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## **Electronic Supplementary Information**

## Efficient adsorption/photodegradation of organic pollutants from aqueous systems using Cu<sub>2</sub>O nanocrystals as a novel integrated photocatalytic adsorbent

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Fig. S1 XRD patterns of the samples obtained with different molar ratios of Cu and  $CuCl_2$  reactants.



Fig. S2. HRTEM (a) image and EDS (b) spectrum of Cu<sub>2</sub>O NCs.



Fig. S3. Linear fit of experimental data obtained using pseudo-second-order kinetic model at different organic pollutants,  $pH = 5.0\pm0.1$ , m/V = 0.1 g/L.

	Organic pollutants	Experimental values $q_e$ (mg/g)	Calculated $q_e$ (mg/g)	$k_2$ (g/(mg·min))	<i>R</i> <sup>2</sup>
	НА	345.22	344.83	0.00350	0.9997
Cu <sub>2</sub> O NCs	CR	84.88	90.91	0.00931	0.9998
	MO	87.53	87.72	0.00241	0.9999
CM Cu <sub>2</sub> O	HA	26.37	26.39	0.0898	0.9999
	CR	27.61	27.78	0.01489	0.9994
	MO	4.97	4.98	1.13055	0.9996

**Table S1.** The experimental and calculated  $q_e$  values, pseudo-second-order rate constants,  $k_2$ , and correlation coefficient values,  $R_2$ .



**Fig. S4.** Digital pictures of HA, CR and MO adsorption on as-prepared Cu<sub>2</sub>O NCs sample. NO. 1 vials are corresponded to HA, CR and MO solution, respectively. NO. 2 and 3 vials are taken at adsorption time of 10 minutes and 30 minutes, respectively.



Fig. S5. (a) Effect of solution pH on the adsorption of HA by  $Cu_2O$  NCs and (b) Zetapotential of  $Cu_2O$  NCs as a function of pH.



Fig. S6. Adsorption isotherms of CR, MO and HA on  $Cu_2O$  NCs (a) and CM  $Cu_2O$  (b).

Samples	Organic	Langmuir			Freundlich		
	pollutants	$q_{ m max} \ ( m mg/g)$	<i>b</i> ( L/mg )	$R^2$	k	п	<i>R</i> <sup>2</sup>
Cu <sub>2</sub> O NCs	НА	405.5	0.935	0.992	199.86	6.449	0.814
	CR	401.4	0.203	0.996	116.12	3.883	0.878
	МО	382.9	0.198	0.995	113.15	3.926	0.921
CM Cu <sub>2</sub> O	HA	212.3	0.0576	0.983	45.62	3.521	0.833
	CR	52.9	0.164	0.963	20.23	5.250	0.911
	МО	22.4	0.044	0.934	3.59	2.955	0.798

**Table S2.** Parameters of the Langmuir and Freundlich isotherm model for organic pollutants on Cu<sub>2</sub>O NCs and CM Cu<sub>2</sub>O

**Table S3.** Comparison of maximum adsorption capacity of organic pollutantsadsorption on  $Cu_2O$  NCs with other different adsorbents.

Adsorbents	Adsorption capacity (mg $\cdot$ g $^{-1}$ )	Ref.
	НА	
Activated carbon	2.15	1
Fly ash	72	2
Coal fly	16.6	3
Mesoporous silica	120	4
Aminopropyl functionalized SBA-15	117	5
Cu <sub>2</sub> O NCs	405.5	This study
	CR	
Activated carbon	52-189	6
waste Fe(III)/Cr(III) hydroxide	44.00	7
waste Orange peel	22.44	8
waste banana pith	20.29	9
Cu <sub>2</sub> O NCs	401.4	This study
	ΜΟ	
Activated carbon	9.94	10
Orange peels	20.50	11
Hyper-cross-linked polymer	70.92	12
Cu <sub>2</sub> O NCs	382.9	This study



Fig. S7 XRD patterns (a) and Cu  $2p_{3/2}$  XPS spectra (b) of CM Cu<sub>2</sub>O and Cu<sub>2</sub>O NCs.



Fig. S8. Photodegradation of CR (a) and MO (c) over Cu<sub>2</sub>O NCs and CM Cu<sub>2</sub>O under visible light irradiation, respectively. UV–Vis spectra of CR (b) and MO (d) in aqueous Cu<sub>2</sub>O NCs dispersions as a function of irradiation time with visible light irradiation ( $\lambda \ge 400$  nm).



Fig. S9. Photocatalytic degradation reaction kinetics of HA, CR and MO over Cu<sub>2</sub>O

NCs, respectively.



Fig. S10. Photodegradation (a) and photocatalytic degradation reaction kinetics (b) of TC over  $Cu_2O$  NCs under visible light irradiation.



Fig. S11 The TOC removal curve of HA photocatalytic degradation by Cu<sub>2</sub>O NCs.



Fig. S12 The UV–Vis diffuse reflectance spectra of  $Cu_2O$  NCs and after HA adsorption.



**Fig. S13.** (a) Digital pictures of the natural groundwater well; (b) the color of the groundwater sample; (c) the characterization of NOM by excitation emission matrix fluorescence spectroscopy and (d) by high performance size exclusion chromatography.

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