Single Source Precursor-based Solvothermal Synthesis of Heteroatom-doped Graphene and Its Energy Storage and Conversion Applications

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Solvothermal processes are considered efficient approaches for the gram-scale production of graphene. Further modification of graphene by chemical doping is an important approach to tailor its properties. In this work, we successfully synthesized sulfur-doped graphene by using a solvothermal method with dimethyl sulfoxide as a precursor, which is a common laboratory reagent. Nitrogen-doped graphene was produced to demonstrate the generality of this process. These heteroatom-doped graphene materials exhibited high surface areas and high contents of heteroatoms. Furthermore, the lithium-ion storage properties and oxygen reduction reaction catalytic activity of these materials were also investigated. The success of this approach might facilitate the development of other advanced graphene-based materials with relative simplicity, scalability, and cost effectiveness for use in various potential applications.
In this research, we describe a novel approach to fabricate S-doped graphene via a solvothermal method using S-containing organic molecules as a precursor. This method directly converts dimethyl sulfoxide (CH₃CH₂SO, DMSO) into S-doped graphene. Furthermore, N-doped graphene was also synthesized using dimethylformamide (CH₃CH₂NCOH, DMF), which is a N-containing organic molecule. These precursors are commonly used as low-cost solvents. It provides the benefits of a single step reaction with relatively mild synthesis conditions, gram-scale products, and higher contents of the heteroatom. The lithium-ion storage properties and electrocatalytic behaviors of heteroatomic-doped graphene were investigated.

Results
Synthesis and characterization of heteroatom-doped graphene. DMSO, the S-containing organic molecule, was heated with NaOH under N₂ gas flow. The mixture was brought to a boil, and maintained the boil with reflux. Under the condition, the colorless liquid became dark brown (see Supplementary Fig. S1). Finally, black cake-like materials were obtained and washed using deionized water then dried in an oven. Two-dimensional sheet-like structures were obtained, which were formed with carbon and sulfur atoms (Fig. 1). The surface morphology of the S-doped graphene was analyzed by using atomic force microscopy (AFM). The AFM images showed crumpled veil-like structures with thickness of around 1 nm (see Supplementary Fig. S2). With 50 ml of DMSO, more than 1.0 g of product per batch was obtained. We believe that this process could be scaled up for larger synthetic yields. To demonstrate that the heteroatom-containing organic molecules could be converted into heteroatom-doped graphene, DMF was chosen to produce N-doped graphene, to represent N-containing molecules. Roughly 2.6 g of product was obtained from 50 mL of DMF. Pristine solvothermal graphene was also prepared using methanol as a precursor (see Supplementary Fig. S3). In a previous study, hexagonal carbon clusters were synthesized by using alkali metals as reducing agents at a low temperature²⁶. In that study, free C=C was stated to be the intermediate during the formation of sp² hybridized carbon. N-doped graphene was also synthesized through the reaction of alkali metal salts (Li₃N) and carbon source (CCl₄)²⁹. Therefore, we propose that C=C, C-S-C, and C-S groups are probably the reaction intermediates, which subsequently assemble into a S-contained sp² hybridized carbon (Fig. 1a).

To investigate the elemental dopant distribution, TEM energy-dispersive X-ray spectroscopy (EDS) mapping was carried out. Fig. 1d shows a homogeneous distribution of elemental S in the graphene sheets. Raman spectroscopy of S-doped graphene, N-doped graphene, and solvothermal graphene was investigated. Broad D and G bands were determined at around 1350 and 1600 cm⁻¹, respectively (Fig. 2). The spectra of the S-doped graphene agreed well with the literature²⁶. X-ray diffraction (XRD) patterns of the solvothermal graphene and graphite were analyzed (Fig. S4). Graphite shows a sharp peak at 26.3° (see Supplementary Fig. S4) corresponding to the (002) diffraction peak (d-spacing = 0.34 nm). The (002) peaks of the S-doped graphene, N-doped graphene and solvothermal graphene are shown at 22.3°, 25.5°, and 23.4°, respectively. The interlayer distance of the S-doped graphene is 0.40 nm, which is larger than that of the N-doped graphene (0.35 nm) and solvothermal graphene (0.38 nm). The conductivity of the S-doped graphene and N-doped graphene is 0.607 × 10⁻² S m⁻¹ and 0.416 × 10⁻² S m⁻¹, which is lower than that of the solvothermal graphene (0.121 S m⁻¹; see Supplementary information). The results show that S or N doping increases the resistance of graphene.

