

**DEPARTMENT OF CHEMISTRY AND ENVIRONMENTAL SCIENCE
SEMINAR SERIES
SPRING 2022**

**DATE: WEDNESDAY, MARCH 30, 2022
LOCATION: TIERNAN HALL – LECT. HALL 2
TIME: 1:00PM-2:20PM**

GUEST SPEAKER

Dr. Xiaodong Michael Shu
Associate Chair
Department of Chemistry
University of South Florida
Tampa, Fl

TOPIC

The Two Interlocked Wheels in Chemistry Research:
Rational Design and Serendipity

ABSTRACT

Homogeneous gold catalysis has been developed explosively during the past decades. Despite the remarkable electrophilic activation of alkynes by cationic Au(I) catalysts, such as PPh₃Au+, one challenge is to overcome their poor stability at high temperature. However, in order to activate some less reactive substrates, such as internal alkynes, harsher reaction conditions are usually required. As a good σ-donor and π-receptor, triazole has been applied as a ligand to improve the stability of cationic Au(I) catalysts. Taking advantage of the good stability of triazole-Au(I) complexes (TA-Au), we successfully achieved good reactivity of intermolecular hydroamination for both terminal and internal alkynes. Unlike previous reported gold catalysts, the TA-Au catalysts activate alkynes selectively over alenes. With this excellent chemoselectivity, TA-Au catalysts showed interesting reactivity in propargyl ester and vinyl ether rearrangement. This facilitates the development of otherwise challenging transformations, for instance, asymmetric synthesis of substituted alenes, Schmittel cyclization, dienal synthesis, and so on.

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Seminar Coordinator:

Dr. Genoa Warner – grw4@njit.edu

Dr. Amir Varkouhi - amir.k.varkouhi@njit.edu