Off-stoichiometric Na\textsubscript{3-3x}V\textsubscript{2+x}(PO\textsubscript{4})\textsubscript{3}/C nanocomposites as cathode materials for high-performance sodium-ion batteries prepared by high-energy ball milling†

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Na\textsubscript{3}V\textsubscript{2}(PO\textsubscript{4})\textsubscript{3} (NVP) is regarded as a promising cathode material for sustainable energy storage applications. Here we present an efficient method to synthesize off-stoichiometric Na\textsubscript{3-3x}V\textsubscript{2+x}(PO\textsubscript{4})\textsubscript{3}/C (x = 0–0.10) nanocomposites with excellent high-rate and long-life performance for sodium-ion batteries by high-energy ball milling. It is found that Na\textsubscript{3-3x}V\textsubscript{2+x}(PO\textsubscript{4})\textsubscript{3}/C nanocomposites with x = 0.05 (NVP-0.05) exhibit the most excellent performance. When cycled at a rate of 1C in the range of 2.3–3.9 V, the initial discharge capacity of NVP-0.05 is 112.4 mA h g\(^{-1}\), which is about 96% of its theoretical value (117.6 mA h g\(^{-1}\)). Even at 20C, it still delivers a discharge capacity of 92.3 mA h g\(^{-1}\) (79% of the theoretical capacity). The specific capacity of NVP-0.05 is as high as 100.7 mA h g\(^{-1}\) after 500 cycles at 5C, which maintains 95% of its initial value (106 mA h g\(^{-1}\)). The significantly improved electrochemical performance of NVP-0.05 is attributed to the decrease of internal resistance and increase of the Na\textsuperscript{+} ion diffusion coefficient.

1. Introduction

Developing clean and renewable energy has become one of the most important scientific and engineering tasks because of increasing environmental concerns and energy shortages.\(^{1,2}\) Among all energy storage systems, lithium ion batteries (LIBs) have been regarded as the best choice because of their high voltage, high energy density, and environmental friendliness.\(^{3-5}\) However, the ability of LIBs has been doubted due to the ever increasing demand and limited lithium reserves in the Earth’s crust. Sodium ion batteries (SIBs) have attracted more attention due to their attractive features, such as high safety, low cost, abundant sodium resources, etc.\(^{6,7}\) Na\textsubscript{3}V\textsubscript{2}(PO\textsubscript{4})\textsubscript{3} (NVP) has been considered as a promising SIB cathode material with an open NASICON framework wherein the sodium ions could diffuse quickly through the interstitial sites reversibly without resulting in a large volume change of the crystalline structure.\(^{8-10}\) These features make NVP a very promising potential cathode material for high-performance SIBs. However, NVP has very low electronic and Na\textsuperscript{+} ion conductivity.\(^{11-13}\) The theoretical capacity is not fully available even at comparatively low rates, which is a major drawback for practical applications. Therefore, many efforts have been made to improve the charge carrier transport kinetics in NVP by modifying NVP by carbon coating and ion doping.\(^{14-18}\) Carbon coating is a common method to improve the electronic conductivity of cathode materials. And it can keep the structural integrity of the electrode materials during the repeated charge and discharge processes, which greatly benefits cycle stability. Recently, many studies on carbon wrapped Na\textsubscript{3}V\textsubscript{2}(PO\textsubscript{4})\textsubscript{3} composites have been reported.\(^{19-21}\) In addition, doping with metallic ions has been proved to be an effective way in improving the intrinsic electronic conductivity in SIBs.\(^{22-26}\) Recently, the effects of off-stoichiometry have been widely studied in cathode materials. Byoungwoo Kang \textit{et al.} proposed that a fast ion-conducting surface phase is formed by controlling the off-stoichiometry in LiFePO\textsubscript{4}.\(^{27}\) The fast kinetic response induced by off-stoichiometry is attributed to the decrease of the battery resistance. Maryia J. Aragón \textit{et al.} studied the effect of sodium excess on Na\textsubscript{3+x}V\textsubscript{2-x}(PO\textsubscript{4})\textsubscript{3} (0 ≤ x ≤ 0.15) as cathode materials for SIBs.\(^{28}\) The synergetic combination of the presence of Na\textsubscript{3}P\textsubscript{2}O\textsubscript{7} impurities and the structure deviation lead to an improvement of the electrochemical performance, especially cycling stability at high rate, compared with the stoichiometric NVP. In the following work, they prepared Na\textsubscript{3-3x}V\textsubscript{2+x}(PO\textsubscript{4})\textsubscript{3} by sol–gel method to detect the effect of vanadium excess on the structure, morphology, and...
electrochemical properties. The rate performance of Na$_{3-x}$V$_{2x+3}$(PO$_4$)$_3$ is significantly improved than that of Na$_{3-y}$V$_{2-y}$ (PO$_4$)$_3$. However, both the high rate performance and cycling stability need further improved, especially the capacity decreased fast even after 300 cycles. High-energy ball milling has been widely used in the field of LIBs and SIBs. In addition, the ball milling process represents a low cost process in a short time and creates fresh surfaces with defects on the surface of the powders, resulting in the modification of the electrochemical performance of electrode material. In this work, we adopt the high-energy ball milling to prepare the off-stoichiometric Na$_{3-x}$V$_{2x+3}$(PO$_4$)$_3$. The high rate performances and more stable cycle properties have been achieved, which is attributed to the decreased internal resistance and increase of Na$^+$ ion diffusion coefficient.

