High-Rate LiTi$_2$(PO$_4$)$_3$@N–C Composite via Bi-nitrogen Sources Doping

Dan Sun, Xia Xue, Yougen Tang, Yan Jing, Bin Huang, Yu Ren, Yan Yao, Haiyan Wang, and Guozhong Cao

†College of Chemistry and Chemical Engineering, Central South University, Changsha, 410083, People’s Republic of China
‡Department of Electrical and Computer Engineering, University of Houston, Houston, Texas 77204, United States
§Department of Materials Science & Engineering, University of Washington, Seattle, Washington 98185, United States
‖Battery Materials, BASF (China) Co., Ltd., Shanghai, 201206, People’s Republic of China
∥State Key Laboratory for Powder Metallurgy, Central South University, Changsha 410083, People’s Republic China

Supporting Information

ABSTRACT: Mesoporous LiTi$_2$(PO$_4$)$_3$@nitrogen-rich doped carbon composites have been synthesized by a novel bi-nitrogen sources doping strategy. Tripolycyanamide (C$_3$H$_6$N$_6$) and urea are proposed for the first time as both nitrogen and carbon sources to achieve a homogeneous nitrogen-doped carbon coating layer via an in situ method. The electrode delivers ultrahigh rate performance and outstanding cycling stability in lithium ion batteries (LIBs). In an organic electrolyte system, the electrode demonstrates high discharge capacities of 120 mAh g$^{-1}$ and 87 mAh g$^{-1}$ at 20C and 50C, respectively. Moreover, 89.5% of initial discharge capacity is retained after 1000 cycles at 10C. When used as an anode for aqueous LIBs, the electrode also demonstrates superior rate capability with the discharge capacity of 103 mAh g$^{-1}$ at 10C, corresponding to 84% of that at 1C. Outstanding cycling stability with capacity retention of 91.2% after 100 cycles at 30 mA g$^{-1}$ and 90.4% over 400 cycles at 150 mA g$^{-1}$ are also demonstrated. The uniform nitrogen-rich carbon coating and unique mesoporous structure play important roles in effectively suppressing the charge-transfer resistance and facilitating Li ion/electron diffusion, thus leading to the superior electrochemical properties.

KEYWORDS: aqueous lithium ion battery, nitrogen doping, mesoporous LiTi$_2$(PO$_4$)$_3$/C, rate performance, cycling performance

INTRODUCTION

During the past decades, LIBs have dominated the portable electronic markets, and they exhibit huge potential for electric vehicles (EVs) and large-scale energy storage systems (ESSs) in the near future. However, the concerns for safety and high cost due to the employment of combustible organic electrolytes greatly narrow their extensive applications. Aqueous lithium ion batteries (ALIBs), which use cheap inorganic salt solution as electrolyte, could fundamentally eliminate the safety issue and rigorous assembly conditions at the same time. Other advantages, such as higher ionic conductivity of electrolyte, more environmental benignity, and lower cost could also be achieved for ALIBs.

Because of the narrow electrochemically stable window of electrolyte, the available electrode materials, especially the anode materials, are seriously confined. Generally, the traditional cathode materials in LIBs like LiFePO$_4$, Li-Ni$_{1/3}$Co$_{1/3}$Mn$_{1/3}$O$_2$, LiCoO$_2$, and LiMn$_2$O$_4$ have been well investigated as the cathodes for ALIBs. However, the anodes for ALIBs are restricted to the electrode materials with Li ion operation potential of 2–3 V versus Li$^+$/Li$^-$. There are only a few suitable candidates, for example, vanadates and Li-Ti$_2$(PO$_4$)$_3$. The first ALIB of VO$_2$/LiMn$_2$O$_4$ was constructed by Dahn et al., cycling for limited cycles. Since then, the ALIB systems, such as LiV$_3$O$_8$/LiMn$_2$O$_4$, NaV$_3$O$_8$/LiMn$_2$O$_4$, LiV$_3$O$_8$/LiNi$_{1/3}$Co$_{1/3}$O$_{2}$, NaV$_2$O$_3$/LiMn$_2$O$_4$ and so on, have been proposed based on vanadates anodes.

