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Effect of hydrogenation on performance of TiO₂(B) nanowire for lithium ion capacitors

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24

25 **Abstract**

26 TiO₂(B) nanowires have shown excellent capacitance and energy density with a very low
27 charge transfer resistance in 80-μm thick lithium ion capacitor electrodes. Nanowires
28 hydrogenated by heat treatment at 500 °C showed improved Li ions diffusion and an increase
29 in capacitance from 148 to 194 F/g as well as energy density from 23 to 30 Wh/kg.
30 Hydrogenation of oxides as a way to improve their capacitance is critically discussed.

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32 *KEYWORDS.* TiO₂(B), nanowire, lithium ion capacitor, hydrogenation, pseudocapacitor

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45 1. Introduction

46 Energy storage demand is increasing from portable electronics to the power grid.
47 Supercapacitors are attractive as energy storage devices due to their high power density and
48 good cyclability; however, the low energy density is the limiting factor. Energy density, E ,
49 can be increased mainly by two ways: 1) by increasing the capacitance, C , via design of high
50 capacity electrodes beyond carbon and 2) by expanding the voltage window, V , of the devices
51 ($E = 1/2 CV^2$) [1]. Pseudocapacitive materials, such as metal oxides [2] or organic redox-
52 active molecules [3], are commonly used to increase the charge storage capacity of the
53 capacitive electrodes via additional redox processes at the electrode-electrolyte interface. One
54 way to design high energy and power density devices is to use pseudocapacitive materials
55 (high capacitance) in organic electrolytes (large voltage window). The concept of lithium-ion
56 capacitor (LIC) provides high energy and power density by using a pseudocapacitive material
57 as an anode and activated carbon as a cathode in an organic electrolyte [4]. LICs can
58 accommodate both faradaic (lithium insertion/extraction) and non-faradaic
59 (adsorption/desorption) processes at the same time with an expanded voltage window in
60 lithium-based organic electrolytes [5].

61 Among various TiO_2 phases, $\text{TiO}_2(\text{B})$ bronze phase has gained attention as a lithium
62 insertion material due to its high theoretical capacity of 335 mAh/g and good rate capability
63 [6]. Compared to other TiO_2 phases involving bulk diffusion-controlled mechanism, $\text{TiO}_2(\text{B})$
64 has an open structure allowing faster lithium ion uptake. However, it still suffers from
65 sluggish ionic transport and poor electronic conductivity. To address this problem, conductive
66 additives like amorphous carbon [7], graphene [8], carbon nanotubes (CNT) [9] and
67 conducting polymers [8,10] were added to different polymorphs of TiO_2 . Since addition of

large amounts of conductive additives contributes little to capacitance of the device, this is not a favorable option. Furthermore, some carbon materials like CNT and graphene are fairly expensive to produce.

To overcome this issue, oxygen defects in the metal oxide lattice structure can be introduced, as they have been reported to improve the conductivity of the material by reducing the bandgap [11]. For instance, yellowish and black TiO_2 nanoparticles after hydrogen treatment showed a significant increase in capacity in lithium-ion batteries [12,13]. However, the origin of this improved electrochemical activity has not been fully understood and even the increase of capacitance has not been supported with sufficient evidence.

Metal oxides (e.g., MnO_2 , RuO_2 , Nb_2O_5 , etc.) are well-known pseudocapacitive materials, which have been studied in various configurations [2,14]. However, only a few reports are available on the $\text{TiO}_2(\text{B})$ phase as a lithium insertion electrode [15–17], and particularly, the effect of hydrogenation of $\text{TiO}_2(\text{B})$ on electrochemical performance for lithium-ion capacitor is not well studied. Recently, partially reduced oxides have shown improvement in photocatalytic activity and electrochemical properties in both lithium-ion batteries and supercapacitors [11,12]. In all cases, the improved electrochemical performance of the oxides was attributed to the defect-induced conductivity due to decrease in the bandgap. Hydrogen plasma treatment [18], sodium borohydride (NaBH_4) reduction [19], vacuum annealing [20] and heat treatment under H_2 at atmospheric pressure [21] have been used to make oxygen-deficient oxides.

