Hydrolysis Precipitation-assisted Solid-state Reaction to Li$_4$Ti$_5$O$_{12}$ and its Electrochemical Properties

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Abstract: Spinel lithium titanate (Li$_4$Ti$_5$O$_{12}$) materials were synthesized by a hydrolysis precipitation-assisted solid-state method in the temperature range from 600 to 900 °C for large-scale production. DSC/TGA, XRD and SEM were used to characterize the as-prepared samples. The optimum synthesis condition was examined in relation to the charge–discharge performance. It was found that when the dry hydrolysis precipitation precursor with 8% Li excess was calcined at 700–800 °C for 12 h in air, a pure Li$_4$Ti$_5$O$_{12}$ phase was obtained. The as-obtained material has the best electrochemical performance due to its narrow size distribution and precise stoichiometry of the oxide.

Keywords: lithium-ion battery; electrochemical performance; Li$_4$Ti$_5$O$_{12}$

1. INTRODUCTION

In recent years, spinel Li$_4$Ti$_5$O$_{12}$ is being considered as one promising alternative to graphite to be an anode material for lithium-ion batteries. Li$_4$Ti$_5$O$_{12}$ has an excellent cycle performance due to its small volume change (the so-called zero-strain material) during charge–discharge cycling [1], excellent lithium insertion–extraction reversibility and a high theoretical capacity of 175 mAh.g$^{-1}$. It has very flat charge and discharge curves and the potential for Li-insertion is about 1.55 V vs. Li$^+$/Li [1-3], which is above the potential range where most electrolytes are reduced. This makes Li$_4$Ti$_5$O$_{12}$ a promising anode material of lithium ion batteries for safety concerns [4]. Although its medium level of discharge voltage discourages the application as a positive electrode, Li$_4$Ti$_5$O$_{12}$ can be coupled with a high-voltage electrode such as LiMn$_2$O$_4$, LiCoO$_2$, LiCoPO$_4$ or LiFePO$_4$ to provide a cell with an operating voltage above 2.0 V [4-8].

The synthesis process plays a major role in improving the physicochemical properties of the electrode materials. And it is well known that the use of nanophase Li$_4$Ti$_5$O$_{12}$ as electrode reduces the diffusion length leading to improved lithium intercalation kinetics [9-11]. Moreover, the use of nanomaterials allows for an increased number of active sites for surface reaction decreasing the local current density which, in turn, lowers the potential and leads to high rate capability [12, 13]. Various techniques for the preparation of nanostructured Li$_4$Ti$_5$O$_{12}$ have been reported in the literatures. These include emulsion gel [14], spray drying [15], hydrothermal [16-18], and sol–gel method [19-25]. However, such wet chemical synthesis methods are complicated and always require a non-aqueous medium which increases the cost. Up to now, the solid-state reaction is still the most attractive process for potential industrial application because it has a series of advantages such as simple process, without using solvent and friendly to the environment. In most of previous works, using bulk TiO$_2$ and Li$_2$CO$_3$/LiOH/LiAc as the raw materials, high-temperature calcination and prolonged time were always necessary to improve the phase purity [26-31]. These synthetic conditions (for example, sintering time and temperature), which require long-range diffusion of the reactants, may result in non-homogeneity, irregular morphology, large particle size, broad particle size distribution, and poor control of stoichiometry. In order to avoid the problem above mentioned, the precision control of starting mixture is very important to decrease the temperature and time of calcination and prepare Li$_4$Ti$_5$O$_{12}$ in nanometer regime.

