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OPEN Large-Area Semiconducting Graphene Nanomesh Tailored b Interferometric Lithography

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Graphene nanostructures are attracting a great deal on erest because of newly emerging properties originating from quantum confinement effects. We report vising interferometric lithography to nanomesh (GNM) with sub-10 nm neck fabricate uniform, chip-scale, semiconducting grap. widths (smallest edge-to-edge distance between two nanoholes). This approach is based on fast, low-cost, and high-yield lithographic technologies and demonstrates the feasibility of cost-effective development of large-scale semicor fucting traphene sheets and devices. The GNM is estimated to have a room temperature energy bar rap o ~30 meV. Raman studies showed that the G band of the GNM experiences a blue shift and b. dening compared to pristine graphene, a change which was attributed to quantum . from and localization effects. A single-layer GNM field effect transistor exhibited promising core current of ~3.9 µA/µm and ON/OFF current ratios of ~35 at room temperature. The N/OFF current ratio of the GNM-device displayed distinct temperature dependence with bout 24 Id enhancement at 77 K.

Graphene has a statistic fascinating two-dimensional (2D) material for its high carrier mobility¹, flexibility², tra marency, and its extraordinary ability to withstand mechanical stress⁴. Due to these properties, graphen is self to many applications in electronics^{5,6}, optoelectronics^{7,8}, sensing, and energy stor-^{9,10}. However, graphene lacks a bandgap, which limits its application in digital electronics and optoele ronics Extensive efforts¹¹⁻²¹ have been made to create semiconducting graphene without disrupting Monal transport properties. One scheme is to obtain bandgap through quantum confinement in phene nanoribbons (GNRs)^{22,23} and graphene nanomesh (GNM)^{20,21,24} with a critical dimension lower than 10 nm. These nanostructures enable potential applications of graphene in electronic and photonic devices, such as highly sensitive sensors^{25,26}, in next-generation spintronics^{27,28}, and in energy harvesting^{29,30} devices. Duan and co-workers²⁰, fabricated semiconducting graphene using copolymer lithography and demonstrated that GNM-based field-effect transistors (FETs) exhibit comparable ON/OFF current ratios but 100 times higher drive currents than those of the similar devices based on individual GNRs. Furthermore, because of its excellent stability, 2D nature, and high electrical sensitivity, GNM is a good candidate to replace traditional solid-state nanoporours materials in fabrication of composite materials³¹ and electrochemical capacitors³² as well as in DNA sequencing³³.

The fabrication of graphene nanomesh for practical applications demands efficient nanopatterning technologies to inexpensively produce nanoscale periodic modulations over large-areas. To our knowledge, a reliable technology with a wafer-scale fabrication capability for graphene nanomesh is still a challenge. E-beam lithography is a highly versatile technique to fabricate graphene nanostructures. However, it is not an economically viable approach when large-scale production is required^{34,35}. Chemical methods,

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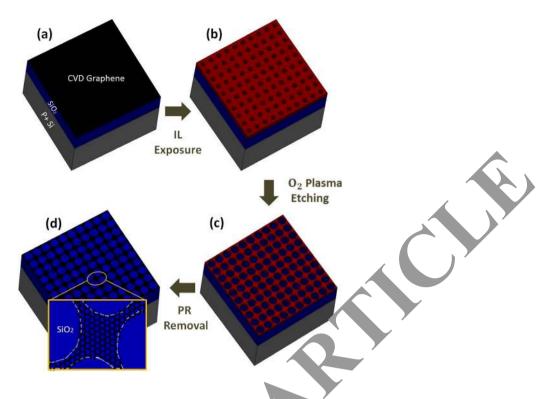


Figure 1. The overall fabrication route of GNM, b sec. Interferometric lithography and O_2 plasma etching. (a) SL-CVD graphene on 285 nm-thick SiO₂ ayer and P + Si substrate. (b) The sample is covered with a spin-coated negative photoresist (P = A square 2D hole array pattern is made in the PR by IL. (c) To define the mesh structure, an isotropic O_2 , usma etching is performed at high pressures. During the O_2 plasma step, the PR line width is recorded, v hich leads to a decrease in the graphene width under the photoresist. (d) The PR mask is removed as a brief soak in acetone solution.

such as copolymer 1 those phy²⁰ and imprint lithography³⁶ can meet the minimum feature size requirement, but they rel, on rather complex operations, and are still limited to microscale areas³⁷.

