobiomagnet S2009/MAT-1726.

References
1Novoselov, K. S.; Geim, A. K.; Morozov, S. V.; Jiang, D.; Zhang, Y.; Dubonos, S. V.; Grigorieva, I. V.; Firsov, A. A. Electric field effect in atomically thin carbon films Science 2004; 306(5696):666-669.
2Novoselov, K. S.; Jiang, Z.; Zhang, Y.; Morozov, S. V.; Stormer, H. L.; Zeitler, U.; Maan, J. C.; Boebinger, G. S.; Kim, P.; Geim, A. K. Room-temperature quantum Hall effect in graphene Science 2007; 315(5817):1379.
3Ozyilmaz, B.; Jarillo-Herrero, P.; Efetov, D.; Abanin, D. A.; Levitov, L. S.; Kim, P. Electronic transport and quantum hall effect in bipolar graphene p-n-p junctions Phys. Rev. Lett. 2007; 99(16):166804-4.
4Zhang, Y.; Tan, Y. W.; Stormer, H. L.; Kim, P. Experimental observation of the quantum Hall effect and Berry’s phase in graphene Nature 2005; 438(7065):201-204.
5Chen, J. H.; Jang, C.; Adam, S.; Fuhrer, M. S.; Williams, E. D.; Ishigami, M. Charged-impurity scattering in graphene Nature Phys. 2008; 4(5):377-381.
6Tan, Y. W.; Zhang, Y.; Bolotin, K.; Zhao, Y.; Adam,
S.; Hwang, E. H.; Das Sarma, S.; Stormer, H. L.; Kim, P. Measurement of Scattering Rate and Minimum Conductivity in Graphene Phys. Rev. Lett. 2007; 99(24):246803-4.

7 Castellanos-Gomez, A.; Smit, R. H. M.; Agrait, N.; Rubio-Bollinger, G. Spatially resolved electronic inhomogeneities of graphene due to subsurface charges Carbon 2012; 50(3):932-938.

8 Castellanos-Gomez, A.; Wojtaszek, M.; Arramel; Tombros, N.; van Wees, B. J. Reversible Hydrogenation and Bandgap Opening of Graphene and Graphite Surfaces Probed by Scanning Tunneling Spectroscopy Small 2012; 8(10):1607-1613.

9 Adam, S.; Hwang, E. H.; Rossi, E.; Das Sarma, S. Theory of charged impurity scattering in two-dimensional graphene Solid State Commun. 2009; 149(27-28):1072-1079.

10 Zhang, Y.; Brar, V. W.; Girit, C.; Zettl, A.; Crommie, M. F. Origin of spatial charge inhomogeneity in graphene Nature Phys. 2009; 5(10):722-726.

11 Castellanos-Gomez, A.; Agrait, N.; Rubio-Bollinger, G. Dynamics of quartz tuning fork force sensors used in scanning probe microscopy Nanotechnology 2009; 20(21):215502-8.

12 Castellanos-Gomez, A.; Agrait, N.; Rubio-Bollinger, G. Force-gradient-induced mechanical dissipation of quartz tuning fork force sensors used in atomic force microscopy Ultramicroscopy 2011; 111(3):186-190.

13 Giessibl, F. J. High-speed force sensor for force microscopy and profilometry utilizing a quartz tuning fork Appl. Phys. Lett. 1998; 73(26):3956-3958.

14 Castellanos-Gomez, A.; Agrait, N.; Rubio-Bollinger, G. Carbon fibre tips for scanning probe microscopy based on quartz tuning fork force sensors Nanotechnology 2010; 21(4):145702.

15 Castellanos-Gomez, A.; Bilan, S.; Zotti, L. A.; Arroyo, C. R.; Agrait, N.; Cuevas, J. C.; Rubio-Bollinger, G. Carbon tips as electrodes for single-molecule junctions Appl. Phys. Lett. 2011; 99(12):123105-3.

