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An investigation of Li_2TiO_3 -coke composite anode material for Li-ion batteries†

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Anode material Li_2TiO_3 -coke was prepared and tested for lithium-ion batteries. The as-prepared material exhibits excellent cycling stability and outstanding rate performance. Charge/discharge capacities of 266 mA h g^{-1} at 0.100 A g^{-1} and 200 mA h g^{-1} at 1.000 A g^{-1} are reached for Li_2TiO_3 -coke. A cycling life-time test shows that Li_2TiO_3 -coke gives a specific capacity of 264 mA h g^{-1} at 0.300 A g^{-1} and a capacity retention of 92% after 1000 cycles of charge/discharge.

Introduction

Except for high energy density, long cycling life-time, and high safety, the application of rechargeable batteries in electric vehicles also requires that the batteries have high power density. In the manufacture of high-power density batteries, high rate capable cathode and anode materials are required.^{1,2} Generally, the cathode materials, such as NCM and LiFePO_4 , are high rate capable. However, an anode material with high rate capability and high energy density is still not available. Presently, graphite is widely employed as an anode material in lithium ion batteries.^{3,4} However, graphite is not high rate capable.⁵ The spinel lithium titanate ($\text{Li}_4\text{Ti}_5\text{O}_{12}$) is a high rate capable anode material, which has an almost zero-strain.⁶ The reduced volume change in lithium ion intercalation/deintercalation guarantees its high reversibility^{7,8} and long cycling life-time. The high lithium insertion potential (about 1.55 V vs. Li^+/Li) could effectively avoid lithium plating over the anode in the charge process even at low temperature or high current density.^{9,10} Hence, regarding safety issues, $\text{Li}_4\text{Ti}_5\text{O}_{12}$ represents a much better option than graphite. However, except the advantages of the spinel $\text{Li}_4\text{Ti}_5\text{O}_{12}$, the high lithium ion insertion potential and the relatively low theoretical capacity (175 mA h g^{-1}) of $\text{Li}_4\text{Ti}_5\text{O}_{12}$ lead to a lower cell energy density, which limits the applications of $\text{Li}_4\text{Ti}_5\text{O}_{12}$ in areas that need the batteries having higher energy density. Hence, searching for high performance anode material that has high specific capacity, high rate capability, long cycling life-time, and high safety is still the goal of researchers in battery investigation.

Monoclinic Li_2TiO_3 has a layer structure and a three-dimensional lithium ion diffusion network.¹¹ In recent years, Li_2TiO_3 is often added into cathode materials, such as

$\text{LiNi}_{1-x-y}\text{Co}_x\text{Mn}_y\text{O}_2$ ($x > 0$, $y > 0$, $x + y < 1$) and LiCoO_2 , to improve their rate capability and stability.^{12,13} Although, Li_2TiO_3 is a good lithium ion conductor, it is almost an electronic insulator. In order to making use of Li_2TiO_3 as an anode material, one must improve the electronic conductivity of Li_2TiO_3 . Investigators usually add carbon materials¹⁴⁻¹⁸ into metal oxide electrode materials or directly prepare the metal oxide electrode materials in metal foam^{19,20} to improve their electronic conductivity. Since the electronic conductivity of Li_2TiO_3 is very poor, we choose to support relatively less amount of Li_2TiO_3 over relatively larger amount of coke to reach good electronic conductivity. Because petroleum coke has medium electronic conductivity and pores that lithium ions could easily diffuse in and out. The coke itself is a potential high rate anode material. In the present work, by making use of the good lithium ion conductivity of Li_2TiO_3 and the electronic conductivity of coke, we intend to use lithium hydroxide, TiO_2 , and petroleum coke to prepare a practically useful high rate anode material Li_2TiO_3 -coke (LTOC) for lithium ion batteries.

Experimental

The synthetic procedure of Li_2TiO_3

Li_2TiO_3 was prepared by the following method. TiO_2 (5.6724 g), $\text{LiOH} \cdot \text{H}_2\text{O}$ (5.9602 g), and H_2O (50.0 ml) were mixed and milled at 180 rpm for 6 h to obtain a slurry. The slurry was transferred into a beaker. The milling container and beads were washed three times with DI water (10 ml for each time) and the washing out mixture was combined with the previous slurry to obtain a diluted slurry. Adding DI water to the diluted slurry to a total volume of 100 ml for spray dry. After spray dry, a powder was obtained, and then the powder was calcined at 850 °C for 4 h in air to obtain Li_2TiO_3 .

