

UNIVERSITY OF RHODE ISLAND
Department of Chemistry

SEMINAR

Room 234 Pastore Hall
3:00 P.M., Monday, Sept. 25, 2017

Prof. Mary P. Watson

Department of Chemistry & Biochemistry
University of Delaware
Newark, DE

"Nickel Catalyzed
Cross-Coupling Reaction of Alkyl
Amine and Alcohol Derivatives"

HOST

Mindy Levine
Department of Chemistry
401-874-4243

Nickel-Catalyzed Cross-Coupling Reactions of Alkyl Amine and Alcohol Derivatives

*Prof. Mary P. Watson
Dept. of Chemistry & Biochemistry
University of Delaware*

Transition metal-catalyzed cross-coupling reactions have revolutionized organic synthesis, particularly the construction of bonds to sp^2 -hybridized carbons. However, the discovery of analogous reactions of C_{sp^3} electrophiles have lagged behind, despite their potential to deliver a range of important targets, including chiral molecules in high enantiopurity. Towards solving this challenge in organic synthesis, we have developed a range of nickel-catalyzed cross-couplings of alkyl amine and alcohol derivatives. In particular, we have developed stereospecific, nickel-catalyzed cross couplings of both benzylic amine and alcohol derivatives. These reactions utilize starting materials that are readily available in high optical purity; proceed with high levels of stereochemical fidelity; employ air-stable, functional group tolerance coupling partners, such as aryl boronic acids; and display excellent functional group tolerance. This strategy is also effective for the arylation of allylic electrophiles, and the use of alternative coupling partners to enable borylations. These reactions provide a range of highly enantioenriched products with tertiary and quaternary stereocenters, including molecules important for their biological activity. We have also developed nickel-catalyzed cross-couplings of alkyl amine derivatives with non-activated alkyl groups (non-benzylic, non-allylic). This exciting new chemistry is particularly useful for late-stage functionalization of alkyl amines. The optimization, scope, and mechanistic studies of these reactions will be presented.

