





Institut de Minéralogie et de Physique des Milieux Condensés Unité Mixte de Recherche 7590 Code 115, 4 Place Jussieu F-75252 Paris CEDEX 05

SÉMINAIRE Mardi 16 octobre, 10h30

Salle de Conférence, 4ème étage, Tour 22-23, Salle 1 IMPMC, Université P. et M. Curie, 4, Place Jussieu, 75005 Paris

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BaFeO₃, A FERROMAGNETIC IRON OXIDE

A small class of oxides containing iron in a high valence state of Fe⁴⁺ (d^4) has been known. The most representative phase is SrFeO₃ (SFO) crystallizing in the cubic perovskite structure (a = 3.850 Å). SFO and related oxides behave very differently from Fe²⁺- and Fe³⁺-oxides like FeO, Fe₃O₄, and LaFeO₃ (LFO). All these Fe²⁺- and Fe³⁺-oxides are antiferromagnetic (or ferrimagnetic) insulators in their ground states, while the Fe⁴⁺-oxides commonly exhibit a shift toward metallicity and ferromagnetism. The specificity of Fe⁴⁺-oxides can be assigned to the fact that the effective charge transfer energy, Δ_{eff} , drastically drops from ~8eV for FeO and ~5.5eV for LFO to ~-3eV for SFO. The characteristics of "Fe⁴⁺"-oxides should thus result from interplay of Fe *d* electrons and O *p* holes.

Very recently we succeeded in preparing BaFeO₃ (BFO) crystallizing in the cubic perovskite structure (a = 3.97106(1) Å). Physical characterization revealed that BFO has a spiral spin structure of the A-type below 111K but turns ferromagnetic with a large atomic moment of 3.5 $\mu_{\rm B}$ /Fe on application of a small external field of ~0.3 T at 5 K (0.2 T at 77K). BFO is the very first Fe-oxide that shows ferromagnetism at ambient pressure. The magnetic transition temperature increases from 111K to more than 300K up to 40GPa.

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