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Low-temperature vapor reduction of graphene oxide electrodes for vertical organic field-effect transistors†

Kun Qiao,  Qing Ma, Junjia Wang and Binghao Wang *

Graphene based vertical organic field-effect transistors (VOFETs) are promising devices for realizing a high current density at a low operation voltage. However, high-quality graphene typically requires a high temperature and a complicated fabrication process, presenting a major barrier to roll-to-roll manufacturing. Here, we report a low temperature (≤ 45 °C), vapor (HI)-assisted process to achieve ultrathin and uniform reduced graphene oxide (rGO) films with a higher electrical conductivity compared with conventional high-temperature reduction methods. Notably, VOFETs with rGO source electrodes exhibited the highest current on/off ratio exceeding 10^4 and a maximum current density of 14.4 mA cm^{-2} owing to the gate-tunable work function characteristics of the prepared rGO. The current density of the device could be further improved by increasing the conductivity of the rGO electrode with additional thermal treatment. The synergy of low-temperature and HI vapor endowed rGO-VOFET devices with excellent current density, on/off ratio, and gate-tunable ability, making them promising candidates for future flexible organic electronics.

Introduction

The ultrashort channel length of the vertical organic field-effect transistors (VOFETs) enables the fabrication of devices in a smaller size with improved electrical performance compared with their lateral counterparts.^{1–4} So far, various VOFETs based on graphene electrodes have been reported due to their excellent conductivity in a two-dimensional form.^{5–14} In these devices, the work function tunability of graphene is exploited *via* an external gate electrical field to modulate the injection barrier height at the graphene/organic semiconductor interface.^{5–7} However, almost all these devices utilized single-layer graphene

obtained by the capital-intensive chemical vapor deposition (CVD) method. Although the CVD method can produce high-quality graphene, the rather complicated transfer process of graphene to the substrate is incompatible with future large-scale roll-to-roll applications.

Reduced graphene oxide (rGO) has a similar structure and properties to graphene.^{15–19} Most importantly, it can be easily mass-produced *via* reduction of low-cost solution-processable graphene oxide (GO). To achieve high-performance devices, the rGO film should possess not only high electrical conductivity, but also ultrathin and uniform film morphology. Although high temperature (> 800 °C) thermal reduction could prepare rGO films with high electrical conductivity, the involved high temperature is not applicable to emerging flexible electronics.^{20–23} In this regard, chemical reduction is preferred because of its low reduction temperature.^{24–28} In particular, chemical reduction by the vapor phase is more promising because dipping the substrate with the GO film in a hot reducing solution would inevitably ruin and even peel off the rGO film from the substrate,^{20,29} which would greatly degrade the device performance. The rGO films prepared by vapor phase reduction, such as with hydrazine vapor, normally suffer from poor electrical conductivity, and further thermal reduction treatment (> 300 °C) is usually required to improve their electrical properties.^{20,30} Therefore, using low-temperature vapor phase reduction to prepare rGO films with good electrical conductivity is highly desirable.

Another critical requirement for using rGO electrodes in VOFETs is ultrathin thickness. The operation of this type of VOFET is based on the gate-tunable work function mechanism,³¹ whereas the work-function tuning induced by the gate electrical field only occurs in an ultrathin rGO film.^{32,33} In addition, the ultrathin rGO electrode benefits the partial screening effect of the gate electrical field,^{4,11} which could further improve the device performance. In a previous work, a simple spin-coating and reduction method was proposed that can effectively prepare highly conductive and ultrathin rGO films.³⁴ By employing hydrazine monohydrate vapor reduction along with 300 °C treatment for 6 h, we achieved a

School of Electronic Science & Engineering, Southeast University, 2 Sipailou Road, Nanjing, Jiangsu Province, 210096, P. R. China. E-mail: binghaowang@seu.edu.cn
† Electronic supplementary information (ESI) available: Fabrication process of the device, optical images of rGO films, a height profile of the GO film, AFM images of the rGO films, and XPS spectra and Raman spectra of the rGO films. See DOI: <https://doi.org/10.1039/d3tc04217k>

