Density Inhomogeneity Driven Percolation Metal-Insulator Transition and Dimensional Crossover in Graphene Nanoribbons

S. Adam, S. Cho, M. S. Fuhrer, and S. Das Sarma

Condensed Matter Theory Center, Department of Physics, University of Maryland, College Park, Maryland 20742-4111, USA
Center for Nanophysics and Advanced Materials, Department of Physics, University of Maryland, College Park, Maryland 20742-4111, USA

(Received 16 April 2008; published 23 July 2008)

Transport in graphene nanoribbons with an energy gap in the spectrum is considered in the presence of random charged impurity centers. At low carrier density, we predict and establish that the system exhibits a density inhomogeneity driven two dimensional metal-insulator transition that is in the percolation universality class. For very narrow graphene nanoribbons (with widths smaller than the disorder induced length scale), we predict that there should be a dimensional crossover to the 1D percolation universality class with observable signatures in the transport gap. In addition, there should be a crossover to the Boltzmann transport regime at high carrier densities. The measured conductivity exponent and the critical density are consistent with this percolation transition scenario.

One of the remarkable experimental findings of the past two years, creating a great deal of activity and controversy, has been the observation that the carrier density dependent dc conductivity of gated 2D graphene layers, while being approximately linear in density at high gate voltage, becomes a nonuniversal constant over a finite gate voltage range \( \Delta V_g \) around the charge neutral Dirac point. While this conductivity minimum plateau formation around the charge neutrality point in 2D graphene is experimentally well established [1], the actual values of \( \sigma_{\text{min}} \approx 2 e^2/h \sim 20 e^2/h \) and \( \Delta V_g \sim 1-15 \text{ V} \) are nonuniversal and depend on the sample quality [2,3]. The minimum conductivity plateau in graphene has been theoretically explained [4,5] to arise from the invariable presence of unintentional random charged impurities at (or near) the graphene-substrate interface which lead to inhomogeneous electron-hole puddle formation in the low gate voltage regime [4–6]. We note that distortions of the graphene membrane and quenched ripples can also give rise to density inhomogeneities [7], and there have been recent theories studying the effect of ripples on graphene conductivity [8]. While we focus here on charged impurity induced inhomogeneities, many of our conclusions are only sensitive to the existence of the inhomogeneous density landscape (i.e., electron-hole “puddles”), and these do not distinguish between mechanisms (e.g., impurities, ripples) producing these puddles. Since graphene is a 2D semimetal (or more appropriately, a zero-gap 2D chiral semiconductor with electron-hole bands touching each other linearly at the charge neutral Dirac point), the conductivity becomes an approximate constant when the gate voltage induced chemical potential is pinned in this electron-hole puddle region around the Dirac point. This inhomogeneous electron-hole puddle based theoretical understanding of the graphene minimum conductivity plateau formation leads immediately to an important fundamental question: Are there situations where this inhomogeneous puddle picture leads to a graphene 2D metal-insulator transition (2D MIT) as is known [9,10] to occur in 2D semiconductor systems?

We show in this Letter that indeed, as a direct consequence of the inhomogeneous puddle formation in graphene, the system will manifest a 2D MIT, which is precisely in the same universality class as the corresponding 2D MIT in electron [9] and hole [10] GaAs systems, provided that there is an energy gap separating the graphene electron and hole bands. The fundamental physics here is that of percolation—for usual 2D zero-gap graphene, percolation through the puddles is allowed at all gate voltages, occurring either through the electron puddles or the hole puddles (or through both [11]), since one or the other is always percolating. If there is a gap, however, there should be a percolation-driven 2D MIT in graphene exactly as found [9,10] in 2D GaAs based semiconductor structures.

The easiest way to introduce an energy gap in graphene, which would then immediately lead to a percolation-induced transport gap (i.e., two separate 2D MIT transitions for electrons and holes), is to consider graphene nanoribbons instead of bulk 2D graphene. In this Letter, we predict and confirm experimentally that graphene nanoribbons exhibit a 2D MIT in the low carrier density regime as a function of the applied gate voltage and that this MIT is in the percolation universality class; furthermore, we predict theoretically that as the ribbons become very narrow, there should be a dimensional crossover to the 1D universality, implying that the observed transport gap would tend to infinity as the ribbon width goes to zero (or in practice, becomes smaller than the typical size of the puddles), reflecting the 1D percolation universality where metallic conduction is completely suppressed. We speculate that such a 2D-1D crossover may have been observed in recent...
three samples are a function of applied gate voltage. The corresponding fit parameters for the graphene samples [3] are consistent with measurements on similarly prepared bulk graphene nanoribbons [3]. Figure 2 shows a similar analysis for three samples fabricated at Maryland and two of the Columbia samples (W = 24 nm and W = 49 nm) reported in Ref. [12]. All three samples show the low-density percolation universality class with critical exponents $\delta = 1.2 \pm 0.2, 1.3 \pm 0.1$, and $1.55 \pm 0.3$, respectively, which are similar to percolation exponents observed in 2D GaAs systems [9,10]. The corresponding fit parameters for the three samples are $W/L = 0.008, 0.002$, and $0.0087$, respectively, and $n_c = -65.08, 170.87$, and $184.23 \times 10^{10}$ cm$^{-2}$, respectively [the units of $A$ are $(10^{10}$ cm$^{-2})^2$25.8 kΩ$^{-1}$].

