

Nanoscale physics

Nanostructural clustering, structure defects and magnetic properties of the magnetoresistance $\text{La}_{0.6}\text{Sr}_{0.15}\text{Bi}_{0.15}\text{Mn}_{1.1-x}\text{Ni}_x\text{O}_{3-\delta}$ ceramics

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According to X-ray diffraction data, all ceramic $\text{La}_{0.6}\text{Sr}_{0.15}\text{Bi}_{0.15}\text{Mn}_{1.1-x}\text{Ni}_x\text{O}_{3-\delta}$ samples with $x = 0, 0.05, 0.1, 0.15, 0.2$ and 0.3 were single-phase and contain a rhombohedral $R\bar{3}c$ type of distortion. The lattice parameter of a perovskite structure slightly changed with increase in x . On the basis of the defect formation mechanism and the obtained experimental data, it has been established that the real structure is a defect and contains variable valence manganese Mn_A^{2+} , Mn_B^{3+} and Mn_B^{4+} ions as well as cationic $\text{V}^{(c)}$ and anionic $\text{V}^{(a)}$ vacancies. The presence of vacancies leads to the appearance of superstoichiometric manganese Mn_A^{2+} ions in A-positions of the perovskite structure with a formation of nanoscale planar clusters of $\sim 10 - 25$ nm, which has been confirmed by the results of magnetic measurements at $T = 77$ K. In the compositions with $x = 0.05$ and 0.1 , the anomalous magnetic hysteresis is due to 90° exchange antiferromagnetic interactions ($\text{Mn}_A^{2+} - \text{O}^{2-} - \text{Mn}_B^{(3,4)+}$) between the nanostructural cluster and the ferromagnetic matrix structure. During substitution of superstoichiometric manganese for Ni ions is a decrease in the phase transition temperatures and the magnitude of magnetoresistance effect as well as an appearance of a wide phase separation region which consists of coexisting inhomogeneous magnetic phases also including the nanoscale antiferromagnetic clusters.