Herein, one-dimensional $\text{TiO}_2(\text{B})$ nanowires were synthesized via hydrothermal method followed by heat treatment under hydrogen gas for the introduction of Ti^{3+} to create oxygen deficiency. The resulting TiO_2 NW and $\text{H}_2\text{-TiO}_2$ NW were tested in organic electrolyte as an anode for LICs. Electrochemical performance is compared with other

hydrogenated oxides and the challenges to achieve higher capacity via the hydrogenation method are elaborated.

2. Experimental

TiO₂ NW/H₂-TiO₂ NW synthesis. 3.7 g of anatase TiO₂ (15 nm, Alfa Aesar) nanoparticles were stirred with 20 ml of 15 M NaOH (Sigma) for 1 h. The suspension was put into a Teflon-lined stainless steel autoclave and heated at 180 °C for 72 h, followed by washing with 0.1 M HCl until the supernatant became neutral. Dried powder was heated to 450 °C for 5 h in air to produce TiO₂(B) phase nanowires (TiO₂ NW). Heating up to 4 h at 500 °C under hydrogen atmosphere (Ar - 10% H₂), following the process conditions reported in the literature [12,21–23] led to hydrogenated TiO₂ NW (H₂-TiO₂ NW).

Electrode preparation. The slurry was made by mixing 80 wt% of TiO₂ NW or H₂-TiO₂ NW with 10 wt% polyvinylidene fluoride (Alfa Aesar, Ward Hill, USA) binder and 10 wt% carbon black (Alfa Aesar, Ward Hill, USA) in 1-methyl-2-pyrrolidinone (Alfa Aesar, Ward Hill, USA). A 200 µm thick coating was deposited on copper foil using a doctor blade and dried overnight resulting in ~80 µm thick electrodes. The coated dried foil was punched into 5 mm diameter disks, which were used for the electrochemical testing.

Electrochemical setup. The electrodes were tested in a three-electrode configuration using a Swagelok[®] cell with TiO₂(B) NW or H₂-TiO₂(B) NW as the working electrode, activated carbon as a counter electrode and silver (Ag) wire as a reference electrode. Two layers of Celgard separator were used and tests were conducted in 1 M solution of lithium perchlorate (Sigma) in 1:1 volume ratio of ethylene carbonate (Sigma) and diethyl carbonate (Sigma) electrolyte. Cyclic voltammetry (CV) was performed on a potentiostat (VMP3,

Biologic, USA) at different scan rates from 1 to 20 mV/s. Electrochemical impedance spectroscopy (EIS) was run in the frequency range of 200 kHz to 10 mHz at a sine wave signal amplitude of +/-10 mV centered at 0 V vs. open circuit potential.

The specific capacitance (C_{sp}) was calculated according to equation 1:

$$C_{sp} = \frac{1}{\Delta E} \frac{\int i dV}{\nu m} \quad (\text{Eq. 1})$$

where, ΔE is the voltage window, i is the discharge current, V is the voltage, ν is the sweep rate, and m is the mass of active material in one electrode. The energy density and power density were obtained from equation 2:

$$E = \frac{1}{2} CV^2 \quad P = \frac{E}{t} \quad (\text{Eq. 2})$$

where, E , C , V , P and t represent energy density, capacitance, voltage window and discharge time, respectively.

Characterization. The morphology of the samples was characterized using a transmission electron microscope (TEM) (JEOL JEM-2100, Japan) with an accelerating voltage of 200 kV. The TEM samples were prepared by placing several drops of the sample dispersion in ethanol onto a copper grid and drying in air. The x-ray diffraction (XRD) analysis was performed on a powder diffractometer (Rigaku Smart Lab, USA) with Cu K α radiation. To investigate the changes in surface area, gas sorption analysis was carried out using quadrosorb gas sorption instrument (Qunatachrome, USA) using N₂ gas adsorbate at -196 °C.

