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# Visualization of a Maze-Like Reconstruction of Graphene on a Copper Surface at the Atomic Scale \*

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*Interaction with the substrate plays an essential role in determining the structure and electronic property of graphene supported by a surface. We observe a maze-like reconstruction pattern in graphene on flat copper foil. With functionalized scanning tunneling microscope tips, a triangular three-for-six structure of graphene and a mixed  $(2\sqrt{2} \times \sqrt{2})R45^\circ$  reconstruction of a Cu(100) surface are separately visualized at the atomic scale. Substrate-induced changes in the structure and electronic property are further illustrated by micro-Raman spectroscopy and scanning tunneling spectroscopy. This finding suggests a new method to effectively induce partial  $sp^3$  hybridization in a single-layer graphene and therefore to tune its electronic property through interaction with the substrate.*

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With a single layer of carbon atoms arranged in a honeycomb lattice,<sup>[1–9]</sup> graphene has become a prominent candidate for future applications, especially novel two-dimensional electronic devices, due to its unique linear energy dispersion at low energies.<sup>[10,11]</sup> Significant research progress has been made to tune the electrical properties of graphene through electrical gates,<sup>[12]</sup> dopants and defects,<sup>[13–16]</sup> local strain,<sup>[17–19]</sup> and substrates.<sup>[20–22]</sup> Associated with the electronic property changes, a three-for-six configuration has been observed on wrinkles of exfoliated graphene layers<sup>[17,18,23]</sup> and on strained graphene layers grown on highly corrugated copper surfaces.<sup>[19]</sup> Some  $sp^3$  hybridization is induced in such configurations to effectively open an energy gap and turn the graphene into a semiconductor or an insulator.<sup>[13–15]</sup> The interaction between graphene and the underlying substrate is crucial in determining the structure transformation and modification of electrical properties in graphene. However, it remains a great challenge to fully understand the graphene-substrate interaction because of the difficulty in probing the atomic structure of the underlying substrate. In addition, the observation of new reconstruction patterns due to graphene-substrate interaction is strongly desired.

In this Letter, we present our investigation of a maze-like reconstruction in graphene on a copper surface by using a low-temperature scanning tunnel-

ing microscope (STM). The reconstruction pattern is composed of a three-for-six triangular lattice and an orthogonal lattice, which are separately visualized with functionalized STM tips. The presence of partial  $sp^3$  hybridization in the triangular lattice is verified by the appearance of a midgap state near the Fermi level in the local electronic density of states. The orthogonal lattice is illustrated as a mixed  $(2\sqrt{2} \times \sqrt{2})R45^\circ$  reconstruction of a Cu(100) surface as supported by x-ray diffraction data. Our results suggest a strong interaction between the graphene and copper surface that has not been reported up until now.

The graphene sample used in this study is synthesized on Cu foil (50- $\mu\text{m}$ -thick) by the ambient chemical vapor deposition (CVD) method.<sup>[24,25]</sup> The growth is carried out at around 1050°C with  $\text{CH}_4$  as the carbon stock. The sample is then further annealed at 400°C for 6 h in an ultra-high vacuum (UHV) chamber (base pressure  $<10^{-10}$  Torr). A scanning electron microscope (SEM) and a home-built low-temperature STM are employed to characterize the sample surface. The local electronic density of states (LDOS) is measured by recording the STM differential conductance ( $dI/dV$ ) signal under open-loop feedback conditions. The sample bias is modulated at 361.1 Hz with an rms amplitude of 10 mV.