The pretreatment of oil coke

The oil coke was placed into a tube furnace and heated from room temperature to 1200 °C at a heating rate of 5 °C min^{-1} in

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argon, then heated at 1200 °C for 6 h, and then cooled down to room temperature in argon to obtain the preheated coke, which was used as anode material and precursor for Li_2TiO_3 -coke (LTOC) preparation.

The synthetic procedure of LTOC

The LTOC was prepared through the following method. The preheated coke (6.5000 g), TiO_2 (1.7067 g), $\text{LiOH}\cdot\text{H}_2\text{O}$ (1.7933 g), surfactant ($[[n\text{-C}_{16}\text{H}_{31}(\text{CH}_2)_3\text{N}]\text{CO}_3\text{CH}_3]$ writing as QA, 0.9457 g), and distilled water (50.0 ml) were added into a ball mill container and milled at 180 rpm for 6 h to obtain a slurry. The slurry was transferred into a beaker. The mill container and beads were washed three times with DI water (10 ml for each time) and the washing out mixture was combined with the previous slurry to obtain a diluted slurry. Adding DI water to the diluted slurry to a total volume of 100 ml for spray dry. After spray dry, a powder was obtained, and then the powder was calcined at 850 °C for 4 h in argon to obtain LTOC.

Cell preparation

The as-prepared anode material, conductive carbon black, carbon nanotubes (dispersed in *N*-methyl-2-pyrrolidone), polyvinylidene fluoride (PVDF) binder (dissolved in *N*-methyl-2-pyrrolidone) were mixed according to the mass ratio of 85 : 10 : 1.5 : 3.5 to prepare a paste. The paste was coated on a copper foil (the thickness of the anode coating layer is 30 μm) with an applicator (frame-type), and then dried in a vacuum oven at 120 °C for 2 h to obtain the anode plate. The dried anode plate was cut into cycles with a diameter of 14 mm. The anode cycles were farther dried in a tubular furnace (filled with argon) at 140 °C for 5 hours, and then transferred into an argon filled glove box (Mikrouna) in argon atmospheres. The anode cycles together with the separator (PP), the lithium metal cycles (as the counter electrode), and the electrolyte solution (1.0 M of LiPF_6 in EC, PC, DMC, and EMC, the volume ratio of EC, PC, DMC, to EMC is 1 : 1 : 1 : 1) were used to assemble coin-type cells.

The crystal structure of the samples was analyzed by X-ray diffraction spectroscopy (Rigaku D/Max-rA, CuK radiation). The surface morphology and microstructure of the samples were characterized by SEM (FE-SEM S-4800, Hitachi) and TEM (Titan G2 60-300, FEI). Laser Raman measurements were conducted on a laser Raman spectroscopy meter with an excitation wavelength of 532 nm (*via* RM10000, Renishaw). The X-ray Photoelectron Spectroscopy (XPS) measurements of samples were conducted on a XPS spectroscopy meter (ESCALAB 250XI, Thermo Fisher). The specific surface areas of the materials were measured by BET method (Nova 2200e, Quantachrome).

The discharge/charge tests of the cells were carried out on a NEWARE CT-3008-5 V-10 mA system. The voltage window is 0.01–2.0 V. In the rate performance test, the cells were tested at current densities 0.033, 0.100, 0.200, 0.300, 0.400, 0.500, 0.600, 0.700, 0.800, 0.900, 1.000, 2.000, 3.000, 4.000, 5.000, and 0.100 A g^{-1} , respectively. In the cycling life-time test, the cells were charge/discharged at 0.300 A g^{-1} . The cycling voltammetry curves were recorded on an electrochemical workstation (CHI 660E, CHENHUA) at a scan rate of 0.1 mV s^{-1} within 0.01–2.0 V.

The EIS measurement of cells was also carried out on the electrochemical workstation (CHI 660E, CHENHUA). All tests were carried out at room temperature.

