

Multi HS Prime Taq Premix

(2X, for Multiplex PCR)

Product Name	Cat. No.	Size	
Multi HS Prime Taq	MH-7100	1.0 ml X 1	
Premix (2X)	IVIN-7 100	1.011111.1	

Package information

MH-7100	2X Multi HS Prime Taq Premix (1.0 ml X 1) - with HS Prime Taq DNA Polymerase, dNTPs mix., reaction buffer, enzyme stabilizer and loading dye
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Description

The Multi HS Prime Taq Premix is for multiplex PCR. This product contains a master mix whose composition and elements were specifically developed for multiplex PCR applications.

This product contains optimized concentrations of Hot-start Taq DNA Polymerase (HS Prime Taq DNA Polymerase, G-7000), dNTPs mixture, MqCl₂ and reaction buffer.

Multiplex PCR is a powerful technique that enables amplification of two or more products in parallel in a single reaction tube.

It is widely used in genotyping applications and different areas of DNA testing in research, forensic, and diagnostic laboratories.

Applications

- Genotyping applications (e.g., STR, VNTP analysis)
- Detection of pathogens/diagnostics
- Qualitative and semi-quantitative gene expression analysis

Protocol

The following 20 µl reaction volume can be used for PCR.

1. Prepare the following components to a PCR tube.

Components	Volume	
DW	add up to 20 μ l	
Multi HS Prime Taq Premix (2X)	10 <i>µ</i> l	
Upstream Primer (10 pmoles/μℓ)	0.5~2.0 <i>µ</i> l	
Downstream Primer (10 pmoles/μℓ)	0.5~2.0µl	
Template DNA*	×μl	

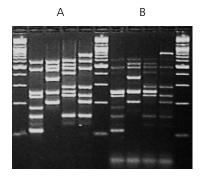
* Amount of template DNA: 10 ng ~ 250 ng

2. PCR cycling

Step	3-step PCR		Cycles
этер	Temp.	Time	Cycles
Initial denaturation	95℃	10 min	1
Denaturation Annealing Extension	95°C x°C 72°C	30 sec 30~60 sec 1 min	30~40
Final Extension	72℃	5 min	1

- 3. Separate the PCR products by agarose gel electrophoresis and visualize with EtBr or any other means.
- ► A DNA fragment which is amplified by Multi HS Prime Taq Premix has A overhang, and it enables you to do cloning by using T-vector.

Performance of Multi HS Prime Taq Premix



A: Multi HS Prime Taq Premix B: Normal Taq premix

Research Use Only