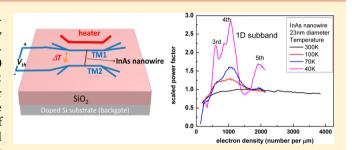


One-Dimensional Quantum Confinement Effect Modulated Thermoelectric Properties in InAs Nanowires

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Supporting Information

ABSTRACT: We report electrical conductance and thermopower measurements on InAs nanowires synthesized by chemical vapor deposition. Gate modulation of the thermopower of individual InAs nanowires with a diameter around 20 nm is obtained over T = 40-300 K. At low temperatures (T <~100 K), oscillations in the thermopower and power factor concomitant with the stepwise conductance increases are observed as the gate voltage shifts the chemical potential of electrons in InAs nanowire through quasi-one-dimensional (1D) subbands. This work experimentally shows the possibility



to modulate semiconductor nanowire's thermoelectric properties through 1D subband formation in the diffusive transport regime for electron, a long-sought goal in nanostructured thermoelectrics research. Moreover, we point out the scattering (or disorder) induced energy level broadening as the limiting factor in smearing out the 1D confinement enhanced thermoelectric

KEYWORDS: Thermopower, one-dimensional, quantum confinement, nanowire, indium arsenide

In the past five decades, much attention has been paid to thermoelectric materials due to their device applications in converting heat to electricity or vice versa. This conversion efficiency is described by one fundamental parameter, the thermoelectric figure of merit, which is defined as $S^2 \sigma T/k$, where S is the Seebeck coefficient or thermopower, σ is the electrical conductivity, T is the absolute temperature, and k is the thermal conductivity. The difficulty in obtaining better ZT values arises from the fact that S, σ , and k are related to each other. Changing one alters the others simultaneously, which makes optimization extremely difficult. Since the first thermoelectric Bi₂Te₃ alloy has been found, the room-temperature ZT of bulk semiconductors has increased only marginally, from about 0.6 to 1.1,2 However, recent studies in nanostructured thermoelectric materials have led to significant advancements, 2,3 such as Bi₂Te₃/Sb₂Te₃ thin film superlattices, 4 PbSeTe/PbTe quantum dot superlattices,⁵ and nanocrystalline BiSbTe bulk alloys⁶ with ZT > 1. There are two basic mechanisms by which ZT is improved in low-dimensional systems: reduced lattice component of thermal conductivity with size effect, and increased power factor $(S^2\sigma)$ by quantum confinement enhanced electronic density of states (DOS) in quasi-1D quantum wires/nanowires or quasi-zero-dimensional (0D) quantum dots or nanoparticles. Most experimental works on the thermoelectric properties of nanostructures have found

greatly reduced thermal conductivity in nanowires or nanoparticle composite materials.^{6–9} However, the full potential of nanomaterials as high figure of merit thermoelectrics can only be achieved when the sharp (or singular) electronic DOS spectra in quantum confined low-dimensional (1D or 0D) systems are exploited, as first calculated by Dresselhaus in 1993 and elaborated further by others using Boltzmann transport formalism for diffusive electrons. ^{10–14} In this regard, combining the peaked 1D electronic DOS and drastically reduced thermal conductivity in semiconductor nanowires appears to be a promising route toward obtaining thermoelectrics with high figure of merit. Unfortunately, despite much progress in the study of thermoelectric properties of nanomaterials, 7-9,15-17 there is no experimental work showing that the sharply peaked DOS in 1D or 0D materials can indeed be used to tailor the thermoelectric properties. In carbon nanotubes (CNTs) where the 1D electron confinement is strong, the thermopower was found to oscillate due to defects induced resonant electron scattering or the Coulomb blockade effect. 18,19 In addition, due to the extremely high thermal conductivity of CNTs, it is not clear if CNTs can work as high ZT thermoelectric materials. In PbSe nanowires, the electrical gating method was used to

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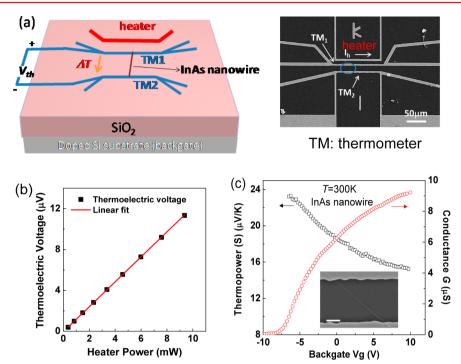


