

Description

dNTP Mix is a mixture of equal concentration (10mM) of dGTP, dTTP, dATP, and dCTP in a buffer (pH 7.5), which purity is above 99%. This mixture is necessary for DNA amplification PCR. The absence of exodeoxyribonucleases and ribonucleases is confirmed by appropriate quality test. The quality stability of dNTP Mix is confirmed through PCR.

Applications

For direct use in PCR, long PCR, RT-PCR, cDNA synthesis, primer extension, DNA sequencing and DNA labeling.

Storage

-20°C, Avoid multiple freeze/thaw cycles. Aliquoting is recommended.

Method	Specification	Result
Purity Assay		
HPLC (column C18; detection UV at 271; mobile phase:A=TEAA 0.1M, pH 7.0; B=60% CH ₃ CN/A)	>99%	dATP 99.2% dGTP 99.2% dCTP 99.3% dTTP 99.4%
LO test(test for detection of exo-, endo-deoxyribonuclease and phosphatase contaminants)	Not detectable	passed
Ribonuclease assay (test for detection of RNase contaminants using [3H]-RNA as a substrate)	Not detectable	passed
Function Assay		
PCR with <i>Pfu</i> and <i>Taq</i> DNA Polymerases	Production of 1000 bp PCR fragment from 2ng of genomic DNA	passed
рН		
	7.0-7.5	7.5
Concentration		
Spectrometry	Each 10mM	Each 10mM

