

NMR Study of Mott Insulator Ba₂NaOsO₆ at High Field

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Introduction

The strong spin-orbit coupling (SOC) effect in Mott insulators, in which the insulating behavior is driven by electron correlations, can give rise to a variety of exotic quantum phases, such as spin liquid, Wely semi-metal, topological insulator, etc. [1]. A representative material of Mott insulating d¹ double perovskites with cubic symmetry is Ba₂NaOsO₆, in which Na and Os ions inhabit alternate cation B sites. In light of the uncommon ferromagnetic state in Ba₂NaOsO₆, quantum models with multipolar magnetic interactions have been proposed. Based on our earlier work [2], a canted ferromagnetic phase preceded by local point symmetry breaking is found at low temperatures, in line with theoretical predictions. To provide further test of the quantum models, we extend our NMR measurement to high magnetic fields up to 34T. We find that at high field, the change of uniform magnetic field $H_{\rm H}$ is within the error bars.

Experimental

The temperature dependence of ²³Na NMR spectra were measured from 5K to 20K at the applied magnetic field of 34T. The measurements were done using the high homogeneity magnet at Cell 14, NHMFL, Tallahassee, FL. A ⁴He variable temperature insert provided the temperature control. The NMR spectrum was obtained from the sum of the Fourier transforms of the standard solid echo sequence using a homemade NMR spectrometer. A high quality single crystal sample of Ba₂NaOsO₆ was mounted with the applied field parallel to the [100] crystalline axis.

Results and Discussion

Figure 1 shows the temperature evolution of ²³Na NMR spectra at 34 T. Above 17 K. the narrow single peak spectrum characterizes a paramagnetic (PM) state with cubic symmetry. At intermediate temperatures, NMR line broadening and triplet splitting indicate non-zero electric field gradient, marking a broken local point symmetry (BLPS) phase. The emergence of two sets of triplets at low temperatures indicates the emergence of two distinct magnetic sites in a two-sublattice canted ferromagnetic (cFM) phase [2]. In cFM phase, the uniform magnetic field H_u is defined as Hu=1/2(<H₁>+<H₂>), where H₁ is the first moment of the left triplet and H_2 is the first moment of the second triplet. In Figure 2 we show that the variation of Hu in applied fields above 20 T is within the error bars.

Conclusions

We measured the ²³Na NMR spectra of a single crystal sample Fig. 2 The uniform magnetic field Hu at Na site at Ba₂NaOsO₆ under an applied magnetic field up to 34 T. A long-range order magnetic phase preceded by local point symmetry breaking is

identified at low temperatures. We studied the relationship of Hu under an applied external magnetic field and find that the observed change at Hu in applied fields above 20 T are within the error bars.

Acknowledgements

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References

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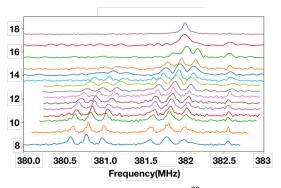
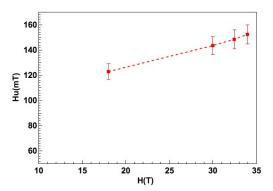


Fig. 1 Temperature evolution of ²³Na spectra at 34T.



different applied external magnetic field.