

Electronic Supporting Information

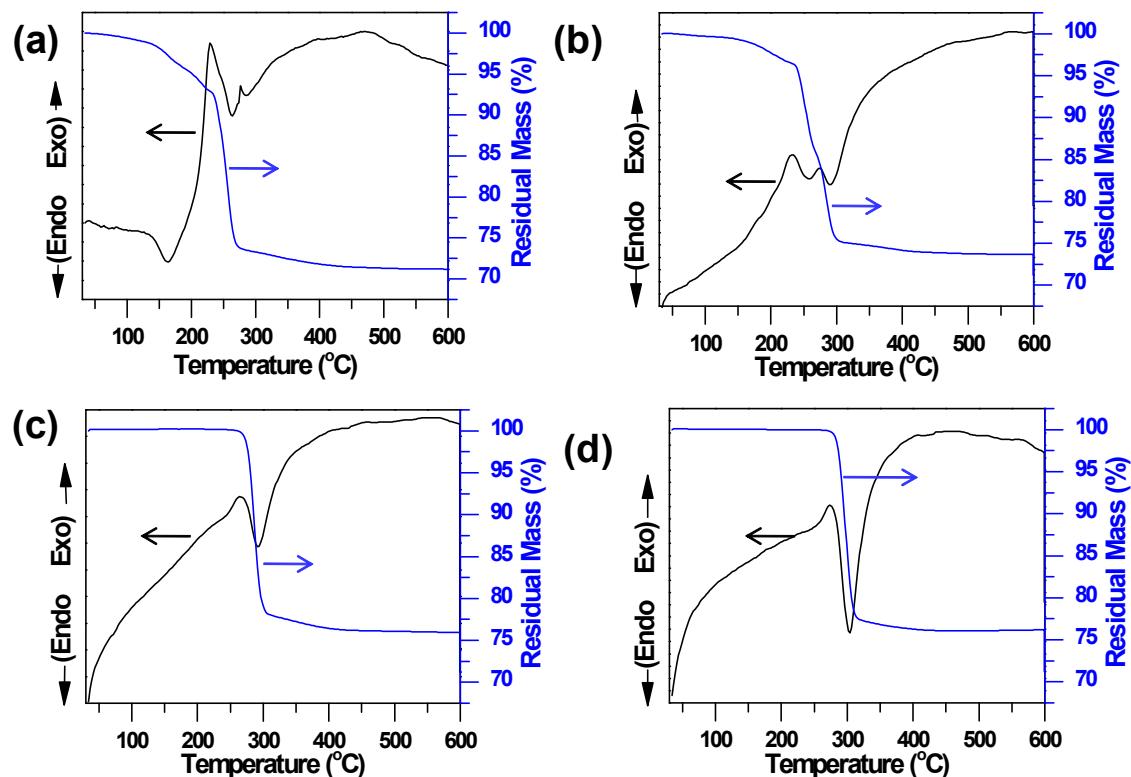


Figure S1: DTA and TG of the as-prepared samples prepared at 150°C for (a) 2h, (b) 5h, (c) 12h and (d) 24h.

