

Electronic Supporting Information (ESI)

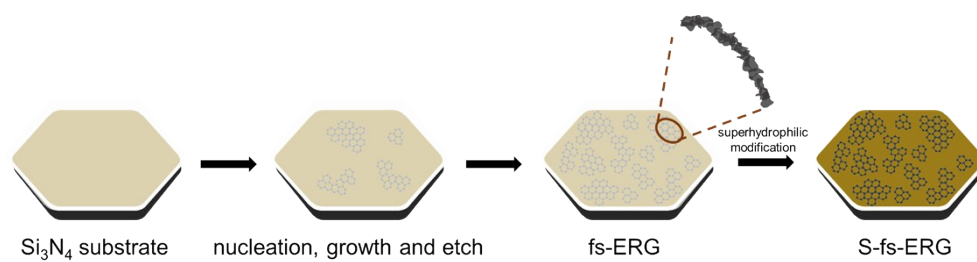
Superhydrophilic edge-rich graphene for the simultaneous and disposable sensing of dopamine, ascorbic acid, and uric acid

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Scheme S1. Schematic diagram for preparation of the S-fs-ERG.

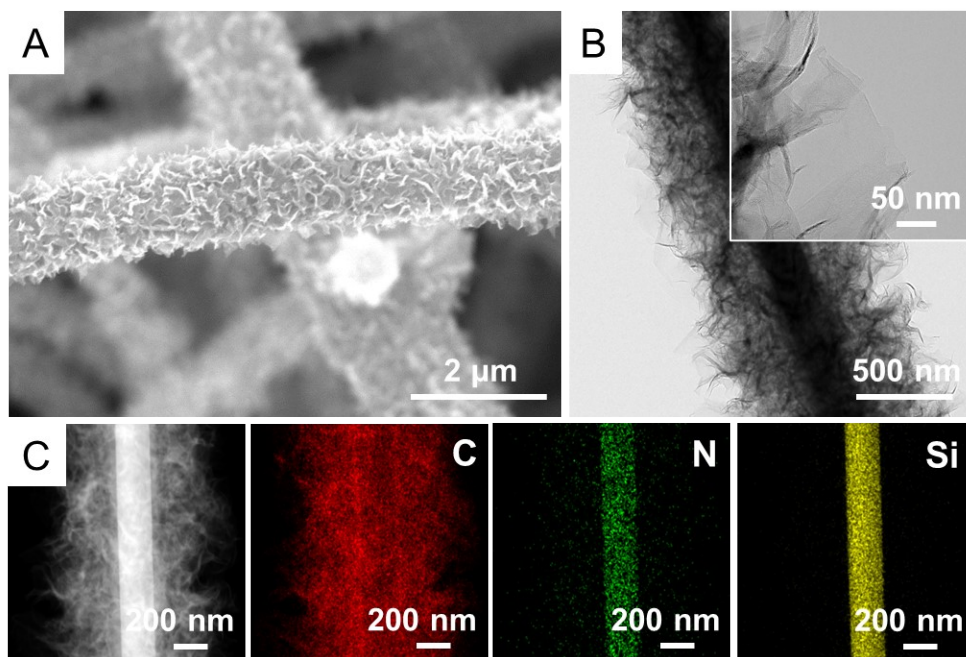


Fig. S1 SEM (A), TEM (B) and corresponding TEM-EDS (C) images of the fs-ERG.

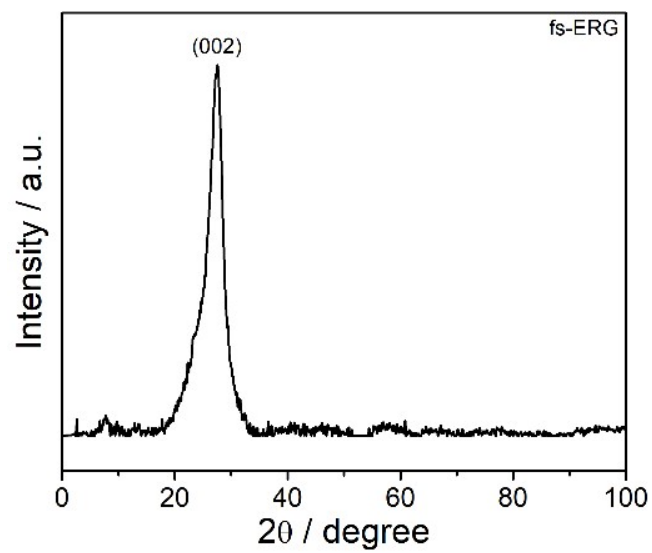


Fig. S2 XRD pattern of the fs-ERG.

