Expanding the library of $S_{eff} = 1/2$ pyrochlore antiferromagnets: Structure and magnetic properties of NaCdCo₂F₇ and NaCdCu₂F₇

A. Kancko,¹ G. Giester,² C. A. Correa,¹ and <u>R. H. Colman¹</u>

¹Charles University, Faculty of Mathematics and Physics, Ke Karlovu 5, 121 16 Prague 2, Czech Republic ²Institute of Mineralogy and Crystallography, University of Vienna, 1090 Vienna, Austria

Materials with a pyrochlore lattice of magnetic ions have experienced decades of intense study due to the frequently exotic electronic properties brought about by the magnetic frustration. Previously these materials were typically rare-earth oxides, but more recently a class of 3d transition metal fluorides have garnered the attention of the condensed matter community. The greater extent of the 3d orbitals, compared to the rare-earth 4f orbitals, leads to greater magnetic exchange and stronger magnetic interaction strengths, leading to mean-field interactions of Θ_{CW} ~ -100 K in all studied members of the family. Despite the strong interactions, no magnetic transitions are observed down to < 4 K, where a spin -glass freezing occurs [1–3]. The spin-glass state is attributed to magnetic bond disorder arising from, fully random, mixed occupancy of the non-magnetic pyrochlore A-site. Theoretical models support this interpretation. The true Hamiltonian of the Co^{2+} pyrochlores is somewhat contentious: inelastic neutron scattering measurements indicate short range correlations with an XY anisotropy [4] and strongly anisotropic g-tensor [5], supported also by PDF analysis of magnetic correlations [6]; despite this, both low-field and high-field magnetisation data for NaCaCo₂F₇ measured along various crystallographic axes show no signs of anisotropy; and ESR measurements indicate a much smaller experimental g-factor of $\tilde{}$ 2, compared to that expected from the INS results. More materials and studies are needed to gain better understanding of these systems.

In this contribution, I will present the structural and magnetic properties of two new members of the family, NaCdCo₂F₇ and NaCdCu₂F₇. I will present a comparison with the previously investigated members. Notably the A-site Na/Cd ion size discrepancy is greater than the previously studied Na/Sr and Na/Ca analogues, leading to greater magnetic bond disorder. In the $S_{eff} = 1/2 \text{ Co}^{2+}$ this leads to a spin-glass transition with an enhanced spin-glass freezing temperature, as expected. The $S = 1/2 \text{ Cu}^{2+}$ pyrochlore appears unique in this family, with avoidance of the spin-glass freezing down to less than our lowest investigated temperatures (0.3 K), making NaCdCu₂F₇ a quantum-spin-liquid candidate.

References:

- [1] J. W. Krizan and R. J. Cava. J. Phys. Condens. Matter 27, 296002 (2015)
- [2] J. W. Krizan and R. J. Cava. Phys. Rev. B 92, 014406 (2015)
- [3] M. B. Sanders et al. J. Phys. Condens. Matter 29, 045801 (2017)
- [4] D. Reig-I-Plessis and A. M. Hallas Physical Review Materials, 5(3), 117 (2021)
- [5] K. W. Plumb *et al.* Nat. Phys. 15, 54 (2019)
- [6] J. Zeisner et al. Phys. Rev. B 99, 1 (2019)