



## Hot Start primer block PCR Mastermix (2x)

**For research use only**

Hot Start primer PCR Master (2x)	1 ml
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Cat No : YT1554

Size : 1 ml

Store at -20 ° c

Hot Start primer block PCR Mastermix (2x) is a ready-to-use mix containing Hot Start Taq- antibody DNA Polymerase and a HotStart Binding Protein. the Hot Start DNA Polymerase combines Taq DNA polymerase and a specific antibody that inhibits the polymerase activity at ambient temperature and activated by heat treatment. HotStart Binding Protein is a novel hot-start technology which inactivate primers. The optimized buffer and hot-start technology used in this premixed enhances the specificity and sensitivity of PCR .The green dye allows direct loading of PCR products on to gels for electrophoresis.

### Features:

- Easy reaction setup at room temperature
- Simply add primers and template
- High specificity due to antibody based hot start
- saves time and reduces contamination
- Minimizes amplification of non-specific products and primer dimers

### Protocol:

- 1- Thaw the master mix and primers
- 2- Mix thoroughly before use to avoid localized concentrations of salts
- 3- Spin vials briefly before use
- 4- For each 25-µL reaction, add the following components

Dye plus hotstart master mix2X	10 µl
Primer F	1 µl
Primer R	1 µl
Template DNA	<200ng
Water	to 20 µl
Total	20 µl

- 5- Gently vortex the samples and briefly centrifuge
- 6- Place the reactions in a thermal cycler.
- 7- Perform PCR using the recommended thermal cycling conditions outlined below:

Cycle	Duration of cycle	Temperature
1	15min	95°C
25-35	20-30 Sec	95°C
	20-50 Sec	50-65°C
	30-90 Sec	72°C
1	5-10min	72°C

\*\*For maximum yield and specificity, temperatures and cycling times should be optimized for each new target or primer pair.

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