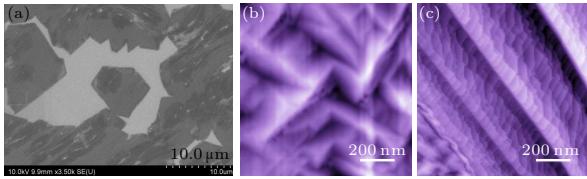
The SEM images in Fig. 1 show the morphology of the sample. The graphene flakes of a size around

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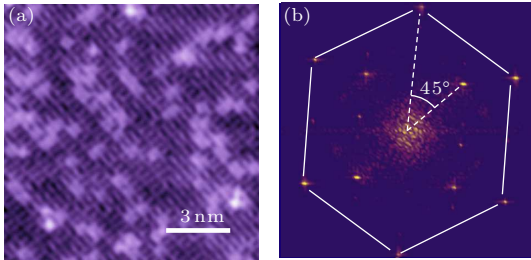
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10  $\mu\text{m}$  can be easily distinguished from the bare copper surface with characteristic hexagonal edges. Second layer graphene can be seen as smaller dark patches, which has been confirmed by micro-Raman studies.<sup>[25,26]</sup> The surface morphology of the graphene-covered area is further illustrated by STM images at the nanometer scale. Two typical topographic features are observed. Some areas appear highly corrugated, as shown in Fig. 1(b), and some show ordered steps and flat terraces of micrometer size, as shown in Fig. 1(c), where the straight steps are several hundreds of nanometers in length and 3–5 atomic layers in height.



**Fig. 1.** The morphology of graphene on a copper surface. (a) SEM image of the sample, and (b) and (c) STM topographic images of graphene on a corrugated surface and on terraces and steps. The images ( $1\ \mu\text{m} \times 1\ \mu\text{m}$ ) are taken with a sample bias of  $-0.7\ \text{V}$  and tunneling current of  $0.1\ \text{nA}$  for (b), and  $-0.5\ \text{V}$  and  $0.1\ \text{nA}$  for (c).

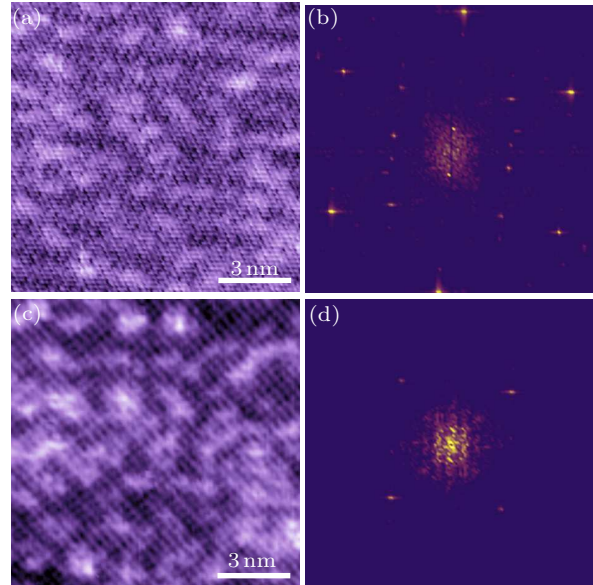


**Fig. 2.** (a) STM topographic image of a maze-like reconstruction pattern of graphene on a copper surface. The image ( $12\ \text{nm} \times 12\ \text{nm}$ ) is taken with a sample bias of  $-0.5\ \text{V}$  and tunneling current of  $15\ \text{pA}$ . (b) FFT of the maze-like reconstruction pattern. The six-fold diffraction spots are connected with white lines. The angle between two lattice sets is about  $45^\circ$ .

When the STM tip is located in the bilayer graphene region, Moiré patterns of bilayer graphene with different twist angles have been observed.<sup>[27]</sup> In the flat region covered with single-layer graphene, a maze-like reconstruction pattern has been observed, as shown in Fig. 2(a). The fast Fourier transform (FFT) of the STM image indicates that the maze-like reconstruction pattern is composed of a six-fold lattice and a four-fold lattice. The lattice constant of the six-fold lattice is  $2.5 \pm 0.05\ \text{\AA}$ , which is equal to that of graphene. The four-fold orthogonal lattice constant is  $3.6 \pm 0.05\ \text{\AA}$ . The angle between two lattice sets is about  $45^\circ$ , as indicated in Fig. 2(b). We notice that the four-fold symmetry is not perfect since the diffraction spots in one direction are stronger than those in the other direction.