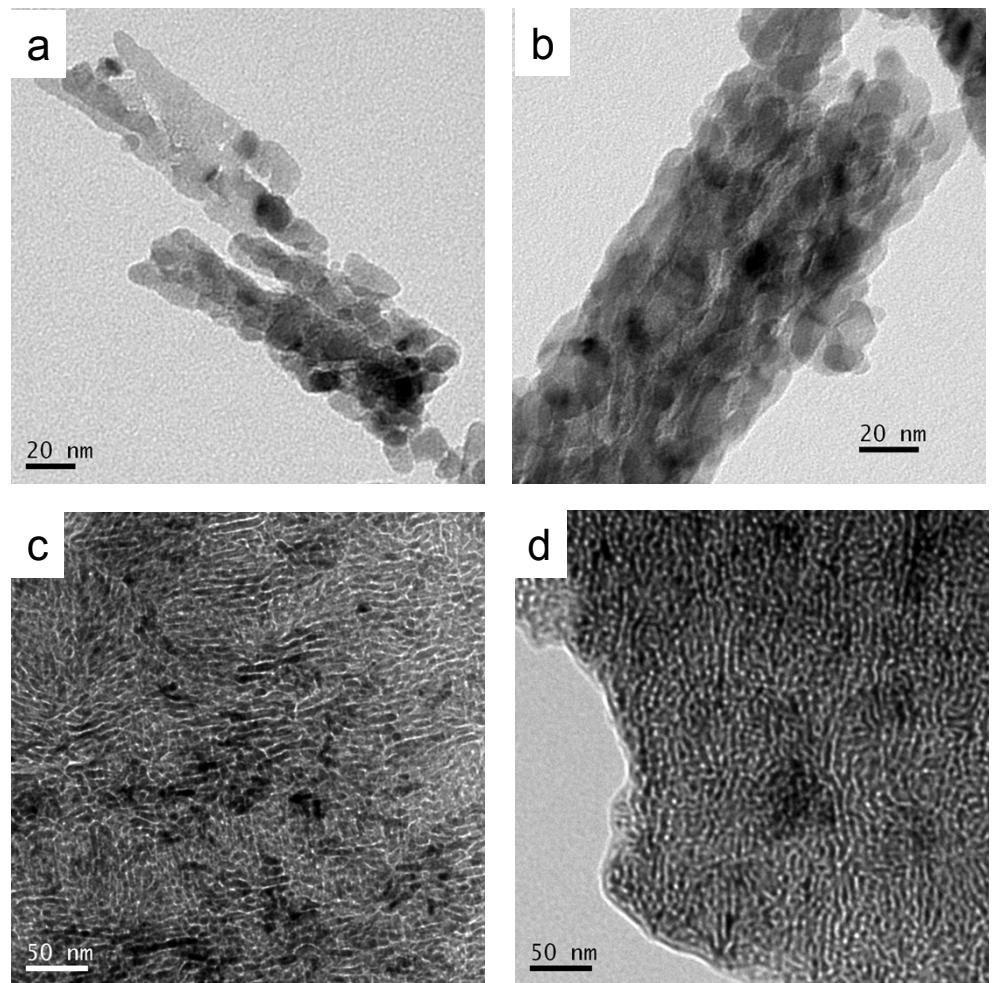


Figure S2: Higher magnification images of TEM.

