

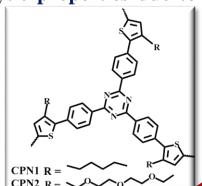
Triazine and Thiophene Based Conjugated Polymer Network: **Electrochromic and Electrocatalytic Properties**

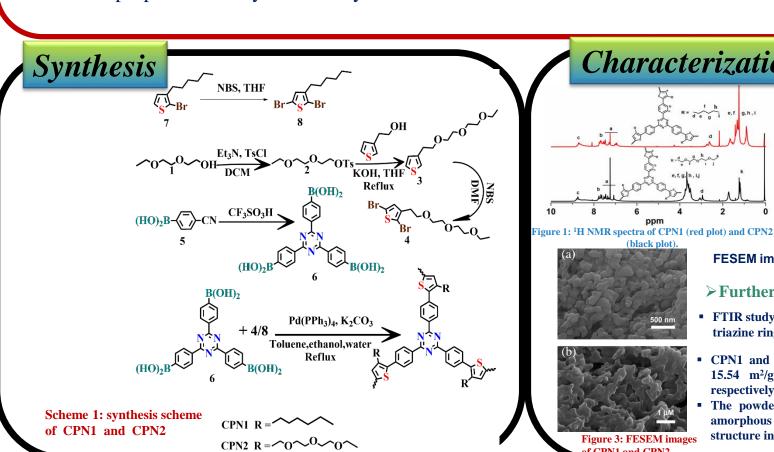
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Introduction

- > Conjugated polymer networks (CPN)^{1,2} which belong to the family of porous organic polymer³, are an emerging class of materials that exhibits promising electrochromic and electrocatalytic properties due to extended π conjugation, interconnected porous structure and high surface area.
- > Herein, we have demonstrated electrochromic (Vis to NIR) and electrocatalytic (Oxygen evolution reaction) property of two triazine and thiophene-based solution processable CPN by coupling the thiophene (donor) unit with a triazine ring (acceptor) through a phenyl ring spacer. The effect of sidechain polarity and functionality on both the properties also systematically studied.





Characterization Mw = 8,861, PDI = 1.75 CPN1 $M_w = 10,089$, PDI =1.94 CPN2 Eluent time (min)

CPN2 **FESEM** image show Interconnected nanospheres

Figure 2: GPC traces of CPN1 and



- triazine ring in the polymer. CPN1 and CPN2 exhibit BET surface area of 34.68 and 15.54 m²/g with average pore of 2.44 and 1.52 nm
- The powder X-ray diffraction (PXRD) analysis revealed amorphous structure indicating interconnected porous structure instead of unimodal pore

Table 2. A summary of the EC parameters of CPN thin-films

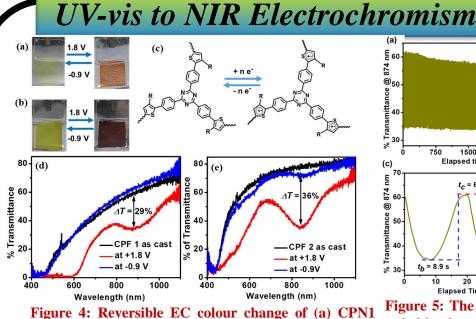
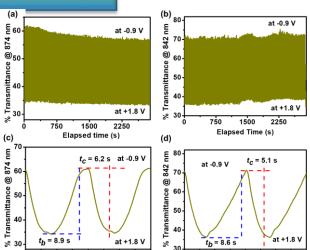
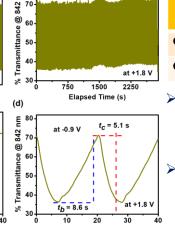


Figure 4: Reversible EC colour change of (a) CPN1 and (b) CPN2 film. (c) Probable mechanism. Transmittance changes of (d) CPN1, and (e) CPN2 film during potential sweeping.





(black plot).

Figure 3: FESEM images

CPN1 62.8 33.8 8.9 116.9 CPN2 74.2 34.9 8.6 163.7

- ➤ The CPN1 exhibits yellow to golden-brown vis-to-NIR EC colour change, whereas, CPN2 exhibits yellow to deep-brown vis-to-NIR EC colour change
- ➤ Both the CPFs exhibited excellent cycling performances with a negligible loss of optical contrast over 150 cycles.

Figure 5: The change in the transmittance with time upon potential switching between +1.8 V and -0.9 V of (a) CPN1 and (b) CPN2 with the electrochromic switching response of (c) CPN1 and (d) CPN2 based ECDs at 874 nm and 842 nm, respectively.

Electrocatalytic activity towards OER (a)

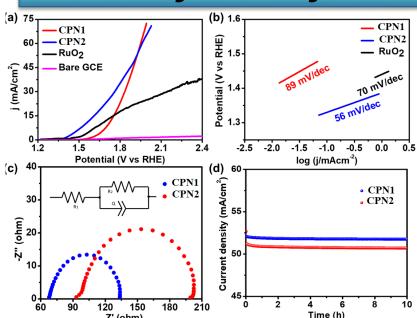


Figure 6: Electrochemical OER activities of CPN1, CPN2, RuO₂, and GCE. (a) LSVs, (b) Tafel plots, (c) EIS Nyquist plot in 1.0 M KOH electrolyte. The corresponding equivalent circuit is given in he inset. (d) The chronoamperometry curves.

(mA/cm²) 0.25 -- CPN2 Current density (51.6

Figure 7: (a) Measurement of the evolved oxygen after 1, 3 and 4 h (b) corresponding chronoamperometry plot.

- > CPN2 with ethylene glycol side chain attached to thiophene moiety shows superior electrocatalytic performance with a current density of 10 mA/cm² at an overpotential of 328 mV compared to CPN1 (10 mA/cm^2 at an overpotential (η) of 488 mV) having hexyl sidechain.
- The achieved OER activity by CPF2 in an alkaline medium is comparable to or even better than that of other well-reported metal-free catalysts with excellent durability.

Conclusion

- Here, we have demonstrated two solution processable CPN based on triazine and thiophene which exhibit UV-vis to NIR electrochromism and OER electrocatalytic activity that are highly dependent on polarity and functionality of the side chain.
- ➤ We believe these materials have potential scope as electrochromic materials for next generation smart window technology and metal free OER electrocatalyst for fuel cell and zinc air battery.

References and Acknowledgement

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- S Halder, N Gupta, RP Behere, B K Kuila, C Chakraborty, Mol. Syst. Des. Eng., 2022,7, 1658-1669. We acknowledge Dr. Chanchal Chakraborty and Sayan Halder for their help for performing electrochemical experiments