

Supplementary Materials

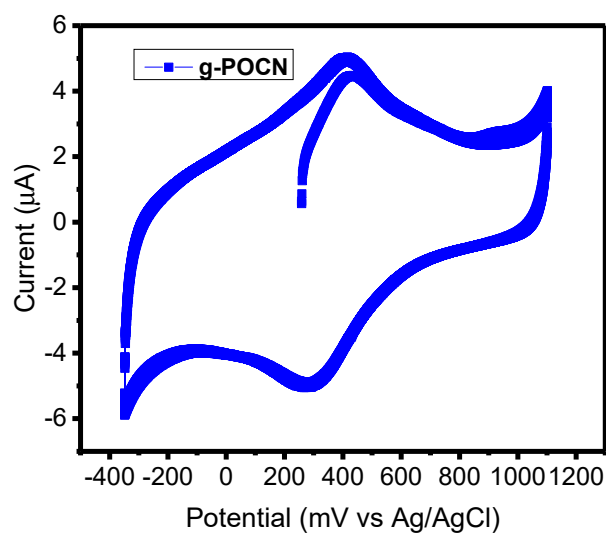


Figure S1. Cyclic voltammetry g-POCN in 0.5 M H_2SO_4 at 100 mV/s (working potential -0.35 V to 1 V).

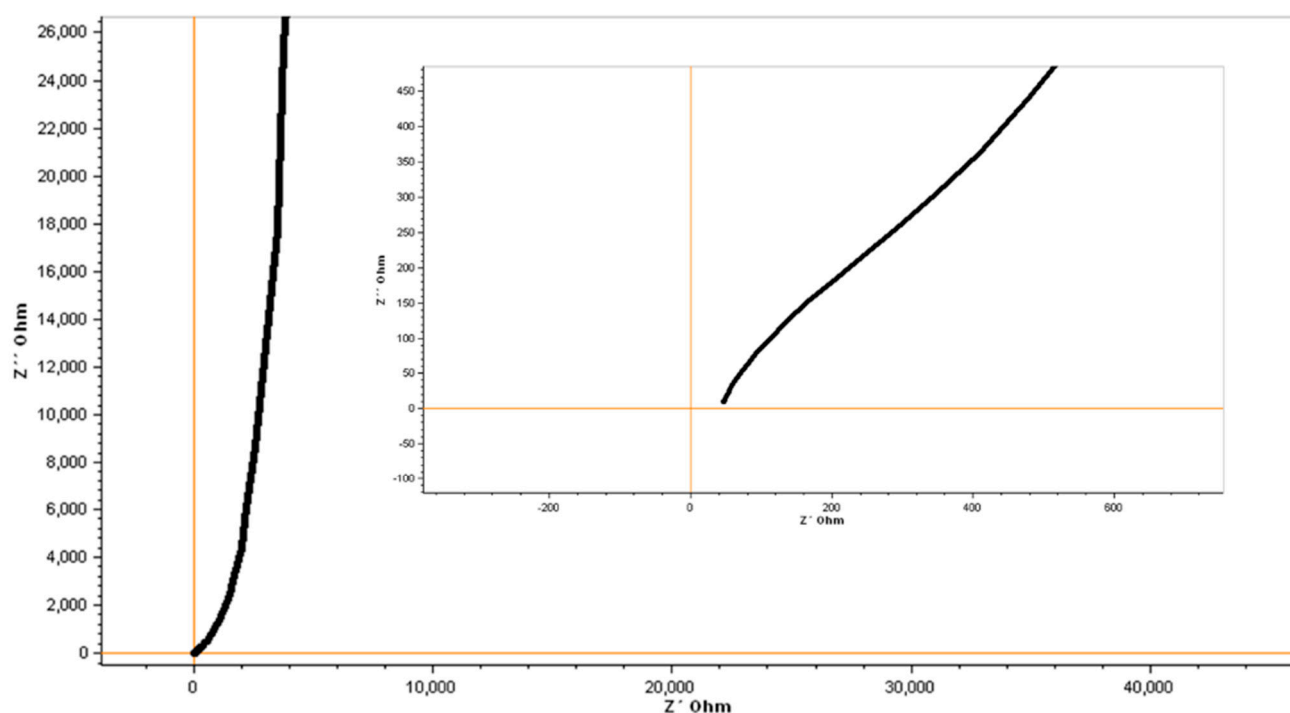


Figure S2. EIS of g-POCN in 0.5 M H_2SO_4 obtained at scan interval from 100 mHz to 100 kHz with amplitude of 10 mV VA to Eoc.