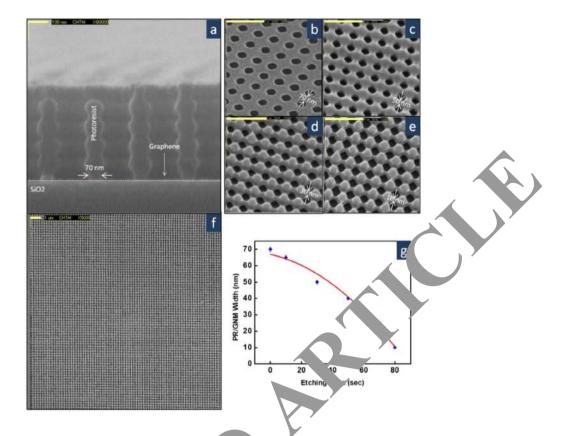
We demon trate the use of interferometric lithography (IL)³⁸⁻⁴⁴ combined with oxygen (O₂) plasma treatment to f oricate G NMs with sub-10 nm neck widths and high uniformity over a 1×1 cm² area. Our approach is factor inexpensive, and high throughput; furthermore it is based on well-established processing step, which have been extensively characterized for decades. Finally our approach is compatible with the integras, oricuits (IC) manufacturing technology⁴⁵. In our work atomic force microscopy (AFM) and Ramon spectroscopy were used to characterize the neck width of GNMs. The relative intensity of the D band increased with decreasing neck width. We also observed a blue shift and drastic broadening in the D band of the narrowest GNM, which can be attributed to the reduced size and edge effects. The GNM channel widths of 2 mm were fabricated to demonstrate its application in electronic decres. Transfer characteristics of the GNM-device were studied at room and cryogenic temperatures. The GNM-FET with neck width of ~10 nm showed promising drive current values and ON/OFF current ratio at room temperature. Bandgap modulation was observed by tuning the source-drain voltage. In addition, the ON/OFF current ratio of the GNM-FET displayed distinct temperature dependence with a significant enhancement at 77 K.

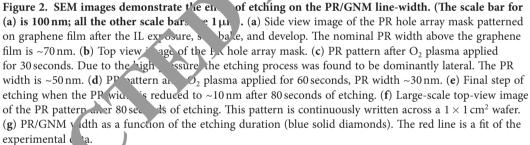
Results

Figure 1 illustrates the process flow for fabrication of the graphene nanomesh (GNM). We used commercially available single-layer (SL), chemical vapor deposited (CVD) graphene on a 285 nm silicon dioxide $SiO_2/p+$ silicon substrate (Fig. 1a). The sample is spin-coated with negative photoresist (PR) and a square 2D hole array pattern is defined in the photoresist by IL (Fig. 1b and Methods). Subsequently, an isotropic O_2 plasma etching is employed to reduce the PR neck width. In addition, the O_2 plasma removes the unprotected graphene regions, which leads to a decrease in the graphene width still coated with PR (Fig. 1c). After the etching process, the PR is removed by soaking the sample in acetone to obtain the patterned GNM (Fig. 1d).