3. Results and discussions

Figure 1 presents a schematic of the synthesis process. Hydrothermal synthesis was considered because it is simple, relatively inexpensive and produces gram-scale amounts of the material [24,25]. The absence of the anatase peaks from the x-ray diffraction (XRD) pattern (Fig. 2a) confirmed the conversion of the TiO₂ anatase phase into TiO₂ bronze and the XRD pattern was found similar to previous reports [26]. As-synthesized NW were 20-40 nm in diameter and 1-5 μm in length (Fig. 2b-d) – thinner and longer compared to the previous reports (100 nm thickness and 1 μm length [15]), most likely due to the use of a nanometer scale initial precursor. Therefore, such ultra-fine NW offer large number of active sites for lithium insertion and short diffusion pathways for the improved electrochemical performance. Moreover, the open-structure NW morphology of TiO₂ (B) may be favorable for Li⁺ intercalation. The surface area of H₂-TiO₂(B) NW was found to be 9.5 m²/g. Therefore, it is believed that surface area doesn't have large impact on its electrochemical performance.

[Insert Figure 1]

Figure 1. Schematic presentation of the hydrothermal synthesis and hydrogenation of TiO₂(B) NW.

[Insert Figure 2]

Figure 2. (a) X-ray diffraction pattern of TiO₂(B) NW, (b-d) Transmission electron microscopy (TEM) images of hydrothermally synthesized TiO₂(B) NW.

We have performed XPS analysis on our pristine and hydrogenated samples (Fig. 3). For example, the oxygen content of pristine and hydrogenated TiO₂(B) NW is 67.0 and 66.6 at. %, as can be seen in Fig. 3. Also, the relative concentrations of Ti³⁺/Ti²⁺ of TiO₂(B) NW and hydrogenated TiO₂(B) NW were 3.1 and 2.7. The very slight reduction of the tested

samples was observed.

[Insert Figure 3]

Figure 3. (a) XPS spectra of TiO₂(B) NW and (b) H₂-TiO₂(B) NW.

Cyclic voltammograms (CVs) at various scan rates were acquired in the voltage window of 2.1 V (Fig. 4). Two broad redox peaks appeared at about -1.5 V and -2.0 V, which correspond to the following lithium insertion/extraction processes:

After hydrogenation, these peaks shifted towards higher potentials (-1.3 V and -1.7 V) and separation between the peaks decreased, which may indicate a change in the electronic structure of the TiO₂(B) NW. The CV shape of H₂-TiO₂ NW appeared wider (more capacitive) than pristine TiO₂ NW (Fig. 4a-b), most likely due to the improved conductivity and change in the oxidation state of the Ti, as shown by in-situ x-ray absorption spectroscopy study [27]. However, in order to better understand the hydrogenation induced changes in the oxidation state of Ti and to correlate such changes with the electrochemical performance, further in-situ studies are required, as conducted on other pseudocapacitive materials [27–29]. The capacitance of H₂-TiO₂ NW was found to be 194 F/g at the lowest scan rate of 1 mV/s – 30% higher than pristine TiO₂ NW at the same scan rate (148 F/g, Fig. 4c). It is worth noting that our as-synthesized TiO₂(B) NW electrodes demonstrated a higher capacitance than in various previous reports [15,17,30]. For instance, TiO₂(B) NW@activated carbon yielded 52 F/g at 0.1 A/g [15] and TiO₂-reduced graphene oxide@activated carbon showed 89 F/g at 2 A/g [30]. Also, our 80 μm titania electrodes (similar to commercial LIC electrodes in

thickness) showed a much higher capacitance even compared to a much thinner nanometer-scale TiO_2 film (150 nm) [4]. Likely, thinner NW, shorter diffusion paths and the increased surface area lead to the higher capacitance.

