

Supporting Information

Uniform Nickel Vanadate ($\text{Ni}_3\text{V}_2\text{O}_8$) Nanowire Arrays Organized by Ultrathin Nanosheets with Enhanced Lithium Storage Properties

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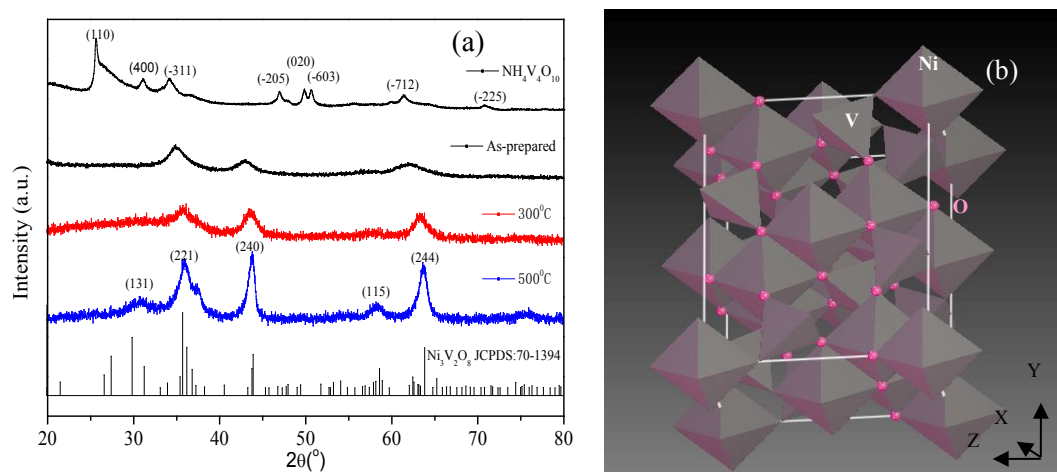


Figure S1. (a) XRD patterns of $\text{NH}_4\text{V}_4\text{O}_{10}$ nanowires and $\text{Ni}_3\text{V}_2\text{O}_8$ nanowires (annealing at different temperatures); and (b) structural model of orthorhombic $\text{Ni}_3\text{V}_2\text{O}_8$.

The crystallite sizes (D) were evaluated by the classical Scherrer equation (Equation S1),¹ the orthorhombic crystallite size (D) of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays annealed at various temperatures was calculated from the X-ray diffraction peaks related to (221), (240) and (244) planes using the Scherrer equation (Equation (2)).^[1]

$$D = k\lambda / \beta \cos \theta \quad (1)$$

$$D = k\lambda / (\beta \div 180 \times 3.14 \cos \theta) \quad (2)$$

Where k is a shape constant, equal to 0.9, λ is the wavelength of the X-rays source used ($\text{Cu K}\alpha = 0.15406 \text{ nm}$), and β and θ are Full width at half maximum (FWHM) and peak position (radians), respectively, determined after Voigt fitting. The results are presented in Table 1. It is evidenced the effect of annealing temperature for crystallite size of $\text{Ni}_3\text{V}_2\text{O}_8$. Overall, the crystallite size increases with the annealing temperature increases.

