

# KlenTaq DNA Polymerase

**Catalog Number: NP041011220 - 500U** 

## **Description:**

This product is suitable for mutation analysis with mutation-specific oligonucleotides. It has a very low background ability to extend a mismatched 3'-oligonucleotide end making it suitable for mutation analysis with mutation-specific oligonucleotides. Amplicons are T/A cloning compatible.

KlenTaq DNA Polymerase has no the N-terminal portion of the gene, encoding Thermus aquaticus (Taq) DNA polymerase, leaving a highly active and even more thermal stable DNA polymerase activity.

KlenTaq has a wide range of optimal MgCl2 concentration. The optimal range of Mg2+ concentration for KlenTaq is broader than for the majority of thermostable polymerases. The mutation rate during polymerization is two-fold lower for KlenTaq in comparison with full-length Taq DNA polymerase.

#### **Contents:**

Components	500U
KlenTaq DNA poly. 5 U/μl	500U
MgCl₂ Solution 25 mM	1 mL
10X Buffer MgCl <sub>2</sub> free	1 mL

### **General Reaction Protocol:**

- 1. Thaw 10X reaction buffer, dNTP mixture.
- 2. Mix the master mix thoroughly and dispense appropriate volumes into PCR tubes or plates.
- 3. Add templates DNA to the individual PCR tubes or wells containing the master mix.

Component	Volume	Final conc.
10X Reaction Buffer	2 μL	1X
MgCl <sub>2</sub> Solution 25 mM	2.4 μL	3 mM
40 mM dNTPs Mix	0.51	0.25 14
(10 mM each)	0.5 μL	0.25 mM
Forward Primer	1l	0.5
(10 pmol/ μL)	1 μL	pmoles/μL
Reverse Primer	11	0.5
(10 pmol/ μL)	1 μL	pmoles/μL
Template DNA	Variable	10 fg to 1 μg
KlenTaq DNA poly.	0.25	
(5 units/μl)	0.25 μL	
PCR grade water	Up to 20μL	
	final volume	<u>-</u>
Total Volume	20 μL	

4. Program the PCR machine according to the program outlined.

Cycle	Time	Temp °C
1	4 min	95
	30 sec	94
30 - 35	30 sec	57
	30-60 sec	72
1	5 min	72

#### Notes:

- # Extension temperature is between 68 and 72 °C. We highly recommend 68 °C for more efficiency of Klen Taq DNA polymerase.
- \* For PCR products longer than 3~4 Kb, use an extension time of approximately 1 min. per Kb DNA.

## **Agarose Gel Electrophoresis:**

Run the total 5-7  $\mu L$  of PCR products alongside 3  $\mu L$  DNA marker on a 2% agarose gel containing Green Viewer Dye DNA safe stain.