

Supporting Information

Immobilization of Dye Pollutant on Iron Hydroxide Coated Substrates:

Kinetics, Efficiency and Adsorption Mechanism

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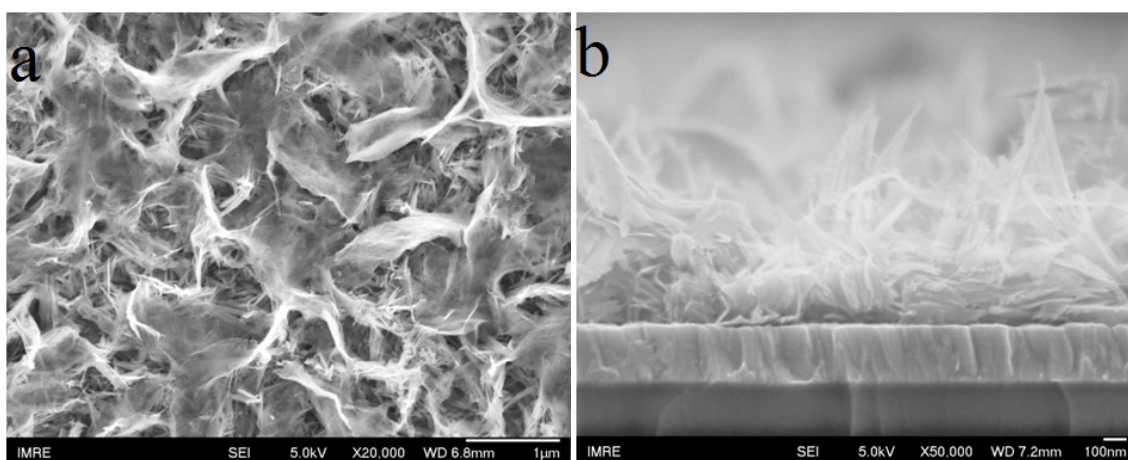


Figure S1 SEM images of (a) surface and (b) cross-section of the FeOOH films deposited with 0.5 M $(\text{NH}_4)_2\text{Fe}(\text{SO}_4)_2$ and 1 M CH_3COONa under a high growth rate. Large and flaky structures with poor adhesion and uniformity are observed.

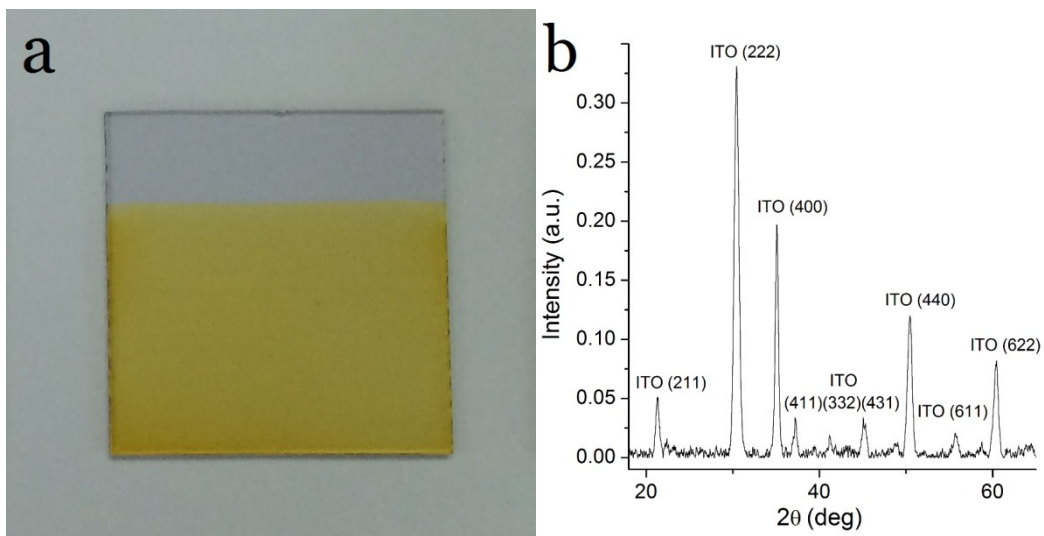


Figure S2 (a) Picture of FeOOH film deposited on ITO under optimized condition showing good uniformity. (b) X-ray diffraction spectrum of the coated FeOOH on ITO. Only diffraction peaks of ITO were observed showing the amorphous nature of the film.