Figure 1. InAs nanowire thermoelectric transport device layout and the room temperature behavior. (a) A schematic and SEM image of nanowire thermoelectric device. Two line electrodes, TM_1 and TM_2 , in contact with a single InAs nanowire, are used as thermometers to calibrate the temperature difference ΔT caused by the heating current I_h . (b) The thermoelectric voltage (V_{th}) across the nanowire is proportional to the heating power (P) due to the proportion of P to ΔT . (c) The gate dependence of InAs nanowire's conductance (red circles) and thermopower (black squares). The inset is a zoomed-in SEM image of the blue circle area in the SEM image of a, showing two thermometer line electrodes connected by a single InAs nanowire. Scale bar: 2 μ m.

modulate the Seebeck coefficient.²⁰ However, the generic behavior of the gate tuned Seebeck coefficient is consistent with bulk behavior, and no sign of sharply peaked 1D DOS was observed. The underlying reason that it has been difficult to achieve the 1D electronic confinement effect enhanced thermoelectric properties in semiconductor nanowires is tied to the stringent experimental requirements: first, nanowires are typically fairly large (10–100 nm) in diameter such that low carrier concentration and low temperatures are generally required to reach the 1D regime for electron transport; second, even in the low carrier concentration/temperature regime, high-quality material is required to avoid a structural defect induced Coulomb blockade which can reduce the conductivity and dominate 1D electrical transport.

Here we report the thermoelectric properties investigated on single indium arsenide (InAs) nanowires. Thanks to the small effective electron mass ($m^* = 0.023m_e$, with m_e as the free electron mass), 1D sub-band splitting is large in InAs nanowires, making them a good candidate to study the 1D confinement effects on the thermoelectric properties at relatively high temperatures (e.g., above liquid nitrogen temperature and possibly even at room temperature). In a recent electrical transport study of InAs nanowire, the effect of 1D subband quantization was clearly seen as steps in conductance vs gate voltage at moderate temperatures (100 K) for nanowires with diameter 10-40 nm in the diffusive transport regime.²¹ Moreover, there is significant interest in the quantum transport effect^{22,23} and nanoelectronic device performance^{24–29} of InAs nanowires. In this work, we demonstrate gate modulated thermopower together with the electrical conductance in InAs nanowires as the Fermi level sweeps through 1D electron subbands. Significantly, below

temperature ~100 K, thermopower exhibits oscillations concomitant with the stepwise conductance increase, as a result of the Fermi energy being tuned through the DOS peaks of 1D subbands. We further find that the gate modulated power factor oscillations correlate well with the corresponding behavior of 1D DOS spectrum. However, together with the thermal smearing of 1D energy levels, the scattering induced level broadening is found to severely suppress/smear out the oscillations in 1D diffusive thermoelectric transport properties, pointing to the imperative need for reduced carrier scattering to achieve quantum confinement enhanced thermoelectric figure of merit at practical temperatures. These findings provide the first evidence of 1D confinement induced thermopower modulation in semiconductor nanowires and serve to provide better insights into tailoring nanomaterials' thermoelectric properties.

InAs nanowires were grown by the chemical vapor deposition (CVD) method on silicon (100) substrates in 10% H₂/Ar mixture gas at 660 °C with 20 nm Au nanoparticles (Ted Pella, Inc.) as catalyst.³⁰ A Lindberg Blue M tube furnace and 1inch diameter quartz tube were used for vapor transport synthesis of nanowires with accurate control of temperature and carrier gas flow rate. The 99% InAs powder (Alfa Aesar) was ground as precursor and placed at the center of the furnace. The substrates were treated with poly L-lysine (Sigma) for 8 min 30 s and covered with 20 nm Au catalyst for 5 min successively. The pretreated substrates were then placed at a distance of 13.5 cm from the source. Detailed growth conditions of InAs nanowires are as follows. At first, maximum flow rate of 100 sccm (standard cubic centimeters per minute) H₂/Ar carrier gas was introduced after the system was pumped down to the base pressure of about 8 mTorr. When the

temperature of the furnace was increased to 660 $^{\circ}$ C, the flow rate was decreased to 4 sccm. The pressure was kept at 1.5 Torr for 5 min and then changed to 100 mTorr for 3 h. After that the furnace was cooled naturally down to room temperature, and dark gray products composed of InAs nanowires were found on the substrates. These InAs nanowires were found to have zincblende crystal structure with $\langle 111 \rangle$ growth direction.³⁰