The narrow neck regions of the GNM strongly affect the charge transport through the structure²⁰. Hence, the ability to precisely control the neck width is crucial for modulating the electronic properties in GNM. For GNMs made by IL, the neck width is tailored independently by varying the laser exposure dose and etching duration (Supplementary Information). We demonstrated the scalability of our

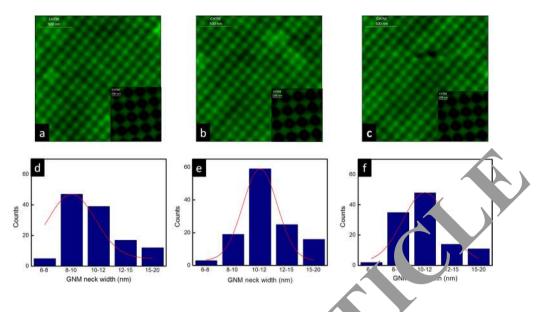


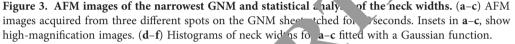




approach by first making a highly uniform PR hole array pattern (serving as an etch mask) on three includual sL-CVD graphene samples and then performing O_2 plasma etching on the processed samples r universe time durations. The initial width and periodicity of the pattern transferred to the PR are \sim mm and \sim 150–200 nm, respectively. Figure 2a,b show the cross section and top-view of a representative hole array mask. An optimized isotropic O_2 plasma etching was carried out separately for 30, 60, and 80 seconds on the processed samples (see Methods). The corresponding SEM images are shown in Fig. 2(c-e). The cross-sectional SEM image in Fig. 2a reveals the corrugated sidewalls of the holes in the PR. Rough PR sidewalls are attributed to standing waves, resulting from the lack of an anti-reflection coating (ARC) layer. Therefore, a narrower GNM is expected compared to the hole array width on the PR surface. Figure 2g plots the average PR/GNM width as a function of etching time. A low magnification SEM image of the PR pattern on graphene after 80 seconds of etching is shown in Fig. 2f to show the uniformity of IL exposure and O_2 plasma etch.

AFM images to characterize GNM neck-width were acquired from different regions of the GNM. GNMs with variable neck widths in the range of 6–20 nm are shown in Fig. 3a-c. More than 75% of the GNMs have neck widths between 8 nm and 12 nm. The standard deviation of the neck width is less than 3 nm over the entire GNM and the smallest measured neck width is $\sim 7 \pm 1$ nm. The AFM height profile of a typical GNM (Supplementary Information) suggests that after the standard cleaning of the sample (see Methods), a thin PR residue $\sim 15-20$ nm is left on the graphene sheet, which potentially introduces some doping⁴⁶ in the GNM and also affects the charge carrier mobility⁴⁷ in the mesh structure. The nanomesh feature, with very minor breaks, was observed over the 1×1 cm² wafer. The AFM profile studies on a set of samples with different etching durations (Supplementary Information) clearly revealed that highly uniform GNMs can be obtained with controllable neck widths using interferometric lithography.





