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Cu-Catalyzed Tandem Reactions of Fluorinated Alkynes with Sulfonyl Azides En Route to 2-Trifluoromethylquinolines


Synthesis of 2-Trifluoromethylquinolines via a [2+3] Cycloaddition

Significance: Reported is the synthesis of 2-trifluoromethyl quinolines 3 and 4 via a copper-catalyzed Huisgen [2+3]-cycloaddition reaction of sulfonyl azides 2 with terminal alkynes 1. The starting material 1 was obtained by the reaction of 2,2,2-trifluoro-N-phenylacetimidoyl chloride with [(trimethylsilyl)ethynyl]magnesium bromide, followed by trimethylsilyl deprotection. Optimum reaction conditions, as presented in the scheme, were used to obtain a variety of quinolines (3). Alkynes 1 containing electron-withdrawing groups as R1 resulted in poor yield; those with electron-donating groups as R1 resulted in moderate to good yields. Compound 3 was isolated as a mixture of isomers (C-5/C-7 from 1:1.7 to 1:1.8) when a meta substituent as R1 on 1 was used. The method was also extended to the synthesis of 2-perfluoroalkylated quinolines (R2 = CF3 (16 examples), C2F5, C3F7, CF2Cl, CF2Br, CF2H (trace yield))

Comment: The importance of the trifluoromethyl group in drug discovery programs has resulted in the development of new and improved methods for the introduction of CF3 groups onto (hetero)arenes. Although recent focus has been on the transition-metal-mediated cross-couplings and direct C–H trifluoromethylations (see Reviews below), methods such as the present one provide competitive alternatives, especially when suitable starting material can be obtained easily. The current method can tolerate a wide range of functional groups, but the yield is lower with electron-poor alkynes. Also, this method was not tested on internal alkynes.


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