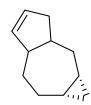
Enantioselective Total Synthesis of ()-Euphorikanin A

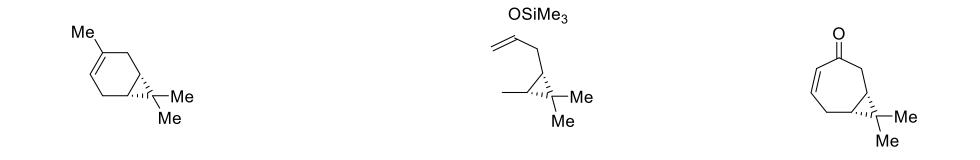
Moritz J. Classen, Markus N. A. Böcker, Remo Roth, Willi M. Amberg, and Erick M. Carreira* *J. Am. Chem. Soc.* 2021, 143, 8261-8265.

- First total synthesis of (+)-euphorikanin A, an ingenane-derived natural product
- Isolated in 2016 and identified as a novel diterpenoid from the roots of *Euphorbia kansui*, commonly known as kansui
- Extracts of the root have been widely used in traditional Chinese medicine. (+)-euphorikanin A has been shown to exhibit cytotoxicity against two human tumor cell lines (NCI-446 and HeLa)
- Features an unprecedented 5/6/7/3-fused tetracyclic skeleton
- " Prepared in 19 steps from (+)-3-carene
- Key step is an Sml2-mediated ketyl-enoate cyclization cascade

Ricky Alvarado November 16th, 2021

Retrosynthesis





Ozonolysis and formation of silyl ketene acetal

Acid-cyclization and dehydration

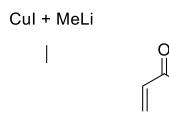
OSiMe**.**

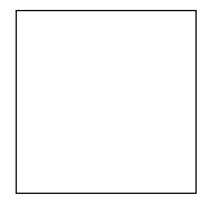
__Me Me

(5) Me₂CuLi Et



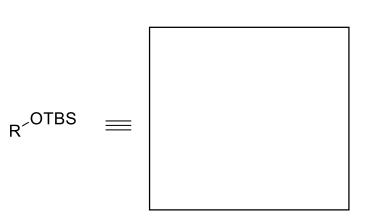
R

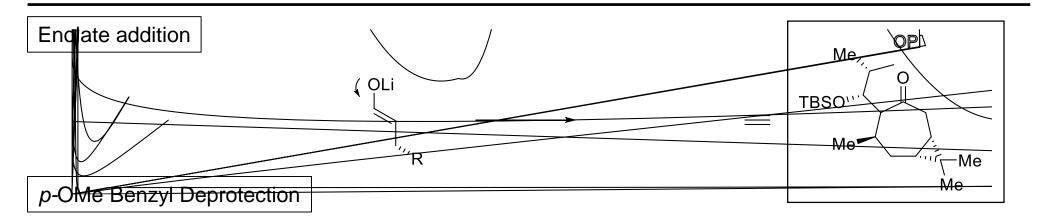


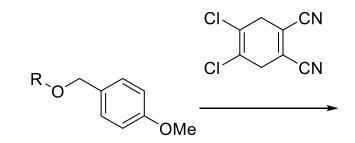


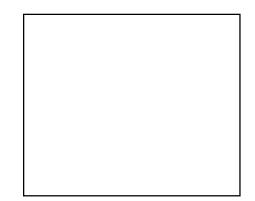
Alcohol protection with TBSOTf

Si



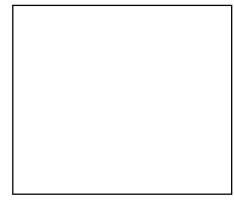






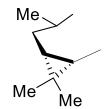
Parikh-Doering Oxidation

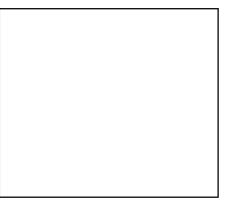




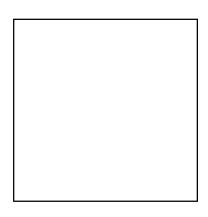
Me		

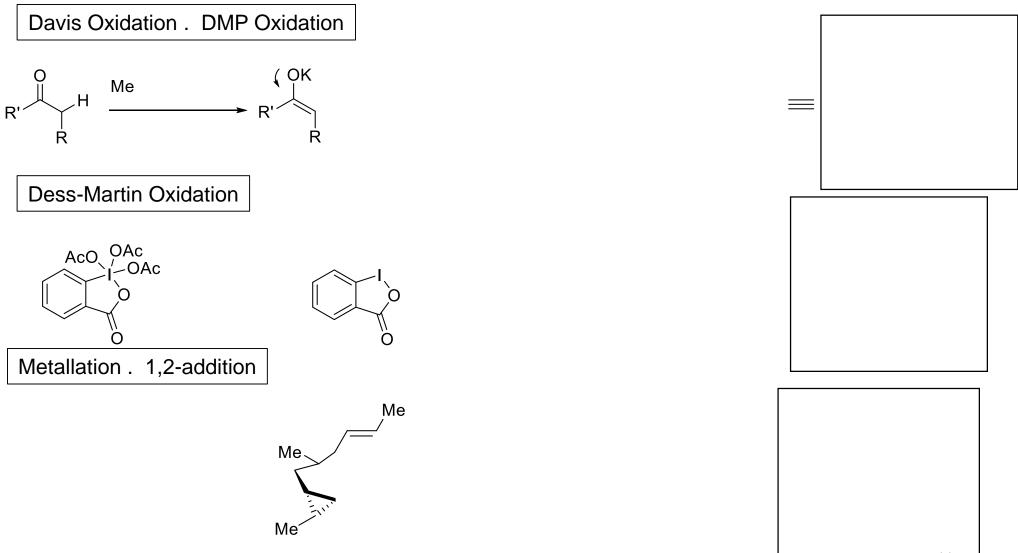
Umpolung Cyclization via Reduction with Sml2. Undesired Stereoisomer





TBS Deprotection





(19) t