Discussion

In order to study the physical and electronic characteristics of the processed samples, GNMs with varying neck widths were characterized using Raman spectroscopy (see Methods). The Raman spectrum acquired from unpatterned graphene (Fig. black line) shows the G band centered at 1587 cm⁻¹, which corresponds to the stretching vibra on of orbon atoms⁴⁸. Also a symmetric 2D band of the double resonance process⁴⁸ is found contered at 2635 cm⁻¹ with a full width at half-maximum (FWHM) of 32 cm^{-1} . For the unpatterned graphene, only a low-intensity or no D band was observed. In contrast, for GNMs with varying neck with two or serve an increase in the relative intensity of the D band (centered at 1352 cm⁻¹) as the mesh neck width decreases (Fig. 4a). We attribute this feature to the increasing of the provide the prov edge-to-surface ratic a. s in samples with smaller neck widths⁴⁹. The I_D/I_G ratio of the unpatterned graphene and GNMs with efferent neck widths are shown in Fig. 4b. As shown in Fig. 4b, the G band characteristic of the GNM with 10 nm neck width is noticeably broader (10 cm⁻¹) and blue shifted by 11 cm⁻¹ comp red to the G band dominating the spectrum of unpatterned graphene. This trend may be attributed to c_1 ntum confinement or localization effects resulting from edge disorders such as variable edge roughness 1.0 M^{50-52} . In addition the GNM mesh was patterned using O₂ plasma under a condition the brown to form oxygen-containing functional groups in graphite⁵³. Because oxygen is more electrone grave than carbon, such functional groups are expected to withdraw π electrons of GNM (i.e., α , with holes) and that potentially can induce a blue shift in the G band⁵⁴. However, for the GNM with widths of ~50 nm, no shift or broadening was observed (Fig. 4b), which suggests that the doping ne⁄ ect is less dominant. Also, because the narrowest GNM in this study has a sub-10 nm neck width, a ne negligible confinement effect is expected. To validate the observed behavior of the G-band, multiple Raman scans were performed at different locations across the narrowest GNM sample. Figure 4c (inset) shows plots of the G-band peaks acquired at four different locations on the sample. The results plotted in Fig. 4c show that all of the G band peaks tend to stay at approximately $1605 \,\mathrm{cm}^{-1}$ by a maximum shift of 17 cm⁻¹ with respect to the unpatterned graphene. Also, the measured FWHM of the G bands at the four locations are within a range of 42-47 cm⁻¹, indicating that, on average, the G band of the GNM is 10 cm⁻¹ broader than that of the unpatterned graphene. The blue shift and broadening of the G band for the narrowest GNM is likely due to the lateral quantum confinement and localization effects and/or chemical doping by functional groups on the GNM edges.

In order to study the transport characteristics of the GNM, a field-effect transistor (Supplementary Information) with the mesh structure serving as the conduction channel was fabricated (see Methods). Due to its 2D nature, the drive current of graphene devices, can be scaled up by increasing the device area. This is of great significance for realizing high-frequency devices with sufficiently high drive current in large circuits⁵⁵. Since our IL technique enables the fabrication of GNMs over of $1 \times 1 \text{ cm}^2$ area, the channel length and width of the transistor are chosen to be $14 \mu m$ and 2 mm, respectively. Figure 5a shows a schematic representation of the GNM-FET. Plots of source-drain current (I_d) as a function of gate voltage (V_g) at a constant source-drain voltage (V_{ds}) are shown in Fig. 5b. The SL GNM-device delivered ~3.9 μ A/ μ m current with an ON/OFF current ratio of ~9 at $V_{ds} = -1$ V. This is due mainly to a large GNM conduction channel filled with high density graphene ribbons capable of driving high currents.



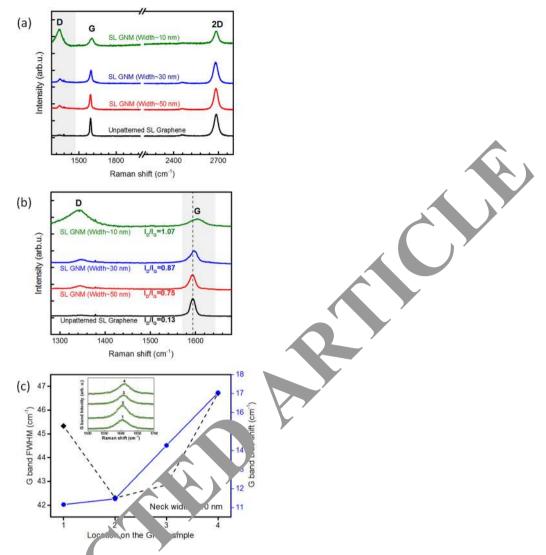


Figure 4. Rar on Spectra of single-layer CVD GNM. (a) Raman spectra of the GNM films with different neck widths concred to pristine single-layer graphene. The spectra were offset for clarity. (b) Close-up view to the spectrum acquired from a showing the G and D bands for the mesh structures compared to the pristine grap. Sheet. (c) Inset: G-band peaks acquired from four Raman scans on four different locations of the nan rowest GNM sample. The intensity bands are fitted using Gaussian functions. The FWHM and blu shift of the G bands of the Raman spectra acquired from four different locations on the narrowest NMM is referred to as the left axis while the G band blue shift is referred to as the using the spectra for those four locations.



