Coulomb Drag of Massless Fermions in Graphene

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Using a novel structure, consisting of two, independently contacted graphene single layers separated by an ultra-thin dielectric, we experimentally measure the Coulomb drag of massless fermions in graphene. At temperatures higher than 50 K, the Coulomb drag follows a temperature and carrier density dependence consistent with the Fermi liquid regime. As the temperature is reduced, the Coulomb drag exhibits giant fluctuations with an increasing amplitude, thanks to the interplay between coherent transport in the graphene layer and interaction between the two layers.

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Bilayer systems formed by two layers of carriers in close proximity are a fascinating testground for electron physics. In particular, the prospect of electron-hole pair (indirect exciton) formation, and dipolar superfluidity has fueled the research of electron-hole bilayers in GaAs/AlGaAs heterostructures. Graphene is a particularly interesting material to explore interacting bilayers. The symmetric conduction and valence bands, and the large Fermi velocity favor correlated electron states at elevated temperatures. The zero energy band-gap allows a seamless transition between electrons and holes in each layer, and obviates the large inter-layer electric field required to simultaneously induce electrons and holes in GaAs bilayers. Coulomb drag, a direct measurement of inter-layer electron-electron scattering can provide insight into the ground state of two- and one- electron systems, as well as correlated bilayer states. Here we demonstrate a novel, independently contacted graphene bilayer, and investigate the Coulomb drag in this system.

Two main ingredients render the realization of independently contacted graphene bilayers challenging. First, an ultra-thin yet highly insulating dielectric is required to separate the two layers. Second, a method to position another graphene layer on a pre-existing device with minimum or no degradation is needed to create the second layer of the structure investigated here. The fabrication of our independently contacted graphene bilayers is described in Fig. 1. First, the bottom graphene layer is mechanically exfoliated onto a 280 nm thick SiO2 dielectric, thermally grown on a highly doped Si substrate. E-beam lithography, metal lift-off, and etching are used to define a Hall bar on the bottom layer [Fig. 1(a)]. A 7 nm thick Al2O3 is then deposited on the bottom layer using a 2 nm oxidized Al interfacial layer, followed by 5 nm of Al2O3 atomic layer deposition. The second, top graphene layer is also mechanically exfoliated on a similar SiO2/Si substrate. A poly methyl metacrylate (PMMA) film is applied on the top layer and cured. Using an NaOH etch, the PMMA film along with the graphene layer, and the alignment marks are detached from the host substrate, forming a free standing membrane. The membrane is placed face down on the substrate containing the bottom graphene layer [Fig. 1(b)], and aligned with it. A Hall bar is subsequently defined on the top layer [Fig. 1(c)]. Ten back-gated, independently contacted graphene bilayers have been fabricated and investigated in this study, all with similar results. We focus here on data collected from three samples, labeled 1, 2, and 3, with mobilities between 4,200-12,000 cm2/Vs for the bottom layer, 4,500-22,000 cm2/Vs for the top layer, and with inter-layer resistances of 1-20 GΩ. These structures are markedly different from graphene bilayers exfoliated from natural graphite, consisting of two graphene monolayers in Bernal stacking.

We now turn to the individual layer characterization. The layer resistivities (ρ) and Hall densities measured for sample 1 at a temperature T=4.2 K, as a function of back-gate bias (VBG) are shown in Fig. 2(a) and 2(b), respectively. The potential of the both layers is held at zero (ground) for all measurements presented in this study. The bottom layer dependence on the applied VBG shows ambipolar conduction and a finite resistance at the charge neutrality (Dirac) point, consistent with the expected response of gated monolayer graphene. More interestingly, the top layer resistivity also changes as a result of the applied VBG. This observation indicates an incomplete screening of the gate-induced electric field by the bottom layer, which is most pronounced in the vicinity of the charge neutrality point, consequence of the reduced density of states in graphene. As we show below, we can quantitatively explain the layer resistivities and densities dependence on VBG.

