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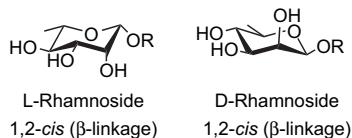
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REPORT

Challenges in the stereocontrolled syntheses of β -rhamnosides

El Sayed H. El Ashry*, Nagwa Rashed, El Sayed I. Ibrahim

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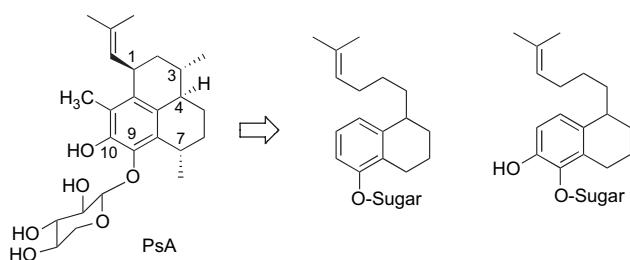
The stereocontrolled syntheses of the β (1,2-cis)-rhamnosides via inter- and intramolecular glycosidation reactions, reductive cleavage of 4,6-acetals, inversion of α -rhamnosidic linkages and modification of β -mannosides have been reviewed. The report contains 88 references.

ARTICLES

Synthesis and evaluation of the bioactivity of simplified analogs of the seco-pseudopterosins; progress toward determining a pharmacophore

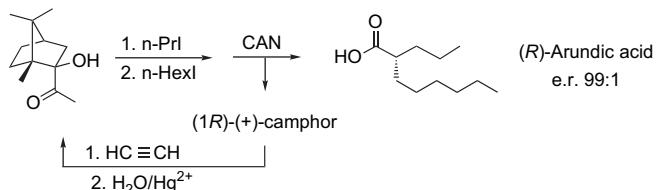
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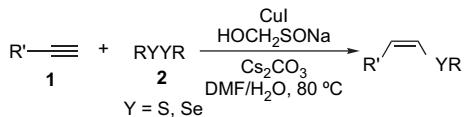
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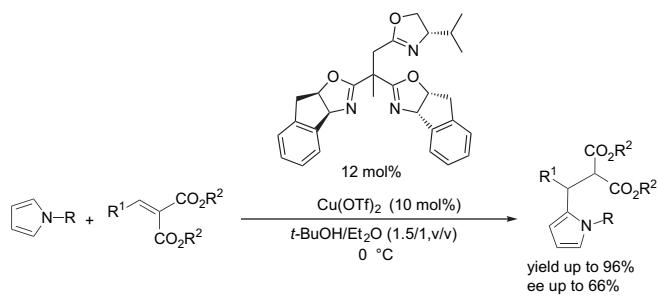
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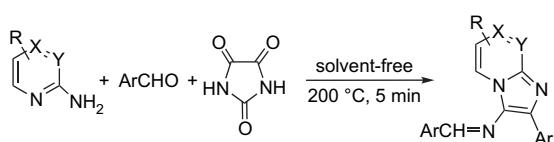
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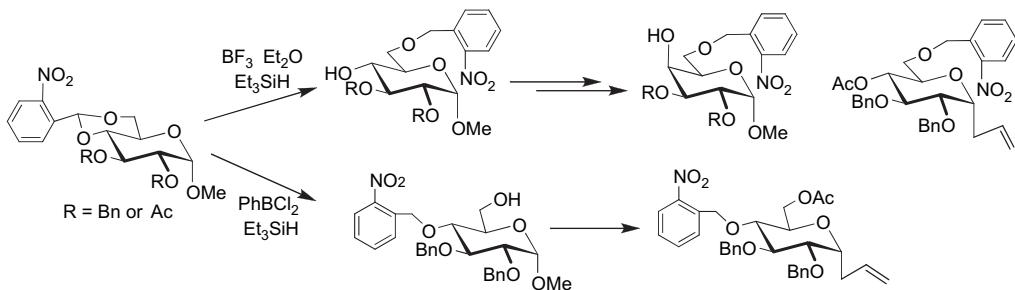
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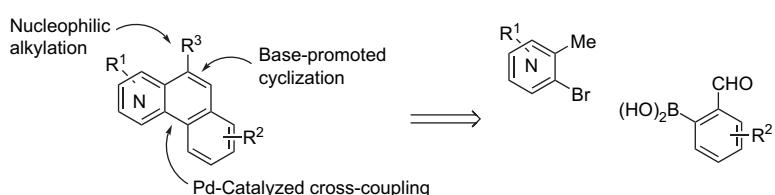