For the 2D percolation universality class, at low density we have

$$\sigma = A(n - n_c)^\delta,$$

where $\delta = 4/3$ is the 2D percolation critical exponent. For graphene ribbons, we expect two such percolation transitions, one for electrons and one for holes, separated by a “transport gap” defined as $\Delta_g = \gamma \sqrt{\pi(n_c^e - n_c^h)}$, where $n_c^e$ (holes) is the critical density for electrons (holes) and $\gamma = h v_F / 2$ is the graphene Fermi velocity. For larger carrier densities, where $|E_F| \gg \Delta_g$, we expect a crossover to a high-density Boltzmann transport regime where [4,5]

$$\sigma = 20 e^2 / h \left( \frac{n - n_p}{n_{imp}} \right),$$

just as for bulk graphene on a SiO$_2$ substrate (where $n_p$ is the charge neutrality point and $n_{imp}$ is the 2D surface impurity density of Coulomb scatterers). If we define $\xi$ as the typical size of the electron or hole puddle, then below we calculate $\xi$ self-consistently using the random phase screening approximation, then so long as the sample width $W \geq \xi$, we would have 2D percolation whereas if $W \leq \xi$ one has 1D percolation, i.e., a chain of approximately $L/\xi$ p-n junctions. Changing $n_{imp}$ (which could be extracted from high-density mobility measurements) would also change $\xi$ and the critical width for which this dimensional crossover is observed. Moreover, we predict that signatures of reduced dimensionality should be apparent in temperature dependent transport measurements.

---

FIG. 1 (color online). Evidence of percolation-driven metal-insulator transition in a graphene nanoribbon. Main panel shows graphene ribbon conductance as a function of gate voltage. Best fits at low density to Eq. (1) give for electrons $A = 1.485, n_c = 26.7485 \times 10^{10}$ cm$^{-2}$, $\delta^e = 1.3 \pm 0.2$ and for holes $A = 1.755, n_c = 18.5 \times 10^{10}$ cm$^{-2}$, $\delta^h = 1.3 \pm 0.1$. The right-hand panel shows the same data in a linear scale. Inset shows the same data in a linear scale, where even by the eye the transition from high-density Boltzmann behavior to the low-density percolation transport is visible.

FIG. 2 (color online). Percolation-driven metal-insulator transition in three additional graphene samples. The left-hand panel shows a naturally occurring graphene nanoribbon with dimensions $W = 200$ nm and $L \approx 11$ $\mu$m and has a critical exponent $\delta^h = 1.2 \pm 0.2$. The center panel is the $W = 24$ nm sample reported in Ref. [12] which has a critical exponent of $\delta^h = 1.3 \pm 0.1$, and the right-hand panel is the $W = 49$ nm sample reported in Ref. [12] which has a critical exponent of $\delta^h = 1.6 \pm 0.3$. 

---

PRL 101, 046404 (2008)  
PHYSICAL REVIEW LETTERS  
week ending 25 JULY 2008
To date, most theories for transport in graphene nanoribbons consider a quasi-1D, rather than the 2D limit. The experimental observation of 2D percolation in these ribbons casts strong doubt onto the relevance of the quasi-1D theories for current graphene nanoribbon experiments. As was already discussed in Ref. [14], many features of the quasi-1D geometry get washed out for $W \approx 10$ nm, which is the case in most experiments on graphene nanoribbons. The length scale controlling the crossover from quasi-1D to 2D behavior in this context may very well be determined by other independent parameters which are unknown at this stage, such as the inelastic scattering length or the phase breaking length (both of which depend on temperature). Whether the transport properties in graphene nanoribbons should be considered using a 2D or a quasi-1D Hamiltonian is at this stage an open question requiring further theoretical and experimental study. Our analyses involving data from two groups, our own and that of the Columbia group [12], clearly establish that depending on the value of $W$ either the 2D or 1D percolation universality class may apply, where we believe this transition to be controlled by the size of the electron and hole puddles induced by charged impurities. We cannot rule out the possibility that further lowering of temperature would lead to quasi-1D behavior [15] and the percolation-driven 2D MIT is only a crossover phenomenon. Although we focus on single-layer graphene, we note that a similar percolation transition should also be seen in graphene bilayers, where, since an electric field induced gap can be introduced into the spectrum without any confinement, the crossover to a quasi-1D regime would not arise. We note that even for bulk graphene, a Boltzmann to percolation crossover could be induced with a magnetic field, where for small field and within the electron-hole puddle model, we expect the $p$-$n$ resistance to be very low justifying the Boltzmann picture, whereas for large magnetic field, the $p$-$n$ junction becomes very resistive [16] inducing a percolation transition. This crossover may have been observed in recent experiments [17].