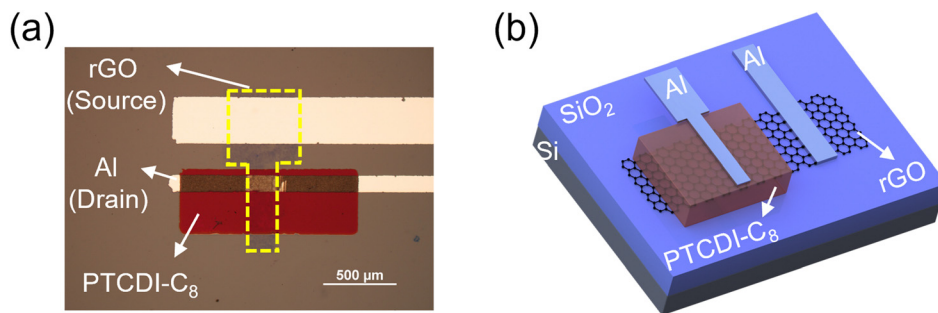


Fig. 1 (a) Top-view optical image and (b) three-dimensional schematic of the device.

conductivity of 10 S cm^{-1} for the rGO film and a current density of 0.73 mA cm^{-2} for VOFETs. However, it involved high temperature long-term thermal annealing, making it difficult for flexible devices.

In this study, we report a low-temperature ($\leq 45 \text{ }^\circ\text{C}$) HI vapor reduction of ultrathin GO films and used them as electrodes for VOFETs. Despite such mild reduction conditions, the resulting rGO film exhibits a sheet resistance of $0.56 \text{ M}\Omega \text{ sq}^{-1}$, which is better than those ($1.1 \text{ M}\Omega \text{ sq}^{-1}$) reduced by $300 \text{ }^\circ\text{C}$ annealing for 6 h. X-ray photoelectron spectroscopy (XPS) also demonstrates a higher reduction level by the HI vapor than thermal treatment. By sandwiching organic semiconductor *N,N'*-dioctyl-3,4,9,10-perylenedicarboximide (PTCDI- C_8) between the rGO and an Al electrode on an $\text{n}^+\text{Si}/\text{SiO}_2$ substrate, the fabricated device (rGO-VOFET) generated the highest current on/off ratio of over 10^4 and a maximum current density of 14.4 mA cm^{-2} .

Results and discussion

A schematic of the device fabrication process is illustrated in Fig. S1 (ESI †). An ultrathin GO film was first formed on an $\text{n}^+\text{Si}/\text{SiO}_2$ substrate by spin-coating a purified GO dispersion. The substrate was placed in a sealed container with HI solution, and the GO film was reduced by exposure to HI vapor. Notably, the reduction temperature was less than $45 \text{ }^\circ\text{C}$. After wet-

patterning of the rGO film, a 350 nm-thick PTCDI- C_8 layer and 50 nm-thick Al electrode were sequentially deposited through shadow masks using a vacuum thermal evaporator, and the final device rGO-VOFETs were fabricated. The top-view optical image and three-dimensional schematic of the device are shown in Fig. 1a and b, respectively. The active area of the device determined by the overlapping area of the top drain and bottom rGO electrode was $200 \times 100 \text{ }\mu\text{m}^2$.

Ultrathin and uniform films with high coverage are required for the gate-tunable work function electrode. 31 The morphology of the GO and rGO films were first evaluated by optical microscopy. As shown in Fig. 2a, the spin-coated GO film on the $\text{n}^+\text{Si}/\text{SiO}_2$ substrate shows a semi-transparent feature and high coverage. After HI vapor reduction, the rGO film (Fig. 2b) on the $\text{n}^+\text{Si}/\text{SiO}_2$ substrate exhibits better contrast due to the improved reflection of incident light. 35 This large-scale high coverage of the rGO film is further demonstrated in Fig. S2 (ESI †). The surface morphology of the GO and rGO films is evaluated by atomic force microscopy (AFM). Fig. 2c shows the spin-coated GO film before reduction, where ultrathin GO flakes uniformly cover the surface. The thickness of the GO film is about 1 nm (Fig. S3, ESI †), indicating its monolayer structure. 26 Importantly, this ultrathin and high-coverage film is maintained after reduction by exposure to HI vapor (Fig. 2d). In contrast, the surface of the rGO film reduced by dipping in HI solution is uneven and contaminated (Fig. S4, ESI †). The results reveal that vapor phase reduction is necessary to keep a clean and smooth surface of the rGO film.