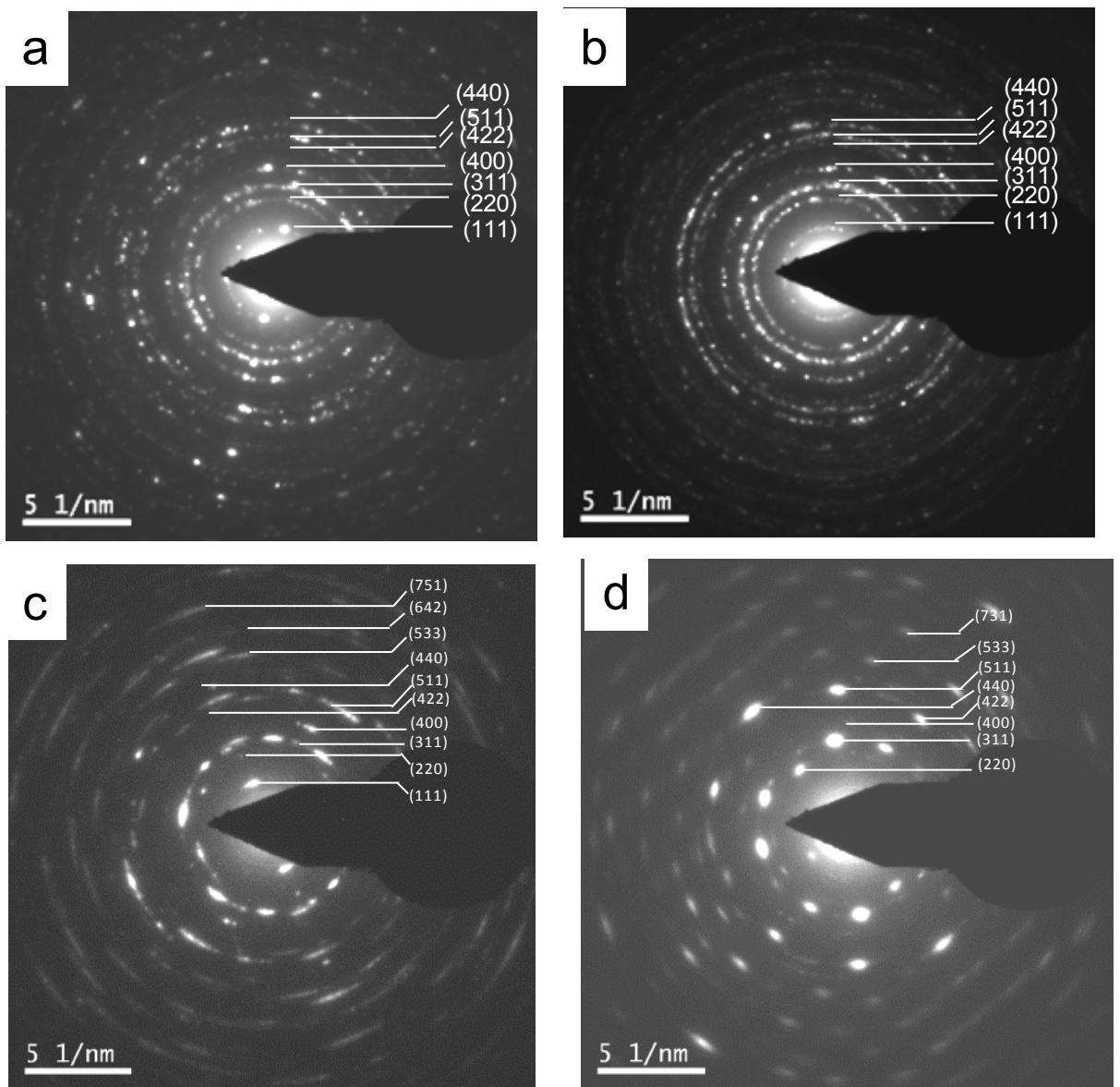


Figure S3: Selected area electron diffraction pattern (SAED) of Co_3O_4 prepared hydrothermally at times (a) 2 h (b) 5 h (c) 12 h and (d) 24 h.

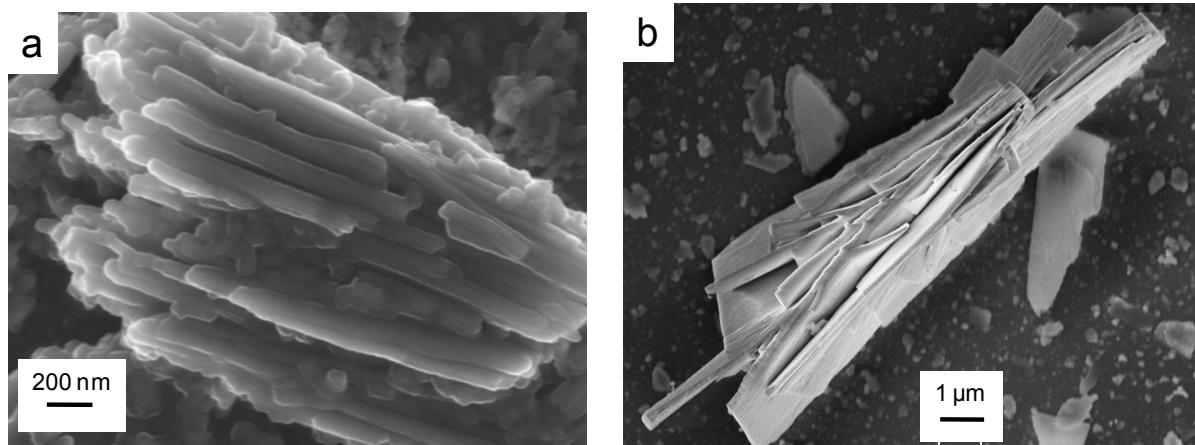


Fig. S4: FESEM images of (a) as-prepared sample synthesized at 150°C/8h, and (b) the corresponding calcined (300°C) sample

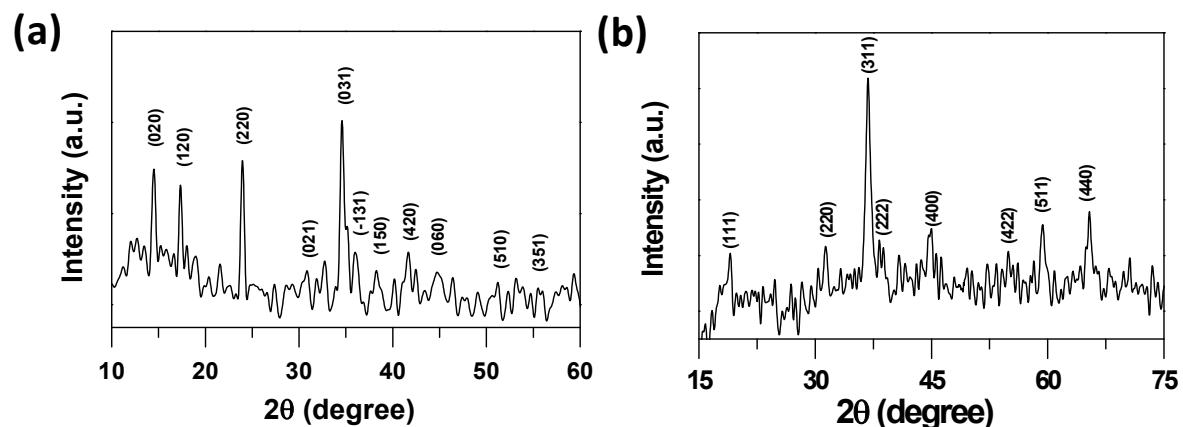


Fig. S5: XRD pattern of (a) as prepared sample synthesized at 150°C/8h, and (b) the corresponding calcined (300°C) sample

