Magnetoresistance in disordered graphene: The role of pseudospin and dimensionality effects unraveled

FRANK ORTMANN\(^1\), ALESSANDRO CRESTI\(^2\), GILLES MONTAMBAUX\(^3\) and STEPHAN ROCHE\(^4,5\)(a)

\(^1\)CEA, INAC, SPRAM, GT - 17 rue des Martyrs, 38054 Grenoble Cedex 9, France, EU
\(^2\)IMEP-LAHC, UMR 5130 (Grenoble INP/UJF/CNRS/Université de Savoie), Minatec - 3 Parvis Louis Néel, 38016 Grenoble, France, EU
\(^3\)Laboratoire de Physique des Solides, UMR 8502 du CNRS, Université Paris-Sud - 91405 Orsay, France, EU
\(^4\)CIN2 (ICN-CSIC) and Universitat Autònoma de Barcelona, Catalan Institute of Nanotechnology, Campus de la UAB - 08193 Bellaterra (Barcelona), Spain, EU
\(^5\)ICREA, Institut Català de Recerca i Estudis Avançats - 08010 Barcelona, Spain, EU

received 15 December 2010; accepted in final form 8 April 2011
published online 6 May 2011

PACS 72.80.Vp – Electronic transport in graphene
PACS 72.15.Rn – Localization effects (Anderson or weak localization)
PACS 73.63.-b – Electronic transport in nanoscale materials and structures

Abstract – We report a theoretical low-field magnetotransport study unveiling the effect of pseudospin in realistic models of weakly disordered graphene-based materials. Using an efficient Kubo simulation method, and simulating the effect of charges trapped in the oxide, different magnetoconductance fingerprints are numerically obtained for system sizes as large as 0.3 \(\mu\)m\(^2\), containing tens of millions of carbon atoms. In two-dimensional graphene, a strong valley mixing is found to irreparably yield a positive magnetoconductance (weak localization), whereas crossovers from positive to negative magnetoconductance (weak antilocalization) are obtained by reducing the disorder strength down to the ballistic limit. In sharp contrast, graphene nanoribbons with lateral size as large as 10 nm show no sign of weak antilocalization, even for very small disorder strength. Our results rationalize the emergence of a complex phase diagram of magnetoconductance fingerprints, shedding new light on the microscopical origin of pseudospin effects.

In 2004, the report on graphene discovery [1] has sparked a great scientific excitement because of both novel type of electronic excitations (so-called massless Dirac fermions) [2] and a promising future of graphene-based technologies [3]. In two-dimensional graphene, the very peculiar nature of low-energy electronic states (encompassing a new pseudospin degree of freedom) yields a wealth of anomalous transport features such as Klein tunneling [4], weak antilocalization [5,6], unconventional quantum Hall effect [7,8], or new ways (supercollimation) to guide charge flows [9].

Besides, the nature of disorder, its effect on electronic properties, and the envisioned defect engineering for novel graphene electronics are currently subjects of great concern [10,11]. However to date, the precise relationship between the underlying disorder features and the onset of graphene unique transport properties remains debated.

The question also applies to graphene nanoribbons which exhibit different electronic properties induced by lateral size confinement effects and edge symmetries [12–16].

One intriguing question is the precise role of pseudospin effects in the modulations of the conductance fingerprints in disordered graphene-based materials (both two-dimensional and nanoribbons). Indeed, in ordinary low-dimensional disordered metals, quantum interferences between multiple scattering paths produce a quantum correction to the semiclassical (Drude) conductivity which can be either reduced (weak localization) [17,18] or enhanced (weak antilocalization-WAL) [19,20], depending on the strength of spin-orbit coupling. Suzuura and Ando [21] first suggested that in graphene, even in the absence of spin-orbit coupling, WAL could occur depending on the long-range nature of the underlying disorder potential. A complete perturbation (diagrammatic) theory was further developed by McCann and coworkers [5], to compute the quantum correction of the (semi-classical) Drude result.

