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Total Synthesis of (±)-Alopecuridine and Its Biomimetic Transformation into (±)-Sieboldine A *Angew. Chem. Int. Ed.* **2011**, *50*, 3916-3919.

## Synthesis of (±)-Alopecuridine

**Significance:** (±)-Alopecuridine A is a Lycopodium alkaloid that is a biological precursor for the significantly cytotoxic (±)-sieboline A. The first total synthesis is achieved here in 13 steps, followed by the conversion into (±)-sieboline A that validates the biogenetic pathway.

**Comment:** Compounds **E** and **F** were obtained as an inseparable mixture of diastereoisomers that were subjected to semipinacol rearrangement conditions. This generated a mixture of diastereomers that could then be separated by chromatography to give the desired intermediate **G** in 45% yield. The tricyclic core **J** was synthesized with a samarium iodide mediated coupling.

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Category

Synthesis of Natural Products and Potential Drugs

**Key words** 

alopecuridine
pinacol coupling
semipinacol
rearrangement

