

CONDENSED MATTER THEORY SEMINAR

Subject: **DFT+DMFT study of $J=1/2$ insulating iridates**

Speaker: **Junior Prof. Hongbin Zhang (TU Darmstadt)**

Date & time: **Friday, April 20th, 2018 at 3.15 p.m.**

Venue: **Seminar room 1.114**

The competition between spin-orbit coupling, crystal field splitting and electron correlations with comparable magnitude gives rise to many interesting phenomena. For instance, the so-called effective $J=1/2$ state has been observed in many iridates compounds, e.g., Ruddlesden-Popper (RP) $\text{Sr}_{n+1}\text{Ir}_n\text{O}_{3n+1}$ and pyrochlore $\text{RE}_2\text{Ir}_2\text{O}_7$ (RE=Bi, Pr, Nd, Sm, Eu, Y) iridates, where metal-insulator transitions occur driven by the interplay of electron correlations with magnetic ordering. Using first-principles methods, for correlated solids based on density functional theory and dynamical mean field theory (DFT+DMFT), we have investigated the metal-insulator transitions in both classes of iridates. We explore the robustness of the effective $J=1/2$ state against band effects due to itineracy, structural distortion, and strain. We show how single-particle spectra, optical conductivities, and orbital and spin moments change with strain, and we demonstrate that the ground state can be well characterized in terms of an effective energy-dependent $J=1/2$ state. Following this line, designing ideal $J=1/2$ states by tailoring the crystal fields will be discussed. For RP compounds, we demonstrate that the crystal field splittings induced by local structural distortions and hybridization are critical to understand previous experimental results. For pyrochlore compounds, the total energies obtained using charge self-consistent DFT+DMFT method reveal that the all-in-all-out magnetic ordering is stable at low temperature in late rare earth pyrochlores, while a bad metallic state is found in early rare earth pyrochlores, in agreement with experiments.