\(^{(a)}\)E-mail: stephan.roche@icn.cat
To determine such a correction, in addition to the usual coherence and magnetic lengths, several phenomenological parameters (intravalley and intervalley elastic scattering times) were introduced to fix the scales beyond which certain classes of interfering loops start to contribute. Unfortunately, the precise values of those elastic scattering times remain out of reach from analytical considerations when considering a realistic disorder potential profile (see also ref. [6] for discussion). After several experimental indications of WAL in both epitaxial [22] and exfoliated graphene layers [23], a complex phase diagram has been reported, exhibiting multiple crossovers from weak antilocalization (negative magnetoconductance) to weak localization (positive magnetoconductance) when tuning the energy of charge carriers or the sample temperature [24].

However, the understanding of the microscopic origin of such a complex phase diagram and its relation to a possible metal-insulator transition remains obscure. In that perspective, it is crucial to clarify the connection between the precise disorder characteristics and the magnetic-field-dependent quantum interference effects (QIE) driving the conductance modulations, as well as the true contribution of the pseudospin degree of freedom, recently questioned by Winkler and Zülicke [25]. Moreover, a recent study pointing towards a deep connection between the pseudospin and the real angular momentum [26] also demands for a careful inspection of pseudospin effects in a complex transport situation. Finally, whereas weak localization in 2D is precursor of a true insulating state in the zero-temperature limit, WAL points towards a robust metallic state, a fact in marked contrast with the usual scaling theory of localization for two-dimensional disordered systems [18,27,28].

Here, by means of an efficient quantum-transport simulation method, the weak-field magnetoconductance of disordered graphene-based materials is explored for the case of long-range disorder potential, which mimics the effect of charges trapped in the underneath oxide layers. This study unveils the true contribution of pseudospin effects in realistic disordered graphene. In two-dimensional graphene, a transition from positive to negative magnetoconductance is confirmed to be related to the decay of the intervalley scattering contribution. Our study provides a clarification of the microscopic origin of the reported experimental phase diagram [24] in terms of the dominant disorder characteristics, whereas weak antilocalization and pseudospin effects are found to be suppressed in the presence of graphene edges (graphene nanoribbons).

We use the π electrons model of graphene well described by the tight-binding Hamiltonian $\mathcal{H} = \sum_\alpha V_\alpha |\alpha\rangle \langle \alpha| + \gamma_0 \sum_{\alpha \beta} e^{-i \phi_{\alpha \beta}} |\alpha\rangle \langle \beta|$, with $\gamma_0 = -2.7$ eV the hopping integral between the nearest-neighbor carbon atoms with distance $a = 0.142$ nm, whereas $V_\alpha$ are the onsite energies of π-orbitals and include the (long-range) disorder potential profile. The effect of the magnetic field is introduced through the Peierls phase [29], with a flux per hexagon being equal to $\phi = J \mathbf{A} \cdot d\mathbf{l} = h/e \sum_{\text{hexagon}} \phi_{\alpha \beta}$. We implement a convenient gauge where $\sum_{\text{hexagon}} \phi_{\alpha \beta}$ can take integer multiples of $1/(N_x N_y)$ to vary the magnetic-field strength and where $N_x$ and $N_y = N_x + 1$ give the sample size. The total density of states in clean graphene at zero field and at $B = 10$ T is found to exhibit the well-known $\sqrt{nB}$ Landau level structure in perfect agreement with theory [30]. We use a common model to describe the long-range disorder [31,32], which is given by the contributions from $N_I$ impurities randomly distributed at $r_i$ (among $N$ sites of the system) with $V_\alpha = \sum_{i=1}^{N_I} \varepsilon_i \exp\left(-|r_\alpha - r_i|^2/(2\xi^2)\right)$, where $\xi$ defines the effective range, and $\varepsilon_i$ are chosen at random within $[-W/2, W/2]$ ($\gamma_0$-unit), such that $W$ gives the strength of the local potential profile (for an illustration see fig. 1 (c)). Different random configurations of graphene samples with same size, $\xi$, $W$, and $n_i = N_i/N$ constitute an ensemble with given disorder strength. Following prior studies [31,32], we fix $\xi = 3a = 0.426$ nm as a typical value for a long-range potential, but vary $W$ to describe different screening situations.

