AccuPower® Multiplex PCR PreMix

[Cat. No.] Please refer to the **Ordering Information**

Introduction

AccuPower® Multiplex PCR PreMix is the powerful technology for convenient and easy performance that allows DNA amplification of two or more products in a single tube. By applying antibody-based HotStart *Top* DNA Polymerase, it provides reduced non-specific reactions such as mis-priming and primer dimer during PCR at a low temperature. This product contains vacuum-dried components including HotStart *Top* DNA Polymerase, dNTPs, reaction buffer, stabilizer, and tracking dye. It simplifies preparation of reaction mixture by adding template DNA and primers without any extra process. After the reaction, samples can be applied directly on agarose gel for analysis.

Applications

- STR analysis
- Molecular diagnostic analysis
- Qualitative, semi-qualitative gene expression assay
- Mutant screening
- Transgenic organism analysis
- Genotyping assay

Features & Benefits

- Multiplex PCR: Generation of 20 multiplexed amplification products in a single tube.
- Specificity & Efficiency: Minimized non-specific amplification and maximized PCR efficiency by using HotStart *Top* DNA Polymerase.
- Sensitivity: Excellent sensitivity and amplification efficiency even with small amounts of DNA.
- User-friendly: Reactants are individually packaged in each of the PCR tubes, it allows any user simply perform PCR by adding template DNA and primers.
- Stability: Included stabilizer enables delivery at room temperature and provides increased stability compared to solution-type products.
- Reproducibility: Mass production under ISO 9001 quality system allows minimized deviation between lots and reproducible results in replicated tests performed under same conditions and variation.

Composition

Composition	Concentration
HotStart *Top* DNA Polymerase	1 U
dNTPs (dATP, dCTP, dGTP, dTTP)	Each 250 μM
Reaction buffer with 2 mM $MgCl_2$	1X
Stabilizer and tracking dye	O

Specifications

HotStart *Top* DNA Polymerase	
5' to 3' exonuclease activity	No
3' to 5' exonuclease activity	No
3'–A overhang	Yes
Fragment size	Up to 1 kb

Storage

Store at -20°C. If stored in the recommended temperature, this product will be stable until the expiration date printed out on the label.

Precautions

This enzyme is specifically optimized for increasing base incorporation rate by inactivating 5' to 3' exonuclease activity. Therefore, this product is not recommended to use for real-time PCR using hydrolysis probe method.

Online Resources

Korean **English**

Visit our **product page** for additional information and protocols

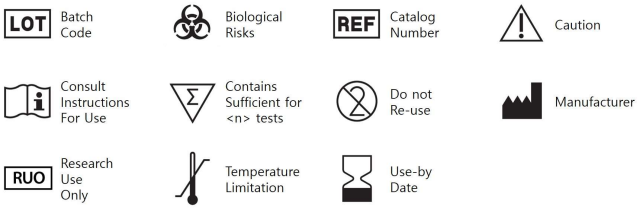
Ordering Information

Description			Cat. No.
0.2 ml thin-wall 8-tube strips with attached cap	96 tubes	20 μl/rxn	K-2111
		50 μl/rxn	K-2112
	480 tubes	20 μl/rxn	K-2113
		50 μl/rxn	K-2114

Notice

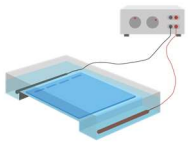
BIONEER corporation reserves the right to make corrections, modifications, improvements and other changes to its products, services, specifications or product descriptions at any time without notice.

Explanation of Symbols



Experimental Procedures

<table>
<tr><th colspan="2">Steps</th><th>Procedure Details</th></tr>
<tr><td>1</td><td> 
Preparation of reaction mixture</td><td>1. Add template DNA, primers, and nuclease-free water into AccuPower® Multiplex PCR PreMix tubes to make a total volume of 20 µl or 50 µl. Do not include the dried pellet.

• Preparation of reaction mixture
<table><tr><th>Components</th><th>20 µl reaction</th><th>50 µl reaction</th></tr><tr><td>Template DNA (1-100 ng)</td><td>Variable (1-10 µl)</td><td>Variable (1-25 µl)</td></tr><tr><td>Forward primer (1-5 pmol/µl)</td><td>0.5-2 µl</td><td>1-5 µl</td></tr><tr><td>Reverse primer (1-5 pmol/µl)</td><td>0.5-2 µl</td><td>1-5 µl</td></tr><tr><td>Nuclease-free water</td><td>Variable</td><td>Variable</td></tr><tr><td>Total volume</td><td>20 µl</td><td>50 µl</td></tr></table>
2. Dissolve the vacuum-dried green pellet by vortexing or pipetting, and briefly spin down.</td></tr>
<tr><td>2</td><td> 
Incubate reactions in a thermal cycler</td><td>3. Perform the reaction under the following conditions.
<table><tr><th>Step</th><th>Temperature</th><th>Time</th><th>Cycles</th></tr><tr><td>Pre-denaturation</td><td>95°C</td><td>10 min</td><td>1 cycle</td></tr><tr><td>Denaturation</td><td>95°C</td><td>30 sec</td><td></td></tr><tr><td>Annealing</td><td>55-65°C†</td><td>30-60 sec</td><td>25-35 cycles</td></tr><tr><td>Extension</td><td>72°C</td><td>1 min/kb</td><td></td></tr><tr><td>Final extension</td><td>72°C</td><td>5 min</td><td>1 cycle</td></tr></table>* Note: Primers are generally designed length of 24-35 nucleotides and ideally have a Tm value range within 5°C.
† Set the annealing temperature to 3-5 degrees lower than the Tm of the primers.</td></tr>
<tr><td>3</td><td> 
Analyze with gel electrophoresis</td><td>4. After the reaction, maintain the reaction mixture at 4-8°C. The samples can be stored at -20°C until use.

5. Load 5 µl of samples on agarose gel without adding a loading-dye mixture, and perform gel electrophoresis for analysis.</td></tr>
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