

Mini-rencontre du GDR Meeticc

Vendredi 05 mars 2021
De 13h30 à 15h30

Le lien générique pour vous inscrire à la réunion du 5 mars est le suivant :

<https://zoom.us/meeting/register/tJ0uf-GopjouHd2VgPKuwiryBK1b2bz7VEe9>

Après avoir cliqué sur ce lien et renseigné vos noms et adresse e-mail vous recevrez un e-mail de confirmation contenant les instructions pour rejoindre la réunion.

Programme

*Animateur : Victor **BALEDENT**, Laboratoire de Physique des Solides, Univ. Paris Saclay

	<i>Exposé (30 min + 10 min)</i>
13:30 – 14:10	Structural characterization of nanomaterials through anomalous diffraction Christophe LEFEVRE IPCMS - CNRS - UMR 7504. Université de Strasbourg
14:10 – 14:30	<i>Pause / discussion libre</i>
	<i>Exposés (15 min + 5 min)</i>
14:30 – 14:50	Complex magnetism in high pressure 3d metal oxides Elena SOLANA-MADRUGA Univ. Lille, CNRS, Centrale Lille, ENSCL,
14:50 – 15:10	Resolving the enigmatic high-field behaviour of CeRhIn₅ Sanu MISHRA Laboratoire National des Champs Magnétiques Intenses, Grenoble
15:10 – 15:30	Strange metal from incoherent bosons Anurag BANERJEE Institut de Physique Théorique, CEA Paris-Saclay.

First NAME : Christophe	last NAME : lefevre
Affiliation : CNRS – IPCMS (UMR 7504)	

Presentation type : <i>Long (30 min)</i>	Talk given in : <i>French</i> (please prepare the of the presentation in english)
I authorize the GDR to record a video of my presentation : <i>YES</i>	

Structural characterization of nanomaterials through anomal diffraction

Resonant X-ray diffraction (REXS), benefiting from the tunable energy of the synchrotron, consists of recording the intensity of a Bragg reflection as a function of the energy of the scattered photons crossing an atomic absorption edge. It combines diffraction with extended X-ray absorption fine structure (EXAFS) or X-ray absorption near- edge structure (XANES) methods and is sensitive to both the local environment of the absorbing atom through the anomalous process and to the long-range order involved in the diffraction process. Anomalous scattering experiments have shown over the past ten years their capabilities for locating metal atoms on different sites, even in cases of small occupancies, and even in mixed metal situations [1-2] This method is well adapted to probe cristallographic parameters of powder or thin films materials such as atomic occupancies or positions (e.g. in GaFeO₃ thin films or in CoFe₂O₄ thin films [3-4]). Moreover, it is also possible to take advantage of the possibilities offered by REXS to evidence structural inversions (i.e. the polarization state) of polar structure [5]. Finally, recent numerical approaches show the possibility to evidence the position of light atoms (e.g. oxygen) in these nanomaterials by performing REXS recording in a wide range of energy.

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- [1] Hodeau, J.-L., Favre-Nicolin, V., Bos, S., Renevier, H., Lorenzo, E. Berar, J.-F. Chem. Rev. 101, 1843–1867, (2001).
- [2] Oeckler, O., Schneider, M. N., Fahrnbauer, F. & Vaughan, G., Solid State Sci. 13, 1157– 1161 (2011).
- [3] C. Lefevre, A. Thomasson, F. Roulland, V.Favre-Nicolin, Y. Joly, Y. Wakabayashi, G. Versini, S., C. Leuvrey, A. Demchenko, N. Boudet, N. Viart, J. Appl. Cryst. 49, 1308–1314 (2016)
- [4] E. Martin *et. al.*, J. Alloys Comp. 836, 155425 (2020).
- [5] C. Lefevre, A. Demchenko, C. Bouillet, M. Luysberg, X. Devaux, F. Roulland, G. Versini, S. Barre, Y. Wakabayashi, N. Boudet, C. Leuvrey, M. Acosta, C.

Complex magnetism in high pressure 3d metal oxides.

Elena Solana-Madruga,^{1,2} Cintli Aguilar-Maldonado,¹ Clemens Ritter,³ Olivier Mentré,¹ J. Paul Attfield² and Ángel M. Arévalo-López¹

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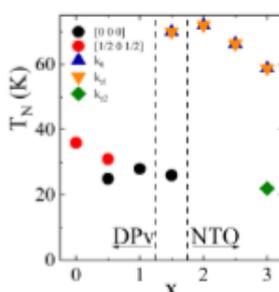
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ABO₃ oxides are intensively studied materials as they exhibit useful chemical and physical properties. Amongst these, the use of high-pressure synthesis techniques allows the small Mn²⁺ cation to be located in the A-site with several structures in competition. The presence of Mn orbital, charge and spin degrees of freedom often induces complex magnetic properties, for instance, the spintronic perovskite MnVO₃-II or the multiferroic MnTiO₃-II with LiNbO₃-type structure.^[1,2]

Tellurium based oxides M₂MTeO₆ (M = Mn and Co) crystallise with corundum-related structures at ambient pressure and show a variety of complex behaviours: Mn₃TeO₆ is a type-II multiferroic,^[3] where two incommensurate magnetic phases (an elliptical helix and a sinusoidal spin density wave) coexist.^[4] Co₃TeO₆ has 5 independent Co sites, providing a rich magnetic phase diagram^[5,6].

