Metal ion-directed solution-phase tailoring: from large-area graphene oxide into nanoscale pieces

Xiansong Wang, Peng Huang, Huiyang Liu, Chao Li, Guangxia Shen* and Daxiang Cui*

Abstract

Due to fascinating electronic properties and great potential in various applications, graphene has attracted great interest. Recently, much work have focused on the synthesis of different sizes and properties of graphene or graphene oxides (GOs), for example, graphene nanoribbons, nanosized graphene pieces, and nanosized triangular and hexagonal graphene sheets terminated by zigzag edges. Herein, we have demonstrated a widely available approach to fabricate the nanoscale GO pieces by directly solution-phase cutting a large-area GO sheet into nanoscale pieces via spontaneous redox reactions at room temperature. In this process, GO acts with dual functions as a model and a reducing reagent. With a typical example of silver ions, we have investigated in detail the influence of the reaction time and concentration of metal ions on yield and size of nanoscale GO pieces. Moreover, we also obtain Ag nanoparticle coating on the GO surface. Finally, a possible mechanism is suggested to explain the formation of nanoscale GO pieces.

Keywords: Graphene oxides, Nanoscale graphene oxide pieces, Spontaneous redox reaction, Metal particles

Background

Since discovered by Andre Geim and Konstantin Novoselov in 2004 [1], graphene has drawn significant attention to different scientific and technical communities due to its unique electrical, chemical, mechanical, optical, and structural properties [2]. However, large-area graphene remains to be a metallic conductor even at the neutrality point which limits its application in nanoelectronic devices and biological science [3-6]. In addition, for the purpose of drug delivery and biological nanoprobe applications, small-sized graphene or graphene oxides (GOs) can easily be swallowed into organs, tissues, and cells [7]. Recently, quite a lot of researchers have reported about the preparation of graphene ribbons with quantum confinement and edge effect properties by directly tailoring large-area graphene via e-beam lithography [8], hydrogen plasma etching [9], scanning tunneling microscope lithography [10], atomic force microscopy [11], chemical stripping, or catalytic tailoring (Fe, Ni, and Co nanoparticles as catalysts) [12-16]. Usually, the technologies used for synthesis of graphene ribbons mostly must be operated under ultrahigh-vacuum and high-energy conditions. So it is very difficult to produce large quantities of water-soluble graphene pieces. Moreover, these extreme synthetic conditions will be ultimately bound to affect the properties of graphene ribbon. Till now, direct soluble-phase formation of nanoscale graphene or graphene oxide pieces has been rarely involved [17]. Generally, through selecting small-sized graphite as raw materials to control the size of GO during the synthesis of GO through the Hummers procedure, subsequently complicated treatment with strong sonication treatment and stepwise centrifugation at 4,000 to 10,000 rpm, a small-sized GO can be obtained [18]. However, the procedures are quite complex and the yield of nanoscale fragments is also very low.

Herein, we report a widely available approach to prepare the nanoscale GO pieces directly utilizing some oxidizing metallic ions (Ag+, Ni2+, Co2+, etc.) via spontaneous redox reactions to cut a large-area GO sheet into nanoscale pieces at room temperature. With an example of silver ions, we have investigated the influence of the reaction time and concentration of metal ions on size and properties of nanoscale GO pieces. Meanwhile, the corresponding silver nanoparticles can also be obtained. Finally, a possible mechanism is put forward for explaining the formation of nanoscale GO pieces.
Methods

Chemicals
All reagents were of analytical grade and purchased from Shanghai Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Natural graphite powder (800 mesh) was provided by Beijing Chemical Reagents (Beijing, China). All aqueous solutions were prepared with ultrapure water (18 MΩ cm).

Preparation of large-area GO
Water-soluble GO was prepared by oxidizing graphite according to a modified Hummers method just as our previous reports [19,20]. Briefly, the graphite powder was first oxidized into graphite oxide using KMnO₄/H₂SO₄, and then the graphite oxide was exfoliated into GO sheets in water under ultrasonication for 1 h, followed by centrifugation at 4,000 rpm for 30 min and dispersion in water. The obtained yellow-brown aqueous suspension of GO was stored at room temperature for further characterization and subsequent reaction.