The (magneto)-transport properties of large and disordered graphene systems are simulated by using an efficient order-N Kubo method [33–35]. From the study of the quantum wave packet dynamics, the elastic mean free path $\ell_r(E)$ can be first derived from the time dependence of the diffusion coefficient $D(E,t)$

$$D(E,t) = \frac{1}{t} \frac{\sum_n \langle \Psi_n | \delta(E - H) \hat{U}^\dagger(t) \hat{x} \hat{U}(t) - \hat{x} \rangle^2 |\Psi_n\rangle}{\sum_m \langle \Psi_m | \delta(E - H) |\Psi_m\rangle}$$

Fig. 1: (Colour on-line) (a) Time-dependent diffusion coefficient at the Dirac point for several impurity densities and $W = 2$. (b) Density of states of pristine graphene at zero and at 10 T. (c) Schematic of the disordered potential in graphene due to long-range Coulomb impurities.
The maximum value of $D(t) = D_{\text{max}}$ allows for the evaluation of the elastic mean free path $\ell_e = D_{\text{max}}/(2v_F)$ (cf. fig. 2) where $v_F$ is the Fermi velocity ($v_F = 8.7 \times 10^5 \text{ m/s}$ at the Dirac point). It is seen that $\ell_e \sim 1/n_i$, in agreement with Fermi’s Golden rule. The dependence of the elastic scattering time $\tau_e = \ell_e/v_F$ on the charge density $n$ follows $\tau_e \sim 1/\sqrt{n}$ for sufficiently small $n$ and $W$ (not shown here) and shows a dip at zero energy for larger $W$ (see fig. 2, inset).

The effect of an applied magnetic field is shown in fig. 3. Top and bottom panels give the time-dependent diffusion coefficients at varying magnetic field and the magnetoconductivity $\Delta \sigma(B) = \sigma(B) - \sigma(B = 0)$, respectively. For $W = 2$ (fig. 3(a)), a steady suppression of QIE-related conductance corrections is clearly seen with increasing $B$ up to $B = 0.274 \text{T}$ at which the decay of $D(t)$ has been fully suppressed. This is in accordance with the weak-localization phenomenon, as confirmed by the positive magnetoconductance reported in panel (d) for four selected energies (averages over 32 different configurations are performed). In contrast, the case $W = 1.5$ shows a clearly different behavior as seen in figs. 3(b) and (e), respectively for $D(t)$ and $\Delta \sigma(B)$ (for which 64 configurations have been averaged). Indeed, a change in the sign of $\Delta \sigma(B)$ is observed, in agreement with a crossover from weak localization to weak antilocalization, which depends not only on the field strength, but also the considered energy of charge carriers. We note that an estimation of the elastic mean free path gives $\ell_e \sim 9–20 \text{ nm}$, which is much smaller than our sample size.

For the case $W = 1$, we first observe that at zero magnetic field, $D(t)$ does not reach a saturation regime within our total computational time (dashed black curve in fig. 3(c)). With increasing the magnetic field, $D(t)$ is found to be reduced and no onset of weak localization manifests. The corresponding $\Delta \sigma(B)$ curves displayed in panel (f) all exhibit a negative magnetoconductance, whose connection to a WAL regime, as discussed here below, is however not straightforward.

