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 Natural Product-Inspired Cascade Synthesis Yields Modulators of Centrosome Integrity
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Indoloquinolizines from a One-Pot, 12-Step, Three-Component Cascade Reaction

Category

Synthesis of Heterocycles

Key words

indoloquinolizines

multicomponent reaction

aza-Claisen rearrangement

Pictet–Spengler cyclization

Michael reaction

retro-Michael reaction

centrocuntins

SYNFACT
of the month

Significance: Reported is the one-pot synthesis of highly functionalized indoloquinolizines **16**, containing a tetracyclic framework found in numerous indole alkaloid natural products. The complex fused heterocycles are generated in isolated yields of 22–91% from commodity materials by a multicomponent reaction (MCR). During the 12-step cascade, two C–C bonds and two C–N bonds are formed within tens of minutes under operationally simple conditions. The cascade was unexpected and required impressive synthetic detective work, logically described and well supported by literature citation and by X-ray structures of intermediates. The 26 reported examples enabled efforts to identify small-molecule modulators of mitosis in HeLa cell lines as well as tool compounds to probe the chemical biology.

Comment: This rapid, complex heterocycle synthesis showcases the power of organic chemistry in addressing important problems in chemical biology. Though the heterocyclic scaffold produced is restricted in scope, the indoloquinolizines are highly functionalized and well poised for diversity-oriented synthesis. Judicious choice of starting materials may quickly afford complex, alkaloid-like templates substituted with esters, halogens, and oxygenation for flexible, orthogonal derivation and library production. This work led to novel compounds, termed centrocuntins, which have been shown to modulate mitotic spindle assembly and may lead to anti-proliferative therapeutics in medicinal oncology research.

Review: J. D. Sunderhaus, S. F. Martin *Chem. Eur. J.* **2009**, *15*, 1300–1308.

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