Proteinase K

For Research Use Only

Cat. No.: MO⁰εγι **Store at:** ۲-Λ °C

Concentration: Y ⋅ mg/ml Quantity: \ml

Description:

Proteinase K is an endolytic protease that cleaves peptide bonds at the carboxylic sides of aliphatic, aromatic or hydrophobic amino acids. The Proteinase K is classified as a serine protease. The smallest peptide to be hydrolyzed by this enzyme is a tetrapeptide.

Applications:

Isolation of genomic DNA from mouse tail. Isolation of genomic DNA from cultured cells. Removal of DNases and RNases when isolating DNA and RNA from tissues or cell lines. Determination of enzyme localization.

Improving cloning efficiency of PCR products.

Source: Pichiapastoriscells with a cloned gene from Tritirachium album.

Molecular Weight: ۲۸,۹kDa monomer

Storage Buffer:

The enzyme is supplied in: $\cdot mM$ Tris-HCl (pH V, \circ), containing calcium acetate and $\circ \cdot (v/v)$ glycerol.

Inhibition and Inactivation:

Inhibitors: Proteinase K is not inactivated by metal chelators, by thiol-reactive reagents or by specific trypsin and chymotrypsin inhibitors. Phenylmethylsulfonyl fluoride and Diisopropyl phosphorofluoridate completely inhibit the enzyme.

Inactivated by heating at $\operatorname{O^{\circ}C}$ for $\Upsilon \cdot$ min.

Note:

The recommended working concentration for Proteinase K is $\cdot, \cdot \circ$ - $\mbox{mg/ml}$. The activity of the enzyme is stimulated by \cdot, \cdot - \mbox{M} SDS or by $1-\xi$ M urea.

 Ca^{+} protects Proteinase K against autolysis, increases the thermal stability and has a regulatory function for the substrate binding site of Proteinase K.

Stable over a wide pH range: $\xi, \cdot - 11, \circ$, optimum pH V, $\circ - \Lambda, \cdot$

Quality Control Assay Data

Endodeoxyribonuclease Assay:

No conversion of covalently closed circular DNA to nicked DNA was detected after incubation of $\xi \cdot \mu g$ of Proteinase K with μg of pUC19 DNA for ξ hours at $\gamma \gamma \circ C$.

Ribonuclease Assay:

No detectable RNA degradation after incubation of $\Lambda \cdot ng$ of Ykb RNA transcript with $\xi \cdot \mu g$ of Proteinase K for ξ hours at $\gamma \gamma \circ C$.

Labeled Oligonucleotide (LO) Assay:

No degradation of single-stranded and double-stranded labeled oligonucleotide was observed after incubation with $\xi \cdot \mu g$ of Proteinase K for ξ hours at $V^{\circ}C$.

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