Unfortunately, most vanadates demonstrated poor cycling stability which may originate from the degradation of crystal structure and vanadium dissolution in aqueous solution. LiTi$_2$(PO$_4$)$_3$/C is believed to be a promising anode material for ALIB by virtues of its good cycling durability and relatively low working potential. Xia’s group reported a piece of milestone work in ALIBs by eliminating the soluble oxygen in Li$_2$SO$_4$ solution. The cycling life of as-constructed LiTi$_2$(PO$_4$)$_3$/LiFePO$_4$ ALIB at relatively high current rate (6C-rate) was significantly improved. Even so, the performance of ALIBs system, particularly at low current densities, is still far from practical applications. LiMn$_2$O$_4$ has been proved to be a promising cathode for ALIBs benefiting from its low cost, high voltage potential and remarkable cycling stability in aqueous solution. In contrast,
the electrochemical properties of as-reported anodes for ALIBs are much insufficient. Electrochemical properties of pristine LiTi$_2$(PO$_4$)$_3$ are seriously restricted by its low electronic conductivity,$^{13}$ which could be well overcome by size reducing and carbon coating.$^{14}$ It is well-known that high-quality carbon coating can not only largely improve the electronic conductivity of materials but also protect inner material from electrolyte flushing, generating better electrochemical performance.$^{15,16}$ In our previous work, LiTi$_2$(PO$_4$)$_3$ nanoparticles coated with high-quality carbon were prepared, which demonstrated outstanding cycling stability over 300 cycles at 0.2C.$^{17}$ Unfortunately, its rate capability was not very good. Recently, intensive researches have revealed that the electronic conductivity of carbon layers could be improved by surface modification (such as doped with N, B, and O), thanks to the enhanced electron properties, pseudocapacitance effect and extra Li$^+$ storage sites after surface atomic modification.$^{18,19}$ The N-modified composites with significantly enhanced electrochemical properties, such as N-doped graphene,$^{18-20}$ N-doped g-$\text{C}_3\text{N}_4$/MoS$_2$,$^{21}$ N-doped graphene/SnO$_2$,$^{22}$ N-doped nanoporous carbon,$^{23}$ N-doped Sn/C,$^{24}$ N-doped rGO/Sn,$^{25}$ N-doped CNT/LiFePO$_4$,$^{26}$ and N-doped Fe$_3$O$_4$/C$^{27}$ have been reported. As far as we know, LiTi$_2$(PO$_4$)$_3$/C has not been reported yet. In addition, mesoporous materials are attracting more and more attention by virtue of the large contact surface area between electrode and electrolyte allowing both Li$^+$ and electron to migrate rapidly, thus resulting in superior rate performance.$^{28-31}$

Herein, a novel bi-nitrogen sources doping approach is proposed to fabricate mesoporous and nitrogen-rich doped LiTi$_2$(PO$_4$)$_3$/C composite. This strategy is completely different from conventional nitrogen doping methods, in which the NH$_3$ gas or amine was used as a single nitrogen source.$^{23,32,33}$ In our work, tripolycyanamide (C$_3$H$_6$N$_6$) is proposed for the first time as both carbon and nitrogen source to achieve homogeneous nitrogen doped carbon coating layer via an in situ method. Moreover, urea (CO(NH$_2$)$_2$) is also introduced to improve the content of nitrogen in carbon layer and generate massive nanopores at the same time. Accordingly, the nitrogen content in carbon is as high as 11.1%. This strategy possesses characteristics of low cost, available and also permits facile scale-up for mass production. Benefiting from high nitrogen doping content and the unique structure, the as-prepared electrode demonstrates ultrahigh rate capability and outstanding cycling life.

## RESULTS AND DISCUSSION

The XRD patterns of as-obtained LTP-T and LTP-T-U composites are presented in Figure 1a. The two samples demonstrate the same diffraction peaks, which are in great agreement with the rhombohedral NASICON type LiTi$_2$(PO$_4$)$_3$ phase (JCPDS #35-0754) with a R$_3$c space group. The absence of impurity peaks reflects the high purity of LiTi$_2$(PO$_4$)$_3$ phase in the samples. The calculated lattice parameters (Table S1) of LTP-T and LTP-T-U match well with previous reports.$^{34,35}$

