

Product Manual

Product Name Hemo Taq DNA Polymerase (Blood-resistant)

Source Recombinantly expressed in Escherichia coli

Catalog Number CSB-DEM044

Physical FormLiquidEnzyme Activity5U/μLStorage Conditions -20 ± 5 °C

Storage Buffer 10 mM Tris-HCl, 100 mM KCl, 1 mM DTT, 0.1 mM EDTA, 0.5% Tween 20,

0.5% NP-40, 50% Glycerol

The amount of enzyme required to incorporate 10 nmol of

Activity Definition deoxynucleotide into acid-insoluble material in 30 minutes at 75°C is

defined as 1 unit (U).

Quality Control No detectable exonuclease and endonuclease activities

Shelf Life 24 months

Product Description

HemoTaq DNA Polymerase (Blood-resistant) is a sensitive and heat-resistant Taq DNA polymerase that has been upgraded and recombinantly expressed. It is modified using an antibody method and possesses both 5'-3' polymerase activity and 5'-3' exonuclease activity. With an optimized buffer system, it exhibits strong tolerance to impurities, especially for direct amplification of blood samples, demonstrating excellent performance.

Product components

Component No.	Component Name	Specifications		
1	5×Hemo buffer	1.2mL	6mL	12mL



2	Hemo Taq DNA Polymerase	500U	2500U	5000U	

Operating instructions

Recommended Reaction System

Composition	Addition amount
ddH ₂ O	Up to 30 μL
5×Hemo buffer	6 μL
10 mM dNTPs	0.6 μL
Upstream primer (10 μM)	0.6 μL
Downstream primer (10 μM)	0.6μL
HemoTaq DNA Polymerase	0.3 μL
Blood template	0.5-5μL

Note:

A.Primer Concentration:

Generally, a primer final concentration of 0.2 µM in the reaction system yields good results. If the reaction performance is poor, adjust the primer concentration within the range of 0.1 μM to 1.0 μM .

B.Template Concentration:

The optimal concentration range for whole blood templates is 0.5%-20%. A recomm-



ended amount of 5% is suggested as an initial trial condition, which is adding 1.5µL of whole blood as a template in a 30µL reaction system. Be cautious to avoid drawing blood clots.

C.Polymerase Concentration:

It is recommended to use $0.05U/\mu L$. The enzyme amount can be adjusted between 0.25 - 1uL. Generally, increasing the enzyme amount can enhance the amplification yield, but it may lead to a decrease in specificity.

D.Quantitative PCR with fluorescence:

When using this product for quantitative PCR with fluorescence, add 0.3 µL of 10μM TaqMan Probe in the recommended system mentioned above.

Recommended PCR reaction procedure

Temperature	Time	Cycles
95°C	5 min	1
95°C	10 s	
45-68°C	30 s	30-35
72°C	1 kb/min	
72°C	5min	1



Note:

E.Quantitative PCR Program with fluorescence (TaqMan Probe method):

95°C for 5 min; 95°C for 10s, 60°C for 30s \star , 40-45 cycles.

F.Annealing temperature and time:

The annealing temperature should be adjusted based on the Tm value of the primers, and the annealing time can be adjusted within 10-30 seconds.