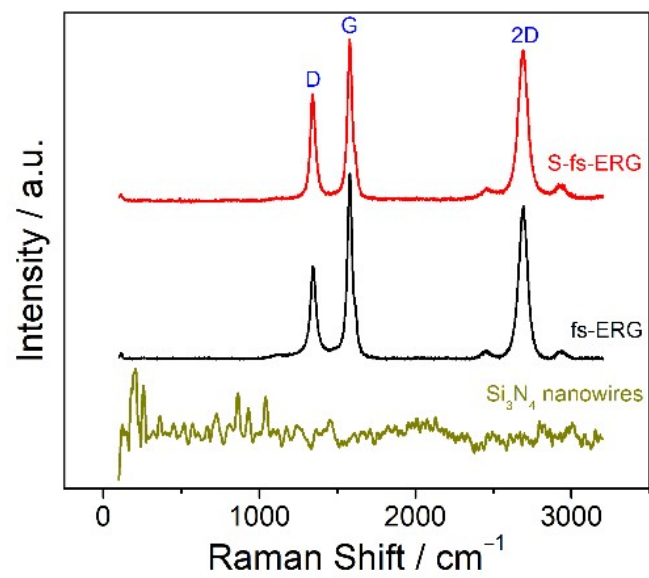


Fig. S3 Raman spectra of the Si₃N₄ nanowires, fs-ERG, and S-fs-ERG.

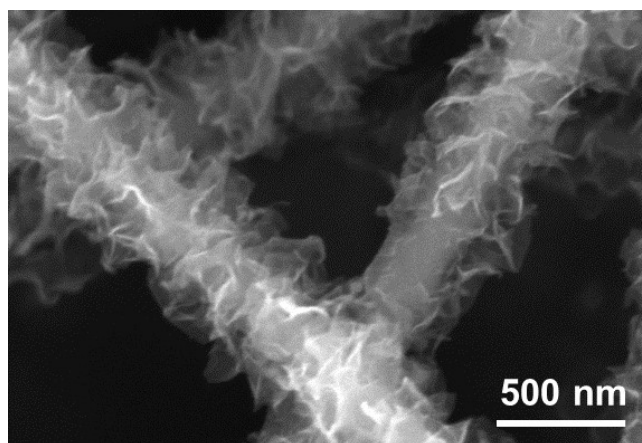


Fig. S4 SEM image of the S-fs-ERG. A small number of fs-ERG nanoplanes were falling-off from the Si₃N₄ nanowires after KOH thermal treatment.

