

Electronic Supplementary Information

T-Nb₂O₅ nanoparticles confined in carbon nanotubes with fast ion diffusion rate as lithium storage

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Material characterization

Compositions of **samples** were characterized by X-ray diffraction (XRD; Bruker D8/Cu K α radiation). Morphology analysis was conducted by utilizing Hitachi's field-emission scanning electron microscopy (SEM, S-4800) and transmission electron microscopy and high resolution transmission electron microscope (HRTEM, JEOL JEM-2010F). Thermogravimetric analysis (TGA) was conducted by the Netzsch STA 449C from room temperature to 800 °C with a heating rate of 10 °C min⁻¹. Brunauer-Emmett-Teller (BET) specific surface area and pore structure analysis (DFT method) **were** measured by nitrogen adsorption/desorption with an Autosorb-iQ Pressure Sorption Analyzer (Quantachrome Instruments USA) at 77 K. The preparation of the working electrodes follows the previous procedure.^{1,2} Cyclic voltammetry curves were accounted on a CHI-660C workstation with a sweep rate of 0.1 mV s⁻¹. Galvanostatic charge and discharge processes were conducted on a Land (CT2001A China) at 0.01-3.00 V (versus Li⁺/Li). The mass loading of the T-Nb₂O₅/CNT active material is 0.968 and 0.992 mg for LIBs and SIBs, respectively.

Explanation on equation:

$$i = av^b \quad (1)$$

a and b are changeable factors. The b value of 0.5 shows that the Li^+ storage is completely regulated by the diffusion controlled process, whereas a b value of 1 means that the surface-controlled process completely occupies a dominant position.

Determination of diffusion coefficient

$$i_p = (2.69 \times 10^5)n^{3/2}AD^{1/2}Cv^{1/2} \quad (2)$$

i_p : peak current (A)

n : the transferred number of electrons in the electrochemical reaction

A : the surface area of the electrode (1.13 cm^2)

D : the Li^+ diffusion coefficient of the active material

C : the concentration of lithium ions in the anode

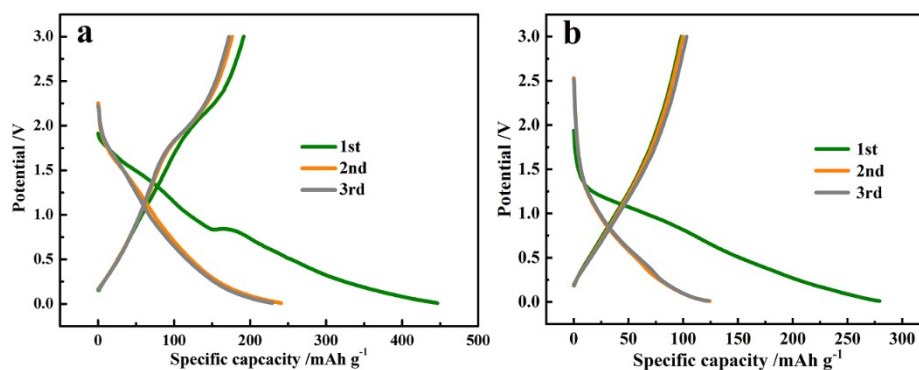


Figure S1. Galvanostatic charge-discharge curves for the first three cycles of T-Nb₂O₅/CNT for LIBs at 0.4 A g⁻¹ (a) and SIBs at 0.1 A g⁻¹ (b).

