Large-Area Graphene Films by Simple Solution Casting of Edge-Selectively Functionalized Graphite

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ABSTRACT We report edge-selective functionalization of graphite (EFG) for the production of large-area uniform graphene films by simply solution-casting EFG dispersions in dichloromethane on silicon oxide substrates, followed by annealing. The resultant graphene films show ambipolar transport properties with sheet resistances of 0.52–3.11 kΩ/sq at 63–90% optical transmittance. EFG allows solution processing methods for the scalable production of electrically conductive, optically transparent, and mechanically robust flexible graphene films for use in practice.

KEYWORDS: edge-selective functionalization · graphene film · solution processing · annealing · sheet resistance · optical transmittance

Exfoliation of graphite, including into individual layers (graphene), is of great interest due to the exceptional mechanical strength, thermal conductivity, and electronic properties characteristics of graphene platelets. As a result, graphene is attractive for a wide range of potential applications in composites, energy related systems, sensors, and others. Graphene is a promising material for replacing indium tin oxide (ITO) in transparent conductive flat panels. Currently, ITO is widely used as the transparent electrode in flat panel displays, because it has a relatively high electrical conductivity and optical transmittance with a very good stability to moisture and harsh environmental conditions. However, indium is expensive and has a limited reserve in nature. In addition, ITO is not appropriate for flexible display devices due to limitations in its mechanical and optical properties. Recently, Hong et al. have demonstrated the feasibility for the use of large-area graphene films as the transparent conductor in flexible polymer film-based flat panel electronics. These graphene films were grown on nickel (Ni) and copper (Cu) substrates at 1000 °C under a reduced pressure by chemical vapor deposition (CVD) process and then transferred onto polymer films. Unfortunately, these methods are not sufficiently cost and time effective to be commercially viable for mass production. To meet commercial demands, new materials or processing methods would be essential. In this regard, graphene oxide (GO) has been investigated as a chemical alternative for mass production of graphene films by solution-based processes. However, GO has poor quality and requires reduction, which has a limited conversion to reduced graphene oxide (rGO). Hence, the original graphene structure from GO cannot be efficiently restored.

Here, we report the formation of large-area, flexible, conductive, and transparent graphene films via a simple solution casting of exfoliated platelets from edge-selectively functionalized graphite (EFG). The newly developed edge-selective functionalization method enabled us to achieve exfoliation of “pristine” graphite into individual graphene and few-layer graphene platelets in a scalable and simple manner. For example, most of the 4-substituted benzoic acids can be used as “molecular wedges” and covalently attached to the edge of graphite. The edge groups on graphite are mainly sp²-bonded C—H groups, a reactive site for “direct” Friedel—Crafts acylation in polyphosphoric acid (PPA)/phosphorus pentoxide (P₂O₅).

RESULTS AND DISCUSSION

From the elemental analysis (Supporting Information, Table S1), the possible available sp²—bonded C—H sites at the edges could be calculated to be about one per every 45 carbons. Hence, a sufficient number

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Received for review March 22, 2011 and accepted May 17, 2011.

Published online May 17, 2011 10.1021/nn201072m

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of “wedge” molecules necessary for exfoliation can be attached around graphite by treating it with 4-ethylbenzoic acid (EBA) in the reaction medium PPA/P₂O₅ at 130 °C (Figure 1a–c). The EFG was then completely worked-up and analyzed (Figure 1d, see Methods, Supporting Information, Figures S1 and S2a) prior to dispersing in organic solvents.

Thermogravimetric analysis (TGA) was used to determine the amount of EBA grafted onto the graphite. As can be seen in Figure 1e, graphite is thermooxidatively stable up to 650 °C in air, while the EFG shows stepwise weight losses. We assign the first weight loss at about 400 °C, which continues until about 720 °C, to the EBA moiety being thermally “stripped off”. Hence, the degree of functionalization can be quantified by the weight loss of EFG up to about 720 °C and it is approximately 15 wt %. The degree of functionalization was 76.5% of the available sp² bonded C–H sites. The temperature where the second weight loss starts is approximately 850 °C, or 200 °C higher than that of pristine graphite (650 °C), suggesting that carbon cations, anions, and/or radicals generated during heating (charring) of the 4-ethylbenzoyl (EB) moiety at the edges of EFG provide an in situ carbon feedstock for “healing” inherent defects on graphite and/or “edge-welding” (cross-linking) graphene platelets. The high char yield (91 wt %) of EFG at 720 °C in a nitrogen atmosphere (Figure 1d) was much higher than that in dry air (85 wt %), suggesting that the formation of large-area graphene films could be possible via solution casting and subsequent “edge-welding” by thermal annealing at elevated temperature. To evaluate whether the “char” at the edges blocks oxidants from accessing the interior of the EFG, and hence keeps the graphitic plane structure intact, TGA thermograms were obtained with a ramp rate of 1 °C/min (Supporting Information, Figure S3).