In this study, we obtained the starting mixture precursor by hydrolysis precipitation using the raw materials of tetrabutyl titanate (Ti(OC$_4$H$_9$)$_4$), lithium acetate (LiAc·2H$_2$O). The conditions to prepare spinel Li$_4$Ti$_5$O$_{12}$ with narrow size distribution were optimized, and the electrochemical properties of the prepared Li$_4$Ti$_5$O$_{12}$ were characterized.
2. EXPERIMENTAL

2.1. Sample preparation and characterization

All chemicals were used as received. Tetrabutyl titanate (CP) was procured from Shanghai Meixing Industries Co. Ltd., China and lithium acetate (AR) from Shanghai Hengxin Chemical Reagent Co. Ltd., China. Typically, 0.5 mol of tetrabutyl titanate was dissolved in ethanol to obtain solution A. A stoichiometric lithium acetate with a Li:Ti molar ratio of 4:5 was dissolved in deionized water to obtain solution B. To compensate for the Li$_2$O sublimation during the thermal treatment, a LiAc.2H$_2$O weight excess from 2% to 10% was adopted. Then, the LiAc aqueous solution was dropped slowly into Ti(OC$_4$H$_9$)$_4$ ethanol solution under magnetic stirring. In this dropping process, the solution turn to white as Ti(OC$_4$H$_9$)$_4$ hydrolyzing. After 1 h stirring, the white solution was treated at 80 °C to remove solvents gradually, and became a precursor finally. Then, the dry precursor was ground and calcined at different temperature for 12 h in air at a heating rate of 5 °C/min, followed by natural cooling in the furnace. Finally, a white powder with amount of about 45 g was obtained. All electrode active samples were prepared according to above sequence.

The thermal analysis (TGA-DSC, SDT Q600) to determine suitable temperatures for calcining the precursor were performed with α-Al$_2$O$_3$ as the reference substance at a heating rate of 5 °C min$^{-1}$ in air. The crystal structures of the as-prepared powders were carried on X-ray diffractometry (XRD, RIGAKU DMAX-33) using Cu Ka radiation ($\lambda$ = 1.5418x10$^{-10}$ m). The morphological characteristics of the powders were investigated using scanning electron microscopy (SEM, Hitachi TM3000).

2.2. Electrode preparation and electrochemical characterization

A two electrode coin-type cell (CR 2032) was fabricated for charge–discharge tests. To make the working electrode, 80 wt.% Li$_4$Ti$_5$O$_12$, 10 wt.% super-P carbon, and 10 wt.% polyvinylidene fluoride (PVdF) were mixed well together in N-methyl pyrrolidone (NMP). The prepared slurry was spread uniformly on current collectors of copper foil (10 μm), and it was dried overnight at 100 °C in vacuum. The test cell is a two-electrode system and all of the cells were assembled in a glove box filled with dry argon gas. A lithium foil was used as the counter electrode, and a microporous polypropylene film (Celgard, 2400) was used as the separator. An equal volume of the ethylene carbonate (EC) and diethyl carbonate (DEC) (1:1) solvent with 1M lithium hexa-fluoro-phosphate (LiPF$_6$) salt was used as an electrolyte. The polypropylene membrane was soaked in the electrolyte for 24 h prior to use. The specific capacity and current density were based on the active material (Li$_4$Ti$_5$O$_12$) only, which was typically 6–7 mg per electrode with a surface area of about 2 cm$^2$.

Galvanostatic charge–discharge test was performed using automatic charge–discharge equipment (Land 2001A) in a potential range of 1.0–3.0 V versus Li$^+$/Li electrode at 0.1 C rate (16.7 mA.g$^{-1}$). All electrochemical experiments were conducted at room temperature with a constant humidity.

3. RESULTS AND DISCUSSION

DSC/TGA was performed on the dry precursor powders prepared from hydrolysis precipitation with 8% Li weight excess, which was provided to compensate for the loss of Li during thermal treatment according to the other researchers’ work [32]. The thermal treatment temperature is in the range of 30 to 1000 °C. The result is shown in Fig.1. The figure clearly shows a step-like pattern of weight loss for the precursor. All electrode active samples were prepared according to above sequence.

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Figure 1. TGA and DSC curves of the dry precursor powders prepared from hydrolysis precipitation with 8% Li excess.

