Communication

Synergistic Effect of a Defect-Free Graphene Nanostructure as an Anode Material for Lithium Ion Batteries

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Abstract: Graphene nanosheets have been among the most promising candidates for a high-performance anode material to replace graphite in lithium ion batteries (LIBs). Studies in this area have mainly focused on nanostructured electrodes synthesized by graphene oxide (GO) or reduced graphene oxide (rGO) and surface modifications by a chemical treatment. Herein, we propose a cost-effective and reliable route for generating a defect-free, nanoporous graphene nanostructure (df-GNS) through the sequential insertion of pyridine into a potassium graphite intercalation compound (K-GIC). The as-prepared df-GNS preserves the intrinsic property of graphene without any crystal damage, leading to micro-/nano-porosity (microporosity: ~10–50 µm, nanoporosity: ~2–20 nm) with a significantly large specific surface area. The electrochemical performance of the df-GNS as an anode electrode was assessed and showed a notably enhanced capacity, rate capability, and cycle stability, without fading in capacity or decaying. This is because of the optimal porosity, with perfect preservation of the graphene crystal, allowing faster ion access and a high amount of electron pathways onto the electrode. Therefore, our work will be very helpful for the development of anode and cathode electrodes with higher energy and power performance requirements.

Keywords: defect free graphene; graphite intercalation compound; lithium ion battery; anode material; nanostructure

1. Introduction

Since graphite, with relatively slow solid-state lithium diffusion, was initially utilized as an anode material in lithium ion batteries (LIBs), active materials with high power and energy density levels have attracted growing interest due to their potential applications in mobile electronics and the electric vehicle market. However, transition metal oxides [1–5], one of the most promising candidates, show relatively high redox potential values, an irreversible phase transition, and rapid capacity decay, in spite of the high theoretical capacity and low cost of these materials.

Graphene, a two-dimensional (2D) honeycomb lattice made of carbon atoms, has been enthusiastically studied, owing to its extraordinary electrical [6] and mechanical [7] properties. Furthermore, graphene has been widely used as a textual additive for the fabrication of nanostructure electrodes and as conducting agents for the improvement of the electrical conductivity between active materials. Graphene oxide (GO) and reduced graphene oxide (rGO), as well as rGO/metal oxide...
composites, have been widely recognized, and their electrochemical performance capabilities have been assessed as anode and cathode electrodes in lithium ion batteries [8-12]. For example, Jiang et al. reported a three-dimensional (3D) porous MoS\textsubscript{2}/GO nanostructure synthesized by a hydrothermal method, showing a significant enhancement of the rate performance [13]. Yu et al. demonstrated that a mesoporous rGO structure with a well-defined porosity, fabricated from colloidal nanocrystals (NCs), provides a considerable performance improvement, with good reproducibility in LIBs [14]. Furthermore, Lee et al. suggested a nanohole-structured architecture synthesized from gelatin-functionalized GO via microwave irradiation, resulting in a high rate performance with stable capacity retention [15]. Furthermore, the Dareeke, Kalantar-zadeh, and Yeo groups have suggested new methods for fabricating layered solid carbonaceous materials collected from CO\textsubscript{2} conversion and their application to capacitors [16], as well as peeling off bulk 3D piezoelectric crystals (WS\textsubscript{2} and MoS\textsubscript{2}) to 2D nanosheets by using high frequency acoustic waves [17]. However, oxygen functional groups ascribed to strong oxidation chemicals (H\textsubscript{2}O\textsubscript{2}, HNO\textsubscript{3}, and KMnO\textsubscript{4}) and imperfect crystal recovery rates have mainly led to limited performance improvements in LIBs. Recently, our group has reported the scalable fabrication of a potassium graphite intercalation compound (K-GIC) and its exfoliation to defect-free graphene [18]. However, the formation of a nanostructure with an optimal porosity level from perfect graphene remains to be achieved, required in order to realize higher energy density and power density levels with good electrochemical performance stability.