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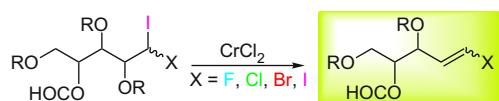
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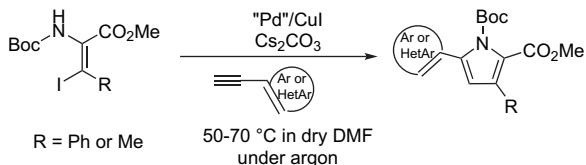
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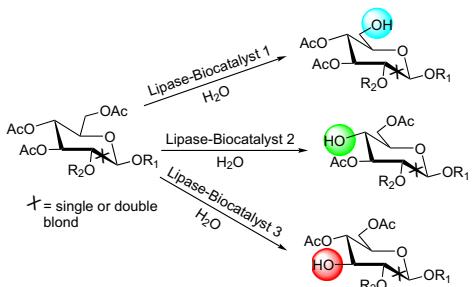
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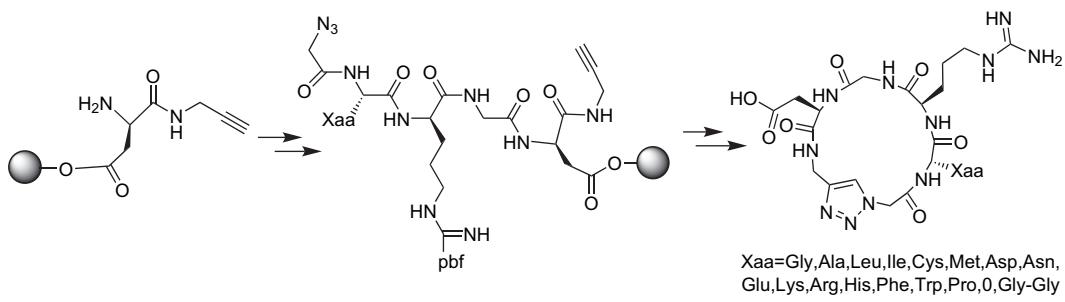
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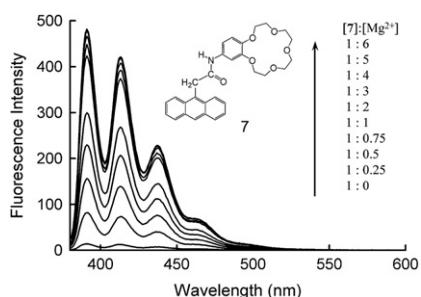
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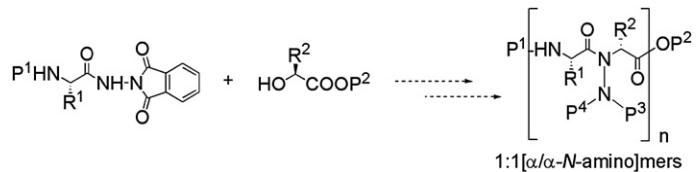
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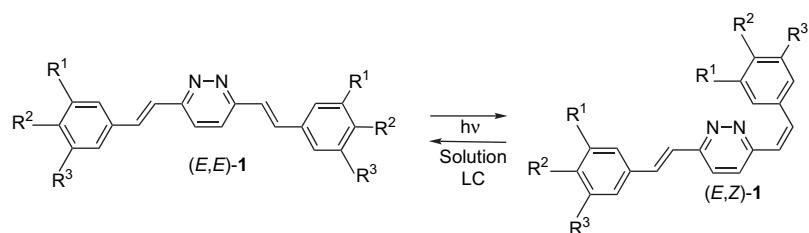
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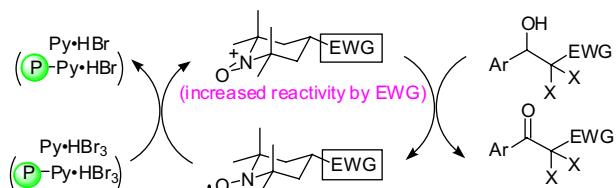
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Zhen-Wu Mei, Takumi Omote, Mounir Mansour, Hiroyuki Kawafuchi, Yutaka Takaguchi, Anny Jutand*, Sadao Tsuboi, Tsutomu Inokuchi*



A combination of Py-HBr₃ as a co-oxidant and the electronically activated TEMPO as a recyclable catalyst is useful for oxidation of not only common alcohols, but also of the electron-deficient secondary alcohols such as ArCH(OH)CFCl₂.



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 ⁱ† Supplementary data available via ScienceDirect



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