Importantly, at $V_{ds} = -100 \text{ mV}$, although the ON-state current density is reduced to ~2.5 μ A/ μ m, the ON/OFF current ratio has increased up to ~35. The modulation of the ON/OFF current ratio by varying the source-drain voltage is a characteristic of the GNM as it has not been reported for pristine graphene devices⁵⁶. The measured dependence of the ON/OFF current ratio on V_{ds} indicates the semiconducting nature of the GNM²². This behavior is attributed to the formation of an electronic bandgap in graphene as a result of quantum confinement, edge and localization effects^{36,57}. The latter also suggests that the blue shift and broadening in the G band for the narrowest mesh is likely due to confinement and localization effects. Multiple minor conduction plateaus are found in the $I_d - V_g$ transfer curves (Fig. 5b), possibly due to the formation of electronic subbands in the GNM³⁶. The $I_d - V_{ds}$ for the GNM transistor (Fig. 5c) is non-linear, regardless of the applied gate voltage. In addition, although the Fermi-level modulation by the gate voltage is present, no saturation regions of the $I_d - V_{ds}$ curves are observed. The ON-state current of the GNM-FET is approximately 9 orders of magnitude lower than the one measured for an unpatterned graphene device (Fig. 5d). The ON current obtained from the GNM-FET is comparable to the results obtained for few-layer GNM-FETs^{36,37,58}, and it is considerably higher than the reported GNR-FETs with similar width^{59,60}. Moreover, the gate voltage sweeping range in the GNM-device is found to be smaller

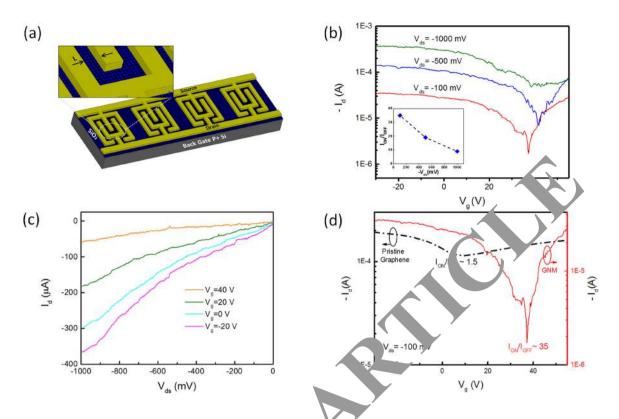


Figure 5. Transfer characteristics of graphene nanomesh. (a) A schematic representation of the GNM-FET. The graphene mesh structure with an encage neck width of ~10 nm was fabricated on heavily doped silicon substrate coated by 285 nm-th. SiO₂ , er as the gate dielectric. The channel length L and width W of the GNM-device are 14 µm and 21. The pectively. (b) I_d versus gate voltage (V_g) recorded at different source-drain voltages obtained at room ten, erature. (c) Source-drain current (I_d) versus source-drain voltage (V_{ds}) recorded at difference are v ltages at room temperature. (d) Comparison of $I_d - V_g$ for bulk (black dashed line) and tNM-device (red solid line) at $V_{ds} = -1$ mV at room temperature. The current for bulk graphene-device and TNM-device are specified on the left and right axis, respectively.

than that of the unpatterned graphene device. This is attributed to an increased leakage through the gate oxide after, attending. The reduced level of current compared to the bulk device is determined by the limit of current pathway and the introduction of edge defects. The comparison (Fig. 5d) also shows that the charge neutrality point for the GNM is shifted toward higher positive gate voltage regions. This mome on is attributed to the absorption of charged impurities at the graphene edges induced by oxi ation, huring the O_2 plasma treatment and polymer residues⁶¹. Despite the fact that absorption of barged impurities by graphene edge terminations in the GNM in principle plays a role in doping of the ctructure, its impact on the semiconductor behavior of the GNM (i.e. bandgap, ON/OFF ratio, etc.) could be affected by many different factors⁶². Such factors may include the type of doping, the position of the dopant with respect to the ribbon edges, the ribbon width, and type of ribbon symmetry (i.e. zigzag and armchair edges). Isolating these various effects would require further systematic investigations. The electronic properties of GNM made by IL along with neck-width and sheet coverage area are compared with previously reported results obtained using different processes in Table 1.

