

Supporting information

For

Rapid gelling of guar gum hydrogel stabilized by copper hydroxide nanoclusters for efficient removal of heavy metal and supercapacitors

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Supplementary materials

Methods

Synthesis of Cu/GG gel: 0.1208g $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ was added into purified GG solution (0.5 wt%, 50 ml), after stirring for 30 min ensures the complete dissolution of salt, the mixture was poured into KOH solution (1 M, 50 ml), the as-obtained gel was named as Cu/GG gelation.

Synthesis of $\text{Cu}(\text{OH})_2$ /GG composite: Typically, 0.1208g $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ was first weighed out and added into KOH solution (1 M, 50 ml), after standing for 30 min with stirring at temperature, the mixture was poured into purified GG solution (0.5 wt%, 50ml), and a gel consisted of $\text{Cu}(\text{OH})_2$ and GG formed immediately.

Synthesis of CuO/GG composite: 0.1208g $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ was added into KOH solution (1 M, 50 ml), then the mixture was transferred into a autoclave with Teflon lining, then it was treated at 100 °C for 12 h, After that, the solution was poured into purified GG solution (0.5 wt%, 50 ml), the final mixture was labelled as CuO/GG composite.



Figure s1 the stability testing of spherical hydrogel after standing for 1 year, the concentration of metal ions is 100 mM.

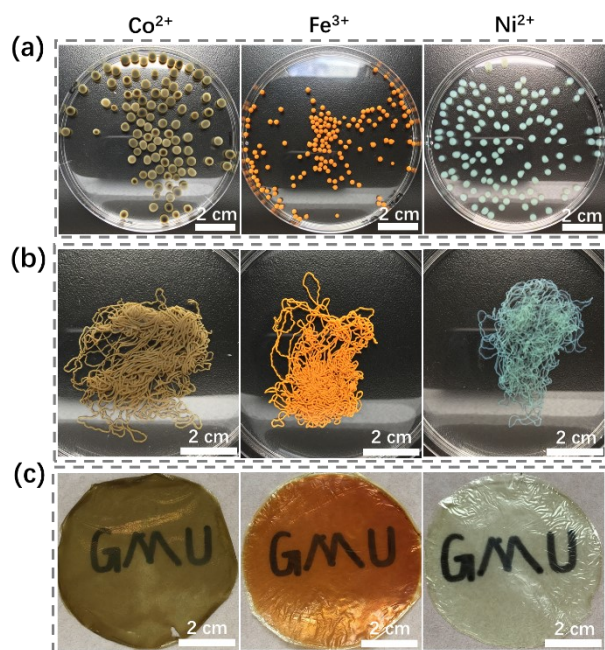


Figure s2 Co/GG, Fe/GG and Ni/GG hydrogel beads (a); fiber (b) and membrane (c)

Table s1 elemental distribution of hydrogel sphere evaluated by EDS

	Elements (wt%)	C	K	Cu	O
10s	Surface	26.56	24.04	13.53	35.87
	Cross-section	31.06	16.27	14.94	37.73
24h	Surface	36.13	2.80	12.54	32.75
	Cross-section	35.80	1.92	13.93	34.09

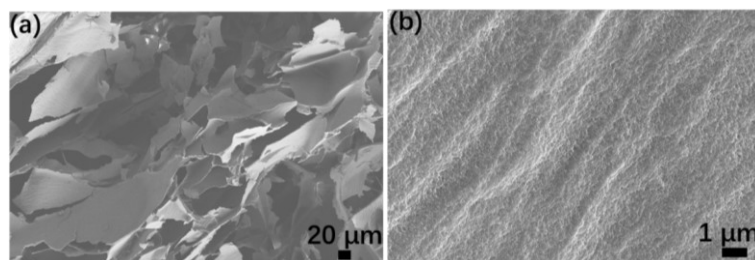


Figure s3 SEM images of the cross-section of Cu/GG sphere after standing for 24h

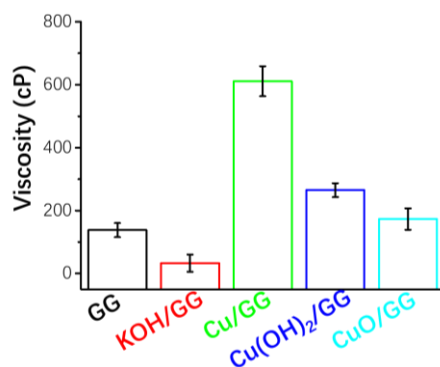


Figure s4 the viscosity transformation affected by different additives at a shear rate of 200 1/s. The concentration of these additives is 10 mM.