By considering the presence of the graphene and copper substrate in the sample, it is clear that we can

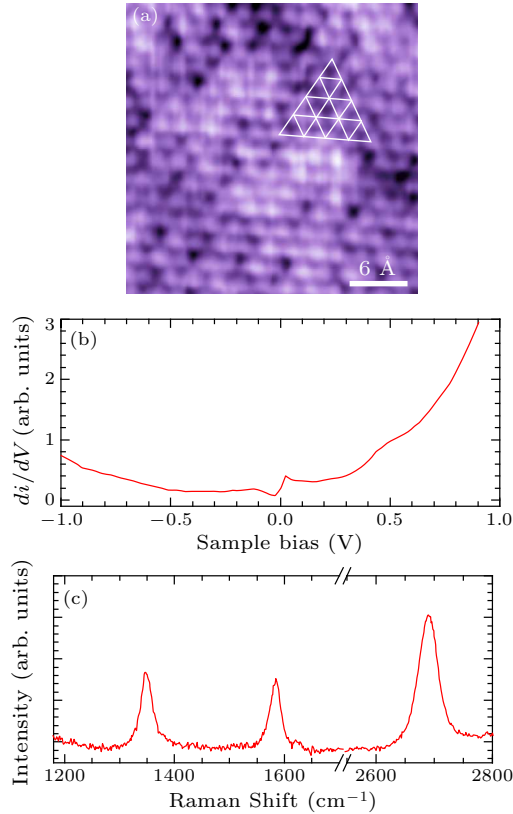
assign the six-fold lattice to the single-layer graphene and the four-fold lattice to the copper surface. To understand the interaction between the graphene layer and the underlying copper surface, atomic resolved images of both components are strongly desired, but this is usually very challenging. Fortunately, both the six-fold pattern and the four-fold pattern can be separately imaged with functionalized STM tips. Figures 3(a) and 3(c) show the STM images of the same area with a much clearer six-fold pattern and with a pure orthogonal four-fold pattern, respectively, as well as the FFT of the images. Although the four-fold lattice is still seen in the FFT image of Fig. 3(a), its intensity is significantly reduced as compared with that in Fig. 2(b), and the four-fold symmetry is barely seen in the STM image (Fig. 3(a)). The FFT image for the four-fold pattern (Fig. 3(d)), on the other hand, only shows a set of four-fold diffraction spots. Selectively imaging different electronic states has also been demonstrated in previous STM studies by manipulating the chemical identity at the apex of an STM tip.<sup>[28–30]</sup> The functionalized STM tips used in our experiment are generated by a gentle poke into the surface or by accidental pick-up of atoms during the scan.



**Fig. 3.** The STM topographic image (a) and its FFT (b) for the six-fold reconstruction pattern, and the STM topographic image (c) and its FFT (d) for the four-fold reconstruction pattern. The STM images ( $12\ \text{nm} \times 12\ \text{nm}$ ) are taken with functionalized tips in the same area, with a sample bias of  $-0.36\ \text{V}$  and tunneling current of  $15\ \text{pA}$  for (a), and  $-0.46\ \text{V}$  and  $15\ \text{pA}$  for (c).

High-resolution STM images further illustrate the details of both patterns at the atomic scale. As shown in Fig. 4(a), a triangular three-for-six pattern is resolved for the single-layer graphene. In such a structure, three carbon atoms in a hexagonal ring are lifted upward relative to the other three atoms, introducing some  $sp^3$  hybridization to the otherwise two-dimensional  $sp^2$  configurations. Previously, such