Results and discussion

The Laser Raman characterization shows that the preheated oil coke has 2 bands (the D band at 1360 cm^{-1} and the G band at 1580 cm^{-1}). The ratio of the D band intensity (I_D) to the G band intensity (I_G) is 0.91 ($I_D/I_G = 0.91$), which indicates that the preheated oil coke is not graphitized (ESI, Section 1 Fig. s1†).^{21–23} The TEM images also show that the preheated coke is not graphitized (ESI, Section 1 Fig. s2 and s3†).

The specific surface areas of Li_2TiO_3 , coke, and LTOC are 2.9, 13.1, and 10.1 $\text{m}^2 \text{g}^{-1}$, respectively. These numbers are close to the specific surface area of regular $\text{Li}_4\text{Ti}_5\text{O}_{12}$.²⁴

In LTOC, the percentage of Li_2TiO_3 is 35.6 wt% (ESI, Section 2†). The EDS analysis indicates that the LTOC contains Al, Si, and S impurities, but not contains nitrogen (ESI, Section 3†).

The XRD analysis shows that the coke has only a broad peak between 20° and 30° (Fig. 1). The broad peak is assigned to the amorphous coke. The peaks of pristine Li_2TiO_3 match that of the standard monoclinic Li_2TiO_3 (PDF#33-0831) with lattice parameters $a = 5.069 \text{ \AA}$, $b = 8.799 \text{ \AA}$, and $c = 9.759 \text{ \AA}$. Hence, the pristine Li_2TiO_3 is monoclinic Li_2TiO_3 . In LTOC, except from the broad peak between 20° and 30° (assigned to amorphous coke), there are peaks that exactly match that of monoclinic Li_2TiO_3 . Hence, monoclinic Li_2TiO_3 is formed in LTOC.

The SEM images of Li_2TiO_3 , coke, and LTOC are given in Fig. 2. The image of Li_2TiO_3 shows that the average size of Li_2TiO_3 crystals is about 1 μm . Petroleum coke has a particle size distribution from nanometers to tens of micrometers. In LTOC (Fig. 2c and d), the crystal size of Li_2TiO_3 is in nanometer level. The SEM images show that under the help of surfactant QA, the Li_2TiO_3 crystals are uniformly distributed on the coke particles. The results indicate that the presence of coke inhibits the growth of Li_2TiO_3 crystals. The attachment of Li_2TiO_3 nanocrystals on coke carbon could offer an excellent interface for electron exchange between Li_2TiO_3 and coke carbon. The EDS mapping of elements C, O, and Ti in LTOC (ESI Section 4 Fig. S4†) shows that the oxygen and titanium are uniformly distributed on coke carbon.

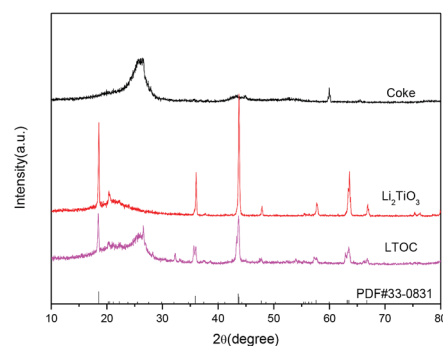


Fig. 1 The XRD patterns of coke carbon, Li_2TiO_3 , and LTOC.



