2-Cyanoethyl \(N,N,N',N'\)-tetraisopropylphosphorodiamidite

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Introduction

2-Cyanoethyl \(N,N,N',N'\)-tetraisopropylphosphorodiamidite is a colorless viscous liquid, which is soluble in most organic solvents. It is a widely used phosphitylating reagent for the preparation of various phosphorylated biomolecules, such as nucleoside carbohydrate conjugates, phospholipids and glycopeptides. In particular, this reagent is highly effective for automated solid-phase DNA/RNA oligonucleotide synthesis.

2-Cyanoethyl \(N,N,N',N'\)-tetraisopropylphosphorodiamidite has shown great utility in the coupling of nucleobases or carbohydrates via their phosphotriesters in the presence of activators such as \(1H\)-tetrazole, in moderate yields under mild conditions.\(^2\) Additionally, 2-cyanoethyl \(N,N,N',N'\)-tetraisopropylphosphorodiamidite is cheaper and more stable than 2-cyanoethyl \(N,N\)-diisopropyl-chlorophosphorodiamidite, the other commonly used phosphinylation reagent.\(^3\)

Abstracts

(A) 2-Cyanoethyl-\(N,N,N',N'\)-tetraisopropylphosphorodiamidite was used by Sheppard and co-workers to prepare carbohydrate phosphoramidites as nucleoglycoconjugate building blocks in good yield in the presence of disopropylammonium tetrazolide under anhydrous conditions. Then, the monosaccharide phosphoramidite was coupled with DNA oligonucleotides by solid-phase chemistry.\(^4\)

(B) Recently, Yamada and co-workers used 2-cyanoethyl \(N,N,N',N'\)-tetraisopropylphosphorodiamidite to synthesize the uridine 3'-phosphoramidite building block in good yield with disopropylammonium tetrazolide as a catalyst under anhydrous conditions, for developing oligonucleotides containing new 2'-O-modified ribonucleosides as nucleic acid based drugs.\(^5\)
(C) Lin and colleagues used 2-cyanoethyl N,N,N′,N′-tetraisopropylphosphoramidite as the phosphorylating reagent in the presence of disopropylammonium tetrazole to couple with 2′,3′-di-O-acetyl-adenosine to generate boron-containing ADP analogues (in an overall yield of 36%).

(D) Smith and co-workers developed an efficient method to prepare aldose phosphate diesters using 2-cyanoethyl N,N,N′,N′-tetraisopropylphosphoramidite. A 5-O-protected diol was firstly reacted with the phosphorylating reagent and 1H-tetrazole as an activator at room temperature, followed by oxidation, generating cyclic phosphate triester diastereoisomers in high yield.

(E) 2-Cyanoethyl- N,N,N′,N′-tetraisopropylphosphoramidite was used to prepare glycoconjugate polymers which carry GGPL analogues, bioactive segments of main cell membrane glycolipids of Mycoplasma fermentas. Therein, Nishida and co-workers reacted 4-nitrophenyl 2,3,4-tri-O-benzyl-α-d-glucopyranoside with 2-cyanoethyl N,N,N′,N′-tetraisopropylphosphoramidite in the presence of 1H-tetrazole, then reacted with choline tosylate, followed by oxidation and removal of the cyanoethyl group, generating 4-nitrophenyl 2,3,4-tri-O-benzyl-6-O-phosphorylcholine-α-D-glucopyranoside (in an overall 54% yield).

(F) Rodriguez and co-workers reported the synthesis of glucose-nucleoside conjugates as anti-HIV produgs by using 2-cyanoethyl N,N,N′,N′-tetraisopropylphosphoramidite as the phosphorylating reagent. Glucosyl phosphoramidite was firstly prepared in the presence of pyridinium trifluoroacetate under anhydrous conditions, and then coupled with nucleosides generating the desired compounds.

References


(8) Smith, J. M.; Borsenberger, V.; Raftery, J.; Sutherland, J. D. Chem. Biol. 2004, 1, 1418.
