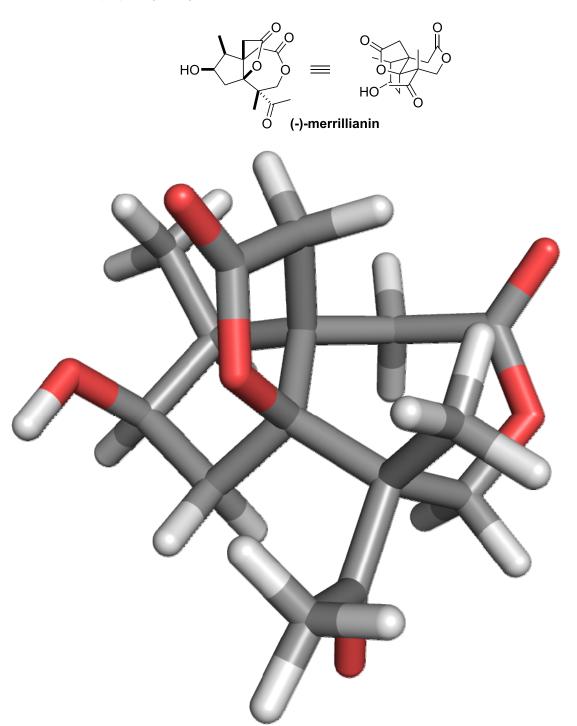
Please propose your synthetic route to merrillianin from a commercial compound.



Synthetic Plan of (-)-Merrillianin

$$HO = \begin{pmatrix} 15 & 10 & 11 & 0 & 0 \\ 15 & 10 & 11 & 0 & 0 \\ 10 & 13 & 0 & 0 & 0 \\ 13 & 0 & 0 & 0 & 0 \\ 13 & 0 & 0 & 0 & 0 \\ 13 & 0 & 0 & 0 & 0 \\ 13 & 0 & 0 & 0 & 0 \\ 13 & 0 & 0 & 0 & 0 \\ 14 & 12 & 0 & 0 & 0 \\ 15 & 0 & 0 & 0 & 0 \\ 16 & 0 & 0 & 0 & 0 \\ 17 & 0 & 0 & 0 & 0 \\ 18 & 0 & 0 & 0 & 0 \\ 19 & 0 & 0 & 0 & 0 \\ 10 & 0 & 0 & 0$$

- **·Isolation**: Isolated from the pericarps of *Illicium merrillianumis* along with (6R)-pseudomajucin, (6R)-pseudomajucinone, (6R)-pseudomajucin, anisatin, anislactone B, cycloparvifloralone, 3α -hydroxycycloparvifloralone.
- J.-M. Huang, C.-S. Yang, M. Kondo, K. Nakade, H. Takahashi, S. Takaoka, Y. Fukuyama. *Tetrahedron* **2002**, *58*, 6937.
- ·Bioactivities: Merrillianin does not have neurotrophic nor toxic effects on rat cortical neurons at 1mM.
- •Structural features: Sesquiterpene having an unusual chemical structure consisting of a cyclopentane fused to a five-membered ring lactone and a seven-membered ring lactone in the propeller form.

The absolute stereochemistry of (6R)-pseudomajucin was determined by the X-ray crystallographic analysis, but that of merrillianin has yet to be determined directly.

·Total synthesis

No report in peer-reviewed articles.

One total synthesis has been reported in a doctoral dissertation: Takashi Iizumi, Ph.D. Thesis, the Tokyo University of Science, 2012 (Shiina group).

(6R)-pseudomajucin

HO $\stackrel{\circ}{\longrightarrow}$ $\stackrel{\longrightarrow}{\longrightarrow}$ $\stackrel{\circ}{\longrightarrow}$ \stackrel

(6R)-pseudomajucinone

(6S)-pseudomajucin

anisatin

cycloparvifloralone: X = H 3α-cycloparvifloralone: X = OH

-1-