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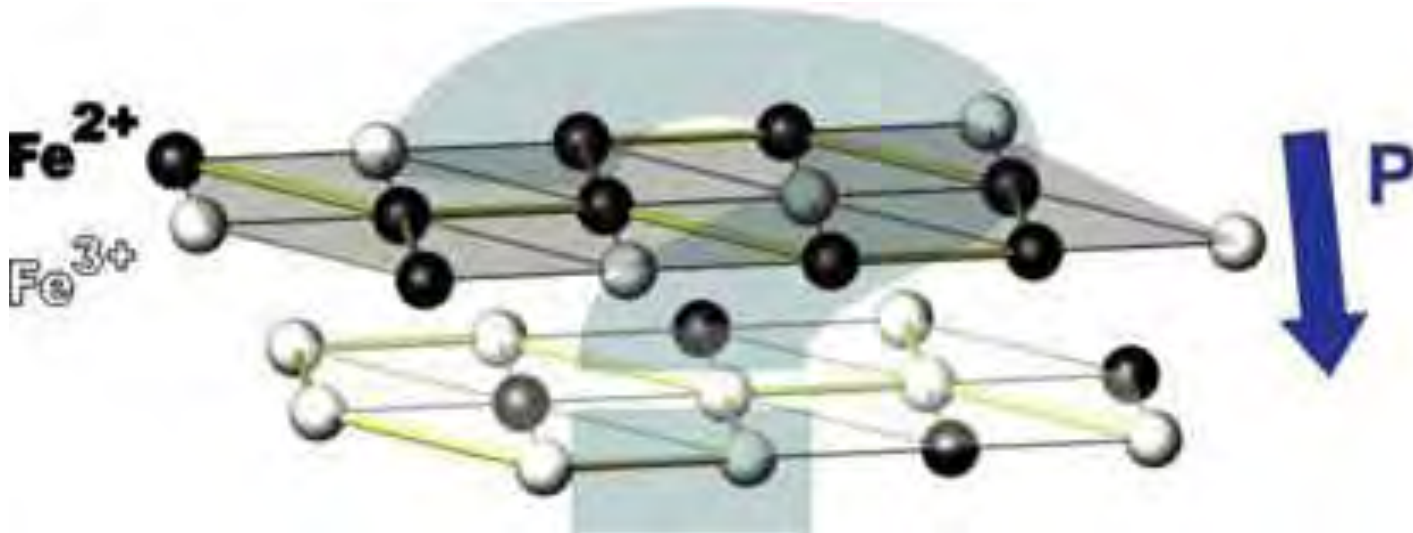
Multiferroicity from charge ordering? A case study

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Magnetolectric multiferroics have a large applications potential. Among possible mechanisms of multiferroicity, ferroelectricity originating from charge ordering (CO) is particularly intriguing because it potentially combines large electric polarizations with strong magnetolectric couplings – but example materials where this is realized are very difficult to find. I will present a case study of such materials, rare earth ferrites, which were recently found to be a non-example. After LuFe_2O_4 had been proposed to be a multiferroic due to ferroelectricity originating from $\text{Fe}^{2+}/\text{Fe}^{3+}$ CO below 320 K, it has become the generally accepted prototypical example of this mechanism, attracting increasing attention. The proposal had been made due to indications of a polar state by dielectric and pyroelectric measurements, and a reasonable model of CO based on the location of superstructure reflections. This model has not been verified though, and the spin-order has been unknown. I will present x-ray and neutron diffraction, and circular dichroism, measurements that allowed i) a full refinement of the CO crystal structure [1], ii) the determination of the spin structures in two nearly degenerate competing magnetic phases [2], and iii) the relation between these orderings [1]. The results reveal a very strong spin-charge coupling. Most importantly, the unambiguously determined arrangement of Fe^{2+} and Fe^{3+} ions excludes any CO-based ferroelectricity – suggesting that LuFe_2O_4 and other rare earth ferrites are not ferroelectric [3]. Time permitting I will also briefly discuss recent time-resolved diffraction experiments on magnetite, another (more likely) candidate material for CO-based ferroelectricity.

[1] J. de Groot, T. Mueller, R. A. Rosenberg, et al., *Phys. Rev. Lett.*, 2012, 108, 187601, [2] J. de Groot, K. Marty, M. D. Lumsden, et al., *Phys. Rev. Lett.*, 2012, 108, 037206, [3] M. Angst, *Phys. Status Solidi RRL*, 2013, 7, 383-400



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