The electrical conductivities of the rGO films are evaluated by measuring their sheet resistance (R_s) via a four-probe method, as it plays a critical role in device performance. For the low-temperature HI vapor reduction method, the impact of the reduction temperature and reduction time on the R_s are systematically investigated. First, the GO films are reduced by exposure to HI vapor at 30, 45, and $60 \text{ }^\circ\text{C}$ for 12 h, respectively. As shown in Fig. 3a, the R_s of the rGO film dramatically decreases from 0.94 to $0.56 \text{ M}\Omega \text{ sq}^{-1}$ as the reduction temperature increases from 30 to $45 \text{ }^\circ\text{C}$. Further increasing the reduction temperature to $60 \text{ }^\circ\text{C}$ does not reduce the R_s too much, suggesting that HI vapor treatment at $45 \text{ }^\circ\text{C}$ is enough to reduce the GO film. Next, the R_s of the rGO film is further evaluated by exposure of the GO film to HI vapor at $45 \text{ }^\circ\text{C}$ for 6, 12, 18, and 24 h, respectively (Fig. 3b). The R_s decreases



Binghao Wang

Binghao Wang is a Professor of Electronic Science and Technology at Southeast University. Prior to this, he was a Project Researcher at The University of Tokyo, and a Joint-PhD student and Postdoctoral Researcher at Northwestern University. He obtained his B.E., M.S., and PhD degrees from Soochow University. His research interests encompass flexible/wearable thin-film transistors and sensors, with a portfolio that includes over 80 co-authored papers and contributions to 2 book chapters.

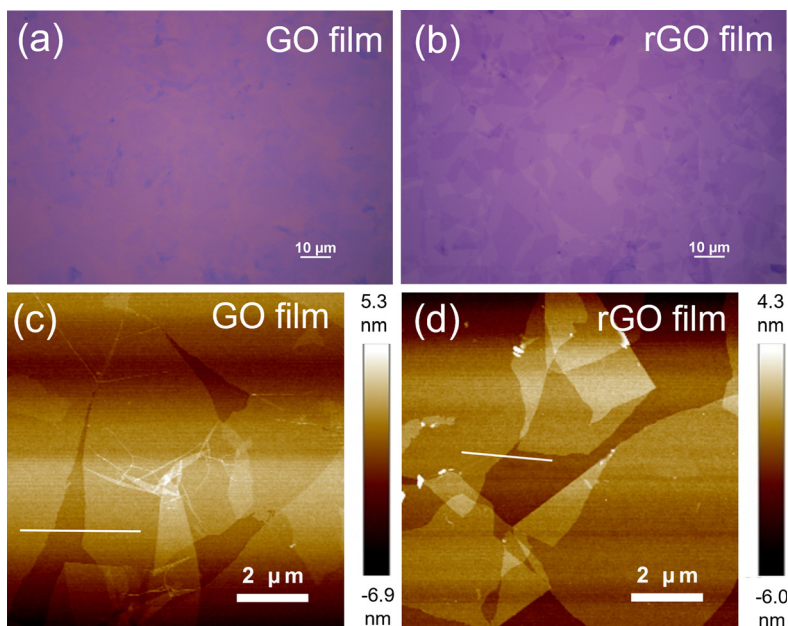


Fig. 2 Optical images of the (a) GO and (b) rGO films. AFM images of the (c) GO and (d) rGO films.

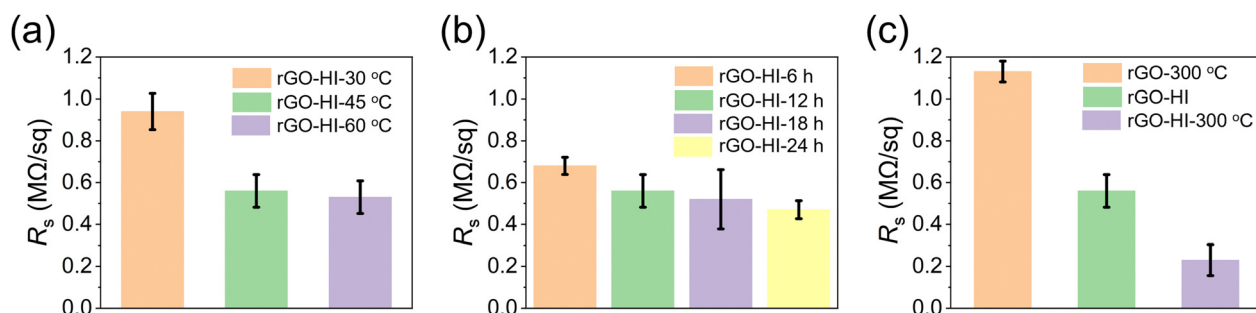


Fig. 3 (a) R_s of the rGO films reduced by HI vapor at 30, 45 and 60 °C for 12 h, respectively. (b) R_s of the rGO films reduced by HI vapor at 45 °C for 6, 12, 18, and 24 h, respectively. (c) R_s of the rGO-300 °C, rGO-HI, and rGO-HI-300 °C films, respectively.

