## Enantioselective Total Synthesis of (-)-Ambiguine







Significance: Isolated from the cultured cyanobacterium Fischerella ambigua in 2010, ambiguine P belongs to the hapalindole alkaloid family. The pentacyclic natural product contains a seven-membered ring and an embedded indole, which pose significant synthetic challenges. The authors report an enantioselective total synthesis of the target employing a stepwise [4+3]-cycloaddition strategy.

Comment: Ketone B, accessible from limonene (A), was transformed into enol ether $\mathbf{D}$ in four steps. A net [4+3] cycloaddition of this diene with indole $\mathbf{E}$ and subsequent intramolecular Friedel-Crafts reaction installed the target's pentacyclic core. Functional group manipulations furnished diene $\mathbf{I}$, which was transformed into the natural product by NBSmediated regioselective oxidation.

## Category

Synthesis of Natural

## Key words

## Friedel-Crafts

reaction
[4+3] cycloaddition
ambiguine $P$
hapalindolealkaloids


