Synthesis of Cassialoin

**Significance:** A significant development in the synthesis of cassialoin is the use of the α-ketol F as a selectively protected stereogenic anthrone surrogate in which the diastereotopic faces are differentiated thereby allowing control of the C-glycosidation reaction (F+G → H) at C10.

**Comment:** A regioisomer (17%) was formed in the cyclocondensation reaction of A and B. The α-ketol F was obtained as a single diastereoisomer. The epoxidation–reduction of glycal K (66%) installed the correct stereochemistry at C1’ and C2’.