Electronic Supplementary Information

Peptide-Templated Synthesis of Branched MnO₂ Nanowires with Improved Electrochemical Performances

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Fig. S1 (a) Molecular structure of I_3K and (b) TEM image of I_3K nanofibers. The three Ile residues are marked in green and the Lys group at the C-terminal in red for clarity. Inset of (b) shows the statistical diameter histogram of the I_3K nanofibers.



Fig. S2 (a) TEM and (b) HR-TEM images of the MnO_2 precipitates formed in the absence of peptide (MP-0). Inset of (a) indicates the size distribution of the formed urchin-like MnO_2 spheres.



Fig. S3 FTIR spectrum of the collected MnO₂ precipitates in the presence of I₃K nanofibers (MP-5).



Fig. S4 The diameter distributions of the formed MnO₂ hybrid nanowires: (a) MP-2, (b) MP-3, (c) MP-4, (d) MP-5, (e) MP-6, and (f) MP-7.



Fig. S5 TGA curves of MP-5 and the urchin-like MnO_2 sphere (MP-0) prepared in the absence of I_3K assemblies. The heating rate was set as 20 °C min⁻¹.



Fig. S6 (a) N₂ adsorption/desorption isotherm and (b) BJH pore size distribution of MP-0.



Fig. S7 (a) N1s XPS spectrum of the branched MnO₂ nanowire (MP-5) and (b) The UV-vis absorbance

at 526 nm as a function of time for different reaction systems.



Fig. S8 The anodic peak current versus the square root of scan rate for the branched MnO_2 nanowire (MP-5), showing a linear relationship between them.



Fig. S9 (a) CV curves of the branched MnO_2 nanowire (MP-5) as the electrode material in aqueous 1 M Na_2SO_4 at different scan rates and (b) Variation of its specific capacitance with the scan rate.