The edge- and basal-plane-specific electrochemistry of a single-layer graphene sheet

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Graphene has a unique atom-thick two-dimensional structure and excellent properties, making it attractive for a variety of electrochemical applications, including electrosynthesis, electrochemical sensors or electrocatalysis, and energy conversion and storage. However, the electrochemistry of single-layer graphene has not yet been well understood, possibly due to the technical difficulties in handling individual graphene sheet. Here, we report the electrochemical behavior at single-layer graphene-based electrodes, comparing the basal plane of graphene to its edge. The graphene edge showed 4 orders of magnitude higher specific capacitance, much faster electron transfer rate and stronger electrocatalytic activity than those of graphene basal plane. A convergent diffusion effect was observed at the sub-nanometer thick graphene edge-electrode to accelerate the electrochemical reactions. Coupling with the high conductivity of a high-quality graphene basal plane, graphene edge is an ideal electrode for electrocatalysis and for the storage of capacitive charges.

Results

High-quality single-layer graphene sheets with lateral dimensions of several centimeter squares were synthesized by chemical vapor deposition (CVD) on a Cu substrate (Methods). In order to perform the edge- and basal plane-selective electrochemistry for a single-layer graphene sheet, we coated the graphene sheet region-specifically with...
Figure 1 | The configurations of plane- and edge-electrodes, and the structure of a CVD graphene sheet. (a) Schematic illustration of the edge-based (left) and basal plane-based (right) electrodes. (b) Raman spectrum of a monolayer CVD graphene sheet on a SiO2/Si substrate. (c) HR-TEM image of a monolayer CVD graphene sheet suspended over a micro grid; inset: a SAED pattern.

Discussion

The electrochemistry at either the edge- or plane-electrode was studied by cyclic voltammetry in an aqueous solution of 0.1 M phosphate buffered saline (PBS) containing 0.1 M KCl. This electrolyte was selected to eliminate the pH effect and obtain data that relevant to bio-systems. In the pure electrolyte, the cyclic voltammograms (CVs) for both electrodes are rectangular in shape, indicating a pure capacitive behavior (Figure 2). These CVs were performed in a narrow potential range to avoid unrequired side reactions. The areal specific capacitances of the edge-electrode was estimated to be about $1.0 \times 10^5 \text{ mF cm}^{-2}$ (the thickness of monolayer graphene was estimated to be 1 nm, see Methods section), and this value is over 4 orders of magnitude higher than that of plane-electrode (4 mF cm$^{-2}$).

The graphene edge was formed by mechanical cutting, leading to the formation of structural defects with dangling bonds. These dangling bonds are unstable upon the exposure to air or solution, and they can be possibly terminated by oxygen containing groups or adsorbing...
other reactive species. The high specific capacitance of the edge is mainly due to the accumulation of ions from the electrolyte at its structural defects to form electrochemical double layers.

The charge transfer processes at the edge- and plane-electrodes were further studied by using potassium ferricyanide, K₃Fe(CN)₆, as an electrochemical probe. Figure 3a and 3c give the CVs of 5 mM K₃Fe(CN)₆ in the same electrolyte and at different potential scan rates. The CV wave current density for edge- or plane-electrode increases linearly with the square root of scan rate (Figure 3b and 3d), indicating the electrochemical processes are controlled by the diffusion of electroactive species. For a linear diffusion-based system, the wave current density \( i_p \) can be given by Randles-Sevcik equation:

\[
i_p = 2.69 \times 10^{3} n^{3/2} A C D^{1/2} v^{1/2}
\]

(1)