To rationalize all those results, we first note that the numerical study reported in [32] suggests that as long as $W \geq 1$, the long-range potential (with $\xi = 0.426 \text{ nm}$) introduces strong valley mixing. This is consistent with our computed positive magnetoconductance for the case $W = 2$ (fig. 3(d)). By further decreasing the disorder strength (from $W = 2$ to $W = 1.5$), WAL occurs as seen in fig. 3(e) which corresponds to the decay of intervalley scattering contribution (see also fig. 4 of [32]). Our study thus exhibits a transition from weak localization (for $W = 2$) to WAL ($W = 1.5$), supporting that the enhancement of intervalley scattering contribution yields predominance of the weak localization effect. The analysis can be deepened by fitting the simulations using the phenomenological law $\Delta \sigma(B) = e^2/\pi h \{ \mathcal{F} (\tau_B^{-1}/\tau_\varphi^{-1}) - 3 \mathcal{F} (\tau_B^{-1}/(\tau_\varphi^{-1} + 2\tau_e^{-1})) \}$,
Fig. 3: (Colour on-line) Top panels: diffusion coefficient at the Dirac point for various magnetic fields and $W$, $n_i = 0.125\%$, $\xi = 0.426\$nm. Bottom panels: $\Delta \sigma(B)$ for four different Fermi level positions ($E_{\text{DP}} = 0$, $E_{\text{I}} = 0.049\$eV$, $E_{\text{II}} = 0.097\$eV$ and $E_{\text{III}} = 0.146\$eV$). Dashed lines are fits as explained in the text. For $W = 1.0$ in (f) data and fit at $E_{\text{DP}}$ have been rescaled by 0.3 for clarity.

where $\mathcal{F}(z) = \ln z + \psi(1/2 + z^{-1})$, $\psi(x)$ is the digamma function and $\tau_B^{-1} = 4eDB/\hbar$ [5]. Here, for simplicity we restrict the model to a single additional elastic scattering time $\tau_\text{e}$ (phenomenological parameter) that contains both contributions of intravalley and intervalley scattering [5]. The least-squares fits (dashed lines) are superimposed to the simulated $\Delta \sigma(B)$ (symbols), taking $\tau_\text{e} = 9\$ps (our maximum computed time). For $W = 2$ and $W = 1.5$, $\tau_\text{e}$ ranges within [1.1–2.3]ps and [1.5–6.3]ps, respectively (increasing values with increasing energy), thus confirming the weak localization regime for lowest $B$, which is fully consistent with [5] ($\tau_\text{e} < \tau_\text{r}$). The case $W = 1$ is more complicated, with a fit (with $\tau_\text{e} > \tau_\text{r}$) only possible at lower magnetic fields and higher energies ($E_{\text{II}}$ and $E_{\text{III}}$) and out of reach for energies nearby the Dirac point. Despite the obtained negative magnetoconductivity, the interpretation in terms of weak antilocalization is actually not obvious. Indeed, as seen in the diffusion coefficient behavior at zero magnetic field, the saturation regime is not reached within our maximum computed time, although the conduction clearly departs from a ballistic regime. By using $\ell_c = D_{\text{max}}/2v_F$, with an extrapolated maximum diffusion coefficient of $\geq 2500\$cm$^2$s$^{-1}$, one gets $\ell_c \geq 330\$nm, which is in the order of our graphene longitudinal size. Accordingly, the transport regime is closer to a quasiballistic case, which jeopardizes the interpretation in terms of WAL. A careful inspection of the experimental data of ref. [24] suggests that as the Fermi level is moved from higher energies down to the Dirac point, the tendency towards WAL is enhanced for a given temperature and magnetic-field strength (note however that intriguingly there is no data at the Dirac point). Surprisingly, this tendency for the energy dependence is actually found opposite in our simulations (see fig. 3(e)). One could first assume that this discrepancy originates from screening effects which are not presently accounted for in a self-consistent manner. On the other hand, other experiments on weakly damaged graphene (with epoxy [36] or fluorine defects [37]) clearly evidence stronger localization contributions close to the Dirac point, as precursor of a transition to the Anderson regime which is more in line with our findings. We also mention that in our simulations, the coherence time is energy-dependent, an assumption that could be relaxed benefiting from the experimental estimations [24].
Different set of curves are identified depending on the ratio $L/\ell$ for a 10 nm wide armchair ribbon (84-aGNR) with $W = 1/2$, $n_s = 0.1\%$ and for different lengths (up to 1.5 $\mu$m) of the disordered region. Main frame: energy-dependent mean free path for $n_s = 0.1\%$ (black curve) and $n_s = 0.2\%$ (red curve). (b) Schematic of a short segment of a disordered nanoribbon with long-range scatters. Bright (dark) spots correspond to positive (negative) values of $V$.

