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Low temperature electron transport spectroscopy of mechanically templated carbon nanotube single electron transistors

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We report electronic transport investigations of mechanically templated carbon nanotube single electron transistors (SETs). The devices were fabricated on a Si/SiO₂ substrate by controllably placing individual single walled carbon nanotubes (SWNTs) between the source and drain electrodes via dielectrophoresis with a 100 nm wide local Al/Al₂O₃ bottom gate in the middle. From the low temperature electronic transport measurements, we show that a quantum dot is formed whose charging energy can be tuned from 10 to 90 meV by varying both the local gate and Si backgate. The temperature dependent measurements show that the Coulomb oscillations persist up to 250 K. The transport properties can be explained by a simple potential configuration, which suggests that two tunnel barriers are formed due to the bending of the SWNT at the local gate edges and that the size of the dot and tunnel barrier transparency can be tuned by the gates allowing the operation of SET in a wide temperature range and thereby realizing a controllable and tunable SET. Our simple fabrication technique and its tunability over a large temperature range could facilitate large scale fabrication of SET for practical applications. © 2013 AIP Publishing LLC.

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I. INTRODUCTION

Single electron transistors (SETs) have attracted considerable attention because of their potential for low powered quantum based nanoelectronic devices such as ultrasensitive sensors, memory and quantum information processing.^{1–4} A SET consists of a small conducting island connected to two metallic leads through tunnel barriers. Electron tunneling can be controlled one by one with a nearby gate electrode, capacitively coupled to the island when its charging energy (e^2/C) is greater than the thermal energy ($k_B T$). By reducing the size of the island, the capacitance decreases leading to a higher charging energy and operating temperature. For the practical realization of SET based devices, controllable and reproducible fabrication of SET is required. Since the first SET was demonstrated about 25 years ago in an aluminum tunnel junction,⁵ it has been realized in a variety of systems including lithographically defined dots in GaAs/AlGaAs heterojunction, direct etching of Si substrate, metallic grains in nanopore, colloidal nanocrystals, nanowires, and nanotubes.^{6–15} Lithography defined dots are often larger in size, requiring sub-Kelvin temperature for operation. In addition, complex fabrication processes make it difficult to control the uniformity and reproducibility. Metallic grains and colloidal nanocrystals give smaller and more uniform quantum dots (QDs) with operating temperature ~ 100 K. However, fabrication of such devices is very challenging due

to difficulty in placement in nanosized gaps, hence leading to extremely low device yield.

Recently, single walled carbon nanotubes (SWNTs)^{16–25} have been considered to be a good candidate for the fabrication of SETs because of their extraordinary electronic properties, extremely small diameter along with unique one dimensional nature. Controllable and reproducible fabrication of SET using SWNT relies on the introduction of tunnel barriers at desired positions along the nanotubes so that charges can be confined in a quantum dot defined by the tunnel barriers. Metal contact induced Schottky barrier has been used to define SWNT based SET.^{22–25} However, the fabrication of SET using this technique is a “hit or miss” and controllability has yet to be demonstrated. It has also been shown that when a SWNT is bent at a selected position, the bend acts as a nanometer sized tunnel barrier.^{19–21} By creating a pair of bends on an individual SWNT using atomic force microscopy (AFM) tip, SETs have been demonstrated.^{19,20} However, AFM manipulation is time consuming and reproducibility of the same sized device can be challenging.

In order to circumvent this and create more controllable and reproducible SETs, we have recently introduced a simple technique to bend SWNT by placing it on top of a local Al/Al₂O₃ gate between the source and drain contact.²⁶ The local Al/Al₂O₃ gate acts as a “mechanical template” and we hypothesized that SWNT bent at the edges of the local gate due to van der Waals interactions with the substrate to introduce two tunnel barriers.²⁷ As a result, a QD is defined by the width of the local gate. Inducing tunnel barriers to the SWNT using this scheme can be advantageous over other

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techniques because of its simplicity of design and compatibility with large scale fabrication.

