## Supporting information

Bimetallic Synergistic Degradation of Chlorophenols by CuCoO<sub>x</sub>–LDH Catalyst in bicarbonate activated hydrogen peroxide system.

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Catalyst	mM of nitrate salt solution used			BET	(Co <sup>2+</sup> +Mg <sup>2+</sup> )/Al <sup>3+</sup>	% Co	рН	% CP
	Co <sup>2+</sup> /other	$Mg^{2+}$	Al <sup>3+</sup>	(m <sup>2</sup> /g)		load		removal
CoO <sub>x</sub> -LDH-1	0.075	15	5	92	3.015	0.60	8.68	72
CoO <sub>x</sub> -LDH -2	0.150	15	5	94	3.030	0.78	8.65	60
CoO <sub>x</sub> -LDH -3	0.225	15	5	94	3.045	1.11	8.66	54
CuO <sub>x</sub> -LDH	1	15	5	84	-	-	8.53	42
MnO <sub>x</sub> -LDH	1	15	5	61	-	-	8.58	25
FeO <sub>x</sub> -LDH	1	15	5	57	-	-	8.49	27
ZnO <sub>x</sub> -LDH	1	15	5	64	-	-	8.51	30
NiO <sub>x</sub> -LDH	1	15	5	81	-	-	8.59	28
MgAl as LDH	-	15	5	82	3.0		8.72	15

**Table S1** Activity of different LDH based metals catalyst on common MgAl support in BAP system.

Conditions: CP 100 ppm, H<sub>2</sub>O<sub>2</sub> 30 mM, NaHCO<sub>3</sub> 30 mM, temperature 40 °C, catalyst amount

1.5 g/L, reaction time 1 h.

## Thermal stability of CuCoOx-LDH-2 catalyst

The DGA analysis as shown in Figure S1 indicates the weight loss on thermal heating. The first mass loss occurred at 250 °C, mainly associated with interlayer and weakly adsorbed water. From temperature 250-500°C, the mass loss is associated with  $CO_3^{2-}$ ,  $NO_3^{-}$  and other intercalated anions chemically bonded with metals layers. The weight loss after 500 °C may be related to traces of  $CO_3^{2-}$  left after calcination.

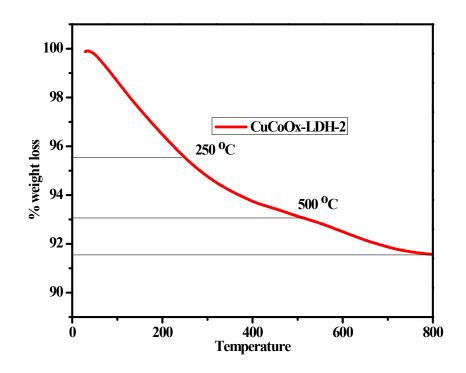
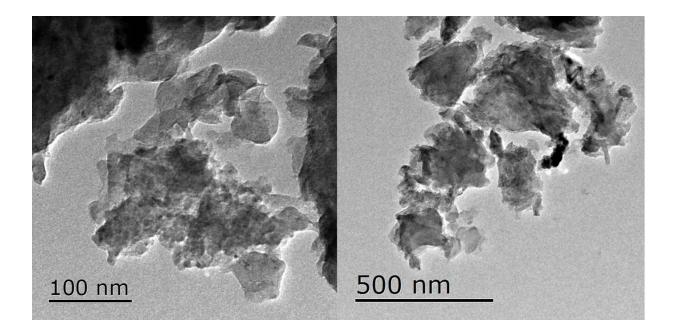


Figure S1 TGA analysis of CuCoO<sub>x</sub>-LDH-2 catalyst.



**Figure S2** TEM images of CuCoO<sub>x</sub>–LDH-2 catalyst.

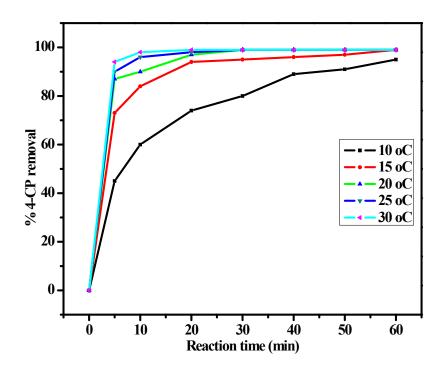
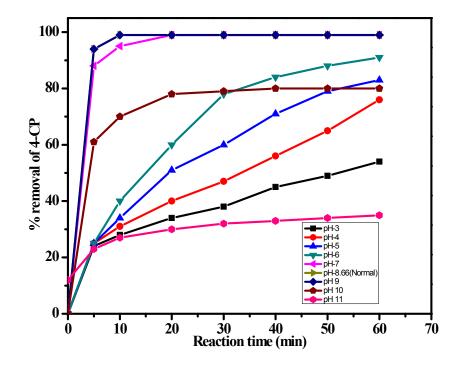