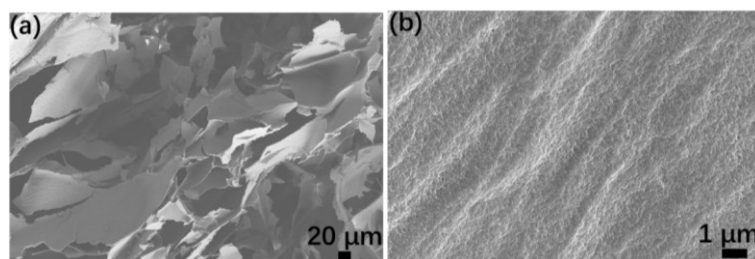


Figure s5 SEM images of the cross-section of Cu/GG sphere after standing for 24h

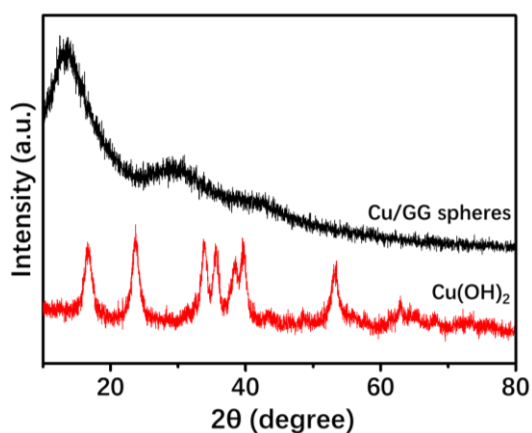


Figure s6 XRD patterns of Cu(OH)₂ and Cu/GG spheres. Cu(OH)₂ was prepared as the following procedure: 0.1208g Cu(NO₃)₂·3H₂O was added into KOH solution (1 M, 50 ml), after reacted for 2 min, the particles were obtained by centrifugation, then, the sample was washed with de-ionized water and freeze-dried.

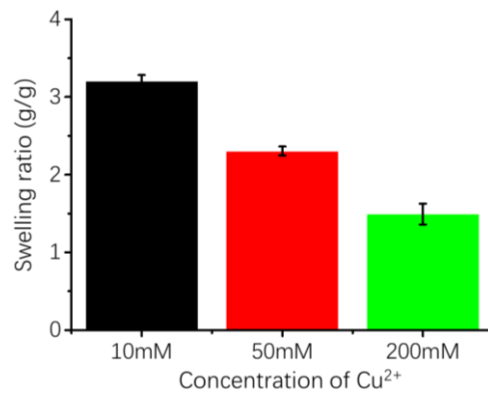


Figure s7 effect of the concentration of Cu^{2+} on the swelling ratio of Cu/GG hydrogel



Figure s8 the encapsulation capacity and stability testing of Cu/GG hydrogel. The mole ratio of pollutant/ Cu^{2+} is 200.

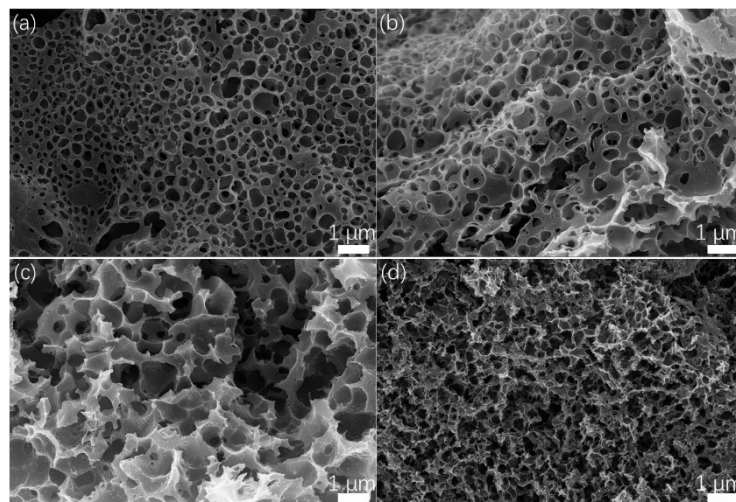


Figure 9 SEM images of HPCS samples prepared under different carbonization temperature

Table s2 pore structure analysis of the samples

Samples	S_{BET} (m^2g^{-1})	$S_{\text{t-plot}}$ (m^2g^{-1})	$S_{\text{t-plot}}/S_{\text{BET}}$	$V_{\text{t-plot}}$ (m^3g^{-1})
HPCS-400	440	334	0.75	0.17
HPCS-600	690	611	0.88	0.32
HPCS-800	1280	930	0.79	0.48
HPCS-900	1524	660	0.44	0.33

