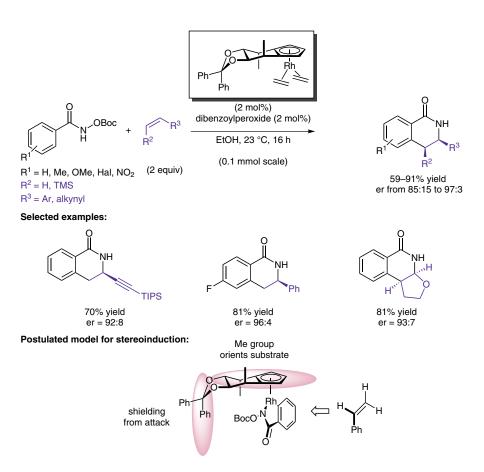
B. YE, N. CRAMER* (ECOLE POLYTECHNIC FÉDÉRALE DE LAUSANNE, SWITZERLAND) Chiral Cyclopentadienyl Ligands as Stereocontrolling Element in Asymmetric C-H Functionalization Science 2012, 338, 504-506.

Chiral Cp Ligands in Rhodium-Catalyzed Asymmetric C-H Functionalization



Significance: A rhodium complex with a chiral Cp ligand that catalyzes an enantioselective synthesis of isoquinolones via a directed C-H bond functionalization is reported. Often, in half-sandwich transition-metal-catalyzed reactions, Cp remains the sole permanent ligand on the metal. Thus, despite the challenges, the development of chiral Cp ligands for inducing enantioselectivity is a powerful approach.

Comment: The highly effective Cp ligand reported is postulated to control the spatial orientation of the coupling partners. For instance, the ligand is C2-symmetric to avoid diastereomeric coordination of the metal. The benzophenone ketal shields one face of the substrate and the equatorial methyl group pushes the bulky Boc group away. The controlled trajectory of the attacking alkene gives rise to the stereo-configuration of the product.

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Category

Metal-Catalyzed Asymmetric Synthesis and Stereoselective Reactions

Key words

rhodium

chiral cyclopentadienyl ligands

isoquinolones

asymmetric C-H functionalization

