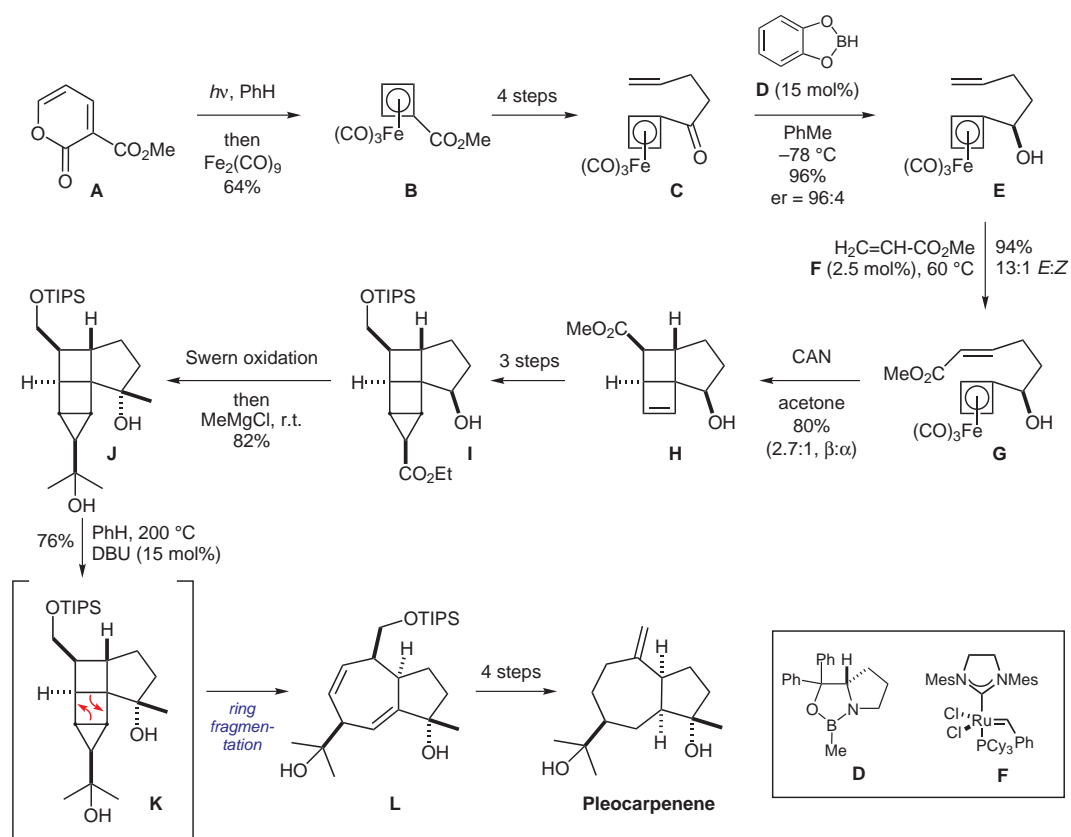


M. J. WILLIAMS, H. L. DEAK, M. L. SNAPPER\* (BOSTON COLLEGE, CHESNUT HILL, USA)  
 Intramolecular Cyclobutadiene Cycloaddition/Cyclopropanation/Thermal Rearrangement: An Effective Strategy for  
 the Asymmetric Syntheses of Pleocarpenene and Pleocarpenone  
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## Synthesis of Pleocarpenene



**Significance:** Pleocarpenene is a guaianene natural product. Snapper and co-workers used (1) an intramolecular cyclobutadiene cycloaddition reaction and (2) a thermal ring fragmentation reaction *en route* to the 5–7-fused ring system.

**Comment:** Asymmetric reduction of iron-cyclobutadiene complex **C** using the Corey oxazaborolidine gave enantioenriched **E** that underwent cross-metathesis giving **G**. Subsequent oxidation with cerium ammonium nitrate (CAN) gave **H** as the major diastereoisomer. Later, thermolysis of **J** in the presence of DBU afforded **L** as the product of ring fragmentation.

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Corey  
 oxazaborolidine

cross-metathesis

ring fragmentation

**SYNFACTS**  
*of the month*

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