Solution-processable graphene nanomeshes with controlled pore structures

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Graphene nanomeshes (GNMs) which can be cheaply produced on a large scale and processed through wet approaches are important materials for various applications, including catalysis, composites, sensors and energy related systems. Here, we report a method for large scale preparation of GNMs by refluxing reduced graphene oxide sheets in concentrated nitric acid solution (e.g., 8 moles per liter). The diameters of nanopores in GNM sheets can be readily modulated from several to hundreds nanometers by varying the time of acid treatment. The porous structure increased the specific surface areas of GNMs and the transmittances of GNM-based thin films. Furthermore, GNMs have large number of carboxyl groups at the edges of their nanopores, leading to good dispersibility in aqueous media and strong peroxidase-like catalytic activity.

Over the last several years, extensive work has been devoted to the syntheses and applications of graphene due to the excellent electrical, optical, mechanical and thermal properties associated with its unique atom-thick two-dimensional structure1–7. The most widely-used method to prepare graphene on a large scale at low cost is the reduction of graphene oxide (GO)8–12. However, the reduced graphene oxide (rGO) is insoluble, infusible and intractable8–14. The poor processability of neutral rGO precludes the possibility of using conventional material processing techniques to fabricate macroscopic rGO-based materials into desired shapes or structures. Furthermore, rGO sheets are normally composed of about 60% intact graphene islands of variable sizes interspersed with planar quasi-amorphous sp2-bonded defect clusters, which predominantly incorporate a large number of carbon pentagons, heptagons, and hexagons15. The defect clusters of rGO sheets can be selectively removed to produce GNMs with more perfect graphene basal planes.

It is highly expected that the preparation of GNM to open the band gap of graphene16,17 and increase its specific surface area (SSA) and transparency18,19, resulting in the formation of a class of new fascinating materials for various applications in graphene-based composites, catalysis, sensors and energy related systems20–22. Up to now, high-quality single-layered GNMs have been prepared only by plasma or photocatalytic oxidation of graphene sheets using solid templates16–18, while a bulk porous graphene material was obtained by chemical vapor deposition23. Unfortunately, only limited amounts of single-layered GNMs can be produced by these methods at high costs, and the resulting GNMs are not solution-processable, for example, by spin- or dip-coating, filtration and printing. Herein, we report a one-step solution method for the large scale production of GNMs by selectively removing the defect clusters of rGO sheets with nitric acid. This technique is readily scalable and conceptually different from those reported for preparing holey graphene sheets16–23. These GNM sheets have uniformly distributed nanopores, ranging from several to hundreds nanometers in diameters, and can be stably dispersed in aqueous media, including pure water.

Results

In a typical experiment, GNMs were prepared by refluxing rGO sheets in 8 mol L−1 nitric acid solution for 4, 6, 9 or 11 hours (see Supplementary Information for details). They are designated as GNM-n, where n refers to the time used for acid treatment (tₐ). Similar technique has previously been used to purify single-walled carbon nanotubes (SWCNTs), removing the residual catalyst and amorphous carbons, at the expense of a significant destruction of SWCNTs24. In this study, we found that the defect clusters of rGO sheets can be selectively removed by the treatment with nitric acid to leave nanopores in the basal planes of rGO sheets due to the relatively higher chemical activities intrinsically associated with the defect clusters. For instance, the GNM sheets prepared by
treating rGO sheets with nitric acid for 6 h (GNM-6) show nanopores with diameters in the range of 0–90 nm (ca. 90% of them ≤ 60 nm) and the pores are uniformly distributed over the whole GNM sheet (Figure 1a, Figure S1). The total area of nanopores within 1.00 μm² GNM-6 sheet was estimated to be about 0.46 μm². The lateral dimensions of GNM-sheets are in the range of 3–5 μm, which are more uniform than those of rGO precursors (0.5–7 μm, Figure S2). The thickness of a GNM-6 sheet is around 1.0 nm according to the height profile of AFM image (Figure 1a), indicating a single-layer nature of rGO sheet 20. High-resolution transmission electron microscope (HR-TEM) images of GNM-6 clearly demonstrate the nanoporous structure, as exemplified by Figure 1b showing a typical image of several nanopores. The nanopores with irregular shapes are surrounded by a graphene basal plane. The distances between neighbouring nanopores are typically in the range of several nanometers to about 50 nm. The selected area electron diffraction (SAED) pattern of the graphene basal plane shows a sharp hexagonal diffraction dots (Figure 1c), indicating a perfect crystalline structure with regular carbon framework. In comparison, the SAED pattern of a rGO sheet usually exhibits a halo ring arising from its defect clusters 25. These results confirm that the defect clusters of rGO have been, at least, partially removed by the treatment with nitric acid.