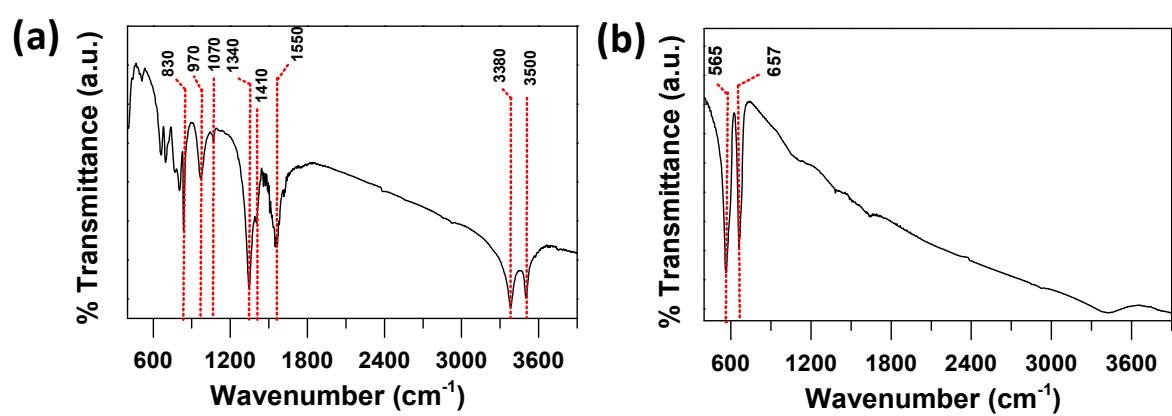


Fig. S6: FTIR of as-prepared sample synthesized at 150°C/8h, and (b) the corresponding calcined (300°C) sample

