In-situ hydroxyl radical generation using the synergism of Co-Ni bimetallic centres of developed nanocatalyst with potent efficiency of degrading toxic water pollutants

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Supporting Information



Fig. S1 Core level spectra of (a) C 1s (b) N 1s (c) O 1s of Co-Ni@CS@Fe₃O₄



Lsec: 200.0 0 Cnts 0.000 keV Det: Octane Plus Det

Fig. S2 EDS spectra of Fe_3O_4 .



Fig. S3Graph plot of degradation efficiency (%) as a function of time (min) (a) 2,4-D (reaction conditions : 50 mL stock solution (100 mg/L), 1 mL H₂O₂, 10 mg catalyst, pH = 7, 60 min, r.t., $\lambda_{max} = 284$ nm) (b) MO (reaction conditions : 50 mL stock solution (10 mg/L), 2 mL H₂O₂, 20 mg catalyst, pH = 7, 60 min, r.t., $\lambda_{max} = 465$ nm).



Fig. S4Effect of catalyst dosage on the catalytic degradation of (a) 2,4-D (reaction conditions : 50 mL stock solution (100 mg/L), x mg catalyst 1 mL H₂O₂, pH = 7, 60 min, r.t., $\lambda_{max} = 284$ nm) (b) MO (reaction conditions : 50 mL stock solution (10 mg/L), x mg catalyst, 2 mL H₂O₂, pH = 7, 60 min, r.t., $\lambda_{max} = 465$ nm).



Fig. S5 Effect of pH on the catalytic degradation of (a) 2,4-D (reaction conditions : 50 mL stock solution (100 mg/L), 10 mg catalyst, 1 mL H₂O₂, 60 min, r.t., $\lambda_{max} = 284$ nm) (b) MO (reaction

conditions : 50 mL stock solution (10 mg/L), 10 mg catalyst, 2 mL H₂O₂, 60 min, r.t., $\lambda_{max} = 465$ nm).



Fig. S6(a) UV-Vis plot of degradation of 2,4-D under optimized conditions (reaction conditions : 50 mL stock solution (100 mg/L), 1 mL H₂O₂, 10 mg catalyst, pH = 7, 60 min, r.t., $\lambda_{max} = 284$ nm) (b) UV-Vis plot of degradation of MO under optimized reaction conditions (50 mL stock solution (10 mg/L), 2 mL H₂O₂, 20 mg catalyst, pH = 7, 60 min, r.t., $\lambda_{max} = 465$ nm).



Fig. S7 Kinetics analysis of oxidative degradation of (a) 2,4-D and (b) MO.



Fig. S8Degradation efficiency versus reaction time in a leaching experiment.



Fig. S9FE-SEM image of recovered catalyst Co-Ni@CS@Fe₃O₄.