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Practical new methods for carbon-carbon bond formation will be described that enable the efficient preparation of amine containing compounds. Additions to *N-tert*-butanesulfinyl imines, enantioselective reactions catalyzed by *N*-sulfinyl ureas, and Rh(I)- and Rh(III)-catalyzed C-H bond functionalization will be presented with an emphasis on regio- and stereoselective entry to pharmaceutically relevant classes of chiral amines and nitrogen heterocycles. The utility of these methods will also be demonstrated with syntheses of bioactive natural products and drugs.

## Leading References:

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