**Discussion**

To reveal the mechanism for the nanopore formation in GNMs, we followed the morphological changes of GNMs with the time of nitric acid treatment. As shown in Figure 2, the number density of nanopores in GNMs decreased from 55 ± 6 μm⁻² through 26 ± 2 to 15 ± 2 μm⁻² with the tₜ increased from 4 through 6 to 9 h. In the meantime, the total area of nanopores within 1.00 μm² of GNM sheet increased from 0.10 through 0.46 to 0.63 μm². At tₜ = 11 h, the GNM sheets were broken up into tiny sheets. The pore sizes of GNMs also increased with the increase of tₜ (Figure S1). For example, the pore diameters of GNM-4 are mainly in the range of 10 to 40 nm (>85%), while those of GNM-9 are mostly larger than 40 nm (>80%). Accordingly, the size and the number density of pores in GNM sheets can be regulated simply by varying tₜ.

The selective removal of the defect clusters in rGO sheets upon the acid treatment was further confirmed by X-ray photoelectron spectroscopy (XPS) and Raman measurements. As can be seen from Figure 3a, the acid-treatment-induced conversion of rGO to GNM reduced the intensities of the peak components associated with C-N, C-O (hydroxyl and epoxide) groups in the C 1s XPS spectra and enhanced the –C=O and –COOH peak intensities significantly 26. The atomic contents of the carbon atoms related to –C=O and –COOH groups were calculated to be 4.63, 7.24, 7.51 and 8.26% for rGO, GNM-6, GNM-9 and GNM-11, respectively. These XPS results reflect that the defect clusters of rGO sheets were partially removed to form the nanopores with defective edges; because it has been previously reported that the C-N, hydroxyl and epoxide groups of rGO sheets are predominantly connected to the defect clusters on their basal planes while the –C=O and –COOH groups are mainly located at their edges 27,28.

The Raman spectrum for each of the GNMs shows a disorder-induced band (D-band) in the range 1300–1370 cm⁻¹ and a first-order-allowed band (G-band) in the region of 1580–1600 cm⁻¹ (Figure 3b)25. The intensity ratio of D/G bands for GNM-6, GNM-9 and GNM-11 were measured to be 0.93, 0.91 and 0.88, respectively. These values are much lower than the corresponding value for rGO (1.11), implying, once again, the removal of the quasi-amorphous sp²-bonded carbons of rGO by the treatment with nitric acid. The
disorder carbons of GNMs are attributed to mainly those carbon atoms at the edges of the nanopores, as indicated by the XPS spectra (Figure 3a).

Owing to the presence of –C=O and –COOH groups at the nanopore edges, the GNMs can form stable dispersions in pure water (Figure 4a). The ionization of carboxy groups enhanced the repulsion force between graphene sheets to increase the solubilities of GNMs, as confirmed by Zeta potential measurements. It is known that a rGO dispersion is stable when its Zeta potential is > –30 mV. In pure water (pH = 6.4), the Zeta potentials of GNM dispersions are in the range of –36 to –46 mV, and hence they are stable (Figure 4b). Indeed, a 2 mg mL⁻¹ dispersion of GNM-6 in pure water did not show participation even after keeping at room temperature for over 1 week (Figure 4a). In contrast, rGO sheets cannot be stably dispersed in the same medium because of their relatively higher Zeta potential (> –30 mV at pH = 6.4, Figure 4b).

To test the solution-processibility of GNMs, we filtrated their stable aqueous dispersions (1 mg mL⁻¹) through porous polytetrafluoroethylene (PTFE) membranes to prepare GNM films (Figure S3). The X-ray diffraction pattern for each of the resultant GNM films shows a peak at 2θ = 24.39°, which is higher than that of the corresponding peak for an rGO film (2θ = 23.22°) prepared under the same condition (Figure 4c). This result indicates that GNM sheets can be assembled into more compact “paper” with a d-space (3.65 Å) smaller than that of an rGO “paper” (3.83 Å). Furthermore, the full width at half-maximum (FWHM) values of the XRD peaks for GNM-6, GNM-9 and GNM-11 papers were measured to be 2.71°, 2.55° and 2.41°, respectively, all being much narrower than that of the rGO paper (4.20°). These results reveal that the graphene sheets in GNM papers have more ordered stacking structures than that in an rGO paper. The cross-section SEM images of the films also indicate that the GNM-6 paper has a more compact stacking structure than that of the rGO paper (Figure S4). Therefore, the partial removal of oxygenated groups from the basal planes of rGO sheets decreased the d-spaces between graphene sheets, facilitating the formation of more ordered GNM films.