As-grown nanowires were suspended in ethanol by ultrasound sonication and then dropped onto a Si substrate with 600 nm thick oxide. The Si substrate was degenerately doped and used as a backgate electrode. Photolithography and thermal evaporation of 80 nm Ni were used to establish the electrodepatterns (Figure 1a) designed for thermoelectric experiment. The sample was dipped in 0.5% hydrofluoric acid (HF) solution for 5 s before evaporation to remove any native oxide layer and ensure Ohmic contacts. Low-frequency lock-in techniques were used to measure the conductance and thermopower in a Quantum Design PPMS cryostat. In the thermopower experiment, a sinusoidal current at frequency ω was applied on the heater to generate a temperature gradient ΔT at frequency of 2ω . The resulting 2ω thermoelectric voltage $V_{\rm th}$ along the nanowire was then detected by lock-in (SR830, Stanford Research). The temperature gradient ΔT was calibrated by using the four-wire resistances of the two electrode lines as thermometer #1 (TM1) and thermometer #1 (TM2) (Supporting Information). For the conductance measurement, a source-drain voltage of 100 µV was applied between the electrodes (TM1 and TM2) at the ends of nanowire, and the current was measured by a lockin (SR830). Multiple devices of InAs NWs with diameter about 25 nm were measured, and the manuscript presents data from the sample #1 on which the most comprehensive data were collected (thermoelectric data on additional devices can be found in Figures S4–7 in Supporting Information).

Figure 1a shows a schematic and scanning electron microscope (SEM) micrograph of a typical device used in our study of thermoelectric properties. The line electrodes on top of InAs nanowire (zoomed in as the inset to Figure 1c) serve also as thermometers. Another Ni electrode placed nearby, but not in contact with, the InAs nanowire serves as a line heater. A current I_h applied to the heater causes a temperature gradient via Joule heating. The heating power (*P*) is calculated as $P = V_h^2/R_h$, in which $V_h = I_h R_h$ is the heater voltage and R_h is the heater resistance. The temperature difference, ΔT , between the two thermometers is proportional to the heating power, which leads to a linear relation of thermoelectric voltage to P (Figure 1b). The thermopower $S \equiv$ $(V_{\rm th}/\Delta T)$ was determined by the measured $V_{\rm th}$ and the ΔT inferred from the resistance vs P dependence of the two thermometers using their temperature-dependent resistance (measured at equilibrium, $P \sim 0$) as thermometry (Figures S1 and S2, Supporting Information). S is expected to be negative for n-type semiconductors, as we have confirmed for our n-type InAs nanowire. For the convenience of presentation, we show the absolute value of S throughout this paper.

The room temperature ($T=300~{\rm K}$) gate voltage ($V_{\rm g}$) dependence of the conductance G and the thermopower S of the InAs nanowire are shown in Figure 1c. The $V_{\rm g}$ dependence of G is typical for a depletion mode n-type field effect transistor (FET). Using the diameter d=23 nm and length of 9.5 μ m measured by SEM, an estimated peak field effect mobility about $2060~{\rm cm^2/(V~s)}$ is extracted by the trans-conductance value. In Figure 1c, it is evident that, as the InAs nanowire is gated

toward depletion, the thermopower increases monotonically. (Similar behavior was observed in devices #2 and #3, as shown in Figures S4 and S6 in the Supporting Information). This behavior of increasing thermopower with decreased conductance or carrier density tuned by gate is consistent with the typical behavior of semiconductors^{20,31} and can be understood using Mott's formula for degenerate semiconductors³²

$$S = -\frac{\pi^2}{3} \frac{k_{\rm B}}{e} k_{\rm B} T \times \frac{\mathrm{d} \ln(\sigma)}{\mathrm{d}E} \bigg|_{E=E_{\rm F}}$$

$$= -\frac{\pi^2}{3} \frac{k_{\rm B}}{e} k_{\rm B} T \times \frac{1}{G} \frac{\mathrm{d}G}{\mathrm{d}V_{\rm g}} \frac{\mathrm{d}V_{\rm g}}{\mathrm{d}E} \bigg|_{E=E_{\rm F}}$$
(1)