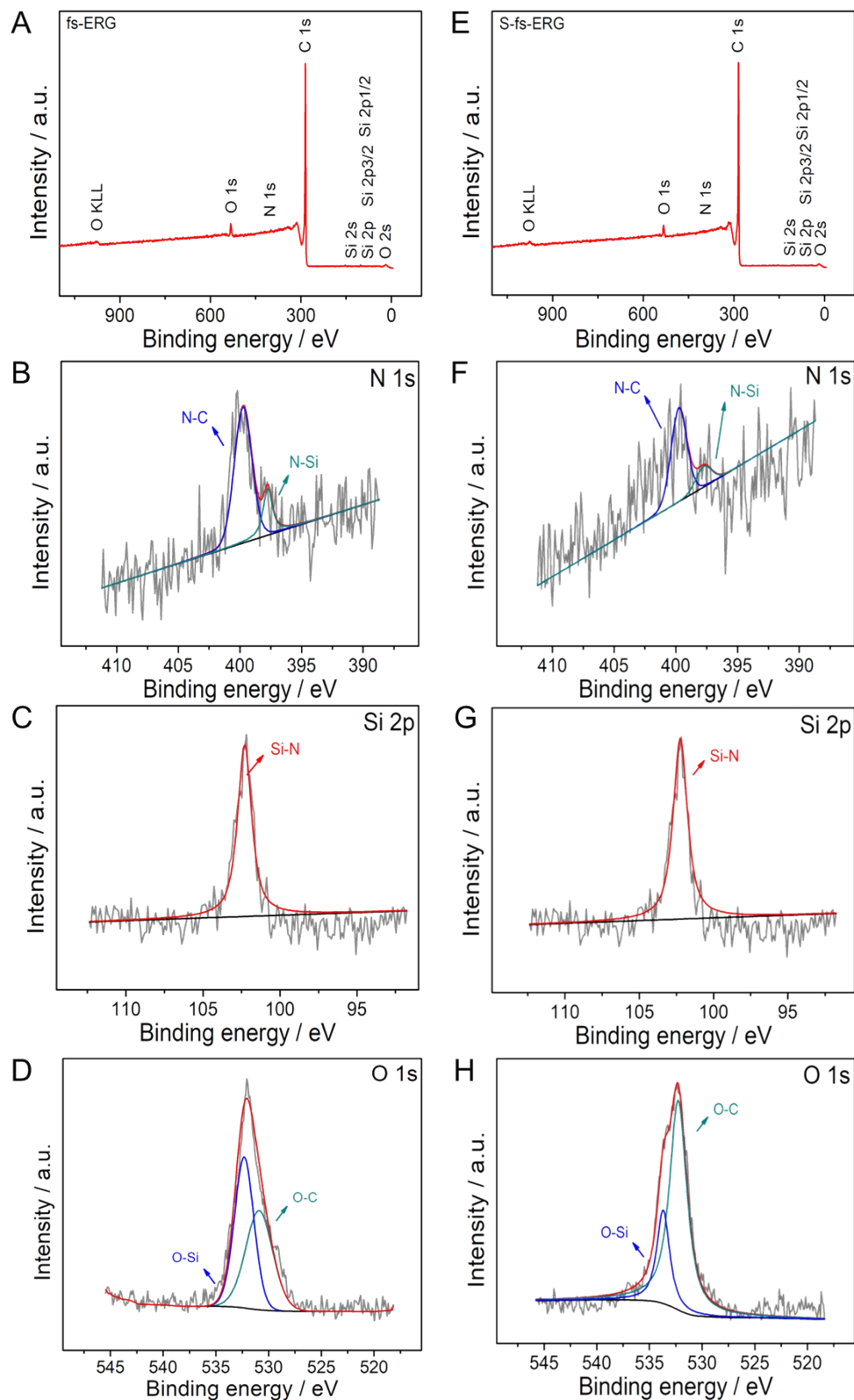


Fig. S5 XPS wide scan (A, E) and high-resolution N 1s (B, F), Si 2p (C, G), and O 1s (D, H) spectra of the fs-ERG (A, B, C, D) and S-fs-ERG (E, F, G, H), respectively.

