Supporting Information

Hierarchical interconnected macro-/mesoporous Co-containing

N-doped carbon for efficient oxygen reduction reactions

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Figure S1. SEM images of (a) PS colloidal spheres and (b) composites of PVA and PS. Inset: an optical photo of the composites of PVA and PS.



Figure S2. SEM images of (a) HP-Co-CN-550, (b) HP-Co-CN-800, and (c) HP-Co-1000.



Figure S3. The thermogravimetric analysis (TGA) of PS, PVA, and PS/PVA.

Table 1. Atomic concentrations (at%) of C, Co, and heterocyclic N components of all as-prepared samples from XPS analysis.

Sample	Co	С	Ν	N1	N2	N3	N4
HP-Co-CN-800	0.22	77.14	10.13	4.39	2.14	2.36	1.24
HP-Co-CN-900	0.35	81.85	9.08	3.07	1.91	3.57	0.53
HP-Co-CN-1000	0.18	83.72	6.80	0.92	2.01	2.70	1.17
HP-CN	0	82.15	9.12	3.15	1.87	3.51	0.59
Co-CN	0.3	83.12	9.65	3.40	1.95	3.47	0.83



Figure S4 EDS elemental mapping indicating the distribution of N, O, and Co in HP-Co-CN-900.



Figure S5. RDE linear sweep voltammograms recorded for HP-Co-CN-800 and HP-Co-CN-1000 supported on a GC electrode in an O_2 -saturated 0.1M KOH solution at a scan rate of 10 mV s⁻¹ and different rotation rates.



Figure S6. (a) Low-magnification and (b) high-magnification SEM images of as-synthesized HP-CNs. (c) SEM images of the Co-CNs. (d) N_2 adsorption–desorption isotherms and the corresponding pore size distribution (inset) of the Co-CNs.



Figure S7. Linear sweep voltammograms recorded for HP-CNs and Co-CNs supported on a GC electrode (c) K-L plots and (d) electrochemical activity given as the kinetic current density ($J_{\rm K}$) at -0.5 V from HP-CNs, Co-CNs, and HP-Co-CNs, respectively.



Figure S8. (a) Cyclic voltammograms of Pt/C and (b)linear sweep voltammograms of HP-Co-CN-900 in O_2 - or N_2 -saturated 0.1 M KOH solutions as well as O_2 -saturated 0.1 M KOH solution with 3 M CH₃OH.



Figure S9. CV of HP-Co-CN-900 before and after stability test (2000 cycles in oxygen-saturated 0.1 M KOH at a scan rate of 100 mV s^{-1}).