

Supporting Information

Green Synthesis of Vanadate Nanobelts at Room Temperature for Superior Aqueous Rechargeable Zinc-Ion Batteries

Zhiqiang Xie^{a,1}, Jianwei Lai^{a,1}, Xiuping Zhu^{b,2}, Ying Wang^{a,*}

^a Department of Mechanical & Industrial Engineering, Louisiana State University, Baton Rouge,
LA 70803, USA.

^b Department of Civil and Environmental Engineering, Louisiana State University, Baton Rouge,
LA 70803, USA.

* Corresponding author: Prof. Ying Wang, E-mail: ywang@lsu.edu.



Figure S1. The SEM image of commercial V_2O_5 powder.

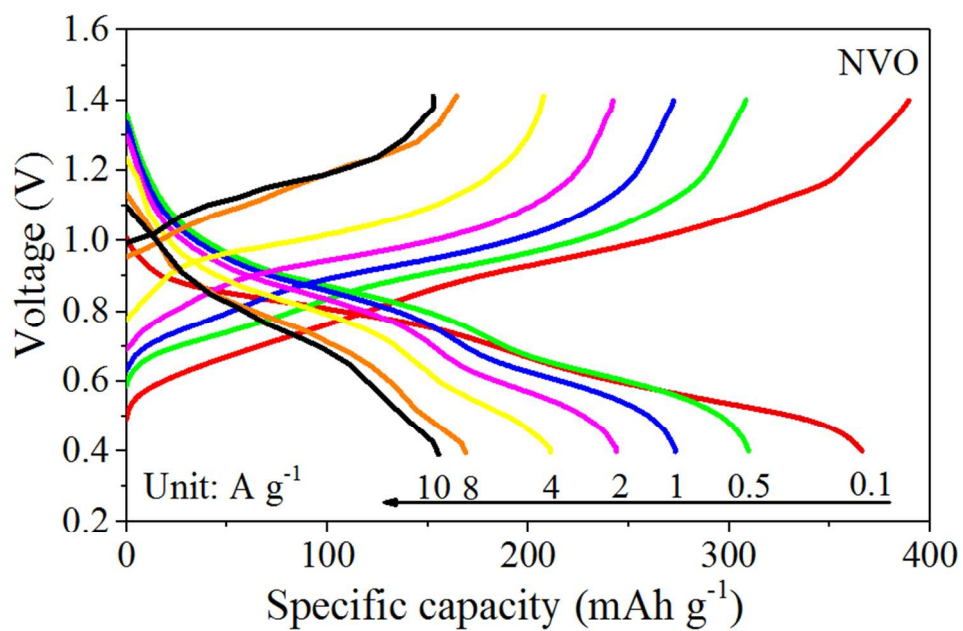


Figure S2. Charge-discharge curves of the NVO at different current densities ranging from 0.1 to 10 A g^{-1} .