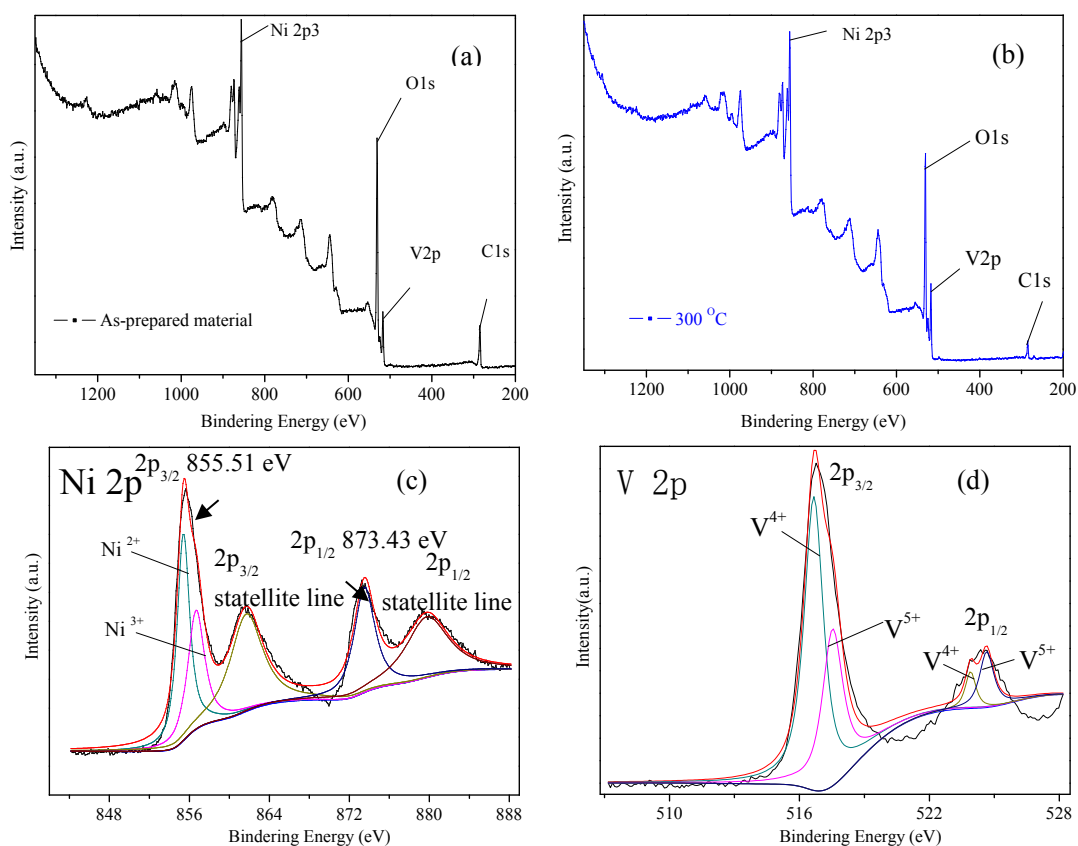


Figure S2. XPS spectra of survey spectrum from as-prepared $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays (a) and that after annealing at 300 °C (b); and the peak deconvolution and fittings of the Ni 2p (c) and V 2p (d).