The N₂ adsorption and desorption isotherms and pore size distributions of the heteroatom-doped graphene were also analyzed (Fig. 2b and Supplementary Fig. S5). The Brunauer–Emmett–Teller (BET) surface area of the S-doped graphene and N-doped graphene were 754 m² g⁻¹ and 317 m² g⁻¹ with the pore volumes of 0.82 cm³ g⁻¹ and 0.91 cm³ g⁻¹, respectively. After annealing at high temperature (900 °C) under an inert atmosphere, the respective BET surface areas of the S-doped graphene and N-doped graphene were increased to 1564 m² g⁻¹ (PV = 1.53 cm³ g⁻¹) and 2255 m² g⁻¹ (PV = 1.73 cm³ g⁻¹).

For elucidating the chemical structure of the S-doped graphene, X-ray photoelectron spectroscopy (XPS) was employed. The XPS spectra of the solvothermal graphene derived from methanol confirmed that carbon and oxygen were the only atoms present. However, sulfur and nitrogen signals could be clearly observed in the XPS spectra of the S-doped graphene and N-doped graphene, respectively (Fig. 3a). The high-resolution Cls peak corresponding to sp²-hybridization (C=C bond, 284.5 eV) was observed, and the samples produced a small peak corresponding to C-O bonds, carbonyls (C=O), and carboxylates (O-C=O) (Fig. 3b)²⁰. The high-resolution S2p spectra of the S-doped graphene can be deconvoluted into three different peaks (Fig. 3c). Lines at 163.9, 165.1 and 168.5 eV correspond to 2p½, 2p¾, and oxidized sulfur groups, respectively. Additionally, there existed an invisible peak corresponding to the thiol group (162.0 eV)²⁰. The high-resolution N1s spectra of the N-doped graphene can also be decomposed into three different peaks that can be
assigned to pyridinic N (398.0 eV), pyrrolic N (400.1 eV), and graphitic N (401.3 eV) (Fig. 3d). The results suggest that the S or N atoms are covalently bonded network of graphene. According to the elemental analysis, the sulfur content of the S-doped graphene was 22.83 wt%, and the nitrogen content of the N-doped graphene was 12.25 wt% (see Supplementary Table S1). The sulfur content of the S-doped graphene is much higher than those previously reported literature (<5%)\(^2\(^7\).\(^2\(^4\).

**Lithium storage properties.** To the best of our knowledge, the battery properties of the graphene family prepared by solvothermal methods have yet to be reported. To evaluate the potential applicability of the S-doped graphene, N-doped graphene and solvothermal graphene as anode materials, electrochemical cells with graphene electrodes were galvanostatically charged and discharged at a current density of 200 mA g\(^{-1}\) at room temperature (Fig. 4). The reversible capacities of the S-doped graphene, N-doped graphene and solvothermal graphene electrodes are reasonably high (500–800 mAh g\(^{-1}\)). Interestingly, the specific capacities decrease during initial 30 cycles and gradually increase as the number of cycles increases in the S-doped graphene and solvothermal graphene. These activation processes can also be observed in other graphene-based electrodes\(^3\(^1\).\(^3\(^2\).\(^ Notably, the S-doped graphene, N-doped graphene and solvothermal graphene electrodes exhibited high
from the 30th to 150th cycle was 97.2%, which is a relatively high value, but those of the S-doped and N-doped graphene were much higher (99.8% and 98.5%, respectively). It was demonstrated that half-cell Coulombic efficiency could affect cycle retention in full lithium ion cells. Additionally, the plateau in the first discharge process that occurs around 1.0–0.7 V in voltage profiles (more clearly seen in the CV curves) is related to the decomposition of electrolyte and the formation of a solid electrolyte interphase (SEI) layer (see Supplementary Fig. S6), this plateau is remarkable in the solvothermal graphene electrode. These findings indicate that the doping of nitrogen and sulfur in graphene can suppress side reactions, which include electrolyte decomposition. The shape of charge and discharge profiles in the N-doped graphene and solvothermal graphene is similar to those in other graphene; however, the S-doped graphene profile is slightly different. Additional cathodic (−1.4 V) and anodic (−2.4 V) peaks were observed in the CV curves of the S-doped graphene (see Supplementary Fig. S6e), which are attributed to the conversion of polysulfides to lithium sulfides and lithium sulfides to polysulfides, respectively. The N-doped graphene electrode shows the highest rate capability. The S-doped graphene, N-doped graphene, and solvothermal graphene electrodes delivered specific capacities of 270.5, 278.9, and 267.5 mAh g⁻¹, respectively, at current density of 2500 mA g⁻¹, which correspond to the respective delivered capacities of 44.9%, 40.7%, and 53.1% at a current density of 200 mA g⁻¹. The theoretical capacity of graphene is 744 mAh g⁻¹ through the formation of LiC₃ in a condition where all the graphene layers present as monolayers. It is reported that the lithium storage properties of graphene have a sensitive dependence on many factors such as synthetic methods and condition, interlayer spacing, disorder degree, and surface functional groups. The reversible capacities of graphene can vary from around 300 mAh g⁻¹ to more than 1500 mAh g⁻¹ (see Supplementary Table S2). The S-doped graphene and N-doped graphene by solvothermal method display high cycle stability with excellent Coulombic efficiency compared to other previous reported graphene electrodes.

