Contribution ID: 15 Type: Oral

Neutron scattering experiments with high magnetic fields in organic magnets

Thursday, 3 November 2022 16:30 (30 minutes)

Purely organic magnets with \(\tilde{\to}\)-electron spins have essentially negligibly small spin-orbit couplings and are attractive materials because they are archetypical Heisenberg spin systems in which the quantum fluctuations play an important role. The spin size and the connectivity of the network is the key factor of the novel magnetic states arising from quantum fluctuation. Among the representative stable organic radical skeleton, nitroxide (-N-O·) radical has the advantage of making antiferromagnetic spin networks. The positive and negative partial charges on the N and O atoms, respectively, easily gives the intermolecular contact between the NO groups on which the singly occupied molecular orbital (SOMO), that is the molecular orbital with the unpaired electron, is distributed. The intermolecular overlap between SOMO's always yields the antiferromagnetic interactions [1,2]. The stacking of planar \(\tilde{\ti

Considering the nitroxide radical as magnetic entity we were able to growth and macroscopically study several compounds with different dimensionalities with different magnetic behaviors (frustrated spin ladder, 3D honeycomb with AF LRO or 3D Kagomé layers, etc...). In this presentation several examples will be shown in order to highlight the crucial role of the high magnetic fields to understand the basic physics of these interesting materials. In some cases, neutron scattering experiments with applied magnetic fields have been proposed to complement this knowledge.

Many times, in organic magnets the first step to understand the magnetic behavior is the study of the overlapping between the magnetic molecular orbitals which is directly related with the experimental spin density determination, and this is often done by using polarized neutrons with applied magnetic fields, what will be also presented.

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Track Classification: Experience with static fields | Science & Techniques