Figure S3 Effect of temperature on the catalytic activity of CuCoO<sub>x</sub>-LDH-2 catalyst.

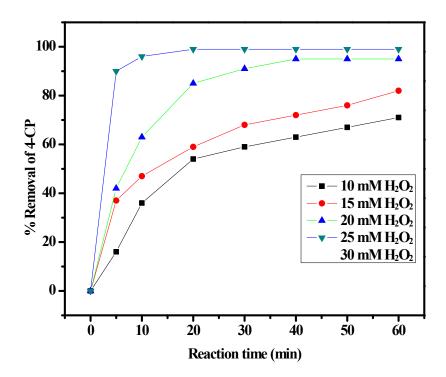
Conditions: 4-CP 100 ppm, H<sub>2</sub>O<sub>2</sub> 25 mM, NaHCO<sub>3</sub> 30 mM, amount of catalyst 0.5 g/L, pH 8.66, temperature 30 °C.

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**Figure S4** Effect of reaction pH on the efficiency of CuCoO<sub>x</sub>-LDH-2 catalyst.

Conditions: CP 100 ppm, H<sub>2</sub>O<sub>2</sub> 25 mM, NaHCO<sub>3</sub> 30 mM, amount of catalyst 0.5 g/L,



**Figure S5** Influence of  $H_2O_2$  on the efficiency of CuCoO<sub>x</sub>-LDH-2 catalyst.

Conditions: CP 100 ppm, NaHCO<sub>3</sub> 30 mM, amount of catalyst 0.5 g/L, temperature 30 °C at pH

 $8.63 \pm 0.3$ .

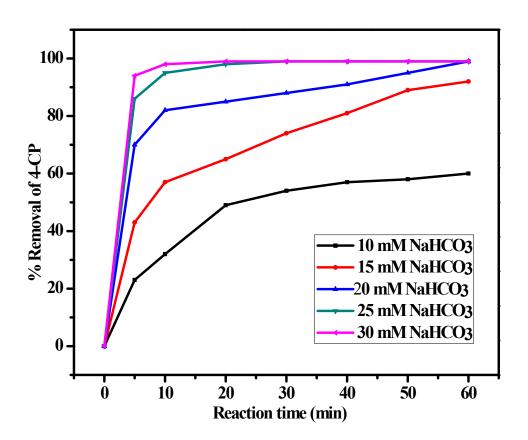
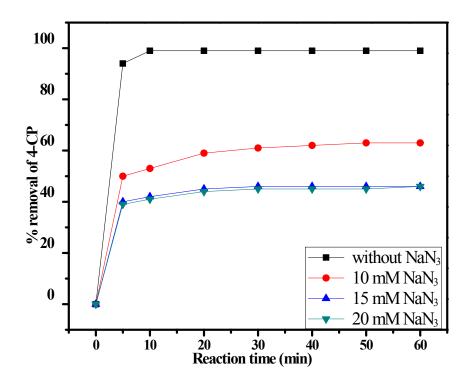


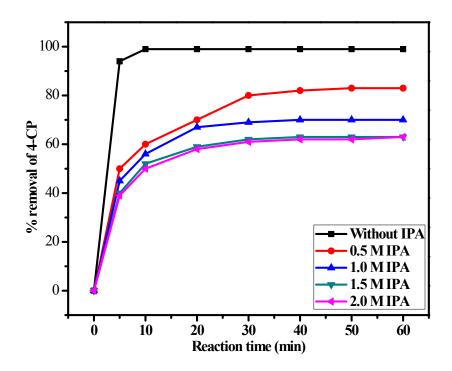
Figure S6 Effect of bicarbonate concentration on the activity of CuCoO<sub>x</sub>-LDH-2 catalyst.