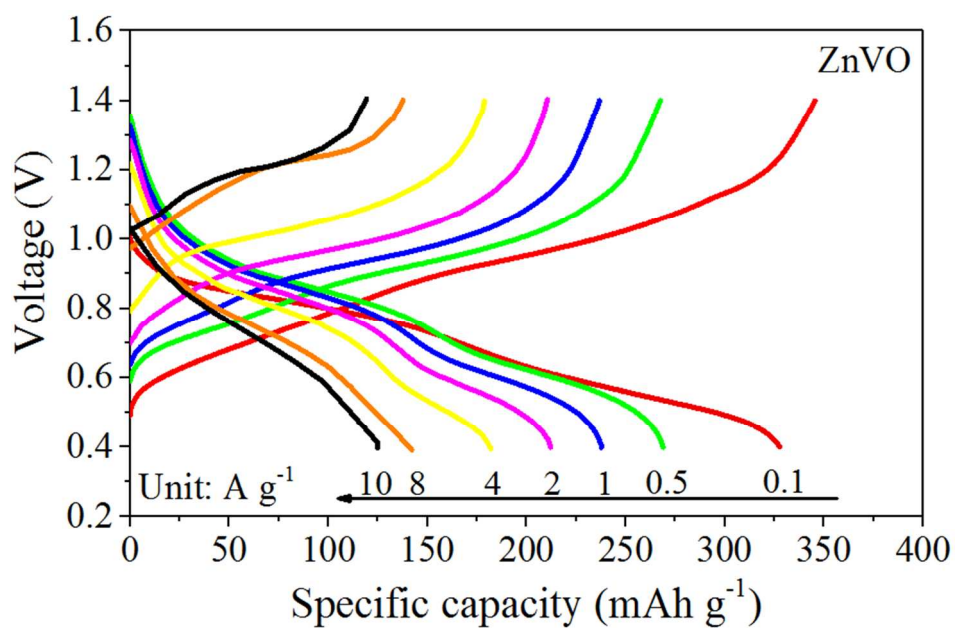


Figure S3. Charge-discharge curves of the ZnVO at different current densities ranging from 0.1 to 10 A g⁻¹.

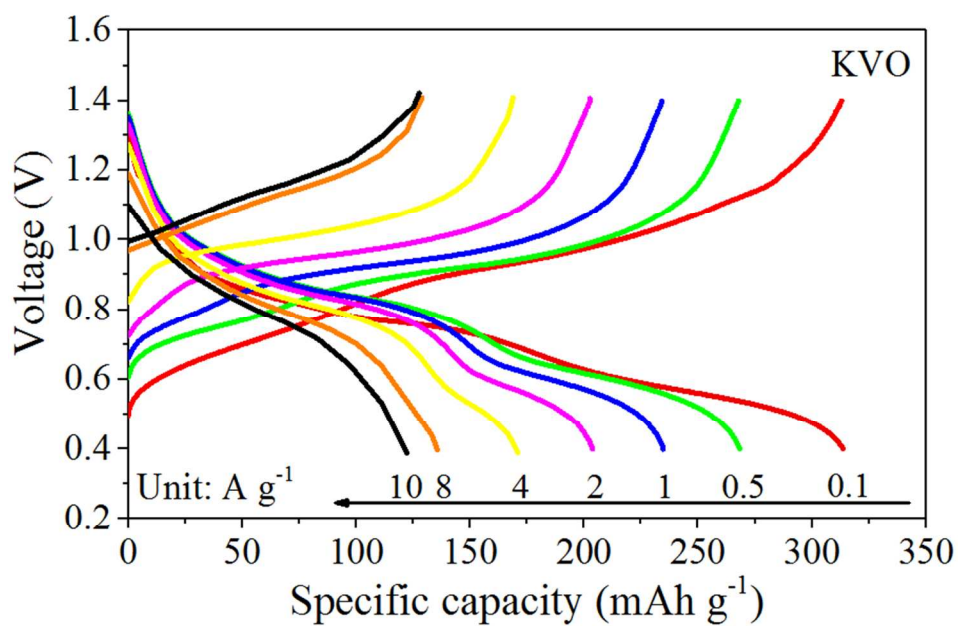


Figure S4. Charge-discharge curves of the KVO at different current densities ranging from 0.1 to 10 A g⁻¹.