Here, we propose a simple and cost-effective method for generating a defect-free graphene nanostructure with optimal micro-/nano-porosity through the controllable expansion of K-GIC. From Raman spectroscopy and high resolution-transmission electron microscopy (HR-TEM), X-ray diffraction (XRD), and X-ray photoelectric spectroscopy (XPS) results, the synthetic df-GNS preserves the unique properties of pristine graphene without the introduction of additional functional groups and without structure damages on the basal plane. Our method is also scalable for the fabrication of the df-GNS (>1 g/batch). Regarding the electrical energy storage performance of LIBs, the df-GNS shows stable capacity retention at different current densities (470 mAh g\textsuperscript{-1} at 0.03 A g\textsuperscript{-1} and 350 mAh g\textsuperscript{-1} at 0.1 A g\textsuperscript{-1} after 100 cycles). Moreover, the rate capability of the df-GNS (retention capacity of ~18% at 5 A g\textsuperscript{-1}) is significantly higher than that of rGO (~11% at 5 A g\textsuperscript{-1}) and graphite (~0.1% at 5 A g\textsuperscript{-1}). Overall, the utilization of df-GNS as an anode electrode, as suggested in this work, will provide a practical solution for LIBs, offering high energy density and power density levels.

2. Materials and Methods

2.1. Material and Fabrication of Defect-Free Graphene Nanostructure

Graphite powder and potassium metal were purchased from Bay Carbon, Inc., Bay, MI, USA. (SP-1 graphite powder) and Kojundo Chemical Inc., Tokyo, Japan (KKE02GB, keep in oil), respectively. For fabrication of potassium intercalation compound, a home-made apparatus was used. Graphite powder (1 g) and potassium metal (0.41 g), filled in Pyrex tubes (K metal: 10 mm in diameter, outside; Graphite powder: 6 mm in diameter, inside), were evacuated for 30 min, and the outer Pyrex tube was sealed and kept at 110 °C for 1 h in a glove box under an Ar atmosphere. Then, 200 mg of a potassium graphite intercalation compound (KC\textsubscript{0}) was put into 10 mL of the selected solvents and kept at 5 °C for 24 h for preparation of the df-GNS with optimal micro-/nano-porosity. The synthetic df-GNS was washed with distilled water, heated to a mild temperature (~40 °C), put through a filtration process, because of the necessity of the removal of residue potassium ions, and then dried at 80 °C for 24 h. To find the optimal conditions, experiments using various solvents and operation temperatures were conducted, and the best condition was to be kept at 0 °C when forming the df-GNS with a larger surface area and high pore volume porous architecture for forming porous df-GNS. For the solvents, hexane, toluene, tetrahydrofuran (THF), ethanol (EtOH), pyridine, dimethylformamide (DMF), and dimethyl sulfoxide (DMSO) were chosen, according to different polarities, and were purchased from Sigma–Aldrich Chemicals Co, St. Louis, Mo, USA.
2.2. Electrochemical Measurement

The working electrode was fabricated by mixing the df-GNS, super-P, and polyvinylidifluoride (PVDF) at a weight ratio of 80:10:10 and then casting on copper foil. The electrodes were dried at 60 °C for 12 h in a vacuum atmosphere. The active material weight of the final electrodes was ~1.0 mg cm⁻². Lithium foil (450 μm), separator (polyethylene, 2400), and electrolyte consisting solutions of LiPF₆ in ethylene carbonate (EC)/dimethyl carbonate (DMC) (1/1 in volume) were chosen. Here, a 2032 type coin cell was selected, and the cell assembly was conducted in a glove box under an Ar atmosphere. The cycling and rate performance was tested at various current densities after a 0.03 A g⁻¹ pre-cycle at a voltage range of 0.01–3.0 V.