Preparation of nanoscale GO pieces
The experiments of cutting large-area GO were carried out as follows: Firstly, 100-mL GO water solution (0.50 mg/mL) was prepared. Homogeneous suspension (20 mL) of GO was mixed with the desired amount of aqueous metallic ion (Ag⁺, Ni²⁺, Co²⁺, etc.) solution (5 mg/mL). Without heating or ultrasonication, the reaction mixtures were kept at room temperature for 48 h. Then the mixtures were centrifuged to remove the nanoparticles and large-scale GO and particle composites at the rate of 8,000 rpm. The upper solution without further purification was detected by atomic force microscopy (AFM), Fourier transform infrared (FTIR) spectroscopy, UV-vision (UV-vis) spectroscopy, and X-ray photoelectron spectroscopy (XPS). In order to investigate the tailoring mechanism, we selected silver ions as a typical example and elaborately investigate the influence of reaction time and concentration of silver ions on the size and properties of nanoscale GO. All experiments were carried out at 25°C ± 2°C.

Characterization of nanoscale GO
AFM images were obtained on a Nanoscope MultiMode V scanning probe microscopy system (Veeco, Plainview, NY, USA) by tapping-mode imaging. Commercially available AFM cantilever probes with a force constant of approximately 48 N/m and resonance vibration frequency of approximately 330 kHz were used. The scanning rate was usually set at 1 to 1.2 Hz. Freshly cleaved mica with atom-level smoothness was used as the substrates. The samples were coated on the mica surface by spin-coating technology. UV-vis spectra were measured at 20°C with a Shimadzu UV-2450 spectrophotometer equipped with a 10-mm quartz cell (Kyoto, Japan). Zeta potentials were measured with NICOMP 380 ZLS Zeta Potential/Particle Size Analyzer. The XPS measurements were performed on an Axis Ultra DLD XPS (Kratos Analytical, Manchester, UK) using a monochromated Al Kα (1,486.6 eV) source at 15 kV. Scanning electron microscopy (SEM) images were taken on a ZEISS-ULTRA 55 SEM (Oberkochen, Germany) equipped with an X-ray energy-dispersive spectroscopy (EDS) at an accelerating voltage of 20 kV (provided in Additional file 1). In addition, the conductive properties of the nanoscale GO film coated on the mica surface were tested using a conductive AFM. The detailed process and results have been given in Additional file 1.

Figure 1 Tapping-mode AFM images of GO and nanoscale GO pieces. (a) GO, (b) Ag⁺, (c) Co²⁺, and (d) Ni²⁺ and corresponding profile analysis: (e) GO, (f) Ag⁺, (g) Co²⁺, and (h) Ni²⁺.
Results and discussion
Tailoring large-area GO by different metal ions
Graphene oxide is very widely generated using natural graphite powder through the Hummers method. The chemically derived GO is soluble in pure water due to hydrophilic functional groups, e.g., carboxyl, hydroxyl, and epoxide groups on the surface [16,21]. Figure 1a shows the AFM image of GO with atom-level smoothness and the sizes in the range of 1 to 10 μm. The height profile of the AFM image in Figure 1e is approximately 1 nm, which is consistent with the data reported in the literature, indicating the formation of a single-layered GO. Figure 1b,c,d depicts that the nanoscale GO pieces with different sizes were tailored utilizing three kinds of metal ions (Ag+, Ni2+, Co2+), respectively. Corresponding profile analysis of these AFM height images (Figure 1f,g,h) has given heights of approximately 1 nm, which were elementally consistent with the thickness of GO. Similarly, in the addition of Ag⁺ ion system, some nanoparticles have been found to be dispersed in the solution or attached on the GO surfaces similar to what we have reported previously [22]. In our previous work, we mainly focused on the synthesis of silver-GO composites. When testing the samples by AFM, some little pieces were occasionally detected in the high-resolution images, which were neglected as contamination before [22]. Thereafter, in order to investigate the tailoring mechanism, we selected the other weak oxidation of metal ions, such as Ni²⁺ and Co²⁺, and obtained results similar to the information given previously. In addition, XPS data have been provided in Additional file 1: Figure S1.