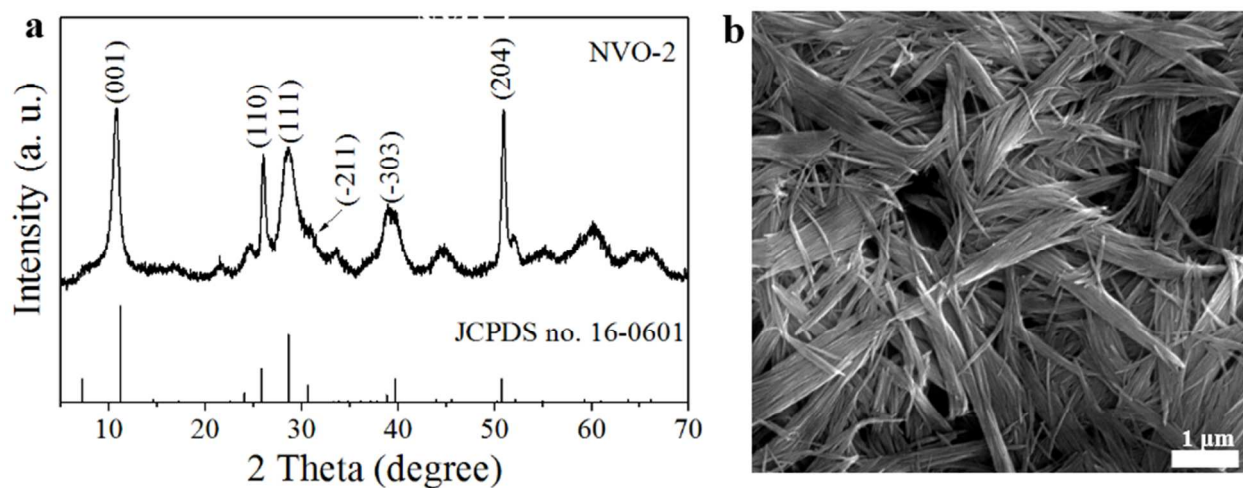


Figure S5. (a) X-ray diffraction pattern of the NVO-2 nanobelts, (b) corresponding SEM image.