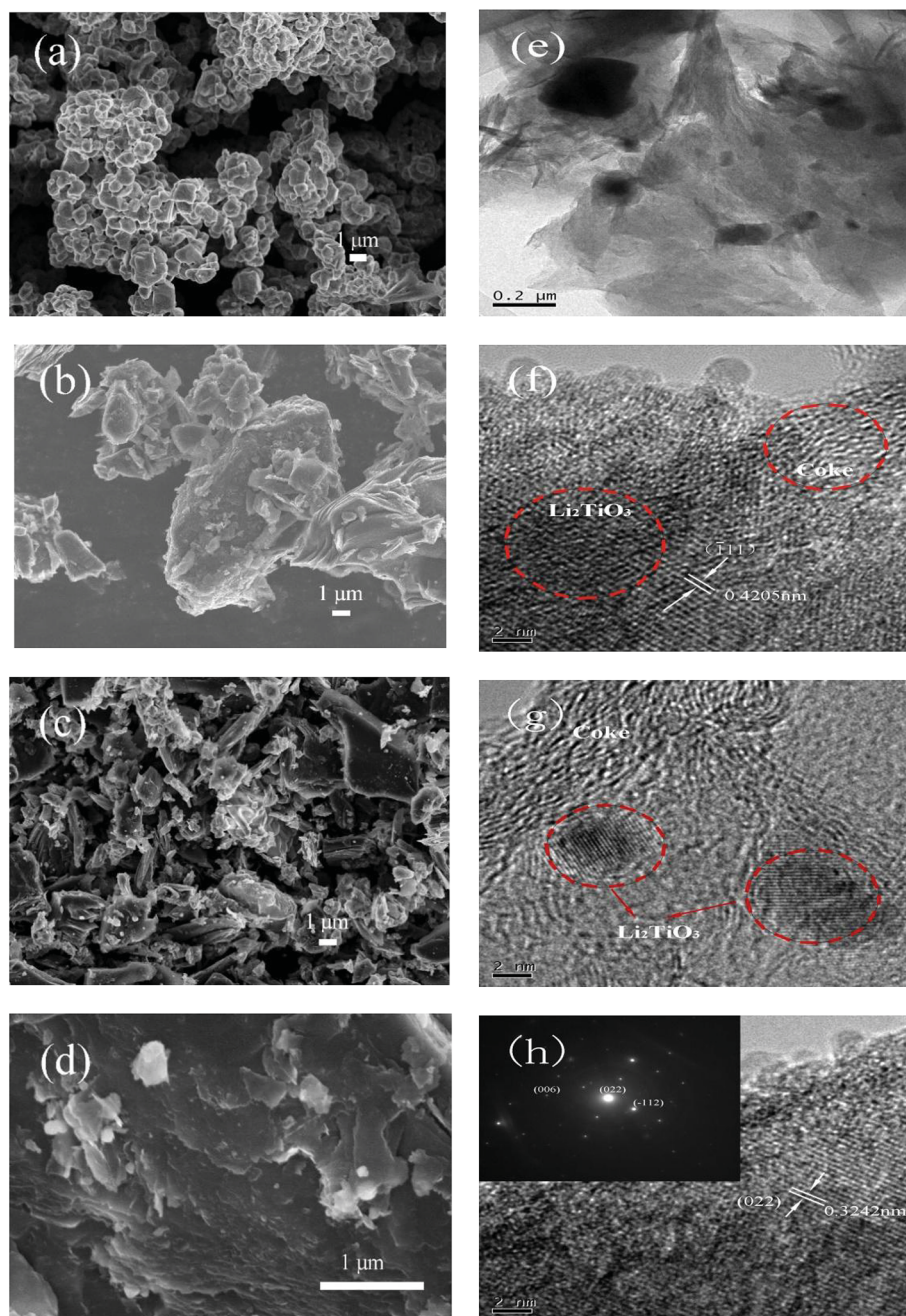


Fig. 2 SEM images of Li_2TiO_3 (a), petroleum coke (b), and LTOC with different resolutions (c) and (d). TEM of LTOC (e), (f), (g), and (h).

The high resolution TEM images (Fig. 2e–h) reveal that the Li_2TiO_3 crystals are formed on coke and the Li_2TiO_3 phase is tightly contacted with the carbon phase. The typical Li_2TiO_3 lattice spacings of 0.3242 and 0.4205 nm can be indexed to (022) and (111) planes, respectively (the inset of image (h) gives the selected area electron diffraction pattern of a Li_2TiO_3 crystal). The results are consistent with that observed in the SEM characterization.

The XPS measurement shows that there is about 12.7% of Ti^{4+} ions reduced to Ti^{3+} ions at high calcination temperature (850 °C). However, the major part of titanium ions is still Ti^{4+} ions (Fig. 3).

The cyclic voltammetry measurements are performed on fresh cells with metal lithium anodes. Fig. S5† (ESI Section 5†) shows the cyclic voltammograms (CVs) of Li_2TiO_3 , coke, and LTOC in their first three cycles. In the reduction process, the CV curves of the three electrodes in their first cycle are significantly different from their subsequent curves, which could be caused from the SEI formation and irreversible structure change. Li_2TiO_3 has weak reduction/oxidation peaks at 1.52/1.63 V (Fig. S5a†), which are assigned to Li_2TiO_3 .²⁵ The major reduction/oxidation of Li_2TiO_3 appears between 0.01 and 1.00 V (vs. metal Li). The CV curves of coke (Fig. S5b†) have strong