To analyze the contribution of sample geometries (edges), we finally simulate numerical transport in graphene nanoribbons [12,15]. As a typical case, we focus on a 10 nm wide armchair ribbon (84-aGNR) with impurity densities $n_s = 0.1\%, 0.2\%$, $\xi = 0.426$ nm and $W = 1/2$. The results for $T(E)$ are averaged over 100 disorder configurations (fig. 4, inset). At the low fields (up to 0.5 T) considered here, the bandwidth structure does not change significantly with respect to the zero-field case. Accordingly, the impact of the field on transport is only determined by the modifications of QIE. Figure 4 (main frame) shows the elastic mean free path of disordered ribbons as a function of electron energy. As in the case of Anderson-type disorder or edge defects [15], $\ell_e$ exhibits important modulations depending on the electron energy, with systematic decay around the energies corresponding to the onset of new subbands. At low energies, $\ell_e$ can even reach a few micrometers for the considered disorder values.

Figure 5 shows the evolution of the quantity $\Delta G(B) = G(B) - G(B = 0)$ (for $n_s = 0.1\%$) for the same set of ribbon lengths considered in fig. 4, and at two different selected energies (dashed vertical lines of fig. 4). Note that the small value of $W = 0.5$ considered here would prohibit valley mixing in the two-dimensional case. The ratio $L/\ell_e$ gives a suitable criterion to classify the transport regime as quasi-ballistic (when $L/\ell_e \leq 1$) and diffusive or localized (when $L/\ell_e > 1$). When $L/\ell_e > 1$, a markedly positive magnetoconductance dominates ($\Delta G(B) > 0$) in agreement with the standard weak localization regime (fig. 5). The magnetoconductance variation increases for larger lengths, giving a stronger field-driven suppression of QIE. The results in the case $n_s = 0.2\%$ are analogous. Our numerical results evidence the absence of weak antilocalization for disordered GNRs because of the strong reintroduction of intervalley scattering in the presence of ribbon edges. One notes that the application of much larger magnetic fields has been recently theoretically predicted and experimentally found to yield very large magnetoresistance signals [38,39]. In the case of zigzag nanoribbons, the two valley structure of two-dimensional graphene is preserved, thus making it reasonable to expect a weak antilocalization behavior. However, our results (not reported here) do not show any clear trend. This might be related to several interrelating factors, such as the existence of a perfectly conducting channel [40] or the raise of intervalley scattering induced by defects close to the edges.

In conclusion, in two-dimensional disordered graphene the crossover between weak localization and weak antilocalization was theoretically demonstrated to be driven by the nature of long-range Coulomb scattering potential. In contrast, predominance of weak localization effects was obtained in graphene nanoribbons. These results allow one to bridge the nature of disorder in graphene with the localization phase diagram explored experimentally [24] and also open new directions for understanding pseudospin effects in graphene-based materials.

***

This work is supported by the NANOSIM-GRAPHENE project (ANR-09-NANO-016-01) of ANR/P3N2009 and a Marie Curie Intra European Fellowship within the 7th European Community Framework Programme. AC acknowledges support from Fondation Nanosciences via the RTRA Dispograph project.

