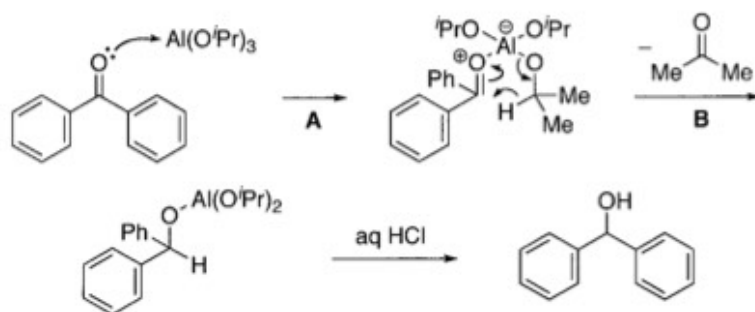
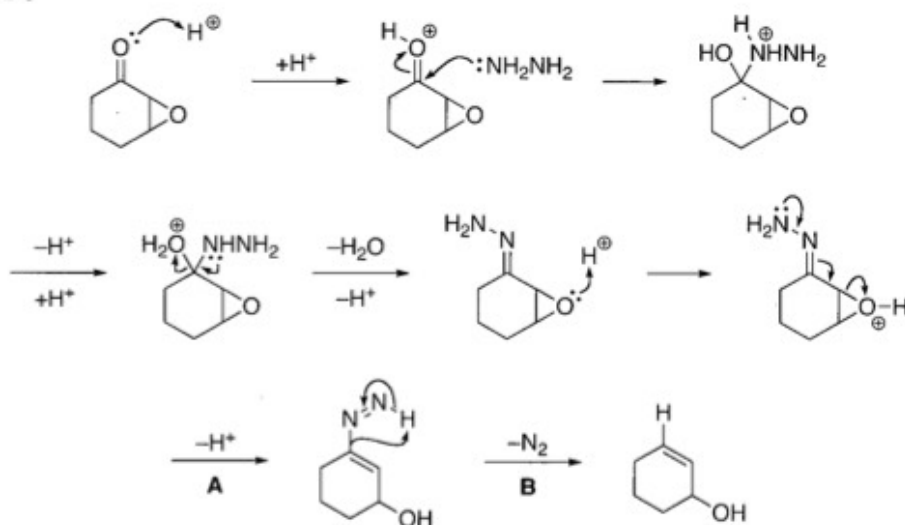


**B002**

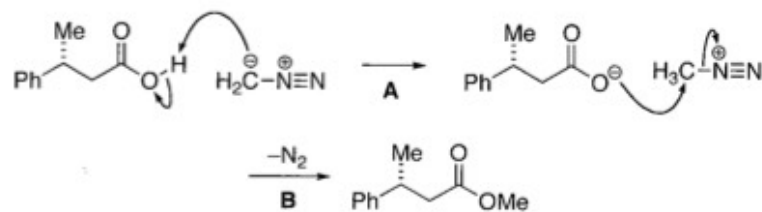
Wilds, A. L. *Org. React.* **1944**, 2, 178.

Meerwein-Ponndorf-Verley reduction. **A**: Formation of an ate complex. **B**: Hydride transfer via a six-membered transition state with formation of acetone.

**B003**

Wharton, P. S.; Bohlen, D. H. *J. Org. Chem.* **1961**, 26, 3615.

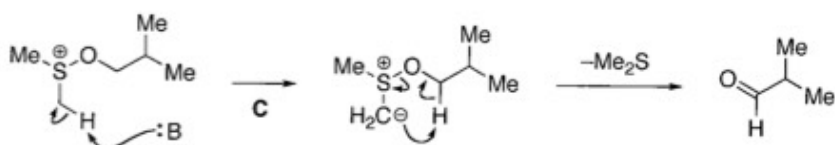
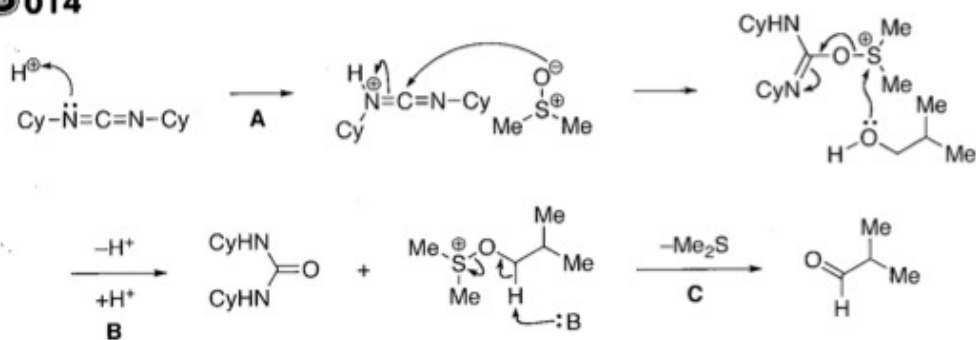
Wharton rearrangement. **A**: Cleavage of the epoxide helped by the nitrogen lone pair of the hydrazone. **B**: Loss of  $\text{N}_2$  (an extremely good leaving group).

**B010**

Black, T. H. *Aldrichimica Acta* **1983**, 16, 3.

