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Efficient synthesis of eudistomin U and evaluation of its cytotoxicity

Chad M. Roggero, Jennifer M. Giulietti, Seann P. Mulcahy

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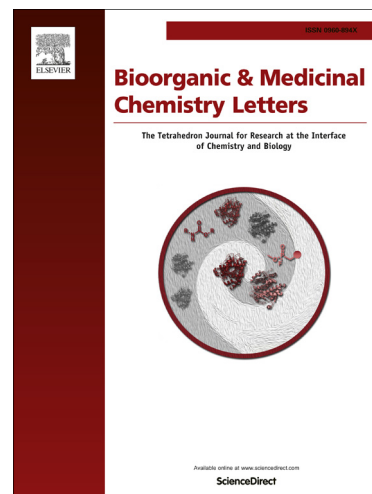
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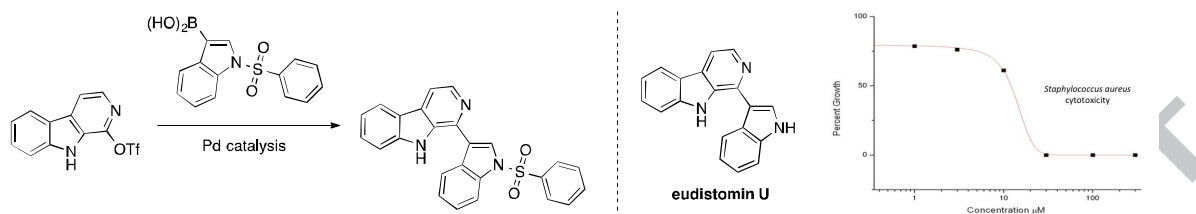
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Graphical Abstract:**Efficient synthesis of eudistomin U and evaluation of its cytotoxicity**

Chad M. Roggero, Jennifer M. Giulietti, and Seann P. Mulcahy*

Providence College, Department of Chemistry and Biochemistry, 1 Cunningham Square, Providence, RI 02918, USA

*Corresponding author. Tel.: +1-401-865-1280; fax: +1-401-865-1438; e-mail: smulcahy@providence.edu

Abstract: Eudistomin U is a member of a subclass of naturally occurring indole alkaloids known as β -carboline. These molecules are reported to have diverse biological activity and high binding affinity to DNA, which make them attractive targets for total synthesis. We describe an efficient, five-step synthesis of eudistomin U by employing two key reactions: a Bischler-Napieralski cyclization and a Suzuki cross coupling. We also describe the cytotoxicity of eudistomin U against various cancer cell lines and human pathogens, in which we observed potent antibacterial activity against Gram-positive bacteria.

Keywords: β -carboline, Suzuki cross coupling, eudistomin U, antibacterial, palladium catalysis

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