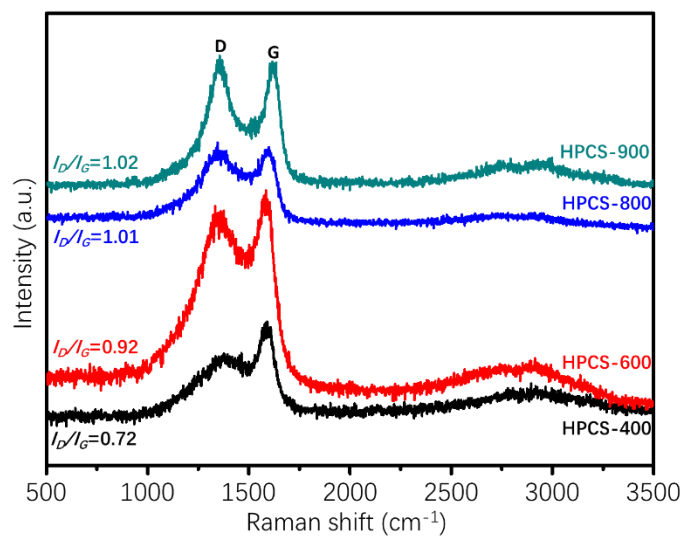


Figure 10 Raman spectra of the samples prepared at different carbonization temperature

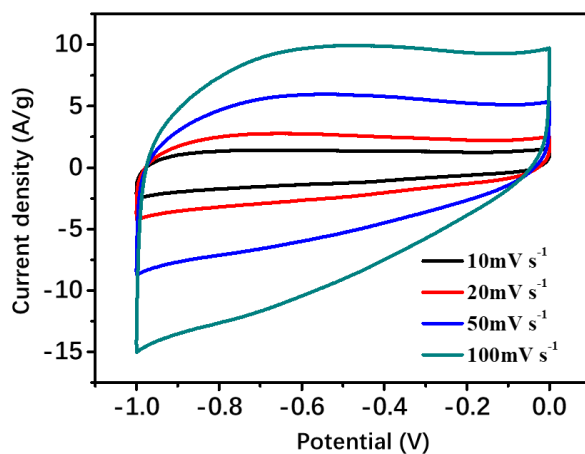


Figure s11 CV curves of HPCS-900 at various scanning rates

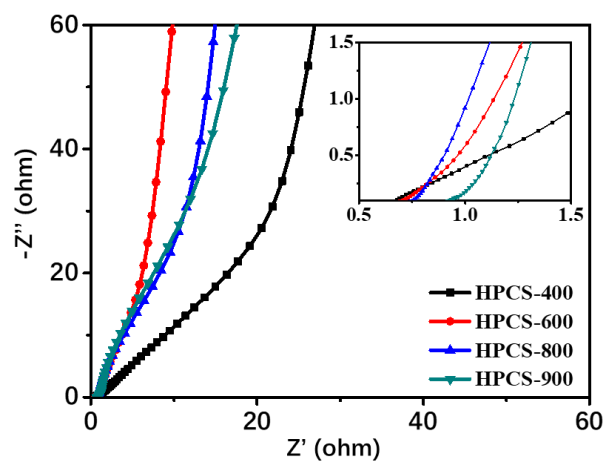


Figure s12 Nyquist plots of EIS of HPCS-900 (three-electrode system)

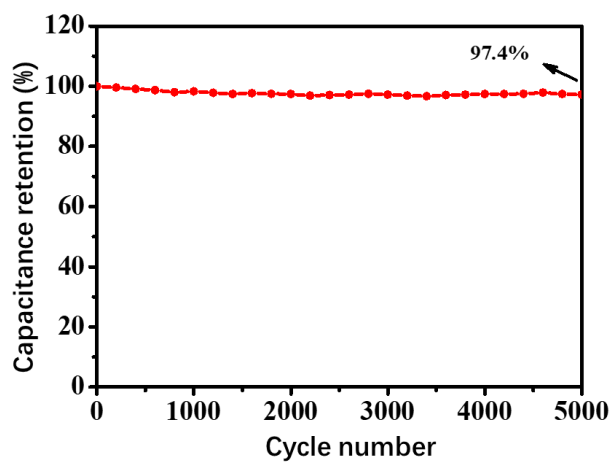


Fig. s13 cycling performance of HPCS-900 at a current density of 10 A/g