

Conduction in carbon nanotubes through metastable resonant states

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Abstract. – We report here on electrical measurements on individual multi-walled carbon nanotubes that show that the presence or movement of impurities or defects in the carbon nanotube can radically change its low-temperature transport characteristics. The low-temperature conductance can either decrease monotonically with decreasing temperature, or show a sudden increase at very low temperatures, sometimes in the same sample at different times. This unusual behavior of the temperature dependence of the conductance is correlated with large variations in the differential conductance as a function of the dc voltage across the wire. The effect is well described as arising from quantum interference of conduction channels corresponding to direct transmission through the nanotube and resonant transmission through a discrete electron state, the so-called Fano resonance.

There is tremendous interest in the transport properties of carbon nanotubes due to their potential for use in future nanodevices [1], and from their role as canonical models of one-dimensional electron transport [2]. While individual single-walled carbon nanotubes (SWNTs) are expected to show either semiconducting or metallic behavior depending on their chirality [3], the presence of impurities, defects and interactions is expected to modify this behavior [4, 5]. However, measuring the transport properties of single or multiwalled carbon nanotubes is complicated by the experimental problem of making electrical contact to the nanotube. Transport measurements on high contact-resistance carbon nanotube devices by a number of groups have elucidated the wide variety of physical problems that can be studied in these systems, including the Coulomb blockade [6], the Kondo effect [7], and Luttinger liquids [2]. In more recent experiments on devices with lower contact resistances, other interesting effects have been reported. For example, by using different techniques to obtain low-resistance contacts, observation of quantization of the conductance [8] and Fabry-Perot

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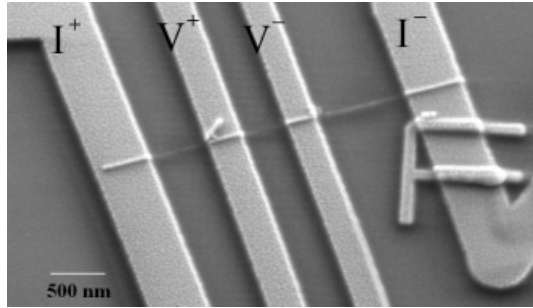


Fig. 1 – Scanning electron microscope (SEM) image of Sample 3, a single MWNT with four electrodes. The legends show the probe configurations used for making the four-terminal resistance measurements. For this sample, each electrode had two external contacts, so that the four-terminal resistance of the two inner metal-nanotube contacts could be directly measured.

interference [9] in ballistic carbon nanotubes has been reported, and there are also suggestions of superconductivity in nanotube rope devices made with low-resistance contacts [10]. However, much interesting physics still remains to be explored.

In this letter, we report on transport measurements on multiwalled carbon nanotubes (MWNTs) with low-resistance contacts. The low-temperature differential conductance $G(V_{dc})$ of these devices as a function of the dc voltage bias V_{dc} across them is highly asymmetric, showing large, reproducible fluctuations that can be as large as 10% of the total conductance. This conductance “fingerprint” can change on thermal cycling to just 2 K, indicating that it might be associated with metastable impurities or mechanical instabilities. The fluctuations in $G(V_{dc})$ are reflected in the zero-bias ($V_{dc} = 0$) conductance as a function of temperature $G(T)$, which can either decrease or increase with decreasing temperatures at low temperatures, sometimes showing both dependences in a single sample upon thermal cycling. The sharp structure in $G(V_{dc})$ can be described as arising from Fano resonances through resonant states in the device. The positions of these peaks and dips frequently change on thermal cycling, indicating that the resonant states may arise from metastable impurities or defects in the device.