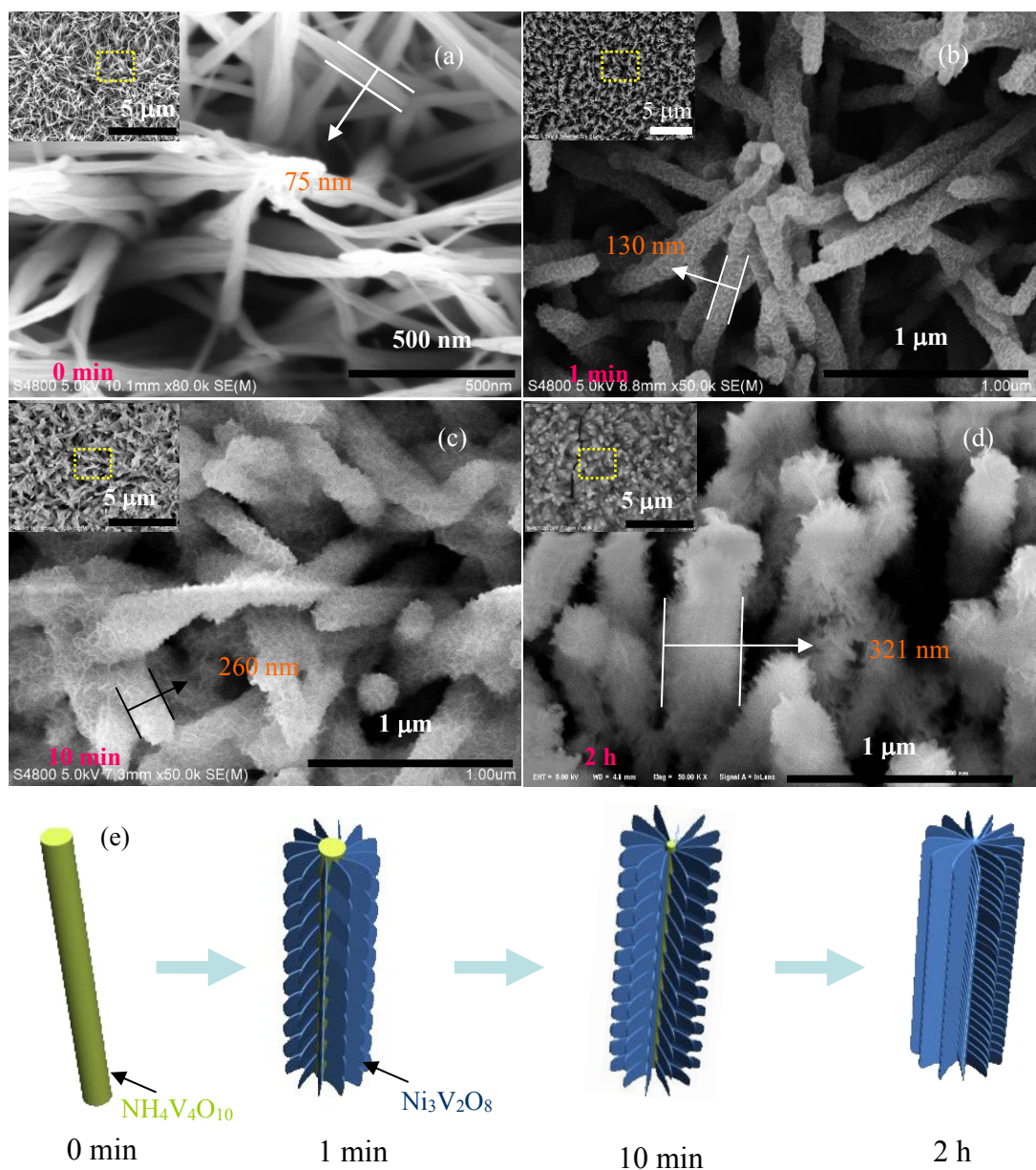


Figure S3. the samples obtained by a second hydrothermal reaction of vanadate ammonia and nickel salt with different reaction time: (a) 0 min, (b) 1 min, (b) 10 min and (b) 2 h.

