

[Cat. No.] **K-2952**

Introduction

AccuPower® P35S PCR Kit is a ready-to-use premix for PCR that can be used to detect virus derived P35S promoter to check whether soybeans or maize are genetically modified (GM).

Genetically modified organisms (GMOs) are organisms whose genetic material has been artificially modified to obtain desired characteristics, such as increased yield and resistance to pests and pathogens. Only those that have the approval of a regulatory agency, such as the Ministry of Food and Drug Safety in South Korea can be sold. Nowadays, many countries/regions require manufacturers to label products that include genetically modified products, leading to an increase in the demand for GMO detection technology.

This product contains vacuum-dried components specific to P35S promoter including DNA polymerase, primers, dNTPs, and reaction buffer required for PCR. This ready-to-use kit simplifies preparation of PCR mixture as the user only has to add template DNA and nuclease-free water. After the reaction, since tracking dye is included, the samples can be applied directly on agarose gel for analysis without adding extra solution.

Features & Benefits

- **Convenience & Reproducibility:** All reactants necessary for PCR including primers are lyophilized in each PCR tube, providing reproducible results in a convenient way.
- **Sensitivity:** By applying the patented PyroHotStart (Enzyme-mediated HotStart) technology that minimizes non-specific reactions and maximizes reaction efficiency, only the target gene can be effectively amplified even with a trace amount of template DNA.
- **Stability:** Included stabilizer in the PCR reaction mixture provides increased stability compared to solution-type products.

Composition

Composition	25 µl reaction
Top DNA Polymerase	1 U
dNTPs (dATP, dCTP, dGTP, dTTP)	Each 250 µM
Reaction buffer with 1.5 mM MgCl ₂	1X
Stabilizer and tracking dye	O
P35S Forward primer	0.5 µM
P35S Reverse primer	0.5 µM

* **Note:** For research use only. Not for use in diagnostic or therapeutic procedures.

Specifications

Top DNA Polymerase	
5'→3' exonuclease activity	No
3'→5' exonuclease activity	No
3'-A overhang	Yes
Fragment size	101 bp

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.

Online Resources



English

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

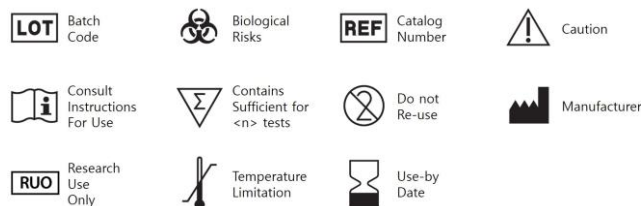
Ordering Information

Description	Cat. No.
AccuPower® P35S PCR Kit, 0.2 ml thin-wall 8-tube strips with attached cap / 96 tubes	K-2952




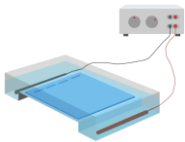
Notice

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Explanation of Symbols



Experimental Procedures

Steps		Procedure Details																								
1	 Add template DNA	<p>1. After preparing the template DNA and nuclease-free water, add the template DNA to the <i>AccuPower®</i> P35S PCR Kit.</p>																								
2	 Preparation of reaction mixture	<p>2. Add nuclease-free water into PCR tubes to make a total volume of 25 μl. (Do not include the volume of the dried premix in the PCR tubes.)</p> <p>3. Completely dissolve the vacuum-dried pellet by vortexing, and briefly spin down.</p>																								
3	 Incubate reactions in a thermal cycler	<p>4. Place PCR tubes on the thermal cycler.</p> <p>5. Perform the reaction under the following conditions.</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="text-align: center;">Step</th> <th style="text-align: center;">Temperature</th> <th style="text-align: center;">Time</th> <th style="text-align: center;">Cycles</th> </tr> </thead> <tbody> <tr> <td>Pre-denaturation</td> <td style="text-align: center;">95°C</td> <td style="text-align: center;">10 min</td> <td style="text-align: center;">1 cycle</td> </tr> <tr> <td>Denaturation</td> <td style="text-align: center;">95°C</td> <td style="text-align: center;">30 sec</td> <td></td> </tr> <tr> <td>Annealing</td> <td style="text-align: center;">60°C</td> <td style="text-align: center;">30 sec</td> <td style="text-align: center;">40 cycles</td> </tr> <tr> <td>Extension</td> <td style="text-align: center;">72°C</td> <td style="text-align: center;">30 sec</td> <td></td> </tr> <tr> <td>Final extension</td> <td style="text-align: center;">72°C</td> <td style="text-align: center;">7 min</td> <td style="text-align: center;">1 cycle</td> </tr> </tbody> </table> <p>* Note: Users can adjust the protocol according to their instrument and template sequences to get optimal results.</p>	Step	Temperature	Time	Cycles	Pre-denaturation	95°C	10 min	1 cycle	Denaturation	95°C	30 sec		Annealing	60°C	30 sec	40 cycles	Extension	72°C	30 sec		Final extension	72°C	7 min	1 cycle
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4	 Analyze with gel electrophoresis	<p>6. After the reaction, maintain the reaction mixture at 4-8°C.</p> <p>7. Load samples on agarose gel without adding a loading-dye mixture, and perform gel electrophoresis for analysis.</p>																								