slowly as the time increases, especially for that treated for more than 12 h. Thus, HI vapor treatment at 45 °C for 12 h was chosen as the optimal condition (rGO-HI). Moreover, we compare the R_s of the rGO films prepared by other methods (Fig. 3c). The rGO films were prepared by direct thermal annealing at 300 °C for 6 h (rGO-300 °C), and post annealing of rGO-HI at 300 °C for 1 h (rGO-HI-300 °C), respectively. Compared with the rGO-300 °C film, the electrical conductivity of the rGO-HI is significantly improved, indicating that this low-temperature HI vapor reduction method is more effective to reduce the rGO film. Moreover, by post annealing at 300 °C for 1 h, the electrical conductivity of the rGO-HI-300 °C film could be further improved.

To precisely reveal the reduction extent of the rGO films by these different methods, XPS analysis is performed. As shown in Fig. S5 (ESI[†]), the C1s XPS spectrum of the GO film can be divided into three peaks at around 288.1, 286.5, and 284.2 eV, arising from the C=O, C-O, and C-C bonds, respectively.^{25,26} The C=O and C-O bonds account for oxygen-containing

groups in the GO film. After reduction treatment, the intensities of the corresponding C=O and C-O peaks dramatically decrease, while the C-C peak becomes the dominant peak. Notably, compared with the C1s XPS spectrum of rGO-300 °C, the intensities of the C-O and C=O peaks of the rGO-HI film are much weaker, indicating a higher reduction level of the HI vapor-prepared rGO film. In addition, the reduction level of the rGO-HI film could be further improved by post annealing at 300 °C for 1 h. The corresponding C=O, C-O, and C-C peak intensities of these rGO films are summarized in Fig. 4. The XPS analysis consists of conductivity tests. Raman spectra are further investigated to study these rGO films (Fig. S6, ESI[†]). Two prominent peaks at 1353 and 1587 cm^{-1} in the Raman spectra correspond to typical D and G peaks, respectively. Compared with rGO-300 °C, the increased I_D/I_G ratios of rGO-HI and rGO-HI-300 °C indicate the increase of the sp^2 domain with decreased size,^{25,26} contributing to higher electrical conductivity.

The electrical conductivity of rGO relies on the recovery of its conjugated structure by eliminating these oxygen-containing

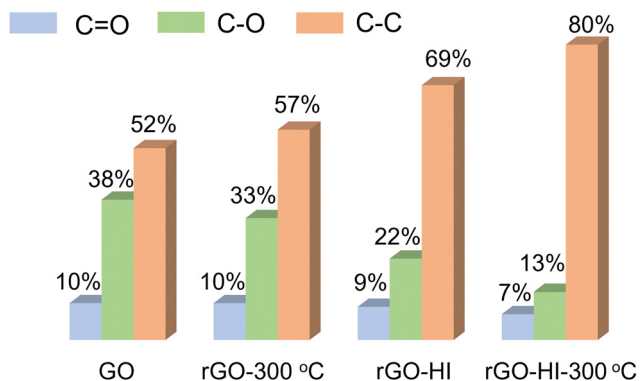


Fig. 4 The intensity ratios of C=O, C-O, and C-C peaks, obtained from the C1s XPS spectra of GO, rGO-300 °C, rGO-HI, and rGO-HI-300 °C, respectively.

groups (epoxy, hydroxyl, and carbonyl groups). For rGO-300 °C, the reduction level of the rGO film mainly depends on the reduction temperature. However, the temperature of 300 °C is

not high enough to considerably eliminate these oxygen functional groups.^{17,35} For the rGO-HI film, the reduction is through the substitution of oxygen functional groups by iodine atoms.^{29,36} Owing to the strong HI acid and weak C-I bond, the oxygen functional groups in the GO film can be effectively removed at a much lower temperature.³⁶