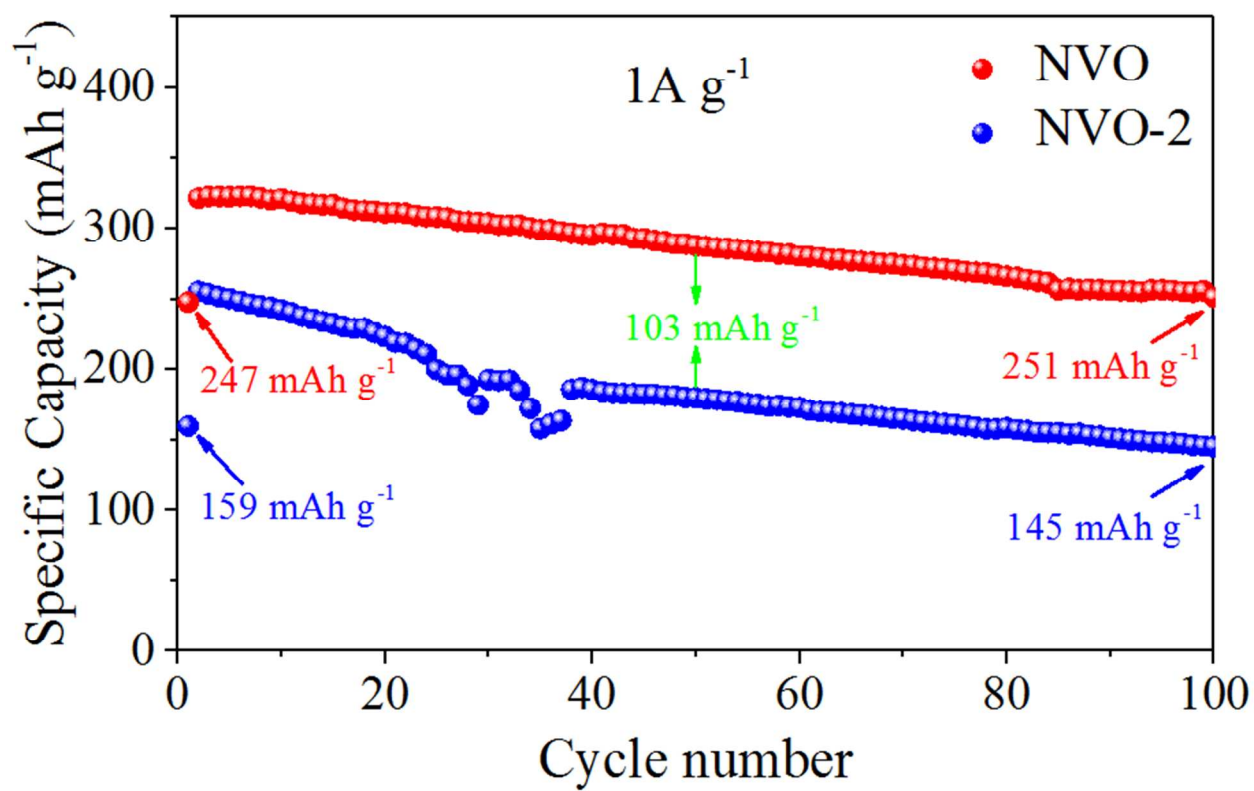


Figure S6. Cycling performances of NVO and NVO-2 at 1 A g⁻¹.

Table S1. Comparison of our work with recently reported vanadium-based cathode materials on the synthesis technique and electrochemical performances for zinc-ion battery.

Materials	Synthesis technique	Precursors	Electrolyte	Specific capacity (mAh g ⁻¹)/ current density (mA g ⁻¹)	Ref.
NaV ₃ O ₈ ·1.35H ₂ O	Room-temperature stirring for 72 h	V ₂ O ₅ , NaCl	1 M ZnSO ₄	366/ 100	Present work
Zn ₃ V ₂ O ₈ ·1.85H ₂ O	Room-temperature stirring for 144 h	V ₂ O ₅ , ZnSO ₄ ·7H ₂ O	1 M ZnSO ₄	328/ 100	Present work
KV ₃ O ₈ ·0.51H ₂ O	Room-temperature stirring for 72 h	V ₂ O ₅ , KCl	1 M ZnSO ₄	316/ 100	Present work
Zn _{0.25} V ₂ O ₅ ·nH ₂ O	Microwave method at 180 °C for 90 min	V ₂ O ₅ , acetone, zinc acetate	1 M ZnSO ₄	300/ 50	1
Zn ₂ V ₂ O ₇	Hydrothermal method at 210 °C for 48 h	Zn(NO ₃) ₂ ·6H ₂ O, NH ₄ VO ₃	1 M ZnSO ₄	250/ 50	2
Zn ₃ V ₂ O ₇ (OH) ₂ ·2H ₂ O	Microwave method at 180 °C for 6 h	Zn(NO ₃) ₂ ·2H ₂ O, NH ₄ VO ₃	1 M ZnSO ₄	200/ 50	3
K ₂ V ₈ O ₂₁	Hydrothermal method at 180 °C for 48 h	V ₂ O ₅ , PEG, KOH	2 M ZnSO ₄	247/300	4
VO _{1.52} (OH) _{0.77}	Hydrothermal method at 150 °C for 24 h	VOCl ₃ , benzyl alcohol	1 M ZnSO ₄	140/ 15	5
LiV ₃ O ₈	Anneal method at 400 °C for 12 h	V ₂ O ₅ , LiCO ₃	1 M ZnSO ₄	280/ 16	6
VS ₂	Hydrothermal method at 180 °C for 20 h	NH ₄ VO ₃ , NH ₃ ·H ₂ O, thioacetamide	1 M ZnSO ₄	190/ 50	7
NaV ₃ O ₈ ·1.5H ₂ O	Stirring at 30 °C for 96 h	V ₂ O ₅ , NaCl	1M ZnSO ₄ / 1M Na ₂ SO ₄	380/ 100	8
Na ₃ V ₂ (PO ₄) ₃	Hydrothermal method at 180 °C for 48 h with post anneal method	V ₂ O ₅ , NH ₄ H ₂ PO ₄ , Na ₂ CO ₃ , citric acid, PEG	0.5 M Zn(CH ₃ COO) ₂	97/ 50	9
Na ₂ V ₆ O ₁₆ · 1.63 H ₂ O	hydrothermal method at 180 °C for 24 h	V ₂ O ₅ , NaOH	3M Zn(CF ₃ SO ₃) ₂	353/ 50	10

References

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