

New Frontiers in Catalytic Heteroatom Transfer: Applications to Late-Stage Oxidation and Small Molecule Synthesis

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Heteroatom transfer reactions - frequently used by Nature for both the biosynthesis of natural products and the metabolism of xenobiotics - are fundamentally useful for the synthesis of pharmaceuticals and other bioactive compounds. As an example of one application, the siteselective oxidation of carbon-hydrogen bonds can be a powerful tool, especially when applied to late-stage functionalization. Transition metal catalysis of site-selective C-H oxidation has rapidly advanced in recent years, but there are nevertheless substantial challenges that remain with regards to achieving catalyst-controlled selectivity. Research in the Hilinski lab has focused on advancing organocatalysis as an alternative approach to transition metal catalysis of C-H functionalization, in order to exploit its potential advantages and complementarity. In this presentation, multiple approaches to organocatalytic C-H oxidation that have resulted in methods for C-H hydroxylation and the first organocatalytic method for C-H amination will be presented, with a focus on catalyst design, selectivity in complex settings, and mechanism. More recently, the lab's focus on applications of catalytic heteroatom transfer has broadened to include other late-stage oxidations relevant to drug discovery (e.g. metabolic N-oxidation) and the discovery of novel cycloaddition reactions initiated by heteroatom transfer. Recent results in these areas will therefore also be presented, with a focus on applications relevant to bioactive small molecule synthesis.

Date: Wed, Nov. 14, 2018

Time: 4:30-5:30 pm Location: 208 Clark Hall

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Students, meet the speaker over coffee and cookies in the Bennett Conference room at 3:30 pm