The thermograms for both graphite and EFG were very similar to those obtained with a ramp rate of 10 °C/min (see Figure 1e). Both samples had degradation temperatures down-shifted approximately 180 °C and the stepwise weight loss of mass in the EFG was evident leaving almost the same amount of char yield (15 wt %) at the plateau region at 540 °C. Hence, the higher thermo-oxidative stability of EFG after charring of the EBA moiety is not due to blocking of the oxidant (kinetic effect). More importantly, the fact that there is no weight loss around 100–200 °C, characteristic of interlamellar water in graphite oxide due to its hygroscopic nature, strongly suggests that the graphite is not oxidized during the edge-functionalization in a mild PPA/P₂O₅ medium.

The covalently linked edge groups provide steric repulsion to open pristine graphite edges (Figure 2a) and thus, the EFG in solid state should consist of many stacks of graphene layers as schematically shown in Figure 2b. The partially edge-delaminated EFG allows efficient solvent penetration for the further exfoliation of EFG into few-layer graphene platelets and graphene in solution (Figure 2c). Wide-angle X-ray diffraction (XRD) patterns (Supporting Information, Figure S4) were applied to the same amount of pristine graphite, PPA/P₂O₅-treated graphite, and EFG to estimate the degree of exfoliation and functionalization of the basal plane; the peak intensity of EFG in the solid state is 10% of that of the precursor graphite with almost the same peak location at 26.58° (0.333 Å), indicating the efficient dispersion in solvents through a kind of “unzipping” mechanism schematically shown in Figure 2a–c and without basal plane functionalization. In a control
experiment, graphite was exposed to the same reaction conditions used for the EFG synthesis in the absence of EBA to test if the reaction medium PPA/P2O5 could damage (oxidize) graphite. XRD patterns show that the peak intensity of PPA-treated graphite is stronger than that of graphite with the same peak location at 26.50° (0.334 Å) (Supporting Information, Figure S4). This implies that the PPA-treated graphite is not chemically altered but has increased crystallinity by selectively etching off of amorphous carbon attached to the surface of graphite. Further evidence for the structural integrity of the EFG basal plane region comes from the elemental (EA, Supporting Information, Table S1) and XPS (Figure S5) analyses, which revealed that there was no additional oxygen content beyond the value calculated on the basis of yield from the edge-selective functionalization. Scanning electron microscopy (SEM) of the precursor graphite powder shows that its surfaces are “clean and smooth” (Figure 2d), while the SEM image of bulk EFG in solid state shows fuzzier surfaces (Figure 2e), indicating that the attachment of EBA expanded the edge. Indeed, EFG is an ideal precursor (Figure 2b) for further exfoliation into scalable and high-quality individual graphene and few layer graphene platelets (Figure 2c).