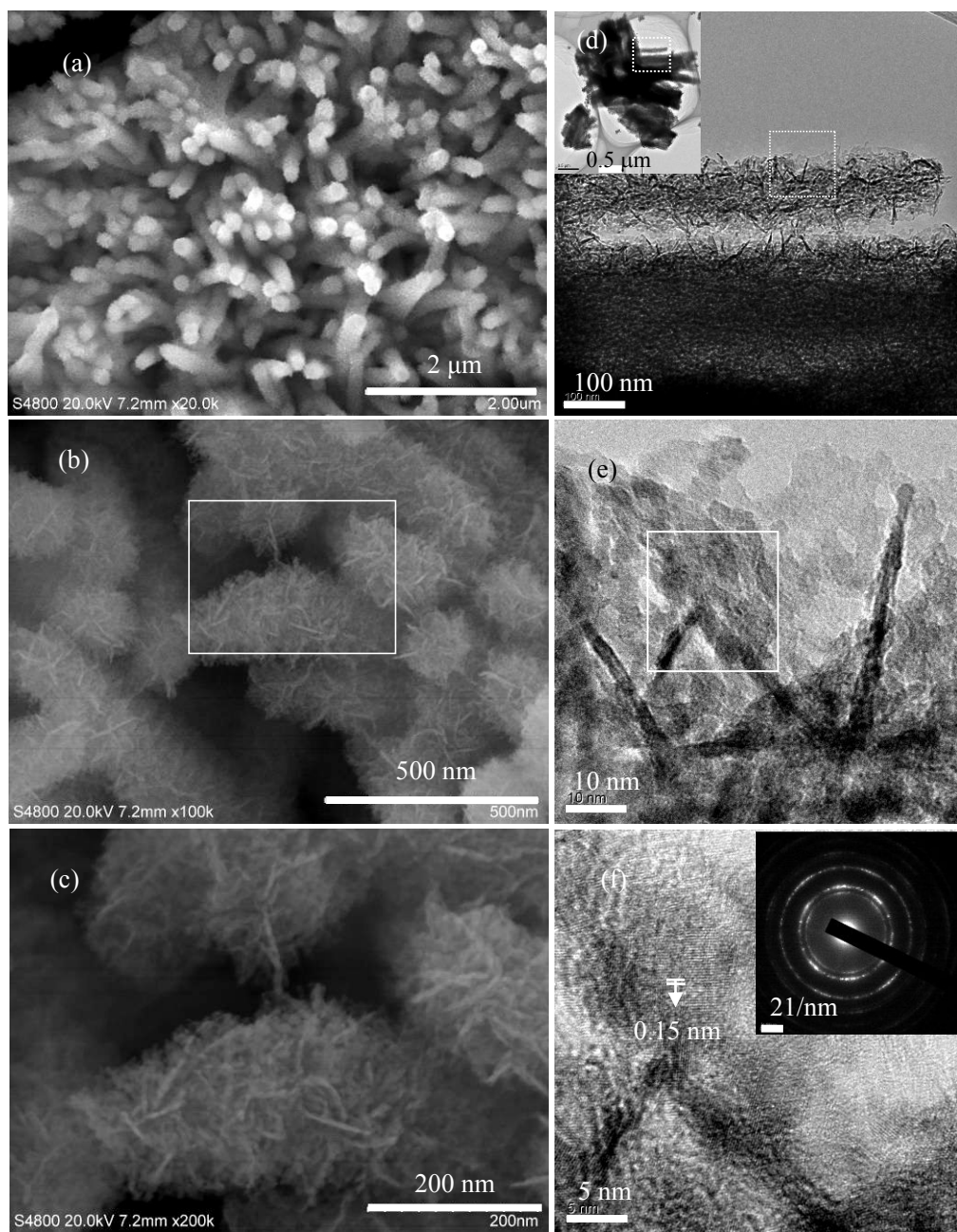


Figure S4. Typical FE-SEM images of the $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays with 300 °C heat treatment at different magnifications. Scale bars, (a) 2 μm ; (b) 500 nm; (c) 200 nm; (d) and (e) TEM images of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays (300 °C) detached from Ti foil, and (f) HR-TEM image (inset: SAED pattern).

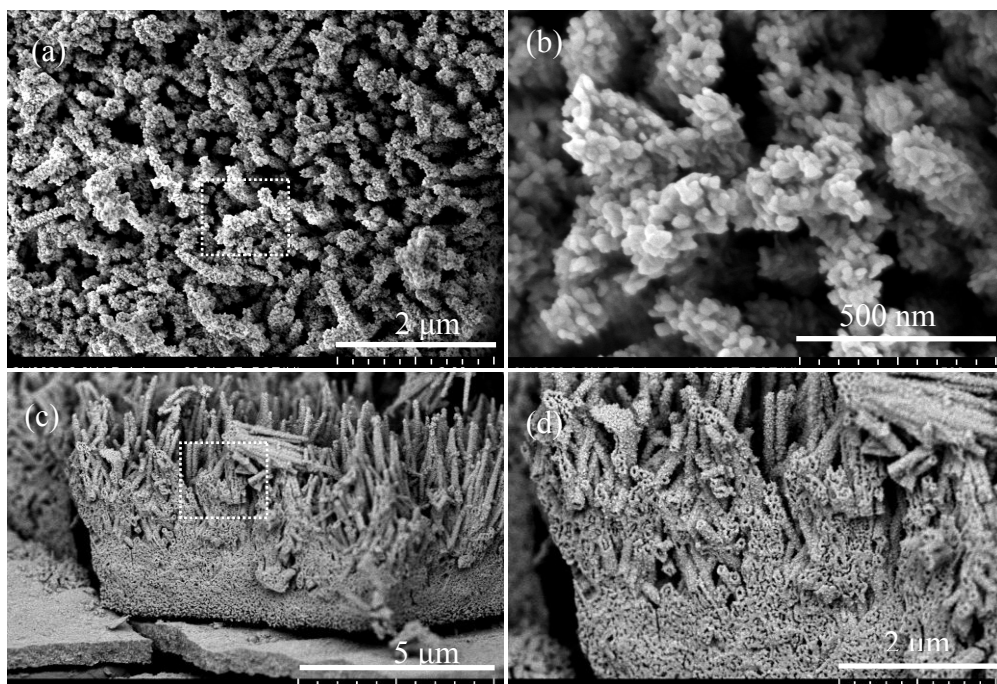


Figure S5. Typical FE-SEM images of the $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays growing on Ti foil with 500 °C heat treatment at different magnifications: a) and b) top-view; c) and d) cross-section view.