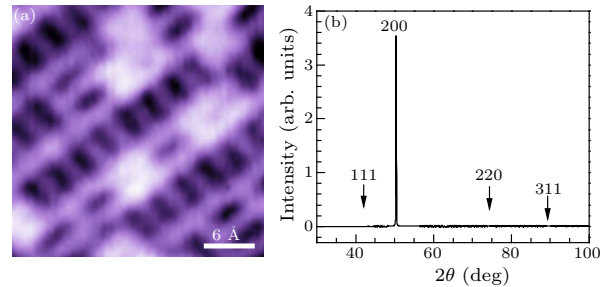
configuration change was observed only in a highly corrugated graphene surface or graphene on SiO<sub>2</sub> substrates,<sup>[17–19,23]</sup> and was attributed to the presence of strong spatially dependent perturbations such as local curvature and trapped charges. The sample surface in our study, however, is flat over 50 nm and no wrinkles or ripples are observed over a 1  $\mu$ m square area. Thus, the origin of the triangular three-for-six structure observed in our study is distinct from local curvature or trapped charges as reported in previous studies.



**Fig. 4.** (a) High-resolution STM image (3 nm  $\times$  3 nm) of the triangular three-for-six pattern of the graphene layer, (b) the  $dI/dV$  spectrum of the sample recorded in the single-layer graphene area, and (c) micro-Raman spectroscopy of the sample. The laser wavelength is 532 nm, and the background luminance from the copper substrate has been subtracted.

The emergence of  $sp^3$  hybridization and the three-for-six reconstruction are further illustrated by the energy-resolved scanning tunneling spectroscopy and micro-Raman spectroscopy. Figure 4(b) presents the  $dI/dV$  spectrum of the sample by scanning sample bias from  $-1$  V to  $1$  V relative to the tip. The Dirac point is located at about  $0.4$  eV below the Fermi energy, indicating the charge transfer from the substrate to the graphene. A depression of LDOS is observed at zero bias due to the lack of excitation of the optical phonon mode at a very low energy in the graphene.<sup>[12,15]</sup> By analyzing the second derivative spectrum, we find that the phonon energy is about  $69$  meV, which is close to the observation in Ref. [12].

In addition, a sharply increasing LDOS that is almost above zero bias is observed in the  $dI/dV$  curve as well. The exact nature of this state is not yet clear. Recent theoretical calculations predict that quasi-bound states localized around defects can induce a bump in LDOS above the Dirac point in graphene.<sup>[32]</sup> Wehling *et al.* stated that a midgap state appears as a universal feature of monovalent impurities bonded to graphene's carbon atoms.<sup>[31]</sup> The midgap states may also be formed due to vacancies, cracks, boundaries, or impurities in the substrate with a high potential difference with respect to the graphene sheet. On the other hand, the homogenous three-to-six pattern in Fig. 4(a) suggests that  $sp^3$  hybridization is not such a local effect as that induced by nitrogen doping.<sup>[14]</sup> Thus, one possible cause of such a state is the bonding between copper atoms in the reconstructed surface and carbon atoms in graphene. To determine the details of such a state, further experiments and theoretical calculation have to be carried out. Nonetheless, our tunneling spectroscopy suggests a significant interaction between the graphene single layer and the reconstructed copper surface, where the copper atoms underneath the graphene play a similar role to the defects in introducing  $sp^3$  hybridization and generating an electronic state near the Fermi level.



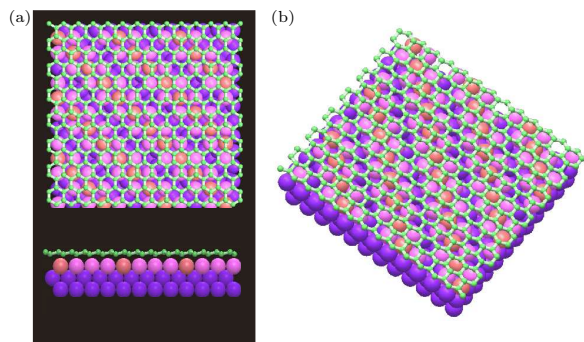
**Fig. 5.** (a) High-resolution STM image (3 nm  $\times$  3 nm) of the four-fold pattern, and (b) the X-ray diffraction pattern of the sample surface.