In this paper, we investigate the mechanisms for the tunnel barrier formation, its tunability, the role of the local gate for controlling the tunnel barrier and the SET, and demonstrate operation at higher temperature. The device is fabricated on Si/SiO₂ substrate by controllably placing individual single walled nanotube (SWNT) between the source and drain electrodes via dielectrophoresis (DEP) with a 100 nm wide local Al/Al₂O₃ gate in the middle. We measured current oscillations at low temperature (down to 4 K) as a function of local gate voltage for different Si back-gate voltages. We show that the charging energy of the SET can be tuned from 10 to 90 meV by tuning both the gates. In addition, the temperature dependent measurements show that the single electron oscillations persist up to 250 K in qualitative agreement with the charging energy calculation. We also extract different device parameters from the slope of Coulomb diamond and explain the transport properties by a simple potential configuration, which suggests that two tunnel barriers are indeed formed due to the bending of the SWNT at the local gate edges and that the size of the dot and tunnel barrier transparency can be tuned by the gates allowing the operation of SET in a wide temperature range and thereby realizing a controllable and tunable SET. Our simple fabrication technique and its tunability over a large temperature range could facilitate large scale fabrication of SET for practical applications.

II. EXPERIMENTAL DETAILS

Figure 1(a) shows a schematic diagram of our device. The devices were fabricated on highly doped Si substrates with a thermally grown 250 nm thick SiO₂ capped layer. Optical lithography was used to define contact pads and position alignment markers using double layer resists (LOR3A/Shipley 1813), developed in CD26, followed by thermal evaporation of 5 nm chromium (Cr) and 45 nm gold and then standard lift-off. The source and drain electrodes with a

channel length of 1 μm were defined using standard electron beam lithography (EBL). After developing in (1:3) methyl isobutyl ketone:isopropyl alcohol (MIBK: IPA), 2 nm Cr and 25 nm thick Pd were deposited by electron beam evaporation followed by lift-off in acetone. We used tapered shaped electrodes as they maximize the electric field at the sharp edge, which increases individual SWNT assembly yield. A second EBL was then performed to define a 100 nm wide local Al gate electrode of 25 nm thickness centered between the source and drain electrodes. During the Al deposition, the sample stage was cooled with liquid nitrogen to minimize the surface roughness of the Al.²⁸ Finally, the sample was treated in oxygen plasma for 10 min to ensure a 2–3 nm thick aluminum oxide (Al₂O₃) layer.

The directed assembly of individual SWNT between the taper shaped electrode was done via DEP using a high quality aqueous solution of SWNT following our previous work.^{29–31} DEP has been used for the assembly of 2D, 1D, and 0D nanomaterials for device applications.^{32–39} DEP can be advantageous over other solution processed techniques because it allows the materials to be directly integrated to prefabricated electrodes. A stable, surfactant free solution with a concentration of SWNTs $\sim 50 \mu\text{g/ml}$ was obtained from Brewer Science.⁴⁰ The average diameter of the SWNTs in the solution was $\sim 1.7 \text{ nm}$ as determined by AFM.⁴¹ For the assembly, the concentration of SWNTs was diluted to $\sim 10 \text{ ng/ml}$ using DI water. A 3 μl drop of solution was cast onto the chip and an AC voltage of 5 V_{p-p} at 1 MHz was then applied between the source and drain electrodes. After 3 min, the AC voltage was turned off and the remaining solution was blown off from the chip with a stream of nitrogen gas. A Zeiss Ultra 55 field emission SEM was used to examine the results of the DEP assembly. Figure 1(b) shows a representative scanning electron micrograph (SEM) image of an individual SWNT assembled between a source and drain electrode pair and on top of the local gate. The devices were then wire bonded and loaded into a ⁴He cryostat for electronic transport measurements. DC charge transport measurements were performed by means of a DL instruments 1211 current preamplifier and a Keithley 2400 source-meter interfaced with LabView.