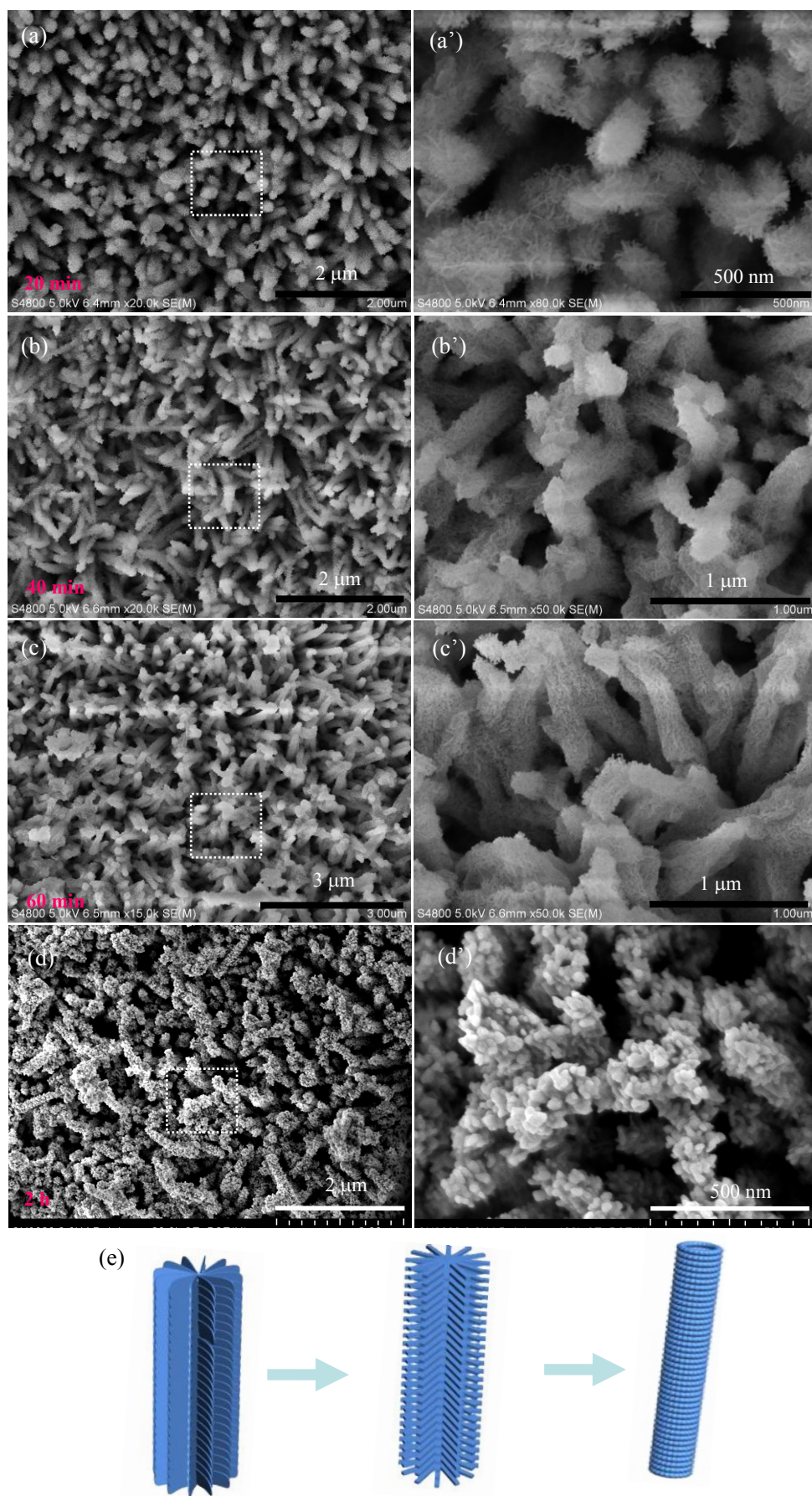


Figure S6. The morphology evolution of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire composed by nanosheets

with increasing the reaction time at 500 °C.

