Cobalt-Catalyzed Aromatic C-H Bond Functionalization/Cyclization Processes

Mingliang Li and <u>(Michael) Fuk Yee Kwong</u>* Department of Chemistry, The Chinese University of Hong Kong, Shatin, New Territories, Hong Kong, China

Email: fykwong@cuhk.edu.hk

Our research group has been interested to develop new method in assembling polycyclic compounds using transition metal catalysts.^[1,2] Here, a cobalt-catalyzed chelation-assisted sequential C–H activation/C–C cleavage/C–H cyclization of aromatic amides with alkylidenecyclopropanes (ACPs) is presented.^[3] This process allows the sequential formation of two C–C bonds, in which it is in different to previous report of using Rh catalyst for the formation of C–N bond. The inexpensive catalyst system exhibits good functional group compatibility and relatively broad substrate scope. The desired products can be easily transformed into polycyclic lactones with *m*-CPBA. Mechanistic studies revealed that the tandem reaction proceeds through a sequential C–H cobaltation, β -carbon elimination and intramolecular C–H cobaltation pathway.



References

- [1] Fu, W. C.; Wang, Z.; Chan, W. T. K.; Lin, Z.; Kwong, F. Y. Angew. Chem. Int. Ed. 2017, 56, 7166.
- [2] Zhao, Q.; Fu, W. C.; Kwong, F. Y. Angew. Chem. Int. Ed. 2018, 57, 3381.
- [3] Li, M.; Kwong, F. Y. Angew. Chem. Int. Ed. 2018, 57, 6512.