The microstructural features of as-prepared composites are displayed in Figure 1. From the TEM picture of LTP-T (Figure 1b), the LTP-T is of individual particles with size range of 50–70 nm. The HRTEM image of LTP-T (Figure 1c) clearly shows that the surface of the LiTi$_2$(PO$_4$)$_3$ particle was coated with an amorphous carbon layer. As deduced from the image, the interplanar spacing is 0.301 and 0.245 nm, corresponding to the d-spacing of the (024) and (300) crystal face of rhombohedral LiTi$_2$(PO$_4$)$_3$, respectively. The N$_2$ adsorption–desorption isotherm of LTP-T is shown in Figure S1a, in which there is a typical IV isotherm, suggesting the existence of mesoporous structure. The BHET surface area of LTP-T is 75.9 m$^2$ g$^{-1}$. Figure 1e gives the TEM image of LTP-T-U, as displayed, the particle size decreases to 30–50 nm. Moreover, better particle dispersion and more obvious pores are observed in comparison with LTP-T. The HRTEM image of LTP-T-U (Figure 1f) displays uniform carbon layer whose thickness is ~5 nm. As reported, the thickness of carbon layer around 4–8 nm on electrode...
materials could reach a good balance of Li\(^+\) diffusion and electron conductivity, promising superior electrochemical properties.\(^\text{15}\) Obviously, it is the in situ coating strategy that creates the uniform carbon layer. In this method, the carbon source could be well coated on the surface of precursor when separating from glycol. The interplanar spacing of 0.269 and 0.276 nm deduced from the HRTEM picture (Figure 1f) corresponds well to the \(d\)-spacing of the (116) and (211) plane of rhombohedral LiTi\(_2\)(PO\(_4\))\(_3\).

The BET surface area of LTP-T-U is 87.6 m\(^2\)·g\(^{-1}\) (Figure S1b), larger than that of LTP-T. Meanwhile, LTP-T-U also displays amounts of pores with larger size range of 5–8 nm and 30–40 nm (Figure 2b) than that of LTP-T (Figure 2a).

![Figure 2. Barrett–Joyner–Halenda (BJH) pore size distribution plot of (a) LTP-T and (b) LTP-T-U.](image)

Obviously, the increase of pore size originates from the addition of urea, which has been widely used as pore-forming agent.\(^\text{36}\) As we know, the pore size significantly affects the rate performance of nanoporous materials.\(^\text{37}\) Generally, the larger pores could enable ions to diffuse into the crystal from the electrolyte more easily, leading to better rate performance.\(^\text{37}\) The carbon contents of LTP-T and LTP-T-U composites are measured by TGA. As recorded in Figure 1e, the amount of coated carbon of LTP-T-U is 6.6 wt %, which is less than that of LTP-T (8.5 wt %). The optimum carbon coating could further increase the electronic conductivity of electrode material without sacrificing tap density and energy density.

X-ray photoelectron spectroscopy (XPS) is applied for the surface element analysis of as-prepared samples. Elements of Ti, C, O, P, and N can be observed in the survey spectrum of both LTP-T (Figure 2a) and LTP-T-U (Figure 2b). Note that the atomic ratio of N/C is 5.4 and 11.1% for LTP-T and LTP-T-U according to the peak area ratio of N 1s to C 1s, respectively. Apparently, the increase of nitrogen content is resulted from the addition of urea. The N/C ratio in LTP-T-U here is higher than those in ref 20 (\(~3.9\%)\), ref 38 (\(~8.9\%)\), ref 39 (\(~5.6\%)\), and ref 40 (\(~10.1\%)\). Apparently, a high quality carbon layer with more nitrogen doping could be achieved via our bi-nitrogen sources doping approach. Although the influence of nitrogen doping to the conductivity of carbon has not been well illustrated, the relatively high content is always expected to be favorable.\(^\text{40}\) The high-resolution XPS spectra of O, P, and Ti of LTP-T and LTP-T-U (Figure S2) are similar and match well the reported LiTi\(_2\)(PO\(_4\))\(_3\).\(^\text{41}\) The high-resolution N 1s XPS spectra of two samples (Figure 3b) can be deconvoluted into four different signals. The peak at \(~402.2\) eV corresponds to the N in N-\(\equiv\)O bond.\(^\text{42}\) The peaks at \(~400.6, \sim 399.3, \sim 398.2\) eV, are attributed to graphitic N, pyrrolic N, and pyridinic N, respectively, as explained in Figure 3d.\(^\text{19}\) Note that the graphic N is dominant in the LTP-T product. While the intensity of graphic N in LTP-T-U decreases clearly in comparison with LTP-T. Theoretical calculations imply that pyrrolic N and pyridinic N doping are more beneficial than graphic N for Li storage.\(^\text{43}\) The high-resolution spectrum of C

