Supporting information

Highly improved rechargeable stability for lithium/silver vanadium oxide battery induced via electrospinning technique

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Supplementary EDS results for β-SVO NRs



Fig. S1 Energy dispersive X-ray spectroscopy (EDS) results for β-SVO NRs of peaks for all possible elements.



Fig. S2 EDS results for β-SVO NRs waving peaks for carbon and copper due to the carbon coated Copper TEM grid.

Spectrum processing: Peaks possibly omitted: 0.931, 1.748, 8.033, 8.899 keV

Quantitation method: Cliff Lorimer thin ratio section.

Processing option: All elements analyzed (Normalised)

Number of iterations = 1

Element	Peak Area	Area Sigma	k factor	Abs Corrn.	Weight%	Weight% Sigma	Atomic%
O K	1608	110	1.403	1.000	28.67	1.52	60.51
V K	4746	115	0.813	1.000	49.06	1.40	32.52
Ag L	1349	100	1.299	1.000	22.26	1.38	6.97

Table S1. Atomic percentages and obtained parameters for β -SVO NRs in EDS measurement.

Complementary images for materials characterization



Fig. S3 XRD pattern for electrospun VOx nanofibers, corresponding to oxide precursors in step b demonstrated in Scheme 1.



Fig. S4 TEM image for the surface on β -SVO NRs presenting nanopores between decorated Ag nanodots.



Supplementary information on the electrochemical performance of β-SVO NRs

Fig. S5 Discharge capacity vs. cycle number plots of cell Li/ β -SVO NRs (stored for more than four months) in the voltages range of 2–3.6 V at 20 mAg⁻¹, 40 mAg⁻¹, 100 mAg⁻¹ and 200 mAg⁻¹, respectively at room temperature.



Fig. S6 *a*. XRD pattern for β -SVO NRs after cycling for more than 30 cycles in charged and discharged state; *b*. TEM images (with SAED pattern) for β -SVO NRs recharged back to 3.6 V.



Electrochemical performance in the comparison study with β-AgVO₃ NW

Fig. S7 Capacity vs. cycle number plots of cell Li/ β -AgVO₃ NW in the voltage range of 2-3.6 V at 20 mAg⁻¹ in comparison with β -SVO NRs.



Fig. S8 Capacity vs. cycle number plots of cell Li/ β -SVO NRs in the voltage window of 1.5-3.6 V and 2-3.6 V, respectively, at 20 mAg⁻¹.