where \( n \) is the number of electrons involved in the redox reaction (here, \( n = 1 \)), \( A \) is the area of the electrode, \( C \) is the concentration of the electrochemical probe, \( D \) is the diffusion coefficient and \( v \) is the scan rate. For the plane-electrode, the wave current density \( (j = i_p/A) \) was measured to be \( 2.2 \times 10^{-4} \text{ A cm}^{-2} \) at a scan rate of \( 5 \text{ mV s}^{-1} \). This current density is in good agreement with both the theoretically calculated value \((j = 2.6 \times 10^{-4} \text{ A cm}^{-2})\), and the wave current density acquired at a conventional glassy carbon (GC) electrode \((2.2 \times 10^{-4} \text{ A cm}^{-2}\); Figure S3). These results confirm that the electrochemical reaction on plane-electrode is controlled by linear diffusion, and the electrochemical property of graphene basal plane is similar to that of conventional GC. For the edge-electrode, the current density calculated from equation 1 is \( 2.6 \times 10^{-4} \text{ A cm}^{-2} \), and this value is about 400 times lower than the experimental CV wave current density \((0.11 \text{ A cm}^{-2})\). The edge-electrode is an ultramicrowave with a width around 1 mm and a length in millimeter scale; thus the diffusion process at its surface is likely different from that at the macroscopic plane-electrode. Actually, the CV curves recorded for the edge-electrode exhibit a quasi-sigmoidal shape even at a high scan rate of \( 100 \text{ mV s}^{-1} \) (Figure 3a). At a slow scan rate of \( 5 \text{ mV s}^{-1} \), a convergent diffusion mode is dominating (Inset of Figure 3a). Convergent diffusion resulted in faster mass transport and stronger wave current density. The steady current \( (i_s) \) for an ultramicrowave electrode can be approximately calculated by equation 2:

\[
i_s = 2 \pi n F D C L \left[1/\left(\ln 4 D t r^2/w^2\right)\right]
\]

(2)

where, \( L \) and \( w \) are the length and width of edge-electrode, respectively. The other terms have their usual electrochemical meanings (cf. equation 1). Accordingly, the steady current density \((j = i_s/Lw)\) was calculated to be \( 10 \text{ A cm}^{-2} \) at a scan rate of \( 5 \text{ mV s}^{-1} \), which is about 2 orders of magnitude higher than the experimental value \((j = 0.11 \text{ A cm}^{-2})\). A possible explanation for the theoretical overestimation given by equation 2 is the limitation of its diffusion law. For example, the flux in memory diffusion that based on the Fick’s first law is generated with a delay from the formation of concentration gradient. Furthermore, a fast electrochemical process can cause a delay of diffusion; the actual diffusion velocity is much slower than the theoretical value. As a result, the concentration of electrochemical probe at the surface of edge-electrode is much lower than that derived by using the theoretical mode of convergent diffusion. It was reported that such deviation was strongly dependent on the size of electrode. For an electrode with a diameter larger than \( 10 \mu \text{m} \), this effect is negligible. For an electrode with a diameter as small as \( 0.11 \mu \text{m} \), however, the experimental current density was more than 1 order of magnitude lower than the theoretical value. In our case, the edge-electrode is only one-atom thick, allowing only one ion to access each site of its surface in the through-thickness direction. The quasi-sigmoidal shaped CV curves (Figure 3a) together with the linear diffusion behavior (Figure 3b) reflect that the redox species diffuse to the surface of the edge-electrode from different directions in a semispherical profile, while only one direction is effective for electrochemical reaction. Nevertheless, the theoretical overestimation caused by a surface blockage effect and/or the adsorption of side products cannot be extensively ruled out.

In spite of the theoretical overestimation discussed above, the observed steady wave current density of ferricyanide at the edge-electrode is at least 2 orders of magnitude stronger than that measured at the plane-electrode (Figure 3), indicating a much faster electron transfer. Indeed, the heterogenous electron transfer rate can be accelerated by increasing the density of electronic states (DOS), relating to the amount of edge defects. The convergent diffusion also induced a faster mass transfer to contribute a stronger wave current density. Our findings have also been supported by a recent work on the electrochemistry at the nanopore made of single-layer CVD-graphene sandwiched between Al₂O₃ dielectric layers. In that case, the nanopores were fabricated by electron beam sculpting and the pore diameters were in the range of 5 to 20 nm. The electrode lengths equal to the pore circumferences; thus, they are in nanometer scale and significantly smaller than that of the edge-electrode used in this study (1–2 mm). As a result, the influence of convergent diffusion is more prominent, leading to a much stronger current density (on the order of \( 10^4 \text{ A cm}^{-2} \)). This high current density is also partly attributed to the high concentration of electrochemical probe (1 M KCl). In comparison with this previous study, our method is more conventional and simpler, avoiding the expensive and complicated lithography process.