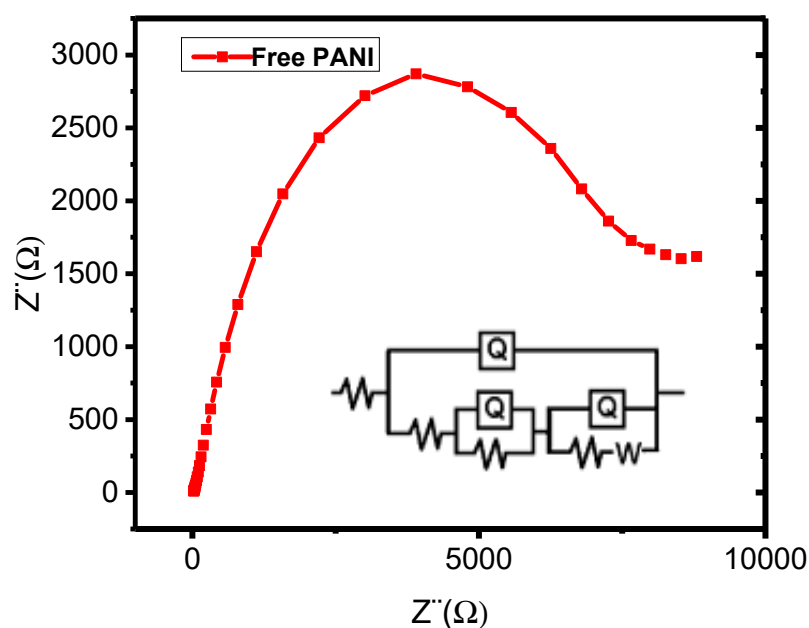


Figure S3. Nyquist diagram of PANI in 0.5 M H₂SO₄ obtained at scan interval from 100 mHZ to 100 kHz with amplitude of 10 mV VA to Eoc.

Table S1. Relevant studies recently reported and their comparison with the present work.

Material	Method of synthesis	Electrolyte	Potential window (V)	Specific capacitance F/g (at current density, A/g, scan rate)	-Capacity retention -Cycle number -Current density (A/g)	Year	Ref
CoFe-LDH/PANI	Hydrothermal synthesis	KOH 6M	-0.1 to 0.7	1686 (1)	98% 10000 20	2022	[1]
PANI/CNT/Elastic polymer fiber	Electro-polymerization	PVA/H ₃ PO ₄	0.0 to 1	111.6 (0.5)	90% 2000 1	2014	[2]
PANI/CNT/Elastic cord	Electro-polymerization	PVA/H ₂ SO ₄	-0.2 to 0.6	394	75.5% 12000 30	2018	[3]
PANI-MWCNT-Ni(OH) ₂	Chemical precipitation	KOH 1 M	0.0 to 0.49	1013 (1)	75% 1000 25	2020	[4]
PANI@Co-Porphyrins	/	H ₂ SO ₄ 1 M	0 to 0.6	823 (0.5)	91% 1000 5	2019	[5]
GO/PANI/CNT-SS	/	KCl 1 M	1.0 to 1.6	331.49 (5 mV/s)	83% 5000 /	2019	[6]
PANI/PEDOT/PANI/	/	PVA/H ₂ SO ₄	0.2 to -0.4	61 F/cm ³ (1 A/cm ³)	73% 1000 /	2019	[7]

UrGO// PEDOT-MoS ₂							
PANI/GO/ PDMS	Micro-emul- sion <i>in-situ</i> polymerization	LiCl 1 M	− 0.2 to 1	576 (1)	/ 10000 5	2018	[8]
BiVO ₄ / PANI	<i>in situ</i> chemical oxidative polymerization	KOH 1 M	0 to 0.66	701 (1)	95.4% 5000 1	2020	[9]
co-MOF/ PANI	<i>in situ</i> chemical oxidative polymerization	KOH 1 M	0 to 0.62 V	504 (1)	90% 5000 2	2020	[10]
CeO ₂ / PANI	/	HCl 1 M	0 to 0.8 V	1452 (2)	~0% 1500 20	2018	[11]
PANI/ g-POCN	Biomimetic Synthesis	H ₂ SO ₄ 0.5 M	−0.2 to 0.9	450.5 (0.2)	80 500 3	2022	This work

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