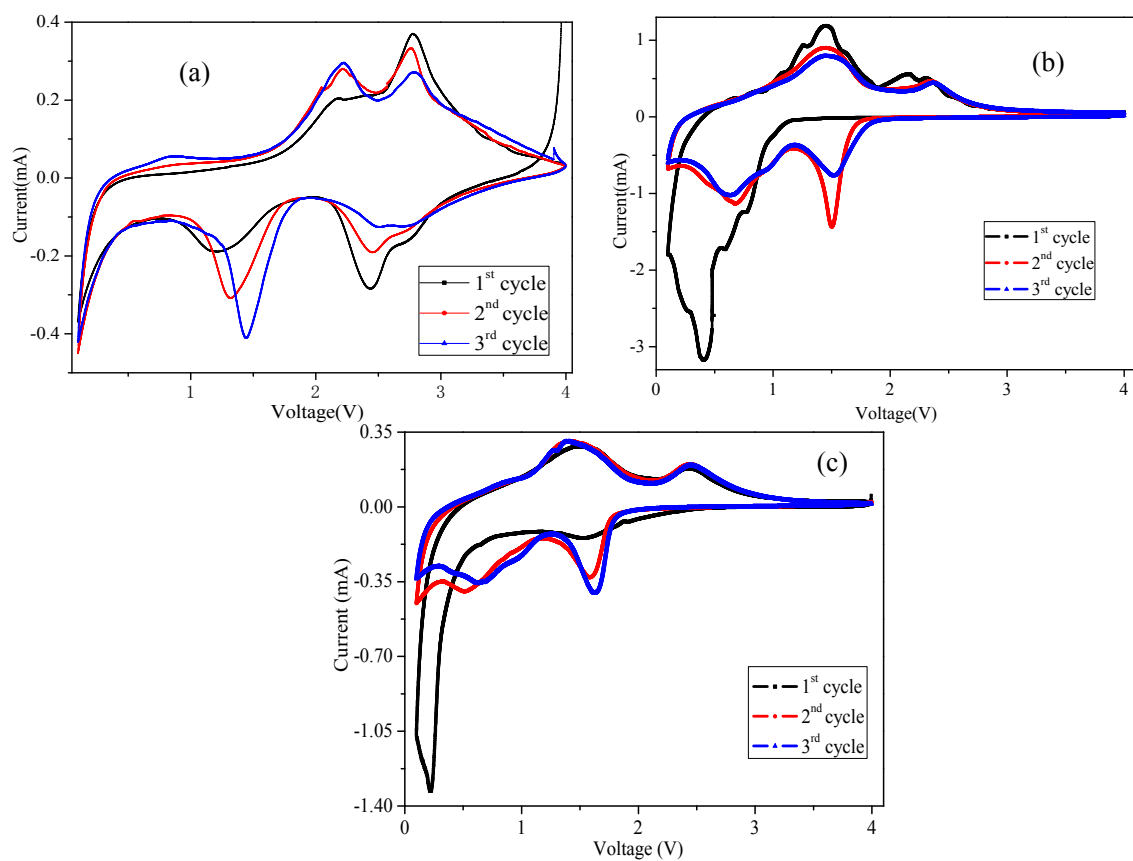


Figure S7. (a)-(c) show plots of the initial three the cyclic voltammogram (CV) curves at 0.1 mV·s⁻¹: (a) NH₄V₄O₁₀ nanowire arrays; (b) the as-prepared Ni₃V₂O₈ nanowire arrays; and (c) Ni₃V₂O₈ nanowire arrays after annealing at 300 °C.

