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Supporting Information



Fig.S1 SEM image of pure Fe₃O₄ nanoparticles prepared by coprecipitation method



Fig.S2 SEM images of the precursors $[NH_3(CH_2)_6NH_3][Fe_{1.5}F_3(SO_4)] \cdot 0.5H_2O$, showing their layered structures.



Fig. S3 TG and DTA curves of the precursor $[NH_3(CH_2)_6NH_3][Fe_{1.5}F_3(SO_4)] \cdot 0.5H_2O$ in Ar gas. There are two distinct weight losses consistent with the molecular formula. A sharp weight loss in the region 573-660 K corresponding to the loss of the water molecules, amine and HF $[0.5H_2O+2NH_3+HF, exp = 17.6\%, cal = 17.3\%]$. The second major weight loss in the region 660-973 K corresponds to the removal of remaining fluorine, hydrogen and the decomposition of SO_4^{2-} moiety, $[5.5H_2 + F_2 + SO_2, exp = 31.8\%, cal = 31.0\%]$.



Fig. S4 Low-magnification TEM image of Fe₃O₄-NPs@C sample.



Fig.S5 Charge/discharge capacities of the Fe $_3O_4$ -NPs@C samples calcinated under various temperatures.



Fig.S6 Charge/discharge capacities of the Fe $_3O_4$ -NPs@C samples calcinated for various time.



Fig.S7 Nyquist plots for of Fe₃O₄-NPs@C composite and pure Fe₃O₄ electrodes measured in the frequency range between 0.01 Hz and 1.0 MHz.