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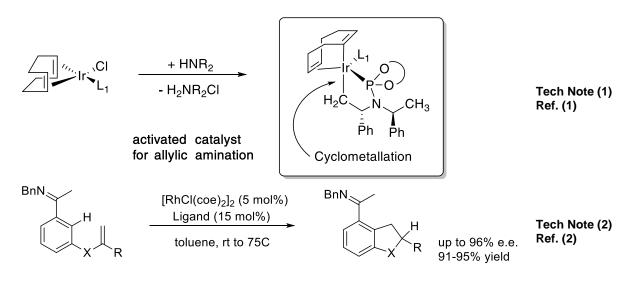


amine

Ph. O P-N Ph Ph

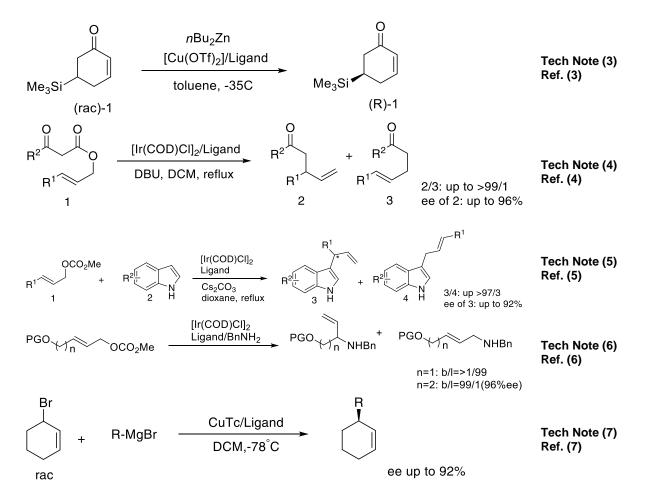
Technical Notes:

- 1. Amination Studies were conducted to determine possible intermediates in the highly enantioselective, iridium-catalyzed amination and etherification of allylic carbonates, and these studies revealed that cyclometalation of this phosphoramidite ligand is likely to generate the active catalyst.
- Cyclization Enantioselective cyclization of aromatic ketimines containing alkenyl groups tethered at the meta position of an imine directing group has been achieved using 5 mol% [RhCl(coe)2]2 and 15 mol% of this (S)-BINOL-derived phosphoramidite ligand.
- 3. Kinetic Resolution A variety of substituted 2-cyclohexenones, such as (R)-1, is obtained enantiomerically pure by employing the chiral copper-phosphoramidite complex [Cu(OTf)2L*] as a highly efficient catalyst for their kinetic resolution.
- Decarboxylative Alkylation [Ir(COD)CI]2/phosphoramidite ligand was found to be an efficient catalytic system for the highly regio- and enantioselective decarboxylative alkylation of γ-substituted allyl βketocarboxylates.
- 5. Friedel-Crafts Alkylation Highly regio- and enantioselective Ir-catalyzed Friedel-Crafts type allylic alkylation of indoles have been realized using [Ir(COD)CI]2/phosphoramidite ligand.
- Allylic Alkylation Ir-catalyzed allylic aminations of (E)-4-benzyloxy-2-butenyl methyl carbonate with benzylamine using Feringa's (Sa,Sc,Sc)-phosphoramidite as a chiral ligand afforded linear-aminated achiral product N,O-dibenzyl-4-amino-2-buten-1-ol regioselectively
- 7. The first dynamic kinetic asymmetric transformation in copper catalyzed allylic alkylation is reported, with enantioselectivities up to 92%.



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