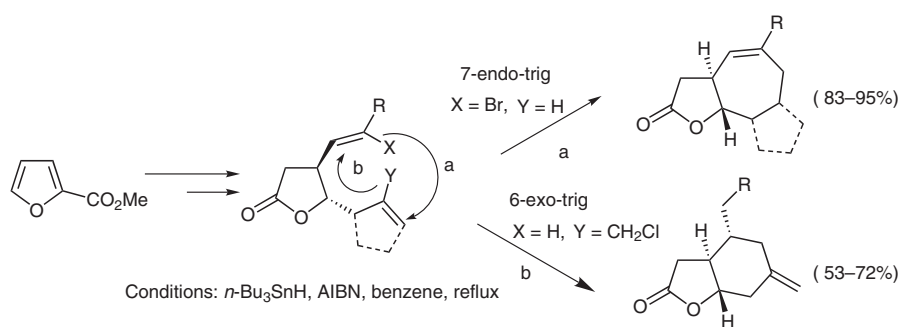


E. JEZEK, A. SCHALL, P. KREITMEIER, O. RESIER* (UNIVERSITY OF REGENSBURG, GERMANY)
Radical Cyclizations as Key Step for the Stereoselective Synthesis of Bi- and Tricyclic Stereoselective Sesquiterpene Lactones
Synlett **2005**, 915-918.

Radical-Mediated Synthesis of Sesquiterpene Lactones



Significance: Starting from inexpensive methyl 2-furoate, via the intermediate cyclopentenylbutenolide (4 steps, 26% overall yield), bi- and tricyclic γ -butyrolactone cores can be formed stereoselectively and in good yields via Y-substituent – directed radical cyclization. Using R = Br, CN, CO₂Et, COMe substitution, reactions proceed in acceptable yields.

Comment: *trans*-Fused seven- or six-membered rings are widespread frameworks in a variety of natural products such as guaianolides. This stereoselective strategy may provide new useful synthetic approaches to such derivatives and other natural products containing similar ring systems which to date have been rather limited.

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Category

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