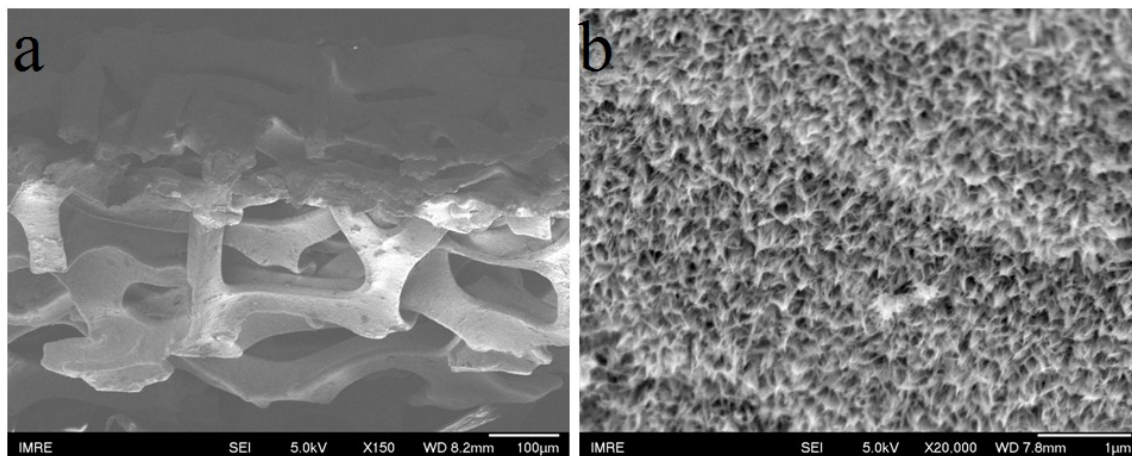


Figure S3 SEM images of the electrodeposited FeOOH films on Ni foam substrate with (a) low and (b) high magnification. The images show the growth of porous structures on the surface of the Ni foam.

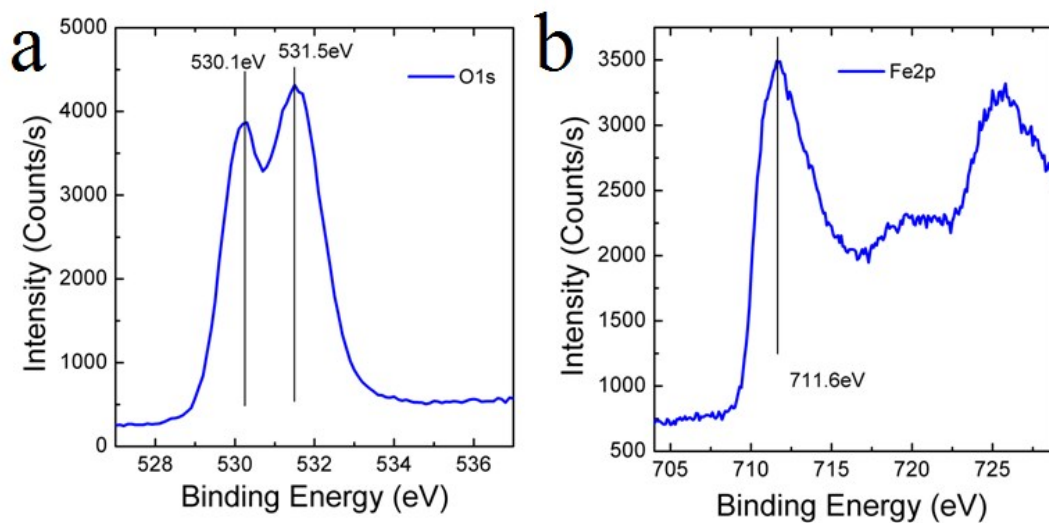


Figure S4 X-ray photoelectron spectra of (a) O 1s and (b) Fe 2p of the FeOOH film deposited on ITO. The binding energy for both O 1s and Fe 2p is consistent with the presence of FeOOH.