Furthermore, VOFETs were fabricated based on the rGO-300 °C, rGO-HI, and rGO-HI-300 °C films. The electrical properties of the devices are estimated at room temperature under dark conditions. The rGO source electrodes are grounded, whereas the top Al drain electrodes and bottom gate electrodes are biased with respect to the ground. In the output characteristics (Fig. 5a-c), a larger drain current (I_{DS}) is observed with a more positive gate voltage (V_{GS}), indicating that they are typical n-type transistors. Notably, the highest I_{DS} values of the devices gradually rise with the increasing electrical conductivity of the rGO electrodes. The transfer characteristics of the devices (Fig. 5d-f) are measured at the drain voltage (V_{DS}) of +30 V. As shown in Fig. 5d, the device based on the rGO-300 °C film shows a relatively low current on/off ratio (~ 100) with large

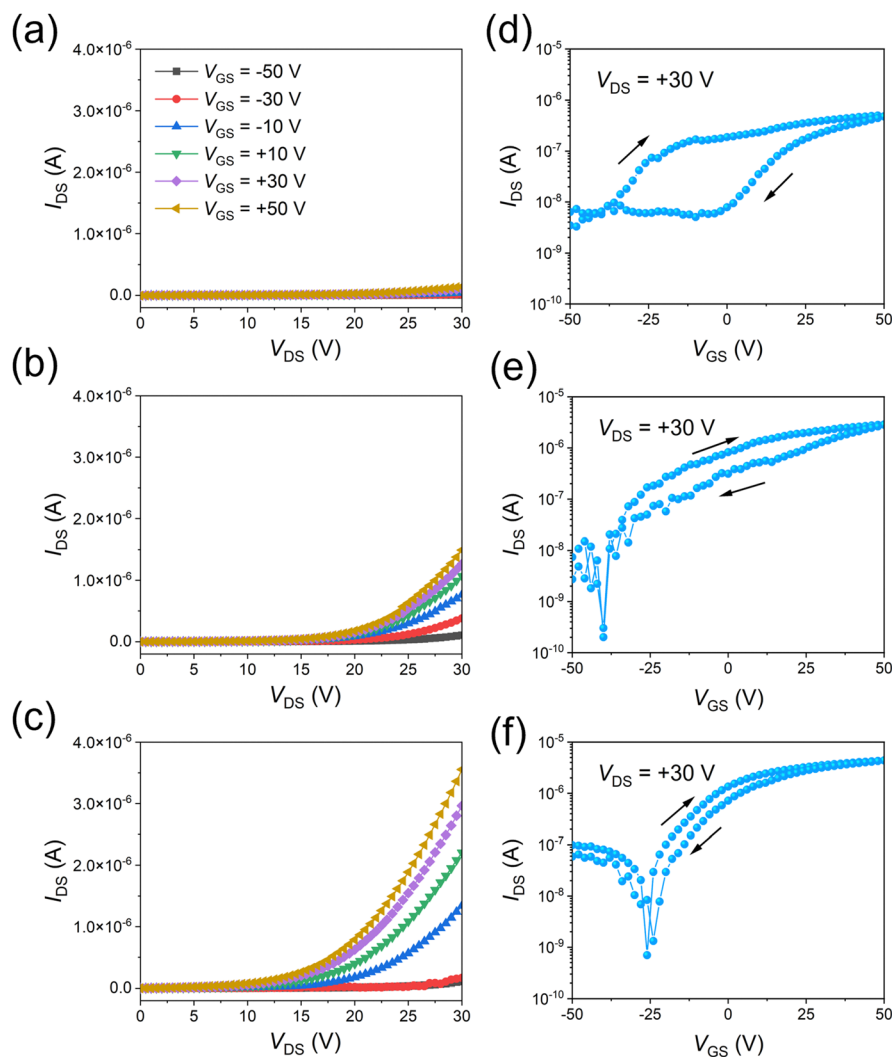


Fig. 5 (a)–(c) Output characteristics and (d)–(f) transfer characteristics of rGO-300 °C, rGO-HI, and rGO-HI-300 °C, respectively.

hysteresis, which could be attributed to the presence of the traps at the rGO/PTCDI-C₈ interface. As discussed before, 300 °C thermal treatment is not enough to reduce the GO film; therefore, there should be many oxygen functional groups remaining in the rGO-300 °C film, resulting in poor device performance. By contrast, the electrical properties of the device based on the rGO-HI film are significantly improved (Fig. 5e), exhibiting the highest current on/off ratio exceeding 10⁴ and a maximum current density of 14.4 mA cm⁻² with small hysteresis. The performance is comparable to those of VOFETs based on CVD-grown graphene electrodes.^{5,12} Fig. 5f illustrates the transfer characteristics of the device based on the rGO-HI-300 °C film. Due to the improved electrical conductivity of the rGO electrode, the highest current density of the device is further improved to 21.9 mA cm⁻².