Our devices consist of an isolated MWNT with four Au/Ti electrodes (fig. 1). The arc-grown MWNTs, typically 2–5 μm long and 25–50 nm wide, were spun onto an Si substrate with a 500 nm thick oxide layer from a suspension in dimethylformamide. After locating the nanotubes with respect to fiducial marks, and patterning electrodes by electron-beam lithography, thin film electrodes (5 nm Ti/50 nm Au) were deposited to make contact to the nanotube. Prior to deposition, a short-time oxygen plasma etch was used to improve the metal-nanotube contact resistance. High-resolution TEM studies of the etching process showed no visible damage to the MWNTs. Without this cleaning, the contact resistances were typically in the range of a few k Ω to a few M Ω , while this process reduced the contact resistance to a few hundred ohms. The presence of four electrodes enables us to make four-terminal resistance measurements on the MWNTs. In some devices, two external contacts were made to each electrode, as shown in fig. 1, enabling direct measurements of the electrode-nanotube contact resistance; in the other devices, the contact resistance was inferred from differences between four-terminal and two-terminal measurements. The typical distance between the voltage probes was 300 nm. The samples were measured in a ^3He refrigerator and a dilution refrigerator. A home-made ac resistance bridge was used to measure the four-terminal differential resistance dV/dI as a function of dc current I_{dc} using the probe configuration shown in fig. 1, with ac excitation (10–100 Hz) in the range of 0.25–1 nA to avoid heating effects. In our early experiments, we found

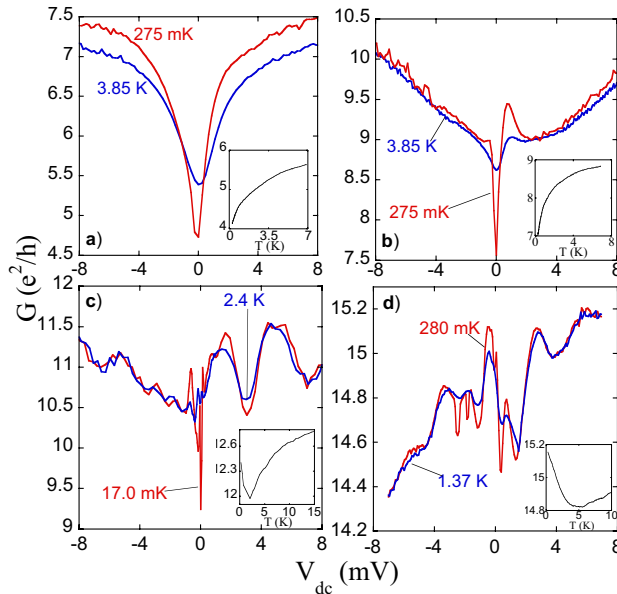


Fig. 2 – $G(V_{dc})$ of 4 samples, in units of e^2/h . The insets show $G(T)$. The average contact resistances (per contact) are a) Sample 1: $1\text{ k}\Omega$ at 4 K ; b) Sample 2: $900\ \Omega$ at 4 K ; c) Sample 3: $133\ \Omega$ and $173\ \Omega$ at 12 K for the two inner contacts; d) Sample 4: less than $100\ \Omega$ at 4 K . Contact resistances were inferred from differences between 2- and 4-terminal measurements of the devices, except for Sample 3, where they could be measured directly.

the conductance had no measurable dependence on the voltage (up to 1 volt) applied to the substrate underneath the oxide layer. This is consistent with the observations of other groups that metallic MWNTs with low-resistance contacts showed no appreciable gate voltage dependence [5]. Consequently, in our later experiments, the substrates were kept floating. In this paper, we shall plot our measurements in terms of the differential conductance $G = 1/(dV/dI)$ as a function of the dc voltage V_{dc} , obtained by numerically integrating the measured dV/dI *vs.* I_{dc} curves.

Figure 2 shows $G(V_{dc})$ for four different samples with contact resistances ranging from $1\text{ k}\Omega$ to less than $100\ \Omega$, at two different temperatures each, and demonstrates the sample-specific behavior that can be seen in our devices. These samples represent behavior seen in over 14 devices that we have measured. $G(V_{dc})$ is highly asymmetric and shows large fluctuations, with the pattern of the fluctuations being different for different devices. In general, the structure in $G(V_{dc})$ becomes sharper as the temperature is lowered: the peaks increase in conductance, while the valleys decrease in conductance. This gives rise to characteristically different temperature dependences for nominally identical samples: for example, if a peak is observed at $V_{dc} = 0$, $G(T)$ will increase with decreasing temperature, while if a valley is observed, it will decrease. (Of course, it is not necessary that either a peak or a valley occur at $V_{dc} = 0$.) This is demonstrated in the insets to the panels in fig. 2, which show the corresponding $G(T)$ for each device. As demonstrated in fig. 2, the structure in $G(V_{dc})$ appears to increase on average as the resistance of the contacts decreases. Indeed, large fluctuations in $G(V_{dc})$ are not seen in our samples with contact resistances in the $\text{k}\Omega$ to $\text{M}\Omega$ range; one sees instead a largely symmetric curve with reduced conductance near $V_{dc} = 0$, and correspondingly, a monotonically decreasing $G(T)$.