where $E_{\rm F}$ is the Fermi energy, $k_{\rm B}$ is the Boltzmann constant, and e is electron charge. In eq 1, $({\rm d}G/{\rm d}V_{\rm g})({\rm d}V_{\rm g}/{\rm d}E_{\rm F})$ is positive and a monotonic function of $V_{\rm g}$ (with increased $V_{\rm g}$, the electron density increases so that both the Fermi energy and conductance increases). In such cases, the absolute value of thermopower S is expected to increase monotonically as the gate tunes the nanowire toward the low carrier density/conductance regime. It should be noted that the measured thermopower values S contain contributions from both the absolute values of the InAs nanowires $S_{\rm InAs}$ and that of the Ni electrode $S_{\rm Ni}$, that is, $S = S_{\rm Ni} - S_{\rm InAs}$. But $S_{\rm Ni}$ is expected to be insensitive to $V_{\rm g}$ due to Ni being a good metal. Therefore the variation of S can be purely attributed to that of InAs nanowire.

The S vs V_g in Figure 1c is in agreement with the general trend of bulk semiconductor's thermopower as a function of electron density (the electron density is proportional to the $V_{
m g}$ in our case). 20,31 This is because the thermal broadening $(k_{\rm B}T)$ and scattering induced level broadening ($\delta E \sim h/\tau$) at 300 K are too large to resolve the separation between quantized 1D subbands, where $k_{\rm B}$ is the Boltmann constant, h is the Planck's constant, and τ is scattering time of electrons. When k_BT and δE are smaller than the energy separation between 1D subbands, the 1D subband quantization and the oscillating 1D DOS should play a dominant role in the thermoelectric transport properties as we demonstrate below. The conductance of the InAs nanowire as a function of the gate voltage $V_{\rm g}$ is shown in Figure 2a for temperatures from 40 to 300 K. As can be seen, when T decreases from 300 K, the overall $G(V_g)$ curve shifts toward right, and the trans-conductance at the low G (electron density) increases, reflecting the carrier freeze-out and increased electron mobility effects when semiconductors are cooled. What is more interesting is that, below 100 K, the curve develops some step-like features. These stepwise conductance increases are due to the population of 1D subbands and the Fermi level of the system lies between two adjacent 1D subbands at these steps.^{21,33} Figure 2b presents a comparison between our experimental $G(V_{\sigma})$ data at 40 K and numerically calculated curves for 1D subband occupation number $N_{\rm 1Dsubbands}$ vs $V_{\rm g}$ following ref 21.³⁴ The 2-fold degeneracy for subbands index higher than 1 was also included in the calculated $N_{\mathrm{1Dsubbands}}$. We found that, without scattering broadening, the thermal broadening itself produces steps (rises) flatter (sharper) than experimental data, consistent with ref 21. A close resemblance can be seen between the data and the theoretical curve (solid line) with both the thermal and the scattering broadening are considered, corroborating the 1D subband filling interpretation of conductance plateaus. Note

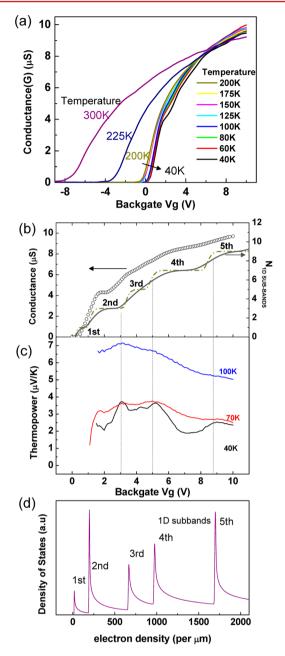


Figure 2. Gate-tuned conductance and thermopower of InAs nanowire (diameter $d=23\,$ nm) at different temperatures. (a) Temperature effect in the gate voltage-dependent conductance G from T=40 to 300 K. Steps due to the electron population of individual 1D subbands as a function of $V_{\rm g}$ are observed below 100 K. (b) Comparing the measured 40 K conductance vs $V_{\rm g}$ data (open circle) with the calculated 1D subband occupation with only thermal broadening (dash dotted line) or both the thermal and the scattering broadening (solid line) considered. (c) Gate modulation of thermopower S at 100, 70, and 40 K. The dashed vertical lines are a guide to the eye, highlighting the appearance of peak in $S(V_{\rm g})$ when a 1D subband starts to be filled. (d) Calculated density of states vs 1D electron density in nanowire with the index of subbands marked.

