

# 5x HOT FIREPol® Blend Master Mix

With 12.5 mM MgCl<sub>2</sub>

Cat. No.	Pack Size	Conc. (MgCl <sub>2</sub> )
04-27-00S25	0.1 ml SAMPLE (25 reactions)	12.5 mM
04-27-00125	1 ml (250 reactions)	12.5 mM
04-27-02025	20 ml (5000 reactions)	12.5 mM
04-27-10025	100 ml (25000 reactions)	12.5 mM

For in vitro use only

#### **Description:**

5x HOT FIREPol® Blend Master Mix is a premixed readyto-use solution containing all reagents required for PCR except template, primers and water.

HOT FIREPol<sup>®</sup> Blend Master Mix contains two carefully optimized enzymes – HOT FIREPol<sup>®</sup> DNA polymerase and a proofreading polymerase. This enzyme blend has both the 5'→3' exonuclease activity as well as the 3'→5' proofreading activity. HOT FIREPol<sup>®</sup> Blend Master Mix exhibits an increased fidelity (up to five fold) compared to HOT FIREPol<sup>®</sup>. Generated PCR products are compatible with blunt-end cloning procedures.

#### **Applications:**

Hot Start PCR

## Mix Composition:

- HOT FIREPol® DNA polymerase
- Proofreading enzyme
- 5x Blend Master Mix Buffer
- 12.5 mM MaCl<sub>2</sub>

1x PCR solution - 2.5 mM MgCl<sub>2</sub>

- 2 mM dNTPs of each
  1x PCR solution 200 μM dATP, 200 μM dCTP,
  200 μM dGTP and 200 μM dTTP
- BSA

#### **Shipping and Storage conditions:**

Routine storage: -20°C

Shipping and temporary storage for up to 1 month at room temperature or storage for up to 6 months at 2-8°C has no detrimental effects on the quality of 5x HOT FIREPol<sup>®</sup> Blend Master Mix.



Reaction setup at room temperature is highly recommended for HOT FIREPol® Blend Master Mix. We recommend using 5x HOT FIREPol® Blend Master Mix in any PCR application that will be visualized by agarose gel electrophoresis and ethidium bromide staining.

### Recommended PCR reaction mix:

Component	Volume	Final conc.
5x HOT FIREPol® Blend Master Mix	4 μΙ	1 x
Forward primer (10 pmol/µl)	0.2-0.6 µl	0.1-0.3 µM
Reverse primer (10 pmol/µl)	0.2-0.6 µl	0.1-0.3 µM
Template DNA	xμl	0.01-10 ng/µl
Add H₂O	Up to 20 µl	

### **Recommended PCR cycles:**

Operation	Temp.	Time	Cycles
Initial activation*	95°C	12-15 min	1
Denaturation	95°C	10-20 s	
Annealing	54-66°C	30-60 s	25-30
Elongation	72°C	20 s - 4 min	
Final elongation	72°C	5-10 min	

<sup>\*</sup> To activate the polymerase, include an incubation step at 95°C for 12-15 minutes at the beginning of the qPCR cycle.

#### Safety warnings and precautions:

This product and its components should be handled only by persons trained in laboratory techniques. It is advisable to wear suitable protective clothing, such as laboratory overalls, gloves and safety glasses. Care should be taken to avoid contact with skin or eyes. In case of contact with skin or eyes, wash immediately with water.

Some applications this product is used in may require a license which is not provided by the purchase of this product. Users should obtain the license if required.