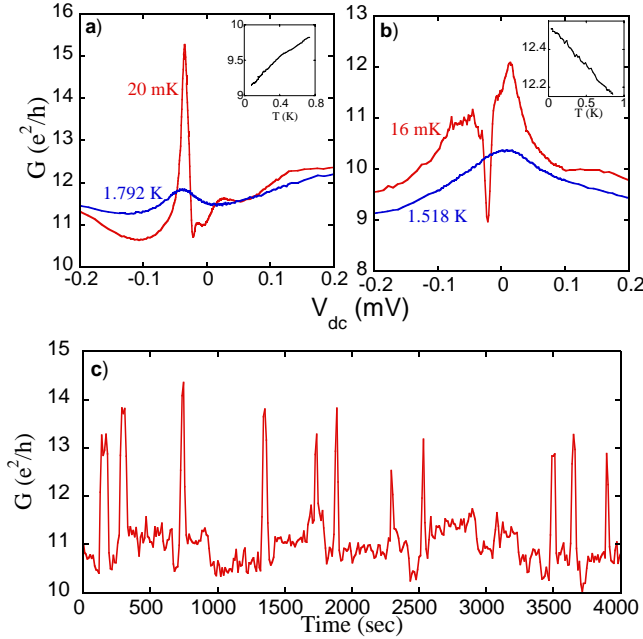


Fig. 3 – a), b) $G(V_{dc})$ of Sample 3 at two temperatures, on two different cooldowns of the dilution refrigerator, in units of e^2/h . a) First cooldown. b) Second cooldown. The insets show the measured temperature dependence of $G(T)$. Note that $G(T)$ does not match $G(V_{dc} = 0)$ exactly, as the sample characteristics changed even on warming to 2 K. c) Time-dependent data taken at a temperature of 35 mK.

$G(V_{dc})$ varies not only from sample to sample, but can also change in a single sample as a function of thermal cycling. Figures 3(a) and (b) show $G(V_{dc})$ for the sample of fig. 2(c), for two different cooldowns of the dilution refrigerator. As can be seen, there is a radical change after warming to room temperature: while the most prominent feature at low temperature in fig. 3(a) is a peak, this feature has changed into a dip in fig. 3(b). As we noted before, this change is also reflected in $G(T)$, as shown in the inset to the panels in fig. 3. Indeed, we found that it was not necessary to warm the samples all the way to room temperature; warming to temperatures on the order of a few kelvins changed the behavior of the sample, although the change was not as dramatic as shown in figs. 3(a) and (b). In our final measurements on this particular device, we also observed that the sample switched between two different conductance states, as shown in fig. 3(c). The conductance difference between the two states is a few e^2/h , much larger than our measurement noise. Similar “telegraph noise” has been observed in other mesoscopic systems where it is associated with an impurity or defect moving between two metastable states [11]. We believe the same phenomenon also exists in our samples.

Sample-specific fluctuations of the conductance are well known in mesoscopic systems. In the case of disordered metals and semiconductors, they are associated with quantum interference of electron waves that are scattered by impurities, defects or grain boundaries in the sample [12]. So long as time-dependent scattering processes that destroy the phase of the electron (processes such as electron-electron or electron-phonon scattering) are negligible, the interference of electrons manifests itself as aperiodic or periodic fluctuations of the conductance as a function of an external parameter such as gate voltage or magnetic field [13].