Here we present the high pressure modifications HP_Mn₃TeO₆, HP_Co₃TeO₆ and their solid solutions, prepared under 8-15 GPa and 1173 K using a Walker –Multianvil high pressure apparatus. HP_Mn₃TeO₆ has P2₁/n double perovskite (DPV) structure and is a collinear AFM with $k = [\frac{1}{2} 0 \frac{1}{2}]$ below 36 K with an unusual large frustration index due to d¹⁰ Te⁶⁺.^[7] HP_Co₃TeO₆^[8] crystallises with R3 Ni₃TeO₆ (NTO) structure. The strong magnetic frustration arising from stacked honeycomb and triangular magnetic sublattices originates an elliptical helix below 58 K with a $k_0 = [0 0 0]$ and a temperature dependent incommensurate $[0 0 k_{z1}]$ component. Below 23 K, k_{z1} splits and originates a coexisting circular helix magnetic structure with $[0 0 k_{z2}]$ propagation vector.

The HP_Mn_{3-x}Co_xTeO₆ oxides show a coherent evolution between DPV (x = 0.5, 1 and 1.5) and NTO (2 and 2.5) solid solutions, as depicted in the phase diagram (Fig.1). The magnetic behaviour of both polymorphs is, however, strongly affected by the presence of Co²⁺ and Mn²⁺ respectively. The clear B-site preference of Co²⁺ among the DPV compounds turns part of the d⁵-d⁵ magnetic interactions into coexisting d⁵-d⁷, which modifies the spin arrangement to a new AFM structure with $k = [0 0 0]$. Among the NTO-type compounds, with complex incommensurate spin structures as that of HP_Co₃TeO₆, the magnetic transition temperature progressively increases with the Mn content, consistently with the preferred cation order. The presence of d⁵ Mn²⁺ stabilises the elliptical modulation of the magnetic moments into the helix and prevents the thermal dependence of the $[0 0 k_z]$ propagation vectors and their splitting. Such dramatic magnetic changes among both HP polymorphs reflect the essential role of t_{2g} orbitals in the magnetic interactions of these compounds.



References:

- [1] M. Markkula, et al. *Phys. Rev. B*. 2011, 84, 094450.
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Figure 1. Phase diagram for the HP-Mn_{3-x}Co_xTeO₆ solid solutions.

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Please check

<i>the type of presentation</i>	LONG (30 min)		SHORT (15 min)	✓
<i>The talk is given in</i>	ENGLISH	✓	FRENCH	

I authorize the GDR to record a video of my presentation: *Y*

Resolving the enigmatic high-field behaviour of CeRhIn₅

The heavy fermion compound CeRhIn₅ is a prototypical system to investigate the rich physics of strongly correlated electrons as it can be readily tuned to a quantum critical point by hydrostatic pressure, chemical doping, and magnetic fields. While the pressure-induced quantum critical point is long accepted to be of the Kondo-breakdown type, the one induced by the magnetic field at $B_c = 50$ T was suggested to be of the spin-density-wave type. This assumption is based on the observation of additional de Haas-van Alphen frequencies deep inside the antiferromagnetic state, above $B^* = 30$ T, where a novel phase of enhanced in-plane electronic anisotropy emerges [1]. These additional frequencies were interpreted as a signature of an abrupt Fermi-surface reconstruction due to the field-induced itineracy of the f electrons at B^* [2].

In this talk, I will present our comprehensive angular-dependent de Haas-van Alphen study of CeRhIn₅ and its non- f reference compound LaRhIn₅, which establishes the localized character of the f electrons inside and outside of the antiferromagnetic phase. Our results rule out any significant field-induced Fermi-surface reconstruction, particularly across B^* , and suggest that the field-induced quantum criticality in CeRhIn₅ does not conform with the established theoretical models [3]. Further, I will also present our high-field specific heat results, where the observation of a distinct anomaly at B^* suggests it to be a thermodynamic phase transition, probably weakly first-order [4].

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Presentation type : 15 mins	Talk given in : <i>English</i> (please prepare the of the presentation in english)
I authorize the GDR to record a video of my presentation : YES	

Strange metal from incoherent bosons

The breakdown of the celebrated Fermi liquid theory in the strange metal phase is the central enigma of correlated quantum matter. Motivated by recent experiments reporting short-lived carriers, along with the ubiquitous observations of modulated excitations in the phase diagram of cuprates, we propose a model for this phase. We introduce bosons emerging from the remnants of a pair density wave as additional current carriers in the strange metal phase. These bosonic excitations are finite momentum Cooper pairs and thus carry twice the electronic charge, and its net spin can either be zero or one arising from the two spin-1/2 electrons. We show that such a model can capture the famous linear relationship of resistivity with temperature and manifests the Drude form of ac-conductivity with a Planckian dissipation rate. Furthermore, such bosons are incoherent and hence do not contribute to the Hall conductivity. The bosons emerging from the electron pairs of spin-triplet symmetry also reproduce the recently observed linear in-field magnetoresistance[1-3].

Références :

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- [2] J. Ayres *et al.*, unpublished (2020)
- [3] A. Banerjee, *et al.* arXiv:2009.09877 (2020)