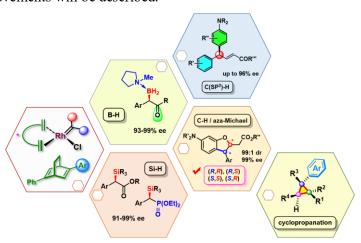
Asymmetric Carbene-Transfer Reactions Catalyzed by Low Coordination State Rhodium(I)

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Transition metal-catalyzed asymmetric carbene insertion reactions have been recognized as one of the most powerful methods for the construction of C-C or C-heteroatom bonds. In recent years, our group has been interested in Rh(I)-carbene chemistry, and achieved the first asymmetric metal-carbene insertion into B-H and Si-H bonds using $Rh(I)/C_1$ -symmetric diene complexes as catalysts. With the established Rh(I)-carbene chemistry, we also discovered that C-H insertion reaction of styryldiazoacetates with aniline derivatives could be achieved in a highly region- and enantio-selective manner. Moreover, the rhodium(I)/chiral diene catalytic system also enables efficient asymmetric Rh(I)-carbene-directed S-H, $C(sp^3)$ -H, N-H, O-H insertions, as well as intramolecular cyclopropanations. In this presentation, these results together with some new achievements will be described.



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