16 Rubio-Bollinger, G.; Castellanos-Gomez, A.; Bilan, S.; Zotti, L.; Arroyo, C.; Agrait, N.; Cuevas, J. C. Carbon fibre tips for scanning probe microscopy and molecular electronics experiments Nanoscope Research Letters 2012; 7(1):254.

17 Moreno-Moreno, M.; Castellanos-Gomez, A.; Rubio-Bollinger, G.; Gomez-Herrero, J.; Agrait, N. Ultralong natural graphene nanoribbons and their electrical conductivity Small 2009; 5(8):924-927.

18 Kobayashi, K.; Kurokawa, S.; Sakai, A. Scanning tunneling microscopy and barrier-height imaging of subsurface dopants on GaAs (110) Jpn. J. Appl. Phys. 2005; 44(12):8619.

19 Kurokawa, S.; Takei, T.; Sakai, A. A search for subsurface dopants on hydrogen-terminated Si (111) surfaces Jpn. J. Appl. Phys. 2003; 42(4):4655.

20 Meepagala, S.; Real, F. Detailed experimental investigation of the barrier-height lowering and the tip-sample force gradient during STM operation in air Phys. Rev. B 1994; 49(15):10761-10763.

21 Sánchez, F.; Infante, I. C.; Lüders, U.; Abad, L.; Fontcuberta, J. Surface roughening by anisotropic adatom kinetics in epitaxial growth of La0.67Ca0.33MnO3 Surf. Sci. 2006; 600(6):1231-1239.

22 Elias, D.; Nair, R.; Mohiuddin, T.; Morozov, S.; Blake, P.; Halsall, M.; Ferrari, A.; Boukhvalov, D.; Katsnelson, M.; Geim, A. Control of graphene’s properties by reversible hydrogenation: evidence for graphene Science 2009; 323(5914):610.

23 Luo, Z.; Shang, J.; Lim, S.; Li, D.; Xiong, Q.; Shen, Z.; Lin, J.; Yu, T. Modulating the electronic structures of graphene by controllable hydrogenation Appl. Phys. Lett. 2010; 97(23):233111-233111-3.

24 Burgess, J. S.; Matis, B. R.; Robinson, J. T.; Bulat, F. A.; Keith Perkins, F.; Houston, B. H.; Baldwin, J. W. Tuning the electronic properties of graphene by hydrogenation in a plasma enhanced chemical vapor deposition reactor Carbon 2011.

25 Wojtaszek, M.; Tombros, N.; Careta, A.; van Loosdrecht, P. H. M.; van Wees, B. J. A road to hydrogenating graphene by a reactive ion etching plasma Arxiv preprint arXiv:1109.1684 2011.

26 www.graphenesupermarket.com.

27 Berger, C.; Song, Z.; Li, T.; Li, X.; Ogbazghi, A.; Feng, R.; Dai, Z.; Marchenkov, A.; Conrad, E.; De Heer, W. Ultrathin epitaxial graphite: 2D electron gas properties and a route toward graphene-based nanoelectronics J. Phys. Chem. B 2004; 108(52):19912-19916.

28 Sutter, P.; Flege, J.; Sutter, E. Epitaxial graphene on ruthenium Nature Mater. 2008; 7(5):406-411.

29 Vázquez de Parga, A.; Calleja, F.; Borca, B.; Passeggi Jr, M.; Hinarejos, J.; Guinea, F.; Miranda, R. Periodically rippled graphene: Growth and spatially resolved electronic structure Phys. Rev. Lett. 2008; 100(5):56807.

30 Ruffieux, P.; Gröning, O.; Bielmann, M.; Mauron, P.; Schlaphbach, L.; Gröning, P. Hydrogen adsorption on sp2-bonded carbon: Influence of the local curvature Phys. Rev. B 2002; 66(24):245416.

31 Muniz, A. R.; Maroudas, D. Hydrogenation effects on the structure and morphology of graphene and single-walled carbon nanotubes J. Appl. Phys. 2010; 108113532.