

Université de Montréal, Montréal, QC, Canada

THIRD INTERNATIONAL SYMPOSIUM
ON C-H ACTIVATION

May 30 – June 2, 2016

IL10 – Functionalization of sp^3 C-H Bonds
with Transition Metal Catalysts and Organocatalysts

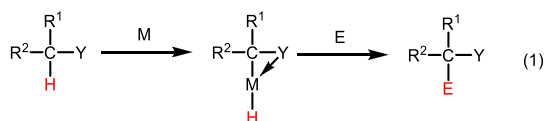
Shun-Ichi Murahashi*

Department of Chemistry, Osaka University, Furuedai, Suita, 1-22-11, 565-0874, Japan

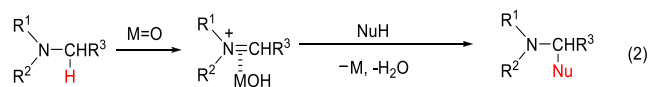
E-mail: murasyt@pearl.ocn.ne.jp

Two methods for functionalization of C–H bonds will be presented.

- 1) The C–H functionalization of substrates such as amines, nitriles, ketones, isocyanates α to heteroatoms with transition-metal catalysts gives α -transition-metalated intermediates, from which various useful methods for unique and selective carbon-carbon bond formation can be constructed (eq 1).¹ Acid-base ambiphilic transition metal catalyst would be a future key catalyst.²
- 2) The C–H functionalization by bio-inspired metal catalysts³ and biocatalysts⁴ with molecular oxygen or peroxides provides useful methods for catalytic C–H oxidative functionalizations of various substrates such as amines, amides, and even non-activated hydrocarbons under mild conditions (eq 2).



M; Ru, Ir, Pd, Rh, Re
Y; NR², CN, C=N, C=O, CF₃



References

1. *J. Am. Chem. Soc.* **1978**, *100*, 348; *J. Am. Chem. Soc.* **2009**, *131*, 10824; *Angew. Chem. Int., Ed.* **2009**, *48*, 2047; *Acc. Chem. Res.* **2000**, *33*, 225-233; *Chem. Rev.* **1998**, *98*, 2599; Ruthenium in Organic Synthesis, Wiley-VCH, S.-I. Murahashi, Ed., **2004**, 53-93.
2. *Angew. Chem. Int., Ed.* **2003**, *42*, 3302.
3. *J. Am. Chem. Soc.* **1988**, *110*, 8256; *J. Am. Chem. Soc.* **1990**, *112*, 7820; *J. Am. Chem. Soc.*, **2003**, *125*, 15321; *Angew. Chem. Int. Ed.* **2005**, *44*, 6931; *J. Am. Chem. Soc.* **2008**, *130*, 11005; (Account review): *Angew. Chem. Int. Ed.* **2008**, *47*, 2079; *Angew. Chem. Int. Ed.* **1995**, *34*, 2443-2465; *Chem. Soc. Rev.* **2008**, *37*, 1490-1501.
4. *J. Am. Chem. Soc.* **1989**, *111*, 5002; *J. Am. Chem. Soc.* **2003**, *125*, 2868; *Angew. Chem. Int. Ed.* **2005**, *44*, 1704; *Chem Commun.* **2014**, *50*, 10295; *Org. Biomol. Chem.* **2015**, *13*, 7999-7613.