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Technological implementation of $La_{0.6}Sr_{0.4}CoO_{3-\delta}$ as porous cathode in SOFCs

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Abstract

Strontium substituted lanthanum cobaltite, La_{1-x}Sr_xCoO₃₋₈ (LSC) is a perovskite-type oxide with high electro-catalytic activity towards oxygen reduction at elevated temperatures which makes fabrication of oxygen electrode for solid oxide fuel cells (SOFC) with very low polarisation resistance (R_P) possible. LSC is also a good mixed electronic and oxide

ion conductor (MIEC) which is believed to reduce R_{P} even further as it expands the catalytically active area of the electrode beyond the triple phase boundary (TPB) i.e. where the electrode and electrolyte meet the gas phase. Further, the high conductivity of LSC reduces losses associated with current collection and current constriction. However, the use of LSC as a cathode on Yttria Stabilised Zirconia (YSZ) is problematic due to the high thermal and stoichiometric expansion coefficient of LSC when compared to that of 8% Y2O3 stabilised ZrO2 (8YSZ). Also ionic and electronic blocking reaction products (SrZrO₃, La₂Zr₂O₇) inhibit the direct use of LSC on YSZ electrolyte at high temperatures and necessitates the use an interdiffusion barrier of rare earth doped Ceria (CGO). In this paper we will report on our efforts to fabricate and implement $La_{0.6}Sr_{0.4}CoO_3$ (LSC40) as oxygen electrode in our anode supported SOFCs and discuss advantages and challenges.









Conclusions

easier.

- Oxygen electrodes based on powder of LSC40 (La_{0.6}Sr_{0.4}CoO₃) can be manufactured with good microstructure and high electrochemical performance as a result.
- Oxygen electrodes of porous LSC40 has substantially better electronic conductivity than conventional LSM:YSZ reducing serial resistance and contacting issues.
- The LSC electrode can withstand thermal cycling reasonably well despite its comparably high TEC.

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