3. Results and Discussion

Graphite powder was chosen to synthesize a highly ordered potassium graphite intercalation compound (K-GIC), with the overall procedure illustrated in Figure 1a. Graphite powder (1 g, 0.083 M) and potassium metal (0.407 g, 0.0104 M) was added to the Pyrex tubes and evacuated for 30 min, then sealed and kept at 110 °C for 1 h in a glove box. The potassium metals were introduced into the graphite crystals and the interlayer spacing between graphene layers was expanded from ~0.34 nm to ~0.57 nm. The potassium graphite intercalation compounds (KCs, gold color) were then very slowly expanded to the df-GNS with micro-/nano-porosity at 0 °C for 24 h in a pyridine solution. In our work, the selected method is the best preparation condition among various expansion conditions, determined after tests with different types of solvents and reaction times in the solvents. Further details of the experimental method are described in our previous work [16] and in the experimental section. Figure 1b shows digital images of the KC₈ (before) and the df-GNS (after) in the pyridine solution before and after the expansion process. After 24 h at 0 °C, the KC₈ powder in the pyridine solution changed to a slurry-like powder, with the disappearance of the gold-colored K-GIC crystals, where it was fully dispersed in the pyridine solvent. From the SEM images (Figure 1c), the morphology of the synthetic df-GNS showed the formation of a nanostructure with a micro-/nano-size distance towards the out-of-plane direction, without a notable size decrease from the initial graphite powder (lateral average size of ~100 μm), presenting microporosity of ~10–50 μm and nanoporosity of ~2–20 nm. In our work, pyridine, which is geometrically similar to the hexagonal lattice of graphene, was found to be the most effective expansion solvent because of its peculiar feature, where it can lead to the formation of micro-/nano-porosity due to the gradual extinction of the van der Waals (vdW) interaction between the graphene layers, with low mechanical stress via the sequential insertion of pyridine molecules into the K-GIC crystal. This result strongly supports the selection of this solvent for the expansion of the graphene layers.
Figure 1. (a) Schematic drawing of the steps used to the defect-free graphene nanostructure. (b) Digital images of the potassium graphite intercalation compound (KCs) and the defect-free, nanoporous graphene nanostructure (df-GNS) dispersed in a pyridine solution. (c) SEM images of the df-GNS (microporosity shown on the left and nanoporosity shown on the right).

Figure 2a shows typical low- and high-magnification TEM images and the selected area diffraction pattern (SAED), clearly indicating the various types of sheet-like multilayered structures and the crystal quality. The df-GNS sample shows a highly-ordered hexagonal lattice (inset of Figure 2a) [19]. These results prove the perfect preservation of the graphene crystal during the expansion process to the df-GNS without the introduction of used chemicals. From studies of the Raman spectroscopy for the graphite, KCs, and df-GNS, we observed a peak shift and broadening for the potassium graphite intercalation compound (KCs) and recovery of the main peaks without an increase of the D band (df-GNS), as shown in Figure 2b. Interestingly, the G band of the KCs was broadened and exhibits two peaks at ~1520 cm⁻¹ and ~1580 cm⁻¹, which were caused by the coupling and interference between the phonon-electron interfaces [20], while its 2D band, known as the effective method used to determine the thickness of the graphene layer, has obviously disappeared. On the other hand, the G band of the df-GNS was slightly red-shifted to ~1582 cm⁻¹, which could be ascribed to change of resonant vibration frequencies by the formation of the porosity, and also from its D band. Here, we confirm that none of the steps have led to any structural damages of the basal planes. Nitrogen adsorption measurement is very useful in determining the textural properties of solids, for instance, for examining the porosity and surface area. Figure 2c shows the N₂ adsorption/desorption isotherm data measured at 77 K. The specific surface area was calculated by the Brunauer–Emmett–Teller (BET) equation. The isotherm of the df-GNS presents a H3 hysteresis loop according to IUPAC classification, and the specific surface area of the df-GNS is 281.6 m² g⁻¹ while that of the graphite is ~10 m² g⁻¹. This result suggests that porosity of the df-GNS is clearly developed during expansion process.