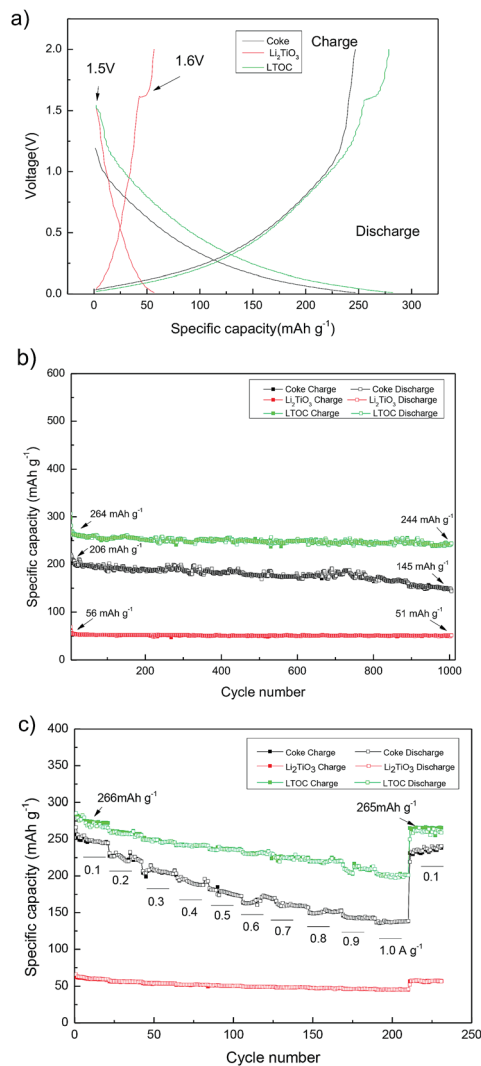


Fig. 5 (a) The discharge/charge curves of the cells with coke, Li₂TiO₃, and LTOC as cathode and metal Li as anode at 0.100 A g⁻¹; (b) discharge/charge cycling life-time test at 0.300 A g⁻¹; (c) discharge/charge rate cycling test at different current densities.

5.000 A g⁻¹, respectively (ESI Section 8†). The specific capacity of LTOC is much higher than that of spinel Li₄Ti₅O₁₂ (about 160 mA h g⁻¹).³⁰

The reversible specific capacities of LTOC and coke are 266 and 245 mA h g⁻¹ (at 0.100 A g⁻¹), respectively (Fig. 5c). The percentage of Li₂TiO₃ in LTOC is 35.6 wt% (ESI, Section 2†). From these data, the specific capacity of Li₂TiO₃ in LTOC is estimated to be 304 mA h g⁻¹, which corresponding to insert/extract 1.25 Li⁺ ions per Li₂TiO₃. The results show that Li₂TiO₃ could host more Li⁺ ions than Li₄Ti₅O₁₂.²⁹

It is known that Li₂TiO₃ is not active as an anode material for lithium ion batteries. The reason is that Li₂TiO₃ is an electronic insulator. After supporting Li₂TiO₃ nanoparticles over coke, better electron transfer pathways between coke and Li₂TiO₃ nanoparticles are established. Since the Li₂TiO₃ particles are in nanometer level, the Li-transport distances are shortened. In the redox reaction of Li₂TiO₃, the insertion of positively charged

Li⁺ could be more easily balanced with the uptake of electrons to compensate Ti³⁺ cations.

Conclusions

In conclusion, the investigation show that the presence of coke carbon could inhibit the crystal growth of Li₂TiO₃ in LTOC composite and the Li₂TiO₃ nanocrystals are attached on the surface of coke. The tight attachment of Li₂TiO₃ nanocrystals on coke offers an excellent interface for electron exchange between Li₂TiO₃ and coke carbon, which might be able to fully activate the Li₂TiO₃ nanocrystals as an anode material, which could contribute large specific capacity (304 mA h g⁻¹). On the other hand, the introduction of Li₂TiO₃ into coke raises the D_{Li^+} of coke, and therefore, improves the rate capability of the material. Hence, the high specific capacity, the low discharge/charge voltage of LTOC vs. Li⁺/Li, the high rate capability, and the simple preparation method of LTOC make it a practical anode material to make high rate and high energy density batteries.

Conflicts of interest

There are no conflicts to declare.

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