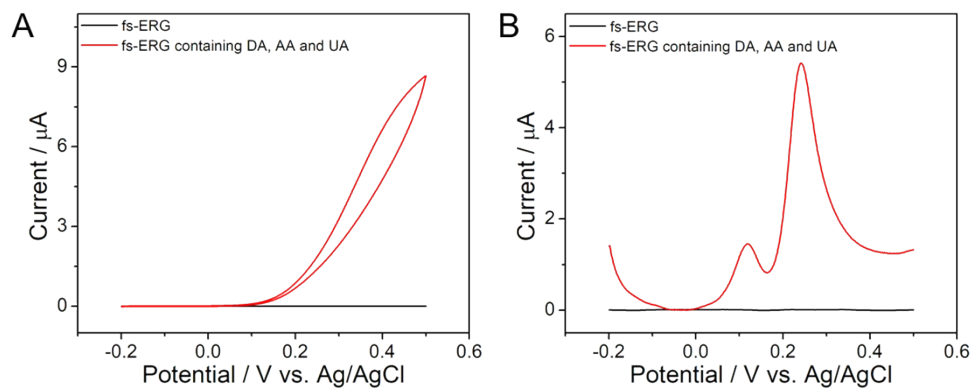


Fig. S6 CV (A) and DPV (B) curves of the fs-ERG electrode in the absence and presence of 0.02 mM DA, 1 mM AA and 0.05 mM UA in 0.1 M PBS (pH 7.4). The scan rate for CV is 50 mV s^{-1} . The pulse amplitude for DPV is 50 mV.

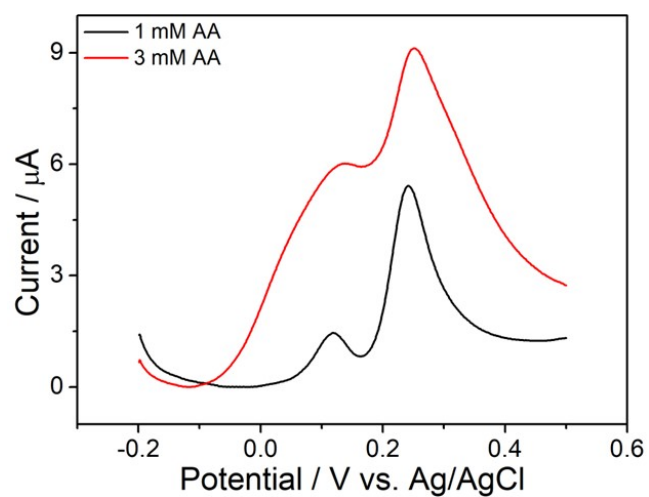


Fig. S7 DPV curves of the fs-ERG electrode containing 0.02 mM DA and 0.05 mM UA with different concentrations of AA in 0.1 M PBS (pH 7.4).

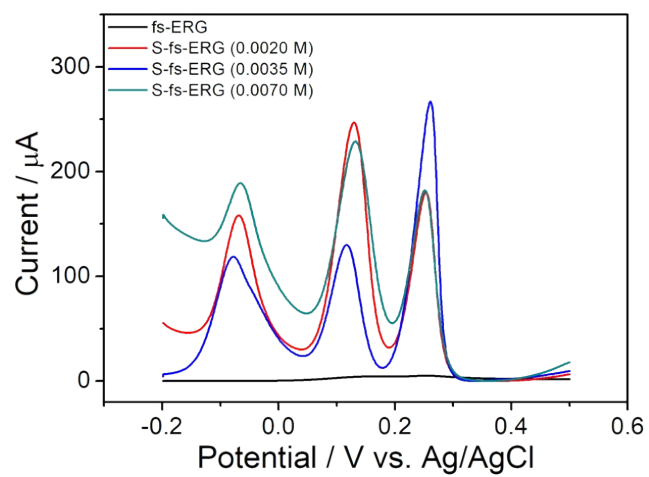


Fig. S8 DPV curves of the fs-ERG and S-fs-ERG electrodes after different concentrations of KOH thermal treatment containing 0.02 mM DA, 1 mM AA and 0.05 mM UA in 0.1 M PBS (pH 7.4).

