



Contribution ID: 173

Type: Poster

Low-temperature phase transitions in $\text{PrAlO}_{3-x}\text{SrTiO}_3$ series

Monday 17 September 2018 17:00 (15 minutes)

Phase and structural behaviour of the continuous perovskite solid solution $\text{Pr}_{1-x}\text{Sr}_x\text{Al}_{1-x}\text{Ti}_x\text{O}_3$ have been studied in the temperature range 20 – 295 K by high-resolution X-ray synchrotron powder diffraction. Superb characteristics of the beamline ID22@ESRF allows to detect either subtle splitting of the main perovskite maxima and/or appearance of weak superstructure reflections, thus proving diverse variants of perovskite structure existing in $\text{Pr}_{1-x}\text{Sr}_x\text{Al}_{1-x}\text{Ti}_x\text{O}_3$ series at different compositions and temperatures. It was revealed that the samples with $x = 0.1$ and 0.2 undergo a sequence of structural phase transitions $R\bar{3}c$ – $\text{Im}\bar{m}b$ – $I2/m$, similar to those observed for the parent PrAlO_3 phase. These LT transitions in praseodymium aluminate are unique among all RAlO_3 perovskites and are considered to be caused by the electronic effects involving Pr^{3+} ions, e.g. a coupling between Pr^{3+} electronic states and phonons and/or cooperative Jahn-Teller effects. It was established that temperatures of both $R\bar{3}c$ – $\text{Im}\bar{m}b$ and $\text{Im}\bar{m}b$ – $I2/m$ transitions in $\text{Pr}_{1-x}\text{Sr}_x\text{Al}_{1-x}\text{Ti}_x\text{O}_3$ series systematically decrease from 205 K and 151 K for PrAlO_3 to 170 K and 90 K for $x = 0.2$ sample. Quite different phase behaviour was observed in the SrTiO_3 -rich part of the system. Simultaneous aliovalent substitution of Sr^{2+} and Ti^{4+} species by 10 % Pr^{3+} and Al^{3+} ions increases the temperature of a $\text{Pm}\bar{3}m$ – $I4/mcm$ transition from 105 K in SrTiO_3 to ~250 K in $\text{Pr}_{0.1}\text{Sr}_{0.9}\text{Al}_{0.1}\text{Ti}_{0.9}\text{O}_3$.

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Session Classification: Poster session 1

Track Classification: P8 Functional materials and materials science