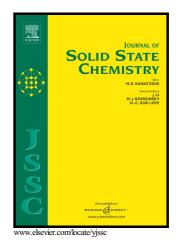
## Author's Accepted Manuscript

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 PII:
 S0022-4596(17)30403-6

 DOI:
 https://doi.org/10.1016/j.jssc.2017.10.005

 Reference:
 YJSSC19967

To appear in: Journal of Solid State Chemistry

Received date: 20 July 2017 Revised date: 23 September 2017 Accepted date: 3 October 2017

Cite this article as: S.Ya. Istomin, A.V. Morozov, M.M. Abdullayev, M. Batuk, J. Hadermann, S.M. Kazakov, A.V. Sobolev, I.A. Presniakov and E.V. Antipov, High-temperature properties of  $(La,Ca)(Fe,Mg,Mo)O_{3-\delta}$  perovskites as prospective electrode materials for symmetrical SOFC, *Journal of Solid State Chemistry*, https://doi.org/10.1016/j.jssc.2017.10.005

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## High-temperature properties of (La,Ca)(Fe,Mg,Mo)O<sub>3-δ</sub> perovskites as prospective electrode materials for symmetrical SOFC

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## Abstract

 $La_{1-v}Ca_vFe_{0.5+x}(Mg,Mo)_{0.5-x}O_{3-\delta}$  oxides with the orthorhombic GdFeO<sub>3</sub>-type perovskite structure have been synthesized at 1573K. Transmission electron microscopy study for selected samples shows the coexistence of domains of perovskite phases with ordered and disordered B-cations. Mössbauer spectroscopy studies performed at 300K and 573K show that while compositions with low Ca-content (La<sub>0.55</sub>Ca<sub>0.45</sub>Fe<sub>0.5</sub>Mg<sub>0.2625</sub>Mo<sub>0.2375</sub>O<sub>3-δ</sub> and La<sub>0.5</sub>Ca<sub>0.5</sub>Fe<sub>0.6</sub>Mg<sub>0.175</sub>Mo<sub>0.225</sub>O<sub>3-δ</sub>) are nearly oxygen stoichiometric, La<sub>0.2</sub>Ca<sub>0.8</sub>Fe<sub>0.5</sub>Mg<sub>0.2625</sub>Mo<sub>0.2375</sub>O<sub>3- $\delta$ </sub> is oxygen deficient with  $\delta \approx$ 0.15. Oxides are stable in reducing atmosphere (Ar/H<sub>2</sub>, 8%) at 1173K for 12h. No additional phases have been observed at XRPD patterns of all studied perovskites and Ce1-xGdxO2-x/2 electrolyte mixtures treated at 1173K-1373K, while Fe-rich compositions ( $x \ge 0.1$ ) react with  $Zr_{1-}$  $_{x}Y_{x}O_{2-x/2}$  electrolyte above 1273K. Dilatometry studies reveal that all samples show rather low thermal expansion coefficients (TECs) in air of 11.4-12.7 ppm K<sup>-1</sup>. In reducing atmosphere their TECs were found to increase up to 12.1-15.4 ppm K<sup>-1</sup> due to chemical expansion effect. Hightemperature electrical conductivity measurements in air and Ar/H<sub>2</sub> atmosphere show that the highest conductivity is observed for Fe- and Ca-rich compositions. Moderate values of electrical conductivity and TEC together with stability towards chemical interaction with typical SOFC electrolytes make novel Fe-containing perovskites promising electrode materials for symmetrical solid oxide fuel cell.

Graphical abstract

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