III. RESULTS AND DISCUSSIONS

Figure 2(a) shows a plot of room temperature drain current as a function of local gate (V_{LG}) and back gate voltages (V_{BG}) for one of our representative DEP assembled device. The highly doped Si was used as a back gate. The device shows a small change in current as a function of gate voltage indicating a small band gap SWNT.⁴² The detailed room temperature property of our DEP assembled devices can be found in our previous publications.^{29–32} In this paper, we only focus on low temperature SET behavior. It should be noted that the local gate shows more efficient gate coupling compared to the back gate due to thinner gate oxide with higher dielectric constant compared to the SiO₂.³¹

Figure 2(b) shows current versus V_{LG} for a fixed source drain voltage of $V_{\text{DS}} = 5 \text{ mV}$ measured at a temperature (T) of 4.2 K and $V_{\text{BG}} = 0 \text{ V}$. Current oscillations are observed as

FIG. 1. (a) Schematic diagram of the mechanical template based SWNT SET device. (b) SEM of a fabricated device. An individual SWNT was placed over the local gate and between the source and drain electrodes by dielectrophoresis. The width of the local gate electrode is $\sim 100 \text{ nm}$. Reprinted with permission from P. Stokes and S. I. Khondaker, Appl. Phys. Lett. **92**, 262107 (2008). Copyright 2008 American Institute of Physics.

FIG. 2. (a) Current (I) versus local gate voltage (V_{LG}) and back gate voltage (V_{BG}) of a representative device measured at room temperature. $V_{DS} = 0.1\text{ V}$ for both curves. (b) I versus V_{LG} measured at $T = 4.2\text{ K}$, $V_{BG} = 0\text{ V}$ and $V_{DS} = 5\text{ mV}$. (c) I versus V_{BG} measured at $T = 4.2\text{ K}$, $V_{LG} = 0\text{ V}$ and $V_{DS} = 5\text{ mV}$. Current oscillations can be seen in (b) and (c), typical of a SET behavior. (d) Color scale plot of I versus V_{BG} and V_{LG} taken at $T = 4.2\text{ K}$ with a constant source-drain voltage of 5 mV . Bright regimes symbolize high current and dark regimes symbolize low current. Diagonal lines running throughout the plot indicating that the conductance of the device can be tuned by using both the local gate and the back gate.

a function of gate voltage, typical of SET behavior.⁴³ Figure 2(c) shows a plot of current versus V_{BG} with $V_{LG} = 0\text{ V}$, and $V_{DS} = 5\text{ mV}$ at $T = 4.2\text{ K}$, showing similar SET behavior. The observation of SET behavior suggests that a quantum dot has formed in the channel and that the SET can be independently controlled by both the back gate and the local gate.

Figure 2(d) displays a color scale plot of current (I) as a function of both V_{BG} and V_{LG} measured at 4.2 K at a fixed bias voltage ($V_{DS} = 5\text{ mV}$). The current was measured by changing V_{BG} from -10 to 10 V at a step of 20 mV , while sweeping the local gate from -2.5 to 2.5 V at every V_{BG} . The darker regions signify Coulomb Blockade, while other bright colors running diagonal to the figure signify peak of current oscillations. The alternating diagonal bright and dark lines demonstrate that, in addition to independent control of the SET by the gates, their combined effect can also control the operating regime of the SET. For example, for a V_{BG} of -7.5 V , the current oscillations persist for a V_{LG} of -2.5 to 2.5 V , while for a V_{BG} of $+5\text{ V}$, there are no current oscillations for any value of V_{LG} . In order to more clearly understand this combined effect of the back gate and local gate on

our device, we present detailed analysis of our data at two constant back gate voltage regimes: (a) $V_{BG} = -7\text{ V}$ where the conductance is high, and (b) $V_{BG} = 1.5\text{ V}$ where the conductance begins to die off. Our analysis is based on color scale plots of the differential conductance (dI/dV_{DS}), as a function of V_{DS} and V_{LG} at 4.2 K (often called a *stability plot*) to determine behavior of the device in these regimes. The stability plots allow us to determine the charging energy of the dot, various capacitances associated with the dot, as well as size of the dots at those regimes.

