

Studies on Alkaloid and Terpenoid Synthesis

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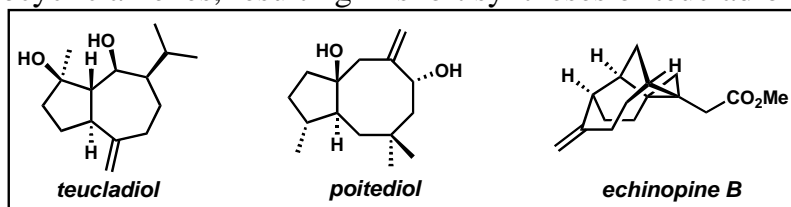
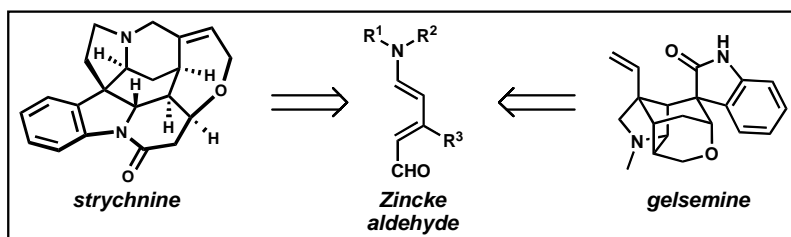
Time: 16:30 - 18:00

**Venue: West Seminar Room, 1F. Faculty of Pharmaceutical Sciences,
the University of Tokyo**



Our research group has been engaged in the synthesis of complex alkaloids and terpenoid natural products. I will describe our efforts toward *Strychnos* and *Gelsemium* alkaloids featuring the strategic use of Zincke aldehydes, which derive from the ring-opening aminolysis of pyridinium salts. Some fascinating and unexpected discoveries from this research program will also be presented.

In the second part of my talk, I will describe our efforts toward several sesquiterpenoid natural products, culminating in our efforts toward the echinopines. These studies inspired the use of the ring-closing metathesis of allylsilanes as a means to access ring structures bearing exocyclic alkenes, resulting in short syntheses of teucladiol and poitediol. Ultimately, our synthesis of the echinopines took advantage of a late-stage enyne cycloisomerization reaction to build its tetracyclic architecture.



**Organizer: GCOE Program Center for Medical System Innovation through Multidisciplinary Integration,
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