The exfoliation/dispersibility of EFG was tested at room temperature in various solvents, namely water, methanol, ethanol, acetic acid, toluene, N,N-dimethylacetamide (DMAC), heptane, acetone, benzene, dichloromethane, tetrahydrofuran (THF), pyridine and N-methyl-2-pyrrolidone (NMP) (Supporting Information, Figure S2c). It was found that the EFG could be dispersed in all of these solvents, and among them the dispersions in dichloromethane yielded the highest concentration at room temperature. Without physical agitation of any kind (e.g., it is known that sonication of certain types can damage the carbon framework20), the EFG was readily exfoliated to achieve stable dispersions with concentrations as high as 0.8 mg/mL in dichloromethane (Supporting Information, Figure S2c and Movie). Mechanical (magnetic) stirring favored more rapid dispersion. The amount of EFG dispersed in each solvent was calculated as described in the experimental section (see Methods). The maximum concentration for a long-term (months) stable dispersion achieved is 8× higher than that reported for dispersions of graphite (0.1 mg/mL) in NMP,21 which was found to be stable only for days after sonication (see below). Unlike aqueous dispersions of graphite oxide (light yellow),22 the color of the EFG dispersion in dichloromethane is dark brown due to high conjugation in the undamaged basal planes (Figure 2f and Supporting Information, Figure S2d). Tyndall scattering (red circle, Figure 2f) was observed with a hand-held laser pointer, demonstrating formation of colloid. After several months of standing in ambient condition, the supernatant from the dichloromethane dispersion at 0.8 mg/mL concentration remained visually unchanged with no sedimentation evident to the eye. On the other hand, the dispersion of graphite in NMP was readily separated,21 indicating that the dispersion was not thermodynamically stable leaving an essentially clear liquid after 1 day (Supporting Information, Figure S2e). Molecular dynamics simulations suggest that the good dispersibility is because the phenyl rings on the EB groups are vertically aligned to the graphene
plane, leading to a strong steric hindrance for reaggregation of the dispersed graphene platelets (Supporting Information, Figure S6 and detailed computation conditions are described in the Methods section). Transmission electron microscope (TEM) images show the presence of graphene platelets (Figure 2g) with functionalized edges (Figure 2h).

The SEM image of a solution-cast EFG film deposited on silicon oxide (SiO2, 300 nm) surface from a diluted solution in dichloromethane shows individualized EFG platelets, as schematically illustrated in Supporting Information, Figure S7a. The lateral width distribution of EFG platelets (Figure S7b) was determined by an image analyzer to be in the range of 0.2–2.2 μm (Figure S7c); the average lateral size was approximately 1.0 μm. AFM images of platelets, deposited from EFG further diluted in dichloromethane on an SiO2 surface, were used to estimate thicknesses of the platelets (Supporting Information, Figure S8). The maximum heights of few-layer graphene platelets are in the range of 1.2–5.0 nm. As a result of the edge functionalization with EBA, the edge of EFG is not as sharp as graphene obtained from micromechanical exfoliation or of reduced graphene oxide (rGO) platelets. Together with Raman spectroscopy (Supporting Information, Figure S9a) and confocal optical microscopy image (Figure S9b), the object shown at the edge of the EFG platelet in Figure S8a is interpreted to be a few-layer EFG with its edge higher than its interior due to the presence of EB moieties at the edge. The height profiles of platelets indicate that the maximum thickness of such platelets is less than 5 nm (Figure S8). The average lateral width from many other AFM images is ∼1.0 μm. The width distribution from the AFM imaging agrees with the SEM result (see Figure S7).

The concentrated EFG dispersion in dichloromethane was drop-coated on an SiO2 surface. The SEM image shows platelets on the surface over a large area with many wrinkles (Figure 3c), as depicted in a schematic illustration shown in Figure 3a. The as-cast film shows high sheet resistance (∼108 kΩ/sq) likely due to the edge groups at the boundaries between platelets contributing high contact resistance. Upon heat treatment at 600 °C in argon for 3 h, the edge EB moieties could act as an in situ feedstock for carbon to “weld” the graphene platelets thus forming a large-area graphene film (Figure 3b, see also Supporting Information, Figure S10). Indeed, the SEM image of the heat-treated EFG (HEFG) film shows a uniform surface (Figure 3d). This film was scratched with a sharp metal (Figure 3d, see also Supporting Information, Figure S10a). The thickness of the HEFG film was determined to be 16 nm by SEM images (Figure 3e,f) and AFM images using tapping mode (Supporting Information,
Figure S10c). Welded boundaries of HEFG could be distinguished from the surface of HEFG thin film (red circle, Figure 3g). The boundary of platelets, where thermal welding was assumed to occur during heat-treatment, can be clearly seen at high magnification (Figure 3h). Selected area electron diffraction (SAED) indicates that the HEFG film has high crystallinity (Figure 3i). These results demonstrate that large-area graphene films can be achieved by solution casting. The 16 nm thick film had an average sheet resistance of 1.3 kΩ/sq. This value is a few orders of magnitude lower than that of the chemically rGO.25,26 The observed reduction in sheet resistance through heating the EFG film supports the concept of edge chemistry “boundary welding” that is in a good agreement with the TGA results (see Figure 1b,c).18