Figure 3(a) shows the stability plot with $V_{BG} = -7\text{ V}$. The V_{DS} was varied from -50 to $+50\text{ mV}$ with a step of 0.5 mV , while the V_{LG} was swept from -1.5 to $+1.5\text{ V}$. Brighter regions (red and white) symbolize high conductance and darker regions (blue) signify Coulomb blockade. Throughout the plot, a series of diamond shaped regions called Coulomb diamonds are observed. Closing of the diamonds indicates measurement of a single quantum dot.⁴³ However, the height of the diamonds and their slopes changes with changing V_{LG} . In order to investigate this in more detail, we focus on two local gate regions: (I) $-1.25 < V_{LG} < -0.75\text{ V}$ and (II) $0.5 < V_{LG} < 1.5\text{ V}$. These

FIG. 3. (a) dI/dV_{DS} versus V_{DS} and V_{LG} taken at 4.2 K with $V_{BG} = -7\text{ V}$. Red regions symbolize high conductance and blue regions symbolize low conductance or Coulomb blockade. (b) Expanded view of the stability plot in region I showing transport through a single QD with charging energy $8-12\text{ meV}$. (c) Expanded view of stability plot in region II showing single QD behavior with charging energy $\sim 20-45\text{ meV}$.

FIG. 4. dI/dV_{DS} versus V_{DS} and V_{LG} taken at 4.2 K with $V_{BG} = 1.5$ V.

regions are expanded in Figures 3(b) and 3(c), respectively. The height of the diamond is a direct measure of charging energy (U_C) of the quantum dot. The dashed lines in Fig. 3(b) are placed for a guide to the eye to outline a diamond from which we obtain $U_C \sim 10$ meV. The capacitance values of the SET can be calculated from the Coulomb diamond using constant interaction model.⁴³ The gate capacitance is calculated from the period ($\Delta V_G^I \sim 50$ mV) between the diamonds using the formula $C_G^I = e/\Delta V_G^I$, which yields $C_G^I \sim 3$ aF. The source and drain capacitances are calculated from the positive slope $\alpha_1 = -C_1^I/C_1^I$ and negative slope $\alpha_2 = C_2^I/(C_1^I + C_2^I)$ of the diamond, respectively (see Fig. 3(b)), which yield $C_1^I \sim 12$ aF and $C_2^I \sim 1.5$ aF. The total capacitance of the SET is $C_\Sigma^I = C_1^I + C_2^I + C_G^I = 16.5$ aF, which can be used to calculate the charging energy of $U_C^I = e^2/C_\Sigma^I \sim 10$ meV in good agreement with the value measured directly from the height of the diamond in this regime. It has been shown that the charging energy of a SWNT quantum dot can be estimated from the formula, $U_C = 5 \text{ eV}/L \text{ (nm)}$, where L is the length the SWNT quantum dot in nm.⁴⁴ Using the value of $U_C = 10$ meV, we calculate a QD of size ~ 500 nm. Figure 3(c) shows an expanded view of the stability plot for $0.5 < V_{LG} < 1.5$ V (region II). Here, the period is ($\Delta V_G^{II} \sim 70$ mV) giving a gate capacitance $C_G^{II} \sim 2.3$ aF. The source and drain capacitances are $C_1^{II} \sim 2.1$ aF and $C_2^{II} \sim 0.35$ aF, respectively, giving a total capacitance of $C_\Sigma^{II} = C_1^{II} + C_2^{II} + C_G^{II} \sim 4.8$ aF and a charging energy of $U_C^{II} = e^2/C_\Sigma^{II} \sim 34$ meV, consistent with the height of the diamond in this regime. From here, we obtain a QD size of ~ 147 nm.

Figure 4 shows the stability plot of the same device at $V_{BG} = 1.5$ V. The charging energy in this case is substantially higher compared to $V_{BG} = -7.0$ V, which is shown in Figure 3. The color scale plot exhibits a large diamond of spacings $\Delta V_G \sim 300\text{--}400$ meV inside which there is a periodic ($\Delta V_G \sim 50$ meV) saw tooth structure. This is a signature of charge transport through a small quantum dot in series with a large quantum dot; however, the smaller QD dominates the transport.^{45,46} The charging energy read off from the diamond is ~ 40 meV for region I ($-2.2 < V_{LG} < -0.75$ V) and is

~ 90 meV in region II ($-0.75 < V_{LG} < 1.5$ V). This corresponds to a QD size of 125 nm in region I and 55 nm in region II, respectively. The capacitances associated with the QD are calculated from the slope of the diamond and is presented in Table I, along with the capacitance, charging energy, and the corresponding size of QD for the regimes shown in Figures 3 and 4. It can be seen that the charging energy increased from 10 to 90 eV as the total capacitance decreased from 16.5 to 1.8 aF under different V_{BG} and V_{LG} conditions demonstrating that the SET is highly tunable in charging energy.