Furthermore, since scattering from each impurity or defect introduces a phase shift, the movement of the impurity by even a very small distance (equivalent to $1/k_F$, where k_F is the Fermi wave vector) will change the phase shift, resulting in a corresponding change in the conductance pattern [14]. Time-dependent conductance changes have been observed in disordered metals, and this mechanism has been shown to be a source of $1/f$ noise in metals at low temperatures [11].

The structure in the conductance of our samples shows a strong temperature dependence only in very specific ranges in applied bias (see fig. 2(b), for example). This rules out conductance fluctuations in the diffusive regime as the source of these fluctuations, as these would not be confined to specific ranges in applied bias. However, a similar interference mechanism can also exist in relatively clean carbon nanotubes with one or a few number of impurities or defects. In this case, the interference can be between electron waves that are directly transmitted, and those that are transmitted via a resonant state. This interference between directly transmitted channels and resonant channels gives rise to a Fano resonance, well known in atomic scattering [15]. For simplicity, we consider the case of a single directly transmitted channel and a single resonant state. Transmission through the resonant channel is described by an amplitude $t_r(\epsilon) = z_r\Gamma/(2(\epsilon - \epsilon_0) + i\Gamma)$ [16]. Without the factor z_r , this gives the usual expression for resonant transmission through a localized state of energy ϵ_0 (measured with respect to the Fermi energy ϵ_F) and intrinsic energy width Γ , $T_r = |t_r|^2 = \Gamma^2/(4(\epsilon - \epsilon_0)^2 + \Gamma^2)$, with a transmission of $T_r = 1$ on resonance ($\epsilon = \epsilon_0$). The transmission amplitude of the direct channel does not depend on ϵ , and can be described by an expression of the form $t_d = \sqrt{T_d}e^{i\alpha_d}$, where T_d is the transmission of the direct channel, and α_d describes the phase difference between the direct and resonant path. The interference between the two paths is taken into account by taking the sum of the transmission amplitudes to calculate the total transmission coefficient, $T_t = |t_r + t_d|^2$. The resulting conductance can be expressed in the Fano form

$$G(\epsilon) = \frac{2e^2}{h}T_t = \frac{2e^2}{h}T_d \frac{|2(\epsilon - \epsilon_0) + q\Gamma|^2}{4(\epsilon - \epsilon_0)^2 + \Gamma^2}, \quad (1)$$

where $q = i + z_r e^{-i\alpha_d}/\sqrt{T_d}$ is the complex Fano parameter [16]. Note that far from resonance ($\epsilon \gg \Gamma$), the conductance reduces to $G = (2e^2/h)T_d$.

A number of characteristics distinguish the resulting Fano resonance from other resonances that might occur in carbon nanotubes. First, depending on the phase difference between the resonant and non-resonant transmission channels, the Fano resonance can give rise to a peak or dip in the conductance (or something in between). Second, the Fano lineshape can be asymmetric about the transmission maximum or minimum. Finally, the position of the resonance is determined by the energy of the resonant state, and does not necessarily occur at the Fermi energy (zero bias). These distinguishing characteristics, which can be clearly seen in our data, rule out other possible mechanisms (such as the Kondo effect) for the structure we observe in the differential resistance of our devices.

A few groups have recently reported the observation of Fano resonances in carbon nanotube devices [17–19]. The Fano resonance appears as a structure in the differential conductance as a function of source-drain bias [17,18]. Kim *et al.* measured the conductance of crossed MWNTs, and observed a Fano resonance in two of the devices [17]. Noting that a Fano resonance was never observed in devices without crossed MWNTs, they associated the presence of the Fano resonance with the MWNT cross, although the mechanism by which a resonant state is created was not discussed. Yi *et al.* measured the conductance of crosses consisting of metal electrodes patterned across MWNT bundles [18]. In two of these devices, they observed non-monotonic behavior of the conductance near zero voltage bias, which they ascribed to a Fano

