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Slater and Mott Insulating States in Os- and Ir-Based Transitional Metal Oxides

A.D. CHRISTIANSON, Oak Ridge National Laboratory

The discovery of a novel $J_{\text{eff}} = 1/2$ electronic configuration and spin-orbit assisted insulating state in Sr_2IrO_4 has stimulated a fresh look at metal-insulator transitions where relativistic effects participate on an even footing with other energy scales such as crystal field splitting and electron-electron correlations. There are several view points on the origin of the insulating state in Sr_2IrO_4 , but the most prominent is that spin-orbit coupling modifies the electronic configuration such that a Mott insulating state emerges despite the relatively modest electron-electron correlations within the $5d$ orbitals. An alternative viewpoint is that magnetic effects enable the opening of the electronic gap giving rise to the insulating state or a Slater metal-insulator transition. Here we describe realizations of both Mott and Slater insulators in the context of Os- and Ir-based $5d$ transition metal oxides. NaOsO_3 , exhibits a continuous phase transition at 410 K where antiferromagnetism appears in conjunction with the onset of insulating behavior. A combination of neutron diffraction and magnetic resonant x-ray scattering enables the conclusion that G-type magnetic order occurs at the metal-insulator transition providing microscopic evidence that NaOsO_3 is the first three dimensional realization of a Slater insulator. On the other hand we have probed the robustness of the $J_{\text{eff}} = 1/2$ Mott insulating state through studies of $\text{Sr}_2\text{Ir}_{1-x}\text{T}_x\text{O}_4$ (T=Mn, Ru). For both Mn and Ru doping we find that despite qualitative changes in the magnetic order the $J_{\text{eff}} = 1/2$ electronic configuration remains robust. In particular, for Ru-doping the signatures of the $J_{\text{eff}} = 1/2$ state are observed for all concentrations where magnetic order is present. Finally, we have investigated Ca_4IrO_6 which appears to exhibit a nearly ideal $J_{\text{eff}} = 1/2$ state which is unperturbed by deviations from cubic crystal field level splitting.