Figure 5 shows the drain current of the SET versus the local gate voltage at different temperatures for fixed value of $V_{BG} = -7$ V (Figure 5(a)) and $V_{BG} = 1.5$ V (Figure 5(b)). The bias voltage was kept fixed at 5 mV. For $V_{BG} = -7$ V, the current oscillations persist up to 120 K, whereas for $V_{BG} = 1.5$ V the oscillations persist up to 250 K. This shows that with increasing V_{BG} , the operating temperature increases, consistent with Figures 3 and 4, where we observed a larger charging energy with increasing V_{BG} . We can also estimate the charging energy of the quantum dot from the temperature dependent data using the formula, $U_C = 4K_B T$.^{47,48} For $T = 250$ K, $U_C = 86$ meV which is close to the charging energy obtained from the stability plot.

The most important observation in our study is that the charging energy and operating temperature can be increased with increasing V_{BG} and V_{LG} . From here, one can conclude that the system is highly tunable in charging energy and temperature. We have measured three other devices and found that this observation is typical for all of our devices. This is due to modification of tunnel barriers with gate voltage which in effect defines how the QDs are localized on the SWNT, which is evident in Table I. In order to understand this, we put forward a generalized potential configuration on

TABLE I. Summary of capacitance and charging energy values for the device at different local gate regions for positive and negative back gate voltages.

V_{BG} (V)	Region	C_G (aF)	C_1 (aF)	C_2 (aF)	C_Σ (aF)	E_C (meV)	L (nm)
-7	I	3	12	1.5	16.5	10	500
-7	II	2.3	2.1	0.35	4.75	34	147
1.5	I	0.6	2.6	0.9	4.1	40	125
1.5	II	0.4	1.3	0.06	1.8	90	55

FIG. 5. Dependence of the drain current on local gate voltage at two different back gate voltages using $V_{DS} = 5$ mV. (a) Temperature dependent I versus V_{LG} at a fixed $V_{BG} = -7$ V. $T = 5, 15, 30, 50, 80$, and 120 from bottom to top. (b) Temperature dependent I versus V_{LG} at a fixed $V_{BG} = 1.5$ V. $T = 15, 30, 50, 80, 120, 200$, and 250 K (bottom to top).

FIG. 6. Potential profile for the quantum dot formation at different local gate and back gate voltage regimes.

how the tunnel barriers are formed by the mechanical templates and modified by the local and backgate voltages, shown in Figure 6.

We first discuss the case for $V_{BG} \ll 0$ V, $V_{LG} \ll 0$ V (Figure 6(a)). In this case, the tunnel barriers defined by the local gate voltage are transparent and the QD is formed by the contact. This is consistent with the fact that the charging energies are small leading to a large QD. However, for $V_{BG} \ll 0$ V and $V_{LG} \gg 0$ V (Figure 6(b)), the tunnel barriers caused by the local gate become opaque and all other potential fluctuations are placed below the Fermi level. As a result, the QD is defined by the local gate only as the QD size in this regime is close to the physical width of the local gate. Now we discuss the case where $V_{BG} \gg 0$ V (right column of Figure 6). In this regime, the tunnel barriers defined by the local gate are opaque and the QD that dominates the transport is defined by the local gate. The observed beating may be due to another quantum dot formed by the charge fluctuations caused by the substrate. This leads to the formation of two QD, the small QD is defined by the local gate edge and the larger one by charge fluctuations.⁴⁹ The increased charging energy with increasing V_{LG} suggests that the tunnel barriers defined by the local gate gotten wider, further reducing the size of the dominant QD.