Figure 2. Characterizations of the graphite, KCs, and df-GNS. (a) Low- (top) and high- (bottom) magnification TEM images of the df-GNS (inset; selected area diffraction pattern (SAED) for 1–210, 0–110, −1010, and −2110). (b) Raman spectroscopy for the graphite, KCs, and df-GNS. (c) N₂ adsorption/desorption isotherm of graphite and df-GNS, determined at 77 K.

To assess df-GNS as an anode material in a lithium ion battery, a working electrode was fabricated by the slurry of a mixture of the active materials (Graphite, rGO, and df-GNS), super-P, and polyvinylidene fluoride (PVDF, Aldrich) at a weight ratio of 80:10:10 on a copper foil. The typical active mass loading was in the range of 1.0–1.5 mg cm⁻². A piece of 450 μm lithium foil (Honjo metal), a polyethylene separator (Asahi Kasei), and an electrolyte containing 1 M LiPF₆ in ethylene carbonate (EC) and dimethyl carbonate (DMC) at a 1:1 volume ratio (Panax ETEC) were chosen. Further details about the electrode preparation and electrochemical measurement methods are described in the experimental section. Figure 3 shows the typical SEM images, Raman spectrum, and survey results from XPS analyses of the graphite, the rGO, and df-GNS samples. For df-GNS, we observed the clear development of in-plane micro-/nano-porosity without a decrease in the particle-
Interestingly, the reversible capacity capacities for the $0.1 \text{ A g}^{-1}$ electrochemical performance, the cycle. The low columbic efficiency in the first cycle may result from the irreversible lithium on plateau during the charge/discharge process. The reversible capacity of the $df$-GNS was stabilized at ~470 mAh g$^{-1}$ during the ten cycles without a noticeable capacity fade. Along with this high capacity, the columbic efficiency was also suitably stabilized from 72.3% at the first cycle to 99.0% in the ten cycles. On the other hand, the rGO sample exhibits an extremely low columbic efficiency of 46.5% in the first cycle. The low columbic efficiency in the first cycle may result from the irreversible lithium loss due to the irreversible reaction and the formation of solid electrolyte interphase (SEI) [21]. To verify the electrochemical performance, the cycling and rate performance outcomes for the graphite, rGO, and $df$-GNS were evaluated. The cycling performance of $df$-GNS at different current densities of 0.03 and 0.1 A g$^{-1}$ (Figure 4c) was assessed, and the results were compared with those of rGO. The reversible capacities for the $df$-GNS were ~470 mAh g$^{-1}$ at 0.03 A g$^{-1}$ and ~350 mAh g$^{-1}$ at 0.1 A g$^{-1}$ after 100 cycles. Interestingly, the reversible capacity at a current density of 0.03 A g$^{-1}$ gradually increased during the initial 40 cycles for the $df$-GNS sample. This peculiar behavior appears to be effective for facilitating lithium ion access, with a significant decrease in the resistance and an increase in the reversible
capacity upon cycling. The df-GNS sample exhibits the best rate capability, in spite of the increasing current density (0.03 to 5 A g$^{-1}$) when compared to that of the graphite and the rGO, as shown in Figure 4d. Specifically, the capacity of the df-GNS at a current density of 5 A g$^{-1}$ is retained, with the capacity retention being ~18% compared to that of the current density at 0.03 A g$^{-1}$ while the rGO and the graphite are 11% and 0.1%, respectively. With a return of the current rate to the initial 0.1 mA g$^{-1}$, after 30 cycles, the df-GNS steadily recovers to a higher capacity rate. Our approach can pave the way for further improving and stabilizing the performance of energy storage devices such as batteries, fuel cells, and capacitors, and the current defect-free materials can be immediately applicable to active materials in other electrochemical energy storage systems suffering from similar problems.