Conclusions

In this work, VOFETs based on a low temperature-prepared rGO electrode were fabricated, achieving a large current density and a high on/off ratio. An ultrathin GO film with high coverage is first prepared by spin-coating a highly purified GO dispersion. Then, low-temperature HI vapor is utilized to reduce the spin-coated GO film to maintain the highly covered film structure and enable its application to future flexible devices. By using these mild reaction conditions, the prepared rGO film possesses very low sheet resistance of 0.56 MΩ sq⁻¹ and gate-tunable work function characteristics. The assembled rGO-VOFET devices exhibit satisfactory performance with the highest current on/off ratio of over 10⁴ and a maximum current density of 14.4 mA cm⁻². Moreover, the current density of the device is closely related to the conductivity of the rGO electrode and could be further improved. Importantly, we anticipate that this low-temperature HI vapor-prepared rGO electrode will help in advancing low-cost, high-performance, roll-to-roll future flexible organic electronics.

Experimental section

Purification of the GO dispersion

The commercial GO dispersion (4 mg mL⁻¹, GaoxiTech Corp, China) was firstly purified by a series of centrifugation treatments. The details of the centrifugation process are as follows: the GO dispersion was first diluted to 1 mg mL⁻¹, the solution was centrifuged at 3800 rpm for 15 min to remove the precipitate. Then, the supernatant liquid was centrifuged at 2300 rpm for 30 min to further remove the precipitate. This process was repeated 4 times. After that, the supernatant liquid was centrifuged at 4000 rpm for 1 h to collect the precipitate. After drying, the precipitate was diluted to 1 mg mL⁻¹ for further use.

Device fabrication process

The fabrication process of the rGO-VOFETs is as follows: the purified GO dispersion (300 μL) was dripped onto n⁺⁺Si/SiO₂

substrates (300 nm-thick), followed by spin-coating at 500 rpm/120 s, 1000 rpm/30 s and 2000 rpm/30 s sequentially to prepare the ultrathin GO film. Then, the spin-coated GO film was placed in a 250 mL sealed container with 3 mL HI solution (55–57%, Adamas, China), and the container was kept at different temperatures for different times. After the reduction treatment, the rGO film was patterned by a wet patterning method.³⁷ Briefly, 100 nm-thick Au was deposited through a shadow mask on the rGO film as a protective layer. Then, the substrate was placed in a beaker filled with bleaching solution (Kaneyo Corp, Japan) under UV light (365 nm, 3 W cm⁻²) irradiation directed on the substrate. After 2 min irradiation, the substrate was carefully rinsed with deionized water and dried by condensed air flow. Then the Au was removed by dipping in KI/I₂ solution for around 1 min. The substrate was rinsed with deionized water to remove the residual KI/I₂ solution and then dried by condensed air flow. KI/I₂ solution was prepared by dissolving 1.6 g KI powder (99%, Adamas, China) and 0.4 g I₂ (99%, Adamas, China) in 50 mL deionized water. Subsequently, 350 nm-thick n-type organic semiconductor PTCDI-C₈ (Luminescence Technology Corp., China) was deposited as an organic layer at a rate of 3 Å s⁻¹ through a shadow mask using a vacuum thermal evaporator. The vacuum chamber pressure was under 3 × 10⁻⁶ torr. Finally, a 50 nm-thick Al electrode was thermally deposited at a rate of 2 Å s⁻¹ under a vacuum pressure of 6 × 10⁻⁶ torr to fabricate the rGO-VOFET devices. The active area of the device determined by the overlapping area of the bottom rGO and top Al drain electrode is 200 × 100 μm².

Measurements and characterizations

The electrical properties of the devices were measured using a semiconductor parameter analyzer (Keithley 4200-SCS, US) at room temperature under dark conditions. The R_s of the rGO film was evaluated by using four-probe equipment (JG ST2258C, Suzhou Jingge Electronic Corp., China). The surface morphology of the GO and rGO films was investigated by using a Bruker Dimensional Icon atomic force microscopy system in tapping mode. XPS was performed on a JEOL JPS-9010MC spectrometer and O(1s) spectra were fitted using three Gaussian–Lorentzian convolution functions after subtracting a linear baseline.

Conflicts of interest

The authors declare no competing financial interest.

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