that the scattering broadening $\delta E = 28$ meV used in the calculation is in good agreement with the uncertainty principle estimation h/τ for the 40 K mobility ($\sim 8000 \text{ cm}^2/(\text{V s})$) of our nanowire. As we discuss later, the scattering induced level broadening is actually the dictating factor in limiting the

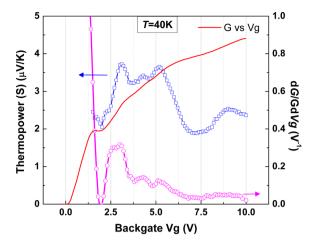


Figure 3. Gate dependence of thermopower, S (blue), conductance G (red), and $\mathrm{d}G/\mathrm{G}dV_{\mathrm{g}}$ (magenta) at 40 K, showing the correspondence between thermopower oscillations and the conductance steps as electrons populate 1D subbands in InAs nanowire.

observation of 1D quantum confinement effect in thermopower and power factor at room temperature in our nanowires.

Gate dependence data of thermopower S at T = 100, 70, and 40 K are presented in Figure 2c. As shown by Figure 2c, the $S(V_{\sigma})$ curve progressively deviates from the monotonically decreasing function as T decreases. At T < 100 K, the gate voltage dependence of S develops multiple oscillations. These oscillations are more discernible at lower T, and the peak positions coincide with the rises in the corresponding conductance vs V_g data when the next 1D subband starts to be filled (Figure 2d). Comparing Figure 2b and c and the calculated 1D density of states in Figure 2d, we associate the peaks in thermopower data at $V_g = 3$, 5, and 8.8 V to be due to the filling of the third, fourth, and fifth 1D subbands. Unfortunately, due to the slow thermal response of the device at low temperatures and the difficulty of using 2ω thermopower measurement technique at frequency lower than 1 Hz, we were unable to measure S reliably down to the single subband regime where the sample impedance was high, although there might be some hint of the second subband filling effect in the lowest V_{σ} regime of our data.

More careful comparison of the conductance and thermopower data further shows a close connection between the electrical transport and thermopower data. In Figure 3, we plot $dG/GdV_g = d \ln(G)/dV_g$ at 40 K together with the thermopower data. It can be seen that there is an excellent match of the peak/dip positions between the two traces: whenever dG/GdV_g peaks/dips, there is a peak or dip in the thermopower. This indicates Mott's formula works at least at the qualitative level for our data. Since conductivity $\sigma = ne^2\tau$ m^* , one sees from the Mott's formula $S \propto d(\ln \sigma)/dE|_{E_v}$ that both $S(V_{\sigma})$ and dG/GdV_{σ} should have characteristics reflecting the DOS, dn/dE, and $d\tau/dE$, or how the scattering time τ varies with energy. From the analysis in Figure 3 and calculated 1D subband population in Figure 2b, we believe that the oscillating 1D DOS is indeed mainly responsible for the oscillating S and $\mathrm{d}G/G\mathrm{d}V_\mathrm{g}.$ We also show the original $G(V_\mathrm{g})$ curve in Figure 3 for comparison. Examining the $G(V_g)$ curve carefully does reveal slope changes in connection with the thermopower oscillations. But since thermopower detects the derivative of energy dependent conductivity, characters of 1D DOS oscillations are more easily seen in $S(V_{\sigma})$ data.

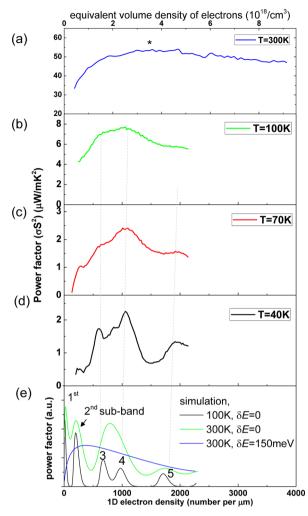


Figure 4. 1D quantum confinement effect in the power factor of InAs nanowire from 40 to 300 K. (a–d) Gate modulated thermoelectric power factor σS^2 at 300, 100, 70, and 40 K plotted as σS^2 vs 1D density (bottom axis) or equivalent volume density (top axis) of electrons in InAs nanowire which has diameter d=23 nm. The 1D subband effect is discernible at 100 K and below. (e) Calculated power factor vs electron density for circular InAs nanowire with d=23 nm, under relaxation approximation and assuming $\tau \sim E^{-1/2}$. Different curves are results from calculations with or without scattering induced level broadening.