IV. CONCLUSION

In conclusion, we studied the electronic transport properties of mechanically templated carbon nanotube SETs. The charging energy of the SET can be varied from 10 to 90 meV by tuning the local gate and back gate voltages, which allowed the operating temperature to be up to 250 K. The transport properties can be explained by a simple potential configuration, which suggests that the bending of the SWNT at the local gate edges creates two tunnel barriers and that

the size of the dot and tunnel barrier transparency can be tuned by the gates allowing the operation of SET in a wide temperature range and thereby realizing a controllable and tunable SET. This simple fabrication technique along with tunability over a large temperature range could facilitate large scale fabrication of SET for practical applications.

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¹M. H. Devoret and C. Glattli, *Phys. World* **11**, 29 (1998).

²K. K. Likharev, *Proc. IEEE* **87**, 606 (1999).

³R. H. Chen, A. N. Korotkov, and K. K. Likharev, *Appl. Phys. Lett.* **68**, 1954 (1996).

⁴R. J. Schoelkopf, P. Wahlgren, A. A. Kozhevnikov, P. Delsing, and D. E. Prober, *Science* **280**, 1238 (1998).

⁵T. A. Fulton and G. D. Dolan, *Phys. Rev. Lett.* **59**, 109 (1987).

⁶J. A. Folk, S. R. Patel, S. F. Godijn, A. G. Huibers, S. M. Cronenwett, C. M. Marcus, K. Campman, and A. C. Gossard, *Phys. Rev. Lett.* **76**, 1699 (1996).

⁷Y. Takahashi, H. Namatsu, K. Kurihara, K. Iwdate, M. Nagase, and K. Murase, *IEEE Trans. Electron Devices* **43**, 1213 (1996).

⁸K. I. Bolotin, F. Kuemmeth, A. N. Pasupathy, and D. C. Ralph, *Appl. Phys. Lett.* **84**, 3154 (2004).

⁹D. Klein, R. Roth, A. K. L. Lim, A. P. Alivisatos, and P. L. McEuen, *Nature (London)* **389**, 699 (1997).

¹⁰S. I. Khondaker, K. Luo, and Z. Yao, *Nanotechnology* **21**, 095204 (2010).

¹¹T. Mori, S. Sato, K. Omura, S. Yajima, Y. Tsuruoka, Y. Achiba, and K. Ishibashi, *J. Vac. Sci. Technol. B* **27**, 795 (2009).

¹²K. Maehashi, H. Ozaki, Y. Ohno, K. Inoue, K. Matsumoto, S. Seki, and S. Tagawa, *Appl. Phys. Lett.* **90**, 023103 (2007).

¹³M. Suzuki, K. Ishibashi, K. Toratani, D. Tsuya, and Y. Aoyagi, *Appl. Phys. Lett.* **81**, 2273 (2002).

¹⁴M. T. Bjork, C. Thelander, A. E. Hansen, L. E. Jensen, M. W. Larsson, L. R. Wallenberg, and L. Samuelson, *Nano Lett.* **4**, 1621 (2004).

¹⁵Z. Zhong, Y. Fang, W. Lu, and C. M. Lieber, *Nano Lett.* **5**, 1143 (2005).

¹⁶M. Bockrath, D. H. Cobden, P. L. McEuen, N. G. Chopra, A. Zettl, A. Thess, and R. E. Smalley, *Science* **275**, 1922 (1997).

