## DIRECTED ENANTIOSELECTIVE COUPLING OF ALDEHYDES AND ALKENES

Alexander Preinfalk, Jing Li, and Nuno Maulide \*

Institute of Organic Chemistry, University of Vienna, Währinger Strasse 38, 1090 Vienna, Austria nuno.maulide@univie.ac.at

In recent years, coupling approaches combining aldehydes and alkenes, readily available feedstocks, have started to emerge as an appealing alternative to classical carbonyl addition reactions (Fig. 1a). However, existing enantioselective catalyst systems invariably deliver the "branched" product and typically require activated olefin reactants. Enantioselective methods to access the highly valuable linear products (Fig. 1b) using a direct redox-neutral coupling of simple unactivated  $\alpha$ -olefins have not yet been developed. In this presentation we will detail an enantioselective redox-neutral coupling that takes advantage of a remote directing group to solve problems of regioand stereoselectivity.

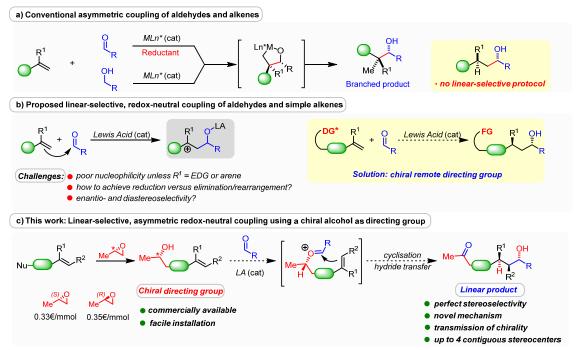


Figure 1. a) Reductive coupling of aldehydes and alkenes; b) and c) a novel linear-selective, enantioselective redox-neutral coupling via a remote directing group.

<sup>[1]</sup> K. D. Nguyen, B. Y. Park, T. Luong, H. Sato, V. J. Garza, M. J. Krische, Science 2016, 354, aah5133.

<sup>[2]</sup> a) M. Kimura, A. Ezoe, K. Shibata, Y. Tamaru, *J. Am. Chem. Soc.* 1998, *120*, 4033–4034; b) H. K. Lin, L. M. Tian, I. J. Krauss, *J. Am. Chem. Soc.* 2015, *137*, 13176–13182.

<sup>[3]</sup> a) J. Li, A. Preinfalk, N. Maulide, J. Am. Chem. Soc. 2019, 141, 143–147; b) J. Li, A. Preinfalk, N. Maulide, Angew. Chem. Int. Ed. 2019, 58, 5887–5890.