ELSEVIER

Tetrahedron

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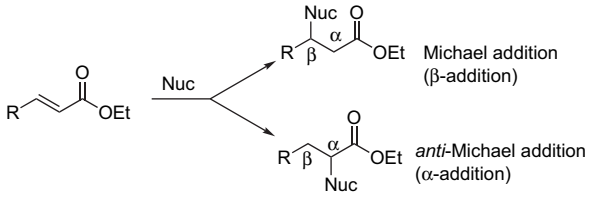
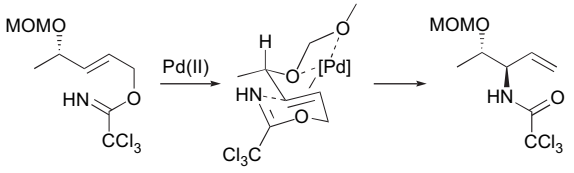
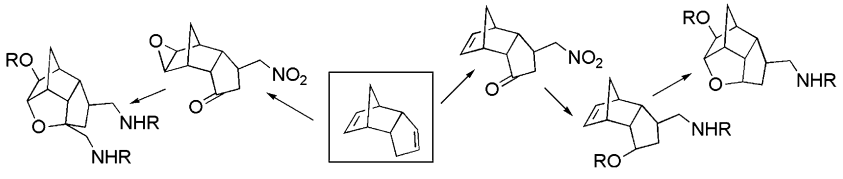
Contents

