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## Structural and Electrical Characterization of Nanostructured SrTi<sub>1-x</sub>Fe<sub>x</sub>O<sub>3-δ</sub>

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Abstract - SrTi<sub>1-x</sub>Fe<sub>x</sub>O<sub>3-0</sub> (x= 0.0 to 0.75) nanostructured samples have been synthesized using a modified polymeric precursor method. The effect of the addition of iron on the structural and electrical properties was investigated by X-ray diffraction, Infrared spectroscopy, X-ray absorption spectroscopy and impedance spectroscopy. It was observed that the presence of Fe<sup>3+</sup> ions in the SrTiO<sub>3</sub> lattice creates oxygen vacancies and that the electrical conductivity increases rapidly with the iron content.

In the last decades compounds with perovskite structure and ABO<sub>3</sub> formula (A and B are cations while O is the oxygen anion) have been intensively studied due their interesting physical properties as for example, the ferroelectricity, piezoelectricity, photoluminescence [1-3]. More recently, perovskite nanostructured compounds have been found to be promising materials in current science and technology. In this context, the SrTi<sub>1-x</sub>Fe<sub>x</sub>O<sub>3- $\delta$ </sub> nanostructured compounds has been the object many studies because the substitution of Ti<sup>4+</sup> by Fe<sup>3+</sup> atom creates different types of defects in the structure which have a significative effect on the sensing properties of this system [4,5].

The conventional way to prepare  $SrTi_{1-x}Fe_xO_{3-\delta}$  compounds is based on the solid-state reaction between SrCO<sub>3</sub>, TiO<sub>2</sub> and Fe<sub>2</sub>O<sub>3</sub> at relatively high processing temperature (> 1100°C) [4,5]. In order to decrease particle sizes to a nanoscale, microsized samples have been subjected to ball milling during up to120 hours, which may introduce impurities into them. On the other hand, the synthesis of nanostructured materials by a chemical route appears to be a good alternative to obtain nanostructured powder samples without such inhomogeneities and at low temperature (< 800°C) [6].

In this work, we present the results concerning the synthesis of nanostructured  $SrTi_{1,x}Fe_xO_{3,\delta}$ powders (x= 0.0 to 0.75) by using a modified polymeric precursors method. Electron scanning microscopy images show that the particle size is around 50 nm. X-ray diffraction measurements indicated that samples present a single cubic SrTiO<sub>3</sub> phase forming a solid solution. Moreover, no evidence of a secondary phase was found by X-ray diffraction measurements. The vibrational band related to the FeO<sub>6</sub> octahedra was identified by infrared spectroscopy indicating the presence of iron ions in the SrTiO<sub>3</sub> lattice. In order to investigate the valence state of iron ions, X-ray absorption (XANES) spectra at Fe K-edge were collected. The analysis of XANES spectra indicates the predominance of Fe<sup>3+</sup> ions which induces the formation of oxygen vacancies (Vö). The presence of Fe<sup>3+</sup> ions contributes to the electrical conductivity which increases with the increase of Fe content. It was observed that, with the increasing Fe<sup>3+</sup>, the materials move from an insulator to a semiconductor.

- [1] Y. Luo, I. Szafraniak, V. Nagarajan, R. B. Wehrspohn and M. Steinhart, Integr. Ferroelectrics 59 (2003) 1513-1516.
- [2] P. R. Choudhury and S. B. Krupanidhi, Appl. Phys. Lett. 92 (2008) 102903.
  [3] S. de Lazaro, J. Milanez, A. T. de Figueiredo and V. M. Longo, Appl. Phys. Lett. 90 (2007) 111904.
- [4] A. Rothschild, W. Menesklou, H. L. Tulley, and E. I. Tiffée, Chem. Mater. 18 (2006) 3651-3659.
- [5] P. Meufles, J. Eur. Ceram. Soc. 27 (2007) 285-290.
- [6] L. F. da Silva, M. I. B. Bernardi, L. J. Q. Maia, G. J. M. Frigo, and V. R. Mastelaro, J. Therm. Anal. Cal. (accepted for publication)