

Growth and characterization of $\text{Dy}_{1-x}\text{Y}_x\text{MnO}_3$ single crystals by optical floating zone technique: A combined X-ray diffraction and DC magnetization study

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Rare earth manganites RMnO_3 ($\text{R} = \text{La}^{3+} - \text{Lu}^{3+}, \text{Y}^{3+}$) represent a fascinating family of multiferroic compounds due to an interplay between their charge, lattice and spin. Pure and Y^{3+} substituted DyMnO_3 samples were synthesized by solid state reaction method and grown as single crystals by optical floating zone method. $\text{Dy}_{(1-x)}\text{Y}_x\text{MnO}_3$ single crystals grown in Argon atmosphere, crystallize in hexagonal structure belonging to the $P6_3cm$ space group. The structural analysis has been carried out by Rietveld refinement using synchrotron X-ray powder diffraction (SXRPD) data. The results obtained after the Rietveld refinement clearly demonstrate that the unit cell volume and lattice parameters decrease with increasing Y^{3+} substitution at Dy^{3+} site in DyMnO_3 . Further, magnetic susceptibility (χ_{dc}) measurements on single crystals of $\text{h-Dy}_{(1-x)}\text{Y}_x\text{MnO}_3$ reveal that the Y^{3+} substitution enhances the antiferromagnetic ordering temperature (T_N) from 67 K for h-DyMnO_3 to 72 K for h-YMnO_3 . Unlike the T_N ordering, the first spin reorientation (SR) transition temperature of Mn^{3+} decreases with increasing Y^{3+} substitutions. The remaining two magnetic transitions at still lower temperatures, i.e. Dy^{3+} ordering temperature ($T_{\text{Dy}^{3+}}$) and second SR temperature (T_{SR2}), remain unaffected by Y^{3+} substitution up to $x = 0.75$. We also present a magnetic phase diagram of $\text{h-Dy}_{(1-x)}\text{Y}_x\text{MnO}_3$ showing stability field regions of different transitions. We believe that our study helps in understanding the magnetic transitions of the other h-REMnO₃.