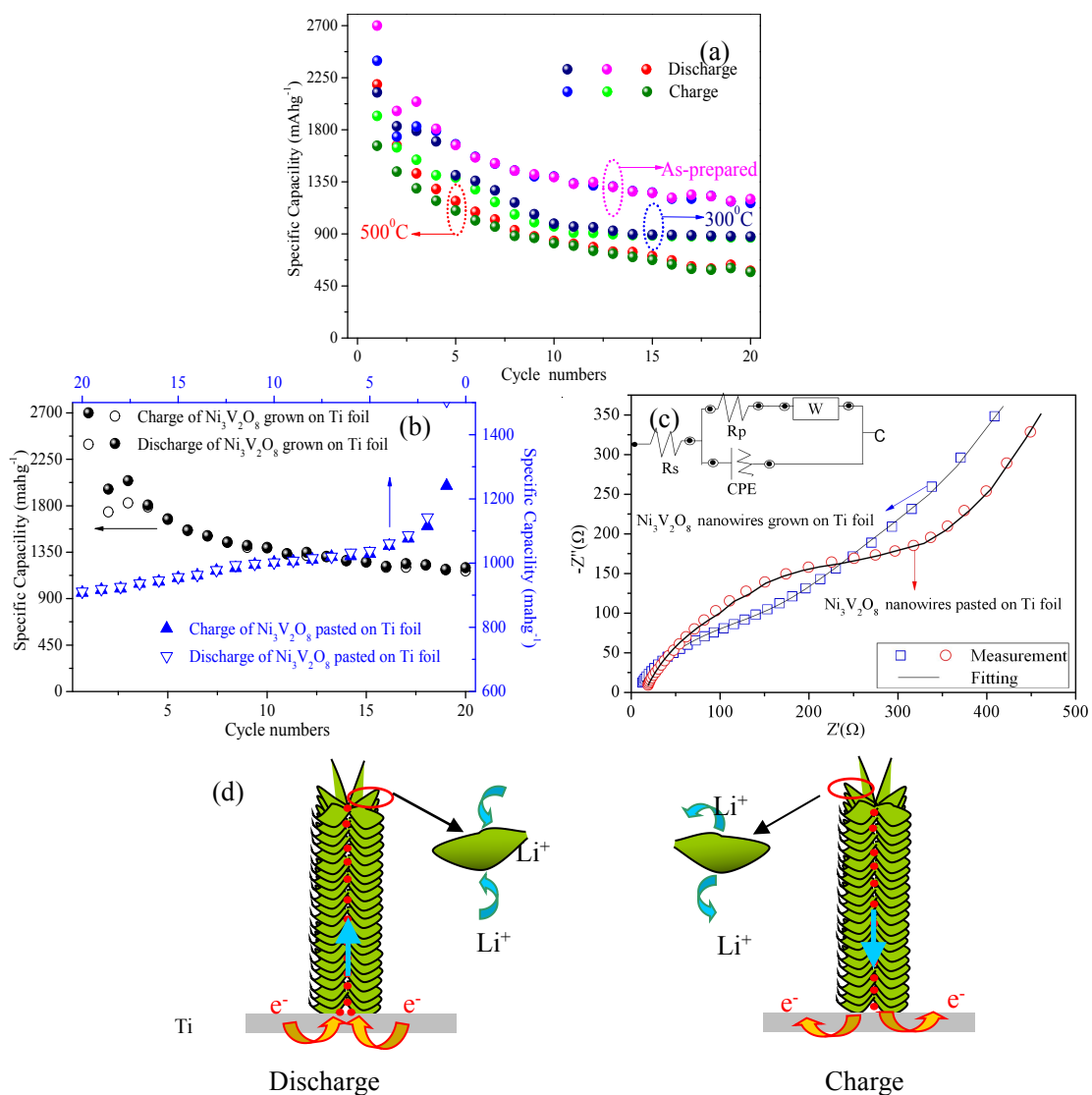


Figure S8. (a) Cycling performances of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays electrodes calcinated at different temperatures at 300 $\text{mA}\cdot\text{g}^{-1}$; (b) Cycling performances of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays grown or pasted on the Ti foil at 300 $\text{mA}\cdot\text{g}^{-1}$; (c) Nyquist plots of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays in the frequency range from 5 kHz to 0.01 Hz, and the inset is the equivalent circuit; and (d) Schematic of charge storage of the $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire array in LIBs.