Tailoring large-area GO by silver ions
For silver ions, a series of systematic experiments have been carried out. In a typical experiment, 0.50 mg/mL of an aqueous GO dispersion (10 mL) was added to 10 mM aqueous AgNO₃ solution (10 mL). As shown in Figure 2a, the large-area GO has been tailored into small fragments after the reaction was kept for approximately 12 h. TEM image and EDS data were given in Additional file 1: Figure S2. They provide the presence of Ag, C, and O elements, which should be attributed to GO and the silicon substrate. Furthermore, the sizes of the little pieces do not become smaller even if the reaction time is beyond 48 h. Meanwhile, there were two kinds of nanoscale GO existing in the mixture: one is the pure nanoscale GO pieces in Figure 2b, and the other is the silver-GO composite pieces in Figure 2c. In addition, the nanoscale...

Figure 2 Tapping-mode AFM image of nanoscale GO pieces using 0.5 mM silver ions for 24 h. a) nanoscale GO, b) and c) the high-resolution images of the labeled area in a.

Figure 3 UV-vis absorption and FTIR spectra of nanoscale GO. a) UV-vis change with reacting time, b) UV-vis change with adding Ag⁺ concentration, c) FTIR spectra of nanoscale GO by adding 0.5mM Ag⁺ after reacting 12h.
GO film cannot be conductive using C-AFM testing (see Additional file 1: Figure S3).

Influence of the reaction time on the sizes and properties of nanoscale GO pieces was monitored by UV-vis spectroscopy (Figure 3a). The UV-vis spectra of GO display two characteristic peaks at 230 and 303 nm, corresponding to $\pi \rightarrow \pi^*$ transition of aromatic C-C bond and $n \rightarrow \pi^*$ transition of C=O bond, respectively [23]. From Figure 2d, it can be found that the two characteristic peaks of GO red-shift to approximately 250 and approximately 310 nm after adding 0.5 mM Ag$^+$ ions into the GO solution for 0.5 h, due to the interaction of GO and silver ions. The peak intensities decayed gradually with prolonged reaction time. Especially the peak intensity in the region approximately 310 nm decreases dramatically after 48 h, providing a first hint that some functional groups in GO may decrease [24]. Similar results can be further achieved by changing the concentration of Ag$^+$ ions in Figure 3b. We can find that there is a distinct difference in wavelengths and intensities of the characteristic peaks of GO with the different concentrations of Ag$^+$ ions in the system after approximately 24 h. At lower concentration, the signal at 310 nm nearly disappears and that at 250 nm becomes distinct, which may mean that the Ag$^+$ ions preferentially attack the sites of $sp^3$ carbon clusters or defective regions on the basal planes and partially restore the $sp^2$ carbon framework. When a higher proportion of Ag$^+$ ions (5 or 0.5 mM) are added into the reaction system, the peak intensity (at approximately 310 nm) of GO seems to be obvious and accompanies a larger red shift with increasing Ag$^+$ ion concentration, gradually close to 360 nm which is for silver plasmon absorption bands [24]. It can be explained that the number of silver nanoparticles fabricated on the GO surface or solution becomes large with the increasing proportion of Ag$^+$ ions in the mixture, which also provides more change for the interaction of Ag nanoparticles and GO. At the same time, we also find that even if the Ag$^+$ concentration is increased to 5 mM, there still exists some nanoscale GO with smooth edges in the mixture.