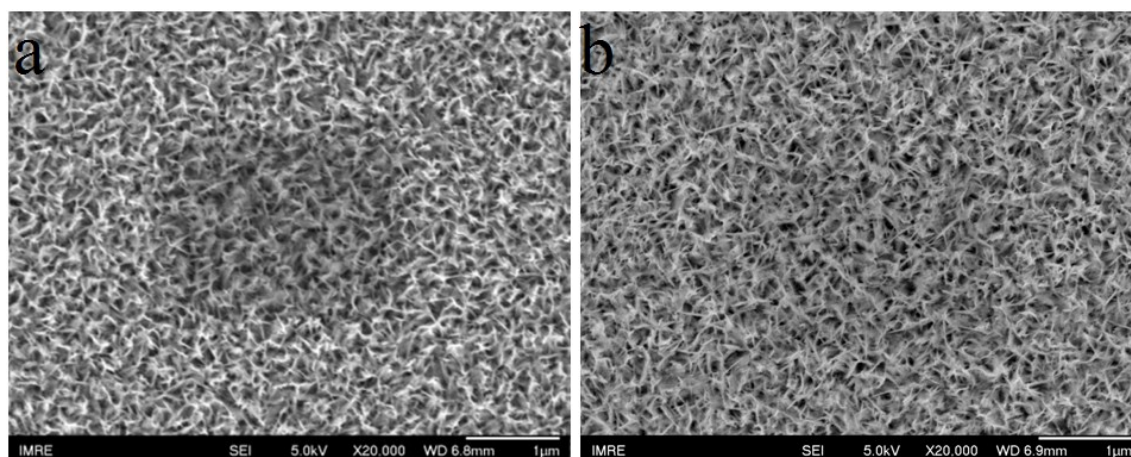


Figure S5 SEM images of FeOOH film (a) before and (b) after the adsorption of Congo red. The images show similar morphology indicating the absence of major structural change.