In view of the above mechanism and structural studies, we further tested the nanoporous structure of GNMs for optical transparency.
enhancement. The transmittances of thin GNM-6 films with different thicknesses deposited on poly(ethylene terephthalate) substrates are plotted in Figure 5a. As can be seen, the transmittance of a GNM-6 film is much higher than that of an rGO film at any given thickness investigated in the present work. For example, a 21-nm thick GNM film has a transmittance of 85% at 550 nm (Figure 5a), but only 74% for its rGO counterpart. Due to its porous structure, the sheet resistance of a GNM-6 film is about one order of magnitude higher than that of a rGO film with the same thickness (Figure S5). However, thermal treatment of the GNM-6 film at 400°C in Ar for 3 h reduced its sheet resistance to a value comparable to that of the rGO film with negligible change in its transparency. More importantly, the nanoporous structure increased the specific surface area (SSAs) of GNMs significantly. The SSAs of GNM-4, GNM-6, GNM-9 and GNM-11 were measured by a standard methylene blue adsorption method to be 810 ± 64, 970 ± 60, 1010 ± 53 m² g⁻¹ (average of 3 samples), respectively. These values are much larger than that of rGO (595 ± 75 m² g⁻¹)³¹.

The high content of –COOH groups and large SSAs of the GNMs make them promising for replacing intrinsic peroxidase catalysts for certain bioreactions or biosensors. As a demonstration, we used GNM-6 or GNM-11 to catalyze the reaction of a peroxidase substrate, 3,3,5,5-tetramethyl benzidine (TMB), in the presence of H₂O₂ to produce a blue color reaction. The absorbance at 652 nm of the reaction system with a constant amount of catalyst is proportional to the reaction conversion or the catalytic activity of the catalyst (Figure 5b). Actually, the reaction conversion for the system with the GNM-6 (GNM-11) peroxidase catalyst is about 4.2 (5.0) times that of the widely-used horse horseradish peroxidase (HRP) and 2.1 (2.5) times that of GO after carrying out the reaction at room temperature for 12 h. This result reflects the unusually high catalytic activities of the newly-developed GNM peroxidase catalysts.

In conclusion, we have developed a cheap, convenient and effective chemical approach for large-scale synthesis of GNMs by refluxing rGO sheets in concentrated nitric acid, which can be readily scalable to industry levels. The porous structures of GNMs can be well controlled simply by regulating the reaction time. Due to its nanoporous nature, the resultant GNMs show high specific surface areas, improved optical transparencies and good solution-processability, making them attractive for various potential applications, including catalysis, energy storage, sensors and environmental protections.

Methods
GO was prepared by the oxidation of natural graphite powder (325 mesh) according to a modified Hummers’ method (see Supplementary Information for details)²⁹,³³,³⁴. rGO was synthesized by reducing the aqueous dispersion of GO with hydrazine hydrate according to the method reported in literature (see Supplementary Information for details)²⁹. GNMs were prepared by mixing 50 mL rGO dispersion (0.1 mg mL⁻¹) with a certain amount of HNO₃ solution to form a mixture containing...
were recorded by using a D8 Advance X-Ray diffractometer with Cu-Kα radiation. The XRD patterns were recorded on the same instrument. The electrical conductivities of rGO or GNM films were measured using a four-probe conductivity test meter (KDY-1, Kunde Technology, Co., Ltd.) at room temperature. The transmittances of the rGO or GNM films were measured by the use of a zeta sizer nanosystem (Malvern Instruments). The HR-TEM spectra were recorded on a RM 2000 Microscopic confocal Raman spectrometer (Renishaw PLC, England) using a 514 nm laser beam. The electrical conductivities of rGO or GNM films were measured using a four-probe conductivity test meter (KDY-1, Kunde Technology, Co., Ltd.) at room temperature. The transmittances of the rGO or GNM films at 550 nm and the UV-visible spectra of the catalytic reaction systems were measured by the use of a U-3010 UV-visible spectrometer (Hitachi, Japan). The zeta potentials of rGO and GNMs aqueous dispersions with various pH values were measured by the use of an electrochemically reduced graphene oxide for ac line-filtering. Adv. Mater. 22, 2206–2210 (2010).

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Author contributions

G.S. and L.D. conceived and designed the experiments. X.W. performed the experiments and analysed the data. L.J., K.S. and C.L. provided helps in the experiments. X.W., G.S., L.D. and L.J. wrote the manuscript. All authors discussed the results and commented on the manuscript.

Additional information

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