Storage Conditions (−20 °C)
Shipping Conditions (−20 °C)

TOYOBO
Ideas & Chemistry

KOD -Plus- Ver.2

Code No. KOD−211

Lot No.

Size 200units

Concentration	:	KOD DNA polymerase	1.0	unit/μl
		anti−KOD DNA polymerase antibody	1.6	mg/ml

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℃.

Assay Condition :

20	mM	Tris−HCl (pH7.5 at 25 ℃)
8	mM	$MgCl_2$
7.5	mM	DTT
2.5	μg	BSA
150	μM	each of dATP,dGTP,dCTP,dTTP(a mix of unlabeled and [^{3}H]−dTTP)
7.5	μg	activated calf thymus DNA per 50μl reaction

Storage Buffer :

50	%	Glycerol
50	mM	Tris−HCl(pH8.0 at 25 ℃)
0.1	mM	EDTA
1	mM	DTT
0.001	%	Tween−20
0.001	%	Nonidet P−40

Materials Provided
10x PCR Buffer for KOD −Plus− Ver.2
25mM $MgSO_4$
dNTPs : 2mM dATP,dGTP,dCTP,dTTP each

Quality Control

1.Nicking Activity : When 15units of this enzyme were incubated with 1μg of pBR322 for 4 hours at 75℃, no nicking activity was observed after agarose gel electrophoresis.

2.PCR Assay : The 6.8 kb cDNA fragment of DNA Polymerase ε gene could be amplified from HeLa total RNA reverse transcript.

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

manual

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