Tetra Butyl Ammonium Fluoride: TBAF

Compiled by Majeti Bharat Kumar

Mr. Bharat received his BSc (Bachelor of Science) degree from Osmania University, India and his MSc degree in chemistry from the University of Hyderabad, India. He is currently pursuing a PhD, carrying out research with Dr. Arabinda Chaudhuri in the area of design, synthesis and transfection biology of novel cationic transfection lipids for use in non-viral gene therapy at Indian Institute of Chemical Technology, Hyderabad, India.
E-mail: mbkumar@iict.ap.nic.in

Tetra butyl ammonium fluoride (TBAF) has been used widely as a reagent for the efficient cleavage of various silyl protecting groups from O-silylated nucleosides,2,3 and pyrophosphates,3 as well as N-silyl,4 and S-silyl derivatives.5 These reactions are carried out under very mild conditions in excellent yields.

Preparation: Aqueous HF is passed through an amberlite IRA 410 OH column, followed by an aqueous solution of tetra-butylammonium bromide. After the resin is washed with H₂O, the combined H₂O fractions are repeatedly evaporated until no water is present. TBAF is collected as an oil in quantitative yield.

Abstracts

Silyl ethers can be converted to esters in one pot when they are treated with TBAF, followed by exposure to acyl chlorides6,7 or anhydride8 in the presence of a base.

The anions generated in situ by desilylation of silylacetylenes,9,10 allylsilanes,11–13 propargylsilanes14 and other silane derivatives can undergo nucleophilic addition to ketones and aldehydes.15

N-tert-Butyloxy carbonyl groups can be removed by using TBAF in refluxing THF.16

Under phase transfer conditions, selective mono- and dialkylation of malononitrile have been achieved by using neat TBAF with potassium carbonate or potassium tert-butoxide.17
References