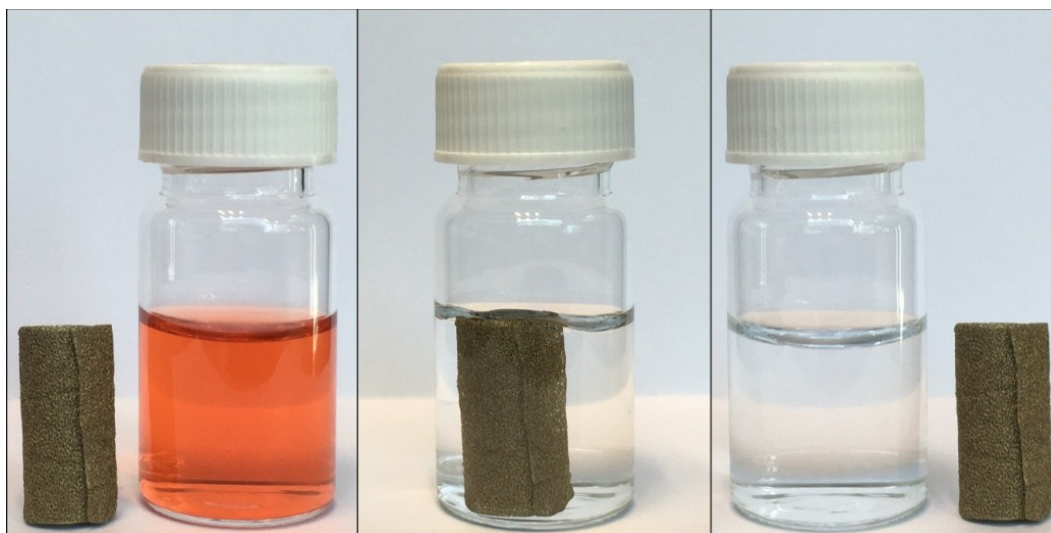


Figure S6 Images of 10 mL of 20 mg/L Congo red solution (left), the solution after 180 minutes immersion of coated FeOOH on Ni foam (middle) and the solution after removal of coated film and contaminants (right). The process above demonstrated easy immobilization and removal of dye pollutant.

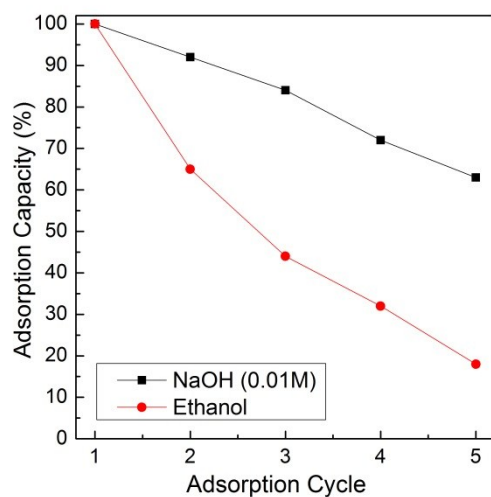


Figure S7 Relative adsorption capacity of the coated FeOOH with respect to the initial adsorption for 5 cycles of adsorption and desorption process. The desorption process was accomplished by a 1 hour soaking in the indicated solutions. We found that each cycle of

adsorption achieved on average 88.9% and 65.1% of the previous adsorption capacity in NaOH solution and ethanol, respectively.

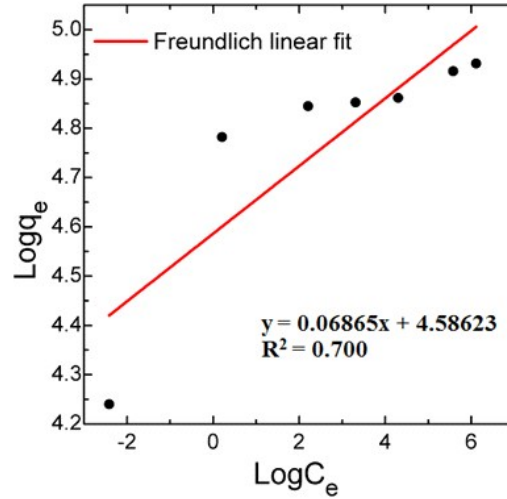


Figure S8 Linearized Freundlich isothermal plots of Congo red adsorption on FeOOH film. A poor correlation coefficient of 0.7 is obtained showing a lack of multilayer adsorption.

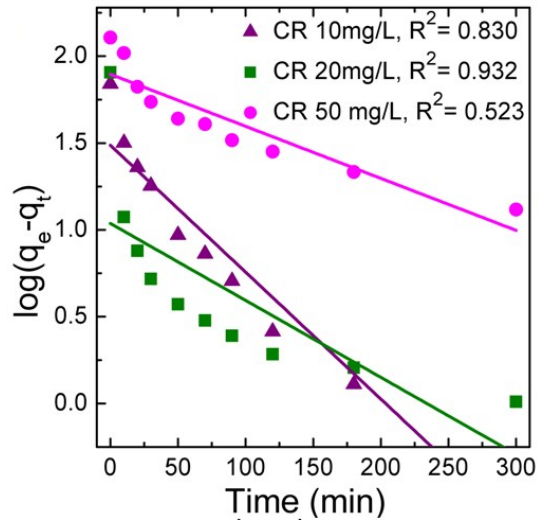


Figure S9 Linearized pseudo-first-order kinetics plots of Congo red adsorption for FeOOH deposited on Ni foam. Poor correlations are obtained for all different Congo red concentrations. The correlation coefficient for each concentration is indicated in the plot.

