Electronic Supplementary Information (ESI) for

Silver vanadium oxide@water-pillared vanadium oxide coaxial nanocables for superior zinc-ion storage properties

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Fig. S2 TEM and HRTEM images of $V_2O_5/V_2O_5 \cdot nH_2O$ (a and b) and $Ag_{0.333}V_2O_5$ (c and d).

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and (c) $V_2O_5/V_2O_5 \cdot nH_2O_1$.

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Fig. S9 SEM images of initial $Ag_{0.333}V_2O_5@V_2O_5.nH_2O$ cathode (a and b); and after cycled $Ag_{0.333}V_2O_5@V_2O_5.nH_2O$ cathode (c and d); The insets in part a and c shows the status of the electrodes.

Fig. S10 The long-time cycling performance of the electrodes for comparison at 3.0 A g^{-1} .

Fig. S11 EDS images of the thick $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ electrode.



Fig. S1 TEM images (a,b and c) of the Ag_{0.333}V₂O₅@V₂O₅·nH₂O coaxial nanocables.



Fig. S2 Typical TEM and HRTEM images of V2O5/V2O5 nH2O (a and b) and Ag0.333V2O5 (c and



Fig. S3 TEM-EDS image of the elemental distributions of O in Ag_{0.333}V₂O₅@V₂O₅·*n*H₂O.

					Spec	run
	•	Ĩ	Element	Weight%	Atomic%	
			ОК	26.97	55.33	
			νк	65.98	42.52	
2			Ag L	7.05	2.15	
	ĩ	9				
2 Il Scale 1105 cts C	4 ursor: 0.000	6	8 10	12 14	16 18	-

Fig. S4 EDS spectrum of the $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ coaxial nanocables.

The EDS result demonstrated the atomic ratio of Ag to V is about 0.05, which is consistent with

the results of XPS (Table S1).

Name	Start BE	Peak BE	End BE	Height CPS	FWHM eV	Area (P) CPS.eV	Area (N) TPP-2M	Atomic %
C1s	293.83	284.79	280.83	10708.22	1.56	22954.87	0.29	23.92
Ag3d	378.48	367.82	364.23	10067.82	1.13	21629.04	0.01	1.05
V2p	528.78	517.58	512.68	57820.53	1.38	147450.9	0.22	17.92
O1s	543.98	530.41	527.88	72386.79	1.54	145134.67	0.69	57.11

Table S1 The detailed analysis data in XPS experiments for varied elements.



Fig. S5 The first cycle discharge-charge profiles of (a) $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$; (b) $Ag_{0.333}V_2O_5$;

and (c) $V_2O_5/V_2O_5 \cdot nH_2O_.$

The first cycle discharge-charge specific capacity of $Ag_{0.333}V_2O_5 @V_2O_5 \cdot nH_2O$ (210.18 mAh g⁻¹ at 0.5 A g⁻¹), $Ag_{0.333}V_2O_5$ (180.56 mAh g⁻¹ at 0.2 A g⁻¹), $V_2O_5/V_2O_5 \cdot nH_2O$ (135.03 mAh g⁻¹ at 0.2 A g⁻¹) are shown in Fig. S4, respectively.



Fig. S6 Nyquist impedance plots of varied electrodes fresh cells.



Fig. S7 Discharge–charge curves of the $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ cathode at different current densities.

The discharge–charge curves of the $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ cathode at different current densities shows in Fig. S6. The charge–discharge curves features are kept well, even in a higher current density of 3.0 A g⁻¹. The specific capacities 373.3, 322.5, 290.62, 274.18, 262.72, 253.76 and 245.9 mAh g⁻¹ are obtained at 0.2, 0.5, 1.0, 1.5, 2.0, 2.5 and 3.0 A g⁻¹, respectively.



Fig. S8 SEM images of $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ cathode after 100 cycles at 0.5 A g⁻¹.



Fig. S9 SEM images of initial $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ cathode (a and b); and after cycled $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ cathode (c and d); The insets in part a and c shows the status of the electrodes.



Fig. S10 The long-term cycling performance of the electrodes for comparison at 3.0 A $g^{-1}.$



Fig. S11 EDS images of the thick $Ag_{0.333}V_2O_5@V_2O_5 \cdot nH_2O$ electrode.