

Taq DNA Polymerase

Cat: C101001 (250 U, 50 μ L) Cat: C101002 (500 U, 100 μ L)

Store at -20°C

Contents:

Component	C101001	C1011002
Taq DNA poly. 5 U/μl	250 U	500U
MgCl ₂ Solution 25 mM	0.5 ml	1 ml
10X Buffer MgCl₂ free	0.5 ml	1 ml

Description:

Taq DNA Polymerase is a high quality purified recombinant enzyme and catalyze 5'→3' synthesis of DNA. The enzyme has no detectable 3'→5' proofreading exonuclease activity. It is provided with 10X reaction buffer that will enable or improve suboptimal PCR caused by templates that have a high degree of secondary structure or that are GC-rich.

Kit storage:

This kit should be stored at -20 °C. Under this condition reagents are stable for two years from the date of production.

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.

Volume	Final Conc.
2 μL	1X
1.2 μL	1.5 mM
0.41	0.2 mM
0.4 μι	
1 μL	0.5
	pmoles/μL
1	0.5
1 μι	pmoles/μL
Variable	10 fg~1 μg
Variable	-
0.25 μL	
20 μL	-
	2 μL 1.2 μL 0.4 μL 1 μL 1 μL Variable Variable 0.25 μL

- 3. Add templates DNA to the individual PCR tubes or wells containing the master mix.
- 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 Pars Tous Tag DNA polymerase.

- * For PCR products longer than 3~4 Kb, use an extension time of approximately 1 min. per Kb DNA.
- * A DNA fragment which is amplified by *Taq* DNA polymerase has A-overhang, and it enables you to do cloning by using T-vector

Agarose gel Electrophoresis:

Run the total 5-7 μ L of PCR products alongside 3μ L DNA marker on a 2% agarose gel containing Green viewer DNA safe stain.

Disclaimers and Addresses:

This product is for **Research Use Only** and should only be used by trained professionals.

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