**A**:  $\text{CH}_3\text{CO}_2\text{H} = 4.8$ ,  $\text{CH}_3\text{N}_2 = 10.2$ . **B**: The  $\text{S}_{\text{N}}2$  reaction occurs in a solvent cage.

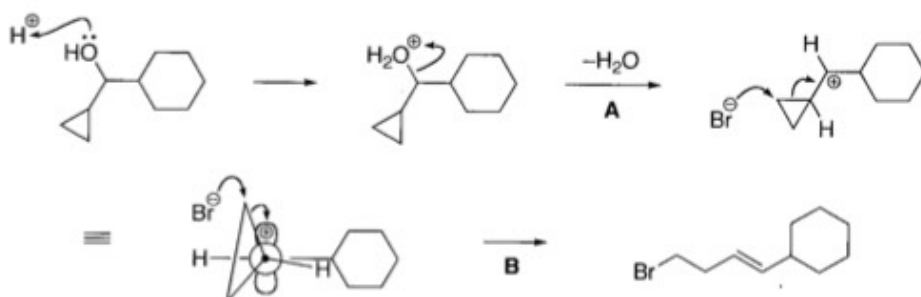
## B014



Tidwell, T. T. *Org. React.* **1990**, 39, 297.

Pfitzner-Moffatt oxidation. **A**: Activation of DCC by protonation. **B**: Nucleophilic substitution at the sulfur atom. **C**:  $\beta$ -Elimination of dimethyl sulfide might proceed either by 1) direct deprotonation with a base or 2) formation and collapse of a sulfur ylide.

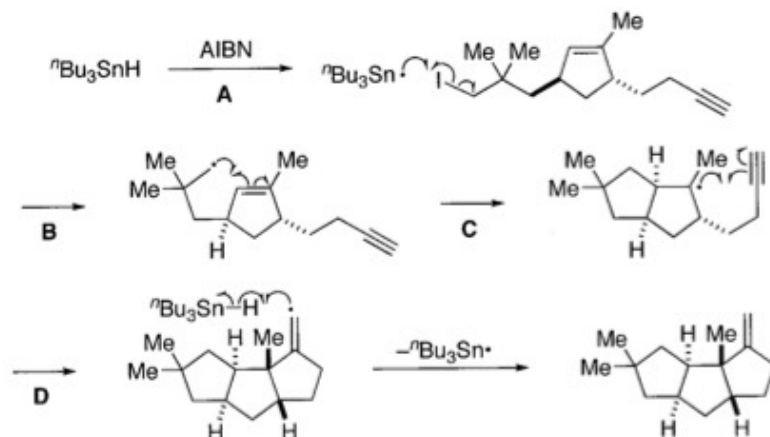
## B015



Ferreri, C.; Ambrosone, M. *Syn. Commun.* **1995**, 25, 3351.

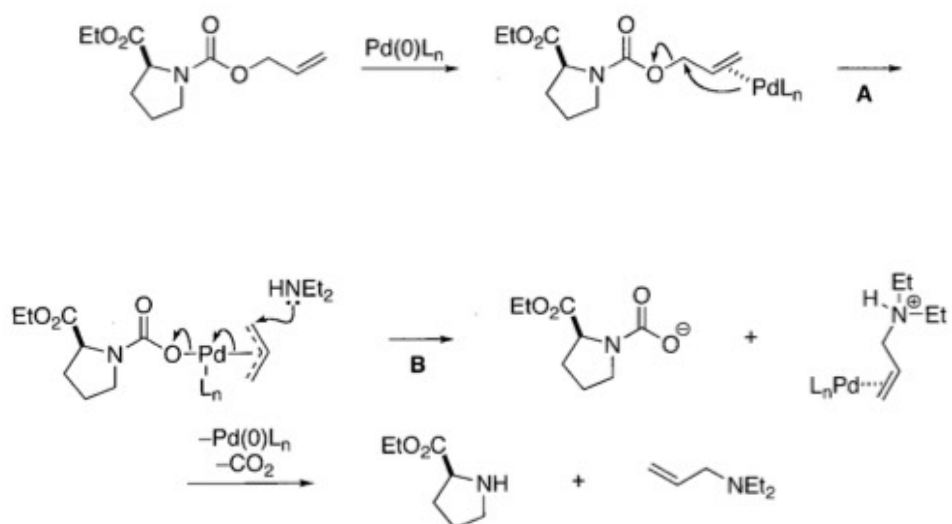
**A**: Generation of a carbocation stabilized by a cyclopropyl group. **B**: Cleavage of the cyclopropane ring occurs by avoiding the steric repulsion to form the *trans*-product.

## B017



Weinges, K.; Reichert, H.; Huber-Patz, U.; Irgartinger, H. *Liebigs Ann. Chem.* **1993**, 403.

**A**: Generation of a tin radical (ref A050). **B**: Attack on the iodide to initiate the radical chain reaction. **C**: 5-*exo*-trig Radical cyclization. **D**: 5-*exo*-dig Radical cyclization.

**B021**

Genet, J. P.; Blart, E.; Savignac, M.; Lemeune, S.; Lemaire-Audoire, S.; Bernard, J. M.  
*Synlett* **1993**, 680.

**A:** Formation of a  $\pi$ -allylpalladium complex. **B:** Attack of  $\text{Et}_2\text{NH}$  to the  $\pi$ -allyl complex.