Figure 2(c) shows the band diagram of the graphene bilayer at VBG=0 V; for simplicity the gate Fermi energy and the charge neutrality point in the two layers are assumed to be at the same energy. Once a finite VBG is applied, finite charge densities are induced in both top (nT) and bottom (nB) layers [Fig. 2(d)]. The difference between the gate and bottom layer Fermi levels is distributed partly across the SiO2 dielectric, and partly on the Fermi energy of the bottom graphene layer: εVBG = e2(nB + nT)/CSiO2 + E_F(nB) (1); E_F(n) = hν_F/√πn is the graphene Fermi energy measured with respect to the charge neutrality point at a carrier density n, ε is the electron charge, ν_F=1.1×106 m/s is the graphene Fermi velocity, and CSiO2 denotes the SiO2 di-
FIG. 1: (color online) Optical micrographs (top) and schematic representation (bottom) of the fabrication process of an independently contacted graphene bilayer. (a) Hall bar device fabrication on the bottom graphene layer, followed by the Al₂O₃ inter-layer dielectric deposition. (b) Top graphene layer isolation on a separate substrate, followed by transfer onto the bottom layer. (c) Top layer Hall bar realization by etching, lithography, metal deposition, and lift-off. The yellow (inner) and red (outer) dashed contours in the optical micrograph represent the top and bottom layers, respectively. The scale bars in all panels are 10 µm.

FIG. 2: (color online) (a) Layer resistivities, and (b) densities vs. $V_{BG}$ measured at $T=4.2$ K in sample 1. Depending on $V_{BG}$, both electrons or holes can be induced in the bottom layer; the top layer contains electrons in the available $V_{BG}$ window, owing to unintentional doping. The symbols in panels (a) and (b) represent experimental data, while the lines represent the calculated values according to Eqs. (1) and (2). (c) Band diagram across the graphene bilayer heterostructure at $V_{BG}=0$ V, and (d) at a positive $V_{BG}$. Both layers are assumed to be at the charge neutrality point, and aligned with the back-gate Fermi level at $V_{BG}=0$. The layers are held at the ground potential, and their thicknesses exaggerated to show the Dirac cones. The applied $V_{BG}$ induces voltage drops $V_{SiO_2}$ and $V_{Al_2O_3}$ across the bottom, and inter-layer dielectrics respectively.

Electric capacitance per unit area. Similarly, the Fermi energy difference between the two layers is responsible for the potential drop across the Al₂O₃ inter-layer dielectric: $E_F(n_B) = e^2 n_T/C_{Al_2O_3} + E_F(n_T)$ (2); $C_{Al_2O_3}$ is the Al₂O₃ dielectric capacitance per unit area. The finite Fermi energy of the bottom layer, $E_F(n_B)$ in Eq. (2), plays the same role with respect to the top layer, as the applied $V_{BG}$ in Eq. (1) for the bottom layer. Equations (1) and (2) allow us to determine $n_B$ and $n_T$ as a function of $V_{BG}$. This model can be adjusted to include finite layer densities at $V_{BG}=0$ V. The layer resistivity dependence on $V_{BG}$ can be understood using a Drude model $\rho_{T,B} = (n_{T,B}^\ast \mu_{T,B})^{-1}$, where $\mu_T$ and $\mu_B$ are the top and bottom layer mobilities, and the layer densities $n_{T,B} = \sqrt{n_{T,B}^2 + n_{0T,0B}^2}$ are adjusted to allow for finite carrier densities ($n_{0T}, n_{0B}$) at the charge neutrality point. The data of Figs. 2(a) and 2(b) show a good agreement between the measured layer resistivities and densities (symbols), and the calculations (solid lines). The layer mobilities, determined from Hall measurements, are $\mu_B = 5,400$ cm²/Vs, and $\mu_T = 4,500$ cm²/Vs at $T=4.2$ K.

Key insight into the physics of the graphene bilayer system can be gained from Coulomb drag measurements. A current ($I_{Drive}$) flown in one (drive) layer leads to a momentum transfer between the two layers, thanks to the inter-layer electron-electron interaction. To counter this momentum transfer, a longitudinal voltage ($V_{Drag}$) builds up in the opposite (drag) layer. The polarity of
V_{\text{Drag}} depends on the carrier type in the two layers, and is opposite (same) polarity as the voltage drop in the drive layer when the both layers have the same (opposite) type of carriers. The drag resistivity is defined as 
\[ \rho_{\text{Drag}} = (W/L)V_{\text{BG}}/I_{\text{Drive}}, \]
where \( L \) and \( W \) are the length and width of the region where drag occurs. \( \rho_{\text{Drag}} \) vs. \( V_{\text{BG}} \) measured at \( T=250 \) K in sample 2 is shown in Fig. 3(a), along with the layer resistivities, \( \rho_T \) and \( \rho_B \). Unlike sample 1 data (Fig. 2), the charge neutrality (Dirac) points of both layers can be captured in the experimentally accessible \( V_{\text{BG}} \) window. Consequently, depending on the \( V_{\text{BG}} \) value, sample 2 can probe three different regimes: a hole-hole bilayer, for \( V_{\text{BG}} < -15 \) V, an electron-hole bilayer for \(-15 \) V < \( V_{\text{BG}} < -2 \) V, and an electron-electron bilayer for \( V_{\text{BG}} > -2 \) V. The dependence of \( \rho_B \) and \( \rho_T \) on \( V_{\text{BG}} \) of Fig. 3 is also in good agreement with the model presented in Fig. 2. Consistent with the above argument, \( \rho_{\text{Drag}} \) is positive in the electron-hole bilayer regime, negative in the hole-hole or electron-electron regime, and changes sign when either the top or the bottom layer are at the charge neutrality point. Standard consistency checks\(^9\) ensured the measured drag is not affected by inter-layer leakage current.

