

Frustrated Magnetism and Spin Liquids: An ideal $S = \frac{1}{2}$ kagomé system?

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The idea of destabilizing a Néel state in favour of a spin liquid state by means of geometrical frustration was proposed long ago by Anderson in the context of triangular $S = \frac{1}{2}$ antiferromagnets [1]. The RVB state (resonating valence bond), fundamental of a spin liquid, has since then been invoked in the case of a large number of systems, such as the high- T_c superconductors, spin ladders, etc. From a theoretical point of view, the RVB state, built from a macroscopic number of singlets, is considered to be the fundamental state of the $S = \frac{1}{2}$ kagomé or pyrochlore lattices. On the experimental side, unconventional dynamics has been highlighted in several corner-sharing compounds, but none proved to be close enough to the ideal system to allow detailed comparison with theory. Recently, the $x = 1$ compound of the Paratacamite family $Zn_xCu_{4-x}(OH)_6Cl_2$ was revealed as a structurally perfect $S = \frac{1}{2}$ kagomé system [2], expected for so long! In this family, the Cu/Zn doping induces a structural change from distorted pyrochlore to perfectly decoupled kagomé planes. Our Muon Spin Relaxation results prove the absence of freezing down to 50 mK $\ll J \sim 190$ K [3]. Nuclear Magnetic Resonance results bring insight into the intrinsic susceptibility and dynamical behaviour at low temperature [4]. A quick comparison will be made with the case of the triangular Heisenberg antiferromagnet $NaCrO_2$ [5].

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