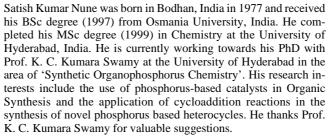
SPOTLIGHT 1221

SYNLETT Spotlight 65

This feature focuses on a reagent chosen by a postgraduate, highlighting the uses and preparation of the reagent in current research

Mitsunobu Reagent [Triphenylphosphine (TPP) and Diethyl Azodicarboxylate (DEAD)/Diisopropyl azodicarboxylate (DIAD)]

Compiled by Satish Kumar Nune



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Dedicated to Professor Mitsunobu who died on the 4th April 2003.



Introduction

The Mitsunobu reaction is perhaps the most favored method for the inversion of chiral secondary alcohols.¹ It involves the reaction of an acid component (RCO₂H) with a mixture of triphenylphosphine (TPP), diethyl azodicarboxylate (DEAD)/diisopropyl azodicarboxylate (DIAD), and an alcohol (Scheme 1).¹ It is supposed that the initial nucleophilic addition of TPP to DEAD/DIAD affords the betaine I;^{2,3} attack of an acid followed by an alcohol on I leads to the formation of a highly reactive alkoxyphosphonium intermediate II which undergoes nucleophilic displacement with virtually complete inversion of configuration at the electrophilic center under mild and nearly neutral conditions. The traditionally used acidic

component can be replaced by metal halides (LiBr), ^{4a} silanols, ^{4b} amides/ imides, ^{4c} nitronates, ^{4d} fluorinated alcohols ^{4e} and compounds possessing an active methylene group, ^{4f} thus rendering the reaction widely applicable in organic synthesis.

Scheme 1

Abstracts

(A) Epoxides are obtained from acyclic 1,2-diols^{5a,b} and cyclic *trans*-1,2-diols^{5c} under the Mitsunobu conditions. 1,1-Disubstituted-1,2-diols lead to the formation of aldehydes or ketones.^{5d}

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(B) Azidation of alcohols^{6a} and α -hydroxy(alkyl)phosphonates^{6b} can also be achieved under Mitsunobu conditions.

$$(EtO)_2P \longrightarrow R \xrightarrow{HN_3/ \text{ Benzene}} (EtO)_2P \longrightarrow F$$

$$OH \longrightarrow R$$

$$TPP/DEAD \longrightarrow R$$

$$N_3$$

(C) The Mitsunobu reaction can be used to distinguish between alcohol and phenol hydroxyls in esterification reactions.⁷

(D) Kinetic resolution of the racemic secondary alcohols^{8a} can be effected by using the enantioselective Mitsunobu reaction of these alcohols with phthalimide in the presence of a chiral tri-coordinate phosphoramidite.^{8b}

(E) Single-step reductive deoxygenation of unhindered alcohols takes place under Mitsunobu conditions.⁹

$$\begin{array}{c} \text{CH}_2\text{OH} & \text{CH}_2 \\ \hline & \text{TPP/DEAD} \\ \hline \text{NDSH} & \\ \hline \\ \text{RCH}_2\text{OH} & \hline \\ \hline \text{NDSH, THF} & \text{RCH}_2\text{N}(\text{NH}_2)\text{SO}_2\text{Ar} \longrightarrow \boxed{\text{RCH}_2-\text{N}=\text{NH}} \\ \hline -N_2 & \text{RCH}_3 \\ \hline \end{array}$$

NDSH = O- Nitrobenzene sulphonyl hydrazine

(F) Intramolecular Mitsunobu reaction of 2-(hydroxyl iminomethyl) benzyl alcohols or 1,2-aminoalcohols gives benzoxazines^{10a} or aziridines^{10b} respectively.

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