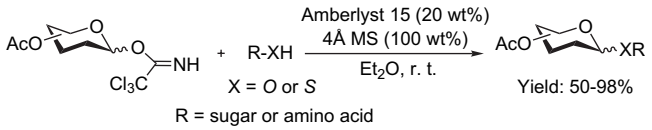
REPORT

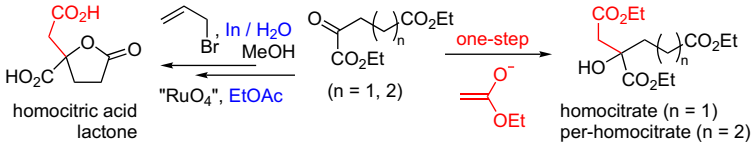


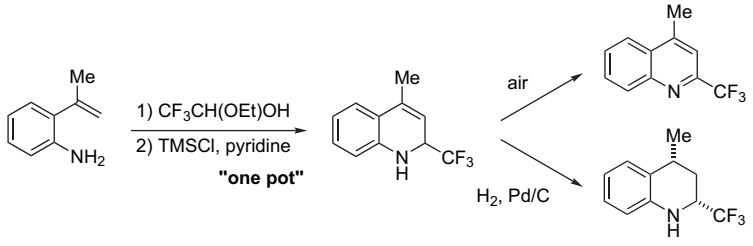
ARTICLES



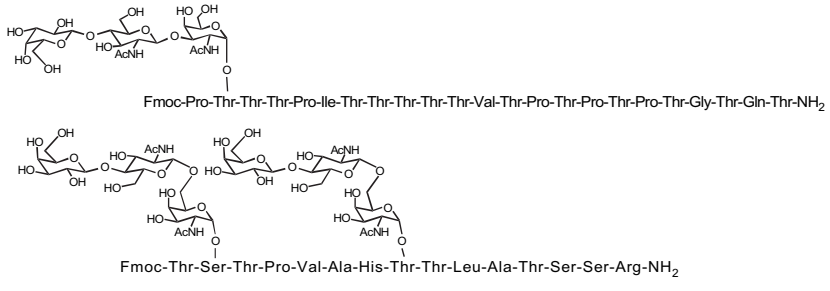


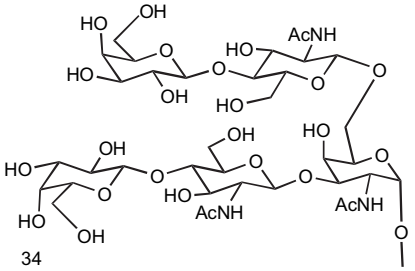






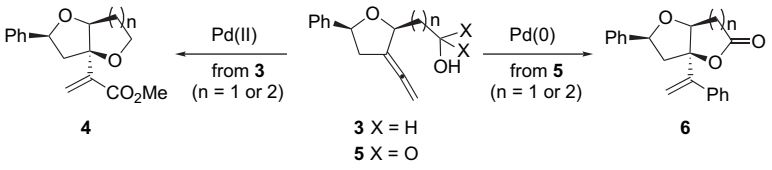
Benzyl- and benzylidene-protected trisaccharide-Ser/Thr derivatives of core 3 and core 6 *O*-glycans were synthesized as the building blocks suitable for the solid-phase synthesis. Syntheses of MUC2 and MUC6 glycopeptides were demonstrated.



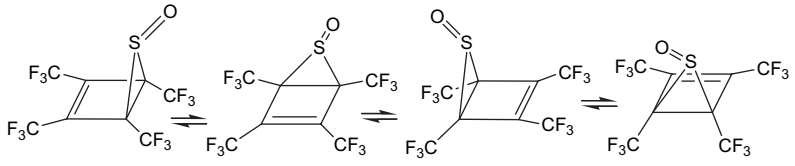


Synthesis of core 4 *O*-glycan and solid-phase synthesis of an *O*-glycan analog of the emmprin (Gly^{34}-Gly^{58}) segment are described. Microwave accelerated Zn/AcOH reduction of trichloroacetamides to acetamides.

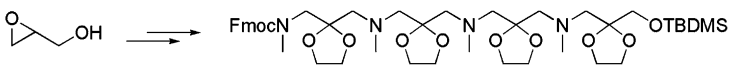
34 58
H-Gly-Ser-Lys-Ile-Leu-Leu-Thr-Cys-Ser-Leu-Thr-Asp-Ser-Ala-Thr-Glu-Val-Thr-Gly-His-Arg-Trp-Leu-Lys-Gly-$SCH_2CH_2CONH_2$

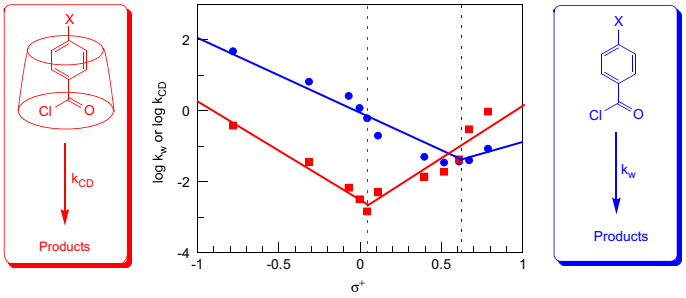


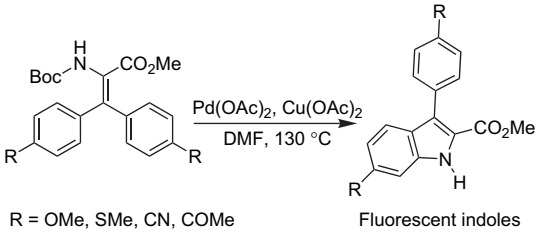
Prins-type cyclization followed by palladium-catalyzed cyclization provides a very good methodology for the synthesis of various sterically congested bicyclic tetrahydrofurans, which are very stereoselective as well.

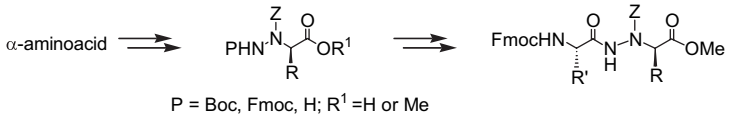


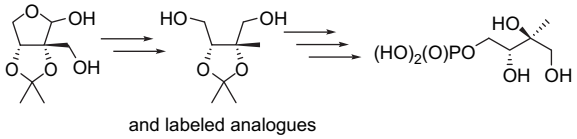
The mechanism of this *walk rearrangement* was theoretically studied. The definition of *pseudopericyclic* reaction was based on the experimental behavior of this compound, which undergoes an extraordinarily facile automerization.