2. Experimental

2.1. Synthesis of Na$_{3-x}$V$_{2x+3}$(PO$_4$)$_3$ (x = 0–0.10) composites (NVP-x)

NVP-x composites were synthesized by solid state reaction. 5 mmol NVP-x composites were synthesized by mixing Na$_2$CO$_3$ (AR, ≥99.8%), NH$_4$VO$_3$ (AR, ≥99.0%), NH$_3$H$_2$PO$_4$ (AR, ≥99.0%) and citric acid (AR, ≥99.5%) with molar ratio of 1.5:1.5x : 2 + x : 3 : 2 (For NVP-0, 0.7949 g Na$_2$CO$_3$, 1.1698 g NH$_4$VO$_3$, 1.7253 g NH$_3$H$_2$PO$_4$ and 2.1014 g citric acid; for NVP-0.05, 0.7552 g Na$_2$CO$_3$, 1.1990 g NH$_4$VO$_3$, 1.7253 g NH$_3$H$_2$PO$_4$ and 2.1014 g citric acid; for NVP-0.10, 0.7154 g Na$_2$CO$_3$, 1.2283 g NH$_4$VO$_3$, 1.7253 g NH$_3$H$_2$PO$_4$ and 2.1014 g citric acid). Then the mixtures were blended and high-energy ball milled for 30 minutes in ethanol media. The mixtures were dried at 80 °C for 12 h and then transferred to a quartz tubular furnace and heated at 400 °C for 6 hours in Ar flow. After that, the precursors were heated in high-energy ball milled again, and finally calcined at 750 °C for 8 hours in Ar flow.

2.2. Structural characterizations

X-ray diffraction (XRD, Rigaku Smartlab3) was employed to characterize the structural properties of the samples. Scanning electron microscope (SEM) and transmission electron microscope (TEM) images were recorded by FEI Inspect F50 equipped with an energy dispersive X-ray spectroscopy (EDS) and FEI Tecnai F20, respectively. X-ray photoelectron spectroscopy (XPS) was carried out by an X-ray photoelectron spectrometer (Thermo Fisher SCIENTIFIC) with Al Kα X-ray source (hv = 1486.6 eV).

2.3. Electrochemical characterizations

The electrochemical performances were measured using CR2025 coin cells assembled in a glove box filled with high pure argon gas. The electrodes were prepared by putting the mixture of 80 wt% NVP-x, 10 wt% acetylene black and 10 wt% polyvinylidene fluoride (PVDF) binder in an appropriate amount of N-methylpyrroliodine (NMP). The slurry of mixture was coated onto an aluminum foil by a manually adjustable film-coating equipment and then dried in a vacuum oven for 12 hours at 120 °C. Pure sodium metal was used as anode, glass fiber membrane as separator, 1 mol L$^{-1}$ NaClO$_4$ in ethylene carbonate (EC): propylene carbonate (PC) (1 : 1 v/v ratio) as the electrolyte. Galvanostatic charging/discharging tests were measured at a voltage window of 2.3–3.9 V (vs. Na/Na$^+$) on a NEWARE battery testing system at room temperature. Cyclic voltammetry (CV) and electrochemical impedance spectroscopy (EIS) spectra were tested with CHI 660A electrochemical analyzer workstation (Chenhua, Shanghai, China).

