



Significance: Luo and co-workers report the synthesis of (-)-zygadenine. Key to their approach is a stereoselective intramolecular Diels–Alder reaction, followed by a radical cyclization to construct the hexacyclic carbon skeleton.

Comment: Cyclization precursor F was prepared from commercially available enone A in five steps. Formation of a mixed anhydride with furan I enabled intramolecular Diels–Alder reaction to dikeione J. Treatment with AIBN and tributyltin hydride, followed by reduction furnished hexacyclic K.