Figure 2. XRD patterns of the samples calcined at 800 °C for 12h with (a) no Li excess, (b) 8% Li excess. The main diffraction peaks of Li$_4$Ti$_5$O$_12$ and rutile phase TiO$_2$ are marked in the diagram.
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The precursor, which comes from the hydrolysis of Ti(OC$_5$H$_9$)$_4$, and another peak centered at 360 °C is attributable to the decomposition of LiAc. Above 500 °C, no obvious weight loss is observed in the TGA curve. But the broad endothermic peak observed in the DSC curve indicates the reaction between Li$_2$O and TiO$_2$ continue to occur higher than 500 °C. In order to investigate the effect of Li excess on the crystal structure, the morphology and the electrochemical performance of the Li$_4$Ti$_5$O$_{12}$ anode powders, the following calcination temperature for the dry precursor powders is firstly carried out at 800 °C, assuring the complete reaction.

The XRD patterns and SEM images of the samples calcined at 800 °C for 12h with no Li excess and 8% Li excess are presented in Fig.2 and Fig.3. It can be seen that the primary product for the sample with no Li excess is a mixture of spinel-type Li$_4$Ti$_5$O$_{12}$ phase.
Figure 6. The initial charge-discharge profiles for the as-prepared oxides after the calcination at between 600 °C and 900 °C for 12h at 0.1C rate.

Figure 7. A comparison of discharge and charge curves of pure Li$_4$Ti$_5$O$_{12}$, anatase TiO$_2$ and the as-prepared oxides after the calcination at 600 °C at 0.1 C rate.

with Ti deficit and rutile-type TiO$_2$. But when 8% excess of Li is considered, very minor amount of rutile TiO$_2$ was detected. The mean size of the powders with no Li excess measured from the SEM image is much larger than that of the sample with 8% Li excess due to the fairly existing TiO$_2$ phase. The initial charge-discharge test shown in Fig.4 presents that the electrode material with 8% Li excess have the best performance, comparing with the other samples with 0%, 2%, 5% and 10% Li excess. So 8% Li excess is considered hereafter.

Fig.5 shows the XRD patterns of the as-prepared samples with 8% Li excess calcined at 600 °C and 750 °C for 12h. After the calcination at 600 °C, the product mainly comprises Li$_4$Ti$_5$O$_{12}$ phase, anatase-type TiO$_2$ and rutile-type TiO$_2$ based on the XRD results. With the increase of calcination temperature, the peak intensity of the spinel Li$_4$Ti$_5$O$_{12}$ increases greatly, indicating that much more Li$_4$Ti$_5$O$_{12}$ is formed as well as the improved crystallinity of the material with temperature. Meanwhile, no obvious diffraction peaks of rutile TiO$_2$ and anatase TiO$_2$ are detected after the calcination at 750 °C. But with increasing temperature to 800 °C, the diffraction peaks corresponding to rutile-type TiO$_2$ phase emerge again, as shown in Fig.2. This may be attributed to the much more loss of Li by evaporation at temperature higher than 750 °C, which is corresponding to the weight loss in the TGA curve as shown in Fig.1. Therefore, it suggests that the precursor powder from the hydrolysis precipitation can be transformed into spinel Li$_4$Ti$_5$O$_{12}$ after calcination at 750 °C, which greatly increases the precise stoichiometry of the oxide. The sample calcined at 750 °C has a much smaller particle size and a more homogenous distribution than that of the sample calcined at 800 °C as shown in Fig.3.

