Transition metal-nitrogen doped porous carbon as efficient oxygen reduction electrocatalysts for anion exchange membrane fuel cells

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Delving for highly active and cost-efficient electrocatalysts for oxygen reduction reaction (ORR) is crucial for the large-scale application of polymer electrolyte fuel cells.¹ Anion exchange membrane fuel cells (AEMFCs) are promising clean energy devices owing to their mild reaction conditions and high probability of utilizing Pt-free catalysts for ORR.^{2,3} Hence, Pt-free ORR catalysts for AEMFC are of great importance. In this regard, great efforts have been made to develop efficient Pt-free cathodes for AEMFCs application.⁴ This study presents the transition-metal-nitrogen-carbon (M-N-C) electrocatalysts by a robust synthesis approach including the melamine-phloroglucinol-formaldehyde (MPF) polymeric networks embedded with metal. The optimized ORR activity was found by varying the metal content, pyrolysis temperature and different metals (Fe, Co, Mn, and Cu). Amongst, MPF/Fe and MPF/Co catalysts perform better in terms of half-wave potential ($E_{1/2}$) values viz. 0.81 and 0.80 V vs. RHE and are comparable to the commercial Pt/C catalyst attributed to the hierarchical porous structure of the catalysts. Superior electrochemical stability and high-power density in an AEMFC (347 mW cm⁻²) made the present work direct to develop highly efficient M-N-C catalysts for diverse electrochemical energy conversion applications.⁵



Figure 1. Graphical illustration of the strategy for developing the M-N-C electrocatalysts; H₂-O₂ single-cell AEMFC using MPF/M (M = Fe, Co, Mn, Cu) as cathode (1 mg cm⁻²) and Pt-Ru/C (0.4 mg_{Pt-Ru} cm⁻²) as anode catalyst. T = 65 °C; 200 kPa backpressure; 1.0 NLPM flow rate.⁵

Acknowledgements

The present work was financially supported by the Estonian Research Council (grants PRG723, PRG4, PRG1509) and M-ERA.Net project "C-MOF.cell" (SLTKT20445). This research was also supported by the EU through the European Regional Development Fund (MOBJD671 and TK141 "Advanced materials and high-technology devices for energy recuperation systems", TK134 "Emerging orders in quantum and nanomaterials", TK143 "Molecular cell Engineering").

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