

Supporting Information

Recovery of uranium from aqueous solutions by the modified honeycomb-like porous carbon material

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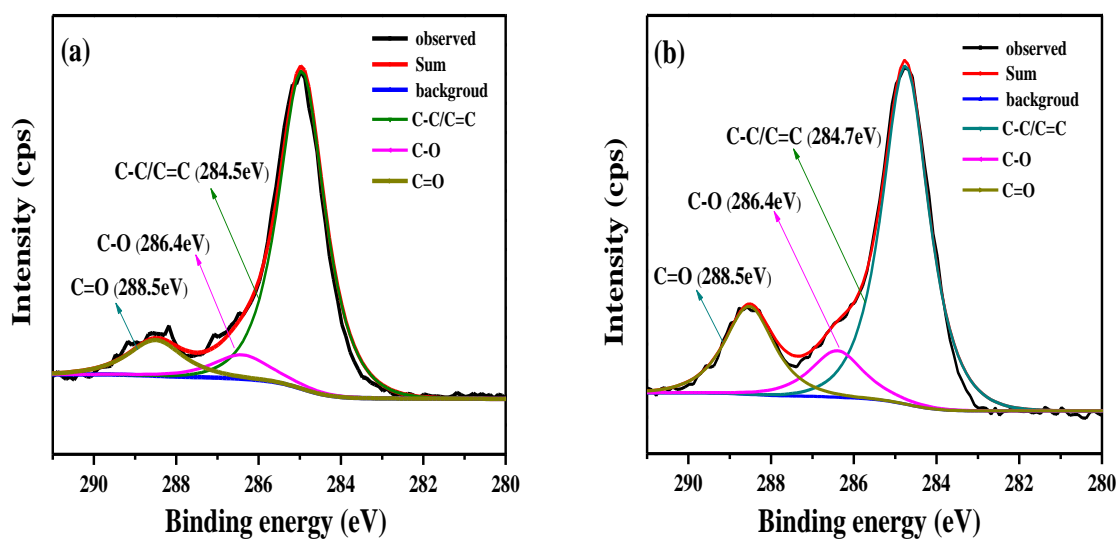


Fig. S1. The C1s spectra of HLPC/MnO₂ (a) and HLPC/MnO₂/U.

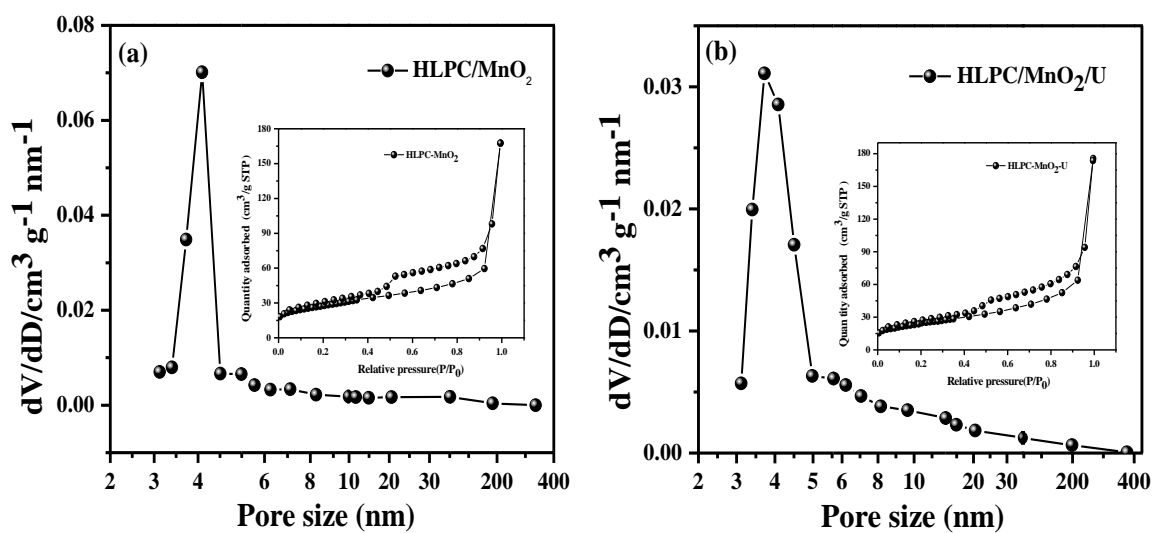


Fig. S2 (a) and (b) the pore size distributions of HLPC/MnO₂ and HLPC/MnO₂/U.

The inset of (a) and (b) is N₂ adsorption/desorption isotherm of HLPC/MnO₂ and HLPC/MnO₂/U.

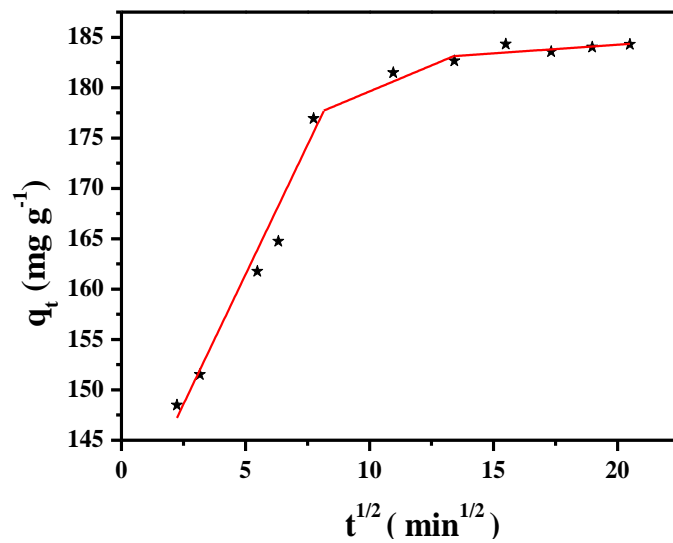


Fig. S3 Intra-particle diffusion kinetics for the adsorption of uranyl ions on HLPC/MnO₂