REFERENCES

[1] GEIM A. K. and NOVOSELOV K. S., Nat. Mater., 6 (2007) 183.
[2] CASTRO NETO A. H., GUINEA F., PERES N. M. R., NOVOSELOV K. S. and GEIM A. K., Rev. Mod. Phys., 81 (2009) 109.
[3] LEMME M., Solid State Phenom., 156-158 (2010) 499.
[4] Katsnelson M. I., Novoselov K. S. and Geim A. K., Nat. Phys., 2 (2006) 620.
[5] McCann E. et al., Phys. Rev. Lett., 97 (2006) 146805.
[6] Morpurgo A. F. and Guinea F., Phys. Rev. Lett., 97 (2006) 296804.
[7] Novoselov K. S. et al., Nature, 438 (2005) 197.
[8] Zhang Y. B. et al., Nature, 438 (2005) 201.
[9] Park C.-H., Son Y.-W., Yang L., Cohen M. L. and Louie S. G., Nano Lett., 8 (2008) 2920.
[10] Stolyarova E. et al., Nano Lett., 9 (2009) 332.
[11] Low T. and Guinea F., Nano Lett., 10 (2010) 3551.
[12] Fujita M., Wakabayashi K., Nakada K. and Kusakabe K., J. Phys. Soc. Jpn., 65 (1996) 1920.
[13] Yang L., Park C.-H., Son Y.-W., Cohen M. L. and Louie S. G., Phys. Rev. Lett., 99 (2007) 186801.
[14] Son Y.-W., Cohen M. L. and Louie S. G., Nature, 444 (2006) 347.
[15] Arushkin D. A., Gunlycke D. and White C. T., Nano Lett., 7 (2007) 204.
[16] Biel B., Triozon F., Blase X. and Roche S., Nano Lett., 9 (2009) 2725.
[17] Akkermans E. and Montambaux G., Mesoscopic Physics of Electrons and Photons (Cambridge University Press) 2007.
[18] Lee P. A. and Ramakrishnan T. V., Rev. Mod. Phys., 57 (1985) 287.
[19] Hikami S., Larkin A. I. and Nagaoka Y., Prog. Theor. Phys., 63 (1980) 707.
[20] Bergman G., Phys. Rev. Lett., 48 (1982) 1046.
[21] Suzuura H. and Ando T., Phys. Rev. Lett., 89 (2002) 266603.
[22] Wu X. et al., Phys. Rev. Lett., 98 (2007) 136801.
[23] Tikhonenko F.V. et al., Phys. Rev. Lett., 100 (2008) 056802.
[24] Tikhonenko F.V. et al., Phys. Rev. Lett., 103 (2009) 226801.
[25] Winkler R. and Zülicke U., Phys. Lett. A, 374 (2010) 4003.
[26] Mecklenburg M. and Regan B. C., Phys. Rev. Lett., 106 (2011) 116803.
[27] Suzuura H. and Ando T., J. Phys. Soc. Jpn., 75 (2006) 024703.
[28] Imura K.-I., Kuranoto Y. and Nomura K., EPL, 89 (2010) 17009.
[29] Luttinger J. M., Phys. Rev., 84 (1951) 814–817.
[30] McClure J. W., Phys. Rev., 104 (1956) 666.
[31] Nomura K. and MacDonald A. H., Phys. Rev. Lett., 96 (2006) 256602.
[32] Zhang Y. Y. et al., Phys. Rev. Lett., 102 (2009) 106401.
[33] Roche S. and Mayou D., Phys. Rev. Lett., 79 (1997) 2518.
[34] Roche S. and Saito R., Phys. Rev. Lett., 87 (2001) 246803.
[35] Lherbier A., Biel B., Niquet Y.-M., Triozon F. and Roche S., Phys. Rev. Lett., 100 (2008) 036803.
[36] Moser J. et al., Phys. Rev. B, 81 (2010) 205445.
[37] Hong X., Cheng S.-H., Herding C. and Zhu J., Phys. Rev. B, 83 (2011) 085410.
[38] Poumirol J.-M. et al., Phys. Rev. B, 82 (2010) 041413(RC).
[39] Bai J. et al., Nat. Nanotechnol., 5 (2010) 655.
[40] Wakabayashi K., Takane Y. and Sigrist M., Phys. Rev. Lett., 99 (2007) 036601.