Electrocatalytic performance. To investigate the electrocatalytic activity of the S-doped graphene and N-doped graphene for the ORR and to confirm the doping effect, we performed rotating-disk electrode (RDE) voltammetry in 0.1 M KOH electrolyte (20 wt% Pt/C commercial E-TEK catalyst). The N-doped graphene shows higher ORR activity than pristine solvothermal graphene. Further, N-doped graphene has a similar onset potential (0.9 V_RHE) as platinum electrode. The activity of doped carbon for the ORR has been improved by a combination of many factors, which have not yet been clarified. However, it is important to note that the enhancement factor can be explained by doping, for example, the presence of N, B, and S in carbon. These results indicate that the electrocatalytic activities of the catalyst for the ORR increase with the introduction of a dopant whose electronegativity is different from that of carbon. Furthermore, Koutechy-Levich analysis by changing the rotating speed reveal that the electron transfer number changes from 2.8 in the pristine solvothermal graphene to 3.9 in the N-doped graphene at 0.2 V_RHE (see Supplementary Fig. S7). For clear emphasis on the importance of proper doping effects on ORR activity in the region of real application, we chose the potential (0.6 V_RHE) to compare the electron transfer number in Fig. Sb, which is very high potential compared to the general viewpoint. In the case of solvothermal graphene, the electron transfer number indicates dominant two-electron pathway that is ineffective for the ORR. However, when doping with elements such as N and S, the ORR pathway would involve both 2-electron and 4-electron transfer which are more favorable for the ORR. Solvothermal heteroatom (N, S)-doped graphene was synthesized by a bottom-up method. It is different from top-down method and contains a significant amount of metallic impurities which may influence the activity. In this work, our solvothermal process uses NaOH or Na metals which do not affect the activity. XPS results also confirmed that there is no other impurity in the as-synthesized graphene. Considering that the doping content in the present metal-free graphene is rather small, the ORR activity of doped graphene is rather high, and the ORR activity of doped graphene is rather high.
optimized, further improvement of ORR activity can be achieved by optimizing the contents and configuration.

**Discussion**

In summary, heteroatom (S or N)-doped graphenes were successfully synthesized by using a simple, single-step solvothermal method with heteroatom-containing organic molecules as precursors. This process provides for a gram-scale production using common laboratory reagents. The solvothermal (S or N)-doped graphenes exhibited high heteroatom content and surface area, and their characterization data indicate that they could be utilized in various energy storage and conversion applications. We believe that heteroatom-doped graphenes can be further extended to applications in sensors, electronics and biological research. We expect that our synthetic approach will be developed to produce doped carbon materials based on other elements (e.g., B, P, and F) which can then increase the method’s potential applications.