To reinforce the point that the 2D MIT in graphene nanoribbons is indeed a percolation transition and not a quantum crossover phenomenon, we calculate the percolation critical density $n_p$ using the nonlinear screening argument of Efros [18] with the basic idea being that the MIT occurs when inhomogeneous density fluctuations created by the charged impurities can no longer be screened by the carriers. This leads to $n_p \approx \sqrt{n_{\text{imp}}}/d$ where the random charged impurities of concentration $n_{\text{imp}}$ are assumed to be located at a distance $d$ from the 2D graphene plane. Taking $d \approx 1$ nm and $n_{\text{imp}} \sim 2-5 \times 10^{11}$ cm$^{-2}$, typical values estimated [2–5] from mobility measurements, we get $n_p \sim 5 \times 10^{12}$ cm$^{-2}$. This is in reasonable agreement with our experimental finding in Figs. 1 and 2.

On the other hand, the quantum localization crossover density $n_q$ can be estimated from the Ioffe-Regel criterion $k_F \ell \sim 1$, where $\ell$ is the mean-free path, to be $n_q \sim 2 \times 10^{10}$ cm$^{-2}$ for the same $n$, and $d$ values. Thus, $n_q \ll n_p$, and our experimental critical density agrees with the percolation critical density, providing further support for a percolation-driven insulating transition in graphene.

Experimentally, one can measure three different gaps. In addition to the transport gap $\Delta_g$ discussed above, the temperature dependence of the conductivity minimum gives an activated gap $\Delta_{\text{act}}$ (we performed this measurement on the 200 nm ribbon, and found $\Delta_{\text{act}} W = 0.1$ eV nm, which is an order of magnitude smaller than theoretical estimates [19] of the confinement induced gap), and finally, Ref. [12] reported the source-drain bias required to induce conduction and found the gap to be orientation independent (contrary to the expectation of the quasi-1D theory that is extremely sensitive to whether the edge is zigzag or armchair). The connection between these three experimental gaps and the theoretical gap in the energy spectrum is beyond the scope of this work, where we focus here only on the transport gap $\Delta_g$ and predict that in the 2D regime, provided that the impurity location is pretty much the same, $|n_q^e - n_q^o| \approx \sqrt{n_{\text{imp}}} \propto \mu^{-1/2}$, where $\mu$ is the high-density mobility in the Boltzmann regime [2–5].

The 2D percolation picture presented above breaks down when the sample width becomes smaller than the typical disorder length scale. Using the self-consistent RPA method of Ref. [5] we can obtain an integral expression for the potential correlation function $\langle V(r)V(0)\rangle$, which for experimentally relevant parameters can be approximated by

$$\langle V(r)V(0)\rangle = \frac{K_0}{2\pi} \frac{\gamma^2}{\xi^2} \exp\left[\frac{-r^2}{2\xi^2}\right].$$

Using $r_s = e^2/\kappa \gamma$, where $\kappa$ is the effective dielectric constant that depends on the choice of substrate, we find

$$K_0 = \frac{1}{4\pi^2} \left(\frac{D_0}{C_0}\right)^2,$$

and

$$\xi = \frac{1}{\sqrt{n_{\text{imp}}}} \frac{1}{4\pi^2 r_s} \left(\frac{C_0}{D_0}\right)^{1/2},$$

where $z = 4k_F d$ and $E_1(z) = \int_1^{\infty} t^{-1} e^{-t} dt$,

$$C_0(z) = -1 + \frac{4E_1(z)}{(2 + \pi r_s)^2} + \frac{2e^{-r_s}}{1 + 2r_s}$$

$$+ \left(1 + 2z r_s\right) e^{2z r_s} (E_1(2z r_s) - E_1(z(1 + 2r_s))),$$

and

$$D_0(z) = 1 - \frac{8r_se^{2z r_s} (E_1(z) + E_1(z(1 + 2r_s)))}{(2 + \pi r_s)^2} + \frac{8e^{-z r_s}}{1 + 2r_s}$$

$$- 2z r_s e^{2z r_s} (E_1(2z r_s) - E_1(z(1 + 2r_s))).$$

This notation is chosen to be consistent with Ref. [5] where the rms density $n^* = 2r_se^{2z r_s} C_0(z) = 4d\sqrt{\pi n^*}$. For typical values of $n_{\text{imp}} \approx 20 \times 10^{10}$ cm$^{-2}$ and $d = 1$ nm, we have
behavior with an effective infinite energy transport gap at $T = 0$, which may have been observed experimentally [12,13], by virtue of the absence of a percolation transition in 1D. A fundamental question of considerable significance that remains open in this context is the experimental absence of quantum localization [21], which may be observable at much lower temperatures than used experimentally so far.

This work is partially supported by U.S. ONR.