## **Experimental report**

Proposal:	5-31-2717		<b>Council:</b> 10/2019		
Title:	Charge and magnetic frustration investigated by neutron diffraction in Tb2NbScO7 pyrochlore.				
Research area: I	Physics				
This proposal is a n	ew proposal				
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	Elsa LHOTEL				
Local contacts:	Claire COLIN				
Samples: Tb2Nt	oScO7				
Instrument		Requested days	Allocated days	From	То
D1B		2	2	19/08/2020	21/08/2020
D2B		2	0		
Abstract:					

**Abstract:** The question of structural/charge disorder and its effect on the frustrated Tb pyrochlore ground state is investigated in the Tb2NbScO7 polycrystalline material. We propose a neutron diffraction experiment using a dilution insert to determine the nature of the magnetic ground state (long-range ordered or short-range correlations) stabilized below about 1 K as suggested by an anomaly of the magnetic susceptibility. This study should allow to better understand the Tb pyrochlore magnetic properties, in particular the not yet elucidated Tb2Ti2O7 ground state, and to unravel novel interplays between disorder, charge and spin correlations in this novel material.



## EXPERIMENTAL REPORT

EXPERIMENT N° 5-31-2717, CRG-2725

INSTRUMENT: DIB & IN6/SHARP

DATES OF EXPERIMENT DIB: 19/08/2020 → 21/08/2020, IN4C: 24/08/2020 → 27/08/2020

TITLE:

D1B: Charge and magnetic frustration investigated by neutron diffraction in Tb2NbScO7 pyrochlore. IN6/SHARP/ Crystal field level transitions infrustrated Tb2NbScO7 pyrochlore prone to charge disorder

EXPERIMENTAL TEAM: (names and affiliation)

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LOCAL CONTACTS: Claire COLIN

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Pyrochlore compounds are the archetype of 3-dimensional frustrated materials with a plethora of exotic behaviors (spin ice, fragmentation, quantum spin liquid, order-by-disorder) depending on the rare-earth and on the transition metal residing on the two interpenetrated pyrochlore (corner-sharing tetrahedra) sublattices. One of the most intriguing members of the family is  $Tb_2Ti_2O_7$  with magnetic rare-earth  $Tb^{3+}$ . Instead of displaying the expected Ising antiferromagnetic order, it presents a puzzling correlated state down to the lowest temperature [1]. Additional effects, such as quadrupolar degrees of freedom, must therefore be at play, and enhance quantum fluctuations.

A side route to investigate the origin of the ground state of this material is to change the metal (ex. Sn and Ir in place of Ti) in order to destabilize the fluctuating state and to reveal hidden ingredients of the Hamiltonian [2,3]. A new proposal is to study Niobate compounds where the Nb can be displaced from their ideal position leading to charge correlations, and can be mixed with another transition metal ion [4]. In Nd<sub>2</sub>NbScO<sub>7</sub> for instance, the Nb/Sc disorder seems to lead to frustrated electric dipoles mimicking the spin ice state [5]. This disorder nevertheless does not affect much the ground state in this case. As previously observed with Ti, Sn, Hf or Ge substitutions, the Tb based compounds are much more sensitive to the nature of the metal, and the presence of disorder on these sites should considerably modify the stabilized ground state.

We have thus started an extensive study of the sister compound  $Tb_2NbScO_7$  properties, in order to evaluate the role of disorder on its ground state. Moreover, the coexistence of magnetic moments on one pyrochlore lattice and of dielectric moments on the other one might lead to interesting magnetoelectric effects in these materials.

Using a solid-state synthesis method, we have synthesized a polycrystalline sample of  $Tb_2NbScO_7$  with a site mixing of  $Nb^{5+}$  and  $Sc^{3+}$  on one pyrochlore sublattice.

The sample was characterized by X-ray diffraction and magnetometry. The magnetization measurements versus temperature do not reveal any anomaly down to about 1 K where a separation between the field-cooled and zero-field-cooled measurements is observed due to the magnetic interactions between the Tb. We performed a neutron diffraction experiment using a dilution insert on D1B in order to check the ground state of our sample (see figure 1). We observed the absence of magnetic Bragg peaks indicating no long-range order. Instead, a diffuse signal was observed in the difference between 50 mK and 60 K, which is very similar to the one measured in Tb<sub>2</sub>Ti<sub>2</sub>O<sub>7</sub> [1], and was interpreted as a spin-liquid signature.

We performed another experiment using a dilution insert on SHARP to measure the transitions between the crystal field (CF) levels of  $Tb^{3+}$  by inelastic neutron scattering. Surprisingly, we observe a shoulder at low energy and low temperature that transforms into a quasielastic signal while heating, but no other excitations that could correspond to crystal field level transitions (see Figure 2). We also checked this absence of inelastic signal on the negative energy transfer side at 200 K. This is at variance with the other Tb pyrochlores where a first crystal field level, responsible for the Ising character of the  $Tb^{3+}$ , is observed around 1 meV, and others are visible at higher energies. This absence of inelastic signal could be due to the Nb/Sc disorder, which would broaden the CF levels as in Nd<sub>2</sub>NbScO<sub>7</sub>. Note however that they are broadened but still visible in this sister compound [5]. It could alternatively be a consequence of the charge distribution (disordered Nb<sup>5+</sup>/Sc<sup>3+</sup>) around the Tb<sup>3+</sup> ions shifting the CF transitions to higher energies.

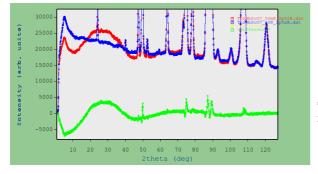


Fig. 1: Powder diffractograms of  $Tb_2NbScO_7$  measured on D1B at 60 K (blue) and 50 mK (red) and there difference (in green) showing the spin liquid like magnetic diffuse scattering.

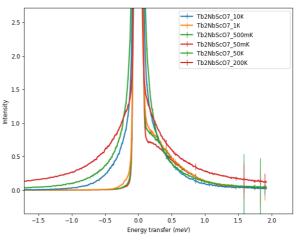


Fig. 2: Thermal variation of the energy spectra of  $Tb_2NbScO_7$  measured on IN6/SHARP at Q=1.2 Å<sup>-1</sup>.

## References:

- [1] J. S. Gardner et al. Phys. Rev. B 64, 224416 (2001).
- [2] I. Mirebeau et al. Phys. Rev. Lett. 94, 246402 (2005).
- [3] E. Lefrançois et al. Phys. Rev. Lett. 114, 247202 2015).
- [5] T. M. McQueen et al. J. Phys.: Condens. Matter 20, 235210 (2008).
- [6] C. Mauws et al., arXiv:1906.10763.