Dual-transition metal and nitrogen co-doped silicon oxycarbide-based catalysts for oxygen reduction at the high temperature PEM fuel cell cathode

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The worldwide transition to renewable energy will rely on the introduction and commercialisation of zeroemission and sustainable energy conversion devices. The high temperature proton-exchange membrane fuel cell (HT-PEMFC) is environmentally friendly energy conversion device suitable for heavy-duty transport, stationary, and aviation applications. One topical concern with HT-PEMFC is the development of efficient Ptgroup-metal (PGM)-free oxygen reduction reaction (ORR) catalyst for the fuel cell cathode [1].

Most promising PGM-free catalysts for the HT-PEMFC cathode are the transition metal (TM) and nitrogen functionalised nanocarbon materials, e.g. Fe-N-C [1,2]. More recently, the dual-TM inclusion (e.g. Fe and Cu) has been investigated to achieve more promising ORR electrocatalysis performance compared to the single metal (e.g. Fe-N-C) catalysts [1]. Here we propose a novel silicon oxycarbide (SiOC)-based [3] dual-TM containing catalysts functionalised with N using the Zeolitic Imidazolate Framework-8. The prepared catalysts show promising results during the rotating ring-disk electrode (RRDE) half-cell ORR testing in 0.5 M H₃PO₄ (Figure 1a) compared to the commercial (com.) Pt/C and com. Fe-N-C (Pajarito Powder, LLC). More importantly, to assess the catalyst ORR performance in the conditions more similar to HT-PEMFC, the gas diffusion electrode (GDE) half-cell tests [4] were performed and the catalyst activity of CuFe-SiOC similar to com. Fe-N-C was observed if the *j*-*E* curves without the iR-drop correction are compared (Figure 1b).

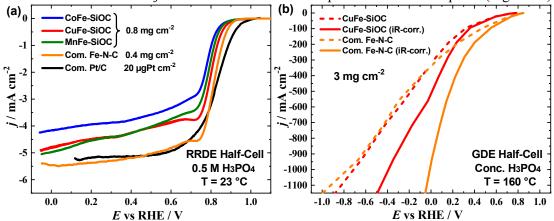


Figure 1: Oxygen reduction reaction studies with (a) RRDE (rotation rate 1600 rpm), (b) GDE half-cell setup with dual-transition metal and nitrogen co-doped silicon oxycarbide-based catalysts and commercial Fe-N-C, Pt/C materials as cathode catalysts.

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References

[1] Q. Meyer, C. Yang, Y. Cheng, C. Zhao, *Electrochem. Energy Rev.*, 2023, 6, 16.

[2] J. Müller-Hülstede, T. Zierdt, H. Schmies, D. Schonvogel, Q. Meyer, C. Zhao, P. Wagner, M. Wark, *J. Power Sources*, 2022, **537**, 231529.

[3] T. Canuto de Almeida e Silva, M. Mooste, E. Kibena-Põldsepp, L. Matisen, M. Merisalu, M. Kook, V. Sammelselg, K. Tammeveski, M. Wilhelm, K. Rezwan, *Catal. Sci. Technol.*, 2019, **9**, 854-866.

[4] T. Zierdt, J. Müller-Hülstede, H. Schmies, D. Schonvogel, P. Wagner, K. A. Friedrich, *ChemElectroChem*, 2024, in press, e202300583.