3. Results and discussion

The diffraction peaks can be indexed to the rhombohedral NVP phase (PDF#53-0018), agreeing well with the previously reported literatures. No other phases appear in NVP-x (x = 0–0.10), confirming the high phase purity. It is observed that a slight shift to smaller angle of diffraction peaks with increasing the deficient Na and excess V concentrations (Fig. 1b), indicating the slight expansion of unit cell. The cell parameters are calculated to be $a = 8.717$ Å, $c = 21.808$ Å, $V = 1435.2$ Å$^3$ for NVP-0, and $a = 8.726$ Å, $c = 21.814$ Å, $V = 1438.4$ Å$^3$ for NVP-0.05 and $a = 8.748$ Å, $c = 21.858$ Å, $V = 1448.7$ Å$^3$ for NVP-0.10 (Table 1). This slight expansion of lattice may make the Na$^+$ ions diffusion into and out of NVP-x particles smoother, inducing the increase of ionic mobility and enhancement of high-rate charge-discharge characteristics. However, the excess V ions may locate at the interstitial sites, instead of the lattice sites, which will impede the transportation of Na$^+$ ions and the deterioration of high-rate charge-discharge characteristics. These two effects may compete with each other and contribute to the cell performance.

Morphological study on NVP-x (x = 0–0.10) has been performed by SEM, and the images are shown in Fig. 2. No significant difference can be seen and all samples show good uniformity in particle shape and size. TEM was conducted to further study the particle morphologies of NVP-x nanocomposites, shown in Fig. 3a–e. It can be seen that the particle size is in a broad range, and the biggest particle sizes of all samples are about 500 nm. No significant dependence on the off-stoichiometric concentration is observed. All the NVP-x particles are separated by amorphous carbon shells with thickness of around 3 nm (Fig. 3d–f). The HRTEM images (Fig. 3g–i) show clear lattice fringes with d spacing of 0.3414,
0.2772 and 0.3831 nm, which correspond to the (024), (116) and (113) planes of rhombohedral NVP, respectively. EDS results of NVP-\(x\) particles are shown in Table 2. It is observed that the actual Na/V, Na/P and V/P ratio are close to that of designed Na/V, Na/P and V/P ratio.

XPS was conducted to probe the valence state of V in NVP-\(x\). V 2p XPS peaks of NVP-\(x\) are shown in Fig. 4a. Two peaks at 515.93 eV and 523.16 eV can be observed for NVP-0, which are attributed to V 2p\(_{3/2}\) and V 2p\(_{1/2}\), respectively. The binding energies are ascribed to V\(^{3+}\), in well agreement with previous reported Na\(_3\)V\(_2\)(PO\(_4\))\(_3\).\(^{39}\) It is observed that V 2p peaks of NVP-0.05 and NVP-0.10 shift to higher binding energy, compared with that of NVP-0, indicating that the valence state of V transfers from V\(^{3+}\) to V\(^{4+}\), which is related to the vanadium locating in a different local environment.\(^{40}\) Since P keeps the valence state of +5, no shift of binding energy can be observed, as shown the XPS spectra in Fig. S1 (ESI†).

The capacity of NVP-0.05 is 112.4 mA h g\(^{-1}\) at 1C, which is about 96% of the theoretical value (117.6 mA h g\(^{-1}\)).\(^{2}\) On the contrary, the discharge capacities of NVP-0 and NVP-0.10 are 97.2 and 104.2 mA h g\(^{-1}\), respectively. Fig. 5b shows the rate performance comparisons of the three nanocomposite electrodes. When cycled at a rate of 0.5C in the window of 2.3–3.9 V, the initial discharge capacity of NVP-0.05 is 114.9 mA h g\(^{-1}\), much higher than that of the other two. The
Fig. 3 TEM images of NVP-\(x\) under different magnifications, (a, d and g) \(x = 0\), (b, e and h) \(x = 0.05\), (c, f and i) \(x = 0.10\).