![Figure 3. Survey XPS spectra of (a) LTP-T and (b) LTP/C-T-U; high-resolution XPS spectra of (c) LTP-T and (d) LTP/C-T-U; (e) Raman spectra of LTP-T and LTP/C-T-U; and (f) illustration of nitrogen doping carbon with different doping sites.](image)
1s in LTP-T-U could be fitted by several peaks corresponding to O–C–O (289.2 eV), C=O (288.0 eV), C–O (286.7 eV), C–N (285.6 eV), and C–C (284.6 eV) further suggests the presence of C–N (Figure S2). The XPS results also reveal that the N atoms are mainly doped in carbon layers instead of LiTi2(PO4)3. To know more about the structure of the carbon layer, Raman spectroscopy was performed (Figure 3c). The two strong bands around 1600 and 1330 cm\(^{-1}\) could be ascribed to the in-plane vibrations of crystalline graphitic carbon (G band) and disordered amorphous carbon (D band), respectively. The relatively high \(I_D/I_G\) value for LTP-T-U (3.08) indicates a more disordered feature in comparison with that of LTP-T (2.86), which may relate to the defects in carbon because of the more N atoms doping. It is well-known that the homogeneity of nitrogen doping and carbon coating significantly affect the properties of electrode. Therefore, the scanning transmission electron microscope-energy dispersive spectrometer (STEM-EDS) elemental mapping was performed. As seen in Figure 4, the appearance of Ti, P, O, C, and N atoms are almost the same, implying the uniformly nitrogen doping and carbon coating. In summary, a homogeneous and nitrogen-rich carbon layer has been successfully achieved.

**Electrochemical Properties in Nonaqueous LIBs.** First, we performed the electrochemical test of as-prepared samples in organic electrolyte cells. The rate performance of LTP-T-U and LTP-T is compared in Figure 5a. LTP-T delivers a discharge capacity of 131 mAh g\(^{-1}\) at 1C and 89 mAh g\(^{-1}\) at 20C, respectively. While LTP-T-U exhibits much better rate performance with a discharge capacity of 128 mAh g\(^{-1}\) at 1C and 120 mAh g\(^{-1}\) at 20C, respectively. When the charge/discharge rate is increased to 40C and 50C, LTP-T-U still delivers 101 and 87 mAh g\(^{-1}\), respectively. In other words, 1.5 min charge can retain a high capacity of 101 mAh g\(^{-1}\). Note that when the current density returns to low value (1C) after high rate test, the capacity could be fully recovered to the original value, revealing that the electrode could be tolerant to ultrafast charge/discharge process. To our best knowledge, no literature has reported such high rate capability for LiTi2(PO4)3/C electrode over 10C. So far the best result for LiTi2(PO4)3/C electrode in organic electrolyte is 103 mAh g\(^{-1}\) at 10C rate, while our material demonstrates the same capacity under 3 times higher current density (40C). Moreover, although the theoretical capacity of LiTi2(PO4)3 is relatively low (138 mAh g\(^{-1}\)), the discharge capacity of LTP-T-U at extremely high current density here is higher than those of graphene-modified LiFePO4 (~86 mAh g\(^{-1}\) at 30C), N-CNTs modified LiFePO4 (~50 mA g\(^{-1}\) at 10C), TiN modified Li4Ti3O12 (~70 mAh g\(^{-1}\) at 20C), LiMn2O4 porous nanorods (~83 mAh g\(^{-1}\) at 30 C), Na1.08V3O8 nanosheets (~72.6 mAh g\(^{-1}\) at 50C). Figure 5b displays the charge/discharge curves of LTP-T-U at various rates. There are a short platform at ~2.75 V (vs Li+/Li) and a long voltage platform at ~2.45 V (vs Li+/Li) at 0.5C. When raising the current rate, flat platforms are well maintained despite of the increase of overpotential. Moreover, remarkable durability at extremely high current density was observed for LTP-T-U. As demonstrated in Figure 5c, LTP-T-U delivers an initial capacity...
of 122 mAh g⁻¹ at 10C and a capacity retention of 89.5% after 1000 cycle. This implies that the LiTi₂(PO₄)₃@C electrode here could tolerate rapidly charge/discharge test without obvious structure degradation after long cycles, which is a desirable characteristic for high power applications. Figure 5d presents the charge/discharge curves of LTP-T-U after different cycles. On the cycles increasing, the voltage platform remains well, though the main platform shrinks and polarization increases slightly, which might be related to the slight capacity fading.