The overall current of the GNM displayed a significant temperature dependence (Fig. 6a), where back-gate modulation at $V_{ds} = -1 V$ increased from ~9 at room temperature to ~215 at 77 K. The ON/OFF current ratio of FET-devices exponentially scales with the bandgap as $I_{ON}/I_{OFF} \propto \exp(-E_g/k_BT)$, where k_B is Boltzmann constant and T is the absolute temperature⁶³. We speculate that, at very low temperatures, the carriers energy ($k_BT < < E_g$) was not sufficient to tunnel through the gap or hop through edge states and therefore the OFF-current state drops markedly. This temperature dependence is in strong contrast to that observed in unpatterned graphene FETs, where the pristine graphene is semi-metallic with a zero bandgap, and the temperature dependence of the transfer curves was negligible (Supplementary Information). Moreover, we noticed that the current plateau features are more pronounced at lower temperatures. The plateau structures are highly reproducible under different thermal cycles, and they are more pronounced at temperatures below 120 K. The former change in the plateau peaks is attributed to either stronger charge hopping through a series of quantum dots or to conduction associated with multiple one dimensional (1D) subbands generated in the mesh structure at lower temperatures⁵⁹.



	Fabrication Technique						
Properties	Block Copolymer Litho ²⁰	Block Copolymer Litho ²¹	Nano Imprint Litho ³⁶	Aluminum Oxide Templating ³⁷	Nano- Sphere Litho ⁵⁷	Nano- Sphere Litho ⁵⁸	Interferometric Litho ¹
Graphene type	SL^*/ML^{Υ}	SL	SL	SL	$N.R^{\intercal}$	SL	SL
Sheet coverage (μm^2)	N.R	106	4	N.R	N.R	N.R	108
Min neck width (nm)	5	18	7	15	20	65	10
ON/OFF Ratio [†]	100	39	80	10	10	2	35
Mobility [†] (cm ² /V.S)	N.R	>1	1000	N.R	N.R	31	80
Band gap [†] (meV)	100	102	N.R	N.R	N.R	0	30
Channel width (µm)	3	N.R	2	5.8	1.5	1	2
Channel length (µm)	2.3	N.R	2	2	20		2,00

Table 1. Comparison of dimensions and electronic properties of GNMs fab icated by a. erent techniques. ¹Present work. *Single layer graphene. ^TMultilayer/Few Layer graphene. ^TNot reported. [†]At room temperature. Note that the ON/OFF current ratio is a function of source train transformed (V_{ds}), and to obtain a reasonable comparison, one should look at the ratios at the same V_{ds} .

The effective mobility of holes in the SL-GNM produce, by IL are estimated to be ~80 cm² V⁻¹ s⁻¹ and ~165 cm² V⁻¹ s⁻¹ at room temperature and 77 K, separately (see Methods and Supplementary Information). The charge carrier mobility in GNM is lim. If by intrinsic scattering, phonon scattering from the supporting substrate, impurity scattering and line edge roughness (LER) scattering^{60,64}. The contributions of these scattering mechanisms comparately vary with the temperature, Fermi-level location, and the neck width of the graphene mesh⁶⁵. Therefore, the observed increase in the hole carrier mobility of the GNM, suggests that the rescattering mechanism is less effective at lower temperatures. These calculated values are relatively higher than reported carrier mobilities for single and few layer GNMs with the same average neck with (Ta le 1)^{21,58}. The electronic bandgap of GNMs inversely scales with the average neck width (ν) by the commentally derived equation $E_g \sim \alpha/w$, where α is between 0.3 and 1.5 with unit [nm eV]⁶⁶. Uso the ON/OFF current ratio of GNM-FETs varies exponentially with the bandgap, as specified by the to average 1.5

$$I_{ON}/I_{OFF} \propto \exp\left(-E_g/k_BT\right).$$
 (1)

