

#### Stereoselective Total Syntheses of Guanacastepenes N and O

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S Supporting Information

**ABSTRACT:** Total syntheses of  $(\pm)$ -guanacastepenes N and O were accomplished with 2-cycloheptenone as starting material. The six- and five-membered rings of the core [5, 7, 6] ring skeleton were constructed with an intramolecular Diels—Alder reaction and  $\alpha$ -carbonyl radical cyclization. The quaternary centers and their stereochemistry were established with sequential Cu(I)-mediated conjugate additions. A sequence with dihydroxylation, conjugate addition, and  $\beta$ -

elimination was devised to incorporate all oxygen functionalities at positions. The total synthesis is adaptable for the synthesis of enantiopure guanacastepenes N and O using chiral intermediate (R)-3-vinyl-2-cycloheptenol obtained from lipase-catalyzed kinetic resolution.

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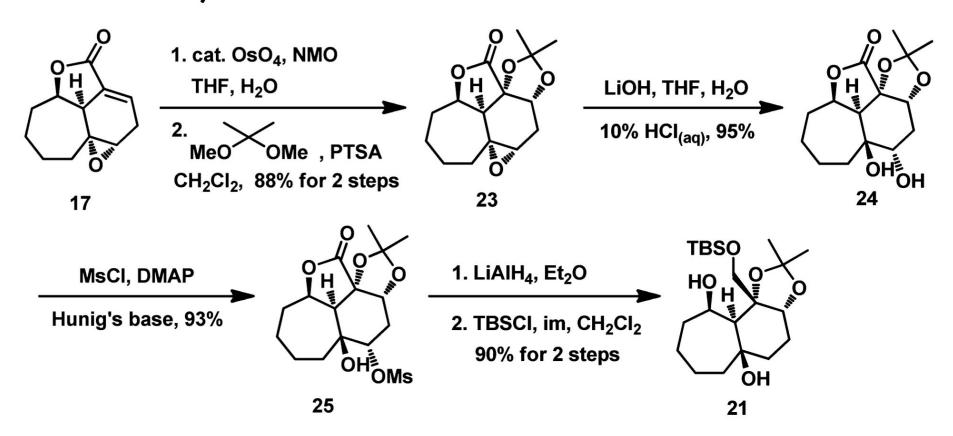
#### Scheme 1. Retrosynthesis of Guanacastepenes N and O

## Scheme 2. Synthesis of Compounds 17 and 18

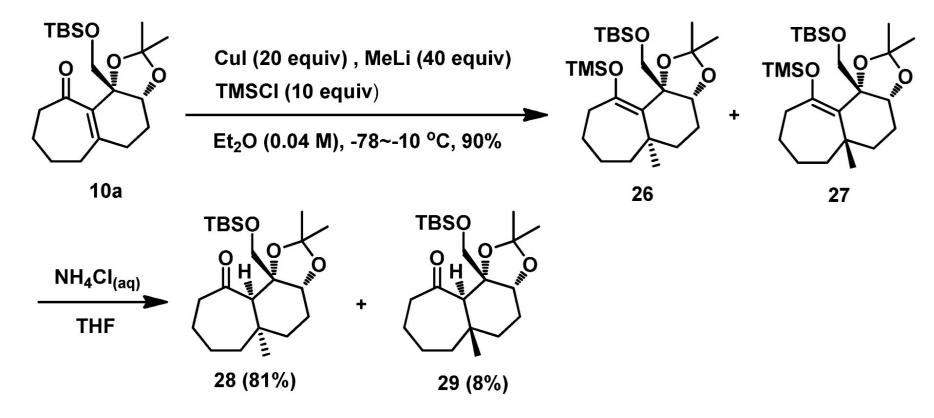
## Scheme 3. Synthesis of Enone 10a

10a

## Scheme 4. Synthesis of Diol 21



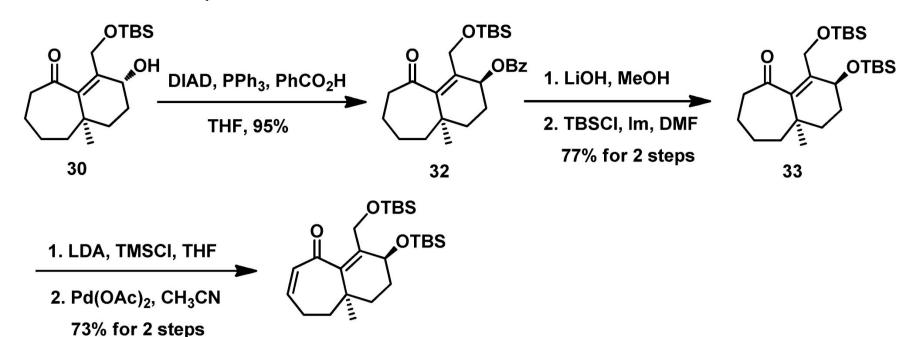
## Scheme 5. Synthesis of Compounds 28 and 29



# Scheme 6. Synthesis of Enones 30 and 31 via $\beta$ -Elimination Reaction

#### Scheme 7. Synthesis of Dienone 9

9



#### Scheme 8. Synthesis of Compounds 37a, 37b, and 38

## Scheme 9. Synthesis of Furan 42

## Scheme 10. Synthesis of Lactones 45 and 46

2. DMDO, CH<sub>2</sub>Cl<sub>2</sub> 62% for 2 steps

#### Scheme 11. Synthesis of Guanacastepenes N and O

## Scheme 12. Synthesis of Chiral Dienol (R)-16