Figure 6a shows the CV curves of LTP-T and LTP-T-U at a scanning rate of 0.2 mV s⁻¹, in which similar redox peaks at 2.3/2.6 V and 2.8/2.9 V are demonstrated, respectively. While the polarization potential of the LTP-T-U is lower than that of LTP-T. Additionally, LTP-T-U has narrower half peak width. These facts indicate better charge transfer kinetics of LTP-T-U and correspond well with its better rate performance. Figure 6b displays the CV curves of LTP-T-U at different sweep rates. As seen, the polarization potential increases with the increasing of sweep rate. At the high scan rate of 0.8 mV s⁻¹, the polarization potential increases with the increasing of scan rate. The square root of scan rate (ν¹/₂) of LTP-T-U is demonstrated in Figure 6c. As seen, the Iₚ/Iₚc is a significant indicator to reveal the reversibility of the Li⁺ insertion/extraction reaction. For a Nernstian wave with a stable product, Iₚ/Iₚc should be equal to unity, and the electrode reversibility is better when the Iₚ/Iₚc is close to unity. Table S2 lists the Iₚ/Iₚc values of the sample LTP-T-U at different sweep rates. The small values confirm its good Li⁺ diffusion kinetics. The relationship between the Iₚ and the ν¹/₂ for LTP-T-U is demonstrated in Figure 6c. As seen, the Iₚ vs ν¹/₂ for the sample approaches to a straight line, indicating the semidiffusion controlled process of the Li⁺ insertion into the sample, which could be regarded as a quasi-reversible system.

To further evaluate the electrochemical interfacial properties, we performed EIS tests of LTP-T-U and LTP-T. The Nyquist plots (Figure 6d) of activated LTP-T and LTP-T-U electrodes after 5 cycles demonstrate two depressed semicircles and a slope line. Generally, the small intercept represents the solution impedance of the cells (R₀). The first semicircle in the high frequency (Rₑ) relates to the interface parameters like porous mature of electrode, the bulk of materials, or surface film contribution, and the semicircle in the mediate frequency is due to the charge-transfer resistance (Rₓ). The slope line at low frequency reflects the Warburg impedance (Zₘ), implying the Li⁺ transfer in the solid electrode. Clearly, the Rₑ and Rₓ of the LTP-T-U electrode are smaller than those of the LTP-T (Table S3). The suppressed impedance is clearly related to the richer mesoporous structure that facilitates the Li⁺ diffusion between electrode and electrolyte. Meanwhile, the remarkably improved electronic conductivity by the uniform nitrogen-rich carbon layer also contributes to the decrease of impedance. The four-point probe test results demonstrate that the electronic conductivity of LTP-T-U (2.86 × 10⁻² S cm⁻¹) is closed to eight times higher than that of LTP-T (3.29 × 10⁻³ S cm⁻¹). Apparently, the superior electrochemical properties of LTP-T-U are mainly attributed to the homogeneous nitrogen-rich carbon coating as well as the unique mesoporous structure.

