Accepted Manuscript

Investigation of Structural and Magnetic Properties of $Nd_{0.7}Ba_{0.3}Mn_{1-x}Ti_xO_3$ (x = 0.05, 0.15 and 0.25) Manganites Synthesized Through a Single-Step Process

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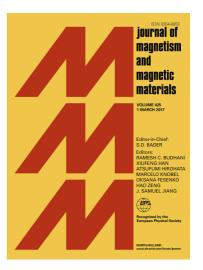
PII: S0304-8853(18)31082-5

DOI: https://doi.org/10.1016/j.jmmm.2018.08.062

Reference: MAGMA 64259

To appear in: Journal of Magnetism and Magnetic Materials

Received Date: 12 April 2018 Revised Date: 31 July 2018 Accepted Date: 22 August 2018



Please cite this article as: D. Kumar, A.K. Singh, Investigation of Structural and Magnetic Properties of $Nd_{0.7}Ba_{0.3}Mn_{1-x}Ti_xO_3$ (x = 0.05, 0.15 and 0.25) Manganites Synthesized Through a Single-Step Process, *Journal of Magnetism and Magnetic Materials* (2018), doi: https://doi.org/10.1016/j.jmmm.2018.08.062

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Abstract: The polycrystalline $Nd_{0.7}Ba_{0.3}Mn_{1-x}Ti_xO_3$ (x = 0.05, 0.15 and 0.25) perovskite manganites have been synthesized using combustion synthesis method followed by singlestep calcination process. Rietveld analysis of the X-ray powder diffraction (XRD) patterns reveals that the samples crystallize in single phase body-centered orthorhombic crystal structure with space group Imma. The lattice parameters and unit cell volume increase exponentially with doping of Ti^{4+} ions for $Nd_{0.7}Ba_{0.3}Mn_{1-x}Ti_xO_3$ (x = 0.05, 0.15 and 0.25). The temperature dependence of magnetization reveals that all samples undergo transition from paramagnetic (PM) to ferromagnetic (FM) phase on cooling below room temperature. The values of Curie temperature (T_C), Curie-Weiss temperature (T_{CW}) and experimental effective paramagnetic moment (μ_{eff}^{exp}) decrease exponentially on the doping of Ti^{4+} ions. The field dependent magnetization measurements reveal coexistence of FM and antiferromagnetic (AFM) states in the low-temperature region. The experimentally observed saturation moment (μ_{sat}^{exp}) of all the samples is lower than the theoretically calculated values and decreases exponentially with doping of Ti⁴⁺, due to AFM component in the FM ordering. The negative slope in the low field region of the M^2 vs. H/M Arrott-plots for all the samples reveals that Nd_{0.7}Ba_{0.3}Mn_{1-x}Ti_xO₃ manganites exhibit a first-order magnetic phase transition. The critical exponents β , γ and δ were estimated for Nd_{0.7}Ba_{0.3}Mn_{1-x}Ti_xO₃ perovskites.

"Keywords: Perovskite Manganite; Combustion Synthesis; X-Ray Diffraction; Rietveld Refinement; Double Exchange Interaction; Critical Exponent."

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