

# INORGANIC SEMINAR

## “Efficient Conversion of Carbon Dioxide to Fuels using Bismuth Materials that Display Tunable Catalytic Profiles”

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Monday, October 22nd, 4:00pm  
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### Abstract:

The electrochemical reduction of CO<sub>2</sub> provides a pathway to address current issues in solar energy storage and the sustainable production of fuels. Many such approaches, however, have been hampered by a lack of affordable platforms that can efficiently promote CO<sub>2</sub> valorization with high selectivity and reasonable kinetics. It is in response to this need that we have developed inexpensive cathode materials comprised of post-transition metals for conversion of CO<sub>2</sub> to CO, HCO<sub>2</sub>H and other value-added reduced carbon compounds. We find that these platforms are easily prepared and the outcome of CO<sub>2</sub> conversion can be tuned in the presence of varied room-temperature ionic liquids (RTILs). For example, we have prepared and studied a class of bismuth-based materials that promotes the reduction of CO<sub>2</sub> to CO with fast kinetics and high efficiencies in the presence of imidazolium ([Im]<sup>+</sup>) based RTILs. A multi-pronged initiative has elucidated the mechanistic pathways and molecular design principles that drive conversion of CO<sub>2</sub> to CO at the Bi/[Im]<sup>+</sup> interface. We will discuss how integrated spectroscopic (XPS, XAS, XR, etc.) electroanalytical and computational methods have been used to interrogate the dynamics between electrocatalyst, RTIL and CO<sub>2</sub> at the cathode/electrolyte interface to provide insight into the pathway(s) by which the Bi/[Im]<sup>+</sup> platform activates CO<sub>2</sub>. We have also evaluated the electrochemistry of the Bi-cathodes in the presence of non-[IM]<sup>+</sup> RTILs. These studies reveal that subtle modification of the RTIL structure, leads to a significantly different catalysis with CO<sub>2</sub>. Electrolysis of CO<sub>2</sub> over Bi in the presence of RTILs generated from the organic base DBU promotes the reduction of CO<sub>2</sub> to yield formate (instead of CO) via an orthogonal CO<sub>2</sub> activation pathway. The ‘catalytic plasticity’ that is displayed by Bi/[RTIL] platforms, along with implications for future discovery of catalyst/electrolyte combinations that can enable CO<sub>2</sub> conversion and solar fuels production will be discussed.

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