Electrochemical Properties in ALIB. Considering the superior electrochemical properties of LTP-T-U in organic electrolyte, the electrochemical behavior in aqueous electrolyte is also investigated. At first, a three-electrode CV method was employed to investigate the Li⁺ ion insertion/extraction behaviors of individual LTP-T-U and LiMn₂O₄, respectively. As displayed in Figure S3, LTP-T-U shows two redox couples at −0.84/−0.72 V and −0.43/−0.42 V versus SCE. Good lithium insertion/extraction behaviors with two redox couples at 0.80/0.78 V and 0.92/0.90 V vs SCE were also observed for
LiMn$_2$O$_4$. Excellent kinetics behaviors demonstrate the promising employment of LTP-T-U and LMO as anode and cathode for ALIB, respectively. The full cell rate performance of LTP-T-U//LiMn$_2$O$_4$ is shown in Figure 7a. As seen, the LTP-T-U demonstrates superior rate capability with a discharge capacity of 123 mAh g$^{-1}$ at 150 mA g$^{-1}$ (based on the mass of LiTi$_2$(PO$_4$)$_3$) and 103 mAh g$^{-1}$ at 1500 mA g$^{-1}$. Figure 7a (inset) gives the corresponding charge/discharge curves of LTP-T-U//LiMn$_2$O$_4$ ALIB at different rates. A long voltage plateau around 1.55 V and a short plateau around 1.0 V are observed at low current densities. Although the discharge voltage decreases slightly on increasing current density, the potential plateau at a high rate of 10C remains well. We did not evaluate much higher rate properties of LTP-T-U as an anode in ALIBs since it is depended on both cathode and anode materials. Note that the rate capability of the used commercial LiMn$_2$O$_4$ (Figure S4) is not so good as LTP-T-U (Table S4). So the rate capability of the full cell would be limited by the cathode in this situation.

Long-term cycling performance of ALIB at various rates was performed. As shown in Figure 7(b–d), the ALIB demonstrates outstanding cycling stability at different rates. At a low rate of 0.2C, it delivers a discharge capacity of 123 mAh g$^{-1}$, and capacity retention of 91.2% after 100 cycles (Figure 6b). As the rate increases to 1C, capacity retention of 90.4% after 400 cycles is demonstrated (Figure 6c). At 5C, LTP-T-U still gives a capacity retention rate of 84.0% even after 1000 cycles (Figure 7d). These results clearly demonstrate that LTL-T-U can be tolerant to various charge/discharge rates. As we know, the poor cycling stability at low current densities is still a critical challenge for ALIBs. The crystal deterioration of electrode, reaction between electrode materials and water or O$_2$, and decomposition of water are considered the main causes for the capacity fading.^{5,7} The cycling performance of ALIB here at low rate (e.g., 0.2C) is much superior to most of the reported ALIBs using vanadium oxides, vanadates or LiTi$_2$(PO$_4$)$_3$/C to date (Table S5).^{5,7,9,11} Note that LiTi$_2$(PO$_4$)$_3$/C with capacity retention of 90% after 300 cycles was also reported by our group.^{17} Low Coulombic efficiency owing to the reactions between electrode surface and aqueous electrolyte and the decomposition of water, is considered as a significant cause of capacity fading for ALIB. As demonstrated, the ALIB here delivers high average Coulombic efficiency at various current densities (>95% at 0.2C and >99% at 1C in Figure 7b,c), in good accordance with the superior cycling performance. The excellent electrochemical properties of LTP-T-U in ALIB indicate that this nitrogen-rich carbon coated LiTi$_2$(PO$_4$)$_3$ composite with unique mesoporous structure could be used as a promising anode for high-power ALIB.

According to the above discussion, the excellent electrochemical properties of LTP-T-U are ascribed to the following factors: (1) The unique mesoporous structure. This structure has been confirmed to be superior to the common ones (e.g., bulk, nanoparticles) with the advantages of larger BET surface, and more abundant nanopores could enlarge the electrolyte/electrode interfacial area, shorten Li$^+$ diffusion distance, relieve the mechanical lattice strain upon Li$^+$ insertion/extraction, and so on. (2) Homogeneous nitrogen-rich carbon coating. It is well accepted that the electrochemical performance of electrode greatly depends on the state of surface carbon layer (e.g., thickness, uniformity, composition, degree of ordering, electron cloud distribution, etc.). Undoubtedly, the well-distributed carbon layer with suitable thickness and the high-content pyridinic- and pyrrolic-type nitrogen in carbon layer could contribute significantly to the electrochemical properties. It has been confirmed that the hybridization of lone pair electrons of N atom with the $\pi$ electrons in carbon will facilitate the electron diffusion, leading to higher electronic conductivity.\(^{40}\)
CONCLUSIONS