**Methods**

**Synthesis of S-doped graphene.** A mixture of 2 g of NaOH and 10 mL of DMSO solution was added into a three-neck flask with a reflux condenser and heated under a N₂ gas flow. The colorless mixture was boiled at around 250°C and was slowly thickened. After boiling for ~24 h, the liquid was converted into a black cake-like substance. Finally, the samples were washed with hydrochloric acid (10 wt%) and deionized water several times, and then dried in an oven at 90°C. The final yield of S-doped graphene is ~35 mg per 1 mL of DMSO (see Supplementary Fig. S1).

**Synthesis of N-doped graphene.** N-doped graphene was synthesized by using a solvothermal method. In a typical synthesis, 30 mL of DMF and 25 g of sodium metal were placed in a 100 mL Teflon-lined autoclave that was sealed and heated at 190°C for 72 h. After cooling to room temperature, the autoclave was opened. The product was removed immediately to a beaker containing 10 wt% of hydrochloric acid. The sample was further washed with deionized water several times, and then dried in an oven at 90°C. (Caution: before washing, the sodium-containing solid state sample can catch fire easily when exposed to air.)

**Synthesis of solvothermal graphene.** Solvothermal graphene were prepared according to a literature procedure using methanol as the precursor. In a typical synthesis, 30 mL of methanol and 25 g of sodium metal were placed in a 100 mL Teflon-lined autoclave that was sealed and heated at 190°C for 72 h. After the autoclave was cooled to room temperature, the solids were annealed at 500°C under a N₂ gas flow. The resulting samples were washed with hydrochloric acid (10 wt%) and deionized water several times, and then dried in an oven at 90°C.

**Characterization.** TEM images and TEM-EDS images were obtained using a JEOL JEM-1010 and JEM-2100F, respectively. AFM images were obtained using a Veeco atomic force microscope. Raman spectroscopy measurements were carried out on a Renishaw 1000R laser spectrometer using 532 nm excitation. BET surface area and BJH pore size distributions were measured by using a Micromeritics TriStar 3000. XPS was carried out using a Kratos AXIS-HS. Elemental analysis was performed using a Thermo Scientific Flash 2000 analyzer at NCIRF (National Center for Inter-university Research Facilities, Seoul National University, Korea).

**Electrochemical characterization (battery application).** Samples were prepared by heat treatment at 600°C in N₂ for battery applications. For the preparation of working electrodes, a well-mixed slurry of synthesized graphene, Super P and poly (vinylidene fluoride) in N-methyl-2-pyrrolidone (at a weight ratio of 70: 10: 20) was coated on a copper current collector. After drying, the electrodes were pressed and dried again at 120°C under vacuum. Two electrode 2016-type coin cells were fabricated with prepared working electrodes and lithium foil (as both counter and reference electrodes) in argon filled glove box. A 1.0 M KOH electrolyte. To calculate the electron transfer number (0.6 V RHE), RDE measurements were conducted using different rotating speeds 400, 900, 1200, and 1600 rpm.