- ¹⁷S. J. Tans, M. H. Devoret, H. Dai, A. Thess, R. E. Smalley, B. L. Geeligs, and C. Dekker, *Nature (London)* **386**, 474 (1997).
- ¹⁸M. Bockrath, D. H. Cobden, J. Lu, A. G. Rinzler, R. E. Smalley, L. Balents, and P. L. McEuen, *Nature* **397**, 598 (1999).
- ¹⁹H. W. C. Postma, T. Teepen, Z. Yao, M. Grifoni, and C. Dekker, *Science* **293**, 76 (2001).
- ²⁰D. Bozovic, M. Bockrath, J. H. Hafner, C. M. Lieber, H. Park, and M. Tinkham, *Appl. Phys. Lett.* **78**, 3693 (2001).
- ²¹M. J. Biercuk, N. Mason, J. M. Chow, and C. M. Marcus, *Nano Lett.* **4**, 2499 (2004).
- ²²P. Jarillo-Herrero, S. Sapmaz, C. Dekker, L. P. Kouwenhoven, and H. S. J. van der Zant, *Nature* **429**, 389 (2004).
- ²³K. Ishibashi, M. Suzuki, T. Ida, and Y. Aoyagi, *Appl. Phys. Lett.* **79**, 1864 (2001).
- ²⁴H. Li, Q. Zhang, and J. Li, *Appl. Phys. Lett.* **88**, 013508 (2006).
- ²⁵P. Stokes and S. I. Khondaker, *ACS Nano* **4**, 2659 (2010).
- ²⁶P. Stokes and S. I. Khondaker, *Appl. Phys. Lett.* **92**, 262107 (2008).
- ²⁷T. Hertel, R. E. Walkup, and P. Avouris, *Phys. Rev. B* **58**, 13870 (1998).
- ²⁸W. J. Liang, M. P. Shores, M. Bockrath, J. R. Long, and H. Park, *Nature* **417**, 725 (2002).
- ²⁹P. Stokes and S. I. Khondaker, *Appl. Phys. Lett.* **96**, 083110 (2010).
- ³⁰K. J. Kormondy, P. Stokes, and S. I. Khondaker, *Nanotechnology* **22**, 415201 (2011).
- ³¹M. R. Islam, K. J. Kormondy, E. Silbar, and S. I. Khondaker, *Nanotechnology* **23**, 125201 (2012).
- ³²S. I. Khondaker, *IEEE Proc.: Circuits Devices Syst.* **151**, 457 (2004).
- ³³P. Stokes and S. I. Khondaker, *Nanotechnology* **19**, 175202 (2008).
- ³⁴P. Stokes, E. Silbar, Y. M. Zayas, and S. I. Khondaker, *Appl. Phys. Lett.* **94**, 113104 (2009).
- ³⁵D. Joung, A. Chunder, L. Zhai, and S. I. Khondaker, *Nanotechnology* **21**, 165202 (2010).
- ³⁶M. R. Islam, D. Joung, and S. I. Khondaker, *New J. Phys.* **13**, 035021 (2011).
- ³⁷D. Joung, L. Zhai, and S. I. Khondaker, *Phys. Rev. B* **83**, 115323 (2011).
- ³⁸B. K. Sarker, S. Shekhar, and S. I. Khondaker, *ACS Nano* **5**, 6297 (2011).
- ³⁹B. K. Sarker, M. R. Islam, F. Alzubi, and S. I. Khondaker, *Mater. Express* **1**, 80 (2011).
- ⁴⁰See <http://www.brewerscience.com> for the physical properties of the SWNT solution.
- ⁴¹S. Shekhar, P. Stokes, and S. I. Khondaker, *ACS Nano* **5**, 1739 (2011).
- ⁴²C. Zhou, J. Kong, and H. Dai, *Phys. Rev. Lett.* **84**, 5604 (2000).
- ⁴³R. Hanson, L. P. Kouwenhoven, J. R. Petta, S. Tarucha, and L. M. K. Vandersypen, *Rev. Mod. Phys.* **79**, 1217 (2007).
- ⁴⁴J. Nygard, D. H. Cobden, M. Bockrath, P. L. McEuen, and P. E. Lindelof, *Appl. Phys. A: Mater. Sci. Process.* **69**, 297 (1999).
- ⁴⁵J. Park and P. L. McEuen, *Appl. Phys. Lett.* **79**, 1363 (2001).
- ⁴⁶H. Li and Q. Zhang, *Appl. Phys. Lett.* **94**, 022101 (2009).
- ⁴⁷J. Kong, C. Zhou, E. Yenilmez, and H. Dai, *Appl. Phys. Lett.* **77**, 3977 (2000).
- ⁴⁸P. L. McEuen, L. P. Kouwenhoven, S. Jauhar, J. Orenstein, K. McCormick, D. Dixon, Y. U. Nazarov, N. C. V. D. Vaart, and C. T. Foxon, *Nanotechnology* **7**, 406 (1996).
- ⁴⁹M. Bockrath, W. Liang, D. Bozovic, J. H. Hafner, C. M. Lieber, M. Tinkham, and H. Park, *Science* **291**, 283 (2001).