Using the neasured ON/OFF current ratio at various temperature and inverting Eq. 1 to calculate the bandgap, e estimale $E_g = 30 \text{ meV}$ for a GNM with an average neck width of ~10 nm. As shown in Table 1, this varies in the range of those previously reported in the literature for a GNM with a comparable "bons width^{21,58}.

In summer we have proposed and demonstrated a simple yet effective approach to produce highly form sraphene nanomesh structures with sub-10 nm neck width across $1 \times 1 cm^2$ areas. The fabricatio of GNM using IL is scalable and can be performed using standard top-down fabrication methods. Learning the semiconducting behavior with an estimated bandgap ~30 meV. GNMs by various neck widths in the range of 50-10 nm have been fabricated on a macroscopic scale with an unprecedented level of control. Raman studies have shown a significant broadening and blue shift of the G band for the narrowest GNM. Such blue shift is attributed to quantum confinement effects. The GNM devices have been fabricated and they have shown drive currents ~3.9 µA/µm, ON/OFF current ratios ~35 at room temperature and ~215 at 77 K, and mobilities suitable for field- effective devices operating at room temperature. The availability of direct bandgap GNMs over large areas provides exciting opportunities for graphene-based photodetectors⁶⁷ and highly interconnected graphene networks⁶⁸. Further narrowing of the GNM neck width is indeed possible and requires optimization on the etching parameters (i.e. gas pressure, RF power, etc.) to provide a finer control over transition from 10 nm neck width to 5nm. Additionally, state of the art techniques such as scanning transmission electron microscopy (STEM) or AFM with carbon nanotube (CNT) tip will offer a more precise characterization for narrower mesh structures. IL parameters could also be used as additional variables in the GNM fabrication. Specifically, the laser wavelength and incident angle could be adjusted to make a structure with smaller periodicities, which in turn improves the drive current of the GNM-device.

Methods

Device fabrication. High-quality SL CVD graphene wafers on $285 \text{ nm SiO}_2/+p$ Si substrate with 95% uniformity were prepared from Graphene Supermarket Inc. The GNM-FETs were fabricated using standard photolithography (Figure S2). The heavily p-doped Si substrate and thermally grown 285 nm-thick



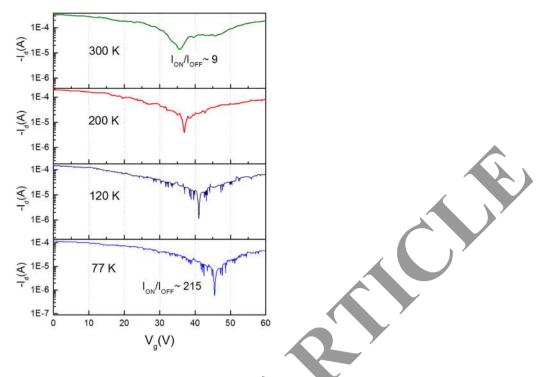


Figure 6. Temperature study of GNM transfer characteristics Comparison of $I_d - V_g$ curves of the GNM-FET with ~10 nm neck width recorded at $V_{ds} = -1$ V and different temperatures 300, 200, 120, and 77 K.