For two closely spaced two-dimensional systems, when the ground state of each layer is a Fermi liquid, and the inter-layer interaction is treated as a perturbation, the \( \rho_{\text{Drag}} \) depends on layer density \( n \) as \( \propto 1/n^{3/2} \), on temperature as \( \propto T^2 \), and inter-layer distance \( d \) as \( \propto 1/d^{1.9} \). Likewise, the Coulomb drag resistivity in graphene, calculated in the Fermi liquid regime using Boltzmann transport formalism and the random phase approximation for the dynamic screening \( k_B T/\epsilon \), can be expressed as:
\[ \rho_{\text{drag}} = -\frac{h}{d^2} \frac{\zeta(3)}{\epsilon^2} \frac{(k_B T)^2}{n_B n_T^2}. \]

The discrepancy, a possible explanation is that Eq. 3 is valid for high densities and/or large inter-layer spacing such as

\[ 10 \]

\[ r_{\text{Dr}} \]

\[ 77K \]

\[ 50K \]

\[ 20K \]

\[ 4.2K \]
that $k_F \cdot d \gg 1$ ($k_F$ denotes the Fermi wave-vector)\textsuperscript{18}, for Fig. 3 data $k_F \cdot d \leq 3$ at all layer densities. The $\rho_{\text{Drag}} \propto (k_BT)^2$ dependence, which stems from the allowed phase space where electron-electron scattering occurs, is followed closely for temperatures between 70 K and 200 K (Fig. 3(b) inset), and softens for $T > 200$ K. Figure 3 data shows a smooth crossover for $\rho_{\text{Drag}}$ through 0 Ω, from the electron-hole to the electron-electron regime [blue (shaded) corridor of Fig. 3(b)]. The crossover can be explained by the co-existence of electron and hole puddles near the charge neutrality point of the bottom layer, which generate drag electric fields of opposite sign, and cancel the $\rho_{\text{Drag}}$.

A remarkable transition in the drag resistance is observed for $T$ lower than 50 K (Fig. 4). As $T$ is reduced, the $\rho_{\text{Drag}}$ data starts to develop fluctuations superposed on the average $\rho_{\text{Drag}}$ vs. $V_{BG}$ dependence of Fig. 3 and Eq.(3), valid for diffusive transport. The $\rho_{\text{Drag}}$ fluctuations, which are reproducible in different measurements, grow in amplitude as $T$ is reduced, and fully obscure the average diffusive drag below 20 K. This manifestation of mesoscopic physics at elevated temperatures is a consequence of the phase coherence length ($L_\varphi$) increasing with reducing $T$, and represents the counterpart of universal conductance fluctuations\textsuperscript{20} in Coulomb drag\textsuperscript{21,22}. Figure 4 data reveal that $\rho_{\text{Drag}}$ fluctuation amplitude reaches a maximum near the charge neutrality point of the bottom layer ($V_{BG} = -1$ V), and increases, albeit slowly as $T$ is decreased. Theoretical arguments\textsuperscript{22} indicate that the drag conductivity ($\sigma_{\text{Drag}} = \rho_{\text{Drag}}/\rho_T \rho_B$) fluctuation amplitude ($\delta \sigma_{\text{Drag}}$) depends on relevant length scales and temperature as $\delta \sigma_{\text{Drag}} \propto T \cdot (L_\varphi^2)/l$; $l$ is the electron mean free path. For the temperature range examined in Fig. 4, $l$ can be considered constant, as the mobility is weakly dependent on $T$. Assuming the electron-electron interaction is the main phase-breaking mechanism in graphene\textsuperscript{23}, hence $L_\varphi = l\sqrt{E_F/2k_BT}$, the $T$ dependence of $\delta \sigma_{\text{Drag}}$ and $\delta \rho_{\text{Drag}}$ should follow a $\propto T^{-1/2}$ dependence, in good agreement with Fig. 4 data.

