**Total Synthesis of (−)-Flueggenine C**

**Significance:** Jeon and Han report the first total synthesis of the dimeric securinega alkaloid (−)-flueggenine C. The key dimerization step is enabled by a bio-inspired Rauhut–Currier reaction accelerated by the presence of an internal nucleophile. This reaction is pertinent in the biosynthesis of dimeric and oligomeric securinega alkaloids.

**Comment:** After efficiently synthesizing intermediate G in seven steps and 14% overall yield, the pivotal Rauhut–Currier dimerization and subsequent acetylation provided D in 74% yield. HWE olefination of F afforded unsaturated lactone G. The final ring closure was facilitated by S_N2 displacement of a mesylate culminating at the natural product.