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**Seminar****Oxides for Energy Storage and Energy-Efficient Devices****Naga Phani Aetukuri****IBM Almaden Research Center & Stanford University, CA**

Metal oxides are a remarkable class of compounds that exhibit almost every known condensed matter phenomenon such as ferroelectricity, ferromagnetism, anti-ferromagnetism, high- $T_c$  superconductivity and metal-insulator transitions. Furthermore, Li-intercalating metal oxides such as  $\text{LiCO}_2$  and  $\text{LiFePO}_4$  are used as cathode materials in Li-ion batteries, which have led to the widespread adoption of portable electronics and to the development of electric automobiles. In this talk, we will present our work on two distinct oxide systems –  $\text{Li}_2\text{O}_2$  and  $\text{VO}_2$  – with applications in energy storage and low power electronic devices, respectively.

First, we will discuss the formation and morphological evolution of  $\text{Li}_2\text{O}_2$  in the context of Li- $\text{O}_2$  batteries, where  $\text{Li}_2\text{O}_2$  is the battery's discharge product. During the battery discharge, electrochemical deposition of  $\text{Li}_2\text{O}_2$  causes electrode passivation because  $\text{Li}_2\text{O}_2$  is an electronic insulator. This limits the maximum deposited thickness of  $\text{Li}_2\text{O}_2$  and results in poor battery capacity. In this work, we show that activating an electrochemical solution-mediated pathway favors the deposition of large  $\text{Li}_2\text{O}_2$  particles. Consequently, there is manifold increase in the battery's discharge capacity. We discuss the design rules for selecting electrolyte solvents that favor this alternate pathway and the implications of this research for metal-air batteries.

Next, we will discuss our work on  $\text{VO}_2$ -based heterostructures and devices.  $\text{VO}_2$  is an archetypal correlated electron system that exhibits a near room temperature metal-insulator transition (MIT) with a concomitant structural phase transition. In this second part of our presentation, we will discuss the control of MIT in liquidelectrolyte gated  $\text{VO}_2$  devices and epitaxially-strained single-crystalline  $\text{VO}_2$  films. We comment on the role of oxygen vacancies in inducing the MIT in electrolyte-gated devices. And, using epitaxially-strained  $\text{VO}_2$  films, we show that the MIT in  $\text{VO}_2$  can be deterministically controlled by tuning the V 3d orbital configuration. We discuss possible device strategies that exploit MIT in  $\text{VO}_2$  and related materials.

***Thursday, Jan 22nd 2015******11:30 AM (Tea/Coffee at 11:15 AM)******Seminar Hall, TCIS***