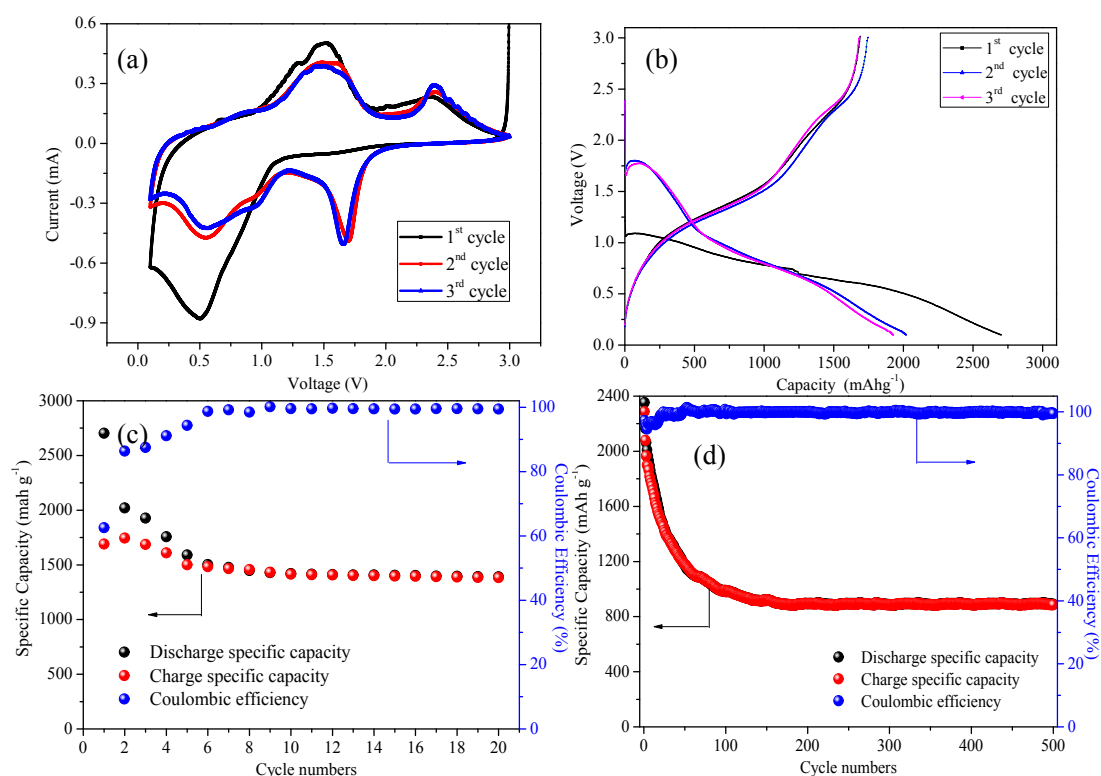


Figure S9. Electrochemical lithium storage properties of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays grown on Ti foil between 3.0-0.1 V: (a) the CV curves at a scan rate of $0.1 \text{ mV}\cdot\text{s}^{-1}$; (b) the initial three charge/discharge voltage profiles at a constant current density of $50 \text{ mA}\cdot\text{g}^{-1}$; and the cycling performance at a current density of (c) $50 \text{ mA}\cdot\text{g}^{-1}$ or (d) $500 \text{ mA}\cdot\text{g}^{-1}$.

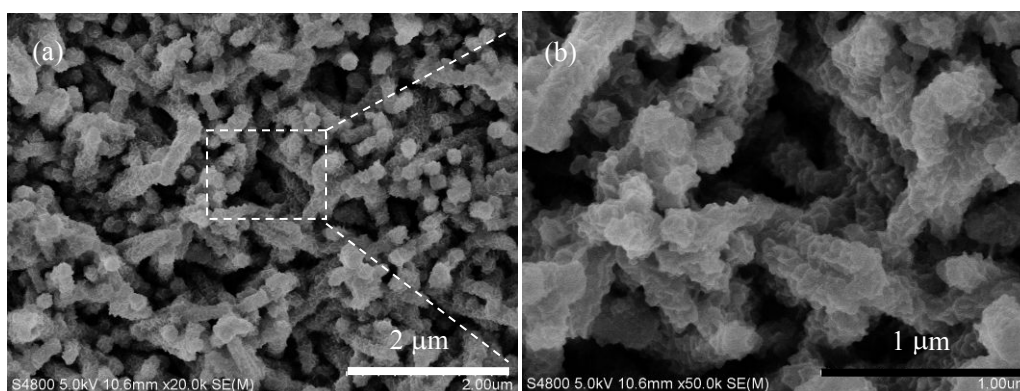


Figure S10. SEM images of the $\text{Ni}_3\text{V}_2\text{O}_8$ electrode after 50 cycles at a charge/discharge current of $500 \text{ mA}\cdot\text{g}^{-1}$.

Table S1. Structure parameters of $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays after annealing at different temperatures

2θ (°)	FWHM	Crystallite size (nm)	d space	Phase
Room Temperature (25 °C)				
35.68	1.80	8.29	2.57	221
43.36	1.54	5.42	2.09	240
63.83	2.29	3.95	1.49	244
300 °C				
35.82	1.54	8.77	2.50	221
43.94	1.12	7.10	2.07	240
63.87	1.41	6.50	1.46	244
500 °C				
35.99	1.35	8.95	2.48	221
43.98	0.86	9.27	2.06	240
63.91	1.07	8.65	1.45	244

Table S2. Electrode resistances for $\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays grown or pasted on Ti file obtained from equivalent circuit fitting of EIS results

	$\text{Ni}_3\text{V}_2\text{O}_8$ nanowire arrays	$\text{Ni}_3\text{V}_2\text{O}_8$ nanowire powder
R_s (Ω)	6.16	15.7
R_p (Ω)	160	272
CPE(Yo) (μmh_0)	18.7	24.6
W (μmh_0)	487	464

[1] P. Soudan, J.P. Pereira-Ramos, G. Gregoire, N. Baffier, The sol-gel mixed oxide $\text{Cr}_{0.11}\text{V}_2\text{O}_{5.16}$: An attractive cathodic material for secondary lithium batteries. *Ionics*, 1997, 3(3-4): 261-264.