# Stereoselective Synthesis of Fluorinated Heterocycles Using Hydroformylation

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Stereoselective synthesis of fluorinated heterocyclic compounds is an active area of research with many applications in the synthesis of bioactive compounds.[1]

In this poster we will discuss a combination of enzymatic resolution and hydroformylation to access some heterocyclic fluorinated scaffolds in a straightforward manner. Finally, the transformation of one of the chiral building blocks into a novel and potentially useful agrochemical intermediate will be described.

[1] (a) Bioorganic and Medicinal Chemistry of Fluorine; Begue, J.-P., Bonnet-Delpon, D., Eds.; Wiley: Hoboken, NJ, 2008. (b) Fluorine In Medicinal Chemistry And Chemical Biology; Ojima, I., Ed.; Wiley-Blackwell: 2009. (c) Nie, J.; Guo, H.-C.; Cahard, D.; Ma, J.-A. Chem. Rev. 2011, 111, 455–529.