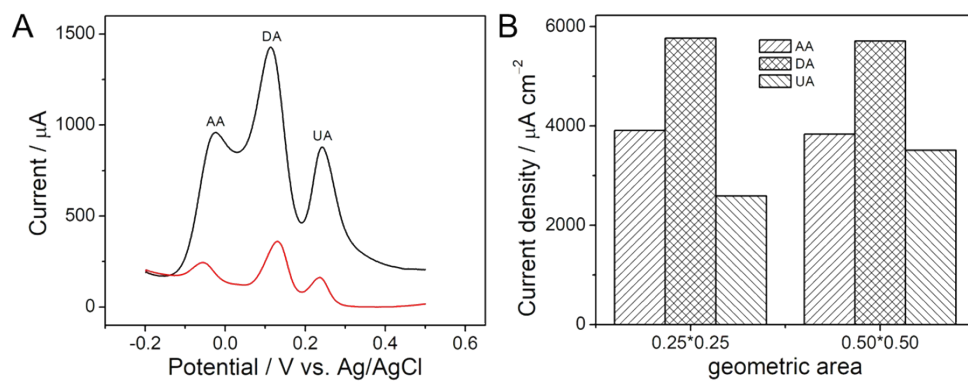


Fig. S9 DPV curves of the S-fs-ERG electrodes with different dimensions of 0.25×0.25 cm (red line) and 0.50×0.50 cm (black line) (A) and the corresponding histograms of the current density responses (B) containing 0.02 mM DA, 1 mM AA and 0.05 mM UA in 0.1 M PBS (pH 7.4).

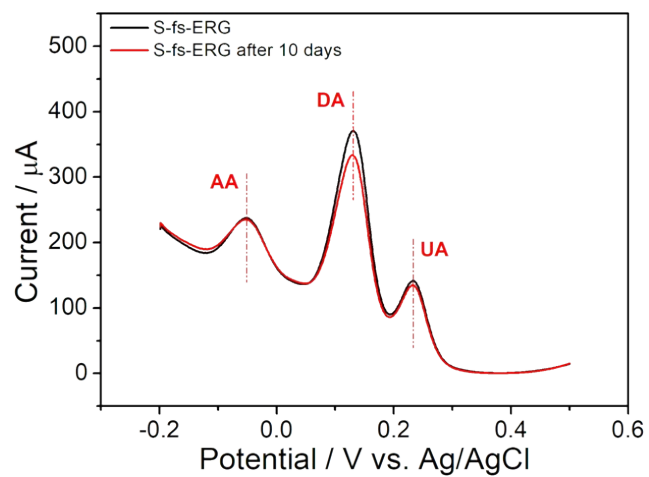


Fig. S10 DPV curves of the S-fs-ERG electrode containing 0.02 mM DA, 1 mM AA and 0.05 mM UA for stability test.

Table S1 The TEM-EDS element analysis of the fs-ERG (A) and S-fs-ERG (B).

(A)	Element (Family)	Atomic Fraction (%)	Mass Fraction (%)	(B)	Element (Family)	Atomic Fraction (%)	Mass Fraction (%)
	C (K)	92.97	85.67		C (K)	96.00	91.80
	N (K)	0.00	0.00		N (K)	0.00	0.00
	Si (K)	7.03	14.33		Si (K)	3.68	7.78
					O (K)	0.32	0.41
	Total	100.00	100.00		Total	100.00	100.00

Table S2 The atomic content changes about C, N, Si and O elements of the fs-ERG and S-fs-ERG.

Atomic contents (%)	C	N	Si	O
fs-ERG	92.46	0.92	3.59	3.04
S-fs-ERG	94.19	0.82	1.85	3.14

Table S3 Comparison of the differently modified electrodes for electrochemical determination of the DA, AA and UA.

Electrode	Linear range / μM			Limit of detection (LOD)			Refs
	DA	AA	UA	DA	AA	UA	
2D-hBN ^a	3–75	N.A.	N.A.	0.65	N.A.	N.A.	1
NOCC ^b	0.3–55	10–1300	N.A.	0.18	3.41	N.A.	2
Pt@NP-AuSn/Ni/CFP ^c	1–10	200–2000	25–800	N.A.	N.A.	N.A.	3
BG-CNPs ^d	5–2535	10–3570	5–220	0.7	1.1	2.4	4
S-fs-ERG	0.1–50	2.5–1000	0.5–50	0.1	2.5	0.5	This work

Note: The low detectable limit as evaluation index in this work.

N.A.: not available.

^a: 2D hexagonal boron nitride

^b: nitrogen/oxygen-codoped carbon cloth

^c: Pt nanoparticles modified nanoporous AuSn(Pt@NP-AuSn) alloy on Ni buffered flexible carbon fiber paper (CFP)

^d: biomass derived ground cherry husks derived carbon nanoplates

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