In summary, a bi-nitrogen sources doping method was proposed for the first time to achieve mesoporous LiTi$_2$(PO$_4$)$_3$@N-rich doped carbon composite. In organic LiBs, the electrode delivered ultrahigh rate performance with a discharge capacity of 101 mAh g$^{-1}$ at 40C and 87 mAh g$^{-1}$ at 50C. When used as an anode for ALIB, striking long cycling performance was also presented. The homogeneous nitrogen-rich carbon coating and the unique mesoporous structure have been verified as important factors. By virtue of its superior electrochemical performance, this composite prepared in the present study could meet the requirement of fast charge/discharge for LiBs and ALIBs. This bi-nitrogen source doping strategy may provide a reference for the fabrication of other electrode materials with nitrogen-rich carbon coating, including Li$_4$Ti$_5$O$_{12}$/C, LiV$_2$(PO$_4$)$_3$/C, LiFePO$_4$/C, and so on.

METHODS

Synthesis of LTP-T and LTP-T Composite. First, 0.887 g of phosphoric acid (H$_3$PO$_4$) and 0.1 g of CO(NH$_2$)$_2$ were dissolved in ethanol. Then, 0.2 g of C$_3$H$_6$N$_6$ was dissolved in glycol with a heating rate of 10$^°$C/min. The four-point probe method was used to measure the electronic conductivity (HORIBA Jobin Yvon) spectrometer was employed to record Raman (HORIBA Jobin Yvon) radiation (Dandong Haoyuan, DX-2008 S-TWIX TEM. Scanning electron microscope (SEM) pictures were taken using XPSPEAK software.

Electrochemical Measurements. The electrochemical tests were all performed in a CR2016-type coin cell. For the fabrication of electrode, the active material, Polytetrafluoroethylene (PTFE), and Super P carbon in a weight ratio of 8:1:1 was mixed first, and distilled water was used as solvent. Then, the mixture was pressed on a stainless steel mesh collector under 20 MPa. After drying at 110 $^°$C under vacuum overnight, the electrodes, whose mass loading of the active material is about 6 mg cm$^{-2}$, were assembled into CR2016 coin-type cells in an Ar-filled Mikrouna glovebox using Li metal as counter electrode. The electrolyte was 1 M LiPF$_6$ in 1:1 v/v ethylene carbonate/dimethyl carbonate (Guangzhou Tinci Materials Technol-ogy Co., Ltd.). The cells were cycled galvanostatically between 1.5 and 3.5 V (vs Li$^+/Li$) using a Neware battery testing system (CT-3008W) at a constant temperature of 30 $^°$C. Cyclic voltammetry (CV) curves were performed by CHI 660d electrochemical station (Shanghai Chenhua, China). Electrochemical impedance spectroscopy (EIS) was tested in a Princeton workstation (PARSTAT2273, EG&G, Salem, MA) over the frequency range from 100 kHz to 10 mHz with an amplitude of 5 mV. Before EIS test, the cells were charged to 2.5 V and then kept at that voltage via a constant voltage mode.

The electrochemical properties in aqueous electrolyte was also performed in a CR2016-type coin, which is composed of commercial LiMn$_2$O$_4$ as cathode, 2 mol L$^{-1}$ Li$_2$SO$_4$ solution as electrolyte and as-prepared LiTi$_2$(PO$_4$)$_3$/C as anode. The LiMn$_2$O$_4$ electrodes, which were purchased from Hunan Reshine New Material Co., Ltd., were made in a similar way as LiTi$_2$(PO$_4$)$_3$/C. An excessive LiMn$_2$O$_4$ with cathode/anode mass ratio of (1.5–2.0)/1 was designed to better evaluate the electrochemical properties of LiTi$_2$(PO$_4$)$_3$.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acsami.5b08697. Lattice parameters, N$_2$ adsorption—desorption isotherm, high-resolution XPS images, $I_p$/I$_0$ value, and impedance values of LTP-T and LTP-T-U. Rate capability of commercial LiMn$_2$O$_4$.

AUTHOR INFORMATION

Corresponding Authors

*E-mail: wanghy419@126.com.
*E-mail: gzcao@u.washington.edu.

Notes

The authors declare no competing financial interest.

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