Although the as-synthesized $\text{TiO}_2(\text{B})$ NW outperformed previous reports, we did not observe a dramatic improvement in the capacitance after hydrogenation of the $\text{TiO}_2(\text{B})$ NW, opposed to some other hydrogenated oxides [21,31–33]. It is reported that hydrogenation induced oxygen defects change the electronic band structure of the oxides, which tune the optical and charge transfer properties of the oxide-based electrodes. This, in turn, significantly improves electrochemical activity of the material. It is most likely that the change in the bandgap will enhance the photocatalytic properties, which are strongly dependent on bandgap variation of oxide semiconductors after hydrogenation [11,18]. However, the relationship between oxygen defects induced by hydrogenation and their effect on the conductivity and capacitance needs deeper understanding.

Nyquist plots measured by the impedance spectroscopy (Figure 4d) of both electrodes showed a negligible semicircle and very small charge transfer resistance (R_{ct}). The ohmic resistance (R_{Ω}), of both electrodes was comparable (0.93 Ω). Although, both electrodes appeared capacitive in the low frequency regime, the Nyquist plot of $\text{H}_2\text{-TiO}_2$ NW appeared more capacitive than pristine electrodes. This is most likely due to the improved charge percolation and low electronic resistance in the hydrogenated electrode [34].

Rate performance of both electrodes shows that $\text{H}_2\text{-TiO}_2$ NW exhibits a higher capacitance across the entire range of sweep rates (1-20 mV/s). High cycling stability is critical for the lithium ion battery electrodes experiencing intercalation/deintercalation. The cycling stability was tested at 2 A/g for 1000 cycles. The $\text{H}_2\text{-TiO}_2$ NW electrode showed a stable cycling behavior with 80 % capacitance retention after 1000 cycles (Fig. 4e).

[Insert Figure 4]

Figure 4. (a) Cyclic voltammograms of $\text{TiO}_2(\text{B})$ NW, (b) CVs of $\text{H}_2\text{-TiO}_2(\text{B})$ NW, (c) corresponding rate performance (d) Nyquist plots of $\text{TiO}_2(\text{B})$ NW and $\text{H}_2\text{-TiO}_2(\text{B})$ NW, (e) Cyclic stability up to 1000 cycles of $\text{H}_2\text{-TiO}_2(\text{B})$ NW.

To conclude, we have synthesized $\text{TiO}_2(\text{B})$ NW with diameters between 20-40 nm and 1-5 μm in length. As-synthesized $\text{TiO}_2(\text{B})$ NW exhibited excellent charge storage capacity of 148 F/g in organic electrolyte, which is attributed to the fine NW morphology, good conductivity and short diffusional pathways. The hydrogenation led to ~30 % increase in capacitance to 194 F/g. The corresponding energy density was 30 Wh/kg – higher than pristine $\text{TiO}_2(\text{B})$ NW. Although hydrogenation improved the charge storage capacity, we did not observe a dramatic increase in capacitance after the hydrogenation process, which was predicted in some studies [21,31–33]. We emphasize that more theoretical and experimental work directed toward understanding the relationship between concentration of the oxygen vacancies, conductivity and hydrogenation conditions of titania is needed to further improve energy storage.

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Figure 1.

Figure 2.

Region	BE [eV]	Fraction	Assigned to
Ti 2p _{3/2} (2p _{1/2})	454.2 (459.8)	0.15	Ti ⁺²
	456.5 (461.4)	0.47	Ti ⁺³
	458.2 (463.8)	0.38	TiO ₂

Region	BE [eV]	Fraction	Assigned to
Ti 2p _{3/2} (2p _{1/2})	454.2 (459.9)	0.18	Ti ⁺²
	456.6 (461.5)	0.50	Ti ⁺³
	458.2 (463.8)	0.32	TiO ₂

Figure 4.

Figure 4.