The electrocatalytic activities of the edge- and plane-electrodes have been studied by using 3 electrochemical probes, including ascorbic acid (AA), beta-nicotinamide adenine dinucleotide (BNAAD).
(NADH) and oxygen. AA plays an important role in health care for human beings, especially essential to skin, connective tissues and immune systems. NADH is an important biomolecule that has wide applications in dehydrogenase-based biosensors. Figure 4a, b show the typical CV curves of 5 mM AA in 0.1 M PBS + 0.1 M NaCl electrolyte. At the edge-electrode, the CV exhibits an oxidation wave centered at ca. 0.30 V (Figure 4a), while the corresponding wave peak at the plane-electrode is positively shifted to 0.41 V (Figure 4b). The CV wave peak of NADH at the edge-electrode is located at ca. 0.44 V (Figure 4c). In comparison, the corresponding wave peak at the plane-electrode is also positively shifted to ca. 0.64 V (Figure 4d). It should be noted here that the peak current on the second CV cycle of NADH oxidation was observed to be much weaker than that on the first cycle (56% lower for the edge and 15% lower for the basal plane, Figure S4). This is possibly due to the adsorption of NADH, the oxidized product of NADH, on the electrode surfaces. The electrochemical results described above indicate that the graphene edge is much superior to its basal plane on the electrocatalytic oxidation of AA and NADH. It has been reported that the oxidation of AA was an inner sphere reaction and the electron transfer kinetics was highly sensitive to the surface properties of electrode. The surface of plane-electrode has a nearly perfect graphitic structure. The more defective sites and oxygen containing groups on the surface of edge-electrode greatly reduced the overpotential for AA oxidation. Furthermore, NADH ions are favor to adsorb on defects, especially those with carboxyl groups. The much stronger passivation of the edge-electrode caused by the adsorption of NADH ions further confirmed that the edge defects are responsible for the electrocatalytic activity of graphene.

In addition to the oxidation reactions of two biomolecules, AA and NADH, the oxygen reduction reaction (ORR) has also been studied at both the graphene edge- and plane-electrodes. In this case, the CVs were performed in 0.1 M KOH solution at a scan rate of 50 mV s⁻¹. Both electrodes showed featureless CV curves in N₂ saturated solutions within the potential range of −0.9 to 0 V (Figure 4e and 4f). After saturating the solutions with O₂, the CVs at both electrodes exhibited strong waves of ORR. The CVs of edge- and plane-electrodes did not have any obvious difference in their onset potentials (−0.22 V vs SCE), while a large difference in their peak potentials was observed. The CV of edge-electrode has an ORR peak centered at −0.48 V (Figure 4e), whereas that at the plane-electrode is negatively shifted to −0.71 V (Figure 4f). Density function theory (DFT) calculations have shown that dangling bonds at the edge of a graphene sheet have strong ability for the adsorption of O₂.
molecules were selectively adsorbed on defects and changed the graphene edge to a p-type semiconductor. As a result, the polarity of the edge is much higher than that of the basal plane. The high polarity along with the existence of oxygen containing groups at graphene edge make the edge-electrode more hydrophilic in alkaline condition, providing a stronger affinity to the electrolyte and dissolved oxygen. The performances of both electrodes on ORR also confirm that the edge of graphene possesses a much stronger electrocatalytic activity than that of graphene basal plane.

In summary, electrodes based on graphene edge and basal plane have been fabricated by region-specific coating CVD-grown mono-layer graphene sheet with epoxy resin. Electrochemical studies indicate that the specific capacitance of graphene edge is 4 orders of magnitude higher than that of graphene basal plane. Furthermore,
the CV waves of the graphene edge-electrode are quasi-sigmoidal in shape, following a convergent diffusion mode. However, the plot of steady current density versus square root of scan rate is linear, following a linear diffusion mode. The current density of the CV wave recorded at the edge-electrode is at least 2 orders of magnitude stronger than that recorded at the plane-electrode at a given scan rate because of a higher electron-transfer rate and the convergent diffusion at the ultramicronband edge-electrode. Graphene edge also has a much stronger electrocatalytic activity than graphene basal plane, not only on the oxidation reactions of AA and NADH but also on ORR. These findings demonstrate that the large basal plane of a monolayer graphene sheet is highly conductive while the graphene edge possesses superb capacitive and electrocatalytic properties.

