

# AddMulti Taq Master (2x Conc.)

Research Use Only

## Product Code

37101

## Component

1. AddMulti Taq Master (2x conc.) 1.0 ml

## Storage Condition

Store at -20°C

## Description

AddMulti Taq Master is supplied as a 2x concentrated master mixture type containing all the reagents needed to perform multiplex PCR. This product contains optimized concentrations of hot-start Taq DNA Polymerase, dNTPs mixture, MgCl<sub>2</sub> and reaction buffer for multiplex PCR.

Multiplex PCR is a powerful technique that enables amplification of more than two target genes 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.

## Components of AddMulti Taq Master as 2x conc.

20mM Tris-HCl (pH8.8), 100mM KCl, 0.2% Triton® X-100, 4mM MgCl<sub>2</sub>. Protein stabilizer, sediment, loading dye and 0.5 mM each of dATP, dCTP, dGTP, dTTP

## Storage and Stability

AddMulti Taq Master (2x conc.) is stable for 2 years when stored in a constant temperature freezer at less than -20°C.

Made in KOREA

This product was manufactured through ISO 9001 & 13485 system.

## Nucleic Acid Amplification Protocol

1. Add the following components to a thin-walled PCR tube:

Nuclease-Free Water	x $\mu$ l
AddMulti Taq Master (2x conc.)	10 $\mu$ l
Forward primer (10 $\mu$ M)	0.25~2.0 $\mu$ l
Reverse primer (10 $\mu$ M)	0.25~2.0 $\mu$ l
DNA template	x $\mu$ l
Total reaction volume	20 $\mu$ l

\* Recommendation for template DNA concentration in a 20  $\mu$ l reaction volume

- 1) Human genomic DNA: 0.1 ng ~ 1  $\mu$ g
- 2) Bacterial genomic DNA: 0.1 ng ~ 100 ng
- 3) Plasmid DNA: 0.01 ng ~ 5 ng

2. PCR cycling

Initial denaturation	95°C, 10 min
	95°C, 15 – 30 sec
PCR cycling (25 – 40 cycles)	55 - 65°C, 15 – 30 sec
	72°C, 30 sec per kb of product length
Final extension	72°C, 5 min
Hold	12°C, $\infty$