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Datasheet for ABIN5519440

EasyTaq® DNA Polymerase for PAGE

Overview	
Quantity:	2500 units
Application:	Polymerase Chain Reaction (PCR)
Product Details	
Purpose:	EasyTaq® DNA Polymerase for PAGE is purified from E. coli expressing a cloned DNA
	polymerase from Thermus aquaticus.
Brand:	EasyTaq®
Specificity:	The enzyme consists of a single polypeptide with a molecular weight of approximately 94 kDa.
	EasyTaq® DNA Polymerase for PAGE has 5'-3' DNA polymerase activity and 5'-3' exonuclease
	activity. It lacks 3'-5' exonuclease activity. This enzyme is supplied with unique buffer, and its
	PCR product is suitable for SDS-PAGE and agarose gel electrophoresis.
Characteristics:	- Extension rate is about 1-2 kb/min.
	- Unique buffer system compatible with PAGE.
	- Template-independent "A" can be generated at the 3' end of the PCR product. PCR products
	can be directly cloned into pEASY®-T vectors.
	- Amplification of genomic DNA fragment up to 3 kb.
Components:	DNA Polymerase, 10X Taq Buffer, 6X DNA Loading Buffer
Unit Definition:	One unit of EasyTaq® DNA Polymerase for PAGE incorporates 10 nmol of deoxyribonucleotide
	into acid-precipitable material in 30 minutes at 74°C.
Application Details	
Application Notes:	Short fragment PCR
Comment:	EasyTaq® DNA Polymerase for PAGE has passed the following quality control assays:
	functional absence of double- and single-strand endonuclease activity, >99% homogeneous

measured by SDS-PAGE. Each batch of EasyTaq® DNA Polymerase for PAGE has been

Application Details

assayed for amplification efficiency to amplify p53 gene from 10 ng of human genomic DNA.
For Research Use only
Storage Buffer: 20 mM Tris-HCl (pH 8.0), 0.1 mM EDTA, 1 mM DTT, 100 mM KCl, 50 % glycerol,
stabilizers
10xEasyTaq® Buffer for PAGE (with Mg2+): 200 mM Tris-HCl (pH 8.3), 200 mM KCl, 100 mM
(NH4)2SO4, 20 mM MgSO4, others
-20 °C
at -20°C for two years
24 months
Johnson, Drugan, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (
1991)