

# **MorreTaq Hot Start DNA Polymeras**

## The Hot Start DNA Polymerase For High sensitivity

MorreTaq Hot Start DNA Polymerase for qPCR is designed for Real-Time PCR and Hot-start PCR. A special inhibition the reaction at room temperature until after the first denaturation step. This prevents primerdimers and other artefacts. The enzyme is a thermostable DNA polymerase that possesses a  $5' \rightarrow 3'$ polymerase activity and a double-stranded specific  $5' \rightarrow 3'$  exonuclease activity. The enzyme consists of a single polypeptide with a molecular weight of 94kDa.

#### **Applications**

- Hot Start and real time PCR
- Multiplex PCR
- · Amplification of complex genomic and cDNA templates

#### **Storage buffer**

50mM Tris-HCl pH7.9, 50mM KCl, 0.1mM EDTA, 1mM DTT, 0.5mM PMSF, 50% lycerol

#### **Storage conditions**

Storage at -20°C

#### **10X reaction buffer**

Buffer A: high quantity (genomic DNA PCR) containing 15mM MgCl<sub>2</sub> . Buffer B: high sensitivity (RT-PCR) containing 15mM MgCl<sub>2</sub>

#### **Unit description**

One unit is defined as the amount of enzyme that will incoporate 10 nmole of dNTP into acidinsoluble material in 30 minutes at 74oC. The reaction conditions are: 50mM Tris-HCl pH8.8, 50mM NaCl, 5mM MgCl<sub>2</sub>, 200uM each of dATP, dCTP, dGTP, H3dTTP, 10 ug activated calf thymus DNA and 0.1mg/ml BSA in a final volume of 50 ul.

#### Storage

50% glycerol (v/v), 20 mM Tris-HCl pH 8.7 at -20°C, 100 mM KCl, 0.1 mM EDTA.

#### Source

E.coli clone

## **Quality control**

The enzyme is free of nicking and priming activities, exonucleases and non-specific endonucleases. SDS/PAGE - 95 kD band. Activity and stability tested via thermo-cycling. The error rate per nucleotide per cycle is ~ 2.5 x 10^-5; the accuracy is ~ 4 x 10^4. Estimated half life at 95°C is 1.5 hours.

#### **PCR reaction mix**

Component	Volume	
MorreTaq Hot Start	0.5.1.1	
DNA polymerase	0.3-141	
10X buffer	10 ul	
10mM dNTP	2 ul	
Primer1 (20 pmol)	2-4 ul	
Primer2 (20 pmol)	2-4 ul	
template	1-10 ul	
ddH <sub>2</sub> O	Up to 100 ul	
Total	100 ul	

### **PCR cycles**

Step	Temperature	Time	Cycle
Initial denaturation	94-95°C	10 mins	1
Denaturation	94-95°C	10-60 sec	
Annealing	50-68°C	10-30 sec	25-35
Extension	72°C	1min/1kb	
Final extension	72°C	1-10 mins	1

#### **IMPORTANT**

Annealing temperature should be 2-6°C lower than the primer melting temperature.

## **Ordering information**

Cat.	Pack	Con.
MHT500	500U	5U/ul
MHT2500	2500U	5U/ul

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#### About MORREBIO

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