To probe the signature of weak localization in Coulomb drag, in Fig. 5 we show an example of $\rho_{\text{Drag}}$ vs. perpendicular magnetic field ($B$) data, measured in sample 3 at $T=0.3$ K and $V_{BG}=0$ V: both layers contain electrons with layer densities $n_T = 1.4 \times 10^{11}$ cm$^{-2}$, and $n_B = 1.5 \times 10^{11}$ cm$^{-2}$. Similar to the $V_{BG}$ dependence of Fig. 4, the $\rho_{\text{Drag}}$ vs. $B$ data shows reproducible mesoscopic fluctuations. The auto-correlation function ($C(\Delta B)$) of Fig. 5 data reveals a correlation field $B_c = 47$ mT, which corresponds to a phase coherence length $L_\varphi = \sqrt{h/eBc} = 300$ nm. Similar $L_\varphi$ values have been extracted from ensemble average measurements using scanning gate microscopy\textsuperscript{24}.

In summary, we demonstrate independently contacted graphene bilayers, and probe the Coulomb drag in this system. At elevated temperatures the drag resistance dependence on density and temperature are consistent with the Fermi liquid theory. At reduced temperatures, the drag exhibits mesoscopic fluctuations which obscure the average drag, a result of the interplay between electron-electron interaction and phase coherent transport.

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\begin{thebibliography}{24}
\bibitem{1} E. Lozovik, V. I. Yudson, JETP Lett. 22, 274 (1975).
\bibitem{2} M. Pohlt et al., Appl. Phys. Lett. 80, 2105 (2002).
\bibitem{3} A. F. Croxall et al., Phys. Rev. Lett. 101, 246801 (2008); J. A. Seamons, C. P. Morath, J. L. Reno, M. P. Lilly, Phys. Rev. Lett. 102, 026804 (2009).
\bibitem{4} K. S. Novoselov et al., Science 306, 666 (2004).
\bibitem{5} K. S. Novoselov et al., Nature 438, 197 (2005); Y. Zhang et al., Nature 438, 201 (2005).
\bibitem{6} H. Min, R. Bistritzer, J.-J. Su, A. H. MacDonald, Phys. Rev. B 78, 121401 (2008).
\bibitem{7} C.-H. Zhang, Y. N. Joglekar, Phys. Rev. B 77, 233405 (2008).
\bibitem{8} P. M. Solomon, P. J. Price, D. J. Frank, D. C. La Tulipe, Phys. Rev. Lett. 63, 2508 (1989).
\bibitem{9} T. J. Gramila, J. P. Eisenstein, A. H. MacDonald, L. N. Pfeiffer, K. W. West, Phys. Rev. Lett. 66, 1216 (1991).
\bibitem{10} M. Yamamoto, M. Stopa, Y. Tokura, Y. Hirayama, S. Tarucha, Science 313, 204 (2006).
\bibitem{11} G. Vignale, A. H. MacDonald, Phys. Rev. Lett. 76, 2786 (1996).
\bibitem{12} E. Tutuc, R. Pillarisetty, M. Shayegan, Phys. Rev. B 79, 041303(R) (2009).
\bibitem{13} S. Kim et al., Appl. Phys. Lett. 94, 062107 (2009).
\bibitem{14} A. Reina et al., J. Phys. Chem. C 112, 17741 (2008).
\bibitem{15} E. McCann, V. I. Fal’ko, Phys. Rev. Lett. 96, 086805 (2006).
\bibitem{16} S. Adam, E. H. Hwang, V. M. Galitski, S. Das Sarma, J. A. Seamons, C. P. Morath, J. L. Reno, M. P. Lilly, Phys. Rev. Lett. 104, 246804 (2010).\end{thebibliography}
Proc. Natl. Acad. Sci. U.S.A. 104, 18392 (2007).
17 H. Schmidt et al., Appl. Phys. Lett. 93, 172108 (2008).
18 W.-K. Tse, B. Y.-K. Hu, S. Das Sarma, Phys. Rev. B 76, 081401 (2007).
19 B. N. Narozhny, Phys. Rev. B 76, 153409 (2007).
20 P. A. Lee, A. D. Stone, Phys. Rev. Lett. 55, 1622 (1985).
21 B. N. Narozhny, I. L. Aleiner, Phys. Rev. Lett. 84, 5383 (2000).
22 A. S. Price, A. K. Savchenko, B. N. Narozhny, G. Allison, D. A. Ritchie, Science 316, 99 (2007).
23 B. L. Altshuler, A. G. Aronov, Electron-electron interactions in disordered systems (North-Holland, Amsterdam, 1985).
24 J. Berezovsky, R. M. Westervelt, Nanotechnology 21, 274014 (2010).