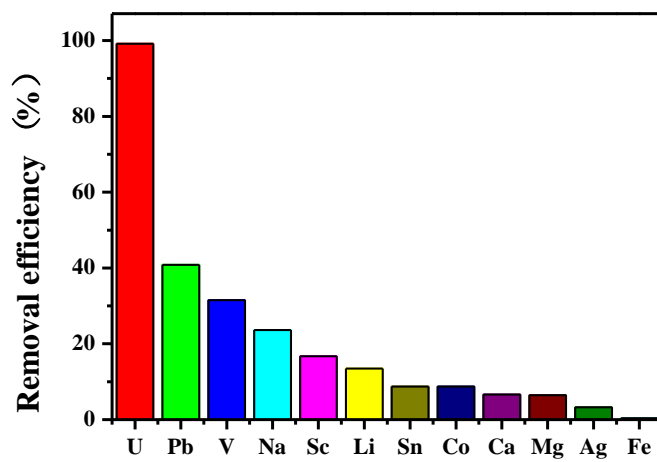


Fig. S4 Selected results of HLPC/MnO₂ for the extraction of uranyl ions from simulating seawater.

Table S1 Elemental composition of the HLPC/MnO₂ and HLPC/MnO₂/U samples determined by XPS spectra.

Sample	XPS				Binding energy			
	C	O	Mn	U	C	O	Mn	U
	at%	at%	at%	at%	ev	ev	ev	ev
HLPC/MnO ₂	47.51	38.02	14.47	0	278.08	523.58	634.08	—
HLPC/MnO ₂ /U	32.92	49.58	16.47	1.03	284.74	530.22	642.11	381.85

Table S2 Parameters of the intra-particle diffusion model of uranium (VI) adsorption on HLPC/MnO₂

T(K)	k _{p1}	R ₁ ²	k _{p2}	R ₂ ²	k _{p3}	R ₃ ²
298	5.148	0.975	1.032	0.876	0.173	0.316

Table. S3 Isotherm constants and values of R² for HLPC/MnO₂.

T(K)	Langmuir isotherm			Freundlich isotherm		
	q _m (mg • g ⁻¹)	b (L mg ⁻¹)	R ²	K (L g ⁻¹)	n	R ²
298	238.09	0.3258	0.998	69.01	0.3248	0.896
308	262.47	0.2679	0.994	73.34	0.3320	0.667
318	271.74	0.6513	0.998	99.26	0.3024	0.919

Table. S4 The maximum adsorption capacity of different adsorbents for uranium (VI).

Adsorbents	Adsorption Capacity mg-U/g-adsorbent	Conditions	Ref.
MnO ₂ -coating of the cactus fibres	62	$C_{\text{initial}} = 9 \times 10^{-6} - 9 \times 10^{-4} \text{ mol L}^{-1}$, m = 10mg, V = 15 ml, T = 298 K, pH=6	[1]
Salicylideneimine-functionalized hydrothermal carbon	261	$C_{\text{initial}} = 0.2 - 1.3 \text{ mmol L}^{-1}$, m = 10 mg, V = 20 ml T = 288.15 K, pH=4.3	[2]
CaCl ₂ -modified Giant Kelp biomass	156.25	$C_{\text{initial}} = 20 - 100 \text{ mg L}^{-1}$, m = 50 mg, V = 100 ml, T = 293 K, pH =5	[3]
imine-functionalized carbon spheres	113.16	$C_{\text{initial}} = 1 - 100 \text{ mg L}^{-1}$, m = 50mg, V = 20 ml, T = 298K, pH=4	[4]
starch/SnO ₂	192	$C_{\text{initial}} = 10 - 150 \text{ mg L}^{-1}$, m = 100 mg, V = 250 ml T = 298 K, pH = 6	[5]
ion-imprinted magnetic chitosan resins	187.26	$C_{\text{initial}} = 15 - 420 \text{ mg L}^{-1}$, m = 50 mg, V = 50 ml T = 298 K, pH = 5	[6]
Graphene oxide-manganese dioxide	185.2	$C_{\text{initial}} = 22.5 - 70 \text{ mg L}^{-1}$, m = 10 mg, V = 20 ml T = 298 K, pH = 3.8	[7]
HLPC-MnO ₂	238.09	$C_{\text{initial}} = 30 - 300 \text{ mg L}^{-1}$, m = 10 mg, V = 20 ml T = 298 K, pH = 5	this work

Table S5. Selected results for the extraction of uranyl ions from simulated seawater

Elements	Ion concentration (ug L ⁻¹)		Removal (%)
	initial	final	
U	3.131	0.026	99.16
Pb	0.424	0.251	40.8
V	0.489	0.335	31.49
Na	1.521×10 ⁴	1.162×10 ⁴	23.60
Sc	249.1	207.4	16.74
Li	17.61	15.24	13.46
Sn	12.35	11.27	8.74
Co	0.63	0.575	8.73
Mg	6.289×10 ³	5.104×10 ³	6.44
Ca	9.169×10 ³	8.563×10 ³	6.61
Ag	0.521	0.504	3.26
Fe	104.3	103.9	0.38

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