SiO₂ layers were used as a bottom ge and g te dielectric, respectively. To make a better back-gate contact, after removing the native adde on esticon substrate by dipping in buffer oxide etch (BOE) solution (6:1 volume ratio of 40° NF F in water to 49% HF in water), a 100 nm-thick Au film was sputtered onto the Si substrate. Then, so e and drain electrodes were deposited using photolithography followed by electron-beam even ration of "Au (5 nm/95 nm). The channel length (L) of the transistor varies from 2–14 μ m and the numel width (W) is 2 mm, providing maximum channel area of 28 × 10³ μ m². Next, interferome tic lithe upphy was performed on the fabricated FET. A negative tone-resist (NR7-500PY) with rominal thickness 500 nm was spun on the sample, followed by two IL exposures using a 355-nm triple YAG later source at an incidence angle $\theta = 30^{\circ}$ with a 90° rotation of the sample. The laser exposure me w/s 10 seconds, with pulse rate of 60 Hz and pulse energy of 100 mJ. After a hard bake at 110° for our econds, the sample was developed in an MF-321 developer for 30 seconds and subbele array pattern was made in the PR. Optimized O₂ plasma etching (O₂ flow, 10 sccm; sequently power, 100W; pressure, 200 mTorr) was performed for 80 seconds to reduce the PR linewidth and remove gruhene in selected areas. The PR/GNM lateral etch rate was 0.8 nm/sec and the vertical etch is estibe 1.8 nm/sec. This etching recipe resulted in significant reduction in the PR width, while the m thickness was only reduced by ~30 percent. Once the etching process is complete, the PR is removed by ample soak in acetone to obtain the final GNM. The samples were annealed at 300 °C in forming gas environment $(5\% H_2, 95\% N_2)$ for 30 min to remove/reduce the polymer resist from the etching mask, and other types of impurities on the mesh surface.

Device characterization. The electronic properties of mesh-patterned graphene devices were characterized by a JANIS ST-500 micro-manipulated probe station under vacuum (10^{-5} Torr) to eliminate/ reduce the hole doping effects due to absorption of oxygen and water molecules at the graphene surface. The probe station was equipped with a Scientific Instruments M9700 temperature controller, and the measurements were carried out with a Keithley 2400 source measure unit (SMU) and Keithley 238 high-current SMU. For comparison purposes, the GNM can be approximated as a mesoscopic honeycomb network of GNR in which the ribbons are interconnected between larger graphene islands. Therefore, the electron and hole mobilities of the GNM were estimated using the formula $\mu = (gL/V_{d-s}C_{ox}W)$, where g is the transconductance (Figure S5), L is the channel length, V_{ds} is the source-drain potential, C_{ox} is the gate capacitance per unit area, and W is the width of channel. $C_{ox} = (\varepsilon_0 \varepsilon_r/d)$, where ε_0 is the vacuum permittivity, ε_r and d are the dielectric constant and thickness of the SiO₂ dielectric layer. It should be noted that in order to study the GNM mobility more accurately, the fringe effect at the ribbon edges of the GNMs and also the exact capacitance of the transistor must be considered, which requires a simulation model based on finite element analysis. SEM imaging was performed on a JEOL- 6500 field-emission microscope. The Raman experiment was carried out by focusing a 532 nm



excitation light (output of laser diode) through a dry objective with numerical aperture 0.95. Prior to the Raman studies, the samples were annealed in forming gas environment $(5\% H_2, 95\% N_2)$ to remove polymer resist from the etching mask, and other types of impurities existed on the mesh surface. The scattered light is collected using the same objective and is directed to a CCD spectrometer (Princeton instruments, IsoPlane). The AFM (Neaspec GmbH Inc.) was operated in tapping mode with resonance frequency of cantilever ~250kHz and tapping amplitude ~50nm. An ultra-sharp AFM tip (Hi'Res-C14/ Cr-Au) with a spike radius of 1 nm was used. Statistical analysis on the acquired AFM images was carried out in AutoCAD software with careful tip corrections.

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Author Cont. tions

ed and designed this project and performed the fabrication of the work. He also wrote the A.K. co main manusci. It text. X.H. was involved in the fabrication process. J.GH. wrote the LabVIEW code for electrical characterization. T.H. performed the Raman spectroscopy. N.D. performed the AFM. S.A., F.C and S.B. provided useful discussions. S.K. was the supervisor and advisor of the project. All authors ntributed to analysis and interpretation of the results and contributed to writing the manuscript

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