Total synthesis of ()-Flueggenine C



An Accelerated Intermolecular Rauhut-Currier Reaction Enables the ()-Flueggenine C

Jeon, S.; Han, S. J. Am Chem. Soc, 2017, 139, 6302–6305.

- Securinega alkaloids consisting of more than 70 natural products and are known since 1956
- Recent isolation of bioactive natural products from Flueggea virosa enabled the isolation of various dimeric and oligomeric alkaloids expanding its structural repertoire
- The biosynthesis of compounds **4** and **5** was reported using a self-catalyzed Baylis-Hilman reaction
- flueggenine A (4) showed modest cytotoxicity against the P-388 tumor cell line
- Flueggenine D and fluevirosinine B exhibited promising anti-HIV activities
- The first asymmetric total synthesis of flueggenine C (6), a C,C-linked dimeric securinega Alkaloid was achieved in this work.

Natural products containing the DMOA (1) core



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Retrosynthetic Analysis of Flueggenine C (6)

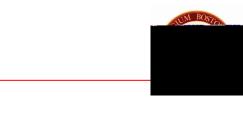


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Possible modes of reactivity in Rauhut-Currier reactions



Conventional Intermolecular RC Reaction









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