This gate-controlled subband filling gives us an opportunity to control and optimize the thermoelectric properties of nanomaterials. Here we consider the power factor σS^2 , a key parameter for the thermoelectrics since it is the numerator in the expression for ZT. Figure 4 displays the gate modulated power factor at 300, 100, 70, and 40 K. To make the discussion more meaningful/universal, we plot the σS^2 as a function of electron density instead of gate voltage. As shown in Figure 4a, power factor at 300 K resembles that of a 3D bulk system; that is, it is a smooth function peaked at some characteristic carrier density ($\sim 1600e/\mu m$ or $3.5 \times 10^{18}/cm^3$, marked by a star in Figure 4a). Measurements on additional devices confirm that the power factor is always optimized at an electron concentration somewhat above $10^{18}/\text{cm}^3$ at 300 K for our In As nanowires, although the absolute value of the maximal σS^2 can vary from sample to sample depending on the mobility of particular nanowire (Figures S4-7, Supporting Information). This optimal electron density for our nanowire is close to the

theoretical value (~1018/cm3) for bulk InAs.12 At 100 K and below (Figure 4b-d), the 1D quantum confinement effect induces oscillatory behavior due to the DOS of 1D subbands being resolved. (Since the thermopower S is roughly proportional to T as shown in eq 1, the absolute value of power factor decreases significantly as the temperature is reduced.) Apparently, the power factor is peaked when the fourth 1D subband starts to be populated at electron density $\sim 1000 \ e/\mu m$, a value slightly lower than the optimal n at 300 K. Similar oscillatory behavior of power factor and the optimal electron density were observed in devices #2 and #3 (Figures S5c and S7c, Supporting Information). We also point out that, when the 1D subbands are resolved, the power factor exhibits much larger modulation (e.g., 500% change at 40 and 70 K vs ~60% modulation at 300 K) over the same carrier density range (Figure S3, Supporting Information).

Before concluding the paper, we remark on additional insights obtained from our gate enabled power factor control via the 1D electron confinement in InAs nanowire. Originally, it was believed that the thermoelectric power factor is always maximized when the Fermi level is placed right at the position of singular DOS and smaller number of subbands is preferred. 10,11 Our experiment indicates that the general idea of utilizing singular DOS to modulate the power factor in nanomaterials is effective. However, several important questions remain to be understood further such as where the optimal Fermi energy lies and if the system has to be in the single subband regime. Here, we found that, although we cannot rule out possible upturn of power factor in the experimentally immeasurable one-subband regime, the power factor peak associated with the third subband (and possibly second subband) is clearly lower than the fourth one. Therefore, it is not clear if less subband filling will give rise to higher power factor. At low carrier density regime, there is also a practical issue of increased contact resistance that can limit the power factor. In addition, our result and analysis emphasize that making small-sized material alone is not sufficient to induce low-dimensional confinement enhanced thermoelectric properties. One also has to reduce carrier scattering (improve the carrier mobility) such that the scattering broadening would not wash out the otherwise sharp DOS spectrum. For example, in Figure 4e we present two simulated curves of σS^2 vs n at 100 and 300 K, assuming no scattering broadening ($\delta E = 0$) and $\tau \sim E^{-1/2}$, a $\tau(E)$ dependence widely used for nanowires.³⁵ The calculation predicts power factor oscillations at locations close to the experimental data, but the predicted oscillation amplitudes are much greater and increase as the number of subbands decreases. Moreover, the theoretical calculation with $\delta E = 0$ exhibits clear oscillations even at 300 K, in contrast to the experimental observation which is closer to the 3D behavior. We found that proper scattering broadening is needed in the calculation to improve the agreement. As shown by the blue line in Figure 4e, adding 150 meV scattering broadening (estimated from the room temperature mobility of nanowire) does wash out the 1D oscillatory features and restores the qualitative 3D behavior. Based on this observation, it appears that obtaining 1D confinement enhanced thermoelectric power factor at room temperature is possible in experimentally accessible parameter ranges for higher mobility InAs nanowires (e.g., 10-20 nm wide InAs/InP core/shell nanowires with $>10^4$ $cm^2/(V s)$ mobility at room temperature²⁶) such that both the thermal and scattering broadening effects are smaller than the

subband separation energy. This may open up a route towards further enhancing ZT in nanowires for practical applications by 1D quantum confinement of carriers, given that the thermal conductivity is known to be strongly reduced in nanowires. Note that the importance of scattering broadening of singular DOS spectrum was recently brought up from the theoretical perspective as well. 14