Large-area graphene films were made with different EFG concentrations in dichloromethane. These solutions were drop-coated onto an SiO2 surface and subsequently heat-treated at 600 °C in argon for 3 h (Figure 4a). To transfer the solution-cast HEFG films onto other substrates, poly(methylmethaacylate) (PMMA) solution in tetrahydrofuran (THF) was spin coated on such a HEFG film (Figure 4b). The SiO2 layer was etched off by floating on aqueous 1 M hydrofluoric acid (HF) solution (Figure 4c). Then, the HEFG on PMMA (HEFG/PMMA) film was transferred to poly(ethylene terephthalate) (PET) film and the PMMA was washed off by immersing in acetone to produce HEFG on PET (HEFG/PET) (Figure 4d). Four different HEFG/PET films with approximately 2 × 2 in. dimensions were tested, though many larger HEFG films on various substrates can be readily prepared through the similar procedure.

The optical transmittances and sheet resistances of the films were measured by using UV−vis spectroscopy and four-point probe, respectively. The optical transmittances are in the range of 63−90% (Figure 4e) with sheet resistances of 0.52−3.11 kΩ/sq (Figure 4f). Although these optoelectronic properties are still not as good as those of the CVD grown graphene film,12 solution casting has many advantages in processing over the CVD process, indicating cost and scalability to very large area. Figure 5 shows I−V characteristics of a field-effect transistor of HEFG film that was transferred onto an SiO2(300 nm)/Si wafer with prepatterned gold electrodes. The film shows ambipolar behavior with a Dirac point around −50 V.27

CONCLUSION

Edge-selectively functionalized graphite (EFG) without damage to the basal plane can be produced with a...
simple one-pot reaction between graphite and 4-ethylbenzoic acid (EBA) in viscous PPA/P2O5 under mild conditions. The EFG could be dispersed in various organic solvents. Among 13 different organic solvents tested in this study, dichloromethane, THF, DMAc, and NMP were good solvents for dispersing EFG and tested in this study, dichloromethane, THF, DMAc, and NMP were good solvents for dispersing EFG and this is attributed to high chemical affinity with these solvents. The EFG was dispersed in dichloromethane and exfoliated into EFG thin platelets. Edge-selective functionalization provides a pathway to exfoliation of graphite into few-layer graphene platelets without damages on their basal plane for better performance. Dispersed platelets were drop-coated to produce a thin film that was then heat-treated at 600 °C under argon atmosphere, yielding a uniformly welded large-area graphene films with low sheet resistance (0.52 – 3.11 kΩ/sq) and comparable optical transmittance (63–90%). The EFG approach is a scalable method for the cost-effective production of high-quality graphene films.

METHODS

Materials. All reagents and solvents were purchased from Aldrich Chemical Inc., and used as received, unless otherwise mentioned. 4-Ethylbenzoic acid as a reactive molecular wedge was purified by recrystallization from heptanes (mp = 112 – 113 °C). Graphite (99.9% purity, stock no. 1250HT, CAS no. 7782-42-5, Lot no. 1250-100907) was obtained from Nanostructured & Amorphous Materials Inc., USA, and used as received.

Instrumentations. Raman spectra were taken with He–Ne laser (532 nm) as the excitation source by using combined AFM and confocal Raman microscope (Alpha 3005, WITec, Germany). Thermogravimetric analysis (TGA) was conducted in air and nitrogen atmospheres at a heating rate of 10 and 1 °C/min using a TA Hi-Res TGA 2950 thermogravimetric analyzer. The field emission scanning electron microscopy (FE-SEM) used in this work was LEO 1530FE and FEI NanoSEM 200. The field emission transmission electron microscope (FE-TEM) employed in this work was a FEI Tecnai G2 F30 S-Twin. The operating voltage was 200 kV. The samples were prepared by dispersion in N-methyl-2-pyrrolidone (NMP). X-ray photoelectron spectroscopy (XPS) was performed with Thermo Fisher K-alpha. Elemental analysis (EA) was conducted with Thermo Scientific Flash 2000. Atomic force microscopy (AFM) analysis was conducted with Veeco Multimode V. X-ray diffraction (XRD) patterns were recorded with Rigaku D/Max2500/PC. The field-effect transistor was fabricated by transferring the HEFG film on prepatterned electrodes, which were defined on SiO2/Si (0.52 sq) and comparable optical transmittance

REFERENCES AND NOTES