Figure 4. (a) Schematic diagram of the rGO and df-GNS electrodes. (b) Charge-discharge voltage profiles for the graphite, rGO, and df-GNS at a current density of 0.03 mA g$^{-1}$. (c) Comparison of the cycling performance outcomes of the rGO and df-GNS at current densities of 0.03 A g$^{-1}$ and 0.1 mA g$^{-1}$ (d) Rate capability of the graphite, rGO, and df-GNS samples.

4. Conclusions

We have introduced a simple and scalable route for creating micro-/nano-structured df-GNS, without crystal damages and the introduction of oxygen functional groups. Here, df-GNS was assessed as an anode electrode during electrical energy storage. From structural/chemical analyses, we have confirmed that the df-GNS, in an optimization condition, preserves the unique properties of graphene and forms an optimal nanostructure with a significantly larger surface area. The electrical energy storage performance of the df-GNS exhibits an excellent capacity retention of ~470 mAh g$^{-1}$ at 0.03 A g$^{-1}$ and ~350 mAh g$^{-1}$ at 0.1 A g$^{-1}$ after 100 cycles. Additionally, the rate capability of the df-GNS (~18% of initial capacity at 5 A g$^{-1}$) is significantly enhanced compared to that of the graphite (~0.1%) and rGO (~11%). The overall results suggest that our approach will bring about the enhanced performance of an anode electrode with higher energy and power performance capabilities in electrochemical energy storage devices.

Supplementary Materials: The following are available online at www.mdpi.com/xxx/s1, Figure S1: (a) SEM images of the graphite, the rGO, and the df-GNS. (b) Raman spectroscopy of the graphite, the rGO, and the df-GNS. (c) Chemical composition of the graphite, the rGO, and the df-GNS analyzed by XPS.
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Conflicts of Interest: The authors declare no conflict of interest.