Conditions: CP 100 ppm, H<sub>2</sub>O<sub>2</sub> 25 mM, amount of catalyst 0.5 g/L, temperature 30 °C at pH

 $8.62 \pm 0.4$ .



**Figure S7** Scavenging role of NaN<sub>3</sub> during degradation of 4-CP. Conditions: CP 200 ppm, H<sub>2</sub>O<sub>2</sub> 25 mM, NaHCO<sub>3</sub> 30 mM, amount of catalyst 0.5 g/L,



**Figure S8** Scavenging action of iso propyl alcohol (IPA) during degradation of 4-CP. Conditions: CP 200 ppm, H<sub>2</sub>O<sub>2</sub> 25 mM, NaHCO<sub>3</sub> 30 mM, amount of catalyst 0.5 g/L, temperature 30 °C.

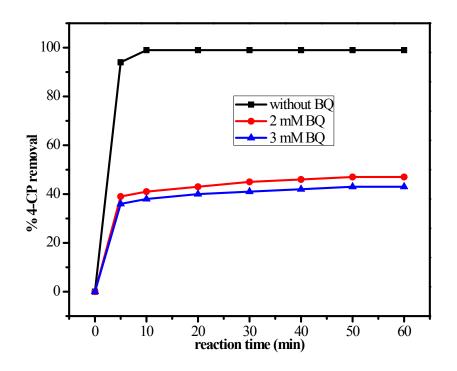
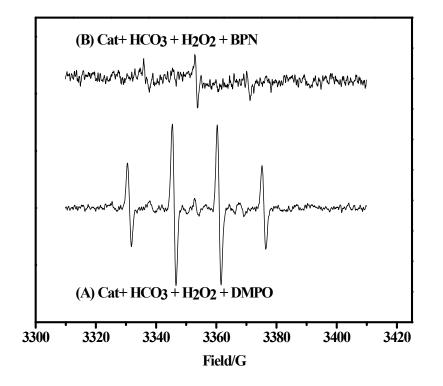


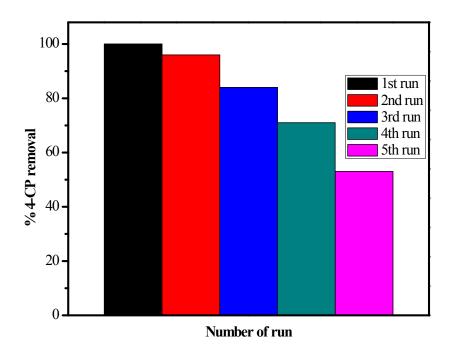
Figure S9 Influence of benzoquinone on the degradation of 4-CP.

Conditions: CP 200 ppm, H<sub>2</sub>O<sub>2</sub> 25 mM, NaHCO<sub>3</sub> 30 mM, amount of catalyst 0.5 g/L,



**Figure S10** Generation of hydroxyl and superoxide radicals with DMPO and BPN Conditions: DMPO 100 mM, BPN 50 mM, NaHCO<sub>3</sub> 30 mM, H<sub>2</sub>O<sub>2</sub> 30 mM, 10 mg/5 mL sample,

10 min reaction time, room temperature.



**Figure S11** Stability of CuO<sub>x</sub>@Co-LDH-2 catalyst in repeated run Conditions: 4-CP 100 ppm, H<sub>2</sub>O<sub>2</sub> 25 mM, NaHCO<sub>3</sub> 30 mM, 0.5 g/L catalyst, time 1 h,

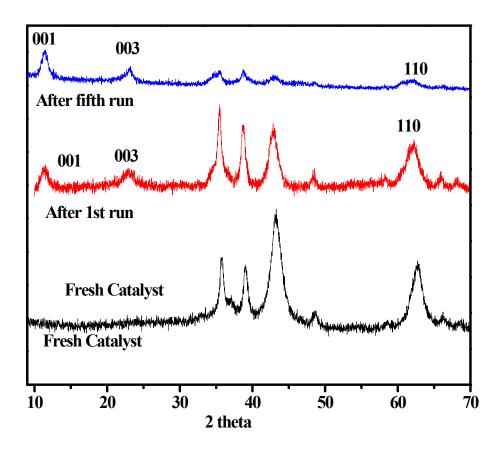


Figure S12 XRD pattern of CuCoO<sub>x</sub>-LDH-2 catalyst during repeated use.