Methods

Monolayer graphene sheets. The monolayer graphene sheets were grown on 25-μm thick Cu foils by CVD technique (Alfa Aesar, #13882) [1]. The details are described as follows. A Cu foil (cleaned by dilute acetic acid and acetone before use) was put into a tri-zone furnace (Lindberg Blue M HT55667C) with 1-inch quartz tube. After filling with H2 (20 scm at 0.30 Torr), the temperature in the furnace was increased from room temperature to 900 °C within 25 min and then to 1000 °C within another 10 min. After a 20 min-annealing process, a mixture of 10 scm CH4 and 5 scm H2 was introduced at 0.35 Torr for the growth of graphene sheet at 1000 °C for 30 min. Then the furnace was firstly cooled to 700 °C at a rate of 15 °C min−1 and finally spontaneously cooled down to room temperature. The gas flow was kept alive throughout the cooling process.

Preparation of the electrodes. 3% solution of PMMA with a molar mass of 996 K (Sigma Aldrich, #182265, dissolved in ethyl lactate) was spin-coated on a graphene/Cu sheet at 2000 rpm (runs per minute) for 60 s and dried at 170 °C for 2 min. Then, air plasma (Femto, Diener Electronics) was used to remove graphene on the back side of Cu foil. The edge-electrode was fabricated by coating the middle region of a graphene/Cu foil with ER (Nanjing Taitline Adhesive CO., LTD, China), leaving its two ending parts uncovered. Successively, the electrode was heated at 60 °C for 1 h to solidify ER. Thereafter one end of this electrode was immersed in a FeCl3 solution (aqueous solution of 0.1 g mL−1 FeCl3 + 0.1 mol L−1 HCl) to regionally etch away the Cu substrate and expose graphene sheet, and the other end was used for connecting potentialist for subsequent electrochemical measurements. After rinsed with distilled water for several times, the graphene end without Cu substrate was further coated with ER and cut vertically to expose its cross-section. The both corner regions of the exposed graphene edge were further coated with ER to leave only a narrow middle region (1.0–2.0 mm) of the graphene edge. The plane-electrode was prepared by partly coating the graphene/Cu sheet with ER, leaving one end free from the ER coating for electrical connection with potentialist and an ER-free window in its middle region. The Cu in the window was etched in FeCl3 solution, and rinsed with distilled water several times. Finally, the window wall was further coated with ER to avoid the percolation of electrolyte (Figure S1).

Surface areas of electrodes. The surface area of an electrode was calculated by the multiplication of its length and width. The length and width of a plane-electrode were measured by a vernier caliper or reading from its SEM image. The length of an edge-electrode was also accurately measured by these techniques. Its width is theoretically identical to the thickness of a single-layer graphene sheet (0.79 nm). Taking the wrinkled structure of graphene edge and its oxygenated groups into account, we evaluated the width of the edge-electrode to be around 1 nm.

Characterizations. All electrochemical tests were performed on an electrochemical analyzer (CHI 660D potentiostat-galvanostat, CH Instruments Inc.) at room temperature. The graphene electrode, a Pt foil and a SCE electrode were used as the working, counter and reference electrodes, respectively. These 3 electrodes were parallel positioned in the electrochemical cell (Figure S5). Raman spectra were recorded on a Renishaw Raman microscope (RM2000) with a 514-nm laser at a power density of 4.7 mW. Scanning electron micrographs (SEM) were performed on a field-emission scanning electron microscope (Sirion 200, Japan). High-resolution transmission electron microscope (HRTEM) images, selected area electron diffraction (SAED) pattern were taken by Tecnai G2 F20-Twin HRTEM. AFM image was taken out by using a SPM-9600 microscope (Shimadzu).

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Author contributions
G.S. and L.D. developed the concept, designed experiments and co-wrote the paper. W.Y. performed experiments, analysed data and wrote the paper. Y.Z., H.P. and Z.L. prepared the CVD graphene sheets. Y.L., C.L., H.P., J.Z. and Z.L. gave helps in the experiments and co-wrote the paper. All authors discussed the results and commented on the manuscript.

Additional information
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