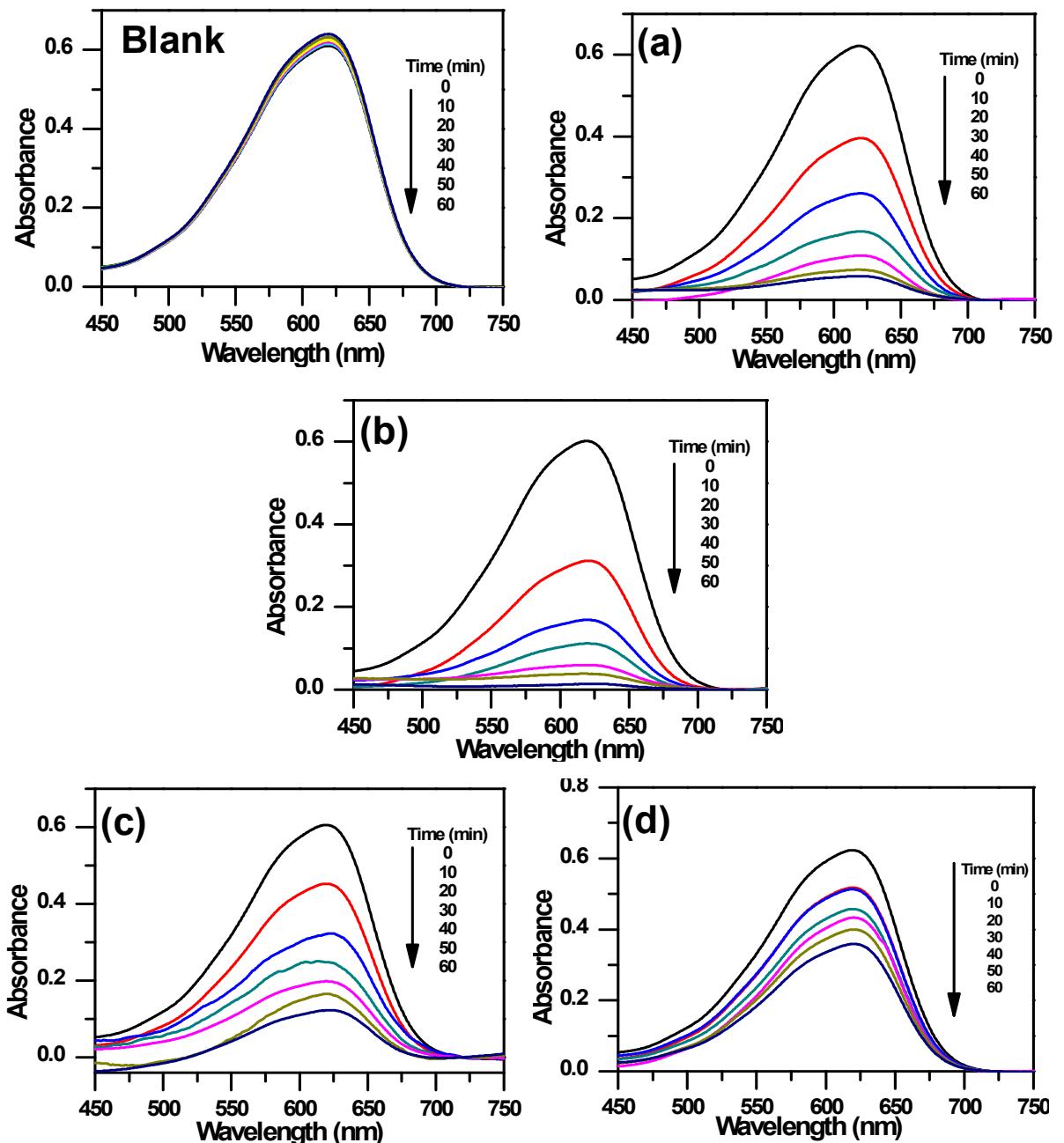


Figure S7: Successive UV-vis absorption spectra with time for the degradation of Chicago sky blue 6B dye at 25 °C using hydrogen peroxide in absence (blank) or in presence of catalyst by Co_3O_4 hydrothermally at 150°C for times (a) 2h (b) 5h (c) 12 h and (d) 24 h.

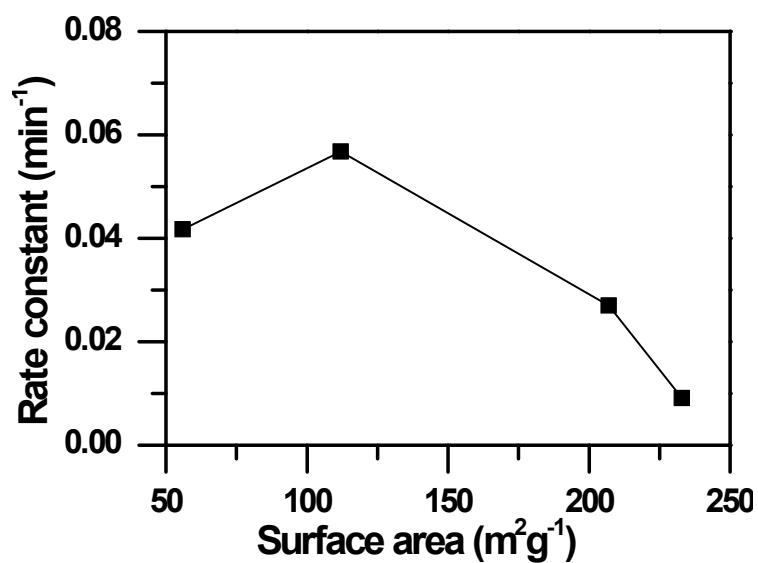


Figure S8: Rate constant vs. surface area plot for the degradation of Chicago sky blue 6B dye using hydrogen peroxide in presence of catalyst Co₃O₄

Table S1: The textural properties of Co_3O_4 particles calcined at 300°C

Synthesis Time (h) at 150°C	S_{BET} (m^2g^{-1}) ^a	S_{External} (m^2g^{-1}) ^b	$S_{\text{Micropore}}$ (m^2g^{-1}) ^c	$V_{\text{p-Total}}$ (cm^3g^{-1}) ^d	$V_{\text{p-Micropore}}$ (cm^3g^{-1}) ^e	Pore diameter (nm) ^f
2	56	56	0	0.46	0	10.1
5	112	112	0	0.50	0	7.4
12	207	165	42	0.30	0.02	3.5
24	233	141	92	0.24	0.04	3.3

^aBET surface area; ^bExternal surface area; ^cMicropore surface area; ^dTotal pore volume;
^eMicropore volume; ^fPore diameter by BJH desorption.

Scheme S1: Mechanism for the degradation of Chicago Sky Blue 6B via decomposition of H₂O₂ in the presence of Co₃O₄ catalyst.