FTIR spectroscopy has been considered as another powerful tool to analyze surface chemical group changes of GO. As shown in Figure 3c, the characteristic peaks of GO (green line) displayed the C=O stretching vibration peak at 1,730 cm$^{-1}$, the vibration and deformation peaks of O-H groups at 3,428 and 1,415 cm$^{-1}$, respectively, the C-O (epoxy groups) stretching vibration peak at 1,220 cm$^{-1}$, and the C-O (alkoxy groups) stretching peak at 1,052 cm$^{-1}$ [25]. After the reaction is conducted for 48 h (red line), the intensities of the FTIR peaks corresponding to the C-O (epoxy groups) stretching vibration peak at 1,220 cm$^{-1}$ and the C-O (alkoxy groups) stretching peak at 1,052 cm$^{-1}$ [25]. After the reaction is conducted for 48 h (red line), the intensities of the FTIR peaks corresponding to the C-O (alkoxy groups) stretching vibration peak at 1,220 cm$^{-1}$ and the C-O (epoxy groups) stretching vibration peak at 1,052 cm$^{-1}$ disappeared nearly, the
C=O stretching vibration peak at 1,730 cm$^{-1}$ decreased dramatically, and the vibration and deformation peaks of O-H groups at 3,428 and 1,415 cm$^{-1}$, respectively, and the C-O (alkoxy groups) stretching peak at 1,052 cm$^{-1}$ increased slightly. These results further confirmed that some active functionalities (epoxide groups) in GO have been removed.

The mechanisms of tailoring GO

Since the appearance of GO, the determination of GO structure has been challenging because of its nonstoichiometric chemical composition, which depends on the synthesis method and the degree of reduction, and the oxygen functional groups in GO have been identified by various kinds of techniques. It is generally agreed that oxygen is present in GO mostly in the form of hydroxyl and epoxide groups on the basal plane, whereas smaller amounts of carboxyl, carbonyl, phenol, lactone, and quinone are present primarily at the sheet edges. The existence of the chemical groups confers new properties on GO such as the perfect monodispersity in water and weak reducibility. Based on the above facts and our experimental results, a probable mechanism is put forward as given in the schematic diagram (Figure 4). Firstly, part of Ag$^+$ ions is preferentially absorbed on the sites of carboxylic groups at the edges of GO by the electrostatic interaction. Then Ag$^+$ ions bonded on GO or freely dispersing in the solution further encounter the reducing groups (e.g., epoxy groups) on the basal plane of other GO sheets. Thus, Ag$^+$ ions themselves are reduced to Ag and then generate Ag nanoparticles; meanwhile, the carbon-carbon skeleton is broken which directly leads to the cutting of GO into little pieces.

Although the feasibility conclusion has been verified through analysis results of UV-vis and FTIR data, we also elaborately investigated the chemical state change of carbon in GO by XPS technology. Figure 5a shows the C$_{1s}$ XPS of GO sheets. There are four different peaks detected that centered at 284.5, 288.4, 293.8, and 296.6 eV, corresponding to C=C/C-C in aromatic rings, C-O (epoxide and alkoxy), C=O, and COOH groups, respectively [26]. After adding Ag$^+$ ions into solution for 48 h, the distinct changes of C$_{1s}$ XPS are detected in Figure 5b. The intensities of C$_{1s}$ peaks of the carbons binding to oxygen, especially the peak of C-O (epoxide and alkoxy), decreased dramatically, which reveals that most epoxide groups reacted with Ag$^+$ ions. Moreover, the diameters and charges of metal ions may have great influence on the sizes and properties of nanoscale GO which will be further confirmed by subsequent work.

Conclusions

In summary, we have demonstrated a very simple strategy to obtain nanoscale GO pieces using metal ions as oxidation reagent at mild condition. Without being heated or treated ultrasonically, two kinds of nanoscale GO pieces: GO pieces and nanoparticle-coated GO piece composites, are obtained. Based on systematic investigations of nanoscale GO piece formation by the addition of Ag$^+$ ions as a tailoring reagent, a probable mechanism is suggested to explain the formation of nanoscale GO pieces, which can be mainly attributed to interaction of metal ions (Ag$^+$, Co$^{2+}$, Ni$^{2+}$, etc.) with the reducing groups (e.g., epoxy groups) on the basal plane of other GO sheets. Obviously, in this progress a large-scale GO acts with dual functions, as a reducing reagent and a nucleation site of metal or metal oxide nanoparticles. This work provides a good way or chance to fabricate nanoscale GO pieces and GO composites in water solution and more widely apply in nanoelectronic devices, biosensors, and biomedicine.
9. Yang R, Zhang LC, Wang Y, Shi ZW, Shi DX, Gao HJ, Wang EG, Zhang GY: An anisotropic etching effect in the graphene basal plane. *Adv Mater* 2010, 22(36):4014–4019.

