

Technical Data Sheet

GENERAL INFORMATION

PRODUCT *Taq* DNA Polymerase, expressed in yeast. Glycerol-free.

Cat. No. MT01U-G1TAQPXA

UNITS 1,000 U

DESCRIPTION *Taq* DNA Polymerase is a highly thermostable DNA polymerase from the thermophilic bacterium *Thermus aquaticus*. This enzyme catalyzes the synthesis of new DNA strands by adding nucleotides in the 5' → 3' direction; while it lacks 3'→5' exonuclease activity. In addition, *Taq* Polymerase exhibits deoxynucleotidyl transferase activity, which frequently results in the addition of extra adenines at the 3'-end of PCR products. *Taq* DNA Polymerase withstand high temperatures, making it suitable for PCR amplification in thermocycling processes.

This product is supplied in a glycerol-free solution and it is free from *E. coli* DNA contaminants.

PRODUCTS PROVIDED

Component		Amount
01U-G1TAQPXA	<i>Taq</i> DNA Polymerase. Glycerol-free.	1 vial x 100 µL
BTAQPXA	10X <i>Taq</i> DNA Polymerase Reaction Buffer.	1 vial x 1.5 mL

DELIVERY CONDITIONS

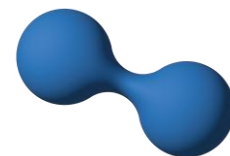
01U-G1TAQPXA 100 µL of *Taq* DNA Polymerase at 10 U/µL in 20 mM Tris pH 7.5, 300 mM KCl, 0.1 mM EDTA, 1 mM DTT, 0.05% Tween-20.

BTAQPXA 1.5 mL of 200 mM Tris pH 8.8, 250 mM KCl, 15 mM MgCl₂, 10 mM DTT, 0.1% Tween-20, 5 mg/mL rBSA.

SHIPPING CONDITIONS This product requires cold shipment conditions. Store the protein in a -20 °C to -80 °C range upon arrival.

STORAGE CONDITIONS Store it at a temperature range from -20 °C to -80 °C for medium and long term. Storage at 4 °C is possible for short term. Avoid multiple freeze/thaw cycles by storing multiple aliquots.





ADDITIONAL INFORMATION

ACTIVITY UNIT DEFINITION

One unit of activity is defined as the amount of enzyme needed to incorporate 15 nmol of dNTPs (deoxyribonucleotides) into a DNA fragment, in 30 minutes at 72 °C.

RECOMMENDED REACTION CONDITIONS

1. Gently vortex and briefly centrifuge all solutions.
2. Prepare the following mix on ice.

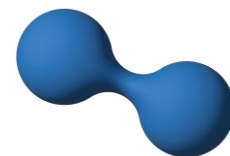
Component	Final concentration
10X <i>Taq</i> DNA Polymerase Reaction Buffer	1X
dNTPs, 10 mM each	0.2 mM each dNPT
10 µM Forward Primer	0.1-1 µM
10 µM Reverse Primer	0.1-1 µM
Template DNA	<0.5 µg/50 µL
<i>Taq</i> DNA Polymerase	5 U/50 µL
Nuclease-free water	To 50 µL

3. Transfer tubes from ice to a PCR machine with a preheated lid to 95 °C. If using a thermocycler without a heated lid, overlay the reaction mixture with mineral oil to prevent evaporation.
4. Perform PCR using recommended thermal cycling conditions:

Step	Temperature	Time	Number of cycles
Initial denaturation	95 °C	3 min	1
Denaturation	95 °C	30 s	25-35
Annealing	42-65 °C*	30 s	
Extension	72 °C	1 min/kb	
Final extension	72 °C	5 min	1

*Annealing temperature should be optimized for each primer set based on primer T_m .





QUALITY CONTROL

Taq DNA POLYMERASE ACTIVITY

Taq DNA Polymerase activity is measured for each batch to verify that the resulting units are the same to the reference units, with an accepted 8% deviation.

E. coli DNA CONTAMINATION

5 U of *Taq* DNA Polymerase is screened for the presence of the specific gene *ybbW* from *Escherichia coli*.¹ A C_q value higher than 35 is accepted.

¹Walker, David I., et al. "A highly specific *Escherichia coli* qPCR and its comparison with existing methods for environmental waters." *Water research* 126 (2017): 101-110.

TECHNICAL SUPPORT

If you have any questions, feel free to contact us at
support@levprot.com

Consult the Safety Data Sheet for information regarding hazards and safe handling practises.

THIS PRODUCT IS INTENDED FOR RESEARCH USE ONLY.

DATE 16/05/2025

REVISION 04

