Supporting Information

Fe–N-doped carbon foam nanosheets with embedded Fe_2O_3 nanoparticles for highly efficient oxygen reduction in both alkaline and acidic media

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Electrochemical measurements

The kinetic parameters can be analyzed with the Koutecky–Levich equation:

$$J^{-1} = J_{\rm K}^{-1} + J_{\rm L}^{-1} = (nFkC)^{-1} + (0.62nFCD^{2/3}v^{-1/6}\omega^{1/2})^{-1}$$

wherein *J*, $J_{\rm K}$ and $J_{\rm L}$ represent the measured current density, the kinetic current density, and diffusion limiting current density, respectively, ω is rotation rate of the electrode, *n* is the electron transfer number, *F* is the Faraday constant, *C* is the bulk concentration of O₂ dissolved in the electrolyte (1.2×10⁻³mol L⁻¹), *D* is the diffusion coefficient of O₂ in the electrolyte (1.9×10⁻⁵ cm⁻² s⁻¹), *v* is the kinematic viscosity of the electrolyte (1.0×10⁻² cm⁻² s⁻¹), and *k* is the electron transfer rate constant.¹ The values of *C*, *D* and *v* are the same in both 0.1 M KOH and 0.5 M H₂SO₄ solution.²

References:

- 1. R. E. Davis, G. L. Horvath, C. W. Tobias, *Electrochim. Acta*, 1967, 12, 287-297.
- J. H. Xue, L. Zhao, Z. Y. Dou, Y. Yang, Y. Yue, Z. Zhu, Rsc Adv, 2016, 6,110820-110830

Table S1. BET surface areas and total pore volumes of Fe₂O₃@Fe-N-C-800-BM and Fe₂O₃@Fe-

Sample	T(°C) ^a	$S_{BET} (m^2 g^{-1})^b$	$V_{total} (cm^3 g^{-1})^c$
Fe ₂ O ₃ @Fe-N-C-700	700	497	0.58
Fe ₂ O ₃ @Fe-N-C-800	800	646	0.85
Fe ₂ O ₃ @Fe-N-C-900	900	527	0.73

N-C obtained at different carbonization temperatures

^a Carbonization temperature.

^b BET specific surface areas obtained from N₂ adsorption isotherm in the range of $P/P_0 = 0.05-0.3$.

^c Total pore volume was obtained at P/P_0 of 0.98.

Fable S2. XPS data for the surface	e species of Fe ₂ O ₃ @Fe-N-C	-800-BM and Fe ₂ O ₃ @Fe-N-C-T
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Sample	Ν	Fe	pyridinic-N	graphitic-N	oxidized-N
	(at%)	(at%)	(%)	(%)	(%)
Fe ₂ O ₃ @Fe-N-C-700	17.8	1.9	53	41	6
Fe ₂ O ₃ @Fe-N-C-800	9.9	2.3	46	47	7
Fe ₂ O ₃ @Fe-N-C-900	4.8	0.7	41	49	10
Fe ₂ O ₃ @Fe-N-C-800-BM	10.6	0.2	43	50	7

materials obtained at different temperatures and their content nitrogen species

Samples	Onset-potential	Half-wave	J ^a	$J_{ m K}{}^{ m b}$
	V (vs. Ag/AgCl)	potential	$(mA cm^{-2})$	(mA cm ⁻²)
Fe ₂ O ₃ @Fe-N-C-700	0.001	-0.278	0.880	9.09
Fe ₂ O ₃ @Fe-N-C-800	0.054	-0.104	3.617	11.73
Fe ₂ O ₃ @Fe-N-C-900	0.052	-0.115	3.406	10.85
Fe ₂ O ₃ @Fe-N-C-800-BM	0.021	-0.270	1.765	5.11
Pt/C	0.056	-0.136	2.565	9.57

Table S3. The data of catalytic activity for Fe₂O₃@Fe-N-C-T in 0.1 M KOH solution

^a The experimental current density (J) at -0.15 V determined at the polarization curve at 1600rpm in 0.1M

