

# KOD Dash

Code No. LDP-101

Lot No. \*\*\*\*\*

Size 250units

Concentration : 2.5 units/ $\mu$ L

Unit Definition : One unit of enzyme is defined as the amount of enzyme that will incorporate 10 nmoles of dNTPs into acid insoluble material in 30 minutes at 75°C.

Assay Condition :

20	mM	Tris-HCl(pH7.5 at 25 °C)
8	mM	MgCl <sub>2</sub>
7.5	mM	DTT
2.5	$\mu$ g	BSA
150	$\mu$ M	each of dATP,dGTP,dCTP,dTTP(a mix of unlabeled and [ <sup>3</sup> H]-dTTP)
7.5	$\mu$ g	activated calf thymus DNA per 50 $\mu$ L reaction

Storage Buffer :

50	%	Glycelrol
50	mM	Tris-HCl(pH8.0 at 25 °C)
0.1	mM	EDTA
50	mM	KCl
1	mM	DTT
0.1	%	Tween <sup>®</sup> -20
0.1	%	Nonidet <sup>®</sup> P-40

Materials Provided : 10  $\times$  PCR Buffer  
 dNTPs : 2mM dATP,dGTP,dCTP,dTTP each

## Quality Control

1. Nicking Activity : When 10units of this enzyme were incubated with 1  $\mu$ g of pBR322 for 1 hour at 75°C ,no nicking activity was observed after agarose gel electrophoresis.
2. Nuclease Activity : When 10units of this enzyme were incubated with 1  $\mu$ g of  $\lambda$  / HindIII-DNA for 1 hour at 75°C ,no nuclease activity was observed after agarose gel electrophoresis.
3. PCR Assay : The 8.5 kbp  $\beta$ -globin gene fragment could be amplified using human genomic DNA.

This enzyme is produced as recombinant in *E.coli* and has very high amplification efficiency. Therefore, there is a possibility to amplify *E.coli* genomic DNA when using rRNA primers.



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