The existence of strong perturbation from the substrate is also suggested by the micro-Raman spectrum as shown in Fig. 4(c). A strong resonance peak of the D band at  $1347$   $\text{cm}^{-1}$  and a weak D' band at  $1620$   $\text{cm}^{-1}$  are observed, and these are usually associated with the presence of lattice distortion, defects, or edge states in single-layer graphene.<sup>[25,33]</sup> The measured high intensity of the D band, comparable to the intensity of the G band, is believed to be due to the three-for-six structure conformation of the graphene in our sample. The 2D band presents a very sharp single resonance peak with an intensity greater than the G resonance, which indicates the existence of a high-quality single-layer graphene.

Figure 5(a) shows a high-resolution STM image of the four-fold orthogonal pattern, which identifies the reconstruction with the aid of an x-ray diffraction pattern of the sample surface. The x-ray diffraction pattern, measured after the STM experiment,



shows a dominant crystal surface orientation along the [100] direction (Fig. 5(b)). It has been reported that copper foil crystallizes during the growth process in the CVD oven,<sup>[34]</sup> since a growth temperature of 1050°C is high enough to induce the crystallization, and graphene can play a significant role during the crystallization and surface reconstruction. Taking the fact that the distance between adjacent atoms on the Cu(100) surface is 2.55 Å, the observed orthogonal pattern with a lattice constant of  $3.6 \pm 0.05$  Å is then determined as a mixed  $(2\sqrt{2} \times \sqrt{2})R45^\circ$  reconstruction on the Cu(100) surface. A similar mixed reconstruction was observed with the chemisorption of oxygen on the Cu(100) crystal surface, as revealed by low-energy electron diffraction (LEED) and STM studies.<sup>[35,36]</sup> For the graphene-covered copper surface as in our study, the carbon atom in graphene is suggested to play a similar role as the chemisorbed oxygen in surface reconstruction, which results in a strong interaction between graphene and the copper substrate.



**Fig. 6.** Atomic model of the reconstructed graphene-copper system. The graphene is shown with green balls and sticks. The copper atom of the  $(2\sqrt{2} \times \sqrt{2})R45^\circ$  reconstruction is represented by red and pink balls, while the copper atoms under the top layer are blue.

An atomic schematic model was constructed based on the atomic-resolved information, as shown in Fig. 6. The upper part on the left is a view from the top, while the lower part is a view from the side and a 3D model is presented on the right. In the graphene overlayer, three atoms in a hexagonal ring are pulled towards the substrate to form the three-for-six configuration. The structure of the copper substrate is constructed following the missing-row model by Zeng *et al.*,<sup>[35]</sup> except that oxygen atoms are removed for clarity. The copper atom of the  $(2\sqrt{2} \times \sqrt{2})R45^\circ$  reconstruction is represented by red and pink balls. We are not able to exclude the existence of oxygen atoms in such a sample system, even though the oxide layer can be removed during annealing.<sup>[34]</sup>

We note that UHV annealing may play a significant role in enhancing the graphene-copper interaction and generating the observed novel structure pattern in our sample system. The Raman spectroscopy on the as-grown sample without UHV annealing shows no peak of D band resonance, indicating a weak perturbation from the substrate. Recent STM studies on similar graphene systems reported stripe and

rhombic patterns of graphene,<sup>[34,37]</sup> where no apparent three-for-six pattern was observed and a weak graphene-substrate interaction was concluded. The nature of various experimentally observed structures in the graphene-substrate system and the role of UHV annealing in modulating graphene-substrate interaction needs further investigation, with both experimental and theoretical calculations.

In summary, prominent interaction between single-layer graphene and a copper substrate has been demonstrated in a maze-like reconstruction pattern. The triangular three-for-six structure in graphene and a mixed  $(2\sqrt{2} \times \sqrt{2})R45^\circ$  reconstruction of the Cu(100) surface are revealed with functionalized STM tips. The electronic property of the graphene is modified as a result of partial  $sp^3$  hybridization. Besides the methods of introducing external impurity and internal structure defects, our results suggest another effective way to induce  $sp^3$  hybridization and thus to tune the electronic properties of a single-layer graphene through its interaction with the substrates.

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