KOH solution

^b The kinetic current densities ($J_{\rm K}$) at -0.10 V determined at the polarization curve at 1600rpm in 0.1M KOH

solution

Samples	Onset-potential	Half-wave	Ja	$J_{\mathrm{K}}^{\mathrm{b}}$
	V (vs. Ag/AgCl)	potential	$(mA cm^{-2})$	(mA cm ⁻²)
Fe ₂ O ₃ @Fe-N-C-700	0.637	0.380	2.805	7.04
Fe ₂ O ₃ @Fe-N-C-800	0.698	0.535	4.656	10.47
Fe ₂ O ₃ @Fe-N-C-900	0.640	0.499	4.405	9.16
Fe ₂ O ₃ @Fe-N-C-800-BM	0.680	0.494	3.523	6.19
Pt/C	0.702	0.569	4.439	12.25

Table S4. The data of catalytic activity for Fe₂O₃@Fe-N-C-Tin 0.5 M H₂SO₄ solution

^a The experimental current density (*J*) at 0.35V determined at the polarization curve at 1600rpm in 0.5M

H_2SO_4 solution

^b The kinetic current densities ($J_{\rm K}$) at 0.60V determined at the polarization curve at 1600rpm in 0.5M H₂SO₄

solution



Fig. S1 (a, b) TEM images of Fe₂O₃@Fe-N-C-700; (c, d) TEM images of Fe₂O₃@Fe-N-C-900.



Fig. S2 (a) STEM image of Fe_2O_3 @Fe-N-C-800, and elemental mapping images (recorded in region 1) of (b) Fe, (c) N, and (d) C.



Fig. S3 (a) Wide XPS survey of theFe₂O₃@Fe-N-C-800. High-resolution (b) C1s, (c) N1s, and (d) Fe 2p spectra of the Fe₂O₃@Fe-N-C-800.



Fig. S4 (a)Nitrogen adsorption-desorption isotherms of Fe_2O_3 @Fe-N-C-T samples prepared at different carbonization temperature of 700,800 and 900°C,respectively; (b) the corresponding pore size distribution curves.



Fig. S5 LSVs of Fe₂O₃@Fe-N-C-800 with different catalyst loadings(100, 200 and 410 μ g cm⁻²) at a scan rate of 5 mV s⁻¹ and a rotation rate of 1600 rpm in O₂-saturated (a) 0.1 M KOH and (b) 0.5 M H₂SO₄, respectively.



Fig. S6 LSV curves at rotation rate from 900 to 2500 rpm and the corresponding K-L plots (inset) of Fe₂O₃@Fe-N-C-800 with the catalyst loading of (a) 100 μ g cm⁻² and (b) 200 μ g cm⁻²in 0.1M KOH solution.



Fig. S7 LSV curves at rotation rate from 900 to 2500 rpm and the corresponding K-L plots (inset) of Fe₂O₃@Fe-N-C-800 with the catalyst loading of (a) 100 μ g cm⁻² and (b) 200 μ g cm⁻²in 0.5M H₂SO₄ solution.



Fig. S8 LSV curves at rotation rate from 900 to 2500 rpm and the corresponding K-L plots (inset) of Pt/C with the catalyst loading of 41μ g Pt cm⁻²in 0.1M KOH solution.



Fig. S9 (a) XRD patterns of Fe_2O_3 @Fe-N-C-800 and Fe_2O_3 @Fe-N-C-800-BM; (b, c) TEM images of Fe_2O_3 @Fe-N-C-800-BM.



Fig. S10 (a)Wide XPS survey of the Fe₂O₃@Fe-N-C-800-BM. High-resolution (b) C1s, (c)N1s,

and (d) Fe 2p spectra of the Fe_2O_3 @Fe-N-C-800-BM.



Fig. S11 LSV curves at rotation rate from 900 to 2500 rpm and the corresponding K-L plots (inset) of the Fe_2O_3 @Fe-N-C-800-BMin (a) 0.1M KOH and (b) 0.5M H₂SO₄ solution.