1. Novoselov, K. S. et al. A roadmap for graphene. Nature 490, 192–200 (2012).
2. Wang, H. & Dai, H. Strongly coupled inorganic–nano-carbon hybrid materials for energy storage. Chem. Soc. Rev. 42, 3088–3113 (2013).
3. Mao, H. Y. et al. Graphene: Promises, Facts, Opportunities, and Challenges in Nanomedicine. Chem. Rev. 113, 3407–3424 (2013).
4. Bo, Z. et al. Green preparation of reduced graphene oxide for sensing and energy storage applications. Sci. Rep. 4, 6844 (2014).
5. Dai, L. Functionalization of Graphene for Efficient Energy Conversion and Storage. Acc. Chem. Res. 46, 31–42 (2012).
6. Yoon, J.-C., Lee, J.-S., Kim, S.-I., Kim, K.-H. & Jang, J.-H. Three-Dimensional Graphene Nano-Networks with High Quality and Mass Production Capability via Precursor-Assisted Chemical Vapor Deposition. Sci. Rep. 3, 1788 (2013).
7. Kali, J. & Wang, Y. Large and fast reversible Li-ion storages in FeO3-graphene sheet-on-sheet sandwich-like nanocomposites. Sci. Rep. 3, 3502 (2013).
8. Kim, H. et al. Scalable functionalized graphene nano-platelets as tunable cathodes for high-performance lithium rechargeable batteries. Sci. Rep. 3, 1506 (2013).
9. Loh, K. P., Bao, Q., Eda, G. & Chhowalla, M. Graphene oxide as a chemically tunable platform for optical applications. Nat. Chem. 2, 1015–1024 (2010).
10. Peng, J. et al. Graphene quantum dots derived from carbon fibers. Nano Lett. 12, 844–849 (2012).
11. Ritter, K. A. & Lyding, J. W. The influence of edge structure on the electronic properties of graphene quantum dots and nanoribbons. Nat. Mater. 8, 235–242 (2009).
12. Han, Z. et al. Hexagonal graphene onion rings. J. Am. Chem. Soc. 135, 10753–10762 (2013).
13. Dimitrakakis, G. K., Tylianakis, E. & Froudakis, G. E. Pillared Graphene: A New 3-D Network Nanostructure for Enhanced Hydrogen Storage. Nano Lett. 8, 3166–3170 (2008).
14. Zhang, L. et al. Porous 3D graphene-based bulk materials with exceptional high surface area and excellent conductivity for supercapacitors. Sci. Rep. 3, 1408 (2013).
15. Wang, H., Majiyalagan, T. & Wang, X. Review on Recent Progress in Nitrogen-Doped Graphene: Synthesis, Characterization, and Its Potential Applications. ACS Catal. 2, 781–794 (2012).
16. Li, X. et al. Boron Doping of Graphene–Silicon p–n Junction Solar Cells. Adv. Energy Mater. 2, 425–429 (2012).
17. Yang, Z. et al. Sulfur-Doped Graphene as an Efficient Metal-free Cathode Catalyst for Oxygen Reduction. ACS Nano 6, 205–211 (2012).
18. Choi, C. H., Chung, M. W., Kwon, H. C., Park, S. H. & Woo, S. I. B- and N- and P-doped graphene as highly active catalysts for oxygen reduction reactions in acidic media. J. Mater. Chem. A 1, 3604–3609 (2013).
19. Zhang, C., Mahmood, N., Yin, H., Liu, F. & Hou, Y. Synthesis of Phosphorus-Doped Graphene and its Multifunctional Applications for Oxygen Reduction Reaction and Lithium Ion Batteries. Adv. Mater. 25, 4932–4937 (2013).
20. Wei, D. et al. Synthesis of N-Doped Graphene by Chemical Vapor Deposition and Its Electrical Properties. Nano Lett. 9, 1752–1758 (2009).
21. Zhang, C. et al. Synthesis, Structure, and Properties of Boron- and Nitrogen-Doped Graphene. Adv. Mater. 21, 4726–4730 (2009).
22. Zhang, C. et al. Synthesis of Nitrogen-Doped Graphene Using Embedded Carbon and Nitrogen Sources. Adv. Mater. 23, 1020–1024 (2011).
23. Poh, H. L., Simsek, P., Sofer, Z. & Pumera, M. Sulfur-Doped Graphene via Thermal Exfoliation of Graphite Oxide in H2SO2, SO2, or CS2 Gas. ACS Nano 7, 5262–5272 (2013).
24. Yang, S. et al. Efficient Synthesis of Heterorotom (N or S) Doped Graphene Based on Ultrathin Graphene Oxide–Porous Silica Sheets for Oxygen Reduction Reactions. Adv. Funct. Mater. 22, 3634–3640 (2012).
25. Wang, R. et al. Controlled Growth of Platinum Nanowire Arrays on Sulfur Doped Graphene as High Performance Electrocatalyst. Sci. Rep. 3, 2431 (2013).
26. Choucivar, M., Thorodson, P. & Stride, J. A. Gram-scale production of graphene based on solvothermal synthesis and sonication. Nat. Nanotechnol. 4, 30–33 (2009).
