

KOD -Plus-Code No. **KOD-201**Lot No. *********Size **200units**

Concentration : KOD DNA polymerase 1.0 unit/ μ l
 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°C.

Assay Condition : 20 mM Tris-HCl(pH7.5 at 25 °C)
 8 mM MgCl₂
 7.5 mM DTT
 2.5 μ g BSA
 150 μ M each of dATP,dGTP,dCTP,dTTP(a mix of unlabeled and [³H]-dTTP)
 7.5 μ g activated calf thymus DNA per 50 μ l reaction

Storage Buffer : 50 % Glycerol
 50 mM Tris-HCl(pH8.0 at 25 °C)
 0.1 mM EDTA
 1 mM DTT
 0.001 % Tween™ 20
 0.001 % Nonidet™ P-40

Materials Provided 10 × PCR Buffer for KOD -Plus-
 25mM MgSO₄
 dNTPs : 2mM dATP,dGTP,dCTP,dTTP each

Quality Control

1. Nicking Activity : When 15units of this enzyme were incubated with 1 μ g of pBR322 for 4 hours at 75°C ,no nicking activity was observed after agarose gel electrophoresis.
2. PCR Assay : The 4.0 kbp fragment of p53 gene could be amplified using 5ng 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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