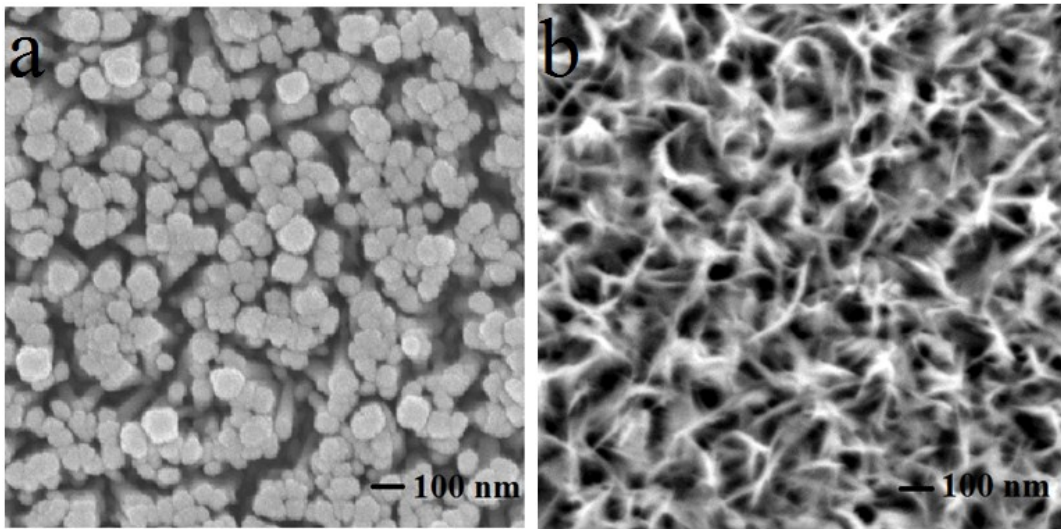


Figure S10 SEM images of (a) hydrothermally grown FeOOH and (b) electrodeposited FeOOH. The hydrothermally grown sample was accomplished by 6 hours of reaction at 100 °C in 10 mL of aqueous solution containing 1.5 mmol FeCl₃•6H₂O and 1.5 mmol NH₂CONH₂. The measured capacity of ~40 mg/g for hydrothermal grown FeOOH is lower than electrodeposited FeOOH reported in this work.

Table S1 A comparison of Congo red adsorption properties of various iron oxide/hydroxide nanostructures

Adsorbents	q_m (mg/g)	References	Year
Coated porous FeOOH thin film	144	This work	2015
Coated FeOOH rods	41	This work	2015
Conventional FeOOH	32	18	2013
Commercial α-Fe₂O₃	42	35	2014
α-FeOOH hierarchical nanostructure by self-assembly	56	18	2013

Urchin-like α-FeOOH hollow spheres	275	16	2012
α-FeOOH hierarchical nanostructure	239	17	2011
Mesoporous Fe₂O₃	53	34	2008
γ-Fe₂O₃ hierarchical nanostructure	58	18	2013
α-Fe₂O₃ hierarchical nanostructure	66	17	2011
Hollow nestlike α-Fe₂O₃	160	29	2012
α-Fe₂O₃ nanoparticles	254	35	2014
α-Fe₂O₃ hollow spheres	195	31	2013

Table S2 Kinetic parameters and correlation coefficients calculated from pseudo-second-order model for adsorption of Congo red on FeOOH film

Congo red concentration (mg L⁻¹)	q_{e,exp} (mg g⁻¹)	q_{e,cal} (mg g⁻¹)	k₂ (g mg⁻¹ min⁻¹)	R²
10	70.351	71.582	1.789× 10 ⁻³	0.999
20	82.637	84.034	6.316× 10 ⁻⁴	0.999
50	128.079	123.001	3.332× 10 ⁻⁴	0.998