α-C-Glycosides via syn Opening of 1,2-Anhydro Sugars with Organozinc Compounds in Toluene/n-Bu₂O

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The diastereoselective addition of organozinc species to 1,2-anhydro sugars in toluene/*n*-dibutyl ether solvent is reported.¹ Compared to the existing methods, the reaction proceeds at 0°C and only a slight excess of nucleophile is required to achieve good yields. Scope was assessed with different *O*-protected glycals along with various nucleophiles (aryl, alkynyl). This methodology was applied to the synthesis of the -anomer of Canagliflozin.

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