References

1. Idota, Y.; Kubota, T.; Matsufuji, A.; Maekawa, Y.; Miyasaka, T. Tin-based amorphous oxide: a high-capacity lithium-ion-storage material. *Science* 1997, 276, 1395–1397.
2. Brousse, T.; Defives, D.; Pasquerneau, L.; Lee, S.; Herterich, U.; Schleich, D. Metal oxide anodes for Li-ion batteries. *Ionics* 1997, 3, 332–337.
3. Poizot, P.; Laruelle, S.; Grugeon, S.; Dupont, L.; Tarascon, J. Nano-sized transition-metal oxides as negative-electrode materials for lithium-ion batteries. *Nature* 2000, 407, 496–499.
4. Grugeon, S.; Laruelle, S.; Dupont, L.; Tarascon, J.M. An update on the reactivity of nanoparticles Co-based compounds towards Li. *Solid State Sci.* 2003, 5, 895–904.
5. Jamnik, J.; Maier, J. Nanocrystallinity effects in lithium battery materials Aspects of nano-ionics. Part IV. *Phys. Chem. Chem. Phys.* 2003, 5, 5215–5220.
6. Geim, A.K.; Novoselov, K.S. The rise of graphene. In *Nanoscience and Technology: A Collection of Reviews from Nature Journals;* World Scientific: Singapore, 2010 11–19.
7. Lee, C.; Wei, X.; Kysar, J.W.; Hone, J. Measurement of the elastic properties and intrinsic strength of monolayer graphene. *Science* 2008, 321, 385–388.
8. Yang, S.; Cui, G.; Pang, S.; Cao, Q.; Kolb, U.; Feng, X.; Maier, J.; Müllen, K. Fabrication of cobalt and cobalt oxide/graphene composites: Towards high-performance anode materials for lithium ion batteries. *ChemSusChem* 2010, 3, 236–239.
9. Yang, S.; Feng, X.; Ivanovici, S.; Müllen, K. Fabrication of graphene-encapsulated oxide nanoparticles: towards high-performance anode materials for lithium storage. *Angew. Chem. Int. Ed.* 2010, 49, 8408–8411.
10. Zhou, G.; Wang, D.W.; Li, F.; Zhang, L.; Li, N.; Wu, Z.S.; Wen, L.; Lu, G.Q.; Cheng, H.M. Graphene-Wrapped FeOx Anode Material with Improved Reversible Capacity and Cyclic Stability for Lithium Ion Batteries. *Chem. Mater.* 2010, 22, 5306–5313.
11. Wei, W.; Yang, S.; Zhou, H.; Lieberwirth, I.; Feng, X.; Müllen, K. 3D Graphene Foams Cross-linked with Pre-encapsulated FeOx Nanospheres for Enhanced Lithium Storage. *Adv. Mater.* 2013, 25, 2909–2914.
12. Wu, Z.S.; Zhou, G.; Yin, L.C.; Ren, W.; Li, F.; Cheng, H.M. Graphene/metal oxide composite electrode materials for energy storage. *Nano Energy* 2012, 1, 107–131.
13. Jiang, L.; Lin, B.; Li, X.; Song, X.; Xia, H.; Li, L.; Zeng, H. Monolayer MoS2-Graphene Hybrid Aerogels with Controllable Porosity for Lithium-Ion Batteries with High Reversible Capacity. *ACS Appl. Mater. Interfaces* 2016, 8, 2680–2687.
14. Yu, H.; Guo, G.; Ji, L.; Li, H.; Yang, D.; Hu, J.; Dong, A. Designed synthesis of ordered mesoporous graphene spheres from colloidal nanocrystals and their application as a platform for high-performance lithium-ion battery composite electrodes. *Nano Research* 2016, 9, 3757–3771.
15. Lee, S.H.; Kotal, M.; Oh, J.H.; Sennu, P.; Park, S.H.; Lee, Y.S.; Oh, I.K. Nanohole-structured, iron oxide-decorated and gelatin-functionalized graphene for high rate and high capacity Li-ion anode. *Carbon* 2017, 119, 355–364.
16. Esralizadeh, D.; Zavabeti, A.; Jalili, R.; Atkin, P.; Choi, J.; Carey, B.J.; Brkljača, R.; O’Mullane, A.P.; Dickey, M.D.; Officer, D.L. Room temperature CO: reduction to solid carbon species on liquid metals featuring atomically thin ceria interfaces. *Nat. Commun.* 2019, 10, 865.
17. Ahmed, H.; Rezk, A.R.; Carey, B.J.; Wang, Y.; Mohiuddin, M.; Berean, K.J.; Russo, S.P.; Kalantar-zadeh, K.; Yeo, L.Y. Ultrafast Acoustofluidic Exfoliation of Stratified Crystals. *Adv. Mater.* 2018, 30, 1704756.
18. Park, K.H.; Lee, D.; Kim, J.; Song, J.; Lee, Y.M.; Kim, H.T.; Park, J.K. Defect-free, size-tunable graphene for high-performance lithium ion battery. *Nano Lett.* **2014**, *14*, 4306–4313.

19. Hernandez, Y.; Nicolosi, V.; Lotya, M.; Blighe, F.M.; Sun, Z.; De, S.; McGovern, I.; Holland, B.; Byrne, M.; Gun’Ko, Y.K. High-yield production of graphene by liquid-phase exfoliation of graphite. *Nat. Nanotechnol.* **2008**, *3*, 563-568.

20. Chacón-Torres, J.; Pichler, T. Defect modulated Raman response of KCs single crystals. *Phys. Status Solidi b* **2011**, *248*, 2744–2747.

21. Wang, A.; Kadam, S.; Li, H.; Shi, S.; Qi, Y. Review on modeling of the anode solid electrolyte interphase (SEI) for lithium-ion batteries. *npj Comput. Mater.* **2018**, *4*, 15.

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