To clarify the effect of impurity on the electrochemical performance of the Li$_4$Ti$_5$O$_{12}$, some electrochemical tests were carried out. Fig.6 shows the charge–discharge curves of the as-prepared oxides after the calcination between 600 °C and 900 °C at 0.1 C rate over the potential range of 1.0-3.0 V (versus Li$^+/Li$). It can be seen that the oxides calcined between 700 °C and 850 °C exhibit extremely flat discharge/charge plateaus at the voltage about 1.5–1.6V (versus Li$^+/Li$), which is an intrinsic electrochemical property of spinel Li$_4$Ti$_5$O$_{12}$. Moreover, the small potential difference between charge and discharge plateau is only 60 mV, indicating that the polarization of all the samples is very low. However, the charge plateau of the sample after the calcination at 900 °C departs from the standard potential gradually. The main reason is perhaps due to the high lithium deficiency in Li$_4$Ti$_5$O$_{12}$ from the evident lithium evaporation at high temperature, which causes the high polarization of the electrode. The sample calcined at 750 °C exhibits the best electrochemical performance with a discharge specific capacity of 152 mAh.g$^{-1}$ and 97% coulombic efficiency, while the sample calcined at 900 °C exhibits a specific capacity of only 92.5 mAh.g$^{-1}$ in the first cycle.

Different from the anodes prepared from the calcination higher than 700 °C, which show only one discharge/charge voltage platform at ~1.5 V~1.6 V, the 600 °C calcined one shows two discharge/charge platforms, one around 1.7 V/1.9 V and the other at ~1.5V/1.6V. As observed in Fig.5, the 600 °C calcined sample is characterized by the presence of noticeable amount of anatase-type TiO$_2$ impurity. And some literatures also testified that the discharge/charge platform at ~1.7 V/1.9 V is an intrinsic electrochemical property of anatase TiO$_2$[33-35]. We also synthesized anatase TiO$_2$ through the hydrolysis precipitation without adding the LiAc and tested the pure anatase-type as the anode at the discharge/charge rate of 0.1 C. As shown in Fig.7, broad discharge and charge platforms are observed at the voltages of 1.8 V and 1.9 V, respectively. Such results strongly support that the second discharge/charge platforms for the 600 °C calcined sample are originated from the lithium intercalation and de-intercalation into the impurity phase of anatase-type TiO$_2$ in the anode.

The cycling stability of the 750 °C calcined sample with 8% Li excess, which has the highest initial discharge capacity, is shown in Fig.8. High cycling stability was observed. At 0.1 C rate, the decay in discharge capacity after the 7 cycles was only 2.6%. It suggests the high reversibility and stability of the Li-intercalation and de-
intercalation through the as-prepared Li$_4$Ti$_5$O$_{12}$.

The spinel Li$_4$Ti$_5$O$_{12}$ with smallest possible particle size and highest possible phase purity exhibits the best electrochemistry performance. So the lowest possible temperature is needed to synthesize phase pure and well-crystallized Li$_4$Ti$_5$O$_{12}$ nanoparticles via a solid state route, which is the most attractive process. Different from the conventional solid-state reaction, the hydrolysis precipitation as the precursor is obtained firstly in this study. The precise control and uniform mixing of the reactants are realized, which is very important to the preparation of pure Li$_4$Ti$_5$O$_{12}$. Meanwhile, the newly formed reactant mixture is highly likely to be interconnected with each other and the reactivity of them is elevated through the hydrolysis precipitation reaction, which will decrease the following calcination temperature finally.

4. CONCLUSION

We have synthesized spinel Li$_4$Ti$_5$O$_{12}$ materials as the negative electrode for Li-ion batteries using a hydrolysis precipitation-assisted solid state method. The influence of hydrolysis precipitation performance is positive to the preparation of Li$_4$Ti$_5$O$_{12}$ at the lowest temperature. A pure Li$_4$Ti$_5$O$_{12}$ phase not containing an impurity phase is obtained based on the XRD measurements, when the precipitation precursor mixture with 8% excess lithium is calcined at 750 °C for 12h. The as-obtained Li$_4$Ti$_5$O$_{12}$ powder is in a narrow size distribution. Accordingly, it shows the higher capacity of ~152 mAh.g$^{-1}$ at 0.1 C rate, comparing with the Li$_4$Ti$_5$O$_{12}$ products containing impurity phases, such as rutile TiO$_2$ and/or anatase TiO$_2$.

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REFERENCES