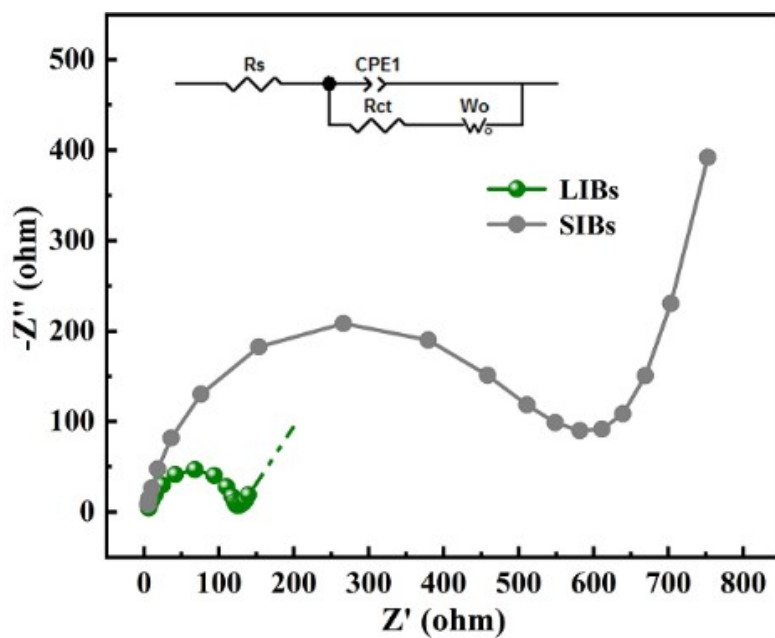


Figure S2. Electrochemical impedance spectra of T-Nb₂O₅/CNT for LIBs and SIBs.

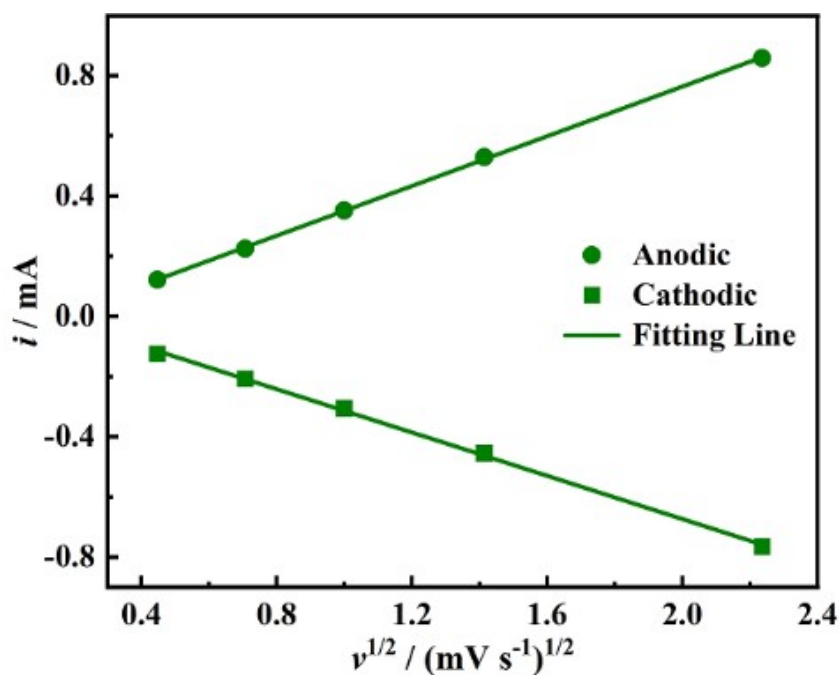


Figure S3. Relationship between peak current (i) and the square root of the scan rate ($v^{1/2}$) of the T-Nb₂O₅/CNT electrode.

Table S1. Comparison of Li⁺ diffusion coefficient for different niobium-based anodes.

Materials	$D_{\text{Li}^+}(\text{cm}^2/\text{s}, \text{lithiation})$	$D_{\text{Li}^+}(\text{cm}^2/\text{s}, \text{delithiation})$	Ref
TiCr _{0.5} Nb _{10.5} O ₂₉ /CNT	3.34×10^{-14}	3.41×10^{-14}	3
s			
TiNb ₂ O ₇ /CNTs	9.27×10^{-16}	1.37×10^{-15}	4
TiNb ₆ O ₁₇	4.28×10^{-14}	5.48×10^{-14}	5
CrNb ₁₁ O ₂₉	1.51×10^{-13}	3.57×10^{-13}	6
Ti ₂ Nb ₁₄ O ₃₉	9.07×10^{-14}	5.52×10^{-14}	7
M-MoNb ₁₂ O ₃₃	4.0×10^{-14}	8.6×10^{-14}	8
N-GeNb ₁₈ O ₄₇	1.596×10^{-15}	1.552×10^{-15}	9
TiNb ₂ O ₇ @C	2.39×10^{-12}	2.13×10^{-12}	10
Mg ₂ Nb ₃₄ O ₈₇	3.23×10^{-13}	6.36×10^{-13}	11
Cu ₂ Nb ₃₄ O ₈₇	3.1×10^{-13}	3.9×10^{-13}	12
T-Nb ₂ O ₅ /CNT	6.61×10^{-12}	4.98×10^{-12}	This

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