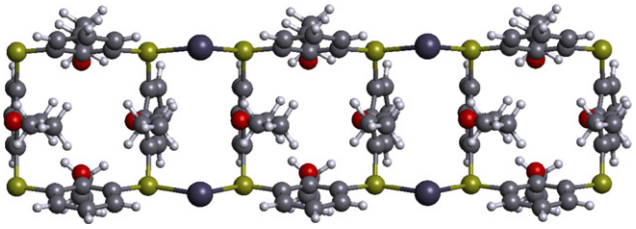


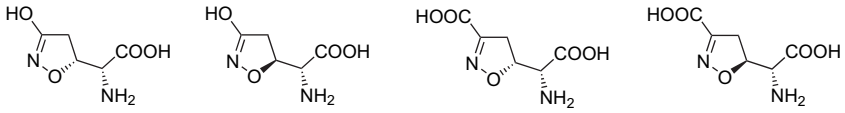


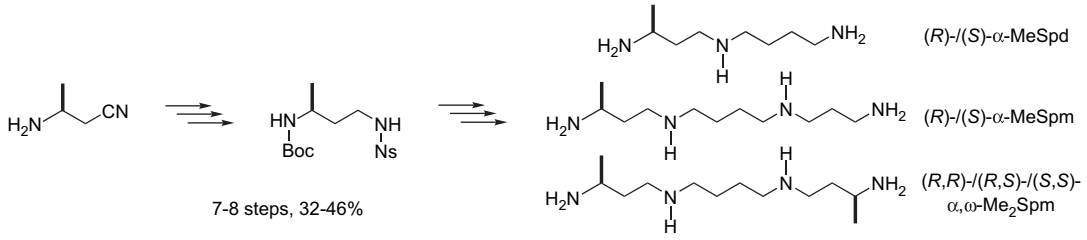


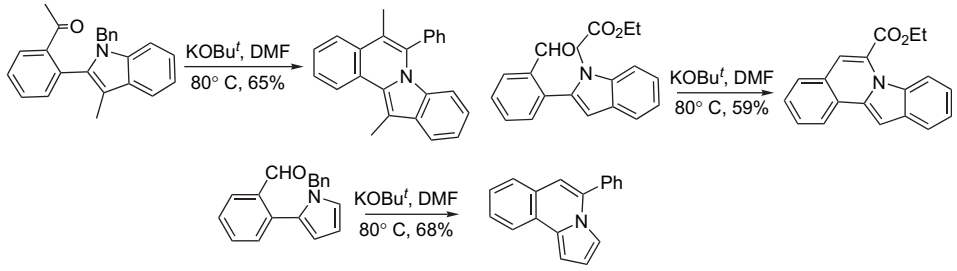


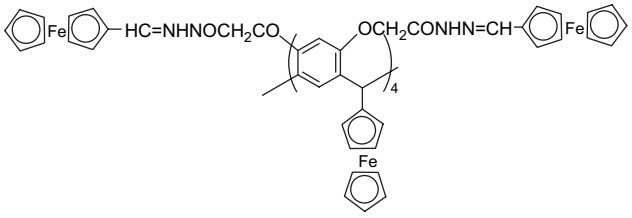
A new synthetic approach toward 2-*C*-methyl-D-erythritol 4-phosphate (MEP), a key intermediate in the mevalonate-independent biosynthetic pathway for isoprenoids, and deuterated analogues of its precursor, 2-*C*-methyl-D-erythritol acetonide, is described. This procedure uses 2-*C*-methyl-D-erythrose acetonide as starting material and delivers, through a mono-protection strategy, the target compounds in a short way and in high yield.