10. Wong HS, Durkan C, Chandrasekhar N: Tailoring the local interaction between graphene layers in graphite at the atomic scale and above using scanning tunneling microscopy. *ACS Nano* 2009, 3(1):3655–3662.

11. Lu G, Zhou XZ, Li H, Yin ZY, Li B, Huang L, Boey F, Zhang H: Nanolithography of single-layer graphene oxide films by atomic force microscopy. *Langmuir* 2010, 26(6):6164–6166.

12. Tsukamoto T, Ogino T: Control of graphene etching by atomic structures of the supporting substrate surfaces. *J Phys Chem C* 2011, 115(17):8580–8585.

13. Gao L, Ren W, Liu B, Wu ZS, Jiang C, Cheng HM: Crystallographic tailoring of graphene by nonmetal SiO_x nanoparticles. *J Am Chem Soc* 2009, 131(39):13934–13936.

14. Campos LC, Manfrinato VR, Sanchez-Yamagishi JD, Kong J, Jarillo-Herrero P: Anisotropic etching and nanoribbon formation in single-layer graphene. *Nano Lett* 2009, 9(7):2600–2604.

15. Datta SS, Strachan DR, Khamis SM, Johnson ATC: Crystallographic etching of few-layer graphene. *Nano Lett* 2008, 8(7):1912–1915.

16. Li Z, Zhang W, Luo Y, Yang J, Hou JG: How graphene is cut upon oxidation? *J Am Chem Soc* 2009, 131(18):6320–6321.

17. Pan DY, Zhang JC, Li Z, Wu MH: Hydrothermal route for cutting graphene sheets into blue-luminescent graphene quantum dots. *Adv Mater* 2010, 22(6):734–738.

18. Wang XL, Bai H, Shi GQ: Size fractionation of graphene oxide sheets by pH-assisted selective sedimentation. *J Am Chem Soc* 2011, 133(16):6338–6342.

19. Zhang J, Yang H, Shen G, Cheng P, Zhang J, Guo S: Reduction of graphene oxide via L-ascorbic acid. *Chem Comm* 2010, 46:1112–1114.

20. Zhang J, Shen G, Wang W, Zhou X, Guo S: Individual nanocomposite sheets of chemically reduced graphene oxide and poly(N-vinyl pyrrolidone): preparation and humidity sensing characteristics. *J Mater Chem* 2010, 20:10824–10828.

21. Eda G, Chhowalla M: Chemically derived graphene oxide: towards large-area thin-film electronics and optoelectronics. *Adv Mater* 2010, 22:2392–2415.

22. Wang X, Huang P, Feng L, He M, Guo S, Shen G, Cui D: Green controllable synthesis of silver nanomaterials on graphene oxide sheets via spontaneous reduction. *RSC Adv* 2012, 2:3816–3822.

23. Eda G, Lin YY, Mattevi C, Yamaguchi H, Chen HA, Chen J, Chen CW, Chhowalla M: Blue photoluminescence from chemically derived graphene oxide. *Adv Mater* 2010, 22(4):505–509.

24. Cassagneau T, Fendler JH: Preparation and layer-by-layer self-assembly of silver nanoparticles capped by graphite oxide nanosheets. *J Phys Chem B* 1999, 103(11):1789–1793.

25. Si Y, Samulski ET: Synthesis of water soluble graphene. *Nano Lett* 2008, 8(6):1679–1682.

26. Diederich DR, Park S, Bialkowski CW, Ruoff RS: The chemistry of graphene oxide. *Chem Soc Rev* 2009, 39(1):228–240.