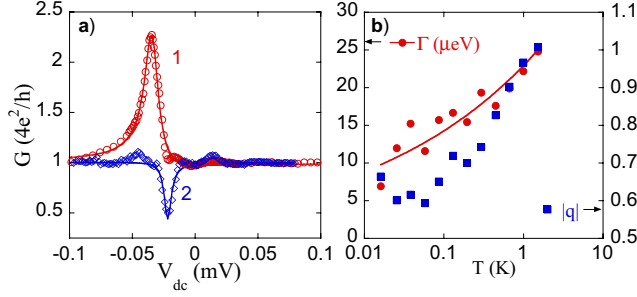


Fig. 4 – a) The Fano resonances in the conductance of Sample 3 from fig. 3(a) at $T = 20$ mK (curve 1), and fig. 3(b) at 16 mK (curve 2). The data (open circles) are normalized to $4e^2/h$, as discussed in the text. The solid lines are fits to the Fano function. b) The temperature dependence of the fitting parameters Γ and $|q|$ of the data from fig. 3(b), on a semilog plot. The solid line is a power law fit to $\Gamma = 22.9 \times T^{0.21} \mu\text{eV}$. The phase of q is nearly $\pi/2$ (1.58 ± 0.06).

resonance arising from interference between a Kondo resonance and non-resonant channels. However, the nature of the localized state giving rise to the Kondo resonance was not made clear. Furthermore, this interpretation is suspect in our opinion, because a Kondo resonance typically arises at the Fermi energy, and the resonances observed by Yi *et al.* were observed at voltage biases of 0.3–0.6 mV. Fano resonances have also been reported in SWNTs [19], where it was speculated that the interference may be between two transport channels in a single SWNT, or between transport channels on different nanotubes in a SWNT bundle. In all papers, the metastable behavior we observe was not reported.

Figure 4(a) shows the measured Fano resonances for the lower temperatures shown in figs. 3(a) and (b), along with fits to the Fano function given above. To obtain these curves, the background conductance outside the region ($V_{\text{dc}} = [-0.1 \text{ mV}, -0.02 \text{ mV}]$ for curve 1, and $V_{\text{dc}} = [-0.05 \text{ mV}, 0.02 \text{ mV}]$ for curve 2) has been fit using a Fourier filter, and subtracted from the experimental data. For the fits, the well-defined position of the resonances indicates that one resonant state participates in the interference. If we assume that only one direct channel participates in the interference, normalization of the data would give a conductance that drops below zero. Consequently, for the fits, we have assumed two direct channels (each with conductance $2e^2/h$) interfering with the resonant channel. These channels might correspond to the two channels of a SWNT, with the assumption that other channels that contribute to the background do not interfere with the resonant state. It can be seen that data are well described by the Fano equation. The change from a peak to a dip in conductance is a result of the annealing process, which presumably causes a change in the interference of the direct transmission states with the resonant state.

Figure 4(b) shows the fitted value of $|q|$ and Γ as a function of temperature for the conductance dip shown in fig. 4(a). Both $|q|$ and Γ increase with increasing temperature. Although the values we obtain for Γ are comparable to those obtained by Kim *et al.*, the temperature dependence is different. Kim *et al.* observed a linear temperature dependence, which they ascribed to thermal broadening of the linewidth, even though Γ was less than $k_B T$. Our temperature dependence is not linear, which is not surprising, since we are in the regime $\Gamma \leq k_B T$. Fitting to a power law gives a dependence $\Gamma \simeq T^{1/5}$, shown as the solid line in fig. 4(b); the origin of this power law is not clear to us. In contrast, while $|q|$ also increases with temperature at higher temperatures, it appears to saturate below 100 mK. The phase of $|q|$ remains essentially constant over the fitted temperature range.

What is the possible origin of the resonant states in our samples? Although we are not certain, the metastability we observe suggests that impurities or structural defects cause variations in the potential seen by the electrons in the nanotube, which in turn lead to localized electronic states with well-defined resonant energies. Movement of the impurities or defects can give rise to changes in the conductance through interference effects. Although the changes we observed in our samples typically occurred on the scale of days at low temperatures, we have also observed changes on the scale of hours. If there are many such impurities moving on a sufficiently rapid time scale, this may be one mechanism for the large $1/f$ noise in carbon nanotubes reported by some groups [20].

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