27. Parasarah, V., Kumar, K., Prakash, R., Pandey, S. K. & Pandey, A. C. Methanol derived large scale chemical synthesis of brightly fluorescent graphene. J. Mater. Chem. A 1, 6506–6509 (2013).
28. Wang, X. et al. A Novel Route to Multilayered Carbon Nanotubes and Carbon Nanorods at Low Temperature. J. Phys. Chem. B 106, 933–937 (2002).
29. Deng, D. et al. Toward N-Doped Graphene via Solvothermal Synthesis. Chem. Mater. 23, 1188–1193 (2011).
30. Lu, J. et al. A nanostructured cathode architecture for low charge overpotential in lithium–oxygen batteries. Nat. Commun. 4 (2013).
31. Al. W. et al. A novel graphene-polylysulphide anode material for high-performance lithium-ion batteries. Sci. Rep. 3, 2341 (2013).
32. Li, X. et al. Superior cycle stability of nitrogen-doped graphene nanosheets as anodes for lithium ion batteries. Electrochem. Commun. 13, 822–825 (2011).
33. Yang, J., Liao, Q., Zhou, X., Liu, X. & Tang, J. Efficient synthesis of graphene-based powder via in situ spray pyrolysis and its application in lithium ion batteries. RSC Adv. 3, 16449 (2013).
34. Ohzuku, T., Ueda, A., Yamamoto, N. & Iwakoshi, Y. Factor affecting the capacity retention of lithium-ion cells. J. Power Sources 54, 99–102 (1995).
35. Wang, G. et al. Sn/graphene nanocomposite with 3D architecture for enhanced reversible lithium storage in lithium ion batteries. J. Mater. Chem. 19, 8378–8384 (2009).
36. Yu, S.-H. et al. Structure-Properties Relationship in Iron Oxide-Reduced Graphene Oxide Nanostructures for Li-Ion Batteries. Adv. Funct. Mater. 23, 4293–4305 (2013).
37. Pan, D. et al. Li Storage Properties of Disordered Graphene Nanosheets. Chem. Mater. 21, 3136–3142 (2009).
38. Yoo, E. et al. Large Reversible Li Storage of Graphene Nanosheet Families for Use in Rechargeable Lithium Ion Batteries. Nano Lett. 8, 2277–2282 (2008).
39. Wu, Z.-S., Ren, W., Xu, L., Li, F. & Cheng, H.-M. Doped Graphene Sheets As Anode Materials with Superhigh Rate and Large Capacity for Lithium Ion Batteries. ACS Nano 5, 5463–5471 (2011).
40. Yu, D., Nagelli, E., Du, F. & Dai, L. Metal-Free Carbon Nanomaterials Become More Active than Metal Catalysts and Last Longer. J. Phys. Chem. Lett. 1, 2165–2173 (2010).
41. Zheng, Y., Jiao, Y., Jaroniec, M., Jin, Y. & Qiao, S. Z. Nanostructured Metal-Free Electrochemical Catalysts for Highly Efficient Oxygen Reduction. Small 8, 3550–3566 (2012).
42. Sharifi, T., Hu, G., Jia, X. & Wågberg, T. Formation of Active Sites for Oxygen Reduction Reactions by Transformation of Nitrogen Functionalities in Nitrogen-Doped Carbon Nanotubes. ACS Nano 6, 8904–8912 (2012).
43. Li, Y. et al. An oxygen reduction electrocatalyst based on carbon nanotube-graphene complexes. Nat. Nanotechnol. 7, 394–400 (2012).
44. Zheng, Y., Jiao, Y., Ge, L., Jaroniec, M. & Qiao, S. Z. Two-Step Boron and Nitrogen Doping in Graphene for Enhanced Synergistic Catalysis. Angew. Chem. Int. Edit. 52, 3110–3116 (2013).
45. Zhang, Y. et al. Manageable N-doped Graphene for High Performance Oxygen Reduction Reaction. Sci. Rep. 3, 2771 (2013).
46. Ambrosi, A. et al. Metallic Impurities in Graphenes Prepared from Graphite Can Dramatically Influence Their Properties. Angew. Chem. Int. Edit. 51, 500–503 (2012).
47. Wang, L., Ambrosi, A. & Pumera, M. “Metal-Free” Catalytic Oxygen Reduction Reaction on Heteroatom-Doped Graphene is Caused by Trace Metal Impurities. Angew. Chem. Int. Edit. 52, 13818–13821 (2013).

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Author contributions
B.Q. proposed the concept and performed the experiment. S.-H.Y. and A.J. carried out lithium storage properties. D.Y.C. carried out electrocatalytic performances. J.H.P. carried out the AFM data. Y.P., B.Q. and S.-H.Y. co-wrote the manuscript. Y.P. and Y.-E.S. coordinated and supervised the work.

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