

Magnetism of novel Nd₂Ir₂O₇-type compound

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Rare-earth A₂Ir₂O₇ oxides constitute a family of compounds that is attractive due to their complex properties and application potential. The geometrically frustrated pyrochlore lattice hosts Ir⁴⁺ ions with strong spin-orbit coupling and electron correlations of comparable strength, thereby paving the way for exotic magnetic and topological phenomena. In combination with crystal-field effects and significant *f-d* exchange between the rare-earth and iridium sites, this leads to the emergence of, e.g., an antiferromagnetic Weyl semimetal state [1], fragmented spin ice with monopole-like excitations [2] and spin liquid states [3,4]. Moreover, the existence of these complex phases is inherently tied to the tetrahedral network of the pyrochlore lattice.

Our present work focusses on the *A* = Nd member of the series. In its pyrochlore form Nd₂Ir₂O₇ exhibits two magnetic transitions – the Ir sublattice orders at ~35 K in the all-in-all-out (AIAO) configuration and induces the same configuration on the Nd sublattice at ~9 K. By introducing Pb in the synthesis process, we prepared single crystals of a new Nd₂Ir₂O₇-type compound, denoted NIO. Although this phase is tetragonal (*I*4₁/*a*), it retains key structural characteristics of the cubic pyrochlore structure (*Fd* $\bar{3}$ *m*). We investigate this compound by means of magnetisation and specific heat measurements, demonstrating close similarities between NIO and Nd₂Ir₂O₇. Furthermore, X-ray and neutron single crystal diffraction are employed to determine the crystallographic and magnetic structures, thereby shedding light on the magnetic properties of NIO.

References:

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