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Diastereoselective Anomeric C(sp^3)–H Cyclization Towards the Design of New Cyclophane-Braced Glycopeptides
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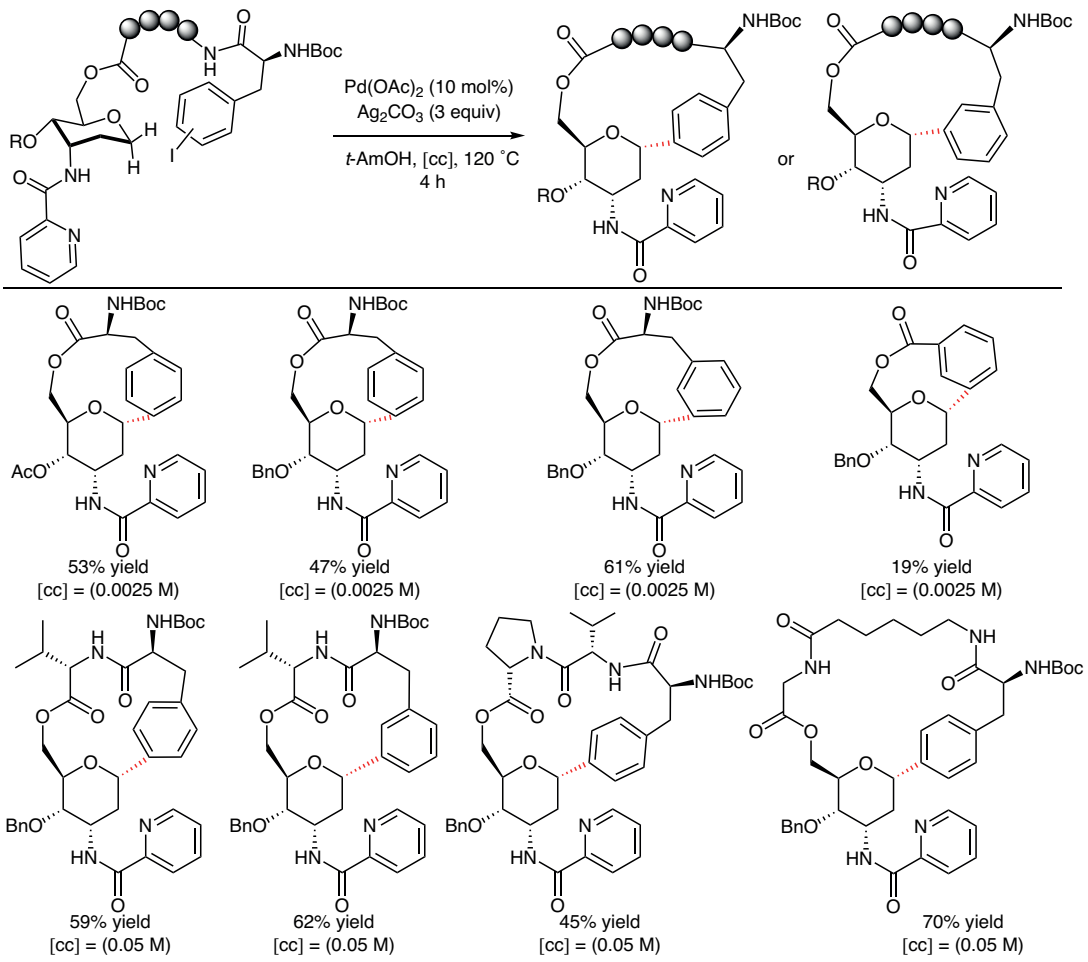
Palladium-Catalyzed Late-Stage Macrocyclization for the Synthesis of Cyclophane-Braced Glycopeptides



Significance: Macrocyclic peptides have gained tremendous attention in peptide drug discovery. In this article, the authors have developed a palladium-catalyzed macrocyclization by selective C(sp^3)–H arylation of the anomeric bonds for the synthesis of cyclophane-braced glycopeptides.

Comment: A series of macrocyclic cyclophane-braced glycopeptides has been synthesized in good to moderate yields by a palladium-catalyzed site-selective anomeric C(sp^3)–H cyclization approach. This protocol showcases the efficiency of a palladium-catalyzed C–H activation strategy for the synthesis of unnatural cyclic peptides.

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Category
Peptide Chemistry

Key words
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glycopeptides

Synfact of the Month