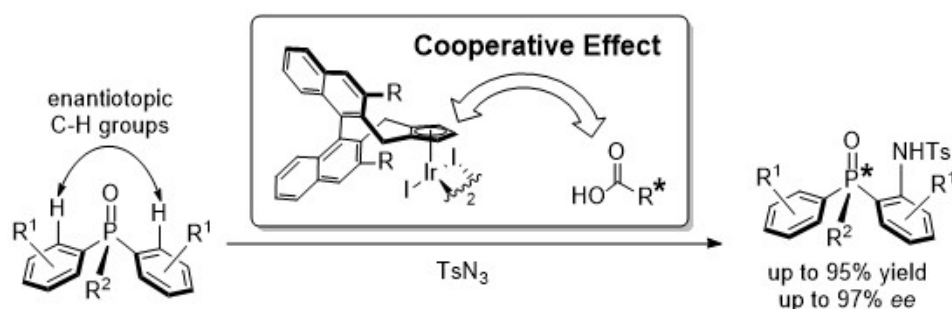


**Chiral Cp<sup>X</sup>Ir(III) Catalyzed C-H Amidation Leading to P-Chiral Arylphosphine Oxides**Y. Jang<sup>1</sup>, M. C. Dieckmann<sup>1</sup>, N. Cramer<sup>1\*</sup><sup>1</sup>Laboratory of Asymmetric Catalysis and Synthesis, EPF Lausanne

Organophosphorus compounds with *P*-stereogenic centers are valuable motifs in pharmaceuticals, agrochemicals, organocatalysts and ligands.<sup>[1]</sup> Only a limited number of catalytic enantioselective approaches have been developed to access molecules with a *P*-stereogenic center.<sup>[2]</sup>

Chang *et al.* reported an Ir(III) catalyzed amidation of arylphosphine oxides that proceeds in modest enantioselectivities.<sup>[3]</sup> We report that our recently developed chiral Cp<sup>X</sup>Ir(III) complex,<sup>[4]</sup> in combination with a chiral carboxylic acid, provides a highly selective C-H amidation process. A very strong cooperative effect between the chiral Cp<sup>X</sup>Ir(III) complex and the carboxylic acid was discovered. This proved to be pivotal for high enantioselectivities and yields.



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