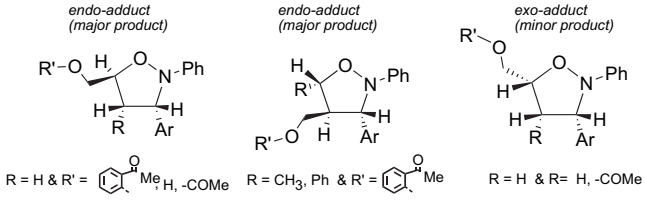


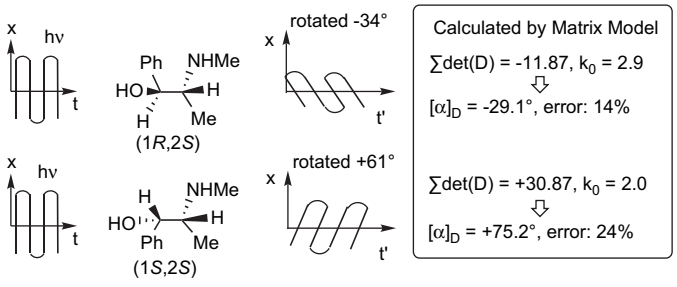


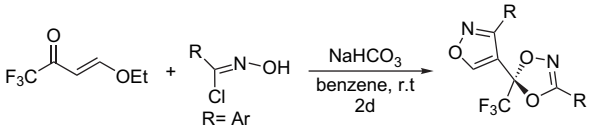




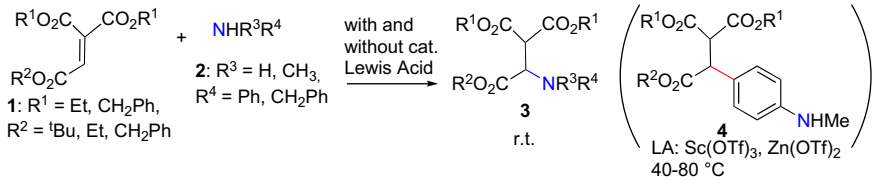




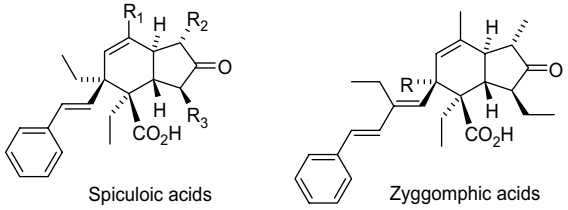


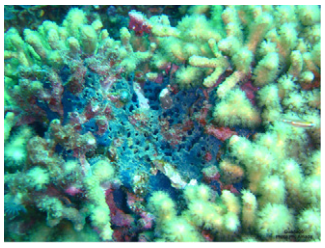


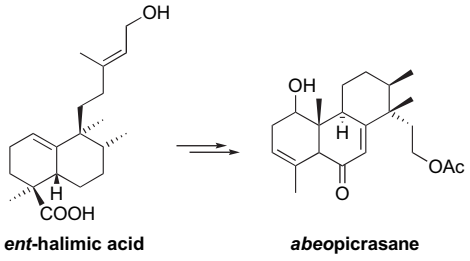
1,3-Dipolar cycloaddition reaction of trifluoromethyl substituted α,β-unsaturated ketone with nitrile oxides was examined and a series of bicycloaddition products were obtained.

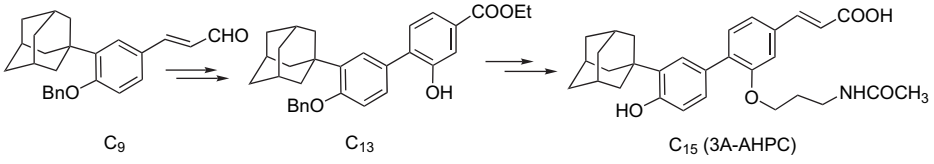


The reaction of ethenetricarboxylates **1** and aromatic amines with and without catalytic Lewis acids gave amine adducts or aromatic substitution products, depending on the conditions.




Plakortis zyggompha





OTHER CONTENT

*Corresponding author
ⓘ+ Supplementary data available via ScienceDirect

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