Table 2 Na/V, Na/P, and V/P ratios of NVP-\(x\) (\(x = 0–0.10\))

| Samples NVP-\(x\) | Na/V Calculated ratio | Measured ratio | Na/P Calculated ratio | Measured ratio | V/P Calculated ratio | Measured ratio |
|------------------|------------------------|----------------|------------------------|----------------|----------------------|----------------|
| \(x = 0\)       | 1.50                   | 1.48           | 1.00                   | 1.02           | 0.67                 | 0.69           |
| \(x = 0.05\)    | 1.39                   | 1.31           | 0.95                   | 1.00           | 0.68                 | 0.70           |
| \(x = 0.10\)    | 1.29                   | 1.29           | 0.90                   | 0.93           | 0.70                 | 0.72           |

Fig. 4 XPS core level spectra of NVP-\(x\) (\(x = 0–0.10\)), (a) V 2p, (b) C 1s.
NVP-0.05 cathode exhibits the smallest capacity decay at each rate. Even at 20C, a discharge capacity of 92.3 mA h g\(^{-1}\) (79% of the theoretical capacity) is still delivered, much higher than that of NVP-0 cathode (84.9 mA h g\(^{-1}\)), indicating the good high-rate electrochemical performance. After rapid changes of the current rate, the NVP-0.05 cathode shows stable capacities at each state. When rate comes back to 0.5C, the capacity of 108.2 mA h g\(^{-1}\) for NVP-0.05 cathode is recovered. Long-term cycling stabilities of samples are subsequently shown in Fig. 5c and d. The NVP-0.05 cathode shows an initial discharge capacity of 112.4 mA h g\(^{-1}\), and delivers a remarkable capacity of 109.5 mA h g\(^{-1}\) after 100 cycles at 1C, corresponding to capacity retention of 97%. Even after 500 cycles at 5C, the specific capacity of NVP-0.05 is still as high as 100.7 mA h g\(^{-1}\), which maintains 95% of its initial capacity. A comparison between our work and the related works is listed in Table 3. It can be clearly seen that the incorporation of high-energy ball milling during the sample preparation significantly improve the high rate performance and cycle stability. SEM experiments were done to compare the morphological changes of the NVP-x electrodes before cycling and after 100 cycles at 1C. As shown in Fig. S2 (ESI†), there is no apparent difference among the SEM images of the electrode before cycling and after 100 cycles at 1C. Before cycling, a compact electrode consisting of the NVP-x particles and acetylene black are observed. After 100 cycles at 1C, the electrodes still maintain integral. Thus, the results suggested that the structures of NVP-x are stable and the amorphous carbons are helpful to assist in buffering volume change of Na ion insertion/extraction.

**Table 3** A comparison between our work and the related works

| Sample                                          | 2C   | 5C   | 20C  | Cycle number | References |
|-------------------------------------------------|------|------|------|--------------|------------|
| \(\text{Na}_3\text{V}_2\text{(PO}_4\text{)}_3\) made by sol–gel | 95   | 88   | 74(67) | 400          | 28         |
| \(\text{Na}_3\text{V}_2\text{(PO}_4\text{)}_3\) made by sol–gel | 105(75) | 96   | 70   | 300          | 29         |
| \(\text{Na}_3\text{V}_2\text{(PO}_4\text{)}_3\) made by ball mill | 107  | 106(101) | 92   | 500          | This work  |
| \(\text{Na}_3\text{V}_2\text{(PO}_4\text{)}_3/C\) made by self combustion | —    | 65   | 65   | —            | 45         |
| Porous \(\text{Na}_3\text{V}_2\text{(PO}_4\text{)}_3/C\) made by sol–gel | —    | 98(87) | —    | 100          | 43         |
| \(\text{Na}_3\text{V}_2\text{(PO}_4\text{)}_3@C\) nanofiber made by electrospinning | 110(97) | —    | —    | 30           | 44         |
| \(\text{Na}_3\text{V}_2\text{(PO}_4\text{)}_3@C\) made by hydrothermal assisted sol–gel | 102  | 95(91) | 91   | 700          | 4          |
To further understand the relationship between electrode kinetics for NVP-x nanocomposites and electrochemical performance, CV and EIS were carried out. The CV profiles of different scan rate for the NVP-x (x = 0–0.10) sodium-ion battery without pre-cycling are displayed in Fig. 6 in the voltage range 2.3–3.9 V vs. Na/Na⁺. The reduction (Na⁺ insertion) peaks and oxidation (Na⁺ extraction) peaks are located at 3.1 and 3.7 V, respectively.45,46 The current of NVP-0.05 is the highest among the three samples, implying its highest reversibility and fastest kinetics during electrochemical reaction. The correlations between the peak current densities $I_p$ and the square root of the scan rate $v^{1/2}$ as presented in Fig. 6d, indicate a diffusion-controllable process for electrode reaction. The corresponding Na⁺ ion diffusion coefficients $D_{Na^+}$ of the three electrodes are calculated by the Randles–Sevcik equation:45