In conclusion, we realized the gate controlled thermoelectric properties of InAs nanowires. Below 100 K, gate voltage is shown to induce pronounced thermopower and power factor oscillations from the filling of 1D subbands for the first time in semiconductor nanowires in the diffusive transport regime. This work demonstrates the feasibility of experimentally optimizing 1D nanostructure's thermoelectric properties by tuning the Fermi level across the peaks of 1D electronic density of states, a long-sought goal in nanostructured thermoelectrics research. Moreover, we point out the importance of reducing scattering induced level broadening in the pursuit of 1D quantum confinement enhanced thermoelectric properties in quasi-1D semiconductors at practical temperatures (e.g., 300 K).

ASSOCIATED CONTENT

S Supporting Information

Detailed description of the calibration procedure of the thermometers and the corresponding temperature gradient along nanowire, gate-dependent conductance, and thermopower and power factor data for additional InAs nanowire devices. This material is available free of charge via the Internet at http://pubs.acs.org.

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Notes

The authors declare no competing financial interest.

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REFERENCES

- (1) Nolas, G. S.; Sharp, J.; Goldsmid, H. J. Thermoelectrics: Basic Principles and New Materials Developments; Springer: New York, 2001.
- (2) Majumdar, A. Science 2004, 303, 777-778.
- (3) Dresselhaus, M. S.; Chen, G.; Tang, M. Y.; Yang, R. G.; Lee, H.; Wang, D. Z.; Ren, Z. F.; Fleurial, J. P.; Gogna, P. *Adv. Mater.* **2007**, *19*, 1043–1053.
- (4) Venkatasubramanian, R.; Siivola, E.; Colpitts, T.; O'Quinn, B. *Nature* **2001**, *413*, 597–602.
- (5) Harman, T. C.; Taylor, P. J.; Walsh, M. P.; LaForge, B. E. Science **2002**, 297, 2229–2232.
- (6) Poudel, B.; Hao, Q.; Ma, Y.; Lan, Y. C.; Minnich, A.; Yu, B.; Yan, X. A.; Wang, D. Z.; Muto, A.; Vashaee, D.; Chen, X. Y.; Liu, J. M.; Dresselhaus, M. S.; Chen, G.; Ren, Z. F. Science 2008, 320, 634–638.
- (7) Li, D.; Wu, Y.; Kim, P.; Shi, L.; Yang, P.; Majumdar, A. Appl. Phys. Lett. 2003, 83, 2934–2936.
- (8) Shi, L.; Hao, Q.; Yu, C. H.; Mingo, N.; Kong, X.; Wang, Z. L. Appl. Phys. Lett. **2004**, 84, 2638–2640.