$$I_p = 2.69 \times 10^5 n^{3/2} A C_0 D^{1/2} v^{1/2}$$

where $I_p$ is the peak current, $n$ is the number of electrons transferred of the reaction, $A$ is the surface area of the electrode, $C_0$ is the bulk concentration of sodium ions in the electrode (2.35 × 10⁻⁶ mol cm⁻³), $D$ is the diffusion coefficient of Na⁺, and $v$ is the scanning rate. From the slope of the fitting line, the apparent diffusion coefficient $D$ of NVP-0.05 is 1.77 × 10⁻⁹ cm² s⁻¹, higher than those of NVP-0 (1.00 × 10⁻⁹ cm² s⁻¹) and NVP-0.10 (1.23 × 10⁻⁹ cm² s⁻¹), respectively.

Fig. 7 shows Nyquist plots of impedance. The impedance spectra of all electrodes consist of a single depressed semicircle in the high frequency region and an inclined line at low frequency, representing the charge-transfer resistance ($R_{ct}$) and the solid-state diffusion of ions in the active materials, respectively. The NVP-0.05 electrode shows the smallest diameter for semicircle in the Nyquist plots than those of NVP-0 and NVP-0.10, indicating the lowest $R_{ct}$ and better electronic conductivity. The fitting parameters are shown in Table 4.

![Fig. 6 Cyclic voltammogram of NVP-x (a) x = 0, (b) x = 0.05, (c) x = 0.10 at different scan rates, (d) linear fitting of $I_p$ vs. $v^{1/2}$ curves.](image)

![Fig. 7 Nyquist plots of NVP-x (x = 0–0.10) before cycles.](image)

| Samples | $R_e$ (Ω) | $R_{ct}$ (Ω) |
|---------|----------|-------------|
| NVP-0   | 4.704    | 1040        |
| NVP-0.05| 3.807    | 441.5       |
| NVP-0.10| 3.943    | 645.4       |

Table 4 Fitting parameters of NVP-x (x = 0–0.10)
Thus, the excellent electrochemical performance of NVP-0.05 is ascribed to the increased $\text{Na}^+$ ion conductivity and decreased resistance. Firstly, lattice volume expansion may make the $\text{Na}^+$ ions diffusion into and out of NVP-0.05 particles easier, leading to the increase of ionic mobility and enhancement of high-rate charge–discharge characteristics. With increasing $V$ excess to 0.10, though the lattice expansion is larger, too many $V$ ions locate at the interstitial sites instead of lattice sites will impede the transportation of $\text{Na}^+$ ions, leading to the deterioration of electrochemical performance. Secondly, the $\text{Na}$ deficiency and $V$ excess may increase the valence state of $V$ ions from +3 towards +4. The proper tuning of the concentration of $V^{3+}$ and $V^{4+}$ may facilitate the electron transport between the $V^{3+}$ and $V^{4+}$ ions through the double exchange interaction, leading to the decrease of resistance. The above structural characterizations demonstrate that $\text{Na}$ deficiency and $V$ excess with $x = 0.05$ is the most appropriate which may induce the highest $\text{Na}^+$ diffusion coefficient and electronic resistance, leading to the best cell performance.

4. Conclusions

In summary, we have developed a simple method using high-energy ball milling to prepare off-stoichiometric $\text{Na}_{3-x}\text{V}_{2.1+x}^3\text{PO}_{12}\text{C}$ ($x = 0–0.10$) nanocomposites with excellent high rate and cycle performance. Among them, $\text{Na}_{3-x}\text{V}_{2.1+x}^3\text{PO}_{12}\text{C}$ nanocomposites with $x = 0.05$ exhibit the best battery performance. The capacity of NVP-0.05 is 112.4 mA h g$^{-1}$ at 1C, which is about 96% of its theoretical value. Long-term cycling performance is conducted at the rate of 5C. Even after 500 cycles, the specific capacity of NVP-0.05 is still as high as 100.7 mA h g$^{-1}$, which maintains 95% of its initial capacity. The significant improved electrochemical performance of NVP-0.05 has been characterized to be due the decrease of internal resistance and increase of $\text{Na}^+$ ion diffusion coefficient.

Conflicts of interest

There are no conflicts to declare.

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