- (9) Zhou, J. H.; Jin, Q.; Seol, J. H.; Li, X.; Shi, L. Appl. Phys. Lett. 2005, 87, 133109.
- (10) Hicks, L. D.; Dresselhaus, M. S. Phys. Rev. B 1993, 47, 16631–16634.
- (11) Mahan, G. D.; Sofo, J. O. Proc. Natl. Acad. Sci. U.S.A. 1996, 93, 7436–7439.
- (12) Mingo, N. Appl. Phys. Lett. 2004, 84, 2652-2654.
- (13) Farhangfar, S. J. Phys. D: Appl. Phys. 2011, 44, 125403.
- (14) Zhou, J.; Yang, R. G.; Chen, G.; Dresselhaus, M. S. Phys. Rev. Lett. 2011, 107, 226601.
- (15) Boukai, A. I.; Xu, K.; Heath, J. R. Adv. Mater. 2006, 18, 864–869.
- (16) Boukai, A. I.; Bunimovich, Y.; Tahir-Kheli, J.; Yu, J. K.; Goddard, W. A.; Heath, J. R. *Nature* **2008**, *451*, 168–171.
- (17) Hochbaum, A. I.; Chen, R. K.; Delgado, R. D.; Liang, W. J.; Garnett, E. C.; Najarian, M.; Majumdar, A.; Yang, P. D. *Nature* **2008**, 451, 163–167.
- (18) Small, J. P.; Perez, K. M.; Kim, P. Phys. Rev. Lett. 2003, 91, 256801.
- (19) Llaguno, M. C.; Fischer, J. E.; Johnson, A. T.; Hone, J. Nano Lett. 2004, 4, 45–49.
- (20) Liang, W. J.; Hochbaum, A. I.; Fardy, M.; Rabin, O.; Zhang, M. J.; Yang, P. D. *Nano Lett.* **2009**, *9*, 1689–1693.
- (21) Ford, A. C.; Bala Kumar, S.; Kapadia, R.; Guo, J.; Javey, A. Nano Lett. 2012, 12 (3), 1340–1343.
- (22) Liang, D.; Sakr, M. R.; Gao, X. P. A. Nano Lett. 2009, 9, 1709–1712.
- (23) Liang, D.; Gao, X. P. A. Nano Lett. 2012, 12 (6), 3263-3267.
- (24) Thelander, C.; Agarwal, P.; Brongersma, S.; Eymery, J.; Feiner, L. F.; Forchel, A.; Scheffler, M.; Riess, W.; Ohlsson, B. J.; Goesele, U.; Samuelson, L. *Mater. Today* **2006**, *9*, 28–35.
- (25) Dayeh, S. A.; Aplin, D. P. R.; Zhou, X. T.; Yu, P. K. L.; Yu, E. T.; Wang, D. L. Small 2007, 3, 326–332.
- (26) Jiang, X.; Xiong, Q.; Nam, S.; Qian, F.; Li, Y.; Lieber, C. M. Nano Lett. 2007, 7, 3214-3218.
- (27) Ford, A. C.; Ho, J. C.; Chueh, Y.; Tseng, Y. C.; Fan, Z.; Guo, J.; Bokor, J.; Javey, A. *Nano Lett.* **2009**, 9 (1), 360–365.
- (28) Du, J.; Liang, D.; Tang, H.; Gao, X. P. A. Nano Lett 2009, 9, 4348–4351
- (29) Sakr, M. R.; Gao, X. P. A. Appl. Phys. Lett. 2008, 93, 203503.
- (30) Lin, P. A.; Liang, D.; Reeves, S.; Gao, X. P. A.; Sankaran, R. M. *Nano Lett.* **2012**, *12*, 315–320.
- (31) Rowe, D. M. CRC Handbook of Thermoelectrics, 1st ed.; CRC Press: Boca Raton, FL, 1995.
- (32) Cutler, M.; Mott, N. F. Phys. Rev. 1969, 181, 1336-1340.
- (33) Colinge, J. P.; Quinn, A. J.; Floyd, L.; Redmond, G.; Alderman, J. C.; Weize, X.; Cleavelin, C. R.; Schulz, T.; Schruefer, K.; Knoblinger, G.; Patruno, P. *IEEE Electron Device Lett.* **2006**, *27*, 120–122.
- (34) It should be noted that there is a slight difference between the 40 K $G(V_g)$ data in Figure 2 parts a and b since Figure 2a data were taken at 0.1 V/s gate sweep rate whereas data in Figure 2b were taken at 0.1 V/10 s sweep rate to be consistent with the thermopower vs V_g experiment in which a long time constant is required. Also, to generate the theoretical $N_{\rm 1Dsubbands}$ vs V_g curve in Figure 2b, we convert the computed $N_{\rm 1Dsubbands}$ vs electron density n curve into a plot of $N_{\rm 1Dsubbands}$ vs V_g by using $n = C_g V_g / e$ where $C_g = 2\pi \varepsilon \varepsilon_0 L/\ln(2h/r)$ with an effective dielectric constant of 3.0. Such a reduced effective dielectric constant of SiO₂ is reasonable for backgated nanowire devices to take into account the vacuum gap and finite size effect (Wunnicke, O. Appl. Phys. Lett. 2006, 89, 083102). Note that, in the calculation of $N_{\rm 1Dsubbands}$ we used 21 nm as the nanowire diameter for the 1D subband energy level calculation, subtracting the native oxide thickness (~2 nm) on the surface of nanowire.
- (35) Seol, J. H.; Moore, A. L.; Saha, S. K.; Zhou, F.; Shi, L.; Ye, Q.; Scheffler, R.; Mingo, N.; Yamada, T. J. Appl. Phys. 2007, 101, 023706.
- (36) Zhou, F.; Moore, A. L.; Bolinsson, J.; Persson, A.; Fröberg, L.; Pettes, M. T.; Kong, H.; Rabenberg, L.; Caroff, P.; Stewart, D. A.; Mingo, N.; Dick, K. A.; Samuelson, L.; Linke, H.; Shi, L. *Phys. Rev. B* **2011**, 83, 205416.